RESTRICTEL X-38(N)-7

ENCLOSURE (B) 8

EFFECT OF REACTION TIME

NAVAL ENGINEER T. OGAWA

Research Period: 1929 - 1930

Prepared for and Reviewed with Author by the U.S. Mayal Technical Mission to Jepan.

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SUMMARY

Studies were made on the effect of varying reaction time on the hydrogenation of Oyama coal, holding reaction conditions constant at 45500; hydrogen pressure at 100 atmospheres and using Fe₂O₃ as a catalyst. It was reaction time that under these conditions, the optimum reaction time was l hour.

INTRODUCTION

in the hydrogenation of coal, reaction time is an important factor, particularly from the standpoint of commercial plant design. Although many reports had previously been made on this subject, accurate conclusions could not be drawn, since the experiments had not been made under comparable conditions. This study was limited to the investigation of the effect of reaction time only, holding other variables constant.

Key research personnel working on project were Naval Eng. T. OGAWA and Naval assist. Eng. I. TAKAHASHI.

II. DETAILED DESCRIPTION

The autoclave and test procedures were the same as described in report No. 3 of this series.

Samples were as follows, (same as in report No. 3):

Coal : Fushum Oyama coal, sized to under 20 mesh.

: 30% topped light oil fraction from Shinbara low-temper-Tar Catalyst : Ferric oxide (commendation type retort.

: Ferric oxide (commercial grade)
: Prepared by electrolysis, purity above 99.5%.

These components were mixed in the following amounts: coal, 100 grams; tar, 40 grams; Fe203, 5 grams; and H2, 19 grams.

Runs were made for reaction times of 0 minutes, 30 minutes, 1 hour, 3 hours, and 5 hours and the other reaction conditions were held as constant as possible. Results of the several runs are summarized in Table I(B)8.

III. CONCLUSIONS

From the preceding experiments, it was noted that a part of the coal was hydrogenation was reliable period of heating to reaction temperature, and the hydrogenation was reliable period of heating to reaction temperature, and the hydrogenation was complete and the yield of liquid products reached a maximum. With increasing reaction time the yield of liquid decreased and the formation of gas increased due to secondary thermal cracking.

ENCLOSURE (B)8

Table I(B)8

	BFFECT OF REAC	TION TIME Run Number					
The first of the second of the			50	76	_43	44	
Reaction Conditions	React. Time (hr-min) React. Temp. (°C) Initial Pressure (atm.) Pressure Drop (atm.) Preheating Time (hr-min)	0-0 455 95 8 1-30	0-30 455 100 21 1-30	1-0 455 101 19 1-55	3-0. 455 99- 23 1-25	5-0 455 • 98 23 1-25	
Reaction Products (gm)	Gas Water 011 Residue	29.3 13.7 64.0 52.8	33.5- 17.8 78.3 29.1	33.6 17:4 85.9 24.9	33.8 17.4 79.2 22.1	40.5 17.5 74.1 25.0	
Cas Analyses (gm)	CO ₂ C _D H ₂ n OO H2 C _D H ₂ n+2	2.2 0.5 0.8 15.8 10.0	1.3 0.5 1.3 13.5 16.9	0.7 0.5 1.6 14.3 16.5	0.6 0.3 1.3 12.8 18.8	1.6 0.4 2.3 23.5 12.7 1.8	
Distillation of Oil (gm)	180°C 180230°C 230360°C Pitch	3.7 60.3	12.3. 5.8 60.2	10.6 5.8 36.0 33.3	11.2 6.4 34.0 27.0	12.0 7.2 32.7 22.1	
Analyses of Residue (gn)	Benzene Soluble Benzene Insoluble Ash	40.1 12.7	15.2 13.9	0.2 12.0 12.9	9.5 12.6	0.1 12.4 12.6	