Novel Synthesis of Ureas: Application of t-Butylureas

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Received September 28, 2009; accepted October 30, 2009; published online November 4, 2009

Otherwise inaccessible tropolonylureas were prepared by reaction of t-butylurea with appropriate amines, with elimination of t-butylamine. This method is also generally applicable for urea synthesis.

Key words urea; t-butylurea; isocyanate; tropolone; substitution

Urea is an important pharmacophore which is present in various pharmaceutical drugs and agrochemicals. For example, terguride is a dopamine agonist, many sulfonlureas are used as therapeutics for diabetes, sorafenib is used to treat renal cell carcinoma, N-(3,4-dichlorophenyl)-N,N'-dimethylurea (diuron) is a herbicide, and N-(2-chloro-4-pyridyl)-N'-phenylurea (forchlorofenuron) is a plant growth regulator. Such compounds may be prepared by conventional methods, i.e., reaction of an amine with an isocyanate (or in situ-formed isocyanate generated by Curtius rearrangement of acid azide), reaction of an amine and a carbamate, or reaction of an amine and phosgene or its equivalents. The last method is convenient, but sometimes encounters difficulties when the amine or intermediate is reactive. In some cases, none of these methods can be easily applied.

We have been investigating the structural and medicinal chemistry of tropolone (2-hydroxy-2,4,6-cycloheptatrien-1-one), which is a nonbenzenoid aromatic compound having a flat seven-membered ring. The tropolone fragment can be used as a pharmacophore instead of benzoic acid, phenol or other heteroaromatic moieties, thujaplicins and colchicines, possess antibacterial, antifungal, antiviral, and antitumor properties. Some tropolone derivatives show significant retinoid activity. Therefore, we set out to synthesize various tropolonylureas from 5-aminotropolone (1). When we attempted to extend tropolone chemistry, we found that compounds such as N,N'-di(5-tropolonyl)urea (2) were not readily accessible from 1, since we could not obtain 5-tropolonyl isocyanate by conventional methods (e.g., using phosgene equivalents). Fortunately, we found a method to prepare compound 2 (Chart 1), and this method can also be applied to general synthesis of urea compounds, as described below.

Results and Discussion

5-Aminotropolone (1) reacts with various isocyanates, such as phenyl, substituted phenyl, and aliphatic isocyanates, leaving the hydroxy group intact. However, when 1 was treated with 1 eq of t-butyl isocyanate at reflux for 51 h in tetrahydrofuran (THF), N,N'-di(5-tropolonyl)urea (2) was unexpectedly obtained as a major product instead of N-t-butyl-N'-tropolonylurea (3). In order to obtain 3, an excess amount of t-butyl isocyanate and a shorter reaction time are required, suggesting that 3 may serve as an intermediate in the formation of 2. Indeed, when isolated 3 was treated with 1 in toluene at 120 °C for 14 h, 2 was smoothly generated in a good yield of 82% (Chart 1).

When N-adamantyl-N'-tropolonylurea was prepared and reacted with 1 under similar conditions, 2 was obtained in a lower yield of 36%. The main reason for this may be that the adamantylamine is not readily removed during the reaction, because of its high boiling point. Therefore, the t-butylamine may be the best leaving alkylamine in terms of both bulkiness (high eliminability) and high volatility. This method for urea preparation, utilizing t-butylurea, should be available not only for 5-tropolonylureas, but also for general preparation of ureas (vide infra).

Reaction of 3 with a variety of primary and secondary amines (e.g., aromatic, heterocyclic, aliphatic amines) in toluene at 120 °C afforded the desired ureas in good yield (Table 1, entries 1—19). A hindered amine, 2,6-dimethylaniline, reacted with 3 to give 4a (entry 1). The reaction proceeded well with amines that are nucleophilic (e.g., entries 2, 4, 6) or weakly nucleophilic (e.g., entries 3, 5, 8, 9, 11). As expected, 3 reacted with piperazine, a bidentate amine, to afford N-piperazinyl-N'-tropolonylurea (4l) in 72% yield, leaving one amino group intact (entry 12). Ureas 4m—p were also readily obtained by the reaction of 3 with other aliphatic secondary amines (entries 13—16). The method was also extended to more complex tropolonylureas (entries 18, 19). The reaction of 3 with bioactive substances, such as histamine and pindolol, provided the corresponding ureas 4r and 4s derivatized from the aliphatic amines, respectively, in high yield. The reactions of N-t-butyl-N'-phenylurea (5) and N-t-butyl-N'-3-pyridylurea (7) with amines are illustrated.
Possible reaction pathways include a stepwise reaction involving an isocyanate intermediate formed by elimination of \( t \)-butylamine, and/or direct nucleophilic substitution of the \( t \)-butylamine by amine \( R'NH \) (Chart 2). It was reported that \( N \)-alkyl-\( N' \)-phenylureas can be prepared from \( N, N' \)-diphenylurea by treatment with aliphatic amines, though this reaction lacks versatility, and the reaction mechanism has not been established. The presence of amine is necessary for the reaction to proceed. The amine may catalyze elimination of the \( t \)-butylamine to afford the isocyanate. Or, nucleophilic attack of the amine on the carbonyl group and subsequent removal of \( t \)-butylamine may occur. In any case, the formed \( t \)-butylamine (bp 44—46 °C) is highly volatile, and can be easily removed from the reaction medium. \( t \)-Butylamine evaporated from the reaction medium can be easily detected with pH test paper.

### Conclusion

A simple and versatile method for the preparation of ureas, based on reaction of \( t \)-butylureas with amines, was developed. This method is especially useful for synthesizing urea compounds for which corresponding isocyanates or their precursors are not available.

### Experimental

#### General Procedures

Melting points were determined on a Yanagimoto micro-melting point apparatus (hot plate) without correction. Mass spectra were recorded on a Hitachi M-80B spectrometer at an ionizing voltage of 70 eV, and figures in parentheses indicate the relative intensities. IR spectra were measured on a Shimadzu IR-460 spectrophotometer. \(^1\)H- and \(^{13}\)C-NMR

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### Table 1. Reaction of 3-Butylureas (3, 5, 7) with Various Amines

<table>
<thead>
<tr>
<th>Entry</th>
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<th>Time (h)</th>
<th>Product</th>
<th>Yield (%)</th>
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<td></td>
<td>6</td>
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\( a \) See Experimental.

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spectra were obtained on a Varian Mercury 300 at 300 MHz and at 75 MHz, respectively. Chemical shifts are recorded in ppm (3) and coupling constants (J) in Hz. Chemical shifts were calculated on the basis of tetramethylsilane (0 ppm for \( ^1H\)-NMR in CDCl₃, CH₃SO (2.49 ppm for \(^1H\)-NMR in DMSO-d₆), CH₃OH (3.59 ppm for \(^1H\)-NMR in CD₃OD). Fuji Davison BW 300 or Merck aluminum oxide 90 was used for flash chromatography, and TLC was carried out on Merck Silica gel 60 PF₅₀.

**Reaction of 5-Aminotropolone (1) with 1 eq of t-Butyl Isocyanate.** A solution of 5-aminotropolone (1, 206 mg, 1.50 mmol) and t-butyl isocyanate (171 μL, 1.50 mmol) in THF (10 mL) was refluxed for 5 h, then cooled to room temperature, and the solvent was evaporated in vacuo. The residue was washed with MeOH to give N,N'-di(tropolonyl)urea (2, 96 mg, 43%) as dark brown powder, mp 258–260 °C (dec.). \(^1H\)-NMR (DMSO-d₆) δ: 9.06 (2H, brs, NH×2), 7.55 (4H, d, J = 12.0 Hz), 7.21 (4H, d, J = 12.0 Hz). IR (KBr) cm⁻¹: 3195, 1612, 1548, 1448, 1419, 1262. MS m/z: 163 (M⁺-137), 137 (65), 135 (59), 109 (80), 57 (52). The filtrate was evaporated to give a residue (184 mg) which consisted of three compounds, N-t-butyl-N'-(tropolonyl)urea (3), N,N'-di(t-butyl)urea, and I in a ratio of ca. 1:0.7:0.5 estimated by the H-NMR integration of the respective NH groups.

**Reaction of 1 with 3 eq of t-Butyl Isocyanate.** A solution of 1 (411 mg, 3.00 mmol) and t-butyl isocyanate (1.06 mL, 9.00 mmol) in THF (10 mL) was refluxed for 5 h, then cooled to room temperature and the solvent was evaporated in vacuo. The residue was washed with MeOH to give N,N,N'-trimethylurea (4a) as a dark brown powder, mp 197–202 °C. \(^1H\)-NMR (DMSO-d₆) δ: 10.98 (2H, brs, NH×2), 7.51 (2H, d, J = 11.0, 1.5 Hz), 7.17 (2H, d, J = 11.0, 1.5 Hz), 6.13 (1H, brs, NH), 1.27 (9H, s, t-Bu). \(^13C\)-NMR (DMSO-d₆) δ: 168.0, 153.8, 140.9, 126.3, 125.2, 49.7, 28.9. IR (KBr) cm⁻¹: 3350, 3170, 3005, 1650, 1446, 1359, 1260. HR-MS m/z: 236.1163 (Calcd for C₁₉H₂₆N₂O₃: 236.1160). MS m/z: 236 (M⁺, 9), 163 (13), 137 (100).

**Reaction of N,N'-Di(t-Butyl)-N'-(tropolonyl)urea (3) with 1 Solution of 1 (253 mg, 1.07 mmol) and 1 (293 mg, 1.24 mmol) in toluene (10 mL) was heated at 120 °C for 14 h, then cooled to room temperature and the solvent was evaporated in vacuo. The residue was washed with MeOH to give 2 (264 mg, 82%).

**Preparation of N-(1-Adamantyl)-N'-tropolonylurea (5a) and N,N'-tropolonylurea (5b).** A solution of 1 (206 mg, 1.50 mmol) and 1-adamantyl isocyanate (531 mg, 3.00 mmol) in THF (10 mL) was refluxed for 28 h, then cooled to room temperature and the solvent was evaporated in vacuo. The residue was flash-chromatographed [silica gel, CH₃Cl₂-1,2-dimethylethane (20:1)] to get N-(1-adamantyl)-N'-tropolonylurea (6a) (168 mg, 36%) as brown scales (MeOH), mp 214–216 °C and 230–235 °C (dec.). \(^1H\)-NMR (DMSO-d₆) δ: 8.51 (1H, brs, NH), 7.50 (2H, d, J = 12.3 Hz), 7.17 (2H, d, J = 12.3 Hz), 6.01 (1H, brs, NH), 2.06–1.98 (3H, m), 1.94–1.89 (6H, m), 1.64–1.59 (6H, m). IR (KBr) cm⁻¹: 3350, 3170, 1529, 1456, 1149, 1142, 1073. MS m/z: 324.1616 (Calcd for C₂₀H₂₅N₂O₃: 324.1619). MS m/z: 314 (M⁺, 0.3), 163 (20), 151 (17), 135 (17), 107 (15), 94 (100).

**Tropolonyleuronea 4a–s; General Procedure.** A solution of 3 (0.5 mmol) and amine (1–3 eq) in toluene (5 mL) was heated at 120 °C. The reaction was monitored by NMR (or TLC). When the reaction reached completion, the solvent was evaporated in vacuo. The residue was washed with MeOH (or CH₃Cl₂, ether, or H₂O), and the crude product was collected by filtration. If necessary, the residue was purified by flash chromatography [silica gel or aluminum oxide].

**N-(2,6-Dimethylphenyl)-N'-tropolonylurea (4a) and N-(2,6-Dimethylphenyl)-N'-tropolonylurea (4b).** Pale yellow needles (MeOH), mp 260–262 °C (dec.). \(^1H\)-NMR (DMSO-d₆) δ: 8.85 (1H, brs, NH), 7.88 (1H, brs, NH), 7.50 (2H, d, J = 12.3 Hz), 7.20 (2H, d, J = 12.3 Hz), 7.01 (3H, s), 2.19 (6H, s, Me₂). HR-MS m/z: 284.1163 (Calcd for C₁₄H₁₄N₂O₃: 284.1160). MS m/z: 284 (M⁺, 4), 163 (53), 149 (49), 121 (34).

**N-(4-Methylphenyl)-N'-tropolonylurea (4b).** Pale yellow needles (MeOH), mp 260–262 °C (dec.). \(^1H\)-NMR (DMSO-d₆) δ: 8.85 (1H, brs, NH), 6.88 (1H, brs, NH), 7.56 (2H, d, J = 12.3 Hz), 7.32 (2H, d, J = 8.7 Hz), 7.20 (2H, d, J = 12.3 Hz), 7.08 (2H, d, J = 8.7 Hz), 2.32 (3H, s, Me). IR (KBr) cm⁻¹: 3300, 3210, 1683, 1605, 1544, 1449, 1425, 1355. HR-MS m/z: 270.0997 (Calcd for C₁₆H₁₃N₂O₃: 270.1004). MS m/z: 270 (M⁺, 2), 163 (49), 121 (34).

**N-(3,4-Difluorophenyl)-N'-tropolonylurea (4c).** N,N'-Di(t-butyl)urea (4d) is Ochre prisms (AcOEt), mp 134–136 °C. \(^1H\)-NMR (DMSO-d₆) δ: 8.37 (1H, brs, NH), 7.46 (2H, dd, J = 11.1, 1.5 Hz), 7.41 (2H, dd, J = 7.2, 1.2 Hz), 7.25 (1H, tt, J = 7.2, 1.2 Hz), 7.14 (2H, dd, J = 11.1, 1.5 Hz), 3.25 (3H, s, OMe). \(^13C\)-NMR (DMSO-d₆) δ: 168.8, 154.7, 143.5, 139.9, 130.1, 129.1, 126.5, 125.9, 124.0, 39.7. IR (KBr) cm⁻¹: 3400, 3225, 1684, 1543, 1496, 1445, 1363, 1262. HR-MS m/z: 270.1015 (Calcd for C₁₁H₈F₂N₂O₃: 270.0944). MS m/z: 270 (M⁺, 27), 163 (29), 134 (82), 106 (1).
3.48 (8H, s). HR-MS (3H, s, Me), 2.00 (2H, m). HR-MS ms/z: 265.1414 (Calcd for C13H19N3O3: 265.1425). MS (3H, s, Me), 2.00 (2H, m). HR-MS ms/z: 265.1414 (Calcd for C13H19N3O3: 265.1425). MS ms/z: 250 (M⁺, 26), 163 (55), 135 (47), 114 (22), 107 (38), 58 (100).

c) 4-(6-Tropolonyl)morpholine (40) Pale yellow powder (MeOH), mp 213—215 °C. 1H-NMR (CDCl3): δ: 7.43 (2H, d, J = 12.3 Hz), 7.29 (2H, d, J = 12.3 Hz), 6.43 (4H, t, J = 4.5 Hz), 4.04 (4H, t, J = 4.5 Hz). HR-MS ms/z: 250.0953 (Calcd for C13H19N3O3: 250.0953). MS ms/z: 250 (M⁺, 26), 163 (55), 135 (47), 114 (22), 107 (38), 58 (100).

N-2-[Dimethylamino]ethyl-N-methyl-N’-(5-tropolonyl)urea (4p) Ocher powder, mp 108—110 °C. 1H-NMR (CDCl3): δ: 11.0 (1H, brs, NH), 7.39 (2H, d, J = 12.6 Hz), 7.30 (2H, d, J = 12.6 Hz). 3.38 (2H, t, J = 4.2 Hz), 2.99 (3H, s, Me), 2.63 (2H, t, J = 4.2 Hz), 2.42 (6H, s, Me×2). HR-MS ms/z: 265.1414 (Calcd for C13H19N3O3: 265.1425). MS ms/z: 265 (M⁺, 14), 163 (15), 135 (16), 107 (13), 79 (19), 58 (100).

N-[4-(4-Morpholinopropyl)]N’-(5-tropolonylurea (4q) Ocher powder, mp 186—188 °C. 1H-NMR (CDCl3): δ: 7.62 (2H, dd, J = 11.1, 1.5 Hz), 7.30 (2H, dd, J = 11.1, 1.5 Hz). 3.69 (4H, t, J = 4.5 Hz), 3.24 (2H, t, J = 7.5 Hz). 6.95 (4H, t, J = 4.5 Hz), 2.43 (2H, t, J = 7.5 Hz). 1.74 (2H, t, J = 7.5, 6.9 Hz). HR-MS ms/z: 307.1544 (Calcd for C14H21N3O3: 307.1531). MS ms/z: 305 (M⁺, 15), 163 (59), 135 (39), 112 (100).

N-[2-(4-Iodoazidomethyl)ethyl]N’-(5-tropolonylurea (4r) Yellow powder (AcOEt), mp 190—192 °C. 1H-NMR (CDCl3): δ: 7.60 (1H, d, J = 12.6 Hz), 7.57 (2H, dd, J = 10.8, 1.5 Hz), 7.27 (2H, d, J = 10.8, 1.5 Hz), 6.87 (1H, d, J = 12.6 Hz), 3.45 (2H, t, J = 6.6 Hz), 2.80 (2H, t, J = 6.6 Hz). IR (KBr) cm⁻¹: 3210, 1665, 1504, 1432, 1344. MS ms/z: 163 (M⁺, -111, 52), 137 (15), 135 (50), 107 (43), 82 (100).

2-N-[2-(4-Iodoazidomethyl)ethyl]-N-(2-propyl)N’-(5-tropolonylurea (4s) White powder. 1H-NMR (CDCl3): δ: 6.85 (1H, d, J = 7.5 Hz), 6.73 (2H, d, J = 7.5 Hz), 6.72 (2H, d, J = 7.5 Hz). 3.47 (2H, t, J = 6.6 Hz), 2.61—2.54 (4H, m), 2.26 (2H, t, J = 7.5 Hz), 2.00 (2H, m). HR-MS ms/z: 324 (M⁺, 3), 205 (88), 159 (57), 113 (84), 92 (100).

References