

thsti ब्रिक—ट्रांसलेशनल स्वास्थ्य विज्ञान और प्रौद्योगिकी संस्थान



BRIC-Translational Health Science and Technology Institute

(An Institute of the Biotechnology Research and Innovation Council, Govt. of India) NCR Biotech Science Cluster, 3rd Milestone, Faridabad – Gurugram Expressway, P.O. Box No. 04, Faridabad - 121001

Rolling Recruitment Notice No.: THS/RN/01/2025/02-II

VACANCY NOTIFICATION

S.	Name of the	Essential Qualification	Job Description	Monthly
No.	post /	and Experience		Emoluments
	No. of post/			
	Age limit			
Proje	ect: Development	of highly infectious BSL4 v	riral agent's pseudo viruses ar	nd its applicability
			ic and emerging zoonotic virus	
PΙ	: Dr. Sweety S			
1.	Research Associate-I	PhD in any branch of Life Sciences from a recognised University.	 Work experience in the area of Virology, Cell culture and immunological assays. 	Rs. 58,000/- + HRA
	One post		Live virus handling and	
	35 years	Desirable: One first author publication	neutralization assay. • Experience in Molecular	
	Date for walk-	in reputed journal.	biological techniques and	
	in interview- 3rd		animal handling.	
	March 2025		 Experienced in different protein purification methods using FPLC / HPLC. Basic understanding and handling of Flow cytometer, confocal and computational biology skills. Data analysis and interpretation. Preference will be given to candidates having work experience in BSL3/ABSL3 	
Duni	at Tanatina Tu	o Company of Circulling of	facility.	
Proje		o Component Signalling sy the tubercle bacilli	stems (TCSs) in M. Tuberculo	osis to counteract
ΡI	: Dr. Ramande			
2.	Project	Master's degree in Life	Experiences in doing DNA	i) Rs. 35,000/-
	Associate-II	Sciences from a recognized	cloning, protein purification,	+HRA Scholars
	One post	University with atleast two (2) years of post-	western blots, FACS experiments and biophysical techniques.	who are selected through (a) National Eligibility
	35 years	qualification research experience.	The candidates should also have experience in handling animals. The candidates should	Tests – CSIR – UGC NET including lectureship or

Proje			also have performed basic microbiology techniques such as growth curves, CFU analysis etc. I, Neonatal and Infant Science Advanced Research on Birth	
DT	INdia Initiati	ve (GARBHINI Phase-II)		
9I 3.	: Dr. Shinjini Bl Senior Project	natnagar/ Dr. Nitya Wadhv Master's degree in		Rs 42 000/- ± HDA
) .	Associate	natural/Agricultural/ Life	ensure that the trial is being	113. TZ,UUU/- + FIKA
		Sciences/M.V.Sc. or	conducted in accordance with	
	One post	equivalent degree from a	the protocol, standard	
	40 4025	recognized University with	operating procedures, good	
	40 years	atleast four (4) years of	clinical practice, and	
	Date for walk-	post-qualification research	applicable regulatory	
	in interview-	experience in R&D.	requirements.	
	28 th February	OR Dectoral degree in Science/	Performs site monitoring throughout the trial which	
	28 th February 2025	Doctoral degree in Science/ Engineering/ Pharma or equivalent from a recognised university.	throughout the trial which involves visiting the trial sites on a regular basis (site initiation to site closeout) in accordance with contracted scope of work. Performs quality functions and executes quality programs (clinical operations, clinical laboratory) as per GCP/GCLP and regulations. Completes appropriate therapeutic, protocol and clinical research training to perform job duties. Setting up the trial sites such that each centre has the trial materials, including the trial drug while ensuring all trial supplies are accounted for. Administers protocol and related trial training to assigned sites and establishes regular lines of communication with sites to	

	T	<u> </u>	manage engoing project
			manage ongoing project
			expectations and issues.
			May provide training and
			assistance to junior clinical
			staff.
			Creates and maintains
			appropriate documentation
			regarding site management,
			monitoring visit findings and
			action plans by submitting
			regular visit reports and other
			required trial documentation.
			 Manages the progress of
			assigned studies by tracking
			regulatory/ IEC submissions
			and approvals, recruitment
			and enrolment, CRF
			completion and submission,
			and data query generation
			and resolution.
			Verifying that data entered on
			to the CRFs is consistent with
			participant clinical notes
			(source data/ document
			verification)
			Writing visit reports.
			Filing and collating trial
			documentation and reports.
			Archiving trial documentation
			and correspondence.
			Evaluate the quality and
			integrity of trial site practices
			related to the proper conduct
			of the protocol and adherence
			to applicable regulations.
			Escalates quality issues to the
			Quality Manager, Project
			Manager and/ or senior
			management.
			Work with Senior
			Management on other
			projects as directed and other
			internal departments on their
			requirements as and when
			required.
4.	Project	For Project Associate-I-	Overall responsibilities are to
	Associate-I/II	Master's degree in Life	
		Sciences or equivalent	conducted in accordance with i) Rs. 31,000/-
	Two posts	1	I ⊥HDΛ Scholars
1	Two posts		the protocol, standard

Date for walk-
in interview-
28th February

35 years

2025

degree from a recognized University.

For Project Associate-IIMaster's degree in Life Sciences or equivalent degree from a recognized University with two (2) years of post-qualification experience.

operating procedures, good clinical practice, and applicable regulatory requirements.

- Conduct regular site visits throughout the trial (from initiation to closeout) to monitor compliance with study protocols.
- Ensure trial sites are fully set up, with necessary materials, including trial drugs, and monitor trial supplies.
- Provide training on the protocol and related trial processes to assigned sites and maintain communication to manage project expectations.
- Monitor the progress of studies, including tracking regulatory approvals, recruitment, CRF completion, and data query resolution.
- Verify data consistency between CRFs and clinical notes (source data verification).
- Document site monitoring activities, visit findings, action plans, and submit regular reports.
- Evaluate trial site practices for protocol adherence and escalate quality issues to management.
- Assist with other projects as needed and support internal departments with trial-related requirements.

who are selected through (a) National Eligibility Tests – CSIR **UGC NET including** lectureship GATE or (b) A selection process through National level examinations conducted Central Department and their agencies and institutions)

ii) Rs.25,000/- + HRA for others who do not fall under (i) above.

For Project Associate-II

- Rs. 35,000/-+HRA Scholars who are selected through (a) National Eligibility Tests – CSIR UGC NET including lectureship GATE or (b) A selection process through National level examinations conducted Central Department and their agencies and institutions)
- ii) Rs.28,000/- + HRA for others who do not fall under (i) above

The interested candidates fulfilling the criteria as mentioned above may walk-in for written test/skill test/interview at 9:00 am on the dates mentioned against each positions at THSTI, NCR Biotech Science Cluster, 3rdMilestone, Faridabad-Gurugram Expressway, Faridabad – 121001.

NOTE: The candidates must bring their latest resume, one set of photocopy of documents in support of their educational qualification and experience along with originals and a valid ID card for verification. Candidates coming after the time slot mentioned will not be entertained.

GENERAL TERMS & CONDITIONS:

- 1. These are the short-term positions and extension will be granted subject to satisfactory performance of the incumbents and tenure of the project for which they are selected. Those appointed to these positions will not have any claim for regularization of their employment.
- 2. All educational, professional and technical qualification should be from a recognized Board/University.
- 3. The experience requirement specified above shall be the experience acquired after obtaining the minimum educational qualifications specified for the post. The candidates are required to satisfy themselves, before applying /appearing for the selection process, that they possess the minimum eligibility criteria as laid down in the recruitment advertisement. No query will be entertained with regard to eligibility criteria.
- 4. The date of Interview of the respective post will be the CRUCIAL DATE for determining eligibility with regard to age, experience, essential qualification etc.
- 5. Age and other relaxations for direct recruits and departmental candidates: 1. By five years for candidates belonging to SC/ST communities. 2. By three years for candidates belonging to OBC communities. 3. For Persons with Benchmark Disabilities (PwBD) falling under the following categories: (i) UR ten years, ii) OBC 13 years (iii) SC/ST 15 4. Age is relaxable for Central Government servants up to five years in accordance with the instructions or orders issued by the Central Government, from time-to-time. 5. Institute employees will get the age relaxation to the extent of the service rendered by them as on closing date of advertisement. 6. For Ex-servicemen upto the extent of service rendered in defence forces (Army, Navy & Air force) plus 3 years provided they have put in a minimum of 6 months attested service.
- 6. All results/notifications will only be published on our website. Therefore, the candidates should essentially visit THSTI website, regularly.
- 7. All communications will only be made through email.
- 8. In case a large number of applications are received, screening will be done to limit the number of candidates to those possessing higher/relevant qualification and experience.
- 9. The no. of vacancy indicated above may change subjected to the actual requirement at the time of Written test/skill test/interview.
- 10. With regard to any provisions not covered in this notification, the bye laws of THSTI / Govt. of India rules / guidelines shall prevail.
- 11. Canvassing wrong information in any form will be a disqualification.

"Government strives to have a work force which reflects gender balance and wome	≥n
candidates are encouraged to apply"	

	(M.V. Santo)
	Head-Administration
======================================	=========