The acid-catalysed synthesis of 7-azaindoles from 3-alkynyl-2-aminopyridines and their antimicrobial activity†

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The synthesis of 7-azaindoles from 3-alkynyl-2-aminopyridines using acidic conditions, namely, a mixture of trifluoroacetic acid (TFA) and trifluoroacetic anhydride (TFAA), is described. This methodology resulted in the synthesis of fifteen 7-azaindoles, with most containing substituents at the 2- and 5-positions. The majority of these were tested for antimicrobial activity against a range of bacteria and yeasts. The 7-azaindoles displayed the best activity against the yeasts, particularly against Cryptococcus neoformans, where activities as low as 3.9 µg ml⁻¹ were observed.

Introduction

The azaindoles are a growing class of indole isosteres that possess promising biological profiles. For example, the natural product variolin B 1 (Fig. 1) isolated from an extremely rare Antarctic sponge is a promising anti-cancer agent.¹

Other examples include the synthetic azaindole derivative of rebecamycin, known as diazarebecamycin 2, which has also been shown to exhibit promising anticancer activity,² and 6-azaindole 3 which has demonstrated activity as an inhibitor of HIV-1 attachment to CD4+ T cells.³

The synthesis of 7-azaindoles is often accomplished by initial Sonogashira coupling of a 3-halosubstituted-2-aminopyridine as shown in Scheme 1.⁴ The intermediate alkynylpyridines are then exposed to a variety of different reaction conditions and reagents to afford the desired azaindoles. Many methods have been developed for this step and include the use of cesium- or potassium-containing bases such as potassium hydride, potassium t-butoxide or cesium t-butoxide⁵a–d or even bases such as DBU⁵e and triethylamine.⁵f Metal-containing catalysts have also been used for this purpose and include amongst others, the use of copper,⁶a an indium salt⁶b or even gold catalysts.⁶c The Cacchi reaction in the presence of a palladium catalyst⁷ has also been successfully used to facilitate both the Sonogashira coupling reaction and the azaindole

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ring forming step to produce 2,3-disubstituted azaindoles directly from the aminopyridines. Even the use of microwaves in the presence of water with\textsuperscript{7a} or without\textsuperscript{7b} added salts has been reported.

As a result of our interest in the synthesis of indole-containing compounds and their antibacterial and antifungal activity,\textsuperscript{8} we initiated a programme directed towards the synthesis of the related compounds, azaindoles. The specific goal described in this publication was to attempt to develop versatile and cheap conditions for the synthesis of 7-azaindoles from the corresponding Sonogashira coupled 3-alkynyl-2-aminopyridine precursors. Thereafter, as azaindoles have interesting biological profiles, we wished to ascertain if the 7-azaindoles we had synthesized would display activity against a range of bacterial and yeast cell lines.

**Results and discussion**

Initially the required halogenated substituted pyridines were prepared or obtained from commercial sources. 5-Chloro-2-aminoypyridine\textsuperscript{5} and 5-bromo-3-iodopyridin-2-amine\textsuperscript{9} were obtained from Sigma-Aldrich. 5-Chloro-2-aminopyridine\textsuperscript{5} was treated with iodine in the presence of Ag\textsubscript{2}SO\textsubscript{4} in ethanol to afford the 3-iodo substituted aminopyridine\textsuperscript{8} in good yield.

Finally, 5-bromo-3-iodopyridin-2-amine\textsuperscript{10} was prepared by initially treating 2-aminopyridine\textsuperscript{5} with NBS\textsuperscript{10} in MeCN to afford 5-bromo-2-aminopyridine\textsuperscript{9}. Then 9 was subjected to potassium iodate, I\textsubscript{2} and a mixture of H\textsubscript{2}SO\textsubscript{4}, acetic acid and water\textsuperscript{a} to provide 5-bromo-3-iodopyridin-2-amine\textsuperscript{10} in good yield (Scheme 2).

We were now in a position to attempt the Sonogashira coupling reaction, for which we believed that coupling would take place at the more reactive iodine substituent in substrates\textsuperscript{8} and\textsuperscript{10}, or at the bromine in the case of aminopyridines\textsuperscript{6} and\textsuperscript{7}. The 2-aminopyridine containing pyridines were treated with catalytic tetrakis(triphenylphosphine)palladium(0), copper(i) iodide, triethylamine as a base and a range of acetylenes to afford the desired compounds (11–14) in good to excellent yield as shown in Scheme 3.

The stage was now set to attempt the formation of the pyrrole ring in order to assemble the 7-azaindole nucleus. For this step we wished to investigate if it was possible to use acidic conditions to affect the ring closure. A variety of acids (HCl, H\textsubscript{2}SO\textsubscript{4}, AcOH and trifluoroacetic acid (TFA)) were initially used. The use of TFA seemed to be promising as a small amount of azaindole was forming but the conversion rate was estimated to be only 10%. Since there are examples in the literature\textsuperscript{4} of the corresponding amide (rather than amine) undergoing ring closure to afford azaindoles, we thought that the addition of trifluoroacetic anhydride (TFAA) to the reaction mixture containing TFA might improve the rate, conversion and yield of the desired azaindoles. After much experimentation we found that in the presence of 1 equivalents of TFA and 1.3 equivalents of TFAA in MeCN as a solvent and heating to reflux for 8 h, the substrate afforded the desired azaindoles (15–18) in generally excellent yields (Scheme 4).

As one final example we wished to take advantage of the different reactivity of the halogens on substrate\textsuperscript{10} to establish if it was possible to introduce two different acetylene substituents using different temperatures. Hence as shown in Scheme 5,\textsuperscript{10} 10 was treated under Sonogashira reaction conditions with phenylacetylene initially at room temperature. After TLC showed that the starting material had all been consumed, the solvent was evaporated under reduced pressure and the crude material was then subjected to the same palladium catalyst using trimethylsilylacetylene as the acetylene coupling partner. After the reaction was deemed to be complete by TLC, work-up of the reaction and analysis of the product by NMR spectroscopy showed that two different acetylene substituents were present on

**Scheme 2**  **Reagents and conditions:**  (i) I\textsubscript{2}, Ag\textsubscript{2}SO\textsubscript{4}, EtOH, rt, 86%; (ii) NBS, MeCN, rt, 98%; (iii) I\textsubscript{2}, KIO\textsubscript{3}, H\textsubscript{2}SO\textsubscript{4}, AcOH, H\textsubscript{2}O, 80 °C, 90%.

**Scheme 3**  **Reagents and conditions:**  (i) RC\textequiv CH, Et\textsubscript{3}N, Cul, cat. Pd(PPh\textsubscript{3})\textsubscript{4}, THF, 70 °C; (ii) RC\equiv CH Et\textsubscript{3}N, Cul, cat. Pd(PPh\textsubscript{3})\textsubscript{4}, THF, rt.

**Scheme 4**
the 3,5-substituted aminopyridine 19. Presumably, this is due to the different reactivities of the iodine and bromine substituents in the two Sonogashira reactions. Exposure of 19 to our acid mediated reaction conditions for azaindole formation resulted in the formation of the desired azaindole 20 in excellent yield.

As we now had a series of substituted azaindoles on hand they were quantitatively evaluated for antimicrobial activity (Table 1). While compound 16b showed promising antibacterial activity (7.8 µg ml⁻¹) against *Pseudomonas aeruginosa*, many of the azaindoles showed very good and selective anti-yeast activity, particularly against *Cryptococcus neoformans*, where all compounds showed noteworthy inhibitory values of between 3.9–15.6 µg ml⁻¹. Previous studies on the antimicrobial effects of other azaindole derivatives include the quantitative study by Kumaran *et al.*,¹¹ and the promising antifungal effects found by Lebouvier.¹² Previous antimicrobial studies on azaindoles for activity against *Cryptococcus neoformans* is, however, lacking and the promising activity noted here warrants further examination, particularly, given that increased reports of infections from *Cryptococcus neoformans* show that this represents a major health threat for HIV infected patients.¹³,¹⁴ Newer antifungals to combat this opportunistic pathogen will certainly be beneficial in future treatments, particularly with emerging trends of resistance evident.

In summary, to the best of our knowledge this is the first report outlining the synthesis of azaindoles from 3-alkynyl-2-aminopyridines under acidic conditions. The azaindoles synthesized display significant anti-yeast activity.

**Experimental**

¹H NMR and ¹³C NMR spectra were recorded either on a Bruker AVANCE 300 spectrometer or a Bruker AVANCE III 500 spectrometer. All chemical shift values are reported in parts per million referenced against TMS which is given an assignment of zero parts per million. Coupling constants (J-values) are given in hertz (Hz). All mass spectroscopy data were collected on a Waters Acquity UPLC system coupled to a Waters HDMS G1 QTOF mass spectrometer. UPLC settings: analytical column: BEH C18 150 × 2.1 mm; column temperature: 60 °C; mobile phase: 90% water (0.1% formic acid): 5% acetonitrile; flow rate: 0.4 ml min⁻¹. MS settings: mode: VTOF; ionisation: ESIPOS and ESINeg; scan range: 100–1000 Da; scan speed: 0.1 second; run time: 10 min. Infrared spectra were recorded on a Bruker Tensor 27 standard system spectrometer with diamond ATR attachment. Macherey-Nagel Kieselgel 60 (particle size 0.063–0.200 mm) was used for conventional silica gel column chromatography with various EtOAc and hexane mixtures as the mobile phase. TLC was performed on aluminum-backed Macherey-Nagel Alugram Sil G/UV254 plates pre-coated with 0.25 mm silica gel 60. The following acetylenes are commercially available: phenylacetylene, trimethylsilylacetylene, 2-ethynylanisole, 4-ethynylanisole, 1-ethynyl-4-methoxy-2-methylbenzene, 1-ethynyl-3,5-bis(trifluoromethyl)benzene and propargylbenzoate. The remaining two acetylenes 1-(prop-2-ynyl)piperidine and 4-(prop-2-ynyl)morpholine were prepared according to ref. 15 and 16.

5-Chloro-3-iodo-2-aminopyridine 8

To a solution of 2-amino-5-chloropyridine 4 (0.700 g, 5.45 mmol) in ethanol was added silver sulfate (1.699 g,
Table 1  Antimicrobial and antifungal activity of selected azaindoles (µg ml⁻¹)

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⁵ Control for bacteria, ciprofloxacin; for the yeasts, amphotericin.

5.45 mmol) in one portion followed by the addition of iodine (1.383 g, 5.45 mmol) portionwise. The heterogeneous mixture was stirred at rt until determined complete by TLC. After the reaction was complete, the mixture was filtered through celite or silica gel and the ethanol removed under reduced pressure. The resulting crude material was re-dissolved in dichloromethane and washed with a saturated solution of aqueous sodium thiosulfate. The separated dichloromethane layer was removed under reduced pressure and the resulting solid was purified by silica gel chromatography to give 2-amino-5-bromopyridine 10 as a white solid (70.11 g, 90%). ¹H NMR (300 MHz, CDCl₃) δ 8.05 (s, 1H), 7.95 (s, 1H), 5.03 (s, 2H); ¹³C NMR (75 MHz, CDCl₃) δ 156.75, 148.72, 148.57, 107.58, 77.93.

General procedure for the Sonogashira reaction to synthesize 11a–c and 12a–d

To a flame-dried round-bottom flask under a nitrogen or argon atmosphere containing 2-amino-3-bromopyridine derivative [6 and 7] was added CuI (2 mol%) and Pd(PPh₃)$_2$ (2 mol%) in one portion followed by the addition of the degassed alkyne solution in THF (20 ml). Triethylamine (5 ml) was then added and the reaction mixture was stirred at 70 °C until no starting material was present as monitored by thin layer chromatography (TLC). After this time, the reaction was quenched with a saturated aqueous ammonium chloride and was then extracted with either dichloromethane or ethyl acetate. The combined organic extracts were dried over magnesium sulfate, filtered through either silica or celite and the excess solvent was removed on a rotary evaporator, followed by purification using flash chromatography (30% EtOAc–hexane) to yield the desired products 11a–c and 12a–d.

3-[(2-Methoxyphenyl)ethynyl]pyridin-2-amine 11a. Using 2-amino-3-bromopyridine (0.450 g, 2.60 mmol), 1-ethyl-2-methoxybenzene (0.343 g, 2.60 mmol), CuI (9.91 mg, 0.0276 mmol) and Pd(PPh₃)$_2$ (0.0212 g, 0.0520 mmol) the product 11a was obtained as a cream white solid (0.351 g, 75%). Mp: 118–120 °C; ¹H NMR (500 MHz, CDCl₃) δ 8.03 (d, J = 5.0, 1.6, 1H), 7.57 (dt, J = 13.5, 6.7, 1H), 7.46 (dd, J = 7.6, 1.6, 1H), 7.32 (td, J = 8.4, 1.7, 1H), 6.98–6.89 (m, 2H), 6.62 (dd, J = 7.5, 5.0, 1H), 5.34 (s, 2H), 3.92 (s, 3H); ¹³C NMR (126 MHz, CDCl₃) δ 159.92, 159.17, 147.83, 138.83, 132.44, 129.95, 120.65, 113.27, 112.09, 110.53, 103.46, 92.26, 89.30, 55.82. IR (cm⁻¹) 3298 (NH str.), 2997 (CH str.), 2312 (alkyne), 1619 (C=N), 1580
Using 2-amino-3-bromopyridine (0.450 g, 2.41 mmol), 1-ethyl-5-methyl-3-(phenylethynyl)pyridin-2-amine 12a. Starting with 2-amino-3-bromo-5-methylpyridine (0.450 g, 2.41 mmol), phenylacetylene (0.2455 g, 2.41 mmol), CuI (9.18 mg, 0.0482 mmol) and Pd(PPh₃)₄ (0.0557 g, 0.0482 mmol) the product 12a(0.450 g, 90%) was obtained as a cream white solid. ¹H NMR (300 MHz, CDCl₃) δ 7.90 (d, J = 1.6, 1H), 7.57–7.49 (m, 2H), 7.47 (d, J = 1.9, 1H), 7.38 (dd, J = 6.5, 2.7, 3H), 4.97 (s, 2H), 2.21 (s, 3H); ¹³C NMR (75 MHz, CDCl₃) δ 156.86, 147.82, 140.62, 131.47, 128.57, 128.45, 122.77, 122.55, 102.87, 95.30, 84.58, 17.27.

3-[2-(Methoxymethyl)-5-methyl]pyridin-2-amine 12b. Using 2-amino-3-bromo-5-methylpyridine (0.450 g, 2.41 mmol), 1-ethyl-2-methoxybenzene (0.3177 g, 2.41 mmol), CuI (9.18 mg, 0.0482 mmol) and Pd(PPh₃)₄ (0.0557 g, 0.0482 mmol), the product 3-(2-(methoxymethyl)-5-methyl)pyridin-2-amine 12b (0.404 g, 70%) was obtained as a cream white solid. Mp: 120–123 °C; ¹H NMR (500 MHz, CDCl₃) δ 7.86 (s, 1H), 7.49–7.39 (m, 2H), 7.31 (d, J = 1.7, 1H), 6.94 (dddd, J = 22.1, 11.0, 4.6, 2H), 5.16 (s, 2H), 3.92 (s, 3H), 2.18 (s, 3H); ¹³C NMR (126 MHz, CDCl₃) δ 147.13, 135.88, 132.49, 129.87, 122.17, 120.62, 112.19, 110.53, 103.13, 92.05, 89.43, 55.80, 17.29; IR (cm⁻¹) 3301 (NH str.), 2961 (CH str.), 1639 (C=O). HRMS (ES⁺) Calculated for C₁₅H₁₂N₂O [M + H⁺]: 239.0953, found 239.0949.

3-[4-(Methoxymethyl)ethyl]-5-methyl]pyridin-2-amine 12c. Using 2-amino-3-bromo-5-methylpyridine (0.450 g, 2.41 mmol), 1-ethyl-4-methoxymethylbenzene (0.3177 g, 2.41 mmol), CuI (9.18 mg, 0.0482 mmol) and Pd(PPh₃)₄ (0.0557 g, 0.0482 mmol), the desired product 12c (0.472 g, 82%) was obtained as a cream white solid. Mp: 94–96 °C; ¹H NMR (500 MHz, CDCl₃) δ 7.86 (d, J = 1.6, 1H), 7.46–7.43 (m, 2H), 7.42 (d, J = 1.9, 1H), 6.91–6.85 (m, 2H), 4.92 (s, 2H), 3.82 (s, 3H), 2.18 (s, 3H); ¹³C NMR (126 MHz, CDCl₃) δ 159.87, 156.86, 147.56, 140.38, 132.97, 122.51, 114.90, 114.12, 103.24, 95.30, 83.30, 55.33, 17.26; IR (cm⁻¹) 3276 (NH str.), 2978 (CH str.), 2238 (alkyne), 1618 (C=O), 1564 (C=C), 1170 (C=O); HRMS (ES⁺) Calculated for C₁₅H₁₄N₂O [M + H⁺]: 239.1184, found 239.1185.

3-[3-(Chlorophenyl)ethyl] 5-methylpyridin-2-amine 12d. Using 2-amino-3-bromo-5-methylpyridine (0.450 g, 2.41 mmol), 1-ethyl-3-chlorobenzene (0.328 g, 2.41 mmol), CuI (9.18 mg, 0.0482 mmol) and Pd(PPh₃)₄ (0.0557 g, 0.0482 mmol), the product 12d (0.482 g, 83%) was obtained as a cream white solid. Mp: 104–106 °C; ¹H NMR (500 MHz, CDCl₃) δ 7.89 (s, 1H), 7.50 (s, 1H), 7.45 (s, 1H), 7.39 (d, J = 7.3, 1H), 7.31 (dt, J = 13.3, 6.8, 2H), 4.93 (s, 2H), 2.20 (s, 3H); ¹³C NMR (126 MHz, CDCl₃) δ 156.84, 148.16, 140.86, 134.35, 131.31, 129.70, 129.58, 128.84, 124.49, 122.68, 102.36, 93.87, 85.78, 17.25; IR (cm⁻¹) 3347 (NH str.), 3005 (CH str.), 2259 (alkyne), 1624 (C=O), 1390 (C=N), 1176 (C=O); HRMS (ES⁺) Calculated for C₁₅H₁₄N₂Cl [M + H⁺]: 243.0689, found 243.0705.

General procedure for the Sonogashira reaction to synthesize 13a-b and 14a-e.

To a flame-dried round-bottom flask under a nitrogen or argon atmosphere containing 2-amino-3-iodopyridine derivative 9 and 10 was added CuI (2 mol%) and Pd(PPh₃)₄ (2 mol%) in one portion followed by the addition of the degassed alkene solution in THF (20 ml). Triethylamine (5 ml) was then added and the reaction mixture was stirred at rt, until no starting material was present as monitored by thin layer chromatography (TLC). After this time, the reaction was quenched with a saturated aqueous ammonium chloride and was extracted with either dichloromethane or ethyl acetate. The combined organic extracts were dried over magnesium sulfate, filtered through either silica or celite and the excess solvent was removed on a rotary evaporator, followed by purification using flash chromatography (30% EtOAc–hexane) to yield the desired products 12a-b and 14a–e.

5-Chloro-3-[trimethylsilyl]ethyl]pyridin-2-amine 13a. Using 2-amino-3-iodo-5-chloropyridine (4.475 g, 17.6 mmol), trimethylsilylacetylene (8.626 g, 88.1 mmol), CuI (0.06704 g, 0.352 mmol) and Pd(PPh₃)₄ (0.4068 g, 0.352 mmol) 13a (3.626 g, 92%) was obtained as a cream white solid; ¹H NMR (300 MHz, CDCl₃) δ 7.96 (d, J = 2.4, 1H), 7.51 (d, J = 2.5, 1H), 5.06 (s, 2H), 0.27 (s, 9H).

5-Chloro-3-[4-methoxy-2-methyl]phenyl]pyridin-2-amine 13b. Using 2-amino-3-iodo-5-chloropyridine (0.300 g, 1.19 mmol), 1-ethyl-4-methoxy-2-methylbenzene (0.1740 g, 1.19 mmol), CuI (4.517 mg, 0.0237 mmol) and Pd(PPh₃)₄ (0.0468 g, 0.0237 mmol) the product 13b was produced (0.2834 g, 87%) as a cream white solid. Mp: 127–130 °C;
1H NMR (500 MHz, CDCl3) δ 7.97 (s, 1H), 7.55 (s, 1H), 7.41 (d, J = 8.4, 1H), 6.79 (s, 1H), 6.74 (d, J = 8.4, 1H), 5.04 (s, 2H), 3.83 (s, 3H), 2.48 (s, 3H); 13C NMR (126 MHz, CDCl3) δ 160.17, 156.92, 145.88, 141.88, 138.66, 133.38, 138.18, 115.32, 114.26, 111.56, 105.04, 95.67, 85.77, 55.31, 21.18; IR (cm⁻¹) 3368 (NH) 2989 (CH str.), 2287 (alkyne), 1605 (C=N), 1558 (C=C), 1164 (C-O); HRMS (ES⁺) Calculated for C15H14N2ClO [M + H⁺]: 273.0795, found 273.0801.

3-(2-Amino-5-bromopyridin-3-yl)prop-2-yn-1-yl benzoate 14a. Using 2-amino-3-iodo-5-bromopyridine (2.500 g, 8.36 mmol), 1-ethyl-2-methoxybenzene (1.105 g, 8.36 mmol), CuI (31.84 mg, 0.167 mmol), Pd(PPh₃)₄ (0.1932 g, 0.167 mmol) furnished the product 14a as a cream white solid (2.013 g, 83%). Mp: 126–128 °C; 1H NMR (300 MHz, CDCl3) δ 8.06 (d, J = 2.3, 1H), 7.67 (d, J = 2.3, 1H), 7.46 (dd, J = 7.5, 1.5, 1H), 7.40–7.29 (m, 1H), 7.00–6.91 (m, 2H), 5.31 (s, 2H), 3.92 (s, 3H); 13C NMR (75 MHz, CDCl3) δ 160.00, 157.82, 148.21, 140.39, 132.47, 130.34, 120.70, 111.56, 110.55, 106.69, 105.26, 93.51, 88.04, 55.8; IR (cm⁻¹) 3310 (NH str.), 2989 (CH str.), 2312 (alkyne), 1627 (C=N), 1571 (C=C), 1181 (C=O); HRMS (ES⁺) Calculated for C14H12N2BrO [M + H⁺]: 303.0133, found 303.0147.

5-Bromo-3-([2-methoxyphenyl]ethyl)pyridin-2-amine 14e. The corresponding 2-amino-3-ethynylpyridine derivatives (40–200 mg) 11a-c, 12a-d, 13a-b and 14a-e were dissolved in MeCN. To the resulting solution was added trifluoroacetic acid (1 eq.) in one portion followed by the addition of trifluoroacetic anhydride (1.3 eq.) dropwise. The resulting mixture was then heated to reflux for 8 h and was allowed to cool to rt. The solvent was then removed on a rotary evaporator and the resulting crude material was re-dissolved in dichloromethane and washed with aqueous sodium carbonate (10%). The separated and dried (MgSO₄) organic phase was purified by flash silica chromatography using ethyl acetate–hexane mixture (10:50%) gave desired azaindoles 15a-c, 16a-d, 17a-d and 18a-e in good yields.

2-(2-Methoxyphenyl)pyrrolo[2,3-b]pyridine 15a. The product 15a was produced as a cream white solid (0.1236 g, 92%). Mp: 125–128 °C; 1H NMR (500 MHz, CDCl3) δ 10.47 (s, 1H), 8.30–8.20 (m, 1H), 7.93–7.87 (m, 1H), 7.84 (dd, J = 7.7, 1.5, 1H), 7.36–7.27 (m, 1H), 7.12–6.98 (m, 3H), 6.84 (d, J = 1.6, 1H), 3.98 (s, 3H); 13C NMR (126 MHz, CDCl3) δ 156.08, 148.60, 141.96, 123.71, 128.31, 123.28, 120.41, 120.09, 116.14, 111.91, 98.09, 55.76; IR (cm⁻¹) 3234 (NH str.), 2933 (CH str.), 1626 (C=N), 1578 (C=C), 1179 (C=O); HRMS (ES⁺) Calculated for C14H12N2O [M + H⁺]: 225.0128, found 225.0142.

2-(2-Methylphenyl)pyrrolo[2,3-b]pyridine 15b. The product 15b was furnished as a cream white solid (0.1542 g, 88%). Mp: 182–184 °C; 1H NMR (500 MHz, CDCl3) δ 12.35 (s, 1H), 8.25 (d, J = 3.3, 1H), 7.92 (d, J = 7.7, 1H), 7.82 (d, J = 8.8, 2H), 7.15–6.97 (m, 3H), 6.67 (s, 1H), 3.90 (s, 3H); 13C NMR (126 MHz, CDCl3) δ 159.83, 149.92, 141.64, 139.57, 128.25, 127.21, 125.15, 122.56, 116.06, 114.32, 96.24, 55.44; IR (cm⁻¹) 3281 (NH str.), 2986 (CH str.), 1671 (C=N), 1594 (C=C), 1176 (C=O); HRMS (ES⁺) Calculated for C14H12N2O [M + H⁺]: 225.0128, found 225.0143.

2-(3-Chlorophenyl)pyrrolo[2,3-b]pyridine 15c. The azaindole 15c was produced as a cream white solid (0.08213 g,
5-Methyl-2-phenyl-1H-pyrrolo[2,3-b]pyridine 16a. The product 16a was formed as a cream white solid (0.1001 g, 94%). 1H NMR (500 MHz, DMSO) δ 11.89 (s, 1H), 8.07 (d, J = 1.3, 1H), 7.93 (d, J = 7.5, 2H), 7.72 (s, 1H), 7.46 (t, J = 7.2, 1H), 7.34 (t, J = 7.4, 1H), 6.84 (d, J = 1.8, 1H), 2.37 (s, 3H); 13C NMR (126 MHz, DMSO) δ 148.33, 143.62, 138.25, 131.71, 128.84, 127.83, 127.56, 125.19, 124.37, 120.73, 96.49, 18.09.

2-(2-Methoxyphenyl)-5-methyl-1H-pyrrolo[2,3-b]pyridine 16b. The desired azaindole 16b was formed as a cream white solid (0.1011 g, 94%). Ms: 169–171 °C; 1H NMR (500 MHz, CDCl3) δ 10.21 (s, 1H), 8.10 (s, 1H), 7.83 (dd, J = 7.8, 1.6, 1H), 7.69 (s, 1H), 7.33–7.26 (m, 1H), 7.05 (d, J = 13.7, 10.3, 4.6, 2H), 6.75 (d, J = 2.0, 1H), 3.98 (s, 3H), 2.42 (s, 3H); 13C NMR (126 MHz, DMSO) δ 155.75, 150.82, 146.03, 142.56, 136.31, 129.14, 128.10, 129.77, 125.04, 121.45, 120.41, 120.14, 119.11, 97.37, 55.75, 18.60; IR (cm−1) 3420 (NH str.), 2981 (CH str.), 1624 (C=O) (NH str.); 3H NMR (126 MHz, CDCl3) δ 155.79, 147.59, 147.46, 143.16, 135.31, 133.16, 129.51, 127.96, 126.28, 126.05, 115.41, 96.61, 55.83, 18.50; IR (cm−1) 3298 (NH str.), 2972 (CH str.), 1635 (C=N), 1575 (C=C), 1168 (C=C–O); HRMS (ESI) Calculated for C11H11C4N3O [M + H]+: 239.1184, found 269.1190.

5-Chloro-1H-pyrrolo[2,3-b]pyridine 17a. The azaindole 17a was yielded as a cream white solid (0.1012 g, 82%). Ms: 239–242 °C; 1H NMR (500 MHz, DMSO) δ 12.07 (s, 1H), 8.10 (s, 1H), 8.03 (s, 1H), 7.90 (d, J = 7.7, 1H), 7.75 (s, 1H), 7.48 (t, J = 7.6, 1H), 7.39 (d, J = 7.8, 1H), 6.97 (s, 1H), 2.38 (s, 3H); 13C NMR (126 MHz, DMSO) δ 148.34, 144.28, 136.58, 133.85, 133.80, 130.69, 127.90, 127.46, 124.75, 124.63, 123.76, 120.52, 97.78, 18.07; IR (cm−1) 3347 (NH str.), 2974 (CH str.), 1630 (C=N), 1587 (C=C); HRMS (ESI) Calculated for C11H12Cl2N2 [M + H]+: 243.0689, found 243.0697.

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3-{Phenylethynyl}-5-{[(trimethylsilyl)ethynyl]pyriddin-2-amine

19. To a flame-dried round-bottom flask under an argon atmosphere containing 2-amino-3-iodopyridine derivative 10 (1.379 g, 4.61 mmol) was added CuI (0.01756 g, 2 mol%) and triethylamine (5 ml) was then added and the reaction mixture was stirred at rt, until no starting material was present as monitored by thin layer chromatography (TLC). After this time, the reaction was quenched with a saturated aqueous ammonium chloride and was extracted with dichloromethane. The combined organic extracts were dried over magnesium sulfate, filtered through celite and the excess solvent was removed on a rotary evaporator. The resulting crude material was taken up in dry dichloromethane and washed with aqueous sodium carbonate (10%). Purification by flash silica using flash chromatography (30% EthOAc-hexane mixture (10:1) afforded 19 as a yellowish solid (1.139 g, 85%). Mp: 155–158 °C; 1H NMR (300 MHz, CDCl₃) δ 8.18 (d, J = 1.9, 1H), 7.71 (d, J = 2.1, 1H), 7.57–7.46 (m, 2H), 7.37 (dd, J = 9.6, 6.1, 3H), 5.34 (d, J = 23.2, 2H), 0.26 (s, 9H); 13C NMR (75 MHz, CDCl₃) δ 157.79, 151.11, 142.72, 131.56, 128.89, 128.53, 122.35, 109.63, 108.85, 101.84, 95.90, 94.99, 83.38, 0.01; IR (cm⁻¹) 3283 (NH str.), 2956 (CH str.), 2144 (alkyne), 1625 (C=–N), 1583 (C= –C); HRMS (ES⁺) Calculated for C₁₉H₁₅N₂Si [M + H⁺]: 303.0133, found 303.0146.

2-Phenyl-5-{[(trimethylsilyl)ethynyl]-1H-pyrrolo[2,3-b]pyridine

20. The forgoing 2-amino-3-ethynylpyridine derivative 19 (0.551 g, 1.90 mmol) was dissolved in MeCN. To the resulting solution was added trifluoroacetic acid (0.2164 g, 0.15 ml, 1.90 mmol) in one portion followed by the addition of trifluoroacetic anhydride (0.5187 g, 0.35 ml, 2.47 mmol) dropwise. The resulting mixture was then heated to reflux for 8 h and then allowed to cool to rt. The solvent was then removed on a rotary evaporator and the resulting crude material was dissolved in dichloromethane and washed with aqueous sodium carbonate (10%). Purification by flash silica using ethyl acetate–hexane mixture (10:50%) gave 2-phenyl-5-{[(trimethylsilyl)ethynyl]-1H-pyrrolo[2,3-b]pyridine 20 (0.4236 g, 91%). Mp: 255–258 °C; 1H NMR (500 MHz, CDCl₃) δ 11.56 (s, 1H), 8.40 (d, J = 1.7, 1H), 8.05 (d, J = 1.6, 1H), 7.85–7.76 (m, 2H), 7.53 (t, J = 7.7, 2H), 7.47–7.38 (m, 1H), 6.75 (d, J = 2.0, 1H), 0.30 (s, 9H); 13C NMR (126 MHz, CDCl₃) δ 148.74, 146.08, 140.24, 131.93, 131.78, 129.19, 126.64, 125.75, 121.46, 112.42, 103.65, 97.73, 94.57, 0.06; IR (cm⁻¹) 3249 (NH str.), 2957 (CH str.), 1609 (C= –N), 1586 (C= –C); HRMS (ES⁺) Calculated for C₁₉H₁₃N₂Si [M + H⁺]: 291.1318, found 291.1327.

Antimicrobial activity

The substituted azaindoles were quantitatively evaluated for antimicrobial activity using the minimum inhibitory concentration (MIC) assay.18 The azaindoles were tested against a number of reference test organisms including Gram-positive (Enterococcus faecalis ATCC 29212 and Staphylococcus aureus ATCC 25923), Gram-negative (Pseudomonas aeruginosa NCTC 9027, Escherichia coli ATCC 8739 and Klebsiella pneumoniae ATCC 13883) and yeasts (Candida tropicalis ATCC 103380, Candida albicans ATCC 10231 and Cryptococcus neoformans ATCC 90112). The positive control used for the bacteria was ciprofloxacin, while that used for the yeasts was amphotericin. The solvent (acetone) was included in the assay as a negative control. All antimicrobial assays were undertaken at least in duplicate or triplicate where necessary.

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Notes and references

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