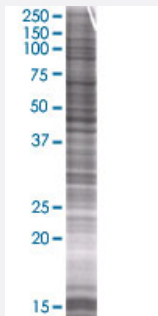


# DFNA5 293T Cell Transient Overexpression Lysate(Denatured)

Catalog # H00001687-T01

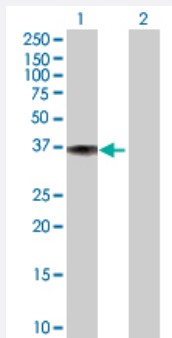
Size 100 uL

## Applications



### SDS-PAGE Gel

DFNA5 transfected lysate.



### Western Blot

Lane 1: DFNA5 transfected lysate ( 32.56 KDa)

Lane 2: Non-transfected lysate.

## Specification

**Transfected Cell Line** 293T

**Plasmid** pCMV-DFNA5 full-length

**Host** Human

**Theoretical MW (kDa)** 32.56

**Quality Control Testing** Transient overexpression cell lysate was tested with Anti-DFNA5 antibody ([H00001687-B01](#)) by Western Blots.  
 SDS-PAGE Gel  
 DFNA5 transfected lysate.  
 Western Blot  
 Lane 1: DFNA5 transfected lysate ( 32.56 KDa)  
 Lane 2: Non-transfected lysate.

<b>Storage Buffer</b>	1X Sample Buffer (50 mM Tris-HCl, 2% SDS, 10% glycerol, 300 mM 2-mercaptoethanol, 0.01% Bromophenol blue)
<b>Storage Instruction</b>	Store at -80°C. Aliquot to avoid repeated freezing and thawing.

## Applications

- Western Blot

## Gene Info — DFNA5

<b>Entrez GeneID</b>	<a href="#">1687</a>
<b>GeneBank Accession#</b>	<a href="#">BC019689</a>
<b>Protein Accession#</b>	-
<b>Gene Name</b>	DFNA5
<b>Gene Alias</b>	ICERE-1
<b>Gene Description</b>	deafness, autosomal dominant 5
<b>Omim ID</b>	<a href="#">600994</a> <a href="#">608798</a>
<b>Gene Ontology</b>	<a href="#">Hyperlink</a>
<b>Gene Summary</b>	Hearing impairment is a heterogeneous condition with over 40 loci described. The protein encoded by this gene is expressed in fetal cochlea, however, its function is not known. Nonsyndromic hearing impairment is associated with a mutation in this gene. Three transcript variants encoding two different isoforms have been found for this gene. [provided by RefSeq]
<b>Other Designations</b>	deafness, autosomal dominant 5 protein inversely correlated with estrogen receptor expression 1 nonsyndromic hearing impairment protein

## Disease

- [Colorectal Neoplasms](#)
- [Genetic Predisposition to Disease](#)
- [Tobacco Use Disorder](#)