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### anti-GTF2E1 antibody (N-Term)

2 Images



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Overview	
Quantity:	400 μL
Target:	GTF2E1
Binding Specificity:	AA 93-121, N-Term
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This GTF2E1 antibody is un-conjugated
Application:	Western Blotting (WB), Immunohistochemistry (Paraffin-embedded Sections) (IHC (p))
Product Details	
Immunogen:	This GTF2E1 antibody is generated from rabbits immunized with a KLH conjugated synthetic
	peptide between 93-121 amino acids from the N-terminal region of human GTF2E1.
Isotype:	lg Fraction
Purification:	This antibody is purified through a protein A column, followed by peptide affinity purification.
Target Details	
Target:	GTF2E1
Alternative Name:	GTF2E1 (GTF2E1 Products)
Background:	Recruits TFIIH to the initiation complex and stimulates the RNA polymerase II C-terminal
	domain kinase and DNA-dependent ATPase activities of TFIIH. Both TFIIH and TFIIE are
	required for promoter clearance by RNA polymerase.

#### **Target Details**

Molecular Weight:	49 kDa
Gene ID:	2960
UniProt:	P29083

#### **Application Details**

Application Notes:	For WB starting dilution is: 1:1000
	For IHC-P starting dilution is: 1:10~50
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Concentration:	0.35 mg/mL

Buffer: Supplied in PBS with 0.09 % (W/V) sodium azide.

Preservative: Sodium azide

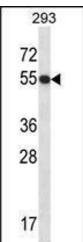
Precaution of Use: This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.

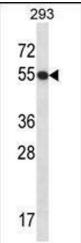
Storage: 4 °C,-20 °C

Storage Comment: Store at 4°C for three months and -20°C, stable for up to one year. As with all antibodies care

should be taken to avoid repeated freeze thaw cycles. Antibodies should not be exposed to

prolonged high temperatures.





# 134597201

#### **Western Blotting**

Image 1. Western blot analysis in 293 cell line lysates (35ug/lane).

#### **Immunohistochemistry**

Image 2. GTF2E1 Antibody immunohistochemistry analysis in formalin fixed and paraffin embedded human stomach tissue followed by peroxidase conjugation of the secondary antibody and DAB staining.