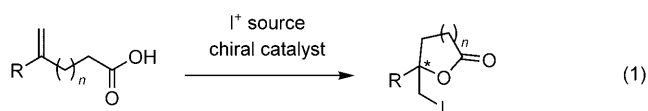


Tertiary Aminourea-Catalyzed Enantioselective Iodolactonization**

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The intramolecular reaction of carboxylic acids with pendant olefins in the presence of a source of I^+ —the iodolactonization reaction—is a powerful method for the generation of five- and six-membered lactones [Eq. (1)]. Diastereoselective variants of this reaction provide efficient access to stereochemically defined lactones, and consequently this reaction has found widespread use in natural products synthesis.^[1] In contrast, the development of catalytic enantioselective variants has proved challenging,^[2,3] a problem likely associated with the inherent difficulty of controlling the reactivity of iodonium ion intermediates through intermolecular interactions.^[4]



The recent discovery of anion-binding mechanisms in H-bonding catalysis^[5] has opened the door to the development of asymmetric catalytic methods that engage reactive cationic intermediates such as *N*-acyliminium ions,^[6] *N*-protoniminium ions,^[7] acylpyridinium ions,^[8] aziridinium ions,^[9] and oxocarbenium ions.^[10] We were intrigued by the possibility that analogous pathways might be available to iodonium ions, thereby providing the control over halonium ion reactivity that is necessary for enantioselective iodolactonization and related reactions. Herein, we report the successful application of such a strategy in the development of a tertiary aminourea-catalyzed asymmetric iodolactonization reaction.

The iodolactonization of hexenoic acid derivative **2a** was selected as a model reaction for catalyst and reagent screening studies. A broad survey of potential H-bond donor catalysts revealed that bifunctional tertiary aminourea derivatives were required to induce useful levels of catalysis. A sharp dependence on the amino group substituents was observed, with di-*n*-pentyl derivative **1** affording highest enantioselectivities.^[11] Whereas *N*-iodoimides or I_2 alone proved poorly reactive (Table 1, entries 1–4), the combination

Table 1: Optimization studies for the enantioselective iodolactonization of **2a**.

Entry ^[a]	I^+ Source	I_2 Additive [mol %]	Yield [%] ^[b]	<i>ee</i> [%] ^[c]
1	4	0	0	–
2	5	0	3	50
3	6	0	25	30
4	I_2	–	12	66
5	4	15	95	92
6	5	15	98	94
7	5	30	85	88

[a] Reactions performed on a 0.05 mmol scale. [b] Determined by 1H NMR analysis using 1,3,5-trimethoxybenzene as the internal standard. [c] Determined by HPLC analysis using commercial chiral columns.

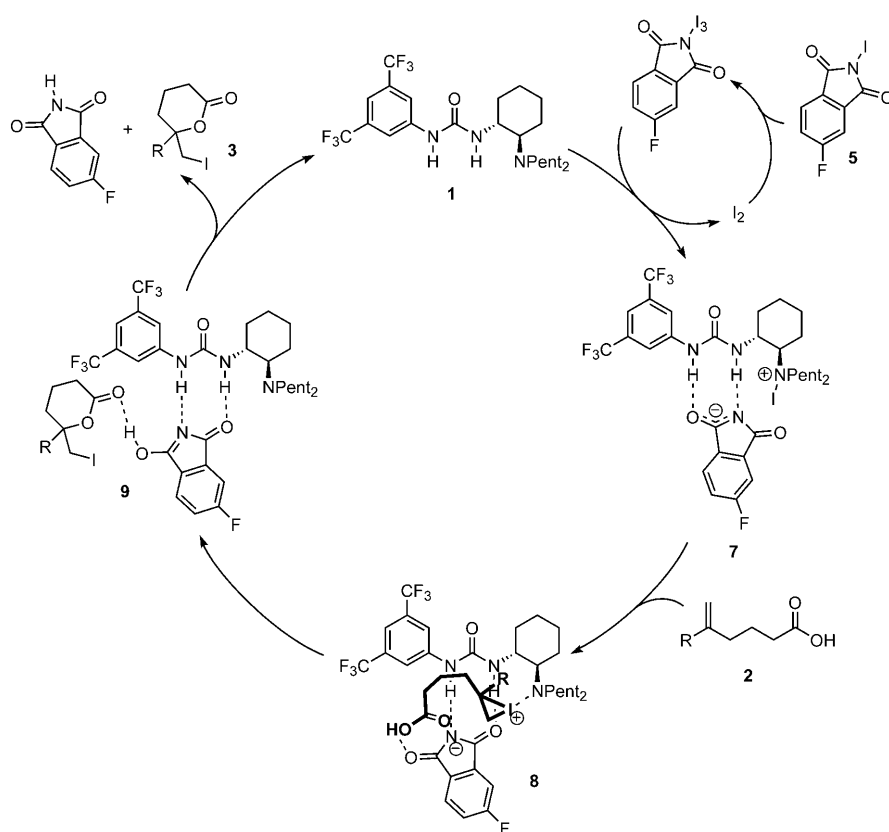
of stoichiometric levels of an *N*-iodoimide derivative and catalytic I_2 was found to produce a high-yielding and highly enantioselective system for iodolactonization (entries 5 and 6). It has been shown recently that *N*-iodoimides undergo conversion to the corresponding triiodide cations upon treatment with I_2 and a protic acid,^[12] and this provides a likely explanation for the synergistic effect of these reagents in the present system. However, increasing the I_2 loading above that of the chiral catalyst (**1**) led to measurable decreases in enantioselectivity (entry 7). Variation of the identity of the *N*-iodoimide resulted in small but measurable changes in the enantioselectivity of the reaction, with *N*-iodo-4-fluorophthalimide derivative **5** proving optimal.^[13] The sensitivity of the product *ee* to the structure of the imide suggests a direct involvement of this counterion in the enantiodetermining step.

Low-temperature 1H NMR studies were performed in an effort to gain insight into the mechanism of the iodolactonization reaction. In the presence of *N*-iodo-4-fluorophthalimide (**5**) and catalytic iodine, catalyst **1** was found to undergo a rapid reaction to yield a compound with spectroscopic and reactivity properties consistent with the *N*-iodo complex **7** (Scheme 1).^[14] Intermediate **7** can be quenched with aqueous sodium thiosulfate to regenerate the starting tertiary amine catalyst **1** as well as the corresponding secondary amine. The

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Scheme 1. Proposed catalytic cycle for the iodolactonization of **2**.

latter is presumably formed from **7** by elimination of HI and subsequent iminium hydrolysis.

A proposal for the mechanism of iodolactonization catalysis is outlined in Scheme 1. Iodonium ion formation from hexenoic acid **2** is presumably induced by *N*-iodo tertiary aminourea **7**. Subsequent cyclization is proposed to take place as the rate- and enantiodetermining step, based on the observation of differing reactivities, under otherwise identical conditions, of pentenoic, hexenoic, and heptenoic acid substrates to generate the corresponding 5-, 6-, and 7-membered lactone products ($5 > 6 \gg 7$). Preliminary computational studies support the intermediacy of iodonium ion complex **8**, which maintains a tertiary amino-iodonium ion interaction.^[15] On the basis of this putative structure, we suggest that urea-bound phthalimide serves as the base to effect deprotonation of the carboxylic acid in the enantiodetermining cyclization event. An alternative mechanism in which deprotonation is induced by the tertiary amino group present in the catalyst is also plausible, although such a mechanism would require significant reorganization of complex **8**.

The optimized reaction conditions developed for the enantioselective iodolactonization of **2a** were applied to a variety of other 5-substituted hexenoic acid derivatives (Scheme 2). In the case of 5-arylhexenoic acids, a clear correlation emerged between the electronic properties of the arene and the observed *ee*, with electron-deficient derivatives undergoing more enantioselective cyclization. For these less

reactive substrates (**3g–3i**), it was necessary to use 2 equivalents of the iodinating agent **5** to achieve useful conversions.

To determine the absolute configuration of **3a**, a radical deiodination was performed to provide the corresponding, known methyl lactone.^[16] This assignment was confirmed by X-ray crystallographic analysis of the 4-bromo derivative **3g**.^[17]

These reaction conditions proved ineffective with the corresponding pentenoic acid substrate **10a** (Table 2, *R* = H), providing the iodolactonization product **11a** in low enantiomeric excess (entry 1). However, enantioselectivities were improved substantially by decreasing the loading of iodine additive, with best results obtained using 0.1 mol% of I_2 (entry 3). The rate of racemic iodolactonization in the absence of **1** is not affected by the concentration of added I_2 , so it appears that the inverse relationship between *ee* and I_2 loading may be due to competing pathways promoted by **1**. In that context, it is

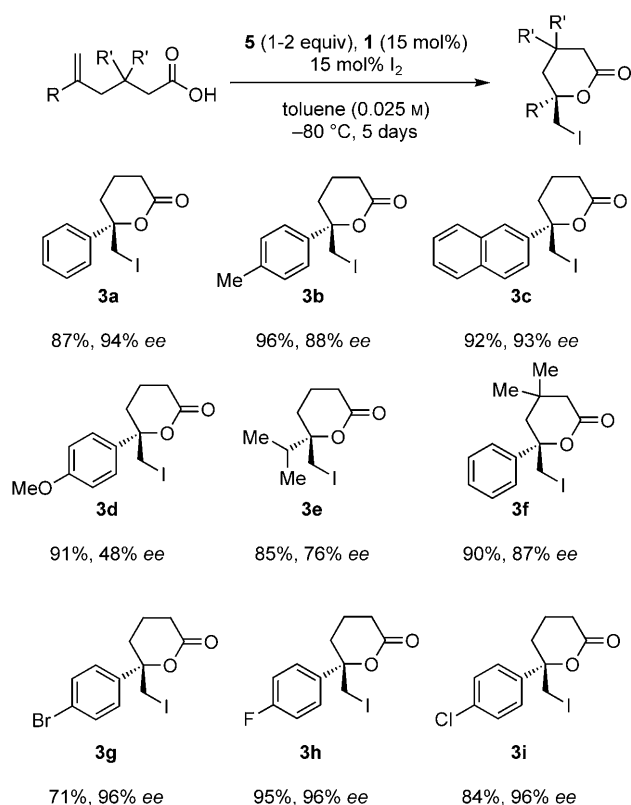
Table 2: Optimization studies for the enantioselective iodolactonization of pentenoic acid derivative **10a**.

Entry ^[a]	Substrate	I_2 Additive [mol %]	Yield [%] ^[b]	<i>ee</i> [%] ^[c]
1	10a	15	86	31
2	10a	1.0	95	76
3	10a	0.1	45	90
4 ^[d]	10a	0.1	82	90
5	10b	0.1	98	0

[a] Reactions performed on a 0.05 mmol scale. [b] Determined by 1H NMR analysis using 1,3,5-trimethoxybenzene as the internal standard. [c] Determined by HPLC analysis using commercial chiral columns. [d] Reaction performed using 2 equivalents of **5**.

interesting to note that the absolute configuration of **10a** was found to be opposite to that of the hexenoic acid cyclization products **3**. The basis for the striking differences in behavior between the pentenoic and hexenoic acid substrates is not understood at this point, but is the subject of current analysis.

Another potentially valuable mechanistic clue is provided by the fact that *gem*-dimethyl-substituted substrate **10b** (*R*' = Me) was found to undergo cyclization to give racemic product



Scheme 2. Substrate scope in the catalytic iodolactonization reaction. Reactions were performed on a 0.2 mmol scale. Yields are of isolated product following purification by column chromatography. Enantiomeric excesses (*ee*) were determined by HPLC or GC (**3e**) analysis on commercial chiral columns. See Supporting Information for full details.

under the conditions optimized for **9a**. Analogous *gem*-dimethyl substitution in the hexenoic acid case also led to significant decrease in *ee* (product **3f**). In general, it was found that any factors expected to lead to more rapid cyclization of the iodonium ion intermediate (**8**, Scheme 1) led to diminished enantioselectivities. If the rates of the two steps (iodonium formation, **7**→**8** and cyclization, **8**→**9**, Scheme 1) are finely balanced, it is possible that acceleration of the second step changes the identity of the rate- and *ee*-determining step to formation of the iodonium ion. Poor face selectivity in the iodination of the alkene would then be responsible for the decreases in enantioselectivity.

In conclusion, under appropriate conditions, tertiary aminourea derivative **1** induces enantioselective iodolactonization reactions to afford 5- and 6-membered iodolactones with high levels of enantioselectivity. Studies to elucidate the mechanism and origin of enantioinduction in this process are underway.

Experimental Section

N-Iodo-4-fluorophthalimide (58 mg, 0.2 mmol) followed by iodine (7.6 mg, 0.03 mmol) were added as solids to a stirred solution of the 5-hexenoic acid (0.2 mmol) and catalyst **1** (15.4 mg, 0.03 mmol) in

toluene (8 mL) at $-80\text{ }^\circ\text{C}$ under a nitrogen atmosphere. After stirring at this temperature for 5 days, the reaction mixture was quenched at $-80\text{ }^\circ\text{C}$ by addition of 10% aqueous sodium thiosulfate solution (4 mL) and partitioned between 1M sodium hydroxide solution (30 mL) and CH_2Cl_2 (30 mL). The organic layer was separated, further washed with 1M sodium hydroxide solution (30 mL), dried ($MgSO_4$), and concentrated in vacuum. Purification by flash column chromatography on silica gel (5–40% ethyl acetate in hexanes) afforded the iodolactone products.

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