## THE TEMPERATURE OF THE CYANOGEN-OXYGEN FLAME

Sir:

We have recently determined<sup>1</sup> the temperature of the H<sub>2</sub>-F<sub>2</sub> flame to be  $4300 \pm 150^{\circ}$ K. at 1 atm. pressure, using a modified line-reversal technique, with the sun in the heavens as the comparison radiator.

A much higher flame temperature is to be expected from thermodynamic considerations for the cyanogen-oxygen flame, according to the reaction

$$(CN)_2(g) + O_2(g) \longrightarrow 2CO(g) + N_2(g)$$

This reaction is exothermic to the extent of 126,680 cal. at 298°K. Furthermore, the thermal stability of the CO and N<sub>2</sub> molecules is even greater than that of the HF molecule.

Although the  $(CN)_2$ - $O_2$  flame was studied many years ago,<sup>2,3</sup> its flame temperature has never been measured directly. This year Thomas, Gaydon and Brewer<sup>4</sup> measured the vibrational "temperature" of the flame molecules and obtained a "temperature" of  $4800 \pm 200^{\circ}$ K. at the stoichiometric composition. The calculated flame temperature reported by these authors was  $4850 \pm 30^{\circ}$ K., using the high value, *i.e.*,  $\Delta H_0^0 = 226,000 \text{ cal./mole}$ or 9.76 ev., for the dissociation energy of  $N_2$ . They pointed out that by using the lower dissociation energy, *i.e.*,  $\Delta H_0^0 = 170,240$  cal./mole or 7.38 ev., the calculated flame temperature was 4325°K., thus fixing the dissociation energy of  $N_2$  at 9.76 ev. However, the vibrational "temperature" is not considered sufficiently representative of the true thermodynamic temperature of the flame. This conclusion was reached in connection with our work on the  $H_2$ - $F_2$  flame. The vibrational "temperature" of this flame was measured in our laboratory by Drs. W. S. Benedict, S. Silverman and B. W. Bullock<sup>5</sup> of the National Bureau of Standards and it was found that the precision and reliability of direct flame measurements by the linereversal method was greater than that of vibrational "temperature" measurements.

Our modified line-reversal method<sup>1</sup> was again employed. The metered  $(CN)_2^6$  and  $O_2$  streams were premixed in a 6-mm. glass tee filled with 20mesh quartz chips. A short length of quartz tubing acted as a torch tip. This tip was also filled with quartz chips and attached to the free end of the premixing tee by means of a piece of rubber tubing. The small oval-shaped opening in the face of the quartz torch tip measured 0.134"  $\times$  0.031". Small needle valves in the inlet gas lines made it possible to vary the rate of either gas.

The shape and structure of the flame varied with feed gas composition. Near the composition corresponding to one mole of cyanogen per mole of

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(3) A. Reis, Z. physik. Chem., 88, 513 (1914).

(4) N. Thomas, A. G. Gaydon and L. Brewer, J. Chem. Phys., 20, 369 (1952).

(5) See W. S. Benedict, B. W. Bullock, S. Silverman and A. V. Grosse, Phys. Rev., 87, 213-214 (1952).

(6) (CN): was prepared by thermal decomposition of Ag(CN), coupled with standard purification.

oxygen the flame consisted of two zones; the inner zone was a brilliant white comparable to that of a carbon arc. The outer zone had the blue color characteristic of the carbon monoxide flame in oxygen. In this work the outer zone was about 1" long and the inner zone 1/4'' to 1/8'' in length. These dimensions correspond to a total gas rate of about 200 cc. per minute. In these instances the flame burned right at the tip of the torch.

The theoretical flame temperature as a function of inlet gas composition was calculated using the latest thermodynamic data<sup>7,8,9,10,11</sup> ( $\Delta H_0^0$  for (CN)<sub>2</sub>dissociation = 114 kcal./mole).<sup>10</sup> These results together with the experimentally determined flame temperatures for various gas compositions are given in Table I.

TABLE I			
Vol. %, O2, (CN)3	Experimental temp., °K.	Theoretical temp., °K.	
<b>25</b> .3,74.7	2580, 2640	۰.	
26.8,73.2	3090, 31 <b>2</b> 0		
33.3,66.7	3270	3200	
36.5,63.5	3333		
<b>50</b> .0 <b>, 5</b> 0.0	<b>46</b> 40	4810	
<b>62</b> .0,38.0	4453		
66.7.33.3	4003, 4043, 4073, 4143	3700	

The maximum experimental flame temperature was observed at the stoichiometric point (50 vol. % (CN)<sub>2</sub>) in line with the above equation and equals  $4640^{\circ} \pm 150^{\circ}$ K. Thus at one atmosphere total pressure it is 340°K. higher than the hydrogenfluorine flame and is therefore the highest directly measured flame temperature recorded to date.

The theoretical flame temperature for the stoichiometric composition is calculated to be 4810  $\pm$  50°K. using the high value for the dissociation energy of nitrogen. A temperature of 4375°K. was obtained using the low value. The experimental flame temperature is in good agreement with the higher theoretical flame temperature and as such represents the first direct measurement of the cyanogen-oxygen flame temperature.

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(8) H. M. Spencer and G. N. Flannagan, THIS JOURNAL, 64, 2511 (1942).

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(10) L. Brewer, L. K. Templeton and F. A. Jenkins, THIS JOURNAL, 73, 1462 (1951).

(11) National Bureau of Standards, Selected Values of Thermodynamic Properties, API Project #44.

THE RESEARCH INSTITUTE OF	J. B. CONWAY
Temple University	R. H. WILSON, JR.
PHILADELPHIA, PA.	A. V. GROSSE
PECETVED OCTOBER 20	1059

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## AMICETIN, A NEW STREPTOMYCES ANTIBIOTIC Sir:

A Streptomyces sp., isolated from a soil sample collected near Kalamazoo, Michigan, was found to inhibit a number of bacteria when grown on an agar streak plate. Mycobacterium tuberculosis var. hominis (ATCC 607), Proteus vulgaris, Escherichia coli, and Staphylococcus aureus were markedly inhibited when cross-streaked against this new isolate. When the organism was grown in submerged