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A new facet of the reaction of nitro heteroaromatic compounds with ethyl isocyanoacetate

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Nitro heteroarenes react with ethyl isocyanoacetate in the presence of 1,8-diazabicyclo[5.4.0]undecene (DBU) to give pyrroles or pyrimidine *N*-oxide depending on the structure of the starting nitro compounds. For example, 4-nitro-2,1,3-benzothiadiazole **3a** reacted with ethyl isocyanoacetate to give ethyl 2,1,3-benzothiadiazolo[3,4-*c*]pyrrole-2-carboxylate **4a** (33%), while a similar reaction with 5-nitro-2,1,3-benzothiadiazole **3b** gave the corresponding compound **4b** (21%) as a sole product. A plausible mechanism for these reactions is presented.


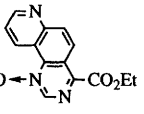
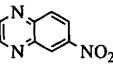
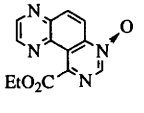
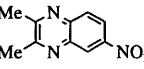
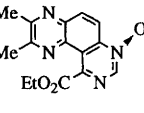
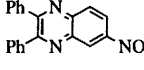
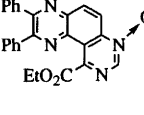
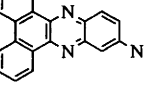
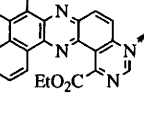

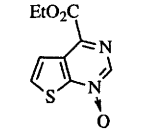
Since substituted pyrroles are of great interest in the preparation of functional compounds such as porphyrins and polypyrroles, many syntheses of them have been developed.¹ Compounds containing a pyrrole ring such as the isoindoles are important for the preparation of conducting polypyrroles with low band-gaps and highly conjugated porphyrins, but there are no good general methods for obtaining them.² Recent pyrrole syntheses from nitroalkenes and isocyanoacetates open a new way to the introduction of various substituents into the β -positions of the pyrrole ring.^{3,4} Furthermore, we and Lash have found that polycyclic aromatic nitro compounds react with isocyanoacetates in the presence of 1,8-diazabicyclo[5.4.0]undecene (DBU) to give the pyrroles fused with aromatic rings, and the pyrroles thus obtained are useful precursors for low band-gap polypyrroles^{3a} or highly conjugated porphyrins.⁵ If this pyrrole synthesis can be extended to nitroalkenes substituted with heteroatoms, pyrroles substituted with heteroatoms can be prepared. However, the reaction of 2-nitro enamines or 2-nitroalkenyl sulfides with ethyl isocyanoacetate failed to give the expected pyrroles, 1-hydroxypyrazoles being obtained instead.⁶ Thus, nitroalkenes having heteroatoms at the β -position react with isocyanoacetates in a different way from those of simple nitroalkenes. In this paper we deal with the reaction of nitro heteroaromatics with isocyanoacetates, in which a third aspect of the reaction of nitro compounds with isocyanoacetates is presented, namely the base-catalysed reaction of heteroaromatic nitro compounds with ethyl isocyanoacetate to give the corresponding pyrroles or pyrimidine *N*-oxides depending on the starting nitro compounds.

Results and discussion

During our study of the synthesis of the annulated pyrroles mentioned above, we found that 5-nitroquinoline **1a** reacted with ethyl isocyanoacetate in the presence of DBU to give the pyrimidine *N*-oxide **2a** (5%) instead of the corresponding and expected pyrroles. In this reaction, most of starting material was recovered unchanged from the reaction mixture. Some similar examples are listed in Table 1.

6-Nitroquinoxaline **1b** and its derivatives **1c,d** gave the pyrimidine *N*-oxides **2b-d** whilst 2-nitrothiophene **1e** under similar condition gave pyrimidine *N*-oxide **2e** (40%). Compounds **2a-e** were identified from their NMR and high

Table 1 The fused pyrimidine *N*-oxides **2a-f** were prepared in THF as solvent at room temperature; the reaction time was 48 h except for **2f** for which it was 24 h

Substrate	Product
 1a	 2a
 1b	 2b
 1c	 2c
 1d	 2d
 1e	 2e
 1f	 2f

resolution mass spectral results; the position of the oxygen was confirmed for compound **2e** by conversion of the latter into pyrimidine on treatment with PCl_3 .

Since pyrimidine bases exist in nature as nucleosides,

the constituents of nucleic acid, they have attracted much attention. Although there are numerous methods for construction of the pyrimidine ring,⁷ this is not so for the direct synthesis of pyrimidine *N*-oxides and the corresponding annulated compounds. An alternative method of preparing pyrimidine *N*-oxides is by *N*-oxidation of the corresponding pyrimidines with peracetic acid, although this gives both a mixture of 1- and 3-oxides⁸ and the possibility of side reactions.⁹

For these reasons, we decided to develop a convenient method of pyrimidine ring synthesis, although in this paper the main stress falls on the diversity of the reactions between aromatic nitro compounds and ethyl isocynoacetate. From this point of view the reactions of the nitro-2,1,3-benzothiadiazoles **3a,b** were particularly noteworthy since the position of the nitro group was the sole factor determining the product. 4-Nitro-2,1,3-benzothiadiazole **3a** reacted with ethyl isocynoacetate in the presence of DBU to give the pyrrole **4a** as the sole product. This reaction was conducted as follows. DBU (1 equiv.) was added dropwise to a stirred solution containing 1 equiv. each of 4-nitro-2,1,3-benzothiadiazole **3a** and ethyl isocynoacetate in dry THF kept at *ca.* 0 °C in an ice bath. After the addition the mixture was allowed to rise slowly to ambient temperature. After 5 h, the reaction was quenched with hydrochloric acid and the mixture worked up to give a brown solid, which was column chromatographed (SiO₂, CHCl₃) to give the pyrrole **4a** (33%). In contrast, use of 5-nitro-2,1,3-benzothiadiazole **3b**¹⁰ as the starting material under similar reaction conditions gave the pyrimidine *N*-oxide **4b** as the sole product with recovery of most of the starting material. HPLC analysis and GC-MS measurement showed that there was no cross-contamination of the two products.

The homologues of **3a** and **3b**, the nitro-2,1,3-benzoselenadiazoles **5a,b** were prepared by a literature procedure (Scheme 1):^{10b} thus, condensation of *o*-phenylenediamine with selenium dioxide followed by nitration with mixed acid gave 4-nitro-2,1,3-benzoselenadiazole **5a** whilst use of 4-nitrophenylene-1,2-diamine in place of *o*-phenylenediamine gave 5-nitro-2,1,3-benzoselenadiazole **5b**.

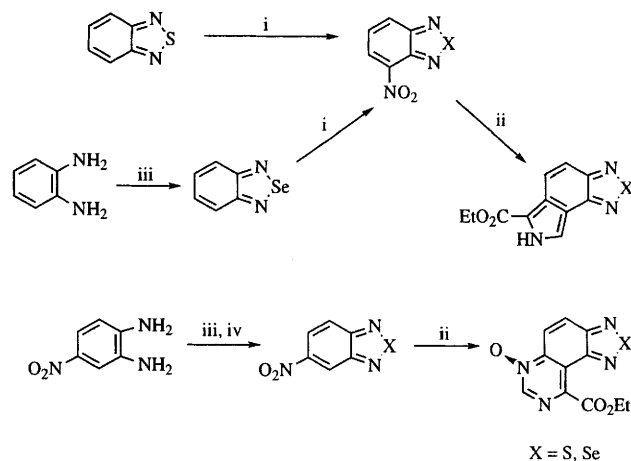
These seleno compounds **5a** and **5b** with ethyl isocynoacetate in the presence of DBU gave results similar to those of the nitro-2,1,3-benzothiadiazoles **3a** and **3b**; see Scheme 1 and Table 2.

Proposed mechanisms for the formation of pyrroles and pyrimidines are illustrated in Scheme 2. Initial attack of the ethyl isocynoacetate anion occurred at the β-position to the nitro groups to form the anionic intermediate **7**. When the nitro group was co-planar with the aromatic ring, this intermediate could be represented by two resonance structures **7** and **8** owing to the ambident character of the nitro group. Subsequent cyclization of the intermediate **7** gave the annulated pyrrole **9** whilst, for **8**, an intramolecular reaction of the carbon atom of the isocyanide moiety and the oxyanion yielded the annulated pyrimidine *N*-oxide **10**.

In order to study these reactions, an equimolar mixture of **3a** and **3b** was allowed to react with an equivalent each of ethyl isocynoacetate and DBU.

Since the pyrimidine *N*-oxide **4b** was formed in lower yield than the pyrrole **4a** (see Table 2), we thought that the major product of this competitive reaction would be **4a**. This, however, was not the case, the product formed being mainly the pyrimidine *N*-oxide **4b** derived from **3b** together with a little **4a** formed from **3a**. These unexpected results suggest that the carbanion of ethyl isocynoacetate attacks the 4-position of **3b** more readily than the 5-position of **3a** and that the Michael adduct **12** from **3b** and ethyl isocynoacetate carbanion is produced with greater facility than **11** (Scheme 3).

Further, in the case of the Michael adduct **11**, subsequent cyclization occurs rapidly to give the corresponding pyrrole **4a** whilst, in contrast, the stability of the Michael adduct **12** slows its cyclization. Such a hypothesis reasonably explains the yields



Scheme 1 Reagents and conditions: i, HNO₃, H₂SO₄, 0 °C, 5 h; ii, CNCH₂CO₂Et, DBU, THF, RT, 5 h; iii, SeO₂, EtOH, AcOH, reflux, 2 h; iv, SOCl₂, Et₃N, RT, 3 h

Table 2 The nitrobenzothiadiazoles **3a,b** and selenadiazoles **5a,b** reacted with ethyl isocynoacetate in THF at room temperature for 5 h

Substrate	Product

of **4a** and **4b** in the independent reaction of **3a** and **3b** with an equimolar proportion of ethyl isocynoacetate.

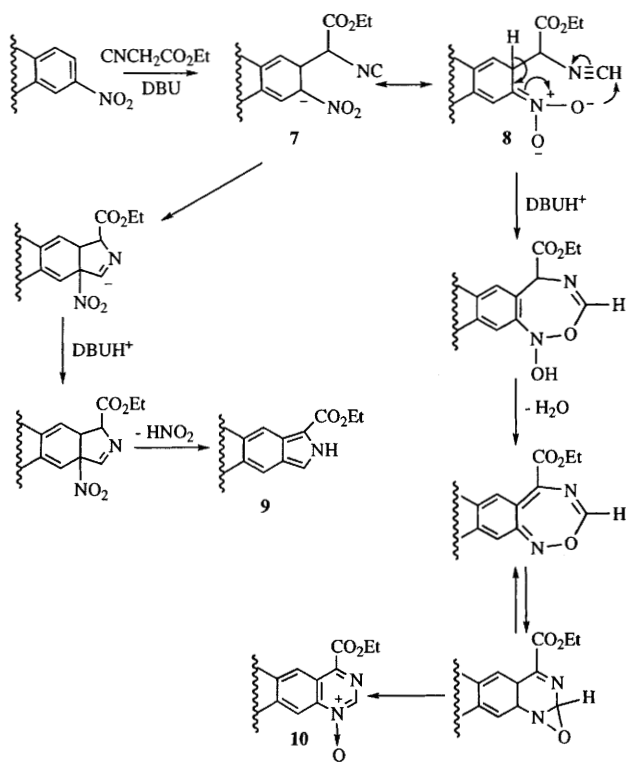
Finally, the pyrimidine *N*-oxides **4b** and **6b** were produced not from the intermediate **12** but from the resonance structure **13**, an intermediate capable of existence only when the nitro group is co-planar with the aromatic ring. Therefore, the pyrroles **4a** and **6a** were the sole products obtained from the highly hindered intermediate **11**. The last assumption is supported by a PM3 calculation.

In summary, we have described the diversity of the reactions of aromatic nitro compounds with ethyl isocynoacetate in a proposed mechanism, the detail of which and the validity of the assumption which underlie it are now under investigation.

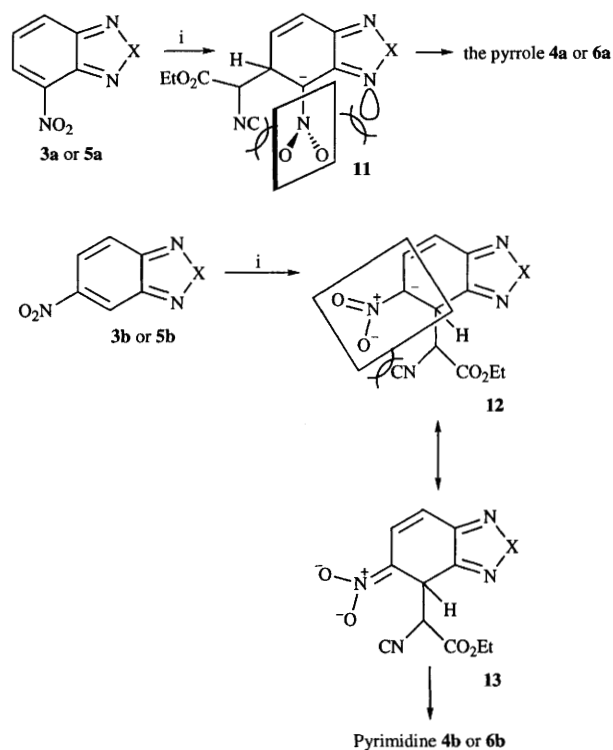
Experimental

General procedures

Melting points were measured on a Yanaco hot-stage apparatus and are uncorrected. ¹H and ¹³C NMR spectra were recorded on a JEOL-JNM-GSX 270 or JNM-EX 400 spectrometer at



Scheme 2 Proposed mechanism for the formation of pyrrole and pyrimidine *N*-oxide



Scheme 3 Reagents and conditions: i, CNCH₂CO₂Et, DBU, THF, RT, 5 h, X = S, Se

ambient temperature with CDCl₃ or [D₆]acetone as a solvent and tetramethylsilane as an internal standard; *J* values are given in Hz. IR and UV–visible spectra were obtained with a Hitachi 270–30 and Shimadzu UV-2200 spectrophotometer, respectively. Mass spectra and high-resolution mass spectra were measured with a Hitachi M80B-LC-API spectrometer under the following ionizing conditions: electron impact, 20 eV; 70 eV for HRMS; high boiling PFK as a standard. THF was freshly distilled from sodium benzophenone ketyl. Ethyl isocyan-

acetate was prepared from ethyl *N*-formylglycinate using POCl₃ and triethylamine.¹¹ Unless otherwise specified, all nitro aromatic compounds not commercially available were prepared by the conventional nitration of heteroaromatics using HNO₃–Ac₂O or HNO₃–H₂SO₄.

Ethyl pyrido[2,3-*h*]quinazoline-4-carboxylate 1-oxide 2a

DBU (1.67 g, 11 mmol) was added dropwise to a solution of 5-nitroquinoline **1a** (1.74 g, 10 mmol) and ethyl isocyanacetate (1.24 g, 11 mmol) in THF (50 cm³) at 0 °C. The resulting mixture was stirred at ambient temperature for 48 h after which it was treated with dilute hydrochloric acid (50 cm³) and extracted with ethyl acetate (3 × 20 cm³). The combined extracts were washed with aq. sodium hydrogen carbonate, water and brine, dried (Na₂SO₄) and evaporated. The residue was purified by silica gel column chromatography (ethyl acetate–hexane) to give the title compound **2a** (0.132 g, 5% yield), mp 176–179 °C; δ_H(CDCl₃) 1.57 (t, 3 H, *J* 7.02), 4.66 (q, 2 H, *J* 7.18), 7.62 (d, 1 H, *J* 8.24), 8.62–8.77 (m, 3 H), 8.88–8.92 (m, 1 H) and 9.55 (s, 1 H); δ_C(CDCl₃) 13.71 (CH₂CH₃), 62.59 (CH₂CH₃), 120.08, 120.39, 121.45, 122.63, 125.53, 127.75, 138.90, 142.33, 150.35, 154.65, 154.67 and 163.58 (C=O); ν_{max}(KBr)/cm⁻¹ 3425, 2930, 2369, 1718, 1548, 1375, 1345, 1293, 1229, 1204 and 1129; λ_{max}(CH₂Cl₂)/nm 384, 344.5, 284.6 and 240 [Found (HRMS): *M*, 269.0776. Calc. for C₁₄H₁₁N₃O₃: *M*, 269.0801].

The pyrimidine *N*-oxides **2b–f** were prepared by a similar procedure to that described for the preparation of **2a**, under the conditions described in Table 1.

Ethyl pyrazino[2,3-*f*]quinazoline-10-carboxylate 7-oxide 2b

Yield 8%; mp 194–197 °C; δ_H(CDCl₃) 1.48 (t, 3 H, *J* 7.33), 4.64 (q, 2 H, *J* 7.17), 8.83–9.12 (m, 4 H) and 9.36 (s, 1 H); δ_C(CDCl₃) 13.96 (CH₂CH₃), 62.66 (CH₂CH₃), 135.57, 135.67, 137.41, 142.00, 142.75, 144.32, 144.63, 144.71, 146.64, 147.21 and 164.81 (C=O); ν_{max}(KBr)/cm⁻¹ 1742, 1522, 1260, 1222, 1198, 1138, 1038 and 1018; λ_{max}(CH₂Cl₂)/nm 345, 285 and 251.5 [Found (HRMS): *M*, 270.0754. Calc. for C₁₃H₁₀N₄O₃: *M*, 270.0754].

Ethyl 2,3-dimethylpyrazino[2,3-*f*]quinazoline-10-carboxylate 7-oxide 2c

Yield 15%; mp 184–187 °C; δ_H(CDCl₃) 1.46 (t, 3 H, *J* 7.32), 2.81 (s, 3 H), 2.83 (s, 3 H), 4.63 (q, 2 H, *J* 7.17), 8.41 (d, 1 H, *J* 9.46), 8.77 (d, 1 H, *J* 9.46) and 9.36 (s, 1 H); δ_C(CDCl₃) 13.91 (CH₂CH₃), 22.93 (Me), 61.45 (CH₂CH₃), 135.75, 136.22, 140.64, 144.35, 144.49, 144.78, 154.21, 156.50, 161.36, 165.92 and 169.41 (C=O); ν_{max}(KBr)/cm⁻¹ 1744, 1576, 1446, 1352 and 1266; λ_{max}(CH₂Cl₂)/nm 345, 285.5 and 251 [Found (HRMS): *M*, 298.1060. Calc. for C₁₅H₁₄N₄O₃: *M*, 298.1067].

Ethyl 2,3-diphenylpyrazino[2,3-*f*]quinazoline-10-carboxylate 7-oxide 2d

Yield 32%; mp 184–187 °C; δ_H(CDCl₃) 1.07 (t, 3 H, *J* 7.02), 4.34 (q, 2 H, *J* 7.18), 7.35–7.47 (m, 5 H), 7.55–7.60 (m, 5 H), 8.56 (d, 1 H, *J* 9.47), 8.88 (d, 1 H, *J* 9.46) and 9.34 (s, 1 H); δ_C(CDCl₃) 13.56 (CH₂CH₃), 62.46 (CH₂CH₃), 120.19, 120.29, 128.21, 128.47, 129.69, 129.83, 130.26, 136.16, 136.39, 137.66, 137.86, 140.93, 142.68, 143.79, 144.70, 145.06, 153.65, 155.59 and 165.84 (C=O); ν_{max}(KBr)/cm⁻¹ 1744, 1360, 1274, 1242, 1186, 1140, 1024, 762 and 714; λ_{max}(CH₂Cl₂)/nm 413.5, 394.5, 328, 261.5 and 228.5; *m/z* 422 (*M*⁺) (Found: C, 71.0; H, 4.5; N, 13.15. C₂₅H₁₈N₄O₃ requires C, 71.08; H, 4.29; N, 13.26%).

Ethyl dibenzo[*a,c*]pyrimido[5,4-*h*]phenazine-15-carboxylate 12-oxide 2e

Yield 25%; mp 292–295 °C; δ_H(CDCl₃) 1.32 (t, 3 H, *J* 7.32), 4.75 (q, 2 H, *J* 7.17), 7.76–7.91 (m, 4 H), 8.59–8.62 (m, 3 H), 8.87 (d, 1 H, *J* 9.76), 9.22 (d, 1 H, *J* 8.24), 9.34 (d, 1 H, *J* 7.93) and 9.38 (s, 1 H); ν_{max}(KBr)/cm⁻¹ 1732, 1272, 1242, 1194 and 770; λ_{max}(CH₂Cl₂)/nm 438, 416, 326.5, 254 and 237 [Found (HRMS): *M*, 420.1227. Calc. for C₂₅H₁₆N₄O₃: *M*, 420.1223].

Ethyl thieno[2,3-*d*]pyrimidine-4-carboxylate 1-oxide 2f

Yield 40%; mp 173 °C; δ_H(CDCl₃) 1.51 (t, 3 H, *J* 7.17), 4.75 (q, 2 H, *J* 7.22), 7.77 (d, 1 H, *J* 5.80), 8.17 (d, 1 H, *J* 5.80) and 9.06 (s, 1 H); λ_{max}(CH₂Cl₂)/nm 341, 289 and 247; *m/z* 226 [*M*⁺ (³⁴S), 6%], 225 [*M*⁺ (³³S), 12], 224 [*M*⁺ (³²S), 100], 152 (96) and 127 (56).

The structure of **2f** was confirmed by treating the compound with PCl_3 and converting it into the corresponding pyrimidine **2g**; mp 79–80 °C; $\delta_{\text{H}}(\text{CDCl}_3)$ 152 (t, 3 H, J 7.0), 4.60 (q, 2 H, J 7.0), 7.76 (d, 1 H, J 2.1), 8.09 (d, 1 H, J 3.2) and 9.28 (s, 1 H) (Found: C, 51.95; H, 3.8; N, 13.6. $\text{C}_9\text{H}_8\text{N}_2\text{O}_2\text{S}$ requires: C, 51.91; H, 3.87; N, 13.45%).

5-Nitro-2,1,3-benzothiadiazole 3b. Thionyl chloride (0.87 cm^3 , 12 mmol) was slowly added at 0 °C to a solution of 5-nitro-*o*-phenylenediamine (1.53 g, 10 mmol) and triethylamine (2.28 cm^3 , 20 mmol) in DMF (15 cm^3). The resulting mixture was stirred for 3 h at room temperature after which it was diluted with water and extracted with chloroform ($2 \times 50 \text{ cm}^3$). The combined extracts were washed with aq. sodium hydrogen carbonate, water and brine, dried (Na_2SO_4) and evaporated under reduced pressure. The residue was purified by silica gel column chromatography (chloroform–hexane) to give the title compound **3b** (0.80 g, yield 44%), mp 133 °C; $\delta_{\text{H}}(\text{CDCl}_3)$ 8.17 (d, 1 H, J 9.27), 8.44 (dd, 1 H, J 9.58, 2.20) and 8.99 (d, 1 H, J 1.95); $\delta_{\text{C}}(\text{CDCl}_3)$ 118.23 (C-7), 122.42 (C-4), 122.86 (C-6), 148.46 (C-5), 153.26 and 156.30; $\nu_{\text{max}}(\text{KBr})/\text{cm}^{-1}$ 1556, 1508, 1344, 1303, 902, 842, 820 and 736; $\lambda_{\text{max}}(\text{CH}_2\text{Cl}_2)/\text{nm}$ 316 and 265; m/z 183 [M^+ (^{34}S), 9%], 182 [M^+ (^{33}S), 5%], 181 [M^+ (^{32}S), 100], 135 ($\text{M}^+ - \text{NO}_2$, 42) and 123 (48) [Found (HRMS): M , 180.9941. Calc. for $\text{C}_6\text{H}_3\text{O}_2\text{S}$: M , 180.9947].

4-Nitro-2,1,3-benzoselenadiazole 5a

Nitration of 2,1,3-benzoselenadiazole with HNO_3 – H_2SO_4 gave the nitro compound **5a** (89%), mp 227 °C; $\delta_{\text{H}}[\text{C}_6\text{H}_5\text{acetone}]$ 7.82 (dd, 1 H, J 9.00, 7.17), 8.27 (dd, 1 H, J 9.01, 1.08) and 8.45 (d, 1 H, J 7.02); $\nu_{\text{max}}(\text{KBr})/\text{cm}^{-1}$ 1526, 1506, 1432, 1320, 1292, 814 and 734; $\lambda_{\text{max}}(\text{CH}_2\text{Cl}_2)/\text{nm}$ 340; m/z 231 [M^+ (^{82}Se), 21%], 230 [M^+ (^{81}Se), 10], 229 [M^+ (^{80}Se), 100], 228 [M^+ (^{79}Se), 7], 227 [M^+ (^{78}Se), 50], 226 [M^+ (^{77}Se), 21], 225 [M^+ (^{76}Se), 21], 192 (42), 183 ($\text{M}^+ - \text{NO}_2$, 37), 171 (38) and 156 (40) [Found (HRMS): M , 228.9405. Calc. for $\text{C}_6\text{H}_3\text{N}_3\text{O}_2\text{Se}$: M , 228.9391].

5-Nitro-2,1,3-benzoselenadiazole 5b

Acetic acid (200 cm^3) was added to a stirred mixture of 5-nitro-*o*-phenylenediamine (7.65 g, 50 mmol) and selenium(IV) oxide (5.55 g, 50 mmol) in ethanol (500 cm^3). The mixture was vigorously stirred at reflux temperature for 2 h after which it was evaporated under reduced pressure. The residue was dissolved in ethyl acetate (200 cm^3) and the solution was washed with aqueous sodium hydrogen carbonate, water and brine, dried (Na_2SO_4), filtered and evaporated under reduced pressure to give the crude product. This could be used in the next reaction without further purification (yield 70%), mp 224 °C; $\delta_{\text{H}}(\text{CDCl}_3)$ 8.00 (dd, 1 H, J 9.77, 0.61), 8.30 (dd, 1 H, J 9.76, 2.14) and 8.84 (dd, 1 H, J 2.44, 0.61); $\nu_{\text{max}}(\text{KBr})/\text{cm}^{-1}$ 1542, 1500, 1338, 1272, 840, 760 and 732; $\lambda_{\text{max}}(\text{CH}_2\text{Cl}_2)/\text{nm}$ 316; m/z 231 [M^+ (^{82}Se), 19%], 230 [M^+ (^{81}Se), 8], 229 [M^+ (^{80}Se), 100], 228 [M^+ (^{79}Se), 4], 227 [M^+ (^{78}Se), 48], 226 [M^+ (^{77}Se), 18], 225 [M^+ (^{76}Se), 19], 192 (42), 183 ($\text{M}^+ - \text{NO}_2$, 37), 171 (18) and 156 (23); [Found (HRMS): M , 228.948. Calc. for $\text{C}_6\text{H}_3\text{N}_3\text{O}_2\text{Se}$: M , 228.9391].

The pyrroles **4a**, **6a** and the pyrimidine *N*-oxides **4b**, **6b** were prepared by a similar procedure to that described in the preparation of **2a**, under the conditions described in Table 2.

Ethyl pyrrolo[3,4-*e*][2,1,3]benzothiadiazole-6-carboxylate 4a. Yield 33%; mp 183 °C; $\delta_{\text{H}}(\text{CDCl}_3)$ 1.49 (t, 3 H, J 7.17), 4.43 (q, 2 H, J 7.12), 7.62 (d, 1 H, J 9.46), 7.96 (d, 1 H, J 3.36), 8.19 (d, 1 H, J 9.46) and 10.1 (s, 1 H); $\delta_{\text{C}}(\text{CDCl}_3)$ 14.39 (CH_2CH_3), 60.20 (CH_2CH_3), 115.24, 115.58, 118.29 (C-5), 119.14 (C-8), 125.00, 125.48 (C-4), 150.39, 153.92 and 160.432 (C=O); $\nu_{\text{max}}(\text{KBr})/\text{cm}^{-1}$ 3212, 1688, 1398, 1338, 1274, 1170 and 1132; $\lambda_{\text{max}}(\text{CH}_2\text{Cl}_2)/\text{nm}$ 358, 308, 295 and 267; m/z 249 [M^+ (^{34}S), 6%], 248 [M^+ (^{33}S), 14], 247 [M^+ (^{32}S), 100] and 201 (Found: C, 53.7, H, 3.7, N, 17.1. $\text{C}_{11}\text{H}_9\text{N}_3\text{O}_2\text{S}$ requires C, 53.43; H, 3.67; N, 16.99%).

Ethyl pyrrolo[3,4-*e*][2,1,3]benzothiadiazole-9-carboxylate 6a. Yield 21%; mp 181 °C; $\delta_{\text{H}}(\text{CDCl}_3)$ 1.48 (t, 3 H, J 7.08), 4.67 (q, 2 H, J 7.16), 8.37 (d, 1 H, J 9.77), 8.77 (d, 1 H, J 9.77) and 9.31 (s, 1 H); $\delta_{\text{C}}(\text{CDCl}_3)$ 13.98 (CH_2CH_3), 63.13 (CH_2CH_3), 116.20, 120.66 (C-4), 128.48 (C-5), 142.80, 145.99, 146.24 (C-7), 148.91, 154.46 and 164.79 (C=O); $\nu_{\text{max}}(\text{KBr})/\text{cm}^{-1}$ 1746, 1276, 1244, 1198, 1184, 1144, 1014 and 832; $\lambda_{\text{max}}(\text{CH}_2\text{Cl}_2)/\text{nm}$ 405, 290 and 254; m/z 278 [M^+ (^{34}S), 7%], 277 [M^+ (^{33}S), 16], 276 [M^+ (^{32}S), 100], 231 ($\text{M}^+ - \text{OEt}$, 47), 204 (79) and 188 (62) [Found (HRMS): M , 276.0314. Calc. for $\text{C}_{11}\text{H}_8\text{N}_4\text{O}_3\text{S}$: M , 276.0318] (Found: C, 47.85, H, 3.0, N, 20.3. $\text{C}_{11}\text{H}_8\text{N}_4\text{O}_3\text{S}$ requires: C, 47.82; H, 2.92; N, 20.28%).

Ethyl pyrrolo[3,4-*e*][2,1,3]benzoselenadiazole-6-carboxylate 6a. Yield 56%; mp 248 °C; $\delta_{\text{H}}(\text{CDCl}_3)$ 1.43 (t, 3 H, J 7.17), 4.43 (q, 2 H, J 7.12), 7.42 (d, 1 H, J 9.77), 8.02 (d, 1 H, J 2.74), 8.13 (d, 1 H, J 9.77) and 12.1 (s, 1 H); $\nu_{\text{max}}(\text{KBr})/\text{cm}^{-1}$ 2896, 1682, 1432, 1322, 1274, 1178 and 1142; $\lambda_{\text{max}}(\text{CH}_2\text{Cl}_2)/\text{nm}$ 378, 294, 288 and 264; m/z 297 [M^+ (^{82}Se), 22%], 296 [M^+ (^{81}Se), 14], 295 [M^+ (^{80}Se), 100], 294 [M^+ (^{79}Se), 7], 293 [M^+ (^{78}Se), 54], 292 [M^+ (^{77}Se), 20], 291 [M^+ (^{76}Se), 19] and 249 ($\text{M}^+ - \text{EtOH}$, 88) [Found (HRMS): M , 294.9853. Calc. for $\text{C}_{11}\text{H}_9\text{N}_3\text{O}_2\text{Se}$: M , 294.9861] (Found: C, 44.2, H, 3.0, N, 14.1. $\text{C}_{11}\text{H}_9\text{N}_3\text{O}_2\text{Se}$ requires C, 44.91; H, 3.08; N, 14.28%).

Ethyl pyrrolo[3,4-*e*][2,1,3]benzoselenadiazole-9-carboxylate 6a. Yield 28%; mp 248 °C; $\delta_{\text{H}}(\text{CDCl}_3)$ 1.47 (t, 3 H, J 7.08), 4.64 (q, 2 H, J 7.16), 8.20 (d, 1 H, J 9.77), 8.67 (d, 1 H, J 9.77) and 9.29 (s, 1 H); $\delta_{\text{C}}(\text{CDCl}_3)$ 14.02 (CH_2CH_3), 63.05 (CH_2CH_3), 117.66, 120.47 (C-4), 130.65 (C-5), 142.31, 146.55, 146.77 (C-7), 153.59, 158.93 and 165.24 (C=O); $\nu_{\text{max}}(\text{KBr})/\text{cm}^{-1}$ 1742, 1278, 1242, 1196, 1016, 828 and 770; $\lambda_{\text{max}}(\text{CH}_2\text{Cl}_2)/\text{nm}$ 419, 319 and 261; m/z 326 [M^+ (^{82}Se), 18%], 325 [M^+ (^{81}Se), 16], 324 [M^+ (^{80}Se), 100], 323 [M^+ (^{79}Se), 7], 322 [M^+ (^{78}Se), 49], 321 [M^+ (^{77}Se), 18], 320 [M^+ (^{76}Se), 20], 279 ($\text{M}^+ - \text{OEt}$, 40), 252 (49) and 236 (15) [Found (HRMS): M , 323.9759. Calc. for $\text{C}_{11}\text{H}_8\text{N}_4\text{O}_3\text{Se}$: M , 323.9762] (Found: C, 41.3, H, 2.7, N, 17.1. $\text{C}_{11}\text{H}_8\text{N}_4\text{O}_3\text{Se}$ requires C, 40.88; H, 2.50; N, 17.34%).

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