UV STELLAR OCCULTATION MEASUREMENTS OF NIGHTTIME EQUATORIAL OZONE

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Abstract. The Princeton University Ultraviolet Spectrometer-Telescope on the NASA Orbiting Astronomical Observatory Copernicus was used for stellar occultation measurements of atmospheric ozone. Two sets of observations of the target star 8-Cen were carried out on 26 July 1975 and 13-14 June 1976 at wavelengths from 2550 Å to 3100 Å. After unfolding of the data, ozone density profiles near the equator within 3 hours of local midnight were obtained at altitudes from 47 to 114 km. A secondary maximum at 97 km has been observed in both sets of data. The ozone density between 47 and 75 km is a factor of 2 to 3 times as large as current models predict. At the lower boundary, about half the ozone destruction should be caused by NOx and ClOx. Above 55 km, virtually all loss is due to HOx. These results suggest an overestimate of HOx and ClOx loss processes or a serious underestimate of the Ox production rate. A minimum in O3 density near 87 km and a maximum at about 97 km are probably caused by reactions of O3 with atomic hydrogen and maximum nocturnal O3 production near 97 km.

Observation Technique

The stellar occultation technique for measurements of atomic species from satellites has been used and described by a number of researchers [Hays and Roble, 1973; Riegler et al., 1976; Atreya et al., 1976]. In the usual applications of the occultation method, the line-of-sight to the star crosses the limb of the earth during a time interval of approximately 10 to 25 seconds depending on the satellite orbit and the atmospheric species observed. In the observations to be discussed here, an observing geometry in which the line-of-sight grazes the earth's limb without actually being occulted by the disc is used. Typical observing times for this approach are 200-300 seconds.

The orbit of the NASA Orbiting Astronomical Observatory OAO-3 Copernicus is nearly circular with 745 km altitude and 35° inclination. These orbital parameters are very favorable for occultation measurements since they permit long occultation times during each orbit, and also reliable long-term predictions of suitable future occultation times.

The OAO-3 star tracker is programmed to track on a portion of the light entering the telescope; another portion is split off to the UV spectrometer. Since the star tracker uses visible light and the spectrometer senses UV light, refraction effects and possible guidance problems limit the range of occultation geometries to tangent altitudes above approximately 47 km.

The star tracker design also restricts occultation measurements to nighttime observations when the point of minimum tangent altitude (i.e., the "tangent point") is within three hours of local midnight. These restrictions and some engineering constraints limit applications of the limb-grazing method from OAO-3 to one or two observations per star per year.

The minimum tangent altitude during each orbit varied by approximately 10 km per orbit because of the precession of the ascending node of the OAO-3 orbit. This means that the observing wavelength for each orbit had to be selected to optimize absorption by atmospheric ozone, i.e., to obtain optical depths between 0.03 and 5. This range of optical depths was chosen so as to minimize the effects of uncertainties in the background and unattenuated counting rates. The set of observations, therefore, consists of a sequence of orbits with overlapping altitude ranges and usually different observing wavelengths between 2550 Å and 3100 Å. The actual data collection during each orbit extended over 240 seconds (limited by the on-board data storage capacity for the observations discussed here). The 1975 measurements were carried out within 25 minutes of local midnight at the tangent point, while the 1976 measurements extended over the range from 0100 to 0300 hours local time.
Figure 1. Line of sight ozone column density from the 1975 and 1976 stellar occultation measurements of E-Gen. Wavelengths and average latitudes and longitudes shown in the parenthesis are for the 1975 observations. Error bars of one standard deviation length are shown only when the error exceeds 10%.

Data Analysis

Examples of the data obtained in the 1975 ozone measurements have been presented by Riegler et al. [1976]. The calculation of column densities from the relative attenuation of the net stellar flux signal is readily obtained after compensation for the small effect of Rayleigh scattering. Since the bandpass of the OAO-3 Ultraviolet Spectrometer is narrow, typically 0.06-0.08 Å, integration over a finite wavelength band is not necessary.

For the conversion from optical depth of column density we have used the ozone absorption cross sections tabulated by Ackerman [1971]. According to a review presented by Hudson [1974], these absorption cross sections can be assumed to have an absolute accuracy of better than 5%. The star, E-Gen, was selected as a light source because previous study had shown its spectrum to be free of structure in the wavelength range of interest in our observations. Furthermore, the O₃ absorption cross section as measured with 0.1 Å resolution varies smoothly in the wavelength region surveyed.

Figure 1 shows ozone column densities as a function of tangent altitude for the July 1975 and June 1976 observations. The relative statistical uncertainty of the calculated column densities, which includes uncertainties in the unattenuated source response and in the instantaneous detector background rate is shown only when it exceeds 10%. As shown in Figure 1, the various sets of data have been obtained with different observing wavelengths and at different times. However, in all data sets the column densities decrease more or less exponentially up to about 85 km, but abruptly change in shape at that altitude, showing a secondary maximum near 97 km tangent altitude before decreasing again.

The relation between the line-of-sight column density and the local ozone density is given by the Abel integral which can be readily inverted analytically. In our computations we have used two approaches: quadratic or exponential local fits to the column density profile [cf., Roble and Hays, 1972], and a numerical iterative procedure [Maser, 1975]. The former method introduces local smoothing which depends on the number of data points used. The latter method can, with certain sets of input data, result in non-physical density profile features. At altitudes below approximately 85 km, a simple Chapman-factor conversion may also be used as a verification of the more elaborate Abel inversion technique.

Calculations with the above-mentioned inversion methods yield comparable results for the ozone density profiles which reflect the spread in the column density profiles given in Figure 1. Figure 2 shows the range of resulting ozone density profiles. The statistical accuracy of the data and the systematic accuracy of the numerical inversion method for the majority of data below 100 km is such that the density is reliable to within ±5% and the altitude to ±0.5 km. The range of ozone densities shown in Figure 2 reflects variations from orbit to orbit as functions of time and geographic location near the equator. The densities can then be compared with average theoretical predictions.

Model Calculations

The model O₃ density profile plotted in Figure 2 was obtained with a diurnal model assuming currently accepted rate constants [Hampson et al., 1977; Margitan, 1977], such as, for example

$$\text{OH} + \text{HO}_2 \rightarrow \text{H}_2\text{O} + \text{O}_2, \quad k = 2 \times 10^{-11} \text{cm}^3\text{s}^{-1} \quad (1)$$

$$\text{O} + \text{HO}_2 \rightarrow \text{OH} + \text{O}_2, \quad k = 3 \times 10^{-11} \text{cm}^3\text{s}^{-1} \quad (2)$$

The ClO volume mixing ratio was assumed to be 2.1 ppb and a tropical model atmosphere was adopted containing 20% more O₂ than the midlatitude spring and fall model [U.S. Standard Atmosphere Supplements, 1966]. In Figure 3, the Oₓ loss rates per day due to dominant processes between 40 and 80 km are plotted.

We divide our discussion into two parts, below 75 km and above it. Below about 75 km the experimental O₃ densities are higher than those in the model by a factor of 2 to 3. This is disturbing, since the dominant sink for Oₓ (O₃ + O + O(³P)) above 35 km is catalytic removal by OH, HO₂, and H while the ClO and NOₓ cycles should become almost equally effective as the HOₓ cycles at 47 km. Throughout the entire altitude regime below 75 km, photochemistry should dominate transport phenomena in determining Oₓ densities. This discrepancy between theory and observation extends into a region where ClO and NOₓ destruction of Oₓ becomes as important as HOₓ destruction. If the Oₓ data are accepted as valid, either the HOₓ and some combination of the ClO and NOₓ loss processes are all fortuitously overestimated or the Oₓ production rate is underestimated.

Above 75 km we have used the recently measured [Clyne, 1974] temperature dependence rate for the reaction
of $1.23 \times 10^{-10} \exp(-562/T)$. At $180^\circ K$ the rate of this reaction is a factor of 5 smaller than the value of $2.6 \times 10^{-11}$ [Hampson et al., 1973] used in an analysis of O$_3$ data previously published [Riegler et al., 1976]. Since reaction (3) is the dominant loss process for O$_3$ above 80 km, the result is an O$_3$ bulge above 90 km. The agreement between the model and observed profiles above 80 km is qualitatively good, but an altitude discrepancy of about 5 km exists and we do not understand.

In fact, the discrepancy between model and observed O$_3$ densities would effectively disappear at all altitudes if the observed profiles could be shifted downward about 5 km, or atmospheric density increased by about a factor of 2 in the model calculations. Given the geometry of the occultation exercises and the fact that a 5 km error in altitude of the tangent ray height would entail about a 30 sec error in timing and no more than a 2 sec error is expected, we have abandoned this explanation of the discrepancy after a painstaking study of the tracking data. The possible time uncertainty of 2 sec corresponds to an uncertainty in the instantaneous tangent altitude of less than 0.5 km for the lowest altitude data points of each orbit, and less than 1 km for the highest altitude data points. Moreover, a factor of 2 increase in the O$_3$ density in the homosphere appears unlikely. An equatorial radius of 6378.15 km was used in the data reduction.

The O$_3$ densities determined in our occultation studies agree well with those measured by Hilsenrath [1974] below 67 km near the equator in a rocket-borne chemiluminescent experiment. They disagree seriously with the data of Hays and Roble [1973] obtained from stellar occultation studies using OAO-2. The geometry of this observation did not permit the long integration time available using the limb grazing technique. The structure in the high-altitude O$_3$ profile has been deduced previously by Evans and Llewellyn [1970] who analyzed O$_2(^{1}Delta)$ and O$_2(^{1}Sigma)$ airglow data obtained in sounding rocket experiments in terms of O$_3$ photolysis. Noxon [1975] deduced similar nighttime ozone profiles above 80 km by observing the twilight enhancement in O$_2(^{1}Sigma)$ airglow emission. Quantitatively the Noxon O$_3$ values, which are somewhat indirectly inferred and are obtained at high latitudes are lower than those reported here. The equatorial observations of O$_2(^{1}Delta)$ emission by Han et al. [1973] present a more difficult problem of reconciliation since their 1.27 $\mu$m emission peak occurred at 80 $\pm$ 5 km. Mass spectrometer measurements of ozone between 90 and 110 km by Brinks [1975] have large experimental uncertainties and show no resemblance to the results presented in this paper.

We conclude that there is a serious problem reconciling photochemical models and observed nocturnal O$_3$ density profiles in the mesosphere. (There is no reason to expect other models to differ significantly from the one we deduce here). We note that a threefold increase in O$_3$ (or O$_2$) densities above 55 km due to a decrease in the HO$_x$ catalyzed O$_2$ recombination rates would require a factor of three decrease in HO$_x$ densities. This would call for a factor of about 30 decrease in H$_2$O and H$_2$ in the mesosphere because HO$_x$ density is proportional to the square root of products of H$_2$O + H$_2$ and O($^3$D). If the high O$_3$ densities reported here are characteristic of the tropics alone a conceivable but extremely unlikely explanation for such low tropical mesospheric HO$_x$ densities would involve a very large flux of sources such as CH$_4$, H$_2$O, H$_2$ from the tropical stratosphere. In analyzing the behavior of O$_3$ above the mesopause it may be useful to keep in mind the evidence obtained by OGO-VI airglow observations that the eddy diffusion coefficient near 100 km at tropical latitudes may be considerably enhanced compared to its value at higher latitudes [Donahue and Carlign, 1976]. Obviously the discrepancy uncovered here would disappear if the ozone absorption cross section were larger by a factor of 2 to 3 than the presently accepted value.

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Figure 3. 24 hours integrated loss rates from diurnal model for odd oxygen.
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