

Supporting Information

Salicylamide derivatives as potent HBV inhibitors: Insights into structure-activity relationships

Jingwen Huo,^{a,†} Jizhen Xiao,^{a,†} Yushi Zhang,^{a,†} Xinhui Qiu,^a Xuechen Huang,^a Ge
Wang,^a Jianhao Wang,^a Kuancheng Liu^{*abc} and Jimin Xu^{*abc}

^a School of Public Health (Shenzhen), Shenzhen Campus of Sun Yat-sen University,
Shenzhen, 518107, China. E-mail: xujm37@mail.sysu.edu.cn,
liukuanc@mail.sysu.edu.cn

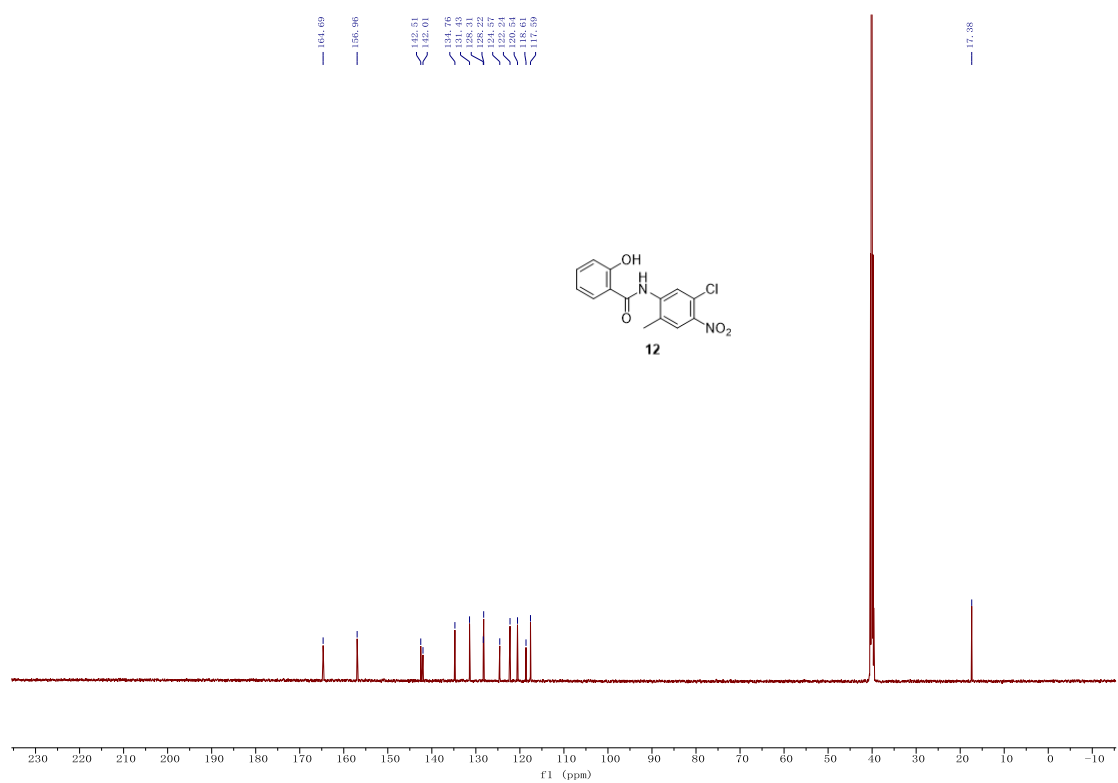
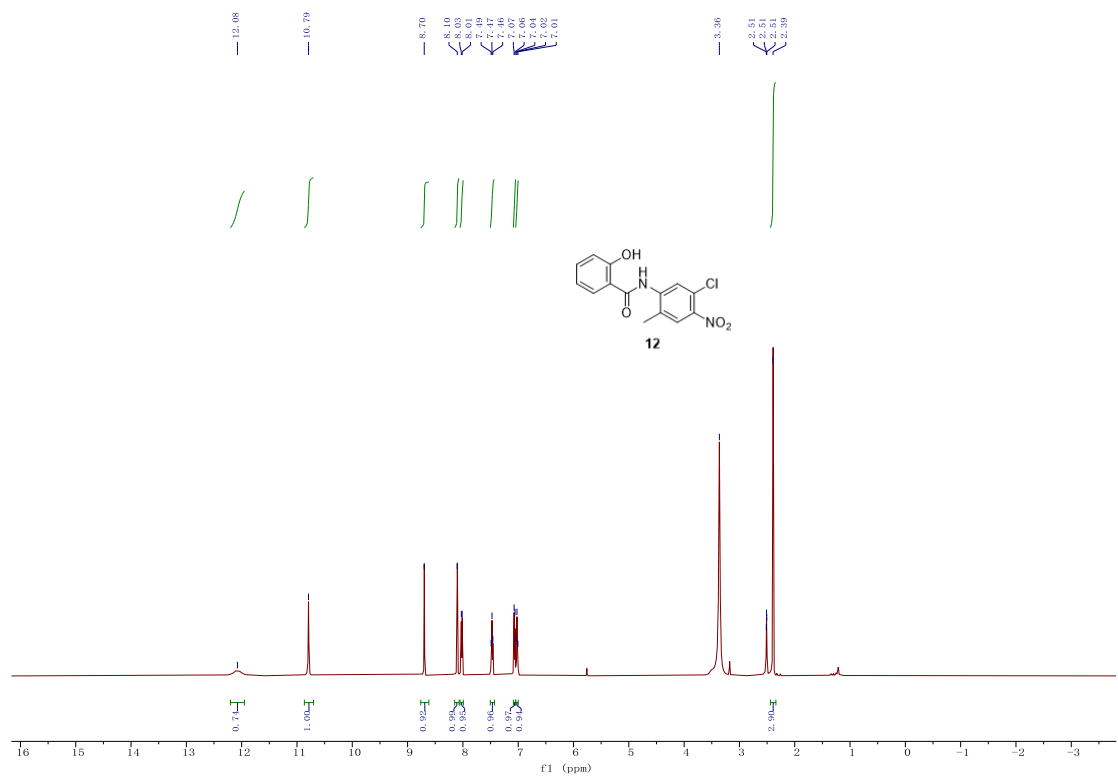
^b School of Public Health (Shenzhen), Sun Yat-sen University, Guangzhou, 510275,
China

^c Shenzhen Key Laboratory of Pathogenic Microbes and Biosafety, Shenzhen Campus
of Sun Yat-sen University, Shenzhen, 518107, China

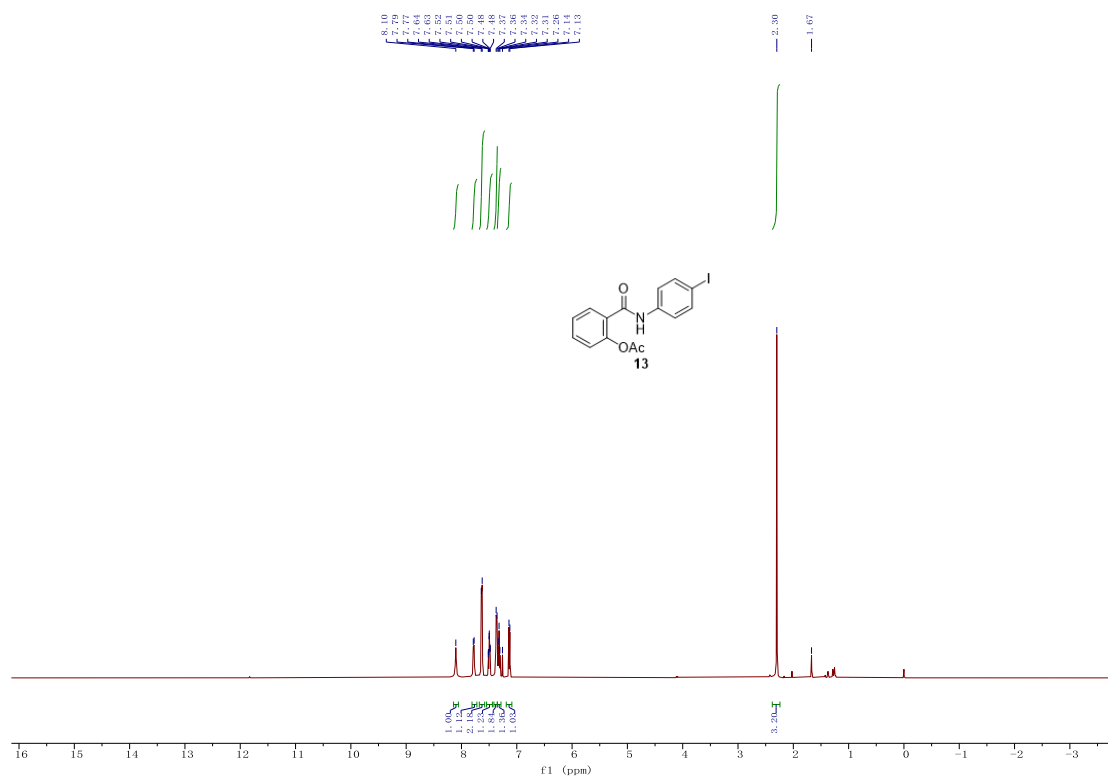
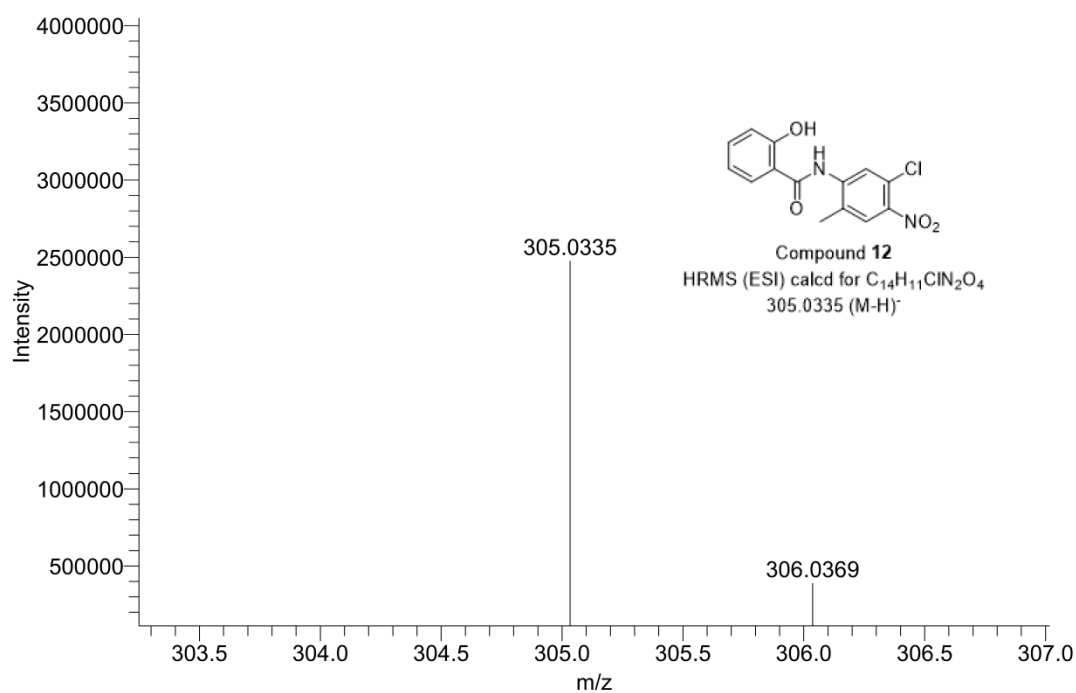
[†] These authors contributed equally to this work

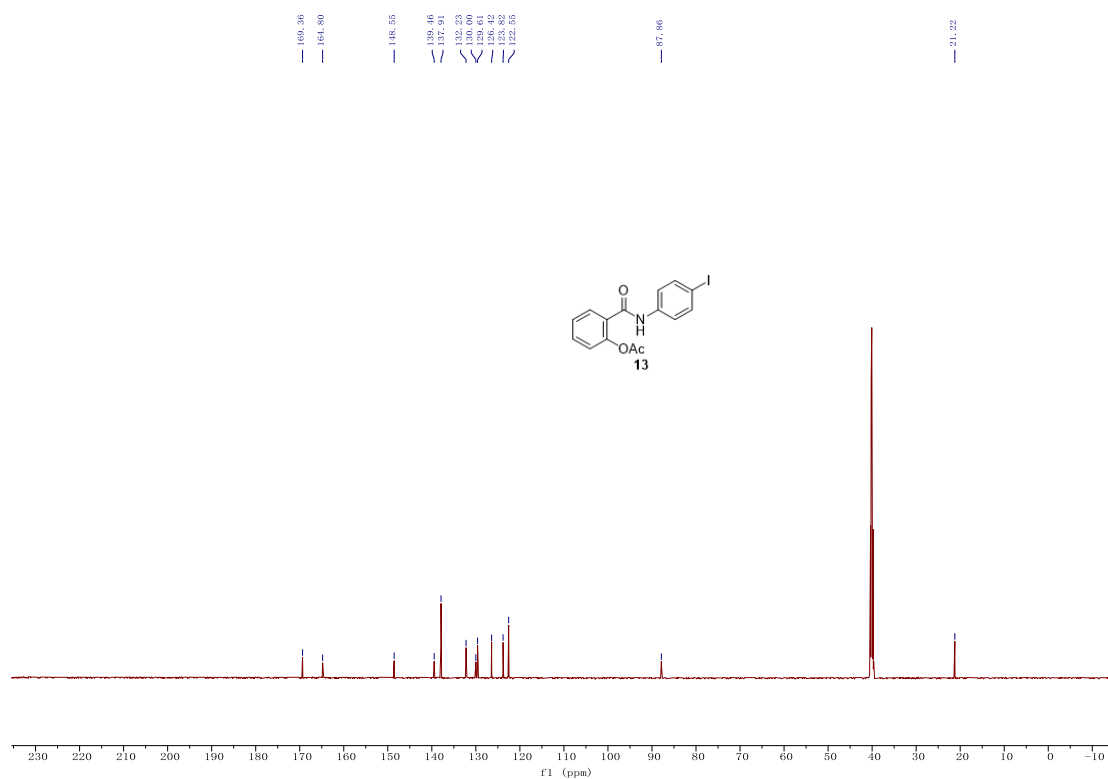
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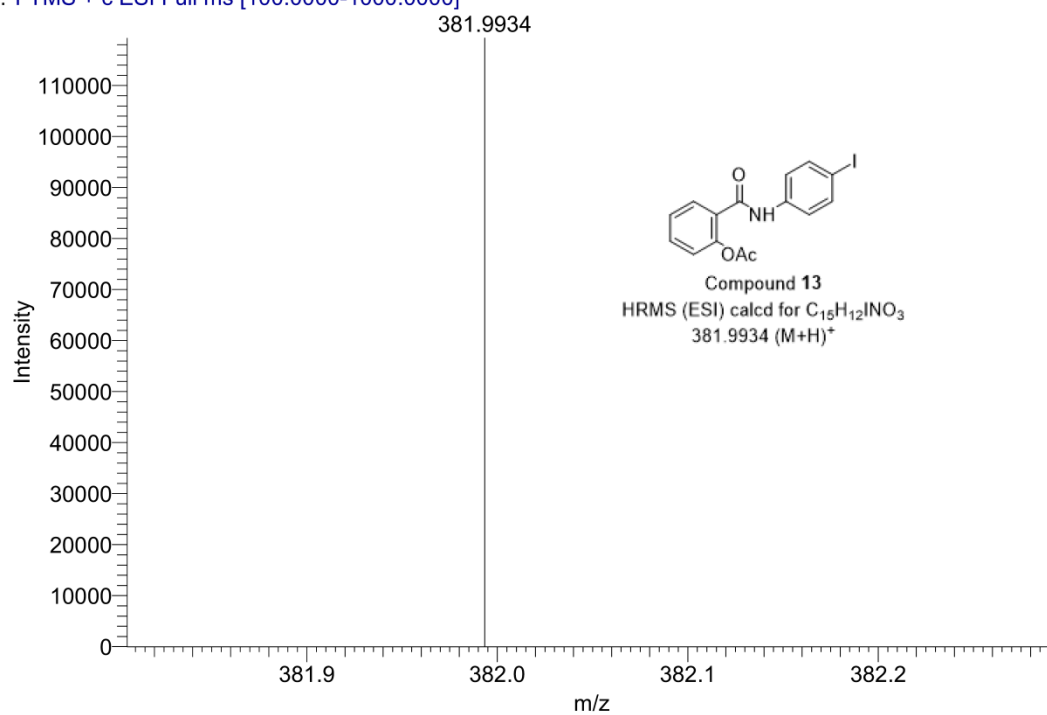


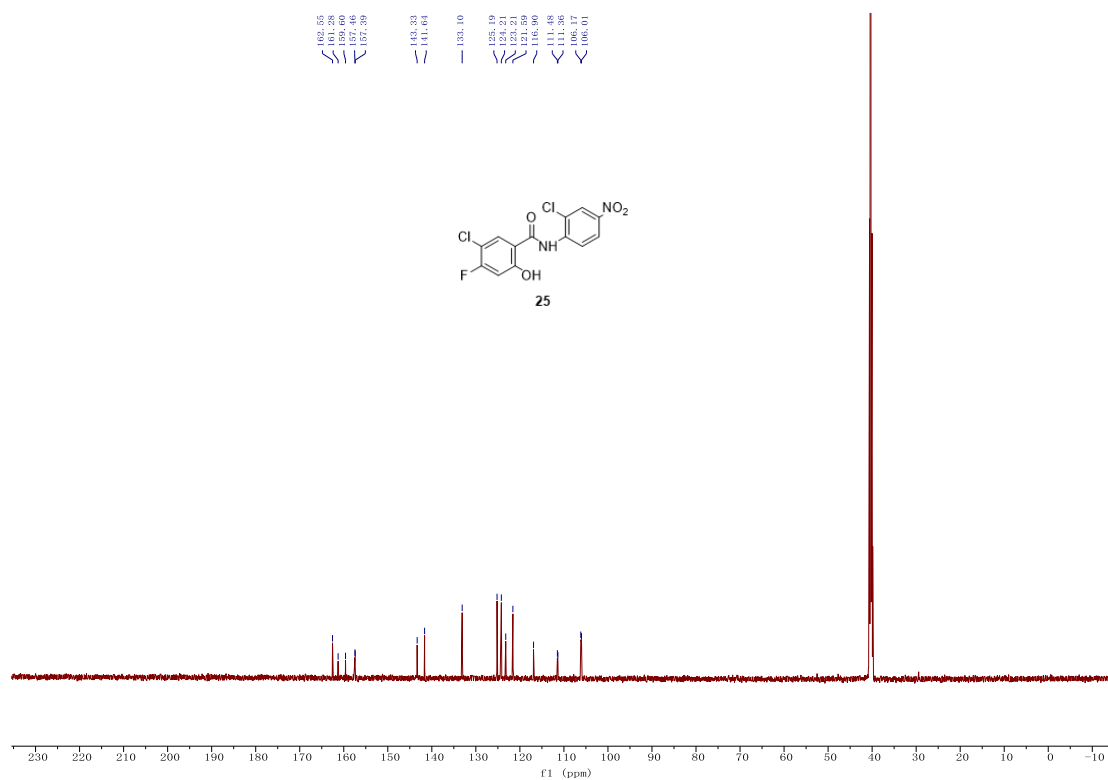
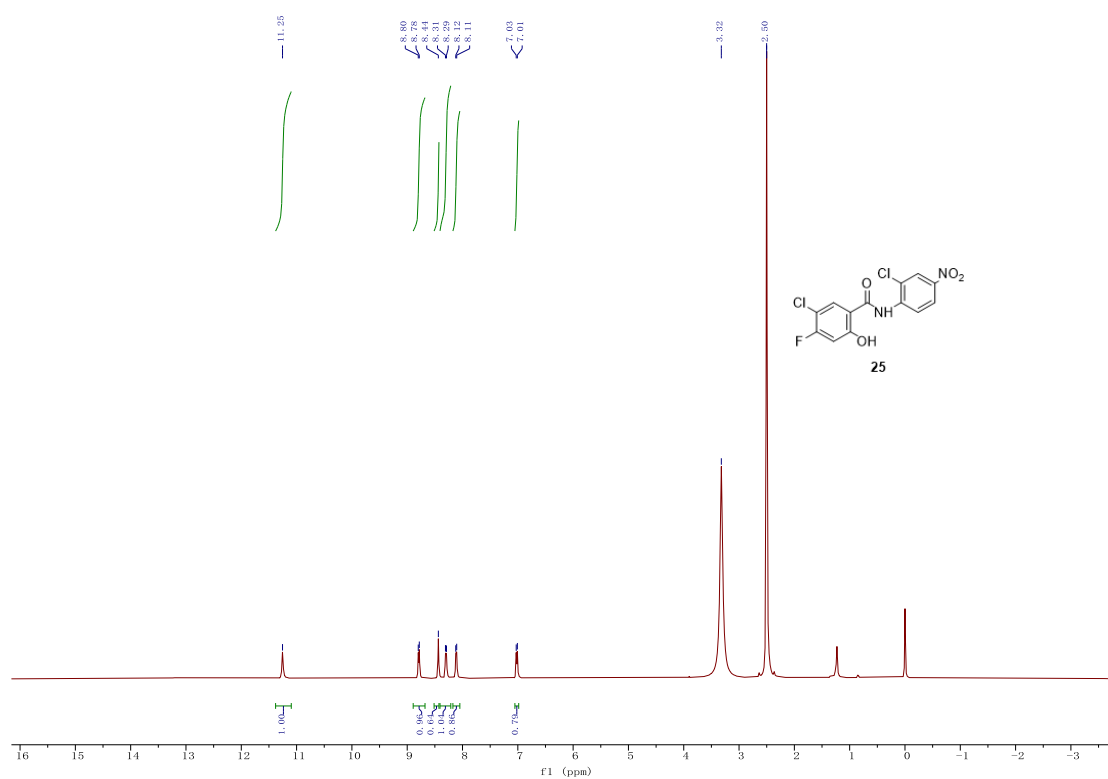
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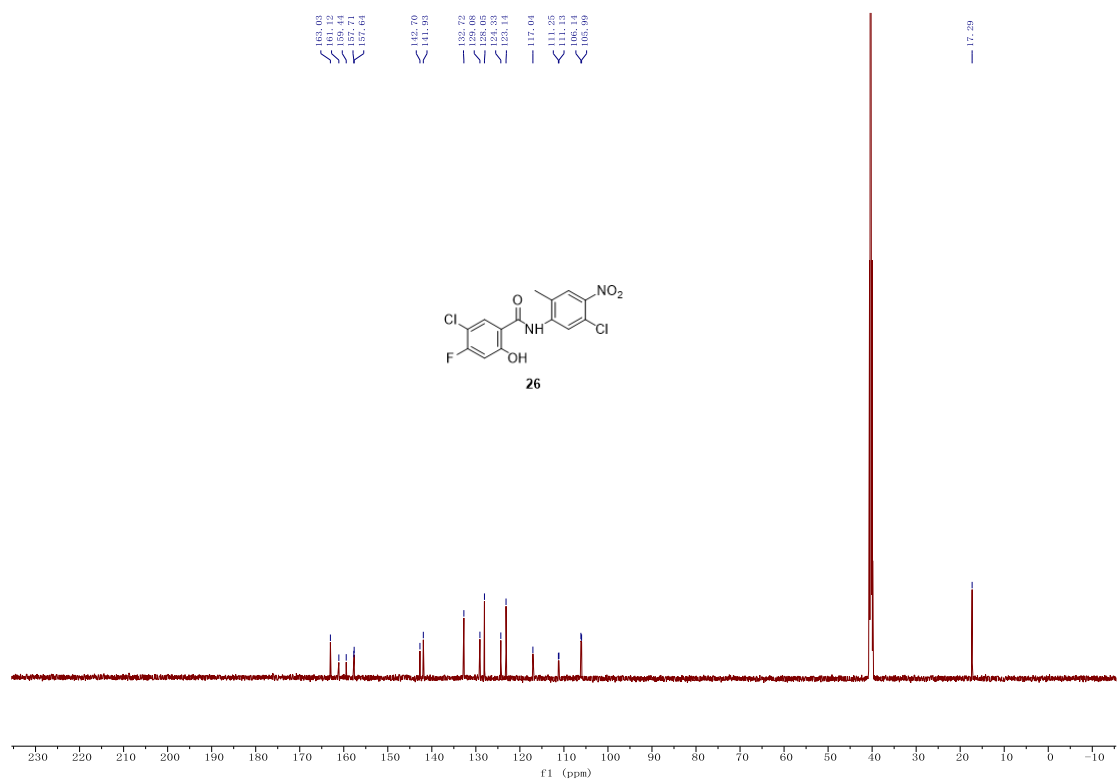
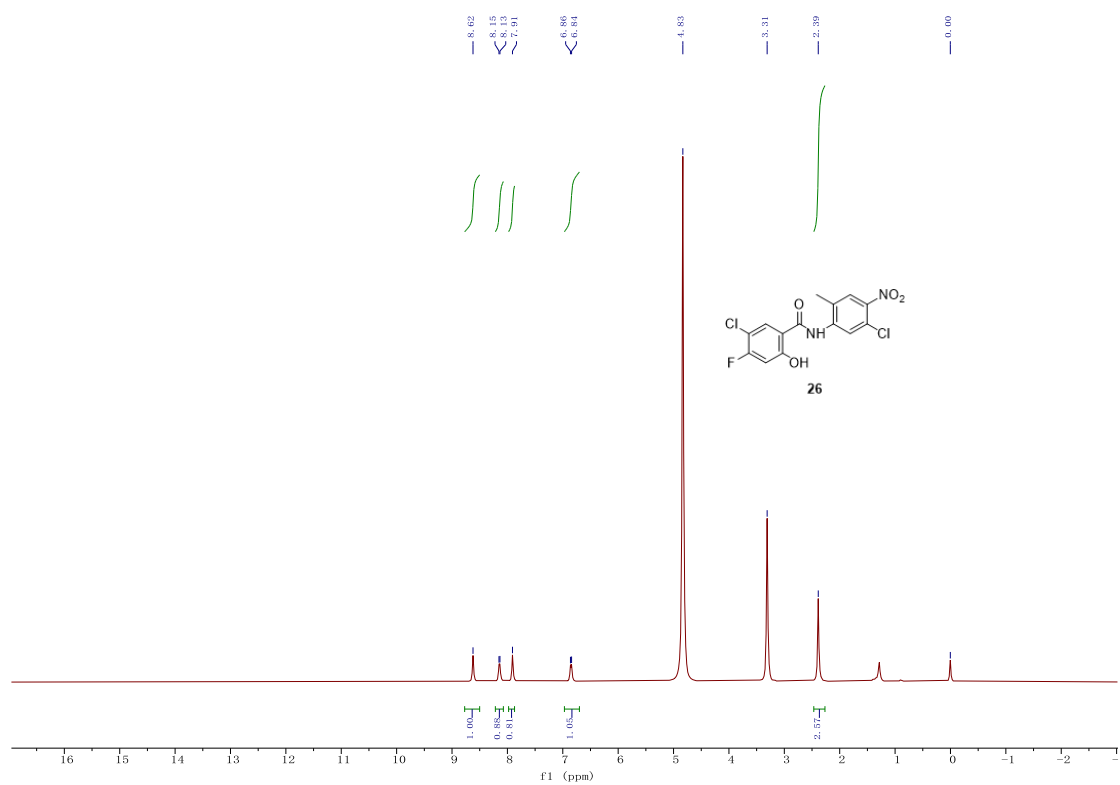




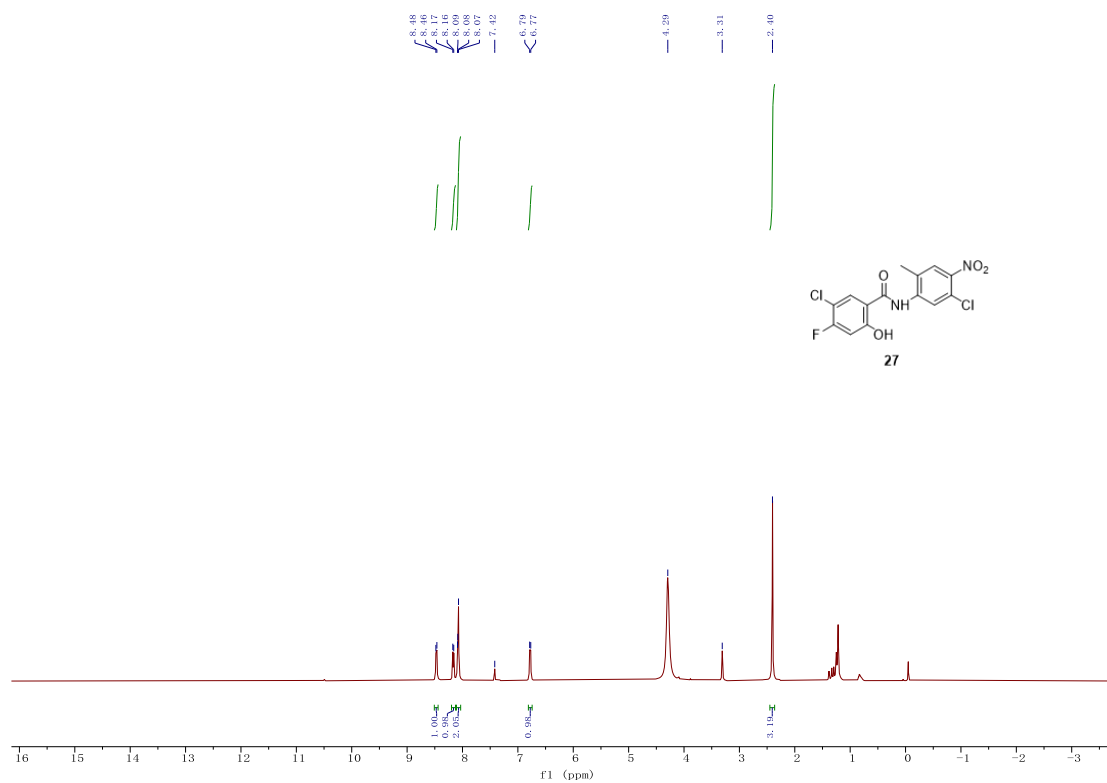
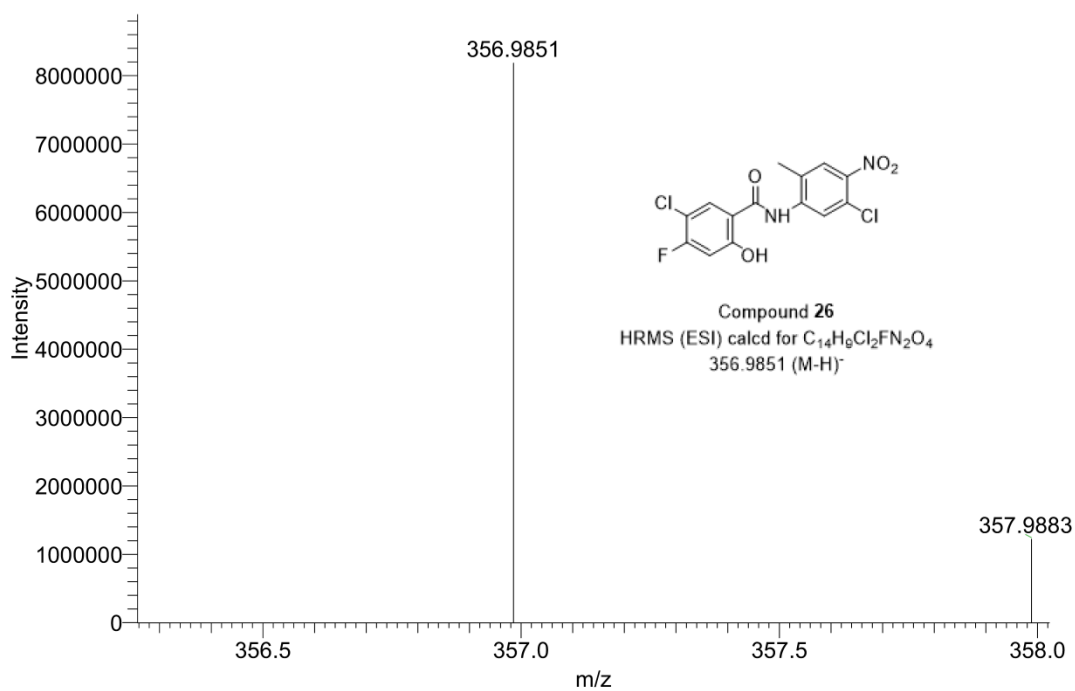
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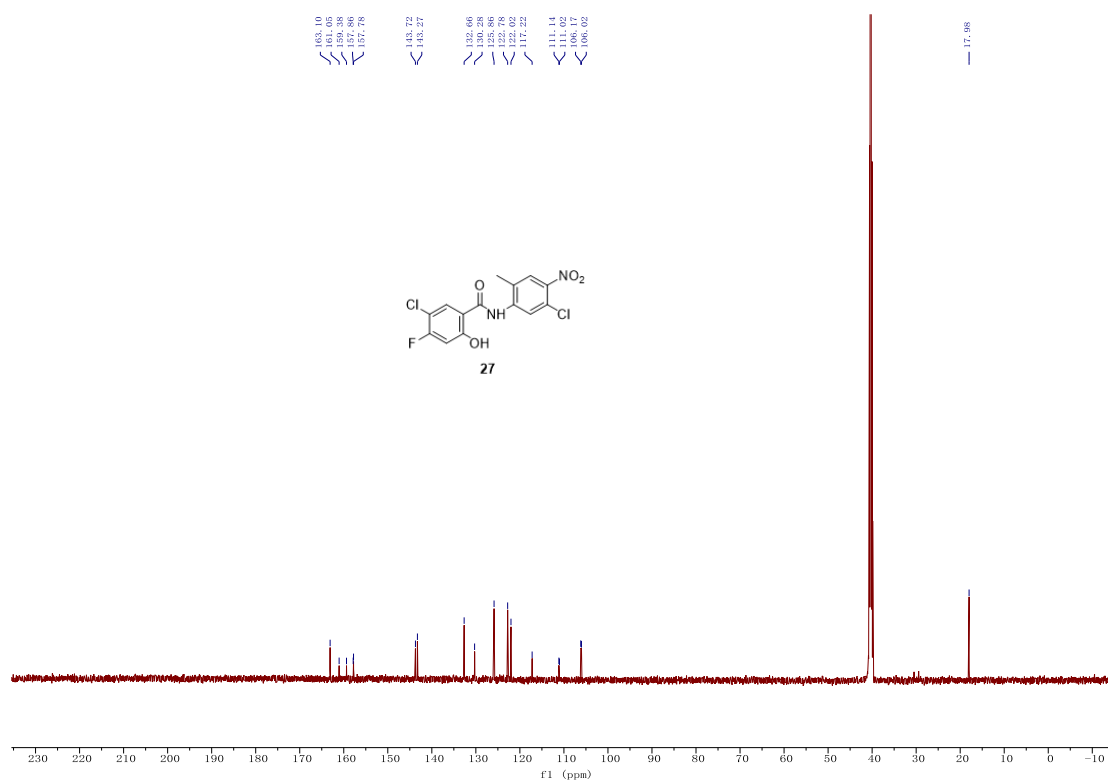




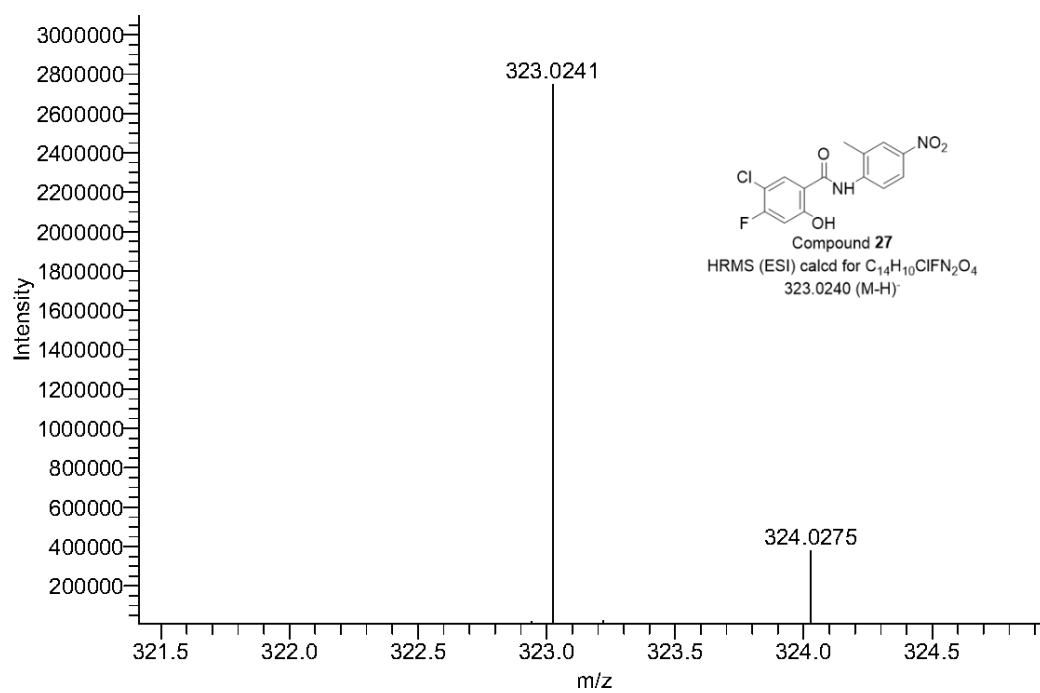


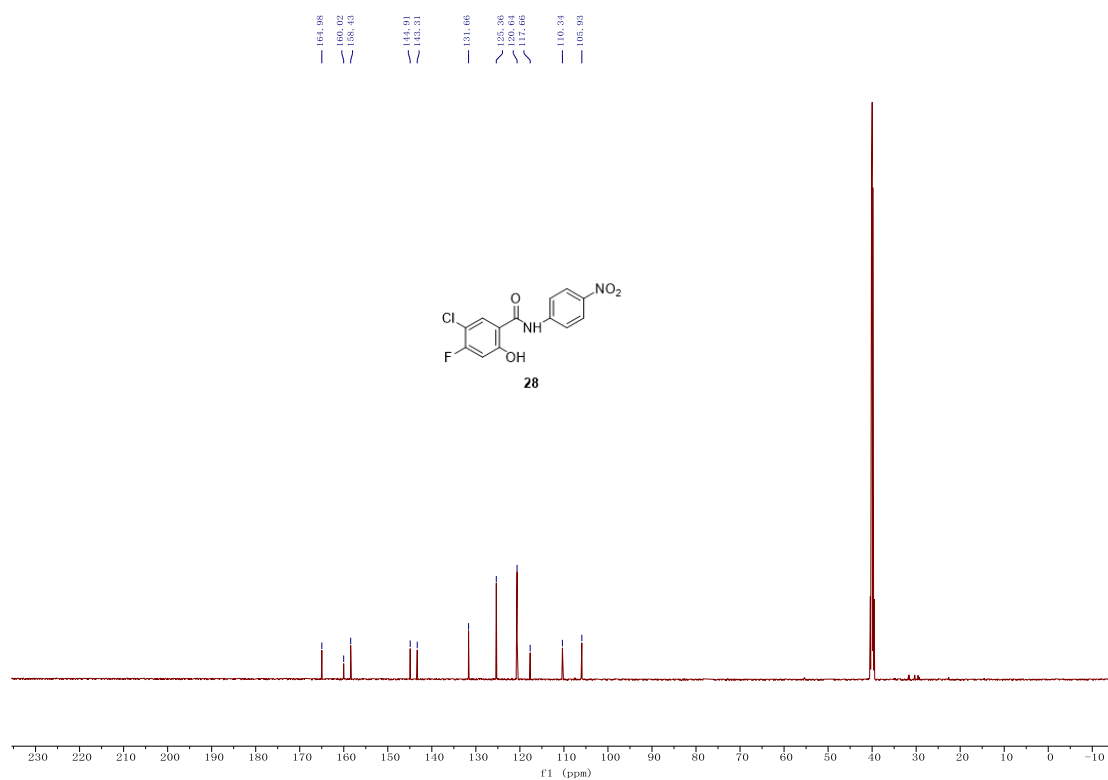
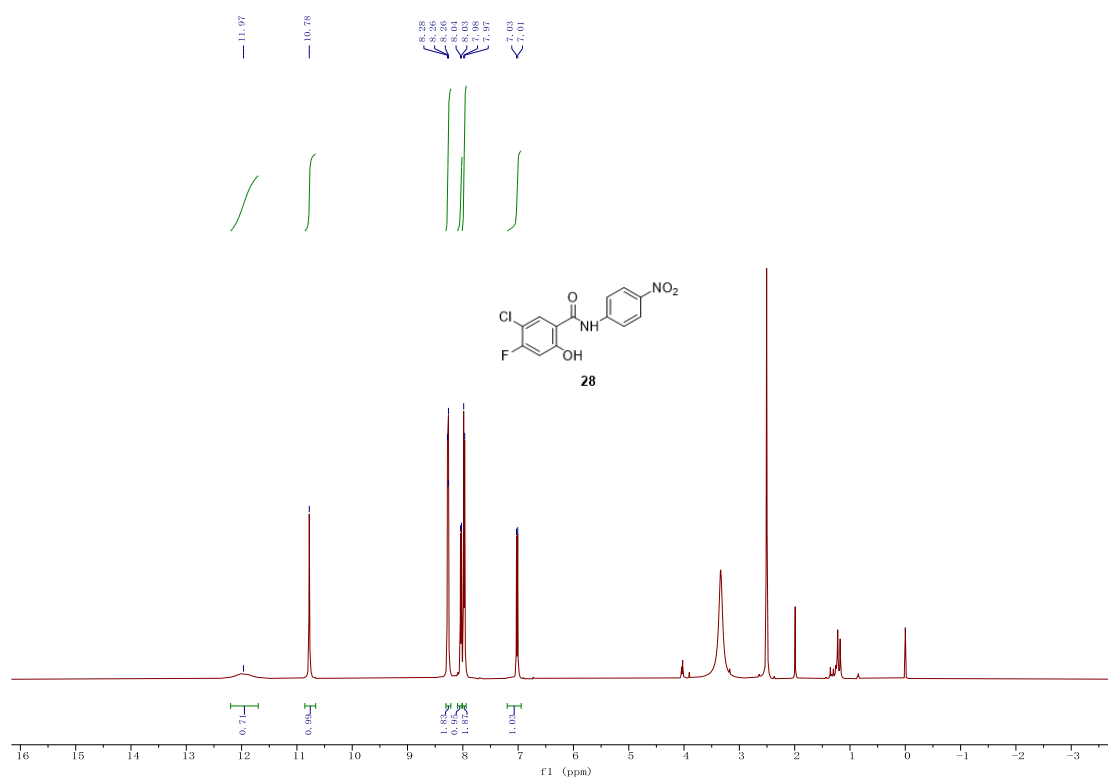
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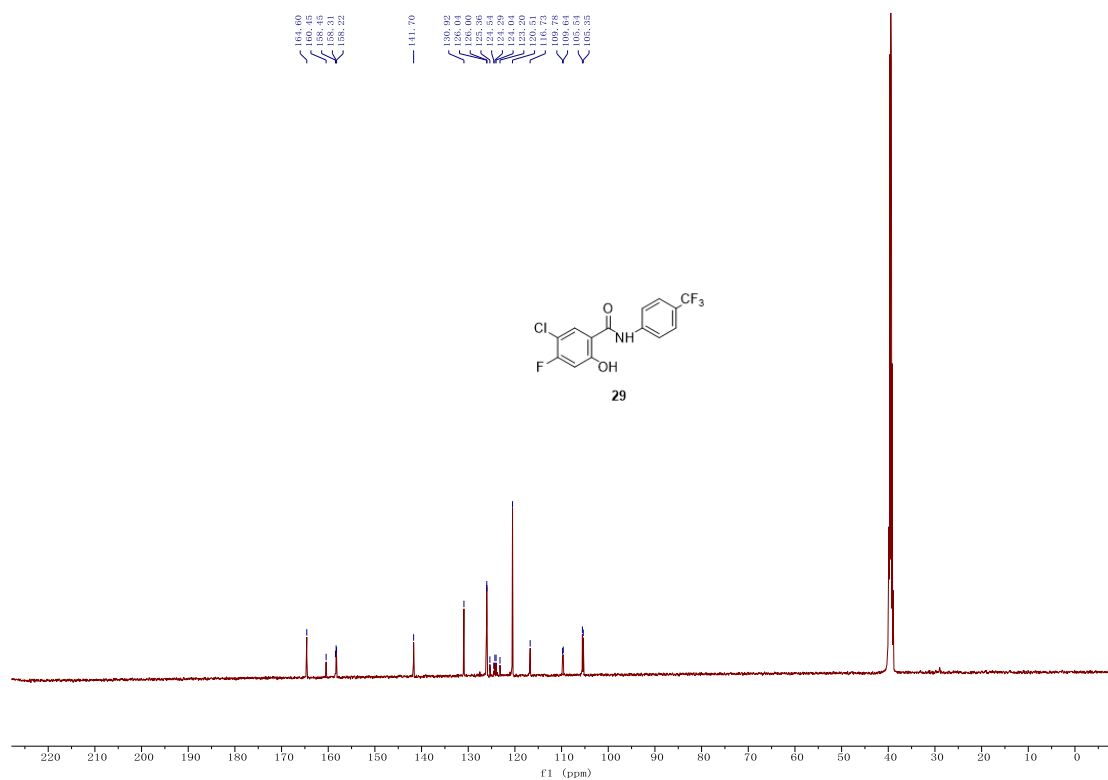
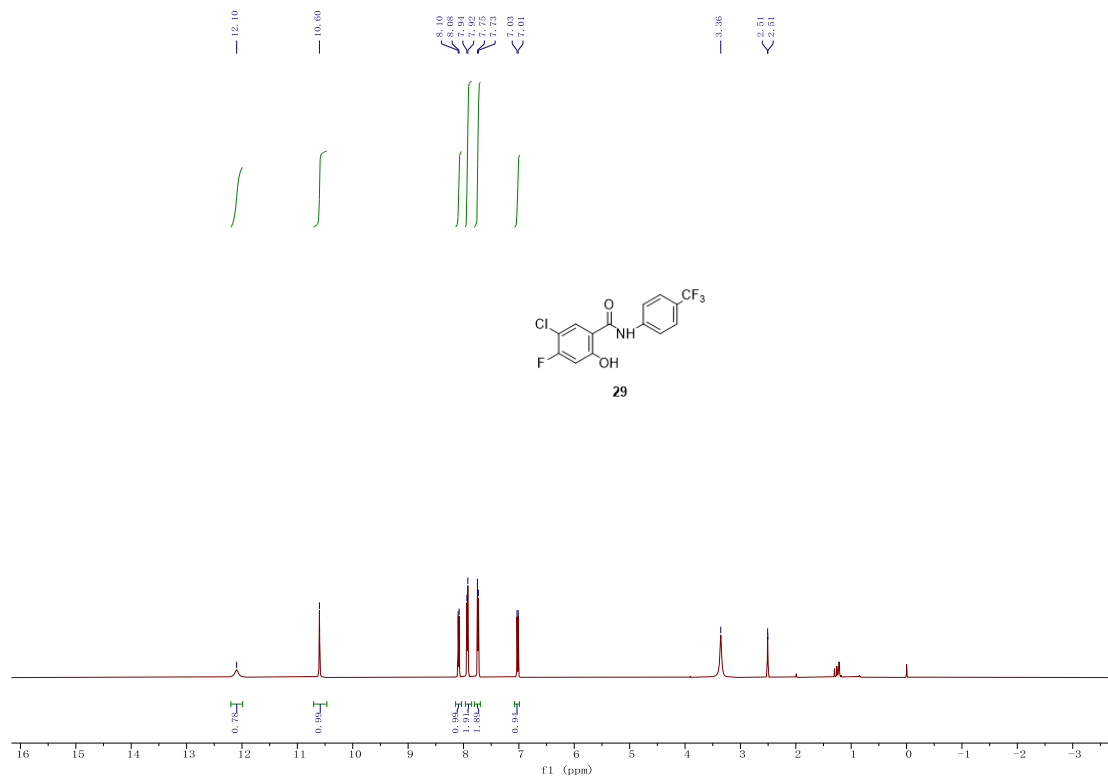


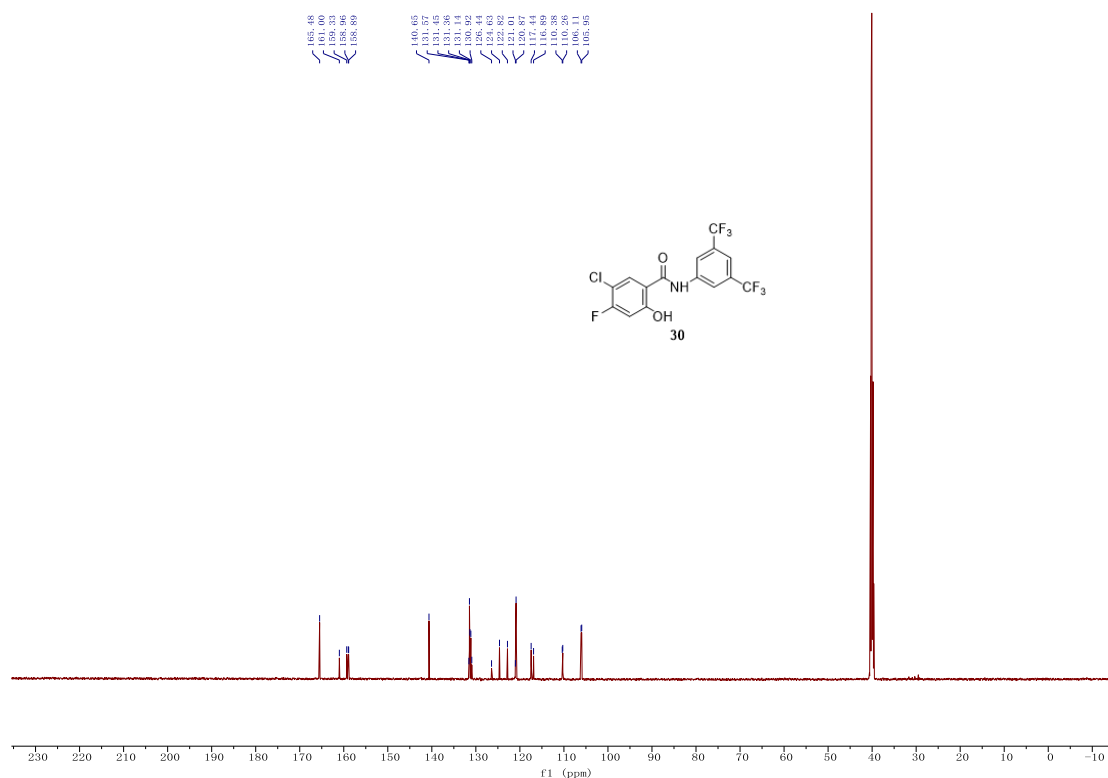
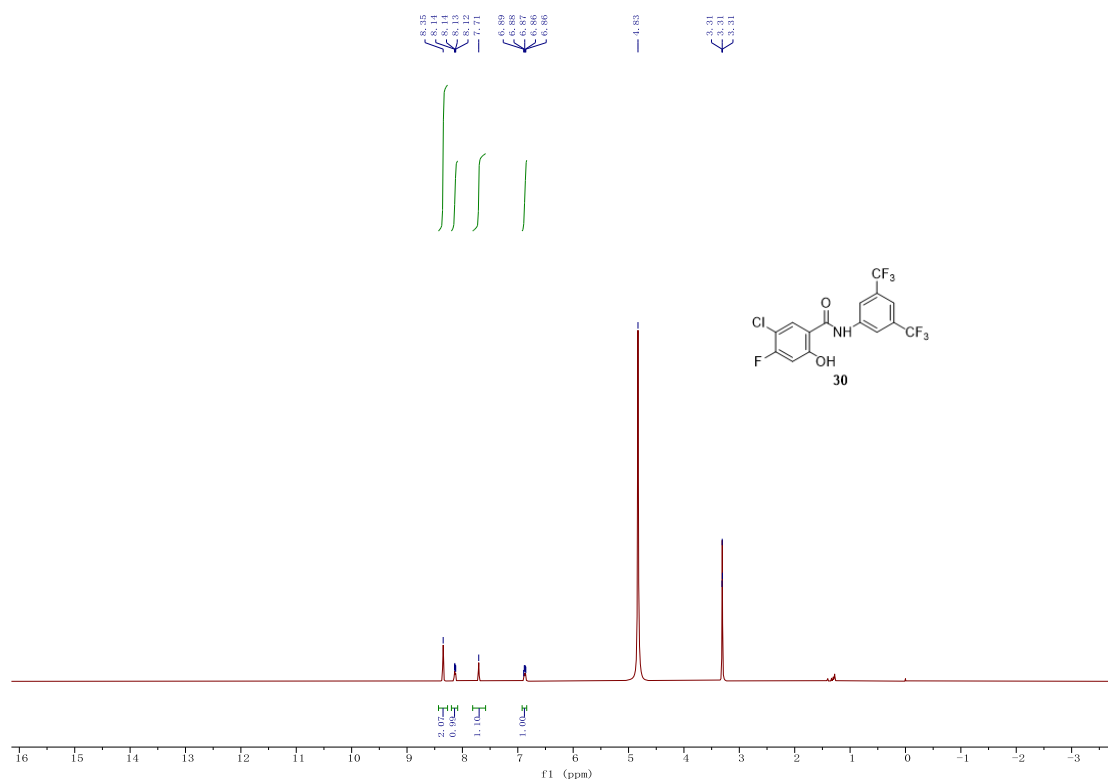


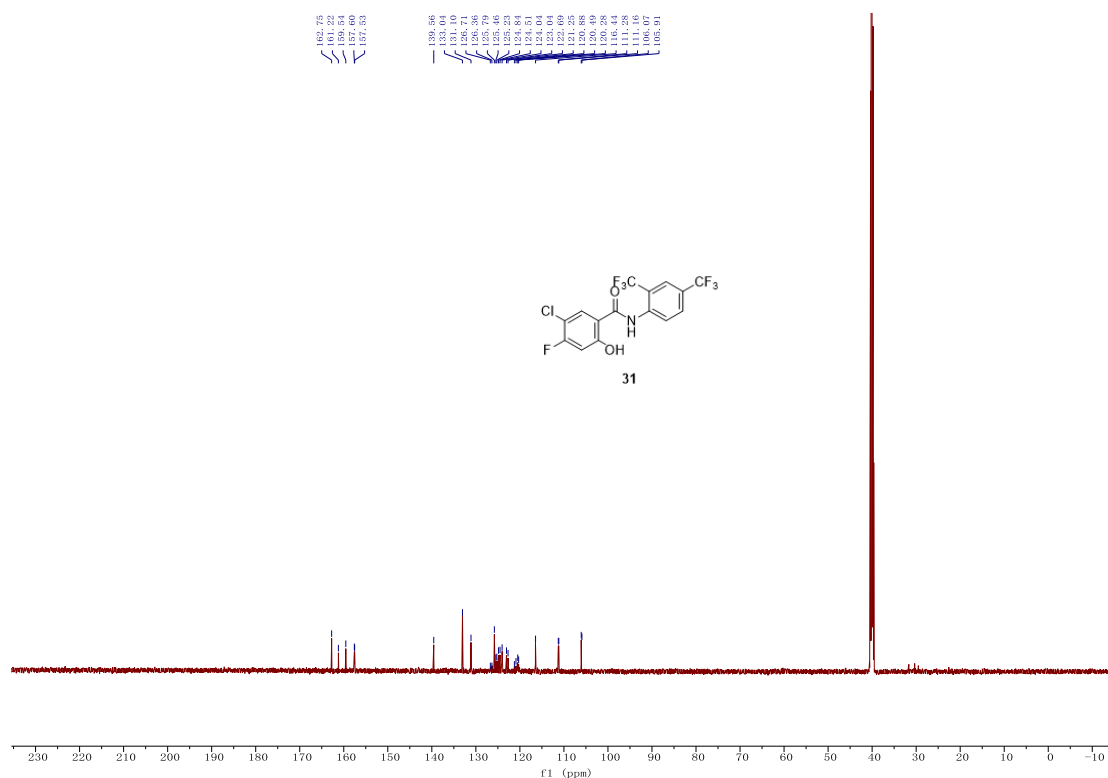
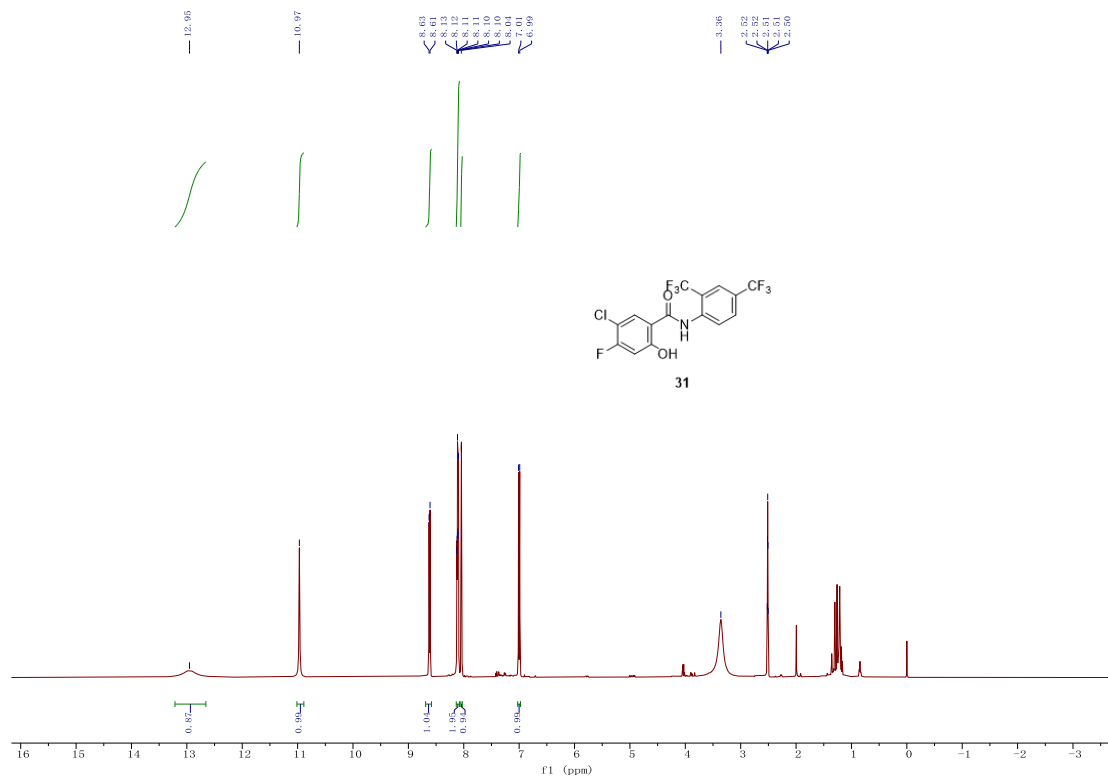
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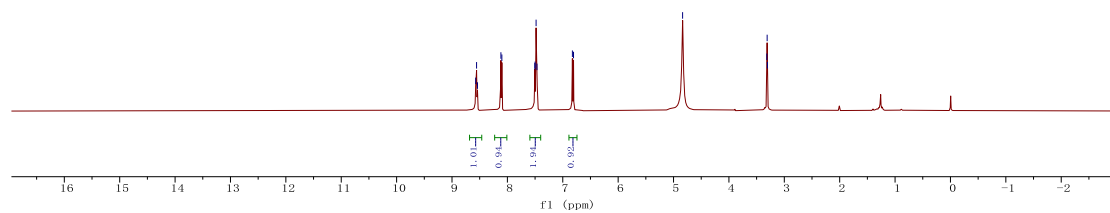
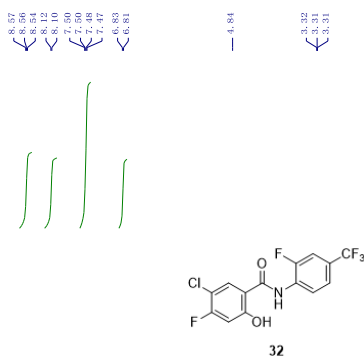
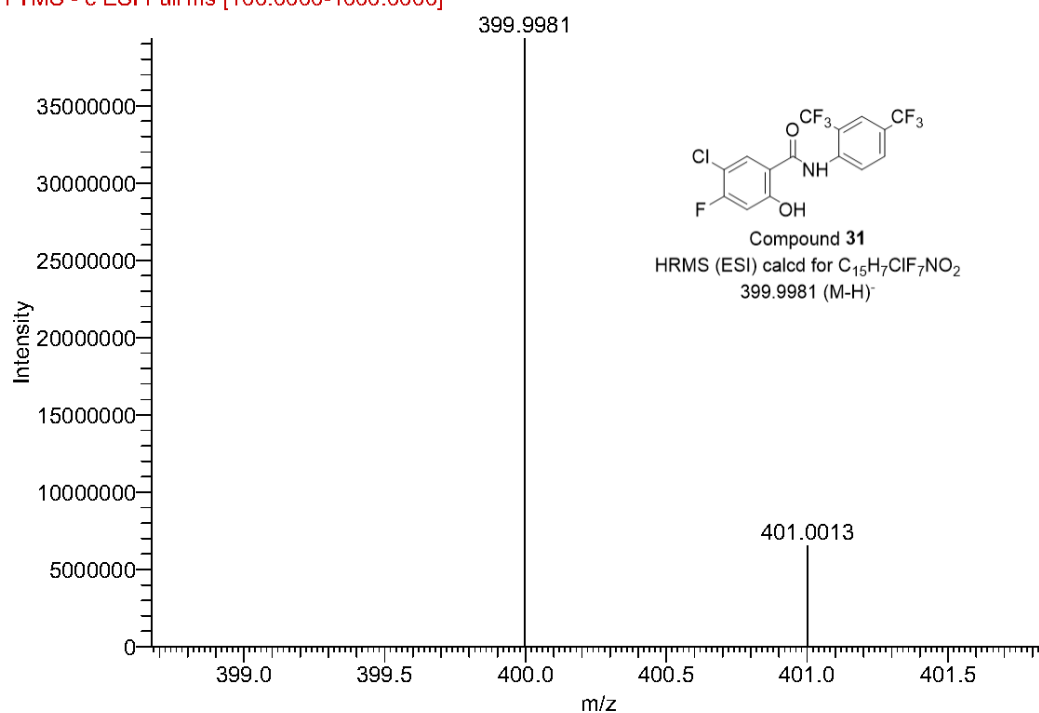


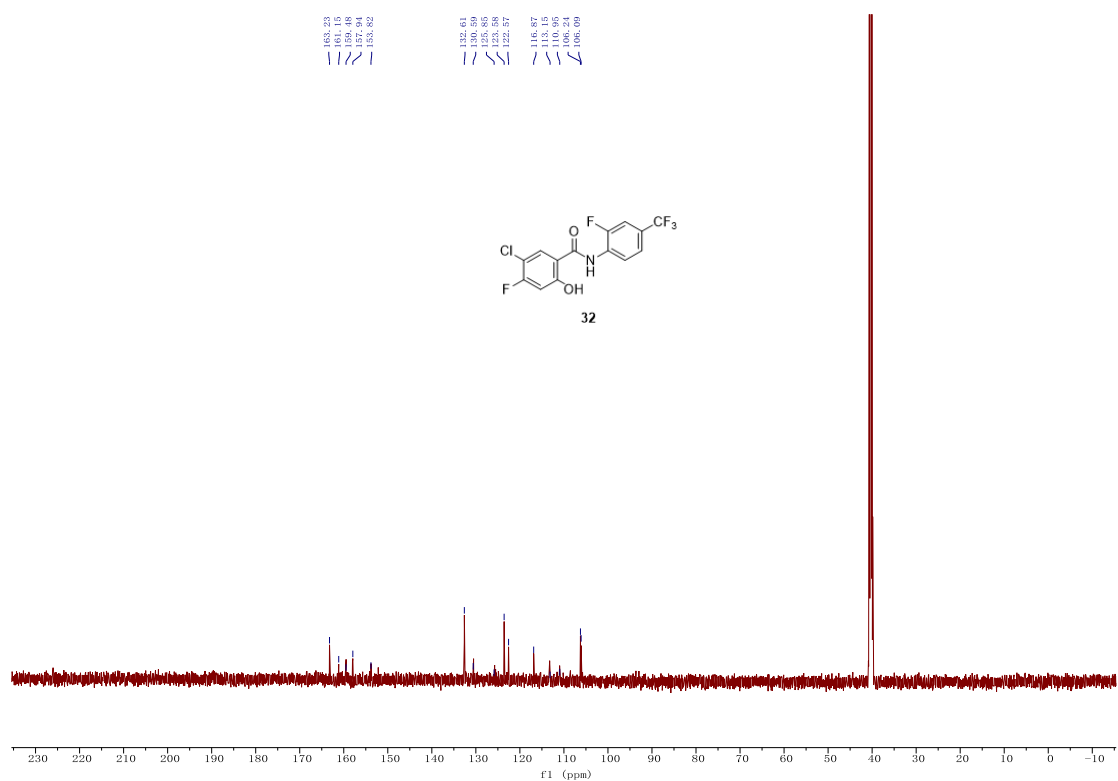




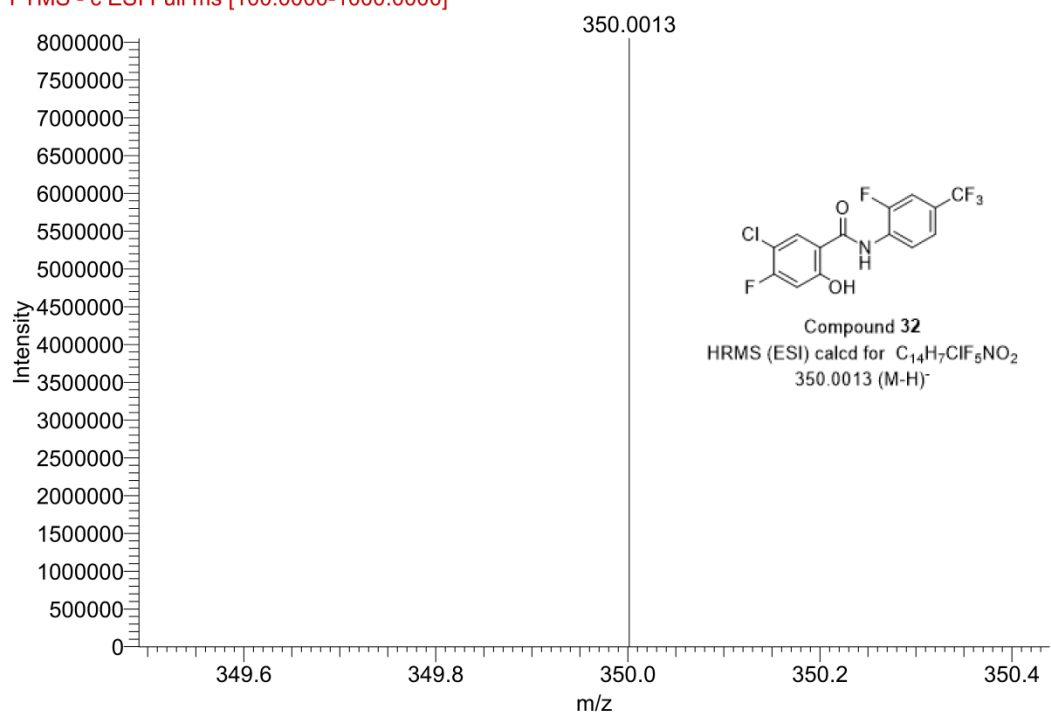
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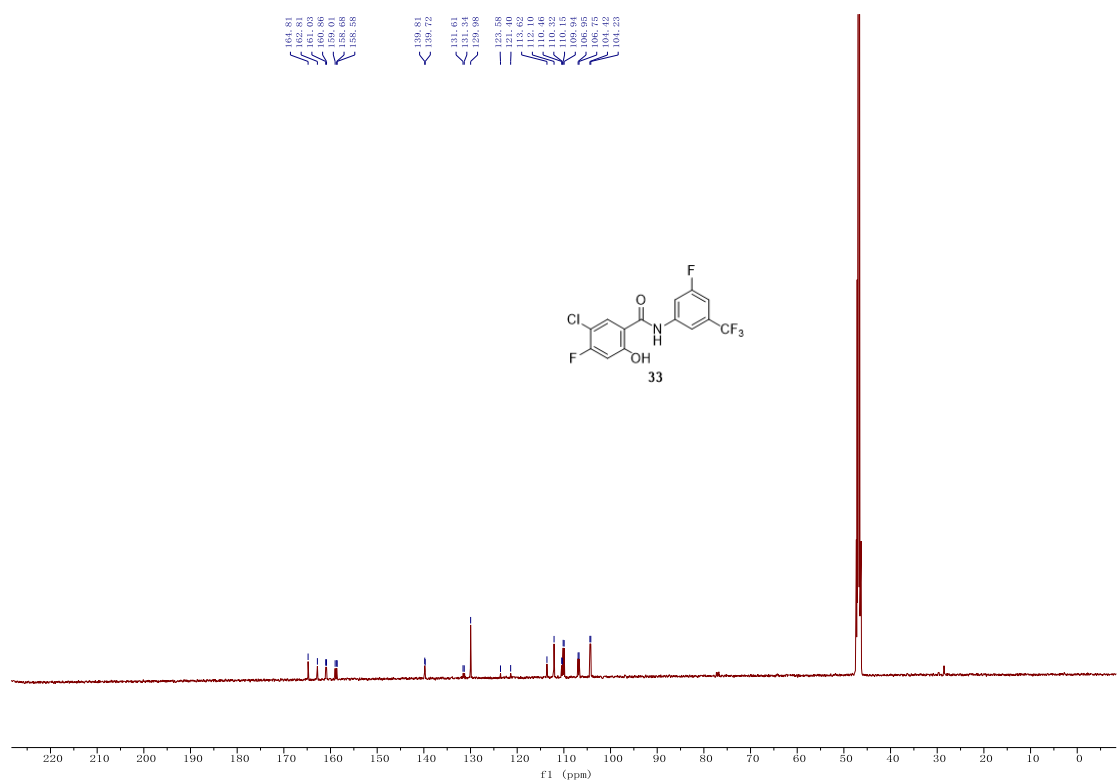
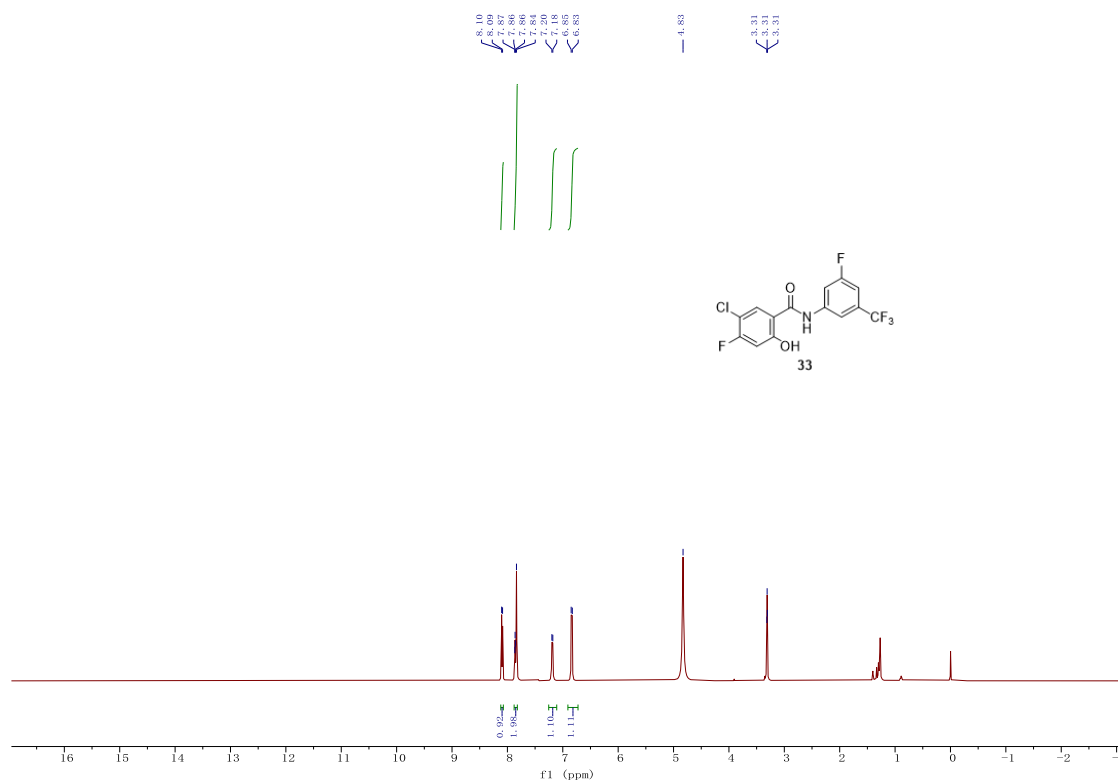
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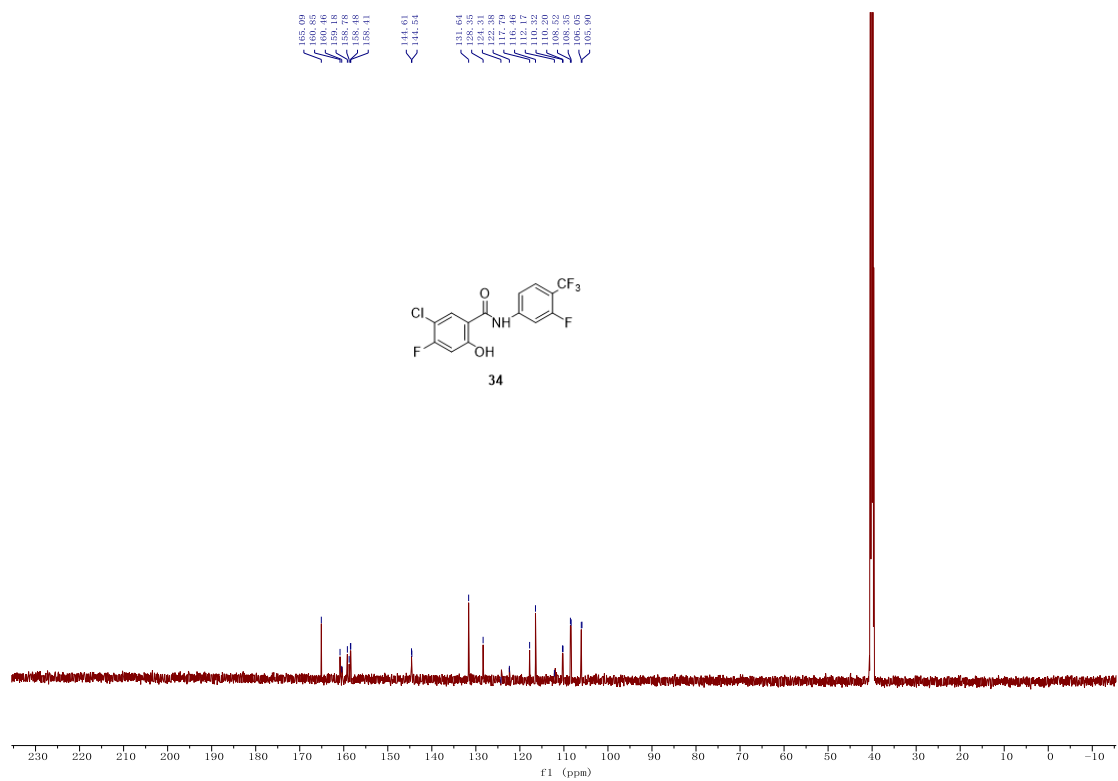
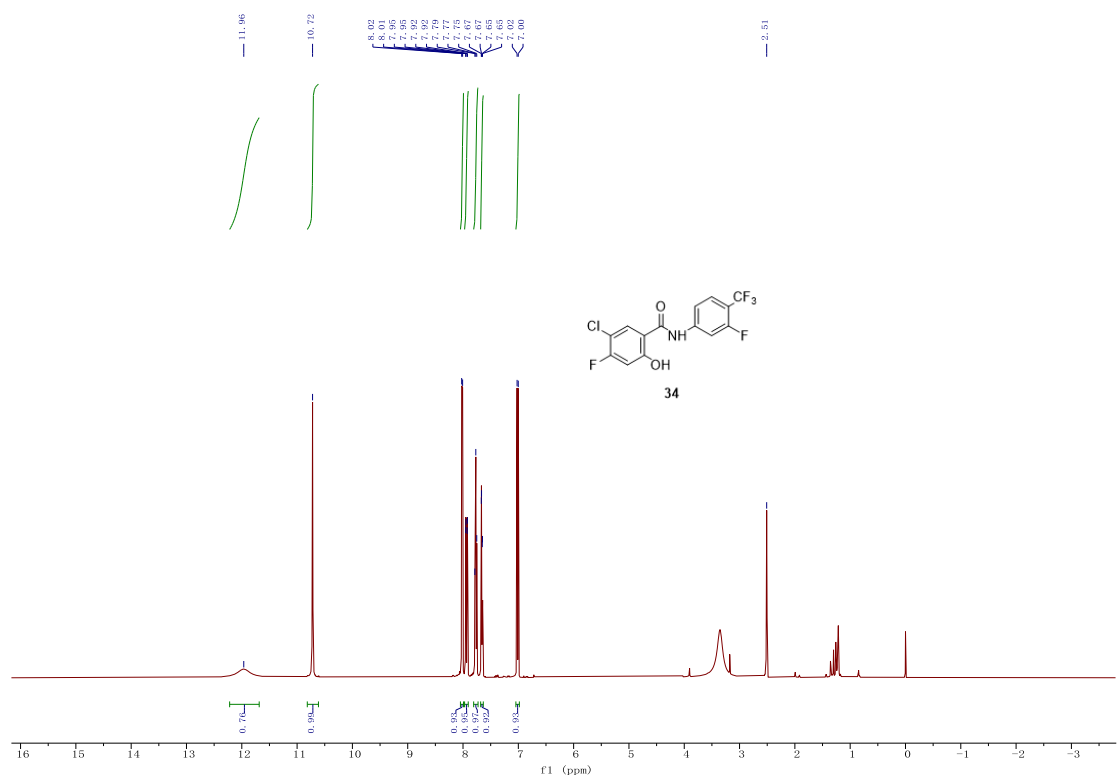


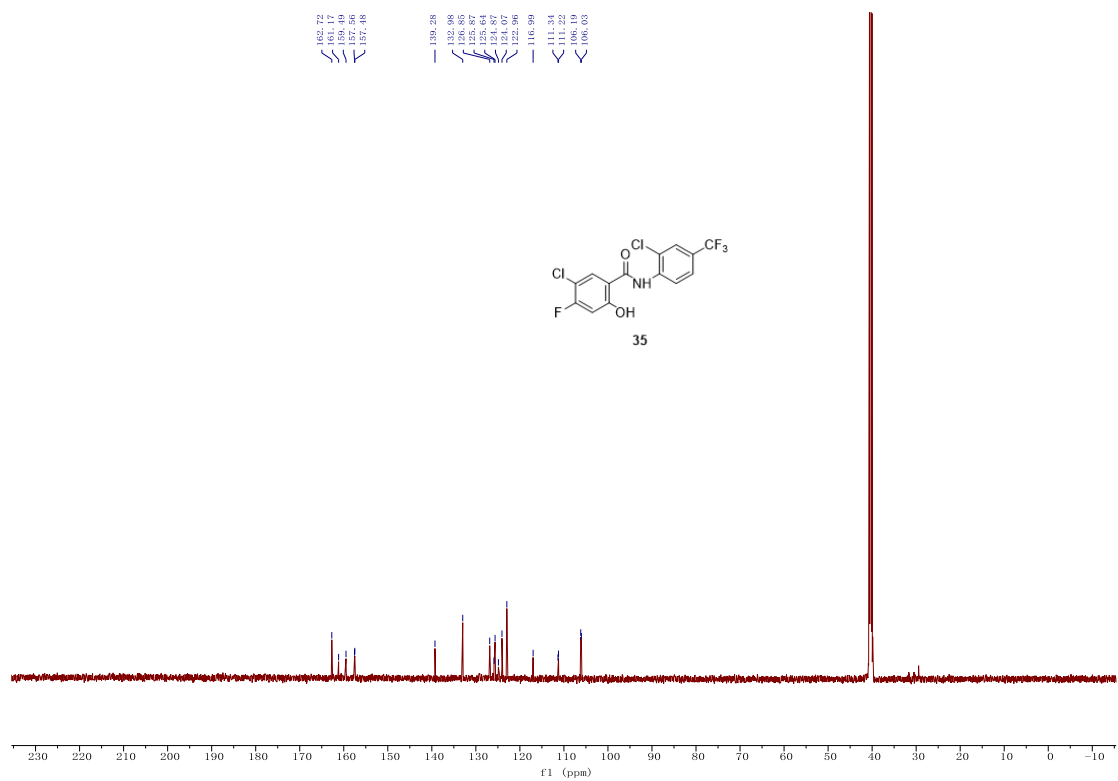
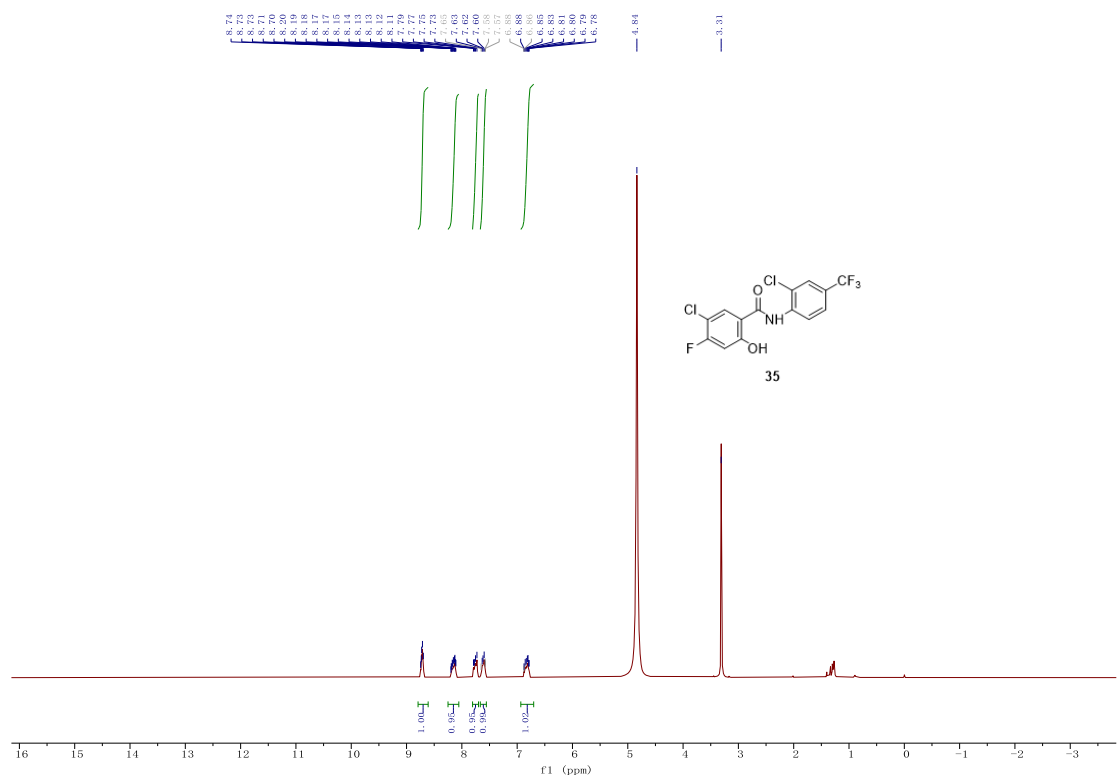


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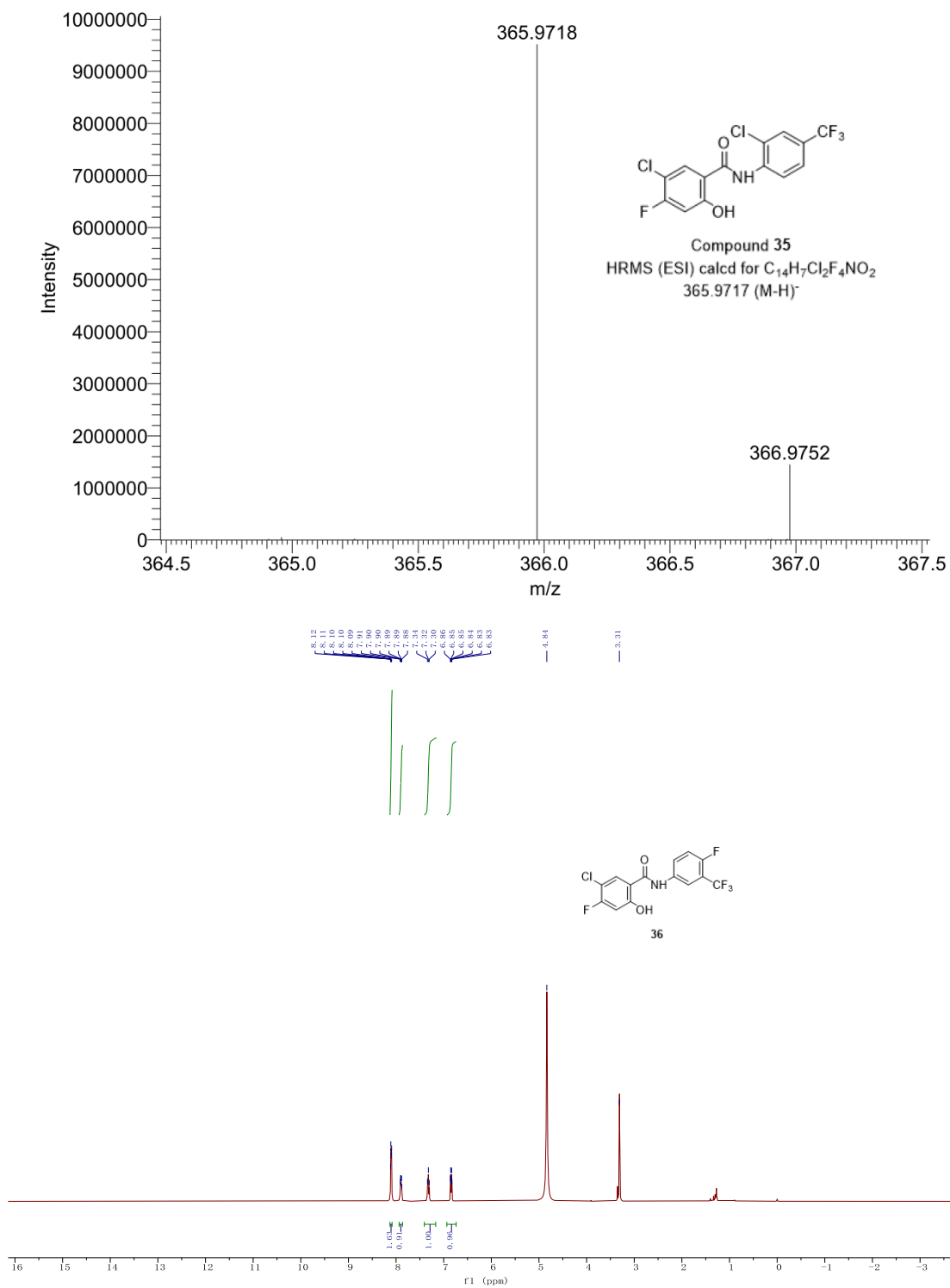


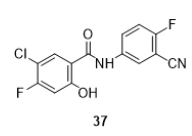
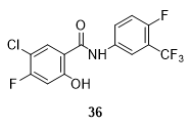


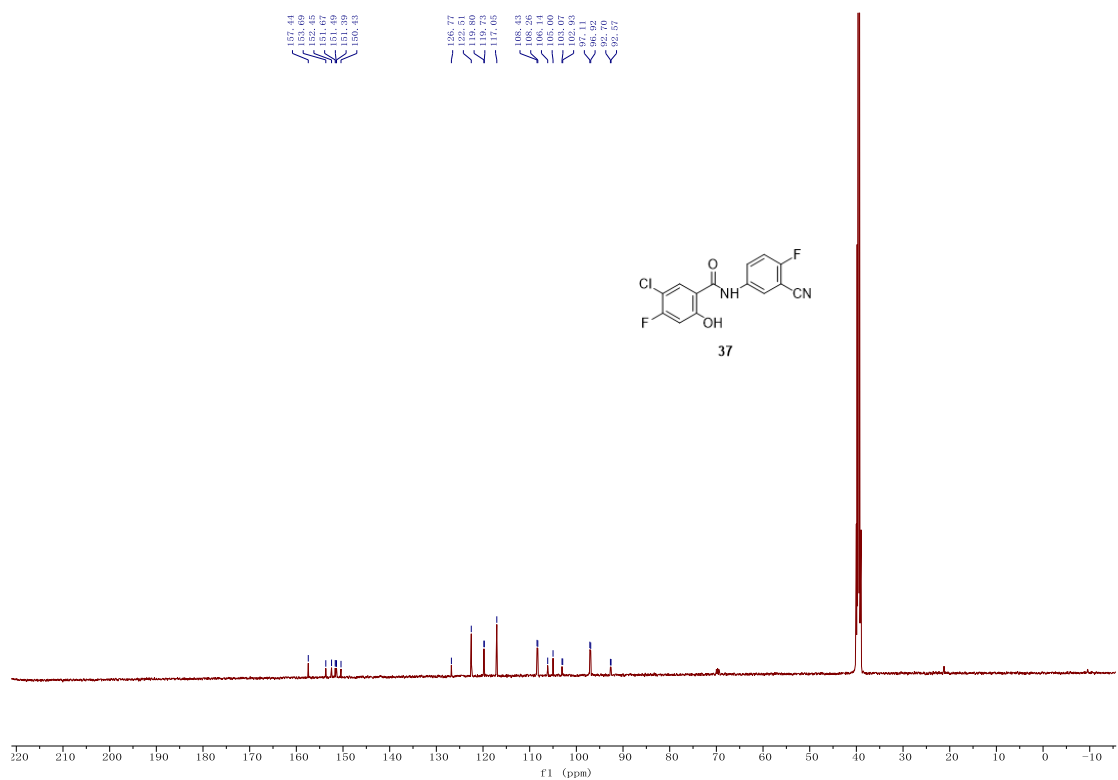




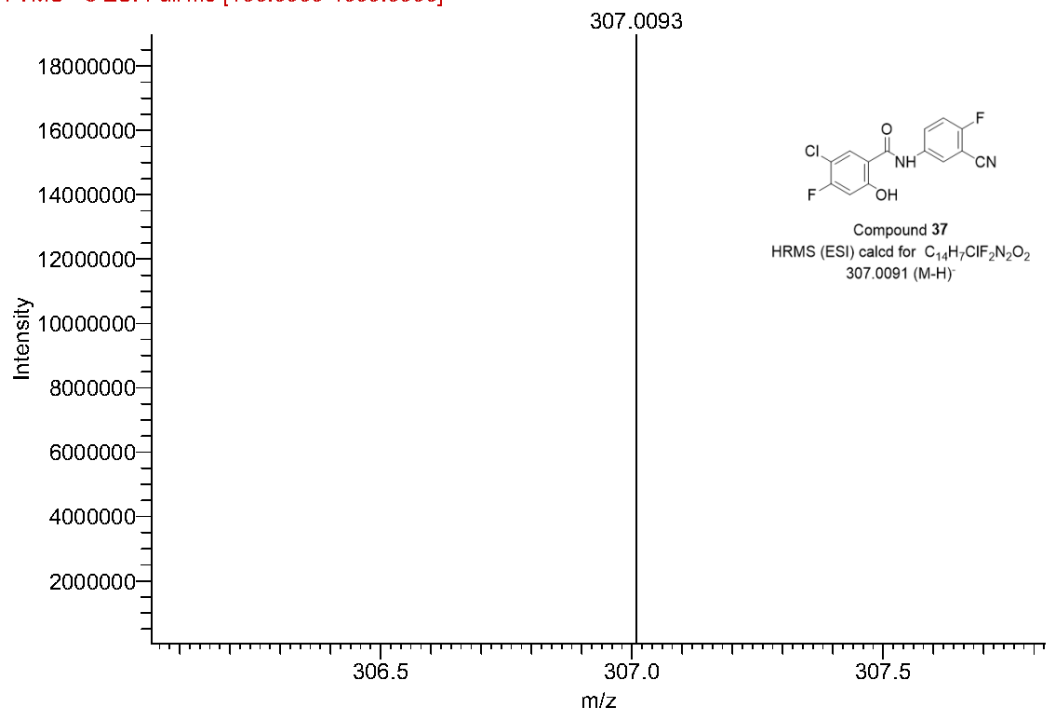
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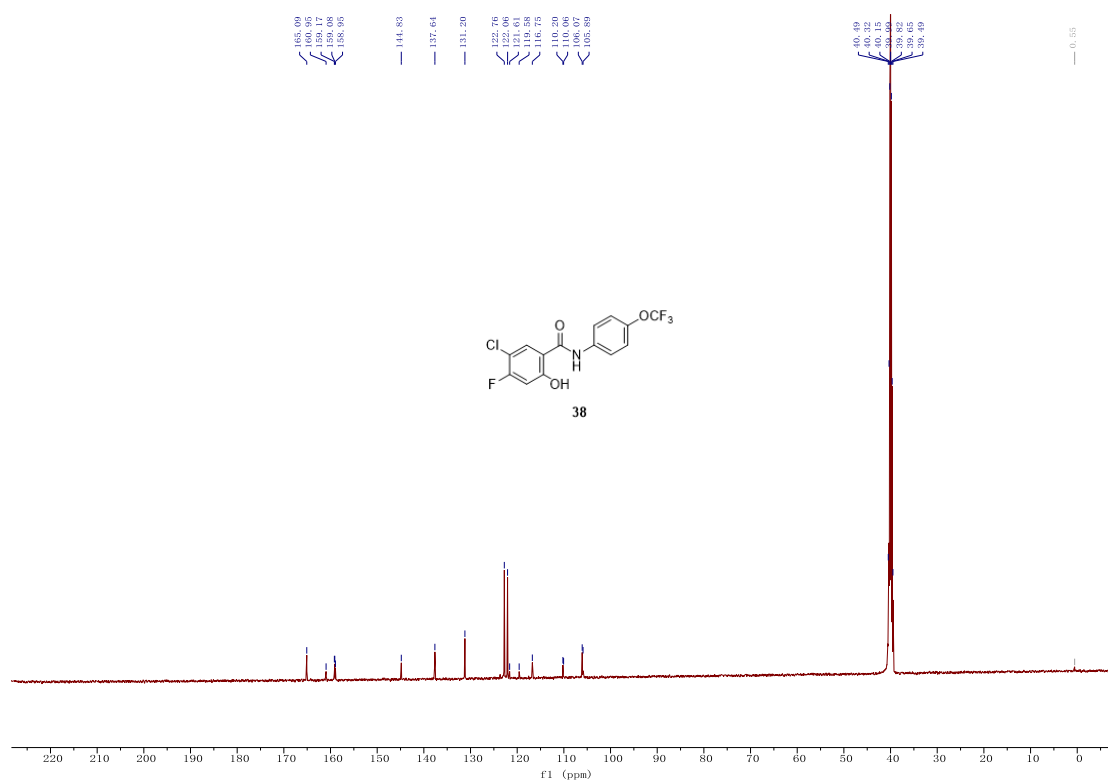
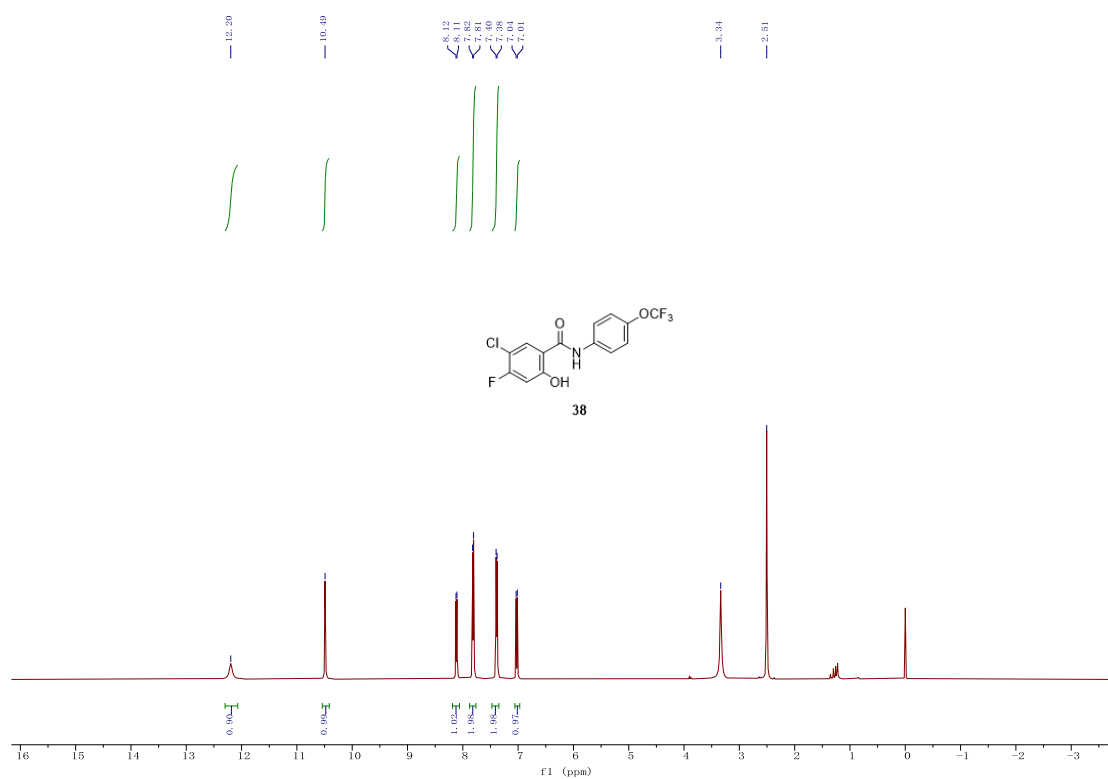


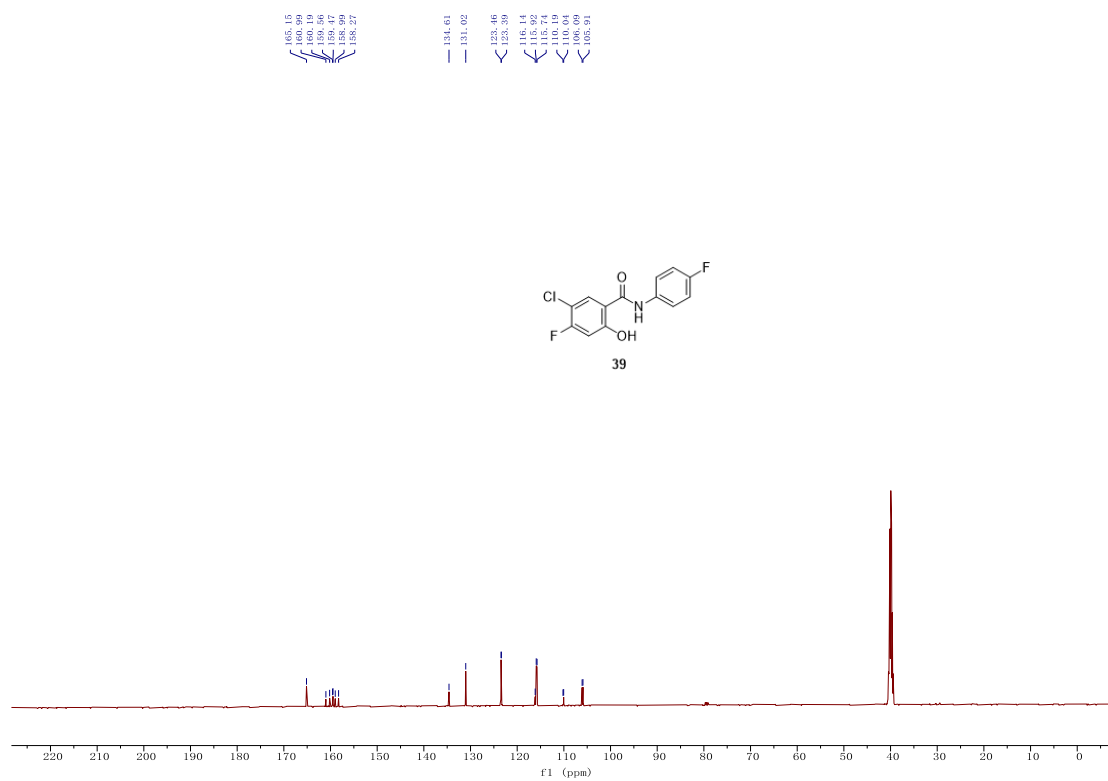
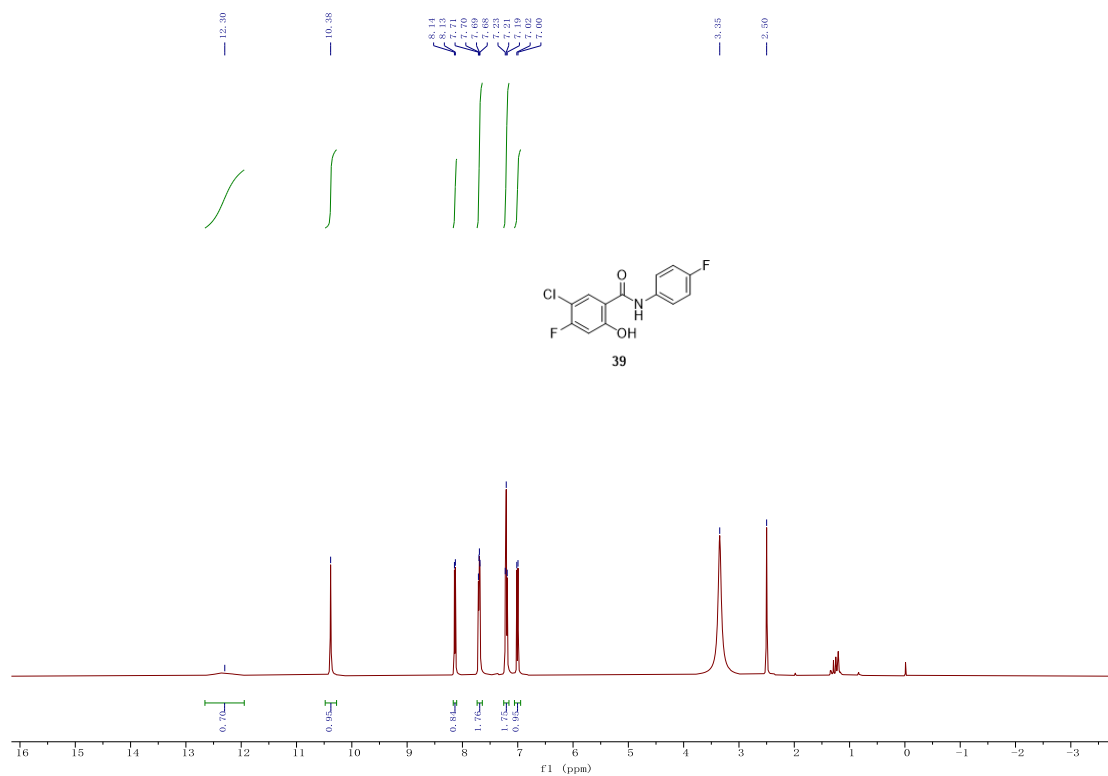




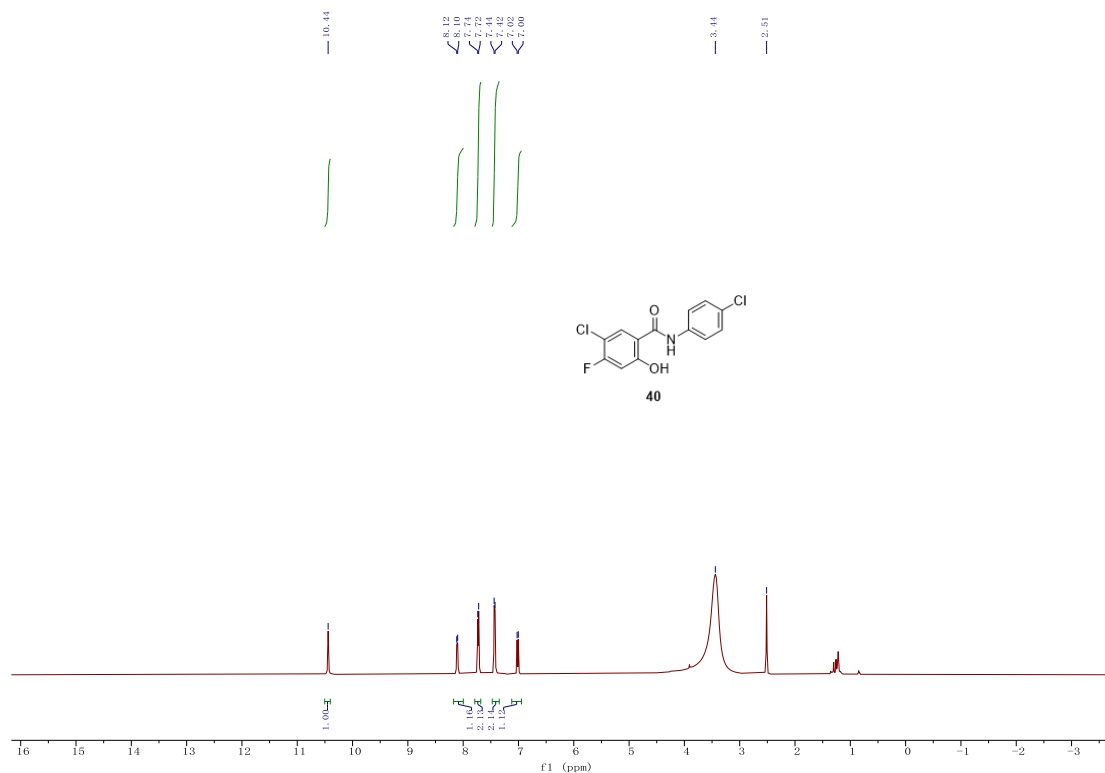
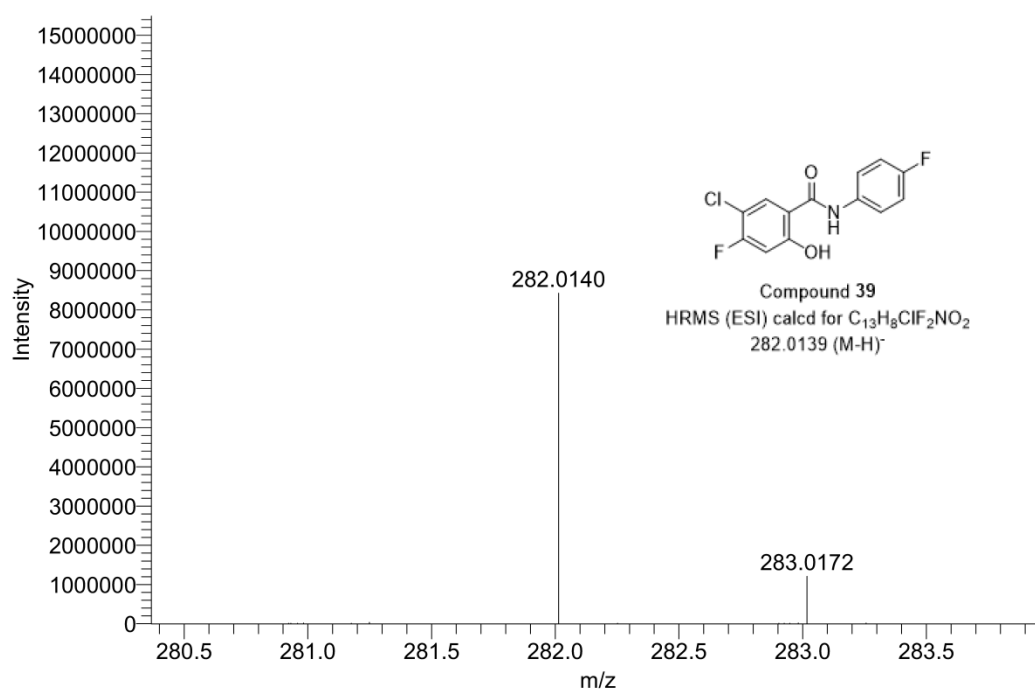
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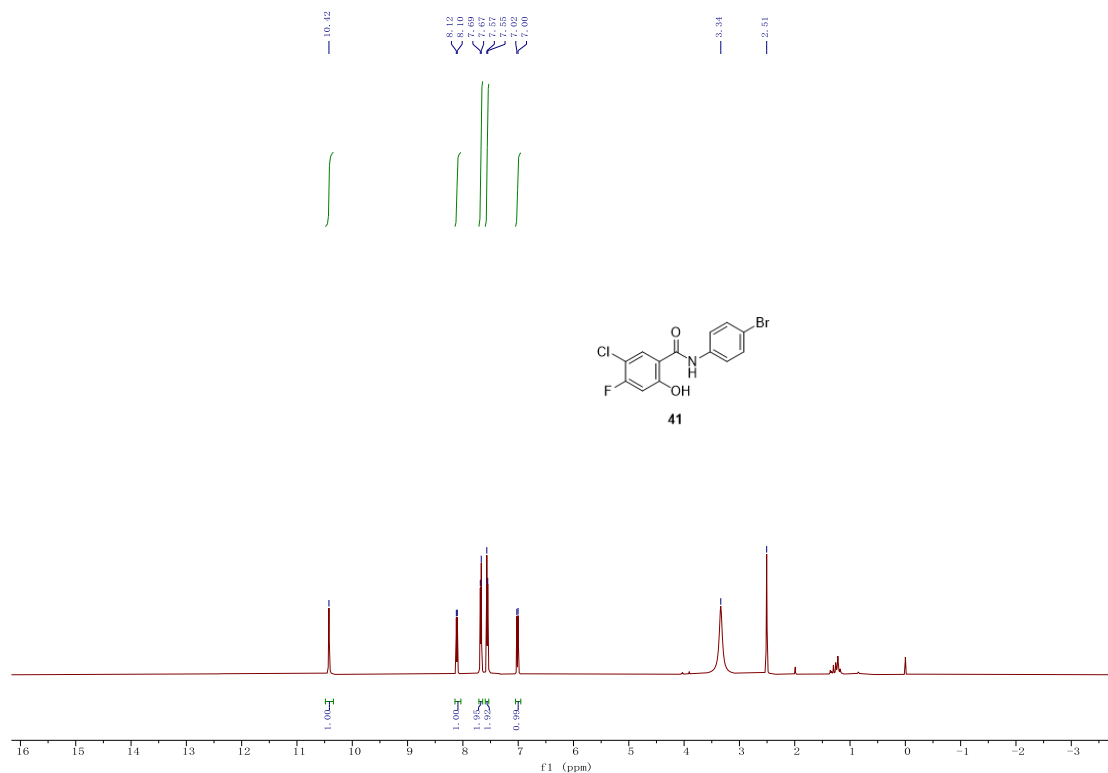
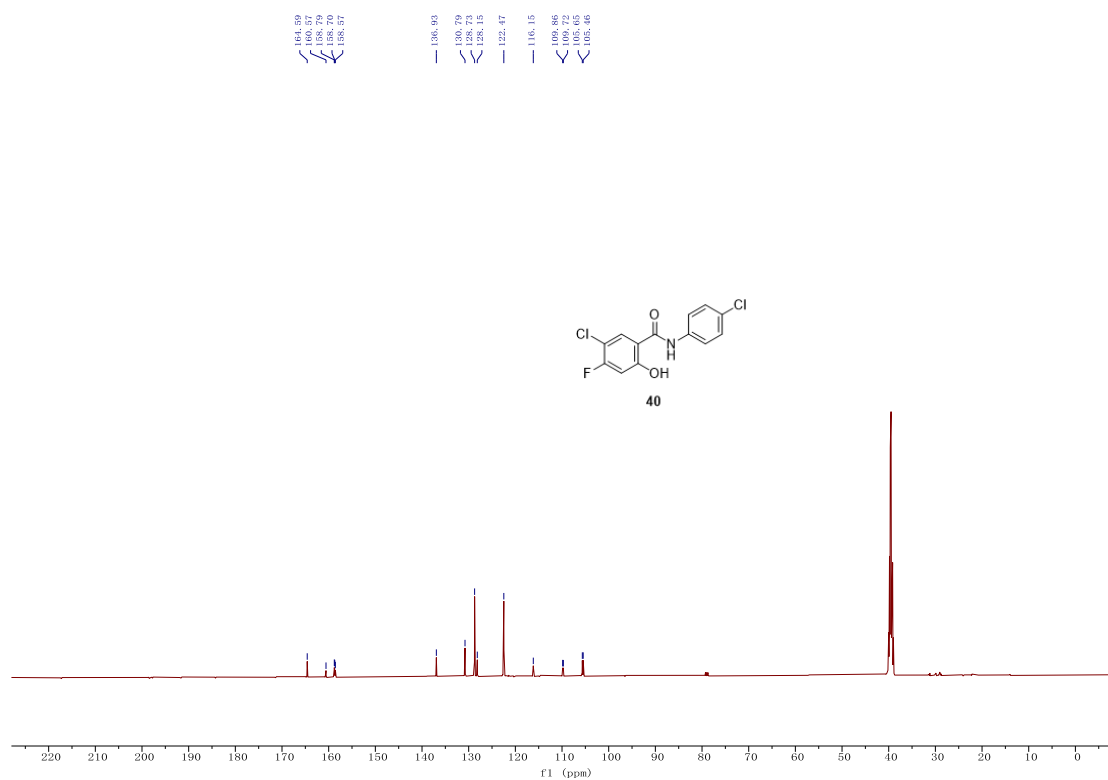


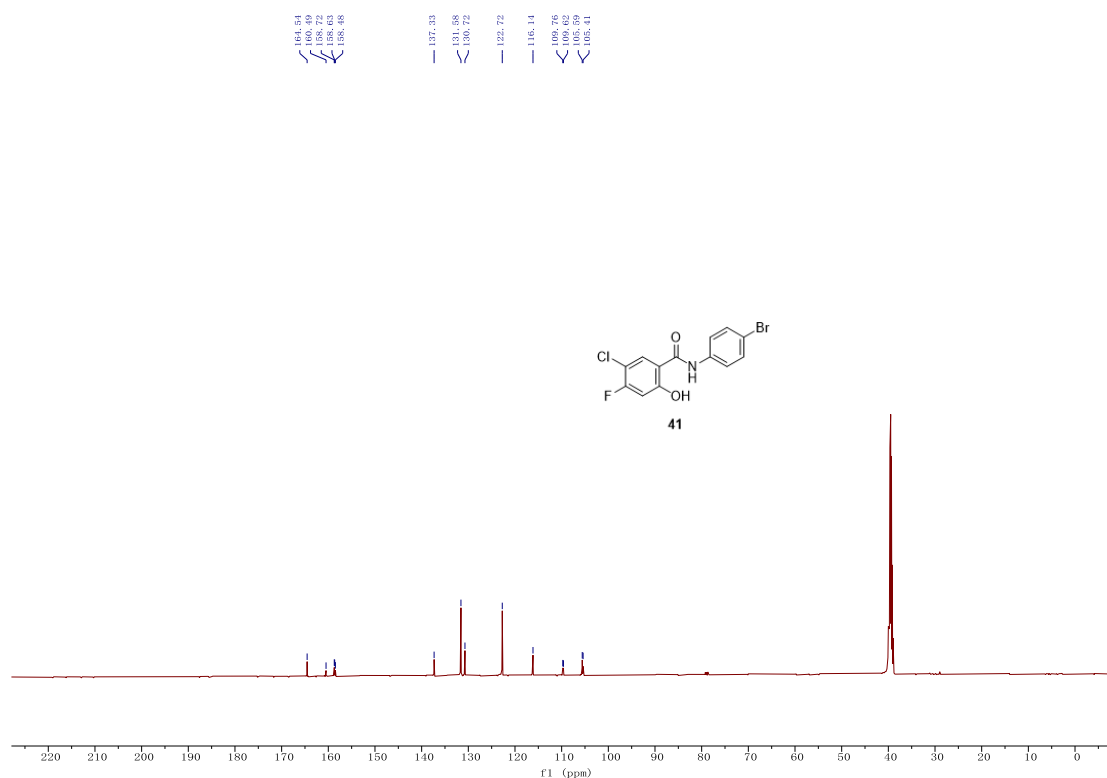




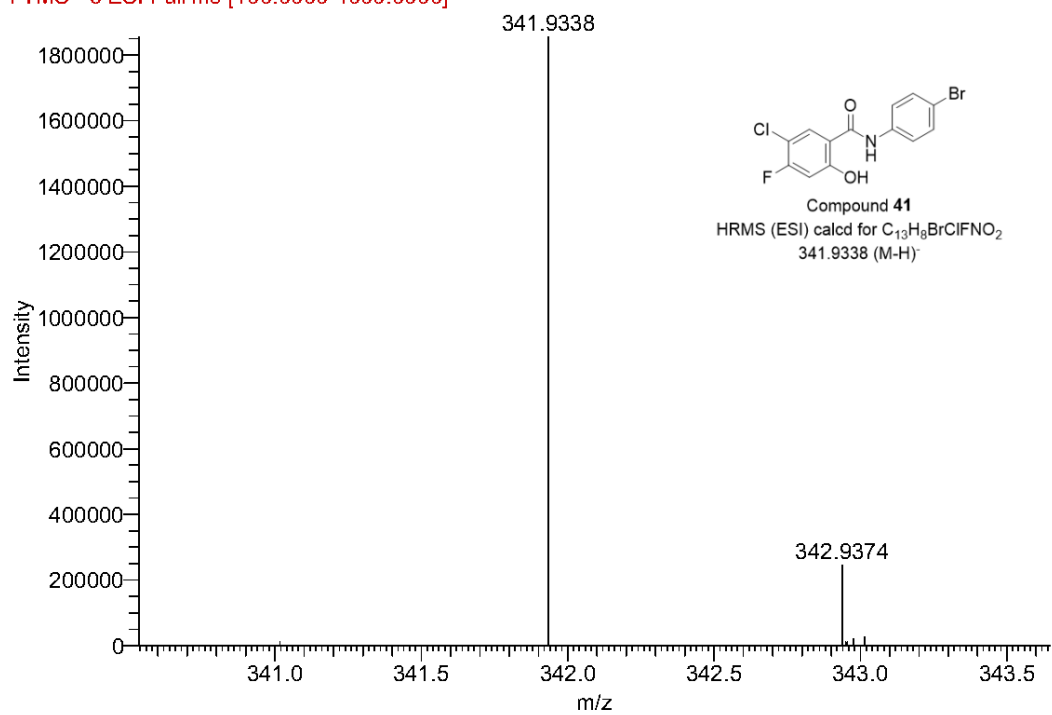
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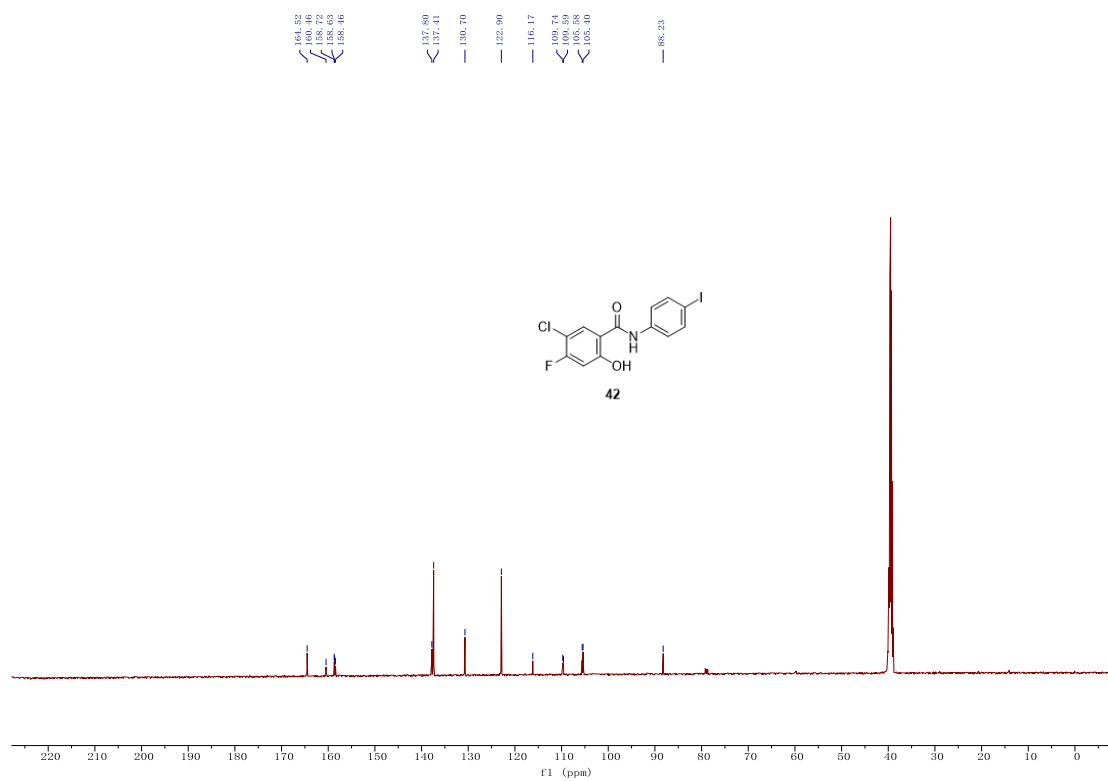
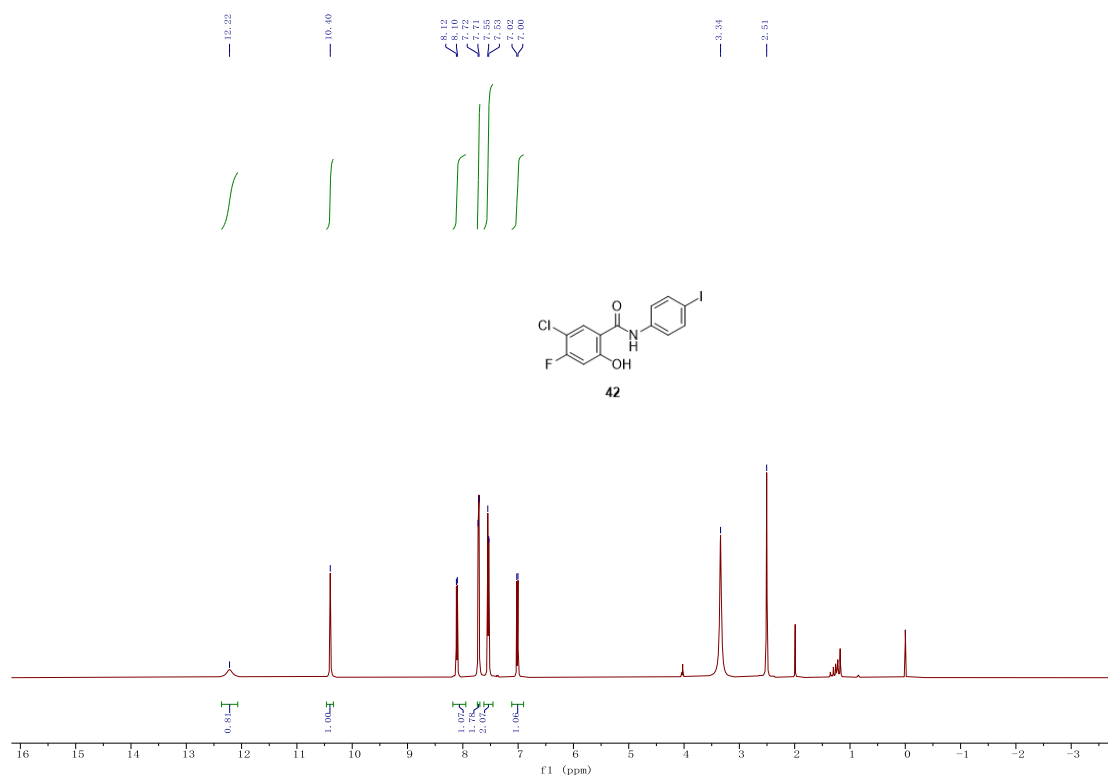


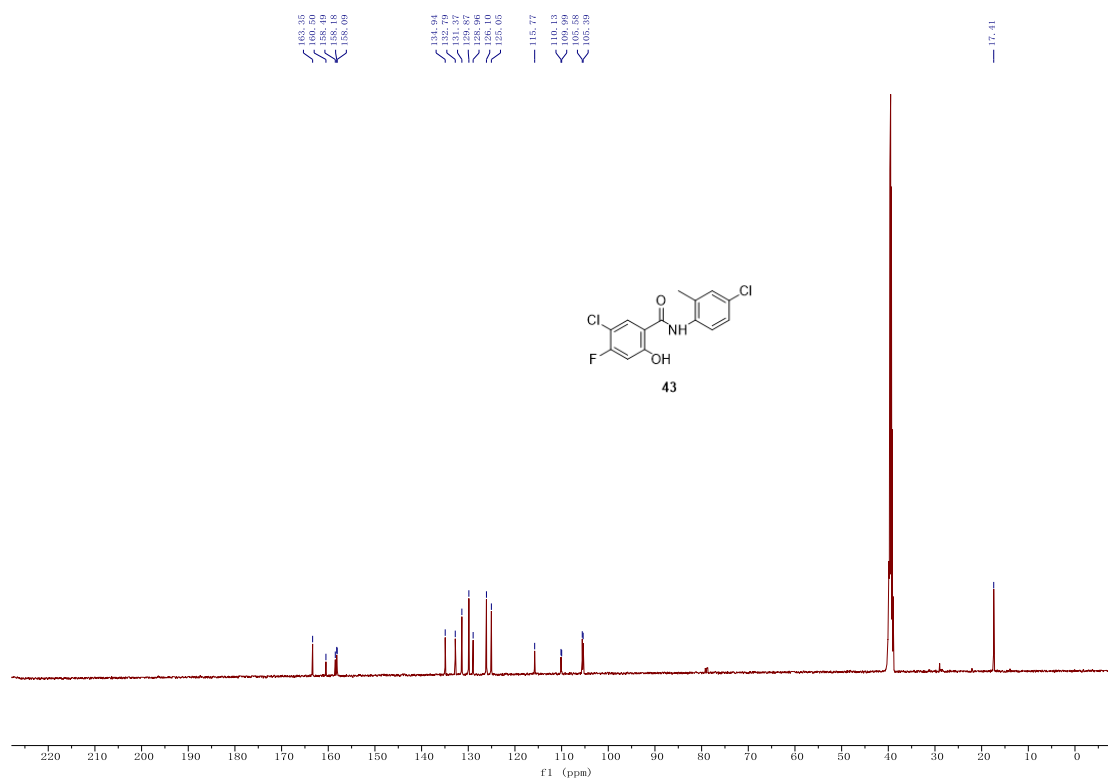
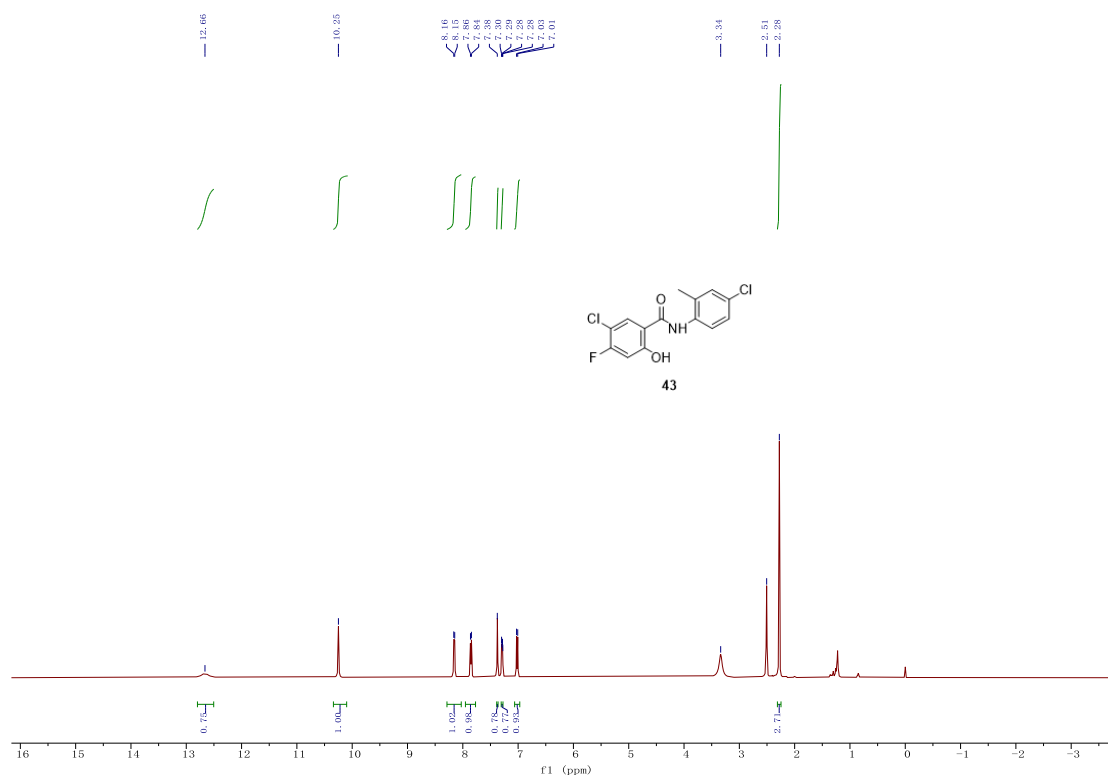




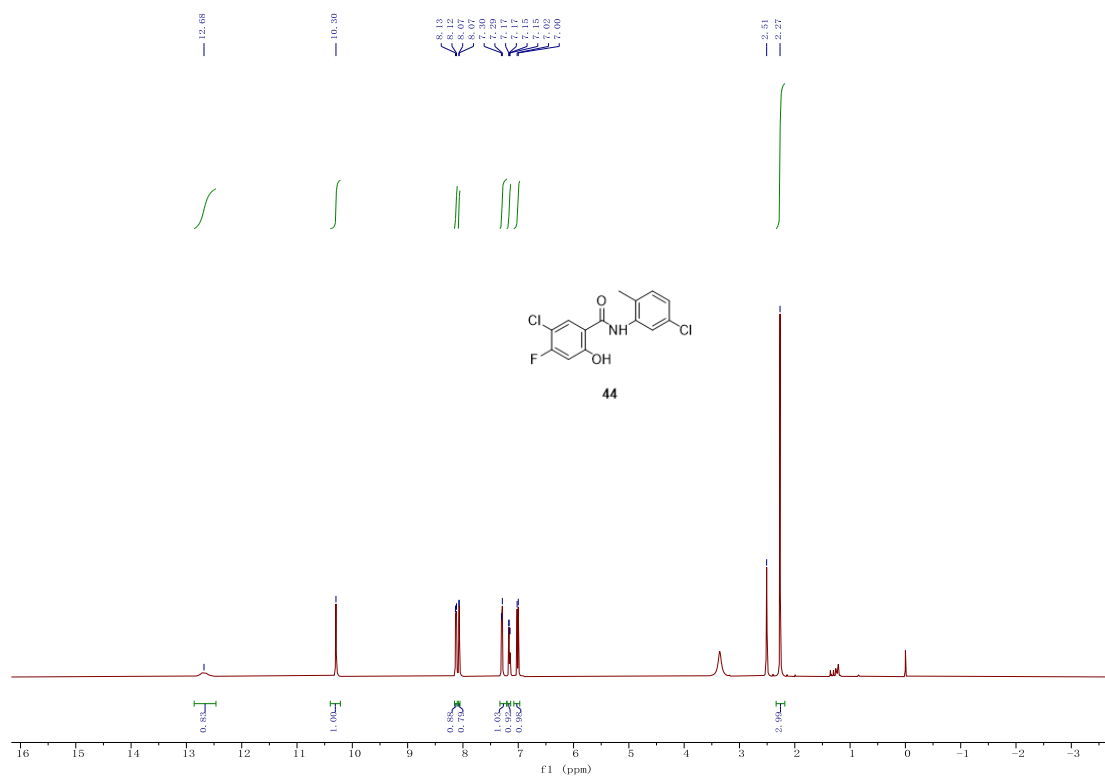
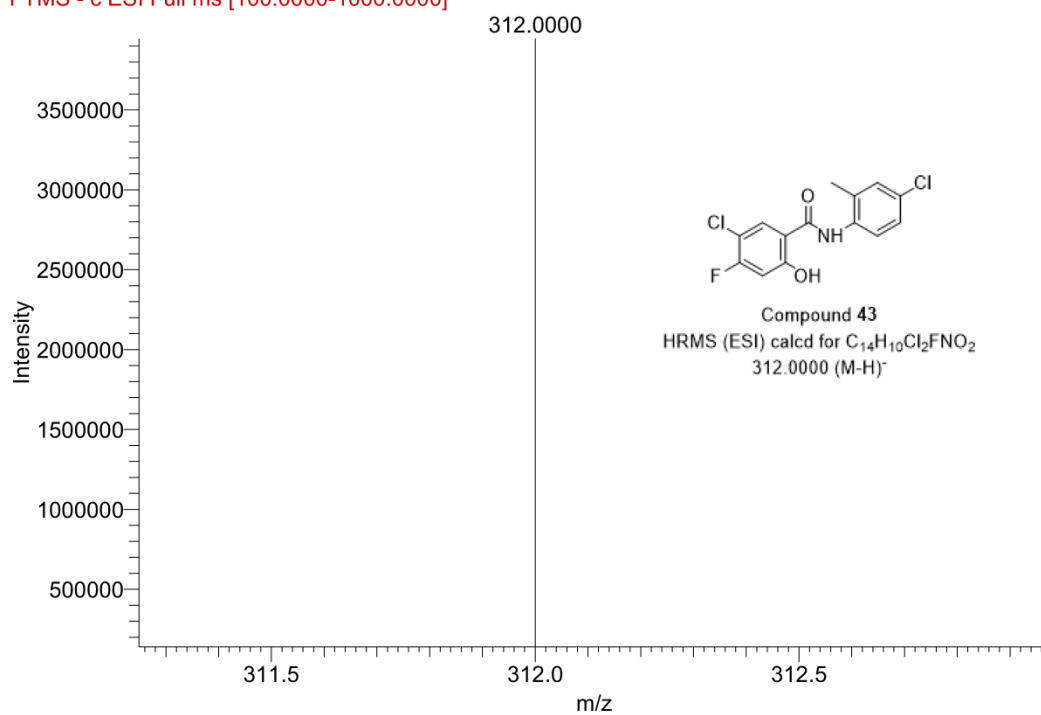
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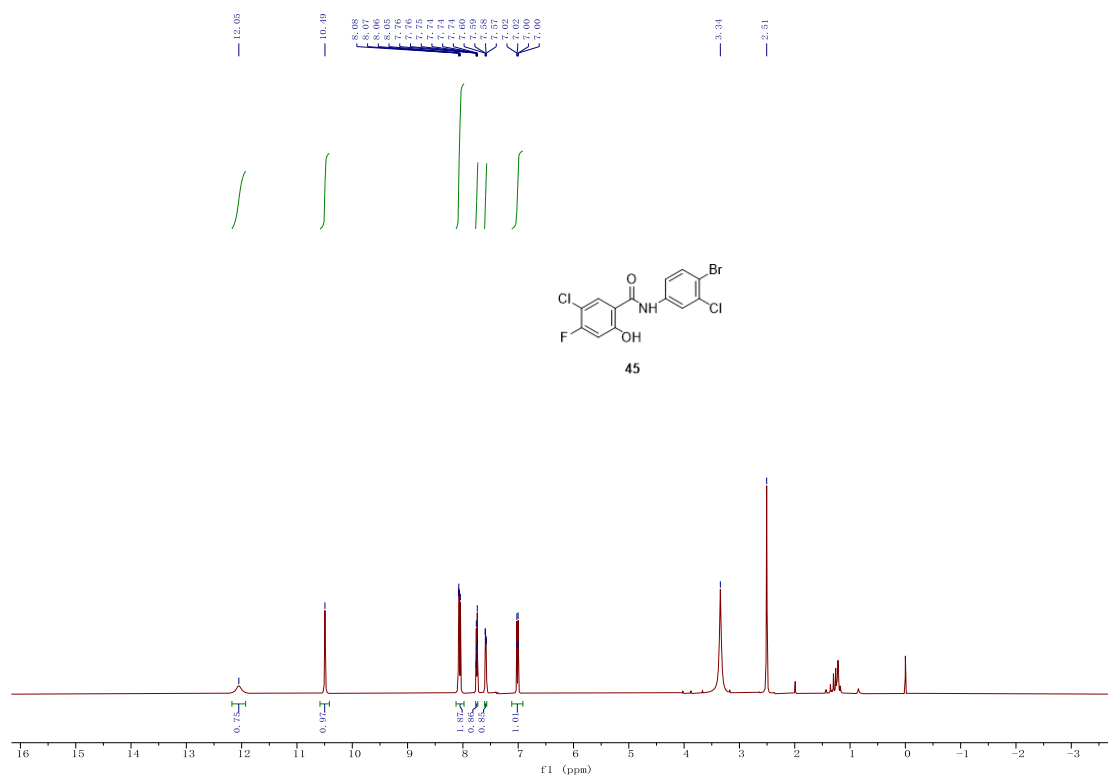
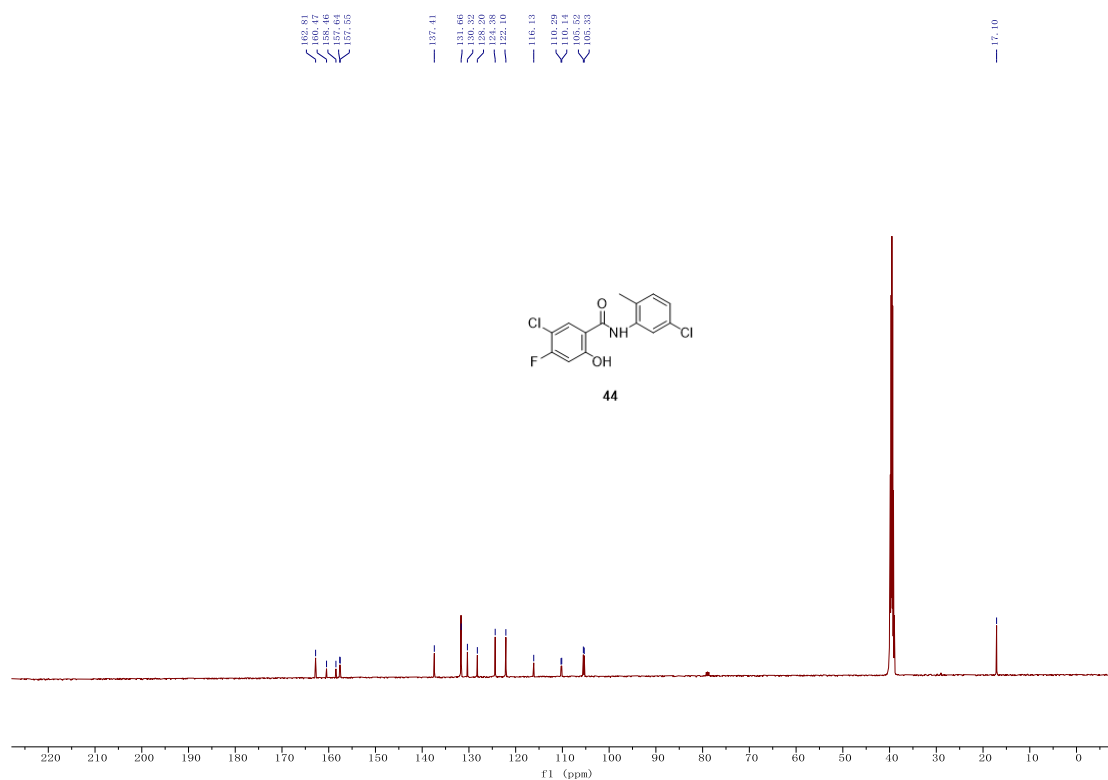


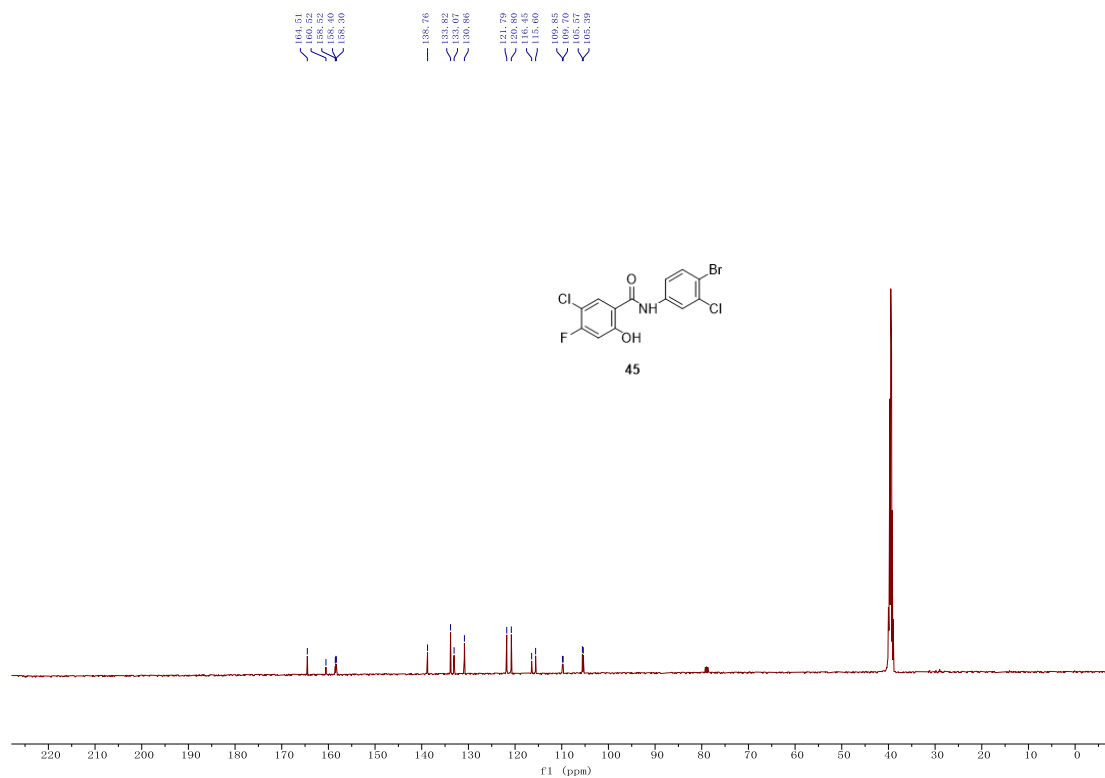




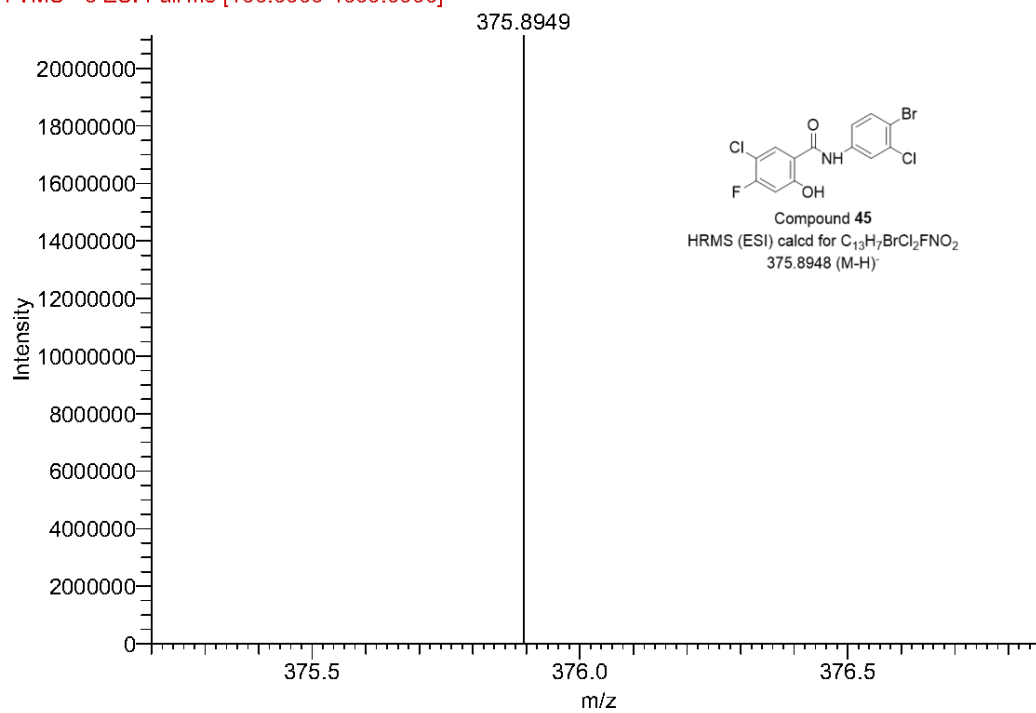
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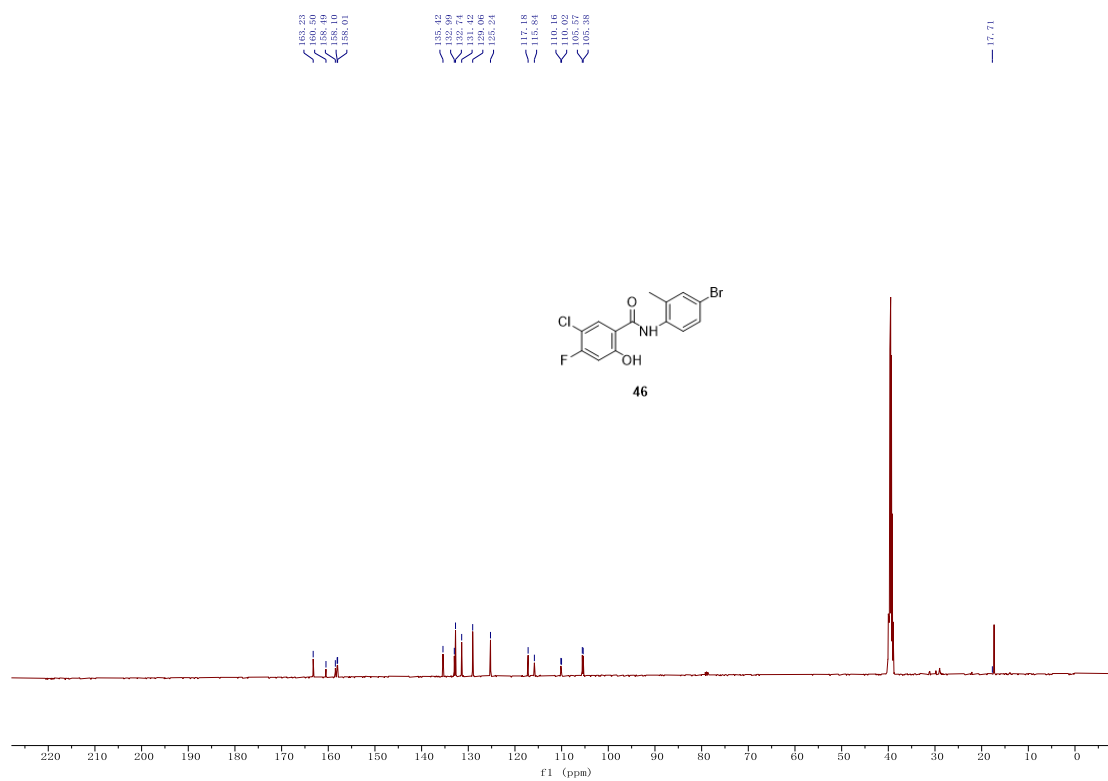
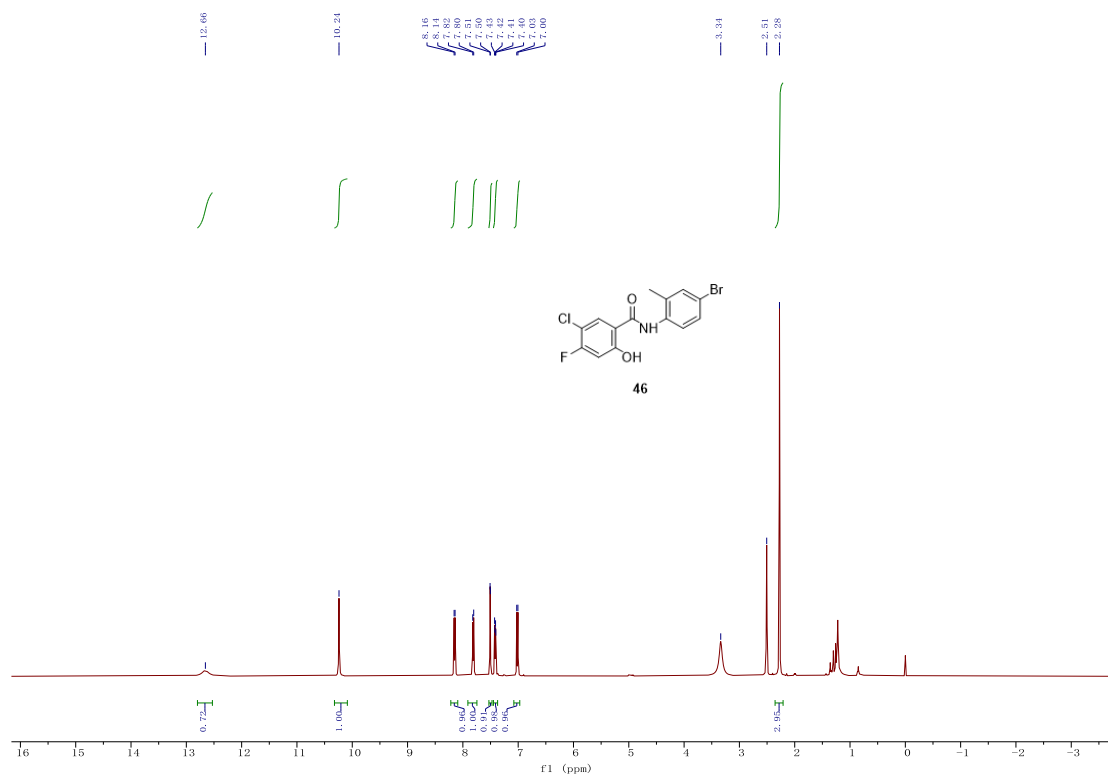




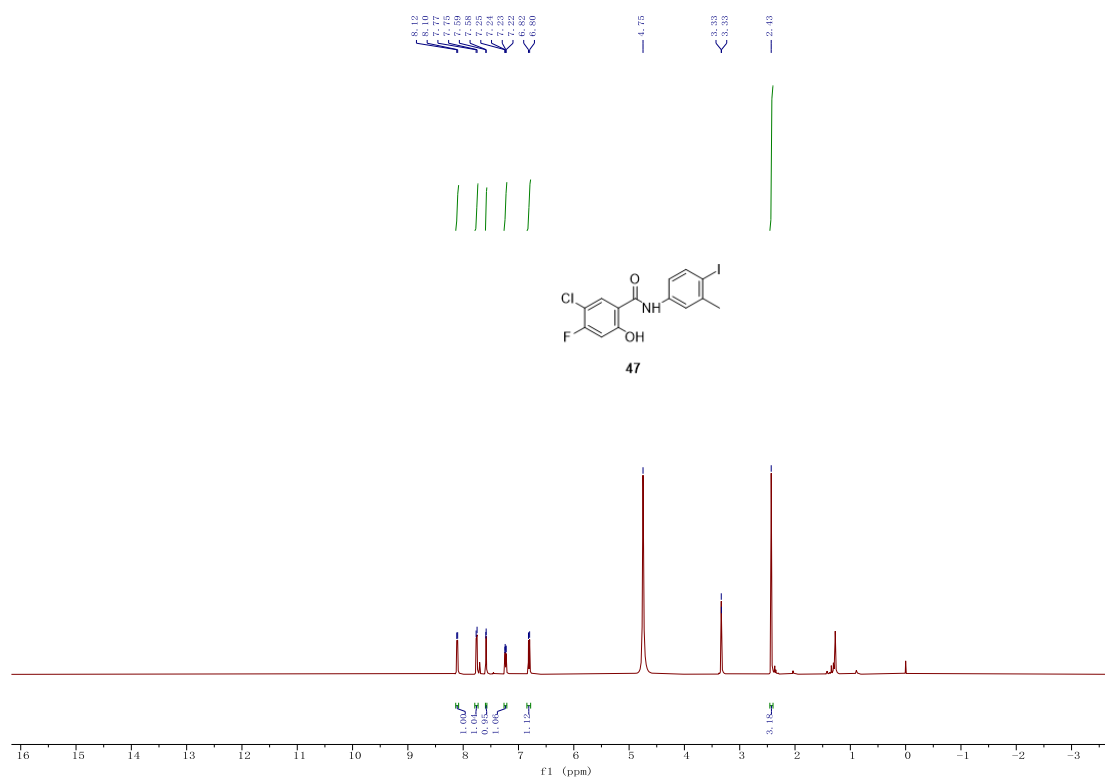
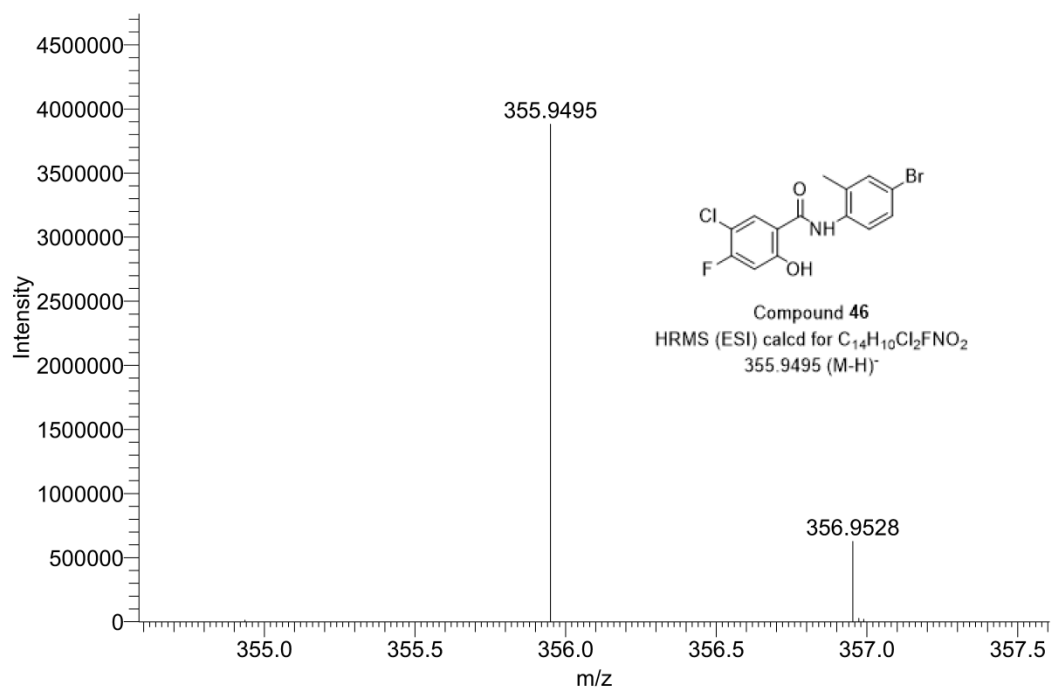


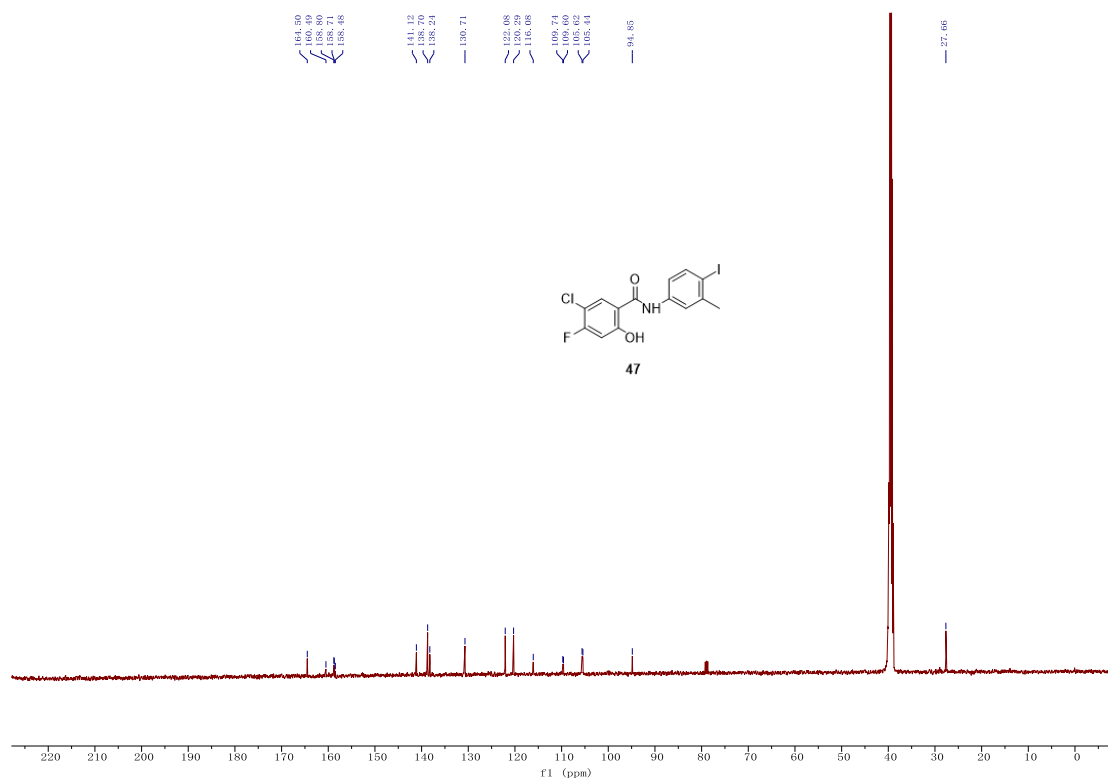
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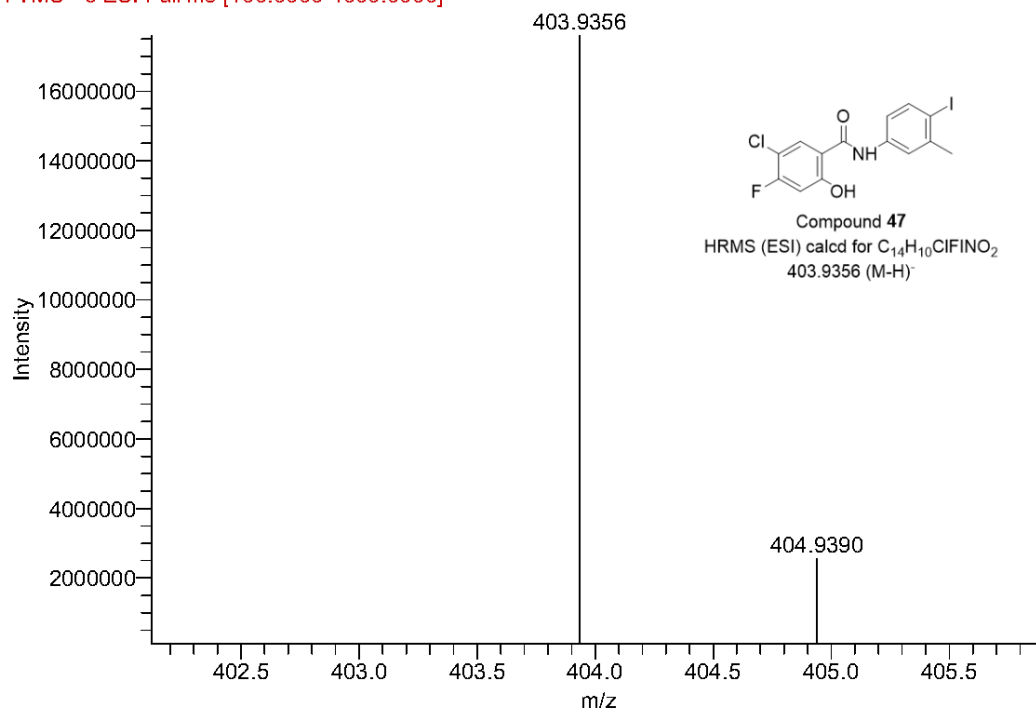


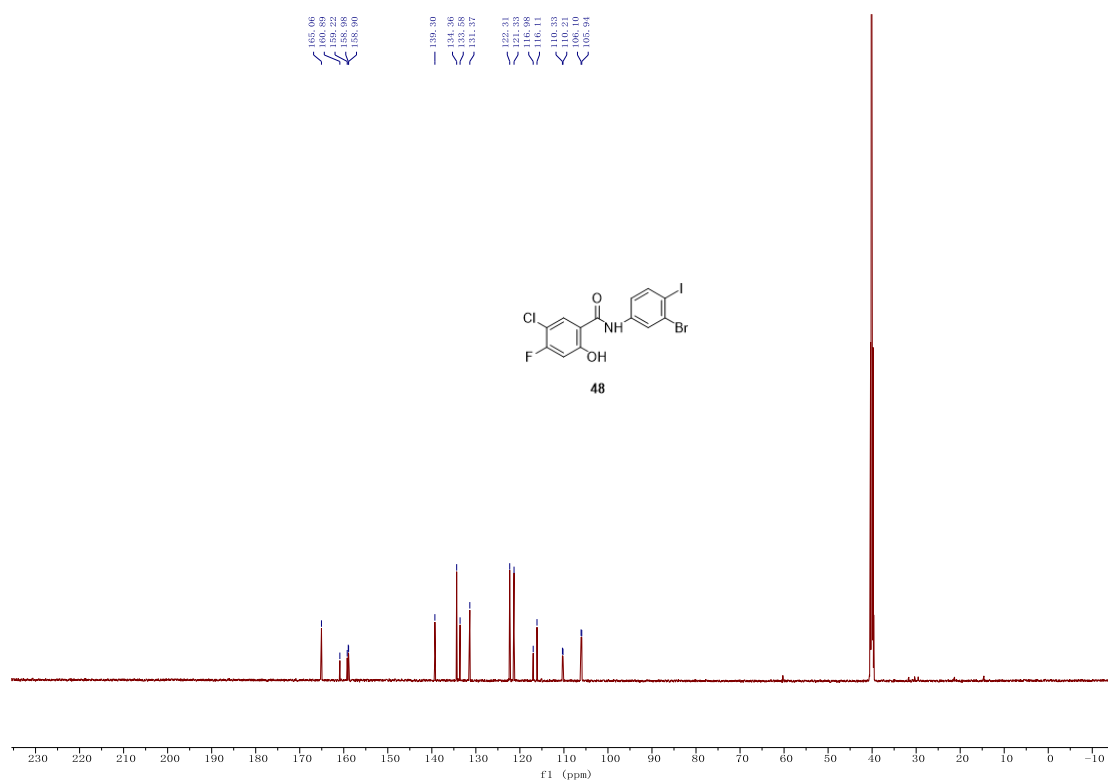
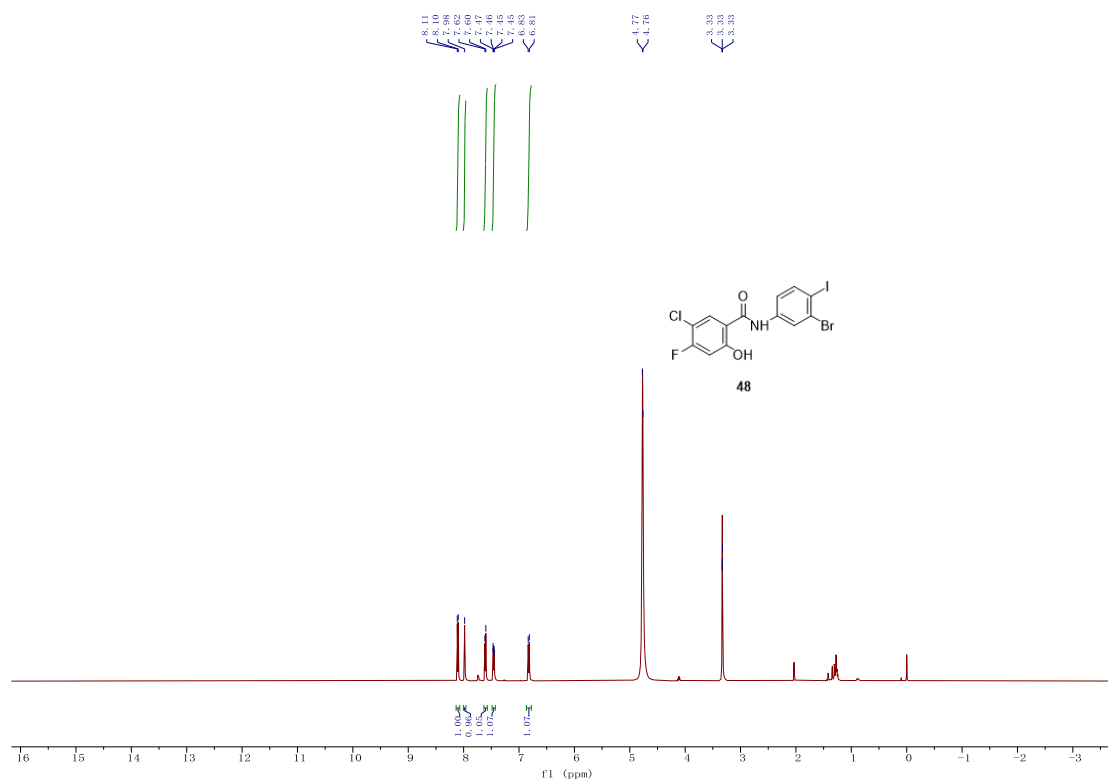
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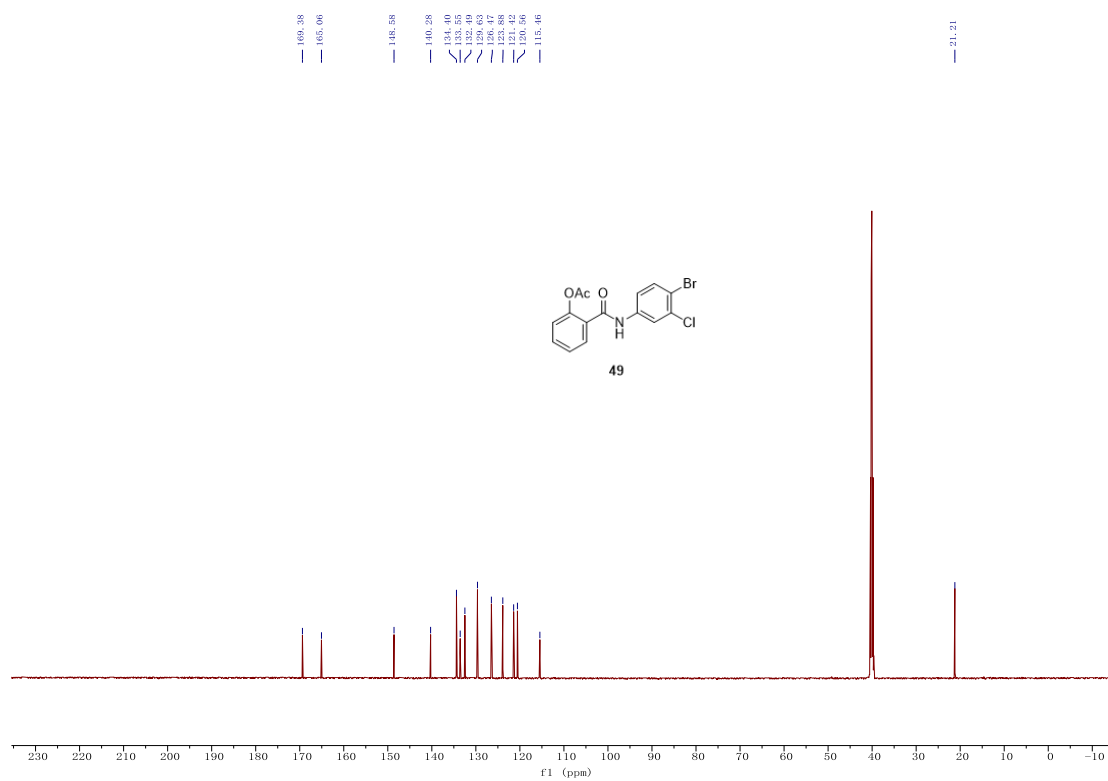
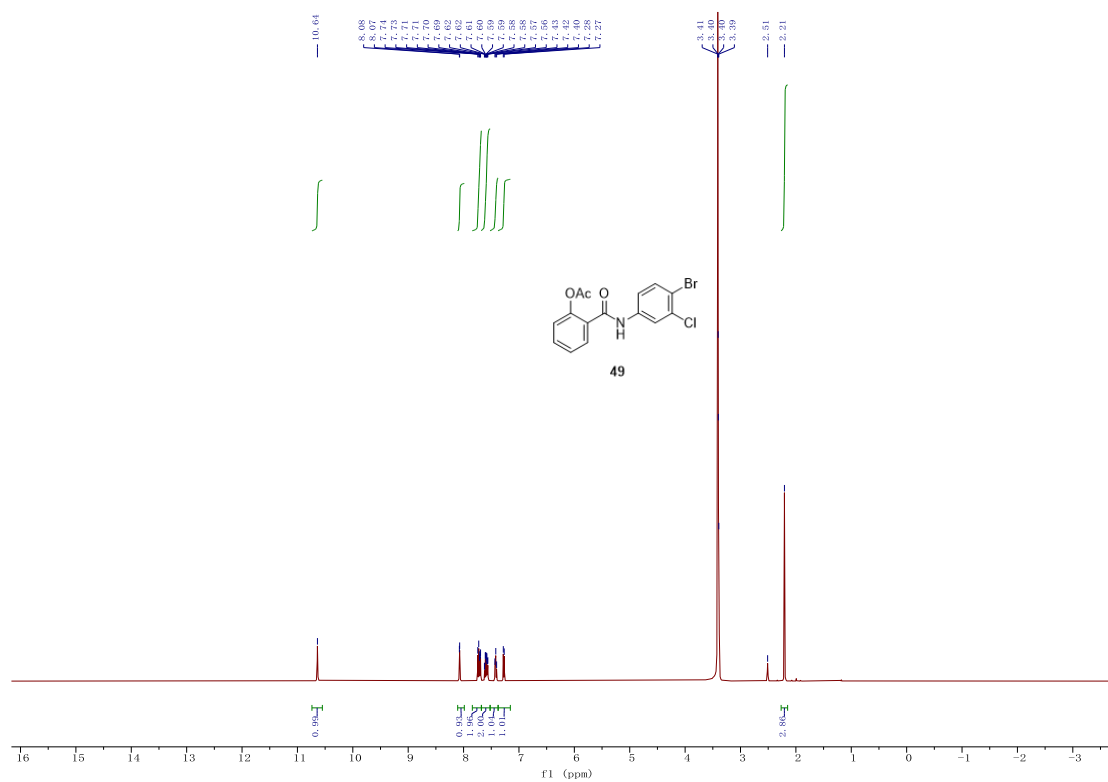


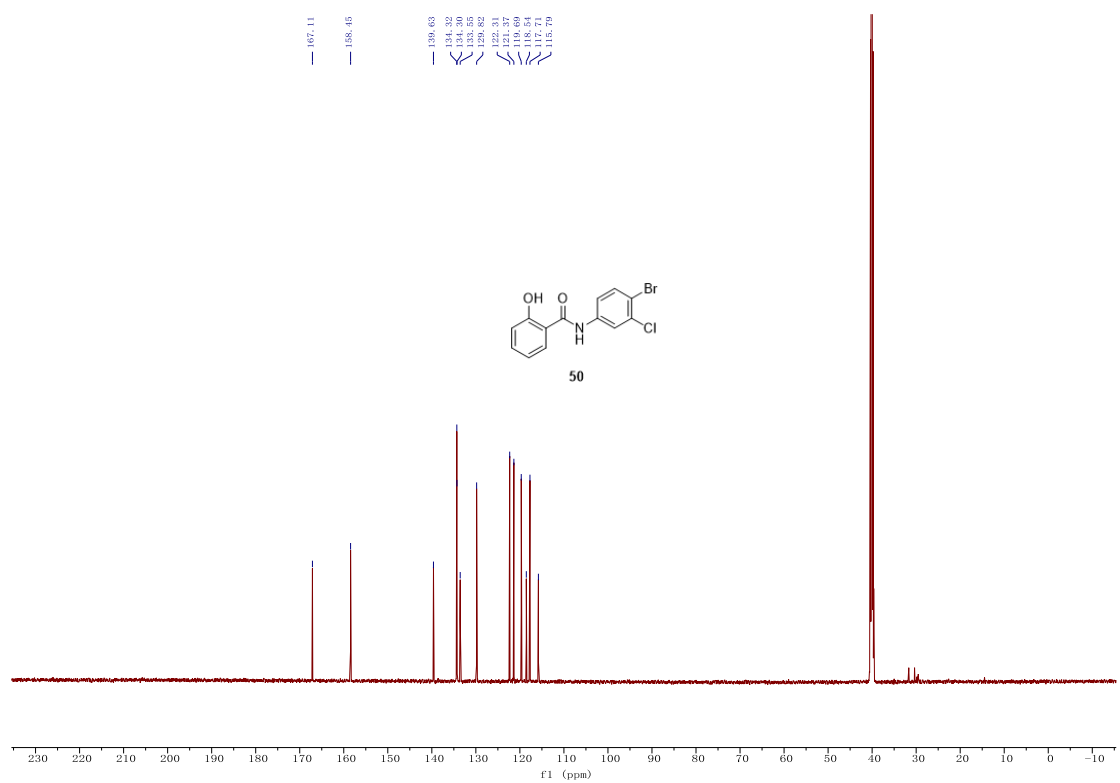
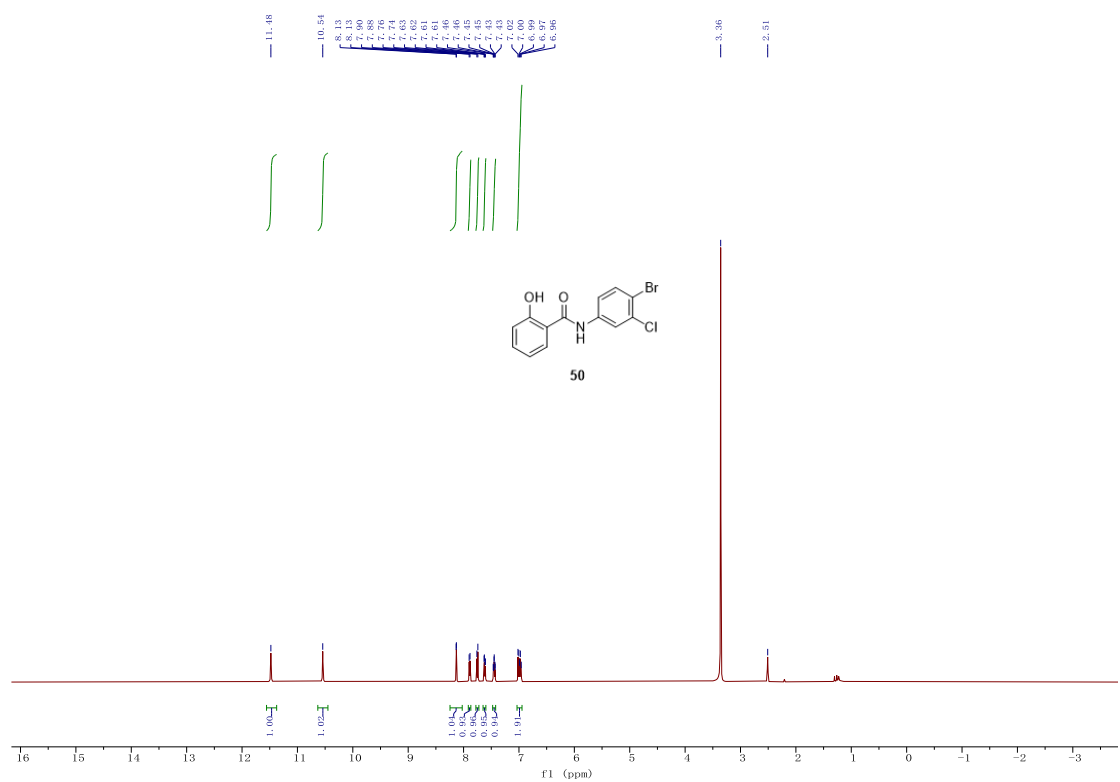


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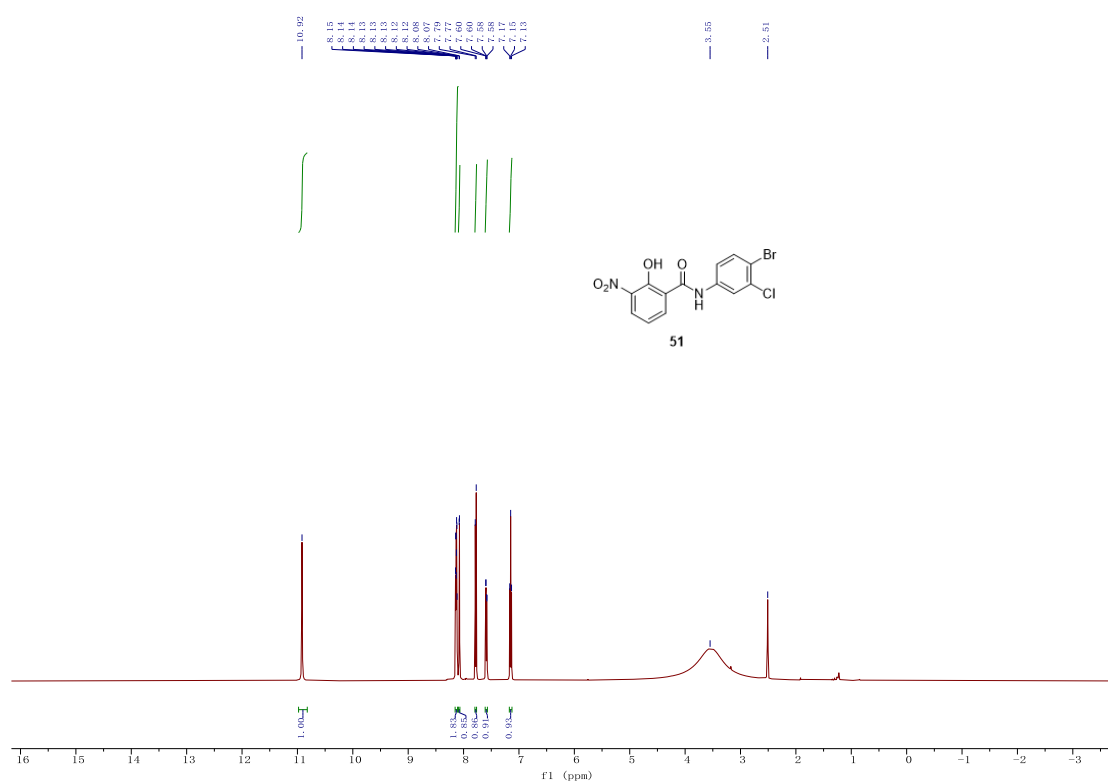
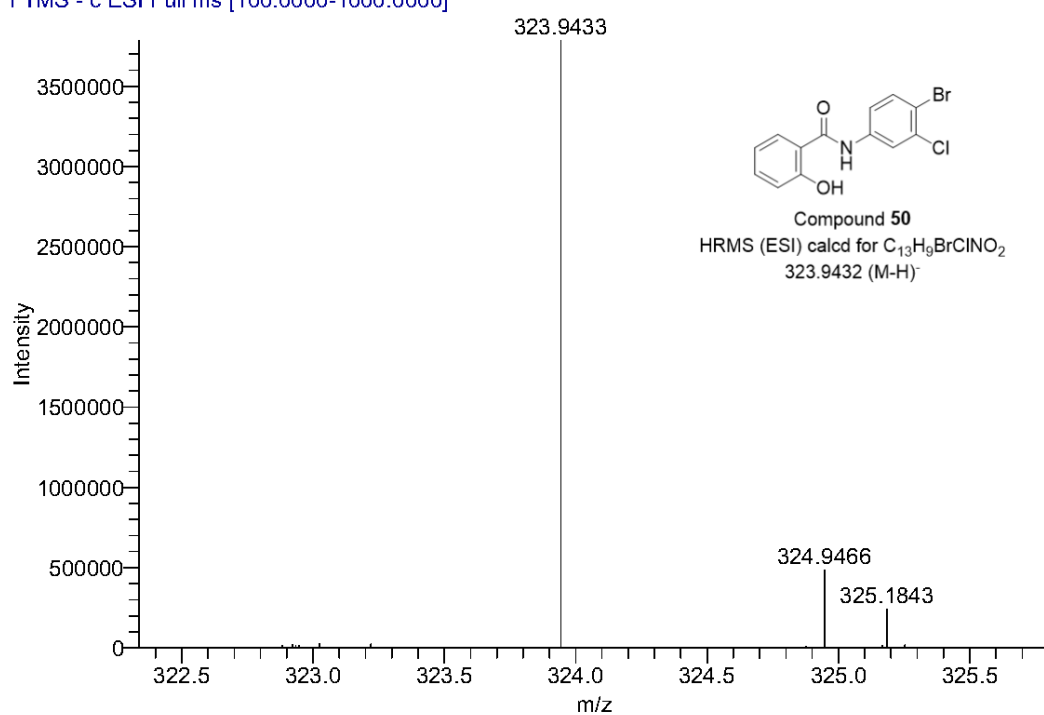


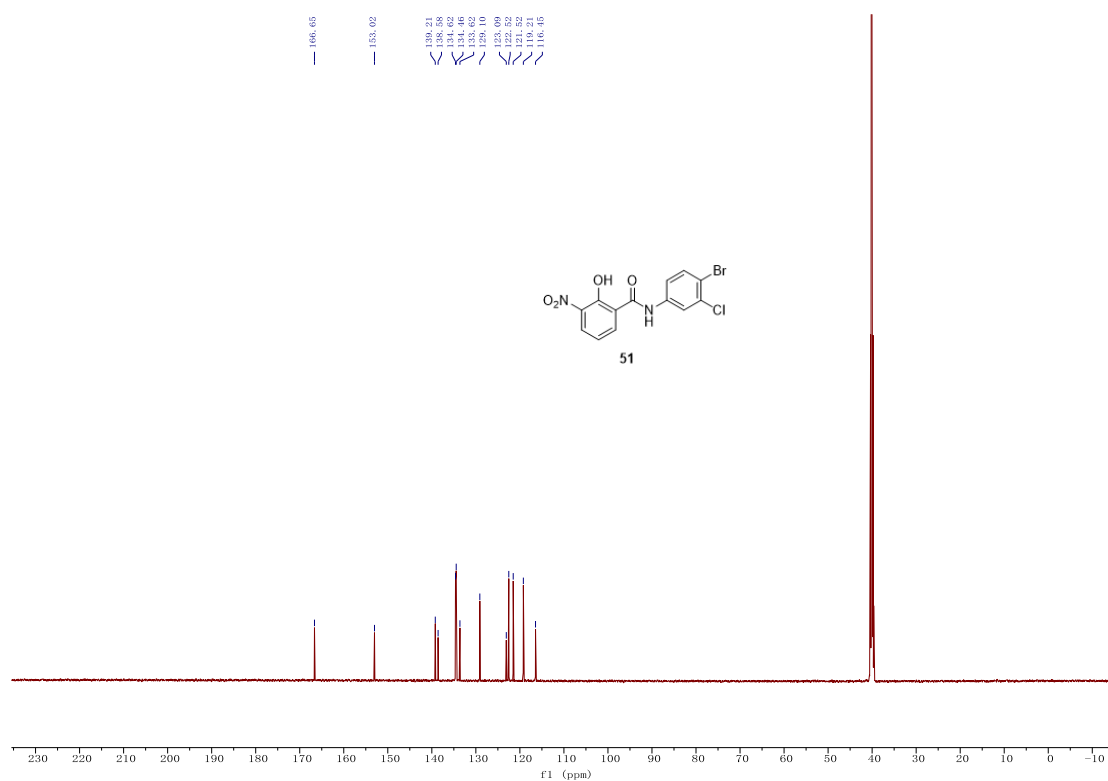




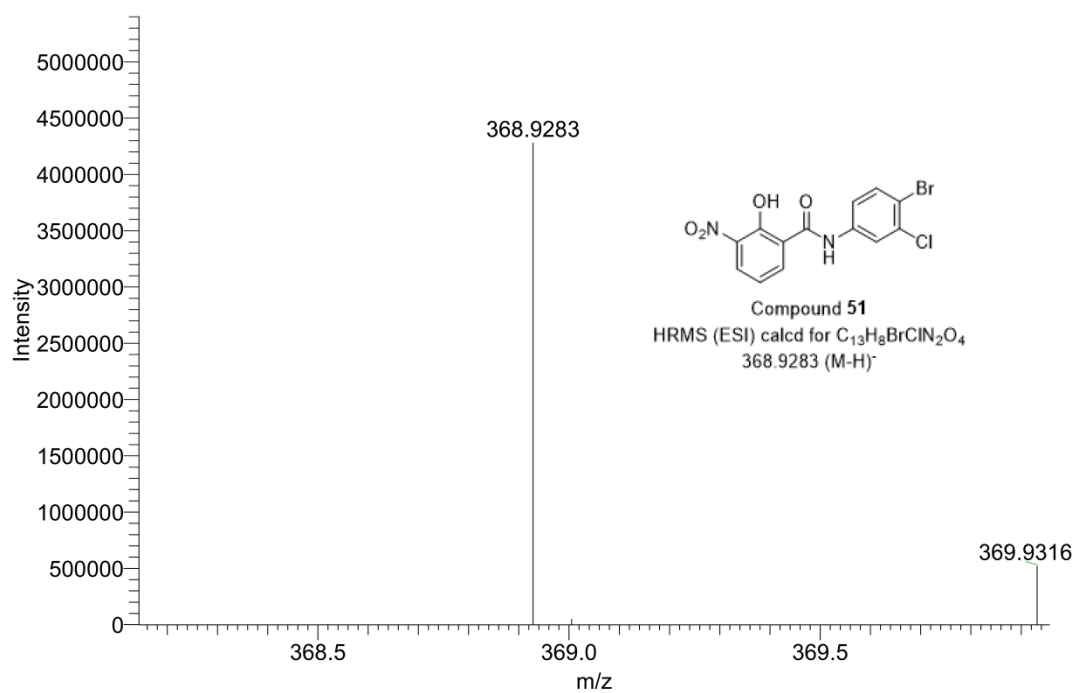


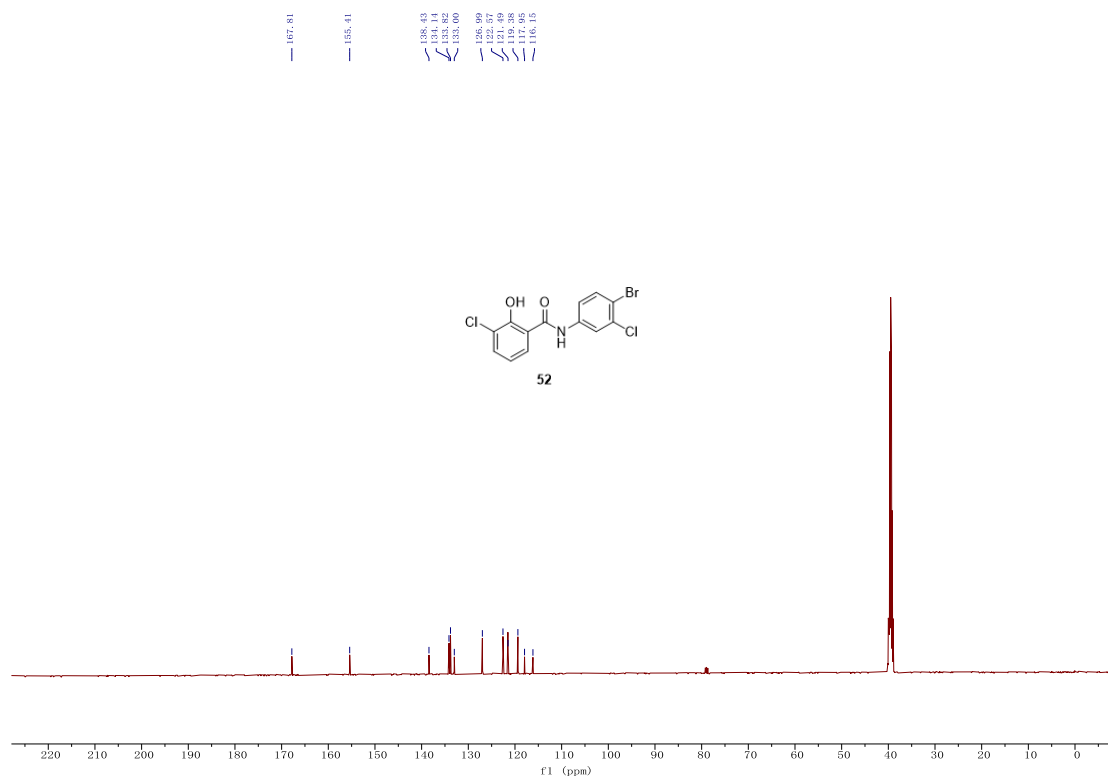
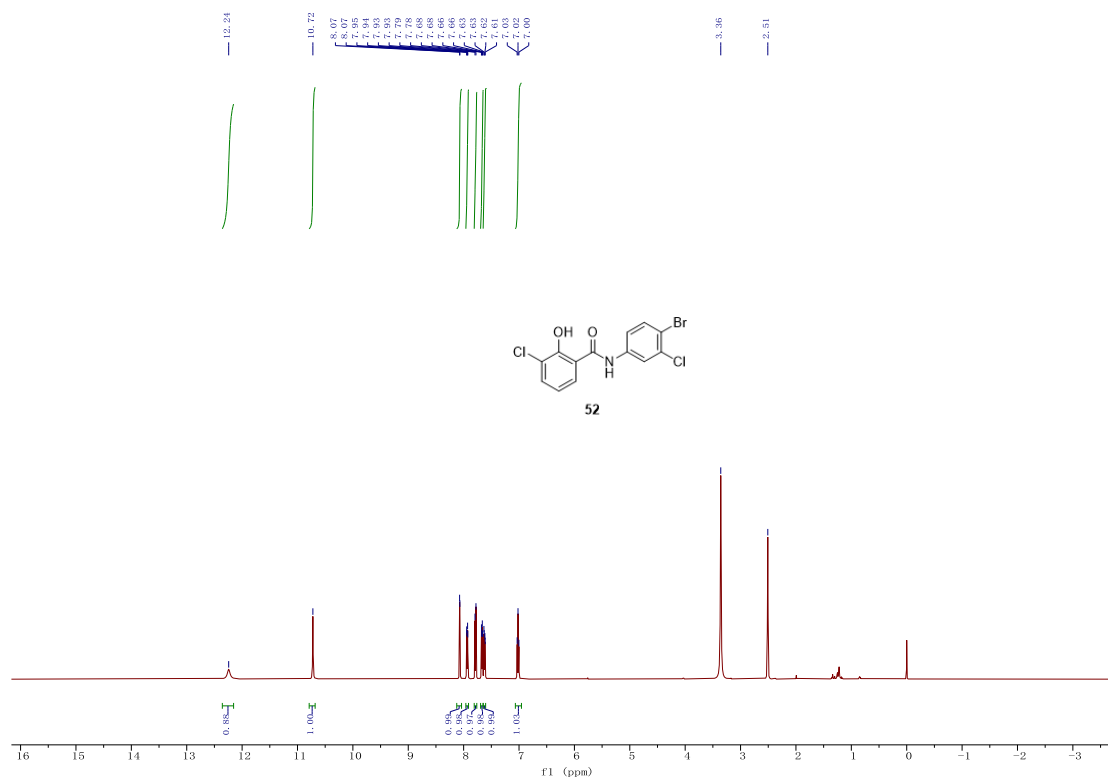
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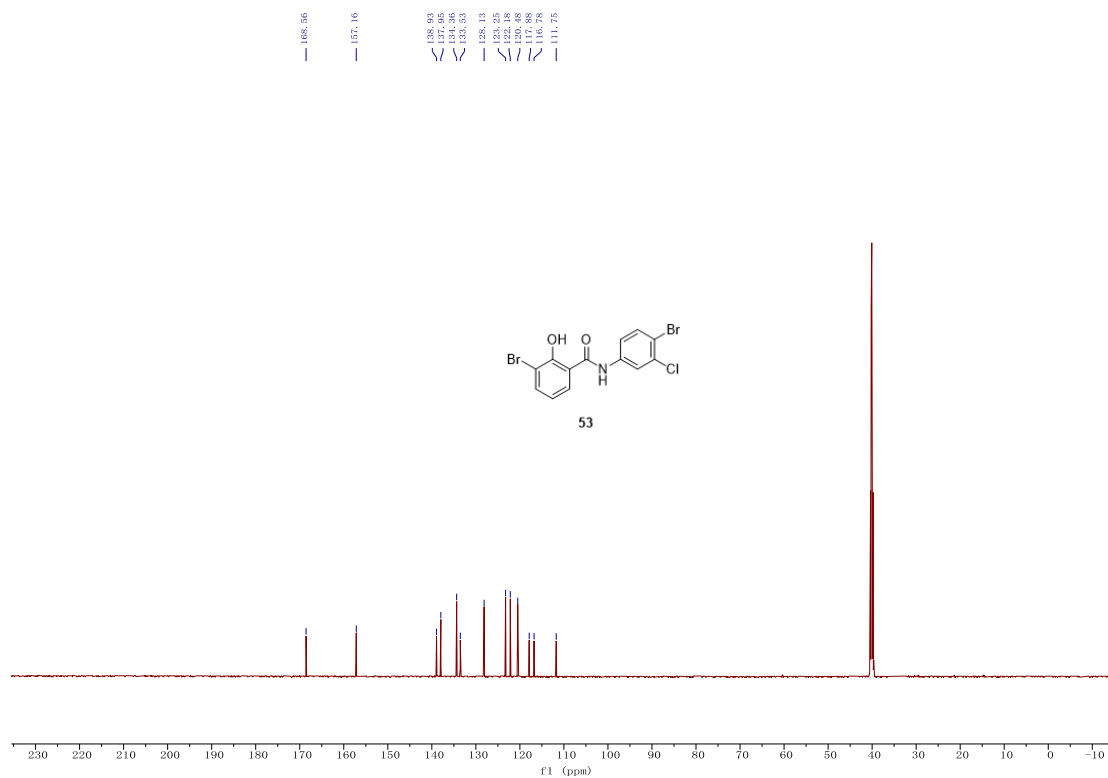
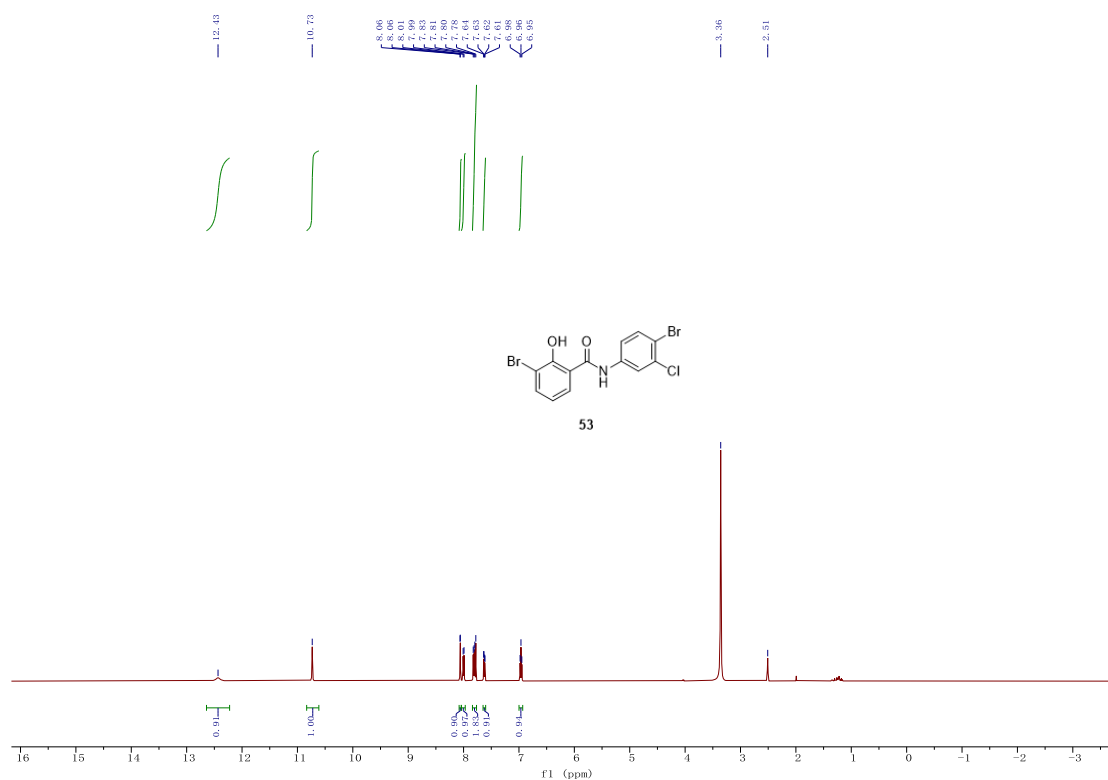


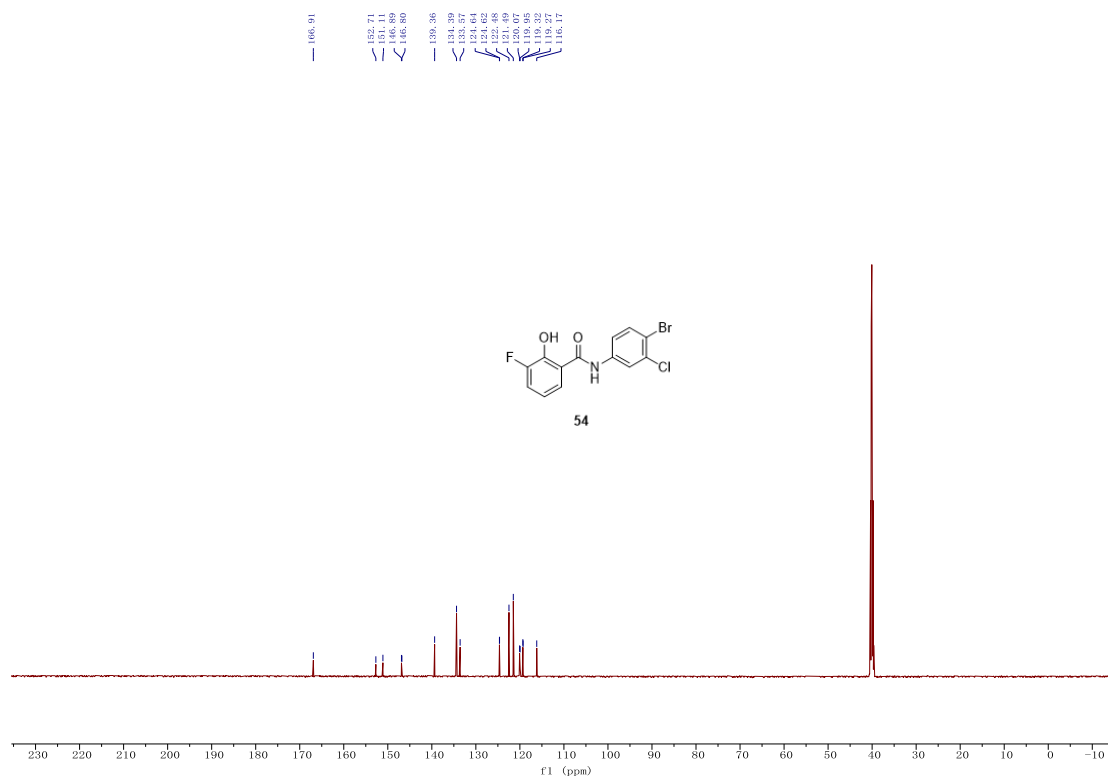
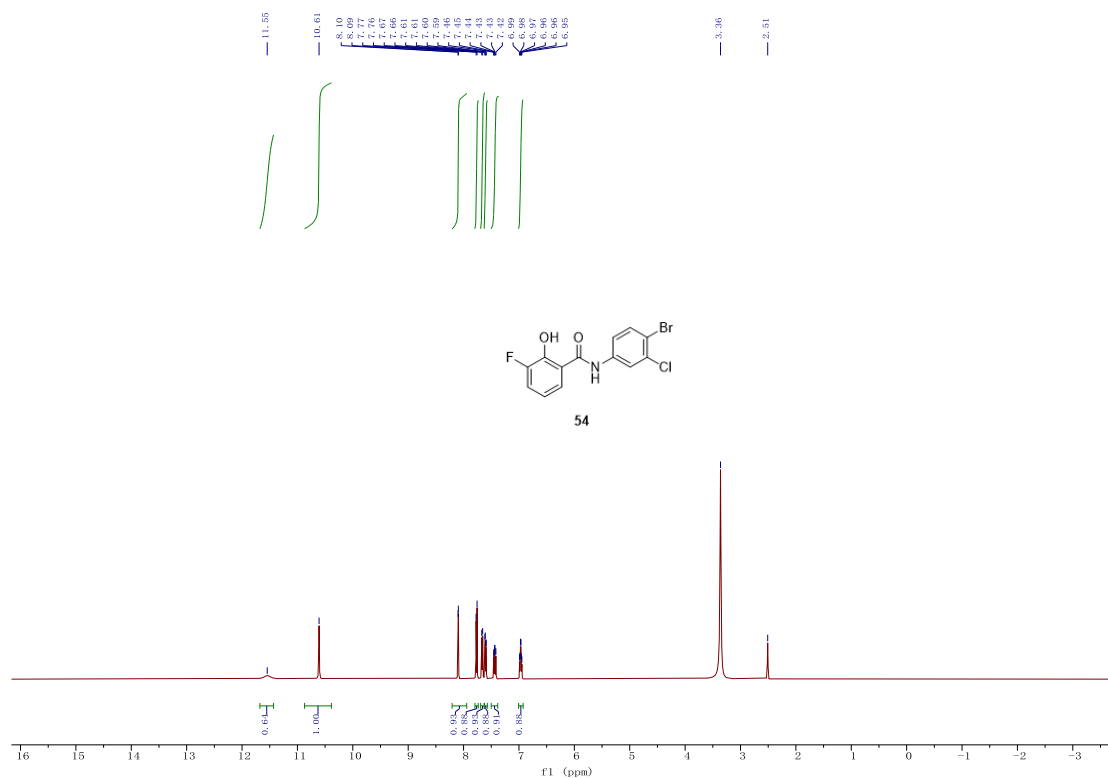


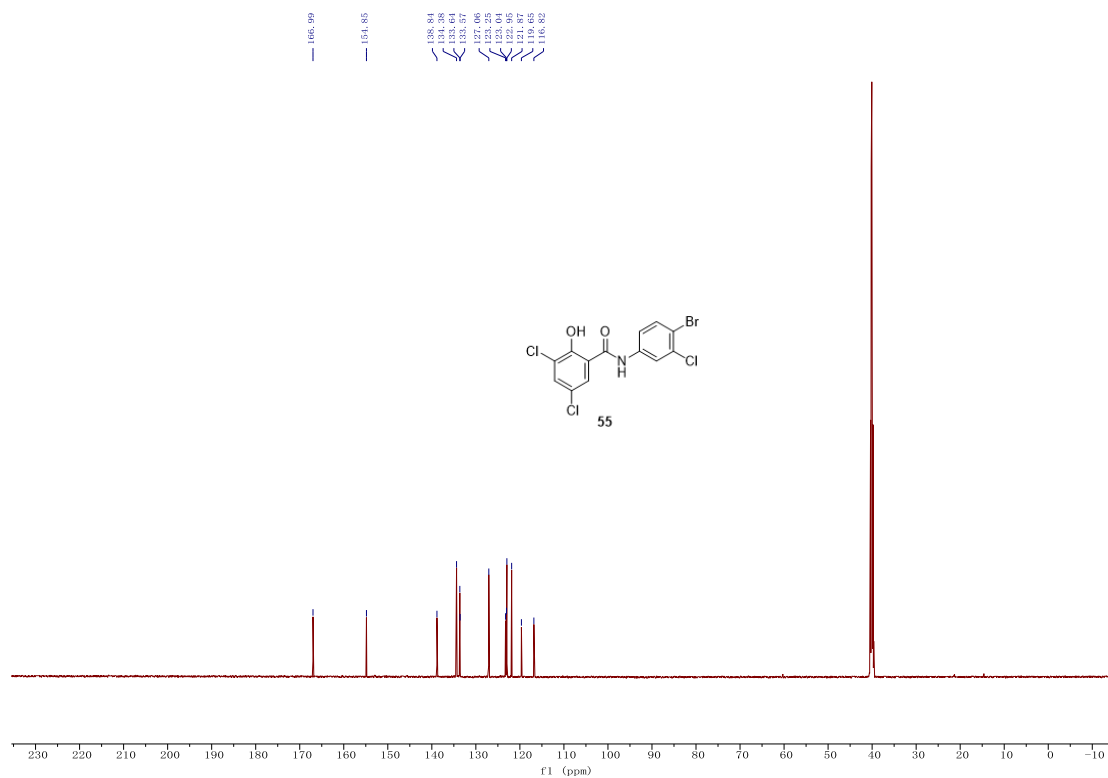
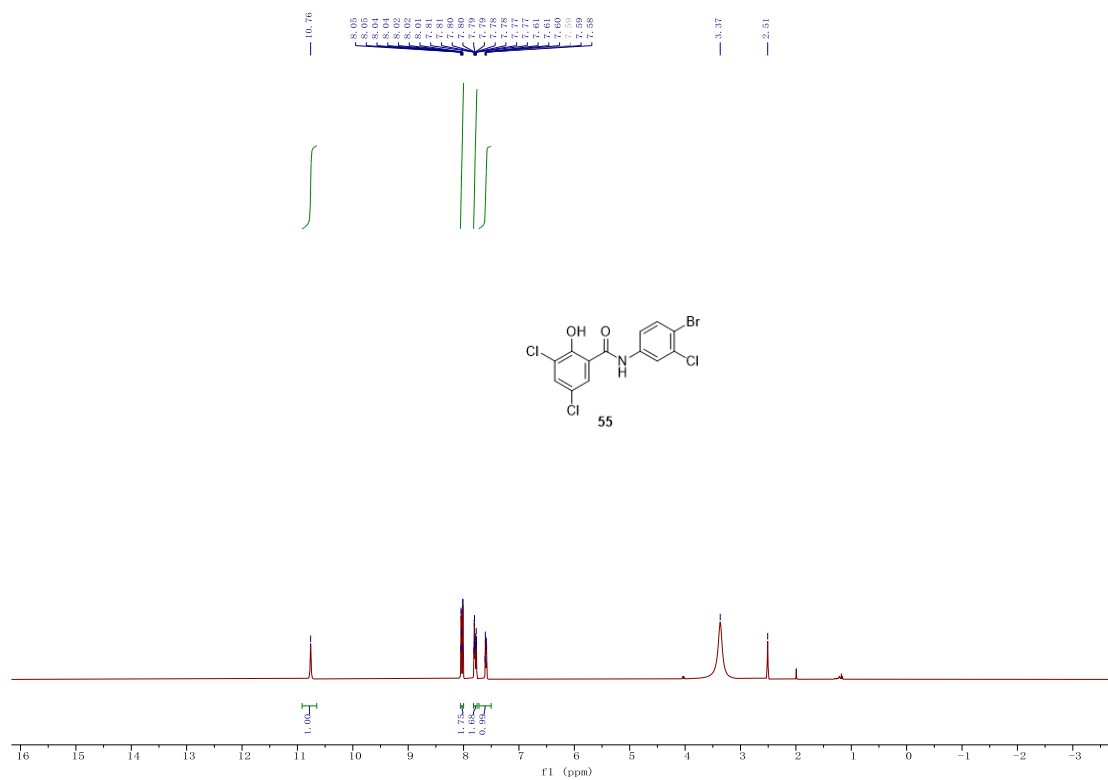
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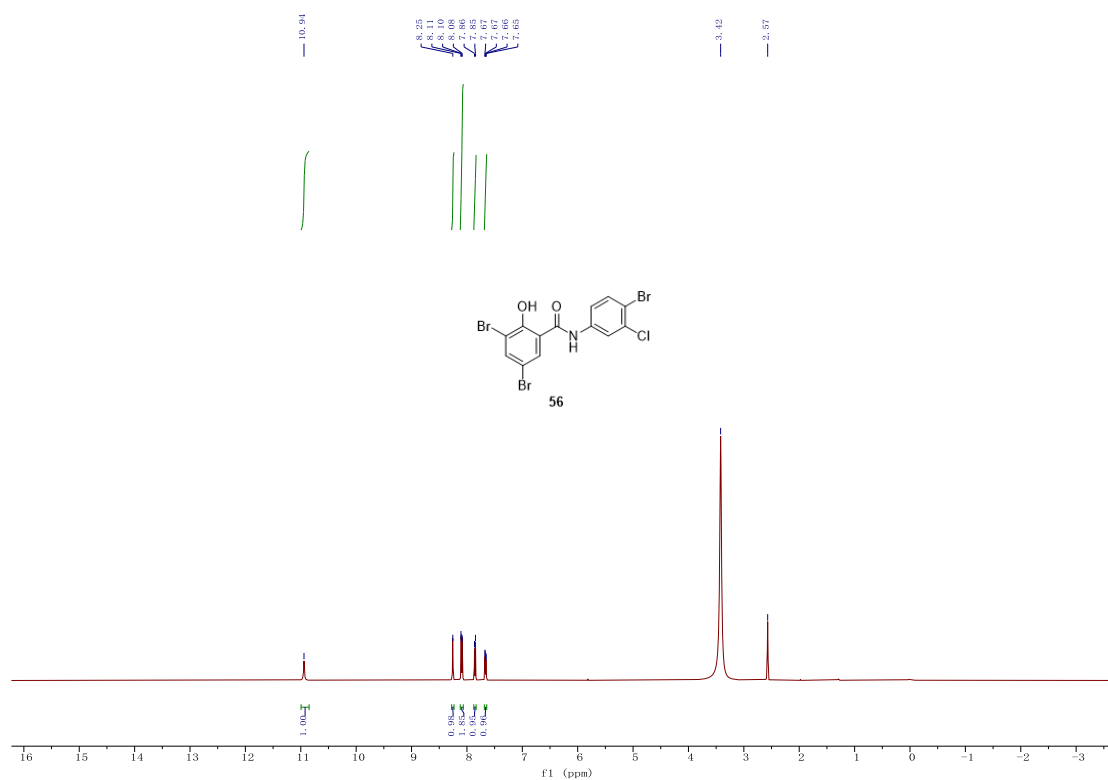
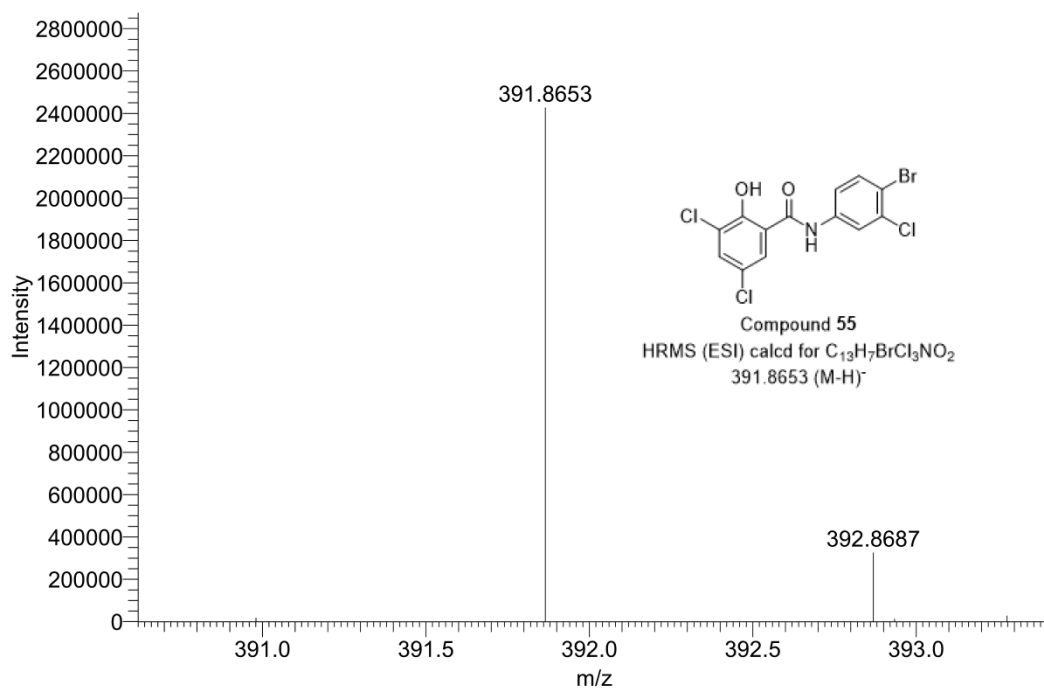


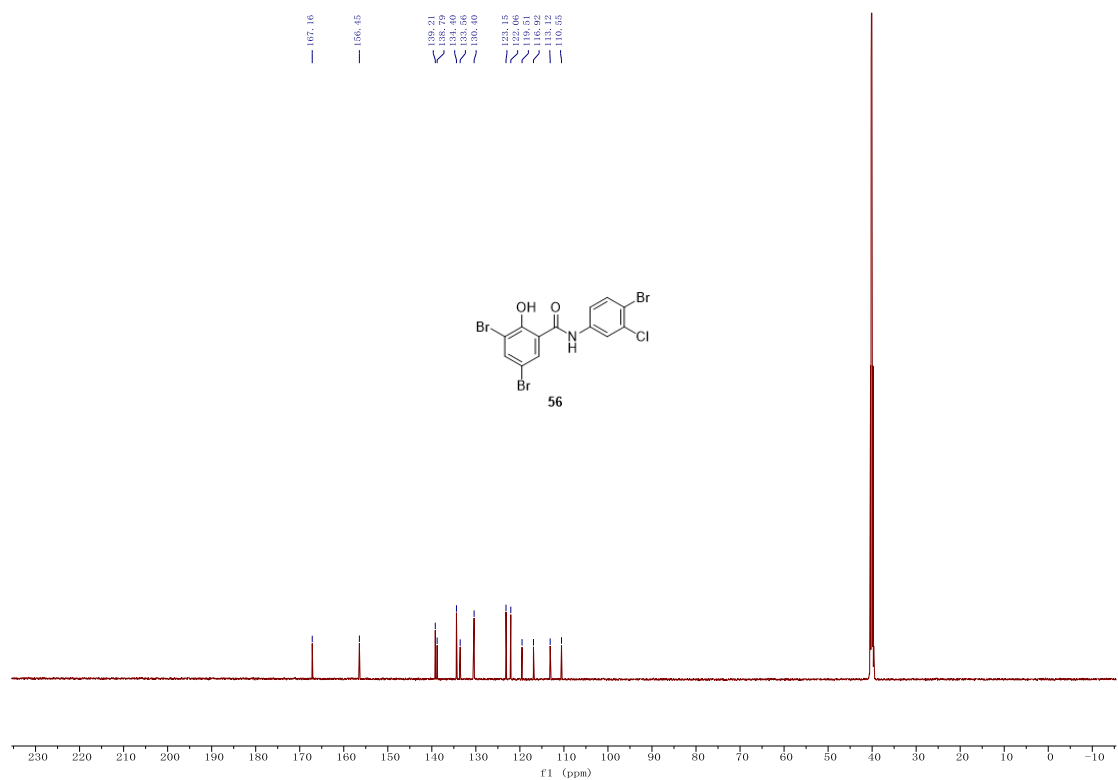






Sample_ESI_2 #24 RT: 0.24 AV: 1 NL: 2.43E6
 F: FTMS - c ESI Full ms [100.0000-1000.0000]





Sample_ESI_4 #40 RT: 0.40 AV: 1 NL: 2.00E5
 F: FTMS - c ESI Full ms [100.0000-1000.0000]

