

Tapas Paul, William P. Malachowski,* and Jisun Lee. Exploration of the Enantioselective Birch–Cope Sequence for the Synthesis of Carbocyclic Quaternary Stereocenters.

Page 935. The absence of *tert*-butyl alcohol in the Birch reduction–allylation of **10a** and **10b**, an important difference in the procedure for these two compounds, was not highlighted in the Experimental Section. A corrected procedure for each compound is described.

((R)-1-Allyl-5-(allyloxymethyl)-2-methoxycyclohexa-2,5-dienyl)((S)-2-(methoxymethyl)-pyrrolidin-1-yl)methanone (11a). Use of the general procedure with **10a** and potassium, but without *tert*-butyl alcohol, afforded a 68% yield.

((R)-1-Allyl-5-((R)-2-(allyloxy)propyl)-2-methoxycyclohexa-2,5-dienyl)((S)-2-(methoxymethyl)-pyrrolidin-1-yl)methanone (11b). Use of the general procedure with **10b** and potassium, but without *tert*-butyl alcohol, afforded a 57% yield.

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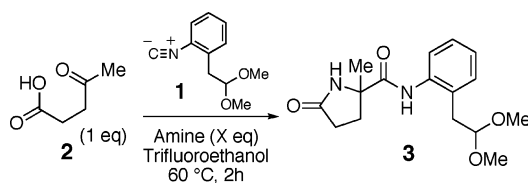
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Jerry Isaacson, Cynthia B. Gilley, and Yoshihisa Kobayashi*. Expedient Access to Unprotected Racemic Pyroglutamic Acids.

Page 3914. Table 1 should be replaced with the following revised one. Two equivalents of isonitrile **1** is used for entry 3.

TABLE 1. Screening of Amine and the Reaction Condition of Ugi Reaction



entry	amine	X (equiv)	isonitrile 1 (equiv)	yield (%) ^a
1	NH ₄ •OAc	1.1	1.1	61
2	NH ₄ •OAc	2	1.1	77
3	NH ₄ •OAc	2	2	82
4 ^b	NH ₄ •OAc	2	1.1	84
5	HMDS ^c	2	1.1	72

^a Isolated yield (1 mmol scale). ^b MS4Å was added (20 mg/mmol). ^c HMDS = hexamethyldisilazane.

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