

# **Teaching rates of reaction post-16: next steps**

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These steps follow those from the graphical representations in the Teacher checklist, which accompanied the first Teaching rates of reaction post-16 article (<u>rsc.li/3yLp1nU</u>).

## Introducing the rate equation

Students should be familiar with transforming a relationship that shows a proportional relationship into an equation by including a constant of proportionality.

### Rate $\propto [x]^n$ becomes Rate = $k[x]^n$

Where k is the rate constant. This is only a constant when the temperature remains the same or when a catalyst doesn't affect the rate.

# Calculating the rate constant

Now take Log<sub>10</sub>

### Log rate = logk + nlog[x]

This equation form can be compared to y = mx + c

Log rate	=	logk	+	nlog[x]
У	=	С	+	тх



So, plotting *log[x]* against *log rate* allows the gradient *n* (order) to be calculated. The intercept is *logk* which is important for calculating the *Ea*.

### Calculating the activation energy $(E_a)$

Applying the Arrhenius equation

#### $k = Ae^{-Ea/RT}$

Take natural logarithms and plot Ink versus 1/T

