Brian S. J. Blagg, Med chem 2013, 3:4
http://dx.doi.org/10.4172/2161-0444.S1.007

OMICS Group Conferences
Accelerating Scientific Discovery

2nd International Conference on
Medicinal Chemistry & Computer Aided Drug Designing

October 15-17, 2013 Hampton Inn Tropicana, Las Vegas, NV, USA

The Hsp90 C-terminal binding site, instructions for and ramifications of inhibition

Brian S. J. Blagg
The University of Kansas, USA

Hsp90 is a molecular chaperone that is responsible for the conformational maturation of more than 200 known substrates, many of which are associated with signaling pathways that are hijacked by transformed cells. As a result, Hsp90 has evolved into a promising anti-cancer target as multiple signaling nodes can be targeted simultaneously through Hsp90 inhibition. More than 15 small molecules that bind to the Hsp90 N-terminal binding pocket have entered clinical trials for evaluation against a number of human malignancies, but unfortunately, a lack of efficacy and/or toxicity has been observed for many of these candidates. In an effort to develop new strategies toward Hsp90 inhibition, we have focused on inhibition of the C-terminal binding site. These methods have resulted in molecules that can segregate induction of the pro-survival heat shock response from client protein inhibition/degradation, and consequently have afforded new methods for the potential treatment of protein misfolding diseases and cancer, respectively. This presentation will provide an overview of the Hsp90 C-terminal binding site and the development of inhibitors that can treat cancer or neurodegenerative diseases.

Biography

Brian S. J. Blagg completed his Ph.D from the University of Utah in 1999,followed by a postdoctoral position at The Scripps Research Institute. He is currently a Professor of medicinal chemistry at The University of Kansas and serves on several Editorial Advisory Boards for Medicinal Chemistry Journals and is an Associate Editor for the Journal of Medicinal Chemistry. The Blagg research Group has won several awards, including the 2009 American Chemical Society's David W. Robertson Award for excellence in Medicinal Chemistry.

bblagg@ku.edu