Microwave Assisted Synthesis of Novel Functionalized Hydantoin Derivatives and Their Conversion to 5-(Z) Arylidene-4H-imidazoles

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Received: 24 May 2011; in revised form: 22 June 2011 / Accepted: 27 June 2011 / Published: 29 June 2011

Abstract: 2-(Alkyl-1-yl)-1H-imidazol-5(4H)-ones 5a–n were synthesized via nucleophilic substitution of the methylsulfanyl group of the corresponding 2-(methylthio)-1H-imidazol-5(4H)-ones 3a–c with suitably substituted secondary amines. The starting 2-thioxo-imidazolidin-4-ones 2a–b were prepared by condensation of thiohydantoin and benzo[b]-thiophene-3-carbaldehyde or benzofuran-3-carbaldehyde under microwave irradiation (MW) conditions. 2-Methylthio derivatives 3a–c were prepared by treatment of 2a–b with methyl iodide in the presence of aqueous sodium hydroxide.

Keywords: MW assisted synthesis; hydantoin; arylidene-4H-imidazoles

1. Introduction

Hydantoin derivatives have achieved considerable success as anticonvulsant agents [1]. The nucleosides of several 5-arylidene-3-arylhydantoins and 2-thiohydantoins show potent activity against human immunodeficiency virus (HIV) [2] and the leukemia subpanel [3]. A recent study showed that S-glucosylated hydantoins (Figure 1) act against herpes simplex virus, type 1 (HSV-1) and type 2 (HSV-2) in Vero cells [4]. A further study reported that thiazolidinediones (TZDs), which are known to have potent enhancing effects on insulin sensitivity, have been developed for the treatment of noninsulin-dependent diabetes mellitus [5,6].
It also been found that TZDs are high-affinity ligands for peroxisome proliferator-activated receptor-γ (PPAR-γ) and inhibit the production of monocyte chemo attractant protein 1 (MCP-1) in some human tissues [6]. Another group of TZDs revealed that troglitazone (Figure 1) acts on acute liver injury induced by ethanol and lipopolysaccharide [7]. For the past few years our group has been working on biologically important compounds [8-10]. Herein we report on the synthesis of variety of benzo[b]thiophene and benzofuran based hydantoin derivatives with the aim of investigating their antimicrobial and neuroprotecting properties.

2. Results and Discussion

As shown in Scheme 1, various substituted hydantion derivatives 5a–n were prepared by the condensation between benzo[b]thiophene-3-carbaldehyde and benzofuran-3-carbaldehyde with thiohydantoin using MW irradiation and a catalytic amount of 2,2,6,6-tetramethyl piperidine in ethanol that yielded (Z)-5-(benzo[b]thiophen-3-ylmethylene)-2-thioxoimidazolidin-4-one (2a) and (Z)-5-(benzofuran-3-ylmethylene)-2-thioxoimidazolidin-4-one (2b), respectively.

Scheme 1. Schematic representation for the synthesis of compounds 5a–n.
Since the starting material for 2b is not commercially available, it was prepared according to the literature procedure [11,12] which involved converting 2-hydroxy acetophenone to 3-methylbenzofuran [11] which was subsequently transformed to benzofuran-3-carbaldehyde by refluxing with SeO₂ in 1,4-dioxane [12].

Although many bases shown in Table 1 can be used as catalysts [e.g., piperidine, pyridine, N-methylpiperidine (NMP), DBU], 2,2,6,6-Tetramethylpiperidine (TMP) works best. The same reaction under conventional reflux condition using ethanol as solvent gave lower yields after longer times (5 h) and/or compounds that required rigorous purification. However the MW reaction provides cleaner reactions, shorter times (15 min) and purification of the products only required washing with cold ethanol. In addition the yields are good to excellent. The optimum temperature and condition for this MW assisted reaction was determined by a series of reactions of appropriate aldehyde 1 with hydantoin. The results are summarized in Table 1. The results indicate that MW irradiation at 90 °C for 15 min in ethanol is the optimum condition for the synthesis of 2a and 2b.

Table 1. Screening of solvents, reaction time and temperature for the synthesis of 2a.

| Entry | Base          | Condition  | Temp. (°C) | Time (min) | Yield (%) b |
|-------|---------------|------------|------------|------------|-------------|
| 1     | -             | No solvent | 90         | 15         | trace       |
| 2     | -             | Ethanol    | 90         | 15         | trace       |
| 3     | Piperidine    | Ethanol    | 90         | 15         | 80          |
| 4     | TMP           | Ethanol    | 90         | 15         | 96          |
| 5     | TMP           | No solvent | 90         | 15         | trace       |
| 6     | DBU/Pyridine  | Ethanol    | 90         | 20         | 20          |
| 7     | TMP           | Acetonitrile | 90   | 15         | 76          |
| 8     | TMP           | Acetonitrile | 130 | 15         | 15          |
| 9     | DBU           | Acetonitrile | 90   | 30         | trace       |
| 10    | TMP           | DMF        | 90         | 15         | 45          |
| 11    | NMP           | DMF        | 90         | 30         | 10          |
| 12    | DBU           | DMF        | 120–140    | 15         | trace       |
| 13    | TMP           | Water      | 90         | 15         | trace       |
| 14    | TMP           | Water      | 130        | 15         | trace       |
| 15    | -             | water      | 130        | 30         | trace       |
| 16    | TMP           | Tolune     | 90         | 15         | trace       |
| 17    | TMP           | Isopropanol | 90   | 15         | 45          |
| 18    | TMP           | THF        | 90         | 15         | 38          |
| 19    | TMP           | n-Butanol  | 90         | 15         | 33          |

a All the reaction was carried out in equimolar amount of each compound in 2 mL of solvent at 150 psi pressure; b Isolated yield.

The Z configuration of the compounds 2a and 2b was confirmed by comparison of previously reported [2] hydantoin derivatives along with nuclear Overhauser effect (NOE) experiments [4]. The vinylic proton in the ¹H-NMR spectra of benzo[b]thiophene and benzofuran derivatives appeared as singlets at 6.72 ppm and 6.58 ppm, respectively. The intermediates 2a and 2b were in turn transformed into the corresponding methylsulfanyl derivatives 3(a–c) by alkylation with methyl iodide in basic medium [13]. The use of 1.3 equiv. of alkyl halide and stirring for 4 h afforded 3a while use of excess
alkylating agent (2.5 equiv.) and stirring for 16 h afforded dialkylation giving 3b in high yield. The $^{13}$C-NMR spectrum of compounds 3a–c showed the absence of C=S signal of 2a, 2b at 179.50 ppm and the appearance of the C=N signal at 170.9 ppm, corresponding to s-alkylation. On the other hand, the $^1$H-NMR spectra of 2a and 2b showed two NH signals at 12.41 and 12.18 which were absent in the spectra of 3a and 3b, in which only a broad peak NH signal at 11.84 ppm appeared. Conversion of 2a to 3a via microwave irradiation was unsuccessful. The intermediate methylsulfanyl derivatives were subsequently converted into compounds 5a–n by nucleophilic substitution of the methylsulfanyl group with suitably substituted secondary amines [14] by using MW irradiation of a solution containing excess amine (>12.5 equiv.) and absolute ethanol at an elevated temperature (100 °C). The same reaction under conventional reflux condition using ethanol as solvent required longer time (16 h) and/or rigorous purification of products. As shown in Table 2, the 5-(Z) arylidene-4H-imidazole compounds 5a–n were formed in good to excellent yields. In most cases, the 5-(Z) arylidene-4H-imidazoles precipitated upon cooling the reaction mixture and only simple trituration with ethanol-hexane followed by recrystallization from ethanol afforded the product 5a–l. But in case of 5m and 5n, the crude reaction mixtures were purified by column chromatography using 40% ethyl acetate-hexane mixture (v/v). All the products were well characterized by $^1$H-NMR, $^{13}$C-NMR, IR and HRMS.

Table 2. MW assisted synthesis of various 5-(Z) arylidene-4H-imidazoles.

| Entry | X = S, R = H | R$_1$ = R$_2$ = R$_3$ = H, Y = CH$_2$ | % Yield a,b |
|-------|-------------|--------------------------------------|-------------|
| 1     |             | 91                                   |             |
| 2     |             | 90                                   |             |
| 3     |             | 86                                   |             |
| 4     |             | 89                                   |             |
| 5     |             | 80                                   |             |
| 6     |             | 92                                   |             |

The same reaction under conventional reflux condition using ethanol as solvent required longer time (16 h) and/or rigorous purification of products. As shown in Table 2, the 5-(Z) arylidene-4H-imidazole compounds 5a–n were formed in good to excellent yields. In most cases, the 5-(Z) arylidene-4H-imidazoles precipitated upon cooling the reaction mixture and only simple trituration with ethanol-hexane followed by recrystallization from ethanol afforded the product 5a–l. But in case of 5m and 5n, the crude reaction mixtures were purified by column chromatography using 40% ethyl acetate-hexane mixture (v/v). All the products were well characterized by $^1$H-NMR, $^{13}$C-NMR, IR and HRMS.

Table 2. MW assisted synthesis of various 5-(Z) arylidene-4H-imidazoles.
Table 2. Cont.

| Entry | Structure | % Yield | Remarks |
|-------|-----------|---------|---------|
| 7     | ![Structure 5g](image) | 88      |         |
| 8     | ![Structure 5h](image) | 85      |         |
| 9     | ![Structure 5i](image) | 86      |         |
| 10    | ![Structure 5j](image) | 86      |         |
| 11    | ![Structure 5k](image) | 91      |         |
| 12    | ![Structure 5l](image) | 90      |         |
| 13    | ![Structure 5m](image) | 88      |         |
| 14    | ![Structure 5n](image) | 77      |         |

* a Isolated yield; b All the products were characterized by IR, $^1$H-NMR, $^{13}$C-NMR, DEPT, HRMS and elemental analysis.

The use of aromatic amines for the same reaction was unsuccessful. From Table 1, it can be seen that in the case of 3,5-dimethylmorpholine (entry 14) the comparatively low yield probably reflects unfavorable steric interaction between the two methyl groups.
3. Experimental

General

The $^{1}$H- and $^{13}$C-NMR spectra were recorded on a 500-MHz Jeol multinuclear NMR spectrometer; chemical shifts were referenced to tetramethylsilane (TMS) as internal standard. Infrared (IR) spectra were obtained on a Varian 3100 Fourier transform (FT)-IR Spectrometer. Melting points were taken on a Melttemp apparatus. All chemicals and reagents were purchased from commercial sources. Mass spectra were obtained from Washington University, St. Louis, MO. Microwave experiments were carried out on CEM Discover microwave instrument.

$^{(Z)}$-5-(Benzo[b]thiophen-3-ylmethylene)-2-thioxoimidazolidin-4-one (2a) under MW irradiation. An equimolar quantity of benzo[b]thiophene-3-carbaldehyde and thiohydantoin were mixed in a mortar pestle and charged into a specially designed MW test tube. After the addition of ethanol (2 mL) followed by 1–2 drops of 2,2,6,6-tetramethylpiperidine, the test tube was sealed then irradiated for 15 min at 90 °C and 150 psi pressure. After cooling, the solid mass was scraped out of the test tube and placed into a flask containing 95% ethanol (20 mL) and filtered. The solid mass was collected then washed with ethanol (20 mL) and dried under reduced pressure and the residue was recrystallized from ethanol to afford 0.49 g (96%) of $^{(Z)}$-5-(benzo[b]thiophen-3-ylmethylene)-2-thioxoimidazolidin-4-one (2a) as yellow floppy crystals. m.p.: 259–261 °C; IR (KBr) $\nu$ (cm$^{-1}$): 3274, 1723, 1649, 1496; $^{1}$H-NMR (DMSO-$d_6$): $\delta$: 12.41 (brs, 1H, NH), 12.18 (brs, 1H, NH), 8.49 (s, 1H, Ar–CH), 8.02 (dd, $J = 7.45$ Hz, 8.00 Hz, 2H, Ar–CH), 7.46–7.42 (m, 2H, Ar–CH), 6.72 (s, 1H, vinylic proton); 13C-NMR (DMSO-$d_6$) $\delta$: 179.5 (C), 166.1 (C), 139.1 (C), 138.4 (C), 129.9 (CH), 129.0 (C), 127.8 (C), 125.7 (CH), 125.6 (CH), 123.6 (C), 122.0 (CH), 102.3 (CH); HRMS m/z: 261.0161 found (calculated for C$_{12}$H$_{8}$N$_{2}$O$_{2}$S$_{2}$, [M+H]$^+$ requires 261.0158).

$^{(Z)}$-5-(Benzofuran-3-ylmethylene)-2-thioxoimidazolidin-4-one (2b). Following the procedure for the preparation of 2a, the microwave-assisted reaction of equimolar quantities of benzofuran-3-carbaldehyde and thiohydantoin gave 0.51 g (96%) of 2b as a dark brown solid. m.p.: 307–308 °C; IR (KBr) $\nu$ (cm$^{-1}$): 3209, 1725, 1648, 1484, 1452; $^{1}$H-NMR (DMSO-$d_6$) $\delta$: 12.39 (brs, 1H, NH), 11.95 (brs, 1H, NH), 8.83 (s, 1H, Ar–CH), 7.88 (d, $J = 7$ Hz, 1H, Ar–CH), 7.61 (d, $J = 8$ Hz, 1H, Ar–CH), 7.38–7.30 (m, 2H, Ar–CH), 6.58 (s, 1H, vinylic proton); 13C-NMR (DMSO-$d_6$) $\delta$: 179.2 (C), 165.7 (C), 154.7 (C), 147.1 (CH), 128.7 (C), 126.6 (C), 125.9 (CH), 124.1 (CH), 120.3 (CH), 114.3 (CH), 112.0 (CH), 99.8 (CH); HRMS m/z: 245.0390 found (calculated for C$_{12}$H$_{8}$N$_{2}$S$_{2}$, [M+H]$^+$ requires 245.0386).

$^{(Z)}$-4-(Benzo[b]thiophen-3-ylmethylene)-(2-methylthio)-1H-imidazol-5(4H)-one (3a). Ethyl iodide (2.15 g, 15.1 mmol) was added to a solution of 2a (2.9 g, 11.1 mmol), in 15% aqueous NaOH (4.5 mL) and 95% methanol (30 mL) and the resulting reaction mixture was stirred for 4 h at room temperature. Most of the solvent was removed by distillation and to the precipitate, was added 15 mL of water and stirred for 10 min and filtered and washed first with water (200 mL) and then with diethyl ether (50 mL) to supply 2.85 g of $^{(Z)}$-4-(benzo[b]thiophen-3-ylmethylene)-(2-methylthio)-1H-imidazol-5(4H)-one (3a) as a bright yellow solid. m.p.: 276–278 °C; IR (KBr) $\nu$ (cm$^{-1}$): 3058, 1701, 1627, 1555, 1492; $^{1}$H-NMR (DMSO-$d_6$) $\delta$: 11.84 (brs, 1H, NH), 8.94 (s, 1H, Ar–CH), 8.11 (d, $J = 7.5$ Hz, 1H, Ar–CH), 7.38 (d, $J = 8$ Hz, 1H, Ar–CH), 7.28–7.24 (m, 2H, Ar–CH), 7.12 (d, $J = 8$ Hz, 1H, Ar–CH), 6.58 (s, 1H, vinylic proton); 13C-NMR (DMSO-$d_6$) $\delta$: 179.5 (C), 165.7 (C), 154.7 (C), 147.1 (CH), 128.7 (C), 126.6 (C), 125.9 (CH), 124.1 (CH), 120.3 (CH), 114.3 (CH), 112.0 (CH), 99.8 (CH); HRMS m/z: 245.0390 found (calculated for C$_{12}$H$_{8}$N$_{2}$S$_{2}$, [M+H]$^+$ requires 245.0386).
8.01 (d, J = 8Hz, 1H, Ar–CH), 7.44–7.40 (m, 2H, Ar–CH), 7.04 (s, 1H, vinylic proton), 2.66 (s, 3H, SCH₃); ¹³C-NMR (DMSO-d₆) δ: 170.9 (C), 165.9 (C), 140.2 (C), 139.4 (C), 138.5 (C), 133.0 (CH), 130.1 (C), 125.4 (CH), 123.5 (CH), 121.9 (CH), 111.2 (CH), 12.9 (SCH₃); HRMS m/z: 275.0322 found (calculated for C₁₃H₁₀N₂O₂S₂, [M+H]^+ requires 275.0315).

(Z)-4-(Benzo[b]thiophen-3-ylmethylene)-1-methyl-2-(methylthio)-1H-imidazol-5(4H)-one (3b). Compound 3b was prepared in same manner as 3a with the exception that an excess (2.5 equiv.) of methyl iodide was used and stirring was carried out for 16 h. This reaction afforded 2.9 g (90% yield) of 3b as a light brown solid. m.p.: 260–263 °C; IR (KBr) ν (cm⁻¹): 1700, 1643, 1540, 1497; ¹H-NMR (DMSO-d₆) δ: 8.99 (s, 1H, Ar–CH), 8.16 (d, J = 7.5 Hz, 1H, Ar–CH), 8.03 (d, J = 7.5 Hz, 1H, Ar–CH), 7.46–7.41 (m, 2H, Ar–CH), 7.14 (s, 1H, vinylic proton), 3.06 (s, 3H, NCH₃), 2.72 (s, 3H, SCH₃); ¹³C-NMR (DMSO-d₆) δ: 169.1 (C), 166.9 (C), 139.4 (C), 139.1 (C), 138.5 (C), 133.6 (CH), 130.0 (C), 125.5 (CH), 123.5 (CH), 122.1 (CH), 112.4 (CH), 26.9 (CH₃), 13.2 (CH₃); HRMS m/z: 289.0483 found (calculated for C₁₄H₁₂N₂O₂S₂, [M+H]^+ requires 289.0471).

(Z)-4-(Benzofuran-3-ylmethylene)-1-methyl-2-(methylthio)-1H-imidazol-5(4H)-one (3c). Compound 3c was obtained as a brown solid in 89% when prepared by the same method used for the preparation of 3a. m.p.: 168–171 °C; IR (KBr) ν (cm⁻¹): 1701, 1642, 1541, 1496 cm⁻¹; ¹H NMR (DMSO-d₆): δ 8.73 (s, 1H, Ar–CH), 8.29 (d, J = 7.45 Hz, 1H, Ar–CH), 7.60 (d, J = 7.45 Hz, 1H, Ar–CH), 7.35–7.29 (m, 2H, Ar–CH), 7.05 (s, 1H, vinylic proton), 3.03 (s, 3H, NCH₃), 2.70 (s, 3H, SCH₃); ¹³C-NMR (DMSO-d₆) δ: 168.5 (C), 165.7 (C), 155.2 (C), 150.4 (CH), 138.7 (C), 126.1 (C), 125.8 (CH), 124.0 (CH), 121.9 (CH), 117.0 (C), 112.3 (CH), 112.0 (CH), 26.8 (CH₃), 13.2 (CH₃); HRMS m/z: 273.0701 found (calculated for C₁₄H₁₂N₂O₂S, [M+H]^+ requires 273.0699).

(Z)-4-(Benzo[b]thiophen-3-ylmethylene)-2-(piperidin-1-yl)-1H-imidazol-5(4H)-one (5a). Compound 3a and piperidine (excess, 12.5 equiv.) were mixed then placed in a specially designed MW test tube. Ethanol (2 mL) was added to the mixture. The test tube was then sealed and then irradiated for 30 min at 100 °C and 150 psi pressure. After cooling, the solid mass was filtered and crashed into 20 mL of 95% ethanol. The solid mass collected was washed with ethanol-hexane mixture (1:5 v/v) and dried over vacuum to get the desired product. The titled compound was obtained as bright yellow solid. m.p.: 275–277 °C; IR (KBr) ν (cm⁻¹): 3111.7, 1695, 1643, 1588; ¹H-NMR (DMSO-d₆) δ: 11.20 (brs, 1H, NH), 8.67 (s, 1H, Ar–CH), 7.98 (dd, J = 4.5 Hz, 7.4 Hz, 2H, Ar–CH), 7.42–7.37 (m, 2H, Ar–CH), 6.63 (s, 1H, vinylic proton), 3.59 (brs, 4H, CH₂ × 2), 1.58–1.55 (m, 6H, CH₂ × 3); ¹³C-NMR (DMSO-d₆) δ: 172.5 (C), 159.3 (C), 139.4 (C), 138.7 (C), 131.4 (C), 128.0 (CH), 125.1 (CH), 125.0 (CH), 123.4 (CH), 121.7 (CH), 101.5 (CH), 45.0 (NCH₂), 25.7 (CH₂), 24.1 (CH₃); HRMS m/z: 312.1173 found (calculated for C₁₇H₁₇N₃OS, [M+H]^+ requires 312.1165) All the compounds 5b–n were prepared in the same way.

(Z)-4-(Benzo[b]thiophen-3-ylmethylene)-2-morpholino-1H-imidazol-5(4H)-one (5b). This compound was obtained as light yellow solid. m.p.: 292–294 °C; IR (KBr) ν (cm⁻¹): 3140, 1686, 1648, 1588, 1423; ¹H-NMR (DMSO-d₆) δ: 11.29 (brs, 1H, NH), 8.72 (s, 1H, Ar–CH), 7.99 (dd, J = 4.5 Hz, 7.6 Hz, 2H, Ar–CH), 7.42–7.37 (m, 2H, Ar–CH), 6.69 (s, 1H, vinylic proton), 5.67–3.62 (m, 8H, CH₂ × 4); ¹³C-NMR (DMSO-d₆) δ: 172.2 (C), 159.4 (C), 139.4 (C), 138.7 (C), 131.2 (C), 128.5 (CH), 125.2 (CH),
123.5 (CH), 121.8 (CH), 102.7 (CH), 66.1 (OCH 2), 40.3 (NCH 2); HRMS m/z: 314.0963 found (calculated for C16H15N3O2S, [M+H] + requires 314.0957).

(Z)-4-(Benzo[b]thiophen-3-ylmethylene)-2-(3-methylpiperidin-1-yl)-1H-imidazol-5(4H)-one (5c). This compound was obtained as white solid. m.p.: 242–244 °C; IR (KBr) ν (cm−1): 3206, 1692, 1616, 1426, 1407; 1H-NMR (DMSO-d6) δ: 11.15 (brs, 1H, NH), 8.66 (s, 1H, Ar–CH), 7.99 (dd, J = 4.3 Hz, 7.9 Hz, 2H, Ar–CH), 7.42–7.37 (m, 2H, Ar–CH), 6.62 (s, 1H, vinylic proton), 4.25 (brs, 2H, CH2), 3.01 (dd, J = 10 Hz, 10.5 Hz, 1H, CH 2), 2.70 (brs, 1H, CH 2), 1.76–1.14 (m, 4H, CH 2 × 2), 1.17–1.13 (m, 1H, CH), 0.87 (d, J = 4.5 Hz, 3H, CH3); 13C-NMR (DMSO-d6) δ: 172.3 (C), 159.5 (C), 139.4 (C), 138.7 (C), 131.4 (C), 127.9 (CH), 125.1 (CH), 125.0 (CH), 123.4 (CH), 121.7 (CH), 101.4 (CH), 51.2 (NCH2), 45.3 (NCH2), 32.6 (CH2), 31.0 (CH), 24.9 (CH2), 18.9 (CH3); HRMS m/z: 326.1334 found (calculated for C18H19N3OS, [M+H] + requires 326.1322).

(Z)-4-(Benzo[b]thiophen-3-ylmethylene)-1-methyl-2-(piperidin-1-yl)-1H-imidazol-5(4H)-one (5d). This compound was obtained as light yellow solid. m.p.: 165–168 °C; IR (KBr) ν (cm−1): 1713, 1617, 1560, 1451; 1H-NMR (DMSO-d6) δ: 8.77 (s, 1H, Ar–CH), 8.02 (dd, J = 7.2 Hz, 7.8 Hz, 2H, Ar–CH), 7.43–7.38 (m, 2H, Ar–CH), 6.84 (s, 1H, vinylic proton), 3.55 (s, 4H, CH 2 × 2), 3.17 (s, 3H, NCH3), 1.62 (s, 6H, CH2 × 3); 13C-NMR (DMSO-d6) δ: 171.5 (C), 161.7 (C), 140.3 (C ), 139.4 (C), 138.6 (C), 131.1 (CH), 129.5 (CH), 125.2 (CH), 125.1 (CH), 123.5 (CH), 121.9 (CH), 105.3 (CH), 48.3 (NCH2), 30.4 (NCH3), 25.6 (CH2), 24.2 (CH2); HRMS m/z: 326.1335 found (calculated for C18H19N3OS, [M+H] + requires 326.1322).

(Z)-4-(Benzo[b]thiophen-3-ylmethylene)-2-(3,5-dimethylmorpholino)-1H-imidazol-5(4H)-one (5e). This compound was obtained as bright yellow solid. m.p.: 240–242 °C. IR (KBr) ν (cm−1): 3112, 1696, 1640, 1584, 1452; 1H-NMR (DMSO-d6) δ: 11.30 (brs, 1H, NH), 8.74 (s, 1H, Ar–CH), 7.99 (dd, J = 4.1 Hz, 7.8 Hz, 2H, Ar–CH), 7.42–7.37 (m, 2H, Ar–CH), 6.66 (s, 1H, vinylic proton), 4.01–3.98 (m, 2H, CH), 3.61–3.60 (m, 2H, CH2), 2.71 (brs, 2H, CH2), 1.13–1.11 (m, 6H, CH3 × 2); 13C-NMR (DMSO-d6) δ: 172.0 (C), 158.8 (C), 142.3 (C), 139.4 (C), 138.7 (C), 131.2 (CH), 129.5 (CH), 125.2 (CH), 125.1 (CH), 123.5 (CH), 121.7 (CH), 102.7 (CH), 71.3 (OCH3), 60.1 (CH) 19.0 (CH3); HRMS m/z: 342.1282 found (calculated for C18H19N3O2S, [M+H] + requires 342.1271).

(Z)-4-(Benzo[b]thiophen-3-ylmethylene)-1-methyl-2-morpholino-1H-imidazol-5(4H)-one (5f). This compound was obtained as bright yellow solid. m.p.: 154–156 °C. IR (KBr) ν (cm−1): 1717, 1615, 1651, 1465; 1H-NMR (DMSO-d6) δ: 8.79 (s, 1H, Ar–CH), 8.05 (dd, J = 7.45 Hz, 7.7 Hz, 2H, Ar–CH), 7.43–7.38 (m, 2H, Ar–CH), 6.89 (s, 1H, vinylic proton), 3.72–3.70 (m, 4H, CH 2 × 2), 3.62–3.60 (m, 4H, CH 2 × 2), 3.18 (s, 3H, NCH3); 13C-NMR (DMSO-d6) δ: 171.2 (C), 161.5 (C), 139.9 (C), 139.3 (C), 138.6 (C), 130.9 (CH), 130.1 (CH), 125.3 (CH), 125.2 (CH), 123.5 (CH), 121.9 (CH), 106.2 (CH), 66.1 (OCH2), 40.3 (NCH2), 30.2 (NCH3); HRMS m/z: 328.1123 found (calculated for C17H17N3O2S, [M+H] + requires 328.1114).

(Z)-4-(Benzo[b]thiophen-3-ylmethylene)-1-methyl-2-(3-methylpiperidin-1-yl)-1H-imidazol-5(4H)-one (5g). This compound was obtained as light yellow solid. m.p.: 150–152 °C; IR (KBr) ν (cm−1): 1709, 1637, 1561, 1457, 1439; 1H-NMR (DMSO-d6) δ: 8.75 (s, 1H, Ar–CH), 8.05 (dd, J = 7.2 Hz, 7.8 Hz, 2H, Ar–CH), 7.43–7.38 (m, 2H, Ar–CH), 6.89 (s, 1H, vinylic proton), 3.72–3.70 (m, 4H, CH 2 × 2), 3.62–3.60 (m, 4H, CH 2 × 2), 3.18 (s, 3H, NCH3); 13C-NMR (DMSO-d6) δ: 171.2 (C), 161.5 (C), 139.9 (C), 139.3 (C), 138.6 (C), 130.9 (CH), 130.1 (CH), 125.3 (CH), 125.2 (CH), 123.5 (CH), 121.9 (CH), 106.2 (CH), 66.1 (OCH2), 40.3 (NCH2), 30.2 (NCH3); HRMS m/z: 328.1123 found (calculated for C17H17N3O2S, [M+H] + requires 328.1114).
2H, Ar–CH), 7.43–7.38 (m, 2H, Ar–CH), 6.83 (s, 1H, vinylic proton), 4.01–3.97 (m, 2H, CH₂), 3.18 (s, 3H, NCH₃), 3.02–2.9 (m, 1H, CH), 2.73–2.70 (m, 2H, CH₂), 1.72–1.69 (m, 4H, CH₂ × 2), 1.17–1.13 (m, 2H, CH₂), 0.90 (d, J = 6.3 Hz, 3H, CH₃); 13C-NMR (DMSO-d₆) δ: 171.5 (C), 161.6 (C), 140.3 (C), 139.4 (C), 138.6 (C), 131.1 (C), 129.5 (CH), 125.3 (CH), 125.2 (CH), 123.5 (CH), 121.9 (CH), 105.2 (CH), 54.3 (CH₂), 47.8 (CH₂), 32.7 (CH₂), 31.0 (CH₂), 30.4 (CH), 25.01 (CH₃), 19.3 (CH₃); HRMS m/z: 340.1485 found (calculated for C₁₉H₂₁N₃OS, [M+H]+ requires 340.1478).

(Z)-4-(Benzothiophen-3-ylmethylene)-2-thiomorpholino-1H-imidazol-5(4H)-one (5h). This compound was obtained as bright yellow solid. m.p.: 269–271 °C; IR (KBr) ν (cm⁻¹): 3110, 1694, 1640, 1585, 1490, 1453; ¹H-NMR (DMSO-d₆) δ: 11.31 (brs, 1H, NH), 8.70 (s, 1H, Ar–CH), 8.00–7.97 (m, 2H, Ar–CH), 7.42–7.37 (m, 2H, Ar–CH), 6.67 (s, 1H, vinylic proton), 3.89 (brs, 4H, CH₂ × 2), 2.68 (s, 4H, CH₂ × 2); 13C-NMR (DMSO-d₆) δ: 172.5 (C), 158.9 (C), 139.5 (C), 138.6 (C), 131.5 (C), 128.7 (CH), 125.5 (CH), 123.5 (CH), 121.3 (CH), 102.1 (CH), 46.3 (NCH₂), 26.3 (SCH₃); HRMS m/z: 330.0737 found (calculated for C₁₆H₁₅N₃OS₂, [M+H]+ requires 330.0729).

(Z)-4-(Benzothiophen-3-ylmethylene)-2-(4-methylpiperazin-1-yl)-1H-imidazol-5(4H)-one (5i). This compound was obtained as yellow solid. m.p.: 234–236 °C; IR (KBr) ν (cm⁻¹): 3108, 1697, 1646, 1591, 1448; ¹H-NMR (DMSO-d₆) δ: 11.15 (brs, 1H, NH), 8.68 (s, 1H, Ar–CH), 7.99 (dd, J = 4 Hz, 8 Hz, 2H, Ar–CH), 7.42–7.37 (m, 2H, Ar–CH), 6.64 (s, 1H, vinylic proton), 3.61 (brs, 4H, CH₂ × 2), 2.37 (s, 4H, CH₂ × 2), 2.19 (s, 3H, NCH₃); 13C-NMR (DMSO-d₆) δ: 172.6 (C), 159.6 (C), 139.3 (C), 138.6 (C), 131.3 (C), 128.4 (CH), 125.1 (CH), 123.5 (CH), 121.7 (CH), 102.2 (CH), 54.4 (NCH₂), 46.0 (NCH₃), 45.4(NCH₃); HRMS m/z: 327.1290found (calculated for C₁₇H₁₈N₄OS, [M+H]+ requires 327.1281).

(Z)-4-(Benzofuran-3-ylmethylen)-1-methyl-2-(piperidin-1-yl)-1H-imidazol-5(4H)-one (5j). This compound was obtained as brown solid. m.p.: 148–150 °C; IR (KBr) ν (cm⁻¹): 1713, 1649, 1556, 1450; ¹H-NMR (DMSO-d₆) δ: 8.61 (s, 1H, Ar–CH), 8.14 (d, J = 7.45 Hz, 1H, Ar–CH), 7.57 (d, J = 7.45 Hz, 1H, Ar–CH), 7.34–7.28 (m, 2H, Ar–CH), 6.70 (s, 1H, vinylic proton), 3.53 (s, 4H, CH₂ × 2), 3.15 (s, 3H, NCH₃), 1.62 (s, 6H, CH₂ × 3); 13C-NMR (DMSO-d₆) δ: 170.8 (C), 161.0 (C), 154.9 (C), 148.0 (CH), 140.0 (C), 126.6 (C), 125.4 (CH), 123.6 (CH), 121.4 (CH), 117.4 (C), 111.9 (CH), 104.4 (CH), 48.3 (NCH₃), 30.3 (NCH₃), 25.6 (CH₂), 24.2 (CH₂); HRMS m/z: 310.1561 found (calculated for C₁₈H₁₉N₃O₂, [M+H]+ requires 310.1557).

(Z)-4-(Benzofuran-3-ylmethylene)-1-methyl-2-morpholino-1H-imidazol-5(4H)-one (5k). This compound was obtained as yellow solid. m.p.: 142–145 °C; IR (KBr) ν (cm⁻¹): 1708, 1644, 1557, 1452; ¹H-NMR (ACETONE-d₆) δ: 8.62 (s, 1H, Ar–CH), 8.06 (d, J = 7.00 Hz, 1H, Ar–CH), 7.51 (d, J = 7.00 Hz, 1H, Ar–CH), 7.35–7.29 (m, 2H, Ar–CH), 6.77 (s, 1H, vinylic proton), 3.77 (s, 4H, OCH₂ × 2), 3.60 (s, 4H, NCH₃), 3.19 (s, 3H, NCH₃); ¹³C-NMR (ACETONE-d₆) δ: 170.1 (C), 160.9 (C), 155.0 (C), 148.0 (CH), 139.7 (C), 126.7 (C), 124.8 (CH), 123.1(CH), 120.4 (CH), 117.1 (C), 111.3 (CH), 105.3 (CH), 65.9 (OCH₂), 47.5 (NCH₂), 28.9 (NCH₃); HRMS m/z: 312.1358 found (calculated for C₁₇H₁₉N₃O₃, [M+H]+ requires 312.1350).

(Z)-4-(Benzofuran-3-ylmethylene)-1-methyl-2-thiomorpholino-1H-imidazol-5(4H)-one (5l). This compound was obtained as yellowish solid. m.p.: 157–159 °C; IR (KBr) ν (cm⁻¹): 1703, 1649, 1541,
1450; \[^1\text{H-NMR (DMSO-} \text{d}_6)\delta: 8.62\ (s, 1H, Ar–CH), 8.14\ (d, J = 7.45 \text{ Hz, 1H, Ar–CH}), 7.58\ (d, J = 7.45 \text{ Hz, 1H, Ar–CH}), 7.33–7.29\ (m, 2H, Ar–CH), 6.76\ (s, 1H, vinylic proton), 3.83–3.81\ (m, 4H, CH\_2 \times 2), 3.15\ (s, 3H, NCH\_3), 2.77–2.75\ (m, 4H, CH\_2 \times 2); \[^1\text{C-NMR (DMSO-} \text{d}_6)\delta: 170.6\ (C), 160.9\ (C), 154.9\ (C), 148.3\ (CH), 139.6\ (C), 126.6\ (C), 125.5\ (CH), 123.7\ (CH), 121.4\ (CH), 117.3\ (C), 111.9\ (CH), 105.5\ (CH), 49.9\ (NCH\_2), 30.2\ (SCH\_2), 26.5\ (NCH\_3); HRMS m/z: 328.1132 found (calculated for C\_17H\_17N\_3O\_2S, \[M+H]\^+ requires 328.1121).

(\text{Z})\text{-4-(Benzofuran-3-ylmethylene)-1-methyl-2-(4-methylpiperazin-1-yl)-1H-imidazol-5(4H)-one (5m). This compound was obtained as a low melting yellow solid. IR (KBr) \nu (\text{cm}^{-1}): 1708, 1644, 1559, 1450; \[^1\text{H-NMR (DMSO-} \text{d}_6 + \text{acetone-} \text{d}_6)\delta: 9.09\ (s, 1H, Ar–CH), 8.57\ (d, J = 7.45 \text{ Hz, 1H, Ar–CH}), 8.00\ (d, J = 7.45 \text{ Hz, 1H, Ar–CH}), 7.81–7.74\ (m, 2H, Ar–CH), 7.21\ (s, 1H, vinylic proton), 4.07\ (t, J = 4.6 \text{ Hz, 4H, CH} \_2 \times 2), 3.66\ (s, 3H, CH\_3), 2.93\ (t, J = 4.6 \text{ Hz, 4H, CH} \_2 \times 2), 2.68\ (s, 3H, CH\_3); \[^1\text{C-NMR (DMSO-} \text{d}_6 + \text{ACETONE-} \text{d}_6)\delta: 170.9\ (C), 161.4\ (C), 155.4\ (C), 148.5\ (CH), 140.3\ (C), 127.1\ (C), 125.6\ (CH), 123.8\ (CH), 121.4\ (CH), 117.7\ (C), 112.0\ (CH), 105.4\ (CH), 54.8\ (CH\_2), 47.6\ (CH\_2), 46.2\ (CH\_3), 30.2\ (CH\_3); HRMS m/z: 325.1670 found (calculated for C\_18H\_20N\_4O\_2, \[M+H]\^+ requires 325.1666).

(\text{Z})\text{-4-(Benzofuran-3-ylmethylene)-2-(3,5-dimethylmorpholino)-1-methyl-1H-imidazol-5(4H)-one (5n). This compound was obtained as bright yellow solid. m.p.: 137–140 °C; IR (KBr) \nu (\text{cm}^{-1}): 1703, 1641, 1557, 1456; \[^1\text{H-NMR (DMSO-} \text{d}_6)\delta: 8.63\ (s, 1H, Ar–CH), 8.15\ (d, J = 7.45 \text{ Hz, 1H, Ar–CH}), 7.58\ (d, J = 7.45 \text{ Hz, 1H, Ar–CH}), 7.33–7.28\ (m, 2H, Ar–CH), 6.74\ (s, 1H, vinylic proton), 3.98\ (d, J = 12.6 \text{ Hz, 2H, CH} \_2), 3.71–3.69\ (m, 2H, CH\_2), 3.18\ (s, 3H, CH\_3), 2.78–2.75\ (m, 2H, CH\_2), 1.13\ (d, J = 12.6 \text{ Hz, 6H, CH} \_3 \times 2); \[^1\text{C-NMR (DMSO-} \text{d}_6)\delta: 170.7\ (C), 160.4\ (C), 154.9\ (C), 148.3\ (CH), 139.6\ (C), 126.6\ (C), 125.8\ (CH), 123.6\ (CH), 121.4\ (CH), 117.3\ (C), 112.0\ (CH), 105.3\ (CH), 71.2\ (OCH\_2), 52.2\ (CH), 30.2\ (CH\_3), 19.0\ (CH\_3); HRMS m/z: 340.1671 found (calculated for C\_19H\_21N\_3O\_3, \[M+H]\^+ requires 340.1663).

4. Conclusions

In summary we have successfully developed a synthetic method that provides ready access to novel biologically important benzo[b]thiophene and benzofuran based thiohydantoin derivatives. We are currently investigating the synthesis of a number of other thiohydantoin-based drug molecules by this method. A detailed biological activity study (antibacterial, antifungal, anticancer and neuroprotective kinase inhibitor activity) of these important compounds is being carried out. Preliminary results indicate that many of the thiohydantoins exhibit excellent neuroprotective properties.

Acknowledgments

The authors are grateful to NIH (1RC2NS064950) for generous financial support.
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Sample Availability: Samples of the compounds 5a–n are available from the authors.

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