

Hervé MEUDAL
IEHC CNRS
Responsable Technique Plateforme MO2VING-RMN
Centre de Biophysique Moléculaire
Rue Charles Sadron CS 80054
45071 ORLEANS cedex 02
02-38-25-55-46 / 55-63
herve.meudal@cnrs-orleans.fr



Liste des publications:

Angewandte Chemie International Edition

Photo-isomerization of azobenzene-extended charybdotoxin for the optical control of Kv1.2 potassium channel activity

Achouba, Y., Peres, B., Ascoët, S., **Meudal, H.**, Caumes, C., Zoukimian, C., Millet, H., Choteau-Bodor, M., Carvalhosa, C., Croyal, M., Bouchama, F., Wulff, H., Téletchéa, S., Béroud, R., Ishow, E., Landon, C., Boumendjel, A., Montnach, J., De Waard, M.
Angew Chem Int Ed Engl. (2025) DOI: 10.1002ange.202423278

Dalton Transactions

Investigation of Ln³⁺ complexation by a DOTA derivative substituted by an imidazothiadiazole: synthesis, solution structure, luminescence and relaxation properties

Caillet, E., Numes, L., Eliseeva, S., Ndiaye, M., Isaac, M., Pallier, A., Morfin, J.F., **Meudal, H.**, Petoud, S., Routier, S., Platas-Iglesias, C., Buron, F., Bonnet, C.
Dalton Transactions (2024) DOI: 10.1039/D4DT00533C

Circulation Research

Optical control of cardiac rhythm by in vivo photoactivation of an ERG channel peptide inhibitor

Montnach, J., Millet, H., Persello, A., **Meudal, H.**, De Waard, S., Mesrica, P., Ribeiro, B., Richard, J., Hivonnait, A., Tessier, A., Lauzier, B., Charpentier, F., Mangoni, M.E., Landon, C., Jopling, C., De Waard, M.
Circulation Research (2023) DOI: 10.1161/CIRCRESAHA.123.322880

biomedicine AND PHARMACOTHERAPY

Structure-function relationship of new peptides activating human Na_v1.1

Lopez, L., De Waard, S., **Meudal, H.**, Caumes, C., Khakh, K., Peigneur, S., Oliveira-Mendes, B., lin, S., De Waele, J., Montnach, J., Cestèle, S., Tessier, A., Johnson, J.P., Mantegazza, M., Tytgat, J., Cohen, C., Béroud, R., Bosmans, F., Landon, C., De Waard, M.
Biomedicine & Pharmacotherapy (2023) DOI: 10.1016/j.biopha.2023.115173

marine drugs

Functional diversification of Oyster Big Defensins generates antimicrobial specificity and synergy against members of the Microbiota

De San Nicolas, N., Asokan, A., Diego Rosa, R., Voisin, S.N., Travers, M.A., Rocha, G., Dantan, L., Dorant, Y., Mitta, G., Petton, B., Charrière, G.M., Escoubas, J.M., Boulo, V., Pouzadoux, J., **Meudal, H.**, Loth, K., Aucagne, V., Delmas, A.F., Bulet, F., Montagnani, C., Destoumieux-Garzón, D.
Marine Drugs (2022) DOI: 10.3390/md20120745

ASIAN JOURNAL OF ORGANIC CHEMISTRY

Synthesis of novel 3',3'-cyclic dinucleotide analogues targeting STING protein

Magand, J., Roy, V., **Meudal, H.**, Rose, S., Quesniaux, V., Chalupska, D., Agrofoglio, L.A.
Asian Journal of Organic Chemistry (2022) DOI: 10.1002/ajoc.202200597

Deltex E3 ligases ubiquitylate ADP-ribosyl modification on protein substrates
 Zhu, K., Suskiewicz, M.J., Hloušek-Kasun, A., **Meudal, H.**, Mikoč, A., Aucagne, V., Ahel, D., Ivan AhelNat, I
 Science Advances. (2022) DOI: 10.1126/sciadv.add4253



In vivo spatiotemporal control of voltage-gated ion channels by using photoactivatable peptidic toxins

Montnach, J., Blömer, LA., Lopez, L., Filipis, L., **Meudal, H.**, Lafoux, A., Nicolas, S., Chu, D., Caumes, C., Béroud, R., Jopling, C., Bosmans, F., Huchet, C., Landon, C., Canepari, M., De Waard, M
 Nat Commun. (2022) DOI: 10.1038/s41467-022-27974-w

Ir-Catalyzed β -C(sp²)–H Borylation of Enamides - Access to 3,3-Dihalogeno-2-Methoxypiperidines

Gillaizeau, I., Dondasse, I., Nicolas, C., Sukach, V., **Meudal, H.**, Mimoun, L
 European J Org Chem. (2021) DOI: 10.1002/ejoc.202101302



Three-dimensional structures of avian beta-microseminoproteins: insight from the chicken egg-specific beta-microseminoprotein 3 paralog

Coste, F., Moreau, T., Labas, V., Chesse, M., Bregeon, M., **Meudal, H.**, Loth, K., Castaing, B., Guyot, N., Rehault-Godbert, S
 FEBS Open Bio. (2021) DOI: 10.1002/2211-5463.13166



Metabolomic NMR studies at presymptomatic and symptomatic stages of Huntington's disease on a Drosophila model

Bertrand, M., Decoville, M., **Meudal, H.**, Birman, S., Landon, C
 J Proteome Res. (2020) DOI: 10.1021/acs.jproteome.0c00335

A Venomics approach coupled to high-throughput toxin production strategies identifies the first venom-derived melanocortin receptor agonists

Reynaud, S., Ciolek, J., Degueldre, M., Saez, NJ., Sequeira, AF., Duhoo, Y., Brás, JLA., **Meudal, H.**, Cabo Diez, M., Fernández Pedrosa, V., Verdенаud, M., Boeri, J., Pereira Ramos, O., Ducancel, F., Vanden Driessche, M., Fourmy, R., Violette, A., Upert, G., Mourier, G., Beck-Sickinger, AG., Mörl, K., Landon, C., Fontes, CMGA., Miñambres Herráiz, R., Rodríguez de la Vega, RC., Peigneur, S., Tytgat, J., Quinton, L., De Pauw, E., Vincentelli, R., Servent, D., Gilles, N
 J Med Chem. (2020) DOI: 10.1021/acs.jmedchem.0c00485

Structure, function and evolution of Gga-AvBD11, the archetype of a new structural avian-double- β -defensin family

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
 Proc Natl Acad Sci USA. (2020) DOI: 10.1073/pnas.1912941117



The ancestral N-terminal domain of big defensins drives bacteria-triggered assembly into antimicrobial nanonet

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
 mBio. (2019) DOI: 10.1128/mBio.01821-19



Synthesis by native chemical ligation and characterization of the scorpion toxin AmmTx3

Zoukimian, C., **Meudal, H.**, De Waard, S., Ouarez, KA., Nicolas, S., Canepari, M., Béroud, R., Landon, C., De Waard, M., Boturyn, D
 Bioorg Med Chem. (2019) DOI: 10.1016/j.bmc.2018.12.009



The unusual resistance of avian defensin AvBD7 to proteolytic enzymes preserves its antibacterial activity

Bailleul, G., Kravtzoff, 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
 PLOS One. (2016) DOI: 10.1371/journal.pone.0161573



**Chemistry
A European Journal**

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

Oukhatar, F., **Meudal, H.**, Landon, C., Logothetis, NK., Platas-Iglesias, C., Angelovski, G., Toth, E
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JBC JOURNAL OF BIOLOGICAL CHEMISTRY

Three-dimensional NMR structure of Hen Egg Gallin (Chicken Ovodefensin) reveals a new variation of the β -defensin fold

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

J Biol Chem. (2014) DOI: 10.1074/jbc.M113.507046

Initial insights into structure-activity relationships of avian defensins

Derache, C., **Meudal, H.**, Aucagne, V., Mark, K.J., Cadène, M., Delmas, AF., Lalmanach, A.C., Landon, C
J Biol Chem. (2012) DOI: 10.1074/jbc.M111.312108

Journal of Peptide Science

Synthesis and analytical investigation of C-terminally modified peptide aldehydes and ketone: application to oxime ligation

Bure, C., Marceau, P., **Meudal, H.**, & Delmas, A.F
J Pept Sci. (2011) DOI: 10.1002/psc.1429

The FEBS Journal

Refined solution structure and backbone dynamics of the archaeal MC1 protein

Paquet, F., Loth, K., **Meudal, H.**, Culard, F., Genest, D., & Lancelot, G
FEBS.J (2010) DOI: 10.1111/j.1742-4658.2010.07927.x

JBC JOURNAL OF BIOLOGICAL CHEMISTRY

Molecular requirements for the insecticidal activity of the plant peptide PA1b

Da Silva, P., Rahoui, I., Laugier, C., Jouvensal, L., **Meudal, H.**, Chouabe, C., Delmas, A.F. & Gressent, F
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European Journal of Medicinal Chemistry

Synthesis and anti-HIV-1 integrase activity of modified dinucleotides

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Antimicrobial Agents and Chemotherapy

Primary Structure and Antibacterial Activity of Chicken Bone Marrow-Derived beta-Defensins

Derache, C., Labas, V., Aucagne, V., **Meudal, H.**, Landon, C., Delmas, A.F., Magallon, T. & Lalmanach, A.C
Antimicrob Agents Chemother. (2009) DOI: 10.1128/AAC.00301-09

Angewandte Chemie International Edition

Detection of enzymatic activity by PARACEST MRI : a general approach to target a large variety of enzymes

Chauvin, T., Durand, P., Bernier, M., **Meudal, H.**, Doan, BT., Noury, F., Badet, B., Beloeil, JC., Tóth, E
Angew Chem Int Ed Engl. (2008) DOI: 10.1002/anie.200800809

Comptes Rendus CHIMIE

NMR studies of telomeric nucleoprotein complexes involving the Myb-like domain of the human telomeric protein TRF2

Bilbille, Y., Paquet, F., **Meudal, H.**, Giraud-Panis, MJ., Lancelot, G
C R Chimie. (2006) DOI: 10.1016/j.crci.2005.06.016

Biopolymers

Solution structures of stomoxyn and spinigerin, two insect antimicrobial peptides with an α -helical conformation

Landon, C., **Meudal, H.**, Boulanger, N., Bulet, P., Vovelle, F
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Biochemical Journal

Structure of the zinc finger domain encompassing residues 13-51 of the nucleocapsid protein from simian immunodeficiency virus

Morellet, N., **Meudal, H.**, Bouaziz, S., Roques, BP
Biochem J. (2006) DOI: 10.1042/BJ20051203

Development of potent inhibitors of botulinum neurotoxin type B
Anne, C., Turcaud, S., Quancard, J., Teffo, F., **Meudal, H.**, Fournie-Zaluski, MC., Roques, BP
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Thio-derived disulfides as potent inhibitors of botulinum neurotoxin type B : Implications for zinc interaction

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

Inguimbert, N., Coric, P., Poras, H., **Meudal, H.**, Teffot, F., Fournie-Zaluski, MC., Roques, BP
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Replacement of glycine with dicarbonyl and related moieties in analogues of the C-terminal pentapeptide of cholecystokinin : CCK2 agonists displaying a novel binding mode

Bellier, B., Million, ME., DaNascimento, S., **Meudal, H.**, Kellou, S., Maigret, B., Garbay, C
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Phosphinic derivatives as new dual enkephalin-degrading enzyme inhibitors : Synthesis, biological properties, and antinociceptive activities

Chen, HX., Noble, F., Mothe, A., **Meudal, H.**, Coric, P., Danascimento, S., Roques, BP., George, P., Fournie-Zaluski, MC
J Med Chem. (2000) DOI: 10.1021/jm9904831

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

David, C., Bischoff, L., **Meudal, H.**, Mothe, A., De Mota, N., DaNascimento, S., Llorens-Cortes, C., Fournie-Zaluski, MC., Roques, BP
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Nucleomimetic strategy for the inhibition of HIV-1 nucleocapsid protein NCp7 activities

Druillennec, S., **Meudal, H.**, Roques, BP., Fournie-Zaluski, MC
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Metallopeptidase inhibitors of tetanus toxin : A combinatorial approach

Martin, L., Cornille, F., Turcaud, S., **Meudal, H.**, Roques, BP., Fournie-Zaluski, MC
J Med Chem. (1999) DOI: 10.1021/jm981066w

Novel constrained CCK-B dipeptoid antagonists derived from pipecolic acid

Bellier, B., Da Nascimento, S., **Meudal, H.**, Gincel, E., Roques, BP., Garbay, C
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Synthesis and biological properties of new constrained CCK-B antagonists : Discrimination of two affinity states of the CCK-B receptor on transfected CHO cells

Bellier, B., McCortTranchepain, I., Ducos, B., Danascimento, S., **Meudal, H.**, Noble, F., Garbay, C., Roques, BP
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Design of orally active dual inhibitors of neutral endopeptidase and angiotensin-converting enzyme with long duration of action

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

Liu, WQ., Carreaux, F., **Meudal, H.**, Roques, BP., Garbay Jaureguiberry, C
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Investigation of the structural parameters involved in the delta-opioid selectivity of several families of families opioid-peptides

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