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Synthesis and Antibacterial Activity of 2,3-dihydro-1H-benzo[f]chromen-1-one and 2,3-dihydro-1H-benzo[f]chromen-1-ol Derivatives

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ABSTRACT

New 2,3-dihydro-1H-benzo[f]chromen-1-one and 2,3-dihydro-1H-benzo[f]chromen-1-ol derivatives (7-14) were prepared from commercially available 2-Hydroxy-acetonaphthone. These compounds were screened for their antibacterial activity against Escherichia coli (MTCC-443), Staphylococcus aureus (MTCC-96), Pseudomonas aeruginosa (MTCC-424) and Streptococcus pyogenes (MTCC-442) bacterial strains by agar well disc diffusion method. It is observed that among the 2,3-dihydro-1H-benzo[f]chromen-1-ol derivatives, compounds 13 and 14 (bearing R_3 = pyrrolidine and morpholine) exhibited excellent activity (zone of inhibition: >20 mm) while the compounds 11 and 12 (bearing R_3 = ethyl and propyl) displayed good activity (zone of inhibition: 16-18 mm).

Keywords: 2,3-dihydro-1H-benzo[f]chromen-1-ol, Aldehydes, Antibacterial activity, Synthesis, *E.coli*, Ampicillin.