

**ENCLOSURE (B) 27**

**STUDIES ON THE OILINESS  
CHARACTERISTICS OF PURE HYDROCARBONS  
BASED ON STATIC FRICTION  
DETERMINATIONS FOR STEEL ON STEEL**

by

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SUMMARY

The oiliness of various cyclic hydrocarbons and a few chain hydrocarbons were measured and the following results were obtained.

1. If aromatic hydrocarbons are hydrogenated step by step, the oiliness of the compounds becomes first poorer and then better.
2. Qualitative correlations seem to exist between the oiliness of the compounds and their molecular volumes and molecular cohesion, i.e., melting point, boiling point, viscosity, etc.
3. The compounds in which the benzene nuclei are combined by single bonds were better lubricants than those having the benzene nuclei combined in condensed form.
4. The oiliness of a cyclic compound was better than that of a chain compound having the same number of carbon atoms.

I. INTRODUCTION

It has been reported in the literature that paraffinic hydrocarbons have better oiliness characteristics than naphthenes (Ref. 1.), and that in the case of straight chain hydrocarbons the oiliness improves as the molecular weight increases (Ref. 2.), and that chain hydrocarbons are better lubricants than cyclic compounds (Ref. 3.), but almost no systematic studies on the oiliness of pure cyclic hydrocarbons have been published. The molecules of mineral lubricants are said to have a cyclic structure with side chains of various lengths, and a systematic study of the oiliness of the pure cyclic hydrocarbons is important in this branch of the investigation of lubricants.

II. DETAILED DESCRIPTION

Studies on these problems were conducted during 1944.

A. Test Procedure and Samples Used in the Test

1. Test Apparatus. The three test pieces of the Doeley machine were changed and three steel balls (dia.  $\frac{1}{4}$  inch) were substituted. The test plate was made of cast steel.

2. Preparation of the Test Plate and the Test Pieces. The test pieces and the test plate were first washed with pure petroleum ether and ethylether, polished with 0.5 emery paper, and then washed with distilled water.

They were next purified by means of electrolytic reduction, applying 20 mA. current, for 30 minutes in a 2% NaOH solution using a carbon anode. Thus treated, the test plate and pieces were washed with water and alcohol, and dried in vacuum.

The oil test sample was rubbed on the surface with a piece of filter paper, and measurement was made 30 minutes after the test pieces were wetted with the oil.

3. Test Procedure. Static coefficients of friction of various

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pure hydrocarbon oils, alone and in 5% benzene solution, were measured at room temperature or at temperatures higher than their melting points.

4. Compounds Used for the Test. Marketed samples of benzene, cyclohexane, naphthalene, tetralin, decalin, and anthracene were purified and used in the test.

Diphenyl and diphenyl benzene were prepared through the thermal condensation of benzene (Ref. 4), and the hydrogenation products of diphenyl, anthracene, and diphenyl benzene were prepared by high pressure hydrogenation (Ref. 5).

The compounds used and their characteristic properties are summarized in Table I(B)27.

#### B. Results

The results are summarized in Table II(B)27 and are represented graphically in Figure 1(B)27.

1. The substances which had better oiliness were also better oiliness compounds in a benzene solution containing 95% by volume of benzene.

2. Benzene Series:- Benzene, cyclohexane. Cyclohexane had better oiliness than benzene as already reported (Ref. 6).

3. Naphthalene Series:- Naphthalene, tetralin, decalin. Contrary to the case of the benzene series, naphthalene had better oiliness than tetralin. Such results may be due to the greater molecular cohesion of naphthalene. In the case of the benzene series, benzene has a greater molecular cohesion than cyclohexane, but the former had poorer oiliness, and such results are attributed to the smaller molecular volume of benzene. Decalin, however, had better oiliness than tetralin. This is similar to the relationship between cyclohexane and benzene. In brief, if naphthalene is hydrogenated step by step, its oiliness becomes first poorer and then better.

#### 4. Other Series

a. Diphenyl Series:- Diphenyl, phenoxylohexane, dicyclohexyl.

b. Anthracene Series:- Anthracene, tetrahydroanthracene, decahydroanthracene, perhydroanthracene.

c. Diphenyl Benzene Series:- Diphenyl benzene, perhydrodiphenyl benzene.

The change of oiliness resulting from stepwise hydrogenation of these other compounds gave results which resembled those of the naphthalene series.

#### 5. General Remarks on the Above Results

a. From the above results, it may be said that, in general, if aromatic hydrocarbons are hydrogenated step by step, the oiliness of the compounds becomes poorer at first and then improves when hydrogenation is complete. Also, qualitative

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correlations seem to exist between the oiliness of the compounds and their molecular volumes and molecular cohesion, i.e. melting point, boiling point, viscosity, etc. Such relations are given in Figures 1(B)27 and 2(B)27.

b. The oiliness of the hydrocarbons becomes gradually better in the following order:

Benzene series → Naphthalene series → Diphenyl series  
Anthracene series → Diphenyl benzene series.

c. The compounds having the benzene nuclei combined with single bonds were better lubricants than the compounds having the same number of nuclei combined in condensed form.

6. Comparison With Chain Compounds. The oiliness of n-hexane, i-hexane, and n-octadecane were measured and the results were compared with those of cyclic compounds having the same number of carbon atoms.

The characteristics of the compounds and the results are given in Table III(B)27 and Figure 3(B)27.

It appears that the oiliness of a cyclic compound is better than that of a chain compound having the same number of carbon atoms.

### III. CONCLUSIONS

From the above results, it was concluded that naphthalene, diphenyl, anthracene, diphenyl benzene, perhydrodiphenyl benzene had good oiliness characteristics, and of these, completely hydrogenated diphenyl benzene was the best.

### References

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6. or. (3)

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Table I (B) 27  
PROPERTIES OF THE RELATED COMPOUNDS

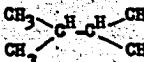
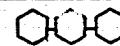
Group	Compound	Molecular Structure	M. C.R.	P. Lit.	B. Obs.	P. Lit.	Elementary Analysis		
							Obs.	Cal.	% Obs.
Benzene Series	Benzene		5.4	5.5	79-80	79.6			
Cyclohexane	Cyclohexane		4.5	4.5	80-81	80.7			
Diphenylbenzene	Diphenylbenzene		79-80	80	217-219	218			
Diphenylbenzene	o-Diphenylbenzene		114	-31	204-200	206			
Diphenylbenzene	m-Diphenylbenzene		114	-125	188-193	189-191			
Diphenylbenzene	p-Diphenylbenzene		114	-	-	-			
Diphenylbenzene	Diphenylbenzene		69-70	68.5-69.5	-	24.8-24.9			
Diphenylbenzene	Diphenylbenzene		114	7	234-236	234-236			
Diphenylbenzene	Diphenylbenzene		114	4	229-235	225-235			
Diphenylbenzene	Diphenylbenzene		217-218	217-218	-	351-342			
Diphenylbenzene	Diphenylbenzene		104.5-107	103-105	-	309-313	92.86	92.26	6.76
Diphenylbenzene	Diphenylbenzene		39	73-77	-	292-295	89.31	90.25	9.63
Diphenylbenzene	Diphenylbenzene		114	114	-	272-277	86.42	87.11	13.26
Diphenylbenzene	Diphenylbenzene		211.5-212	212-213	-	253	-	-	-
Diphenylbenzene	Diphenylbenzene		160.161.5	162-163	-	-	85.32	87.03	22.36
Diphenylbenzene	Diphenylbenzene		160.161.5	162-163	-	-	85.32	87.03	22.97

Table III(B)27  
STATISTICAL INFLATION CORRELATION COEFFICIENTS AND OTHER PROPERTIES

Compound	Molecular Structure	Percentages of Compounds				$\frac{1}{M_p}$ (c)	Vis. (Poise) of 5 mol solution in benzene		
		100%		5% solution in benzene					
		Static G.F. (%)	$T_{\text{bp}}^{\text{mp}}$ (°C)	Static G.F. (%)	$T_{\text{bp}}^{\text{mp}}$ (°C)				
Benzene	<chem>C</chem>	0.170	13	0.164	22	0.0127	0.00358		
Cyclohexane	<chem>C1CCCCC1</chem>	0.153	13	0.160	22.5	0.0116	0.00358		
Heptahexane	<chem>C1=CC=CC=C1</chem>	0.144	90	0.147	22.5	0.00980	0.00272		
Tetralin	<chem>C1=CC=CC=C1</chem>	0.159	90	0.159	24.2	0.00956	0.00243		
Diphenyl	<chem>c1ccc(cc1)-c2ccc(cc2)C</chem>	0.154	90	0.153	23.2	0.00723	0.00676		
Diphenyl-1	<chem>c1ccc(cc1)-c2ccc(cc2)C3=CC=CC=C3</chem>	0.134	75	0.152	25.2	0.00649	0.00292		
Phenylhexane	<chem>Cc1ccccc1Cc2ccccc2</chem>	0.149	24	0.153	22.6	0.00625	0.00357		
Diphenyl-2	<chem>c1ccc(cc1)-c2ccc(cc2)C3=CC=CC=C3</chem>	0.156	75	0.153	23.0	0.00601	0.00361		
Acetophenone	<chem>CC(=O)c1ccccc1</chem>	0.121	230	0.129	17.6	0.00561	0.00204		
Tetraphenyl- acetophenone	<chem>CC(=O)c1ccccc1c2ccccc2c3ccccc3</chem>	0.135	105	0.153	18	0.00557	0.00204		
Decaphenyl- acetophenone	<chem>CC(=O)c1ccccc1c2ccccc2c3ccccc3c4ccccc4c5ccccc5c6ccccc6</chem>	0.139	105	0.156	9.8	0.00536	0.00302		
Penta-phenyl- acetophenone	<chem>CC(=O)c1ccccc1c2ccccc2c3ccccc3c4ccccc4</chem>	0.142	105	0.162	27.6	0.00520	0.00370		
Decaphenyl benzene	<chem>c1ccc(cc1)-c2ccc(cc2)C3=CC=CC=C3c4ccccc4c5ccccc5c6ccccc6</chem>	0.122	160	0.141	23.5	0.00442	0.00206		
Penta-phenyl benzene	<chem>c1ccc(cc1)-c2ccc(cc2)C3=CC=CC=C3c4ccccc4</chem>	0.104	170	0.126	25.2	0.00403	0.00230		
Penta-phenyl benzene	<chem>c1ccc(cc1)-c2ccc(cc2)C3=CC=CC=C3c4ccccc4</chem>	0.104	170	0.126	25.2	0.00403	0.00230		

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Table III(B)27  
COMPARISON OF OILINESS OF CHAIN AND CYCLIC COMPOUNDS

Name	Chemical Structure	Density (d <sub>4</sub> <sup>20</sup> )	M.P. (°C)	B.P. (°C)	Static Coef.*	Temp. (°C)
n-Hexane	$\text{CH}_3(\text{CH}_2)_4\text{CH}_3$	0.6992		68-70	0.161	12.8
i-Hexane		0.6630		57-59	0.160	12.6
Cyclohexane		0.7783		79-81	0.153	13.0
n-Octadecane	$\text{CH}_3(\text{CH}_2)_{16}\text{CH}_3$		28	158-162 15.5mm	0.132	45.0
Parahydrodiphenyl benzene			160-161		0.104	170

\*By modified Deesley machine. Steel ball on steel plate.

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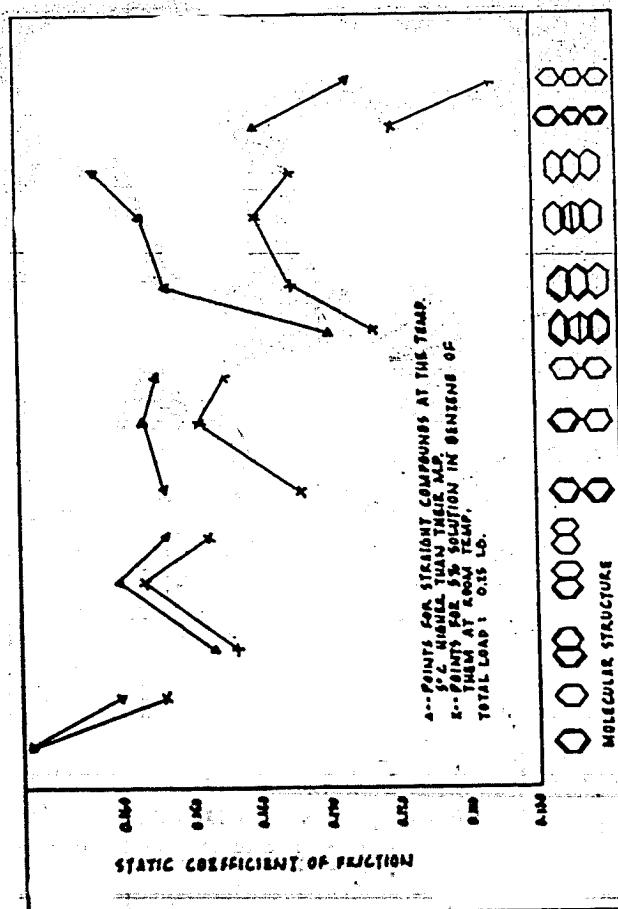


Figure 1(B)27  
STATIC COEFFICIENT OF FRICTION BY MODIFIED DEELEY MACHINE  
(Point Contact, Steel on Steel)

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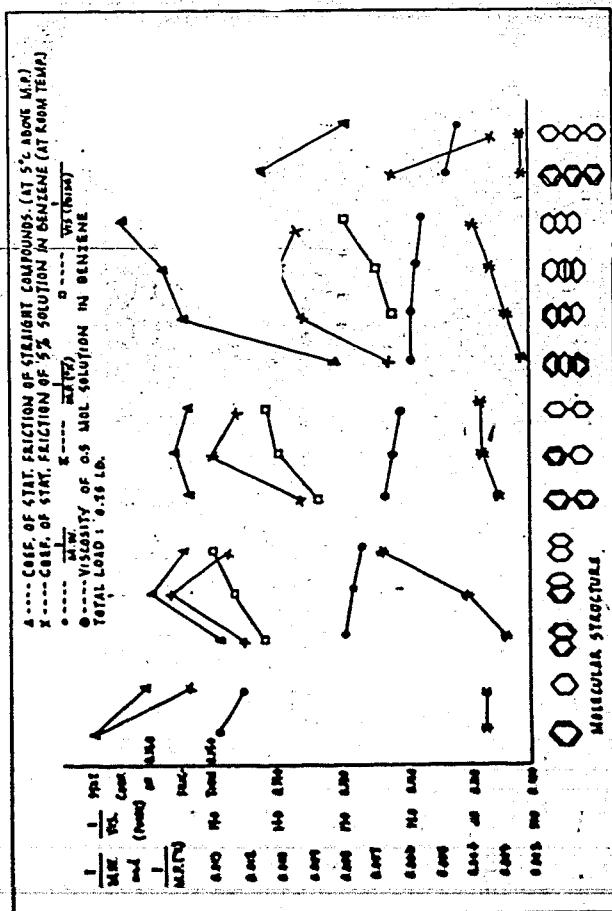


Figure 2(B)27  
CORRELATION OF THE STATIC COEFFICIENTS OF FRICTION BY DEARLY (MODIFIED) MACHINE  
(Point contact, steel on steel--with the molecular weight and viscosity of the compounds).

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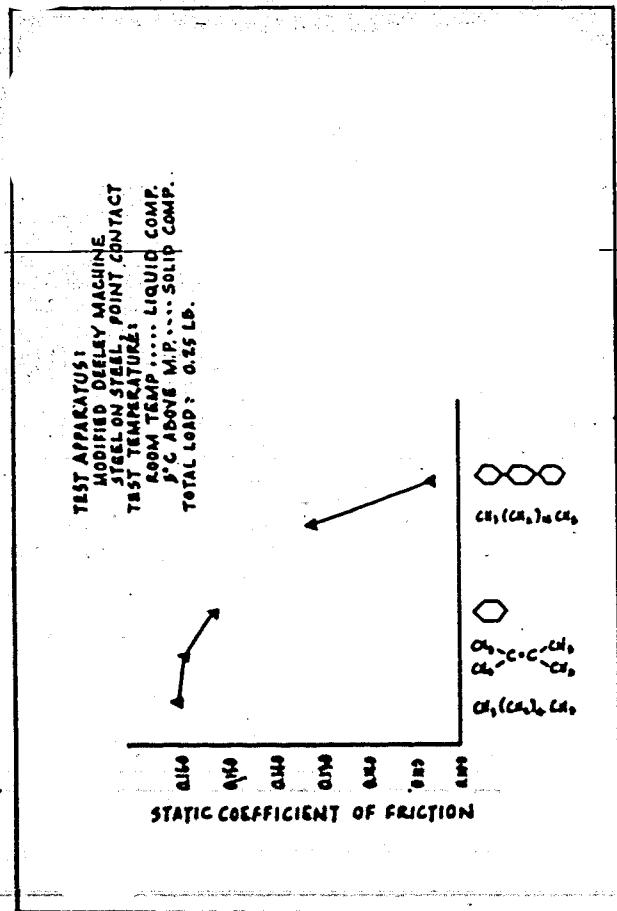


Figure 3(B)27  
COMPARISON OF CYCLIC COMPOUNDS WITH CYCLIC COMPOUNDS  
(Static Coefficient of Friction)