

DNAxPAb

Hard-to-Find Antibody

## TFF1 DNAxPab

Catalog # H00007031-W01P Size 200 ug

Specification	
Product Description	Rabbit polyclonal antibody raised against a full-length human TFF1 DNA using DNAx™ Immune tech nology.
Technology	DNAx™ Immune
Immunogen	Full-length human DNA
Sequence	MATMENKVICALVLVSMLALGTLAEAQTETCTVAPRERQNCGFPGVTPSQCANKGCCFDDTVRG VPWCFYPNTIDVPPEEECEF
Host	Rabbit
Reactivity	Human
Purification	Protein A
Quality Control Testing	Antibody reactive against mammalian transfected lysate.
Storage Buffer	In 1x PBS, pH 7.4
Storage Instruction	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

## **Applications**

Western Blot (Transfected lysate)

Protocol Download

- Immunofluorescence (Transfected cell)
- Flow Cytometry (Transfected cell)



Gene Info — TFF1	
Entrez GenelD	7031
GeneBank Accession#	NM_003225.2
Protein Accession#	NP_003216.1
Gene Name	TFF1
Gene Alias	BCEI, D21S21, HP1.A, HPS2, pNR-2, pS2
Gene Description	trefoil factor 1
Omim ID	113710
Gene Ontology	<u>Hyperlink</u>
Gene Summary	Members of the trefoil family are characterized by having at least one copy of the trefoil motif, a 40 -amino acid domain that contains three conserved disulfides. They are stable secretory proteins e xpressed in gastrointestinal mucosa. Their functions are not defined, but they may protect the muc osa from insults, stabilize the mucus layer, and affect healing of the epithelium. This gene, which is expressed in the gastric mucosa, has also been studied because of its expression in human tumo rs. This gene and two other related trefoil family member genes are found in a cluster on chromos ome 21. [provided by RefSeq
Other Designations	breast cancer estrogen-inducible sequence gastrointestinal trefoil protein pS2

## Disease

- Cerebral Hemorrhage
- Chronic Disease
- Genetic Predisposition to Disease
- Hypertension
- Intracranial Hemorrhages
- Kidney Calculi
- Kidney Diseases
- Lung Neoplasms
- Pulmonary Disease



- Stroke
- Subarachnoid Hemorrhage
- Urinary Bladder Neoplasms
- Werner syndrome