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### Mouse B7-H4 Biotinylated Antibody

**R**Dsystems

Antigen Affinity-purified Polyclonal Goat IgG Catalog Number: BAF2154

DESCRIPTION	
Species Reactivity	Mouse
Specificity	Detects mouse B7-H4 in Western blots. In Western blots, less than 1% cross-reactivity with recombinant mouse (rm) B7-1, rmB7-2, rmB7-H1, rmB7-H2, rmB7-H3, and rmPD-L2 is observed.
Source	Polyclonal Goat IgG
Purification	Antigen Affinity-purified
mmunogen Mouse myeloma cell line NS0-derived recombinant mouse B7-H4 Phe29-Pro258 Accession # Q7TSP5	
Formulation	I vophilized from a 0.2 µm filtered solution in PBS with BSA as a carrier protein. See Certificate of Analysis for details

### APPLICATIONS

Please Note: Optimal dilutions should be determined by each laboratory for each application. General Protocols are available in the Technical Information section on our website.				
	Recommended Concentration	Sample		
Western Blot	0.1 µg/mL	Recombinant Mouse B7-H4 (Catalog # 2154-B7)		
Flow Cytometry	0.25 μg/10 <sup>6</sup> cells	See Below		

DATA		
Flow Cytometry	Detection of B7-H4 in HEK293 Human Cell Line Transfected with Mouse B7-H4 and eGFP by Flow Cytometry. HEK293 human embryonic kidney cell line transfected with either (A) mouse B7-H4 or (B) irrelevant protein and eGFP was stained with Goat Anti-Human/Mouse B7-H4 Biotinylated Polyclonal Antibody (Catalog # BAF2154) followed by APC-conjugated Streptavidin (Catalog # F0050). Quadrant markers were set based on Goat IgG control antibody staining (Catalog # Catalog # BAF108). View our protocol for Staining Membrane-associated Proteins.	
PREPARATION AND S	TORAGE	
Reconstitution	Reconstitute at 0.2 mg/mL in sterile PBS.	
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below.	
Stability & Storage	<ul> <li>ge Use a manual defrost freezer and avoid repeated freeze-thaw cycles.</li> <li>12 months from date of receipt, -20 to -70 °C as supplied.</li> <li>1 month 2 to 8 °C under starile conditions after reconstitution.</li> </ul>	

6 months, -20 to -70 °C under sterile conditions after reconstitution.

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**Global** bio-techne.com info@bio-techne.com techsupport@bio-techne.com TEL +1 612 379 2956 USA TEL 800 343 7475 Canada TEL 855 668 8722 China TEL +86 (21) 52380373 Europe | Middle East | Africa TEL +44 (0)1235 529449

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### BACKGROUND

**R**Dsystems

B7-H4, also known as B7x and B7S1, is a member of the B7 family of immune co-stimulatory proteins (1). Mature B7-H4 is a 50 kDa - 80 kDa glycosylated molecule with a 28 kDa protein core (2). Partial sensitivity to cleavage by PI-PLC suggests a possible GPI linkage of mouse B7-H4 to the cell membrane (3). The 230 amino acid (aa) extracellular region of B7-H4 contains one Ig-like V-set domain and one Ig-like C2-set domain which is followed by a hydrophobic C-terminal region (3 - 5). Within the ECD, mouse B7-H4 shares 90% and 99% aa sequence identity with human and rat B7-H4, respectively. It shares 21% - 29% aa sequence identity with B7-1, B7-2, B7-H1, B7-H2, B7-H3, and PD-L2. B7-H4 expression is induced on mitogen- or LPS-activated B cells, T cells, dendritic cells, monocytes, and macrophages (3, 4) and blocked by GM-CSF or IL-4 (6). It is also expressed on various normal epithelia and upregulated in several carcinomas and renal tubule epithelial cell lesions (2, 6 - 10). B7-H4 is expressed on the surface of macrophages but intracellularly in ovarian and breast cancer cells (2, 6 - 7). It is found in the serum and ascites fluid of cancer patients (11). B7-H4 binds an unidentified ligand on activated T cells which is distinct from BTLA, CD28, CTLA4, ICOS, and PD-1 (3 - 5, 12). Exposure to B7-H4 inhibits antigen-dependent induction of T cell proliferation and activation (3 - 5). Alternatively, B7-H4 expressing renal tubular epithelial cells promote T cell activation (8). Regulatory T cells, IL-6, and IL-10 induce B7-H4 expression on antigen presenting cells which fosters tumor growth by dampening the anti-tumor immune response (6, 13). B7-H4 also promotes the malignant transformation of epithelial cells by protecting them from apoptosis (2). Despite evidence for the involvement of B7-H4 in immune regulation, mice deficient in its expression do not show significant immune deficiencies, suggesting compensation by other molecules *in vivo* (14).

#### References:

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