

Unveiling the choline chloride–thiourea (1:1) DES as a greener medium and reagent for pyrimidinethione synthesis from α,β -unsaturated carbonyl compounds

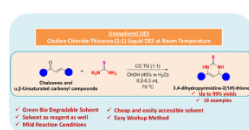


Apeksha Singhal¹, Ankit Yadav^{1,2}, Shashwat Vashistha¹, Khushi Tago¹, T. M. Raoarajan³ and Sharda Pasricha^{1,2}

Author affiliations

Abstract

Widespread studies on developing greener organic chemistry techniques were prompted by growing concerns about the health and environmental issues brought on by the effects of organic compounds. Since solvents and catalysts are necessary for many organic reactions, greener solvents and catalysts could remedy this problem. 3,4-Dihydropyrimidine-2(1H)-thiones (DHPMs) are an essential class of biomolecules. Several methods for synthesizing DHPMs have been reported. Nevertheless, most of these methods were linked to one or more disadvantages, such as laborious catalyst production, use of expensive catalysts, and severe reaction conditions like longer reaction times, higher temperatures and strong base, and reduced yields. Therefore, a simple, effective, and environmentally friendly protocol for the synthesis of 4,6-diaryl-3,4-dihydropyrimidine-2(1H)-thiones from α,β -unsaturated carbonyl compounds using the biodegradable, cheap, and easily accessible choline chloride:thiourea (CC:TU) 1:1 room temperature deep eutectic solvent (DES) is reported. The pyrimidinethione products were obtained from structurally and electronically diverse α,β -unsaturated carbonyl compounds in good to excellent (45–99%) yields under the optimized reaction conditions. The CC:TU (1:1) DES was used for the first time in a synthetic protocol, and it was found to be an augmenting medium and a reactant for the reaction. Hopefully, this paper will open doors to developing new DESs for synthesis. It will also encourage scientists to explore using CC:TU (1:1) as a solvent and reactant for several organic transformations involving thiourea.



* Exclusion of taxes
This article contains 9 pages(s)

Other ways to access this content

Log in
Using your institution credentials



Sign in
With your membership or subscriber account



Supplementary files

Supplementary information
PDF (425/K)

Article information

<https://doi.org/10.1039/D5OB00182J>

Article type Paper
Submitted 31 Jan 2025
Accepted 15 Apr 2025
First published 16 Apr 2025

Citation *Org. Biomol. Chem.*, 2025, **23**, 4951–4959
BibTex Go

Permissions [Request permissions](#)

Social activity Abstracts 0

Tweet

Share