TO: C.I.O.S. Secretariat, 32 Bryanston Square, London, W.1.

SUBJECT: Visit of CIOS Teams 551 and 551a to Oil Targets in Leuna, Schkopau. Zeitz (Troglitz), Stassfurt, Heiligenstadt, Leverkusen, Sterkrade, Hochst and Heidelberg.

. and 55la - 25 June-15 July 1945:

rip 551.

Dr. W.F. Faragher, U.S. Team Leader. Capt. C.C. Chaffee, U.S. Ord., Deputy Leader.

Lieut. R.J. Osol, U.S. Ord.

Dr. Hans Schindler, U.S.

Dr. W.A.Horne, U.S.

Mr. J.G.Allen, U.S.

Dr. G.S.Bays, U.S.

.w Mr. B.L. MacKusick, U.S.

Trip 55la

Major D.A. Howes, British.

I.G. Farbenindustrie A.G., Leuna (Target 30/4.02).

Synthetic Lube Oil Manufacture.

Additional information has been obtained in the recovery of oil and related products from the aluminumchloride complex formed in the polymerization reaction.

The AlCl3 sludge obtained by centrifuging the autoclave product is treated with methanol, the oil obtained is neutralized, dried and treated again with AlCl3 (at about 70°C and atmospheric pressure). The AlCl3 sludge is again removed by centrifuging and the oil is neutralized, washed and distilled. The product so obtained is identified as "R" oil (Vis. at 100°C 2.5-3.0°E) and is used for less exacting lubricating purposes. The production of "R" oil amounted to about 700 tons per year.

Decomposition of the AlCl3 sludge obtained in the course of the refining of "R" oil results in the formation of a drying oil used in the protective-coating industry. This product is identified as "RR" oil.

For the manufacture of SS 906, aluminum chloride containing not more than 1% iron is used, whereas in the manufacture of SS 903, aluminum chloride with an iron content of 1.0-2.5% iron is suitable.

The following samples were obtained for further study:-

SS 906 (finished oil)
Light ends from SS 906 operation
Total product from SS 906 operation
Aluminum chloride used for catalyst
"R" oil
Acetylene hydrogenation catalyst used
in ethylene purification.

Toluene Manufacture from Benzene and Methanol.

Development work on the manufacture of toluene from benzene and methanol was carried out at Leuna and a plant operating in accordance with this method had been erected at Waldenburg. The Waldenburg plant had initially a capacity of 40,000 tons/year, but was later enlarged to a capacity of 50,000 tons/year.

The process consists in reacting 4 mols of benzene and one mol of methanol at 350°C and 35 atm., using as catalyst zinc phosphate on kieselguhr. The space velocity is about 0.25, but is reduced with increasing age of the catalyst. The lifetime of the catalyst is 4-6 weeks, during which time a carbon deposit of 30-40% is formed. The catalyst is not regenerated and removal of the catalyst from the reactor is difficult. Attempts to regenerate the catalyst were unsuccessful. A sample of this catalyst was secured for further study.

The methanol used is freed-from ammonia and amines by an organic cation exchanger. The benzene used is refined and is low in sulfur but frequently contained acetonitrile which is undesirable but could not be removed in a satisfactory way. The crude reaction product contains about 16-17% by volume toluene, 7% xylenes, 7% heavy residue, besides unreacted benzene. The amount of dimethylether is about 0.3% by volume (calculated as liquid product). The crude product was distilled by means of three columns packed with Raschig rings; the toluene was separated subsequently in a bubble tray column. The xylene fraction was

obtained by means of a column packed with Raschig rings. No detailed information on the distillation equipment was available. The maximum monthly toluene production in Waldenburg was 3800 tons of nitration-grade toluene, but decreased because of raw material and transport difficulties. The amount of xylene produced was about 25-30% by wt. of that of toluene and the quantity of alkylbenzenes produced was about the same as that of xylene.

DHD and HF Processes.

Samples of DHD catalysts and additional process details missed in the earlier visits were obtained. One drum (200 liters) of 5931, the standard DHD catalyst made at Leuna, and one drum (200 liters) of 7935, a slightly different preparation made at Ludwigshafen and not yet tested on a commercial scale, were collected. Details of the preparation of 5931 were also obtained, but similar information on 7935 was not available because it was not manufactured at Leuna.

The HF process is essentially the same as the DHD process except for the fact that the former operation is carried out at lower pressures and for shorter reaction periods between regenerations. The HF catalysts are the same as the DHD catalysts, except for lower MoO3 content. The Germans have found that the higher the paraffin content of the feed stock, the lower the operating pressure should be in order to minimize cracking of paraffins to The highly naphthenic feeds from hydrogenation were dehydrogenated at about 30-40 atm. in the DHD plant. Because of the low paraffin conent, and therefore of the low coke formation, an operating period of 160-200 hours between regenerations is practicable, and no automatic cycle-control equipment need be employed. For dehydrogenating crude-oil distillates of far lower naphthene content. a pressure of about 15 atm. is used. Goke yields are higher and regeneration must be more frequent. Reaction periods of only 10 hours are employed, which calls for automatic cycle controls; hence the HF process.

There is no HF plant at Leuna, but the catalyst is made there. Preparation details were obtained for 5% MoO3 content HF catalyst, but no samples were taken because of its similarity to the 10% HF-DHD material.

Lubricants and Inhibitors.

As a result of the inspection of plant and laboratory Tacilities at the Ammoniakwerk Merseburg at Leuna and the interrogation of Dr. Lisa Rossig, the following information was obtained on miscellaneous lubricants and inhibitors.

Miscellaneous Lubricants included machine-gun oil, Waffenel Blau 44; torpedo oil T-1; SS 1631, reported to be a "V" weapon oil; a special low-temperature oil used in the Russian campaign; coded K-10 and K-19; Y-axle oil, red in Color; and aviation hydraulic oil, coded Do 2000. The essential characteristics of these lubes follow:

<u>Lubricant</u> .			Specifications.			
	Visc. Eng.	Pour Pt.	V.I.	Flash Pt.oc	Blend Formula.	
Waffenol Blau —	1.9 at 20° C.				E-455 VT 120 Mesulfol-II	45% 45 10
Torpedo 1 Oil T-l a	1.5-12.5 t 20°C.	Below -50			E-515 KSE SS 903	63% 2 35
V weapon Oil SS 1631	2°E at 50°C.	Below -45°C		145°C	E-515 KSE SS 903	72% 3% 25%
Lube 0il- K-10		Below 50°C.	135	• -	E-515 SS 903	50% 50%
Lube Oil K-19	2.0 at 990C.	Below 45°C.	130		E-515 SS 903	25% 75%
Y Axle Oil		-49 to 58°C.	118- 122		E-504 R-oil Dye	20% 80% 1%
Hydrau- lic Oil Do 2000	1.75 at 20°C. 650 at -60°C.	-70°C.	<u>-</u> - 1	20°C.	E-3022 or E-3023 KSE V120 Dye 0	12% 4% 84% •006%

Inhibitors and Additives: E-455, Mesulfol II, E515, KSE, E504, E3022, and E3023. Information on these inhibitors follows:

Remarks. Chemical ${ t Product}$ Formula. (ROOC(CHg)z)2 Adipic acid ester is used in R is alcohol machine-gun oil in fairly from isobutanol large percentage. synthesis, BP 140-180°C -Mesulfol II (R-0-C-S-CH₂)₂ The additive is used in machine-gun oil and is used as an extreme pressure additive. ine-gun oil and is used as an R is C5 or C6 Addition of Mesulfol II is controlled to obtain a sulfur content of 3%, and copper test shows no discoloration after three days. E515 (RO-C(CH2)2)2 Adipic acid ester has been used in fairly large proportions in Torpedo Oil, R is alcohol SS K31, K-9, K-10 which are from isobutanol synthesis BP low-temperature lubes. --180-250°C∙ RSO2NHCH2COOR Additive is used as a corros-KSE ion preventive in Torpedo Oil, T-1; SS1631 and hydraulic oil. R is alcohol from isobutan-Should prove of interest as_ ol synthesis BP inhibitor in recoil oils. 180-250°C (ROOC(CH₂)₂)₂ Additive is added to R oil made R is alcohol from neutralized AlCl₃ complex, from isobutan obtained in the manufacture of E504 ol BP 160-200°C.synthetic lube. CH₃ ROOCCH₂CH Used in hydraulic oil Do2000 & ROOCCH2CH2 replaced by E3023

R = m-methyl cyclohexyl.

(ROOC(CH2)2)2 Used in hydraulic oil Do2000

R =cyclohexyl

Samples were secured of the following:

E-455 E-504
Mesulfol II E-3023
KSE Y-Axle Oil

Rust preventive information obtained is believed to be of importance to the Ordnance Department. In addition to the above data, a number of ZWB reports were found and are listed under "Documents".

Catalysts.

More complete analyses and details of the preparation and production of catalysts made or used by the Organic Division of Ammoniakwerk-Merseburg were obtained.

<u> Catalyst - Leur</u>	na No. Use. Production Tons/Mo.
616	Methanol synthesis 30 from CO and Hg
1132	Isobutanol synthesis 60 from CO and H2
1750	Hydrogenation of higher alcohols. Mild hydrogen- ation catalyst.
2493	Dehydrogenation of alcohols - to aldehydes.
2730	Polymerization of Isobuty lenes to isooctylenes.
* 3076	Hydrogenation of diisobuty- 1 lene to iso-octane.
* \$390	Strong hydrogenation cata- 1 lyst, e.g. hydrogenation of phenol to cyclohexanol.
3510	Production of gasoline from - middle oil.
4577	Production of stearyl amine 8-9 kg/from stearic acid. Mo.

atalyst - Leun	a No. Use.	Production Tons/Mo.
* 4788	Hydrogenation of acety	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
	to ethylene	
* 4821	Polymerization of isobution to iso-octylene.	1tyl- 10
* 5058	Hydrogenation of Middle to gasoline (Gas phase	e Oil 25).
_	Heavy Water - Army Sec	ret -
5436;	Dehydrogenation and aromatization.	10
* 5623	Synthesis of toluene f benzene & methyl alcoh	rom 150 ol
5655	Hydroforming and dehyd genation	r o-
5780	Dehydramation of alc Basic carrier for othe catalysts	ohols. 100
6067	Amine synthesis	very lit
_6069	Catalyst for the react of methane and ammonia form methyl amine	ion 10
* 6448	Dehydrogenation of n-b	outane 100
6523	Conversion of phenol to	to -
* 6545 -	Heavy Water - Army Sec	eret -
* 6853	Arobin catalyst. Used version of heavy aromalighter aromatics.	for con- atics to
* 7187	Catalytic-cracking ca	
10927 26 —	Tar & Sump-phase hydr Ammonia catalyst	ogenation -

Samples of the catalysts marked with an asterisk (*) were secured for further study. In addition, samples were secured of the following catalysts not mentioned elsewhere:

Hydrogenation catalyst No. 6434 - 200 liters.
Hydrogenation catalyst No. 8376 - 800 liters.
Water Gas Shift Catalyst - 25 liters.

Synol Process.

Additional information was obtained on the operation of this process. No satisfactory pilot-plant runs had ever been completed, due to interruptions by bombings. Previously obtained flow schemes and product distributions were based solely on small-scale bench models, extrapolated to pilot-plant scale.

A 100-liter drum of the Synol catalyst was obtained.

OXO Process. Previously described flow schemes for this process had been attempted but proved unsatisfactory. A modified scheme as the plant was finally built and typical run data on a cracked-oil fraction were obtained.

A loo-liter drum of the OXO catalyst was obtained.

Other Samples.

The following hydrocarbon samples were secured at Leuna from processes previously discussed.

Aviation Alkylate

Heavy Alkylate Feed to Arobin Process (HF residue from Moosbierbaum) "TTH" (Tief-Temperatur Hydrierung) process, feed stock & products. Crude Isobutanol product

I.G. Farbenindustrie A.G., Schkopau, Target 30/4.02.

One liter of Kybol (all that was available), 3 onepound samples of aluminum-chloride used in manufacturing lubricating oils SS 903 and SS 906, three drums (each 200 ltr) of SS 903, and three drums of SS 906 (each 200 ltr), were obtained. One small drum each of new and regenerated acetylene hydrogenation used in ethylene manufacture was also secured.

Inquiry was made to harmonize a discrepancy that was found in the statements made on an earlier visit concerning the production of Zahlenbuna (Buna 85). The capacity of the polymerizer with screw conveyor is 75-100 tens per month, as was stated. In the plant production summary, the total polymer sold as Zahlenbuna is likewise correct; viz., 270 tons per month. The 170-tons are Buna 32.

Completion of data on an earlier flow-sheet for the hydrogenation of acetylene to ethylene was effected.

CIOS Target No. 30/4.07, Braunkohle Benzin A.G. (Brabag IV)

Hydrogenation Plant, Troglitz near Zeitz.

The plant was visited to obtain samples of hydrogenation catalyst No. 5058 and also samples of lubricating oils previously manufactured from the brown-coal tar hydrogenation-separator product. The lube oil samples include one spindle oil obtained by distillation of the dewaxed separator residue and also a neutral oil. Raw oils were made during February and March of 1944. Lube oil manufacture is not currently practised because of bomb damage to the dewaxing and distillation plants.

It was found that 2 hydrogenation stalls with a total of 5 reactor chambers were in operation. The process used is of the low-temperature type (350-400°C) at 300 atm. and brown coal tar is charged.

Target of Opportunity CIOS Group 30, WIFO Blending Station, Stassfurt.

Several members of the team made the trip to Stassfurt, but the presence of the Russians prevented their making any investigation.

Target of Opportunity for CIOS Group 30, Wirtschaftliche Forschungsgesellschaft m.b.H. (WIFO), Heiligenstadt.

Since the MAIN DEPOT Wife at Stassfurt, known to include a special laboratory for Rust Preventive Research, and in the underground storage, miscellaneous fuels and lubricants, as well as the records of the Central Technical Division, Berlin, could not be investigated, the smaller ARMY COMMAND DEPOT at Heiligenstadt was investigated. Information along the following lines was obtained:

- 1. Fuel and Lubricant specifications on the following materials handled by the Wifo were obtained:

 - #) Army Delivery Motor Fuel
 b) Benzol for Motor Fuel
 c) Automotive Lubricants (Summer & Winter grades)
 - d) Gear Oils (Summer and Winter grades)
- for tests were obtained in quantities sufficient for large-scale tests. These samples include:

Motoremoil (Summer) Nachlaufschmierstoff T 42 (rust preventive oil) Nachtaulschmierstoll T 42 (rust preventive oil All-Purpose Grease Gear Oil Methanol Glysantin (a glycerol substitute)

3. General operational information regarding the

- Wife system and the present location of key personnel and offices was obtained.
- I.G.Farbenindustrie A.G., Leverkusen CIOS Target of Opportunity, Group 30.

Recoil Fluid.

The recoil fluid used regularly by the German Army (triglycol, ethylene glycol and water) was found unsuitable in the winter campaign in Russia. The addition of low molecular weight sulfonamides eliminated this difficulty since, in this way, it was possible to obtain a recoil fluid with not too high viscosity at low temperatures and still maintain the high specific gravity reguired for satisfactory brake action.

The sulfonamides manufactured at Leverkusen for the above purpose were the monomethylamide of methanesulfonic acid and the methylnydroxyethylamide of methane--1-bb - Lar sulfonic acid.

Additive for Break-in Oil.

Product 891, dichlorodiphenylphosphorous acid, has been manufactured at Leverkusen as an additive for break-in oil for aircraft engines. To increase the solubility in oil,

the compound is used in the form of its stearylamine salt; and to further increase ease of handling, the additive was distributed as a solution in alcohol-benzol (50:50). The solution which contains 50% by weight of the stearylamine salt of Product 891 was termed J7, and samples were secured for further study. The break-in oil for aircraft engines consisted of Rotring oil (Red Band) with 2% by weight of J7. By use of the additive, it was possible togreduce the break-in time of German aircraft engines from 70 hours to 20 hours and to eliminate rejects.

Synthetic Lubricating Oil from Tetrahydrofuran.

Synthetic oils of good viscosity index, but poor thermal stability have been prepared at Leverkusen by copolymerization of tetrahydrofuran and ethylene oxide in the presence of FeCl3 and SOCl2. Engine tests with an oil of a viscosity of about 83 sis at 210°F resulted in ring sticking after short running time, and indicated that the piston was not properly lubricated, probably because of decomposition of the lubricant. The synthetic oil is not miscible with petroleum oil. The product has been used as gear oil and for the lubrication of machinery operating below 280°C. Production of the synthetic oil was irregular and on a small scale, not exceeding 1-2 tons per month.

Samples of this synthetic oil (called M-620) were secured for further study.

CIOS Target No. 30/5.01, Ruhrchemie A.G., Sterkrade-Holten. Synthetic Lubricating Oil Manufacture.

Automotive lube oil of a viscosity of 6-80E at 50°C was manufactured at the Ruhrchemie plant by polymerizing olefins in the gasoline-boiling range, using ACL; as catalyst. The olefins were obtained by separately cracking Fischer-Tropsch gas oil and "Sweat oil" from the manufacture of wax from Fischer-Tropsch material. Cracking was carried out in a Dubbs unit with steam injection between furnace outlet and reaction chamber inlet. The plant had a design capacity of 1000 tons of oil/month, but due to good operational practices, produced 1500 tons/month.

Development work on the manufacture of aviation-grade brightstock led to the design of a plant, the construction of which was begun in 1943. Due to the air attacks, the

equipment was moved to Willingen near Kassel, where the plant was to be erected underground. However, this was not accomplished. The brightstock was to have a viscosity of 38°E at 50°C and should have a higher V.I and better oxidation resistance than the automotive lube oil. This was to be accomplished by polymerizing only olefins in the range to be accomplished by polymerizing only olefins in the range to Co to Co and incorporating an inhibitor (phenthizin) into the olefin feed stock. The plant was designed for a capacity of about 1000 tons of finished oil per month.

Manufacture of Toluene.

Ruhrchemie developed a process to convert n-heptane to toluene by using a Cr203-Al203 catalyst. The reaction takes place at 480-5300C, and no external heating is required for the reactor, since the heat of reaction from the regeneration step is sufficient for the aromatization step. The cycle time planned for the full-scale plant was 60 minutes. Regeneration of the catalyst comprises two steps, viz., 1) burning off the carbon formed during the reaction, 2) reduction of six-valent or to three-valent or by means of hydrogen liberated during the conversion process. Catalyst life is said to be at least 2 years.

The yield of liquid product amounts to 90-92% by wt. of the feed and the toluene content of the liquid product is 50% by wt.

Development work on the process was finished in October 1944 and a plant for the production of 24,000 tons of toluene per year was designed. Plant construction was abandoned, however, on order of the German government when the plant was about 20% complete.

Miscellaneous.

Supplementary data on the oxidation of high melting wax to high mol. wt. acids (H2SO4 and gas stream from the NH3 oxidation unit) was obtained.

A description of the laboratory method for determining the activity of catalysts was obtained.

Production of high molecular weight alcohols from wax was found not be be a direct operation, but rather only a laboratory reduction by usual methods of the acids made from high-melting wax

A description and flow sheet of the methanization of coke-oven gas was obtained.

In order to preserve for a later team samples that were still in the plant, arrangements were made with Prof. Pritz Martin to have the samples collected and packaged. For identification, the packages were to be marked "Trip 551". Samples requested were:

Sample.

1 200-liter drum.

Catalytic Cracking Catalyst (Granosil, an activated clay obtained from Bleichton-Gesellschaft, m.b.H., Roman Mayr-Haus, Munchen 2)

Fe-Cu on kieselguhr catalyst for all available up to 1 200-L. drum.

operation at 220°C. Ni catalyst for methanization all available up to of coke-oven gas. all available up to 1 200-1. drum. of coke-oven gas.

Cr203 - Al203 catalyst for Up to 1 200-liter cyclizing nC7 etc.

Cobalt Catalyst.

3 tons.

Feinreinigungsmasse

1 200-liter drum.

No samples of products from Co catalyst operation were available at the plant or in any of the Ausweichslager.

I.G. Farbenindustrie, Hechst - CIOS 22/lg. Target of Opportunity for Group 30.

- I.G. Farbenindustrie, Hochst, was visited by a detachment from CIOS party 551 on 9 July to obtain further information on synthetic additives used in cutting and metaldrawing oils. In addition to development and manufacturing information already available from previous visits on Hochst compounds of this type, the following additional information and samples were obtained:
- 1) The Metal-Drawing Oil, Saure E, is MEPASIN sulfamido acetic acid, which is obtained by acetone extraction of the unreacted MEPASIN from Bohrmittel H#. A 1-1.5% water

suspension of the acid(pH=4) is used as the drawing agent.

This material has been used with partial success on non-bonderized metals and was being considered as a substitute for the soap solution previously used in the final drawing process.

Plant-scale tests have been made on the drawing of bonderized metals, starting with the metal discs. Full-scale production tests were also made on the final draw of 3.7 cm. steel shells. The tests were only partly successful and no definite conclusion on the efficiency of the material can be drawn at present. The difficulties encountered when working with Saure E included the following points:

- a) Material treated with Sture E must be drawn immediately since the effective film tends to shrink, thereby leaving some metal areas uncovered;
- b) The solution of Saure E causes reddening and peeling of the skin.

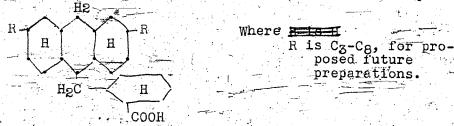
According to the limited experience gained, it seemed to be possible to overcome these difficulties by adjusting the pH of the working solution to values between 5 and 7. This was done by the addition of zincate solution (sufficient to neutralize one-half of the Saure E used) to a 0.5% solution of Saure E.

Better results without the above-mentioned disadvantic ages are expected from the use of isopropyleyclohexyl acid which can be used in form of its Na salt and in exactly the same way as the usual soap. No production scale tests on this material have been made, but laboratory tests indicate that it is about 5 times as efficient (calculated from the amount of chemical required for covering the metal surface) as the usual soap solution.

2) - The raw material for the emulsifying agents made at Hochst was KOGASIN II from Fischer-Tropsch synthesis. In order to be less dependent upon this source of material, a research program was in progress along the lines of condensing alkyl benzene or anthracene with phthalic anhydride in the presence of AlCl3, and then hydrogenating this type of compound. The chemical structure of the compound from ALKYL benzene was of the type -

$$R - \left\langle \frac{H}{H} \right\rangle - \frac{C}{H_2} - \left\langle \frac{H}{COOH} \right\rangle$$
 where R is C_3 - C_8

The compound from anthracene had the formula:



Laboratory-scale work on the preparation of these compounds was under way and no conclusions as to the properties of these compounds as additives was available.

3) - A drum sample of Bohrmittel Hihas been secured for test, either as a cutting oil, as a gasoline additive, or in the extracted-acid form as a metal-drawing oil.

CIOS Target 30/4.03 -- I.G. Farbenindustrie, Ludwigshafen, (Heidelberg).

An interrogation of Pier and his staff was made at Heidelberg for the purpose of obtaining additional information on the subject of hydrogenation catalyst composition and preparation. A complete list of these catalysts and details of their preparation were obtained. This is summarized below.

War research in the hydrogenation field had consisted largely in development of the Arobin Process, DHD Process, double DHD Process for toluene production, and a process to obtain higher isoparaffin content fuels from bituminous coal middle oil by a prehydrogenation over catalyst 7360 to remove oxygen compounds and reduce the nitrogen compounds to amines which were then washed out with dilute sulfuric acid, followed by a hydrogenation cracking over hydrogenfluoridetreated clay.

Catalyst No.

Use.

Aromatization and hydrogenation.

5058 Prehydrogenation and TTH.

6434 Hydrogenation of middle oil.

8376 (-7846_{w250}) General hydrogenation.

	Catalyst No.	<u>Use</u> .
	7846	Prehydrogenation catalyst
		(replaced by 8376)
	5615 (=6718)	Dehydrogenation of gasoline,
		hydrogenation of di-isobutylene,
		reduction of higher alcohols.
	7019	Aromætization.
	7360	Denyerogenation.
	7965 s	Dehydrogenation.
		Methane-splitting catalyst.
		Water-gas shift catalyst.
		Grude-molybdenum catalyst
		(Liq.phase hydrogenation).
	10927	Grude-Iron catalyst.
		(Lig.phase hydrogenation).
	6448	Butane-Dehydrogenation.
,	6752	Catalytic Cracking.

Samples and Documents.

A total of approximately ten tons of samples of oil and catalyst were secured on this trip. These samples were delivered to the "Toot Sweet" railhead at Mainz, packed and labelled for shipment to the U.K. and U.S. as shown in the attached list. A detailed list of samples is attached. The further transport of these samples from Mainz is being arranged by the Ordnance Department through Captain Chaffee.

There are also attached lists of documents secured from I.G. at Leuna and from Ruhrchemie at Sterkrade. These latter documents were secured at Nuttlar where they had been stored.

DETAILED SAMPLE-LIST CIOS Trips 551 and 551a.

As a result of CIOS Trips 551 and 551a, the following samples were collected for distribution to the indicated agencies:

Number. Description. Qu	antity.	<u>To</u> 🎝
1. Aviation Alkylate ET 120 (line)	100 L	TIIC
2. Aviation Alkylate ET 120 (storage)	100 I	TIIC
3. Arobin Feed (HF bottoms)	200 L	TIIC
4. Light ends from SS 906, Leuna	200 L	TIIC
5. SS 906 Leuna	200 L	TIIC
6. SS 903 Schkopau	200 L	Ordnance
7. Y Axle Oil (dved red)	200 L	
7. Y Axle Oil (dyed red) 8. Ester Oil - 504 (Adipate)	200 L	Ord.TIIC Ord.TIIC
9. Iso-octane Hydro.Cat (used) 3076?	100°L	
10. Acetylene Hydro Cat - S (regn)	*24 Kg.	TIIC \\TIIC
11. Arobin Cat (used 3 Mo) 2 full 685	~ & ± .Ng•	TITO
12. Acetylene Hydro.Cat. S, New	24 Kg.	TIIC
13. SS 903 Schkopau	-9A + A-6.	
14. SS 903 Schkopau	200 L	Br.
15. SS 906 Schkopau	200 L	TIIC
16. SS 906 Schkopau	200 L	Br
17. Isobutanol Cat. No.1132 ?	-200 L	TIIC
18. Isobutanol Cat. No.1182 7	100 L	TIIC
	100 L	TIIC
	200 L	TIIC
The state of the s	100 L	TIIC
	l L b	
	l box	
	100 L	TIIC
	100 L	TIIC
25. Ester Oil No. 504	200 L	\mathtt{Br} .
26. Y Axle oil	200 L	\mathtt{Br}_{ullet}
27.— Spindle oil - Zeitz (Box)	2 L	TIIC
28. Lube 011 Zeitz	2 L	TIIC
29. SS 906 Schkopau	200 L	Ord.
30. Hydro Cat No. 5058, Zeitz	100 L	TIIC
31.	100 L	TIIC
	100 J	TIIC
33.	100/1	TIIC
34. Isobutylene Poly.Cat.4821 Schkolen		TIIC
35.	100 L	TIIC
36. Cat.Cracking Cat. No.7187 Schkolen		TIIC
37.	200 L	TIIC
38. Heavy alkylate - Leuna	2 L-	TIIC
39. Hydro Cat. No. 8376	200 L	TIIC
40. Hydro Cat. No. 8376	200 L	TIIC

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Number. <u>Description</u>.
                                                Quantity.
                                                                                TIIC
  41. Hydro Cat. No. 8376
                                                                  200 L
   ## 6434 ## 200 L

44. DHD catalyst No. 5931 ## 200 L

45. ## ## 7935 ## 200 L

46. Distillation feed TTH (Leuna) I ## 200 L

47. Charge to TTH (Leuna) I ## 200 L

48. Gasoline feed TTH (Leuna) I ## 200 L
                                                                200 L
                                                                                   TIIC.
                                                                                   TIIC
                                                                                 TIIC
                                                                                 TILC
  46. Distillation feed TTH (Leuna) I 200 L TIIC
47. Charge to TTH (Leuna) II 200 L TIIC
48. Gasoline from TTH (Leuna) III 200 L TIIC
49. Diesel Oil TTH (Leuna) IV 200 L TIIC
50. Crude Isobutanol " 200 L TIIC
51. Light Syn.Lub.Oil 120°C Fl. 200 L Br.
-52. Ident Syn.Lub.Oil 170°C Fl. 200 L Ord.
53. KSE Blending Agent (packed wi 56 & 58) 1 L (box) Br. 54. " " ( " 55 & 57) 1 L (box) Ord. -
   except where noted. The following samples were
        secured from the WIFO Blending Station at
    Heiligenstadt:
         Fl. Nachlaufs#chmierstoff T42 50 L
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ا میں اسے رائی میں				
Number.	Description.	Quan	tity.	To.
82. Fl. 83. 84. 85. Gear 86. " 87. Summ 88. " 89. Glys 90. " 91. Abso 92. Meth 93. " 94. Glys 95. " 96. Abso 97. Meth 98. "	Nachlaufschmierstof """ Oil, can ner Motor Oil (10 1 santin chmierfett TL 6014B hanol Anti-freeze santin chmierfett TL 6014B hanol Anti-freeze santin chmierfett TL 6014B	f T42 50 " 50 " 50 " 50 Lit.cartons) " " 2 in box 1 with 87 1 & 103 1 in box 1 with 87 1 a 103 1 lin box 1 l	L 0 L B L B L Gal. Gal. Gal. Gal. Gal. Qt. Qt. Qt. Qt. Qt. Qt. Qt. Qt. Qt. Qt	rdn. pit. Br. Br. d-TIIC d-TIIC d-TIIC d-TIIC d-TIIC d-TIIC d-TIIC d-TIIC d-TIIC Br. Br. Br. Br.
Unde	erlined destination walso interested.	indicates lab	eling - ot	her
62,	70, 72 and 73 are a	ll packed in	one box.	9 <u>, </u>
Samples from				
99. Prod 100. " 101. Prod 102. " 103. *"	luct M-620 (Packed " (Packed	with 99) with 100) with 87)	2 Kg. (Gla 2 Kg. (Gla 1 lb. 2 lb. 2 lb.	

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	14 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1		.)	
	1.	No.	1722	Beeinflussung der Klopfgrenzkurve von aromat-
	¥.		/-	enhaltigen Kraftstoffen durch verschiedene Ole.
			893/2	Das Siedenverhalten von Flugmotorenkraftstoffen
	3.	No.		Werhalten von Kraftstoffen in SG-Behältern.
			893/3	Ruckstandsbildung von bleihaltigen Kraftstoffen
	5.	No.	690-	Beschreibung der russischen Flugmotoren "AM 35" und "AM 38".
-	6.	No.	1150	Klopfmessung mit dem DVL-Verfahren der Bruck- beschleunigung.
	7	Mo	~1382	Prufung von Laboratoriumsverfahren zur Bestim-
•	· 🔍	_NO.	TOOS	mung des Bleigehaltes in Kraftstoffen.
•••	8.	No	1327	Uber Laboratoriums- und Motorverfahren zur Pru-
		· 140 • 1	TODI	fung der Klopffestigkeit von Otto-Kraftstoffen.
-	ä 📉	NO F	523/7	Einfluss der Triebwerksgestaltung und der Be-
	÷,	110.0		triebsbedingungen auf das Klopfverhalten von
				Kraftstoffen. 7. Teilbericht.
``	10.	NO.	688	Einfluss des Bleigehalts von Kraftstoffen auf
	T (1)			Bauteile von Flugmorotren (2 copies).
	11.	No.]	.053	Regelung von Zustandsgrössen, insbesondere von
,				Gastemperaturen am Ende Längerer Rohrleitungen
-	200	4.5	. 1.	1. Teilbericht.
٠.	12.	No.	695	Messung der Kolbentemperatur am laufenden Motor.
	13.	No. F	523/5	Einfluss der Triebswerksgestaltung und der Be-
	10.	11000		triebsbedingungen auf das Klopfverhalten von
			٠,	Kraftstoffen. 爱. Teilbericht: Versuche am BMV
				VI-Einzylinder-Motor, Reihe 9 mit Vergaser-
				betrieb.
	14.	No.	L679	Der Einfluss der Betriebsbedingungen auf die
٠.	7/2	= -	•	Kolbentemperatur.
	15.	No.	523/2	Einfluss der Triebwerksgestaltung und der Be-
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	1			zylinder-Prufstand mit Kraftstoffeinspritzung.
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	•	Te	echnis	che Hochschule, Dresden, 10 Januar, 1945.
	18.	Ber	icht u	ber die Untersuchung von 9 definierten Ölen
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Dr.H. Zorn, Heidebroek, Technische Hochschule, Dresden,

10 Februar, 1945.

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- 1. Efficiency and Economics of Gas-Driven Vehicles. HEFT 3.
- 2. (i) Investigations on Automotive Diesels \((ii) \) Fresh Oil Lubrication of Main Bearings \((iii) \) Investigation of Radiation in Combustion \(\) space of high speed Diesel engine.
- 3. Research om the combustion process in high speed HEFT 5.
- 4. Cylinder and Piston Ring wear. HEFT 29.

 5. Measurement of knock ratings in Otto engines HEFT 31.
- 6. Determination of knock noise in Otto engines by electroacoustic instruments. HEFT 33.
- 7. Mechanical losses in high speed Diesel engines HEFT 34.
- 8. Comparison of Bearing metals_ HEFT 52.
- 9. (i) Injection of fuel into Diesel engines
 (ii) Ignition delay measurements by photocells of
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- 10. Determination of Lubricating Oil film strength HEFT 54.
- 11. Effect of fuel and engine on the starting of Diesel engines HEFT 55.
- 12. Mixture formation in the injection nozzle HEFT 57.
- 13. Lubrication of glass in the boundary region HEFT 59.
- 14. Operation of 2 stroke engines with liquefied gas HEFT 60.
- 15. Scavenger action explained on the basis of a new thoery of expansion flow
- 16. Investigation of the suction stroke in Diesel engines HEFT 62.

- 17. (i) Ignition delay and rating of fuels
 - (ii) Ignition delay in Diesel & Otto fuels HEFT 63.
- 18. The effect of air flow on fuel distribution in Diesel engines HEFT 76
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- 19. Review of theory of mixture formation in Otto and Diesel engines.
- 20. Motor ratings for Diesel fuels.
- 21. Knocking in multi-cylinder engines.
- 22. Motor Testing of synthetic Otto fuels.
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- 27. Experiments on the use of liquefied gas for Diesel operation.
- 28. Experiments on a carburettor engine with auto ignition
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 - C. Miscellaneous.
- 31. Instructions for the operation of a cathode ray oscillograph.
- 32. Investigations on the development of operating a mixture-fed engine by auto ignition (TH Stuttgart) (FKFS392)
- _33. Cold starting tests with Wehrmacht All Purpose 0.11 (Adam Opel).
- 34. Thesis Determination of pressure characteristics of lube oils.

- 35. Thermal stability of gear oils (T.H.Stuttgart).
- 36. Knock testing of aviation gasolines in single cylinder engines (I.G. Uppau) Report 474.
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- 38. Behavior of aviation fuels at high altitudes.
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- 40. Fuel-rating with respect to vapor lock.
- 41. Minutes of meeting on knock testing in I.G. and CFR engines etc.

42. Memorandum on fuel problems for aviation.

- 43. History and uses of IG engine "k" (Diesel).
- 44. Determination of cetane numbers of Diesel fuels by method of Dr. Neumann.
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- 46. Comparison of engines for determination of cetane numbers.
- 47. The IG Test_Diesel.

 48. Minutes of meeting on standardisation of Diesel Fuel Testing.
- 49. Standardisation of engine testing of Diesel fuels.
- 50. Sludge formation in aviation lubes.
- 51. Lubricating oil testing in single cylinder JUMO 205.
- 52. Intava Report attempted evaluation of oil tests in the BMW engine.
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- 54. Gear tests at oil temperature of 150°C (T.H. stuttgart)
- 55. Thermal stability of gear oil Wehrmacht 8E.
- 56. Oil Testing with the Almen machine
- 57. 10 drawings from Oppau report 478 Wear Testing Mach.
- 58. 3 11 11 11 542 11
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- 60. Testing of synthetic oil in Humboldt-Dentz engine.
- _61. Report on lubricating qualtities of oils.
- 62. 2 Vacuum reports on Hypoid gear oils.
 - 63. Testing of gear oils.
- 64. ditto.
 - 65. Test_for the cold properties of lubricating oils.
- 66. Determination of minimum flow temperature of lubricating oils.
- 67. Cranking tests with new Wehrmacht winter oils.
 - 68. Development of low temperature viscosimeter.
 - 69. Procedure for cranking tests at low temperatures.
 - 70. Pumping of low pour lubes in engines.
- 71. Determination of pumpability of bunker oils.
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- 76. Pumpability of gear oils at low temperatures.
- 77. Cooperation tests on vapor lock.

- 78. Report on incomplete gear oil tests.
 - 79. Investigation of increase of boiling range of carburettor fuels.
- 80. Addition of various EP agents to synthetic oil.
 - 81. Tests on effect of Bright Stock on blends containing synthetic residues and distillates.
 - 82. Oxidation resistance of aviation oil blend K2025.
- 83. Evaluation of lubricants by engine wear.
 - 84. Thermal stability tests of gear oils.
- 85. Starting of Otto engines at low temperatures. 86. Tests on aviation oil K2015.
- 87. Foaming of lubricating oils.
- 88. Ring sticking tests on SS 1060.
- 89. Cracking tests on Roumanian oil.
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- 100. Effect of Oppanol on engine wear.
- 101. Effect of Oppanol on behavior of engine oil.
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- 105. Effect of viscosity on oil consumption.
- 106. Testing of motor oil 3370.
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- 110. Aviation oil K1951.
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- 116. Ignition varue of RCH cetane.
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- 120. Engine testing of aviation oil 1929.
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- 129. Development of engine test for aviation oils in
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- 135. Effect of engine condition in testing lubricating
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- 137. Tests with mixtures of Gas 81 and gasoline.
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- 140. Test to classify discrepancies in octane number determination of Ruhr Benzine Fuels.
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