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The Discovery of Novel Protein Tyrosine Phosphatase Inhibitors Using a High-throughput Screening Approach

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Abstract

Protein tyrosine phosphatase epsilon (PTP ϵ) is important for signal transduction in osteoclasts, and is considered to be an attractive drug target for the treatment of osteoporosis. We identified 11 potent PTP ϵ inhibitors based on three chemical scaffolds through the high-throughput screening of a chemical library. As these compounds are structurally diverse with high bioavailability, they warrant further investigation in the near future. The discovery of these inhibitors and the relationship between their structure and inhibitory activity toward PTP ϵ is discussed in detail.

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- Bonsu Ku, Hye-Yeoung Yun, Kyung Won Lee, Ho-Chul Shin, Sang-Rae Lee, Chang Hyen Kim, Hwangseo Park, Kyu Yang Yi, Chang Hoon Lee and Seung Jun Kim, Identification of N-(5-(phenoxyethyl)-1,3,4-thiadiazol-2-yl)acetamide derivatives as novel protein tyrosine phosphatase epsilon inhibitors exhibiting anti-osteoclastic activity, *Bioorganic & Medicinal Chemistry*, 10.1016/j.bmc.2018.09.022, **26**, 18, (5204-5211), (2018).

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Date()).getTime()]];function(){var s=document.getElementsByTagName('script')[0],p=document.creat  
eElement('script');p.async='async';p.src='//rum-  
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