

# Datasheet for ABIN857419

# **IMPDH2 ELISA Kit**





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Quantity:	96 tests
Target:	IMPDH2
Reactivity:	Human
Method Type:	Sandwich ELISA
Detection Range:	23.5-1500 pg/mL
Minimum Detection Limit:	23.5 pg/mL
Application:	ELISA

## **Product Details**

Sample Type:	Serum, Plasma, Cell Culture Supernatant, Tissue Homogenate
Analytical Method:	Quantitative
Detection Method:	Colorimetric
Specificity:	This assay has high sensitivity and excellent specificity for detection of Human IMP (Inosine Monophosphate) Dehydrogenase 2 (IMPDH2).
Cross-Reactivity (Details):	Limited by current skills and knowledge, it is impossible for us to complete the cross-reactivity detection between the target antigen and all analogues for other species. Therefore, cross reaction may still exist.
Sensitivity:	5.8 pg/mL
Components:	<ul> <li>Assay plate (12 × 8 coated Microwells)</li> <li>Standard (freeze dried)</li> </ul>

• Biotin-antibody (100 × concentrate)

- HRP-avidin (100 × concentrate)
- · Biotin-antibody Diluent
- HRP-avidin Diluent
- · Sample Diluent
- Wash Buffer (25 x concentrate)
- · TMB Substrate
- · Stop Solution
- · Adhesive Strip (for 96 wells)
- · Instruction manual

## **Target Details**

Target:	IMPDH2	
Alternative Name:	IMP (Inosine Monophosphate) Dehydrogenase 2 (IMPDH2) (IMPDH2 Products)	
Background:	Synonyms: hCG_2002013, IMPD2, IMPDH-II, IMP oxireductase 2 inosine 5 phosphate dehydrogenase 2 inosine monophosphate dehydrogenase 2 inosine monophosphate dehydrogenase type II	
UniProt:	P12268	
Pathways:	Ribonucleoside Biosynthetic Process, SARS-CoV-2 Protein Interactome	

### **Application Details**

### **Application Notes:**

- The supplier is only responsible for the kit itself, but not for the samples consumed during the assay. The user should calculate the possible amount of the samples used in the whole test.
   Please reserve sufficient samples in advance.
- Samples to be used within 5 days may be stored at 2-8°C, otherwise samples must be stored at -20°C (≤ 1 month) or -80°C (≤ 2 months) to avoid loss of bioactivity and contamination.
- Grossly hemolyzed samples are not suitable for use in this assay.
- If the samples are not indicated in the manual, a preliminary experiment to determine the validity of the kit is necessary.
- Please predict the concentration before assaying. If values for these are not within the range
  of the standard curve, users must determine the optimal sample dilutions for their particular
  experiments.
- Tissue or cell extraction samples prepared by chemical lysis buffer may cause unexpected ELISA results due to the impacts of certain chemicals.
- Owing to the possibility of mismatching between antigens from another resource and antibodies used in this supplier's kits (e.g., antibody targets conformational epitope rather than linear epitope), some native or recombinant proteins from other manufacturers may not be recognized by this supplier's products.
- · Influenced by factors including cell viability, cell number and cell sampling time, samples

from cell culture supernatant may not be recognized by the kit.

 Fresh samples without long time storage are recommended for the test. Otherwise, protein degradation and denaturalization may occur in those samples and finally lead to wrong results.

#### Comment:

Detection wavelength: 450 nm

Information on standard material:

Depending on the antigen to be detected, standards can be either native or recombinant protein. The recombinant proteins are being expressed in CHO cells in most cases. Please inquire for more information. The formulation of auxiliary material in the standard is considered proprietary information, however it does not contain any poisonous substance. Proclin 300 (1:3000) is used as preservative.

#### Information on reagents:

In most cases the stop solution provided is 1 N H2SO4. The formulation of wash solution is proprietary information. None of the components contain (sodium) azide, thimerosal, 2-mercaptoethanol (2-ME) or any other poisonous materials. For the sandwich method kits, the sample diluent, antibody diluent, enzyme diluent and standard all contain BSA.

#### Information on antibodies:

The antibodies provided in different kits vary in regards to clonality and host. Some antibodies are affinity purified, some are Protein A

### Assay Time:

1 - 4.5 h

### Plate:

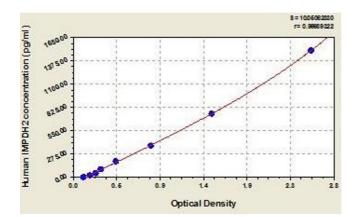
Pre-coated

### Protocol:

This assay employs the quantitative sandwich enzyme immunoassay technique. Antibody specific for IMP (Inosine Monophosphate) Dehydrogenase 2 (IMPDH2) has been pre-coated onto a microplate. Standards and samples are pipetted into the wells and any IMP (Inosine Monophosphate) Dehydrogenase 2 (IMPDH2) present is bound by the immobilized antibody. After removing any unbound substances, a biotin-conjugated antibody specific for IMP (Inosine Monophosphate) Dehydrogenase 2 (IMPDH2) is added to the wells. After washing, avidin conjugated Horseradish Peroxidase (HRP) is added to the wells. Following a wash to remove any unbound avidin-enzyme reagent, a substrate solution is added to the wells and color develops in proportion to the amount of IMP (Inosine Monophosphate) Dehydrogenase 2 (IMPDH2) bound in the initial step. The color development is stopped and the intensity of the

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## **ELISA**

Image 1. Typical standard curve