Year 12 Chemistry

# Unit 3 Revision Lecture Monash University presented by Peter Skinner

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## 1 Course structure

- Unit 3: 2 SACs, 16% of study score
  - 1. Energy production
    - Obtaining energy from fuels
    - Fuel choices
    - Galvanic cells
    - Fuel cells
  - 2. Optimising yield
    - Rate of reactions
    - Extent of reactions
    - Production via electrolysis
    - Rechargable batteries
- Unit 4: 3 SACs, 24% of study score
- Exam: 60% of study score
  - -15 minutes reading time, 2.5 hours writing time
  - 30 multiple choice questions (spend 30—45 minutes, **do last**) harder in new study design
  - 90 marks written questions (spend 1 hr 45 m—2 hr)
  - Last year:
    - \* 23% calculations (21 marks)
    - \* 44% extended answer (40 marks)
    - $\ast~32\%$  short answer (29 marks)
  - 5—10 marks on writing chemical equations
  - Same marking panel as last year
  - Indirect assessment of pracs
  - $\ge 1$  mark for significant figures
  - Importance of written communication
  - First parts are important, no consequential marks
  - Use dot points (short form) especially in rates & concentration

#### Key points

- Spend 30—45 minutes on multiple choice
- Focus on redox reactions
- Use data book
- Multiple choice questions are hard
- Memorise oxidation numbers

# 2 Energy production

- $C = n \div v$  or  $C = m \div V$  (concentration in g L<sup>-1</sup>)
- Gases: PV = nRT and  $n = V \div V_m$
- Past exams before 2017 use different SLC
- Renewability *reasonable* timeframe
- Fuel choices consider:
  - External temperature
  - Viscosity (intermolecular forces)
  - Hygroscopic properties (attracts water  $\implies$  forms H-bonds)
  - Cloud point
- Blended fuels use energy per mass not energy per mol

# 3 Yield & rate

- Equilibrium constant  $K_C$  needs units
- $K_C \equiv K$
- Example question for rates: limiting factor for rate, given a set (equal) rate of both reactants consumption/production
- Collision theory:
  - 1. Particles must collide
  - 2. Particles must collide with sufficient energy to overcome  $E_A$
- Increase of rate with temperature:
  - 1.  $\uparrow$  temperature  $\implies \uparrow$  energy  $\implies$  more frequent collisions
  - 2.  $\uparrow$  temperature  $\implies \uparrow$  energy  $\implies$  collisions occur with greater energy
    - (  $\implies$  greater *proportion* of particles that can react per unit time)
- $\uparrow c(\text{reactants}) \implies \text{more collisions}$
- Definition of *rate*: more products per unit time  $\longrightarrow$  faster rate
- Cause and effect: propose hypothesis and prove by induction
- Maxwell-Boltzmann distributions  $x_{\text{peak}}$  is constant for different concentrations
- Memorise definition of *catalyst*: provides a reaction with an alternative energy pathway which has a lower activation energy

### 3.1 Equilibria

- all reactants and products are present at equilibrium
- $K_C$  is fixed at a constant temperature and reaction
- $K_C$  changes with concentrations (relative)
- If reaction equation is reversed,  $K_C$  value will be the reciprocal
- If temperature changes,  $K_C$  will change (but not necessarily proportionally)
- Le Chatelier's principle: If a change is made to a system at equilibrium, the system will partially oppose this change if it is possible

- Accuracy of graph drawing use **clear** plastic ruler
  - Label vertical ratios
- Use concentration table format for calculating equilibrium constant  $K_C$

#### Important

 $K_C$  is **not** related to the rate of reaction  $\implies$  we cannot say how fast a reaction os going to occur from the  $K_C$  value

#### 3.2 Exothermic & endothermic reactions

- All combustion reactions are exothermic
- Data book: molar heat of combustion =  $|\Delta H|$
- Endothermic reactions rarely occur naturally (creates instability/entropy)
- $E_A = |E_{\max} E_{\text{initial}}|$
- If coefficients of a thermochemical equation are changed,  $\Delta H$  also changes
- Possible data discrepencies in theoretical results:
  - State of  $H_2O$
  - Incomplete combustion
  - Heat loss to environment
- Analogy with simultaneous equations
- Calorimetry insulate *sides* of can not bottom. Be specific.

Multiple choice question examples (features of **exothermic** reactions):

a) Products are as they have le	ss chemical energy than reactants	(more stable)
b) required to break bonds in	n products compared to reactants	(more chemical energy)
Multiple choice question examples (features	of <b>endothermic</b> reactions):	
a) Transformation of energy from	om surroundings into	(thermal, chemical)
b) : Surroundings and reaction becomes		(colder)

# 4 Oxidation numbers (memorise)

Species	Rule
Elements	Always 0
Ions	Same as common ion
Hydrogen	+1 (unless present as $H_2O$ - O.N. = 0; or as hydride - O.N. = -1)
Oxygen	-2 (unless present as $O_2$ - O.N. = 0; or as peroxide - O.N. = -1)
Molecules	Sum of O.N. must equal zero
Molecular ions	Sum of O.N. must equal overall charge on ion

# 5 Redox reactions

- Verify equations: check charge of each side independently: charge(LHS) = charge(RHS)
- Electrochemical series always has oxidants on left
- Top left and bottom right always react spontaneously
- For electrochemical cell questions: first parts are important, no consequential marks
- Non-standard conditions can alter positions of half-equations on electrochemical series and change  $E^0$  values
- Secondary cells polarity is constant, but reaction at each electrode swaps

#### 5.1 Galvanic cells

- Value of  $E^0$  is *not* a reliable indicator for rate of reaction
- Half cells are physically separate
- Products must remain in contact with electrodes

#### 5.2 Electrolytic cells

- Possible question: name observations
  - Bubbles
  - O<sub>2</sub> would *not* be visible
  - Cannot see  $\uparrow$  [H<sup>+</sup>]
  - Can see Cu(s) deposit on electrode
  - Can see colour change (pH)  ${\rm Cu_2}^+$  solution can be an indicator
- Less side reactions in e.g. lithium ion cells (efficiency)
- Lower reactions in electrochemical series do not occur forwards  $(L \rightarrow R)$
- Check state of H<sub>2</sub>O can it be liquid at that temperature?

### 5.3 Fuel cells

- Galvanic cells are primary cells, fuel cells are not primary are they secondary?
- Major disadvantage of fuel cells: expensive electrodes (they must also function as catalysts)
- Fuel cells same overall reaction as combustion
- Reactangs must not come into contact
- Highly efficient

#### 5.4 Electrochemical series

Strongest oxidant will always react preferentially with best reductant Always identify *all* chemicals present in reaction on electrochemical series