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NATIONAL ADVISORY COMMITTEE FOR AERONAUTICS 1176 00091 1827 **WARTINE REPORT** ORIGINALLY ISSUED June and September 1945 as Memorandum Reports E5F20 & E5I12 SMOKING CHARACTERISTICS OF VARIOUS FUELS AS DETERMINED BY OPEN-CUP AND LABORATORY-BURNER SMOKE TESTS

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# NACA

#### WASHINGTON

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#### NACA MR Mos. E5F20, E5I12

#### MATIONAL ADVISORY COMMITTEE FOR AEROFAUTICS

MEMORANDUM REPORT

for the

Air Technical Service Command, Army Air Forces

SYOKING CHARACTELISTICS OF VARIOUS FUELS AS DETERMINED

BY OPER-CUP AND LABORATORY-BURNER SMOKE TESTS

By Earl R. Ebersole and Henry C. Barnett

#### ILTRODUCTION

At the request of the Air Technical Service Command, Army Air Forces, the Cleveland laboratory of the MACA is conducting tests in an I-15 combustion chamber to obtain performance data on a variety of hydrocarbon fuels.

Inasmuch as the smoking tendency of jet-propulsion fuels is of interest, a laboratory program has been conducted to obtain data on the smoking behavior of fuels. The results of open-cup smoke tests of 25 hydrocarbon fuels and of two series of prepared blends are presented. In order to determine the effect of fuel-air ratio as well as hydrocarbon class on the smoking behavior of fuels, 21 hydrocarbons were tested in a laboratory burner at fuel-air ratios ranging from 0.062 to 0.119. The results of these controlled-burning smoke tests are also presented. Correlation with data from full-scale jetpropulsion tests will be necessary to determine the applicability of the results.

#### APPARATUS AND TES" PROCEDURE

<u>Open-cup tests</u>. - The apparatus used in the open-cup tests is shown schematically in figure 1. A 3-milliliter sample of the fuel to be tested was placed in the open cup and ignited with an open flame. Snoke passing up the chimney absorbed a portion of the light beam from the light source. The effect of this light absorption on the photoelectric cell was registered by the recording microammeter. Before each test, the recording microammeter was adjusted to zero, if necessary, by varying the resistance in the lamp circuit. A check run was made on each fuel. The readings in microamperes were converted to percentage light absorbed by means of the calibration curve shown in figure 2. In the calibration of the recording instrument the light beam was interrupted with screens of varying mesh. The transmission through these screens was measured in a standard spectrophotometer using light with a wavelength of approximately 6000 A. The percentage light absorbed was then plotted against microamperb readings to obtain the calibration curve.

Each of 25 hydrocarbon fuels was burned in an open cup in order to determine snoking characteristics. In each test the burning rate of the fuel was determined by measuring the time required to burn a 3-milliliter sample.

The fuels tested included representative hydrocarbons of the paraffinic, cycloparaffinic, olefinic, and aromatic classes as well as mixed fuels. Kerosene and the same kerosene from which aromatics had been removed were included in the mixed fuels tested. Two series of binary blends consisting of toluene in hydrogenated triisobutylene and triisobutylene in hydrogenated triisobutylene were tested over the range of concentrations from 0 to 100 percent.

Laboratory-burner tests. - The burner and the auxiliary apparatus used in the laboratory-burner tests are shown schematically in figure 3. Primary air was passed through the preheater at a constant flow rate of  $4.1 \pm 0.1$  liters per minute. Secondary air was introduced at the base of the burner at a constant flow rate of  $10.4 \pm 0.2$  liters per minute. The fuel to be tested was placed in a graduated burette, which was modified to give low and constant flow rates. (See reference 1.) The fuel was then admitted to the primary air stream and was vaporized and mixed with the primary air in the preheater at a temperature of  $180^{\circ}$  to  $200^{\circ}$  F above the boiling point of the hydrócarbon. After the fuel was ignited through the small hole in the side of the chimney, the hole was sealed off by means of a slip ring.

The flow rate of the air was maintained at  $1^{l_2}.5 \pm 0.3$  liters per minute for all fuel-air ratios. The fuel-air ratio was varied by changing the flow rate of the fuel from the graduated burette. The rate of fuel flow was measured by determining to the nearest 0.01 minute the time required for a given volume (0.825 ml) to flow from the burette.

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Each of 21 fuels including representative hydrocarbons of paraffinic, cycloparaffinic, olefinic, aromatic, and dicyclic classes was tested for smoking tendency at various fuel-air ratios. Smoke particle size was not measured in these tests.

#### DISCUSSION OF RESULTS

<u>Open-cup tests</u>. - The results of the open-cup smoke tests are presented in table I. Smoke is reported as percentage light absorbed. Total smoke from a 3-milliliter samplo was determined by measuring the area under the curve traced by the recording microammeter. Values of average smoke were computed by dividing the total smoke by the burning time. Values of peak smoke were obtained by visually averaging the deflections in the highost portion of the curve.

The data in table I show that, with the exception of the cycloparaffins, the fuels of a given hydrocarbon class gave readings within a range of 8 porcont light absorbed and that the argumetics gave higher readings than the other classes. The greater smoking tordency of the aromatics can be further shown by the tests on kerosene and dearonatized kerosene. Keroseno gave a pack value of 21 percent light absorbed as compared with 5 percent for dearomatized kerosene.

In order to ascertain the offect of burning rate on the smoking tendency, average smoke ( porcentage light absorbed) was plotted against burning rate (grams per min) in figure 4. Both test run and check run for each fuel are plotted. The five-carbon ring cycloparaffins are not shown in this figure because data were obtained for only two compounds in this series. It can be seen in table I that considerable difference in smoking tendency was found between the five-carbon ring and the six-carbon ring cycloparaffins. The fuels tested (fig. 4) show that the type of hydrocarbon exerts a much greater influence on the smoking tendency than does the burning rate of the fuel.

Figure 5 shows the variation of the burning rats of the fuels with boiling point. As in figure 4 both test and chock runs are plotted. For the fuels testod, the burning rate of hydrocarbons within a givon class decreased with increasing boiling point. It is of interost that two fuels of different classes with different boiling points will burn at the same rate; for example, from the curves of figure 5, a paraffinic hydrocarbon with a boiling point of 176° F will burn at approximately the same rate as an aromatic hydrocarbon with a boiling point of  $305^{\circ}$  F.

In order to show the relation of smoking tendency and composition for fucls having equal boiling points, tests were made on blends of triisobutylene in hydrogenated triisobutylene. A linear relation was obtained. (See fig. 6.) Figure 7 shows the results of tests of two fuels (toluene and hydrogenated triisobutylene) having different boiling points.

Laboratory-burner tests. - The results of the laboratoryburner smoke tests are presented in table II together with the boiling point and the stoichiometric fuel-air ratio of each fuel tested. At the maximum fuel-air ratios that would support combustion in the burner no smoke was obtained for parafrinic, cycloparaffinic, or olefinic hydrocarbons with the exception of triisobutylene, which gave detectable smoke at a fuel-air ratio of 0.102.

Aromatic and dicyclic hydrocarbons began smoking at approximately their stoichiometric fuel-air ratios. Figures 8 and 9 show the variation of smoke (percentage light absorbed) with fuel-air ratio for aromatic and dicyclic hydrocarbons, respectively. Korosene is also included in figure 9. Within exporimental error the smoking tendency appears to be a linear function of the fuel-air ratio for aromatic and dicyclic hydrocarbons. Comparison of figures 8 and 9 indicates that the smoking tendency of nayhthalenic hydrocarbons is greater than that of aromatic hydrocarbons.

#### SULLARY OF RESULTS

The data obtained from an investigation of the smoking characteristics of 25 hydrocarbons indicate that for uncortrolled burning in an open cup:

1. The smoking tendency of a hydrocarbon fucl is more dependent upon the type of hydrocarbon than upon its boiling point or burning rate.

2. The burning rate of hydrocarbons within a given class tended to decrease with increasing boiling point.

3. The smoking tendency of a commercial kerosene was about four times that of the same kerosene from which aromatics had been removed. Results obtained from controlled-burning smoke tests indicate that:

1. The smoking tendency of hydrocarbon fuels is dependent upon both the class of hydrocarbon and the fuel-air ratio.

2. Mithin experimental error the smoking tendency of aromatic and dicyclic hydrocarbons is a linear function of the fucl-air ratio.

The following table summarizes the data obtained:

Class	Boiling point (°F)	Hydrogen- carbon retio	Foak smoke (porcont light absorbed)	Burning rate (grams/min)
Paraffins Six-carbon ring cyclopareffins	122-230 175-2.9	0.159-0.196 .1(7	10–15 8–13	0.69-1.07 .6285
Five-carbon ring cycloparaffirs	12?-151	.167	23-31	.97-1.13
Olofins Arom: tica	101-172 230-323	.167 .09 <u>5</u> 111	цо-47 68-75	.97-1.33 .75-1.36

OPEN-CUP SMOKE TESTS

#### LABORATORY-BURGER SMOKE CESTS

Hydrocarbon class	Bciling- point range (°F)	Stoichiometric fuel-air ratio	Fuel-air-ratio range	Smoke (norcent light absorbed)
Paraffins	175-350	0.035-0.066	0.064-0.114	0
Cyclohexanes	178-270	.067	.073115	0
Olofins	133-350	.067	.062109	0-3
Aromatics	175-465	.070075	.063108	0-47
Dicyclics	365-605	.069077	.069119	3-80

Aircraft Engine Research Laboratory,

National Advisory Committie for Aeronautics,

Cleveland, Ohio, June 20, 1945; Sept. 12, 1945.

#### REFERENCE

 Zentner, E. Thomas: Delivory of Liquids at Low and Constant Ratos. Ind. and Eng. Chem. (Anal. ed.), vol. 16, no. 7, July 21, 1944, pp. 471-472.

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#### NACA MR No. E5F20

#### TABLE I. - RELATIVE SNOKING CHARACTERISTICS OF

Class of hydro- carbon	Compound	Peak amoke observed (percent light absorbed)	Average smoke (percent light absorbed)	(area in sq	burning (grams/		Hydrogen- carbon ratio
Paraffins	2,2-Dimethyl-	11	10	1.4	1.052	122	0.196
	butane 2,3-Dimethyl-	13 11	11 11	1.7 2.0	1.076	137	.196
	butane 2,2,3-Tri- methylbutane	13 15 15	10 11 11	1.7 2.2 2.4	.978 .805 .779	1 <b>7</b> 8	.192
	2,3-Dimethyl- pentane	10 10	76	1.4	.772	193	.192
	2,2,3-Tri- methylpentane	13 11	10 8	2.5 2.0	.679 .688	230	-189
Cyclo- paraffins	Cyclopentane	26 23	20 18	3.6	1.125	122	0.167
F	Methylcyclo- pentane	31 31	26 24	5.4 5.8	.965 .965	161	.167
	Cyclohexane Methyl-	8 10 11	8 7 8	1.8 1.5 2.0	.825 .849 .755	178 212	.167 .167
	cyclohexane Ethyl- cyclohexane	13 11 13	10 9 8	2.5 2.8 2.3	.787 .617 .517	269	.167
Olefins	Trimethyl-	42	37	5.8	1.325	101	0.167
	ethylene 2,3-Dimethyl-	44 44	37 37	5.8 6.4	1.325	133	.167
	butene-1 2,3-Dimethyl- butene-2	43 40 42	37 33 35	6.6 6.9 7.6	1.223 1.026 .968	164	.167
	Triptene	46 47	38 38	8.2 8.2	1.063 1.063	172	.167
Cyclo- olefins	Nethyl- cyclohexene	50 47	41 37	8.0 8.1	1.323 1.142	216	0.143
Aromatics	Toluene	71 68	50 49	10.7	1.363	230	0.095
	Xylene isomers	68 75	52 57	18.3 19.1	.844 .893	279-288	
	n-Propyl- benzene	70 70	52 54	18.7	.816 .816	320 323	•111 •111
	m-Ethyl- toluene	70 72	51 55	19.4 22.7	•763 •745	525	•111
Mixed fuels	Virgin base stock	21			0.698	115-298	
	Hot-acid octanes Triiso-	13 35			.652	174-257 340-350	
	butylene <sup>a</sup> Hydrogenated	16				335-350	
	triiso- butylene <sup>®</sup> Kerosene Kerosene	21 5				302-486	
	(dearo- matized)	5			NAT COMMITTI	IONAL ADV	I SORY RONAUTICS

#### VARIOUS FUELS BURNED IN OPEN CUP

<sup>a</sup>Listed as mixed fuels because their purity is questionable.

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### NACA MR No. E5II2

Class of hydrocarbon	Compound	Boiling point (°F)	Stoichi- ometric fuel-air ratio	Fuel-air- ratio range	Hydrogen- carbon ratio	Smoke (percent light absorbed)
Paraffins	Triptane	178	0.065	0.067-0.103	0.191	0
	Diisopropyl	137	.065	.064071	.195	0
	Neohexane	122	.065	.070085		0
	2,2,3-Trimethyl- pentane	230	•066	.074083	.188	0
	Hydrogenated triisobutylene	335-350	• 066	.083114	.181	0
Cyclohexanes	<b>Cyclohexane</b>	178	.067	.073093	.167	o
•	Ethylcyclo- hexane	269	•067	.073115	.167	o
Olefins	2,3-Dimethyl- butene-1	133	.067	.080096	.167	o
	2,3,3-Trimethyl- butene-1	172	.067	.062090	.167	0
	Triisobutylene	340-350	.067	.092109	.167	0-3
Aromatics	Benzene	176	.075	.063104	.063	0-37
	<b>Sthylbenzene</b>	277	.073	.076100		3-34
	Isopropylbenzene		.072	.063100		0-32
	1,2,4-Trimethyl- benzene	337	.072	.076108	.111	3-47
	Triisopropyl- benzene	465	.070	.073107	.133	5-44
Dicyclics	l-Methyl- naphthalene	473	.077	.073114	.076	3-80
	Dimethyl- naphthalene	511	.075	.078102	.083	11-59
	Noncamyl- naphthalene	535-608	.073	.069097	.101	9-50
	Tetralin	405	.073	.076108	.101	7-61
	Decalin	365-882	.069	.084119	.150	3-18
Mixed fuels	Kerosene	302-486		.060106		0-5

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#### TABLE I. - SMOKING CHARACTERISTICS OF HYDROCARBON FUELS

DETERMINED FROM CONTROLLED-BURNING TESTS

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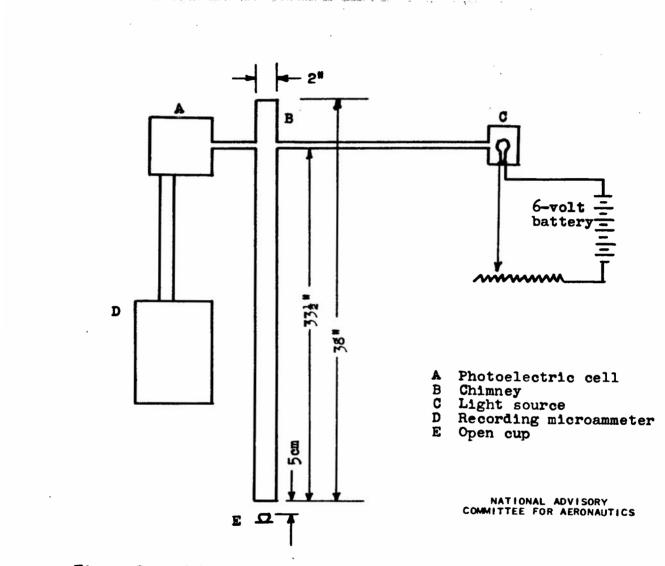


Figure 1. - Schematic diagram of apparatus for open-cup smoke tests.

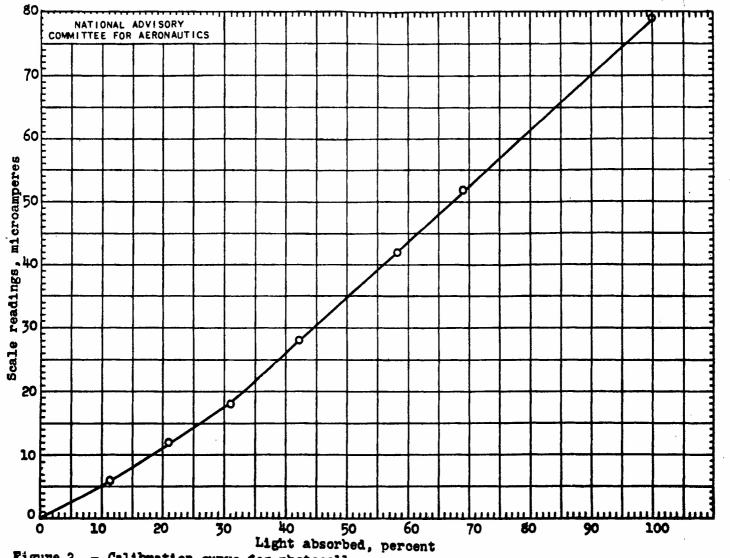
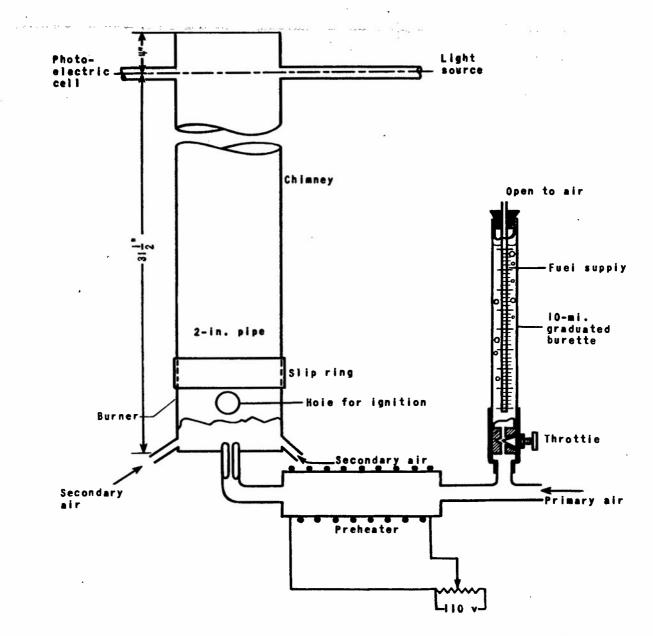


Figure 2. - Calibration curve for photocell.

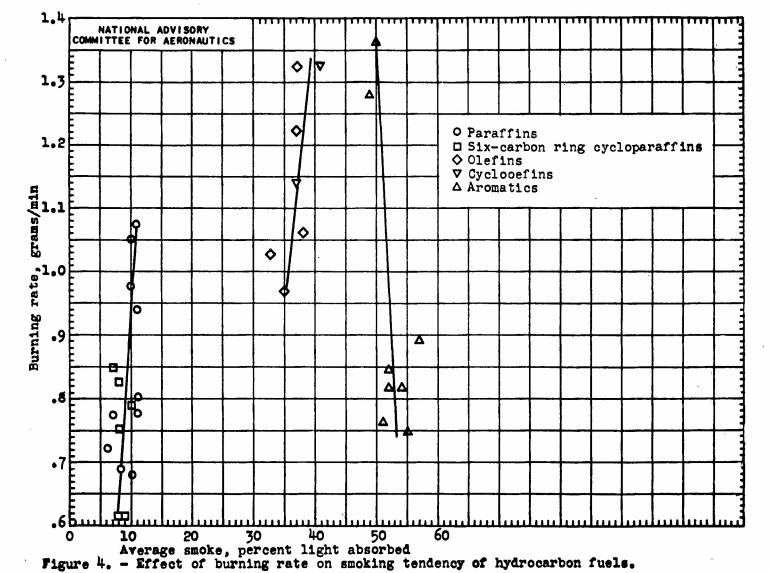
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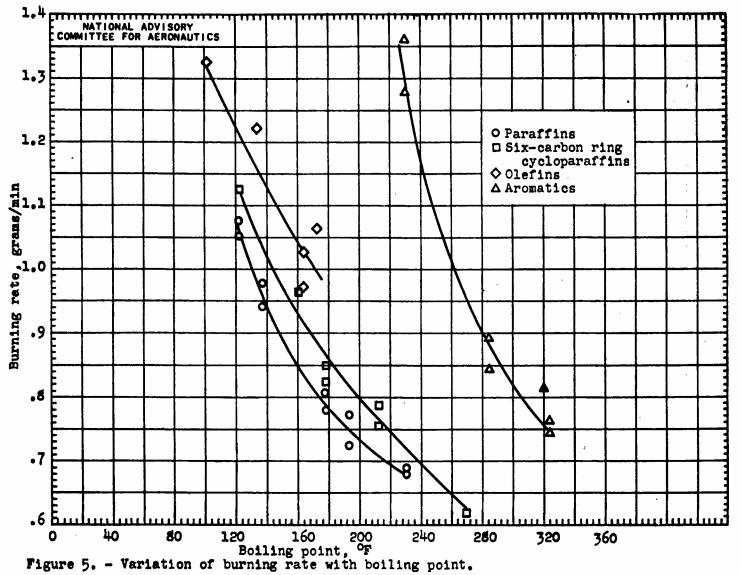


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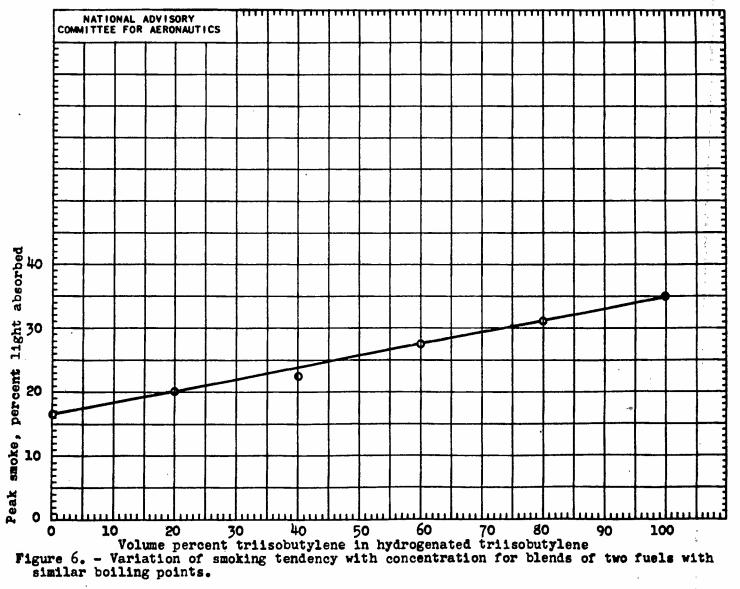
Figure 3. - Schematic diagram of burner and auxiliary apparatus.



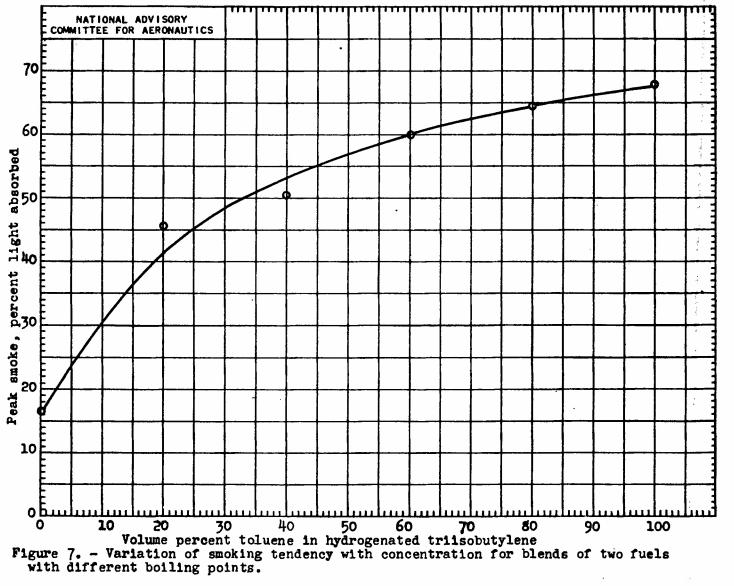
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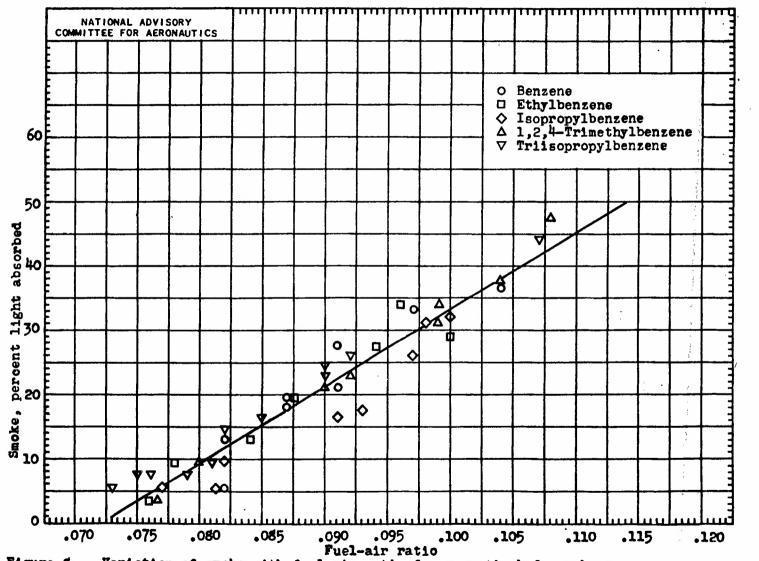


Figure 8. - Variation of smoke with fuel-air ratio for aromatic hydrocarbons.

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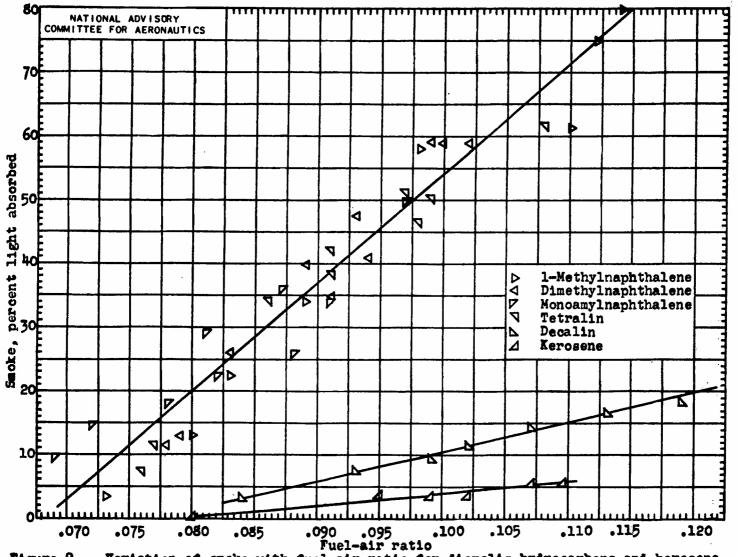


Figure 9. - Variation of smoke with fuel-air ratio for dicyclic hydrocarbons and kerosene.

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