

# Recombinant Human Dynein intermediate chain 1/DNAI1 protein ab153279

[1 References](#)   [1 Image](#)

Description	
Product name	Recombinant Human Dynein intermediate chain 1/DNAI1 protein
Expression system	Wheat germ
Protein length	Full length protein
Animal free	No
Nature	Recombinant
Species	Human
Sequence	MIPASAKSPHKQPHKQSSISIGRGTRKRDEDSGTEVGEGTD EWAQSKATVR PPDQLELTD AELKEEFTRILTANNPHAPQNVRYSFKEGTY KPIGFVNQL AVHYTQVGNLIPKDSDEGRRQHYRDELVAGSQESVKVISE TGNLEEDEEP KELETEPGSQTDVPAAGAAEKVTEEELMTPKQPKERKLT NQFNFSERASQ TCNNPVRDRECQTEPPPRTNFSATANQWEYDAYVEELE KQEKTKKEKA KTPVAKKSGKMAMRKLTSMESQTDDLIKLSQAAKIMERM VNQNTYDDIAQ DFKYDDAADEYRDQVGTLLPLWK FQNDKAKRLSVTALC WNPKYRDLFAV GYGSYDFMKQSRGM LLLYSLKNPSFPEYMFSSNSGVMCL DIHVDHPYLVA VGHYDGNVAINLKKPHSQPSFCSSAKSGKHSDPVWQV KWQKDDMDQNLN FFSVSSDGRIVSWTLVKRKL VHIDVIKLKVEGSTTEVPEGL QLHQVGC GT AFDFHKEIDY MFLVGTEEGKIYKCSKSYSSQFLDTYDAHN MSVDTVSWNP YHTKVFMS C SSDWTVKIWDHTIKTPMFIYDLNSAVGDVAW APYSSTVFAA VTTDGAHIFDLANKYEAICNQPVAAKKNRLTHVQFNLIHPI IIVGDDR GHIISLKLSPNLRKMPKEKKGQEVQKGPAVEIAKLDKLLNL

<b>Amino acids</b>	1 to 699
<b>Tags</b>	GST tag N-Terminus

## Specifications

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Our **Abpromise guarantee** covers the use of **ab153279** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

<b>Applications</b>	ELISA Western blot
<b>Form</b>	Liquid
<b>Additional notes</b>	This product was previously labelled as Dynein intermediate chain 1.

## Preparation and Storage

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<b>Stability and Storage</b>	Shipped on dry ice. Upon delivery aliquot and store at -80°C. Avoid freeze / thaw cycles. pH: 8.00 Constituents: 0.31% Glutathione, 0.79% Tris HCl
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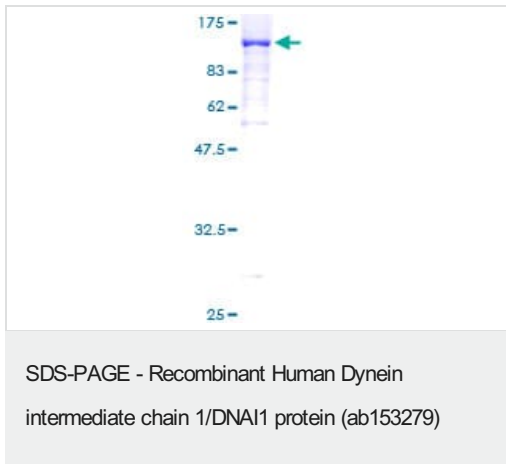
## General Info

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<b>Function</b>	Part of the dynein complex of respiratory cilia.
<b>Involvement in disease</b>	Defects in DNAI1 are the cause of primary ciliary dyskinesia type 1 (CILD1) [MIM:244400]. CILD1 is an autosomal recessive disorder characterized by axonemal abnormalities of motile cilia. Respiratory infections leading to chronic inflammation and bronchiectasis are recurrent, due to defects in the respiratory cilia; reduced fertility is often observed in male patients due to abnormalities of sperm tails. Half of the patients exhibit situs inversus, due to dysfunction of monocilia at the embryonic node and randomization of left-right body asymmetry. Primary ciliary dyskinesia associated with situs inversus is referred to as Kartagener syndrome. Defects in DNAI1 are the cause of Kartagener syndrome (KTGS) [MIM:244400]. KTGS is an autosomal recessive disorder characterized by the association of primary ciliary dyskinesia with situs inversus. Clinical features include recurrent respiratory infections, bronchiectasis, infertility, and lateral transposition of the viscera of the thorax and abdomen. The situs inversus is most often total, although it can be partial in some cases (isolated dextrocardia or isolated transposition of abdominal viscera).
<b>Sequence similarities</b>	Belongs to the dynein intermediate chain family. Contains 5 WD repeats.
<b>Cellular localization</b>	Cytoplasm > cytoskeleton > cilium axoneme.

## Images

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ab153279 on a 12.5% SDS-PAGE stained with Coomassie Blue.

**Please note:** All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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