

Liste des publications

2020 Références trouvées : 1

Guyot, N., **Meudal, H.**, Trapp, S., Iochmann, S., Silvestre, A., Jousset, G., Labas, V., Reverdiau, P., Loth, K., Hervé, V., Aucagne, V., Delmas, A.F., Rehault-Godbert, S., Landon, C (2020)

[Structure, function and evolution of Gga-AvBD11, the archetype of a new structural avian-double- \$\beta\$ -defensin family](#)

[Proc. Natl. Acad. Sci. USA.](#) 117, 337-345

2019 Références trouvées : 2

Loth, K., Vergnes, A., Barreto, C., Voisin, S.N., **Meudal, H.**, Da Silva, J., Bressan, A., Belmadi, N., Bachère, E., Aucagne, V., Cazevielle, C., Marchandin, H., Rosa, R.D., Bulet, P., Touqui, L., Delmas, A.F., Destoumieux-Garzon, D (2019)

[The ancestral N-terminal domain of big defensins drives bacteria-triggered assembly into antimicrobial nanonets](#)

[mBio](#) 10(5) e01821-19

Zoukimian, C., **Meudal, H.**, De Waard, S., Ouares, KA., Nicolas, S., Canepari, M., Bérout, R., Landon, C., De Waard, M., Boturyn, D (2019)

[Synthesis by native chemical ligation and characterization of the scorpion toxin AmmTx3](#)

[Bioorganic & medicinal chemistry](#) 27 (1) 247-253

2016 Références trouvées : 1

Bailleul, G., Kravtsoff, A., Joulin-Giet, A., Lecaille, F., Labas, V., **Meudal, H.**, Loth, K., Teixeira-Gomes, AP., Gilbert, FB., Coquet, L., Jouenne, T., Brömme, D., Schouler, C., Landon, C., Lalmanach, G., Lalmanach, AC (2016)

[The unusual resistance of avian defensin AvBD7 to proteolytic enzymes preserves its antibacterial activity](#)

[PLOS ONE](#) 11 (8) 1-20

2015 Références trouvées : 1

Oukhatar, F., **Meudal, H.**, Landon, C., Logothetis, NK., Platas-Iglesias, C., Angelovski, G., Toth, E (2015)

[Macrocyclic Gd\(3+\) complexes with pendant crown ethers designed for binding zwitterionic neurotransmitters](#)

Chemistry 21 (31) 11226-37

2014 Références trouvées : 1

Herve, V., **Meudal, H.**, Labas, V., Réhault-Godbert, S., Gautron, J., Berges, M., Guyot, N., Delmas, A.F., Nys; Y., Landon, C (2014)

[Three-dimensional NMR structure of Hen Egg Gallin \(Chicken Ovodefensin\) reveals a new variation of the \$\beta\$ -defensin fold](#)

J. Biol. Chem. 289 (10) 7211-20

2012 Références trouvées : 1

Derache, C., **Meudal, H.**, Aucagne, V., Mark, KJ., Cadène, M., Delmas, AF., Lalmanach, A.C., Landon, C (2012)

[Initial insights into structure-activity relationships of avian defensins](#)

J. Biol. Chem. 287 (10) 7746-55

2011 Références trouvées : 1

Bure, C., Marceau, P., **Meudal, H.**, & Delmas, A.F (2011)

[Synthesis and analytical investigation of C-terminally modified peptide aldehydes and ketone: application to oxime ligation](#)

J. Pept. Sci. 18 (3) 147-54

2010 Références trouvées : 2

Paquet, F., Loth, K., **Meudal, H.**, Culard, F., Genest, D., & Lancelot, G (2010)

[Refined solution structure and backbone dynamics of the archaeal MC1 protein](#)

FEBS. J. 277 (24) 5133-45

Da Silva, P., Rahioui, I., Laugier, C., Jouvensal, L., **Meudal, H.**, Chouabe, C., Delmas, A.F. & Gressent, F (2010)

[Molecular requirements for the insecticidal activity of the plant peptide PA1b](#)
J. Biol. Chem. 285 (43) 32689-32694

2009 Références trouvées : 2

Aubert, Y., Chassignol, M., Roig, V., Mbemba, G., Weiss, J., **Meudal, H.**, Mouscadet, J.F. & Asseline, U (2009)

[Synthesis and anti-HIV-1 integrase activity of modified dinucleotides](#)
Eur. J. Med. Chem. 44 (12) 5029-5044

Derache, C., Labas, V., Aucagne, V., **Meudal, H.**, Landon, C., Delmas, A.F., Magallon, T. & Lalmanach, A.C (2009)

[Primary Structure and Antibacterial Activity of Chicken Bone Marrow-Derived beta-Defensins](#)
Antimicrob. Agents. Chemother. 53 (11) 4647-4655

2008 Références trouvées : 1

Chauvin, T., Durand, P., Bernier., M., **Meudal, H.**, Doan, BT., Noury, F., Badet, B., Beloeil, JC., Tóth, E (2008)

[Detection of enzymatic activity by PARACEST MRI : a general approach to target a large variety of enzymes](#)
Angew. Chem. Int. Ed. Engl. 47 (23) 4370-4372

2006 Références trouvées : 3

Bilbille, Y., Paquet, F., **Meudal, H.**, Giraud-Panis, MJ., Lancelot, G (2006)

[NMR studies of telomeric nucleoprotein complexes involving the Myb-like domain of the human telomeric protein TRF2](#)
C. R. Chimie. 9 (3-4) 452-458

Landon, C., **Meudal, H.**, Boulanger, N., Bulet, P., Vovelle, F (2006)

[Solution structures of stomoxyn and spinigerin, two insect antimicrobial peptides with an \$\alpha\$ -helical conformation](#)
Biopolymers 81 (2) 92-103

Morellet, N., **Meudal, H.**, Bouaziz, S., Roques, BP(2006)

[Structure of the zinc finger domain encompassing residues 13-51 of the nucleocapsid protein from simian immunodeficiency virus](#)

Biochem. J. 393 (3) 725-732

2003 Références trouvées : 2

Anne, C., Turcaud, S., Quancard, J., Teffo, F., **Meudal, H.**, Fournie-Zaluski, MC., Roques, BP (2003)

[Development of potent inhibitors of botulinum neurotoxin type B](#)

J. Med. Chem. 46 (22) 4648-4656

Anne, C., Blommaert, A., Turcaud, S., Martin, AS., **Meudal, H.**, Roques, BP (2003)

[Thio-derived disulfides as potent inhibitors of botulinum neurotoxin type B : Implications for zinc interaction](#)

Bioorg. Med. Chem. 11 (21) 4655-4660

2002 Références trouvées : 1

Inguibert, N., Coric, P., Poras, H., **Meudal, H.**, Teffot, F., Fournie-Zaluski, MC., Roques, BP (2002)

[Toward an optimal joint recognition of the S-1 ' subsites of endothelin converting enzyme-1 \(ECE-1\), angiotensin converting enzyme \(ACE\), and neutral endopeptidase \(NEP\)](#)

J. Med. Chem. 45 (7) 1477-1486

2000 Références trouvées : 2

Bellier, B., Million, ME., DaNascimento, S., **Meudal, H.**, Kellou, S., Maignet, B., Garbay, C (2000)

[Replacement of glycine with dicarbonyl and related moieties in analogues of the C-terminal pentapeptide of cholecystokinin : CCK2 agonists displaying a novel binding mode](#)

J. Med. Chem. 43 (20) 3614-3623

Chen, HX., Noble, F., Mothe, A., **Meudal, H.**, Coric, P., Danascimento, S., Roques, BP., George, P., Fournie-Zaluski, MC (2000)

**Phosphinic derivatives as new dual enkephalin-degrading enzyme inhibitors :
Synthesis, biological properties, and antinociceptive activities**

J. Med. Chem. 43 (7) 1398-1408

1999 Références trouvées : 3

David, C., Bischoff, L., **Meudal, H.**, Mothe, A., De Mota, N., DaNascimento, S., Llorens-Cortes, C., Fournie-Zaluski, MC., Roques, BP (1999)

Investigation of subsite preferences in aminopeptidase A (EC 3.4.11.7) led to the design of the first highly potent and selective inhibitors of this enzyme

J. Med. Chem. 42 (25) 5197-5211

Druillenec, S., **Meudal, H.**, Roques, BP., Fournie-Zaluski, MC (1999)

Nucleomimetic strategy for the inhibition of HIV-1 nucleocapsid protein NCp7 activities

Bioorg. Med. Chem. Lett. 9 (4) 627-632

Martin, L., Cornille, F., Turcaud, S., **Meudal, H.**, Roques, BP., Fournie-Zaluski, MC (1999)

Metallopeptidase inhibitors of tetanus toxin : A combinatorial approach

J. Med. Chem. 42 (3) 515-525

1998 Références trouvées : 1

Bellier, B., Da Nascimento, S., **Meudal, H.**, Gincel, E., Roques, BP., Garbay, C (1998)

Novel constrained CCK-B dipeptoid antagonists derived from pipercolic acid

Bioorg. Med. Chem. Lett. 8 (11) 1419-1424

1997 Références trouvées : 3

David, C., Bischoff, L., **Meudal, H.**, Llorens-Cortes., C., Roques, BP., Fournie-Zaluski, MC (1997)

Characterization and inhibition of aminopeptidase A by a-mercapto-β-amino acyl dipeptides

Lett. Pept. Sci. 4 (4-6) 411-414

Bellier, B., McCortTranchepain, I., Ducos, B., Danascimento, S., **Meudal, H.**, Noble, F., Garbay, C., Roques, BP (1997)

**Synthesis and biological properties of new constrained CCK-B antagonists :
Discrimination of two affinity states of the CCK-B receptor on transfected CHO
cells**

J. Med. Chem. 40 (24) 3947-3956

Bischoff, L., David, C., Martin, L., **Meudal, H.**, Roques, BP., FournieZaluski, MC (1997)
**2,4-Dinitrophenyl 4-methoxybenzyl disulfide : A new efficient reagent for the
electrophilic sulfenylation of β -amino ester enolates**

J. Org. Chem. 62 (14) 4848-4850

1996 Références trouvées : 3

Fournie Zaluski, MC., Coric, P., They, V., Gonzalez, W., **Meudal, H.**, Turcaud, S., Michel, JB., Roques, BP (1996)

**Design of orally active dual inhibitors of neutral endopeptidase and angiotensin-
converting enzyme with long duration of action**

J. Med. Chem. 39 (13) 2594-2608

Coric, P., Turcaud, S., **Meudal, H.**, Roques, BP., Fournie Zaluski, MC (1996)

**Optimal recognition of neutral endopeptidase and angiotensin-converting
enzyme active sites by mercaptoacyldipeptides as a means to design potent dual
inhibitors**

J. Med. Chem. 39 (6) 1210-1219

Liu, WQ., Carreaux, F., **Meudal, H.**, Roques, BP., Garbay Jaureguiberry, C (1996)

**Synthesis of constrained 4-(phosphonomethyl)phenylalanine derivatives as
hydrolytically stable analogs of O-phosphotyrosine**

Tetrahedron 52 (12) 4411-4422

1993 Références trouvées : 1

Guis, C., Bruetschy, L., **Meudal, H.**, Roques, BP., Gacel, GA (1993)

**Investigation of the structural parameters involved in the delta-opioid
selectivity of several families of families opioid-peptides**

Int. J. Pept. Prot. Res. 41 (6) 576-586
