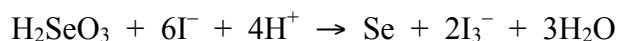


- The following reaction is run from 4 different starting positions.



Marks
6

Experiment Number	Initial $[\text{H}_2\text{SeO}_3]$ (mol L ⁻¹)	Initial $[\text{I}^-]$ (mol L ⁻¹)	Initial $[\text{H}^+]$ (mol L ⁻¹)	Initial rate of increase of $[\text{I}_3^-]$ (mol L ⁻¹ s ⁻¹)
1	0.100	0.100	0.100	1.000
2	0.100	0.075	0.100	0.422
3	0.075	0.100	0.100	0.750
4	0.100	0.075	0.075	0.237

Determine the rate law for the reaction.

The rate law has the form:

$$\text{rate} = k[\text{H}_2\text{SeO}_3]^x[\text{I}^-]^y[\text{H}^+]^z$$

Between experiments (1) and (2), only $[\text{I}^-]$ is varied:

$$\frac{\text{rate}_{(1)}}{\text{rate}_{(2)}} = \frac{([\text{I}^-]_{(1)})^y}{([\text{I}^-]_{(2)})^y} \quad \text{so} \quad \frac{1.000}{0.422} = \left(\frac{0.100}{0.075}\right)^y$$

Solving this gives $y = 3$.

Between experiments (1) and (3), only $[\text{H}_2\text{SeO}_3]$ is varied:

$$\frac{\text{rate}_{(1)}}{\text{rate}_{(3)}} = \frac{([\text{H}_2\text{SeO}_3]_{(1)})^x}{([\text{H}_2\text{SeO}_3]_{(3)})^x} \quad \text{so} \quad \frac{1.000}{0.750} = \left(\frac{0.100}{0.075}\right)^x$$

Solving this gives $x = 1$.

Between experiments (2) and (4), only $[\text{H}^+]$ is varied:

$$\frac{\text{rate}_{(2)}}{\text{rate}_{(4)}} = \frac{([\text{H}^+]_{(2)})^z}{([\text{H}^+]_{(4)})^z} \quad \text{so} \quad \frac{0.422}{0.237} = \left(\frac{0.100}{0.075}\right)^z$$

Solving this gives $z = 2$.

Rate law: $\text{rate} = k[\text{H}_2\text{SeO}_3][\text{I}^-]^3[\text{H}^+]^2$

ANSWER CONTINUES ON THE NEXT PAGE

Calculate the value of the rate constant.

Using experiment (1):

$$(1.000 \text{ mol L}^{-1} \text{ s}^{-1}) = k \times (0.100 \text{ mol L}^{-1}) \times (0.100 \text{ mol L}^{-1})^3 \times (0.100 \text{ mol L}^{-1})^2$$

$$k = 1.00 \times 10^6 \text{ mol}^{-5} \text{ L}^5 \text{ s}^{-1}$$

Answer: $k = 1.00 \times 10^6 \text{ mol}^{-5} \text{ L}^5 \text{ s}^{-1}$