

SYBR™ Safe DNA Gel Stain

Catalog Number S33100, S33101, S33102, S33103, S33110, S33111, S33112

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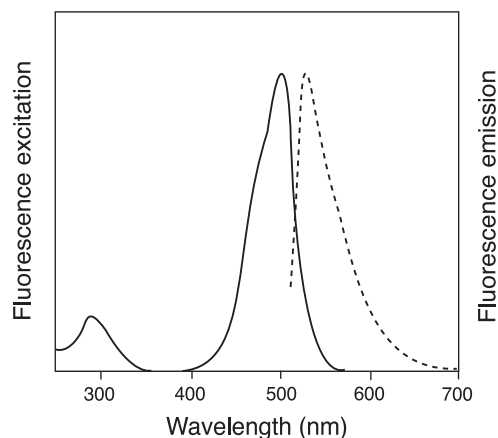
Table 1 Contents and storage

Material	Cat. No.	Amount	Concentration	Storage	Stability
SYBR™ Safe in TBE buffer	S33100	1 L*	0.5X, 45 mM Tris-borate, 1 mM EDTA, pH ~8.3	<ul style="list-style-type: none"> • 2–25°C • Protect from light 	~6 months, when stored as directed
	S33101	4 L†			
SYBR™ Safe in TAE buffer	S33111	1 L*	1X, 40 mM Tris-acetate, 1 mM EDTA, pH ~8.3		
	S33112	4 L†			
SYBR™ Safe in DMSO	S33102	400 µL	10,000X		
	S33103	50 µL			
SYBR™ Safe DNA Gel Stain Starter Kit	S33110	1 L of SYBR™ Safe gel stain, and one SYBR™ Safe photographic filter	0.5X TBE		
Number of labelings: * Provides sufficient material to stain ~20 minigels. † Provides sufficient material to stain ~80 minigels. 4 L unit size is packaged in a cube-shaped container with a removable spigot for easy dispensing and storage.					
Approximate fluorescence excitation and emission maxima: 280, 502/530 nm, bound to DNA					

Introduction

SYBR™ Safe DNA gel stain has been specifically developed for reduced mutagenicity, making it safer than ethidium bromide for staining DNA in agarose or acrylamide gels. SYBR™ Safe stain comes either as a concentrate or as a ready-to-use solution that can be used like an ethidium bromide solution, and the detection sensitivity with SYBR™ Safe stain is comparable to that obtained with ethidium bromide. DNA bands stained with SYBR™ Safe DNA gel stain can be detected using a standard UV transilluminator, a visible-light transilluminator, or a laser-based scanner. The stain is also suitable for staining RNA in gels. Bound to nucleic acids, SYBR™ Safe stain has fluorescence excitation maxima at 280 and 502 nm, and an emission maximum at 530 nm (Figure 1, page 2).

Figure 1 Normalized fluorescence excitation and emission spectra of SYBR™ Safe DNA gel stain, determined in the presence of DNA



Before starting

Storage, handling, and disposal

You may store the SYBR™ Safe DNA gel stain at any temperature between 2°C to 25°C. SYBR™ Safe in DMSO freezes at low temperatures; therefore, the product must be completely thawed and mixed before using. Repeated freeze-thawing has minimal impact on product performance.

SYBR™ Safe DNA gel stain showed no or very low mutagenic activity when tested by an independent, licensed testing laboratory, and this stain is not classified as hazardous waste under U.S. Federal regulations. The safety testing included 3 well-established mammalian cell-based tests (Table 2, page 3), a battery of well-established Ames-test bacterial strains (Figure 2, below), and extensive testing for environmental safety (Tables 3 and 4, page 3). Use care when using this reagent and dispose of the stain in compliance with all pertaining local regulations.

Figure 2 Summary of Ames test results for mutagenicity. Samples were pre-treated with a mammalian S9 fraction and then tested using the indicated Ames test strain. With strains TA97a, TA98, TA100, and TA102, a result of less than two-fold above background suggests that the compound is nonmutagenic; whereas, a result of greater than this value suggests that the compound is mutagenic. With strains TA1535, TA1537, and TA1538, a result of less than three-fold above background suggests that the compound is nonmutagenic; whereas, a result of greater than this value suggests that the compound is mutagenic. All tests were performed by Covance Laboratories, Inc., Vienna, VA, an independent testing laboratory.

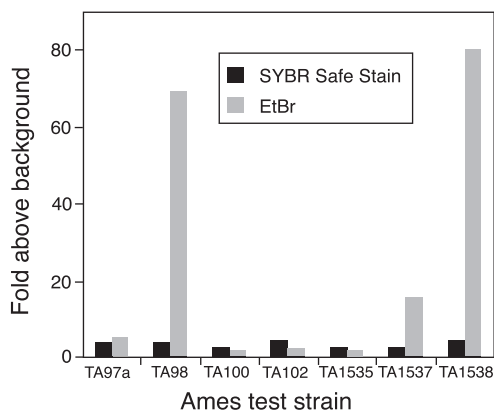


Table 2 Summary of mammalian cell-based tests for DNA genotoxicity

Test*	Cell type	Test result with S9 activation [†]	Test result without S9 activation [†]
Transformation test ¹	Syrian hamster embryo (SHE) cells	Not applicable	Negative
Chromosomal aberration test ²	Cultured human peripheral blood lymphocytes	Negative	Negative
Forward-mutation test ^{3,4}	L5178Y TK mouse lymphoma cells	Negative	Negative
<p>* All tests were performed by Covance Laboratories, Inc., Vienna, VA, an independent testing laboratory. † S9, a mammalian extract obtained from Aroclor™ 1254-induced rat liver.</p> <p>1. Fundamental and Molecular Mechanisms of Mutagenesis 356:1 (1996); 2. Evans, H.J., in <i>Chemical Mutagens, Principles and Methods for their Detection Vol 4</i>, A. Hollaender, Ed., Kluwer Academic/Plenum Publishers (1976) pp. 1–29; 3. Mutation Res 72, 447 (1980); 4. Mutation Res 59, 61 (1979).</p>			

Table 3 Summary of environmental safety test results

Test*	Method	Results
Aquatic toxicity	Fathead minnow CA Title 22 acute screening	Not classified as hazardous or toxic to aquatic life
Ignitability	EPA 1010	Not ignitable (>100°C)
Corrosivity	EPA 150.1	Not corrosive (pH = 8.25)
Corrosivity (by Corrositex)	DOT-E 10904	Category 2 noncorrosive
Reactivity	EPA 9010B/9030A	No reactivity detected
<p>* All tests were independently confirmed by AMEC Earth and Environmental San Diego Bioassay Laboratory, San Diego, CA.</p>		

Table 4 Summary of pollutant discharge test results

Test*	SYBR™ Safe Stain [†]	0.5X TBE
pH (150.1)	8.45	8.48
Total cyanide (335.2)	None detected	None detected
Chemical oxygen demand (COD; 410.1)	7020	6840
Ammonia as nitrogen (350.1)	253	248
Total organic carbon (415.1)	2480	2360
Total phenolics (420.1)	None detected	None detected
Organochlorine pesticides and PCBs (608M)	None detected	None detected
Semi-volatile organic compounds (625)	None detected	None detected
Volatile organic compounds (624)	None detected	None detected
Metals (6010B, 7060A, 7421, 7470A, 7740, 7841)	None detected	None detected
<p>* Code of Federal Regulations Title 40, Part 136; † 1X SYBR™ Safe stain (Lot X40023) in 0.5X TBE.</p>		

Experimental protocols

Staining nucleic acids after electrophoresis

- 1.1 Soak the gel in SYBR™ Safe stain. If using SYBR™ Safe gel stain concentrate, dilute 10,000X in TAE or TBE buffer (as appropriate) prior to use. Place the gel in a plastic container, such as a pipet-tip box lid or a household food-storage container. Do not use a glass container, because the dye in the staining solution may adsorb to the walls of the container, resulting in poor gel staining. Add sufficient SYBR™ Safe DNA gel stain to cover the gel. A 50 mL volume is sufficient for staining most standard minigels. To stain larger gels, increase the volume of staining solution in proportion to the increased gel volume, and ensure that the gel is fully immersed during staining.
- 1.2 Incubate for 30 minutes. Cover the gel and the staining solution with aluminum foil or place them in the dark to protect from light. Continuously agitate the gel on an orbital shaker at 50 rpm. No destaining is required.

Precasting SYBR™ Safe Stain in agarose gels

- 2.1 Prepare the agarose gel directly in SYBR™ Safe DNA gel stain. SYBR™ Safe stain is provided in buffer; simply substitute SYBR™ Safe stain for the buffer when preparing the molten agarose. If using the 10,000X SYBR™ Safe stain concentrate, dilute the concentrated stain 1:10,000 in agarose gel buffer (e.g., 1X TBE or 1X TAE) and add the buffer plus stain mixture to the powdered agarose. For example if you run TBE gels and require 30 mL of molten agarose for your tray, add 3 µL of 10,000X SYBR™ Safe stain concentrate to 30 mL of 1X TBE, mix well, and add to the powdered agarose.

Note: You can heat the agarose/SYBR™ Safe stain mixture in the microwave. As with precasting gels with ethidium bromide, the mobility of nucleic acid fragments in the gel may be somewhat slower when run in these gels compared to their mobility in the gel without stain.

- 2.2 Run the gel. Use a running buffer appropriate to the SYBR™ Safe gel stain formulation. No post-staining or destaining is needed.

Viewing and photographing the gel

You can view stained gels using a standard 300 nm transilluminator, a 254 nm epi- or transilluminator, or a blue-light transilluminator such as the Safe Imager™ 2.0 Blue-Light transilluminator (Cat. no. G6600). DNA stained with SYBR™ Safe stain can also be visualized and analyzed using imaging systems equipped with an excitation source in the UV range or between 470–530 nm. Refer to Table 5 on page 5 to determine the optimal filter sets to use, or contact the instrument manufacturer for advice.

Note: If bands from the SYBR™ Safe stained gel are to be excised and used in a ligation reaction, we recommend that the gel is illuminated using blue-light source (i.e., Safe Imager™ 2.0 Blue-Light transilluminator) and not a UV light source. In some instances, UV light sources in combination with SYBR™ Safe stain can lead to reduced cloning efficiencies.

You can photograph stained gels using Polaroid™ 667 black-and-white print film and SYBR™ Safe photographic filter (Cat. no. S37100). SYPRO™ photographic filter (Cat. no. S6656) or a Kodak™ Wratten #9 filter also work well. Using this film and one of these filters, SYBR™ Safe DNA gel stain provides the same detection sensitivity as ethidium bromide and an appropriate filter. Do not use a standard ethidium bromide photographic filter with SYBR™ Safe DNA gel stain. Gels stained with SYBR™ Safe stain can also be imaged using a CCD camera or a laser-based scanner.

Table 5 Filter selection guide for use with SYBR™ Safe stain

Instrument (Manufacturer)	Excitation source	Emission filter
Alphalmager (Alpha Innotech)	302 nm	SYB-500
Alphalmager HP (Alpha Innotech)	302 nm	SYB-500
AlphaDigiDoc RT (Alpha Innotech)	UV transilluminator	
Shroud, Camera Stand (Alpha Innotech)	UV transilluminator	SYB-100
DE500 or DE400 light cabinet 2.17" diameter (Alpha Innotech)	UV transilluminator	SYB-500
DE500 or DE400 light cabinet 2" diameter (Alpha Innotech)	UV transilluminator	SYB-400
VersaDoc Imaging Systems (Bio-Rad)	Broadband UV	520LP
Molecular Imager FX Systems (Bio-Rad)	488 nm	530 nm BP
Gel Doc Systems (Bio-Rad)	302 nm	520DF30 (#170-8074)
Typhoon 9400/9410 (GE Healthcare)	488 nm	520 BP 40
Typhoon 9200/9210/8600/8610 (GE Healthcare)	488 nm	526 SP
FluorImager (GE Healthcare)	488 nm	530 DF 30
Storm (GE Healthcare)	Blue (fluorescence mode)	
VDS-CL (GE Healthcare)	Transmission	UV Low
Ultracam/Gel Imager (Ultra-Lum)	UV	Yellow Filter (#990-0804-07)
Omega Systems (Ultra-Lum)	UV	520 nm
Polaroid Camera (Polaroid)	UV	SYBR™ Safe Photographic Filter (S37100)
FOTO/Analyst Express/Investigator/Plus/Luminary (FOTODYNE)	UV	Fluorescent Green (#60-2034)
FOTO/Analyst Minivisionary (FOTODYNE)	UV	Fluorescent Green (#62-4289)
FOTO/Analyst Apprentice (FOTODYNE)	UV	Fluorescent Green (#62-2535)
FOTO/Analyst Luminary (FOTODYNE)	UV	Fluorescent Green (#60-2056)
FCR-10 (Polaroid)	UV	#3-4218
FUJI FLA-3000 (FUJI Film)	473 nm	520LP
BioDocIt/AC1/EC3/BioSpectrum (UVP)	302 nm	SYBR™ Green (#38-0219-01) or SYBR™ Gold (#38-0221-01)
Gel Logic (Kodak)	UV	535 nm WB50
Syngene Instruments (Syngene)	UV	500–600 nm Shortpass filter

Product list

Current prices may be obtained at www.thermofisher.com or from our Customer Service Department.

Cat no.	Product name	Unit size
G6600	Safe Imager™ 2.0 Blue-Light Transilluminator	each
S33100	SYBR™ Safe DNA gel stain in 0.5X TBE	1 L
S33101	SYBR™ Safe DNA gel stain in 0.5X TBE	4 L
S33102	SYBR™ Safe DNA gel stain *10,000X concentrate in DMSO*	400 µL
S33103	SYBR™ Safe DNA gel stain *10,000X concentrate in DMSO* *sample size*	50 µL
S33110	SYBR™ Safe DNA Gel Stain Starter Kit *with 1 L of SYBR™ Safe DNA gel stain in 0.5X TBE (S33100) and one photographic filter (S37100)*	1 kit
S33111	SYBR™ Safe DNA gel stain in 1X TAE	1 L
S33112	SYBR™ Safe DNA gel stain in 1X TAE	4 L
S37100	SYBR™ Safe photographic filter	each
G5218-01	E-Gel™ 1.2% with SYBR™ Safe	18 gels
G6206-01	E-Gel™ 1.2% with SYBR™ Safe Starter Kit	1 kit
G5218-02	E-Gel™ 2.0% with SYBR™ Safe	18 gels
G6206-02	E-Gel™ 2.0% with SYBR™ Safe Starter Kit	1 kit

Purchaser notification

These high-quality reagents and materials must be used by, or directly under the supervision of, a technically qualified individual experienced in handling potentially hazardous chemicals. Read the Safety Data Sheet provided for each product; other regulatory considerations may apply.

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