

## Q1.

- 2 (a)  $\times$   
 $\times$   
 $\checkmark$   
 $\checkmark$  (-1 for each error or omission) ..... B2 [2]
- (b) heat lost by liquid gold =  $0.95m \times 129 \times \Delta T$  ..... C1  
 heat gained (silver) =  $0.05m \times 235 \times (1340 - 300) + 0.05m \times 105\,000$  ..... C1, C1  
 $122.5m\Delta T = 17\,470m$   
 $\Delta T = 143\text{ K}$  ..... C1  
 temperature =  $143 + 1340 = 1483\text{ K}$  ..... A1 [5]
- (c) e.g. thermocouple/resistance thermometer ..... B1 [1]

## Q2.

6	0	-	+	(-1 for each error)	B2	
	+	+	0	(-1 for each error)	B2	
	+	+	0	(-1 for each error)	B2	[6]
Total						[6]

## Q3.

- 3 (a) (thermal) energy/heat required to convert unit mass/1 kg of solid to liquid with no change in temperature/at melting point ..... M1  
 ..... A1 [2]
- (b) (i) energy required to warm ice =  $24 \times 10^{-3} \times 2.1 \times 10^3 \times 15$  (= 756 J) ..... C1  
 energy required to melt ice at  $0^\circ\text{C}$  =  $24 \times 10^{-3} \times 330 \times 10^3$  (= 7920 J) ..... C1  
 total energy = 8700 J ..... A1 [3]
- (ii) energy lost by warm water =  $200 \times 10^{-3} \times 4.2 \times 10^3 \times (28 - T)$  ..... C1  
 $200 \times 4.2 \times (28 - T) = 24 \times 4.2 \times T + 8676$  ..... C1  
 $T = 16^\circ\text{C}$  ..... A1 [3]  
 [allow 2 marks if  $\Delta T$  calculated]  
 [allow 2 marks if  $(24 \times 4.2 \times T)$  omitted]  
 [allow 1 mark for  $224 \times 4.2 \times (28 - T) = 8676$ ,  $T = 19^\circ\text{C}$ ]

## Q4.

- 3 (a) gradient of graph is (a measure of) the sensitivity the gradient varies with temperature ..... M1  
 ..... A1 [2]
- (b)  $2040 \pm 20\ \Omega$  corresponds to  $15.0 \pm 0.2^\circ\text{C}$  ..... C1  
 $T/\text{K} = T/^\circ\text{C} + 273.15$  (allow 273.2) ..... C1  
 temperature is 288.2 K ..... A1 [3]



## Q5.

- 2 (a) (on melting,) bonds between molecules are broken/weakened  
or molecules further apart/are able to slide over one another  
kinetic energy unchanged so no temperature change  
potential energy increased/changed so energy required  
B1  
B1  
B1 [3]
- (b) thermal energy/heat required to convert unit mass of solid to liquid  
with no change in temperature/ at its normal boiling point  
M1  
A1 [2]
- (c) (i) thermal energy lost by water =  $0.16 \times 4.2 \times 100$   
= 67.2 kJ  
 $67.2 = 0.205 \times L$   
 $L = 328 \text{ kJ kg}^{-1}$   
C1  
C1  
A1 [3]
- (ii) more energy (than calculated) melts ice  
so, (calculated)  $L$  is lower than the accepted value  
M1  
A1 [2]

## Q6.

- 3 (a) increasing separation of molecules / breaking bonds between molecules  
(allow atoms/molecules, overcome forces)  
doing work against atmosphere (during expansion)  
B1  
B1 [2]
- (b) (i) 1 either bubbles produced at a constant rate / mass evaporates/lost at  
constant rate  
or find mass loss more than once and this rate should be constant  
or temperature of liquid remains constant  
B1 [1]  
B1 [1]  
2 to allow/cancel out/eliminate/compensate for heat losses (to atmosphere)  
(do not allow 'prevent'/'stop')
- (ii) use of  $\text{power} \times \text{time} = \text{mass} \times \text{specific latent heat}$   
 $(70 - 50) \times 5 \times 60 = (13.6 - 6.5) \times L$   
 $L = 845 \text{ J g}^{-1}$   
C1  
C1  
A1 [3]

## Q7.

- 3 (a) (i) 1 deg C corresponds to  $(3840 - 190) / 100 \Omega$   
for resistance  $2300 \Omega$ , temperature is  $100 \times (2300 - 3840) / (190 - 3840)$   
temperature is  $42^\circ\text{C}$   
C1  
A1 [2]
- (ii) either  $286 \text{ K} \equiv 13^\circ\text{C}$  or  $42^\circ\text{C} \equiv 315 \text{ K}$   
thermodynamic scale does not depend on the property of a substance  
so change in resistance (of thermistor) with temperature is non-linear  
B1  
M1  
A1 [3]
- (b) heat gained by ice in melting =  $0.012 \times 3.3 \times 10^5 \text{ J}$   
= 3960 J  
C1  
heat lost by water =  $0.095 \times 4.2 \times 10^3 \times (28 - \theta)$   
 $3960 + (0.012 \times 4.2 \times 10^3 \times \theta) = 0.095 \times 4.2 \times 10^3 \times (28 - \theta)$   
 $\theta = 16^\circ\text{C}$   
C1  
C1  
A1 [4]  
(answer  $18^\circ\text{C}$  – melted ice omitted – allow max 2 marks)  
(use of  $(\theta - T)$  then allow max 1 mark)

**Q8.**

- |         |   |                |     |
|---------|---|----------------|-----|
| 4       | (a) $+\Delta U$ : increase in internal energy<br>+q: thermal energy / heat supplied to the system<br>+w: work done on the system                                | B1<br>B1<br>B1 | [3] |
| (b) (i) | (thermal) energy required to change the state of a substance per unit mass without any change of temperature  | M1<br>A1<br>A1 | [3] |
| (ii)    | when evaporating<br>greater change in separation of atoms/molecules<br>greater change in volume<br>identifies each difference correctly with $\Delta U$ and $w$ | M1<br>M1<br>A1 | [3] |

**Q9.**

- |            |  |                      |     |
|------------|--|----------------------|-----|
| 3          | (a) (numerically equal to) quantity of (thermal) energy required to change the state of unit mass of a substance without any change of temperature<br>(Allow 1 mark for definition of specific latent heat of fusion/vaporisation) | M1<br>A1             | [2] |
| (b) either | energy supplied = $2400 \times 2 \times 60 = 288000 \text{ J}$<br>energy required for evaporation = $106 \times 2260 = 240000 \text{ J}$<br>difference = $48000 \text{ J}$<br>rate of loss = $48000 / 120 = 400 \text{ W}$         | C1<br>C1<br><br>A1   |     |
| or         | energy required for evaporation = $106 \times 2260 = 240000 \text{ J}$<br>power required for evaporation = $240000 / (2 \times 60) = 2000 \text{ W}$<br>rate of loss = $2400 - 2000 = 400 \text{ W}$                               | (C1)<br>(C1)<br>(A1) | [3] |

**Q10.**

- |         |   |                |     |
|---------|---|----------------|-----|
| 3       | (a) sum of potential energy and kinetic energy of atoms/molecules/particles reference to random (distribution)  | M1<br>A1       | [2] |
| (b) (i) | as lattice structure is 'broken'/bonds broken/forces between molecules reduced (not molecules separate)<br>no change in kinetic energy, potential energy increases<br>internal energy increases                                       | B1<br>M1<br>A1 | [3] |
| (ii)    | either molecules/atoms/particles move faster/ $\langle c^2 \rangle$ is increasing<br>or kinetic energy increases with temperature (increases)<br>no change in potential energy, kinetic energy increases<br>internal energy increases | B1<br>M1<br>A1 | [3] |

**Q11.**

**MEGA LECTURE**

- 1 (a) (i)  $Q = mc\Delta\theta$  ..... C1  
 $2300 = 0.75 \times c \times (100 - 20) / 120$  ... (if uses  $\pm 273$ , then -2)..... C1  
 $c = 4600 \text{ J kg}^{-1} \text{ K}^{-1}$  ... (allow 1 sf)..... A1  
(ii)  $Q = mL$  ..... C1  
 $2300 = (0.375 / 420) \times L$   
 $L = 2.6 \times 10^6 \text{ J kg}^{-1}$  ... (allow 1 sf)..... A1 [5]
- (b) e.g. heat losses, power not constant etc ..... M1  
(do not allow if related to s.h.c., rather than l.h.c.)  
effect on value for  $L$  ..... A1 [2]

**Q12.**

- 3 (a)  $pV/T = \text{constant}$ ..... C1  
 $T = (6.5 \times 10^6 \times 30 \times 300) / (1.1 \times 10^5 \times 540)$ ..... C1  
 $= 985 \text{ K}$ ..... A1 [3]  
(if uses  $^{\circ}\text{C}$ , allow 1/3 marks for clear formula)
- 3 (b) (i)  $\Delta U = q + w$  ..... M1  
symbols identified correctly ..... A1 [2]  
directions correct.....
- (ii)  $q$  is zero ..... B1  
 $w$  is positive OR  $\Delta U = w$  and  $U$  increases ..... B1  
 $\Delta U$  is rise in kinetic energy of atoms ..... M1  
and mean kinetic energy  $\propto T$  ..... A1 [4]  
(allow one of the last two marks if states ' $U$  increases so  $T$  rises')

**Q13.**

- 7 (a) variation is non-linear ..... 1  
two possible temperatures ..... 1 [2]
- (b) e.g. 1. small thermal capacity/measure  $\Delta\theta$  of small object  
/short response time  
2 readings taken at a point/physically small  
3 can be used to measure temperature difference  
4 no power supply required  
etc. (any two, 1 mark each) ..... 2 [2]

**Q14.**





3	(a)	correct statement, words or symbols .....	B1	[1]
	(b)(i)	$w = p\Delta V$ ..... $= 1.03 \times 10^5 \times (2.96 \times 10^{-2} - 1.87 \times 10^{-5})$ $= (-) 3050 \text{ J}$ .....	C1 A1	[2]
	(ii)	$q = 4.05 \times 10^4 \text{ J}$ .....	B1	[1]
	(iii)	$\Delta U = 4.05 \times 10^4 - 3050 = 37500 \text{ J}$ ...no e.c.f. from (a)..... penalise 2 sig.fig. once only	A1	[1]
	(c)	number of molecules = $N_A$ ..... energy = $37500 / (6.02 \times 10^{23})$ $= 6.2 \times 10^{-20} \text{ J}$ (accept 1 sig.fig.) .....	C1 A1	[2]

### Q15.

2	(a)	(i)	idea of heat lost (by oil) = heat gained (by thermometer) $32 \times 1.4 \times (54 - t) = 12 \times 0.18 \times (t - 19)$ $t = 52.4^{\circ}\text{C}$	C1 C1 A1	[3]
		(ii)	either ratio (= $1.6/54$ ) = 0.030 or (= $1.6/327$ ) = 0.0049	A1	[1]
	(b)	thermistor thermometer (allow 'resistance thermometer') because small mass/thermal capacity	B1 B1	[2]	
	(c)	boiling point temperature is constant further comment e.g. heating of bulb would affect only rate of boiling	M1 A1	[2]	

### Q16.

2	(a)	(Thermal) energy / heat required to convert unit mass of solid to liquid at its normal melting point / without any change in temperature (reference to 1 kg or to ice → water scores max 1 mark)	M1 A1	[2]
	(b)	(i) To <u>make allowance</u> for heat gains from the atmosphere	B1	[1]
		(ii) e.g. constant rate of production of droplets from funnel constant mass of water collected per minute in beaker (any sensible suggestion, 1 mark)	B1	[1]
		(iii) mass melted by heater in 5 minutes = $64.7 - \frac{1}{2} \times 16.6 = 56.4 \text{ g}$ $56.4 \times 10^{-3} \times L = 18$ $L = 320 \text{ kJ kg}^{-1}$ (Use of $m = 64.7$ , giving $L = 278 \text{ kJ kg}^{-1}$ , scores max 1 mark use of $m = 48.1$ , giving $L = 374 \text{ kJ kg}^{-1}$ , scores max 2 marks)	C1 C1 A1	[3]

### Q17.



- 4 (a)  $\Delta U = q + w$  (allow correct word equation) B1 [1]
- (b) *either* kinetic energy constant because temperature constant M1  
 potential energy constant because no intermolecular forces M1  
 so no change in internal energy A1 [3]  
*or* kinetic energy and potential energy both constant (M1)  
 so no change in internal energy (A1)  
 reason for *either* constant k.e. or constant p.e. given (A1)

### Q18.

- 3 (a) e.g. two objects of different masses at same temperature (M1)  
 same material would have different amount of heat (A1)  
 e.g. temperature shows direction of heat transfer (M1)  
 from high to low regardless of objects (A1)  
 e.g. when substance melts/boils (M1)  
 heat input but no temperature change (A1)  
 any two, M1 + A1 each, max 4 ..... [4]
- (b) (i) energy losses (to the surroundings) ..... M1  
*either* increase as the temperature rises ..... A1 [2]  
*or* rise is zero when heat loss = heat input ..... A1
- (ii) idea of input power = maximum rate of heat loss ..... C1  
 power =  $m \times c \times \Delta\theta / \Delta t$   
 $54 = 0.96 \times c \times 3.7 / 60$  ..... C1  
 $c = 910 \text{ J kg}^{-1} \text{ K}^{-1}$  ..... A1 [3]
- [Total: 9]

### Q19.

- 2 (a) sum of kinetic and potential energies of molecules / particles / atoms ..... M1  
 random (distribution) ..... A1 [2]
- (b)  $+\Delta U$ : increase in internal energy ..... B1  
 $+q$ : heating of / heat supplied to system ..... B1  
 $+w$ : work done on system ..... B1 [3]
- (c) (i) work done =  $p\Delta V$  ..... C1  
 $= 1.0 \times 10^5 \times (2.1 - 1.8) \times 10^{-3}$   
 $= 30 \text{ J}$  ..... M1  
 $w = 30 \text{ J}, q = 0$  so  $\Delta U = 30 \text{ J}$  ..... A1 [3]
- (ii) these three marks were removed, as insufficient data was given in the question.
- [Total: 8]

### Q20.



- 2 (a) temperature scale calibrated assuming linear change of property with temperature  
neither property varies linearly with temperature B1 B1 [2]
- (b) (i) does not depend on the property of a substance B1 [1]  
(ii) temperature at which atoms have minimum/zero energy B1 [1]
- (c) (i) 323.15 K A1 [1]  
(ii) 30.00 K A1 [1]

### Q21.

- 3 (a) temperature of the spheres is the same  
no (net) transfer of energy between the spheres B1 B1 [2]
- (b) (i) power =  $m \times c \times \Delta\theta$  where  $m$  is mass per second  
3800 =  $m \times 4.2 \times (42 - 18)$  C1  
 $m = 38 \text{ gs}^{-1}$  C1 A1 [3]  
(ii) some thermal energy is lost to the surroundings  
so rate is an overestimate M1 A1 [2]

### Q22.

- 3 (a) *either* change in volume =  $(1.69 - 1.00) \times 10^{-3}$   
*or* liquid volume  $\ll$  volume of vapour M1  
work done =  $1.01 \times 10^5 \times 1.69 = 1.71 \times 10^5 \text{ (J)}$  A1 [2]
- (b) (i) 1. heating of system/thermal energy supplied to the system B1 [1]  
2. work done on the system B1 [1]
- (ii)  $\Delta U = (2.26 \times 10^6) - (1.71 \times 10^5)$  C1  
 $= 2.09 \times 10^6 \text{ J (3 s.f. needed)}$  A1 [2]

### Q23.



3 (a) resonance B1 [1]

(b)  $Pt = mc \Delta\theta$  C1  
 $750 \times 2 \times 60 = 0.28 \times c \times (98 - 25)$  C1  
 $c = 4400 \text{ J kg}^{-1} \text{ K}^{-1}$  A1 [3]

(use of  $\Delta\theta = 73 + 273$  max. 1/3)

(use of  $t = 2 \text{ s}$  not  $120 \text{ s}$  max. 2/3)

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(c) e.g. some microwave leakage from the cooker  
 e.g. container for the water is also heated  
 (any sensible suggestion) B1 [1]

## Q24.

3 (a) initially,  $pV/T = (2.40 \times 10^5 \times 5.00 \times 10^{-4})/288 = 0.417$  M1  
 finally,  $pV/T = (2.40 \times 10^5 \times 14.5 \times 10^{-4})/835 = 0.417$  M1  
 ideal gas because  $pV/T$  is constant A1 [3]  
 (allow 2 marks for two determinations of  $V/T$  and then 1 mark for  $V/T$  and  $p$  constant, so ideal)

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(b) (i) work done  $= p\Delta V$   
 $= 2.40 \times 10^5 \times (14.5 - 5.00) \times 10^{-4}$  C1  
 $= 228 \text{ J}$  (ignore sign, not 2 s.f.) A1 [2]

(ii)  $\Delta U = q + w = 569 - 228$  M1  
 $= 341 \text{ J}$  A1 [2]  
 increase





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