भारत सरकार GOVERNMENT OF INDIA MINISTRY OF SCIENCE AND TECHNOLOGY



CORRIGENDUM

Fellowship in Genetic Diagnostics

CALL FOR APPLICATIONS

With an aim to expand National Inherited Disorders Administration Kendras (NIDAN Kendras) to provide comprehensive clinical care including diagnosis, management, multidisciplinary care, counseling, prenatal testing at Government Hospitals/ Medical Colleges, Department of Biotechnology (DBT), Government of India has launched a 6 month 'Fellowship in Genetic Diagnostics' for Faculty/ Clinicians only from Government Medical Colleges and Hospitals. Four (4) trainees in a year (in two batches) will be taken by each centre.

Advertisement in this regards has been issued on 1st July 2019.

The **REVISED** list of the centres where this 'Fellowship in Genetic Diagnostics' is available may be noted as follows:

Sr No	Name of the Centre	Thrust areas	Clinical services	Laboratory facilities	Number of seats	Contact address of Training Co-ordinator
					per year	U
1.	Department of Medical	Birth defects, Genetic	Clinical genetics,	Cytogenetics, Cell		Dr Shubha Phadke,
	Genetics, Sanjay Gandhi	Hematology,	Prenatal Diagnosis,	Culture, Molecular		Head, Department of
	Postgraduate Institute of	Dysmorphology, IEM,	Disease Mangement,	Cytogenetics including		Medical Genetics
	Medical Sciences,	Skeletal Dysplasias,	Fetal malformation	Microarray, MLPA,	4	shubharaophadke@gma
	Raibarelly Road,	Neurological and muscular	diagnosis,	Molecular Genetics,		<u>il.com</u>
	Lucknow, 226014 [UP]		Reproductive	Sanger Sequencing,		0522 249 4334 / 4325

	क्षिक्तत्तर आसून्य विव स्थान राष्ट्र स्थान राष्ट्र स्थान स्यान स्थान स्थान स्थान स्या स्यान स्यान स्या स्यान स्या स्या स्या स्या स्या स्या स्या स्या	disorders, Clinical Genetics	genetics, Genetic counseling, Management of genetic disorders, Fetal autopsy	NGS Analysis, Enzyme Assays, Newborn screening		
2.	Department of Haematology, Christian Medical College, Vellore, Tamilnadu	Hemoglobinopathies, Disorders of Haemostasis, Bone marrow Failure syndromes Rare haematological disorders, Primary immunodeficiency Diseases	All inherited haematological disorders-Diagnosis and Management including Haematopoietic Stem cell transplant	State of the art molecular lab, Molecular Genetics- Identification of genetic variants, Sanger Sequencing, NGS Analysis, Interpretation of results and clinical correlation, Prenatal Diagnosis	4	Dr. Eunice Sindhuvi Department of Haematology eunice@cmcvellore.ac.i n 0416 -228- 3569/3577
3	Department of Clinical Genetics Christian Medical College, Vellore, Tamilnadu	Dysmorphology, Rare syndromes, Neuromuscular and neurodegenerative disorders, Genodermatosis, Cardiac and Renal, Autoimmune, Multifactorial disorders	Clinical genetics, Prenatal Diagnosis, Adult Genetics, Disease Management	Molecular Genetics laboratory for genetic variants by Sanger,genescan, NGS, Bioinformatics, Protein analysis, Relevant enzyme assays	4	Dr. Sumita Danda Head, Department of Clinical Genetics. sdanda@cmcvellore.ac. in 0416-228-3161/2304
4	Centre for Genetic Studies and Research, The Madras Medical Mission, Chennai – 600037, Tamilnadu	Cardiac genetics (Paediatric and adult), Reproductive genetics, Common genetic disorders /syndromes, Rare genetic diseases	Clinical genetics – prenatal, postnatal and adult genetics. Counselling and Management of genetic disorders.	Cell culture, Cytogenetics, Molecular cytogenetics (FISH), Aneuploidy screening and testing, , Sperm aneuploidy test (SAT),	4	Dr.Bibhas Kar Consultant and Head Centre for Genetic Studies and Research drbibhaskar65@gmail.c om

				Single gene disorder testing		0 44 – 26561801 / 4259
5	Diagnostics Division, Centre for DNA Fingerprinting and Diagnostics, Hyderabad, Telangana	Cytogenetic testing for chromosomal disorders and copy number variations, Molecular testing for Thalassemia, DMD, SMA, other single gene disorders, NGS panel testing, Whole exome/ genome sequencing, enzyme assay for lysosomal storage diseases, inborn errors of metabolism testing	Not available	Cytogenetics, Cell Culture, Molecular Cytogenetics including FISH, Microarray analysis, Molecular Genetics, Sanger Sequencing, MLPA, Genotyping, NGS Analysis for whole exome/genome, Enzyme Assays	4	Dr Ashwin Dalal Head, Diagnostics Division, Centre for DNA Fingerprinting and Diagnostics adalal@cdfd.org.in 040-27216147/6148
6	Division of Genetics, Department of Pediatrics, AIIMS, New Delhi 110029	Dysmorphology, IEM, LSD, Skeletal dysplasia, birth defects, Neurodevelopmental disorders	Management of IEMs Enzyme replacement therapy for LSDs Genetic counselling And Prenatal diagnosis Fetal autopsy	Cytogenetics, Cell Culture, Molecular Genetics, Sanger Sequencing, MLPA, QF PCR, Real time PCR, Bioinformatics Analysis (NGS, CMA), Enzyme Assays, HPLC, Prenatal screening and testing	4	Dr Madhulika Kabra Dr Neerja Gupta madhulikakakbra@hot mail.com, neerja17aiims@gmail.c om

7	ICMR-National Institute of Immunohaematology, 13th Floor, KEM Hospital, Parel, Mumbai, 400015	Primary Immunodeficiency Diseases, Transfusion medicine, Hemoglobinopathies, Red cell membranopathies and enzymopathies,	Specialized diagnostic services for various inherited haematological and immunological disorders, Carrier detection and Prenatal diagnosis	Sanger sequencing, Cytogenetics, Karyotyping and FISH, HPLC system, Cell culture facility, Flowcytometry, Red cell enzyme assays, Automated	4	Dr Manisha Madkaikar, Director, madkaikarmanisha@ya hoo.co.in directorniih@gmail.co m 02224132928 022 24138518/19
	ICMS INDIAN COUNCIL OF MEDICAL RESEARCH NATIONAL INSTITUTE OF IMMUNOHAEMATOLOGY	Haemophilia and other inherited bleeding disorders, Inherited thrombotic disorders, Cytogenetics	Trenatar diagnosis	coagulometer and thromboelastograph		022 24136316/17
8	Division of Medical Genetics & Metabolism, Genetic lab, Department of Pediatrics, Maulana Azad Medical College, New Delhi	IEM, Newborn Screening, Dysmorphology, Skeletal Dysplasias, Neurological and muscular disorders, Clinical Genetics, Neurotramsmitter disorders	Clinical genetics, Prenatal Diagnosis, Disease Mangement, Emergency sick day IEM management	Mass spectrometry, Cytogenetics, Cell Culture, Molecular Cytogenetics, Molecular Genetics, Sanger Sequencing,, Enzyme Assays, Newborn screening & Confirmation	4	Dr Seema Kapoor, Head , Division of Medical Genetics drseemakapoor@gmail. com 011232397417, 01123221927

The other details of the advertisement will remain same.

Last date for submission of applications: 31st July 2019.