Product name:	FoxE1 Rabbit Polyclonal Antibody
Cat number:	ABN11084
Conjugate:	Unconjugated
Size:	100µL
Clone:	Polyclonal
Concentration:	1mg/ml
Host:	Rabbit
Isotype:	IgG
Immunogen:	The antiserum was produced against synthesized peptide derived from human TTF2. AA range:10-59
Reactivity:	Human,Mouse,Rat
Applications:	WB 1:500-1:2000,IHC 1:100-1:300,ICC/IF 1:50-1:200,ELISA 1:5000-1:20000
Molecular Weight:	34kDa
Purification:	Affinity purification
Form:	Liquid
Buffer:	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% New type preservative N.
Storage:	Store at 4°C short term. Aliquot and store at -20°C for 12 months. Avoid freeze/thaw cycles.
Background:	This intronless gene belongs to the forkhead family of transcription factors, which is characterized by a distinct forkhead domain. This gene functions as a thyroid transcription factor which likely plays a crucial role in thyroid morphogenesis. Mutations in this gene are associated with congenital hypothyroidism and cleft palate with thyroid dysgenesis. The map localization of this gene suggests it may also be a candidate gene for squamous cell epithelioma and hereditary sensory neuropathy type I. [provided by RefSeq, Jul 2008],disease:Defects in FOXE1 are the cause of Bamforth-Lazarus syndrome [MIM:241850]. A disease associated with thyroid agenesis, cleft palate and choanal atresia.,function:Probable transcription factor. Could be involved in thyroid gland organogenesis.,polymorphism:An alanine stretch that varies from 12 to 19 residues is present. This polymorphisms can be used as a marker to study the role of FOXE1 in other cases of thyroid dysgenesis, especially in familial cases.,PTM:Phosphorylated.,sequence caution:Several conflicts.,similarity:Contains 1 fork-head DNA-binding domain.,tissue specificity:Detected in adult brain, placenta, lung, liver, skeletal muscle, kidney, pancreas, heart, colon, small intestine testis and thymus. Expression was strongest in heart and pancreas.,