

Light of Very Short Wave-Length (2100 Å.) in the Solar Spectrum

FABRY and Buisson¹ have shown that the short wave-length limit of the spectrum of the sun near 2900 Å. is due to the absorption by ozone in the atmosphere of the earth. This absorption of the ultra-violet Hartley band of ozone begins at the wave-length 3100 Å. and reaches a maximum at 2540 Å. (Läuchli²). It is important to notice that the absorption falls rapidly on the short wave-length side of the band. Edgar Meyer³ first pointed out that this property of ozone gives an opportunity of observing sunlight in the region of 2100 Å. Several investigators searched for this short wave-length radiation, but without any success⁴. The reason for this is mainly given by the fact that below 2600 Å., the atmospheric oxygen is strongly absorbing over long distances⁵, so that it is necessary to use extremely sensitive apparatus to detect any radiation of this wave-length.

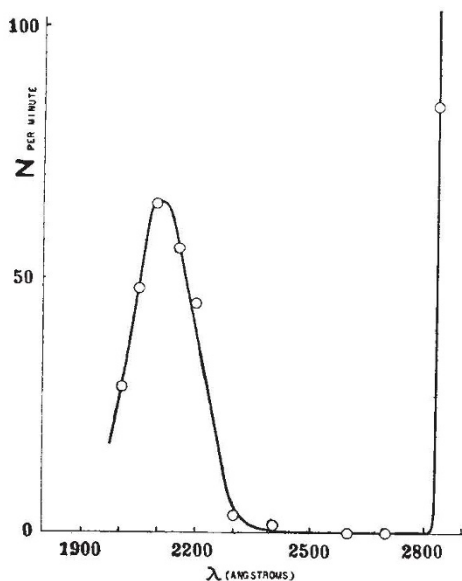


FIG. 1.

The researches of Schein and Stoll⁶ show that it is possible to build extremely sensitive light counters for the ultra-violet, which enable very weak radiations to be measured. Such counters have been used to search for the above-mentioned radiation of the sun. On account of the absorption by oxygen in this spectral region the main experiments were carried out at great heights; at first in Arosa (1860 m.), afterwards at the Jungfrauoch (3460 m.).

A quartz monochromator was directed to the sun, and the radiation measured by a sensitive photon counter. The number of photons, which are proportional to the intensity of the radiation, were registered. A full description of the experiments will be given later in a more detailed paper.

The curve shown (Fig. 1) was obtained at the Jungfrauoch and shows the ultra-violet spectrum of the sun from 2000 Å. to 2850 Å. For these measurements a photon counter was used with a sensitive layer of platinum sputtered in an atmosphere of hydrogen. In the curve the number of registered photons is plotted against the wave-length in angstroms.

The curve obtained shows distinctly that in the

short wave-length region of the ultra-violet a certain amount of sunlight reaches the earth's surface. The maximum intensity of this radiation lies at a point where the combined absorption of ozone and oxygen gives the greatest relative transparency of the atmosphere.

The following facts are characteristic of the intensity of this new radiation:

(1) It falls rapidly as the sun moves from the meridian to lower altitudes.

(2) It increases very much with the height above sea-level; at the Jungfrauoch we measured an intensity about a thousand times greater than at Arosa.

The more detailed discussion of these problems will shortly be published in a paper by M. Schein and B. Stoll in the *Helvetica Physica Acta*.

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¹ Ch. Fabry and H. Buisson, *Astrophys. J.*, **54**, 297; 1921.

² A. Läuchli, *Helv. Phys. Acta*, **1**, 208; 1928. *Z. Phys.*, **53**, 92; 1929.

³ Edgar Meyer, *Ann. Phys.*, **12**, 859; 1903.

⁴ Edgar Meyer, *Ann. Phys.*, **12**, 859; 1903. Edgar Meyer, *Verh. d. Klimat. Tagung in Davos*, 1925. P. Lambert, G. Déjardin and D. Chalonge, *J. Phys. et le Rad.*, **4**, 536; 1923.

⁵ H. Buisson, G. Jausseran and P. Rouard, *C.R.*, **190**, 808; 1930. **194**, 1477; 1932. F. W. P. Götz and H. Maier-Leibnitz, *Z. Geophys.*, **9**, 253; 1933.

⁶ M. Schein and B. Stoll, *Helv. Phys. Acta*, **7**, 485; 1934.

Steric Hindrance and Geometrical Isomerism

It is not often that evidence in the form of steric hindrance is available for confirming the configurations of geometrical isomers among ethylenic compounds. The following observations are thus of interest. *cis*- $\alpha\beta$ -Dibromocinnamic acid¹ (m.p. 100°) esterifies to the extent of only 6 per cent in one hour, and 33 per cent in 16 hours, by the Fischer-Speier method, whereas the *trans*-acid¹ (m.p. 136°) is esterified completely in one hour. This is in conformity with my earlier observation² in regard to the different modes of addition of bromine to β -phenylpropionic acid and its methyl ester.

cis-*o*-Nitro- α -bromocinnamic acid (m.p. 159°) has recently been prepared for the first time, in these laboratories, by Mr. J. D. Vasavada and me, as dense yellow rhombic tablets, by crystallising from benzene the unesterified portion of the acid product of the direct nitration of *cis*- α -bromocinnamic acid³ (m.p. 120°). It has also been made from *o*-nitrocinnamic acid dibromide by the action of alcoholic potash and pyridine. In ten per cent chloroform solution with a trace of free bromine, it is transformed with remarkable ease, by a few minutes exposure to direct sunlight, into the insoluble needle-shaped crystals of the *trans*-isomer⁴ (m.p. 212°), which is capable of complete esterification in 1 hour. This and further work in this series will shortly be published elsewhere.

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¹ *Annalen*, **247**, 139.

² *Ind. Sci. Congr. Chem. Abstr.*, No. 75, 1934.

³ *J. Chem. Soc.*, **83**, 668.

⁴ *Ber.*, **24**, 251.