भारत सरकार GOVERNMENT OF INDIA विज्ञान और प्रौद्योगिकी मंत्रालय MINISTRY OF SCIENCE AND TECHNOLOGY



Fellowship in Genetic Diagnostics

CALL FOR APPLICATIONS FOR 3rd BATCH OF TRAINEES

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.

In the current era of genomics, genetics has become an integral component of medicine. This platform would offer a great opportunity to Govt. Hospitals and Medical Colleges by training their permanent faculty, physicians and scientists who are keen to develop Genetic Facilities to cater the need of patients. The program will provide training in cytogenetic and molecular genetic techniques along with the understanding about their applications in patient care. Under this program well-established centres with facilities for diagnosis and management of genetic disorders have been selected to provide training. The list of centres where the Fellowship is available is given at the end. Interested and eligible candidates should send application to the centre of their interest in the format provided below. **Applicant should not apply to more than two centres at a time.** The fellowship will provide an opportunity to get training in the exciting field of genetics as applied for patient care. Further, the faculty members trained under this program will be eligible to apply for DBT grant for Establishment of National Inherited Disorders Administration Kendras (NIDAN Kendras) at their respective institutes/colleges/hospitals after successful completion of their training.

Eligibility and Mode of selection of trainees: Post-graduate degree (MD/MS/DNB) in Pediatrics, Medicine or Obstetrics & Gynaecology, Pathology, Microbiology, Biochemistry, Laboratory Hematology, Laboratory Medicine, Anatomy, Physiology, Dermatology, Hematology, Radiotherapy, Endocrinology, Ophthalmology, Oncology or any other clinical / paraclinical specialty/Superspecialty and holding regular position in Government medical college/hospital.

Mode of selection: Screening and selection based on scrutiny of Application form and telephonic interview. The commitment to learn and the possibilities of implementation at the workplace will be judged.

Award Support: The trainee/clinicians from outstation institutes will be paid a displacement allowance of Rs 30, 000 per month. The trainee/clinicians from outstation institutes will be paid a displacement allowance of Rs 30, 000 per month' (The trainee will be eligible for the displacement allowance. Only if the trainee is not from the city where the training is planned)

Fellowship Schedule: 3rd Batch from July 2021 to December 2021

Fellowship of 3rd Batch will start from 1st July 2021.

Last date for submission of applications: 31st march 2021.

How to Apply:

Application form (in the format given below) along with self-attested photocopies of degree certificates, MCI registration certificate and experience certificates should be sent to the Training Co-ordinator of particular institution.

The application should be compulsorily forwarded by the Head (Director / Principal) of the institute.

Soft copy of the application is to be sent by email to the Training Co-ordinator of the concerned centre (s) and it should be followed by duly forwarded hard copy of the application along with copies of necessary certificates (If there is delay in submitting the application through proper channel, candidate can submit advance copy within last date of application to consider candidature; and application through proper channel can be submitted so as to reach to the concerned training centre coordinator by 10th April 2021). While sending soft copy of the application to the concerned centre, copy of the email may please also be marked to Dr. Onkar N. Tiwari, Scientist 'E', Department of Biotechnology, Govt. of India in the email ID: onkar.dbt@nic.in for information.

LIST OF THE CENTRES WHERE THE FELLOWSHIP IN GENETIC DIAGNOSTICS IS AVAILABLE IS AS FOLLOWS:

Sr	Name of the Centre	Thrust areas	Clinical services	Laboratory facilities	Number	Contact address of
No					of seats	Training Co-ordinator
					per year	
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@gm
		Neurological and muscular	diagnosis,	Molecular Genetics,		<u>ail.com</u>

	Raibarelly Road, Lucknow, 226014 [UP]	disorders, Clinical Genetics	Reproductive genetics, Genetic counseling, Management of genetic disorders, Fetal autopsy	Sanger Sequencing, NGS Analysis, Enzyme Assays, Newborn screening		0522 249 4334 / 4325
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	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	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
5	Division of Genetics, Department of Pediatrics, AIIMS, New Delhi 110029	metabolism testing 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

6	ICMR-National Institute of Immunohaematology, 13 th Floor, KEM Hospital, Parel, Mumbai, 400015	Primary Immunodeficiency Diseases, Transfusion medicine, Hemoglobinopathies, Red cell membranopathies and enzymopathies, Haemophilia and other inherited bleeding disorders, Inherited thrombotic disorders, Cytogenetics	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 coagulometer and thromboelastograph	4	Dr Manisha Madkaikar, Director, madkaikarmanisha@ya hoo.co.in; directorniih@gmail.co m 02224132928 022 24138518/19
7	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

Last date for submission of applications: 31st march 2021

Application form is on the next page

Application form for the Fellowship in Genetic Diagnostics

1.	Full Name (in Block lette	rs) :			
	(As per SSC / X Class Reco	ord)			Stick passport size
2.	Gender:				photograph
3.	Date of birth:		Age:		
4.	Qualifications:				
5.	Medical council registrat	tion number:			
6.	Present designation:				
7.	Complete address:				
8.	Contact number(s):		Email id:		
9.	Educational qualification	ns (starting with graduation):		
	Degree	College/ University		Year of passing	Awards/ distinctions/ honours

attach the list

13. Mention briefly (in not more than 250 word medical practice:	ds) your reasons for applying for this fellowship a	nd how you think it would help you in your
Date: Place:	Signature of the applicant	
Comment by forwarding authority:		
Director / Principal / Dean of the Institute / Medical	College (Signature and Stamp)	