

Identity and Purity

Proton NMR Spectrum

Identity: the signals are consistent with the proposed structure and in accord with literature where available.

Residual Solvents: no residual solvents are observed.

Impurities: no significant impurities are evident in the spectrum.

Carbon-13 NMR Spectrum

Identity: the signals are consistent with the proposed structure and in accord with literature where available.

High-resolution Mass Spectrum (ESI+)

Found m/z 561.1539. $C_{27}H_{29}F_3NaO_6S$ $[M+Na]^+$ requires m/z 561.1535. The deviation of 0.7 ppm is within normally accepted limits for the establishment of identity by HRMS.

HPLC

A sharp, symmetrical peak is observed (99.6 %). Note: in the absence of reference materials for preparing calibration curves, it is assumed that all peaks have the same detector response. Where possible, the conditions of analysis follow a pharmacopeial or literature method, or have been adapted from same.

Elemental Analysis

	Found:	C 60.20, H 5.54 %
$C_{27}H_{29}F_3O_6S$	Requires:	C 60.21, H 5.43 %

The elemental analyses fall within generally accepted limits for establishing the molecular formula given. The results may also be taken to imply the absence of significant quantities of water or inorganic salts (which have not been elsewhere tested for because of sample size limitations).

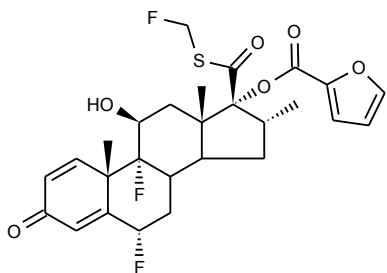
The available quantity of custom-synthesised material is always small, and this limits the extent and type of analytical data which can be obtained. This Certificate is presented in descriptive format for use by analytical chemists who are trained in the use of custom-synthesised materials. Custom materials often contain higher levels of residual solvents and/or water, and we urge you to use the corrected purity where needed rather than the raw HPLC purity. This compound is intended for use as an analytical reference material and it is not for human administration. Structures are shown with relative stereochemistry unless otherwise specified.

The re-test date is assigned from experience gained with the material in the laboratory and/or on storage. It is not possible to perform formal storage studies because of the small amount of material available.

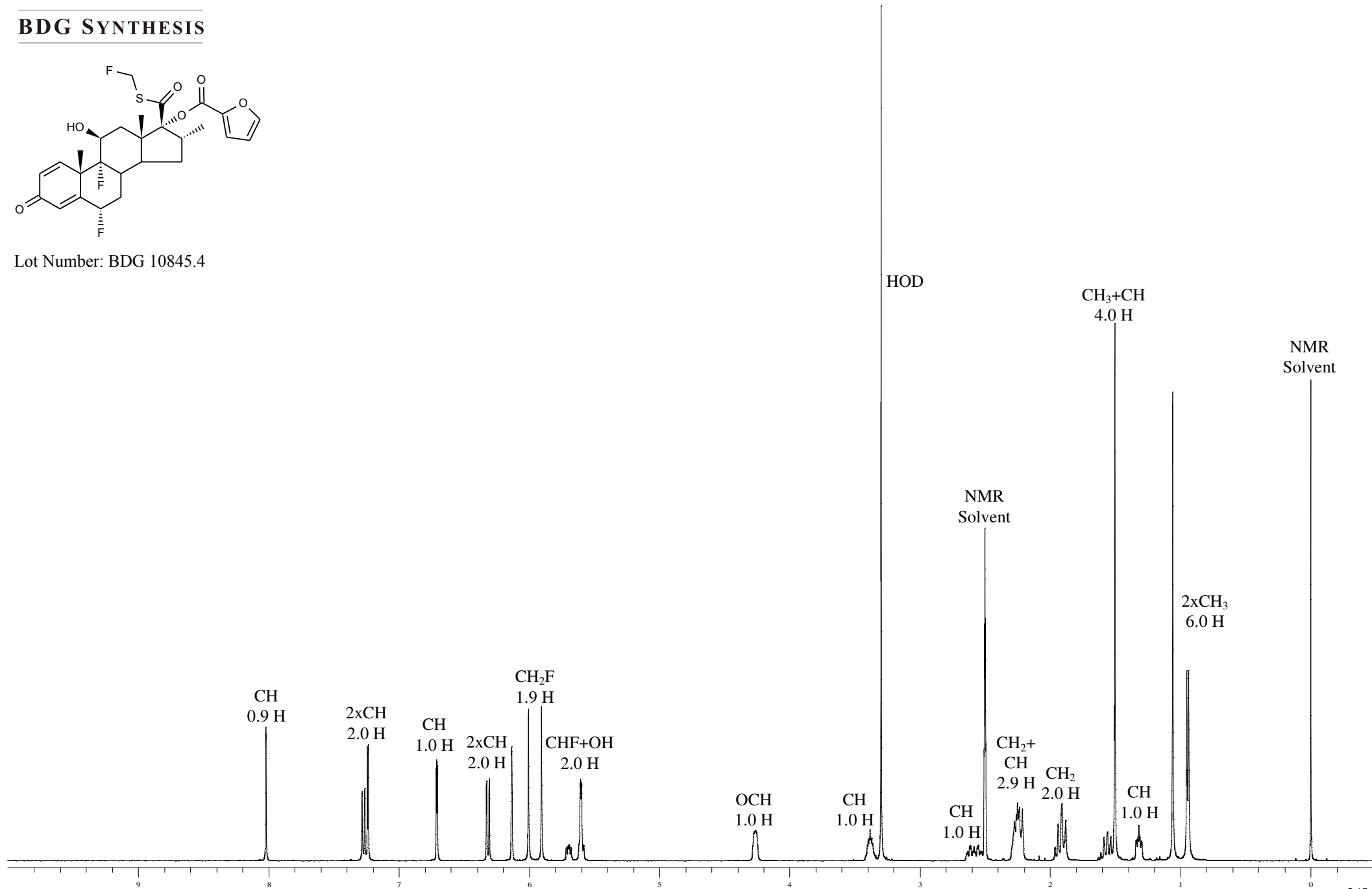


Proton NMR Spectrum of Fluticasone Furoate in DMSO-d₆

BDG SYNTHESIS



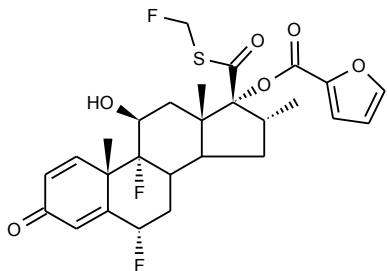
Lot Number: BDG 10845.4



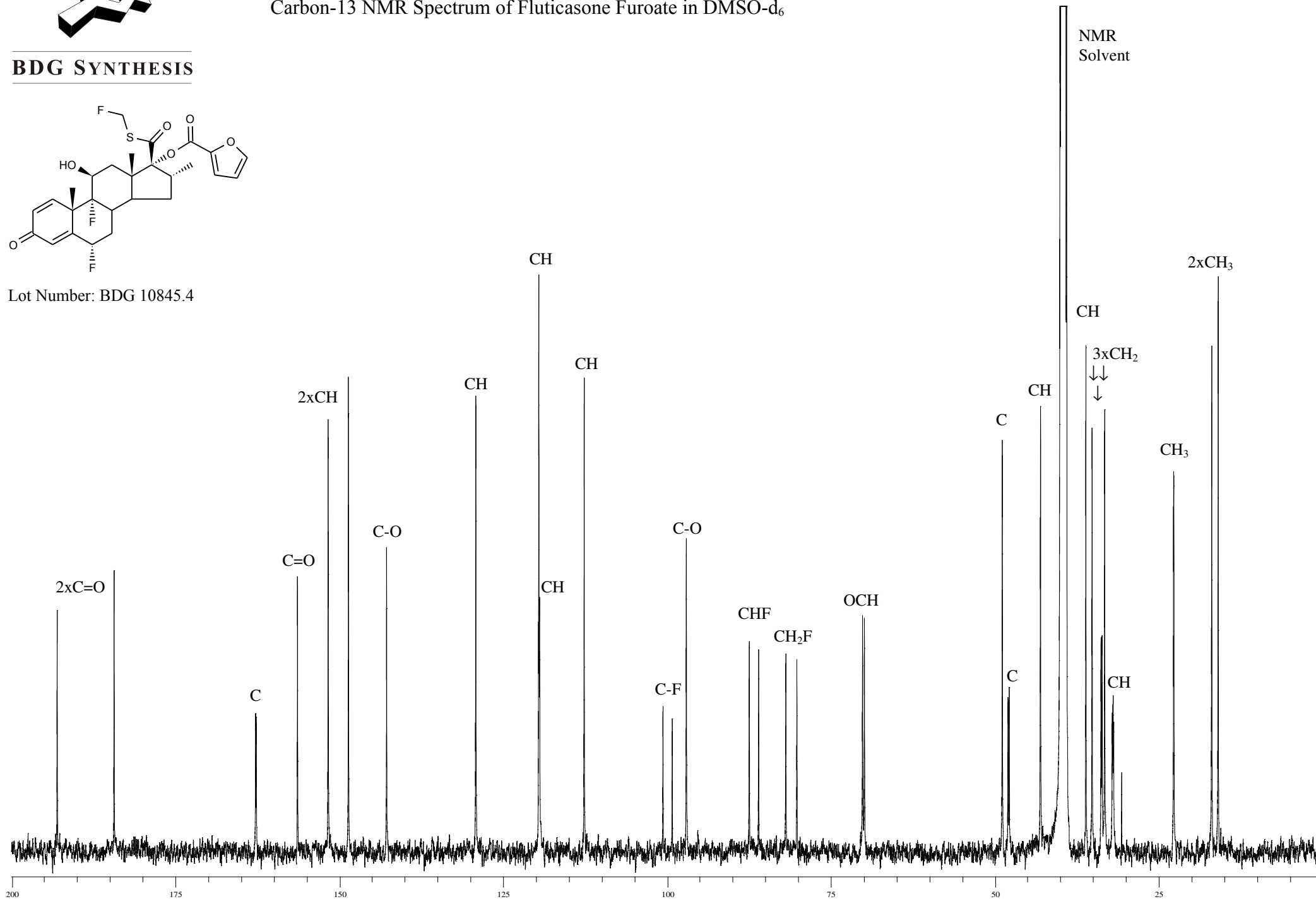


Carbon-13 NMR Spectrum of Fluticasone Furoate in DMSO-d₆

BDG SYNTHESIS



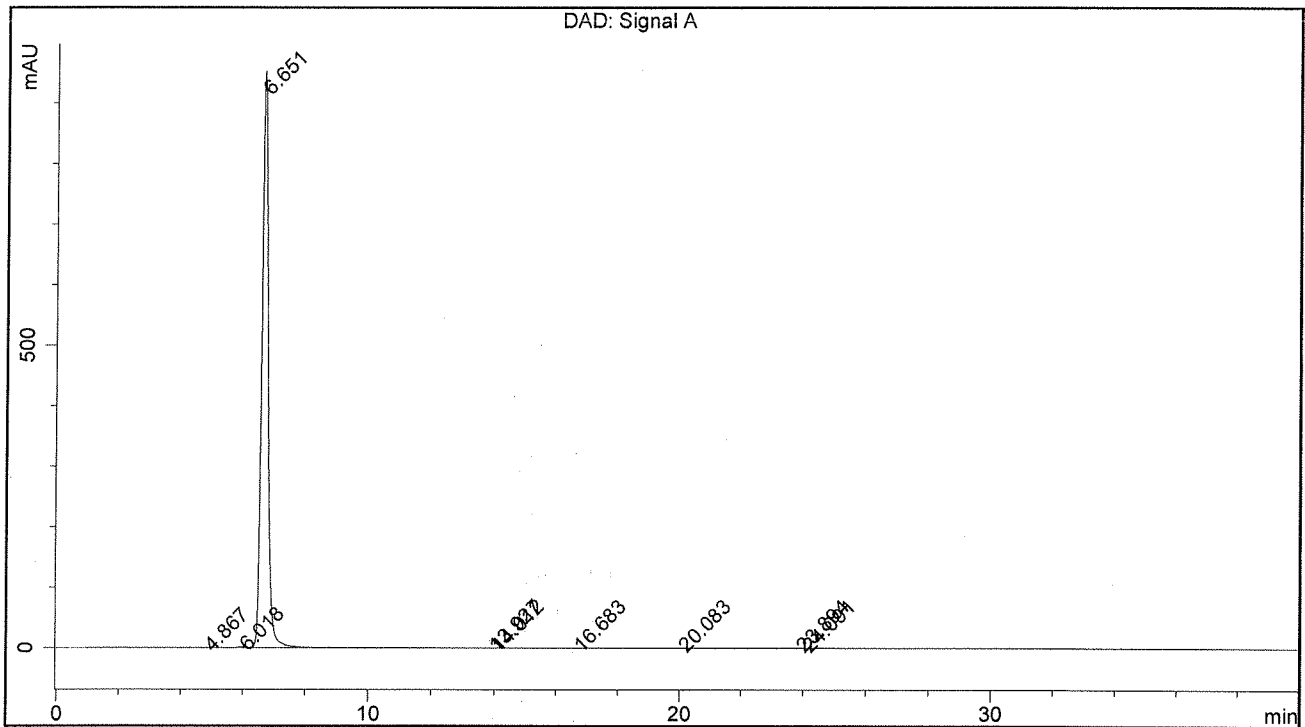
Lot Number: BDG 10845.4



BDG - Analysis of Fluticasone Furoate

Column : Phenomenex Luna C18 5um 250 x 4.6 mm
 Guard : Phenomenex Security Guard C18 4 x 3 mm
 Mobile Phase : 30:70 Water : Acetonitrile Flow Rate : 1.0 mL/min
 Sample Solvent : Mobile Phase Injection Volume : 10 uL
 Column Temperature : 30C Detection : UV at 241 nm

Sample Name	BDG 10845.4	Instrument	AnalyticalLC01
Acquisition	29/04/2014, 18:36:41	Method (rev.)	LC10394b (6)
Sequence	BDG_29Apr2014b - Reprocessed	Vial Position	32
Operator	solvation010\cerityadmin	Injection	1 of 1



Area Percent Report

Peak#	RT	Peak Height	Peak Area	Width	Area %
1	4.87 min	0.1984	2.7859	0.2118 min	0.020 %
2	6.02 min	1.6430	39.9155	0.3374 min	0.286 %
3	6.65 min	952.4092	13885.4642	0.2241 min	99.593 %
4	13.93 min	0.1083	0.6863	0.0859 min	0.005 %
5	14.01 min	0.1062	1.4300	0.1671 min	0.010 %
6	16.68 min	0.1181	2.8981	0.3043 min	0.021 %
7	20.08 min	0.1453	3.9758	0.3292 min	0.029 %
8	23.89 min	0.1365	1.9598	0.1835 min	0.014 %
9	24.09 min	0.1462	3.1407	0.2647 min	0.023 %