## Anti-mouse IgG (H+L), F(ab')<sub>2</sub> Fragment (Sepharose<sup>®</sup> Bead Conjugate)



877-616-CELL (2355) Orders:

orders@cellsignal.com

Support: 877-678-TECH (8324)

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cellsignal.com

3 Trask Lane | Danvers | Massachusetts | 01923 | USA

<b>Applications:</b> IP	Reactivity:	Source/Isotype: Rabbit
Product Usage Information		Add 10-20 $\mu$ l of well-vortexed beads to 200 $\mu$ l of cell lysate at 1 mg/ml in 1X Cell Lysis Buffer (10X) #9803. See protocol for more details.
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA, 50% glycerol. Store at –20°C. Do not aliquot the antibodies.
Specificity/Sensitiv	vity	Assay by immunoelectrophoresis resulted in a single precipitin arc against anti-goat serum, mouse $IgG$ and mouse serum. No reaction was observed against anti-pepsin, anti-goat $IgG$ $F_c$ , or human serum proteins.
Source / Purification	on	The Anti-mouse $IgG (H+L)$ , $F(ab')_2$ Fragment is produced by immunizing goats with mouse $IgG$ whole molecules. The Anti-mouse $IgG (H+L)$ , $F(ab')_2$ Fragment is prepared from monospecific antiserum by immunoaffinity chromatography using mouse $IgG$ coupled to agarose beads followed by solid phase adsorption to remove any unwanted reactivities, pepsin digestion, and chromatographic separation.
Description		This Cell Signaling Technology antibody is immobilized via covalent binding of primary amino groups to N-hydroxysuccinimide (NHS)-activated sepharose beads. Anti-mouse IgG (H+L), F(ab') <sub>2</sub> Fragment (Sepharose Bead Conjugate) is useful for the immunoprecipitation of antibodies raised in mice.
Background		Anti-mouse IgG (H+L), $F(ab')_2$ Fragment (Sepharose Bead Conjugate) is ideal for immunoprecipitation. This secondary antibody will detect mouse antibodies. Since this secondary antibody is made with $F(ab')_2$ fragments, non-specific binding through $F_c$ receptors present on cells will be eliminated.
Species Reactivity		Species reactivity is determined by testing in at least one approved application (e.g., western blot).
Applications Key		IP: Immunoprecipitation
Cross-Reactivity K	еу	M: Mouse
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