

Supporting Information

Functional chromatographic technique for natural product isolation

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Table of Contents

A. Recombinant protein expression and purification	S3
B. Protein-resin preparation	S3-S4
C. Functional chromatography	S4
D. HPLC-MS analyses	S4
E. ATPase activity assays	S4
F. His ₆ -p97 stability tests	S4-S5
G. MDH assays	S5
H. Ub ^{G76V} GFP and TCR α -GFP degradation assays	S5
Figure S1. Resin loading efficiency	S6
Figure S2. His ₆ -p97 resin stability	S6
Figure S3. UbG76VGFP assay	S7
Figure S4. TCR α -GFP assay	S7
Figure S5. ¹ H NMR spectrum of oxaspirol C (1) in DMSO- <i>d</i> ₆	S8
Figure S6. ¹³ C NMR spectrum of oxaspirol C (1) in DMSO- <i>d</i> ₆	S9
Figure S7. ¹ H- ¹ H COSY NMR spectrum of oxaspirol C (1) in DMSO- <i>d</i> ₆	S10
Figure S8. ¹ H- ¹ H ROESY NMR spectrum of oxaspirol C (1) in DMSO- <i>d</i> ₆	S11
Figure S9. ¹ H- ¹³ C HSQC NMR spectrum of oxaspirol C (1) in DMSO- <i>d</i> ₆	S12
Figure S10. ¹ H- ¹³ C HMBC NMR spectrum of oxaspirol C (1) in DMSO- <i>d</i> ₆	S13
Figure S11. ¹ H NMR spectrum of oxaspirol B (2) in CDCl ₃	S14
Figure S12. ¹³ C NMR spectrum of oxaspirol B (2) in CDCl ₃	S15
Figure S13. ¹ H- ¹ H COSY NMR spectrum of oxaspirol B (2) in CDCl ₃	S16
Figure S14. ¹ H- ¹ H ROESY NMR spectrum of oxaspirol B (2) in CDCl ₃	S17
Figure S15. ¹ H- ¹³ C HSQC NMR spectrum of oxaspirol B (2) in CDCl ₃	S18
Figure S17. ¹ H- ¹³ C HMBC NMR spectrum of oxaspirol B (2) in CDCl ₃	S19

A. Recombinant protein expression and purification. All proteins and used in this study were prepared by conventional recombinant protein expression followed by immobilized metal affinity chromatography (IMAC) or native purification in the case of p97.

p97. *E. coli* BL21(DE3) cells containing pET14b-p97 were grown in Luria Broth (LB) medium containing 100 µg/mL ampicillin at 37°C to an OD₆₀₀ of 0.8, followed by induction with 0.5 mM isopropyl β-D-1-thiogalactopyranoside (IPTG) for 4 h at 37°C. Cells were collected by centrifugation (3000 × g for 10 min), resuspended in 40 mL lysis buffer (50 mM HEPES pH 7.4, 150 mM KCl, 5 mM MgCl₂, 5% glycerol, 2 mM β-mercaptoethanol (BME), one complete EDTA-free protease inhibitor cocktail (Roche) per 50 mL of buffer), and lysed by single passage through an M110-T microfluidizer (Microfluidics). The lysate was clarified by centrifugation (118,834 × g, 1 h, 4°C) and the resulting supernatant was incubated for 1 h at 4°C with Talon metal affinity resin (Clontech) equilibrated in 50 mM HEPES, pH 7.4, 150 mM KCl, 5 mM MgCl₂, 5% glycerol. The resin and supernatant were then loaded into a 25 mL disposable column (BioRad), washed with 10 column volumes of wash buffer (50 mM HEPES, pH 7.4, 150 mM KCl, 5 mM MgCl₂, 5% glycerol, 2 mM BME, 5 mM imidazole), followed by 10 column volumes of stringent wash buffer (50 mM HEPES, pH 7.4, 1 M KCl, 5 mM MgCl₂, 5% glycerol, 2 mM BME, 20 mM imidazole), and eluted with elution buffer (50 mM HEPES, pH 7.4, 1 M KCl, 5 mM MgCl₂, 5% glycerol, 2 mM BME, 250 mM imidazole). Fractions were analyzed by 12% SDS PAGE and those containing p97 were pooled and dialyzed into storage buffer (20 mM HEPES, 150 mM KCl, 5 mM MgCl₂, 5% glycerol, 2 mM BME, pH 7.4), concentrated using a 30 kDa Ultra-15 centrifugal filter (Amicon) to yield a solution of p97 at 12 mg/mL, which was aliquoted, frozen in liquid N₂, and stored at -80°C until needed.

HSPA1A and HSC70. *E. coli* BL21(DE3) cells containing pSpeedET-HSPA1A or pSpeedET-HSC70 were grown in Luria Broth (LB) medium containing 50 µg/mL kanamycin at 37°C to an OD₆₀₀ of 0.5, cooled in 16°C for 1 h, followed by induction with 0.2 mM isopropyl β-D-1-thiogalactopyranoside (IPTG) for 48 h at 16°C. Cells were collected by centrifugation (3000 × g for 10 min), resuspended in 40 mL lysis buffer (50 mM HEPES, pH 7.4, 150 mM KCl, 5 mM MgCl₂, 5% glycerol, 2 mM BME, one complete EDTA-free protease inhibitor cocktail (Roche) per 50 mL of buffer), and lysed by single passage through an M110-T microfluidizer (Microfluidics). The lysate was clarified by centrifugation (118,834 × g, 1 h, 4°C) and the resulting supernatant was incubated for 1 h at 4°C with Talon metal affinity resin (Clontech) equilibrated in 50 mM HEPES pH 7.4, 150 mM KCl, 5 mM MgCl₂, and 5% glycerol. The resin and supernatant were then loaded into a 25 mL disposable column (BioRad), washed with 10 column volumes of wash buffer (50 mM HEPES pH 7.4, 150 mM KCl, 5 mM MgCl₂, 5% glycerol, 2 mM BME, 5 mM imidazole), followed by 10 column volumes of stringent wash buffer (50 mM HEPES, pH 7.4, 1 M KCl, 5 mM MgCl₂, 5% glycerol, 2 mM BME, 20 mM imidazole), and eluted with elution buffer (50 mM HEPES pH 7.4, 1 M KCl, 5 mM MgCl₂, 5% glycerol, 2 mM BME, 250 mM imidazole). Fractions were analyzed by 12% SDS PAGE and those containing HSPA1A or HSC70 were pooled and dialyzed into storage buffer (20 mM HEPES pH 7.4, 150 mM KCl, 5 mM MgCl₂, 5% glycerol, 2 mM BME), concentrated using a 30 kDa Ultra-15 centrifugal filter (Amicon) to yield a solution of HSPA1A at 2.0 mg/mL or HSC70 at 9.8 mg/mL, which was aliquoted, frozen in liquid N₂, and stored at -80°C until needed.

B. Protein resin preparation. An aliquot of a slurry of Affigel-15 resin (Bio-Rad) was transferred to a microcentrifuge tube, collected by centrifugation on a minicentrifuge at 2000×g, and the aqueous phase was removed. Next, the resin was washed three times with an equal volume of buffer (20 mM HEPES pH 7.4, 150 mM KCl, 1 mM MgCl₂, 2 mM BME). The protein was added in an equal volume of buffer (comparable volumes of resin and aqueous

phase are used) and the resin was incubated by inverted shaking at 4°C. Aliquots were taken every 10-15 min for 5 h to monitor coupling. After protein has finished coupling, the resin bound protein is washed three times with two volumes of buffer (twice the volume of resin). The activity of each batch of resin was checked using either the ATPase activity assay (p97 and HSC70) as described in Section G or a L-malate dehydrogenase assay (MDH) as described in Section H. HSPA1A could not be evaluated by the ATPase assay as described in the manuscript text.

C. Functional chromatography. The following provides a general procedure for the functional chromatographic purification: a 200 µL aliquot of solution of *Parmotrema tinctorium* extract (2 mg/mL) in buffer (20 mM HEPES pH 7.4, 150 mM KCl, 1 mM MgCl₂, 2 mM BME) containing 2% DMSO was added to 200 µL of resin bearing 5 mg/mL of protein in a 2 mL Eppendorf tube. The tube shaken on an inversion rotor for 12 h at 4°C. After, the supernatant was discarded and the resin washed three times with 500 µL of buffer. After the second wash, the resin was suspended in 500 µL of buffer and transferred to a ½ dram glass vial. After removal of the supernatant, 200 µL of 95% EtOH was added and the mixture was incubated for at rt for 10 min. The EtOH was collected and dried by airflow.

D. HPLC-MS analyses. The following outlines the general conditions and methods used for the LC-MS analyses included in Figure 4.

Column Dimensions: Symmetry C18 4.6 × 75.0 mm Column. Particle Size = 3.5 µM

Solvent A: 25:925 CH₃CN:H₂O containing 0.1% formic acid

Solvent B: 95:5 CH₃CN:H₂O containing 0.1% formic acid

Flow rate: 0.2 mL/min

Injection Volume: 35 µL

Gradient: 5% - 95% Line B

Method: Sample loading was conducted with 95% solvent A and 5% solvent B running. Over a 36 min period, a gradient was run from 95% solvent A and 5% solvent B to 5% solvent A and 95% solvent B. The mixture of 5% solvent A and 95% solvent B was then held for an additional 12 min. Total run time is 60 min. Plots were cropped to display between 0 – 34 min for presentation in Figure 4.

0-12 min = 95% A and 5% B

12-48 min = gradient to 95% B

48-60 min = hold 95% B

E. ATPase activity assays. ATPase activity was evaluated using Malachite green as an indicator for each protein. Assay buffer (50 mM HEPES pH 7.4, 150 mM KCl, 10 mM MgCl₂, 2 mM BME) was prepared in 100 µL reaction volume and protein was added to 1 µM followed by incubation at 37°C for 10 min, ATP hydrolysis was initiated by adding 2 mM ATP to each microcentrifuge tube, followed by incubation at 37°C. Aliquots (20 µL) were taken at 2 min, 4 min, 6 min, 8 min, and 10 min and added immediately into 800 µL of malachite green solution (9.3 µM malachite green, 53 mM (NH₄)₂MoO₄, 1M HCl, 10% Tween 20). After 1 min, 10 µL of 34% sodium citrate was added and the OD₆₆₀ was read on a Genesys 10S Vis Spectrophotometer (Thermo Scientific).

F. His₆-p97 stability test. His₆-p97 was coupled onto Affigel-15 resin and stored at 4°C. Three preparations were tested, uncoupled H6p97, ethanolamine capped H6p97, and uncapped H6p97. These were assayed over the course of 31 days using the ATPase activity assay.

G. MDH assay. L-Malate dehydrogenase (Roche) purchased as a 3.2 M ammonium sulfate suspension was dialyzed into (20 mM HEPES pH 7.4, 150 mM KCl, 1 mM MgCl₂, 2 mM BME) using a Slide-A-Lyzer Dialysis Cassette (Thermo Scientific). Uncoupled Malate Dehydrogenase, Affigel-10-MDH, Affigel-15-MDH were prepared and assayed in Methacrylate cuvettes. The MDH Assay buffer contains (50 mM HEPES pH 7.4, 2 mM ketomalonnate, 180 μM NADH). Cuvette was blanked with assay solution without NADH. NADH was then added and a baseline of the assay buffer was taken. A 1 μL aliquot of each MDH preparation was added and read at 340 nM at 1 sec intervals for 60 sec using a Biomate3 Spectrophotometer (Thermo Scientific).

H. Ub^{G76V}GFP and TCRα-GFP assays. HEK293 cells stably expressing Ub^{G76V}-GFP or TCRα-GFP were kindly provided by Dr. Ron Kopito (Stanford University). For live cell imaging, HEK293 cells stably expressing Ub^{G76V}GFP and TCRα-GFP were seeded at 2 × 10⁶ cells in phenol-red free Dulbecco's Modified Eagle Medium (DMEM) supplemented with 10% fetal bovine serum (FBS) on 35 mm glass-bottom dishes (Bio Express). After incubation at 37°C under a 5% CO₂ atmosphere for 18 h, each dish was treated with either 10 μM, 20 μM, or 50 μM of **1** or **2** from DMSO stock solutions (0.1% final DMSO concentration). DMSO (0.1%) was used as a negative control. The cells were imaged in phenol red free DMEM supplemented with 10% FBS. The fluorescence intensity was determined by flow cytometry on a FACScan flow cytometer (BD Biosciences) equipped with an air-cooled 15 mW Argon ion laser tuned to 488 nm. The emission fluorescence of GFP was detected and recorded through a 530/30bandpass filter. List mode data files consisting of 10,000 events gated on FSC (forward scatter) versus SSC (side scatter) were acquired and analyzed using CellQuest PRO software (BD Biosciences) at a rate of 200-400 events per second. Data from these studies are provided in Figures S3 and S4.

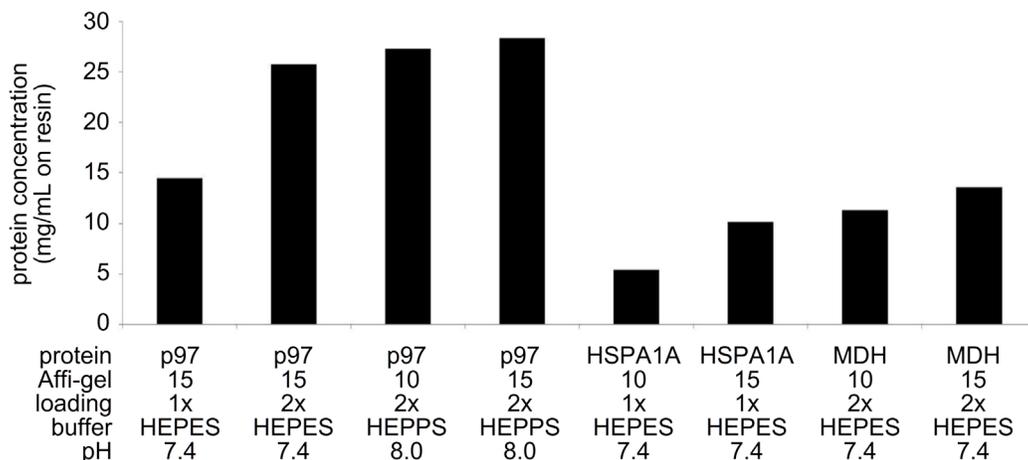


Figure S1. Resin loading efficiency. Affi-gel resins allows up to 25 mg/mL protein loading. In this study, we evaluated to loading of His₆-p97, His₆-HSPA1A, and MDH (malate dehydrogenase) on to either Affi-gel 10 or Affi-gel15 resin.

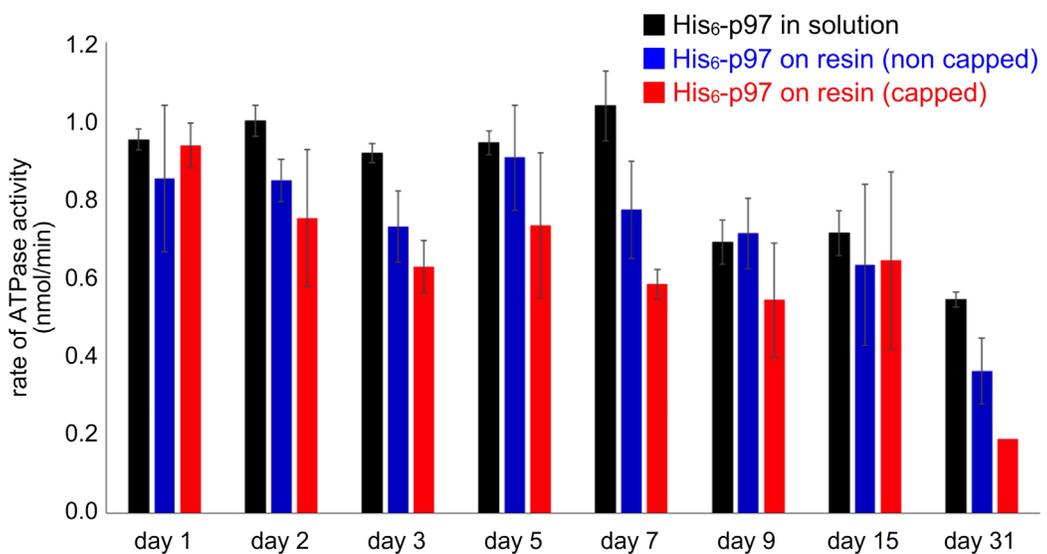


Figure S2. His₆-p97 resin stability. The ATPase activity of free His₆-p97 and Affi-gel 15 resin bearing 11.4 mg/mL of His₆-p97 was measured over the course of a month at 4 °C. APTase activity was monitored by the Malachite green assay as described in the experimental procedures.

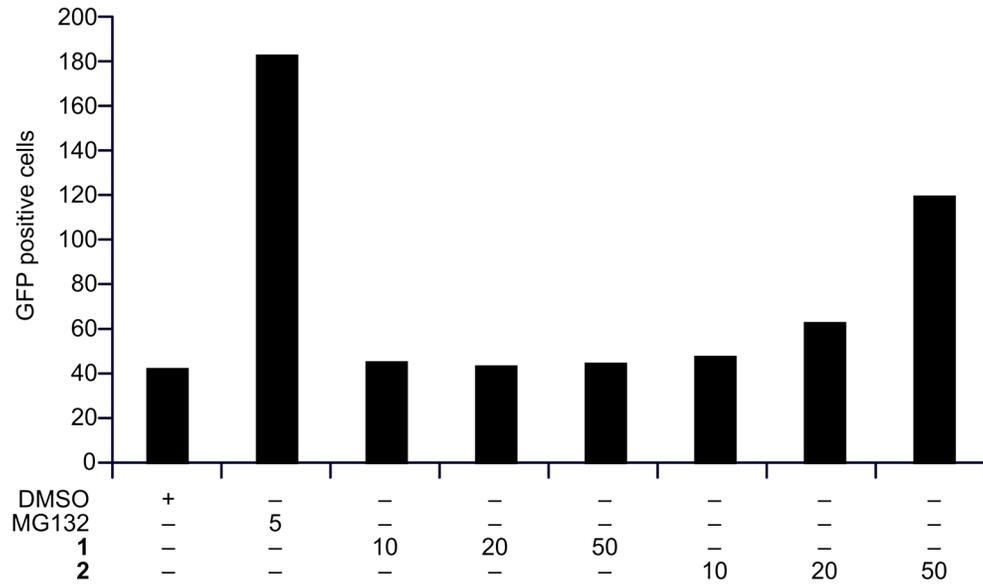


Figure S3. Ub^{G76V}GFP assay. Oxaspirol B (2) prevents degradation of Ub^{G76V}GFP in HEK-293 cells as compared to MG132, a proteasome inhibitor, while oxaspirol C (1) does not. Concentrations are provided in μ M.

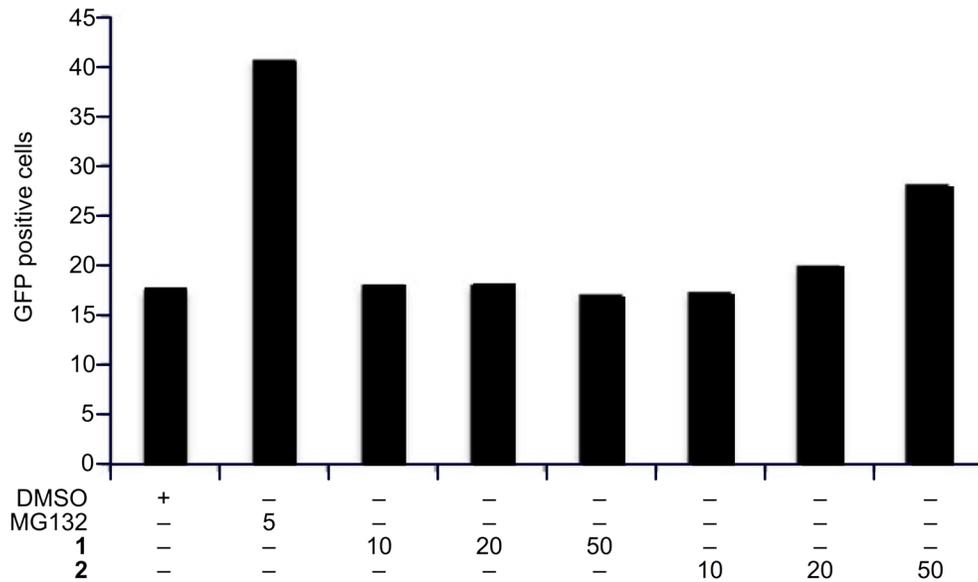


Figure S4. TCR α -GFP assay. Oxaspirol B (2) prevents degradation of TCR α -GFP in HEK-293 cells as compared to MG132, a proteasome inhibitor, while oxaspirol C (1) does not. Concentrations are provided in μ M.

Figure S5. ^1H NMR spectrum of oxaspirol C (**1**) in $\text{DMSO-}d_6$

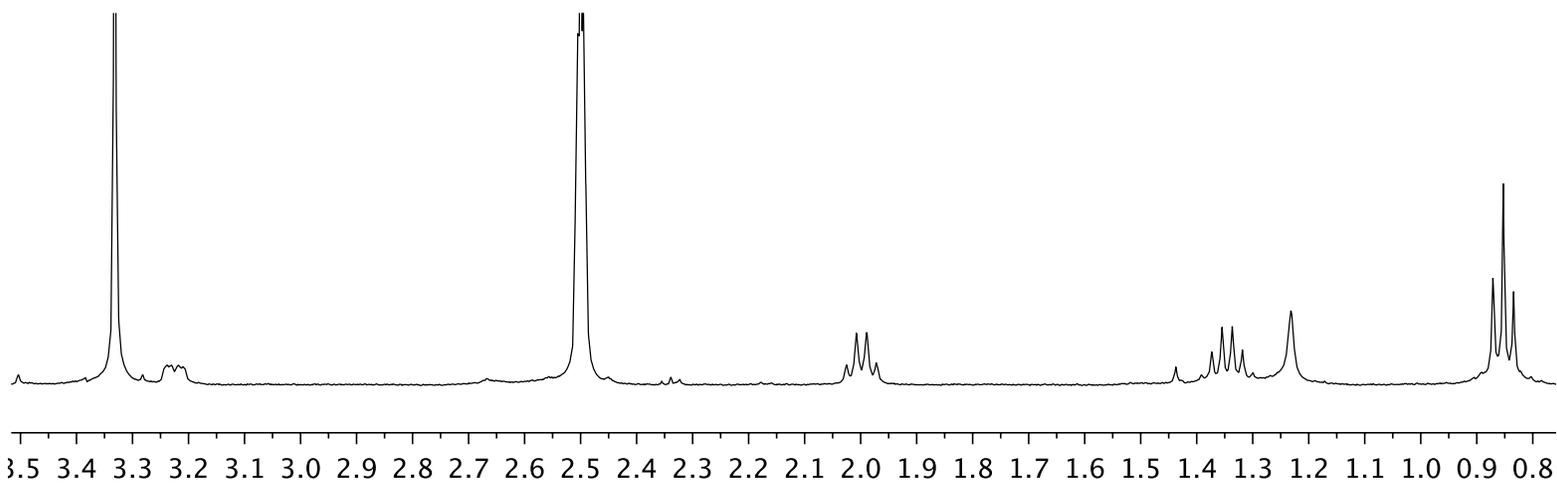
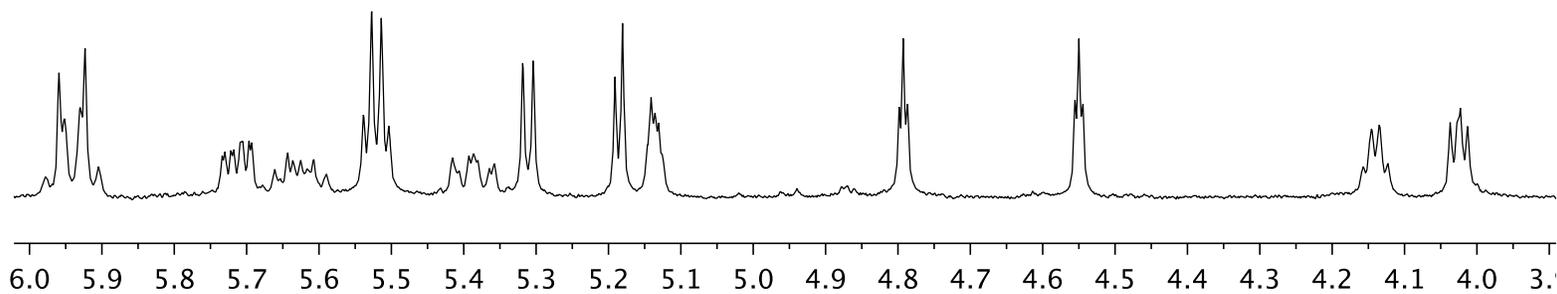
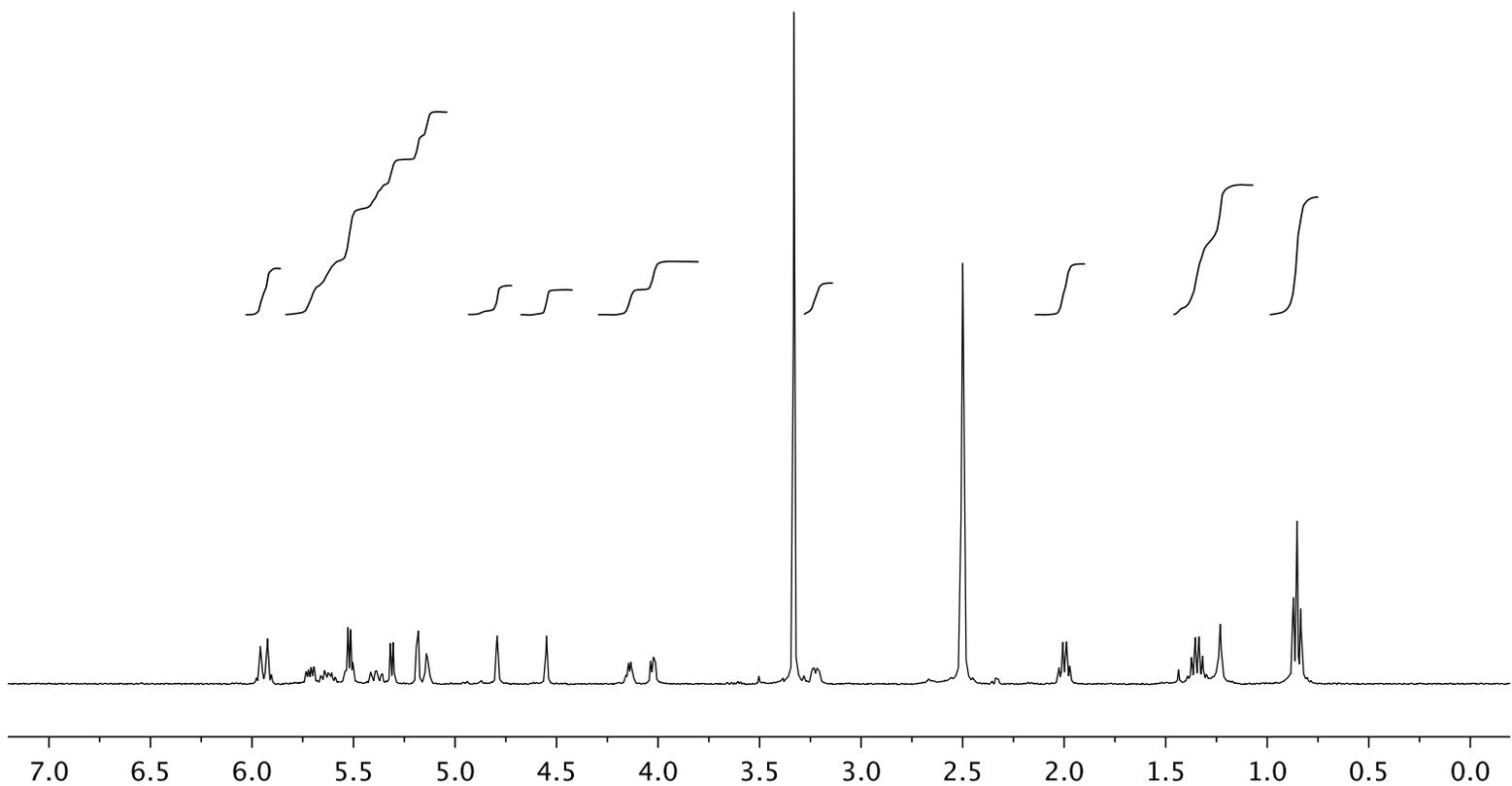


Figure S6. ^{13}C NMR spectrum of oxaspirol C (**1**) in $\text{DMSO-}d_6$

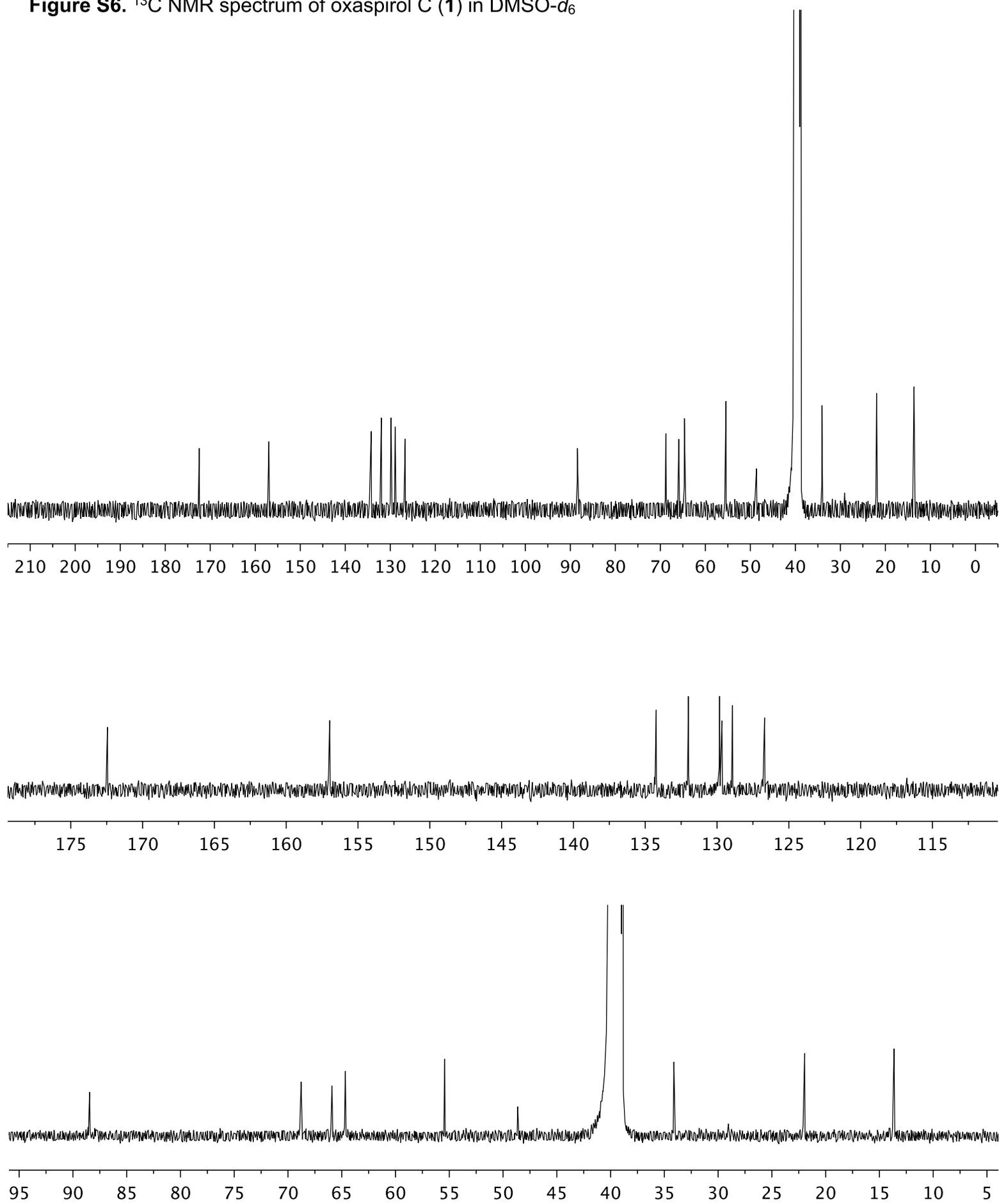


Figure S7. ^1H - ^1H gCOSY spectrum of oxaspirol C (**1**) in $\text{DMSO-}d_6$

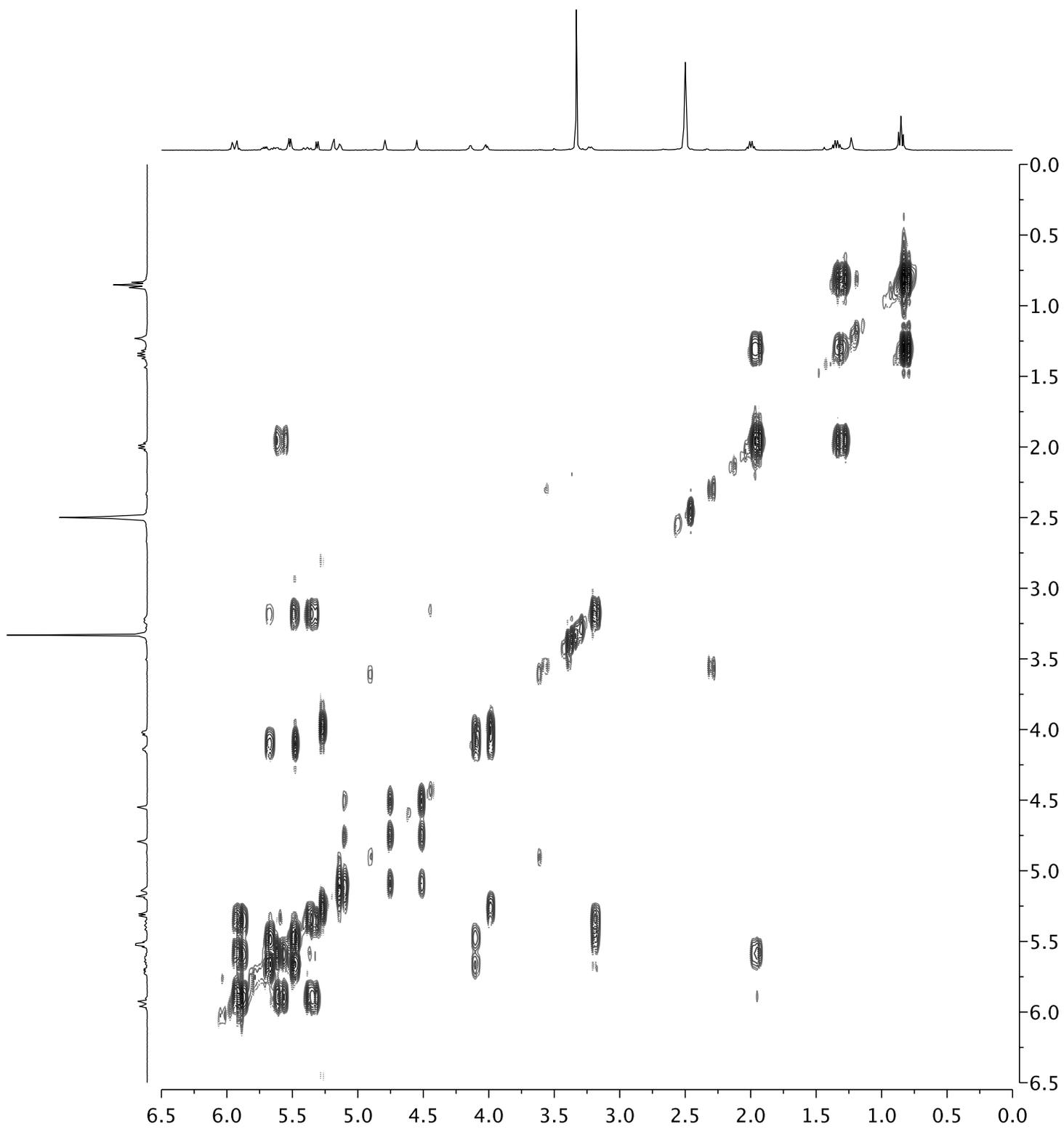


Figure S8. ^1H - ^1H ROESY spectrum of oxaspirol C (**1**) in $\text{DMSO-}d_6$

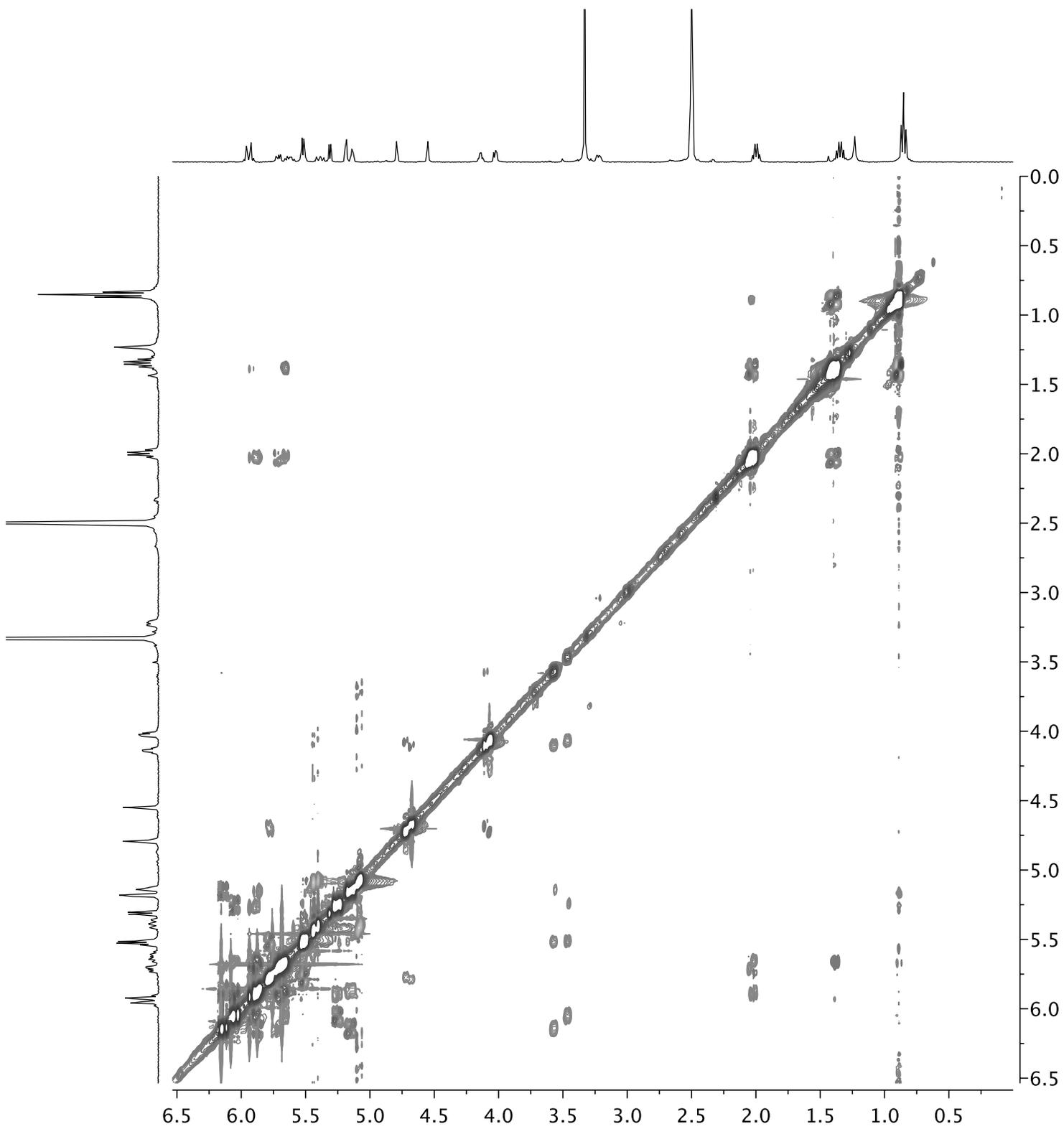


Figure S9. ^1H - ^{13}C HSQC spectrum of oxaspirol C (**1**) in $\text{DMSO-}d_6$

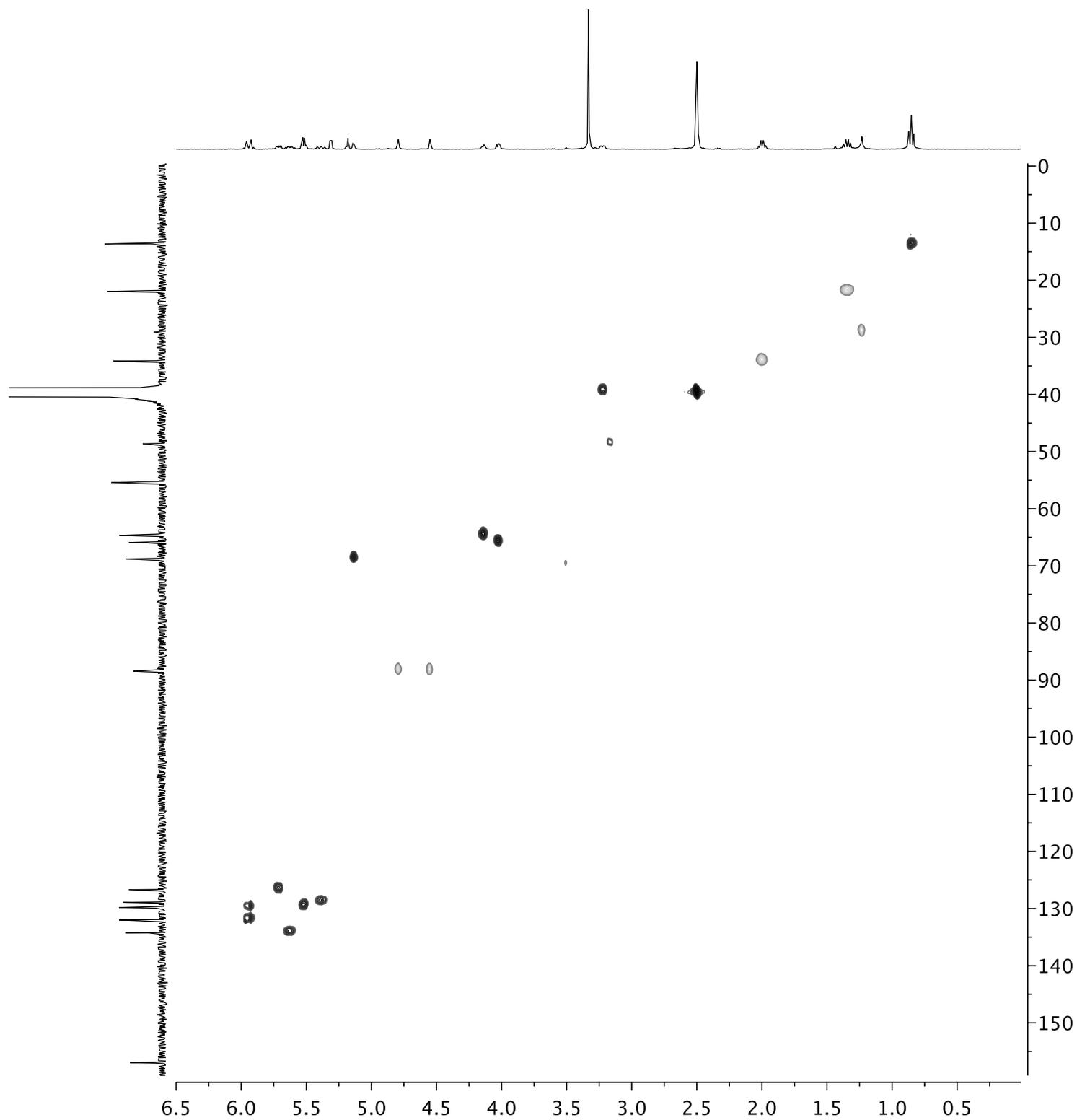


Figure S10. ^1H - ^{13}C HMBC spectrum of oxaspirol C (1) in $\text{DMSO-}d_6$

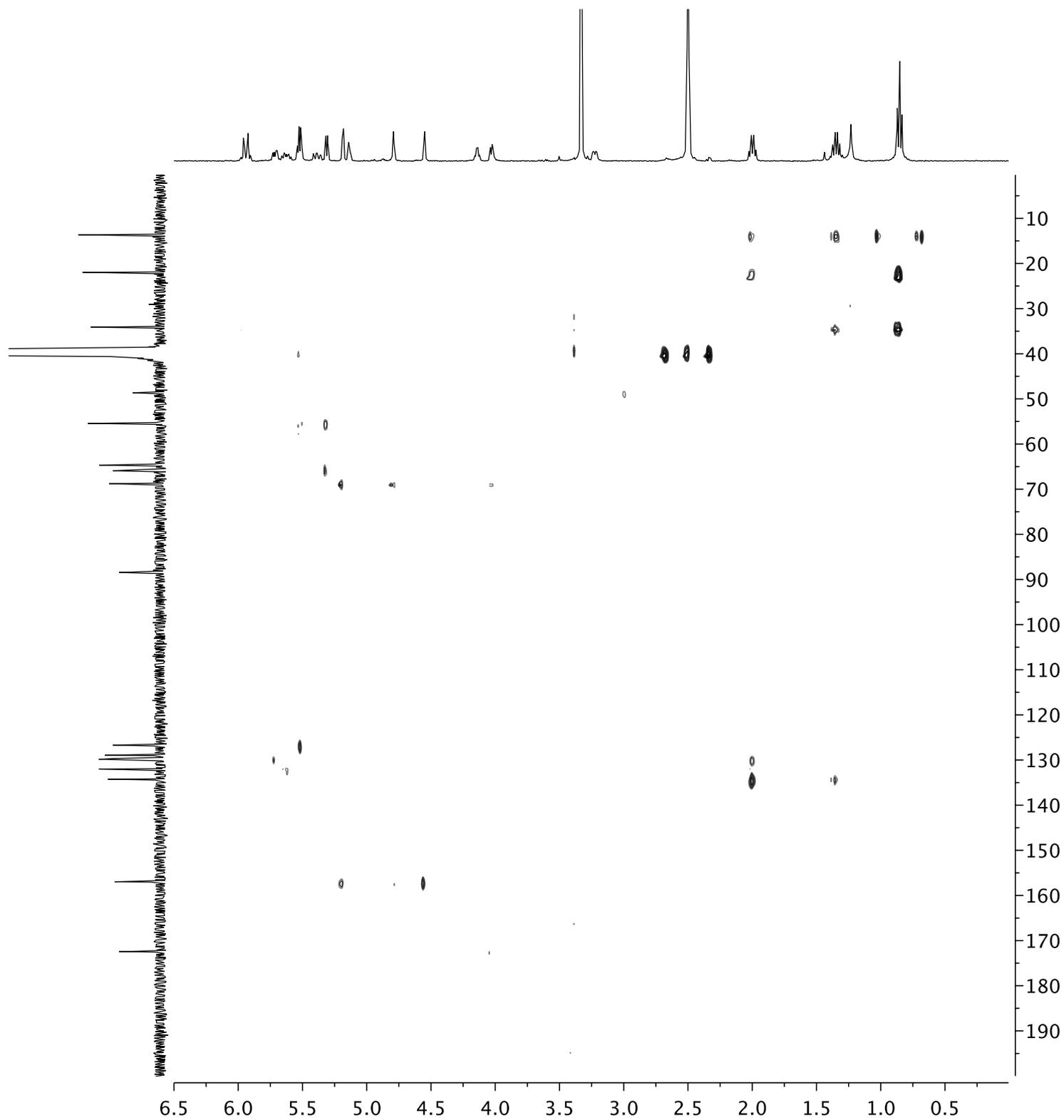


Figure S11. ^1H NMR spectrum of oxaspirol B (**2**) in CDCl_3

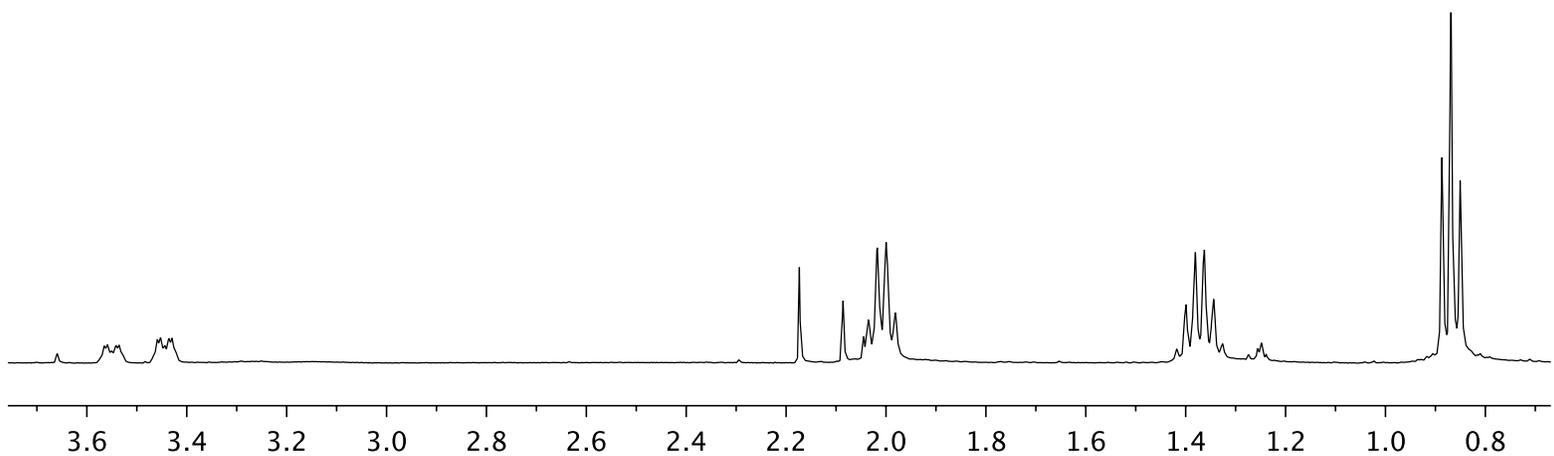
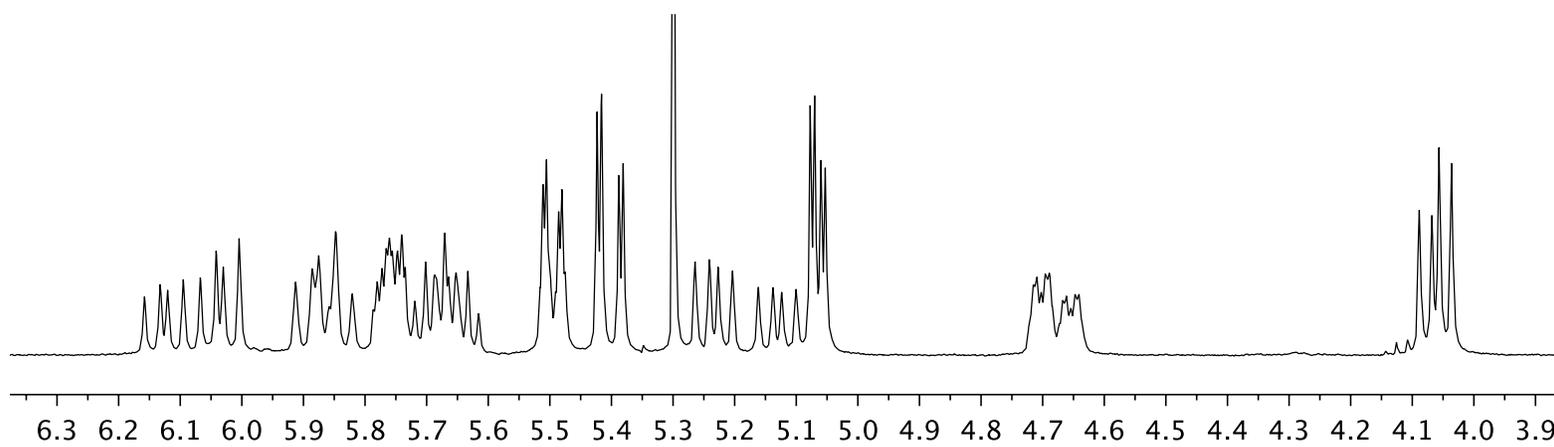
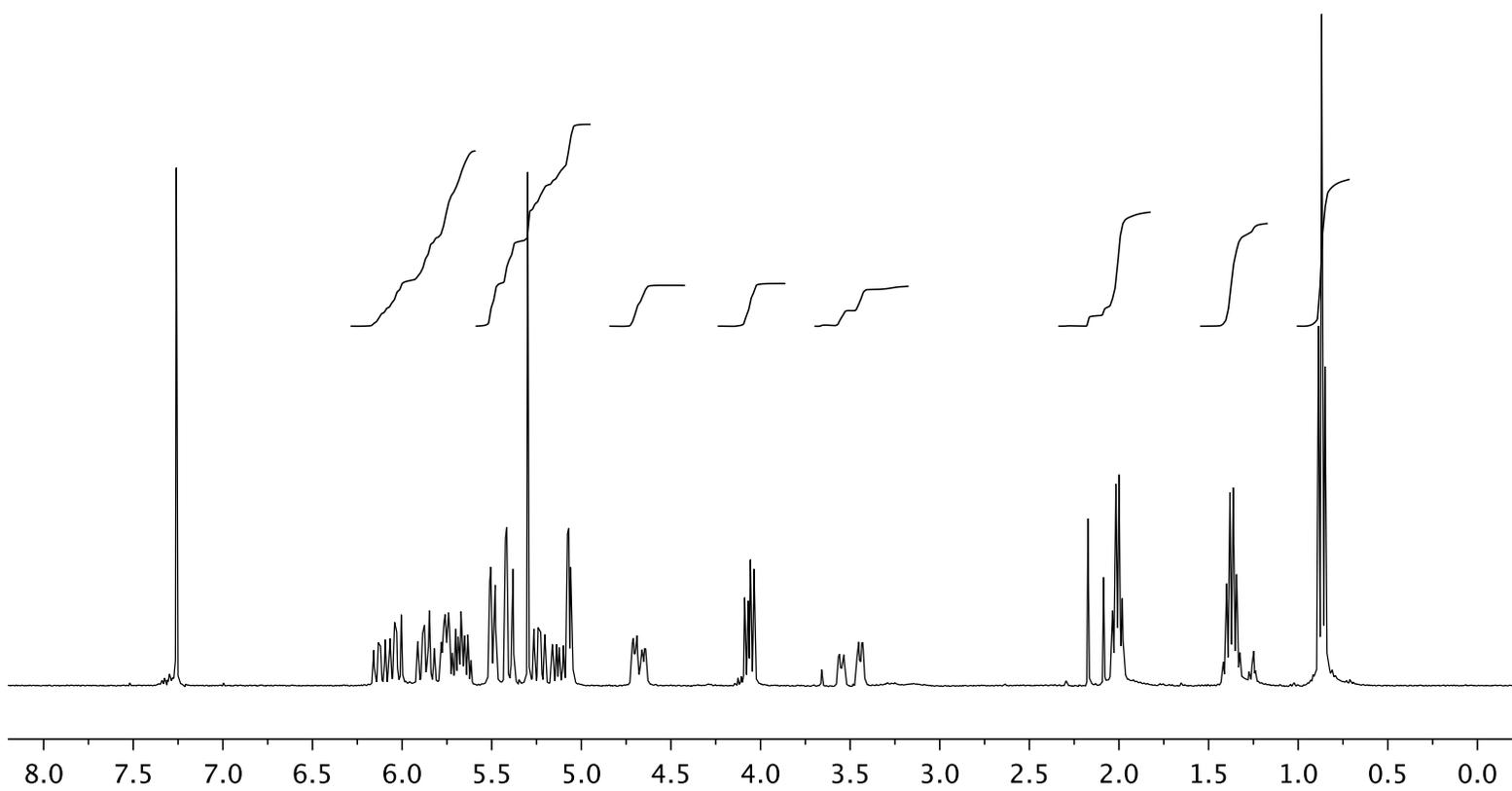


Figure S12. ^{13}C NMR spectrum of oxaspirol B (**2**) in CDCl_3

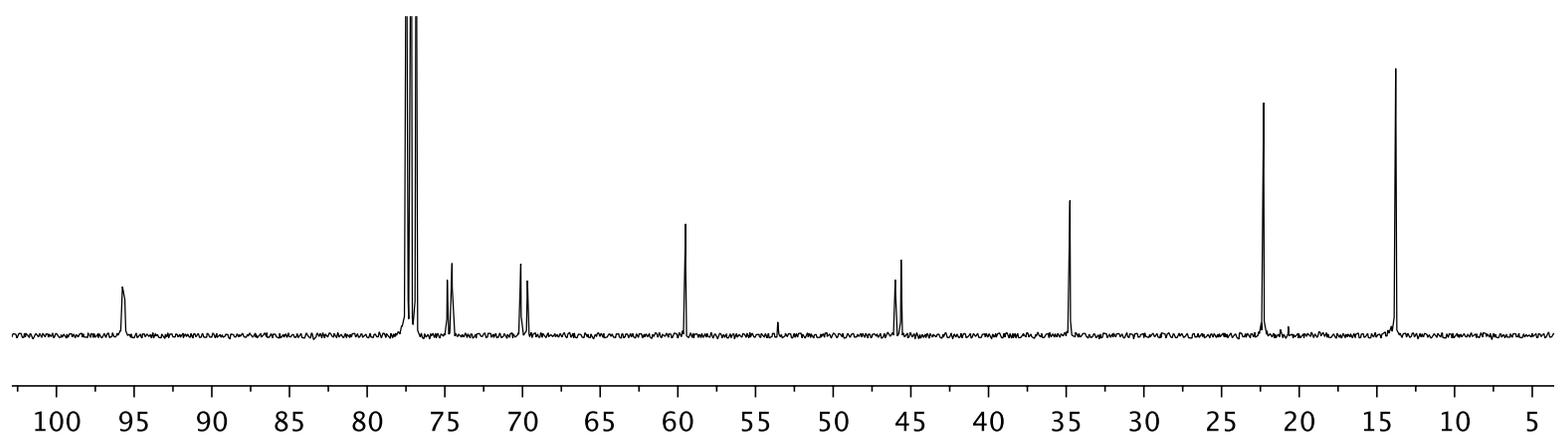
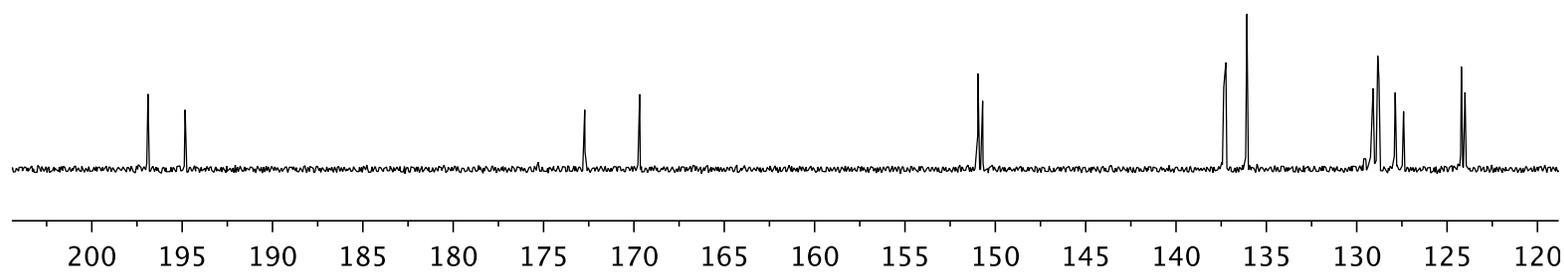
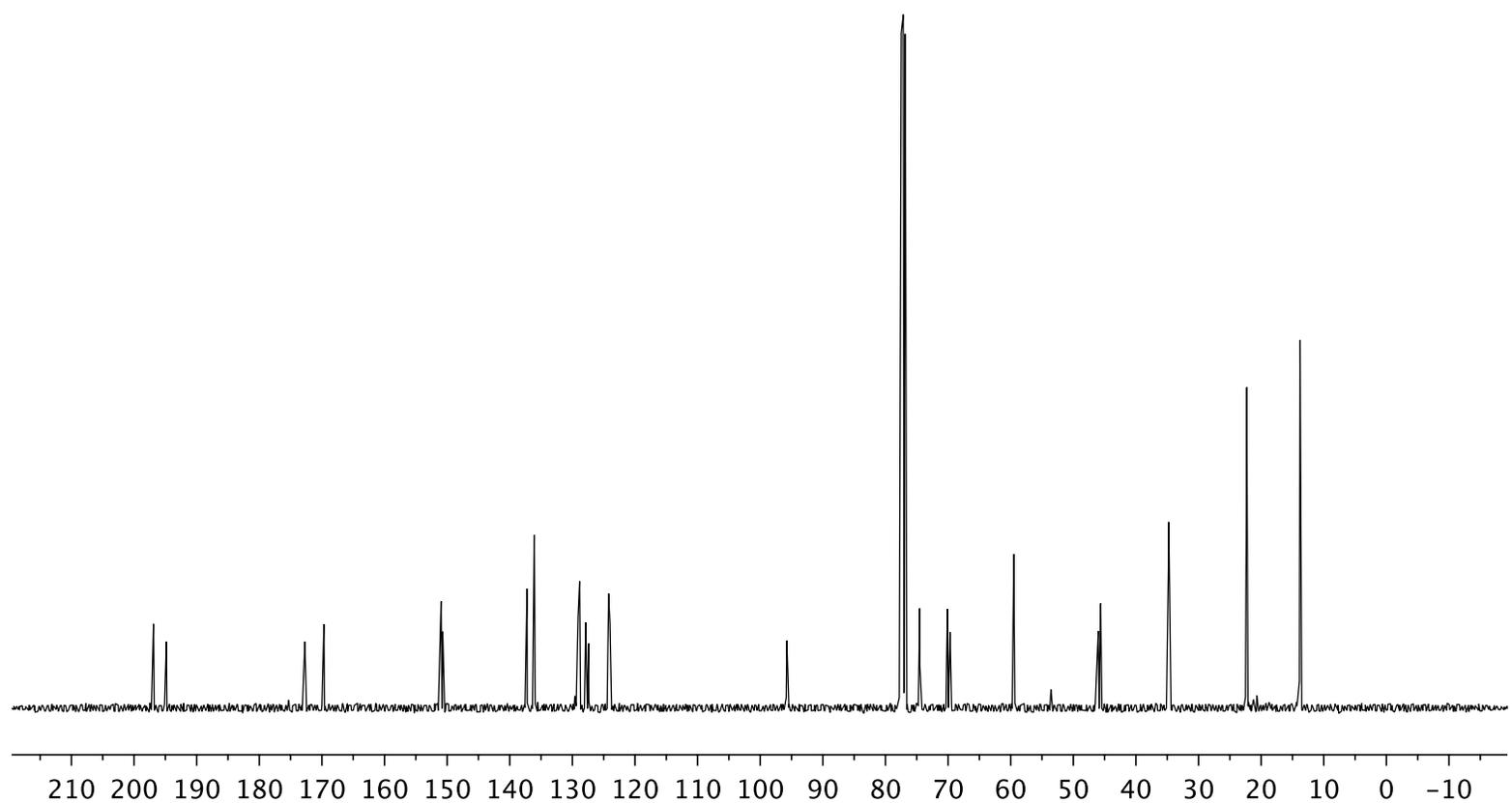


Figure S13. ^1H - ^1H gCOSY spectrum of oxaspirol B (**2**) in CDCl_3

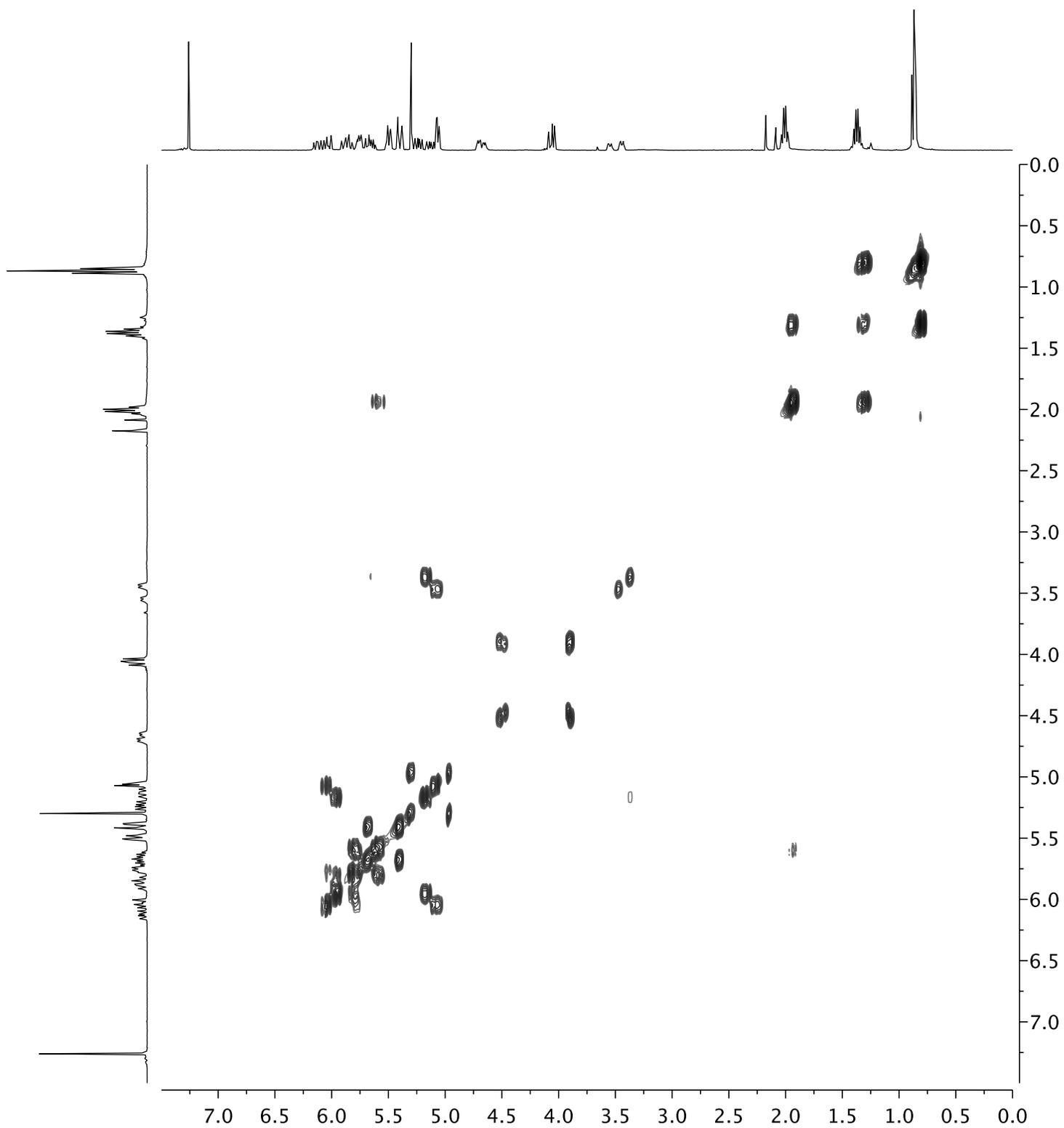


Figure S14. ^1H - ^1H ROESY spectrum of oxaspirol B (**2**) in CDCl_3

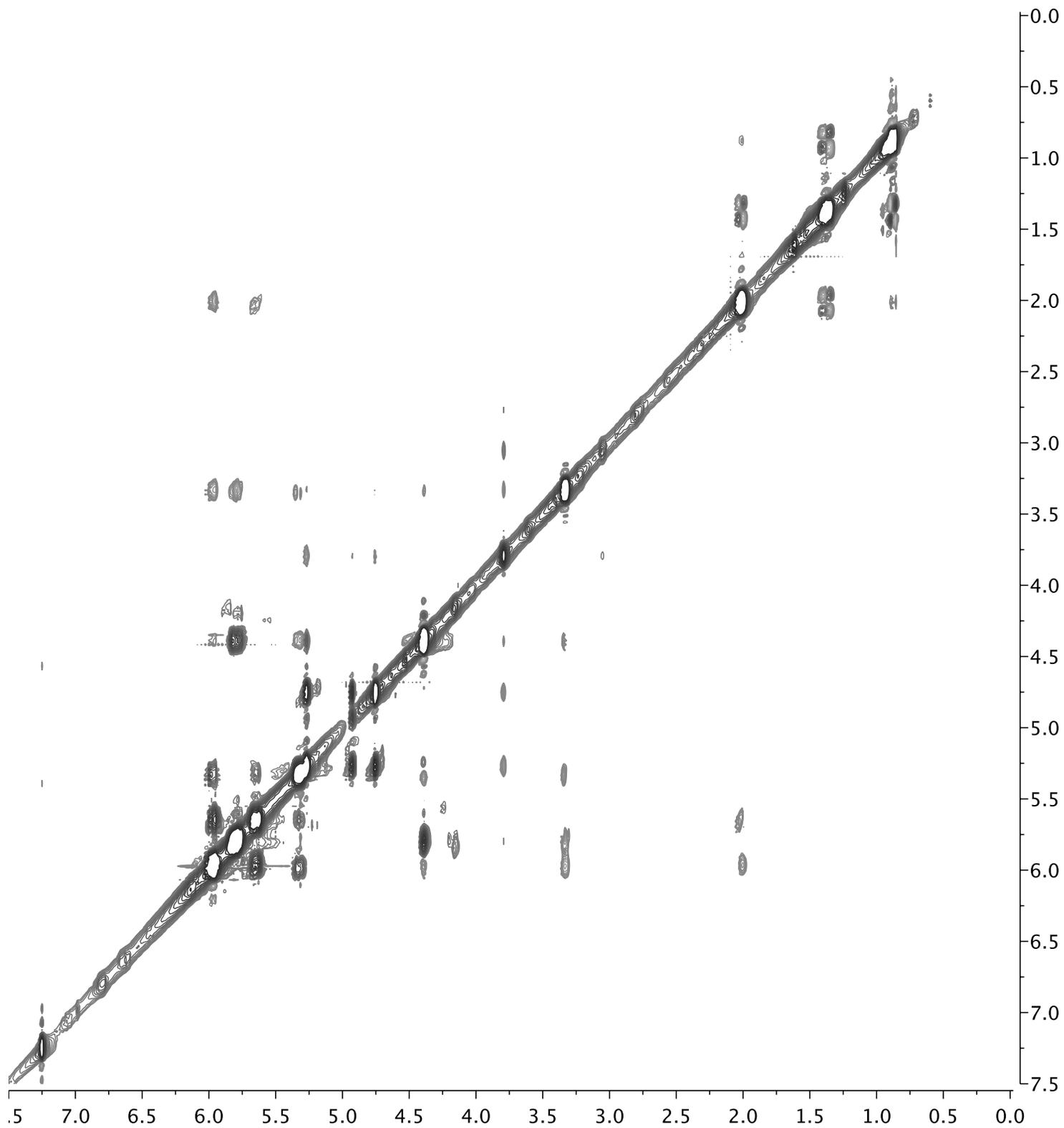


Figure S15. ^1H - ^{13}C HSQC spectrum of oxaspirol B (**2**) in CDCl_3

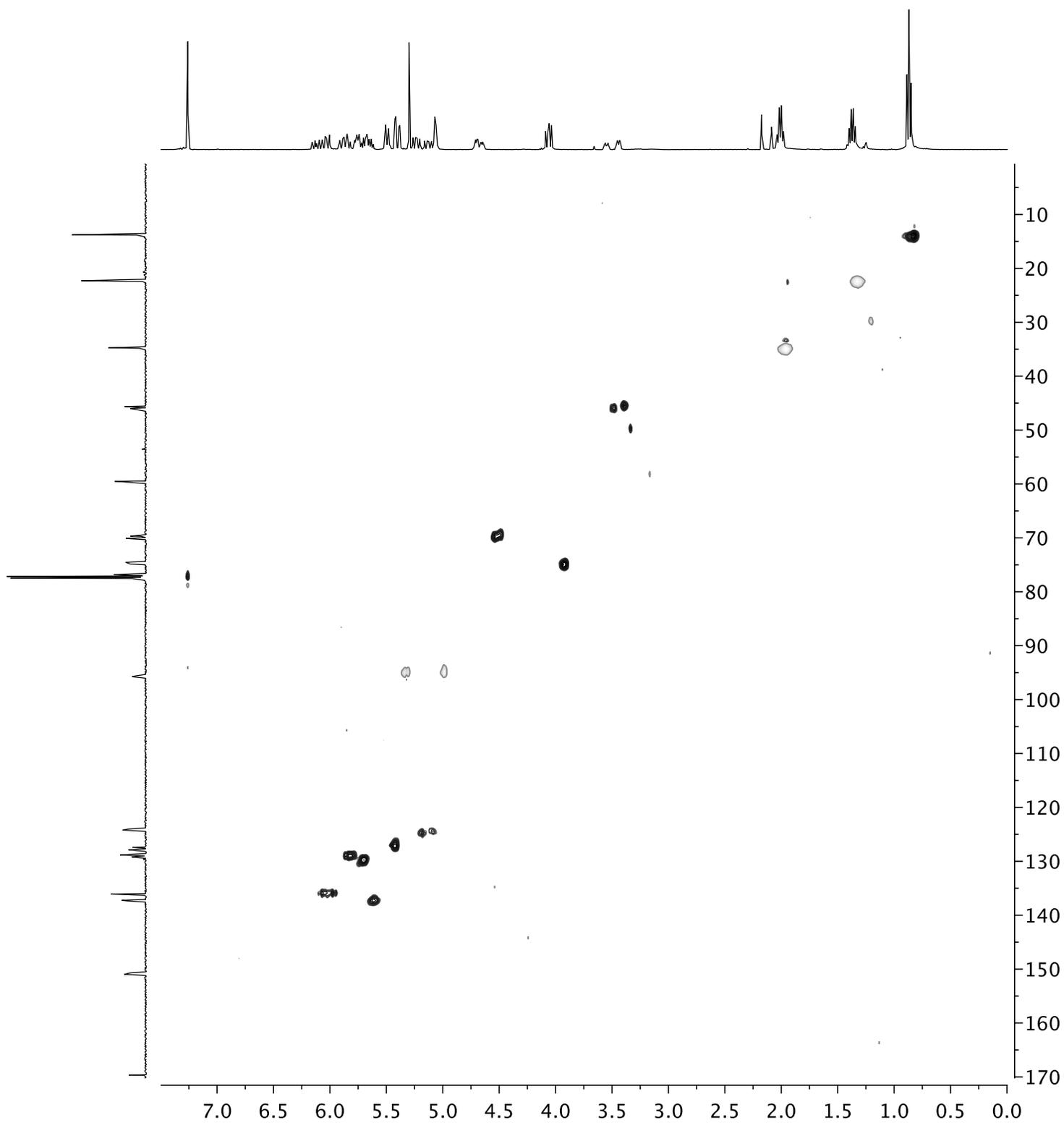


Figure S16. ^1H - ^{13}C HMBC spectrum of oxaspirol B (**2**) in CDCl_3

