# **CATALOG**14<sup>th</sup> Edition

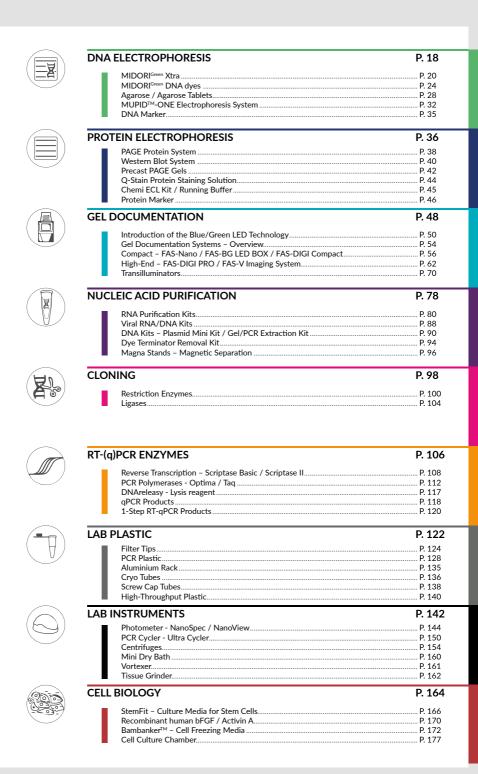








# CONTENT



#### **COMPANY PROFILE**

NIPPON Genetics EUROPE GmbH



#### NIPPON Genetics EUROPE - Our Company

NIPPON Genetics is a Japanese Life-Tech company, focused on cutting edge products for molecular and cell biology laboratories. We have been working to help researchers in making the bright future they envision a reality.

NIPPON Genetics EUROPE GmbH was founded in 2004. We are a growing group of highly motivated people with a strong background in life-science research. In the past few years, we have become a team that can support you not only with innovative products but also with advice on applications. This background also allows us to understand the needs of the customer and to develop new and exciting products for you.

We provide a wide variety of products, including gel documentation systems without UV-light, safe DNA-stains, cell freezing media, RT- (a) PCR enzymes, lab plastics and many more. We deliver these products to researchers around the globe working at universities, research institutes, pharmaceutical and biotech companies and/or clinical laboratories. Continuously growing our portfolio and expanding the cooperation with companies and scientists across the world, NIPPON Genetics EUROPE wants to provide the best service and solution for your research requirements.



#### **COMPANY PROFILE**

NIPPON Genetics EUROPE GmbH





#### **Our Philosophy**

The foundation of our business is contribution.

Things that researchers need for their research, things that help them, solutions for problems. We devote ourselves to providing researchers with these things, and this forms the foundation of our company.

As for the employees who support this foundation, we look for people with the human power required to make a positive contribution. Putting in the right kind of effort so that we can grow in the right way and help our customers make the important decisions when it matters. That's the kind of organization we are building.

#### Kazuo Yamazaki

CEO of NIPPON Genetics Co. Ltd in Japan

2011	2014	2016	2018	2020
•	•	•	•	•
Introduction of	Introduction of the	Introduction of the	ISO 9001:2015	New Office
the Cell Freezing	unique Blue/Green	RNA-Purification	Certification	Building with
Media Bambanker	Excitation Light	Kits	for Quality	increase in

Management

Warehouse size

#### Website

www.nippongenetics.eu



#### **Our Website**

Our website is the central information source for you! Here you can find all the information you need about our products. Whether you need a manual, MSDS, safety report or other material, just visit our website and download everything you need. We are always happy to receive your feedback about our service and products.

You can also find Technical Notes about many of our products, which we create with scientific enthusiasm in our laboratory. Furthermore, we get great feedback from the scientific society, which leads to the creation of various Application Notes.

Customers from Germany, Austria and the Netherlands can directly order products in our webshop. Every product page is in English or alternatively in German and a large number of product pages is also available in French.





# **Quality Management**

ISO 9001:2015 certified



#### ISO 9001:2015 certification

NIPPON Genetics EUROPE has always been a quality driven company: Quality of our products but also the quality of our service. It was important for us to show this quality-driven ideology and therefore we have decided to certify our quality management system according to ISO 9001:2015. We are very proud to announce that we were certified straight away in the first attempt, showing that our idea of analyzing, reflecting and improving is key to maintain our high-standard.

We see the ISO 9001:2015 as a central tool to improve our quality continuously. Hence, we will be performing a regular audit to ensure that the company still works according to the standard definide in ISO 9001:2015.



# **Management Team**

NIPPON Genetics EUROPE GmbH







#### Our Management Team

Dr. Jürgen Lünzer founded the company in 2004, with the support of our Japanese colleagues, mainly focused on international sales. In the beginning, the main objective was to start cooperations with other brands and distributors.

In the year 2010, Jürgen invited his long-term colleague Dr. Oliver Schwarz to join the company. Similar to Jürgen, Oliver had acquired deep knowledge about the Life-Science industries over the years and was therefore the perfect candidate to lead the German sales team.

In 2014, Dr. Marcelo Lanz joined the team for Product Management and International Sales. Due to the continuous successful growth of the NIPPON Genetics EUROPE product portfolio, Dr. Manuel Franke became part of the team in 2018 and is now operating as Tech Support Manager and in Research & Development of new products. Dr. Verena Krieger joined the company as a Marketing Manager in 2020. Since 2021, Dr. Jakob Maciejko contributes to the team as a new Product Manager.

We at NIPPON Genetics EUROPE see us a science-driven, technology-loving and innovation-seeking team still continuously looking for new opportunities to keep this success story going.

# Int. Sales and Marketing Team

NIPPON Genetics EUROPE GmbH









#### Become a GeneF@n member

GeneF@n is our exclusive club to get special promotions for our satisfied customers. Only GeneF@n members have the big advantage to receive continuous discounts and promotions on our products. Furthermore, as a GeneF@n member you can order free samples for our consumables.

Learn about new Application and Technical Notes. Here you will find helpful information to improve your experiments, which make your daily lab routine much easier.

GeneF@n





#### **National Sales Team**

NIPPON Genetics EUROPE GmbH



Postcode area	Supervisor	Contact information
01000 - 34999, 37000 - 39999, 99000 - 99999	Nenad Marić	nmaric@nippongenetics.de +49 (0)170 3422593
40000 - 54999	Dr. Kristina Kux	kkux@nippongenetics.de +49 (0)1713352548
35000 - 36999, 55000 - 79999, 88000 - 89999	Ina Dutzi-Kirchner	idutzi@nippongenetics.de +49 (0)1712012220
80000 - 87999, 90000 - 96999, 97000 -98999, Austria	Björn Krietemeyer	bkrietemeyer@nippongenetics.de +49 (0)15142116899

#### We care about you personally

We would be pleased to advise you personally on site and demonstrate to you the products in which you are interested. For Germany, Austria and the Netherlands, we offer our products directly to our customers. Just contact your personal product specialist of our national sales team and make an appointment for a product demonstration.

# Office & Logistic Management

NIPPON Genetics EUROPE GmbH









#### Our commitment

Our dedicated logistic team will prepare your shipment very carefully and fast to guarantee short delivery times. We deliver our products worldwide from our warehouse in Düren, Germany.

We educate young people to work with us and help our customers all over the world, as we believe that a central objective of a company is to transfer established knowledge to the next generation.



# Research & Development

NIPPON Genetics EUROPE GmbH



#### Our passion

We care about developing new products for our customers and commit to listening to their needs and requirements in the lab. Specified new products are tested, developed and continuously improved in our own Biosafety Level 1 laboratory.

In our workshop, we manufacture a selected group of our products with high quality awareness. Furthermore, we repair our instruments, in the rare cases of defects, by ourselves.

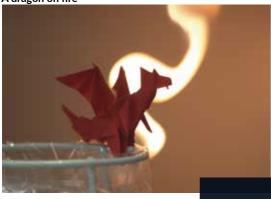




# **Origami - Competition**

NIPPON Genetics EUROPE GmbH

A dragon on fire



A dragon on Blue/Green light





#### Our winners of the Origami-Competition

Last year we organized our first Origami-Competition, many thanks to everyone who participated! Our customers folded their desired origami and sent us photos of them in a lab environment. Because of so many lovely and creative pictures, it was very difficult for us to select 3 winners. We hope that the lucky winners are happy with their prizes. Yes, we love dragons.



#### MIDORI Green Xtra Agarose Tablet

MIDORI<sup>Green</sup> Xtra is our latest, highly sensitive green fluorescent stain for safe visualization of DNA and RNA in agarose gels. It now comes in agarose tablet format. Just dissolve the tablet in buffer or water and cast your gel.

DNA Electrophoresis

Page: 28



#### FastGene® PAGE Protein System

The FastGene® PAGE Protein System contains all the necessary components for PAGE protein analysis and includes a hand-cast set for convenient gel casting. The system is also compatible with a wide range of pre-cast gels

Protein Electrophoresis

Page: 38

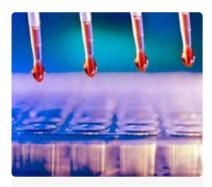


#### FastGene® Restriction Enzymes

The FastGene® Restriction Enzymes are perfect for routine cloning experiments. Use our Enzyme-Finder to search within 115 different enzymes for a suitable enzyme by name recognition sequence or overhang.

Cloning

Page: 100



#### FastGene® TP Filter Tips

FastGene® TP Filter Tips are transparent, for observable liquid control inside the pipette. For clear observation of the aspirated volume, the smallest and the largest tips come with volume marks. TP Filter Tips show high compatibility with a wide range of pipettes.

Lab Plastic

Page: 126



#### FastGene® FAS-DIGI Compact

The FAS-DIGI Compact is our new stand-alone imaging system for the detection of DNA and RNA in agarose gels, equipped with a light-sensitive 24 MPixel camera. It can be upgraded to the FAS-DIGI PRO, operated by our newly developed full control imaging software.

Gel Documentation

Page: 60



# FastGene® NanoSpec and NanoView (spectro)photometer

The NanoSpec (spectrophotometer) and NanoView (photometer) are powerful devices with a microvolume drop reader and a cuvette reader, designed for versatile applications. This includes quantitative measurements of nucleic acids, proteins, protein assays or bacterial cultures.

Lab Instruments

Page: 144



#### FastGene® Mini Dry Bath Advance

The FastGene® Mini Dry Bath Advance is a microprocessor controlled block heater. It provides excellent temperature control for various applications and can be used with exchangeable thermo blocks for different tube sizes, ranging from 0.2 ml PCR tubes to 50 ml culture tubes.

Lab Instruments

Page: 160

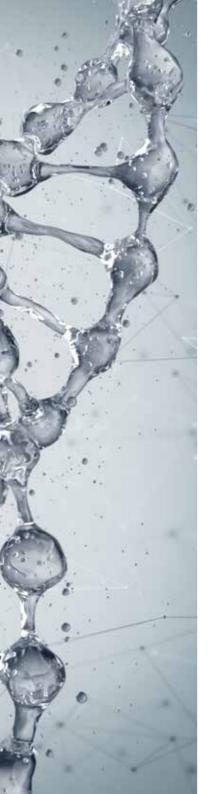


#### StemFit Basic 04 CT®

StemFit® is a culture medium for embryonic stem cells and iPS cells. This medium is recommended by the nobel prize winner Shin'ya Yamanaka and allows very reproducible growth rates under feeder-free and xeno-free conditions.

Cell Biology

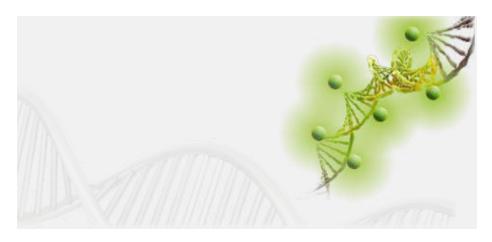
Page: 166



MIDORI <sup>stee</sup> Atra	P. 20
MIDORI <sup>Green</sup> DNA dyes	P. 24
Agarose and Agarose Tablets	P. 28
MUPID™-ONE Electrophoresis System	P. 32
DNA Marker	P. 35

#### MIDORI Green Xtra

DNA stain with the strongest signal



- Staining of DNA/RNA in agarose gels
- **Ultra-sensitive**
- Safe DNA dye
- Optimal for Blue/Green LED and Blue LED light
- Almost no background

#### MIDORI Green Xtra: The revolution

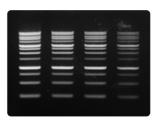
MIDORI<sup>Green</sup> Xtra is a new highly sensitive green fluorescent stain for a safe visualization of DNA and RNA in agarose gels. This DNA stain is a safe and better alternative to the traditional nucleic acid stain ethidium bromide (EtBr). Remarkably, agarose gels stained with MIDORI<sup>Green</sup> Xtra have a very low background fluorescence, which makes the identification of low amounts of DNA very easy.

#### Proven safety

MIDORI<sup>Green</sup> Xtra delivers even better DNA/RNA signals than ethidium bromide. However, this innovative DNA stain is non-carcinogenic, non-mutagenic and non-toxic and therefore not harmful for your health. Independent laboratories confirmed its safety: Both the mutagenicity test (Ames-Test) and the cytotoxicity test were negative.

#### Optimal for Blue/Green LED technology

MIDORI<sup>Green</sup> Xtra is optimized for Blue/Green and Blue LED light, leading to unbeatable fluorescence signals of DNA and RNA in agarose gels. In addition, UV-light is also suitable for the detection of nucleic acid, but less efficient than non-damaging visible light. Remarkably, MIDORI<sup>Green</sup> Xtra did not stain the agarose gel, leading to an excellent signal to noise ratio.



Ultra-high sensitivity of DNA bands detected with MIDORI<sup>Green</sup>
Xtra (dilution factor 1:25000) using with a Blue/Green LED
transilluminator.

Safe DNA stain with an unbeatable sensitivity

Ames-Test

Cytotoxicity Test

#### MIDORI Green Xtra

DNA stain with the strongest signal



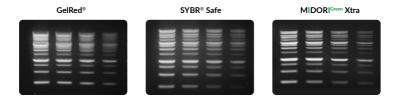


#### Simply the best

MIDORI<sup>Green</sup> Xtra leads to unbeatable fluorescence signals of nucleic acids. The direct comparison with the popular DNA dyes GelRed® and SYBR® Safe shows, that MIDORI<sup>Green</sup> Xtra reaches new levels of sensitivity: Even the detection of the smallest quantities of DNA or RNA is possible. But don't take our word for it - try them for yourself! Just contact us, and get your free sample.

# No changes in electrophoresis mobility and band distortion

The in-gel staining of agarose gels can cause a distortion of DNA bands and can result in a change of the migration pattern. However, with MIDORI<sup>Green</sup> Xtra you never have problems with distorted DNA bands and you obtain always the same migration pattern, even at different DNA concentrations. Look at the Tech Note at the next page for more information.



Comparison of the sensitivity between GelRed®, SYBR® Safe and MIDORI Green Xtra using a Blue/Green LED transilluminator. For each gel we loaded different amounts of our DNA Marker MWD1P (Page 29) (10  $\mu$ L, 7.5  $\mu$ L, 5  $\mu$ L and 2.5  $\mu$ L).

Cat. No.	Product	Content
MG10	MIDORI <sup>Green</sup> Xtra	1 ml (staining up to 25 liters of agarose)



#### Happy customers

MIDORI<sup>Green</sup> Xtra is a new and safe stain for the detection of DNA in agarose gels. This dye has been already used very successfully by several laboratories. The feedback from the scientific society is very positive. Especially with Blue/Green LED, MIDORI<sup>Green</sup> Xtra leads to fantastic results.

#### Still destroying your DNA with UV-light? Try the new Blue/Green LED technology!

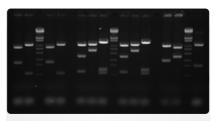
MIDORI<sup>Green</sup> Xtra (4 µl, 100 ml agarose gel)

(7 µl, 100 ml agarose gel)

SYBR® Safe

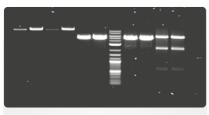
German Researcher University of Göttingen

Images were taken with a phone on a Blue/Green LED Transilluminator (FG-09).



#### Sandra Gebauer University Medical Center Göttingen

Images were taken with the FAS-V gel doc system. 2 μl MIDORI<sup>Green</sup> Xtra in 150 ml TAE buffer (1% gel).



#### German Researcher University of Hannover

Images were taken with a Blue/Green LED gel doc system. 2 µl MIDORIGreen Xtra in 100 ml liquid 1% agarose.



#### Martinsried, Germany

Images were taken with the FAS Digi. 4 µl MIDORIGreen Xtra in 100 ml 1% agarose.

#### **Customer Testimonial**

"Overwhelming results with Blue LED light. Much better than Ethidium bromide!"



#### Japanese Researcher

Jichi Medical University Department of Regenerative Medicine, Shimotsuke, Japan

\*\*\*\*

**Technical Data** 

#### Product evaluation of MIDORI Green Xtra in DNA staining

#### Purpose

Evaluate the performance of the new staining reagent MIDORI Green Xtra by using the in-gel staining method.

#### Background

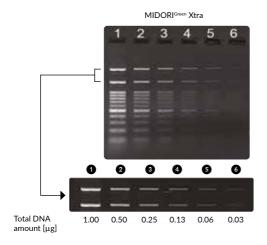
One method of staining DNA separated by gel electrophoresis is the "in-gel" staining method. For in-gel staining, electrophoresis is carried out using a gel containing nucleic acid staining reagent. Therefore, it is possible to observe the electrophoresis result without requiring DNA staining process. However, it can come to a distortion of the bands and there is a risk of causing a change in migration pattern, which should be molecular weight dependent. For this reason, in addition of being able to detect the band with high sensitivity, the reagent used for in-gel staining should precisely separate the DNA by size.

#### Experimental procedure

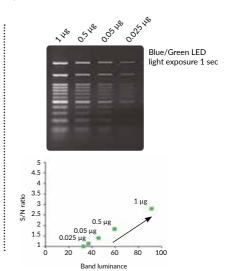
- 1) Gel preparation 2.0% TAE agarose gel with MIDORI Green Xtra (4 µl for a 100 ml gel)
- 2) DNA sample: 100 bp DNA ladder, 0.1 μg/μl (FastGene® MWD100)
- 3) Agarose gel electrophoresis: 100 V, 30 min
- 4) Gel doc system: FAS-Digi (GP-05LED) with Blue/Green LED light
- 5) Images were analyzed with Image J and the band luminance and S/N ratio were calculated for the 100 bp band

#### Result

#### 1 Influence on band formation



#### 2 Band luminance and S/N ration



#### Summary

- MIDORI<sup>Green</sup> Xtra is a reagent with no changes in electrophoretic mobility and band distortion.
- MIDORI<sup>Green</sup> Xtra is a DNA staining reagent that enables lower background and higher signal-to-noise ratio.
- → MIDORI<sup>Green</sup> Xtra has the ideal properties for the in-gel staining method with Blue/Green LEDs.



#### MIDORI Green Advance

#### Safe In-gel staining

MIDORI<sup>Green</sup> Advance is a safe alternative to the traditional nucleic acid stain ethidium bromide. It is a non-carcinogenic and less mutagenic dye for detecting dsDNA, ssDNA and RNA in agarose gels with a very high sensitivity. MIDORI<sup>Green</sup> Advance can be used with UV-light or with our innovative Blue/Green LED technology.

- Perfect staining of DNA/RNA in agarose gels
- Non-toxic, non-carcinogenic
- Safe alternative to ethidium bromide
- High fluorescence
- Optimal for UV-light



- Ames-Test
- Acute Oral Toxicity Test
- Chromosome Aberration Test
- Mouse Bone Marrow Micronucleus Test
- Latex and Nitrile Gloves Penetration Test

#### Safe alternative to ethidium bromide

MIDORI<sup>Green</sup> Advance is a non-carcinogenic dye. Optimised for a brighter signal when excited by UV-light or Blue/Green light. It has the advantages, such as being non-carcinogenic and having an excellent signal-to-noise ratio. MIDORI<sup>Green</sup> Advance shows a very high sensitivity even for small DNA fragments. The dilution factor of MIDORI<sup>Green</sup> Advance can be as high as 1:25000. Hence, 4-6 μl are enough for the staining of a 100 ml agarose gel, resulting in ~17 to 25 liters of stained agarose gels.



Comparison of sensitivity between MIDORI<sup>Green</sup> Advance and ethidium bromide using a UV-transilluminator.

#### **Proven Safety**

It is essential for a good replacement of the mutagenic DNA stain ethidium bromide to deliver strong signals. MIDORI<sup>Green</sup> Advance delivers signals with a comparable intensity. Nonetheless, the safety of the user must not be compromised. Hence, several tests were performed with MIDORI<sup>Green</sup> Advance and according to those tests, MIDORI<sup>Green</sup> Advance is safe.

#### Staining RNA with MIDORI Green Advance

#### Method:

RNA samples were separated on a 1% agarose gel stained with MIDORI Green Advance (Fig. 1) or with ethidium bromide (Fig. 2). Lane 1 and 2: 0.5  $\mu$ g of RNA. Lane 3: 0.3  $\mu$ g of RNA. Lane 4: 0.7  $\mu$ g of RNA. The separation of the RNA was performed using a 1x TBE Buffer and 100 V for 1 hour.

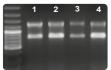


Fig. 1: RNA stained using MIDORI<sup>Green</sup> Advance. The two bands represent the major rRNA of 28S and 18S.

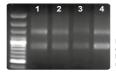


Fig. 2. RNA stained using ethidium bromide. The two bands represent the major rRNA of 28S and 18S.

#### Results/Conclusion:

MIDORI<sup>Green</sup> Advance delivered superior image quality and very distinctive bands indicating the presence of the expected 28S and 18S rRNA bands. Bands intensity correspond to the amount of RNA and the predicted bands were visible and distinctive.

 ${\sf Data\;kindly\;provided\;by\;Ms\;Kirstin\;Linsmeier,\;University\;of\;Heidelberg,\;Germany}$ 

Cat. No.	Product	Content
MG04	MIDORI <sup>Green</sup> Advance	1 ml (staining up to 25 liters of agarose)



#### In-sample staining for the strongest signal

MIDORI Green Direct stain represents a safe nucleic acid stains for visualisation of double-stranded DNA, single-stranded DNA and RNA in agarose gels. In contrast to most other non-ethidium bromide based dyes, MIDORI Green Direct is just added to your samples.

- Direct staining of DNA/RNA
- Non-toxic, non-carcinogenic
- Safe alternative to ethidium bromide
- Loading dye is included
- Very low background

#### Safety first

MIDORI<sup>Green</sup> Direct is non-carcinogenic and less mutagenic compared to ethidium bromide. Furthermore, we can state that MIDORI<sup>Green</sup> Direct is impenetrable to latex gloves and cell membranes (Fig 1.). MIDORI<sup>Green</sup> Direct is classified as non-hazardous to aquatic life, under CCR Title 22 regulation. Thus, small amounts of MIDORI<sup>Green</sup> Direct stain can be safely released into the environment.

# SYBR® Green MIDORIGneen Direct

Fig. 1: HeLa cells were incubated at 37°C with SYBR® Green I and MIDORI<sup>Green</sup> Direct. Images were taken following incubation for 30 min. SYBR® Green I entered into cells rapidly as evident from the bright green nuclear staining. However, MIDORI<sup>Green</sup> Direct was unable to cross cell membranes as shown by the lack of any fluorescence staining.

- Ames-Test
- Cytotoxicity Test
- Cell Membrane Permeability
- Hazardous Waste Screening
- Latex Gloves Penetration Test

#### Ordering information

# Cat. No. Product Content MG06 MIDORI<sup>Green</sup> Direct (with loading dye) 1 ml

#### The best signal to noise ratio

The direct staining of the DNA rather than the gel eliminates the background staining, providing a perfect signal (Fig. 2). MIDORI Green Direct was developed to work with Blue LED light and Blue/Green LED light transilluminators, but you can also use it with a regular UV transilluminator.

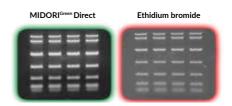


Fig 2.: MIDORI<sup>Green</sup> Direct detected by Blue/Green LED light vs. ethidium bromide detected using UV-light.

#### No extra steps necessary

MIDORI<sup>Green</sup> Direct is provided in form of a 10X sample loading dye and is only added to your samples. You do not need to add any other dyes to the gel matrix nor to the running buffers. MIDORI<sup>Green</sup> Direct contains a mixture of orange G and xylene xyanol. In the case you want to add your own loading dye, there is a loading dye free version of MIDORI<sup>Green</sup> Direct as well.

You do not need to add any other loading dye to both gel matrix and running buffer

#### Better results for downstream applications

The isolation of DNA from agarose gels enables downstream applications. It is well known that many dyes, such as ethidium bromide or even SYBR™ Green are strong enzyme inhibitors due to their intercalating properties. MIDORI<sup>Green</sup> dyes bind to the DNA backbone. This results in a much higher efficiency for downstream applications, i.e. cloning (Fig. 3), sequencing, PCR etc.

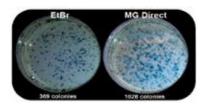


Fig. 3: DNA was isolated from agarose gels stained with ethidium brig. 3: DNA was isolated from agarose, where MIDORI<sup>com</sup> Direct was used. The isolated DNA was transformed into E. coli. The transformed bacteria were plated on selective media and incubated for 16 hours at 37°C. Hours at 30°C.



**Technical Data** 

MIDORI Green Advance: Long term storage test (3 months) of prestained gels

#### Purpose

MIDORI<sup>Green</sup> Advance was used to prepare a prestained gel. One was used "on the day of making", another one was used "after 3 months". Each gel was subjected to electrophoresis. Gel images were taken under the same conditions and were compared afterwards.

#### Method

- 1. A prestained gel was prepared:
  - 1.5% TAE agarose gel | 12.5 ml mini gel | 0.5 µl MIDORI<sup>Green</sup> Advance
- 2. The prestained gel was used for electrophoresis:
  - Condition 1: Used for electrophoresis on the day of creation
  - Condition 2: Store at 4°C, after 3 months the gel was used for electrophoresis
- 3. Electrophoresis and gel imaging conditions:
  - $\bullet$  DNA sample: Bioline Easy ladder I (Bio-33045) 5  $\mu$ l / lane Conc. (250 ng / 5  $\mu$ l)
  - Electrophoresis: SafeBlue Electrophoresis system (MBE-150 Plus) 100V 30min
  - Gel imaging: FAS-Digi (Pentax MX-1) Blue/Green LED transilluminator

#### Prestained gel storage method

Usually, when an agarose gel is refigerated and stored at 4°C, it is ideal to store it in a container containing "the same buffer solution used for gel preparation" in order to prevent drying. However, in the case of a prestained gel, in order to prevent dilution of the staining reagent, it is necessary to add the same concentration of the staining reagent to the storage buffer. Therefore, we did not use buffer for storage this time. We wrapped the gel as it was, shielded with aluminium foil, to avoid light exposure and tried a method to store it with a plastic bag with zipper.



 Wrap each gel together with tray.



All gels are covered with aluminium foil.



3. Packed in a sealable plastic bag and stored at 4 °C.

#### Result

#### On the day of creation



#### After 3 months of storage



#### Summary

The result of this study shows, that even after refigerating a gel which was stained with MIDORI<sup>Green</sup> Advance for 3 months at 4°C there was no difference in the detection of sensitivity observed and it was possible to use it for electrophoresis without problems.

#### For UV-light

#### For Blue/Green & Blue LEDs

#### **MIDORI**Green Advance Strongest signals with UV-light



Cat. No.: MG04 / Content: 1ml Sufficient to stain 25 liters of agarose

#### MIDORI Green Xtra Unbeatable with visible light



Cat. No.: MG10 / Content: 1ml Sufficient to stain 25 liters of agarose

#### MIDORIGE Direct

In-sample staining for the strongest signal



Cat. No.: MG06 / Content: 1ml Stain up to 2000 samples

In-gel & post-staining

Direct staining of DNA/RNA

# Free Sample?

Would you like to test our DNA dyes or the Agarose Tablets? No problem! Just give us a call or write us an email and get your free sample very soon.



+49 2421 554960



www.nippongenetics.eu

# **SFast Gene™** MIDORI<sup>Green</sup> Agarose Tablets



- Simple and safe gel pouring
- DNA dye is already in the tablet (MIDORI<sup>Green</sup> Xtra or Advance)
- High fluorescence
- Only water or buffer needed
- Increase your reproducibility and safe time

#### Don't waste time preparing gels!

MIDORI<sup>Green</sup> Agarose Tablets are a fast, clean solution for preparing agarose gels without any additional time-consuming steps, such as weighing or adding different components. Just add the tablet to pure cold water or buffer, heat, and pour. That's it! Once the gel hardens, it's ready for loading. Each tablet contains the perfect amount of the DNA dye MIDORI<sup>Green</sup> Xtra or Advance.

If you're tired of preparing agarose gels for your lab, this is the quickest and easiest solution to reduce effort and improve the quality of your gels.

#### Easy workflow

The fastest workflow to make agarose gels: 1. Add the tablet to pure cold water (when using the tablets with TBE or TAE) or in cold buffer (when using the tablets without buffer); 2. Dissolve the tablet by shaking your solution; 3. Heat the solution until it is clear; 4. Add the solution to your gel tray; 5. Run the gel and detect your DNA bands.

Add tablet Shake it Heat it up Add to gel tray Run the gel



# *ூ Fast போட* MIDORI<sup>Green</sup> Agarose Tablets

#### Choose your tablet

Depending on your needs NIPPON Genetics EUROPE provides 5 different MIDORI<sup>Green</sup> Agarose Tablets: First choose the DNA dye: We offer tablets with MIDORI<sup>Green</sup> Xtra and MIDORI<sup>Green</sup> Advance. You are using TBE or TAE as a buffer? Then use MIDORI<sup>Green</sup> Tablets with TBE or TAE. You want to use your own buffer or a different running buffer? No problem, just use the MIDORI<sup>Green</sup> Agarose Tablets without buffer.



MIDORI<sup>Green</sup> Advance TBE Agarose Tablets 75 tablets (0.5 g agarose each) Cat. No.: AG09



MIDORI<sup>Green</sup> Advance TAE Agarose Tablets 75 tablets (0.32 g agarose each) Cat. No.: AG10



MIDORI<sup>Green</sup> Advance Agarose Tablets (without buffer) 100 tablets (0.5 g agarose each) Cat. No.: AG11



MIDORI<sup>Green</sup> Xtra
TAE Agarose Tablets
100 tablets (0.5 g agarose each)
Cat. No.: AG13



MIDORI<sup>Green</sup> Xtra Agarose Tablets (without buffer) 100 tablets (0.32 g agarose each) Cat. No.: AG12

#### The perfect gel concentration

With the MIDORI<sup>Green</sup> Agarose Tablets (Xtra or Advance) you can get your desired gel percentage: Just use the instructions from the tables. Please note that the MIDORI<sup>Green</sup> Agarose Tablets with TAE buffer contain less agarose.

## Gel concentration for AG09 / AG11 and AG13 (Tablets with TBE and without buffer)

	1		
Gel	1 Tablet	2 Tablets	
1%	50 ml H <sub>2</sub> O	100 ml H <sub>2</sub> O	
1.5%	$33~\mathrm{ml~H_2O}$	67 ml H <sub>2</sub> O	
2%	25 ml H <sub>2</sub> O	50 ml H <sub>2</sub> O	

## Gel concentration for AG10 and AG12 (Tablets with TAE buffer)

Gel	1 Tablet	2 Tablets
1%	32.5 ml H <sub>2</sub> O	65 ml H <sub>2</sub> O
1.5%	21.5 ml H <sub>2</sub> O	43 ml H <sub>2</sub> O
2%	16.25 ml H <sub>2</sub> O	32.5 ml H <sub>2</sub> O

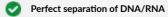
Cat. No.	Product	Content
AG09	MIDORI <sup>Green</sup> Advance TBE Agarose Tablets	75 Tablets (0.5 g Agarose each)
AG10	MIDORI <sup>Green</sup> Advance TAE Agarose Tablets	75 Tablets (0.65 g Agarose each)
AG11	MIDORI <sup>Creen</sup> Advance Agarose Tablets (without buffer)	100 Tablets (0.5 g Agarose each)
AG12	MIDORI <sup>Green</sup> Xtra Agarose Tablets (without buffer)	100 Tablets (0.5 g Agarose each)
AG13	MIDORI <sup>Green</sup> Xtra TAE Agarose Tablets	100 Tablets (0.325 g Agarose each)

# **SFast Gene**™ Agarose

#### Molecular grade agarose

The FastGene® Agarose was developed for an accurate separation of DNA fragments, such as PCR products and plasmid DNA, as well as RNA. The very high quality allows all experiments for molecular biology. The purity of the agarose leads to an excellent transparency and a low background. This is especially important to obtain sharp and well defined DNA and/or RNA bands with the highest sensitivity in the low molecular weight range.





Sharp and well defined DNA bands

Electroendosmosis (EEO): 0.14-0.16

Concentrations from 0.75 - 2%

#### Agarose tablets - no weighing required

With the FastGene® Agarose Tablets you can create agarose gels without time consuming weighing. Just add one tablet to 50 ml of gel running buffer and heat — the result is a 1% agarose gel. It's that simple. With our Agarose Tablets there's no powder to weigh and no mess to make!



The tablets can be dissolved in your favorite running buffer.

#### Quality agarose for small products

The detection of small bands is only possible with high quality agarose. Two gels were prepared using the FastGene® Agarose and a low quality agarose from competitor C. The ladder was stained with MIDORI<sup>Green</sup> Direct and separation of the bands was done using the Mupid™-ONE electrophoresis system (MU2, next page).





Detection of small bands using high quality FastGene® Agarose and Competitor C's Agarose. All seven bands from the ladder are visible when using FastGene® Agarose. When comparing the green box to the red box, Competitor C's Agarose does not show the lowest three bands.

#### Choose a suitable agarose concentration

High gel strength allows you to use gel concentrations from 0.75% - 2%. Blotting experiments will run perfectly, and separation efficiency up to 23 kb is guaranteed. Every batch of our agarose is tested with different-sized DNA fragments, and the background fluorescence is measured with ethidium bromide or non-toxic stains to assure the cleanest signals. But don't take our word for it — try them for yourself!

Cat. No.	Product	Content
AG01	FastGene® Agarose	100 g
AG02	FastGene® Agarose	500 g
AG05-100	FastGene® Agarose Tablets	100 Tablets (0.5 g Agarose each)

# சிச்சி Agarose Gel Band Cutter

#### Safe time for cutting DNA bands

This easy-to-use tool will facilitate your daily work. Now you can excise your DNA bands easily without risking contamination or scratching the glass surface of the transilluminator. The FastGene® Agarose Gel Band Cutter is a ready-to-use and disposable tool for cutting agarose gel bands. This affordable tool simplifies fragment purification and eliminates wasted effort using razor blades. The size of the excised gel band will always be 6 mm x 3 mm, and you can collect multiple bands in a single FastGene® Cutter — making large-scale purifications that much easier.



FastGene® Agarose Gel Band Cutter is the best way to excise DNA bands from an agarose gel.



Easily excise DNA bands



No razor blades necessary



Safe time

#### Ordering information

Cat. No	э.	Product	Content
FG-83	)	FastGene® Agarose Gel Band Cutter	50 Units

#### **Customer Testimonial**

"We are very happy using the FastGene® Gel Band Cutter and have successfully implemented it in our practical course. In the past, our students had issues cutting out the correct band without adding too much unnecessary agarose when using a scalpel and a tweezer. This is important since during the next step the same amount of extraction buffer has to be added to the agarose material. This problem was solved by using this product. We have tested similar products but they could not convince us."



#### Zeynep Weninger

Laboratory Biochemistry - Faculty of applied chemistry Nürnberg Institute of Technology Georg Simon Ohm, Germany



#### **Stock Solutions**

# All you need for a perfect agarose gel electrophoresis

Running and sample buffers as ready solutions. The highly concentrated stock solution is industrially produced and tested and therefore a secure and convenient alternative for selfmade agarose buffers.

Cat. No.	Product	Content
ID1521	50X TAE Buffer	500 ml
ID1531	10X TBE Buffer	500 ml
ID1654	6x Nucleic Acid Loading Buffer	10 ml





**TAE Buffer** 



6x DNA Loading Buffer

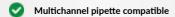


# Mupid<sup>™</sup>-ONE Electrophoresis System











Gel casting set included

#### Advanced DNA separation

The Mupid™-One Electrophoresis system is one of the most convenient DNA separation systems on the market. It has many novel features including a separated power supply, a simple buffer drainage system, support for multi-channel pipettes, and seven output voltage settings (18, 25, 35, 50, 70, 100 and 135 V) as well as a timer function for delivering the perfect run every time.

#### What makes this system so safe?

For prevention of an electric shock, the system is running only if both parts (chamber and power controller) are connected and if the lid is closed! With an open lid the main power can not be switched on.

#### What makes this system so popular?

The Mupid<sup>TM</sup>-One gel trays are heat resistant: By using a novel polymeric material (PPHOX), gel solution up to  $100^{\circ}$ C can be poured into the tray without turning milky or britle. The clean up of the used gel trays can be performed with boiling water.



The Mupid $^{\text{TM}}$ -One is a CE labeled electrophresis system for agarose gels.

#### What makes this system so easy-to-use?

The power controller is easy to use. Seven conventional voltages (18, 25, 35, 50, 70, 100 and 135 V) are available. The peak voltage is constant (140 V) and output level changes through pulse control. A timer with an alarm function is included. All parameters of the last run are memorised and automatically saved. The power controller can be disconnected easily. Therefore, the chamber is safe and easy to clean up.

# Mupid<sup>™</sup>-ONE Electrophoresis System

#### Everything you need for a perfect gel!

The Mupid<sup>™</sup>-One comes with the Gel casting set standard (Cat.No. ON-MS). This casting set includes 4 combs, which can be used from both sides (13 wells and 26 wells). Furthermore, this set comes with different gel trays: Two gel trays small (S) for the preparation of mini gels and one gel tray large (L) for larger gels. The optional gel casting set GM-HR includes two large combs and two different gel trays: Four gel trays mini and two gel trays small (S).



#### Gel casting set standard (ON-MS), included with the Mupid™-One.

#### All accessories are included

SPECIFICATIONS		
Compact design	<b>~</b>	Overall dimensions (H x D x W): 5.9 x 16.2 x 18.3 cm Bath volume: 270 - 320 ml
Integrated power supply	<b>~</b>	Input voltage: AC100 V - 240 V, 50-60 Hz Output voltage: 8 V, 25 V, 35 V, 50 V, 70 V, 100 V and 135 V
Memory function	<b>~</b>	Automatic memory function from the last use
Safety lid	<b>~</b>	Without the lid, main power can not be operated
Multi-channel pipette compatible	<b>~</b>	The included combs are multi-channel pipette compatible
Optimal gel tray size	<b>~</b>	Small gel tray: 130 mm (B) x 16.5 mm (H) x 59.5 mm (L) Large gel tray: 130 mm (B) x 24 mm (H) x 122 mm (L)
Optimal comb size	<b>~</b>	Number of wells: 13 or 26 Spacing size: 9 mm (13 wells)

#### Ordering information

Cat. No.	Product	Content
MU2	Mupid™-One	Mupid $^{TM}$ -One electrophoresis system with $1x$ gel chamber, $1x$ power controller, $1x$ gel casting set, $4x$ combs, $2x$ gel trays $S$ , $1x$ gel tray $L$
MU4	Mupid™-One LED Illuminator	Mupid™-One LED Illuminator (see next page)

#### Accessories for the Mupid<sup>™</sup>-One

Cat. No.	Product	Content
ON-MS	Gel casting set standard	1x Mupid <sup>™</sup> -One gel casting set, 4x combs, 2x gel trays S, 1x gel tray L
GM-HR	Gel casting set large	1x Mupid <sup>™</sup> -One gel casting set large, $2x$ large combs, $4x$ gel trays mini, $2x$ gel trays L
ON-GL	Large gel trays	2 gel trays L
ON-GS	Small gel trays	4 gel trays S
ON-SD	Gel casting stand standard	1 gel casting stand standard
AC-C1	Gel combs	2 combs for the Mupid $^{\!TM}\!$ -One electrophoresis system

# Mupid<sup>™</sup>-ONE LED Illuminator



#### Watch the DNA run

The MUPID™-One LED Illuminator allows the visualization and detection of DNA fragments during the run. The illuminator substitutes the MUPID lid and includes an orange coloured filter to allow you easily check the results without wearing goggles.



Real time electrophoresis



Safe Blue LED light



Add-on for the Mupid<sup>™</sup>-One electrophoresis chamber

#### Blue LED light for a safe detection of DNA

The MUPID™-One LED Illuminator produces blue light with an emission peak at 470 nm, effective for the excitation of safe nucleic acid stains such as MIDORI<sup>Green</sup>Xtra and SYBR® dyes. The separated DNA is not damaged by the LED light, because it contains no short-wave UV-light.





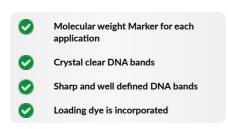
SPECIFICATIONS		
Safe Blue LED light	<b>~</b>	Blue LED light for the safe detection of green DNA dyes (wavelength of 470 nm)
Compact design	<b>~</b>	Dimensions (H x D x W): $5.1 \times 16.6 \times 17$ cm Viewing area: $15 \times 6$ cm
Compatible	<b>~</b>	MUPID™-One, Mupid™ exU and Mupid™ ACE

Cat. No.	Product	Content
MU4	Mupid™-One LED Illuminator	Mupid™-One LED Illuminator with black gel trays

# **Grast Gene™** DNA Marker

#### For each application the right ladder

The FastGene® DNA ladders were developed for different applications: The MWD50 was designed for the most accurate discrimination of small PCR products, from 50 bp up to 1,500 bp. The MWD100 is the perfect ladder for everyday use. The ladder with 12 fragments starts at 100 bp, therefore being suitable even for small qPCR products, and goes up to 3,000 bp so that the sizes of small plasmids and big PCR products can be determined. For very large products and plasmids, NIPPON Genetics EUROPE offers the MWD1P. The ladder starts at 100 bp and goes up to 10,000 bp.



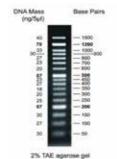
#### Stable even at room temperature

The DNA Ladders MWD100 and MWD1P are extremly stable. The stability tests show that the ladders are stable for at least 12 months at 25°C. For long term storage, store at 4°C to -20°C.

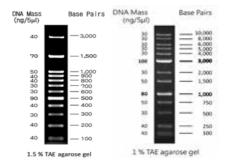
#### Dyes for easy tracking

All our ladders have tracking dyes and loading dyes included, so that the movement of the DNA can be tracked and the optimal stopping point can be determined.

#### MWD50



MWD1P



MWD100

Best quality/price ratio

SPECIFICATIONS			
Cat. No.	MWD50	MWD100	MWD1P
Description	50 bp Ladder	100 bp Ladder	1 kb Ladder
Range / bp	50 - 1,500	100 - 3,000	100 - 10,000
Number of bands	17	12	13
Reference bands	3 (200, 500, 1,200)	2 (500 & 1,500)	2 (1,000 & 3,000)
Loading dye	Orange G	Orange G & Xylene cyanol FF	Bromphenol blue
Content	56 μg in 500 μl	50 μg in 500 μl	50 μg in 500 μl
Recommended load	5 ul		

Cat. No.	Product	Content
MWD50	FastGene® 50 bp Standard DNA Marker	500 μΙ
MWD100	FastGene® 100 bp Standard DNA Marker	500 μΙ
MWD1P	FastGene® 1 kb Standard DNA Marker Plus	500 μΙ

# **PROTEIN ELECTROPHORESIS**

	80	00	O	
.0.	J	0	0	
0 (	)°	0.0	0"	
°°°	۵			c
0	0	0	c	

PAGE Protein System	P. 38
Western Blot System	P. 40
Precast PAGE Gels	P. 42
Q-Stain Protein Staining Solution	P. 44
Chemi ECL Kit and Running Buffer	P. 45
Protoin Markor	D 44

# **SFast Gene™** PAGE Protein System

All you need for PAGE protein analysis



- Complete set for PAGE protein analysis
- Hand-cast gel set included (1 mm gels)
- Compatible with different pre-cast gels
- 0.75 mm / 1.5 mm hand-cast sets available
- Includes glass plate holder and tube holder

## Powerful protein analysis via PAGE

PAGE stands for Polyacrylamide Gel Electrophoresis and describes an analytical method in biochemistry for the separation of differently sized protein mixtures in an electric field. During PAGE, proteins migrate through a gel matrix in response to an applied electric field. Smaller proteins travel faster through the gel than larger proteins, leading to a size dependent separation.

The most popular form of PAGE is SDS-PAGE. The detergent sodium dodecyl sulfate (SDS) is added for PAGE sample preparation and is also part of the buffer composition. The treatment with the harsh detergent leads to full protein denaturation and unfolding. SDS binds to hydrophobic parts of the unfolded protein and masks the intrinsic charge of the protein with its own negative charge. As a consequence, SDS-protein complexes migration is predominantly dependent on the size of the protein, allowing an estimation of its molecular weight.

## **Broad Gel compatibility**

The FastGene® PAGE Protein System is compatible with a wide range of pre-cast gels (e.g. FastGene® gels, Bio-Rad TGX™ gels or ThermoFisher mini gels). It also runs hand-cast gels included in the PAGE Protein System Set (for casting 1 mm thick gels). Also available are FastGene® hand-cast gel sets for gel preparation with 0.75 mm or 1.5 mm gel thickness.



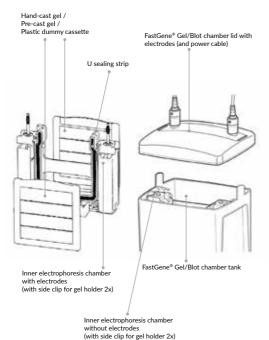
The FastGene® PAGE Protein System contains all the necessary equipment to conveniently perform protein electrophoresis.

# **SFast Gene™** PAGE Protein System

All you need for PAGE protein analysis

## Complete set for PAGE protein separation

The FastGene® PAGE Protein System contains all the necessary components you need for PAGE protein analysis. Three types of sealing strips are included in the set, allowing compatibility with different gel types and gel sizes. The electrophoresis tank can hold a maximum of 4 gels simultanously for electrophoretic protein separation. The hand-cast set comes with everything needed for convenient gel casting.



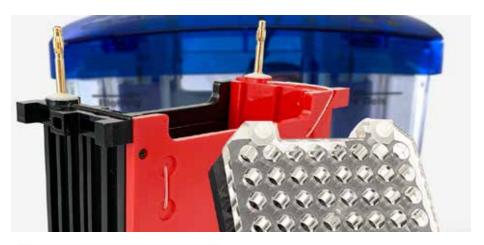
## PAGE Protein System (PG01) content

F	astGene® PAGE Protein System (PG01)			Qty
	Inner electrophoresis chamber with electrodes (with side clip for gel holder 2x)			1
	Inner electrophoresis chamber without electrodes (with side clip for gel holder 2x)			1
	U sealing strip long for 10 x 10 cm gels (e.g. ThermoFisher $^{\text{TM}}$ mini gels)	5		4
tem	U sealing strip short for Bio-Rad TGX™ (10 x 8 cm)	gels '		4
PAGE System	U sealing strip FastGene® (10 x 8 cm)			4
PAG	Plastic dummy cassette short (10 x 8 cm)			1
	Plastic dummy cassette long (10 x 10 cm)			
	FastGene® Gel/Blot chamber lid with electrodes (and power cable)			1
FastGene® Gel/Blot chamber tank (PG05)				1
	Gel shovel			5
(SS)	Comb 1 mm 10 wells	nm)	m)	5
ickne	Comb 1 mm 10 Wells  Comb 1 mm 15 wells  Comb 1 mm 15 wells		.5 m	5
gel th	Glass spacer long 1 mm	0) 7(	PG03 (1.5 mm)	5
E E	Glass spacer long 1 mm  Glass plates short (PG04)		PGC	10
et (1	Comb 1 mm 15 wells Comb 1 mm 15 wells Glass spacer long 1 mm Glass spates short (PG04) Gel casting base Gel casting clip (PG06) Sealing gaskets (PG07) Glass plate holder Tube holder			4
ing se	ຜ Gel casting clip (PG06)			4
-cast	Sealing gaskets (PG07)			5
hand	Glass plate holder			1
Ge	Tube holder			1

Cat. No.	Product	Content
PG01	FastGene® PAGE Protein System	Complete protein PAGE set (see table above for content)
PG02	FastGene® Comb Set 075	Gel hand-casting set (for 0.75 mm gel thickness) (see table above for content)
PG03	FastGene® Comb Set 150	Gel hand-casting set (for 1.5 mm gel thickness) (see table above for content)
PG04	FastGene® Glass plates short	Short flat glass plates for hand-cast gels (10 glass plates)
PG05	FastGene® Gel/Blot chamber tank	Large gel electrophoresis buffer tank (1 chamber tank)
PG06	FastGene® Gel casting clip	Frame for holding hand-cast gels (4 pieces)
PG07	FastGene® Sealing gaskets	Sealing gaskets for gel casting (5 pieces)

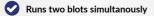
## \$ Fast சோட் Western Blot System

Efficient wet-transfer device



Enicient and renable protein transfer		Efficient and reliable protein transfer
---------------------------------------	--	---





## Biochemical protein analysis via blotting

Blotting of proteins is a powerful biochemical method for the detection of proteins. Protein bands that were separated by size after PAGE are transferred and immobilized on a carrier membrane. The firm binding of the proteins to the membrane allows a subsequent protein detection by choosing from a variety of staining or immunological methods.

## Wet-transfer Western Blot system

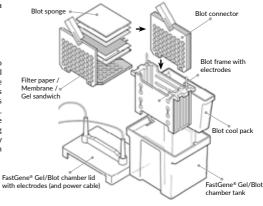
The FastGene® Western Blot System is an efficient device to perform the first Western Blot protein transfer step from gel to membrane via a wet electrophoretic transfer technique. The Western Blot Systems contains all the necessary components for reliable protein transfer and comes with detailed guidelines to successfully perform transfer and Western Blot analysis. The system includes a cool pack, which is installed during the run. The cool pack absobs generated running heat, avoiding the possibility of power loss. Gel and membrane can be easily sandwiched in the blot connector, which is assembled via an easy-to-use lock mechanism.

## Ordering information

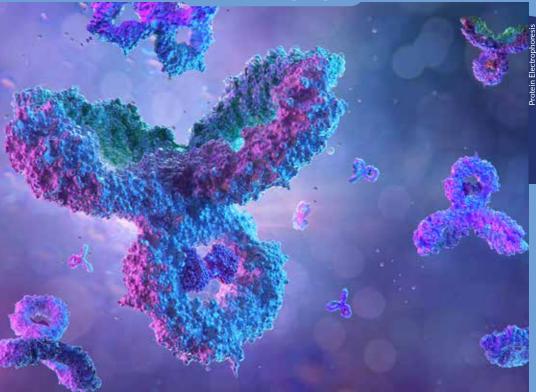
Cat. No.	Product	Content
PG08	FastGene® Western Blot System	Complete Western Blot set (see table above for content)
PG09	FastGene® Western Blot components	Components for Blotting System (see table above for content)

## Western Blot System (PG08) content

FastGene® Western Blot System (PG08)		Qty
Blot frame with electrodes		1
Blot connector	60	2
Blot sponge	PG	5
Blot cool pack		2
FastGene® Gel/Blot chamber lid with electrodes (and power cable)		1
FastGene® Gel/Blot chamber tank (PG05)		1



**Product Highlight** 



# Do you have questions about our PAGE or blotting systems?

Please do not hesitate to ask us! We offer product demonstrations online or in your lab. Just arrange an appointment with us!



+49 2421 554960

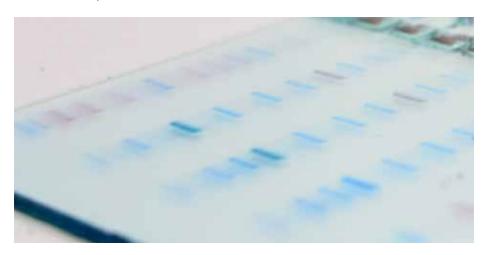


info@nippongenetics.de
 info@nippongenetics.de

www.nippongenetics.eu

## **SFast** Gene™ Precast PAGE Gels

Get the best separation



- 8 x 10 cm PAGE Gels
- Homogenous and gradient gels
- No special buffers required
- Superior protein band resolution and stability
- Long shelf live

## Get the best separation

Pouring hand-cast gels for protein separation can be time consuming and prone to error. FastGene® Precast Protein Gels are the perfect replacement, facilitating lab work immensely. Due to the proprietary gel casting method, which is more uniform than any self-cast gel, the FastGene® Precast Protein Gels have the advantage of being more consistent, having therefore a much higher reproducibility.

## Homogenous or gradient PAGE gels

FastGene® Precast Protein Gels are available in a variety of homogenous and gradient gels. They are formulated for denaturing as well as native gel electrophoresis – depending only on the used running buffer. Our gels are compatible with MOPS or MES buffer.



Each box of our FastGene® Precast Protein Gels comes with 10 gels, a cassette opener and spacers. You also need buffer? No problem, just order our MOPS buffer packs.

Load up to 60 µl sample on each lane

## Superior running performance

FastGene® Precast Protein Gels are casted at a neutral pH environment. The hydrolysis of polyacrylamide is reduced, resulting in an increased gel stability and superior band resolution. Further advantages are optimised running performance and a larger loading volume (up to 60 µl). The extra large wells also prevent a lane-to-lane overflow and a higher transfer efficiency.

## **Grast** Gene™ Precast PAGE Gels

Get the best separation



## New quality standards

The FastGene® Precast Protein Gels have a revolutionary highperformance. The unique buffer formulation that maintains a low operating pH during the electrophoresis eliminates the "smilies" and poor resolution of self made gels and many competitor Precast gels.

# SFast Gene" Competitor B

Direct comparison of a FastGene® Precast Protein Gel (12%) with a very common competitor gel manufacturer.

## Free sample

You would like to test our Precast Protein Gels? No problem! All gels are available as a sample with all necessary components for protein electrophoresis, including MOPS buffer! Just contact us, and get your free sample.

#### Compatibility

The gels are compatible with different common protein electrophoresis gel tanks like from Bio-Rad and our FastGene® Electrophoresis Unit.

Manufacturer	Product
All manufacturer	s are using 8 x 10 cm gels
NGE	FastGene® PAGE Protein System
BioRad	Mini PROTEAN II & 3 Mini PROTEAN Tetra System
Hoefer	SE 250 Mighty Small II SE260 Mighty Small II Deluxe

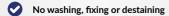
Cat. No.	Product	Content
PG-S012	FastGene® PAGE Gel 8 x 10 cm - 12%	10 gels
PG-S412	FastGene® PAGE Gel 8 x 10 cm - 4-12%	10 gels
PG-S420	FastGene® PAGE Gel 8 x 10 cm - 4-20%	10 gels
PG-S816	FastGene® PAGE Gel 8 x 10 cm - 8-16%	10 gels
PG-MOPS10	FastGene® MOPS Buffer Pouches	10 Pouches for 1 L each

## \$ Fast:போட் Q-Stain Protein Stain

#### Like Coomassie Blue only simpler

The FastGene® Q-Stain is a single-step, modified Coomassie Blue protein gel stain for polyacrylamide gels. This protein staining solution eliminates the need to fix, wash or destain your protein gel. Just run your protein gel, add the FastGene® Q-Stain, and watch your bands appear in several seconds. The FastGene® Q-Stain does not stain the polyacrylamide gel. The result is a crystal-clear background with clearly visible protein bands. Unlike many other stains, the FastGene® Q-Stain is a water-based product, free of Methanol and Acetic acid.





High sensitivity - 10 ng bands detectable

Free of methanol and acetic acid

No oversaturation

## Ideal for mass spectrometry

The FastGene® Q-Stain Protein Stain is 100% compatible with mass spectrometry. Just follow the procedure below and analyse your protein:

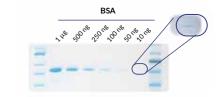
- Incubate the excised protein band in 1 ml 30% EtOH or 30% acetone for 30 min at room temperature
- 2. Repeat step 1 until the stain is removed
- 3. Continue with a typical mass spectrometry protocol

Never Wash or Destain again!



## One-step protein staining in 10 minutes

The special formulation of the FastGene® Q-Stain enables a very quick stain procedure of protein gels. The protein bands will be visible in less than 10 minutes. Very low amounts of proteins (down to 10 ng) can be detected by longer staining. It is impossible to over-saturate proteins with the FastGene® Q-Stain, so longer incubation times have no harmful effects. Save time by using Q-Stain for a safe and efficient detection of proteins in polyacrylamide gels.



Detection of 10 ng of protein after 30 minutes incubation. For a better visualization the 10 ng protein band is shown with a stronger contrast.

#### Staining a protein gel was never so easy

The entire staining procedure can be completed in about 10 minutes (for typical protein amounts). Just remove the gel after the electrophoresis, add the stain, wait for 10 minutes and watch your protein bands become visible.



Cat. No.	Product	Content
FG-QS1	FastGene® Q-Stain	1 liter

## *் F் துட்பு பாட்* Western ECL Kit



# FastGene Western ECL Kit Competitor M Competitor B Competitor T

Comparison between the FastGene® Western ECL Kit with 3 competitor products. All ECL kits were used under the same experimental workflow. An IL-6 fusion protein was detected using a primary antibody against IL-6 from mouse and a secondary anti-mouse peroxide (POD) antibody. 2 seconds exposure for the Western Blox.

## Chemiluminescent Western Blot Detection

The FastGene® Western ECL Kit is a luminol-based enhanced chemiluminescent substrate and sensitive with conducting immunoblots with horseradish peroxidase (HRP) – conjugated secondary antibodies. Due to the excellent substrate sensitivity and long signal duration, the FastGene® Western ECL Kit enables the detection of antigens with a very low concentration. Furthermore, its long chemiluminescent signal duration makes both digital and film-based imaging possible without any loss of the signal. Appropriate primary and secondary antibody dilutions are suggested to attain optimal signal intensity and duration.



Workflow using the FastGene® Western ECL Kit: Mix the luminol solution and peroxide solution in a 1:1 ratio, and thoroughly agitate the chemiluminescent substrate solution for preparing the 0.1 ml of solution per cm² of membrane. Place the membrane with the protein side up and remove the membrane from the chemiluminescent substrate solution. Take an image the membrane with a digital imager.

### Ordering information

Cat. No.	Product	Content
FG-CH01	FastGene® Western ECL Kit	50 ml each Buffer

## **Running Buffer**

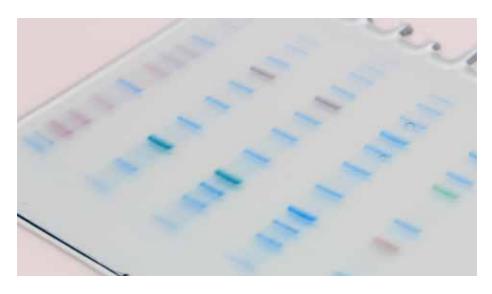
## All you need for perfect protein electrophoresis

The running buffer is available as a ready solution or as a measured powder in aluminium foil to make 1 liter of buffer. This eliminates the tedious weighing of SDS and other components.



Cat. No.	Product	Content
PG-MOPS10	FastGene® MOPS Buffer Pouches	10 Pouches for 1 L each
ID1501	Running Buffer Tris-Glycine-SDS	10x 500 ml

## *GFastGene* Protein Marker



- Huge size range (6.5 270 kDa)
- Ready-to-use
- Sharp bands
- Reference bands
- Quality tested

## One, two or three colours?

You have the freedom of choice: Five different protein Markers with different colours and distinct size ranges. All of our Protein Markers are supplied in a loading buffer for a direct loading on gels. The FastGene® Protein Markers have sharp bands with an excellent accuracy. They are designed for monitoring protein separation during SDS-polyacrylamide gel electrophoresis, verification of Western transfer efficiencies (PVDF, nylon, or nitrocellulose membranes) and for approximating the size of proteins.

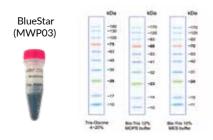
Supplied in a loading buffer for direct loading on gels

Cat. No.	Product	Content
MWP03	BlueStar Prestained Protein Marker (500 µl)	Sufficient for 100 mini gels or 50 large gels
MWP04	BlueStar PLUS Prestained Protein Marker (500 $\mu$ l)	Sufficient for 100 mini gels or 50 large gels
MWP05	JustBlue Prestained Protein Marker (500 μl)	Sufficient for 100 mini gels or 50 large gels
MWP06	BlueEasy Prestained Protein Marker (500 μl)	Sufficient for 100 mini gels or 50 large gels
MWP07	Unstained Protein Marker (500 μl)	Sufficient for 100 mini gels or 50 large gels

## *் F் த்து பிசாட*் Protein Marker

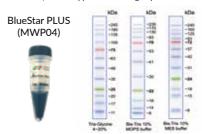
#### BlueStar Prestained Protein Ladder

The BlueStar Prestained Protein Ladder is a three colour protein standard with 10 prestained proteins covering a wide range of molecular weights from 10 to 180 kDa.



#### BlueStar PLUS Prestained Protein Ladder

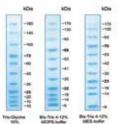
The BlueStar PLUS is a three colour protein standard with 12 prestained proteins covering a wide range of molecular weights from 10 to 245 kDa. The Prestained Protein Ladder is designed for monitoring protein separation during SDS-polyacrylamide gel electrophoresis, verification of Western transfer efficiency on membranes (PVDF, nylon, or nitrocellulose) and for approximate sizing of proteins.



#### JustBlue Prestained Protein Ladder

The JustBlue Protein Marker is a reasonable priced alternative to the BlueStar Marker. Eleven proteins are coupled with a blue chromophor. 3-5  $\mu l$  per loading for clear visualization during electrophoresis on 10-15 well gels. We recommend 2-3  $\mu l$  per well for Western transfer. Two reference bands (25 and 72 kDA) are enhanced in intensity when separated on SDS-PAGE. Stable for up to 3 months at 4°C, long term storage at -20 °C.





## BlueEasy Prestained Protein Ladder

The BlueEasy Prestained Protein Ladder (MWP06) is a three colour protein standard with 10 prestained proteins. It has the largest range of molecular weights from 6.5 to 270 kDa. The BlueEasy Prestained Protein Ladder is designed for monitoring protein separation during SDS-polyacrylamide gel electrophoresis, verification of Western transfer efficiency on membranes (PVDF, nylon or nitrocellulose) and for approximating the size of proteins.



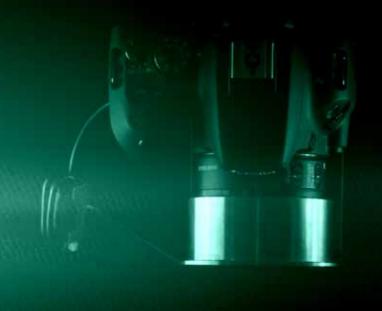


#### Unstained Protein Ladder

The Unstained Protein Ladder is a mixture of 12 unstained recombinant proteins covering a wide range of molecular weights from 10 to 200 kDa. Please note that this ladder is not stained. The bands will become visible only once you stain your gel with a generic method, such as Coomassie blue or Q-stain.







# GEL DOCUMENTATION

Introduction of the Blue/Green LED Technology	P. 50
Gel Documentation Systems - Overview	P. 54
Compact - FAS-Nano / FAS-BG LED BOX / FAS-DIGI Compact	P. 56
High-End - FAS-DIGI PRO / FAS-V Imaging System	P. 62
Transilluminators	P. 70



## The Blue/Green LED Technology



#### The Danger of UV-Light

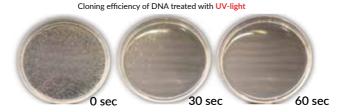
Detection of nucleic acids is mainly performed using light in the UV-range. However, DNA is able to absorb light in the UV-spectrum. This property leads to DNA modifications and DNA degradation when exposed to UV-light. It damages sample DNA and is also dangerous for the user.

## Blue/Green LED - The Revolution

Instead of using a single wavelength, the Blue/Green LED technology uses a combination of wavelengths in a spectrum of light from 470 nm to 520 nm. This Blue/Green light is able to excite all common red and green DNA dyes with a very high intensity. This intensity can be achieved by the accumulated energy absorption of the dyes.

## Comparing the influence of UV-light to Blue/Green LED light on the cloning efficiency

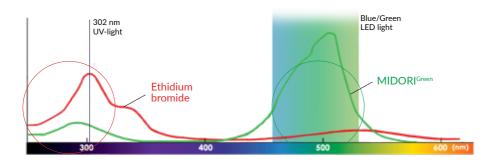


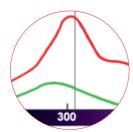






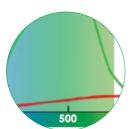
## The Revolution





## **UV-Light:** Good detection, insecure signal

UV-light transilluminators use just a single wavelength for the visualization of DNA. Red and green DNA dyes, like ethidium bromide or the MIDDRI of the Susually have a good absorption in the UV-light spectrum. This results in DNA bands with a sufficient intensity. However, UV-light is dangerous for the user and for the sample DNA. Just 30 sec of UV-light exposure significantly reduces the cloning efficiency and has consequences for further downstream applications. For this reason, the visualization of DNA with UV-light is not the state-of-the-art method anymore.

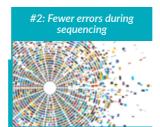


## Blue/Green light: Safe detection of all Red and Green DNA dyes

In contrast to UV-light, Blue/Green LED technology uses a spectrum of light between 470 nm and 520 nm. This light is not harmful for the DNA or for the user. Even ethidium bromide or other red DNA dyes with a low absorption in this spectral area show DNA band intensity comparable to UV-light. The reason for that is the accumulated energy absorption of the DNA in the Blue/Green spectrum. Green DNA dyes show very high absorption intensity in the Blue/ Green light spectrum, leading to DNA bands with superb intensity.

## Try Blue/Green light - Your Benefits:









## State-of-the-Art Method









## Blue/Green LED Technology conquers the world

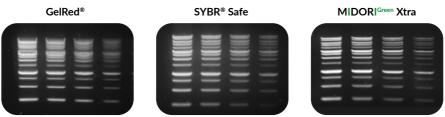
Gel documentation systems with Blue/Green LED technology are already used in more than 1000 laboratories around the world. No damage of DNA, for much better results of downstream applications (ligation, cloning, sequencing). Together with all common DNA dyes, Blue/Green LED light leads to fantastic intensities of DNA/RNA bands in agarose gels. Gel documentation with Blue/Green LED light is therefore the new state-of-the-art method for the visualization of nucleic acids.

1000+ Instruments around the world with Blue/Green LED light



## Blue/Green LED Technology





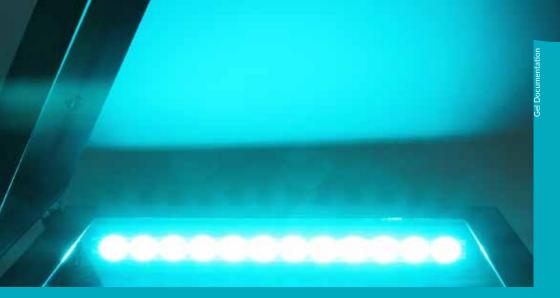
Comparison of the sensitivity between GelRed®, SYBR® Safe and MIDORI<sup>Green</sup>Xtra using the FAS-DIGI Pro. In combination with MIDORI<sup>Green</sup>Xtra, the FAS-DIGI Pro leads to an unbeatable image quality of agarose gels. Each gel was loaded with different amounts of our DNA marker MWD1P (Page 35).

# **⑤** Fast Gene™ Gel Documentation Systems

Find the right gel doc system for your needs

	FAS-Nano (GP-06LED)	FAS-BG LED BOX (GP-04LED)	FAS-DIGI Compact (GP-08LED)	FAS-DIGI PRO (GP-07LED)	FAS-V (GP-FAS-V)
Safe Blue/Green LED light	•	•	•	•	•
Detection of Green DNA dyes	•	•	•	•	•
Detection of Red DNA dyes	•	•	•	•	•
White Light Imaging	0	•	0	•	•
High Resolution Camera	0	0	•	•	•
Parfocal Lens	0	0	0	0	•
Software included	0	•	0	•	•
Networkable	0	0	0	•	•
Stand-Alone System	0	•	•	•	•
Large illuminated Area	0	0	•	•	•
Quantification of DNA and RNA	0	0	0	•	•
CE Certification	•	•	•	•	•

<sup>\*</sup> Operation also possible without Computer



# Talk to the experts and enjoy a product demonstration

Finding the right gel doc system or transilluminator can be difficult. We can help you! Just arrange an appointment with us and enjoy a product demonstration.



+49 2421 554960



www.nippongenetics.eu

## Fast Gene™ FAS-Nano



- Gel documentation with safe Blue/Green LED light
- Detection of red and green DNA dyes
- Compact footprint and light weight
- Image acquisition with smart device
- Amber shield included

## Take gel images with your phone

The FastGene® FAS-Nano LED system is the most compact gel illumination system on the market. Ideally suited for tight spaces on a bench-top, the system operates both as an illuminator and, if equipped with a smartphone or tablet, a documentation system that captures an image of your gel.

## The first portable gel imaging system!

## The perfect personal illuminator

Its very small footprint and light weight make the FastGene® FAS-Nano a perfect personal illuminator. An array of Blue/ Green LEDs positioned around the periphery of the glass plate provide excitation light for both red and green DNA dyes without UV-light damage.



Combine your smartphone or tablet with the FastGene® FAS-Nano and turn the illuminator to a gel documentation system. The recording of the gel image is easily done by taking a picture.

Cat. No.	Product	Content
GP-06LED	FastGene® FAS-Nano	Illuminator, amber shield, dark hood, adaptor for mobile camera.

## Fast Gene™ FAS-Nano



SPECIFICATIONS		
Safe Blue/Green LED light	~	Blue/Green light spectrum from 470 nm to 520 nm No risk of damaging DNA or harming your skin and eyes
Smartphone lens	~	Ultrawide angle lens
Compact footprint	<b>~</b>	Dimensions (H x D x W): 12.8 x 21.6 x 16.8 cm   Illuminated area: 10 x 10 cm Weight: 1.2 kg
Accessory	~	Ultrawide angle lens

## **GFastGene**™ FAS-BG LED BOX



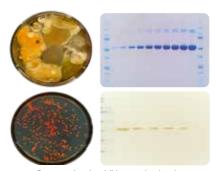
- Gel documentation with safe Blue/Green LED light
- Very compact footprint
- Detection of red and green DNA dyes
- Occumentation of protein gels, membranes and petri dishes
- High resolution camera with 9 MPixel

## Compact imaging System with Blue/Green LED

The FastGene® FAS-BG LED BOX comes with the advantages of the Blue/Green LED technology combined with a compact footprint. All red and green DNA dyes are easily detectable with this system.

## One imaging system - multiple applications

The Blue/Green LED technology permits the detection of DNA with highest sensitivity and without harming your eyes, skin or your sample. With the white LED array you can image protein gels stained with coomassie or silver staining. The white LED epi-illumination allows the documentation of opaque surfaces such as petri dishes or membranes.



Documentation of petri dishes, protein gels and Western Blot images.

Cat. No.	Product	Content
GP-04LED	FastGene® FAS-BG LED BOX	LED imaging box with a high resolution CMOS camera (9 MPixel)

## **GrastGene**™ FAS-BG LED BOX

#### **Customer Testimonial**

"We have been using the FAS-BG LED BOX already for four months as the main device for the detection of DNA bands in agarose gels. The FAS-BG LED BOX has a compact design, is easy to use and produces images with a high quality. In order to make the USB memory accessible in the network, we use a USB switch that connects the stick to a neighboring PC. We are very satisfied with the Blue/Green LED technology and have replaced our entire UV devices to protect our samples and colleagues."



#### **Thorben Detering**

Institute of Food Chemistry, Leibniz University Hannover, Germany



#### Easy connection to a monitor or PC

The FastGene® FAS-BG LED BOX can be directly connected to an external monitor (via VGA) or to a thermal printer (via USB). By using a USB-switch you are also able to transfer your gel pictures from the FAS-BG LED BOX to a PC, where you can share them via network.



The large touch screen display an inbuild software allows easy navigation and image capturing.

## Excise your DNA fragments the easy way

With the FastGene® FAS-BG LED BOX and our MIDORI<sup>Green</sup> dyes it is simple to excise your DNA fragments from gels. There is no need for protective eyewear — just switch on the Blue/Green LEDs and excise your DNA fragment. You can also obtain perfect signals if you are still using ethidium bromide or other red DNA dyes.



The FastGene® FAS-BG LED BOX comes with Epi White LED light for petri dishes and membranes and white back light for easy detection of protein gels.

SPECIFICATIONS				
Safe Blue/Green LED light	<b>~</b>	Blue/Green light spectrum from 470 nm to 520 nm No risk of damaging DNA or harming your skin and eyes		
Easy image capture	<b>~</b>	CMOS 9 MPixel camera   Exposure time: 11 exposure scales Image types: JPEG, TIFF, BMP   Image Storage: USB 2.0		
2 White light sources	<b>~</b>	Epi white light for petri dishes and membranes White back light for protein gels		
Compact footprint	~	Dimensions (H x D x W): 23 x 25.4 x 23 cm   Illuminated area: 16 x 11.5 cm Weight: 3.2 kg		
Connectable to a monitor or PC	~	Direct connection to an external monitor (via VGA) or to a thermal printer (via USB), easy connection to a PC using a USB-switch		

## Fast Gene™ FAS-DIGI Compact



- Gel documentation with safe Blue/Green LED light
- Scientific grade camera
- Detection of red and green DNA dyes
- Large illuminated area
- Compact and easy to use stand alone system

The FAS-DIGI Compact comes with a huge Blue/ Green LED transillumianor, giving you a large illuminated are for any gel size

## Strong Blue/Green LED transilluminator - for all common DNA dyes

The FastGene® FAS-DIGI Compact is our smaller sister model of the FAS-DIGI PRO. Its compact design is perfect for labs that need a powerful system but have limited bench-top space. The strong Blue/Green LED transilluminator was designed for ultra safe and ultra sensitive DNA/RNA detection. The FastGene® FAS-DIGI Compact can be used for all common red and green DNA dyes, without the harmful properties of UV-light. You are able to excite ethidium bromide, GelRed®, MIDORI Green or SYBR® Green with very high intensity. The inbuilt Amber Filter viewing window allows you to look at your illuminated gel and easily cut out bands for further applications.



## Ethidium Bromide



#### MIDORI Green Advance



#### MIDORI Green Direct



# Fast Gene™ FAS-DIGI Compact





The FAS-DIGI Compact is equipped with a high resolution scientific



You can upgrade the FAS-DIGI Compact to the FAS-DIGI PRO by simply replacing the dark box

## High resolution scientific grade camera

Take gel pictures with the highest image quality using a 24 MPixel Canon EOS DSLR camera with a huge APS-C CMOS sensor. With the 3x zoom (focal length of 18 mm to 55 mm) you can easily enlarge the area of interest. The high resolution allows you to extract small gel image areas in best quality. Exposure times up to 30 sec enable you to detect even faint bands.

## Simply upgrade to the PRO model

The FAS-DIGI Compact is the perfect gel documentation device for limited lab space. However, if you decide at one point to upgrade your system to the even more powerful FAS-DIGI PRO, the switch will be easily performed. All you need to do is replace the FAS-DIGI Compact dark box with the FAS-DIGI PRO dark box, that comes with an intuitive and easy to use imaging software.

SPECIFICATIONS		
Safe Blue/Green LED light	~	Blue/Green light spectrum from 470 nm to 520 nm No risk of damaging DNA or harming your skin and eyes
Scientific grade camera	<b>~</b>	24 MPixel (Resolution: 6000 x 4000), APS-C sensor, F/4-5.6 aperture, 18-55 mm zoom lens, 0.00025 to 30 seconds exposure time
High-quality material	<b>~</b>	Coated aluminium metal
Huge transilluminator	<b>~</b>	View area: 26 x 21 cm
Viewing window	<b>~</b>	Amber filter window
Very Compact design	<b>~</b>	Dimensions (H x D x W): 50 x 35 x 32.5 cm, Weight: 7.4 kg
Upgradable	<b>~</b>	Upgradable with the FAS-DIGI PRO dark box
Integrated power supply	<b>~</b>	100-240 V~, 50/60 Hz

Cat. No.	Product	Content
GP-08LED	FastGene® FAS-DIGI Compact	LED imaging box with amber filter window, B/G transilluminator, high resolution camera

## Fast Gene™ FAS-DIGI PRO



- Gel documentation with safe Blue/Green LED light
- Scientific grade camera
- Image software with comprehensive features for image acquisition
- Fully networkable
- Compatible with all common DNA dyes

#### Touch the revolution

The FastGene® FAS-DIGI PRO is our powerful imaging system for the detection of DNA and RNA in agarose gels. Equipped with a light-sensitive 24 MPixel camera, the FAS-DIGI PRO is controlled completely by an innovative imaging software. With the live view mode, all camera settings, the exposure time, the lens' aperture, and digital zoom are displayed in real-time. The FAS-DIGI PRO is a fully networkable gel doc system allowing simple transfer of images when connected to a PC.



## Blue/Green LED light for a safe detection of DNA and RNA

The FastGene® FAS-DIGI PRO is composed of a strong transilluminator equipped with the unique Blue/Green LED technology. The LEDs emit light at a wavelength from 470 nm to 520 nm without damaging nucleic acids. The Blue/Green LED light enables the detection of all common green dyes, such as MIDORI<sup>Green</sup> or SYBR® Green, yellow dyes e.g. SYBR® Safe and red dyes, e.g. ethidium bromide or GelRed®.

## Still destroying your DNA with UV-light? Try the new Blue/Green LED light!



The FAS-DIGI PRO is composed of a huge transilluminator with an illuminated area of  $26 \times 21 \, \mathrm{cm}$ . The dark hood can be easily removed, which allows a very easy excision of DNA bands. Otherwise, just use the amber shield which is magnetically attached to the box.

## **GFast Gene™** FAS-DIGI PRO

## Camera for high quality agarose gel

The documentation of agarose gels with highest image quality can be obtained using a 24 MPixel camera with an immense APS-C CMOS sensor. The sensor produces no visible noise from ISO 100 all the way up to ISO 1600. Furthermore, the 24 MPixel allows the detection of lowest light signals in agarose gels. The exposure time of the sensor can be set from 1/4000 sec up to 30 sec. The 3x zoom (focal length of 18 mm to 55 mm) allows a perfect enlargement of the area of interest.



The camera is directly connected to the power supply adapter of the FAS-DIGI PRO. Replacing batteries is not necessary.

## Huge and strong transilluminator

The imaging area of the transilluminator has a size of  $26 \times 21$  cm, which allows the imaging of multiple agarose gels of various sizes. Additionally, the FAS-DIGI PRO comes with an amber shield, which can be attached with magnets inside the box. This makes cutting out DNA bands very easy.



Use the included amber shield to cut out DNA bands

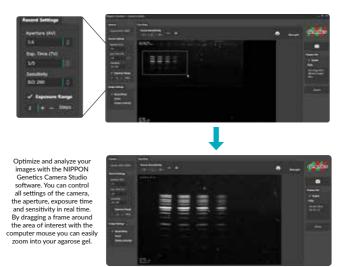
SPECIFICATIONS		
Safe Blue/Green LED light	~	Blue/Green light spectrum from 470 nm to 520 nm No risk of damaging DNA or harming your skin and eyes
Scientific grade camera	~	24 MPixel (Resolution: 6000 x 4000), APS-C sensor, F/4-5.6 aperture, 18-55 mm zoom lens, 0.00025 to 30 seconds exposure time
High-quality material	<b>~</b>	Coated aluminium metal
White light source	~	White LED transilluminator: Documentation of protein gels
Imaging software	~	NIPPON Genetics Camera Studio, Windows 10, Saved image format TIFF and JPEG
Huge transilluminator	<b>~</b>	View area: 26 x 21 cm
Integrated power supply	<b>~</b>	100-240 V~, 50/60 Hz
Compact design	~	Dimensions (H x D x W): 57 x 35 x 32.5 cm, Weight: 14 kg

Cat. No.	Product	Content
GP-07LED	FastGene® FAS-DIGI PRO	LED imaging box, B/G transilluminator, imaging software, high resolution camera, White LED transilluminator, Magnetic amber filter shield, Magnetic amber filter for the camera lens

## **⑤**Fast Gene™ FAS-DIGI PRO - The Software

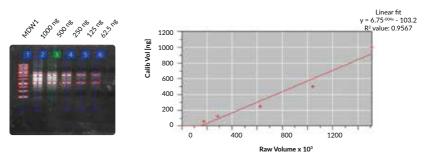
## Easy-to-use control imaging software

The FastGene® FAS-DIGI PRO comes with the intuitive NIPPON Genetics Camera Studio software. With this software you can control all necessary parameters of the camera to analyze and optimize any gel image. These four settings will provide the highest quality images your lab has ever seen for DNA gels: aperture, exposure time, sensitivity and focus. Mouse-driven control makes image optimization a click away! Images are saved as TIFF and JPEG format, and can be printed directly by a printer connected to your PC.



#### Quantification of nucleic acids with the FAS-DIGI PRO

For the quantification of DNA or RNA in agarose gels, it is necessary that the light signals received by the camera are proportional to the DNA/RNA concentration. Usually, researchers are using a gel doc system with an integrated CCD camera for the quantification of their DNA/RNA signals. However, modern scientific grade CMOS cameras are so accurate that they can be used for the quantification of nucleic acids, too. The price tag of a CMOS camera is much lower than of a CCD camera. The FAS-DIGI PRO uses very modern CMOS cameras of the highest quality. Generated pictures can be quantified by using the Total LAB 1D software (not part of the NIPPON Genetics Camera Studio software).



Quantification of RNA with the FAS-DIGI PRO using the Total LAB 1D software (Cat. No.: GP-QS1). A 1% agarose gel was stained with 3 µl of MIDORI Far No. 100 ml of agarose. The gel was loaded with MWD1P (5 µl), and human total RNA (Agilent Cat No.: 750500) in different concentrations (1000, 500, 250, 125, 62.5 ng). The CMOS sensor of the Canon scientific grade camera is able to generate pictures, which can be quantified by using the Total LAB 1D software. MIDORI Form Xtra shows a low background and produces crystal clear bands. This stain is detected with a linear signal to noise ratio and is therefore suitable for quantification.

## **SET SET SET OF THE Accessories FAS-DIGI PRO - The Accessories**

#### Amber Filter Shield

The FastGene® FAS-DIGI PRO comes with an amber filter shield that can be used to excise DNA bands without having to wear amber filter goggles. The amber filter shield is positioned inside the box and held in place with magnets. When not in use, the amber filter shield can be stored on the inside of the door.



The amber filter shield enables a quick and easy excision of DNA.



Use the amber shield to cut out DNA bands

## White light LED Illuminator

The documentation of protein gels and Petri dishes is possible with the white LED illuminator used inside the FastGene® FAS-DIGI PRO. The white light LED illuminator is powered by battery. Hence, the procedure is very simple: (1) Remove the amber filter shield, (2) remove the amber filter in front of the camera lens (3) place the illuminator in the central area and take your image.



Place the amber filter shield inside the door.



Remove the amber filter in front of the lens.



Position the white LED illuminator in the central area and adjust the recording settings in the NIPPON Genetics Camera Studio.

# Fast Gene™ FAS-V Imaging System



- Safe Blue/Green LED light -No damaging of your DNA
- Stand-alone system
- White LED light for the documentation of protein gels
- Easy-to-use software
- Best image quality

### Stand alone documentation

The FastGene® FAS-V is our most advanced imaging system, working with the innovative Blue/Green LED excitation light technology for the detection of DNA/RNA. This imaging platform combines a powerful CCD camera, brilliant touchscreen display and the superior technology of ultra-bright Blue/Green LEDs. The illumination hits the sweet spot for exciting common red and green DNA dyes such as EtBr and MIDORI<sup>Green</sup>. You can always expect at least equivalent results as compared to UV-light transilluminators, but without the risk of damaging DNA or harming your skin and eyes. Lower energy photons from Blue/Green LEDs will not crosslink or damage DNA unlike short wave UV-light.

#### **Customer Testimonial**

"We decided to purchase the FAS-V gel documentation system with its CCD sensor, because we were very interested in the integrated Blue/Green LED technology. This light protects the skin and eyes of the user and is also safe for the examined samples. [...] The FAS-V has no disadvantages compared to conventional UV-based gel documentation systems. Even at low concentration it is possible to visualize DNA bands. Furthermore, with the integrated white light it is possible to document finished Western blots. Working with the FAS-V is simpler and faster than with our previous device. Because of the simultaneously recording of 6 images with different exposure values you always get very good images of your gel in a very short time. We are very satisfied with the FAS-V and can recommend the device in good conscience.



#### German Researcher

Institute of Technical Chemistry, Leibniz University Hanover, Germany

\*\*\*\*

# Fast Gene™ FAS-V Imaging System



SPECIFICATIONS		
Safe Blue/Green LED light	~	Blue/Green light spectrum from 470 nm to 520 nm No risk of damaging DNA or harming your skin and eyes
CCD camera	~	CCD camera (1600 x 1200 - UXGA)   Exposure time: 0.001 to 30 sec Sensitivity: 0.13 Lux   Aperture: f/1.2   Focal distance: 12.5 - 75 mm   6x zoom
2 White light sources	~	White LED room light: Documentation of membranes and petri dishes White LED transilluminator: Documentation of protein gels
Huge transilluminator	~	Illuminated area: 26 x 21 cm   Dimensions (H x D x W): 78.5 x 40 x 38.2 cm
Networkable computer	~	Integrated computer, which can be directly connected to a network
User friendly software	<b>~</b>	Intuitive imaging software for image acquisition Controlled by a 10.4" touchsreen
Easy image capture	<b>~</b>	Save images in JPEG, TIFF, PNG or BMP format   Image storage: 16 GB SSD
Integrated power supply	<b>~</b>	100-240 V~, 50/60 Hz, 2 A

# **SFast Gene™** FAS-V Imaging System

## Very sensitive camera (CCD technology)

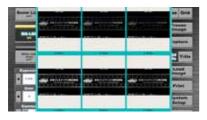
The FastGene® FAS-V has a big CCD-Sensor with a diameter of 1/1.8". The pixel size of  $4.4~\mu m$  allows the detection of lowest light signals. Images can be recorded in the common TIFF and JPEG as well as in BMP and PNG format. The files can be stored on the 16~GB internal SSD storage or on a USB-stick

#### Very bright parfocal lens - no more focusing

The lens of the FastGene® FAS-V is parfocal, as most microscopic lenses, and prefocused on the imaging area, enabling you to zoom in to the area of interest without readjusting the focus. The huge aperture of f/1.2 enables the maximum transmission of light to the sensor. The aperture can be regulated steplessly until complete closure, making it your decision which aperture delivers the best image.

The 6x zoom, with a focal length from 12.5 mm to 75 mm, allows a perfect enlargement of the area of interest. Setting the lens at 12.5 mm allows the imaging of the complete illuminated area, while zooming into 75 mm will eliminate any unnecessary area.





Easy to control imaging software: Activate or deactivate all three light sources. Or change the image capturing settings. With the multiple exposure mode you can capture 6 images simultaneously with different exposure times.



No more focusing with the parfocal lens.

## Touchscreen operation

The FastGene® FAS-V is easily controlled by a gorgeous, colour 10.4" touchscreen display. All three light sources can be activated and deactivated by the touchscreen. Additionally, the exposure time and gain can be easily adjusted.

The FAS-V system will take up to six pictures simultaneously, using different exposure times. The user can then view and choose which one to use.

A captured image can be edited on site. The image editor starts when an image is loaded from the internal or an external storage. The image can be mirrored horizontally as well as vertically or turned by 90° clockwise or anticlockwise. The contrast of the image can be adjusted and the unimportant parts of the image can be removed using the cropping function.

## White LED transilluminator

The detection of protein bands is performed with the bright white LED transilluminator plate. The white light LED transilluminator has a huge working area of 26 cm x 21 cm, enabling the documentation of very large protein gels or of multiple protein gels at once. The additional white LED room light allows the positioning of gels and to capture images of membranes and petri dishes.

Cat. No.	Product	Content
GP-FAS-V	FastGene® FAS-V	Imaging unit with built-in computer, B/G LED slide table, high resolution CCD camera, 2 MPixel, Lens 12.5 - 75 mm, lens hood, amber filter, software, stylus and manual

# Fast Gene™ FAS-V Imaging System

## Blue/Green transilluminator for perfect gel imaging

The FastGene® FAS-V has the biggest transilluminator with the Blue/Green LED technology. The imaging area of 26 cm x 21 cm has a superb uniform illumination. This light uniformity is due to two perfectly positioned fields of 12 LED arrays at each side with excitation of 470 nm - 520 nm. This enables the detection of green dyes, such as MIDORI<sup>Green</sup> or SYBR™ Green, yellow dyes, e.g. SYBR™ Safe and red dyes, e.g. ethidium bromide or GelRed™. The special amber filter, optimised for green, yellow and red DNA dyes leads to the strongest signal with the lowest background.



# **⑤** Fast Gene™ Imaging System Accessories



## FastGene® Amber Goggles

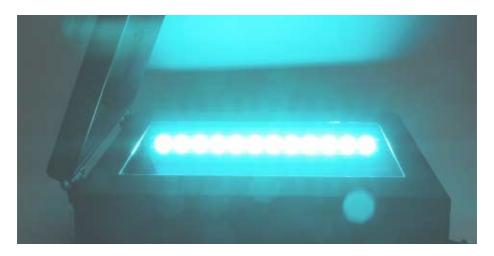
Glasses for watching agarose gels under Blue and Blue/Green Light.

## **TotalLab Quantification Software**

Image analysis tool for the quantification of your DNA, RNA and protein samples.

Cat. No.	Product	Content	
GPG	Amber Goggle	1x Amber Goggle	
GP-QS1	TotalLab Quantification Software	1- License of the Quantification Software	

## **SFast Gene™** Transilluminators



- **✓** Very high life expectancy
- Amber Filter / UV-filter included
- Safe Blue/Green LED or Blue LEDlight
- Compatible with MIDORI<sup>Green</sup> dyes
- Easy excision of DNA bands

## Blue/Green LED Transilluminators

The Blue/Green LED Transilluminators enable safe detection of DNA and RNA in agarose gels They emit light from 470 nm to 520 nm and are compatible with all common green and red DNA dyes, such as MIDORI<sup>Green</sup> and ethidium bromide.

## Blue LED Transilluminators

Our Blue LED Transilluminators also enable a safe and damage-free detection of nucleic acids. They produce light with a narrow emission peak at ~470 nm, effective for the visualisation of green DNA stains such as MIDORI<sup>Green</sup> and SYBR®. Blue LEDs are not compatible for the detection of red DNA dyes.

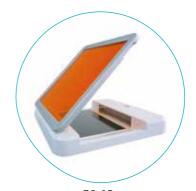
Cat. No.	FG-09	FG-11	FG-12	FG-05	FG-06	FG-300
Light source	Blue/Green LED	Blue/Green LED	Blue LED (470 nm) White light LED	Blue LED (470 nm)	Blue LED (470 nm)	UV-light (302 nm)
Compatible DNA dyes	Green and Red dyes	Green and Red dyes	Green dyes & Proteins	Green dyes	Green dyes	Red and Green dyes
Imaging area	21 cm x 26 cm	n.a.	18 cm x 12 cm	12 cm x 7 cm	20 cm x 16 cm	26 cm x 21 cm
Dimensions (H x D x W)	13 x 33 x 32 cm	2.5 x 19 x 3.9 cm	30 x 18.5 x 22 cm	3 x 21 x 21 cm	8 x 28 x 34 cm	8 x 28 x 34 cm
Weight	6.3 kg	0.17 kg	2.4 kg	2.1 kg	3 kg	4.3 kg
Power	AC adapter, 2 A	AC adapter, 18 V / 1 A	AC adapter 24 V, 1 A	24 V, 1.67 A	24 V, 1.67 A	48 W
Filter	Amber filter (~520 nm)	Amber filter (~520 nm)	Amber filter (~520 nm)	Amber filter (~520 nm)	Amber filter (~520 nm)	UV-blocking shield

# **ਓ** Fast போட் Transilluminators

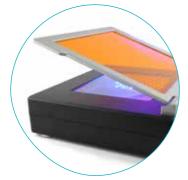








FG-05
FastGene® Blue LED Illuminator



FG-06
FastGene® Blue LED Transilluminator



**FG-300** FastGene® UV Transilluminator

# \$ Fast சோட் Blue/Green LED Transilluminator XL



## Say goodbye to UV-light

The FastGene® Blue/Green LED Transilluminator XL is a Blue/ Green LED-based transilluminator for a safe detection of DNA in agarose gels. The Blue/Green LEDs emit light from 470 nm to 520 nm, enabling the excitation of all common green and red DNA dyes, such as the MIDORI<sup>Green</sup> dyes and ethidium bromide.



No UV-light and no DNA degradation



Huge illuminated area



Excitation of all common green and red DNA dves

## Get your DNA the easy way

With the FastGene® Blue/Green LED Transilluminator XL it becomes extremely simple to cut your DNA fragment out of gels. You don't need to wear protective eyewear, or worry about DNA degredation. It is the 21st century way of working with DNA.

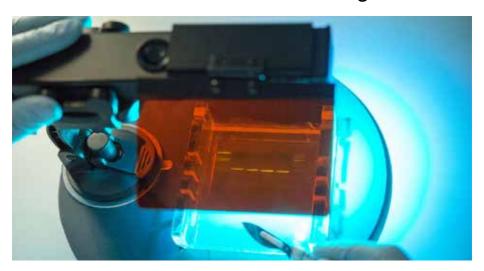




SPECIFICATIONS		
Huge illuminated area	~	Dimensions (H x D x W): 13 x 33 x 32 cm   Illuminated area: 26 x 21 cm Weight: 6.3 kg
Safe Blue/Green LED light	<b>~</b>	Spectrum of light with Blue/Green light from 470 nm to 520 nm No risk of damaging DNA or harming your skin and eyes
Amber filter included	~	Amber shield for a clear detection of DNA/RNA bands

Cat. No.	Product
FG-09	FastGene® Blue/Green LED Transilluminator XL

# & Fast Gene™ Blue/Green LED Flashlight



#### The "new" indispensable lab tool

Visualise your gel with the new Blue/Green LED flashlight. It is equipped with a Blue/Green LED light source for safe detection of all red and green DNA dyes. This novel Blue/Green Flashlight has all the benefits you would expect from using LED light. It is harmless to your skin, eyes and DNA samples, but provides signal intensity previously only seen with research quality UV transilluminators.



Portable transilluminator



Safe Blue/Green LED light



Detection of fluorescent proteins in living organisms





Detection of fluorescent proteins in plants and animals.

#### **Ordering Information**

Cat. No.	Product
FG-11	FastGene® Blue/Green LED Flashlight (stand & cutting board are included)

#### Stand and cutting board are included

The Blue/Green LED Flashlight comes with a stand, cutting board and an attached amber filter shield so you can view and cut out your DNA bands. Also, you can easily take images of your gel with a smart device. Whether you need to visualize your DNA in a gel or are confirming GFP expression post transfection, this convenient yet powerful illuminator will become an invaluable lab too!



#### Detect DNA everywhere

The Blue/Green LED flashlight was originally designed to visualize DNA in agarose gels. However, the flashlight can also be used to detect fluorescent protein expression (e.g. GFP, YFP) in living plants, animals and bacteria. Furthermore, the relative expression level of fluorescent proteins can also be estimated by the brightness of the fluorescence.

### **⑤Fast Gene™** Blue/White LED Tab



#### The "coolest" way to detect DNA

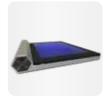
The FastGene® Blue/White LED Tab is a portable easy-touse transilluminator in tablet format. It is equipped with Blue LEDs, which produce light with a narrow emission peak at 470 nm. This wavelength leads to an effective visualization of green nucleic acid stains such as MIDORI<sup>Green</sup> and SYBR® dyes without destroying your DNA.

### Ideal for protein gels

White LEDs are also included in the FastGene® Blue/White LED Tab. This allows easy documentation of protein gels. A white plate is used to diffuse the white light, creating a homogeneous illumination. The illuminated area has the same size as the blue LED light.

- No damage of your DNA
- Easy detection of green DNA stains
- Portable and easy handling
- Occumentation of protein gels





The FastGene® Blue/White LED Tab comes with a large illuminated area and contains a Blue LED and a White LED light source.

Cat. No.	Product
FG-12	FastGene® Blue/White LED Tab

### **⑤Fast Gene™** Blue/White LED Tab

#### Brightness according to your needs

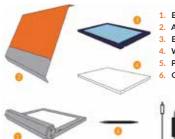
You can adjust the brightness of the FastGene® Blue/White LED Tab in both modes, the Blue LED mode and the White LED mode. Just press the brightness control button and choose one of the three possible brightness levels.

#### Best performance with MIDORI Green dyes

The FastGene® Blue/White LED Tab is ideal for the detection and documentation of green DNA dyes. Get fantastic DNA signals with the MIDORI<sup>Green</sup> dyes.



Adjust the brightness to your needs with 3 brightness levels.



- 1. Blue/White LED Tab
- 2. Amber shield
- 3. Blue uniform plate
- 4. White uniform plate
- 5. Power cord
- 6. Gel cutting knife





MIDORI<sup>Green</sup> Direct





DNA stained with MIDORI<sup>Green</sup> Advance and MIDORI<sup>Green</sup> Direct detected with the Blue/White LED tab.



SPECIFICATIONS		
Tablet format	~	Dimensions (H x D x W): 30 x 18.5 x 22 cm   Illuminated area: 18 x 21 cm Weight: 2.4 kg
Safe Blue LED light	<b>~</b>	Blue light with a wavelength of 470 nm No risk of damaging DNA or harming your skin and eyes
White light source	~	White LEDs for the documentation of protein gels
3 brightness levels	<b>~</b>	Adjust the brightness to your needs with 3 different brightness levels

# \$ Fast சோட் Blue LED (Trans-)Illuminators

#### A better alternative to UV-light

The FastGene® LED Illuminator and the FastGene® LED Transilluminator are using Blue LEDs. These LEDs produce light with a narrow emission peak centered at 470 nm, effective for the visualisation of green nucleic acid stains such as MIDORI<sup>creen</sup> and SYBR® dyes.

#### Safe for your DNA and your health

The biggest advantage for using these LED instruments is the fact that the light does not affect skin and eyes and most importantly, it does not damage your sample DNA. This is especially important if the excised DNA fragment will be used for cloning experiments. All instruments come with an amber filter, which allows the examination of separated DNA without any goggles.







The FastGene® LED Illuminator (Cat. No. FG-05).





The FastGene® Blue LED Transilluminator (Cat. No. FG-06)

#### **Ordering Information**

Cat. No.	Product
FG-05	FastGene® Blue LED Illuminator
FG-06	FastGene® Blue LED Transilluminator



#### High quality UV-light table

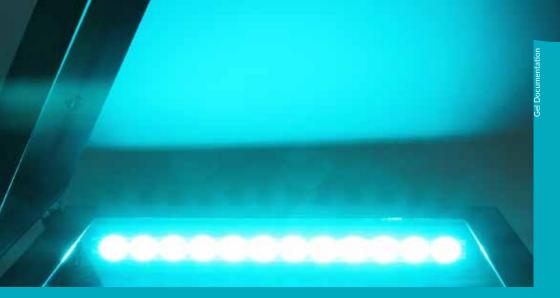
The achievable sensitivity of DNA detection, stained with ethidium bromide, is strongly dependent on the quality of the UV lamps and filter material. High quality UV tables show almost no visible light. The quality of the UV-light source and of the filter can be easily tested: UV-light is invisible to the human eye. If the position of the UV lamps is easily detected, then the quality of the light bulbs and filter are inferior. The FastGene® UV Transilluminator passes this test without a problem. Furthermore, the FastGene® UV Transilluminator includes a specifically manufactured filter system and shows a very effective protection against harmful UV radiation.

#### UV filter with 0.01% UV transmission!



The FastGene® UV Transilluminator.

Cat. No.	Product
FG-300	FastGene® UV Transilluminator



# Talk to the experts and enjoy a product demonstration

Finding the right gel doc system or transilluminator can be difficult. We can help you! Just arrange an appointment with us and enjoy a product demonstration.



+49 2421 554960



www.nippongenetics.eu



# NUCLEIC ACID PURIFICATION

RNA Purification Kits	P. 80
Viral RNA/DNA Kits	P. 88
DNA Kits - Plasmid Mini Kit and Gel/PCR Extraction Kit	P. 90
Dye Terminator Removal Kit	P. 94
Magna Stands - Magnetic Separation	P. 96

### \$Fast Gene™ RNA Kits

New level of RNA purity



- Fast procedure delivering high-quality RNA in minutes
- Consistent RNA yields
- Ready-to-use RNA for any downstream application
- Basic Kit for an easy and fast purification, even of fatty tissue
- Premium Kit for ultrapure & concentrated RNA free of genomic DNA

#### RNA of the highest quality

The FastGene® RNA Kits deliver RNA of the highest grade. The quality of RNA is determined by the RNA integrity number (RIN). According to the manufacturer's instructions, the RIN gives an idea of the integrity of the RNA. High quality RNA will give a RIN above 8, with 10 being the maximum value. The FastGene® RNA Basic and Premium Kits purify RNA to a grade comparable to market leaders. Therefore, the purified RNA is in an ideal state for downstream applications, such as reverse transcription.

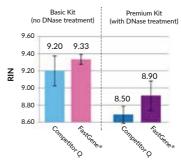
#### New level of RNA purity

The FastGene® RNA Premium Kit was developed for rapid, efficient, and clean purification of RNA from tissues and cells for challenging applications such as next-generation sequencing (NGS). The silica membrane based technology does not use phenols.

#### Very quick procedure

The FastGene® RNA Basic Kit has a very quick and easy procedure, optimal for a high number of samples. Additionally, a special protocol for large inputs was developed. The buffer necessary for large inputs can be purchased separately without the need of buying a whole kit.

#### Quality (RIN Score)



RNA quality determination using an Agilent Bioanalyzer. The FastGene® RNA Kits repeatedly deliver high quality RNA.

### \$Fast Gene™ RNA Kits

New level of RNA purity



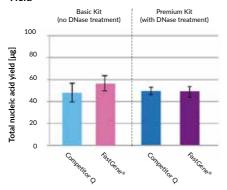
#### Large yields

Obtaining a good yield from the RNA purification is essential. The FastGene® RNA Basic and Premium Kits deliver very large total yields, therefore enabling mutliple analyses of a single RNA purification. When compared to the market leaders, the FastGene® RNA Basic kit delivers a higher yield showing the optimised purification procedure.

#### Basic or premium?

The FastGene® RNA Kits come in two different versions. The FastGene® RNA Basic Kit is ideal for purification of RNA where small amounts of copurified DNA can be neglected. Whereas the FastGene® RNA Premium Kit ensures the complete elimination of genomic DNA.

#### Yield



The FastGene® RNA Kits deliver very high yield.



Choose the Basic Kit when small amounts of copurified DNA are not a problem for you.



Choose the Premium Kit for an optimal DNA removal for very pure RNA.

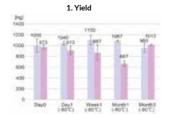
#### RNA stability of samples stored in RL-lysis buffer (3 months)

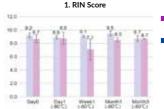
#### Background

Ideally, the extraction of the RNA and the subsequent analysis should be carried out as quickly as possible, since the RNA degrades rapidly. Depending on the timing of an experiment, it can be necessary to store the RNA before extraction.

#### Method

Here, we investigated the possibility of storing the RNA in the cell lysis buffer RL at -80 °C and compared the yield of RNA and the RIN score to RNA extracted from directly frozen cells.





#### FastGene® RNA Basic Kit

Competitor Q

#### Results/Conclusion:

The RNA yield and RIN score of a freshly prepared RNA isolation did not change when stored at -20 °C or at -80 °C for up to 3 months. Both yield and RIN score showed equivalent or better results than competitor Q's RNA kit.

### **SFast Gene™** RNA Kits - Sizes

### **BASIC**



FastGene® RNA Basic (6 Preps)



FastGene® RNA Basic (50 Preps)



FastGene® RNA Basic (250 Preps)

### **PREMIUM**



FastGene® RNA Premium (6 Preps)



FastGene® RNA Premium (50 Preps)



FastGene® RNA Premium (250 Preps)

#### **Customer Testimonial**

"I can highly recommend the FastGene® RNA Kits:

- easy protocol
- easy procedure without long waiting times
- good price-performance ratio"



# **Jennifer Truong**Physiological Institute, University of Munich, Germany

More testimonials on our website www.nippongenetics.eu



Cat. No.	Product	Content
FG-80006	FastGene® RNA Basic Kit	6 Preps
FG-80050	FastGene® RNA Basic Kit	50 Preps
FG-80250	FastGene® RNA Basic Kit	250 Preps
FG-81006	FastGene® RNA Premium Kit	6 Preps
FG-81050	FastGene® RNA Premium Kit	50 Preps
FG-81250	FastGene® RNA Premium Kit	250 Preps
FG-80L025	FastGene® RNA Lysis Buffer	25 ml
FG-80L125	FastGene® RNA Lysis Buffer	125 ml

# \$ Fast சோட் RNA Kits - Procedure

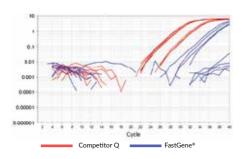
	FastGene®	RNA Basic	FastGene® RNA Premium		RNA Premium
Step	Standard protocol	Large input protocol	Stan	dard protocol	Large input protocol
Sample quantity	< 5 * 10° cultured cells <10 mg animal tissue	< 10 <sup>7</sup> cultured cells <20 mg animal tissue		06 cultured cells g animal tissue	< 10 <sup>7</sup> cultured cells <20 mg animal tissue
Resuspension   lysis of the cells	350 μl buffer RL (with final concentration of 20 mM DTT or TCEP)	600 μl buffer RL (with final concentration of 20 mM DTT or TCEP)	fina	D μl buffer RL (with al concentration of mM DTT or TCEP)	600 μl buffer RL (with final concentration of 20 mM DTT or TCEP)
Filtration of cellular debris					FastGene® RNA filter column O x g for 1 min at room temp.
Optimize RNA binding conditions	350 μl 70% ethanol Mix thoroughly	600 μl 70% ethanol Mix thoroughly		350 μl 70% ethanol Mix thoroughly	600 μl 70% ethanol Mix thoroughly
RNA binding	Load mix onto FastGe Centrifuge at ≥ 10,00 1 min	ene® RNA binding column 0 x g		Load mix onto Fast Centrifuge at ≥ 10, 1 min	Gene® RNA binding column 000 x g
Protein elimination	Add 600 µl of buffer l Centrifuge at ≥ 10,00 30 s			Add 600 µl of buffe Centrifuge at ≥ 10, 30 s	
Desalination	Add 700 µl of buffer l Centrifuge at ≥ 10,00 30 s			Add 700 µl of buffe Centrifuge at ≥ 10, 30 s	
Removal of RW2	Centrifuge at full spec Transfer spin column	ed 1 min to new 1.5 ml collection tube	Ī	Centrifuge at full sp Transfer spin colum collection tube	
Elution of RNA	Add 50 µl of buffer R Centrifuge at ≥ 10,00 1 min	E to membrane center 0 x g		Add 50 µl of buffer Centrifuge at ≥ 10,	RE to membrane center 000 x g 1 min
Optimize DNase I conditions				Add 5 μl 10 x DNa the eluate	se I reaction buffer to
DNA Digestion			Ì	Add 1 µl of DNase Incubate for 10 mir	
RNA rebinding optimization				Add 250 µl of buffe Mix thoroughly by	er RBD to the mixture pipetting
RNA binding				Transfer the mix int column Centrifuge 1 min	to FastGene® RNA mini-elute at ≥ 10,000 x g
Desalination   Elimination of digested DNA				Add 700 µl buffer I Centrifuge at ≥ 10, Transfer spin colum	
Removal of RW2			Ī	Centrifuge at full sp Transfer spin colum collection tube	
Elution of RNA			Í		iffer RE to the membrane at ≥ 10,000 x g 1 min

### *ூ F்\_ு பாட்* RNA Premium Kit

#### No more DNA contamination

Purification of RNA will always have the possibility of genomic DNA contamination. The FastGene® RNA Premium Kit comes with an enzyme, which specifically degrades DNA: DNase I.

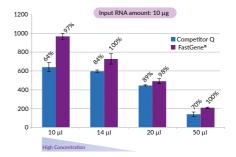
Many RNA purification kits perform the DNA degrading step on the silica membrane. Nonetheless, this step is much more efficient when performed in solution. The FastGene® RNA Premium Kit comprises a DNA degrading step after elution of nucleic acids from the silica membrane.



Detection of genomic DNA contamination using DNA specific primer. RNA isolated using Competitor Q's kit shows considerably earlier Cq values when compared to the FastGene® RNA Premium Kit.

### Mini elute column for the highest concentration and perfect recovery

The FastGene\* RNA Premium Kit comes with a mini elution column. It has a unique design, allowing elution volumes of as little as  $10~\mu$ l. This generates highly concentrated RNA stocks, essential for low amount of tissue or cellular material. The recovery rate is very high (>95%) even at very low elution volumes. At these volumes, not even the market leader can achieve our yield and recovery rates.



Extremely good recovery rates of the FastGene® RNA Premium Kit compared with a RNA kit from a competitor. The mini elution columns of the FastGene® RNA Premium Kit allow a very low elution volume of 10 µl.

#### DNase I treatment after elution vs. on column

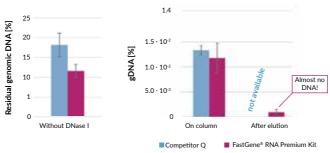
#### Background

Treatment with DNase I is important to remove copurified DNA after RNA purifications. There are two ways for DNase I treatment when working with silica membranes:

- 1. DNase I treatment after elution: this is the standard protocol of the FastGene® RNA Premium Kit.
- 2. DNase I treatment on column: standard protocol for most other RNA purification kits. (this option is also availabe for the FastGene® RNA Basic Kit)

#### Method:

Here, we investigated the effect of two different DNase I treatment options: 1. After elution and 2. On column



#### Results/Conclusion:

DNase I treatment after elution showed the lowest amount of residual genomic DNA with a higher reproducibility when compared to the other tested conditions.

#### **Application**

#### Contamination of DNA in purified RNA: Comparative evaluation of RNA extraction kit, by analyzing the lowest level of contamination

Product

FastGene® RNA Premium Kit (FG-81050, FG-81250)

Manufacturer

NIPPON Genetics EUROPE

The following data is kindly provided by Mr. Tetsuro Ariyoshi, RIKEN Center for Biosystems Dynamics Research, Laboratory of Cell Polarity Regulation, Japan. Thank you for your kind publication.

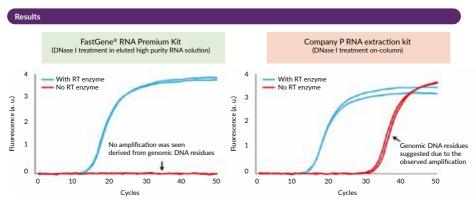
#### Background

Quantitative analysis of expression of RNA is carried out, but since we are conducting experiments in which the detection of genomic DNA cannot be avoided in the design of gPCR primer, we should suppress DNA contamination in the purified RNA as much as possible. It is necessary to analyse RNA extraction kits for this applicability. In this case, FastGene® RNA Premium Kit which adopts "DNase I treatment in eluted high-purity RNA solution" and can expect high DNA removal efficiency as a standard protocol, is compared to a commercially available RNA extraction kit with "DNase I treatment on-column" as the standard protocol, and the comparative evaluation was conducted. As an evaluation method, we examined "whether amplification by residual genomic DNA with qPCR is observed" under the reaction condition "without addition of reverse transcripase (without RT enzyme)".

#### Method

RNA is analysed using two kinds of RNA extraction kits, "when reverse transcriptase treatment was performed" and "when reverse transcription reaction was not performed". qPCR was performed respectively, and the amplification curves were compared.

- 1. Type of initial sample (per Prep): Animal cells (HEK293T 5×105 cells)
- 2. Final elution buffer volume during RNA extraction: 30 µl
- 3. Reverse transcription and qPCR reaction reagents: TaKaRa One Step TB Green PrimeScript PLUS RT-PCR Kit (RR096A)
- 4. Input amount of RNA: Total RNA 60 ng
- 5. One Step RT-qPCR reaction with and without RT enzyme



Signals derived from contamination of genomic DNA are hardly detected from purified RNA using FastGene® RNA Premium Kit.

#### Mr. Tetsuro Ariyoshi:



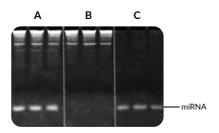
Customers comment

Quantitative analysis of expression of RNA is carried out. However, we are conducting experiments in which the detection of genomic DNA cannot be avoided in the design of qPCR primer. I was looking for an extraction kit in which DNA contamination in purified RNA can be suppressed as much as possible. In purified RNA using FastGene® RNA Premium Kit, almost no signal derived from contamination of DNA was detected, and the amount of expressed RNA could be more accurately quantified compared with purified RNA using competitor's products. In experiments where it is necessary to minimize DNA contamination as much as possible, such as when primers with junctions cannot be designed, the FastGene® RNA Premium Kit is the product of choice. 85

## சிர் திர்க்க திரிக்க த



- Simply add to your RNA kit and enrich miRNA molecules
- Suitable with competitors and FastGene RNA extraction kits
- 3x higher amount of pure miRNA compared to competitors
- Easy-to-use, only one additional step in the protocol



(A) Purification of total RNA + miRNA by using the miRNA Enhancer together with the RNA Basic Kit. (B) Purification of RNA by using only the RNA Basic Kit without the miRNA Enhancer. (C). Purification of pure miRNA together with the RNA Premium Kit.

### Enrichment of miRNA using a standard RNA purification kit

The FastGene® miRNA Enhancer allows the binding of small RNA to the column of your standard RNA purification protocol. With a single additional step you just have to add the RNA enhancer solution during the purification. Dependent on the kit or protocol you are using, you obtain pure miRNA or total RNA + miRNA. Use the miRNA Enhancer togehter with our FastGene® RNA Premium Kit and you will get pure miRNA. Or use the FastGene® RNA Basic Kits and you will obtain total RNA together with miRNA. The miRNA Enhancer is also compatible with other RNA extraction kit.



Use the mitRNA Enhancer with FastGene® RNA Basic kit or FastGene® RNA Premium Kit for the recovery of small RNA

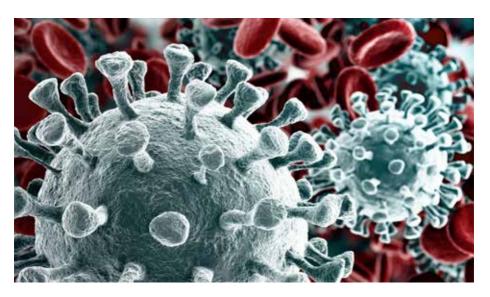
Cat. No.	Product	Content
FG-RNAE-25	FastGene® miRNA Enhancer Kits (4 times×25)	100 Preps



# The importance of miRNA

Micro RNAs (miRNAs) form a group of about 2,500 molecules known for humans so far, which bind to proteins from the Argonaute family (AGO). A small-sized miRNA-AGO complex binds to specific mRNA sites, that is determined by the sequence of the miRNA. The Argonaute protein either blocks protein translation of the mRNA or eliminates the mRNA by cleaving it. Therefore, if a miRNA-AGO complex interacts with a specific mRNA, the gene is silenced and its corresponding protein will no longer be produced.

## **SFastGene™** RNA Viral Kit



	Quick pro	cedure
--	-----------	--------

Easy protocol

Consistent performance

Suitable for SARS-CoV-2 purification

SPECIFICATIONS		
Parameter	High copy plasmid	
Max. sample volume	300 μl / prep	
Preparation time	20 min	
Maximum loading volume	800 μΙ	
Maximum elution volume	30 μΙ	
Operation	Spin column centrifugation	
Storage temperature	Room temp.	

#### Ordering information

Cat. No.	Product	Content
FG-82050	FastGene® RNA Viral Kit	50 preps
FG-82300	FastGene® RNA Viral Kit	300 preps

#### Highest quality viral RNA

The FastGene® RNA Virus kit purifies RNA to a grade comparable to market leaders and delivers RNA of the highest quality. The RNA is therefore ideal for downstream applications, such as reverse transcription and qPCR.

#### Fast and convenient procedure

The FastGene® RNA Virus kit is a silica-membrane based RNA purification method optimized for viral RNA. The procedure was designed to be straight forward and easy to perform. The kit employs the silica membrane technology for the fastest and most convenient high purity RNA isolation, instead of conventional alcohol precipitation or phenol / chloroform extraction.

#### Virus RNA extraction from different sources

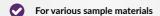
- Cell-free fluid
- Plasma and serum
- Urine
- Virus infected samples
  - Cell culture supernatant

#### Compatible with SARS-CoV-2 Diagnostic Kits

The FastGene® RNA Virus kits purifies RNA to a quality that it can be used with commercially available virus detection diagnostic kits.

# \$ Fast சோட் Mag Virus Kit





Simple and short protocol

Easy to automate

Suitable for SARS-CoV-2 purification

#### Purification of viral RNA and DNA

The FastGene® Mag Virus Kit is based on magnetic bead technology. It has been developed for the isolation of total nucleic acids (DNA and RNA) from a variety of sample materials:

- Blood samples
- Liquid samples (e.g. plasma, serum, urine, swab wash)
- Tissue samples
- Feces

#### Very quick procedure

The FastGene® Mag Virus kit has a quick protocol and the procedure is very easy to perform. The obtained nucleic acids are of highest quality and can be used directly as a template for downstream applications such as PCR, qPCR, qRT-PCR or for any other kind of enzymatic reaction.

#### Easy to automate

The kit is suitable for manual use but also can easily be used in automated systems. Automation protocols are available for KingFisher\* Flex and BioSprint\* 96 systems. The kit is also compatible with liquid handling roboters, e.g. Hamilton\* or Tecan\*.

#### Compatible with SARS-CoV-2 Diagnostic Kits

The FastGene® Mag Virus kit purifies RNA and DNA to a quality that it can be used with commercially available virus detection diagnostic kits. The kit also enables handling of potentially infectious samples.

Cat. No.	Product	Content
FG-84096	FastGene® Mag Virus Kit	96 preps
FG-84960	FastGene® Mag Virus Kit	10 x 96 preps
FG-845K	FastGene® Mag Virus Kit	5000 preps
FG-8425K	FastGene® Mag Virus Kit	25000 preps

## *FastGene*<sup>™</sup> Plasmid Mini Kit



- High yields of plasmid DNA
- Cost effective preparations
- Optimum lysis protocol
- LB-Broth capsules included

#### One kit with all components

Each kit comes with ready-to-use LB-Broth capsules. Add one LB-Broth capsule in 40 ml water, autoclave your solution and start your cloning experiment. The kit includes everything that is needed for a plasmid preparation.



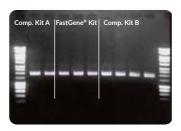
Each Plasmid Mini Kit comes with the easy-to-use LB-Broth capsules.

#### High and low-copy plasmid DNA preparation

The FastGene® Plasmid Mini Kit is designed for rapid small scale isolation and purification of high copy and low copy plasmid DNA. The purified plasmid DNA is of high quality and ready to use in low-salt Tris buffer, suitable for typical downstream applications: Cloning, sequencing, PCR, transformation and restriction analysis.

#### Fast protocol and high yield

The FastGene® Plasmid Mini Kits are faster than competitors with comparable yield. This allows you to save time and perform your downstream application quicker.



pBluescript plasmid DNA was isolated from a 1.4 ml E. coli culture according to the recommended procedures of the different kits and eluted in 50 µle lution buffer. 2 µl of each eluate were loaded on a 0.7% TAE agarose gel. FastGene® Plasmid Mini Kits yield an equal amount of plasmid DNA in a faster time compared to other suppliers. The preparation with the FastGene® Plasmid Mini Kit was performed by using the Fast Protocol (next page).

Cat. No.	Product Content	
FG-90402	FastGene® Plasmid Mini Kit	100 preps + 10 LB-Broth capsules
FG-90502	FastGene® Plasmid Mini Kit	300 preps + 10 LB-Broth capsules

# \$ Fast சோட் Plasmid Mini Kit

1	High cop	y plasmid	Low copy plasmid
	Fast protocol	Standard protocol	Low copy protocol
Harvest of bacteria	ON culture 1 - 3ml >10,000ipm ; Imin. Remove the supernature	ON culture 1 - 5ml >10,000rpm ; 2min. Remove the supernatant	ON culture 5 · 10mi >10,000epm ; 2min. Remove the supernatant
Lysis	200yl of mP1 Vortoxing 200yl of mP2 Invert the tube 2min at room temperature 300yl of mP3 Invert the tube	200pl of nP1 Vortexing 200pl of nP2 Invert the tube 2min at room temperature 200pl of nP3 Invert the tube	400µl of mP1   Vortoxing 400µl of mP2   Invert the tube 2min at room temperature 600µl of mP3   Invert the tubel
Lysate clarification	13,000rpm ; 2min	13,000rpm ; 2min	13,000rpm; 3min
Sample loading	Load the supernature 13,000/pm; 30sec	Load the supernature 13,000 rpm; 30 sec	Load 750yl of the supernatant 13,000rpm; 30sec
Membrane washing	150µl mP4	400µl of mP4 13,000rpm; 30sec 600µl of mP5 13,000rpm; 30sec	400µl of mP4 13,000rpm; 30sec 600µl of mP5 13,000rpm; 30sec
Membrane drying	300µl mP5 13,000rpm; 3min	13,000rpm; 2min	13,000rpm; 2min
Elution	50µl of mP6 2min at room temperature 13,000rpm ; 2min	50µt of miP6 2min at room temperature 13,000rpm ; 2min	50µi of preheaded (70°C) mP6 2min at norm temperature 13,000rpm; 2min

SPECIFICATIONS					
Parameter	High copy plasmid	Low copy plasmid			
Max. sample volume	1-5 ml over-night culture	5-10 ml over-night culture			
Typical yield	< 25 μg	< 25 μg			
Elution volume	50 μΙ	50 μl			
Binding capacity	40 μg	40 μg			
Size of vector	< 15 kb	< 15 kb			
Prep time	26 min / 12 samples	36 min / 12 samples			
Format	spin column	spin column			

# \$ Fast சோட் Gel/PCR Extraction Kit

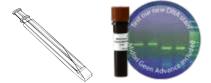


- Very high recovery rate
- Cost effective preparations
- Fast and convenient procedure
- MIDORI<sup>Green</sup> Advance and Gel Band Cutter are included

SPECIFICATIONS						
Parameter	Gel Extraction	PCR Clean-up				
Max. sample volume	300 mg agarose gel	100 μl PCR mix				
Gel	< 2,5% TAE or TBE					
Typical Recovery	70-80%	80-90%				
Binding capacity	10 μg	10 μg				
DNA fragment size	50 bp - 10 kb	50 bp - 10 kb				
Primer removal		< 25 bp				
Elution volume	20-50 μΙ	20-50 μΙ				
Prep time	20 minutes	20 minutes				

### Two in one - DNA cleanup from agarose gels and PCR

The FastGene® Gel/PCR Extraction Kit is designed for the extraction of DNA from agarose gels and for the purification of PCR products. DNA fragments purified with FastGene® Gel/PCR Extraction Kits are ready for direct use in all common downstream applications, like sequencing, ligation and transformation, restriction digestion, microarray analysis, PCR and in vitro transcription.



Each Gel/PCR Extraction Kit contains 5 Agarose Gel Band Cutters and 50  $\mu$ l MIDORI Green Advance. Everything you need to cut out your DNA fragments!

#### Get a free sample!

Convince yourself and test the Gel/PCR Extraction Kit for free. Just contact us and get your free sample very soon!

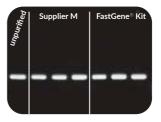
Cat. No.	Product	Content
FG-91202	FastGene® Gel/PCR Extraction Kit	100 preps + 50 μl MIDORI <sup>Green</sup> Advance + 5 Gel Band Cutter
FG-91302	FastGene® Gel/PCR Extraction Kit	300 preps + 50 $\mu$ l MIDORI Green Advance + 5 Gel Band Cutter
FG-830	FastGene® Agarose Gel Band Cutter	50 pieces

### டு *F் துட்டு و* Gel/PCR Extraction Kit



#### Easy workflow

The FastGene® Gel/PCR Extraction Kit provides spin columns, buffers, and collection tubes for silica-membrane-based purification of DNA fragments from agarose gels and PCR products. With a simple and fast bind-wash-elute procedure you can purify DNA ranging from 15 bp to 10 kb with an elution volume of 20-50 µl.



PCR fragments of 300 bp were purified from 40 µJ of a PCR stock solution using FastGene® Gel/ PCR Extraction Kit and a competitor kit, according to manufacturers protocol. 5 µJ of eluted DNA were analyzed on a 1.5% TAE agarose gel. The figure demonstrates that the FastGene® Gel/PCR Extraction Kit shows up to 90% of DNA recovery.

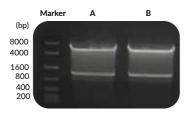
#### Extraction of large DNA fragments with the FastGene® Gel/PCR Extraction Kit

#### Background:

It is a well-known problem that the recovery of DNA fragments larger than 1 kb proves to be difficult and leads to the loss of large amounts of DNA. In this AppNote the FastGene® Gel/PCR Extraction Kit was used for the isolation of two DNA bands after restriction digestion.

#### Method:

A 6.9 kb large plasmid was digested with a restriction enzyme. The restriction digest was analysed by agarose gel electrophoresis at 100 V for 20 min. The 0.7% agarose gel was produced using 1x TAE buffer (Fig 1.). The target fragments were excised out of the gel and transferred in a 1.5 ml tube. The fragments were purified with the FastGene® Gel/PCR Extraction Kit. 100 ng of each purified DNA fragment were electrophoresed again at 100 V for 20 min (Fig. 2).



**Fig. 1:** Identification of two restriction sites (5.4 kb and 1.5 kb) of the plasmid after restriction.

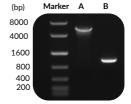


Fig. 2: Clear identification of the two DNA fragments after extraction with the FastGene® Gel/PCR Extraction Kit.

#### Results/Conclusion:

Both fragments show a good recovery rate after extraction. The customer also highlighted the fast preparation, easy handling, high recovery rate for large fragments and the unproblematic performance in downstream applications.

# சி சிக்க பிர்க்க நிருந்தின் இது பிருந்தின் இது பி



- Remove dyes from sequencing products
- Avoid sequencing blobs
- Easy protocol

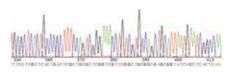
Resin preparation	8.0 ml Buffer DT Vortexing >30 min at room temperature
Column	Apply 750 μl hydrated resin into the column 750 x g; 3 min
preparation	Transfer the column into new tube
Loading sample	Up to 20 μl of sample solution
Sample purification	750 x g ; 3 min

Workflow of the FastGene® Dye Terminator Removal Kit.

#### Optimized sequencing results

The FastGene® Dye Terminator Removal Kit removes dye terminator molecules from sequencing samples, using an efficient and reasonably priced gel filtration. The kit includes a bottle of gel filtration matrix, resuspension buffer and filter spin columns. At first, an aliquot of the matrix is mixed with resuspension buffer. Thereafter, the equilibrated matrix is transferred into filter spin columns. After a brief spin, the sequencing reaction can be loaded onto the column. The last centrifugation step elutes the purified sample. All components can be stored at room temperature.

SPECIFICATIONS					
Parameter	Dye Terminator Removal				
Max. sample volume	20 μl sequencing reaction				
Recovery	> 90%				
Prep time	5 minutes				
Storage	room temperature, 12 months				



DNA purified by the FastGene® Dye Terminator Removal Kit shows a very good performance in sequencing experiments, no dye blobs are detected.

Cat. No.	Product	Content
FG-9411	FastGene® Dye Terminator Removal Kit	50 preps



# Free Sample?

You would like to test our RNA or DNA kits? No problem! Just give us a call or write us an email and get your free sample very soon.



info@nippongenetics.de

 info@nippongenetics.de

www.nippongenetics.eu

# **SFast Gene™** Magna Stands



- Neodymium magnets for optimal separation
- Very quick separation of magnetic beads from the solution
- Stable pellet even during resuspension
- Complete collection of magnetic beads for less material loss

#### Adjustable side position magnets

By securing a pellet (with neodymium magnets) on the side of the tube walls rather than at the bottom, the MagnaStands allow complete removal of the supernatant without touching the pellet. Additionally, with the MagnaStand 1.5, the vertical position is adjustable allowing the magnets to be precisely placed on the tube according to volume used in the purification. Leaders in NGS are recommending the MagnaStand for use with their products!

#### No more carry-over effects

Magnetic beads have long been used to isolate nucleic acids as well as recombinant proteins. These purifications can be problematic, often resulting in carry-over contaminants when the pellet is disturbed or not allowing to completely remove the supernatant. The design of the FastGene® MagnaStand elegantly solves both issues.

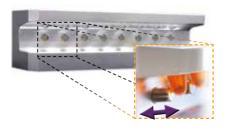


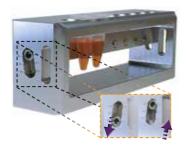
FastGene® MagnaStands are the best tool to easily purify magnetic beads from small volumes. The magnetic beads are firmly held in one position of the tube wall. This prevents accidental aspiration of magnetic beads.

Cat. No.	Product	Size
FG-SSMAG96	96-Well FastGene® MagnaStand	96 Wells
FG-SSMAG96LV	96-Well FastGene® MagnaStand low volume	96 Wells
FG-SSMAG2	FastGene® 0.2 ml MagnaStand	8 x 0.2 ml
FG-SSMAG1.5	FastGene® 1.5 ml MagnaStand	8 x 1.5 ml

# Fast Gene™ Magna Stands







#### 96-Well MagnaStand

- Magnetic stand for reliable high-throughput purification
- Optimal positioning of full- and half-skirted 96-well plates
- Purification from very small volumes (5 μl)
- Ultra low elution volume version (3 μl)

#### 0.2 ml MagnaStand

- 8 magnets for 0.2 ml reaction tubes
- Perform up to 8 purifications in parallel
- $\bullet$  Purification of DNA from very small volumes (down to 3  $\mu l)$
- Each magnet position is adjustable for close contact

#### 1.5 ml MagnaStand

- 8 Ultra strong extra large magnets for larger volumes
- Suitable e.g. for the purification of recombinant proteins
- Adjust the magnet position to the volume of your sample

#### Test the MagnaStands for free!

Each FastGene® MagnaStand uses neodymium magnets, the strongest type of permanent magnet commercially available.

Convince yourself and contact us for free testing.



#### **Customer Testimonial**

"NIPPON Genetics EUROPE provided us a 0.2 MagnaStand for testing. I was totally excited and ordered several MagnaStands for the whole lab. I tested several comparison products, but none of them was so convincing as the FastGene® MagnaStand. The beads are kept punctually, and the magnets can be positioned exactly with the supplied allen key. Meanwhile, the magnetic stands are present in the whole institute. Each laboratory table has now a MagnaStand."



Carolin Dreier Institute of Human Genetics, University Hospital Münster, Germany More testimonials on our website www.nippongenetics.eu







P. 104

P. 100

# **Section** Restriction Enzymes

Cut your DNA the easy way



- 115 restriction enzymes for all your needs
- FastCut digestion in just 5-15 minutes
- No Star activity
- Highest activity and purity

#### What are Restriction Endonucleases?

Restriction enzymes cleave double-stranded DNA at or near a specific recognition site. These enzmyes are classified into four types, based on their subunit structure, cofactor requirements and specificity of cleavage.

#### Perfect for your cloning experiments

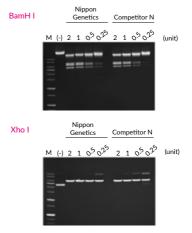
Our restriction enzymes belong to type II, which cleave DNA within or near the recognition sequence. The restriction enzymes cleave double stranded DNA either at the center of both strands to yield "blunt ends" or at a staggered position leaving overhangs called "sticky ends".





#### Excellent activity - compare it to the world market leader

Our restriction enzymes show an excellent activity, at least as high as the world market leader, competitor N. Let's ask you this simple question: Why paying higher prices for the same results? Our restriction enzymes show superb activity, purity, no star activity and promise satisfying results. It's that simple.



Our restriction enzymes (BamH I and Xho I) show at least the same activity as competitor N. M (Marker), (-) negative control. Cut your DNA the easy way

#### Find the restriction enzymes you need - with the Enzyme-Finder

Find your suitable restriction enzyme with our practical Enzyme-Finder by either typing in the name, recognition sequence or overhang sequence. The enzyme list shows you all important properties of our 115 restriction enzymes. Also listed are Isochizomers, which have the exact same recognition and cutting site. Check our the Enzyme-Finder: www.nippongenetics.eu/en/enzyme-finder/

#### Choose between 115 different restriction enzymes for your cloning experiments:



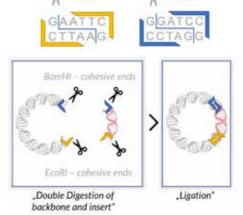


### Digestion in just 5-15 minutes - The FastCut protocol

Most of our restriction enzmyes are supplied with the FastCut buffer, enabling a digestion in just 5-15 minutes. Take 5-10 units of the FastCut restriction enzyme per  $\mu g$  DNA and incubate at the recommended temperature for at least 5 minutes.

#### Double digestion for easy cloning

A vector and insert DNA can be cloned by cleaving both DNA fragments with two different restriction enzymes. This generates two different overhang ends for each fragment. The double digestion strategy prevents the vector from self-ligation and increases cloning efficiency. Most of our restriction enzymes are 100% active in the FastCut buffer, making double digestion simple.



#### 1. Double digestion using colour-coded buffers:

If possible, use the buffer in which both enzymes have 100% activity. Example: For performing a double digestion reaction using Not I and Pst I, simply select Buffer III, because both enzymes are 100% active in Buffer III.

If there is no optimal buffer for both enzymes, use a non-optimal Buffer and adjust the number of units or incubation time for the slower rate of cleavage. Example: For performing a double digestion reaction using Not I and Pvu II, we recommend to select Buffer II (100% activity of Pvu II) and double the units of Not I, as Not I shows 50 % activity in Buffer II.

#### 2. Double digestion using the FastCut buffer:

Most restriction enzymes are 100% active in FastCut buffer, making the buffer choice for double digestion very simple.

#### 3. Setting up a double digestion with a unique buffer

Some restriction endonucleases require a unique buffer for maximum activity: If a restriction enzyme requires a unique buffer, please refer to the table "activity chart of common restriction enzymes in the five unique buffers", for selection of the ideal double digestion buffer.

# **Section** Restriction Enzymes

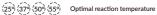
					Activity in FastGene Buffer [%]		
Enzyme	Cat. #	Sequence 5' → 3'	Enzyme Properties	1	П	III	IV
Aat II	FG-AatII	GACGT↓C	(97) <b>60</b> (10 (10 (10 (10 (10 (10 (10 (10 (10 (10	0	25	25	100
Acc I	FG-Accl	GT↓MKAC	<b>⊕∞</b> ∨ <b>⊝ ⊚</b>	75	100	100	100
Acc III*	FG-AccIII	T↓CCGGA	69 No @ Gm	0	25	100	0
Acu I	FG-Acul	CTGAAGN <sub>15</sub> ↓	® <b>0 0</b>	50	50	75	100
Afl II	FG-AfIII	C↓TTAAG	® <b>0 0</b>	75	100	75	100
Age I	FG-Agel	A↓CCGGT	⊕ 60 00 60	100	50	0	100
Alw I	FG-Alwi	GGATCNNNN↓N	(37) 68 (V (m) 68	50	50	10	100
Alw26 I	FG-Alw26I	GTCTCN NNNN	(ii) (ii) (iii) (	75	100	50	100
Apa I	FG-Apal	GGGCC1C	900000	100	25	0	100
ApaL I	FG-ApaLI	G↓TGCAC	®® 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	50	100	50	100
Apo I	FG-Apol	R↓AATTY	@ <b>@</b> • <b>@</b>	10	75	100	75
Asc I	FG-Ascl	GG\$CGCGCC	⊕••••	0	0	0	100
Ava I	FG-Aval	C↓YCGRG	£3000000000000000000000000000000000000	25	100	100	100
Ava II	FG-Avail	G\$GWCC		100	100	50	100
Avr II	FG-Avril	C↓CTAGG		100	50	50	100
Bal I*	FG-AVIII FG-Ball		(#) <b>(6)</b> (1)	0	75	25	75
BamH I*		TGG↓CCA	(9) 69 60 60 C	75	100	100	100
	FG-BamHI	G↓GATCC	<b>⊕</b> ®■ <b>◎</b>		100	100	
Bcl I	FG-Bcll	T↓GATCA	(6) No 11 (10) No	50			75
Bgl I	FG-BgII	GCCNNNN\\NGGC	⊕⊕⊕⊕	75	75	100	50
Bgl II	FG-BgIII	A↓GATCT	<b>⊕⊚</b> □ <b>⊚</b>	10	75	100	10
Bsa I	FG-Bsal	GGTCTCN↓NNNN	<u>⊕⊚00</u> ⊖⊕	50	100	100	100
BsaW I	FG-BsaWI	W↓CCGGW	@ <b>@</b> ( ) @	50	100	100	100
BsiW I	FG-BsiWI	C↓GTACG	<b>⊕●●</b>	50	75	100	50
BsmB I	FG-BsmBI	CGTCTCN↓NNNN	<b>⊚⊚</b> • • • • •	10	50	100	25
BsoB I	FG-BsoBI	C↓YCGRG	⊕ ⊕ ∪ ❷	10	100	100	100
BspE I	FG-BspEI	T↓CCGGA	<b>⊕</b> ⊕⊕ <b>⊚</b>	10	10	100	10
BsrF I	FG-BsrFI	R↓CCGGY	☞•••	75	100	100	100
BstY I	FG-BstYI	R↓GATCY	@®□@	50	100	75	100
BtsC I	FG-BtsCl	GGATGNN↓	⊕ 😡 🕡 🚳	75	100	100	100
Cfr10 I*	FG-Cfr10I	R↓CCGGY	☞ • • • •	10	10	10	25
Cfr42 I	FG-Cfr42I	cccc↓gc	###	100	50	25	75
Cfr9 I	FG-Cfr9I	C↓CCGGG	③* <b>6</b> 9 □ Θ	0	0	100	0
Cla I	FG-Clal	AT↓CGAT	☞ 💇 🕜 😔 🕞 🔘	50	75	75	
CviA I	FG-CviAl	<b>↓</b> GATC	③ 69 0 0 0 0 0 0	10	50	10	
Dde I	FG-Ddel	C↓TNAG	<b>∞</b> 60000	25	50	100	50
Dpn I	FG-DpnI	GA↓TC	⊕ 👓 🕜 🚳	75	100	100	100
Dpn II*	FG-DpnII	<b>↓</b> GATC	<b>3760000</b>	25	75	100	75
Dra I	FG-Dral	TTT↓AAA	③◎ ◎ ◎	75	100	50	100
Eag I	FG-Eagl	C↓GGCCG	ℬே	10	25	100	10
Eco47 I	FG-Eco47I	G↓GWCC	3760 □ 00 00 00	100	100	100	100
EcoN I	FG-EcoNI	CCTNN\$NNNAGG	(37) 63 O 60	50	100	75	100
EcoO109 I	FG-EcoO109I	RG↓GNCCY	3769 ₩ 60	50	75	100	100
EcoR I*	FG-EcoRI	G↓AATTC	Ø <b>6 6 6</b>	50	100	75	100
EcoR V	FG-EcoRV	GAT↓ATC	(27) (20 10 10 10 10 10 10 10 10 10 10 10 10 10	0	100	100	50
EcoT38 I	FG-EcoT38I	GRGCY↓C	(37) <b>69 0 0</b>	75	100	0	100
Esp3 I	FG-Esp3I	CGTCTCN NNNN	®® <b>©</b> ⊚ <b>®</b>	25	50	10	100
Fok I	FG-Fokl	GGATGN <sub>12</sub>	®® ∨ ⊚ ⊝ <b>⊚</b>	100	100	10	100
Fsp I	FG-Fspl	TGC↓GCA	(37) <b>80</b> (2 (9) <b>60</b>	75	100	50	100
Hae II	FG-Haell	RGCGC LY	®® ∨ ⊚ <b>®</b>	10	100	100	100
Hae III	FG-Haell	GG↓CC	@@ 0 @	50	100	75	100
Hga I	FG-Hgal	GACGCN <sub>4</sub> N <sub>4</sub>	@ <b>@</b> 0@	100	75	10	100
Hinc II	FG-HincII	GACGCN <sub>4</sub> ↓N <sub>4</sub> GTY↓RAC	£ 60 € 60 € 60 € 60 € 60 € 60 € 60 € 60	75	50	50	100
Hind II	FG-Hindll	GTY\$RAC	®® <b>0</b> ⊚ <b>0</b>	100	100	50	100
			90000				
Hind III	FG-HindIII	A\$AGCTT		25	100	75	100
Hinf I	FG-Hinfl	G\$ANTC	<b>₩₩₩₩</b>	50	100	100	100
HinP1 I	FG-HinP1I	e∱cec	<b>⊕⊕⊕⊕</b>	50	100	100	75
Hpa I	FG-Hpal	GTT↓AAC	⊕® <b>0</b> ⊕ <b>3</b>	0	50	25	100
Hpa II	FG-Hpall	C↓CGG	⊕ ⊕ ∪ ⊝ ⊚	100	75	50	100
Hph I	FG-HphI	GGTGAN7↓	⊕⊕00	100	75	10	100
Hpy188 I	FG-Hpy188I	TCN↓GA	<u>@@∪</u> @	50	75	50	100
Hpy99 I	FG-Hpy991	CGWCG↓	🕾 🛛 🖂 🚳 📗	100	25	10	

# **Section** Restriction Enzymes

_					Activity in FastGene Buffer [%]				
Enzyme	Cat. #	Sequence 5' → 3'	Enzyme Properties		Ш	Ш	IV		
HpyCH4 V	FG-HpyCH4V	TG↓CA	⊕⊚ 0 ⊚	75	100	25	100		
Kpn I	FG-Kpnl	GGTAC↓C	ூ № • •	100	50	0	100		
Kpn2 I	FG-Kpn2I	T↓CCGGA	<b>⊛</b> •••	100	25	75	50		
Lsp1109 I	FG-Lsp1109l	GCAGCN7↓N3	⊕ ⊕ ⊕ ⊚	25	75	100	100		
Mbo I	FG-Mbol	↓GATC	®®®®⊝⊚	75	100	100	100		
Mbo II	FG-Mboll	GAAGAN7↓	® ⊕ ⊕ ⊚	100	100	50	100		
Mlu I	FG-Mlul	A↓CGCGT	<b>?</b> •••••	25	75	100	50		
Mnl I	FG-Mnll	CCTCN6↓	ூ @ ● ●	75	100	75	100		
Mse I	FG-Msel	T↓TAA	ூ ⊕ ∨ ⊚	75	100	100	100		
Msp I	FG-Mspl	C↓CGG	ூ № 00 00	75	100	75	100		
MspA1 I	FG-MspA1I	CMG↓CKG	⊕⊕•	0	100	75	100		
Mun I	FG-Munl	C↓AATTG	ூ ⊕ ⊕ ⊚	100	100	10	100		
Nae I	FG-Nael	GCC↑GGC	⊕ • • • •	100	100	25	100		
Nco I	FG-Ncol	C↓CATGG	ூ⊚ • •	50	100	100	75		
Nde I	FG-Ndel	CA↓TATG	ூ 🚱 🕜 🚳	75	100	100	100		
NgoM IV	FG-NgoMIV	G↓CCGGC	ூ⊚⊚	25	75	0	100		
Nhe I	FG-Nhel	G↓CTAGC	®® <b>®</b> ●	100	100	10	100		
Nla IV	FG-NlaIV	GGN↓NCC	⊕⊕⊚	0	10	10	100		
Not I	FG-NotI	GC↓GGCCGC	⊕60	0	50	100	0		
Nru I	FG-Nrul	TCG↓CGA	⊕ № ӨӨӨ	0	50	100	75		
Nt.BstNB I	FG-NtBstNBI	GAGTCNNNN↓	<b>⊛</b> •••	0	10	100	0		
PaeR7 I	FG-PaeR7I	C↓TCGAG	⊕ № 00 00	25	100	10	100		
PflM I	FG-PfIMI	CCANNNN↓NTGG	3769 □ 0−0	0	100	100	50		
Ple I	FG-Plel	GAGTCNNNN↓N	(37) 🚳 🕠 (29)	75	75	50	100		
PluT I	FG-PluTI	GGCGC↓C	⊕ № 00 ⊕ 60	75	25	10	100		
PspG I	FG-PspGI	↓ccwgg	(37) No (1) (cm)	25	100	75	100		
Pst I	FG-PstI	CTGCA <b>J</b> G	(37) 60 m (4)	100	100	100	75		
Pvu I	FG-Pvul	CGAT↓CG	€ 000	25	75	100	50		
Pvu II	FG-Pvull	CAG↓CTG	37 No (1) O	75	100	25	10		
Rsa I	FG-Rsal	GT↓AC	37 No O GO	100	100	75	100		
Sac I	FG-SacI	GAGCT↓C	⊕ ⊕ ⊕ ⊚	100	75	25	75		
Sac II	FG-SacII	CCGC↓GG	37) 68 W 69 60	50	100	50	100		
Sal I	FG-Sall	GTCGAC	€ 60	0	0	100	0		
Sau96 I	FG-Sau96I	G↓GNCC	∞∞∞∞∞	50	100	100	100		
Sbf I	FG-Sbfl	CCTGCA↓GG	③ ⊕ ∪ ⊚	50	25	10	100		
Sca I	FG-Scal	AGT↓ACT	(37) <b>60</b> (11) <b>6</b> 0	0	0	100	0		
Sda I	FG-Sdal	CCTGCA↓GG	@ <b>@</b>	75	75	0	100		
Sfi I	FG-Sfil	GGCCN3\$NGGCC	<b>⊕⊚⊚⊚⊚</b>	25	100	25	100		
SgrA I	FG-SgrAl	CR↓CCGGYG	⊕6000	100	100	0	100		
Sma I	FG-Smal	CCC1eee	⊕60000	0	0	0	100		
SnaB I	FG-SnaBl	TAC↓GTA	⊕⊚00	100	75	25	100		
Spe I	FG-Spel	ALCTAGT	⊕••••	50	100	75	100		
Sph I	FG-SphI	GCATG↓C	<b>⊕⊚ 0 0</b>	50	100	50	75		
Sse9 I	FG-Sse9l	↓AATT	9000	100	50	50	75		
Ssp I	FG-Sspl	AAT↓ATT	⊕ 🚱 🕡 🚳	50	100	25	100		
Stu I	FG-Stul	AGG↓CCT	⊕⊗00⊖⊚	75	100	75	100		
StyD4 I	FG-StyD4I	↓CCNGG	⊕600⊖6	10	100	100	100		
Swa I	FG-Swal	ATTT↓AAAT	8000	75	75	100	25		
Taq I	FG-Taql	T↓CGA	@ <b>@</b>	50	100	100	100		
TspM I	FG-TspMI	C↓CCGGG	€ 60 00 60	50	75	50	100		
Tth1111	FG-Tth111I	GACN I NNGTC	@ <b>@</b> @	25	100	100	100		
Xba I	FG-Xbal	T↓CTAGA	⊕6000	0	100	100	100		
Xho I	FG-Xhol	C↓TCGAG	€ 60 00 00 00 00 00 00 00 00 00 00 00 00	50	100	100	100		
Xma I	FG-Xmal	C↓CCGGG	⊕600	50	75	25	100		

#### **Chart Legend**











Supplied buffer





Thermal inactivation condition



Not heat inactivatable



FastCut protocol available

# Fast Gene™ Ligases



- T4 Ligase joins blunt ends and sticky ends
- Quick ligation of DNA fragments in 5 min using the Kickspeed ligation kit
- Supplied with optimal buffer conditions for an efficient ligation
- Highest activity and purity

#### **T4 DNA Ligase**

The FastGene® T4 DNA Ligase catalyzes the formation of a covalent bond between the 5'-phosphate and 3'-OH in nicked duplex DNA or between two DNA ends. This activity is very useful to ligate DNA fragments with either cohesive or blunt ends, that are generated by restriction enzyme digestion.

The T4 DNA Ligase can also ligate RNA with DNA or RNA in a double helix with lower efficiency. The enyzme is expressed and purified in E. coli and is free of endonuclease, exonuclease and phosphatase.

### Ligation in 5 minutes using the Kickspeed Ligation kit

The FastGene® Kickspeed DNA Ligation kit is formulated for quick ligation of DNA fragments with cohesive ends within 5 min at room temperature.

Or use the Kickspeed 2X DNA Ligation Mix. This is a ready-touse solution. This master mix enables quick ligation in a short incubation time (< 5min) at room temperature.

#### Applications

- Vector construction
- Linker ligation
- Fragment assembly
- Routine cloning

Cat. No.	Product	Content
FG-T4	FastGene® T4 DNA Ligase	20,000 units (400 U/μl)
FG-T4BP	FastGene® T4 DNA Ligase	100,000 units (400 U/μl)
FG-T4HC	FastGene® T4 DNA Ligase	100,000 units (2000 U/μl)
FG-LK30	FastGene® Kickspeed DNA Ligation kit	30 reactions
FG-LK60	FastGene® Kickspeed DNA Ligation kit	60 reactions
FG-LM50	FastGene® Kickspeed 2x DNA Ligation kit	50 reactions



# Would you like to try it?

You would like to test our Restriction Enzymes or Ligases? No problem! Just give us a call or write us an email and get your enzymes very soon.



+49 2421 554960

www.nippongenetics.eu





Reverse Transcription – Scriptase Basic and Scriptase II	P. 108
PCR Polymerases - Optima / Taq	P. 112
DNAreleasy - Lysis reagent	P. 117
qPCR Products	P. 118
1-Step RT-aPCR Products	P. 120

# Fast Gene™ Scriptase Basic

Perfect Enzymes for Reverse Transcription



- Reverse transcriptase for the quantification of gene expression
- RNase inhibitor included
- For high DNA concentrations
- Enzyme only or cDNA Synthesis Kit

#### 

The FastGene® Scriptase Basic shows higher sensitivity when compared to wildtype MuLV. The Scriptase Basic is able to produce a template from RNA concentrations as low as 0.1 ng.

#### Enzyme only or cDNA Synthesis Kit

The FastGene® Scriptase Basic is available in two forms: Enzyme only contains the enzyme, buffer and dNTPs. The cDNA Synthesis Kit, additionally comes with Oligo dTs, random hexamers and RNase inhibitor.

#### Designed for endpoint RT-PCR

The FastGene® Scriptase Basic was designed for large RNA quantities, typically used in an endpoint RT-PCR. Nonetheless, it is also able to process lower RNA concentrations.

#### Optimized for better performance

The FastGene® Scriptase Basic is an enhanced version of the Murine Leukemia Virus (MuLV) reverse transcriptase. Like the wildtype, it has the ability to synthesize a cDNA strand, with a reduced RNase H activity and processivity. The robustness of the enzyme was greatly increased. It is perfectly suited for large RNA quantities and easy templates.

### No inhibition - Even at large RNA concentration

The special buffer formulation permits a high RNA concentration. Other reverse transcriptases are not able to process such large quantities.

Cat. No.	Product	Content
LS52	FastGene® Scriptase Basic (20,000 units at 200 U/ $\mu$ I) with buffer and dNTPs	100 rxn
LS62	FastGene® Scriptase Basic cDNA Synthesis Kit containing Oligo dTs, random hexamer and RNase inhibitor	100 rxn



Perfect Enzymes for Reverse Transcription



- Reverse transcriptase for the quantification of gene expression
- Very low RNase H activity
- High yield of full-length cDNA
- Synthesis of cDNA from very low amounts of RNA

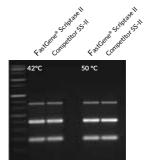
# Everything you need for your reverse transcription

The FastGene® Scriptase II cDNA Synthesis Kit includes all necessary components to perform a reverse transcription. The kit contains the Scriptase II, Oligo dTs, random hexamer and RNase inhibitor.

#### Also available as a 5x ReadyMix

# Engineered enyzme for better performance

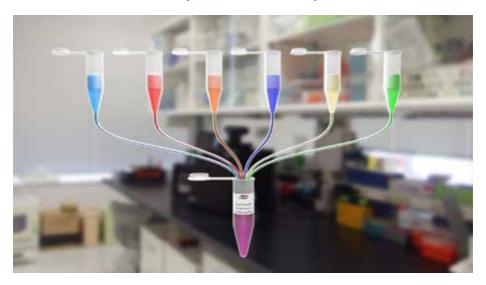
The reverse transcriptase Scriptase II from FastGene® allows the stynthesis of cDNA from very low RNA quantities. The FastGene® Scriptase II contains mutations within the RNase H domain of the MuLV's reverse transcriptase. By reducing the degradation of the RNA during the first-strand synthesis, a higher yield of full-length cDNA is obtained.



Comparison of multiplex PCR using cDNA produced by Competitor SS-II enzyme and FastGene\* Scriptase II at 42°C and 50 °C.

Cat. No.	Product	Content
LS53	FastGene® Scriptase II (20.000 units at 200 U/μl)	100 rxn
LS63	FastGene® Scriptase II cDNA Synthesis Kit containing random hexamer and RNase inhibitor	100 rxn
LS65	FastGene® Scriptase II cDNA Synthesis Kit containing Oligo dTs, random hexamer and RNase inhibitor	100 rxn
LS64	FastGene® Scriptase II cDNA Synthesis 5x ReadyMix	100 rxn

# & Fast செட் Scriptase II - ReadyMix



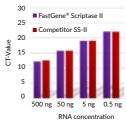
# Reverse transcription: Ready-to-use

The FastGene® Scriptase II cDNA Synthesis 5x ReadyMix is ready-to-use with all necessary ingredients in just one vial. Just add the Scriptase II ReadyMix to your template and you are ready to go.

## Engineered enzymes - optimized for qPCR

The FastGene® Scriptase II delivers superior cDNA templates for downstream applications, e.g. qPCR and NGS. The resulting full length cDNA gives a complete picture of the gene and is able to show modifications, e.g. splicing variants.

# Comparison - GAPDH - qPCR



Comparison of qPCR results using primers for GAPDH produced by using different RNA starting concentration by FastGene® Scriptase II and competitor SS-II enzyme at 42°C

#### **Customer Testimonial**

"I especially like that the Scriptase II leads to stable results. As a result of performing RT-PCR using tumor derived RNA, we were able to detect the expression of genes, where the amplification was unstable with other RT reagents. The amplification of full-length cDNA has also been confirmed. I would love to also try the 5x ReadyMix."



#### Haruko Hayasaka

Department of Bioscience and Biotechnology, Kinki University, Osaka, Japan



Cat. No.	Product	Content
LS53	FastGene® Scriptase II (20.000 units at 200 U/μl)	100 rxn
LS63	FastGene® Scriptase II cDNA Synthesis Kit containing random hexamer and RNase inhibitor	100 rxn
LS64	FastGene® Scriptase II cDNA Synthesis 5x ReadyMix	100 rxn
LS65	FastGene® Scriptase II cDNA Synthesis Kit containing Oligo dTs, random hexamer and RNase inhibitor	100 rxn

# **Technical Data**

## Very fast reverse transcription reactions

#### Purnose

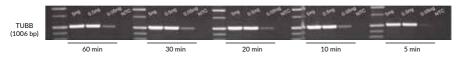
FastGene® Scriptase II is an engineered reverse transcriptase, able to deliver highest quality cDNA from a small amount of RNA. Optimization of enzymatic design has led to one of the most reactive RT-enzymes. This technical note shows the investigation of the minimum time possible of a reverse transcription. We were able to shorten time to 5 minutes with different concentrations of RNA. The resulting cDNA was used in endpoint PCR as well as in qPCR experiments.

#### Method

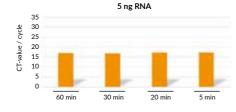
- FastGene® Scriptase II cDNA Synthesis Kit (LS63)
- RNA: Universal Human Reference RNA (Agilent Technologies) Input RNA amount: 5 ng, 0.5 ng, 0.05 ng
- Primer
- TUBB (1006 bp): Endpoint PCR
- GAPDH (138 bp): qPCR

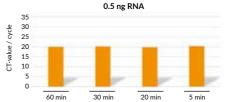
#### Result

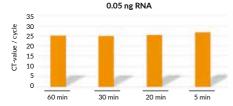
1 Endpoint PCR



(2) (Quantitative PCR







#### Conclusion

FastGene® Scriptase II was able to produce cDNA in 5 minutes.

Result 1: For large PCR products, the band of 0.05 ng RNA after 5 min was slightly weaker. Hence, for products of 1000 bp a 10 min RT step is recommended for low RNA amounts.

Result 2: No difference in CT-value exceeding ± 1 cycle was detected.

FastGene® Scriptase II can therefore be recommended for short-term reverse transcripton reactions.

# *ூ Fast செட*் Optima HotStart ReadyMix



- Proofreading DNA polymerase
- ReadyMix Just add your template and primer
- For very complex templates (up to 20 kb)
- Extremely high fidelity
- Problem solver

# Optimal robustness for very complex samples

The FastGene® Optima can easily handle very complicated templates. The highly purified Taq polymerase gives great efficiency while the proof-reading polymerase guarantees the fidelity. The robustness of both enzymes makes the amplification of complex tissue, such as liver tissue (Fig. 1), possible.



Fig. 1: The comparison between (A) the "best-selling" blended Tag mix and (B) FastGene" Optima polymerase mixture uses with catshark liver DNA (hard to amplify) as a template. The PCR product with a size of 1030 bp was separated on a 1.2% agarose gel. The FastGene" Optima produces less primer dimers and has a higher amplification efficiency.

# Processivity, fidelity and big fragments

The FastGene® Optima polymerase is a mixture of two different types of PCR enzymes – a Taq polymerase and a modified type-B polymerase with excellent proof-reading abilities. Each enzyme is purified using three different chromatography technologies. This results in very high enzyme purity and activity. Optima is extremely robust, making it ideal for a broad range of PCR applications. Standard PCR, challenging PCR, and very long amplicons (over 7.5 kb) are easily handled by this enzyme mixture.

## Optimal efficiency for GC-rich templates

Most polymerases have a low amplification efficiency, if the template DNA is GC-rich. As seen in Fig. 2, the FastGene® Optima has an excellent amplification efficiency even with GC rich templates. The efficiency of the FastGene® Optima is even higher than the efficiency of polymerases specially designed for GC-rich templates (Fig. 2).

	Competitor T			Competitor T FastGene® Optim		® Optima
	1	2	GAPDH M	1	2	GAPDH

Fig. 2: Comparing the ability of Competitor T and FastGene® Optima polymerase mixture to amplify GC-rich DNA fragments. Two fragments with 60.7% GC and 64.3% GC were amplified, resulting in two products of 1839 bp and 1260 bp, respectively. FastGene® Optima had a higher efficiency compared to Competitor Ts polymerase mixture.



Robustness "of a Rhino" is the key advantage of the Optima DNA polymerase. Do you have any problems with your PCR? Just try the Optima - you will get reliable and reproducible results. Anytime!

Cat. No.	Product	Content
LS29	FastGene® Optima HotStart ReadyMix	$500 \times 25~\mu l$ reactions (6.25 ml total volume)

# *ூ Fast செட*் Optima HotStart ReadyMix

# Optimal for SNP-typing

The detection of single nucleotide polymorphism (SNP) requires extreme fidelity, which is guaranteed by the proof-reading activity of the FastGene® Optima (Fig. 3).

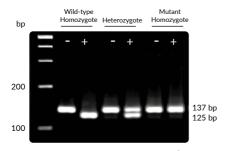


Fig. 3: SNP typing of the ALDH gene using FastGene® Optima polymerase. The ALDH gene, classified as human sensitivity to alcohol, was analysed for presence of SNP by digesting the amplification of homoand heterozygotes using Mboll.

## HotStart - It is your decision when to start

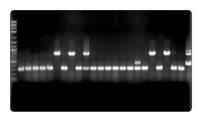
For labs preferring low primer-dimers and an easy room temperature set-up, the HotStart-version of the FastGene® Optima is the best choice. Designed as a master mix, the Optima HotStart ReadyMix combines the superb efficiency and robustness of the Optima enzyme mix with a proprietary antibody that inhibits a preliminary unspecific reaction. This antibody is permanently denatured during the primary PCR activation step. The HotStart ReadyMix comes with all the necessary ingredients for an optimal PCR. Just add your template and primers. Additionally, the ReadyMix includes a loading dye, so that the PCR sample can be directly loaded onto an agarose gel.

# **Applications**

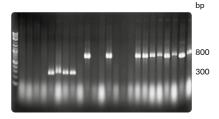
- RT-PCR
- Very complex templates
  - GC-rich templates
- SNP Analysis
- Multiplex PCR
- Any standard PCR application



## Direct PCR from E. coli colonies



VS.



Direct PCR from E. coli colonies using FastGene® Optima HotStart ReadyMix (left gel) or "best-selling" blended Taq mix (right gel). The ReadyMixes were used to determine the presence or absence of inserts. The Optima HotStart ReadyMix yielded a clear electrophoretic pattern without smearing. In addition, Optima was able to amplify clean product from EVERY colony. The competitor was not able to amplify 10 colonies.

#### **Customer Testimonial**

"We tested very successfully the HotStart ReadyMix for duplex-PCR of cDNAs from our knock-down mutants. The PCR reactions show no unspecific products. Additionally this product has an excellent price performance ratio"



**Dr. Matthias Schmidt**Institute for Molecular Life Science
Goethe University, Frankfurt, Germany



# *ூ Fஊ் பேட்* apTaq HotStart Polymerase



- HotStart: Very fast PCR
- Aptamer technology: Reversible enzyme activation or inactivation
- Maximal specificity, sensitivity and yield
- Robust and reliable reaction
- For a wide range of templates

# Redefine your PCR

The FastGene® apTaq DNA polymerase is a recombinant and thermostable Taq-Polymerase using the aptamer based HotStart activation technology. The aptamer allows a reversible and immediate activation of the polymerase, leading to specific priming and a very fast PCR.

# Reversible polymerase activation – The aptamer principle

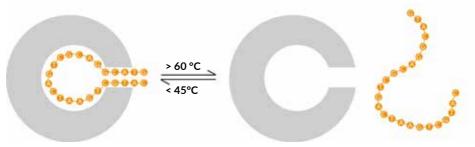
The FastGene® apTaq DNA-Polymerase is a HotStart enzyme, which is completely inactive at room-temperature and only becomes active at higher temperatures. Random primer annealing and unspecific amplification that might occur with standard PCR enzymes are yesterday's problems. In contrast to antibody-based methods, the apTaq DNA-Polymerase includes synthetically manufactured aptameroligonucleotides. At low temperatures, the polymerase is inactivated through reversible aptamer binding. The aptamer acts as a molecular switch, changing its tertiary structure at higher temperatures. Temperatures below 45 °C deactivate the polymerase, whereas temperatures above 60 °C fully activate the enzyme. Therefore, the FastGene® apTaq DNA-Polymerase is less temperature-sensitive and reduces the risk of contamination

## **Applications**

- Fast PCR
- Routine PCR
- PCR using complex templates
  - SNP Analysis
- Any standard PCR application

Inactive apTaq Polymerase inhibited by aptamer





 $The polymerase \ oligonucle otide-aptamer\ mixture\ is\ a\ reversible, temperature-dependent\ HotStart\ system.$ 

Cat. No.	Product	Content
LS34	FastGene® apTaq HotStart Polymerase	500 Units

# *ூ F்சுட்ப் E* BAC-free Taq



- ONA polymerase with no bacterial contamination
- Prevents false positive PCR results from bacterial DNA
- Perfect for bacterial genome analysis

# Conventional Taq-Enzyme # - - | # - - - | syntisco d system and system syntisco d syntisco d system syntisco d syntisc

Amplification of a non-ribosomal gene using E. coli DNA (+) or no template control. The no template control (-) was amplified with standard Taq and FastGene® BAC-free HS Taq. The conventional Taq produced a PCR-product despite no tempate being present, while no product was detected using the FastGene® BAC-free HS Taq. This indicates a bacterial genomic DNA contamination of the conventional Taq polymerase.

# Free of any bacterial contamination

The FastGene® BAC-free HotStart Taq DNA polymerase is based on the single-subunit, wild-type Taq DNA polymerase of the thermophilic bacterium Thermus aquaticus, but is purified from an eukaryotic recombinant expression system. Contaminating DNA, present in most other polymerase preparations, often precludes the accurate interpretation of results, especially when targeting conserved sequences (e.g. the bacterial 165 rRNA region).

# Eukaryotic expression system - No more false positive

Performing PCR with bacterial templates could lead to a false positive result, when using Taq enzymes purified from E. coli expression systems due to a contamination of the Taq enzyme with prokaryotic genomes. The FastGene® BAC free HotStart Taq DNA Polymerase is produced using eukaryotic cells. Hence, no bacterial genome is present.

## **Applications**

- Bacterial genome analysis
- Pathogen detection
- Amplification of low copy DNA templates
- Multiplex PCR
- Specific amplification of complex templates
- RT-PCR



Best choice for 16S/23S microbial screening, E. coli contamination and forensic studies.

Cat. No.	Product	Content
LS33	FastGene® BAC-free HotStart Taq Polymerase	500 Units

# **ூ** Fast சோட் Taq DNA Polymerase



# Taq polymerase with a high purity

The FastGene® DNA Polymerase is based on the single subunit, wild-type Taq DNA polymerase of the thermophilic bacterium Thermus aquaticus. The enzyme is purified using three different chromatography technologies and results in a very high purity and activity.

#### Two different reaction buffers

The enzyme comes with 2 different reaction buffers. Buffer A is a "high yield" buffer, for most amplicons. Buffer B is a standard KCI-based Taq buffer with a higher sensitivity.

#### **Customer Testimonial**

"We are happily using the FastGene® Taq DNA polymerase for over 12 months for routine SNP-analysis. We have chosen FastGene® Taq DNA polymerase since we needed a robust and reliable polymerase. We are very happy with it and the price-performance ratio is excellent!"

Dr. J. Wagner

PlantaLyt GmbH, Hannover, Germany

# \*\*\*\*

# Ordering information

Cat. No.	Product	Content
LS21	FastGene® TAQ DNA polymerase	500 Units
LS21dNTP	FastGene® TAQ DNA polymerase + dNTPs	500 Units + 1 ml dNTPs [2 mM each]
LS22	FastGene® TAQ DNA polymerase	2000 Units





# Everything you need for your PCR

The FastGene® Taq ReadyMix (2X) is a ready-to-use cocktail with two inert tracking dyes and containing all components for PCR, except for primers and template. The 2X ReadyMix contains FastGene® Taq DNA polymerase, Taq buffer, dNTPs, MgCl, and stabilizers.



FastGene® Taq reactions with 1X loading dye reaction buffer. (A) Volumes above wells indicate the volume of the PCR reaction loaded on the gel. (B) On a 1% agarose gel, the blue dye migration corresponds to a 5 kb DNA fragment, and the yellow dye migrates at 75 bp.

Cat. No.	Product	Content
LS26	FastGene® Taq Ready Mix PCR Kit	50 x 50 μl reactions
LS27	FastGene® Taq Ready Mix PCR Kit	250 x 50 μl reactions

# **DNAreleasy Advance**



- PCR done the easy way
- Successful lysis of different biological material
- Very easy-to-use

## From cells to PCR in 15 minutes

Are you tired of the time-consuming extraction processes and costly spin columns that you've been using to prepare samples for DNA amplification? With the DNAreleasy Advance Direct Lysis Kit, we now offer a better solution. The new cell lysing reagent only requires a 15 minutes incubation in a thermal cycler before the DNA is ready-to-use directly for your PCR — without any further sample processing!

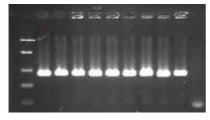
# Successfully used samples

- Saliva
- Hair roots
- Animal tissue (horse, pig liver, etc.)
- Mouse tails and ears
- Plants (leaf, blossom, pollem): Cabbage, maize, canola, soy, sugar beet, etc.
- Drosophila
  - Yeast
- Mollusca

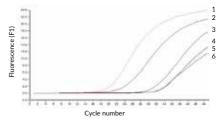
#### Easy procedure



Using DNAreleasy Advance is really easy. Just mix cells with 20 µl of the reagent, place in a a thermal cycler or incubator and heat at 65°C for 5 minutes, followed by 96°C for 5 minutes before holding at 20 °C for 5 minutes. After the lysis, a part or all of the lysate can be added directly to your PCR mix or it can be stored at -20 °C for future use.



Genomic DNA from scallops was isolated with DNAreleasy Advance, and a part of the supernatant was directly added to the PCR reaction. The agarose gel shows the high yield obtained.



Genomic DNA was isolated using DNAreleasy Advance and analyzed by qPCR: (1) positive control human DNA, (2) saliva, (3) hair root, (4) pig liver, (5) drosophila melanogaster, (6) horse meat.

Cat. No.	Product	Content
LS05	DNAreleasy Advance	300 μl, 10 reactions
LS06	DNAreleasy Advance	1.5 ml, 50 reactions

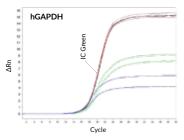
# & Fast சோட் IC Green qPCR Universal Kit



#### No inhibition - For highest sensitivity

It is well-known that SYBR® Green is extensively inhibiting the qPCR. This fact led to the development of SYBR® resistant enzymes. An alternative approach is to develop a dye that does not inhibit the reaction. This dye is named FastGene® IC Green. FastGene® IC Green is an intercalating dye, only detecting double stranded DNA. By not inhibiting the reaction, the FastGene® IC Green Kit is able to detect genes at a lower CT-value, creating a higher sensitivity!

The superior buffer chemistry enables the detection of low copy number genes, which could not be detected with other dyes. The comparison to competitors shows that FastGene® IC Green is one of the best qPCR mixes available. This has been confirmed by customers analysing various genes.



## Universal - for any qPCR instrument

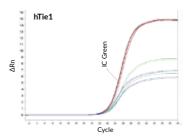
The FastGene® IC Green Kit is universal. The reference dyes come in a separate vial and can be added to the master mixes once. Hence, this kit can be used with qPCR instruments which need a high ROX™ concentration as well as instruments that need a low concentration or no ROX™. A special version with fluorescein is also available

# Robust chemistry for faster results

The FastGene® IC Green buffers were designed to have a superior robustness. This guarantees the linearity of the qPCR and creates a better accuracy, essential for reproducible results. Additionally, qPCRs can be performed at shorter amplification times, for example using fast protocols.

## **Applications**

- Quantification of gene expression
- Quantification of gene copy number
  - Melt-curve analysis
- Detection of gene expression (knock-out analysis)



Comparison of FastGene® IC Green (black & red) with the market leading competitors KB (green) and T (blue). The differences of the C<sub>r</sub>-values were under 1 cycle.

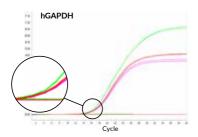
Cat. No.	Product	Content
LS4001	FastGene® 2x IC Green Universal (ROX™)	100 reactions
LS4005	FastGene® 2x IC Green Universal (ROX™)	500 reactions
LS4050	FastGene® 2x IC Green Universal (ROX™)	5000 reactions
LS4101	FastGene® 2x IC Green Universal (Fluorescein)	100 reactions
LS4105	FastGene® 2x IC Green Universal (Fluorescein)	500 reactions
LS4150	FastGene® 2x IC Green Universal (Fluorescein)	5000 reactions

# *ூ Fஊ்டு் - Probe qPCR Universal Kit*



# Robust chemistry for multiplexing

The robustness of the buffer ensures the ability to perform multiplex qPCR. Get the highest sensitivity for multiple targets using the FastGene® Probe Universal Kit. The FastGene® Probe Universal Kit is compatible with all real time PCR instruments.



# Perfect efficiency

For the FastGene® Probe qCPR, use hydrolysis probes, enabling multiplex, and leading to very specific signal and low to none background fluorescence. The buffer chemistry, combined with optimal primer design, is the most important part of a Probe assay based reaction. Here we present the superior buffer system of the FastGene® Probe Universal Kit.

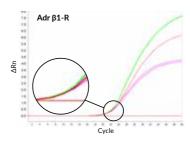
Get a very high dynamic range and reproducible results by using the FastGene® Probe Universal mix. Achieve higher efficiencies and more accurate results.

## Save time with fast protocols

The unique buffer composition enables a faster reaction: apply a fast protocol, available on many modern qPCR instruments, and save plenty of time.

## **Applications**

- Quantification of gene expression
- Quantification of gene copy number
- Multiplex qPCR
- SNP genotyping
- NGS validation



Reactions (25 μl) were set up according to manufacturer's instructions, with 25 ng of hgDNA as template, and 0.5 μM of each primer. PCR was performed for a total of 35 cycles. Green: Competitor KB. Red: Competitor T. Pink: Probe qPCR Universal Kit.

Cat. No.	Product	Content
LS4501	FastGene® 2x Probe Universal (ROX™)	100 reactions
LS4505	FastGene® 2x Probe Universal (ROX™)	500 reactions
LS4550	FastGene® 2x Probe Universal (ROX™)	5000 reactions

# *ூ F⊒stப்பாட*் IC Green 1-Step RT-qPCR



# Robust chemistry for 2 reactions in one tube

The FastGene® IC Green 1-Step mix contains a reverse transcriptase and a DNA polymerase. Having a 1-tube reaction setup for the reverse transcription and for the quantitative PCR has many advantages: 1) The 2x master mix ensures the same concentration of buffer and enzyme when performing the experiment multiple times, 2) it is less prone to wrong mixtures of the reaction mix contents, 3) higher convenience due to less preparation time, and many more.

# Applications

- Quantification of gene expression
- Quantification of gene copy number
  - Melt-curve analysis
- Detection of gene expression (knock-out analysis)

# Ordering information

Cat. No.	Product	Content
LS4301LR	2x FastGene® IC Green 1-Step Mix (low ROX™)	1 ml (100 reactions)
LS4305LR	2x FastGene® IC Green 1-Step Mix (low ROX™)	5 x 1 ml (500 reactions)
LS4301HR	2x FastGene® IC Green 1-Step Mix (high ROX™)	1 ml (100 reactions)
LS4305HR	2x FastGene® IC Green 1-Step Mix (high ROX™)	5 x 1 ml (500 reactions)

# *ூ F் த்து பிதாட* Probe 1-Step RT-qPCR



# High-performance enzymes for incredible sensitivity

The FastGene® Probe 1-Step Mix was developed for the rapid detection of multiple gene expressions using multiplex qPCR directly from RNA. The optimal conditions for the reverse transcription as well as for the DNA polymerisation ensures highest sensitivity and the detection of low copy genes.

#### **Applications**

- Quantification of gene expression
- Quantification of gene copy number
- Multiplex qPCR
- SNP genotyping
- NGS validation

Cat. No.	Product	Content
LS4701LR	2x FastGene® Probe 1-Step Mix (low ROX™)	1 ml (100 reactions)
LS4705LR	2x FastGene® Probe 1-Step Mix (low ROX™)	5 x 1 ml (500 reactions)
LS4701HR	2x FastGene® Probe 1-Step Mix (high ROX™)	1 ml (100 reactions)
LS4705HR	2x FastGene® Probe 1-Step Mix (high ROX™)	5 x 1 ml (500 reactions)



# Free Sample?

You would like to test our DNA polymerases or our qPCR reagents? No problem!



+49 2421 554960



info@nippongenetics.de
 info@nippongenetics.de

www.nippongenetics.eu





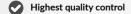
Filter Tips	P. 124
PCR Plastic	P. 128
Aluminium Rack	P. 135
Cryo Tubes	P. 136
Screw Cap Tubes	P. 138
High-Throughput Plastic	P. 140

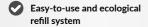
# \$ Fast Gene™ Filter Tips

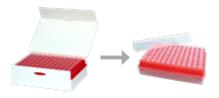
Quality Made in Japan











## Japanese quality

The filter tips are made in Japan. Quality control is one of the highest priorities. This ensures that our tips are free of faults, such as misplacement of the filter, broken tips, missing tips, endotoxins, etc.. All of our tips are free of RNase, DNase, genomic DNA and proteins.

# Easy-to-use PaperRefill system

The FastGene® PaperRefill System is a more ecological way compared to standard refill systems. Just insert the new pipette tip rack in your filter tip box without any tip wobbling during refill.

## More than just plastic

To improve precision and handling, NIPPON Genetics EUROPE provides high quality and modern filter tips. Their maximum compatibility and conformity with a large number of pipettes enable accuracy and comfortability for the daily laboratory work.

## **Customer Testimonial**

"We have been using the refillable filter tips from Nippon Genetics for a broad spectrum of molecular biology techniques, including NGS and array-CGH. We are impressed by their high manufacturing quality and ease of use. The tips are long and thin and the filter does not come in contact with the liquid even if you fill it to the maximum. They also exhibit minimum retention of liquids. It is very easy to refill the empty tip boxes (spare tips come in pre-filled and sterile racks) and by using that system you produce less plastic waste! We highly recommend these tips to all researchers looking for excellent quality, value-for-money filter tips!"



**D. Palaiologou, PhD** Genesis Genoma Lab, Chalandri, Greece

\*\*\*\*

# \$\int\text{Gene\text{\textit{\text{Gene\text{\text{\text{Tiller}}}}}\text{Filter Tips}

Quality Made in Japan



## Precision for lowest volume

The FastGene® Filter Tips come in three different versions for the lowest volumes:

- Filter Tips 10 μl short: The small size guarantees easier handling in such small wells (e.g. 96-well plates or PCR Tubes).
- Filter Tips 10 µl long: Very useful to avoid contamination of the pipette with samples stored in larger tubes.

10 racks with 96 tips

10 racks with 96 tips

• Filter Tips 20 µl: The best option for larger volumes with a small tip.



Cat. No.	Product	Content
FG-FT10S	10 μl short	10 racks with 96 tips
FG-FT10SRF	10 μl short refill	10 racks with 96 tips



Cat. No.	Product	Content
FG-FT20	20 μΙ	10 racks with 96 tips
FG-FT20RF	20 μl refill	10 racks with 96 tips

## Precision for larger volumes

10 μl long

10 μl long refill

FG-FT10L

FG-FT10LRF





Cat. No.	Product	Content
FG-FT100	100 μΙ	10 racks with 96 tips
FG-FT100RF	100 ul refill	10 racks with 96 tips

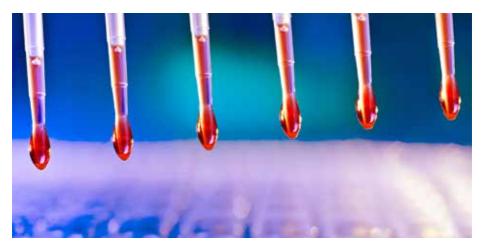


Cat. No.	Product	Content
FG-FT200	200 μΙ	10 racks with 96 tips
FG-FT200RF	200 μl refill	10 racks with 96 tips



Cat. No.	Product	Content
FG-FT1000	1000 μΙ	10 racks with 96 tips
FG-FT1000RF	1000 μl refill	10 racks with 96 tips

# **SFast Gene™** TP Filter Tips



- Compatible with pipettes from Eppendorf, Gilson, Rainin, Thermo, Socorex and many more
- Highest quality control
- Efficient contamination protection

# Highest quality plastic

Quality control is one of the highest priorities. This ensures that our FastGene TP Filter Tips are free of faults, such as misplacement of the filter, broken tips, missing tips, etc.. All of our tips are free of RNase, DNase, genomic DNA and proteins.

## Best volume control

FastGene® TP Filter Tips are transparent, so it is easy to always control the liquid inside the pipette. For clear observation of the aspirated volume, the smallest and the largest tips (10  $\mu$ l, short and 1000  $\mu$ l) come with volume marks.

# Compatibility

The FastGene® TP Filter Tips are compatible with the following single channel pipettes:

		FastGene® TP line Filter Tips					
		FG-TP-10	FG-TP-10L	FG-TP-20	FG-TP-100	FG-TP-200	FG-TP-1000
	Eppendorf (Reference, Research V)	0.5-10 μΙ	0.1-25 µl 0.5-10 µl	2-20 μΙ	10-100 μΙ	20-200 μΙ	100-1000 μΙ
	Finnpipette (Digital, Focus)	1-10 μl 2-20 μl	1-10 μl 2-20 μl	2-20 μl 10-100 μl	10-100 μΙ	20-200 μΙ	100-1000 µl
ettes	Gilson (Pipetman)	P-2, P-10	P-2, P-10	P-20	P-100	P-200	P-1000
nel Pip	Rainin (SL-PL-Series)	0.5-10 μΙ	0.5-10 μΙ	2-20 μΙ	10-100 μΙ	20-200 μΙ	100-1000 μΙ
Single Channel Pipettes	Rainin (XLS-Series)	0.5-10 μΙ	0.5-10 μΙ	20-200 μΙ	20-200 μΙ	20-200 μΙ	100-1000 μΙ
Single	Brand	0.1-2.5 μl 0.5-10 μl	0.1-2.5 μΙ	2-20 μl 5-50 μl, 10-100 μl		20-200 μΙ	100-1000 μΙ
	DragonLab	0.5-10 μΙ	0.5-10 μΙ	2-20 μΙ	10-100 μΙ	20-200 μΙ	100-1000 μΙ
	SOCOREX	0.5-10 μΙ	0.5-10 μΙ	20-200 μΙ	20-200 μΙ	20-200 μΙ	100-1000 μΙ

# \$FastGene™ TP Filter Tips

# Filter protection

The tips are equipped with an anti-aerosol filter. Crosscontamination between samples is avoided, giving you maximum security with your results.





Cat. No.	Product	Content
FG-TP-10	10 μl short	10 racks with 96 tips





Cat. No.	Product	Content
FG-TP-20	20 μΙ	10 racks with 96 tips

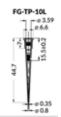


Cat. No.	Product	Content
FG-TP-200	200 μΙ	10 racks with 96 tips

# No DNA adsorption

FastGene® TP Filter Tips are made of tested polypropylene plastic, which stop the adsoprtion of DNA and prevents concentration fluctuations.





Cat. No. Product		Content
FG-TP-10L	10 μl long	10 racks with 96 tips





Cat. No.	Product	Content
FG-TP-100	100 μΙ	10 racks with 96 tips





Cat. No.	Product	Content
FG-TP-1000	1000 µl	10 racks with 96 tips

# SFast Gene™ PCR Tubes





Compatible with most thermal cyclers



Reproducible PCR results



Free of RNase, DNase and human genomic DNA

## No evaporation

The evaporation of samples is a well-known error factor which depends on the quality of the PCR plastic. Preventing evaporation is important especially for users that perform low volume (5 - 10 ul) PCR. FastGene® PCR Tubes and strips are intensively tested, under very stringent conditions.

# **Guaranteed quality**

The performance and reproducibility of your PCR result is significantly influenced by plastics. As a result of a unique manufacturing process, our FastGene® PCR single tubes and 8-well strips fulfill the highest requests of quality. All FastGene® PCR plastic products are manufactured by using ultra pure polypropylene. Proteins are not able to bind to the surface. The tubes and strips have very thin walls but are extremely stable and robust. Because of a very stringent QC procedure the "batch-to-batch" reproducibility of all plastics is extremely high.



#### Test our PCR tubes for free!

We offer PCR tubes with certified quality standards. Convince yourself and contact us for free testing!

#### 0.1 ml PCR Tubes



0.1 ml PCR single tubes with flat caps (1000) Cat. No.: FG-011F



0.1 ml PCR 8-well strips and flat cap strips (120) Cat. No.: FG-017FC

# **SFast Gene™** PCR Tubes



0.1 ml PCR 8-well strips with single flat caps (120) Cat. No.: FG-018WF



0.1 ml flat cap strips (120) Cat.No.: FG-008FCP



0.1 ml PCR 8-well white tube strips with flat cap strips (125) Cat. No.: FG-019FC



0.1 ml PCR 8-well strips (120) Cat. No.: FG-018

## 0.2 ml PCR Tubes



0.2 ml PCR single tubes with flat caps (1000) Cat. No.: FG-021F



0.2 ml PCR single tubes with domed caps (1000) Cat. No.: FG-021D



0.2 ml PCR 8-well strips with single flat caps (120) Cat. No.: FG-088WF



0.2 ml PCR 8-well strips with single domed caps (120) Cat. No.: FG-088WD



0.2 ml PCR 8-well strips without caps (120) Cat. No.: FG-028



0.2 ml flat cap strips (120) Cat.No.: FG-008FC



0.2 ml domed cap strips (120) Cat.No.: FG-008DC



0.2 ml PCR 8-well strips and flat cap strips (120) Cat. No.: FG-016FC



0.2 ml PCR 8-well strips and domed cap strips (120) Cat. No.: FG-016DC



#### Application

# Comparing PCR tubes for Multiplex Probe-based Assay on a Rotor-Gene® Q

Product

FastGene® 0.2 ml PCR tubes with flat caps (FG-021F)

Manufacturer

NIPPON Genetics EUROPE

The following data is kindly provided by Dr. Birgit Klinkert, ARDEYPHARM GmbH, Herdecke, Germany

#### Background

The detection of the non-pathogenic Escherichia coli strain Nissle 1917 (EcN) in stool samples is standardly performed in this laboratory using strain specific TaqMan® Probes. Here, the signal of the manufacturer's original plastic was compared to the FastGene® PCR Tubes from NIPPON Genetics EUROPE.

#### Method

- PCR tubes
- 1) FastGene® 0.2 ml PCR tubes with flat caps (Cat. No.: FG-021F)

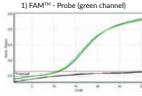
2) Original 0.2 ml Rotor-Gene® tubes (Cat. No.: 981005)

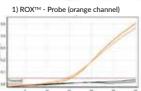
- qPCR Instrument
- QIAGEN® Rotor-Gene® Q Mdx 5plex



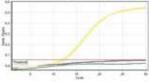
- Probes-labels
- 1) FAM™ (green channel) - Reporter primer designed to detect specific EcN plasmids 2) HEX
  - (yellow channel) Reporter primer designed to detect specific EcN plasmids
- 3) ROX™ 4) Cy5 (red channel)
- (orange channel) Reporter primer designed to detect specific regions in the EcN genome - Reporter primer as a positive PCR control and designed to detect
  - common enterobacteriae sequences

## Results





2) HEX-Probe (vellow channel)



2) Cy5 - Probe (red channel)



Dr. Birgit Klinkert:

The lid of the FastGene® 0.2 ml PCR tubes are different from the original. Nonetheless, the lock mechanism of the Rotor-Gene® Q Mdx worked perfectly with them. The fluorescence of the probes in the reaction are measured at the tip of the tubes. We can recommend to replace the original tubes for the here tested fluorescent probes without any restriction.

# \$\overline{\ove

## For capping and decapping

The CapEasy was created to keep your daily labwork simple. Sealing or removing the caps on 8-well and 12-well PCR strip tubes can often lead to loss of sample, cross contamination, and sore fingers.

This can be avoided by using the CapEasy Tool. The uniformly distributed pressure ensures perfect sealing of all wells regardless if domed or flat lids are used. When access to the contents of the strip tubes is needed, removing the lids is done in a single smooth motion without the chance of knocking the sample out of the tube.



## PCR tube recommendation

0.1 ml PCR 8-well strips and flat cap strips Cat. No.: FG-017FC

0.2 ml PCR 8-well strips and flat cap strips (120)

Cat. No.: FG-016FC

0.2 ml PCR 8-well strips and domed cap strips

Cat. No.: FG-016DC

# Ordering information

Cat. No.	Product
FG-CDC02	FastGene® CapEasy

# Free Sample?

You would like to test our Lab Plastic, e.g. our Filter Tips or our PCR tubes? No problem! Just give us a call or write us an email and get your free sample very soon.



**L** +49 2421 554960



info@nippongenetics.de

 info@nippongenetics.de

# **SFast Gene™** PCR Plates



- Compatible with most thermal cyclers
- Raised well rims to avoid cross contamination
- Free of RNase, DNase and human genomic DNA
- Compatible with heat sealing foils

# **Guaranteed quality**

The FastGene® PCR Plates are manufactured and tested for compatibility with leading manufacturers thermal cyclers. Due to the thin-walled design optimizing heat transfer, the reaction volume becomes realibly tempered.

All FastGene® PCR plastic products are manufactured by using ultra pure polypropylene. Proteins are not able to bind to the surface. Because of a very stringent QC procedure the "batch-to-batch" reproducibility of all PCR Plates is extremely high.

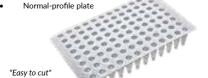
## DNA adsorption test with the FastGene® 96-well PCR plate (FG-170225) Method: Comparison of the DNA concentration after the incubation of different temperatures (4°C, 37°C, 65°C) compared to control DNA amount before incubation. DNA concentration was determined using qPCR quantification. 34 32 Before incubation 30 After incubation (30 min at 4°C) 28 Before incubation 26 After incubation (30 min at 37°C) 24 Before incubation 22 After incubation (30 min at 65°C) 20 ng / reaction Result/Conclusion: The incubation tests under all conditions showed no significant decrease of DNA concentration after incubation in comparison with the control DNA amount. This means that the FastGene® plastic shows no binding of DNA.

# **SFastGene™** PCR Plates



#### FastGene® 96-well Plate

- Non-skirted





Cat. No.: FG-170225 (25 plates)

#### FastGene® 96-well Plate

- Non-skirted
- Low-profile plate Stable Design



Cat. No.: FG-170350 (50 plates)

"Very high plate stabiliy through cross-connected wells"

## FastGene® 96-well Plate

- Semi-skirted
- Normal-profile plate



Cat. No.: FG-190250 (50 plates)

#### FastGene® 96-well Plate FROSTED ABI® style

- Semi-skirted
- Normal-profile plate
- Frosted plastic



Cat. No.: FG-200250 (50 plates)

#### FastGene® White 96-well Plate Roche® style

- Semi-skirted
- Low-profile plate
- For LightCycler™



Cat. No.: FG-210250 (50 plates)

#### FastGene® Fast 96-well Plate

- Semi-skirted
- Low-profile plate



.....

Cat. No.: FG-03890-50 (50 plates)

#### FastGene® 96-well Plate

- Full-skirted
- Low-profile plate



Cat. No.: FG-180250 (50 plates)

#### FastGene® 384-well Plate

- FastGene® Silicon sealing mat (soft)
- Full-skirted

- Volume 50 µl
- For ABI 7500 FAST



Cat. No.: FG-300150 (50 Plates) Cat. No.: FG-3110MS (10 Mats soft)

# **SFast Gene**™ PCR adhesive seal

- 138 x 79 mm (with edge), 118 x 79 mm (without edge)
- Suitable for Real-Time PCR applications
- Suitable for PE, PS and PP plates
- Highest quality prevents evaporation during PCR or storage
- Peelable without sticky residue after the seal is peeled
- End tabs for easy removal
- Resistant to DMSO
- Can be used at temperatures from -80 °C to +120 °C



Cat. No.: FG-93AC2 (100 sheets)

# Ordering information

Cat. No.	Product	Content
FG-93AC2	FastGene® Adhesive PCR Foil	100 sheets
FG-93AF	FastGene® Adhesive Seal Aluminium	100 sheets



Cat. No.: FG-014 (500 Units)

- Frosted lid and frosted side writing surface
- Graduations every 500 µl
- Thumb-friendly beveled lip, easy to open and close
- Autoclavable when open
- Compatible with all common micro centrifuges

# சார் சிர் சிர் பிர்கள் இது பிர்கள் இது

- Frosted lid and frosted side writing surface
- Graduations every 100 µl
- Thumb-friendly beveled lip, easy to open and close
- Autoclavable when open
- Compatible with all common micro centrifuges



Cat. No.: FG-015 (500 Units)

# Free Sample?

Just give us a call or write us an email if you would like to test our Lab Plastic and receive a free sample!



+49 2421 554960



info@nippongenetics.de

# **SFastGene™** Aluminium Rack





- Made in Germany from high quality aluminium
- Perfect for cooling tubes and plates on ice
- Holds semi-skirted or non-skirted 96 well plates
- Holds 6 reaction tubes and 8 PCR tubes

# Keep PCR plates and tubes cool and secure

The FastGene® Aluminium Rack is a high quality rack holding 1x 96 well PCR plate (semi or non skirted), for reaction tubes and 8x PCR tubes. The rack was designed for convenient cooling of PCR and qPCR plates as well as all the necessary reaction components. The premium quality aluminium block keeps plates and tubes cool and secure on ice. The high thermal conductivity of aluminium allows reliable, fast and uniform cooling of plates and tubes, which is especially important for sensitive samples, such as PCR enzymes. Keep your samples safe and secure on ice and improve your results - no more tipping over tubes in the ice box and getting ice contaminations.



The FastGene® Aluminium Rack conveniently and securly holds semiskirted or non-skirted 96-well plates - also at room temperature.



The high quality aluminium block is the perfect tool to keep sensitive samples cool and secure on ice.

Cat. No.	Product	Content
FG-AR01	FastGene® Aluminium Rack	1x Rack for 1x 96 Well plate, 6 reaction tubes and 8 PCR tubes

# **G**Fast Gene™ Cryo Tubes



- Cryopreservation of cells and tissues
- Temperature resistant: -196°C to +121°C
- Separate and unremovable 2-D barcode
- Wide opening for easy tissue storage

# Cell and tissue cryopreservation

The FastGene® Cryo Tubes are the optimal solution for a safe and long storage of cryopreserved samples. Three different sizes with 0.5 ml, 1 ml and 2 ml volume, the choice of an external or internal lid and the possibility of introducing a FastGene® 2-D barcode offer the right tube for every demand. The FastGene® 2-D barcode inserts are separately available and once attached to the tube they are so firmly embedded that they can't be lost. The Cryo Tubes are designed self-standing and with an extra protection against leakage.

## Automation friendly by using SBS format

The FastGene® Cryo Racks were designed to be automation friendly. This is ensured by the SBS format, which is widely used in an automated environment.



The FastGene® Cryo Tubes were designed to accomodate different volumes. The large tubes are able to store 2 ml, while the smaller tubes can handle a volume of 1 ml or 0.5 ml. All FastGene® Cryo Tubes have a perfect inlay that enables FastGene® 2-0 barcode insert to be introduced.



The FastGene® Cryo Racks are SBS format compatible.

# \$ Fast சேடா Cryo Tubes

## Cryo Tubes with external lid



500 tubes (20 bags of 25 tubes) Cat. No.: FG-CRY-05S

0.5 ml Cryo Tubes with external lid



1.0 ml Cryo Tubes with external lid 500 tubes (20 bags of 25 tubes)

Cat. No.: FG-CRY-10S



2.0 ml Cryo Tubes with external lid 500 tubes (20 bags of 25 tubes)

Cat. No.: FG-CRY-20S

# Cryo Tubes with internal lid



0.5 ml Cryo Tubes with internal lid 500 tubes (20 bags of 25 tubes)

Cat. No.: FG-CRY-In-05S



1.0 ml Cryo Tubes with internal lid 500 tubes (20 bags of 25 tubes)

Cat. No.: FG-CRY-In-10S



2.0 ml Cryo Tubes with internal lid 500 tubes (20 bags of 25 tubes)

Cat. No.: FG-CRY-In-205

## Cryo Racks



Cryo Tube Racks in SBS format for 0.5 ml (10 racks, 40 tubes per rack)

Cat. No.: FG-CRY-05RC



Cryo Tube Racks in SBS format for 1.0 ml (10 racks, 40 tubes per rack)

Cat. No.: FG-CRY-10RC



Cryo Tube Racks in SBS format for 2.0 ml (10 racks, 40 tubes per rack)

Cat. No.: FG-CRY-20RC



2-D Inserts 500 pcs.

Cat. No.: FG-CRY-2D



# Fast Gene™ Screw Cap Tubes



- Small packaging units prevent contamination
- High temperature resistance -80 °C to +121°C
- Compatible with 2D barcode inserts

# Screw Cap Tubes for cell storage

The storage of prokaryotic and eukaryotic cells gives the researcher the possibility to perform experiments over an extended time period and save cells that delivered experimental success. The FastGene® Screw Cap Tubes were developed to tolerate very high temeperature fluctuations.



The FastGene® Screw Cap Tubes are available in 5 different cap colours and 2 different volumes. Dark tubes for light protection are also available.

The 2D barcodes enable easy tracking of stored tubes.

# Perfect design - Japanese precision

The FastGene® Screw Cap Tubes were designed for different storage volumes. The large tubes are able to store 2 ml, while the small tubes have a volume of 0.5 ml. Both versions of the FastGene® Cell Storage Tubes have a convex inlay that enables the introduction of FastGene® 2D barcode inserts.



The FastGene® Screw Cap Tubes come in small packaging units (25 tubes per bag) to prevent contaminations.



The FastGene® Tube Racks are SBS format compatible.

# **G** Fast Gene™ Screw Cap Tubes

# **DNA** adsorption test

#### Purpose:

FastGene® Screw Cap Tubes (0.5 ml) were tested for their DNA adsorption over the cryopreservation.

#### Method:

Measurement of DNA concentration in FastGene® Screw Cap Tubes before and after cryopreservation.

- 1: Human genomic DNA was 20-fold diluted in order to create a 250 μl solution with a concentration of 5.0 ng/μl.
- 2: Each time 50 µl of this DNA were dispensed to 0.5 ml Screw Cap Tubes. As result, 5 mother tubes were created.
- 3: For the measurement of DNA concentration in the mother tubes after dispensing, a Qubit® was used.
- $4:10~\mu l$  from each mother tube were dispensed to three daughter tubes.
- 5: After dispensing the daughter tubes were stored for 24 h at 4°C.
- 6: The concentration of each daughter tube (5 x 3 daughter tubes = 15 tubes) was measured with Qubit®.
- 7: The mother tubes' DNA concentration was determined as 100%.

DNA conc. mother tube [ng/µl]	DNA conc. daughter tube [ng/μl]	DNA conc. changing [%]
5.08	5.18	
	5.22	101.84
	5.12	

#### Results/Conclusion:

The DNA concentration in the FastGene® 0.5 ml Screw Cap Tubes shows little or no change, so that the tubes could be used without problems for DNA experiments.

Cat. No.	Product	Description	Сар	Content
FG-SCT05-S	0.5 ml Screw Cap Tubes	clear tube, natural colour cap, sterilized		500 tubes (20 x 25 tubes)
FG-SCT05-RS	0.5 ml Screw Cap Tubes	clear tube, red colour cap, sterilized		500 tubes (20 x 25 tubes)
FG-SCT05-GS	0.5 ml Screw Cap Tubes	clear tube, green colour cap, sterilized		500 tubes (20 x 25 tubes)
FG-SCT05-BS	0.5 ml Screw Cap Tubes	clear tube, blue colour cap, sterilized		500 tubes (20 x 25 tubes)
FG-SCT05-YS	0.5 ml Screw Cap Tubes	clear tube, yellow colour cap, sterilized		500 tubes (20 x 25 tubes)
FG-SCT05-SHS	0.5 ml Screw Cap Tubes	dark tube with light protection, sterilized		500 tubes (20 x 25 tubes)
FG-SCT20-S	2 ml Screw Cap Tubes	clear tube, natural colour cap, sterilized		500 tubes (20 x 25 tubes)
FG-SCT20-RS	2 ml Screw Cap Tubes	clear tube, red colour cap, sterilized		500 tubes (20 x 25 tubes)
FG-SCT20-GS	2 ml Screw Cap Tubes	clear tube, green colour cap, sterilized		500 tubes (20 x 25 tubes)
FG-SCT20-BS	2 ml Screw Cap Tubes	clear tube, blue colour cap, sterilized		500 tubes (20 x 25 tubes)
FG-SCT20-YS	2 ml Screw Cap Tubes	clear tube, yellow colour cap, sterilized		500 tubes (20 x 25 tubes)
FG-SCT20-SHS	2 ml Screw Cap Tubes	dark tube with light protection, sterilized		500 tubes (20 x 25 tubes)
FG-SCR-RC	Tube Rack	Empty rack (red)		10 Racks for 48 tubes
FG-SCR-2D	2D Inserts	2D barcode Inserts		500 pcs.

# 

## Accessories for high-throughput applications

Our FastGene® High-Throughput Plastic includes deep-well plates, elution plates and tip comb plates, compatible with KingFisher™ automated sample purification systems. The accessories are used for the purification of nucleic acids with the help of automatic magnetic separation in high throughput.

FastGene® 96 Deep-well Plate for King Fisher™



- Deep well plates developed for magentic-bead based, automated nucleic acid purification workflows
- High quality polypropylene (PP)
- No nucleic acid or protein adherence
- Excellent recovery of magnetic beads due to welldesign
- Well shape: Deep wells (up to 2.2 ml storage volume per well), square wells, V-bottom

# Ordering information

Cat. No.	Product	Content
FG-250150	FastGene® 96 Deep-well Plate for KingFisher™	50 pcs

#### FastGene® 96-well Tip Comb for King Fisher"



- Tip combs for magnetic separation platforms
- Serve as a cover for automation platform magnets and prevent sample carryover during purification procedure
- High quality polypropylene (PP)
- No nucleic acid or protein adherence
- Excellent recovery of magnetic beads due to welldesign
- Size: 127.2 mm x 85.2 mm x 43.9 mm
- Well shape: round wells, V-bottom

# Ordering information

Cat. No.	Product	Content
FG-2502100	FastGene® 96-well Tip Comb for KingFisher™	100 pcs

### FastGene® 96-well Elution Plate for King Fisher™



- Elution plates developed for magnetic-bead based, automated nucelic acid purifiaction workflows
- Low binding Medical-grade quality polypropylene (PP)
- No nucleic acid or protein adherence
- Size: 127.5 mm x 85.35 mm x 15.1 mm
- Well shape: 0.5 ml, square wells, V-bottom

Cat. No.	Product	Content
FG-250350	FastGene® 96-well Elution Plate for KingFisher™	50 pcs



# Free Sample?

You would like to test our Lab plastic? No problem! Just give us a call or write us an email and get your free sample very soon.



info@nippongenetics.de

 info@nippongenetics.de

www.nippongenetics.eu

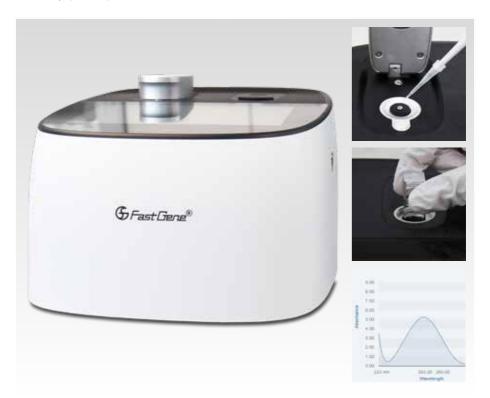




Photometer - NanoSpec / NanoView	P. 144
PCR Cycler - Ultra Cycler	P. 150
Centrifuges	P. 154
Mini Dry Bath	P. 160
Vortexer	P. 161
Tissue Grinder	P. 162

# \$ Fast சோட் NanoSpec Photometer

Powerful, reliable, intuitive



- Best device performance
  Microvolume drop and cuvette reader
- Powerful Full analysis (190 - 850 nm)
- Highest reliability 20+ measurement modes
- Convenient usage
  Simple and intuitive menu
- Cost-saving
  Affordable stand-alone device

# The All-round UV-Vis Spectrophotometer

The FastGene® NanoSpec is a powerful, all-round UV-Vis spectrophotometer characterised by highest user-friendliness.

The integration of a microvolume drop reader and a cuvette reader enable the use of small sample volumes and large sample volumes, respetively.

Over 20 preset measuring modes operating within a highend full spectrum analysis (190 - 850 nm) allow fast and easy sample analysis.

Versatile applications include quantitative measurements of nucleic acids, proteins, protein assays or bacterial cultures.

Easy performance of the FastGene® NanoSpec is ensured by an integrated control unit with a simple to use software, navigated by a glove-compatible touch-screen.

Cat. No.	Product
FG-NP01	FastGene® NanoSpec Photometer

### \$ Fast போட் NanoSpec Photometer

Powerful, reliable, intuitive

SPECIFICATIONS	
Light source	Xenon Flash Lamp
Wave length spectrum	190 - 850 nm
Measurement time	< 8 Sec
Spectral Resolution	1.0 nm (FWHM at Hg 253.7 nm)
Minimum Sample Size	1 μL (microvolume mode)
Cuvette Center Height	15 mm
Connectivity, Data storage	4 x USB Ports, Ethernet, RS-232, 32 GB Internal storage
Integrated power supply	100-240V, 4 A (50/60 Hz) automatic voltage sense, standard IEC Inlet plug
Display	7-inch widescreen 1200 x 800 HD colour touch display
Footprint (D x W), Weight	29 x 21.6 cm, 3.0 kg

#### Testimonial from our R&D

"The FastGene $^{\odot}$  NanoSpec photometer is characterized by its simple and intuitive operation system. The desired measurement protocol can be found quickly using the touchscreen and the good menu navigation.

The Droplet has an automatic mode, which means that many samples can be measured very quickly without having to make settings again."



**Dr. Manuel Franke**Head of Research and Development
NIPPON Genetics EUROPE





The light weight of the FastGene® NanoSpec combinded with a small footprint of roughly a DIN A4 page size, make it a compact and uncomplicated yet powerful stand-alone device, even with limited benchtop space.

### \$ Fast போட் NanoSpec Photometer

Powerful, reliable, intuitive

#### Two sources - One instrument

The FastGene® NanoSpec comes with a microvolume drop reader and a cuvette reader. You have the possibility to easily measure smallest sample quantities or large sample quantities.



Two sources: microvolume (µL) drop reader and cuvette reader.

FastGene® NanoSpec comes with both reading modes so that you can

#### Intuitive and easy to use Software

The software of the FastGene® NanoSpec was designed having the user in mind. The programs are easy to identify and straight-forward to use.

The device software comprises over 20 preset measurement modes organized in distinct tabs for versatile applications. The creation of new and personalized programs is easy and self-explanatory.



Nucleic Acids

The Nucleic Acid tab allows you to choose the measurement mode for different types of DNA and RNA. A customized Nucleic Acid measurement mode can also be applied.



Protein UV

The Protein UV tab supports 8 modes at 280 nm with different measurement factors or customizable input of extinction coefficients.



Protein Assay

The Protein Assay tab allows quantification of protein concentration at a specific wavelegth, after staining with colorimetric reagents. The protein concentration can be measured after generating a standard curve.



More

The More Application tab supports various measurement modes used in general-purpose UV-Vis spectrophotometry.

Cat. No.	Product	
FG-NP01	FastGene® NanoSpec Photometer	

Product Highlight



# Talk to the experts and get a free product demonstration

Finding the perfect Spectrophotometer can be difficult. We can help you! Just arrange an appointment with us and get a product demonstration.



+49 2421 554960



info@nippongenetics.de

www.nippongenetics.eu

### **SFast** Gene™ NanoView Photometer

Compact, easy to use, affordable



- Best device performance
  Microvolume drop and cuvette reader
- Specialist
  For DNA, RNA, Proteins and OD600
- Convenient User Interface
  10 preset measurement modes
- Smart
  Compact, silent and easy to use
- Cost-saving
  Affordable stand-alone device

Drop reader or cuvette reader - Why not both in one device?

Both measurement modes are smartly integrated under one lid and allow you to take precise measurements of DNA, RNA, protein or OD600.

#### The Compact Specialist

The FastGene® NanoView is our smart and compact specialist photometer designed for precise applications.

The microvolume drop reader and cuvette reader are both integrated under one lid making the FastGene® NanoView ultra-small, silent and an absolute lightweight.

Ten preset modes ensure easy quantitative analysis of nucleic acids (260 nm), proteins (280 nm) and OD600 cuvette measurements.

Easy performance of the FastGene® NanoView is ensured by an integrated control unit with a simple to use software, navigated by a glove-compatible touch-screen.



### **SFast** Gene™ NanoView Photometer

Compact, easy to use, affordable

#### Convenient and optimized User Interface

The software UI of the FastGene® NanoView was designed for ease of use with large and self-explanatory icons.

The specialist device comes with 10 preset measurement modes enableing convenient DNA, RNA, protein or OD600 analysis.



Keeping it clean? Easy!

Not only the software is easy to operate. The microvolume drop reader is quickly cleaned with a dust-free laboratory tissue, also between multiple sample measurements, for effortless device maintenence.



Main Menu tab composition
The main menu enables easy navigation through distinct measurement modes with intuitive icons.



Favorites Screen

The preferred and often used measurement modes can be stored in the Favorites tap for fast navigation and conveient use.

SPECIFICATION	
Light source	LEDs
Wave lengths	260 nm, 280 nm / 600 nm (Cuvette) / 360 nm (Baseline)
Measurement time	< 10 Sec
Minimum Sample Size	2 μL
Cuvette Center Height	15 mm
Connectivity, Data storage	2 x USB ports, USB-B, RS-232, 8 GB internal storage
Integrated power supply	100-240V, 4 A (50/60 Hz) automatic voltage sense, standard IEC Inlet plug
Display	4.3-inch 480 x 272 colour touch display
Footprint (D x W), Weight	$19 \times 14.5$ cm, $1.4$ kg

Cat. No.	Product
FG-NP02	FastGene® NanoView Photometer

### \$*F்±ட்டோட*் Ultra Cycler

Efficient, intuitive, smart



- Very fast ramp rates for a quick PCR
- 96-well PCR instrument with a gradient
- Touchscreen with a very easy-to-use software
- Compatible with most PCR tubes and 96-well microplates

#### It's simple - Affordable gradient PCR

A thermal gradient of up to 24°C can be precisely generated in the FastGene® Ultra Cycler. The temperature of each well is stable and reliably repeatable from cycle to cycle. The fully adjustable heated lid is compatible with a wide range of 0.2 ml PCR tubes (standard and low profile) and 96-well PCR microplates (fully-skirted, semi-skirted, and non-skirted). With a huge touchscreen interface and simple programming, this system is convenient and easy-to-use. As a small and robust device, the FastGene® Ultra Cycler is perfect for any lab environment by providing years of reliable amplification!



# \$*Fast:போட* Ultra Cycler

Efficient, intuitive, smart



SPECIFICATIONS	
Gradient	24°C over the whole block width
High temperature range, accuracy and resolution	Temperature range: $4^{\circ}\text{C}$ - $99^{\circ}\text{C}$   Temperature accuracy: $\pm$ 0.25°C Temperature resolution: 0.1°C increments
Very fast ramp rates	Heating rate: 7°C / per second Cooling rate: 5°C / per second
Compatible	0.2 ml tubes or strip tubes with flat or domed caps 96- well high or low skirted plates with strip caps, adhesive seal
Condensation control	Automatic utilising applied pressure heated lid
Heated lid	Heated lid with a temperature range of 60 °C - 115°C
Compact design	Dimensions (H x D x W): 19 x 28.5 x 18 cm Weight: 5.5 kg
Integrated power supply	100-240V, 4 A (50/60 Hz) automatic voltage sense, standard IEC Inlet plug
Huge touchscreen	7-inch widescreen colour touch display

#### **Customer Testimonial**

"We own a Nippon FastGene® Ultra Cycler since December 2017. The Cycler is used every day. So far, the device works very reliable. It is relatively small, very quiet and has very short run times, because of fast heating and fast cooling of the samples. The handling and programming of the cycler is very simple with a clear menu navigation. Everyone can save own programs with a personal avatar"



**Researcher** Institute of Molecular Botany, University Ulm, Germany



Cat. No.	Product	
FG-TC01	Gradient UltraCycler PCR Thermocycler with touchscreen	

### \$*Fast'ப்பட* Ultra Cycler

#### The gradient advantage

The gradient function allows you to optimise your reactions. Discover the best annealing temperature over a range of 24°C. The block system is designed for the use of 96 individual PCR tubes (0.2 ml), 12 PCR 8-well strips (0.2 ml) or 96-well PCR plates (0.2 ml). The UltraCycler combines the latest electronics and peltier technology with an remarkably high operating comfort

#### Quickstart with Albert



Albert enables the user to configure easy to moderate complexity profiles in just a few moments. Every step of a routine PCR is available (even a 1-Step RT-PCR can be performed

#### Touchscreen graphical user interface

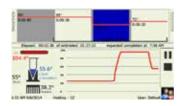
A high performance graphical processor with a large 7 inch, vivid colour touchscreen display allows for an easy run setup and monitoring. The powerful yet intuitive software makes the creation of even the most complex thermal profiles a child's play.

#### Heated lid evaporation control

The FastGene® Ultra Cycler employs an applied pressure heated lid design to keep the air contained within the tube hotter than the reaction volume. This causes any evaporation to condense back into the cooler reaction liquid, thereby eliminating the need for an oil or wax condensation overlay.

#### USB connectivity

A front USB port allows for fast, easy file transfer to a USB memory stick enabling the sharing of thermal profiles between instruments and users. The use of a USB mouse is also supported.



Watch your PCR
The software allows you to watch all steps during the PCR cycle with precise temperature information.

Exact temperatures
The temperature gradient selected with the Albert PCR assistant

In temperature gradient selected with the Albert PCR assistant is used to calculate the exact temperature in each lane. Hence, a better determination of the optimal temperature is possible.



Genious configuration

The Albert PCR assistant enables the user to configure easy to moderate complexity profiles in just a few moments. All the thermal steps which occur in a typical profile are included and the parameters and can be adjusted in just a few clicks.



The user accounts section

Allows to create up to 99 user profiles each with dedicated file storage directory and personalized Icon. When a user is selected, thermal profiles will be loaded or saved to a directory specific to that user providing easy navigation of the data.

Cat. No.	Product
FG-TC01	Gradient UltraCycler PCR Thermocycler with Touchscreen



# Talk to the experts and enjoy a free product demonstration

Finding the perfect Thermal Cycler can be difficult. We can help you! Just arrange an appointment with us and enjoy a product demonstration.



✓ info@nippongenetics.de

www.nippongenetics.eu

### \$ F கர் சேர்ச் Mini Centrifuge



- Four different colours with a compact design
- Supplied with standard microtube, slide and strip tube rotor
- Ideal for quick spin down and microfiltration

The FastGene® Mini Centrifuge adaptors and rotors.

#### The ideal lab companion

The FastGene® Mini Centrifuges come in four different colours and are supplied with three rotors. The first rotor is designed to centrifuge up to six individual 1.5 ml plastic micro centrifuge tubes. It will also spin down 0.5 ml tubes and 0.2 ml tubes with the adapters supplied with the unit. The second rotor can load two 8-well strips (tube capacity 0.2 ml). The rotors are designed for applications requiring relatively low g-forces, such as microfiltration, cell separation and quick spin downs of liquid from the tube lids and tube walls.



The rotors of the Mini Centrifuges can be easily replaced.

# \$ Fast பேடாம்™ Mini Centrifuge





FastGene® Mini Centrifuge in Pink



FastGene® Mini Centrifuge in Green



FastGene® Mini Centrifuge in Blue



FastGene® Mini Centrifuge in Red

SPECIFICATIONS		
Four different colours	<b>~</b>	Pink, Blue, Green and Red
Three different rotors included	<b>✓</b>	Standard angle rotor for 6x 1.5/2.0 ml tubes   Slide rotor 0.2 ml strip tube rotor
Adaptors included	<b>~</b>	6x adaptors for 0.5 ml tubes 6x adaptors for 0.2 ml tubes
High speed	<b>✓</b>	Centrifugal force: 2,000 x g Speed: 6000 rpm
Compact design	<b>~</b>	Dimensions (H x D x W): 11.8 x 17.5 x 14.8 cm
Integrated power supply	<b>~</b>	100-240 V ~ 0.5 A

Cat. No.	Product	Content
NG002P	FastGene® Mini Centrifuge (Pink)	Pink Mini Centrifuge   3 rotors   6 adaptors for 0.2 ml and 0.5 ml tubes
NG002B	FastGene® Mini Centrifuge (Blue)	Blue Mini Centrifuge   3 rotors   6 adaptors for 0.2 ml and 0.5 ml tubes
NG002G	FastGene® Mini Centrifuge (Green)	Green Mini Centrifuge   3 rotors   6 adaptors for 0.2 ml and 0.5 ml tubes
NG002R	FastGene® Mini Centrifuge (Red)	Red Mini Centrifuge   3 rotors   6 adaptors for 0.2 ml and 0.5 ml tubes

# \$ Fast செட்் High Speed Mini Centrifuge



#### Fast and silent

The FastGene® High Speed Mini Centrifuge comes with a microrotor with a capacity of 12 tubes and can centrifuge up to 12,300 xg respectively 13,500 rpm. The centrifuge shows a very compact design. After centrifugation, the door will be opened automatically. The high quality of the centrifuge is supported by the fact that all mechanical parts are made of strong steel components. The air flow system guarantees that the noise during centrifugation is very low. The included rotor is autoclavable and a rotor for 8-well PCR strips is also available separately.





The standard rotor of the FastGene® High Speed Mini Centrifuge can load 12x 2 ml or 1.5 ml tubes. The optional PCR rotor (Cat.No.: NG004) is suitable for 0.2 ml single tubes or 4x 8-well strips.

SPECIFICATIONS				
High speed	Standard rotor	Centrifugal force: 12,300 x g Speed: 13,500 rpm	PCR strip rotor	Centrifugal force: 1,850 x g Speed: 6,000 rpm
High capacity	Standard rotor	12x 2 ml or 1.5 ml tubes	PCR strip rotor	4 x 8well PCR strips
Time control		Pulse or timed ≤ 30 min Blue LCD display		
RPM/RCF conversion	Easy conversion to RPM/RCF			
Very silent	Less than <56 dB			
Compact size		Dimensions (H x D x W): $14.5$ x $24.5$ x $20.8$ cm Weight: $4.4$ kg		
Integrated power supply	220V, 50-60 Hz			

Cat. No.	at. No. Product Content	
NG003	FastGene® High Speed Mini Centrifuge	Centrifuge comes with the standard rotor
NG004	PCR rotor	Optional rotor for 0.2 ml single tubes or 4x 8-well strips

# **⑤** Fast Gene<sup>™</sup> Plate Centrifuge





Plate Centrifuge with two plate carriers



Convenient, silent and easy-to-use



Spinning down liquids in 96- and 384-well plates

#### Spin down your plates - the easy way

Our FastGene® Plate centrifuge was designed for quick and gentle centrifugation of various plate and tube types. Its handling could not be easier! All you need to do is set the spinning time (up to 10 min) on the illuminated display and you are good to go. The compact footprint, simple handling and affordable price make the FastGene® Plate Centrifuge the perfect alternative to more complex and expensive high-speed plate centrifuges.

#### Optimize your results

The FastGene® Plate centrifuge was made for reliable and reproducible results. It is important to have the entire reaction volume present in the bottom of the wells and to eliminate the chance of cross-contamination with other wells. Droplets or condensation on the sides of the wells can cause assay failure due to inadequate volumes or reaction separation. A reliable and properly-designed plate-centrifugation should therefore be a part of every GLP-protocol.

#### Centrifugation was never more silent

We know that everyday laboratory life can be loud and stressful. The FastGene Plate Centrifuge is equipped with an extra silent rotor and extra tight lid sealing. It is the perfect device to support a more silent lab environment - keep your work focus high and get better results!

## **SFast** Gene™ Plate Centrifuge

#### Convenient and versatile

The FastGene® Plate Centrifuge is very versatile: It comes with 2 plate adapters, making it compatible with all types of 96-well plates such as full-skirted, half-skirted or non-skirted plates. These adapters can also be used for individual reaction tubes and 8-well strips, making it the perfect centrifuge for spinning down liquids in life science laboratories. Even 384-well plates (full-skirted) are no problem for the plate centrifuge.

#### Optimize your high throughput applications

Precise high throughput applications need reliable equipment. For genuine analysis it is extremely important to have the entire reaction volume in the bottom of the well, without any droplets on the wall side. The FastGene Plate Centrifuge helps to spin down your plates for:

- · PCR and qPCR applications
- ELISA
- Colorimetric or fluorescence based high throughput screenings
- Next Generation Sequencing applications
- Bacterial growth cultures

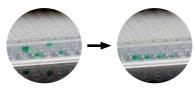
Prevent assay failures with a reliable and properly designed plate centrifuge.

#### Easy to clean

Spilling biological material can be a major source of infection. The rotors of the centrifuge are easily removable, enabling complete cleaning of the internal area of the centrifuge.



Adapter plates for semi- and non-skirted 96-well plates as well as single reaction tubes or 8-tube strips.



Before and after centrifugation of a full-skirted 384-well plate.

SPECIFICATIONS				
Compatible for many well plates	<b>✓</b>	96-well plates (full-skirted, semi-skirted and non-skirted) 384-well plates (full-skirted)		
High speed	<b>✓</b>	Centrifugal force: 480 x g Speed: 2200 rpm		
Compact size	<b>✓</b>	Dimensions (H x D x W): $14 \times 36 \times 29$ cm Weight: $1.3 \text{ kg}$		
Adapter plates included	<b>✓</b>	Adapter plate for semi-and non-skirted 96-well plates Adapter plate for single reaction tubes or 8-tube strips		
Integrated power supply	<b>✓</b>	200-240 V, 50-60 Hz Additional version with 110 V, 50-60 Hz is also available		

Cat. No.	Product	Content
NG040	FastGene® Plate Centrifuge	Plate Centrifuge, 4 x Adapters (200-240 V)
NG040US	FastGene® Plate Centrifuge (US version)	Plate Centrifuge, 4 x Adapters (110 V)



# Do you have questions about our centrifuges?

We understand that, as a scientist, you would like to test our centrifuges before buying them. That is why we offer a product demonstration online or in your lab. Just arrange an appointment with us!



info@nippongenetics.de

 info@nippongenetics.de

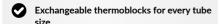
www.nippongenetics.eu

# \$ Fast செட் Mini Dry Bath Advance









#### Precise sample heating for every tube size

The FastGene® Mini Dry Bath Advance is a microprocessor controlled block heater, operated via nine setable programs. It provides excellent temperature control for a wide variety of applications and delivers accurate and reliable experimental results. Applications include sample and reaction tempering, sample heating, denaturation of electrophoresis samples, serum coagulation and many more. The compact, yet powerful design fulfills all your incubation needs. The mini dry bath is used with exchangeable thermo blocks for different tube sizes, ranging from 0.2 ml PCR tubes to 50 ml culture tubes.

SPECIFICATIONS	
Temperature Control Range	room temp. (+5 °C) - ~100 °C)
Heating Time (20 °C to 100 °C)	≤ 18 min
Temperature Accuracy Discrepancy	± 0.3 °C
Time Setting	0-99 h, 0-99 min, 0-99 sec
Max. temperature	100 ℃
Cooling Time	Natural Cooling
Dimensions (H x D x W)	13.6 x 16 x 11 cm
Cover	With Transparent Plastic Cover

### \$ Fast சொ்ச Mini Dry Bath Advance



FastGene® Mini Dry Bath Advance, equipped with the 15x 1.5 ml tube thermoblock. Nine different programs can be set, including three temperature steps for each program.



Seven thermoblocks for common tube sizes are available for the FastGene® Mini Dry Bath Advance, ranging from 0.2 ml PCR tubes to 50 ml culture tubes.

#### Ordering information

Cat. No.	Product	Content
NG020A	FastGene® Mini Dry Bath Advance	Mini dry bath main device
NG025A	Metal thermo block	For 32x 0.2 ml reaction tubes
NG029A	Metal thermo block	For 24x 0.5 ml reaction tubes
NG026A	Metal thermo block	For 15x 1.5 ml reaction tubes
NG030A	Metal thermo block	For 15x 2 ml reaction tubes
NG024A	Metal thermo block	For 12x 5 ml reaction tubes
NG027A	Metal thermo block	For 6x 15 ml reaction tubes
NG028A	Metal thermo block	For 2x 50 ml reaction tubes

### *GFastGene*<sup>™</sup> Vortexer Mini



#### Ordering information



in single tubes, falcon tubes and beakers. The adjustable rotational speed of 0-4000 rpm enables a very gentle up to vigorous shaking. The stability is ensured by a heavy base
preventing the vortexer from dancing and avoiding a slipping of the sample tube.

The FastGene® Vortexer Mini is suitable for mixing samples

Compact with a high performance

SPECIFICATIONS	
Construction material	Chemical resistant plastic
Support system	Heavy base
Operational mode	Touch
Speed setting	Analogue
Speed	0 - 4000 rpm

Cat. No.	Product	Content
VX2	FastGene® Vortexer Mini	Main Unit

### Mixy Professional Tissue Grinder



- Grinder for resuspending pellets and disrupting tissue
- Simply to use and cordless
- Homogenization of animal tissue, bones, plant tissue and food

#### Mixy homogenizes almost everything

The Tissue Grinder Mixy Professional is a motor-driven grinder for resuspending pellets or disrupting soft tissue in microcentrifuge tubes. The motor is powered by a 3.7 V battery and can be used cordlessly for up to 4 hrs.

#### The easiest way to homogenize tissue

Take the grinder out. Place the pestle on the pestle adapter. Insert the pestle into the microcentrifuge tube, and use the button to start mixing. Release the button after the mixing operation is completed.

SPECIFICATIONS	
Speed:	12,000 rpm
Successfully homogenized tissues	Animal, bacteria, plants (root and leaf) and bones
Life time of the rechargeable battery	4 hours
Dimension (H x W):	15.5 x 2.5 cm
Weight:	0.2 kg

Cat. No.	Product	Content
NG010	Mixy Professional	Tissue Grinder with lithium battery and 10 plastic pestles
NG011	Metal Pestle	Autoclavable steel pestle
NG006	Plastic Pestles	100 disposable plastic pestles 1.5 cm <sup>3</sup>

#### Application

#### Improving RNA extraction from arterial tissue

Product

Mixy Professional Tissue Grinder (NG010)

Manufacturer

NIPPON Genetics EUROPE

The following data is kindly provided by Daniel Schick, University Medical Centre in Aachen, Germany.

#### Background

Isolation of RNA from arterial tissue is difficult. Disrupting the tissue before starting the isolation of the nucleic acid can enhance the RNA yield. Here, we present the isolation of RNA performed with and without the use of the homogenizer Mixy Professional. The RNA was used to analyse the expression of metalloproteins in cardiovascular diseases.

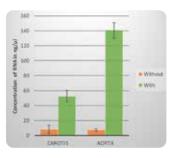
#### **Experimental Condition**

- Type and amount of tissue: Aorta (30 µg) and carotis (5 µg) isolated from mice (1 12 months old, stored at -80 °C)
- · Condition of tissue: Intact tissue morphology
- Methods:
- Homogenizing the tissue in lysis buffer using the Mixy Professional Tissue Grinder
- 2. Incubation with Proteinase K
- 3. Column based mRNA isolation
- 4. Spectrometric determination of concentration and quality
- 5. RNA quality determination using agarose gel electrophoresis
- 6. Reverse transcription
- 7. Quantification (relative) of gene expression using qPCR assays

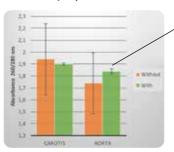


#### Results

#### Concentration of RNA



#### Quality of RNA



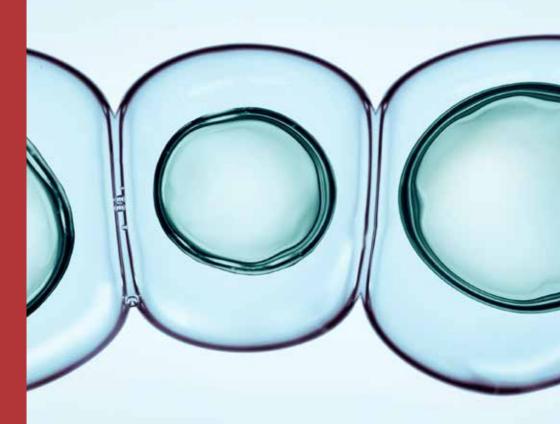
Using the Mixy Professional Tissue Grinder reduces the variability of the RNA quality immensely.



#### Customers comment

#### Daniel Schick:

The RNA yield was extensively increased by 6 - 20 fold when using the Mixy Professional Tissue Grinder. Additionally, the quality of the RNA measured spectrometrically showed considerably less variation when using the Grinder (absorbance at 260/280 nm is  $1.94 \pm 0.30$  without vs  $1.90 \pm 0.01$  with the Mixy Grinder Professional).



# **CELL BIOLOGY**



StemFit - Culture Media for Stem Cells	P. 166
Recombinant human bFGF and Activin A	P. 170
Bambanker <sup>™</sup> - Cell Freezing Media	P. 172
Cell Culture Chamber	P 177



Recommended by leading Scientist



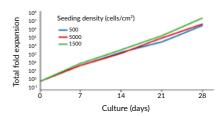
- Recommended by the nobel prize winner Shin'ya Yamanaka
- Enables single cell cloning
- Feeder-free and Xeno-free
- Less media volume needed
- Reproducible and fast replication rates

### No feeder cells and xeno-free culture medium

StemFit\* medium was developed to produce a reliable and well-defined growth condition for human stem cells. It has all necessary components for the culture of embryonic stem cells (ES) as well as induced pluripotent stem cells (ipS), it has a xeno-free composition and only contains human components. StemFit\* also eliminates the need for feeder cells. These important benefits lead to a reduction of variation in growth, and reduces concerns for contamination in the cultivation of stem cells.

#### Very reproducible growth rates

The cultivation of stem cells using StemFit® results in a very reproducible growth rate, allowing a perfect planning of experiments. No more variation due to different starting conditions caused by the natural variation when culturing on feeder cells. Analysing the morphology of stem cells cultivated in StemFit® shows that the colony shape and size are very similar to the cells grown on feeder cells.



Human 201B7 iPSCs were cultured on iMatrix-511 with StemFit® for 4 weeks without weekend feeding.



Recommended by leading Scientist





#### Less media needed

Due to the high-quality components and the ideal concentration of nutrients, the volume of StemFit\* required per plate is lower than that of conventional media, and even lower than that of other feeder-free media. For each well of a six-well plate you need just 1.5 ml, instead of 2 ml. A further 50% reduction in media consumption is the direct result of far fewer feeding steps during the week. This means that you are saving in reagents, time, and money.



The volume of StemFit® can be reduced by 25% per well. The reduced amount of media changes leads to a further volume reduction of more than 50%.

#### Free up your weekends

The cultivation of stem cells is very complicated and laborintensive, including feeding steps during the weekend. StemFit® allows a weekend free from stem cell media changes. The recommended weekly workflow minimizes the hands-on time.



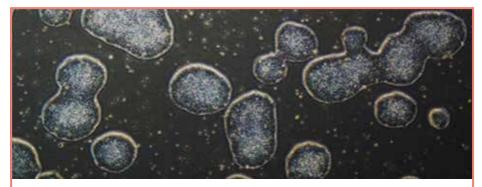
Weekend-free workflow with StemFit®. (Black circle = cell passage; Pink circle = media change (MC); white circle = maintenance-free day).

Let StemFit feed your cells while you enjoy your weekend!

#### Combined medium during the whole process

StemFit® contains no bFGF, so you can choose the best bFGF concentration for your needs. Therefore, you can use the same medium for reprogramming, cultivation and differentiation.



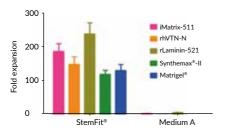


# Recommended by the nobel prize winner Shin'ya Yamanaka

"StemFit, a newly developed Xf-medium, was the best medium for hESC and hiPSC culture with rLN511E8"

#### Superior growth performance on any matrix

StemFit\* has been tested on many different matrices. As can be seen below, the fold expansion rate of cells cultured in StemFit\* is much higher when compared to Medium A from the market leading competitor.



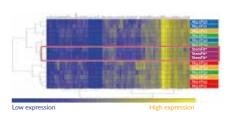
Fold expansion of human 201B7 iPS-cells, transitioned to feeder-free conditions with StemFit® or commercially available medium A. Cells were cultered for one week.

#### Start from a single cell

StemFit\* enables a superior colony forming efficiency from a single cell clone, which minimizes the effects of stress and results in reliable cells for downstream applications. Furthermore, with StemFit\* you can easily determine the efficiency of your cell production and duplicate individual clones.

#### Consistent gene expression profile

The cultivation of stem cells is very stressful for the cells. Every passage and growth period could therefore introduce unwanted changes in the genome expression profile. Hence, the CGI Catapult Institute in London investigated the genomic profile of StemFit\* after 1 passage, 3 and 5 passages and compared it to 4 commercially available media. The most consistent gene expression was obtained using StemFit\*.



The expression profile using the TaqMan ScoreCard™ assay (n=3) showed that the most consistent gene expression of after 5 passages was obtained using StemFit®.

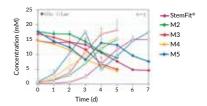


Human embryonic stem cells were dissociated into single cells and cultered with StemFit®. Cells show a normal cell morphology.



#### Less production of lactate

The production of lactate is the result of hypoxic stress. The consequences can be changes in the genome expression profile or lead to unwanted differentiation of the stem cells. The CGI Catapult Institute in London showed that there is considerably less lactate production when the cells are grown in StemFit\* (pink line with white circles).







StemFit Basic03

StemFit Basic04CT (complete type)

Single-cell culture	✓	✓
Xeno-free	✓	✓
Animal-origin free	✓	✓
Clinical research	✓	✓
bFGF	-	✓
cGMP	-	in preparation
Number of bottles	2	1

#### Reference in Literature

- Nakagawa, M. et al. (2014). A novel efficient feeder-free culture system for the derivation of human induced pluripotent stem cells. Sci. Rep., 4, 3594.
- Desai N. et al. (2015). Human embryonic stem cell cultivation: historical perspective and evolution of xeno-free culture systems. Reprod Biol Endocrinol., 13:9.
- Morizane, R. and Bonventre, J. (2017). Generation of nephron progenitor cells and kidney organoids from human pluripotent stem cells. Nature Protocols, 12 No.1.
- many others...

Cat. No.	Product	Content
Basic03	StemFit® Medium (Clinical grade)	500 ml (400 ml Liquid A, 100 ml Liquid B)
Basic04CT	StemFit® Medium Complete Type (Clinical grade, incl. bFGF)	500 ml (one bottle composition)

### Recombinant human bFGF

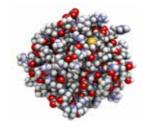


#### Take care of your stem cells

Basic fibroblast growth factor (bFGF) is a prototypic member of the fibroblast growth factor family. Proteins of this family play a central role during prenatal development, postnatal growth and regeneration of a variety of tissues, by promoting cellular proliferation and differentiation.

bFGF is a critical component for maintaining embryonic stem cells and iPS cells in culture in an undifferentiated state. Human bFGF from Ajinomoto is a bioactive protein intended for use in cell culture applications.

- Manufactured under cGMP compliant facility
- Animal-origin free
- **⊘** Great performance with StemFit®
- High purity and activity
- High batch homogeneity



Molecular structure of the basic fibroblast growth factor.

#### Ordering information

Cat. No.	Product	Content
bFGF-1mg	Basic Fibroblast Growth Factor (bFGF).	1 mg

### **Activin A**



#### Differentiate stem cells into endoderm or mesoderm cells

Activin A is a member of the TGF-beta superfamily of cytokines and is involved in a wide range of biological processes including tissue morphogenesis and repair, fibrosis, inflammation, neural development, hematopoiesis, reproductive system function, and carcinogenesis. Human Activin A is a 26.0 kDa disulfide-linked homodimer of two  $\beta$ A chains, each containing 116 amino acid residues.

Activin A is mainly used for stem cell cultivation in order to differentiate the stem cells into endoderm or mesoderm.

The perfect combination with StemFit®

Cat. No.	Product	Content
BasicAA10	Recombinant human Activin A (0.1 mg/ml)	100 μΙ, 10 μg
BasicAA50	Recombinant human Activin A (0.1 mg/ml)	500 μl, 50 μg



# Stem cell therapy moves towards the clinic

Stem cell research is leading to potential new therapies to treat disease, with several applications in clinical trails or expected to enter trials in the coming months. These new discoveries are transforming how we think about the future of medicine.

Due to its high potential, the global regenerative medicine market is likely to expand considerably in the coming years. According to a report published by Fortune Business Insights, titled "Regenerative Medicine: Global Market Analysis, Insights and Forecast, 2019-2026," the market was valued at US\$ 23,841.5 Mn in 2018. Fortune Business Insights states that the market will reach US\$ 151,949.5 Mn by the end of 2026.

### Bambanker™



- Higher survival rate
- No programmed or sequential freezing required
- Serum-free no risk of contamination
- Usable for all known cell lines

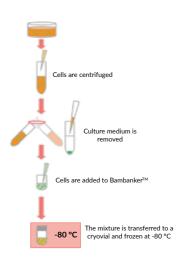
#### Long-term storage of cultured cells

Cryopreservation of mammalian cells is extremely valuable and common in biological research. Once transferred from growth media to freezing medium, the cells are usually frozen at a controlled rate and stored in liquid nitrogen or at -130 °C in a mechanical deep freezer. Although freezing a cell line is a commonly performed procedure, problems arise when suitable freezers are not available, or undefined variables are introduced by the presence of serum, extra-wash or complicated freezing algorithms.

#### Save time while saving your cells

The cell freezing media Bambanker™ permit cryopreservation of cells at -80 °C (or in liquid nitrogen), avoiding the need for an additional and expensive ultra-low freezer or time consuming and complicated controlled freezing protocols. Simply 1) harvest cells, 2) aspirate medium, 3) resuspend in Bambanker™, 4) transfer to a cryovial and 5) store at -80 °C. No programmed or sequential freezing is required! Bambanker™ is a serum-free cryopreservation medium that is delivered ready-to-use and can be kept in the refrigerator for up to two years. Convenient 20 ml bottles are available, making Bambanker™ freezing medium ideal for use by individual members of your lab.

Ready-to-use medium for preservation of cells



### Bambanker<sup>™</sup>



#### Higher number of intact cells after thawing

Bambamker™ freezing medium offers the combination of a simple protocol, leading to a high survivability. This product has been used in many labs worldwide, with a broad range of cell types. Recovery rates, even of sensitive cells, are much higher compared to regular cell freezing media. The vast majority of cell types show a survivability of more than 50% with many approaching 90% or more. Hybridoma cells can reach 100% recovery after long-term storage.

#### Serum adds variation to long-term storage

All Bambanker™ products are produced with no serum. Cryopreservation media which contain serum have an undefined composition and the disadvantage of fluctuations in recovery rates. Reproducibility of experiments with cells that were frozen in a serum-containing-medium could be affected by lot-to-lot variation of the serum. The composition and concentration of proteins and other biological molecules may vary with each batch. This can result in unexpected issues when thawing and using the cells. Every ingredient of Bambanker™ is precisely defined. Your cells will therefore behave and recover in a reproducible manner.

### Bambanker<sup>™</sup> prevents undesired differentiation

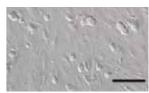
Before freezing

10

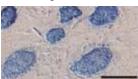
100µm

2 days after thawing

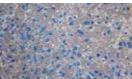


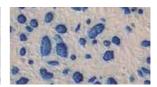


ALP staining



3 days after thawing





Cell viability and ALP staining of pluripotent stem cells. Upper row: A great number of cells are detected two days after thawing. The cells show no morphological change after thawing. Lower row: Bambanker<sup>th</sup> does not cause cell differentiation as all stem cells frozen down are still producing high levels of alkaline phosphatase, a reporter for pluripotent are cells.

#### Cells successfully frozen with Bambanker™

Bambanker<sup>TM</sup> is suitable for all known cell lines. The JCRB cell bank stores over 1,400 different cell lines with great success with Bambanker<sup>TM</sup> (take a look at the Application Note on the next page). Furthermore, there are plenty of published scientific research papers, describing Bambanker<sup>TM</sup> as the freezing medium of choice.

Cat. No.	Product	Content
BB01	Bambanker™	120 ml
BB02	Bambanker <sup>TM</sup>	5 x 20 ml
BB03	Bambanker™	20 ml



#### Application

#### Comparing Bambanker<sup>™</sup> with another cryopreservation medium for the cultivation of 1.400 different cell lines

The following data were kindly provided by Dr. Arihiro Ohara, National Institute of Biomedical Laboratories JCRB cell bank.

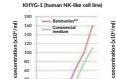
#### Background

The JCRB cell bank handles approximately 1,400 different cell lines. A low survival rate after thawing frozen cell lines (KHYG-1, KAI3, HL60, OVMANA) has let us to test Bambanker<sup>™</sup> and compare it to the previously used preservation medium for the four cell lines. The growth efficiency after thawing was compared for cells stored with the currently used commercially available preservation medium and Bambanker™.

#### Method

All cultured cells were harvested in the logarithmic growth phase. 1 ml preservation medium was added to approximately 1 x 106 cells in a storage tube. The cells were stored for 2 weeks at -80 °C. The frozen cells were thawed in an 37°C water bath and incubated at 37°C and 5% CO<sub>2</sub> in a 96-well plate. Every day the viable cell number was determined.

#### Results

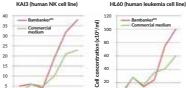


Cell lines in suspension

60

40 <u></u>

20

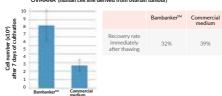


Days of cultivation





Days of cultivation



The survival rate after thawing the four cell lines (KHYG-1, HL60, KAI3, OVMANA) was very low, either with the previously used commercially available product or with Bambanker™. However, after thawing, the cell proliferation of all four cell lines was improved with Bambanker™ when compared to the previously used commercially available product.

#### Dr. Arihiro Ohara:



Customers comment

JCRB cell bank has carried out cell bank business for 30 years and we currently store 1,400 types of cell lines. [...] Due to the high number and the wide variety of cell lines, we had some problems. Some users complained that their cell lines died after thawing, resulting in unsuccessful cultivation. Especially four types of cell lines were a problem which had to be urgently improved. Therefore, we compared Bambanker™ with our currently used commercial preservation medium in a cryopreservation test. The cell lines, which were stored with Bambanker™, showed much higher cell proliferation than cells, which were stored with our currently used commercially available product. Surprisingly, with Bambanker™ we got for all four cell lines very reproducible results. [...] In the future, we will completely change to Bambanker™ in order to improve the survival rate and growth of our cells. We are thankful for resolving that long-standing problem and recommend Bambanker™ to all domestic researchers and foreign cell banks.

→ The JCRB cell bank has been using Bambanker<sup>™</sup> since 2014 for all their cell lines.

### Bambanker<sup>TM</sup> - HRM

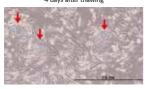


- Optimal for the cryopreservation of primate ES and iPS cells
- Made with human serum albumin
- No animal components only human albumin

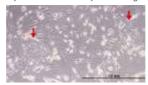
### A new hope for the cryopreservation of ES and iPS cells

Primate embryonic stem cells and induced pluripotent stem cells are extremely sensitive to cryopreservation which presents many difficulties when compared to murine cells or other cells. The slow-freezing method using DMSO has been popular for a wide variety of cell lines. Currently, the vitrification method is considered to be superior for freezing primate ES/iPS cells. Vitrification is the rapid cooling of freezing media to a glass-like crystalline state. The vitrification method requires impeccable timing, a high level of skill, and still can yield poor results. In addition, this method shows sensitivity to dry ice transportation. To address these problems, a new freezing medium called Bambanker<sup>TM</sup> HRM has been developed. What makes it so special is the removal of bovine serum albumin (which can lead to cell differentiation in some cases) and any animal-derived material (xeno-free). Both improve the storage and survivability of primate ES/iPS cells while greatly simplifying the protocol.

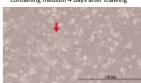
Cells cryopreserved with Bambanker<sup>™</sup> HRM 4 days after thawing



Cells cryopreserved with vitrification freezing preservation solution 4 days after thawing



Cells cryopreserved with 10% DMSO containing medium 4 days after thawing



IPS cells (20187) were cryopreserved in Bambanker<sup>™</sup> HRM, in a conventional vitrification medium or in a 10% DMSO/culture medium. After 3 days, the cells were thawed, according to protocol, and then plated. In case of 10% DMSO containing medium only small colonies could be recovered what suggests a small survival rate, while with Bambanker<sup>™</sup> HRM it was possible to recover large colonies with almost the same size as achieved with vitrification freezing preservation solution. These results indicate that the same storage efficiency can be achieved with Bambanker<sup>™</sup> HRM as with the vitrification preservation solution with the additional advantage of an easier handling.

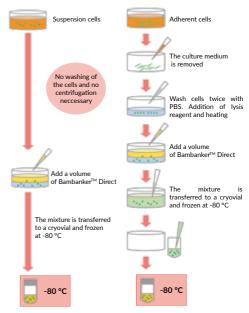
Cat. No.	Product	Content
BBH01	Bambanker <sup>™</sup> HRM	20 ml

### Bambanker<sup>™</sup> - Direct



#### Bambanker<sup>™</sup> Direct for hybridomas

There are certain cell types, such as hybridomas, which show increased sensitivity to external stress. These cells often have higher death rates or unwanted differentiation after long-term storage and freeze-thaw cycles. Bambanker™ Direct was created especially for these types of cells. They can now be frozen for long-term storage immediately upon addition of the cryoprotectant, eliminating the need for a centrifugation step. Bambanker™ Direct is added one-to-one with the cell medium, mixed and directly placed in the freezer. Bambanker™ Direct does not contain serum components. This is advantageous for cells for which animal-derived serum could be an issue.



#### Ordering information

Cat. No.	Product	Content
BBD01	Bambanker <sup>™</sup> Direct	20 ml

### Bambanker<sup>™</sup> - DMSO-Free

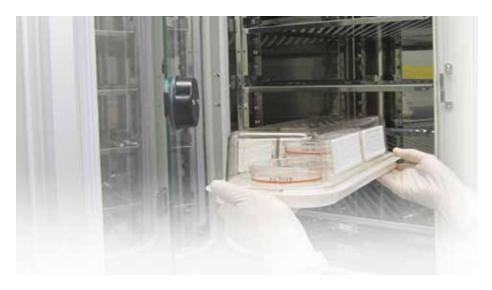


#### No more DMSO - for the most sensitive cells

DMSO is added to most freezing reagents to help avoid formation of ice crystals, which harm the cells during freezing. Unfortuantely, DMSO is cytotoxic and can reduce the survival rate of certain sensitive cell lines. Bambanker™ DMSO Free is made without DMSO. Instead, it uses a unique formula to avoid the formation of ice crystals. This makes Bambanker™ DMSO Free especially suitable for cell lines that are sensitive to DMSO under long-term storage conditions.

Cat. No.	Product	Content
BBF01	Bambanker <sup>™</sup> DMSO-Free	20 ml

### *ூட்டாட* Cell Culture Chamber



- Cell culture protective tray for the protection of your cells
- Versatile Usable with all standard plates, dishes and flasks
- Essential to avoid contamination of your cells

#### Protect your valuable cells

Tissue and cell culture are indispensable tools for modern biology. Nevertheless, many cell biologists had the frustrating experience of dealing with microbial infections or cross-contamination with other cell lines. This problem can become a disaster when primary cells or stem cells are affected. Cell culture vessels (flasks, plates or dish) are unknowingly exposed to pathogens in three primary locations: (1) cell culture hood, (2) cell incubator, and (3) during transport between them. It is critical to maintain a sterile environment and clean cell culture technique. However, accidents happen and once a culture container is infected, it can cause the infection to spread to other cell culture vessels throughout the incubator. This can cause massive issues for the scientific research in your lab!

#### No contamination and no infections

With the FastGene® Cell Culture Chamber, contamination is now a problem of the past. These simple-to-use chambers deliver a sterile environment for your culture plates, dishes, and flasks. The chambers provide protection while in the incubator, under the cell culture hood, or during transportation. Stop the risk of infecting your cells!





Each FastGene $^{\circ}$  Cell Culture Chamber can fit numerous plates, dishes, and flasks — making them ideal for labs with multiple users.

Cat. No.	Product	Content	
CC01	FastGene® Cell Culture Chamber	1 x autoclavable chamber including filters	
CC01F	Filters	Replacement set containing 4 x filters	

lotes	

Bambanker™ is a registered trademark of Lymphotec, Tokyo, Japan FastGene® is a registered trademark of NIPPON Genetics EUROPE GmbH, Dueren Germany GelRed® is a registered trademark of Biotium, Hayward, USA IDsol™ is a registered trademark of Cyrusbioscience, Taiwan Illumina® is a registered trademark of Illumina, San Diego, USA KingFisher™ is a registered trademark of Thermo Fisher Scientific, Waltham, USA LightCycler® is a registered trademark of Hoffmann-La-Roche, Basel, Switzerland Matrigel® is a registered trademark of Corning, Corning, USA Mupid™ is a registered trademark of Advance, Tokyo, Japan Qubit® is a registered trademark of Thermo Fisher Scientific, Waltham, USA Rotor-Gene® is a registered trademark of Qiagen, Hilden, Germany ROX™ is a registered trademark of Applied Biosystems, Waltham, USA StemFit® is a registered trademark of Ajinomoto, Tokio, Japan SYBR® is a registered trademark of Thermo Fisher Scientific, Waltham, USA Synthemax® is a registered trademark of Corning, Corning, USA TaqMan® is a registered trademark of Applied Biosystems, Carlsbad, USA TGX™ is a registered trademark of Bio-Rad, Hercules, USA







