

Solar Irradiance Measurement and Estimation of Aerosol and Ozone in Antarctica During Austral Summer of 1991-92

S.D. SHARMA

**National Physical Laboratory
K.S. Krishnan Road, New Delhi -110 012**

Abstract

Solar irradiance has been measured in Antarctica during the eleventh Indian expedition using a number of instruments covering a range from 295nm to 14.5 μ m i.e. UVB, visible and infrared. The measurements have been analysed to estimate the aerosol content, aerosol size distribution and gross total ozone in the Antarctic atmosphere. Results analysed by computer, have been discussed separately depending upon the wavelength range under contention. The results show increasing trend of aerosols over the past years, higher ozone presence compared to that of Delhi and ozone values comparable to those obtained from TOMS.

Introduction

Recently the world has seen a sudden awakening to global changes caused by increase of green house gases and depletion of ozone catalyzed by anthropogenic gases. The discovery of "ozone-hole" by Farman et al.(1985) has proved a milestone in global awareness of the ecological problems faced by the planet. During the eleventh Indian expedition to Antarctica, solar irradiance covering a range from 295nm (UVB) to 14.5 (μ m (IR) was recorded by using a number of instruments like Sunphotometer (368nm, 500nm, 675nm, 778nm), UVB Sunphotometer (295nm, 392nm, 310nm), Spectroradiometer (280nm to 368nm) and IR Sunphotometer (2.5m - 14.5m). Different parts of the spectrum provide information about different aspects of the problem.

Experimental Set-up

The following apparatus were used for the observations.

(a) Sunphotometer

It is 'EKO'Japan make instrument which can operate at four different wavelengths viz., 368 nm (UV), 500 nm (visible), 675 nm (visible) and 778 nm (IR). It can be operated by holding in hand or fixing the instrument on a stand. It has four narrow band filters to measure solar irradiance at the above mentioned four wavelengths. The filter positions can be changed by rotating a knob which also indicates the kind of filter in line. In order to take the observations the instrument is pointed towards the Sun, orientation is adjusted so that the light spot falls inside the specified circle. At this position the Sun is set in the optimum alignment. The instrument is set in 'Hold' position so that it records the maximum value of the signal around the optimum setting. Depending upon the intensity of the signal, instrument can be set at gain/gain10 positions. Gain10 position multiplies the incoming signal by 1000. The measurements have been taken at hourly interval for 8 hours on clear days. The instrument is also provided with an in-built thermometer to note the ambient temperature at the time of observation which is required for temperature correction while deriving the turbidity.

ih) UV - B Sunphotometer

The instrument is designed and developed at the National Physical Laboratory, New Delhi. It consists of three filters with peak wavelengths at 310nm, 302nm and 295nm. The radiation after passing through filters falls on a photo-multiplier tube. Filters are brought into the field of view by a lever as it is pushed stepwise into the instrument. The P.M. tube is operated at 900VDC to record the signal. The instrument can be aligned towards the Sun using the movements along horizontal and vertical axes. Power required for the instrument is provided by a D.C. power supply unit 'Aplab 7333' make. Two types of neutral density filters were used to reduce the intensity of solar radiations to observational level. The output is read on a sensitive microvoltmeter at the maximum output position, which is finally converted into $\text{Watt/cm}^2/\text{nm}$ by multiplying by a calibration factor. The measurements were taken on sunny days at hourly interval between 0800 and 1700 hrs. During the austral summer of 1991 -92 a total of 23 days' observations were taken:

(c) Spectroradiometric measurements

The spectroradiometric Model 742 is procured from Optronic Labs. Inc. USA. it consists of three parts:

- i) Optical head
- ii) Electronic control
- iii) Data logger

It is a versatile instrument which can measure solar irradiance from 200nm to 800nm in a continuous fashion. First, the zero level is adjusted to about 6.0V. Then the optical head is aligned to the Sun and desired wavelength adjusted at the electronic unit. This tunes the optical head to the desired wavelength. Data is printed by the data logger. Because of the partial failure of the system in our case, the data was recorded manually from the display at electronic unit.

Measurements were taken from 04.1.92 to 12.2.92 on sunny days (23 days), five times a day. They were limited to UV - B range of the spectrum viz., 280nm, 290nm, 295nm, 300nm, 305nm, 310nm, 315nm, 320nm and 368nm for cross

(d) Infrared Sunphotometer

The transmission of the atmosphere in IR region is of great importance in view of the study of heat balance, thermal structure, abundance of constituents etc. Principal absorbers of IR are H₂O, CO₂, O₃. In addition to these, CH₄, CO and N₂O occur as minor constituents. N₂ and O₂ have no dipole moment and so possess no IR rotation-vibration bonds. Molecular scattering of radiations is important for smaller wavelengths near visible but for longer wavelengths absorption by atmospheric gases like H₂O, CO₂, O₃, N₂O, CO, CH₄ is important whereas absorption and scattering by aerosols is low. This IR Sunphotometer studies the constituents of air by recording their absorption spectra falling in IR region.

The block diagram of the system is shown in Fig 1. It consists of Sun tracker, circular variable filter, chopper, pyroelectric detector, lock-in amplifier and recorder. The filter is motor driven and allows continuous scanning of wavelengths from 2.5 μm to 14.5 μm in 20 steps. The signal is synchronously detected using lock-in amplifier. A heliostat mounted outside is used to direct sunlight into a 'Newtonian Telescope' which concentrates it on the apparatus constantly.

Results and Discussion

(a) The radiation coming from Sun suffers extinction as it travels through the atmosphere. Atmospheric turbidity is defined as the extinction of direct solar radiation by existing aerosols. The direct solar intensity varies with wavelength, amount of suspended particles and their radii. Aerosol extinction coefficient can be calculated as :-

$$I_{\lambda} = I_0 \lambda e^{-(T_m \lambda + T_O \lambda + T_R \lambda) m} \quad \dots (1)$$

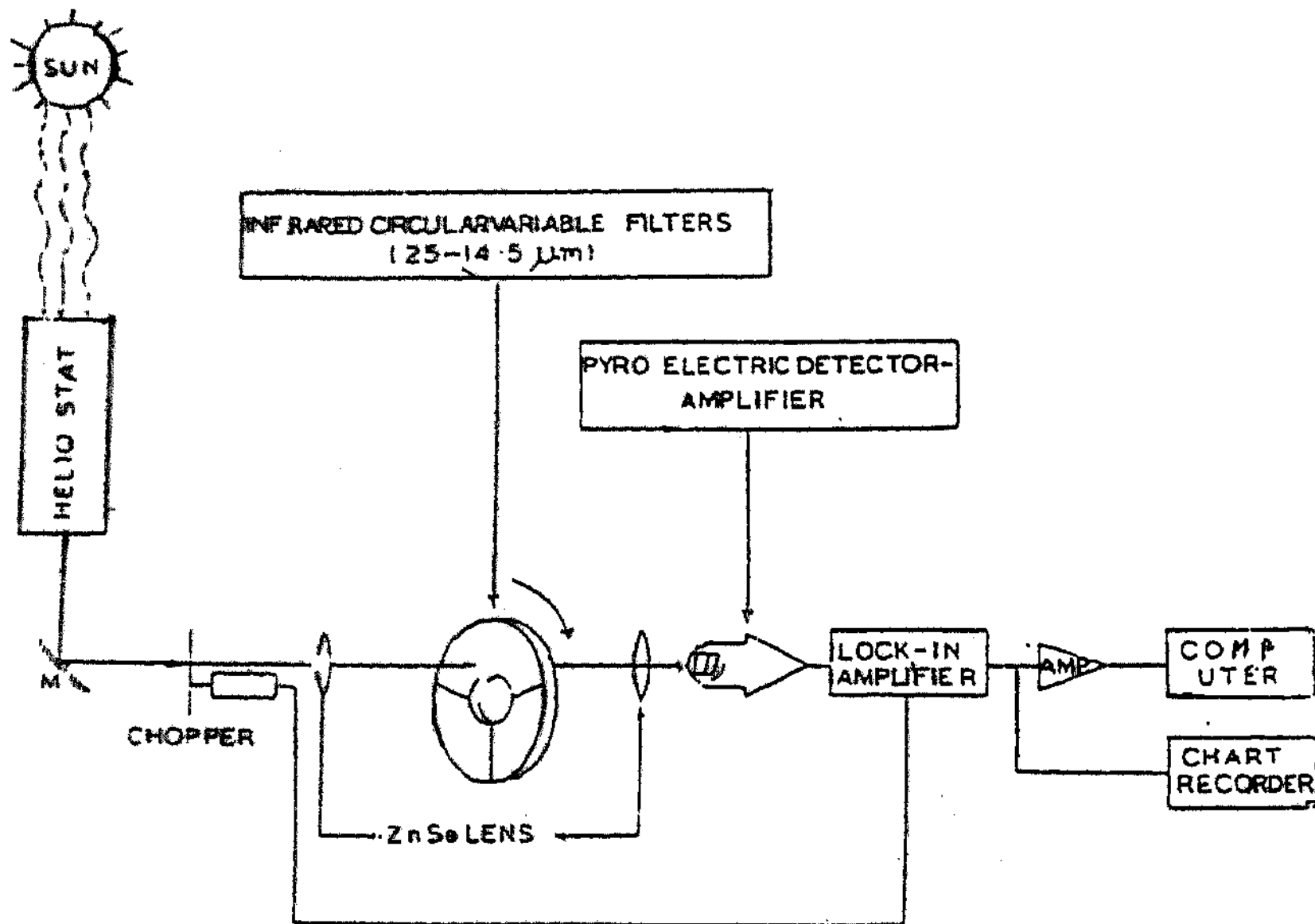


Fig 1; The block diagram of infra-red radiation monitoring system.

where,

I_{λ} = The irradiance at the observing point.

$I_0\lambda$ = The extra-terrestrial irradiance at the mean Sun-Earth distance.

$T_m\lambda$ = The extinction coefficient due to aerosols.

$T_o\lambda$ = The absorption coefficient due to ozone.

$T_R\lambda$ = The Rayleigh scattering coefficient.

m = The absolute airmass.

Sunphotometer does not measure absolute irradiance but a reading proportional to ' I_{λ} '. $I_0\lambda$ is the meter reading at airmass '0'. Values of $T_R\lambda$ and $T_o\lambda$ obtained from literature are as follows:

Wavelength (nm)	368	500	675	778
T_r	0.4945	0.1391	0.0410	0.0231
T_o	0	0.0114	0.0144	0

Also,

$$m = \frac{1}{\text{Sinh } +0.15(h - 3.885)^{-1.253}} \dots\dots(2)$$

where h = solar elevation in degree &

$$\sin h = \sin L \cdot \sin S + \cos L \cdot \cos S \cdot \cos t \quad \dots (3)$$

where L = Latitude of the place

S = Declination of the Sun

t = true solar time = (local time - local noon) $\times 15^\circ$

From equation (1) $T_{m\lambda}$ can be calculated as follows:

$$T_{m\lambda} = 1/m \ln (E_{\lambda} / I_{\lambda} - (T_{o\lambda} + T_{R\lambda})) \dots (4)$$

' λ ' should be such that no absorption takes place by CO₂ and water vapours

For practical determination of T_m , the following equation is more useful than that of (4).

$$T_{m\lambda} = \frac{\ln (E_{\lambda} / E_{\lambda} S)}{m} - (P/P_o T_{R\lambda} + T_{o\lambda})$$

where

$E_{o\lambda}$ = Calibration factor

E_{λ} = Meter reading

P = The station pressure

P_o = The standard pressure at sea level viz. 1013.2 mb

S = Correction factor for mean Sun - Earth distance
(provided with instrument)

The value of $E_{o\lambda}$ is provided with the instrument but it is liable to change with time. It was recalibrated in Antarctica as well as later in N.P.L. The results are as follows :-

Wavelength	Calibration factors		Available with the instrument
	At NPL	At Antarctica	
368 nm	0.016	0.032	0.128
500 nm	0.35	0.40	0.533
675 nm	0.11	0.13	1.054
778 nm	0.59	0.70	0.637

Values at NPL and Antarctica are slightly different on account of haze present over Delhi at the time and also because of vast temperature difference at the two places.

Angstrom's wavelength exponent ' α ' is related to aerosol size distribution. ' α ' can be calculated from extinction coefficients for any pair of filters :

$$\alpha = (\ln T_{m\lambda_1}/T_{m\lambda_2}) / \ln \lambda_2/\lambda_1 \quad \dots (6)$$

27 sunny days for sunphotometric measurements were available during summer of XI Indian expedition to Antarctica. The measurements have been used to calculate extinction coefficients $T_{m\lambda}$ for all the four wavelengths. Six possible ' α ' values have been calculated at different times of day for different values of m . The general trend in ' α ' values has been found as:

$$\alpha_{12} > \alpha_{13} > \alpha_{14} > \alpha_{23} > \alpha_{24} > \alpha_{34}$$

Diurnal variations of $T_{m\lambda}$ vs local time have been plotted for a number of days and shown in Fig 2. The minimum and maximum values of $T_{m\lambda}$ are found as:

Wavelength (nm)	Min. $T_{m\lambda}$	Max. $T_{m\lambda}$
368	0.2883	1.06
500	0.1984	0.9407
675	0.1062	1.03
778	0.1544	1.06

In contrast, during the VI expedition, $T_{m\lambda}$ 368 values varied from 0.04 to 0.16 and for higher wavelengths the atmosphere was clear over Antarctica.

(b) UV - B Sunphotometer measurements were taken for 23 sunny days at Maitri, Antarctica. UV - B energies were calculated from the output in millivolts by multiplying by calibration factors. Calculations were made for estimation of O₃ using the modified expression:

$$I_{\lambda} = I_0 \lambda e^{-(T_{m\lambda} + T_{R\lambda} + B_x)m} \quad \dots (7)$$

which gives

$$x = I/B_{\lambda} (1/m \ln I_0 \lambda / I_{\lambda} - T_{m\lambda} - T_{R\lambda}) \quad \dots (8)$$

Here.

I_{λ} = measured energies

$I_0 \lambda$ = energies at the top of atmosphere

$60.74 \times 10^6 \text{ w/cm}^2 \text{ at } 310 \text{ nm}$

(Ref. Table B - 5. The Stratosphere, 1981)

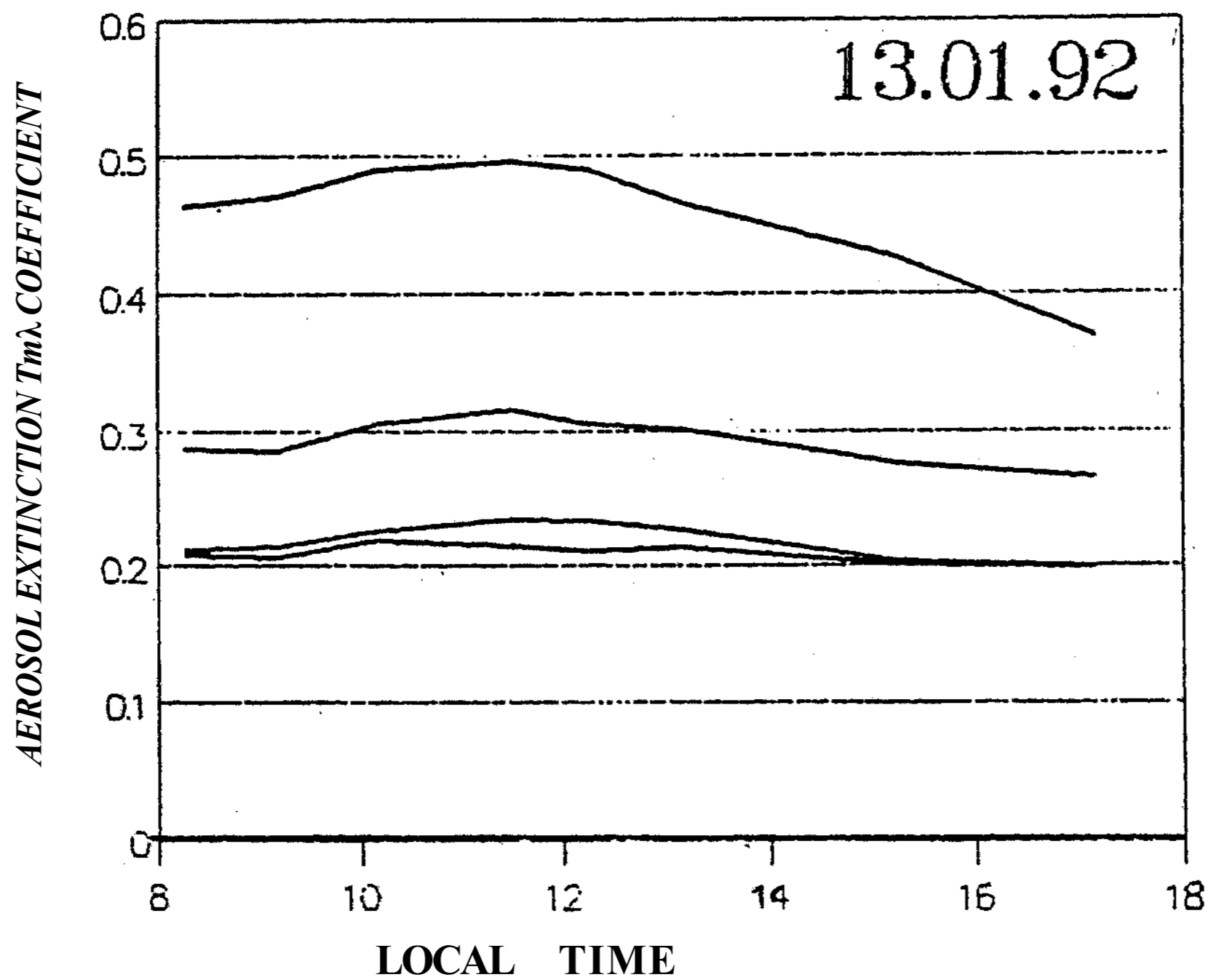


Fig 2 : Diurnal variation of $T_{m\lambda}$ at Maitri, Antarctica.

$B = 3.82$ (M.C.Sharma et al.,1986)

TR310= 1.0614

$h = 30^\circ$ (Required for calculation of m)

x = total ozone values

$T_{m\ 310}$ can be calculated using equation (6) and $T_{m\ 368}$ values.

The results are shown as follows (Table 1)

Diurnal variations and seasonal variations are shown in Figs. 3 & 4.

(c) Spectroradiometric measurements were recorded manually because of the failure of the computer part. The intensity of solar irradiance at a particular wavelength is indicated in terms of voltage at the screen. Then, as per instrument literature :

$$I = K(10^{-(v+5)} - 10^{-11})$$

where

V = output at the screen of electronic unit

$K\lambda$ = calibration constants as follow:

Table I

Date	Time (hrs)	Tm368	Tm310	I310	X	
					(x 10 ⁻⁸)	(atm-cm)
30.1.92	1451	0.3811	1.634	0.6398	13.1	0.3582
31.1.92	1500	0.3781	1.679	0.5043	15.8	0.3692
01.2.92	0805	0.4505	2.020	0.6370	223	0.2890
03.2.92	1400	0.4679	2.280	0.6918	16.0	0.3187
05.2.92	1415	0.4544	1.995	0.6398	638	0.4523
07.2.92	1415	0.4490	2092	0.6428	11.1	0.3792
08.2.92	1400	0.3942	1.922	0.5481	143	0.3709
12.2.92	0905	0.4343	1.553	0.5668	15.3	0.3571

λ nm	K λ
280	2.450 X10 ³
285	2.396 X 10 ³
290	2.354 x 10 ³
295	2.365 x 10 ³
300	2.308 x10 ³
305	2.470 x10 ³
310	2.712 x10 ³
315	2.951 x10 ³
320	3.999 x 10 ³
368	3.305 x10 ³

Calculations for all the $I\lambda$ values have been made. Data for $\lambda = 310$ nm have been used for estimation of O₃ as in the previous case. Results are shown in Table II. Ozone values in February'92 are normal but January'92 values are quite low (to be explained).

(d) Infrared sunphotometric observations were recorded on sunny days to study qualitative and quantitative presence of H₂O, CO₂, O₃, CO, CH₄, N₂O etc. in the atmosphere. A typical run is shown in Fig 5.

As indicated, the water vapour absorption bands and CO₂ absorption bands dominate the spectrum in 2.5 to 4.5 μ m range. In this range the absorptions of CO₂, H₂O, CH₄, N₂O overlap each other. 2.7 μ m is a moderately strong absorption band of H₂O. If we observe the spectra at New Delhi, it shows

Table II

Day Number	O3 (=310)
4	0.140(Atm-cm)
5	0.104
6	0.113
7	0.132
8	0.123
9	0.118
10	0.113
11	0.156
12	0.124
13	0.121
14	0.127
17	0.123
20	0.130
24	0.108
31	0.324
33	0.341
34	0.382
35	0.373
36	0.355
37	0.344
38	0.330
39	0.321
43	0.321

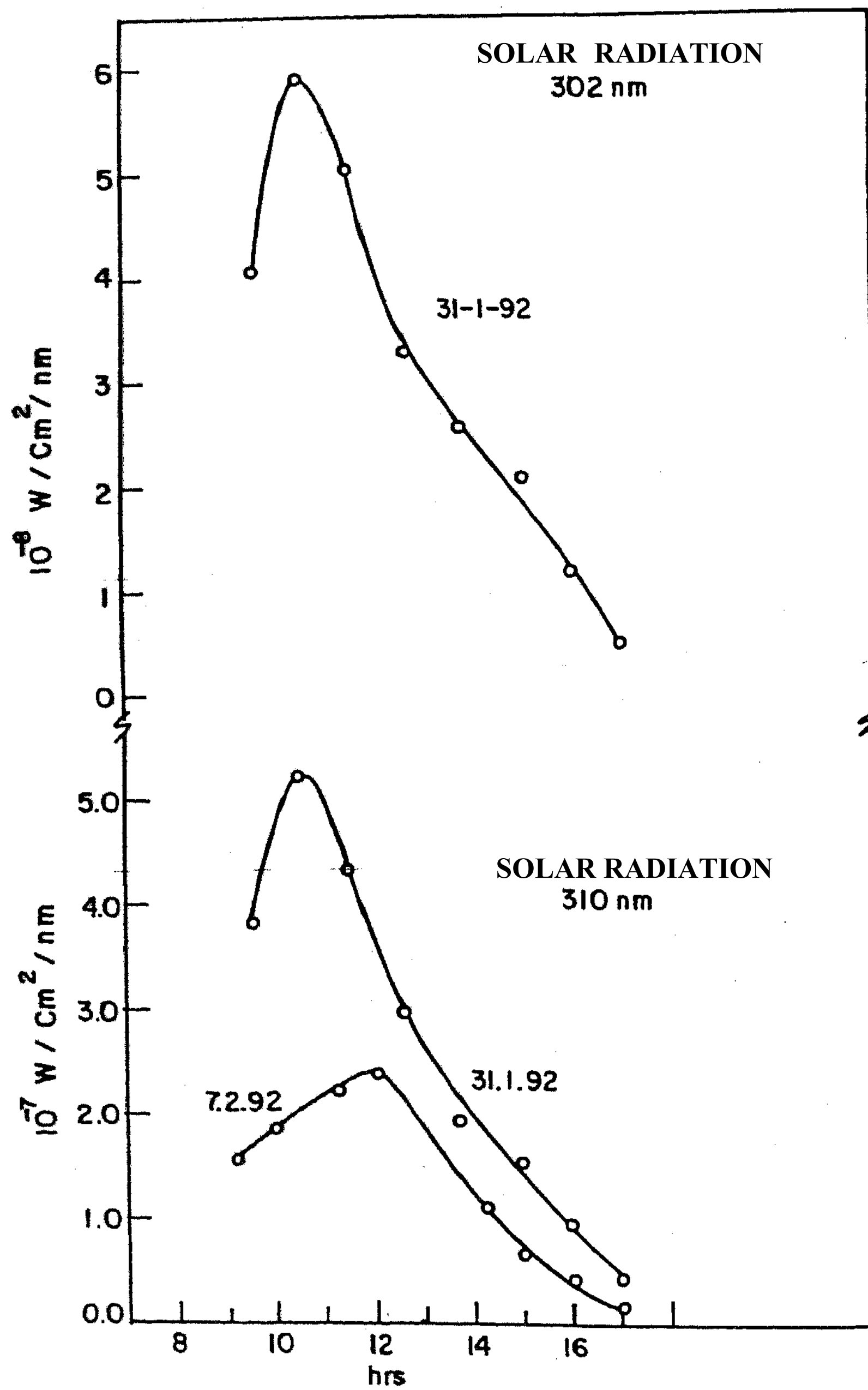


Fig 3 : Diurnal variation of solar irradiance at Maitri, Antarctica.

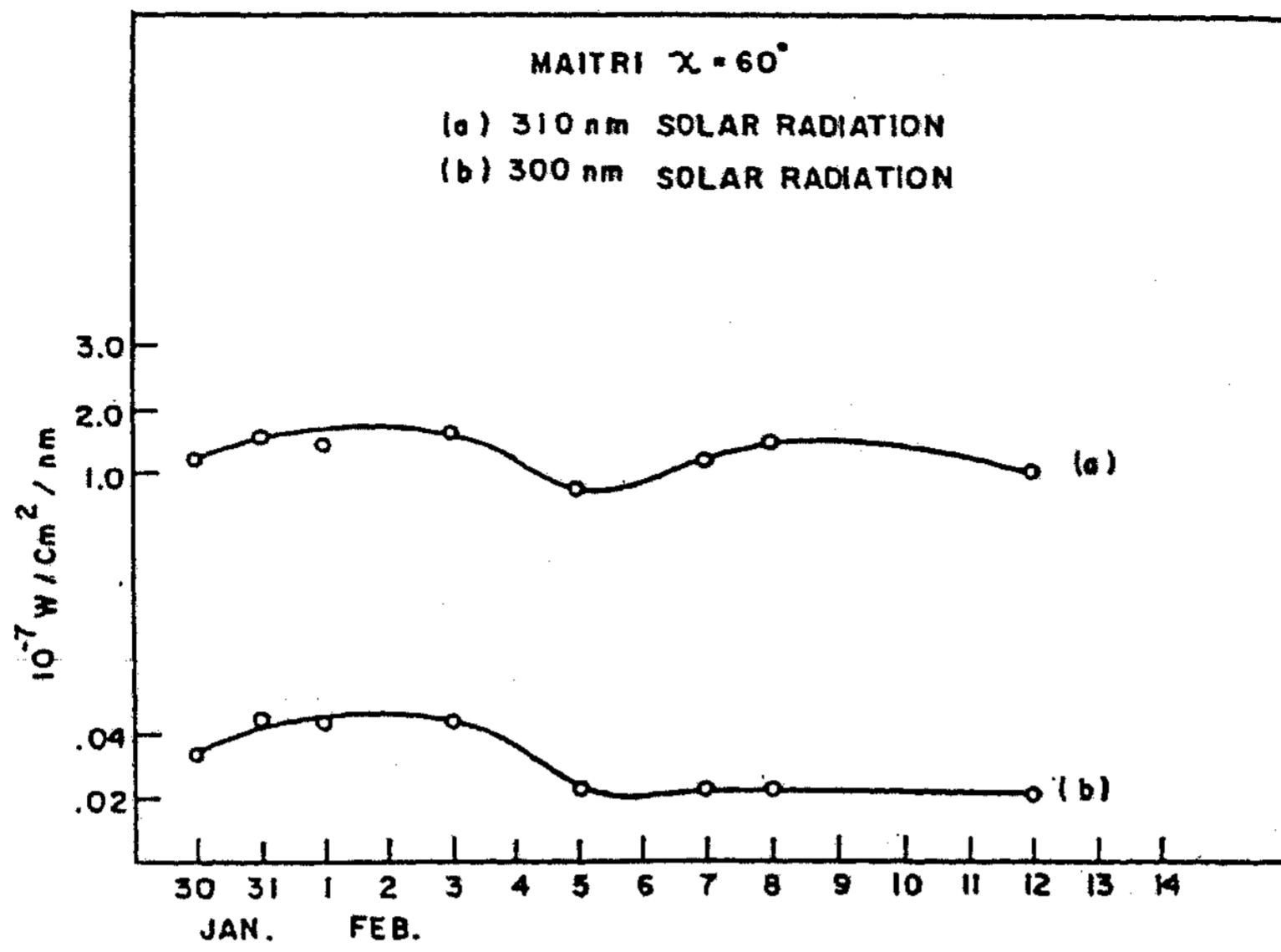


Fig 4 : Seasonal variation of solar irradiance at Maitri, Antarctica during austral summer of 1991-92.

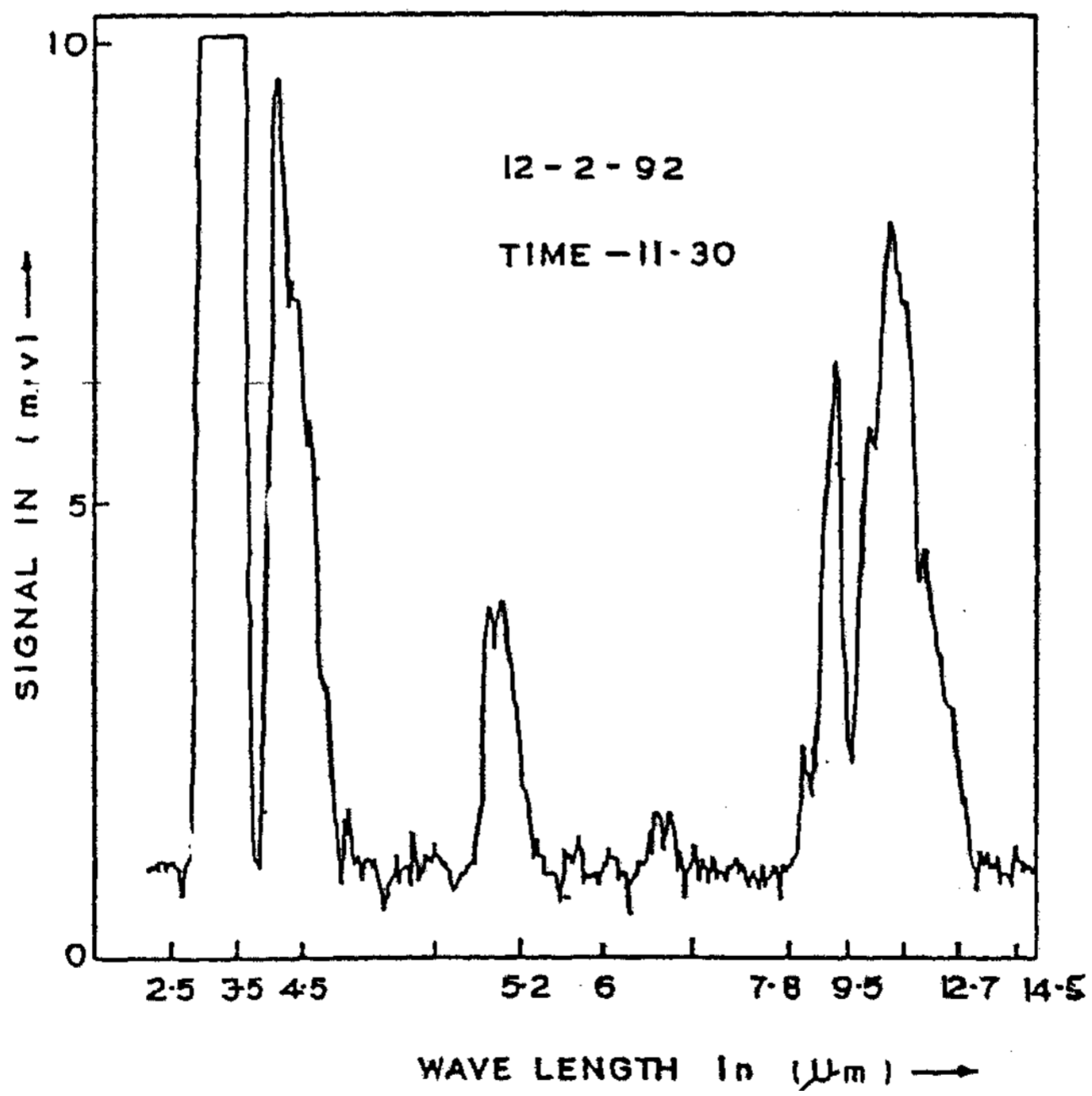


Fig 5: Infrared absorption (2.5 - 14.5 μm) spectra recorded at Antarctica.

signature of water vapour absorption but the absorption signature of $2.7\mu\text{m}$ is *absent in Antarctica* which shows low water vapour content of Antarctic atmosphere.

Also in Fig 5, $9.6\mu\text{m}$ ozone band is clearly seen. While comparing with New Delhi record, it shows more optical depth of ozone at Antarctica. This supports the early observations that at poles ozone concentration is more than at lower latitudes.

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- The Stratosphere(1981), theory and measurments Table B5.