Effective C-N bond formation on the 1-methyl-2-quinolone skeleton

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Abstract
Electron-deficient 1-methyl-3,6,8-trinitro-2-quinolone is highly reactive, and readily reacted with
two equivalents of primary amines at room temperature to afford ammonium salts of 4-
alkylamino-3,4-dihydro-1-methyl-3,6,8-trinitro-2-quinolones. 4-Alkylamino-6,8-dinitro-1-
methyl-2-quinolones were also prepared when the reaction was conducted at elevated
temperature.

Keywords: 1-Methyl-2-quinolone, cine-substitution, dihydroquinolone

Introduction

The 1-methyl-2-quinolone (MeQone) framework is often found in quinoline alkaloids isolated
from the Rutaceae family, and numerous naturally occurring MeQones have been synthesized.1,2
In addition, development of preparative methods for unnatural MeQone derivatives is also one of
the important projects for researchers studying drug design.3 From this viewpoint, we have paid
attention to high reactivity of 1-methyl-3,6,8-trinitro-2-quinolone (1), and have studied the
functionalization of the MeQone skeleton. The 4-position of 1 is highly reactive because of steric
repulsion between 1-methyl and 8-nitro groups.4 The C-C bond formation at this position is
easily performed accompanied by elimination of an adjacent nitro group when 1 is treated with
nucleophiles such as 1,3-dicarbonyl compounds,5 ketones6 and enamines.6 The present cine-
substitution is initiated by addition of the nucleophile at the 4-position, and the following
aromatization together with elimination of nitrous acid affords 4-substituted 6,8-dinitro-1-
methyl-2-quinolones 2 (Scheme 1). On the basis of these results, the C-N bond formation is
attempted by use of amine as the nucleophile since introduction of a nitrogen function into the 4-
position enables further functionalization of the MeQones skeleton.7
Results and Discussion

When 1 was treated with methylamine in acetonitrile at room temperature, the solution was immediately turned to reddish yellow, and precipitation of ammonium salt 3a was observed during the reaction. In the $^1$H NMR of 3a, three singlet signals were observed at 2.95, 8.41, 8.56 ppm, however the other signals were considerably broadened. It is considered that amine attacks at the 4-position of 1 to give adduct intermediate 4. The proton on the amino group was intramolecularly transferred to the 3-position affording dihydroquinolone 5. As a result of deprotonation at the 3-position (H_a) by another amine, ammonium 3 is formed (Scheme 2, route a). As another route, direct conversion from 4 to 3 is also possible.

The present reaction was applicable to other primary amines having a more bulky substituent. The reaction of 1 with propylamine similarly furnished ammonium salt 3b in a good yield. It was also possible to employ sterically hindered amines such as isopropylamine, isobutylamine, sec-butylamine, tert-butylamine and benzylamine, and leading to corresponding ammonium salts 3c-g, respectively (Table).
Table 1. Preparation of ammonium salt 3

<table>
<thead>
<tr>
<th>R</th>
<th>Yield / %</th>
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<tbody>
<tr>
<td>Me</td>
<td>a 91</td>
</tr>
<tr>
<td>Pr</td>
<td>b 71</td>
</tr>
<tr>
<td>i-Pr</td>
<td>c 79</td>
</tr>
<tr>
<td>i-Bu</td>
<td>d 74</td>
</tr>
<tr>
<td>sec-Bu</td>
<td>e 56</td>
</tr>
<tr>
<td>tert-Bu</td>
<td>f 74</td>
</tr>
<tr>
<td>PhCH2</td>
<td>g 79</td>
</tr>
</tbody>
</table>

Preparation of cine-substituted quinolones 6 was attempted. When a solution of 1 and propylamine in acetonitrile was heated under reflux, 6,8-dinitro-1-methyl-4-propylamino-2-quinolone (6b) was obtained 36% yield. Isobutylamino derivative 6d was also prepared in 29% yield in the reaction of 1 with isobutylamine. In cases of more bulky amines such as sec-butylamine and tert-butylamine, 6e and 6f were not formed. cine-Substituted quinolones 6 were produced by elimination of nitrous acid from the 3- and the 4-positions (Scheme 2, route b), however sterically hindered alkylamino groups prevented deprotonation at the vicinal position (Hb).

The regioselective C-N bond formation on the MeQone skeleton was readily performed. It is expected to prepare new MeQone derivatives by use of this method.

Experimental Section

General Procedures. The melting points were determined on a Yanaco micro-melting-points apparatus, and were uncorrected. 1H and 13C NMR spectra were measured on a Bruker DPX-400 at 400 MHz and at 100 MHz respectively with TMS as an internal standard. Assignments of 13C NMR spectra (s, d, t and q) were made from DEPT experiments. IR spectra were recorded on a Horiba FT-200 IR spectrometer. Elemental microanalyses were performed using a Yanaco MT-3 CHN corder. In some cases, satisfactory analytical data of salt 3 were not obtained because of unstable and hygroscopic properties. All the reagents and solvents were commercially available and used as received. All of reactions were carried out under ambient atmosphere. Column chromatography was performed using Wakogel C-200.

1-Methyl-3,6,8-trinitro-2-quinolone (1). Following the procedure described for 1-methyl-2-pyridone, MeQone was prepared by oxidation of 1-methylquinolinium ion using potassium ferricyanide (III) under alkaline conditions after methylation of quinoline with dimethyl sulfate using the three times diluted solution. Nitration of MeQone with fuming nitric acid (d = 1.52) afforded 1 in 90% yield.
Ammonium salts of 4-alkylamino-3,4-dihydro-1-methyl-3,6,8-trinitro-2-quinolone (3). To a solution of 1 (300 mg, 1 mmol) in acetonitrile (15 mL), was added over 5 minutes a solution of amine (5 mmol) in acetonitrile (5 mL). The resultant solution was stirred at room temperature for 2 hours. The precipitated yellow solid during the reaction was collected by filtration, and successively washed with small amounts of acetonitrile and hexane to afford ammonium salt 3.

**Methylammonium 3,4-dihydro-6,8-dinitro-1-methyl-4-methylamino-2-quinolone-3-nitronate (3a).** Yellow powder, mp 103-111 °C (dec.). IR (KBr / cm⁻¹) 3600-3300 (br), 1659, 1603, 1539, 1524, 1338, 1215, 1074, 974; ¹H NMR (400 MHz, DMSO-d₆) δ 2.07 (br s, 3H), 2.40 (br s, 3H), 2.95 (s, 3H), 5.34 (br s, 1H), 5.5-6.5 (br, 4H), 8.41 (s, 1H), 8.56 (s, 1H). Anal. Calcd. for C₁₂H₁₆N₆O₇: C, 40.45; H, 4.53, N, 23.59. Found: C, 40.42; H, 4.47; N, 23.23.

**Propylammonium 3,4-dihydro-6,8-dinitro-1-methyl-4-propylamino-2-quinolone-3-nitronate (3b).** Yellow powder, mp 99-107 °C (dec.). IR (KBr / cm⁻¹) 3500-3300 (br), 1653, 1608, 1537, 1338, 1221, 1072, 980; ¹H NMR (400 MHz, DMSO-d₆) δ 0.7-0.9 (br, 6H), 1.3-1.4 (br, 2H), 1.5-1.6 (br, 2H), 2.2-2.3 (br, 1H), 2.3-2.4 (br, 1H), 2.6-2.8 (br, 3H), 2.96 (s, 3H), 5.38 (br s, 1H), 4.5-7.0 (br, 3H), 8.40 (s, 1H), 8.56 (s, 1H). Anal. Calcd. for C₁₂H₂₄N₆O₇: C, 46.60; H, 5.87; N, 20.38. Found: C, 46.82; H, 5.60, N, 20.14.

**Isopropylammonium 3,4-dihydro-6,8-dinitro-4-isopropylamino-1-methyl-2-quinolone-3-nitronate (3c).** Yellow powder, mp 110-120 °C (dec.). IR (KBr / cm⁻¹) 3500-3300 (br), 1657, 1605, 1533, 1336, 1223, 1074, 978; ¹H NMR (400 MHz, DMSO-d₆) δ 0.5-1.5 (br, 12H), 2.7-2.8 (br, 1H), 2.9-3.0 (br, 3H), 3.0-3.2 (br, 2H), 5.1-5.3 (br, 1H), 4.5-7.0 (br, 3H), 8.2-8.3 (br, 1H), 8.5-8.6 (br, 1H). Anal. Calcd. for C₁₂H₂₄N₆O₇: C, 46.60; H, 5.87; N, 20.38. Found: C, 46.32; H, 5.87, N, 20.65.

**Isobutylammonium 3,4-dihydro-6,8-dinitro-4-isobutylamino-1-methyl-2-quinolone-3-nitronate (3d).** Yellow powder, mp 90-110 °C (dec.). IR (KBr / cm⁻¹) 3500-3300 (br), 1657, 1605, 1527, 1336, 1227, 1072, 980. Since this salt was decomposed in DMSO during the measurement, satisfactory NMR data could not be obtained. Anal. Calcd. for C₁₈H₂₈N₆O₇: C, 49.09; H, 6.41; N, 19.08. Found: C, 48.51; H, 6.22, N, 19.22.

**sec-Butylammonium 4-sec-butylamino-3,4-dihydro-6,8-dinitro-1-methyl-2-quinolone-3-nitronate (3e).** Yellow powder, mp 90-100 °C (dec.). IR (KBr / cm⁻¹) 3500-3300 (br), 1660, 1606, 1527, 1336, 1227, 1074, 980; ¹H NMR (400 MHz, DMSO-d₆) δ 0.4-1.9 (br, 16H), 2.7-3.2 (br, 5H), 3.5-3.5 (br, 1H), 4.0-7.0 (br, 3H), 8.30 (br s, 1H), 8.56 (br s, 1H). Anal. Calcd. for C₁₈H₂₈N₆O₇: C, 49.09; H, 6.41; N, 19.08. Found: C, 48.66; H, 6.32, N, 19.13.

**tert-Butylammonium 4-tert-butylamino-3,4-dihydro-6,8-dinitro-1-methyl-2-quinolone-3-nitronate (3f).** Yellow powder, mp 100-120 °C (dec.). IR (KBr / cm⁻¹) 3500-3300 (br), 1674, 1605, 1518, 1331, 1225, 1074, 974; ¹H NMR (400 MHz, DMSO-d₆) δ 0.4-1.9 (br, 18H), 2.7-3.9 (br, 3H), 5.51 (br s, 1H), 3.9-6.2 (br, 4H), 8.8-9.5 (br, 2H).

**Benzylammonium 4-benzylamino-3,4-dihydro-6,8-dinitro-1-methyl-2-quinolone-3-nitronate (3g).** Yellow powder, mp 130-136 °C (dec.). IR (KBr / cm⁻¹) 3500-3300 (br), 1653, 1606, 1531,
1338, 1221, 1072, 982; $^1$H NMR (400 MHz, DMSO-$d_6$) $\delta$ 2.6-3.3 (br, 3H), 3.3-4.4 (br, 5H), 4.7-6.8 (br, 4H), 6.8-7.7 (br, 10H), 8.8-9.6 (br, 2H). Anal. Calcd. for C$_{24}$H$_{24}$N$_6$O$_7$: C, 56.69; H, 4.76; N, 16.53. Found: C, 56.74; H, 4.69, N, 16.66.

4-Alkylamino-6,8-dinitro-1-methyl-2-quinolone (6). To a solution of 1 (300 mg, 1 mmol) in acetonitrile (15 mL), was added over 5 minutes a solution of amine (5 mmol) in acetonitrile (5 mL). After purge with nitrogen gas, the mixture was heated under reflux for 3 hours. The precipitated yellow solid was collected by filtration after cooling down to room temperature, and successively washed with small amounts of acetonitrile and hexane to afford cine-substituted product 6. The filtrate was concentrated, and the residue was treated with column chromatography on silica gel to give 6 (eluted with toluene).

6,8-Dinitro-1-methyl-4-propylamino-2-quinolone (6b). Yellow needles, mp 263-268 °C (dec.). IR (KBr / cm$^{-1}$) 3375, 1626, 1539, 1525, 1336, 1279; $^1$H NMR (400 MHz, CDCl$_3$) $\delta$ 1.09 (t, $J$ = 7.4 Hz, 3H), 2.09 (triple septet, $J$ = 7.4, 7.4 Hz, 1H), 3.26 (dt, $J$ = 7.4, 5.3 Hz, 2H), 3.41 (s, 3H), 4.98 (br s, 1H), 5.86 (s, 1H), 8.60 (d, $J$ = 4.7 Hz, 1H), 8.73 (d, $J$ = 4.7 Hz, 1H); $^{13}$C NMR (100 MHz, CDCl$_3$) $\delta$ 11.6 (q), 21.7 (t), 34.2 (q), 45.4 (t), 94.2 (d), 118.7 (s), 119.6 (d), 122.3 (d), 125.9 (s), 139.1 (s), 148.6 (s), 153.1 (s), 163.1 (s). Anal. Calcd. for C$_{13}$H$_{14}$N$_4$O$_5$: C, 50.98; H, 4.61; N, 18.29. Found: C, 50.87; H, 4.51, N, 18.35.

6,8-Dinitro-4-isobutylamino-1-methyl-2-quinolone (6d). Yellow needles, mp 188-189 °C (dec.). IR (KBr / cm$^{-1}$) 3365, 1626, 1547, 1522, 1335, 1277; $^1$H NMR (400 MHz, CDCl$_3$) $\delta$ 1.07 (d, $J$ = 6.7 Hz, 6H), 2.09 (triple septet, $J$ = 6.7, 6.7 Hz, 1H), 3.0-3.1 (br, 2H), 3.41 (s, 3H), 5.00 (br t, 1H), 5.86 (s, 1H), 8.60 (d, $J$ = 6.7 Hz, 1H), 8.73 (d, $J$ = 6.7 Hz, 1H). Anal. Calcd. for C$_{14}$H$_{16}$N$_4$O$_5 \cdot 1 / 2$ toluene: C, 57.37; H, 5.50; N, 15.29. Found: C, 57.44; H, 5.43, N, 15.30.

References

