## VectaFluor™ Excel Amplified Kit

Anti-Rabbit IgG, DyLight™ 488



Cat. No.	DK-1488
Storage	Store reagents in original bottles at 2–8 °C. Do not freeze
Description	Instructions for immunofluorescent staining
	VectaFluor Excel Amplifed Fluorescent Staining System offers a non-biotin amplification method for fluorescence applications. This system uses a

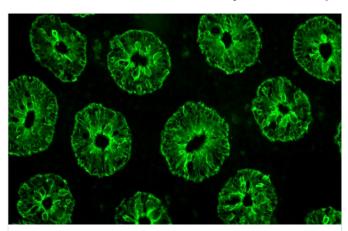
detection system.

Components

Product Name	Volume
Normal Horse Serum, 2.5%	15 ml
Amplifier Antibody (Goat Anti-Rabbit IgG)	15 ml
VectaFluor DyLight 488 Horse Anti-Goat IgG	15 ml

proprietary, ready-to-use (R.T.U.) amplifier antibody, followed by a R.T.U. VectaFluor DyLight dye-labeled

The VectaFluor Excel Amplified Kit will stain approximately 150 sections based on 100  $\mu$ l per section.



Colon: Cytokeratin detected with VectaFluor Excel Amplified Kit, Anti-Rabbit IgG, DyLight 488 (green). Mounted in VECTASHIELD® HardSet™ Antifade Mounting Medium.

## **Preparation of Working Solution**

VectaFluor Excel Amplified Kit reagents are ready-to-use—no mixing or titering is necessary to obtain optimal staining.

The staining procedure should be performed at room temperature (20–25°C). VectaFluor Excel Amplified Kit reagents should be equilibrated to room temperature for optimal performance.

A number of different wash buffers can be used. One of the most common is 10 mM sodium phosphate, pH 7.5, 0.9% saline (PBS). 0.1% Tween® 20 detergent may be added to the wash buffer and is especially recommended for use with automated stainers.

## Staining Procedure

- For paraffin sections, deparaffinize and hydrate tissue sections through xylenes or other clearing agents and graded alcohol series.
  - For frozen sections or cell preparations fix with acetone or an appropriate fixative for the antigen under study, if required.
- 2. If antigen unmasking is required, perform this procedure using a Vector® Antigen Unmasking Solution, Citrate-based (H-3300) or Tris-based (H-3301).

- 3. Wash in buffer for 5 minutes.
- 4. Incubate for 20 minutes with 2.5% Normal Horse Serum.
- 5. Tip off excess serum from sections.
- Incubate with rabbit primary antibody diluted in an appropriate diluent.
- 7. Wash in buffer for 5 minutes.
- 8. Incubate for 15 minutes with Amplifier Antibody.
- 9. Wash in buffer for 5 minutes .
- 10. Incubate for 30 minutes with VectaFluor Reagent.
- 11. Wash for  $2 \times 5$  minutes in buffer.
- 12. Mount in a media suitable for fluorescence, such as one of the VECTASHIELD Antifade Mounting Media.

Detailed product listing, specifications, protocols and additional information are available on our website; vectorlabs.com