Measurement of Pressure Effect on Viscosity of Steam

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The viscosity of steam at subcritical pressures is measured by a capillary viscometer in the temperature range 250-600°C and at pressures up to 200 bar. In this region the gradient of isotherms in the viscosity-pressure diagram changes its sign from negative to positive. Results show that about 600°C is the lowest limit of the temperature range where the excess viscosity can be expressed as a function of density only. The measured viscosity agrees reasonably well with the correlation based on the authors' previous measurements at higher pressures.

A negative pressure effect on the viscosity of steam at subcritical temperatures was first discovered by Kestin and coworkers and recorded in several experimental reports (1). However, the highest temperature of these observations was limited to 450°C where the residual effect is still significant, and it is not clear at what temperature the effect dies out and becomes negligible.

The authors previously reported measurements of viscosity of water and steam up to 900°C and 1000 bar (3), but few measurements in the region of lower pressure were included in it. This paper describes some results at temperatures from 250° to 600°C and at pressures below 200 bar.

Method and Apparatus

Measurements were performed with a high-pressure capillary viscometer which is similar in principle to that used in the previous investigation (3) but different in detail. At subcritical pressures, liquid-gas interfaces in the flow system often cause trouble in flow rate measurement. The major differences compared with the previous apparatus were: addition of preheaters and coolers in order to fix the position of the interfaces in the flow circuit; and use of a shorter capillary for better temperature distribution.

Figure 1 shows a schematic outline of the experimental setup. The capillary is installed in the pressure vessel (4), which is heated by a main heater (6) through a thermostat cylinder (5) made of copper. The test fluid is circulated by a high-pressure injector system (13); it flows into the test section via the cooler (2) and subheater (3) and returns to the other end of the injector system. The flow rate is calculated from the record of an electric counter (14), while the pressure drop between both ends of the capillary is measured by a mercury manometer (12) and a traveling microscope. The capillary is made of Pt-Rh (5%) alloy and has the dimensions of 100 mm length and about 0.27 mm i.d.; it was calibrated with water and nitrogen (2). Because the inner surface of the metallic capillary was not perfectly smooth and also because the capillary was comparatively short, the critical Reynolds number was about 220, which was lower than the normal value of 2300 for a smooth circular tube. Variation of the capillary constant against the Reynolds number was checked with nitrogen. At low Reynolds numbers, the calibrated constant with respect to nitrogen agreed with those with respect to water (1.002 cP for water at 1 atm, 20°C).

Calculation

Viscosity values were calculated by use of the following modified Poiseuille equation (3):

$$\eta = \frac{\pi C}{8 \, lQ} \frac{P_1^2 - P_2^2}{2 \, P_1} (1 + 3 \, \alpha \Delta t) - \frac{\rho Q}{8 \, \pi l} \frac{(m + \ln P_1 / P_2)}{1 + \alpha \Delta t}$$
(1)

where η is the viscosity in μ P, C is a capillary constant in cm⁴, *l* is the length of the capillary in cm, *Q* is the volumetric flow rate in cm^3/s , P_1 and P_2 are pressures at the inlet and exit of the capillary in dyne/cm², α is the expansion coefficient of platinum (1.02 \times 10⁻⁵ C⁻¹), Δt is the temperature difference between experimental temperature and 20°C, ρ is the density of the fluid in g/cm³, and m is the kinetic energy correction factor, 1.12. The density of steam was calculated with the aid of the equation of state approved by ICPS (1967) (5).

Estimated Error

The largest error in the viscosity comes from the measurement of the pressure difference owing to the small value of $P_1 - P_2$ (keeping the Reynolds number as low as possible). The flow rate Q, calculated from the injector speed, was measured with a possible maximum error of 0.5%. The pressure was measured to 0.1 bar by a Bourdon-type pressure gage calibrated against a standard dead-weight gage, and the temperature was measured with the aid of a platinum resistance thermometer. The error in viscosity owing to the temperature measurement, including fluctuations and imperfect distribution, was estimated to be less than 0.5%. The length of the capillary was measured to 0.02%. The error owing to the uncertainty in the factor m was estimated to be less than 0.1%. The estimated error of viscosity is thus about 1.5%.

Results and Conclusion

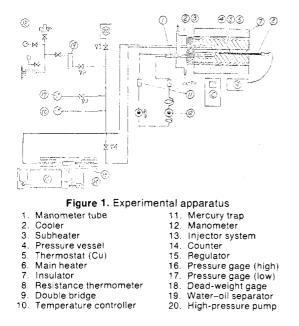
Experimental results are given in Table I. All data taken with a Reynolds number greater than 400 were disregarded. As seen in Figure 2, discrepancies between measured data and the ICPS equation (4) are quite large along the 375° and 400°C isotherms. The ICPS equation expresses the residual viscosity as a function of density only. The discrepancy of the ICPS equation and Equation 2 becomes smaller than experimental error above about 600°C. Measured data agree with the authors' equation in ref. 3:

$$\frac{2.39474 \times 10^5 \times (1 - \rho)^4}{1 + t + 1.75439 \times 10^{-8} t^4} + \frac{5.44832 \times 10^8}{(t + 90)^3 \times (1 - \rho)^2 + 4.64786 \times 10^{-2}}$$
(2)

In ref. 3 a constant in the equation was misprinted as -2.39474×10^4 , which should be corrected as above.

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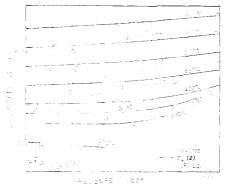


Figure 2. Measured results

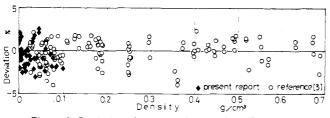


Figure 3. Deviation of measured points from Equation 2

The standard deviation between Equation 2 and the measured data is about 1.5%. The above expression describes the viscosity of water and steam up to $1000^{\circ}C$ and 1000 bar with the aid of a single equation and agrees with Rivkin's data (1). Because of the trouble caused by water-steam interfaces in the connecting tubes, the reproducibility obtained during the present investigation was less satisfactory than that in the authors' previous investigation at higher pressures (3). A deviation plot of the results, as well as of those of ref. 3 with respect to Equation 2, is shown in Figure 3.

At temperatures higher than 600° C, the excess viscosity of steam may be treated as a function of density only. Below about 600° C, the residual effect of the negative pressure gradient should be taken into account as given by Equation 2.

| t, °C | P, bar | ρ, g/cm³ | η, μΡ | t, °C | P, bar | ρ, g/cm³ | η, μΡ |
|----------|--------|-------------|----------|----------|---------------|-------------|----------|
| <u> </u> | | 10-6 | | | | 10-6 | |
| 250.7 | 4.9 | 2049 | 181.6 | 400.8 | 13.3 | 4367 | 240.6 |
| 250.2 | 10.2 | 4366 | 184.0 | 400.9 | 13.4 | 4400 | 238.4 |
| 251.1 | 15.8 | 6930 | 180.1 | 402.8 | 102.0 | 38497 | 239.6 |
| 251.0 | 37.0 | 18147 | 174.4 | 400.4 | 200.0 | 100210 | 255.0 |
| 250.8 | 37.0 | 18216 | 172.5 | 400.4 | 200.0 | 100210 | 200.0 |
| 250.9 | 37.9 | 18751 | 171.1 | 450.4 | 23.3 | 7162 | 263.1 |
| | 57.5 | 10/01 | 1,1.1 | 450.5 | 47.0 | 14782 | 259.9 |
| 301.2 | 16.0 | 6283 | 200.5 | 450.4 | 103.0 | 34737 | 262.5 |
| 301.2 | 16.2 | 6361 | 200.4 | 451.0 | 161.3 | 59161 | 267.6 |
| 302.9 | 25.7 | 10340 | 196.2 | 431.0 | 101.5 | 55101 | 207.0 |
| 301.2 | 50.4 | 22194 | 198.8 | 503.1 | 2 1 .1 | 5998 | 283.5 |
| 301.2 | 50.5 | 22242 | 195.6 | 500.8 | 25.9 | 7399 | 282.2 |
| | 50.5 | 22272 | 199.0 | 501.8 | 52.2 | 15241 | 280.0 |
| 350.7 | 29.3 | 10761 | 221.3 | 500.8 | 53.7 | 15714 | 286.1 |
| 350.4 | 50.1 | 19267 | 218.0 | 500.8 | 146.1 | 46580 | 291.1 |
| 350.3 | 83.4 | 35137 | 225.1 | 500.0 | 140.1 | 40000 | 232.2 |
| 350.5 | 87.3 | 37229 | 226.1 | 551.0 | 31.3 | 8384 | 306.2 |
| 349.7 | 115.7 | 55054 | 219.6 | 551.4 | 51.1 | 13857 | 300.8 |
| 349.9 | 115.7 | 55011 | 221.7 | 552.0 | 111.8 | 31566 | 313.0 |
| 350.8 | 150.0 | 86493 | 226.4 | 552.1 | 111.8 | 31561 | 314.9 |
| | 100.0 | 00150 | 22011 | 551.2 | 112.8 | 31906 | 317.4 |
| 377.3 | 6.7 | 2248 | 235.8 | 550.6 | 205.9 | 62410 | 320.6 |
| 377.5 | 7.0 | 2346 | 238.8 | | 20010 | | |
| 377.5 | 7,1 | 2380 | 237.4 | 601,4 | 16.7 | 4164 | 329.8 |
| 376.7 | 22.9 | 7899 | 237.4 | 601.3 | 56.8 | 14463 | 322.5 |
| 377.2 | 57.2 | 21038 | 234.2 | 601.3 | 58.4 | 14897 | 321.1 |
| 378.3 | 111.8 | 46524 | 242.8 | 601.3 | 98.1 | 25536 | 328.5 |
| 377.9 | 112.8 | 47149 | 241.5 | 601.4 | 99.7 | 25926 | 327.3 |
| 377.0 | 158.8 | 78182 | 240.5 | 601.3 | 135.3 | 35905 | 330.5 |
| | | | 2 | 601.3 | 157.9 | 42382 | 332.6 |
| | | | | 599.3 | 207.8 | 57533 | 340.1 |
| | | | | | 207.0 | | |

Table I. Measured Data

The authors express appreciation to J. Kestin for his kind advice on preparation of the manuscript.

Nomenclature

- $C = capillary constant, cm^4$
- / = length of the capillary, cm
- m = kinetic energy correction factor
- P_1 , P_2 = pressures at inlet and outlet of the capillary, dyne/cm²
- $Q = volumetric flow rate, cm^3/s$
- $t = \text{temperature}, ^{\circ}\text{C}$
- Δt = temperature difference, t = 20, °C
- α = thermal expansion coefficient of platinum, 1.02 × 10⁻⁵, °C⁻¹
- ρ = density, g/cm³
- $\eta = \text{viscosity}, \mu P$

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