

the pipette was adjusted to the desired pressure and sealed by closing the needle valve B. The system was then opened to the gas burette lines. After this had been done the mercury and the gas enclosed in the pipette were allowed to expand to constant temperature in an oil bath. The volume of gas was then measured on a mercury manometer. From volume and number of moles of gas initially enclosed the density at the high pressure calculated. The correction was applied in these measurements to the temperature.

The apparatus contained liquid nitrogen and fitted with a connecting line to the pipette. Temperature was measured in a pocket immersed in the liquid nitrogen. A vacuum pump was also provided for evacuation to a vacuum.

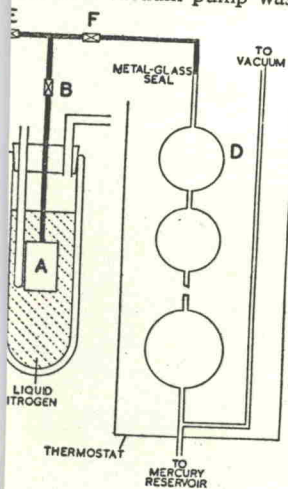


Diagram of apparatus.

so obtain temperatures below the normal

allowing the gas pipette be outside the cryostat. The proportion of the enclosed gas was approximately 10% of the total. Care was taken to ensure that the gas was dry; it was about 10% of the total. The number of moles of gas was determined from the compressibility data of Michels and Goudek 7 and of Johnson 8.

action of this volume which was immersed in the cryostat. These volume measurements of the compressibility of Michels and Goudek 7 and of Johnson 8.

erg standard test gauges of the Bourdon type were used for the purpose of the investigation. The gauges were calibrated by the method of Hainsworth at a pressure of 100-150 atm through a liquid nitrogen trap at 110° C and then through a steel trap at 110° C.

by a method similar to that used by Schiff and Steacie 10 was immersed in Dry Ice and alcohol. It was next opened and 50 ml of D₂O was added.

placed. The reactor was closed and returned to the Dry Ice bath for about 20 minutes to ensure "deep freezing" of the D₂O. After this period it was opened and 100 g of calcium was added. The vessel was again evacuated, then sealed and the Dry Ice bath removed. After a short while an extremely rapid exothermic reaction took place, the pressure rose to 100 atm in about half a minute. The deuterium was then passed at about 40

TABLE 1.—ISOTHERMS OF HYDROGEN

| T = 64.5° K | | T = 78.9° K | |
|----------------|----------------------------------|----------------|----------------------------------|
| pressure (atm) | density (mole cm ⁻³) | pressure (atm) | density (mole cm ⁻³) |
| 350 | 0.0344 | 300‡ | 0.0299‡ |
| 500 | 0.0387 | 500 | 0.0358 |
| 790 | 0.0440 | 600‡ | 0.0385‡ |
| 1000 | 0.0467 | 700 | 0.0404 |
| 1250 | 0.0493 | 800‡ | 0.0422‡ |
| | | 950 | 0.0442 |
| | | 1250 | 0.0478 |

‡ measurements using hydrogen prepared by reaction of H₂O with calcium.

TABLE 2.—ISOTHERMS OF DEUTERIUM

| T = 64.5° K | | T = 78.9° K | |
|----------------|----------------------------------|----------------|----------------------------------|
| pressure (atm) | density (mole cm ⁻³) | pressure (atm) | density (mole cm ⁻³) |
| 150 | 0.0265 | 150 | 0.0215 |
| 200 | 0.0296 | 200 | 0.0258 |
| 300 | 0.0348 | 300 | 0.0313 |
| 350 | 0.0369 | 350 | 0.0335 |
| 400 | 0.0381 | 400 | 0.0352 |
| 500 | 0.0408 | 500 | 0.0380 |
| 700 | 0.0446 | 700 | 0.0424 |
| 900 | 0.0474 | 900 | 0.0456 |

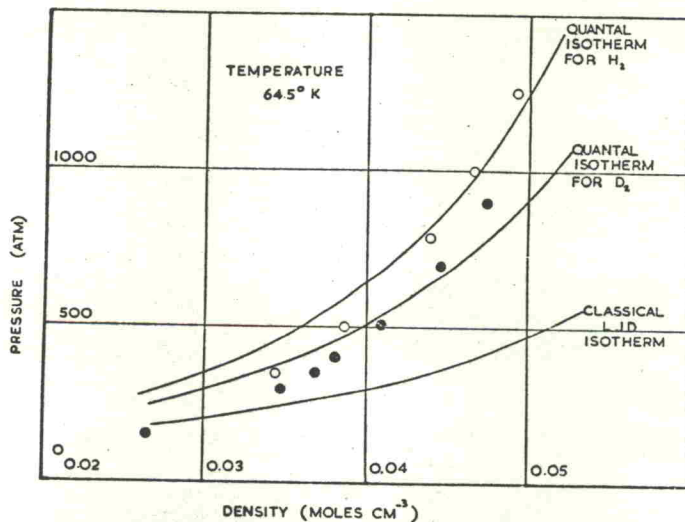


FIG. 2.—Theoretical and experimental isotherms at 64.5° K. The open circles are the experimental points for H₂, the filled circles are for D₂.

atm through a liquid nitrogen trap into the gas compressor. It is, perhaps, noteworthy that Schiff and Steacie 10 carried out the reaction at 260° C; in our case the reaction started while the reactor was still below 0° C. A few density measurements were made on a sample of hydrogen prepared by the same method, using H₂O instead of D₂O.