Exam 4A Chem 1142 Spring 2019

Na	me: KEY	
MUL	TIPLE CHOICE. [2.5 pts ea.] Record the best response on the scantron sheet. [50 pts total.]	
Assur	ne all solutions are aqueous and at a temperature of 25 °C, unless stated otherwise.	
Q1.	The second law of thermodynamics states that: A) The entropy of the reaction always increases B) The entropy of the universe always increases C) The entropy of the surroundings always increases D) The entropy of the system always increases	
Q2.	Which substance would we expect to have the greatest molar entropy at 25 °C? A) $NaF(s)$ B) $N_2(g)$ C) $H_2O(l)$ D) $C(s, graphite)$	
Q3.	Which chemical equation would most likely have $\Delta S^{\circ}_{rxn} < 0$? A) $CH_4(g) + 2O_2(g) \rightarrow CO_2(g) + 2H_2O(g)$ B) $Na(s) + \frac{1}{2} Br_2(l) \rightarrow NaBr(s)$ C) $H_2(g) + \frac{1}{2} O_2(g) \rightarrow H_2O(g)$ D) $C(s, graphite) + O_2(g) \rightarrow CO_2(g)$	
Q4.	A chemical reaction has $\Delta H_{\rm rxn} = -5.00$ kP at 250 K. What will the entropy change of the surroundings by $\Delta H = 20 \text{J/K}$ B) -20J/K C) $+1300 \text{kJ} \cdot \text{K}$ D) $-1300 \text{kJ} \cdot \text{K}$ An endothermic reaction has $\Delta S_{\rm rxn} > 0$. What can you say about its spontaneity? $\Delta G = \Delta H - T\Delta S$	ę
Q5.	An endothermic reaction has $\Delta S_{rxn} > 0$. What can you say about its spontaneity? A) It will always be spontaneous B) It will always be non-spontaneous C) It will be spontaneous at low temperatures, but non-spontaneous at high temperatures	

5=K. InW

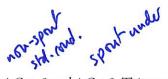
(D) It will be non-spontaneous at low temperatures, but spontaneous at high temperatures

C) When it is a substance under 1 atm pressure at its normal melting point

Q6. When will the entropy of a substance be zero?

(D) When it is a perfect crystal at 0 K

A) When it is an ion at a molar concentration of 1 M B) When it is an element in its standard state at 25 °C



-	1	-	A	- 1	. 1	reaction	1	10		α	1	10	6 02	0	TT .		. 1
)	/	A	C	nemical	reaction	has	// (7	· >	()	and	/\(\)	<	()	Inis	means	that:

- A) It is spontaneous under standard conditions, but non-spontaneous under current conditions
- (B) It is non-spontaneous under standard conditions, but spontaneous under current conditions
- C) It will always be non-spontaneous under any condition
- D) It will always be spontaneous under any condition

Q8. A chemical reaction with $\Delta G^{\circ} \ll 0$ will likely have an equilibrium constant, K, such that: AZB KVP



- B) $K \ll 1$
- C) K=1
- D) K=0
- Q9. Which of the following chemical equations corresponds to the standard Gibbs energy of formation of $NH_3(g)$?
 - A) $NH_3(g) + O_2(g) \rightarrow NO_2(g) + \frac{3}{2} H_2(g)$
 - B) $2NH_3(g) \rightarrow N_2(g) + 3H_2(g)$
 - C) $\frac{1}{2}$ N₂(g) + $\frac{3}{2}$ H₂(g) \rightarrow NH₃(g) D) NH₂⁻(aq) + H⁺(aq) \rightarrow NH₃(g)

plement. ___ NH3/q)

reduced : sain es.

- Q10. Which pair of chemical equations, when coupled (added), give rise to a spontaneous reaction?
 - (i) $A \longrightarrow B$ $\Delta G^{\circ} = -20 \text{ kJ}$
 - $B \rightarrow C$ $\Delta G^{\circ} = +25 \text{ kJ}$ (ii)
 - $A \rightarrow C$ $\Delta G^{\circ} = +10 \text{ kJ}$ (iii)
 - (iv) $B \longrightarrow D$ $\Delta G^{\circ} = -15 \text{ kJ}$
 - A) (ii) & (iii) **
 - B) (ii) & (iv) +10
 - C) (i) & (ii) 15
 - (D) (i) & (iii) 10
- Q11. Which of the following half-reactions is properly balanced?
 - A) $Br_2(1) \rightarrow 2Br^-(aq)$
 - B) $2H^{+}(aq) + e^{-} + IO^{+}(aq) \longrightarrow H_{2}O(1) + \frac{1}{2}I_{2}(s)$
 - C) $2e^- + 2Cl^-(aq) \rightarrow Cl_2(g) \times$
 - (D) $2e^- + H^+(aq) + Pb(OH)_3^+ \rightarrow Pb(OH)_2 + H_2O(I)$
- Q12. Where does reduction take place in a voltaic (galvanic) cell?
 - A) The salt bridge
 - B) The voltmeter
- red (at
- C) The cathode
- an ot
- D) The anode
- Q13. Given the following two electrode potentials:

$$Cu^{2+} + 2e^- \rightarrow Cu$$

$$E^{\circ} = +0.34 \text{ V}$$

$$\mathbb{Z}n^{2+} + 2e^{-} \longrightarrow \mathbb{Z}n$$

$$E^{\circ} = -0.76 \text{ V}$$

The best reducing agent would be:

- A) Cu2+
- B) Cu
- $C) Zn^{2+}$
- D) Zn

(2=+ Mg2+ -) Mg) x3 2(r + 3Mg2+ 6=5 2C13+ + 5Mg (at nd. conde, ox Q14. What is the cell reaction for the voltaic cell: $Cr(s)|Cr^{3+}(aq)||Mg^{2+}(aq)|Mg(s)$ A) $2Cr(s) + 2Cr^{3+}(aq) \rightarrow 3Mg^{2+}(aq) + 3Mg(s)$ (B) $2Cr(s) + 3Mg^{2+}(aq) \rightarrow 2Cr^{3+}(aq) + 3Mg(s)$ C) $2Cr^{3+}(aq) + 3Mg(s) \rightarrow 2Cr(s) + 3Mg^{2+}(aq)$ D) $3Mg^{2+}(aq) + 2Cr^{3+}(aq) \rightarrow 3Mg(s) + 2Cr(s)$ Q15. Which of the following statements is true about a voltaic (galvanic) cell that has an Eo value of -0.50 V? (i) The reaction is spontaneous X (ii) $K > 1 \times$ (iii) $\Delta G^{\circ} > 0$ A) (i) & (ii) B) (iii) only C) (ii) & (iii) D) (i) only Q16. How would E_{cell} change if the concentration β f Ag⁺(aq) was increased in the following voltaic (galvanic) cell? $Ag(s)|Ag^{+}(aq)||Au^{3+}(aq)|Au(s)$ 3=+ An -) An A) It would increase B) It would not change D) There is not enough information to decide 3450) + Au(s) Q17. The reaction MgCl₂ \rightarrow Mg + Cl₂ has a cell potential (E_{cell}) of -3.5 V. This means that: A) The reaction is spontaneous and will produce 3.5 V in a cell The reaction is non-spontaneous, and will produce 3.5 V in a cell C) The reaction can be driven by applying a voltage of > 3.5 V 🗸 D) The reaction is spontaneous, but can be stopped by applying a voltage of $< -3.5 \text{ V} \checkmark$ $Q = I \cdot t$ = 12.0 \(\text{C} \) \(\text{24.0 min} \) \(\text{1 min} \)
= 17,280C Q18. How many moles of electrons are produced when 12.0 A flows for 24.0 min? A) 288 mol B) 5.58 mol C) 0.500 mol (D) 0.179 mol Q19. Copper can be plated out of a solution containing Cu²⁺ according to the half-reaction: $Cu^{2+}(aq) + 2e^{-} \rightarrow Cu(s)$ What mass of copper is formed when 15.0 A of current flows for 10.0 min? 0.093 mole + 1 (n 355g(n = 2.969 (n A) 2.96 g B) 5.92 g C) 8.15 g D) 12.6 g Q20. Which electrode is assigned a standard electrode potential of 0.00 V? A) The standard hydrogen electrode

B) The standard pH electrode
 C) The standard acidic electrode
 D) The standard platinum electrode

Short Response.

Show ALL work to receive credit.

<u>Using the half-reaction method</u>, balance the following redox reaction that occurs in **basic** solution:

(o) (4)(-1) (4) (-1)(+1)

$$AI + NO_3^- \rightarrow AI(OH)_4^- + NH_3$$

Be sure to identify oxidation states of each atom as part of your answer. Clearly indicate whether

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Q22. [13 pts.]

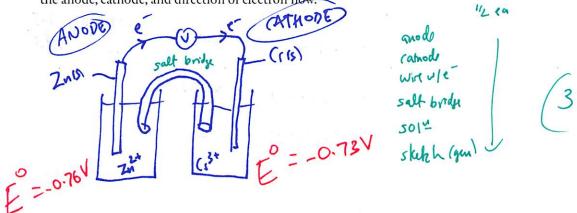
Consider the voltaic cell

 $|Zn(s)|Zn^{2+}(aq)||Cr^{3+}(aq)|Cr(s)|$

(i) Write the half-cell reactions and the overall cell reaction.

(Zn(s) -> Zn'(as) +2e-) x3 (3e-+ Cr3+(as) -> Cr(s)) +2 3Zn(s) +2(r3+(as) 6e-) 3Zn'(as) + 2Cr(s).

(ii) Make a sketch of this voltaic cell and label it. At a minimum, be sure to include labels showing the anode, cathode, and direction of electron flow.



(iii) Calculate
$$E_{cell}$$
. $E_{cell} = E_{calhode} - E_{aude} = (-0.78V) - (-0.76V) = +0.03V$

(iv) What is
$$E_{\text{cell}}$$
 if $[Zn^{2+}] = [Cr^{3+}] = 0.10$ M? Assume a temperature of 298 K.
$$E = E^{\circ} - \frac{RT}{\kappa F} \cdot \ln Q \qquad \qquad Q = \frac{[Zn^{3+}]^2}{[Cr^{3+}]^2} = \frac{0.10^3}{0.10^3} = 0.10$$

Q23. [12 pts.] For the chemical reaction:

$$H_2O(g) + Cl_2(g) \longrightarrow 2HCl(aq) + \frac{1}{2}O_2(g)$$

(i) Calculate ΔH° , ΔS° , and ΔG° at 25 °C. (Hint: be sure to look at the data table at the bottom of this question!)

$$\Delta H^{\circ} = -92.6 \text{ KJ}$$

$$\Delta S^{\circ} = -196.3 \frac{7}{\text{K}}$$

$$\Delta G^{\circ} = \Delta H^{\circ} - T \Delta S^{\circ}$$

$$= -92.6 \text{ KJ} - 298.15 \text{ K} \times \frac{-196.37}{\text{K}} \times \frac{\text{KJ}}{10^{3} \text{J}}$$

$$= -92.6 \text{ KJ} + 58.5 \text{ KJ}$$

$$= -34.1 \text{ KJ}$$

(ii) Use this information to calculate the equilibrium constant at 25 °C.

$$\Delta G^{\circ} = -\frac{1}{2} \ln K$$

$$\Rightarrow \ln K = -\frac{\Delta G^{\circ}}{2} = +\frac{34,100}{8.3145} = \frac{13.7}{100.100} = 13.7$$

$$\Rightarrow K = e^{13.7} = 9.3 \times 10^{5} = 1$$

(iii) Predict how the spontaneity of this reaction will change (if at all!) at low vs. high temperature. Be sure to explain your answer!

as TT, -TAS dominate, making No +ve

Substance	$\Delta H_{\rm f}^{o} ({\rm kJ/mol})$	So (J/mol·K)
H ₂ O(g)	-241.8	188.8
Cl ₂ (g)	0	223.1
HCl(aq)	-167.2	56.5
O ₂ (g)	0	205.2

Q24. [13 pts.] Consider the reaction:

$$CO(g) + 2H_2(g) \rightleftharpoons CH_3OH(g)$$

Calculate ΔG_{rxn} at 25 °C under each of the following conditions: (Hint: be sure to look at the data table at the bottom of this question!)

i) standard conditions
$$\Delta G = \sum_{p} \Lambda_{p} \cdot \Delta G_{p}(p) - \sum_{p} \Lambda_{r} \cdot \Delta G_{p}(R) = \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] - \left[|mol \times -137 \cdot 2 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] - \left[|mol \times -137 \cdot 2 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] - \left[|mol \times -137 \cdot 2 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot 3 \frac{R-7}{mol} \right] + \left[|mol \times -162 \cdot$$

ii) at equilibrium

iii) under conditions of $p_{CH_3OH} = 1.5$ atm, and $p_{CO} = p_{H_2} = 0.020$ atm.

$$\Delta G = \Delta G + RT \ln Q$$

$$= -25.1 \frac{kJ}{MO!} + 8.3145 \frac{J}{1000J} + 298.15 \frac{k}{4} \ln \frac{P_{CH_{J}OM}}{P_{ro} + P_{L_{Z}}^{2}}$$

$$= -25.1 \frac{kJ}{MO!} + 2.478968 \frac{kJ}{MO!} \ln \frac{1.5}{0.020 \times 0.020^{2}}$$

$$= -25.1 \frac{kJ}{MO!} + 30.1 \frac{kJ}{MO!} = \frac{1.5}{MO!}$$

Substance	$\Delta G_{\rm f}^{\rm o} ({ m kJ/mol})$
CH ₃ OH(g)	-162.3
$H_2(g)$	0
CO(g)	-137.2

IA	IIA			Perio	dic T	able o	of the	Elem	ents			IIIA	IVA	VA	VIA	VIIA	VIIIA
- 1	1																2
н																	He
1,008	2											13	14	15	16	17	4.003
3	4											5	6	7	8	9	10
Li	Be											В	C	N	0	F	Ne
6.941	9.012											10.81	12.01	14.01	16.00	19.00	20.18
11	12											13	14	15	16	17	18
Na	Mg											Al	Si	P	S	CI	Ar
22.99	24.31	3		5	- 6	7	8	9	10	11	12	26.98	28.09	30.97	32.07	35.45	39.95
19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36
K	Ca	Sc	Ti	V	Cr	Mn	Fe	Co	Ni	Cu	Zn	Ga	Ge	As	Se	Br	Kr
39.10	40.08	44.96	47.87	50.94	52.00	54.94	55.85	58.93	58.69	63.55	65.39	69.72	72.61	74.92160	78.96	79.90	83,60
37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54
Rb	Sr	Y	Zr	Nb	Mo	Tc	Ru	Rh	Pd	Ag	Cd	l ln	Sn	Sb	Te	1	Xe
85.47	87.62	88.91	91.22	92.91	95.94	(98)	101.1	102.9	106.4	107.9	112.4	114.8	118.7	121.8	127.60	126.9	131.3
55	56	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86
Cs	Ba*	Lu	Hf	Та	W	Re	Os	Ir	Pt	Au	Hg	TI	Pb	Bi	Po	At	Rn
132.9	137,3	175.0	178.5	180.9	183.8	186.2	190.2	192.2	195.1	197.0	200.6	204.4	207.2	209.0	[210]	[210]	[222]
87	68	103	104	105	106	107	108	109	110	111	112	113	114	115	116	117	118
Fr	Ra**	Lr	Rf	Db	Sg	Bh	Hs	Mt		l		İ					
[223]	[226]	(262)	[261]	[262]	[266]	[264]	[265]	[268]	[269]	[272]	[277]	<u> </u>	[285]		[269]		[293]
																,	
		57	58	59	60	61	62	63	64	65	66	67	68	69	70		
	•	La	Ce	Pr	Nd	Pm	Sm	Eu	Gd	Tb	Dy	Ho	Er	Tm	Yb		
		138.9	140.1	140.9	144.2	[145]	150.4	152.0	157.3	158,9	162.50	164.9	167.3	168.9	173.0		
	**	89	90	91	92	93	94	95	96	97	98	99	100	101	102		
	**	Ac	Th	Pa	U	Np	Pu	Am	Cm	Bk	Cf	Es	Fm	Md	No		
		[227]	232.0	231.0	238.0	[237]	[244]	[243]	[247]	[247]	[251]	[252]	[257]	[258]	[259]	J	

 $N_{\rm A} = 6.022 \times 10^{23} \, \rm mol^{-1}$

$$R = 8.3145 \frac{J}{\text{mol} \cdot \text{K}} = 0.08206 \frac{\text{atm} \cdot \text{L}}{\text{mol} \cdot \text{K}}$$

$$M_1V_1=M_2V_2$$

$$\Delta G = -nFE_{\text{cell}}$$

$$\Delta G^{\circ} = -nFE^{\circ}_{cell}$$

$$E_{\text{cell}}^{\prime\prime} = \frac{RT}{nF} \ln K$$

$$E_{\text{cell}} = E_{\text{cell}}^{\circ} - \frac{RT}{nF} \ln Q$$

$$E^{\circ}_{\text{cell}} = E^{\circ}_{\text{cathode}} - E^{\circ}_{\text{anode}}$$
 $F = 96,500 \text{ C/mol e}^{-}$

$$F = 96.500 \text{ C/mol e}^{-1}$$

1 V = 1 J/C

$$R = 8.3145 \text{ J} \cdot \text{mol}^{-1} \cdot \text{K}_{\cdot}^{-1}$$

$$Q(\text{charge}) = I \cdot t$$

$$\Delta G = \Delta H - T \Delta S$$

$$\Delta S_{\rm surr} = q_{\rm surr}/T$$

$$Q ext{ (charge)} = I \cdot t$$
 $\Delta G = \Delta H - T \Delta S$
 $\Delta G = \Delta G^{\circ} + RT \ln Q$ $\Delta G^{\circ} = -RT \ln K$

$$\Delta G^{\circ} = -RT \ln K$$

Standard reduction potentials

		E°(V)
	2 F⁻(aq)	2.87
$H_2O_2(aq) + 2 H^+(aq) + 2 e^-$	2 H ₂ O(I)	1.78
$PbO_2(s) + 4 H^+(aq) + SO_4^{2-}(aq) + 2 e^-$	\longrightarrow PbSO ₄ (s) + 2 H ₂ O(I)	1.69
$MnO_4^-(aq) + 4 H^+(aq) + 3 e^-$	\longrightarrow MnO ₂ (s) + 2 H ₂ O(l)	1.68
$MnO_4^-(aq) + 8 H^+(aq) + 5 e^-$	\longrightarrow Mn ²⁺ (aq) + 4 H ₂ O(I)	1.51
Au ³⁺ (aq) + 3 e ⁻	→ Au(s)	1.50
$PbO_2(s) + 4 H^+(aq) + 2 e^-$	$\longrightarrow Pb^{2+}(aq) + 2 H_2O(l)$	1.46
Cl ₂ (g) + 2 e ⁻	2 Cl⁻(aq)	1.36
$Cr_2O_7^{2-}(aq) + 14 H^*(aq) + 6 e^-$	\longrightarrow 2 Cr ³⁺ (aq) + 7 H ₂ O(I)	1.33
$O_2(g) + 4 H^*(aq) + 4 e^-$	→ 2 H ₂ O(I)	1.23
MnO ₂ (s) + 4 H*(aq) + 2 e	\longrightarrow Mn ² *(aq) + 2 H ₂ O(I)	1.21
O ₃ ~(aq) + 6 H*(aq) + 5 e~	$\longrightarrow \frac{1}{2}I_2(aq) + 3 H_2O(I)$	1.20
3r ₂ (/) + 2 e ⁻	2 Br"(aq)	1.09
/O ₂ *(aq) + 2 H*(aq) + e	VO ²⁺ (aq) + H ₂ O(I)	1.00
NO ₃ ⁻ (aq) + 4 H ⁺ (aq) + 3 e ⁻	\longrightarrow NO(g) + 2 H ₂ O(I)	0.96
CIO ₂ (g) + e ⁻	→ CIO ₂ (aq)	0.95
Ag+(aq) + e-	→ Ag(s)	0.80
Fe ³⁺ (aq) + e	Fe ²⁺ (aq)	0.77
$O_2(g) + 2 H^*(aq) + 2 e^-$	$\longrightarrow H_2O_2(aq)$	0.70
MnO ₄ (aq) + e	$\longrightarrow Mn0_4^{2^-}(aq)$	0.56
₂ (s) + 2 e ⁻	2 l⁻(aq)	0.54
Cu ⁴ (aq) + e ⁻	Cu(s)	0.52
$O_2(g) + 2 H_2O(I) + 4 e^-$	→ 4 OH~(aq)	0.40
Cu ²⁺ (aq) + 2 e ⁻	Cu(s)	0.34
60_4^2 (aq) + 4 H ⁺ (aq) + 2 e	$\longrightarrow H_2SO_3(aq) + H_2O(l)$	0.20
Cu ²⁺ (aq) + e ⁻	→ Cu ⁺ (aq)	0.16
Sn ⁴⁺ (aq) + 2 e	Sn ²⁺ (aq)	0.15
2 H*(aq) + 2 e		0
e³+(aq) + 3 e ⁻	→ Fe(s)	-0.036
² b ²⁺ (aq) + 2 e ⁻	> Pb(s)	-0.13
Sn ²⁺ (aq) + 2 e	Sn(s)	-0.14
Ni ²⁺ (aq) + 2 e ⁻	→ Ni(s)	-0.23
Cd ²⁺ (aq) + 2 e ⁻	→ Cd(s)	-0.40
$e^{2}+(aq)+2e^{-}$	Fe(s)	-0.45
Cr³+(aq) + e⁻	Cr ²⁺ (aq)	-0,50
Cr ³⁺ (aq) + 3 e ⁻		-0.73
$2^{-2}(aq) + 2e^{-}$	— Zn(s)	-0.76
2 H ₂ O(I) + 2 e ⁻		-0.83
Mn ²⁺ (aq) + 2 e	→ Mn(s)	-1.18
$N^{3+}(aq) + 3e^{-}$	→ Al(s)	-1.66
$Mg^{2+}(aq) + 2e^{-}$	→ Mg(s)	-2.37
Va*(aq) + e	→ Na(s)	-2.71
Ca ² *(aq) + 2 e ⁻	→ Ca(s)	-2.76
$3a^{2}+(aq)+2e^{-}$	→ Ba(s)	-2.90
(+(aq) + e	→ K(s)	-2.92