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SuperKine™ West Pico PLUS Chemiluminescent Substrate

Cat #: BMU101-EN Size: 100 mL/500 mL

[-]	Product Name: SuperKine™ West Pico PLUS Chemiluminescent Substrate		
REF	Catalog Number: BMU101-EN	LOT	Lot Number: Refer to product label
	Formulation: Liquid, ready to use		Applications: WB, IP
Å	Storage: Protected from light, store at 4°C,	\triangle	Note: No
	stable for 12 months from date of shipment		

Product Application

Assay principle: After years of development on chemiluminescence technology, there have been a variety of luminescence systems, but the most commonly used one in laboratories is still a technology system based on Luminol or its derivatives (isoluminol, etc.). Western luminescent detection reagent ECL immunoblotting chemiluminescence solution is a non-radioactive (horseradish peroxidase) luminescence system designed to detect trace protein immobilized on a solid membrane (such as NC, PVDF, etc.). It is an experimental auxiliary reagent for the photosensitive recording of its immunoblotting by X-ray film (radiograph).

Advantages	Mechanism	Application
High sensitivity	Add unique immune signal	Easier to obtain detection signals, better detection of
	enhancement components	low-abundance proteins (picogram level)
Signal stability	The signal is strong and lasts for 4	The signal is more stable, the signal can still be
	hours	obtained after 4h color development of the signal film
Easy operation	Contains optimized antibody	Increase the storage time of the diluted antibody, and
	stabilization components	can be used repeatedly
High compatibility	Suitable for PVDF membrane and	Good compatibility with a variety of membrane types,
	NC membrane	worry-free use
Stable performance	Stored at 4°C for one year, there is	Can be stored for more than one year without affection
	no difference in color rendering effect	on use
Universal detection	Can be detected with X-ray film and	Can get good results through a variety of instruments
	chemiluminescence imager	

Application suggestion:

- 1. Routine electrophoresis, membrane transfer, HRP-labeled antibody incubation, and membrane washing. It is recommended to use SuperKine™ Enhanced Antibody Dilution Buffer (BMU103-EN) to dilute the antibody. Universal Loading Control Antibody Cocktail (KTD101-EN) is used for the sample internal reference detection. It is recommended to use HRP-labeled IgG, such as, HRP goat anti-rabbit secondary antibody (A21020).
- 2. The diluted antibody should be stored at 4°C immediately after the antibody incubation, so that it can be reused later. While washing the HRP-labeled secondary antibody on the membrane, freshly prepare the luminescence working solution, mixing the two reagents at 1:1 ratio to prepare the working solution.



- 3. If the size of the blotting membrane is 1 cm², it is recommended to use 0.1-0.2 ml of SuperKine™ West Pico PLUS Chemiluminescent Substrate working solution.
- 4. Incubate in the ECL working solution for 1-5 minutes.
- 5. Clamp the membrane with tweezers, and gently touch the lower edge of the membrane with the filter paper to remove excess luminescent liquid on the membrane. Cover the blotting membrane with a transparent plastic wrap.
- 6. Expose to X-ray film or take photo by chemiluminescence imager.

Highlight moment: Except the enhanced ECL luminescent solution, there are many ways to improve the signal for immunological experiments, such as using SuperKine™ Enhanced Antibody Dilution Buffer (BMU103-EN) in WB experiment, choosing DyLight, IFKine™ specialized secondary antibodies in IF experiment, or adopting high-quality animal serum, these are all pretty good options. Scan the QR code on the right side to view more Abbkine product information.



Experiment results display:



Fig. The sample is Mouse TNF-alpha protein (PRP1113, 17KD), the primary antibody is TNF- α Polyclonal Antibody (ABP0127, 1:2500), and the secondary antibody is HRP, Goat Anti-Rabbit IgG (A21020, 1:10000). The exposure time is 30s.

Precautions:

- 1. Do not mix components from different batch numbers and different manufacturers; otherwise, it may cause abnormal results.
- 2. In order to obtain the best experiment results, you need to optimize all of your experiment elements, including the number of samples, antibody concentration, as well as the use of membranes and blocking reagents.
- 3. Mix the two substrate components at 1:1 ratio to prepare a substrate working solution. Please pay attention to change the tips during the aspiration process of A and B solution.
- 4. SuperKine™ West Pico PLUS Chemiluminescent Substrate has a long luminescent duration, but it is best to perform compression or imaging within 30 minutes of color development.
- 5. It is recommended that every 1 cm² membrane corresponds to 0.1-0.2 mL SuperKine™ West Pico PLUS Chemiluminescent Substrate working solution (picogram level).

FAQ:

- 1. Can BMU101-EN be reused? A: Repeated use is not recommended. The correct way is to add the ECL working liquid drops to the film, make it evenly covered on the film and then expose it on the machine.
- 2. What is the protein detection range of the luminescent solution? A: Our company verified that the color developing solution had the best effect in the protein expression range of 6.25-50 ng.
- 3. What is the reason for the weak exposure band? Is there any good solution? A: Low abundance of protein expression or low loading amount and high dilution ratio of antibody can lead to long exposure time and weak exposure band. It is recommended to use TBST to re-wash the exposed film for 2-3 times, 1 min each time, and then replace it with a more sensitive ECL (such as BMU102-EN), which can be re-exposed without the need to transfer the film with a new sample.

Recommended Products:

Catalog No.	Product Name	Recommended Reason
BMU102-EN	SuperKine™ West Femto	Sensitive, efficient, stable signal
	Maximum Sensitivity Substrate	
BMU103-EN	SuperKine™ Enhanced	Sensitive, stable performance, wide application
	Antibody Dilution Buffer	

<u>Disclaimer:</u> The reagent is only used in the field of scientific research, not suitable for clinical diagnosis or other purposes.

