Supporting Information

Supporting Information 1:

HPLC Analytical Method Chromatographic condition Column: -size: l = 0.15 m, $\Phi = 4.6 \text{ mm}$ -stationary phase: octadecylsilyl silica gel for chromatography (5 µm) Mobile phase: mix 50 volume of methanol and 50 volume of 0.03 mol/L monopotassium phosphate buffer (adding 2 mL of triethylamine to 1000 mL of 0.03 mol/L mono potassium solution, ajust pH to 4.0 by phosphoric acid). Injection volumn: 10 µL Flow rate: 1.2 mL/min Detection: spectrophotometer at 238 nm Test solution: with the aid of ultrasound, dissolve 25 mg of the substance to be examined in 40 mL of mobile phase and dilute to 50.0 mL with the same solvent.

Supporting information 2: Chromatograms and spectrograms





Sorted By:SignalMultiplier:1.0000Dilution:1.0000Use Multiplier & Dilution Factor with ISTDs

Signal 1: VWD1 A, Wavelength=238 nm

Peak #	RetTime [min]	Туре	Width [min]	Area mAU *s	Height [mAU]	Area %
1	3.279	BV	0.0604	71.04477	17.00593	0.4145
2	3.587	VV	0.1888	220.21754	16.85555	1.2850
3	3.810	VB	0.1063	114. 43090	16.08986	0.6677
4	8.158	BB	0.2565	225.03416	13.34277	1.3131
5	9.398	BB	0.3573	77.53874	3.08524	0.4524
6	13.854	BV	0.4416	503.77908	17.01027	2.9395
7	15.263	VB	0.5940	1.59261e4	392.69418	92.9278
Total	s :			1.71381e4	476.08381	



Figure S2. MS spectrogram of Impurity 2.



Figure S3. ¹H-NMR spectrogram of Impurity 2 in CDCl₃.



Figure S4. ¹³C-NMR spectrogram of Impurity **2** in CDCl₃.



Figure S5. HPLC Chromatogram of Impurity 3.

Area Percent Report

Sorted By : Signal Multiplier : 1.0000 Dilution : 1.0000 Use Multiplier & Dilution Factor with ISTDs

Signal 1: VWD1 A, Wavelength=238 nm

Peak #	RetTime [min]	Туре	Width [min]	Area mAU *s	Height [mAU]	Area %
1	2. 197	BV	0. 1488	289. 90604	30. 10201	1. 7338
2	2.951	BB	0.1204	55.88548	6.81855	0.3342
3	3.794	VB	0.1239	22.99636	2.78789	0.1375
4	6.748	MM	0.4624	119.61582	4.31179	0.7154
5	8.462	MM	0.5695	18.56132	5.43189e-1	0.1110
6	11.660	MM	1.0091	39.07751	6.45429e-1	0.2337
7	14.649	MM	0.3939	10.86778	4.59875e-1	0.0650
8	17.480	MM	0.6898	34.83944	8.41755e-1	0.2084
9	19.747	BB	0.4542	254.21104	8.38398	1.5204
10	23.343	MM	0.9893	1.58171e4	266. 45694	94.5969
11	26.989	MM	0.6619	23.79769	5.99192e-1	0.1423
12	41.584	MM	1.1595	33.66652	4.83932e-1	0.2013
Totals :				1.67205e4	322. 43453	



Figure S6. MS spectrogram of Impurity 3.



Figure S7. ¹H-NMR spectrogram of Impurity 3 in CDCl₃.



Figure S8. ¹³C-NMR spectrogram of Impurity **3** in CDCl₃.





Area Percent Report

Sorted By:SignalMultiplier:1.0000Dilution:1.0000Use Multiplier & Dilution Factor with ISTDs

Signal 1: VWD1 A, Wavelength=238 nm

Peak #	RetTime [min]	Туре	Width [min]	Area mAU *s	Height [mAU]	Area %	
1	19. 513	MM	0. 4490	9. 78214	3. 63075e-1	0. 0557	
2	37.526	MM	1.0145	41.08688	6.74983e-1	0.2338	
3	43.478	BB	0.9987	1.75203e4	272.64084	99.7105	
Total	s:			1.75711e4	273, 67890		



Figure S10. MS spectrogram of Impurity 4.



Figure S11. ¹H-NMR spectrogram of Impurity **4** in DMSO-*d*₆.



Figure S12. ¹³C-NMR spectrogram of Impurity 4 in DMSO- d_6 .



Figure S13. MS spectrogram of Compound 17.



Figure S14. ¹H-NMR spectrogram of Compound **17** in DMSO-*d*₆.



Figure S15. ¹³C-NMR spectrogram of Compound 17 in DMSO-*d*₆.



Figure S16. MS spectrogram of Compound 18.



Figure S17. ¹H-NMR spectrogram of Compound **18** in DMSO-*d*₆.



Figure S18. ¹³C-NMR spectrogram of Compound **18** in DMSO- $d_{6..}$



Figure S19. MS spectrogram of Compound 19.



Figure S20. ¹H0-NMR spectrogram of Compound 19 in DMSO-*d*₆.



Figure S21. ¹³C-NMR spectrogram of Compound 19 in DMSO- d_6 .





Area Percent Report

Percent Report

Sorted By : Signal Multiplier : 1.0000 Dilution : 1.0000 Use Multiplier & Dilution Factor with ISTDs

Signal 1: VWD1 A, Wavelength=238 nm

Peak #	RetTime [min]	Туре	Width [min]	Area mAU *s	Height [mAU]	Area %
1	13.861	MM	0.4149	42.41013	1.70344	0.3161
2	24.287	BB	0.7832	1.31751e4	251.99783	98.1915
3	28.227	MM	0.9895	165.30312	2.78439	1.2320
4	32.329	MM	0.8895	34.95171	6.54885e-1	0.2605
Totals	5:			1.34178e4	257.14055	



Figure S23. MS spectrogram of Impurity 5.



Figure S24. ¹H-NMR spectrogram of Impurity **5** in CDCl₃.



