



EDA Monoclonal Antibody

Catalog No	YP-mAb-16071
Isotype	IgG
Reactivity	Human;Mouse;Rat
Applications	WB
Gene Name	EDA
Immunogen	The antiserum was produced against synthesized peptide derived from the Internal region of human EDA. AA range:120-170
Specificity	EDA Monoclonal Antibody detects endogenous levels of EDA protein.
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source	Monoclonal, Mouse,IgG
Purification	The antibody was affinity-purified from mouse antiserum by affinity-chromatography using epitope-specific immunogen.
Dilution	WB 1:500-1:2000
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Synonyms	ED1;ectodysplasin A;Ectodysplasin-A;membrane form;secreted form;EDA;EDA2; Ectodermal dysplasia protein;EDA protein
Observed Band	42kD
Calculated Molecular Weight	42kD
Cell Pathway	Cell membrane ; Single-pass type II membrane protein .; [Ectodysplasin-A, secreted form]: Secreted .
Tissue Specificity	Not abundant; expressed in specific cell types of ectodermal (but not mesodermal) origin of keratinocytes, hair follicles, sweat glands. Also in adult heart, liver, muscle, pancreas, prostate, fetal liver, uterus, small intestine and umbilical chord.
Function	alternative products:Additional isoforms seem to exist,disease:Defects in EDA are a cause of hypodontia [MIM:300606]. Hypodontia is agenesis of two or more permanent teeth without associated systemic disorders. Hypodontia due to EDA defects is an X-linked recessive disorder. Affected individuals have normal hair, skin, and nails, but lack primary and permanent teeth.,disease:Defects in EDA are the cause of ectodermal dysplasia, type 1 (ED1) [MIM:305100]; also known as Christ-Siemens-Touraine syndrome or X-linked hypohidrotic ectodermal dysplasia (XLHED). Ectodermal dysplasia defines a heterogeneous group of



disorders due to abnormal development of two or more ectodermal structures. ED1 is a disease characterized by sparse hair (atrachosis or hypotrichosis), abnormal or missing teeth and the inability to sweat due to the absence of sweat glands. ED1 is the most common form of over 150 cli

Background

The protein encoded by this gene is a type II membrane protein that can be cleaved by furin to produce a secreted form. The encoded protein, which belongs to the tumor necrosis factor family, acts as a homotrimer and may be involved in cell-cell signaling during the development of ectodermal organs. Defects in this gene are a cause of ectodermal dysplasia, anhidrotic, which is also known as X-linked hypohidrotic ectodermal dysplasia. Several transcript variants encoding many different isoforms have been found for this gene. [provided by RefSeq, Jul 2008],

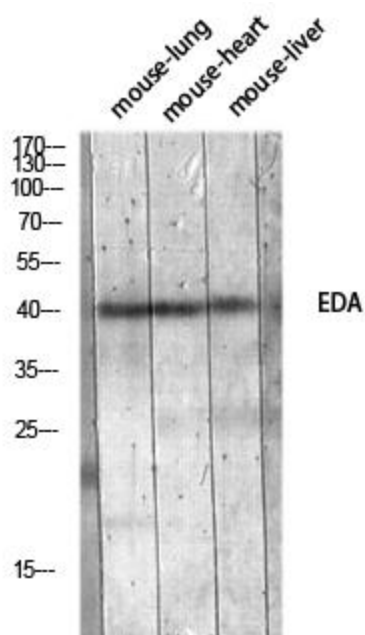
matters needing attention

Avoid repeated freezing and thawing!

Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

Products Images



Western Blot analysis of various cells using EDA Monoclonal Antibody