Supporting Information

for

Economical and scalable synthesis of 6-amino-2-cyanobenzothiazole

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Copies of \textsuperscript{1}H and \textsuperscript{13}C NMR spectra, details of calorimetry experiments and comparison of routes to ACBT 8
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1. Spectral Data

1.1 2-Chloro-6-nitro-1,3-benzothiazole (6)

**Figure S1:** $^1$H NMR spectrum of 2-chloro-6-nitro-1,3-benzothiazole 6

**Figure S2:** $^{13}$C NMR spectrum of 2-chloro-6-nitro-1,3-benzothiazole 6
1.2 6-Nitro-1,3-benzothiazole-2-carbonitrile (13)

**Figure S3:** $^1$H NMR spectrum of 6-nitro-1,3-benzothiazole-2-carbonitrile 13

**Figure S4:** $^{13}$C NMR spectrum of 6-nitro-1,3-benzothiazole-2-carbonitrile 13
1.3 6-Amino-1,3-benzothiazole-2-carbonitrile (8)

Figure S5: $^1$H NMR spectrum of 6-amino-1,3-benzothiazole-2-carbonitrile 8

Figure S6: $^{13}$C NMR spectrum of 6-amino-1,3-benzothiazole-2-carbonitrile 8
1.4 Crude mixture from cyanation reaction in ethanol

Figure S7: $^1$H NMR spectrum of a mixture of 6-nitro-1,3-benzothiazole-2-carbonitrile (13), 2-ethoxy-6-nitro-1,3-cyanobenzothiazole (14) and 6-nitro-2-benzothiazolecarboxylic acid ethyl ester (15). The assignment of the three components of the mixture was supported by LC–ESIMS analysis. The $^1$H NMR peaks for 13 match those in Figure S3, and the $^1$H NMR peaks for 14 match literature data as described in: Barlin, G.B. and Ireland, S.J. Heterocyclic Amplifiers of Phleomycin. IX* Some Derivatives of Fused and Unfused Mono- and Diaza Heterocycles. *Aust. J. Chem. 1985, 38, 1685–1691.
2. Calorimetry

2.1 Calorimetry Procedures

2.1.1 Addition of NaCN (aq) to 6-nitro-1,3-benzothiazole-2-carbonitrile (13) and DABCO in MeCN.

In a 50 mL vessel, 6-nitro-1,3-benzothiazole-2-carbonitrile (13, 200 mg, 0.932 mmol) and DABCO (15.7 mg, 0.14 mmol) were dissolved in MeCN (20.0 mL). Aqueous NaCN (0.49 M) was then fed into the reaction as described below, and power compensation calorimetry was used to determine heat changes.

**Steady State 1:**

| Stir  | 300 rpm |
| Reactor Temperature | 21 °C |
| Circulator Temperature | 1 °C |
| Time | 60 min |

**Feed 1:**

| Stir  | 300 rpm |
| Reactor Temperature | 21 °C |
| Circulator Temperature | 1 °C |
| Feed | 0.49 M NaCN (aq) at 0.2 mL min⁻¹ for 10 min (2.00 mL, 0.978 mmol NaCN) |
| Time | 10 min |

**Steady State 2:**

| Stir  | 300 rpm |
| Reactor Temperature | 21 °C |
| Circulator Temperature | 1 °C |
| Time | 1080 min |

**Feed 2:**

| Stir  | 300 rpm |
| Reactor Temperature | 21 °C |
| Circulator Temperature | 1 °C |
| Feed | 0.003 M FeCl₃ (aq) at 2 mL min⁻¹ for 10 min (20.0 mL, 0.060 mmol FeCl₃) |
| Time | 10 min |
Steady State 3:
Stir 300 rpm
Reactor Temperature 21 °C
Circulator Temperature 1 °C
Time 60 min

The IQ software requires input of the specific heat capacity of the feed for the calculation of compensatory power. For this, the specific heat capacity of water (4.18 J g\(^{-1}\) K\(^{-1}\)) was used.
2.1.2 Addition of water to 6-nitro-1,3-benzothiazole-2-carbonitrile (13) and DABCO in MeCN

In a 50 mL vessel, 6-nitro-1,3-benzothiazole-2-carbonitrile (13, 200 mg, 0.932 mmol) and DABCO (15.7 mg, 0.14 mmol) were dissolved in MeCN (20.0 mL). Water (2.0 mL) was then fed into the reaction as described below, and power compensation calorimetry was used to determine heat changes.

**Steady State 1:**

- **Stir** 300 rpm
- **Reactor Temperature** 21 °C
- **Circulator Temperature** 1 °C
- **Time** 90 min

**Feed 1:**

- **Stir** 300 rpm
- **Reactor Temperature** 21 °C
- **Circulator Temperature** 1 °C
- **Feed** Water at 0.2 mL min⁻¹ for 10 min (2.00 mL)
- **Time** 10 min

**Steady State 2:**

- **Stir** 300 rpm
- **Reactor Temperature** 21 °C
- **Circulator Temperature** 1 °C
- **Time** 50 min

The software requires input of the specific heat capacity of the feed for the calculation of compensatory power. For this, the specific heat capacity of water (4.18 J g⁻¹ K⁻¹) was used.
2.2 Calorimetric Data

2.2.1 Addition of NaCN (aq) to 6-nitro-1,3-benzothiazole-2-carbonitrile (13) and DABCO in MeCN.

The raw calorimetry data show an increase in the power demand of the internal heater upon addition of the aqueous NaCN solution. This indicates an endothermic process in the reactor requiring greater heating power from the heater to maintain the set reactor temperature of 21 °C.

The IQ software integrates the power output (W) during the reaction to give the total energy ($Q_{Total}$ in kJ) using the equation:

$$Q_{Total} = Q_{Comp} - Q_{Dose}$$

$$Q_{Dose} = m \times C_p \times (T_f - T_i)$$

Where
- $Q_{Comp} =$ Heat compensated for by internal heater i.e. the inverse of the heat produced by the reaction
- $Q_{Dose} =$ Heat added due to feed being at a different temperature to the reaction temperature
- $m =$ total mass of feed (g s$^{-1}$)
- $C_p =$ specific heat capacity of feed solution (J g$^{-1}$ °C$^{-1}$)
- $T_i =$ final temperature (°C)
$T_i =$ initial temperature (°C)

It should be noted that the calculated energies given relate to the output of the heater coil rather than the reaction, i.e. a negative value indicates release of energy from the heater coil and consumption of that energy by the reaction (endothermic reaction). The signs of the energy is therefore inverted to give the true value for the reaction.

There is a very short delay upon the start of the feed before the endotherm is observed. Following the end of the feed there is a further spike, which is due to manual feeding of the remaining aqueous NaCN solution from the syringe. According to these data, 0.37 kJ is consumed during the feed and 0.05 kJ is consumed upon addition of the remaining solution from the syringe, giving a total energy of 0.42 kJ.
2.2.2 Addition of water to 6-nitro-1,3-benzothiazole-2-carbonitrile 13 and DABCO in MeCN.

The raw calorimetry data show an increase in the power demand of the internal heater upon addition of water. This indicates an endothermic process in the reactor requiring greater heating power from the heater to maintain the set reactor temperature of 21 °C.
It should be noted that the calculated energies and molar heat given relate to the output of the heater coil rather than the reaction, i.e. a negative value indicates release of energy from the heater coil and consumption of that energy by the reaction (endothermic reaction). The signs of the energy and molar heat values are therefore inverted to give the true value for the reaction.

There is a short delay upon the start of the feed before the endotherm is observed. Following the end of the feed there is a further spike, which is due to manual feeding of the remaining water in the syringe. According to these data, 0.49 kJ is consumed during the feed and 0.02 kJ is consumed upon addition of the remaining water from the syringe, giving a total energy of 0.51 kJ.

The calculated molar heat in this experiment ($\Delta H = +4.57$ kJ mol$^{-1}$) is the energy per mole water added.
2.2.3 Calculations

Molar heat calculation for addition of NaCN to the reaction mixture:

\[ E_{(NaCN)} = E_{(NaCN+water)} - E_{(water)} = 0.42 - 0.51 = -0.09 \text{ kJ} \]

\[ \Delta H_{(NaCN)} = \frac{E_{(NaCN)}}{n_{(NaCN)}} = \frac{-0.09}{0.001} = -90 \text{ kJ mol}^{-1} \]

Where:

- \( E_{(NaCN)} \): Total energy released from addition of NaCN (calculated)
- \( E_{(NaCN+water)} \): Total energy released from addition of NaCN in water (0.5 M) (measured)
- \( E_{(water)} \): Total energy released from addition of water (measured)
- \( \Delta H_{(NaCN)} \): Heat of reaction for the addition of 1 mole of NaCN (calculated)
- \( n_{(NaCN)} \): Moles of NaCN added to reaction (measured)

It is important to note that this calculation of molar heat of reaction is an approximation, as the properties (specific heat capacity, ionic strength) of the feeds in the reaction and control experiment are different.
3. Comparison of routes to ACBT 8

A: Takakura et al. 2011 and Wang et al. 2013

\[
\text{S} \quad \text{16} \xrightarrow{\text{KNO}_3, \ H_2SO_4, \ rt, \ 18 \ h} \text{O}_2\text{N} \quad \text{N} \quad \text{S} \quad \text{Cl} \quad \text{6} \xrightarrow{\text{SnCl}_2 \cdot 2\text{H}_2\text{O}, \ \text{HCl (aq), EtOH, reflux}} \text{N} \quad \text{S} \quad \text{Cl} \quad \text{7} \xrightarrow{\text{KCN, DMSO, } 120^\circ C} \text{N} \quad \text{S} \quad \text{Cl} \quad \text{8, ACBT}
\]

this work

mp 193-194 °C (EtOH)

mp 58-61%

B: McCutcheon et al. 2015

\[
\text{O}_2\text{N} \quad \text{N} \quad \text{S} \quad \text{Cl} \quad \text{9} \xrightarrow{\text{NH}_2\text{NH}_2\text{H}_2\text{O, EtOH}} \text{O}_2\text{N} \quad \text{N} \quad \text{S} \quad \text{Cl} \quad \text{10}
\]

\[
\text{Cl} \quad \text{N} \quad \text{S} \quad \text{S} \quad \text{Cl} \quad \text{11} \xrightarrow{\text{S}_2\text{Cl}_2, \ \text{MeCN, THF, 80%}} \text{Cl} \quad \text{N} \quad \text{S} \quad \text{S} \quad \text{Cl} \quad \text{12, Appel's salt}
\]

\[
\text{O}_2\text{N} \quad \text{N} \quad \text{S} \quad \text{Cl} \quad \text{13} \xrightarrow{\text{Zn, NH}_2\text{Cl, MeOH, reflux, 62%}} \text{O}_2\text{N} \quad \text{N} \quad \text{S} \quad \text{Cl} \quad \text{8, ACBT, 95%}
\]

C: This work

\[
\text{S} \quad \text{N} \quad \text{Cl} \quad \text{16} \xrightarrow{\text{KNO}_3, \ H_2SO_4, \ rt, \ 18 \ h} \text{O}_2\text{N} \quad \text{N} \quad \text{S} \quad \text{Cl} \quad \text{6} \xrightarrow{\text{DABCO (15 mol%), NaCN (1.05 equiv.), MeCN: H}_2\text{O (10:1), rt, 24 h}} \text{O}_2\text{N} \quad \text{N} \quad \text{S} \quad \text{Cl} \quad \text{13} \xrightarrow{\text{Fe, AcOH, rt, 24 h, 60-71%}} \text{O}_2\text{N} \quad \text{N} \quad \text{S} \quad \text{Cl} \quad \text{8, ACBT, mp 219-220 °C (EtOH)}
\]

mp 193-194 °C (EtOH)

mp 83-93%

mp 164-165 °C (MeOH)
Table S1: Comparison of routes to ACBT

<table>
<thead>
<tr>
<th>Route</th>
<th>Starting material(s)</th>
<th>Number of steps</th>
<th>Purification steps</th>
<th>Reported overall yield</th>
<th>Estimated costs of reagents to produce 10g of ACBT</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>6</td>
<td>2</td>
<td>6 → 7: extraction and column 7 → 8: (extensive) extraction and column</td>
<td>15–33%</td>
<td>6: £751–£1706  SnCl₂·H₂O: £37–£85  Total: £788–£1791</td>
<td>Dilute conditions of cyanation of 7 cause significant problems during work-up. This, together with the danger of using KCN in DMSO at high temperature, make this reaction unsuitable for scale-up beyond ca. 1g scale (leading to only ca. 500 mg of 8) in a typical research lab.</td>
</tr>
<tr>
<td>B</td>
<td>16b</td>
<td>3</td>
<td>16 → 6: precipitation and crystallisation 6 → 7: extraction and column 7 → 8: (extensive) extraction and column</td>
<td>6–27%</td>
<td>16: £30–£80  KNO₃: &lt;£1  SnCl₂·H₂O: £36–£85  Total: £67–£166</td>
<td>In our attempts, reduction of 13 with zinc powder partially stalled at the corresponding nitroso intermediate; the source/particle size of zinc powder may be critical.</td>
</tr>
<tr>
<td>B</td>
<td>9 and 11</td>
<td>4</td>
<td>9 → 10: precipitation 11 → 12: precipitation 10 + 12 → 13: column 13 → 8: filtration and column</td>
<td>47%</td>
<td>11: £1  S₂Cl₂: £1  9: £69  Hydrazine: £12  Zinc powder: £5  Total: £88</td>
<td>For the DABCO-catalysed cyanation of 6, the use of MeCN as solvent, slow addition of aqueous NaCN (to prevent an exotherm), and quench of unreacted cyanide with aqueous FeCl₃ make this step more appropriate for scale-up. In our hands, the reduction of 13 with iron powder was more straight-forward than reduction with zinc powder (as described by McCutcheon et al.).</td>
</tr>
</tbody>
</table>

*Costs of starting materials, reagents and catalysts were estimated based on the best catalogue prices listed on ScienceWarehouse on 3 August 2016; costs of bulk chemicals (sulfuric acid, HCl (aq), FeCl₃, KCN, NaCN), solvents and consumables required for purifications have not been included. **Calculations for 16 → 6 were based on reaction/purification conditions and yield described in this work.