

# DDO oxidizes D-Asp to OA

D'Eustachio, P., Jassal, B.

European Bioinformatics Institute, New York University Langone Medical Center, Ontario Institute for Cancer Research, Oregon Health and Science University.

The contents of this document may be freely copied and distributed in any media, provided the authors, plus the institutions, are credited, as stated under the terms of [Creative Commons Attribution 4.0 International \(CC BY 4.0\) License](#). For more information see our [license](#).

29/11/2020

## Introduction

Reactome is open-source, open access, manually curated and peer-reviewed pathway database. Pathway annotations are authored by expert biologists, in collaboration with Reactome editorial staff and cross-referenced to many bioinformatics databases. A system of evidence tracking ensures that all assertions are backed up by the primary literature. Reactome is used by clinicians, geneticists, genomics researchers, and molecular biologists to interpret the results of high-throughput experimental studies, by bioinformaticians seeking to develop novel algorithms for mining knowledge from genomic studies, and by systems biologists building predictive models of normal and disease variant pathways.

The development of Reactome is supported by grants from the US National Institutes of Health (P41 HG003751), University of Toronto (CFREF Medicine by Design), European Union (EU STRP, EMI-CD), and the European Molecular Biology Laboratory (EBI Industry program).

## Literature references

- Fabregat, A., Sidiropoulos, K., Viteri, G., Forner, O., Marin-Garcia, P., Arnau, V. et al. (2017). Reactome pathway analysis: a high-performance in-memory approach. *BMC bioinformatics*, 18, 142. [↗](#)
- Sidiropoulos, K., Viteri, G., Sevilla, C., Jupe, S., Webber, M., Orlic-Milacic, M. et al. (2017). Reactome enhanced pathway visualization. *Bioinformatics*, 33, 3461-3467. [↗](#)
- Fabregat, A., Jupe, S., Matthews, L., Sidiropoulos, K., Gillespie, M., Garapati, P. et al. (2018). The Reactome Pathway Knowledgebase. *Nucleic Acids Res*, 46, D649-D655. [↗](#)
- Fabregat, A., Korninger, F., Viteri, G., Sidiropoulos, K., Marin-Garcia, P., Ping, P. et al. (2018). Reactome graph database: Efficient access to complex pathway data. *PLoS computational biology*, 14, e1005968. [↗](#)

Reactome database release: 74

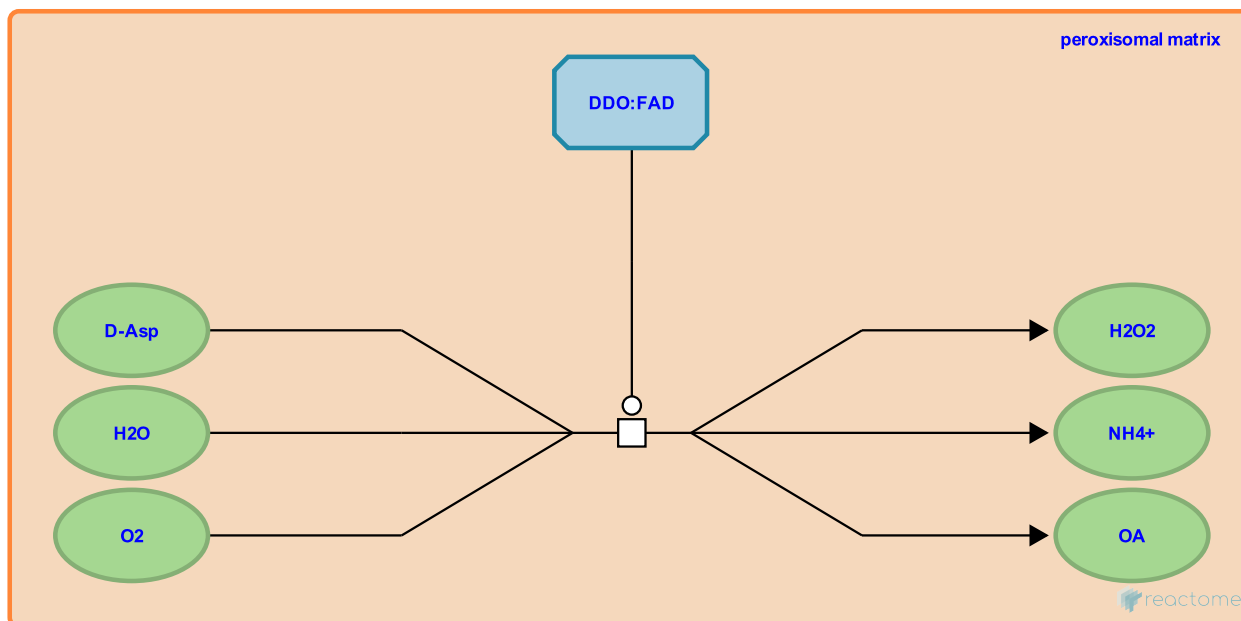
This document contains 1 reaction ([see Table of Contents](#))

## DDO oxidizes D-Asp to OA ↗

**Stable identifier:** R-HSA-6810076

**Type:** transition

**Compartments:** peroxisomal matrix



Peroxisomal DDO (D-aspartate oxidase) catalyzes the oxidation of D-Asp (D-aspartate) to OA (oxaloacetate) with the formation of H<sub>2</sub>O<sub>2</sub>. The human enzyme is a monomer with an FAD cofactor (Katane et al. 2010, 2015; Setoyama & Miura 1997), as is its well-characterized bovine homolog (Negri et al. 1992). Its peroxisomal location is inferred from studies in cultured cells of fusion proteins containing the carboxy-terminal peptide sequence of DDO (Amery et al. 1998).

### Literature references

- Katane, M., Kawata, T., Nakayama, K., Saitoh, Y., Kaneko, Y., Matsuda, S. et al. (2015). Characterization of the enzymatic and structural properties of human D-aspartate oxidase and comparison with those of the rat and mouse enzymes. *Biol. Pharm. Bull.*, 38, 298-305. ↗
- Amery, L., Brees, C., Baes, M., Setoyama, C., Miura, R., Mannaerts, GP. et al. (1998). C-terminal tripeptide Ser-Asn-Leu (SNL) of human D-aspartate oxidase is a functional peroxisome-targeting signal. *Biochem. J.*, 336, 367-71. ↗
- Katane, M., Saitoh, Y., Hanai, T., Sekine, M., Furuchi, T., Koyama, N. et al. (2010). Thiolactomycin inhibits D-aspartate oxidase: a novel approach to probing the active site environment. *Biochimie*, 92, 1371-8. ↗
- Setoyama, C., Miura, R. (1997). Structural and functional characterization of the human brain D-aspartate oxidase. *J. Biochem.*, 121, 798-803. ↗
- Negri, A., Cecilian, F., Tedeschi, G., Simonic, T., Ronchi, S. (1992). The primary structure of the flavoprotein D-aspartate oxidase from beef kidney. *J. Biol. Chem.*, 267, 11865-71. ↗

### Editions

2015-11-20	Authored, Edited	D'Eustachio, P.
2015-11-30	Reviewed	Jassal, B.