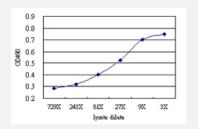
# ALDH4A1 (Human) Matched Antibody Pair

Catalog # H00008659-AP51 Size 1 Set

## Applications



Sandwich ELISA detection sensitivity ranging from approximately 729x to 3x dilution of the ALDH4A1 293T overexpression lysate (non-denatured).

| Specification                    |  |
|----------------------------------|--|
| Product Description              | This antibody pair set comes with a matched antibody pair to detect and quantify the protein level of human ALDH4A1.   |
| Reactivity                       | Human  |
| Interspecies Antigen<br>Sequence | Mouse (91); Rat (91)   |
| Quality Control Testing          | Standard curve using ALDH4A1 293T overexpression lysate (non-denatured) as an analyte.<br>Sandwich ELISA detection sensitivity ranging from approximately 729x to 3x dilution of the ALDH4A<br>1 293T overexpression lysate (non-denatured).                   |
| Supplied Product                 | Antibody pair set content:<br>1. Capture antibody: mouse monoclonal anti-ALDH4A1 (100 ug)<br>2. Detection antibody: rabbit purified polyclonal anti-ALDH4A1 (50 ug)<br>*Reagents are sufficient for at least 3-5 x 96 well plates using recommended protocols. |
| Storage Instruction              | Store reagents of the antibody pair set at -20°C or lower. Please aliquot to avoid repeated freeze tha w cycle. Reagents should be returned to -20°C storage immediately after use.  |

### Applications

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• ELISA Pair (Transfected lysate)

Protocol Download

| Gene Info — ALDH4A1 |  |
|---------------------|--|
| Entrez GenelD       | <u>8659</u>  |
| Gene Name           | ALDH4A1  |
| Gene Alias          | ALDH4, P5CD, P5CDh, P5CDhL, P5CDhS   |
| Gene Description    | aldehyde dehydrogenase 4 family, member A1   |
| Omim ID             | <u>239510 606811</u>   |
| Gene Ontology       | Hyperlink  |
| Gene Summary        | This protein belongs to the aldehyde dehydrogenase family of proteins. This enzyme is a mitocho ndrial matrix NAD-dependent dehydrogenase which catalyzes the second step of the proline degr adation pathway, converting pyrroline-5-carboxylate to glutamate. Deficiency of this enzyme is ass ociated with type II hyperprolinemia, an autosomal recessive disorder characterized by accumulat ion of delta-1-pyrroline-5-carboxylate (P5C) and proline. Alternatively spliced transcript variants e ncoding different isoforms have been identified for this gene. [provided by RefSeq |
| Other Designations  | OTTHUMP0000002544 OTTHUMP0000002545 P5C dehydrogenase aldehyde dehydrogenas<br>e 4A1 mitochondrial delta-1-pyrroline 5-carboxylate dehydrogenase   |

### Pathway

- <u>Alanine</u>
- <u>Arginine and proline metabolism</u>
- Metabolic pathways

#### Disease

- <u>Adenocarcinoma</u>
- Esophageal Neoplasms
- Hearing Loss