## **Accepted Manuscript**

Design, synthesis and SAR analysis of potent BACE1 inhibitors: Possible lead drug candidates for Alzheimer's disease

Hamadeh Tarazi, Raed Abu Odeh, Raed Al-Qawasmeh, Imad Abu Yousef, Wolfgang Voelter, Taleb H. Al-Tel

PII: S0223-5234(16)30959-X

DOI: 10.1016/j.ejmech.2016.11.021

Reference: EJMECH 9054

To appear in: European Journal of Medicinal Chemistry

Received Date: 6 June 2016

Revised Date: 12 October 2016
Accepted Date: 10 November 2016

Please cite this article as: H. Tarazi, R.A. Odeh, R. Al-Qawasmeh, I.A. Yousef, W. Voelter, T.H. Al-Tel, Design, synthesis and SAR analysis of potent BACE1 inhibitors: Possible lead drug candidates for Alzheimer's disease, *European Journal of Medicinal Chemistry* (2016), doi: 10.1016/j.ejmech.2016.11.021.

This is a PDF file of an unedited manuscript that has been accepted for publication. As a service to our customers we are providing this early version of the manuscript. The manuscript will undergo copyediting, typesetting, and review of the resulting proof before it is published in its final form. Please note that during the production process errors may be discovered which could affect the content, and all legal disclaimers that apply to the journal pertain.



#### ACCEPTED MANUSCRIPT

#### **Graphical Abstract**

Design, Synthesis and SAR Analysis of Potent BACE1 Inhibitors: Possible Lead Drug Candidates for Alzheimer's Disease

Hamadeh Tarazi,<sup>a,b</sup> Raed Abu Odeh,<sup>b,c</sup> Raed Al-Qawasmeh,<sup>d</sup> Imad Abu Yousef,<sup>e</sup> Wolfgang Voelter<sup>f</sup> and Taleb H. Al-Tel,<sup>\*a,b</sup>

<sup>a</sup>College of Pharmacy, University of Sharjah, P.O. Box 27272, Sharjah, UAE

<sup>b</sup>Sharjah Institute for Medical Research, University of Sharjah, P.O.Box 27272, Sharjah, UAE

<sup>c</sup>College of Health Sciences, Department of Medical Laboratory Sciences, University of Sharjah, P.O. Box 27272, Sharjah, United Arab Emirates

<sup>d</sup>Department of Chemistry, The University of Jordan, Amman 11942, Jordan

<sup>e</sup>College of Arts and Sciences, Department of Biology, Chemistry and Environmental Sciences,

American University of Sharjah, Sharjah, United Arab Emirates

<sup>f</sup>Interfakultäres Institut für Biochemie, Eberhard-Karls-Universität Tübingen, Hoppe-Seyler-Straße 4, D-72076 Tübingen

New potent isophthalic acid derivatives armed with imidazolyl and indolyl groups as  $\beta$ -secretase inhibitors have been synthesized. The most potent compound, **11b**, displayed an EC<sub>50</sub> value of 0.29  $\mu$ M.

### Download English Version:

# https://daneshyari.com/en/article/5159087

Download Persian Version:

https://daneshyari.com/article/5159087

<u>Daneshyari.com</u>