

(A Corporate Statutory Body Constituted by an Act of State Legislature)

ALTO BETIM - GOA 403 521

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GBSHSE/ACAD/2012/1857

Date. 16th July, 2012

To, The Heads of all recognised Higher Secondary Schools, within the jurisdiction of this Board.

Sub: Evaluation Scheme/Design of question paper of Std. XII – 2012 – 13 in Psychology/ Mathematics/Practicals

Sir/Madam,

With reference to the subject cited above kindly note the changes made in the Evaluation Scheme / Design of Question Paper for the following subjects of Std XII for this Academic year 2012 - 13.

- Psychology: Design of Question paper Mid Term 20 marks Weightage to concept / Subject units
 At Sr. No. 2: pages 32 to 46 may be deleted, for 1st Mid Term Test 2012.
- **2.** <u>Mathematics:</u> Design of the Question Paper Second Term 80 marks <u>Weightage to learning outcomes.</u>

S. No	Learning outcomes	Marks	% of Marks
1.	Knowledge	20	25%
2.	Understanding	36	45%
3.	Application	20	25%
4.	Skill	04	05%
	Total	80	100%

Practical subjects: viz Physics, Chemistry, Biology and Computer Science;ADDENDUM:

Refer Circular No. 19 dated 25th April 2012 in respect of Internal Assessment and its weightage in H.S.S.C. Examination to be conducted by the Board in March 2013.

- 20 marks for Assignment /Project will be as follows:
- a. 10 marks will be allotted for project work and same shall be added to practical component.
- b. 10 marks will be allotted for assignment on theory and it shall be added to the theory component.

You are requested to bring to the notice of all concerned.

sd/-(Bhagirath G. Shetye) Secretary

Enclosures:

- Pattern and design of the question paper for Mid Term, 1st Term, 2nd Tern and Final Exam of Physics and Chemistry.
- 2. Clarification regarding syllabus of practical in Physics and Chemistry.

	MID TERM TEST	-	FIRST TERM		SECOND TERM		FINAL	
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MAX MAKKS NO OF	2		9		9		5	
1		\dagger	2007	†	Knowledge: 30%		Knowledge: 30 %\	
WEIGHTAGE K	Knowledge: 40 %\		Knowledge: 50 %	-			.00	
	hin		oi.		Amiliation: 20%		Amplication . 20%	
OBJECTIVES Ap	Application: 10%		Application: 20 %	1		90	1 3	4
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<u>ರ</u>		æ 8	Current Electricity	8 9	Atoms	0.2	Moving Charges and Magnetism	05
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	1		Wave Optics	80	fagnetism	05	Electromagnetic waves	2,5
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		1	T A (41)	80	T A (4 m/s)	80	LA (4 mks) x 2	80
T)	LA(4mks)	\$ 8	LA (+ IIIKs) SAII(3 mks)	3 ~	(8)	18	SAII (3 mks) x 5	15
	SAII (3 mks)	3 8	SAL(3 mks)	22		22	SAI (2 mks) x 11	22
QUESTIONS S	SAI (2 mks)	3 5	VSA (1mks)	12		12	VSA (1mks) x 10	2
OF	NO OPTIONS		OPTIONS IN TWO LA TYPE + 1		OPTIONS IN TWO LA TYPE + 1 SAII	SAII	OPTIONS IN TWO LA TYPE + $1 \text{ SAII TYPE} = 20 \%$	
OPTIONS			SAII TYPE		IIFE			
DIEDICH TV E	Fosy 30%		Fasv 30%		Easy 30%			
	age		age iit		Average 60% Difficult 10%		Average 60% Difficult 20%	
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PRACTICALS

FIRST TERM 20 MKS (BASED ON EXPTS AND ACTIVITIES PERFORMED DURING THE TERM)

1 Expt (to be performed during the	VITIES PEI
1 Expt (to be performed during the exam) Record of 4 Activities	10 mks
Viva on Activities/Expts	04 mk
Journal	02 mk
Break Lin of Montre to 1	04 mks

Break Up of Marks to be awarded for Expt: Setting/ Diagram Observations Calculations/Graph Result	021	Break Up of Marks to be awarded for Journal Completeness Neatness Regularity	02 mks 01 mks 01mks	

SECOND TERM 20 MKS (BASED ON EXPTS AND ACTIVITIES PERFORMED DURING THE TERM)

1 Expt (to be performed during all	CTIVITIES I
1 Expt (to be performed during the exam) Record of 4 Activities	10 mks
Viva on Activities/Expts	04 mk
Journal	02 mk
Break Up of Marke to be 110	04 mks

Break Up of Marks to be awarded for Expt: Setting/ Diagram Observations Calculations/Graph Result	03 mks 04 mks	Break Up of Marks to be awarded for Journal Completeness Neatness Regularity	02 mks 01 mks 01mks

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GOA BOARD OF SECON	DARY AND HIGHER SECONDA	RY EDU	CATION		- Marie - Array - Arra
	ALTO BETIM – GOA 403:	521			
CLASS: XII	Ş	SUBJECT	:СНЕМІ	STRY	
PATTERN AND DESIGN OF	F QUESTION PAPERS (2012-2013 O	NWARDS	3)		
		MID TERM TEST	FIRST TERM	SECOND TERM	FINAL
TIME IN MINS		60	150	150	150
MAX MARKS		20	60	60	55
NO OF MAIN QUESTIONS		2	6	6	5
WEIGHTAGE	KNOWLEDGE	6	18	18	17
	UNDERSTANDING	9	24	24	21
то	APPLICATION	3	12	12	11
OBJECTIVE	SKILL	2	6	6	6
WEIGHTAGE	THE SOLID STATE	4	4	†	3
TO CONTENT	GENERAL PRINCIPLES AND PROCESSES OF ISOLATION OF ELEMENTS	3	3		3
	CHEMISTRY IN EVERYDAY LIFE	3	3		2
	CHEMICAL KINETICS	5	5		4
	HALOALKANES AND HALOARENES	5	5		3
	SURFACE CHEMISTRY		8	2	3
	ELECTROCHEMISTRY	 	8	3	4
	THE d-AND f-BLOCK ELEMENTS	<u> </u>	9	3	4
	ALCOHOLS,PHENOLS AND ETHERS		8	2	3
	BIOMOLECULES		7	2	3
	SOLUTIONS		1	8	4
	THE P-BLOCK ELEMENTS			10	6
	COORDINATION COMPOUNDS			8	3
	ALDEHYDES, KETONES AND CARBOXYLIC ACIDS			10	5
	AMINES		1	7	3
	POLYMERS			5	2
WEIGHTAGE TO TYPE OF	VSA	3	10	10	10
QUESTIONS	SAI	10	30	30	22
	SA II	3	12	12	15
	LA	4	8	8	8
SCHEME OF OPTIONS		<u> </u>	J:		
DIFFICULTY LEVEL	EASY	20%	20%	20%	20%
	AVERAGE	60%	60%	60%	60%
•	DIFFICULT	20%	20%	20%	20%

MARKS DISTRIBUTION FOR FIRST TERM PRACTICAL EXAM IN CHEMISTRY FOR STD. XII SC.

MAX. MKS : 20 MKS. DURATION 2 HRS.

Q1. Volumetric Analysis 07 mks

Q2. Inorganic salt (container D) 07 mks

Q3. Functional Group determination (container F) 02 mks

Q4. Journal + Viva 02 + 02 mks

Marking scheme for Volumetric analysis and Inorganic salt analysis to be followed as earlier.

MARKS DISTRIBUTION FOR SECOND TERM PRACTICAL EXAM IN CHEMISTRY FOR STD. XII SC.

MAX. MKS : 20 MKS. DURATION: 3 HRS.

Q1. Volumetric Analysis

06 mks

Q2. Physical Experiments 03 mks

Q3. Journal + Viva 01+01 mk.

Q4. Inorganic salt analysis (container D) 06 mks

Q5. Natural organic compound (container E) 01 mk

Q6. Functional Group determination (container F) 02mks

NOTE: The second term practical Examination also includes investigatory project carrying 10 mks

Marks for the project to be allotted as follows:

Neatly hand written project
 Oral presentation
 O3 mks

3. Viva on Project 02 mks

NOTE: Marking scheme for Volumetric analysis and Inorganic salt analysis will be like that of the Final Examination.

MARKS DISTRIBUTION FOR FINAL PRACTICAL EXAM IN CHEMISTRY FOR STD. XII SC MAX. MARKS: 25 **DURATION: 3HRS.** Q1. Volumetric Analysis 06mks Q2. Physical Experiment 03mks Q3. Journal + Viva 01 +02 mks Q4. Inorganic salt (container D) 06 mks Q5. Functional Group determination (container F) 02 mks Q6.Identification of natural organic compound (container E) 01 mk Q7. Project + Viva 02+02 mks Marking scheme Q1. Volumetric analysis 06 mks Observation 01 mk Correct volume of stock solution 01 mk Readings 03 mks Calculations 01 mk as follows: i) Calculations of N/M of solution in container B using $N_3V_1=N_2\,V_2$ or $a_1\,m_1\,v_1=a_2\,m_2\,v_2-\mathcal{V}_2\,mk$ ii) Percentage purity of the solution in container B ______ g of which have been dissolved per Note: Calculations should be either in terms of normality or molarity throughout . Q4. Inorganic salt analysis (container D) 06 mks Dry tests 2 1/2 mks Cation 2mks as follows Group separation 01 mk Identification of cation ½ mk

½ mk

Anion

01mk as follows

Identification

½ mk

C.T

½ mk

Correct formula of the compound

½ mk

Internal Examiner shall evaluate

Q1.

06mks

Q2.

03 mks

Q3.

01+ 2 mks

Total

12 mks

External Examiner shall evaluate

Q4

06 mks

Q5

02 mks

Q6

01 mk

Q7

02 +02 mks

Total

13 mks

Clarification Regarding Syllabus of Practical's In Chemistry For Std XII Science For The Academic Year 2012-2013.

With respect to sets (A), (B), (C), (D) & (E) following experiments are to be performed and evaluated for Board Exam.

- 1. Preparation of lyophilic sol of starch
- 2. Study of role of emulsifying agent (soap) on stabilizing (o/w) emulsions of:
 - Mustard Oil.
 - ii. Castor Oil
 - iii. Coconut Oil.
- Effect of concentration on the rate of reaction between sodium thiosulphate and HCL.
- 4. Enthalpy of neutralisation of strong acid (HCL) and strong base (NaOH).

From sets D & E the following experiments are to be demonstrated only:

- Variation of cell potential in Zn|Zn²⁺|| Cu²⁺| Cu with a change in concentration of electrolytes (CuSo₄ or ZnSo₄) at room temperature.
- Separation of pigments from extracts of leaves by paper chromatography and determination of R_f values.

From sets F and G following preparation are to be demonstrated only:

- 1. Preparation of double salt of ferrous ammonium sulphate (1st term).
- Preparation of Acetanilide
 (2ndterm).

With respect to set J students will be required to prepare standard solution from the stock solution provided to them (as done earlier)."

With respect to **set K** compounds of As3+ are not to be given.

III. GALVANIC CELL

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Introduction: Electrochemical cells can be used to convert chemical energy into electrical energy by the chemical reactions taking place at the two electrodes in the cell.

It is an oxidation reduction reaction in the cell which produces electric current. For this purpose the oxidising and reducing reagents are to be kept separate so that the electron transfer occurs through a conducting wire. The development of the potential at the electrode depends on temperature and concentration.

The electrochemical cell which converts chemical energy into electrical energy is called a galvanic or voltaic cell after Galvani (1780) who invented the cell. A typical galvanic cell is the Daniel cell.

Theory: A galvanic cell can be constructed by taking two electrodes, zinc and copper, and dipping them in their respective salt solutions of the same strength, i.e. ZnSO₄ and CuSO₄, and connecting the two solutions by a salt—idge and the two electrodes by a conducting wire when the current flows from zinc to copper as long as the circuit is closed. The reactions taking place at the zinc and copper electrodes are as follows:

At the Zinc electrode:

$${\rm Zn}_{(s)}$$
 ----> ${\rm Zn}^{2+}_{(aq)}$ +2e (half cell oxidation reaction)

Zinc has a greater tendency to ionise than copper since zinc is placed higher than copper in the Electrochemical series and hence it leaves its electrons on the zinc electrode and goes as Zn^{++} ions in its salt solution rendering the zinc electrode negative. Since the electrons are lost to the copper electrode, Zinc electrode is the anode and the process occurring at this electrode is oxidation.

At the copper electrode:

$$\mathrm{Cu^{2+}}_{(aq)}$$
 + 2e $^-$ ----> $\mathrm{Cu_{(s)}}$ (half cell reduction reaction)

The electrons released by the zinc atoms pass to the copper electrode where the Cu⁺⁺ ions from the solution pick up these electrons and are discharged at the copper electrode, making this electrode positively charged and as such the copper electrode is the cathode and the process occuring at this electrode is reduction.

The overall cell reaction = half cell oxidation reaction + half cell reduction reaction which is written as:

$$Zn_{\{s\}} + Cu^{2+}_{\{aq\}} -----> Zn^{2+}_{\{aq\}} + Cu_{\{s\}}$$

represented as ${}^-Z_\Pi\backslash Z_\Pi^{2+}\backslash\backslash Cu^{2+}\backslash Cu^{+}$ (IM) (IM)

Where $E_{cell}=1.1$ volts at 25°C when the concentrations of the two salt solutions are 1 M

Electromotive force of a cell (E_{cell}):

Current flows from one electrode to the other due to the difference in potential between the two electrodes. The flow of current from the electrode of higher potential to the electrode of lower potential causes this difference in potential which is called **Experiment**:

Aim : To set up a simple galvanic cell and determine its emf using salt solutions of three different concentrations and to find the change of emf with time.

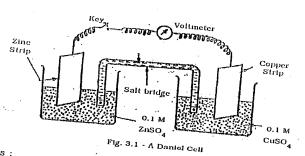
Principle: Since the potential of an electrode depends upon concentration, the emf of the cell will change with different concentrations of the salt

Apparatus: 2 beakers (100 mi), salt bridge, filter paper strip, connecting wires, sand paper, high resistance voltmeter, 2 pinch-cocks, potentiometer.

Materials and chemicals : Zinc strip, copper strip, zinc sulphate, copper sulphate, distilled water, sulphuric acid, filter paper.

Procedure: Propage 200 ml. of 1 M and 0.1 M solutions of CuSO₄ and ZnSO₄ two 100 ml. beakers, and prepare three sets of the above solutions as shown in the observation table below. Clean properly the zinc and the ZnSO₄ and CuSO₄ solutions, respectively. Connect the solutions in the two beakers by a salt bridge or a filter paper saturated with ICCl which acts as an external circuit in which is inserted a key and a high arranged as shown in the Fig. 3.1.

Use the first set of solutions. Press the key to complete the electrical circuit. On pressing the key a deflection is observed in the voltmeter. Take the voltmeter reading and record it at regular intervals of time (every 5 minutes) as shown in the observation table. Repeat the experiment such that the cell potential values and their change at regular intervals of time is recorded for the other two sets.



Observations:

Sr. No.	Concentrations of Solutions	Cell Potential	Voltmeter reading for time in minutes
1.	IM ZnSO ₄ and	-	0 5 10 15
	IM CuSO4	1	18
2.	0.1M ZnSO		
з.	O.1 M CuSO ₄		
	0.1 M ZnSO ₄ and 1 M CuSO ₄		
: Dit			1 1 1

Note: Different concentrations of ${\rm ZnSO_4}$ and ${\rm CuSO_4}$ may also be considered in the Calculations:

1)
$$E^{o}_{cell} = E^{o} - E^{o}$$

cathode anode (RHC) (LHC)

$$= E^{o} - E^{o}$$

$$Cu^{2+}/Cu = Zn^{2+}/Zn$$

$$= +0.34 V - (-0.76 V)$$

$$= +1.10V \text{ at } 298^{o}K$$
2) $E = E^{o} - 2.303RT \text{ Log } Q$
Cell Cell , nF

Now log
$$Q = \frac{[Zn^{2+}][Cu]}{[Cu^{2+}][Zn]} = \frac{[Zn^{2+}]}{[Cu^{2+}]}$$
 Since $[Cu] = [Zn] = 1$

$$= \frac{0.1}{0.1} = 1$$

$$\therefore$$
 log Q = 1, n = 2. E_{Cell}^0 = +1.10 Volts and log I = 0

T = Room temperature

$$\therefore E_{\text{cell}} = E_{\text{cell}}^{0} - \frac{2.303 \times 8.3143 \times T \times 0}{2 \times 96,500}$$

$$E_{\text{cell}} = E_{\text{cell}}^{\text{o}}$$
$$= +1.10 \text{ V}$$

3]
$$\cdot Q = \frac{(Zn^{2+})}{(Cu^{2+})} = \frac{0.1}{1.0} = 10^{-1} \therefore \log Q = \log 10^{-1} = -1$$

$$\therefore$$
 n = 2, E_{coll}^{o} = + 1.10 V and T-Room temperature

$$E_{\text{cell}} = E_{\text{cell}}^{0} - 2.303 \text{ RT log Q}$$

$$n F$$

$$= E_{\text{cell}}^{0} - 2.303 \times 8.3143 \times T \times (-1)$$

$$2 \times 96,500$$

= X Volts

Result: The emf of the cell is -V.

The cell potential depends on the concentration of the solutions and decreases with time.

- **Precautions:** i) The terminals of the connecting wires and those of the electrodes, voltmeter, key should be cleaned with sand paper.
 - Zinc and copper rods should be cleaned properly before dipping them in their respective solutions.
 - iii) The connections should be tightly fixed and should be checked before pressing the key.

Object. To compare the effectiveness of a number of oils in forming the emulsions.

Theory. Not all oils form emulsions with water with the same case using the same emulsifier. The extent of emulsification depends upon the nature of oil and the nature of emulsifier used.

Apparatus required. Measuring cylinder 25 mL (one), Graduated pipette 10 mL (four), Stop watch (one), Test tubes (eight).

Reagents required. Four different oils e.g., Mustard oil, Cotton seed oil, Castor oil, and Kerosene oil, 1% solution of sodium oleate (soap

Procedure. This experiment may be performed in two parts.

- (i) When oil is in larger proportions. In this case, the resulting emulsion is water-in-oil (W/O) type.
- (ii) When water is in larger proportions. In this case the resulting emulsion is oil-in-water (O/W) type.

Preparation of Water-in-oil (W/O) Emulsion

Pipette out 5 mL of each oil in a separate test tube, label each test tube and arrange them on a test tube stand. Add 2 mL of distilled water and 0.5 mL of 1% soap solution to each test tube. Shake the contents of each tube thoroughly and place it back on the test tube stand. Observe and record the time taken by each system for breaking into two layers, in the following table. The emulsion taking longer time for separating into oil and water is more stable. Each oil can thus be rated in terms of time its emulsion takes for breaking up into two layers.

Preparation of Oil-in-water (O/W) Emulsion

Pipette out 2 mL of each oil separately in test tubes. Label them all. Add 5 mL of distilled water and 0.5 mL of 1% soap solution to each test tube. Shake the contents of each test tube well and place them on a test tube stand.

Observe and record the time taken by the emulsion to break up i.e., separates into two layers. Rate each oil according to the time taken by each system. The most stable emulsion takes the longest to separate into layers. The corresponding oil thus is easily emulsified. Observations.

10.	Name of the oil	Time taken for the separation of two layers, minutes	Rating of the oil
	Fa	or case (i): W/O emulsion	
	Mustard oil Cotton seed oil	tį	****
Ć	Castor oil	<i>t</i> ₂ <i>t</i> ₃	*****
	For	14 case (ii) : O/W emulsion	*****
	Mustard oil Cotton seed oil	t_1'	*****
ر د ت	Castor oil	t'2 t'3	
	Its. The rating of	1'4	

Results. The rating of the oils on the basis of the stability of the emulsions formed is,

For case (i) oil oil oil oil For case (ii) oil oil oil oil

Discussion. Try to correlate the rating of the oils with their structures.

Precautions. (i) Shaking of the contents of each test tube should be done for the same period of time, and with almost similar force.

(ii) Breaking up time of an emulsion should be the time taken by an emulsion to separate completely into two layers.

Experiment 3.2

Object. To compare the effectiveness of various emulsifying agents in stabilizing emulsions.

Theory. The emulsifying capacity of an emulsifier is governed by it tendency to lower down the interfacial tension between the dispersed phase and the dispersion medium. Obviously, it is related to its structural parameters like chain-length, nature of functional group and that of the oil. The hydrocarbon end of the emulsifier is oil-soluble and the ionizable group gets oriented towards water. Thus, an emulsifier forms a layer around the droplets of the dispersed phase and prevents coalescence of the dispersed phase.

Experiment 5.2

Object. To investigate the separation of the coloured components present in a given flower/grass extract by paper chromatography. Determine their R_f values.

Theory. Flowers and grass contain certain highly coloured pigments. These pigments can be extracted with acetone. The extract can be concentrated by leaving the extract in a china dish, when the volatile solvent evaporates.

The mixture of pigments is then allowed to get partitioned between immobile and mobile phases. Due to difference in their adsorption characteristics, different pigment molecules travel with different speeds when made to migrate on chromatographic paper.

Apparatus required. Same as in Experiment 5.1.

Reagents required. Flowers or grass, Petroleum ether, Acetone, Filter paper no.1, and the Solvent (a mixture of petroleum ether and acetone in the ratio 3:1).

Procedure. Grind the flower or grass with acetone in a pastle and mortar, till the coloured components get extracted. Filter the solution and preserve it in a corked test tube. Cut a filter paper strip of suitable dimensions, draw a pencil line about 3 cm from the lower edge, and put a cross (x) near at the middle of the line. With the help of a fine capillary, place a drop of the flower/grass extract (obtained above) at the cross mark. Let the spot dry up. Place another drop of this extract at the same place. Dry it in air. Pour the solvent into the jar so as to form a layer of about 2-3 cm height. Clip the upper end of the filter paper into the hook so that the lower end just dips into the solvent. Cover the jar and let the solvent move up. Remove the filter paper strip from the jar as soon as liquid rises to a height of about 20 cm.

Mark the solvent front with a pencil. Dry the filter paper in air and mark the coloured spots with a pencil. Measure the distance of each spot and that of the solvent front from the base-line. Record your observations and calculate the R_{ℓ} values for each component.

Observations and calculations. Temperature = t° C Distance of solvent from the base line = p cm

Sr.no.	Colour of the spot	Distance moved, cm	R_f
1	Blue	x	x/p
2	Green	у	y/p
3	Red	z	z/p

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Results. The R_f values calculated above, follow the order

Hence, pigment molecules move fastest, and that of the pigment slowest.

AZJ PREFARACION OF ACETANILIDE

Acetanilide is an acetyl derivative of aniline. Its molecular formula is C₆H₅NH.COCH₃.

Experiment 12.3

Object. To prepare a sample of acetanilide.

Apparatus required. Round bottomed flask (100 mL), Reflux condenser, Measuring cylinder (25 mL), Beakers and funnel.

(Fin = .	Heakers	and funnel.
Reagents required.		5 ml.
	Acetic anhydride	5 mL
	Glacial acetic acid	5 mL
	Zinc dust	1 g
	Animal charcoal	l g

Theory. Acctanilide can be obtained by acetylation of aniline using

acetic anhydride in glacial acetic acid at refluxing temperature.

Reaction.

```
heat C<sub>6</sub>H<sub>5</sub>NH.CO.CH<sub>3</sub> + CH<sub>3</sub>COOH
C_6H_5NH_2 + (CH_3CO)_2O
  aniline
                acetic anhydride
```

Procedure. Take 5 mL of aniline in a clean dry round bottomed flask (100 mL). To it, add slowly a mixture containing 5 mL each of acetic anhydride and glacial acetic acid. After complete addition, add 1 g of zinc dust and fit up the flask with a reflux condenser. Reflux it for about 10 minutes on a sand bath.

Remove the condenser and pour the hot reaction mixture in 100 mL of cold water (taken in a 250 mL beaker). Care should be taken not to allow zinc dust to pass into the beaker. Stirt the mixture thoroughly with a glass rod. Colourless crystals of acetanilide separate out. Filter the crystalline material on a Buchner funnel under suction, and dry the material on filter paper pads.

Purification by recrystallization. Transfer the crude sample of acetanilide to a 250 mL beaker. Add 100 mL of boiling water and 5 drops of ethyl alcohol. Boil the solution to obtain a clear solution. Add 1 g of animal charcoal, boil for a minute and filter the hot solution. Allow the filtrate to cool. Shining crystals of acetanilide separate out. Filter these crystals on a Buchner funnel under suction. Dry the crystals between the folds of the filter paper. Determine the melting point of the purified and dried sample.

Results.	Total yield of the crude sample	sample.
	Colour of the crude sample	= g
	Colour of the recrystallized sample	
	Melting point of the many this	=
Precautio	Melting point of the recrystallized sample	= °C

Precautions. (i) If possible, the freshly distilled aniline should be used. (ii) While transferring the contents of the flask into water taken in a beaker, care should be taken not to allow zinc dust to pass into the beaker.

Fig. 1.6. Drying or me crystais, (4) pressing occur (b) in a dessicator.

Object. To prepare a crystalline sample of ferrous ammonium sulphate Experiment 1.1 (Molur's salt), FcSO4.(NH4)2SO4.6 H2O.

Theory. When a saturated aqueous solution containing equimolar amounts of ferrous sulphate and ammonium sulphate is cooled, crystals of Mohr's salt are obtained. The crystalline material corresponds to a stoichiometry represented by the formula FeSO₄.(NH₄)₂SO₄.6H₂O. These crystals have a light green colour.

Stoichiometry. The stoichiometric equation and the relative amounts of various salts involved in this preparation are,

FeSO_{4.7}H₂O + (NH₄)₂SO₄ + x H₂O \rightarrow FeSO₄.(NH₄)₂SO_{4.6}H₂O + x¹ H₂O [56+32+64+2(14+4)+32+64+6(2+16)] 1 mole [56+32+64+(7x18)] [2(14+4)+32+64] 392 g / mol

Thus, for preparing 392 g of the Mohr's salt, one would need 278 g of 278 g/mol 132 g/mol FcSO_{4.7H2O} (ferrous sulphate heptahydrate) and 132 g of (NH₄)₂SO₄. In order to prepare a certain desired amount of the Mohr's salt, say W g, the relative amounts of the two constituent salts can be calculated as follows, viz.,

of the two constituent sails can be calculated.

Mass of FeSO₄.7H₂O =
$$\frac{278 \times W}{392}$$
 g

Mass of (NH₄)₂SO₄ = $\frac{132 \times W}{392}$ g

Thus, for preparing a 10 g sample of the Mohr's salt, one would need as follows:

Mass of FeSO_{4.7}H₂O required =
$$\frac{278 \times 10}{392} \approx 7.1 \text{ g}$$

Mass of (NH₄)₂SO₄ required = $\frac{132 \times 10}{392} \approx 3.4 \text{ g}$

Apparatus required. 3 Beakers 100 mL, Porcelain (china) dish (3"diameter), Glass funnel, Glass rod, and an ice bath.

Reagents required. Ferrous sulphate (heptahydrate) crystals = 7 g= 3.5 gAmmonium sulphate = 1-2mLDil. sulphuric acid

Procedure. Boil about 40-50 mL of distilled water in a beaker. In another beaker, place the required amounts of ferrous sulphate and ammonium sulphate, and add 2 mL of dil.H₂SO₄. Now, add boiling water in small amounts to the beaker containing the two salts while continuously stirring the contents. The contents of the beaker may be heated to nearly boiling, and if there is still some salt left undissolved, then add a little more warm distilled water from the other beaker and stirr it well to dissolve the salts completely. Filter the hot solution through a filter paper cone, and collect the filtrate in the china dish. Heat the contents of the china dish gently to concentrate the solution up to crystallization point.

To check the crystallization point, one end of the glass rod is dipped into the solution in the china dish and cooled in air by blowing air on to it. If fine crystals appear at the end of the glass rod, the china dish is removed from the tripod stand and allowed to cool in air. After some fine crystals have appeared, the china dish is placed in an ice-bath. When the crystallization is complete, the mother liquor is drained off, and the crystals in the china dish are washed with a very fine spray of ice cold water-ethyl alcohol mixture. The china dish is then kept slanting so that the excess of water gets drained out. The semi-dry crystals are then dried by pressing them gently between the folds of filter papers. The dry crystals are then weighed, and a sample is scaled in a polythene pack. The packet is then stapled in your record book.

Results. The practical yield of the Mohr's salt is gram.

The colour of the Mohr's salt crystals is (light green)

The shape of the Mohr's salt crystals is (monoclinic)

Percentage yield. Since, it is not always possible to recover the stoichiometric amount of the salt from the solution by crystallization, hence success of any preparation is judged from the percentage yield, which is defined as,

Percentage yield = Experimental yield in grams × 100
Stoichiometric (expected) yield in grames

Precautions. (i) Do not use too much of the solvent while preparing the saturated solution of the salt(s).

- (ii) Do not boil the solution for a long.
- (iii) Do not concentrate the solution too much.
- (iv) Cooling of the saturated solution should be slow.

(v) During drying, do not press the crystals hard. Points to Remember. (f) Sulphuric acid is added to prevent the hydrolysis of ferrous ion.

(ii) Noiling the solution for a long time results in partial exidation (1



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GBSHSE/ACAD/ 2012/ 1492

Date: 20-06-2012

Υo,

The Heads of all recognised Higher Secondary Schools, within the jurisdiction of this Board.

Sub:- Distribution of portion in Marathi Lang. I, Marathi Lang.II, Konkani Lang.II, Hindi Lang.II Urdu Lang.II & Sanskrit Lang for II for Std XII.

Sir/Madam,

This is in continuation of the circular no.21 dated 26th April, 2012. Please find enclosed herewith the Distribution of portion in Marathi Lang. I, Marathi Lang.II, Konkani Lang.II, Hindi Lang.II, Urdu Lang.II & Sanskrit Lang for II Std XII.

The contents of the letter may be brought to the notice of the Teachers concerned.

Yours faithfully,

(Dr. Thomas Mathew)
Joint Secretary

Encl: As above



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Subject: MARATHI Lang. I Std XII

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First -Term Portion
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जृत्तान्तलेखन, पत्रलेखन, गर्णा, निर्देध
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Second -Term Partion
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उत्तरः ,रसरहरः



DISTRIBUTION OF PORTION

Subject: SANSKRIT Lang. II S. 4 XII

Term wise portion distribution

A. First midterm examination (20 marks)

- 1. Text Lessons from 1 to 3
- 2. Grammar- सन्धिः, समासः (अव्ययीभावः, तत्पुरुषः, कर्मधारयः, द्वन्द्वः), पर्यायवाचिनः, शब्दरूपाणि, धातुरूपाणि, टाप्, कारकम्, प्रश्ननिर्माणम्
- 3. Non Textual

Short notes on 1. वैदिकसाहित्यम् 2. डॉ. श्री. भा. वर्णेकरः 3. भासः अपठितगद्यस्य अनुवादः

4. संवादपूर्तिः

B. 1st Terminal Examination (80 marks)

- 1. Text- Lesson from 1 to 6
- 2. Grammer- सन्धिः, समासः (अव्ययीभावः, तत्पुरुषः, कर्मधारयः, द्वन्द्वः, द्विगुः, बहुव्रीहिः), पर्यायवाचिनः, शब्दरूपाणि, धातुरूपाणि, टाप्,डीप्, ठक्, तल्, कारकम्, प्रश्निर्माणम्, कालपरिवर्तनम्, प्रयोगः
- 3. Non Textual

Short notes on 1. वैदिकसाहित्यम् 2. डॉ. श्री. भा. वर्णेकरः 3. भासः 4. कालिदासः, 5. अम्बिकादत्तव्यासः अपिठतगद्यस्य अनुवादः, अनुच्छेदाधारिताः प्रश्नाः

4. चित्रवर्णनम्, पत्रलेखनम् , संवादपूर्तिः

C.II terminal Examination (80 marks)

Text-lessons from 7 to 10

Core content- 20 % of the First Terminal Portion

(1 Prose & 1 poetry)

Grammer-सन्धिः, समासः (अव्ययीभावः, तत्पुरुषः, कर्मधारयः, इन्द्रः, द्विगुः, बहुव्रीहिः), पर्यायवाचिनः, शब्दरूपाणि, धातुरूपाणि, टाप्,डीप्, ठक्, तल्, मतुप्, कारकम्, प्रश्ननिर्माणम्, कालपरिवर्तनम्,वचनपरिवर्तनम्, वाक्यसंशुद्धिः, प्रयोगः



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Subject:- HINDI Lang. II

Std XII

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Subject :- URDU Lang.II

Std XII

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1) Suggested Assignment / Project (any One)

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2) The criteria for the evaluation of the assignment should based on following:

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First -Term Portion
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व्याकरपः शब्दांच्या जाती, भाववाचक नाने , दिरुखाधी शब्द, दिशपण,
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