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APPENDICES

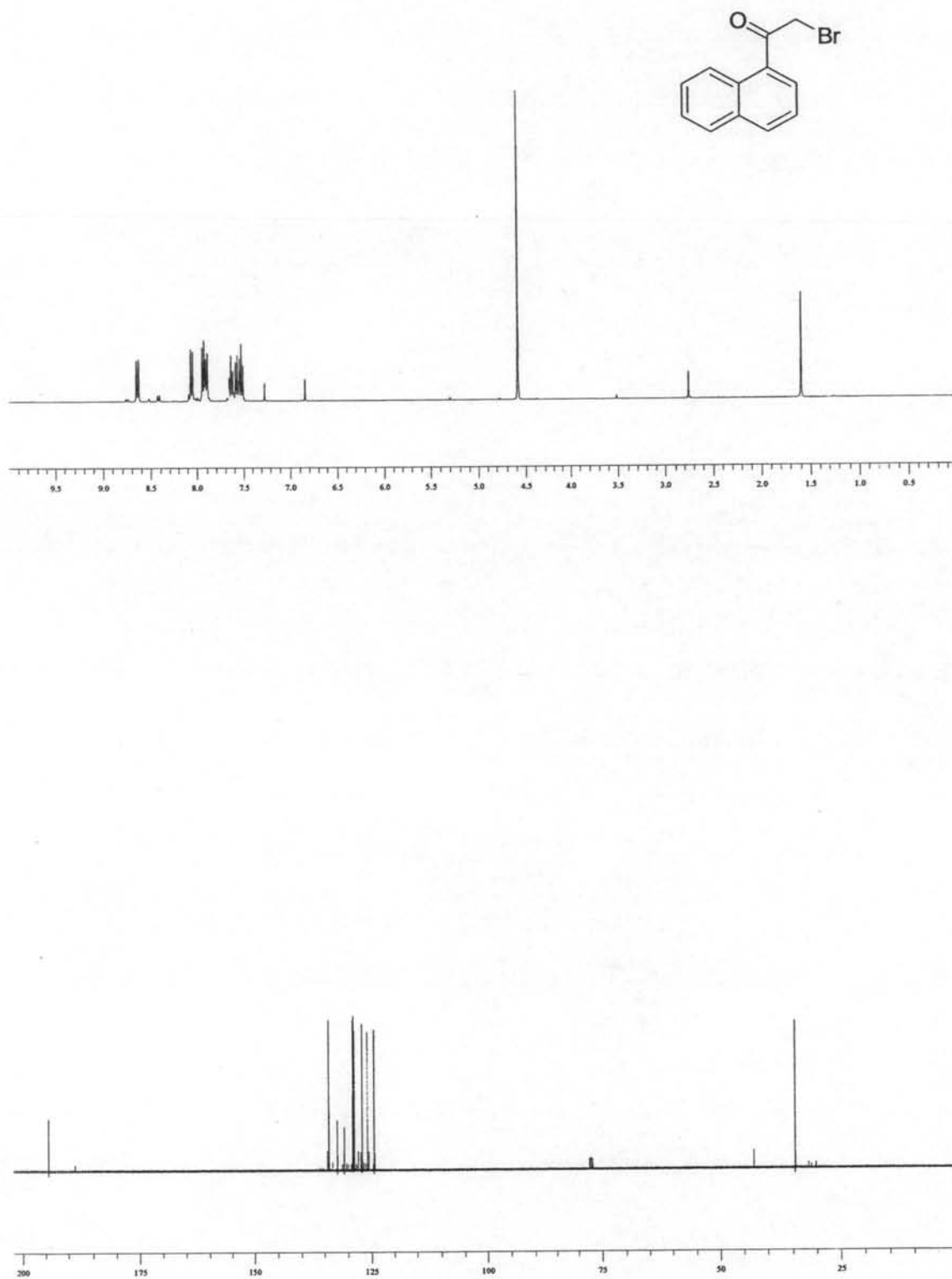


Figure A.1 ^1H NMR (CDCl_3 , 400 MHz) and ^{13}C NMR (CDCl_3 , 100 MHz) spectra of 2-bromo-1-(naphthalen-1-yl)ethanone (**42a**).

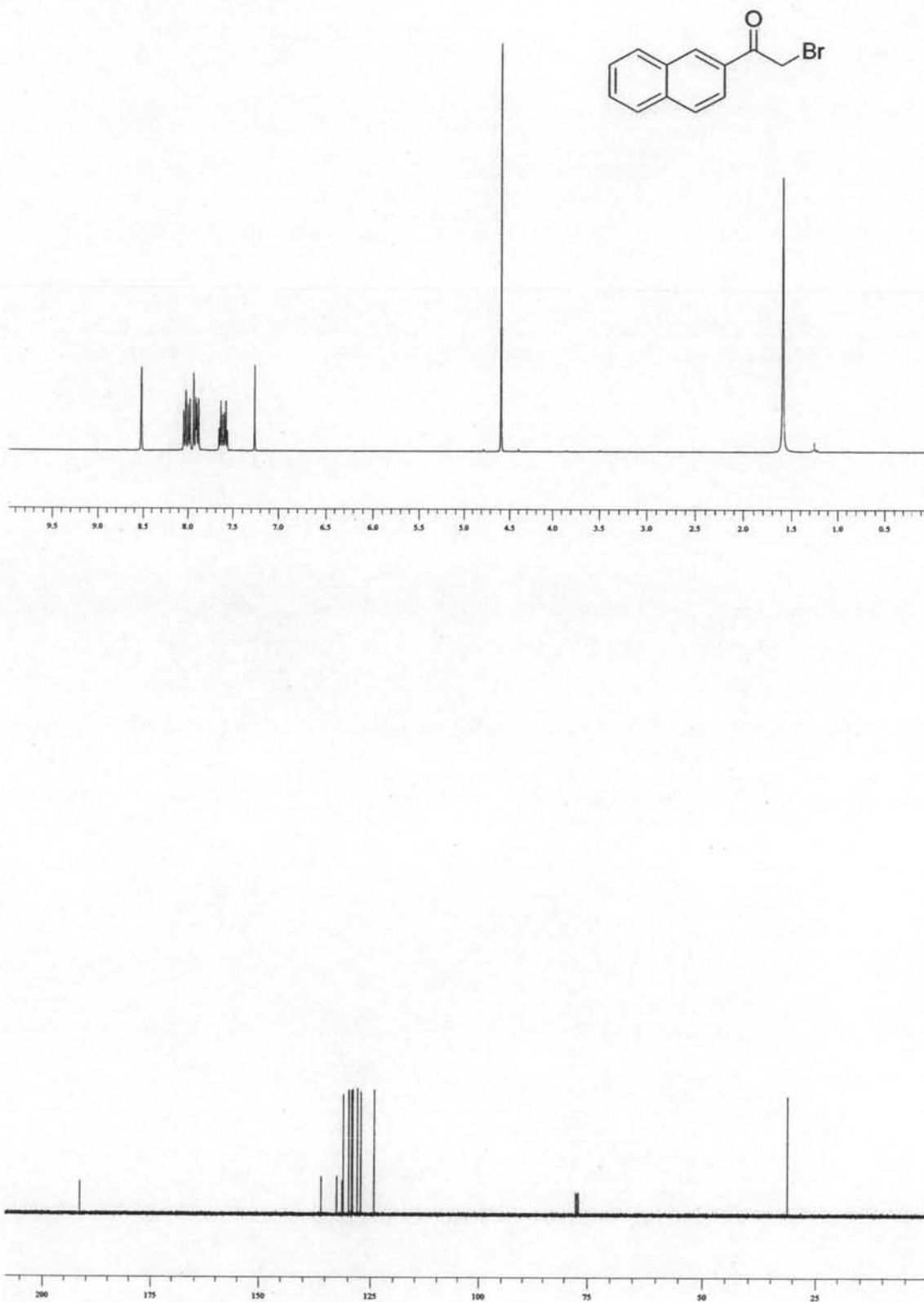


Figure A.2 ^1H NMR (CDCl_3 , 400 MHz) and ^{13}C NMR (CDCl_3 , 100 MHz) spectra of 2-bromo-1-(naphthalene-2-yl)ethanone (**42b**).

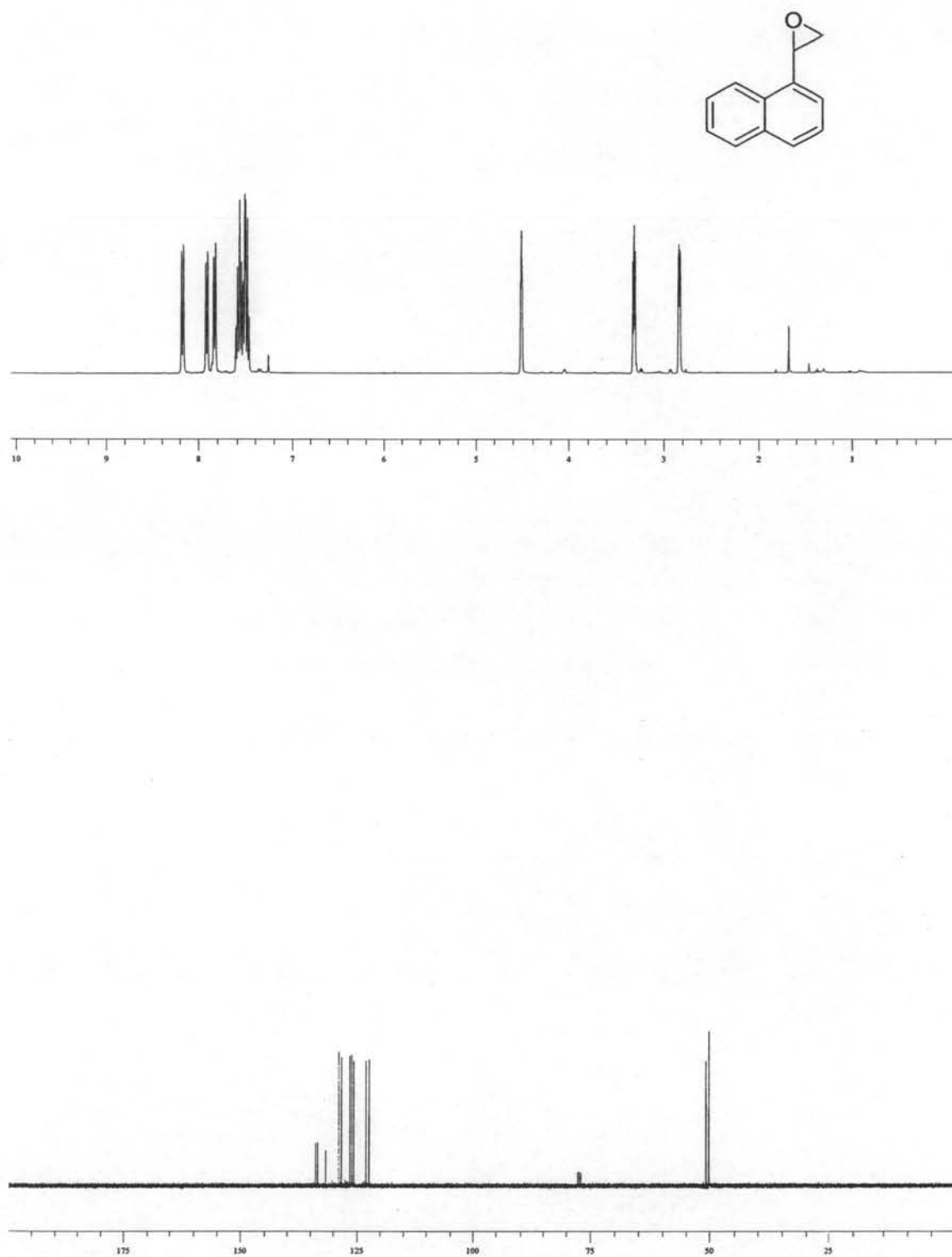


Figure A.3 ^1H NMR (CDCl_3 , 400 MHz) and ^{13}C NMR (CDCl_3 , 100 MHz) spectra of racemic-2-(naphthalen-1-yl)oxirane (**43a**) and (*R*)-2-(naphthalen-1-yl)oxirane (**44a**)

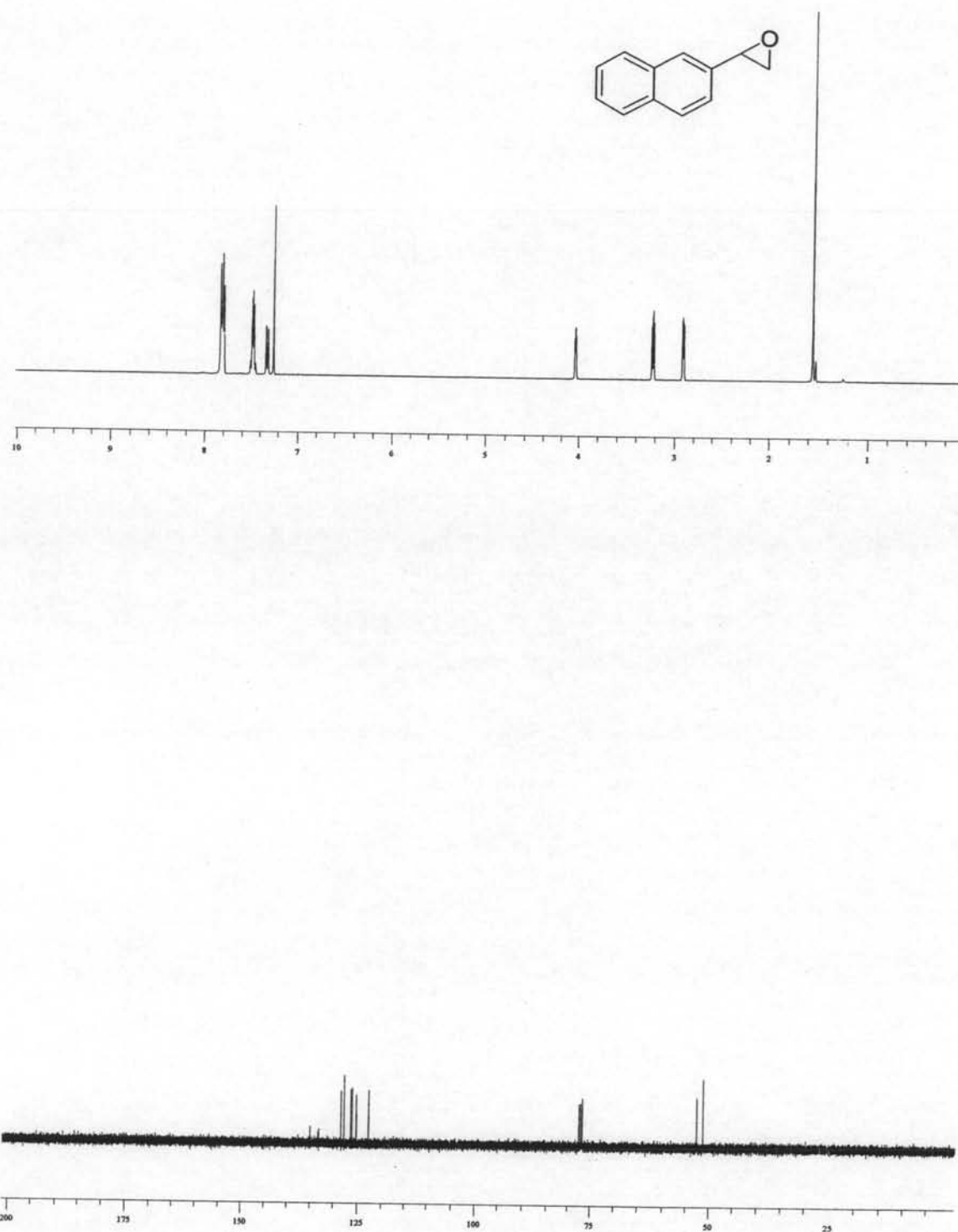


Figure A.4 ^1H NMR (CDCl_3 , 400 MHz) and ^{13}C NMR (CDCl_3 , 100 MHz) spectra of racemic-2-(naphthalen-2-yl)oxirane (**43b**) and (*R*)-2-(naphthalen-2-yl)oxirane (**44b**)

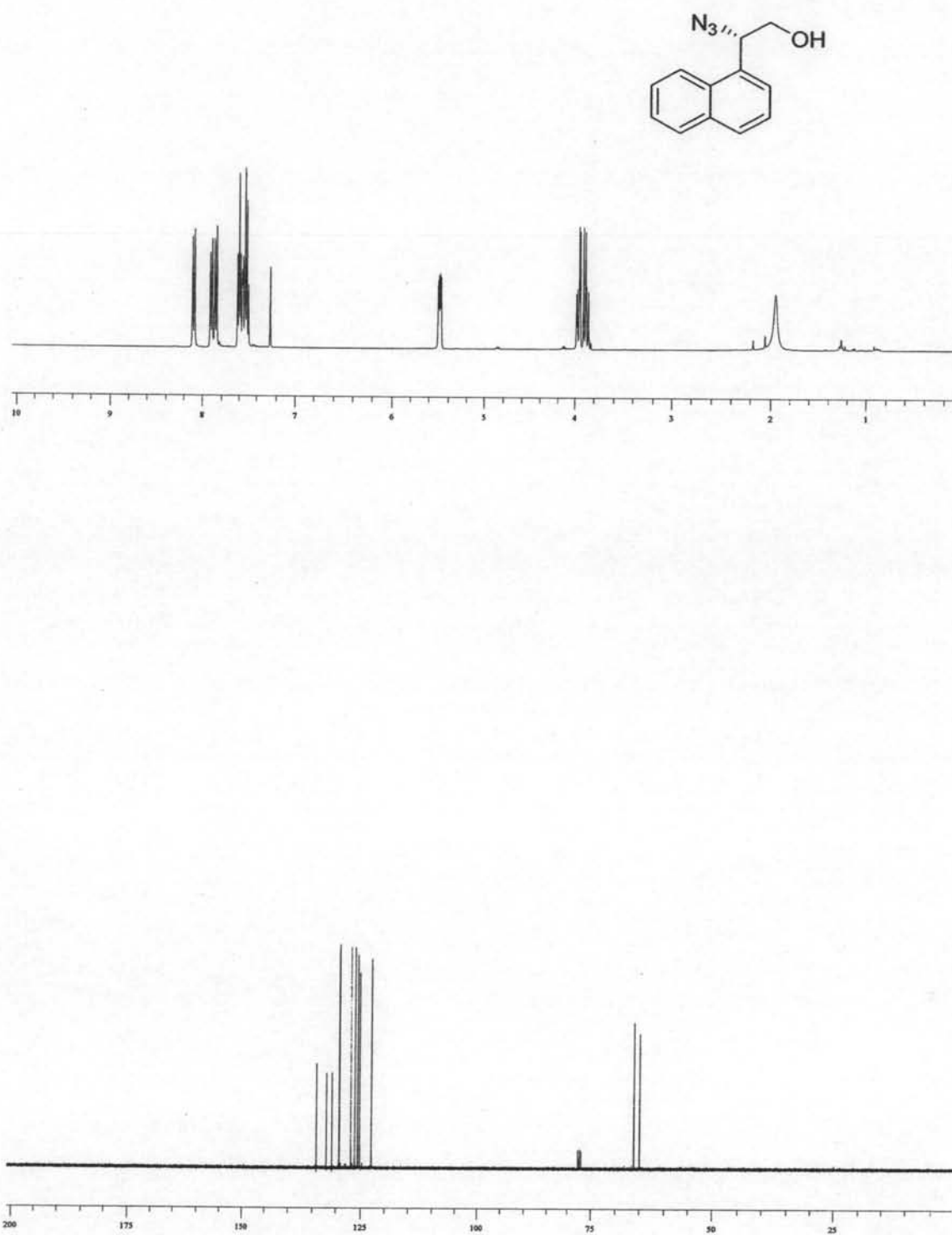


Figure A.5 ^1H NMR (CDCl_3 , 400 MHz) and ^{13}C NMR (CDCl_3 , 100 MHz) spectra of (*S*)-2-azido-2-(naphthalen-1-yl)ethanol (**45a**)

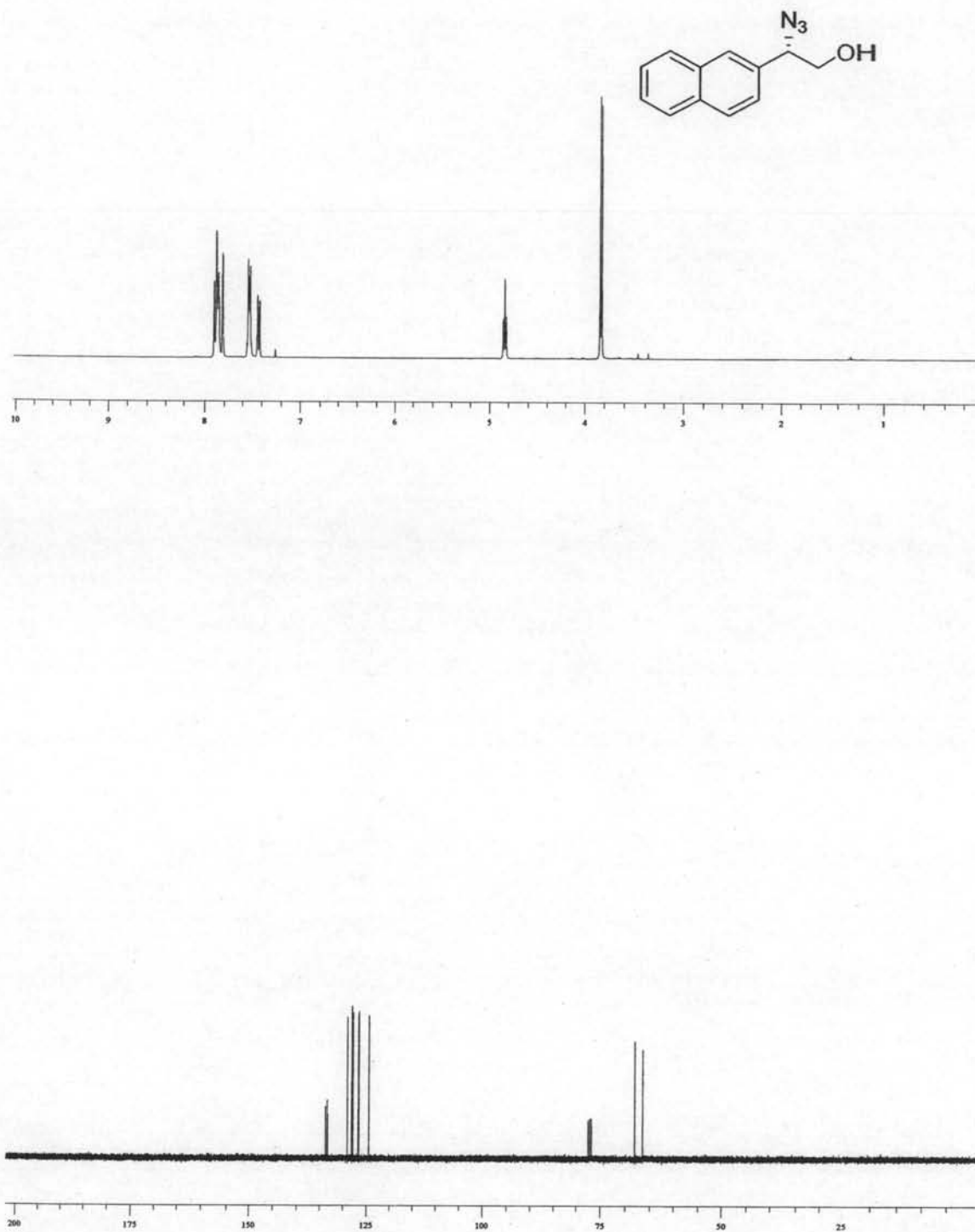


Figure A.6 ¹H NMR (CDCl₃, 400 MHz) and ¹³C NMR (CDCl₃, 100 MHz) spectra of (*S*)-2-azido-2-(naphthalen-2-yl)ethanol (**45b**)

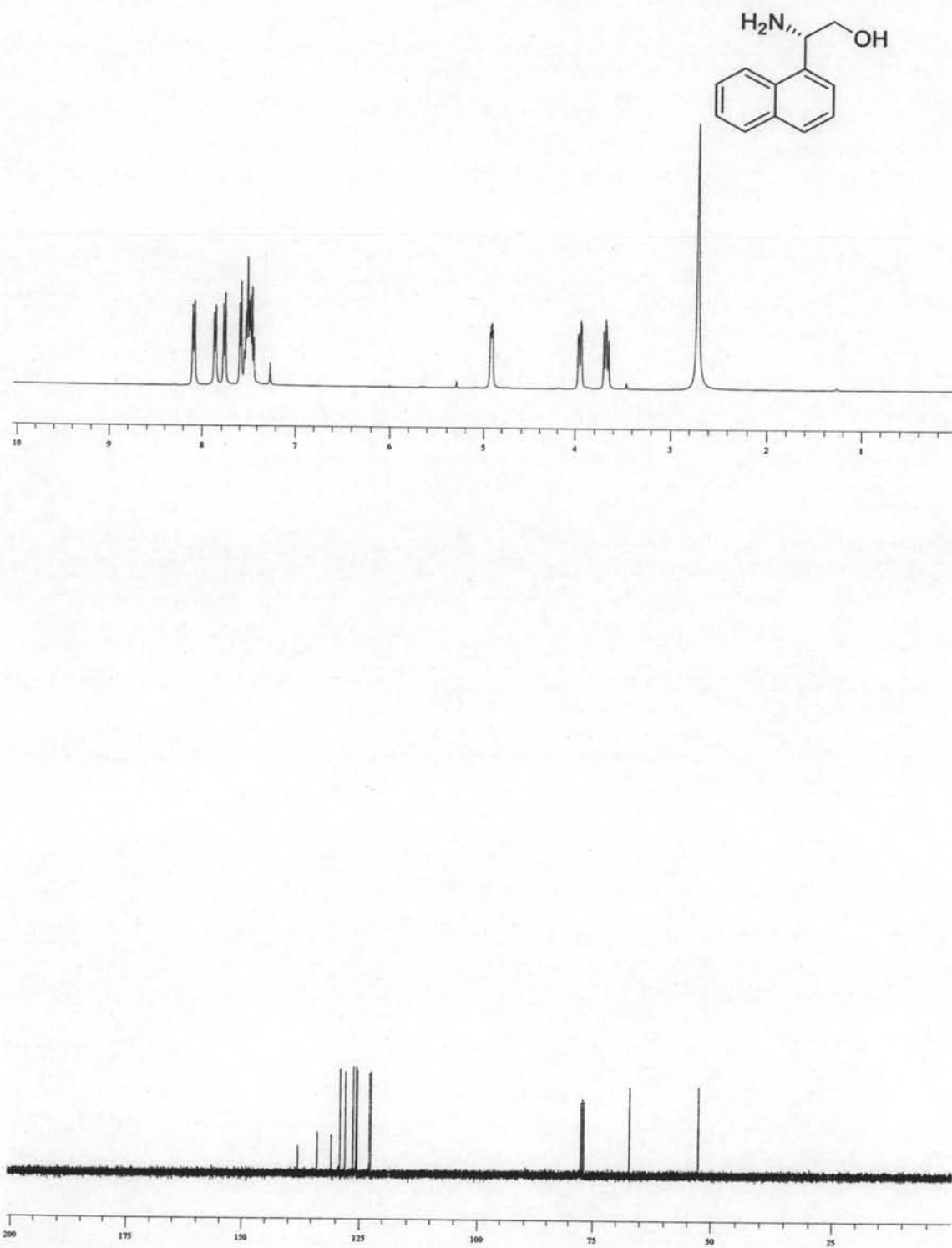


Figure A.7 ^1H NMR (CDCl_3 , 400 MHz) and ^{13}C NMR (CDCl_3 , 100 MHz) spectra of (*S*)-2-amino-2-(naphthalen-1-yl)ethanol (**46a**)

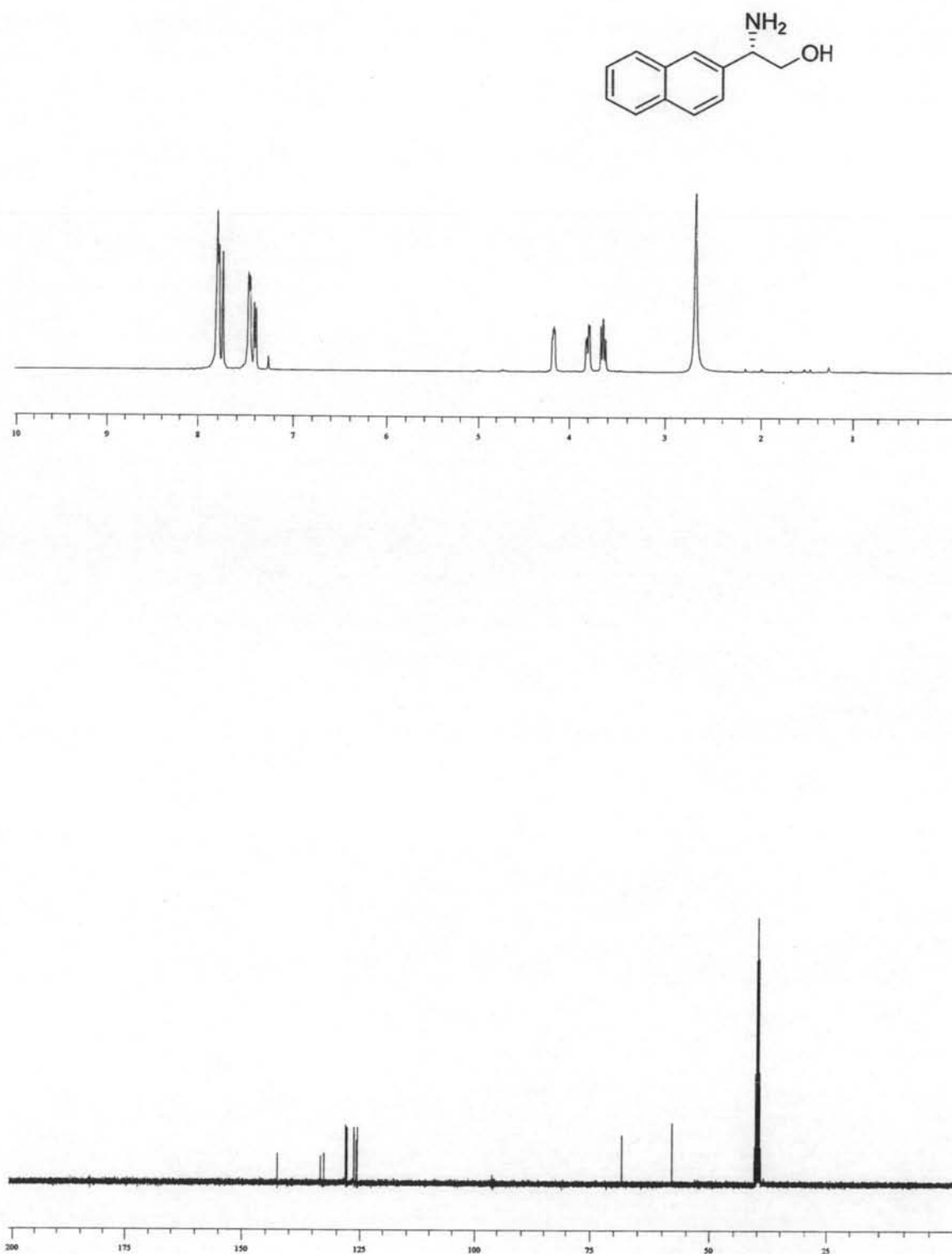


Figure A.8 ^1H NMR (CDCl_3 , 400 MHz) and ^{13}C NMR (CDCl_3 , 100 MHz) spectra of (*S*)-2-amino-2-(naphthalen-2-yl)ethanol (**46b**)

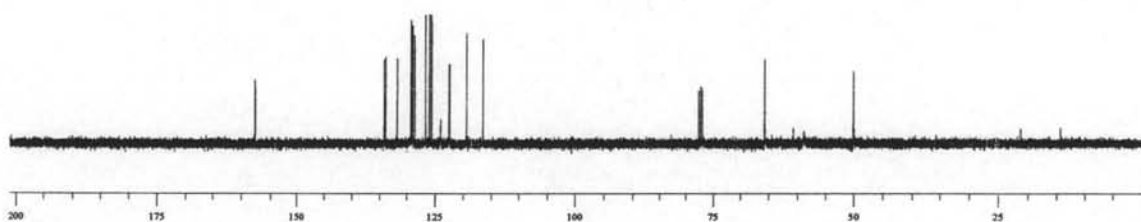
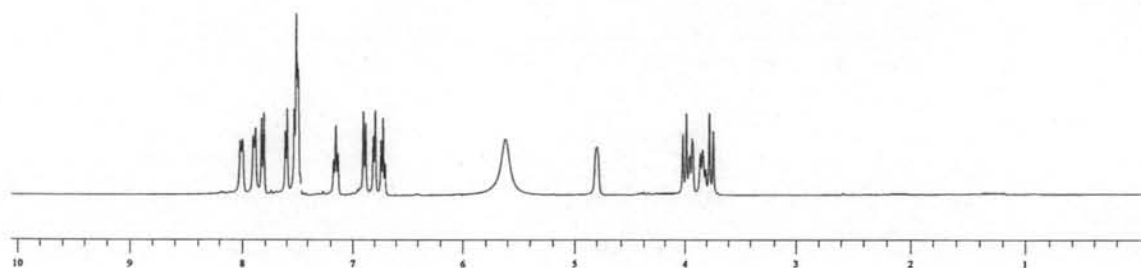
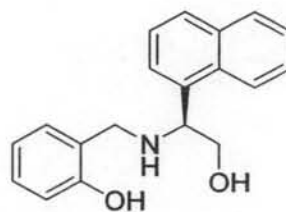


Figure A.9 ^1H NMR (CDCl_3 , 400 MHz) and ^{13}C NMR (CDCl_3 , 100 MHz) spectra of (*S*)-2-((2-hydroxy-1-(naphthalen-1-yl)ethylamino)methyl)phenol (**47a**)

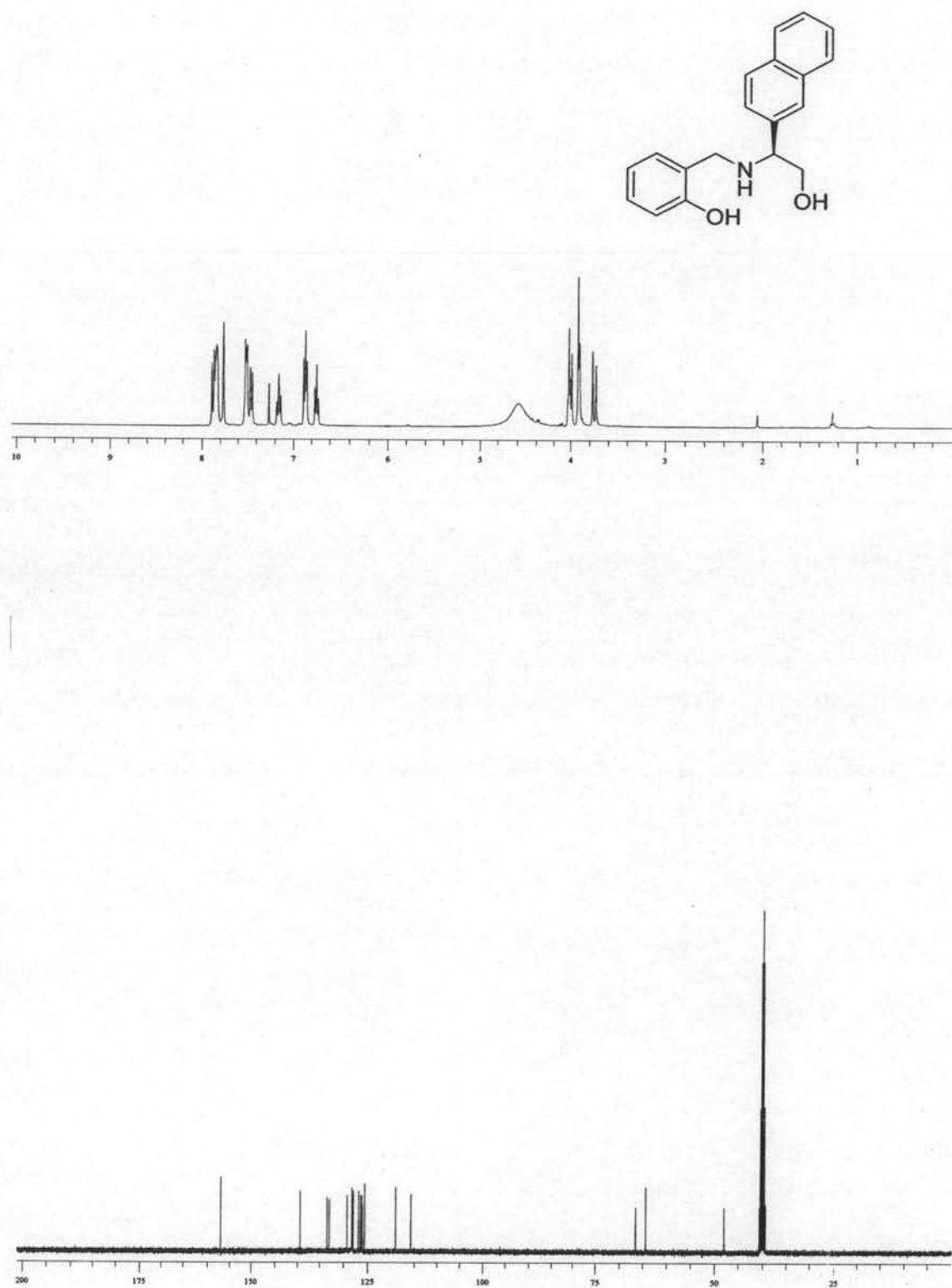


Figure A.10 ^1H NMR (CDCl_3 , 400 MHz) and ^{13}C NMR ($\text{DMSO-}d_6$, 100 MHz,) spectra of (*S*)-2-((2-hydroxy-1-(naphthalen-2-yl)ethylamino)methyl)phenol (**47b**)

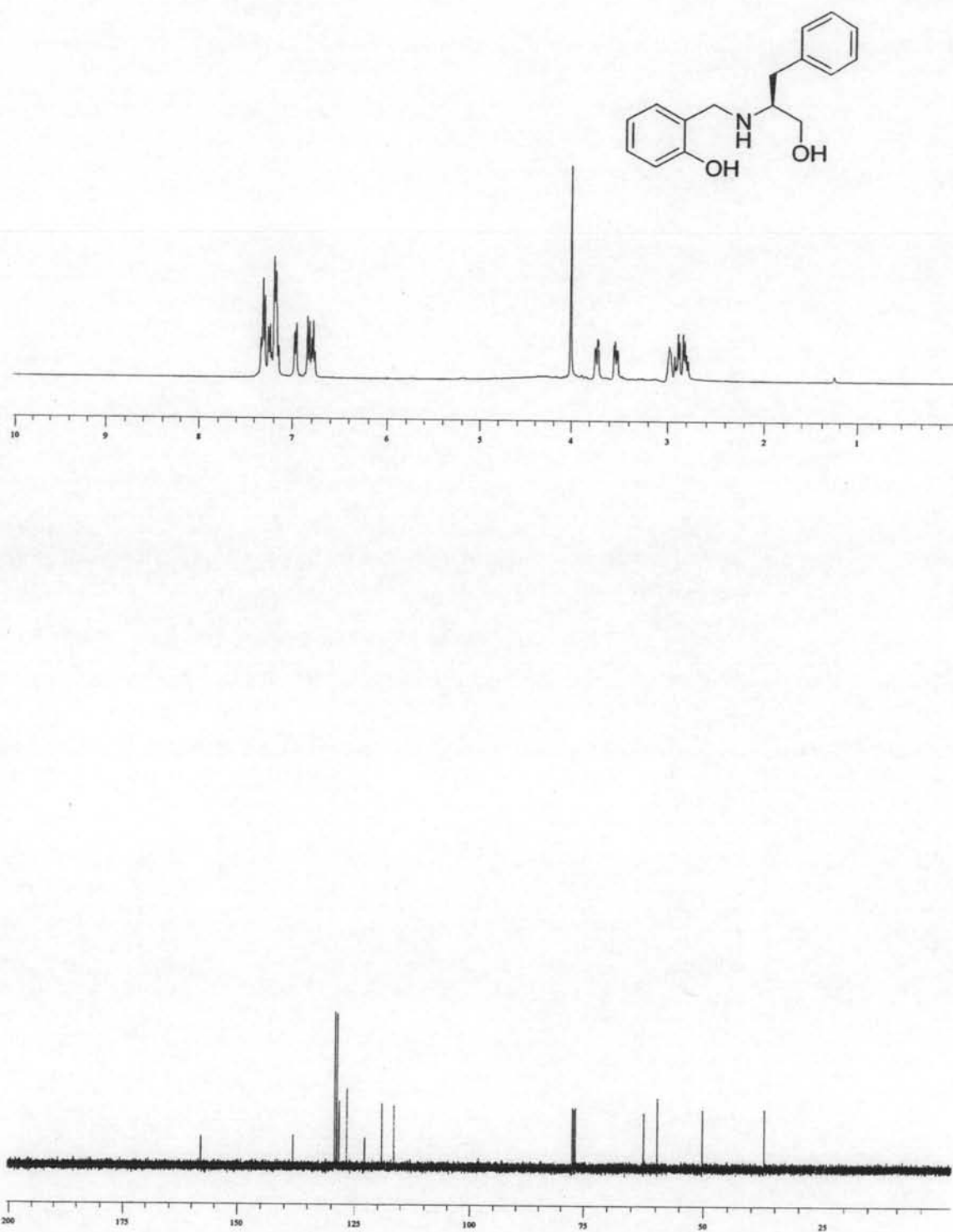


Figure A.11 ¹H NMR (CDCl₃, 400 MHz) and ¹³C NMR (CDCl₃, 100 MHz) spectra of (*S*)-2-((1-hydroxy-3-phenylpropan-2-ylamino)methyl)phenol (**26**)

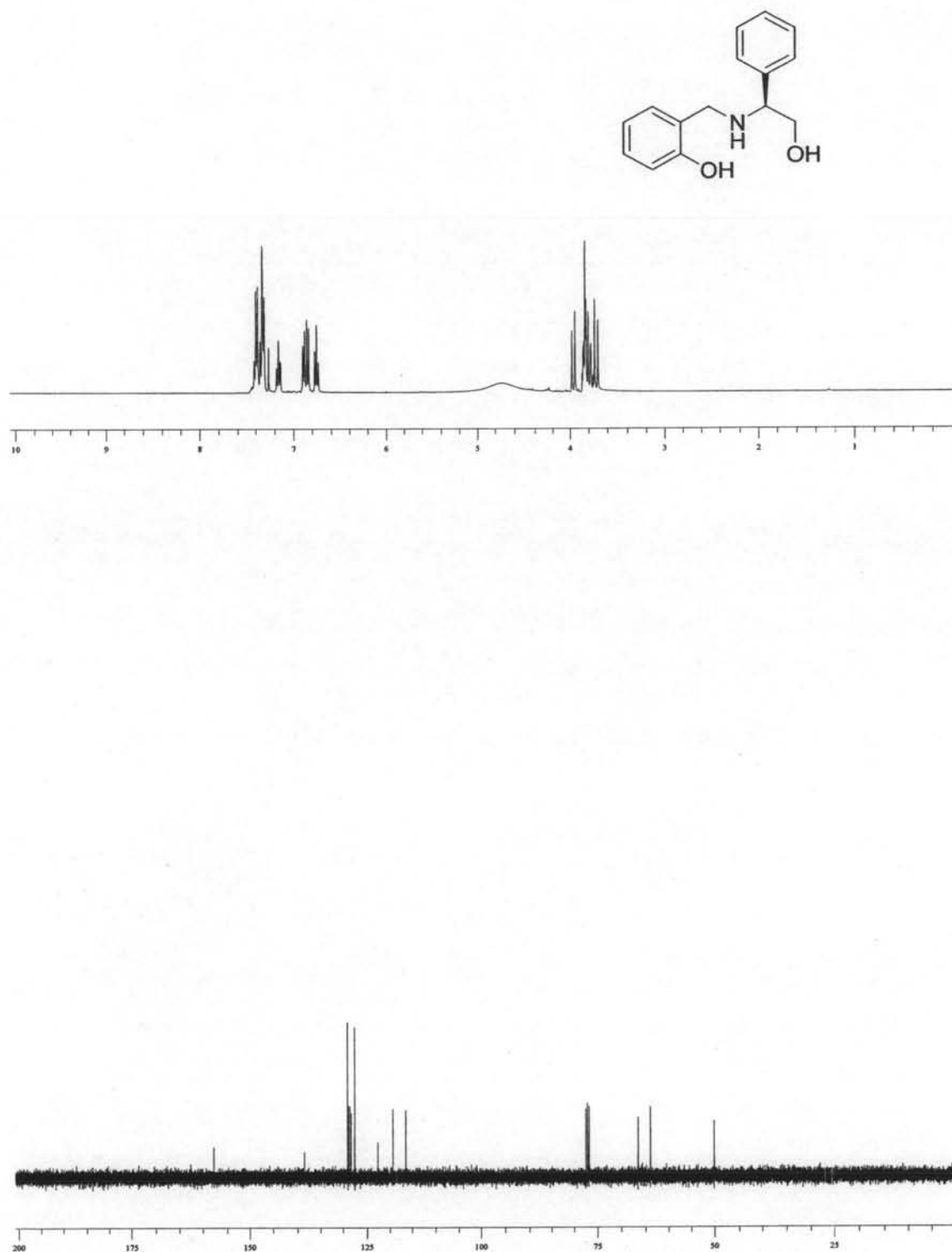


Figure A.12 ¹H NMR (CDCl₃, 400 MHz) and ¹³C NMR (CDCl₃, 100 MHz) spectra of (*S*)-2-((2-hydroxy-1-phenylethylamino)methyl)phenol (**47c**)

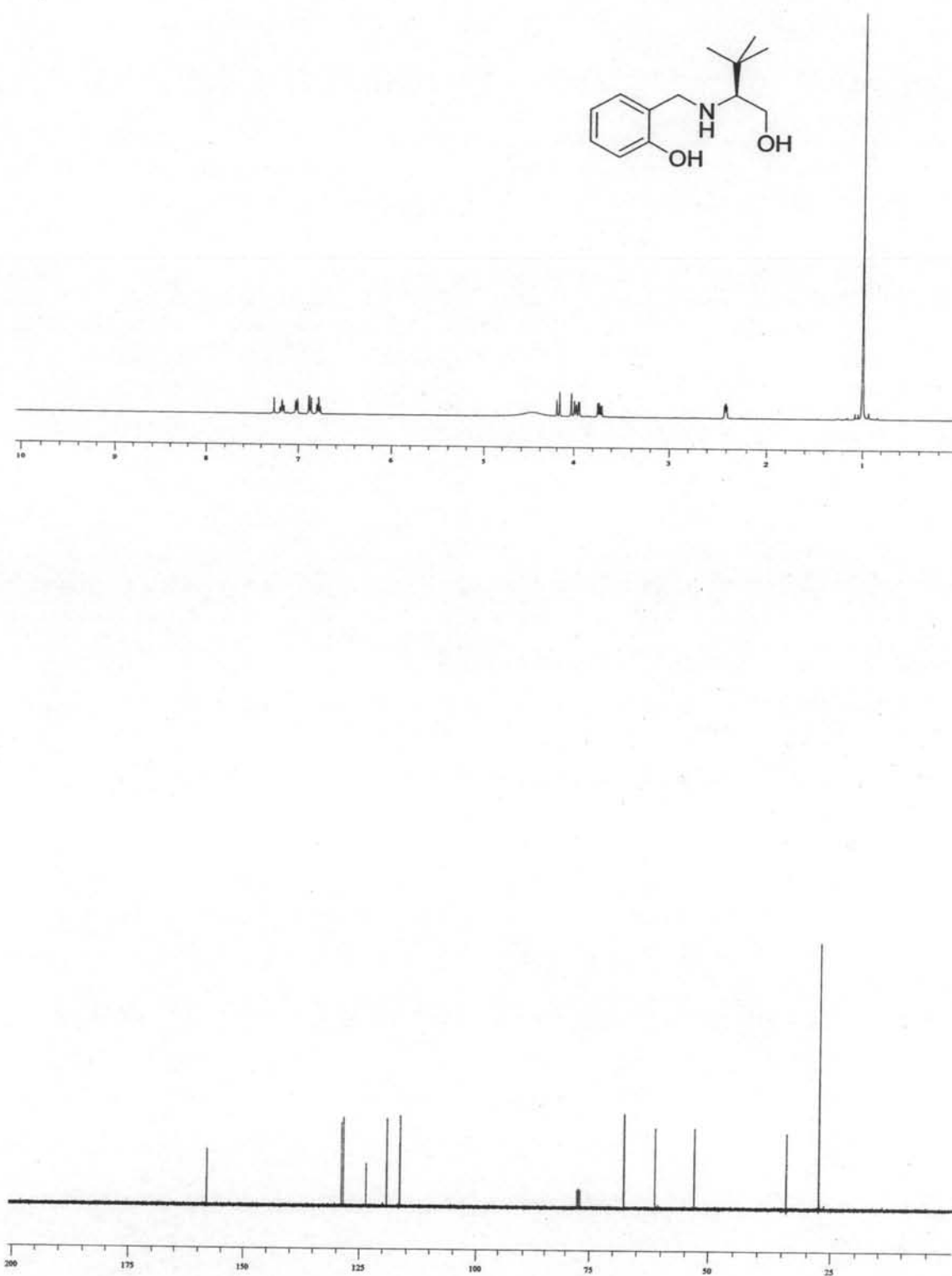


Figure A.13 ^1H NMR (CDCl_3 , 400 MHz) and ^{13}C NMR (CDCl_3 , 100 MHz) spectra of (*S*)-2-((1-hydroxy-3,3-dimethylbutan-2-ylamino)methyl)phenol (**25**)

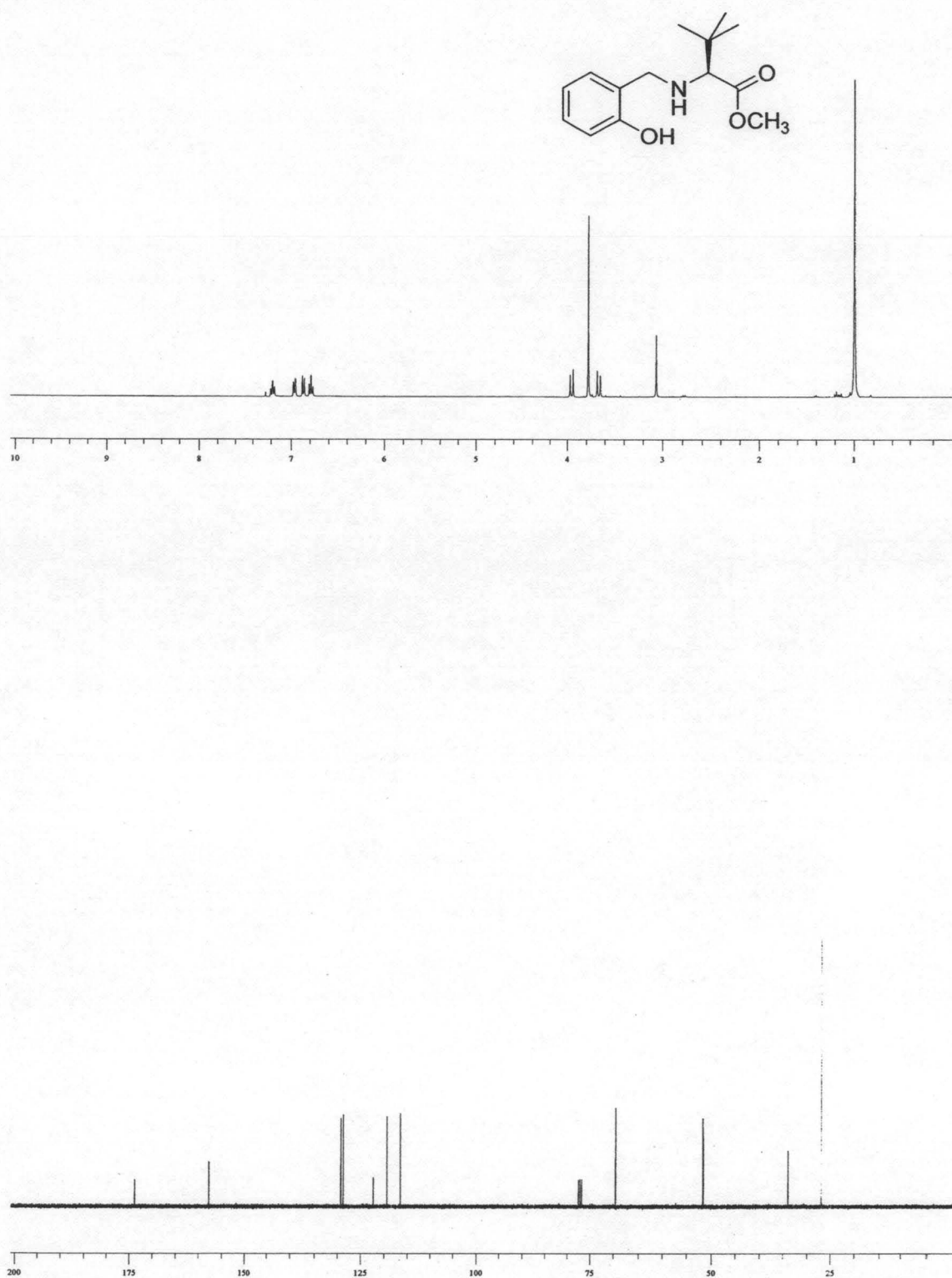


Figure A.14 ¹H NMR (CDCl₃, 400 MHz) and ¹³C NMR (CDCl₃, 100 MHz) spectra of (*S*)-methyl 2-(2-hydroxybenzylamino)-3,3-dimethylbutanoate (**48**)

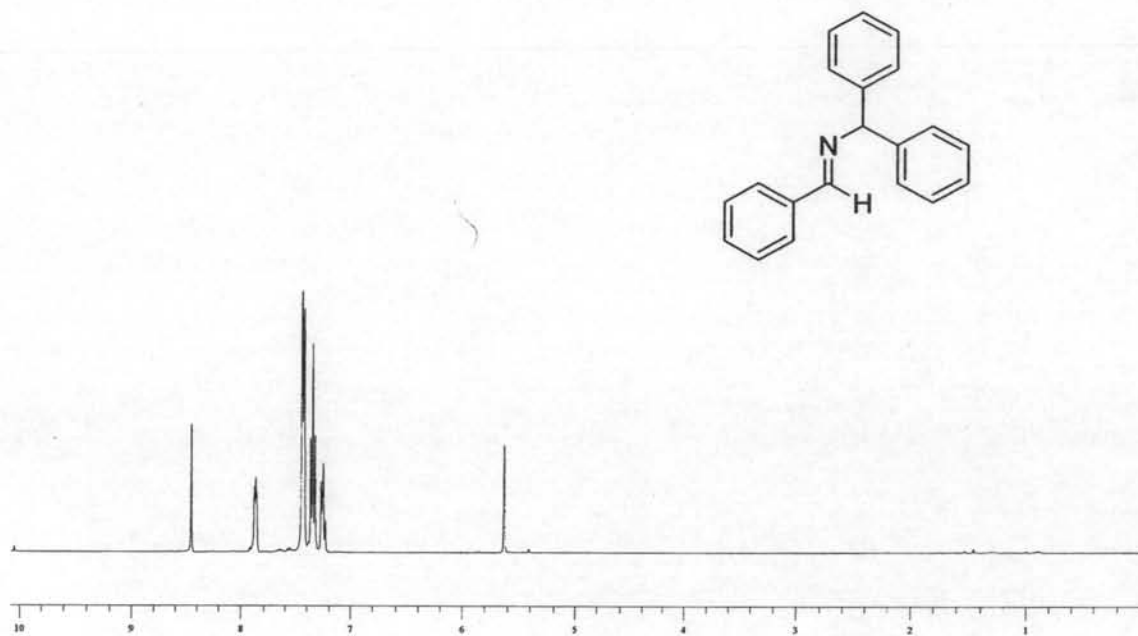


Figure A.15 ¹H NMR (CDCl₃, 400 MHz) spectrum of (*E*)-*N*-benzylidenediphenylmethanamine (**49**)

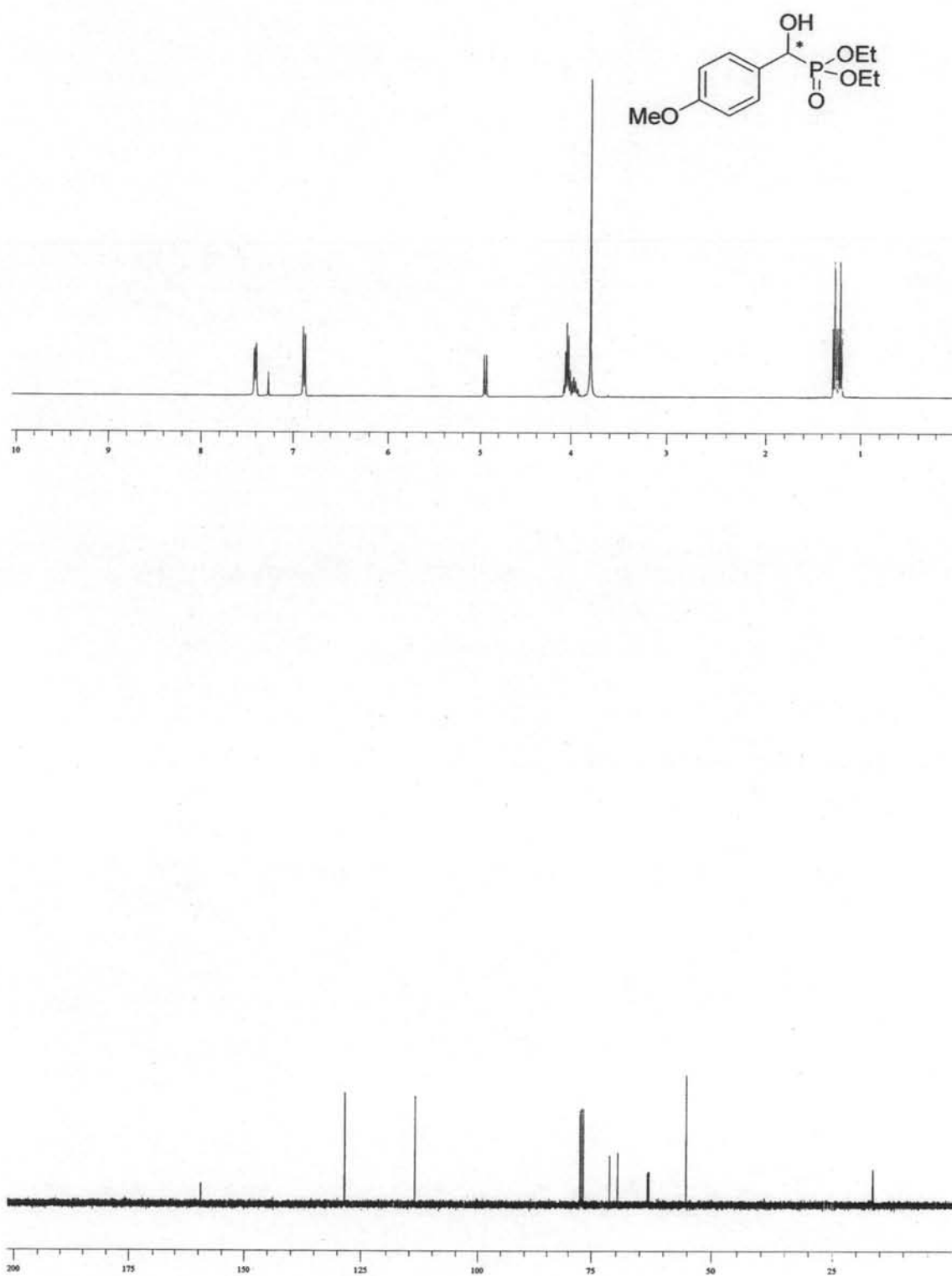


Figure A.16 ^1H NMR (CDCl_3 , 400 MHz) and ^{13}C NMR (CDCl_3 , 100 MHz) spectra of Diethyl 1-hydroxy-(4-methoxyphenyl)methylphosphonate (**50**)

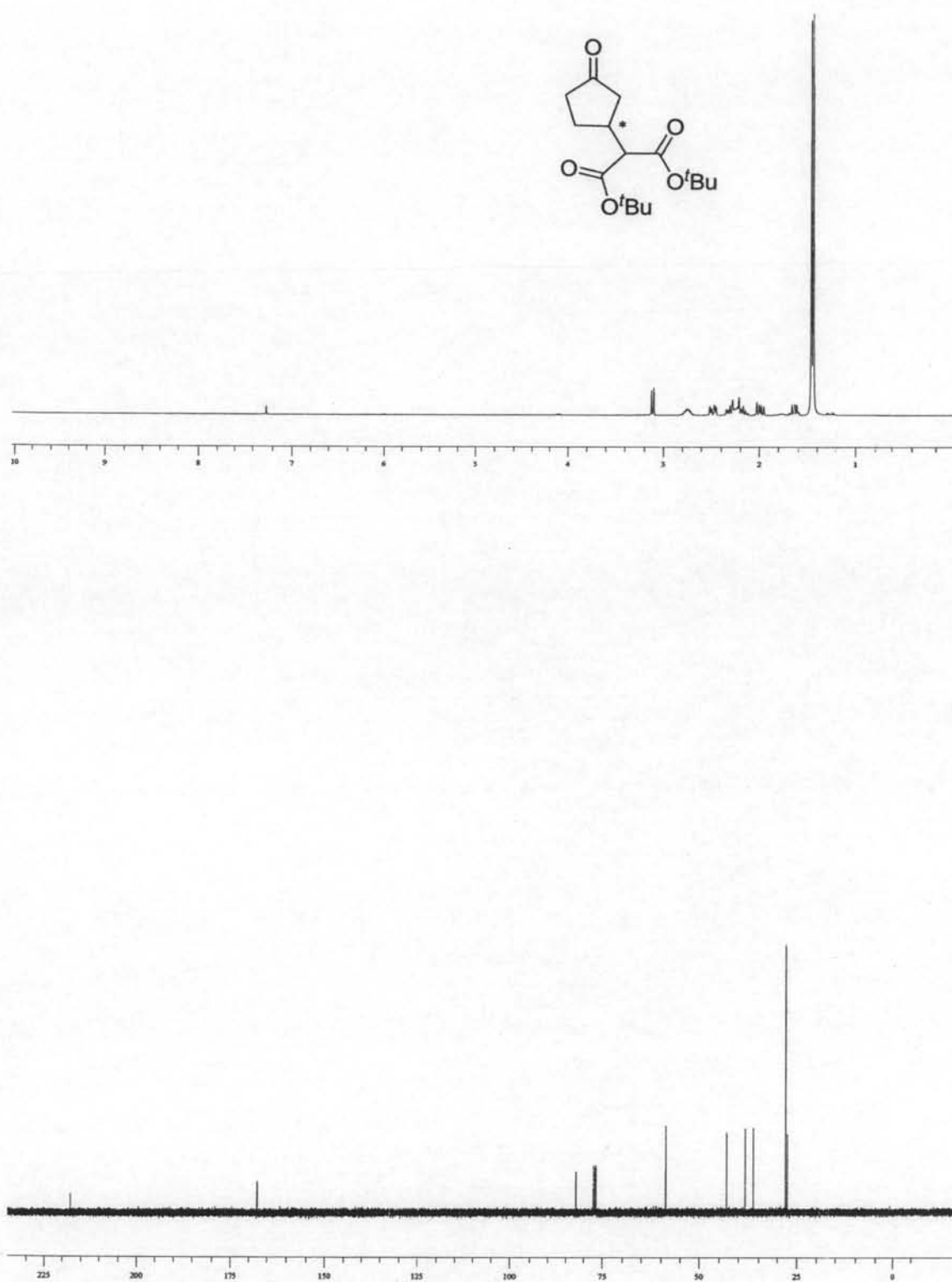


Figure A.17 ^1H NMR (CDCl_3 , 400 MHz) and ^{13}C NMR (CDCl_3 , 100 MHz) spectra of di-*tert*-butyl 2-(3-oxocyclopentyl)malonate (**51a**)

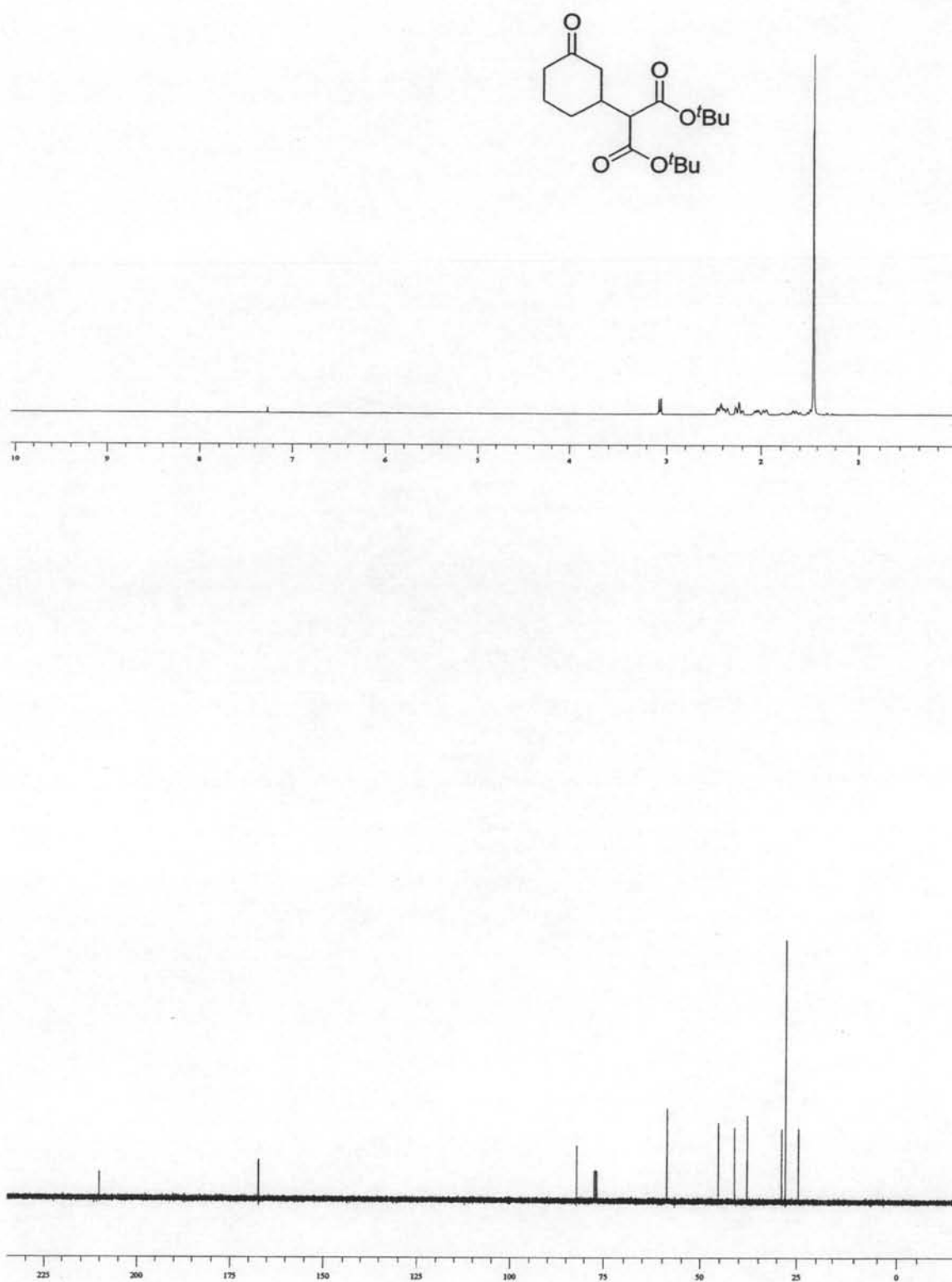


Figure A.18 ^1H NMR (CDCl_3 , 400 MHz) and ^{13}C NMR (CDCl_3 , 100 MHz) spectra of di-*tert*-butyl 2-(3-oxocyclohexyl)malonate (**51b**)

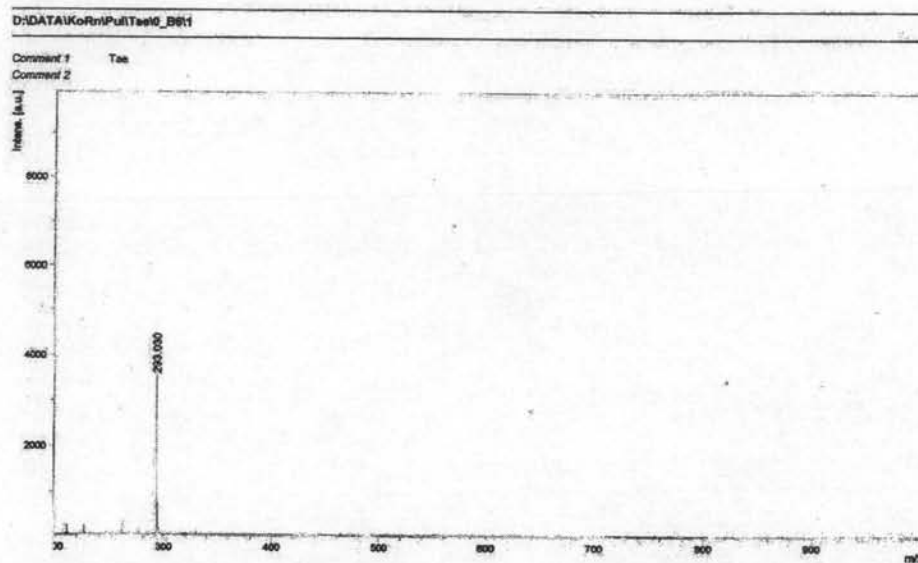


Figure A.19 MALDI-TOF spectrum of (*S*)-2-((2-hydroxy-1-(naphthalen-1-yl)ethylamino)methyl)phenol (**47a**)

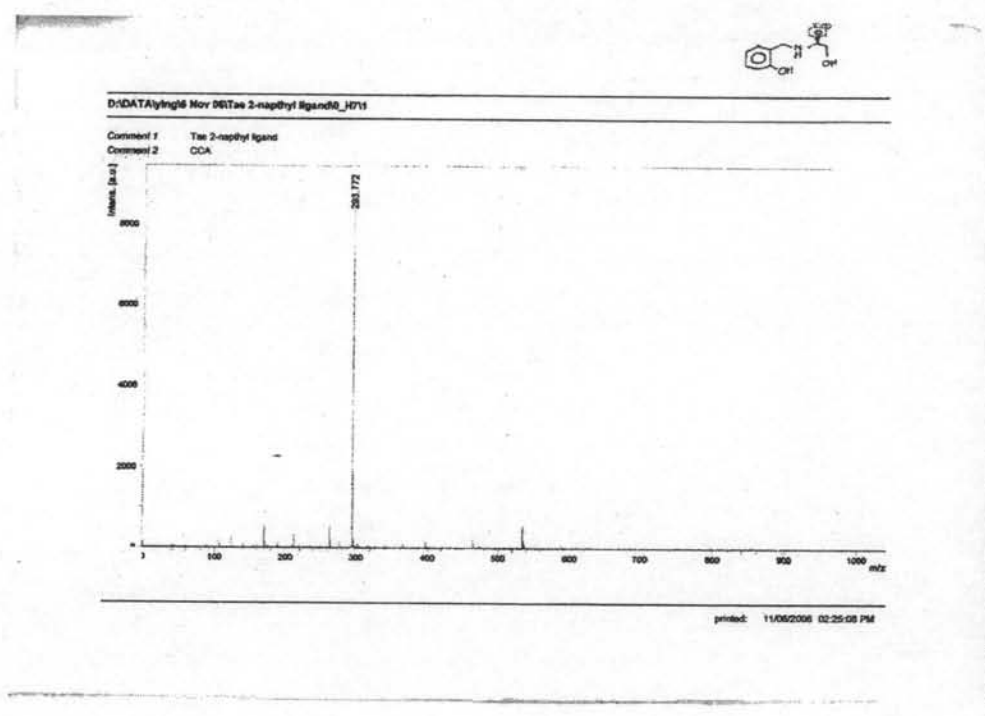
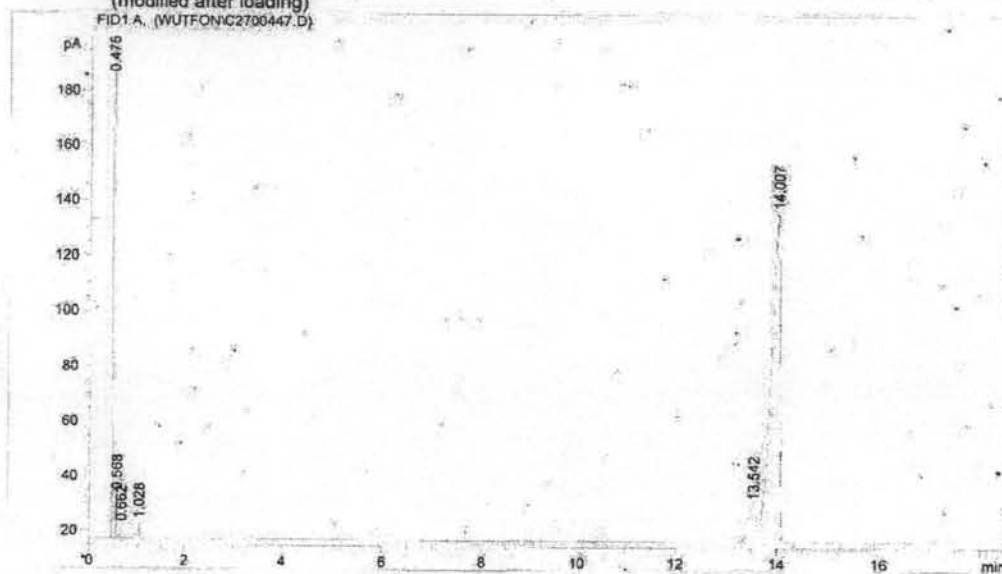


Figure A.20 MALDI-TOF spectrum of (*S*)-2-((2-hydroxy-1-(naphthalen-2-yl)ethylamino)methyl)phenol (**47b**)

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 Inj Volume : Manually
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 (modified after loading)
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 Last changed : 7/31/06 4:30:39 PM by Wut
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 FID1.A, (WUTFONIC2700447.D)



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 Area Percent Report
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Sorted By : Signal
 Multiplier : 1.0000
 Dilution : 1.0000

Signal 1: FID1.A

Peak #	RetTime [min]	Type	Width [min]	Area [pA*s]	Height [pA]	Area %
1	0.475	BB S	9.00e-3	2.65053e4	5.15343e4	94.17810
2	0.568	BV X	8.42e-3	6.92723	14.90146	0.02461
3	0.662	VV X	0.0106	2.80152	4.30797	0.00995
4	1.028	VV X	0.0120	4.28436	5.56145	0.01522
5	13.542	BV	0.1539	163.42863	13.82763	0.58069
6	14.007	VP	0.1629	1461.06348	119.26054	5.19142

Totals : 2.81438e4 5.16922e4

Results obtained with enhanced integrator!
 =====

Instrument 1 7/31/06 4:30:51 PM Wut

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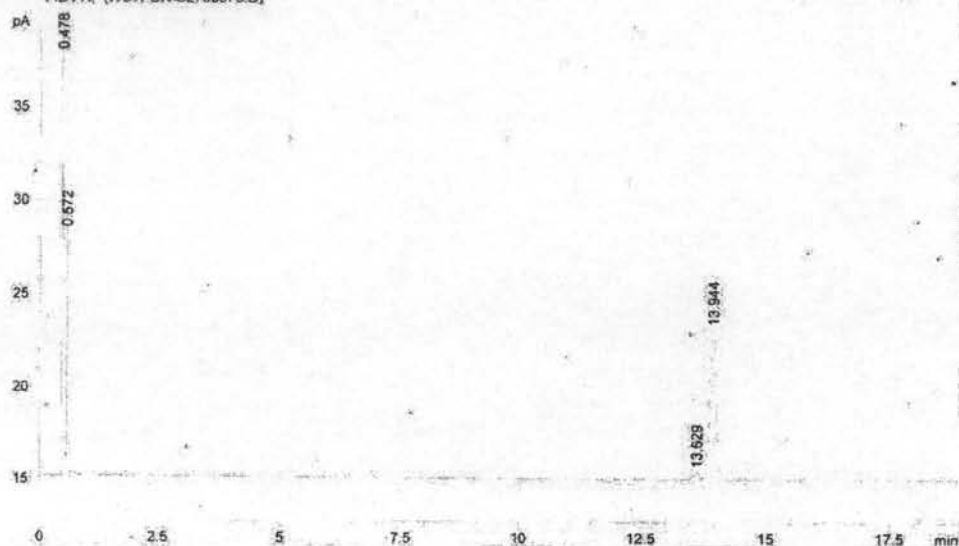
Figure A.21 GC Chromatogram of (*R*)-**51b** (80 %*ee* from using **25** as a chiral ligand)

Data File C:\HPCHEM1\DATA\WUTFONIC2700375.D

```

=====
Injection Date : 3/20/06 2:10:23 PM
Sample Name    :                               Vial : -
Acq. Operator : wut                          Inj : 1
                                           Inj Volume: Manually
Acq. Method   : C:\HPCHEM1\METHODS\ISO40.M
Last changed  : 3/20/06 1:59:54 PM by wut
                (modified after loading)
Analysis Method : C:\HPCHEM1\METHODS\ISO40.M
Last changed  : 3/20/06 4:02:38 PM by wut
                (modified after loading)
FID1 A, (WUTFONIC2700375.D)

```



```

=====
Area Percent Report
=====

```

```

Sorted By      : Signal
Multiplier    : 1.0000
Dilution      : 1.0000

```

Signal 1: FID1 A,

Peak #	RetTime [min]	Type	Width [min]	Area [pA*s]	Height [pA]	Area %
1	0.478	BB S	8.61e-3	1.34830e4	2.80263e4	99.37638
2	0.572	BV X	7.90e-3	5.93667	12.52309	0.04376
3	13.629	BV	0.1070	3.55795	4.14971e-1	0.02622
4	13.944	VB	0.1242	75.11564	8.09810	0.55364

```
Totals :                1.35676e4  2.80473e4
```

Results obtained with enhanced integrator!

```

=====
*** End of Report ***
=====

```

Instrument 1 3/20/06 4:02:51 PM wut

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Figure A.22 GC Chromatogram of (*R*)-**51b** (91 %*ee* from using **47a** as a chiral ligand)

Data File C:\HPCHEM\1\DATA\WUTFONC\2700377.D

```

=====
Injection Date : 3/20/06 2:50:30 PM
Sample Name   :                               Vial : -
Acq. Operator : wut                           Inj : 1
                                           Inj Volume : Manually
Acq. Method   : C:\HPCHEM\1\METHODS\ISO40.M
Last changed  : 3/20/06 1:59:54 PM by wut
               (modified after loading)
Analysis Method : C:\HPCHEM\1\METHODS\ISO40.M
Last changed  : 3/20/06 4:04:10 PM by wut
               (modified after loading)
FID1 A. (WUTFONC\2700377.D)

```



```

=====
Area Percent Report
=====

```

```

Sorted By      : Signal
Multiplier     : 1.0000
Dilution       : 1.0000

```

Signal 1: FID1 A.

Peak #	RetTime [min]	Type	Width [min]	Area [pA*s]	Height [pA]	Area %
1	0.479	BB S	8.92e-3	1.55020e4	3.05463e4	98.23233
2	0.573	BV X	8.30e-3	5.69778	11.44421	0.03611
3	1.039	BP	0.0125	5.04122e-1	6.15998e-1	0.00319
4	13.634	BV	0.1194	17.88722	1.94338	0.11335
5	13.981	VB	0.1268	254.86694	27.32519	1.61503

```
Totals:      1.57810e4  3.05876e4
```

Results obtained with enhanced integrator!

```

=====
*** End of Report ***
=====

```

Instrument 1 3/20/06 4:04:15 PM wut

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Figure A.23 GC Chromatogram of (R)-51b (87 %*ee* from using 47b as a chiral ligand)

Data File C:\HPCHEM\1\DATA\WUTFONC2700430.D

```

=====
Injection Date : 5/26/06 3:19:21 PM
Sample Name   :
Acq. Operator : wutfon
Vial         :
Inj Volume   : Manually
Acq. Method  : C:\HPCHEM\1\METHODS\ISO40.M
Last changed : 5/26/06 1:49:05 PM by wutfon
              (modified after loading)
Analysis Method: C:\HPCHEM\1\METHODS\ISO40.M
Last changed : 5/26/06 5:07:20 PM by wutfon
              (modified after loading)
FID1 A.: (WUTFONC2700430.D)

```



```

=====
Area Percent Report
=====

```

```

Sorted By      : Signal
Multiplier    : 1.0000
Dilution      : 1.0000

```

Signal 1: FID1 A,

Peak #	RetTime [min]	Type	Width [min]	Area [pA*s]	Height [pA]	Area %
1	0.480	BB S	8.64e-3	2.10946e4	4.06893e4	99.11435
2	0.575	BB X	8.87e-3	7.28379	14.47647	0.03422
3	13.672	BV	0.1162	140.56395	15.29068	0.66045
4	13.961	VP	0.1262	40.64538	4.50828	0.19098

```
Totals :          2.12831e4  4.07235e4
```

Results obtained with enhanced integrator

```

=====
*** End of Report ***

```

Figure A.24 GC Chromatogram of (*R*)-**51b** (55 %*ee* from using **48** as a chiral ligand)

Data File C:\HPCHEM\1\DATA\WUTFONC2700396.D

IS 231

=====
 Injection Date : 4/27/06 11:23:04 AM
 Sample Name : Vial: -
 Acq. Operator : Wut Inj: 1
 Inj Volume : Manually
 Acq. Method : C:\HPCHEM\1\METHODS\ISO40.M
 Last changed : 4/27/06 11:16:57 AM by Wut
 (modified after loading)
 Analysis Method : C:\HPCHEM\1\METHODS\ISO40.M
 Last changed : 4/28/06 12:41:49 PM by Wut
 (modified after loading)
 FID1 A, (WUTFONC2700396.D)



=====
 Area Percent Report
 =====

Sorted By : Signal
 Multiplier : 1.0000
 Dilution : 1.0000

Signal 1: FID1 A,

Peak #	RetTime [min]	Type	Width [min]	Area [pA*s]	Height [pA]	Area %
1	0.481	BB S	8.99e-3	2.31039e4	4.50369e4	99.14151
2	0.667	BB T	7.96e-3	5.52272	10.81826	0.02370
3	1.789	BP	0.0216	6.84693e-1	5.11720e-1	0.00294
4	27.054	BV	0.2100	33.20890	1.87181	0.14250
5	27.728	VB	0.2239	160.64743	8.64063	0.68936

Totals : 2.33040e4 4.50587e4

Results obtained with enhanced integrator!

=====
 *** End of Report ***

Instrument 1 4/28/06 12:41:55 PM Wut

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Figure A.25 GC Chromatogram of (*S*)-**51a** (66 %*ee* from using **47a** as a chiral ligand)

Data File C:\HPCHEM\1\DATA\WUTFONIC2700477.D

=====
 Injection Date : 8/31/06 10:10:29 AM
 Sample Name : Vial : -
 Acq. Operator : WUT Inj : 1
 Inj Volume : Manually
 Acq. Method : C:\HPCHEM\1\METHODS\ISO40.M
 Last changed : 8/31/06 10:04:18 AM by WUT
 (modified after loading)
 Analysis Method : C:\HPCHEM\1\METHODS\ISO40.M
 Last changed : 9/1/06 5:23:44 PM by WUT
 (modified after loading)



=====
 Area Percent Report
 =====

Sorted By : Signal
 Multiplier : 1.0000
 Dilution : 1.0000

Signal 1: FID1 A.

Peak #	RetTime [min]	Type	Width [min]	Area [pA*s]	Height [pA]	Area %
1	0.476	BB S	8.32e-3	2.35528e4	4.71643e4	98.97970
2	26.715	BV	0.2103	62.62929	3.55832	0.26320
3	27.379	VB	0.2239	180.15704	9.82674	0.75710

Totals : 2.37956e4 4.71776e4

Results obtained with enhanced integrator!
 =====

Instrument 1 9/1/06 5:23:52 PM WUT

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Figure A.26 GC Chromatogram of (*S*)-**51a** (48 %*ee* from using **25** as a chiral ligand)

Data File C:\HPCHEM1\DATA\WUTFONIC2700379.D

```

=====
Injection Date : 3/20/06 3:25:07 PM
Sample Name   :                               Vial : -
Acq. Operator : wut                          Inj : 1
                                           Inj Volume : Manually
Acq. Method  : C:\HPCHEM1\METHODS\ISO40.M
Last changed : 3/20/06 3:07:30 PM by wut
              (modified after loading)
Analysis Method : C:\HPCHEM1\METHODS\ISO40.M
Last changed  : 3/20/06 4:04:56 PM by wut
              (modified after loading)
FID1 A, (WUTFONIC2700379.D)

```



```

=====
Area Percent Report
=====

```

```

Sorted By      : Signal
Multiplier     : 1.0000
Dilution       : 1.0000

```

Signal 1: FID1 A,

Peak #	RetTime [min]	Type	Width [min]	Area [pA*s]	Height [pA]	Area %
1	0.477	PB S	8.80e-3	2.43557e4	4.89517e4	98.83817
2	0.538	BV X	7.72e-3	7.10523	15.34604	0.02883
3	0.572	VB X	8.65e-3	74.16885	142.84396	0.30099
4	0.982	BB	0.0114	139.76581	192.97353	0.56719
5	10.194	BP	0.1075	32.76448	4.68880	0.13296
6	10.572	BB	0.0950	32.49281	4.68038	0.13186

```
Totals:                2.46420e4  4.93123e4
```

Results obtained with enhanced integrator!

Instrument 1 3/20/06 4:05:21 PM wut

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Figure A.27 GC Chromatogram of 50 (0 %ee from using 47a as a chiral ligand)

Data File C:\HPCHEM\1\DATA\WUTFONC2700380.D

```

=====
Injection Date : 3/20/06 3:39:30 PM
Sample Name    :                               Vial : -
Acq. Operator : wut                          Inj : 1
                                           Inj Volume : Manually
Acq. Method   : C:\HPCHEM\1\METHODS\ISO40.M
Last changed  : 3/20/06 3:07:30 PM by wut
               (modified after loading)
Analysis Method : C:\HPCHEM\1\METHODS\ISO40.M
Last changed  : 3/20/06 4:04:56 PM by wut
               (modified after loading)
FID1 A, (WUTFONC2700380.D)

```



```

=====
Area Percent Report
=====

```

```

Sorted By      : Signal
Multiplier     : 1.0000
Dilution       : 1.0000

```

Signal 1: FID1 A,

Peak #	RetTime [min]	Type	Width [min]	Area [pA*s]	Height [pA]	Area %
1	0.477	PB S	8.47e-3	2.80713e4	5.52309e4	98.33550
2	0.538	BV X	8.46e-3	7.91159	16.90996	0.02771
3	0.572	VB X	8.83e-3	130.31720	260.85913	0.45651
4	0.982	BB	0.0115	243.27515	333.90021	0.85221
5	10.206	BB	0.0922	46.17417	6.25184	0.16175
6	10.589	BB	0.1051	47.47813	6.74736	0.16632

```
Totals:                2.85465e4  5.58555e4
```

Results obtained with enhanced integrator!

Instrument 1 3/20/06 4:05:47 PM wut

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Figure A.28 GC Chromatogram of 50 (0 %ee from using 47b as a chiral ligand)

VITA

Mr. Ittiphol Saengswang was born on June 6th, 1981 in Bangkok, Thailand. He was received a Bachelor's Degree of Science, majoring in chemistry from Chulalongkorn University in 2003. Since 2003, he has been a graduate student studying Organic chemistry as his major course at Chulalongkorn University. During his studies, he was promoted by a research grant and the teaching assistant fellowship from the graduate school, Chulalongkorn University.

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