## Accepted Manuscript

Synthesis, characterization and anticancer activity of novel ferrocene containing quinolinones: 1-Allyl-2-ferrocenyl-2,3-dihydroquinolin-4(1*H*)-ones and 1-allyl-2-ferrocenylquinolin-4(1*H*)-ones

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PII: S0022-328X(18)30696-X

DOI: 10.1016/j.jorganchem.2018.08.004

Reference: JOM 20531

To appear in: Journal of Organometallic Chemistry

Received Date: 20 April 2018

Revised Date: 2 August 2018

Accepted Date: 7 August 2018

Please cite this article as: A. Pejović, Jó. Drabowicz, M. Cieslak, J. Kazmierczak-Baranska, K. Królewska-Golińska, Synthesis, characterization and anticancer activity of novel ferrocene containing quinolinones: 1-Allyl-2-ferrocenyl-2,3-dihydroquinolin-4(1*H*)-ones and 1-allyl-2-ferrocenylquinolin-4(1*H*)-ones, *Journal of Organometallic Chemistry* (2018), doi: 10.1016/j.jorganchem.2018.08.004.

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## Synthesis, characterization and anticancer activity of novel ferrocene containing quinolinones: 1-allyl-2-ferrocenyl-2,3-dihydroquinolin-4(1*H*)-ones and 1-allyl-2-ferrocenylquinolin-4(1*H*)-ones

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Abstract: A two new series of ferrocene containing quinolinones – 1-allyl-2-ferrocenyl-2,3dihydroquinolin-4(1*H*)-ones and 1-allyl-2-ferrocenylquinolin-4(1*H*)-ones – were prepared and characterized by standard spectroscopic techniques and cyclic voltammetry. The *in vitro* antitumor activity of all synthesized compounds was investigated against Human Cervix Carcinoma (HeLa), Chronic Myelogenous Leukemia (K562) and normal endothelial (HUVEC) cell lines using the MTT method. Quinolone derivative **5c** exhibited the highest cytotoxic activity in the cell growth inhibition of HeLa cell line while **5f** was the most toxic against K562 cells.

**Key words:** Ferrocene, Quinolinones, 2,3-Dihydroquinolin-4(1H)-ones, Quinolin-4(1H)-ones, Electrochemistry, Anticancer activity

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