ORIGINAL PAPER



Ultrasound-assisted synthesis and in vitro antimicrobial activity of novel 5-oxo-2-pyrrolidinecarboxamides and 7-oxo-2-azepanecarboxamides

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Abstract

A facile one-pot reaction has been set up for the synthesis of novel 5-oxo-2-pyrrolidinecarboxamides and 7-oxo-2-azepanecarboxamides **4a-j** from three-component reaction of keto carboxylic acids **1a-d**, various primary amines **2a-b** and isocyanides **3a-b** under ultrasound irradiation. The present protocol offers attractive characteristics such as easy handling methodology, good-to-excellent yields, environmental friendliness, clean reaction, higher atom economy, convenient operation, and shorter reaction time. The newly synthesized 5-oxo-2-pyrrolidinecarboxamides and 7-oxo-2-azepanecarboxamides **4a-j** have been screened for their antimicrobial activity and the majority of these cyclic amides exhibited weak antimicrobial activity.

 $\textbf{Keywords} \ \ Ultrasound \ synthesis \cdot Antimicrobial \ activity \cdot Ugi \ reaction \ (U-4C-3CR) \cdot 5-Oxo-2-pyrrolidine carboxamides \ and \ 7-oxo-2-azepane carboxamides$