

# m-IgG $\lambda$ BP-CFL 594: sc-516192

## BACKGROUND

Mouse IgG $\lambda$  light chain binding protein (m-IgG $\lambda$  BP) conjugated to CruzFluor™ 594 is a strongly recommended alternative to conventional anti-mouse IgG secondary antibodies for Western blotting (WB), immunofluorescence (IF) and flow cytometry (FCM) signal enhancement. Mouse IgG $\lambda$  light chain binding protein is a highly specific detection reagent that provides strong signal with minimal background and virtually complete elimination of lot-to-lot variation associated with conventionally generated secondary antibodies. Mouse IgG $\lambda$  light chain binding protein (m-IgG $\lambda$  BP) is suitable for binding to mouse IgG $\lambda$  light chain immunoglobulins; not suitable for use with mouse monoclonal IgG $\kappa$  light chain primary antibodies. CruzFluor™ 594 (CFL 594) is a red fluorescent dye that is an excellent substitute for AlexaFluor® 594, offering comparable photostability and the ability to resist protein quenching. Suitable for use with RGB imaging systems, such as Invitrogen/iBright and other comparable systems.

## SOURCE

m-IgG $\lambda$  BP-CFL 594 is a purified recombinant mouse IgG $\lambda$  light chain binding protein conjugated to CruzFluor™ 594 (CFL 594).

## PRODUCT

Each vial contains 200  $\mu$ g mouse IgG $\lambda$  binding protein-CFL 594 in 0.5 ml of PBS containing 0.1% gelatin and 0.1% sodium azide.

## APPLICATIONS

m-IgG $\lambda$  BP-CFL 594 is recommended for detection of mouse IgG $\lambda$  light chain by RGB Western Blotting (starting dilution: 1:1000, dilution range: 1:500-1:2000), immunofluorescence (starting dilution 1:50, dilution range 1:50-1:200), immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500) and flow cytometry (0.5-1  $\mu$ g per 1 x 10<sup>6</sup> cells). Optimal dilution to be determined by titration.

## RECOMMENDED SUPPORT PRODUCTS

- CrystalCruz® Cover Glasses, 22 x 50 mm, precleaned: sc-24975
- PBS (Phosphate Buffered Saline), powder, 1 packet: sc-24947
- Formaldehyde, 37% formaldehyde solution, 25 ml: sc-203049
- Hydrogen Peroxide, 30% solution, 100 ml: sc-203336
- FCM Lysing solution: sc-3621
- FCM Fixation Buffer: sc-3622
- FCM Permeabilization Buffer: sc-3623
- FCM Wash Buffer: sc-3624
- Intracellular FCM System: sc-45063

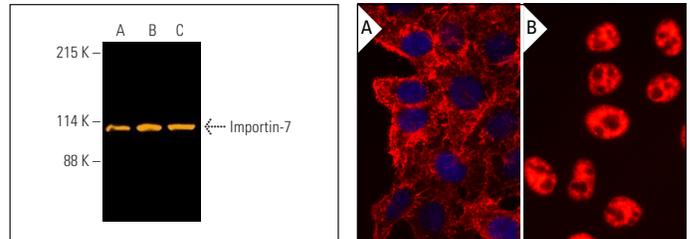
## STORAGE

Store at 4° C, \*\*DO NOT FREEZE\*\*. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

## RESEARCH USE

For research use only, not for use in diagnostic procedures.

## DATA



Importin-7 (E-2): sc-365231. Fluorescent western blot analysis of Importin-7 expression in K-562 (A), HeLa (B) and SK-N-MC (C) whole cell lysates. Blocked with UltraCruz® Blocking Reagent: sc-516214. Detection reagent used: m-IgG $\lambda$  BP-CFL 594: sc-516192.

PPP2R4 (C-10): sc-398242. Immunofluorescence detection of PPP2R4 in formalin-fixed HeLa cells showing cytoplasmic and membrane localization and nuclear DAPI counterstain (A). hnRNP F/H (1G11): sc-32310. Immunofluorescence staining of formalin-fixed HeLa cells showing nuclear localization (B). Detection reagent used: m-IgG $\lambda$  BP-CFL 594: sc-516192.

## CRUZFLUOR™ SPECTRAL PROPERTIES

PRODUCT	CAT. #	EXCITATION MAXIMUM	EMISSION MAXIMUM
m-IgG $\kappa$ BP-CFL 488	sc-516176	488 nm	514 nm
m-IgG $\lambda$ BP-CFL 488	sc-516190		
m-IgG $\kappa$ BP-CFL 555	sc-516177	556 nm	569 nm
m-IgG $\lambda$ BP-CFL 555	sc-516191		
m-IgG $\kappa$ BP-CFL 594	sc-516178	587 nm	603 nm
m-IgG $\lambda$ BP-CFL 594	sc-516192		
m-IgG $\kappa$ BP-CFL 647	sc-516179	654 nm	669 nm
m-IgG $\lambda$ BP-CFL 647	sc-516193		
m-IgG $\kappa$ BP-CFL 680	sc-516180	683 nm	700 nm
m-IgG $\lambda$ BP-CFL 680	sc-516194		
m-IgG $\kappa$ BP-CFL 790	sc-516181	786 nm	811 nm
m-IgG $\lambda$ BP-CFL 790	sc-516195		

## PROTOCOLS

See our web site at [www.scbt.com](http://www.scbt.com) for detailed protocols and support products.