

# THE COLLEGES OF MEDICINE OF SOUTH AFRICA

Incorporated Association not for gain Reg No 1955/000003/08

#### Final Examination for the Fellowship of the College Of Pathologists of South Africa - Anatomical

## 27 February 2020

### Paper 1

(3 hours)

All questions to be answered. Each question to be answered in a separate book (or books if more than one is required for the answer)

- 1 a) In the interpretation of immunohistochemical assays
  - List potential causes of background or artifactual staining that you would consider for troubleshooting. (4)
  - ii) Tabulate the advantages and disadvantages of monoclonal and polyclonal antibodies in the interpretation of immunohistochemical assays. (4)
  - b) Writes notes on test validation in surgical pathology laboratory under the following subheadings
    - i) 3 parameters evaluated in qualitative test validation giving one example. (2)
    - ii) 5 parameters evaluated in quantitative test validation giving one example. (3)
  - c) About PCR in surgical pathology
    - i) List 8 major applications, giving some examples.
    - ii) List 4 reasons why false negative results may occur.
  - d) Discuss the reasons for the diagnosis of Atypical Small Acinar Proliferation (ASAP) in prostatic biopsies.
     (6)
    - [25]

(4)

(4)

(2)

- 2 a) With regards to medulloblastoma, discuss the following
  - i) Histologically defined classification.
  - ii) Genetically defined classification, including immunohistochemical stains utilised in this classification. (6)
  - b) In the routine evaluation of breast carcinoma, outline the reporting of oestrogen receptor, progesterone receptor and HER2 immunohistochemistry, detailing technical and interpretive aspects of these tests.
  - c) When performing an autopsy, what macroscopic findings might a pathologist encounter
    - i) In the heart in Systemic Lupus Erythematosus?
    - ii) In the urogenital tract in schistosomiasis?
    - iii) In the heart of a neonate with Tetralogy of Fallot?
    - iv) In the lung, in the 4 stages of lobar pneumonia?
    - v) In the small intestine and liver in Typhoid?

(2) [25]

(2)

(2)

(2)

(2)

3 Provide a panel of immunohistochemical markers you can use to distinguish renal clear a) cell carcinoma from the following tumours (14)i) Clear cell tumour (sugar tumour) of the lung. ii) Hepatic clear cell carcinoma. iii) Ovarian/endometrial clear cell carcinoma. b) Briefly discuss the biosafety principles for infection control during autopsies. (11)[25] 4 a) i) Discuss the molecular pathogenesis of Hereditary medullary thyroid carcinoma. (7) List 6 immunohistochemical stains expected to be positive in medullary thyroid ii) carcinoma. (3)Discuss the pitfalls of using proteolytic enzyme digestion antigen retrieval and microwave b) antigen retrieval in immunohistochemistry. (4) Describe the renal electron microscopic findings in post-infectious glomerulonephritis. (2) C) Discuss cell-rich subepidermal blisters under the following headings d) List 5 examples of a cell rich subepidermal blister. (2.5)i) ii) For each of the above examples, state the exact level of the basement membrane split which would be seen on electron microscopy. (3.5) e) List 6 histochemical stains that highlight Michaelis Gutmann bodies. (3)

[25]



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Paper 2

(3 hours)

All questions to be answered. Each question to be answered in a separate book (or books if more than one is required for the answer)

1	a) b)	Write notes on the glomerular capillary endotheliosis under the following subheadingsi)Definition.ii)Patho-mechanism.iii)Microscopic features.iv)Electron microscopic appearances.v)Immunofluorescence.v)Immunofluorescence.Write notes on lymphomatoid granulomatosis under the following subheadings
		i)Definition.(3)ii)Microscopic features in the lung.(3)iii)Grading.(2)
	c)	List the most useful cytomorphologic features in the diagnosis of malignancy in bile duct brushings specimens. (7)
	d)	List four conditions associated with nodular regenerative hyperplasia of the liver. (2) [25]
2	a)	Compare ovarian low-grade serous carcinoma and ovarian high-grade serous carcinoma under the following headings
		i)Origin, including precursor lesion.(2)ii)Molecular abnormalities.(2)
		iii) Prognosis and response to chemotherapy. (2)
	b)	List the conditions in which squamous metaplasia of endometrial glands is most likely to be found. (2)
	C)	Discuss atypical polyploid adenomyoma. (3)
	d)	Outline your approach to a spindle cell lesion of the bladder with cytologic atypia in an adult patient with regards to
		<ul> <li>i) Differential diagnosis. (3)</li> <li>ii) Potentially useful immunohistochemical stains. (3)</li> </ul>
	e)	Discuss the use of the immunohistochemical marker CD30 in the evaluation of lymphomas. (3)
	f)	Discuss the grading of soft tissue sarcomas. Including potential pitfalls of this grading system. (5) [25]

PTO/Page 2 Question 3...

- 3 a) Write short notes on intraductal papillary mucinous neoplasm (IPMN) under the following headings
  - Gross and light microscopic features. (6) i) Molecular markers/mutations. ii) (2)
  - Differential diagnosis and name distinguishing feature. iii)
  - (1) Briefly discuss Castleman disease, hyaline vascular type under the following headings b) Microscopy. (5) i)
    - Immunohistochemistry. ii)

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- (3) Write short notes on the gross and microscopic features of goblet cell adenocarcinoma C) (goblet cell carcinoid). (8)
  - [25]

a)	Describe NUT carcinomas under the following headings	
	<ol> <li>List the known molecular aberrations of this malignancy.</li> </ol>	(2)
	ii) Describe the characteristic light microscopic features.	(4)
	iii) Compare and contrast the immunohistochemical profile of NUT carcin	nomas and
	SMARCB1 deficient sinonasal carcinoma.	(3)
b)	atic tumour	
,	(PHAT).	(5)
c)	Describe the Weiss criteria for the classification of adrenal cortical tumours.	(4)
d)	Describe the light microscopic features of chondromyxoid fibroma.	(4)
e)		
,	i) List the two (2) HLA haplotypes associated with Coeliac disease.	(1)
	ii) List one (1) antigenic environmental trigger of this condition.	(0.5)
	iii) List the factors assessed in the Modified Marsh-Oberhuber classification	
	disease.	(1.5)
		[25]