

# Anticancer Activity and Cellular Repression of c-MYC by the G-Quadruplex-Stabilizing 11-Piperazinylquindoline Is Not Dependent on Direct Targeting of the G-Quadruplex in the c-MYC Promoter

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## Supporting Information

**ABSTRACT:** This G-rich region of the c-MYC promoter has been shown to form a G-quadruplex structure that acts as a silencer element for c-MYC transcriptional control. In the present work, we have synthesized a series of 11-substituted quindoline analogues as c-MYC G-quadruplex-stabilizing compounds, and the cell-free and in vitro activity of these compounds were evaluated. Two lead compounds (**4** and **12**) demonstrated good cell-free profiles, and compound **4** (2-(4-(10*H*-indolo[3,2-*b*]quinolin-11-yl)piperazin-1-yl)-*N,N*-dimethylethanamine) significantly down-regulated c-MYC expression. However, despite the good cell-free activity and the effect of these compounds on c-MYC gene expression, we have demonstrated, using a cellular assay in a Burkitt's lymphoma cell line (CA46-specific), that these effects were not mediated through targeting of the c-MYC G-quadruplex. Thus, caution should be used in assigning the effects of G-quadruplex-interactive compounds that lower c-MYC to direct targeting of these promoter elements unless this assay, or similar ones, demonstrates direct targeting of the G-quadruplex in cells.

## INTRODUCTION

The MYC family of proteins [c-, N-, L-] is involved in promoting cell growth, proliferation, and apoptotic processes under cell stress. c-MYC performs these functions through the regulation of gene expression.<sup>1–3</sup> Overexpression of the c-MYC oncogene is linked with cellular proliferation and inhibition of differentiation, leading to its association with a wide range of human cancers, including colon, breast, small-cell lung, osteosarcomas, glioblastomas, lymphoma, and myeloid leukemia.<sup>4–6</sup> The transcriptional regulation of c-MYC expression involves multiple promoters and transcriptional start sites, with P1 and P2 being the predominant promoters. The nuclease hypersensitive element (NHE) III<sub>1</sub> of the P1 promoter controls 85–90% of c-MYC transcription, and the guanine-rich strand of the promoter element has been shown to form an intramolecular G-quadruplex structure that appears to act as a silencer element when stabilized by the G-quadruplex-interactive compound TMPyP4.<sup>7,8</sup>

Recently, a variety of small molecules have been synthesized and tested for their ability to interact with the c-MYC G-quadruplex. Some of these compounds include quindolines, berberine, methylene blue, and bisaryl diketene analogues.<sup>9–12</sup> Quindoline compounds were first reported as telomeric G-quadruplex-interactive agents by Guyen and co-workers.<sup>13</sup> Ou et al. synthesized and tested a series of 11-substituted quindoline analogues for G-quadruplex stabilization and c-MYC down-regulation.<sup>11</sup> Quindoline-*i* (*N*<sub>1</sub>,*N*<sub>1</sub>-diethyl-*N*<sub>2</sub>-(10*H*-indolo[3,2-*b*]quinolin-11-yl)ethane-1,2-diamine) exhibited the best c-MYC G-quadruplex stabilization, and this correlates with c-MYC down-regulation and effects on cell

proliferation in HepG2 cell lines. Targeted effects of the quindoline analogues were supported by the differential effects in Ramos and CA46 Burkitt's lymphoma cell lines.

While quindoline-*i* was an effective down-regulator of c-MYC expression, recently published evidence does not support intracellular stabilization of the NHE III<sub>1</sub> promoter G-quadruplex as its mechanism of action.<sup>14</sup> In comparison, there is much stronger evidence to show that an ellipticine compound exerts its intracellular effects through the c-MYC G-quadruplex, confirming the in vitro existence and targetability of the G-quadruplex structure.<sup>14</sup> Here we describe the design, synthesis, and screening of 11-piperazinylquindolines as c-MYC G-quadruplex-stabilizing compounds in an effort to gain target specificity, with activity in cancer cell lines, and to confirm the mechanism of action. We propose that the addition of a piperazine ring would provide steric bulk to the planar quindoline ring, resulting in increased G-quadruplex binding and reduced duplex DNA binding, which would lead to better selectivity and provide an opportunity to add substituents with varying properties and linker lengths. We synthesized a series of disubstituted analogues with the addition of a second side chain at two different positions on the quindoline ring, which generally increased G-quadruplex stabilization while maintaining cellular potency. A number of these compounds demonstrated good thermal stabilization of the c-MYC G-quadruplex, with minimal effects on other promoters. We also examined the mechanism of action of these compounds using

Received: February 28, 2012

Published: June 12, 2012

the newly described “CA46 exon-specific” assay, which is presently the only cellular assay that can discriminate between lowering of c-MYC through direct binding to the G-quadruplex in the NHE versus a secondary effect.

## MATERIALS AND METHODS

**Chemical Synthesis.** Polyphosphoric acid, phosphorus oxychloride, and 3-(piperazin-1-yl)propan-1-ol were purchased from Alfa Aesar. 1-(3-Methoxypropyl)piperazine and 1-(2-(piperidin-1-yl)ethyl)piperazine were bought from Oakwood Chemicals, and 4-(2-(piperazin-1-yl)ethyl)morpholine was purchased from AK Scientific. 4-(2-Morpholinoethyl)phenylboronic acid was obtained from CombiBlocks. All other chemicals were purchased from Sigma-Aldrich. All the solvents were obtained from Fischer Scientific. Flash chromatography was performed with silica gel (230/400 mesh, Fisher Scientific). All anhydrous reactions were carried out under positive pressure of nitrogen or argon. HPLC–MS analyses were performed on an Agilent 1100 series instrument with a Zorbax C18 reverse-phase column. HRMS results were obtained on an apex-Qe instrument. All  $^1\text{H}$  NMR and  $^{13}\text{C}$  NMR spectra were recorded on a Bruker 300 MHz or DRX 500 MHz NMR spectrometer, using deuterated solvents. The spectra are reported in ppm and referenced to deuterated DMSO (2.49 ppm for  $^1\text{H}$ , 39.5 ppm for  $^{13}\text{C}$ ) or deuterated chloroform (7.26 ppm for  $^1\text{H}$ , 77 ppm for  $^{13}\text{C}$ ). All compounds were analyzed for purity by HPLC using either MS or UV absorbance detectors. All final compounds showed  $\geq 95\%$  purity.

**Synthesis of Quindoline-i (N1,N1-Diethyl-N2-(10H-indolo[3,2-b]quinolin-1-yl)ethane-1,2-diamine).** Quindoline-i was synthesized from 11-chloroquindoline and *N,N*-diethylethane-1,2-diamine. Detailed procedures and spectral data have been reported earlier.<sup>14</sup>

**General Procedure for the Synthesis of Compounds 2–10.** A mixture of 11-chloro-10H-indolo[3,2-b]quinoline **1** and the corresponding amine was heated at an appropriate temperature for a specific time. After cooling, the reaction mixture was poured into ice-cold water. The precipitated solid was filtered, washed with water, and then dried. The crude product was purified using 2–5% methanol in chloroform as a solvent by silica gel column chromatography.

**11-(Piperazin-1-yl)-10H-indolo[3,2-b]quinoline (2).** Compound **1** (100 mg, 0.39 mmol) was reacted with piperazine (1 g) at 100 °C for 24 h to obtain 97 mg (80%) of compound **2** as a yellow solid.  $^1\text{H}$  NMR (300 MHz, DMSO- $d_6$ ):  $\delta$  10.91 (s, 1H, NH), 8.43–8.24 (m, 2H, ArH), 8.14 (d,  $J = 8.4$  Hz, 1H, ArH), 7.70–7.48 (m, 4H, ArH), 7.25 (t,  $J = 7.5$  Hz, 1H, ArH), 3.64–3.45 (m, 4H), 3.14–2.97 (m, 4H), 2.75 (br s, 1H, NH).  $^{13}\text{C}$  NMR (75 MHz, DMSO- $d_6$ ):  $\delta$  161.68, 148.14, 145.87, 144.78, 136.70, 130.32, 129.97, 127.23, 127.01, 125.12, 124.21, 122.12, 121.94, 120.21, 112.89, 51.25, 46.23. MS (ESI):  $m/z = 303.2$  [100%, (M + H) $^+$ ]. HRMS calcd for  $\text{C}_{19}\text{H}_{19}\text{N}_4$  [M + H] $^+$  303.1604, found 303.1603. HPLC MS purity 99.02%.

**11-(4-Methylpiperazin-1-yl)-10H-indolo[3,2-b]quinoline (3).** Compound **3** was obtained from **1** (500 mg, 1.98 mmol) and 1-methylpiperazine (5 mL) heated at 100 °C for 12 h. Yield of compound **3** was 530 mg (84%).  $^1\text{H}$  NMR (300 MHz, DMSO- $d_6$ ):  $\delta$  10.88 (s, 1H, NH), 8.35–8.27 (m, 2H, ArH), 8.14 (d,  $J = 8.4$  Hz, 1H, ArH), 7.68–7.48 (m, 4H, ArH), 7.25 (t,  $J = 7.1$  Hz, 1H, ArH), 3.62–3.38 (m, 4H), 2.85–2.56 (m, 4H), 2.35 (s, 3H).  $^{13}\text{C}$  NMR (75 MHz, DMSO- $d_6$ ):  $\delta$  148.16, 145.83, 144.78, 136.78, 130.30, 129.90, 127.26, 127.02, 125.10, 124.28, 124.12, 122.11, 121.94, 120.20, 112.89, 56.31, 51.36, 46.99. MS (ESI):  $m/z = 317.2$  [100%, (M + H) $^+$ ]. HRMS calcd for  $\text{C}_{20}\text{H}_{21}\text{N}_4$  [M + H] $^+$  317.1761, found 317.1757. HPLC MS purity 98.05%.

**2-(4-(10H-Indolo[3,2-b]quinolin-11-yl)piperazin-1-yl)-N,N-dimethylethanamine (4).** Compound **4** was synthesized following the general procedure using **1** (1 g, 3.96 mmol) and 1-[2-(dimethylamino)ethyl]piperazine (5 mL) heated at 100 °C for 16 h to give 1.1 g (79%) of **4** as a yellow solid.  $^1\text{H}$  NMR (300 MHz, DMSO- $d_6$ ):  $\delta$  10.88 (s, 1H, NH), 8.38–8.27 (m, 2H, ArH), 8.14 (d,  $J = 8.4$  Hz, 1H, ArH), 7.69–7.49 (m, 4H, ArH), 7.25 (t,  $J = 7.8$  Hz, 1H, ArH), 3.58–3.44 (m, 4H), 2.90–2.65 (m, 4H), 2.62–2.53 (m, 2H),

2.48–2.40 (m, 2H), 2.19 (s, 6H).  $^{13}\text{C}$  NMR (75 MHz, DMSO- $d_6$ ):  $\delta$  148.17, 145.80, 144.83, 136.74, 130.28, 129.90, 127.25, 127.00, 125.09, 124.30, 124.13, 122.08, 121.93, 120.19, 112.92, 56.85, 56.11, 54.61, 51.49, 45.88. MS (ESI):  $m/z = 374.4$  [100%, (M + H) $^+$ ]. HRMS calcd for  $\text{C}_{23}\text{H}_{28}\text{N}_5$  [M + H] $^+$  374.2339, found 374.2334. HPLC MS purity 99.11%.

**4-(2-(4-(10H-Indolo[3,2-b]quinolin-11-yl)piperazin-1-yl)-ethyl)morpholine (5).** Compound **1** (200 mg, 0.79 mmol) was heated with 1.5 mL of 4-(2-(piperazin-1-yl)ethyl)morpholine at 110 °C for 24 h to obtain 294 mg (89%) of compound **5** as a solid.  $^1\text{H}$  NMR (300 MHz, DMSO- $d_6$ ):  $\delta$  10.86 (s, 1H, NH), 8.38–8.25 (m, 2H, ArH), 8.14 (d,  $J = 8.1$  Hz, 1H, ArH), 7.68–7.50 (m, 4H, ArH), 7.25 (t,  $J = 7.8$  Hz, 1H, ArH), 3.65–3.56 (m, 4H), 3.54–3.42 (m, 4H), 2.91–2.68 (m, 4H), 2.68–2.58 (m, 2H), 2.57–2.52 (m, 2H), 2.48–2.38 (m, 4H).  $^{13}\text{C}$  NMR (75 MHz, DMSO- $d_6$ ):  $\delta$  148.19, 145.83, 144.82, 136.73, 130.28, 129.91, 127.25, 126.99, 125.08, 124.29, 124.13, 122.12, 121.93, 120.19, 112.90, 67.10, 56.78, 56.16, 54.80, 54.61, 51.56. MS (ESI):  $m/z = 416.3$  [100%, (M + H) $^+$ ]. HRMS calcd for  $\text{C}_{25}\text{H}_{29}\text{N}_5\text{O}$  [M + H] $^+$  416.2445, found 416.2448. HPLC MS purity 100%.

**11-(4-(2-(Piperidin-1-yl)ethyl)piperazin-1-yl)-10H-indolo[3,2-b]quinoline (6).** Compound **6** was obtained from compound **1** (350 mg, 1.38 mmol) and 3 mL of 1-(2-(piperidin-1-yl)ethyl)piperazine heated at 120 °C for 12 h. Yield, 330 mg (57%).  $^1\text{H}$  NMR (300 MHz, DMSO- $d_6$ ):  $\delta$  10.87 (s, 1H, NH), 8.38–8.25 (m, 2H, ArH), 8.13 (d,  $J = 8.1$  Hz, 1H, ArH), 7.68–7.50 (m, 4H, ArH), 7.25 (t,  $J = 6.6$  Hz, 1H, ArH), 3.60–3.43 (m, 4H), 2.90–2.67 (m, 4H), 2.65–2.53 (m, 2H), 2.50–2.45 (m, 2H), 2.44–2.30 (m, 4H), 1.58–1.45 (m, 4H), 1.45–1.35 (m, 2H).  $^{13}\text{C}$  NMR (75 MHz, DMSO- $d_6$ ):  $\delta$  148.18, 145.83, 144.82, 136.74, 130.27, 129.90, 127.24, 126.99, 125.07, 124.29, 124.13, 122.11, 121.92, 120.18, 112.90, 57.03, 56.39, 55.26, 54.79, 51.55, 26.34, 24.82. MS (ESI):  $m/z = 414.5$  [100%, (M + H) $^+$ ]. HRMS calcd for  $\text{C}_{26}\text{H}_{31}\text{N}_5$  [M + H] $^+$  414.2652, found 414.2658. HPLC MS purity 100%.

**2-(4-(10H-Indolo[3,2-b]quinolin-11-yl)piperazin-1-yl)-N,N-diethylethanamine (7).** Compound **7** was synthesized in 92% yield (735 mg) from compound **1** (500 mg, 1.98 mmol) and 3 mL of *N,N*-diethyl-2-(piperazin-1-yl)ethanamine heated at 100 °C for 48 h.  $^1\text{H}$  NMR (500 MHz, DMSO- $d_6$ ):  $\delta$  10.90 (s, 1H, NH), 8.33 (d,  $J = 8.5$  Hz, 1H, ArH), 8.29 (d,  $J = 7.5$  Hz, 1H, ArH), 8.14 (d,  $J = 8.5$  Hz, 1H, ArH), 7.68–7.52 (m, 4H, ArH), 7.25 (t,  $J = 8.0$  Hz, 1H, ArH), 3.60–3.40 (m, 4H), 2.90–2.70 (m, 4H), 2.67–2.60 (m, 2H), 2.58–2.48 (m, 6H), 1.00 (t,  $J = 7.0$  Hz, 6H).  $^{13}\text{C}$  NMR (75 MHz, DMSO- $d_6$ ):  $\delta$  148.18, 145.83, 144.82, 136.75, 130.27, 129.90, 127.25, 126.98, 125.07, 124.30, 124.13, 122.12, 121.92, 120.18, 112.90, 57.12, 54.86, 51.57, 50.93, 47.64, 12.68. MS (ESI):  $m/z = 402.3$  [65%, (M + H) $^+$ ], 201.6 (100%). HRMS calcd for  $\text{C}_{25}\text{H}_{31}\text{N}_5$  [M + H] $^+$  402.2652, found 402.2645. HPLC MS purity 100%.

**3-(4-(10H-Indolo[3,2-b]quinolin-11-yl)piperazin-1-yl)-N,N-dimethylpropan-1-amine (8).** Compound **8** was synthesized from **1** (500 mg, 1.98 mmol) and *N,N*-dimethyl-3-(piperazin-1-yl)propan-1-amine (2 mL) heated at 110 °C for 24 h to obtain 633 mg (82%) of **8** as a yellow solid.  $^1\text{H}$  NMR (300 MHz, DMSO- $d_6$ ):  $\delta$  10.86 (s, 1H, NH), 8.40–8.25 (m, 2H, ArH), 8.14 (d,  $J = 8.4$  Hz, 1H, ArH), 7.68–7.50 (m, 4H, ArH), 7.25 (t,  $J = 7.4$  Hz, 1H, ArH), 3.60–3.45 (m, 4H), 2.85–2.60 (m, 4H), 2.47 (t,  $J = 7.2$  Hz, 2H), 2.29 (t,  $J = 7.2$  Hz, 2H), 2.16 (s, 6H), 1.73–1.58 (m, 2H).  $^{13}\text{C}$  NMR (75 MHz, DMSO- $d_6$ ):  $\delta$  148.18, 145.82, 144.82, 136.76, 130.28, 129.90, 127.24, 127.00, 125.08, 124.29, 124.14, 122.11, 121.93, 120.19, 112.90, 58.20, 57.11, 54.51, 51.59, 46.09, 25.41. MS (ESI):  $m/z = 388.3$  [100%, (M + H) $^+$ ]. HRMS calcd for  $\text{C}_{24}\text{H}_{29}\text{N}_5$  [M + H] $^+$  388.2496, found 388.2496. HPLC MS purity 99.08%.

**11-(4-(3-Methoxypropyl)piperazin-1-yl)-10H-indolo[3,2-b]quinoline (9).** Compound **9** was prepared from **1** (200 mg, 0.79 mmol) and 2 mL of 1-(3-methoxypropyl)piperazine heated at 110 °C overnight to obtain 264 mg (88%) of compound **9**.  $^1\text{H}$  NMR (500 MHz, DMSO- $d_6$ ):  $\delta$  10.87 (s, 1H, NH), 8.33 (d,  $J = 8.5$  Hz, 1H, ArH), 8.30 (d,  $J = 7.5$  Hz, 1H, ArH), 8.14 (d,  $J = 8.5$  Hz, 1H, ArH), 7.65–7.58 (m, 3H, ArH), 7.55 (t,  $J = 7.5$  Hz, 1H, ArH), 7.26 (t,  $J = 8.0$  Hz, 1H, ArH), 3.60–3.46 (m, 4H), 3.43 (t,  $J = 6.5$  Hz, 2H), 3.27 (s, 3H),

2.90–2.60 (m, 4H), 2.56–2.50 (m, 2H), 1.90–1.74 (m, 2H). <sup>13</sup>C NMR (125 MHz, DMSO-*d*<sub>6</sub>): δ 147.81, 145.44, 144.44, 136.36, 129.90, 129.52, 126.87, 126.62, 124.71, 123.92, 123.75, 121.73, 121.54, 119.81, 112.52, 70.78, 58.39, 55.57, 54.11, 51.20, 27.11. MS (ESI): *m/z* = 375.2 [40%, (M + H)<sup>+</sup>]. HRMS calcd for C<sub>23</sub>H<sub>26</sub>N<sub>4</sub>O [M + H]<sup>+</sup> 375.2179, found 375.2178. HPLC MS purity 99.4%.

**3-(4-(10H-Indolo[3,2-*b*]quinolin-11-yl)piperazin-1-yl)propan-1-ol (10).** Compound 10 was prepared from 1 (500 mg, 1.98 mmol) and 1-(3-hydroxy propyl)piperazine (1.5 mL) by heating at 100 °C for 24 h to obtain 620 mg (86%) of 10 as a yellow solid. <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>): δ 10.88 (br s, 1H, NH), 8.42–8.25 (m, 2H, ArH), 8.14 (d, *J* = 8.1 Hz, 1H, ArH), 7.76–7.50 (m, 4H, ArH), 7.25 (t, *J* = 6.4 Hz, 1H, ArH), 5.10–4.30 (m, OH), 3.70–3.40 (m, 6H), 2.92–2.63 (m, 4H), 2.58–2.50 (m, 2H), 1.80–1.58 (m, 2H). <sup>13</sup>C NMR (75 MHz, DMSO-*d*<sub>6</sub>): δ 148.17, 145.80, 144.81, 136.73, 130.30, 129.89, 127.23, 127.02, 125.10, 124.27, 124.14, 122.09, 121.94, 120.20, 112.89, 60.29, 56.25, 54.53, 51.55, 30.54. MS (ESI): *m/z* = 361.2 [100%, (M + H)<sup>+</sup>]. HRMS calcd for C<sub>22</sub>H<sub>25</sub>N<sub>4</sub>O [M + H]<sup>+</sup> 361.2023, found 361.2022. HPLC MS purity 100%.

***N,N*-Dimethyl-3-(11-(4-methylpiperazin-1-yl)-10H-indolo[3,2-*b*]quinolin-10-yl)propan-1-amine (11).** To a solution of compound 3 (100 mg, 0.31 mmol) in DMF at 0 °C, sodium hydride (60% in mineral oil) (22 mg, 0.93 mmol) was added. After the reaction mixture was stirred for 1 h, 3-chloro-*N,N*-dimethylpropan-1-amine hydrochloride (98 mg, 0.62 mmol) was added. The stirring was continued for 3 days at room temperature. After completion of the reaction, the mixture was poured into ice-cold water. The reaction was quenched with saturated ammonium chloride solution, and extraction was with CHCl<sub>3</sub> (2 × 50 mL). The organic layer was dried over anhydrous MgSO<sub>4</sub> and concentrated to give the product. This crude product was purified by silica gel column chromatography, eluting with chloroform to give 93 mg (73%) of 11 as a light yellow solid. <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>): δ 8.42–8.31 (m, 2H, ArH), 8.23 (d, *J* = 8.4 Hz, 1H, ArH), 7.75–7.54 (m, 4H, ArH), 7.36–7.28 (m, 1H, ArH), 4.84 (t, *J* = 6.9 Hz, 2H), 3.68–3.18 (m, 4H), 3.00–2.52 (m, 4H), 2.46–2.30 (m, 5H), 2.19 (s, 6H), 2.03–1.87 (m, 2H). <sup>13</sup>C NMR (75 MHz, DMSO-*d*<sub>6</sub>): δ 148.27, 145.92, 145.59, 136.49, 130.90, 130.83, 129.94, 127.56, 126.64, 125.22, 124.79, 122.00, 121.91, 120.61, 110.51, 57.11, 55.90, 51.36, 47.22, 46.03, 42.95, 28.16. MS (ESI): *m/z* = 402.4 [100%, (M + H)<sup>+</sup>]. HRMS calcd for C<sub>25</sub>H<sub>32</sub>N<sub>5</sub> [M + H]<sup>+</sup> 402.2652, found 402.2647. HPLC MS purity 100%.

**3-(11-(4-(2-(Dimethylamino)ethyl)piperazin-1-yl)-10H-indolo[3,2-*b*]quinolin-11-yl)-*N,N*-dimethylpropan-1-amine (12).** Compound 4 (500 mg, 1.3 mmol) was reacted with sodium hydride (60% in mineral oil) (288 mg, 12 mmol) and 3-chloro-*N,N*-dimethylpropan-1-amine hydrochloride (612 mg, 3.9 mmol) to give compound 12 (523 mg, 85%) as a solid. <sup>1</sup>H NMR (300 MHz, CDCl<sub>3</sub>): δ 8.49 (d, *J* = 7.8 Hz, 1H, ArH), 8.42–8.28 (m, 2H, ArH), 7.68–7.55 (m, 2H, ArH), 7.52–7.41 (m, 2H, ArH), 7.29 (t, *J* = 7.5 Hz, 1H, ArH), 4.82 (t, *J* = 6.3 Hz, 2H), 4.45–3.65 (m, 2H), 3.60–2.80 (m, 4H), 2.75–2.50 (m, 4H), 2.50–2.29 (m, 10H), 2.28–2.10 (m, 6H), 2.06–1.90 (m, 2H). <sup>13</sup>C NMR (75 MHz, CDCl<sub>3</sub>): δ 148.71, 146.01, 145.46, 135.92, 130.76, 130.13, 129.97, 127.49, 126.10, 124.47, 124.44, 122.18, 120.12, 109.43, 57.51, 57.40, 57.24, 54.63, 51.45, 46.32, 46.03, 42.82, 28.40. MS (ESI): *m/z* = 459.2 [100%, (M + H)<sup>+</sup>]. HRMS calcd for C<sub>28</sub>H<sub>39</sub>N<sub>6</sub> [M + H]<sup>+</sup> 459.3231, found 459.3231. HPLC MS purity 100%.

**2-(4-(2-(3,4-Difluorophenyl)-10H-indolo[3,2-*b*]quinolin-11-yl)piperazin-1-yl)-*N,N*-dimethylethanamine (13).** A mixture of 2-(4-(2-bromo-10H-indolo[3,2-*b*]quinolin-11-yl)piperazin-1-yl)-*N,N*-dimethylethanamine (200 mg, 0.44 mmol), 3,4-difluorophenylboronic acid (104 mg, 0.66 mmol), sodium carbonate (141 mg, 1.32 mmol), and Pd(dppf)<sub>2</sub>Cl<sub>2</sub> (10 mol %) in 1,2-dimethoxyethane/water (4:1) was heated at 90 °C for 6 h. After completion of the reaction, the reaction mixture was cooled to room temperature, poured into ice-cold water, and extracted with ethyl acetate. The organic layer was separated, dried over anhydrous MgSO<sub>4</sub>, and evaporated under vacuum to give the product. The crude product was purified by silica gel column chromatography using 5% MeOH in CHCl<sub>3</sub> as an eluent to afford 186 mg (86%) of pure product 13 as a yellow solid. <sup>1</sup>H NMR

(300 MHz, DMSO-*d*<sub>6</sub>): δ 11.00 (s, 1H, NH), 8.51 (d, *J* = 2.1 Hz, 1H, ArH), 8.30 (d, *J* = 7.5 Hz, 1H, ArH), 8.22 (d, *J* = 8.7 Hz, 1H, ArH), 7.98–7.84 (m, 2H, ArH), 7.70–7.56 (m, 4H, ArH), 7.27 (t, *J* = 7.2 Hz, 1H, ArH), 3.63–3.48 (m, 4H), 2.93–2.65 (m, 8H), 2.44 (br s, 6H). <sup>13</sup>C NMR (75 MHz, DMSO-*d*<sub>6</sub>): δ 148.44, 148.27, 145.35, 144.93, 139.12, 137.07, 134.52, 130.75, 130.47, 127.55, 126.15, 124.58, 124.22, 122.08, 121.99, 121.93, 120.35, 119.16, 118.94, 116.93, 116.70, 112.99, 56.35, 55.46, 54.58, 54.18, 51.54, 45.44. MS (ESI): *m/z* = 486.2 [100%, (M + H)<sup>+</sup>]. HRMS calcd for C<sub>29</sub>H<sub>29</sub>F<sub>2</sub>N<sub>5</sub> [M + H]<sup>+</sup> 486.2463, found 486.2465. HPLC MS purity 100%.

***N,N*-Dimethyl-2-(4-(2-(4-(2-morpholinoethyl)phenyl)-10H-indolo[3,2-*b*]quinolin-11-yl)piperazin-1-yl)ethanamine (14).** Compound 14 was prepared following a procedure similar to that for compound 13, using 2-(4-(2-bromo-10H-indolo[3,2-*b*]quinolin-11-yl)piperazin-1-yl)-*N,N*-dimethylethanamine (200 mg, 0.44 mmol), 4-(2-morpholinoethyl)phenylboronic acid (156 mg, 0.66 mmol), sodium carbonate (141 mg, 1.32 mmol), and Pd(dppf)<sub>2</sub>Cl<sub>2</sub> (10 mol %) to give 14 (217 mg, 87%) as a yellow solid. <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>): δ 10.89 (s, 1H, NH), 8.51 (d, *J* = 1.2 Hz, 1H, ArH), 8.29 (d, *J* = 7.8 Hz, 1H, ArH), 8.20 (d, *J* = 8.7 Hz, 1H, ArH), 7.92 (dd, *J* = 8.7 Hz, 1H, ArH), 7.72 (d, *J* = 8.1 Hz, 2H, ArH), 7.68–7.56 (m, 2H, ArH), 7.40 (d, *J* = 8.1 Hz, 2H, ArH), 7.26 (t, *J* = 7.5 Hz, 1H, ArH), 3.66–3.58 (m, 4H), 3.57–3.46 (m, 4H), 2.87–2.74 (m, 4H), 2.66–2.56 (m, 4H), 2.55–2.51 (m, 4H), 2.48–2.40 (m, 4H), 2.24 (br s, 6H). <sup>13</sup>C NMR (75 MHz, DMSO-*d*<sub>6</sub>): δ 148.17, 145.20, 144.85, 140.58, 138.97, 136.87, 136.57, 130.58, 130.36, 127.65, 127.50, 126.26, 124.36, 122.17, 121.92, 121.18, 120.26, 112.93, 67.07, 60.85, 57.43, 56.67, 54.86, 54.15, 51.61, 46.27, 32.87. MS (ESI): *m/z* = 563.4 [100%, (M + H)<sup>+</sup>]. HRMS calcd for C<sub>35</sub>H<sub>42</sub>N<sub>6</sub>O [M + H]<sup>+</sup> 563.3492, found 563.3491. HPLC MS purity 98.17%.

***N,N*-Dimethyl-2-(4-(2-(pyridin-4-yl)-10H-indolo[3,2-*b*]quinolin-11-yl)piperazin-1-yl)ethanamine (15).** Compound 15 was synthesized following a procedure similar to that for compound 13, using 2-(4-(2-bromo-10H-indolo[3,2-*b*]quinolin-11-yl)piperazin-1-yl)-*N,N*-dimethylethanamine (100 mg, 0.22 mmol), pyridin-4-ylboronic acid (40.5 mg, 0.33 mmol), sodium carbonate (69.9 mg, 0.66 mmol), and Pd(dppf)<sub>2</sub>Cl<sub>2</sub> (10 mol %) to obtain 93 mg (93%) of compound 15 as a yellow solid. <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>): δ 11.10 (s, 1H, NH), 8.72 (d, *J* = 5.1 Hz, 2H, ArH), 8.68 (s, 1H, ArH), 8.31 (d, *J* = 7.5 Hz, 1H, ArH), 8.27 (d, *J* = 9.0 Hz, 1H, ArH), 8.05 (d, *J* = 8.7 Hz, 1H, ArH), 7.86 (d, *J* = 5.1 Hz, 2H, ArH), 7.72–7.58 (m, 2H, ArH), 7.28 (t, *J* = 6.9 Hz, 1H, ArH), 3.70–3.55 (m, 4H), 3.26–3.13 (m, 2H), 2.98–2.82 (m, 6H), 2.76 (s, 6H). <sup>13</sup>C NMR (75 MHz, DMSO-*d*<sub>6</sub>): δ 151.28, 148.86, 148.14, 145.93, 145.02, 137.37, 133.47, 131.01, 130.63, 127.47, 125.60, 124.11, 122.55, 122.27, 122.09, 122.04, 120.44, 113.03, 54.53, 54.22, 53.52, 51.52, 44.09. MS (ESI): *m/z* = 451.3 [70%, (M + H)<sup>+</sup>]. HRMS calcd for C<sub>28</sub>H<sub>30</sub>N<sub>6</sub> [M + H]<sup>+</sup> 451.2605, found 451.2602. HPLC MS purity 100%.

**4-(4-(10H-Indolo[3,2-*b*]quinolin-11-yl)phenethyl)morpholine (16).** A mixture of 1 (200 mg, 0.79 mmol), 4-(2-morpholinoethyl)phenylboronic acid (279 mg, 1.19 mmol), sodium carbonate (252 mg, 2.38 mmol), and Pd(dppf)<sub>2</sub>Cl<sub>2</sub> (10 mol %) in 1,2-dimethoxyethane/water (3:1) was heated at 90 °C overnight. After completion of the reaction, the reaction mixture was cooled to room temperature, poured into ice-cold water, and extracted with ethyl acetate (2 × 100 mL). The organic layer was separated, dried over anhydrous MgSO<sub>4</sub>, and evaporated under vacuum to give the crude product. The crude product was purified by silica gel column chromatography using 2% MeOH in CHCl<sub>3</sub> as an eluent to afford (235 mg, 72%) of pure product 16. <sup>1</sup>H NMR (300 MHz, DMSO-*d*<sub>6</sub>): δ 10.97 (s, 1H, NH), 8.37 (d, *J* = 7.5 Hz, 1H, ArH), 8.25 (d, *J* = 8.7 Hz, 1H, ArH), 7.80 (d, *J* = 8.1 Hz, 1H, ArH), 7.67 (t, *J* = 8.1 Hz, 1H, ArH), 7.63–7.42 (m, 7H, ArH), 7.29 (t, *J* = 7.2 Hz, 1H, ArH), 3.74–3.50 (m, 4H), 3.02–2.83 (m, 2H), 2.80–2.63 (m, 2H), 2.60–2.50 (m, 4H). <sup>13</sup>C NMR (75 MHz, DMSO-*d*<sub>6</sub>): δ 146.33, 145.30, 144.66, 141.52, 132.44, 131.26, 131.03, 130.47, 130.27, 130.09, 126.74, 126.16, 125.93, 125.73, 125.23, 122.15, 122.02, 120.30, 112.78, 67.06, 60.80, 54.16, 33.08. MS (ESI): *m/z* = 408.3 [40%, (M + H)<sup>+</sup>]. HRMS calcd for C<sub>27</sub>H<sub>25</sub>N<sub>3</sub>O [M + H]<sup>+</sup> 408.2070, found 408.2071. HPLC MS purity 98.2%.

Table 1. Oligonucleotide Sequences (5′–3′)

d(AT)	ATATATATATATATATATATATATATATATATATAT
d(GC)	GCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG
c-MYC G4	TGGGGAGGGTGGGGAGGGTGGGGAAAGG
Bcl-2 G4	AGGGGCGGGCGCGGGAGGAAGGGGGCGGGAGCGGGGGC
PDGFA G4	GGAGGCGGGGGGGGGGGCGGGGGCGGGGGCGGGGGAGGGGGCGGGG
PDGFR- $\beta$ G4	GCTGGGAGAAGGGGGGGCGGGGGCAGGGAGGGTGGGA
hTERT G4	GGGGAGGGGCTGGGAGGGCCCCGAGGGGGCTGGGCCGGGGACCCGGGAGGGGTGGGACGGGGCGGGG
HIF-1 $\alpha$ G4	GCGCGGGAGGGGAGAGGGGGCGGGAGCGCG

**Thermal Stabilization.** Spectral characteristics and thermal stability of the c-MYC G-quadruplex (Table 1) in the absence and presence of compounds were recorded on a J-810 spectropolarimeter (Jasco, Easton, MD). Briefly, the c-MYC G-quadruplex was diluted to 5  $\mu$ M in a 50 mM Tris-HCl solution (pH 7.4), incubated at 25 °C for 10 min, heated to 95 °C for 10 min, and allowed to cool in the air over ~30 min before the addition of compound at 1 equiv. Positive molecular ellipticity at the parallel G-quadruplex peak (262 nm) was confirmed by spectral examination before each mixture was heated from 5 to 95 °C at 2 °C/min. Molecular ellipticity as a function of temperature was used to calculate a  $T_m$  (the temperature at which 50% of the formed higher order DNA structure was melted) for each condition using GraphPad Prism software and a nonlinear regression model with variable slope.  $\Delta T_m$  values were calculated as  $T_{m\text{-compound}} - T_{m\text{-control}}$ .

**Competition Dialysis.** All DNA (75  $\mu$ M) were prepared for the experiment in 50 mM Tris-HCl and 25 mM KCl solution. Duplex DNA sequences were annealed by incubating complementary strands together for 10 min at room temperature, heating at 95 °C for 10 min, and slowly (1 °C/min) decreasing temperature to the  $T_m$ , holding at that temperature (°C) for 10 min, and slowly (1 °C/min) decreasing temperature to room temperature. G-Quadruplexes were formed by heating to 95 °C for 10 min and then rapidly cooling (in air) to room temperature. All topologies were confirmed by circular dichroism before continuing with the assay. An amount of 250  $\mu$ L of DNA, in their formed and confirmed topologies, was added to prewet minidialysis units in a flotation device, placed in a beaker containing 250 mL of BPES buffer (6 mM Na<sub>2</sub>HPO<sub>4</sub>, 2 mM NaH<sub>2</sub>PO<sub>4</sub>, 1 mM disodium EDTA) with 25 mM KCl plus 2  $\mu$ M compound 4 or 12, and placed in a cold room on a stirring device for 48 h. Following the published protocol,<sup>15</sup> the concentration of compound bound to DNA was determined by measuring UV absorbance and using a standard absorbance curve. Binding per putative site value was calculated based on two binding sites per G-quadruplex and intercalation between every other base pair in dsDNA. The experiment was performed in duplicate.

**Cellular Viability.** Cells were maintained in a 37 °C, 5% CO<sub>2</sub> incubator, in exponential growth, for the duration of experimentation in either DMEM (HCT-116) or RPMI-1640 (Raji and CA46) medium supplemented with 10% fetal bovine serum and 1 $\times$  penicillin/streptomycin. Cytotoxicity of compounds in cells (ATCC, Manassas, VA) was determined using the MTS assay<sup>16</sup> with compound diluted over a 5–6 log range in 0.5 log steps. IC<sub>50</sub> values were determined with GraphPad Prism software using nonlinear regression modeling. The experiments were performed with biological triplicates.

**Transcriptional Regulation.** CA46 or HCT-116 cells were seeded in a T-25 flask or six-well plates, respectively, at a density of (0.5–1.0)  $\times$  10<sup>6</sup> cells/mL in 1 mL overnight before incubation with the appropriate IC<sub>50</sub> concentration of 4, 12, or quindoline-i for the prescribed times. Cells were washed with PBS (2 $\times$ ) and lysed, and RNA was isolated with an RNeasy Mini kit (Qiagen, Valencia, CA) using the Qiacube automated system. Then 200–500 ng of cDNA was reverse transcribed with the Quantitect cDNA synthesis kit (Qiagen) prior to quantitative real-time PCR detection on the BioRad MyIQ thermocycler. FAM-labeled TaqMan primers were obtained from Applied Biosystems (Carlsbad, CA) for GAPDH, HIF-1 $\alpha$ , VEGFA, PDGFR- $\beta$ , hTERT, and c-MYC (exon 1 and exon 2).<sup>14</sup> The  $\Delta\Delta Ct$  method was used to calculate changes in expression, normalized to the appropriate time-matched DMSO vehicle-treated control cells. Experi-

ments were performed in at least duplicates, with duplicate measurements within each qPCR reaction; two-tailed unpaired Student's *t* tests were utilized to determine statistical significance.

## RESULTS AND DISCUSSION

**Chemistry.** Quindoline analogues (Table 2) were synthesized from 11-chloroquindoline 1 (Schemes 1 and 4) or substituted 11-chloroquindoline (Schemes 2 and 3). Anthranilic acid and aniline were used in a multistep procedure to synthesize 11-chloroquindoline as reported in literature.<sup>17,18</sup>

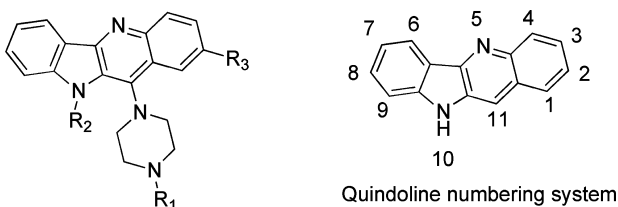
**Synthesis of 11-Substituted Quindoline Analogues.** 11-Substituted quindoline analogues 2–10 were synthesized from 11-chloroquindoline and commercially available piperazines (Schemes 1–4). Reactions were performed by heating two reactants together under neat conditions. Products were isolated in 60–80% yields. It was expected that the addition of a piperazine ring would increase the binding to G-quadruplex. The piperazine N4 atom also provides a site for the addition of groups with varying properties.

**Disubstituted Quindoline Analogues.** A second set of quindoline analogues was prepared using compound 4 as a lead molecule and substituting at two different positions: indole NH and C2 on the quindoline ring (Scheme 2). It was expected that substitution of a second side chain would lead to an increase in the G-quadruplex binding and cellular activity of the analogues. Compound 12 was prepared by alkylation at indole –NH of compound 4. Compounds 13–15 were synthesized using a palladium-catalyzed Suzuki reaction (Scheme 3). A 3-bromo analogue was reacted with corresponding boronic acids to obtain compounds 13–15 in ~90% yield.

**Cell-Free and Cellular Evaluation. Evaluation of Monosubstituted Quindoline Analogues.** *Thermal Stabilization.* Quindoline analogues substituted at the R<sub>1</sub> position were tested for their ability to thermally stabilize the c-MYC G-quadruplex, resulting in an increase in melting temperature ( $\Delta T_m$ ) (Table 3). This increase in  $T_m$  value for each compound can be directly correlated with the strength of binding with the G-quadruplex structure.<sup>19,20</sup>  $\Delta T_m$  values ranged from 1.9 to 11.4 °C. The lead compound, quindoline-i, shows a  $\Delta T_m$  of 5.2 °C, while compound 2 (11-piperazinylquindoline) shows a moderate  $\Delta T_m$  at 6.7 °C. Compounds 4, 5, and 9 demonstrated comparable thermal stabilization of the c-MYC G-quadruplex, whereas compounds 3, 10, 11, and 16 were less efficacious, and compounds 6–8 demonstrated higher  $\Delta T_m$  in the range 8.8–11.4 °C.

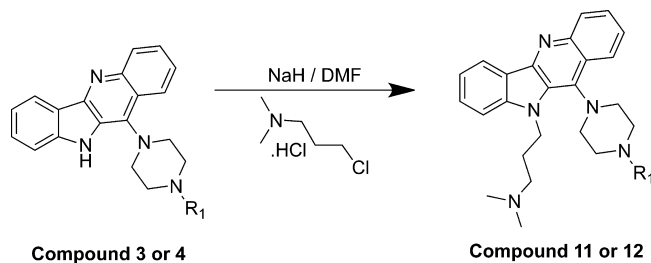
*Cellular Cytotoxicity.* Both the colon cancer cell line HCT-116 and the Burkitt's lymphoma cell line Raji exhibit high levels of c-MYC expression. As an initial indication of both intracellular accumulation of the compound and potential c-MYC down-regulation, cytotoxicity in each of these two cell lines was measured after 96 h of exposure (Table 3). With the exception of compound 16, all compounds demonstrated

Table 2. Structures for Substituted Quindoline Analogues



Compound	R <sub>1</sub>	R <sub>2</sub>	R <sub>3</sub>
2	-H	-H	-H
3	-CH <sub>3</sub>	-H	-H
4		-H	-H
5		-H	-H
6		-H	-H
7		-H	-H
8		-H	-H
9		-H	-H
10		-H	-H
11	-CH <sub>3</sub>		-H
12			-H
13		-H	
14		-H	
15		-H	
16			
Quindoline-i			

Scheme 2. Synthesis of compounds 11 and 12



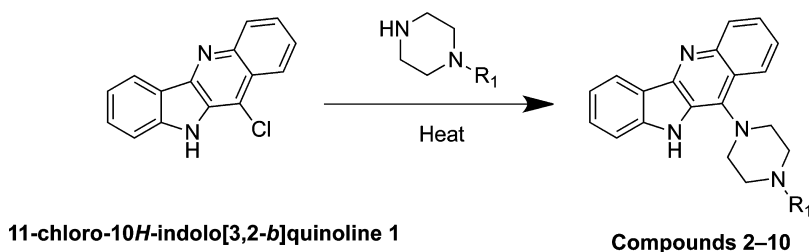
greater than 10  $\mu\text{M}$  potency in the HCT-116 cell line, with 96 h  $\text{IC}_{50}$  values ranging from 0.5 to 9.1  $\mu\text{M}$  and compounds 4, 6, and 8 showing high nanomolar efficacy. Raji cells, on the other hand, generally have higher  $\text{IC}_{50}$  values, ranging from 2.33 to 100  $\mu\text{M}$ . The most potent analogue tested in these cells is compound 4. Thus, considering the moderate thermal stabilization by this compound and the cellular potency, second-generation molecules were derived from compound 4.

**Evaluation of Disubstituted Compounds. Thermal Stabilization.** In comparison to compound 4, all disubstituted quindoline analogues demonstrated comparable or improved thermal stabilization of the parallel c-MYC G-quadruplex structure (Table 4). For compounds 12–15,  $\Delta T_m$  values ranged from 6.5 to 16.5  $^{\circ}\text{C}$ , with compound 12 showing the greatest stabilization ( $\Delta T_m = 16.5$   $^{\circ}\text{C}$ ). This compound is better when compared to both compounds 4 and 11, since it is a combination of the two side chains, and there is apparent cooperativity with the double substitutions that is greater than either alone ( $\Delta T_m$  of 6.7 and 2.8  $^{\circ}\text{C}$ , respectively).

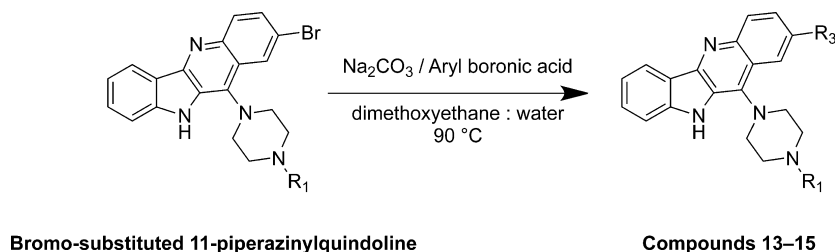
**Cellular Cytotoxicity.** Each of the four double-substituted compounds displayed a greater than 3-fold decrease in potency in the HCT-116 colon cancer cell line, with compound 15 showing the least potency at an  $\text{IC}_{50}$  (96 h) of 20.9  $\mu\text{M}$  (Table 4). Accordingly, this compound showed poor efficacy in the Raji lymphoma cell line, whereas the other three compounds showed moderate activity with 96 h  $\text{IC}_{50}$  values of 3.1–10.8  $\mu\text{M}$ . Again, the best activity is seen with compound 12, with a 96 h  $\text{IC}_{50}$  of 3.4 and 3.1  $\mu\text{M}$  in HCT-116 and Raji cells, respectively. Thus, compound 12 was selected as the disubstituted quindoline analogue for further examination.

**Comparison of a Monosubstituted Compound 4 with the Disubstituted Compound 12. Competition Dialysis.** Competition dialysis shows that compound 4 binds with higher affinity to G-quadruplexes than compound 12. Competition dialysis results on compounds 4 and 12 are shown in Figure 1. Both compounds bind to folded G-quadruplex structures with greater affinity than double-stranded DNA (dGdC or dAdT). Overall, compound 4 showed higher binding to G-quadruplex structures than compound 12, although inclusion of a second substitution appeared to increase G-quadruplex selectivity but

Scheme 1. Synthesis of compounds 2–10



Scheme 3. Synthesis of compounds 13–15



Scheme 4. Synthesis of compound 16

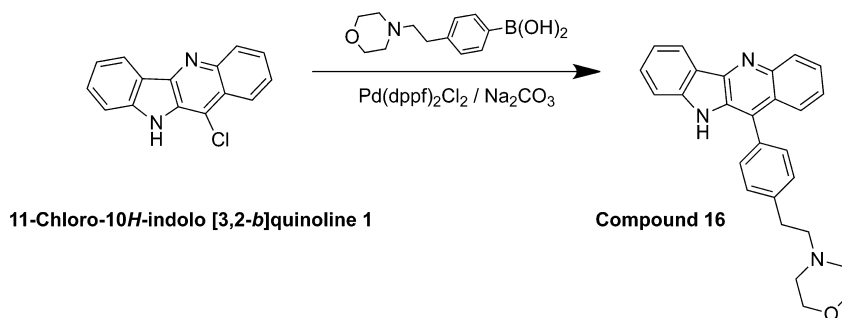


Table 3. Melting Temperature and Cytotoxicity of Single-Substituted Compounds against HCT-116 (Colon) and Raji (Lymphoma) Cell Lines

compd	$\Delta T_m$ in °C (1 equiv) <sup>a</sup>	96 h IC <sub>50</sub> ( $\mu$ M)	
		HCT-116	Raji
quindoline-i	5.2	0.11	0.70
2	6.7	4.3	33.9
3	3.0	3.5	5.44
4	6.7	0.97	2.33
5	5.9	9.1	52.9
6	8.8	0.5	56.1
7	10.3	3.4	26
8	11.4	0.9	11.8
9	6.3	6.3	21.3
10	1.3	2.4	>100
11	2.8	2.98	7.03
16	1.9	>100	>100

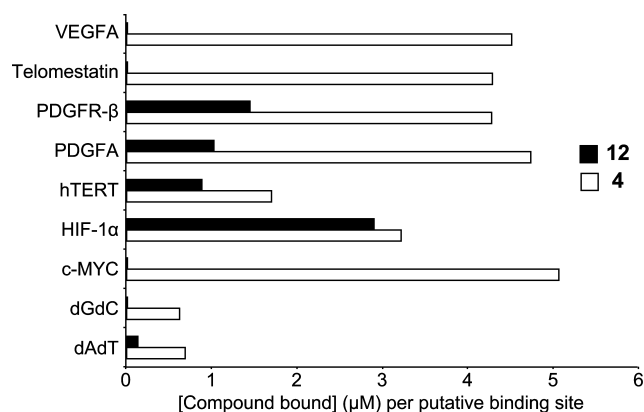
<sup>a</sup>Increase in melting temperature of c-MYC G-quadruplex in the presence of 1 equiv of ligand (DNA/ligand = 1:1).

Table 4. Melting Temperature and Cytotoxicity of Double-Substituted Compounds against HCT-116 (Colon) and Raji (Lymphoma) Cell Lines

compd	$\Delta T_m$ in °C (1 equiv) <sup>a</sup>	96 h IC <sub>50</sub> ( $\mu$ M)	
		HCT-116	Raji
4	6.7	0.97	2.33
12	16.5	3.4	3.1
13	6.5	3.7	5.8
14	13.1	3.8	10.8
15	8.6	20.9	45

<sup>a</sup>Increase in melting temperature of c-MYC G-quadruplex in the presence of 1 equiv of ligand (DNA/ligand = 1:1).

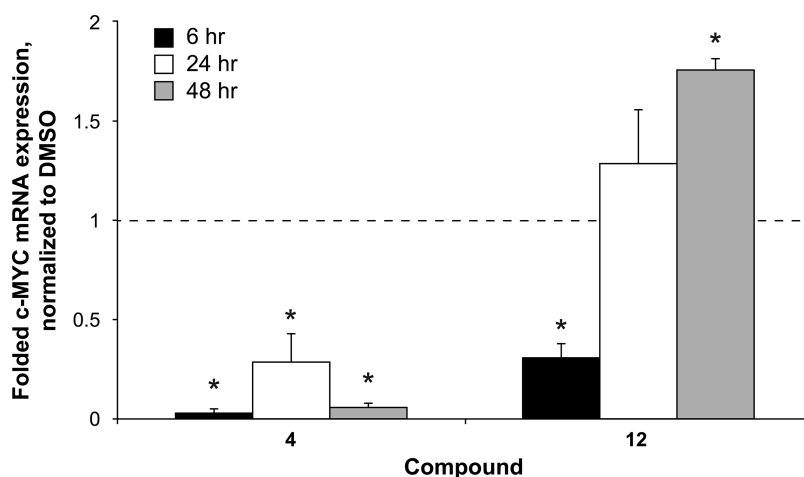
in this case not for the c-MYC G-quadruplex structure. Overall, compound 12 shows the highest binding affinity for the HIF-1 $\alpha$  G-quadruplex, which has a parallel structure similar to that of



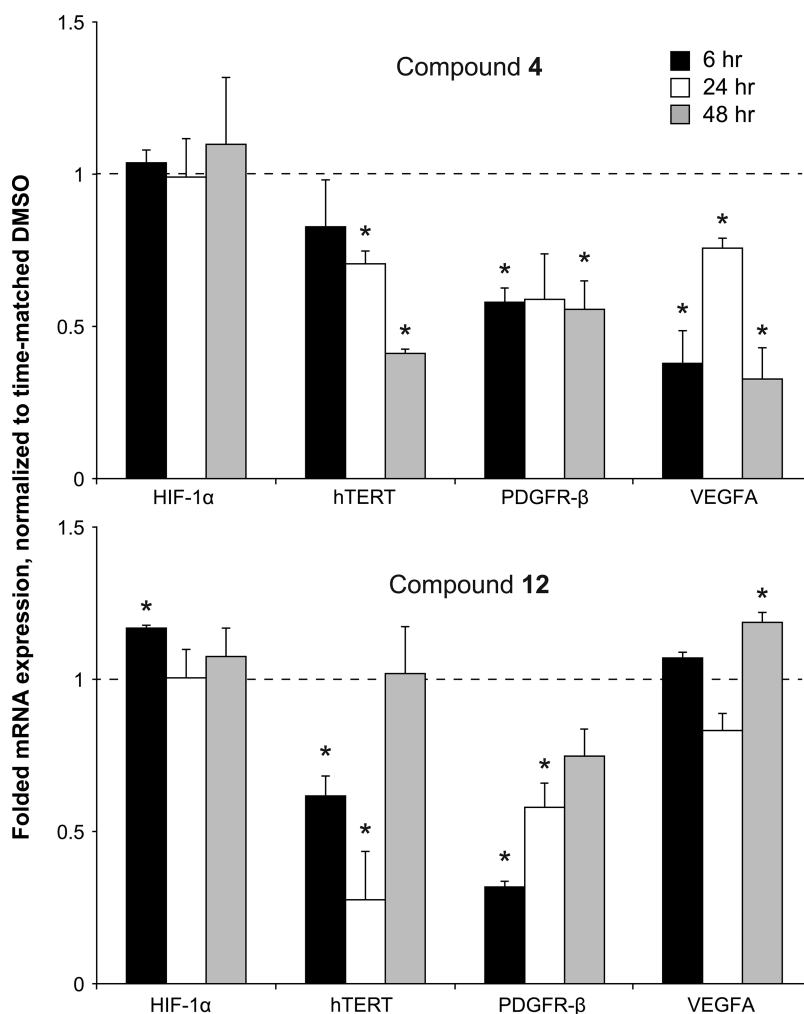
**Figure 1.** Affinity of compounds 4 and 12 to various DNA topologies. DNA structures (G-quadruplex or dsDNA) were induced as described, placed in individual dialysis tubes, and incubated with 2  $\mu$ M compound for 48 h. [Compound bound] was determined for each binding site of the relevant structure. Compound 4 (white bars) demonstrates promiscuous binding to G-quadruplex structures with the strongest affinity for c-MYC, whereas compound 12 (black bars) shows notable binding to the parallel HIF-1 $\alpha$  structure and minor affinity toward the more complicated hTERT and PDGFR- $\beta$  formations.

the c-MYC G-quadruplex, whereas compound 4 shows more promiscuous binding properties, with the highest affinity for the c-MYC G-quadruplex.

**Transcriptional Regulation.** Compounds 4 and 12 were examined for their effect on transcriptional regulation of c-MYC in the HCT-116 colon cancer cell line (Figure 2). By use of their respective 24 h IC<sub>50</sub> doses (23 and 31  $\mu$ M), mRNA was quantified at 6, 24, and 48 h post-treatment, and gene expression was normalized to the DMSO vehicle control. At all time points, compound 4 significantly lowered c-MYC expression, whereas compound 12 demonstrated an intriguing pattern of modulation of c-MYC expression with a significant down-regulation early (i.e., 6 h), which was reversed to a significant increase by 48 h.



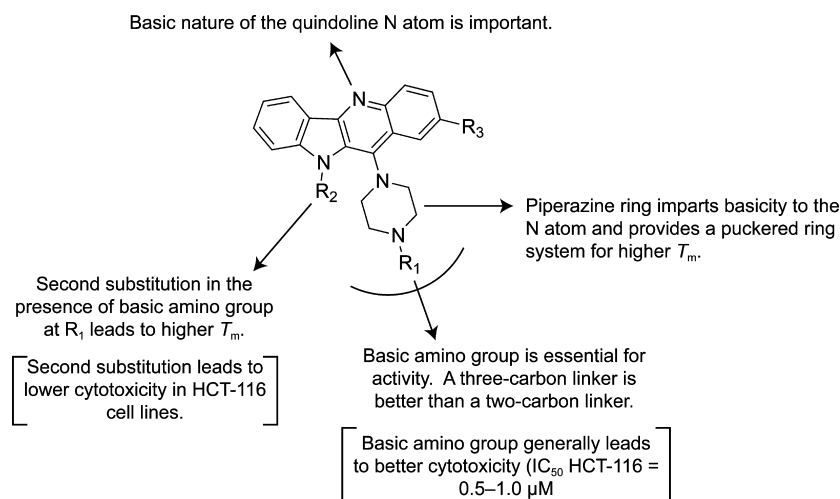
**Figure 2.** Transcriptional regulation of c-MYC by compounds 4 and 12 in HCT-116 colon cancer cells. Cells were exposed for a day to their respective 24 h  $IC_{50}$  concentrations of 23 and 31  $\mu$ M and examined for transcriptional regulation of c-MYC over time, normalized to GAPDH, and compared to the DMSO-vehicle control (6–48 h). Expression was decreased early by both compounds 4 and 12, but a sustained decrease was only noted with compound 4: (\*)  $p < 0.05$  as compared to time-matched vehicle control.



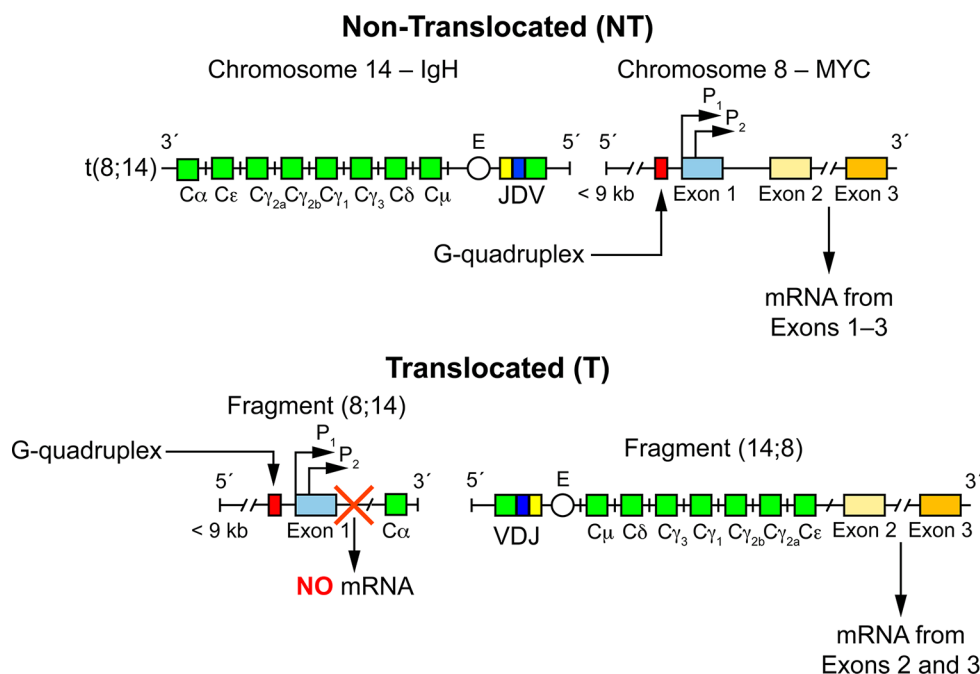
**Figure 3.** Transcriptional regulation of different genes by compounds 4 (top) and 12 (bottom) in HCT-116 colon cancer cells. Cells were exposed for a day to their respective 24 h  $IC_{50}$  concentrations and examined at 6, 24, and 48 h for transcriptional regulation by qPCR: (\*)  $p < 0.05$  as compared to time-matched vehicle control.

Compounds 4 and 12 showed nonselective binding to the different G-quadruplex structures, as shown in the competition dialysis experiments (Figure 1). To correlate the cytotoxicity

noted with compound 4 with alterations in c-MYC expression and to elucidate potential mechanisms of cell killing with compound 12, both compounds were also examined for their



**Figure 4.** Schematic depiction of structure–activity relationships for the quindoline analogues. Text in brackets is a description of the structure–cytotoxicity (HCT-116) relationships for the quindoline analogues.

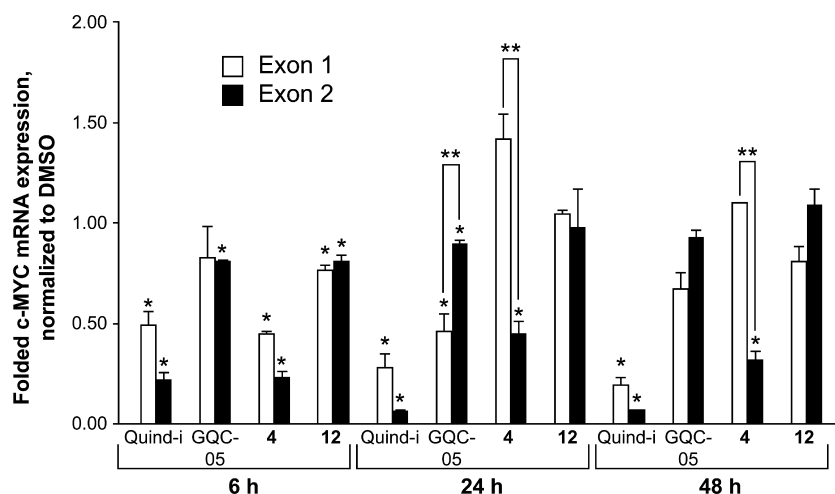


**Figure 5.** Exon-specific expression in CA46 cells based on translocation events. (A) Because of the reciprocal translocation between chromosomes 8 and 14, there are varying resultant c-MYC mRNAs produced. The NT products are normal, with a functional c-MYC under the control of a G-quadruplex, whereas the functional c-MYC produced from the fragment (14;8) on the T allele lacks G-quadruplex-mediated control. The G-quadruplex was removed, along with exon 1, and produces no known product from the fragment (8;14). Measurements of mRNAs containing exon 1 will mirror the NT allele. mRNAs containing exon 2 will show both the T and the NT products. This figure and caption are reproduced in accordance with the guidelines of the American Society for Biochemistry and Molecular Biology and were originally published in *Journal of Biological Chemistry* (Brown, R. V.; Danford, F. L.; Gokhale, V.; Hurley, L. H.; Brooks, T. A. Demonstration that drug-targeted down-regulation of MYC in non-Hodgkins lymphoma is directly mediated through the promoter G-quadruplex. *J. Biol. Chem.* 2011, 286, 41018–41027<sup>14</sup>). Copyright 2011, the American Society for Biochemistry and Molecular Biology.

effect on transcription of various genes: HIF-1 $\alpha$ , VEGFA, PDGFR- $\beta$ , and hTERT in HCT-116 cell lines. Compounds were tested at their 24 h IC<sub>50</sub> values, mRNA was quantified, and gene expression was normalized to DMSO vehicle control (Figure 3). Although compounds 4 and 12 both bind to the HIF-1 $\alpha$  G-quadruplex in the competition dialysis assay, these compounds do not show any notable effect on HIF-1 $\alpha$  expression. Compound 4 demonstrates moderate binding to the hTERT G-quadruplex, with concurrent, significant, time-dependent lowering of mRNA expression, whereas the lower

binding affinity compound 12 significantly lowers hTERT mRNA at 6 and 24 h, but expression normalizes by 48 h. This pattern is again seen with PDGFR- $\beta$  by compound 12, which is in accordance with similar binding profiles as measured by competition dialysis. This compound does not demonstrate binding to the VEGFA G-quadruplex, and qPCR reveals only a late, but significant, increase in expression. Compound 4 shows strong binding to the G-quadruplexes in both the PDGFR- $\beta$  and VEGFA promoters and significant effects on mRNA expression of each target. While there is no direct correlation





**Figure 6.** Exon-specific assay data for compounds 4, 12, GQC-05, and quindoline-i. CA46 Burkitt's lymphoma cells were exposed to 19.4, 2.8, 13, and 10  $\mu$ M of each compound, respectively, for 6–48 h. If an exon-specific effect in c-MYC expression were noted, a preferential decrease in the nontranslocated exon 1 would be expected, as the G-quadruplex is maintained. Compound 12 served as the negative control in this experiment, as predicted by the lack of c-MYC lowering ability in HCT-116 cells, and accordingly there was no noted change in expression. Quindoline-i significantly decreased expression from both exons, whereas compound 4 led to a significant decrease in exon 2 through 24 h, with significant increases in exon 1 from 24 to 48 h. Exon 1 is expressed  $1/1000$  compared to exon 2, so while this increase is significant from the baseline, there will be no notable increase in intracellular c-MYC because of this. GQC-05 significantly decreased MYC mRNA expression in exon 1 but not in exon 2 at the 24 h time point: (\*)  $p < 0.05$  as compared to time-matched vehicle control; (\*\*)  $p < 0.05$  between exons.

between G-quadruplex affinity, as measured by the competition dialysis assay, and intracellular transcriptional regulation, it is clear that these compounds do not have specific effects on c-MYC transcription.

**Structure–Activity Relationships. Melting Temperature.** Melting temperature data ( $\Delta T_m$ ) of quindoline compounds 2–16 are correlated with the ability of these compounds to stabilize and interact with the c-MYC G-quadruplex and can be used to deduce some preliminary conclusions about structure–activity relationships as shown in Figure 4.

- (1) The quindoline N5 atom is highly basic because of the presence of a piperazine ring at position C11. This nitrogen atom is protonated at physiological pH, and this ionization is important for G-quadruplex binding. Compound 16, a phenyl analogue of compound 5, shows very low  $\Delta T_m$ , 1.9  $^{\circ}$ C.
- (2) The presence of a basic amino group in the side chain is important for binding. Compound 4 shows better stabilization than compound 3. Similarly, compound 8 (dimethylamino group) shows higher  $\Delta T_m$  than 9 or 10 (methoxy or hydroxyl).
- (3) A compound with a longer linker length, compound 8 (three carbons), shows higher  $\Delta T_m$  than the two-carbon linker, compound 4.
- (4) Substitution of a second alkylamino chain on indole nitrogen coupled with the presence of a basic amino group on piperazine increases G-quadruplex stabilization. Compound 12 shows higher  $\Delta T_m$  than compound 4. However, compound 11, an analogue of compound 3, does not show any increase in  $\Delta T_m$  after the addition of a second group at  $R_2$ .

Similar SAR conclusions can be drawn for cytotoxicity data in HCT-116 cell lines. A schematic depiction is shown in Figure 4. Compounds 4, 6, and 8, with amino side chains at  $R_1$ , lead to better cytotoxicity in HCT-116 cell lines. Substitution at the  $R_2$  position (indole nitrogen) leads to lower cytotoxicity.

**Exon-Specific Assay in CA46 Cells.** Historically, compounds with purported mechanisms of action attributable to the c-MYC G-quadruplex were confirmed in vitro with a pair of Burkitt's cell lines that were oncogenically transformed and driven by a translocation between an immunoglobulin gene and the c-MYC gene. This translocation either maintains, in the case of RAJI cells, or loses, in the case of CA46 cells, the endogenous promoter region of c-MYC, including the G-quadruplex-forming NHE III<sub>1</sub> region through exon 1. While these cell lines harbor similar basal levels of c-MYC expression and are a good indication of G-quadruplex-mediated effects on c-MYC, they are not an isogenic pair and are fundamentally different in a number of aspects unrelated to either c-MYC or the G-quadruplex.

Recently, on the basis of the observation that one can separate the mRNA products from the translocated and nontranslocated chromosomes in the CA46 cells, we described the “CA46 exon-specific” test.<sup>14</sup> In this test, amplification of exon 1 is used to monitor the effects on the nontranslocated, G-quadruplex-maintaining c-MYC gene, while amplification of exon 2 will predominantly represent c-MYC expression from the translocated, G-quadruplex-lost chromosome (Figure 5). To determine if the c-MYC-lowering effect of compound 4 is mediated through the c-MYC G-quadruplex and to further examine the mechanism of compound 12, we tested the compounds in this exon-specific assay in CA46 cells. If the mechanism of c-MYC transcriptional modulation is mediated through stabilization of the G-quadruplex, a preferential decrease in exon 1, but not exon 2, would be noted, as previously reported for GQC-05.<sup>14</sup> While this alone is not conclusive for a G-quadruplex-mediated compound effect and further analysis with chromatin immunoprecipitation for displacement of transcription factors would be needed as confirmatory data, an absence of this exon-specific effect would enable a confident refusal of a c-MYC G-quadruplex-mediated action.

Although the initial report of quindoline-i suggested targeting of this element,<sup>11</sup> the CA46 test (measured at 24 h in a previous report) did not.<sup>14</sup> Here we describe a more thorough time-course with quindoline-i as the parent compound for these new analogues, as well as for compounds **4** and **12** (Figure 6). Each compound was tested at its respective 24 h IC<sub>50</sub> in CA46 cells (19.4, 2.8, and 10  $\mu$ M for **4**, **12**, and quindoline-i, respectively). As anticipated by the previous results shown in Figure 2, there was no change in c-MYC expression, on either exon, induced by compound **12** at 24 and 48 h, but contrary to the results in Figure 2, no lowering of c-MYC was seen at 6 h. For compound **4**, which showed a significant change in c-MYC expression (Figure 2), there was no reduction in c-MYC expression in exon 1 corresponding to the nontranslocated allele still maintaining the G-quadruplex, whereas c-MYC lowering was evident in the translocated allele. Thus, the c-MYC lowering seen by compound **4** in any cell line is due to a non-G-quadruplex-specific effect.

For quindoline-i the interpretation of the results is more complex. There is clearly no exon-specific effect, as we have observed previously for GQC-05, because lowering of c-MYC expression occurred for both exon 1 and exon 2 at all three times evaluated. In each case the extent of suppression of gene expression was about the same. Thus, the explanation for the decrease in c-MYC mRNA expression in the nontranslocated allele where the G-quadruplex is retained can most likely be explained by the same, but as yet unknown, non-G-quadruplex-mediated effect seen in the translocated allele rather than a direct effect through the G-quadruplex in the c-MYC promoter.

11-Piperazinyl-substituted quindoline analogues were primarily designed as c-MYC G-quadruplex-interactive compounds. However, as evident from competition dialysis experiments, transcriptional data, and the exon-specific assay in CA46 cells, these compounds are predominantly non-selective. As observed with other G-quadruplex-interactive compounds, their anticancer activity in cells is due to the interaction with G-quadruplex in multiple oncogenes and cannot be attributed only to c-MYC or any other single gene.

## CONCLUSIONS

We have synthesized a series of 11-piperazinyl-substituted quindoline analogues and tested two of these compounds further for cytotoxicity in HCT-116 and Raji cell lines. The most active compound (**4**) shows an IC<sub>50</sub> of 0.97  $\mu$ M in HCT-116 cell lines and an IC<sub>50</sub> of 2.33  $\mu$ M in Raji cell lines. This compound also exhibits reduction in c-MYC mRNA expression in HCT-116 cells at 6 h, and this continues through 48 h. Despite the significant effect on c-MYC expression, further analysis using the exon-specific effect in CA46 did not support the mechanism of action of this compound to be through the G-quadruplex within the NHE III<sub>1</sub>. Compounds **4** and **12** are both nonselective and also affect the expression of other target genes. This work suggests that these G-quadruplex-interactive compounds are not selective and their anticancer activity could be due to a combination of effects on different genes in colon and lymphoma cell lines. While initial results led to excitement about new and potent compounds, complete cell-free and in vitro analyses enabled us to conclude that they are not working in the cell through the expected target mechanism of action. Thus, it is imperative to fully characterize compounds using appropriate cellular systems before claims are made for putative G-quadruplex-interactive compounds that target specific G-quadruplexes in promoter elements. Proof of principle in

cellular systems similar to the CA46 exon-specific assay described here is critical if such claims are to be made.

## ASSOCIATED CONTENT

### Supporting Information

Spectral data (<sup>1</sup>H and <sup>13</sup>C NMR, HRMS, and HPLC–MS) for compounds **2**–**16**. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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### Notes

The authors declare no competing financial interest.

## ACKNOWLEDGMENTS

This research was supported by grants from Science Foundation Arizona (Grant SBC 0016-07), the National Institutes of Health (Grant CA95060), and the National Foundation for Cancer Research (Grant VONHOFF0601). We thank David Bishop for preparing, proofreading, and editing the final version of the text and figures.

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