

## Detailed scientific publications:

### - EDITION OF BOOKS (published = 11)

#### ***Ion Channels as Marine Drug Targets.***

Special issue of *Marine Drugs*, MDPI Publisher (2018-19).

#### ***Natural toxins/molecules (and derivatives) from animal venoms: from basic research to therapeutic applications.***

Special issue of *Molecules*, MDPI Publisher (2019).

#### ***Natural toxins/molecules (and derivatives) from animal venoms: from basic research to therapeutic applications.***

Special issue of *Molecules*, MDPI Publisher (2018).

#### ***Structure-Activity Relationship of Natural Products.***

Special issue of *Molecules*, MDPI Publisher (2017).

#### ***Ion Channel Neurotoxins.***

Special issue of *Toxins*, MDPI Publisher (2014).

#### ***Animal Venom Toxins and their Therapeutic Applications.***

Special issue of *Inflammation and Allergy – Drug Targets*, Bentham Science Publishers (2011).

#### ***Drug Targets in Viral Infections.***

Special issue of *Infectious Disorders - Drug Targets*, Bentham Science Publishers (2009).

#### ***Animal Venoms.***

Special issue of *Toxins*, MDPI Publisher (2009).

#### ***Handbook of Biologically-Active Peptides.***

Section Editor for book sections on « Venom Peptides », 1<sup>st</sup> and 2<sup>nd</sup> Editions, Elsevier Publishers (2006 & 2012).

#### ***Animal Toxins and Potassium Channels.***

*Perspectives in Drug Discovery and Design*, Vols. 15/16, Kluwer Academic Publishers (1999).

#### ***HIV Infection in CD4<sup>+</sup> cells.***

*Perspectives in Drug Discovery and design*, Vol. 5, Escom Science Publishers (1996).

### - BOOK CHAPTERS (published = 4)

#### ***Animal Toxins in the World of Modern Biotechnologies.***

Sabatier, J.-M., De Waard, M., « *Handbook of Biologically-Active Peptides*», 2<sup>nd</sup> Edition, Kastin A. Ed., Elsevier Publishers (2012).

#### ***Structure-Function Strategies to Improve the Pharmacological Value of Animal Toxins.***

De Waard, M., Sabatier, J.-M., « *Handbook of Biologically-Active Peptides*», Kastin A. Ed., Elsevier Publishers (2006).

#### ***Methodological Approaches to the Study of Ion Channels using Peptide Toxins : Proposed Comprehensive Guidelines.***

De Waard, M., Sabatier, J.-M., Rochat, H., « *Perspectives in Molecular Toxinology* », Ménez A. Ed., John Wiley & Sons Ltd., pp 255-269 (2002).

#### ***Chemical Synthesis and Characterization of Small Proteins: Example of Scorpion Toxins.***

Sabatier, J.-M., *Handbook of Toxinology*, "Animal Toxins: Tools in Cell Biology", Chapman & Hall, Birkhäuser Verlag Basel/Switzerland, pp 196-216 (2000).

- ARTICLES: specialised journals  
(published = 152)

**20-1 Breakthroughs in medicinal chemistry: New targets and mechanisms, new drugs, new hopes-6.**

Vanden Eynde, J.J., Mangoni, A.A., Rautio, J., Leprince, J., Azuma, Y.T., García-Sosa, A.T., Hulme, C., Jampilek, J., Karaman, R., Li, W., Gomes, P.A.C., Hadjipavlou-Litina, D., Capasso, R., Geronikaki, A., Cerchia, L., Sabatier, J.-M., Ragno, R., Tuccinardi, T., Trabocchi, A., Winum, J.Y., Luque, F.J., Prokai-Tatrai, K., Spetea, M., Gütschow, M., Kosalec, I., Guillou, C., Vasconcelos, M.H., Kokotos, G., Rastelli, G., de Sousa, M.E., Manera, C., Gemma, S., Mangani, S., Siciliano, C., Galdiero, S., Liu, H., Scott, P.J.H., de Los Ríos, C., Agrofoglio, L.A., Collina, S., Guedes, R.C., Munoz-Torrero, D., *Molecules*, 25, 119-143 (2020).

**19-1 Breakthroughs in medicinal chemistry: New targets and mechanisms, new drugs, new hopes-4.**

Mangoni, AA., Guillou, C., Vanden Eynde, J.J., Hulme, C., Jampilek, J., Li, W., Prokai-Tatrai, K., Rautio, J., Collina, S., Tuccinardi, T., de Sousa, M.E., Sabatier, J.-M., Galdiero, S., Karaman, R., Kokotos, G., Torri, G., Luque, F.J., Vasconcelos, M.H., Hadjipavlou-Litina, D., Siciliano, C., Gütschow, M., Ragno, R., Gomez, P.A.C., Agrofoglio, L.A., Munoz-Torrero, D., *Molecules*, 24, 130-142 (2019).

**19-2 Review: Therapeutic potential of carbonic anhydrase inhibitors**

Rahman, S., Bibi, S., Javed, T., Alam, F., Ali, A., Qureshi, Z.R., Ali, S., Ullah, M., Asad, M.H.B., Hasan, S.M.F., Sabatier, J.-M., Rizvanov, A.A., *Pak. J. Pharm. Sci.*, 32, 709-720 (2019).

**19-3 Breakthroughs in medicinal chemistry: New targets and mechanisms, new drugs, new hopes-5.**

Mangoni, AA., Vanden Eynde J.J., Jampilek, J., Hadjipavlou-Litina, D., Liu, H., Reynisson, J., de Sousa, M.E., Gomez, P.A.C., Prokai-Tatrai, K., Tuccinardi, T., Sabatier, J.-M., Luque, F.J., Rautio, J., Karaman, R., Vasconcelos, M.H., Gemma, S., Galdiero, S., Hulme, C., Collina, S., Gütschow, M., Kokotos, G., Siciliano, C., Capasso, R., Agrofoglio, L.A., Ragno, R., Munoz-Torrero, D., *Molecules*, 24, 2415-2428 (2019).

**19-4 Proteomics study of southern Punjab Pakistani cobra (*Naja naja*: formerly *Naja naja karachiensis*) venom.**

Bin Asad, M.H., McCleary, R., Salafutdinov, I., Alam, F., Shah, H.S., Bibi, S., Ali, A., Khalid, S., Hasan, S.M.F., Sabatier, J.-M., De Waard, M., Hussian, I., Rizvanov, A.A., *Toxicology & Environmental Chemistry*, 101, 91-116 (2019).

**19-5 Venoms of Iranian Scorpions (*Arachnida, Scorpiones*) and their potential for drug discovery.**

Kazemi, S.M., Sabatier, J.-M., *Molecules*, 24, 2670-2690 (2019).

**19-6 Bee venom: overview of main compounds and bioactivities for therapeutic interests.**

Wehbe, R., Frangieh, J., El Obeid, D., Sabatier, J.-M., Fajloun, Z., *Molecules*, 24, 2997-3009 (2019).

**19-7 A nanobody-derived mimotope against VEGF inhibits cancer angiogenesis.**

Karami, E., Sabatier, J.-M., Irani, S., Behdani, M., Shahbazzadeh, D., Kazemi-Lomedasht, F., *Artificial Cells, Nanomedicine, and Biotechnology*, in press (2019).

**19-8 Development of a human scFv antibody targeting the lethal Iranian cobra (*Naja oxiana*) snake venom.**

Kazemi-Lomedasht, F., Yamabhai, M., [Sabatier, J.-M.](#), Behdani, M., Reza, M., Shahbazzadeh, D., *Toxicon*, 171, 78-85 (2019).

**18-1 Action mechanism of melittin-derived antimicrobial peptides, MDP1 and MDP2, de novo designed against multidrug-resistant bacteria.**

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**18-2 Lethal toxic dose (i.p. LD50), total protein contents and comparative hemolytic potential of (99mTc-labeled & non-labeled) *Naja naja karachiensis* venom.**

Bin Asad, M.H., Asad, A.F., Bibi, S., Ullah, K., Javed, T., Ullah, M., Ali, A., Qureshi, Z.R., Amirzada, M.I., Al-Kahraman, Y.M., Hasan, S.M.F., [Sabatier, J.-M.](#), Rizvanov, A. *Pak. J. Pharm. Sci.*, 31, 685-689 (2018).

**18-3 Membrane-active peptide PV3 efficiently eradicates multidrug-resistant *Pseudomonas aeruginosa* in a mouse model of burn infection.**

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**18-4 Vipers of the middle east: a rich source of bioactive molecules.**

Rima, M., Alavi-Naini, S.M., Karam, M., Sadek, R., [Sabatier, J.-M.](#), Fajloun, Z. *Molecules*, 23, 2721-2737 (2018).

**18-5 Breakthroughs in medicinal chemistry: New targets and mechanisms, new drugs, new hopes-3.**

Mangoni, AA., Tuccinardi, T., Collina, S., Vanden Eynde, J.J., Munoz-Torrero, D., Karaman, R., Siciliano, C., de Sousa, M.E., Prokai-Tatrai, K., Rautio, J., Guillou, C., Gütschow, M., Galdiero, S., Liu, H., Agrofoglio, L.A., [Sabatier, J.-M.](#), Hulme, C., Kokotos, G., You, Q., Gomez, P.A.C. *Molecules*, 23, 1596-1606 (2018).

**17-1 Breakthroughs in medicinal chemistry: New targets and mechanisms, new drugs, new hopes-2.**

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**17-2 Characteristics and lethality of a novel recombinant dermonecrotic venom phospholipase D from *Hemiscorpius lepturus*.**

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**17-3 Treating autoimmune disorders with venom-derived peptides.**

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**17-4 Preface.**

[Sabatier, J.-M.](#) *Infect. Disorders Drug Targets*, 17(1), 2 (2017).

**17-5 Special issue 'Structure-activity relationship of natural products'.**

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**17-6 Consequences of *Androctonus mauretanicus* and *Buthus occitanus* scorpion venoms on electrolyte levels in rabbits.**

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**17-7 Peptide screen identifies a new NADPH oxidase inhibitor: impact on cell migration and invasion.**

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**16-1 Genetic characterization of lactic acid bacteria from Tunisian milk waste and their antimicrobial activity against some bacteria implicated in nosocomial infections.**

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**16-2 Mechanism of action and in vitro activity of short hybrid antimicrobial peptide PV3 against *Pseudomonas aeruginosa*.**

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**16-3 Editorial.**

Sabatier, J.-M. *Infect. Disorders Drug Targets*, 16(2), 78 (2016).

**15-1 Comparison of the neurotoxic and myotoxic effects of two Moroccan scorpion venoms and their neutralization by experimental polyclonal antivenom.**

Oukkache, N., Ahmad Rusmili, M.R., Othman, I., Ghalim, N., Chgoury, F., Boussadda, L., El Mdaghri, N., Sabatier, J.-M., *Life Sciences*, 124, 1-7 (2015).

**15-2 Correspondences between the binding characteristics of a non-natural peptide, *Lei-Dab7*, and the distribution of SK subunits in the rat central nervous system.**

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**15-3 Endogenous animal toxin-like human beta-defensin 2 inhibits own K<sup>+</sup> channels through interaction with channel extracellular pore region.**

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**15-4 Chlorotoxin: a helpful natural scorpion peptide to diagnose glioma and fight tumor invasion.**

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**15-5 Small-conductance Ca<sup>2+</sup>-activated potassium type 2 channels regulate the formation of contextual fear memory.**

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**15-6 Characterization of Am IT, an anti-insect  $\beta$ -toxin isolated from the venom of scorpion *Androctonus mauretanicus*.**

Oukkache, N., El Jaoudi, R., Chgoury, F., Rocha, M.T., Sabatier, J.-M., *Acta Physiologica Sinica*, 67, 295-304 (2015).

**15-7 Bacteriocins active against multi-resistant Gram-negative bacteria implicated in nosocomial infections.**

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**14-1 Unusual binding mode of scorpion toxin BmKTX onto potassium channels relies on its distribution of acidic residues.**

Chen, Z., Hu, Y., Hu, J., Yang, W., Sabatier, J.-M., De Waard, M., Cao, Z., Li, X., Han, S., Wu, Y., *Biochem. Biophys. Res. Comm.*, 447, 70-76 (2014).

**14-2 Protein content analysis and antimicrobial activity of the crude venom of *Montivipera Bornmuelleri*, a viper from Lebanon.**

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**14-3 Evaluation of the lethal potency of scorpion and snake venoms and comparison between intraperitoneal and intravenous injection routes.**

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**13-1 Peptide binding to Ochratoxin A mycotoxin: a new approach in conception of biosensors.**

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**13-4 Two conserved arginine residues from the SK3 channel outer vestibule control selectivity of recognition by scorpion toxins.**

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13-5 **The deciphered genome of *Mesobuthus martensii* uncovers the resistance mysteries of scorpion to its own venom and toxins at the ion channel level.**

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12-1 **Lacticin LC14, a new bacteriocin produced by *Lactococcus lactis* BMG6.14: isolation, purification and partial characterization.**

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12-2 **Small efficient cell-penetrating peptides derived from scorpion toxin maurocalcine.**

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12-3 **Strategies toward structural and functional 'optimization' of animal peptide toxins.**

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11-1 **Animal venoms: from deadly arsenals (toxins) to therapeutic drug candidates.**

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11-2 **Analysis of the interacting surface of maurotoxin with the voltage-gated Shaker B K<sup>+</sup> channel.**

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10-2 **Protein-protein recognition control by modulating electrostatic interactions.**

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10-3 **Therapeutic value of peptides from animal venoms.**

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10-4 **Structure-function relationships of KTS Disintegrins and design of antiangiogenic drugs.**

Kallech-Ziri, O., Luis, J., Fajloun, Z., Sabatier, J.-M., Lehmann, M., El Ayeb, M., Marrakchi, N., Loret, E., *Letters in Drug Design & Discovery*, 7, 36-40 (2010).

**09-1 Small-conductance Ca<sup>2+</sup>-activated potassium type 2 channels regulate the formation of contextual fear memory.**

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**09-2 Drug targets in viral infections.**

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**08-1 Chemical synthesis and 1H-NMR 3D structure determination of AgTx2-MTX chimera, a new potential blocker for Kv1.2 channel, derived from MTX and AgTx2 scorpion toxins.**

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**08-2 Animal toxins acting on voltage-gated potassium channels.**

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**08-3 Effect of Cu<sup>2+</sup> on the oxidative folding of synthetic Maurotoxin in vitro.**

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**08-4 Chemical synthesis and characterization of J46 peptide, an atypical class IIa bacteriocin from *Lactococcus lactis* subsp. *Cremonis* J46 strain.**

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**08-5 Design of a disulfide-less, pharmacologically-inert and chemically-competent analog of Maurocalcine for the efficient transport of impermeant compounds into cells.**

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**06-3 CD26 modulates nociception in mice via its dipeptidyl-peptidase IV activity.**

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**06-4 Transient loss of voltage control of Ca<sup>2+</sup> release in the presence of maurocalcine in skeletal muscle.**

Pouvreau, S., Csernoch, L., Allard, B., Sabatier, J.-M., De Waard, M., Ronjat, M., Jacquemont, V., *Biophys. J.*, 91, 2206-2215 (2006).

**06-5 Resistance of Hepatitis C virus to NS3-4A protease inhibitors: molecular mechanism of drug resistance induced by R155Q, A156T, D168A and D168V mutations.**

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**06-6 Block of neural Kv1.1 potassium channels for neuroinflammatory disease therapy.**

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**05-1 K<sup>+</sup> channel types targeted by synthetic OSK1, a toxin from *Orthochirus scrobiculosus* scorpion venom.**

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**05-2 Contribution of the functional dyad of animal toxins acting on voltage-gated Kv1-type channels.**

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**05-3 Maurocalcine and domain A of the II-III loop of the dihydropyridine receptor Ca<sub>v</sub>1.1 subunit share common binding sites on the skeletal ryanodine receptor.**

Altafaj, X., Cheng, W., Estève, E., Urbani, J., Grunwald, D., Sabatier, J.-M., Coronado, R., De Waard, M., Ronjat, M., *J. Biol. Chem.*, 280, 4013-4016 (2005).

**05-4 Transduction of the scorpion toxin maurocalcine into cells. Evidence that the toxin crosses the plasma membrane.**

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**05-5 Molecular modeling and docking simulations of scorpion toxins and related analogues on human SKCa2 and SKCa3 channels.**

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**05-7 Differential effects of maurocalcine on Ca<sup>2+</sup> release events and depolarisation-induced Ca<sup>2+</sup> release in rat skeletal muscle.**

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