

# TGFB2 Rabbit pAb

货号: B11375

## 产品信息

反应	Human,Mouse,Rat
宿主	Rabbit
克隆性	Polyclonal
预测反应	<b>WB:</b> Homo sapiens , Mus musculus , Gallus gallus <b>IF:</b> Gallus gallus
应用	<a href="#">WB</a>
推荐浓度	<b>WB:</b> 1:500 - 1:1000
理论分子量	47kDa/50kDa
实测分子量	48KDa
形式	Liquid
保存条件	Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.01% thiomersal,50% glycerol,pH7.3.
偶联物	Unconjugated
阳性对照	Mouse lung,Mouse heart,Mouse uterus,Rat lung,Rat heart
细胞定位	Secreted
纯化	Affinity purification

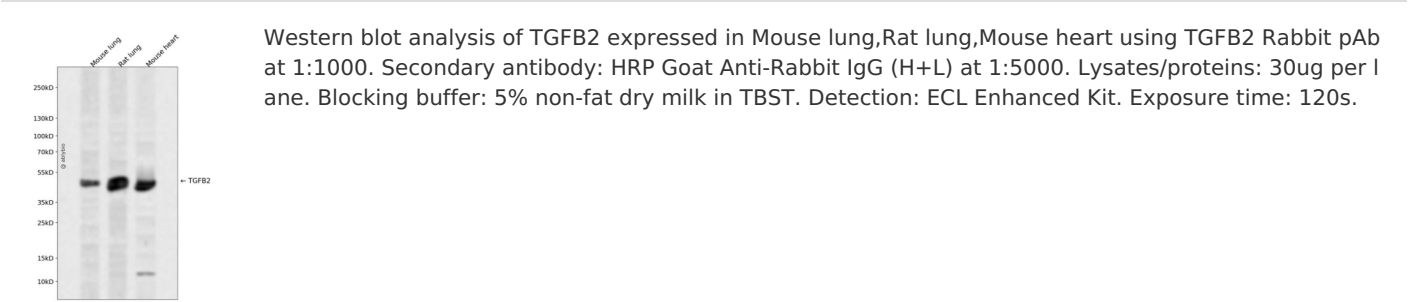
## 抗原信息

抗原信息	Recombinant fusion protein containing a sequence corresponding to amino acids 60-220 of human TGFB2 (NP_003229.1).
序列	PEEVPPEVISIYNSTRDLLQEASRRAAACERERSDEEYYAKEVYKIDMPPFFPSENAIPPTFYRPHYFRIVRFDVSAMEKNASN LVKAEFRVFRQLQNPKEVPEQRIELYQILKSKDLTSPTQRYIDSKVVKTRAEGEWLSFDVTDVAVHEWLHHKDRNLGF

## 靶点信息

研究背景	This gene encodes a member of the transforming growth factor beta (TGFB) family of cytokines, which are multifunctional peptides that regulate proliferation, differentiation, adhesion, migration, and other functions in many cell types by transducing their signal through combinations of transmembrane type I and type II receptors (TGFB1 and TGFB2) and their downstream effectors, the SMAD proteins. Disruption of the TGFB/SMAD pathway has been implicated in a variety of human cancers. The encoded protein is secreted and has suppressive effects of interleukin-2 dependent T-cell growth. Translocation t(1;7)(q41;p21) between this gene and HDAC9 is associated with Peters' anomaly, a congenital defect of the anterior chamber of the eye. The knockout mice lacking this gene show perinatal mortality and a wide range of developmental defects, including cardiac defects. Alternatively spliced transcript variants encoding different isoforms have been identified.
基因ID	7042
基因名	TGFB2
Swiss	P61812
别名	TGFB2;G-TSF;LDS4;TGF-beta2

产品验证



实验步骤

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