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Please use this identifier to cite or link to this item: http://hdl.handle.net/123456789/3464 Title: Stereoselective synthesis of trans-3-functionalized-4-pyrazolo[5,1-b]thiazole-3-carboxylate substituted β-lactams: Potential synthons for diverse biologically active agents Authors: Mandal, S.K. (/jspui/browse?type=author&value=Mandal%2C+S.K.) Keywords: Cycloaddition Pyrazolo[51-b]thiazole-3-carboxylate Single crystal X-ray Stereoselective Trans-β-Lactams Issue Date: 2020 Taylor and Francis Inc. Publisher: Citation: Synthetic Communications, 50(19), pp.2969-2980. Abstract: An efficient protocol for the stereoselective synthesis of pyrazolo[5,1-b]thiazole-3-carboxylate tethered β-lactam conjugates 8a-j from novel pyrazolo [5,1-b]thiazole-3-carboxylate substituted Schiff's bases 6a-f is reported here. The reaction between various ketene precursors and novel Schiff's bases 6a-f afforded exclusive formation of trans-β-lactams 8a-j. The substrate scope of this approach was investigated extensively by varying different groups (R, Z). All the novel compounds were characterized using various spectroscopic techniques, such as FT-IR, 1H NMR, 13C NMR, elemental analysis, 13C NMR (DEPT-135), and mass spectrometry in representative cases. Single crystal X-ray crystallographic study of trans-ethyl 7-(1-(4-methoxyphenyl)-4-oxo-3phenoxyazetidin-2-yl)-6-methyl-2-(methylthio)pyrazolo[5,1-b]thiazole-3-carboxylate 8a has confirmed the molecular structure and the stereochemical outcome. To the best of our knowledge, the synthesis of such types of Schiff's bases and  $\beta$ -lactam conjugates has not been reported so far. Description: Only IISERM authors are available in the record. URI: https://www.tandfonline.com/doi/full/10.1080/00397911.2020.1788599 (https://www.tandfonline.com/doi/full/10.1080/00397911.2020.1788599) http://hdl.handle.net/123456789/3464 (http://hdl.handle.net/123456789/3464) Appears in Research Articles (/jspui/handle/123456789/9)

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