Joule-Thomson effect



P2320600

Physics	Thermodynamics	Temperature & Heat	
Difficulty level	QQ Group size	C Preparation time	Execution time
hard	-	10 minutes	30 minutes
This content can also be found online at:		第回 2017 記録 記録	

http://localhost:1337/c/6475b4cadd89390002768e25



Informations

Introduction

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Fig. 1: Experimental set-up (without PC)

A stream of gas is fed to a throttling point, where the gas (CO2 or N2) undergoes adiabatic expansion. the differences in temperature established between the two sides fo the throttle point are measured at various pressures and the Joule-Thomson coefficients of the gases in qestion are calculated. The principle of the Joule-Thomson effect is also applied, for example, in snow cannons for freezing water.





Other information (2/2)

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Safety instructions 1/2

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Carbon dioxide

H280: Contains gas under pressure; may explode if heated.

P403: Store in a well ventilated place.

Nitrogen

H280: Contains gas under pressure; may explode if heated.

P403: Store in a well ventilated place.

Fig. 2: safety instructions

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Setup and Procedure

Safety instructions 2/2

The set-up of the experiment is shown in Fig. 1.

If necessary, screw the reducing valves onto the steel cylinders and check the tightness of the main valves. Secure the steel cylinders in their location. Attach the PVC-tubing between the reducing valve and the Joule-Thomson apparatus with hose tube clips.On each side of the glass cylinder, introduce a temperature probe up to a few millimetres from the frit and attach with the union nut. Connect the temperature probe on the pressure side to inlet 1 and the temperature probe on the unpressurised side to inlet 2 of the temperature measurement apparatus.

Safety instructions 2/2

Important

The experimenting room and the experimental apparatus must be in a thermal equilibrium at the start of the measurement. The experimental apparatus should be kept out of direct sunlight and other sources of heating or cooling. Set the temperature measurement apparatus at **temperature difference** measurement. The temperature meter should be switched on at least 30 min before performing the experiment to avoid thermal drift.



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Safety instructions 2/2

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Important

Read the operating instructions for further explanations of the temperature meter. Open the valves in the following order: steel cylinder valve, operating valve, reducing valve, so that an initial pressure of 100 kPa is established. Reduce the pressure to zero in stages, in each case reading off the temperature difference one minute after the particular pressure has been established. Perform the measurement for both gases, and determine the atmospheric pressure and ambient temperature.



Ausrüstung

Position	Material	Item No.	Quantity
1	Joule-Thomson apparatus	04361-00	1
2	Universal power supply, 600mA 3/4.5/5/6/7.5/9/12V, incl. 9 adaptors	11078-99	1
3	Reducing valve for CO2 / He	33481-00	1
4	Reducing valve f.nitrogen	33483-00	1
5	Wrench for steel cylinders	40322-00	1
6	Gas-cylinder Trolley for 2 Cyl.	41790-20	1
7	Steel cylinder,CO2, 10l, full	41761-00	1
8	Steel cylinder,nitrogen,10l, full	41763-00	1
9	Hose clip f.12-20 diameter tube	40995-00	2
10	PVC tubing webbed, inner dia. = 8 mm, I = 1 m	47528-01	2
11	Cobra SMARTsense High Precision Temperature - Sensor for measuring temperature, -50 +150 °C (Bluetooth + USB)	12950-00	1
12	measureLAB, multi-user license	14580-61	1



Zusätzliche Ausrüstung

Position Material

Menge

1 PC with Windows XP® oder higher 1

Theory and evaluation

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Fig. 2: Throttling and the Joule-Thomson effect

The staate for real gases is given by the van der Waals equation

$$\left(p+rac{a}{V^2}
ight)*(V-b)=RT$$

where p ist the pressure, V the molar volume and T the temperature of the gas.

R ist the universal gas constant, a und b are the

caracteristic Van der Waalas coefficients of the gas. The additional

pressure by intermolecular forces of attraction is described by a, b

represents the volume of molecules



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Theory and evaluation In real gases the intrinsic energy U is composed of a thermocinetic and a potential component. The total change of the intrinsic energy U of a real gas therefore depends not only on the

temperature and the molar heat CV of a gas but also on the volume. The potential energy -a/V in the gas is given by the work against the intermolecular forces

 $U = C_v * T - \frac{a}{V}$

Theory of evaluation

The total didfferential of the intrinsic energy is given by

$$\mathrm{d} U \;=\; \left(rac{\partial U}{\partial T}
ight)_V \cdot \mathrm{d} T + \left(rac{\partial U}{\partial V}
ight)_T \cdot \mathrm{d} V$$

with

$$\left(rac{\partial U}{\partial T}
ight) \, = \, C_V \; \, ext{for} \; \, \Delta V \, = \, 0$$

and

$$\left(rac{\partial U}{\partial V}
ight) \, = \, + rac{a}{V^2} \; \, ext{for} \; \, \Delta T \, = \, 0 \; .$$

The expansion of the gas at the throttle-point is adiabatic (delta Q=0). If external heat losses and friction during the flow of the gas are excluded, the total energy H of the process is constant:

$$H \,=\, U + p \cdot V \,=\, U_1 + p_1 \cdot V_1 = U_2 + p_2 \cdot V_2$$

see Fig. 2

For further calculations, pressure p can be substituted by using van der Waals equation:

$$H \,=\, C_V \cdot T - rac{a}{V} + \left(rac{RT}{V-b} - rac{a}{V^2}
ight) \cdot V$$

Theory of evaluation

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After calculating the differential of (1) and substituting the total differentials by finite differences again, one obtains a relation between ΔT and ΔV :

$$\Delta H = C_V \cdot \Delta T + rac{2a}{V^2} \cdot \Delta V + rac{RV}{V-B} \cdot \Delta T - rac{RTV}{\left(V-b
ight)^2} \Delta V + rac{RT}{V-b} \cdot \Delta V$$

$$\Delta H \,=\, \left(C_V + rac{RV}{V-b}
ight) \cdot \Delta T + \left(rac{2a}{V^2} - rac{RTb}{\left(V-b
ight)^2}
ight) \cdot \Delta V$$

 $\Delta H = 0$ because H is constant.

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In the next step, b is considered to be small compared to V. The molar heat at constant volume CV can be replaced with the molar heat at constant pressure Cp via the ideal gas equation and under the assumption of even gas particle masses m:

 $C_{\mathrm{p,m}}~=~C_{\mathrm{V,m}}+R$

Putting (3) into equation (2) results in

$$rac{\Delta T}{\Delta V} = rac{R \, T b - 2 a}{V^2 \cdot C_{P,m}}$$

Theory of evaluation

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At the throttle point a pressure gradient p1–p2 and a temperature gradient T1–T2 are established. This effect is named the Joule-Thomson effect and is described by the coefficient

$$\mu \,=\, rac{T_1-T_2}{p_1-p_2}\,=\, rac{\Delta T}{\Delta p}$$

To derive a formula of this coefficient with Van der Waals forces involved, the difference ΔV in (4) has to be expressed in terms of Δp . This can be approximately done via the ideal gas equation again under the assumption of even gas particle masses m:

$$V = \frac{nRT}{p}$$



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After differentiating this equation to get an expression for dV, the total differentials are substituted by the finite differences, which results in the following:

$$\Delta V = rac{RT}{p^2} \Delta p$$

Solving equation (4) for ΔT gives

 $\Delta T \,=\, rac{\Delta V}{V^2 \cdot C_{{\scriptscriptstyle {\it v.m}}}} \cdot (R \, Tb - 2a)$

After replacing ΔV in (7) with the term in (6)

$$\Delta T = rac{-RT\Delta p}{p^2} \cdot rac{p^2}{R^2 \cdot T^2} \cdot rac{RTb-2a}{C_{p,m}}$$

Theory of evaluation

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Putting this into (4), the equation for the Joule-Thomson coefficient results in

$$\mu \,=\, rac{\Delta T}{\Delta p} \,=\, rac{1}{C_{p,m}} \,\cdot\, igg(rac{2a}{RT} - b igg)$$



Safety instructions 1/2



Fig. 3 shows the temperature differences measured at various pressures. The measurement values in Fig. 3 give the straight line gradients.

$$\mu_{
m CO_2}~=~(1.084~\pm~0.050)\cdot 10^{-5}~{
m K\over Pa}$$

and

$$\mu_{
m N_2}~=~(0.253~\pm~0.030)\cdot 10^{-5}~{
m rac{K}{Pa}}$$

where N2 is an approximation for air.

Theory of evaluation

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The two temperature probes may give different absolute values for the same temperature. But this is no problem, as only the temperature difference is important for the determination of the Joule-Thomson coefficients.

The literature values are

$$\mu_{\rm CO_2} \; = \; 1.16 \cdot 10^{-5} \; \tfrac{\rm K}{\rm Pa}$$

at 20 °C and 10^–5 Pa, and $\mu_{
m air}~=~0.23\cdot 10^{-5}~{
m K\over Pa}$

at 20 °C and 10^–5 Pa.

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For CO2, with

a= 3.60 Pa m^6 /mol^2

b= 42.7 cm^3/mol

cp= 366.1 J/mol K

equation (8) gives the coefficient

 $\mu_{\rm CO_2} = 0.795 \cdot 10^{-5} \, {{
m K}\over {
m Pa}}$

Theory of evaluation

For air, with

a= 40 Pa m^6 /mol^2

b= 39,1 cm^3/mol

cp= 9 J/mol K

equation (8) gives the coefficient

 $\mu_{
m air} \, = \, 0.387 \cdot 10^{-5} \; {
m K \over Pa}$

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Remarks

The formula for the Joule-Thomson coefficient indicates the condition for a cooling process

$$T_i = \frac{2a}{R \cdot b}$$

where Ti is the inversion temperature. If $\Delta p < 0$ the temperature T has to be lower then Ti to cool the gas with the process:

 $\Delta p < 0$ and T < Ti then $\Delta T < 0$.

For air (N2) or CO 2 cooling is observed at room temperature. H2 or He have a positive coefficient at room temperature, for H2 this can be hazardous because self ignition is possible.

<image><complex-block><form>



Measurement 2/2

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Fig.4: measureLAB software interface with the high precision sensor and the two temperature sensors

- $\circ~$ start the sensor and press it for at least 3 seconds.
- the sensor will be displayed on the left side of the measureLAB control panel. Please checkmark it.
- $\circ\;$ start the measurement using this button



 $\circ~$ end the measurement using this button.

