

On Capillary Viscosity Measurements: How Far do Surface Tension Effects go?

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Groups V, IX and X

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**1. Background
and Purpose**



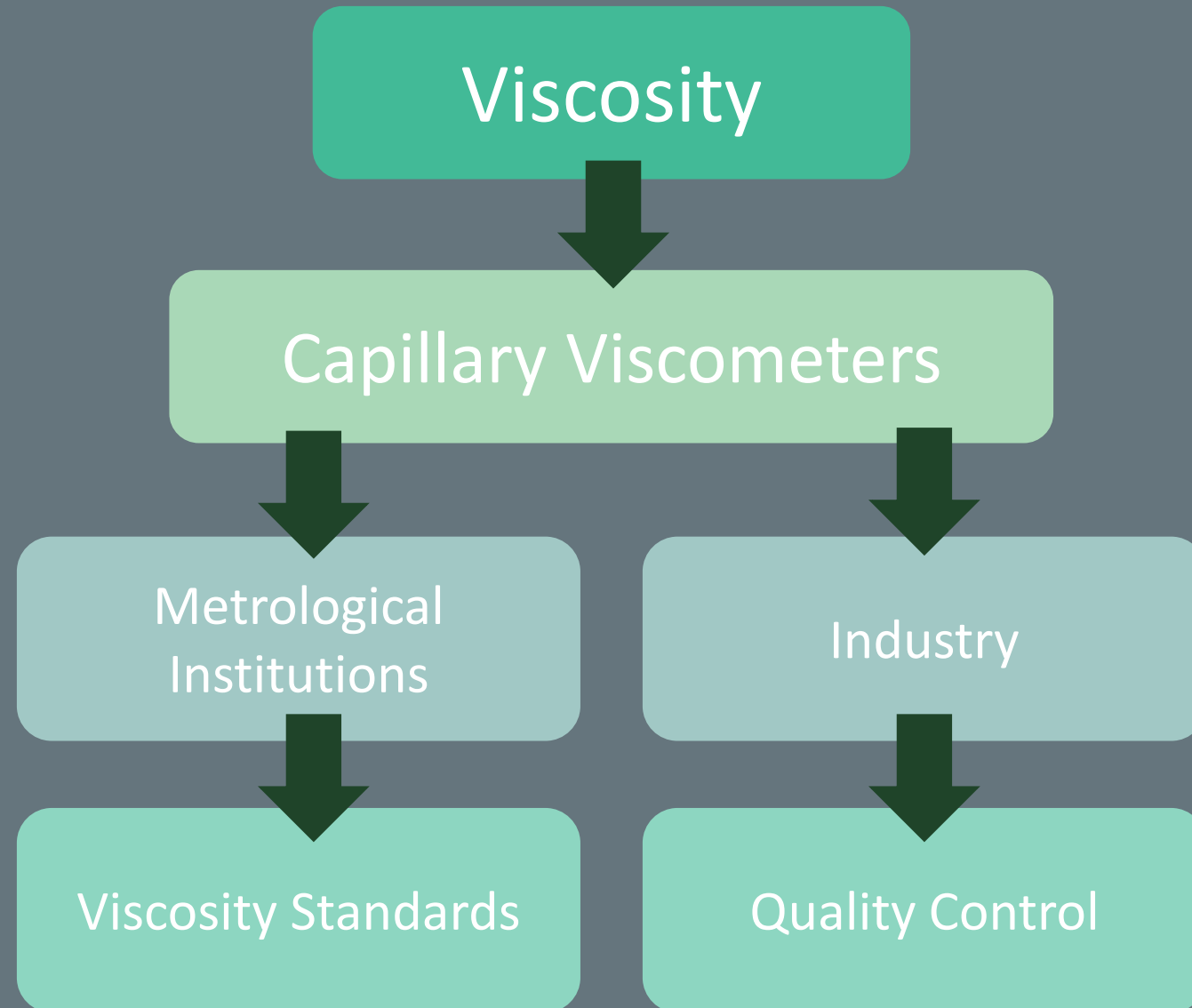
2. Experimental Work



3. Conclusions



1. Background and Purpose



Capillary Viscosity Measurements

Hagen – Poiseuille Equation:

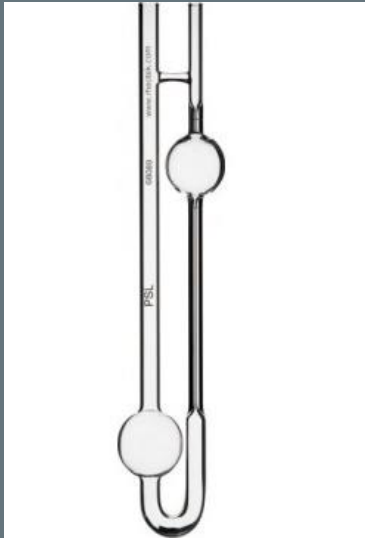
$$\Delta p = \frac{8\pi\mu LQ}{A^2}$$


$$v = \frac{\pi R^4 g h_m}{8VL} \cdot t$$

$$v = K \cdot t$$

- Characteristic for each capillary
- Kinetic Energy effects are not considered
- Archimedes' Principle is not considered

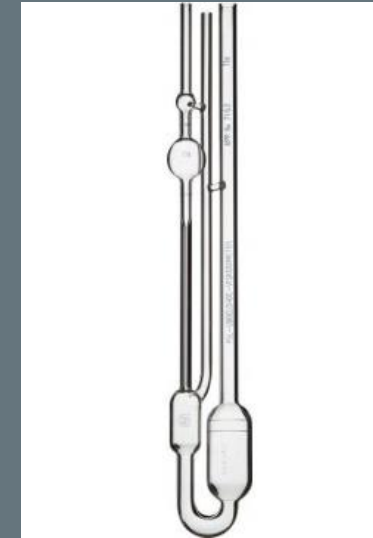
Types of Capillary Viscometers



Ostwald/U-tube
Viscometer



Canon-Fenske
Viscometer



Ubbelohde
Viscometer

Restrictions to the Hagen-Poiseuille Equation

Kinetic Energy Effects

The Hagen-Poiseuille equation is based on the following assumptions:

- Constant velocity along the tube of the capillary
- Parabolic distribution of the flow velocity through the cross section

Surface Tension Effects

- Ubbelohde Capillary Viscometers are designed to compensate the effect of the surface tension in the measuring bulb by means of the appropriate curvature of the suspended level
- Water as the primary viscosity standard

Purpose



Obtain the lowest uncertainty for the
Ubbelohde capillary viscometer 541
01/IA



Understand the need for the
corrections that must be considered
when using capillary viscometers and
how they should be applied



2. Experimental Work

Results

Capillary Viscometer Calibration

According to German
Standard DIN 51562 –
4 with 3 viscosity
standards



New value for K:
 $K = 0.006088 \text{ mm}^2/\text{s}^2$
($u_k = \pm 1.0\%$ 95%confidence level)



Old K (Schott Geräte):
 $K = 0.004998 \text{ mm}^2/\text{s}^2$
($u_k = \pm 1.2\%$ 95%confidence level)



Deviation of 21.8%



The glass conditions
change over time!

Results

Kinetic Energy Effects Assessment

For *n*-Tetradecane at 293.95 K

According to the
supplier Schott Geräte



Kinetic Energy Correction
vs flow time



Kinetic Energy Correction for
n-C₁₄ at 293.95 K



Correction of 0.03%

Results

Surface Tension Effects Assessment¹

For *n*-Tetradecane at 293.95 K

$$\mathbf{v}_{\text{corr}} = \mathbf{K} \cdot \mathbf{t} \cdot \left[\mathbf{1} - \chi \left(\frac{\sigma_0}{\rho_0} - \frac{\sigma_i}{\rho_i} \right) \right]$$



$$\chi = \frac{1 - \left(\frac{v_A}{K_0 t_A} \right)}{\left(\frac{\sigma_0}{\rho_0} \right) - \left(\frac{\sigma_A}{\rho_A} \right)}$$



Correction of 2.1%

$$\Delta_{\text{ST}} = \left[\chi \left(\frac{\sigma_i}{\rho_i} - \frac{\sigma_0}{\rho_0} \right) \right]$$

0, A – Viscosity Standards
i – fluid sample

Discussion

Table 1 – Comparison of the experimental results for *n*-tetradecane at 293.95 K.

$\eta_{\text{capillary}} / \text{mPa}\cdot\text{s}$ (before calibration)	$\eta_{\text{capillary}} / \text{mPa}\cdot\text{s}$ (after calibration)	$\eta_{\text{VW}} / \text{mPa}\cdot\text{s}$ (Vibrating wire) ²	$\eta_{\text{K}} / \text{mPa}\cdot\text{s}$ (Calibration + Kinetic Energy)	$\eta_{\text{ST}} / \text{mPa}\cdot\text{s}$ (Calibration + Surface Tension)
2.440 (8.44%)	2.262 (0.53%)	2.250	2.261 (0.48%)	2.311 (2.71%)

²Santos et al. 2017 (doi: 10.1016/j.fluid.2017.08.025)



3. Conclusions



The calibration showed that the current state of the capillary is very different from the initial one



The assesement of the kinetic energy and surface tension effects was performed



The viscosity value corrected for kinetic energy effects approximates the virating-wire viscosity value (not affected by any of the mentioned effects)



It is not accurate to calculate the surface tension effect for a fluid that is very similar to the one used to calibrate the viscometer



Understand the need for the corrections that must be considered when using capillary viscometers and how they should be applied

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