MedChemComm



CORRECTION

View Article Online
View Journal | View Issue



Cite this: *Med. Chem. Commun.*, 2016, **7**, 2192

Correction: Design and microwave assisted synthesis of novel 2-phenyl/2-phenylethynyl-3-aroyl thiophenes as potent antiproliferative agents

Rupinder Kaur Gill, abc Ramandeep Kaur, Virender Kumar, Vivek Gupta, Gagandeep Singh and Jitender Bariwal (1988)

DOI: 10.1039/c6md90049f

www.rsc.org/medchemcomm

Correction for 'Design and microwave assisted synthesis of novel 2-phenyl/2-phenylethynyl-3-aroyl thiophenes as potent antiproliferative agents' by Rupinder Kaur Gill *et al.*, *Med. Chem. Commun.*, 2016, 7, 1966–1972.

The authors regret the following errors in their paper: (1) The compound numbers shown below the structures in Table 1 should be corrected to show 10d and 12d, instead of 10a and 12a.

(2) On page 1967, section 2.1, first paragraph, when referring to optimization studies shown in Table 1 10a should be replaced by 10d and 12a should be replaced by 12d, as follows:

Earlier, we had reported the efficient formation of biaryl moiety via Suzuki–Miyaura cross-coupling reaction under microwave irradiation conditions; using tetrakis(triphenylphosphane)palladium(0) [Pd(PPh₃)₄] as a catalyst and Cs₂CO₃ as a base under MWI at 140 °C and 100 W for 10 min,²⁹ therefore, we have synthesized our targeted compound (4,5,6,7-tetrahydro-2-phenylbenzo[b]thiophen-3-yl)(phenyl)methanone 12d by coupling of 2-iodo thiophene derivative 10d with phenylboronic acid 11 via Suzuki–Miyaura cross-coupling reaction (Scheme 1) by following the same protocol as employed earlier; however, the yield obtained was very low (Table 1, entry 1). Further, the optimization of this reaction under MW irradiation was carried out using Pd(OAc)₂ and [Pd(PPh₃)₄] as catalysts and Cs₂CO₃ and K₂CO₃ as bases in order to increase the yield of target compounds. It was observed that the use of 5 mol% of Pd(PPh₃)₄ as a catalyst and 3.0 eq. of K₂CO₃ in DMF-H₂O under microwave irradiation (100 W) for 20 min at 140 °C furnished the desired compound 12d in 91% yield (Table 1, entry 3).

The Royal Society of Chemistry apologises for these errors and any consequent inconvenience to authors and readers.

^a Department of Pharmaceutical Chemistry, ISF College of Pharmacy, Moga-142001, Punjab, India

^b I. K. Gujral Punjab Technical University, Kapurthala, Jalandhar-144 601, Punjab, India

^c Department of Pharmaceutical Sciences, Guru Nanak Dev University, Amritsar-143 005, Punjab, India

^d Department of Pharmaceutical Sciences, University of Nebraska Medical Center, Omaha, Nebraska, 68198 USA

^e Post-Graduate Department of Physics & Electronics, University of Jammu, Jammu Tawi-180 006, India

f Bio-Organic and Photochemistry Laboratory, Department of Pharmaceutical Sciences, Guru Nanak Dev University, Amritsar-143 005, Punjab, India

g Satiate Research & Anatech Pvt. Ltd., HSIIDC, Barwala, Panchkula-134118, Haryana, India. E-mail: jitender.bariwal@gmail.com; Fax: +91 1636 239515; Tel: +91 1636 324200