

# THE OPEN UNIVERSITY OF SRI LANKA B.Sc DEGREE/STAND ALONE COURSE IN SCIENCE - LEVEL 5 Final Examination—2015/2016

CHU 3238/CMU 3233 - POLYMER CHEMISTRY

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# THE OPEN UNIVERSITY OF SRI LANKA

### B.Sc. Degree Program -Level 5

## Final Examination -2015/2016

# CMU3233/CHU3238-Polymer Chemistry

Date: 21.01.2017

9.30am-12.30pm

### Instructions to candidates:

- This paper consists of two parts, Part I (MCQ) and part II (essay type).
- Part I consists of 25 MCQs, recommended time to complete this part is 1 hour.
- Part II consists of six essay type questions. You are expected to answer **four** questions among six. Recommended time to complete this part is 2 hours.

#### For MCOs

- Answer all questions.
- Choose the most correct answer to each question and mark a cross" X" over the answer on the given answer sheet.
- Use a PEN (not a pencil) in answering.
- Any answer with more than one cross will not be counted.
- 1/6<sup>th</sup> marks will be deducted for each incorrect answer
- The use of a non programmable electronic calculator is permitted.

PART I (35 mark				
01. What is the 1) 1	functionality of O	H-CH <sub>2</sub> -CH <sub>2</sub> -COOH		•
1) 1	2) 2	3) 3	4) 5	
02. What polys 1) Cationic 5) Free radical and	2) Anionic	can be applied to po 3) Free radical	olymerize viny 4) Anionic ar	
<ol> <li>Three basic step</li> <li>Repeating units</li> <li>Monomer conce</li> <li>By products such</li> </ol>	s are involved in po are added one at a ntration drops stead h as H <sub>2</sub> O are forme	time. dily throughout the i	reaction.	
04. HClO <sub>4</sub> can 1) cationic initiator 5) Zeiglar Natta cat	2) anionic initi	ator 3) free radica	al initiator	4) inhibitor
05. Ethylene an 1) same physical pr 4) same solubility.	d polyethylene hav coperties. 2) 5) different p	same melting point	s. 3) sar other.	me boiling points.
06. Above flow 1) glassy state. 5) visco-fluid state.	2) rubbery state.	he polymer is in the 3) visco-elas	tic state.	4) solid state.
07. Degree of co	rystallinity (χ) can l	be expressed as		
1) $\chi = \frac{Density of \{1\}}{Density of \{1\}}$	semicrystalline – 10 00%crystalline – 1	00%amorphous} pol 00%amorphous} pol	lymer lymer	
2) $ \chi = \frac{Densityof\{}{Densityof\{1\}} $	100% crystalline – 5	semicrystalline} poly	ymer	
3) $\chi = \frac{Densityof\{1\}}{Densityof\{s\}}$	00%crystatime – 1 00%amorphous – s	00%amorphous} pol semicrystalline} poly	lymer vmer	
Density of {s	emicrystalline – 10 00% crystallina – 11	0%amorphous} poly	vmer	
4) $\chi = \frac{Density of \{1\}}{Density of \{5\}}$	semicrystalline – 10	00%amorphous} pol	vmer	
5) $\chi = \frac{Density of \{s\}}{Density of \{1\}}$	emicrystalline – 10 00%crystalline – se	0%crystalline} polyi emicrystalline} polyi	mer mer	
08. Density of soll) Dilatmetry. 5) End group analys	2) Viscometry.	ymers can be determ 3) X-Ray diffra	_	4) Cryoscopy

a) Polyethylene	er following poly e. b) Polyp Illowing polymei		c) Polystyrene  active?	
1) a only.				5) b and c only.
1) polymer mol	ar mass only.	ation depends on 2) rate of mix 5) all of abov	ing only. 3) n	ature of the polymer only.
1) weight avera	ge molar mass.	ed to determined 2) number avelative viscosity.	verage molar mass.	3) specific viscosity.
a) It is a separat	r following state ion technique. a poly dispersed	b) Chromato	Permeation Chroma graphic column is fi	ntography? illed with non-porous beads.
The correct state 1) a only.		3) c only.	4. a and b only.	5. all of above.
13. In molai poly dis <sub>l</sub>	mass distribution mass distrib	on curve, relative sample can be rep	e portions of various	s average molar masses of a
1) $\bar{M}_n > \bar{M}_V > 1$	$\bar{M}_W$ 2)	$\bar{M}_W > \bar{M}_V > \bar{M}_R$	3. $\bar{M}_n =$	$= \bar{M}_V = \bar{M}_W$
4) $\bar{M}_V > \bar{M_W} >$	$\bar{M}_n$ 5. $\bar{M}_{\bar{\nu}}$	$v = \bar{M}_W > \bar{M}_n$		
a) It is compatib	r following states ble with rubber. as an accelerator.	b) Can I	ZnO in rubber vulca be used as an activa	unization process. tor.
The correct state 1) a only.		3) c only.	4. a and b only.	5. all of above.
	degradation may radiation.	y occur by 3) mechanical str	ress. 4) atmo	ospheric oxygen.
<ol> <li>They are irrev</li> <li>Pre-polymers</li> <li>Final product</li> </ol>	versible reactions having reactive obtained are ins phenol-formalde	functional groups	s undergo these reac	ctions.

- 17. What statement is **not true** about polystyrene? 1) It is produced by free radical polymerization.
- 2) It is produced by coordination polymerization.
- 3) It has a low softening point.
- 4) It is insoluble in aromatic hydrocarbons.
- 5. It is widely used as a packing material.
  - 18. Extent of polymerization is given by

1) 
$$1 - \frac{N_0}{N}$$
 2)  $1 + \frac{N_0}{N}$  3)  $1 - \frac{N}{N_0}$  4)  $1 + \frac{N}{N_0}$  5)  $\frac{N}{N_0} - 1$ 

2) 
$$1 + \frac{N_0}{N}$$

3) 
$$1 - \frac{N}{N_0}$$

4) 
$$1 + \frac{N}{N_0}$$

5) 
$$\frac{N}{N_0} - 1$$

19. A-B represents a monomer having a COOH group (A) and OH group (B). When extent of polymerization is 95%, the number average degree of polymerization is,

1) 35

20. Rate equation for anionic polymerization of styrene is

1. 
$$\frac{-d[M]}{dt} = \frac{k_{tr}[NH_3]}{k_{p}[I][M]}$$
2. 
$$\frac{-d[M]}{dt} = \frac{k_{i}[M]^2}{k_{p}[NH_3]}$$
3. 
$$\frac{-d[M]}{dt} = \frac{k_{i}k_{p}[I][M^2]}{k_{tr}[NH_3]}$$
4. 
$$\frac{-d[M]}{dt} = \frac{k_{p}[NH_3]}{k_{i}[I]}$$
5. 
$$\frac{-d[M]}{dt} = \frac{k_{tr}[I]^2[M]}{k_{i}||NH_3||}$$

2. 
$$\frac{-d[M]}{dt} = \frac{k_i[M]^2}{k_p[NH_3]}$$

3. 
$$\frac{-d[M]}{dt} = \frac{k_i k_p[I][M^2]}{k_{II}[NH_3]}$$

$$4. \frac{-d[M]}{dt} = \frac{k_p[NH_3]}{k_i[I]}$$

$$5. \frac{-d[M]}{dt} = \frac{k_{tr}[I]^{2}[M]}{k_{t}||NH_{3}||}$$

21. Rate equation for cationic polymerization is

1. 1st order w.r.t monomer concentration.

2. 2<sup>nd</sup> order w.r.t. monomer concentration.

3. 1st order w.r.t. catalyst concentration.

4. 0<sup>th</sup> order w.r.t. monomer concentration.

5. independent of catalyst concentration.

22. Beilstein test is used to identify

1) HDPE

2) PP

3) PVC

4) LDPE

5) PE

23. What statement is true about Raman spectroscopy?

- 1) It is a very popular method than IR spectroscopy in polymer characterization.
- 2) Raman spectra are indicative of asymmetric bond stretching and bending.
- 3) It can be used to determine the composition of copolymers.
- 4) It can be used to identify tacticity of PP.
- 5) It can be used to determine the crystallinity of polymers.

24. Melt flow index (MFI) of a given polymer is not affected by

1) Molecular weight distribution.

2) The degree of chain branching.

3) Crystallinity.

4) Viscosity.

5) Type of inter molecular forces.

25. Consider following parameters obtained from a thermogram of a given polymer.

a) The enthalpy of crystallization and melting.

b) Glass transition temperature.

c) Intrinsic viscosity.

The correct parameter/s is/are

1) a only.

2) b only.

3) c only.

4) a and b only.

5. none of above.

## PART II (65 marks)

01. a) i. What is meant by functionality of a monomer?

ii. Explain the differences among mono functional, bi functional and tri functional monomers by giving examples.

b) i. What are the different modes of additions in chain polymerization of CH<sub>2</sub>=CH-CH<sub>3</sub>. Explain.

ii. Which one is the most favored mode of additions among above mentioned modes? Give reasons.

c) i. What are the constituents of Ziegler-Natta catalyst?

ii. How does CH<sub>2</sub>=CH-CH<sub>3</sub> monomer bind to the active center of Ziegler-Natta catalyst through mono metallic mechanism? Explain by drawing the structure of the transition state.

d) i. What are copolymers?

ii. "Properties of polymers can be changed by copolymerization with other monomers". Justify this statement by taking appropriate examples.

e) i. Describe the chemical process of solution polymerization.

ii. State two major disadvantages of this technique over other polymerization techniques.

02. a) i. What is meant by stereoregular polymers? Give examples.

ii. Explain different types of stereoisomers arise due to asymmetric carbon.

iii. "Polypropylene oxide shows optical isomerism". Justify your answer by drawing the structure.

b) Describe how specific volume of an amorphous polymer varies with temperature.

c) i. What is meant by glass transition temperature (Tg)? What is the importance of it? ii. Tg values of polyethylene and polystyrene are 188K and 373K, respectively. Explain the difference.

iii. What are plasticizers? Explain the relationship between plasticizers and the glass transition temperature.

iv. What is the major difference between amorphous and crystalline polymers?

d) Briefly explain the factors that affect the crystallinity of a polymer?

e) Polyisobutyl contains two chains of eight repeat units in each unit cell. Unit cell dimension of polyisobutyl is approximately 16Å×11Å×8Å. Calculate the density of 100% crystalline polyisobutyl.

03. a) i. How do you differentiate a monodispersed system from a polydispersed system? ii. A polymer sample is made up of a mixture having molar masses 12,000, 90,000 and 60,000 g/mol with molar ratio of 1:2:1, respectively. Calculate the number average molar

Mass,  $\bar{M_n}$  . If the weight average molar mass,  $\bar{M_W}$  of above polymer is 120,000 g/mol, calculate the polydispersity factor of the polymer sample.

b) i. Explain reasons for having slow solubility of polymers compare to simple compounds.

ii. What is meant by swelling of polymers?

iii. How do you enhance the swelling process of polymers? Explain.

iv. "Acrylonitrile is resistant for swelling in organic solvents". Explain.

c) i. Explain the difference among specific viscosity, intrinsic viscosity and relative viscosity.

ii. Describe an experimental technique to determine intrinsic viscosity.

- 04. a) i. What is meant by sulphur vulcanization?
  - ii. What are the essential ingredients necessary for above process? Explain their role.
  - iii. How do you obtain vulcanized rubber with high degree of monosulphide linkages?
  - iv. What are the importances of monosulphide linkages in vulcanized rubber?
  - b) i. What is meant by oxidative degradation? How does it happen?
    - ii. What precautions can be taken to overcome this degradation? Explain.
  - c) i. What is meant by autocoagulation of field latex? How does it happen?
    - ii. What is the purpose of adding bases to prevent autocoagulation? Explain.
    - iii. What are the common preservatives used in latex industry? Describe their functions.
  - d) What do you mean by yellow discolouration? Describe two ways of preventing it.
  - e) Write brief accounts on
    - i. Thermosets.
- ii. Foaming agents.
- iii. Fillers.
- iv. Antidegredants.
- 05. a) i. Derive the Modified Carother's equation given below starting from  $N_A$  and  $N_B$  as initial moles of molecules of the monomers type A-A and type B-B, available in reaction medium respectively. Define the terms in the equation below and consider  $N_A < N_B$ .

$$X_n = \frac{1+r}{1+r-2rP}$$

- ii. Percentage conversion of diacid-dialcohol polymerization reaction is 99.6%. If 8% of more diacid is used, calculate the degree of polymerization.
- b) How do you differentiate chain length of a polymer from kinetic chain length?
- c) i. What is meant by auto acceleration?
  - ii. Describe the kinetic interpretation of above phenomenon.
- 06. a) i. What is the basis of Infrared spectroscopy?
  - ii. How can it be used a qualitative tool to identify polymers? Explain.
  - iii. Describe the importance of FTIR as a quantitative tool in polymer chemistry.
  - iv. Following table summarizes absorbance values recorded at 2250 cm<sup>-1</sup> for nitrile stretching and 1600 cm<sup>-1</sup> for polystyrene ring stretching for number of styrene/acrylonitrile copolymers. For the same copolymer with unknown composition showed stretching frequencies at 2250 cm<sup>-1</sup> and 1600 cm<sup>-1</sup> are 0.205 and 0.121 respectively. Calculate the unknown composition of the copolymer.

% Acrylonitrile	Polystyrene stretching	Nitrile stretching		
concentration	frequency, cm <sup>-1</sup>	frequency, cm <sup>-1</sup>		
10	0.383	0.230		
20	0.177	0.223		
30	0.120	0.230		
40	0.091	0.235		
50	0.070	0.227		
60	0.059	0.231		

b) 75g of Polystyrene is mixed with 10g of plasticizers; the  $T_g$  was lowered from 373K to 273K. What would be the expected  $T_g$  when the amount of plasticizers increased up to 25g?