

Water-Mediated Ceric Ammonium Nitrate Catalyzed C-C/C-N Bond Formation: Convenient Access to Polyfunctionalized Pyrazoles *via* Multicomponent Reaction

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Received: 22 January 2019;

Accepted: 25 February 2019;

Published online: 28 March 2019;

AJC-19353

An efficient multicomponent approach has been developed in the environmentally green aqueous medium for synthesis of substituted polyfunctionalized pyrazoles. The simple and readily available aldehydes, malononitrile and phenylhydrazines substrates afforded polysubstituted imidazoles (15 examples) up to 96 % yield. The polyethylene glycol is playing the dual role of solvent and promoter with water in this reaction catalyzed by ceric ammonium nitrate.

Keywords: Multicomponent reaction, Pyrazole, Water-polyethylene glycol media, Ceric ammonium nitrate.

INTRODUCTION

Pyrazole is a privileged heterocyclic skeleton due to its incorporation into pharmaceuticals, agrochemicals and materials as well as its natural occurrence [1,2]. Pyrazoles have been reported to exhibit a wide range of biological properties and immense significance in the pharmacological agents of diverse therapeutic categories [3-5]. Especially, the several blockbuster drugs and pesticides, including celecoxib (celebrex) [6], sildenafil (viagra) [7], zometapine [8], fenpropimox [9], fipronil [10] and tebufenpyrad [11] have made pyrazole a popular synthetic target. Accordingly, they have been extensively studied in the last few decades as a prominent class with great attention for both pharmaceutical and agricultural benefits.

Fascinatingly, ceric ammonium nitrate (CAN) catalyzed organic transformations are rapidly developing due to its well-known versatility and stability, commercial availability, reasonably priced and simple handling attracted much attention in carbon-carbon and carbon-heteroatom bond formation [12-15]. As might be expected of very powerful one-electron oxidants, the chemistry of cerium(IV) oxidation of organic molecules is dominated by radical and radical cation chemistry [16-19] and emerged as a vital feed-stock in diverse chemical industries [20]. As stated in a recent literature, use of CAN in a site-selective fashion might provide a pathway for building addressable libraries of

scaffolds [21-26]. Herein, we are going to report a novel method offering efficient formation of polysubstituted pyrazoles with high productivity from readily available and simple starting materials, namely aldehydes, malononitrile and phenylhydrazines.

EXPERIMENTAL

All commercially available reagents were used directly without purification unless otherwise stated. All the solvents utilized in the reactions were distilled for purity. ¹H NMR and ¹³C NMR spectra were recorded at 400 and 100 MHz, correspondingly. Chemical shifts are expressed in parts per million (ppm) downfield from tetramethylsilane (TMS) as an internal reference standard. Infrared spectra were recorded and band positions are reported. Thin-layer chromatography (TLC) was performed by using commercially prepared 0.25 mm silica gel plates and visualization was effected with a short wavelength UV lamp (254 nm). The relative proportions of solvents in chromatography solvent mixtures are referred volume to volume ratio.

General procedure for the synthesis of 5-amino-pyrazole-4-carbonitriles (4): To a round bottom flask, aldehyde (1) (1 mmol), malononitrile (2) (1.1 mmol), phenylhydrazine (3) (1 mmol), CAN (10 mol %) and PEG-400:H₂O (5 mL) were added with ratio 4:1 (v/v). The resulting reaction mixture was stirred at room temperature for 5 min. After completion of the reaction, stirred reaction mixture further diluted with excess water (2 ×

10 mL) and then filtered to afford the crude product, which was used further by water washing and drying properly without purification to afford the desired product **4**.

5-Amino-1-phenyl-3-phenyl-1H-pyrazole-4-carbonitrile (4a): Pale yellow solid, yield 93 %; m.p.: 159-160 °C; IR (KBr, $\nu_{\max}$, cm^{-1}): 3293, 2363, 1601, 1256; ^1H NMR (400 MHz, CDCl_3): δ = 7.70 (m, 2H, ArH), 7.68 (s, 2H, NH_2), 7.40 (m, 2H, ArH), 7.33 (m, 3H, ArH), 7.29 (m, 2H, ArH), 6.91 (m, 1H, ArH) ppm; ^{13}C NMR (100 MHz, CDCl_3): δ 144.68, 137.33, 135.33, 129.79, 129.33, 129.03, 128.63, 128.44, 128.37, 126.21, 120.13, 112.79 ppm; Anal. calcd. (found) % for $\text{C}_{16}\text{H}_{12}\text{N}_4$: C, 73.83 (73.84); H, 4.65 (4.63); N, 21.52 (21.54).

5-Amino-1-phenyl-3-*p*-tolyl-1H-pyrazole-4-carbonitrile (4b): Pink powder solid, yield 90 %; m.p.: 117-119 °C; IR (KBr, $\nu_{\max}$, cm^{-1}): 3482, 3319, 3095, 2925, 2359, 1598, 1415, 1257, 1127, 1113, 1096; ^1H NMR (400 MHz, CDCl_3): δ 2.41 (s, 3H), 6.91 (dd, J = 3.5 Hz and J = 7.3 Hz, 1H), 7.15 (d, J = 7.6 Hz, 2H), 7.22 (d, J = 7.9 Hz, 2H), 7.29-7.33 (m, 2H), 7.59 (d, J = 7.9 Hz, 2H), 7.70 (s, 2H) ppm; ^{13}C NMR (100 MHz, CDCl_3): δ 153.20, 148.82, 145.20, 138.97, 138.14, 132.95, 129.77, 129.71, 126.62, 120.44, 113.22, 104.65, 21.90 ppm; Anal. calcd. (found) % for $\text{C}_{17}\text{H}_{14}\text{N}_4$: C, 74.43 (74.45); H, 5.14 (5.13); N, 20.42 (20.43).

5-Amino-1-phenyl-3-(4-methoxyphenyl)-1H-pyrazole-4-carbonitrile (4c): Cream coloured solid, yield 91 %; m.p.: 106-108 °C; IR (KBr, $\nu_{\max}$, cm^{-1}): 3313, 2363, 1595, 1244; ^1H NMR (400 MHz, $\text{DMSO}-d_6$): δ = 7.81 (s, 2H, NH_2), 7.57 (d, J = 8.76 Hz, 2H, ArH), 7.19 (t, J = 7.84, 2H, ArH), 7.03 (d, J = 8.00 Hz, 2H, ArH), 6.95 (d, J = 8.76 Hz, 2H, ArH), 6.71 (t, J = 7.22 Hz, 1H, ArH), 3.77 (s, 3H, OCH_3) ppm; ^{13}C NMR (100 MHz, CDCl_3): δ 159.26, 145.53, 136.53, 133.34, 129.01, 128.46, 126.98, 118.28, 114.13, 111.76, 55.13 ppm; Anal. calcd. (found) % for $\text{C}_{17}\text{H}_{14}\text{N}_4\text{O}$: C, 70.33 (70.35); H, 4.86 (4.84); N, 19.30 (19.31); O, 5.51 (5.50).

5-Amino-1-phenyl-3-(4-hydroxyphenyl)-1H-pyrazole-4-carbonitrile (4d): Brown solid, yield 89 %; m.p.: 210-212 °C; IR (KBr, $\nu_{\max}$, cm^{-1}): 3398, 2181, 1688, 1257; ^1H NMR (400 MHz, $\text{DMSO}-d_6$): δ = 9.61 (s, 1H, OH), 7.77 (s, 2H, NH_2), 7.46 (d, J = 8.56 Hz, 2H, ArH), 7.18 (t, J = 7.84 Hz, 2H, ArH), 7.01 (d, J = 7.72, 2H, ArH), 6.78 (d, J = 8.56, 2H, ArH), 6.69 (t, J = 7.24 Hz, 1H, ArH) ppm; ^{13}C NMR (100 MHz, CDCl_3): δ 157.64, 145.65, 137.08, 127.12, 126.86, 118.09, 115.50, 112.04, 111.68 ppm; Anal. calcd. (found) % for $\text{C}_{16}\text{H}_{12}\text{N}_4\text{O}$: C, 69.55 (69.56); H, 4.38 (4.39); N, 20.28 (20.27); O, 5.79 (5.78).

5-Amino-1-(4-chlorophenyl)-3-phenyl-1H-pyrazole-4-carbonitrile (4e): Pink solid, yield 92 %, m.p.: 133-135 °C; IR (KBr, $\nu_{\max}$, cm^{-1}): 3420, 3322, 3095, 2365, 1598, 1487, 1267, 1134, 1096; ^1H NMR (400 MHz, CDCl_3): δ = 7.70 (s, 1H), 7.66 (d, J = 7.6 Hz, 2H), 7.56 (s, 1H), 7.42 (t, J = 7.6 Hz, 2H), 7.33-7.36 (m, 1H), 7.25 (d, J = 8.4 Hz, 2H), 7.06 (d, J = 8.8 Hz, 2H) ppm; ^{13}C NMR (100 MHz, CDCl_3): δ 155.92, 143.67, 138.37, 135.41, 132.28, 129.52, 129.09, 129.02, 126.60, 125.08, 114.22, 112.05 ppm; Anal. calcd. (found) % for $\text{C}_{16}\text{H}_{11}\text{ClN}_4$: C, 65.20 (65.22); H, 3.76 (3.75); Cl, 12.03 (12.02); N, 19.01 (19.03).

5-Amino-1-phenyl-3-(4-bromophenyl)-1H-pyrazole-4-carbonitrile (4f): Pink solid, yield 93 %; m.p.: 165-167 °C;

IR (KBr, $\nu_{\max}$, cm^{-1}): 3302, 2371, 1592, 1254; ^1H NMR (400 MHz, $\text{DMSO}-d_6$): δ = 7.81 (s, 2H, NH_2), 7.57 (q, J = 7.91 Hz, 4H, ArH), 7.21 (t, J = 7.78 Hz, 2H, ArH), 7.06 (d, J = 7.96, 2H, ArH), 6.75 (t, J = 7.27 Hz, 1H, ArH) ppm; ^{13}C NMR (100 MHz, CDCl_3): δ 145.12, 135.12, 134.12, 131.51, 129.09, 127.39, 120.61, 118.95, 112.03 ppm; Anal. calcd. (found) % for $\text{C}_{16}\text{H}_{11}\text{N}_4\text{Br}$: C, 56.66 (56.64); H, 3.27 (3.28); Br, 23.56 (23.55); N, 16.52 (16.54).

5-Amino-3-(4-nitrophenyl)-1-phenyl-1H-pyrazole-4-carbonitrile (4g): Red solid, yield 95 %; m.p.: 164-166 °C; IR (KBr, $\nu_{\max}$, cm^{-1}): 3465, 3355, 3105, 2354, 1610, 1417, 1456, 1345, 1256, 1133, 1108, 1094; ^1H NMR (400 MHz, CDCl_3): δ = 8.25 (d, J = 7.6 Hz, 2H) 8.03 (s, 1H), 7.74-7.77 (m, 3H), 7.20-7.34 (m, 2H), 7.18 (d, J = 7.6 Hz, 2H), 6.96 (s, 1H) ppm; ^{13}C NMR (100 MHz, CDCl_3): δ 156.53, 149.56, 145.26, 137.73, 135.32, 131.72, 130.93, 129.90, 122.78, 122.15, 123.47, 113.43, 112.36 ppm; Anal. calcd. (found) % for $\text{C}_{16}\text{H}_{11}\text{N}_5\text{O}_2$: C, 62.95 (62.96); H, 3.63 (3.61); N, 22.94 (22.96); O, 10.48 (10.47).

5-Amino-3-(2-chlorophenyl)-1-phenyl-1H-pyrazole-4-carbonitrile (4h): Yellow solid, yield 90 %; m.p.: 139-141 °C; IR (KBr, $\nu_{\max}$, cm^{-1}): 3475, 3425, 3150, 2520, 2335, 1660, 1575, 765, 720; ^1H NMR (400 MHz, CDCl_3): δ = 7.61 (s, 2H) 7.58 (d, J = 8.4 Hz, 2H), 7.35 (d, J = 8.4 Hz, 2H), 7.28-7.32 (m, 2H), 7.15 (d, J = 7.6 Hz, 2H), 6.94 (t, J = 7.6 Hz, 1H) ppm; ^{13}C NMR (100 MHz, CDCl_3): δ 154.82, 144.87, 144.43, 139.44, 138.52, 137.90, 129.41, 128.54, 127.04, 119.34, 117.25, 112.84, 114.04, 99.48 ppm; Anal. calcd. (found) % for $\text{C}_{16}\text{H}_{11}\text{N}_4\text{Cl}$: C, 65.20 (65.22); H, 3.76 (3.75); Cl, 12.03 (12.02); N, 19.01 (19.02).

5-Amino-3-(4-cyanophenyl)-1-phenyl-1H-pyrazole-4-carbonitrile (4i): Yellow solid, yield 94 %; m.p. 158-160 °C; IR (KBr, $\nu_{\max}$, cm^{-1}): 3434, 3316, 3275, 2353, 2229, 1583, 1472, 1265, 1155, 1127, 1096; ^1H NMR (400 MHz, CDCl_3): δ 10.72 (s, 1H, ArH), 7.85 (s, 1H, NH), 7.77-7.81 (m, 4H, ArH), 7.24 (t, J = 7.2 Hz, 2H, ArH), 7.13 (d, J = 7.6 Hz, 2H, ArH), 6.82 (t, J = 7.2 Hz, 1H, ArH) ppm; ^{13}C NMR (100 MHz, CDCl_3): δ 152.34, 146.19, 137.83, 135.75, 129.70, 128.15, 127.09, 126.74, 120.56, 117.92, 115.61, 113.25, 112.82 ppm; Anal. calcd. (found) % for $\text{C}_{17}\text{H}_{11}\text{N}_5$: C, 71.57 (71.58); H, 3.89 (3.88); N, 24.55 (24.57).

5-Amino-1-(3-chlorophenyl)-3-phenyl-1H-pyrazole-4-carbonitrile (4j): Pale yellow solid, yield 92 %; m.p.: 172-175 °C; ^1H NMR (400 MHz, CDCl_3): δ = 7.72-7.68 (m, 1H, ArH), 7.64 (s, 2H, NH_2), 7.43-7.35 (m, 3H, ArH), 7.22-7.20 (m, 2H, ArH), 7.18 (s, 1H, ArH), 6.93 (dd, J = 8.4 & 1.6 Hz, 1H, ArH), 6.86 (dd, J = 8.0 & 1.2 Hz, 1H, ArH) ppm; ^{13}C NMR (100 MHz, CDCl_3): δ 145.77, 138.36, 135.20, 134.88, 130.25, 128.81, 128.67, 126.37, 119.96, 112.80, 110.89 ppm; Anal. calcd. (found) % for $\text{C}_{16}\text{H}_{11}\text{N}_4\text{Cl}$: C, 65.20 (65.22); H, 3.76 (3.78); Cl, 12.03 (12.05); N, 19.01 (19.03).

5-amino-3-(5-bromo-2-hydroxyphenyl)-1-(4-bromophenyl)-1H-pyrazole-4-carbonitrile (4k): Yellow solid, yield 87 %; m.p.: 180-182 °C; ^1H NMR (400 MHz, CDCl_3): δ 9.23 (s, 1H, OH, Ar-OH), 7.83 (s, 1H, ArH), 7.23-7.34 (m, 5H), 6.83 (d, 3H, ArH) ppm; ^{13}C NMR (100 MHz, CDCl_3): δ 155.9, 142.71, 139.59, 132.17, 132.10, 131.22, 120.64, 118.31, 113.98, 112.13, 111.0 ppm; Anal. calcd. (found) % for $\text{C}_{16}\text{H}_{10}\text{N}_4\text{OBr}_2$:

C, 44.27 (44.28); H, 2.32 (2.31); Br, 36.81 (36.82); N, 12.91 (12.92); O, 3.69 (3.70).

5-Amino-3-(5-bromo-2-hydroxyphenyl)-1-(4-chlorophenyl)-1H-pyrazole-4-carbonitrile (4l): Yellow solid, yield 88 %; m.p.: 176-178 °C; ¹H NMR (400 MHz, CDCl₃): δ 7.79 (s, 1H, ArH), 7.60 (s, 1H, OH, Ar-OH), 7.27-7.35 (m, 5H), 6.89-6.93 (m, 3H, ArH) ppm; ¹³C NMR (100 MHz, CDCl₃): δ 156.03, 141.61, 139.94, 132.73, 131.47, 129.54, 125.95, 120.03, 118.53, 113.87, 111.18 ppm; Anal. calcd. (found) % for C₁₆H₁₀N₄OBrCl: C, 49.32 (49.32); H, 2.59 (2.61); Br, 20.51 (20.49); Cl, 9.10 (9.12); N, 14.38 (14.40); O, 4.11 (4.13).

5-Amino-1-(3-chlorophenyl)-3-(4-chlorophenyl)-1H-pyrazole-4-carbonitrile (4m): Yellow solid, yield 90 %; m.p.: 97-99 °C; ¹H NMR (400 MHz, CDCl₃): δ 7.61-7.62 (m, 2H, ArH), 7.59 (s, 2H, NH₂), 7.35-7.38 (m, 2H, ArH), 7.17-7.21 (m, 2H, ArH), 6.85-6.93 (m, 2H, ArH) ppm; ¹³C NMR (100 MHz, CDCl₃): δ 145.52, 136.30, 135.23, 134.41, 133.44, 130.29, 128.91, 127.45, 120.19, 112.82, 110.92 ppm; Anal. calcd. (found) % for C₁₆H₁₀N₄Cl₂: C, 58.38 (58.36); H, 3.06 (3.08); Cl, 21.54 (21.56); N, 17.02 (17.04).

5-Amino-3-(2-chloro-4-fluorophenyl)-1-(4-chlorophenyl)-1H-pyrazole-4-carbonitrile (4n): Yellow solid, yield 86 %; m.p.: 134-136 °C; ¹H NMR (400 MHz, CDCl₃): δ 8.10 (s, 2H, NH₂), 7.84 (s, 1H, ArH), 7.20 (m, 2H, ArH), 7.1 (s, 1H, ArH), 7.0 (m, 1H, ArH), 6.90 (m, 2H, ArH) ppm; ¹³C NMR (100 MHz, CDCl₃): δ 163.67, 161.16, 145.37, 135.26, 133.65, 133.29, 133.19, 130.31, 128.74, 120.36, 116.97, 116.73, 114.93, 114.72, 112.86, 110.96 ppm; Anal. calcd. (found) % for C₁₆H₉N₄Cl₂F: C, 55.35 (55.36); H, 2.61 (2.60); Cl, 20.42 (20.43); F, 5.47 (5.46); N, 16.14 (16.15).

5-Amino-3-(2-chloro-6-fluorophenyl)-1-(4-chlorophenyl)-1H-pyrazole-4-carbonitrile (4o): Yellow solid, yield 85 %; m.p.: 75-78 °C; ¹H NMR (400 MHz, CDCl₃): δ 7.70 (m, 2H, ArH), 7.68 (s, 2H, NH₂), 7.33 (m, 2H, ArH), 7.29 (m, 1H, ArH), 6.91 (m, 2H, ArH) ppm; ¹³C NMR (100 MHz, CDCl₃): δ 163.57, 161.07, 142.87, 133.23, 133.21, 133.18,

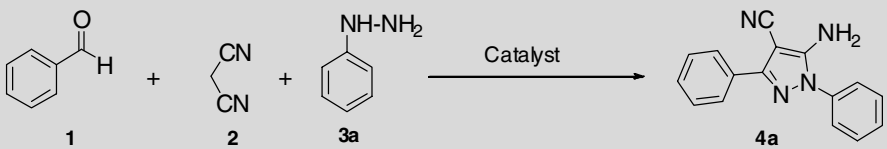
133.08, 129.26, 128.91, 128.87, 125.10, 116.97, 114.89, 114.68, 113.93 ppm; Anal. calcd. (found) % for C₁₆H₉N₄Cl₂F: C, 55.35 (55.37); H, 2.61 (2.59); Cl, 20.42 (20.44); F, 5.47 (5.45); N, 16.14 (16.16).

RESULTS AND DISCUSSION

Our studies directed toward inexpensive, non-toxic cerium(IV) mediated importantly more sustainable reaction media poly(ethylene glycol) (PEG) [27-30] for organic transformations pleasing into account of its popularity. At the outset of our investigation, we selected benzaldehyde (**1**), malononitrile (**2**) and phenylhydrazine (**3**) as model substrates to optimize reaction condition in presence of CAN (20 mol %) using nonvolatile and nontoxic polyethylene glycol (PEG-200) medium. We observed 58 % of the corresponding desired product 5-amino-1-phenyl-3-phenyl-1H-pyrazole-4-carbonitrile (**4a**) after 5 min at room temperature (Table-1, entry 1). To improve the yields of desired product, various types of polyethylene glycol solutions such as PEG-300, PEG-400 and PEG-600 were examined and obtained the corresponding product 5-amino-1-phenyl-3-phenyl-1H-pyrazole-4-carbonitrile (**4a**) (Table-1, entries 2-4) in yields of 59, 65 and 60 %, respectively. Among them, PEG-400 turned out to be the most efficient medium, giving **4a** in higher yield in comparisons to others.

On the other hand, PEG solutions as green reaction media was spurred by several factors *e.g.*, green separation chemistry [31-37], study of PEG metal ion coordination [36,38-41], aqueous biphasic solvent (ABS) solvent properties [42-46]. However, the integration of solvent properties of polyethylene glycol (PEG) by aqueous biphasic reactive extraction (ABRE) concept and its additional phase-transfer characteristics into a single efficient system which facilitates the separation of reactants and catalysts from products. Also, environmentally benign properties of these systems in comparison to the current use of organic solvents in extraction and reactive extraction have increased interest. Under these state of affairs, the effect of

TABLE-1
OPTIMIZATION OF THE REACTION CONDITIONS^a

			
Entry	Catalyst	Solvent	Yield ^b (%)
1	Ceric ammonium nitrate	PEG-200	58
2	Ceric ammonium nitrate	PEG-300	59
3	Ceric ammonium nitrate	PEG-400	65
4	Ceric ammonium nitrate	PEG-600	60
5	Ceric ammonium nitrate	PEG-400:H ₂ O [4:1]	84
6	Ceric ammonium nitrate	PEG-400:H ₂ O [1:1]	66
7	Ceric ammonium nitrate	PEG-400:H ₂ O [1:4]	63
8 ^c	Ceric ammonium nitrate	PEG-400:H ₂ O [4:1]	84
9 ^d	Ceric ammonium nitrate	PEG-400:H ₂ O [4:1]	84
10 ^e	Ceric ammonium nitrate	PEG-400:H ₂ O [4:1]	77
11 ^f	Ceric ammonium nitrate	PEG-400:H ₂ O [4:1]	69
12 ^{d,g}	Ceric ammonium nitrate	PEG-400:H ₂ O [4:1]	—
13 ^{d,g}	Ceric ammonium nitrate	PEG-400:H ₂ O [4:1]	—

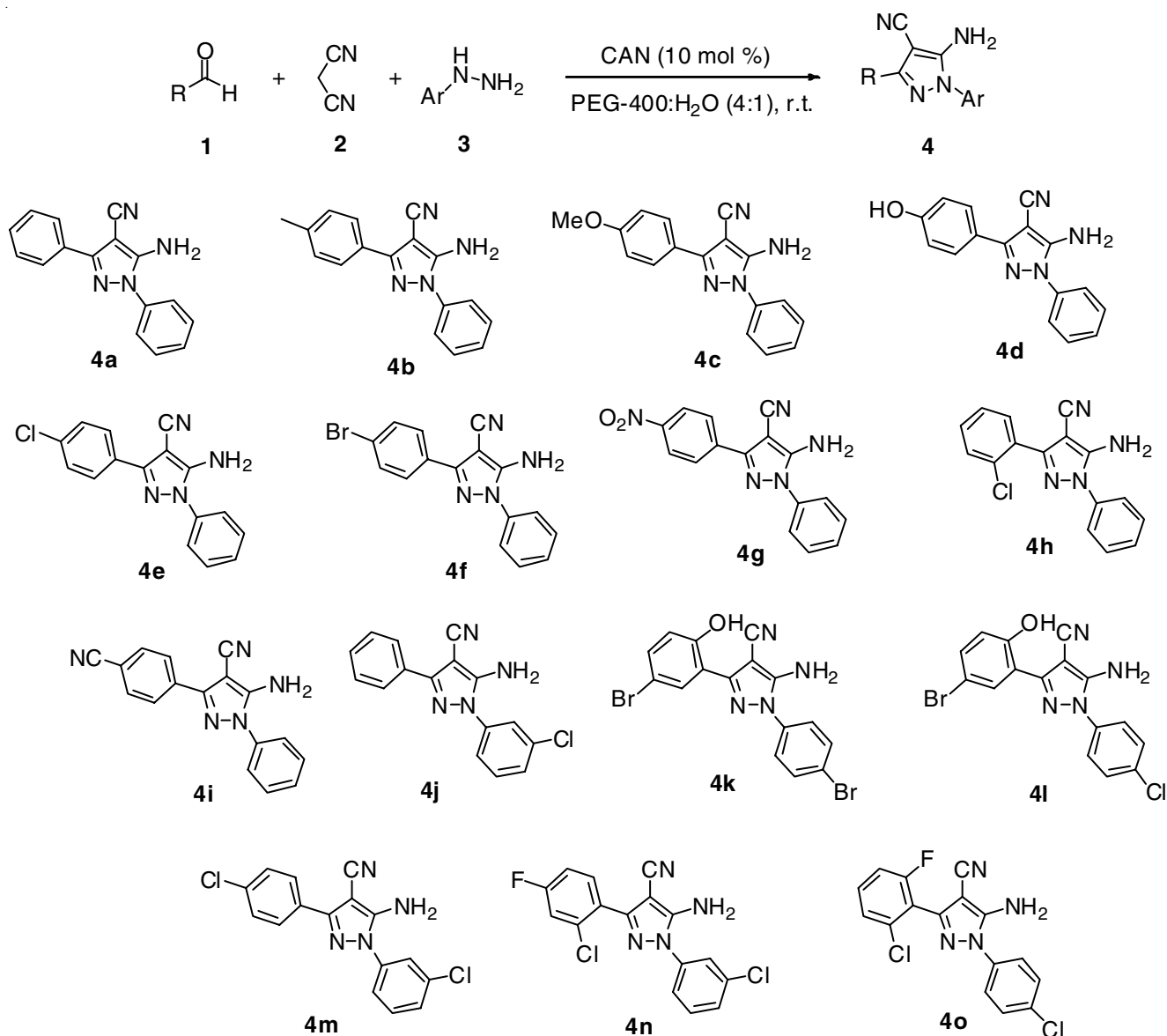
^aReaction conditions: **1a** (0.5 mmol, 1.0 equiv), **2** (0.5 mmol, 1.0 equiv), **3a** (0.5 mmol, 1.0 equiv), CAN (20 mol %) and solvent (5 mL) stirred for 5 min at room temperature. ^bIsolated yields, ^cdef CAN (15, 10, 7 and 5 mol %), ^eTemp. (°C) = 60, 80 in entries 12 and 13, respectively, CAN = ceric ammonium nitrate, PEG = polyethylene glycol.

different PEG-400:H₂O (v/v) ratios at room temperature was studied using CAN (20 mol %). The yields of desired product **4a** dramatically increased, *i.e.* 84, 66 and 63 %, respectively (Table-1, entries 5-7). After that, reaction conditions were optimized by variation of catalyst loading and reproduced same results with 15 and 10 mol % of CAN as well (Table-1, entries 8-9). In next, we find that further decrease in catalyst loading from 10 to 5 mol % desired product was obtained in lower yields (Table-1, entries 10-11). Furthermore, the same results were obtained for different temperatures such as 60, 80 °C (Table-1, entries 12-13), respectively.

With the optimized reaction condition in hand, we investigated the various functional group substituted on the benzene ring of aromatic benzaldehydes (Fig. 1), we found that compounds **4b-4d** have electron-donating functional groups bearing benzaldehydes **1b-1d** (4-Me, 4-MeO and 4-OH) afforded slightly higher yields. Then, considerably decreased in the yields (com-

pounds **4e-4h**) for electron withdrawing functional group was observed. The scope of the reaction was further extended to different aromatic aldehydes with malononitrile and substituted phenylhydrazines under the optimized reaction conditions and the products obtained (compounds **4i-4o**). The electronic effect introduced by the substitution of the aromatic aldehydes had insignificant influence on the yields of the corresponding pyrazole derivatives. The products were obtained as white to dark yellow solid and data matches well with the literature values. In general, the CAN catalyzed progression is taking place well under mild conditions for a wide range of aldehydes and phenylhydrazines moieties with electron-withdrawing and electron-donating substituents for synthesis of polysubstituted pyrazoles in good to excellent yields.

On the basis of the above-mentioned experimental results, a possible reaction mechanism is proposed for the developed multicomponent protocol in presence of CAN as an activator



Reaction conditions: **1** (1.1 mmol), **2** (1.0 mmol), **3** (1.0 mmol), 10 mol % CAN in 5 mL PEG-400:H₂O (4:1) at room temperature

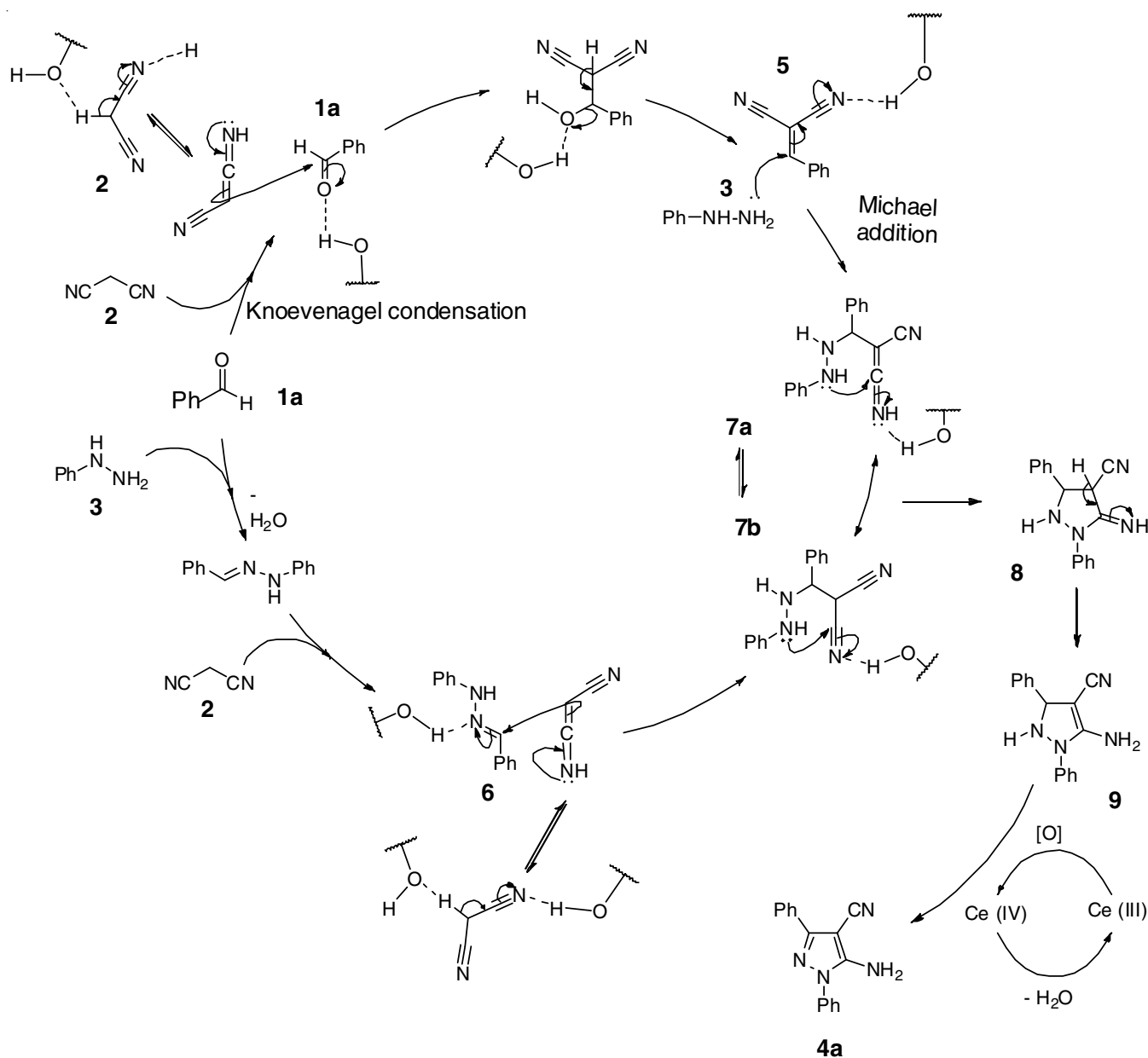
Fig. 1. Synthesis of substituted polyfunctionalized pyrazoles

and oxidizing agent depicted in **Scheme-I** (**4a** as an example). Initially, aromatic benzaldehyde (**1a**) reacted with activated nitrile group from malononitrile (**2**) in presence of water and polyethylene glycol 400 (PEG-400:H₂O) medium with ratio 4:1 (v/v), leads to the formation of the benzylidene-malononitrile adduct (**5**) *via* condensation with the formation of C=C bond. Afterward, phenylhydrazine (**3**), to be able to react easily with formed adduct benzylidene-malononitrile (**5**) gives rise to intermediate (**6a**). On the other hand, aromatic benzaldehyde (**1a**) and phenylhydrazine (**3**) possibly leads to the formation of intermediate imine (**7**), subsequently reacts with malononitrile (**2**) furnishes 2-(phenyl(2-phenylhydrazinyl)-methyl)malononitrile (**6b**) *via* Michael addition. Both of the forms, **6a** and **6b** have equilibrium *via* a tautomeric 1,3-H shift. Such equilibrium responsible to interconvert them into one another ascertained that reaction time and yield of the product are not seemed to be dependent on the order of addition of reactants in reaction. As a result, both (**6a**) and (**6b**) affords

five-membered ring construction of 5-imino-1,3-diphenylpyrazolidine-4-carbonitrile (**9**) *via* C-N bond formation. In next step, owing to the imine-enamine tautomerism, 5-imino-1,3-diphenylpyrazolidine-4-carbonitrile (**9**) get transform to 2,3-dihydropyrazole derivative (**10**) with tautomeric proton shift. Simultaneously, 2,3-dihydropyrazole derivative (**10**) converts to the product (**4a**) efficiently with C-C and C-N bond formation in presence of CAN under aerobic oxidation (**Scheme-I**).

Conclusion

In summary, an efficient and regioselective method has been developed having facile and practical reaction condition to furnish polysubstituted pyrazoles, providing easy access to the broad library. In this, the intramolecular C-N bond formation *via* oxidation of C-H and N-H bonds in construction of nitrogen-containing heterocyclic ring system has various advantages, such as direct functionalization of C-H bonds with NHR groups without pre-activation of the reaction centers;



facilitative preparation of structurally diverse substrates and no generation of the unwanted stoichiometric amounts of byproducts derived from the leaving groups in substrates for the classic coupling reaction. Hence, it is general, inexpensive and environmentally benign approach having operational simplicity (reaction, work-up and purification) could make it potentially attractive for library construction of polyfunctionalized pyrazoles, particularly in the area of pharmaceuticals.

CONFLICT OF INTEREST

The authors declare that there is no conflict of interests regarding the publication of this article.

REFERENCES

- J.-J. Liu, M. Zhao, X. Zhang, X. Zhao and H.-L. Zhu, *Mini Rev. Med. Chem.*, **13**, 1957 (2013); <https://doi.org/10.2174/13895575113139990078>.
- V. Kumar, K. Kaur, G.K. Gupta and A.K. Sharma, *Eur. J. Med. Chem.*, **69**, 735 (2013); <https://doi.org/10.1016/j.ejmech.2013.08.053>.
- J. Dwivedi, S. Sharma, S. Jain and A. Singh, *Mini Rev. Med. Chem.*, **18**, 918 (2018); <https://doi.org/10.2174/1389557517666170927160919>.
- S. Ganguly and S.K. Jacob, *Mini Rev. Med. Chem.*, **17**, 959 (2017); <https://doi.org/10.2174/1389557516666151120115302>.
- M.F. Khan, M.M. Alam, G. Verma, W. Akhtar, M. Akhter and M. Shaquiquzzaman, *Eur. J. Med. Chem.*, **120**, 170 (2016); <https://doi.org/10.1016/j.ejmech.2016.04.077>.
- T.D. Penning, J.J. Talley, S.R. Bertenshaw, J.S. Carter, P.W. Collins, S. Docter, M.J. Graneto, L.F. Lee, J.W. Malecha, J.M. Miyashiro, R.S. Rogers, D.J. Rogier, S.S. Yu, G.D. Anderson, E.G. Burton, J.N. Cogburn, S.A. Gregory, C.M. Koboldt, W.E. Perkins, K. Seibert, A.W. Veenhuizen, Y.Y. Zhang and P.C. Isakson, *J. Med. Chem.*, **40**, 1347 (1997); <https://doi.org/10.1021/jm960803q>.
- N.K. Terrett, A.S. Bell, D. Brown and P. Ellis, *Bioorg. Med. Chem. Lett.*, **6**, 1819 (1996); [https://doi.org/10.1016/0960-894X\(96\)00323-X](https://doi.org/10.1016/0960-894X(96)00323-X).
- H.A. DeWald, S. Lobbestael and B.P.H. Poschel, *J. Med. Chem.*, **24**, 982 (1981); <https://doi.org/10.1021/jm00140a013>.
- M. Kim, C. Sim, D. Shin, E. Suh and K. Cho, *Crop Prot.*, **25**, 542 (2006); <https://doi.org/10.1016/j.cropro.2005.08.010>.
- R.E. Sammelson and J.E. Casida, *J. Org. Chem.*, **68**, 8075 (2003); <https://doi.org/10.1021/jo034520z>.
- D. Marcic, *Exp. Appl. Acarol.*, **36**, 177 (2005); <https://doi.org/10.1007/s10493-005-3579-2>.
- V. Nair, J. Mathew, J. Prabhakaran and R. Ruzziconi, *Chem. Soc. Rev.*, **26**, 127 (1997); <https://doi.org/10.1039/CS9972600127>.
- V. Nair, L. Balagopal, R. Rajan and J. Mathew, *Acc. Chem. Res.*, **37**, 21 (2004); <https://doi.org/10.1021/ar030002z>.
- S.B. Panicker, L.G. Nair, T.G. George and A. Augustine, *Synlett.*, **2003**, 156 (2003); <https://doi.org/10.1055/s-2003-36775>.
- A. Dhakshinamoorthy, *Synlett.*, **2005**, 3014 (2005); <https://doi.org/10.1055/s-2005-921893>.
- G.A. Molander, *Chem. Rev.*, **92**, 29 (1992); <https://doi.org/10.1021/cr00009a002>.
- B.B. Nayak, S. Sahu, S. Patel, S. Dash and B.K. Mishra, *Indian J. Chem.*, **47A**, 1486 (2008).
- V. Nair, L. Balagopal, R. Rajan and J. Mathew, *Acc. Chem. Res.*, **37**, 21 (2004); <https://doi.org/10.1021/ar030002z>.
- V. Nair and A. Deepthi, *Chem. Rev.*, **107**, 1862 (2007); <https://doi.org/10.1021/cr068408n>.
- T.J. Demars, M.K. Bera, S. Seifert, M.R. Antonio and R.J. Ellis, *Angew. Chem. Int. Ed.*, **54**, 7534 (2015); <https://doi.org/10.1002/anie.201502336>.
- X. Zeng, S. Ji and S. Wang, *Tetrahedron*, **61**, 10235 (2005); <https://doi.org/10.1016/j.tet.2005.08.040>.
- S.Y. Wang and S.J. Ji, *Tetrahedron*, **62**, 1527 (2006); <https://doi.org/10.1016/j.tet.2005.11.011>.
- G. Savitha and P.T. Perumal, *Tetrahedron Lett.*, **47**, 3589 (2006); <https://doi.org/10.1016/j.tetlet.2006.03.046>.
- V. Nair, K. Mohanan, T.D. Suja and E. Suresh, *Tetrahedron Lett.*, **47**, 705 (2006); <https://doi.org/10.1016/j.tetlet.2005.11.115>.
- R. Varala, R. Enugala, S. Nuvula and S.R. Adapa, *Synlett.*, **2006**, 1009 (2006); <https://doi.org/10.1055/s-2006-939066>.
- V. Sridharan, C. Avendaño and J.C. Menéndez, *Tetrahedron*, **63**, 673 (2007); <https://doi.org/10.1016/j.tet.2006.11.002>.
- D.E. Bergbreiter, J. Tian and C. Hongfa, *Chem. Rev.*, **109**, 530 (2009); <https://doi.org/10.1021/cr8004235>.
- N.R. Candeias, L.C. Branco, P.M.P. Gois, C.A.M. Afonso and A.F. Trindade, *Chem. Rev.*, **109**, 2703 (2009); <https://doi.org/10.1021/cr800462w>.
- Z. Andrade and L. Alves, *Curr. Org. Chem.*, **9**, 195 (2005); <https://doi.org/10.2174/1385272053369178>.
- J. Chen, S.K. Spear, J.G. Huddleston and R.D. Rogers, *Green Chem.*, **7**, 64 (2005); <https://doi.org/10.1039/b413546f>.
- R.D. Rogers, A.H. Bond, C.B. Bauer, J. Zhang, M.L. Jezl, D.M. Roden, S.D. Rein and R.R. Chomko, eds.: R.D. Rogers and M.A. Eiteman, *Aqueous Biphasic Separations: Biomolecules to Metal Ions*, Plenum: New York (1995).
- J.G. Huddleston, H.D. Willauer, K.R. Boaz and R.D. Rogers, *J. Chromatogr. B: Biomed. Appl.*, **711**, 237 (1998); [https://doi.org/10.1016/S0378-4347\(97\)00662-2](https://doi.org/10.1016/S0378-4347(97)00662-2).
- R.D. Rogers, H.D. Willauer, S.T. Griffin and J.G. Huddleston, *J. Chromatogr. B: Biomed. Appl.*, **711**, 255 (1998); [https://doi.org/10.1016/S0378-4347\(97\)00661-0](https://doi.org/10.1016/S0378-4347(97)00661-0).
- R.D. Rogers, A.H. Bond, J. Zhang and C.B. Bauer, *Appl. Radiat. Isot.*, **47**, 497 (1996); [https://doi.org/10.1016/0969-8043\(95\)00335-5](https://doi.org/10.1016/0969-8043(95)00335-5).
- R.D. Rogers, A.H. Bond, C.B. Bauer, J. Zhang, S.D. Rein, R.R. Chomko and D.M. Roden, *Solvent Extr. Ion Exch.*, **13**, 689 (1995); <https://doi.org/10.1080/07366299508918298>.
- R.D. Rogers, A.H. Bond and C.B. Bauer, *Sep. Sci. Technol.*, **28**, 139 (1993); <https://doi.org/10.1080/01496399308019483>.
- R.D. Rogers, C.B. Bauer and A.H. Bond, *Sep. Sci. Technol.*, **30**, 1203 (1995); <https://doi.org/10.1080/01496399508010341>.
- R.D. Rogers, A.H. Bond and J.L. Wolff, *J. Coord. Chem.*, **29**, 187 (1993); <https://doi.org/10.1080/00958979308037425>.
- R.D. Rogers, A.H. Bond, S. Aguinaga and A. Reyes, *J. Am. Chem. Soc.*, **114**, 2967 (1992); <https://doi.org/10.1021/ja00034a032>.
- R.D. Rogers, A.H. Bond and D.M. Roden, *Inorg. Chem.*, **35**, 6964 (1996); <https://doi.org/10.1021/ic960587b>.
- R.D. Rogers, J. Zhang and C.B. Bauer, *J. Alloys Compd.*, **249**, 41 (1997); [https://doi.org/10.1016/S0925-8388\(96\)02640-0](https://doi.org/10.1016/S0925-8388(96)02640-0).
- H.D. Willauer, J.G. Huddleston and R.D. Rogers, *Ind. Eng. Chem. Res.*, **41**, 1892 (2002); <https://doi.org/10.1021/ie010598z>.
- J.G. Huddleston, H.D. Willauer and R.D. Rogers, *J. Chromatogr. B: Biomed. Sci. Appl.*, **743**, 137 (2000); [https://doi.org/10.1016/S0378-4347\(00\)00230-9](https://doi.org/10.1016/S0378-4347(00)00230-9).
- H.D. Willauer, J.G. Huddleston, S.T. Griffin and R.D. Rogers, *Sep. Sci. Technol.*, **34**, 1069 (1999); <https://doi.org/10.1080/01496399908951081>.
- H.D. Willauer, J.G. Huddleston and R.D. Rogers, *Ind. Eng. Chem. Res.*, **41**, 2591 (2002); <https://doi.org/10.1021/ie0107800>.
- J.G. Huddleston, H.D. Willauer, R.D. Rogers, R.D. Rogers, P. Gaillard, P.-A. Carrault, B. Testa and M.H. Abraham, *Phys. Chem. Chem. Phys.*, **4**, 4065 (2002); <https://doi.org/10.1039/B203018G>.