

DCD(63-110) is processed to DCD(63-109)

Hains, DS., Jupe, S., Shamovsky, V.

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 <u>Creative Commons Attribution 4.0 International (CC BY 4.0)</u> <u>License</u>. For more information see our <u>license</u>.

19/05/2024

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. 7
- Sidiropoulos, K., Viteri, G., Sevilla, C., Jupe, S., Webber, M., Orlic-Milacic, M. et al. (2017). Reactome enhanced pathway visualization. *Bioinformatics*, 33, 3461-3467. A
- 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. *オ*

This document contains 1 reaction (see Table of Contents)

DCD(63-110) is processed to DCD(63-109) 7

Stable identifier: R-HSA-6803060

Type: omitted

Compartments: extracellular region



A 47aa dermcidin (DCD)-derived peptide (DCD(63-109), also known as as DCD-1) is an antimicrobial peptide with a negative net charge and acidic pI (Schittek B 2012). Like other antimicrobially active DCD-derived peptides, DCD(63-109) is produced in human eccrine sweat through proteolytic processing of a 110-amino acid (aa) precursor protein (Schittek B et al. 2001; Rieg S et al. 2006). DCD-derived peptides are able to bind to the bacterial surface, however they do not exert their activity by permeabilizing bacterial membranes (Senyürek et al. 2009, Steffen H et al. 2006). The negative net charge of DCD(63-109) did not significantly affected the peptide binding to bacterial-mimetic membranes (Jung et al. 2010, Steffen et al. 2006; Senyurek et al. 2009). Spin-down assays of DCD(63-109) and other DCD peptides revealed that the affinity with which dermcidin binds to bacterial-mimetic membranes is primarily dependent on its amphipathic alpha-helical structure and its length (>30 residues)(Jung et al. 2010).

DCD(63-109) shows a broad spectrum of antimicrobial activity against Gram-positive (Staphylococcus aureus, Enterococcus faecalis, Staphylococcus epidermidis, Listeria monocytogenes) and Gram-negative bacteria (Escherichia coli, Pseudomonas putida, Salmonella typhimurium) as well as Candida albicans (Cipakova I et al. 2006, Lai YP et al. 2005, Schittek B et al. 2001, Steffen H et al. 2006, Vuong C et al. 2004). The activity of the DCD(63-109) was maintained over a broad pH range and in high salt concentrations that resembled the conditions in human sweat (Schittek B et al. 2001).

Literature references

- Baechle, D., Kalbacher, H., Steffen, H., Cansier, A., Flad, T., Mueller, GA. et al. (2006). Cathepsin D is present in human eccrine sweat and involved in the postsecretory processing of the antimicrobial peptide DCD-1L. J. Biol. Chem., 281, 5406-15. ↗
- Schittek, B. (2012). The multiple facets of dermcidin in cell survival and host defense. J Innate Immun, 4, 349-60. 🛪
- Kimura, A., Seeber, S., Rieg, S., Schittek, B., Kalbacher, H., Steffen, H. et al. (2006). Generation of multiple stable dermcidin-derived antimicrobial peptides in sweat of different body sites. J. Invest. Dermatol., 126, 354-65.

Editions

2015-10-05	Authored	Shamovsky, V.
2016-04-15	Reviewed	Jupe, S.
2016-08-02	Reviewed	Hains, DS.
2016-08-15	Edited	Shamovsky, V.