Sern II Organic Chem - III March CBSCIS QP Code: 16112 2015

(2½ Hours)

N.B.: 1. All Questions are compulsory

Total Marks: 70

2. Figures to right indicate full marks

	Q1. (a) Give IUPAC nomenclature of the following (any three):	(03)
	H ₃ C S COOCH ₃	
	H H ₃ CHN	
	HN HN	
	Br (ii) Et (iii)	
	(b) Compare and comment on basicity of imidazole and pyridine.	(02)
	(c) Explain disrotatory motion with example.	(01)
	(d) Explain the term Functional Group Addition with suitable example.	(01)
	(e) What is the strategy used for disconnection in pyrrole.	(01)
	(f) Give structures of the following in chair form:	(03)
	(i) 5α-progestane (ii) Cortisone (iii) 3β-hydroxy-6β-acetoxy-5α-androstane	(05)
	(g) Give product formed for reaction of Cholesterol with H ₂ O ₂ and justify	(01)
	(h) Draw reaction to depict classical and non-classical route for Friedel Craft alkylation.	
	(i) Define term Atom efficiency. If atom efficiency of a reaction corresponds to 40% then	
8	be its theoretical E-factor?	(02).
	be its disordical B-factor:	(02).
	02 (a) Write the following montions with mechanism (any two):	(04)
	Q2. (a) Write the following reactions with mechanism (any two):	(04)
	(i) Doebner Miller synthesis	
	(ii) Pomeranz - Fritsch synthesis	
	(iii) Fischer indole synthesis	
	(b) Using orbital diagram, explain whether $(4\pi + 2\pi)$ cycloaddition photochemical reactions of the second	
	suprafacial or antarafacial by giving suitable example.	(04)
	(c) Compare biocatalytic route of penicillin G with chemical synthesis.	(03)
		(0.4)
	Q3. (a) Attempt the following conversions (Any Four)	(04)
	(i) Pyrimidine to 5-bromopyrimidine	
	(ii) Pyridine to 2-phenylpyridine	
	(iii) 2,3-butanedione to 2,4,5-trimethylimi lazəle	
	(iv) Pyridine to 1,4-dihydropyridine	
	(v) 2,4,6-trichloropyrimidine to pyrimidine	7 a 13
	(b) Using synthon approach devise scheme for synthesis of atenolol.	(04)
	(c) Discuss classical and non-classical route for synthesis of Hydroquinone.	(03)
	Q4. (a) Write structures of products formed for the following reactions (any eight):	(08)
	(i) Furan + Maleic anhydride	
	(ii) Pyridine Liq.NH ₃	
	1.11 110110 111110	
	Thiophene fumily HNO3	
	(iv) Ac ₂ O	
	(v) Isoquinoline.	
	Ac20	- A"
	Pyrrcle Triethylamine	
	(vi) Triethylamine	
	Br ₂ , 130°C	4
	(VII) Pyrimidine	¥
	H ₂ SO ₄ /SO ₃	
	Imidazole	

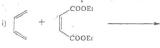
WG-Con.: 7652-15.

(ix) Quinoline Sulfur

[TURN OVER

QP Code: 16112

(b) Write structures of products formed in the following reactions:



Q5. (a) Write the following reactions with mechanism (any two):

(04)

(03)

- (i) Knorr pyrrole
- (ii) Hinsberg synthesis
- (iii) Paal Knorr synthesis for furan.

- (b) Give reasonable explanation for the following (Any seven) (i) Protonation of Nitrogen atom present in pyridine ring is easier than pyrrole.
 - (ii) Which is the preferred position for Electrophilic aromatic substitution reaction in imidazole and why?
 - (iii) Why pyrimidine (pKa: 1.30) is much less basic than pyridine (pKa: 5.2).
 - (iv) 5β-cholestane-3-one forms 4-bromo derivative.
 - (v) Acetates of 5α -cholestane- 3β -ol get more rapidly hydrolysed than 5α -cholestane- 3α -ol.
 - (vi) Cholesterol gives trans product upon oxidation with H₂O₂
 - (vii) Electrophilic substitution in quinoline takes place at 5 and &-position
 - (viii) Furan undergoes Diels Alder reaction whereas thiophene does not

Q6. (a) Draw resonating structures for the following

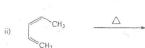
(04)

- (i) Pyridine
- (ii) Indole
- (iii) Imidazole (iv) Quinoline

(b) Explain following reactions with mechanism:

(04)





(c). Give simplest retrosynthetic pathway for the following (any 3)

- (i) p-methoxyacetophenone
 - (ii) Ethyl p-aminobenzoate
 - (iii) 1-butene
 - (iv) 2,4-dimethyl-2-pentanol

WG-Con.: 7652-15.