

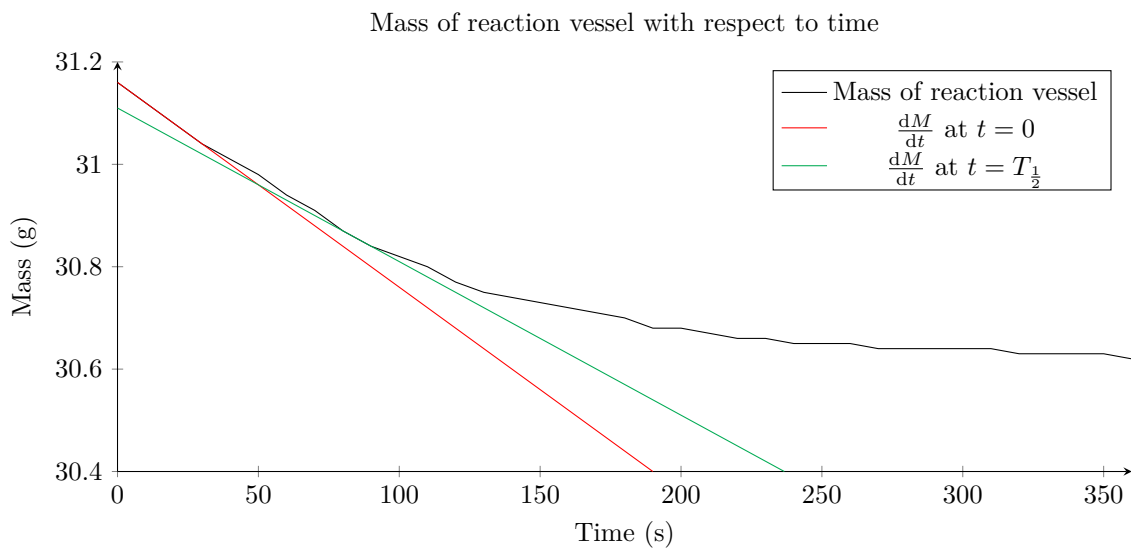
## PAG 9.2

The rate of reaction of calcium carbonate and hydrochloric acid

Rosie Bartlett

### 1 Mass experiment

#### 1.1 Results



#### 1.2 Analysis of results

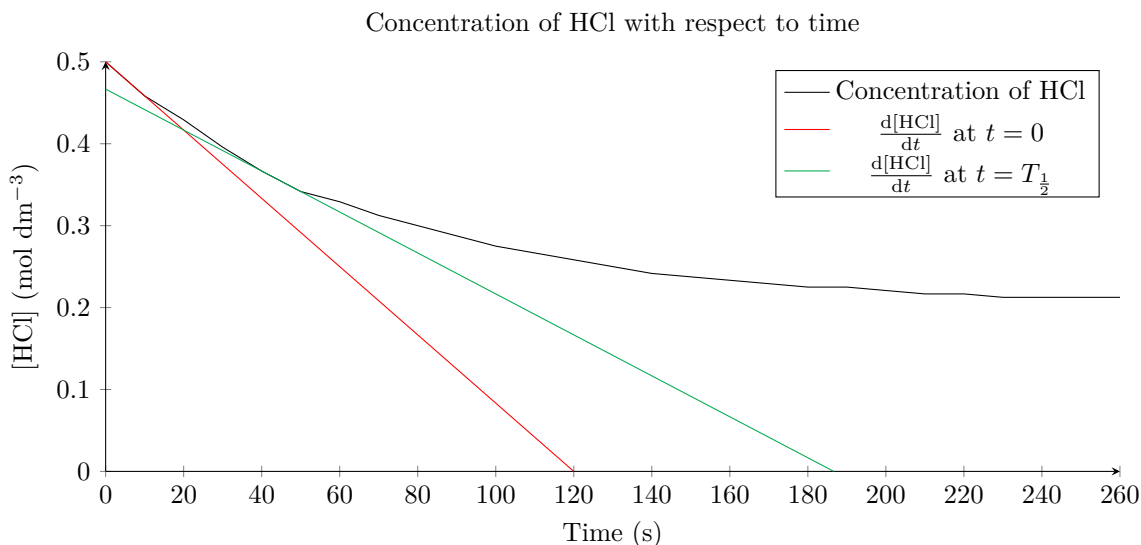
1. Half life 1  $\rightarrow$  75s  
Half life 2  $\rightarrow$  55s  
Half life 3  $\rightarrow$  55s
2. The half live values above include one anomaly, which when excluded give a constant half life.

### 2 Gas reaction

#### 2.1 Results

By using the formula below, we can calculate the concentration of HCl from the volume of gas produced. Where  $V(\text{CO}_2)$  is the volume of  $\text{CO}_2$  produced.

$$[\text{HCl}] = \frac{120 - V(\text{CO}_2)}{240}$$



## 2.2 Analysis of results

- Half life 1  $\rightarrow$  43.5s  
Half life 2  $\rightarrow$  50s  
Half life 3  $\rightarrow$  45.5s
- Since all the half lives are similar, we can suggest that the reaction is first order.

## 3 Extension opportunities

- (a) Mass lost:  
Since there was an excess of  $\text{CaCO}_3$ , when the reaction reached completion, there would have been 0.04 mol of  $\text{CO}_2$  produced. this would hav had a mass of 0.88g, meaning 0.88g would have been lost. In the reaction we only saw a loss of 0.5g, so the reaction did not go to completion.  
Gas produced:  
Again the  $\text{CaCO}_3$  was in excess, so at the end of the reaction, 0.01 mol of  $\text{CO}_2$  would have been produced, with a volume of 120cm<sup>3</sup>. we only saw a maximum volume of 69cm<sup>3</sup>, so the reaction did not go to completion.
- (b) As the reaction progresses, the concentration of HCl is constantly decreasing, which constantly decreases the rate, meaning a very long long time would be required to run the reaction to completion.
- (c) For the reaction to go to completion, we would need a larger gas syringe to account for the extra 20cm<sup>3</sup> of  $\text{CO}_2$  produced.
- Mass lost:  
At  $t = 0$ ,  $\frac{dM}{dt}$  was  $-4 \times 10^{-3} \text{ g s}^{-1}$ . At  $t = T_{\frac{1}{2}}$ ,  $\frac{dM}{dt}$  was  $-3 \times 10^{-3} \text{ g s}^{-1}$ .  
Gas produced:  
At  $t = 0$ ,  $\frac{d[\text{HCl}]}{dt}$  was  $\frac{1}{240} \text{ mol dm}^3 \text{ s}^{-1}$  or approximately  $4.17 \times 10^{-3} \text{ mol dm}^3 \text{ s}^{-1}$ . At  $t = T_{\frac{1}{2}}$ ,  $\frac{d[\text{HCl}]}{dt}$  was  $2.5 \times 10^{-3} \text{ mol dm}^3 \text{ s}^{-1}$ .  
In both cases, the gradient at  $T_{\frac{1}{2}}$  is approximately half of what t was at  $t = 0$ , giving further evidence that the reaction is first order.