

Synthesis and biological activity of new benzothiazole derivatives with potent anticancer activity

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The MAPK pathway is normally involved in the regulation of cell proliferation and survival, while oncogenic mutations of its components are frequently observed in several types of cancer. BRAF mutations are present in 5-10% of colorectal cancer patients while overall in 8% of human tumors. Due to the rapid development of resistance to the currently available drugs and the various side effects that come along with therapy, the interest in this field has focused in the discovery of new selective BRAF inhibitors that overcome the problems mentioned above. In order to contribute in this field, a series of benzothiazole derivatives were designed and synthesized as potent BRAF^{V600E} inhibitors. In this presentation, the synthesis of new compounds that contain the benzothiazole ring will be presented along with their biological evaluation which includes in vitro and in vivo experiments.

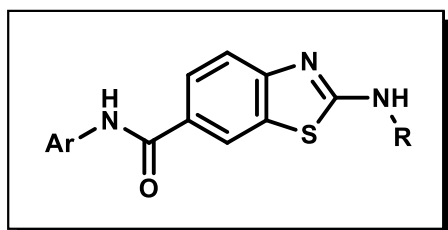


Figure 1: General structure of novel benzothiazole derivatives.

References:

1. B. Agianian, E. Gavathiotis, J. Med. Chem., 61 (2018) 5775-5793

Acknowledgements:

This research is co-financed by Greece and the European Union (European Social Fund- ESF) through the Operational Programme «Human Resources Development, Education and Lifelong Learning» in the context of the project “Strengthening Human Resources Research Potential via Doctorate Research” (MIS-5000432), implemented by the State Scholarships Foundation (IKY).

Also, we acknowledge support of this work by the project “STHENOS-b: Targeted therapeutic approaches against degenerative diseases with special focus on cancer and ageing-optimisation of the targeted bioactive molecules” (MIS 5002398).