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**Record 1 of 1****Title:** Synthesis of Chromen-4-One-Oxadiazole Substituted Analogs as Potent -Glucuronidase Inhibitors**Author(s):** Taha, M (Taha, Muhammad); Rahim, F (Rahim, Fazal); Ali, M (Ali, Muhammad); Khan, MN (Khan, Muhammad Naseem); Alqahtani, MA (Alqahtani, Mohammed A.); Bamarouf, YA (Bamarouf, Yasser A.); Gollapalli, M (Gollapalli, Mohammed); Farooq, RK (Farooq, Rai Khalid); Shah, SAA (Shah, Syed Adnan Ali); Ahmed, QU (Ahmed, Qamar Uddin); Zakaria, ZA (Zakaria, Zainul Amiruddin)**Source:** MOLECULES **Volume:** 24 **Issue:** 8 **Article Number:** 1528 **DOI:** 10.3390/molecules24081528 **Published:** APR 2 2019**Times Cited in Web of Science Core Collection:** 0**Total Times Cited:** 0**Usage Count (Last 180 days):** 4**Usage Count (Since 2013):** 4**Abstract:** Chromen-4-one substituted oxadiazole analogs 1-19 have been synthesized, characterized and evaluated for -glucuronidase inhibition. All analogs exhibited a variable degree of -glucuronidase inhibitory activity with IC50 values ranging in between 0.8 +/- 0.1-42.3 +/- 0.8 M when compared with the standard d-saccharic acid 1,4 lactone (IC50 = 48.1 +/- 1.2 M). Structure activity relationship has been established for all compounds. Molecular docking studies were performed to predict the binding interaction of the compounds with the active site of enzyme.**Accession Number:** WOS:000467765700083**PubMed ID:** 31003424**Language:** English**Document Type:** Article**Author Keywords:** chromen-4-one; oxadiazole; synthesis; -glucuronidase inhibition; molecular docking; SAR**KeyWords Plus:** URINARY BETA-GLUCURONIDASE; ALPHA-GLUCOSIDASE INHIBITION; IN-VITRO EVALUATION; MOLECULAR DOCKING; HYBRID MOLECULES; AKT INHIBITORS; DESIGN; DERIVATIVES; BENZOTHAZOLE; ANTIBACTERIAL**Addresses:** [Taha, Muhammad] Imam Abdulrahman Bin Faisal Univ, Dept Clin Pharm, IRMC, POB 1982, Dammam 31441, Saudi Arabia.

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