• Consider the results of the following set of experiments studying the rate of the reaction of nitric oxide with hydrogen at 1280 °C.

$2\mathrm{INO}(g) + 2\mathrm{II}_2(g) \rightarrow \mathrm{IN}_2(g) + 2\mathrm{II}_2\mathrm{O}(g)$			
Experiment #	[NO] / M	$[H_2]/M$	Initial Rate / M s <sup>-1</sup>
1	$5.0 \times 10^{-3}$	$2.0 \times 10^{-3}$	$1.3 \times 10^{-5}$
2	$1.0 \times 10^{-2}$	$2.0 \times 10^{-3}$	$5.2 \times 10^{-5}$
3	$1.0 \times 10^{-2}$	$4.0 \times 10^{-3}$	$1.0  imes 10^{-4}$

 $2NO(g) + 2H_2(g) \rightarrow N_2(g) + 2H_2O(g)$ 

Write the rate law expression.

Between experiments 1 and 2,  $[H_2]$  is kept constant. Doubling [NO] (from  $5.0 \times 10^{-3}$  to  $1.0 \times 10^{-2}$  M) leads to the rate quadrupling. The reaction is second order with respect to [NO].

Between experiments 2 and 3, [NO] is kept constant. Doubling  $[H_2]$  (from  $2.0 \times 10^{-3}$  to  $4.0 \times 10^{-3}$  M) leads to the rate doubling. The reaction is first order with respect to  $[H_2]$ . Thus,

rate  $\alpha$  [NO]<sup>2</sup>[H<sub>2</sub>] = k[NO]<sup>2</sup>[H<sub>2</sub>]

Rate =  $k[NO]^2[H_2]$ 

Calculate the rate constant, k. Include units in your answer.

Using experiment 1 and rate =  $k[NO]^2[H_2]$ :

$$(1.3 \times 10^{-5} \text{ M s}^{-1}) = k \times (5.0 \times 10^{-3} \text{ M})^2 \times (2.0 \times 10^{-3} \text{ M})$$
 so  $k = 260 \text{ M}^{-2} \text{ s}^{-1}$ 

 $(M s^{-1}) = (units of k) \times (M)^2 \times (M)$  so the units of k are  $M^{-2} s^{-1}$ 

 $k = 260 \text{ M}^{-2} \text{ s}^{-1}$ 

What is the rate of the reaction when [NO] is  $1.2 \times 10^{-2}$  M and [H<sub>2</sub>] is  $6.0 \times 10^{-3}$  M?

Using rate = 
$$(260 \text{ M}^{-2} \text{ s}^{-1})[\text{NO}]^{2}[\text{H}_{2}]$$
:  
rate =  $(260 \text{ M}^{-2} \text{ s}^{-1}) \times (1.2 \times 10^{-2} \text{ M})^{2} \times (6.0 \times 10^{-3} \text{ M}) = 2.2 \times 10^{-4} \text{ M s}^{-1}$ 

Rate =  $2.2 \times 10^{-4} \text{ M s}^{-1}$ 

• What is the value of the equilibrium constant for the following reaction at 298 K?

 $2Fe^{3+}(aq) + Sn(s) \implies Sn^{2+}(aq) + 2Fe^{2+}(aq)$ 

The reduction half cell reactions and  $E^0$  values are:

Fe<sup>3+</sup>(aq) + e<sup>-</sup> → Fe<sup>2+</sup>(aq)  $E^0 = +0.77 \text{ V}$ Sn<sup>2+</sup>(aq) + 2e<sup>-</sup> → Sn(s)  $E^0 = -0.14 \text{ V}$ 

In the reaction, Sn is being oxidized and so the overall cell potential is:

 $E^0 = ((+0.77) - (-0.14))$  V = +0.91 V

The reaction involves 2 electrons so, using  $E^0 = \frac{RT}{nF} \ln K$ :

$$\ln K = E^{0} \times \frac{nF}{RT} = (+0.91 \text{ V}) \times \left(\frac{2 \times 96485 \text{ C mol}^{-1}}{8.314 \text{ J K}^{-1} \text{ mol}^{-1} \times 298 \text{ K}}\right) = 70.9$$
$$K = e^{70.9} = 6.05 \times 10^{30}$$

Answer:  $6.05 \times 10^{30}$