EXPERIMENTAL INVESTIGATION OF THE PHYSICAL PROPERTIES 
OF MEDIUM AND HEAVY OILS, THEIR VAPORIZATION 
AND USE IN EXPLOSION ENGINES 
By Fritz Heinlein 

PART I 

From "Der Motorwagen," October 10, 1925 

MEMORIAL A. L. MORTON 

Washington 
May, 1926
EXPERIMENTAL INVESTIGATION OF THE PHYSICAL PROPERTIES
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PART I.

Introduction

With the rapidly increasing use of motor cars and the extensive employment of explosion engines in agriculture, the consumption of light oils has increased enormously and the prices of these oils have continually risen. Efforts have accordingly not been lacking to increase the fuel-oil yield of the coal abundantly available in Germany and especially to transform the heavy oils obtained from the distillation of coal and lignite into light fuel oils. Parallel with these purely chemical processes, engine designers have sought by constructive means to remove the difficulties which now interfere with the use of heavy oils in explosion engines.

*From "Der Motorwagen," October 10, 1925, pp. 609-616.

This article is only a slightly condensed reprint of a graduation thesis at the Munich Technical High School. The experiments were performed in the physical laboratory of the M.T.H.S. (Dr. Oscar Knoblauch, Principal), and were rendered possible by the financial support of the Augsburg-Nuremberg Engine Works and of the German Petroleum Company.
While little has yet been learned from chemistry concerning profitable methods for obtaining an abundant supply of light oils from coal and heavy oils, engine designers have already made considerable progress toward the solution of the fuel problem. This progress has been made in two fundamentally different directions, which may be briefly distinguished as follows:

1. The group of engines with medium and high-pressure carburetion in the cylinder;

2. The group of engines with low-pressure carburetion of the heavy oils before reaching the cylinder.

Carburetion does not here mean the complete transformation of liquid fuels into the chemical condition of simple gaseous hydrocarbons, but into as thorough a mixture as possible of air and fuel, which, beginning with the atomization of the latter must, as a result of the partial evaporation of the atomized fuel, be so far advanced, up to its ignition, as to produce a sootless combustion of the mixture.

In the first case, the solution is sought by the employment of auxiliary pumps for delivering and vaporizing the fuel on its admission to the cylinder or by the use of hot-bulb engines or their modifications. For this purpose a new structural form of the engine is desirable: The recently perfected method of solid injection has afforded a new incentive to ren-
der heavy oils available for motor-car engines. A comprehensive article on this subject by F. E. Bielefeld ("Schwerölmotoren für Fahrzeuge unter besondere Berücksichtigung ihrer Wirtschaftlichkeit." "Autotechnik" 1924, No. 17), gives a good idea of the present status of this engine class and is further supplemented by the recent publication of two articles by Professor A. Loschge on the Acro engine ("Der Acro-Motor," "Autotechnik" 1924, No. 26, and 1925, No. 3) and by R. C. Von Gorrison, Hannover on a Dorner heavy-oil engine ("Der Schwerölmotorwagen von Dorner," "A.D.A.C. Sportzeitung," No. 1, Jan. 6, 1925).

According to both these publications, we have come, in this direction, very close to a solution of the heavy-oil problem for automobile engines, if we have not already reached a definite utilizable solution.

Along with the development of these high-pressure and medium-pressure heavy-oil engines, we have sought from the beginning to adapt the low-pressure carburetor engine, which works with suction intake and low compression up to about 5 atm. \((5 \text{ kg/cm}^2) = 71 \text{ lb./sq.in.}\), to heavy fuels without changing their internal structure. The entire work of producing an ignitible mixture of heavy oil and air is left to the carburetor in this engine class, the same as in light-oil engines. The carburetor alone has undergone many changes and seems to be capable of perfect adaptation to the complicated loading and operating conditions of automobile engines. Hence nearly
N.A.C.A. Technical Memorandum No. 362

All carburetor manufacturers have placed on the market carefully constructed heavy-oil carburetors (Dr. A. Heiller, "Schwerölbetrieb bei Kraftwagen," "Zeitschrift des Vereines deutscher Ingenieure" 1923, No. 10; W. A. Ostwald, "Schwerölbetrieb mit Doppelpatrin-Vergaser," "Autotechnik" 1923, No. 19) based on the carburetors with spraying nozzles almost universally used for light oils.

For a comprehensive solution of the problem of heavy-oil carburetion in low-pressure carburetor engines, there is still a lack of experiments on the physical properties of the heavy oils, which affect their vaporization, and a critical discussion of how far the vaporization can be carried and the atomization of the fuels must be carried, in order that the mixtures of air and heavy oils may attain the same degree of perfection as the light-oil mixtures and burn without freeing carbon.

We will therefore attempt in this article to define the technically important physical properties of the heavy oils, in so far as these properties affect the vaporization under the pressure and temperature conditions which exist during the intake phase of the engine. From the results of these experiments, conclusions will then be drawn (with mathematical treatment of the vaporization process of the oils) on the state of the vaporization, as it takes place under the conditions existing in the engine. By comparison with the light oils under like conditions, it is then obvious as to how far the atomiza-
tion of the heavy oils must be carried, in order to employ them successfully in carburetor engines.

In order to make the experiments more comprehensive, five heavy oils, two medium oils and two light oils were tested.

I. Heavy oils.

a) Gas oil (petroleum distillate),
b) Rositzer oil ( " " ),
c) Paraffin oil ( lignite " ),
d) Coal-tar oil I ( coal " ),
e) Coal-tar oil II ( " " ).

II. Medium oils.

a) Solar oil,

b) Kerosene

III. Light oils.

a) Gasoline,
b) Benzol (benzene).

Both the light oils (gasoline and benzol) were included in the investigation, because they have hitherto been the principal fuels used in carburetor engines and because conclusions can be drawn from the properties of these fuels concerning the conditions required for the vaporization of even heavy oils for their successful use as engine fuels.

General statements concerning liquid fuels.— The liquid fuels, from the volatile gasoline to the heavy oils, are to be regarded as mixtures of saturated and unsaturated hydrocarbons. According to their source they belong, for the most part,
either to the aliphatic series or to the aromatic series of hydrocarbons, the latter occurring chiefly in the fuel oils derived from the distillation of coal. They differ fundamentally from the chemically homogeneous substances, such as alcohol and pure benzol (benzene) \( C_6 H_6 \), and their behavior, while being converted from the liquid to the gaseous state, is characterized by the phenomena of physical mixtures.

1. The liquid fuels show no uniform boiling point at first, but boil within certain prescribed limits.

2. In evaporating the fuels at constant external pressure, the lighter components evaporate first and the temperature continually rises. Consequently, the vapor tension decreases as the evaporation proceeds.

This behavior of the fuel oils involves an accurately determinable vapor pressure only when the liquid component with the lowest boiling point is still present in the liquid form in the Torricellian vacuum at the temperature of the experiment. With a spatial increase of the vacuum, it happens, however, that the light liquid components completely evaporate and occur above the heavy components as unsaturated vapors and consequently no longer show the maximum saturation pressure of the fuels at the same temperature. This fact must be taken into account in the determination of the vapor pressures and cannot be disregarded in estimating the rate of evaporation.
Physical properties of liquid fuels.— The physical properties affecting the vaporization of liquid fuels are of prime importance in judging the possibility of their utilization in explosion engines. In addition to their chemical analysis, which is necessary for computing the quantity of air theoretically required, the following data must therefore be determined.

1. Specific gravities,
2. Boiling curves and limits,
3. Vapor pressures,
4. Specific volumes and vapor densities,
5. Vaporization heats,
6. Specific heats,
7. Vaporization rates and diffusion coefficients.

Chemical analysis.— The process does not differ from the usual method and was carried out by me for all the fuels with the electrical combustion furnace belonging to the inorganic chemical laboratory of the Munich Technical High School.

The combustion tube was filled to two-thirds of its length with copper oxide as the oxidizing agent. A boat-shaped platinum capsule, containing fine copper oxide in which a previously weighed quantity of the oil was absorbed, was then introduced into the empty end of the tube. Adjoining the platinum capsule, a copper spiral was also introduced. In order to extract any halogens which might be present, a silver spiral was intro-
duced into the opposite end of the tube. The tube was then placed in the heating jacket of the electric furnace. The platinum-boat end of the combustion tube was connected (with the interposition of a drying apparatus) with an air-filled gasometer. By turning the cock of a drying apparatus, connection was made with a second gasometer, which was filled with oxygen. All the connections had to be air-tight. A regulating cock was introduced into the circuit before the drying apparatus, in order to regulate the flow of the air or oxygen. The silver-spiral end of the combustion tube was connected with a calcium chloride tube and a potash apparatus, which were accurately weighed before the experiment. The weight of the carbon dioxide was found in the potash apparatus and the weight of the water in the calcium chloride tube and thence from the amounts of carbon and hydrogen contained in the fuel were calculated. It was necessary to inclose gasoline or benzol in a small bulb of thin glass, sealed by melting, and to place the bulb in a separate tube which was introduced into the circuit before the combustion tube. After the bursting of the bulb, the air stream was saturated with the fuel vapors and carried them into the combustion tube.

Three analyses were made with each fuel and the mean results are given in
TABLE I.

<table>
<thead>
<tr>
<th>Fuel</th>
<th>Sp. grav. at 15°C (59°F)</th>
<th>Chemical analysis</th>
<th>Boiling limits at 730 mm (28.35 in.) Hg</th>
<th>Remarks</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>% C</td>
<td>% H</td>
<td>from</td>
</tr>
<tr>
<td>Gas oil</td>
<td>0.864</td>
<td>86.92</td>
<td>12.94</td>
<td>190</td>
</tr>
<tr>
<td>Rositzer oil</td>
<td>0.907</td>
<td>84.30</td>
<td>11.05</td>
<td>140</td>
</tr>
<tr>
<td>Paraffine oil</td>
<td>0.906</td>
<td>86.94</td>
<td>12.91</td>
<td>200</td>
</tr>
<tr>
<td>Coal-tar oil I</td>
<td>0.910</td>
<td>87.50</td>
<td>12.01</td>
<td>90</td>
</tr>
<tr>
<td>Coal-tar oil II</td>
<td>0.852</td>
<td>86.98</td>
<td>12.62</td>
<td>120</td>
</tr>
<tr>
<td>Solar oil</td>
<td>0.850</td>
<td>86.30</td>
<td>13.40</td>
<td>155</td>
</tr>
<tr>
<td>Kerosene</td>
<td>0.813</td>
<td>85.88</td>
<td>14.10</td>
<td>150</td>
</tr>
<tr>
<td>Gasoline</td>
<td>0.739</td>
<td>85.70</td>
<td>14.30</td>
<td>60</td>
</tr>
<tr>
<td>Benzol</td>
<td>0.870</td>
<td>91.35</td>
<td>8.30</td>
<td>75</td>
</tr>
</tbody>
</table>

The specific gravities were determined with a specific-gravity bottle.

**Quantity of air theoretically required for the combustion.**

This can be determined from the chemical analysis of the fuel. The following computation accords with the combustion formulas.

\[
2\text{H}_2 + \text{O}_2 = 2\text{H}_2\text{O} \quad \text{C}_2 + 2\text{O}_2 = 2\text{CO}_2
\]

\[
4\text{kg} + 32\text{kg} = 36\text{kg} \quad 24\text{kg} + 64\text{kg} = 88\text{kg}
\]

From these formulas we obtain, for coal-tar oil I, the volumetric proportions at 15°C (59°F) and 760 mm (29.92 in.) Hg as follows:
\[
\begin{align*}
\text{The weight of the combustion water is } &\quad 0.875 \text{ kg of } H_2O = 0.806 \text{ kg of } H_2O \\
&\quad 0.1201 \text{ kg of } H_2 + 0.744 \text{ kg of } H_2O = 1.488 \text{ kg of } H_2O
\end{align*}
\]

\[
\begin{align*}
0.488 \text{ kg of } C_2H_2 + 2.4 \times 0.875 \times 760 \times 2.88 &= 0.02 \text{ m of } CO_2 \\
22.4 \times 0.875 \times 760 \times 2.88 &= 3.8 \text{ m of } C_2H_2
\end{align*}
\]

\[
\begin{align*}
0.1201 \text{ kg of } H_2 + 22.4 \times 0.1201 \times 760 \times 2.88 &= 0.02 \text{ m of } H_2O \\
22.4 \times 0.1201 \times 760 \times 2.88 &= 3.8 \text{ m of } H_2
\end{align*}
\]

Give the following values:

In determining the gas volumes, the mean barometer reading of 725 mm (28.54 in.) was taken, with which the experiment was performed.

The theoretical combustion equation is

\[
\text{I kg fuel} + 2.55 \text{ m of } O_2 = 1.486 \text{ m of } H_2O + 1.806 \text{ m of } CO_2
\]

Note: The above (etc.) not included in the above.

On addressing the weight of the combustion (cases)

0.9961 kg fuel + 2.550 m of O_2 = 1.856 m of NO + 1.806 m of CO_2

0.876 kg of O + 1.806 m of CO_2

0.1201 kg of H_2

N. A. C. A. Technological Memorandum No. 365
and the volumetric share of the CO$_2$ in the noncondensable portion of the combustion products is

$$\frac{1.806 \times 100}{1.806 + 9.59} = \frac{180.6}{11.396} = 15.87\%$$

In the cylinder combustion of heavy-oil fuel mixtures, according to previous experiments, a greater excess of air is necessary for complete combustion. Conclusions may be reached regarding the lack or excess of air in the cylinder from the above computation of the theoretical air requirement and from the determination of the volumetric content of CO$_2$ and of unburned CO and O in the exhaust gases of an engine.

Table II gives the theoretical air consumption, the weight of the combustion water and the percentage of CO$_2$ in the total volume of the combustion products for the theoretically required amount of air.
## TABLE II.

<table>
<thead>
<tr>
<th>Fuel</th>
<th>Chemical analysis</th>
<th>Theoretical consumption of air</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>% C</td>
<td>% H</td>
</tr>
<tr>
<td>Gas oil</td>
<td>86.92</td>
<td>12.94</td>
</tr>
<tr>
<td>Rositzer oil</td>
<td>84.3</td>
<td>11.05</td>
</tr>
<tr>
<td>Paraffine oil</td>
<td>86.94</td>
<td>12.91</td>
</tr>
<tr>
<td>Coal-tar oil I</td>
<td>87.5</td>
<td>12.01</td>
</tr>
<tr>
<td>Coal-tar oil II</td>
<td>86.98</td>
<td>12.62</td>
</tr>
<tr>
<td>Solar oil</td>
<td>86.3</td>
<td>13.40</td>
</tr>
<tr>
<td>Kerosene</td>
<td>85.88</td>
<td>14.10</td>
</tr>
<tr>
<td>Gasoline</td>
<td>85.70</td>
<td>14.30</td>
</tr>
<tr>
<td>Benzol</td>
<td>91.35</td>
<td>8.30</td>
</tr>
</tbody>
</table>
TABLE II (Cont.)

<table>
<thead>
<tr>
<th>Fuel</th>
<th>Combustion with theoretical quantity of air</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Wt. of combustion water</td>
</tr>
<tr>
<td>--------------</td>
<td>-------------------------</td>
</tr>
<tr>
<td>Gas oil</td>
<td>kg</td>
</tr>
<tr>
<td></td>
<td>1.287</td>
</tr>
<tr>
<td>Rositzer oil</td>
<td></td>
</tr>
<tr>
<td></td>
<td>1.099</td>
</tr>
<tr>
<td>Paraffine oil</td>
<td></td>
</tr>
<tr>
<td></td>
<td>1.283</td>
</tr>
<tr>
<td>Coal-tar oil I</td>
<td></td>
</tr>
<tr>
<td></td>
<td>1.196</td>
</tr>
<tr>
<td>Coal-tar oil II</td>
<td></td>
</tr>
<tr>
<td></td>
<td>1.256</td>
</tr>
<tr>
<td>Solar oil</td>
<td></td>
</tr>
<tr>
<td></td>
<td>1.331</td>
</tr>
<tr>
<td>Kerosene</td>
<td></td>
</tr>
<tr>
<td></td>
<td>1.403</td>
</tr>
<tr>
<td>Gasoline</td>
<td></td>
</tr>
<tr>
<td></td>
<td>1.421</td>
</tr>
<tr>
<td>Benzol</td>
<td></td>
</tr>
<tr>
<td></td>
<td>0.828</td>
</tr>
</tbody>
</table>

Evaporation curves.— These show the evaporated quantities of fuel as plotted against the boiling temperatures and are plotted by taking the evaporated volumes in % of the original volume as the ordinates and the corresponding temperatures of the oil vapor as the abscissas.

Hence 100 cm³ (6.1 cu.in.) of each fuel was evaporated in the apparatus made by Runge of Berlin for the Berlin "Physikalisch-Technischen Reichs-Anstalt" and the distillates were collected in graduated vessels. The evaporated volumetric percentages were obtained directly from the quantities of the distillates collected at the different temperatures.
The curves in Fig. 1, for all the fuels tested, show that the medium and heavy fuels do not evaporate at atmospheric pressure within so narrow temperature limits and so uniformly as gasoline and benzol. Furthermore, they leave, within the given boiling limits, residues which are decomposed only at the high combustion temperatures in the engine cylinder. The evaporation temperatures are moreover considerably higher than for the light oils.

Vapor tensions.—The vapor tension of a liquid at a given temperature depends, in chemically homogeneous substances, only on this temperature and rises and falls with it. The vapor-tension curve, which represents this dependence of the vapor pressures on the temperatures is, at the same time, the limiting curve between the wet and superheated vapor. It is definitely determined only when the liquid has a constant boiling temperature at the existing external pressure and does not change as the evaporation proceeds. From the vapor-tension curve there can be found directly the temperature at which the complete evaporation of a given quantity of liquid (to a predetermined volume) is possible. In this connection, however, the saturation pressure must be determined as a partial pressure in the given volume of vapor and air for the evaporation. The greater the air space is, in which the evaporation takes place, just so much smaller will be the partial pressure of the liquid and just so much lower the temperature at which complete evaporation is possible.
The vapor tensions cannot be so readily determined in liquid mixtures as in chemically homogeneous substances. The process of evaporation has already been briefly described under the heading "General statements concerning liquid fuels" and will be more fully discussed before we proceed to the determination of the vapor-pressure curves.

In the isothermal evaporation of the fuel in a Torrecellian vacuum, we will observe that the fuel continues to evaporate until equilibrium is established between the fuel liquid and the fuel vapor. The latter contains lighter hydrocarbons than the former, because the lighter components of the fuel evaporate at a lower temperature than the heavier. At a given temperature we will obtain a physically correct vapor pressure, which is composed of the pressures of all the liquid components, when the lightest liquid component is present in the vacuum in both the vapor and liquid form. As soon as the lighter liquid components are no longer present in the liquid form, they cease to appear at their full saturation pressures in the total pressure of the fuel. The vapor pressure is then smaller than the true saturation pressure of the fuel. This phenomenon was thoroughly investigated in a preliminary experiment with the apparatus for determining the vapor pressure which will be described later. An increase in the size of the vacuum, due to the sinking of the mercury and of the liquid fuel, resulted in a further evaporation of the fuel at constant temperature, but simultaneously also in lowering the
vapor pressure. It was only with a vacuum of less than 15 cm$^3$
(0.915 cu. in.) for the heavy and medium oils and 20 cm (1.22
cu. in.) for both the light oils, that a constant vapor pressure
at constant temperature was observed, up to temperatures of re-
spectively 60 and 30°C (140 and 86°F), for the 0.1 gram of fuel
used.

In order to be able to establish, however, the vapor ten-
sions in a practical manner for the future consideration of the
evaporation process, I refrained from determining the absolute
values of the vapor tensions at the small volumes of 15–20 cm
(0.915-1.22 cu. in.) which never occur in the engine cylinders.

These values would have been much too high for the actual evapora-
tion process at the given volume in the engine. In all the vapor-
pressure experiments, I have therefore determined the vapor vol-
umes corresponding to the vapor pressures for the temporarily
constant temperature; established this dependence in pressure-
volume curves for every temperature; and plotted therefrom the
vapor-pressure curves for predetermined volumes, whereby the ob-
erved vapor volumes were converted to 0.1 gram fuel. This made
it possible to calculate the vapor pressures under like condi-
tions for all the fuel oils.

The vapor-pressure curves for each fuel form a group corre-
sponding to the dependence of the vapor pressures on the volume.
The experimental results are given in Tables III - XI and Figs.
3a - 11b, (Part II, Technical Memorandum No. 363).
The vapor tensions for liquid fuels had often been previously determined without taking account, however, of the above-described phenomenon. On the vapor tensions of heavy oils we have Neumann's experiments (K. Neumann, "Untersuchung an der Dieselmachine," "Zeitschrift des Vereines deutscher Ingenieure" 1918, p. 708), which were performed, however, at constant volume and served as a basis for the discussion of evaporation in constant-pressure oil engines.

**Experimental procedure and apparatus.**—The experiments were performed according to the isothermal method, which is the best for liquid mixtures. Static and dynamic methods give accurate results only with chemically homogeneous substances. The apparatus used in the experiments (Fig. 2) was made of changeable length and can be used both to determine the vapor pressures and the vapor densities at constant pressure.

A seamless thick-walled steel tube d, of 6 mm (2.36 in.) inside diameter, with three welded steel-tube connections, is fastened with iron straps to an iron pan and a small table which supports the whole apparatus. Welded to each tubular connection there is a threaded piece for receiving a packing box, into which is inserted a glass tube well tightened with rubber gaskets: 1. experiment tube a, provided with a cubic-centimeter scale; 2. barometer tube b; 3. pressure-gauge c, which is needed for experiments above atmospheric pressure. A thin bent steel tube is welded to the tube d which joins the three glass tubes. The
bent tube is connected with a glass tube \( g \), about 700 mm (27.6 in.) long, the latter being connected with the rubber tube \( e \), and the glass mercury reservoir \( R \). The glass tube \( g \) carries a tightly-fitted cock \( h \). The mercury reservoir \( R \) is a cylindrical glass receptacle with a calcium chloride tube at the top and a thick nipple for connecting with the tube \( e \). The reservoir rests in a felt-lined oak sheath which can be raised or lowered on a sheet-steel guide. The arresting device is a steel spring, which is screwed to the oak sheath and engages in the notches in the steel guide. The latter is screwed to an oak support, which has a wood base and carries a wooden measuring stick with a millimeter scale. On the measuring stick there is a steel slide with a spring stop. This slide carries a glass tube \( f \), for reading the levels of the three tubes \( a \), \( b \), and \( c \).

Since the room was too small for reading the mercury meniscus with a telescope, the tube \( f \) had to be resorted to. This was secured to the slide by iron straps and branched toward the experiment tubes in two arms and terminated in front of the measuring stick in a vertical tube about 6 cm (2.36 in.) high. Likewise, at the distances of the observation tubes \( a \), \( b \), and \( c \) on both arms of the glass-tube fork before and behind the observation tubes there are 6 mm vertical tubes, all communicating with one another and, of course, also with the vertical tube in front of the measuring stick.

If this glass fork is filled with a liquid, preferably
Xylol on account of its concave meniscus, and the meniscus of the fork is made to coincide with the meniscus of each observation tube, we obtain the desired reading on the measuring stick. Since all the readings are made on the same scale, which undergoes no expansion, there are no reading errors to be corrected.

The tube \( c \) serves to determine the level of the lower mercury surface, which is below atmospheric pressure. The cock \( h \) provides an airtight shut-off for tube \( c \) during the experiment. The upper glass tube with the mercury seal comes into use only at high vapor pressures. It was not needed in these experiments. The glass tube \( b \) serves as an ordinary barometer and has, at the top, a perfectly fitted glass stopper with mercury seal. A thermometer gives the temperature of the barometer tube, which is affected by the higher experiment temperature in the tube \( a \) and necessitates a correction of the barometer column. The experiment tube \( a \) has an inside diameter of 16 mm (0.63 in.) and is provided with a cubic-centimeter scale for reading the vapor volumes.

After the vapor tensions are determined at the different temperatures, provision is made for the experiment tube \( a \) to be heated in each experiment by hot vapors of uniform temperature. For this purpose, tube \( a \) is surrounded with a glass cylinder, which is closed at the bottom by the cork \( p \). In order to prevent the bursting of the glass jacket at high temperatures, the cork is surrounded by a glass ring containing, in the
space thus formed, about 1 cm (0.4 in.) of mercury q, thus forming a perfect seal and preventing all tension in the glass cylinder. Two glass tubes k and l pass through the cork and terminate above the mercury in the heating jacket. Tube l introduces the heating vapors and extends about 5 cm (2 in.) into the heating jacket, while tube k returns the liquid from the condensed vapors to the heating flask. For convenient demounting, each of these tubes is in two parts with rubber connections capable of withstanding temperatures up to 200°C (392°F).

The heating is accomplished indirectly by means of electricity on account of the fire danger. A copper flask containing the heating liquid stands in an electrically heated vessel filled with rape oil and is closed with a three-hole cork. The tube k reaches to the bottom of the flask; the tube l, only into the vapor space at the top. A third glass tube m, reaching to the bottom of the flask, returns the condensed vapors to the flask. The heating vapors pass from the flask through the tube l into the heating jacket, from which they pass through the upper cork, to a return cooler connected with the water pipe, unless they condense before reaching the cooler. The glass tube m returns the condensed liquid to the copper flask. The circuit of the heating vapors is therefore a closed one and the loss of heating liquid is small. The apparatus is assembled by rubber connections, so that it can be readily taken apart and put together again.
The regulation of the electric heating is readily accomplished by switched-in resistances, so that the vapor production proceeds very uniformly. The temperature of the heating vapors is taken in the heating jacket by a thermometer calibrated at the "Phys.-Techn. Reichsanstalt" in Berlin. The thermometer is suspended on a wire, so that it can be easily raised and lowered from without.

The following chemically pure substances, with constant boiling points, were employed as heating liquids:

<table>
<thead>
<tr>
<th>Substance</th>
<th>Boiling point</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ether</td>
<td>0°C 64°F</td>
</tr>
<tr>
<td>Carbon disulphide</td>
<td>44°C 111.2°F</td>
</tr>
<tr>
<td>Methyl alcohol</td>
<td>63°C 145.4°F</td>
</tr>
<tr>
<td>Benzol (benzene)</td>
<td>73°C 172.4°F</td>
</tr>
<tr>
<td>Water</td>
<td>100°C 312.0°F</td>
</tr>
<tr>
<td>Amyl alcohol</td>
<td>130°C 266.0°F</td>
</tr>
<tr>
<td>Turpentine</td>
<td>160°C 320.0°F</td>
</tr>
<tr>
<td>Aniline</td>
<td>178°C 352.4°F</td>
</tr>
</tbody>
</table>

The boiling points changed somewhat during the course of the experiments, partly due to changes in the barometric pressure and partly due to residues of the heating liquid last used. This circumstance did not affect the results, however, because the temperature of the heating vapors was known.
Preparation and assembling of the apparatus.—Before inserting the three glass tubes, a, b, and c, the apparatus was carefully dried by drawing through it for about 30 minutes, air which had been previously dried by calcium chloride. Mercury that had been well boiled and then cooled in a dessicator was now poured into the mercury reservoir until it rose above the three packing boxes. By lowering the mercury reservoir, the apparatus could be entirely filled, without too much mercury rising above the packing boxes. The mercury reservoir was then closed with a rubber stopper, into which a calcium chloride tube was inserted, so that no moisture could enter. The tubes were then inserted, after having been previously heated to expel any moisture adhering to their walls. The level of the mercury in the reservoir R could be verified by comparison with the standard barometer in the physical laboratory. The large cork for the lower mercury seal was then put in place; the glass tubes were inserted in the packing boxes; the latter were tightened; and the apparatus was closed. Raising and lowering the mercury reservoir several times showed whether there were any air bubbles left in the connecting tube as far as the glass tubes a and b. In case there were, they could be allowed to escape by opening the glass stoppers at the tops of the tubes a and b. After this preliminary testing of the apparatus, the cock i was closed with the mercury reservoir raised and the mercury seals were produced for the tubes a and b.
After these preparations, the oil to be tested could be introduced into the experiment tube a. For this purpose a thinly blown glass bulb, ending in a capillary tube about 1 cm (0.4 in.) long, was used. After the bulb was weighed empty, it was warmed slightly and immersed in the oil, which was drawn into it as a result of the cooling. By repeating this process several times, the bulb and capillary were entirely filled. The weight of the oil was then found by weighing the filled bulb and deducting the weight of the empty bulb. The mercury reservoir was then raised so high that on removing the stopper at the top, the mercury flowed into the widened part of tube a. The filled bulb, together with the stopper, could then be easily pressed into the tube.

After making sure that the stopper was tight and that the mercury seal was all right, the mercury reservoir was lowered, thereby producing the vacuum in which the evaporation immediately began. The capillary tube was left open so that the evaporation did not depend on the bursting of the bulb. The evaporation proceeded smoothly in the vacuum without disturbance or delay. After the liquid had been introduced into the tube, the heating jacket was put in place, the mercury seal was produced at the bottom, the cork was inserted at the top and the connections were made with the heating flask and the return cooler.

In beginning the experiments, readings of the mercury level were taken at the room temperature for the different posi-
tions of the mercury receptacle. The experiment was then begun with ether for the heating liquid and with a delay of about 30 minutes before each reading, for the whole heating jacket to become evenly heated and until it could be assumed that the experimental fluid in the vacuum had acquired the same temperature. Before each experiment the barometer stand in the apparatus was found by reading the lower mercury level (which communicated with the mercury reservoir) on the tube c with the cock i open and also the position of the mercury in the barometer tube b at the top. The difference between the level of the mercury in the barometer tube and in the experiment tube gave the vapor pressure at the temperature of the heating jacket. Since this vapor pressure did not remain constant on lowering the mercury reservoir, even though liquid fuel were still present in the vacuum (which, according to the explanation already given, could not be expected to be otherwise), the vapor volumes had to be read on the experiment tube simultaneously with the vapor pressures. The following readings were accordingly made:

a) For the vapor-pressure determinations:

1) Mercury level in barometer b,
2) " " experiment tube a,
3) Vapor volume (cm³) in tube a,
4) Temperature in heating jacket,
5) " " barometer tube,
6) " of room,
7) Barometer.
b) For the vapor-density determination:

1) Vapor volume \( (\text{cm}^3) \), after all the liquid was evaporated,

2) Weight \( G \) of the evaporated fuel.

Translation by Dwight M. Miner,
National Advisory Committee
for Aeronautics.
1 Gasoline  
2 Benzol  
3 Coal–tar oil I  
4 Solar oil  
5 Kerosene  
6 Rositzer fuel oil  
7 Coal–tar oil II  
8 Gas oil  
9 Paraffine oil

**Fig. 1** Evaporation curves.
Fig. 2  Apparatus for determining vapor pressure and vapor density