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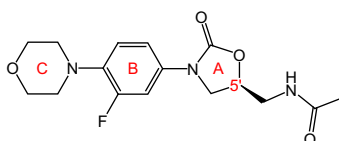
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Recent development of potent analogues of oxazolidinone antibacterial agents

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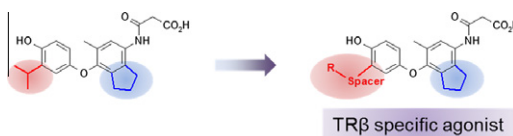


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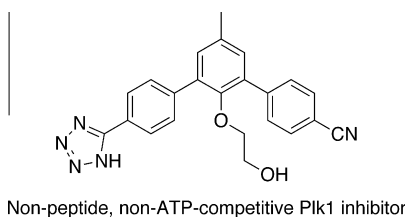
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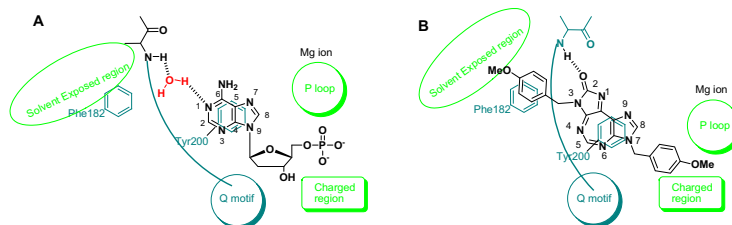
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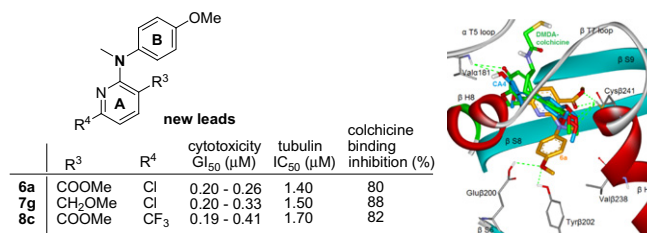


(A) Model for interactions between AMP and DDX3. (B) Proposed model for interactions between **1** and DDX3.

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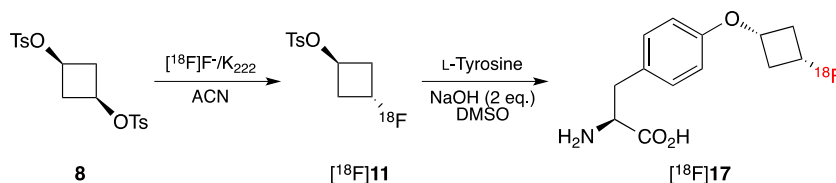
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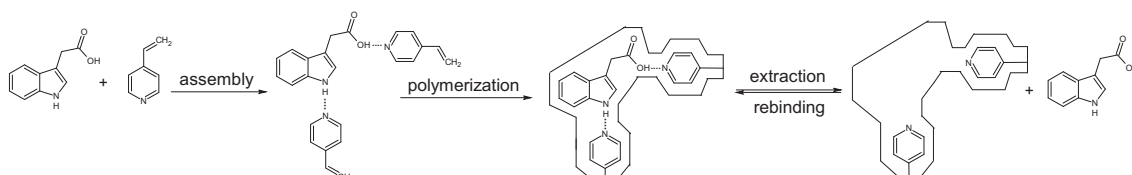
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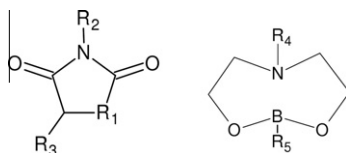
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Ivana Porobić, Darko Kontrec, Milan Šoškić*



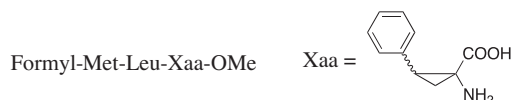
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A formyl peptide substituted with a conformationally constrained phenylalanine residue evokes a selective immune response in human neutrophils pp 668–675

Ryo Hayashi, Masaya Miyazaki, Satoshi Osada, Hiroshi Kawasaki, Ichiro Fujita, Yuhei Hamasaki, Hiroaki Kodama*

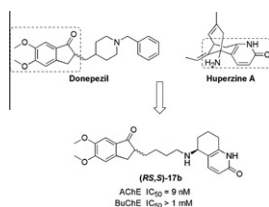


Peptide	Chemotaxis	O ₂ ⁻ production	[Ca ²⁺] _i	Priming
(+)E analog	+	-	+++	-
(-)E analog	+++	-	+++	-
(+)Z analog	+	+	+++	+
(-)Z analog	+++	+++	+++	+



Design, synthesis and evaluation of novel heterodimers of donepezil and huperzine fragments as acetylcholinesterase inhibitors pp 676–683

Yueqing Hu, Jun Zhang, Oormila Chandrashankra, Fanny C. F. Ip, Nancy Y. Ip*

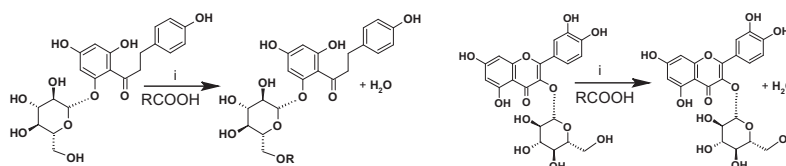


Heterodimers of donepezil and huperzine A fragments tethered with a tetramethylene linker exhibited high potency and selectivity towards acetylcholinesterase inhibition.



Biocatalytic synthesis, structural elucidation, antioxidant capacity and tyrosinase inhibition activity of long chain fatty acid acylated derivatives of phloridzin and isoquercitrin pp 684–692

Ziaullah, Khushwant S. Bhullar, Sumudu N. Warnakulasuriya, H. P. Vasantha Rupasinghe*



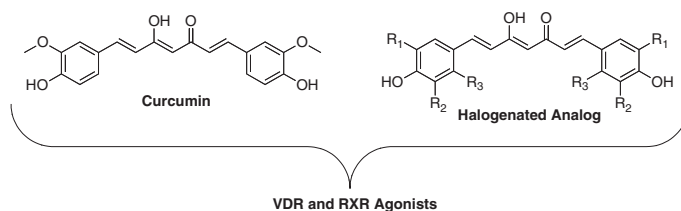
Our present investigation describes the biocatalytic preparation, detailed NMR structural assignment, antioxidant capacity and tyrosinase inhibitory activity of long chain acylated derivatives of phloridzin and isoquercitrin. (i) Acetone, 3 Å molecular sieves, Novozyme 435®, 45 °C, Stirring, 24 h; R = Oleic, Stearic, Linoleic, Linolenic, Eicosapentaenoic (EPA), Docosahexaenoic Acids (DHA) or their corresponding esters.



Synthesis and biological evaluation of halogenated curcumin analogs as potential nuclear receptor selective agonists

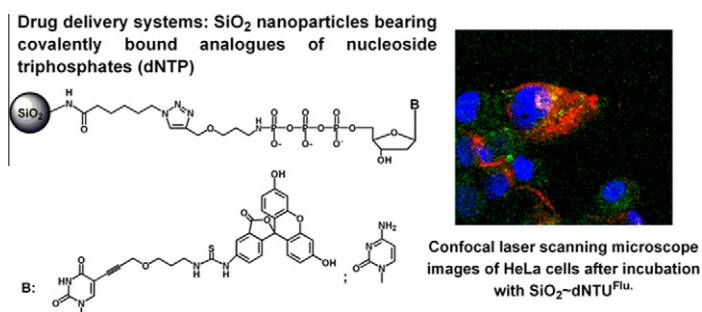
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Shane Batie, Jamie H. Lee, Rabia A. Jama, Drew O. Browder, Luis A. Montano, Chanh C. Huynh, Lisa M. Marcus, Dorian G. Tsosie, Zeynab Mohammed, Vu Trang, Pamela A. Marshall, Peter W. Jurutka*, Carl E. Wagner*

**SiO₂ nanoparticles as platform for delivery of nucleoside triphosphate analogues into cells**

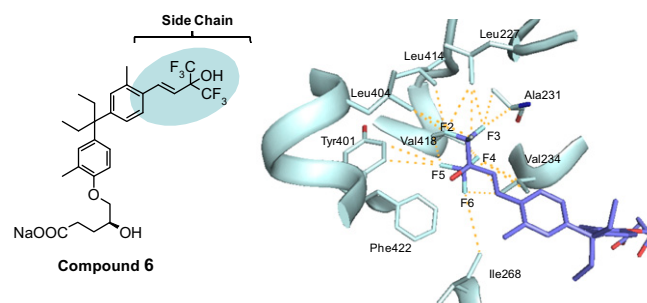
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Svetlana V. Vasilyeva*, Vladimir N. Silnikov, Natalia V. Shatskaya, Asya S. Levina, Marina N. Repkova, Valentina F. Zarytova

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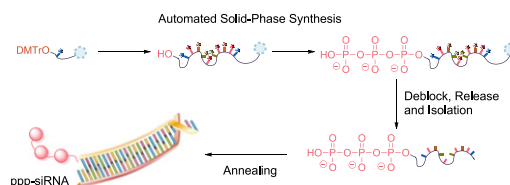
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Hirotaka Kashiwagi*, Masateru Ohta, Yoshiyuki Ono, Kenji Morikami, Susumu Itoh, Hideki Sato, Tadakatsu Takahashi

**Automated parallel synthesis of 5'-triphosphate oligonucleotides and preparation of chemically modified 5'-triphosphate small interfering RNA**

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Ivan Zlatev*, Jeremy G. Lackey, Ligang Zhang, Amy Dell, Kathy McRae, Sarfraz Shaikh, Richard G. Duncan, Kallanthottathil G. Rajeev, Muthiah Manoharan*



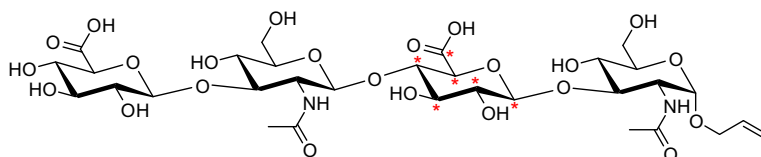
Automated parallel and high-throughput synthesis of 5'-triphosphate oligonucleotides enables the preparation of chemically modified 5'-triphosphate small interfering RNAs designed to address RNA interference gene knockdown and directed immunostimulation.



Synthesis of ^{13}C -labeled and functionalized Hyaluronan derivatives for biophysical studies and surface modifications

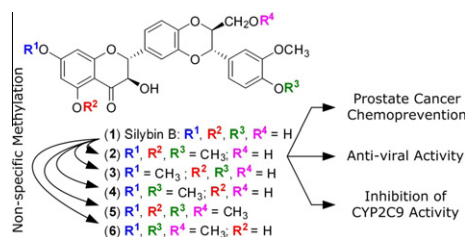
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Stephan Rigol, Liang Xia, Athanassios Giannis*

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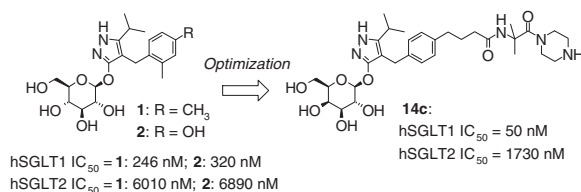
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**Design, synthesis, and structure–activity relationships of a series of 4-benzyl-5-isopropyl-1H-pyrazol-3-yl β -D-glycopyranosides substituted with novel hydrophilic groups as highly potent inhibitors of sodium glucose co-transporter 1 (SGLT1)**

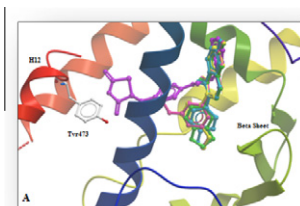
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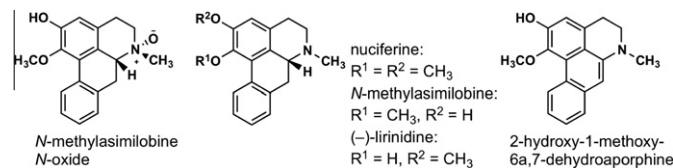
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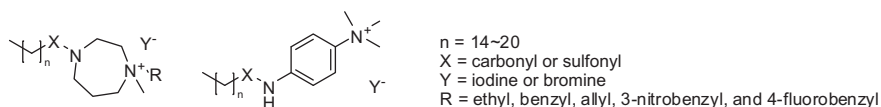
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Design, synthesis and biological evaluation of novel aliphatic amido/sulfonamido-quaternary ammonium salts as antitumor agents

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Doona Song, Jee Sun Yang, Seo Joong Kim, Bo-Kyung Kim, Song-Kyu Park, Misun Won, Kiho Lee, Hwan Mook Kim, Kang-Yell Choi, Kyeong Lee*, Gyoonee Han*



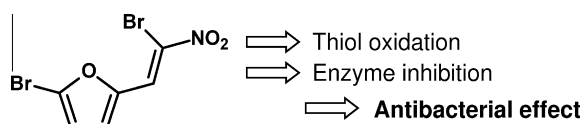
A novel class of aliphatic amido/sulfonamido-quaternary ammonium salts exhibited potent growth inhibitory activities via RhoB-mediated pathway.



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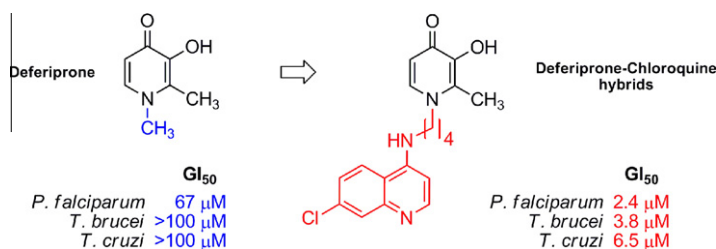
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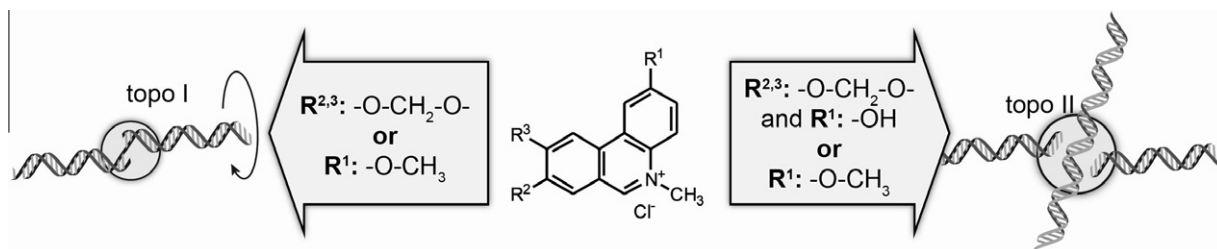
Sebastian S. Gehrke, Erika G. Pinto, Dietmar Steverding, Karin Pleban, Andre G. Tempone, Robert C. Hider, Gerd K. Wagner*



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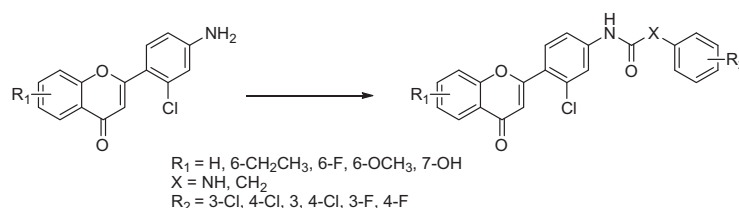
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Simone A. Baechler, Markus Fehr, Michael Habermeyer, Andreas Hofmann, Karl-Heinz Merz, Heinz-Herbert Fiebig, Doris Marko*, Gerhard Eisenbrand

**Exploration of 1-(3-chloro-4-(4-oxo-4H-chromen-2-yl)phenyl)-3-phenylurea derivatives as selective dual inhibitors of Raf1 and JNK1 kinases for anti-tumor treatment**

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Feng Jin, Dan Gao, Cunlong Zhang, Feng Liu, Bizhu Chu, Yan Chen, Yu Zong Chen, Chunyan Tan*, Yuyang Jiang*



A series of 1-(3-chloro-4-(4-oxo-4H-chromen-2-yl)phenyl)-3-phenylurea derivatives were designed, synthesized and explored for their inhibitory activities against kinases and tumor cell lines.

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*Corresponding author

Supplementary data available via SciVerse ScienceDirect

COVER

Botulinum neurotoxins are the most lethal toxins known to man and are considered by the Centers for Disease Control and Prevention (CDC) to be a “Category A” agent placing them as one of the six highest priority bioterrorist agents. There are no approved pharmacological treatments for botulinum neurotoxication. The work detailed combines studies using synthesis, crystallography, modeling, kinetic and cellular research to advance pharmacological intervention against this neurotoxin. [Šilhár, P.; Silvaggi, N.R.; Pellett, S.; Čapková, K.; Johnson, E.A.; Allen, K.N.; Janda, K.D. *Bioorg. Med. Chem.* DOI: 10.1016/j.bmc.2012.12.001]

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