

FINAL EXAM DECEMBER 2017

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MEDICAL GENETICS PAPER-III

GENE/D/17/53 /III

TIME : 3 HOURS MAX. MARKS : 100

- Attempt all questions in order.
- Each question carries 10 marks.
- Read the question carefully and answer to the point neatly and legibly.
- Do not leave any blank pages between two answers.
- Indicate the question number correctly for the answer in the margin space
- Answer all the parts of a single question together.
- Start the question to a question on a fresh page or leave adequate space between two answers.
- Draw table/diagrams/flowcharts wherever appropriate.

Write Short notes on:

1.	b.	What are lysosomal storage disorders (LSD's)? What is ERT (Enzyme-replacement therapy) for LSDs? What are novel strategies/ other strategies to treat LSDs?	4+3+3
2.		What are the pros and cons of stem cell banking? How will you counsel a couple whose previous baby has beta-thalassemia and now they have come to gets your advice for stem cell banking in future pregnancies?	5+5
3.		What are SiRNAs and their mechanisms of action? What is the role of SiRNA in therapeutics?	5+5
4.	b.	What are the different principles and methods used for NGS (Next- generation sequencing)? Make a flow chart showing principle of NGS data analysis. When will you prefer Exome sequencing by NGS over Sanger Sequencing for single gene disorders?	3+3+4
5.		What are the different 'in-silico' tools used for pathogenicity testing of a sequence variation? What are the guidelines you will follow and how will you denote the pathogenicity of sequence variation?	6+4
6.		What are different gene-transfer vehicles used in gene therapy? Name some disorders amicable to gene therapy.	6+4



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7.		How will you investigate a child with suspected DMD (Duchenne muscular dystrophy) What are the recent treatment strategies for DMD?	6+4
8.	b. c.	What is pharmacogenetics? How do genes affect the metabolism of Warfarin? What are the fetal effects of maternal warfarin intake? Name some teratogenic drugs and mention their effects on fetus.	2+2+2+4
9.	ра	hat are the functional studies used to detect thogenicity of a novel sequence variation detected in a nogle gene disorder?	10
10.		What is osteogenesis imperfecta (OI)? What are the inheritance patterns? How will you investigate? How will you treat a child with OI?	6+4