



Real-Time PCR Equipment Set Up and Preperation

Store at 2-8°C in the dark for long time storage.



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- · HiQ Pro-Block

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A Introduction

HiQ Pro-Block is a highly effective blocking reagent capable of reducing non-specific background staining associated with immunodetection techniques such as immunocytochemistry, ELISA, and Western blotting. Antibodies tend to bind non-specifically to membranes and tissues. To minimize this nonspecific binding, a blocking buffer is used to enhance signal-to noise ratio.

HiQ Pro-Block shows no cross-reactivity and it blocks more effectively than serum albumins. The blocking potency of HiQ Pro-Block is superior to most other conventional formulations in producing a high signal-to-noise ratio.

HiQ Pro-Block may also be used as effective diluents for antibodies.

HiQ Pro-Block eliminates the need for matching species with the link antibody and is often, more effective at reducing non-specific background staining than normal serum.

HiQ Pro-Block does not contain biotin. The reagent is filtered at 0.2 microns.

Appearance and pH

Buffered solution of serum-free protein and proprietary additives in a phosphate buffer, pH 7.4-7.6.

Uses and Limitations

Not to be taken internally.

For In-Vitro Diagnostic use only.

Histological applications.

Immunologic applications.

Do not use if reagents become cloudy.

Do not use past expiration date.

Use caution when handling reagents.

Immunohistochemistry

- 1. Apply enough HiQ Pro-Block to cover the tissue section.
- 2. Incubate tissue section for 5 minutes at either room temperature or 37° C prior to application of the primary antibody. After incubation, rinse once in buffer

Note: do not incubate tissue sections in excess of 10 minutes or a reduction in desired staining may occur.

Note: For bulk staining, pour HiQ Pro-Block in a covered staining tray and dip slides for 5 minutes. Replace with fresh HiQ Pro-Block after 5-10 uses. This step can be performed at the time of deparaffinization is desired.

Note: For antibodies with particularly high background staining, dilute HiQ Pro-Block in PBS (1:5-10) and use as a wash buffer in addition to the blocking step.

📤 EIA/ELISA

- 1. Add 100-300 ul of HiQ Pro-Block for 96 well plates.
- 2. Incubate microtiter well for 2-10 minutes prior to addition of sample. Rinse, and continue procedure.

Note: Do not incubate in excess of 10 minutes.

Membrane Blots

1. Incubate membranes on a shaker in 0.5-1mL/cm² of HiQ Pro-Block for 15 min to 1 hr at room temperature, or 8-30 min at 37°C.

Precautions

Avoid contact with skin and eyes.

Harmful if swallowed.

Follow all Federal, State, and local regulations regarding disposal.