

Stratospheric Methane Measurements and Predictions

By M. ACKERMAN, D. FRIMOUT, C. MULLER¹⁾ and D. J. WUEBBLES²⁾

Abstract – A new determination of stratospheric methane from 22 km to 35 km altitude with implications on the abundance of this constituent at greater heights is presented. Previous measurements, some of which showed large discrepancies with currently admitted values, have been reinterpreted and brought into agreement. The results are in contradiction according to present theories with the *in situ* determined CH₄ abundances at the upper edge of the stratosphere. Implications on the vertical transport coefficient used in one dimensional models are discussed.

Key words: Stratosphere; Methane.

Introduction

The first determinations of the abundance of CH₄ in the stratosphere were published by BAINBRIDGE and HEIDT [1] in 1966. They showed a decrease of the mixing ratio above the tropopause and were based on *in situ* sampling with subsequent gas chromatography. Using infrared absorption measurements in the 7.7 μm band, KYLE *et al.* [2] also showed a decrease of the mixing ratio above the tropopause. ACKERMAN *et al.* [3] indicated also a decrease of concentration with altitude in the stratosphere as well as CUMMING and LOWE [4] and BURKERT *et al.* [5]. From aircraft borne spectrometer and interferometer data FARMER *et al.* [6] and LOWE and MCKINNON [7] showed a decrease of the CH₄ mixing ratio from low to high latitude in the low stratosphere. The analysis of air samples collected by means of rockets in 1968 and in 1972 have led to determinations of the methane abundance near the stratopause (EHHALT *et al.* 1975 [8]).

Since CH₄ cannot be produced in the stratosphere, its abundance at various altitudes depends on the rate of its chemical destruction and the strength of vertical mixing. This latter property is a basic data for photochemical modeling of the stratosphere. Its knowledge allows, for instance, the evaluation of the rate of transport of aircraft effluents and other contaminants to the chemically ozone sensitive regions of the upper stratosphere. Several authors (NICOLET and PEETERMANS [9],

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WOFSY and McELROY [10], HUNTEN [11]) have attempted to deduce values of the vertical transport parameters using the available methane data. CHANG [12] has analysed the various values and shown that the improvement of experimental data can reduce the present uncertainty and consequently the uncertainties in the models and in their predictions.

The present work is an attempt to reduce the uncertainty. It consists of the determination of the abundance of CH_4 from 22 to 40 km, using high resolution infrared absorption spectroscopic measurements, interpreted on the basis of the presently available laboratory spectroscopic data, followed by a discussion of other results and of theoretical evaluations.

Instrumentation and observation

A 60 cm focal length grille spectrometer was flown on 2 October 1975 from the CNES range in Aire sur l'Adour to measure the vertical distribution of HCl in the stratosphere (ACKERMAN *et al.* [13]). Observations of the solar spectrum from an altitude of 35 km took place in the wave numbers range from 2916 cm^{-1} to 2970 cm^{-1} , which includes the P_5 to P_9 multiplets of the $3.3 \mu\text{m}$ band of CH_4 (Fig. 1). The range of solar zenith angles from $80^\circ 5'$ to $95^\circ 5'$ was covered at 43°N and 2°E . The methane absorptions shown in Fig. 1 were always present in the spectra. Those due to P_9 are perturbed by solar absorption features and are not considered in this work. The measured equivalent widths of the multiplets are shown in Fig. 2 versus elevation angles for observation above the horizontal and listed in Table 1 versus grazing altitude of the solar rays for observations below the horizontal.

Interpretation of the data

(a) Observations above the horizontal

Values of the number density, and of the scale height, have been assumed to compute equivalent widths by dividing the atmosphere in layers of 1 km thickness to take into account the change of path length, pressure and temperature as a function of height in the successive layers above float altitudes according to the earth geometry, and to the mid-latitude Spring-Fall model of the U.S. Standard Atmosphere supplement, 1966. A Voigt profile has been introduced in the computation for each line of the multiplets of which the spectroscopic data, taken from TOTH *et al.* [14] and from McCLATCHEY *et al.* [15], are given in Table 2. The change of equivalent width with solar elevation angle for the four multiplets has been computed and is shown in Fig. 2 with the data. The shaded curves marked 1 represent the equivalent widths

that should be observed if the number density at 35 km was equal to $(1.46 \pm 0.12) \times 10^{11} \text{ cm}^{-3}$ and if the scale height was equal to 4.54 km. These values correspond to the mixing ratio values published by EHHALT *et al.* [8] and used in models to evaluate the strength of the vertical transport of methane. The range of values taken for the number density at 35 km is obtained from the mixing ratio value published by EHHALT *et al.* [8] and from the extreme values of total number densities given in the

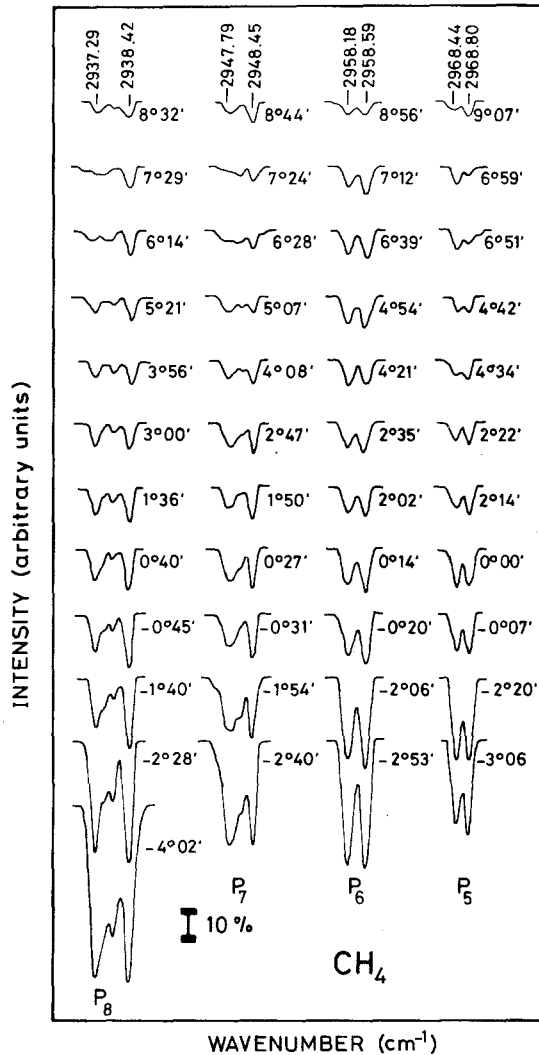


Figure 1

Portions of absorption spectra recorded on 2 October 1975, from 35 km altitude at 43°N and 2°E in the 3.3 μm CH₄ band. Multiplets P₅ to P₈ are shown with some identification wavenumbers in cm^{-1} . The angular values shown on the right-hand side of each portion of spectrum are solar elevation and solar depression angles for positive and negative values respectively.

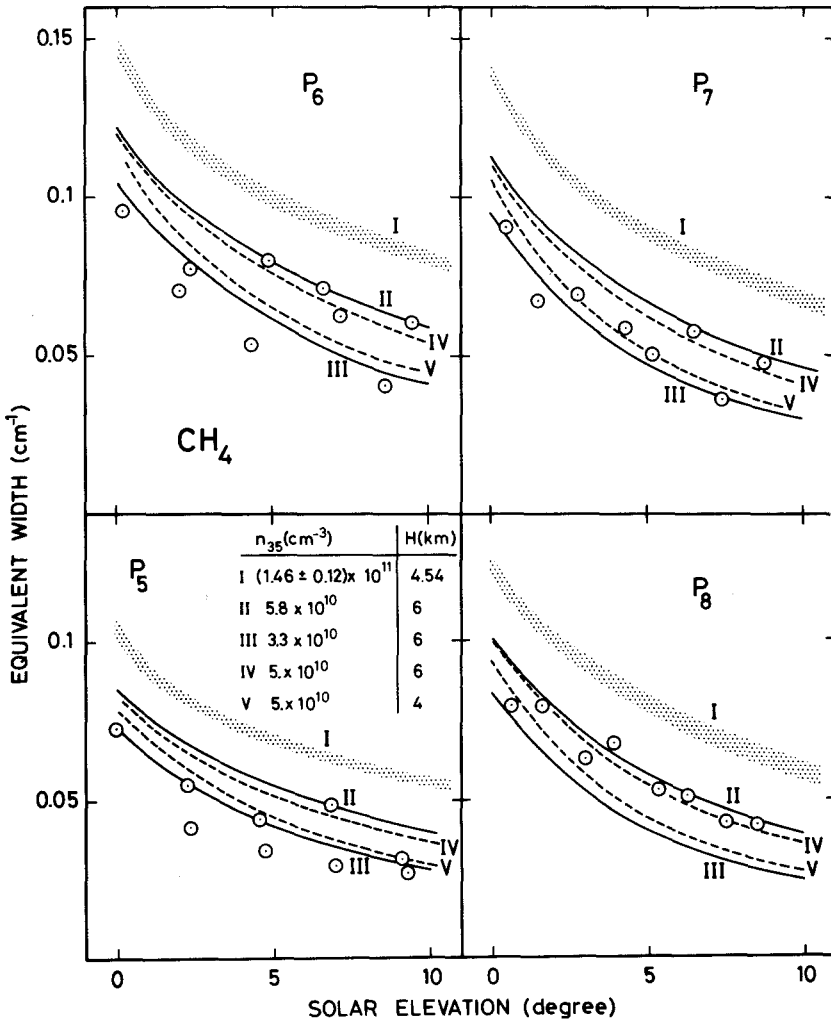


Figure 2

Equivalent width versus solar elevation angle. The circles represent the measurements. The shaded curves correspond to the mixing ratio values represented by the solid broken curve of Fig. 4. The other curves correspond to the CH₄ number densities at 35 km [$n_{35}(\text{cm}^{-3})$] and to the scale heights above this altitude H (km) indicated on the figure.

U.S. Standard Atmosphere Supplement, 1966. The computed equivalent widths do not fit the experimental data points. The computation has then been made for various sets of n_{35} and scale heights. Changing the scale height values from 4.5 km to 6.0 km has the most appreciable effect on the change of equivalent width with elevation angle at small angle. The scatter of the measured values does not allow us to make a clear choice. However, there is a tendency indicating that the scale height above 35 km is at least over a few kilometers altitude closer to 6 km, so that the

Table 1

Equivalent widths in cm^{-1} of the $3.3 \mu\text{m}$ P branch multiplets of CH_4 versus minimum altitudes, z , reached by the solar rays

z (km)	P_5	P_6	P_7	P_8
35	7.1×10^{-2}	1.06×10^{-1}	6.8×10^{-2}	8.5×10^{-2}
33	1.16×10^{-1}	1.64×10^{-1}	1.33×10^{-1}	1.10×10^{-1}
31	1.53×10^{-1}	2.19×10^{-1}	2.00×10^{-1}	1.52×10^{-1}
29	2.11×10^{-1}	2.76×10^{-1}	2.75×10^{-1}	2.10×10^{-1}
27			3.70×10^{-1}	2.82×10^{-1}
25				3.66×10^{-1}
23				4.91×10^{-1}
21				6.41×10^{-1}

Table 2

Line positions λ , integrated absorption cross sections, S , full width at half height W and ground level Energies E for the CH_4 multiplets considered in the treatment of the data.

	λ (cm^{-1})	S ($\text{cm}^2 \text{cm}^{-1}$) 297°K	W (cm^{-1})	E (cm^{-1})
P_5	2968.885	7.06×10^{-20}	0.122	157
	2968.770	1.27×10^{-21}	0.094	157
	2968.738	7.06×10^{-20}	0.122	157
	2968.473	7.06×10^{-20}	0.122	157
	2968.404	7.06×10^{-20}	0.122	157
P_6	2958.683	4.40×10^{-20}	0.114	219
	2958.651	6.58×10^{-20}	0.118	219
	2958.537	1.10×10^{-19}	0.102	219
	2958.233	6.62×10^{-20}	0.118	219
	2958.120	6.42×10^{-20}	0.118	219
	2958.017	1.12×10^{-19}	0.102	219
	P_7	2948.478	5.13×10^{-20}	0.114
2948.421		5.00×10^{-20}	0.114	293
2948.130		1.13×10^{-21}	0.102	219
2948.107		8.47×10^{-20}	0.114	293
2947.912		4.88×10^{-20}	0.114	293
2947.810		3.28×10^{-20}	0.114	293
2947.700		1.13×10^{-21}	0.102	219
2947.668		5.00×10^{-20}	0.114	293
P_8		2938.248	6.46×10^{-20}	0.110
	2938.206	3.88×10^{-20}	0.110	376
	2938.206	2.59×10^{-20}	0.110	376
	2937.769	3.88×10^{-20}	0.110	376
	2937.494	3.88×10^{-20}	0.110	376
	2937.307	2.70×10^{-20}	0.110	376
	2937.234	3.95×10^{-20}	0.110	376

values corresponding to curves II and III of Fig. 2 have been represented in Figs. 3 and 4 where envelopes are shown at and above 35 km for the CH_4 number densities and mixing ratios respectively with the results of other investigators.

(b) *Observations below the horizon*

The amount of methane above 35 km has been taken into account by choosing a scale height equal to 6 km and a number density of 4.5×10^{10} at 35 km, both values being the result of the preceding section. The number densities below float altitude in successive layers of 2 km thickness have been determined from the equivalent widths listed in Table 1. The same atmospheric model and spectroscopic data as in the previous sections have been introduced in the computation. The results are shown in Figs. 3 and 4 as number densities and volume mixing ratios respectively.

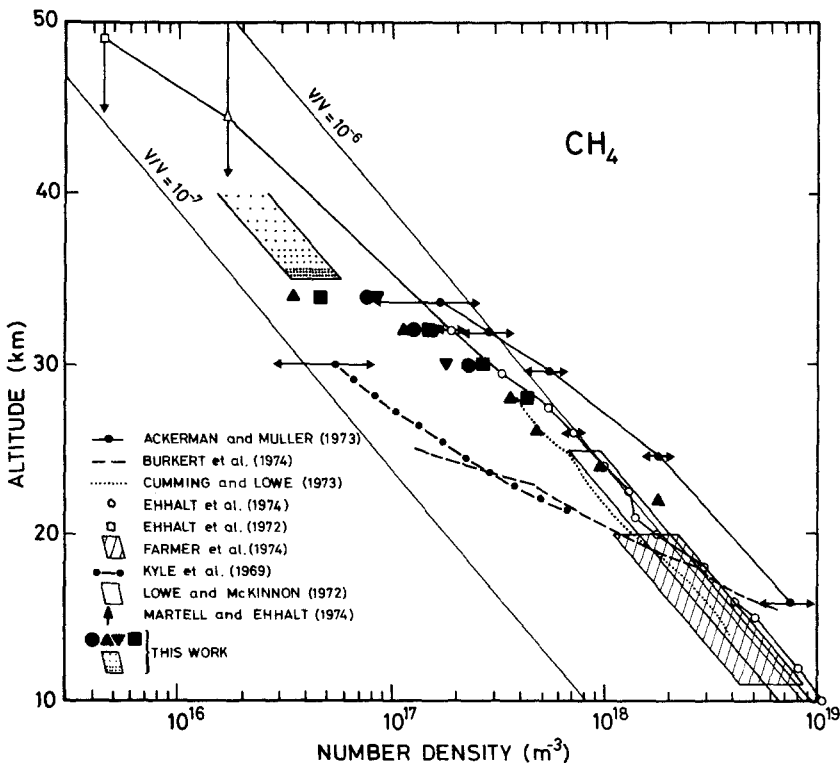


Figure 3

Methane number densities as published by various groups of authors and reported by HARD [19] with the values deduced from the observations reported in the present work. The arrowed vertical lines centered on the values between 40 and 50 km indicate the altitude range of vertical rocket sampling.

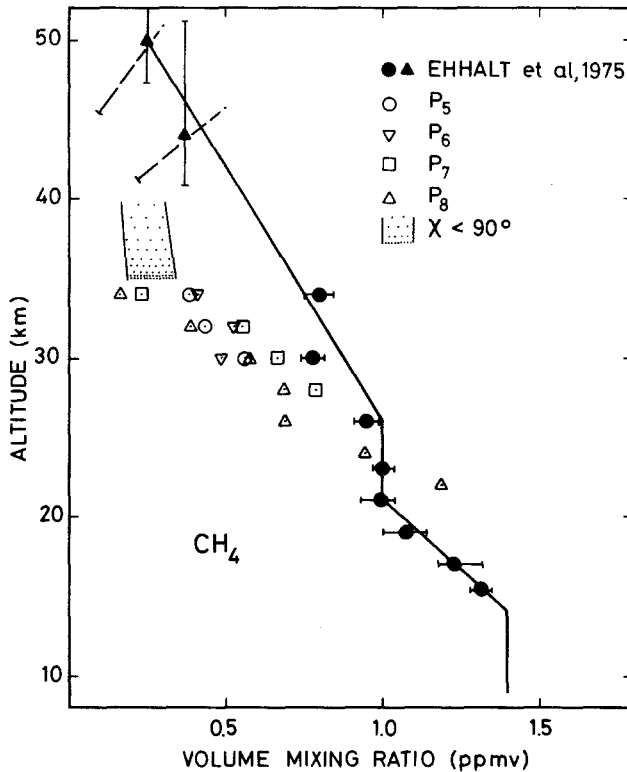


Figure 4

Methane mixing ratio as a function of altitude. The presently reported observations are shown with the values from EHHALT *et al.* [8]. Tilted dashed bars are shown with the rocket determinations in order to keep this figure consistent with the previous one (Fig. 3).

Comparison with other results

Four sets of spectroscopic data are available about stratospheric methane. They cover a period of 10 years since the measurements published by KYLE *et al.* [2] were performed in December 1967 at 33°N. The ν_4 band, from 1299 cm^{-1} to 1306 cm^{-1} , was observed in absorption during a balloon ascent for solar elevation angles from 31° to 5° . The authors give all information required to reinterpret their observations. After ten years the quality of spectroscopic parameters has greatly improved. This situation justifies a new analysis. This has been made, following the method already described previously on the basis of the spectroscopic data of MCCLATCHEY *et al.* [15] listed in Table 3. However, nitrous oxide absorbs in the region from 1299.1 cm^{-1} to 1307 cm^{-1} which is considered. For this reason, the analysis has been limited to the balloon float period. A number density of N_2O at 28 km equal to $6 \times 10^{10}\text{ cm}^{-3}$ has been assumed with a scale height of 3.7 km leading to a maximum contribution to

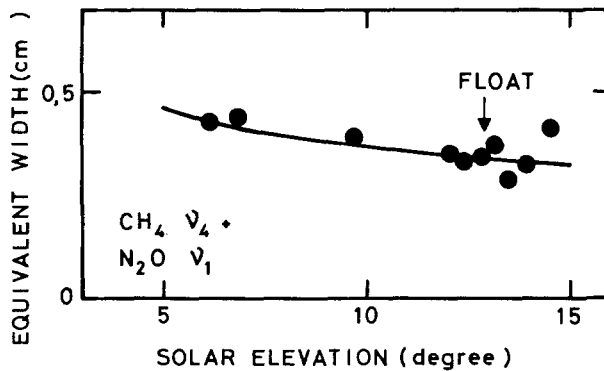
Table 3
*Spectroscopic data used for the ν_4 CH₄ band. The symbols
 have the same meaning as in Table 2.*

λ (cm ⁻¹)	S_{297} (cm ² cm ⁻¹)	W (cm ⁻¹)	E (cm ⁻¹)
1299.638	2.35×10^{-20}	0.112	293.10
1299.678	1.71×10^{-21}	0.098	950.27
1299.903	2.35×10^{-20}	0.112	293.11
1299.903	8.32×10^{-21}	0.104	575.14
1300.282	6.21×10^{-21}	0.130	10.48
1300.467	8.33×10^{-21}	0.104	575.23
1300.467	1.71×10^{-21}	0.098	950.30
1300.748	2.99×10^{-22}	0.096	1251.70
1301.208	2.05×10^{-21}	0.100	814.61
1301.290	4.48×10^{-22}	0.096	1251.80
1301.371	1.94×10^{-20}	0.112	219.90
1301.500	8.37×10^{-21}	0.106	470.77
1301.550	2.92×10^{-20}	0.116	219.92
1301.550	3.07×10^{-21}	0.100	814.97
1301.825	1.25×10^{-20}	0.106	470.78
1302.040	4.86×10^{-20}	0.100	219.90
1302.040	9.01×10^{-22}	0.098	1095.60
1302.451	5.21×10^{-21}	0.102	689.93
1302.774	1.78×10^{-20}	0.110	376.80
1302.774	3.35×10^{-20}	0.120	157.11
1302.774	2.85×10^{-21}	0.098	950.35
1302.945	2.09×10^{-20}	0.106	470.81
1303.198	1.39×10^{-20}	0.104	575.19
1303.290	3.67×10^{-22}	0.122	000.00
1303.373	3.35×10^{-20}	0.120	157.12
1303.567	3.93×10^{-20}	0.112	293.13
1303.709	5.13×10^{-21}	0.100	815.05
1303.709	5.90×10^{-20}	0.092	104.77
1303.709	7.49×10^{-22}	0.096	1252.00
1303.951	1.78×10^{-20}	0.110	376.71
1304.223	3.54×10^{-20}	0.124	104.77
1304.332	5.22×10^{-21}	0.102	689.99
1304.463	2.36×10^{-20}	0.112	293.15
1304.602	9.03×10^{-22}	0.098	1095.60
1304.602	2.36×10^{-20}	0.108	104.77
1304.602	8.35×10^{-21}	0.104	575.15
1304.847	1.71×10^{-21}	0.098	950.45
1304.847	3.38×10^{-20}	0.124	62.87
1305.002	2.92×10^{-20}	0.116	219.90
1305.002	1.57×10^{-20}	0.112	293.15
1305.092	1.26×10^{-20}	0.106	470.83
1305.286	3.08×10^{-21}	0.100	814.83
1305.286	5.57×10^{-21}	0.104	575.02
1305.413	3.38×10^{-20}	0.124	62.87
1305.454	1.88×10^{-20}	0.102	31.43
1305.454	4.50×10^{-22}	0.096	1252.00
1305.563	2.24×10^{-20}	0.120	157.13
1305.563	1.14×10^{-21}	0.098	950.47

Table 3—continued.

λ (cm ⁻¹)	S_{297} (cm ² cm ⁻¹)	W (cm ⁻¹)	E (cm ⁻¹)
1305.635	2.92×10^{-20}	0.116	219.93
1305.635	1.19×10^{-20}	0.110	376.71
1305.688	2.81×10^{-20}	0.130	31.44
1305.809	3.48×10^{-21}	0.102	690.01
1305.809	1.87×10^{-20}	0.130	10.48
1305.809	1.26×10^{-20}	0.106	470.84
1305.905	3.36×10^{-20}	0.120	157.13
1305.989	3.55×10^{-20}	0.124	104.77
1305.989	6.03×10^{-22}	0.098	1095.20
1305.989	3.08×10^{-21}	0.100	814.61
1306.021	1.78×10^{-20}	0.110	376.76
1306.106	2.36×10^{-20}	0.112	296.16
1306.106	5.64×10^{-20}	0.112	62.87
1306.254	4.88×10^{-20}	0.100	219.93
1306.254	4.50×10^{-22}	0.096	1252.00
1306.254	8.36×10^{-21}	0.104	575.02
1306.254	5.23×10^{-21}	0.102	690.02
1306.428	2.10×10^{-20}	0.106	470.85
1306.428	1.72×10^{-21}	0.098	950.49
1306.627	9.04×10^{-22}	0.098	1095.20
1306.627	5.14×10^{-21}	0.100	814.61
1306.832	7.51×10^{-22}	0.096	1252.00

the equivalent width equal to 0.1 cm^{-1} . The observations are then, as shown in Fig. 5, well represented if methane number density and scale height are respectively taken equal to $2 \times 10^{11} \text{ cm}^{-3}$ and 4 km. This represents an increase by a factor of almost 4 of the original values published by KYLE *et al.* [2] bringing them in agreement with others.



Equivalent width observed at balloon float altitude by KYLE *et al.* [2] before sunset versus solar elevation angle. The curve corresponds to a methane number density at 30 km equal to $2 \times 10^{11} \text{ cm}^{-3}$ associated with a scale height of 4 km. The N_2O absorption of 0.1 cm^{-1} equivalent widths has been taken into account, corresponding to an N_2O number density at 30 km equal to $6 \times 10^{10} \text{ cm}^{-3}$ associated with a scale height of 3.7 km.

Enough flight data were given by CUMMING and LOWE [4] in their publication to reinterpret their observation made at 47°N in August 1965 and at 28 km float altitude. A computation was made using the now available spectroscopic data on the components of the R_5 multiplet of the $3.3 \mu\text{m}$ ν_3 band. As shown in Fig. 6, a CH_4 number density and a scale height equal to $3 \times 10^{11} \text{ cm}^{-3}$ and 4.5 km respectively at 28 km, fit the data points and are in good agreement with the original result of the authors.

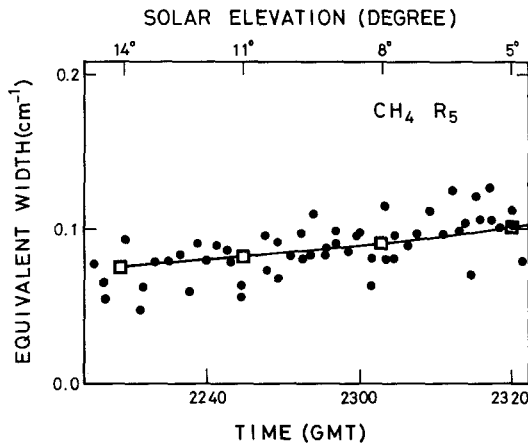


Figure 6

Equivalent width due to the $R_5\text{CH}_4$ multiplet observed by CUMMING and LOWE [4] versus time during the float period of the gondola. The circles and the solid line are from the original publication. The squares correspond to the reinterpretation with a CH_4 number density at 28 km altitude equal to 3×10^{11} associated with a scale height equal to 4.5 km, both being in good agreement with the original interpretation.

In the spectra obtained by ACKERMAN *et al.* [3] in October 1971 at 43°N, the Q branch of the $3.3 \mu\text{m}$ band of CH_4 is the most prominent feature which at the time of the publication could not be analysed due to the lack of availability of a powerful enough computer. This has now partly been done. The equivalent width equal to $0.8 \pm 0.1 \text{ cm}^{-1}$ observed at 35 km for a zenith angle equal to 90° is represented by a CH_4 number density equal to $(4.0 \pm 1.5) \times 10^{10} \text{ cm}^{-3}$ associated with a scale height equal to 4 km. This brings downward the values published in 1972 which have always appeared to be systematically too high.

Discussion

Four sets of data obtained at various latitudes and seasons are now available at 30 km altitude for a period of ten years. The values derived on the basis of a consistent set of spectroscopic parameters are listed in Table 4 with *in situ* sampling results.

Table 4
Methane number densities at 30 km altitude.

CH ₄ number density (cm ⁻³)	Time	Latitude	Ref.
1.9×10^{11}	August 1965	47°N	[4]
2×10^{11}	December 1967	33°N	[2]
2×10^{11}	October 1971	43°N	[3]
2.2×10^{11}	October 1975	43°N	This work
3×10^{11}	—	32°N	[8]

No significance in terms of geographic, time or season variability can be attributed to the differences appearing between the various values, since the uncertainties are most probably larger than these differences. Methane number density at 30 km appears to be very stable within experimental error limits.

The shaded area in Fig. 7 shows the range of the methane mixing ratio presented in this paper, along with theoretical values predicted by the LLL one-dimensional model. This model is based on a unified chemical scheme including 84 reactions among 23 species. Results for three different profiles of the vertical transport coefficients represented in Fig. 8 are shown. The curve labelled CHANG [20] is based on the transport coefficient derived in the NAS Report (1976). The curves labelled CRUTZEN and ISAKSEN [21] and HUNTEN [22] are also shown for comparison.

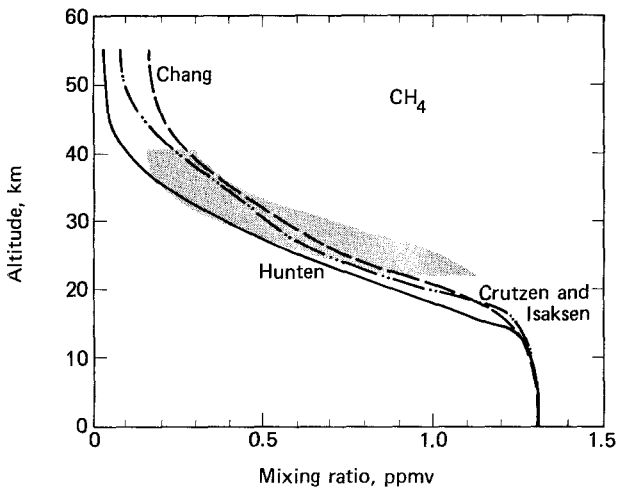


Figure 7

Vertical distribution of the CH₄ volume mixing ratio computed on the basis of various values of vertical transport coefficient [20–22].

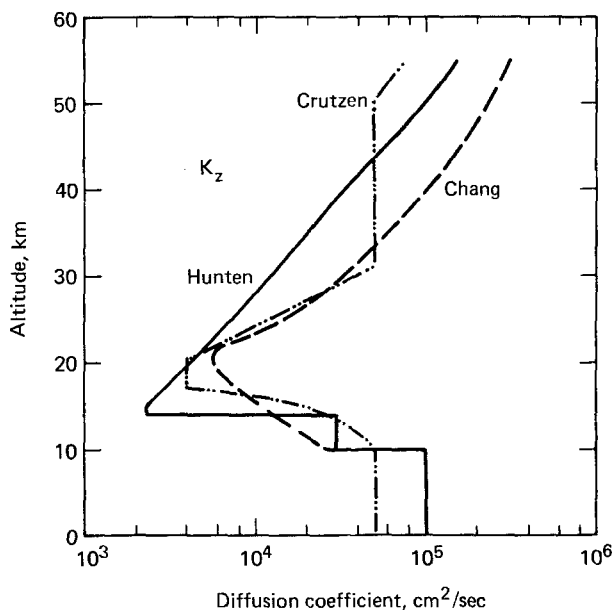
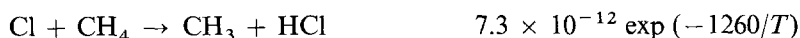
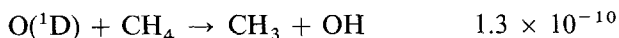
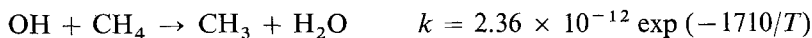


Figure 8
Vertical transport coefficient proposed by various authors [20-22].

The important chemical reactions determining the theoretical distribution of methane are



The reaction rates, k , are those recommended by NASA [23]. The concentrations of the reactants are given in Table 5. Only a few measurements of OH and Cl exist in the upper stratosphere and no measurements exist at all of O(¹D) to confirm the model derived concentration for these species. Thus, the relative fit to the CH₄ profile of results with different vertical transport coefficients is only of limited use in determining which set of coefficients is most satisfactory.

Low stratospheric methane abundances will have various consequences on model calculation of other species. As an example, they will have the effect of reducing the HCl/ClO ratio.

Conclusion

New stratospheric methane measurements have been presented with previous measurements reinterpreted on the basis of a consistent set of spectroscopic para-

Table 5
Selected concentrations (cm⁻³) for important species to CH₄ loss rate as derived from the LLL 1-D model with Chang, 1976 K₂.

Altitude (km)	OH	Cl	O(¹ D)
10	8.56(+5)	1.08(+3)	1.67(-2)
11	1.03(+6)	1.83	2.30
12	1.57	3.47	3.16
13	2.16	5.78	4.35
14	2.23	7.41	6.01
15	2.27	9.15	8.35
16	2.25	1.08(+4)	1.17(-1)
17	2.18	1.24	1.65
18	2.06	1.37	2.35
19	1.92	1.49	3.40
20	1.81	1.62	5.00
21	1.75	1.74	7.49
22	1.76	1.88	1.12(+0)
23	1.84	2.05	1.63
24	2.00	2.27	2.29
25	2.24	2.58	3.19
26	2.57	2.96	4.44
27	3.00	3.46	6.21
28	3.56	4.11	8.70
29	4.27	4.95	1.22(+1)
30	5.14	6.06	1.71
31	6.20	7.52	2.38
32	7.48	9.40	3.26
33	8.94	1.18(+5)	4.39
34	1.06(+7)	1.46	5.74
35	1.25	1.79	7.33
37.5	1.78	2.80	1.25(+2)
40	2.31	4.02	2.02
42.5	2.54	4.97	2.91
45	2.34	4.88	3.35

meters. Discrepancies with values obtained at the upper edge of the stratosphere by means of *in situ* sampling calls for new measurements, particularly at those altitudes.

Acknowledgement

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