

DNAxPAb

Hard-to-Find
Antibody

UTS2 DNAxPab

Catalog # H00010911-W01P

Size 200 ug

Specification

Product Description	Rabbit polyclonal antibody raised against a full-length human UTS2 DNA using DNAx™ Immune technology.
Technology	DNAx™ Immune
Immunogen	Full-length human DNA
Sequence	MYKLASCCLLFIGFLNPLLSLPLLDREISFQLSAPHEDARLTPEELERASLLQILPEMLGAERGDILRKADSSTNIFNPRGNLRKFQDFSGQDPNILLSHLLARWKPYPKKRETPDCFWKYCV
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 — UTS2

Entrez GeneID [10911](#)

GeneBank Accession# [NM_006786.2](#)

Protein Accession# [NP_006777.1](#)

Gene Name UTS2

Gene Alias PRO1068, U-II, UCN2, UII

Gene Description urotensin 2

Omim ID [604097](#)

Gene Ontology [Hyperlink](#)

Gene Summary This gene encodes a mature peptide that is an active cyclic heptapeptide absolutely conserved from lamprey to human. The active peptide acts as a vasoconstrictor and is expressed only in brain tissue. Despite the gene family name similarity, this gene is not homologous to urocortin, a member of the sauvagine/corticotropin-releasing factor/urotensin I family. Most of the proprotein is cleaved to make the mature peptide. Transcript variants encoding different preproprotein isoforms have been described for this gene. [provided by RefSeq]

Other Designations OTTHUMP00000001357|OTTHUMP00000001358

Pathway

- [Neuroactive ligand-receptor interaction](#)

Disease

- [Diabetes](#)
- [Diabetes Mellitus](#)
- [Genetic Predisposition to Disease](#)
- [Glucose Intolerance](#)
- [Hypercholesterolemia](#)
- [Hypertension](#)

- [Insulin Resistance](#)
- [Kidney Failure](#)
- [Metabolic Syndrome X](#)
- [Myocardial Infarction](#)
- [Pre-Eclampsia](#)
- [Renal Insufficiency](#)
- [Werner syndrome](#)