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Synthesis and application of dual chiral [2.2] paracyclophane-based N-heterocyclic carbene in enantioselective β -boration of acyclic enones

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1. General information22. Copies of ¹H NMR and ¹³C NMR Spectrums33. HPLC spectra7

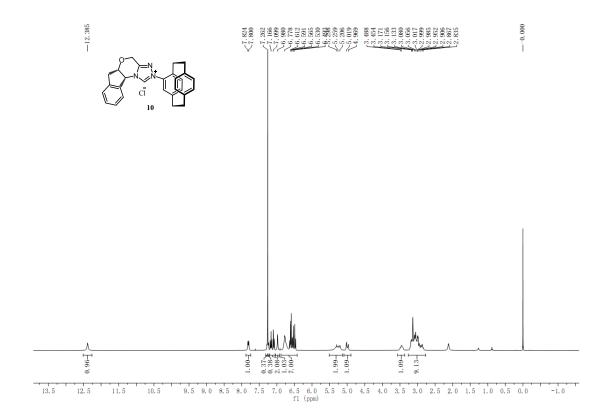
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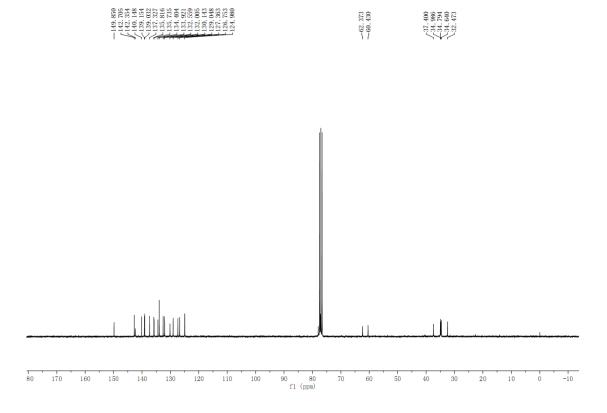
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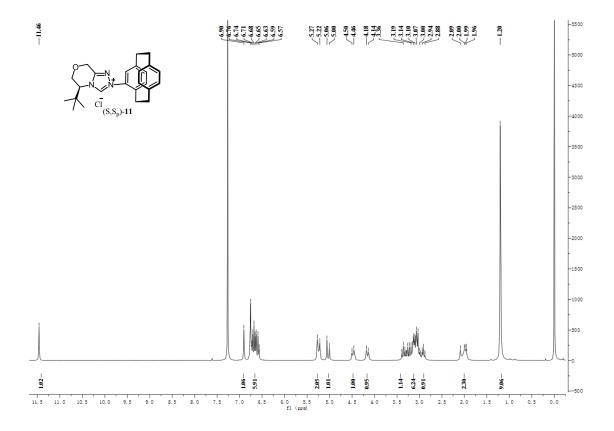
[†] These authors contributed equally to this work.

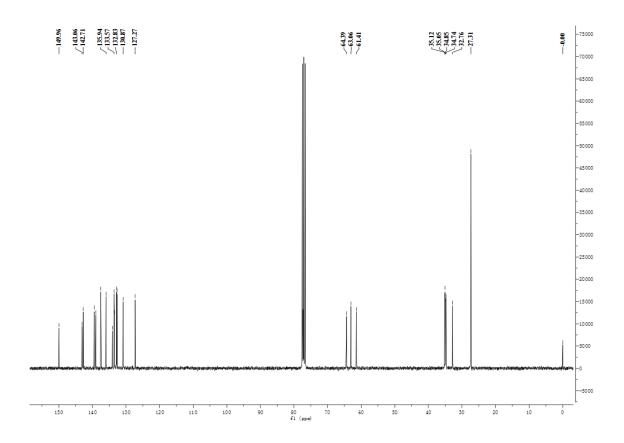
1.General information

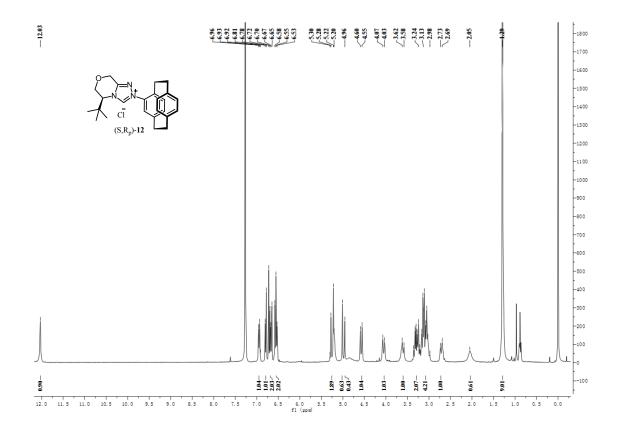
All manipulations of oxygen- and moisture-sensitive materials were conducted with a standard Schlenk technique under a purified argon atmosphere. Solvents were reagent grade and purified by standard techniques. ^{1}H NMR and ^{13}C NMR spectra were recorded on a 300 MHz spectrometer. Chemical shifts were reported as δ values in ppm and referenced to using tetramethylsilane as an internal. All reactions were monitored by TLC with silica gel-coated plates and visualized with a UV light at 254 nm. Optical rotations were reported as follows: $[\alpha]_{D}^{T}$ (c = g/100 mL, solvent).

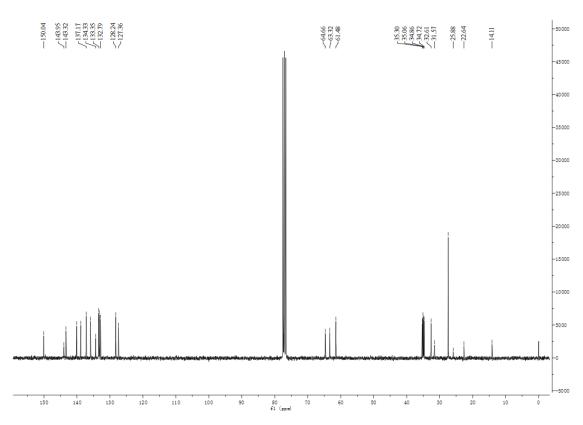


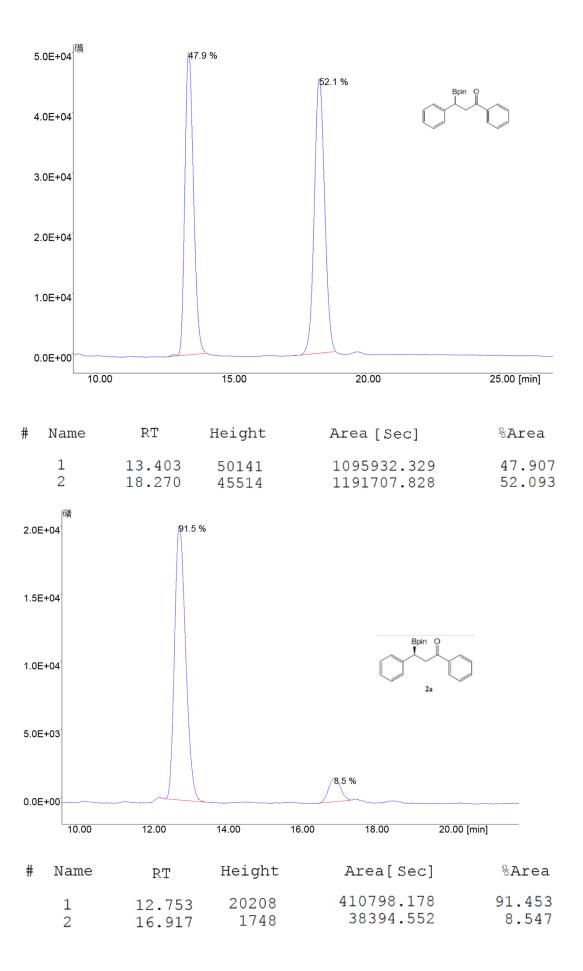


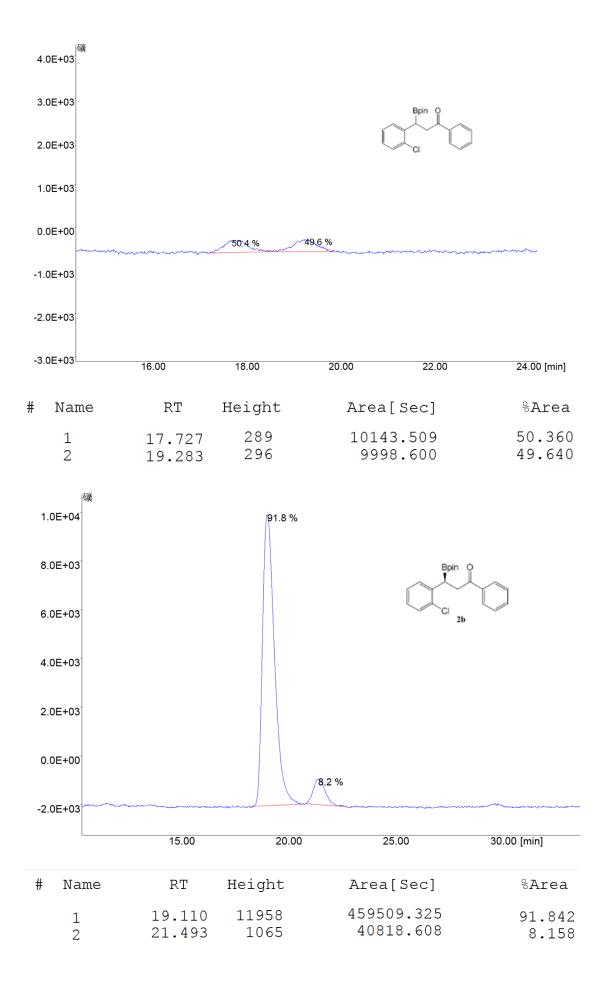


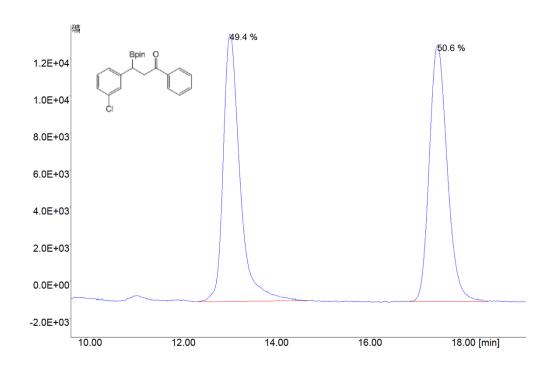




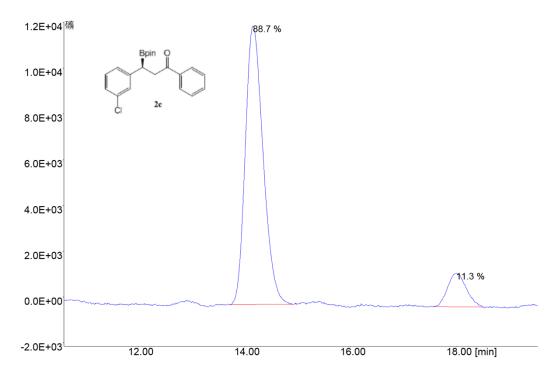




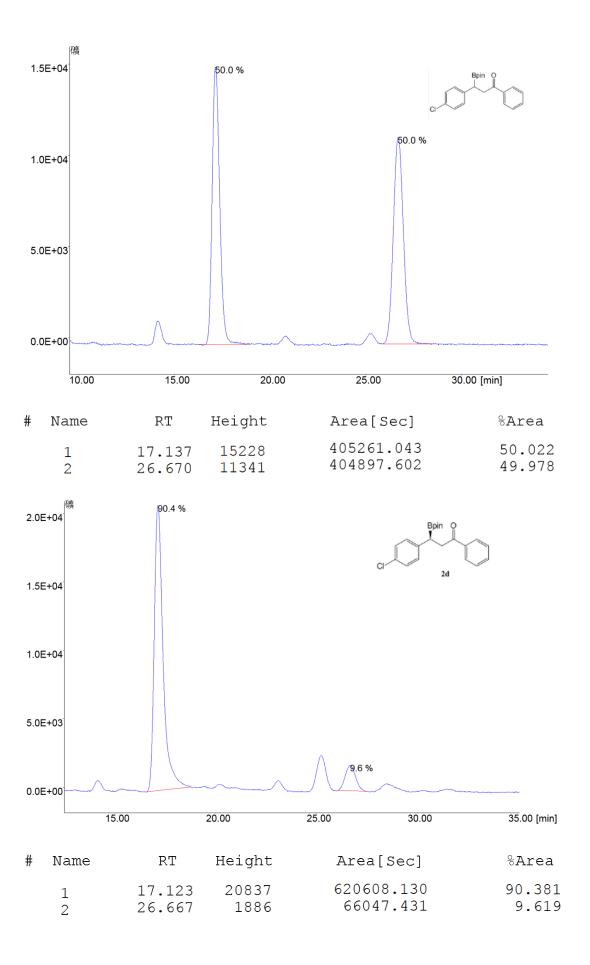


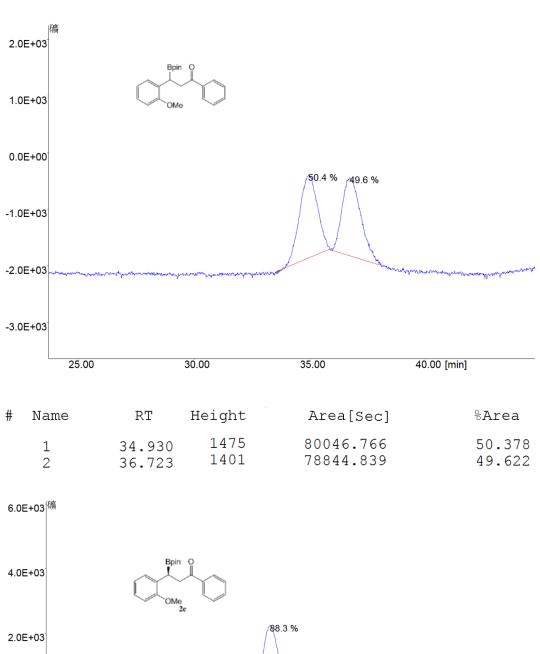


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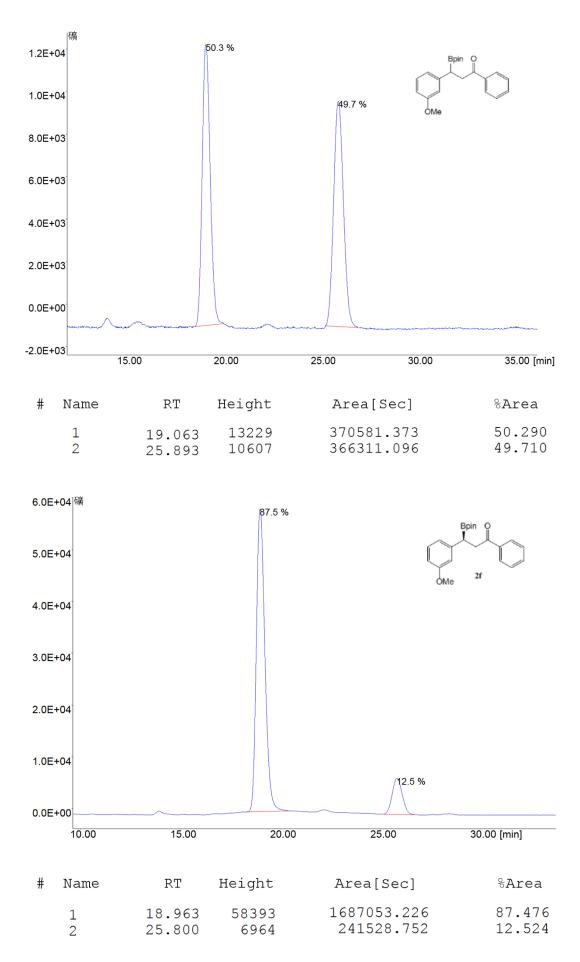


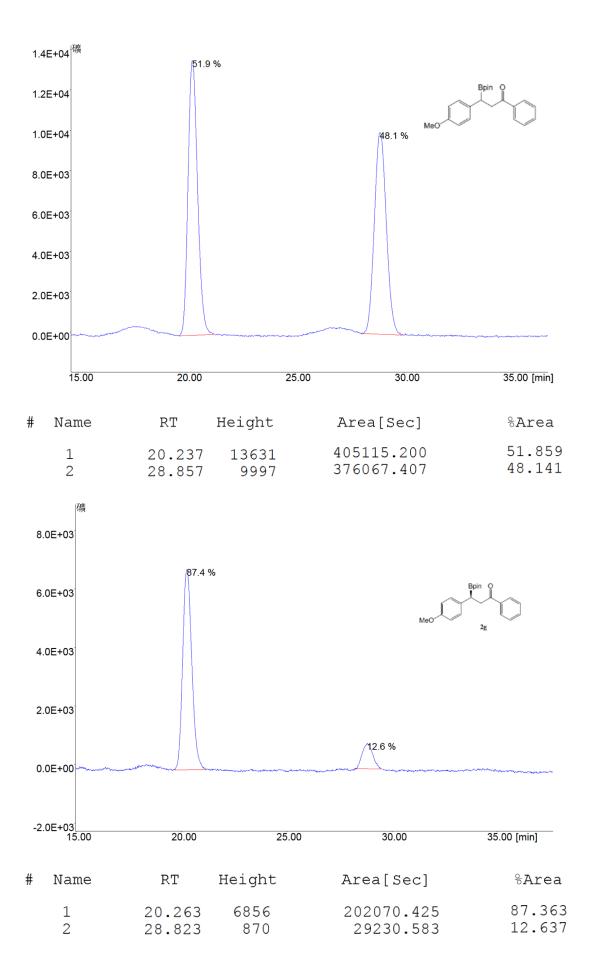
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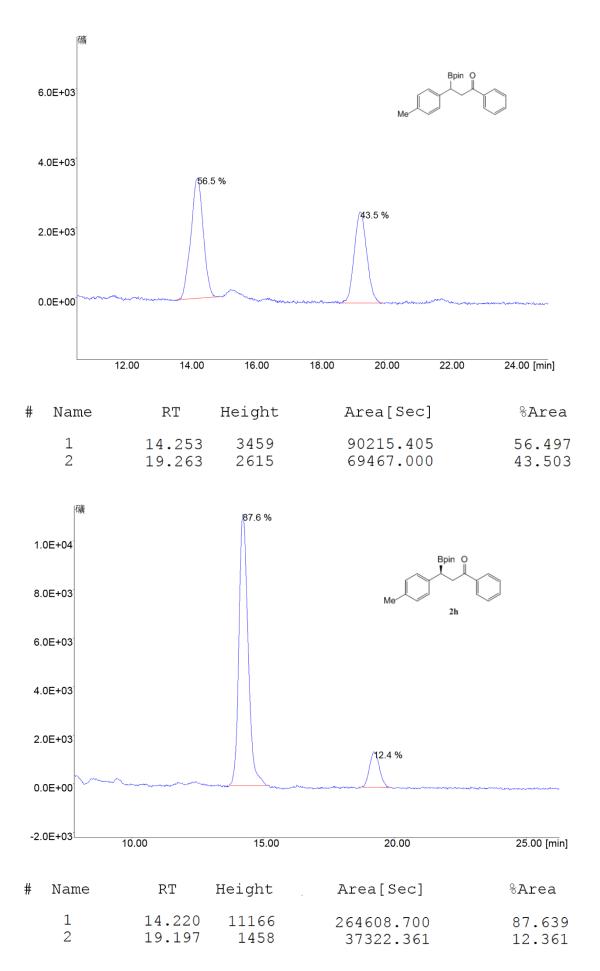


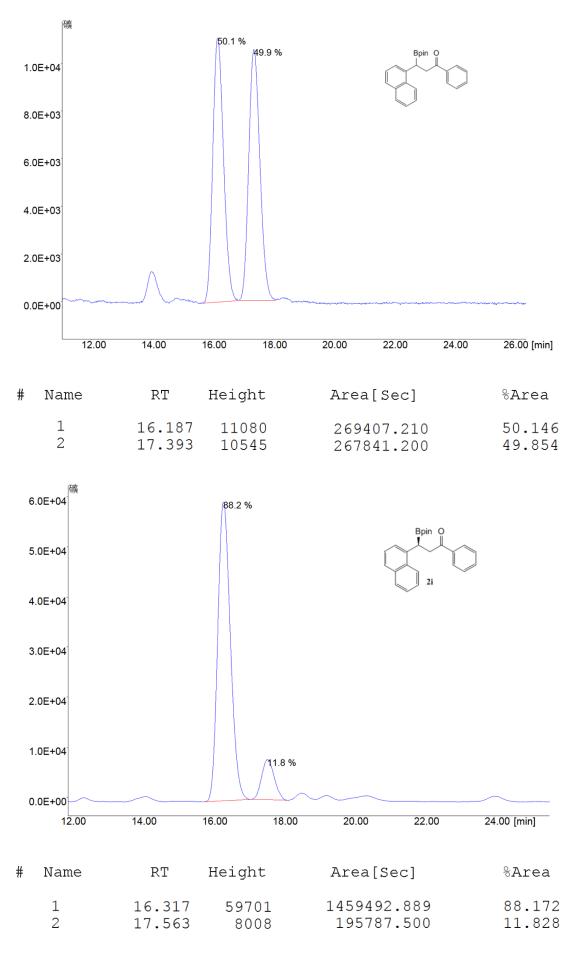


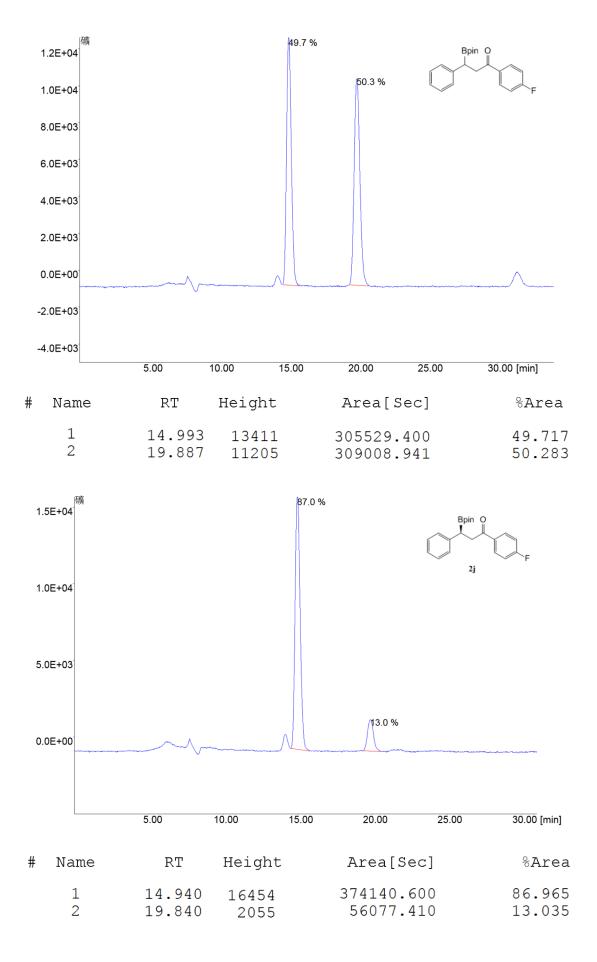
4.0E+	-03		Bpin OMe 2e						
2.0E+	-03				88.3 9	%			
0.0E+	-00					11.7 %			
-2.0E+	-03	and the second seco	aggerten er Arbeitage (1900) e	and the same of th				nganakhinghangunganhgaghtai	
		28.00	30.00	32.00	34.00	36.00	38.00	40.00	42.00 [min]
#	Name		RT	Height	t	Area[[Sec]		%Area
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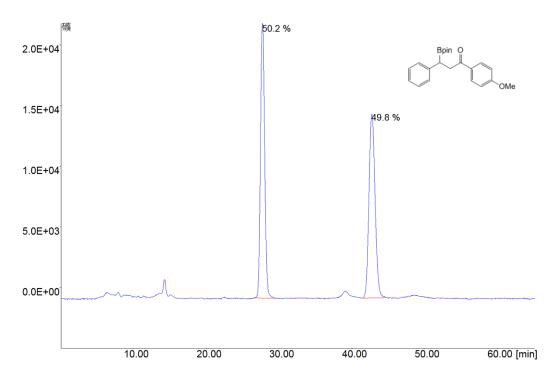




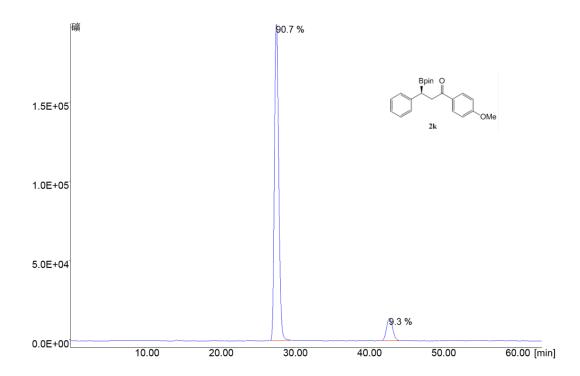




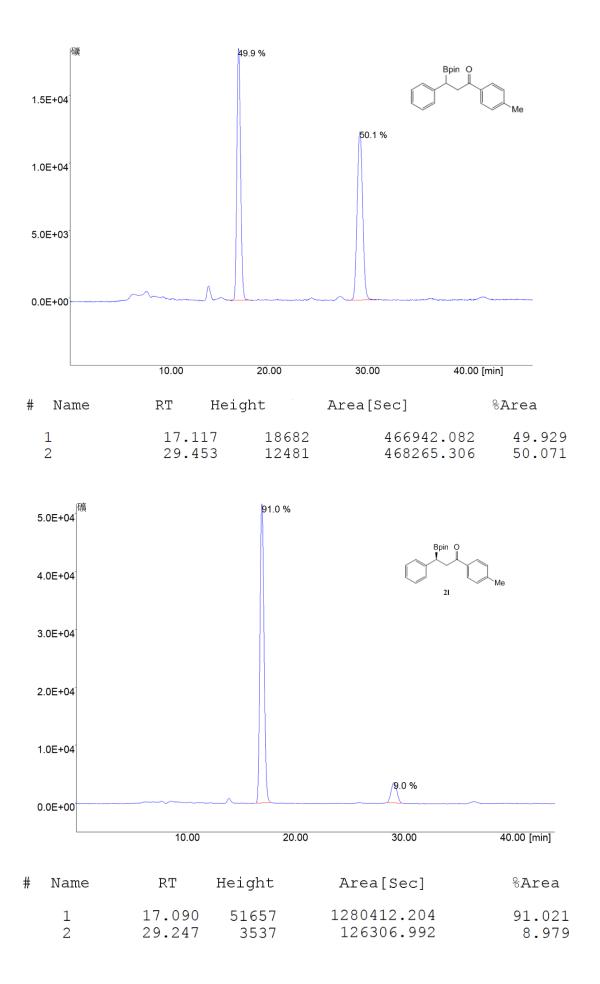


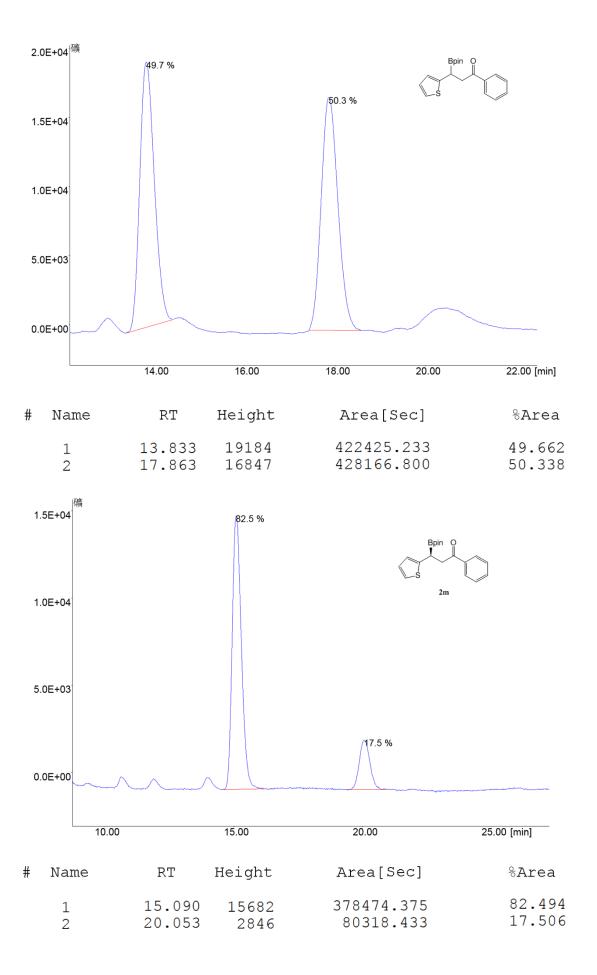


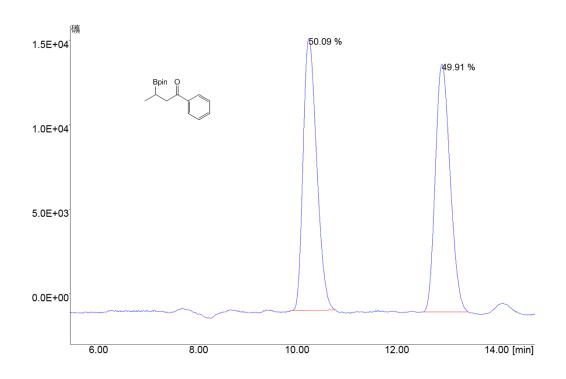
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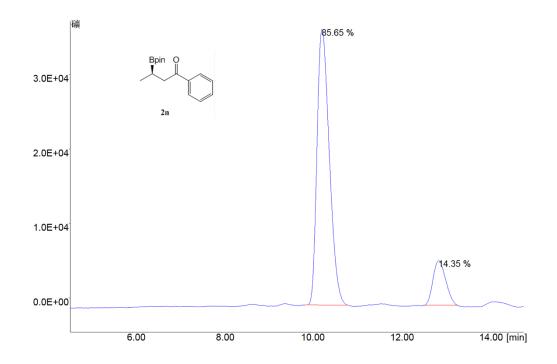
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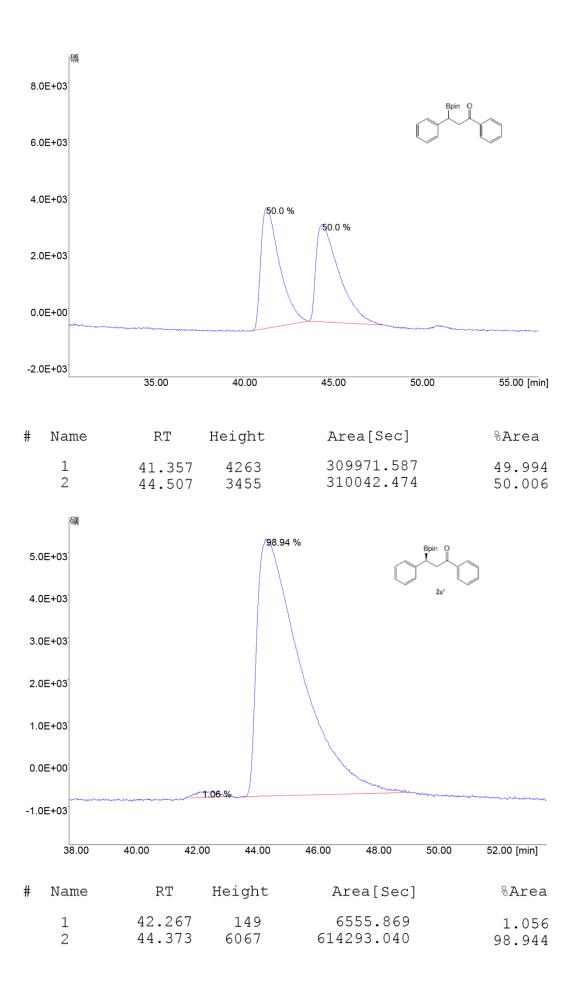


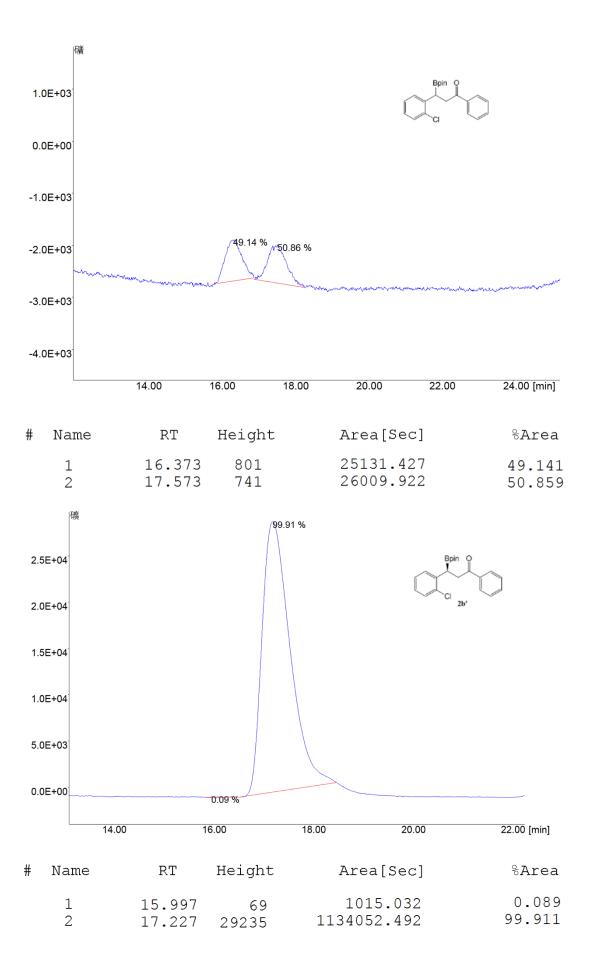


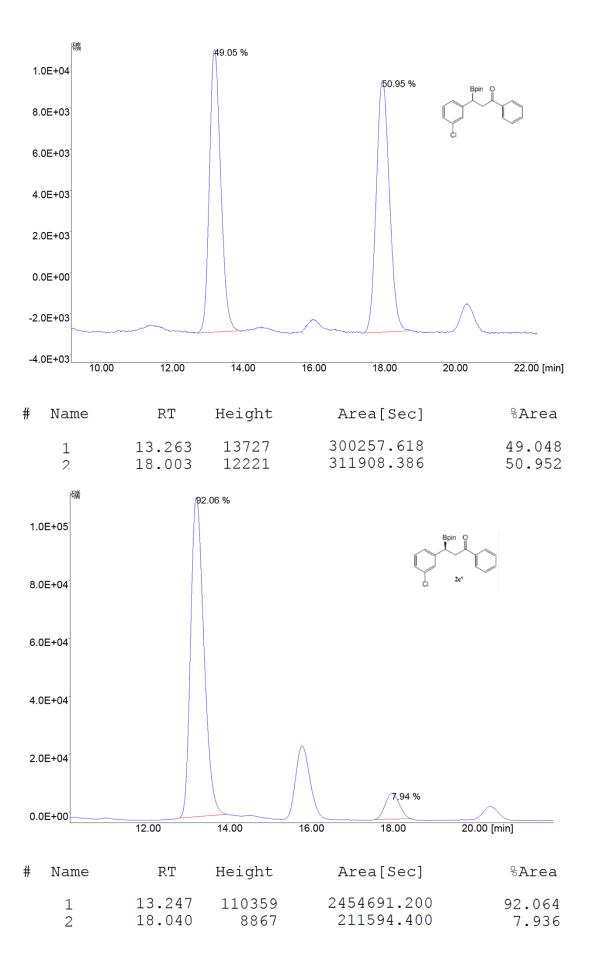
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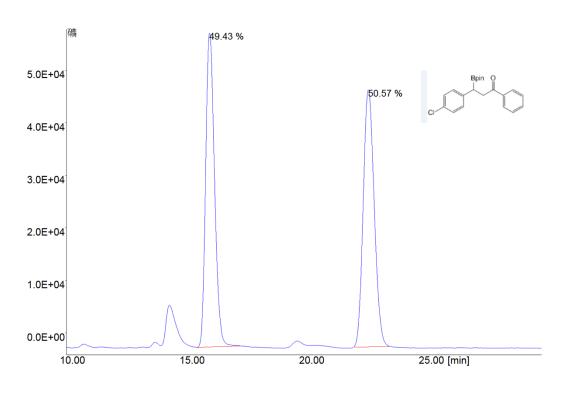


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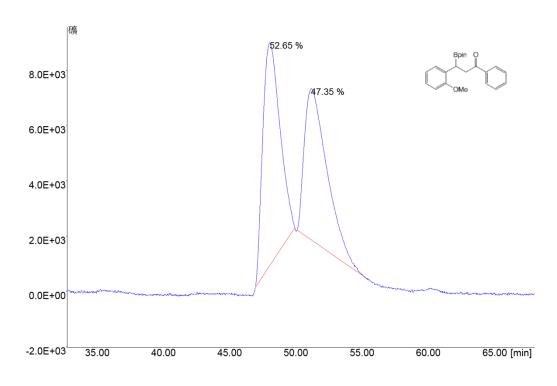




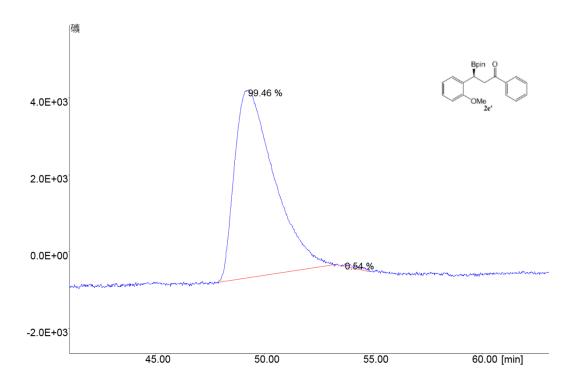




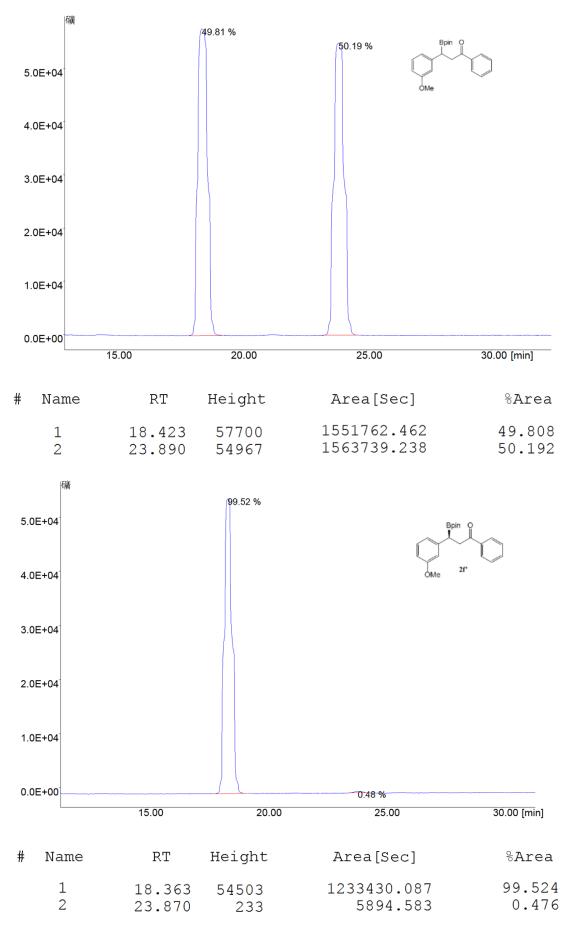
#	Name		RT	Heig	ht	Are	a[Sec]		%Area
	1 2		15.830 22.460	597 489			8430.10 8060.73		49.429 50.571
8.0	礦 DE+04			92.63 %				Bpin O	
6.0	DE+04						Q/	2d'	
4.0	0E+04								
2.0	DE+04								
0.0	0E+00						7.37	%	
		12.00	14.00	16.00	18.00	20.00	22.00	24.00	26.00 [min]
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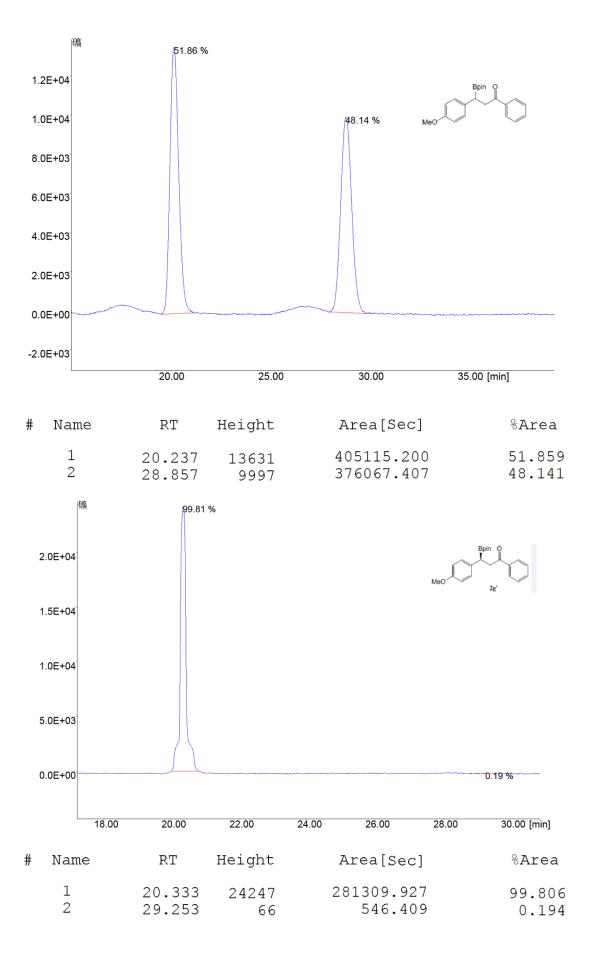


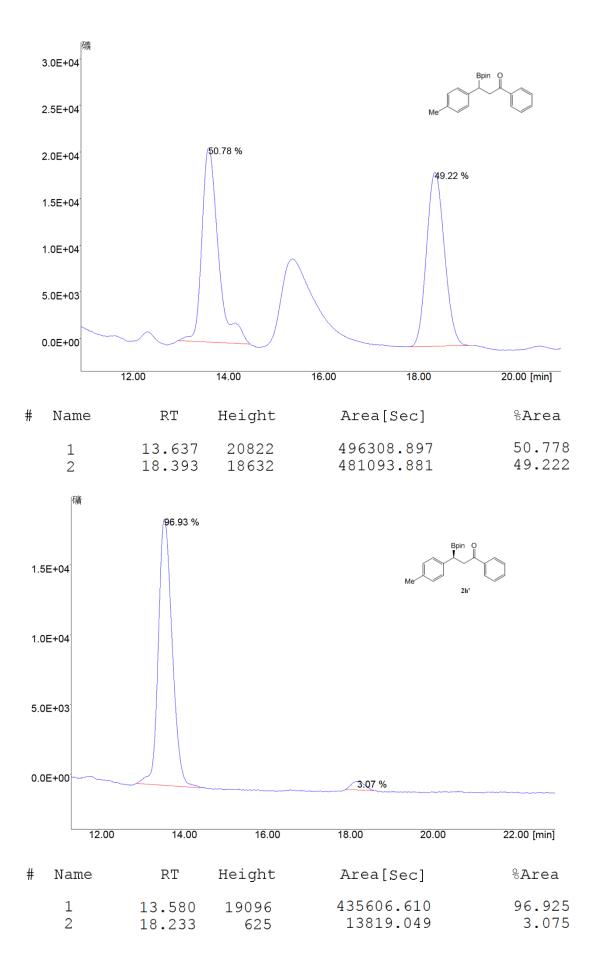
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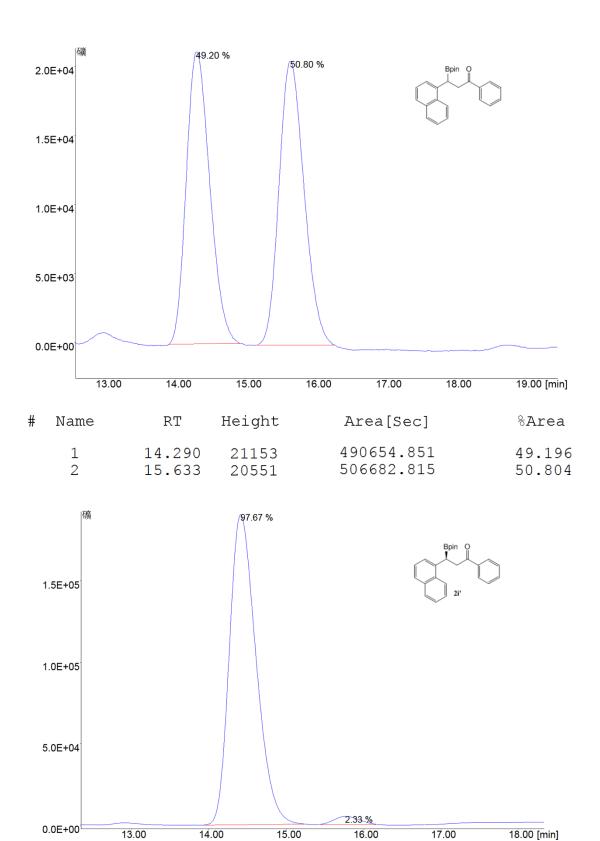


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