



# Rabbit Anti-Human IYD Polyclonal antibody (DPABH-07839)

This product is for research use only and is not intended for diagnostic use.

## PRODUCT INFORMATION

<b>Immunogen</b>	IYD fusion protein, sequence: EPRTAEARPWVDEDLKDSSDLHQAEEADAWEQSEENVEHIPFSHNHYPEKEMVKRSQE FYELLNKRRSVRFISNEQVPMEVIDNVIRTAGTAPSGAHTEPWTFVVVKDPDVKHKIRKI IEEEEEINYMKRMGHRWVTDLKKLRTNWIKEYLDTAPILILIFKQVHGFAANGKKKVHYY NEISVSIACGILLAALQNAGLVTVTTTPLNCGPRLRVLLGRPAHEKLPMLLPVGYPSEKA TVPDLKRKPLDQIMVTV (33-289aa encoded by BC056253)
<b>Isotype</b>	IgG
<b>Source/Host</b>	Rabbit
<b>Species Reactivity</b>	Human
<b>Purification</b>	Antigen affinity purification
<b>Conjugate</b>	Unconjugated
<b>Applications</b>	WB, ELISA
<b>Positive Control</b>	human liver tissue
<b>Format</b>	Liquid
<b>Size</b>	50 µl, 100 µl
<b>Buffer</b>	PBS with 0.02% sodium azide and 50% glycerol pH 7.3.
<b>Preservative</b>	0.02% Sodium Azide
<b>Storage</b>	Store at -20°C. Aliquoting is unnecessary for -20°C storage.

# BACKGROUND

## Introduction

This gene encodes an enzyme that catalyzes the oxidative NADPH-dependent deiodination of mono- and diiodotyrosine, which are the halogenated byproducts of thyroid hormone production. The N-terminus of the protein functions as a membrane anchor. Mutations in this gene cause congenital hypothyroidism due to dysmorphogenesis type 4, which is also referred to as deiodinase deficiency, or iodotyrosine dehalogenase deficiency, or thyroid morphogenesis type 4. Alternative splicing results in multiple transcript variants.

## Keywords

IYD; iodotyrosine deiodinase; TDH4; DEHAL1; C6orf71; dJ422F24.1; iodotyrosine dehalogenase 1; IYD-1;

# GENE INFORMATION

## Entrez Gene ID

[389434](#)

## UniProt ID

[Q6PHW0](#)