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- Diagnostik & molekulare Diagnostik
- Laborgeräte & Service

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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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FANCA 293T Cell Transient Overexpression Lysate(Denatured)

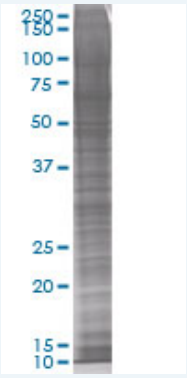
Catalog # : H00002175-T01 **規格 :** [100 uL]

List All

Specification

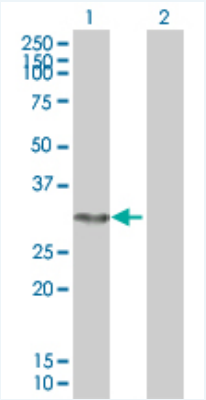
Transfected Cell Line:	293T
Plasmid:	pCMV-FANCA full-length
Host:	Human
Theoretical MW (kDa):	32.78
Quality Control Testing:	Transient overexpression cell lysate was tested with Anti-FANCA antibody (H00002175-B01) by Western Blots.

SDS-PAGE Gel



FANCA transfected lysate

Western Blot



Lane 1: FANCA transfected lysate (32.78 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

Application Image

Western Blot

Western Blot

Gene Information

Entrez GeneID:	2175
GeneBank Accession#:	BC008979
Protein Accession#:	AAH08979
Gene Name:	FANCA
Gene Alias:	FA,FA-H,FA1,FAA,FACA,FAH,FANCH,MGC75158
Gene Description:	Fanconi anemia, complementation group A
Omim ID:	227650, 607139
Gene Ontology:	Hyperlink
Gene Summary:	The Fanconi anemia complementation group (FANC) currently includes FANCA, FANCB, FANCC, FANCD1 (also called BRCA2), FANCD2, FANCE, FANCF, FANCG, FANCI, FANCJ (also called BRIP1), FANCL, FANCM and FANCN (also called PALB2). The previously defined group FANCH is the same as FANCA. Fanconi anemia is a genetically heterogeneous recessive disorder characterized by cytogenetic instability, hypersensitivity to DNA crosslinking agents, increased chromosomal breakage, and defective DNA repair. The members of the Fanconi anemia complementation group do not share sequence similarity; they are related by their assembly into a common nuclear protein complex. This gene encodes the protein for complementation group A. Alternative splicing results in multiple transcript variants encoding different isoforms. Mutations in this gene are the most common cause of Fanconi anemia. [provided by RefSeq
Other Designations:	Fanconi anemia, complementation group H,Fanconi anemia, type 1

Related Disease

Adenocarcinoma Breast cancer Breast Neoplasms Cardiovascular Diseases Diabetes Mellitus, Type 2 Disease Progression Edema Esophageal Neoplasms Fanconi Anemia Genetic Predisposition to Disease Kidney Failure, Chronic Lung Neoplasms Meningeal Neoplasms Meningioma Neoplasms, Glandular and Epithelial Ovarian cancer Ovarian Neoplasms Pancreatic Neoplasms Papillomavirus Infections

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