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M. Taguchi, K. Sugimoto, K. Goda, T. Akama, K. Yamamoto, T. Suzuki, Y. Tomishima, M. Nishiguchi, K. Arai, K. Takahashi and T. Kobori	1963	Sphingomyelin Analogues as Inhibitors of Sphingomyelinase
K. Takabe, N. Mase, H. Hashimoto, A. Tsuchiya, T. Ohbayashi and H. Yoda	1967	Chemoenzymatic Synthesis of (<i>E</i>)-3,7-Dimethyl-2-octene-1,8-diol Isolated from the Hairpencils of Male <i>Danaus chrysippus</i> (African Monarch)
D. J. Bennett, A. Anderson, K. Buchanan, A. Byford, A. Cooke, D. K. Gemmell, N. M. Hamilton, M. S. Maidment, P. McPhail, D. F. M. Stevenson, H. Sundaram and P. Vijn	1971	Novel Water Soluble 2,6-Dimethoxyphenyl Ester Derivatives with Intravenous Anaesthetic Activity
S. El Fangour, L. Balas, J.-C. Rossi, A. Fedenyuk, N. Gretskeya, M. Bobrov, V. Bezuglov, C. J. Hillard and T. Durand	1977	Hemisynthesis and Preliminary Evaluation of Novel Endocannabinoid Analogues
S. Jaroch, P. Hölscher, H. Rehwinkel, D. Sülzle, G. Burton, M. Hillmann and F. M. McDonald	1981	Dihydroquinolines with Amine-Containing Side Chains as Potent n-NOS Inhibitors
K. L. Seley, S. Quirk, S. Salim, L. Zhang and A. Hagos	1985	Unexpected Inhibition of <i>S</i> -Adenosyl-L-homocysteine Hydrolase by a Guanosine Nucleoside
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H. Choo, Y. Chong and C. K. Chu	1993	The Role of 2',3'-Unsaturation on the Antiviral Activity of Anti-HIV Nucleosides against 3TC-Resistant Mutant (M184V)
E. Altmann, J. Green and M. Tintelnot-Blomley	1997	Arylaminoethyl Amides as Inhibitors of the Cysteine Protease Cathepsin K—Investigating P ₁ ' Substituents
S. C. Turner, W. A. Carroll, T. K. White, M. Gopalakrishnan, M. J. Coghlan, C.-C. Shieh, X.-F. Zhang, A. S. Parihar, S. A. Buckner, I. Milicic and J. P. Sullivan	2003	The Discovery of a New Class of Large-Conductance Ca ²⁺ -activated K ⁺ Channel Opener Targeted for Overactive Bladder: Synthesis and Structure-Activity Relationships of 2-Amino-4-azaindoles
S. Oh, I. H. Jeong, W.-S. Shin and S. Lee	2009	A Study on the Synthesis of Antiangiogenic (+)-Coronarins A and Congeners from (+)-Sclareolide
J. Howarth and D. Wilson	2013	1,4-Dihydroxy-2,3-dioxatricyclo[8.4.0.0 ^{4,9}]tetradecane and Derivatives with In Vitro Activity Against <i>Plasmodium falciparum</i> , <i>Trypanosoma b. brucei</i> , <i>Trypanosoma cruzi</i> , and <i>Leishmaniasis infantum</i>
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