STUDIES ON SYNTHESIS
OF HYDROGEN PERCYIDE FROM
HYDROGEN-OXYGEN MIXTURE BY
ELECTRIC ARC DISCHARGE METHOD

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### SUMMARY

when the flow rate of the hydrogen-oxygen mixture (02 about 5%) through the electric are exceeded 100 m/sec, the power consumed by the hydrogen peroxide synthesis was about 300 kwh/kg H2O2.

To obtain the synthesized hydrogen peroxide, the following three methods were used. Among them, Method 1 was the best.

- 1. Distilled water at the end of the reaction tube was used to absorb the  $H_2O_2$ .
- 2. Distilled water was allowed to contact the reacted gas directly and absorb the  $\rm H_{2}O_{2}$ .
- 3. Synthesized H<sub>2</sub>O<sub>2</sub> was condensed directly by cooling.

#### i. INTRODUCTION

The synthesis of hydrogen peroxide from hydrogen-oxygen mixtures by electric discharge previously had been effected with an ozonizer tube. However, the treated volume per unit in this method, was so small that to improve this point, are discharge was tried by Lieutenant T. KONOSU from September to December 1944. When the are discharge was used, the flow speed in the reaction tube was most important, and it was found necessary that the flow speed be above 100 m/sec. To obtain the synthesized products, distilled water was injected at the end of the reaction tube, and the products absorbed.

#### II. DESCRIPTION

#### A. Apparatus

The apparatus is shown in Figure 1(B)4

#### B. Procedure

In Figure 1(B)4, the mixed hydrogen oxygen gases are stored in the gas tank (T) and the  $O_2$  content is maintained at about  $\mathcal{H}$ . By means of the gas recycling pump (P), the gases are forced into the reaction tube (R). Their volume is measured by the flow meter  $(F_1)$  and their pressure by the manometer  $(P_1)$ . In the reaction tube they contact the arc and the cooling water, which is recycled by the recycling pump (I), and flow into the cooling tube (C). Then the cooling water flows into the receiver (E), and residual gases are recycled into the gas tank (T) by the gas recycling pump (P) after their volume is measured by the flow meter  $(F_2)$ , and their pressure by the manometer  $(P_2)$ . When the experiment is over, the product is removed and titrated with 1/10 N KMnO4 to determine the concentration of  $H_2O_2$ .

#### C. Results

The results are shown in Table I(B)4

The apparatus in which the cooling water was used so as to contact directly with reacted gases, is shown in Figure 2(B)4, and its results are shown in Table II(B)4.

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RESTRICTED

ENCLOSURE (B)4

The apparatus in which the synthesized H<sub>2</sub>O<sub>2</sub> was condensed directly by cooling is shown in Figure 3(B)4, and its results are shown in Table III(B)4.

# II. CONCLUSIONS

When the cooling method was used as shown in Figure 1(B)4, the power consumption was about 300 kwh/kg H<sub>2</sub>O<sub>2</sub>, and the H<sub>2</sub>O<sub>2</sub> concentration was about 0.5%. This concentration may be increased by reducing the volume of initial cooling water. In this case, however, the materials of the cooling water recycling pump, which is made of iron and copper, are not suitable. Hence, it will be necessary to select a material that will minimize the decomposition of H<sub>2</sub>O<sub>2</sub> synthesized.

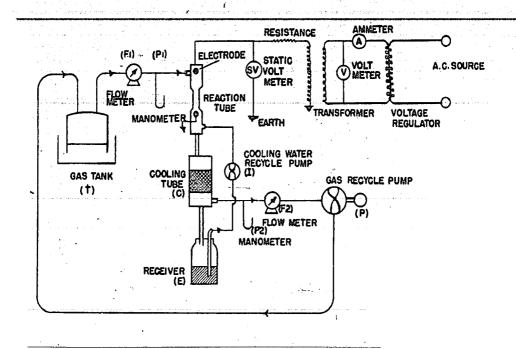
It was the object of this research to develop a commercial method of producing H<sub>2</sub>O<sub>2</sub> which would eliminate the use of platinum and which could be used as a substitute for the (NH<sub>2</sub>)<sub>2</sub>S<sub>2</sub>O<sub>3</sub> method. In this experiment it was hoped that the power consumption would be below 100 kwh/kg H<sub>2</sub>O<sub>2</sub> and that the concentration of H<sub>2</sub>O<sub>2</sub> produced be at least one percent. Additional experiments will be necessary to develop this method commercially.

ENCLOSURE (B)4

H202 Concent-ration (%) 0.00 0.00 0.01 6.3 7.0 5.0 7.0 Power Consumption (KHH/kgH2O<sub>2</sub>) 1,000 2,800 2,000 8 350 38 33 Total H202 (FH/hr) 0.19 0.26 0.48 1.42 1.56 1.98 1.86 GAS ×12 Power (watts) 8 97 25 QQ 7 27 528 8 8 Table I(B)4. RESULTS OF ARC DISCHARGE ON MIXED H2, Power Factor 9.0 9.0 9.0 9.0 9.0 9.0 9.0 Current (milli-amps) 35 45 7 ß ያ R 45 Secondary Voltage (kilo – volts) ä 15.6 0.11 10.6 22.0 10.5 13.3 Current (amps) ଷ ଯ 13 8 ଷ 8 ଷ Primary Voltage (volts) 45 3 42 # 45 S ß Pressure (P<sub>1</sub>) (m Hg abs) **2**‡0 2,0 8 8 8 8 8 Gas volumes (F<sub>1</sub>) (14t/min) 8 8 335 7 8 R 22

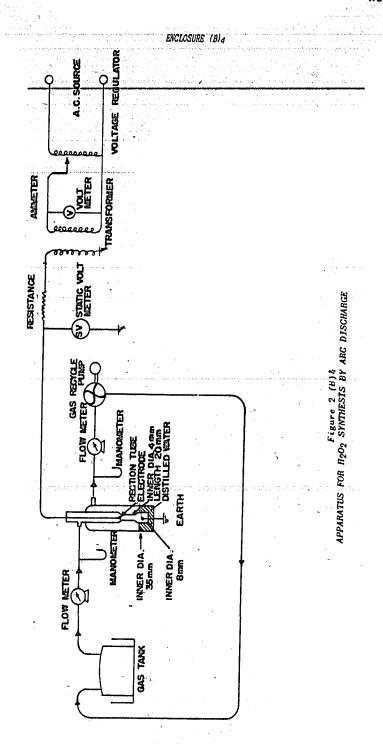
ENCLOSURE (B)4

	1	Total   Power   Hy02   Power   Hy02   Consumption   Fation   Fation   Fation   Fation   Fation   Fation   Fation   (3)		8 0 007				Consumption concent- (NWH/kgH2O <sub>2</sub> ) ration				550 1.0
Table II(B)4 RESULTS OF ARC DISCHARGE ON LIXED H2, 02 GAS				1.37				Total consu	(TIM ATT)			0.1
	.	Power (watts)		552	576	O <sub>2</sub> GAS		Power (watts)	2,50	3	7#X	2
		Power		9.0	9.0	, II (B)4 MIXED H2,		Power factor	9.0	90	2 0	;
	Secondary	Current (millid- amps)	50	S	83	Table III(B)4,	ary	Current (milli- amps)	82	Ç.	\$ 8	
		Voltege (kilo- volts)	10.8	11.0	11.5	Table III(B)4 TESULTS OF ARC DISCHARGE ON MIXED H2,	Secondary	Voltage (kdlo- volts)	10.8	10.8	11.5	
	Primary	Current (amps)	8	ล	20	RESULTS C	£.	Current (amps)	8	श्च	8	
		Voltage (volts)	4.5	947	877		Primary	Voltage (volts)	45	45	877	
		Pressure (mm Hg abs)		07/2	740			Fressure (mm Hg abs)	07/2	740	07/2	
		Gas volumes (lit/min)		100	027			Gas volumes (1/min)	75	100	120	



# DETAIL OF REACTION TUBE (QUARTZ TUBE) Rmm (OUTER DIA.IOMM 4mm (OUTER DIA.6mm)

Figure 1 (B)4
APPARATUS FOR H<sub>2</sub>O<sub>2</sub> SYNTHESIS BY ARC DISCHARGE



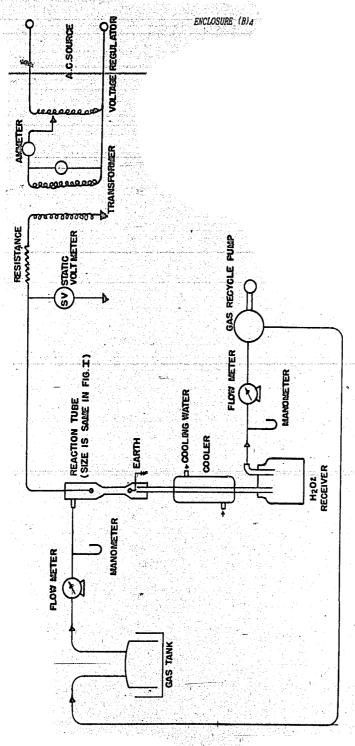


Figure 3 (B)4
APPARATUS FOR H202 SYNTHESIS BY ARC DISCHARGE