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See the following pages for more information!



Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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CTSK 293T Cell Transient Overexpression Lysate(Denatured)

Catalog # : H00001513-T01

規格 : [100 uL]

List All

Specification

Transfected Cell Line: 293T

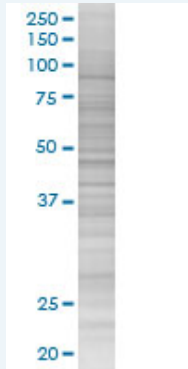
Plasmid: pCMV-CTSK full-length

Host: Human

Theoretical MW (kDa): 36.3

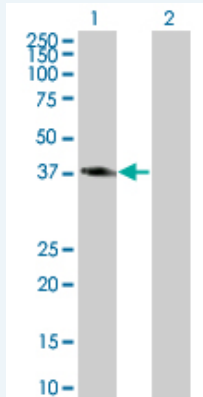
Quality Control Testing: Transient overexpression cell lysate was tested with Anti-CTSK antibody (H00001513-B01) by Western Blots.

SDS-PAGE Gel



CTSK transfected lysate.

Western Blot



Lane 1: CTSK transfected lysate (36.3 KDa)

Lane 2: Non-transfected lysate.

Storage Buffer: 1X Sample Buffer (50 mM Tris-HCl, 2% SDS, 10% glycerol, 300 mM 2-mercaptoethanol, 0.01% Bromophenol blue)

Storage Instruction: Store at -80°C. Aliquot to avoid repeated freezing and thawing.

MSDS:  [Download](#)

Applications

Western Blot

Gene Information

Entrez GeneID: [1513](#)

GeneBank Accession#: [NM_000396.2](#)

Protein Accession#: =

Gene Name: CTSK

Gene Alias: CTS02,CTSO,CTSO1,CTSO2,MGC23107,PKND,PYCD

Gene Description: cathepsin K

Omim ID: [265800](#), [601105](#)

Gene Ontology: [Hyperlink](#)

Gene Summary: The protein encoded by this gene is a lysosomal cysteine proteinase involved in bone remodeling and resorption. This protein, which is a member of the peptidase C1 protein family, is predominantly expressed in osteoclasts. However, the encoded protein is also expressed in a significant fraction of human breast cancers, where it could contribute to tumor invasiveness. Mutations in this gene are the cause of pycnodysostosis, an autosomal recessive disease characterized by osteosclerosis and short stature. This gene may be subject to RNA editing. [provided by RefSeq]

Other Designations: OTTHUMP00000032938,cathepsin O1,cathepsin O2,cathepsin X

Gene Pathway

[Lysosome](#)

Related Disease

[Alzheimer Disease](#) [Alzheimer disease](#) [Body Weight](#) [Cardiovascular Diseases](#)
[Diabetes Complications](#) [Diabetes Mellitus, Type 2](#) [Edema](#)
[Genetic Predisposition to Disease](#) [Metabolic Syndrome X](#) [Neoplasms](#) [Osteoporosis](#)
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