

Short Note

(2E)-3-(4-Dimethylaminophenyl)-1-(2,5-dimethylfuran-3-yl)-prop-2-en-1-one

Abdullah M. Asiri ^{1,2,*} and **Salman A. Khan** ¹

¹ Chemistry Department, Faculty of Science, King Abdul Aziz University, P.O. Box 80203, Jeddah, Saudi Arabia

² The Center of Excellence for Advanced Materials Research, King Abdul Aziz University, Jeddah, P.O. Box 80203, Saudi Arabia

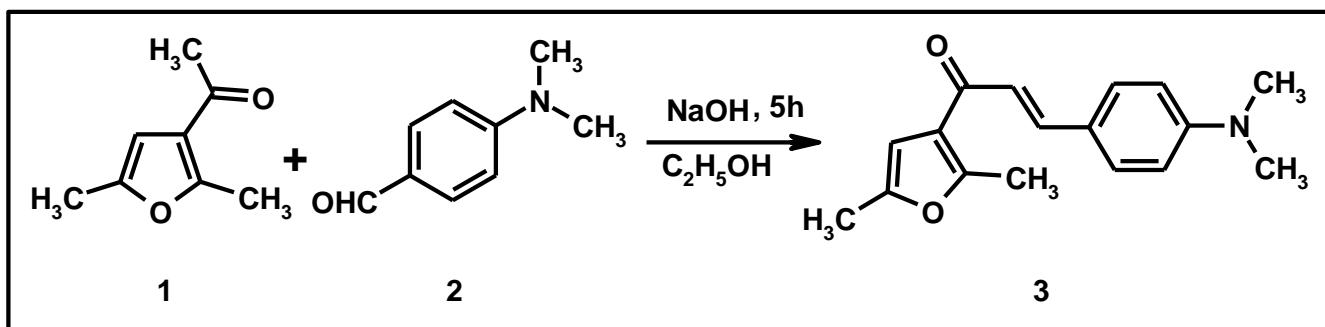
* Author to whom correspondence should be addressed; E-Mail: aasiri2@kau.edu.sa; Tel.: +966 2 6952292; Fax: +966 2 6952292.

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Abstract: The title compound, (2E)-3-(4-dimethylaminophenyl)-1-(2,5-dimethylfuran-3-yl)-prop-2-en-1-one (**3**) was synthesized in high yield by reaction of 3-acetyl-2,5-dimethylfuran and 4-dimethylaminobenzaldehyde in the presence of 30% NaOH solution. The compound was fully characterized from its IR, ¹H NMR, ¹³C NMR, GC-MS data and elemental analysis.

Keywords: chalcone; condensation; 4-dimethylaminobenzaldehyde

Chalcones are characterized by the α,β -unsaturated carbonyl system [1], which is important in elucidating the mechanism of transamination and racemisation reactions in biological systems. Chalcones have been studied as antimalarial [2], antifungal [3], anticancer [4], antioxidant [5], tyrosinase inhibitory [6], antiinflammatory [7] and antibacterial agents [8]. Beyond these very important applications in biological chemistry, chalcones have attracted some attention in the field of material sciences including non-linear optics (NLO) [9], optical limiting [10], electrochemical sensing [11] and Langmuir film [12]. They are also used as intermediates for the formation of various heterocyclic compounds such as pyrimidines, pyrazolines, pyrazoles, thiazines [13]. These observations led us to synthesize a new chalcone from 3-acetyl-2,5-dimethylfuran and 4-dimethylaminobenzaldehyde.

Figure 1. Synthesis of compound (3).

A solution of 3-acetyl-2,5-dimethylfuran (0.46 g, 0.0033 mol) and 4-dimethylaminobenzaldehyde (0.5 g, 0.0033 mol) in an ethanolic solution of NaOH (3.0 g in 10 mL of ethanol) was stirred for 16 h at room temperature. The solution was poured into ice-cold water of pH ~2 (pH adjusted by HCl). The semi-solid separated was collected.

Yellow solid: yield: 86%; semi-solid.

GC-MS m/z (rel. int.%): 270 (58) [M+1]⁺.

IR (KBr) ν_{max} cm⁻¹: 3061 (Ar-H), 2903 (C-H), 1640 (C=O), 1559 (C=C).

¹H NMR (Bruker, 600 MHz, CDCl₃): δ (ppm) 7.63 (d, 2H, J = 8.0 Hz), 6.72 (d, 2H, J = 8.0 Hz), 7.55 (d, 1H, C=CH, J = 15.4 Hz), 7.21 (d, 1H, CH=C, J = 15.4 Hz), 6.74 (s, 1H, Ar-H), 3.17 (s, NCH₃), 3.04 (s, NCH₃), 2.26 (s, CH₃), 2.08 (s, CH₃).

¹³C NMR (150 MHz, CDCl₃): δ (ppm) 184, 156, 152, 150, 142, 130, 124, 122, 123, 119, 112, 113, 105, 40, 39, 14, 13.

Anal. calcd. for C₁₇H₁₉NO₂: C, 75.81, H, 7.11, N, 5.20. Found: C, 75.76, H, 7.09, N, 7.16.

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References

- Asiri, A.M.; Khan, A.S. 2E,2'E-3,3'-(1,4-Phenylene)bis(1-(2,5-dimethylfuran-3-yl)prop-2-en-1-one. *Molbank* **2010**, 2010, M694.
- Bhattacharya, A.; Mishra, L.C.; Sharma, M.; Awasthi, S.M.; Bhasin, V.K. Antimalarial pharmacodynamics of chalcone derivatives in combination with artemisinin against Plasmodium falciparum *in vitro*. *Eur. J. Med. Chem.* **2009**, 44, 3388–3393.
- Lopez, S.N.; Castelli, M.V.; Zaccino, S.A.; Dominguez, J.N.; Lobo, G.; Charris-Charris, J.; Cortes, J.C.G.; Ribas, J.C.; Devia, C.; Rodriguez, A.M.; Ricardo D. Enriz, R.D. *In vitro* antifungal

- evaluation and structure–activity relationships of a new series of chalcone derivatives and synthetic analogues, with inhibitory properties against polymers of the fungal cell wall. *Bioorg. Med. Chem.* **2001**, *9*, 1999–2013.
- 4. Ducki, S. The development of chalcones as promising anticancer agents. *IDrugs* **2007**, *1*, 42–46.
 - 5. Yaylı, N.; Ucuncu, O.; Aydın, E.; Gok, Y.; Yaşar, A.; Baltacı, C.; Yıldırım, N.; Kucuk, M. Stereoselective photochemistry of heteroaryl chalcones in solution and the antioxidant activities. *J. Photochem. Photobiol.* **2005**, *169*, 229–234.
 - 6. Nerya, O.; Musa, R.; Khatib, S.; Tamir, S.; Vaya, J. Chalcones as potent tyrosinase inhibitors: The effect of hydroxyl positions and numbers. *Phytochemistry* **2004**, *65*, 1389–1395.
 - 7. Cheng, J.-H.; Hung, C.-F.; Yang, S.-C.; Wang, J.-P.; Won, S.-J.; Lin, C.-N. Synthesis and cytotoxic, anti-inflammatory, and anti-oxidant activities of 2',5'-diakoxylchalcones as cancer chemopreventive agents. *Bioorg. Med. Chem.* **2008**, *16*, 7270–7276.
 - 8. Avila, H.P.; Smania, E.F.A.; Monache, F.D.; Junior, A.S. Structure–activity relationship of antibacterial chalcones. *Bioorg. Med. Chem.* **2008**, *16*, 9790–9794.
 - 9. Holla, B.S.; Veerendra, B.; Shivananda, M.K. Non-linear optical properties of new arylfuranylpropenones. *J. Crystal Growth* **2004**, *263*, 532–535.
 - 10. Poornesh, P.; Shettigar, S.; Umesh, G.; Manjunatha, K.B.; Kamath, K.P.; Sarojini, B.K.; Narayana, B. Nonlinear optical studies on 1,3-disubstituent chalcones doped polymer films. *Optical Mater.* **2009**, *31*, 854–859.
 - 11. Delavaux-Nicot, B.; Maynadie, J.; Lavabre, D.; Fery-Forgues, S. Ca^{2+} vs. Ba^{2+} electrochemical detection by two disubstituted ferrocenyl chalcone chemosensors. Study of the ligand–metal interactions in CH_3CN . *J. Organomet. Chem.* **2007**, *692*, 874–886.
 - 12. Lu, Z.; Zhang, F.; Lei, X.; Yang, L.; Xu, S.; Duan, X. In situ growth of layered double hydroxide films on anodic aluminum oxide/aluminum and its catalytic feature in aldol condensation of acetone. *Chem. Eng. Sci.* **2008**, *63*, 4055–4062.
 - 13. Asiri, A.M.; Khan, S.A. Synthesis and Anti-bacterial Activities of a Bis-chalcone Derived from Thiophene and Its Bis-cyclized Products. *Molecules* **2011**, *63*, 523–531.