

**DEPARTMENT OF CHEMISTRY
UNIVERSITY OF SWAZILAND**

C304

ANALYTICAL CHEMISTRY I I

DECEMBER 2013 FINAL EXAMINATION

Time Allowed:

Three (3) Hours

Instructions:

1. This examination has six (6) questions and one (1) data sheet. The total number of pages is five (5), including this page.
2. Answer any four (4) questions fully; diagrams should be clear, large and properly labeled. Marks will be deducted for improper units and lack of procedural steps in calculations.
3. Each question is worth 25 marks.

Special Requirements

1. Data sheet.
2. Graph paper.

YOU ARE NOT SUPPOSED TO OPEN THIS PAPER UNTIL PERMISSION TO DO SO HAS BEEN GIVEN BY THE CHIEF INVIGILATOR.

QUESTION 1 [25]

- a) A spectroscopy source emits 1500cm^{-1} radiation.
- What type of transition would it induce in a sample? [1]
 - What is its wavelength in μm ? [1]
 - What is its energy in J? [1]
 - What is its frequency in Hz? [1]
 - What is its energy in eV? [1]
- b) Explain the difference in sample placement between IR and uv-visible spectroscopy [4]
- c) Gratings have a very good resolving power in spectroscopy.
- Physically, how does a grating look like (2)
 - Use equations to explain how a grating works (3)
 - Calculate the second order resolving power of a grating which is 5cm long with 1180 lines per mm (3)
- d) One of the applications of GC is the separation of benzene from its mixture with cyclohexane, followed by quantification of the benzene.
- In GC, use diagrams to explain what is meant by longitudinal diffusion [2]
 - State the equation that relates longitudinal diffusion in a packed chromatographic bed, to linear velocity [2]
 - In GC, use diagrams to explain what is meant by Eddy diffusion [2]
 - State the equation that relates bandbroadening due to Eddy Diffusion, to linear velocity [2]

QUESTION 2 [25]

- a) State Beer's Law as applied to spectroscopy, and explain all terms appearing in it [3]
- b) Stray light is problematic in spectroscopy
- What is meant by "stray light" in spectroscopy? [1]
 - Use equations to explain why stray light leads to negative deviations from Beer's Law [3]
 - How is stray light eliminated in spectroscopy? [1]
- c) Draw and label a vacuum phototube and explain how it works. [3]
- d) The mobile phase is a critical component in chromatography.
- Explain the role of the mobile phase in gas chromatography [1]
 - List and discuss any two (2) desirable properties of a mobile phase in gas chromatography [4]
 - Explain how silanol groups are deactivated in chromatography [4]
- e) Use diagrams to explain how UV-visible detection of ink isomers is carried out after separation in an HPLC instrument. [5]

Question 3 [25]

- a) Use diagrams to explain why it was not possible to perform atomic absorption spectroscopy using traditional broadband sources prior to the invention of the hollow cathode lamp [4]
- b) Explain why in atomic absorption spectroscopy
 - (i) 1000 ppm La is added to all solutions in the analysis of Pb [2]
 - (ii) Excess EDTA is added to all solutions in the analysis of Ca [2]
- c) Explain the role of background correction in flame atomic absorption spectroscopy, and explain using diagrams how this is achieved [4]
- d) Chromatography is semi-batch differential migration, phase distribution technique. Explain the meaning of this phrase. [3]
- e) Draw the Van Deemter Plot for gas chromatography and explain how it is used to optimize linear velocity of the carrier gas. [3]
- f) What is the major difference between a typical Gc Van Deemter Plot and a typical LC one ? [2]
- g) Use diagrams to explain
 - (i) Why N₂ is preferred over He as carrier gas in chromatography [2]
 - (ii) The effect liquid loading of stationary phase on bandbroadening [3]

Question 4 [25]

- a) With the aid of a diagram briefly describe the IPC torch [3]
- b) Describe the path taken by a Ca atom in a sample containing CaCl₂ solution from aspiration stage to atomization stage in ICP [4]
- c) Discuss two major advantages of ICP over flame or graphite furnace atomic absorption. [2]
- d) The ICP has many excitation lines which must be resolved using a grating.
 - i) Physically, how does a grating look like? [1]
 - ii) Use equations to explain how it works as a monochromator [3]
- e) With the aid of a diagram, briefly but informatively explain the function of each of the following detectors
 - i) Electron Capture Detector [4]
 - ii) Flame Ionization Detector. [4]
 - iii) Thermal Conductivity Detector [4]

Question 5 [25]

- a) State Beer's Law and explain all terms appearing in it [3]
- b) Explain how an unstable power supply gives rise to deviation in Beer's Law, and use a diagram to explain how this deviation is eliminated [3]
- c) Use diagrams to explain how UV-visible spectroscopy is used to determine stoichiometry by the Job's Method [3]

- d) In liquid chromatography, two solvent reservoirs are usually used. Explain the reason for this. [2]
- e) In gas chromatography, dual columns are often used simultaneously. Explain the reason for this. [2]
- f) One of the applications of GC is the separation of benzene from its mixture with cyclohexane, followed by quantification of the benzene.
 - (i) In GC, what is meant by lateral diffusion? [3]
 - (ii) State the equation that relates resistance to mass transfer in the mobile phase to bandbroadening [3]
 - (iii) In GC, what is meant by resistance to mass transfer in the stationary phase? [3]
 - (iv) State the equation that relates resistance to mass transfer in the stationary phase to bandbroadening [3]

Question 6 [25]

- a) Prisms are used as monochromatours for spectroscopy
 - i) Draw the prism (1)
 - ii) Use equations to explain how the prism works (3)
 - iii) Draw and label the Bunsen arrangement of optical components in a spectrometer (3)
- b) The Nernst Glower is a useful source of radiation in infrared spectroscopy.
 - i) Describe the Nernst Glower as used in IR spectroscopy. [1]
 - ii) Which of the molecules oxygen and hydrogen chloride is IR active and why? [2]
 - iii) Why is it not possible to carry out quantitative analyses on dispersive IR? [3]
- c) Nebulization is a very wasteful approach to atomization.
 - i) What does the term "nebulization" mean? [1]
 - ii) Use diagrams to explain how nebulization is carried out in atomic spectroscopy [3]
 - iii) Use your answer in (c) ii above to explain why nebulization is considered inefficient [2]
- d) Bandbroadening is important for peak resolution in HPLC.
 - i) Use a drawing to explain the importance of linear velocity on HETP [2]
 - ii) On this drawing, indicate the optimum linear velocity [1]
 - iii) Use diagrams to explain the phenomenon of "race track effect", how it affects bandbroadening, and how it is eliminated. [3]

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A value in brackets denotes the mass number of the longest lived or best known isotope.

$\text{Ag}(\text{CN})_2^-$	5×10^{20}
$\text{Ag}(\text{NH}_3)_2^+$	1.6×10^7
$\text{Ag}(\text{S}_2\text{O}_3)_2^{3-}$	4.7×10^{13}
$\text{Al}(\text{OH})_4^-$	1.0×10^{33}
$\text{Ca}(\text{EDTA}) =$	1.0×10^{11}
$\text{Cd}(\text{CN})_4 =$	8.3×10^{17}
$\text{Cd}(\text{NH}_3)_4^{++}$	5.5×10^6
$\text{Co}(\text{NH}_3)_6^{+3}$	2×10^{35}
$\text{Cr}(\text{OH})_4^-$	4×10^{28}
$\text{Cu}(\text{CN})_4^{-3}$	1×10^{23}
$\text{Cu}(\text{NH}_3)_4^{++}$	1.2×10^{11}
$\text{Fe}(\text{CN})_6^{-3}$	4.0×10^{43}
$\text{Fe}(\text{CN})_6^{-4}$	2.5×10^{35}
$\text{Fe}(\text{SCN})^{++}$	1.0×10^3
$\text{HgCl}_4 =$	1.3×10^{15}
$\text{Hg}(\text{CN})_4 =$	8.3×10^{38}
$\text{Hg}(\text{SCN})_4 =$	5.0×10^{20}
$\text{HgI}_4 =$	6.3×10^{29}
$\text{Mg}(\text{EDTA}) =$	1.3×10^9
$\text{Ni}(\text{NH}_3)_4^{++}$	4.7×10^7
$\text{Pb}(\text{OH})_3^-$	7.9×10^{13}
$\text{Zn}(\text{CN})_4 =$	4.2×10^{16}
$\text{Zn}(\text{NH}_3)_4^{++}$	7.8×10^8
$\text{Zn}(\text{OH})_4 =$	6.3×10^{14}

▲ Actinide

58 Ce 140.12	59 Pr 140.9077	60 Nd 144.24	61 Pm (145)	62 Sm 150.36	63 Eu 151.96	64 Gd 157.25	65 Tb 158.9254	66 Dy 162.50	67 Ho 164.9304	68 Er 167.26	69 Tm 168.9342	70 Yb 173.04	71 Lu 174.967
90 Th 232.0381	91 Pa 231.0359	92 U 238.0289	93 Np 237.0482	94 Pu (244)	95 Am (243)	96 Cm (247)	97 Bk (247)	98 Cf (251)	99 Es (252)	100 Fm (257)	101 Md (258)	102 No (259)	103 Lr (260)

5. FIRST IONIZATION ENERGIES, e.v.

1A	2A	14										3A	4A	5A	6A	7A	25	
5.4	9.3											8.3	11	15	14	17	22	
5.1	7.6	3B	4B	5B	6B	7B	8B				1B	2B	6.0	8.1	11	10	13	16
4.3	6.1	6.6	6.8	6.7	6.8	7.4	7.9	7.9	7.6	7.7	9.4	6.0	8.1	10	9.8	12	14	16
4.2	5.7	6.6	7.0	6.8	7.2		7.5	7.7	8.3	7.6	9.0	5.8	7.3	8.6	9.0	10	12	14
3.9	3.2	5.0 6.9	5.5	6	8.0	7.9	8.7	9.2	9.0	9.2	10	6.1	7.4	8			11	

1A	2A											3A	4A	5A	6A	7A	
1.0	1.5											2.0	2.5	3.0	3.5	4.0	
0.9	1.2	3B	4B	5B	6B	7B	8B	1B	2B	1.5	1.8	2.1	2.5	3.0			
0.8	1.0	1.3	1.5	1.6	1.6	1.5	1.8	1.8	1.8	1.9	1.6	1.8	1.8	2.0	2.4	2.8	
0.8	1.0	1.2	1.4	1.6	1.8	1.9	2.2	2.2	2.2	1.9	1.7	1.7	1.8	1.9	2.1	2.5	
0.7	0.9	1.1	1.3	1.5	1.7	1.9	2.2	2.2	2.2	2.4	1.9	1.8	1.8	1.9	2.0	2.2	

[illegible]

9. LATTICE ENERGIES

Li ⁺	60	Sr ⁺²	113	S ⁻²	184	(All negative)				kJ/mol	
Na ⁺	95	Ba ⁺²	135	Se ⁻²	198	F	Cl	Br	I		
K ⁺	133	B ⁺³	20	Te ⁻²	221	Li	1030	840	781	718	
Rb ⁺	148	Al ⁺³	50	F ⁻	136	Na	914	770	728	681	
Be ⁺²	31	N ⁺³	171	Cl ⁻	181	K	812	701	671	632	617
Mg ⁺²	65	P ⁺³	212	Br ⁻	195	Rb	780	682	654	612	
Ca ⁺²	99	O ⁻²	140	I ⁻	216	Cs	744	630	613	585	

H ³	12.3 years	K ⁴⁰	1.28 × 10 ⁹ y	I ¹³¹	8.1 days
F ²⁰	11.4 secs	Ca ⁴⁵	165 days	Cs ¹³⁷	30 years
C ¹⁴	5730 years	Fe ⁵⁹	45 days	Au ¹⁹⁸	2.69 days
Na ²⁴	15.0 hours	Co ⁶⁰	5.26 y	Ra ²²⁶	1620 yrs.
P ³²	14.3 days	Br ⁸²	35.5 hours	U ²³⁵	7.1 × 10 ⁸ y
S ³⁵	88 days	Sr ⁹⁰	28 years	U ²³⁸	4.51 × 10 ⁹ y
Cl ³⁶	3.1 × 10 ⁵ y	I ¹²⁹	1.7 × 10 ⁷ y	Pu ²³⁰	24,400 y

AgBr	4×10^{-13}	BaC ₂ O ₄	2×10^{-8}	KClO ₄	2×10^{-2}
Ag ₂ CO ₃	6×10^{-12}	BaSO ₄	1×10^{-10}	MgCO ₃	1×10^{-5}
AgCl	1×10^{-10}	CaCO ₃	5×10^{-9}	MgC ₂ O ₄	9×10^{-5}
Ag ₂ CrO ₄	2×10^{-12}	CaF ₂	4×10^{-11}	MgNH ₄ PO ₄	2×10^{-13}
Ag[Ag(CN) ₂]	4×10^{-12}	CaC ₂ O ₄	2×10^{-9}	Mg(OH) ₂	1×10^{-11}
AgI	1×10^{-16}	CdS	1×10^{-28}	MnS	1×10^{-15}
Ag ₃ PO ₄	1×10^{-19}	Cu(OH) ₂	2×10^{-20}	PbCrO ₄	2×10^{-14}
Ag ₂ S	1×10^{-50}	CuS	1×10^{-36}	PbS	1×10^{-28}
AgCNS	1×10^{-12}	Fe(OH) ₃	1×10^{-36}	PbSO ₄	2×10^{-8}
Al(OH) ₃	2×10^{-32}	Hg ₂ Br ₂	3×10^{-23}	SrCrO ₄	4×10^{-5}
BaCO ₃	5×10^{-9}	Hg ₂ Cl ₂	6×10^{-19}	Zn(OH) ₂	3.6×10^{-16}
BaCrO ₄	1×10^{-10}	HgS	1×10^{-52}	ZnS	1×10^{-24}

11. ACID-BASE INDICATORS AT 25°C

Indicator	pH range	pK _{in}	Acid	Base
Thymol blue	1.2-2.8	1.6	red	yellow
Methyl yellow	2.9-4.0	3.3	red	yellow
Methyl orange	3.1-4.4	4.2	red	yellow
Bromocresol green	3.8-5.4	4.7	yellow	blue
Methyl red	4.2-6.2	5.0	red	yellow
Chlorophenol red	4.8-6.4	6.0	yellow	red
Bromothymol blue	6.0-7.6	7.1	yellow	blue
Phenol red	6.4-8.0	7.4	yellow	red
Cresol purple	7.4-9.0	8.3	yellow	purple
Thymol blue	8.0-9.6	8.9	yellow	blue
Phenolphthalein	8.0-9.8	9.7	colorless	red
Thymolphthalein	9.3-10.5	9.9	colorless	blue

14. DATA REJECTION—Q TABLE

n	Q ₉₀	n	Q ₉₀	n	Q ₉₀
3	0.94	6	0.56	9	0.44
4	0.76	7	0.51	10	0.41
5	0.64	8	0.47		

19. t TABLE

D.F.	t ₅₀	t ₉₀	t ₉₅	t ₉₉
1	1.0	6.3	13	64
2	0.82	2.9	4.3	9.9
3	0.76	2.35	3.2	5.8
4	0.74	2.13	2.8	4.6
5	0.73	2.02	2.57	4.0
6	0.72	1.94	2.45	3.7
7	0.71	1.90	2.36	3.5
8	0.71	1.86	2.31	3.36
9	0.70	1.83	2.26	3.25
10	0.70	1.81	2.23	3.17
20	0.69	1.72	2.09	2.84
30	0.68	1.70	2.04	2.75
∞	0.67	1.64	1.96	2.58

15. Bond Enthalpies

kJ mol ⁻¹ at 25°C (i.e. Bond Energies)						
Single	O	N	C	S	F	Cl
H	463	391	413	368	563	432
C	358	305	346	272	489	328
N	222	163	MISC.	275	192	
S-S	251	H-H	436	C=C	615	
S-F	327	N=N	946	C≡C	812	
S-Cl	271	N=O	607	C=O	749	

12. ELECTRODE POTENTIALS, E°

Na ⁺ + e ⇌ Na	-2.713
Mg ²⁺ + 2e ⇌ Mg	-2.37
Al ³⁺ + 3e ⇌ Al	-1.66
Zn ²⁺ + 2e ⇌ Zn	-0.763
Fe ²⁺ + 2e ⇌ Fe	-0.44
Cd ²⁺ + 2e ⇌ Cd	-0.403
Cr ³⁺ + e ⇌ Cr ²⁺	-0.38
Tl ⁺ + e ⇌ Tl	-0.336
V ³⁺ + e ⇌ V ²⁺	-0.255
Sn ²⁺ + 2e ⇌ Sn	-0.14
Pb ²⁺ + 2e ⇌ Pb	-0.126
2H ⁺ + 2e ⇌ H ₂	0.000
S ₄ O ₆ ²⁻ + 2e ⇌ 2S ₂ O ₃ ²⁻	0.09
TiO ²⁺ + 2H ⁺ + e ⇌ Ti ³⁺ + H ₂ O	0.10
S + 2H ⁺ + 2e ⇌ H ₂ S	0.14
Sn ⁴⁺ + 2e ⇌ Sn ²⁺	0.14
Cu ²⁺ + e ⇌ Cu ⁺	0.17
SO ₄ ²⁻ + 4H ⁺ + 2e ⇌ H ₂ O + H ₂ SO ₃	0.17
AgCl + e ⇌ Cl ⁻ + Ag	0.222
Saturated calomel	(0.244)
Hg ₂ Cl ₂ + 2e ⇌ 2Cl ⁻ + 2Hg	0.268
Bi ³⁺ + 3e ⇌ Bi	0.293
UO ₂ ²⁺ + 4H ⁺ + 2e ⇌ U ⁴⁺ + 2H ₂ O	0.33
VO ²⁺ + 2H ⁺ + e ⇌ V ³⁺ + H ₂ O	0.34
Cu ²⁺ + 2e ⇌ Cu	0.34
Fe(CN) ₆ ³⁻ + e ⇌ Fe(CN) ₆ ⁴⁻	0.355
Cu ⁺ + e ⇌ Cu	0.52
I ₃ ⁻ + 2e ⇌ 3I ⁻	0.545
H ₃ AsO ₄ + 2H ⁺ + 2e ⇌ H ₃ AsO ₃ + H ₂ O	0.56
I ₂ + 2e ⇌ 2I ⁻	0.621
2HgCl ₂ + 2e ⇌ Hg ₂ Cl ₂ + 2Cl ⁻	0.63
O ₂ + 2H ⁺ + 2e ⇌ H ₂ O ₂	0.69
Quinone + 2H ⁺ + 2e ⇌ Hydroquinone	0.70
Fe ³⁺ + e ⇌ Fe ²⁺	0.771
Hg ₂ ²⁺ + 2e ⇌ 2Hg	0.792
Ag ⁺ + e ⇌ Ag	0.799
Hg ²⁺ + 2e ⇌ Hg	0.851
2Hg ²⁺ + 2e ⇌ Hg ₂ ²⁺	0.907
NO ₃ ⁻ + 3H ⁺ + 2e ⇌ HNO ₂ + H ₂ O	0.94
HNO ₂ + H ⁺ + e ⇌ NO + H ₂ O	0.98
VO ₂ ⁺ + 2H ⁺ + e ⇌ VO ²⁺ + H ₂ O	0.999
Br ₂ + 2e ⇌ 2Br ⁻	1.08
2IO ₃ ⁻ + 12H ⁺ + 10e ⇌ 6H ₂ O + I ₂	1.19
O ₂ + 4H ⁺ + 4e ⇌ 2H ₂ O	1.229
MnO ₂ + 4H ⁺ + 2e ⇌ Mn ²⁺ + 2H ₂ O	1.23
Cr ₂ O ₇ ²⁻ + 14H ⁺ + 6e ⇌ 7H ₂ O + 2Cr ³⁺	1.33
Cl ₂ + 2e ⇌ 2Cl ⁻	1.358
2BrO ₃ ⁻ + 12H ⁺ + 10e ⇌ 6H ₂ O + Br ₂	1.50
MnO ₄ ⁻ + 8H ⁺ + 5e ⇌ 4H ₂ O + Mn ²⁺	1.51
Ce ⁴⁺ + e ⇌ Ce ³⁺	1.61

13. MEAN ACTIVITY COEFFICIENTS

M	KCl	Na ₂ SO ₄	ZnSO ₄
0.001	0.965	0.89	0.70
0.01	0.901	0.72	0.39
0.1	0.769	0.45	0.15
1.0	0.606	0.20	0.045

16. HEATS OF FORMATION

ΔH° in kJ mol⁻¹ at 25°C
All ions in H₂O solution except as noted

All Elements = 0

H ₂	218	H ⁺	0.0	H ₂ O _g	-242
O ₂	249	Na ⁺	-240	H ₂ O _l	-286
C _g	717	Ag ⁺	106	CO _g	-111
N ₂	473	NH ₄ ⁺	-133	CO _{2g}	-394
F ₂	79	OH ⁻	-230	NH _{3g}	-46
Cl ₂	122	F ⁻	-333	NO _g	90
Br ₂	112	Cl ⁻	-167	NO _{2g}	33
I ₂	107	Br ⁻	-122	N ₂ O _{4g}	9
S ₈	279	I ⁻	-55	SO _{2g}	-297
P ₄	315	S=	33	SO _{3g}	-396
Na ₂	107	SO ₄ ²⁻	-909	H ₂ S _g	-21
K ₂	88	CO ₃ ²⁻	-677	NaF _s	-574
Na ⁺	609	HF _g	-271	NaCl _s	-411
K ⁺	514	HCl _g	-92	KF _s	-567
F ₂	-255	HBr _g	-36	KCl _s	-437
Cl ₂	-233	HI _g	26	AgCl _s	-127
CH _{4g}	-75	HCN _g	135	AgBr _s	-100
C ₂ H _{2g}	227	PH _{3g}	5	PCl _{3g}	-287
C ₂ H _{4g}	52	C ₆ H _{6l}	49	PCl _{5g}	-375
C ₂ H _{6g}	-85	CH ₃ OH _l	-238		
C ₃ H _{8g}	-105	C ₂ H ₅ OH _g	-235		
nC ₄ H _{10g}	-127	C ₂ H ₅ OH _l	-278		
nC ₈ H _{18g}	-209	COCl _{2g}	-219		
CCl _{4l}	-135	CH ₃ Cl _g	-81		

17. ABS. ENTROPY S°

J mol⁻¹ K⁻¹ at 25°C

H ₂	131	P _{4wh}	164	SF _{6g}	292
N ₂	192	HF _g	174	NO _g	211
O ₂	205	HCl _g	187	NO _{2g}	240
Cl ₂	223	H ₂ O _g	189	N ₂ O _{4g}	304
F ₂	203	CO _g	198	NH _{3g}	192
Cgra	5.7	CO _{2g}	214	PCl _{3g}	312
S _{8r}	254	SO _{2g}	248	PCl _{5g}	365
CH _{4g}	186	SO _{3g}	256	BF _{3g}	254
C ₂ H _{6g}	229	CH ₃ OH _l	127		
C ₃ H _{8g}	270	C ₂ H ₅ OH _g	283		
C ₂ H _{2g}	201	C ₂ H ₅ OH _l	161		
C ₂ H _{4g}	219	(CH ₃) ₂ O _g	266		
C ₆ H _{6g}	269	CH ₃ COOH _g	282		

18. ΔG° FORMATION

kJ mol⁻¹ at 25°C

H ₂	203	HF _g	-273	H ₂ O _g	-229
F ₂	62	HCl _g	-95	H ₂ O _l	-237
Cl ₂	106	HBr _g	-54	SO _{2g}	-300
O ₂	232	HI _g	1.7	SO _{3g}	-371
NO _g	87	NH _{3g}	-16	PCl _{3g}	-268
NO _{2g}	51	CO _g	-137	PCl _{5g}	-305
N ₂ O _{4g}	98	CO _{2g}	-394	CH _{4g}	-51
C ₂ H _{4g}	68	C ₂ H _{2g}	209	C ₂ H _{6g}	-33
C ₆ H _{6l}	125	CH ₃ OH _l	-162		
CCl _{4l}	-65	C ₂ H ₅ OH _l	-175		
BF _{3g}	-1120	CHCl _{3g}	-70		
SF _{6g}	-1105	CH ₃ COOH _g	-374		
NH ₄ Cl _s	-203	(CH ₃) ₂ O _g	-113		

20. CONC. ACIDS AND BASES

	M.W.	Density	Wt. %	Molar-ity
Acetic	60.05	1.05	99.5	17.4
H ₂ SO ₄	98.07	1.83	94	17.6
HF	20.01	1.14	45	25.7
HCl	36.46	1.19	38	12.4
HBr	80.91	1.52	48	9.0
HNO ₃	63.01	1.41	69	15.4
HClO ₄	100.46	1.67	70	11.6
H ₃ PO ₄	98.00	1.69	85	14.7
NaOH	40.00	1.53	50	19.1
NH ₃	17.03	0.90	28	14.8

21. DENSITIES (g cm⁻³)

Water at	Air (70 cm)	0.0011
0°C 0.9168	Glass	2.7
10° 0.9997	Na ₂ CO ₃	2.5
20° 0.9982	NaCl	2.2
22° 0.9978	BaSO ₄	4.5
24° 0.9973	AgCl	5.6
26° 0.9968	Aluminum	2.7
28° 0.9963	Iron	7.9
30° 0.9956	Brass	8.4
90° 0.9653	Mercury	13.6
100° 0.0006	Platinum	21.4

22. MOBILITIES (m²V⁻¹s⁻¹ × 10⁹)

Li ⁺	39	H ₃ O ⁺	350	1/2Ba ²⁺	64
Na ⁺	50	NH ₄ ⁺	73	1/2La ³⁺	70
K ⁺	74	Ag ⁺	62	1/2SO ₄ ²⁻	80
Cl ⁻	76	OH ⁻	198	1/2PO ₄ ³⁻	80
Br ⁻	78	I ⁻	77	NO ₃ ⁻	71

23. WATER V.P. (torr)

0°C	4.6	25°	23.8
15°	12.8	30°	31.8
20°	17.5	50°	92.5

24. MISCELLANEOUS

Std. dev. = $\sqrt{\frac{1}{n} \sum (X_i - \bar{X})^2}$ (n-1)
 Conf. limits = $\bar{X} \pm t_s / \sqrt{n}$
 $E = E^\circ - (0.0592/n) \log ([Red]/[Ox])$
 $\log I_2/I_1 = abc = A = \log 1/T$
 $\log N_T = \log N_0 - 0.301T/T_i$
 $x = (-b \pm \sqrt{b^2 - 4ac})/2a$
 $n\lambda = 2d \sin \theta$
 $2.303 \log_{10} a = \log_e a$
 $h = 6.626 \times 10^{-34} \text{ J s}$
 $e = 1.602 \times 10^{-19} \text{ C}$
 $N_A = 6.022 \times 10^{23} \text{ mol}^{-1}$
 $F = 96487 \text{ C}$
 $g = 9.807 \text{ m s}^{-2}$
 $c = 2.998 \times 10^8 \text{ m s}^{-1}$
 $1 \text{ amu} = 1.661 \times 10^{-27} \text{ kg}$
 $R = 1.987 \text{ cal mol}^{-1} \text{ K}^{-1}$
 $= 0.08206 \text{ litre atm mol}^{-1} \text{ K}^{-1}$
 $= 8.314 \text{ J mol}^{-1} \text{ K}^{-1}$
 $= 8.314 \text{ kPa dm}^3 \text{ mol}^{-1} \text{ K}^{-1}$
 $0^\circ \text{C} = 273.15 \text{ K}$
 $1 \text{ eV} = 1.602 \times 10^{-19} \text{ J}$
 $1 \text{ cal} = 4.1840 \text{ J}$
 $760 \text{ torr} = 101.3 \text{ kPa}$