



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-सी/आरएन/20/2026

दिनांक: 29 अक्टूबर 2026

RECRUITMENT NOTICE NO.: THS-C/RN/20/2026

Dated: 29 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 ट्रांसलेट 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 क्लिनिकल डेवलपमेंट सर्विसेज एजेंसी (CDSA) केंद्र में परियोजना पदों की रिक्तियों को भरने के लिए की जा रही है। CDSA, THSTI का एक विशेष केंद्र है, जिसे सार्वजनिक स्वास्थ्य रोगों के लिए किफायती स्वास्थ्य उत्पादों के विकास को सुविधाजनक बनाने के उद्देश्य से स्थापित किया गया है। यह देश का एकमात्र सार्वजनिक केंद्र है जिसे लाभ-न कमाने वाले तकनीक-आधारित प्रीक्लिनिकल और क्लिनिकल उत्पाद विकास के साथ-साथ सार्वजनिक एजेंसियों द्वारा किए जाने वाले क्लिनिकल अनुसंधान को समर्थन और पोषण देने के उद्देश्य से बनाया गया है। यह प्रशिक्षण और सीखने के एक इको-सिस्टम के विकास की दिशा में काम करता है और सार्वजनिक क्षेत्र की संस्थाओं तथा छोटे और मध्यम उद्यमों (SME) के साथ मिलकर नवाचारपूर्ण तकनीकों को जनहित में चिकित्सीय उत्पादों में बदलने का कार्य करता है। This recruitment is to fill up the vacancies for project positions at Clinical Development Services Agency (CDSA) center. CDSA is a niche center of THSTI established to facilitate development of affordable healthcare products for public health diseases. It is the only public Centre in the country created with a mandate to support and nurture cost-effective, high quality, not-for-profit technology-based preclinical and clinical product development as well as support clinical research conducted by public agencies. It works towards development of an eco-system for training and learning and work with public sector institutions, and small and medium enterprises (SME) to translate innovative technologies into medical products for public good.

CDSA के मुख्य उद्देश्य निम्नलिखित हैं:

- एक अकादमिक क्लिनिकल रिसर्च यूनिट के रूप में, अध्ययन योजना, सेटअप, संचालन, परियोजना प्रबंधन, निगरानी, डेटा प्रबंधन, सुरक्षा रिपोर्टिंग, विश्लेषण और रिपोर्ट लेखन में अन्वेषकों और SMEs को अंत-to-अंत क्लिनिकल अध्ययन समर्थन प्रदान करना।
- क्लिनिकल विकास/प्रयोजन और नियमन के क्षेत्र में उच्च गुणवत्ता वाले प्रशिक्षण के माध्यम से शोध क्षमता और क्षमता का निर्माण करना।
- देश में क्लिनिकल रिसर्च पर्यावरण का समर्थन और सुदृढ़ करना।
- नियामक विज्ञान और नीति समर्थन: शोधकर्ताओं, नियामकों, स्वास्थ्य नीति निर्माताओं और उद्योग को समर्थन देने के लिए उपकरण और दृष्टिकोण प्रदान करना।

The main objectives of CDSA are:

- As an academic Clinical Research Unit, to undertake & provide end -to- end clinical study support for investigators and SMEs in study planning, set up, conduct: project management, monitoring, data management, safety reporting, analysis and report writing
 - Build research capacity and capability through high quality training in the area of clinical development/trials and regulation
 - Support and strengthen clinical research environment in the country
 - Regulatory science and policy support: provide tools and approaches to support researchers, regulators, health policy makers & industry.
4. 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:

1.	पद का नाम/Name of the post	डाटा प्रबंधक/ Quality Manager
	पदों की संख्या/Number of the post	01
	परियोजना का नाम/Name of the Project	Sepsis related mortality in neonates in India: A multi-disciplinary, multi-institutional research program for context specific solutions
	वेतन/Emoluments	Rs. 56,000/- + HRA
	उम्र/Age	Up to 35 years
	न्यूनतम शैक्षिक योग्यता और अनुभव/Minimum Educational Qualification and Experience	<p>Essential qualification and work experience:</p> <ul style="list-style-type: none"> Master's degree in any field preferably in science, with 6 years of experience in clinical data management/clinical research/operations/MIS/data analysis/IT/computer science/healthcare field <p style="text-align: center;">OR</p> <ul style="list-style-type: none"> Graduation degree in any field preferably in science, with 8 years of experience in clinical data management/clinical research/operations/MIS/data analysis/IT/computer science/healthcare field <p>Desirable qualification and work experience:</p> <ul style="list-style-type: none"> Diploma in Clinical research and clinical data management. Familiarity with industry standard CDMS and some programming skills: Preparation of Clinical Study Data Management documents.
	नौकरी का प्रोफाइल/Job profile	<ul style="list-style-type: none"> Clinical Study Protocol understanding and experience in the preparation of Data Management documents - DMP (Data Management Plan), DVP (Data Validation Plan/ Edit Checks Document), Annotated CRF, Data Entry Guideline etc. Quality Check of Database Design, Validation Program, Annotated CRF, Data Extract Views, Laboratory Details, Site and Investigators and Final Data Listings Good experience and working knowledge of CRF designing Preparation of Data transfer guidelines for external data load and self-evident correction chart. Working knowledge of Query management, data cleaning, data freezing and data archival. Change Requests and takes approval from sponsor Global library review and approval for changes Interact with other project team members to support the set-up, maintenance, and closure of the Data Management aspects of the project Sound knowledge of Clinical Database Development tools, logics and techniques and GCDMP Working knowledge of database standards and study development process, CDM SOPs, CDISC & SDTM standards E/SAE reconciliation Knowledge of medical dictionaries integration and medical coding in the databases. Preparing interim reports and review of listings of data for clinical trial status and data extraction in collaboration with the statistician

		<ul style="list-style-type: none"> • Maintaining and archiving of clinical study related documents • Participates in cross functional team meetings & external client meetings as DM representative • Generate ad-hoc reports as needed • The data manager will ensure that security of all data is maintained and confidentiality of participants is protected. • Managing requests for data from external third parties – including liaising with internal staff and external collaborators to provide data in a timely and appropriate manner and maintenance of a database detailing the status of such external data requests. • Knowledge of Laboratory Information Management Systems (LIMS) to track requisition and specimen status to ensure prompt turnaround time of sample results. • Maintain stringent standards for quality, identifying any issues which might adversely impact the quality of test results and immediately communicating these to the appropriate management representatives to ensure prompt resolution • Create charts, graphs, dashboards, and other visual representations to present data • Coordinate the archiving of study databases and related documents • Effective interaction with intra-departments to ensure all required, vital information and documentation is acquired in a timely manner. • Lead in preparation of datasets for analysis including data cleaning and ensuring compliance with the data protection. • Develop Standard Operation Procedures and train study staff • Supervise DM activities at the clinical site.
	कौशल /Skills	<ul style="list-style-type: none"> • Good management & leadership skills • IT literate (experience with Microsoft based applications, data visualization and other CDMS applications) • Must have experience in handling databases and query management • Client focused approach to work • Meticulous attention to detail. • Must have understanding of clinical trials and familiarity with clinical data management functions. • Good interpersonal, verbal and written communication skills. • Client focused approach to work. • A flexible attitude with respect to work assignments and new learning. • Meticulous attention to detail. • Effective time management in order to meet metrics or team objectives. • Commitment to project and team goals. • Must be able to work independently but seek guidance when necessary. • Team player with outstanding inter-personal, negotiation skills and organizational skills. • Sense of urgency in completing assigned tasks. • Exhibits a sense of urgency about solving problems and completing work. • Shows commitment to and performs consistently high quality work. • Ability to model behaviors and ethics in line with CDSA Mission and Vision
वॉक-इन साक्षात्कार की तिथि/ Date of walk-in interview:		17 November 2026 @09:00 AM at THSTI, NCR Biotech Science Cluster, 3rd Milestone, Faridabad-Gurugram Expressway, Faridabad – 121001

2.	पद का नाम/Name of the post	परियोजना अनुसंधान वैज्ञानिक – I (गैर-चिकित्सीय) /Project Research Scientist - I (Non-Medical)
	पदों की संख्या/Number of the post	01
	परियोजना का नाम/Name of the Project	Sepsis related mortality in neonates in India: A multi-disciplinary, multi-institutional research program for context specific solutions
	वेतन/Emoluments	Rs. 56,000/- + HRA
	उम्र/Age	Up to 35 years
	न्यूनतम शैक्षिक योग्यता और अनुभव/Minimum Educational Qualification and Experience	Essential Qualification: <ul style="list-style-type: none"> Post Graduate Degree, including the integrated PG degrees Desirable Qualification and Experience: <ul style="list-style-type: none"> Postgraduate degree in Clinical Research, Clinical Trials, Epidemiology, Public Health, or a related discipline from a recognised university Additional qualification or prior training in Medical Laboratory Technology or a related laboratory discipline. Experience in clinical research, preferably involving biological sample collection, processing, storage, transport, and associated documentation. Experience in preparing standard operating procedures (SOPs), study workplans, and coordinating study activities. Good Clinical Practice (GCP) certification.
	नौकरी का प्रोफाइल/Job profile	<ul style="list-style-type: none"> Support the coordination and management of clinical studies, including ethics and institutional approvals. Monitor study related laboratory processes including sample collection, processing, labeling, storage, transport, tracking, testing workflow, and documentation, to ensure compliance with protocol requirements, SOPs, and applicable quality standards, including GCLP principles, where applicable. Develop, review and maintain trial documents, including protocols, SOPs, CRFs, trial manuals, and operational guidelines. Support trial close-out activities, including database lock, site close-out, reconciliation and archiving of essential trial documents, and completion of trial close-out requirements. Coordinate the transfer, storage, or destruction of trial materials and biological samples in accordance with the trial protocol and applicable guidelines. Contribute to scientific writing, including manuscripts, trial protocols, grant reports, and conference presentations. Ensure adherence to the protocol, ethical requirements, and applicable guidelines, and support identification and escalation of protocol deviations and site-level issues Conduct site monitoring visits from site initiation through closeout, ensuring trials are conducted in compliance with the study protocol, GCP guidelines, SOPs, and applicable regulatory requirements. Coordinate with investigators, collaborators, and external stakeholders to ensure effective study execution Provide training and operational guidance to study teams on protocol, processes, and documentation requirements, as applicable

		<ul style="list-style-type: none"> Perform any other project related duties assigned by the Principal Investigator.
	कौशल /Skills	<ul style="list-style-type: none"> Proficient in computer literacy in Excel, Word, PowerPoint, and Access. Excellent documentation, communication, and organizational skills. Detail-oriented with effective time management skills and ability to manage multiple tasks and priorities efficiently. Ability to work independently with minimal guidance as well as collaboratively within a team setting
वॉक-इन साक्षात्कार की तिथि/ Date of walk-in interview:		12 November 2026 @09:00 AM at THSTI, NCR Biotech Science Cluster, 3rd Milestone, Faridabad-Gurugram Expressway, Faridabad – 121001

नोट:1) उपरोक्त उल्लिखित पद हेतु आवेदन करने वाले उम्मीदवार अपने नवीनतम रिज्यूमे, शैक्षिक योग्यता और अनुभव के समर्थन में दस्तावेजों की एक प्रति, मूल दस्तावेज और सत्यापन के लिए एक वैध आईडी कार्ड लाना होगा। **2)** जो उम्मीदवार निर्धारित समय के बाद आएंगे, उन्हें प्रवेश नहीं दिया जाएगा। **3)** लिखित परीक्षा/कौशल परीक्षण/साक्षात्कार के लिए आने वाले सभी उम्मीदवारों को अनिवार्य रूप से अपनी मोबाइल फोन और वैध पहचान प्रमाण रिसेप्शन पर जमा करना होगा, और यह केवल चयन प्रक्रिया पूरी होने के बाद ही वापस किया जाएगा।

NOTE: 1) The candidates applying for the above mentioned post 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 cards for verification. 2) Candidates coming after the time slot mentioned will not be entertained. 3) All the candidates coming for written test/skill test/interview will be mandatorily required to deposit their mobile phone along with a valid Identity proof at the reception and the same will only be returned back on completion of the entire selection process.

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 qualification 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.
- 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/Department 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.

- i) The no. of vacancy indicated above may change subjected to the actual requirement at the time of Written test/skill test/interview.
- j) With regard to any provisions not covered in this notification, the bye laws of THSTI / Govt. of India rules/ guidelines shall prevail.
- k) Canvassing wrong information in any form will be a disqualification.

"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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