Supporting Information
for
Studies on the interaction of isocyanides with imines: reaction scope and mechanistic variations

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Experimental procedures

General information

Unless stated otherwise, all reactions were carried out under argon atmosphere in dried glassware. Commercially available reactants were used without further purification. Thin-layer chromatography was performed on pre-coated Merk silica gel 60 F254 plates and visualized under an UV lamp. $^1$H, and $^{13}$C NMR spectra were recorded on a Varian Mercury 400 (at 400 MHz, and 100 MHz respectively). Unless otherwise quoted, NMR spectra were recorded in CDCl$_3$ solution with TMS as an internal reference. Data for $^1$H NMR spectra are reported as follows: chemical shift (δ ppm), multiplicity, integration and coupling constants (Hz). Data for $^{13}$C NMR spectra are reported in terms of chemical shift (δ ppm). IR spectra were recorded using a Thermo Nicolet Nexus spectrometer and are reported in frequency of absorption (cm$^{-1}$). High resolution mass spectrometry was performed by the University of Barcelona Mass Spectrometry Service.

The microwave-assisted reactions were performed by setting the temperature at reported number and the maximum power and pressure at 250 W and 200 psi respectively. The power is automatically altered by the device to maintain the set temperature for the reported duration of time. A ramp period of 10 min is set to reach the desired temperature and the device automatically starts the emission period as soon as the set temperature is obtained. During the reactions the stirring option is on and a stream of nitrogen cools the reactor. The Max Power option is also on to maintain the reaction temperature while cooling. This option should not be used for polar solvents.

The $N$-aryl-aromatic imines 1 were prepared by mixing the amine and carbonyl components in ACN, in the presence of 4 Å MS at room temperature, according to a standard method. $N$-alkyl imines and activated substrates were prepared following the described procedures (see below).

Preparation of imines and related compounds (1)

General procedure

Equimolar amounts of the corresponding aldehyde and amine (1:1, 2 mmol) together with 2 g of activated MS 4 Å were placed in a 50 mL flask, 10 mL of dry ACN were added and the solution was stirred under nitrogen for 24 h or until all the starting material was consumed. The reaction progress was monitored by TLC or HPLC. After the reaction was complete, 10 mL of saturated aqueous NaHCO$_3$ solution were added. Molecular sieves were filtered and washed with 20 mL of EtOAc. The aqueous layer was extracted with $3 \times 10$ mL of EtOAc. The combined organic layers were dried over Na$_2$SO$_4$, filtered and concentrated in vacuo. The resulting crude imine was used without further purification.
General procedure for isatin imines

Equimolar amounts of isatin and the corresponding amine (1:1, 5 mmol) were placed in a 100 mL flask, 20 mL of absolute EtOH and 0.2 mL glacial acetic acid were added and the mixture was refluxed for 12 h or until all the starting material was consumed. The reaction progress was monitored by TLC or HPLC. After the reaction was complete, the solution was set aside to gradually cool down overnight. The formed crystals were filtered, washed with cold EtOH and dried in vacuo [1].

The following compounds were prepared according to literature procedures:

N-(4-Chlorobenzylidene)propan-1-amine [2]

(E)-Ethyl 2-((4-methoxyphenyl)imino)acetate [3]

(S)-(+)N-(4-Chlorobenzylidene)-4-methylbenzenesulfinamide [4]

1-(4-Chlorobenzylidene)-2-(4-methoxyphenyl)hydrazine [5]

4-Chlorobenzaldehyde oxime [6]
General procedures for the azetidine formation

**General procedure A**
0.5 mmol (1 equiv) of imine 1 were dissolved in 2 mL of dry THF. To this solution was dropwise syringed 0.5 mmol (1 equiv) of BF$_3$·Et$_2$O while stirring. In case of using a solid isocyanide, it was first dissolved in 1 mL of THF. After 2 min, 1.5 mmol (3 equiv.) of the corresponding isocyanide was gradually added. The solution was stirred for 24 h or until the reaction was complete or showed no evolution. The reaction was quenched with 10 mL of saturated NaHCO$_3$ solution and extracted with $3 \times 10$ mL of AcOEt. The combined organic phases were dried over Na$_2$SO$_4$, concentrated in vacuo and purified by flash chromatography (Hexanes/EtOAc).

**General procedure B**
In a microwave reactor tube, 0.5 mmol (1 equiv) of imine 1 were dissolved in 2 mL of dry THF. To this solution was dropwise syringed 0.5 mmol (1 equiv) of BF$_3$·Et$_2$O while stirring. After 2 minutes, 1.5 mmol (3 equiv) of the corresponding isocyanide was gradually added. In case of using a solid isocyanide, it was first dissolved in 1 mL of THF. The solution was irradiated in the microwave for 30 min at 65 °C. Then it was allowed to cool to room temperature. The reaction was quenched with 10 mL of saturated NaHCO$_3$ solution and extracted with $3 \times 10$ mL of EtOAc. The combined organic phases were dried over Na$_2$SO$_4$, concentrated in vacuo and purified by flash chromatography (hexanes/EtOAc).

**Characterization data**

![Structure of 3a](image)

* $N,N'$-(4-(4-Chlorophenyl)-1-(4-methoxyphenyl)azetidine-2,3-diylidene)bis(2-methylpropan-2-amine) (3a).

According to procedure B, azetidine 3a was obtained as a pale yellow solid (48%).

$C_{24}H_{30}ClN_3O$ ($M = 411.97$ g·mol$^{-1}$).

IR (neat, cm$^{-1}$): 3417, 2969, 2930, 2860, 1719, 1668, 1508, 1380, 1303, 1245, 1207, 1130, 1028, 932, 829.
$^{1}$$H$ NMR (400 MHz, CDCl$_3$) $\delta$ 7.42 (d, $J = 9.1$ Hz, 2H), 7.33 (s, 4H), 6.76 (d, $J = 9.1$ Hz, 2H), 5.38 (s, 1H), 3.71 (s, 3H), 1.49 (s, 9H), 1.09 (s, 9H).

$^{13}$$C$ NMR (101 MHz, CDCl$_3$) $\delta$ 154.6, 153.9, 152.3, 136.8, 134.4, 134.0, 129.36, 129.35, 117.2, 114.2, 70.3, 58.1, 55.5, 53.9, 53.5, 30.8, 30.2.

HRMS (ESI): calculated for C$_{24}$H$_{31}$ClN$_3$O [M + H]$^+$: 412.215, found 412.215.

$N,N'$-(4-(4-Chlorophenyl)-1-(4-methoxyphenyl)azetidine-2,3-diylidene)dicyclohexanamine (3b).

According to procedure B, using 20% BF$_3$·Et$_2$O, 3b was obtained as an amorphous solid (41%).

C$_{28}$H$_{34}$ClN$_3$O (M = 464.04 g·mol$^{-1}$).

IR (neat, cm$^{-1}$): 2930, 2853, 1713, 1674, 1508, 1367, 1297, 1245, 1137, 1008, 823, 733.

$^{1}$$H$ NMR (400 MHz, CDCl$_3$) $\delta$ 7.29 (d, $J = 9.1$ Hz, 2H), 7.25 (d, $J = 9.6$ Hz, 2H), 6.69 (d, $J = 9.1$ Hz, 2H), 5.39 (s, 1H), 4.21 (m, 2H), 3.64 (s, 3H), 3.18 (m, 16H). 3.18 (m, 16H).

$^{13}$$C$ NMR (101 MHz, CDCl$_3$) $\delta$ 157.1, 154.5, 152.2, 135.6, 134.1, 133.8, 129.3, 128.2, 116.9, 114.3, 69.3, 61.2, 57.1, 55.49, 55.44, 35.34, 35.26, 33.6, 32.8, 25.9, 25.5, 24.8, 23.87, 23.72.

HRMS (ESI): calculated for C$_{28}$H$_{35}$ClN$_3$O [M + H]$^+$ 464.2463, found 464.2452.

$2,3$-Bis(benzylimino$)$-4-(4-chlorophenyl)-1-(4-methoxyphenyl)azetidine (3c).

According to procedure A, using 20% BF$_3$·Et$_2$O, azetidine 3c was obtained as an amorphous solid (63%).
\[
\text{C}_{30}\text{H}_{26}\text{ClN}_3\text{O} \quad (M = 480.00 \text{ g\cdot mol}^{-1}).
\]

**IR** (neat, cm\(^{-1}\)): 3058, 3026, 2930, 1726, 1687, 1508, 1450, 1245, 1130, 1028, 829.

**\(^1\)H NMR** (400 MHz, CDCl\(_3\)) \(7.41\) (ddd, \(J = 6.2, 1.3, 0.7 \text{ Hz}, 2\text{H}\), 7.35 – 7.31 (m, 2H), 7.28 (cs, 6H), 7.21 – 7.12 (m, 4H), 6.96 (dd, \(J = 7.9, 1.6 \text{ Hz}, 2\text{H}\), 6.70 (d, \(J = 9.1 \text{ Hz}, 2\text{H}\), 5.51 (s, 1H), 5.09 (s, 2H), 4.45 (d, \(J = 14.4 \text{ Hz}, 1\text{H}\), 4.41 (d, \(J = 14.4 \text{ Hz}, 1\text{H}\), 3.64 (s, 3H).

**\(^{13}\)C NMR** (101 MHz, CDCl\(_3\)) \(\delta 160.0, 154.0, 152.3, 140.7, 136.7, 133.8, 133.2, 132.1, 128.6, 127.6, 127.4, 127.2, 126.6, 126.47, 126.1, 125.3, 116.2, 113.3, 68.9, 55.3, 54.4, 52.4.

**HRMS** (ESI): calculated for \(\text{C}_{30}\text{H}_{27}\text{ClN}_3\text{O} [M + H]^+ 480.1837\), found 480.1835.

\[
N,N'-(4-(4-Chlorophenyl)-1-(p-tolyl)azetidine-2,3-diylidene)bis(4-methoxyaniline) (3d)
\]

According to procedure B, azetidine 3d was obtained as a yellow amorphous solid (19%).

\[
\text{C}_{30}\text{H}_{27}\text{ClN}_3\text{O}_2 \quad (M = 495.99 \text{ g\cdot mol}^{-1}).
\]

**IR** (neat, cm\(^{-1}\)): 1656, 1504, 1499, 1378, 1294, 1241, 1030, 830.

**\(^1\)H NMR** (400 MHz, CDCl\(_3\)) \(7.54\) (d, \(J = 8.9 \text{ Hz}, 2\text{H}\), 7.44 (d, \(J = 8.5 \text{ Hz}, 2\text{H}\), 7.15 (d, \(J = 8.5 \text{ Hz}, 2\text{H}\), 7.07 (d, \(J = 8.0 \text{ Hz}, 4\text{H}\), 6.89 (d, \(J = 8.9 \text{ Hz}, 2\text{H}\), 6.74 (bs, 4H), 5.84 (s, 1H), 3.81 (s, 3H), 3.78 (s, 3H), 2.28 (s, 3H).

**\(^{13}\)C NMR** (101 MHz, CDCl\(_3\)) \(\delta 159.1, 158.1, 156.6, 150.2, 139.71, 139.58, 136.8, 134.4, 133.1, 132.8, 129.7, 129.1, 128.4, 124.9, 122.6, 116.7, 114.05, 113.87, 69.3, 55.65, 55.58, 21.1.

**HRMS** (ESI): calculated for \(\text{C}_{30}\text{H}_{27}\text{ClN}_3\text{O}_2 [M + H]^+ 496.1786\), found 496.1787.
According to procedure B, azetidine 3e was obtained as a cream amorphous solid (19%).

\[ \text{C}_{24}\text{H}_{30}\text{ClN}_{3}\text{O} \text{ (M = 411.97 g mol}^{-1} \text{).} \]

IR (neat, cm\(^{-1}\)): 2962, 2861, 1722, 1671, 1477, 1203, 837.

\(^1\)H NMR (400 MHz, CDCl\(_3\)) \( \delta \) 7.51 (t, \( J = 2.1 \) Hz, 1H), 7.34 (bs, 4H), 7.04 (t, \( J = 8.2 \) Hz, 1H), 6.72 (dd, \( J = 8.2, 1.2 \) Hz, 1H), 6.45 (ddd, \( J = 8.2, 2.1, 1.2 \) Hz, 1H), 5.39 (s, 1H), 3.74 (s, 3H), 1.48 (s, 9H), 1.09 (s, 9H).

\(^{13}\)C NMR (101 MHz, CDCl\(_3\)) \( \delta \) 159.9, 153.6, 152.3, 141.1, 136.5, 134.4, 129.32, 129.26, 129.20, 107.90, 107.81, 101.9, 70.4, 58.1, 55.0, 53.9, 30.6, 30.0.

HRMS (ESI): calculated for \( \text{C}_{24}\text{H}_{31}\text{ClN}_{3}\text{O} \text{ [M + H]}^+ \): 412.215, found 412.215.

According to procedure B, azetidine 3f was obtained as a cream solid (12%).

\[ \text{C}_{24}\text{H}_{30}\text{ClN}_{3}\text{O} \text{ (M = 411.97 g mol}^{-1} \text{).} \]

IR (neat, cm\(^{-1}\)): 3084, 2952, 2819, 1721, 1660, 1507, 1233, 1121, 754.

\(^1\)H NMR (400 MHz, CDCl\(_3\)) \( \delta \) 7.47 (d, \( J = 9.0 \) Hz, 2H), 7.40 (dd, \( J = 6.9, 2.9 \) Hz, 1H), 7.35 (dd, \( J = 6.9, 3.6 \) Hz, 1H), 7.21 (m, 2H), 6.77 (d, \( J = 9.0 \) Hz, 2H), 5.99 (s, 1H), 3.71 (s, 3H), 1.48 (s, 9H), 1.09 (s, 9H).
$^{13}$C NMR (101 MHz, CDCl$_3$) δ 154.7, 154.4, 153.4, 152.5, 135.8, 133.66, 133.56, 131.8, 129.89, 129.79, 129.69, 128.9, 127.8, 122.5, 117.1, 114.2, 65.7, 57.6, 55.4, 53.7, 30.7, 29.6.

HRMS (ESI): calculated for C$_{24}$H$_{31}$ClN$_3$O [M + H]$^+$: 412.215, found 412.215.

The compound is depicted in the image. The structure of $N$-(4-(2-Chlorophenyl)-3-imino-1-(4-methoxyphenyl)-azetidin-2-yldene)-2-methylpropan-2-amine (3f$^\prime$)

According to procedure B, azetidine 3f$^\prime$ was obtained as yellow oil (15%).

C$_{20}$H$_{22}$ClN$_3$O (M = 355.86 g·mol$^{-1}$).

IR (neat, cm$^{-1}$): 3379, 2962, 2820, 1640, 1508, 1233, 1040, 806.

$^1$H NMR (400 MHz, CDCl$_3$) δ 7.47 – 7.43 (m, 1H), 7.37 – 7.33 (m, 1H), 7.31 – 7.14 (m, 2H), 6.73 (d, $J$ = 9.0 Hz, 2H), 6.59 (d, $J$ = 9.0 Hz, 2H), 5.68 (d, $J$ = 5.4 Hz, 1H), 5.25 (d, $J$ = 5.4 Hz, 1H), 3.71 (s, 3H), 1.43 (s, 9H).

$^{13}$C NMR (101 MHz, CDCl$_3$) δ 152.7, 139.7, 137.8, 135.2, 134.6, 130.2, 129.8, 128.8, 127.7, 115.1, 114.8, 111.2, 61.3, 58.7, 55.7, 29.3.

HRMS (ESI): calculated for C$_{20}$H$_{23}$ClN$_3$O [M + H]$^+$: 356.1524, found 356.1513.

The compound is depicted in the image. The structure of N,N'-(4-(4-Chlorophenyl)-1-(o-tolyl)azetidine-2,3-diyldene)bis(2-methylpropan-2-amine) (3g)

According to procedure B, azetidine 3g was obtained as a brown paste (31%).

C$_{24}$H$_{30}$ClN$_3$ (M = 395.97 g·mol$^{-1}$).

IR (neat, cm$^{-1}$): 2952, 2861, 2850, 1671, 1488, 1203, 837.
1H NMR (400 MHz, CDCl₃) δ 7.45 (d, J = 8.1 Hz, 1H), 7.25 (d, J = 8.7 Hz, 2H), 7.20 (d, J = 8.7 Hz, 2H), 7.04 (m, 2H), 6.94 (m, 1H), 5.56 (s, 1H), 2.29 (s, 3H), 1.47 (s, 9H), 1.11 (s, 9H).

13C NMR (101 MHz, CDCl₃) δ 154.9, 153.0, 137.0, 136.4, 134.2, 132.0, 131.2, 129.4, 129.0, 125.9, 125.0, 122.8, 118.6, 114.9, 71.6, 58.2, 53.6, 30.7, 30.0, 20.1.

HRMS (ESI): calculated for C₂₄H₃₁ClN₃ [M + H]⁺: 396.2201, found 396.2194.

N-(4-(4-Chlorophenyl)-3-imino-1-propylazetidin-2-ylidene)-2-methylpropan-2-amine (3h’)

According to procedure B, azetidine 3h’ was obtained as a lemon amorphous solid (32%).

C₁₆H₂₂ClN₃ (M = 291.82 g·mol⁻¹).

IR (neat, cm⁻¹): 3359, 3318, 2952, 2861, 2149, 1579, 1091, 837.

1H NMR (400 MHz, CDCl₃) δ 7.42 (d, J = 8.4 Hz, 2H), 7.32 (d, J = 8.4 Hz, 2H), 5.79 (s, 1H), 2.75 (cs, 2H), 1.57 (s, 1H), 1.41 (cs, 2H), 1.17 (s, 9H), 0.76 (t, J = 7.4 Hz, 3H).

13C NMR (101 MHz, CDCl₃) δ 161.1, 136.0, 131.1, 130.5, 129.1, 123.4, 54.7, 46.1, 29.6, 29.1, 24.5, 11.3.

HRMS (ESI): calculated for C₁₆H₂₃ClN₃ [M + H]⁺: 292.1575, found 292.1575.

Ethyl 3,4-bis(tert-butylimino)-1-(4-methoxyphenyl)azetidine-2-carboxylate (3i)

According to procedure A, using 20% BF₃, Et₂O, azetidine 3i was obtained as a white amorphous solid (9%).

C₂₁H₃₁N₃O₃ (M = 373.49 g·mol⁻¹).
IR (neat, cm⁻¹): 2960, 1737, 1717, 1674, 1187.

¹H NMR (400 MHz, CDCl₃) δ 7.46 (d, J = 9.1 Hz, 2H), 6.85 (d, J = 9.1 Hz, 2H), 4.95 (s, 1H), 4.19 (cs, 2H), 3.77 (s, 3H), 1.42 (s, 9H), 1.30 (s, 9H), 1.20 (t, J = 7.1 Hz, 3H).

¹³C NMR (126 MHz, CDCl₃) δ 169.6, 154.8, 151.4, 147.7, 134.2, 116.7, 114.5, 68.5, 62.0, 58.5, 55.7, 54.0, 30.7, 29.9, 14.1.

HRMS (ESI): calculated for C₂₁H₃₂N₃O₃ [M + H]⁺: 374.2438, found 374.2441.

Ethyl 3,4-bis(cyclohexylimino)-1-(4-methoxyphenyl)azetidine-2-carboxylate (3j)

According to procedure A, using 20% BF₃·Et₂O, azetidine 3j was obtained as an amorphous white solid (34%).

C₂₅H₃₅N₃O₃ (M = 425.56 g·mol⁻¹).

IR (neat, cm⁻¹): 2930, 2854, 1720, 1687, 1514.

¹H NMR (400 MHz, CDCl₃) δ 7.27 (d, J = 8.0, Hz, 2H), 6.79 (d, J = 8.0 Hz, 2H), 4.98 (s, 1H), 4.16 (m, Hz, 2H), 4.07 (m, 2H), 3.70 (s, 3H), 3.38 (m, 4H), 1.72 - 1.25(m, 16H), 1.18 (t, J = 7.0 Hz, 3H).

¹³C NMR (101 MHz, CDCl₃) δ 206.5, 154.5, 134.0, 116.5, 114.4, 67.0, 62.00, 61.98, 61.9, 57.3, 55.5, 35.1, 33.6, 33.4, 30.9, 25.8, 25.5, 24.6, 23.8, 23.7, 14.1 (three quaternary carbon signals not detected)

HRMS (ESI): calculated for C₂₅H₃₀N₄O [M + H]⁺: 426.2751, found 426.2745.

3,4-bis(tert-Butylimino)-1-(4-methoxyphenyl)spiro[azetidine-2,3'-indolin]-2'-one (3k)
According to procedure B, azetidine 3k was obtained as a brown paste (14%).

\[ \text{C}_{25}\text{H}_{30}\text{N}_{4}\text{O}_{2} \ (M = 418.53 \text{ g mol}^{-1}) \].

\(^1\text{H NMR} \ (400 \text{ MHz, CDCl}_3) \delta 9.05 \text{ (bs, 1H), 7.45 (m, 1H), 7.35 (m, 1H), 7.31 – 7.14 (m, 2H), 6.73 (d, \ J = 9.0 \text{ Hz, 2H}), 6.59 (d, \ J = 9.0 \text{ Hz, 2H}), 3.71 \text{ (s, 3H), 1.43 (s, 9H), 0.98 (s, 9H).} \]

\text{LRMS (HPLC-MS): calculated for C}_{25}\text{H}_{31}\text{N}_{4}\text{O}_{2} [M + H]^+: 419.24, found 419.}

\[ 3,4\text{-bis(tert-Butylimino)-1-(p-tolyl)spiro[azetidine-2,3'}\text{-indolin}-2'}\text{-one (3l) } \]

According to procedure A, azetidine 3l was obtained as an amorphous cream solid (28%).

\[ \text{C}_{25}\text{H}_{30}\text{N}_{4}\text{O} \ (M = 402.53 \text{ g mol}^{-1}) \].

\( \text{IR (neat, cm}^{-1} \): 2962, 2861, 1742, 1691, 1508, 1355, 1183, 928. \)

\(^1\text{H NMR} \ (400 \text{ MHz, CDCl}_3) \delta 9.42 \text{ (s, 1H), 7.27 (m, \ J = 7.8, 1.2 \text{ Hz, 1H}), 7.22 (cs, 3H), 7.01 (m, \ J = 7.8, 0.8 \text{ Hz, 1H}), 6.93 – 6.90 (m, 2H), 6.89 (s, 1H), 2.17 (s, 3H), 1.50 (s, 9H), 1.02 (s, 9H).} \)

\(^{13}\text{C NMR} \ (101 \text{ MHz, CDCl}_3) \delta 175.4, 150.7, 147.3, 140.4, 136.9, 131.6, 130.5, 129.4, 125.44, 124.5, 123.5, 116.0, 111.3, 77.4, 58.5, 54.1, 30.6, 29.5, 20.8. \)

\text{HRMS (ESI): calculated for C}_{25}\text{H}_{31}\text{N}_{4}\text{O} [M + H]^+: 403.2492, found 403.249.}

\[ N\text{-}(\text{tert-Butyl)-2-(4-chlorophenyl)-N'}\text{-}(4\text{-methoxyphenyl)-2-((4-methoxyphenyl) amino)acetimidamide (4)}} \]

Amidine 4a was obtained by direct stirring aniline 1a, aldehyde 2a and tert-butyl
isocyanide, with Sc(OTf)₃ (20% mol) at rt. for 14 h in ACN. After standard aqueous work up and EtOAc extraction, the crude was purified through column chromatography (SiO₂, Hexanes-EtOAC 50:50) to yield amidine 4 (28%).

\[
C_{26}H_{30}ClN_3O_2 \quad (M = 451.99 \text{ g·mol}^{-1}).
\]

\[
^{1}H \text{ NMR} \quad (400 \text{ MHz, CDCl}_3) \quad \delta \quad 7.25 \quad (d, \quad J = 8.5 \text{ Hz}, \quad 2\text{H}), \quad 7.08 \quad (d, \quad J = 8.5 \text{ Hz}, \quad 2\text{H}), \quad 6.83 \quad (d, \quad J = 9.0 \text{ Hz}, \quad 2\text{H}), \quad 6.65 \quad (\text{cs}, \quad 4\text{H}), \quad 6.40 \quad (d, \quad J = 8.8 \text{ Hz}, \quad 2\text{H}), \quad 6.00 \quad (m, \quad 1\text{H}), \quad 4.86 \quad (s, \quad 1\text{H}), \quad 3.78 \quad (s, \quad 3\text{H}), \quad 3.73 \quad (s, \quad 3\text{H}), \quad 3.46 \quad (\text{bs}, \quad 1\text{H}), \quad 1.46 \quad (s, \quad 9\text{H}).
\]

\[
^{13}C \text{ NMR} \quad (101 \text{ MHz, CDCl}_3) \quad \delta \quad 154.5, 153.4, 143.8, 141.0, 138.5, 134.0, 129.4, 128.9, 122.9, 116.4, 114.93, 114.80, 114.8, 113.96, 59.7, 55.7, 55.5, 50.8, 28.4.
\]

LRMS (HPLC-MS): calculated for \(C_{26}H_{31}ClN_3O_2\) [M + H]^+: 452.20, found 452.

\[
N,N',N''-(5-(4-Chlorophenyl)-1-(4-methoxyphenyl)-pyrrolidine-2,3,4-triylidene)tris(2-methylpropan-2-amine) \quad (5)
\]

According to procedure B, pyrrolidine 5 was obtained as a brown paste (9%).

\[
C_{29}H_{39}ClN_4O \quad (M = 495.10 \text{ g·mol}^{-1}).
\]

IR (neat, cm⁻¹): 2962, 1691, 1640, 1508, 1355, 1233, 816.

\[
^{1}H \text{ NMR} \quad (400 \text{ MHz, CDCl}_3) \quad \delta \quad 7.32 \quad (d, \quad J = 9.2 \text{ Hz}, \quad 2\text{H}), \quad 7.26 \quad (d, \quad J = 8.5 \text{ Hz}, \quad 2\text{H}), \quad 7.16 \quad (d, \quad J = 8.5 \text{ Hz}, \quad 2\text{H}), \quad 6.73 \quad (d, \quad J = 9.2 \text{ Hz}, \quad 2\text{H}), \quad 5.59 \quad (s, \quad 1\text{H}), \quad 3.74 \quad (s, \quad 3\text{H}), \quad 1.45 \quad (s, \quad 9\text{H}), \quad 1.43 \quad (s, \quad 9\text{H}), \quad 1.29 \quad (s, \quad 9\text{H}).
\]

HRMS (ESI): calculated for \(C_{29}H_{40}ClN_4O\) [M + H]^+: 495.2885, found 495.2875.
2-(4-Chlorophenyl)-6-methoxy-1H-indol-3-amine (6)

According to procedure B, indole 6 was obtained as an slightly unstable amorphous cream solid (27%).

C\textsubscript{15}H\textsubscript{13}ClN\textsubscript{2}O (M = 272.72 g·mol\textsuperscript{-1}).

IR (neat, cm\textsuperscript{-1}): 3237, 2952, 2881, 2393, 2220, 1488, 1152, 1020, 827.

\textsuperscript{1}H NMR (400 MHz, DMSO) δ 12.45 (m, 1H), 7.96 (d, J = 8.6 Hz, 2H), 7.71 (d, J = 8.6 Hz, 2H), 7.54 (d, J = 8.7 Hz, 1H), 7.01 (d, J = 2.2 Hz, 1H), 6.93 (dd, J = 8.7, 2.2 Hz, 1H), 3.84 (s, 3H), 1.25 (s, 2H).

\textsuperscript{13}C NMR (101 MHz, DMSO) δ 157.8, 142.8, 137.0, 134.6, 129.8, 128.7, 122.5, 119.7, 112.9, 95.8, 55.9.

Ethyl 3-(tert-butilamino)-2-(4-methoxyphenylamino)-3-oxopropanoate (7)

According to procedure A, aminoamide 7 was obtained as light brown paste (41%).

C\textsubscript{16}H\textsubscript{24}N\textsubscript{2}O\textsubscript{4} (M = 308.37 g·mol\textsuperscript{-1}).

IR (neat, cm\textsuperscript{-1}): 3381, 2968, 1741, 1699, 1619, 1512, 1464, 1367, 1244, 1179, 1033.

\textsuperscript{1}H NMR (400 MHz, CDCl\textsubscript{3}) δ 6.79 (d, J = 8.9 Hz, 2H), 6.64 (s, 1H), 6.60 (d, J = 8.9 Hz, 2H), 4.71 (s, 1H), 4.36 – 4.20 (m, 2H), 3.75 (s, 3H), 1.32 (s, 9H), 1.30 (t, J = 7.1 Hz, 3H).

\textsuperscript{13}C NMR (101 MHz, CDCl\textsubscript{3}) δ 169.4, 165.5, 153.4, 139.6, 115.3, 114.8, 64.2, 62.4, 55.5, 51.3, 28.4, 14.0.
HRMS (ESI): calculated for $\text{C}_{16}\text{H}_{25}\text{N}_{2}\text{O}_{4} [\text{M + H}]^+: 309.1809$, found 309.1809.

3',4'-bis(tert-Butylimino)-6'-methoxy-3',4'-dihydro-1'H-spiro[indoline-3,2'-quinolin]-2-one (8)

According to procedure B, tetrahydroquinoline 8 was obtained as a brown solid (17%).

$\text{C}_{25}\text{H}_{30}\text{N}_{4}\text{O}_{2}$ (M = 418.53 g·mol$^{-1}$).

IR (neat, cm$^{-1}$): 3420, 3227, 2962, 1681, 1508, 1223, 1040, 755.

$^1\text{H NMR}$ (400 MHz, CDCl$_3$) $\delta$ 7.84 (dd, $J$ = 7.8, 1.5 Hz, 1H), 7.70 (m, 1H), 7.42 (dd, $J$ = 7.6, 1.3 Hz, 1H), 7.32 (m, $J$ = 7.6, 1.3 Hz, 1H), 7.20 (bs, 2H), 7.12 (dd, $J$ = 8.0, 1.0 Hz, 1H), 6.45 (bs, 1H), 5.27 (bs, 1H), 3.77 (s, 3H), 1.60 (s, 9H), 1.20 (s, 9H).

$^{13}\text{C NMR}$ (101 MHz, CDCl$_3$) $\delta$ 167.2, 154.8, 153.4, 140.7, 130.3, 129.6, 128.4, 127.1, 125.7, 123.7, 121.5, 120.7, 119.6, 115.7, 105.7, 59.8, 55.6, 51.7, 29.7, 29.1.

HRMS (ESI): calculated for $\text{C}_{25}\text{H}_{31}\text{N}_{4}\text{O}_{2} [\text{M + H}]^+: 419.2442$, found 419.2438.

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Copies of the NMR spectra

\[ \text{N,N'-}(4\text{-}(4\text{-Chlorophenyl})-1\text{-}(4\text{-methoxyphenyl})\text{azetidine-2,3-diylidene)}\text{bis}(2\text{-methylpropan-2-amine}) \ (3a) \]
$N,N'-(4-(4-Chlorophenyl)-1-(4-methoxyphenyl)azetidine-2,3-diylidene)dicyclohexanamine (3b)$
$N,N'-(4-(4-Chlorophenyl)-1-(4-methoxyphenyl)azetidine-2,3-diylidene)bis(1-phenylmethanamine)$ (3c)
(N,N',N,N')-N,N'-[4-(4-Chlorophenyl)-1-(p-tolyl)azetidine-2,3-diylidene]bis(4-methoxyaniline) (3d)
$N,N'-(4-(4-Chlorophenyl)-1-(3-methoxyphenyl)azetidine-2,3-diylidene)bis(2-methylpropan-2-amine)$ (3e)
$N,N'-(4-(2-Chlorophenyl)-1-(4-methoxyphenyl)azetidine-2,3-diylidene)bis(2-methylpropan-2-amine) \ (3f)$
$N(4\text{-}(2\text{-Chlorophenyl})\text{-}3\text{-}\text{imino}\text{-}1\text{-}(4\text{-methoxyphenyl})\text{azetidin}-2\text{yldene})\text{-}2\text{-methylpropan-2-amine}$ ($3^f$)
$N,N'-(4-(4-Chlorophenyl)-1-o-tolylazetidine-2,3-diylidene)bis(2-methylpropan-2-amine)$ (3g)
N-(4-(4-Chlorophenyl)-3-imino-1-propylazetidin-2-ylidene)-2-methylpropan-2-amine (3h')
Ethyl 3,4-bis(tert-butylimino)-1-(4-methoxyphenyl)azetidine-2-carboxylate (3i)
Ethyl 3,4-bis(cyclohexylimino)-1-(4-methoxyphenyl)azetidine-2-carboxylate (3i)
3,4-bis(tert-Butylimino)-1-(4-methoxyphenyl)spiro[azetidine-2,3'-indolin]-2'-one (3k)
3,4-bis(tert-Butylimino)-1-p-tolylspiro[azetidine-2,3'-indolin]-2'-one (3l)
N-(tert-Butyl)-2-(4-chlorophenyl)-N'-(4-methoxyphenyl)-2-((4-methoxyphenyl)amino)acetimidamide (4)
$N,N',N''-(5$-(4-Chlorophenyl)-1-(4-methoxyphenyl)pyrrolidine-2,3,4-
triylidene)tris(2-methylpropan-2-amine) \ (5)$
2-(4-Chlorophenyl)-6-methoxy-1H-indol-3-amine (6)
Ethyl 3-(tert-butylamino)-2-(4-methoxyphenylamino)-3-oxopropanoate (7)
3',4'-bis(tert-Butylimino)-6'-methoxy-3',4'-dihydro-1'H-spiro[indoline-3,2'-quinolin]-2-one (8)
X-ray Structure of bis-aminoazetidine 3a

(CCDC 963354)

$N,N'-(4-(4-Chlorophenyl)-1-(4-methoxyphenyl) azetidine \cdot 2,3-$
diylidene)bis(2-methylpropan-2-amine)

Cell views of bis-iminoazetidine 3a