

Supporting Information

for

Synthesis of thiazolidines via regioselective addition of unsymmetric thioureas to maleic acid derivatives

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Experimental Section

General Information: NMR spectroscopic data were recorded with Bruker DPX-300, Bruker Avance 400 and Bruker Avance 500 spectrometers (300.13, 400.13 and 500.03 MHz for ^1H , 75.47, 100.61 and 125.76 MHz for ^{13}C , respectively) in CDCl_3 and $\text{DMSO}-d_6$ and were referenced to residual solvent proton signals ($\delta_{\text{H}} = 7.26$ and 2.50 ppm, respectively) and solvent carbon signals ($\delta_{\text{C}} = 77.00$ and 39.52 ppm, respectively). DEPT spectra were used for carbon atoms signals assignment. The NMR spectroscopic signals of the thiazolidines atoms on the phenyl ring of former thiourea are referred to as “A” and the atoms on the phenyl ring of former maleimide to as “B”. The diastereotopic protons of the CH_2 -group of thiazolidines are marked by H^{a} and H^{b} . Melting points were determined with a Stuart SMP30 instrument and are uncorrected. Mass spectra were recorded with a Bruker Maxis HRMS-ESI-qTOF spectrometer (electrospray ionization mode). Single crystal X-ray data were obtained using an Agilent Technologies SuperNova Atlas and an Agilent Technologies Xcalibur Eos diffractometers. The samples were measured at a temperature of 100 K. The structures have been solved by the direct methods and refined by means of the SHELXL-97 program¹ incorporated in the OLEX2 program package.² The carbon-bound H atoms were placed in calculated positions and were included in the refinement in the ‘riding’ model approximation, with $U_{\text{iso}}(\text{H})$ set to $1.2U_{\text{eq}}(\text{C})$ and C–H 0.97 Å for the CH_2 groups and $U_{\text{iso}}(\text{H})$ set to $1.5U_{\text{eq}}(\text{N})$ and C–H 0.96 Å for the CH_3 groups. Empirical absorption correction was applied in CrysAlisPro program complex³ using spherical harmonics, implemented in SCALE3 ABSPACK scaling algorithm. Column chromatography was performed with Merck silica gel 60 (230-400 mesh). For TLC analysis Alugram SIL G/UV₂₅₄ (Macherey-Nagel) plates were used. *N*-Benzyl-*N'*-phenylthiourea⁴ was already available in our laboratory.

General procedure for the preparation of thioureas 3a-j.⁵ Stirred mixture of appropriate amine (23 mmol) and isothiocyanate (23 mmol) in ethanol (25 mL) was refluxed until the reaction completion indicated by TLC. The reaction mixture was cooled in an ice bath, a precipitate was filtered and recrystallized from ethanol.

***N*-Ethyl-*N'*-(4-methoxyphenyl)thiourea (3a).** Pale beige crystals, yield 3.61 g (75%), mp 130-132 °C (lit.⁶ 147 °C). ^1H NMR (400 MHz, CDCl_3): $\delta = 1.15$ (t, $J = 7.2$ Hz, 3 H, CH_2CH_3); 3.65 (dq, $J = 5.5, 7.2$ Hz, 2 H, CH_2); 3.82 (s, 3 H, OCH_3); 5.74 (br s, 1 H, NH); 6.92-6.96 (m, 2 H, H^{m}); 7.12-7.16 (m, 2 H, H^{o}); 7.55 (br s, 1 H, NH) ppm.

***N*-Ethyl-*N'*-(4-methylphenyl)thiourea (3b).**⁷ Colorless crystals, yield 3.26 g (73%), mp 97-98 °C (lit.⁸ 95-96 °C). ^1H NMR (400 MHz, CDCl_3): $\delta = 1.16$ (t, $J = 7.2$ Hz, 3 H, CH_2CH_3); 2.36 (s, 3 H, CH_3); 3.65 (dq, $J = 5.4, 7.3$ Hz, 2 H, CH_2); 5.89 (br s, 1 H, NH); 7.08 (d, $J = 8.2$ Hz, 2 H, H^{m}); 7.22 (d, $J = 8.2$ Hz, 2 H, H^{o}); 7.72 (br s, 1 H, NH) ppm.

***N*-Ethyl-*N'*-phenylthiourea (3c).**⁹ Colorless crystals, yield 2.57 g (62%), mp 100-101 °C (lit.¹⁰ 101-102 °C). ¹H NMR (400 MHz, CDCl₃): δ = 1.18 (t, *J* = 7.2 Hz, 3 H, CH₃); 3.67 (dq, *J* = 6.8, 7.2 Hz, 2 H, CH₂); 5.97 (br s, 1 H, NH); 7.20 (d, *J* = 7.5 Hz, 2 H, H^o); 7.30 (t, *J* = 7.5 Hz, 1 H, H^p); 7.41-7.45 (m, 2 H, H^m); 7.79 (br s, 1 H, NH) ppm.

***N*-(4-Chlorophenyl)-*N'*-ethylthiourea (3d).**⁷ Colorless crystals, yield 3.80 g (77%), mp 116-117 °C (lit.¹¹ 119 °C). ¹H NMR (400 MHz, CDCl₃): δ = 1.19 (t, *J* = 7.3 Hz, 3 H, CH₃); 3.65 (dq, *J* = 7.3, 7.3 Hz, 2 H, CH₂); 5.90 (br s, 1 H, NH); 7.16 (d, *J* = 8.5 Hz, 2 H, H^o); 7.39 (d, *J* = 8.5 Hz, 2 H, H^m); 7.95 (br s, 1 H, NH) ppm.

***N*-Ethyl-*N'*-(4-fluorophenyl)thiourea (3e).** Pale beige crystals, yield 3.18 g (70%), mp 79-80 °C (lit.¹² 81-82 °C). ¹H NMR (400 MHz, CDCl₃): δ = 1.16 (t, *J* = 7.2 Hz, 3 H, CH₃); 3.65 (dq, *J* = 7.0, 7.3 Hz, 2 H, CH₂); 5.84 (br s, 1 H, NH); 7.08-7.13 (m, 2 H, H^o); 7.19-7.22 (m, 2 H, H^m); 8.13 (br s, 1 H, NH) ppm.

¹³C NMR (100 MHz, CDCl₃): δ = 14.2 (CH₃); 40.2 (CH₂); 117.0 (d, *J* = 22.7 Hz, C^m); 127.8 (d, *J* = 8.7 Hz, C^o); 132.0 (d, *J* = 2.6 Hz, Cⁱ); 161.4 (d, *J* = 248.2 Hz, C^p); 180.7 (C=S) ppm.

***N*-Ethyl-*N'*-(4-nitrophenyl)thiourea (3f).** Yellow crystals, yield 3.39 g (66%), mp 174-175 °C (lit.⁵ 159-160 °C). ¹H NMR (400 MHz, DMSO-*d*₆): δ = 1.15 (t, *J* = 7.2 Hz, 3 H, CH₃); 3.48-3.54 (m, 2 H, CH₂); 7.79-7.82 (m, 2 H, H^o); 8.15-8.18 (m, 2 H, H^m); 8.26 (br s, 1 H, NH); 10.06 (br s, 1 H, NH) ppm.

¹³C NMR (100 MHz, DMSO-*d*₆): δ = 13.7 (CH₃); 38.8 (CH₂); 120.3, 124.5 (C^{m,o}); 141.7, 146.4 (C^{i,p}); 179.8 (C=S) ppm.

***N*-(2,6-Dimethylphenyl)-*N'*-methylthiourea (3g).**¹³ Pink crystals, yield 3.13 g (70%), mp 137-139 °C (lit.¹⁴ 140-141 °C). ¹H NMR (400 MHz, CDCl₃): δ = 2.26 (s, 6 H, 2 CH₃); 3.09 (d, *J* = 4.7 Hz, 3 H, NCH₃); 5.36 (br s, 1 H, NH); 7.14 (d, *J* = 7.4 Hz, 2 H, H^m); 7.18-7.22 (m, 1 H, H^p); 7.35 (br s, 1 H, NH) ppm.

***N*-Isopropyl-*N'*-phenylthiourea (3h).**¹⁵ Colorless crystals, yield 3.44 g (77%), mp 101-102 °C (lit.¹⁶ 101 °C). ¹H NMR (400 MHz, CDCl₃): δ = 1.20 (d, *J* = 6.6 Hz, 6 H, CH(CH₃)₂); 4.60 (dsept, *J* = 6.6, 6.6 Hz, 1 H, CH(CH₃)₂); 5.79 (br s, 1 H, NH); 7.18 (d, *J* = 7.6 Hz, 2 H, H^o); 7.28-7.32 (m, 1 H, H^p); 7.42-7.46 (m, 2 H, H^m); 7.67 (br s, 1 H, NH) ppm.

***N*-Cyclohexyl-*N'*-phenylthiourea (3i).**¹⁷ Colorless crystals, yield 4.67 g (87%), mp 151-153 °C (lit.¹⁸ 150.1-150.9 °C). ¹H NMR (400 MHz, CDCl₃): δ = 1.07-1.17 (m, 3 H, c-Hex); 1.34-1.44 (m, 2 H, c-Hex); 1.57-1.68 (m, 3 H, c-Hex); 2.03-2.07 (m, 2 H, c-Hex); 4.24-4.29 (br s, 1 H, NCH); 5.89 (br s, 1 H, NH); 7.18 (d, *J* = 7.6 Hz, 2 H, H^o); 7.27-7.31 (m, 1 H, H^p); 7.41-7.45 (m, 2 H, H^m); 7.75 (br s, 1 H, NH) ppm.

***N*-tert-Butyl-*N'*-phenylthiourea (3j).**¹⁹ Colorless crystals, yield 3.96 g (83%), mp 120-122 °C (lit.¹⁹ 120.5-121.5 °C). ¹H NMR (400 MHz, DMSO-*d*₆): δ = 1.47 (s, 9 H, 3 CH₃); 7.04-7.08 (m, 1 H,

H^p); 7.26-7.30 (m, 2 H, H^o); 7.35 (br s, 1 H, NH); 7.43 (d, *J* = 7.6 Hz, 2 H, H^m); 9.26 (br s, 1 H, NH) ppm.

***N*-Ethyl-*N'*-isopropylthiourea (3k).** Isopropylamine (1.0 mL, 0.69 g, 11.7 mmol) was added to a stirred solution of ethylisothiocyanate (1.0 mL, 1.02 g, 11.7 mmol) in diethyl ether (5 mL). After 2 h a precipitate was filtered, washed with diethyl ether (10 mL) and dried. Colorless crystals, yield 1.42 g, 83%, mp 108-109 °C.

¹H NMR (400 MHz, DMSO-*d*₆): δ = 1.04 (t, *J* = 7.2 Hz, 3 H, CH₂CH₃); 1.08 (d, *J* = 6.5 Hz, 6 H, CH(CH₃)₂); 3.34 (br s, 2 H, CH₂); 4.22 (br s, 1 H, CH); 7.11 (br s, 1 H, NH); 7.16 (br s, 1 H, NH) ppm.

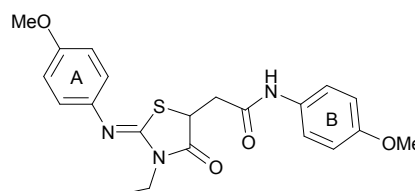
¹H NMR (400 MHz, CDCl₃): δ = 1.22 (t, *J* = 7.2 Hz, 3 H, CH₂CH₃); 1.23 (d, *J* = 6.5 Hz, 6 H, CH(CH₃)₂); 3.42 (br s, 2 H, CH₂); 4.23 (br s, 1 H, CH); 5.57 (br s, 1 H, NH); 5.73 (br s, 1 H, NH) ppm.

¹³C NMR (100 MHz, CDCl₃): δ = 14.1 (CH₂CH₃); 22.5 (CH(CH₃)₂); 38.9 (CH₂); 46.0 (CH); 180.1 (C=S) ppm.

HRMS (ESI), *m/z*: calcd for C₆H₁₄N₂S [M+H]⁺ 147.0950, found 147.0954.

General procedure for the preparation of thiazolidines 5aa-ff, 5ga-gf. Stirred mixture of thiourea **3a-g** (1 mmol) and *N*-arylmaleimide **4a-g** (1 mmol) in *i*-PrOH (5 mL) was refluxed for 3-5 h, and then was poured into water (30 mL). A precipitate was filtered, washed with water and recrystallized from ethanol/water (1:1) mixture.

2-(3-Ethyl-2-(4-methoxyphenyl)imino-4-oxo-1,3-thiazolidin-5-yl)-*N*-(4-methoxyphenyl)-acetamide (5aa). Beige crystals, yield 320 mg (77%), mp 171-172 °C.



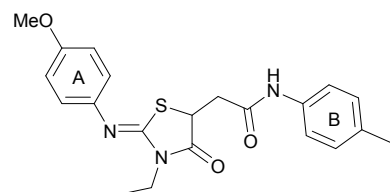
¹H NMR (300 MHz, CDCl₃): δ = 1.29 (t, *J* = 7.1 Hz, 3 H, CH₂CH₃); 2.76 (dd, *J* = 16.1, 10.0 Hz, 1 H, H^a); 3.32 (dd, *J* = 16.1, 3.8 Hz, 1 H, H^b); 3.76 (s, 3 H, OCH₃); 3.78 (s, 3 H, OCH₃); 3.86-3.99 (m, 2 H, CH₂CH₃); 4.42 (dd, *J* = 10.0, 3.8 Hz, 1 H, SCH);

6.79-6.94 (m, 6 H, H^{mB}, H^{o,mA}); 7.34 (d, *J* = 8.9 Hz, 2 H, H^{oB}); 7.65 (br s, 1 H, NH) ppm.

¹³C NMR (75 MHz, CDCl₃): δ = 12.6 (CH₂CH₃); 38.2, 40.3 (CH₂); 43.8 (SCH); 55.4 (2 OCH₃); 114.1, 114.5 (C^{mA}, C^{mB}); 121.8, 122.1 (C^{oA}, C^{oB}); 130.4 (C^{iB}); 141.1 (C^{iA}); 153.2 (C=N); 156.6, 156.7 (C^{pA}, C^{pB}); 166.9, 173.9 (C=O) ppm.

HRMS (ESI), *m/z*: calcd for C₂₁H₂₃N₃O₄S [M+H]⁺ 414.1482, found 414.1484.

2-(3-Ethyl-2-(4-methoxyphenyl)imino-4-oxo-1,3-thiazolidin-5-yl)-*N*-(4-methylphenyl)-acetamide (5ab). Beige crystals, yield 301 mg (76%), mp 169-170 °C.



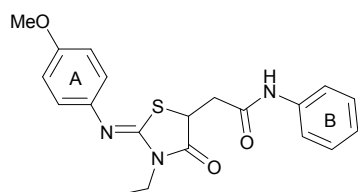
¹H NMR (400 MHz, CDCl₃): δ = 1.30 (t, *J* = 7.0 Hz, 3 H, CH₂CH₃); 2.29 (s, 3 H, CH₃); 2.79 (dd, *J* = 16.2, 9.9 Hz, 1 H, H^a); 3.35 (dd, *J* = 16.2, 4.0 Hz, 1 H, H^b); 3.79 (s, 3 H, OCH₃); 3.87-3.99 (m, 2 H, CH₂CH₃); 4.44 (dd, *J* = 9.9, 4.0 Hz, 1 H, SCH); 6.84-6.87

(m, 2 H, H^{mA}); 6.91-6.93 (m, 2 H, H^{oA}); 7.10 (d, $J = 8.2$ Hz, 2 H, H^{mB}); 7.34 (d, $J = 8.2$ Hz, 2 H, H^{oB}); 7.46 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, CDCl_3): $\delta = 12.6$ (CH_2CH_3); 20.8 (CH_3); 38.2, 40.4 (CH_2); 43.8 (SCH); 55.4 (OCH_3); 114.4 (C^{mA}); 119.9 (C^{oB}); 122.1 (C^{oA}); 129.5 (C^{mB}); 134.3, 134.7 ($C^{i,pB}$); 141.1 (C^{iA}); 153.3 ($\text{C}=\text{N}$); 156.7 (C^{pA}); 167.0, 173.9 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $[\text{M}+\text{H}]^+$ 398.1533, found 398.1539.

2-(3-Ethyl-2-(4-methoxyphenyl)imino-4-oxo-1,3-thiazolidin-5-yl)-*N*-phenylacetamide (5ac).



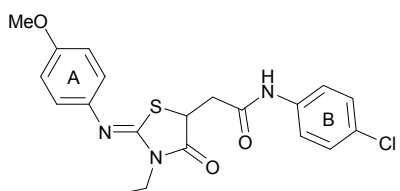
Pale yellow crystals, yield 295 mg (80%), mp 168-169 °C.

^1H NMR (300 MHz, CDCl_3): $\delta = 1.30$ (t, $J = 7.1$ Hz, 3 H, CH_2CH_3); 2.80 (dd, $J = 16.3, 10.0$ Hz, 1 H, H^a); 3.37 (dd, $J = 16.3, 3.6$ Hz, 1 H, H^b); 3.78 (s, 3 H, OCH_3); 3.87-3.99 (m, 2 H, CH_2CH_3); 4.43 (dd, $J = 10.0, 3.6$ Hz, 1 H, SCH); 6.84-6.94 (m, 4 H, $H^{m,oA}$); 7.07-7.13 (m, 1 H, H^{pB}); 7.26-7.32 (m, 2 H, H^{mB}); 7.46 (d, $J = 8.1$ Hz, 2 H, H^{oB}); 7.80 (br s, 1 H, NH) ppm.

^{13}C NMR (75 MHz, CDCl_3): $\delta = 12.6$ (CH_2CH_3); 38.3, 40.5 (CH_2); 43.7 (SCH); 55.4 (OCH_3); 114.5 (C^{mA}); 119.9 (C^{oB}); 122.1 (C^{oA}); 124.6 (C^{pB}); 129.0 (C^{mB}); 137.3 (C^{iB}); 141.1 (C^{iA}); 153.2 ($\text{C}=\text{N}$); 156.8 (C^{pA}); 167.1, 173.9 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{20}\text{H}_{21}\text{N}_3\text{O}_3\text{S}$ $[\text{M}+\text{H}]^+$ 384.1376, found 384.1373.

***N*-(4-Chlorophenyl)-2-(3-ethyl-2-(4-methoxyphenyl)imino-4-oxo-1,3-thiazolidin-5-yl)-acetamide (5ad).** Colorless crystals, yield 284 mg (68%), mp 174-175 °C.

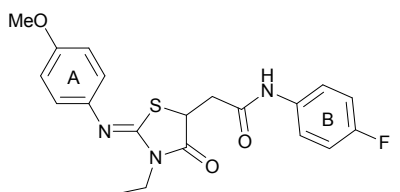


^1H NMR (400 MHz, $\text{DMSO}-d_6$): $\delta = 1.20$ (t, $J = 7.0$ Hz, 3 H, CH_2CH_3); 2.93 (dd, $J = 16.6, 9.4$ Hz, 1 H, H^a); 3.24 (dd, $J = 16.6, 3.5$ Hz, 1 H, H^b); 3.73 (s, 3 H, OCH_3); 3.74-3.86 (m, 2 H, CH_2CH_3); 4.54 (dd, $J = 9.4, 3.5$ Hz, 1 H, SCH); 6.85-6.93 (m, 4 H, $H^{m,oA}$); 7.34 (d, $J = 8.8$ Hz, 2 H, H^{mB}); 7.55 (d, $J = 8.8$ Hz, 2 H, H^{oB}); 10.25 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, $\text{DMSO}-d_6$): $\delta = 12.2$ (CH_2CH_3); 37.4, 38.7 (CH_2); 43.3 (SCH); 55.2 (OCH_3); 114.4 (C^{mA}); 120.6, 122.0 ($C^{oA,B}$); 126.9 (C^{pB}); 128.7 (C^{mB}); 137.6 (C^{iB}); 141.2 (C^{iA}); 154.1, 156.1 ($\text{C}=\text{N}$, C^{pA}); 167.9, 173.5 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{20}\text{H}_{20}\text{ClN}_3\text{O}_3\text{S}$ $[\text{M}+\text{H}]^+$ 418.0987, found 418.0983.

2-(3-Ethyl-2-(4-methoxyphenyl)imino-4-oxo-1,3-thiazolidin-5-yl)-*N*-(4-fluorophenyl)-acetamide (5ae). Colorless crystals, yield 320 mg (80%), mp 147-148 °C.

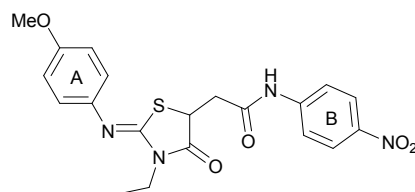


^1H NMR (400 MHz, CDCl_3): $\delta = 1.30$ (t, $J = 7.1$ Hz, 3 H, CH_2CH_3); 2.80 (dd, $J = 16.2, 9.9$ Hz, 1 H, H^a); 3.34 (dd, $J = 16.2, 4.0$ Hz, 1 H, H^b); 3.79 (s, 3 H, OCH_3); 3.86-4.00 (m, 2 H, CH_2CH_3); 4.43 (dd, $J = 9.9, 4.0$ Hz, 1 H, SCH); 6.85-6.93 (m, 4 H, $H^{m,oA}$); 6.97-7.01 (m, 2 H, H^{mB}); 7.41-7.46 (m, 2 H, H^{oB}); 7.64 (br s, 1 H, NH) ppm.

^{13}C NMR (75 MHz, CDCl_3): δ = 12.6 (CH_2CH_3); 38.3, 40.4 (CH_2); 43.7 (SCH); 55.4 (OCH_3); 114.5 (C^{mA}); 115.7 (d, J = 22.2 Hz, C^{mB}); 121.7 (d, J = 8.1 Hz, C^{oB}); 122.1 (C^{oA}); 133.3 (d, J = 2.4 Hz, C^{iB}); 141.0 (C^{iA}); 153.0, 156.8 ($\text{C}=\text{N}$, C^{pA}); 159.5 (d, J = 244.0 Hz, C^{pB}); 167.1, 173.9 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{20}\text{H}_{20}\text{FN}_3\text{O}_3\text{S}$ $[\text{M}+\text{H}]^+$ 402.1282, found 402.1280.

2-(3-Ethyl-2-(4-methoxyphenyl)imino-4-oxo-1,3-thiazolidin-5-yl)-N-(4-nitrophenyl)-acetamide (5af). Yellow crystals, yield 334 mg (78%), mp 168-169 °C.



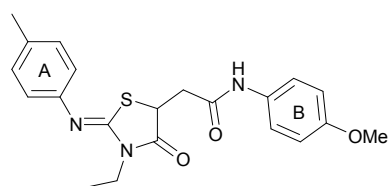
^1H NMR (300 MHz, CDCl_3): δ = 1.31 (t, J = 7.0 Hz, 3 H, CH_2CH_3); 2.89 (dd, J = 16.5, 9.1 Hz, 1 H, H^{a}); 3.42 (dd, J = 16.5, 4.4 Hz, 1 H, H^{b}); 3.79 (s, 3 H, OCH_3); 3.87-4.03 (m, 2 H, CH_2CH_3); 4.46 (dd, J = 9.1, 4.4 Hz, 1 H, SCH); 6.83-6.96 (m, 4 H, $\text{H}^{\text{m,oA}}$); 7.69

(d, J = 9.0 Hz, 2 H, H^{oB}); 8.19 (d, J = 9.0 Hz, 2 H, H^{mB}); 8.48 (br s, 1 H, NH) ppm.

^{13}C NMR (75 MHz, CDCl_3): δ = 12.6 (CH_2CH_3); 38.5, 40.8 (CH_2); 43.3 (SCH); 55.4 (OCH_3); 114.5 (C^{mA}); 119.9, 122.0, 125.1 (C^{oA} , $\text{C}^{\text{o,mB}}$); 140.9, 143.3, 143.7 (C^{iA} , $\text{C}^{\text{i,pB}}$); 152.5 ($\text{C}=\text{N}$); 156.9 (C^{pA}); 167.6, 174.1 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{20}\text{H}_{20}\text{N}_4\text{O}_5\text{S}$ $[\text{M}+\text{H}]^+$ 429.1227, found 429.1234.

2-(3-Ethyl-2-(4-methylphenyl)imino-4-oxo-1,3-thiazolidin-5-yl)-N-(4-methoxyphenyl)-acetamide (5ba). Colorless crystals, yield 275 mg (69%), mp 155-157 °C.



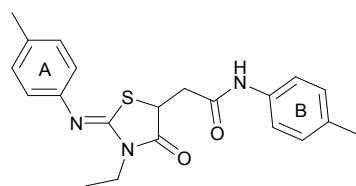
^1H NMR (400 MHz, CDCl_3): δ = 1.30 (t, J = 7.2 Hz, 3 H, CH_2CH_3); 2.31 (s, 3 H, CH_3); 2.76 (dd, J = 16.2, 9.9 Hz, 1 H, H^{a}); 3.32 (dd, J = 16.2, 3.9 Hz, 1 H, H^{b}); 3.76 (s, 3 H, OCH_3); 3.87-3.99 (m, 2 H, CH_2CH_3); 4.42 (dd, J = 9.9, 3.9 Hz, 1 H, SCH); 6.81-6.87 (m,

4 H, H^{mB} , H^{oA}); 7.12 (d, J = 8.1 Hz, 2 H, H^{mA}), 7.33-7.35 (m, 2 H, H^{oB}); 7.59 (br s, 1H, NH) ppm.

^{13}C NMR (100 MHz, CDCl_3): δ = 12.6 (CH_2CH_3); 20.9 (CH_3); 38.2, 40.3 (CH_2); 43.8 (SCH); 55.4 (OCH_3); 114.1 (C^{mB}); 120.8, 122.0 (C^{oA} , C^{oB}); 129.8 (C^{mA}); 130.3, 134.1 (C^{pA} , C^{iB}); 145.4 (C^{iA}); 153.3 ($\text{C}=\text{N}$); 156.6 (C^{pB}); 166.9, 173.9 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{21}\text{H}_{23}\text{N}_3\text{O}_3\text{S}$ $[\text{M}+\text{H}]^+$ 398.1533, found 398.1533.

2-(3-Ethyl-2-(4-methylphenyl)imino-4-oxo-1,3-thiazolidin-5-yl)-N-(4-methylphenyl)-acetamide (5bb). Colorless crystals, yield 304 mg (80%), mp 184-185 °C.



^1H NMR (400 MHz, CDCl_3): δ = 1.30 (t, J = 7.0 Hz, 3 H, CH_2CH_3); 2.30 (s, 3 H, CH_3); 2.32 (s, 3 H, CH_3); 2.78 (dd, J = 16.0, 9.9 Hz, 1 H, H^{a}); 3.35 (dd, J = 16.0, 4.0 Hz, 1 H, H^{b}); 3.89-3.98 (m, 2 H, CH_2CH_3); 4.44 (dd, J = 9.9, 4.0 Hz, 1 H, SCH); 6.86 (d, J = 8.1 Hz, 2 H, H^{oA});

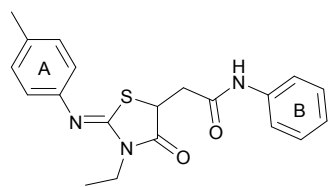
7.01-7.13 (m, 4 H, H^{mA} , H^{mB}); 7.33 (d, J = 8.2 Hz, 2 H, H^{oB}); 7.44 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, CDCl_3): δ = 12.6 (CH_2CH_3); 20.8, 20.9 (CH_3); 38.2, 40.5 (CH_2); 43.8 (SCH); 119.9, 120.8 (C^{oA} , C^{oB}); 129.5, 129.8 (C^{mA} , C^{mB}); 134.1, 134.3, 134.7 (C^{pA} , C^{iPB}); 145.4 (C^{iA}); 153.3 ($\text{C}=\text{N}$); 166.9, 173.9 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $[\text{M}+\text{H}]^+$ 382.1583, found 382.1586.

2-(3-Ethyl-2-(4-methylphenyl)imino-4-oxo-1,3-thiazolidin-5-yl)-N-phenylacetamide (5bc).

Colorless crystals, yield 268 mg (73%), mp 191-192 °C.

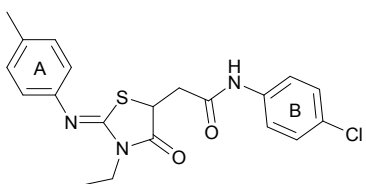


^1H NMR (300 MHz, CDCl_3): δ = 1.30 (t, J = 7.1 Hz, 3 H, CH_2CH_3); 2.32 (s, 3 H, CH_3); 2.79 (dd, J = 16.3, 10.0 Hz, 1 H, H^{a}); 3.36 (dd, J = 16.3, 3.7 Hz, 1 H, H^{b}); 3.85-4.02 (m, 2 H, CH_2CH_3); 4.43 (dd, J = 10.0, 3.7 Hz, 1 H, SCH); 6.86 (d, J = 8.2 Hz, 2 H, H^{oA}); 7.05-7.15 (m, 3 H, H^{pB} , H^{mA}); 7.27-7.33 (m, 2 H, H^{mB}); 7.45 (d, J = 7.8 Hz, 2 H, H^{oB}); 7.72 (br s, 1 H, NH) ppm.

^{13}C NMR (75 MHz, CDCl_3): δ = 12.6 (CH_2CH_3); 20.9 (CH_3); 38.3, 40.6 (CH_2); 43.8 (SCH); 119.9, 120.8 (C^{oA} , C^{oB}); 124.6 (C^{pB}); 129.0, 129.8 (C^{mA} , C^{mB}); 134.2 (C^{pA}); 137.3 (C^{iB}); 145.4 (C^{iA}); 153.2 ($\text{C}=\text{N}$); 167.1, 173.9 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{20}\text{H}_{21}\text{N}_3\text{O}_2\text{S}$ $[\text{M}+\text{H}]^+$ 368.1427, found 368.1423.

N-(4-Chlorophenyl)-2-(3-ethyl-2-(4-methylphenyl)imino-4-oxo-1,3-thiazolidin-5-yl)-acetamide (5bd). Colorless crystals, yield 307 mg (76%), mp 175-176 °C.

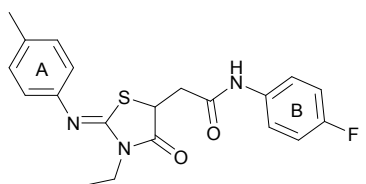


^1H NMR (400 MHz, $\text{DMSO}-d_6$): δ = 1.20 (t, J = 7.0 Hz, 3 H, CH_2CH_3); 2.25 (s, 3 H, CH_3); 2.94 (dd, J = 16.6, 9.5 Hz, 1 H, H^{a}); 3.26 (dd, J = 16.6, 3.7 Hz, 1 H, H^{b}); 3.72-3.87 (m, 2 H, CH_2CH_3); 4.54 (dd, J = 9.5, 3.7 Hz, 1 H, SCH); 6.81 (d, J = 8.2 Hz, 2 H, H^{oA}); 7.13 (d, J = 8.2 Hz, 2 H, H^{mA}); 7.34 (d, J = 8.8 Hz, 2 H, H^{mB}); 7.55 (d, J = 8.8 Hz, 2 H, H^{oB}); 10.25 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, $\text{DMSO}-d_6$): δ = 12.2 (CH_2CH_3); 20.4 (CH_3); 37.4, 38.7 (CH_2); 43.3 (SCH); 120.6, 120.8 (C^{oA} , C^{oB}); 128.7, 129.6 (C^{mA} , C^{mB}); 126.9 (C^{pB}); 133.1 (C^{pA}); 137.6 (C^{iB}); 145.7 (C^{iA}); 154.2 ($\text{C}=\text{N}$); 167.9, 173.5 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{20}\text{H}_{20}\text{ClN}_3\text{O}_2\text{S}$ $[\text{M}+\text{H}]^+$ 402.1038, found 402.1035.

2-(3-Ethyl-2-(4-methylphenyl)imino-4-oxo-1,3-thiazolidin-5-yl)-N-(4-fluorophenyl)acetamide (5be). Colorless crystals, yield 284 mg (74%), mp 160-161 °C.



^1H NMR (400 MHz, CDCl_3): δ = 1.30 (t, J = 7.0 Hz, 3 H, CH_2CH_3); 2.32 (s, 3 H, CH_3); 2.79 (dd, J = 16.2, 9.6 Hz, 1 H, H^{a}); 3.34 (dd, J = 16.2, 4.0 Hz, 1 H, H^{b}); 3.87-4.00 (m, 2 H, CH_2CH_3); 4.43 (dd, J = 9.6, 4.0 Hz, 1 H, SCH); 6.86 (d, J = 8.2 Hz, 2 H, H^{oA}); 6.95-7.01 (m, 2 H, H^{mB}); 7.12 (d, J = 8.2 Hz, 2 H, H^{mA}); 7.39-7.44 (m, 2 H, H^{oB}); 7.69 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, CDCl_3): δ = 12.6 (CH_2CH_3); 20.9 (CH_3); 38.3, 40.4 (CH_2); 43.7 (SCH); 115.7 (d, J = 22.7 Hz, C^{mB}); 120.8 (C^{oA}); 121.7 (d, J = 7.8 Hz, C^{oB}); 129.8 (C^{mA}); 133.3 (d, J = 2.9 Hz, C^{iB}); 134.3 (C^{pA}); 145.3 (C^{iA}); 153.1 ($\text{C}=\text{N}$); 159.5 (d, J = 244.2 Hz, C^{pB}); 167.1, 174.0 ($\text{C}=\text{O}$) ppm.
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HRMS (ESI), m/z : calcd for $\text{C}_{20}\text{H}_{20}\text{FN}_3\text{O}_2\text{S}$ $[\text{M}+\text{H}]^+$ 386.1333, found 386.1337.

2-(3-Ethyl-2-(4-methylphenyl)imino-4-oxo-1,3-thiazolidin-5-yl)-N-(4-nitrophenyl)acetamide (5bf). Beige crystals, yield 326 mg (79%), mp 177-179 °C.

^1H NMR (400 MHz, CDCl_3): δ = 1.31 (t, J = 7.2 Hz, 3 H, CH_2CH_3); 2.32 (s, 3 H, CH_3); 2.89 (dd, J = 16.5, 9.3 Hz, 1 H, H^{a}); 3.43 (dd, J = 16.5, 4.4 Hz, 1 H, H^{b}); 3.88-4.02 (m, 2 H, CH_2CH_3); 4.45 (dd, J = 9.3, 4.4 Hz, 1 H, SCH); 6.86 (d, J = 8.1 Hz, 2 H, H^{oA}); 7.13 (d, J = 8.1 Hz, 2 H, H^{mA}); 7.69 (d, J = 8.8 Hz, 2 H, H^{oB}); 8.18 (d, J = 8.8 Hz, 2 H, H^{mB}); 8.52 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, CDCl_3): δ = 12.6 (CH_2CH_3); 20.9 (CH_3); 38.5, 40.8 (CH_2); 43.3 (SCH); 119.2, 120.7 (C^{oA} , C^{oB}); 125.1 (C^{mB}); 129.9 (C^{mA}); 134.4 (C^{pA}); 143.3, 143.7 ($\text{C}^{\text{i,pB}}$); 145.1 (C^{iA}); 152.6 ($\text{C}=\text{N}$); 167.6, 174.1 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{20}\text{H}_{20}\text{N}_4\text{O}_4\text{S}$ $[\text{M}+\text{H}]^+$ 413.1278, found 413.1281.

Ethyl 4-(((3-ethyl-2-(4-methylphenyl)imino-4-oxo-1,3-thiazolidin-5-yl)acetyl)amino)benzoate (5bg). Beige crystals, yield 244 mg (56%), mp 179-180 °C.

^1H NMR (400 MHz, CDCl_3): δ = 1.30 (t, J = 7.2 Hz, 3 H, NCH_2CH_3); 1.38 (t, J = 7.2 Hz, 3 H, OCH_2CH_3); 2.32 (s, 3 H, CH_3); 2.84 (dd, J = 16.5, 9.8 Hz, 1 H, H^{a}); 3.40 (dd, J = 16.5, 4.0 Hz, 1 H, H^{b}); 3.86-4.00 (m, 2 H, NCH_2CH_3); 4.35 (q, J = 7.2 Hz, 2 H, OCH_2CH_3); 4.44 (dd, J = 9.8, 4.0 Hz, 1 H, SCH); 6.86 (d, J = 8.1 Hz, 2 H, H^{oA}); 7.12 (d, J = 8.1 Hz, 2 H, H^{mA}); 7.56 (d, J = 8.5 Hz, 2 H, H^{mB}); 7.97 (d, J = 8.5 Hz, 2 H, H^{oB}); 8.20 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, CDCl_3): δ = 12.6 (OCH_2CH_3); 14.3 (NCH_2CH_3); 20.9 (CH_3); 38.3, 40.7 (COCH_2 , NCH_2); 43.6 (SCH); 60.9 (OCH_2); 118.8, 120.8 (C^{oA} , C^{mB}); 126.2 (C^{iB}); 129.8, 130.8 (C^{mA} , C^{oB}); 134.3 (C^{pA}); 141.5 (C^{pB}); 145.3 (C^{iA}); 153.0 ($\text{C}=\text{N}$); 167.1, 166.4, 174.0 (3 $\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{23}\text{H}_{25}\text{N}_3\text{O}_4\text{S}$ $[\text{M}+\text{H}]^+$ 440.1638, found 440.1633.

2-(3-Ethyl-4-oxo-2-phenylimino-1,3-thiazolidin-5-yl)-N-(4-methoxyphenyl)acetamide (5ca).

Beige crystals, yield 271 mg (71%), mp 181-183 °C.

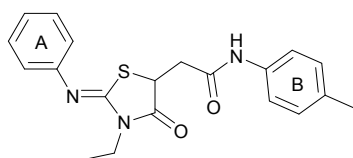
^1H NMR (400 MHz, CDCl_3): δ = 1.31 (t, J = 7.2 Hz, 3 H, CH_2CH_3); 2.77 (dd, J = 16.2, 10.0 Hz, 1 H, H^{a}); 3.43 (dd, J = 16.2, 3.8 Hz, 1 H, H^{b}); 3.76 (s, 3 H, OCH_3); 3.87-3.99 (m, 2 H, CH_2CH_3); 4.42

(dd, $J = 10.0, 3.8$ Hz, 1 H, SCH); 6.82 (d, $J = 8.8$ Hz, 2 H, H^{mB}); 6.96 (d, $J = 7.6$ Hz, 2 H, H^{oA}); 7.11 (t, $J = 7.3$ Hz, 1 H, H^{pA}); 7.30-7.36 (m, 4 H, H^{mA} , H^{oB}); 7.59 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, CDCl_3): $\delta = 12.6$ (CH_2CH_3); 38.2, 40.2 (CH_2); 43.9 (SCH); 55.4 (OCH_3); 114.1 (C^{mB}); 121.0, 121.8 (C^{oA} , C^{oB}); 124.6 (C^{pA}); 129.2 (C^{mA}); 130.3 (C^{iB}); 148.0 (C^{iA}); 153.6 ($\text{C}=\text{N}$); 156.6 (C^{pB}); 166.9, 173.9 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{20}\text{H}_{21}\text{N}_3\text{O}_3\text{S}$ $[\text{M}+\text{H}]^+$ 384.1376, found 384.1375.

2-(3-Ethyl-4-oxo-2-phenylimino-1,3-thiazolidin-5-yl)-*N*-(4-methylphenyl)acetamide (5cb).



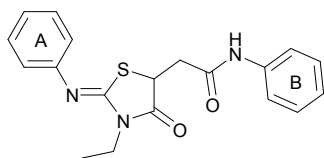
Colorless crystals, yield 287 mg (78%), mp 179-181 °C.

^1H NMR (400 MHz, CDCl_3): $\delta = 1.30$ (t, $J = 7.2$ Hz, 3 H, CH_2CH_3); 2.30 (s, 3 H, CH_3); 2.79 (dd, $J = 16.2, 10.1$ Hz, 1 H, H^a); 3.35 (dd, $J = 16.2, 3.8$ Hz, 1 H, H^b); 3.88-4.00 (m, 2 H, CH_2CH_3); 4.44 (dd, $J = 10.1, 3.8$ Hz, 1 H, SCH); 6.96 (d, $J = 7.3$ Hz, 2 H, H^{oA}); 7.01-7.13 (m, 3 H, H^{pA} , H^{mB}); 7.30-7.34 (m, 4 H, H^{mA} , H^{oB}); 7.48 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, CDCl_3): $\delta = 12.5$ (CH_2CH_3); 20.8 (CH_3); 38.2, 40.3 (CH_2); 43.9 (SCH); 120.0, 121.0 (C^{oA} , C^{oB}); 124.6 (C^{pA}); 129.2, 129.5 (C^{mA} , C^{mB}); 134.3, 134.7 ($\text{C}^{i,pB}$); 147.9 (C^{iA}); 153.7 ($\text{C}=\text{N}$); 167.0, 173.9 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $[\text{M}+\text{H}]^+$ 368.1427, found 368.1431.

2-(3-Ethyl-4-oxo-2-phenylimino-1,3-thiazolidin-5-yl)-*N*-phenylacetamide (5cc).



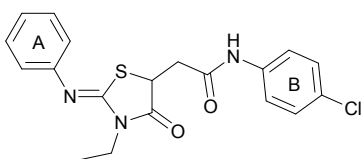
Colorless crystals, yield 243 mg (69%), mp 166-168 °C.

^1H NMR (400 MHz, $\text{DMSO}-d_6$): $\delta = 1.21$ (t, $J = 7.0$ Hz, 3 H, CH_2CH_3); 2.94 (dd, $J = 16.8, 9.8$ Hz, 1 H, H^a); 3.25 (dd, $J = 16.8, 3.5$ Hz, 1 H, H^b); 3.73-3.90 (m, 2 H, CH_2CH_3); 4.56 (dd, $J = 9.8, 3.5$ Hz, 1 H, SCH); 6.93 (d, $J = 7.6$ Hz, 2 H, H^{oA}); 7.01-7.05 (m, 1 H, H^p), 7.08-7.13 (m, 1 H, H^p); 7.26-7.36 (m, 4 H, H^{mA} , H^{mB}); 7.52 (d, $J = 7.6$ Hz, 2 H, H^{oB}); 10.11 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, $\text{DMSO}-d_6$): $\delta = 12.2$ (CH_2CH_3); 37.4, 38.7 (CH_2); 43.5 (SCH); 119.1, 121.0 (C^{oA} , C^{oB}); 123.4, 124.2 (C^{pA} , C^{pB}); 128.2, 129.2 (C^{mA} , C^{mB}); 138.7 (C^{iB}); 148.2 (C^{iA}); 154.6 ($\text{C}=\text{N}$); 167.7, 173.6 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{19}\text{H}_{19}\text{N}_3\text{O}_2\text{S}$ $[\text{M}+\text{H}]^+$ 354.1271, found 354.1270.

***N*-(4-Chlorophenyl)-2-(2-phenylimino-3-ethyl-4-oxo-1,3-thiazolidin-5-yl)acetamide (5cd).**



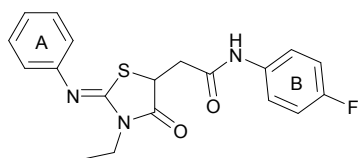
Colorless crystals, yield 283 mg (73%), mp 172-174 °C.

^1H NMR (400 MHz, CDCl_3): $\delta = 1.31$ (t, $J = 7.1$ Hz, 3 H, CH_2CH_3); 2.82 (dd, $J = 16.3, 10.0$ Hz, 1 H, H^a); 3.36 (dd, $J = 16.3, 3.7$ Hz, 1 H, H^b); 3.88-4.00 (m, 2 H, CH_2CH_3); 4.44 (dd, $J = 10.0, 3.7$ Hz, 1 H, SCH); 6.96 (d, $J = 8.6$ Hz, 2 H, H^{oA}); 7.10-7.14 (m, 1 H, H^{pA}); 7.25-7.27 (m, 2 H, H^{mB}); 7.30-7.35 (m, 2 H, H^{mA}); 7.42 (d, $J = 8.8$ Hz, 2 H, H^{oB}); 7.65 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, CDCl_3): δ = 12.6 (CH_2CH_3); 38.3, 40.4 (CH_2); 43.7 (SCH); 120.9, 121.0 (C^{oA} , C^{oB}); 124.7 (C^{pA}); 129.0, 129.2 (C^{mA} , C^{mB}); 129.7 (C^{pB}); 135.9 (C^{iB}); 147.9 (C^{iA}); 153.3 ($\text{C}=\text{N}$); 167.0, 173.9 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{19}\text{H}_{18}\text{ClN}_3\text{O}_2\text{S}$ $[\text{M}+\text{H}]^+$ 388.0881, found 388.0880.

2-(3-Ethyl-4-oxo-2-phenylimino-1,3-thiazolidin-5-yl)-N-(4-fluorophenyl)acetamide (5ce).



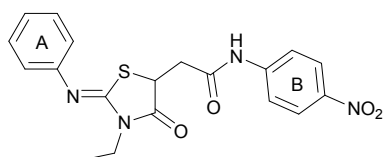
Colorless crystals, yield 285 mg (77%), mp 151-153 °C.

^1H NMR (400 MHz, CDCl_3): δ = 1.31 (t, J = 7.2 Hz, 3 H, CH_2CH_3); 2.80 (dd, J = 16.3, 9.9 Hz, 1 H, H^{a}); 3.35 (dd, J = 16.3, 3.8 Hz, 1 H, H^{b}); 3.86-4.01 (m, 2 H, CH_2CH_3); 4.43 (dd, J = 9.9, 3.8 Hz, 1 H, SCH); 6.95-7.01 (m, 4 H, H^{oA} , H^{mB}); 7.10-7.14 (m, 1 H, H^{pA}); 7.30-7.34 (m, 2 H, H^{mA}); 7.39-7.44 (m, 2 H, H^{oB}); 7.78 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, CDCl_3): δ = 12.6 (CH_2CH_3); 38.3, 40.3 (CH_2); 43.7 (SCH); 115.7 (d, J = 22.6 Hz, C^{mB}); 121.0 (C^{oA}); 121.7 (d, J = 8.1 Hz, C^{oB}); 124.7 (C^{pA}); 129.2 (C^{mA}); 133.2 (d, J = 2.2 Hz, C^{iB}); 147.9 (C^{iA}); 153.4 ($\text{C}=\text{N}$); 159.5 (d, J = 243.5 Hz, C^{pB}); 167.1, 173.9 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{19}\text{H}_{18}\text{FN}_3\text{O}_2\text{S}$ $[\text{M}+\text{H}]^+$ 372.1177, found 372.1181.

2-(3-Ethyl-4-oxo-2-phenylimino-1,3-thiazolidin-5-yl)-N-(4-nitrophenyl)acetamide (5cf).



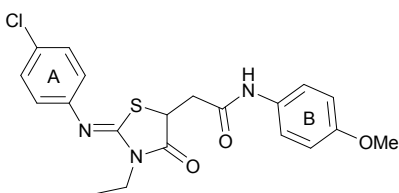
Yellow crystals, yield 311 mg (78%), mp 165-167 °C.

^1H NMR (400 MHz, $\text{DMSO}-d_6$): δ = 1.22 (t, J = 7.0 Hz, 3 H, CH_2CH_3); 3.05 (dd, J = 17.1, 9.3 Hz, 1 H, H^{a}); 3.32 (dd, J = 17.1, 3.6 Hz, 1 H, H^{b}); 3.74-3.89 (m, 2 H, CH_2CH_3); 4.58 (dd, J = 9.3, 3.6 Hz, 1 H, SCH); 6.92 (d, J = 7.6 Hz, 2 H, H^{oA}); 7.09-7.13 (m, 1 H, H^{pA}); 7.32-7.36 (m, 2 H, H^{mA}); 7.76 (d, J = 9.2 Hz, 2 H, H^{oB}); 8.20 (d, J = 9.2 Hz, 2 H, H^{mB}); 10.76 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, $\text{DMSO}-d_6$): δ = 12.3 (CH_2CH_3); 37.6, 38.9 (CH_2); 43.3 (SCH); 119.0, 120.1 (C^{oA} , C^{oB}); 124.4 (C^{pA}); 125.1 (C^{mB}); 129.4 (C^{mA}); 142.5, 144.8 ($\text{C}^{\text{i,pB}}$); 148.3 (C^{iA}); 154.6 ($\text{C}=\text{N}$); 169.0, 173.6 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{19}\text{H}_{18}\text{N}_4\text{O}_4\text{S}$ $[\text{M}+\text{H}]^+$ 399.1121, found 399.1121.

2-(2-(4-Chlorophenyl)imino-3-ethyl-4-oxo-1,3-thiazolidin-5-yl)-N-(4-methoxyphenyl)acetamide (5da). Colorless crystals, yield 260 mg (62%), mp 188-189 °C.

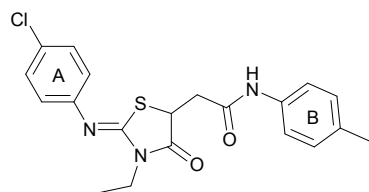


^1H NMR (300 MHz, CDCl_3): δ = 1.30 (t, J = 7.1 Hz, 3 H, CH_2CH_3); 2.80 (dd, J = 16.2, 9.9 Hz, 1 H, H^{a}); 3.34 (dd, J = 16.2, 3.8 Hz, 1 H, H^{b}); 3.78 (s, 3 H, OCH_3); 3.85-3.97 (m, 2 H, CH_2CH_3); 4.43 (dd, J = 9.9, 3.8 Hz, 1 H, SCH); 6.82 (d, J = 8.9 Hz, 2 H, H^{mB}); 6.90 (d, J = 8.6 Hz, 2 H, H^{oA}); 7.27 (d, J = 8.6 Hz, 2 H, H^{mA}); 7.35 (d, J = 8.9 Hz, 2 H, H^{oB}); 7.52 (br s, 1 H, NH) ppm.

^{13}C NMR (75 MHz, CDCl_3): δ = 12.5 (CH_2CH_3); 38.3, 40.1 (CH_2); 44.0 (SCH); 55.4 (OCH_3); 114.2 (C^{mB}); 121.9, 122.5 (C^{oA} , C^{oB}); 129.3 (C^{mA}); 129.9, 130.2 (C^{pA} , C^{iB}); 146.5 (C^{iA}); 154.3 ($\text{C}=\text{N}$); 156.7 (C^{pB}); 166.7, 173.8 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{20}\text{H}_{20}\text{ClN}_3\text{O}_3\text{S}$ $[\text{M}+\text{H}]^+$ 418.0987, found 418.0991.

2-(2-(4-Chlorophenyl)imino-3-ethyl-4-oxo-1,3-thiazolidin-5-yl)-N-(4-methylphenyl)-acetamide (5db). Pale beige crystals, yield 282 mg (70%), mp 202-203 °C.



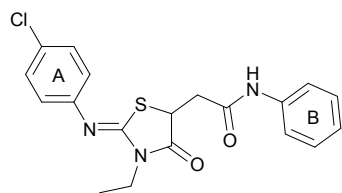
^1H NMR (400 MHz, CDCl_3): δ = 1.30 (t, J = 7.0 Hz, 3 H, CH_2CH_3); 2.30 (s, 3 H, CH_3); 2.81 (dd, J = 16.3, 9.9 Hz, 1 H, H^{a}); 3.35 (dd, J = 16.3, 3.8 Hz, 1 H, H^{b}); 3.86-3.98 (m, 2 H, CH_2CH_3); 4.44 (dd, J = 9.9, 3.8 Hz, 1 H, SCH); 6.90 (d, J = 8.7 Hz, 2 H, H^{oA}); 7.10 (d, J = 8.2

Hz, 2 H, H^{mB}); 7.27-7.34 (m, 4 H, H^{mA} , H^{oB}); 7.38 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, CDCl_3): δ = 12.5 (CH_2CH_3); 20.8 (CH_3); 38.3, 40.2 (CH_2); 43.9 (SCH); 120.0, 122.5 (C^{oA} , C^{oB}); 129.3, 129.5 (C^{mA} , C^{mB}); 129.9 (C^{pA}); 134.4, 134.6 ($\text{C}^{\text{i,pB}}$); 146.5 (C^{iA}); 154.4 ($\text{C}=\text{N}$); 166.8, 173.8 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $[\text{M}+\text{Na}]^+$ 424.0857, found 424.0855.

2-(2-(4-Chlorophenyl)imino-3-ethyl-4-oxo-1,3-thiazolidin-5-yl)-N-phenylacetamide (5dc).



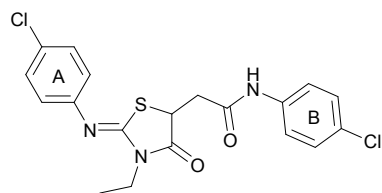
Pale yellow crystals, yield 271 mg (70%), mp 211-213 °C.

^1H NMR (400 MHz, $\text{DMSO}-d_6$): δ = 1.21 (t, J = 6.9 Hz, 3 H, CH_2CH_3); 2.95 (dd, J = 16.8, 9.4 Hz, 1 H, H^{a}); 3.26 (dd, J = 16.8, 3.4 Hz, 1 H, H^{b}); 3.74-3.89 (m, 2 H, CH_2CH_3); 4.55 (dd, J = 9.4, 3.4 Hz, 1 H, SCH); 6.92 (d, J = 7.9 Hz, 2 H, H^{oA}); 7.08-7.13 (m, 1 H, H^{pB}); 7.31-7.36 (m, 4 H, H^{mA} , H^{mB}); 7.55 (d, J = 7.9 Hz, 2 H, H^{oB}); 10.25 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, $\text{DMSO}-d_6$): δ = 12.2 (CH_2CH_3); 37.4, 39.5 (CH_2); 43.4 (SCH); 120.6, 121.0 (C^{oA} , C^{oB}); 124.1 (C^{pB}); 126.9 (C^{pA}); 128.2, 129.2 (C^{mA} , C^{mB}); 137.6 (C^{iB}); 148.2 (C^{iA}); 154.5 ($\text{C}=\text{N}$); 167.9, 173.5 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{19}\text{H}_{18}\text{ClN}_3\text{O}_2\text{S}$ $[\text{M}+\text{H}]^+$ 388.0881, found 388.0885.

N-(4-Chlorophenyl)-2-(2-(4-chlorophenyl)imino-3-ethyl-4-oxo-1,3-thiazolidin-5-yl)-acetamide (5dd). Colorless crystals, yield 340 mg (81%), mp 166-167 °C.



^1H NMR (400 MHz, $\text{DMSO}-d_6$): δ = 1.20 (t, J = 7.0 Hz, 3 H, CH_2CH_3); 2.97 (dd, J = 16.8, 9.4 Hz, 1 H, H^{a}); 3.25 (dd, J = 16.8, 3.7 Hz, 1 H, H^{b}); 3.72-3.88 (m, 2 H, CH_2CH_3); 4.57 (dd, J = 9.4, 3.7 Hz, 1 H, SCH); 6.95 (d, J = 8.5 Hz, 2 H, H^{oA}); 7.34 (d, J = 8.8 Hz, 2 H, H^{mB}); 7.38 (d, J = 8.5 Hz, 2 H, H^{mA}); 7.54 (d, J = 8.8 Hz, 2 H, H^{oB}); 10.26 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, DMSO- d_6): δ = 12.2 (CH_2CH_3); 37.4, 38.5 (CH_2); 43.5 (SCH); 120.6, 122.9 (C^{oA} , C^{oB}); 126.9, 128.2 (C^{pA} , C^{pB}); 128.7, 129.1 (C^{mA} , C^{mB}); 137.6 (C^{iB}); 147.1 (C^{iA}); 155.5 ($\text{C}=\text{N}$); 167.9, 173.5 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{19}\text{H}_{17}\text{Cl}_2\text{N}_3\text{O}_2\text{S}$ $[\text{M}+\text{H}]^+$ 422.0491, found 422.0496.

2-(2-(4-Chlorophenyl)imino-3-ethyl-4-oxo-1,3-thiazolidin-5-yl)-N-(4-fluorophenyl)acetamide (5de). Colorless crystals, yield 257 mg (63%), mp 200-201 °C.

^1H NMR (400 MHz, CDCl_3): δ = 1.29 (t, J = 7.0 Hz, 3 H, CH_2CH_3); 2.82 (dd, J = 16.3, 9.8 Hz, 1 H, H^{a}); 3.37 (dd, J = 16.3, 3.8 Hz, 1 H, H^{b}); 3.84-4.00 (m, 2 H, CH_2CH_3); 4.43 (dd, J = 9.8, 3.8 Hz, 1 H, SCH); 6.88-6.92 (m, 2 H, H^{oA}); 6.97-7.01 (m, 2 H, H^{mB}); 7.27-7.30 (m, 2 H, H^{mA}); 7.39-7.44 (m, 2 H, H^{oB}); 7.70 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, CDCl_3): δ = 12.5 (CH_2CH_3); 38.3, 40.2 (CH_2); 43.8 (SCH); 115.7 (d, J = 22.7 Hz, C^{mB}); 121.8 (d, J = 8.1 Hz, C^{oB}); 122.4 (C^{oA}); 129.3 (C^{mA}); 130.0 (C^{pA}); 133.2 (d, J = 2.2 Hz, C^{iB}); 146.5 (C^{iA}); 154.2 ($\text{C}=\text{N}$); 159.6 (d, J = 244.3 Hz, C^{pB}); 166.9, 173.8 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{19}\text{H}_{17}\text{ClFN}_3\text{O}_2\text{S}$ $[\text{M}+\text{H}]^+$ 406.0787, found 406.0783.

2-(2-(4-Chlorophenyl)imino-3-ethyl-4-oxo-1,3-thiazolidin-5-yl)-N-(4-nitrophenyl)acetamide (5df). Colorless crystals, yield 293 mg (68%), mp 173-174 °C.

^1H NMR (300 MHz, CDCl_3): δ = 1.31 (t, J = 7.1 Hz, 3 H, CH_2CH_3); 2.93 (dd, J = 16.6, 9.3 Hz, 1 H, H^{a}); 3.43 (dd, J = 16.6, 4.1 Hz, 1 H, H^{b}); 3.86-4.01 (m, 2 H, CH_2CH_3); 4.46 (dd, J = 9.3, 4.1 Hz, 1 H, SCH); 6.89 (d, J = 8.4 Hz, 2 H, H^{oA}); 7.29 (d, J = 8.4 Hz, 2 H, H^{mA}); 7.68 (d, J = 8.8 Hz, 2 H, H^{oB}); 8.19 (d, J = 8.8 Hz, 2 H, H^{mB}); 8.31 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, CDCl_3): δ = 12.5 (CH_2CH_3); 38.5, 40.6 (CH_2); 43.5 (SCH); 119.2 (C^{oB}); 122.4 (C^{oA}); 125.1 (C^{mB}); 129.4 (C^{mA}); 130.1 (C^{pA}); 143.1, 143.8 ($\text{C}^{\text{i,pB}}$); 146.3 (C^{iA}); 153.7 ($\text{C}=\text{N}$); 167.5, 173.9 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{19}\text{H}_{17}\text{ClN}_4\text{O}_4\text{S}$ $[\text{M}+\text{H}]^+$ 433.0731, found 433.0732.

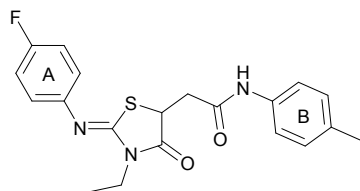
2-(3-Ethyl-2-(4-fluorophenyl)imino-4-oxo-1,3-thiazolidin-5-yl)-N-(4-methoxyphenyl)acetamide (5ea). Colorless crystals, yield 290 mg (72%), mp 172-173 °C.

^1H NMR (400 MHz, DMSO- d_6): δ = 1.20 (t, J = 7.0 Hz, 3 H, CH_2CH_3); 2.90 (dd, J = 17.7, 9.6 Hz, 1 H, H^{a}); 3.22 (dd, J = 17.7, 3.4 Hz, 1 H, H^{b}); 3.74-3.85 (m, 2 H, CH_2CH_3); 4.54 (dd, J = 9.6, 3.4 Hz, 1 H, SCH); 6.86 (d, J = 9.0 Hz, 2 H, H^{mB}); 6.93-6.97 (m, 2 H, H^{oA}); 7.14-7.20 (m, 2 H, H^{mA}); 7.42 (d, J = 9.0 Hz, 2 H, H^{oB}); 9.97 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, DMSO- d_6): δ = 12.2 (CH_2CH_3); 37.4, 38.5 (CH_2); 43.6 (SCH); 55.1 (OCH_3); 113.9 (C^mB); 115.9 (d, J = 22.2 Hz, C^mA); 120.6 (C^oB); 122.6 (d, J = 8.4 Hz, C^oA); 131.8 (C^iB); 144.6 (d, J = 2.2 Hz, C^iA); 155.3 ($\text{C}=\text{N}$, C^pB); 159.0 (d, J = 239.9 Hz, C^pA); 167.1, 173.6 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{20}\text{H}_{20}\text{FN}_3\text{O}_3\text{S}$ $[\text{M}+\text{H}]^+$ 402.1282, found 402.1284.

2-(3-Ethyl-2-(4-fluorophenyl)imino-4-oxo-1,3-thiazolidin-5-yl)-*N*-(4-methylphenyl)acetamide (5eb). Colorless crystals, yield 289 mg (75%), mp 197-198 °C.



^1H NMR (400 MHz, CDCl_3): δ = 1.30 (t, J = 7.2 Hz, 3 H, CH_2CH_3); 2.30 (s, 3 H, CH_3); 2.81 (dd, J = 16.2, 10.1 Hz, 1 H, H^a); 3.36 (dd, J = 16.2, 3.8 Hz, 1 H, H^b); 3.86-3.97 (m, 2 H, CH_2CH_3); 4.43 (dd, J = 10.1, 3.8 Hz, 1 H, SCH); 6.91-6.95 (m, 2 H, H^oA); 6.98-7.03 (m, 2H, H^mA);

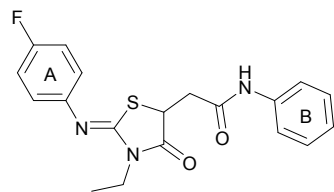
7.10 (d, J = 8.2 Hz, 2 H, H^mB); 7.33 (d, J = 8.2 Hz, 2 H, H^oB); 7.43 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, CDCl_3): δ = 12.5 (CH_2CH_3); 20.8 (CH_3); 38.3, 40.2 (CH_2); 43.9 (SCH); 115.9 (d, J = 22.0 Hz, C^mA); 120.0 (C^oB); 122.4 (d, J = 8.1 Hz, C^oA); 129.5 (C^mB); 134.4, 134.6 (C^ipB); 144.0 (d, J = 2.2 Hz, C^iA); 154.3 ($\text{C}=\text{N}$); 159.9 (d, J = 243.5 Hz, C^pA); 166.9, 173.8 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $[\text{M}+\text{H}]^+$ 386.1333, found 386.1338.

2-(3-Ethyl-2-(4-fluorophenyl)imino-4-oxo-1,3-thiazolidin-5-yl)-*N*-phenylacetamide (5ec).

Colorless crystals, yield 318 mg (86%), mp 183-185 °C.



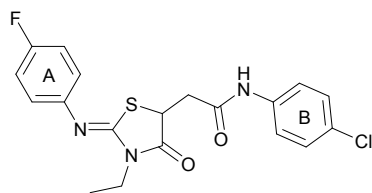
^1H NMR (400 MHz, CDCl_3): δ = 1.30 (t, J = 7.0 Hz, 3 H, CH_2CH_3); 2.82 (dd, J = 16.3, 10.0 Hz, 1 H, H^a); 3.37 (dd, J = 16.3, 3.7 Hz, 1 H, H^b); 3.82-4.02 (m, 2 H, CH_2CH_3); 4.43 (dd, J = 10.0, 3.7 Hz, 1 H, SCH); 6.90-

6.94 (m, 2 H, H^oA); 7.00-7.03 (m, 2 H, H^mA); 7.08-7.13 (m, 1 H, H^pB); 7.27-7.31 (m, 2 H, H^mB); 7.46 (d, J = 7.8 Hz, 2 H, H^oB); 7.73 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, CDCl_3): δ = 12.5 (CH_2CH_3); 38.3, 40.4 (CH_2); 43.8 (SCH); 115.9 (d, J = 22.7 Hz, C^mA); 119.9 (C^oB); 122.4 (d, J = 8.1 Hz, C^oA); 124.7 (C^pB); 129.1 (C^mB); 137.2 (C^iB); 144.0 (d, J = 2.2 Hz, C^iA); 154.2 ($\text{C}=\text{N}$); 160.0 (d, J = 242.8 Hz, C^pA); 167.0, 173.8 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{19}\text{H}_{18}\text{FN}_3\text{O}_2\text{S}$ $[\text{M}+\text{H}]^+$ 372.1177, found 372.1174.

***N*-(4-Chlorophenyl)-2-(3-ethyl-2-(4-fluorophenyl)imino-4-oxo-1,3-thiazolidin-5-yl)-acetamide (5ed).** Colorless crystals, yield 298 mg (73%), mp 190-192 °C.



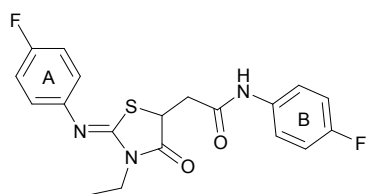
^1H NMR (400 MHz, CDCl_3): δ = 1.30 (t, J = 7.0 Hz, 3 H, CH_2CH_3); 2.82 (dd, J = 16.3, 9.7 Hz, 1 H, H^a); 3.35 (dd, J = 16.3, 4.0 Hz, 1 H, H^b); 3.85-4.01 (m, 2 H, CH_2CH_3); 4.42 (dd, J = 9.7, 4.0 Hz, 1 H, SCH); 6.90-6.94 (m, 2 H, H^oA); 6.99-7.04 (m, 2H, H^mA); 7.25 (d, J =

8.7 Hz, 2 H, H^mB); 7.42 (d, J = 8.7 Hz, 2 H, H^oB); 7.77 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, CDCl_3): δ = 12.5 (CH_2CH_3); 38.4, 40.4 (CH_2); 43.7 (SCH); 116.0 (d, J = 22.8 Hz, C^{mA}); 121.1 (C^{oB}); 122.4 (d, J = 8.1 Hz, C^{oA}); 129.1 (C^{mB}); 129.8 (C^{pB}); 135.8 (C^{iB}); 143.9 (d, J = 2.2 Hz, C^{iA}); 153.9 ($\text{C}=\text{N}$); 160.0 (d, J = 243.5 Hz, C^{pA}); 167.0, 173.8 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{19}\text{H}_{17}\text{ClFN}_3\text{O}_2\text{S}$ $[\text{M}+\text{H}]^+$ 406.0787, found 406.0791.

2-(3-Ethyl-2-(4-fluorophenyl)imino-4-oxo-1,3-thiazolidin-5-yl)-N-(4-fluorophenyl)acetamide (5ee). Colorless crystals, yield 276 mg (71%), mp 174-176 °C.



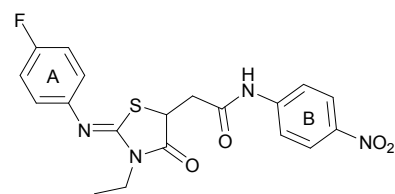
^1H NMR (300 MHz, CDCl_3): δ = 1.29 (t, J = 7.1 Hz, 3 H, CH_2CH_3); 2.82 (dd, J = 16.3, 9.9 Hz, 1 H, H^{a}); 3.36 (dd, J = 16.3, 3.9 Hz, 1 H, H^{b}); 3.84-3.99 (m, 2 H, CH_2CH_3); 4.43 (dd, J = 9.9, 3.9 Hz, 1 H, SCH); 6.89-7.05 (m, 6 H, $\text{H}^{\text{m,oA}}$, H^{mB}); 7.39-7.45 (m, 2 H, H^{oB}); 7.72

(br s, 1 H, NH) ppm.

^{13}C NMR (75 MHz, CDCl_3): δ = 12.5 (CH_2CH_3); 38.3, 40.2 (CH_2); 43.8 (SCH); 115.7 (d, J = 22.7 Hz, C^{mB}); 116.0 (d, J = 22.7 Hz, C^{mA}); 121.8 (d, J = 7.6 Hz, C^{oB}); 122.4 (d, J = 8.1 Hz, C^{oA}); 133.2 (d, J = 2.7 Hz, C^{iB}); 143.9 (d, J = 2.6 Hz, C^{iA}); 153.9 ($\text{C}=\text{N}$); 159.6 (d, J = 243.5 Hz, C^{p}); 160.0 (d, J = 244.6 Hz, C^{p}); 167.0, 173.8 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{19}\text{H}_{17}\text{F}_2\text{N}_3\text{O}_2\text{S}$ $[\text{M}+\text{H}]^+$ 390.1082, found 390.1087.

2-(3-Ethyl-2-(4-fluorophenyl)imino-4-oxo-1,3-thiazolidin-5-yl)-N-(4-nitrophenyl)-acetamide (5ef). Beige crystals, yield 309 mg (74%), mp 211-213 °C.



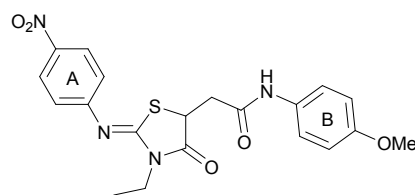
^1H NMR (400 MHz, $\text{DMSO}-d_6$): δ = 1.21 (t, J = 7.0 Hz, 3 H, CH_2CH_3); 3.06 (dd, J = 17.1, 9.3 Hz, 1 H, H^{a}); 3.33 (dd, J = 17.1, 3.7 Hz, 1 H, H^{b}); 3.73-3.88 (m, 2 H, CH_2CH_3); 4.59 (dd, J = 9.3, 3.7 Hz, 1 H, SCH); 6.94-6.96 (m, 2 H, H^{oA}); 7.14-7.20 (m, 2 H, H^{mA}); 7.77 (d,

J = 9.3 Hz, 2 H, H^{oB}); 8.20 (d, J = 9.3 Hz, 2 H, H^{mB}); 10.74 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, $\text{DMSO}-d_6$): δ = 12.2 (CH_2CH_3); 37.5, 38.8 (CH_2); 43.2 (SCH); 115.9 (d, J = 22.6 Hz, C^{mA}); 118.8 (C^{oB}); 122.6 (d, J = 8.4 Hz, C^{oA}); 125.0 (C^{mB}); 142.3, 144.7 ($\text{C}^{\text{i,pB}}$); 144.6 (d, J = 2.9 Hz, C^{iA}); 155.1 ($\text{C}=\text{N}$); 159.0 (d, J = 240.6 Hz, C^{pA}); 168.9, 173.4 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{19}\text{H}_{17}\text{FN}_4\text{O}_4\text{S}$ $[\text{M}+\text{H}]^+$ 417.1027, found 417.1026.

2-(3-Ethyl-2-(4-nitrophenyl)imino-4-oxo-1,3-thiazolidin-5-yl)-N-(4-methoxyphenyl)-acetamide (5fa). Pale yellow crystals, yield 155 mg (36%), mp 205-206 °C.



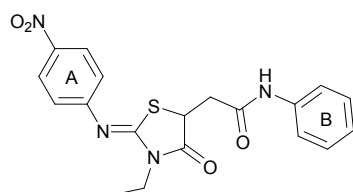
^1H NMR (400 MHz, $\text{DMSO}-d_6$): δ = 1.22 (t, J = 7.0 Hz, 3 H, CH_2CH_3); 2.95 (dd, J = 16.6, 9.6 Hz, 1 H, H^{a}); 3.24 (dd, J = 16.6, 3.7 Hz, 1 H, H^{b}); 3.70 (s, 3 H, OCH_3); 3.76-3.84 (m, 2 H, CH_2CH_3); 4.61 (dd, J = 9.6, 3.7 Hz, 1 H, SCH); 6.85 (d, J = 9.0 Hz, 2 H, H^{mB});

7.17 (d, J = 8.8 Hz, 2 H, H^{oA}); 7.41 (d, J = 9.0 Hz, 2 H, H^{oB}); 8.23 (d, J = 8.8 Hz, 2 H, H^{mA}); 10.00 (s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, DMSO- d_6): δ = 12.2 (CH_2CH_3); 37.5, 38.2 (CH_2); 44.0 (SCH); 55.1 (OCH_3); 113.9 (C^{mB}); 120.6 (C^{oB}); 122.1 (C^{oA}); 125.2 (C^{mA}); 131.8 (C^{iB}); 143.7 (C^{pA}); 154.6, 155.3, 156.8 ($\text{C}=\text{N}$, C^{iA} , C^{pB}); 167.1, 173.6 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{20}\text{H}_{20}\text{N}_4\text{O}_5\text{S}$ $[\text{M}+\text{H}]^+$ 429.1227, found 429.1232.

2-(3-Ethyl-2-(4-nitrophenyl)imino-4-oxo-1,3-thiazolidin-5-yl)-N-phenylacetamide (5fc).



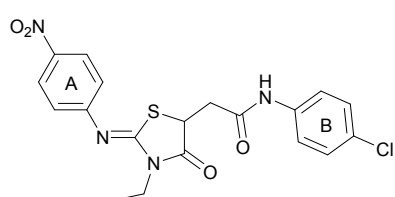
Colorless crystals, yield 248 mg (62%), mp 204-205 °C.

^1H NMR (400 MHz, DMSO- d_6): δ = 1.22 (t, J = 7.0 Hz, 3 H, CH_2CH_3); 3.00 (dd, J = 16.8, 9.6 Hz, 1 H, H^{a}); 3.28 (dd, J = 16.8, 3.5 Hz, 1 H, H^{b}); 3.75-3.90 (m, 2 H, CH_2CH_3); 4.63 (dd, J = 9.6, 3.5 Hz, 1 H, SCH); 7.04 (t, J = 7.3 Hz, 1 H, H^{pB}); 7.17 (d, J = 8.8 Hz, 2 H, H^{oA}); 7.26-7.30 (m, 2 H, H^{mB}); 7.51 (d, J = 7.9 Hz, 2 H, H^{oB}); 8.23 (d, J = 8.8 Hz, 2 H, H^{mA}); 10.13 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, DMSO- d_6): δ = 12.1 (CH_2CH_3); 37.5, 38.3 (CH_2); 43.9 (SCH); 119.1 (C^{oB}); 122.1 (C^{oA}); 123.4 (C^{pB}); 125.2 (C^{mA}); 128.7 (C^{mB}); 138.1 (C^{iB}); 143.7 (C^{pA}); 154.6, 156.7 ($\text{C}=\text{N}$, C^{iA}); 167.7, 173.5 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{19}\text{H}_{18}\text{N}_4\text{O}_4\text{S}$ $[\text{M}+\text{H}]^+$ 399.1121, found 399.1123.

N-(4-Chlorophenyl)-2-(3-ethyl-2-(4-nitrophenyl)imino-4-oxo-1,3-thiazolidin-5-yl)-acetamide (5fd). Pale yellow crystals, yield 310 mg (72%), mp 241-243 °C.

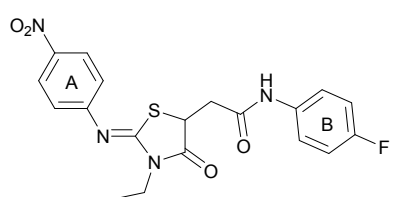


^1H NMR (400 MHz, DMSO- d_6): δ = 1.22 (t, J = 7.0 Hz, 3 H, CH_2CH_3); 3.01 (dd, J = 16.9, 9.3 Hz, 1 H, H^{a}); 3.27 (dd, J = 16.9, 3.5 Hz, 1 H, H^{b}); 3.74-3.85 (m, 2 H, CH_2CH_3); 4.63 (dd, J = 9.3, 3.5 Hz, 1 H, SCH); 7.16 (d, J = 8.8 Hz, 2 H, H^{oA}); 7.34 (d, J = 8.8 Hz, 2 H, H^{mB}); 7.54 (d, J = 8.8 Hz, 2 H, H^{oB}); 8.23 (d, J = 8.8 Hz, 2 H, H^{mA}); 10.28 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, DMSO- d_6): δ = 12.2 (CH_2CH_3); 37.5, 38.3 (CH_2); 43.8 (SCH); 120.6 (C^{oB}); 122.1 (C^{oA}); 125.2 (C^{mA}); 127.0 (C^{pB}); 128.7 (C^{mB}); 137.6 (C^{iB}); 143.7 (C^{pA}); 154.6, 156.7 ($\text{C}=\text{N}$, C^{iA}); 167.9, 173.5 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{19}\text{H}_{17}\text{ClN}_4\text{O}_4\text{S}$ $[\text{M}+\text{H}]^+$ 433.0731, found 433.0735.

2-(3-Ethyl-2-(4-nitrophenyl)imino-4-oxo-1,3-thiazolidin-5-yl)-N-(4-fluorophenyl)acetamide (5fe). Colorless crystals, yield 324 mg (78%), mp 227-228 °C.



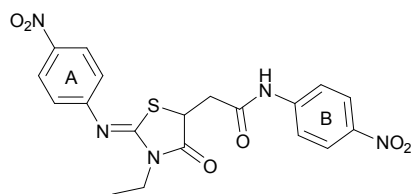
^1H NMR (400 MHz, DMSO- d_6): δ = 1.22 (t, J = 6.8 Hz, 3 H, CH_2CH_3); 3.00 (dd, J = 16.8, 9.4 Hz, 1 H, H^{a}); 3.28 (dd, J = 16.8, 3.2 Hz, 1 H, H^{b}); 3.71-3.89 (m, 2 H, CH_2CH_3); 4.62 (dd, J = 9.4, 3.2 Hz, 1 H, SCH); 7.10-7.18 (m, 4 H, H^{oA} , H^{mB}); 7.51-7.55 (m, 2 H, H^{oB}); 8.23 (d, J = 8.7 Hz, 2 H, H^{mA}); 10.20 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, DMSO- d_6): δ = 12.1 (CH_2CH_3); 37.2, 38.2 (CH_2); 43.9 (SCH); 115.1 (d, J = 22.0 Hz, $\text{C}^{m\text{B}}$); 120.9 (d, J = 8.1 Hz, $\text{C}^{o\text{B}}$); 122.1 ($\text{C}^{o\text{A}}$); 125.2 ($\text{C}^{m\text{A}}$); 135.0 (d, J = 2.3 Hz, $\text{C}^{i\text{B}}$); 143.7 ($\text{C}^{p\text{A}}$); 154.6, 156.7 ($\text{C}=\text{N}$, $\text{C}^{i\text{A}}$); 158.0 (d, J = 239.9 Hz, $\text{C}^{p\text{B}}$); 167.6, 173.5 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{19}\text{H}_{17}\text{FN}_4\text{O}_4\text{S}$ $[\text{M}+\text{H}]^+$ 417.1027, found 417.1023.

2-(3-Ethyl-2-(4-nitrophenyl)imino-4-oxo-1,3-thiazolidin-5-yl)-N-(4-nitrophenyl)acetamide

(5ff). Yellow crystals, yield 307 mg (69%), mp 234-235 °C.



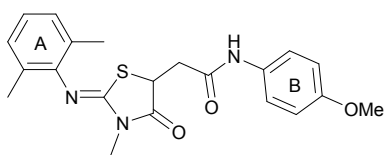
^1H NMR (400 MHz, DMSO- d_6): δ = 1.23 (t, J = 6.9 Hz, 3 H, CH_2CH_3); 3.11 (dd, J = 17.1, 9.2 Hz, 1 H, H^{a}); 3.36 (dd, J = 17.1, 3.4 Hz, 1 H, H^{b}); 3.78-3.84 (m, 2 H, CH_2CH_3); 4.66 (dd, J = 9.2, 3.4 Hz, 1 H, SCH); 7.17 (d, J = 8.8 Hz, 2 H, $\text{H}^{o\text{A}}$); 7.76 (d, J = 9.2 Hz, 2 H, $\text{H}^{o\text{B}}$); 8.19-8.25 (m, 4 H, $\text{H}^{m\text{A}}$, $\text{H}^{m\text{B}}$); 10.75 (s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, DMSO- d_6): δ = 12.1 (CH_2CH_3); 37.6, 38.5 (CH_2); 43.6 (SCH); 118.8 ($\text{C}^{o\text{B}}$); 122.1 ($\text{C}^{o\text{A}}$); 125.0, 125.2 ($\text{C}^{m\text{A}}$, $\text{C}^{m\text{B}}$); 142.3, 143.7, 144.7 ($\text{C}^{p\text{A}}$, $\text{C}^{i,p\text{B}}$); 154.5, 156.6 ($\text{C}=\text{N}$, $\text{C}^{i\text{A}}$); 168.9, 173.4 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{19}\text{H}_{17}\text{N}_5\text{O}_6\text{S}$ $[\text{M}+\text{H}]^+$ 444.0972, found 444.0971.

2-(2-(2,6-Dimethylphenyl)imino-3-methyl-4-oxo-1,3-thiazolidin-5-yl)-N-(4-methoxyphenyl)acetamide (5ga).

Beige crystals, yield 261 mg (66%), mp 192-193 °C.



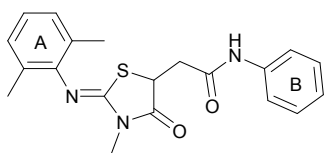
^1H NMR (400 MHz, CDCl_3): δ = 2.09 (s, 6 H, 2 CH_3); 2.83 (dd, J = 16.2, 9.2 Hz, 1 H, H^{a}); 3.25 (dd, J = 16.2, 3.8 Hz, 1 H, H^{b}); 3.39 (s, 3 H, NCH_3); 3.76 (s, 3 H, OCH_3); 4.40 (dd, J = 9.2, 3.8 Hz, 1 H, SCH); 6.81 (d, J = 8.9 Hz, 2 H, $\text{H}^{m\text{B}}$); 6.92 (t, J = 7.4 Hz, 1 H, $\text{H}^{p\text{A}}$); 7.01 (d, J = 7.4 Hz, 2 H, $\text{H}^{m\text{A}}$); 7.33 (d, J = 8.9 Hz, 2 H, $\text{H}^{o\text{B}}$); 7.52 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, CDCl_3): δ = 17.7 (2 CH_3); 29.6 (NCH_3); 40.4 (CH_2); 43.9 (SCH); 55.4 (OCH_3); 114.1 ($\text{C}^{m\text{B}}$); 121.8 ($\text{C}^{o\text{B}}$); 124.1 ($\text{C}^{p\text{A}}$); 128.09 ($\text{C}^{o\text{A}}$); 128.13 ($\text{C}^{m\text{A}}$); 130.3 ($\text{C}^{i\text{B}}$); 145.5 ($\text{C}^{i\text{A}}$); 154.1 ($\text{C}=\text{N}$); 156.6 ($\text{C}^{p\text{B}}$); 166.7, 174.2 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{21}\text{H}_{23}\text{N}_3\text{O}_3\text{S}$ $[\text{M}+\text{H}]^+$ 398.1533, found 398.1534.

2-(2-(2,6-Dimethylphenyl)imino-3-methyl-4-oxo-1,3-thiazolidin-5-yl)-N-phenylacetamide

(5gc). Pale pink crystals, yield 274 mg (75%), mp 167-168 °C.

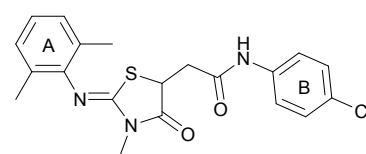


^1H NMR (400 MHz, CDCl_3): δ = 2.10 (s, 6 H, 2 CH_3); 2.86 (dd, J = 16.3, 9.3 Hz, 1 H, H^{a}); 3.28 (dd, J = 16.3, 3.8 Hz, 1 H, H^{b}); 3.40 (s, 3 H, NCH_3); 4.41 (dd, J = 9.3, 3.8 Hz, 1 H, SCH); 6.92 (t, J = 7.4 Hz, 1 H, $\text{H}^{p\text{A}}$); 7.01 (d, J = 7.4 Hz, 2 H, $\text{H}^{m\text{A}}$); 7.08-7.12 (m, 1 H, $\text{H}^{p\text{B}}$); 7.25-7.31 (m, 2 H, $\text{H}^{m\text{B}}$); 7.44 (d, J = 7.6 Hz, 2 H, $\text{H}^{o\text{B}}$); 7.65 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, CDCl_3): δ = 17.7 (2 CH_3); 29.5 (NCH_3); 40.3 (CH_2); 44.0 (SCH); 119.8 (C^{oB}); 124.2, 124.7 (C^{pA} , C^{pB}); 128.0 (C^{oA}); 128.2 (C^{mA}); 129.0 (C^{mB}); 137.2 (C^{iB}); 145.4 (C^{iA}); 154.1 ($\text{C}=\text{N}$); 166.9, 174.2 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{20}\text{H}_{21}\text{N}_3\text{O}_2\text{S}$ $[\text{M}+\text{H}]^+$ 368.1427, found 368.1426.

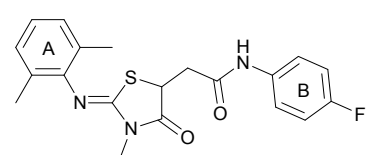
***N*-(4-Chlorophenyl)-2-(2-(2,6-dimethylphenyl)imino-3-methyl-4-oxo-1,3-thiazolidin-5-yl)acetamide (5gd).** Colorless crystals, yield 287 mg (71%), mp 193-195 °C.

 ^1H NMR (300 MHz, CDCl_3): δ = 2.09 (s, 6 H, 2 CH_3); 2.87 (dd, J = 16.3, 9.0 Hz, 1 H, H^{a}); 3.27 (dd, J = 16.3, 4.0 Hz, 1 H, H^{b}); 3.40 (s, 3 H, NCH_3); 4.41 (dd, J = 9.0, 4.0 Hz, 1 H, SCH); 6.93 (t, J = 7.4 Hz, 1 H, H^{pA}); 7.01 (d, J = 7.4 Hz, 2 H, H^{mA}); 7.25 (d, J = 8.7 Hz, 2 H, H^{mB}); 7.40 (d, J = 8.7 Hz, 2 H, H^{oB}); 7.71 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, CDCl_3): δ = 17.7 (2 CH_3); 29.6 (NCH_3); 40.4 (CH_2); 43.9 (SCH); 121.1 (C^{oB}); 124.2 (C^{pA}); 128.0 (C^{oA}); 128.2 (C^{mA}); 129.0 (C^{mB}); 129.7 (C^{pB}); 135.8 (C^{iB}); 145.4 (C^{iA}); 153.9 ($\text{C}=\text{N}$); 166.9, 174.2 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{20}\text{H}_{20}\text{ClN}_3\text{O}_2\text{S}$ $[\text{M}+\text{H}]^+$ 402.1038, found 402.1037.

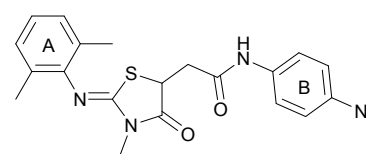
2-(2-(2,6-Dimethylphenyl)imino-3-methyl-4-oxo-1,3-thiazolidin-5-yl)-*N*-(4-fluorophenyl)acetamide (5ge). Colorless crystals, yield 253 mg (66%), mp 186-188 °C.

 ^1H NMR (300 MHz, CDCl_3): δ = 2.09 (s, 6 H, 2 CH_3); 2.86 (dd, J = 16.3, 9.3 Hz, 1 H, H^{a}); 3.28 (dd, J = 16.3, 3.7 Hz, 1 H, H^{b}); 3.40 (s, 3 H, NCH_3); 4.40 (dd, J = 9.3, 3.7 Hz, 1 H, SCH); 6.91-7.03 (m, 5 H, $\text{H}^{\text{m,pA}}$, H^{mB}); 7.37-7.43 (m, 2 H, H^{oB}); 7.42 (br s, 1 H, NH) ppm.

^{13}C NMR (75 MHz, CDCl_3): δ = 17.7 (2 CH_3); 29.6 (NCH_3); 40.2 (CH_2); 43.9 (SCH); 115.7 (d, J = 22.7 Hz, C^{mB}); 121.7 (d, J = 8.1 Hz, C^{oB}); 124.2 (C^{pA}); 128.0 (C^{oA}); 128.2 (C^{mA}); 133.2 (d, J = 2.8 Hz, C^{iB}); 145.4 (C^{iA}); 153.9 ($\text{C}=\text{N}$); 159.5 (d, J = 233.2 Hz, C^{pB}); 166.9, 174.2 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{20}\text{H}_{20}\text{FN}_3\text{O}_2\text{S}$ $[\text{M}+\text{H}]^+$ 386.1333, found 386.1338.

2-(2-(2,6-Dimethylphenyl)imino-3-methyl-4-oxo-1,3-thiazolidin-5-yl)-*N*-(4-nitrophenyl)acetamide (5gf). Pale yellow crystals, yield 333 mg (81%), mp 204-206 °C.

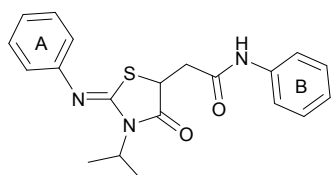
 ^1H NMR (400 MHz, CDCl_3): δ = 2.09 (s, 6 H, 2 CH_3); 2.96 (dd, J = 16.6, 8.6 Hz, 1 H, H^{a}); 3.35 (dd, J = 16.6, 4.2 Hz, 1 H, H^{b}); 3.42 (s, 3 H, NCH_3); 4.45 (dd, J = 8.6, 4.2 Hz, 1 H, SCH); 6.94 (t, J = 7.4 Hz, 1 H, C^{pA}); 7.01 (d, J = 7.4 Hz, 2 H, H^{mA}); 7.66 (d, J = 9.1 Hz, 2 H, H^{oB}); 8.15 (br s, 1 H, NH); 8.19 (d, J = 9.1 Hz, 2 H, H^{mB}) ppm.

^{13}C NMR (100 MHz, CDCl_3): δ = 17.7 (2 CH_3); 29.7 (NCH_3); 40.6 (CH_2); 43.6 (SCH); 119.1 (C^{oB}); 124.3 (C^{pA}); 125.1 (C^{mB}); 128.0 (C^{oA}); 128.2 (C^{mA}); 143.2, 143.7, 145.3 (C^{iA} , $\text{C}^{\text{i,pB}}$); 153.5 ($\text{C}=\text{N}$); 167.5, 174.2 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $C_{20}H_{20}N_4O_4S$ $[M+H]^+$ 413.1278, found 413.1276.

Reaction of thioureas **3h,i with *N*-phenylmaleimide (**4c**).** Stirred mixture of *N*-alkyl-*N'*-phenylthiourea **3h** or **3i** (1 mmol) and *N*-phenylmaleimide (**4c**) (173 mg, 1 mmol) in *i*-PrOH (5 mL) was refluxed for indicated time (17 h for **3h**, 28 h for **3i**). Solvent was evaporated under reduced pressure and the residue was separated by column chromatography (hexane/EtOAc 6:1) to obtain thiazolidines **6h,7h** and **6i,7i**, respectively.

2-(3-Isopropyl-4-oxo-2-phenylimino-1,3-thiazolidin-5-yl)-*N*-phenylacetamide (6h**).** Pale beige crystals, yield 93 mg (25%), mp 157-159 °C.

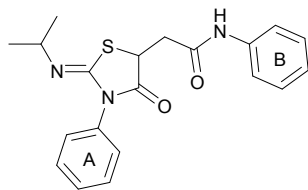


1H NMR (400 MHz, $CDCl_3$): δ = 1.53 (d, J = 7.0 Hz, 3 H, CH_3); 1.54 (d, J = 7.0 Hz, 3 H, CH_3); 2.80 (dd, J = 16.0, 9.5 Hz, 1 H, H^a); 3.32 (dd, J = 16.0, 4.1 Hz, 1 H, H^b); 4.37 (dd, J = 9.5, 4.1 Hz, 1 H, SCH); 4.90 (sept, J = 7.0 Hz, 1 H, $CH(CH_3)_2$); 6.94 (d, J = 7.3 Hz, 2 H, H^{oA}); 7.09-7.13 (m, 2 H, H^{pA} , H^{pB}); 7.28-7.34 (m, 4 H, H^{mA} , H^{mB}); 7.44-7.47 (m, 3 H, H^{oB} , NH) ppm.

^{13}C NMR (100 MHz, $CDCl_3$): δ = 18.8 (2 CH_3); 40.7 (CH_2); 43.3 (SCH); 48.0 (NCH); 119.9, 120.9 (C^{oA} , C^{oB}); 124.6, 124.7 (C^{pA} , C^{pB}); 129.0, 129.3 (C^{mA} , C^{mB}); 137.3 (C^{iB}); 148.2 (C^{iA}); 153.4 (C=N); 167.0, 174.2 (C=O) ppm.

HRMS (ESI), m/z : calcd for $C_{20}H_{21}N_3O_2S$ $[M+H]^+$ 368.1427, found 368.1428.

2-(2-Isopropylimino-4-oxo-3-phenyl-1,3-thiazolidin-5-yl)-*N*-phenylacetamide (7h**).** Colorless crystals, 142 mg (39%), mp 208-209 °C (decomp.).

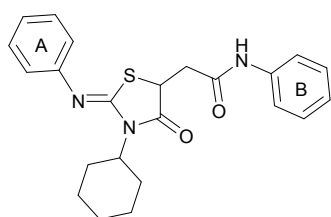


1H NMR (400 MHz, $CDCl_3$): δ = 1.09 (d, J = 6.1 Hz, 3 H, CH_3); 1.10 (d, J = 6.1 Hz, 3 H, CH_3); 2.94 (dd, J = 16.2, 9.8 Hz, 1 H, H^a); 3.45 (dd, J = 16.2, 4.0 Hz, 1 H, H^b); 3.49 (sept, J = 6.1 Hz, 1 H, $CH(CH_3)_2$); 4.60 (dd, J = 9.8, 4.0 Hz, 1 H, SCH); 7.08 (m, 2 H, H^{pB}); 7.28-7.32 (m, 4 H, H^{oA} , H^{mB}); 7.35-7.39 (m, 1 H, H^{pA}); 7.35-7.45 (m, 4 H, H^{mA} , H^{oB}); 7.72 (br s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, $DMSO-d_6$): δ = 23.38, 23.41 (2 CH_3); 39.1 (CH_2); 43.1 (SCH); 52.9 (NCH); 119.1 (C^{oB}); 123.4 (C^{pB}); 128.0 (C^{pA}); 128.4, 128.7, 129.8 ($C^{m,oA}$, C^{mB}); 135.8 (C^{iA}); 138.7 (C^{iB}); 150.4 (C=N); 167.8, 173.4 (C=O) ppm.

HRMS (ESI), m/z : calcd for $C_{20}H_{21}N_3O_2S$ $[M+H]^+$ 368.1427, found 368.1433.

2-(3-Cyclohexyl-4-oxo-2-phenylimino-1,3-thiazolidin-5-yl)-*N*-phenylacetamide (6i**).** Colorless crystals, yield 75 mg (18%), mp 180-182 °C (decomp.).

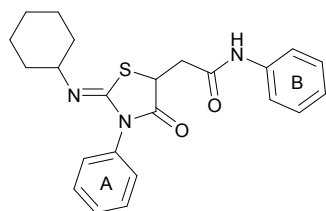


1H NMR (400 MHz, $CDCl_3$): δ = 1.18-1.42 (m, 3 H, c-Hex); 1.64-1.87 (m, 5 H, c-Hex); 2.35-2.45 (m, 2 H, c-Hex); 2.79 (dd, J = 16.2, 9.6 Hz, 1 H, H^a); 3.33 (dd, J = 16.2, 4.0 Hz, 1 H, H^b); 4.37 (dd, J = 9.6, 4.0 Hz, 1 H, SCH); 4.46-4.52 (m, 1 H, NCH); 6.93 (d, J = 7.4 Hz, 2 H, H^{oA}); 7.09-7.13 (m, 2 H, H^{pA} , H^{pB}); 7.28-7.34 (m, 4 H, H^{mA} , H^{mB}); 7.45-7.50 (m, 3 H, H^{oB} , NH) ppm.

^{13}C NMR (100 MHz, CDCl_3): δ = 25.1, 26.0, 28.1, 28.2 (c-Hex); 40.7 (CH_2); 43.2 (SCH); 56.0 (NCH); 119.9, 120.9 (C^{oA} , C^{oB}); 124.5, 124.6 (C^{pA} , C^{pB}); 129.0, 129.2 (C^{mA} , C^{mB}); 137.3 (C^{iB}); 148.3 (C^{iA}); 153.8 ($\text{C}=\text{N}$); 167.1, 174.4 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{23}\text{H}_{25}\text{N}_3\text{O}_2\text{S}$ $[\text{M}+\text{H}]^+$ 408.1740, found 408.1747.

2-(2-Cyclohexylimino-4-oxo-3-phenyl-1,3-thiazolidin-5-yl)-*N*-phenylacetamide (7i). Colorless crystals, yield 145 mg (36%), mp 206-207 °C (decomp.).



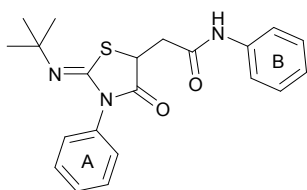
^1H NMR (400 MHz, CDCl_3): δ = 1.14-1.19 (m, 1 H, c-Hex); 1.23-1.35 (m, 4 H, c-Hex); 1.54-1.56 (m, 1 H, c-Hex); 1.64-1.70 (m, 4 H, c-Hex); 2.93 (dd, J = 16.2, 10.0 Hz, 1 H, H^{a}); 3.14-3.19 (m, 1 H, NCH); 3.45 (dd, J = 16.2, 3.8 Hz, 1 H, H^{b}); 4.60 (dd, J = 10.0, 3.8 Hz, 1 H, SCH); 7.08-7.12 (m, 1 H, H^{pB}); 7.26-7.30 (m, 4 H, H^{oA} , H^{mB}); 7.33-7.37 (m, 1 H, H^{pA}); 7.40-7.45 (m, 4 H, H^{mA} , H^{oB}); 7.80 (br s, 1 H, NH) ppm.

^1H NMR (400 MHz, $\text{DMSO}-d_6$): δ = 1.13-1.28 (m, 5 H, c-Hex); 1.53-1.66 (m, 5 H, c-Hex); 3.07-3.13 (m, 2 H, H^{a} , NCH); 3.32 (dd, J = 16.5, 3.7 Hz, 1 H, H^{b}); 4.70 (dd, J = 9.0, 3.7 Hz, 1 H, SCH); 7.04-7.08 (m, 1 H, H^{pB}); 7.30-7.34 (m, 4 H, H^{oA} , H^{mB}); 7.36-7.40 (m, 1 H, H^{pA}); 7.45-7.49 (m, 2 H, H^{mA}); 7.61 (d, J = 7.9 Hz, 2 H, H^{oB}); 10.2 (s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, CDCl_3): δ = 24.5, 25.5, 29.7, 33.1, 33.2 (c-Hex); 40.8 (CH_2); 43.6 (SCH); 61.7 (NCH); 119.8 (C^{oB}); 124.4 (C^{pB}); 128.5 (C^{pA}); 128.0, 128.9, 129.0 ($\text{C}^{\text{m,oA}}$, C^{mB}); 135.1 (C^{iA}); 137.5 (C^{iB}); 149.2 ($\text{C}=\text{N}$); 167.3, 174.1 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{23}\text{H}_{25}\text{N}_3\text{O}_2\text{S}$ $[\text{M}+\text{H}]^+$ 408.1740, found 408.1745.

2-(2-*tert*-Butylimino-4-oxo-3-phenyl-1,3-thiazolidin-5-yl)-*N*-phenylacetamide (7j). Reaction of thiourea **3j** with *N*-phenylmaleimide (**4c**). Stirred mixture of *N*-*tert*-butyl-*N'*-phenylthiourea (**3j**) (208 mg, 1 mmol) and *N*-phenylmaleimide (**4c**) (173 mg, 1 mmol) in benzene (5 mL) was refluxed for 25 h. Solvent was evaporated under reduced pressure and the residue was separated by column chromatography (hexane/EtOAc 8:1). Colorless crystals, yield 208 mg (55%), mp 207-208 °C (decomp.).

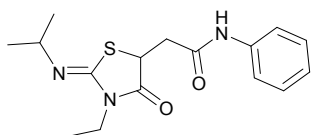


^1H NMR (400 MHz, $\text{DMSO}-d_6$): δ = 1.18 (s, 9 H, $(\text{CH}_3)_3$); 3.05 (dd, J = 16.4, 9.0 Hz, 1 H, H^{a}); 3.29 (dd, J = 16.4, 3.7 Hz, 1 H, H^{b}); 4.68 (dd, J = 9.0, 3.7 Hz, 1 H, SCH); 7.04-7.08 (m, 1 H, H^{pB}); 7.24 (d, J = 7.2 Hz, 2 H, H^{oA}); 7.29-7.38 (m, 3 H, H^{pA} , H^{mB}); 7.43-7.47 (m, 2 H, H^{mA}); 7.61 (d, J = 7.8 Hz, 2 H, H^{oB}); 10.2 (s, 1 H, NH) ppm.

^{13}C NMR (100 MHz, $\text{DMSO}-d_6$): δ = 29.2 (3 CH_3); 39.3 (CH_2); 43.9 (SCH); 53.9 ($\text{C}(\text{CH}_3)_3$); 119.2 (C^{oB}); 123.4 (C^{pB}); 127.7 (C^{pA}); 128.4, 128.6, 128.8 ($\text{C}^{\text{m,oA}}$, C^{mB}); 136.3 (C^{iA}); 138.8 (C^{iB}); 146.3 ($\text{C}=\text{N}$); 167.8, 172.5 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{21}\text{H}_{23}\text{N}_3\text{O}_2\text{S}$ $[\text{M}+\text{H}]^+$ 382.1583, found 382.1589.

2-(3-Ethyl-2-isopropylimino-4-oxo-1,3-thiazolidin-5-yl)-N-phenylacetamide (7k). Reaction of



thiourea 3k with N-phenylmaleimide (4c). Stirred mixture of *N*-ethyl-*N'*-isopropylthiourea (**3k**) (146 mg, 1 mmol) and *N*-phenylmaleimide (**4c**) (173 mg, 1 mmol) in isopropyl alcohol (5 mL) was refluxed for 15 h.

Solvent was evaporated under reduced pressure and the residue was recrystallized from ethanol/water (1:1) mixture. Colorless crystals, yield 264 mg (83%), mp 165-166 °C.

¹H NMR (400 MHz, CDCl₃): δ = 1.15-1.20 (m, 9 H, 3 CH₃); 2.78 (dd, *J* = 16.0, 9.7 Hz, 1 H, H^a); 3.39 (dd, *J* = 16.0, 4.0 Hz, 1 H, H^b); 3.45 (sept, *J* = 6.2 Hz, 1 H, CH(CH₃)₂); 3.71-3.85 (m, 2 H, NCH₂); 4.43 (dd, *J* = 9.7, 4.0 Hz, 1 H, SCH); 7.10-7.14 (m, 1 H, H^p); 7.30-7.34 (m, 2 H, H^m); 7.51 (d, *J* = 7.7 Hz, 2 H, H^o); 7.81 (br s, 1 H, NH) ppm.

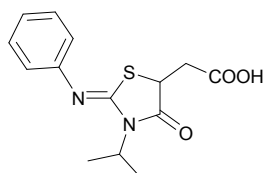
¹³C NMR (100 MHz, CDCl₃): δ = 12.4 (CH₂CH₃); 23.50, 23.54 (CH(CH₃)₂); 37.9, 41.0 (2 CH₂); 43.5 (SCH); 53.5 (CH(CH₃)₂); 119.9 (C^o); 124.6 (C^p); 129.0 (C^m); 137.5 (Cⁱ); 148.2 (C=N); 167.4, 173.8 (C=O) ppm.

HRMS (ESI), *m/z*: calcd for C₁₆H₂₁N₃O₂S [M+H]⁺ 320.1427, found 320.1418.

General procedure for the preparation of thiazolidinylacetic acids 9h,i,l. The stirred mixture of thiourea **3h,i,l** (1 mmol) and maleic anhydride (**8**) (98 mg, 1 mmol) in benzene (5 mL) was refluxed for the indicated amount of time. The solvent was evaporated under reduced pressure and the residue was recrystallized from CCl₄/*i*-PrOH mixture.

2-(3-Isopropyl-4-oxo-2-phenylimino-1,3-thiazolidin-5-yl)acetic acid (9h). During 21 h.

Colorless crystals, yield 207 mg (71%), mp 143-144 °C.



¹H NMR (400 MHz, DMSO-*d*₆): δ = 1.42 (d, *J* = 6.8 Hz, 3 H, CH₃); 1.44 (d, *J* = 6.8 Hz, 3 H, CH₃); 2.88 (dd, *J* = 17.6, 8.4 Hz, 1 H, H^a); 2.99 (dd, *J* = 17.6, 3.9 Hz, 1 H, H^b); 4.40 (dd, *J* = 8.4, 3.9 Hz, 1 H, SCH); 4.75 (sept, *J* = 6.8

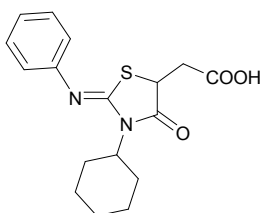
Hz, 1 H, NCH); 6.90 (d, *J* = 7.4 Hz, 2 H, H^o); 7.12 (t, *J* = 7.4 Hz, 1 H, H^p); 7.33-7.37 (m, 2 H, H^m); 12.7 (br s, 1 H, COOH) ppm.

¹³C NMR (100 MHz, DMSO-*d*₆): δ = 18.3 (CH₃); 18.5 (CH₃); 36.7 (CH₂); 42.6 (SCH); 46.9 (NCH); 120.9 (C^o); 124.1 (C^p); 129.2 (C^m); 148.4 (Cⁱ); 154.3 (C=N); 171.5 (C=O); 173.6 (C=O) ppm.

HRMS (ESI), *m/z*: calcd for C₁₄H₁₆N₂O₃S [M+H]⁺ 293.0954, found 293.0955.

2-(3-Cyclohexyl-4-oxo-2-phenylimino-1,3-thiazolidin-5-yl)acetic acid (9i). During 21 h.

Colorless crystals, yield 290 mg (87%), mp 201-202 °C.



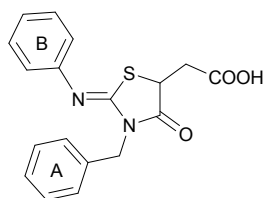
¹H NMR (400 MHz, DMSO-*d*₆): δ = 1.10-1.17 (m, 1 H, c-Hex); 1.25-1.34 (m, 2 H, c-Hex); 1.60-1.68 (m, 3 H, c-Hex); 1.79-1.82 (m, 2 H, c-Hex); 2.23-2.36 (m, 2 H, c-Hex); 2.86 (dd, *J* = 17.5, 8.4 Hz, 1 H, H^a); 2.99 (dd, *J* = 17.5, 3.6 Hz, 1 H, H^b); 4.31-4.41 (m, 2 H, SCH, NCH); 6.88 (d, *J* = 7.8 Hz, 2 H, H^o);

7.10-7.13 (m, 1 H, H^p); 7.33-7.37 (m, 2 H, H^m); 12.7 (br s, 1 H, COOH) ppm.

^{13}C NMR (100 MHz, $\text{DMSO-}d_6$): δ = 24.5 (c-Hex- CH_2); 25.6 (c-Hex- CH_2); 25.7 (c-Hex- CH_2); 27.6 (c-Hex- CH_2); 27.7 (c-Hex- CH_2); 36.8 (CH_2); 42.6 (SCH); 54.9 (NCH); 120.9 (C^o); 124.1 (C^p); 129.2 (C^m); 148.5 (C^i); 154.6 ($\text{C}=\text{N}$); 171.5 ($\text{C}=\text{O}$); 173.7 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{17}\text{H}_{20}\text{N}_2\text{O}_3\text{S}$ $[\text{M}+\text{H}]^+$ 333.1267, found 333.1268.

2-(3-Benzyl-4-oxo-2-phenylimino-1,3-thiazolidin-5-yl)acetic acid (9l). During 21 h. Colorless crystals, yield 258 mg (76%), mp 197-199 °C.

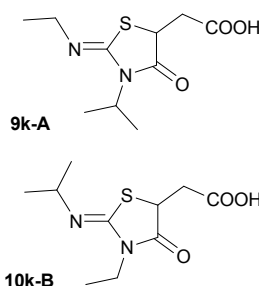


^1H NMR (400 MHz, $\text{DMSO-}d_6$): δ = 2.94 (dd, J = 17.6, 8.8 Hz, 1 H, H^a); 3.08 (dd, J = 17.6, 3.8 Hz, 1 H, H^b); 4.62 (dd, J = 8.8, 3.8 Hz, 1 H, SCH); 4.90 (d, J = 15.0 Hz, 1 H, NCH_2); 4.97 (d, J = 15.0 Hz, 1 H, NCH_2); 6.88 (d, J = 7.5 Hz, 2 H, H^{oB}); 7.11 (t, J = 7.5 Hz, 1 H, H^{pB}); 7.26-7.29 (m, 1 H, H^{pA}); 7.32-7.40 (m, 6 H, $\text{H}^{m,oA}$, H^{mB}); 12.8 (br s, 1 H, COOH) ppm.

^{13}C NMR (100 MHz, $\text{DMSO-}d_6$): δ = 36.5 (CH_2COO); 43.3 (SCH); 45.5 (NCH_2); 120.9 (C^{oB}); 124.3 (C^{pB}); 127.3 (C^{pA}); 127.5, 128.4 ($\text{C}^{m,oA}$); 129.3 (C^{mB}); 136.1 (C^{iA}); 147.9 (C^{iB}); 154.4 ($\text{C}=\text{N}$); 171.6 ($\text{C}=\text{O}$); 173.6 ($\text{C}=\text{O}$) ppm.

HRMS (ESI), m/z : calcd for $\text{C}_{18}\text{H}_{16}\text{N}_2\text{O}_3\text{S}$ $[\text{M}+\text{H}]^+$ 341.0954, found 341.0953.

Reaction of thiourea 3k with maleic anhydride (8). The stirred mixture of thiourea **3k** (146 mg, 1 mmol) and maleic anhydride (**8**) (98 mg, 1 mmol) in benzene (5 mL) was



refluxed for 15 h. The solvent was evaporated under reduced pressure and the residue was washed with a small portion of cold benzene. Mixture of two isomers **2-(2-ethylimino-3-isopropyl-4-oxo-1,3-thiazolidin-5-yl)acetic acid (9k-A)** and **2-(3-ethyl-2-isopropylimino-4-oxo-1,3-thiazolidin-5-yl)acetic acid (10k-B)** in the ratio 1:1.8 was obtained. Colorless solid, yield 220 mg (90%).

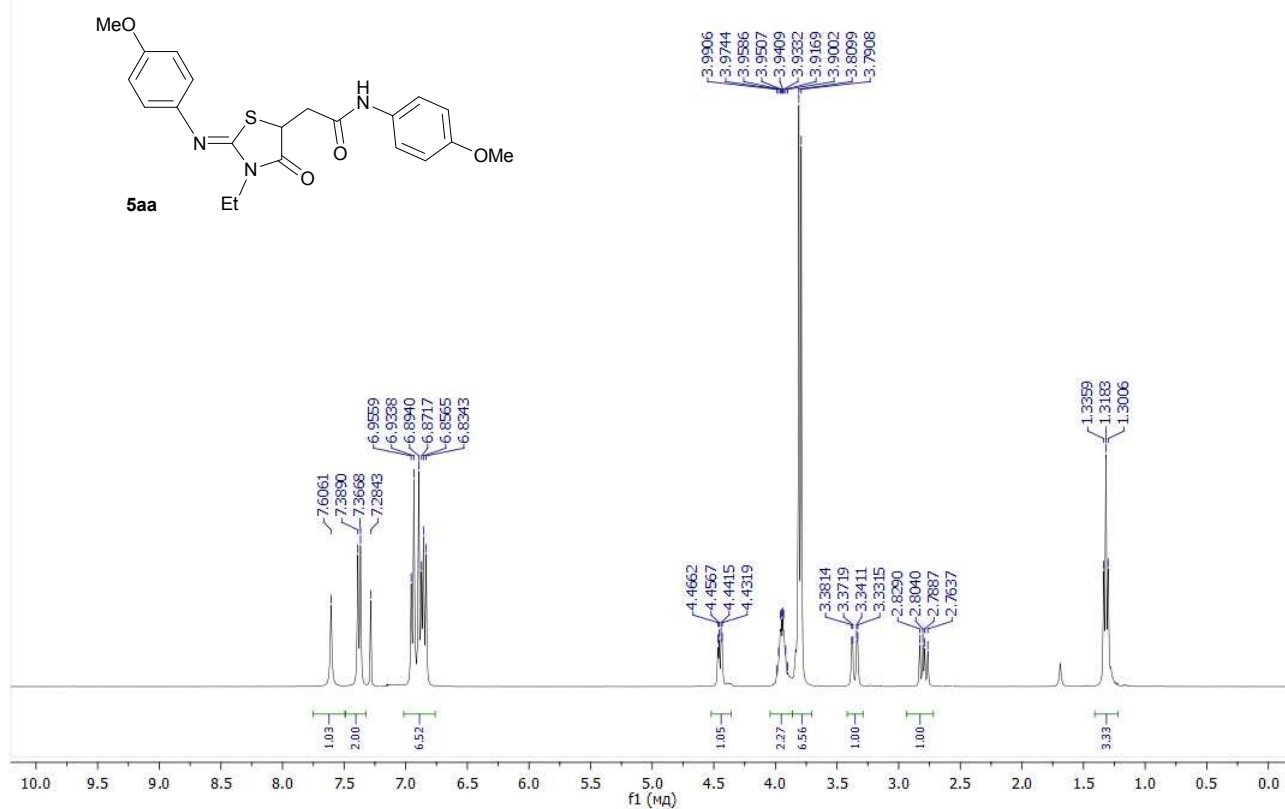
^1H NMR (400 MHz, $\text{DMSO-}d_6$) of isomer **9k-A**: δ = 1.16 (t, J = 7.2 Hz, 3 H, CH_2CH_3); 1.31 (d, J = 6.9 Hz, 3 H, $\text{CH}(\text{CH}_3)$); 1.33 (d, J = 6.9 Hz, 3 H, $\text{CH}(\text{CH}_3)$); 2.82 (dd, J = 17.5, 8.5 Hz, 1 H, H^a); 2.99 (dd, J = 17.5, 3.8 Hz, 1 H, H^b); 3.19 (q, J = 7.2 Hz, 2 H, CH_2CH_3); 4.35 (dd, J = 8.5, 3.8 Hz, 1 H, SCH); 4.62 (sept, J = 6.9 Hz, 1 H, $\text{CH}(\text{CH}_3)$); 12.7 (br s, 1 H, COOH) ppm.

^1H NMR (400 MHz, $\text{DMSO-}d_6$) of isomer **10k-B**: δ = 1.06 (t, J = 7.0 Hz, 3 H, CH_2CH_3); 1.10 (d, J = 6.2 Hz, 3 H, $\text{CH}(\text{CH}_3)$); 1.11 (d, J = 6.2 Hz, 3 H, $\text{CH}(\text{CH}_3)$); 2.83 (dd, J = 17.5, 8.9 Hz, 1 H, H^a); 3.04 (dd, J = 17.5, 3.8 Hz, 1 H, H^b); 3.38 (sept, J = 6.2 Hz, 1 H, $\text{CH}(\text{CH}_3)$); 3.54-3.67 (m, 2 H, CH_2CH_3); 4.43 (dd, J = 8.9, 3.8 Hz, 1 H, SCH); 12.7 (br s, 1 H, COOH) ppm.

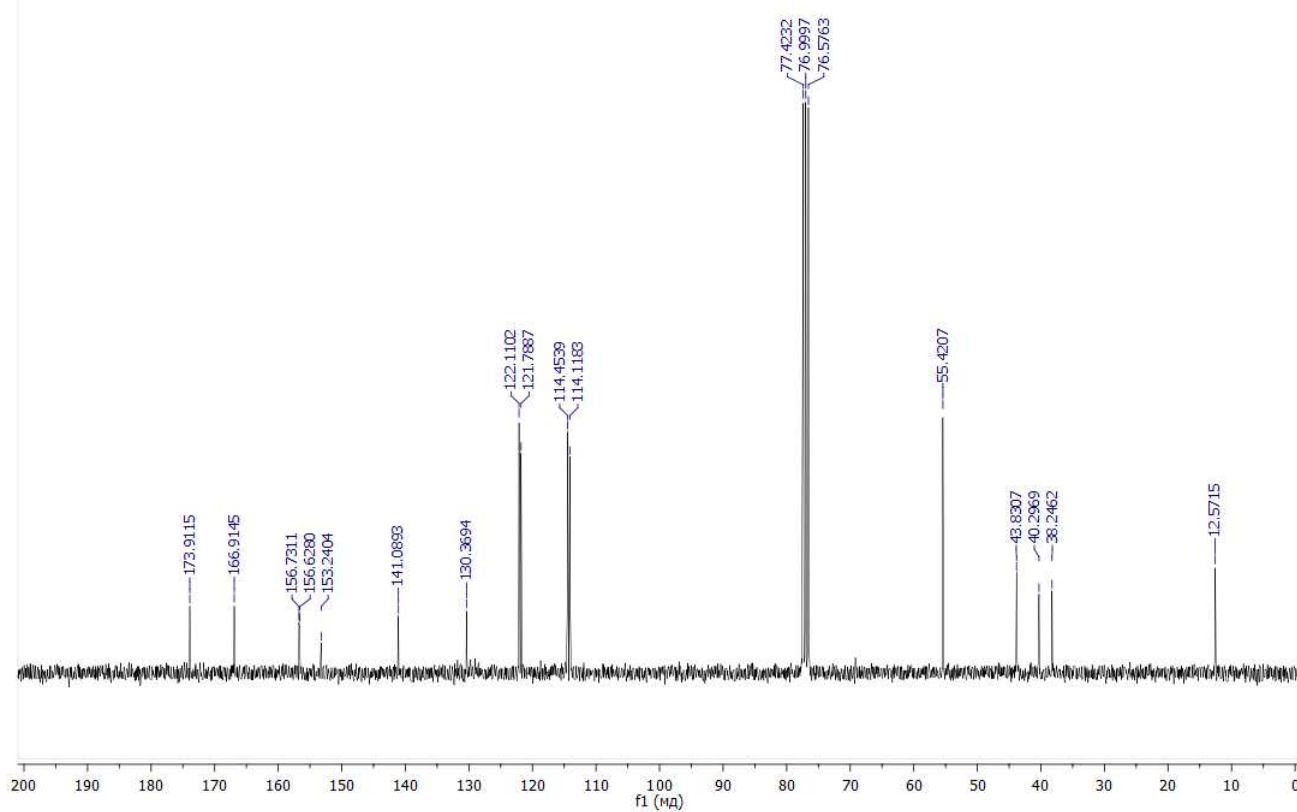
^{13}C NMR (100 MHz, $\text{DMSO-}d_6$) of isomer **9k-A**: δ = 15.8 (CH_2CH_3); 18.3, 18.4 ($\text{CH}(\text{CH}_3)_2$); 37.1 (CH_2COO); 42.4 (SCH); 46.2 (CH_2CH_3); 46.3 ($\text{CH}(\text{CH}_3)_2$); 149.0 ($\text{C}=\text{N}$); 171.5 ($\text{C}=\text{O}$); 173.1 ($\text{C}=\text{O}$) ppm.

^{13}C NMR (100 MHz, DMSO- d_6) of isomer **10k-B**: δ = 12.0 (CH_2CH_3); 23.57, 23.63 ($\text{CH}(\text{CH}_3)_2$); 37.0, 37.1 (2 CH_2); 42.9 (SCH); 52.7 ($\text{CH}(\text{CH}_3)_2$); 150.7 (C=N); 171.6 (C=O); 172.8 (C=O) ppm.
HRMS (ESI), m/z : calcd for $\text{C}_{10}\text{H}_{16}\text{N}_2\text{O}_3\text{S}$ $[\text{M}+\text{H}]^+$ 245.0954, found 245.0957.

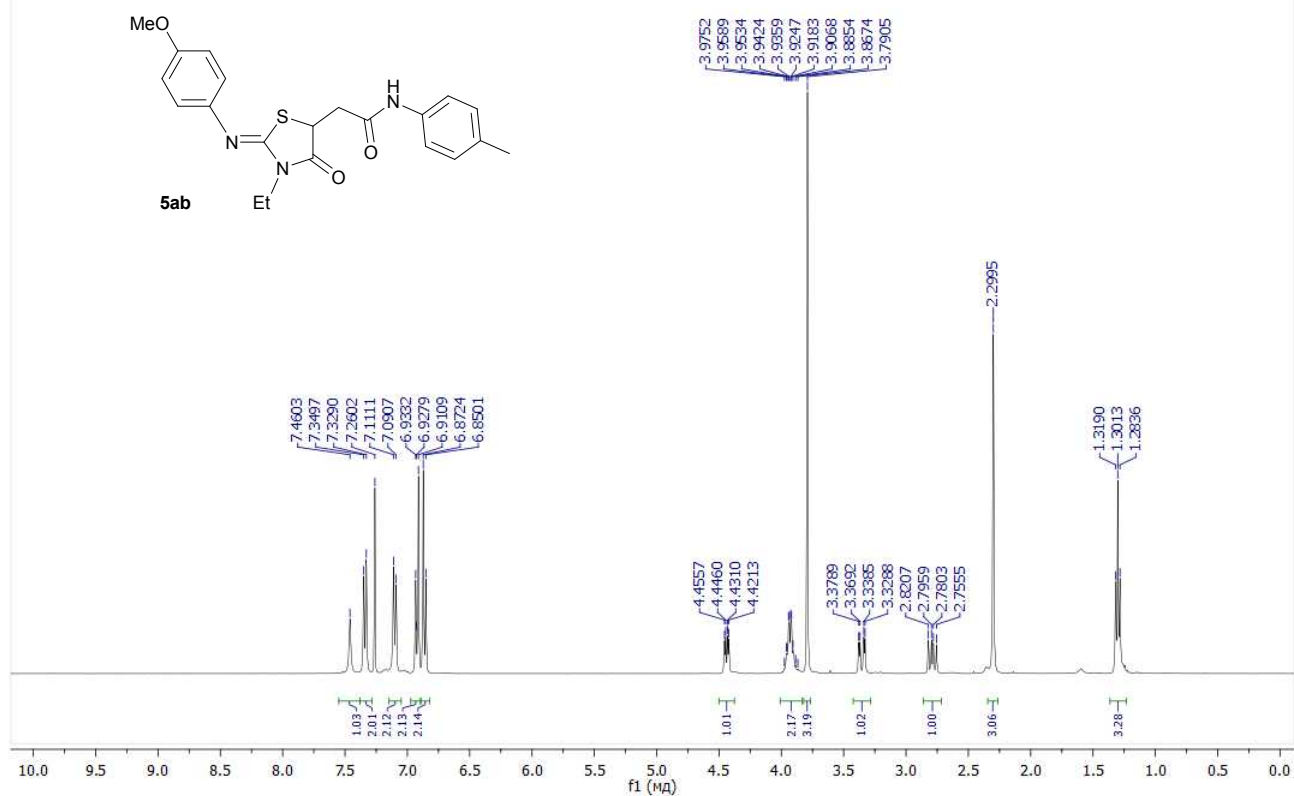
MAK, 9, BF = 400.13 MHz, Solvent - CDCl₃, 18 Jul 2013 T=298 K



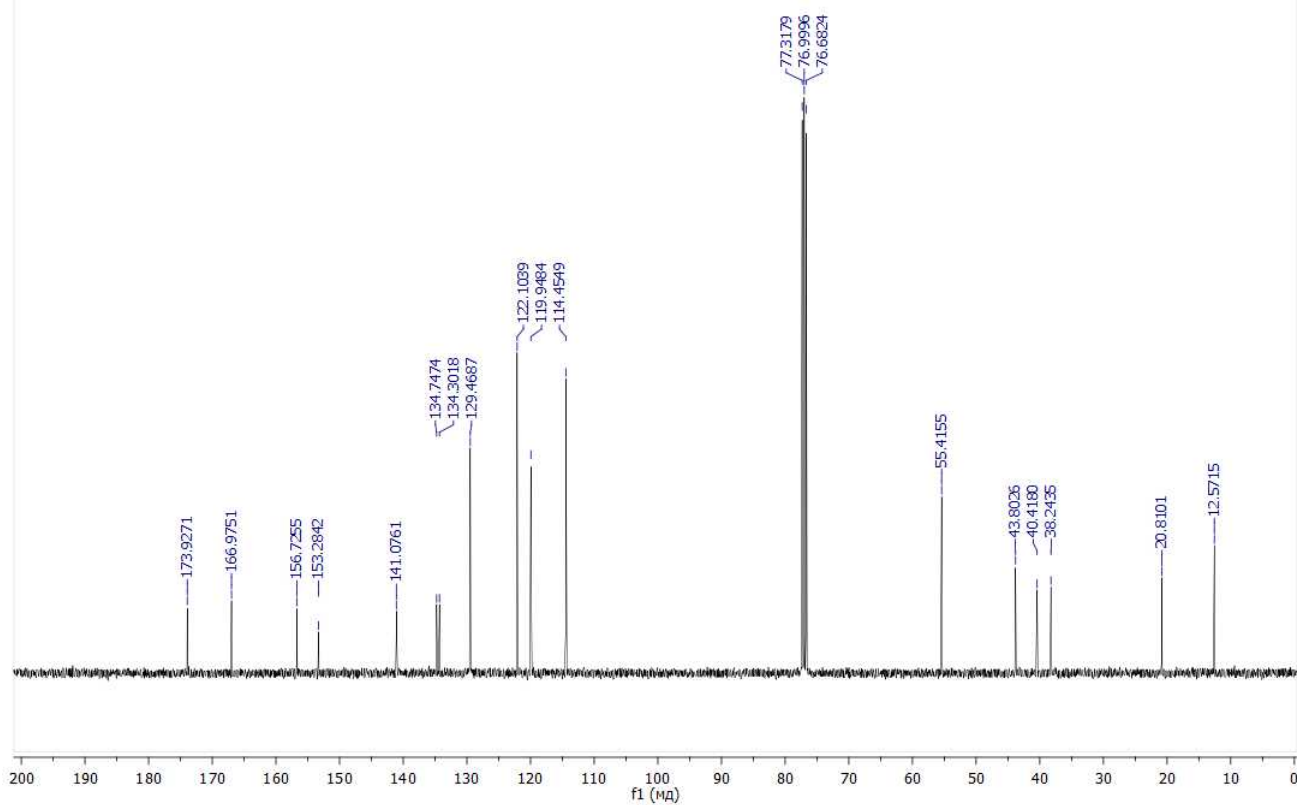
MAKc, 764, BF = 75.47 MHz, SOLVENT - CDCl₃, 3 Apr 2013



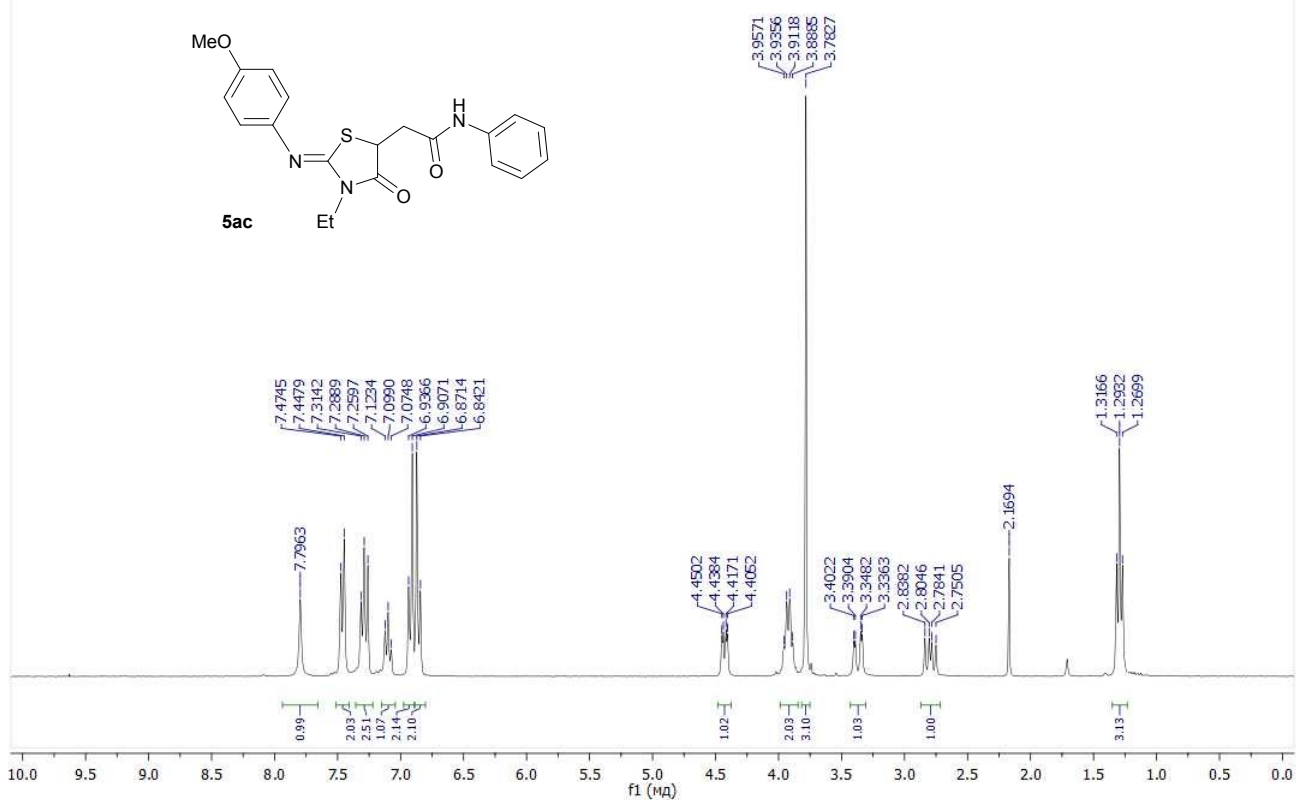
MAK, 181, BF = 400.13 MHz, Solvent - CDD3, 07 Nov 2013 T=299 K



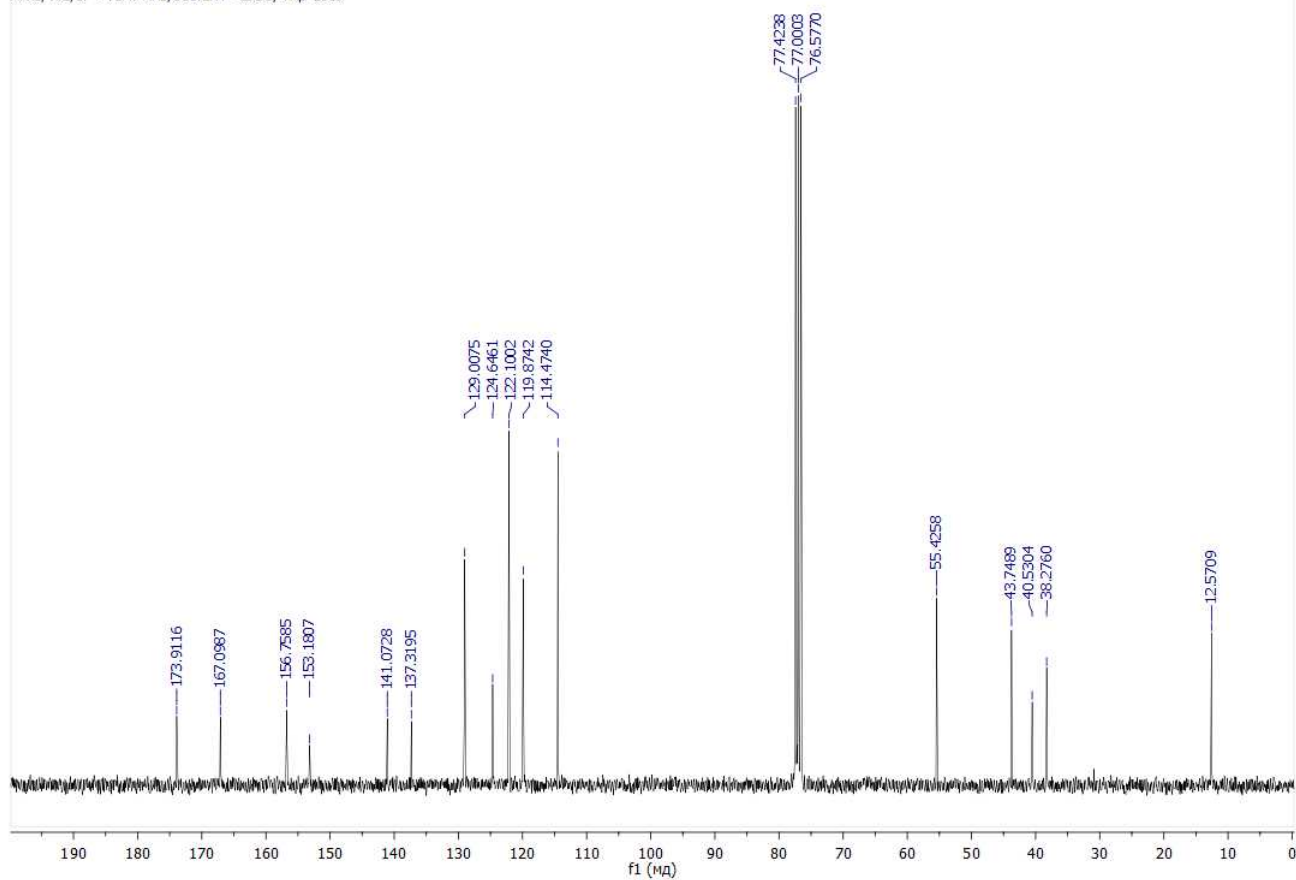
MAKc, 181, BF = 100.612769 MHz, Solvent - CDD3, 11 Nov 2013 T=298 K



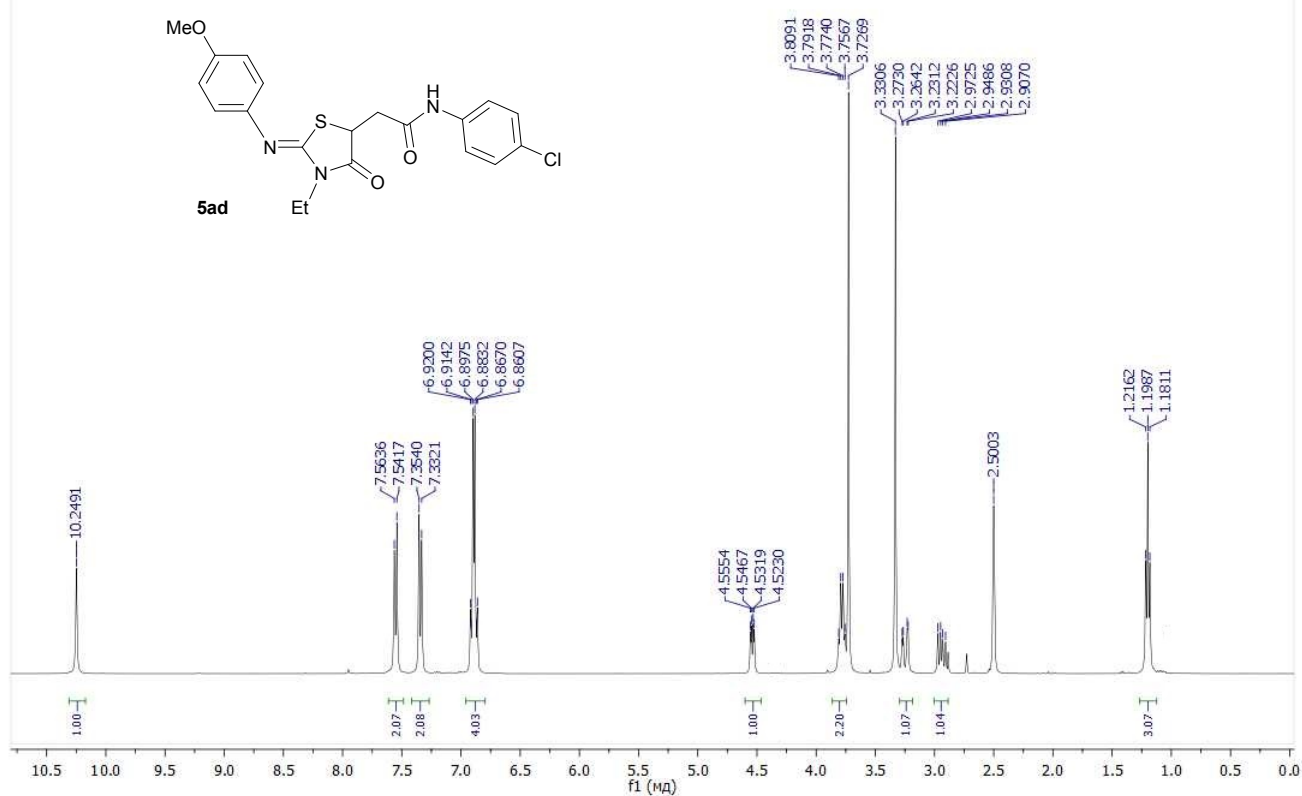
MAK, 772, BF = 300.13 MHz, SOLVENT - CDCl₃, 4 Apr 2013



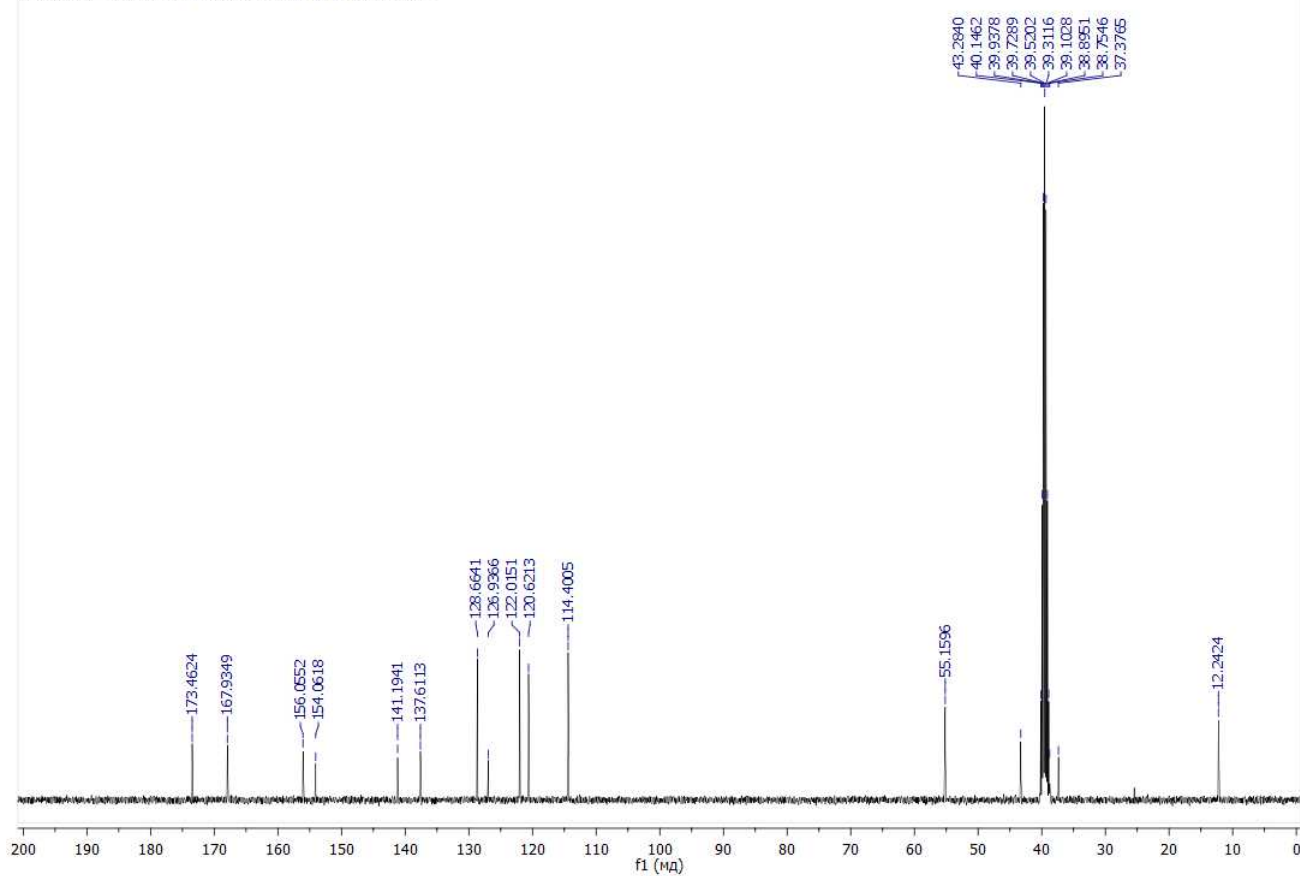
MAKc, 772, BF = 75.47 MHz, SOLVENT - CDCl₃, 4 Apr 2013



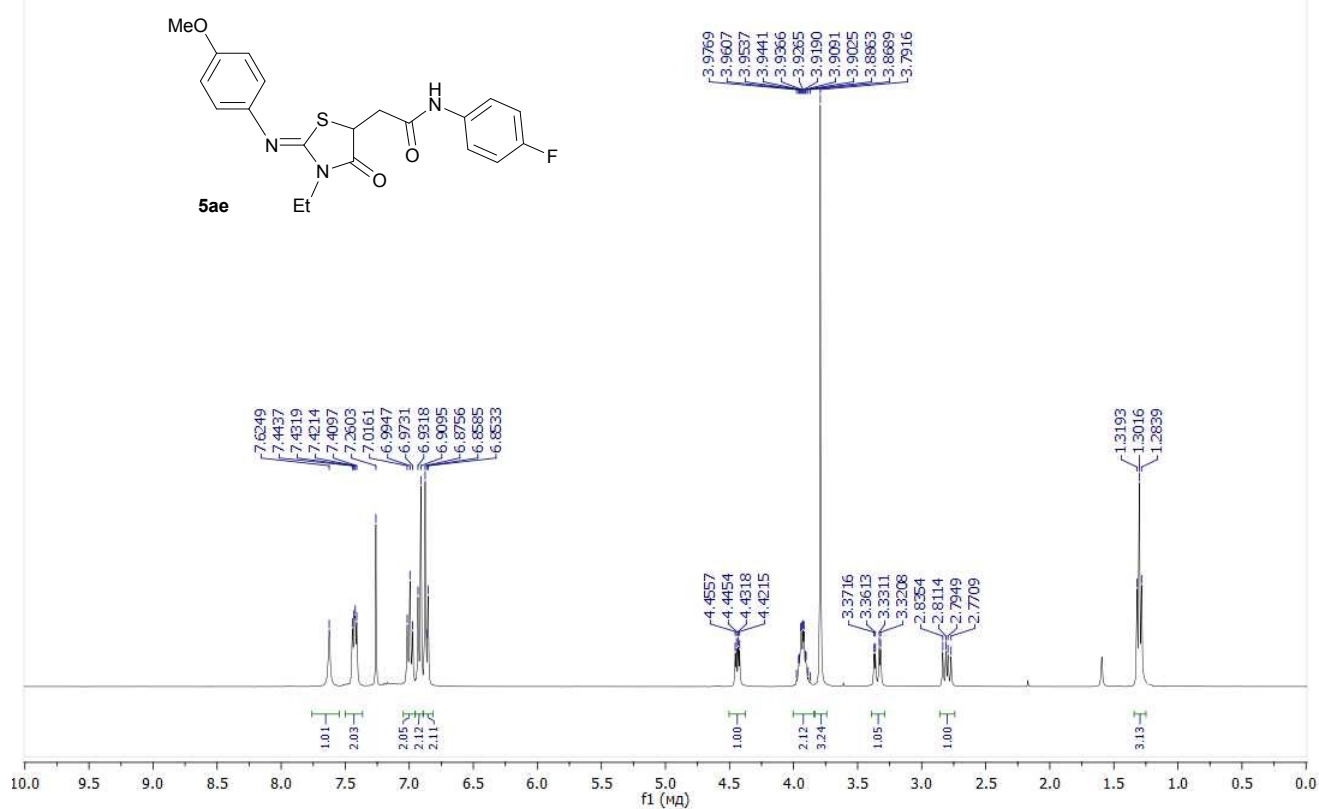
MAK, 857, BF = 400.13 MHz, Solvent - DMSO, 18 Apr 2013 T=300 K



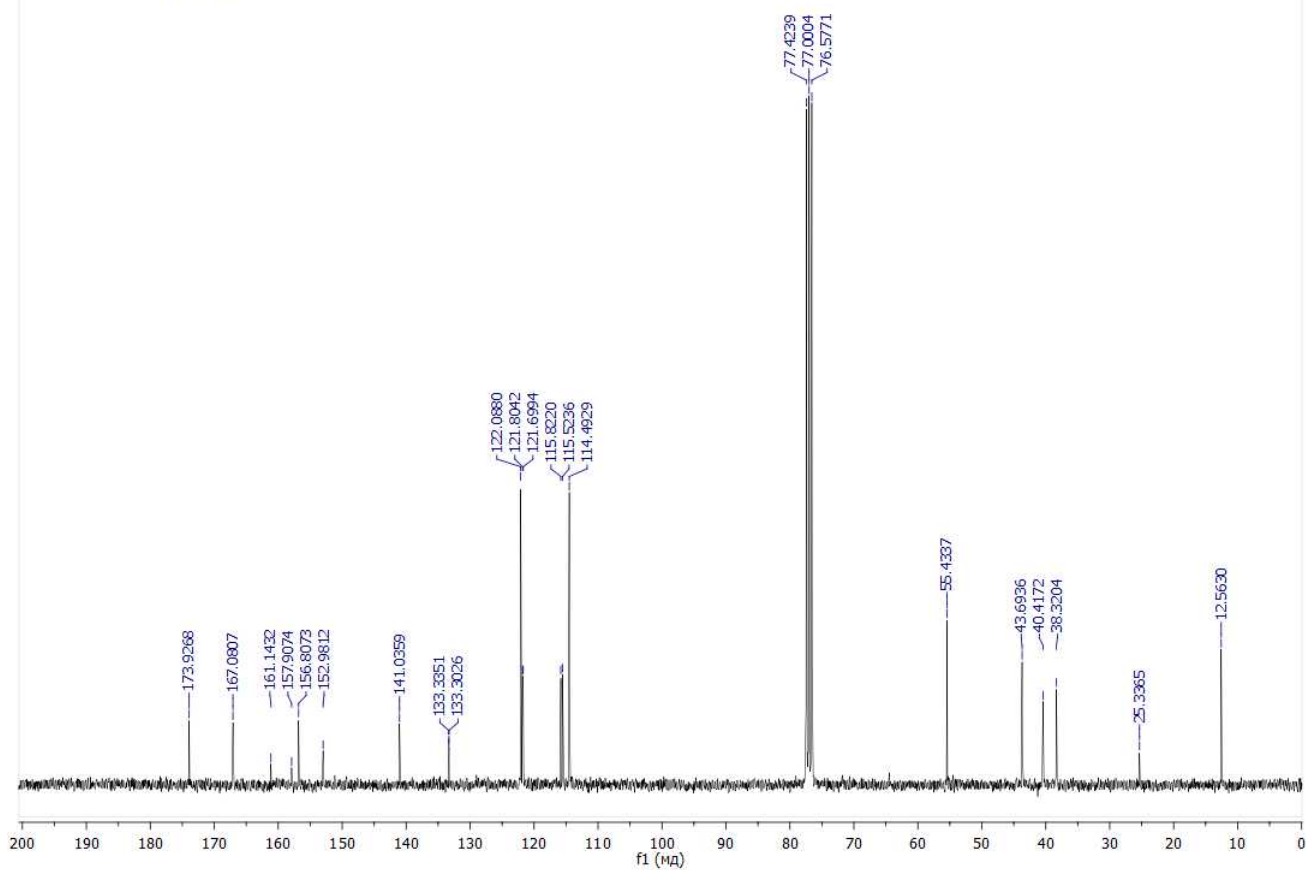
MAKc, 857, BF = 100.612769 MHz, Solvent - DMSO, 18 Apr 2013 T=300 K



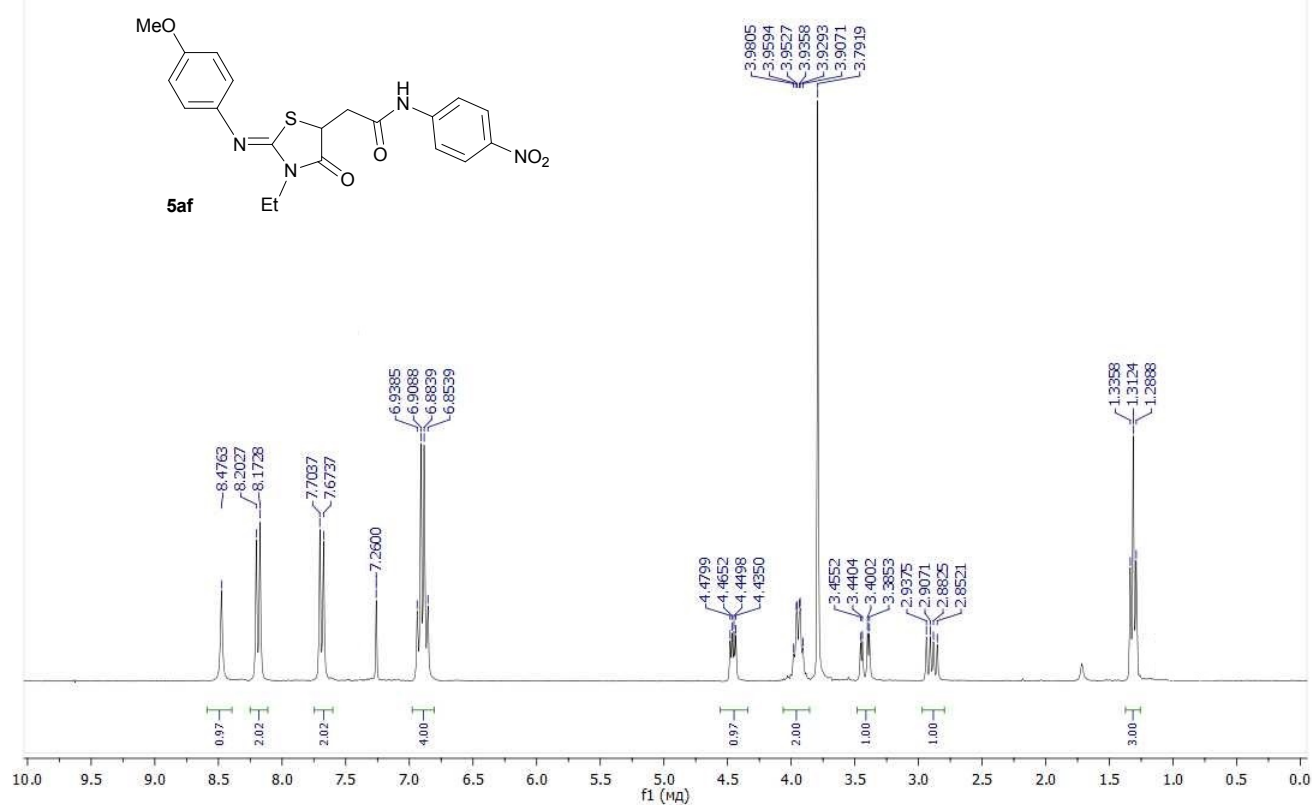
MAK, 970, BF = 400.13 MHz, Solvent - CDCl₃, 21 Jun 2013 T=300 K



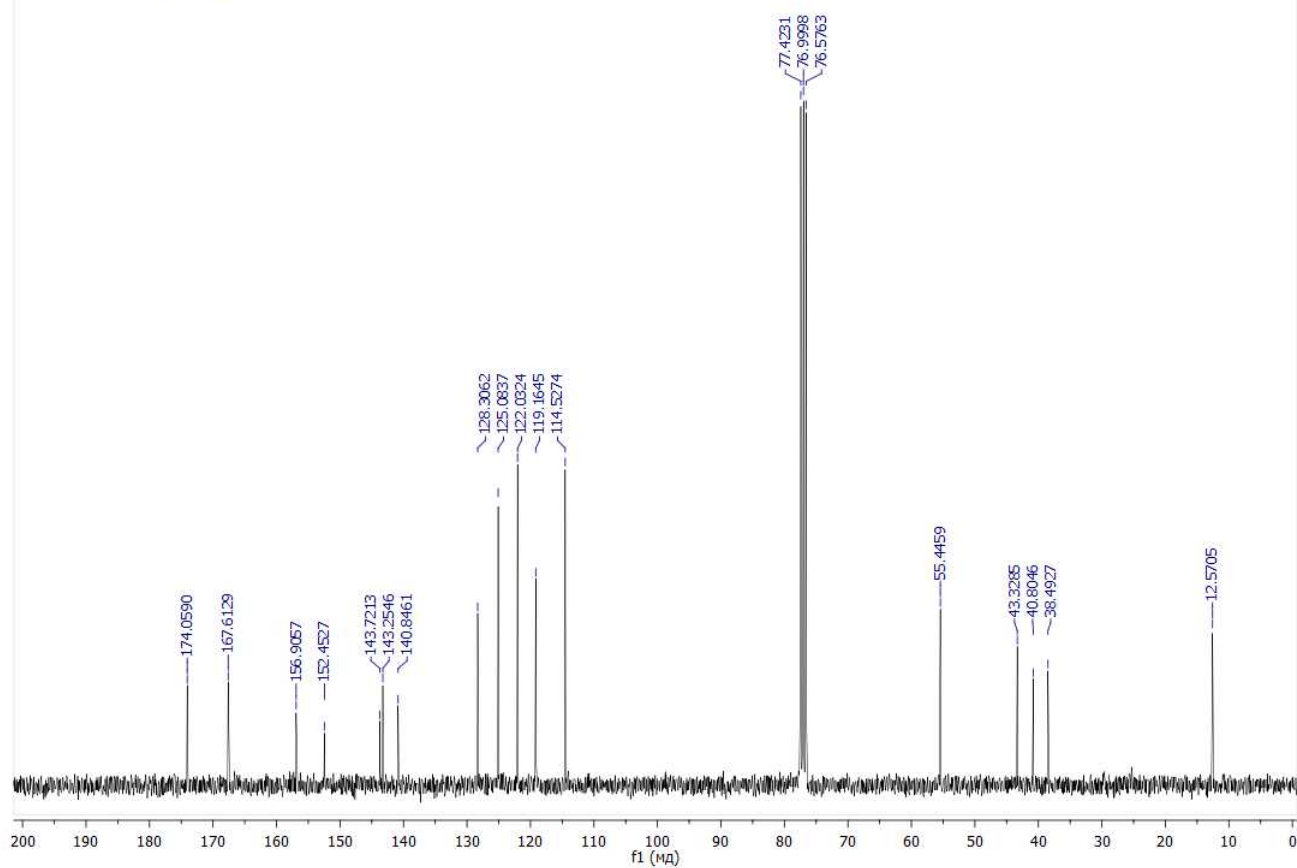
MAKc, 820, BF = 75.47 MHz, SOLVENT - CDCl₃, 11 Apr 2013



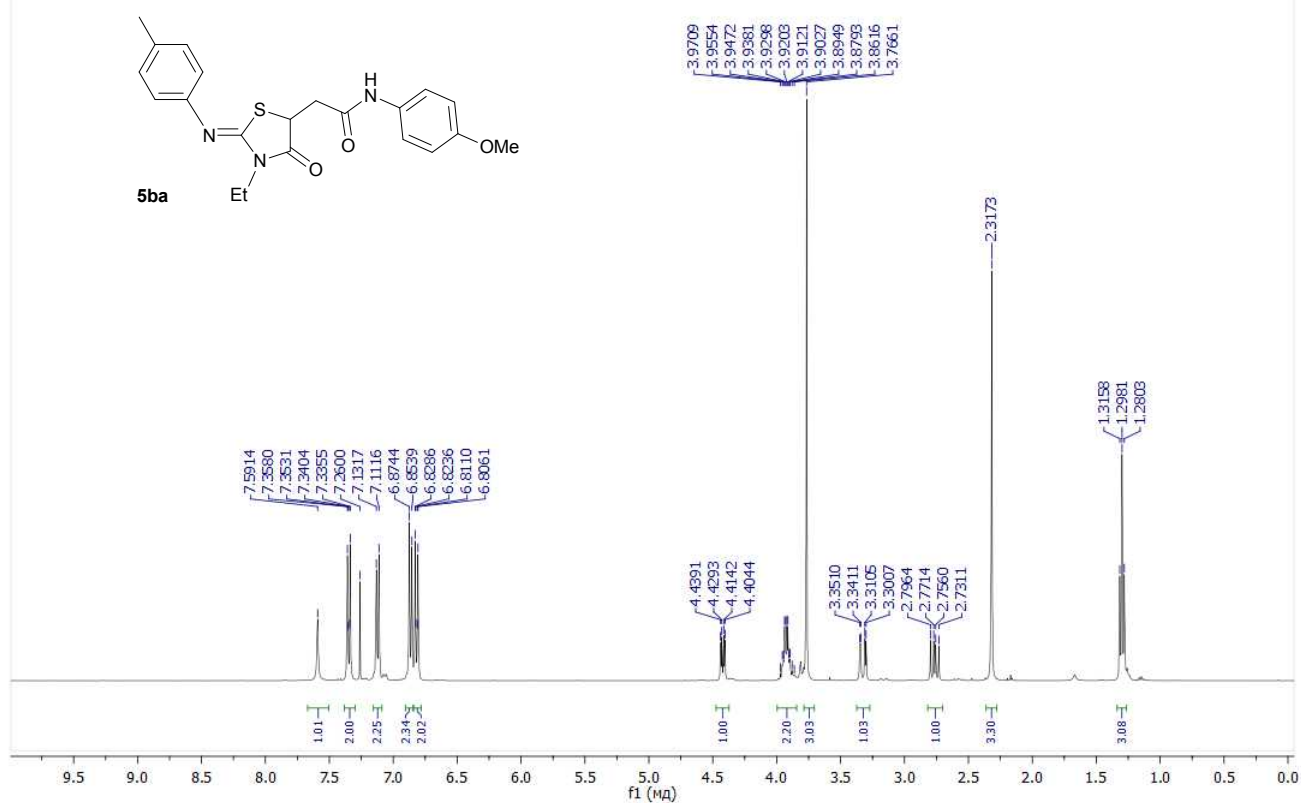
MAK, 763, BF = 300.13 MHz, SOLVENT - CDCl₃, 3 Apr 2013



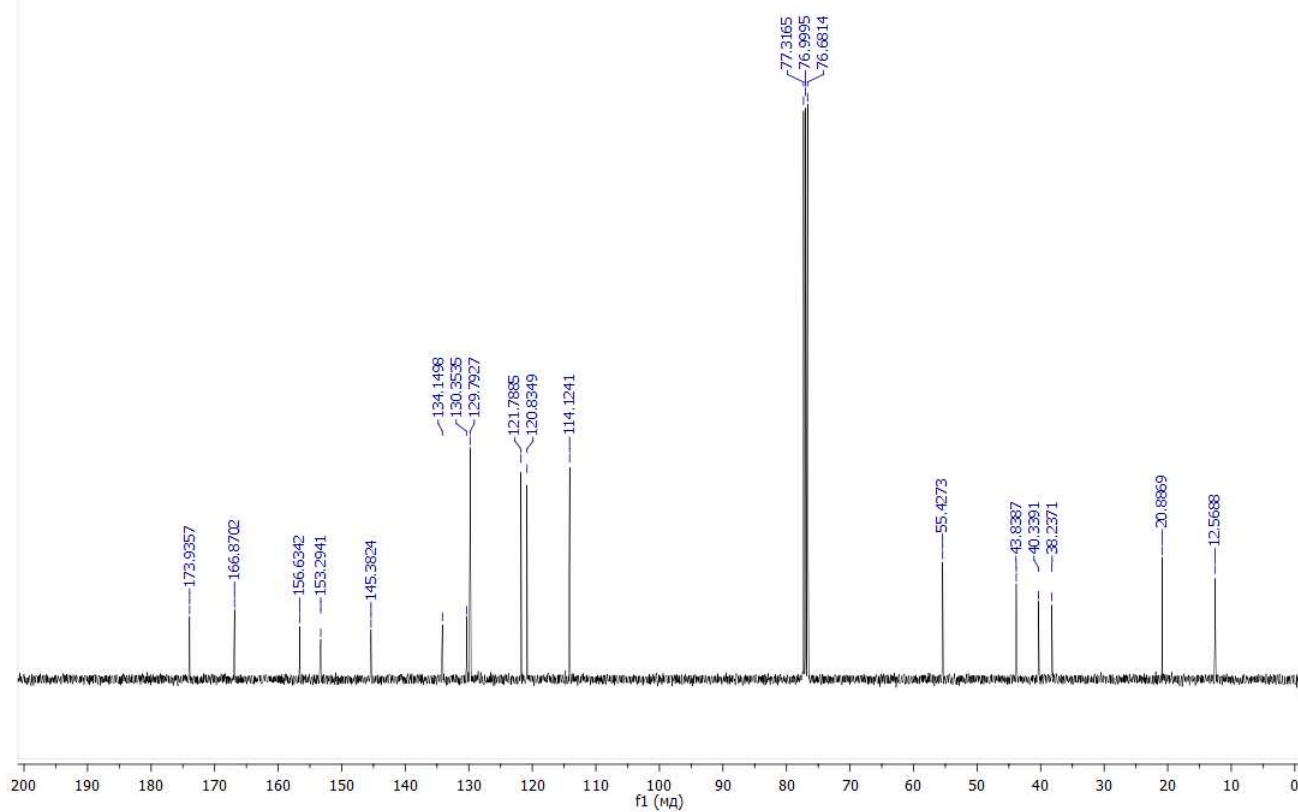
MAKc, 763, BF = 75.47 MHz, SOLVENT - CDCl₃, 3 Apr 2013



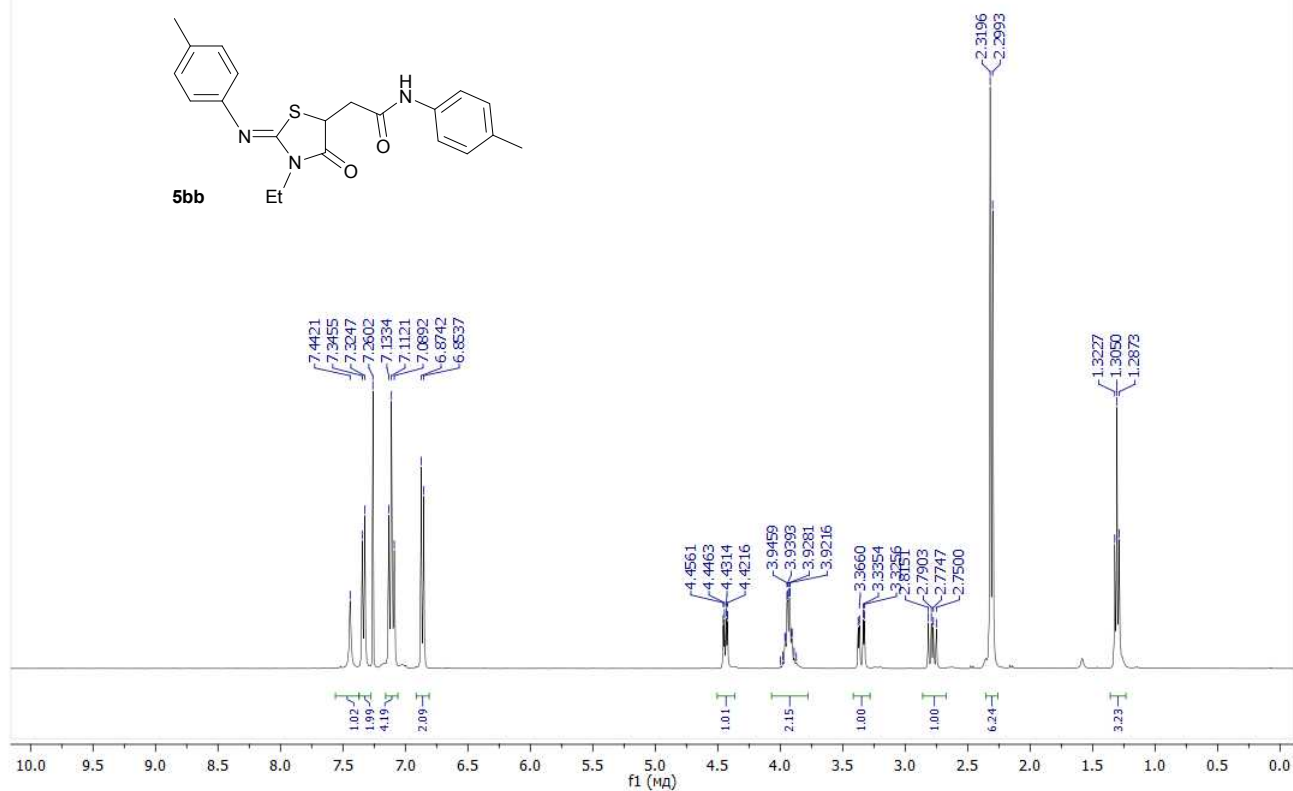
MAK, 792, BF = 400.13 MHz, Solvent - CDCl₃, 08 Apr 2013



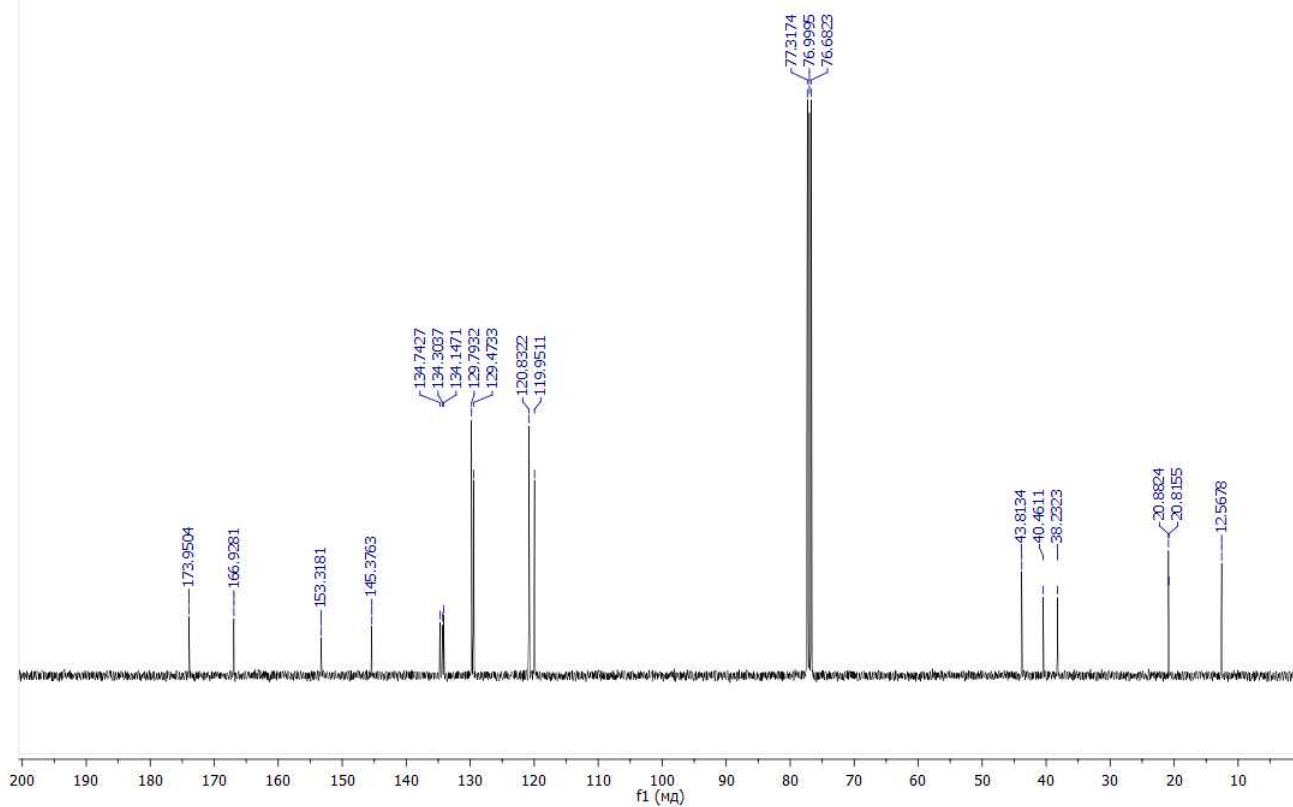
MAKc, 792, BF = 100.612769 MHz, Solvent - CDCl₃, 08 Apr 2013



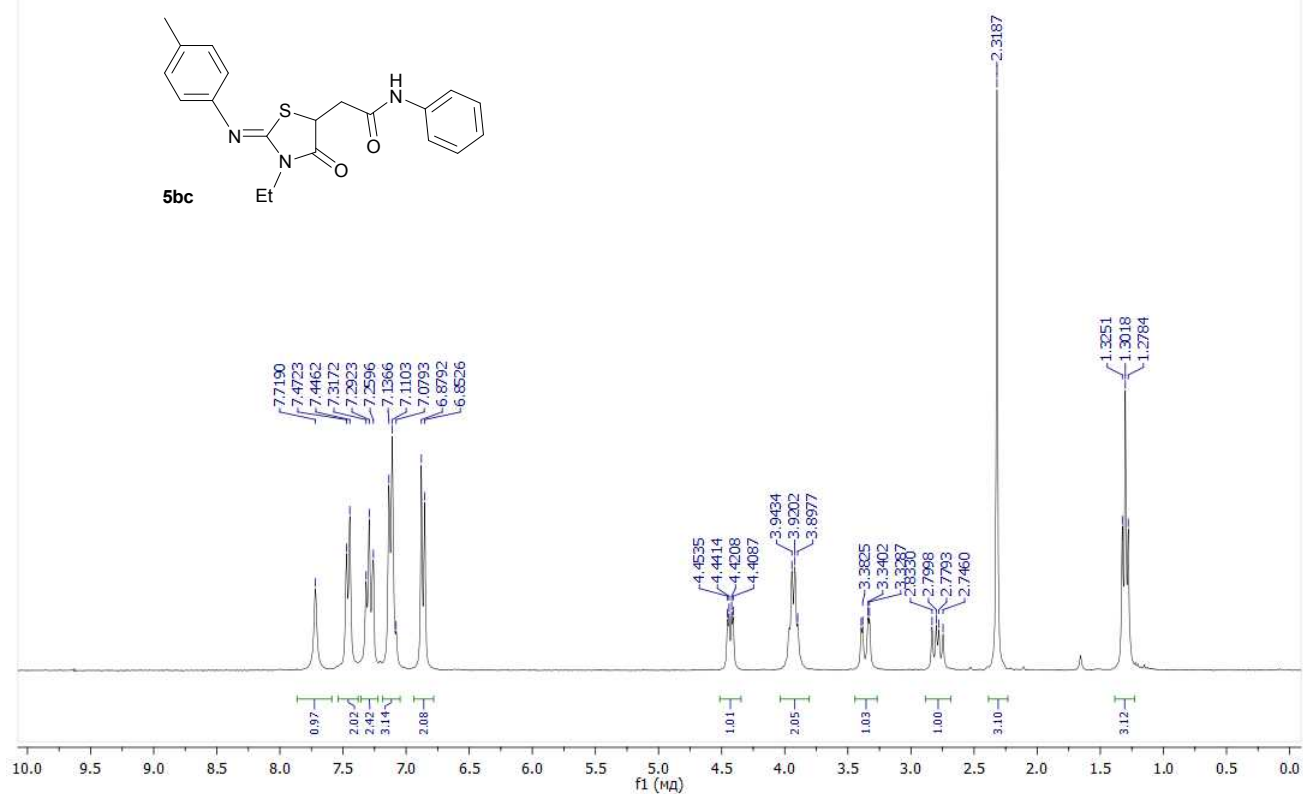
MAK, 171, BF = 400.13 MHz, Solvent - CDCl₃, 05 Nov 2013 T=298 K



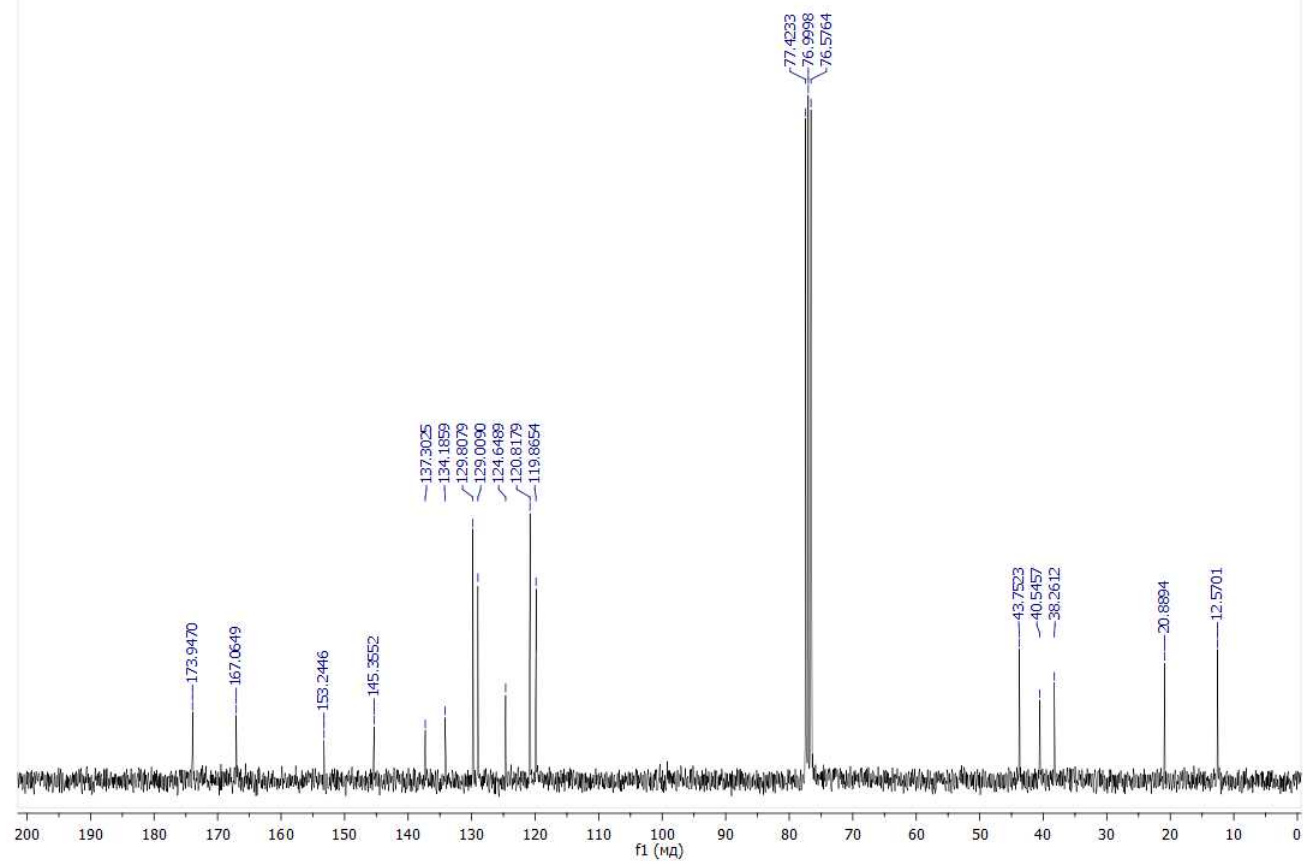
MAKc, 171, BF = 100.612769 MHz, Solvent - CDCl₃, 06 Nov 2013 T=300 K



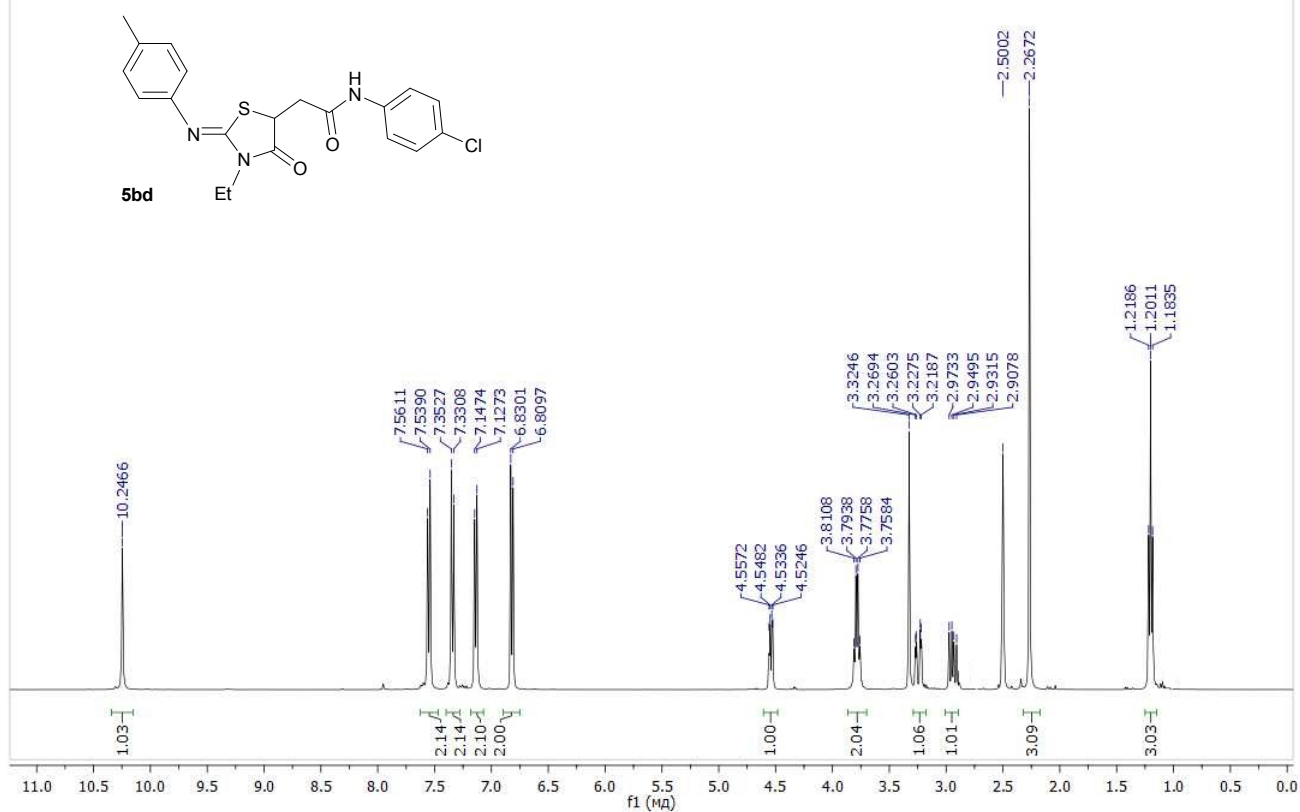
MAK, 741, BF = 300.13 MHz, SOLVENT -CDCl₃, 26 Mar 2013



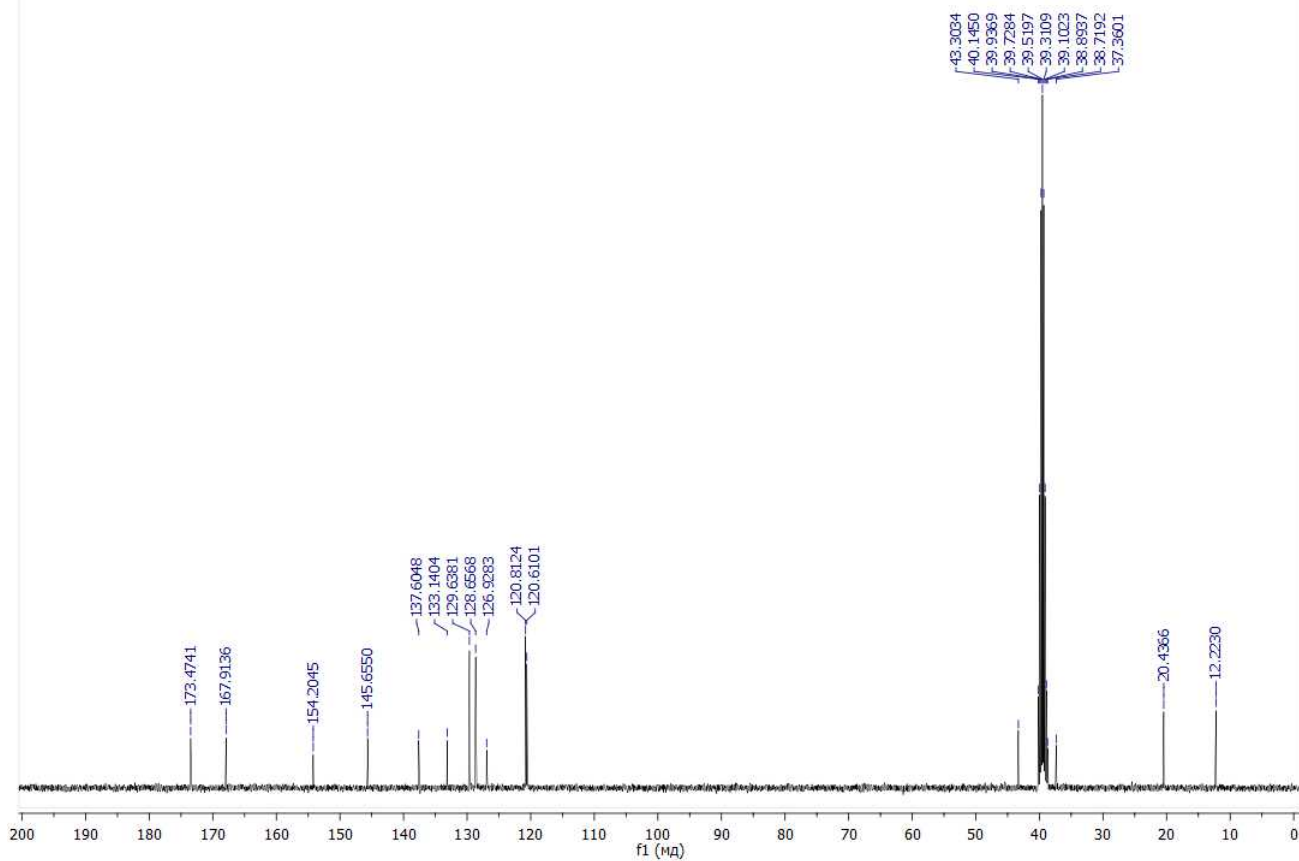
MAKc, 741, BF = 75.47 MHz, SOLVENT - CDCl₃, 26 Mar 2013



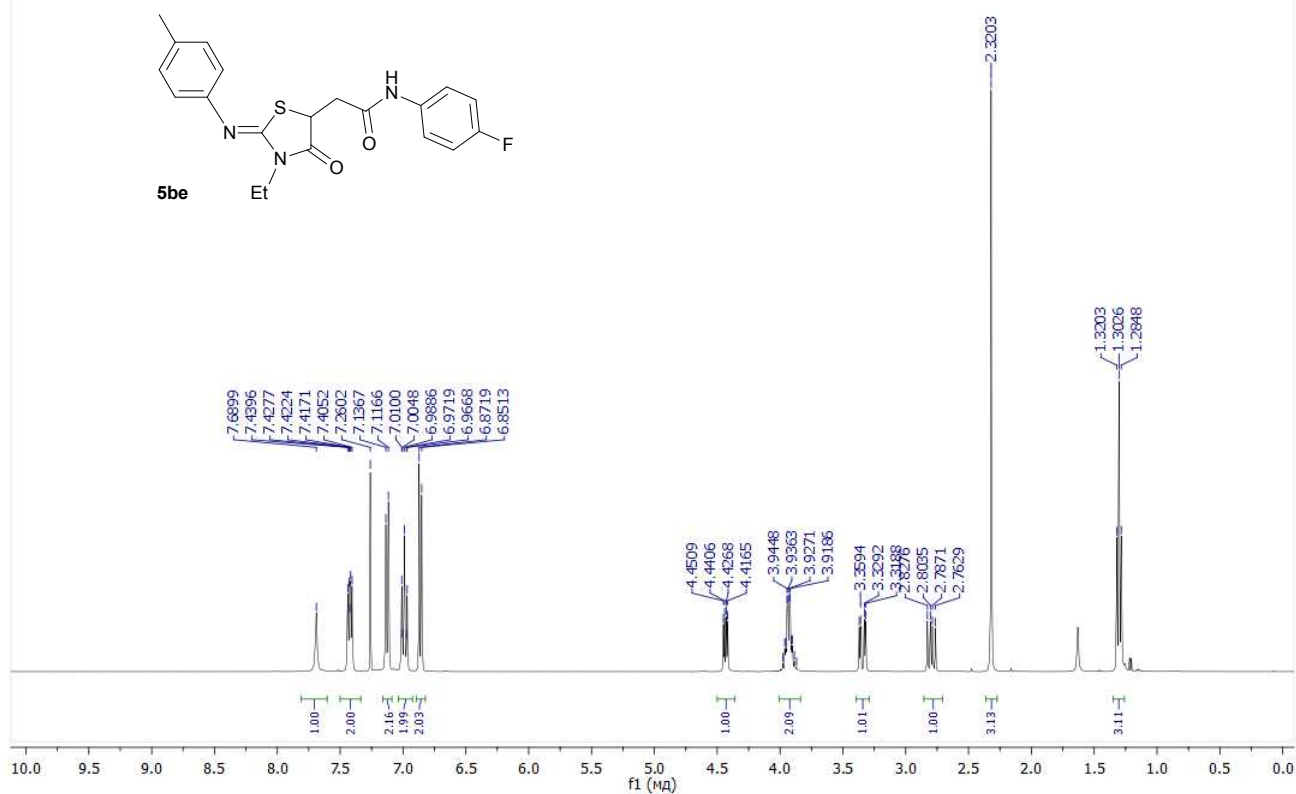
MAK, 848, BF = 400.13 MHz, Solvent - DMSO, 17 Apr 2013 T=299 K



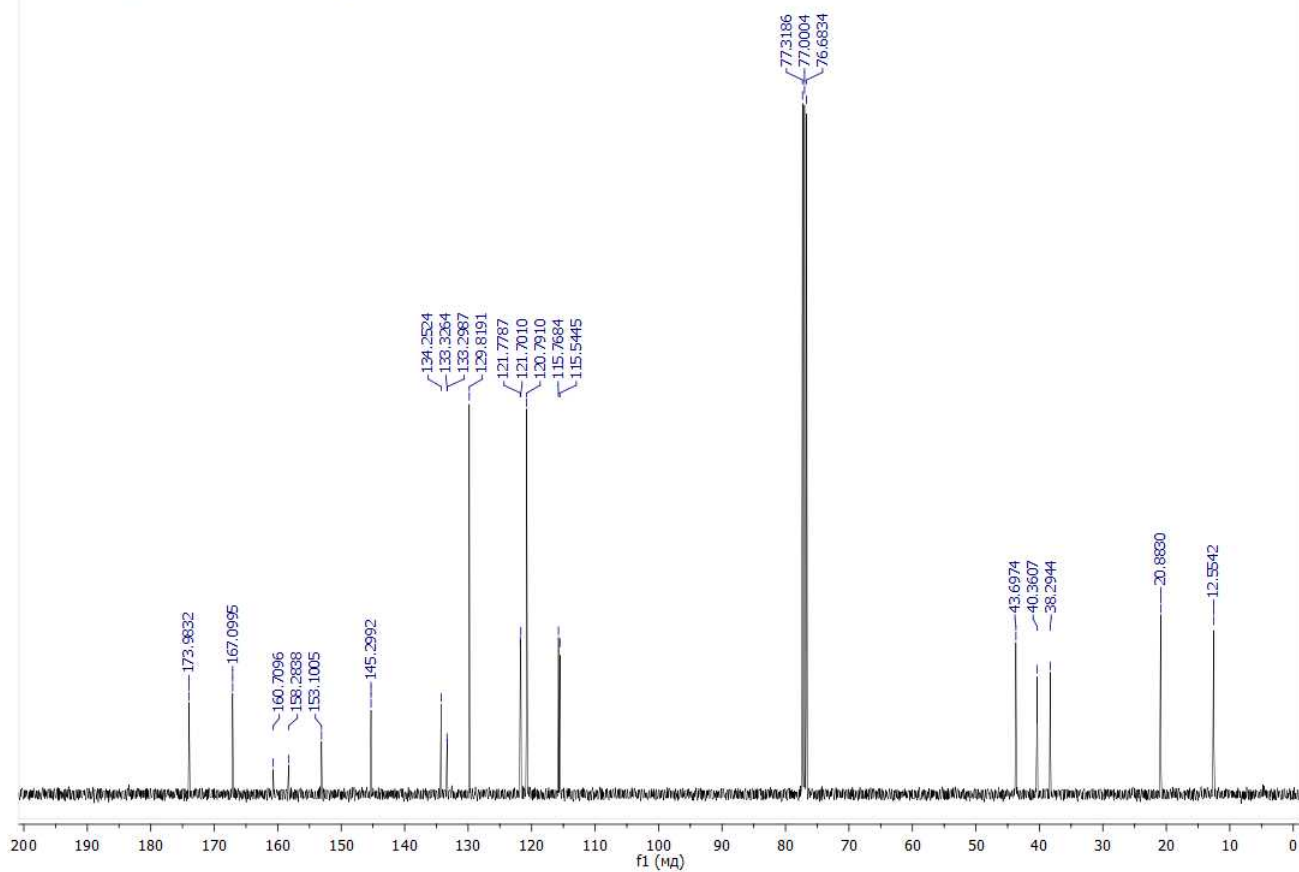
MAKc, 848, BF = 100.612769 MHz, Solvent - DMSO, 17 Apr 2013 T=300 K



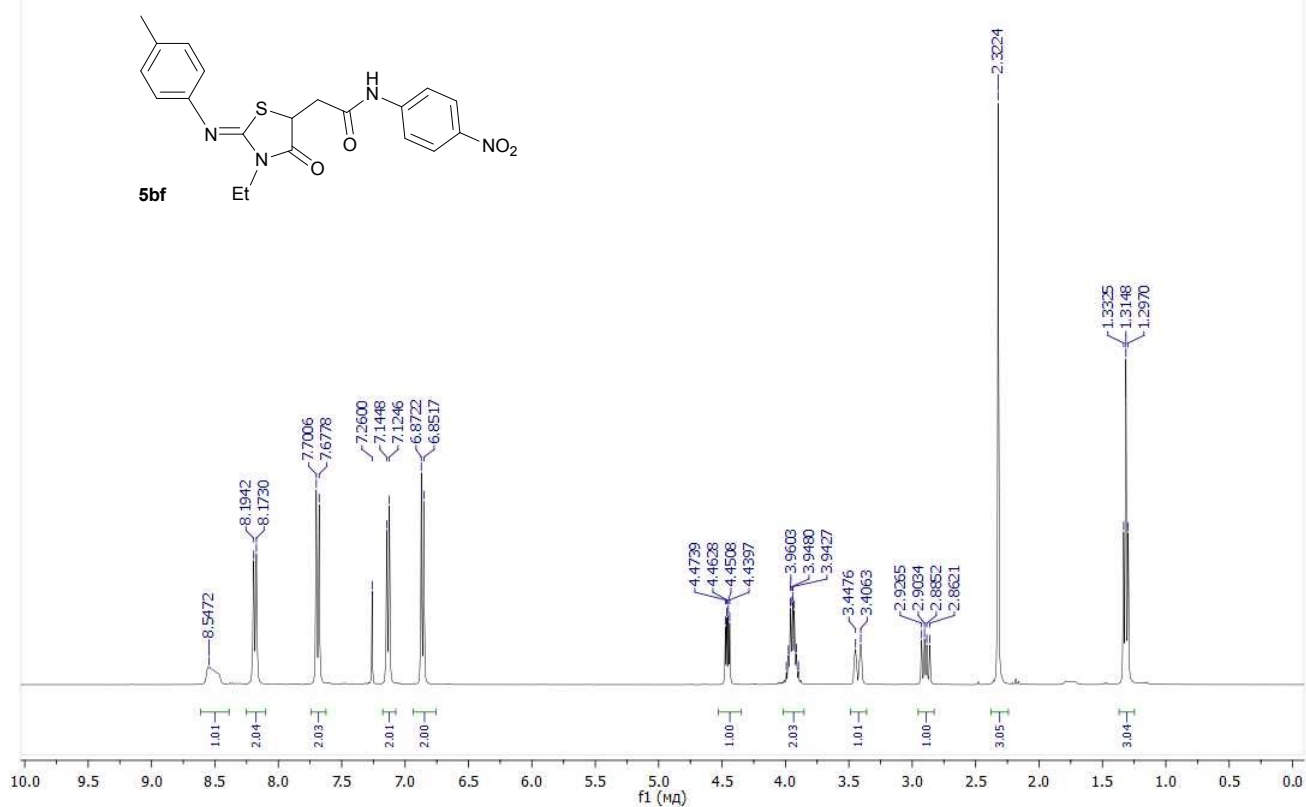
MAK, 989, BF = 400.13 MHz, Solvent - CDD3, 08 Jul 2013 T=298 K



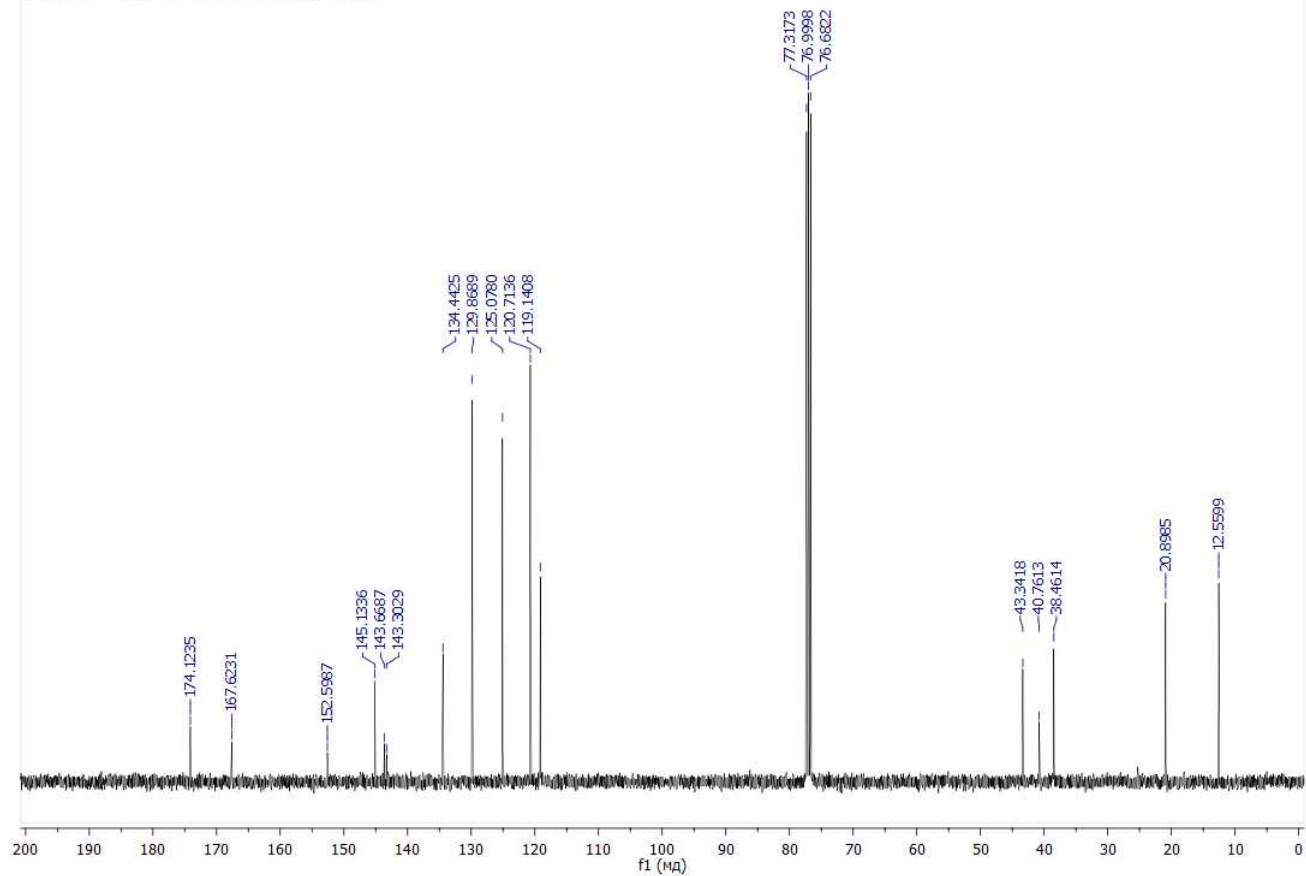
MAKc, 989, BF = 100.612769 MHz, Solvent - CDD3, 10 Jul 2013 T=298 K



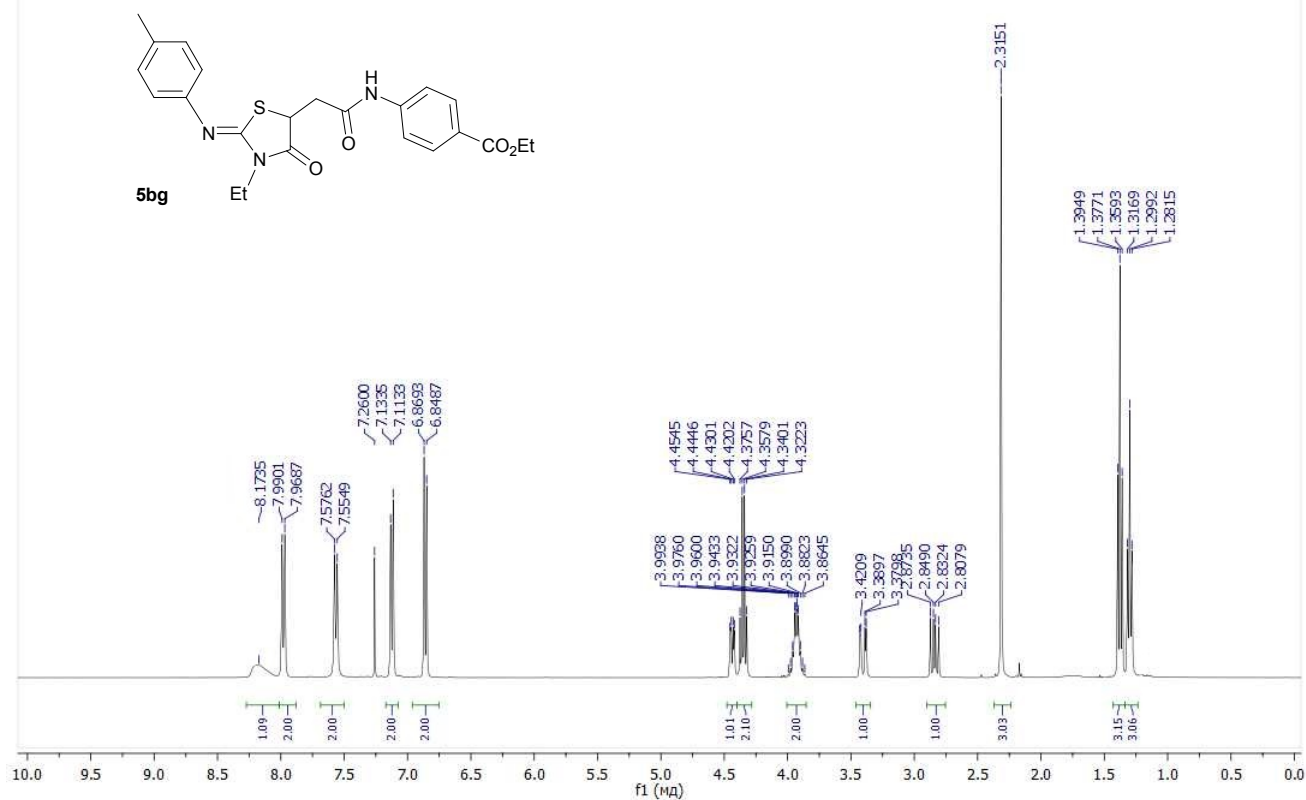
MAK, 791, BF = 400.13 MHz, Solvent - CDD3, 08 Apr 2013



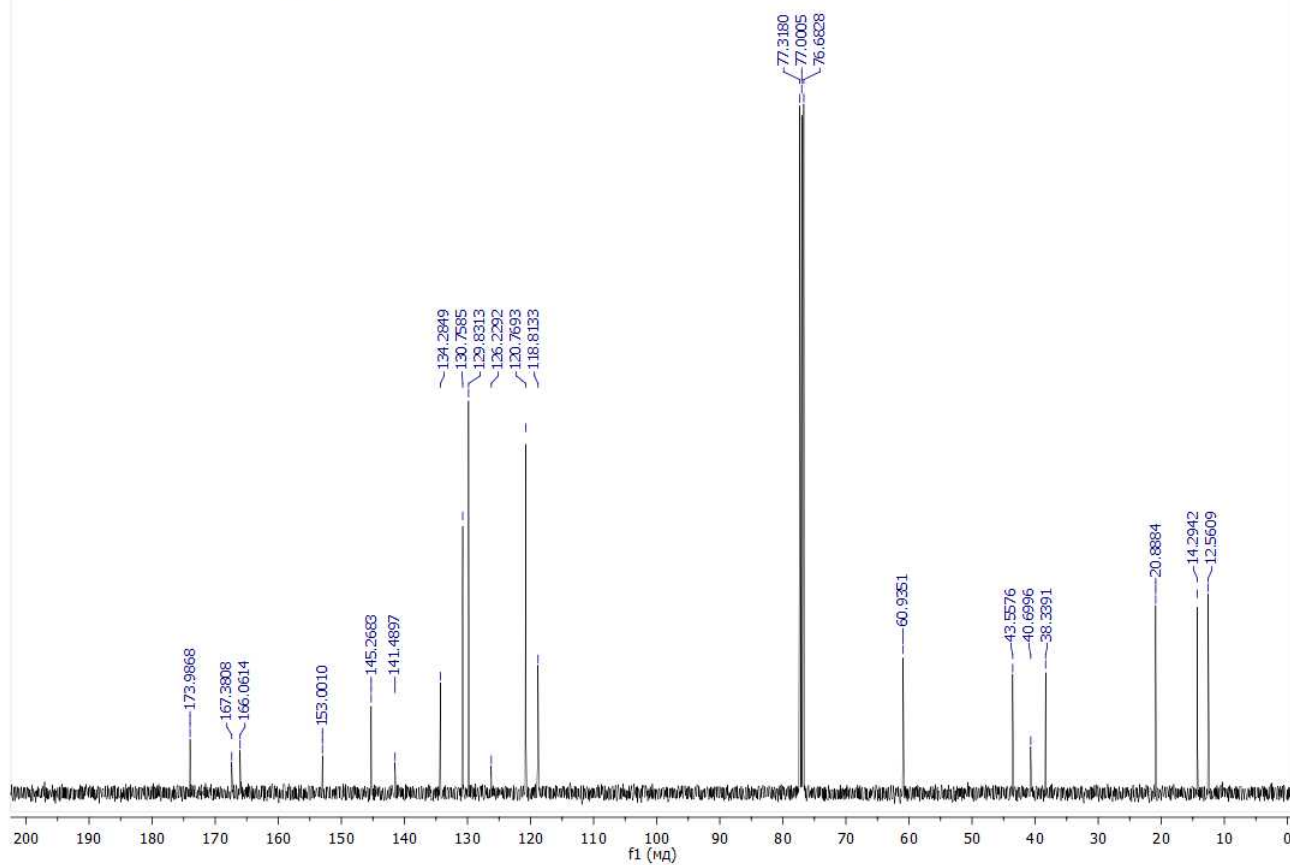
MAKc, 791, BF = 100.612769 MHz, Solvent - CDD3, 08 Apr 2013



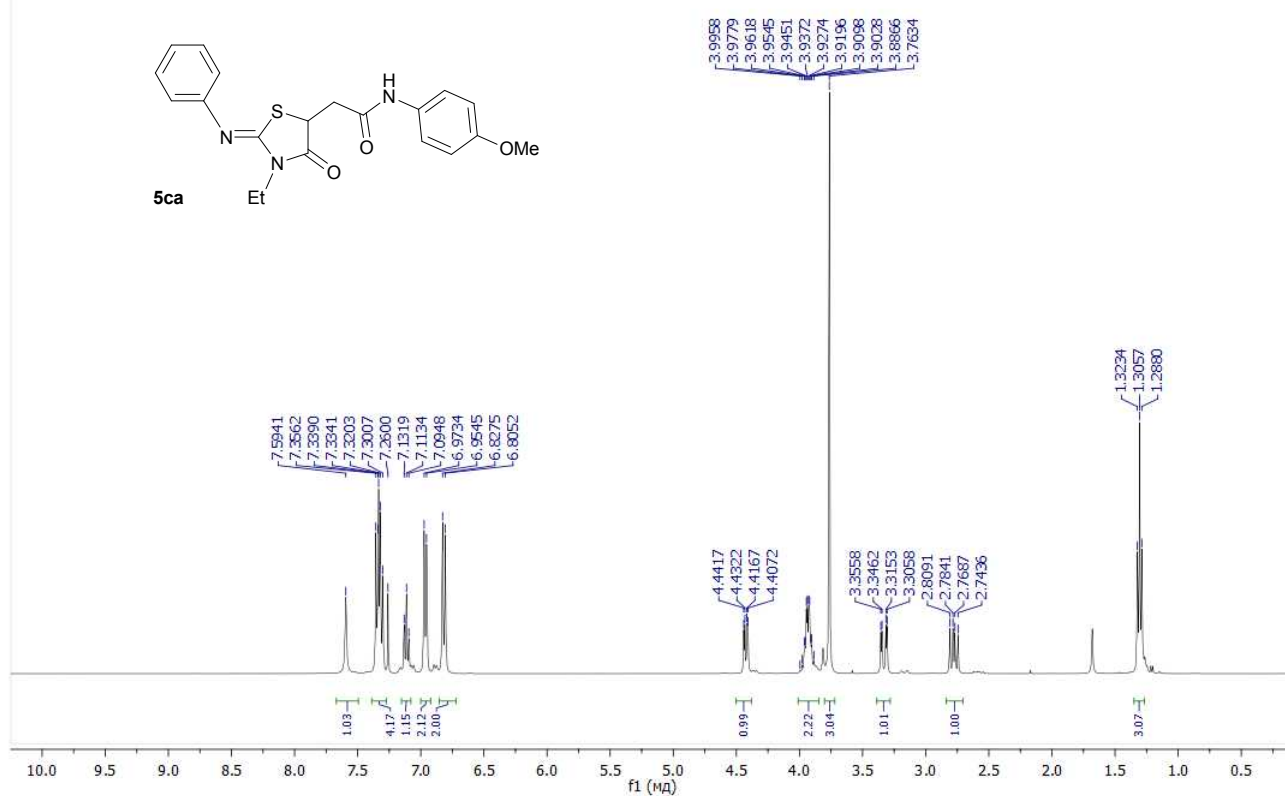
MAK, 817, BF = 400.13 MHz, Solvent - CDCl₃, 11 Apr 2013 T=299 K



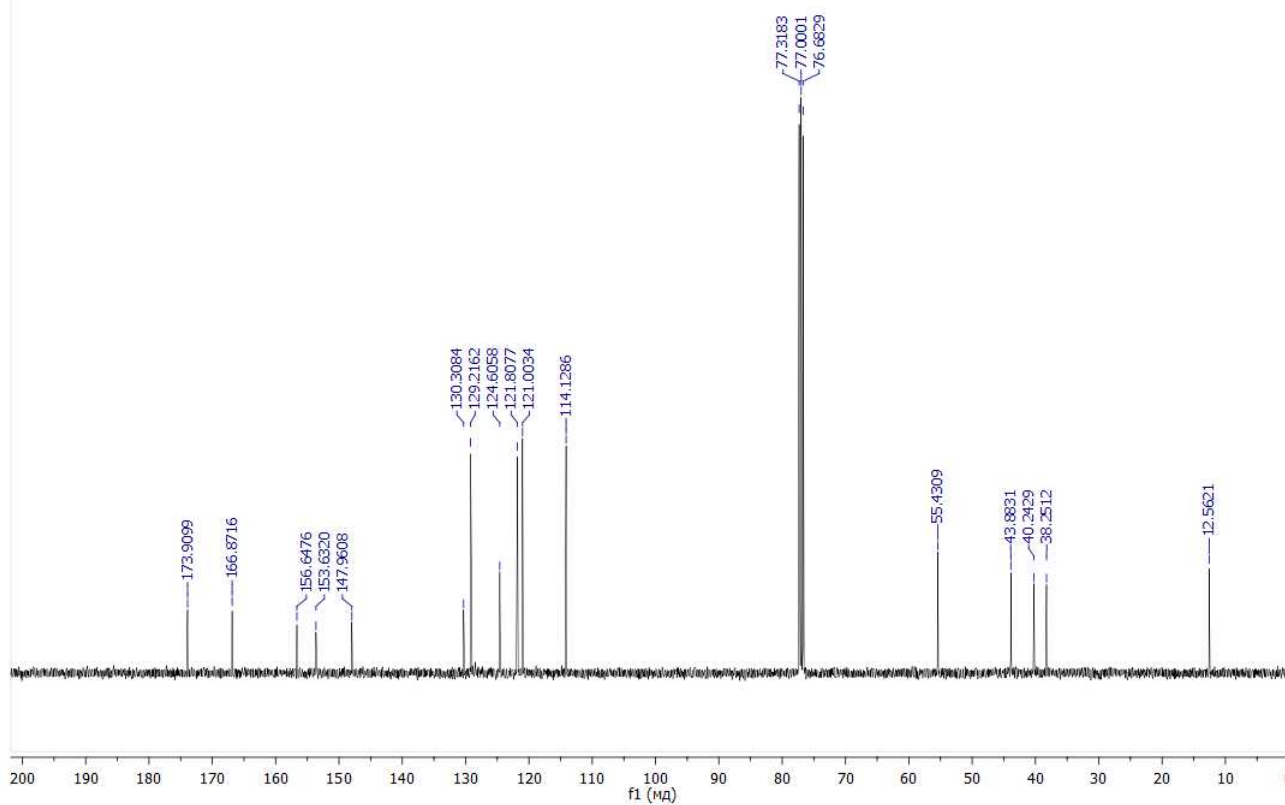
MAKc, 817, BF = 100.612769 MHz, Solvent - CDCl₃, 11 Apr 2013 T=299 K



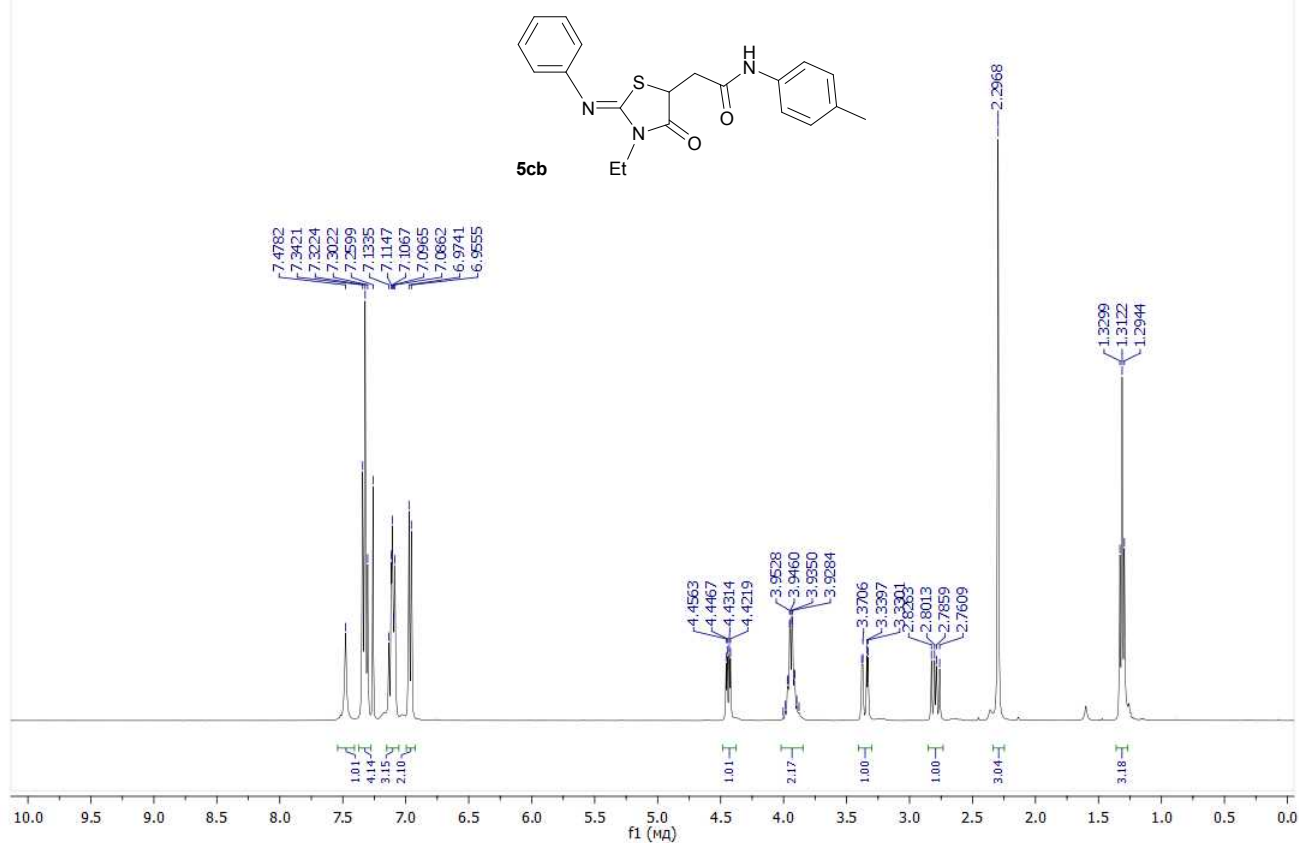
MAK, 871, BF = 400.13 MHz, Solvent - CDCl₃, 23 Apr 2013 T=298 K



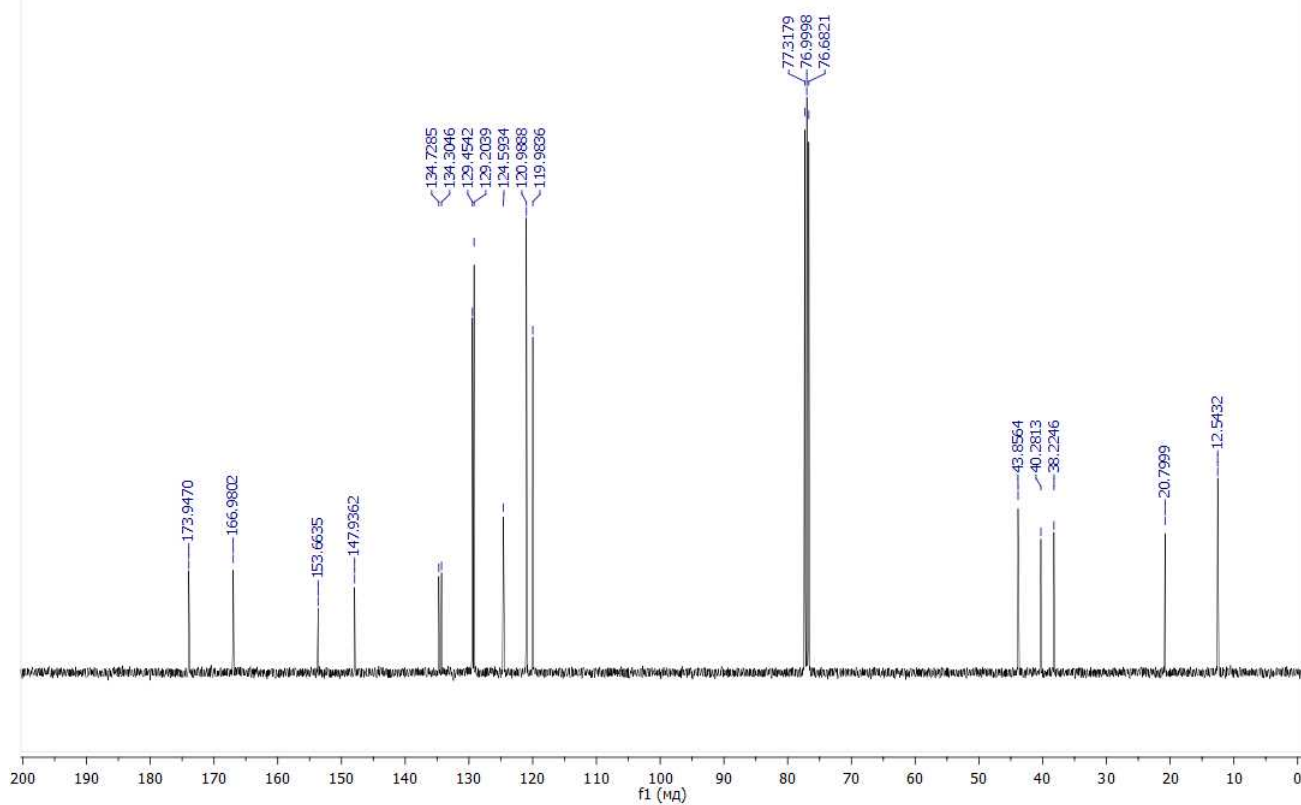
MAKc, 871, BF = 100.612769 MHz, Solvent - CDCl₃, 23 Apr 2013 T=298 K



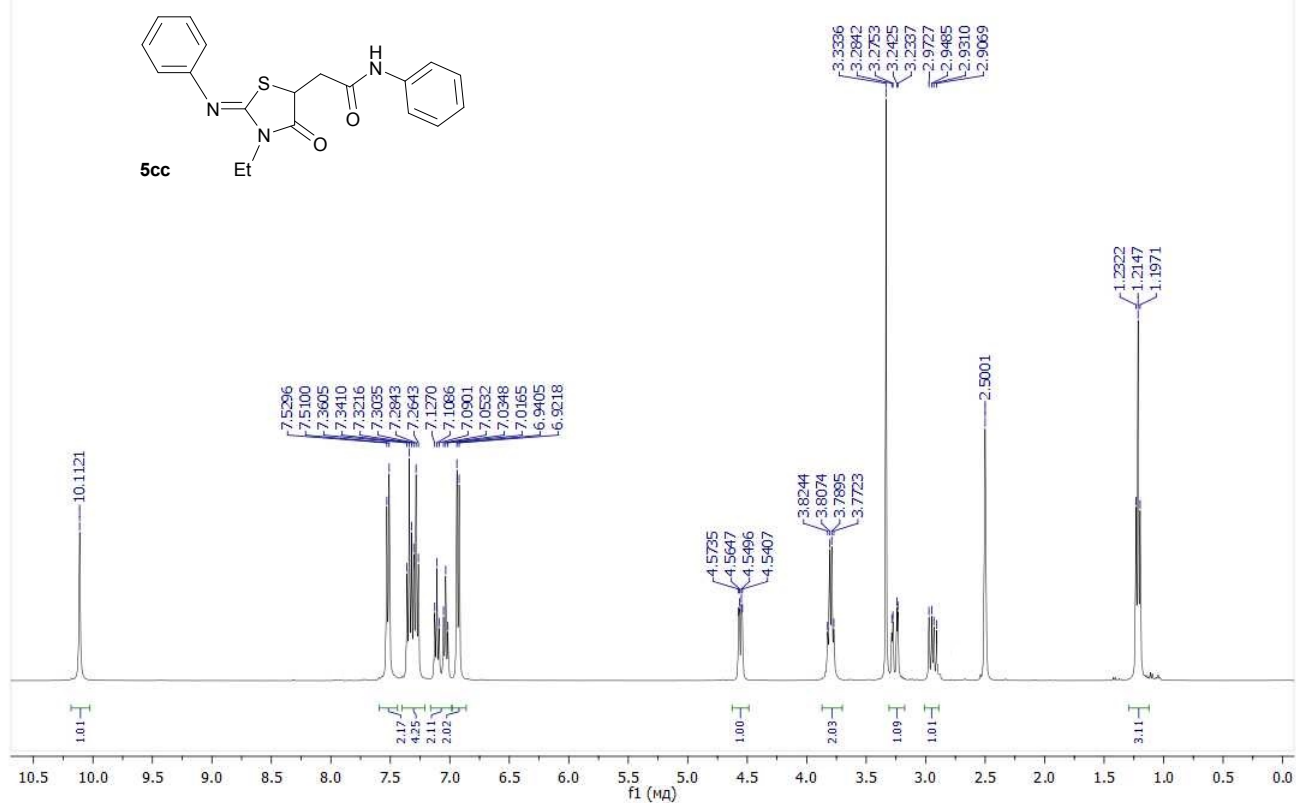
MAK, 177, BF = 400.13 MHz, Solvent - CDCl₃, 06 Nov 2013 T=299 K



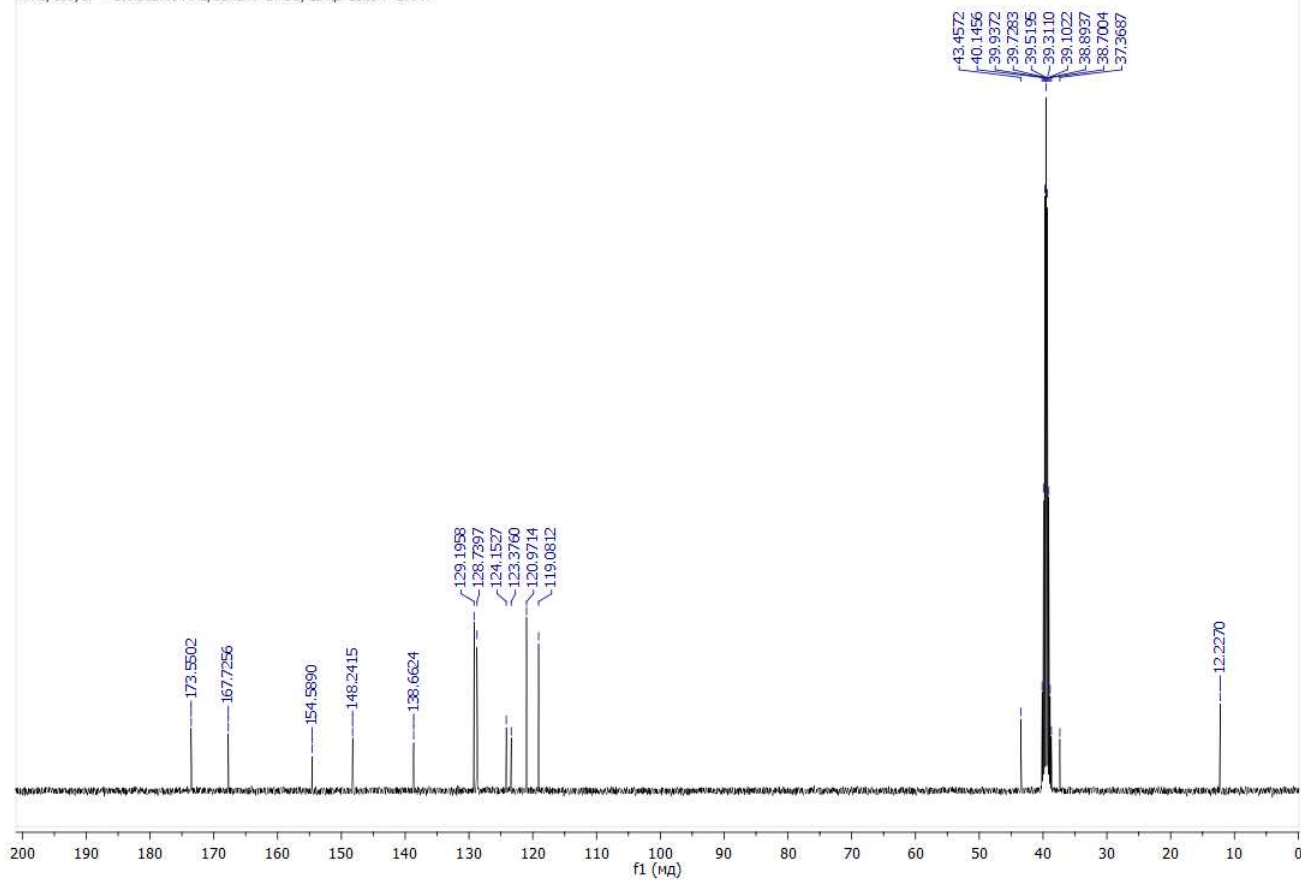
MAKc, 177, BF = 100.612769 MHz, Solvent - CDCl₃, 07 Nov 2013 T=299 K



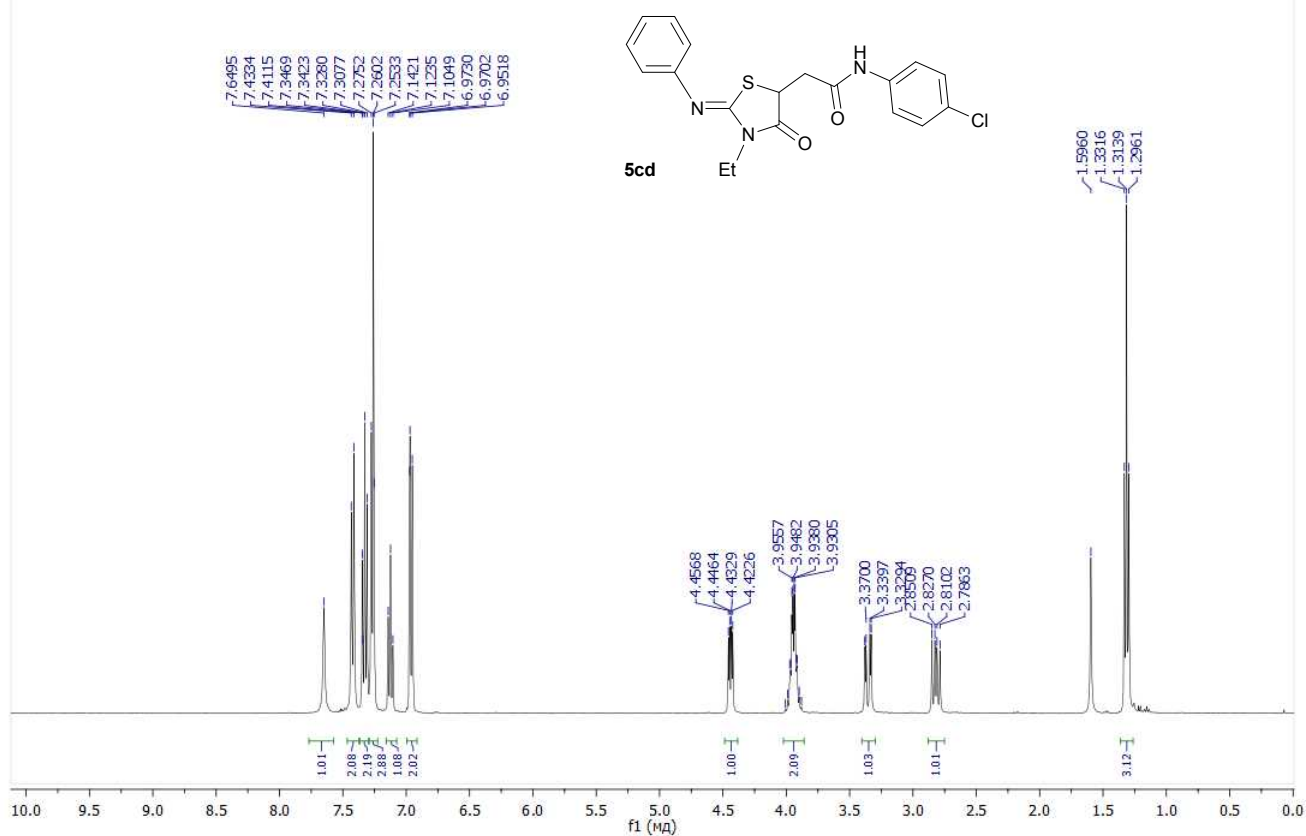
MAK, 865, BF = 400.13 MHz, Solvent - DMSO, 22 Apr 2013 T=299 K



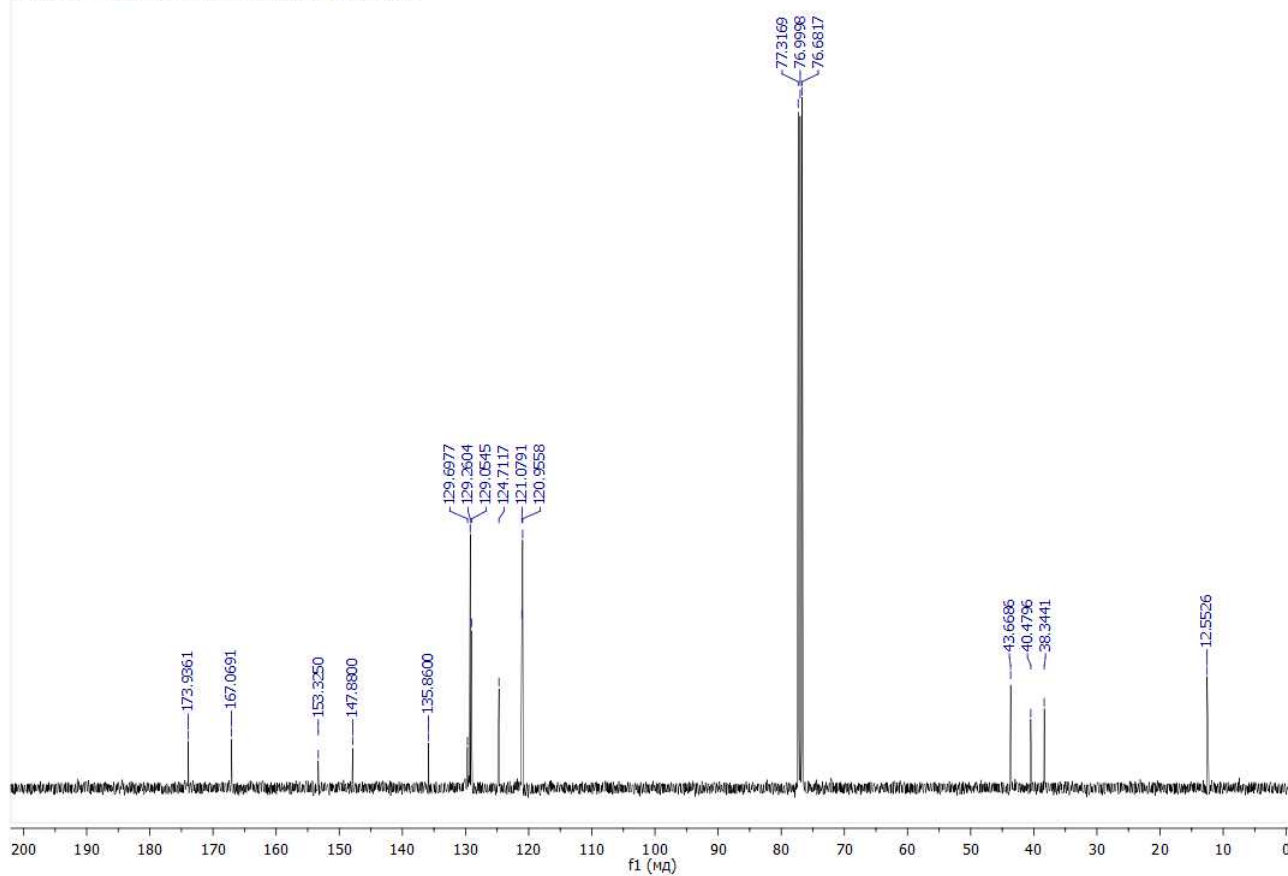
MAKc, 865, BF = 100.612769 MHz, Solvent - DMSO, 22 Apr 2013 T=299 K



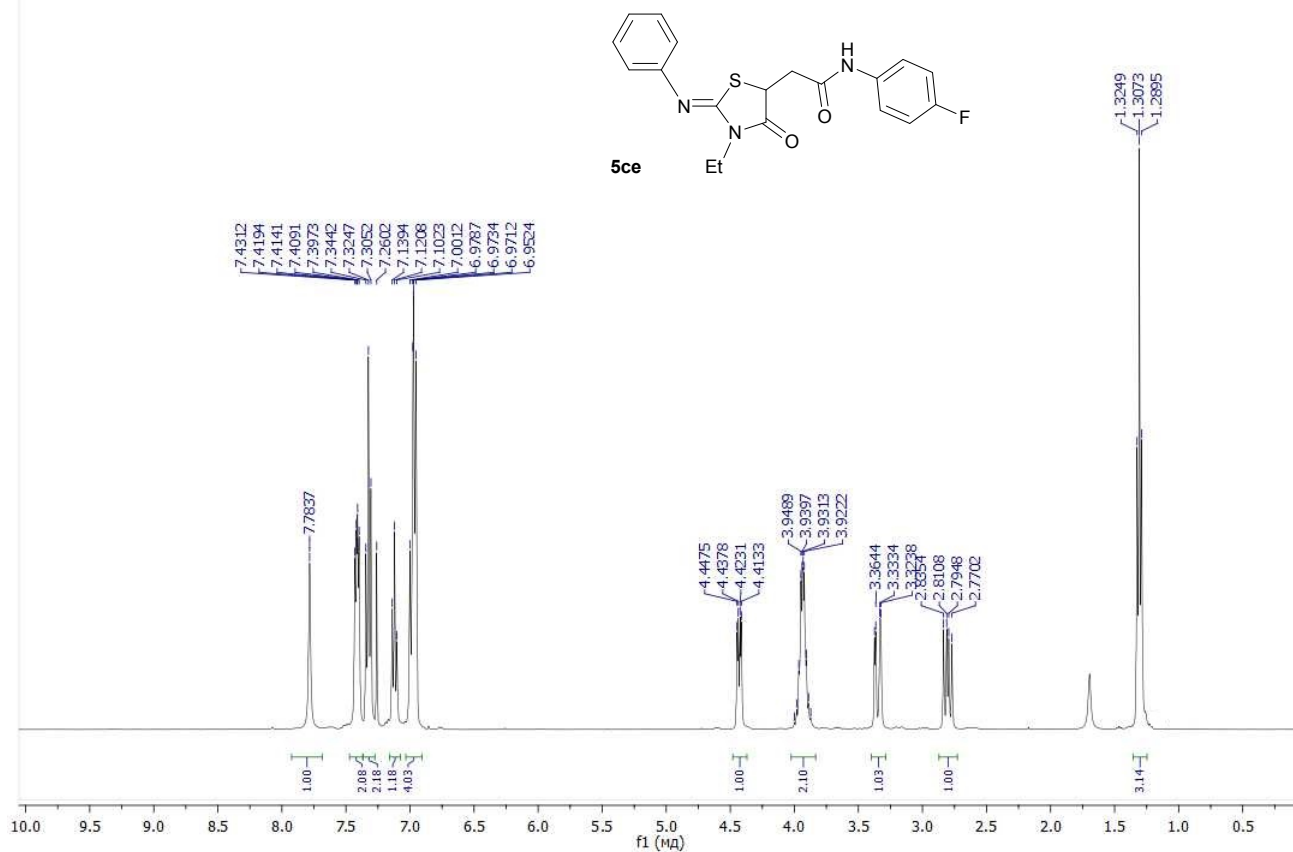
MAK, 991, BF = 400.13 MHz, Solvent - CDD3, 08 Jul 2013 T=298 K



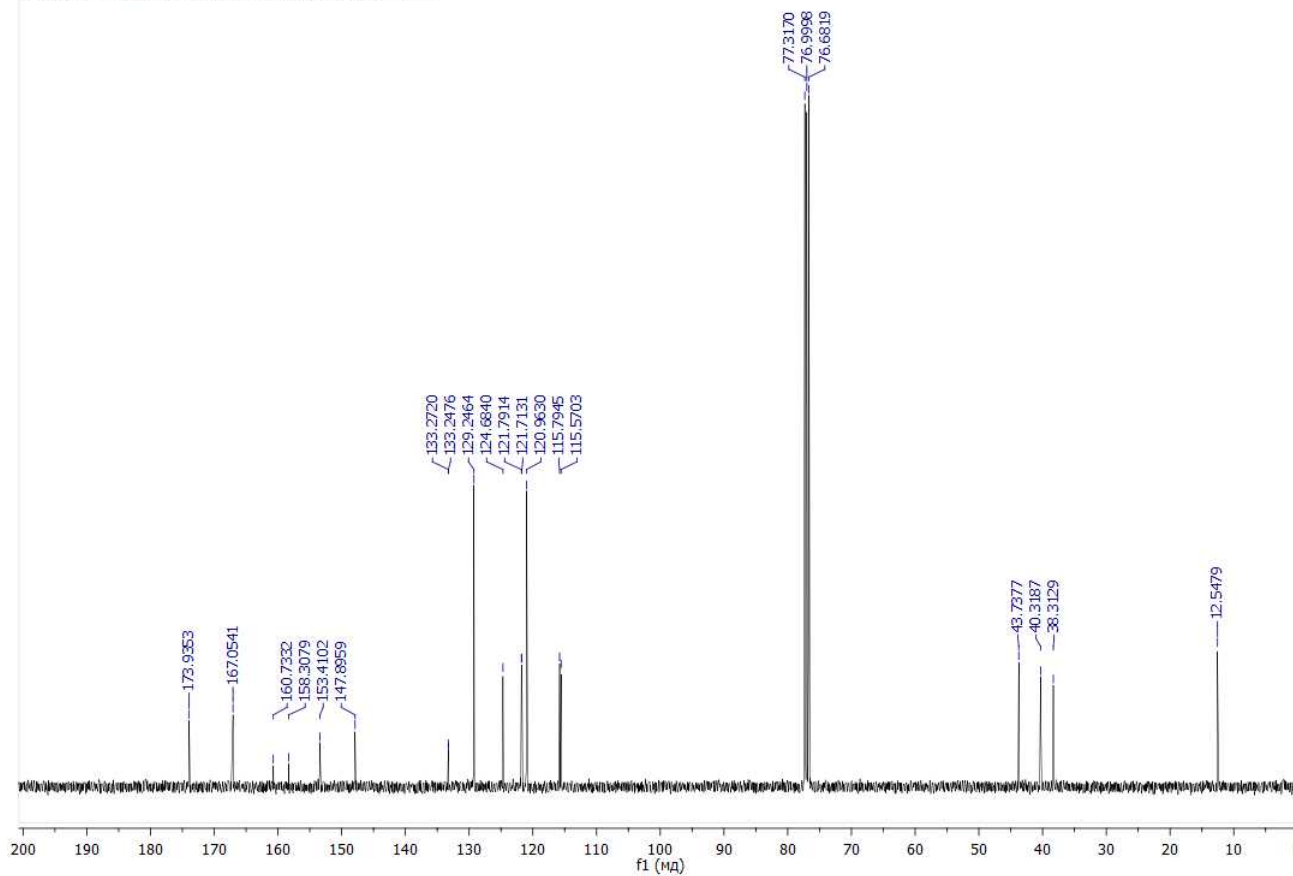
MAKc, 991, BF = 100.612769 MHz, Solvent - CDD3, 10 Jul 2013 T=298 K



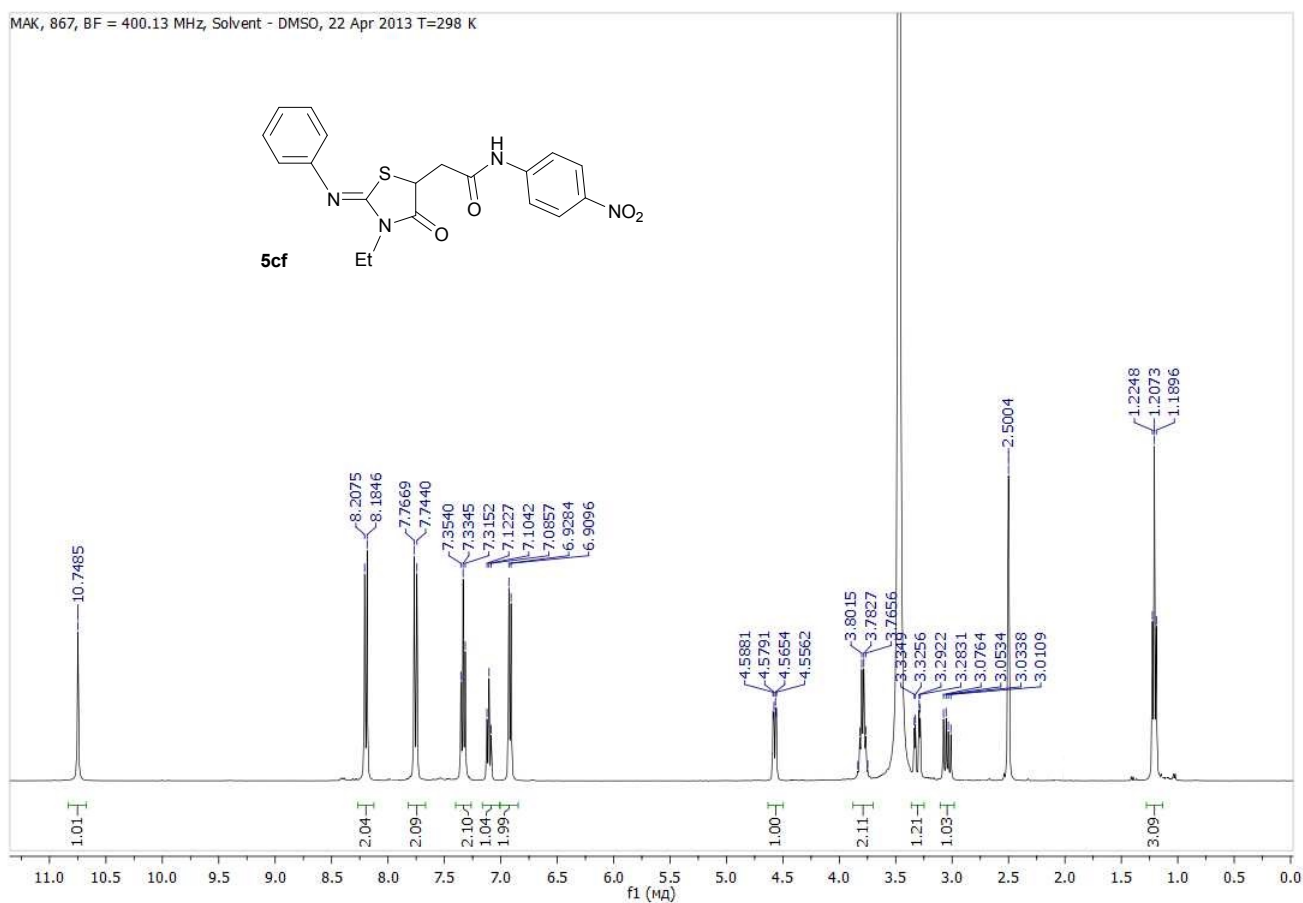
MAK, 872, BF = 400.13 MHz, Solvent - CDD3, 23 Apr 2013 T=298 K



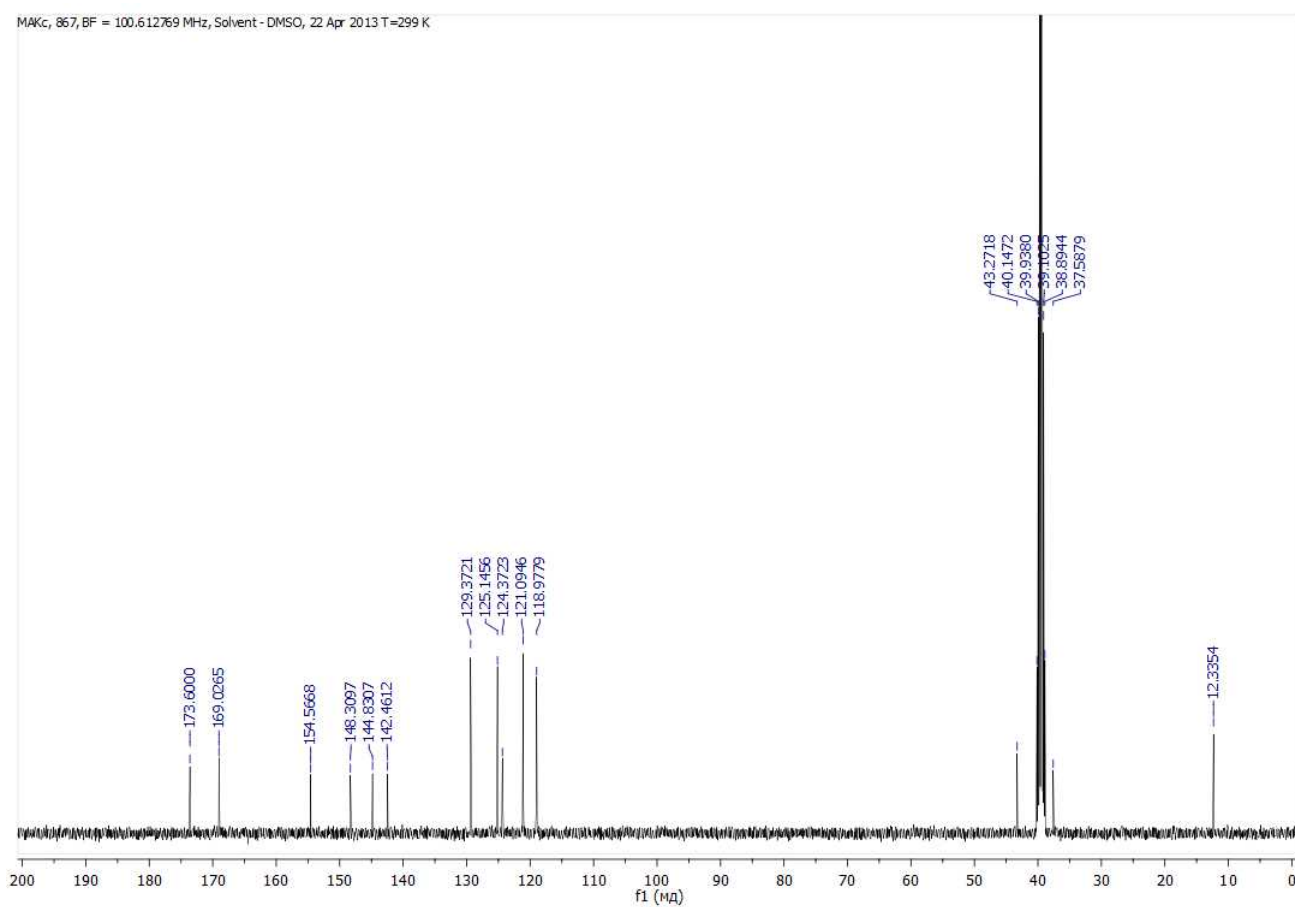
MAKc, 872, BF = 100.612769 MHz, Solvent - CDD3, 23 Apr 2013 T=299 K



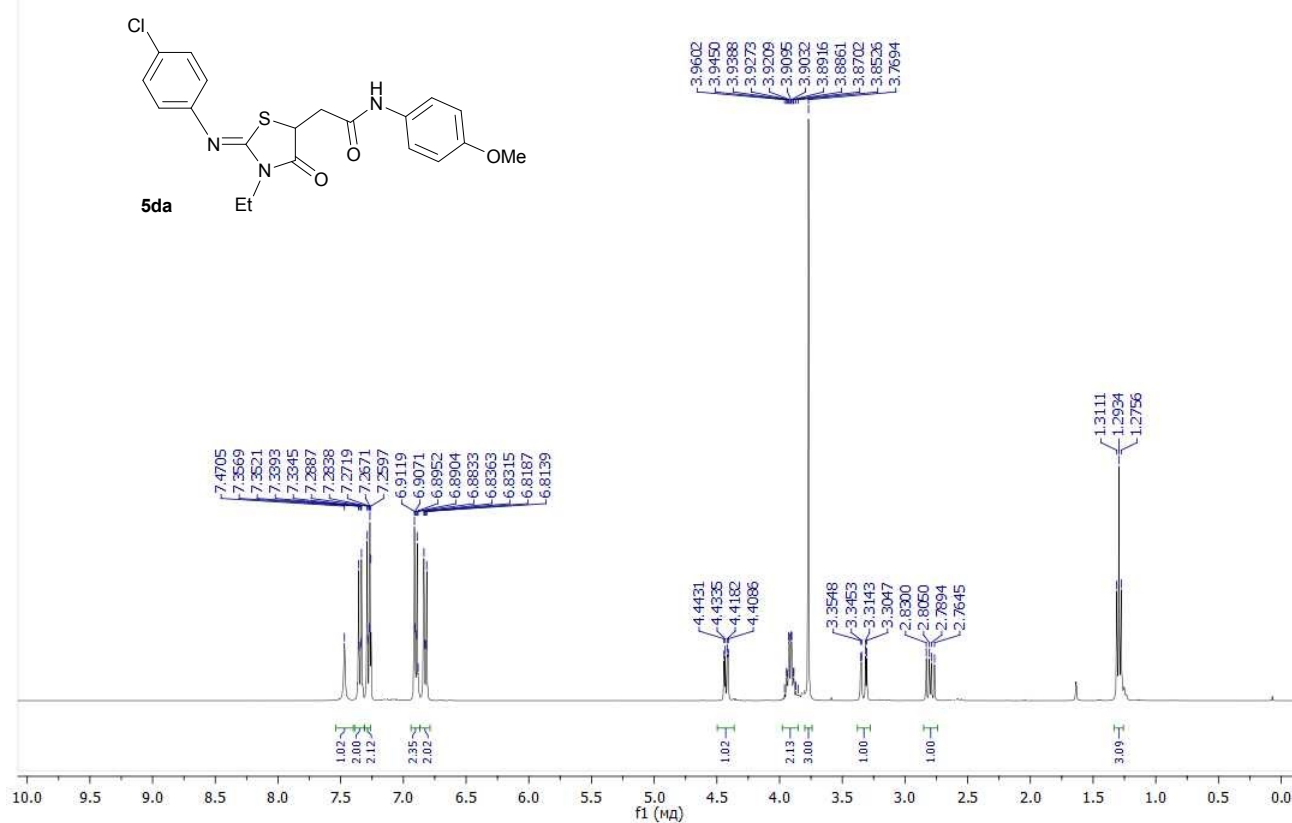
MAK, 867, BF = 400.13 MHz, Solvent - DMSO, 22 Apr 2013 T=298 K



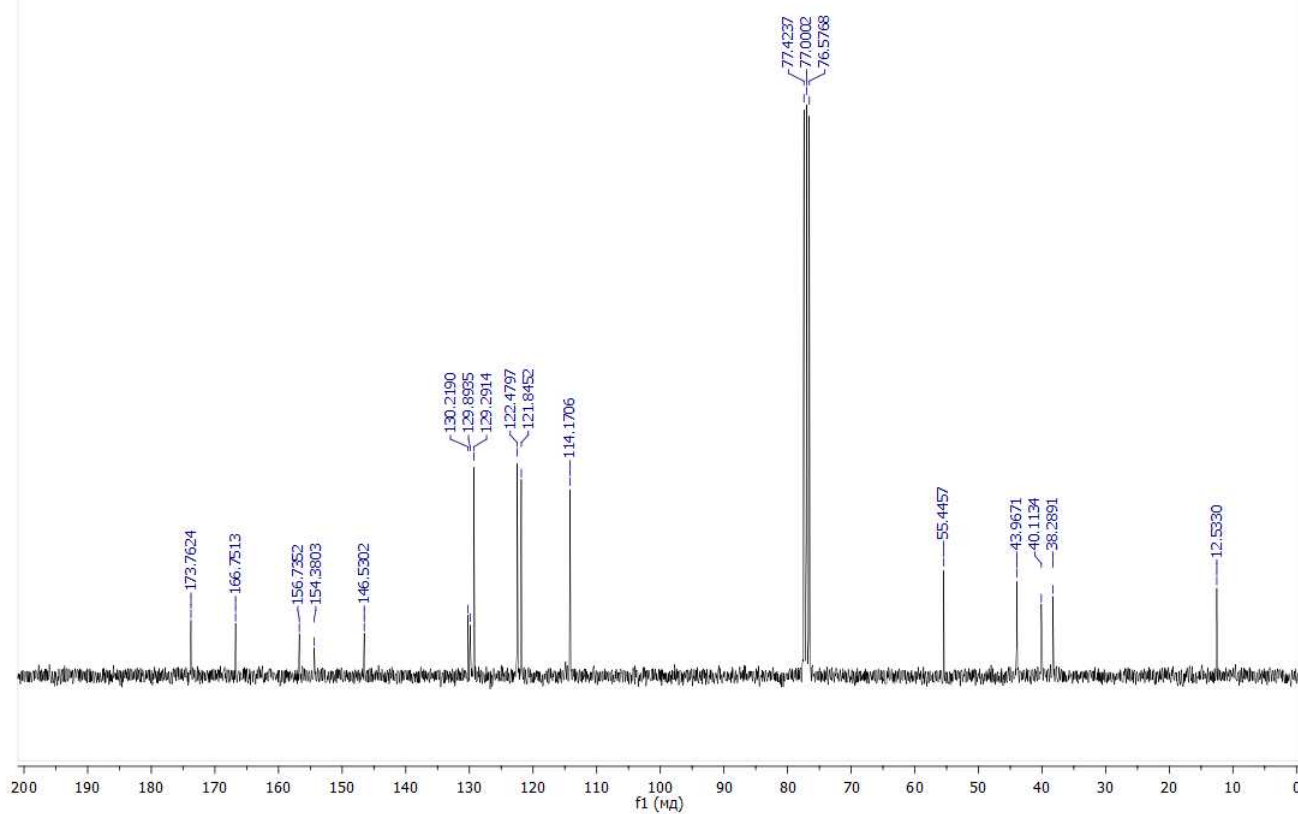
MAKc, 867, BF = 100.612769 MHz, Solvent - DMSO, 22 Apr 2013 T=299 K



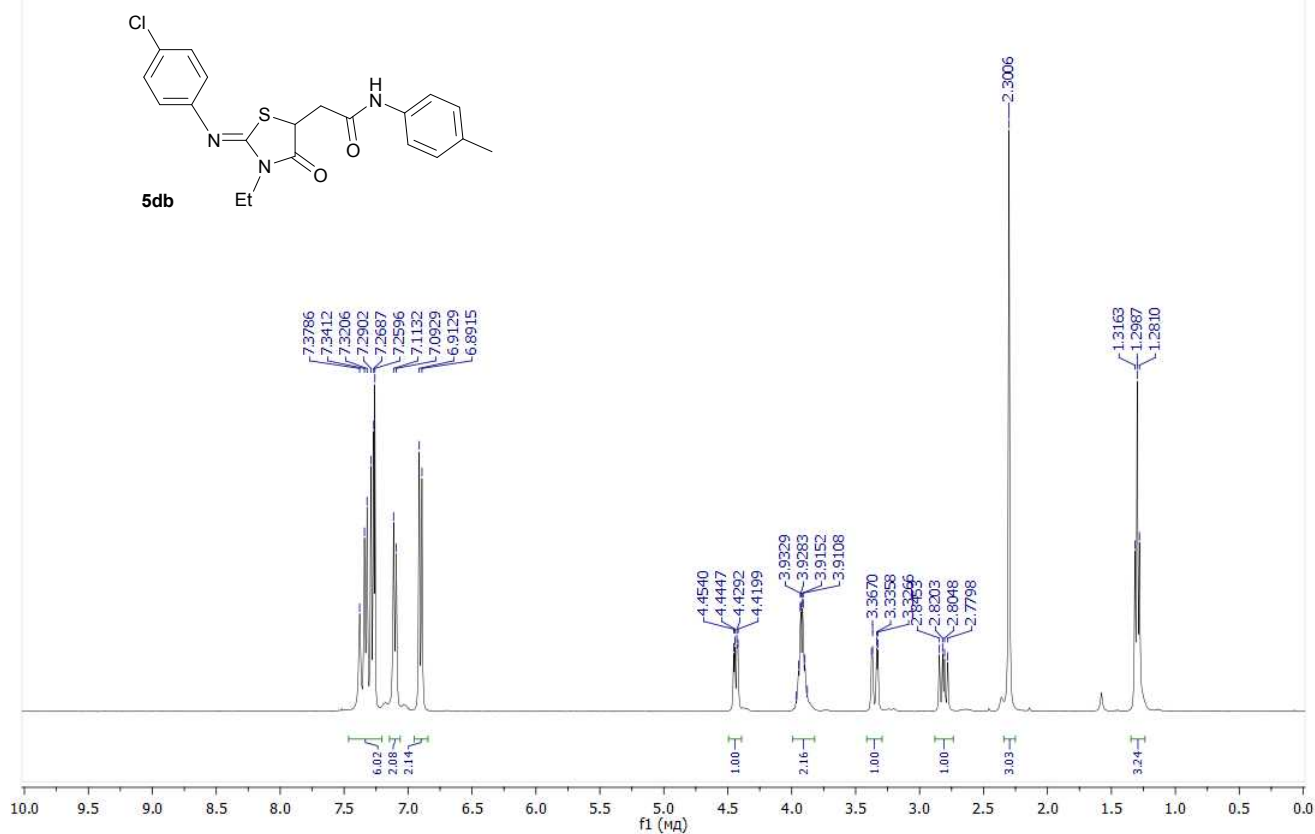
MAK, 3, BF = 400.13 MHz, Solvent - CDCl₃, 12 Jul 2013 T=299 K



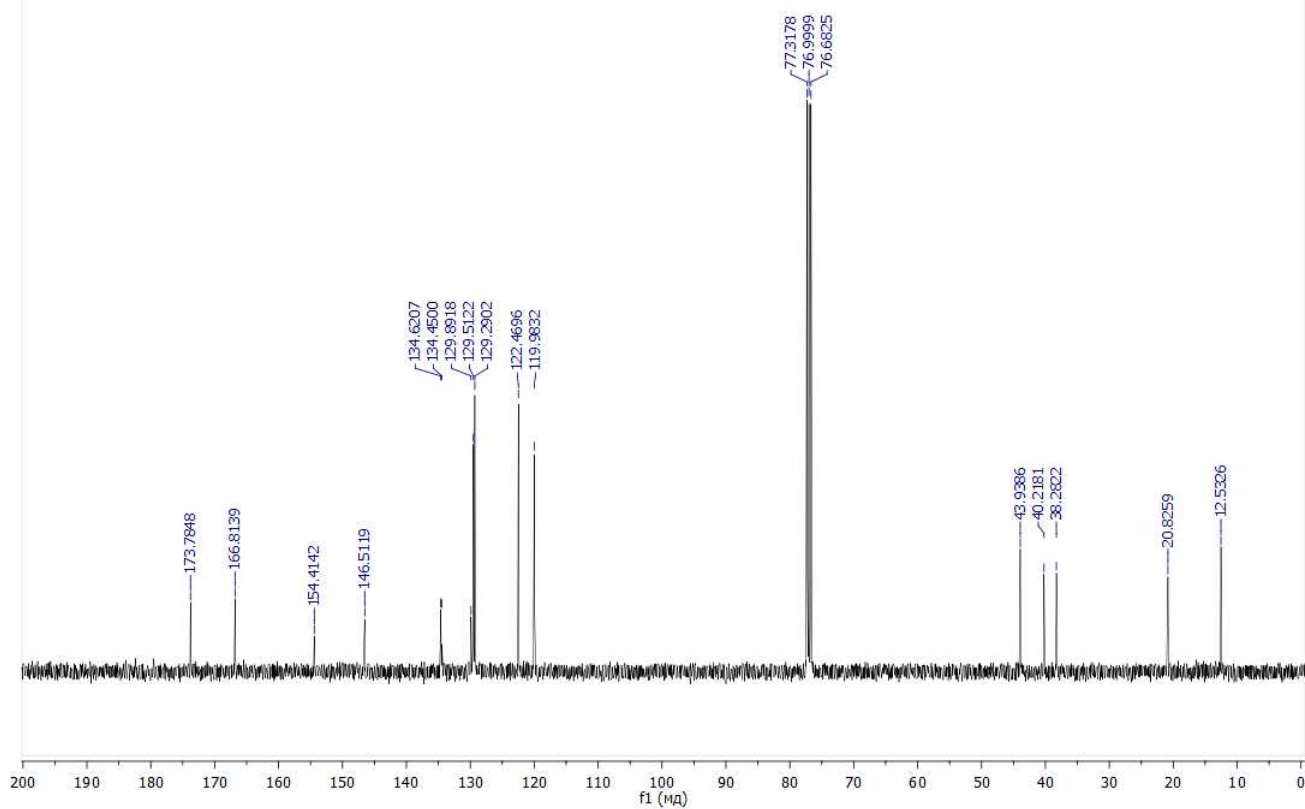
MAKc, 757, BF = 75.47 MHz, SOLVENT - CDCl₃, 29 Mar 2013



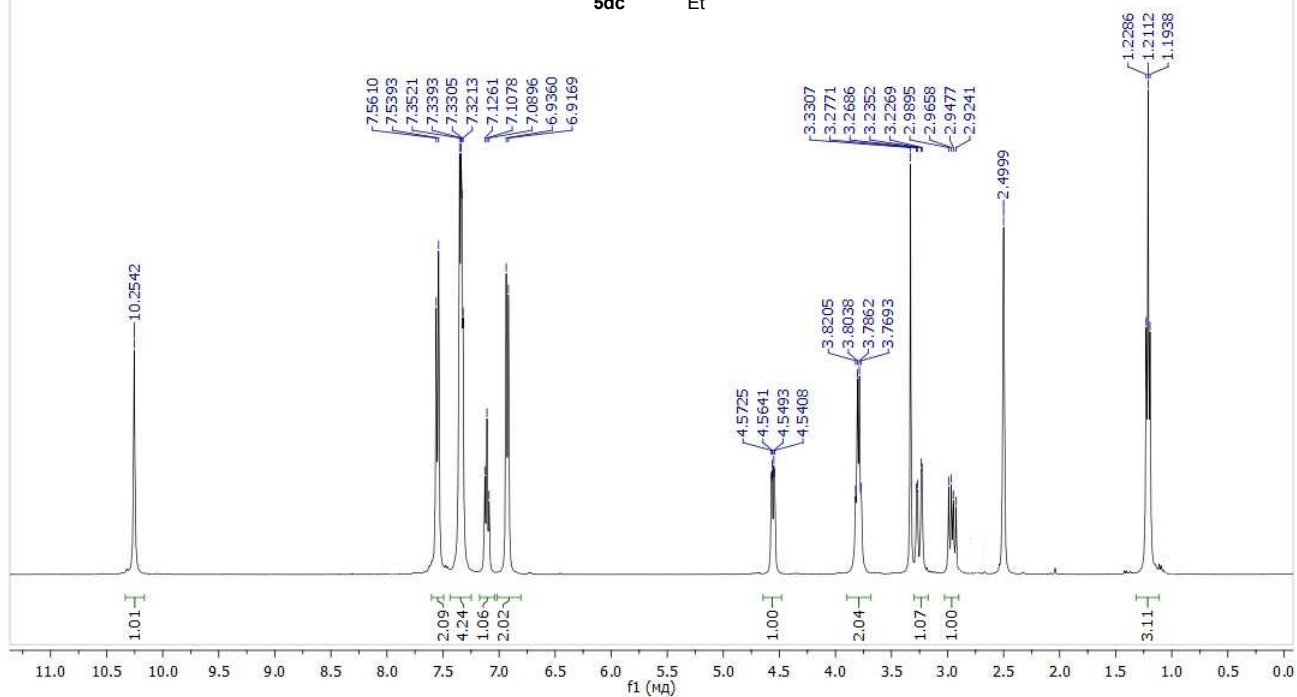
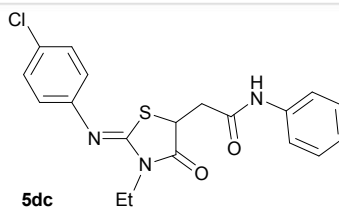
MAK, 184, BF = 400.13 MHz, Solvent - CDCl₃, 08 Nov 2013 T=298 K



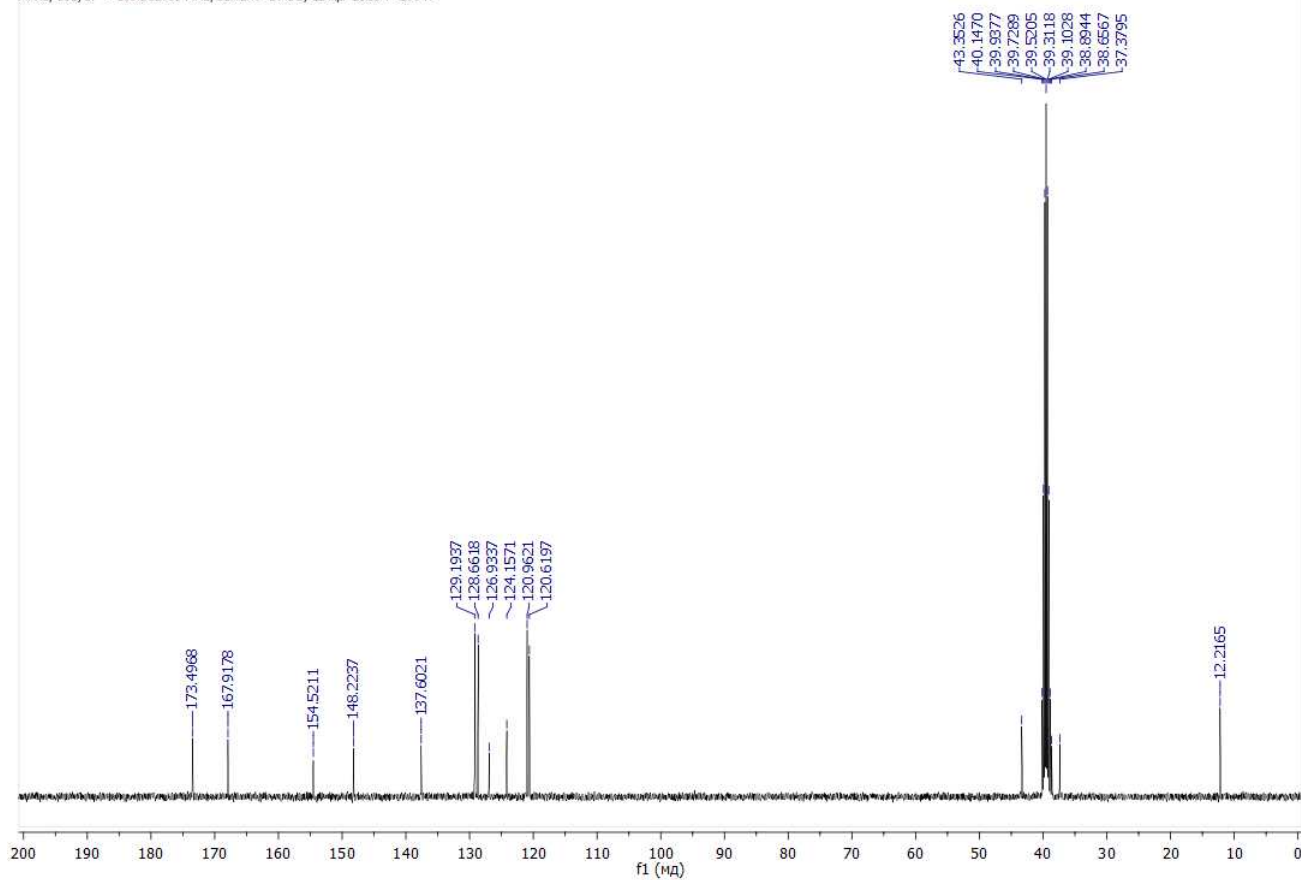
MAK, 184, BF = 100.612769 MHz, Solvent - CDCl₃, 11 Nov 2013 T=298 K



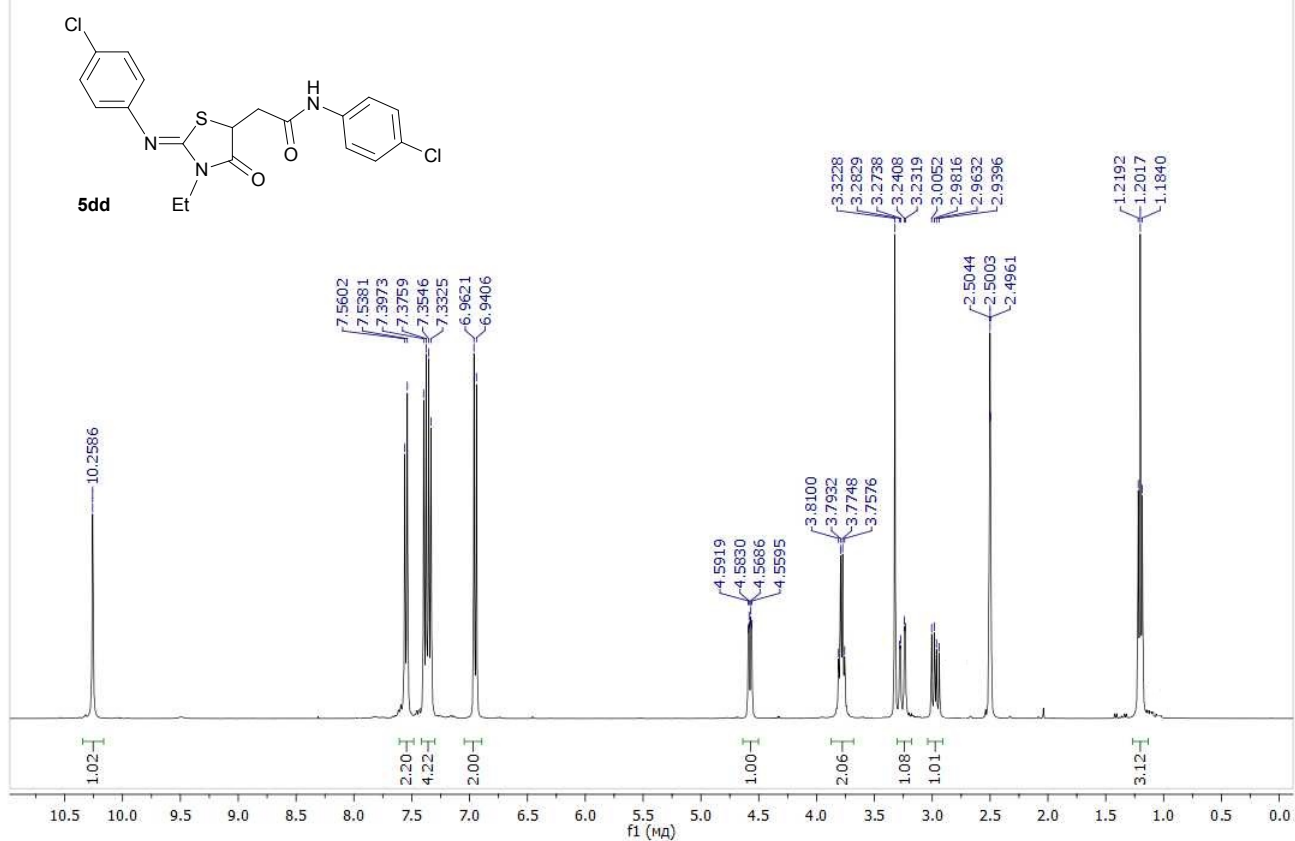
MAK, 866, BF = 400.13 MHz, Solvent - DMSO, 22 Apr 2013 T=299 K



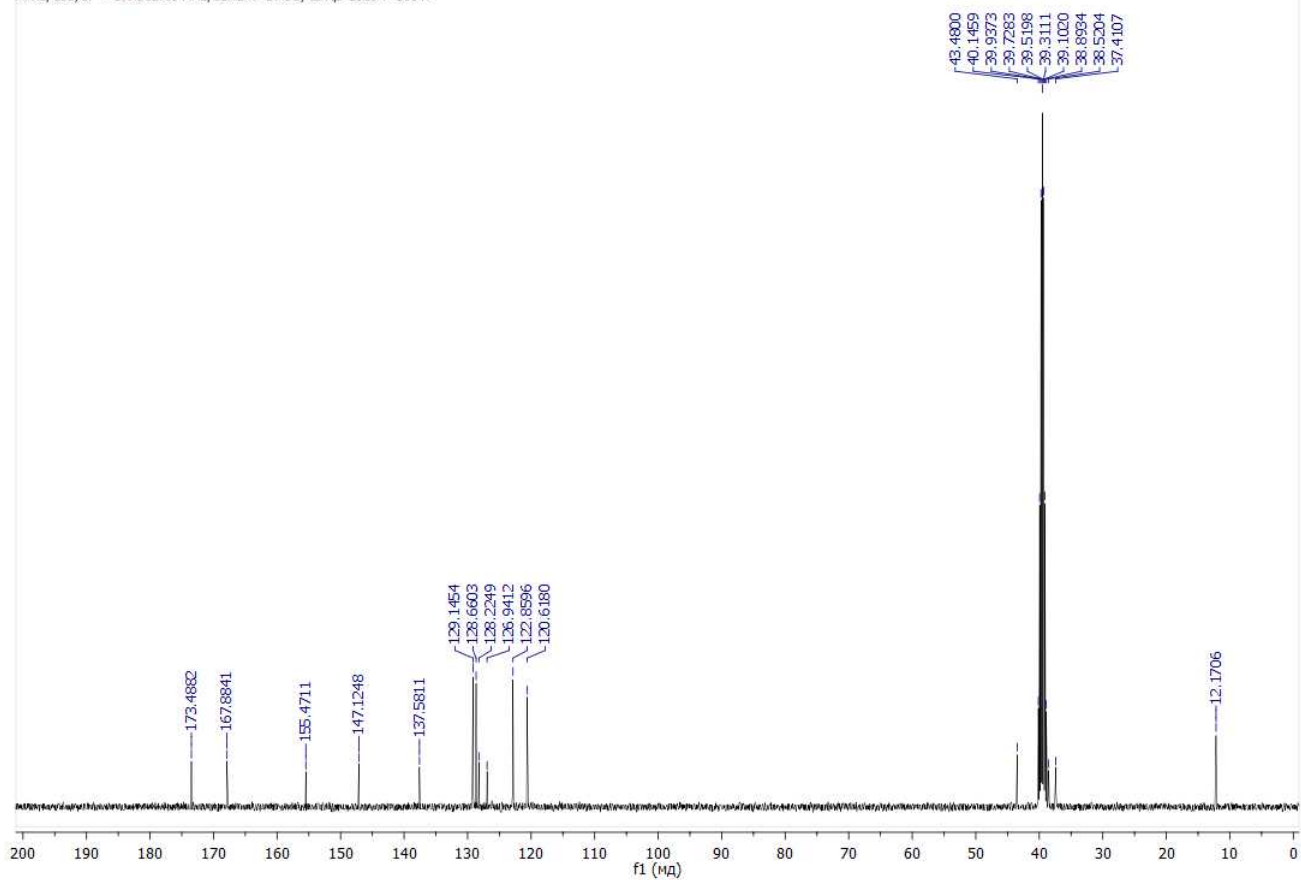
MAK, 866, BF = 100.612769 MHz, Solvent - DMSO, 22 Apr 2013 T=299 K



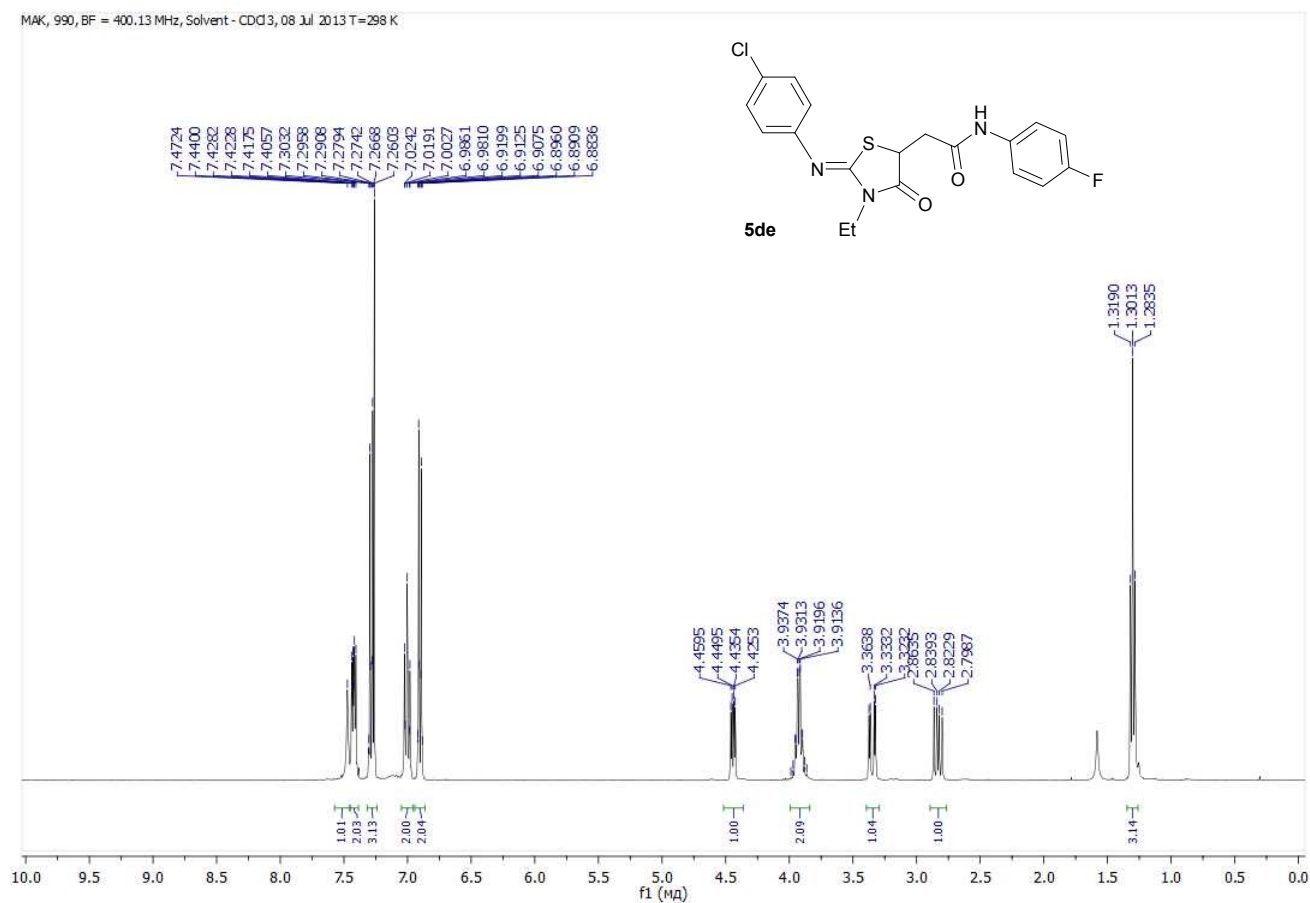
MAK, 858, BF = 400.13 MHz, Solvent - DMSO, 18 Apr 2013 T=300 K



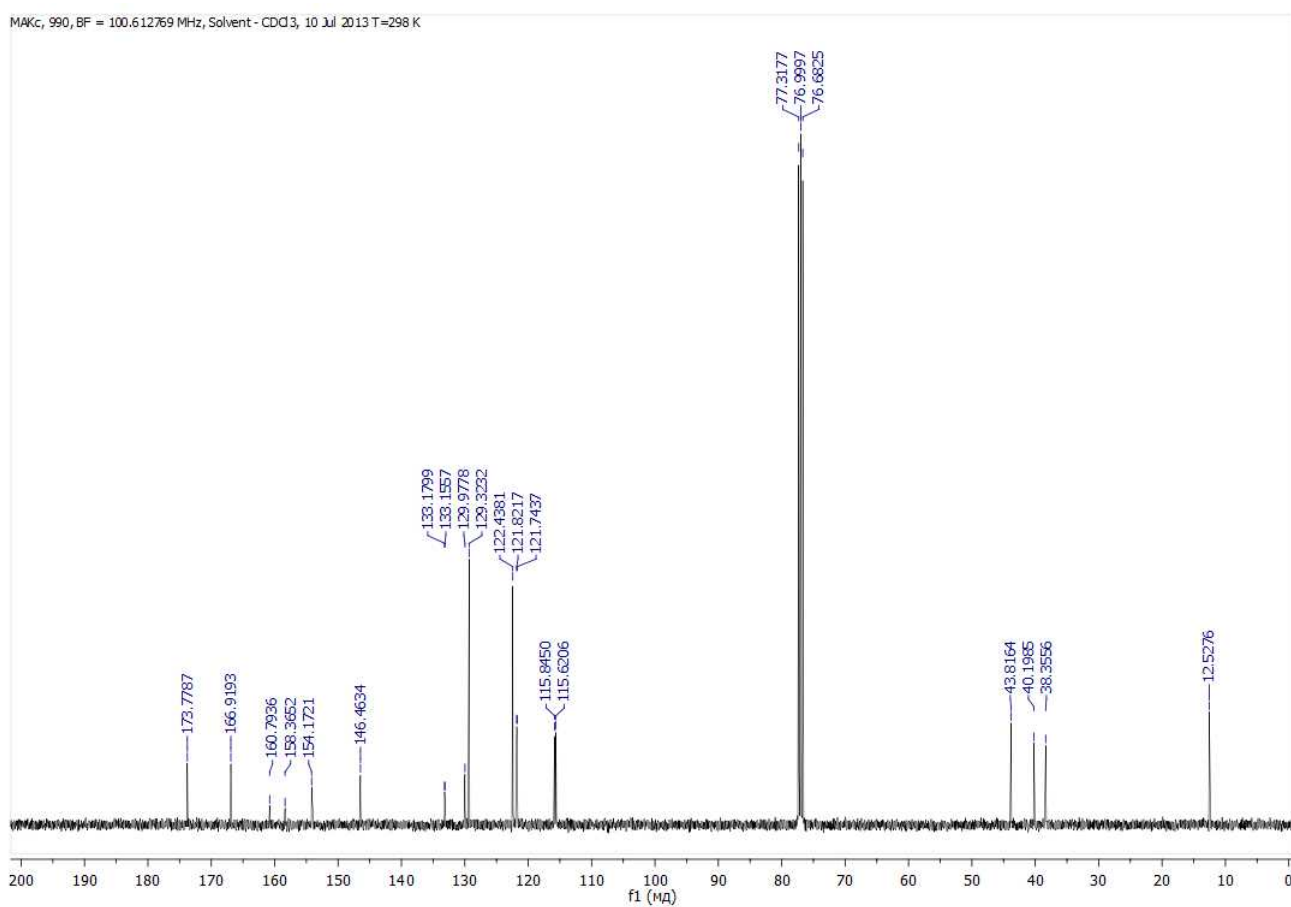
MAKc, 858, BF = 100.612769 MHz, Solvent - DMSO, 18 Apr 2013 T=300 K



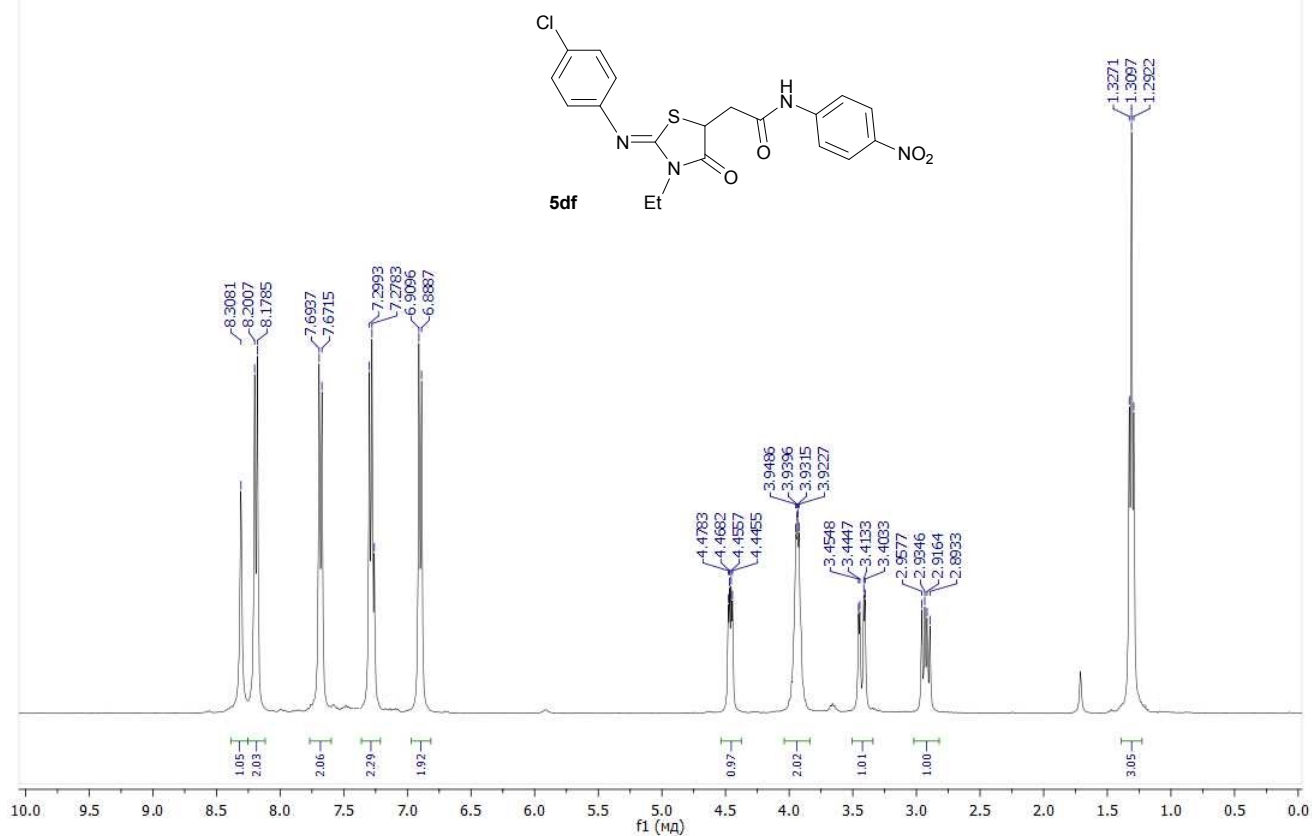
MAK, 990, BF = 400.13 MHz, Solvent - CDD3, 08 Jul 2013 T=298 K



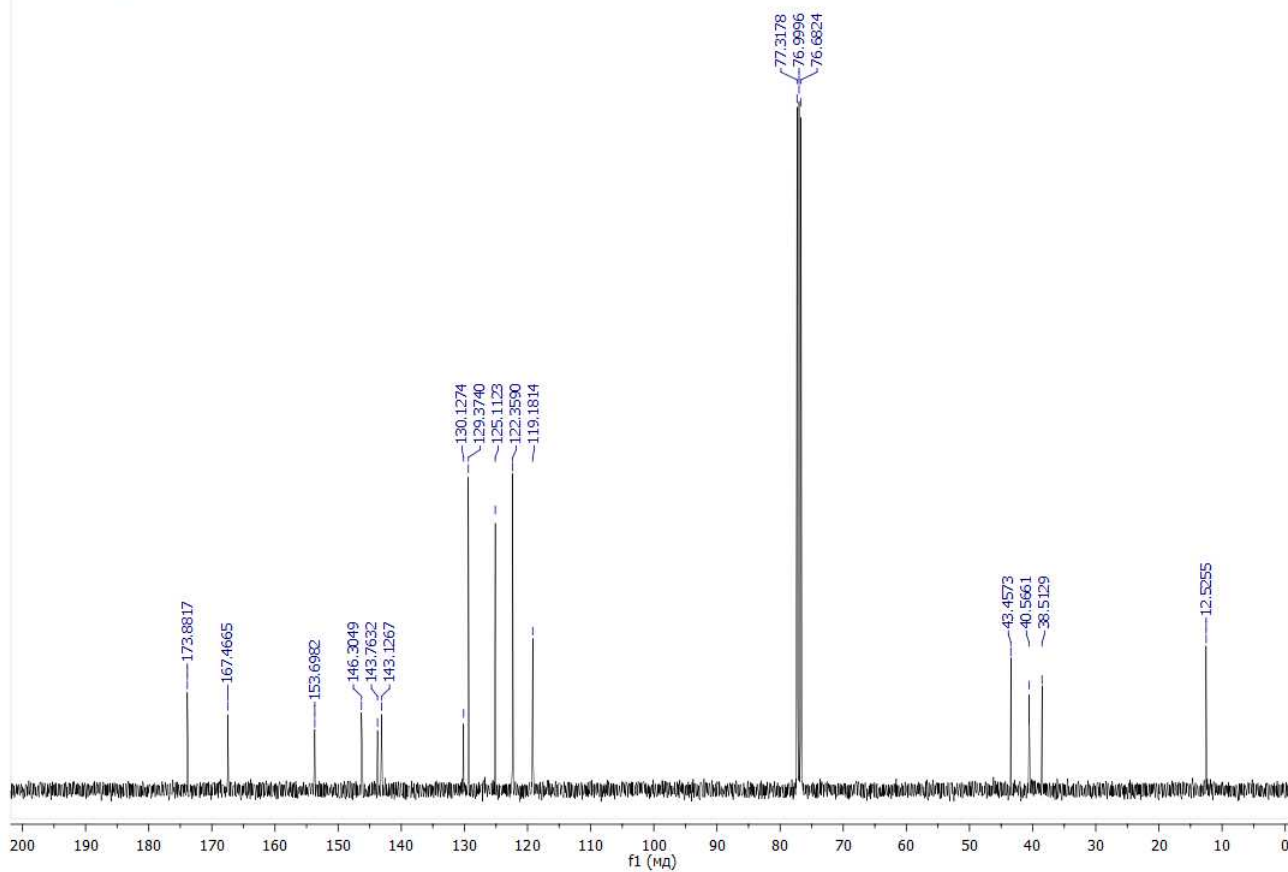
MAKc, 990, BF = 100.612769 MHz, Solvent - CDD3, 10 Jul 2013 T=298 K



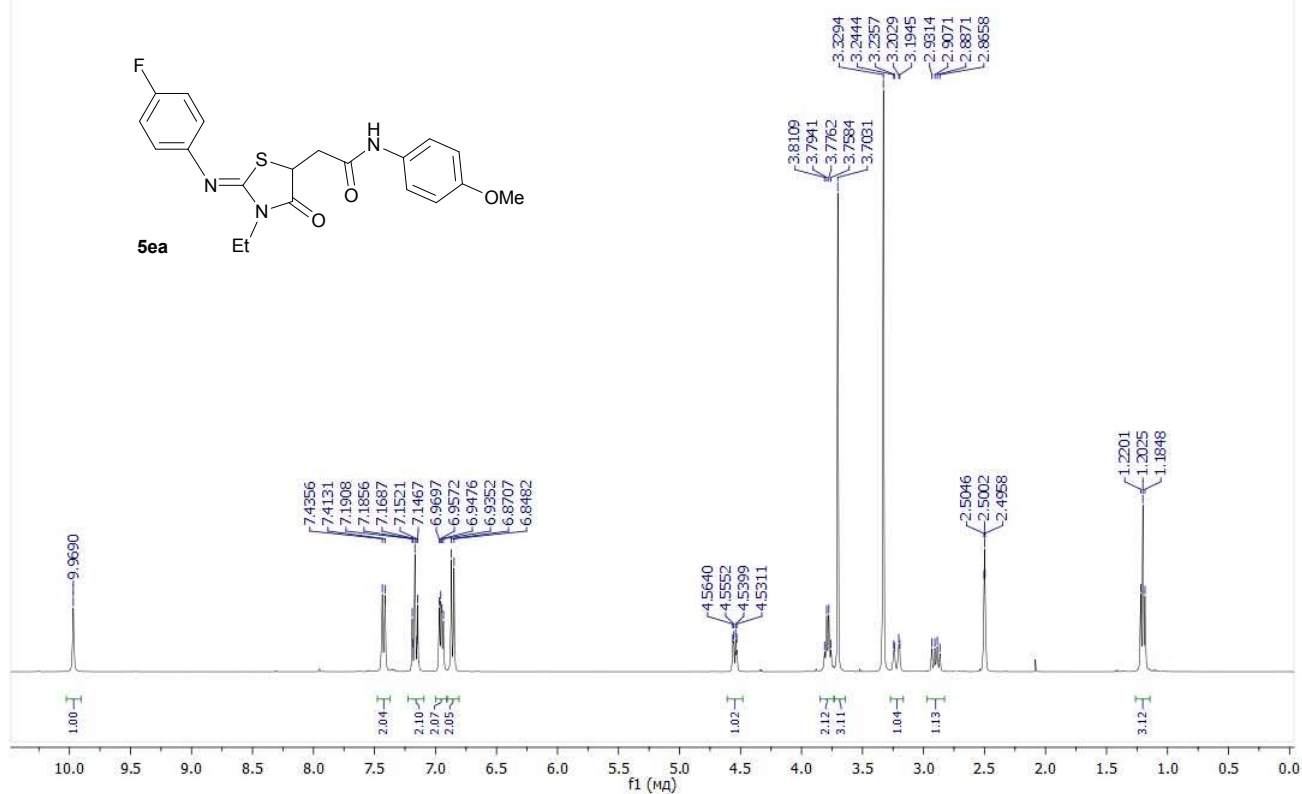
MAK, 778, BF = 400.13 MHz, Solvent - CDD3, 05 Apr 2013



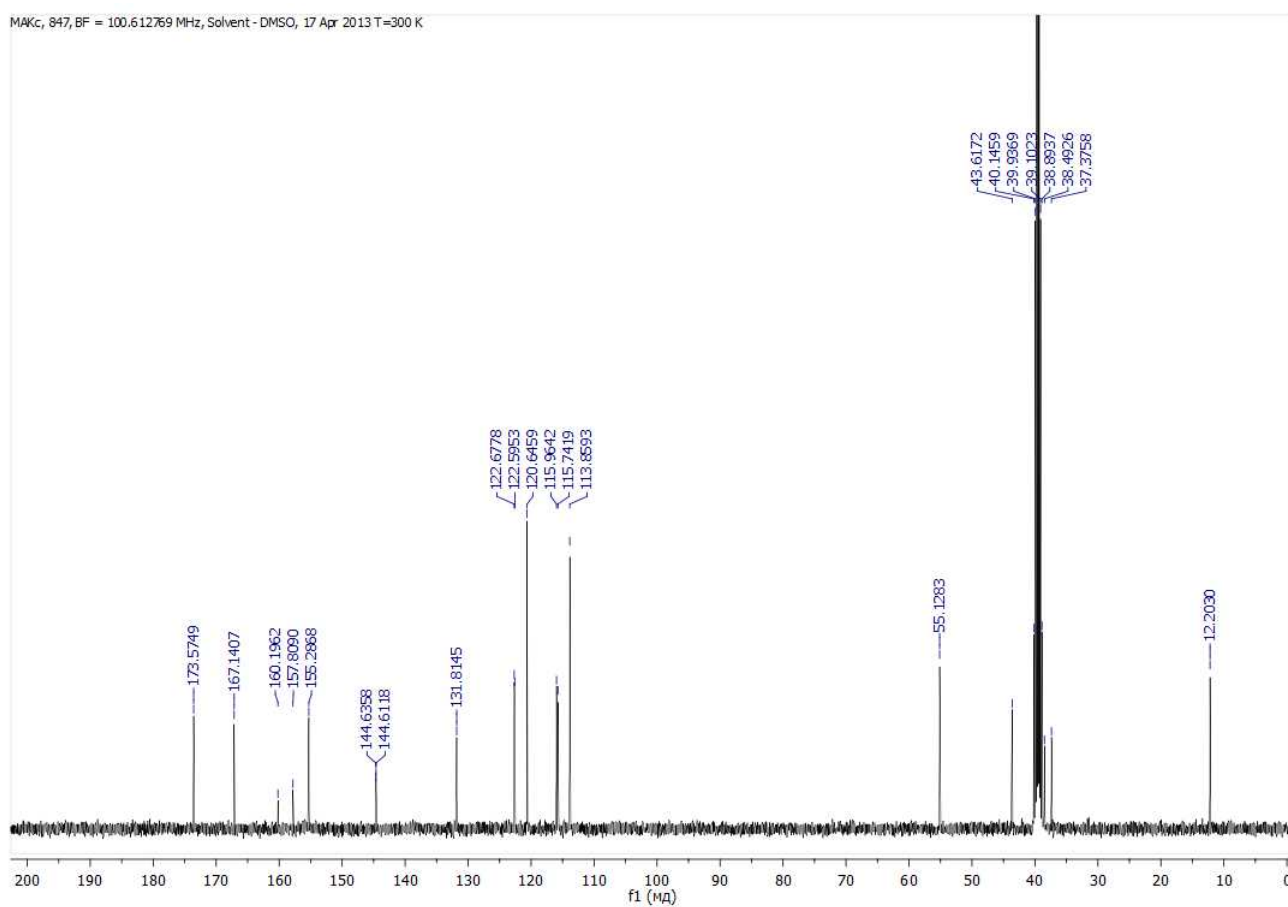
MAKc, 778, BF = 100.612769 MHz, Solvent - CDD3, 05 Apr 2013



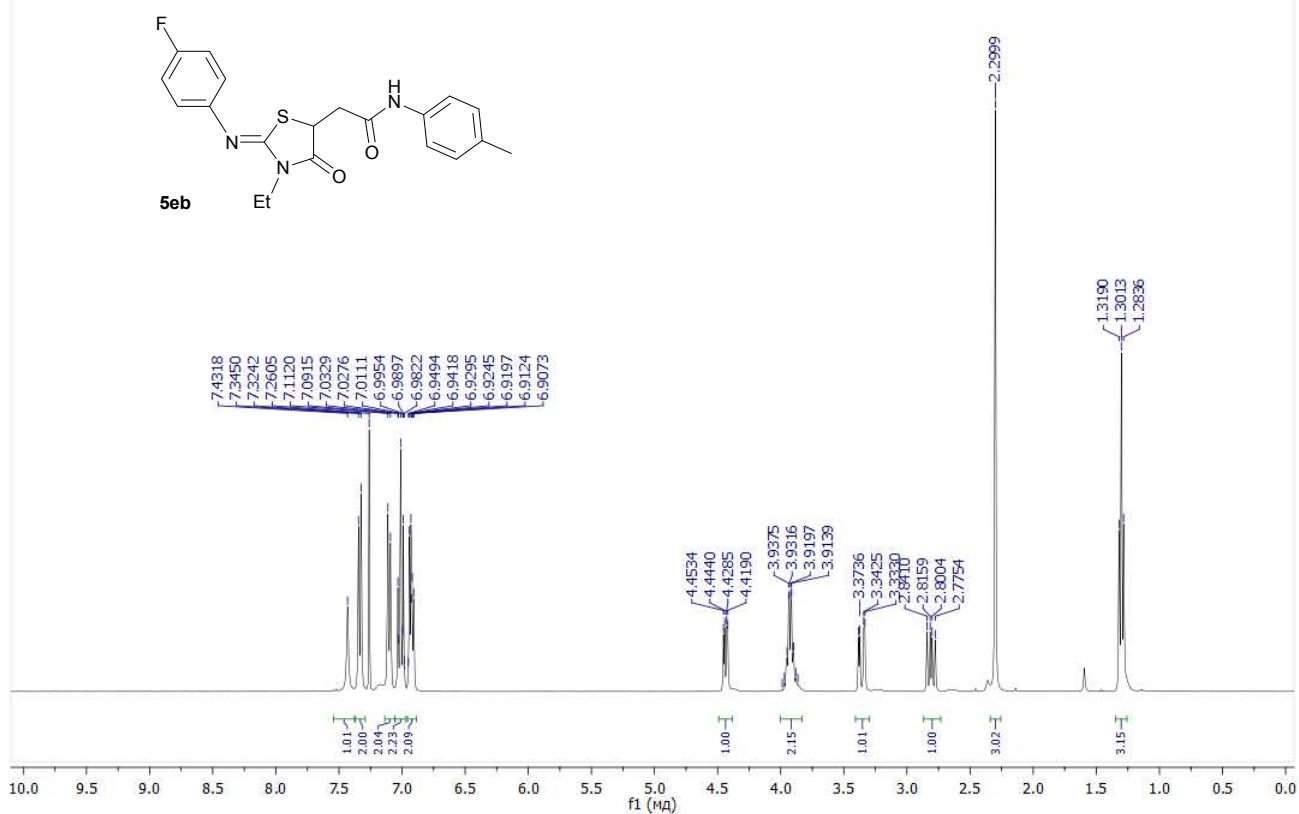
MAK, 847, BF = 400.13 MHz, Solvent - DMSO, 17 Apr 2013 T=300 K



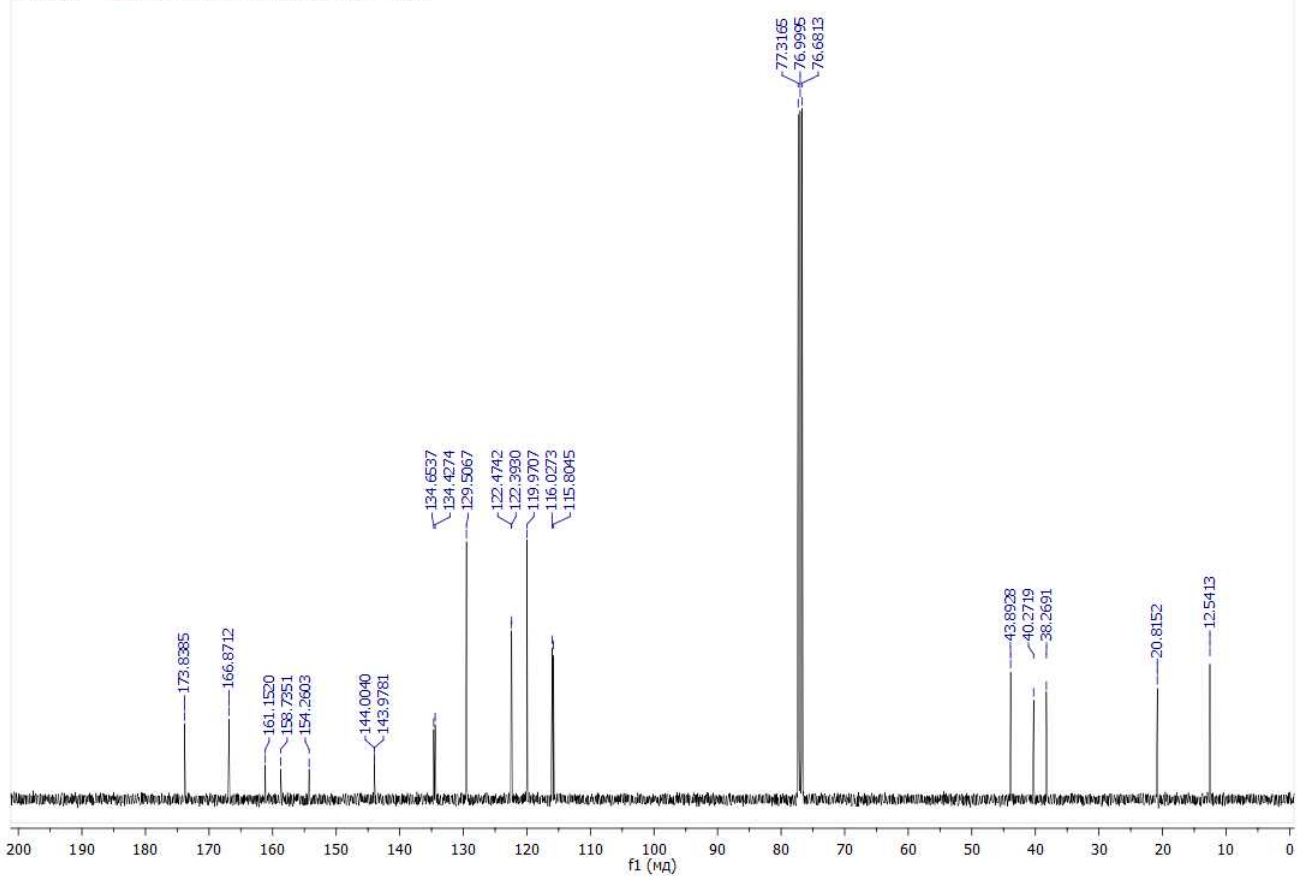
MAKc, 847, BF = 100.612769 MHz, Solvent - DMSO, 17 Apr 2013 T=300 K



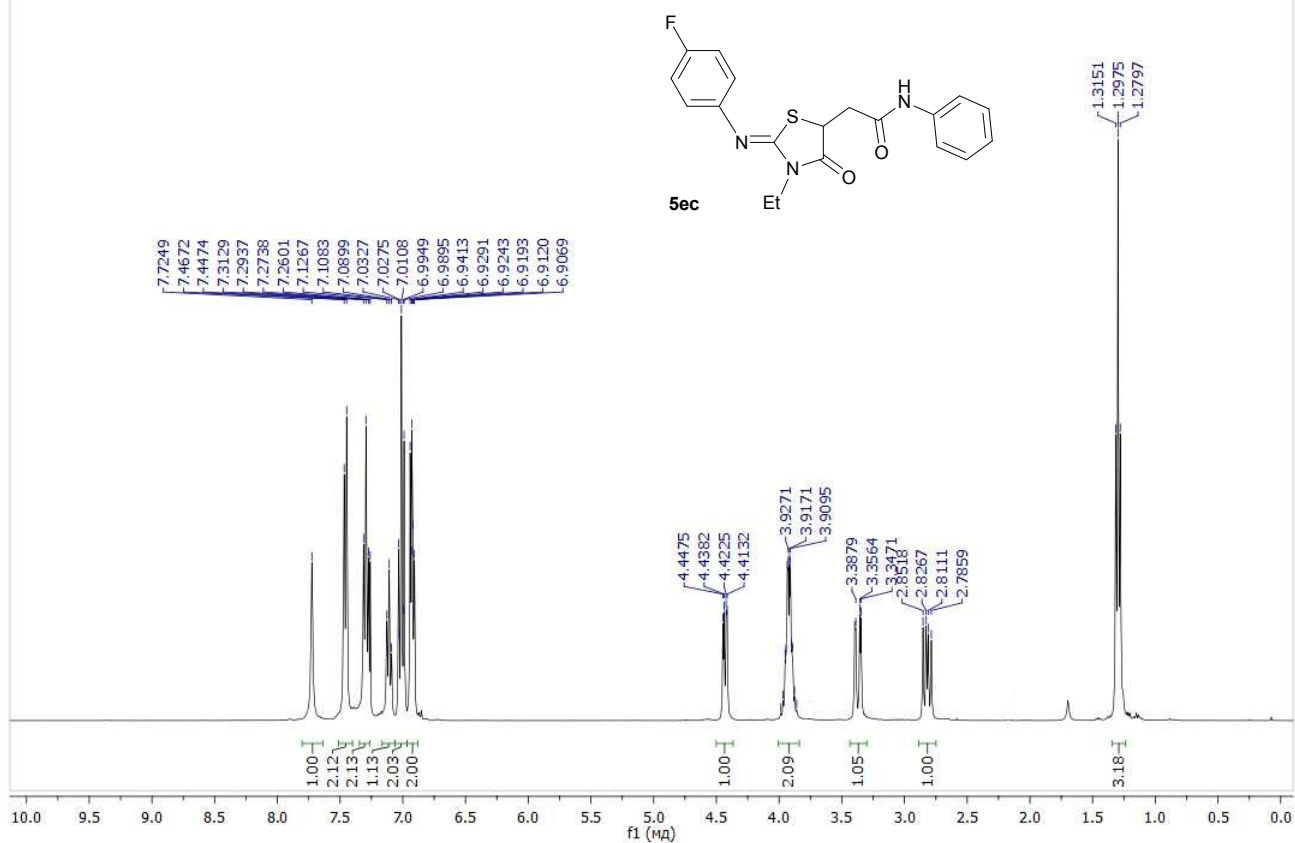
MAK, 185, BF = 400.13 MHz, Solvent - CDCl₃, 08 Nov 2013 T=298 K



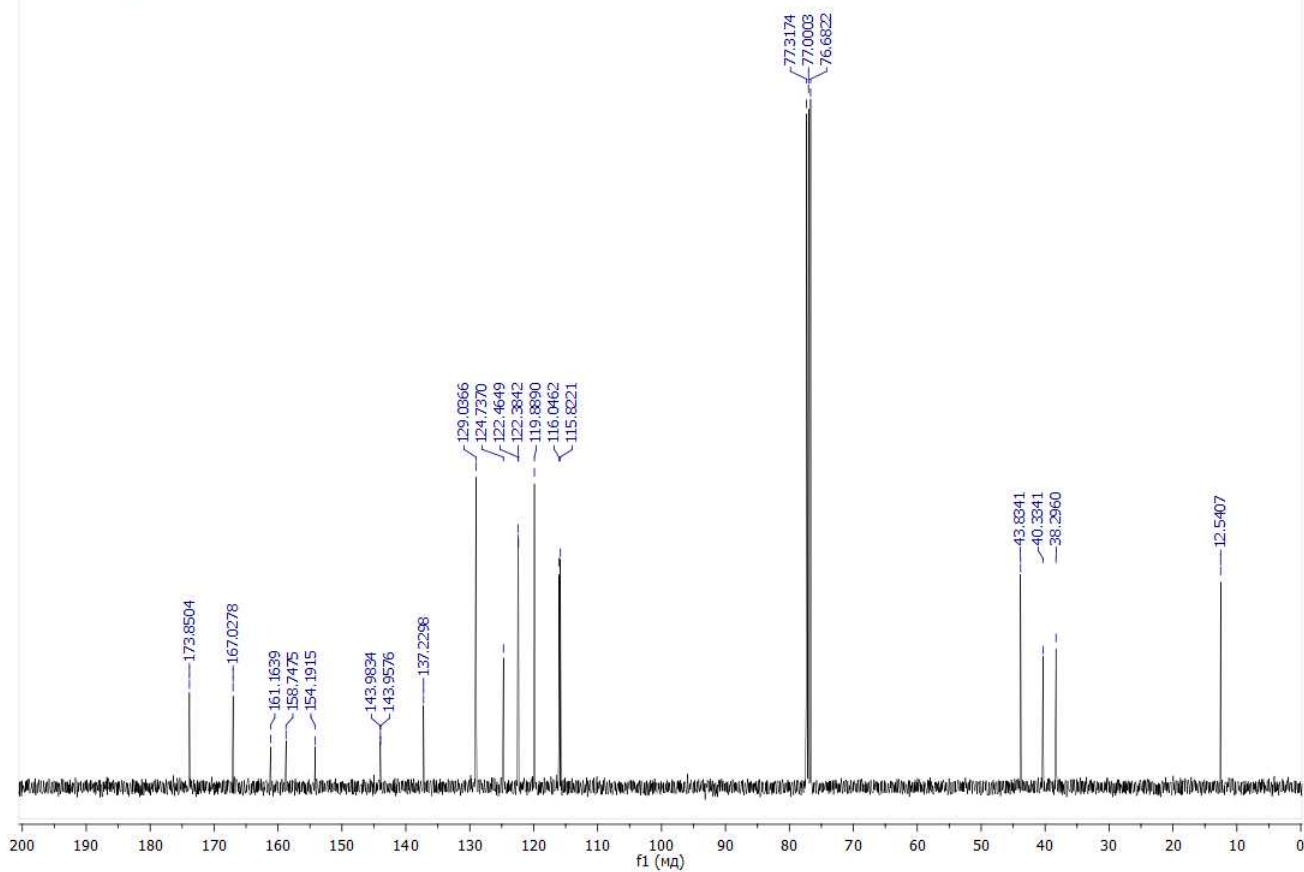
MAKc, 185, BF = 100.612769 MHz, Solvent - CDCl₃, 11 Nov 2013 T=298 K



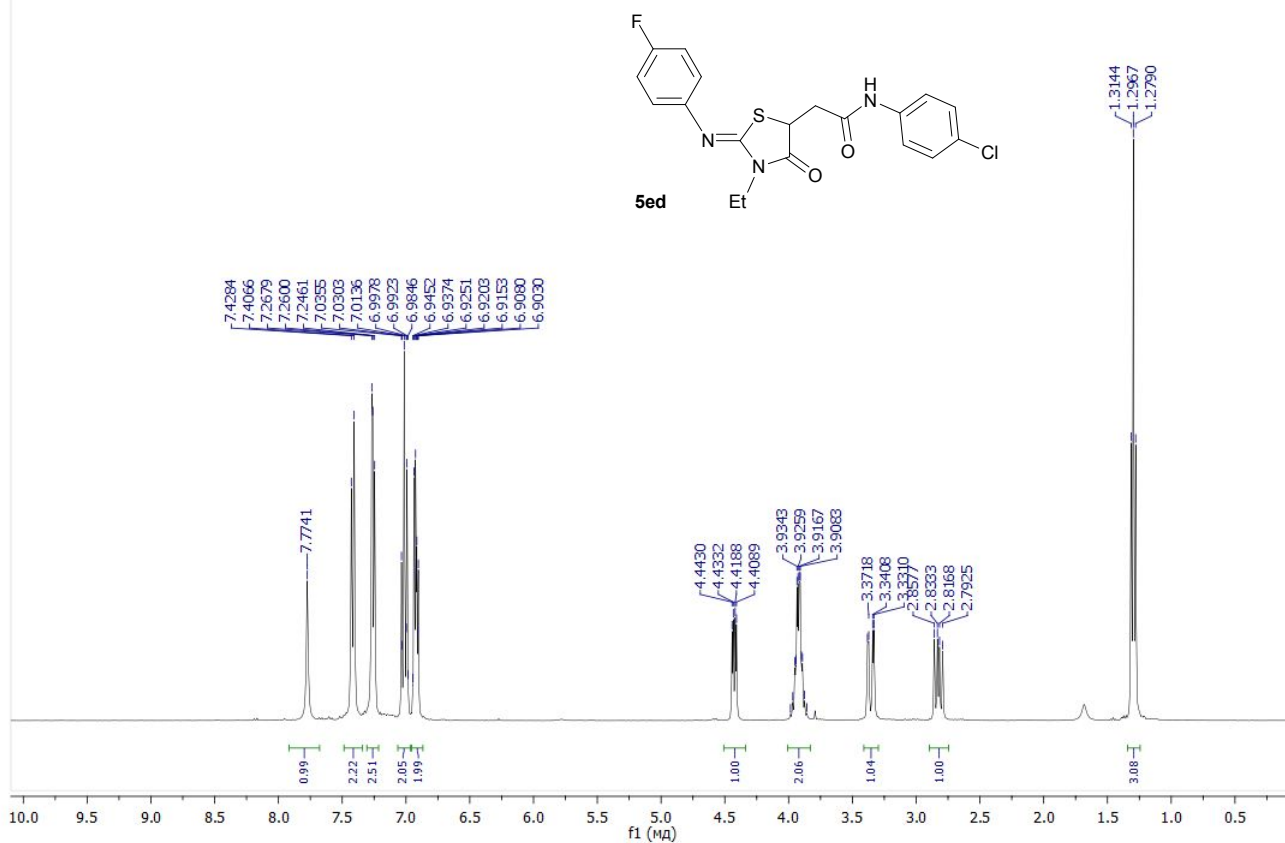
MAK, 776, BF = 400.13 MHz, Solvent - CDCl₃, 05 Apr 2013



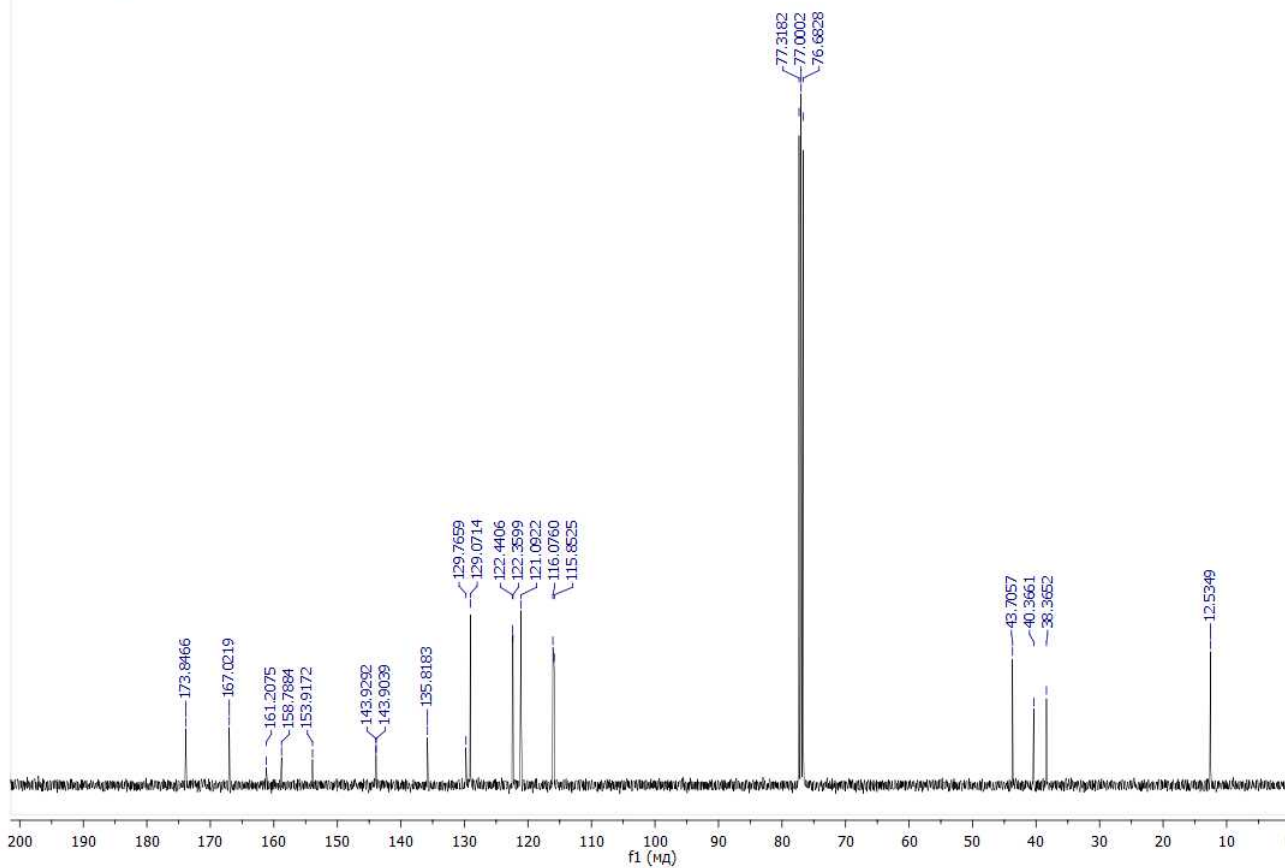
MAKc, 776, BF = 100.612769 MHz, Solvent - CDD3, 05 Apr 2013



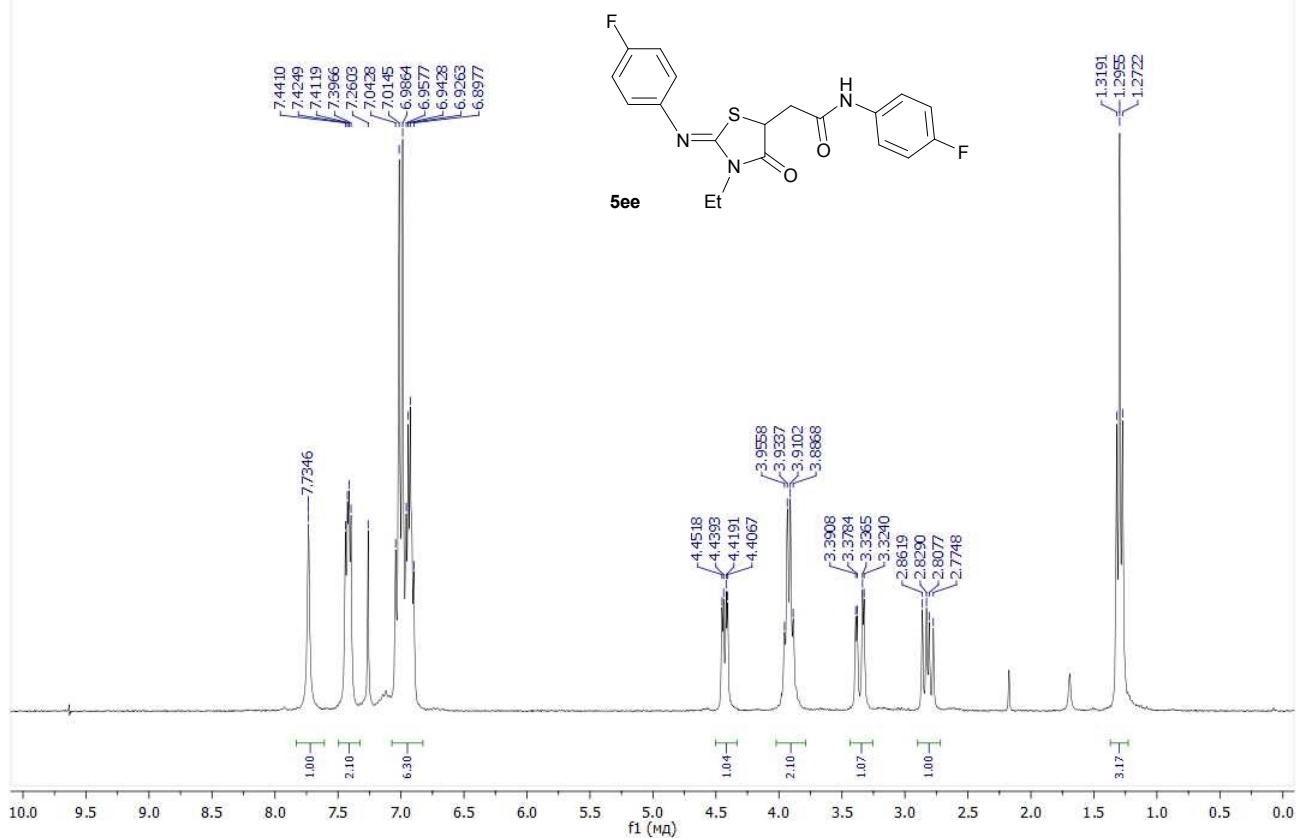
MAK, 840, BF = 400.13 MHz, Solvent - CDCl₃, 16 Apr 2013 T=300 K



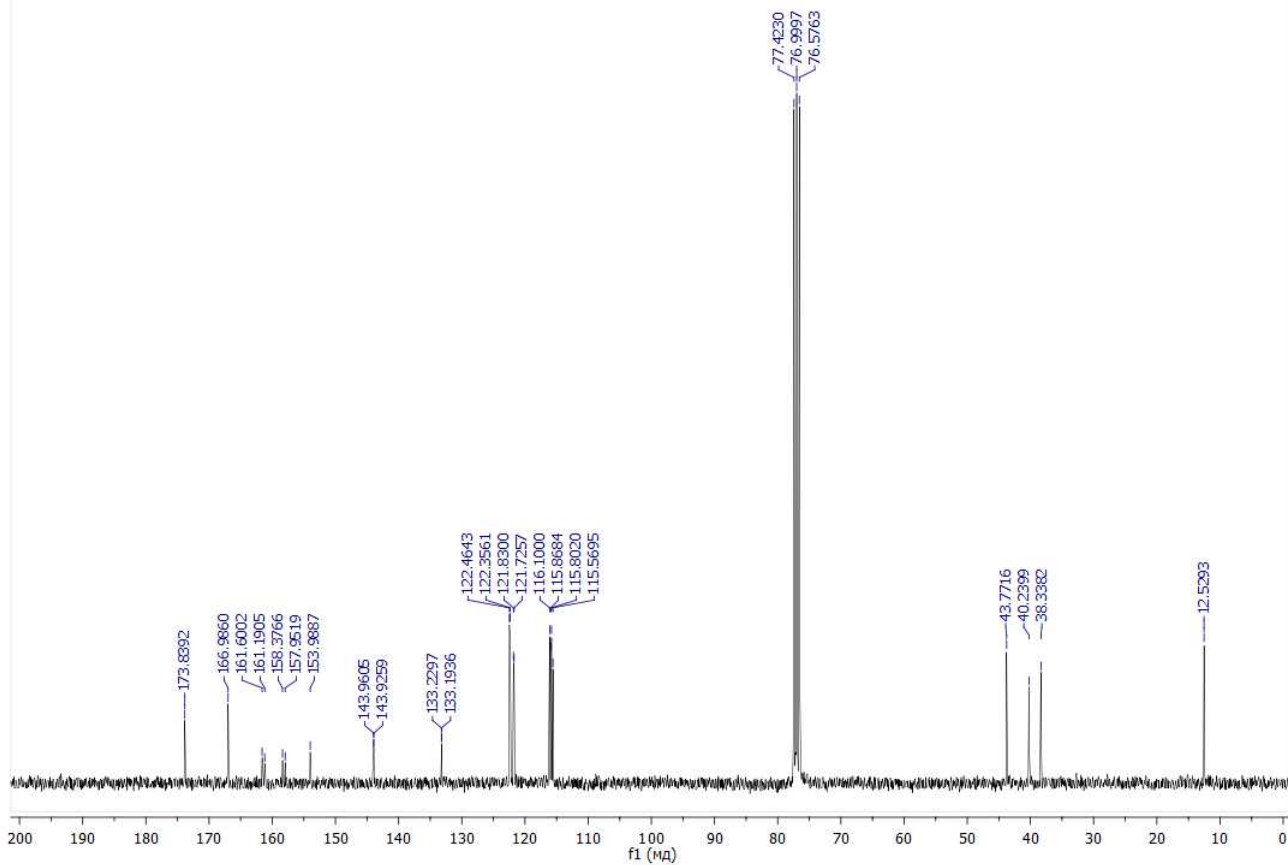
MAKc, 840, BF = 100.612769 MHz, Solvent - CDCl₃, 16 Apr 2013 T=301 K



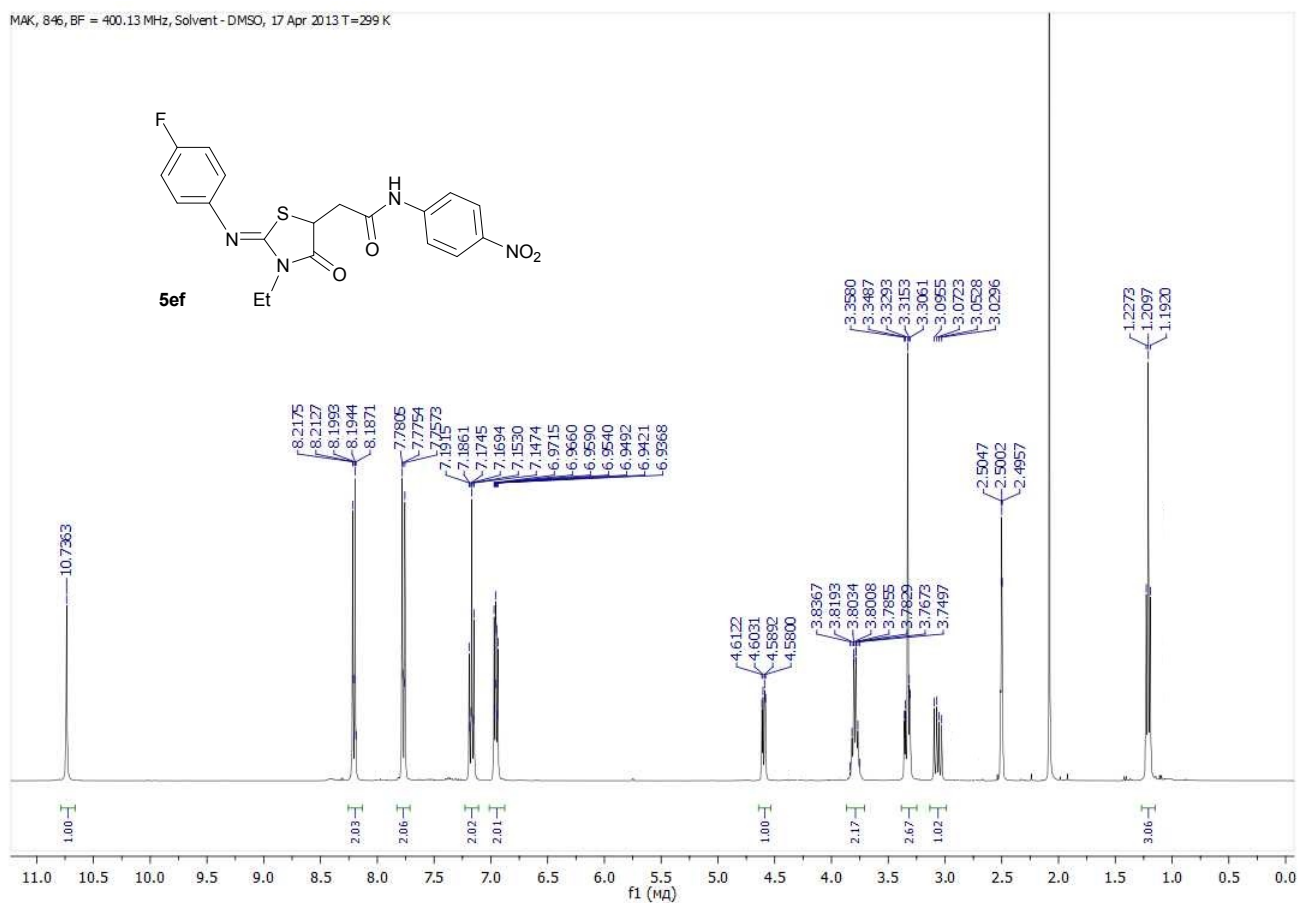
MAK, 771, BF = 300.13 MHz, SOLVENT - CDCl₃, 4 Apr 2013



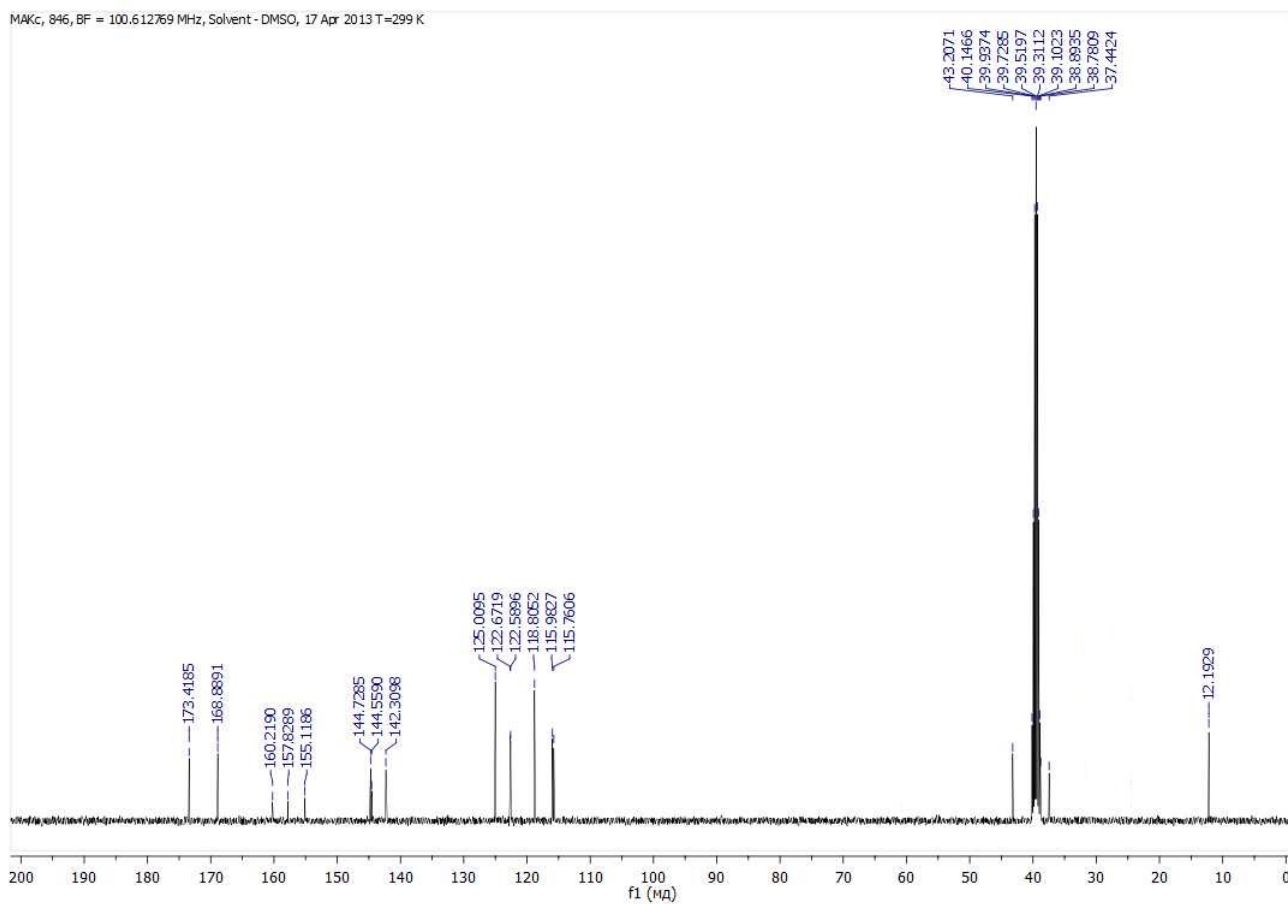
MAK, 771, BF = 75.47 MHz, SOLVENT - CDCl₃, 4 Apr 2013



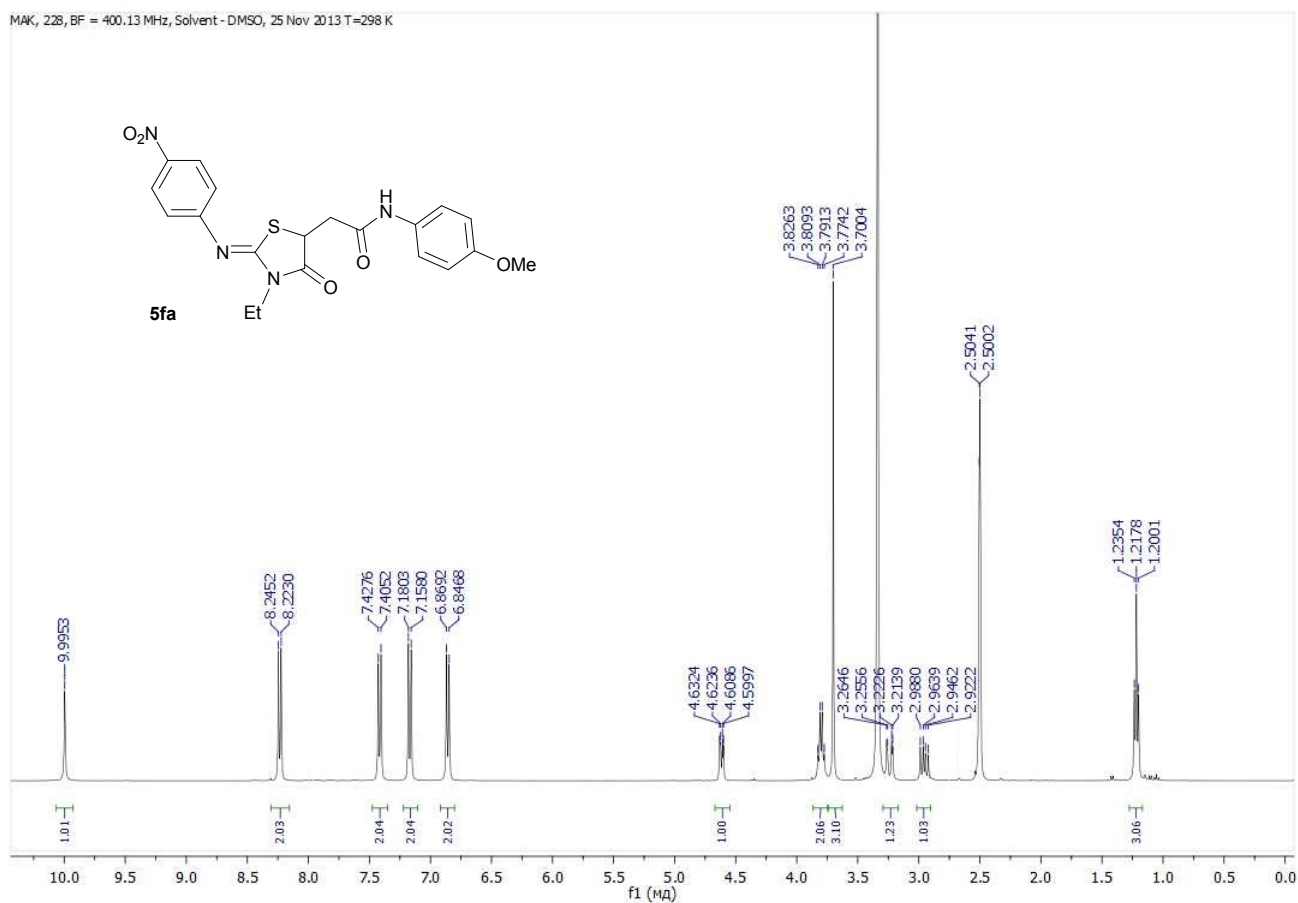
MAK, 846, BF = 400.13 MHz, Solvent - DMSO, 17 Apr 2013 T=299 K



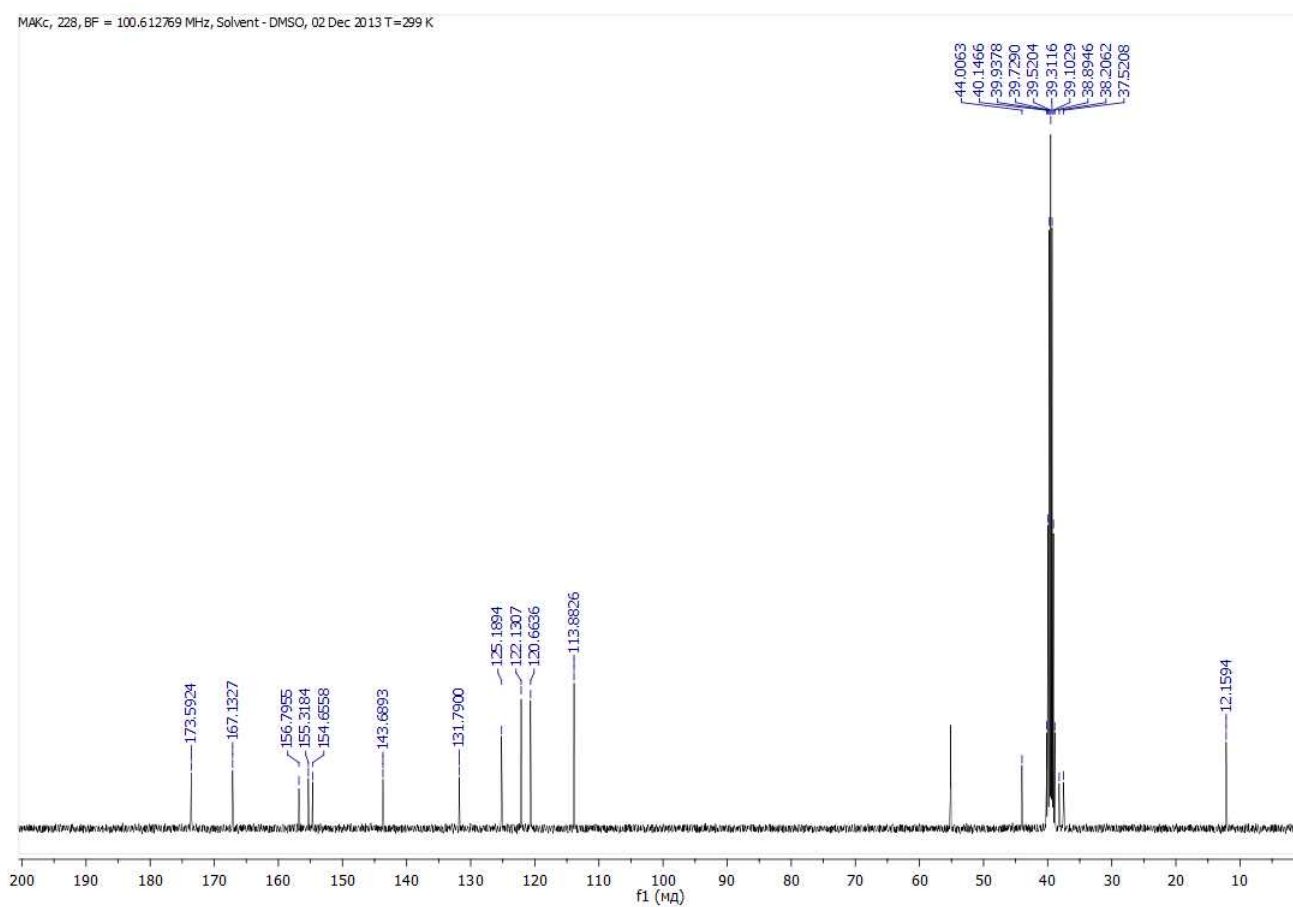
MAKc, 846, BF = 100.612769 MHz, Solvent - DMSO, 17 Apr 2013 T=299 K



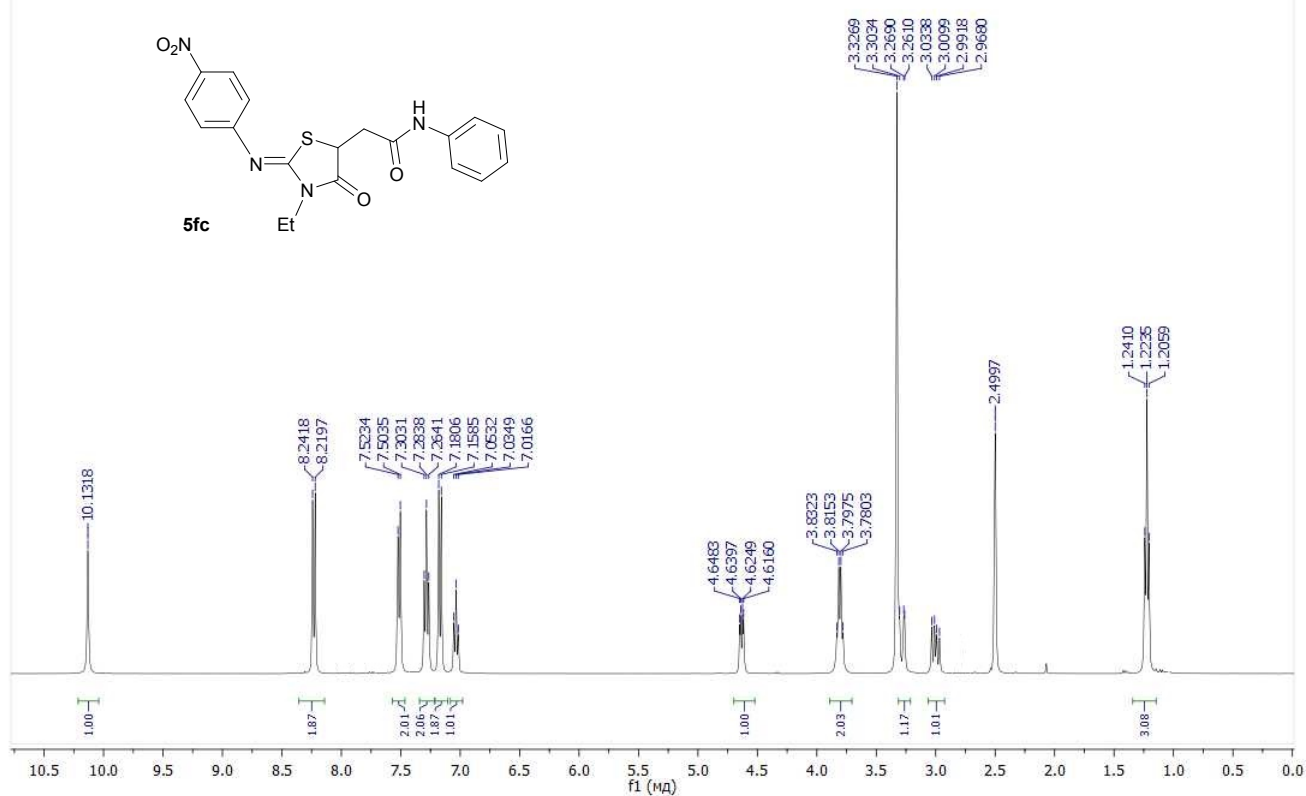
MAK, 228, BF = 400.13 MHz, Solvent - DMSO, 25 Nov 2013 T=298 K



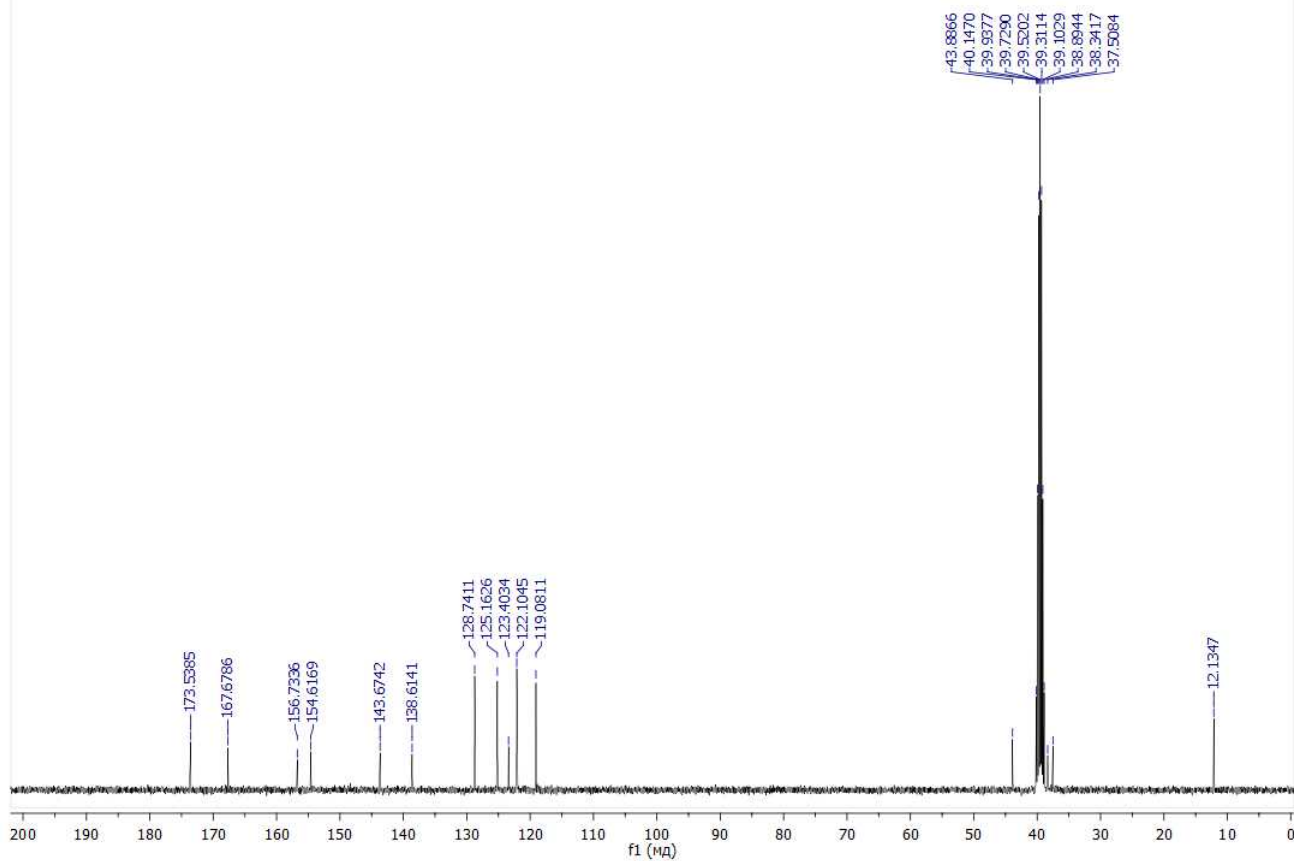
MAKc, 228, BF = 100.612769 MHz, Solvent - DMSO, 02 Dec 2013 T=299 K



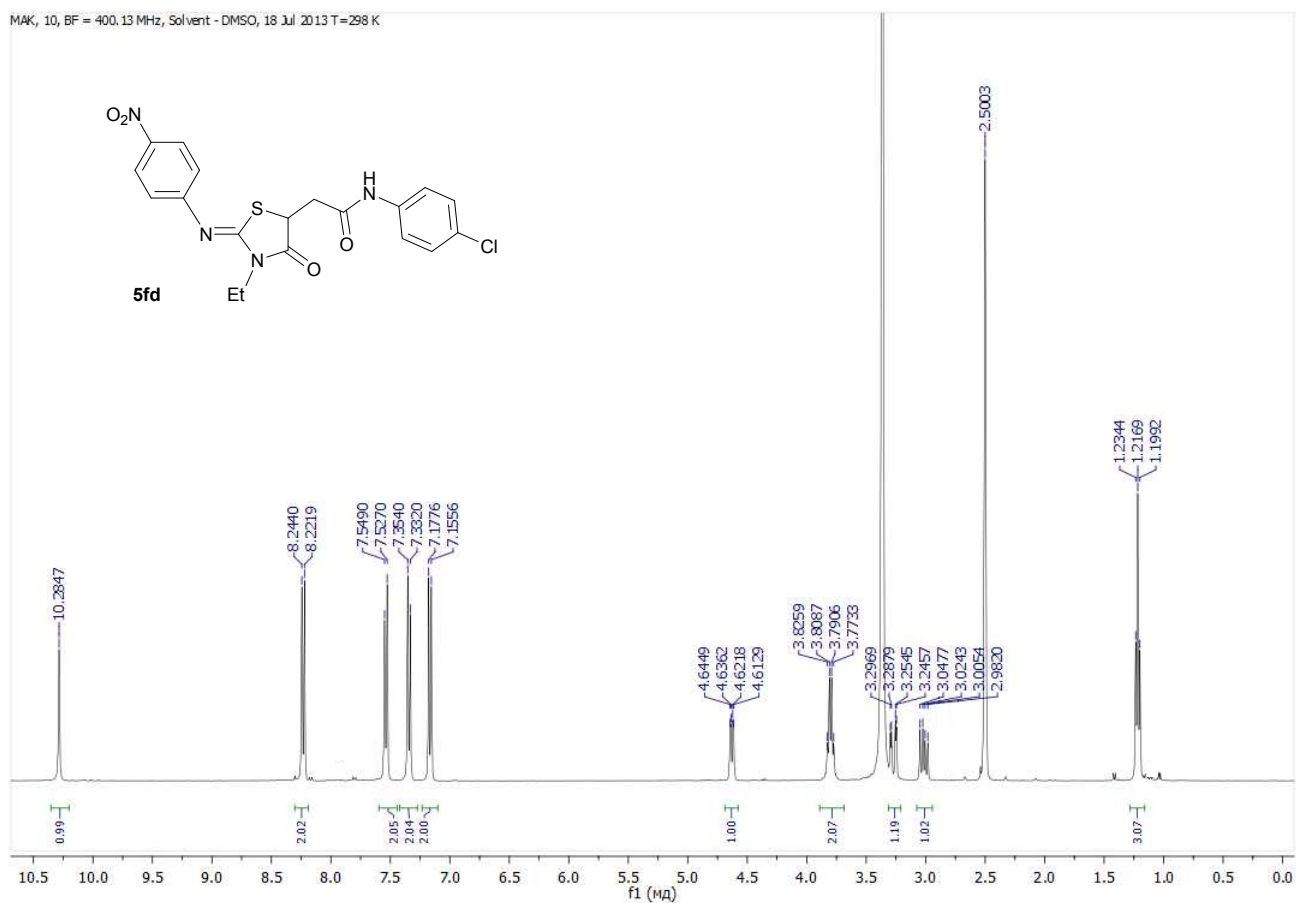
MAK, 833, BF = 400.13 MHz, Solvent - DMSO, 15 Apr 2013 T=300 K



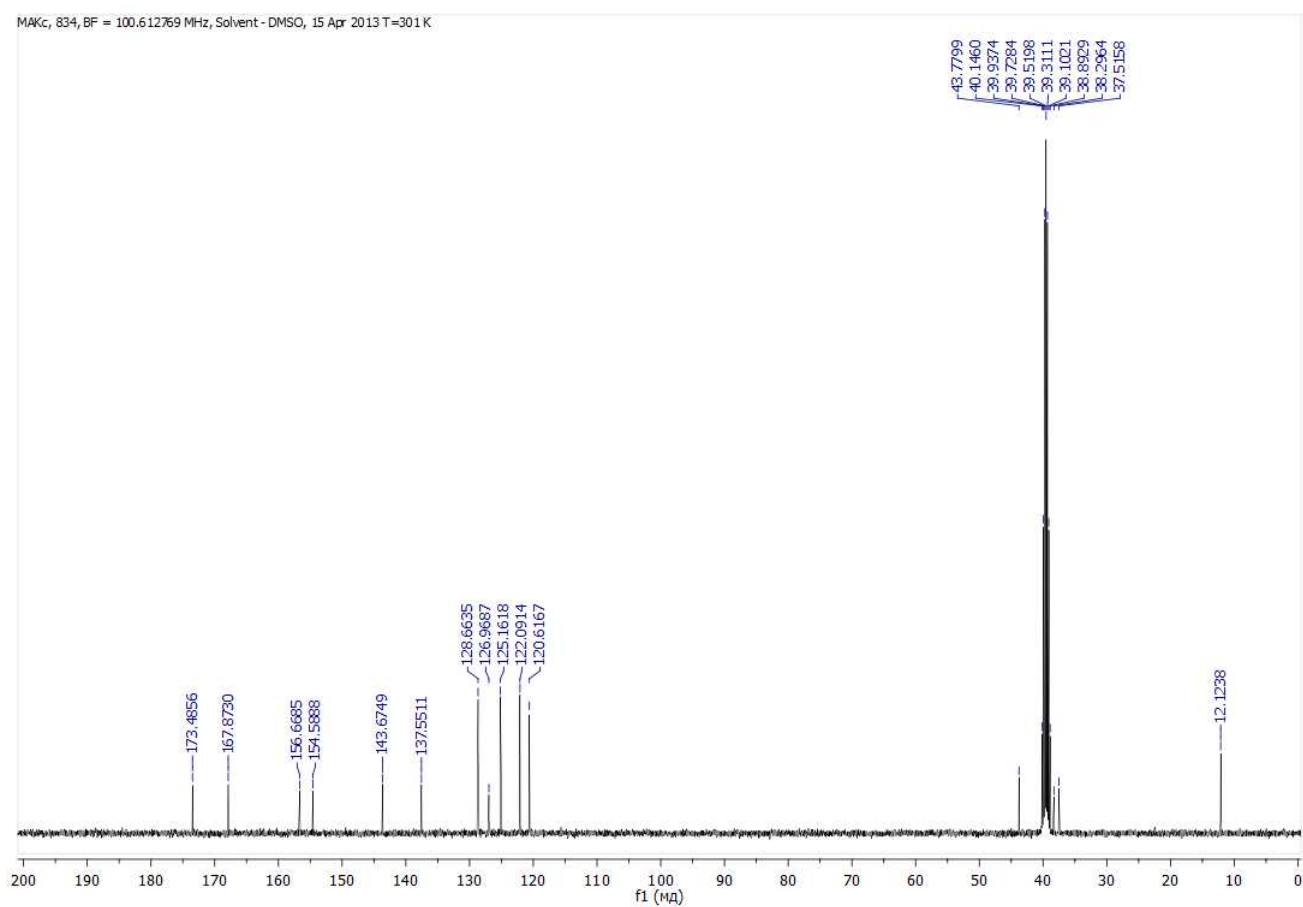
MAKc, 833, BF = 100.612769 MHz, Solvent - DMSO, 15 Apr 2013 T=301 K



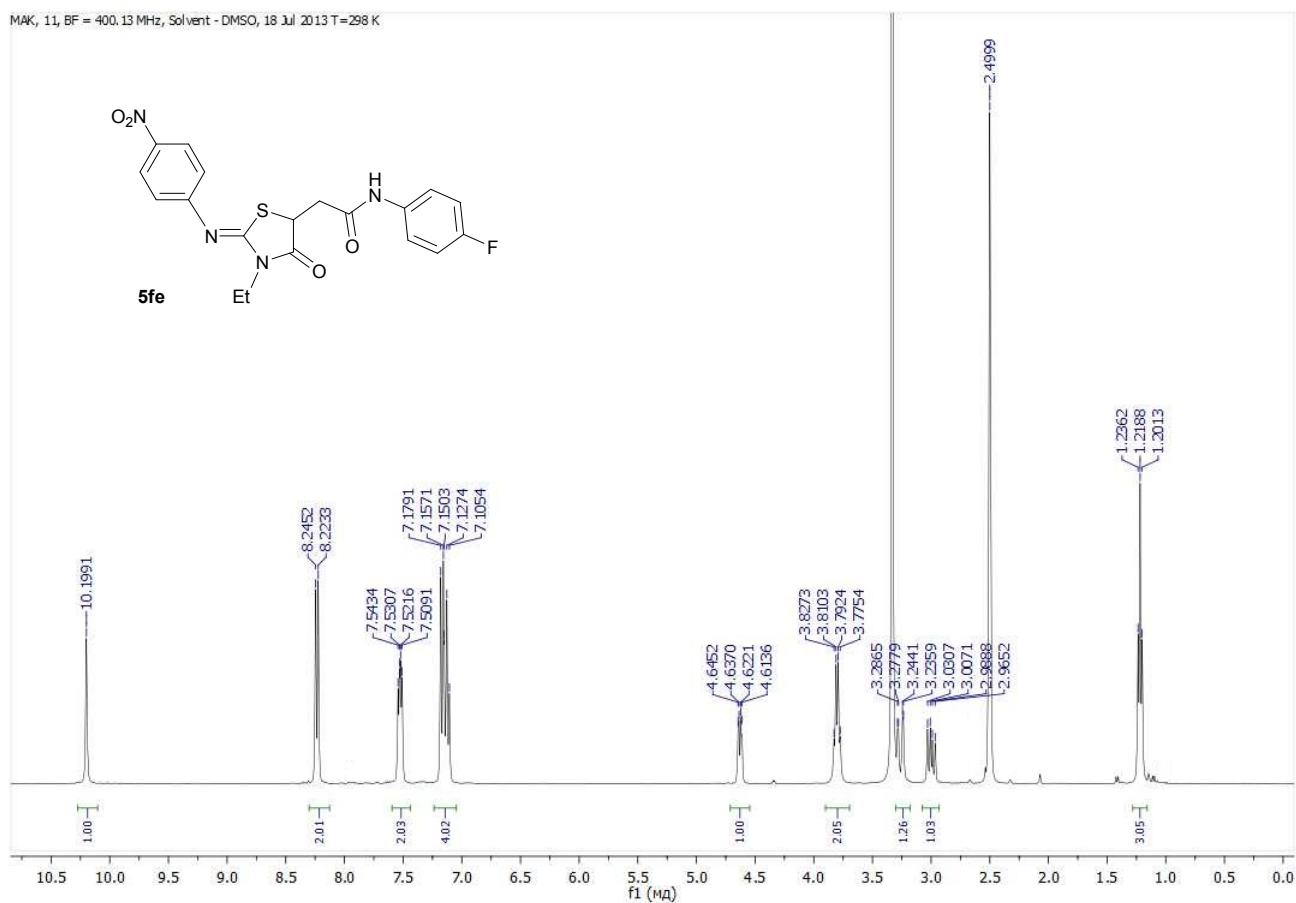
MAK, 10, BF = 400.13 MHz, Solvent - DMSO, 18 Jul 2013 T=298 K



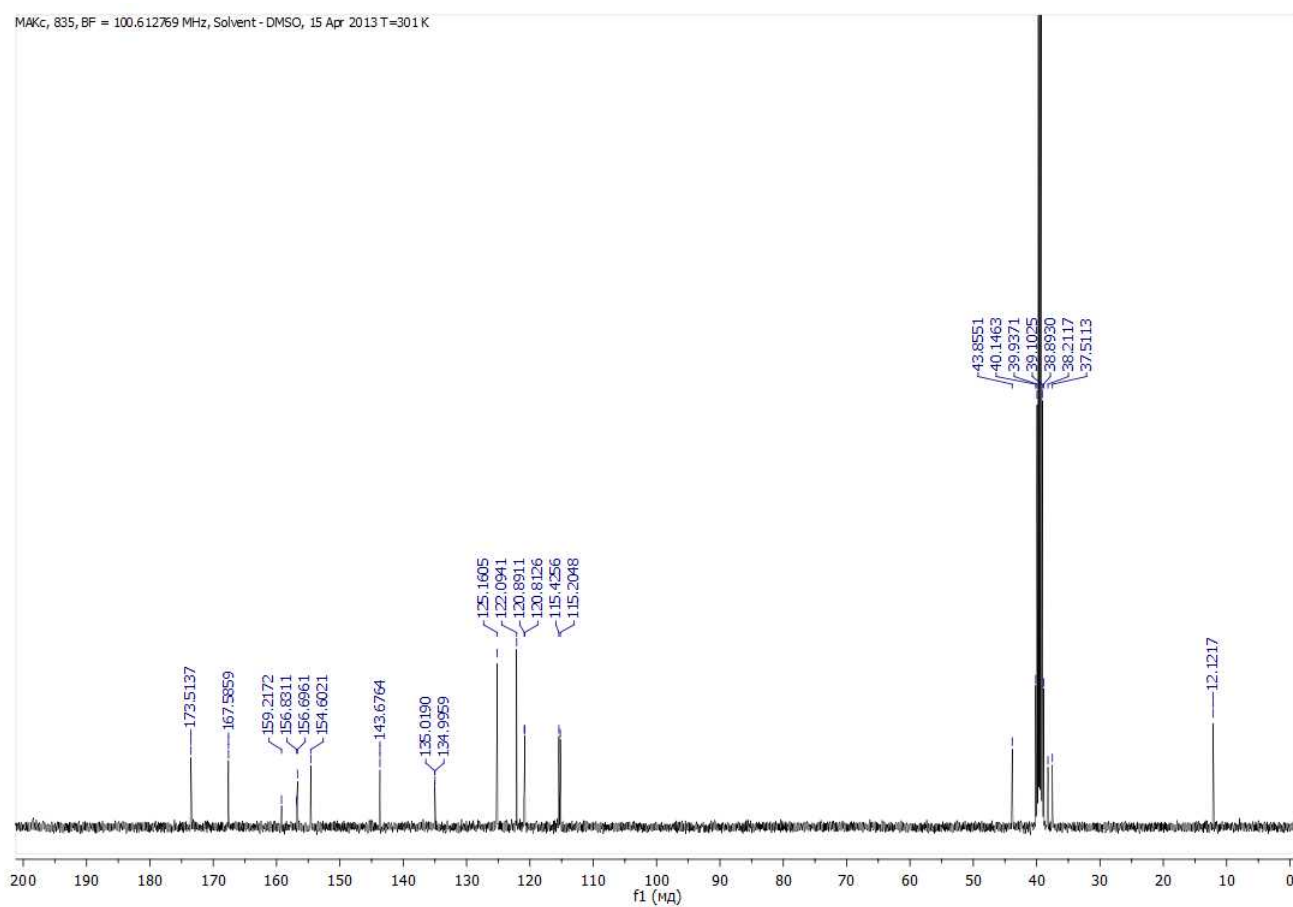
MAKc, 834, BF = 100.612769 MHz, Solvent - DMSO, 15 Apr 2013 T=301 K



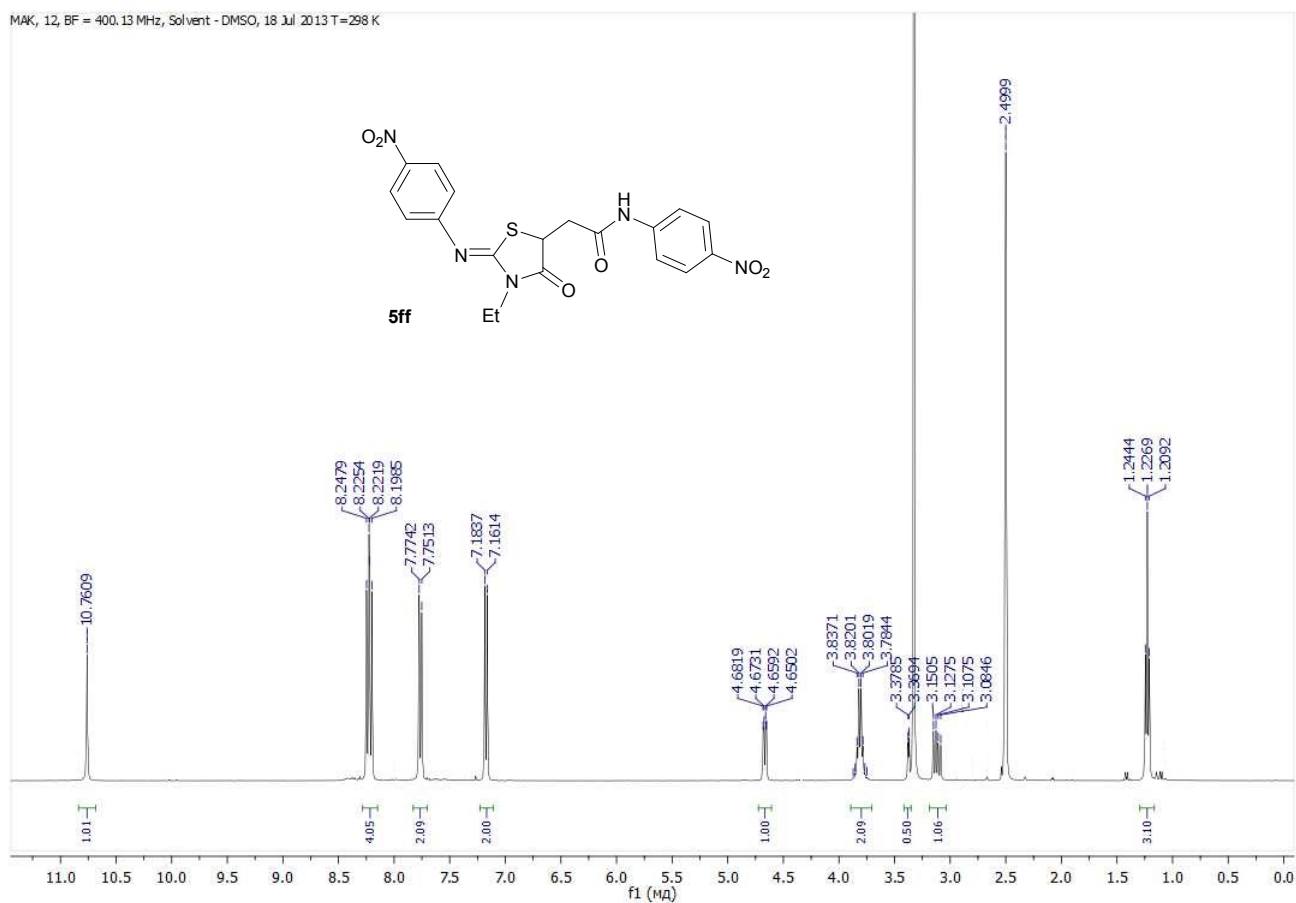
MAK, 11, BF = 400.13 MHz, Solvent - DMSO, 18 Jul 2013 T=298 K



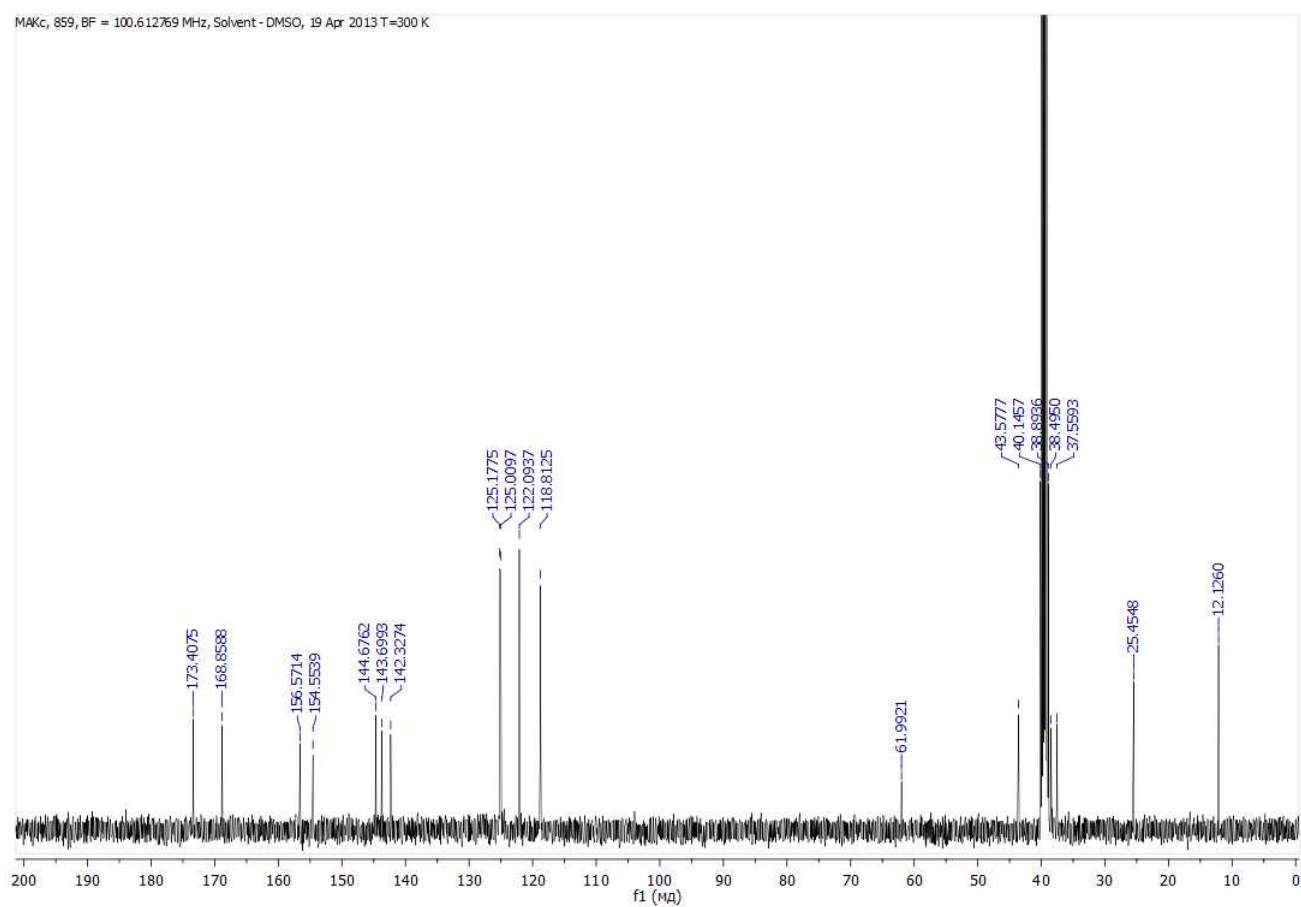
MAKc, 835, BF = 100.612769 MHz, Solvent - DMSO, 15 Apr 2013 T=301 K



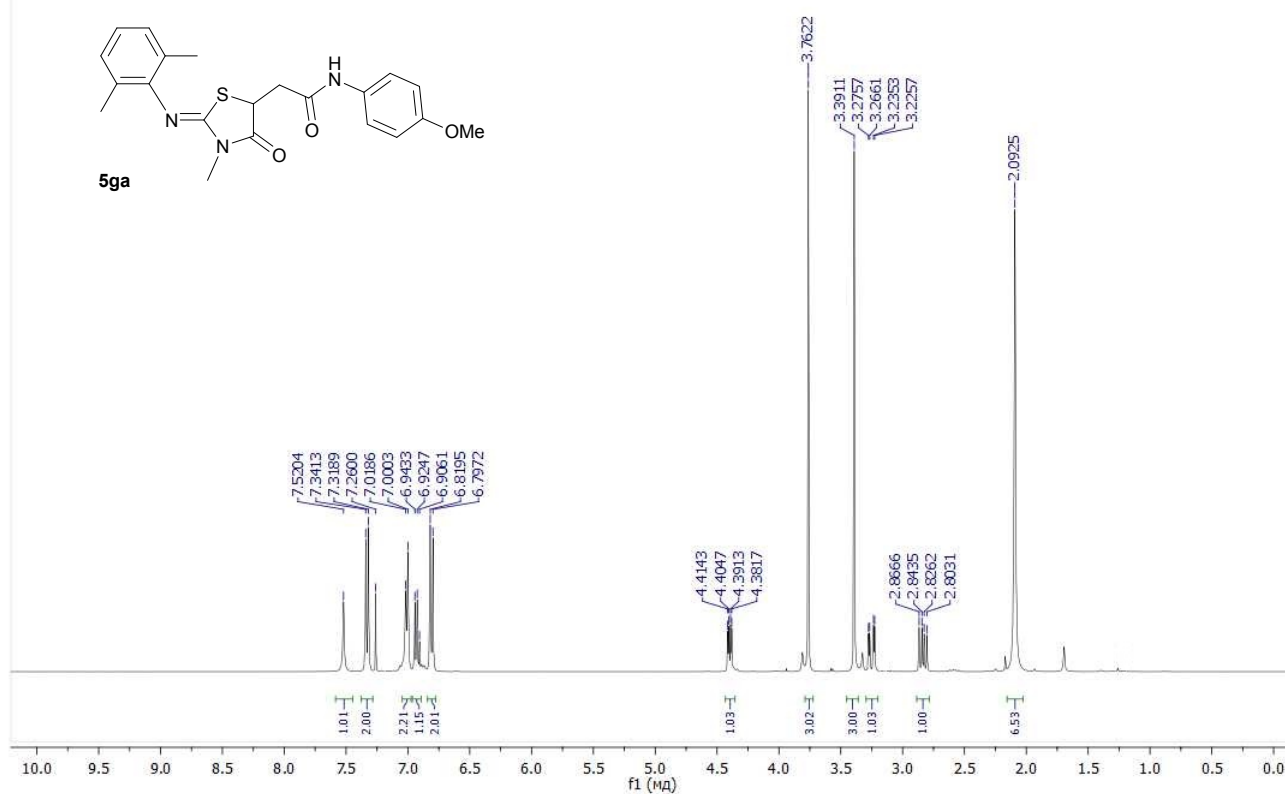
MAK, 12, BF = 400.13 MHz, Solvent - DMSO, 18 Jul 2013 T=298 K



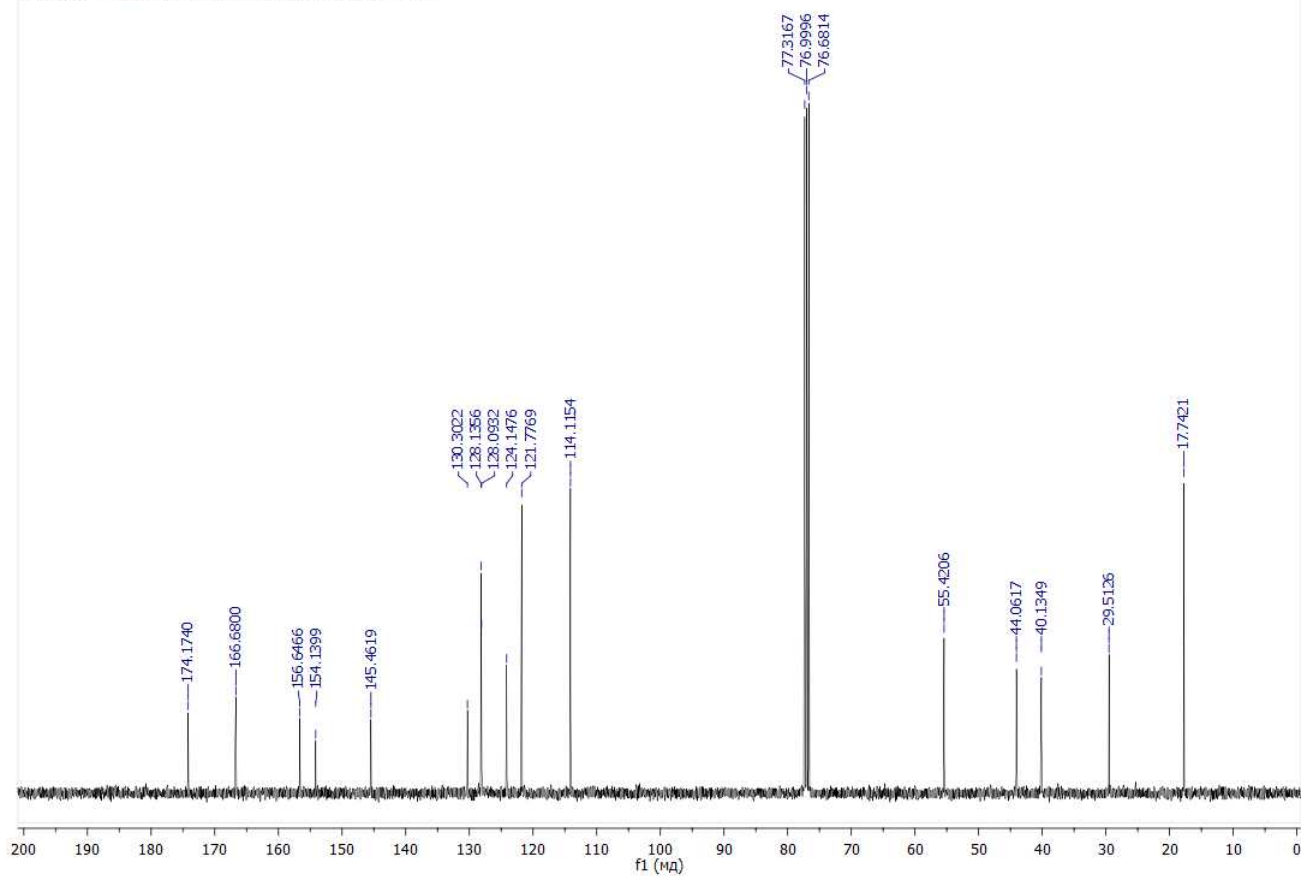
MAKc, 859, BF = 100.612769 MHz, Solvent - DMSO, 19 Apr 2013 T=300 K



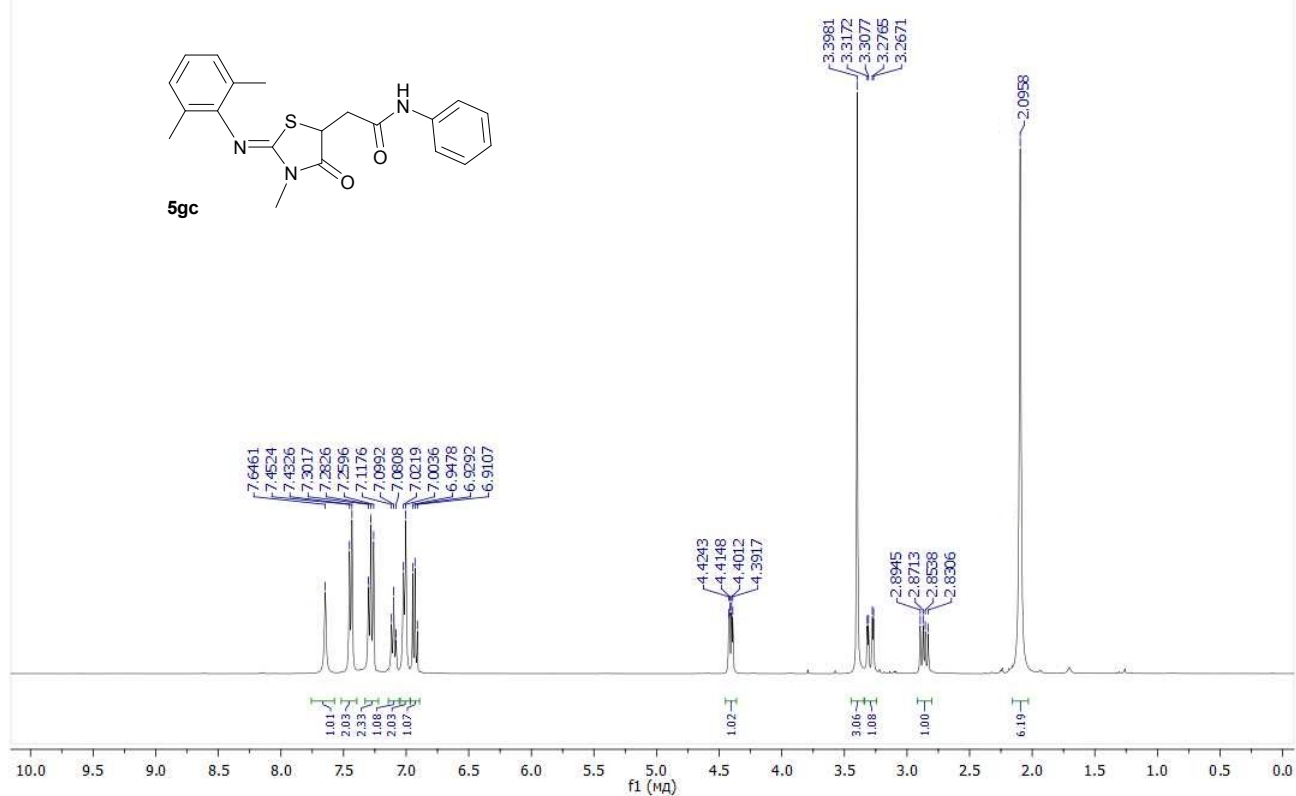
MAK, 839, BF = 400.13 MHz, Solvent - CDCl₃, 16 Apr 2013 T=300 K



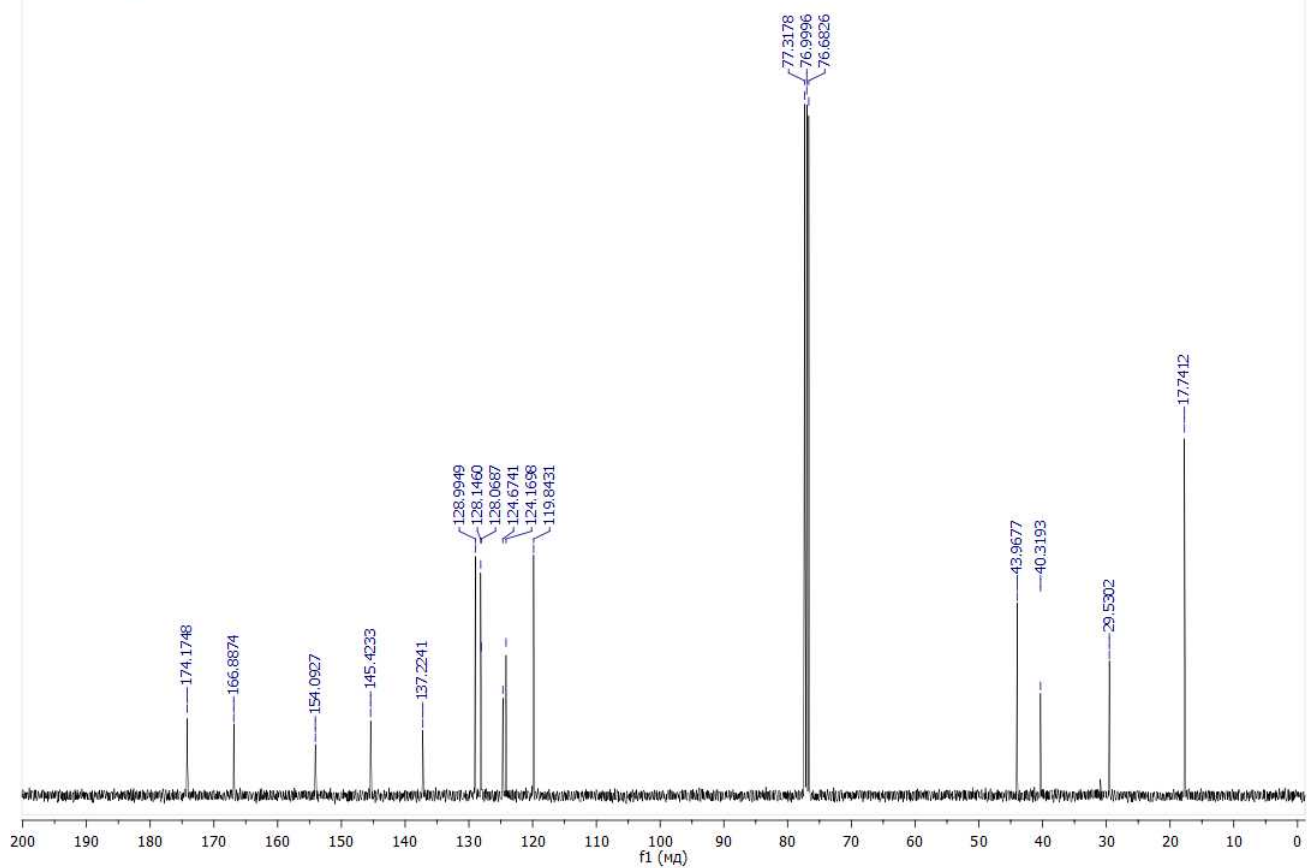
MAKc, 839, BF = 100.612769 MHz, Solvent - CDCl₃, 16 Apr 2013 T=301 K



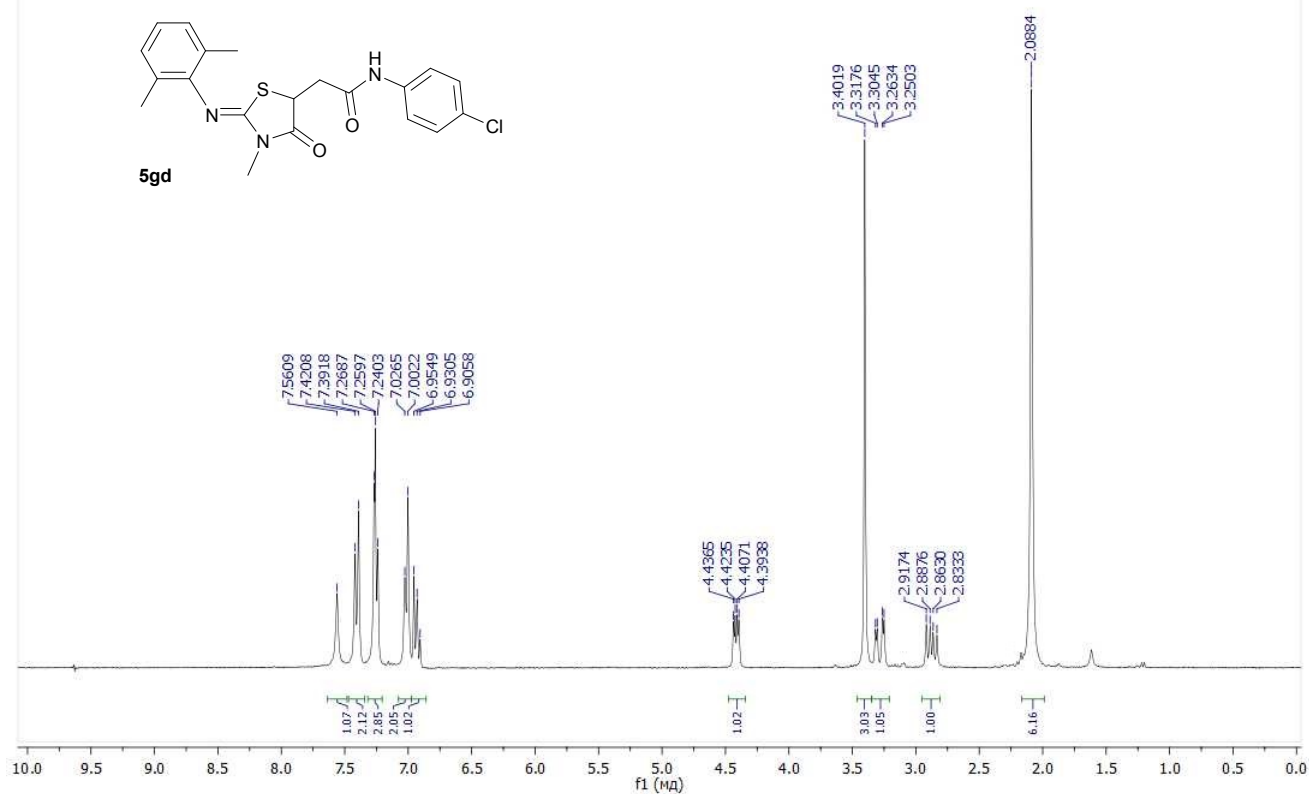
MAK, 773, BF = 400.13 MHz, Solvent - CDD3, 04 Apr 2013



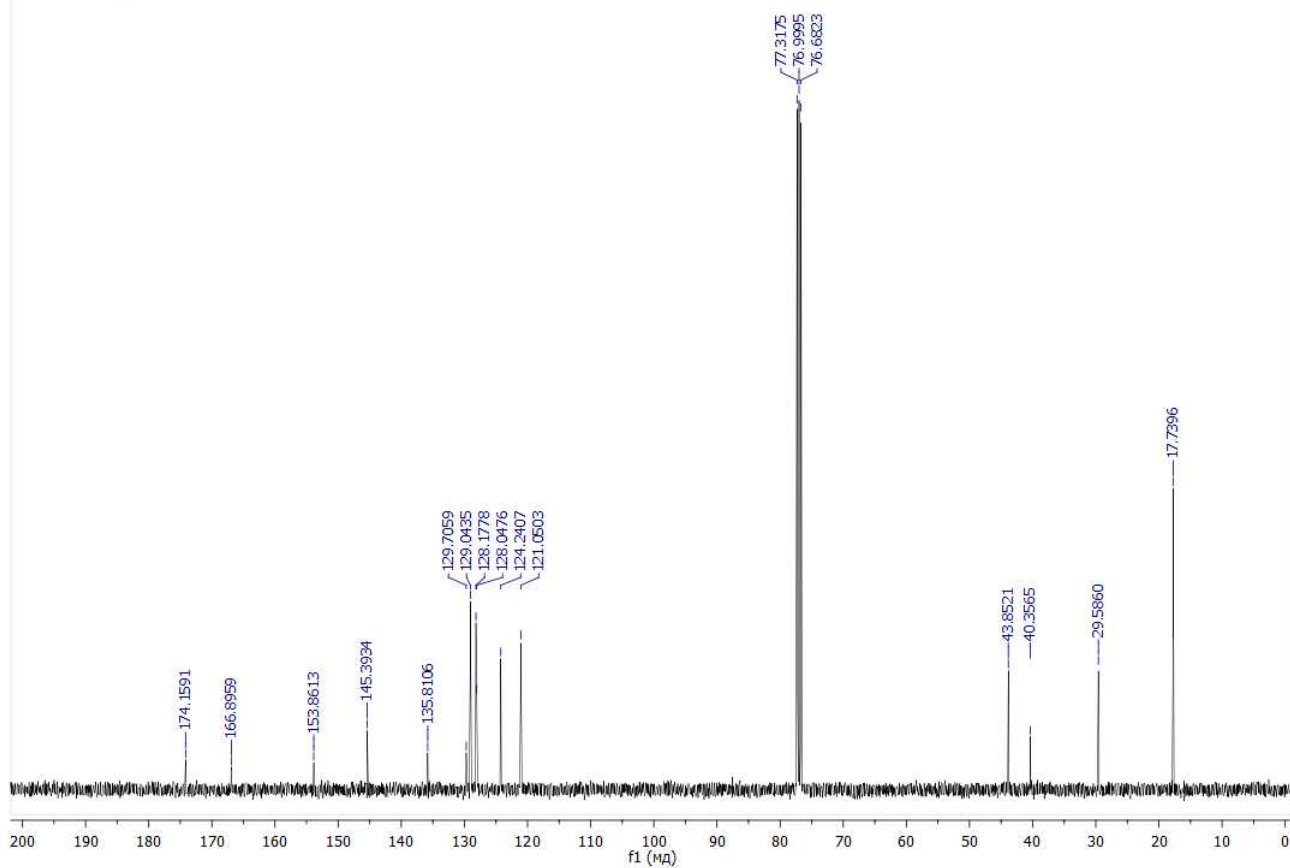
MAKc, 773, BF = 100.612769 MHz, Solvent - CDD3, 04 Apr 2013



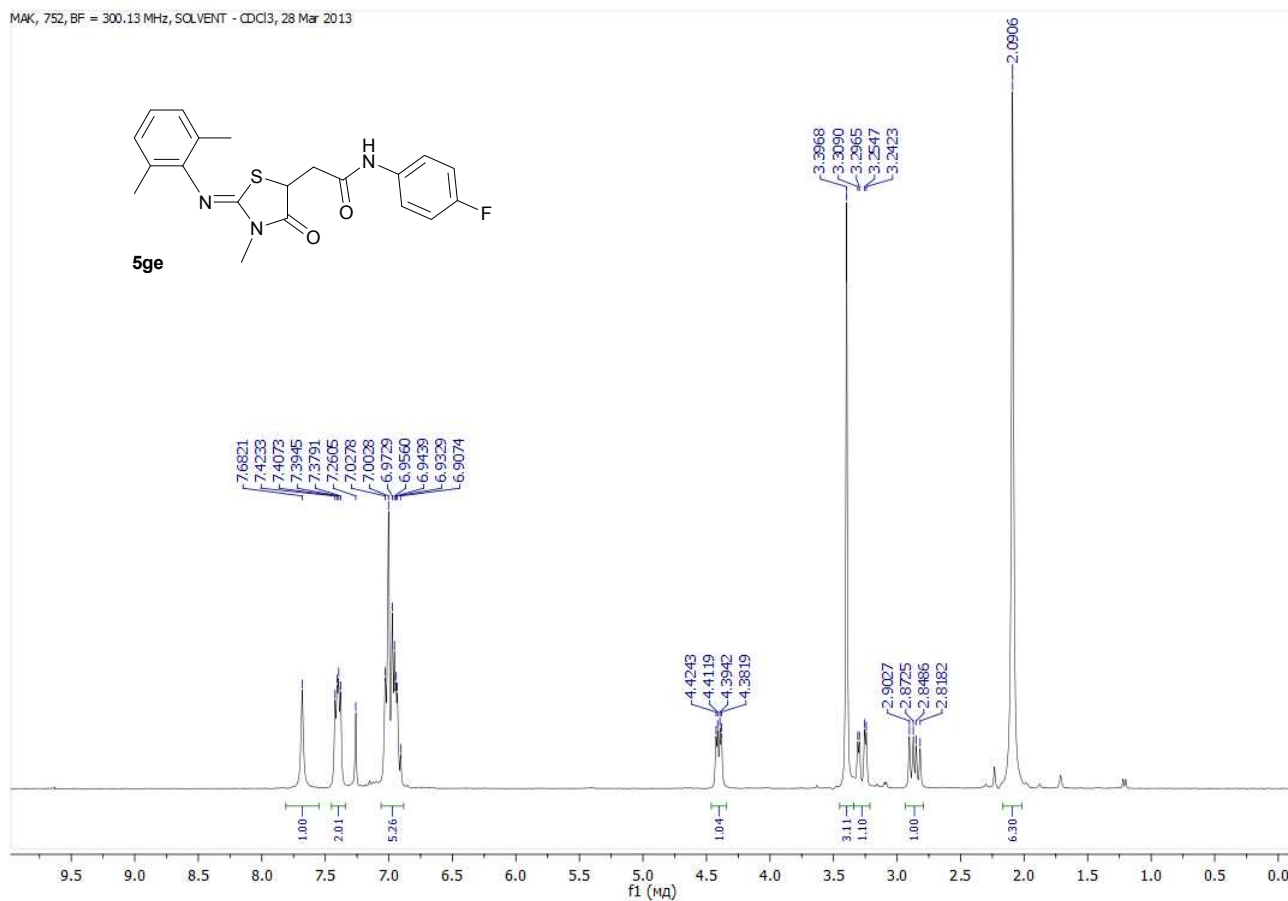
MAK, 767, BF = 300.13 MHz, SOLVENT - CDCl₃, 3 Apr 2013



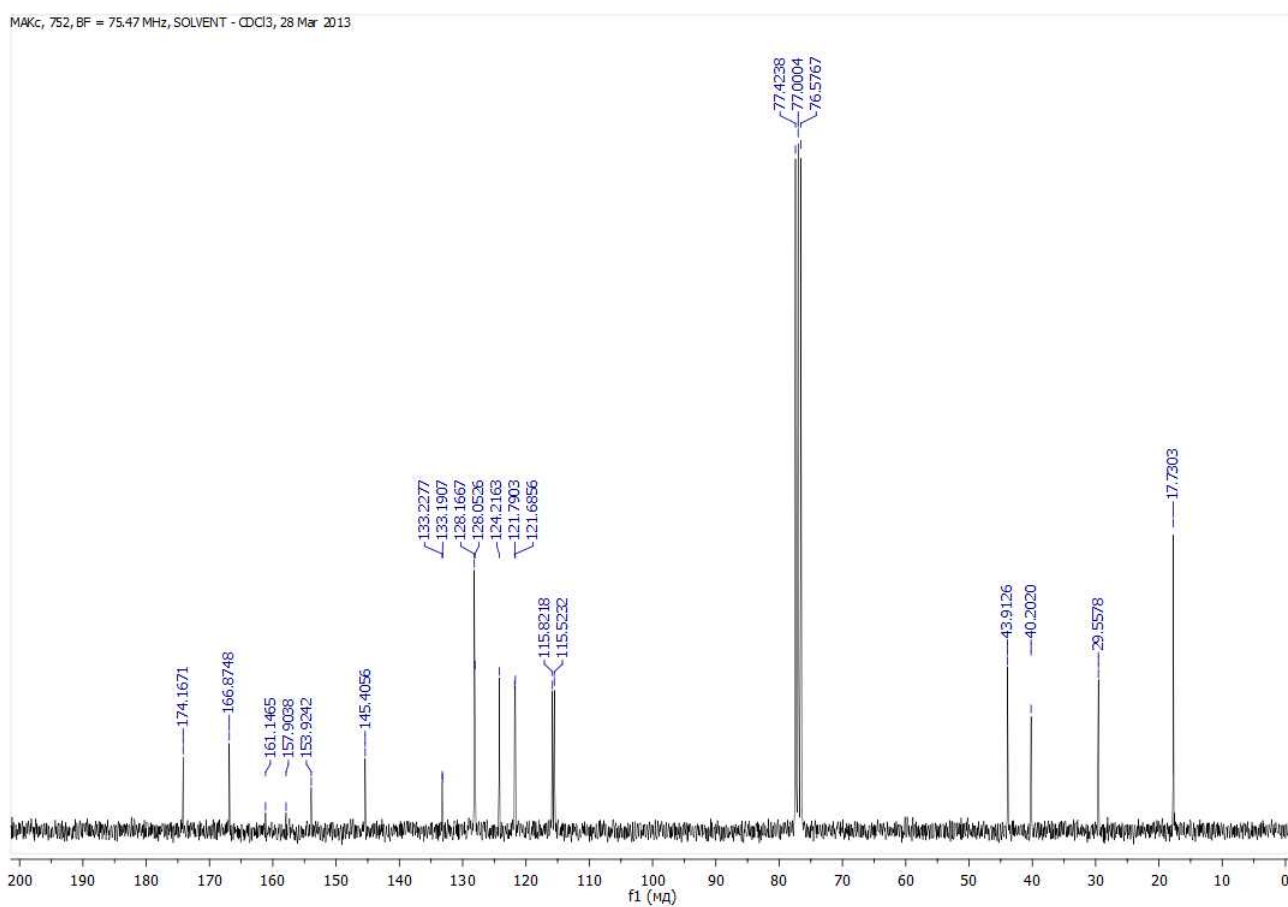
MAKc, 838, BF = 100.612769 MHz, Solvent - CDCl₃, 16 Apr 2013 T=301 K



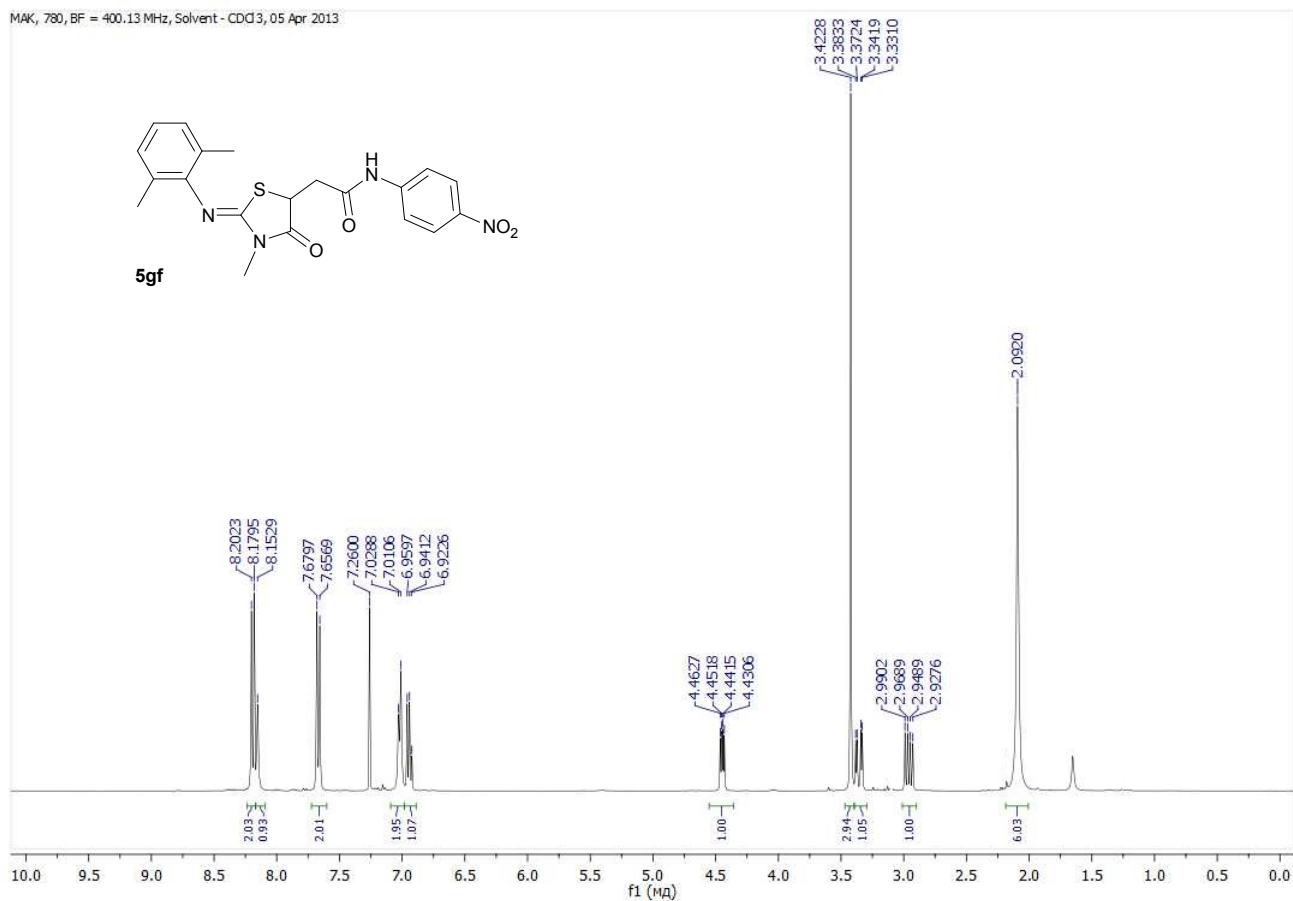
MAK, 752, BF = 300.13 MHz, SOLVENT - CDCl₃, 28 Mar 2013



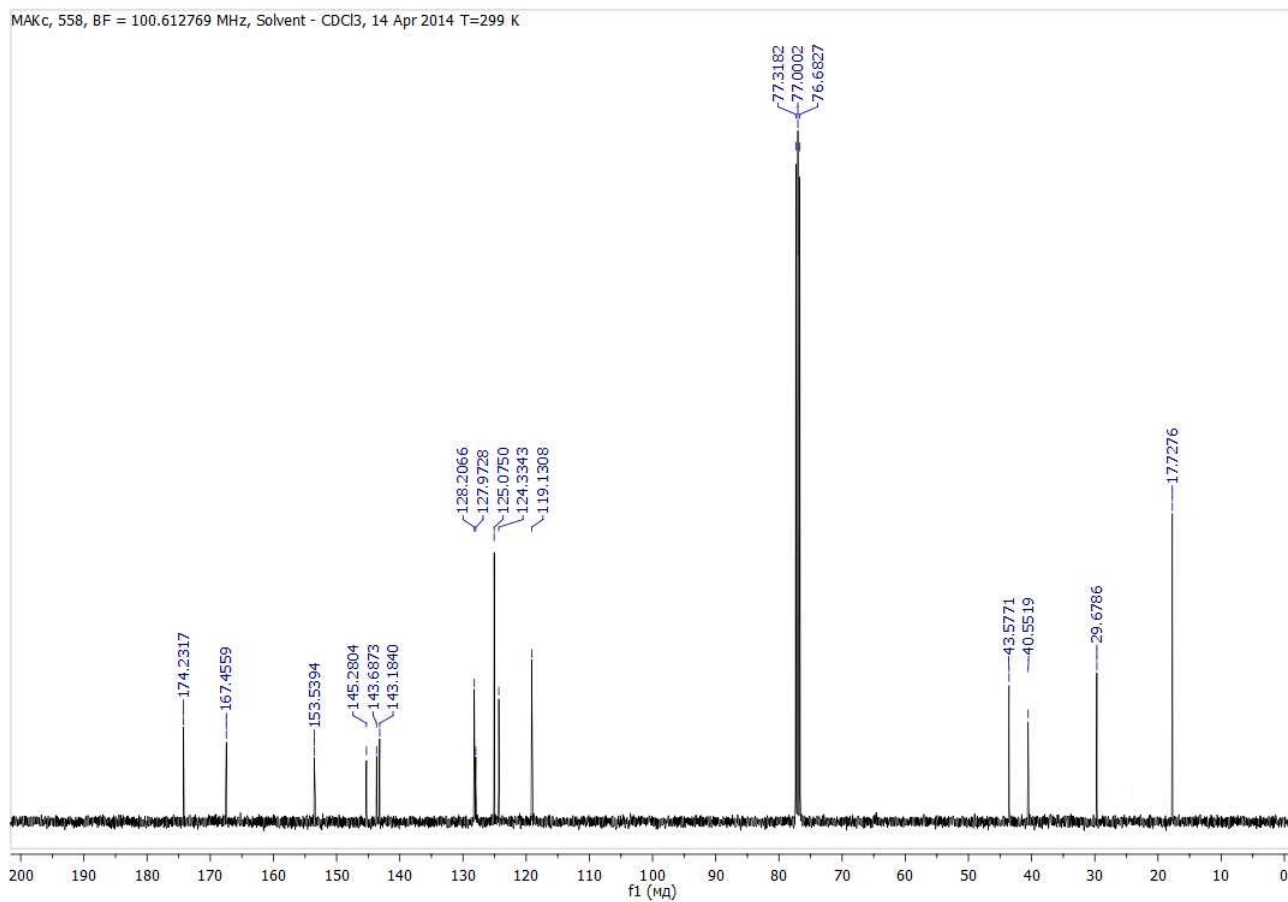
MAKc, 752, BF = 75.47 MHz, SOLVENT - CDCl₃, 28 Mar 2013



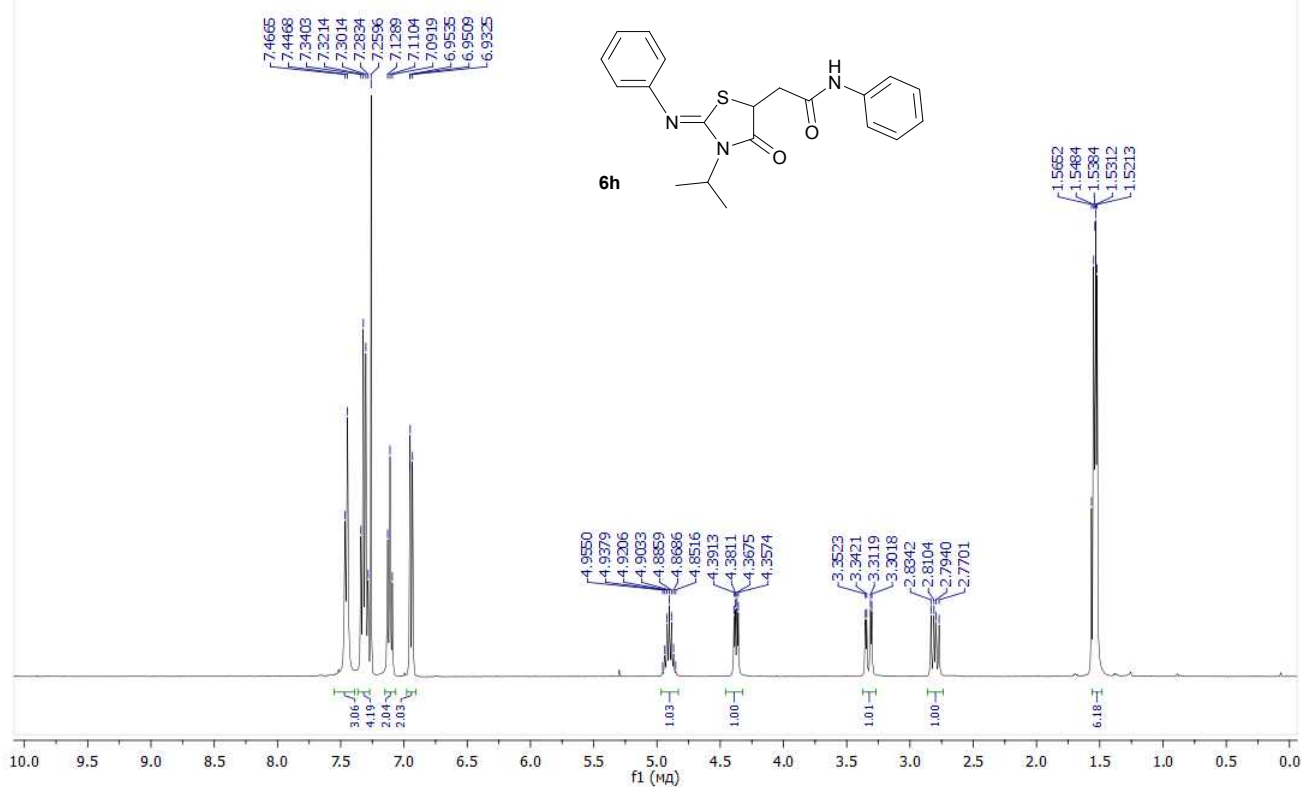
MAK, 780, BF = 400.13 MHz, Solvent - CDCl₃, 05 Apr 2013



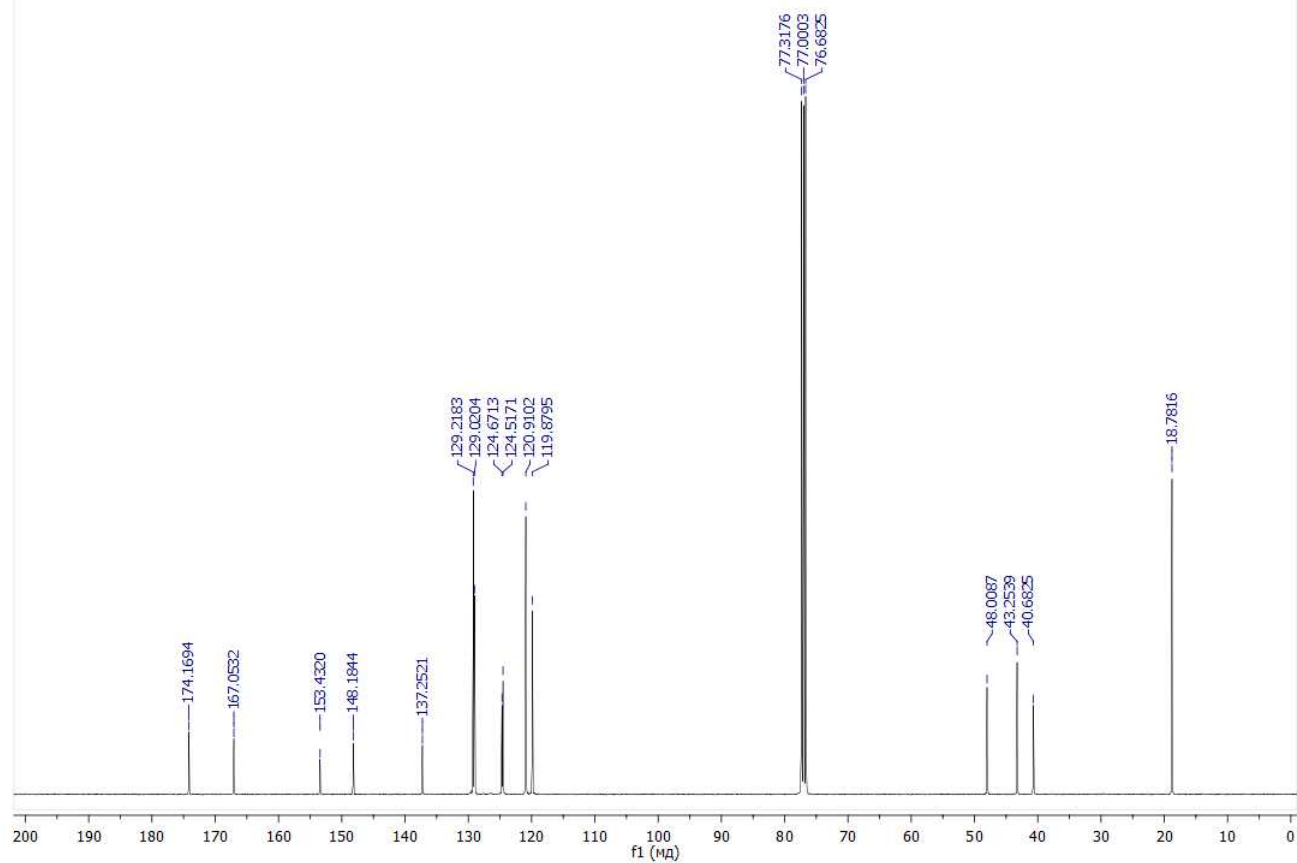
MAKc, 558, BF = 100.612769 MHz, Solvent - CDCl₃, 14 Apr 2014 T=299 K



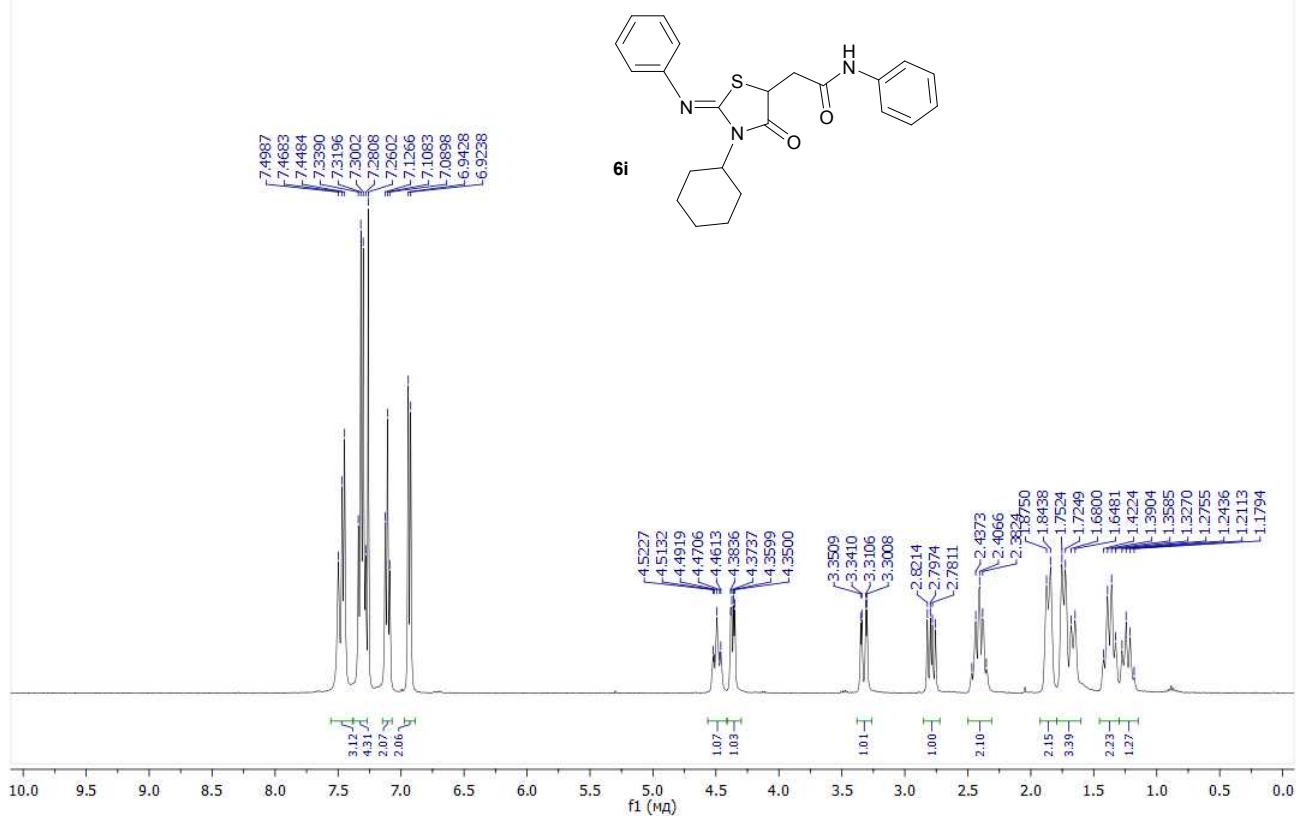
MAK, 64, BF = 400.13 MHz, Solvent - CDCl₃, 16 Sep 2013 T=297 K



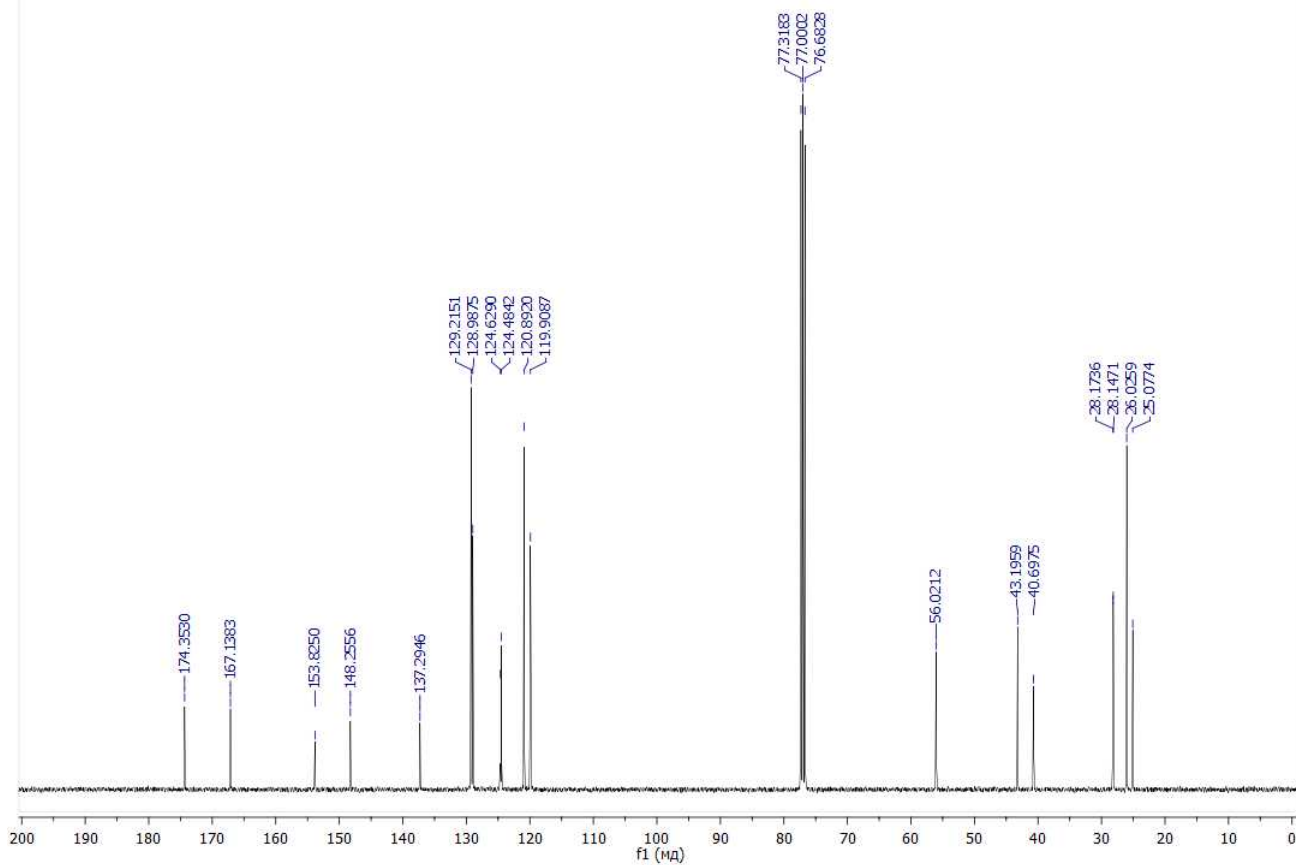
MAKc, 559, BF = 100.612769 MHz, Solvent - CDD3, 14 Apr 2014 T=299 K

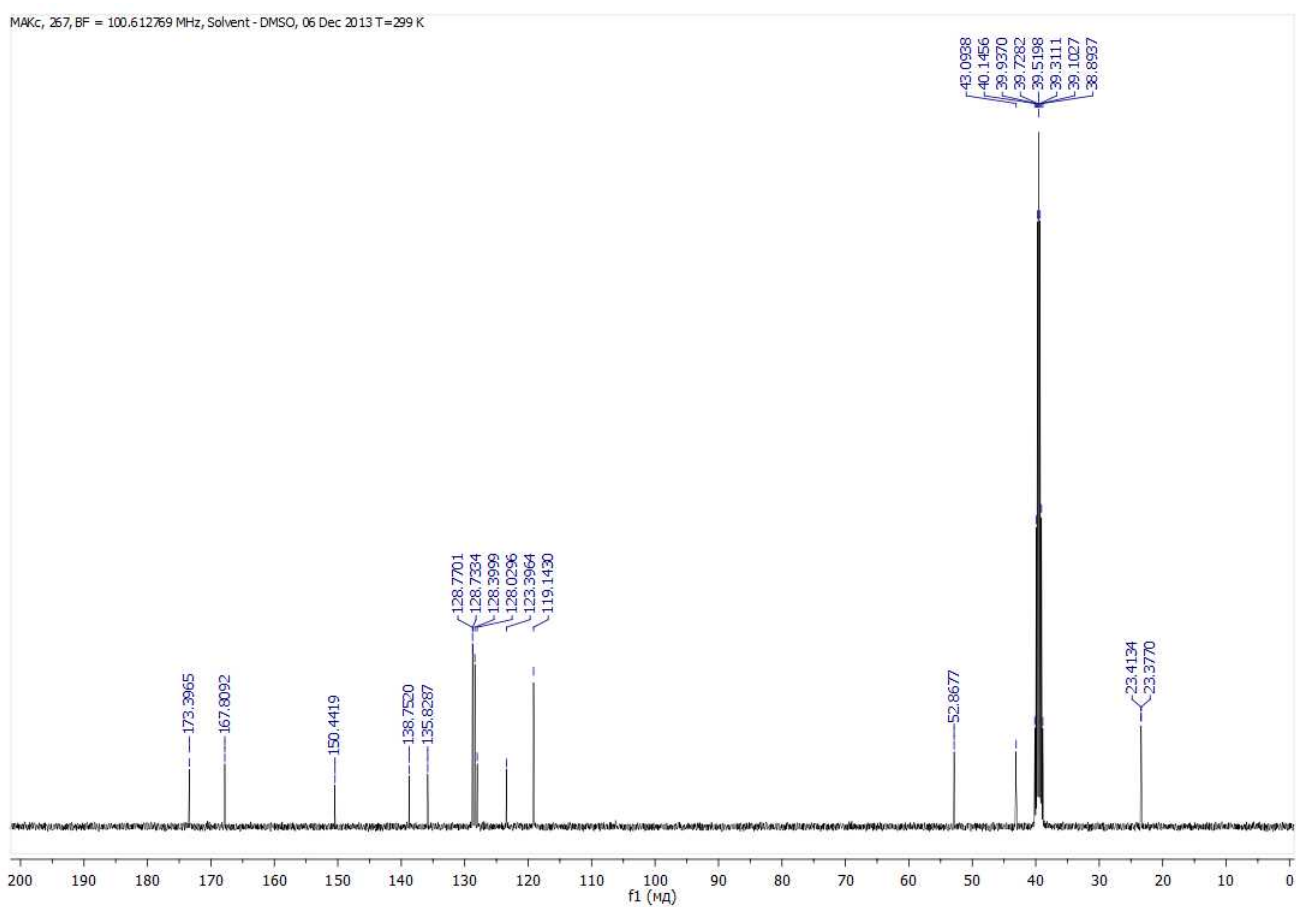
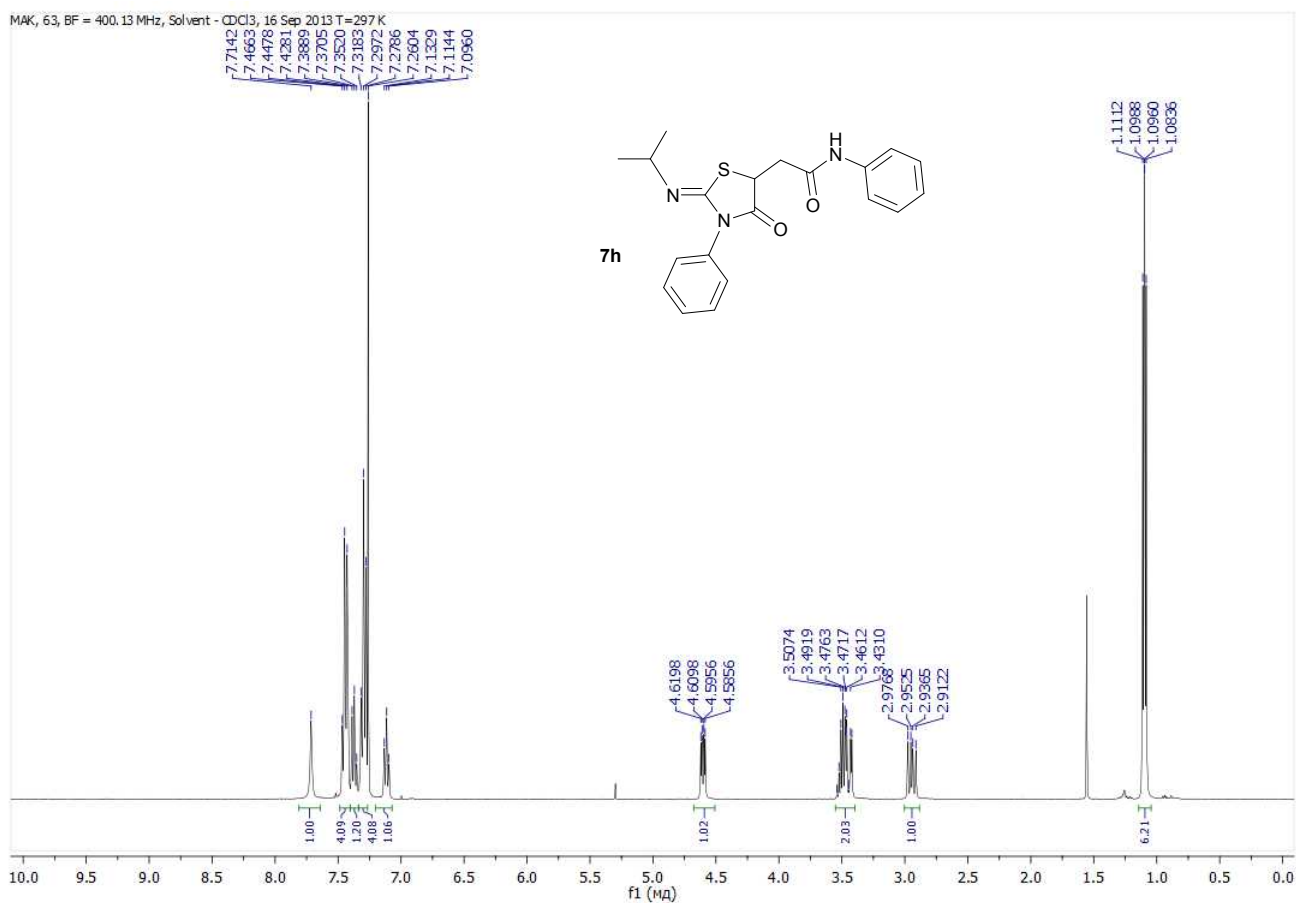


MAK, 93, BF = 400.13 MHz, Solvent - CDCl₃, 27 Sep 2013 T=297 K

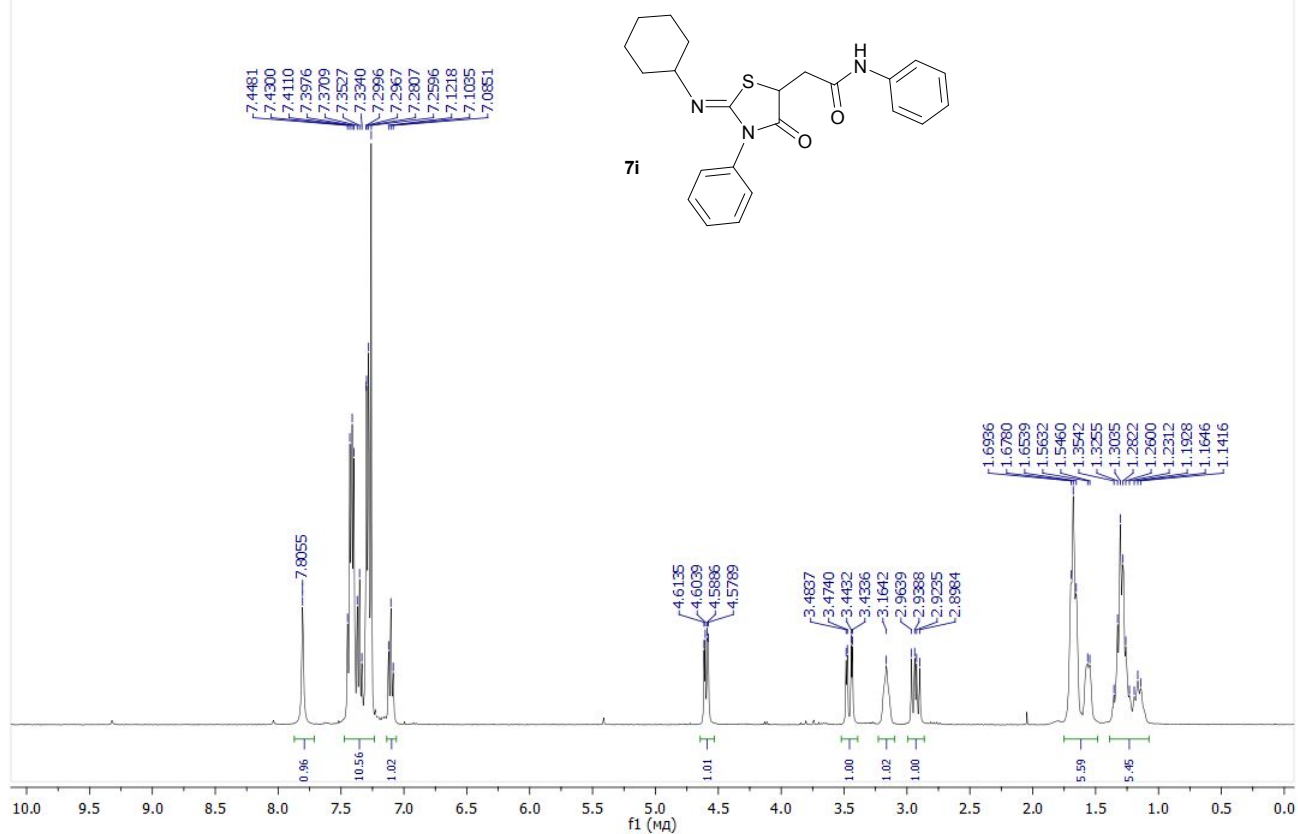


MAKc, 93, BF = 100.612769 MHz, Solvent - CDCl₃, 04 Oct 2013 T=298 K

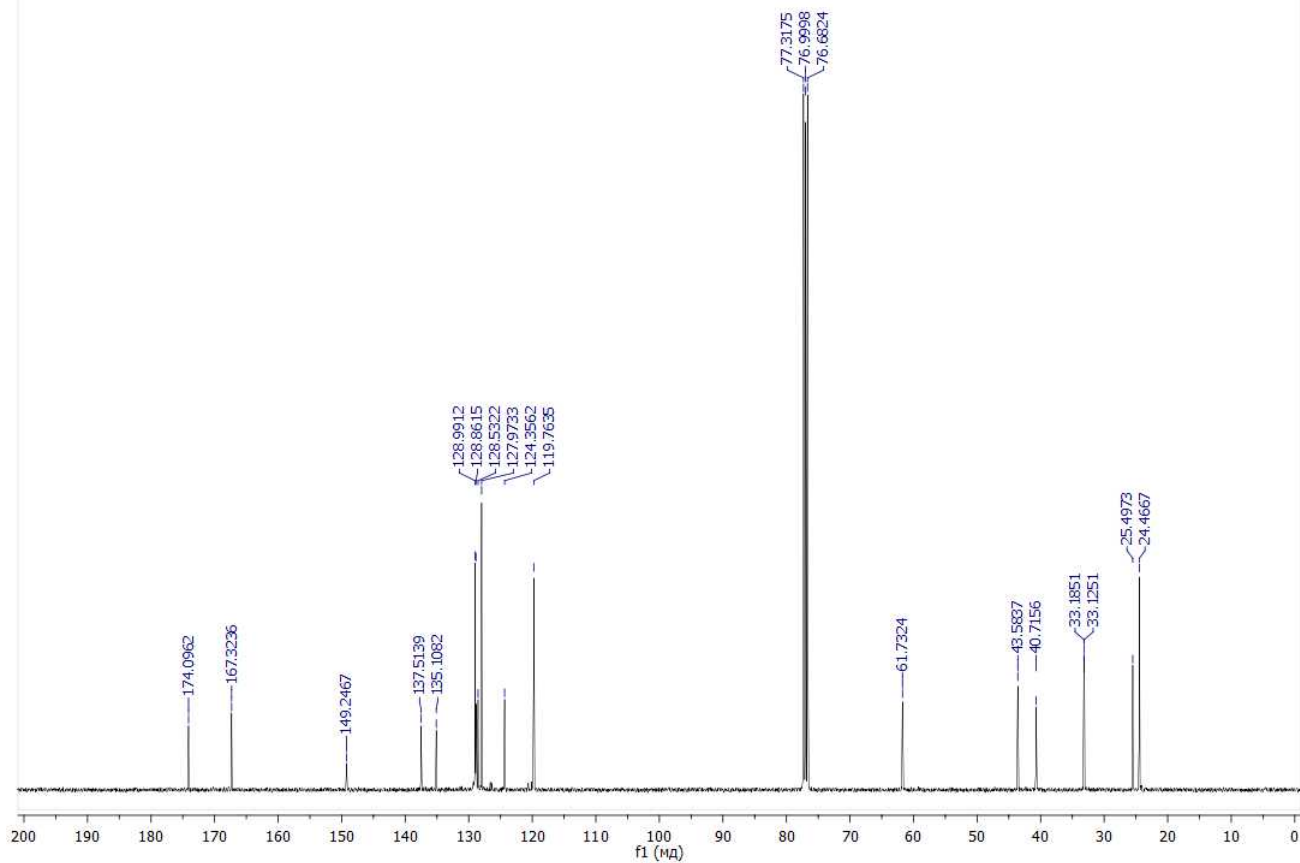




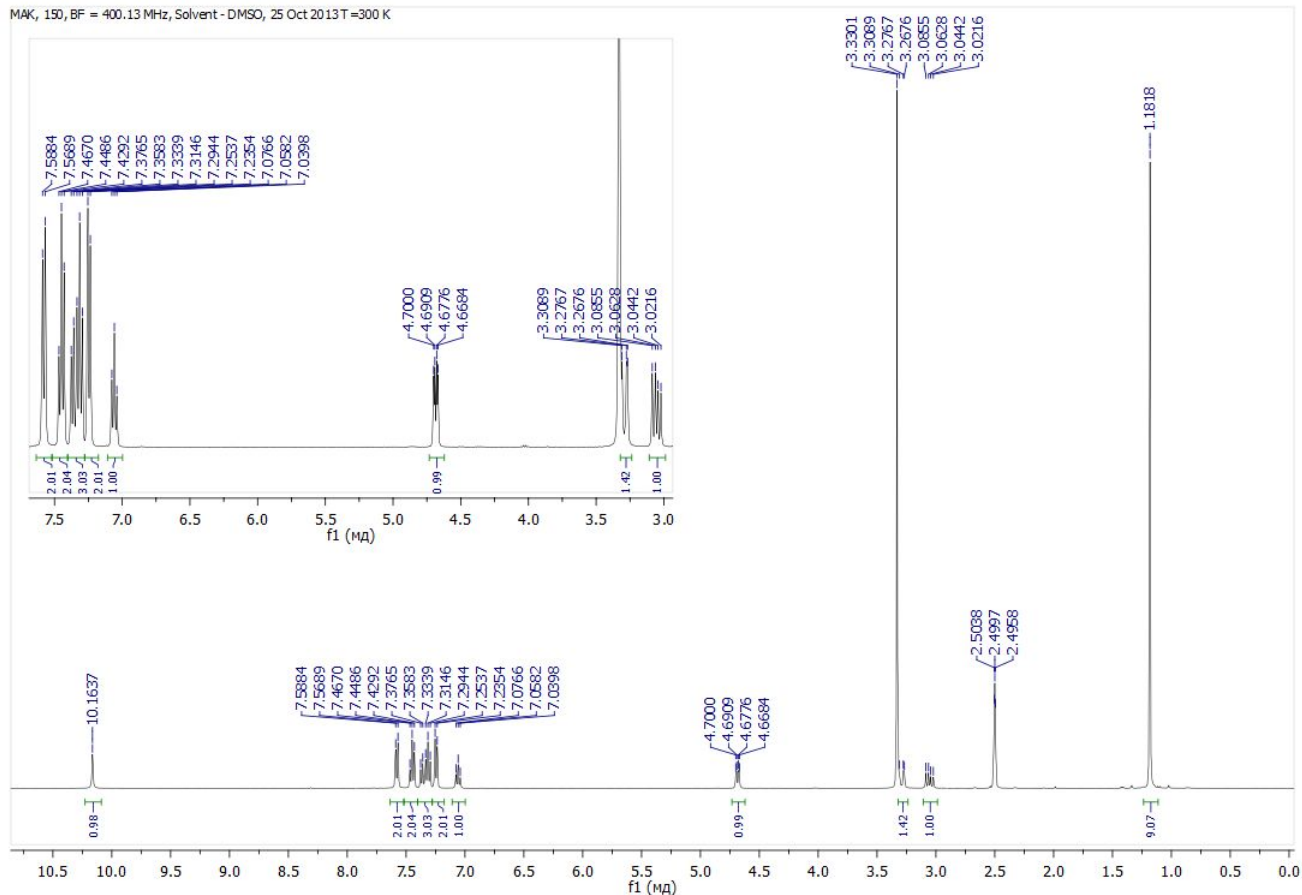
MAK, 92, BF = 400.13 MHz, Solvent - CDCl₃, 27 Sep 2013 T=297 K



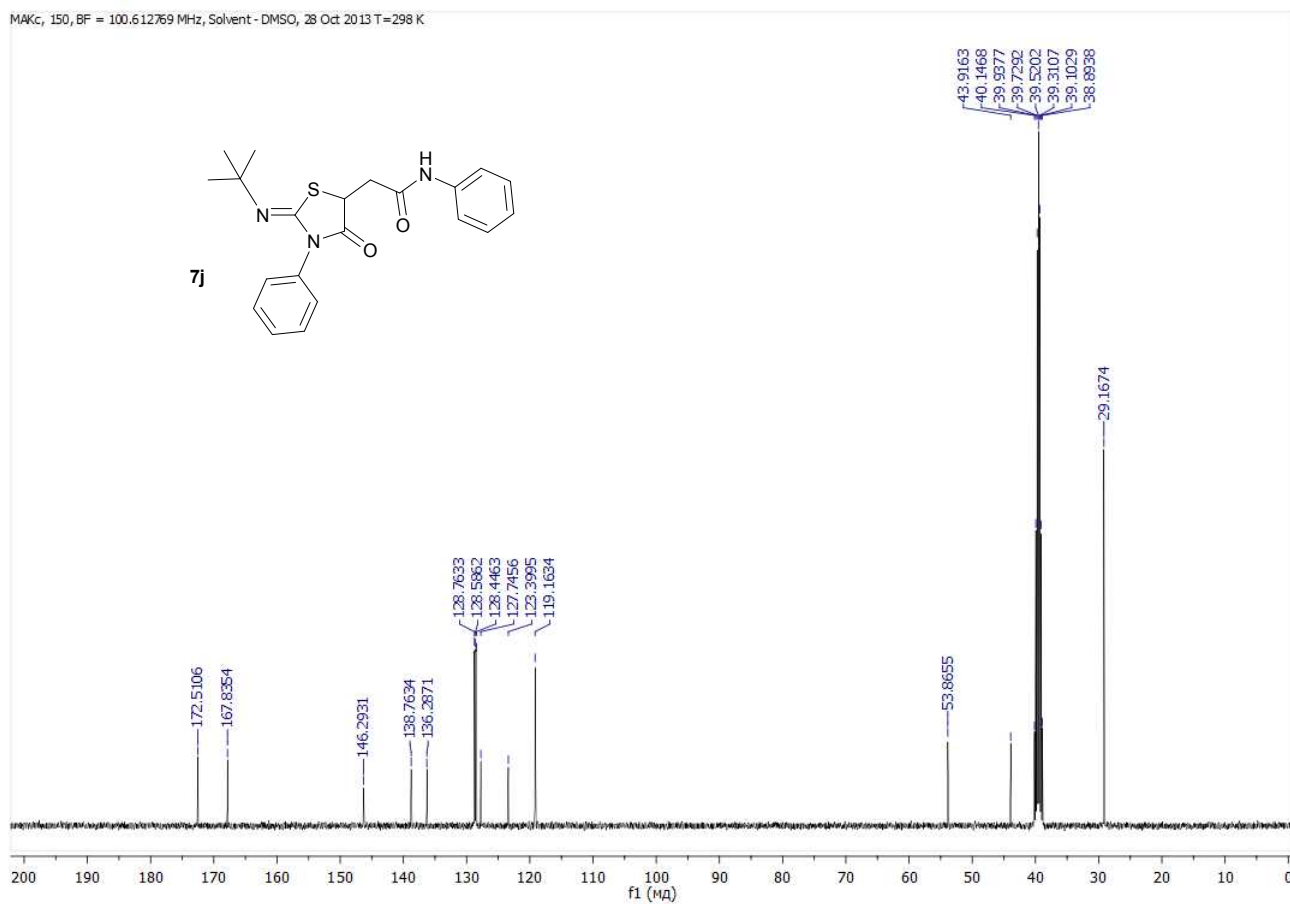
MAKc, 92, BF = 100.612769 MHz, Solvent - CDCl₃, 04 Oct 2013 T=298 K



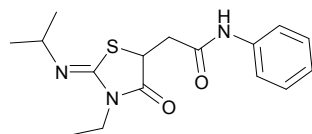
MAK, 150, BF = 400.13 MHz, Solvent - DMSO, 25 Oct 2013 T=300 K



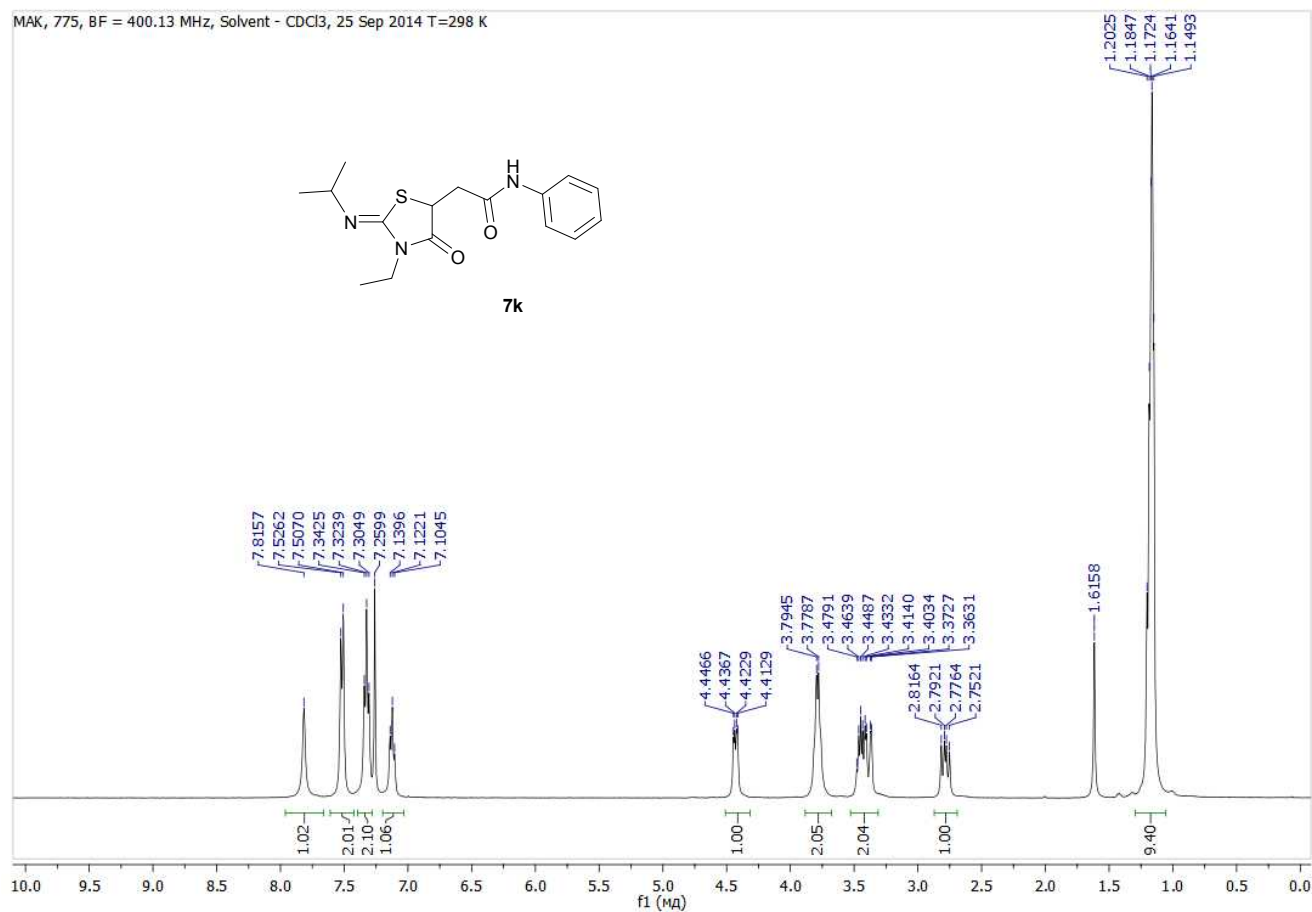
MAKc, 150, BF = 100.612769 MHz, Solvent - DMSO, 28 Oct 2013 T=298 K



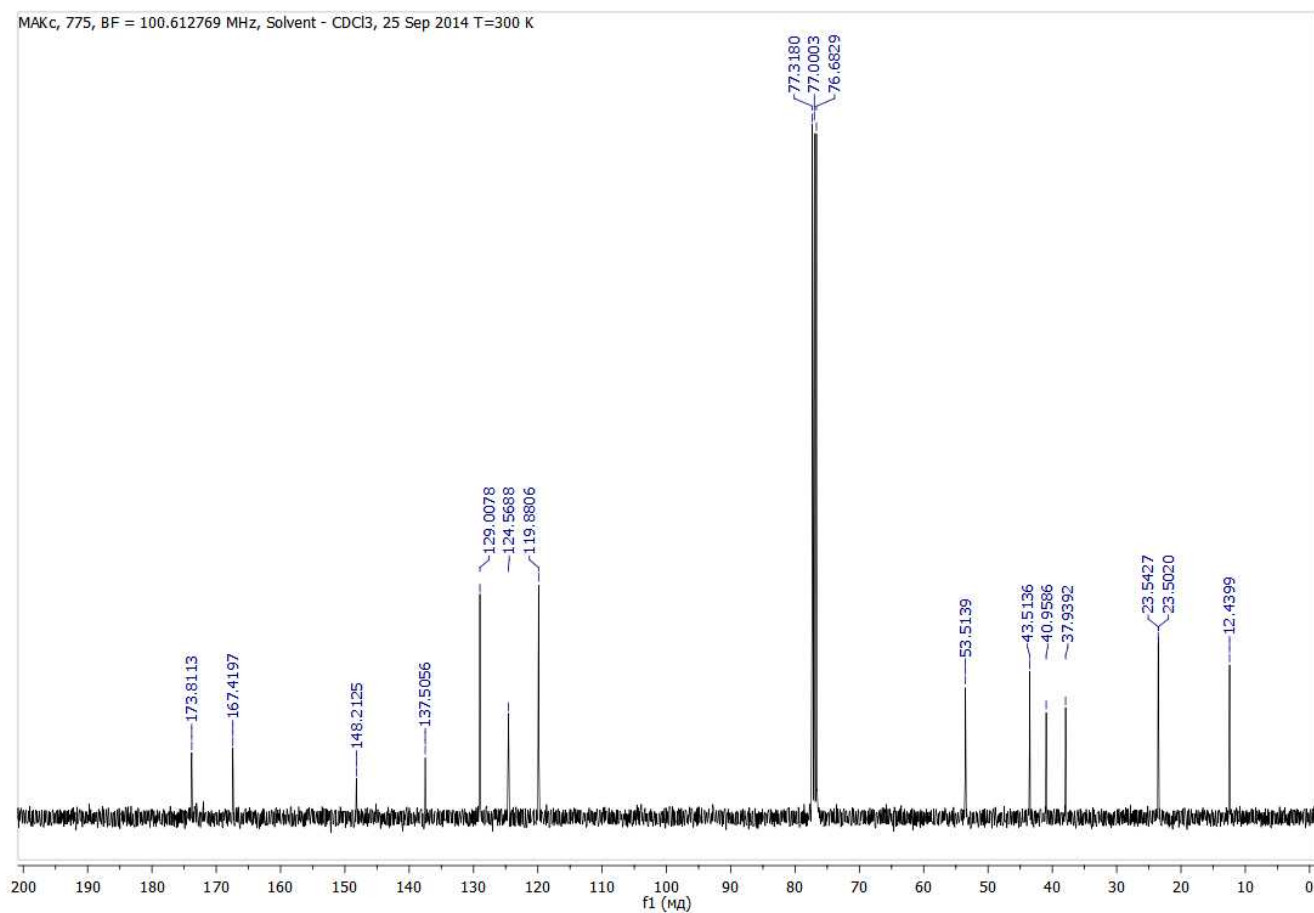
MAK, 775, BF = 400.13 MHz, Solvent - CDCl₃, 25 Sep 2014 T=298 K



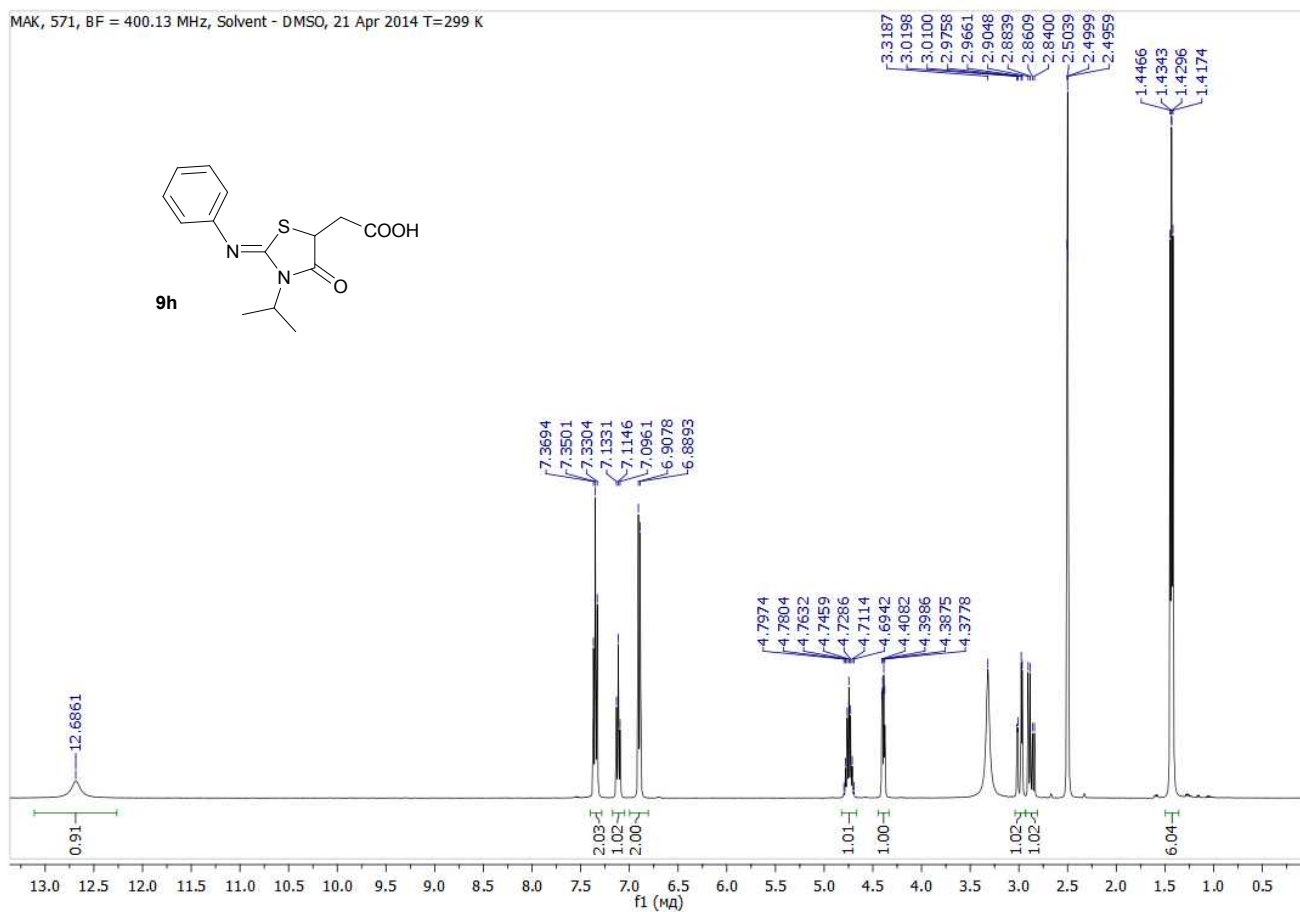
7k



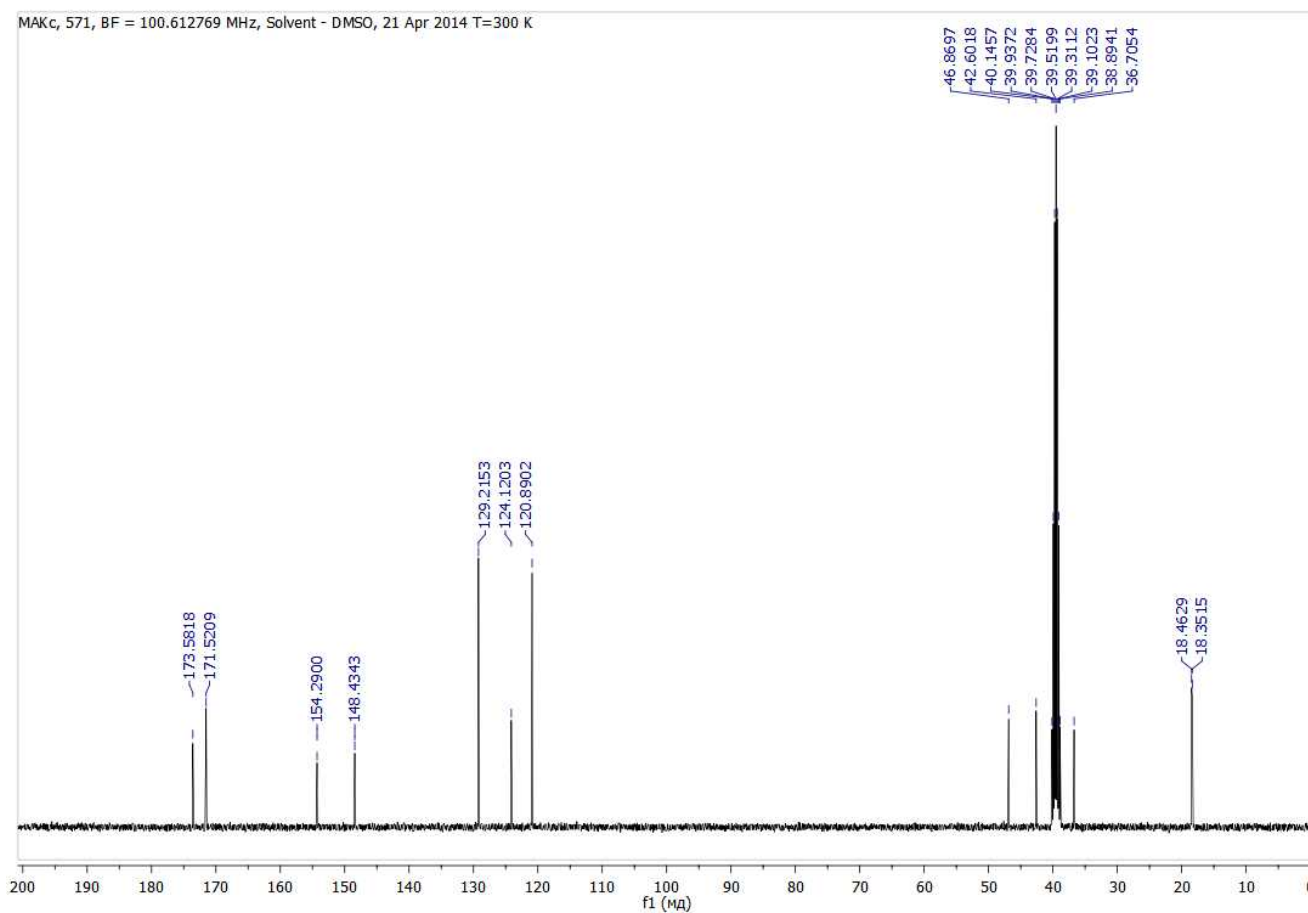
MAKc, 775, BF = 100.612769 MHz, Solvent - CDCl₃, 25 Sep 2014 T=300 K



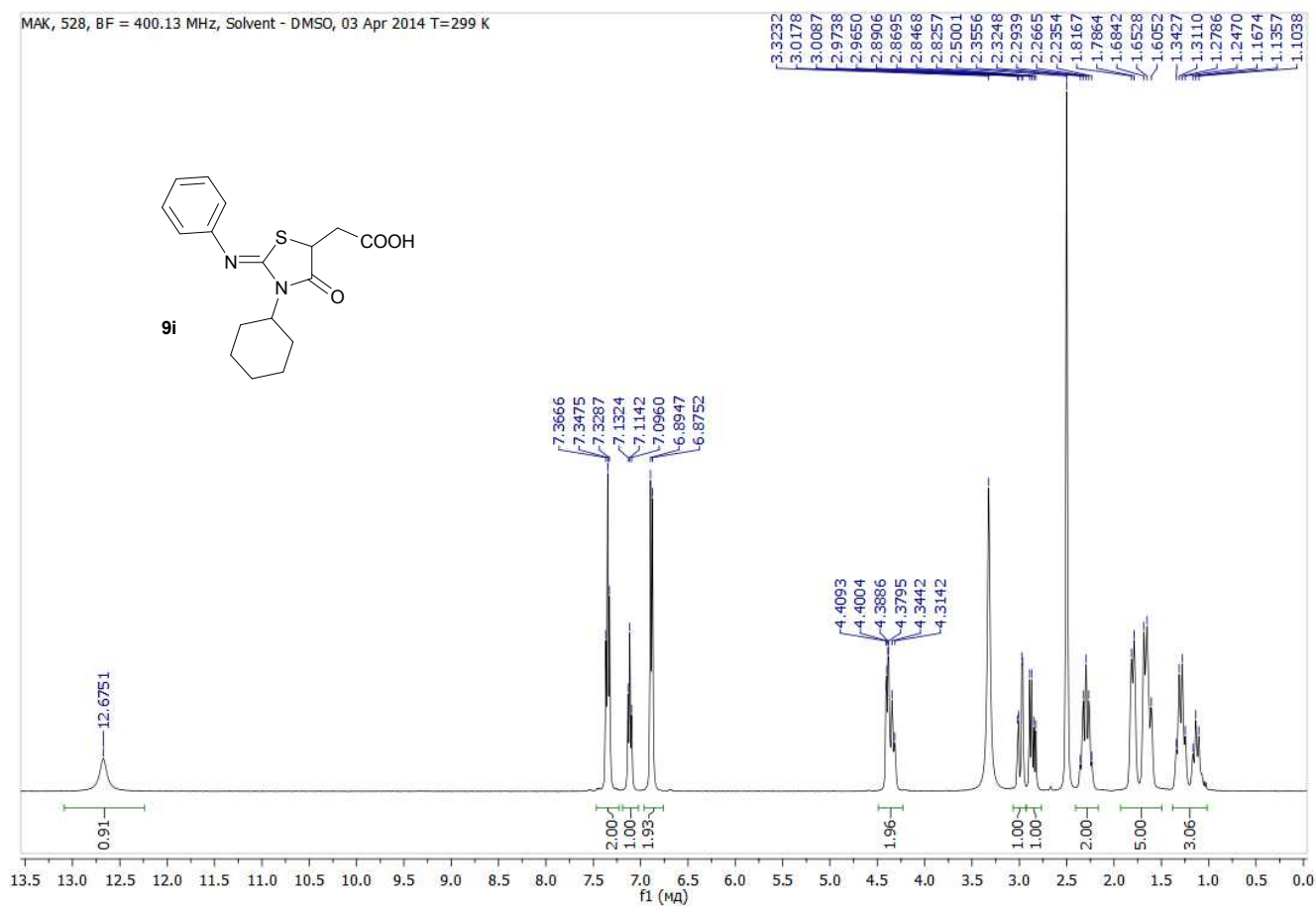
MAK, 571, BF = 400.13 MHz, Solvent - DMSO, 21 Apr 2014 T=299 K



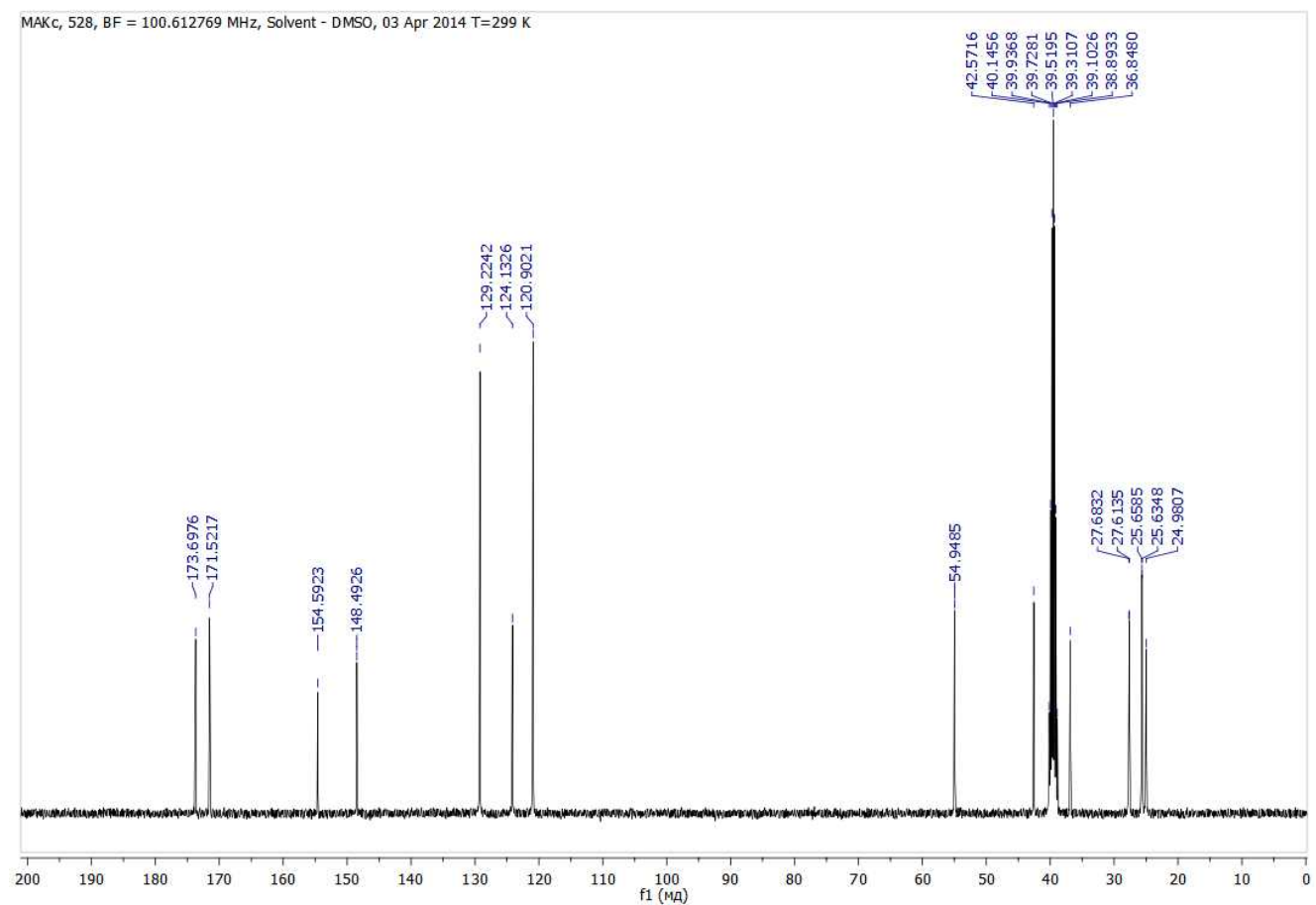
MAKc, 571, BF = 100.612769 MHz, Solvent - DMSO, 21 Apr 2014 T=300 K



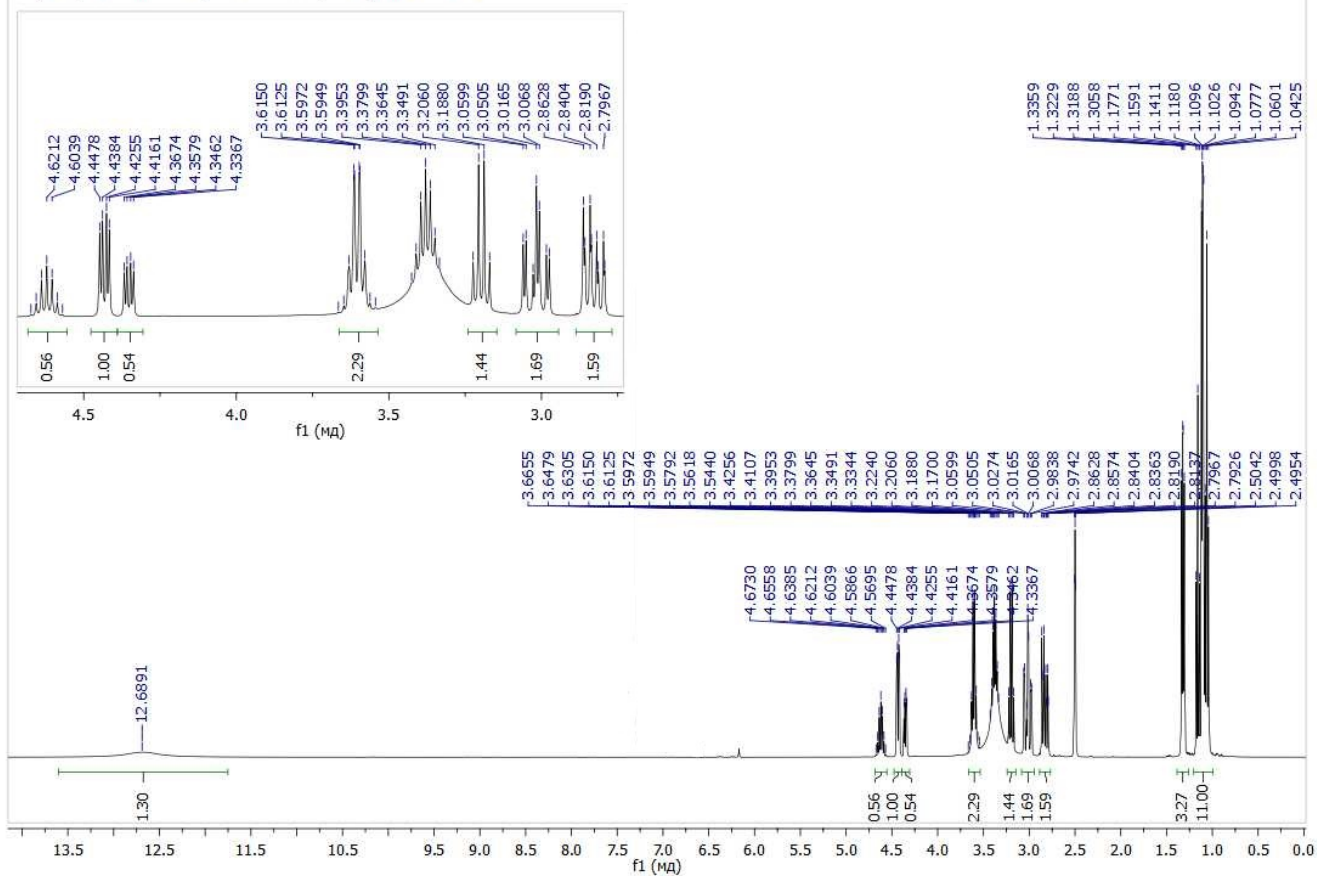
MAK, 528, BF = 400.13 MHz, Solvent - DMSO, 03 Apr 2014 T=299 K



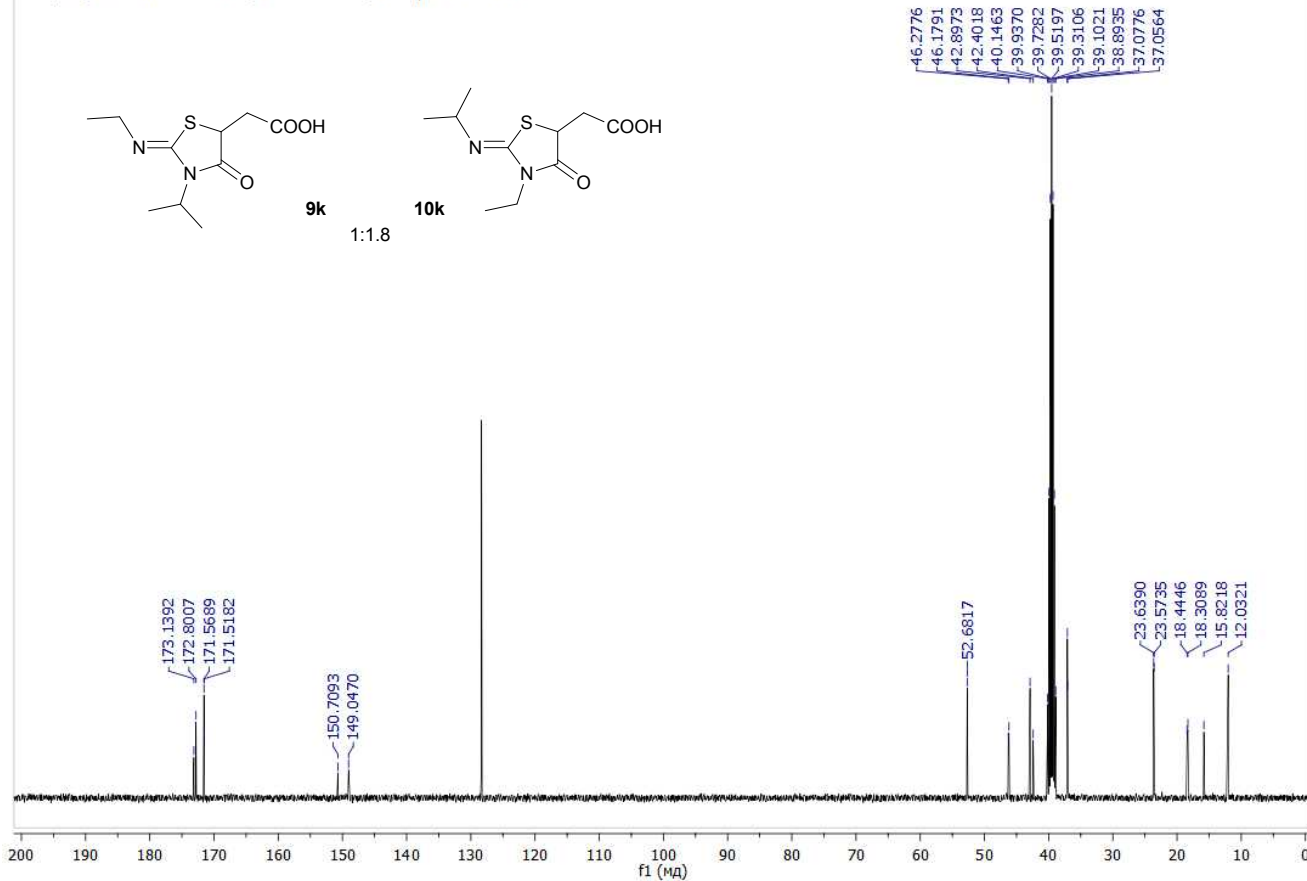
MAKc, 528, BF = 100.612769 MHz, Solvent - DMSO, 03 Apr 2014 T=299 K



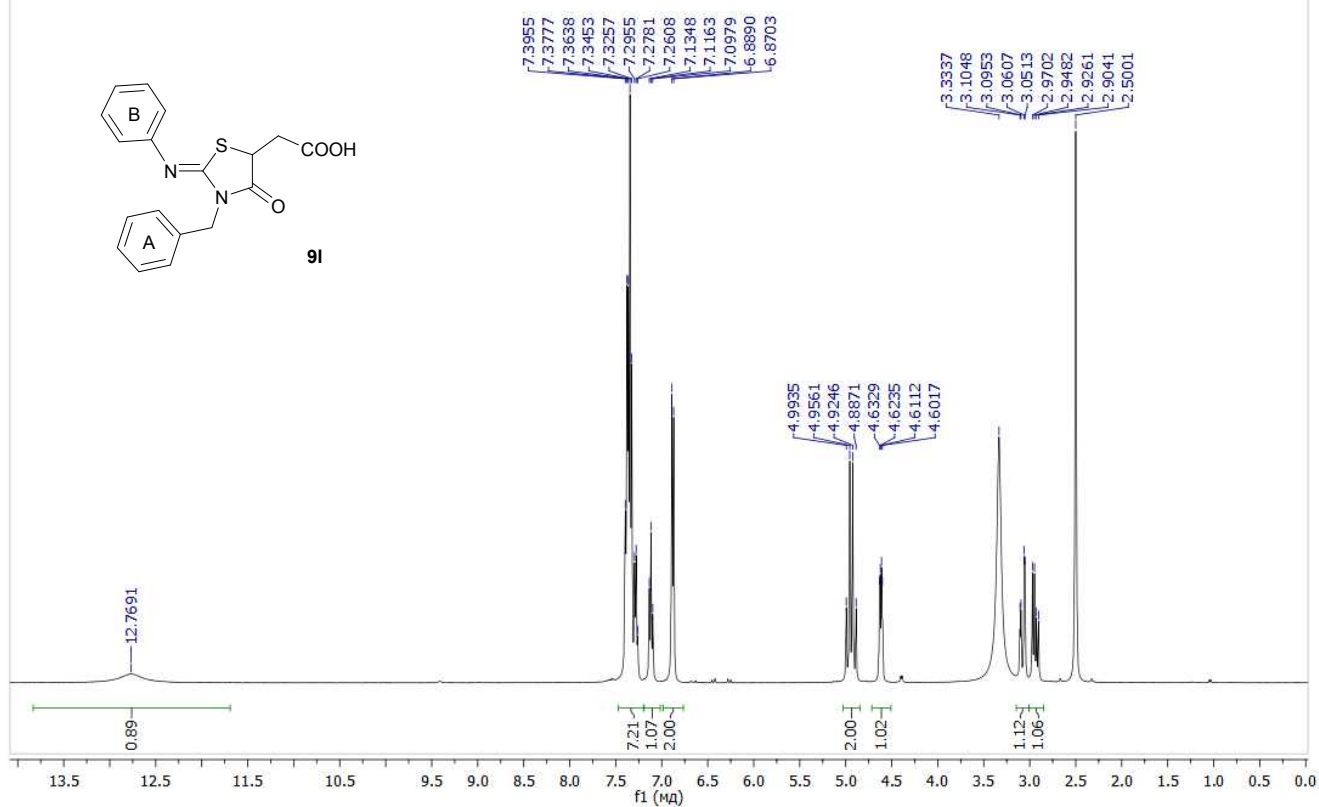
MAK, 786, BF = 400.13 MHz, Solvent - DMSO, 29 Sep 2014 T=298 K



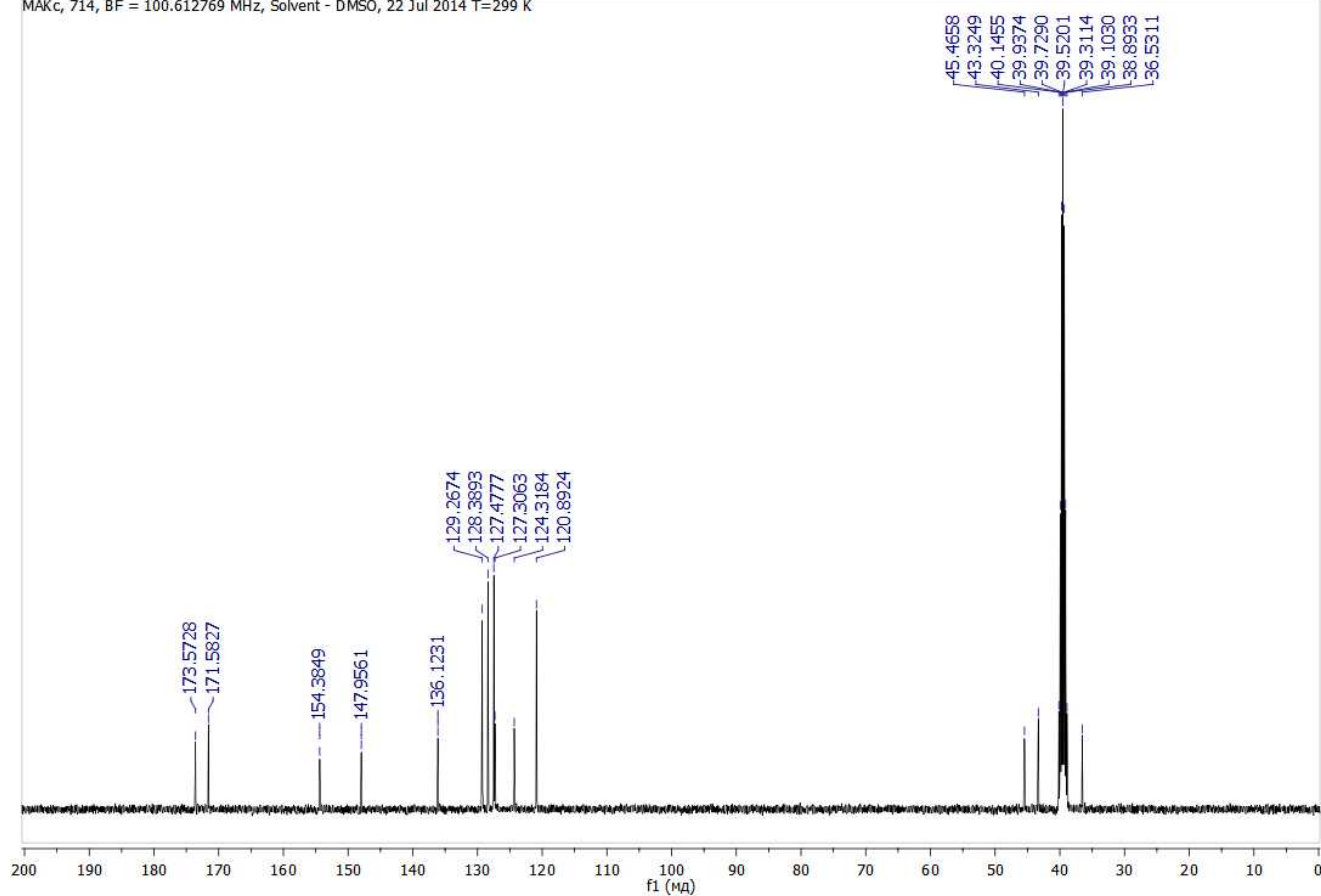
MAKc, 786, BF = 100.612769 MHz, Solvent - DMSO, 30 Sep 2014 T=300 K

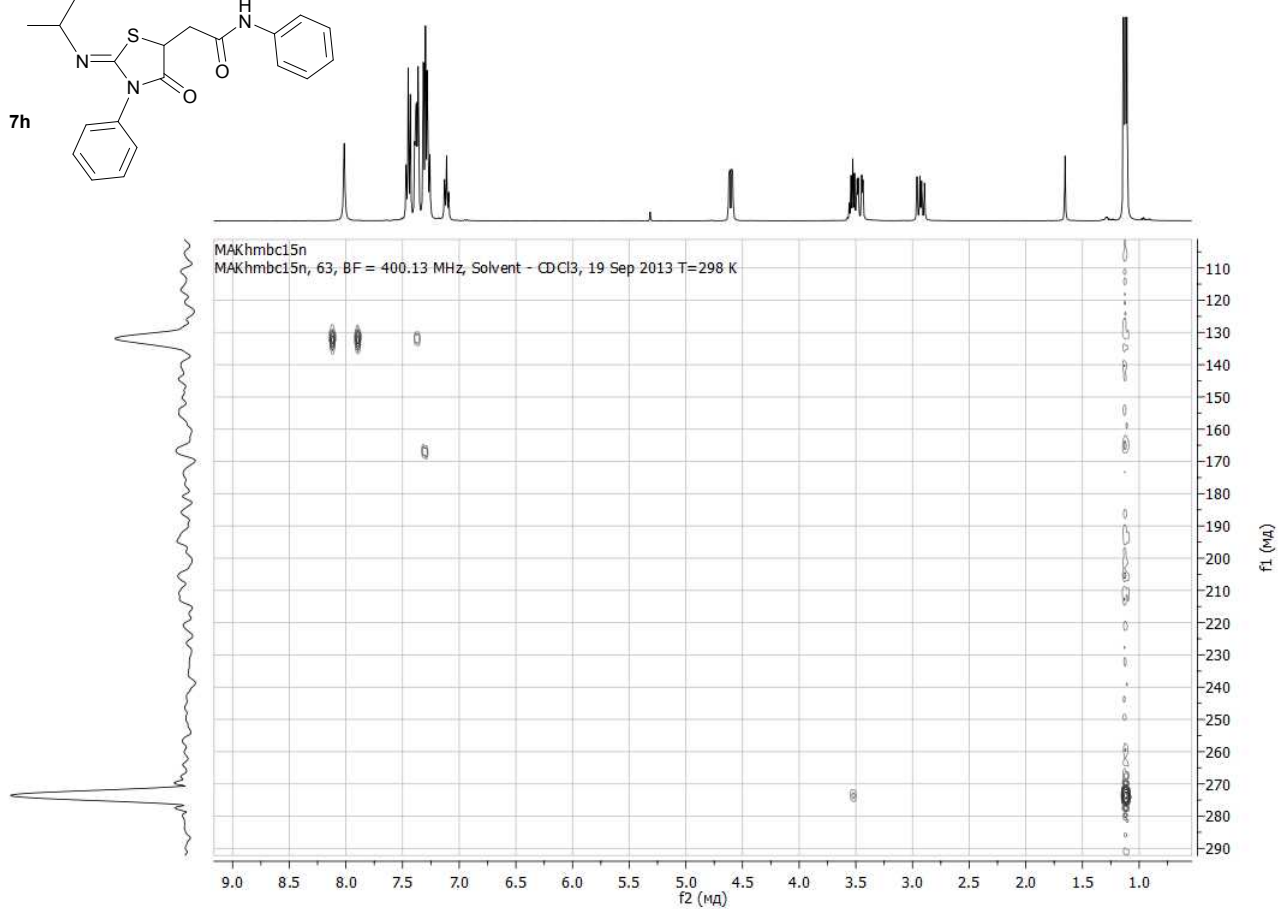
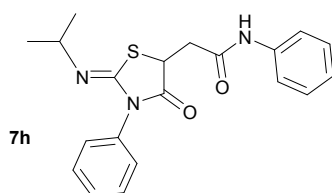
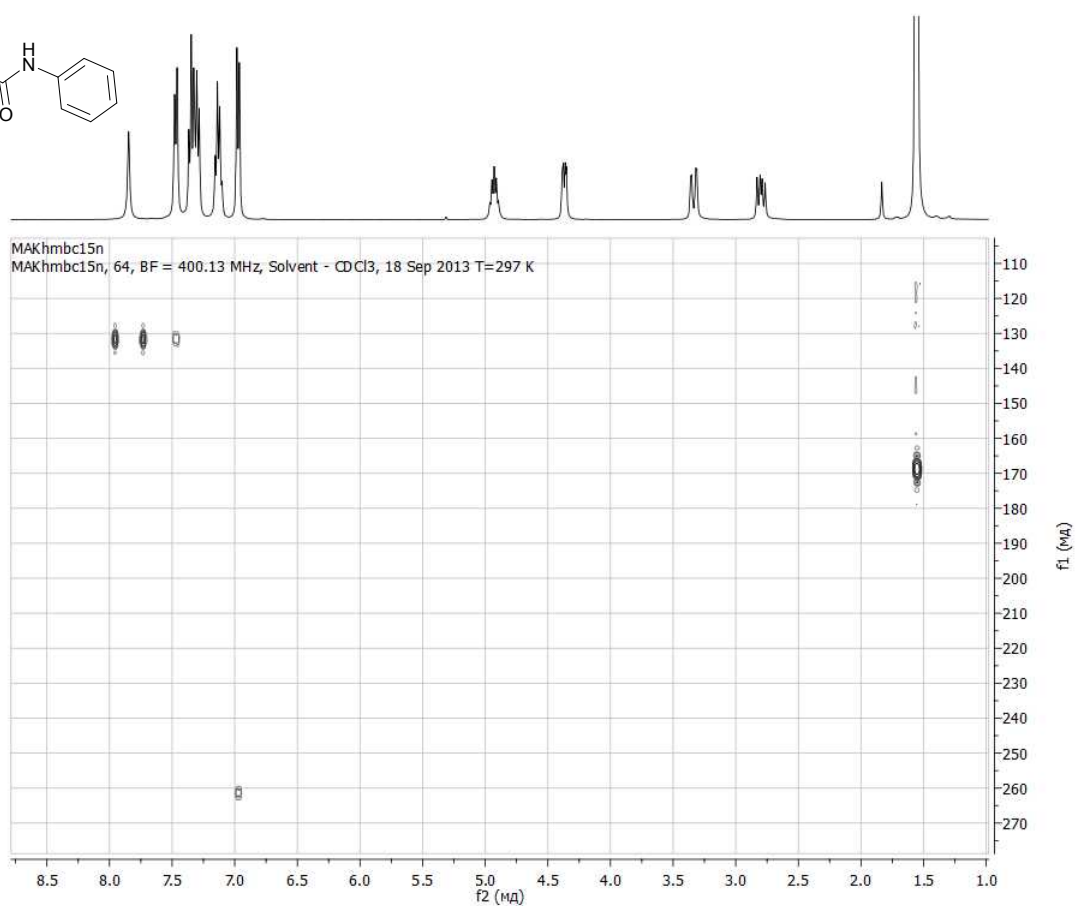
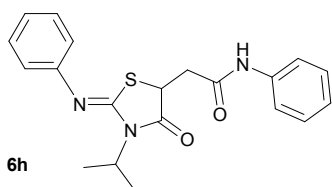


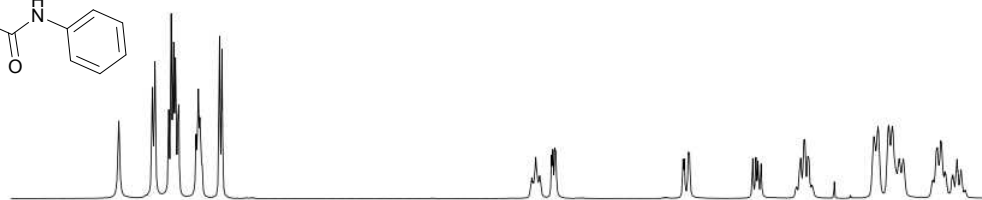
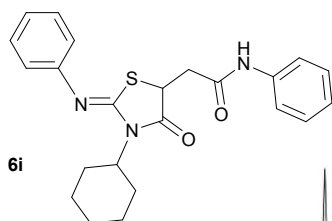
MAK, 714, BF = 400.13 MHz, Solvent - DMSO, 21 Jul 2014 T=299 K



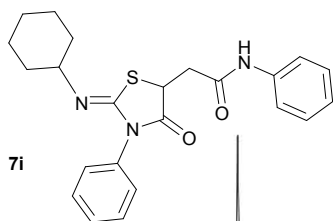
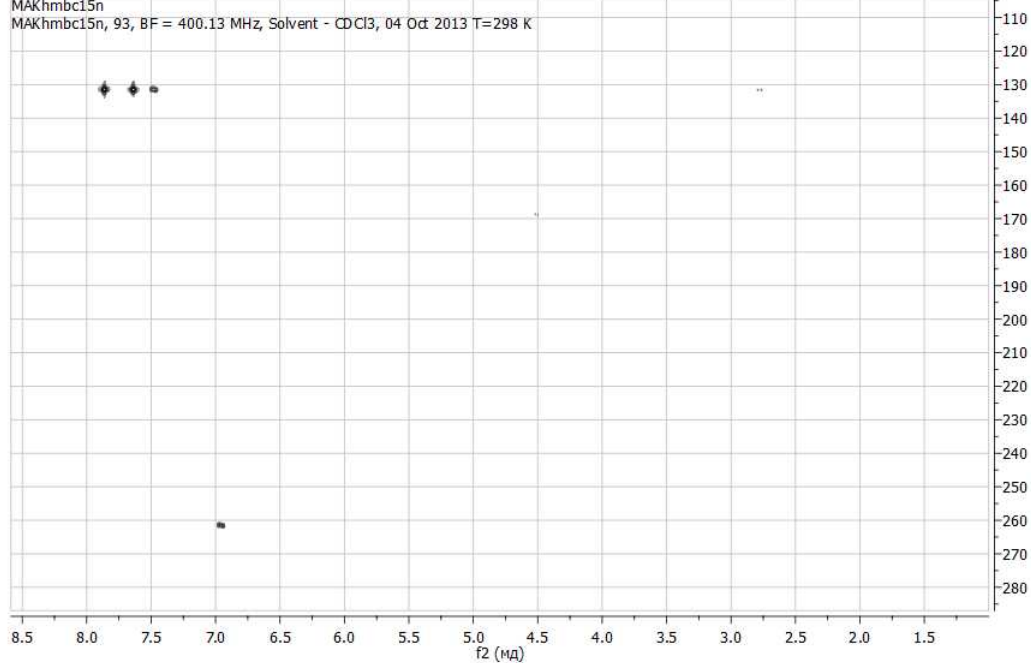
MAKc, 714, BF = 100.612769 MHz, Solvent - DMSO, 22 Jul 2014 T=299 K



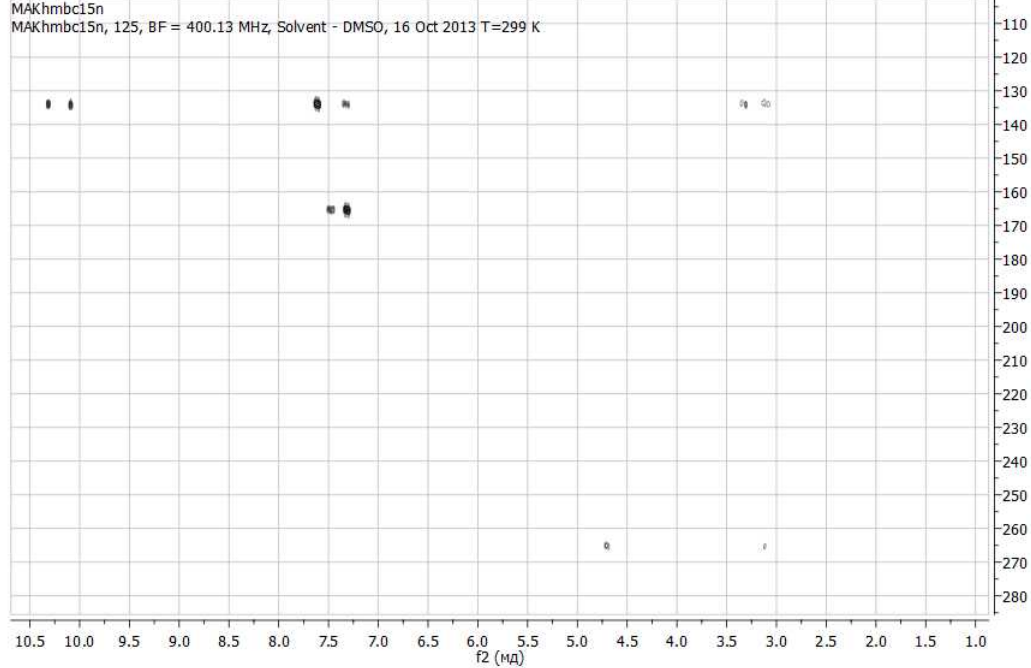


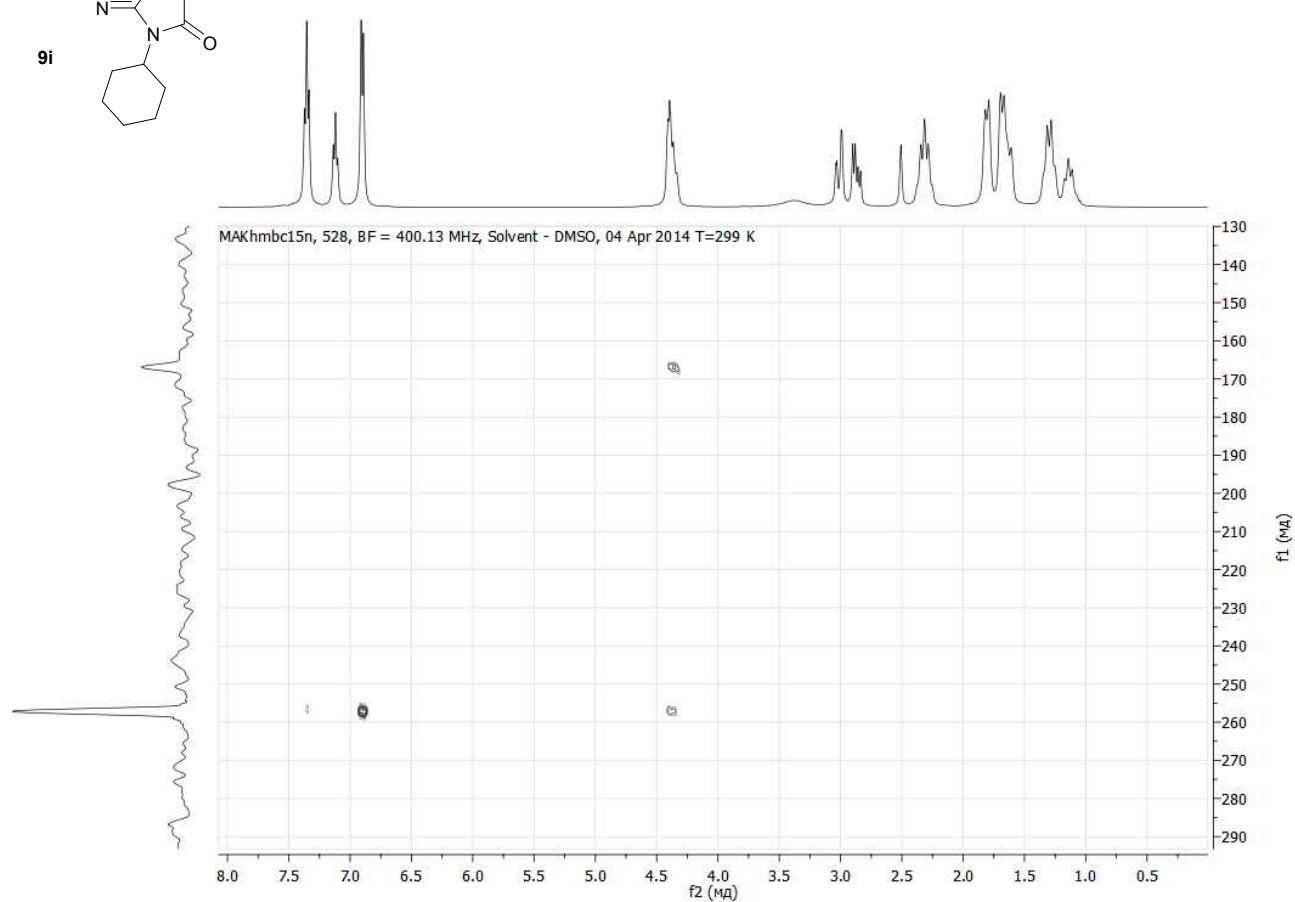
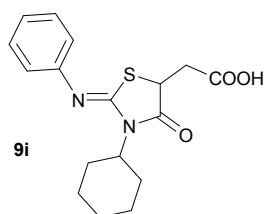


MAKhmbc15n
MAKhmbc15n, 93, BF = 400.13 MHz, Solvent - CDCl₃, 04 Oct 2013 T=298 K



MAKhmbc15n
MAKhmbc15n, 125, BF = 400.13 MHz, Solvent - DMSO, 16 Oct 2013 T=299 K





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