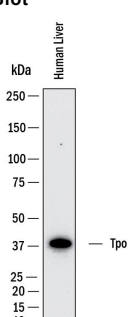


DESCRIPTION	
Species Reactivity	Human
Specificity	Detects recombinant protein specific for human Thrombopoietin in Direct ELISA.
Source	Monoclonal Mouse IgG ₁ Clone # 1065103
Purification	Protein A or G purified from ascites
Immunogen	<i>Spodoptera frugiperda</i> , Sf 21 (baculovirus)-derived human Thrombopoietin/Tpo Ser22-Gly353 Accession # P40225
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose.

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	2 µg/mL	Human liver tissue

DATA

Western Blot



Detection of Human Thrombopoietin/Tpo by Western Blot. Western Blot shows lysates of human liver tissue. PVDF membrane was probed with 2 µg/ml of Mouse Anti-Human Thrombopoietin/Tpo Monoclonal Antibody (Catalog # MAB11609) followed by HRP-conjugated Anti-Mouse IgG Secondary Antibody (Catalog # HAF018). A specific band was detected for Thrombopoietin/Tpo at approximately 38 kDa (as indicated). This experiment was conducted under reducing conditions and using Western Blot Buffer Group 1.

PREPARATION AND STORAGE	
Reconstitution	Reconstitute lyophilized material at 0.2 mg/ml in sterile PBS. For liquid material, refer to CoA for concentration.
Shipping	Lyophilized product is shipped at ambient temperature. Liquid small pack size (-SP) is shipped with polar packs. Upon receipt, store immediately at the temperature recommended below.
Stability & Storage	Use a manual defrost freezer and avoid repeated freeze-thaw cycles. <ul style="list-style-type: none"> • 12 months from date of receipt, -20 to -70 °C as supplied. • 1 month, 2 to 8 °C under sterile conditions after reconstitution. • 6 months, -20 to -70 °C under sterile conditions after reconstitution.

BACKGROUND

Thrombopoietin (Tpo), is a key regulator of megakaryocytopoiesis and thrombopoiesis. It is principally produced in the liver and is bound and internalized by the receptor Tpo R/c-mpl. Defects in the Tpo-Tpo R signaling pathway are associated with a variety of platelet disorders (1-3). The 353 amino acid (aa) human Tpo precursor is cleaved to yield the 332 aa mature protein. Mature human Tpo shares approximately 70% aa sequence homology with mouse and rat Tpo. It is an 80-85 kDa protein that consists of an N-terminal domain with homology to Erythropoietin (Epo) and a C-terminal domain that contains multiple N-linked and O-linked glycosylation sites (4, 5). Tissue specific alternate splicing of human Tpo generates multiple isoforms with internal deletions, insertions, and/or C-terminal substitutions (6). Tpo promotes the differentiation, proliferation, and maturation of megakaryocytes (MK) and their progenitors (4, 5, 7). Several other cytokines can promote these functions as well but only in cooperation with Tpo (8, 9). Notably, IL-3 independently induces MK development, although its effects are restricted to early in the MK lineage (8, 9). Tpo additionally promotes platelet production, aggregation, ECM adhesion, and activation (10 - 13). It is cleaved by platelet-derived thrombin following Arg191 within the C-terminal domain and subsequently at other sites upon extended digestion (14). Full length Tpo and shorter forms circulate in the plasma (4, 5). The C terminal domain is not required for binding to Tpo R or inducing MK growth and differentiation (5). Aside from its hematopoietic effects, Tpo is expressed in the brain where it promotes the apoptosis of hypoxia-sensitized neurons and inhibits neuronal differentiation by blocking NGF induced signaling (15, 16).

References:

1. Deutsch, V.R. and A. Tomer (2006) *Br. J. Haematol.* **134**:453.
2. Kaushansky, K. (2005) *J. Clin. Invest.* **115**:3339.
3. Li, J. *et al.* (1999) *Br. J. Haematol.* **106**:345.
4. Bartley, T.D. *et al.* (1994) *Cell* **77**:1117.
5. de Sauvage, F.J. *et al.* (1994) *Nature* **369**:533.
6. Marcucci, R. and M. Romano (2008) *Biochim. Biophys. Acta* **1782**:427.
7. Kaushansky, K. *et al.* (1994) *Nature* **369**:568.
8. Kaushansky, K. *et al.* (1995) *Proc. Natl. Acad. Sci.* **92**:3234.
9. Broudy, V.C. *et al.* (1995) *Blood* **85**:1719.
10. Lok, S.I. *et al.* (1994) *Nature* **369**:565.
11. Chen, J. *et al.* (1995) *Blood* **86**:4054.
12. Oda, A. *et al.* (1996) *Blood* **87**:4664.
13. Van Os, E. *et al.* (2003) *Br. J. Haematol.* **121**:482.
14. Kato, T. *et al.* (1997) *Proc. Natl. Acad. Sci.* **94**:4669.
15. Ehrenreich, H. *et al.* (2005) *Proc. Natl. Acad. Sci.* **102**:862.
16. Samoylenko, A. *et al.* (2008) *Cell. Signal.* **20**:154.