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Title Source rock analysis well 15/12-5		
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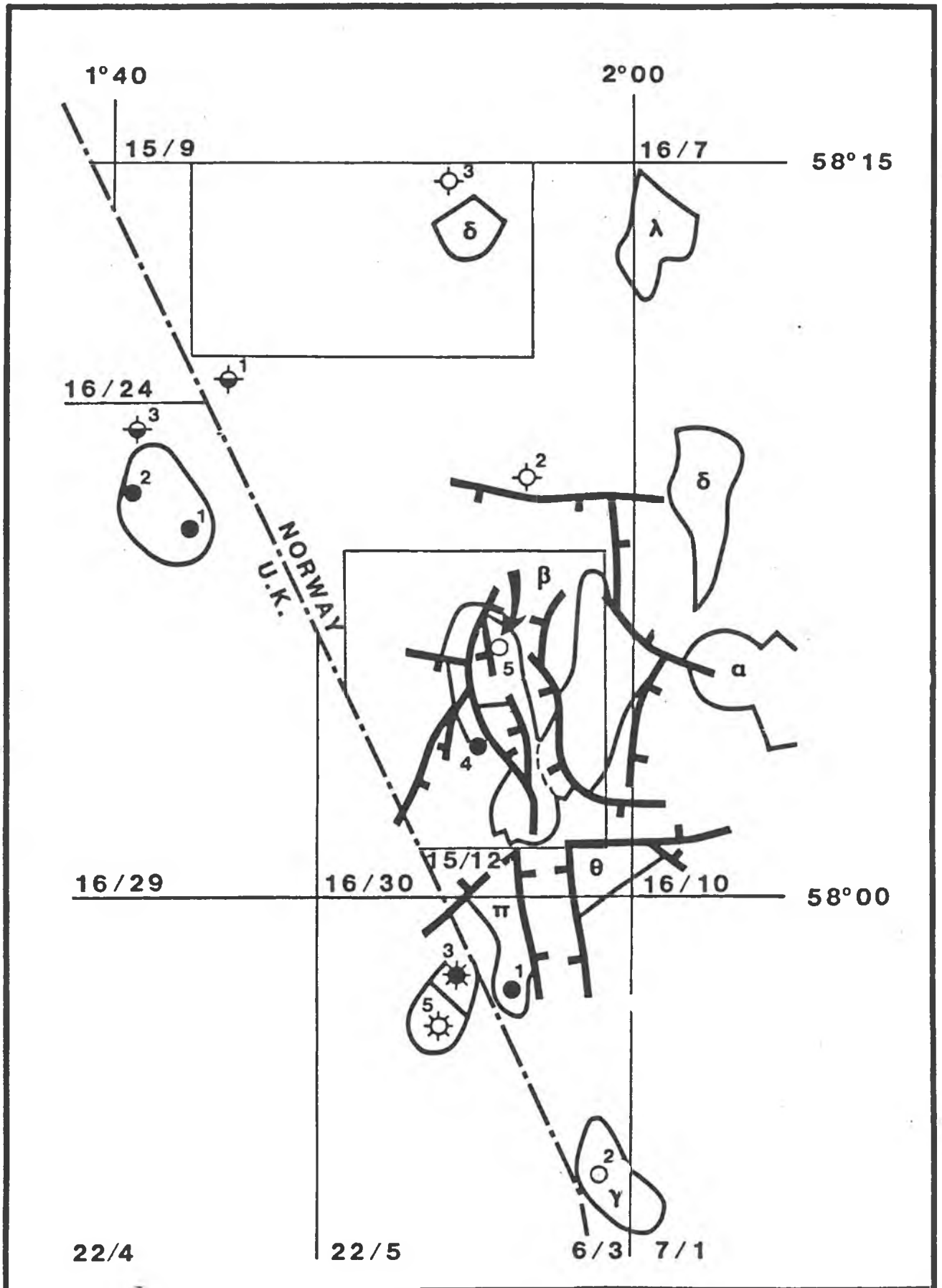
Abstract The Upper Jurassic shales are fairly immature on the 15/12-5 structure, and seem to have a somewhat limited source rock quality. Migrated hydrocarbons extracted from the reservoir sandstone cores are definitely more mature, but might have been generated from the same Upper Jurassic source rock in a more mature situation and probably with a slight change in facies.

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LOCATION MAP



INTRODUCTION

The analytical work presented in this report have been performed by IKU.

The added evaluation have been done at section for organic geochemistry, GEOLAB, Statoil.

On page 2 the location map is giving the well position within block 15/12.

UPPER JURASSIC SOURCE ROCKS

Cuttings, SWC's and conventional cores have been analysed from the Draupne (47 cm thick) and Heather (20 m thick) formations.

An approximate average for the total organic carbon in the Heather formation has been estimated to 1.5 %. On structure, the formation seems to be gradually more silty towards the bottom, and a more shaly facies off structure might have a better source rock quality.

The Draupne formation seems to have a more limited quality on this locality than normally observed for the Upper Jurassic hot shale. The total organic carbon is slightly below 2.5 % on average (1.9 - 2.8 %), while the production potential varies between 2 and 7.6 kg hydrocarbons/tonne rock. The HI-values are below 300 mg HC/gTOC (107 - 247). This seems, at the best, to classify the formation as a medium good source rock at an expected maturity level below $R_o = 0.7$. Data from quantified pyrolysis also indicate a somewhat reduced kerogen quality. Only up to 22 % of the produced hydrocarbons was within the $C_{15}+$ range. Qualitatively the amount of alkane/alkene pairs are significant, but not dominating. The optical analysis indicates over 80 % amorphous material. The biogenic source was however not possible to establish, but a smaller part of liptinitic material (5 - 10 %) contained significant amounts of terrestrial palynomorphs. Very high values for the pristane/phytane ratio in extracts from the formation and the low TOC-values seem to confirm a more coastal, less reducing environment for the deposition than what is usually the case. This goes especially for the Draupne formation. Significant amounts of higher normal alkanes in the C-25 to C-35 range also contributes to an impression of a significant element of higher land-plants in the kerogen. The amount of extractable material varies between 2000 and 4500 ppm.

The level of maturity, is however too low on structure to allow for any significant contribution to the observed accumulated hydrocarbons. A maturity level lower than the first main oil generating phase for kerogen type II is confirmed by a spore colour index between 2.5 - 3 corresponding to a vitrinite reflection value of $R_o = 0.40$ (in accordance to IKU correlation scheme). Biomarkers in extracts from the source rock confirms this level by 55 % $\beta\beta$ (m/z) 217, low values for % 20S (38 %), low $\alpha\beta$ / $\alpha\beta + \beta\alpha$ (85 %), high Tm/Ts values (3.5 - 7.4).

The chromatograms of the saturate fractions are also pointing at a low maturation level. This is especially indicated by the relatively high concentration of isoprenoids and a small but significant bimodale distribution. Correspondingly, the aromatic functions are showing values for the methyl phenanthrene index indicating low maturity. However, in the coresamples near the reservoir interval the component distribution is very abnormal. The chromatograms show a strong dominance of light hydrocarbons. The content of normal alkanes decreases drastically towards n-C19, followed by a low but constant concentration of heavier homologs towards n-C32. Whether this has any relationship to the observed close to 100 % "possible additives" (lignite) (see IKU p. 17) in cutting samples is difficult to prove. "Possible additives" was discovered in samples from the interval 2884 - 2900 m, 2900 - 2915 m and with decreasing amounts in samples from the interval 2915 - 2960 m.

Still, the isotopic composition of extracts and practions from the source rocks does not diverge from extracted hydrocarbons from the reservoir zone. This probably indicate that the accumulated hydrocarbons are generated from the same or highly similar source rocks at a higher level of maturity. Surprisingly, however, a fairly large difference was observed in the isotopic composition between the kerogen concentrate and the total extract.

CONCLUSION

The upper Jurassic source rocks on the 15/12-5 structure are fairly immature. The production potential classifies the source rocks as medium good. The kerogen seems influenced by significant amounts of terrestrial material, possibly deposited in a somewhat less reductive and more coastal than usually is the case. The accumulated hydrocarbons are therefore not produced on structure by the upper Jurassic shales.

ACCUMULATED/MIGRATED HYDROCARBONS

Headspace gas from sealed canned samples containing mud and cuttings have been analysed for the composition of light hydrocarbons. As table 1a - c (IKU) indicate, the total amount of methane and heavier gas-components is significant in parts of the sealing rock. The gas has probably mainly been generated at a higher maturity level than on structure, as indicated by a relatively low $i-C_4/n-C_4$ ratio and a significant wetness.

Similar gas compositions as observed in the sealing source rock has been observed through the hole reservoir zone. The definitely highest gas concentration was indicated in samples from 2915 to 2960 m.

Core samples from the reservoir interval have been extracted, and the total amount of extractable hydrocarbons and ratios of various fractions are given in table 6 -8. The highest concentration of hydrocarbons was observed near 2927.55 m, with a sharp decrease at 2952.65 m. The pristane/phytane ratio seems to vary little, between 1.2 and 1.5, typical values for North Sea oils generated from mature Upper Jurassic source rocks. For comparison, extracts from the source rocks show values between 2.4 and 3.5, and probably confirming.

- a) the significantly higher maturity level in the migrated hydrocarbons relatively to the source rock on the structure and
- b) that the environment of deposition has been more reductive in the drainage area for the migrated hydrocarbon.

The ratio seems to exclude any significant contribution for possible coal or coaly shale source rocks.

The biomarkers indicate a significantly higher level of maturity for the migrated hydrocarbons than for the extracts from the source rock.

Isotopically, the migrated hydrocarbons seem to have been generated by the Upper Jurassic shales, but at a higher level of maturity.

REPORT

REPORT TITLE:

DATAREPORT

SOURCE ROCK EVALUATION AND CHARACTERISATION OF
HYDROCARBONS FROM WELL 15/12-5

REPORT NO.: 22.1850.00/02/87

AUTHORS:

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REPORT

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CLIENT:
Statoil, Geolab, v/ Steinar Ulvøen. Open contract T.4533, job no. 25.

SUMMARY:

This datareport presents the data of source rock evaluation and characterisation of hydrocarbons from well 15/12-5.

<i>KEY WORDS:</i> Well 15/12-5	
Source rock	
Asphaltenes	

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1. SUMMARY

Samples from well 15/12-5 were sent from Statoil to IKU for organic geochemical analysis. The aim of this project was to give an indication of quality and saturation of the hydrocarbons in the reservoir zone from 2918 m, and unconformities in the cap rock which is Draupne Fm (2841 m) and Heather Fm (2888 m). The job was assigned the IKU project code 22.1850.00 and was carried out according to the terms in the open contract T-4533 job no. 25.

The samples received for analysis were as follows:

- 199 canned cutting samples
- 13 sidewall cores (swc)
- 16 core samples

The canned cutting samples covered a depth interval from 234 m to T.D., while the samples which should be chosen for this project were in the interval from 2840 m to T.D.

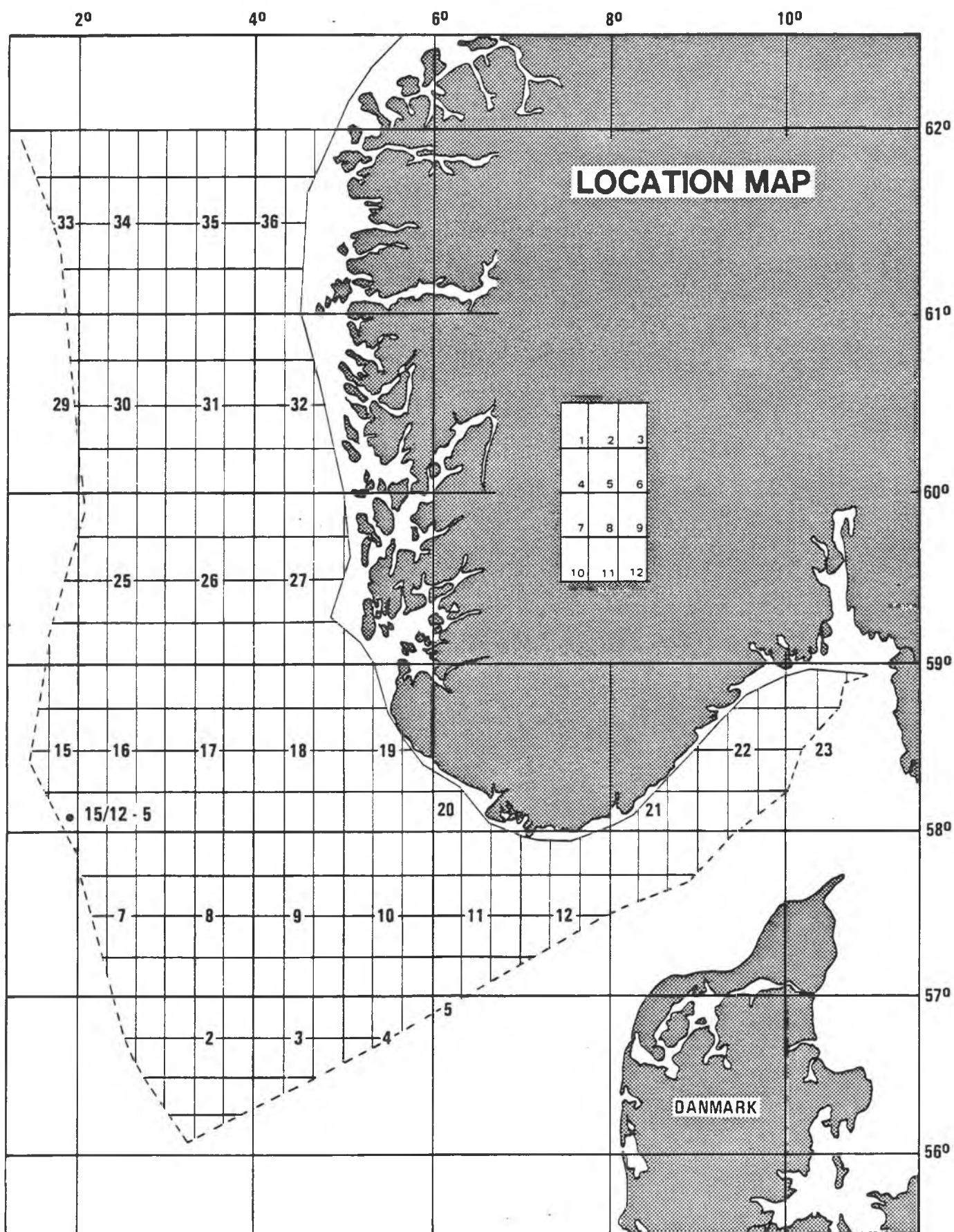
The following analysis were performed:

- | | |
|--|------------|
| - Headspace and occluded gas analysis | 20 samples |
| - Sample preparation, i.e. washing, sieving and handpicking, lithological description | 20 samples |
| - Total organic carbon | 15 samples |
| - Rock-Eval pyrolysis | 15 samples |
| - Kerogen description, TAI | 4 samples |
| - Preparation of kerogen concentrates | 4 samples |
| - Pyrolysis-GC quantification | 5 samples |
| - Extraction | 20 samples |
| - Asphaltene precipitation | 20 samples |
| - Iatroscan quantification | 20 samples |
| - MPLC separation | 20 samples |
| - GC of saturated hydrocarbons | 20 samples |
| - GC of aromatic hydrocarbons | 10 samples |
| - GC-MS of biomarkers | 10 samples |
| - $^{13}\text{C}/^{12}\text{C}$ isotope ratios of extracts, saturates, aromatics, NSOs and asphaltenes | 10 samples |
| - $^{13}\text{C}/^{12}\text{C}$ isotope ratios of kerogen concentrates | 4 samples |

Samples from interval 2840-2889 m were chosen for characterisation of the source rock, and 15 samples (10 cuttings and 5 sidewall cores) from this interval were analysed by TOC and Rock-Eval pyrolysis. The sample material from this interval was of poor quality and for preparation of kerogen concentrates and kerogen description only 4 samples were selected. For extraction and chromatographic separation 4 sidewall cores (swc) from this interval were analysed, while 16 sandstone cores from the reservoir zone were analysed for characterisation of the reservoir hydrocarbons.

The weight of saturated and aromatic hydrocarbons, NSOs' and asphaltenes after chromatographic separation of extracts are not satisfactory as commented by Statoil. The difference between the amount of NSOs' and asphaltenes given in Table 6 and the amount of nonhydrocarbons given in Table 7 are too high. The testing of our chromatographic system i.e. precolumns, regeneration, reproducibility tests etc., has started and the results of these analysis will be given to Statoil as soon as they are available.

Five copies of the data report has been sent to Statoil and five copies are stored at IKU.



2. EXPERIMENTAL METHODS

2.1 Gas analyses

·Headspace gas:

A septum was attached to the can and a sample of the headspace gas was taken for analysis of C₁, C₂, C₃, i-C₄, nC₄ and C₅₊ hydrocarbons.

This analysis was performed on Carlo Erba Fractovap 2150 and 2350 gas chromatographs equipped with 2 m x 1/8" stainless steel columns filled with Porapak Q on Chromosorb using nitrogen as carrier gas. An oven temperature of 150°C was used. After elution of n-butane, the column was back-flushed and C₅₊ compounds were recorded as one peak. A standard gas sample containing methane, ethane, propane, n-butane, n-pentane and n-hexane (100 ppm each) was used for quantification.

The can was then opened and headspace volume, water volume, and sample weight were measured. The canned samples were washed with warm water (30-40°C) on 4.0, 2.0 and 0.125 mm sieves to remove drilling mud and were then dried at 35°C.

Occluded gas:

Prior to drying, an aliquot of the 2 mm fraction of each sample was crushed in water for 10 minutes using a gas-tight ball mill. The evolved gas was analysed as described for headspace gas.

2.2 Lithological descriptions

Lithological examinations were carried out using a binocular microscope (maximum 50x magnification). Colours were described on dry samples according to the "Rock Colour Chart" published in 1979 by the Geological Society of America, Boulder, Colorado. Handpicking of the cuttings for organic geochemical analyses was based on the lithological descriptions.

2.3 Total Organic Carbon

Bulk samples were crushed in a mortar. Aliquots of the samples were then weighed into Leco crucibles, treated three times with hot 10% HCl

to remove carbonate, and washed 4 times with distilled water to remove any traces of HCl. The crucibles were then placed on a hot plate (60°C) and dried for 24 hours. The total organic carbon (TOC) content of the dried samples was determined using a Leco CS 244 carbon analyser.

The instrument was calibrated using a Triassic shale (4.96% TOC) as a standard. Calibration and blank runs were carried out every 15th measurement.

2.4 Rock-Eval pyrolysis

Powdered sample (100 mg shale, 5-10 mg coal) was weighed into a platinum crucible (the base and cover of which are made of sintered steel), and analysed on a Rock-Eval II pyrolyser (Girdel/IFP) under standard conditions (Carrier gas: Helium; Flow Rate: 100 ml/min.; Temp. Program: 300°C (3 min.) - 25°C/min. - 550°C).

2.5 Preparation of kerogen concentrates

The powdered and pre-extracted rock sample was treated overnight with concentrated HCl (200 ml), followed by treatment with hot 40% HF (100 ml, 50°C, overnight). The acids were removed by decanting and repeated washing with hot water (2 hours), 10% ammonium carbonate solution (> 2 hours) and cold water (2x2 hours).

The kerogen was further purified by extraction with methanol, methanol/DCM (1:1) and hexane/DCM (1:1). The dried kerogen sample was then analysed for Total Carbon content.

2.6 Evaluation of kerogen type and thermal maturity in transmitted light

The rock samples were crushed and treated with hydrochloric acid and hydrofluoric acid in order to remove the mineral matter and isolate the kerogen. The isolated organic matter/kerogen was mounted in glycerine jelly on microscope slides as a strew mount.

The treatment of the isolated kerogen varied according the requirements and preservation of the samples. This included:

- T-slide: represented the total acid residue
- O-slide: represented the kerogen debris remaining after flotation in $3nBr_2$ to remove heavy minerals
- N-slide: represented a sieved residue (15 μm mesh)
- X-slide: represented oxidised residue.

T and/or O slides were necessary in order to evaluate the kerogen composition/palyno-facies of the samples.

N and/or X slides were required if it was necessary to concentrate larger kerogen fragments or spores/pollen for age determination or spore colour (NB: oxidised residue is not used for spore colour determination as oxidation will alter the colour).

The samples were studied using a Leitz Dialux microscope with a white halogen light source. Objective lenses of 10x and 63x magnification were used. The higher magnification allowed a more detailed description of the kerogen, with resolution of particles down to 2 μm diameter. The thermal alteration index (TAI) measurements were obtained from the colour of spores and pollen. The techniques used in visual kerogen study are adapted from Staplin (1969) and Burgess (1974).

COMPARISON OF MICROSCOPIC THERMAL MATURATION SYSTEMS AT IKU

MATURATION LEVEL	SPORE COLOUR		VITRINITE REFLECTANCE	SPORINITE FLUORESCENCE
	TAI	NEWTAI	%	
IMMATURE	1 - 2-	1.0- 3.5	0.25-0.40	1-4
EARLY MATURITY	2- - 2	3.5- 4.0	0.40-0.60	4-6
MODERATE MATURITY	2 - 2+	4.0- 4.5	0.60-0.80	6-7
MATURITY	2+ - 3-	4.5- 5.5	0.80-1.00	7-8
LATE MATURITY	3- - 3	5.5- 6.0	1.00-1.20	8-9
POST-MATURE	3 - 5	6.0-10.0	>1.20	Absent

2.7 Pyrolysis Gas Chromatography (Py-GC Programmed)

Thermal extraction (S1)

The whole rock (20-30 mg) or kerogen sample (<1 mg) was placed in a boat-shaped sample probe and thermo-extracted in a stream of helium at 340°C for 4 minutes.

Pyrolysis-Gas Chromatography (S2)

The thermo-extracted sample was pyrolysed in a helium atmosphere using a furnace type pyrolyzer as described by Solli et al. (1984). The pyrolysis temperature program was 340°C - 50°C/min. - 550°C. The outlet of the furnace was directly connected to a splitter (30:1) which enabled the pyrolysis products to be simultaneously recorded as a bulk "S2" peak by an FID and passed into a gas chromatograph. The pyrolysis products were trapped on the chromatographic column by cooling the front part of the column with liquid nitrogen. On completion of the pyrolysis programme, the collected pyrolysis products were injected onto the column by removing the nitrogen batch. The GC conditions are outlined below.

GC conditions

Column: 30 m DB-1, i.d. 0.32 mm, fused silica capillary column.

Carrier gas: Helium, inlet pressure 7.5 psi; flow rate 1.5 ml/min.

Temp. programme: -10°C (1 min.) - 5°C/min. - 310°C (10 min.)

2.8 Extractable Organic Matter

Powdered rock was extracted by Soxhlet technique with boiling dichloromethane (DCM) containing 7% methanol (v/v) as solvent.

DCM of organic geochemical grade was used and blank analyses indicated that the occurrence of contaminating hydrocarbons was negligible. Activated copper filings were used to remove any free sulphur from the samples.

After extraction, the solvent was evaporated off using a Büchi Rotavapor and the amount of extractable organic matter (EOM) was determined.

2.9 Separation of extractable organic matter (EOM) fractions

Iatroscan

The extractable organic material was diluted with DCM and ca. 20-30 μg of the solution were applied to a precleaned and activated Chromarod (type SII). Three replicates of each sample were analysed. The rods were fully developed with n-hexane (10-11 cm), then dried and redeveloped to half-rod height (5.5 cm) in toluene, and finally developed in DCM+MeOH (93:7v/v, 2.5 cm).

The rods were passed through the flame-ionisation detector (FID) of an Iatroscan TH-10 ($\text{H}_2 = 160 \text{ ml/min}$; air = 2 l/min.; scan speed = 0.38 cm/sec. (gear no. 30)). The FID response was recorded and integrated using a DEC Minichrom system. The retention factors of the aliphatic and aromatic hydrocarbons were then determined by comparison with standards (nC_{20} and fluorene).

Medium-pressure liquid chromatography (MPLC)

The EOM/oil ($>210^\circ\text{C}$) sample was diluted in DCM (1:3 $\text{mg}/\mu\text{l}$) and the asphaltenes were precipitated using excess n-pentane (40:1 pentane:(DCM+EOM)). The asphaltene fraction was weighed after drying at 50°C for 12 hours.

The remaining maltenes were separated into saturated, aromatic and non-hydrocarbon fractions using an MPLC system with n-hexane as eluant (Radke et al., 1980). The various fractions were concentrated using a Büchi Rotavapor, transferred to glass vials and the remaining solvent removed.

2.10 Gas chromatographic analysis

The saturated hydrocarbon fractions were diluted with n-hexane and analysed on a Carlo Erba Fractovap GC. The GC is equipped with 15 m DB-1 fused silica column, and hydrogen is used as carrier gas with a flow rate of about 1.5 ml/min. Injections were performed in split mode (split ratio 1:10). The temperature programme used was 80°C (2 min) - $4^\circ\text{C}/\text{min}$ - 280°C .

The total aromatic fractions were diluted with n-hexane and analysed on an HP 5730A gas chromatograph, fitted with a DB-5 fused silica column

(15 x 0.25 mm i.d.), using a hydrogen carrier gas with a flow rate of 2.5 ml/min. The injection split ratio was 1:10.

The temperature programme used was 80°C (2 min.) - 4°C/min - 280°C.

Data processing for all the GC analyses was performed on a VG Multi-chrom lab data system.

2.11 Gas chromatography - mass spectrometry (GC-MS)

GC-MS analysis were performed on a VG Micromass 70-70H GC-MS-DS system. The Varian Series 3700 GC was fitted with a fused silica OV-1 capillary column (30m x 0.3 mm i.d.). Helium (1.5 ml/min) was used as carrier gas and the injections were performed in split mode (1.5 µl, split ratio 1:15).

The GC oven was programmed from 120°C (2 min.) to 280°C at 4°C/min.

The saturated hydrocarbons were analysed in multiple ion mode (MID) at a scan cycle time of approximately 2 secs. The mass spectrometers operated at 70eV electron energy with an ion source temperature of 200°C. Data acquisition was performed using VG data systems.

Peaks were identified by comparison with elution patterns in certain mass chromatograms. Peak ratios were calculated from peak heights in the appropriate mass chromatograms.

2.12 $\delta^{13}\text{C}$ isotope analysis

The $\delta^{13}\text{C}$ isotope analyses were performed by mass spectrometry at the Institute for Energy Technology (IFE) in Oslo according to their method. Their reference value for the standard NBS-22 is -29.8 ‰ (PDB).

The isotope ratios are given as delta-values (del):

$$\text{del (\%)} = ((R \text{ sample} - R \text{ stand.}) / (R \text{ stand.})) * 1000$$

$^{13}\text{C}/^{12}\text{C}$ - isotope ratios are calculated versus PDB.

Project no.: 22.1850.00
 Well ident.: 15/12-5
 DATE : 14 - 4 - 87.

Table 1a:

CONCENTRATION (ul Gas / kg dry Rock) OF C1 - C5+ HYDROCARBONS IN HEADSPACE

IKU no.	DEPTH m	C1	C2	C3	iC4	nC4	C5+	SUM C1-C4	SUM C2-C4	WET-NESS (%)	iC4 nC4
C 5870	2855	4144	1184	1324	230	434	694	7317	3172	43.36	0.53
C 5871	2870	12520	5983	10283	2262	5528	11025	36577	24056	35.77	0.41
C 5872	2885	33552	12329	13701	2966	7803	17516	70351	36799	52.31	0.38
C 5873	2889	21984	9516	14542	3504	9424	29384	58971	36987	62.72	0.37
C 5874	2900	1709	566	678	111	314	1038	3378	1669	49.41	0.35
C 5875	2915	1425	672	866	121	343	1031	3428	2002	58.41	0.35
C 5876	2930	9918	2992	4293	1018	2110	3520	20330	10413	51.22	0.48
C 5877	2945	6013	1700	2988	925	2306	6218	13932	7919	56.84	0.40
C 5878	2960	476	210	417	111	294	2379	1527	1031	67.51	0.38
C 5879	2975	177	60	132	31	81	336	482	305	63.32	0.39
C 5881	3005	1071	296	540	179	467	1729	2554	1482	58.05	0.38
C 5882	3020	2781	1213	1937	482	1176	2900	7589	4808	63.35	0.41
C 5883	3035	4755	2040	3189	815	2028	5127	12827	8072	62.93	0.40
C 5884	3050	3695	1376	2462	721	1909	5468	10162	6467	63.64	0.38
C 5885	3065	6012	2520	3193	656	1593	4465	13975	7963	56.98	0.41
C 5886	3080	4354	1762	2219	419	920	1676	9674	5320	54.89	0.46
C 5887	3095	749	336	571	127	348	1135	2131	1382	64.85	0.37
C 5888	3110	763	340	547	121	325	817	2095	1332	63.57	0.37
C 5890	3140	751	350	504	103	274	1025	1983	1231	62.09	0.38
C 5891	3180	1156	437	587	122	316	932	2628	1442	54.33	0.39

Project no.: 22.1850.00
 Well ident.: 15/12-5
 DATE : 14 - 4 - 87.

Table 1b:

CONCENTRATION (ul Gas / kg dry Rock) OF C1 - C5+ HYDROCARBONS IN OCCLUDED

IKU no.	DEPTH m	C1	C2	C3	iC4	nC4	C5+	SUM C1-04	SUM C2-04	WET-NESS (%)	iC4 / nC4
5870	2855	200	117	564	140	596	3426	1616	1417	87.65	0.24
5871	2870	203	96	566	156	722	3627	1742	1539	88.37	0.22
5872	2885	531	1612	6598	619	3040	11601	10400	11670	99.72	0.20
5873	2889	1081	459	1521	284	1518	10936	5863	4732	81.56	0.19
5874 *	2900	116	55	321	95	350	1600	937	821	87.61	0.27
5875	2915	1579	262				12457	1841	262	14.25	
5876	2930	10391	553	2162	938	3238	29001	17281	6890	39.87	0.29
5877	2945	7940	247	468		699	14936	9354	1414	15.12	0.00
5878	2960	2471	294	404		362	14124	3551	1080	30.42	0.00
5879	2975	1648	197	293		263	11305	2401	753	31.35	0.00
5881	3005	911	80	119	35	92	1311	1027	315	25.77	0.27
5882	3020	1067	117	384	155	641	7885	2364	1297	54.88	0.24
5883	3035	1055	127	408	169	633	7623	2392	1337	55.90	0.27
5884	3050	857	96	178	45	227	5054	1403	547	38.95	0.20
5885	3065	706	146	562	134	591	4221	2140	1434	67.00	0.25
5886	3080	497	106	402	94	384	2220	1483	936	66.51	0.24
5887	3095	570	62	130	30	30	1919	322	252	30.58	1.00
5888	3110	385	89	147		107	1904	1228	345	27.94	0.00
5890	3140	564	43	74	14	54	945	730	164	24.75	0.07
5891	3150	453	51	149	34	175	1366	362	409	47.53	0.15

* The 125 µm fraction (normally 2 mm) was crushed for occluded gas.

Project no.: 22.1850.00
 Well ident.: 15/12-B
 DATE : 14 - 4 - 87.

Table 1c:

CONCENTRATION (ul Gas / kg dry Rock) OF C1 - C5+ HYDROCARBONS SUMMATION

IKU no.	DEPTH m	C1	C2	C3	iC4	nC4	C5+	SUM C1-C4	SUM C2-C4	WET-NESS (%)	iC4 nC4
C 5870	2855	4344	1300	1888	370	1030	4119	8933	4589	51.37	0.36
C 5871	2870	12723	6079	10649	2419	6250	16650	38319	25596	66.80	0.39
C 5872	2885	34083	13941	20299	3585	10844	29017	62751	48668	68.81	0.33
C 5873	2889	23066	9975	17064	3788	10942	40319	64834	41768	64.42	0.35
C 5874 *	2900	1825	621	998	206	664	2638	4315	2490	57.70	0.31
C 5875	2915	3004	934	866	121	343	13489	5269	2265	42.98	0.35
C 5876	2930	20309	3545	6455	1955	5349	32521	37612	17303	46.00	0.37
C 5877	2945	13983	1947	3486	925	3005	21156	23286	9334	40.08	0.31
C 5878	2960	2967	504	821	104	676	16503	5078	2111	41.57	0.16
C 5879	2975	1825	257	425	31	344	11943	2882	1058	36.70	0.09
C 5881	3005	1982	376	659	205	560	3540	3791	1799	47.57	0.37
C 5882	3020	3843	1330	2321	537	1818	10786	9953	6105	61.34	0.35
C 5883	3035	5810	2168	3597	984	2660	12748	15219	9409	61.83	0.37
C 5884	3050	4551	1471	2640	766	2137	10522	11565	7014	60.85	0.36
C 5885	3065	6718	2666	3756	791	2184	8686	16104	9396	58.31	0.36
C 5886	3080	4850	1868	2621	513	1304	3896	11157	6306	56.62	0.39
C 5887	3095	1319	397	701	157	378	3054	2953	1634	55.34	0.42
C 5888	3110	1648	429	694	121	431	2722	3323	1675	50.41	0.23
C 5890	3145	1316	393	573	117	328	1970	2732	1417	51.85	0.36
C 5891	3130	1639	468	716	155	491	2795	3490	1881	53.85	0.32

* The 125 µm fraction (normally 2 mm) was crushed for occluded gas.



Lithology and Total Organic Carbon measurements

TABLE NO.: 2.
WELL NO.: 15/12-5

Sample	Depth (m)	TOC	Lithology
C-5870	2840-2855	0.41	*60% <u>Claystone</u> , greyish green, waxy, blocky to subfissile, calcareous
		1.92	*20% <u>Claystone</u> , light olive grey to brownish grey, micromicaceous, blocky, noncalcareous.
			10% <u>Claystone</u> , greyish red to dark reddish brown, blocky to subfissile and stringy calcareous
			10% <u>Chalk/Limestone</u> , white, occasionally pink Sm.am. Sand; Pyrite; Claystone, black, fissile to subfissile; Claystone, grey, subfissile to blocky
C-5871	2855-2870	2.45	*40% <u>Claystone</u> , sandy and silty, black, dark, grey, dark brownish grey, micromicaceous, fissile to subfissile, slightly calcareous to calcareous
		0.51	*30% <u>Claystone</u> , greyish green, waxy, blocky to subfissile, calcareous
			20% <u>Claystone</u> , greyish red to dark reddish brown, blocky to subfissile and stringy, calcareous 10% <u>Chalk, Limestone</u> , white, brownish white; Sand; Pyrite
C-5872	2870-2885	2.77	*30% <u>Claystone</u> , sandy and silty, black, dark grey, dark brownish grey, calcareous
		0.41	*30% <u>Claystone</u> , greyish green, waxy, subfissile, calcareous
		1.72	*20% <u>Claystone</u> , light olive grey, blocky, non-calcareous 20% <u>Claystone</u> , greyish red to dark reddish brown, calcareous
			Sm.am. Coal; <u>Chalk/Limestone</u> , white, pink; Sand; Pyrite



Lithology and Total Organic Carbon measurements

TABLE NO.: 2.
WELL NO.: 15/12-5

Sample	Depth (m)	TOC	Lithology
C-5873	2885-2889	2.62	*30% <u>Claystone</u> , sandy and silty, black, dark grey, dark brownish grey, calcareous
		0.44	*20% <u>Claystone</u> , greyish green, waxy, subfissile to blocky, calcareous
		1.83	*20% <u>Claystone</u> , light olive grey to greyish brown, non-calcareous to slightly calcareous
			20% <u>Claystone</u> , greyish red to dark reddish brown, calcareous
			10% <u>Sand</u> , fine, single grains, subangular to subrounded
		Sm.am. Coal, black; Limestone, white	
C-5874	2889-2900		100% Possibly additives (lignite) Sm.am. Sand, fine, angular to subrounded; Claystone, similar to above lithology
C-5875	2900-2915		100% Possibly additives (lignite) Sm.am. Sand; Claystone
C-5876	2915-2930		80% <u>Sandstone</u> , fine to medium, well cemented, silica cement 20% Possibly additives (lignite) Sm.am. Claystones
C-5877	2930-2945		90% <u>Sand/Sandstone</u> , medium, silica and calcite cemented, subangular to subrounded 10% Possibly additives (lignite) Sm.am. Claystones; Chalk, white
C-5878	2945-2960		90% <u>Sand/Sandstone</u> , medium to coarse, subangular to subrounded, no staining observed 10% Possibly additives (lignite)



Lithology and Total Organic Carbon measurements

TABLE NO.: 2.
WELL NO.: 15/12-5

Sample	Depth (m)	TOC	Lithology
C-5879	2960-2975		100% <u>Sandstone/Sand</u> , medium to very coarse, subangular to subrounded, some brown coating on grains Sm.am. Possibly additives; Sandstone, grey, very fine to fine
C-5881	2990-3005		100% <u>Sandstone</u> , medium to fine, angular to subangular, well cemented, calcite and silica cement, occasionally some coarse and very coarse grains, occurring as single grains, possibly caved material Sm.am. Possibly additives (lignite)
C-5882	3005-3020		100% <u>Sandstone</u> , medium to fine/Sand, medium to very coarse Sm.am. ?Additives; Chalk, white
C-5883	3020-3035		100% <u>Sandstone</u> , medium to fine, well cemented/Sand, medium to very coarse, single grains Sm.am. ?Lignite additives; Pyrite
C-5884	3035-3050		100% <u>Sandstone</u> , very fine to fine, well cemented with silica, calcite and clay cement Sm.am. ?Lignite additive; Pyrite
C-5885	3050-3065		100% <u>Sandstone</u> , very fine to fine, well cemented with silica, calcite and clay cement, some medium to very coarse single grains Sm.am. ?Lignite additive; Coak, black; Pyrite; Chalk, white



Lithology and Total Organic Carbon measurements

TABLE NO.: 2.
WELL NO.: 15/12-5

Sample	Depth (m)	TOC	Lithology
C-5886	3065-3080		100% <u>Sandstone</u> , fine to medium, angular to subangular, well cemented, calcite, silica and clay cement, some medium to coarse, subangular to subrounded grains Sm.am. ?Lignite additive; Claystone, grey
C-5887	3080-3095		100% <u>Sand/Sandstone</u> , mainly as single grains, medium to very coarse, subangular to subrounded, cementing seems to be poor Sm.am. Claystone, grey to dark grey, micromicaeous
C-5888	3095-3110		100% <u>Sand/Sandstone</u> , medium to very coarse, subangular to subrounded, mainly as single grains, cementing seems to be poor. No staining Sm.am. ?Lignite additive; Claystone, grey
C-5890	3125-3140		100% <u>Sand/Sandstone</u> , both medium to very coarse, single grains, (poorly cemented), and well cemented, fine Sm.am. ?Lignite additive
C-5891	3190-3150		<u>Sand/Sandstone</u> , both as medium to very coarse, single grain and as well cemented, fine, angular to subangular Sm.am. ?Lignite additive; Pyrite

Table 3:

DATA FROM ROCK EVAL PYROLYSIS

IKU No.	DEPTH m/ft	S1 (mg/g ROCK)	S2 (mg/g ROCK)	S3 (mg/g ROCK)	TOC (%)	HYDR. INDEX (mg/g TOC)	OXYGEN INDEX (mg/g TOC)	PETROLEUM POTENTIAL S1+S2	PROC. INDEX S1+S2	TEMP. MAX (C)	
C 5837	2846	0.68	5.30	1.29	2.48	214	52	5.98	0.11	436	
		:SWC									
C 5839	2850	0.75	7.23	1.54	2.29	316	67	7.98	0.09	432	
		:SWC									
C 5870	2855	0.11	0.13	1.22	0.41	32	298	0.24	0.46	440	
		:Clst. gy grn									
C 5870	2855	0.23	2.74	1.54	1.92	143	30	2.97	0.08	433	
		:Clst. lt olv gy									
C 5841	2870	0.66	4.48	0.81	2.18	206	37	5.14	0.13	434	
		:SWC									
C 5871	2870	0.19	0.25	1.50	0.51	49	294	0.44	0.43	434	
		:Clst. gy grn									
C 5871	2870	0.59	6.35	0.86	2.45	259	35	6.94	0.09	433	
		:Clst. blk dk gy									
C 5843	2879	0.23	0.44	0.52	1.05	42	50	0.67	0.34	433	
		:SWC									
C 5846	2885	0.40	2.25	0.53	1.47	153	36	2.65	0.15	434	
		:SWC									
C 5872	2885	0.20	1.84	1.71	1.72	107	99	2.04	0.10	431	
		:Clst. lt olv gy									
C 5872	2885	0.79	6.83	1.44	2.77	247	52	7.62	0.10	433	
		:Clst. blk dk gy									
C 5872	2885	0.14	0.08	1.65	0.41	20	402	0.22	0.64	443	
		:Clst. gy grn									
C 5873	2889	0.16	0.13	2.09	0.44	30	475	0.29	0.55	331*	
		:Clst. gy grn									
C 5873	2889	0.32	2.42	2.32	1.83	132	127	2.74	0.12	432	
		:Clst. lt olv gy									
C 5873	2889	0.81	6.18	0.79	2.62	236	30	6.99	0.12	434	
		:Clst. blk dk gy									

* The low Tmax value (331°C) is probably due to a low flat S2 peak.

DATE : 22 - 4 - 87.

Sample	Depth (m)	Composition of residue	Particle size	Preservation palynomorphs	Thermal maturation index	Remarks
C-5870	2840-2855	Am: 80% Lm: 5-10% W: 5-10% C: <5%	F-M-(L)	Poor-Fair	2,5 - 3	Organic residue dominated by degraded amorphous matter. The small liptinitic fraction contains mostly terrestrial palynomorphs.
C-5871	2855-2870	Am: >90% Lm: <5% W: <5% C: <5%	F-M-L	Poor	?	Organic residue totally dominated by fine-particulate amorphous matter and dark granular aggregates. The aggregates contain frequent pyrite framboids.
C-5872	2870-2885 Clst. lt. olv. gy	Am: 80-85% Lm: 5-10% W: 5-10% C: <5%	F-M-L	Fair	2.5-3	Mainly as sample C-5870 above.
C-5872	2870-2885 Clst. blk. dk gy	Am: 80% Lm: 5-10% W: 5-10% C: 5%	F-M-L	Fair	2.5-3?	Mainly as sample C-5872 above, but with a slightly higher proportion of woody and coaly material. Microforams recorded

Abbreviations

Am Amorphous

Al Algal material

W Woody material

F Fine

Lm Liptinitic material

C Coaly fragments

M Medium

L Large

Table 5:

Semiquantitative data from pyrolysis gas chromatography

AREA
 STATOIL 15/12-5

IKU NO	DEPTH (m)	SAMPLE TYPE	C1	C2-C5	C6-C14	C15+
C-5870	2855.00	Cut Clst lt olv gy	695216	1196463	4949633	651465
C-5871	2870.00	Cut Clst blk dk gy	932317	810198	7756713	2690137
C-5872	2885.00	Cut Clst blk dk gy	841703	2506810	7573573	2532992
C-5873	2889.00	Cut Clst blk dk gy	809294	2365808	6623119	1910691
C-5873	2889.00	Cut Clst lt olv gy	620824	1900635	4583342	595184

HEIGHT
 STATOIL 15/12-5

IKU NO	DEPTH (m)	SAMPLE TYPE	C1	C2-C5	C6-C14	C15+
C-5870	2855.00	Cut Clst lt olv gy	3891	627031	1050775	91072
C-5871	2870.00	Cut Clst blk dk gy	6247	319405	1627106	322762
C-5872	2885.00	Cut Clst blk dk gy	5503	1527059	1561222	324279
C-5873	2889.00	Cut Clst blk dk gy	5608	1512648	1272212	257393
C-5873	2889.00	Cut Clst lt olv gy	3710	1070860	1072825	92016

PERCENT
 STATOIL 15/12-5

IKU NO	DEPTH (m)	SAMPLE TYPE	01	02-05	06-014	015+
C-5870	2855.00	Cut Clst lt olv gy	9.28	15.97	66.06	8.69
C-5871	2870.00	Cut Clst blk dk gy	7.65	6.65	63.62	22.07
C-5872	2885.00	Cut Clst blk dk gy	6.24	18.59	56.17	18.79
C-5873	2889.00	Cut Clst blk dk gy	6.75	19.72	55.20	15.92
C-5873	2889.00	Cut Clst lt olv gy	8.06	24.68	59.52	7.73

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TABLE 7b:

CONCENTRATION OF EOM AND CHROMATOGRAPHIC FRACTIONS

(Weight ppm of rock)

MPLC-data

IKU-No	DEPTH (m)	EOM	Sat.	Aro.	HC	Non HC
C 5836	2844.00 SWC	3752	720	968	1689	2064
C 5840	2860.00 SWC	4291	2298	681	2979	1312
C 5844	2880.00 SWC	4517	98	229	327	4190
C 5847	2886.30 SWC	2289	163	235	398	1892
C 5848	2892.60 Core	2379	368	402	769	1610
C 5849	2911.75 Core	1736	257	247	504	1232
C 5850	2917.50 Core	6557	2782	1669	4451	2106
C 5851	2918.70 Core	6706	2709	1584	4293	2414
C 5852	2921.20 Core	9772	3349	1886	5234	4538
C 5853	2922.25 Core	13291	4709	3363	8072	5219
C 5854	2924.40 Core	14787	5587	3435	9022	5765
C 5855	2926.60 Core	16727	6029	4156	10185	6542
C 5856	2927.55 Core	21650	8679	5858	14537	7114
C 5857	2931.50 Core	16582	5255	4132	9387	7195
C 5858	2935.60 Core	16769	7254	4826	12080	4689
C 5859	2937.55 Core	21444	7934	5576	13510	7934
C 5860	2940.20 Core	18319	6601	3896	10497	7822
C 5861	2942.55 Core	20868	7243	4748	11989	6878
C 5862	2943.70 Core	14639	3304	3199	8503	6336
C 5863	2952.65 Core	764	187	55	242	521

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TABLE 7c:

CONCENTRATION OF EOM AND CHROMATOGRAPHIC FRACTIONS

(mg/g TOC)

MPLC-data

IKU-No	DEPTH (m)	EOM	Sat.	Aro.	HC	Non HC
C 5836	2844.00	171.3	32.9	44.2	77.1	94.2
	SWC					
C 5840	2860.00	170.9	91.5	27.1	118.7	52.3
	SWC					
C 5844	2880.00	501.9	10.9	25.4	36.3	465.6
	SWC					
C 5847	2886.30	186.1	13.2	19.1	32.3	153.8
	SWC					
C 5848	2892.60	146.0	22.6	24.6	47.2	98.8
	Core					
C 5849	2911.75	149.7	22.2	21.3	43.5	106.2
	Core					
C 5850	2917.50	799.6	339.2	203.5	542.8	256.9
	Core					
C 5851	2918.70	779.8	315.0	184.1	499.1	280.7
	Core					
C 5852	2921.20	1110.5	380.5	214.3	594.8	515.7
	Core					
C 5853	2922.25	1165.9	413.1	295.0	708.1	457.8
	Core					
C 5854	2924.40	1540.3	582.0	357.8	939.8	600.5
	Core					
C 5855	2926.60	1480.2	533.5	367.8	901.3	578.9
	Core					
C 5856	2927.55	1691.4	678.0	457.7	1135.7	555.7
	Core					
C 5857	2931.50	1480.6	469.2	369.0	838.1	642.4
	Core					
C 5858	2935.60	1612.4	697.5	464.1	1161.6	450.8
	Core					
C 5859	2937.55	1553.9	575.0	404.0	979.0	575.0
	Core					
C 5860	2940.20	1514.0	545.5	322.0	867.5	646.4
	Core					
C 5861	2942.55	1557.3	540.5	354.2	894.7	662.6
	Core					
C 5862	2943.70	1667.3	596.0	359.4	955.4	711.9
	Core					
C 5863	2952.65	381.8	93.5	27.7	121.2	260.6
	Core					

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T A B L E 7d:

COMPOSITION OF MATERIAL EXTRACTED FROM THE ROCK

MPLC-data

I	I	I	I	I	I	I	I	I	I
I	IKU-No	DEPTH	Sat	Aro	HC	SAT	Non HC	HC	I
I			---	---	---	---	---	---	I
I			EOM	EOM	EOM	Aro	EOM	Non HC	I
I		(m)	%	%	%	x 100	%	x 100	I
I									I
I	C 5836	2844.00	19.2	25.8	45.0	74.4	55.0	81.8	I
I		SWC							I
I	C 5840	2860.00	53.6	15.9	69.4	337.5	30.6	227.0	I
I		SWC							I
I	C 5844	2880.00	2.2	5.1	7.2	42.9	92.8	7.8	I
I		SWC							I
I	C 5847	2886.30	7.1	10.3	17.4	69.2	82.6	21.0	I
I		SWC							I
I	C 5848	2892.60	15.5	16.9	32.3	91.5	67.7	47.8	I
I		Core							I
I	C 5849	2911.75	14.8	14.2	29.0	103.9	71.0	40.9	I
I		Core							I
I	C 5850	2917.50	42.4	25.5	67.9	166.7	32.1	211.3	I
I		Core							I
I	C 5851	2918.70	40.4	23.6	64.0	171.1	36.0	177.8	I
I		Core							I
I	C 5852	2921.20	34.3	19.3	53.6	177.6	46.4	115.4	I
I		Core							I
I	C 5853	2922.25	35.4	25.3	60.7	140.0	39.3	154.7	I
I		Core							I
I	C 5854	2924.40	37.8	23.2	61.0	162.7	39.0	156.5	I
I		Core							I
I	C 5855	2926.60	36.0	24.8	60.9	145.1	39.1	155.7	I
I		Core							I
I	C 5856	2927.55	40.1	27.1	67.1	148.1	32.9	204.4	I
I		Core							I
I	C 5857	2931.50	31.7	24.9	56.6	127.2	43.4	130.5	I
I		Core							I
I	C 5858	2935.60	43.3	28.8	72.0	150.3	28.0	257.6	I
I		Core							I
I	C 5859	2937.55	37.0	26.0	63.0	142.3	37.0	170.3	I
I		Core							I
I	C 5860	2940.20	36.0	21.3	57.3	169.4	42.7	134.2	I
I		Core							I
I	C 5861	2942.55	34.7	22.7	57.5	152.6	42.5	135.0	I
I		Core							I
I	C 5862	2943.70	35.7	21.6	57.3	165.8	42.7	134.2	I
I		Core							I
I	C 5863	2952.65	24.5	7.3	31.8	337.9	68.3	46.5	I
I		Core							I

Project no.: 22.1850.00
 Well ident.: 15/12-5
 DATE : 3 - 2 - 87.

T A B L E 8b:

CONCENTRATION OF EOM AND CHROMATOGRAPHIC FRACTIONS

(Weight ppm of rock)

Iatroscan-data

IKU-No	DEPTH (m)	EOM	Sat.	Aro.	RES 1	RES 2
C 5836	2844	3735	241	663	2084	747
C 5840	2860	4250	350	650	2300	975
C 5844	2880	4516	129	452	1935	1968
C 5847	2886.30	2283	63	283	1031	906
C 5848	2892.60	2381	271	336	1559	216
C 5849	2911.75	1736	228	257	902	348
C 5850	2917.50	6486	3081	1324	1351	730
C 5851	2918.70	6667	3107	1427	1307	827
C 5852	2921.20	9832	4420	2168	1647	1571
C 5853	2922.25	13400	5960	3200	1820	2400
C 5854	2924.40	14800	6740	3300	1840	2780
C 5855	2926.60	16842	7605	3316	2500	3421
C 5856	2927.55	21463	8878	4829	3439	4268
C 5857	2931.50	16585	7537	2976	2463	3585
C 5858	2935.60	16875	7375	3828	2625	3031
C 5859	2937.55	21351	9000	4351	6486	4730
C 5860	2940.20	14762	6143	3095	2238	3310
C 5861	2942.55	20645	7355	4129	3000	6161
C 5862	2943.70	14737	5658	2737	2079	4263
C 5863	2952.65	295	125	75	32	63

Project no.: 22.1850.00
 Well ident.: 15/12-5
 DATE : 3 - 2 - 87.

T A B L E 8c:

CONCENTRATION OF EOM AND CHROMATOGRAPHIC FRACTIONS

(mg/g TOC)

Iatroscan-data

IKU-No	DEPTH (m)	EOM	Sat.	Aro.	RES 1	RES 2
C 5836	2844	170.5	11.00	30.26	95.18	34.11
C 5840	2860	169.3	13.94	25.90	91.63	38.84
C 5844	2880	501.8	14.34	50.18	215.05	218.64
C 5847	2886.30	185.6	5.12	23.05	83.86	73.62
C 5848	2892.60	146.1	16.62	20.63	95.63	13.23
C 5849	2911.75	149.6	19.69	22.12	77.76	30.00
C 5850	2917.50	791.0	375.74	161.50	164.80	88.99
C 5851	2918.70	775.2	361.24	165.89	151.94	96.12
C 5852	2921.20	1117.3	502.29	246.37	187.17	178.57
C 5853	2922.25	1175.4	522.81	280.70	159.65	210.53
C 5854	2924.40	1541.7	702.08	343.75	191.67	289.58
C 5855	2926.60	1490.5	673.03	293.43	221.24	302.75
C 5856	2927.55	1676.8	693.60	377.29	268.67	333.46
C 5857	2931.50	1480.8	672.91	265.68	219.95	320.12
C 5858	2935.60	1622.6	709.13	368.09	252.40	291.47
C 5859	2937.55	1547.2	652.17	315.32	470.04	342.73
C 5860	2940.20	1220.0	507.67	255.80	184.97	273.51
C 5861	2942.55	1540.7	548.87	308.14	223.88	459.80
C 5862	2943.70	1655.8	635.72	307.51	233.59	479.01
C 5863	2952.65	147.4	62.50	37.31	15.89	31.25

Project no.: 22.1850.00
 Well ident.: 15/12-5
 DATE : 3 - 2 - 87.

T A B L E 8d:

COMPOSITION OF MATERIAL EXTRACTED FROM THE ROCK

Iatroscan-data

I		:	Sat	Aro	RES 1	SAT	RES 2	RES 1	I
I	IKU-No	DEPTH	---	---	---	---	---	---	I
I		:	EOM	EOM	EOM	Aro	EOM	RES 2	I
I		(m)	%	%	%	x 100	%	x 100	I
I		:							I
I	C 5836	2844	6.45	17.74	55.81	36.36	20.00	279.03	I
I		:							I
I	C 5840	2860	8.24	15.29	54.12	53.85	22.94	235.90	I
I		:							I
I	C 5844	2880	2.86	10.00	42.86	28.57	43.57	98.36	I
I		:							I
I	C 5847	2886.30	2.76	12.41	45.17	22.22	39.66	113.91	I
I		:							I
I	C 5848	2892.60	11.37	14.12	65.45	80.54	9.05	723.04	I
I		:							I
I	C 5849	2911.75	13.16	14.78	51.96	89.00	20.05	259.19	I
I		:							I
I	C 5850	2917.50	47.50	20.42	20.83	232.65	11.25	185.19	I
I		:							I
I	C 5851	2918.70	46.60	21.40	19.60	217.76	12.40	158.06	I
I		:							I
I	C 5852	2921.20	44.96	22.05	16.75	203.88	15.98	104.81	I
I		:							I
I	C 5853	2922.25	44.48	23.88	13.58	186.25	17.91	75.83	I
I		:							I
I	C 5854	2924.40	45.54	22.30	12.43	204.24	18.78	66.19	I
I		:							I
I	C 5855	2926.60	45.16	19.69	14.84	229.37	20.31	73.08	I
I		:							I
I	C 5856	2927.55	41.36	22.50	16.02	183.84	19.89	80.57	I
I		:							I
I	C 5857	2931.50	45.44	17.94	14.85	253.28	21.62	68.71	I
I		:							I
I	C 5858	2935.60	43.70	22.69	15.56	192.65	17.96	86.60	I
I		:							I
I	C 5859	2937.55	42.15	20.38	30.38	206.83	22.15	137.14	I
I		:							I
I	C 5860	2940.20	41.61	20.97	15.16	198.46	22.42	67.63	I
I		:							I
I	C 5861	2942.55	35.62	20.00	14.53	178.12	29.84	48.69	I
I		:							I
I	C 5862	2943.70	38.39	18.57	14.11	206.73	28.93	48.77	I
I		:							I
I	C 5863	2952.65	42.41	25.32	10.78	167.51	21.21	50.84	I
I		:							I

TABLE 9:

TABULATION OF DATA FROM THE GASCHROMATOGRAMS

I I I I	IKU No.	DEPTH (m)	PRISTANE	PRISTANE	PHYTANE	A	n-C17	CP1	I
			PHYTANE	A = n-C17	B = n-C18	B	n-C27		
I		:							I
I	C 5836	2844	2.4	2.3	1.2	1.9	2.4	1.1	I
I		SWC							I
I	C 5840	2860	2.5	1.8	0.9	2.0	1.7	1.1	I
I		SWC							I
I	C 5844	2880	3.1	2.1	0.8	2.7	0.9	1.1	I
I		SWC							I
I	C 5847	2886.30	3.3	2.5	0.8	3.0	0.6	1.1	I
I		SWC							I
I	C 5848	2892.60	3.0	1.5	0.9	1.8	2.5	1.3	I
I		Core							I
I	C 5849	2911.75	3.5	1.1	0.8	1.3	14.5	1.4	I
I		Core							I
I	C 5850	2917.50	1.5	0.7	0.6	1.3	3.5	1.0	I
I		Core							I
I	C 5851	2918.70	1.2	0.8	0.7	1.2	2.4	1.0	I
I		Core							I
I	C 5852	2921.20	1.4	0.7	0.6	1.3	5.9	1.1	I
I		Core							I
I	C 5853	2922.25	1.2	0.7	0.6	1.2	2.4	1.1	I
I		Core							I
I	C 5854	2924.40	1.4	0.8	0.6	1.3	2.2	1.0	I
I		Core							I
I	C 5855	2926.60	1.2	0.8	0.7	1.2	2.7	1.0	I
I		Core							I
I	C 5856	2927.55	1.4	0.7	0.6	1.3	2.6	1.1	I
I		Core							I
I	C 5857	2931.50	1.4	0.7	0.6	1.3	2.5	1.0	I
I		Core							I
I	C 5858	2935.60	1.4	0.7	0.5	1.3	3.0	1.0	I
I		Core							I
I	C 5859	2937.55	1.3	0.7	0.6	1.2	2.3	1.0	I
I		Core							I
I	C 5860	2940.20	1.3	0.8	0.6	1.3	2.4	1.1	I
I		Core							I
I	C 5861	2942.55	1.3	0.8	0.6	1.2	2.5	1.0	I
I		Core							I
I	C 5862	2943.70	1.3	0.8	0.6	1.3	1.7	1.0	I
I		Core							I
I	C 5863	2952.65	3.0	1.0	0.9	1.2	6.4	1.0	I
I		Core							I

DATE : 16 - 2 - 87.

TABLE 10:

AROMATIC HYDROCARBON RATIOS.

I	:	Depth	:					I
I	:	IKU No.	:	MNR	DMNR	MPI 1	MPI 2	I
I	:	(m)	:					I
I	:		:					I
I	:	C-5836	:	0.96	2.0	0.65	0.51	I
I	:	SWC	:					I
I	:	C-5840	:	0.87	1.9	0.63	0.53	I
I	:	SWC	:					I
I	:	C-5844	:	0.72	1.9	0.59	0.54	I
I	:	SWC	:					I
I	:	C-5848	:	1.16	2.3	0.58	0.55	I
I	:	Core	:					I
I	:	C-5849	:	1.19	2.4	0.54*	0.48*	I
I	:	Core	:					I
I	:	C-5852	:	1.12	2.0	0.65	0.67	I
I	:	Core	:					I
I	:	C-5855	:	1.08	2.1	0.82	0.80	I
I	:	Core	:					I
I	:	C-5858	:	0.98	2.0	0.85	0.83	I
I	:	Core	:					I
I	:	C-5861	:	0.91	1.8	0.82	0.84	I
I	:	Core	:					I
I	:	C-5863	:	1.00	2.4	0.60*	0.55*	I
I	:	Core	:					I

* Erroneous ratio, because of small elution pattern.

MNR = methylnaphthalene ratio $2\text{-MN}/1\text{-MN}$
 DMNR = dimethylnaphthalene ratio $2,6\text{-DMN}+2,7\text{-DMN}/1,5\text{-DMN}$
 MPI 1 = methylphenanthrene index 1 $1.5*(3\text{-MP}+2\text{-MP})/(P+9\text{-MP}+1\text{-MP})$
 MPI 2 = methylphenanthrene index 2 $3*(2\text{-MP})/(P+9\text{-MP}+1\text{-MP})$

Table 11a: Isotope data.

IKU no.	Depth (m)	Extract	SAT	ARO	NSO	Asphaltenes
C-5836 SWC	2844.00	-29.5	-29.4*	-29.6	-29.2	-28.7
C-5840 SWC	2860.00	-28.5	-28.3*	-28.6	-28.3	-28.1
C-5844 SWC	2880.00	-28.8	-28.2*	-28.7	-28.6	-28.5
C-5848 Core	2892.60	-28.7	-27.4*	-28.2	-29.1	-28.2
C-5849 Core	2911.75	-28.6	-29.4	-27.9	-28.3	-28.0
C-5852 Core	2921.20	-28.8	-29.1	-28.3	-28.7	-28.9
C-5855 Core	2926.60	-28.8	-29.1	-28.3	-28.2	-28.9
C-5858 Core	2935.60	-28.7	-29.1	-28.2	-28.3	-29.0
C-5861 Core	2942.55	-28.7	-29.1	-28.2	-28.5	-28.9
C-5863 Core	2952.65	-28.7	-29.3	-27.8	-28.6	-28.4

* The value could be too heavy because of a small sample amount.

Table 11b: Isotope data of kerogen concentrates.

IKU no.	Depth (m)	Kerogen concentrates
C-5870 Clst.lt.olv.gy	2840-2855	-26.0
C-5871 Clst.blk.dk.gy.	2855-2870	-27.3
C-5872 Clst.blk.dk.gy.	2870-2885	-27.9
C-5872 Clst.lt.olv.gy.	2870-2885	*

* Too little sample material (kerogen concentrate).

Table 12: Molecular ratios from sterane and terpane mass chromatograms.
Maturity ratios.

IKU code	Depth (m)	$\alpha\beta/\alpha\beta+\beta\alpha$ ¹⁾	%22S ²⁾	% $\beta\beta$ ³⁾	%20S ⁴⁾
C-5836 SWC	2844.00	0.85	63.6	55.8	39.6
C-5840 SWC	2860.00	0.80	62.0	55.5	37.8
C-5844 SWC	2880.00	0.85	57.8	51.5	42.0
C-5848 Core	2892.60	0.86	63.5	57.8	38.9
C-5849 Core	2911.75	0.82	55.4	68.5	37.0
C-5852 Core	2921.20	0.92	65.9	76.1	55.1
C-5855 Core	2926.60	-	58.8	77.3	41.3
C-5858 Core	2935.60	0.92	58.5	73.5	43.2
C-5861 core	2942.55	-	59.9	75.5	47.2
C-5863 Core	2952.65	0.92	59.9	67.6	41.7

1) E/E+F in m/z 191

2) Average % distribution between first and second eluting isomers of extended hopanes (G-M in m/z 191)

3) $2(r+s)/(q+t+2(r+s))$ in m/z 217

4) $q/q+t$ in m/z 217

167/zz/an1/12

Table 13: Molecular ratios from terpane and sterane mass chromatograms.
Maturity and source characteristic ratios.

IKU no.	Depth (m)	Q/E ¹⁾	T _m /T _s ²⁾	X/E ³⁾	Z/E ⁴⁾	a/a+j ⁵⁾
C-5836 SWC	2844.00	0.06	5.40	0.07	-	0.74
C-5840 SWC	2860.00	0.09	-	-	-	0.75
C-5844 SWC	2880.00	0.07	7.40	0.04	-	0.75
C-5848 Core	2892.60	0.05	3.47	0.07	0.04	0.77
C-5849 Core	2911.75	0.08	-	0.08	-	0.75
C-5852 Core	2921.20	0.11	-	-	-	0.85
C-5855 Core	2926.60	0.09	0.50	-	-	0.82
C-5858 Core	2935.60	0.08	0.63	0.06	-	0.83
C-5861 Core	2942.55	0.09	0.55	-	-	0.84
C-5863 Core	2952.65	0.08	1.06	-	-	0.76

1) Relative abundance of tricyclic terpanes (Q/E in m/z 191)

2) B/A in m/z 191

3) Relative abundance of unknown (X/E in m/z 191)

4) Relative abundance of bisnorhopane (Z/E in m/z 191)

5) Relative abundance of C₂₇ rearranged steranes (a/a+j in m/z 217)

Table 14a:

Peak heights and areas from
m/z 191 mass chromatograms

Sample identity: C-5836, 2844.00 m (swc)

Peak heights and peak areas from m/z 191.

Peak identities	Peak heights	m/z 191	Peak areas
A	7.8		12.5
B	90.1		388.9
C	187.4		1116.2
X	15.2		33.1
D	23.7		86.5
E	285.4		1522.5
F	42.8		187.7
G	103.9		578.5
H	68.1		377.7
I	20.8		73.0
J ₁	43.8		226.9
J ₂	30.5		121.8
K ₁	30.8		161.0
K ₂	14.0		85.2
P	50.2		247.8
Q	18.4		83.5
R	14.4		110.7
S	21.3		98.6
T	5.1		27.5

Sample identity: C-5840, 2860.00 m (swc)

Peak heights and peak areas from m/z 191.

Peak identities	Peak heights	m/z 191	Peak areas
B	32.6		123.3
C	61.6		357.0
E	88.8		440.6
F	20.1		66.4
G	40.3		189.9
H	27.3		110.6
I	8.9		19.2
J ₁	13.3		65.3
J ₂	6.5		30.1
P	24.9		105.2
Q	10.1		36.4
R	5.9		35.0
S	11.7		51.2

Sample identity: C-5844, 2880.00 m (swc)

Peak heights and peak areas from m/z 191.

Peak identities	Peak heights	Peak areas
A	30.1	104.8
B	302.8	1354.1
C	465.7	3139.0
D	66.3	300.5
E	725.1	4092.9
F	124.0	567.3
G	278.8	1704.3
H	185.0	1090.1
I	52.1	289.8
J ₁	108.3	756.5
J ₂	82.9	512.4
K ₁	56.7	357.4
K ₂	30.8	134.0
L ₁	19.2	114.6
L ₂	12.0	62.7
P	122.6	573.9
Q	49.9	233.2
R	31.5	236.7
S	67.5	318.4
T	12.2	78.5

Sample identity: C-5848, 2892.60 m (Core)

Peak heights and peak areas from m/z 191.

Peak identities	Peak heights	m/z 191	Peak areas
A	44.5		151.0
B	210.1		911.1
Z	14.3		46.1
C	390.2		2725.9
X	37.0		105.8
D	53.1		193.4
E	752.5		4150.1
F	125.4		599.0
G	274.1		1748.6
H	197.4		1089.8
I	59.6		356.6
J ₁	128.1		794.0
J ₂	75.3		493.4
K ₁	64.2		234.2
K ₂	28.7		218.9
L ₁	33.5		204.3
L ₂	12.0		51.7
P	87.8		406.2
Q	30.6		142.1
R	19.6		145.3
S	39.8		164.4
T	7.9		54.4

Sample identity: C-5849, 2911.75 m (Core)

Peak heights and peak areas from m/z 191.

Peak identities	Peak heights	m/z 191	Peak areas
B	79.7		337.2
C	111.2		576.6
D	18.6		66.9
E	156.9		763.9
F	34.7		154.9
G	62.7		335.0
H	53.9		228.4
I	12.5		69.2
J ₁	21.0		119.0
J ₂	14.1		55.0
P	26.9		130.3
Q	10.0		36.3
R	7.1		49.5
S	11.9		58.6

Sample identity: C-5852, 2921.20 m (Core)

Peak heights and peak areas from m/z 191.

Peak identities	Peak heights	m/z 191	Peak areas
A	9.5		28.1
C	45.9		302.4
E	96.8		550.3
G	37.6		183.9
H	21.0		110.2
J ₁	25.8		146.3
J ₂	16.2		62.4
K ₁	19.9		62.7
K ₂	7.1		10.2
P	14.1		53.0
Q	10.8		48.6
R	5.7		25.3
S	7.6		25.2

Sample identity: C-5855, 2926.60 m (Core)

Peak heights and peak areas from m/z 191.

Peak identities	Peak heights	Peak areas
A	26.5	96.9
B	12.9	26.3
C	86.2	655.3
E	210.2	1198.1
G	72.1	377.6
H	45.4	258.3
J ₁	46.0	270.1
J ₂	25.2	124.6
K ₁	30.6	156.4
K ₂	18.6	103.7
L ₁	12.5	38.1
L ₂	8.7	33.2
P	27.3	118.8
Q	19.2	91.3
R	10.9	62.7
S	15.8	53.3

Sample identity: C-5858, 2935.60 m (Core)

Peak heights and peak areas from m/z 191.

Peak identities	Peak heights	m/z 191	Peak areas
A	65.1		256.4
B	34.4		146.5
C	165.3		1334.0
E	359.9		2147.5
F	19.5		63.9
G	112.8		695.8
H	77.2		399.4
J ₁	72.5		448.7
J ₂	46.6		279.0
K ₁	35.2		227.0
K ₂	26.7		133.5
P	37.2		160.5
Q	29.4		130.5
R	14.2		92.4
S	23.5		99.8
T	9.9		61.6

Sample identity: C-5861, 2942.55 m (Core)

Peak heights and peak areas from m/z 191.

Peak identities	Peak heights	m/z 191	Peak areas
A	25.3		92.6
B	12.7		40.0
C	70.6		536.4
E	181.0		374.6
G	55.8		317.4
H	35.7		177.5
J ₁	33.3		232.2
J ₂	22.3		110.8
K ₁	16.1		102.5
K ₂	10.3		25.9
P	18.2		71.5
Q	17.9		80.3
R	7.5		26.3
S	8.4		34.3

Sample identity: C-5863, 2952.65 m (Core)

Peak heights and peak areas from m/z 191.

Peak identities	Peak heights	m/z 191	Peak areas
A	17.8		69.8
B	18.2		80.9
C	73.1		459.6
E	126.7		681.6
F	9.4		35.6
G	33.1		160.2
H	26.7		118.0
J ₁	14.9		65.3
J ₂	11.0		31.3
P	23.0		103.7
Q	11.3		50.7
R	6.5		46.4
S	8.2		34.1

Table 14b:

Peak heights and areas from
m/z 217 and 218 mass chromatograms

Sample identity: C-5836, 2844.00 m (swc)

Peak heights and peak areas from m/z 217 and 218 mass chromatograms.

Peak identities	m/z 217		m/z 218	
	Peak heights	Peak areas	Peak heights	Peak areas
a	92.8	492.6		
b	55.2	246.7		
c	19.2	83.0		
f	22.1	272.9		
g	27.0	255.1		
h	141.3	889.3	36.3	235.6
i	-	-	16.5	98.6
j	33.0	161.2		
k	72.7	465.5		
n	30.6	224.4	15.5	88.9
o	12.9	79.8	12.0	79.0
p	14.9	94.8		
q	36.4	197.9		
r	30.4	174.0	28.0	129.1
s	15.9	57.9	18.7	139.1
t	48.1	321.4		
u	36.8	222.3		
v	12.9	58.7		

Sample identity: C-5840, 2860.00 m (swc)

Peak heights and peak areas from m/z 217 and 218 mass chromatograms.

Peak identities	m/z 217		m/z 218	
	Peak heights	Peak areas	Peak heights	Peak areas
a	31.2	150.7		
b	16.6	81.1		
c	6.6	31.1		
f	8.9	89.5		
g	10.3	75.1		
h	44.8	272.8	12.3	63.8
i	-	-	4.3	27.6
j	10.5	55.5		
k	17.8	123.2		
n	8.4	66.5	4.0	25.3
o	4.2	26.2	4.1	21.5
p	3.5	19.7		
q	10.1	49.8		
r	9.7	44.3	8.0	35.2
s	4.6	20.9	5.0	23.9
t	16.8	105.9		
u	15.1	85.5		
v	5.4	25.4		

Sample identity: C-5844, 2880.00.m (swc)

Peak heights and peak areas from m/z 217 and 218 mass chromatograms.

Peak identities	m/z 217		m/z 218	
	Peak heights	Peak areas	Peak heights	Peak areas
a	182.2	904.4		
b	109.1	496.4		
c	37.9	161.9		
f	38.3	502.5		
g	45.9	354.4		
h	284.0	1839.8	63.8	453.8
i	-	-	42.0	229.5
j	59.8	288.9		
k	111.8	847.5		
n	42.0	445.6	27.0	197.0
o	31.8	185.4	27.1	156.6
p	18.4	112.2		
q	59.2	302.4		
r	41.0	261.2	37.5	187.6
s	26.0	138.9	31.6	242.2
t	90.7	563.5		
u	95.7	512.9		
v	26.7	133.5		

Sample identity: C-5848, 2892.60 m (Core)

Peak heights and peak areas from m/z 217 and 218 mass chromatograms.

Peak identities	m/z 217		m/z 218	
	Peak heights	Peak areas	Peak heights	Peak areas
a	254.9	1325.8		
b	159.6	715.9		
c	46.5	195.5		
f	56.1	726.7		
g	72.5	628.5		
h	308.3	1836.4	72.7	485.0
i	-	-	43.7	242.8
j	73.8	419.5		
k	170.5	1106.6		
n	62.3	592.6	34.5	215.3
o	42.8	296.1	36.6	188.0
p	38.4	205.9		
q	66.9	426.5		
r	62.0	370.1	49.0	257.3
s	44.9	203.8	41.9	306.9
t	111.3	696.6		
u	64.3	353.7		
v	22.8	128.3		

Sample identity: C-5849, 2911.75 m (Core)

Peak heights and peak areas from m/z 217 and 218 mass chromatograms.

Peak identities	m/z 217		m/z 218	
	Peak heights	Peak areas	Peak heights	Peak areas
a	16.2	82.5		
b	10.9	48.9		
c	4.2	13.6		
f	4.3	48.5		
g	5.4	48.4		
h	22.0	140.1	7.0	44.2
i	-	-	3.8	25.1
j	5.7	20.7		
k	9.5	78.4		
n	4.3	39.5	3.5	16.0
o	2.5	13.5	2.5	10.4
p	2.2	8.9		
q	4.7	27.4		
r	7.7	50.7	4.4	24.7
s	4.2	17.7	3.4	22.4
t	7.7	51.9		
u	11.0	73.7		
v	3.9	16.8		

Sample identity: C-5852, 2921.20 m (Core)

Peak heights and peak areas from m/z 217 and 218 mass chromatograms.

Peak identities	m/z 217		m/z 218	
	Peak heights	Peak areas	Peak heights	Peak areas
a	43.1	224.6		
b	23.4	124.6		
c	5.7	21.1		
f	10.3	123.6		
g	8.8	75.9		
h	45.4	280.2	19.7	128.6
i	-	-	17.7	98.4
j	8.0	38.2		
k	18.4	136.9		
n	17.0	138.3	11.6	70.0
o	17.9	97.9	15.4	72.6
p	4.4	26.3		
q	6.7	55.1		
r	16.6	103.6	12.9	67.3
s	15.8	97.0	12.8	91.7
t	10.8	78.7		
u	19.4	108.0		
v	11.4	63.2		

Sample identity: C-5855, 2926.60 m (Core)

Peak heights and peak areas from m/z 217 and 218 mass chromatograms.

Peak identities	m/z 217		m/z 218	
	Peak heights	Peak areas	Peak heights	Peak areas
a	60.2	309.3		
b	33.0	174.0		
c	7.4	30.4		
f	16.0	147.8		
g	12.6	123.5		
h	64.5	394.1	29.9	159.7
i	-	-	25.1	133.0
j	13.5	57.6		
k	27.1	208.2		
n	24.6	185.7	20.9	122.8
o	20.3	148.3	20.3	126.9
p	6.2	27.5		
q	9.5	64.2		
r	22.7	143.0	18.7	96.6
s	17.9	117.4	18.1	128.8
t	13.8	116.0		
u	41.5	214.7		
v	22.7	108.9		

Sample identity: C-5858, 2935.60 m (Core)

Peak heights and peak areas from m/z 217 and 218 mass chromatograms.

Peak identities	m/z 217		m/z 218	
	Peak heights	Peak areas	Peak heights	Peak areas
a	72.3	366.0		
b	39.5	204.9		
c	9.4	33.3		
f	18.6	183.7		
g	16.7	143.1		
h	69.8	458.7	35.8	222.5
i	-	-	30.0	186.3
j	15.9	82.4		
k	32.9	298.8		
n	33.7	212.4	28.7	154.2
o	29.1	170.9	27.7	164.2
p	9.5	50.7		
q	16.8	126.5		
r	30.1	186.3	24.7	140.4
s	26.8	183.3	25.5	178.9
t	19.6	145.9		
u	46.7	262.1		
v	23.9	121.4		

Sample identity: C-5861, 2942.55 m (Core)

Peak heights and peak areas from m/z 217 and 218 mass chromatograms.

Peak identities	m/z 217		m/z 218	
	Peak heights	Peak areas	Peak heights	Peak areas
a	35.6	180.1		
b	21.6	107.5		
c	5.0	21.9		
f	9.5	111.1		
g	8.8	85.5		
h	41.1	245.9	17.9	117.8
i	-	-	16.0	93.7
j	5.6	32.6		
k	17.6	119.1		
n	17.1	96.6	13.2	79.5
o	17.1	83.3	14.4	86.2
p	4.1	20.4		
q	8.5	61.5		
r	14.7	84.1	13.9	65.9
s	12.8	85.5	14.8	98.5
t	9.1	65.5		
u	24.6	144.0		
v	12.6	55.7		

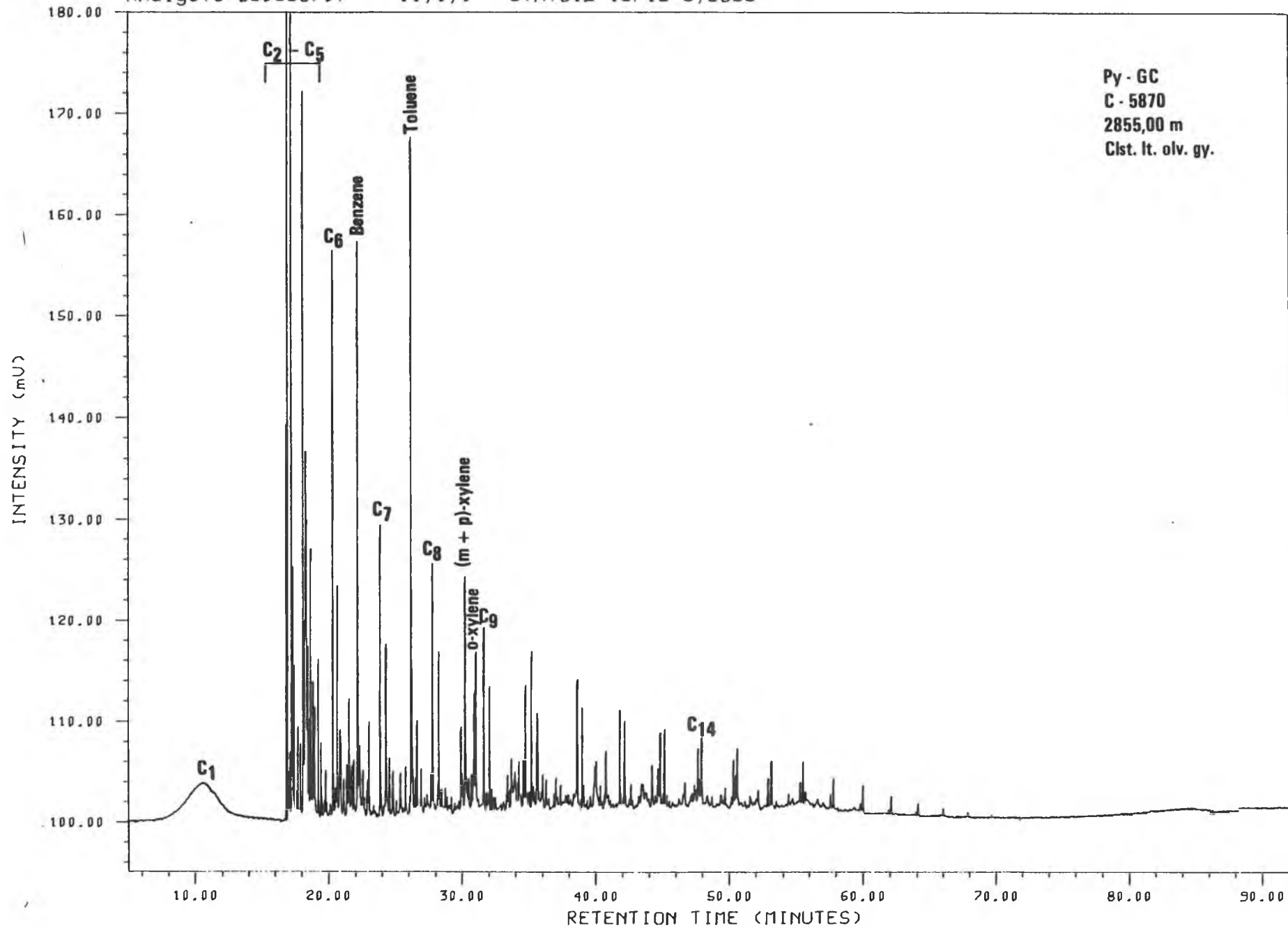
Sample identity: C-5863, 2962.65 m (Core)

Peak heights and peak areas from m/z 217 and 218 mass chromatograms.

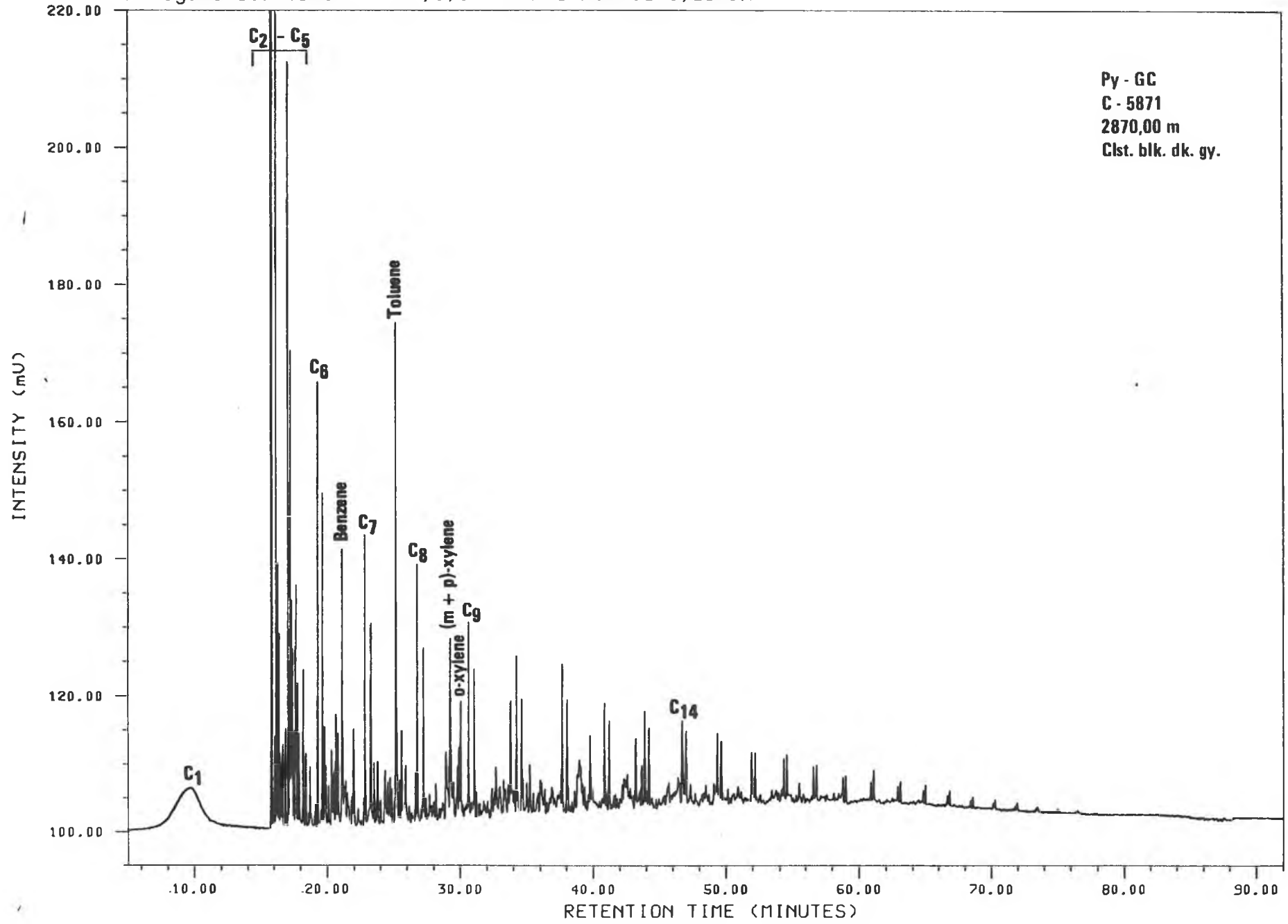
Peak identities	m/z 217		m/z 218	
	Peak heights	Peak areas	Peak heights	Peak areas
a	14.9	66.5		
b	8.7	41.4		
c	1.1	4.6		
f	4.3	43.5		
g	4.1	23.3		
h	18.7	115.1	9.0	57.1
i	-	-	7.1	37.4
j	4.9	15.5		
k	8.1	50.5		
n	7.8	53.6	6.0	34.2
o	7.3	42.4	4.8	27.2
p	2.3	9.1		
q	4.2	19.2		
r	4.4	25.1	4.7	22.8
s	4.1	25.1	4.0	30.2
t	5.2	28.4		
u	13.4	63.5		
v	5.7	30.6		

FIGURE 1
PYROLYSIS GAS CHROMATOGRAMS

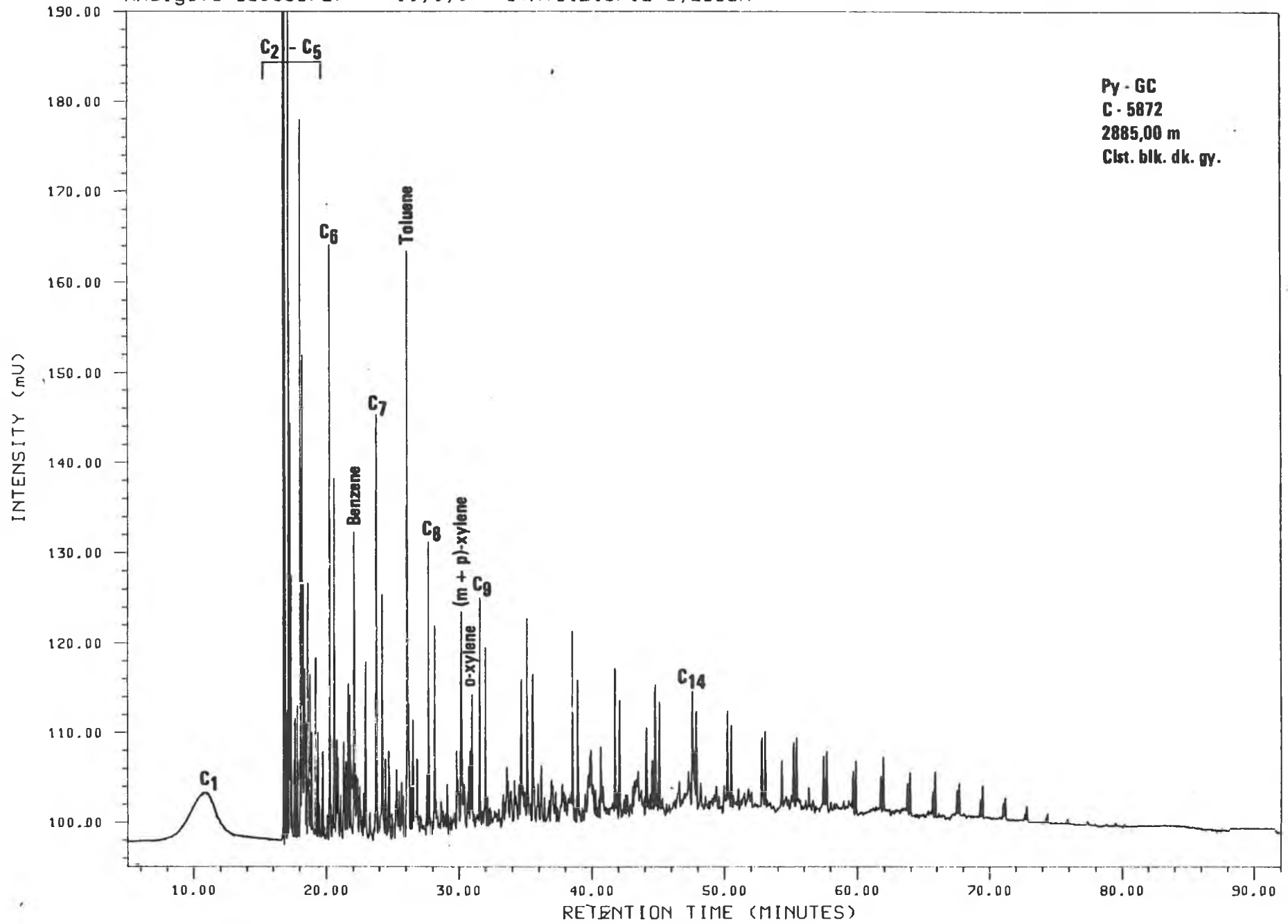
Analysis 850C5870P 11,1,1 STATOIL 15/12-5,2855



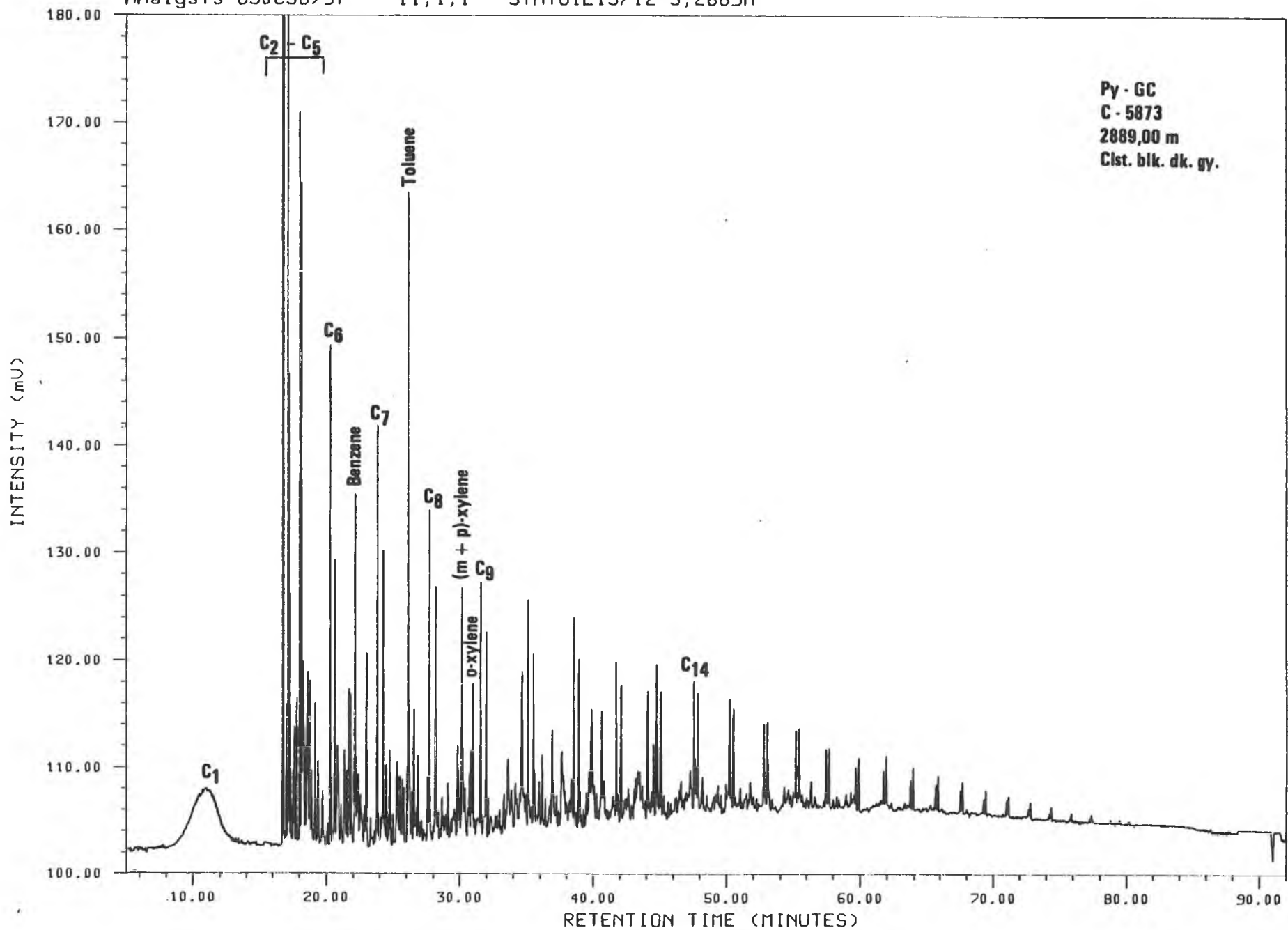
Py - GC
C - 5870
2855,00 m
Clst. lt. olv. gy.



Py - GC
C - 5871
2870,00 m
Clst. blk. dk. gy.



Py - GC
C - 5872
2885,00 m
Clst. blk. dk. gy.

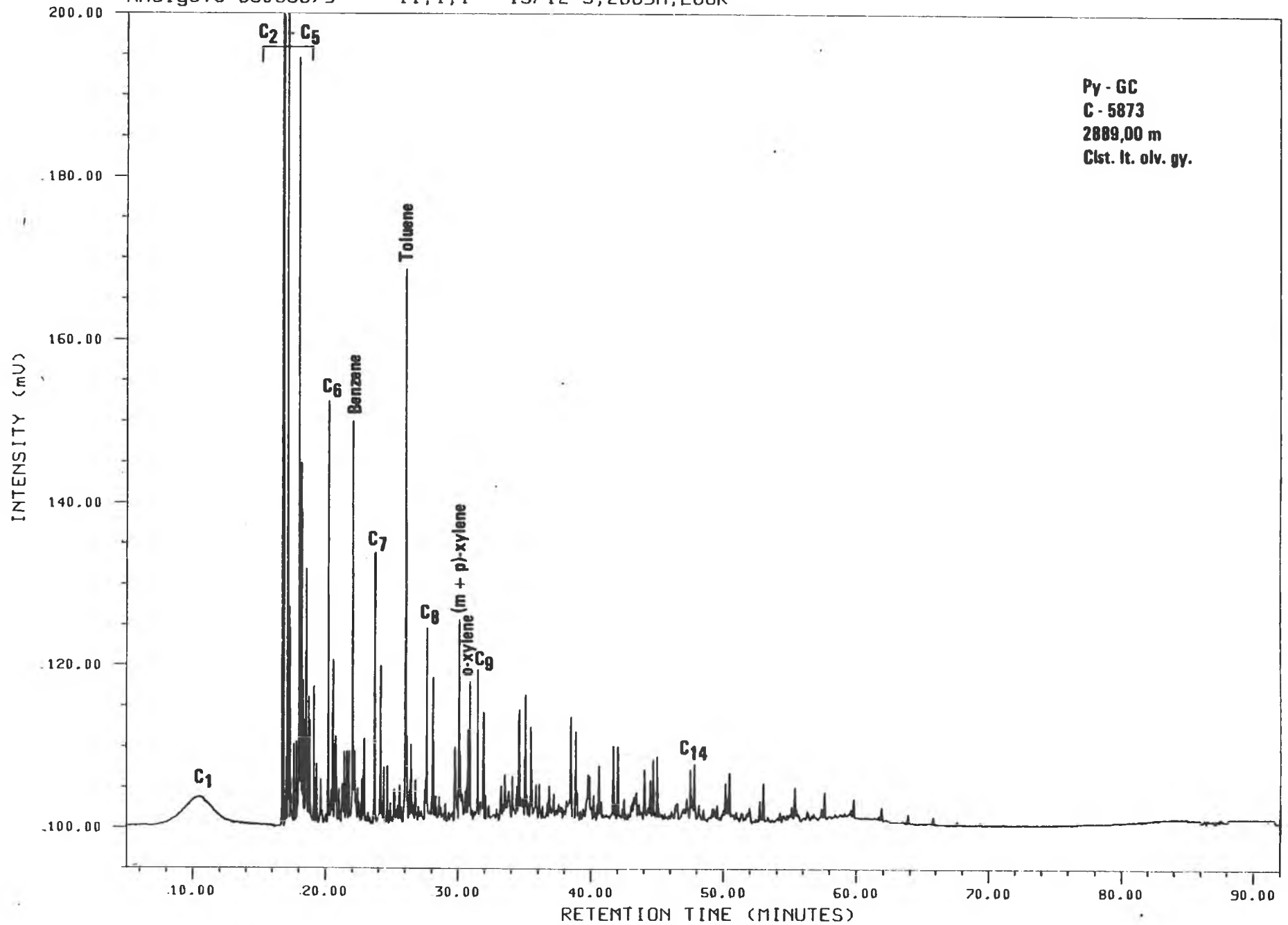


Py - GC
C - 5873
2889,00 m
Clst. blk. dk. gy.

Analysis 850C5873

11,1,1

15/12-5,2889M,LOGR



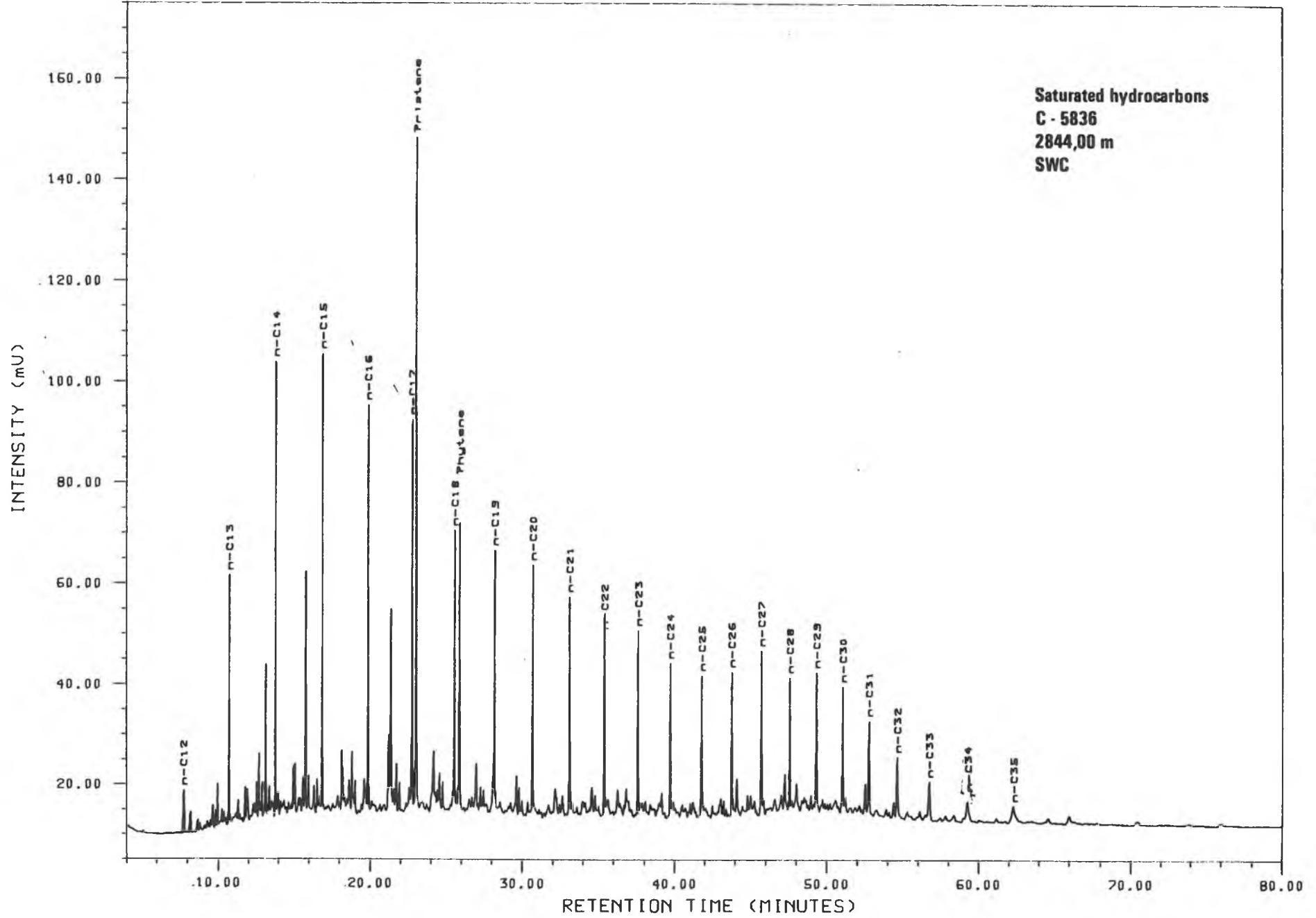
Py - GC
C - 5873
2889,00 m
Clst. lt. olv. gy.

FIGURE 2

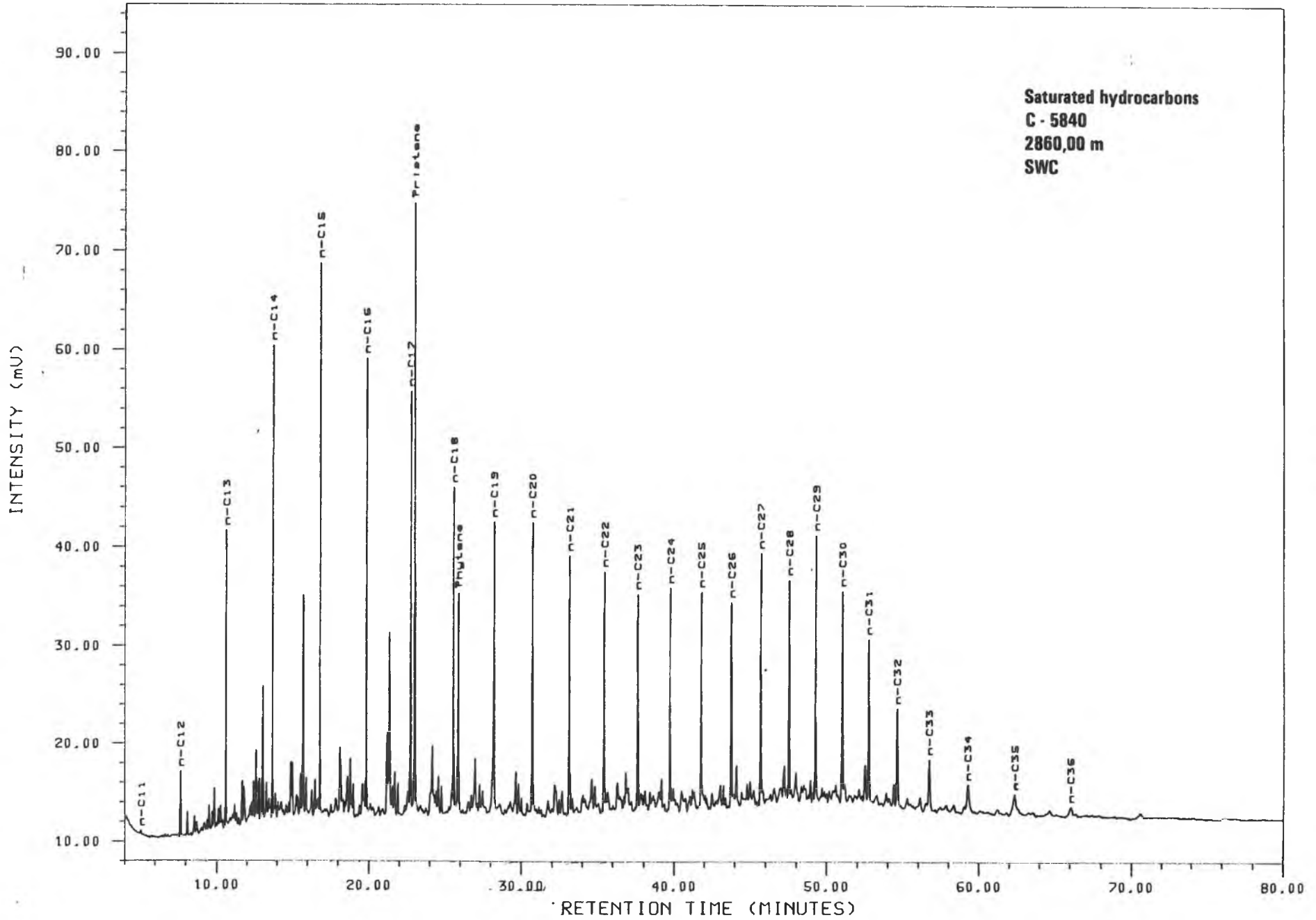
SATURATED HYDROCARBON GAS CHROMATOGRAMS

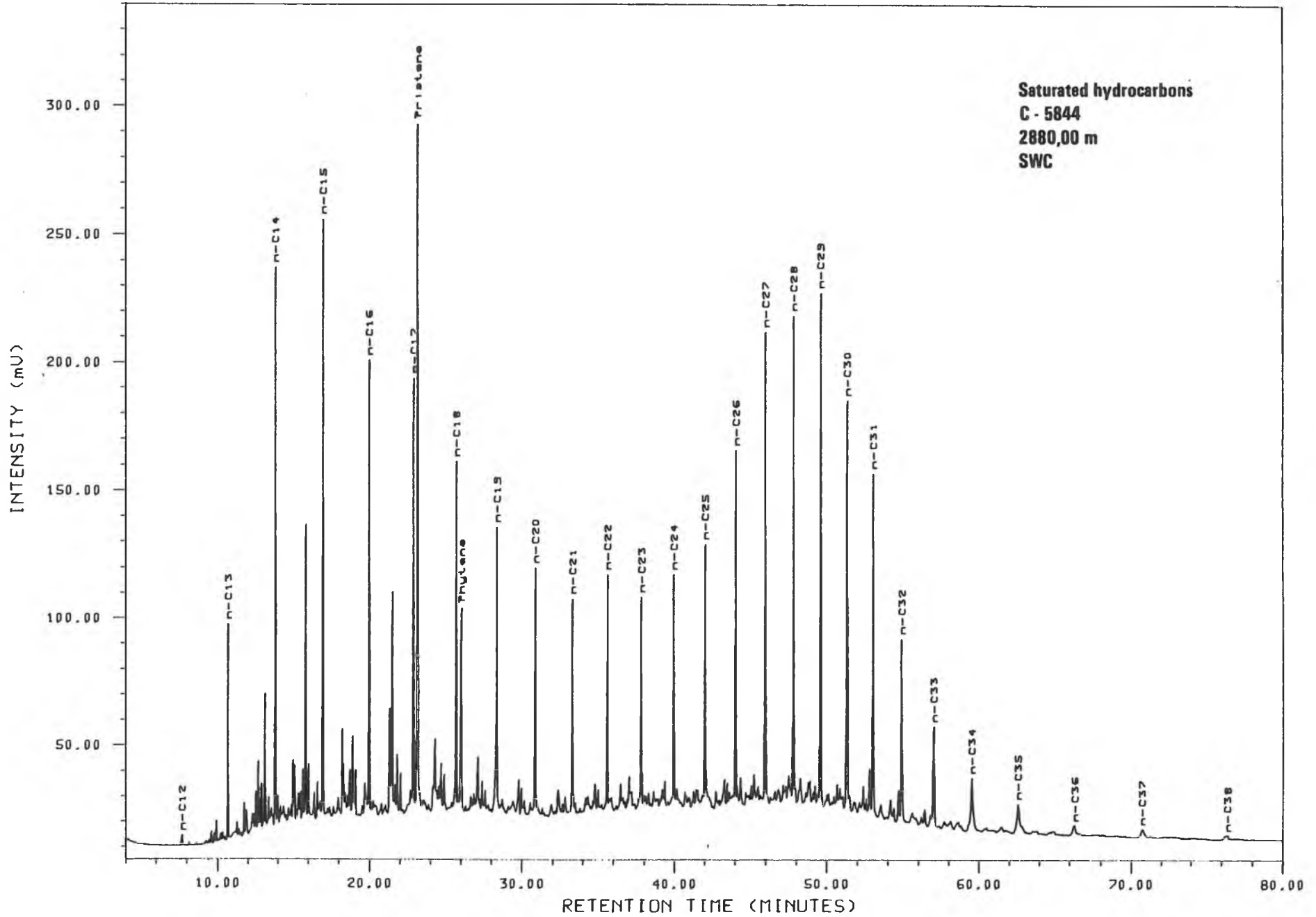
n-C₁₅ etc. - n-alkanes
Pr - pristane
Ph - phytane
* - other acyclic
isoprenoids

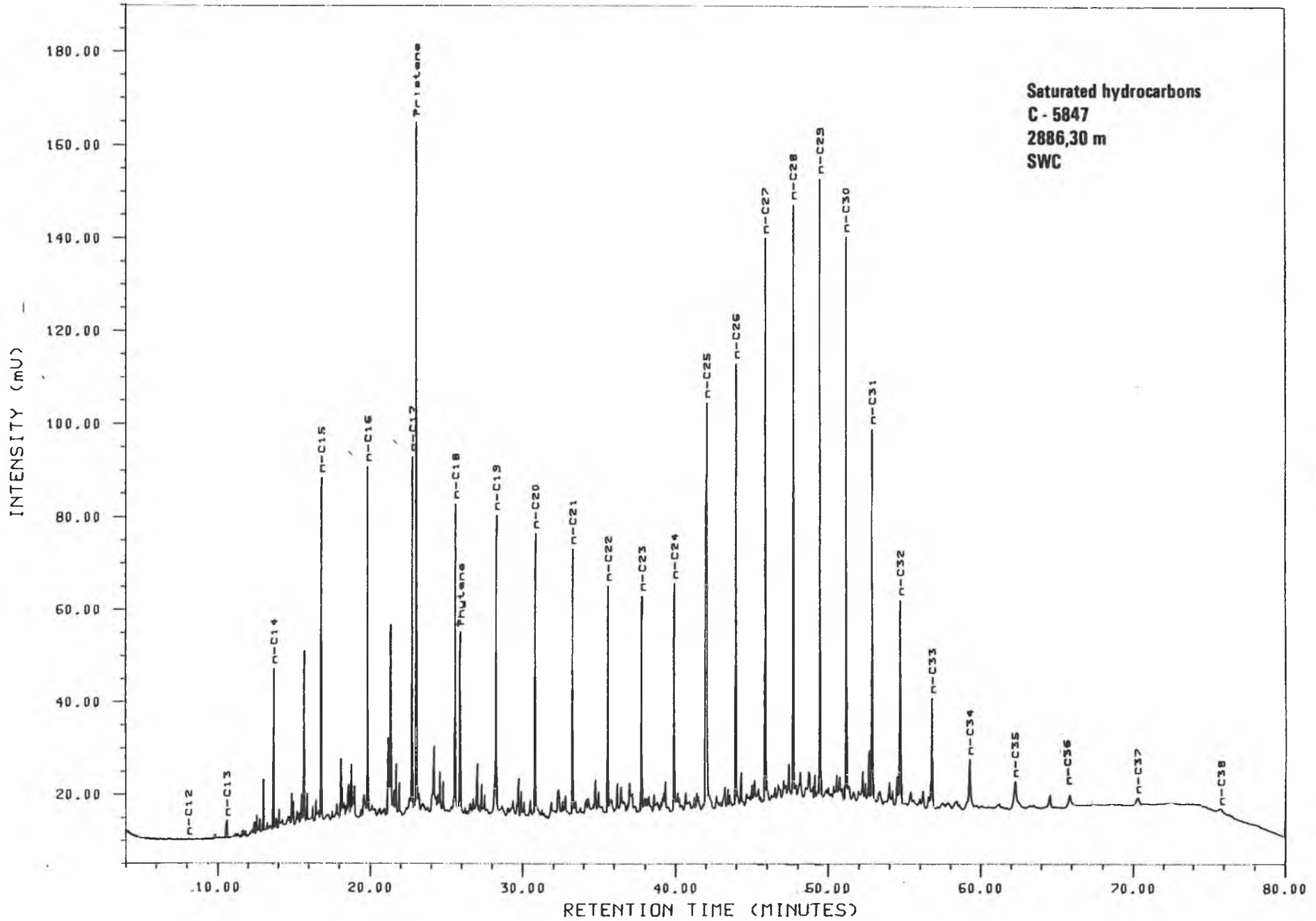
Analysis 850C58365 4,1,1 C-5836,15/12-5,SAT.

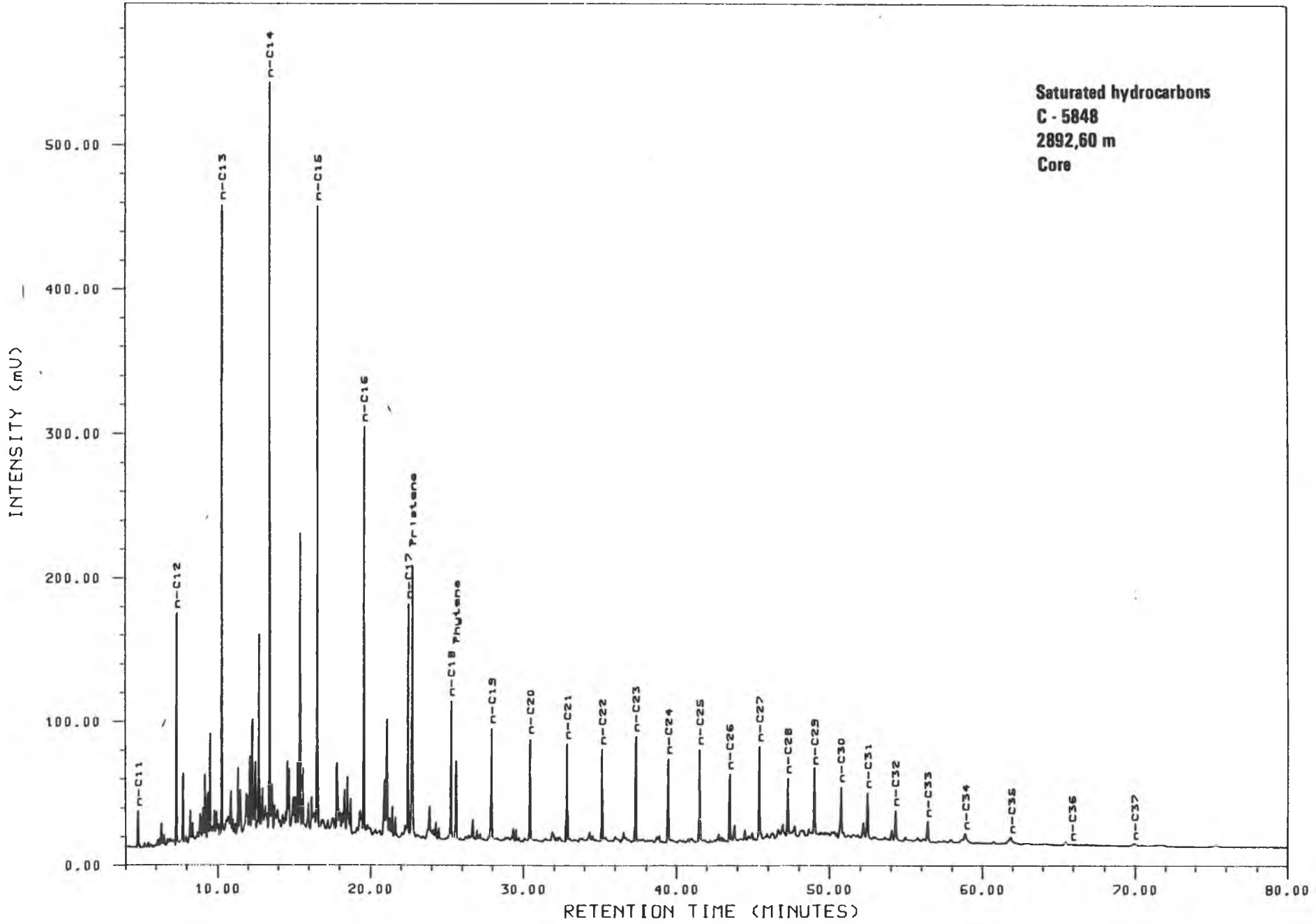


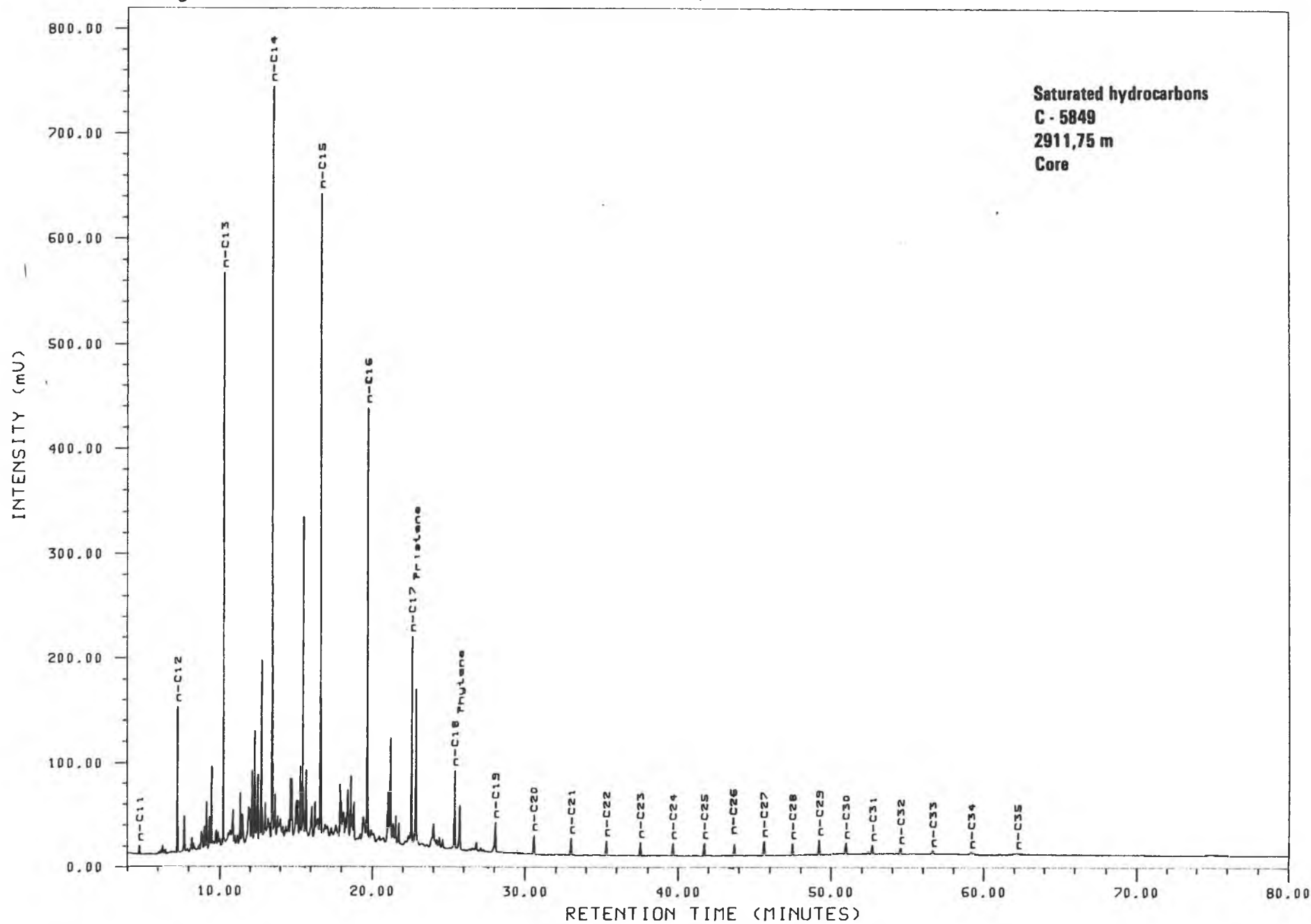
Analysis 850C5840S 4,1,1 C-5840,15/12-5,SAT.

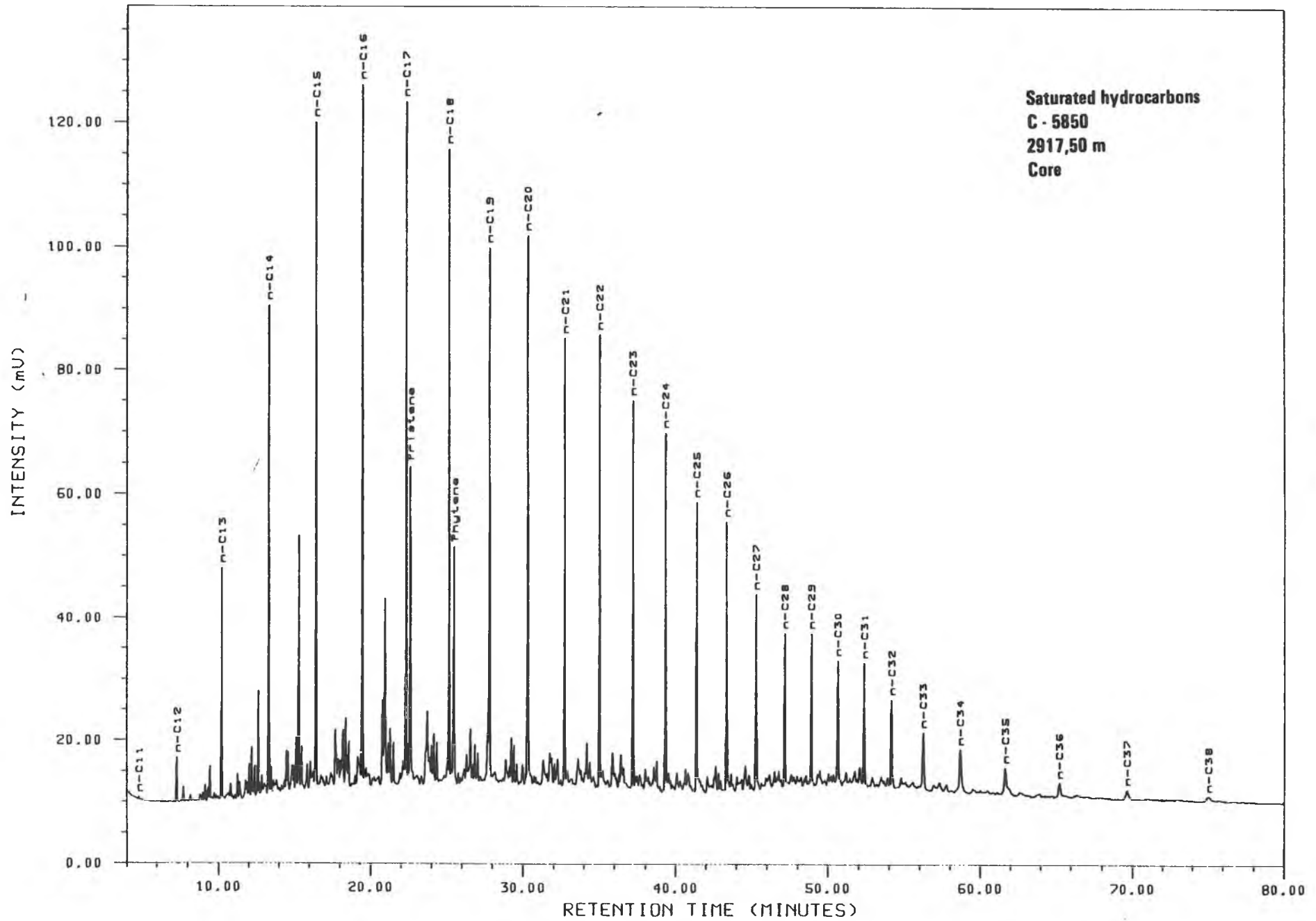


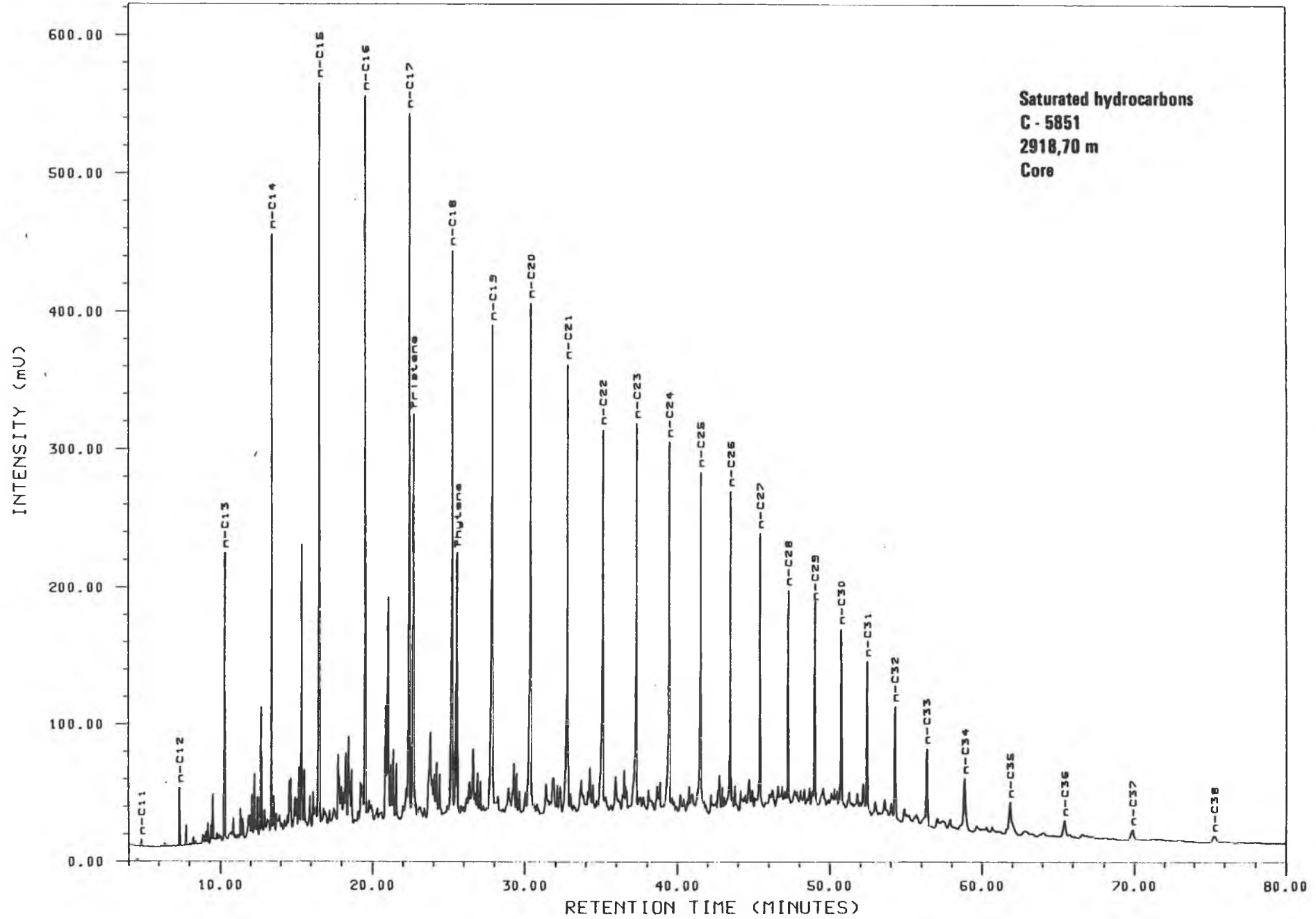


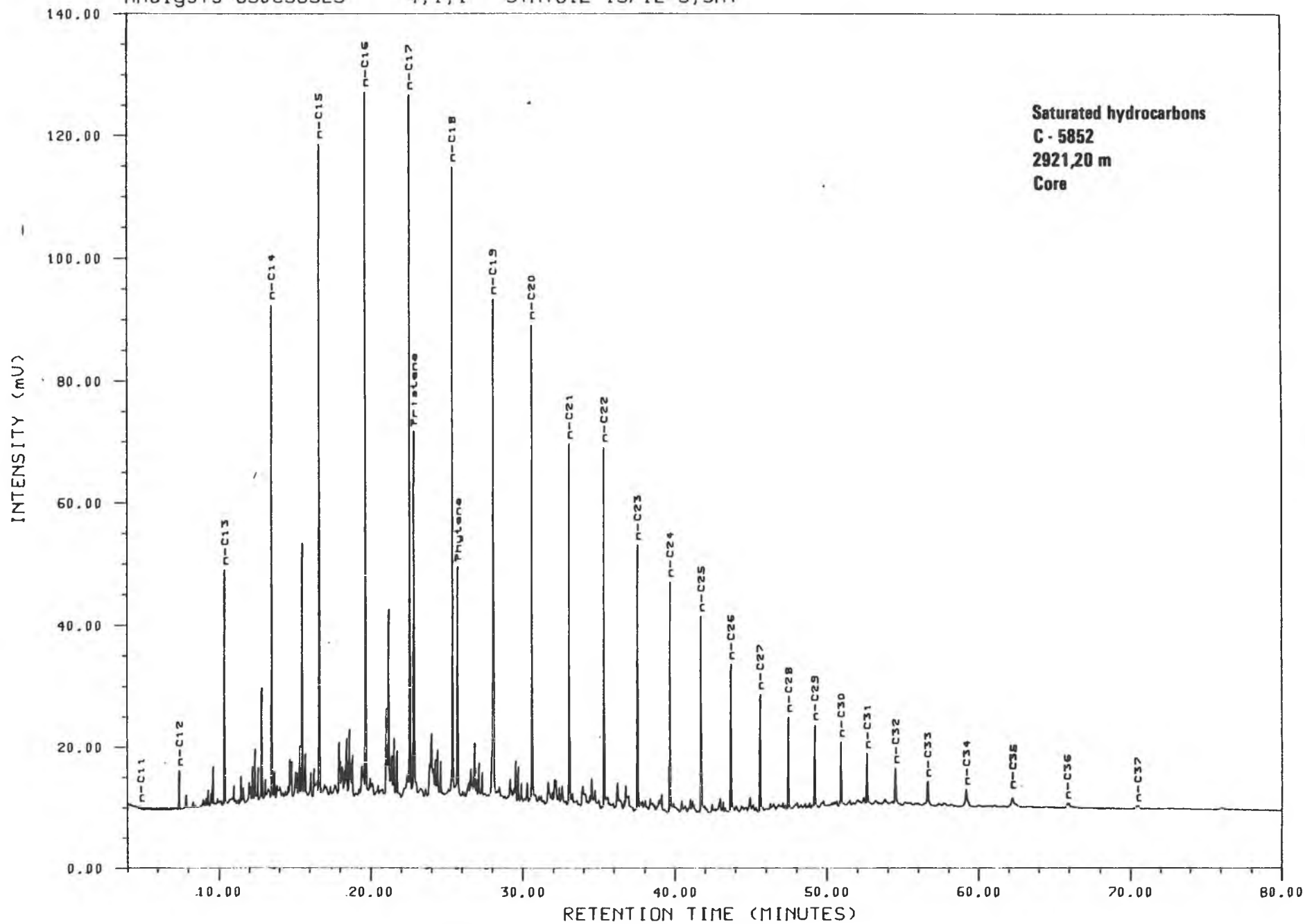






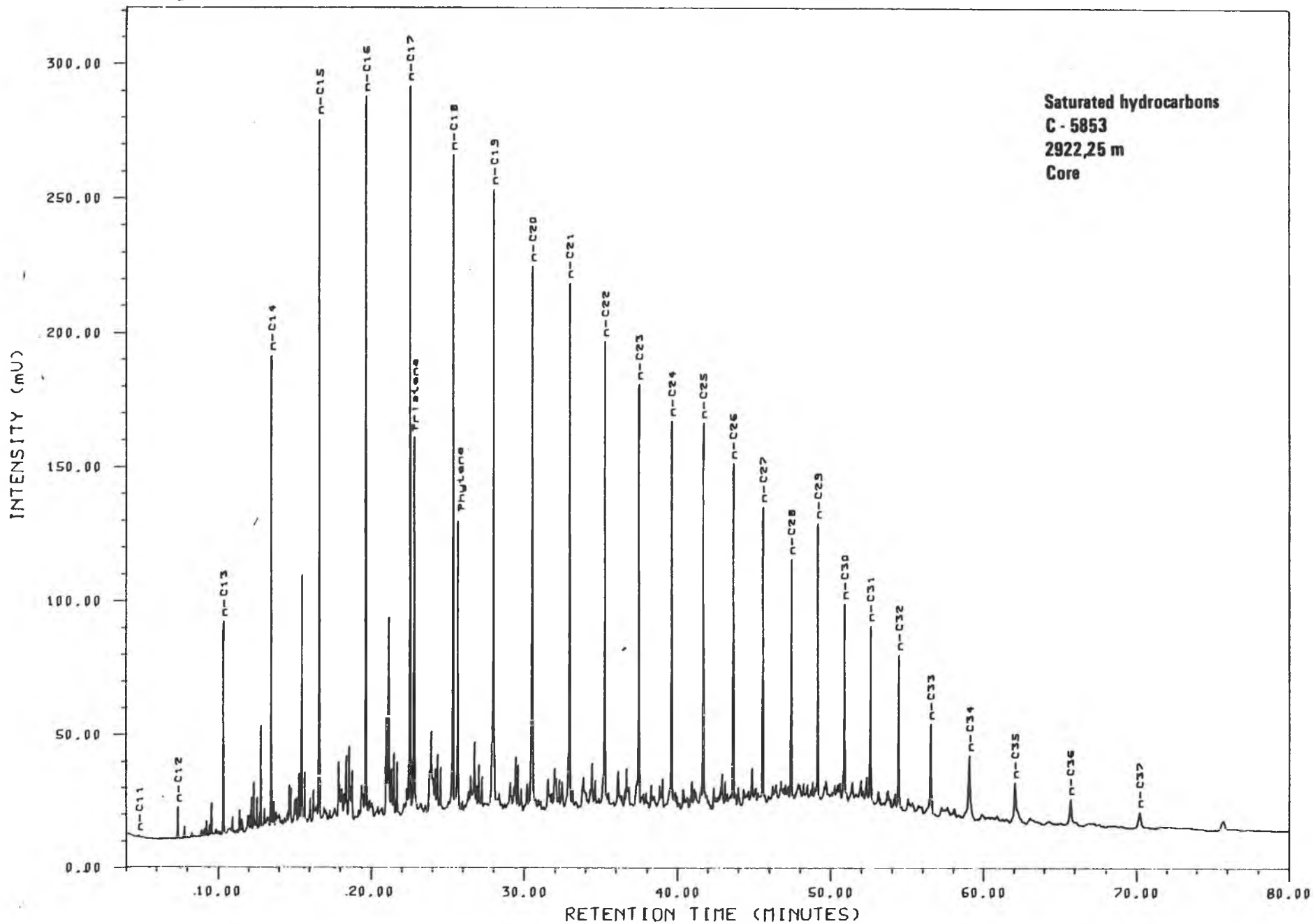


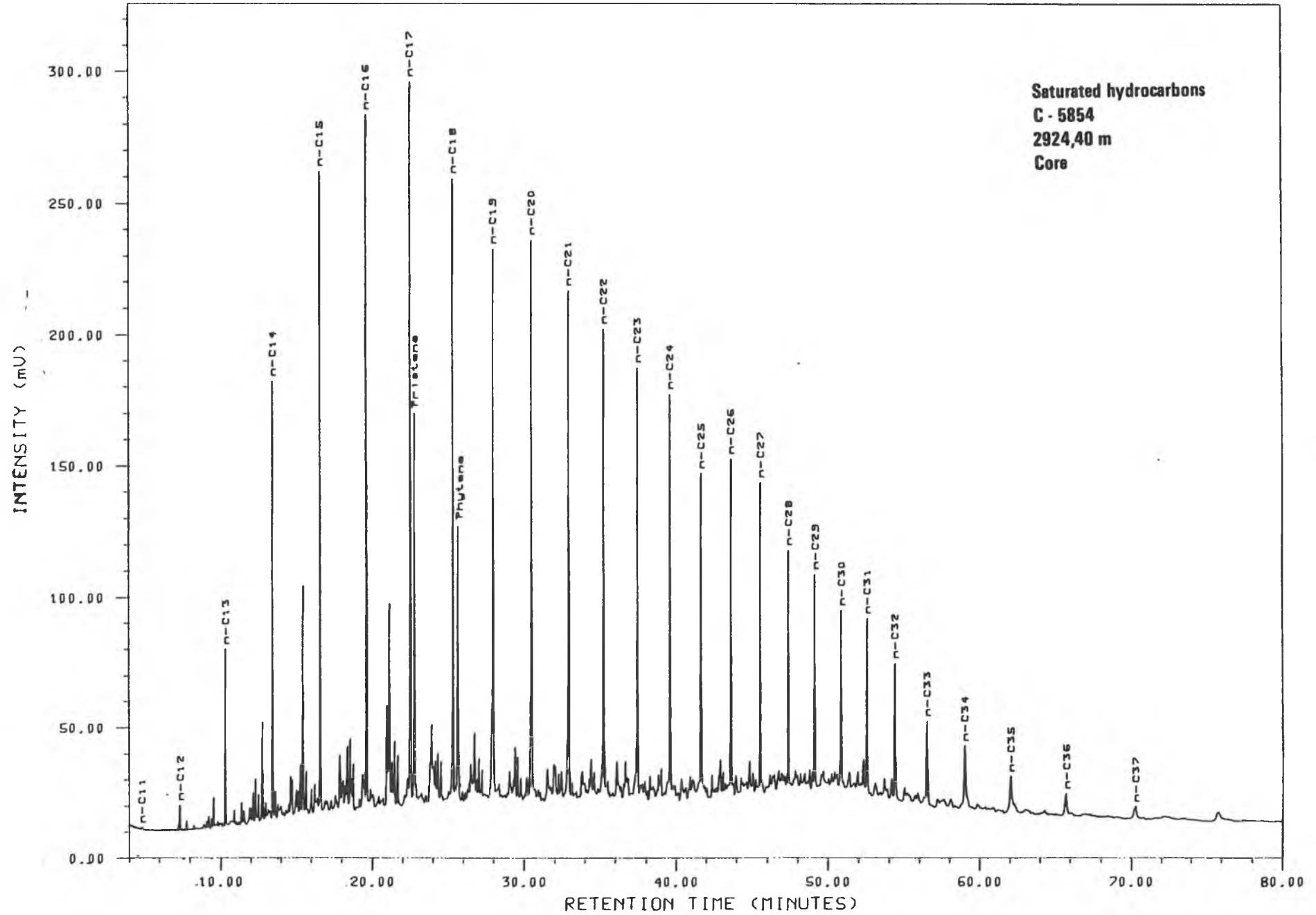


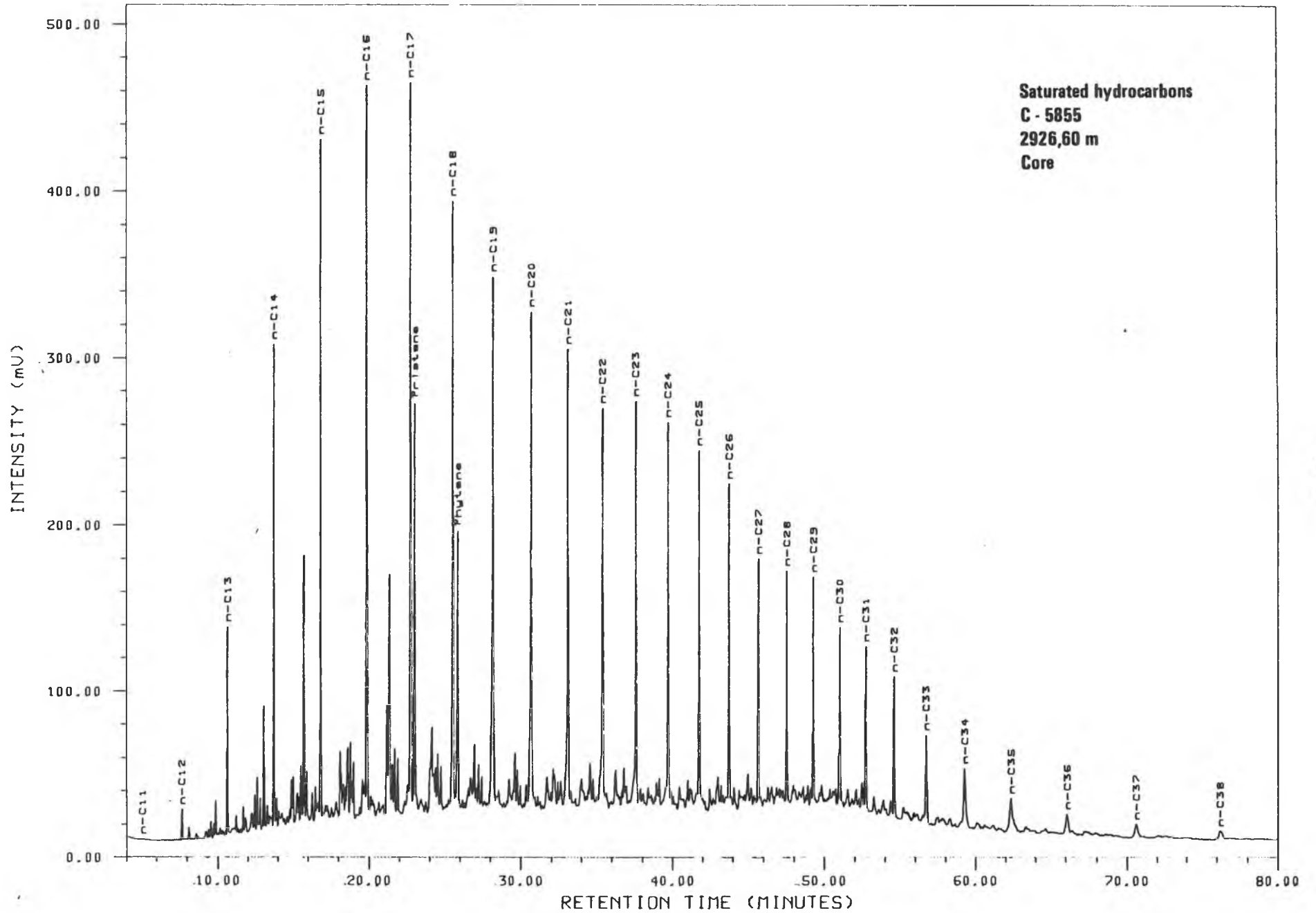


Analysis 850C5853S

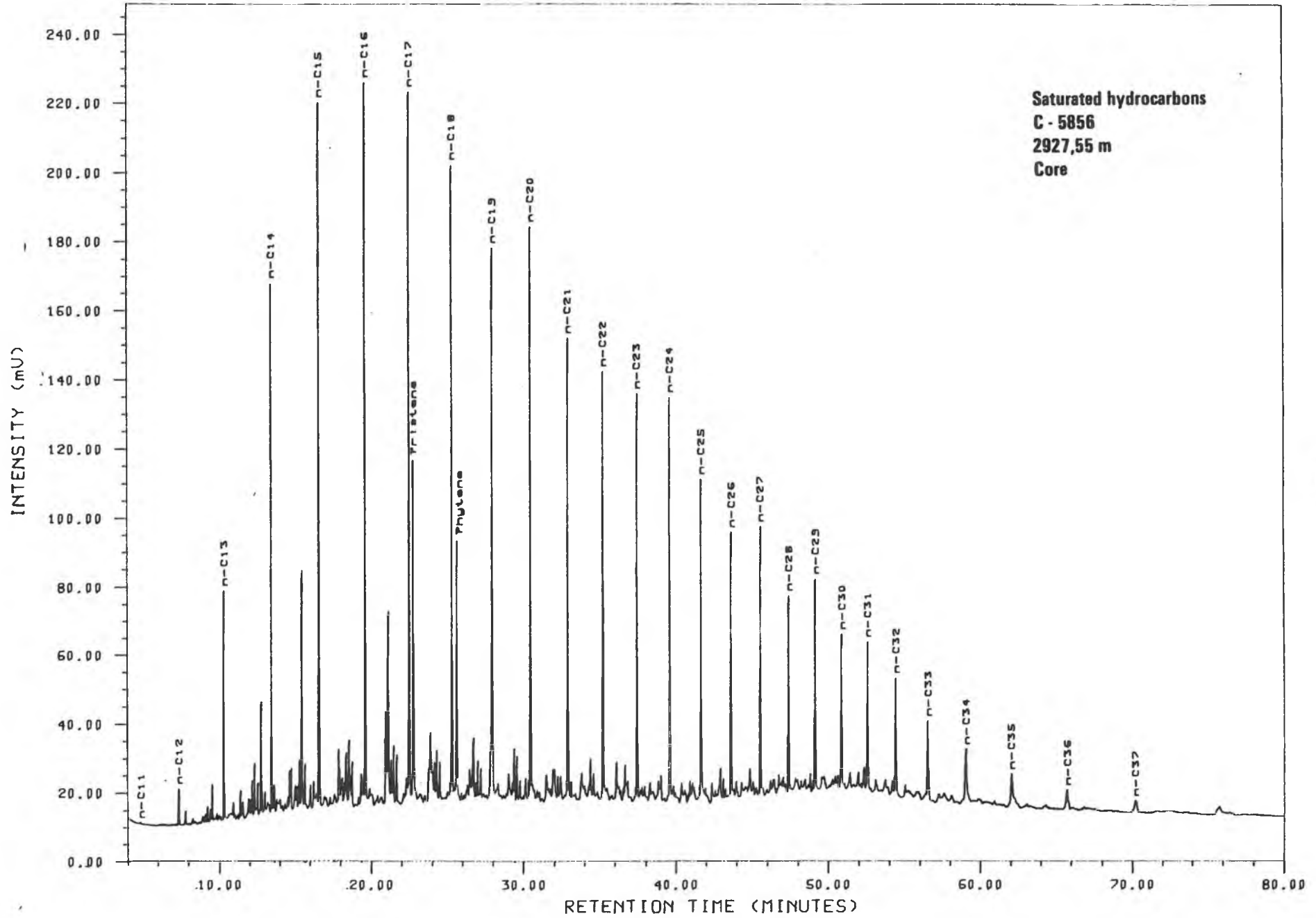
4,1,1 STATOIL 15/12-5,SAT

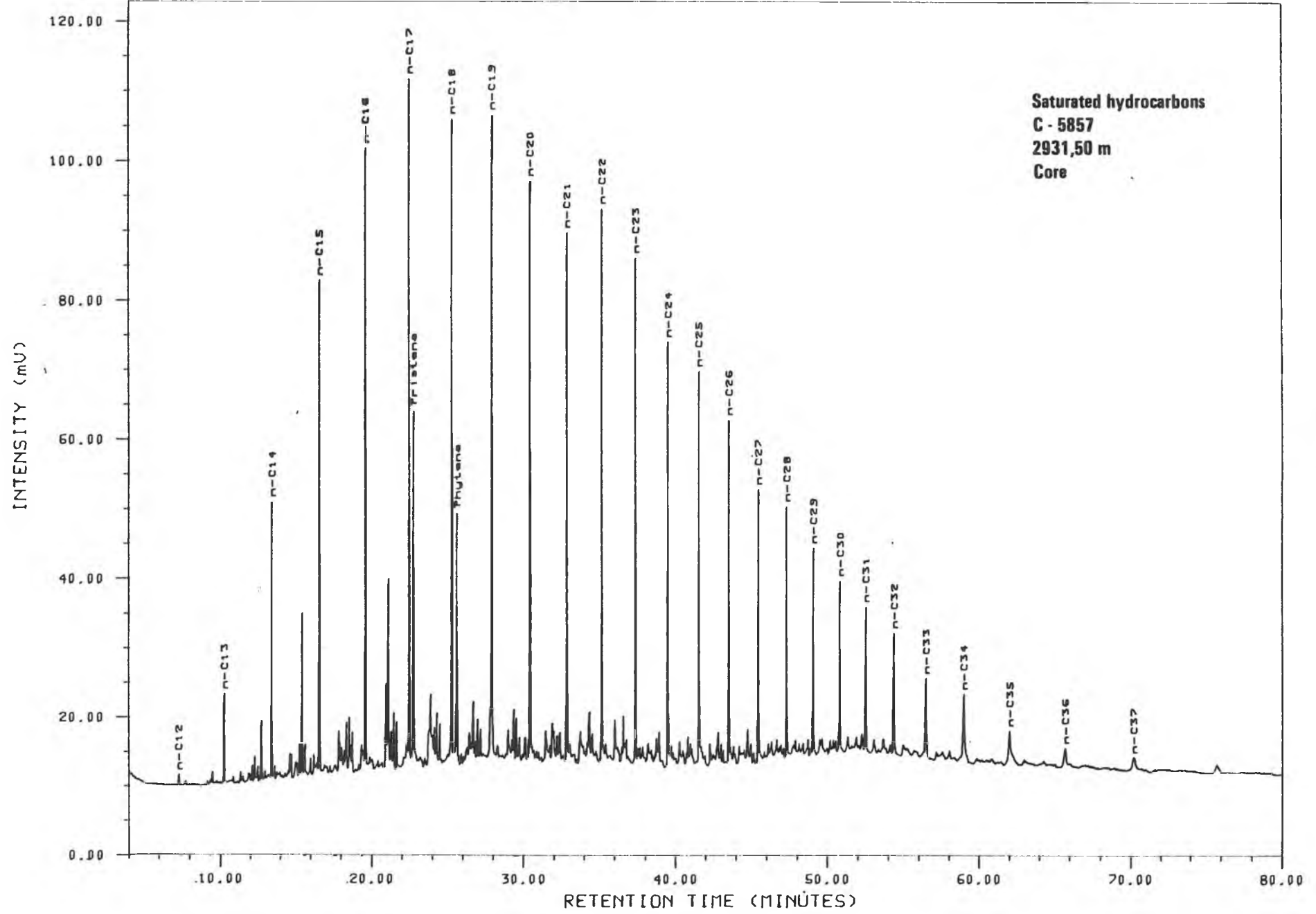


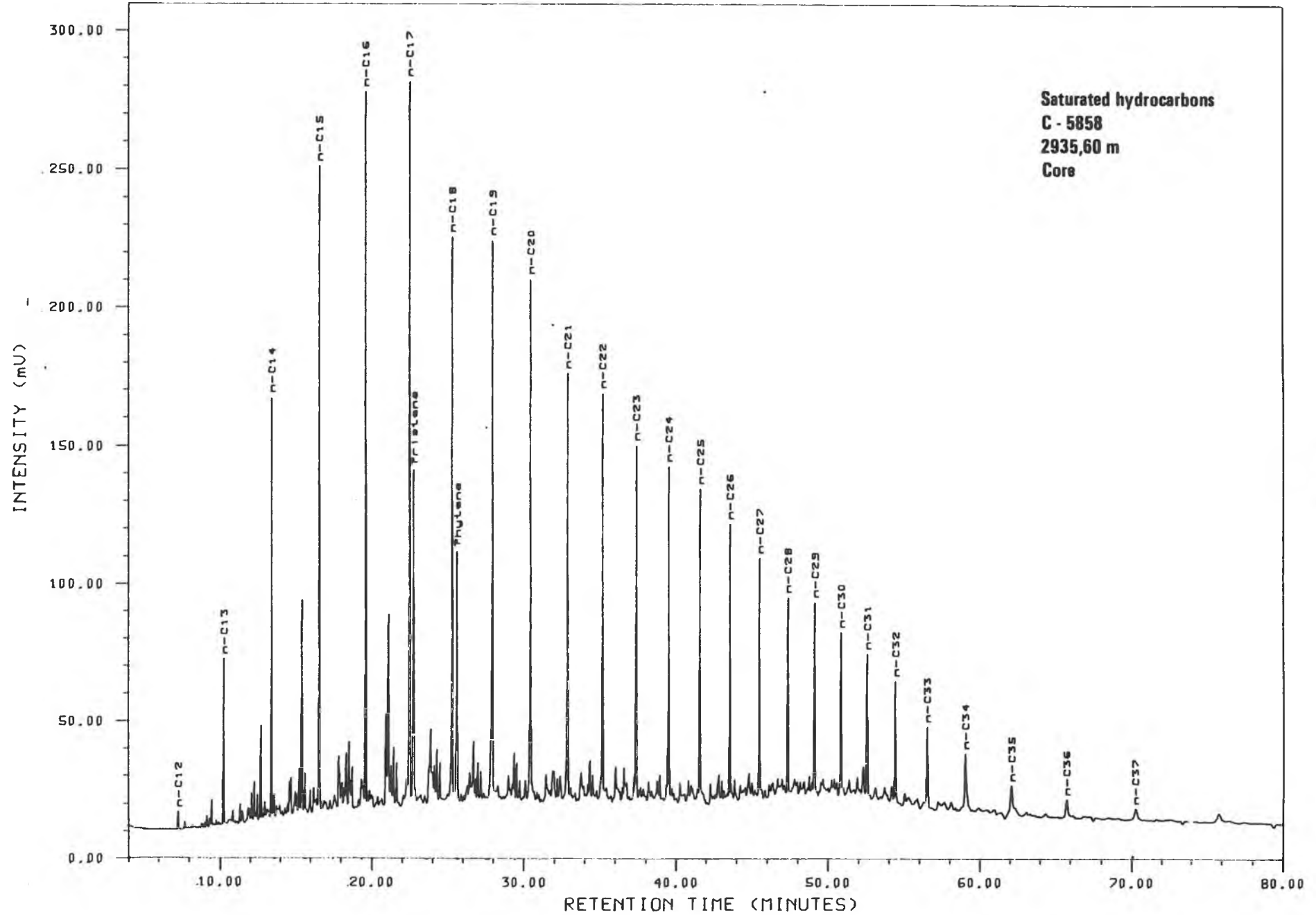




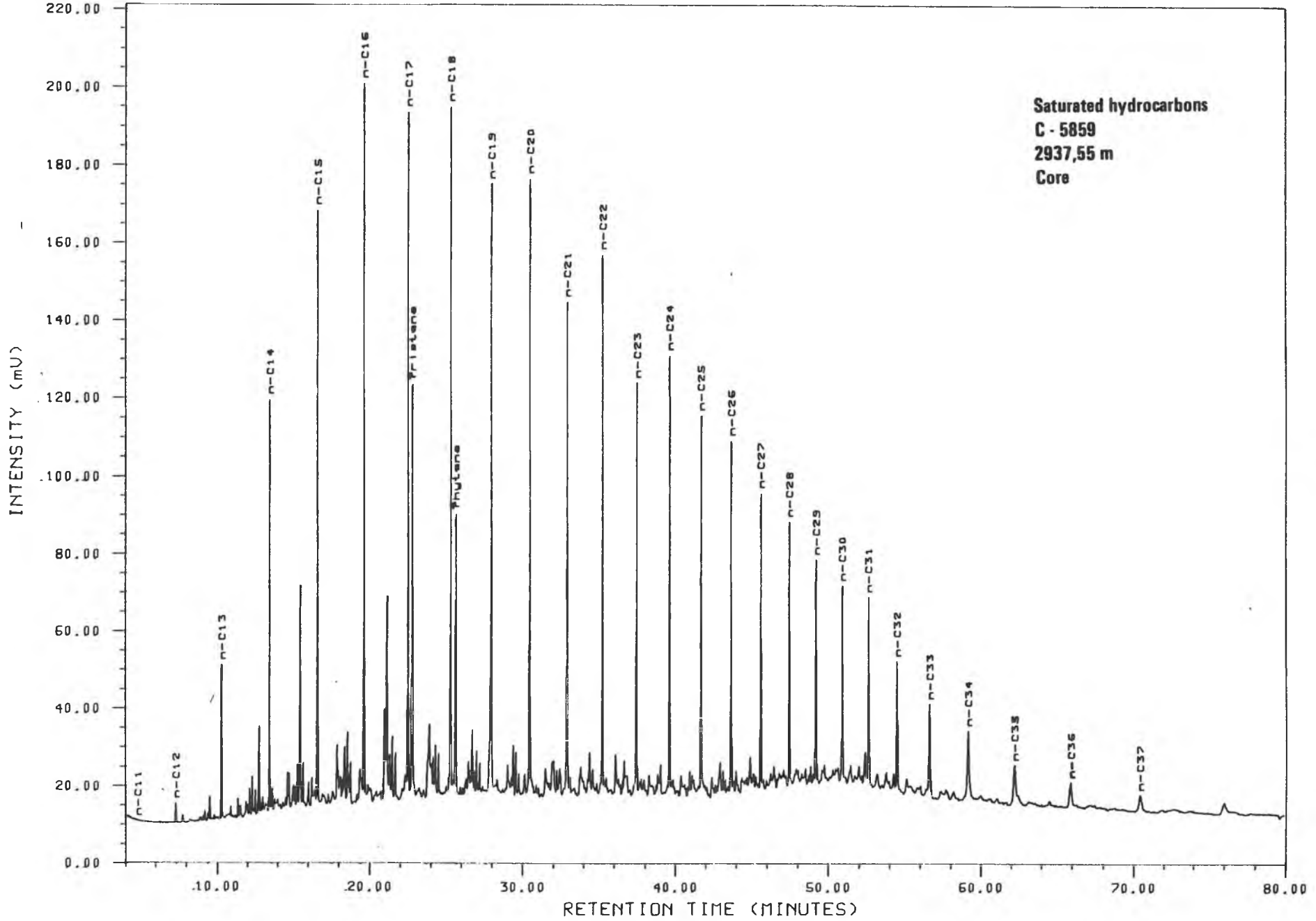
Analysis 850C5856S 4,1,1 STATOIL 15/12-5,SAT







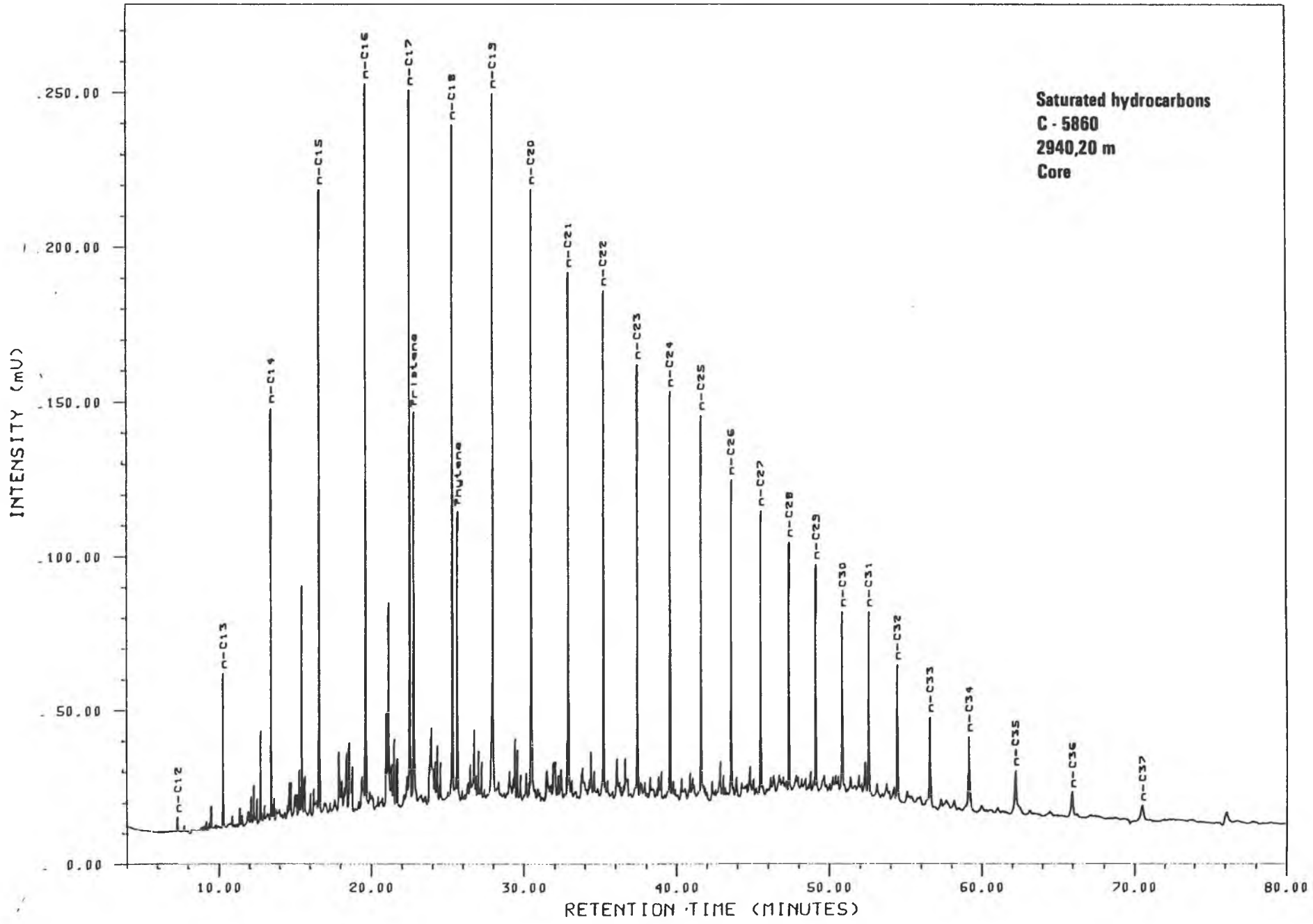
Analysis 850C5859S 4,1,1 STATOIL 15/12-5,SAT

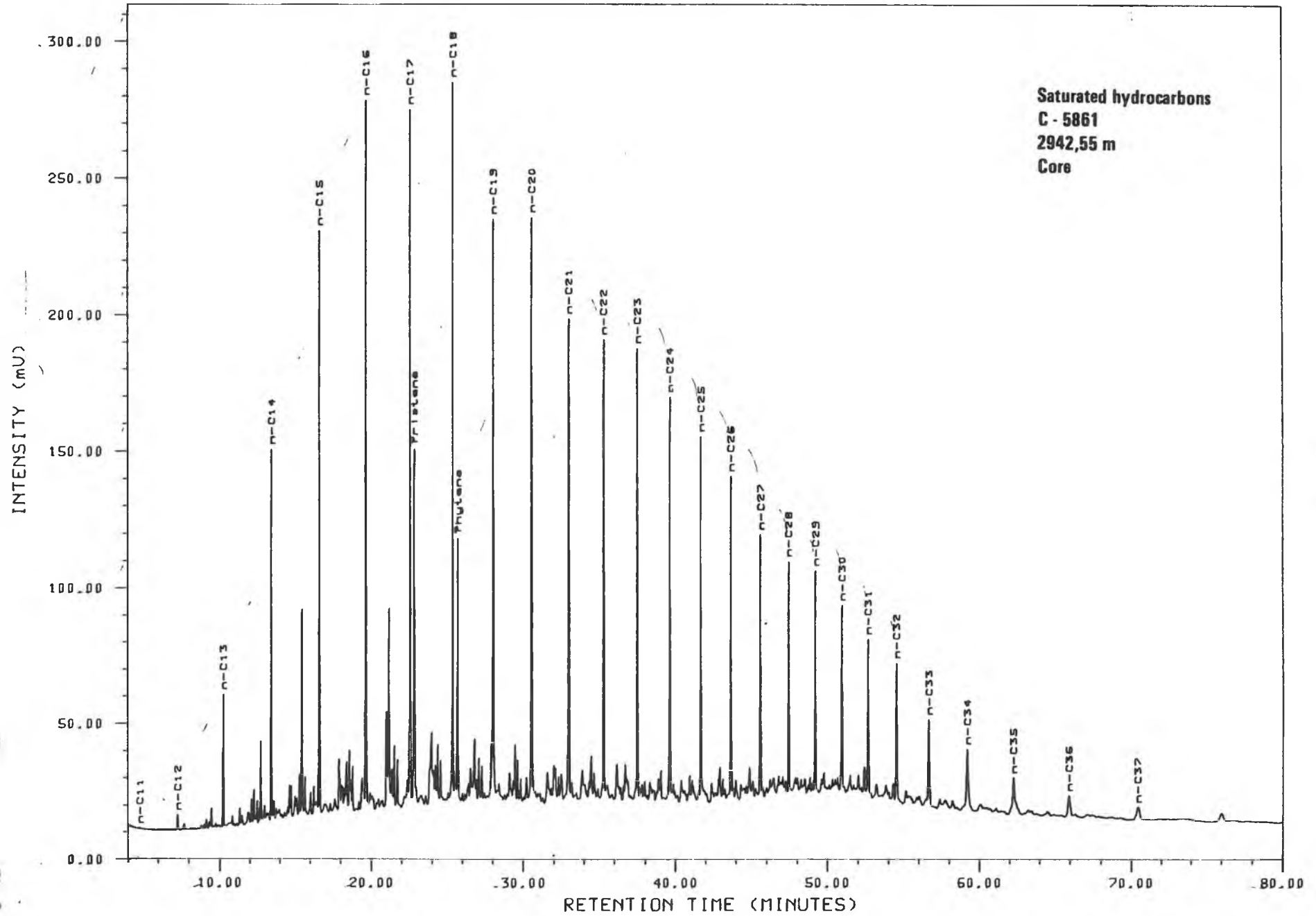


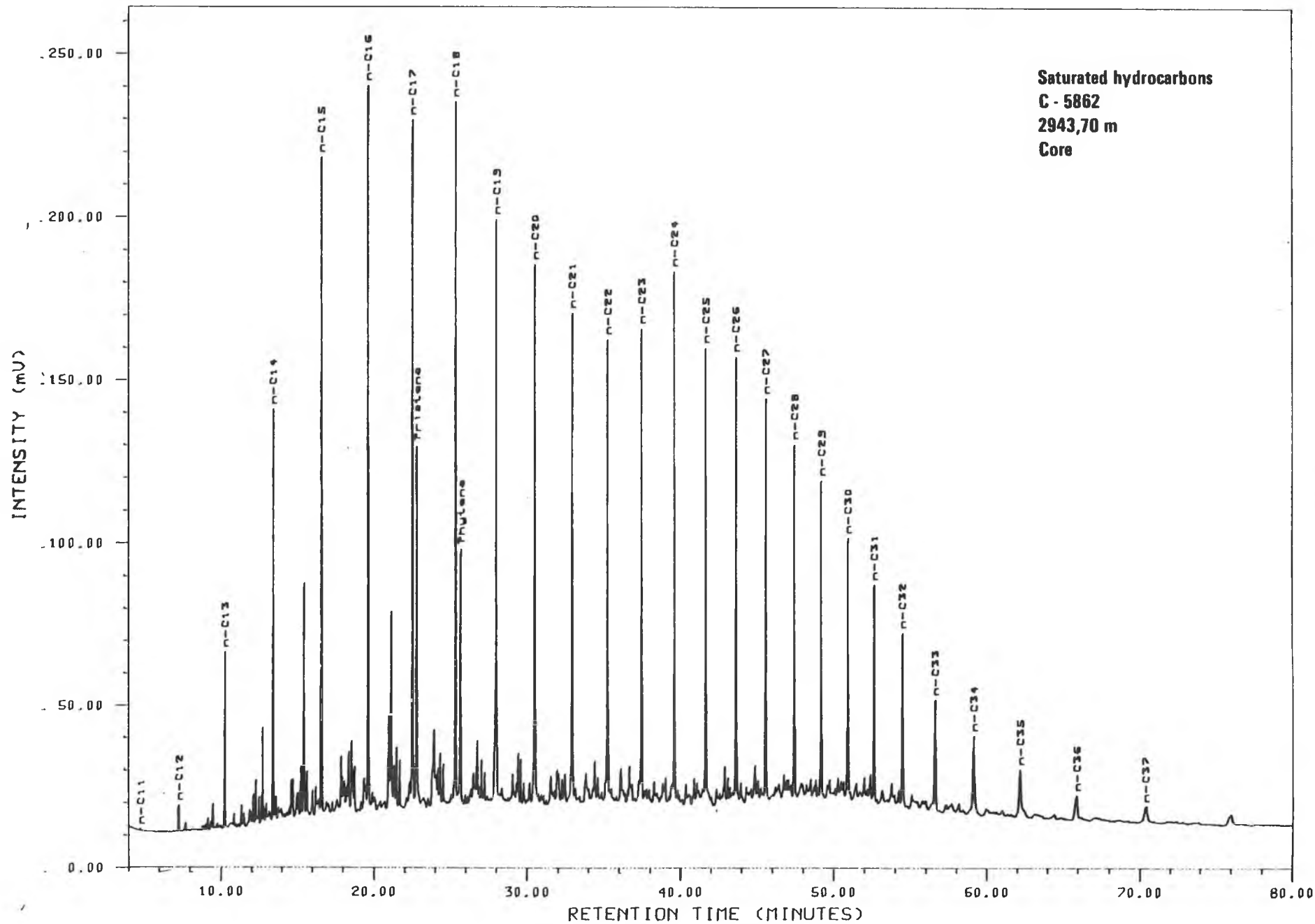
Analysis 850C5860S

4,1,1

STATOIL 15/12-5, SAT







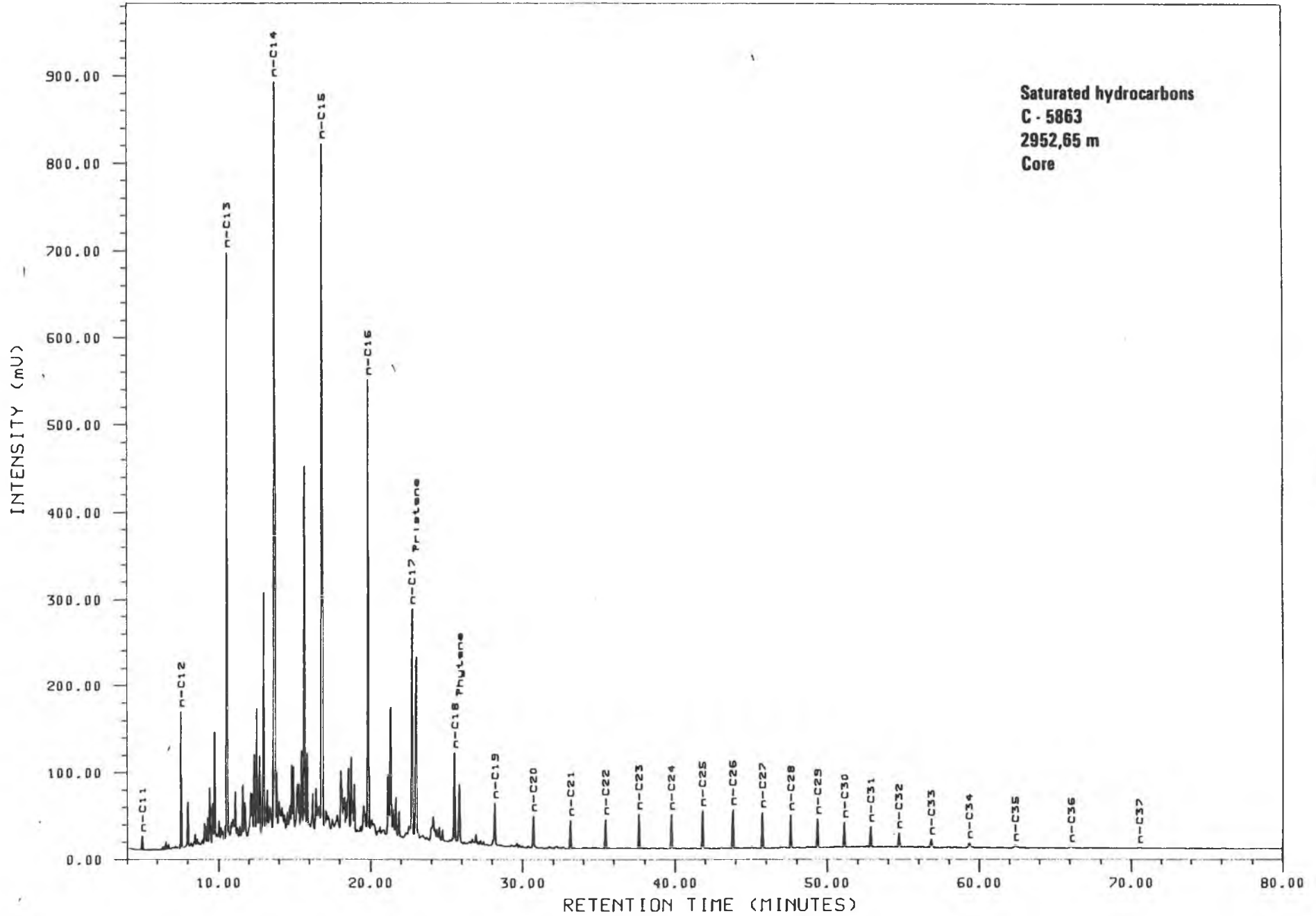
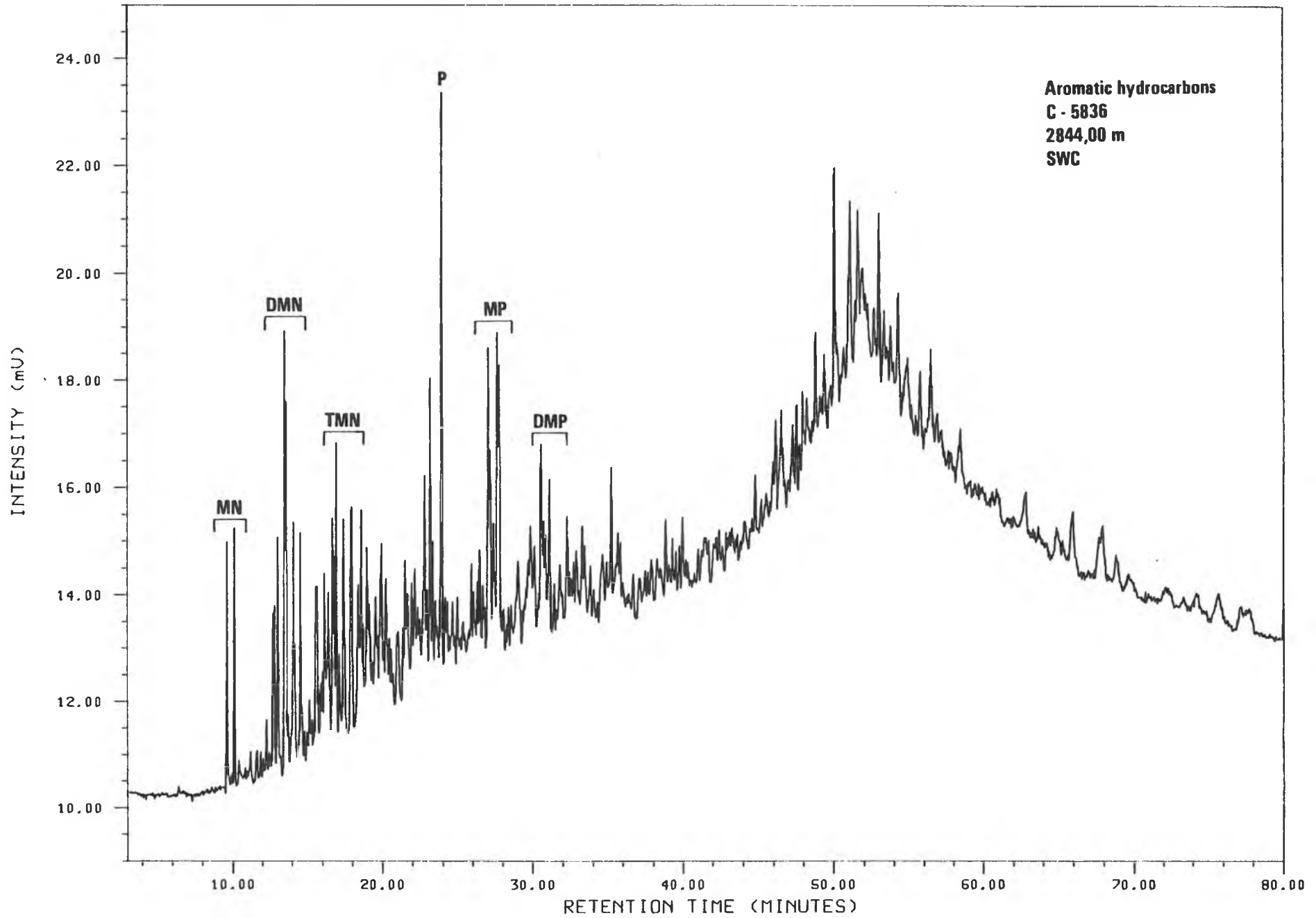


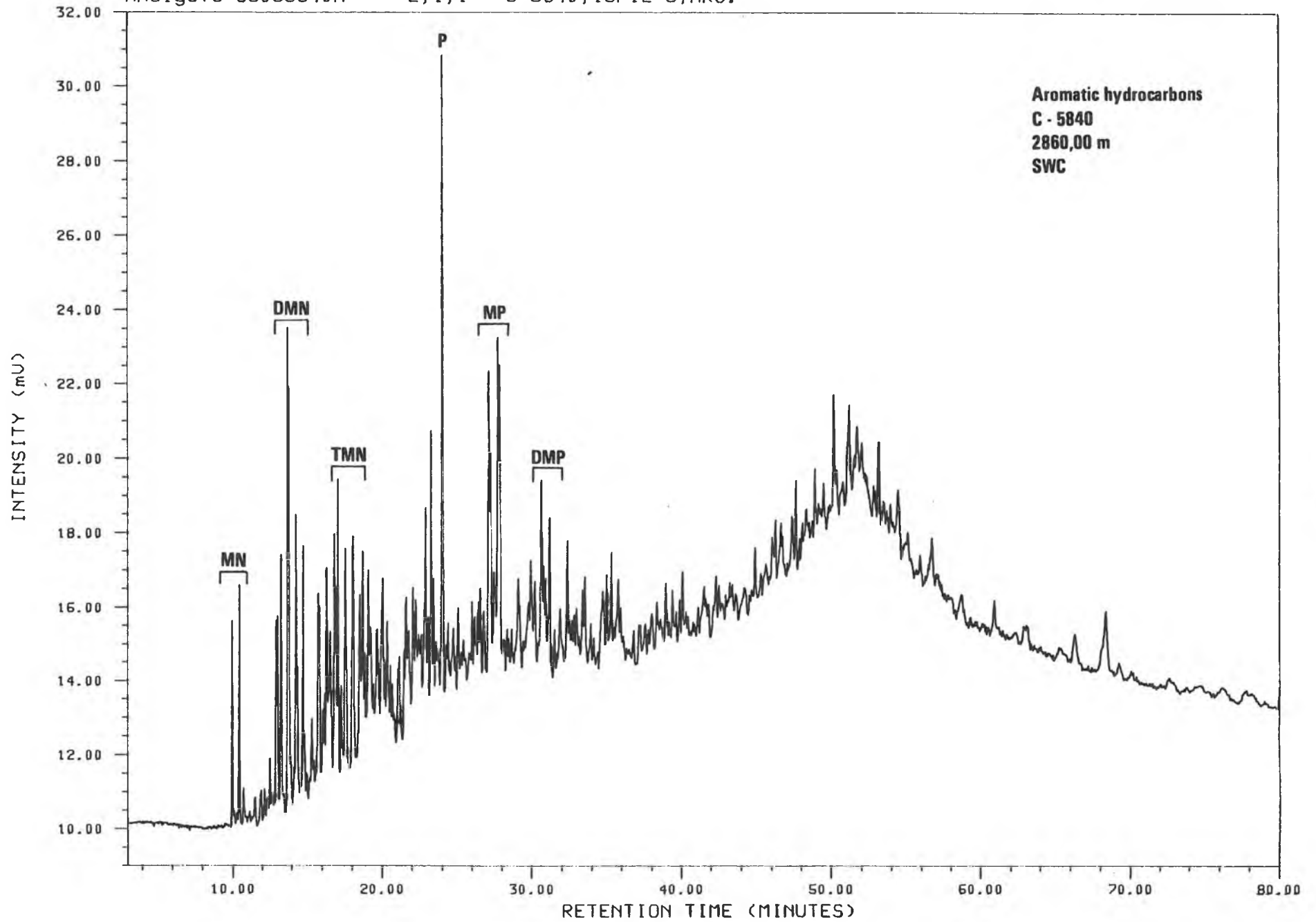
FIGURE 3

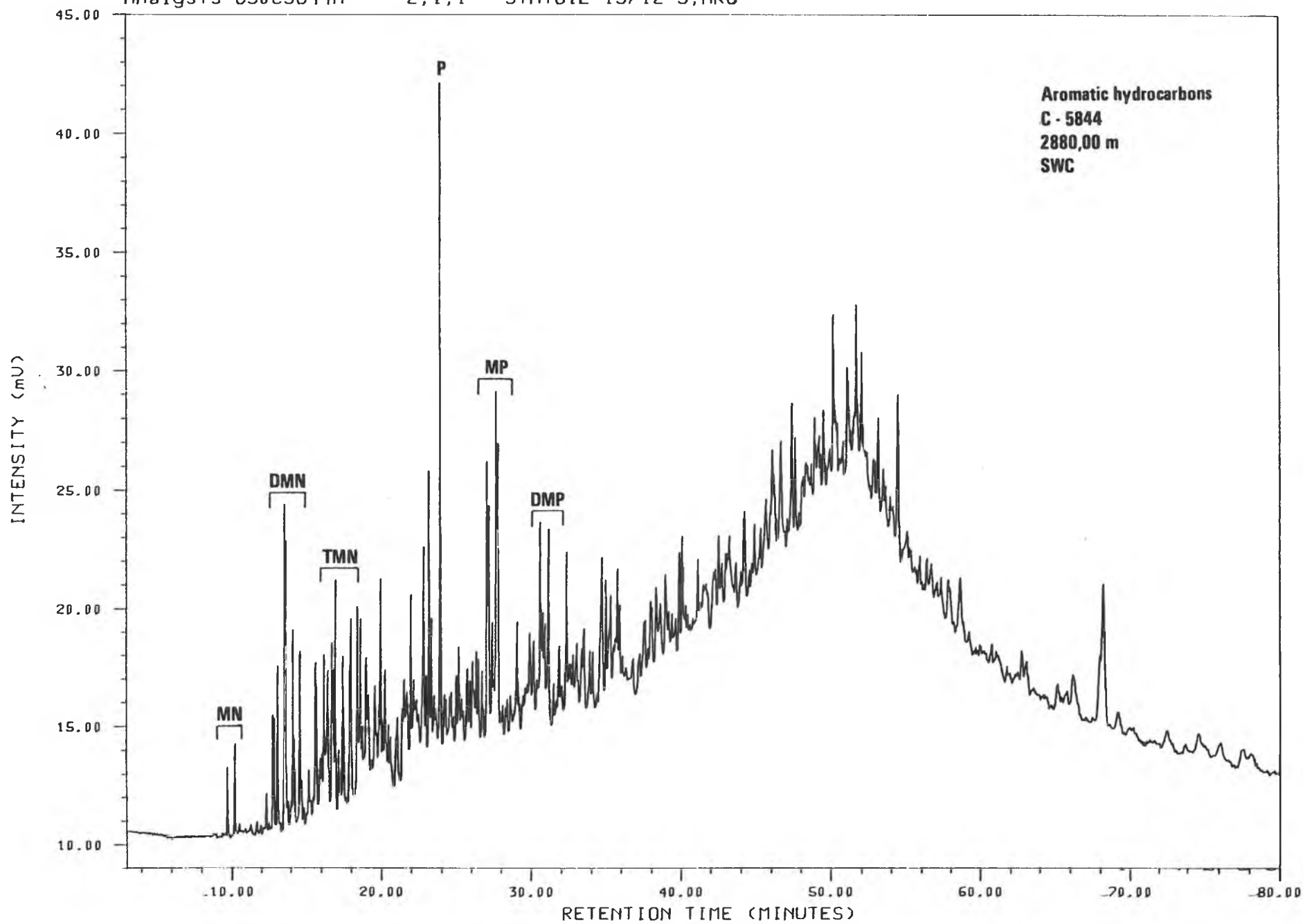
AROMATIC HYDROCARBON GAS CHROMATOGRAMS
FID DETECTION

N,MN,DMN,TMN	- naphthalene and alkylated homologs
P,MP,DMP	- phenanthrene and alkylated homologs

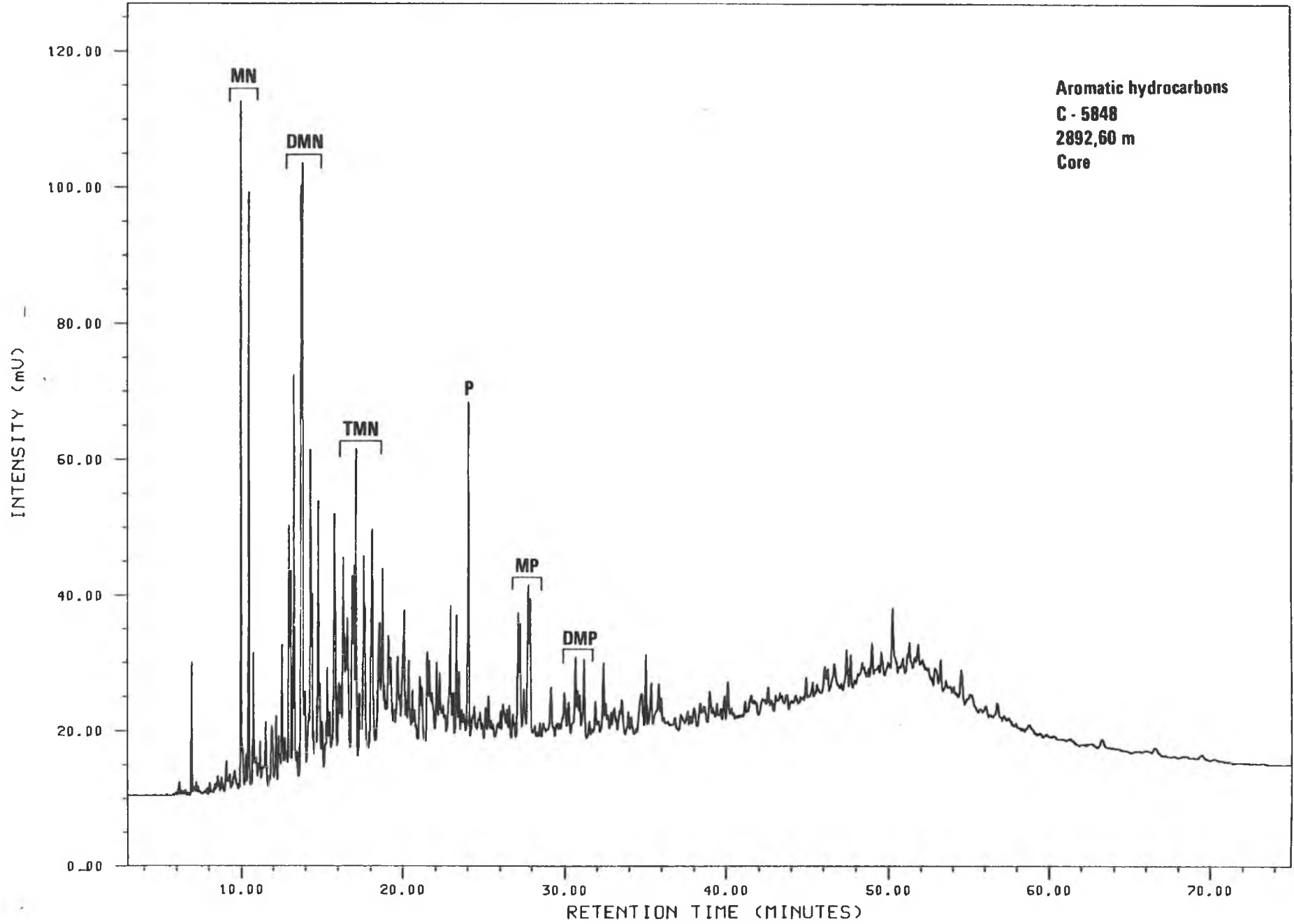


Analysis 850C5840A 2,1,1 C-5840,15/12-5,ARO.



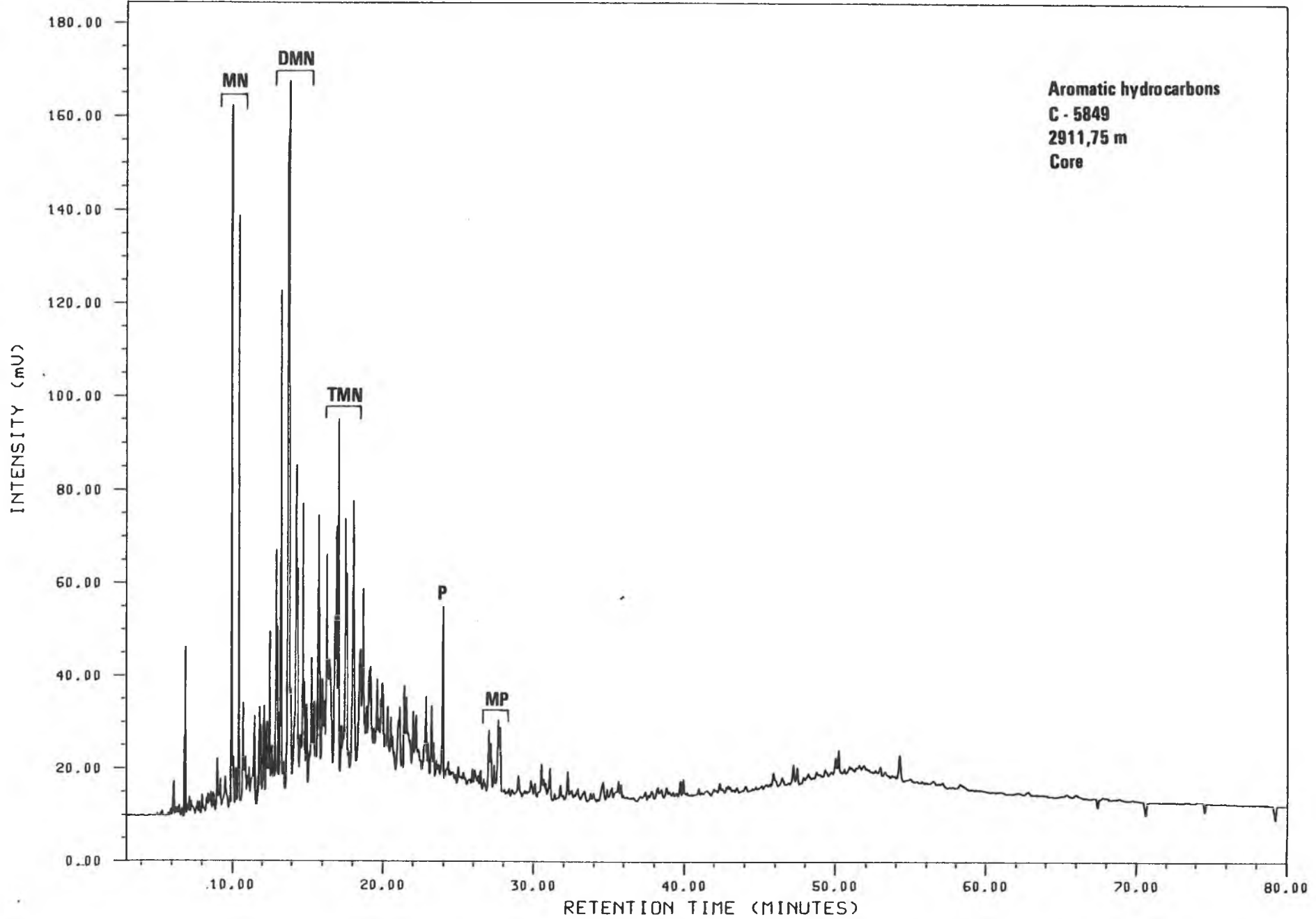


Analysis 1850AUT01 2,1,1 C-5848A

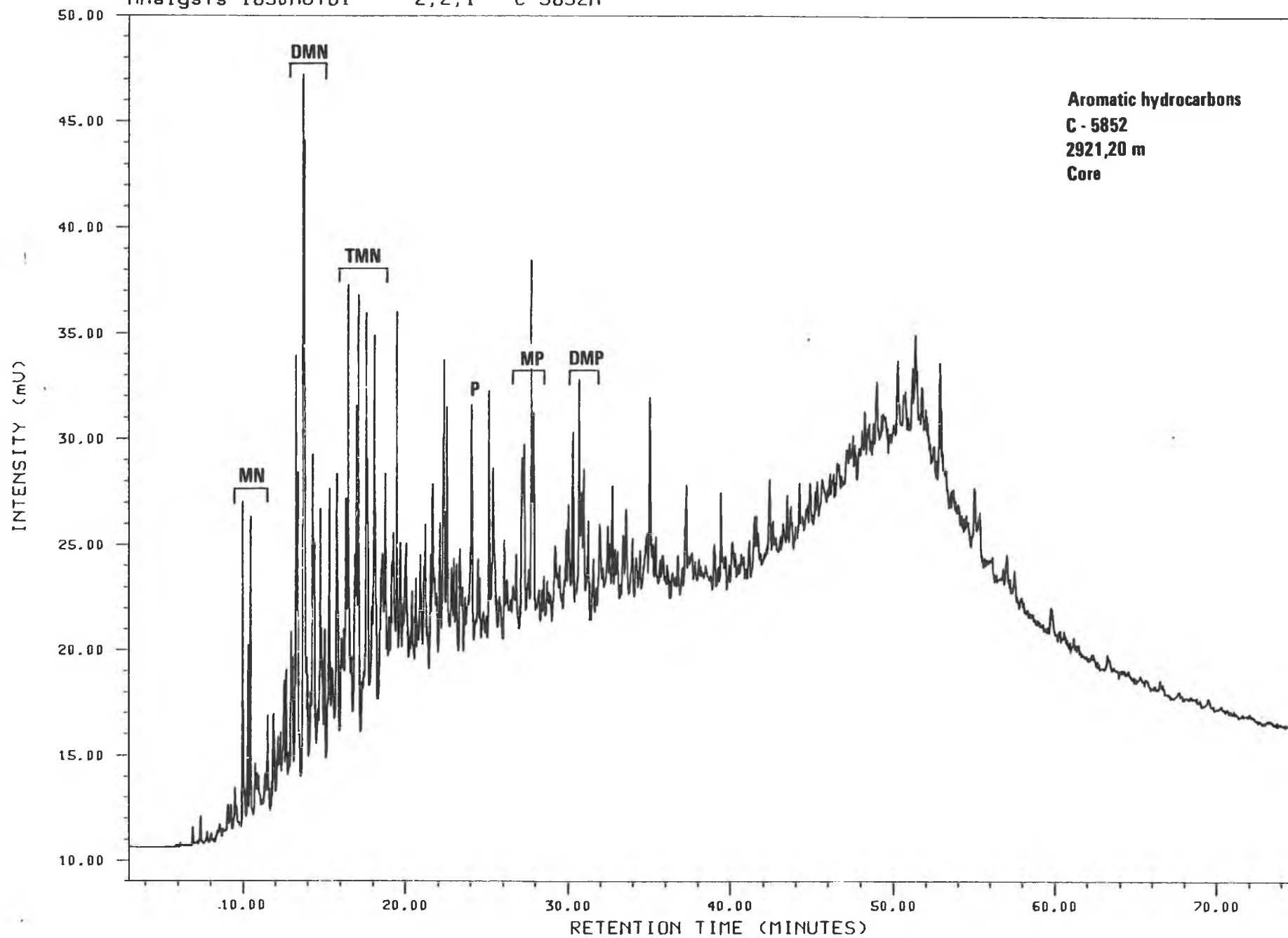


Aromatic hydrocarbons
C - 5848
2892,60 m
Core

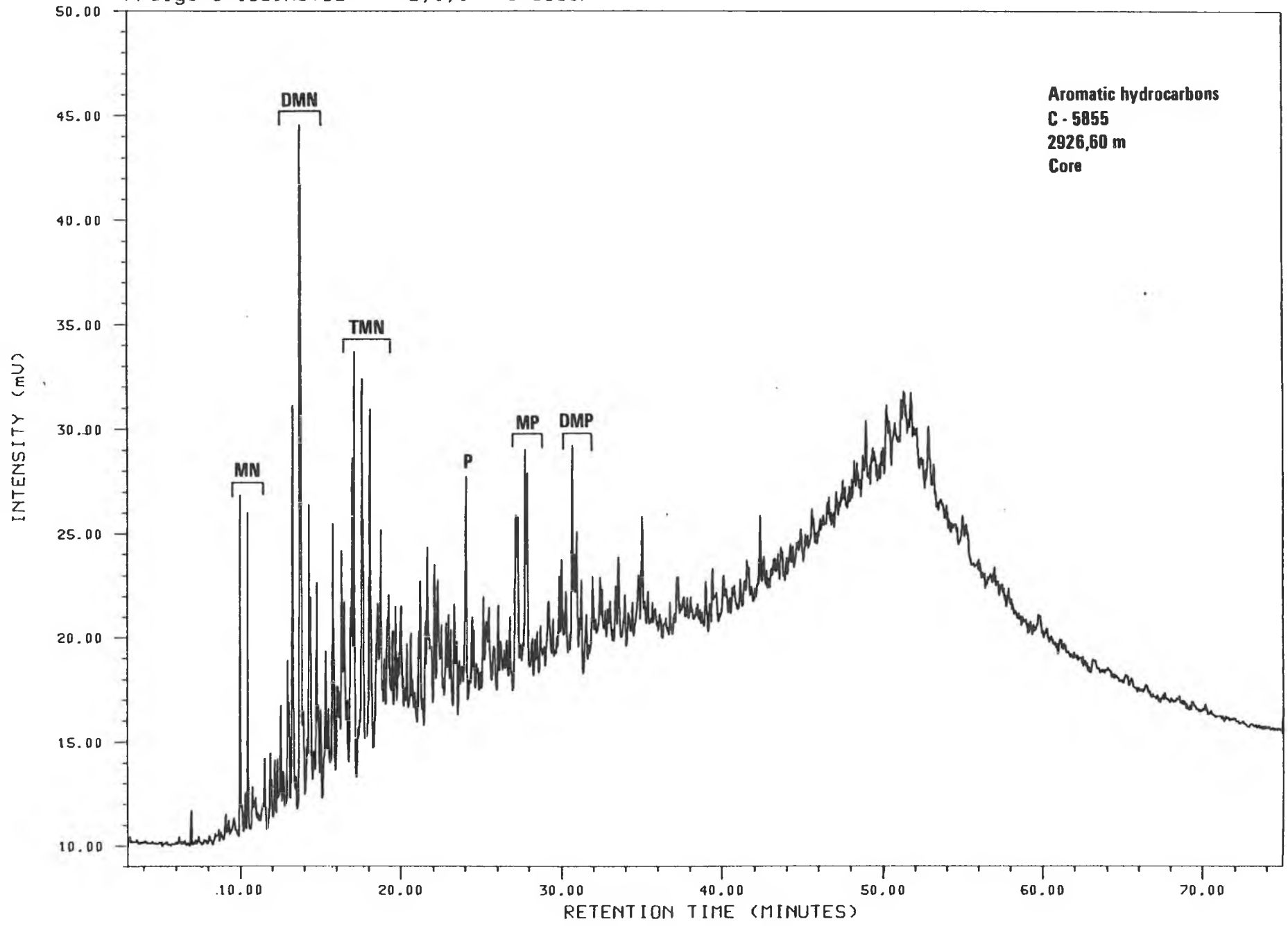
Analysis 850C5849A 2,1,1 STATOIL 15/12-5,ARO



Aromatic hydrocarbons
C - 5849
2911,75 m
Core

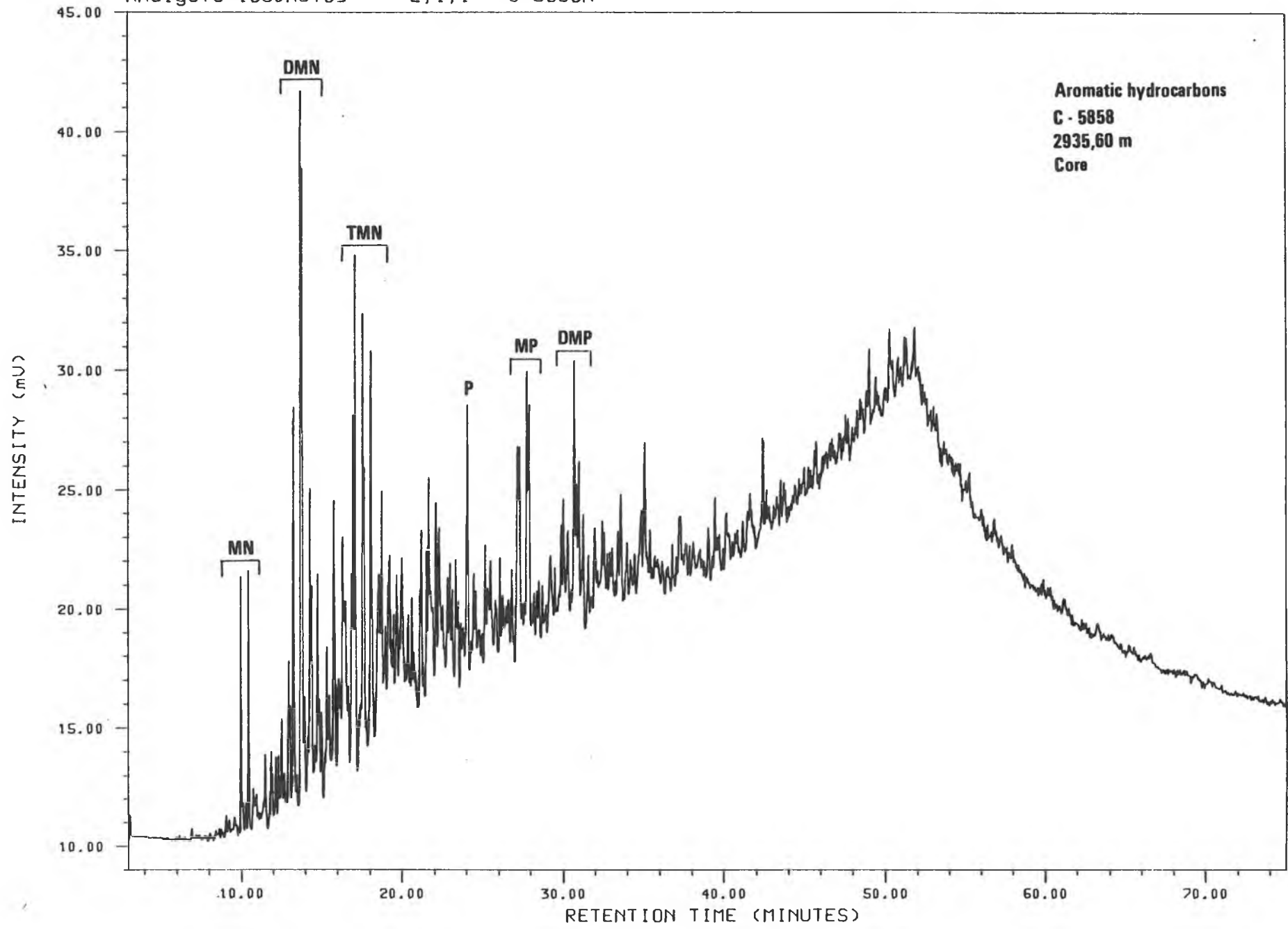


Aromatic hydrocarbons
C - 5852
2921,20 m
Core

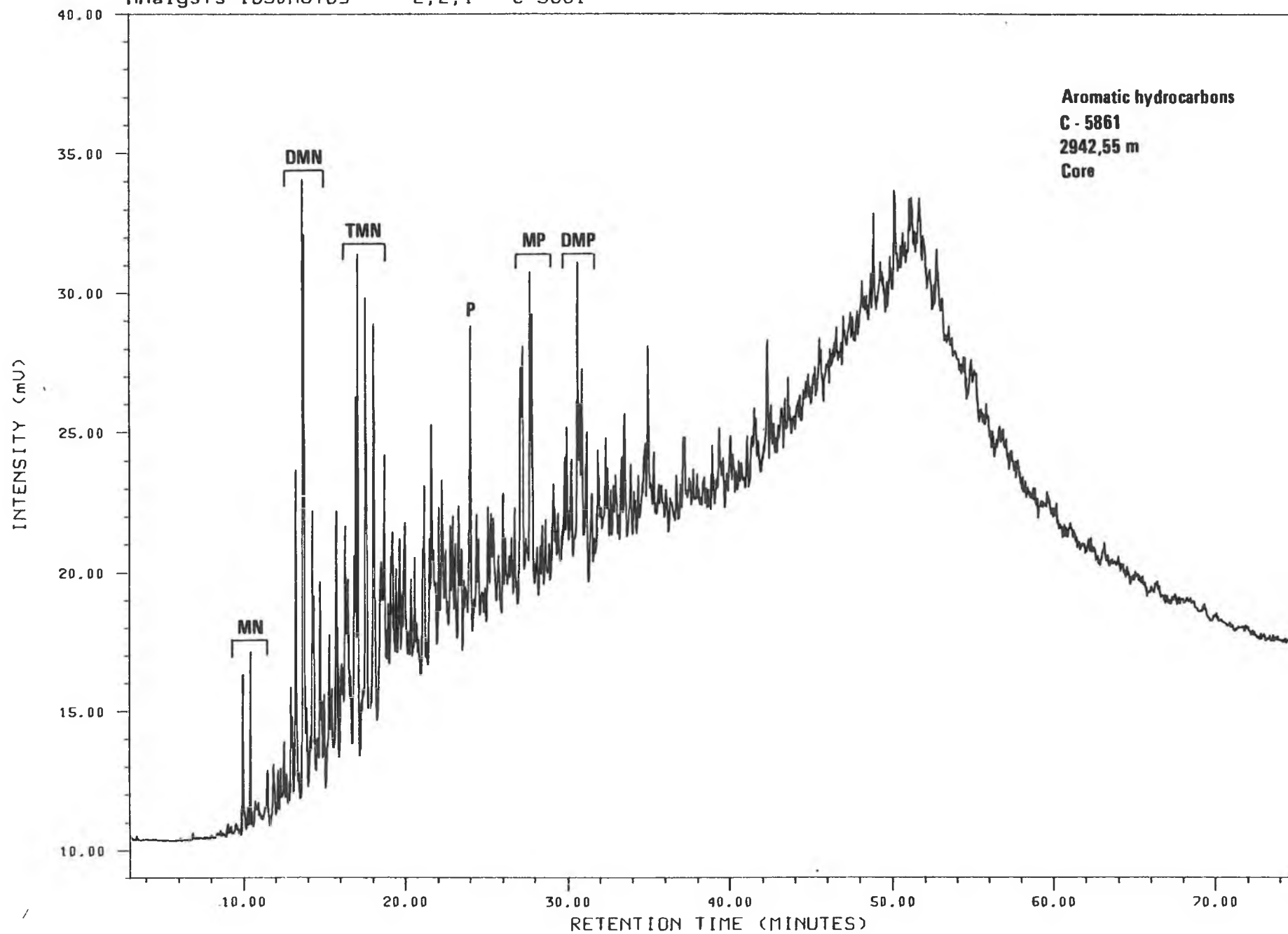


Aromatic hydrocarbons
C - 5855
2926,60 m
Core

Analysis 1850AUT03 2,1,1 C-5858A



Aromatic hydrocarbons
C - 5858
2935,60 m
Core



Aromatic hydrocarbons
C - 5861
2942,55 m
Core

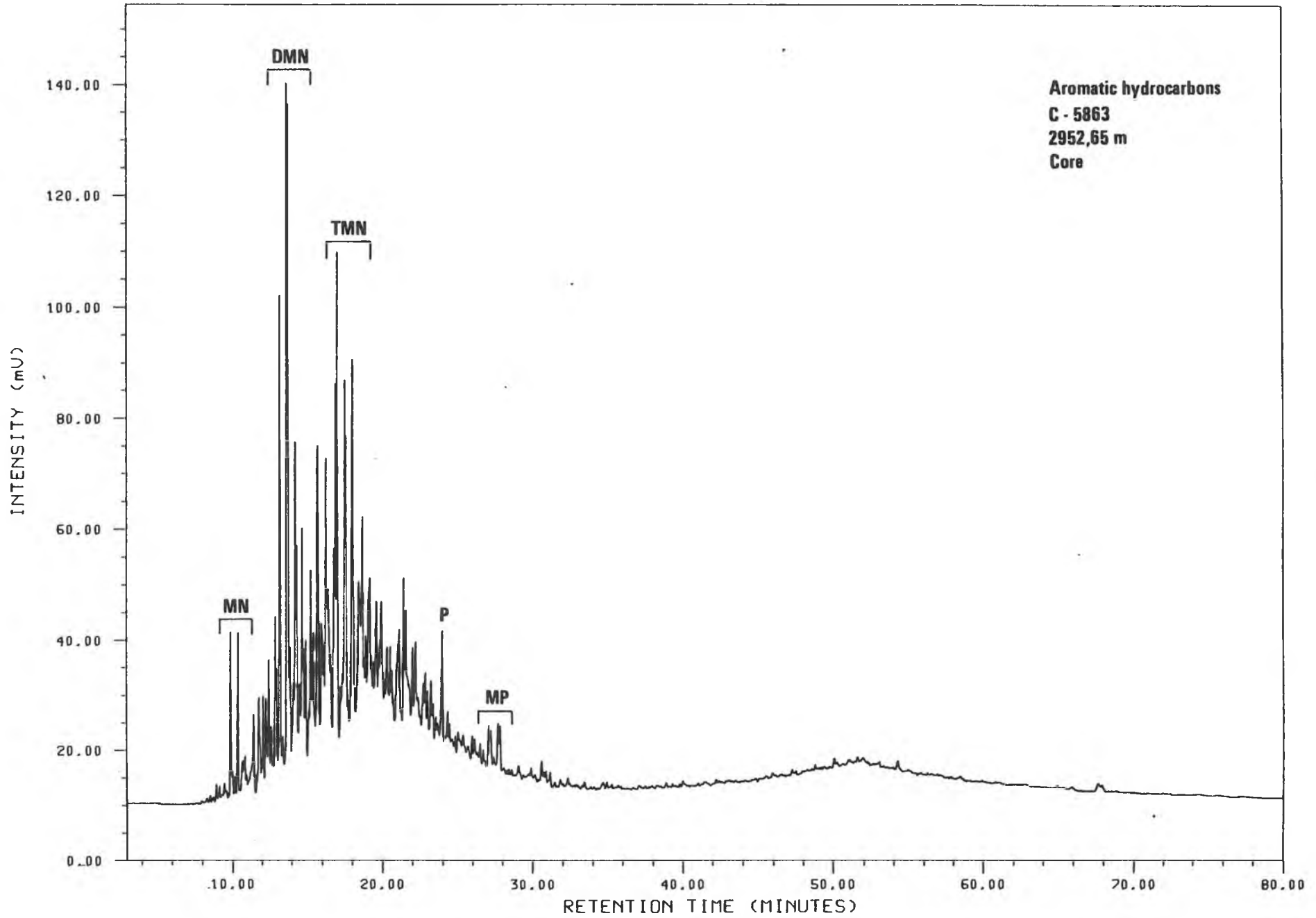
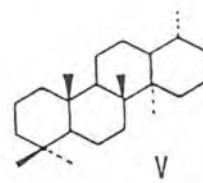
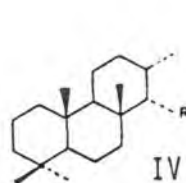
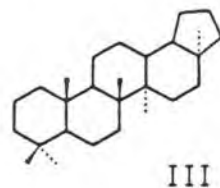
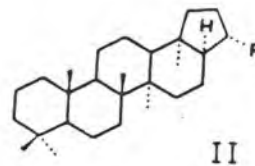
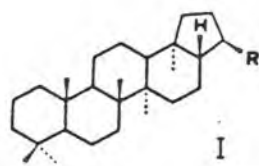


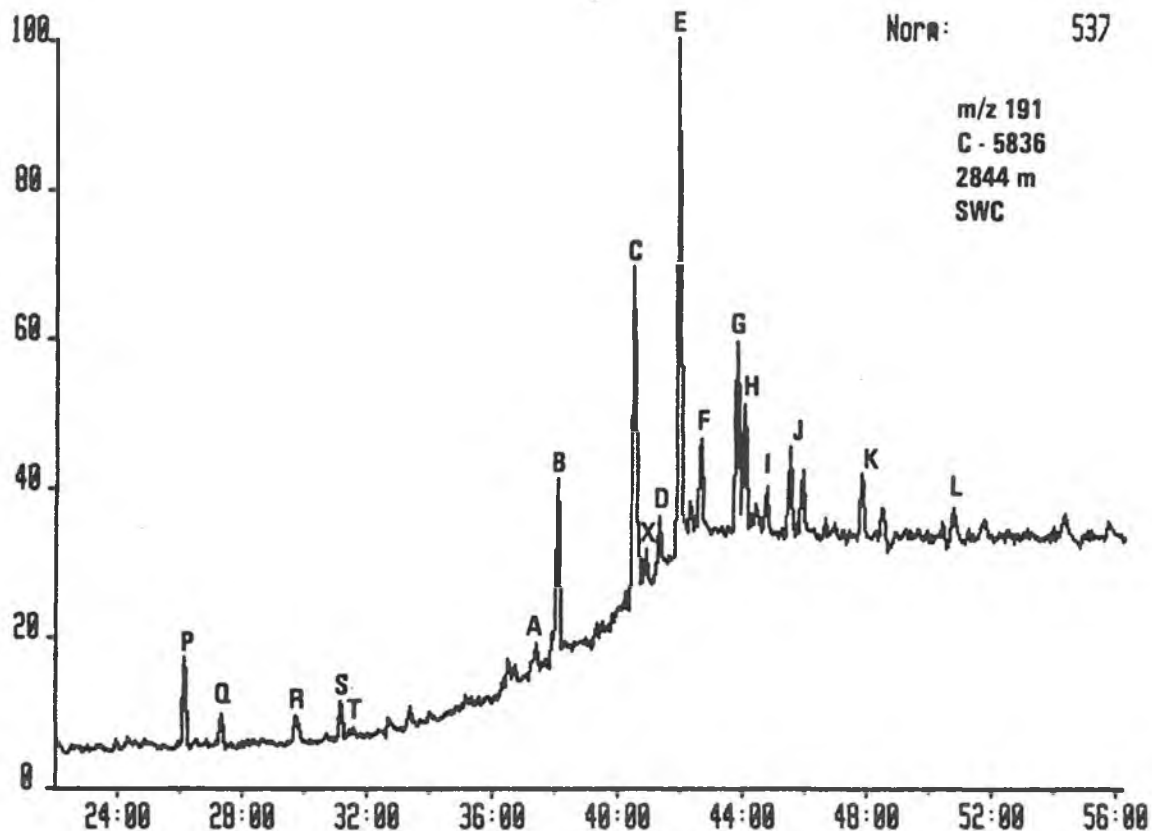
Figure 4.

Mass chromatograms representing terpanes (m/z 191) and m/z 205.

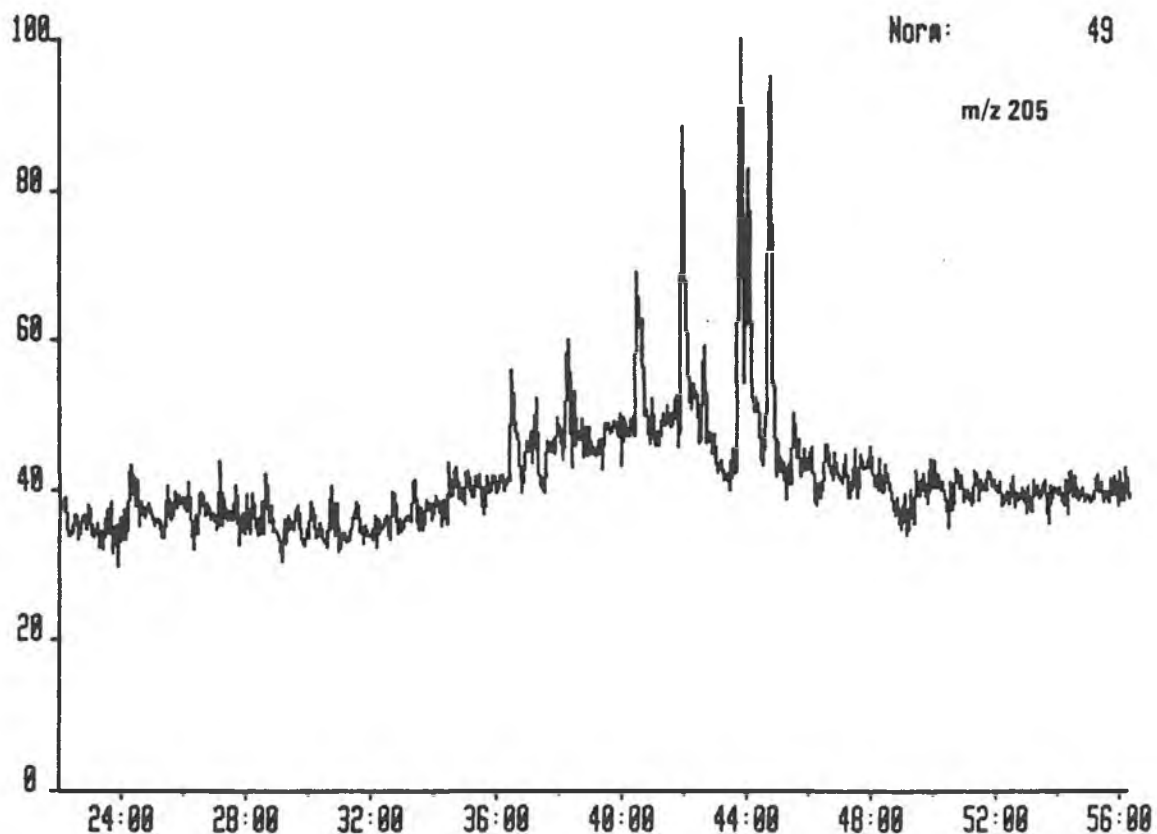
A	T_s , 18 α (H)-trisorneohopane	$C_{27}H_{46}$	(III)
B	T_m , 17 α (H)-trisnorhopane	$C_{27}H_{46}$	(I, R=H)
C	17 α (H)-norhopane	$C_{29}H_{50}$	(I, R= C_2H_5)
D	17 β (H)-normoretane	$C_{29}H_{50}$	(II, R= C_2H_5)
E	17 α (H)-hopane	$C_{30}H_{52}$	(I, R= C_3H_7)
F	17 β (H)-moretane	$C_{30}H_{52}$	(II, R= C_3H_7)
G	17 α (H)-homohopane (22S)	$C_{31}H_{54}$	(I, R= C_4H_9)
H	17 α (H)-homohopane (22R)	$C_{31}H_{54}$	(I, R= C_4H_9)
	+ unknown triterpane (gammacerane?)		
I	17 β (H)-homomoretane	$C_{31}H_{54}$	(II, R= C_4H_9)
J	17 α (H)-bishomohopane (22S,22R)	$C_{32}H_{56}$	(I, R= C_5H_{11})
K	17 α (H)-trishomohopane (22S,22R)	$C_{33}H_{58}$	(I, R= C_6H_{13})
L	17 α (H)-tetrakishomohopane (22S,22R)	$C_{34}H_{60}$	(I, R= C_7H_{15})
M	17 α (H)-pentakishomohopane (22S,22R)	$C_{35}H_{62}$	(I, R= C_8H_{17})
Z	bisnorhopane	$C_{28}H_{48}$	
X	unknown triterpane	$C_{30}H_{52}$	
P	tricyclic terpene	$C_{23}H_{42}$	(IV, R= C_4H_9)
Q	tricyclic terpene	$C_{24}H_{44}$	(IV, R= C_5H_{11})
R	tricyclic terpene (17R,17S)	$C_{25}H_{46}$	(IV, R= C_6H_{13})
S	tetracyclic terpene	$C_{24}H_{42}$	(V)
T	tricyclic terpene (17R,17S)	$C_{26}H_{48}$	(IV, R= C_7H_{15})



C5836SAT 5-FEB-07 Sir:Voltage 7070H Acnt:IKU System:TRIT
 Sample 1 Injection 1 Group 1 Mass 191.1000
 Text:

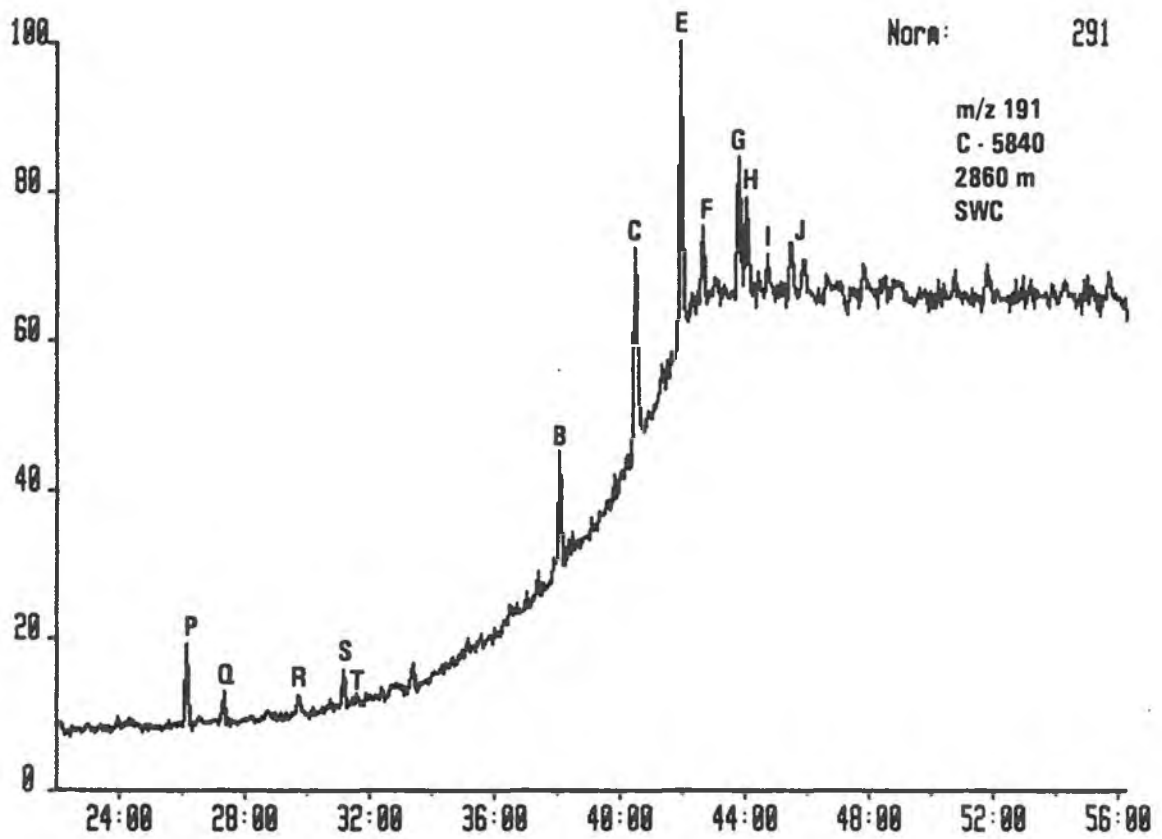


C5836SAT 5-FEB-07 Sir:Voltage 7070H Acnt:IKU System:TRIT
 Sample 1 Injection 1 Group 1 Mass 205.1000
 Text:



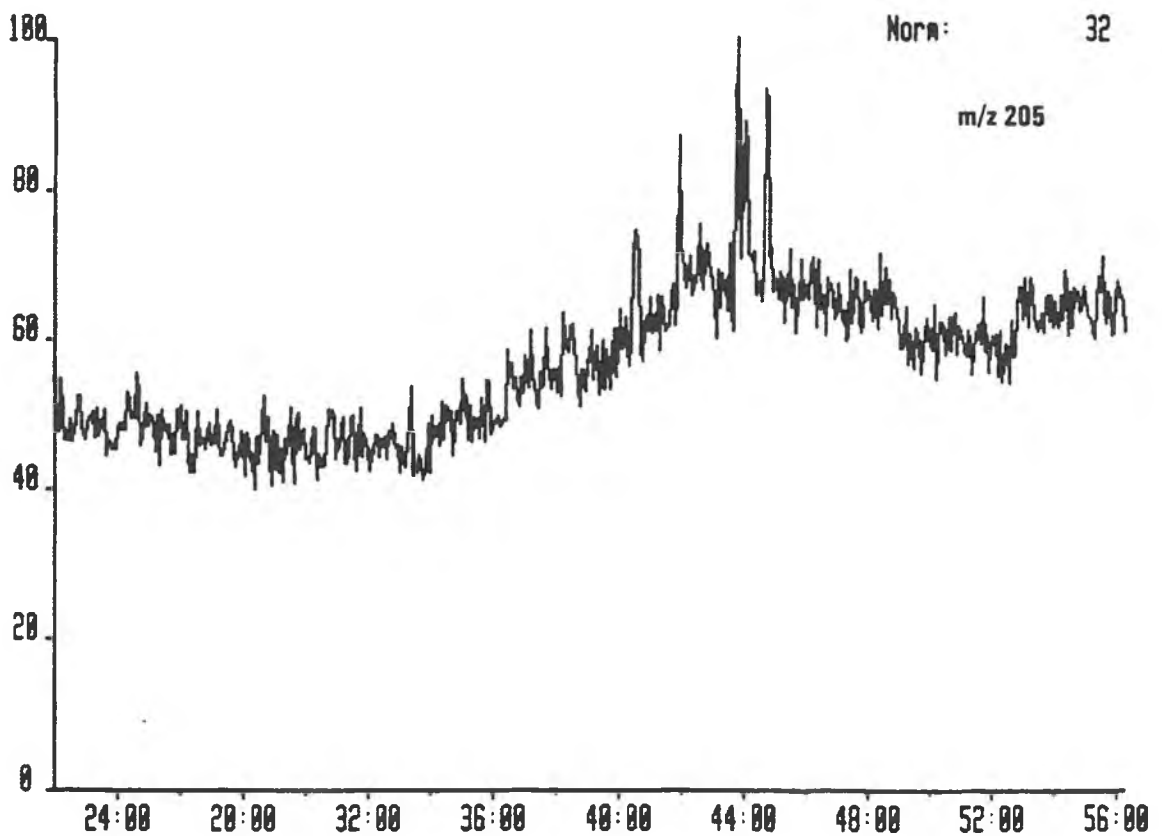
C5840SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU
 Sample 1 Injection 1 Group 1 Mass 191.1000
 Text:

System:TRIT



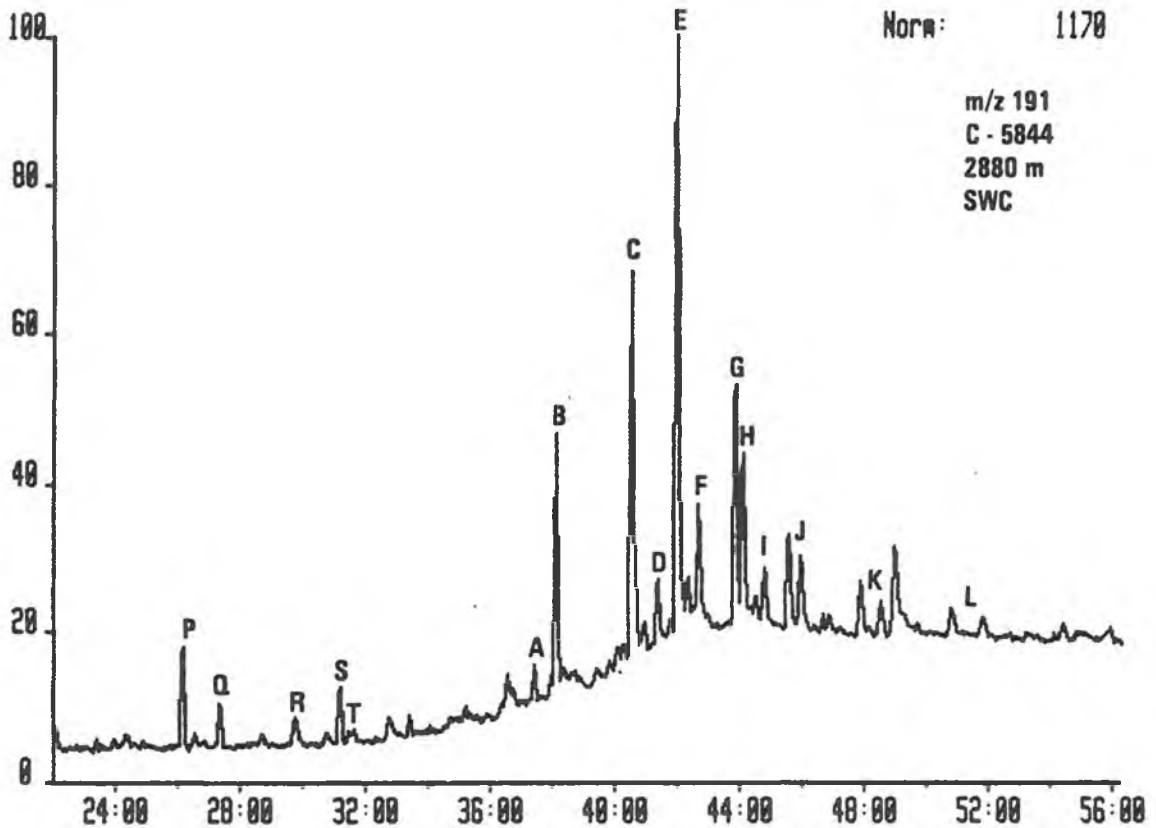
C5840SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU
 Sample 1 Injection 1 Group 1 Mass 205.1000
 Text:

System:TRIT



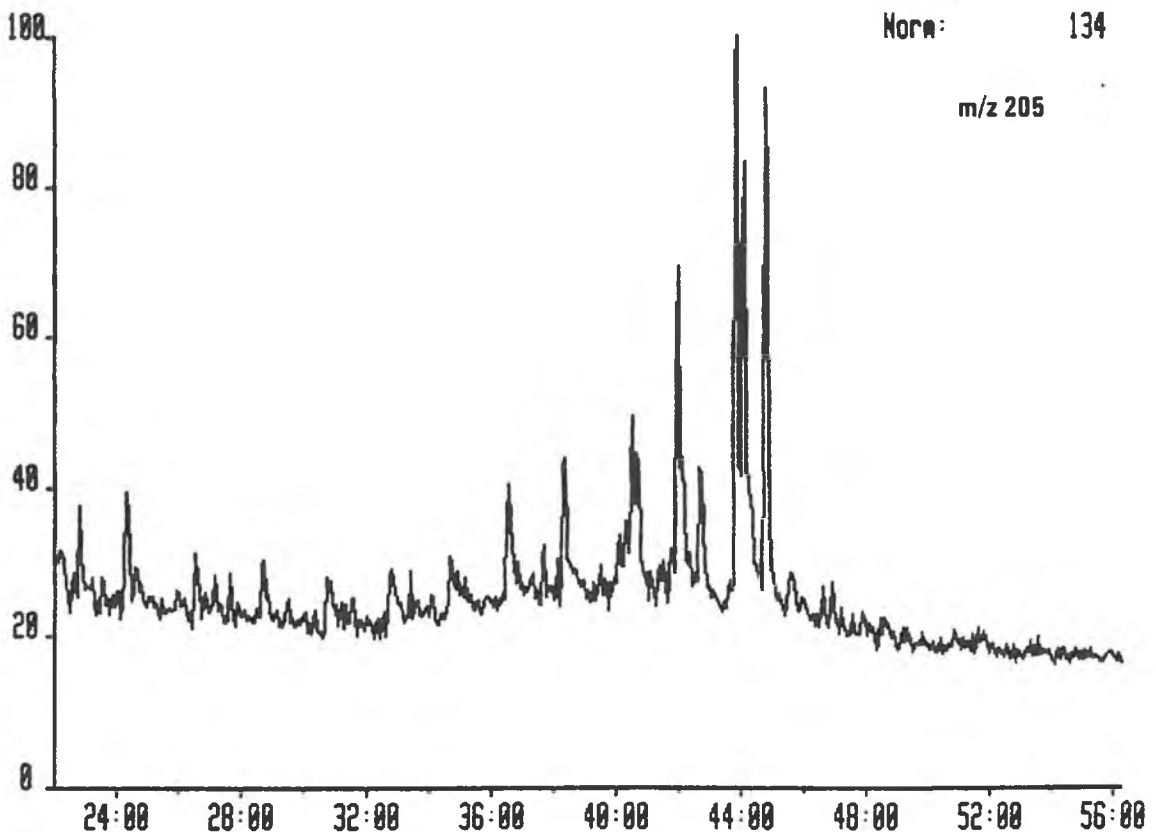
C5844SAT 6-FEB-87 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 191.1000
Text:

System:TRIT



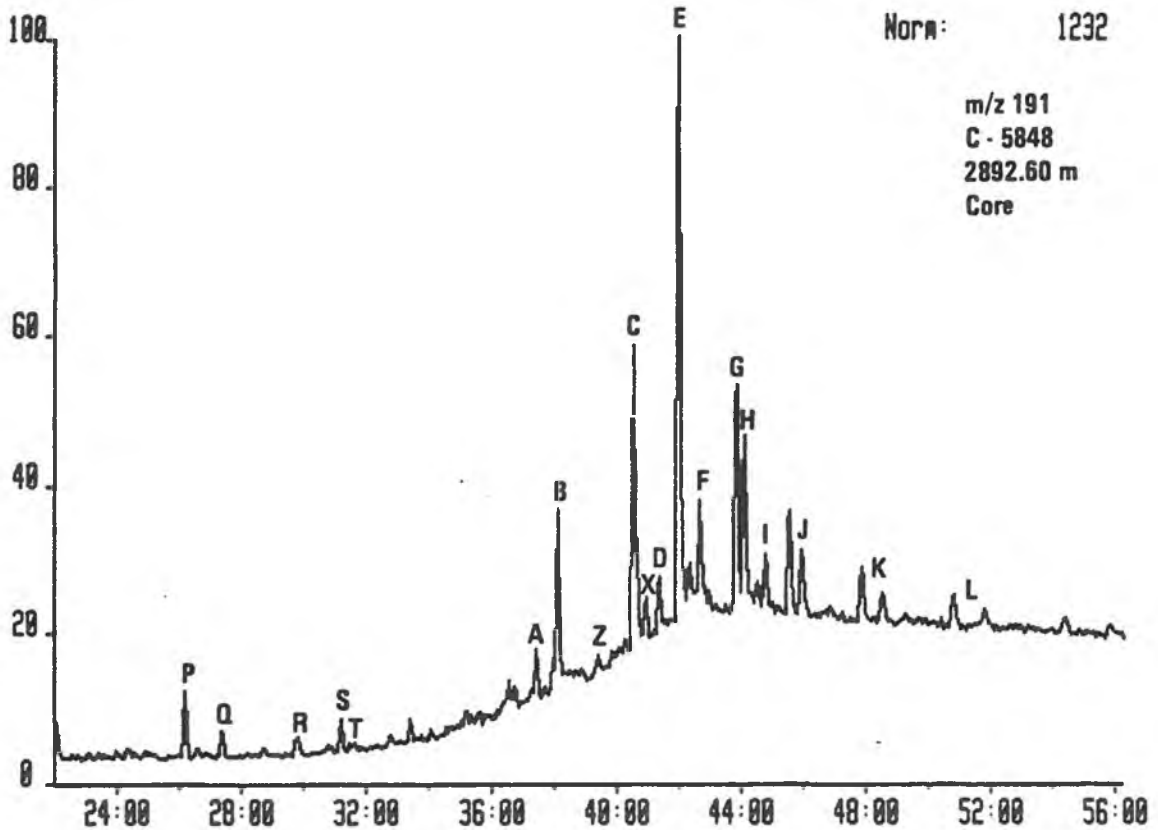
C5844SAT 6-FEB-87 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 205.1000
Text:

System:TRIT



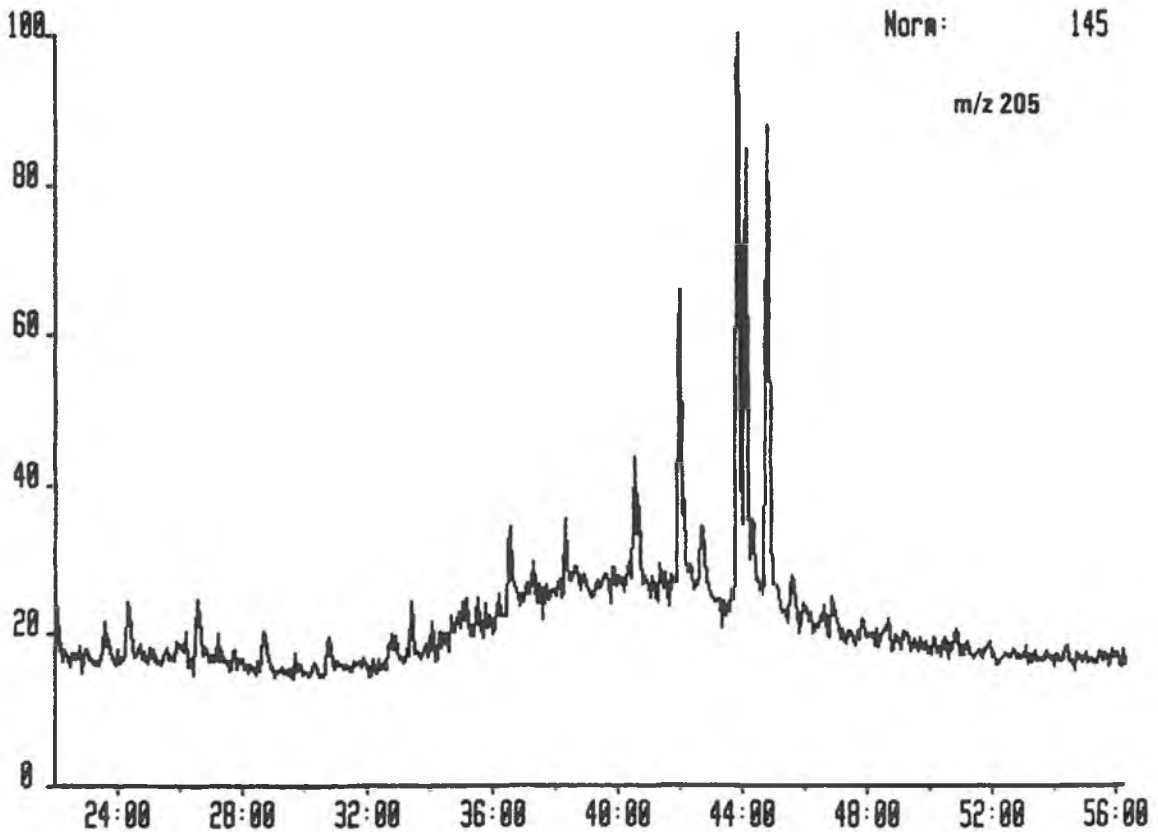
C5848SAT 5-FEB-07 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 191.1000
Text:

System:TRIT

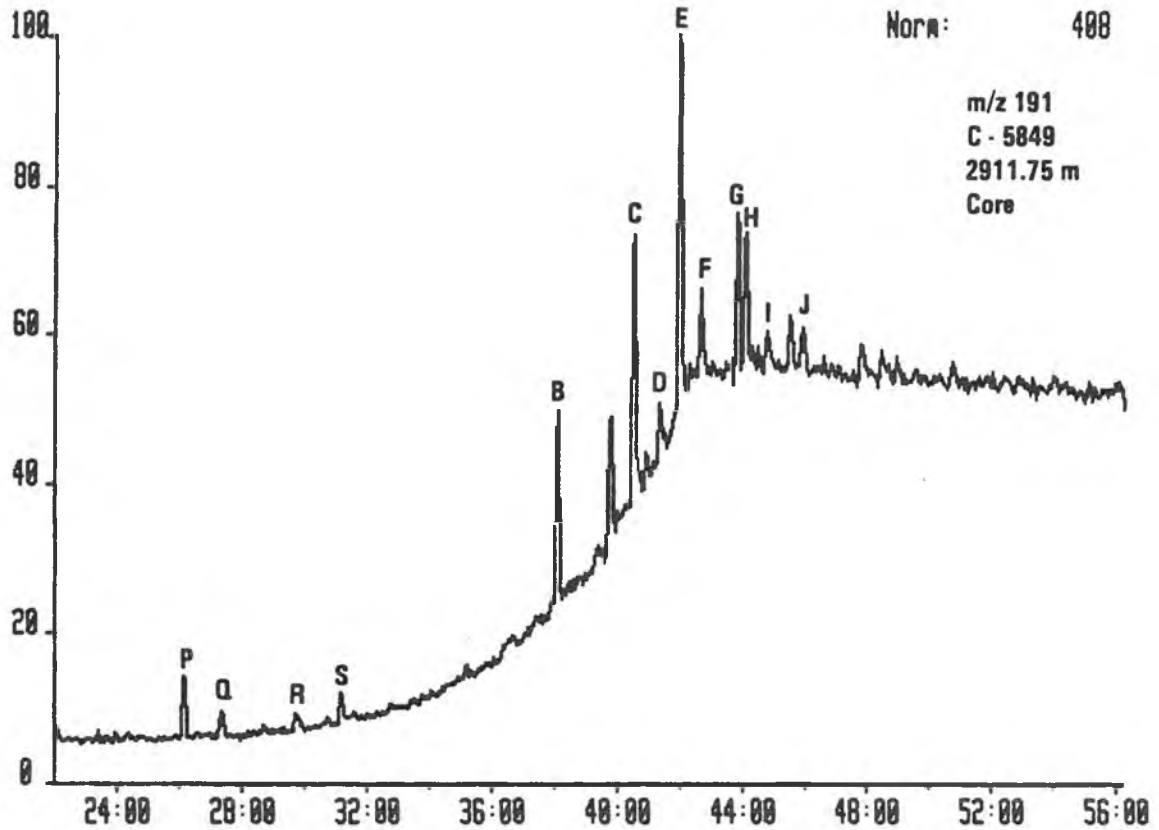


C5848SAT 5-FEB-07 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 205.1000
Text:

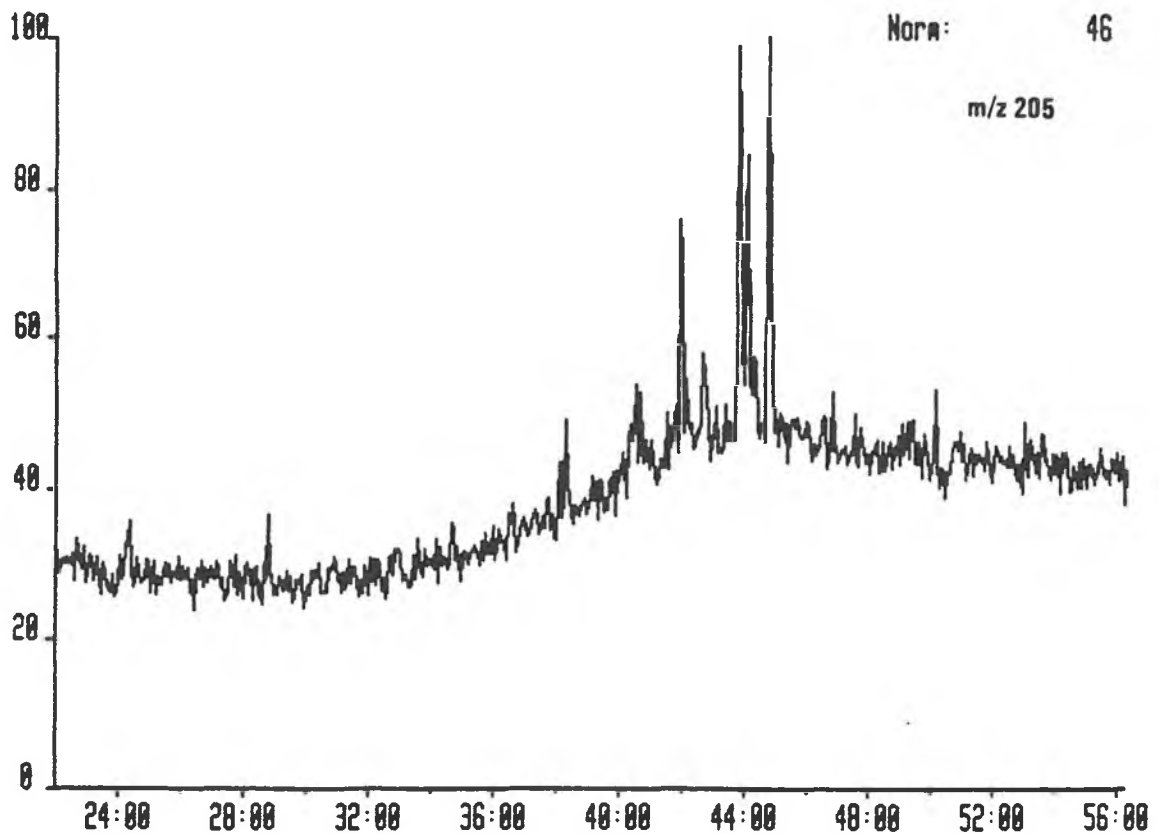
System:TRIT



C5849SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU System:TRIT
 Sample 1 Injection 1 Group 1 Mass 191.1000
 Text:

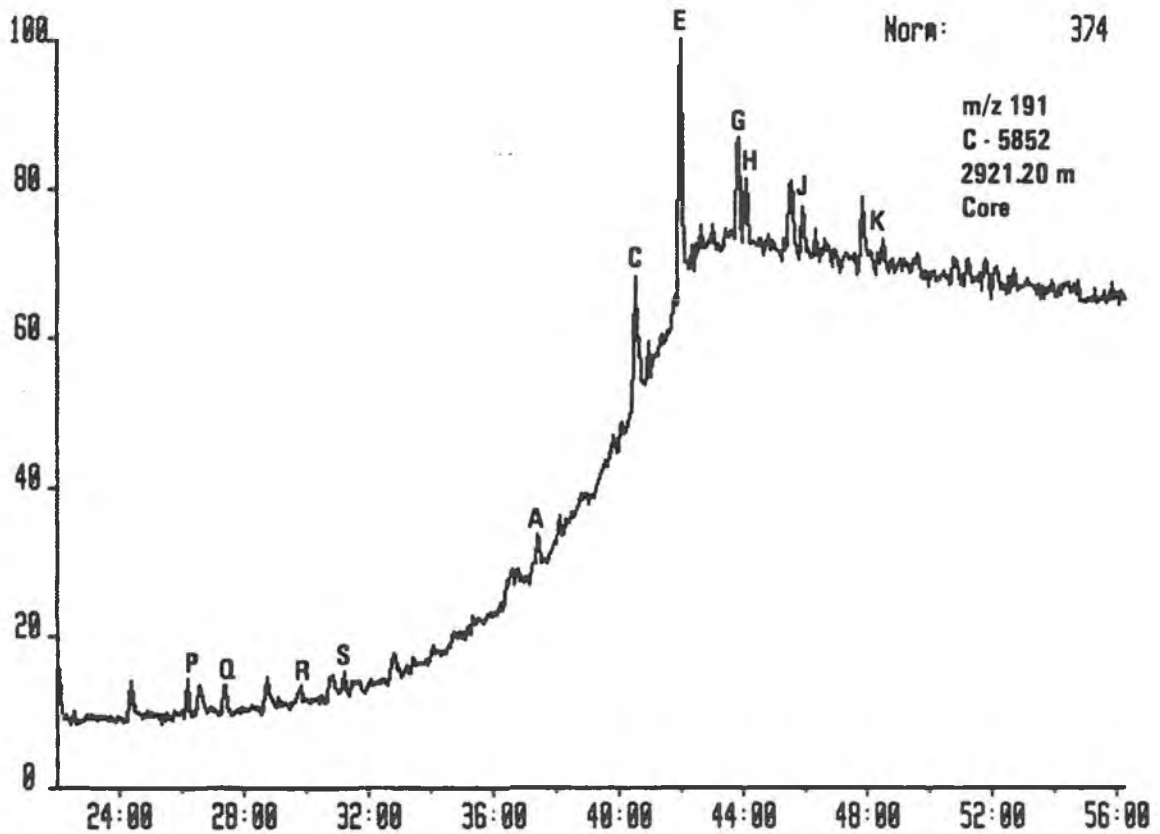


C5849SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU System:TRIT
 Sample 1 Injection 1 Group 1 Mass 205.1000
 Text:



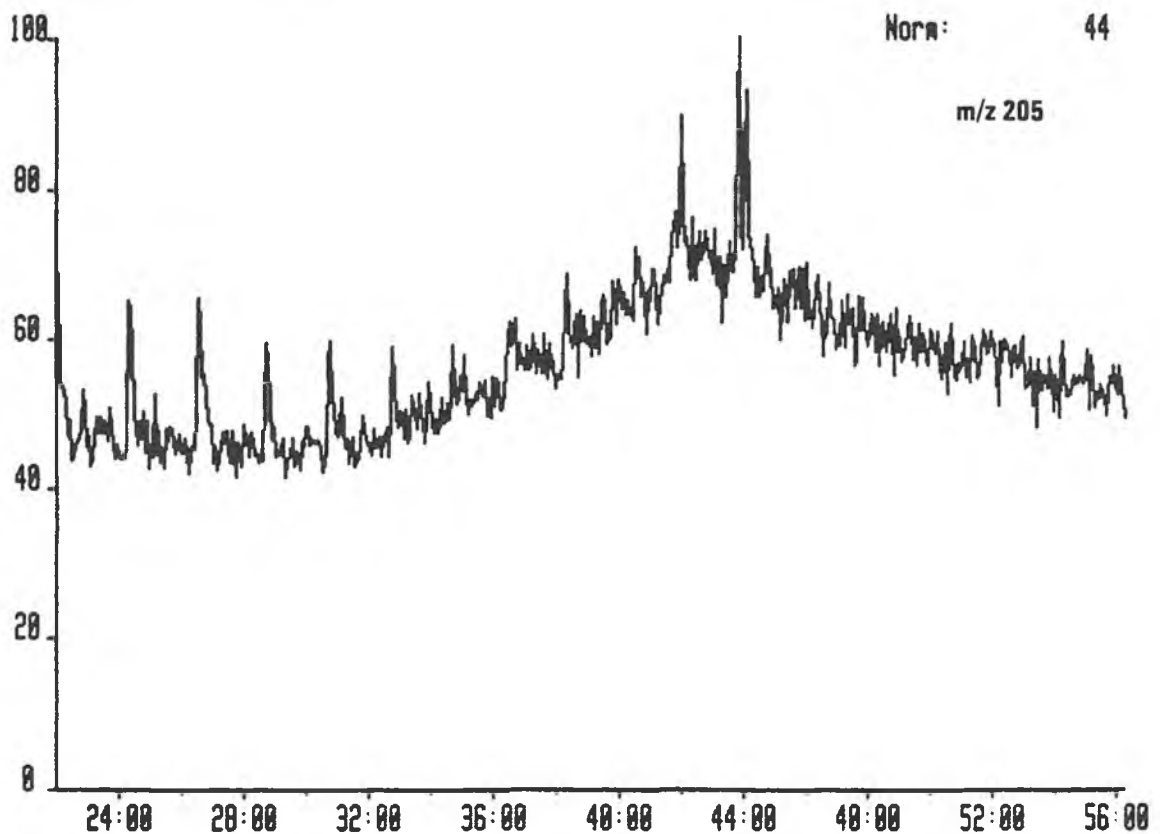
C5852SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU
 Sample 1 Injection 1 Group 1 Mass 191.1000
 Text:

System:TRIT

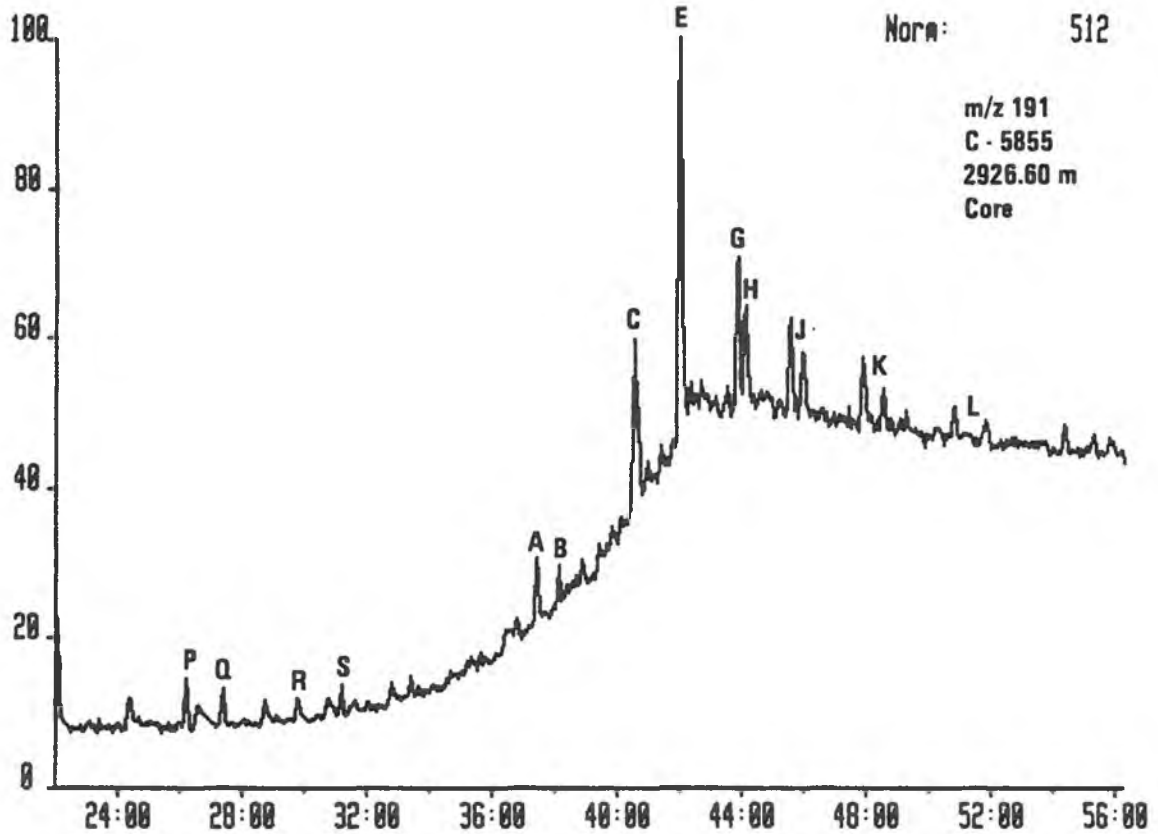


C5852SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU
 Sample 1 Injection 1 Group 1 Mass 205.1000
 Text:

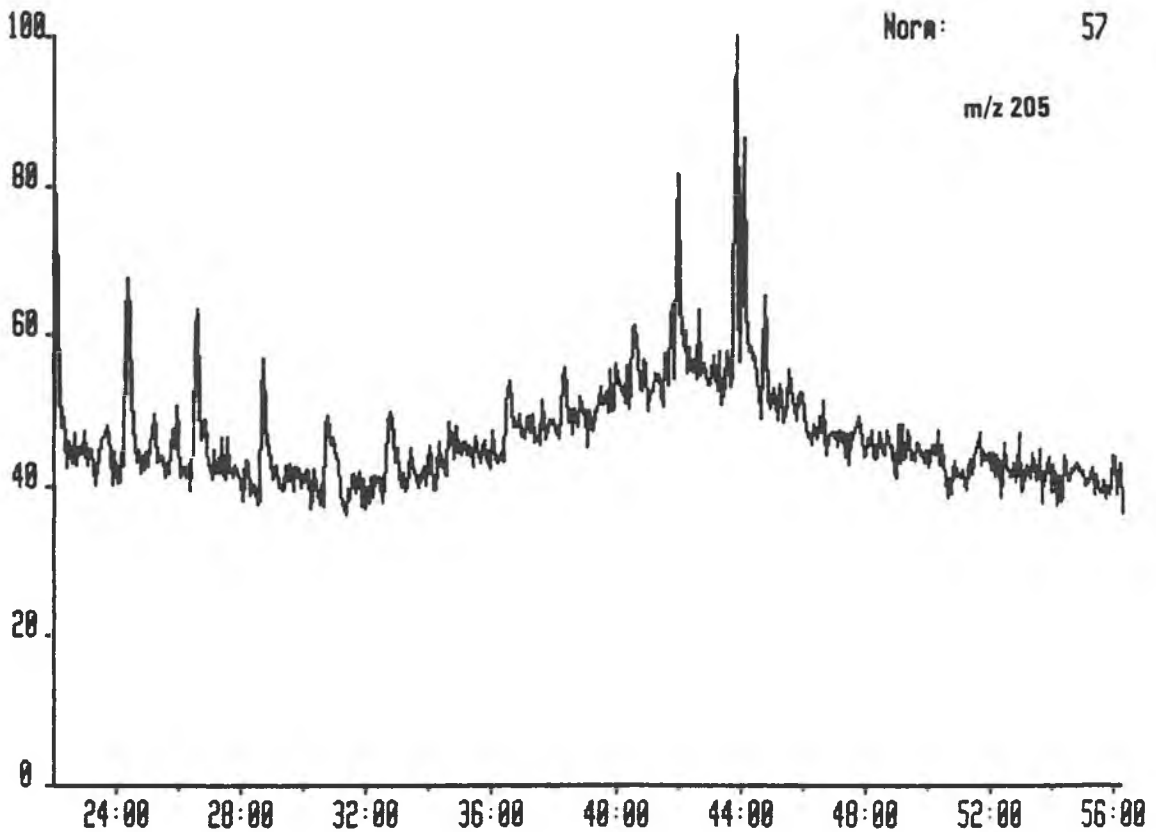
System:TRIT



C5855SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU System:TRIT
Sample 1 Injection 1 Group 1 Mass 191.1000
Text:

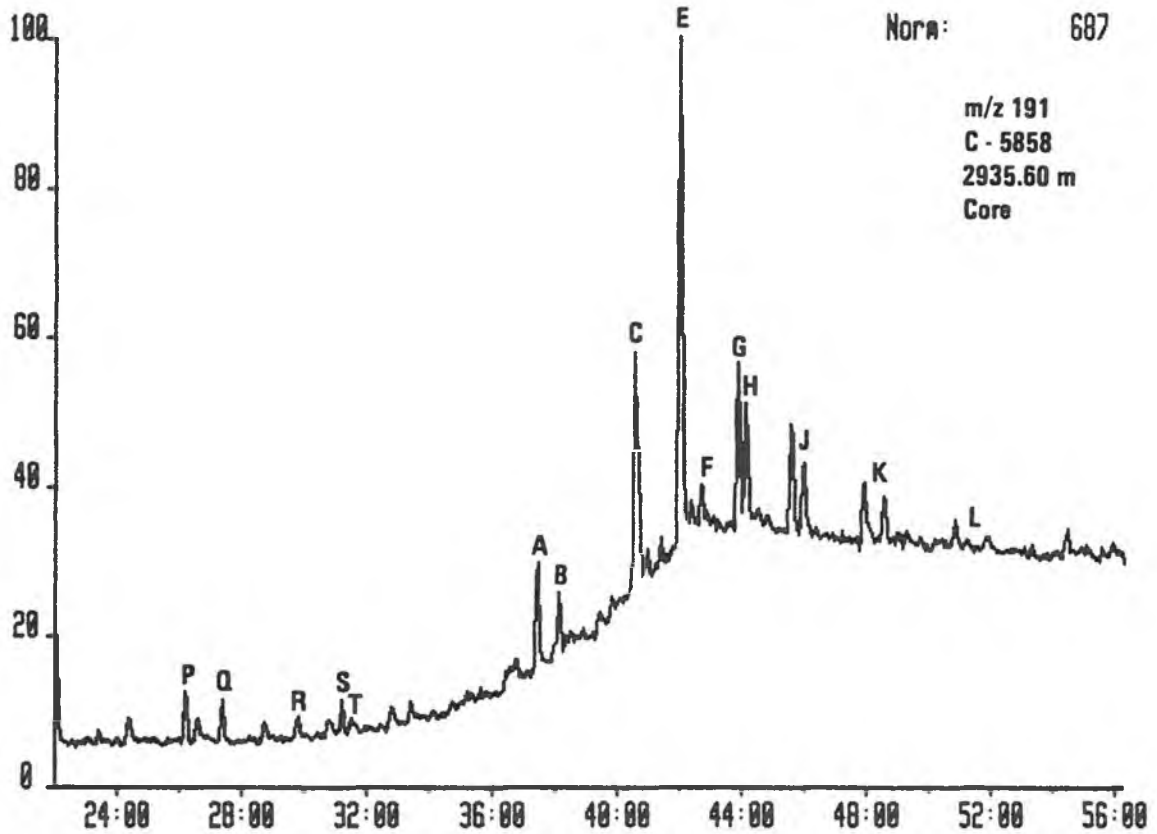


C5855SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU System:TRIT
Sample 1 Injection 1 Group 1 Mass 205.1000
Text:



C5858SAT 6-FEB-07 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 191.1000
Text:

System:TRIT

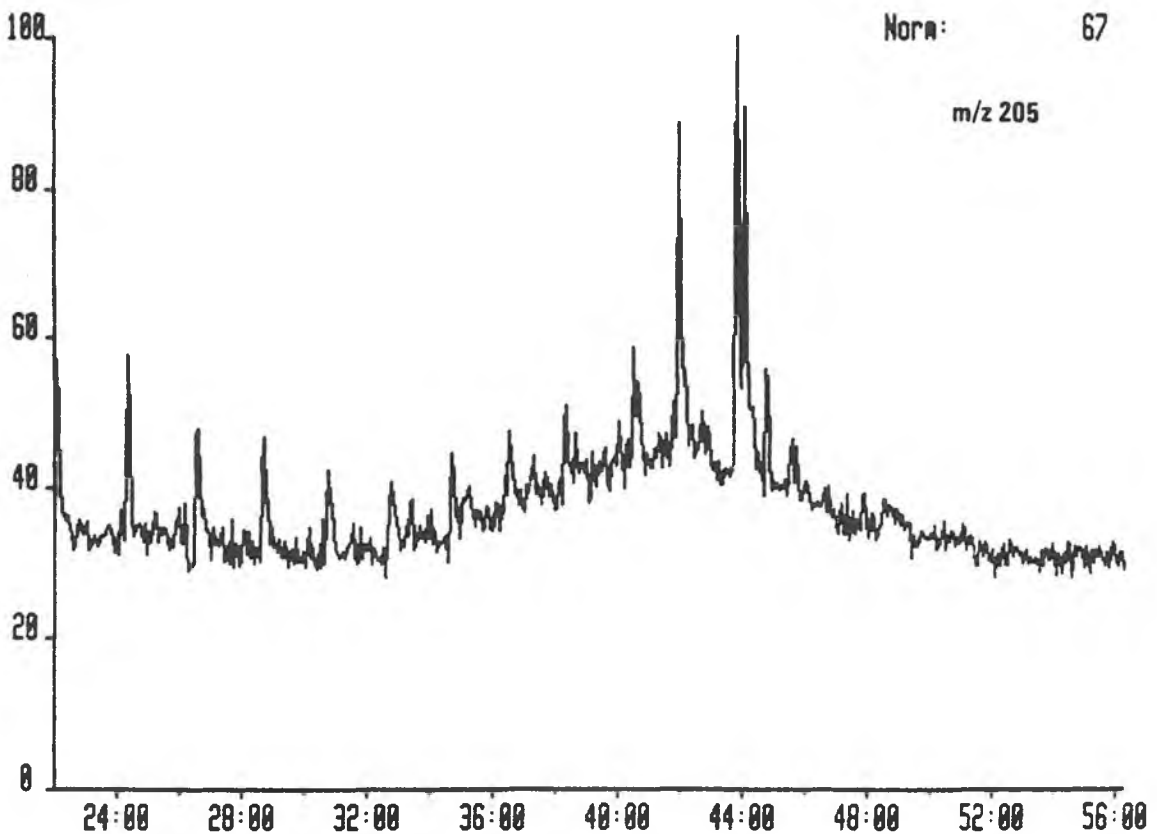


Norm: 687

m/z 191
C - 5858
2935.60 m
Core

C5858SAT 6-FEB-07 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 205.1000
Text:

System:TRIT

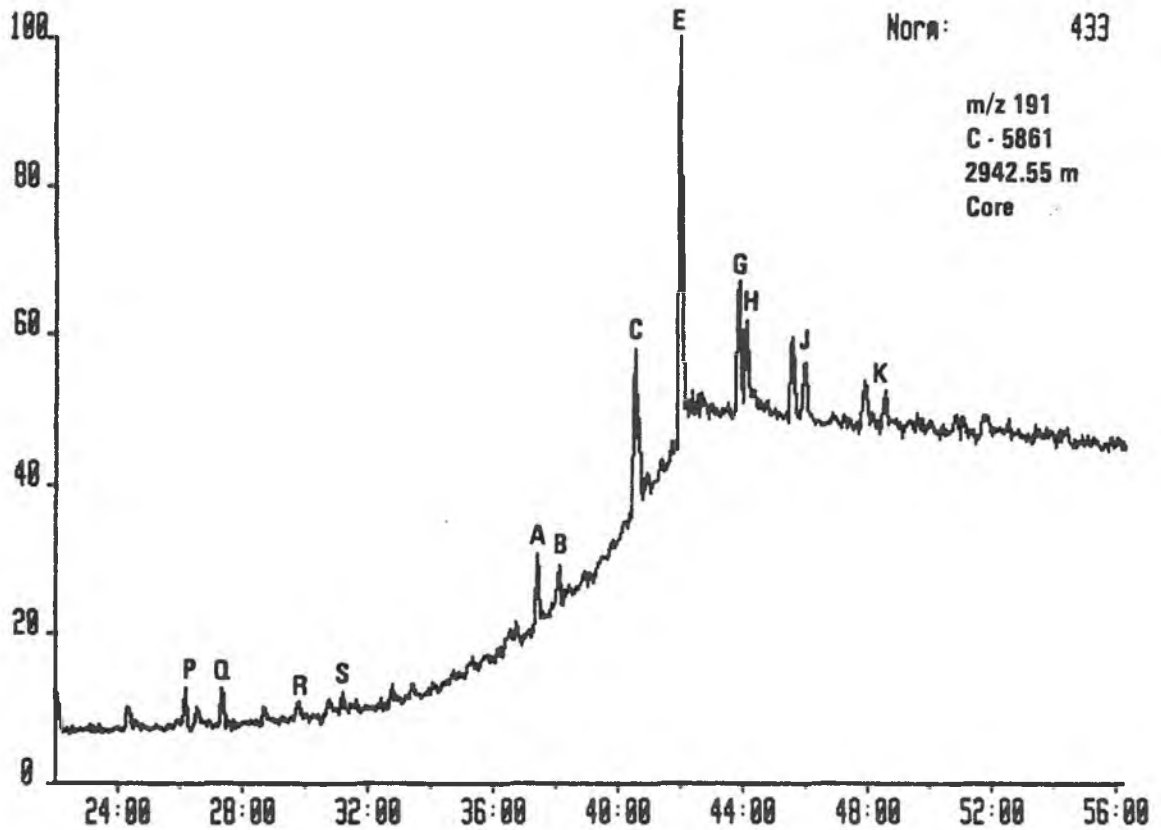


Norm: 67

m/z 205

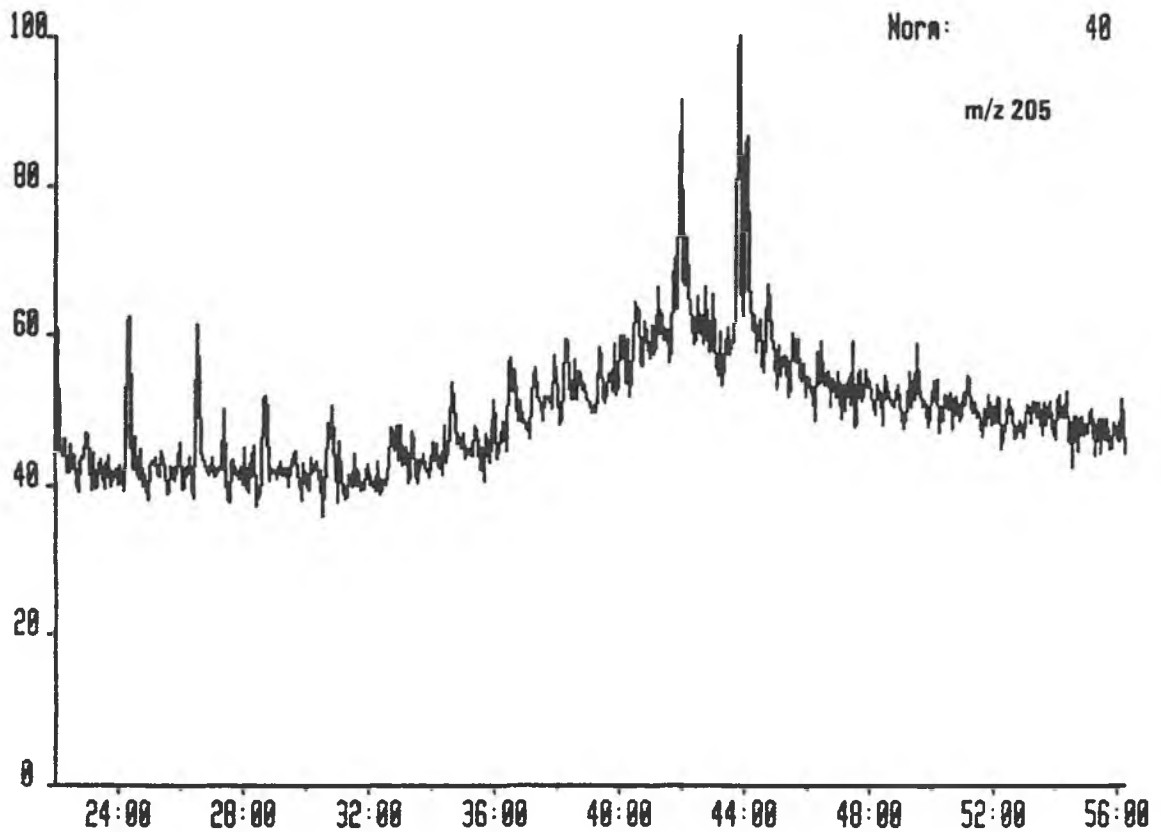
C5061SAT 6-FEB-07 Sir:Voltage 7070H Acnt:IKU
 Sample 1 Injection 1 Group 1 Mass 191.1000
 Text:

System:TRIT



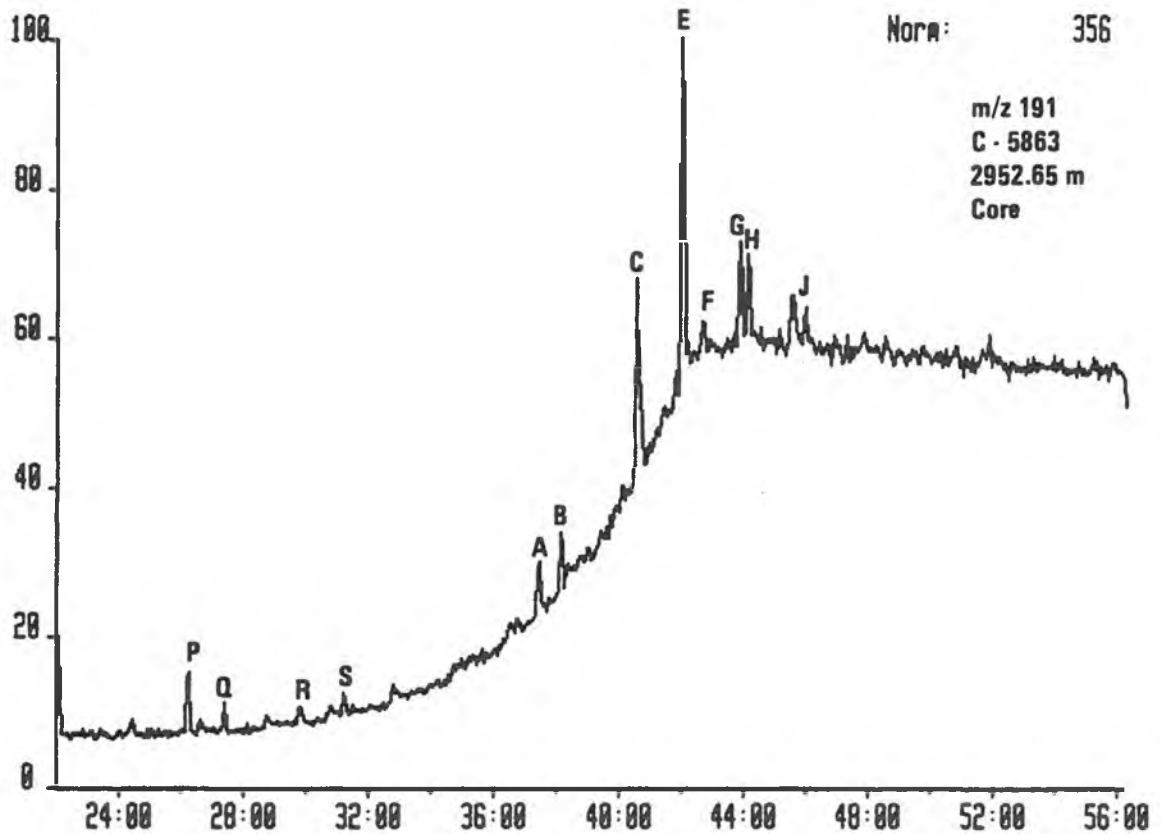
C5061SAT 6-FEB-07 Sir:Voltage 7070H Acnt:IKU
 Sample 1 Injection 1 Group 1 Mass 205.1000
 Text:

System:TRIT



C5063SAT2 6-FEB-07 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 191.1000
Text:

System:TRIT



C5063SAT2 6-FEB-07 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 205.1000
Text:

System:TRIT

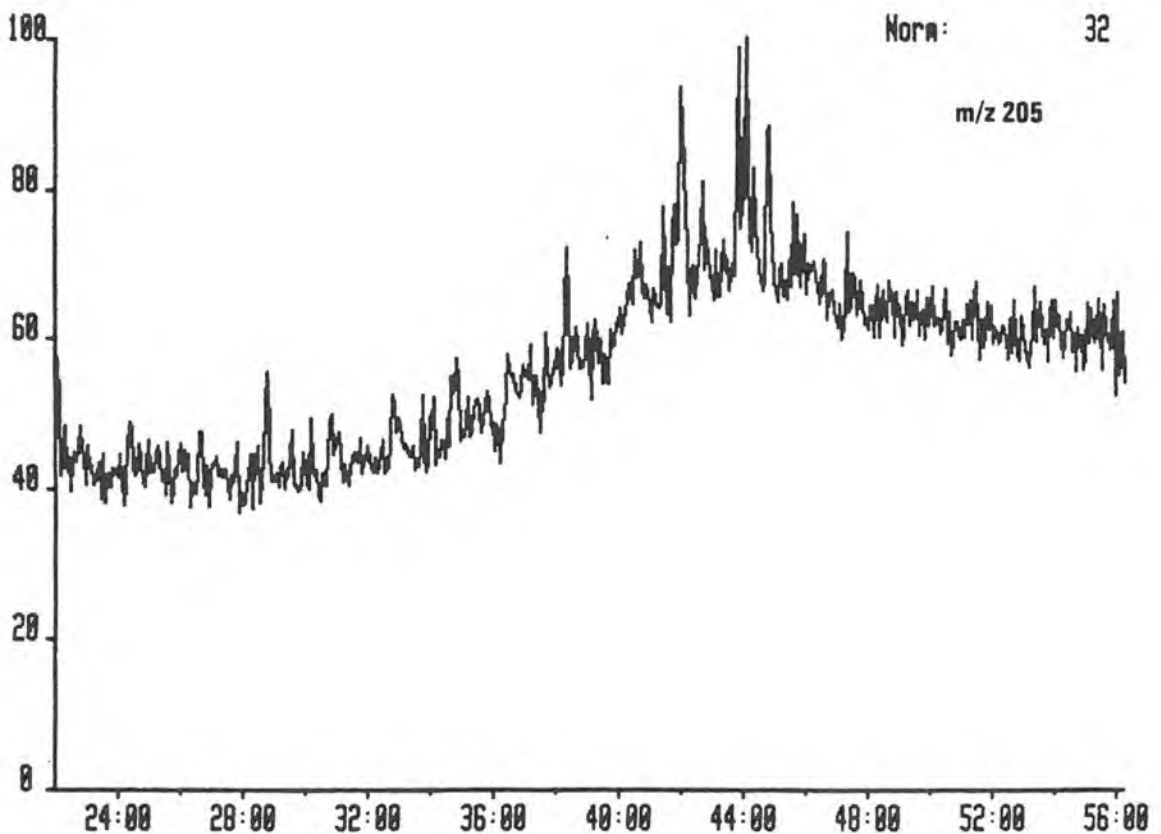
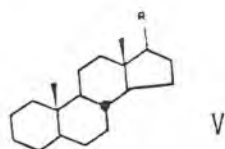
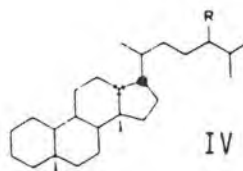
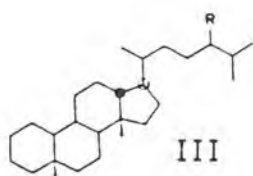
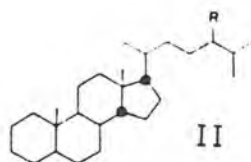
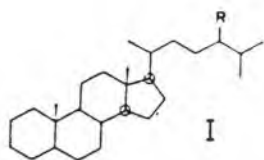


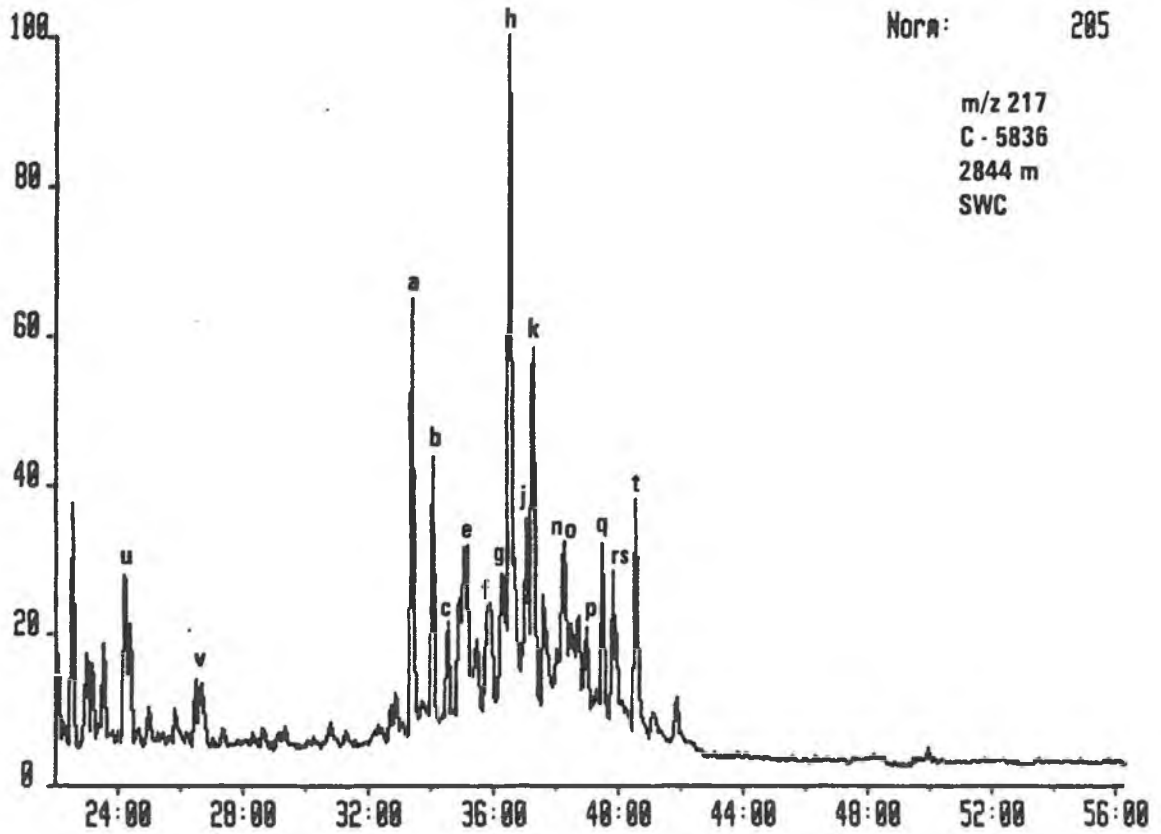
Figure 5.

Mass chromatograms representing steranes (m/z 217 and 218)

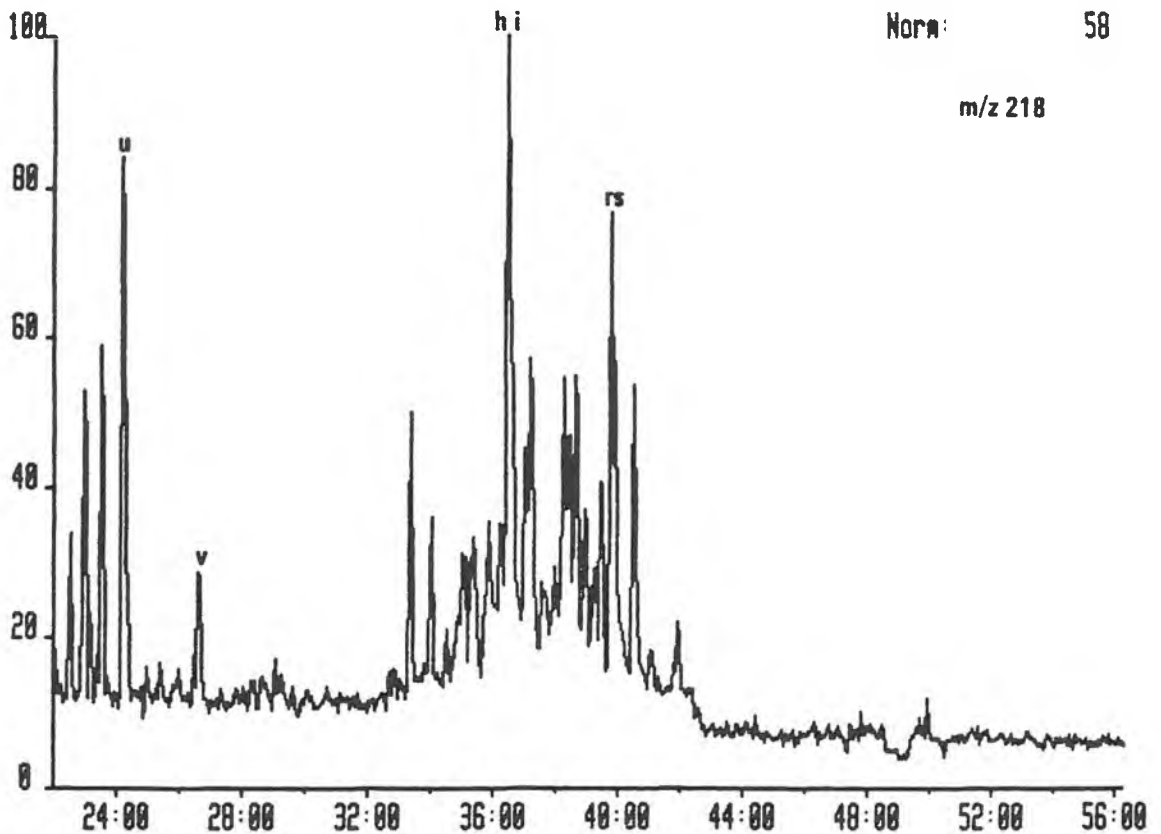
a	13 β (H),17 α (H)-diasterane (20S)	C ₂₇ H ₄₈	(III, R=H)
b	13 β (H),17 α (H)-diasterane (20R)	C ₂₇ H ₄₈	(III, R=H)
c	13 α (H),17 β (H)-diasterane (20S)	C ₂₇ H ₄₈	(IV, R=H)
d	13 α (H),17 β (H)-diasterane (20R)	C ₂₇ H ₄₈	(IV, R=H)
e	13 β (H),17 α (H)-diasterane (20S)	C ₂₈ H ₅₀	(III, R=CH ₃)
f	13 β (H),17 α (H)-diasterane (20R)	C ₂₈ H ₅₀	(III, R=CH ₃)
g	13 α (H),17 β (H)-diasterane (20S)	C ₂₈ H ₅₀	(IV, R=CH ₃)
	+ 14 α (H),17 α (H)-sterane (20S)	C ₂₇ H ₄₈	(I, R=H)
h	13 β (H),17 α (H)-diasterane (20S)	C ₂₉ H ₅₂	(III, R=C ₂ H ₅)
	+ 14 β (H),17 β (H)-sterane (20R)	C ₂₇ H ₄₈	(II, R=H)
i	14 β (H),17 β (H)-sterane (20S)	C ₂₇ H ₄₈	(II, R=H)
	+ 13 α (H),17 β (H)-diasterane (20R)	C ₂₈ H ₅₀	(IV, R=CH ₃)
j	14 α (H),17 α (H)-sterane (20R)	C ₂₇ H ₄₈	(I, R=H)
k	13 β (H),17 α (H)-diasterane (20R)	C ₂₉ H ₅₂	(III, R=C ₂ H ₅)
l	13 α (H),17 β (H)-diasterane (20S)	C ₂₉ H ₅₂	(III, R=C ₂ H ₅)
m	14 α (H),17 α (H)-sterane (20S)	C ₂₈ H ₅₀	(I, R=CH ₃)
n	13 α (H),17 β (H)-diasterane (20R)	C ₂₉ H ₅₂	(III, R=C ₂ H ₅)
	+ 14 β (H),17 β (H)-sterane (20R)	C ₂₈ H ₅₀	(II, R=CH ₃)
o	14 β (H),17 β (H)-sterane (20S)	C ₂₈ H ₅₀	(II, R=CH ₃)
p	14 α (H),17 α (H)-sterane (20R)	C ₂₈ H ₅₀	(I, R=CH ₃)
q	14 α (H),17 α (H)-sterane (20S)	C ₂₉ H ₅₂	(I, R=C ₂ H ₅)
r	14 β (H),17 β (H)-sterane (20R)	C ₂₉ H ₅₂	(II, R=C ₂ H ₅)
	+ unknown sterane		
s	14 β (H),17 β (H)-sterane (20S)	C ₂₉ H ₅₂	(II, R=C ₂ H ₅)
t	14 α (H),17 α (H)-sterane (20R)	C ₂₉ H ₅₂	(I, R=C ₂ H ₅)
u	5 α (H)-sterane	C ₂₁ H ₃₆	(V, R=C ₂ H ₅)
v	5 α (H)-sterane	C ₂₂ H ₃₈	(IV, R=C ₃ H ₇)



C5836SAT 5-FEB-07 Sir:Voltage 7070H Acnt:IKU System:TRIT
 Sample 1 Injection 1 Group 1 Mass 217.1000
 Text:

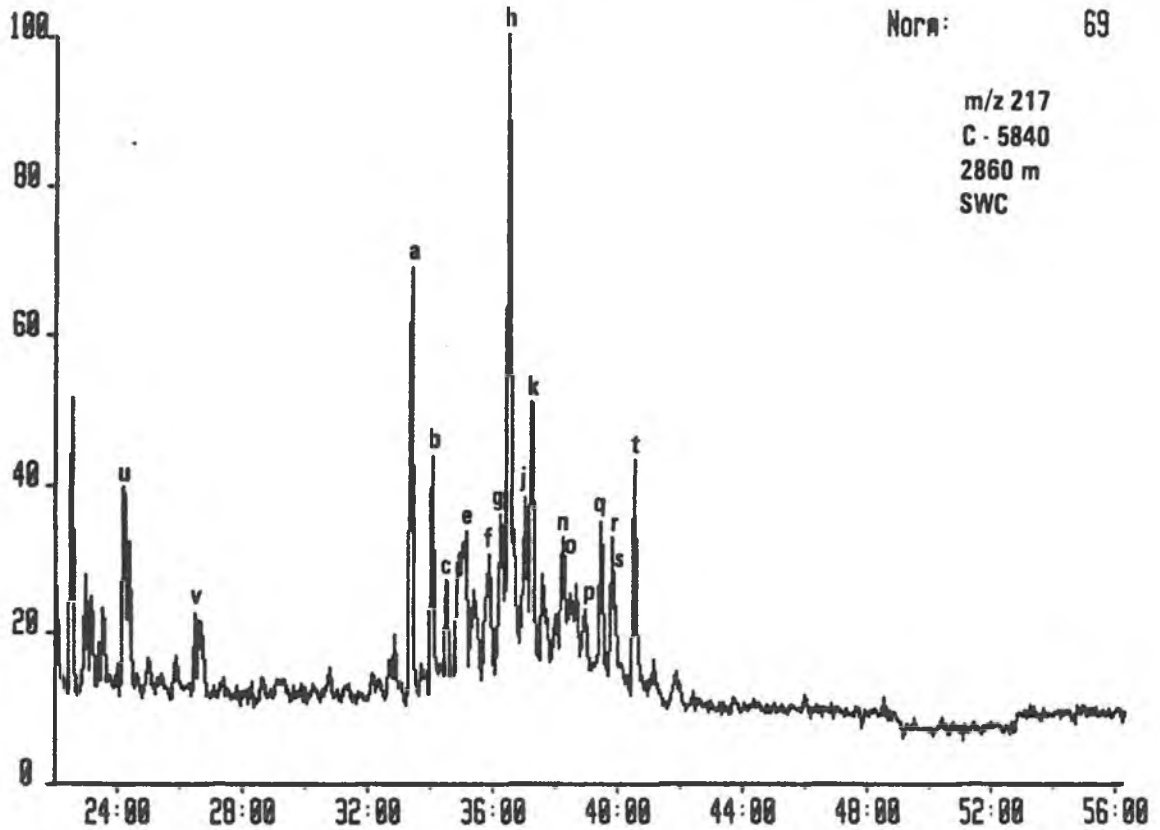


C5836SAT 5-FEB-07 Sir:Voltage 7070H Acnt:IKU System:TRIT
 Sample 1 Injection 1 Group 1 Mass 218.1000
 Text:



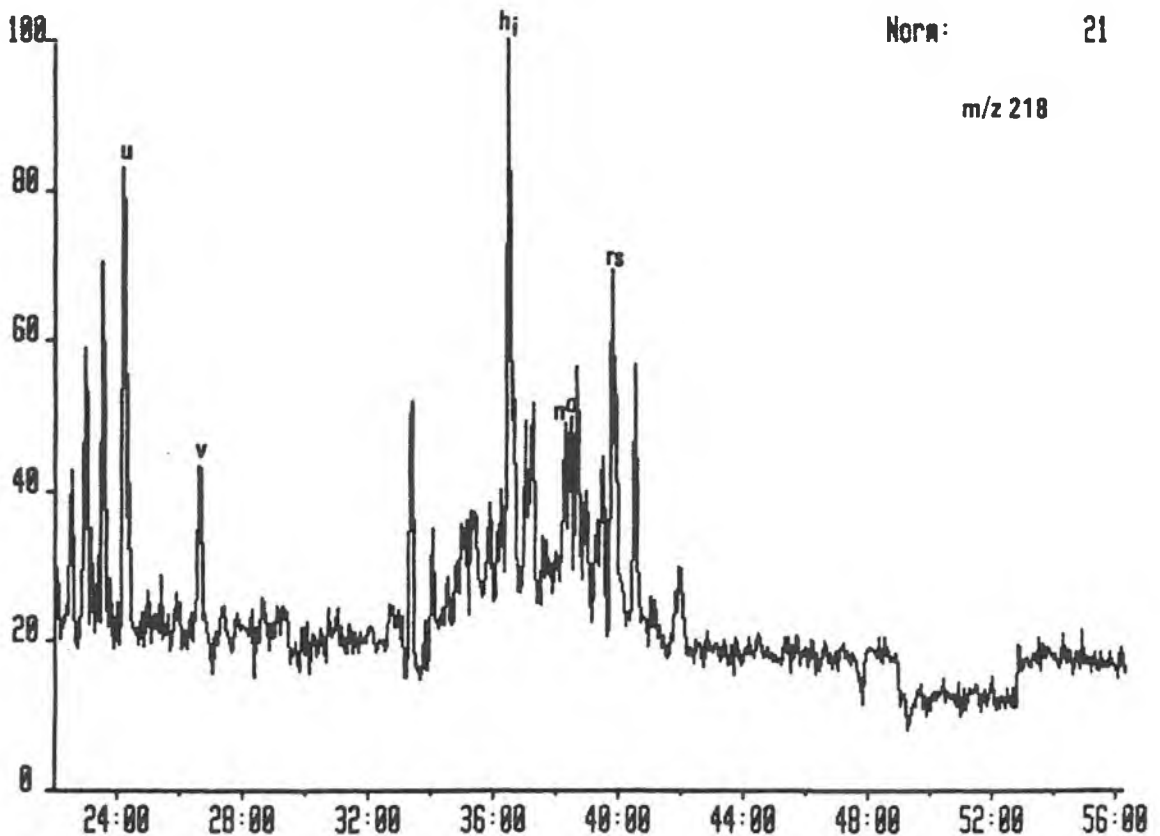
C5840SAT 5-FEB-07 Sir:Voltage 7070H Acnt:IKU
 Sample 1 Injection 1 Group 1 Mass 217.1000
 Text:

System:TRIT



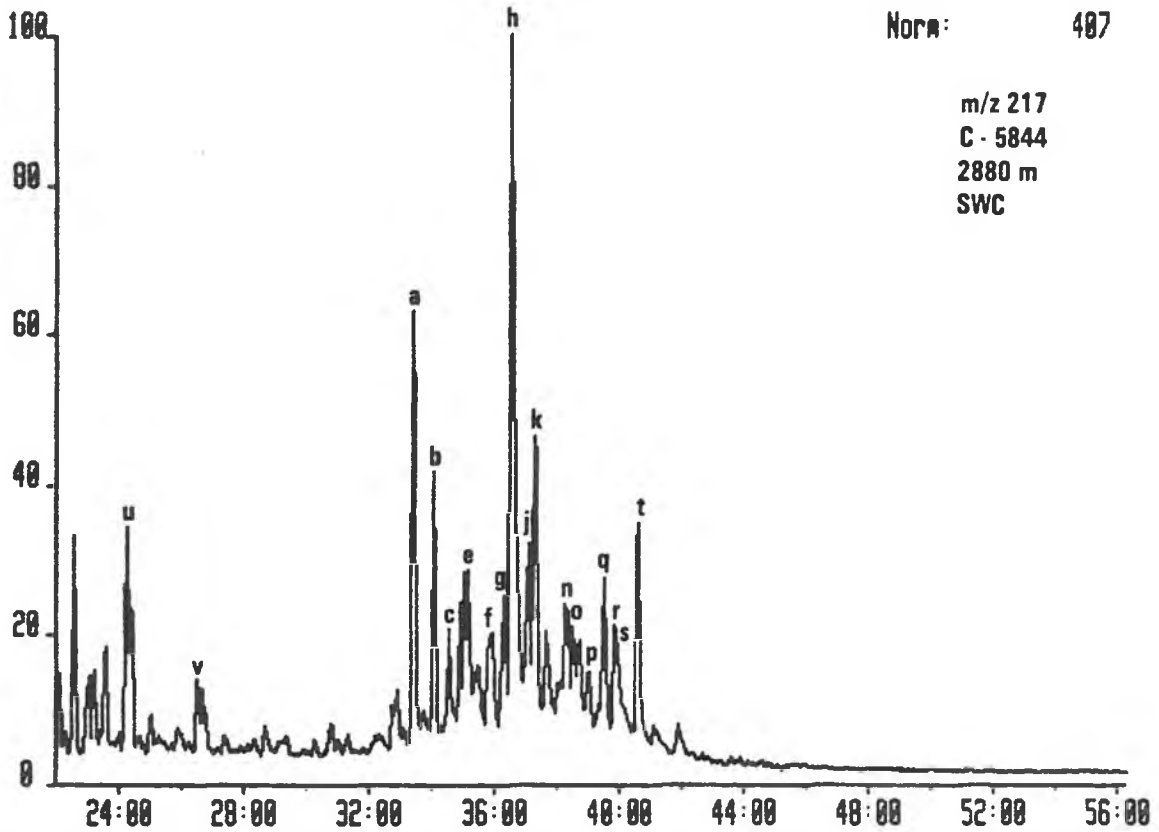
C5840SAT 5-FEB-07 Sir:Voltage 7070H Acnt:IKU
 Sample 1 Injection 1 Group 1 Mass 218.1000
 Text:

System:TRIT



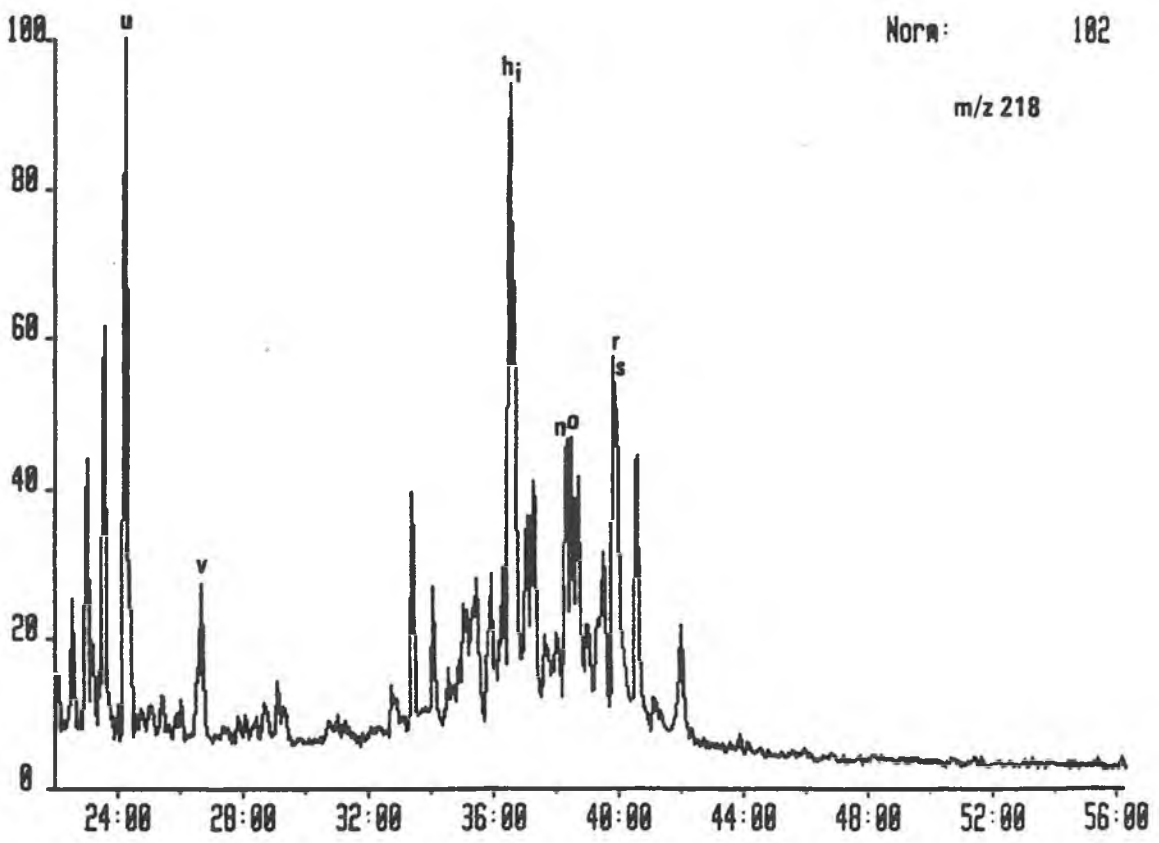
C5844SAT 6-FEB-87 Sir:Voltage 7070H Acnt:IKU
 Sample 1 Injection 1 Group 1 Mass 217.1000
 Text:

System:TRIT



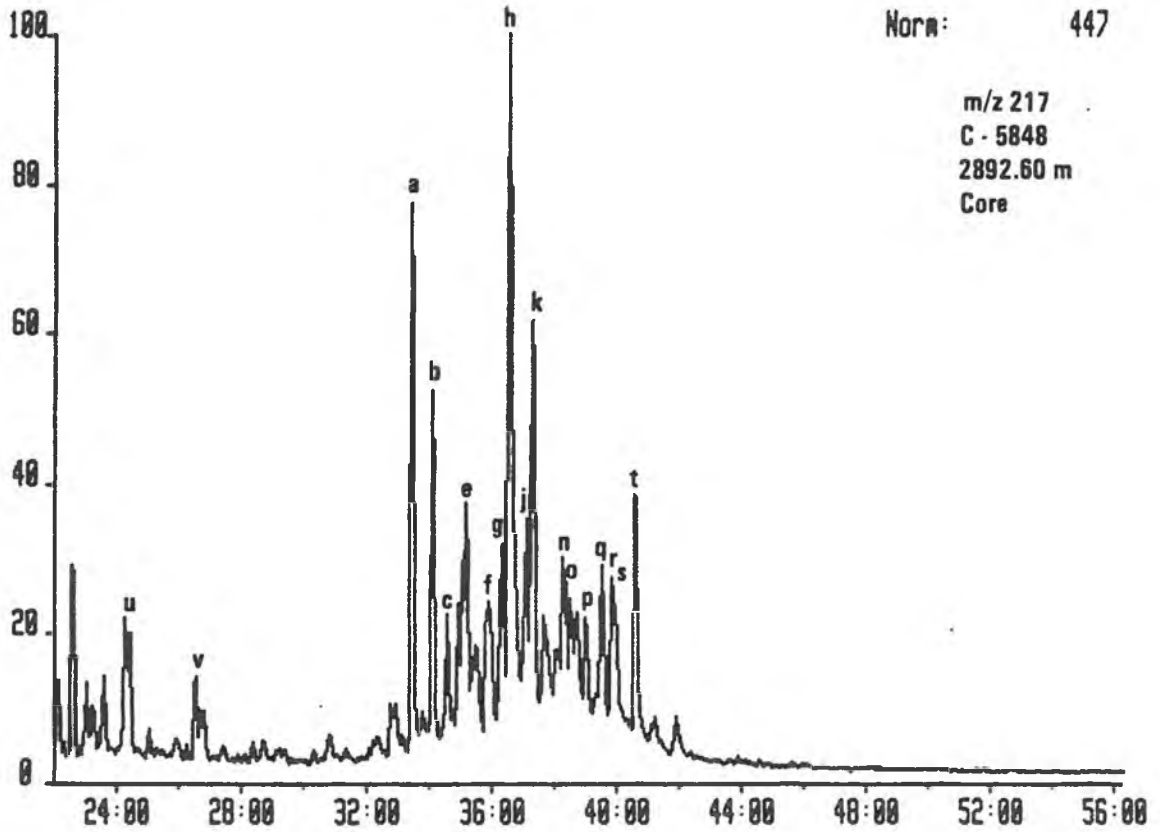
C5844SAT 6-FEB-87 Sir:Voltage 7070H Acnt:IKU
 Sample 1 Injection 1 Group 1 Mass 218.1000
 Text:

System:TRIT



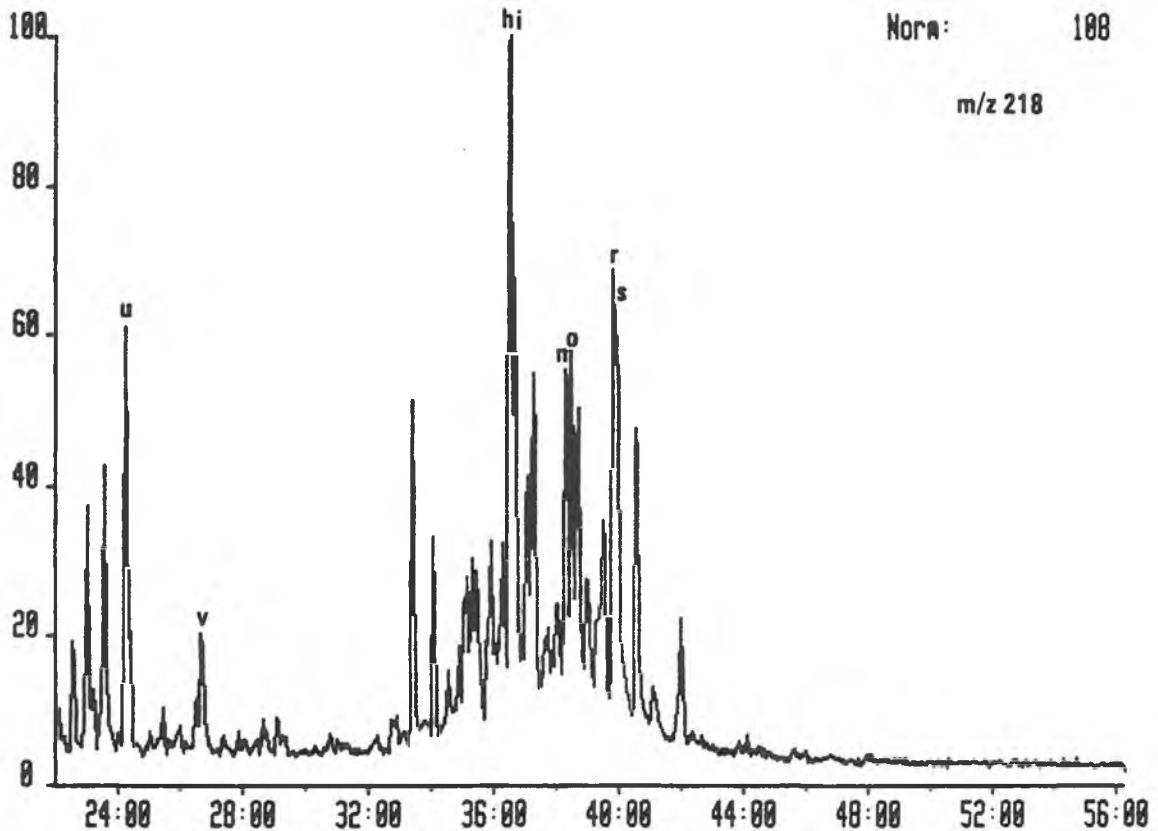
C5848SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU
 Sample 1 Injection 1 Group 1 Mass 217.1000
 Text:

System:TRIT



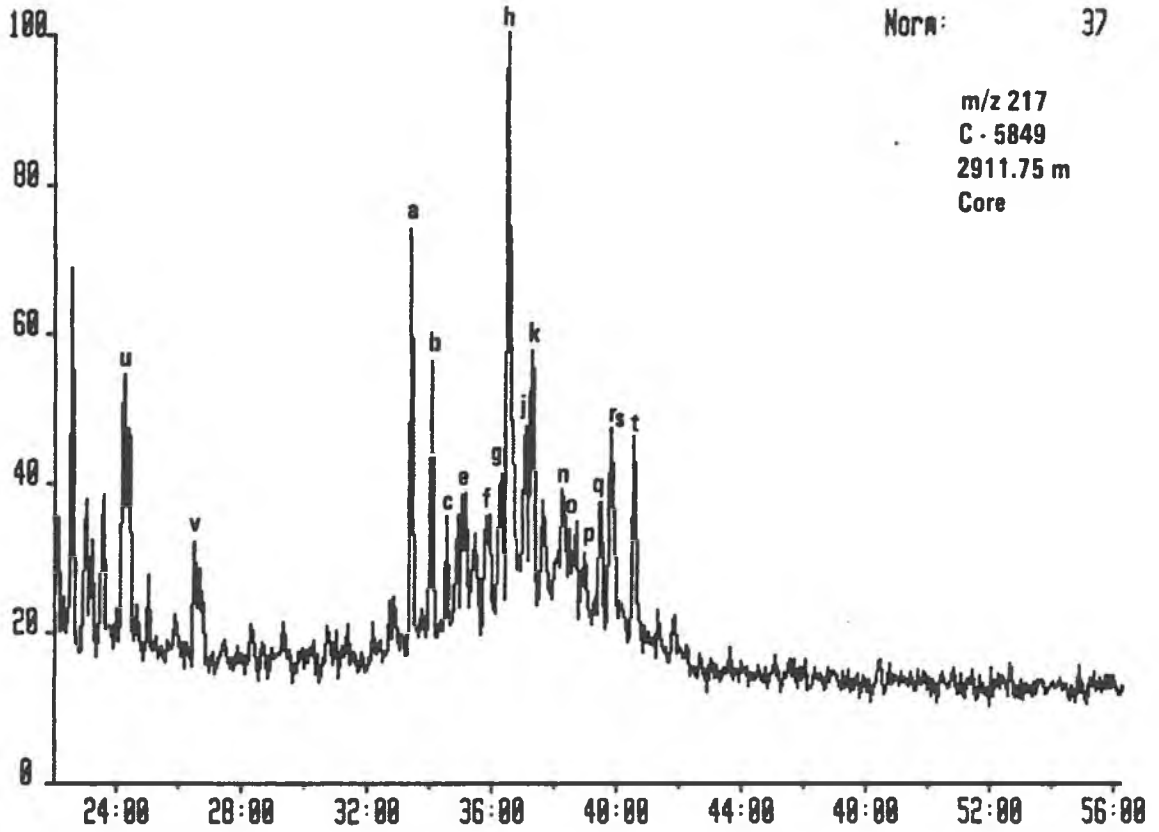
C5848SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU
 Sample 1 Injection 1 Group 1 Mass 218.1000
 Text:

System:TRIT



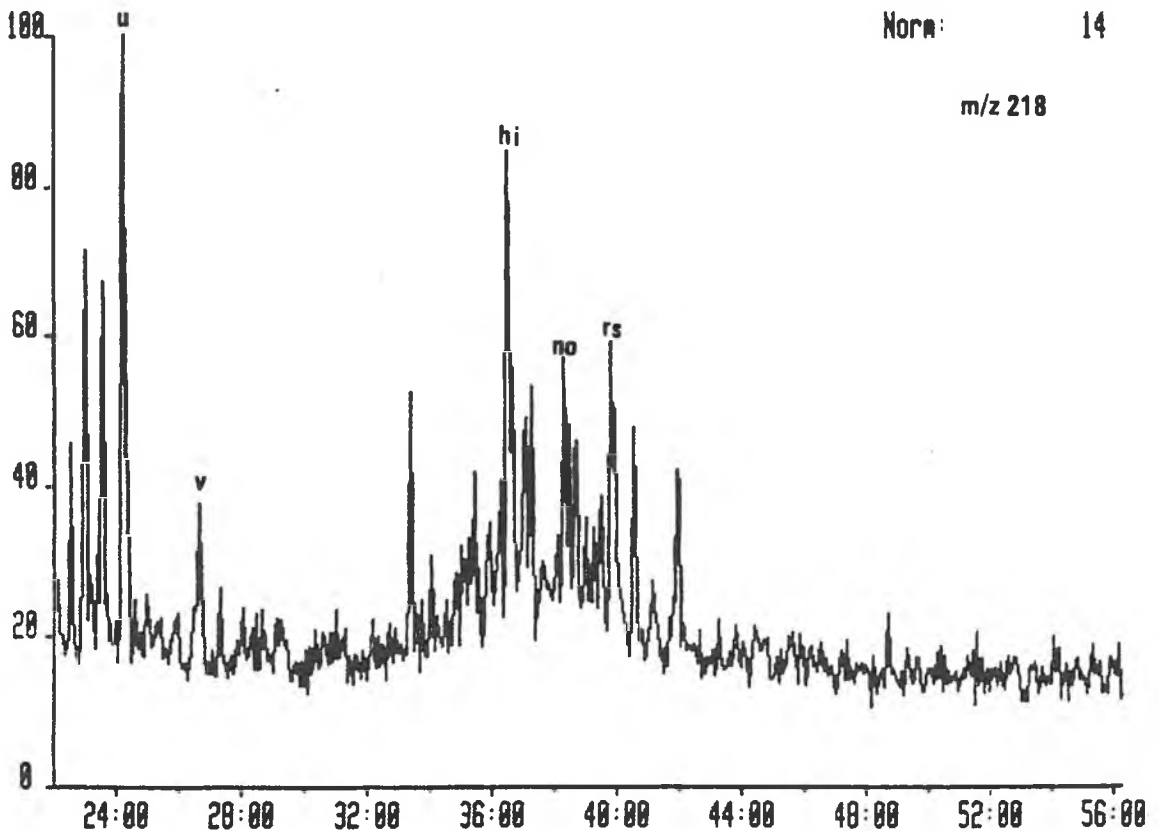
C5849SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU
 Sample 1 Injection 1 Group 1 Mass 217.1000
 Text:

System:TRIT



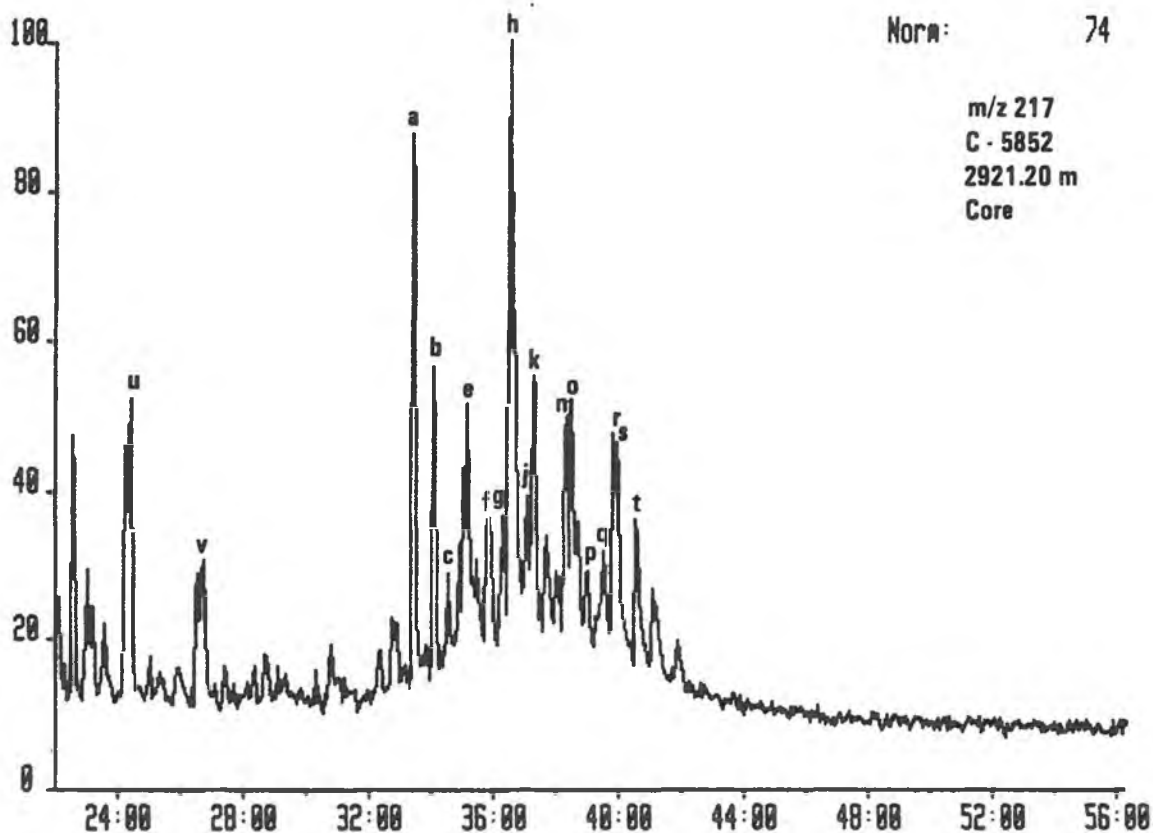
C5849SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU
 Sample 1 Injection 1 Group 1 Mass 218.1000
 Text:

System:TRIT



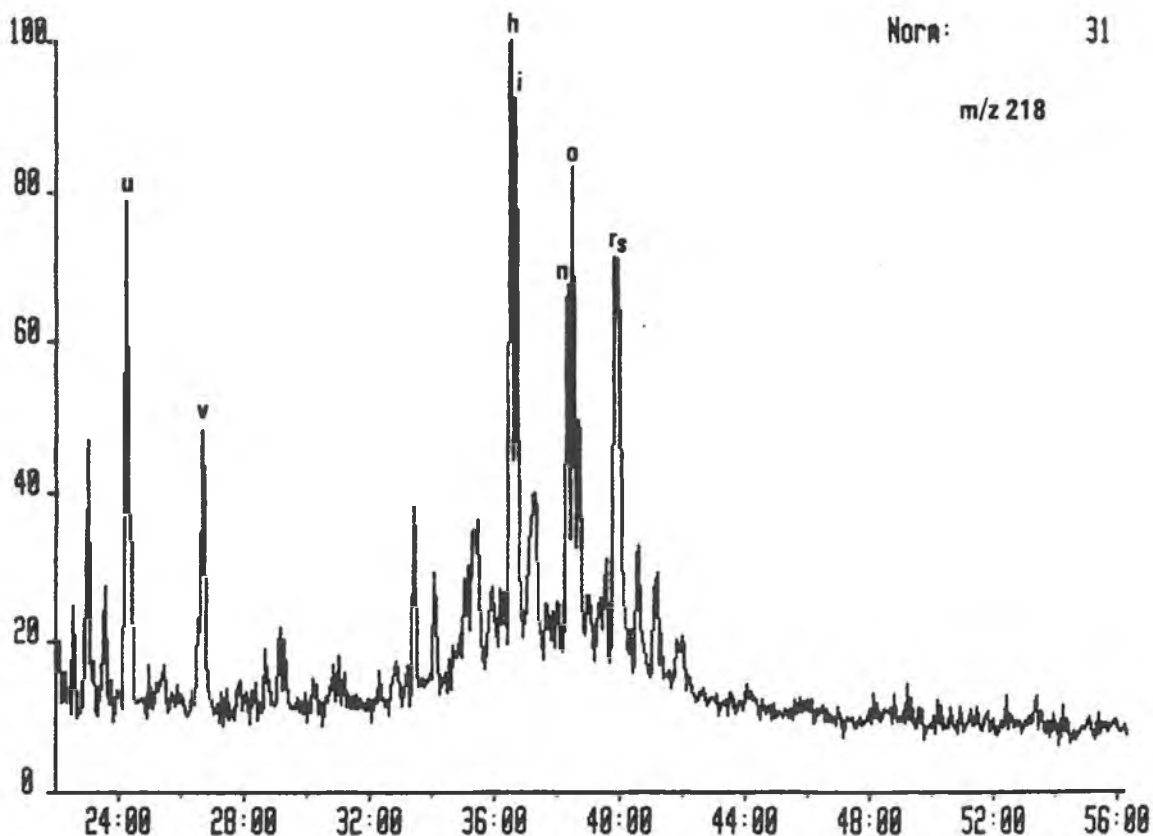
C5852SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU
 Sample 1 Injection 1 Group 1 Mass 217.1000
 Text:

System:TRIT

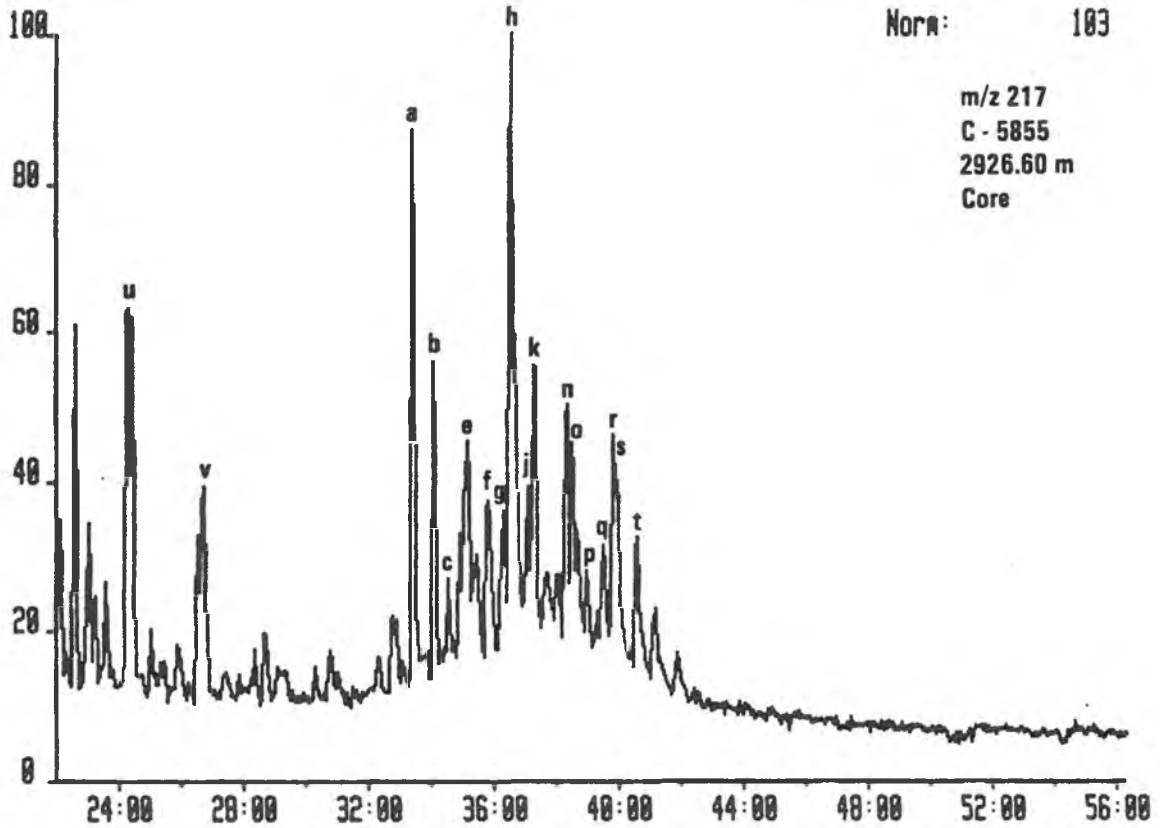


C5852SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU
 Sample 1 Injection 1 Group 1 Mass 218.1000
 Text:

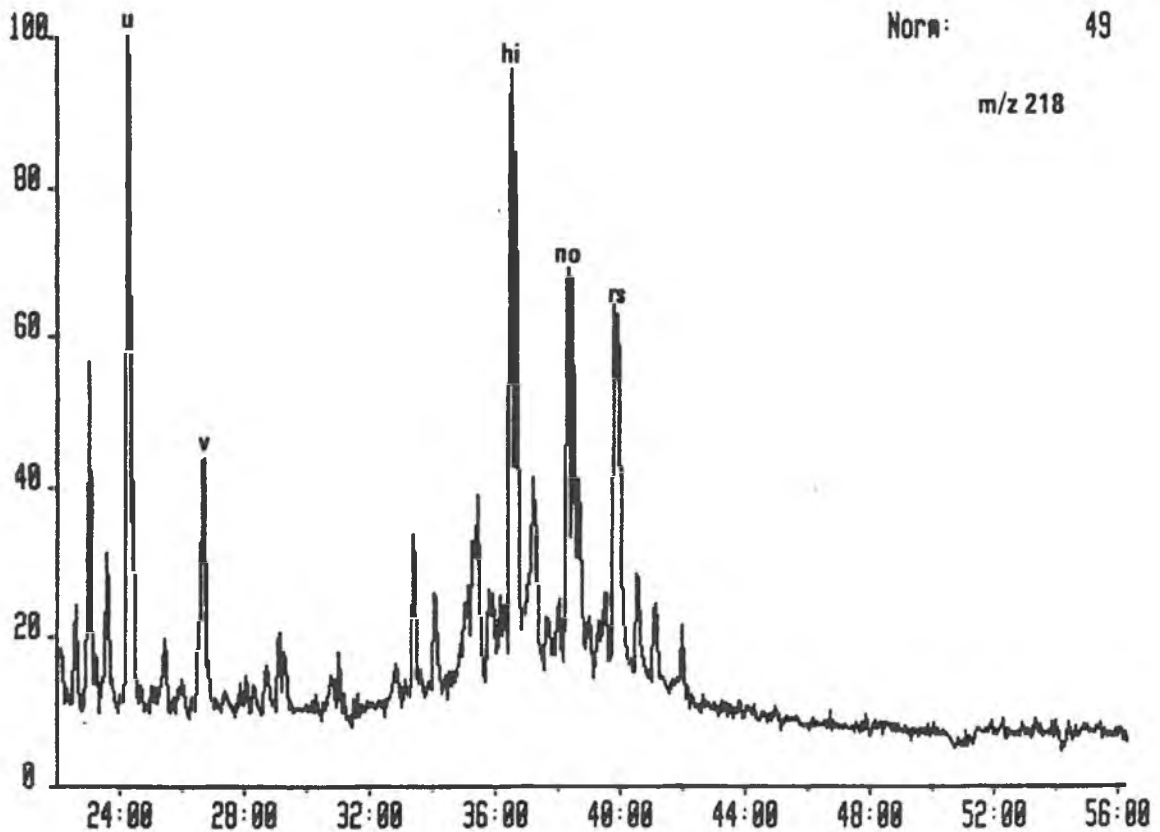
System:TRIT



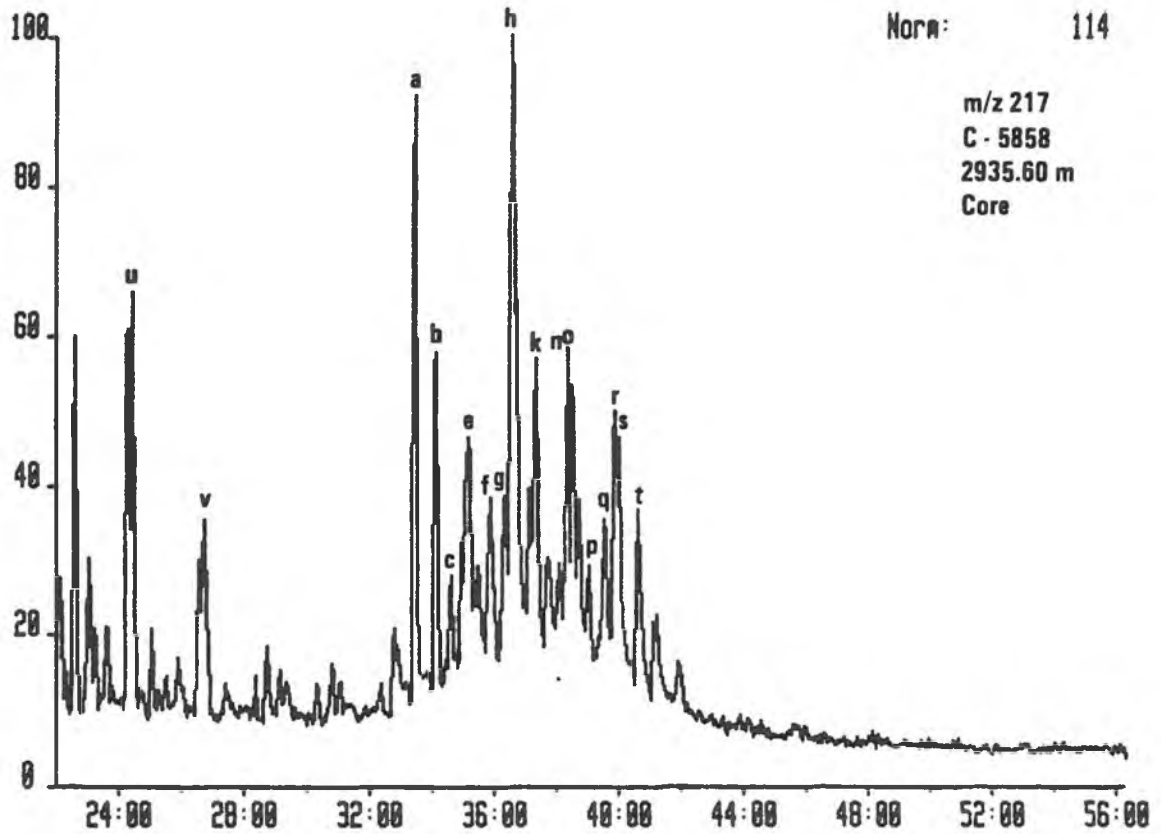
C5855SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU System:TRIT
 Sample 1 Injection 1 Group 1 Mass 217.1000
 Text:



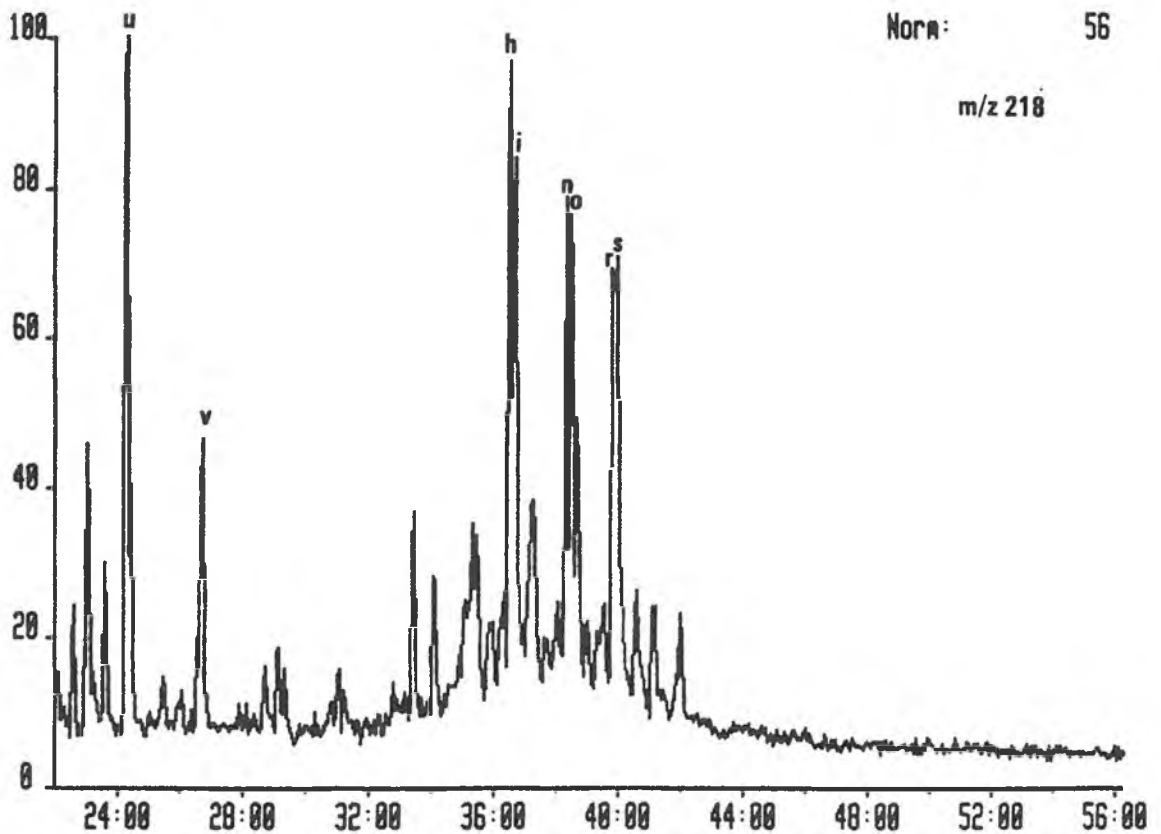
C5855SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU System:TRIT
 Sample 1 Injection 1 Group 1 Mass 218.1000
 Text:



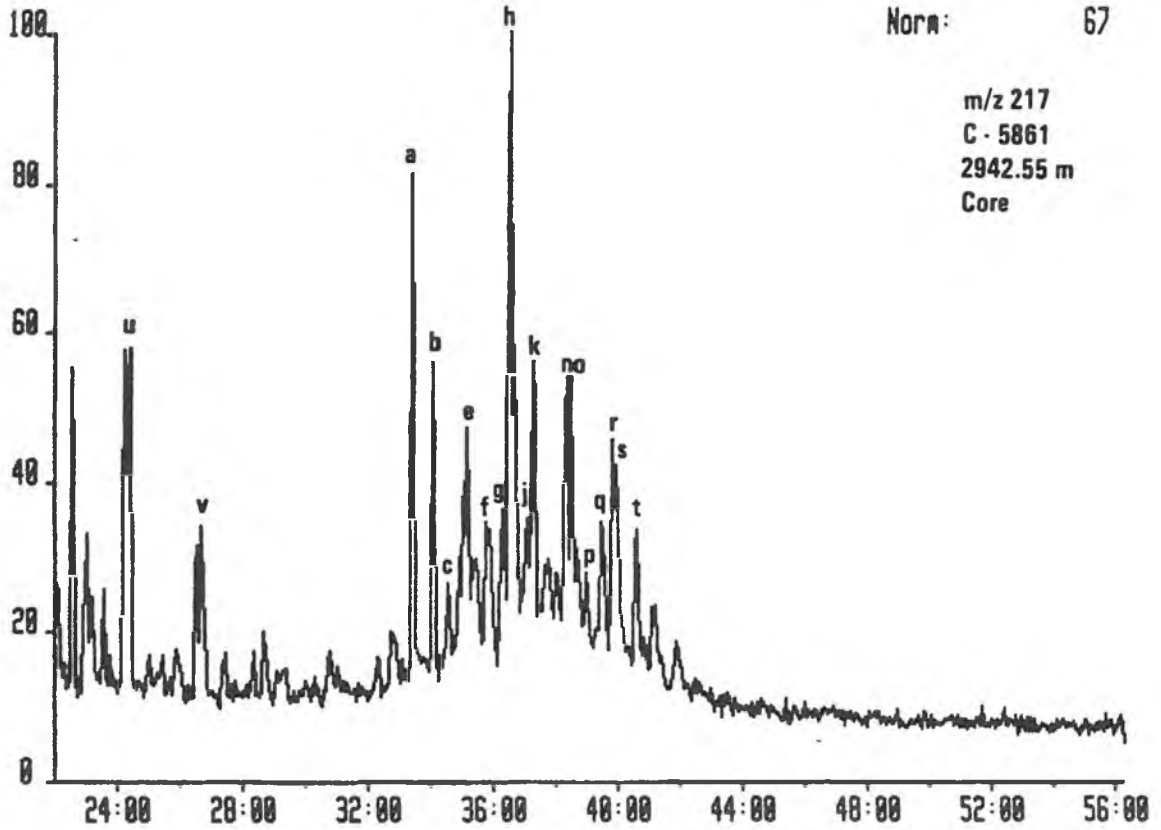
C5050SAT 6-FEB-87 Sir:Voltage 7070H Acnt:IKU System:TRIT
 Sample 1 Injection 1 Group 1 Mass 217.1000
 Text:



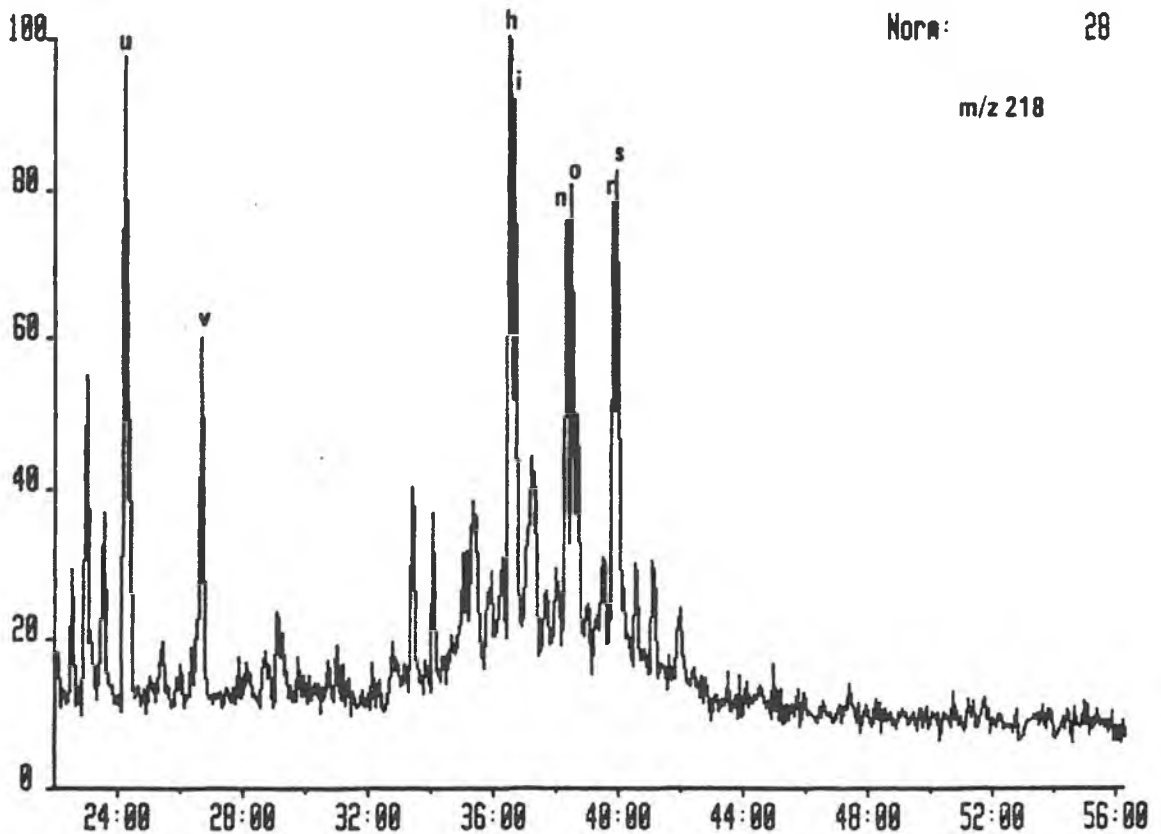
C5050SAT 6-FEB-87 Sir:Voltage 7070H Acnt:IKU System:TRIT
 Sample 1 Injection 1 Group 1 Mass 218.1000
 Text:



C5861SAT 6-FEB-87 Sir:Voltage 7070H Acnt:IKU System:TRIT
 Sample 1 Injection 1 Group 1 Mass 217.1000
 Text:

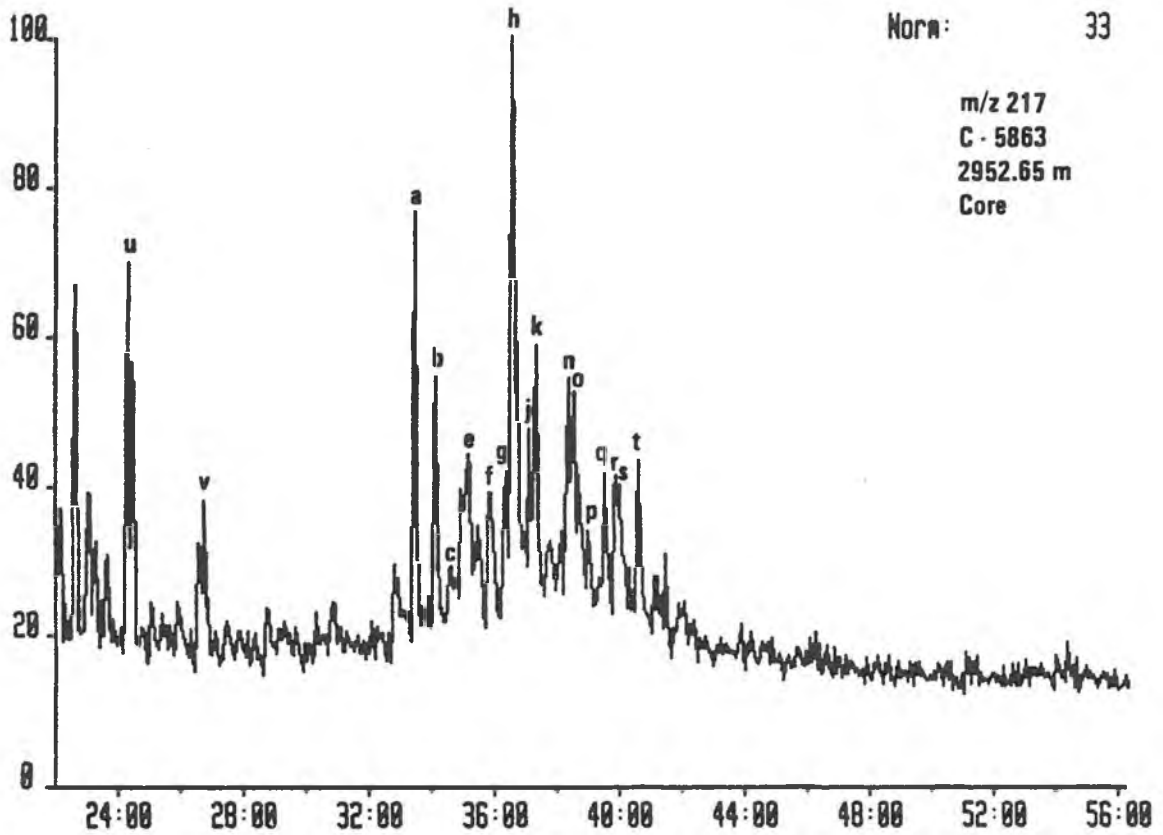


C5861SAT 6-FEB-87 Sir:Voltage 7070H Acnt:IKU System:TRIT
 Sample 1 Injection 1 Group 1 Mass 218.1000
 Text:



C5863SAT2 6-FEB-07 Sir:Voltage 7070H Acnt:IKU
 Sample 1 Injection 1 Group 1 Mass 217.1000
 Text:

System:TRIT



C5863SAT2 6-FEB-07 Sir:Voltage 7070H Acnt:IKU
 Sample 1 Injection 1 Group 1 Mass 218.1000
 Text:

System:TRIT

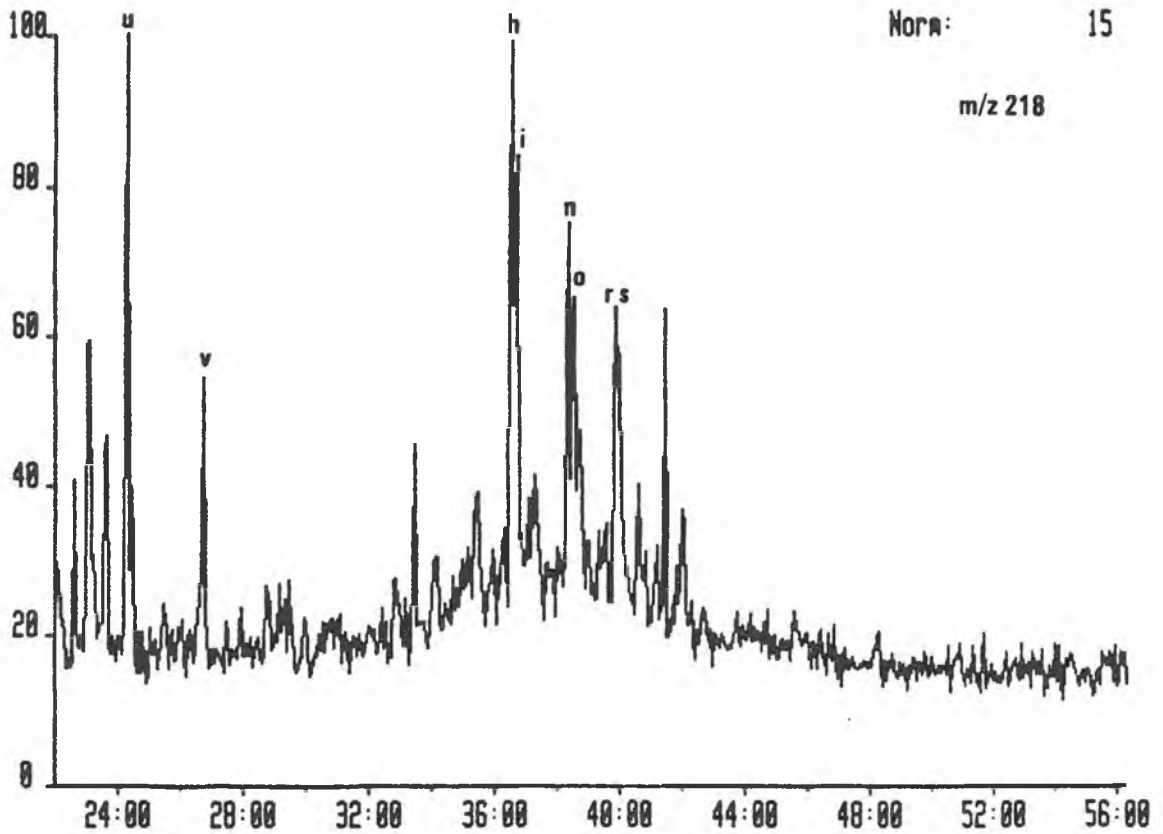
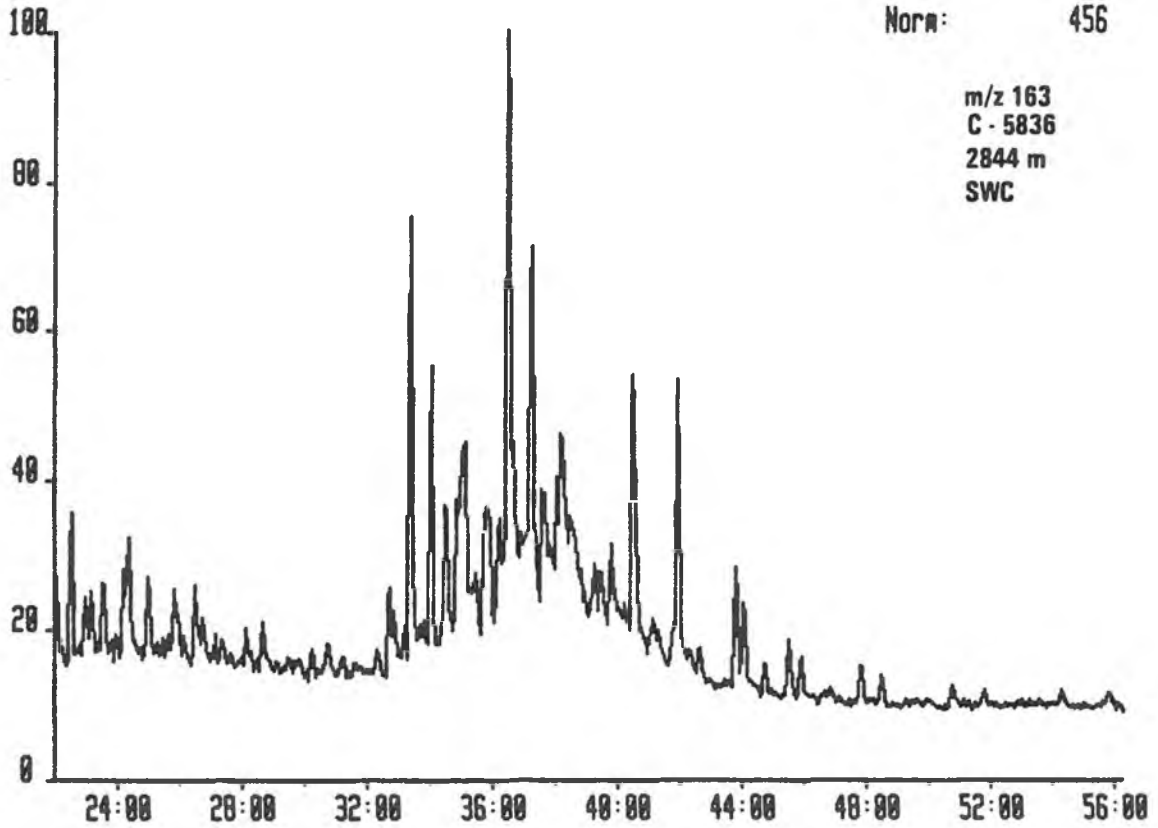


FIGURE 6

MASS CHROMATOGRAMS OF M/Z 163 AND 177

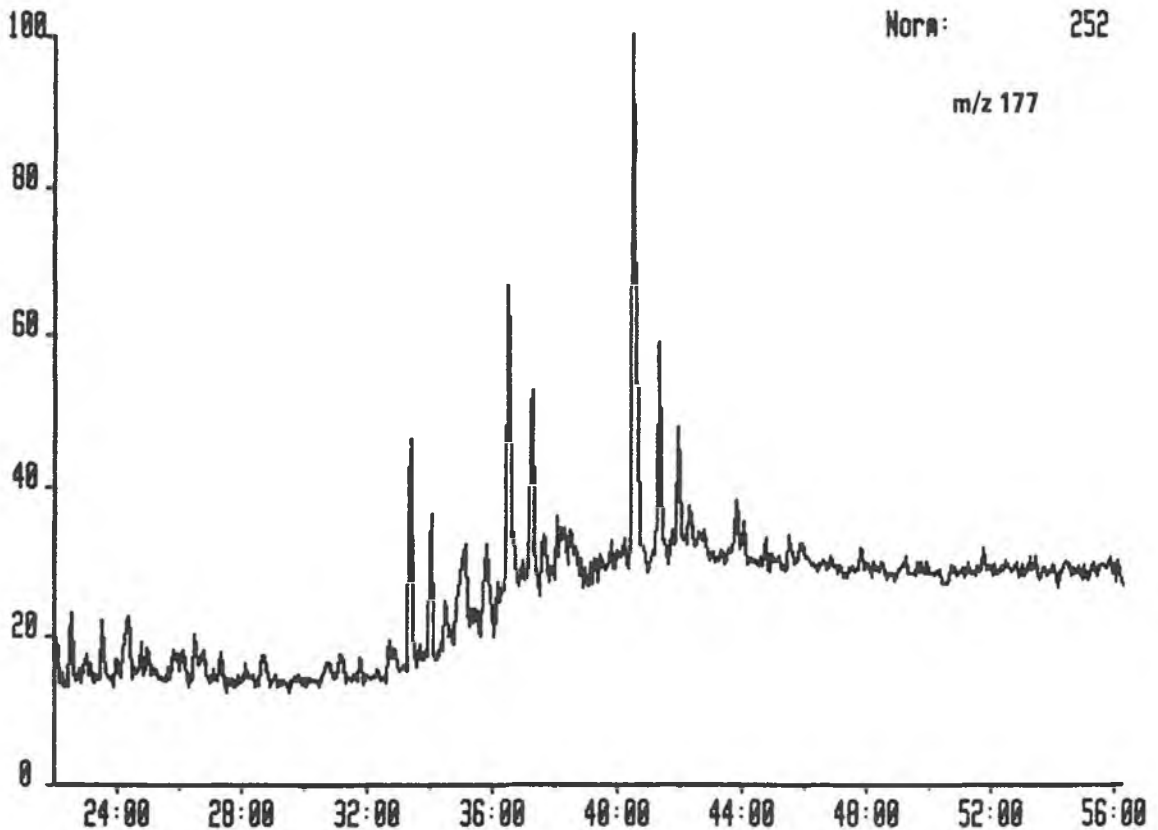
C5836SAT 5-FEB-07 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 163.1000
Text:

System:TRIT

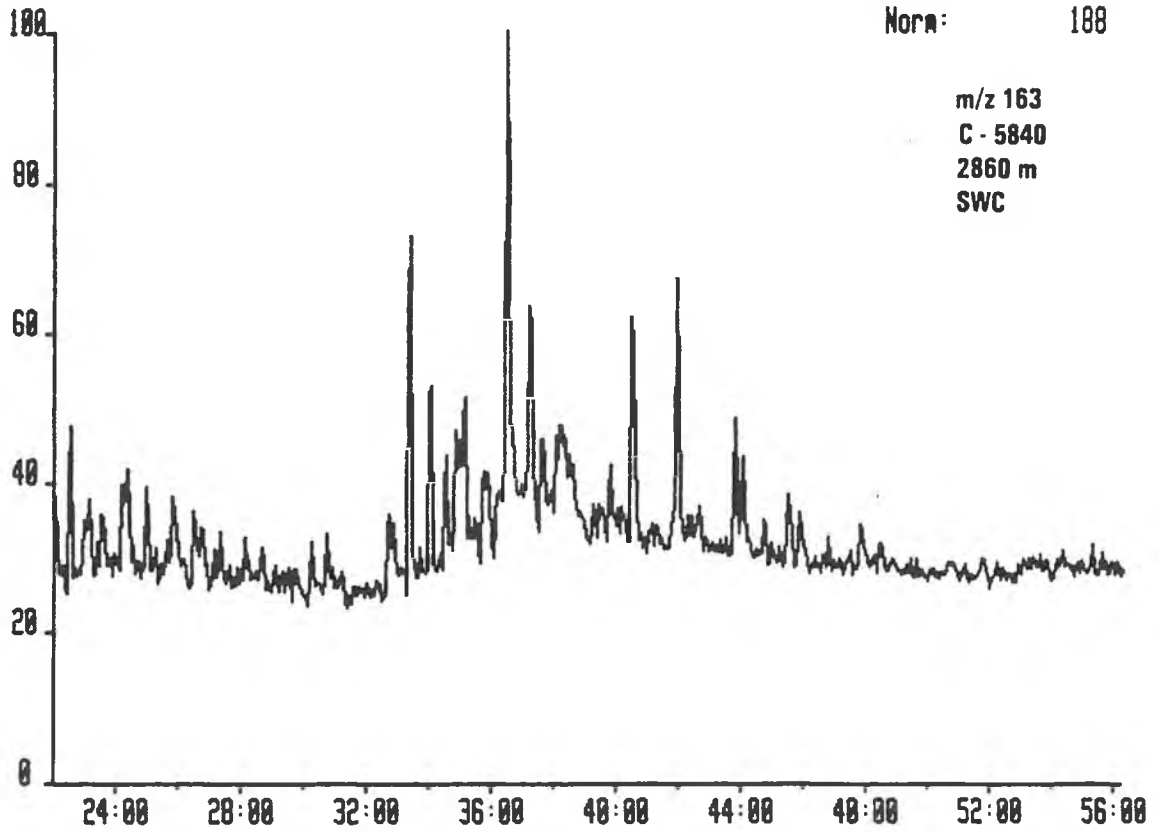


C5836SAT 5-FEB-07 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 177.1000
Text:

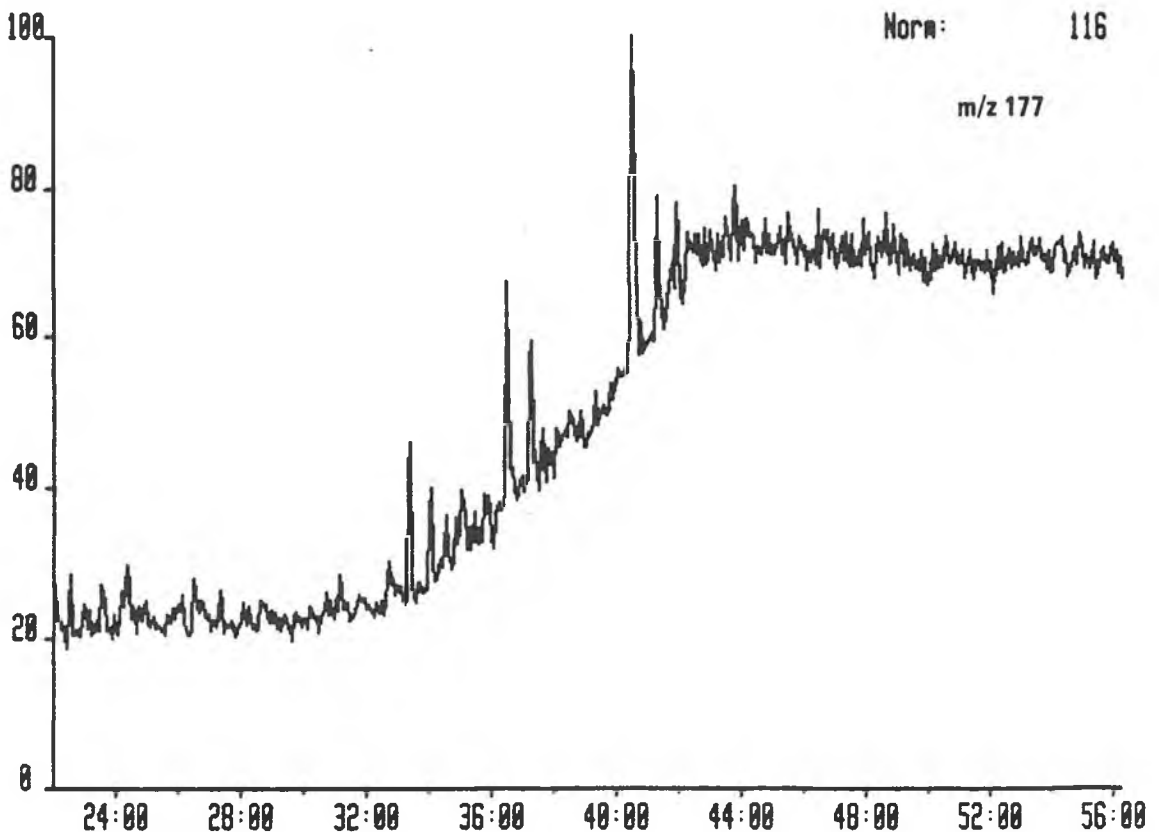
System:TRIT



C5840SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU System:TRIT
Sample 1 Injection 1 Group 1 Mass 163.1000
Text:

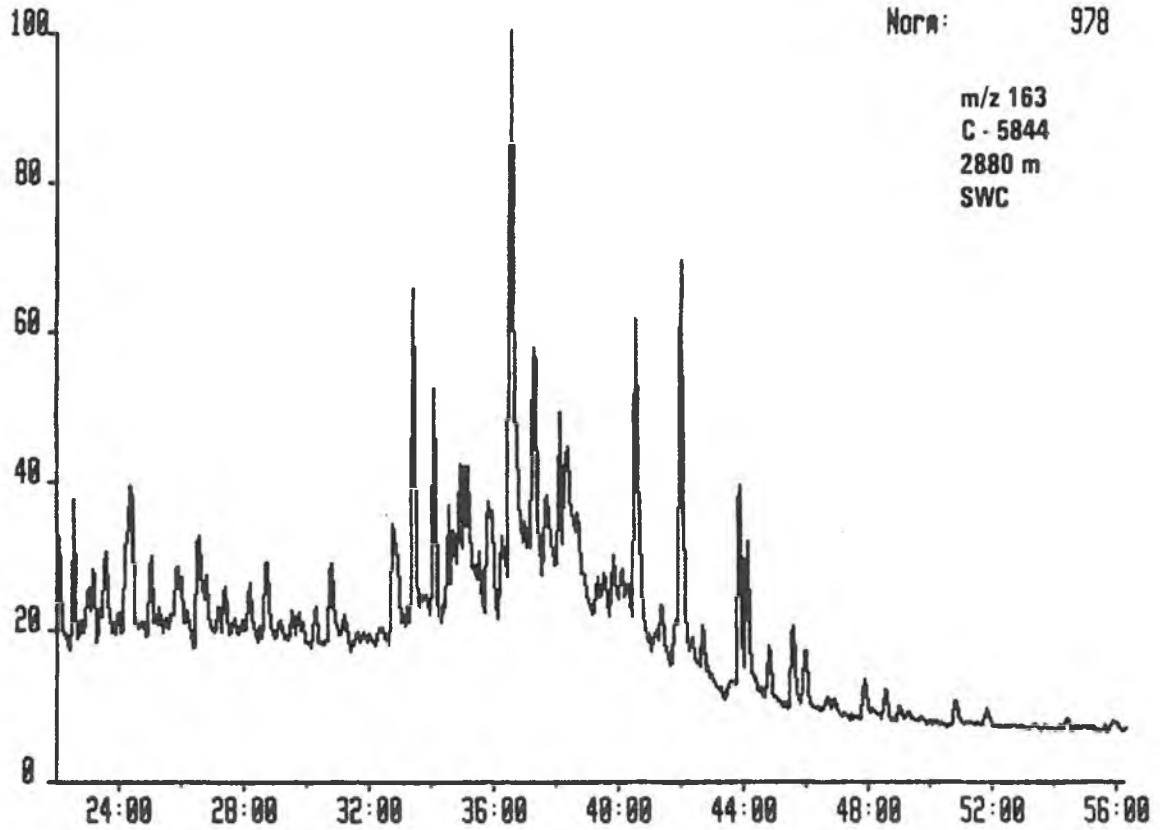


C5840SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU System:TRIT
Sample 1 Injection 1 Group 1 Mass 177.1000
Text:



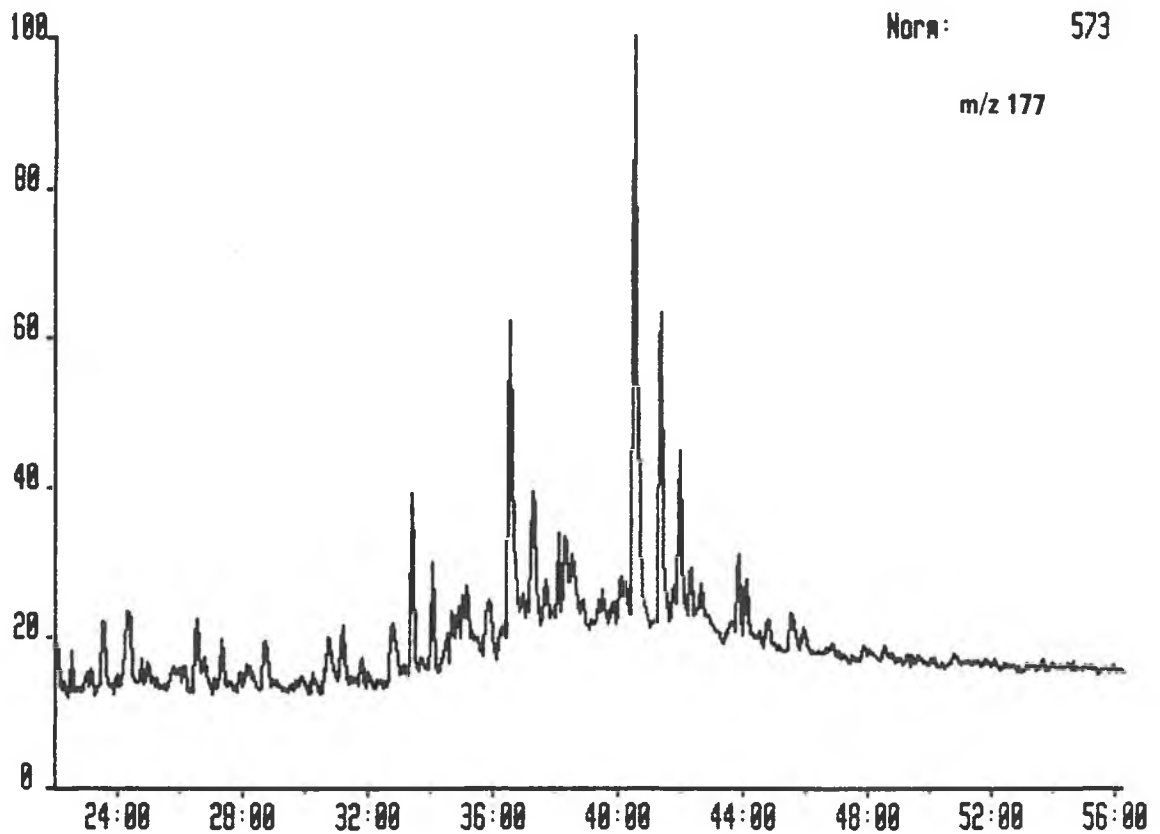
C5044SAT 6-FEB-07 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 163.1000
Text:

System:TRIT

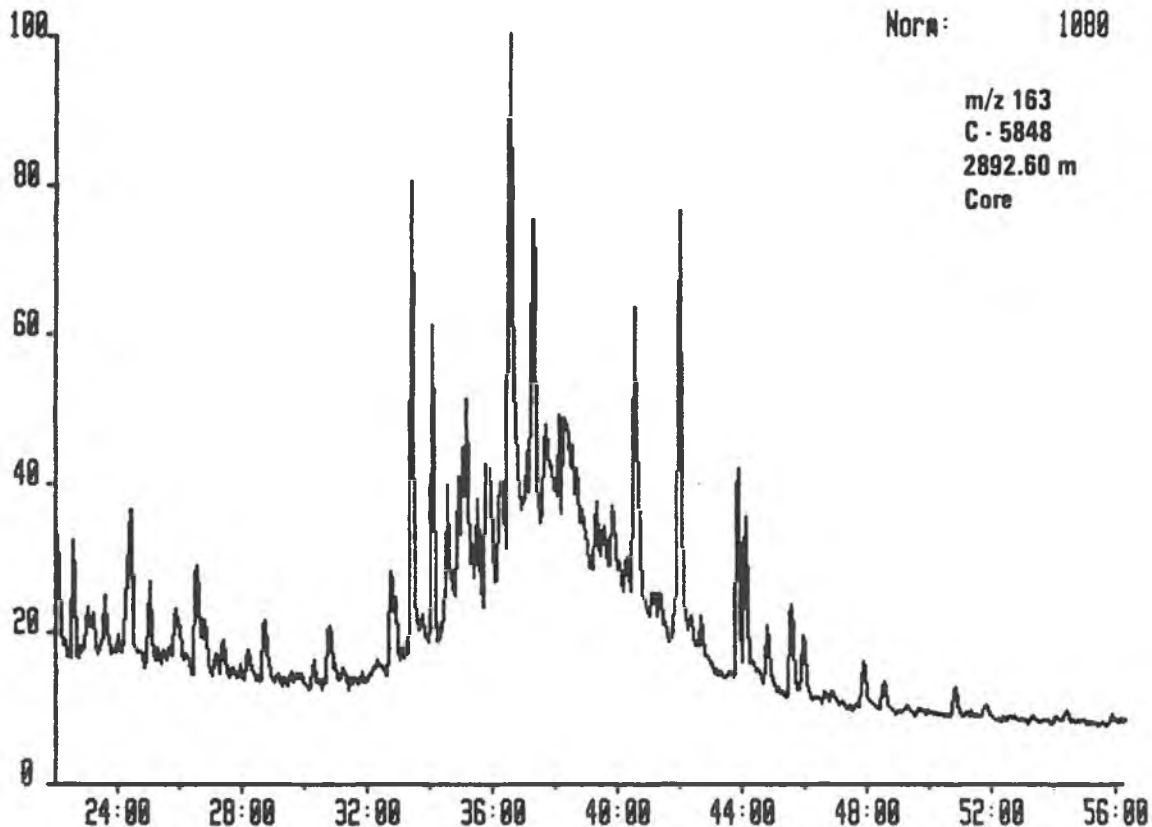


C5044SAT 6-FEB-07 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 177.1000
Text:

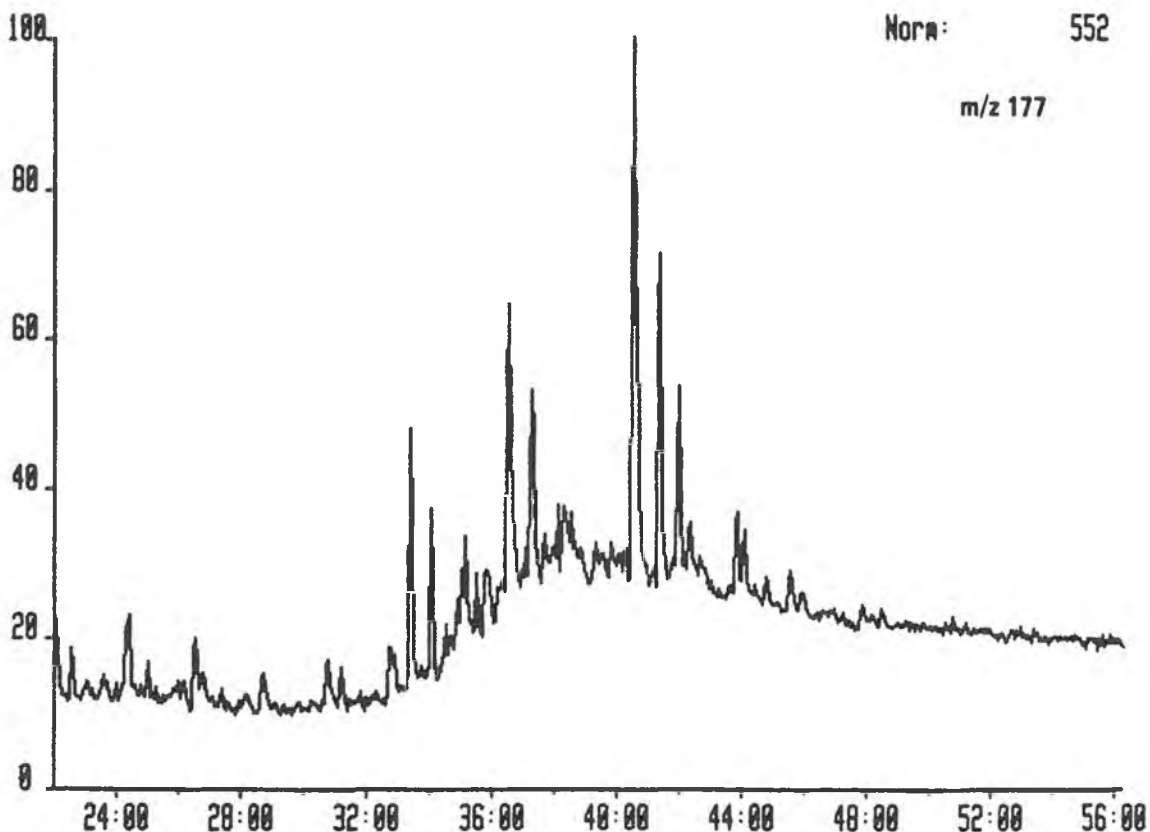
System:TRIT



C5848SAT 5-FEB-07 Sir:Voltage 7070H Acnt:IKU System:TRIT
Sample 1 Injection 1 Group 1 Mass 163.1000
Text:

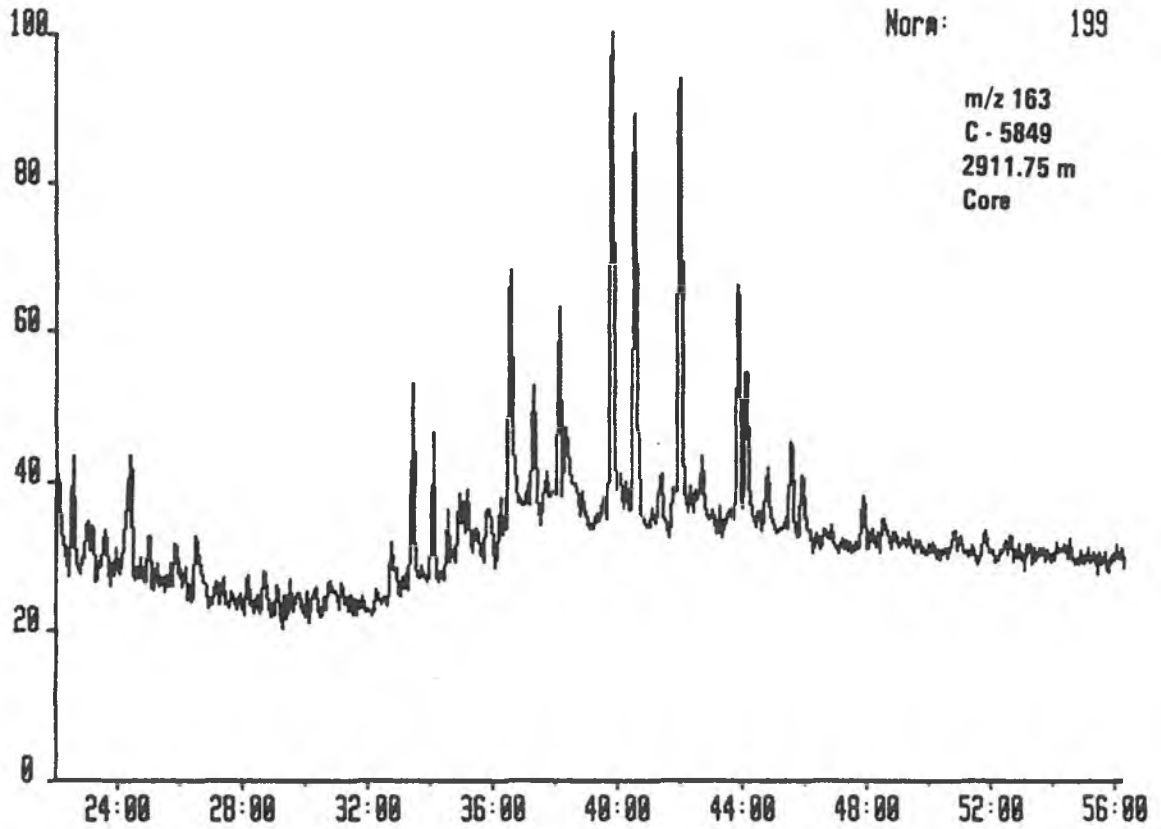


C5848SAT 5-FEB-07 Sir:Voltage 7070H Acnt:IKU System:TRIT
Sample 1 Injection 1 Group 1 Mass 177.1000
Text:



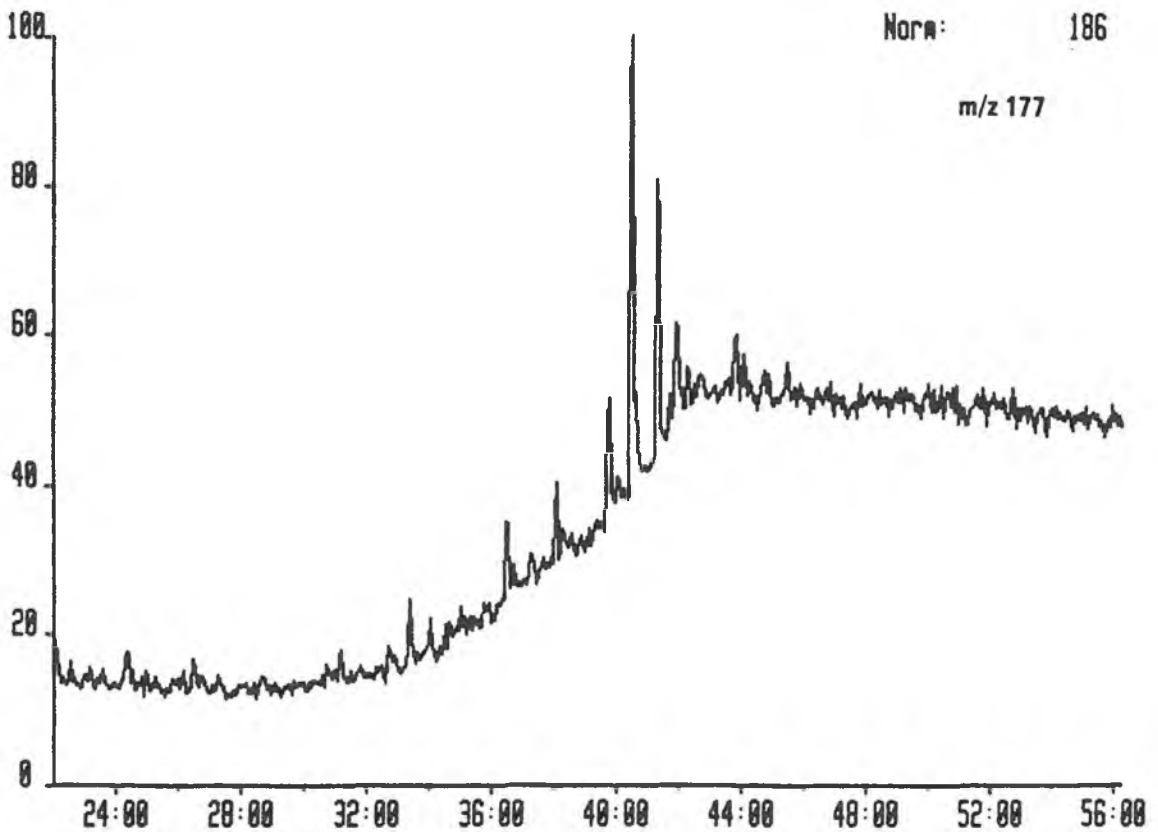
C5849SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 163.1000
Text:

System:TRIT



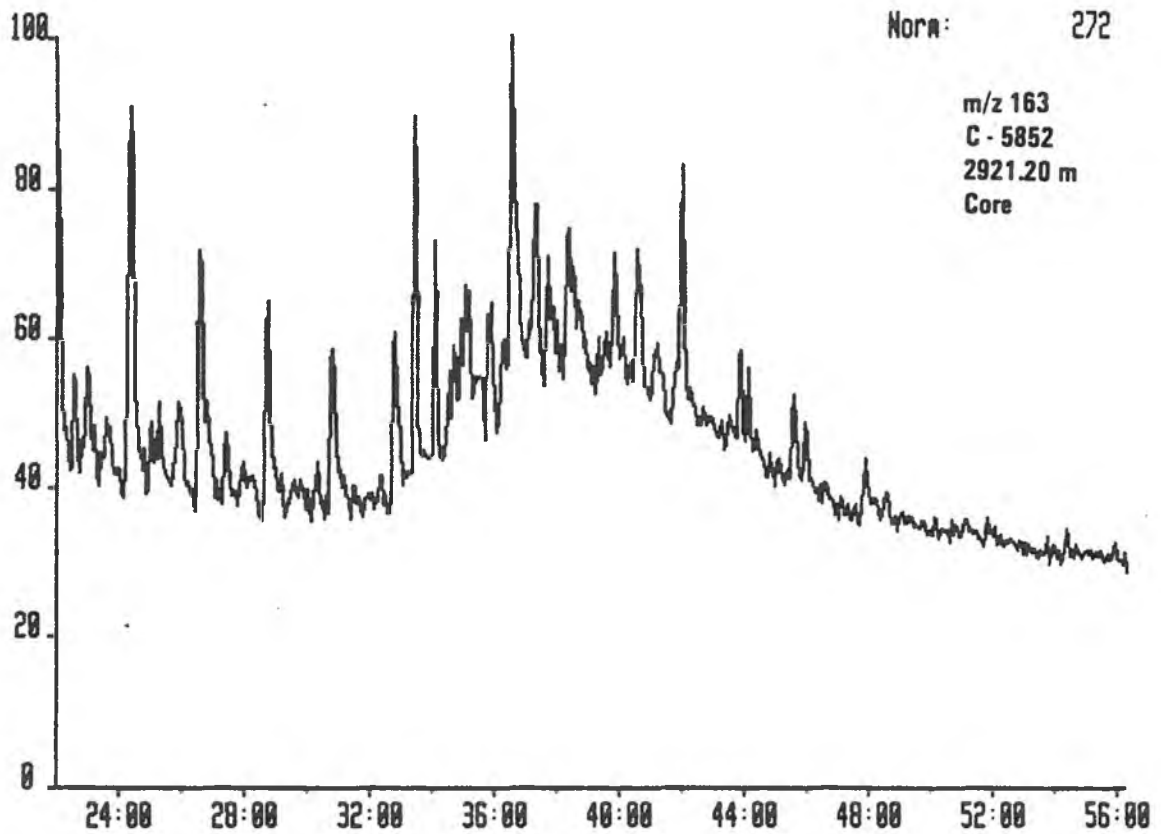
C5849SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 177.1000
Text:

System:TRIT



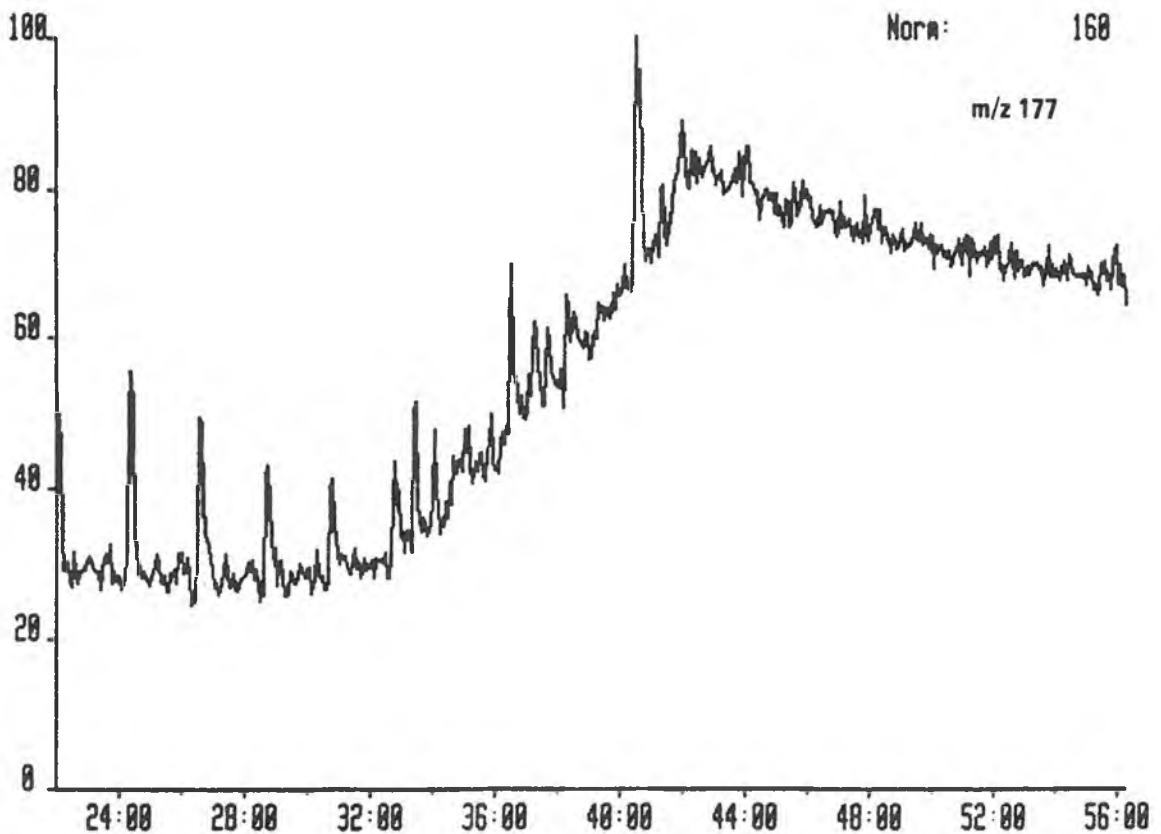
C5852SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 163.1000
Text:

System:TRIT

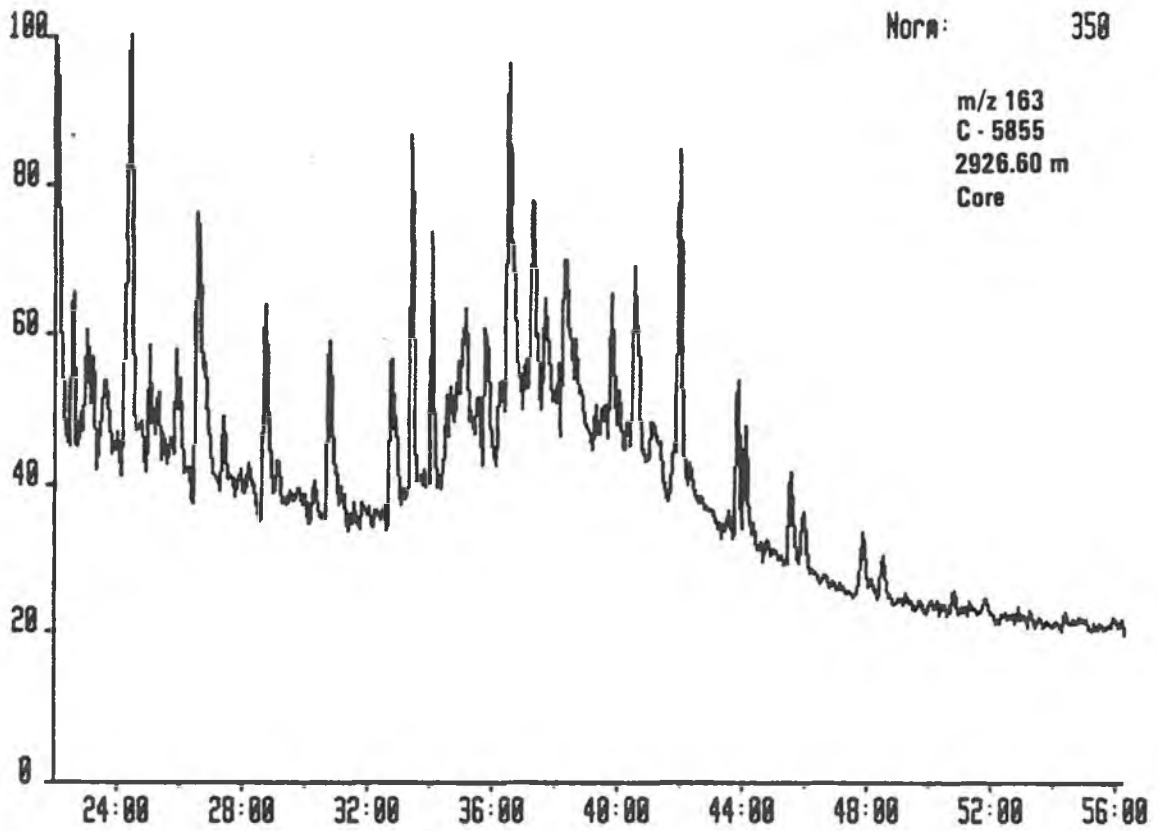


C5852SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 177.1000
Text:

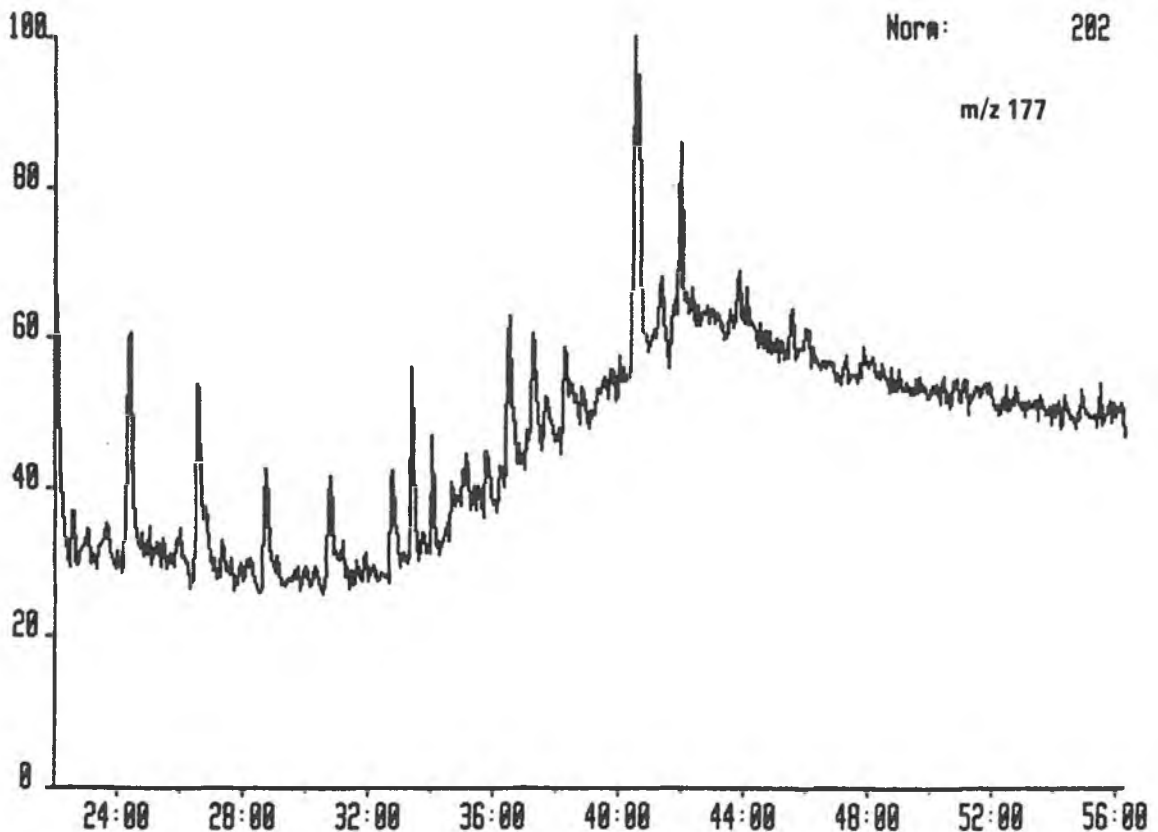
System:TRIT



C5855SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU System:TRIT
Sample 1 Injection 1 Group 1 Mass 163.1000
Text:

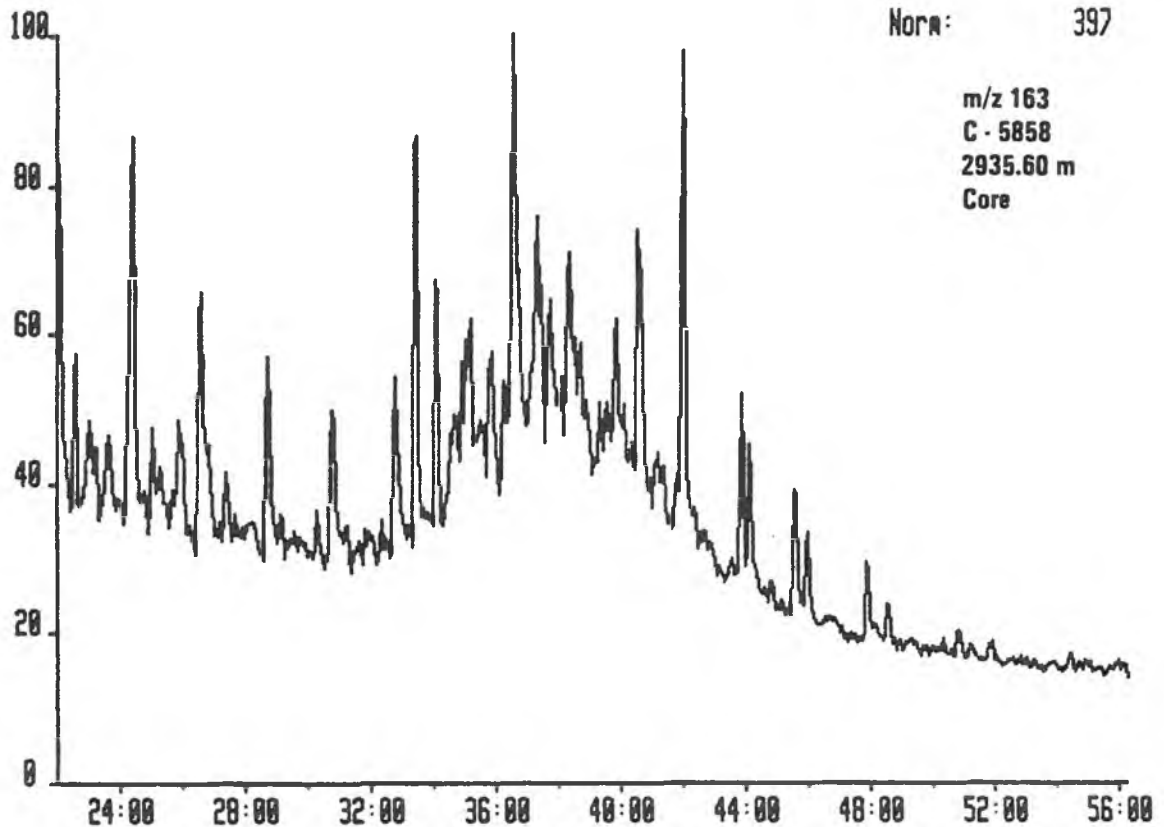


C5855SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU System:TRIT
Sample 1 Injection 1 Group 1 Mass 177.1000
Text:



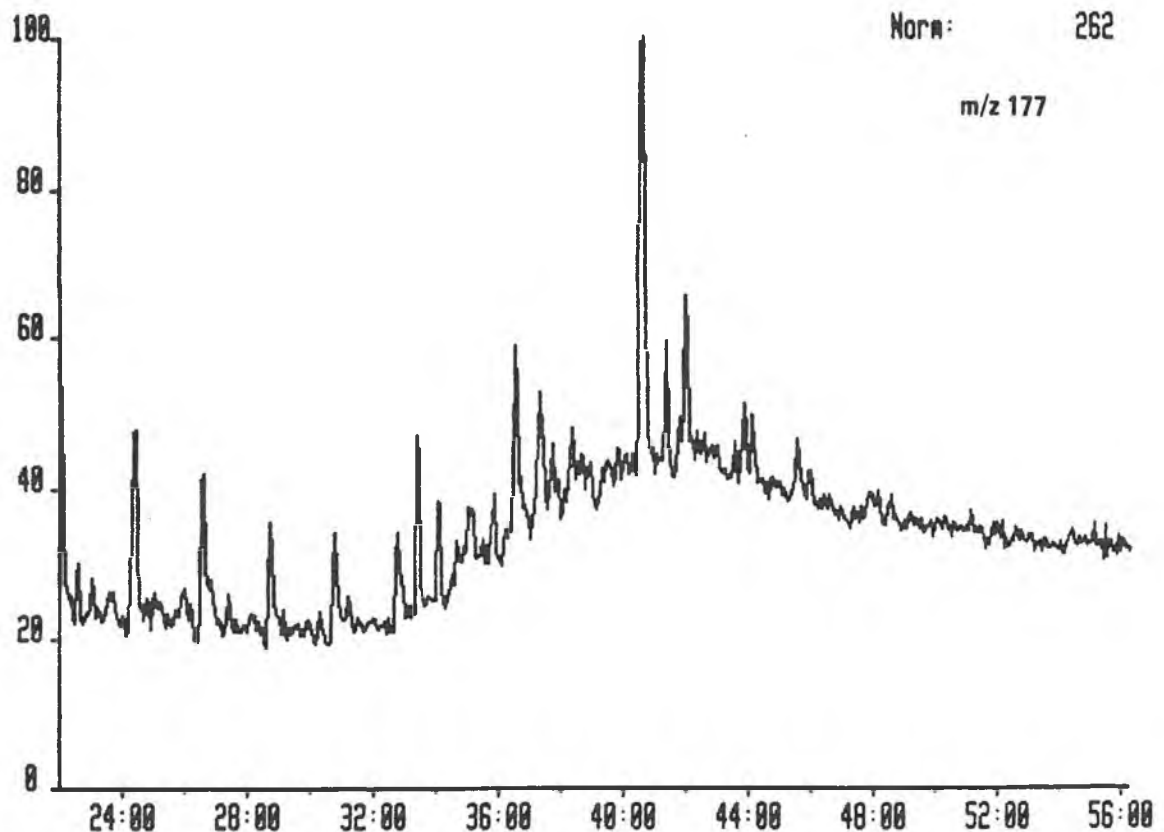
C5858SAT 6-FEB-87 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 163.1000
Text:

System:TRIT



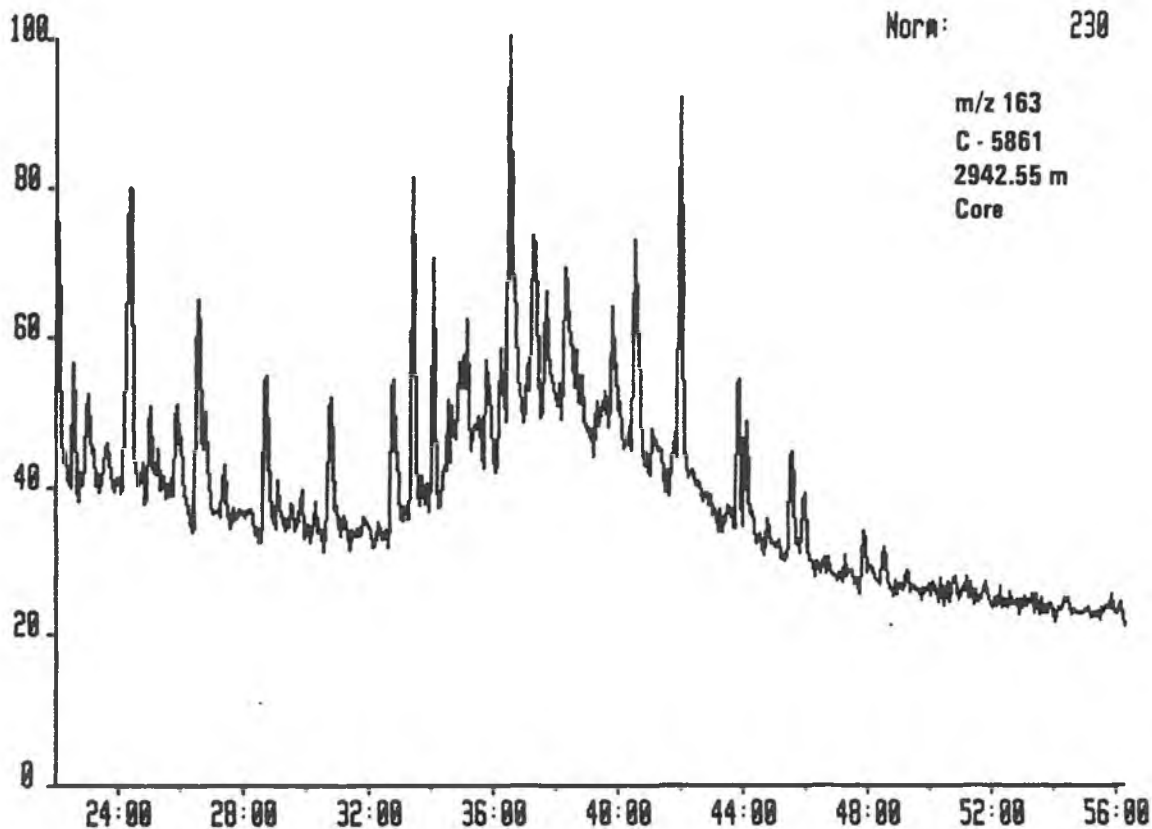
C5858SAT 6-FEB-87 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 177.1000
Text:

System:TRIT



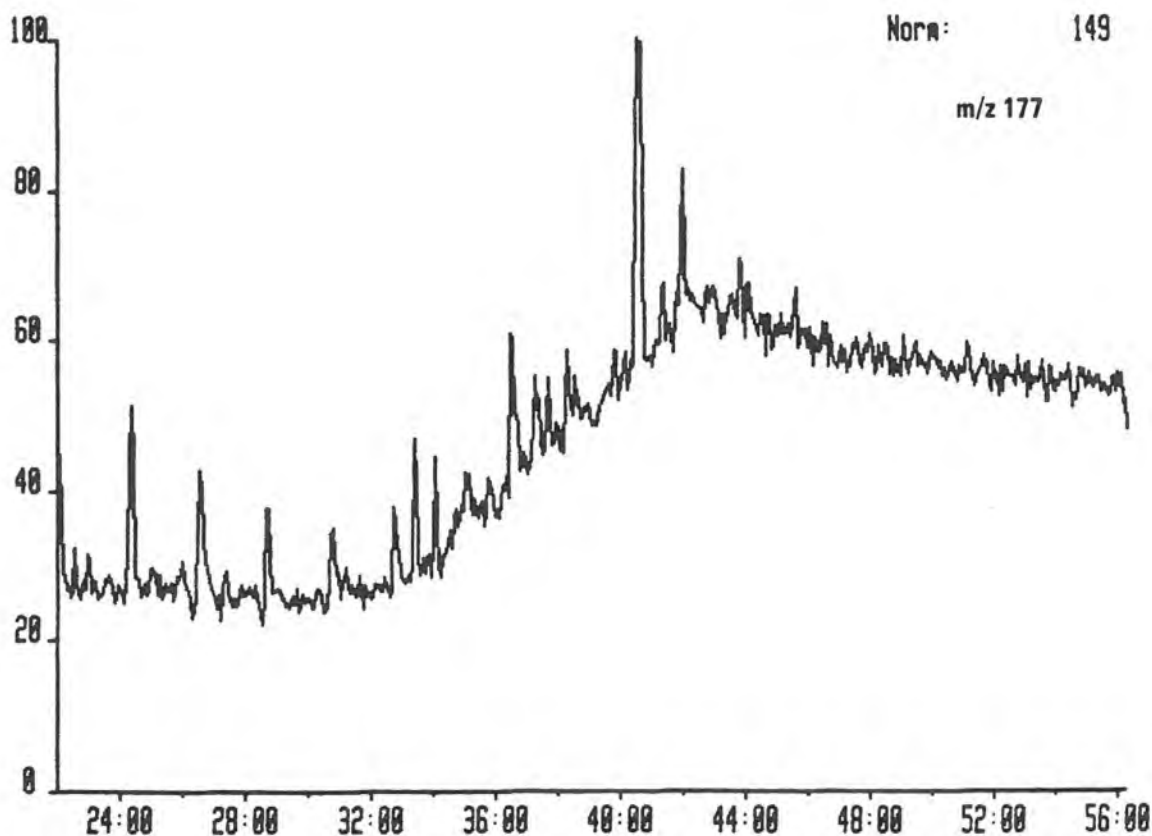
C5861SAT 6-FEB-87 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 163.1000
Text:

System:TRIT

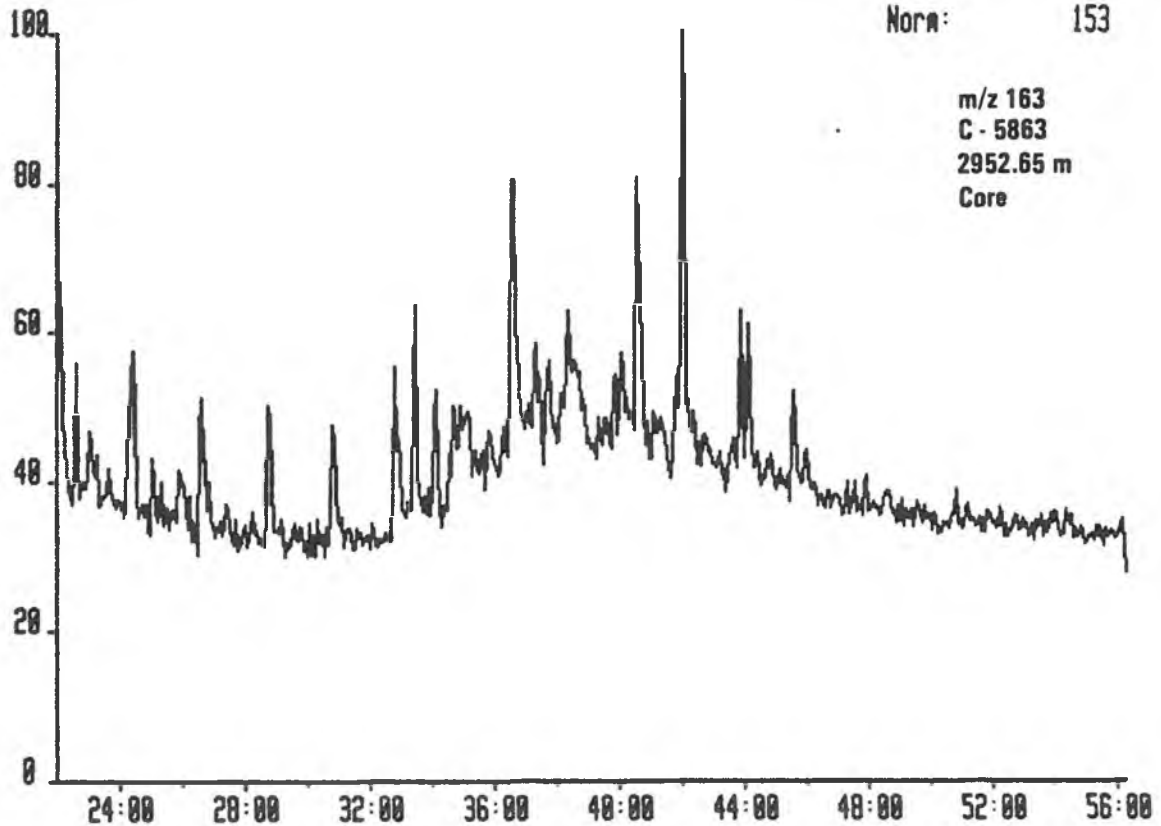


C5861SAT 6-FEB-87 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 177.1000
Text:

System:TRIT



C5863SAT2 6-FEB-87 Sir:Voltage 7070H Acnt:IKU System:TRIT
Sample 1 Injection 1 Group 1 Mass 163.1000
Text:



C5863SAT2 6-FEB-87 Sir:Voltage 7070H Acnt:IKU System:TRIT
Sample 1 Injection 1 Group 1 Mass 177.1000
Text:

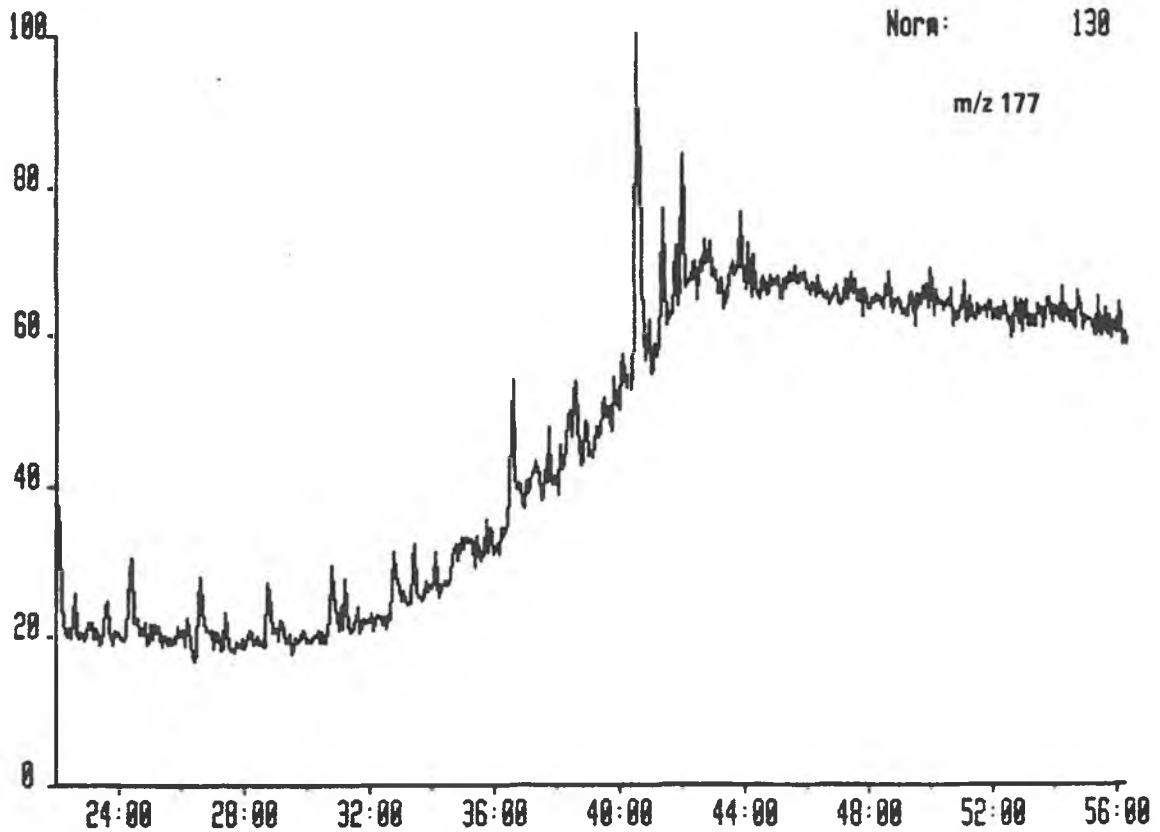
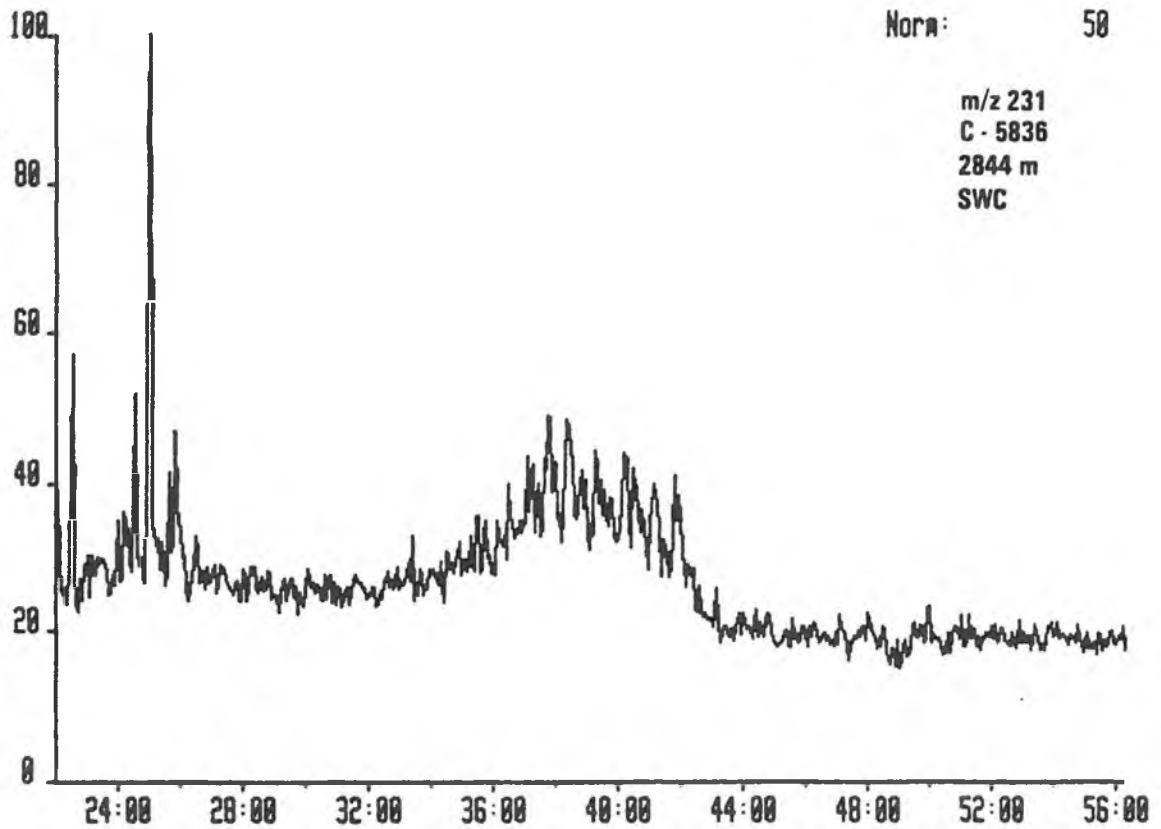


FIGURE 7

MASS CHROMATOGRAMS OF M/Z 231 AND 259

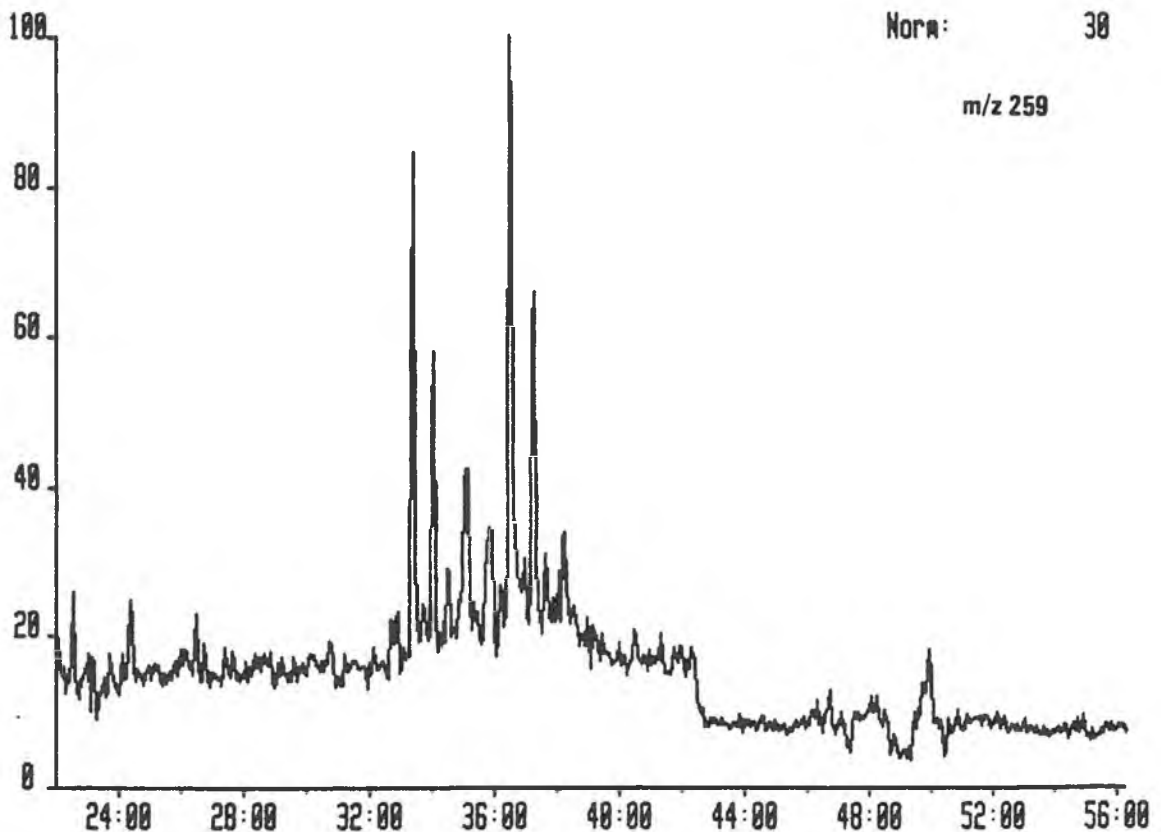
C5836SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 231.1000
Text:

System:TRIT

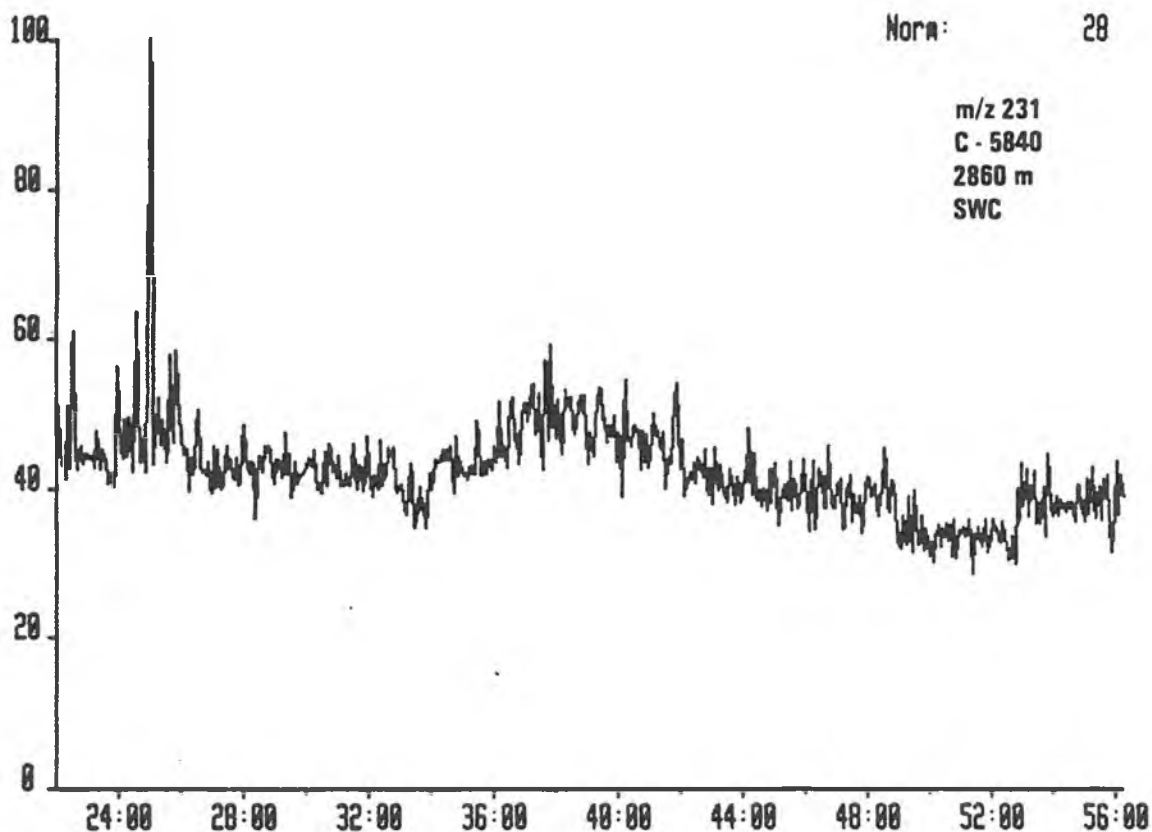


C5836SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 259.1000
Text:

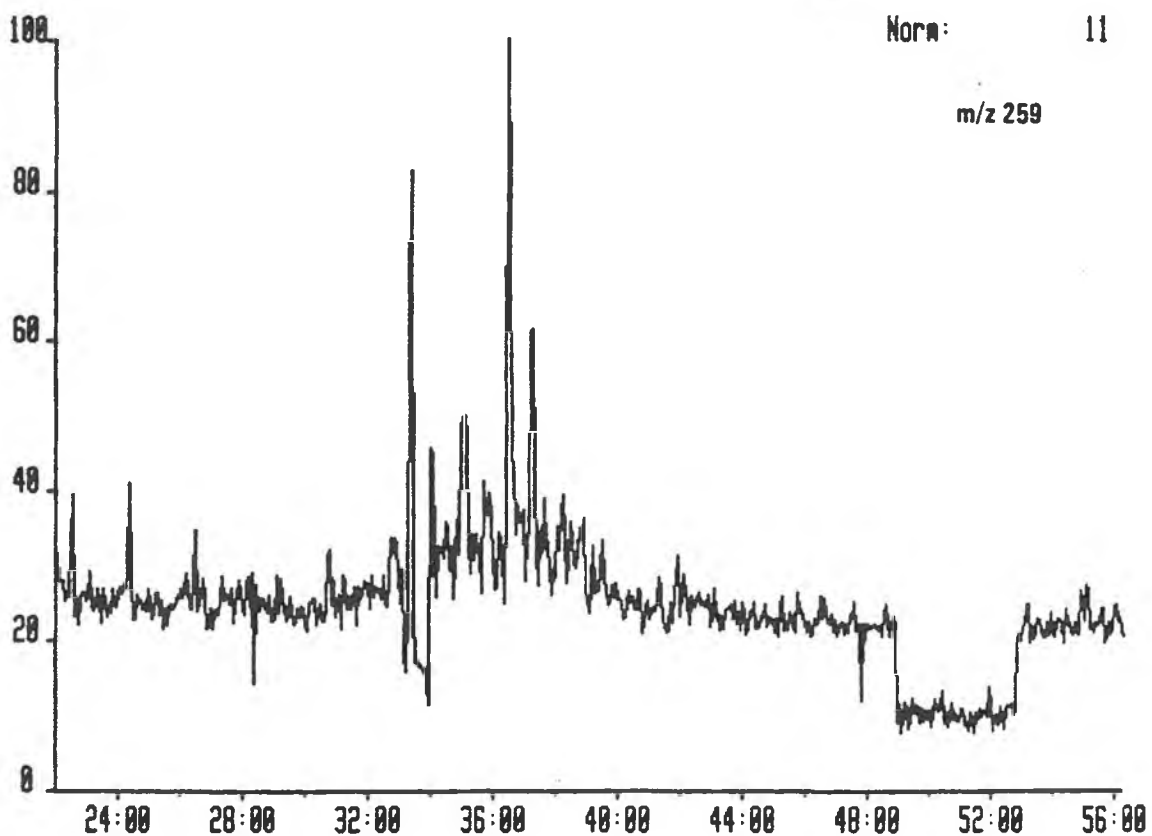
System:TRIT



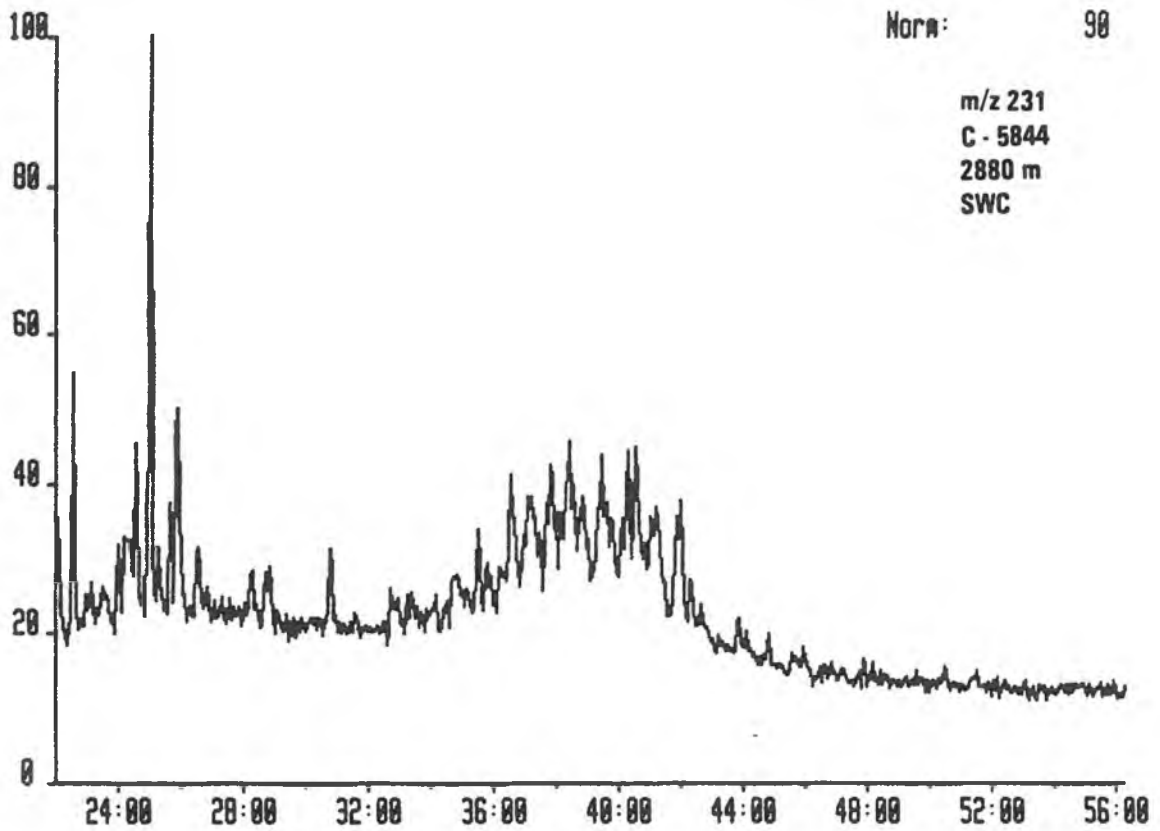
C5840SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU System:TRIT
Sample 1 Injection 1 Group 1 Mass 231.1000
Text:



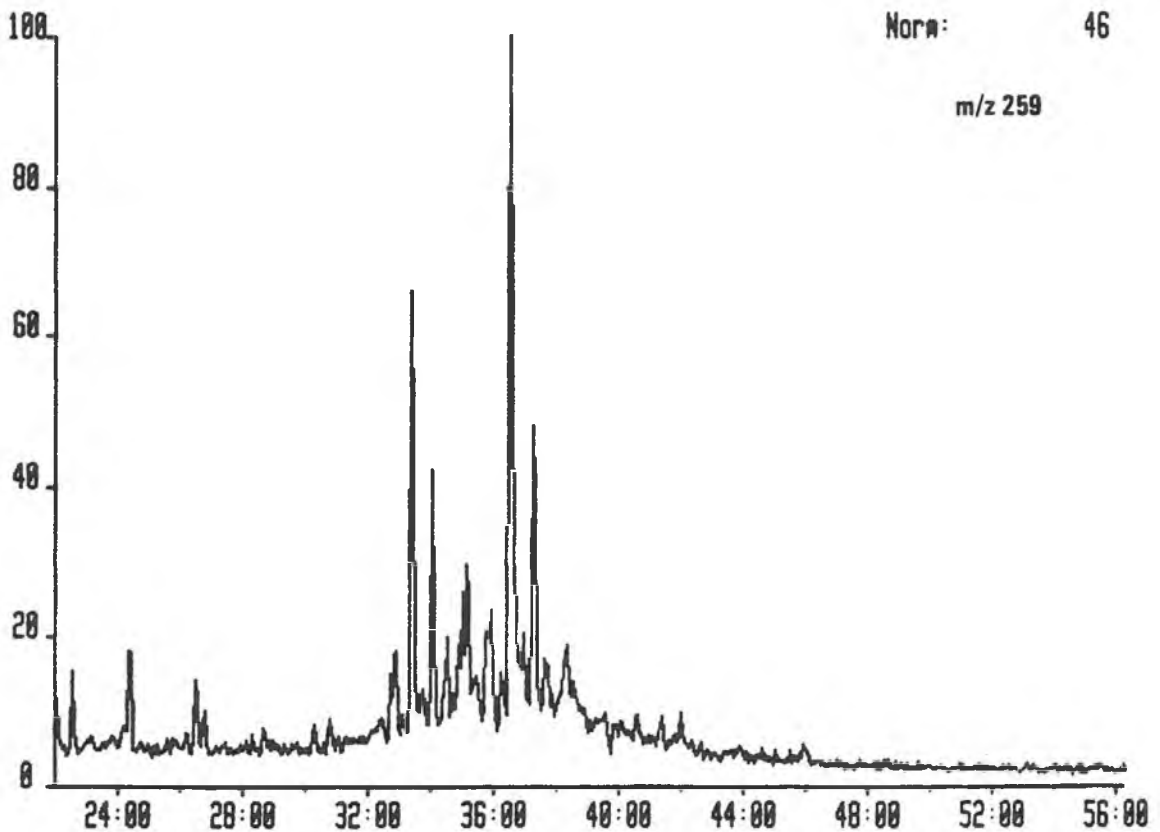
C5840SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU System:TRIT
Sample 1 Injection 1 Group 1 Mass 259.1000
Text:



C5844SAT 6-FEB-87 Sir:Voltage 7070H Acnt:IKU System:TRIT
Sample 1 Injection 1 Group 1 Mass 231.1000
Text:

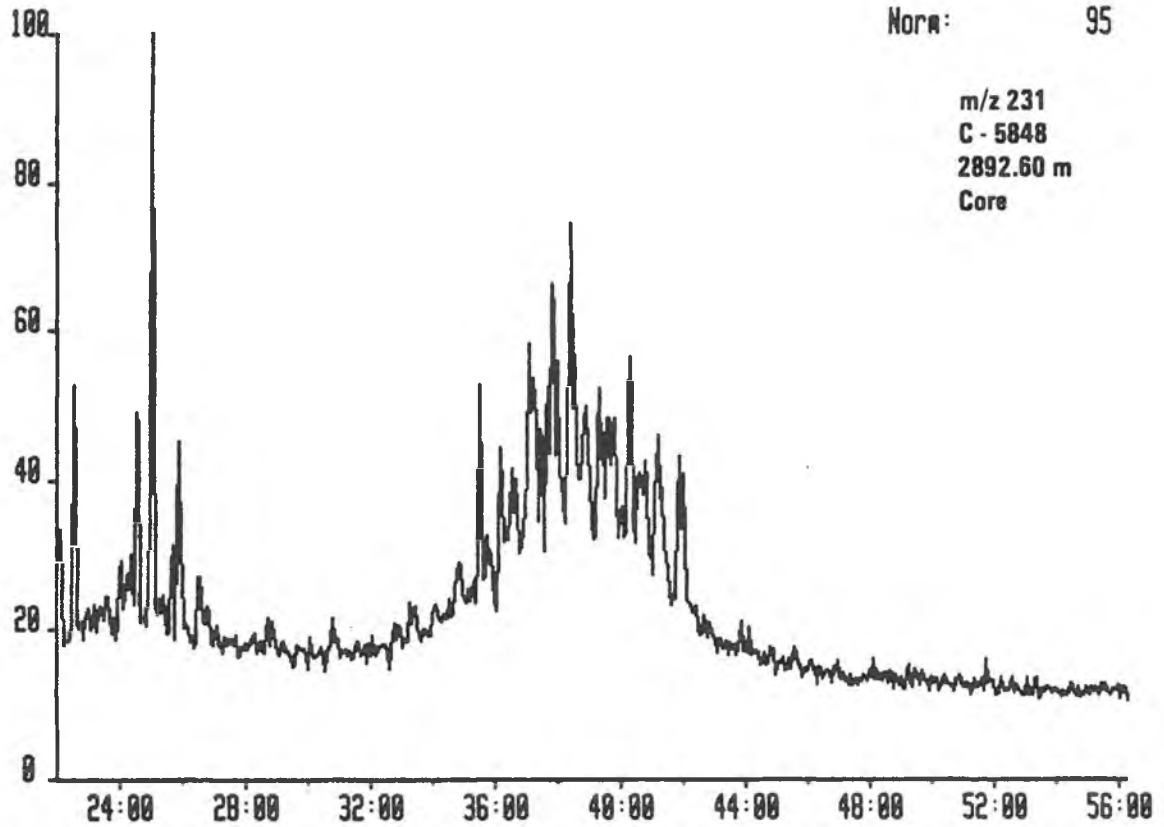


C5844SAT 6-FEB-87 Sir:Voltage 7070H Acnt:IKU System:TRIT
Sample 1 Injection 1 Group 1 Mass 259.1000
Text:



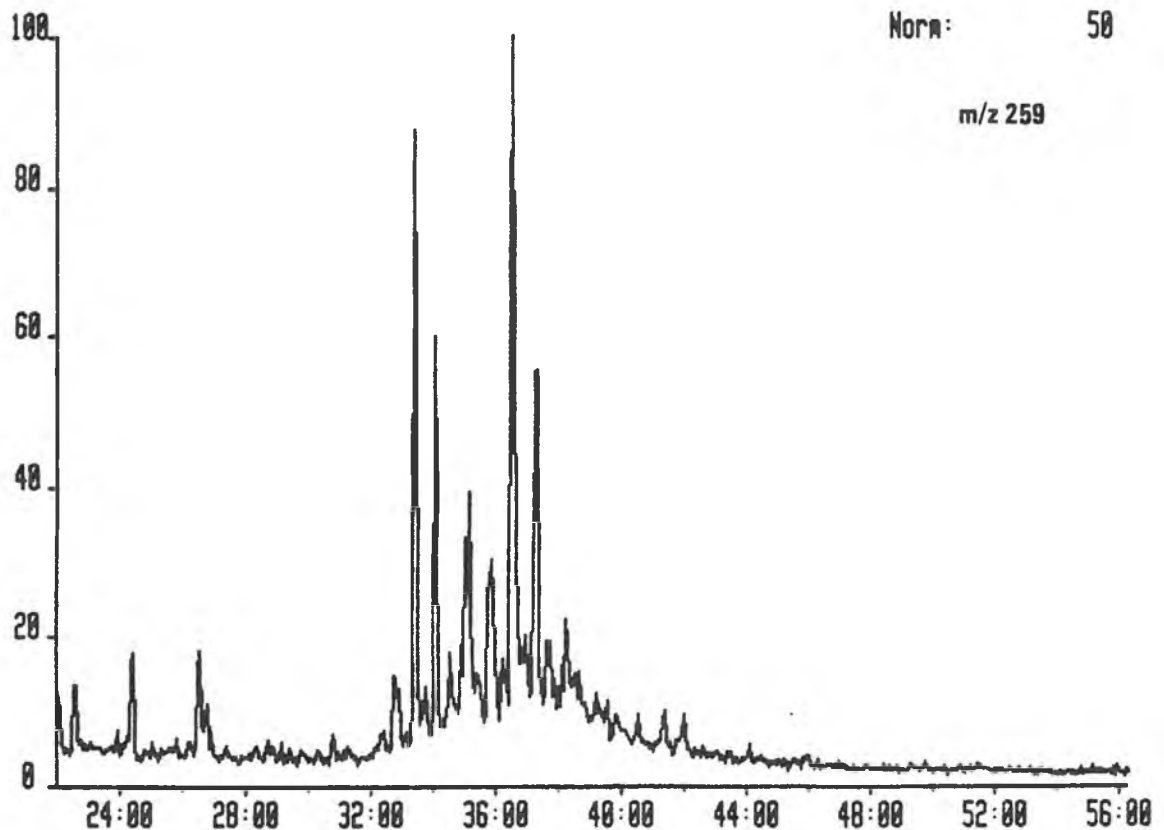
C5848SAT 5-FEB-07 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 231.1000
Text:

System:TRIT

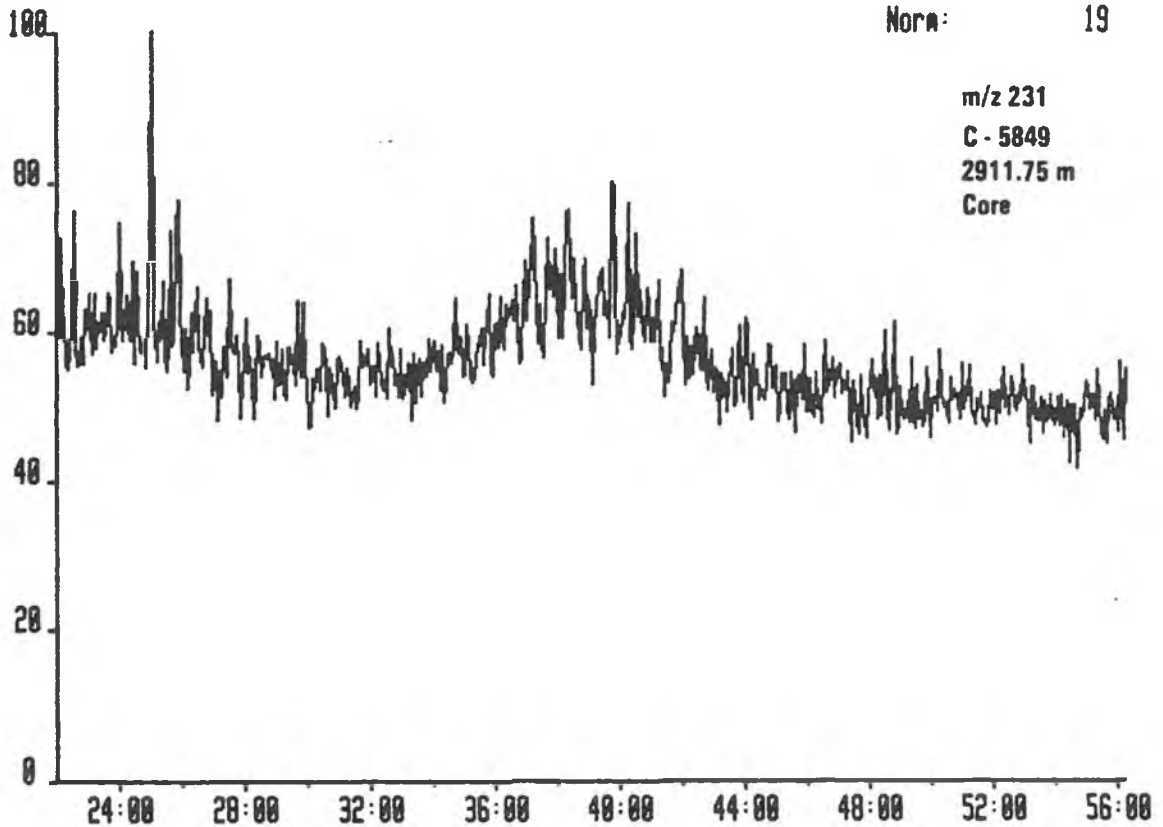


C5848SAT 5-FEB-07 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 259.1000
Text:

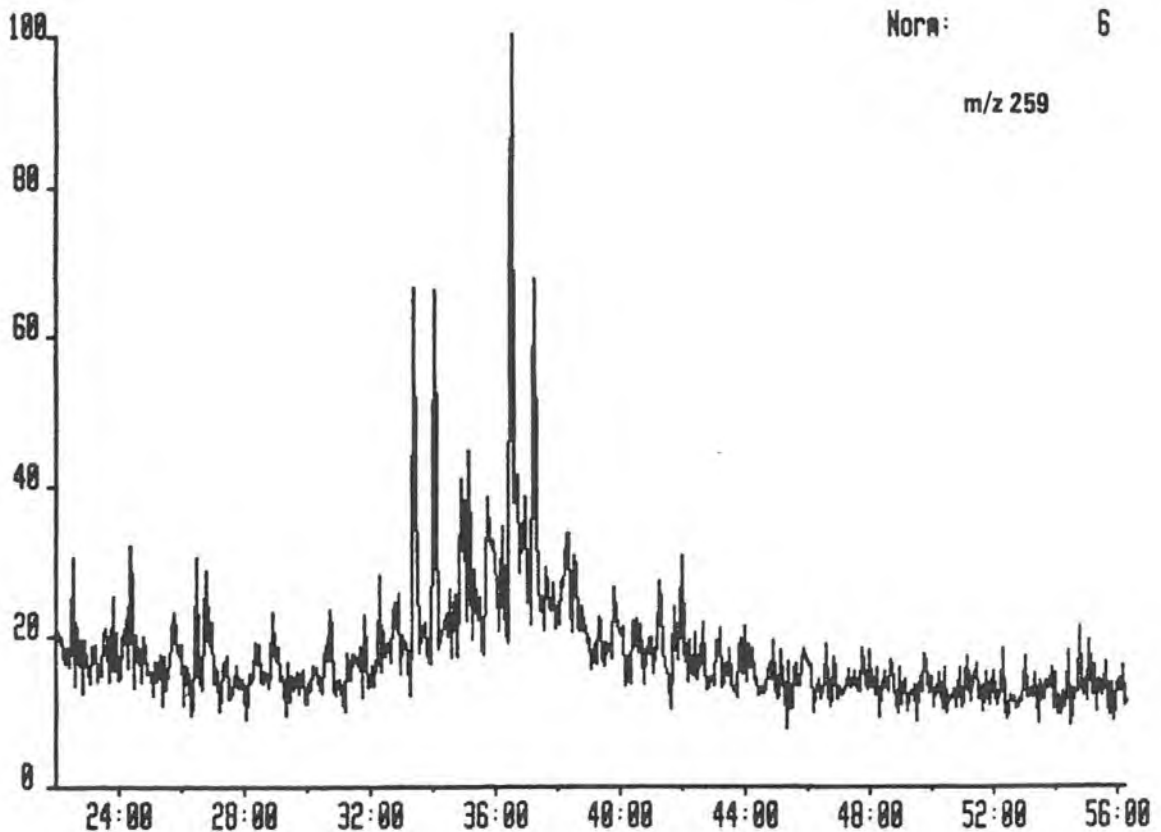
System:TRIT



C5849SAT 5-FEB-07 Sir:Voltage 7070H Acnt:IKU System:TRIT
Sample 1 Injection 1 Group 1 Mass 231.1000
Text:

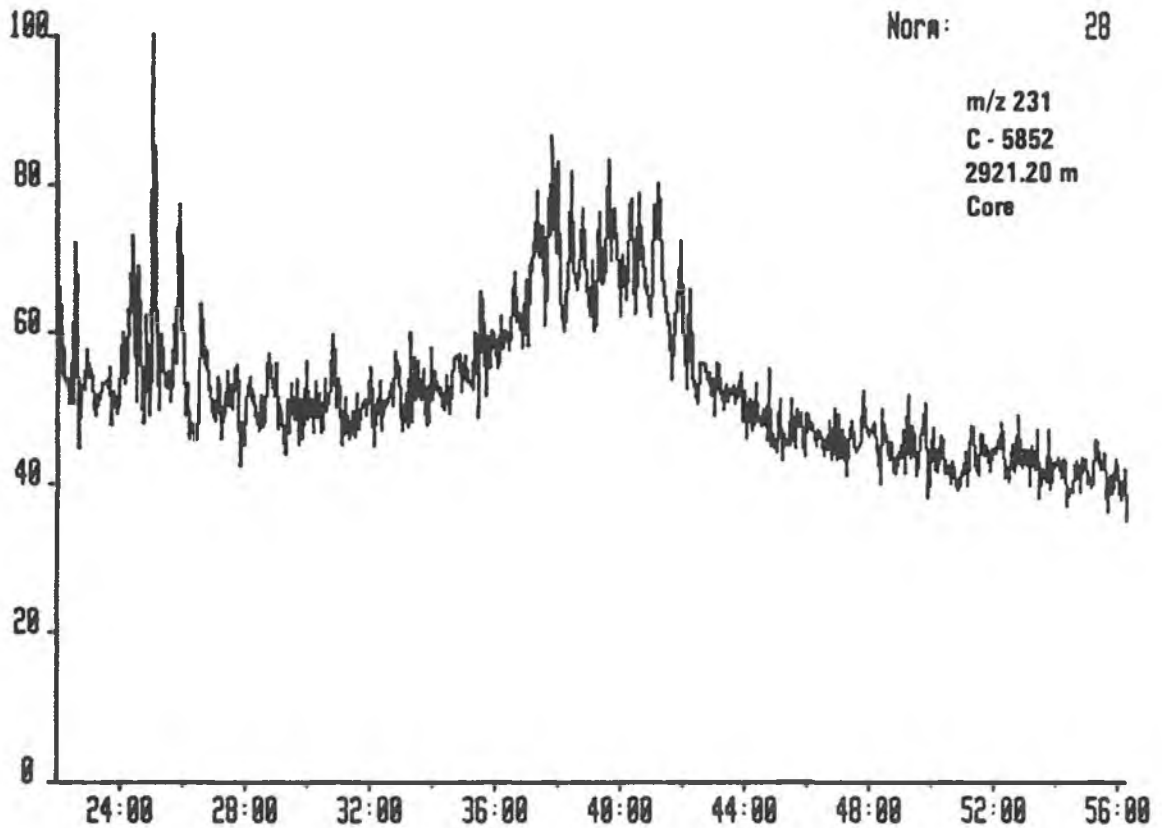


C5849SAT 5-FEB-07 Sir:Voltage 7070H Acnt:IKU System:TRIT
Sample 1 Injection 1 Group 1 Mass 259.1000
Text:



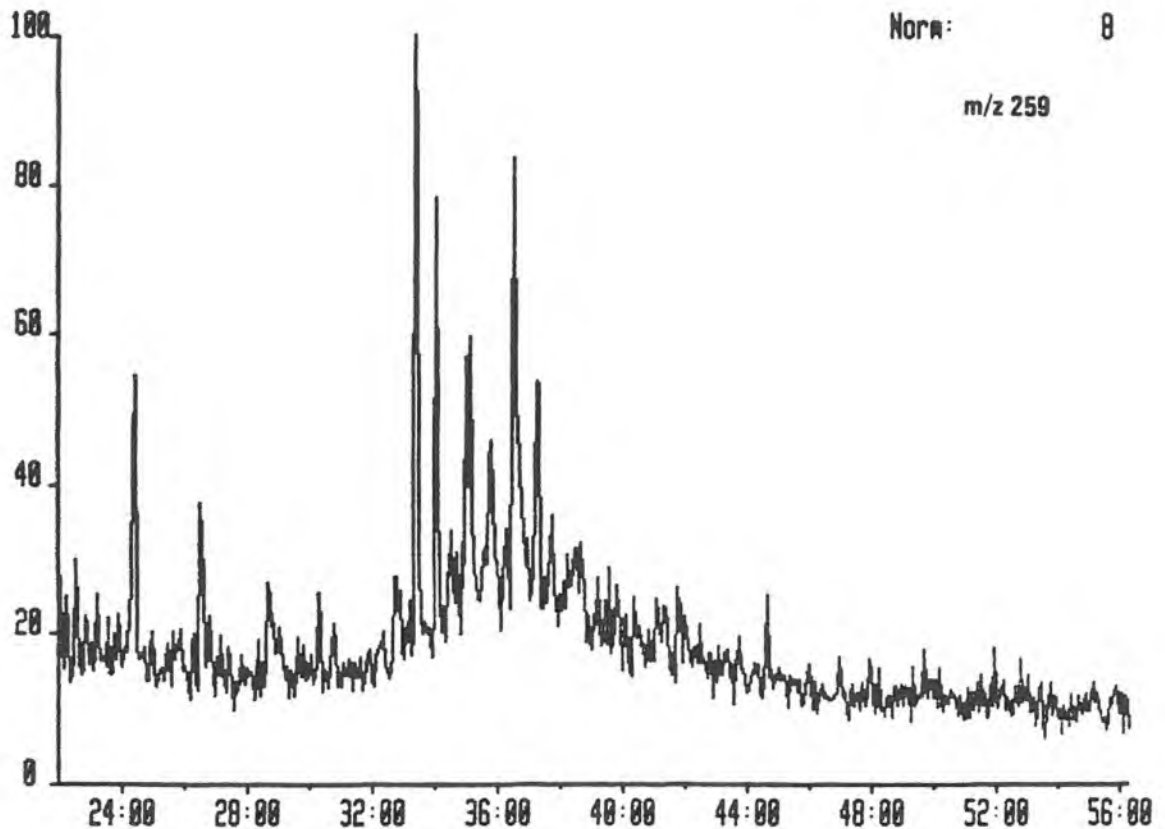
C5852SAT 5-FEB-07 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 231.1000
Text:

System:TRIT



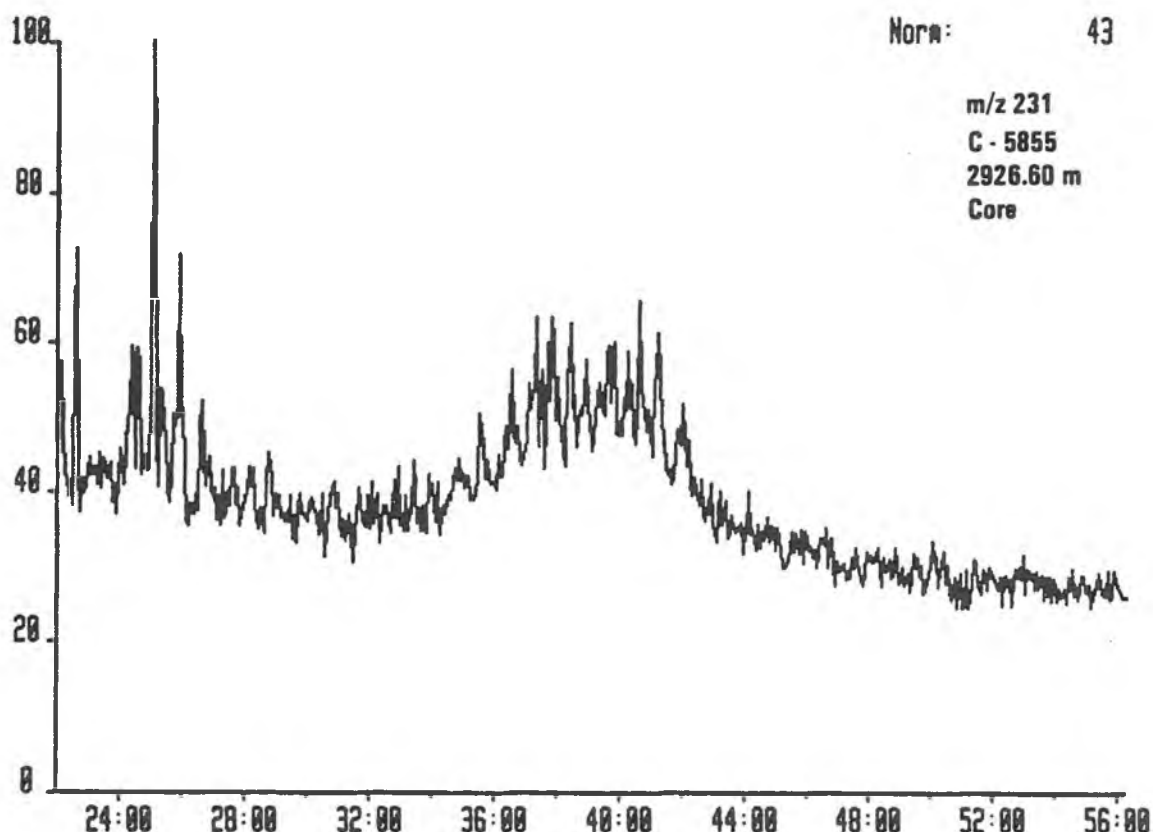
C5852SAT 5-FEB-07 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 259.1000
Text:

System:TRIT



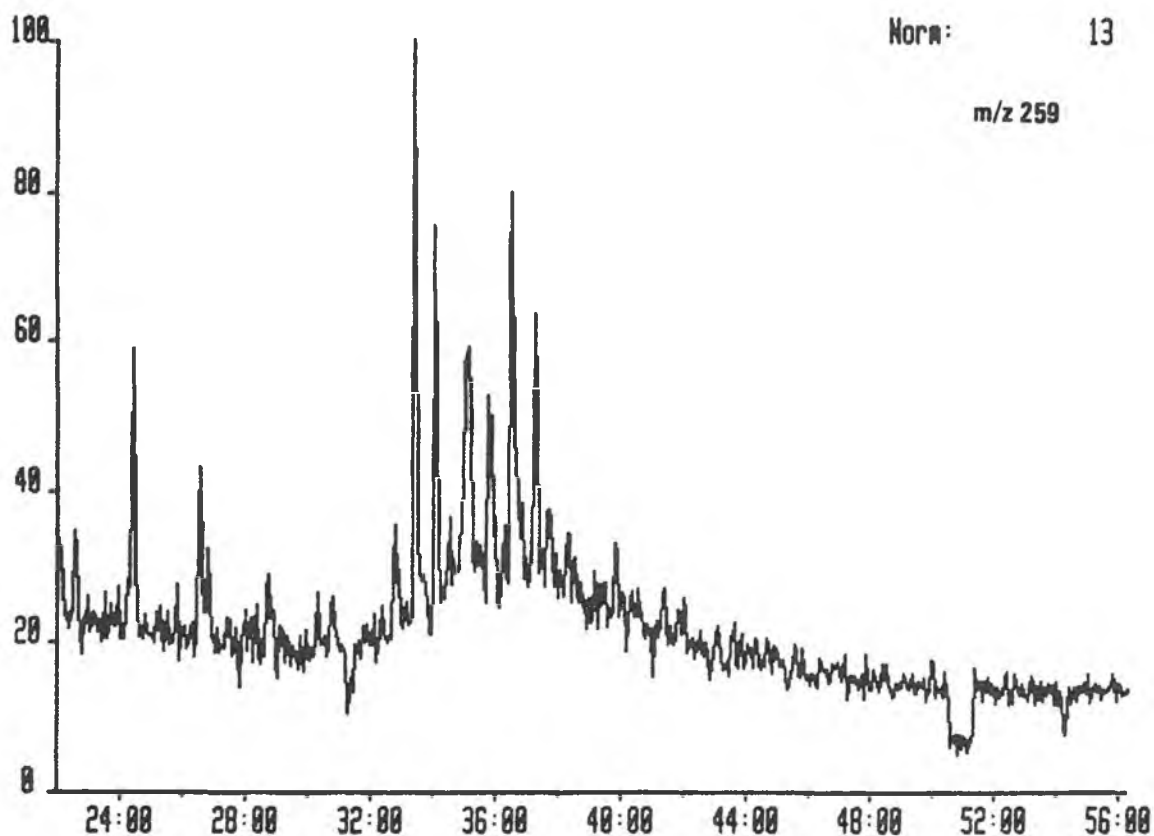
C5855SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 231.1000
Text:

System:TRIT



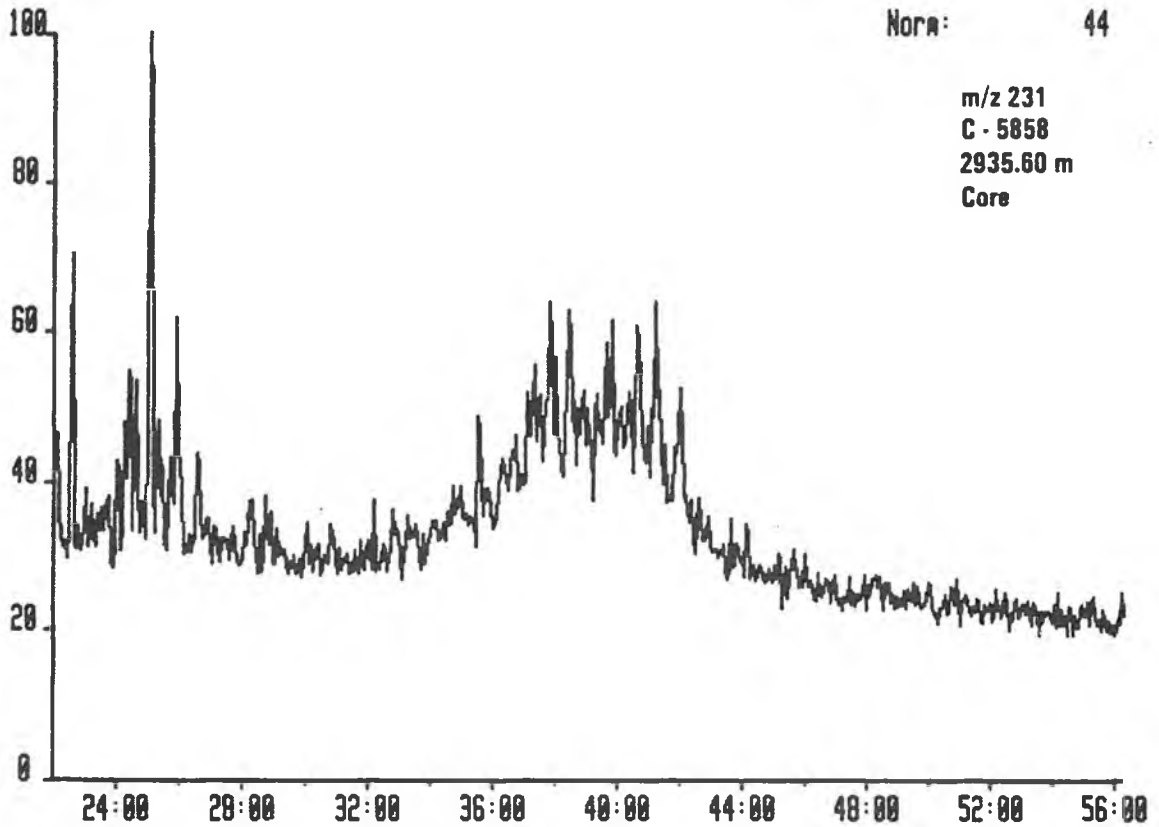
C5855SAT 5-FEB-87 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 259.1000
Text:

System:TRIT



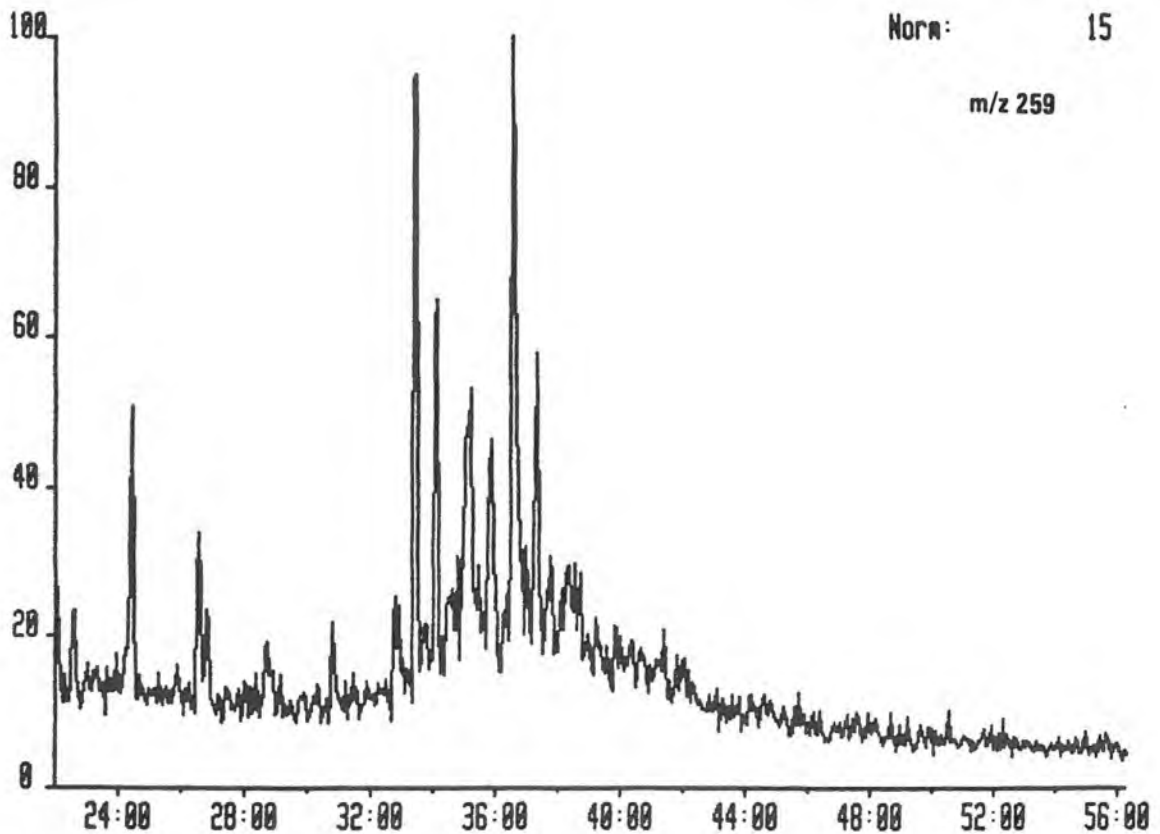
C5858SAT 6-FEB-87 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 231.1000
Text:

System:TRIT

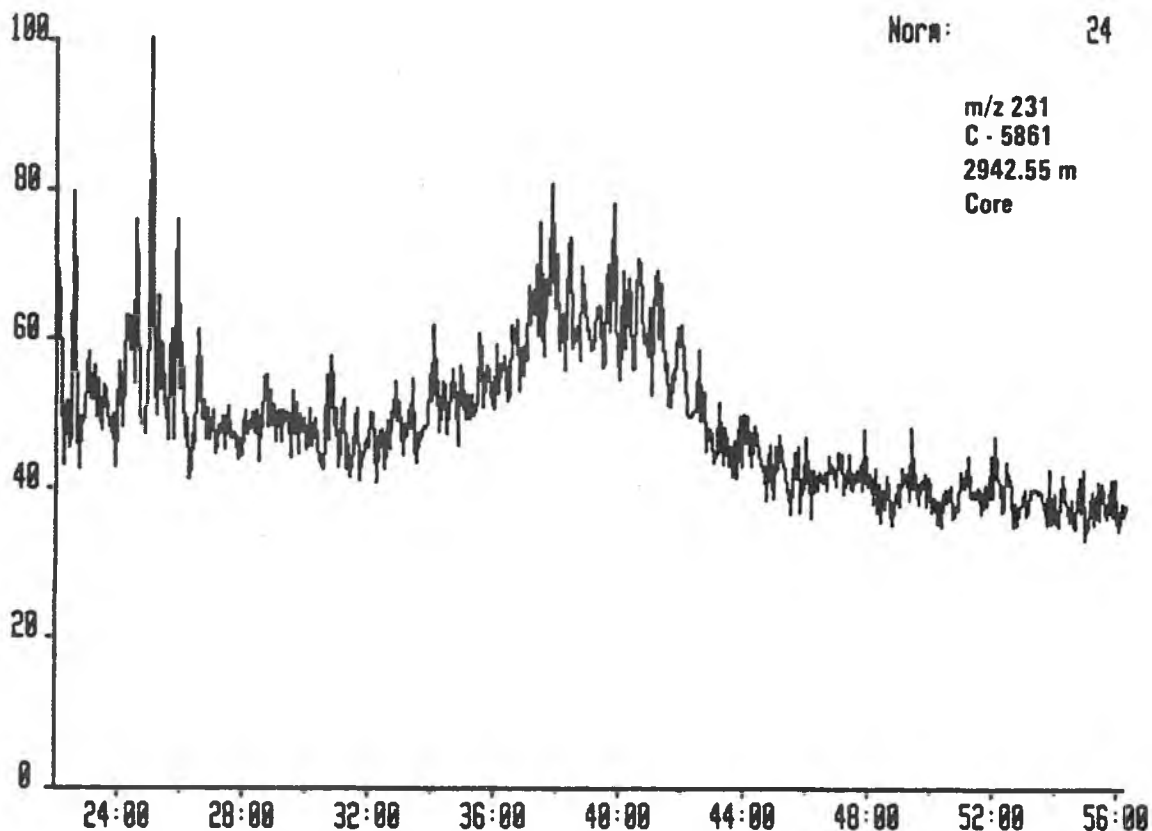


C5858SAT 6-FEB-87 Sir:Voltage 7070H Acnt:IKU
Sample 1 Injection 1 Group 1 Mass 259.1000
Text:

System:TRIT



C5861SAT 6-FEB-07 Sir:Voltage 7070H Rcnt:IKU System:TRIT
Sample 1 Injection 1 Group 1 Mass 231.1000
Text:



C5861SAT 6-FEB-07 Sir:Voltage 7070H Rcnt:IKU System:TRIT
Sample 1 Injection 1 Group 1 Mass 259.1000
Text:

