# 15th Tetrahedron Symposium 

CHALLENGES IN BIOORGANIC AND ORGANIC MEDICINAL CHEMISTRY 24-27 June 2014 | London, UK

## Programme Book

## [P1.95]

Novel compound from Limonium myrianthum possessing antioxidant activity
A.V. Gadetskaya ${ }^{1}$, G.E. Zhussupova ${ }^{1}$, M.K. Murzakhmetova ${ }^{2}$, S.A. Ross ${ }^{3}$, A.I. Zhussupova* ${ }^{* 1}$
${ }^{1}$ Al-Farabi Kazakh National University, Kazakhstan, ${ }^{2}$ Institute for Human and Animal Physiology MES RK, Kazakhstan, ${ }^{3}$ University of Mississippi, USA

Plants of the Limonium Mill genus (Plumbagenaceae family) are represented by 300 species world-wide and 18 in Kazakhstan. Two species, L. gmelinii and L. myrianthum, are present in stocks exceeding 50 thousand tons in Kazakhstan. The current studies are focused on L. myrianthum. Specimens of L. myrianthum were collected in South Kazakhstan in August 2010. Finely ground, air-dried roots ( 300 g ) were subjected to sequential extraction with $n$-hexane ( 0.3 $\mathrm{L} \times 2 ; 48 \mathrm{~h}$ ), acetone ( $0.3 \mathrm{~L} \times 3 ; 48 \mathrm{~h}$ ) and $\mathrm{MeOH}(0.3 \mathrm{~L} \times 3 ; 48 \mathrm{~h})$ at $30^{\circ} \mathrm{C}$ with constant mixing. Extracts were evaporated to dryness under reduced pressure at $35-37^{\circ} \mathrm{C}$. Acetone extracts of the roots showed the highest antioxidant activities and were combined then subjected to silica gel chromatography with elution successively with $100 \% \mathrm{CH}_{2} \mathrm{Cl}_{2}$, then $\mathrm{CH}_{2} \mathrm{Cl}_{2}$ : MeOH mixtures ( $5 \%, 10 \%, 15 \%, 25 \%, 30 \%, 35 \%$ ) and finally $100 \% \mathrm{MeOH}$ to yield 8 fractions. Active fractions were combined and subjected to Sephadex LH-20 chromatography to yield 33 fractions of decreasing molecular mass. A yellow compound isolated from fractions 12 and 13 was identified as epigallocatechin-2-O-p-phenoxy $\left(\mathrm{C}_{21} \mathrm{H}_{18} \mathrm{O}_{8} ;\right.$ mol wt 398.36). It showed potent antioxidant activity in assays of liver microsomal lipid peroxidation. At $4 \mu \mathrm{~g} / \mathrm{mL}$. It reduced the LPO level to $62 \%$ of control and at $20 \mu \mathrm{~g} / \mathrm{mL}$, it reduced the level of LPO to $12 \%$ of control. The antioxidant properties of epigallocatechin-2-O-p-phenoxy require further investigation. Twodimensional spectra were obtained on a Bruker DRX-500 spectrometer. GC-MS spectra were obtained using an HP 6890 gas chromatograph.

Keywords: novel compound, antioxidant activity

| [P1.83] | Silver/ThioClickFerrophos catalyzed enantioselective 1,4-conjugate addition of glycine imino ester to aryl- and alkylidene phosphonates <br> M. Kimura*, S. Fukuzawa, Chuo University, Japan |
| :---: | :---: |
| [P1.84] | N -heterocyclic carbene mediated transformation of cinnamils to vinylfulvenes C.R. Sinu*, V. Nair, National Institute for Interdisciplinary Science and Technology, India |
| [P1.85] | Reducing agent promotes hydroxylation of fatty acid by lipoxygenase T. Itoh*, T. Saito, K. Yamamoto, Showa Pharmaceutical University, Japan |
| [P1.86] | Design and synthesis of antagonist to prevent the active conformation of vitamin D receptor A. Kato*, N. Yoshimoto, T. Itoh, K. Yamamoto, Showa Pharmaceutical University, Japan |
| [P1.87] | Anti-oxidant and tyrosinase inhibitory compounds from the aerial parts of Aruncus dioicus var. kamtschaticus <br> B.T. Zhao*, S.Y. Jeong, Q.H. Vo, P.H. Nguyen, M.H. Woo, Catholic University of Daegu, Republic of Korea |
| [P1.88] | Design, synthesis and characterization of selenium-containing inhibitors of activated thrombinactivatable fibrinolysis inhibitor (TAFIa) <br> K. Yamamoto*, N. Yoshimoto, T. Itoh, H. Ishii, Showa Pharmaceutical University, Japan |
| [P1.89] | Synthesis of functionalized $\alpha$-trifluoroethyl amine scaffolds via addition of highly functionalized organomagnesium and zinc reagents to N -aryl hemiaminal ethers <br> A. Deutsch*, C. Wagner, A. Hoffmann-Röder, Ludwig Maximilian University of Munich, Germany |
| [P1.90] | Chemistry of antitumor 1,2,3,4-tetrahydroisoquinoline natural productsPreparation of tricyclic model compounds with improved cytotoxicity profiles <br> N. Saito*, K. Nakai, M. Yokoya, Meiji Pharmaceutical University, Japan |
| [P1.91] | Novel approach towards the stereoselective synthesis of protected inositols L.H. Sayer*, G.J. Florence, T.K. Smith, University of St Andrews, UK |
| [P1.92] | New anti-inflammatory caffeoylglycerides from the grains of Sorghum bicolor P.H. Nguyen* ${ }^{1}$, B.T. Zhao ${ }^{1}$, Q.H. Vo ${ }^{1}$, J.H. Lee ${ }^{2}$, Y.H. Kim ${ }^{3}$, M.H. Woo ${ }^{1}$, ${ }^{1}$ Catholic University of Daegu, Republic of Korea, ${ }^{2}$ Kangwon National University, Republic of Korea, ${ }^{3}$ Kyungbuk National University, Republic of Korea |
| [P1.93] | New protein tyrosine phosphatase 1B inhibitors from the aerial parts of Selaginella tamariscina P.H. Nguyen* ${ }^{1}$, B.T. Zhao ${ }^{1}$, Q.H. Vo ${ }^{1}$, J.S. Choi ${ }^{2}$, M.H. Woo ${ }^{1}$, ${ }^{1}$ Catholic University of Daegu, Republic of Korea, ${ }^{2}$ Pukyong National University, Republic of Korea |
| [P1.94] | Tablets on the basis of herbal substance "Limonidin" <br> G.E. Zhussupova ${ }^{1}$, K.B. Murzagulova ${ }^{2}$, A.I. Zhussupova* ${ }^{1}$, A.V. Gadetskaya ${ }^{1}$, Z.A. Abilov ${ }^{1},{ }^{1}$ AI-Farabi Kazakh National University, Kazakhstan, ${ }^{2}$ Romat Pharmaceutical Company, Kazakhstan |
| [P1.95] | Novel compound from Limonium myrianthum possessing antioxidant activity <br> A.V. Gadetskaya ${ }^{1}$, G.E. Zhussupova ${ }^{1}$, M.K. Murzakhmetova ${ }^{2}$, S.A. Ross ${ }^{3}$, A.I. Zhussupova ${ }^{*}{ }^{1}$, ${ }^{1}$ Al-Farabi Kazakh National University, Kazakhstan, ${ }^{2}$ Institute for Human and Animal Physiology MES RK, Kazakhstan, ${ }^{3}$ University of Mississippi, USA |
| [P1.96] | Lanostane triterpenoids from Ganoderma lucidum and their biological activities V.T. Nguyen* ${ }^{1}$, J.H. Lee ${ }^{1}$, B.S. Min ${ }^{1}$, ${ }^{1}$ Catholic University of Daegu, Republic of Korea, ${ }^{2}$ Kangwon National University, Republic of Korea |
| [P1.97] | Synthesis and biological evaluation of novel 23-hydroxybetulonic acid derivatives as potential antitumor agents <br> H-Y. Zhang*, P-Q. Zhu, S-T. Xu, X-M. Wu, J-Y. Xu, China Pharmaceutical University, China |
|  | Poster Session II 25 June 2014: 17:00-17:30 26 June 2014: 10:40-11:20 \& 14:00-14:50 |
| [P2.01] | Methoxymethyl ether (MOM) as protecting hydroxyl groups of benzyl ß-L-arabinopyranoside C. Ferris*, M.V. de Paz, B. Begines, E. Galbis, J.A. Galbis, University of Seville, Spain |
| [P2.02] | Functional N -fused imidazole as building block: a novel efficient entry to biologically relevant versatile heterocyclic-condensed purines <br> V. Chaudhary*, S. Guchhait, National Institute of Pharmaceutical Education and Research, India |
| [P2.03] | Development of ERK5 inhibitors for anti-cancer therapy <br> N.C. Martin ${ }^{1}$, ${ }^{1}$ Northern Institute for Cancer research, UK, ${ }^{2}$ Cancer Research Discovery Laboratories, UK, <br> ${ }^{3}$ The Beatson Institute for Cancer Research, UK |
| [P2.05] | Novel asymmetric catalytic intramolecular approach to bioactive chiral cycloalkanols: comparison of batch and flow-techniques <br> H. Viana* ${ }^{1}$, C. Marques ${ }^{1}$, P.H. Seeberger ${ }^{2}$, K. Gilmore ${ }^{2}$, A.J. Burke ${ }^{1}$, ${ }^{1}$ Universidade de Évora, Portugal, ${ }^{2}$ Max Planck Institute of Colloids and Interfaces, Germany |
| [P2.06] | Tandem ATRP-Diels Alder synthesis of polyHEMA-based hydrogels |

