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Addition/Correction

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Correction to “Organocatalytic Enantioselective Intramolecular Oxa-Michael Reaction of Enols: Synthesis of Chiral Isochromene”

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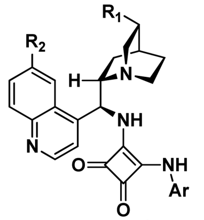
The catalyst structure in Table 1 was omitted from the final publication. The full table is provided below.

Table 1. Optimization of the Reaction Conditions

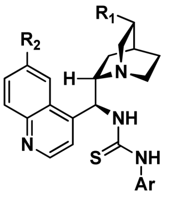
entry	2	solvent	temp (°C)	time (h)	yield[b] (%)	ee[c] (%)
1	**2a**	1,2-DCE	rt	4	3	79
2	**2b**	**1,2-DCE**	**rt**	**4**	**42**	**94**
3	**2c**	1,2-DCE	rt	4	16	91
4	**2d**	1,2-DCE	rt	4	12	78
5	**2e**	1,2-DCE	rt	4		
6	**2f**	1,2-DCE	rt	4	21	87
7	**2g**	1,2-DCE	rt	4	10	89
8	**2b**	toluene	rt	4	29	94
9	**2b**	PhCl	rt	4	38	92
10	**2b**	PhH	rt	4	36	87
11	**2b**	CH_3CN	rt	4	22	81
12	**2b**	F_3CCH_2OH[f]	rt	4		
13	**2b**[d]	1,2-DCE	rt	4	9	94
14	**2b**[e]	1,2-DCE	rt	4	43	86
15	**2b**	1,2-DCE	10	4	36	92

[a] All of the reactions are carried out on 0.02 mmol scale. [b] The conversion was calculated on the basis of ^1H NMR spectroscopy of the crude reaction mixture using anisole as the internal standard. [c] Enantiomeric excess were determined by HPLC analysis on a chiral stationary phase. [d] 5 mol % catalyst was used. [e] 15 mol % catalyst was used. [f] CF_3CH_2OH is non-nucleophilic, as MeOH gave the exoacetal.[12c]

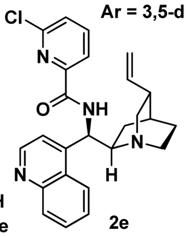
Ar = 3,5-di CF_3-C_6H_3



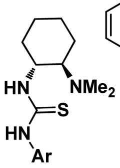
2a, R_1 = Et, R_2 = OMe
2b, R_1 = C_2H_3, R_2 = OMe



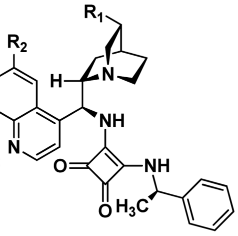
2c, R_1 = C_2H_3, R_2 = H
2d, R_1 = Et, R_2 = OMe



2e



2f



2g, R_1 = C_2H_3, R_2 = OMe

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