

Very Wide Array

Fluorescent/Radioisotope



Science Imaging

System



ur FLA-3000, with its wide format Imaging Plate, gives you the power to do more. Now you can quickly perform gene expression profiling

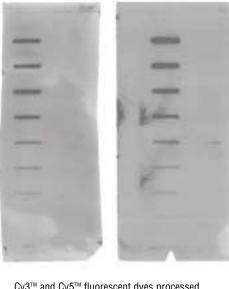
using commercially available cDNA membranes up to 22 x 22cm in size. How quickly? A complete 23 x 40cm Imaging Plate can be scanned in six minutes.

not only efficiently detect and discriminate between Cy3TM and Cy5, TM but also between $Cy2^{TM}$ and $Cy5^{TM}$ labeled samples. Plus, the FLA-3000 continues to push the boundaries of performance for a multifunctional scanner in speed, throughput, sensitivity and resolution. All that, and a 23cm wide-format Imaging Plate.

The power of multiformat detection

And the FLA-3000 allows you to configure it to process samples labeled with either radioisotopes or fluores-

cent dyes – all within a single research tool. Its combination of two solidstate Second Harmonic Generation lasers (blue: 473nm, green: 532nm) and a He-Ne laser (red: 633nm) makes the FLA-3000 truly unique. For example, it is the only system that can



22 x 22cm cDNA macroarray radioisotope image

Image courtesy of Norio Komatsu, M.D., Ph.D., Jichi Medical School and Masao Seto. M.D., Ph.D., Aichi Cancer Center Research Institute, cDNA membrane (Genome Systems, Inc.) hybridized with 32P-labeled probe and exposed for 70 hours to a BAS-MS 2340 Imaging Plate. (FLA-3000 settings: 50µm pixel, \$4000 and L5)

Cy3™ and Cy5™ fluorescent dyes processed on a single sample

respective diluted protein on the membrane was excited by either



apturing images of samples labeled with fluorescent dyes has never been easier. At the heart of our FLA-3000 lies an exclusive, patented Second Harmonic Generation (SHG) laser assembly. Its SHG532nm

green laser allows for numerous applications to satisfy even the most demanding laboratories. For instance, the use of RITC, SYPRO® Red, HNPP and $Cy3^{TM}$ is now a practical reality. The FLA-3000 also gives you speed. Images of $50\mu m$, $100\mu m$ or $200\mu m$ pixel size, to a maximum size of $23 \times 40 cm$, can be obtained in six minutes. Smaller areas, of course, take even less time to scan.

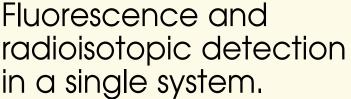
The FLA-3000 can also be configured with just the SHG473nm laser and a He-Ne (633nm) laser, making it ideal for SYBR®

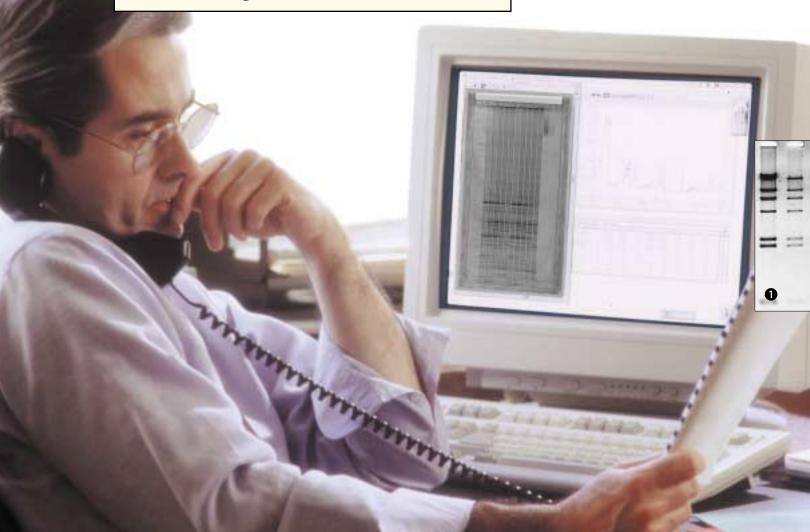
Green I and SYBR® Green II, ethidium bromide, FITC, Phycoerythrin and Cy5, $^{\text{TM}}$ not to mention alkaline phosphatase substrates such as AttoPhos $^{\text{TM}}$ and DDAO-phosphate.

Capturing radioisotopic images with proven, patented technology.

Fujifilm had more than fifteen years of experience with phosphor Imaging Plate (IP) technology for medical radiography before they expanded its use to bioresearch. And the FLA-3000 makes good use of this development. The Imaging Plate in the FLA-3000 is approximately 100 times more sensitive to emitted radiation than conventional X-ray film. It reduces sample exposure times by as much as 90% and the amount of radioisotope needed for detection by as much as 90%. What's more, Imaging Plates are easy to use. They are reusable. They do not require a dark room or chemicals, and they can be exposed at room temperature.

Imaging Plates have a flexible polyester base





coated with highly dispersed barium fluorohalide phosphor crystals (BaF(Br,I):Eu²⁺). When a radioactively labeled sample is exposed to an IP, the energy from the sample is transferred to the phosphor crystals and stored as trapped electrons. In short, the electrons become trapped to form 'F-centers' in the BaF(Br,I) matrix and Eu²⁺ ions trap holes. These F-centers have an absorption band at about 600nm. So scanning the exposed IP with the He-Ne (633nm) laser releases the trapped electrons to the conduction band. The trapped electrons recombine with the holes trapped by Eu,²⁺ and photons at about 400nm are released. This process is known as photo-stimulated luminescence. The blue light emitted is collected to produce the digitized image.

This Imaging Plate technology makes the FLA-3000 a fast and accurate means for quantitative analysis of all alpha, beta, and gamma-emitting isotopes (including tritium). With the BAS-ND type IP, neutron ray detection is also possible. The FLA-3000 is ideal for a broad spectrum of applications, from gene array analysis to drug metabolism.

Collecting images from radioactivelylabeled samples is as easy as generating images from fluorescent-labeled samples.

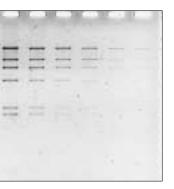
Three lasers mean more applications.

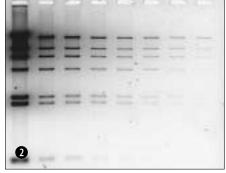
The FLA-3000 performs a wide array of imaging applications, from fluorescence to radioisotope (IP method). It's ideal for SYBR® Green I and II, SYPRO® Orange, SYPRO® Ruby, ethidium bromide, FITC, Phycoerythrin and Cy5.™ It's fast. You get six-minute scanning at even the highest resolution for areas as large as 23 x 40cm. Our patented SHG laser is the only laser technology that efficiently detects and discriminates not only between Cy3™ and Cy5,™ but also Cy2™ and Cy5™ labeled samples. And light emitted from our SHG laser is coherent,

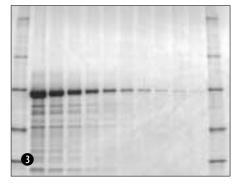
which means it's not dispersed when you use thick gels. So, unlike with other light sources, you get good image quality even with thick gels.

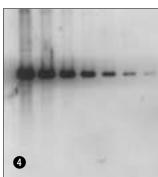
Speed, image quality, flexibility and extraordinary capabilities make the FLA-3000 a remarkable research tool, with much of the credit going to the revolutionary SHG Laser. The laser diode assembly in the FLA-3000 uses our newly developed, periodically poled MgO/LiNbO₃ (PPMGLN) crystal as a wavelength transformer. Inside the crystal, the direction of the dipole is periodically inverted. Hundreds of inverted regions, with a width of approximately 2.5µm, are aligned within the 1mm thick crystal. The wave-











Data Images, left to right:

On monitor: SYBR® Green I-stained Differential Display. Sample: Mouse Testis. Courtesy of Dr. Yasuyuki Kurihara, Yokohama National University.

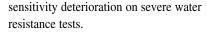
- 1. EtBr-stained DNA agarose gel. Sample: λ DNA/Hind III. Applied Amount: μ g/100ng/50ng/20ng/10ng/5ng/2ng/1ng
- 2. $SYBR^{\circ}$ Green I-stained DNA agarose gel. Sample: λ DNA/Hind III. Applied Amount: μ g/100ng/50ng/20ng/10ng/5ng/2ng/1ng
- 3. SYPRO® Orange-stained protein SDS-PAGE. Sample: Bovine Serum Albumin. Applied Amount: MW Marker//µg/500ng/200ng/100ng/50ng/20ng/10ng/5ng/2ng/1ng/MW Marker

length of the secondary light beam is then freely controlled by varying the width of the poled regions. The use of this new crystal along with other SHG technology led to the world's first blue, solid-state laser diode.

A Configuration for Your Laboratory

The FLA-3000 is available in various configurations, so you select the system that best meets the demands of your laboratory today.

Tomorrow, if your research or application needs change, you can add additional capabilities.



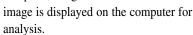
The wider detection area of this high water resistant Imaging Plate in combination with ArrayGauge software (Windows®) makes the

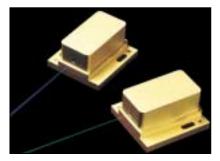
FLA-3000 ideal for macroarray analysis.

The BAS-MS Imaging Plate also has a soft ferrite backing layer, so you can use it with all the scanners in the BAS series.

How to capture fluorescent images:

- 1. Put the sample on the FLUOR-Stage.
- 2. Open the cover and place the FLUOR-Stage on the loading unit.
- 3. Select desired operating parameters and initiate scan.
- 4. When scanning is complete, a digitized

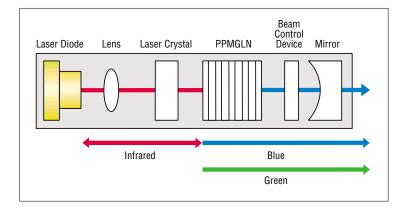


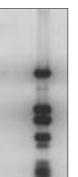


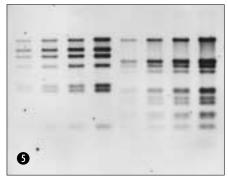
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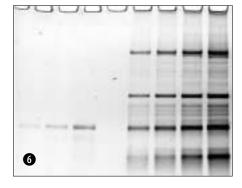
New BAS-MS Imaging Plate

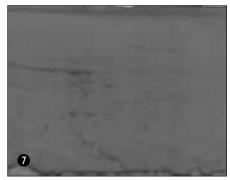
With its high sensitivity and high water resistance, our newly developed photo stimulable phosphor Imaging Plate gives you a higher level of performance. In fact, there's no











- 4. Chemifluorescence Southern blotting. Detected with AttoPhos™ as alkaline phosphatase substrate. Sample: pBR328/ Bam HI. Applied amount: 100pg/30pg/10pg/30pg/10pg/30pg/100fg/30fg/100fg/30fg/100fg/MW Marker
- 5. Chemifluorescence Southern blotting. Detection with DDAO-phosphate (1,3-dichloro-9, 9-dimethylacridin-2-one-7-yl). Sample: Left four lanes: \(\lambda\) DNA/Hind III. Right four lanes: \(\lambda\) DNA/Eco RI+Hind III. Applied amount: Left four lanes; 10ng/20ng/50ng/100ng, Right four lanes; 10ng/20ng/50ng/100ng
- 6. FITC (Fluorescein isothiocyanate): Conventional SDS-PAGE of protein. Sample: Monoclonal antibody labeled with FITC and MW marker labeled with FITC. Method: SDS-PAGE
- 7. SYPRO® Orange-stained 2D protein electrophoresis. Sample: Yeast. Courtesy of Dr. Yoshihiro Yamamoto, Kyoto Municipal Institute of Industrial Research.







Specifications and Applications

Specifications

Scanner	
Excitation Lasers	473nm (SHG)
	532nm (SHG)
	633nm (He-Ne)
Detectable Nuclides	¹⁴ C, ³² P, ³³ P, ³⁵ S, ¹²⁵ I, ³ H, and others
Fluorescence	Cy2 [™] , ECF, ECL Plus [™] , FITC, Fluorescein,
	SYBR® Green I, SYBR® Green II, SYBR®
	Gold, SYPRO® Ruby, SYPRO® Orange, Vistra
	Green [™] Alexa 532, Cy3 [™] Ethidium
	Bromide, Rhodamine B, SYPRO® Red, Texas
	Red,® Cy5,™ DDAO-Phosphate
Chemifluorescence	AttoPhos,™ AttoPhos Plus,™ ECL Plus,™
	HNPP, DDAO-phosphate, etc.
Scanning Area	23 x 40cm (max.) (scanning area is to be
	designated)
Pixel Size	50μm/100μm/200μm (selectable)
Gradation	65, 536 or 256 (selectable)
Dynamic Range	5 orders of magnitude
Sensitivity for RI Detection	S 4,000
Sensitivity for Fluor. Detection	F 1, 10, 100, 1000 (selectable)
Scanning Time	ca. 6 min. (50µm, 23 x 40cm)
Software for Scanning	ImageReader (MacOS/Windows®)

Image Analysis		
Computer Hardware	Macintosh™/IBM Compatible (selectable)	
Operating System	MacOS/Windows® 95, Windows® 98,	
	Windows® NT	
Main Memory	128MB or More	
HDD	4GB or More	
17-inch Color Monitor		
3.5-inch MO Disk, 640MB Type (1	28/230/540MB compatible)	
Laser Beam Printer		
Software for Image Analysis	Science Lab (MacOS/Windows®)	
Environment		
Power Supply	AC100V±10%, AC120V±10%,	
	AC240V±10%	
Frequency	50/60Hz	

Operating Environment 15°C - 30°C, 30%~70% Humidity (avoid condensation)

Power Consumption About 1.1kVA

Options

High-Quality Pictrography 3000/4000 Macroarray Analysis Software High-Sensitivity, Water Resistance IP IP Cassette

High-Sensitivity, Water Resistance IP BAS-MS 2325/2340/2025/2040
IP Cassette BAS Cassette 2325, 2025, 2040
Imaging Plates BAS-SR 2025/2040, BAS-TR 2025/2040,

and BAS-ND 2025/2040

ArrayGauge (Windows®)

IP Eraser

Accessories



FLUOR-Stage
For capturing fluorescent or chemifluorescent images. 23x40cm.



IP Stage
For capturing radioisotopic images. Exposed IP attaches to the magnetic stage using a soft-ferrite backed surface. 23x40cm



Multi-Stage
For detecting fluorescence in 20 x
40cm glass plate for DD method or
in microtiter plates. 23 x 40cm.



IP Cassette 2340
Used when IP is exposed to RIlabeled sample. Available in: 23x25cm, 23x40cm, 20x25cm, 20x40cm.



Imaging Plates Available types: BAS-MS 2325, BAS-MS 2340, BAS-MS 2025, BAS-MS 2040.



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