

**E.M.D. N°01**

**Exercise 1 (10pt)**

For magnesium (Mg), the following data are given at 298 K:

	O <sub>2</sub> (G)	Mg(S)	MgO(S)
$\Delta_f H^0$ (en KJmol <sup>-1</sup> )	0	0	-601,5
$S^0$ (en J K <sup>-1</sup> mol <sup>-1</sup> )	205	32,7	27

Melting temperature  $T_F = 923$  K with  $L_F = 9.2$  kJ.mol<sup>-1</sup>; Boiling (vaporization) temperature  $T_{VAP} = 1393$ K with  $L_{VAP} = 131,8$  kJ.mol<sup>-1</sup>

- Calculate  $\Delta_r G^\circ(T)$  and draw the Ellingham diagram of magnesium.

**Exercise 2 (4pt)**

At 30°C, the vapor in equilibrium with an ideal solution of a kerosene (K) / MIBK mixture (MIBK = methyl isobutyl ketone, used in liquid–liquid extraction) has:

- Total pressure:  $P_{tot} = 0.420$  atm
- Vapor-phase mole fraction of MIBK:  $y_{MIBK} = 0.780$

At this temperature, the saturated vapor pressure of MIBK is:

- $P_{MIBK}^{sat} = 0.610$  atm

Calculate the mole fraction of MIBK in the liquid solution  $x_{MIBK}$ .

**Exercise 3 (6pt)**

A quantity of 0.6 mol of calcium carbonate (CaCO<sub>3</sub>) is introduced into a 30 L vessel kept at constant temperature, initially evacuated (vacuum).

At 1480 K, the standard Gibbs free energy of reaction is  $\Delta G^\circ = -8.5$  kJ.mol<sup>-1</sup>.

1. Write the expression of the equilibrium constant and calculate its value at 1480 K.
2. Deduce the equilibrium CO<sub>2</sub> pressure at 1480 K.
3. Calculate the equilibrium composition of the system.

Given:  $R = 8,31$  J mol<sup>-1</sup> K<sup>-1</sup>

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**Exercise (10 pts)**

Pour923> T>298 Mg is solid and MgO is solid (2.5 pts)

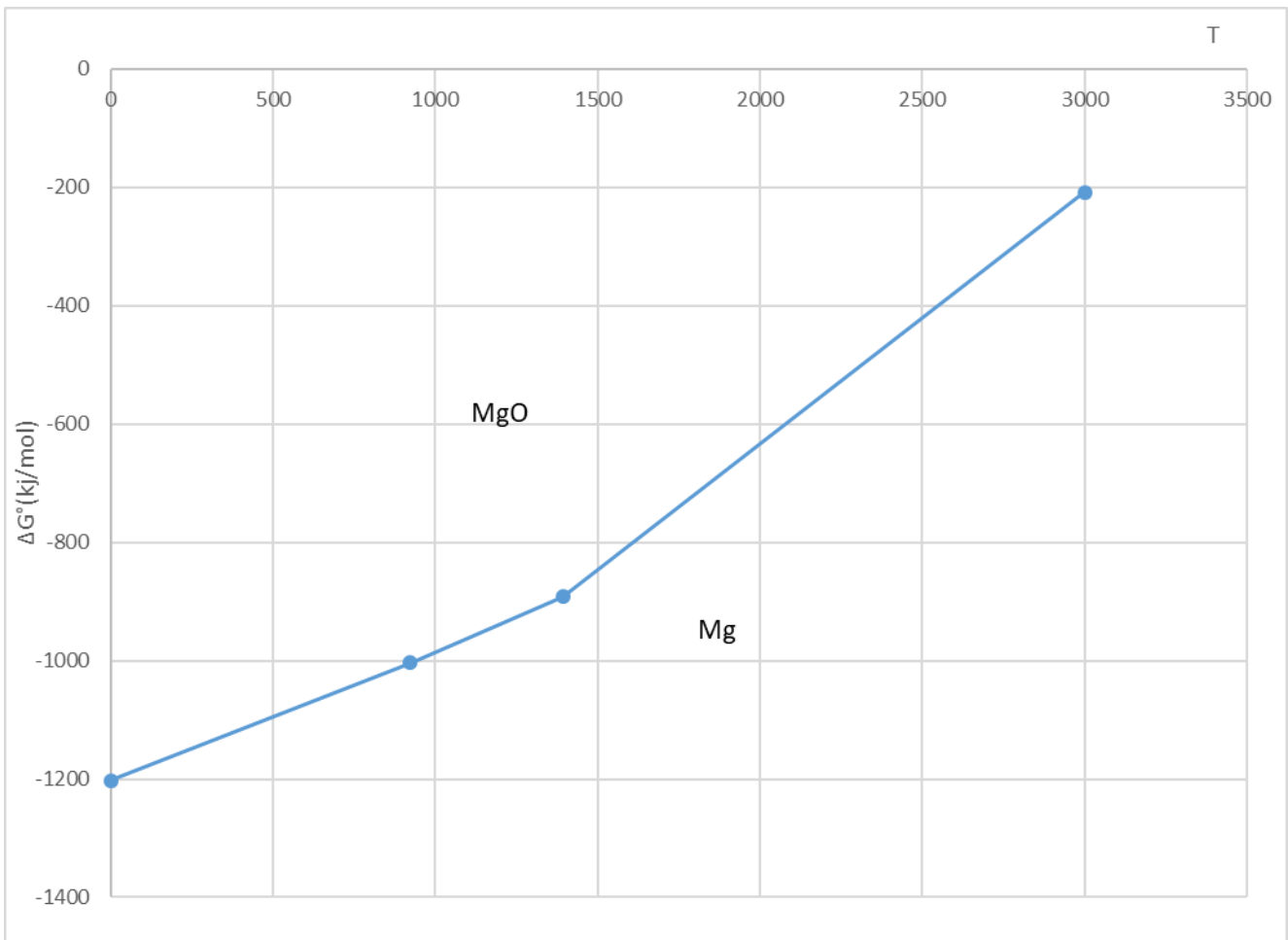
d'où  $\Delta_r G^\circ(T) = -1203 + 216,4 \cdot 10^{-3} \cdot T$

Pour923< T<1393 Mg is liquid and MgO is solid (2.5 pts)

d'où  $\Delta_r G^\circ(T) = -1221,4 + 236,3 \cdot 10^{-3} \cdot T$

Pour1393< T Mg is gas and MgO is solid (2.5 pts)

d'où  $\Delta_r G^\circ(T) = -1485 + 425,5 \cdot 10^{-3} \cdot T$



(2.5 pts)

## Exercise 2 (4pt)

### 1) Vapor phase (Dalton's law):

$$y_{\text{MIBK}} = P_{\text{MIBK}} / P_{\text{tot}}$$

$$P_{\text{MIBK}} = y_{\text{MIBK}} \cdot P_{\text{tot}} = 0,780 \times 0,420$$

$$P_{\text{MIBK}} = 0,328 \text{ atm} \quad (2 \text{ pts})$$

### 2) Liquid phase (Raoult's law):

$$P_{\text{MIBK}} = x_{\text{MIBK}} \cdot P_{\text{MIBK}}^{\text{sat}}$$

$$0,328 = x_{\text{MIBK}} \cdot 0,610$$

$$x_{\text{MIBK}} = 0,328 / 0,610 = 0,538 \quad (2 \text{ pts})$$

## Exercise (6pts)

1.

$$K_0^T = e^{\frac{-\Delta_r G_{1480}^0}{RT}} \quad (1 \text{ pt})$$

$$K_0^T = e^{\frac{8500}{8,31 \times 1480}} = e^{0,691} = 1,996 \quad (1 \text{ pt})$$

2.  $P_{\text{CO}_2} = K_{1480}^0 x P_0 = 1,996 \times 1 = 1,996 \text{ bar} \quad (1 \text{ pts})$

3. Equilibrium composition (ideal gas):.

CO<sub>2</sub> is treated as an ideal gas.  $x = PV/(RT) = 1,996 \times 10^5 \times 0,030 / (8,31 \times 1480) = 0,487 \text{ mol.}$   
(1.5 pt)

	extent of reaction (mol)	CaCO <sub>3</sub> (s)	CaO(s)	+CO <sub>2</sub> (g)
Initial (mol)	0	0,6	0	0
At equilibrium (mol)	x 0,487	0,6-x 0.113	x 0,487	x 0,487

(1.5 pt)