

KEEMIAÜLESANNETE LAHENDAMISE LAHTINE VÕISTLUS

Noorem rühm (9. ja 10. klass)

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Ülesannete lahendused

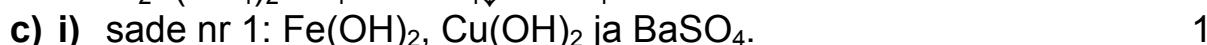
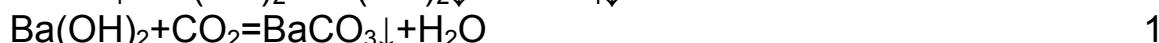
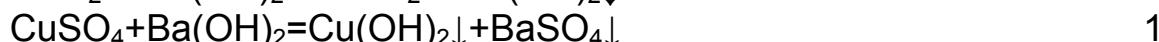
1. a) i) $\%(\text{CuSO}_4) = \frac{56,1 \text{ g} \cdot 159,5 \frac{\text{g}}{\text{mol}} \cdot 100}{249,5 \text{ g/mol} \cdot (56,1 \text{ g} + 457 \text{ g})} = 6,99$ 1

ii) $n(\text{CuSO}_4) = \frac{342 \text{ g} \cdot 0,0699}{159,5 \text{ g/mol}} = 0,150 \text{ mol}$ 1

iii) $n(\text{Ba(OH)}_2) = \frac{2,29 \text{ kg} \cdot 1000 \text{ g/kg} \cdot 0,05}{171 \text{ g/mol}} = 0,67 \text{ mol}$ 1

iv) $n(\text{CO}_2) = \frac{7,2 \text{ dm}^3}{22,4 \text{ dm}^3/\text{mol}} = 0,32 \text{ mol}$ 1

v) $n((\text{NH}_4)_2\text{SO}_4) = \frac{118 \text{ cm}^3 \cdot 1,115 \text{ g/cm}^3 \cdot 0,2}{132 \text{ g/mol}} = 0,2 \text{ mol}$ 1



d) Baariumhüdroksiidi jäääb peale esimest sadestamist alles 0,67 mol – 0,2 mol – 0,15 mol = 0,32 mol. Tekib 0,2 mol BaCl_2 . Süsihaptegaasiga sadestatakse 0,32 mol Ba^{2+} ja neutraliseeritakse ära hüdroksiid. Lahusesse jäääb alles 0,2 mol baariumkloriidi, mis ammoniumsulfaadiga reageerides annab 0,2 mol baariumsulfaadi sadet. Kloriidist ja ammoniumioonidest jäääb lahusesse alles 0,4 mol NH_4Cl . Kuna pärast lahuse aurutamist veel kuumutati jääki ja ammoniumkloriid kuumutades laguneb, siis ei jäää portselankaussi suurt midagi.

1
12p



b) $\Delta H = 1 \cdot (-393,5 \text{ kJ/mol}) + 2 \cdot (-241,8 \text{ kJ/mol}) - 1 \cdot (-74,8 \text{ kJ/mol})$

$\Delta H = -802,3 \text{ kJ/mol}$ 2

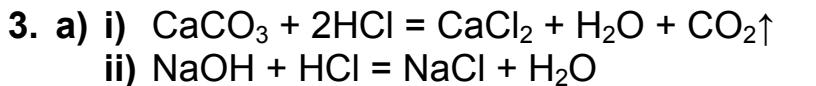
c) $n(\text{CH}_4) = \frac{1}{2} \cdot 502 \frac{\text{mg}}{\text{dm}^3} \cdot 250000 \text{ dm}^3 \cdot \frac{1 \text{ g}}{1000 \text{ mg}} \cdot \frac{1}{30 \text{ g/mol}} = 2090 \text{ mol}$ 2

$E = 802,3 \text{ kJ/mol} \cdot 2090 \text{ mol} \cdot \frac{1 \text{ GJ}}{10^6 \text{ kJ}} = 1,68 \text{ GJ}$ 1

$$d) E = \frac{1}{2} \cdot 12 \cdot 300 \text{ m}^3 \cdot 1000 \frac{\text{dm}^3}{\text{m}^3} \cdot \frac{1 \text{ mol}}{22,4 \text{ dm}^3} \cdot 802,3 \frac{\text{kJ}}{\text{mol}} \cdot \frac{1 \text{ GJ}}{10^6 \text{ kJ}}$$

$$E = 64,5 \text{ GJ} \approx 65 \text{ GJ}$$

2
8p



1

1

$$b) n(\text{HCl}) = \left(25,00 \text{ cm}^3 \cdot 1,101 \frac{\text{mol}}{\text{dm}^3} - 24,31 \text{ cm}^3 \cdot 0,3603 \frac{\text{mol}}{\text{dm}^3} \right) \frac{1 \text{ dm}^3}{1000 \text{ cm}^3}$$

$$n(\text{HCl}) = 0,01877 \text{ mol}$$

1

$$c) \%(\text{CaCO}_3) = \frac{0,01877 \text{ mol} \cdot \frac{1}{2} \cdot \frac{100,09 \text{ g}}{1 \text{ mol}}}{1,02 \text{ g}} \cdot 100\% = 92,1\%$$

1

d) Tiitrimisel tekkivate vigade vähendamiseks tuleks kogu eksperimenti veel vähemalt kaks korda korrrata ja lugeda töeliseks kaltsiumkarbonaadi sisalduseks kolme eksperimendi põhjal saadud kokkulangeva protsendilise sisalduse keskmist.

1

e) Kuna viga sõltub ainult arvutatud HCl moolide arvu erinevusest, siis:

$$n(\text{HCl}) = \left(25,00 \text{ cm}^3 \cdot 1,101 \frac{\text{mol}}{\text{dm}^3} - 24,37 \text{ cm}^3 \cdot 0,3603 \frac{\text{mol}}{\text{dm}^3} \right) \cdot \frac{1 \text{ dm}^3}{1000 \text{ cm}^3} = 0,01874 \text{ mol}$$

$$\%(\text{viga}) = \frac{0,01874 - 0,01877}{0,01877} \cdot 100 = -0,2\%$$

(“–“tähistab, et CaCO_3 sisaldus tuleb õigest väärustusest 0,2% väiksem)

2
7p

4. a) A – FeO (+arvutus) 0.5
 B – $\text{Fe}(\text{CO})_5$ (+arvutus) 0.5
 C – $\text{H}_2[\text{Fe}(\text{CO})_4]$ 0.5
 D – $\text{Na}_2[\text{Fe}(\text{CO})_4]$ 0.5
 E – $\text{Fe}_3(\text{CO})_{12}$ (+arvutus) 0.5
 F – $\text{Fe}_2(\text{CO})_9$ (+arvutus) 0.5
 G – $[\text{Fe}(\text{CO})_4]\text{I}_2$ 0.5
 H – FeI_2 (+arvutus) 0.5

Ühendite üldvalem Fe_xZ_y

$$A: 56x/0,78 = 56x + 16y \Rightarrow x = 1, y = 1 \quad \text{FeO}$$

$$B: 56x/0,29 = 56x + 28y \Rightarrow x = 1, y = 5 \quad \text{Fe}(\text{CO})_5$$

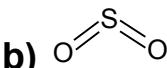
$$E: 56x/0,33 = 56x + 28y \Rightarrow x = 1, y = 4 \quad \text{Fe}_3(\text{CO})_{12}$$

$$F: 56x/0,31 = 56x + 28y \Rightarrow x = 2, y = 9 \quad \text{Fe}_2(\text{CO})_9$$

$$H: 56x/0,18 = 56x + 127y \Rightarrow x = 1, y = 2 \quad \text{FeI}_2$$

b) $\text{FeO} + \text{H}_2 = \text{Fe} + \text{H}_2\text{O}$	1
$\text{Fe} + 5\text{CO} = \text{Fe}(\text{CO})_5$	1
$\text{Fe}(\text{CO})_5 + \text{Ba}(\text{OH})_2 = \text{H}_2[\text{Fe}(\text{CO})_4] + \text{BaCO}_3$	1
$\text{H}_2[\text{Fe}(\text{CO})_4] + 2\text{Na} = \text{Na}_2[\text{Fe}(\text{CO})_4] + \text{H}_2$	1
$\text{H}_2[\text{Fe}(\text{CO})_4] + 2\text{I}_2 = [\text{Fe}(\text{CO})_4]\text{I}_2 + 2\text{HI}$	1
$\text{Fe}(\text{CO})_5 + \text{I}_2 = [\text{Fe}(\text{CO})_4]\text{I}_2 + \text{CO}$	1
$[\text{Fe}(\text{CO})_4]\text{I}_2 = \text{FeI}_2 + 4\text{CO}$	1
$2\text{Fe}(\text{CO})_5 = \text{Fe}_2(\text{CO})_9 + \text{CO}$	1
$3\text{Fe}_2(\text{CO})_9 = 2\text{Fe}_3(\text{CO})_{12} + 3\text{CO}$	1
	13p

5. a) Klaasi summaarne valem $\text{Na}_2\text{O}\cdot\text{CaO}\cdot 6\text{SiO}_2$ 1
- b) $\text{C}_x\text{H}_y\text{O}_z + (2x+y/2-z)/2\text{O}_2 = x\text{CO}_2 + y/2\text{H}_2\text{O}$ 1
- c) i) $n(\text{O}_2)_{\text{alg}} = \frac{1000 \text{ dm}^3}{22,4 \text{ dm}^3/\text{mol}} = 44,64 \text{ mol}$
- $n(\text{O}_2)_{\text{reag}} = 44,64 \text{ mol} \cdot 0,1344 = \mathbf{6,00 \text{ mol}}$ 1
- ii) $n(\text{gaasid}) = \frac{1000 \text{ dm}^3}{21,0 \text{ dm}^3/\text{mol}} = 47,6 \text{ mol}$
- $n(\text{gaasid})_{\text{nt}} = \frac{1000 \text{ dm}^3}{22,91 \text{ dm}^3/\text{mol}} = 43,64 \text{ mol}$ 1
- Seega $n(\text{H}_2\text{O}) = 47,6 \text{ mol} - 43,6 \text{ mol} = \mathbf{4,0 \text{ mol}}$ 1
- $y = 2 \cdot 4 = 8$
- iii) $n(\text{CO}_2) = n(\text{gaasid})_{\text{nt}} - [n(\text{O}_2)_{\text{alg}} - n(\text{O}_2)_{\text{reag}}] = \mathbf{5,00 \text{ mol}}$ 1
- seega $x = 5$
- d) $(10 + 4 - z)/2 = 6$, siit $z = 2$ MMA summaarne valem on $\text{C}_5\text{H}_8\text{O}_2$ **1**
7p

6. a) A on väävel (S_8) 1 E on vääveltrioksiid (SO_3) 1
- B on divesiniksulfiid (H_2S) 1 F on väävelhape (H_2SO_4) 1
- C on vääveldioksiid (SO_2) 1 G on kaltsiumsulfaat (CaSO_4) 1
- D on väävlishappe (H_2SO_3) 1
- b)  1
- c) $\text{S} + \text{H}_2 = \text{H}_2\text{S}$ 0,5
- $2\text{H}_2\text{S} + 3\text{O}_2 = 2\text{H}_2\text{O} + 2\text{SO}_2$ 1
- $2\text{H}_2\text{S} + \text{O}_2 = 2\text{H}_2\text{O} + 2\text{S}$ 1
- $\text{O}_2 + 2\text{SO}_2 = 2\text{SO}_3$ 1
- $\text{SO}_2 + \text{H}_2\text{O} = \text{H}_2\text{SO}_3$ 0,5
- $\text{SO}_3 + \text{H}_2\text{O} = \text{H}_2\text{SO}_4$ 0,5
- $\text{Ca}(\text{OH})_2 + \text{H}_2\text{SO}_4 = \text{CaSO}_4 \downarrow + 2\text{H}_2\text{O}$ 0,5
- 13p**