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# **Chemistry of Carbonate-Sulfur Flux**

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Reactions of alkaline metal carbonates with sulfur are investigated in detail. The evolution of CO and a trace of  $SO_2$  were observed in the course of reaction with major component of polysulfides. Some evidences that the reaction proceeds with breaking of terminal sulfur-sulfur bond in the sulfur polymer, and forming CO,  $SO_2$  and polysulfide are presented. Polysulfides have the role of keeping free sulfur and allow it to react with other chemicals to rather high temperatures.

#### Introduction

The reaction of alkaline metal carbonate and sulfur has been known to produce alkaline metal polysulfides.

 $3Na_2CO_3 + 12S \rightarrow 2Na_2S_5 + Na_2S_2O_3 + 3CO_2$  (1)<sup>1</sup>

 $4K_2CO_3 + 16S \rightarrow 3K_2S_5 + K_2SO_4 + 4CO_2$  (2)<sup>2</sup>

$$3K_2CO_3 + 2(x+1)S \rightarrow 2K_2S_x + K_2S_2O_3 + 3CO_2$$
 (3)<sup>3</sup>

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(the reaction of Na<sub>2</sub>CO<sub>3</sub> with S begins at 275-280 ℃; that



Figure 1. Gas evolution for the reaction of  $Na_2CO_3$  and S. The composition of the reactants:  $Na_2CO_3$ , 0.03 mole; S, 0.1 mole.

of  $K_2CO_3$  with S begins at 108 °C and completes below 180 °C)

On the other hand, sodium thiosulfate disproportionates at temperatures above 223  $^{\circ}\mathrm{C}$  as

$$4Na_2S_2O_3 \rightarrow 3Na_2SO_4 + Na_2S_5 \tag{4}^4$$

and is formed readily at lower temperatures by the reaction

$$Na_2SO_3 + S \rightarrow Na_2S_2O_3 \tag{5}^5$$

Alkaline metal carbonate-sulfur flux was used for preparing sodium chromium sulfide (NaCrS<sub>2</sub>) from NaCrO<sub>2</sub> or Cr<sub>2</sub>  $O_3^{6}$ . Recently yttrium oxysulfide (Y<sub>2</sub>O<sub>2</sub>S) has been prepared in the same flux<sup>7</sup>.

In the present work, further details of reactions taking place in the alkaline metal carbonate-sulfur flux have been studied.

#### Experimentals

All of the chemicals used in this investigation were from commercial products of chemical pure grade without further purification.

The reaction mixture was loaded in a quartz tube and heated in an electric furnace. The gaseous products were collected above water in a Mariott bottle, to measure volume and to sample for IR spectral analysis.

The molten mass remaining in the quartz tube was washed out with water and analyzed for anions in it: Water soluble sulfides, sulfites, and thiosulfates of the reaction products were analyzed by iodometry, and sulfates by conductometry with barium chloride. Water insoluble solids were analyzed by XRD and IR.

#### **Results and Discussion**

**Reaction of Na<sub>2</sub>CO<sub>3</sub> with S.** Gas evolution is represented in Figure 1. IR spectra shown in Figure 2 reveal that the gaseous product is mainly  $CO_2$  and a small amount of COS, indicating production of CO gas in the course of



**Figure 2.** Infrared spectra of evolved gas for the reaction of: a)  $Na_2CO_3$  (0.09 mole) and S (0.22 mole), and gas is collected at temperatures from room temperature to 470 °C; b) CaCO\_3 (0.001 mole),  $Na_2CO_3$  (0.0015 mole) and S (0.01 mole), and gas is collected at temperatures from 450 °C to 650 °C; c) CaCO\_3 (0.04 mole) and  $Na_2S_2O_3$  (0.08 mole), and gas is collected at temperatures from 500 °C to 600 °C.

reaction. Since  $CO + CO_2$  mixture of total pressure 1 atm can contain less than 0.1% CO below 400 °C due to the Bourdourd reaction

$$2CO \rightarrow C + CO_2$$
 (6)

Since the reaction (6) is rather slow at lower temperatures, most of CO would be converted to  $C+CO_2$  and small amount of CO may react with sulfur to form COS. The volume of gas evolved corresponds to 0.8 mole per mole of sodium carbonate used. Therefore, it is plausible to assume the reaction proceedes as follows:

$$Na_2CO_3 + S_{(x+1)} \rightarrow Na_2S_x + SO_2 + CO$$
 (7)

followed by the reaction (7),

$$Na_2CO_3 + SO_2 \rightarrow Na_2SO_3 + CO_2 \tag{8}$$

$$Na_2SO_3 + S \rightarrow Na_2S_2O_3 \tag{5}$$

followed by the reaction (4).

Overall reaction can be described as

$$2Na_2CO_3 + 7S \rightarrow 1\frac{1}{4} Na_2S_5 + \frac{3}{4} Na_2SO_4 + \frac{1}{2} C + \frac{1}{2} CO_2(9)$$

which would predict that 1.5 mole of  $CO_2$  will be produced from 2 moles of  $Na_2CO_3$ , but part of CO from reaction (7) would be followed as:

$$CO+S\rightarrow COS$$
 (10)

Actual total gas volume was measured to 1.61 mole, corresponding to conversion of 20% of CO to COS.



**Figure 3.** Gas evolution for the reaction of CaCO<sub>3</sub>, Na<sub>2</sub>CO<sub>3</sub> and S. The composition of the reactants: CaCO<sub>3</sub>, 0.02 mole; Na<sub>2</sub>CO<sub>3</sub>, 0.03 mole; S, 0.1 mole.

Gas evolution starts at about 300  $^{\circ}$  c and is completed at around 400  $^{\circ}$  c.

**Reaction of CaCO<sub>3</sub> with Na<sub>2</sub>CO<sub>3</sub> and S.** The course of reaction for this mixture is shown in Figure 3, which indicates that CaCO<sub>3</sub> reacts with sulfur after completion of Na<sub>2</sub>CO<sub>3</sub>+S reaction, from about 450 °C. The IR spectra indicates the gaseous product is constituted mainly of CO<sub>2</sub> with a fair amount of COS and small amount of SO<sub>2</sub>. The volume of gas produced at each step corresponds to nearly 3/4 moles per mole of metal carbonate present; the first stage: 0.81 mole per mole of Na<sub>2</sub>CO<sub>3</sub>, the second stage: 0.89 mole per mole of CaCO<sub>3</sub>. Therefore, the reaction mechanism for CaCO<sub>3</sub> +S reaction appears to be the same as for Na<sub>2</sub>CO<sub>3</sub>. The differences of this reaction with the previous sodium carbonate reaction are that CO is produced at higher temperature, so COS formation proceedes faster and that greater stability of solid CaS may prevent formation of calcium polysulfides.

$$CaCO_3 + S_{x+1}^{2-} \rightarrow CaS + SO_2 + CO + S_{x-1}$$

$$CaCO_3 + SO_2 \rightarrow CaSO_3 + CO_2$$

$$CaSO_3 + S \rightarrow CaS_2O_3$$

$$4CaS_2O_3 \rightarrow CaS + 3CaSO_4 + S_4$$

$$CaSO_4 + S \rightarrow CaS + 2SO_2$$

Furthermore, the reaction

$$2Na_2SO_4 + S \rightarrow 2Na_2SO_3 + SO_2$$

takes place below 550  $^\circ\!\!\!\mathbb{C}$  . At temperature above 600  $^\circ\!\!\!\mathbb{C}$  , the reaction

$$Na_2SO_4 + 2S \rightarrow Na_2S + 2SO_2$$

may be predominant, but supply of free sulfur from the flux will become negligible at higher temperatures. In fact, gaseous product obtained beyond 550  $\degree$  does not contain detectable amount of SO<sub>2</sub>.

**Reaction of Na<sub>2</sub>CO<sub>3</sub> with Na<sub>2</sub>S<sub>2</sub>O<sub>3</sub>.** The total amount of gaseous product evolved corresponds to 0.31 mole per Na<sub>2</sub>S<sub>2</sub>O<sub>3</sub> used. Gas evolution starts at 420  $^{\circ}$ C and ends at about 600  $^{\circ}$ C (Figure 4). Since 1 mole of free sulfur as Na<sub>2</sub>S<sub>5</sub> is produced by disproportionation reaction (4) from 4 moles



Figure 4. Gas evolution for the reaction of  $Na_2CO_3$  and  $Na_2S_2O_3$ . The composition of the reactants:  $Na_2CO_3$ , 0.04 mole;  $Na_2S_2O_3$ , 0.04 mole.



**Figure 5.** Gas evolution for the reaction of  $CaCO_3$ , and  $Na_2S_2O_3$ . The composition of the reactants:  $CaCO_3$ , 0.002 mole;  $Na_2S_2O_3$ , 0.005 mole.

of Na<sub>2</sub>S<sub>2</sub>O<sub>3</sub>, more than 0.25 mole of CO<sub>2</sub> per mole of Na<sub>2</sub>S<sub>2</sub>O<sub>3</sub> is expected to be produced assuming the disproportionation reaction (4) and decomposition of Na<sub>2</sub>S<sub>2</sub>O<sub>3</sub>. When Na<sub>2</sub>S<sub>2</sub>O<sub>3</sub> alone is heated small amount of SO<sub>2</sub> evolves up to 600  $\degree$  .

**Reaction of CaCO<sub>3</sub> with Na<sub>2</sub>S<sub>2</sub>O<sub>3</sub>.** The gas evolution starts at about 475  $^{\circ}$ C and ends at about 800  $^{\circ}$ C. The volume of gas produced is about 0.32 mole per mole of Na<sub>2</sub>S<sub>2</sub>O<sub>3</sub> used (Figure 5). From Figure 3 and 4, supply of free sulfur by the disproportionation reaction (4) of sodium thiosulfate is fast enough for CaCO<sub>3</sub>+S reaction at about 450  $^{\circ}$ C.

**Reaction of Na<sub>2</sub>S with Y<sub>2</sub>O<sub>3</sub>.** It has been known that  $Y_2O_2S$  can be prepared by the reaction of  $Y_2O_3$  in Na<sub>2</sub>CO<sub>3</sub>+S flux (starts at 375 °C °), but the reaction of Na<sub>2</sub>S with  $Y_2O_3$  did not lead to formation of  $Y_2O_2S$ . The reaction of  $Y_2O_3$  in Na<sub>2</sub>CO<sub>3</sub>+S flux is very complex and the detailed reaction will be presented in a later paper.

Reaction of CaCO<sub>3</sub> with Na<sub>2</sub>S. no reaction was ob-

served.

### Conclusion

In conclusion, the carbonates may be assumed to attack the terminal sulfur-sulfur bond in the sulfur polymer to form CO, SO<sub>2</sub> and sulfide, and leaving free sulfur chain to form polysulfide with sulfide. Thus, the role of  $Na_2CO_3 + S$  flux is to keep free sulfur in the form of sodium polysulfide allowing the latter to react with other chemicals up to rather high temperature<sup>9</sup>. In case of carbonates reacting at higher temperatures, COS may also contribute to sulfurization reactions.

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