



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

भर्ती नोटिस सं. : टीएचएस/आरएन/21/2026

दिनांक: 14 अक्टूबर 2026

RECRUITMENT NOTICE NO.: THS/RN/21/2026

Dated: 14th October 2026

भर्ती अधिसूचना/ RECRUITMENT NOTIFICATION

1. BRIC-Translational Health Science and Technology Institute (THSTI), जैव प्रौद्योगिकी अनुसंधान और नवाचार परिषद, जैव प्रौद्योगिकी विभाग, विज्ञान और प्रौद्योगिकी मंत्रालय, भारत सरकार का एक संस्थान है। भारत का यह संस्थान फरीदाबाद में स्थित इंटरडिसिप्लिनरी एनसीआर बायोटेक साइंस क्लस्टर का एक अभिन्न अंग है, जिसमें अभिनव ट्रांसलेशनल अनुसंधान करने और मानव स्वास्थ्य में सुधार के लिए अवधारणाओं को उत्पादों में ट्रांस्लेट करने के लिए विषयों और व्यवसायों में अनुसंधान सहयोग विकसित करने का मिशन है।

BRIC-Translational Health Science and Technology Institute (THSTI) is an Institute of the Biotechnology Research and Innovation Council, Department of Biotechnology, Ministry of Science & Technology, Govt. of India. The institute is an integral part of the interdisciplinary NCR Biotech Science Cluster located at Faridabad, with the mission to conduct innovative translational research and to develop research collaborations across disciplines and professions to translate concepts into products to improve human health.

2. ब्रिक-टीएचएसटीआई ने अनुसंधान और प्रयोगशाला कर्मचारियों की प्रशिक्षित टीमों द्वारा समर्थित उद्योग के साथ कई अंतर-संस्थागत सहयोग और कनेक्टिविटी का निर्माण किया है। टीएचएसटीआई ने विभिन्न केंद्रों की स्थापना की है जैसे (क) मातृ और बाल स्वास्थ्य केंद्र, (ख) वायरस अनुसंधान, चिकित्सा और टीका केंद्र (ग) तपेदिक अनुसंधान केंद्र (घ) माइक्रोबियल अनुसंधान केंद्र, (ङ) इम्युनोबायोलॉजी और इम्युनोथेरेपी केंद्र (च) ड्रग डिस्कवरी केंद्र (छ) नैदानिक विकास सेवा एजेंसी (ज) कम्प्यूटेशनल और गणितीय जीव विज्ञान केंद्र (झ) बायो-डिजाइन और निदान केंद्र। इन केंद्रों को कई मुख्य सुविधाओं द्वारा मजबूत किया गया है जैसे कि बायोएसे लेबोरेटरी, बायोरेपोजिटरी, बायोसेफ्टी लेवल-3 लैब, डेटा मैनेजमेंट सेंटर, इम्युनोलॉजी कोर लेबोरेटरी, मल्टी-ओमिक्स सुविधा, प्रयोगात्मक पशु सुविधा, वैक्सीन डिजाइन और विकास सुविधा, बायोडिजाइन में नवाचार का स्कूल आदि। जो THSTI के अनुसंधान कार्यक्रमों और राष्ट्रीय राजधानी क्षेत्र बायोटेक साइंस क्लस्टर और अन्य शैक्षणिक और औद्योगिक भागीदारों के लिए विशाल संसाधनों के रूप में काम करते हैं। ब्रिक-टीएचएसटीआई कई महत्वाकांक्षी और वैश्विक रूप से प्रतिस्पर्धी शैक्षणिक पाठ्यक्रमों के माध्यम से वैज्ञानिक लीडर की अगली पीढ़ी को प्रशिक्षित करता है जो बहु-विषयक शिक्षाविदों-उद्योग साझेदारी के माध्यम से अनुसंधान और नवाचार को बढ़ावा देता है।

BRIC-THSTI has built several inter-institutional collaborations and connectivity with industry supported by well-trained teams of research and laboratory staff. THSTI has established various centres namely (a) Centre for Maternal and Child Health, (b) Centre for Virus Research, Therapeutics and Vaccines (c) Centre for Tuberculosis Research (d) Centre for Microbial Research, (e) Centre for Immunobiology and Immunotherapy (f) Centre for Drug Discovery (g)

Clinical Development Services Agency (h) Computational and Mathematical Biology Centre (i) Centre for Bio-design and Diagnostics. These centres are strengthened by many core facilities viz. Bioassay Laboratory, Biorepository, Biosafety Level-3 Lab, Data Management Centre, Immunology Core laboratory, Multi-Omics facility, Experimental Animal Facility, Vaccine design and Development facility, School of Innovation in Bio design etc. that serve as huge resources for the research programmes of THSTI and also the National Capital Region Biotech Science Cluster and other academic and industrial partners. BRIC-THSTI trains the next generation of scientific leaders through many ambitious and globally competitive academic courses which promotes research and innovation through multi-disciplinary academia-industry partnerships.

3. यह भर्ती निम्नलिखित परियोजनाओं के तहत ब्रिक-टीएचएसटीआई की रिक्तियों को भरने के लिए है:

This recruitment is to fill up the vacancies of BRIC-THSTI under the following projects:

पद के लिए आवश्यक शैक्षिक योग्यता और अनुभव / Educational Qualification and Experience required for the post:

क्रम संख्या/ S.No.	पद का नाम/Name of the Post/ पदों की संख्या/ No. of posts/ मासिक समेकित परिलब्धियां/ Monthly consolidated emoluments/ आयु सीमा/ Age Limit	आवश्यक और वांछनीय योग्यता और अनुभव/ Essential & Desirable qualifications & Experience	नौकरी का विवरण / कौशल आवश्यक Job description/ Skills required
प्रोजेक्ट/Project : Placenta Health Project - Adaptive Dose Finding Platform in India पीआई/ PI : Dr. Nitya Wadhwa			
1.	क्लीनिकल परियोजना प्रबंधन / Clinical Project Manager एक पद/ One post Rs. 1,50,000/- 45 वर्ष/ 45 years	MBBS with MD/ DNB, preferably in Obstetrics & Gynaecology/ Pediatrics/ Community Medicine/ or a related discipline from a recognized university. OR MBBS with a PG Diploma in Obstetrics & Gynaecology/ Child Health/ Pediatrics/ or a related discipline from a recognized university, with at least one (1) year of relevant experience in clinical research and/or maternal and child health. OR MBBS/ MDS from a recognized university, with at least four years (4) of relevant experience in clinical research and/or maternal and child health. OR MBBS with a PG degree in Public Health/ Clinical Research/ or a	The Clinical project Manager (CPM) will be leading the study team and will be primary point of contact for operational aspects of implementation of the clinical trial activities from study start-up through database lock, ensuring compliance with GCP and applicable guidance. He/ she will be the primary link between study coordination unit and study investigators. The CPM will have an oversight responsibility for activities undertaken at hospital sites. He/she will be responsible for: • Providing input into and/or developing study related material such as clinical operations plan, SOPs, CRF completion guidelines, informed consent, study logs/forms and other study related documents;

		related discipline from a recognized university, with at least two (2) years of relevant experience in clinical research and/or maternal and child health. Desirable: • Work experience in a clinical trial • Understanding of GCP, regulations and guidelines • Demonstrated ability to develop and implement monitoring plans, SOPs • Computer skills including proficiency in use of Microsoft Office applications • Familiarity with statistical/analytical software (R, Stata) and data management platforms (Red Cap) • Ability to build effective project teams, ability to motivate others, delegation, drive and timely/quality decision making • Knowledge of adverse medical event investigation, analysis, and reporting procedures and standards • Effective communication skills, the provision of timely and accurate information to stakeholders • Good organisational behaviour and problem-solving skills • Effective time management skills and ability to manage competing priorities.	• Supporting the submissions for relevant government / ethics approvals; • Developing training module and planning the initial and retraining sessions for the research study staff along with the site CROs (called clinical research officers) • Contribute through operational inputs in protocol and study budget related decisions; • Structuring and supervising compliance for the study management plans; Ensuring compliance with the project requirements and cascading the issues/ updates to the relevant stakeholders; • Supervising the site preparation, study implementation at site and ongoing study and QC activities; • Overseeing the PK-PD substudy conducted at sites • Reviewing protocol deviations and loss to follow up to ensure quality data is delivered; • Communicating with site supervisor and site investigator for tracking patient recruitment and progress to study timelines; maintaining and reporting metrics for clinical site performance • Reviewing and exploring study data on an ongoing basis to identify patterns, trends, inconsistencies, outliers, and potential data-quality or operational issues • Having the ability to undertake basic exploratory data analysis, generate and review descriptive summaries and study indicators, and appropriately interpret study data in collaboration with the investigators, data management team, and statisticians • Providing input and support to maintain appropriate documentation for adverse event safety monitoring, and
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			collaborating in submission of safety reports to sponsor, Ethics Committees and other applicable authorities; • Liaisoning with the QM team to ensure good quality of study data; • Providing support to site team to prepare for clinical audits and to respond to audit findings conducted by internal QA and external agencies; • Supervising the data management progress with data manager and the DM team; • Work with coordinating PIs to ensure that the trial is meeting its targets, is producing meaningful output and to predict and plan any changes that warrant requests to changes in protocol, funding, or timelines; • Keeping stakeholders informed on study progress, risks and accomplishments. • Knowledge of adverse medical event investigation, analysis, and reporting procedures and standards <i>The CPM will be based at the coordinating centre at THSTI, Faridabad with regular site visits to AIIMS, Delhi or GTB Hospital, Delhi or LHMC, Delhi (the 3 participating hospital sites).</i>
2.	क्लीनिकल अनुसंधान अधिकारी / Clinical Research Officer एक पद/ One post Rs. 1,25,000/- 45 वर्ष/ 45 years	MBBS from a recognised university with at least three (3) years of relevant experience in clinical research and/or maternal and child health. OR MBBS with a PG degree in Public Health/ Clinical Research/ or a related discipline from a recognized university, with at least one (1) year of relevant experience in clinical research and/or maternal and child health. OR MDS from a recognised university with at least three (3) years of	The selected candidate will be responsible for the day-to-day coordination and supervision of DREAM Trial activities at participating hospital site(s), ensuring compliance with the study protocol, SOPs, GCP, and applicable regulatory and ethical requirements. The role will involve coordination with investigators, the Study Coordination Unit, site teams, Data Management and Quality Management teams. She/He will be responsible for:

		relevant experience in clinical research and/or maternal and child health. OR BDS/ BAMS/ BHMS/ BPT or equivalent degree plus a PG degree in Public Health/ Clinical Research/ or a related discipline from a recognised university, with at least three (3) years of relevant experience in clinical research and/or maternal and child health. OR BDS/ BAMS/ BHMS/ BPT or equivalent degree from a recognised university, with at least five (5) years of relevant experience, after completion of internship, in clinical research and/or maternal and child health. Desirable: • Work experience in a clinical trial • Conversant with Good Clinical Practice • Demonstrated ability to develop and implement monitoring plans, SOPs • Computer skills including proficiency in use of Microsoft Office applications • Ability to build effective project teams, ability to motivate others, delegation, drive and timely/ quality decision making • Good organisational behaviour and problem-solving skills • Effective time management skills and ability to manage competing priorities.	• Coordinating and supervising screening, enrolment, study visits and participant follow-up as per protocol. • Supervising study procedures, including clinical assessments, sample collection, investigational product-related activities and adherence assessments. • Overseeing implementation of the PK-PD substudy at the site. • Supporting timely AE/SAE documentation, follow-up and safety reporting. • Reviewing CRFs/eCRFs and study data for completeness, consistency and timely resolution of queries. • Reviewing protocol deviations, missing/inconsistent data, loss to follow-up and site performance indicators and supporting corrective actions. • Coordinating biological sample collection, processing, storage and transportation, as applicable. • Ensuring appropriate investigational product accountability, dispensing, return and adherence documentation. • Liaising with Data Management and Quality Management teams to ensure quality and timely flow of study data. • Training and supervising research staff in study procedures, data collection, follow-up and CRF/eCRF completion. • Supporting monitoring visits, audits and inspections and resolution of observations. • Tracking recruitment, retention and study progress and providing regular updates to the CRC and investigators. • Ensuring availability and appropriate maintenance of study equipment, supplies, documents and logs.
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			Performing any other study-related work assigned by the Principal Investigator/Coordinating Investigator. <i>The clinical research officers will be based either at AIIMS, Delhi or GTB Hospital, Delhi or LHMC, Delhi (the 3 participating hospital sites)</i>
उपरोक्त पदों के लिए/ For posts mentioned above- ➤ ऑनलाइन आवेदन प्राप्त करने की अंतिम तिथि: 04 नवंबर 2026 / Last date for receipt of online application: 4th November 2026 ➤ आवेदनों की जांच/शॉर्टलिस्टिंग की जाएगी और आगे के चयन के लिए प्रक्रिया की जाएगी। The applications will be scrutinised/shortlisted and processed for further selection.			

सामान्य नियम व शर्तें/ GENERAL TERMS & CONDITIONS:

- 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.
- All educational, professional and technical qualifications should be from a recognized Board/University.
- 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 the eligibility criteria.
- Closing date of online application will be the **CRUCIAL DATE** for determining eligibility with regard to age, essential qualification, experience etc.
- The age limit, qualification, experience and other requirements may be relaxed at the discretion of the competent authority, in case of candidates who are otherwise suitable.
- 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.
- All results/notifications will only be published on our website. Therefore, the candidates should essentially visit THSTI website, regularly.
- All communications will only be made through email.
- 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.
- The no. of vacancy indicated above may change subjected to the actual requirement at the time of Written test/skill test/interview.
- With regard to any provisions not covered in this notification, the bye laws of THSTI / Govt. of India rules/ guidelines shall prevail.
- Canvassing wrong information in any form will be a disqualification.

उपरोक्त तालिका में उल्लिखित पदों के लिए आवेदन कैसे करें/ HOW TO APPLY FOR POSTS MENTIONED IN ABOVE TABLE:

1. **Documents to be kept handy before filling up the online application:** (all the documents except (i) should be in pdf format):

- i) A soft copy of your passport size photo and signature. (jpeg/jpg/png format)
- ii) A comprehensive CV containing details of qualification, positions held, professional experience / distinctions etc.
- iii) Matriculation certificate (equivalent to 10th Standard) / Mark sheet
- iv) Intermediate certificate (equivalent to 12th Standard) / Mark sheet
- v) Graduation/Diploma degree certificate / Mark sheet
- vi) Post-Graduation degree certificate & Mark sheet (if applicable)
- vii) PhD degree/certificate (if applicable)
- viii) Relevant experience certificates (if applicable)
- ix) Caste / Disability certificate in the format prescribed by the Govt. of India, if applicable

2. **Procedure for filling up online application:**

- i) The eligible and interested candidates may apply online at the Institute's website. Applications through any other mode will not be accepted.
- ii) The following will be the step wise procedure-
 - A) Step 1 : Details of applicant
 - B) Step 2 : Uploading of documents
 - C) Step 3 : Payment of application fee
 - The payment can be made by using Debit Card / Credit Card / Internet Banking/ UPI.
 - Once payment is made, no correction / modification is possible
 - Candidates are requested to keep a copy of the provisional receipt for future reference.
 - Fee once paid shall not be refunded under any circumstances.
 - Details of fees to be paid are as shown below:

S. No	सीधी भर्ती पर आवेदन करना/ Applying on direct recruitment	आवेदन शुल्क राशि/ Application fee amount
1.	Unreserved, OBC & EWS candidates	Rs. 590/-
2.	SC/ST/Women/PwBD	Rs 118/-

D) Step 4 : Submission of application form

- iii) On successful submission of application, an auto-generated email containing the reference number will be sent to the email address provided. Please keep a note of the reference number for future correspondence.
- iv) Candidates are required to keep a printout of the online application form by using the print button on the dashboard for future reference.
- v) Candidates must ensure that he / she fulfils all the eligibility criteria as stipulated in the advertisement. If it is found that he / she does not fulfil the stipulated criteria during the recruitment process, the candidature of the candidate will be cancelled. If the same is noticed after the appointment, the candidate will be terminated following due process.
- vi) Incomplete applications shall be summarily rejected and no correspondence in this regard shall be entertained.
- vii) In case of difficulty in filling up the online form, please send e-mail to **personnel@thsti.res.in** along with the screenshot of the error displayed (if any).

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

-s/d-
Head-Administration

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