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/21/2026 **दिनांक: 26 नवंबर 2026**

RECRUITMENT NOTICE NO.: THS-C/RN/21/2026 **Dated: 26 November 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 ट्रान्स्लेट 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, (ड़) 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:

a. 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।
b. क्लिनिकल विकास/प्रयोजन और नियमन के क्षेत्र में उच्च गुणवत्ता वाले प्रशिक्षण के माध्यम से शोध क्षमता और क्षमता का निर्माण करना।
c. देश में क्लिनिकल रिसर्च पर्यावरण का समर्थन और सुदृढ़ करना।
d. 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:

a. 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
b. Build research capacity and capability through high quality training in the area of clinical development/trials and regulation
c. Support and strengthen clinical research environment in the country
d. 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:

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

1.	**पद का नाम/Name of the post**	क्लिनिकल अनुसंधान सहयोगी/ Clinical Research Associate
	पदों की संख्या/Number of the post	02
	परियोजना का नाम/Name of the Project	Efficacy. Safety, and Tolerability of Ayurveda regimen as an adjunct to anti-tuberculosis treatment (ATT) and macronutrient supplementation on body weight, nutritional outcomes, and quality of life in adults with newly diagnosed drug-sensitive NAAT-positive Pulmonary Tuberculosis: An open-label randomized controlled parallel-arm clinical study
	वेतन/Emoluments	Rs. 67,000/- + HRA
	उम्र/Age	40 years
	न्यूनतम शैक्षिक योग्यता और अनुभव/Minimum Educational Qualification and Experience	• Master's degree in any branch of life sciences or pharmacy or public health or health related discipline with minimum three (3) years of relevant clinical trial monitoring experience. • MBBS/ BDS/ BHMS/ BAMS/ BPT with relevant clinical trial monitoring experience.
	नौकरी का प्रोफ़ाइल/ Job profile	• The Clinical Research Associate (CRA) is responsible for overseeing clinical trial sites from initiation to closeout, ensuring compliance with study protocols, ICH-GCP, applicable regulations, and internal SOPs. Responsibilities include: • Conduct site monitoring visits from initiation through closeout, ensuring trials are conducted in compliance with the study protocol, GCP guidelines, SOPs, and applicable regulatory requirements. • Set up trial sites, ensuring that investigational products and essential trial supplies are delivered, stored, and documented appropriately. • Perform quality checks and execute quality assurance process across clinical operations and clinical laboratories in accordance with GCP/GCLP standards. • Provide training on protocols and trial procedures to site staff and maintain ongoing communication to support study execution and address issues. • Support clinical staff through guidance and training as and when needed. • Create, maintain, and submit all required documentation related to site management, monitoring visits, findings, and follow-up actions. • Track and manage study progress, including regulatory and ethics submissions, patient recruitment and enrolment, CRF completion, and data query resolution. • Verify data accuracy through source data/document verification to ensure consistency between CRFs and clinical records. • Prepare detailed monitoring visit reports and contribute to the preparation and archiving of essential trial documents. • Assess trial site compliance and escalate quality or protocol deviations to the Project Manager, or Senior Leadership as appropriate. • Collaborate with Clinical Portfolio Management and other internal departments on cross-functional initiatives and project requirements • Collaborate cross-functionally with clinical operations, data management, safety, and regulatory teams.

		• Maintain effective communication with investigators and site staff to ensure study success. • Frequently travel to assigned trial/study sites by eligible modes of travel, including city and state public transportation, own transportation, train travel, or private mass transport services, including standard and luxury buses.
	कौशल /Skills	• Proficient in computer applications, with demonstrated expertise in Microsoft Office Suite (Word, Excel, PowerPoint, Outlook). • Strong knowledge of ICH-GCP, GCLP, and regulatory guidelines. • Excellent documentation, communication, and organizational skills. • Ability to travel frequently to assigned trial sites. • Detail-oriented with effective time management skills and ability to manage multiple tasks and priorities efficiently.
वॉक-इन साक्षात्कार की तिथि/ Date of walk-in interview:		**16th December 2026 @09:00 AM at THSTI, NCR Biotech Science Cluster, 3rd Milestone, Faridabad-Gurugram Expressway, Faridabad – 121001**
4.	**पद का नाम/Name of the post**	परियोजना तकनीकी सहयोग - III /Project Technical Support - III
	पदों की संख्या/Number of the post	01
	परियोजना का नाम/Name of the Project	Improving maternal and neonatal outcomes using imaging data science
	वेतन/Emoluments	Rs. 28,000 + HRA
	उम्र/Age	35 years
	न्यूनतम शैक्षिक योग्यता और अनुभव/Minimum Educational Qualification and Experience	**Essential qualification:** Three Years Graduate degree in relevant subject / field + three Years post qualification experience OR Post Graduate in life science.
	नौकरी का प्रोफ़ाइल/ Job profile	The **Project Technical Support – III** plays a key role in supporting the execution and management of clinical trials and research studies across all phases. Reporting to the Chief–CPM and Project Manager, the role ensures efficient day-to-day research operations, accurate maintenance of regulatory and study documentation, and effective coordination of communication and logistics among study stakeholders. The incumbent is expected to work independently on assigned tasks, anticipate and address potential challenges, and contribute to process improvements. • Review and develop familiarity with study protocols, including procedures, timelines, eligibility criteria, confidentiality, and privacy protections. • Perform data and process monitoring support; ensure maintenance of essential documents at trial sites and CDSA. • Support on-site co-monitoring visits with the Project Manager (PM) and Quality Manager (QM); willingness to travel across trial sites as required.

		• Assist the study team in communicating study requirements to investigators and site staff; provide appropriate training materials and maintain study-specific training logs. • Support the development of clinical study documents for start-up and other project requirements; assist in preparing submissions for regulatory and ethics committee (EC) approvals. • Create, maintain, and periodically review the Trial Master File (TMF), Investigator Site Files (ISF), and related systems/tools for accuracy, completeness, and GCP compliance. • Participate in project-related meetings; assist with agenda preparation, presentations, minutes, and tracking of action items. • Assist with preparation, handling, distribution, filing, version control, and archiving of study documentation as per SOPs and project requirements. • Maintain and update regulatory/ethics submission trackers; follow up on approvals and site compliance documentation. • Support safety reporting by tracking and forwarding SAEs/AEs to the relevant study teams. • Coordinate study logistics, including shipment and tracking of clinical supplies, kits, equipment, and courier activities. • Assist with eCRF/EDC management by monitoring data entry, resolving queries, and coordinating with data management. • Maintain delegation of authority logs and ensure investigator/site staff training certifications are up to date. • Support audit and inspection readiness; assist with CAPA documentation and tracking. • Provide general administrative and project support, including maintaining trackers, study contact lists, meeting schedules, and inter-departmental coordination
	कौशल /Skills	• Strong written and verbal communication skills in English. • Proficiency in MS Office (Word, Excel, PowerPoint, Outlook) and working knowledge of project management tools. • Ability to work independently with minimal supervision as well as collaboratively in a team environment. • Flexible, organized, and able to work effectively under pressure while maintaining integrity.
वॉक-इन साक्षात्कार की तिथि/ Date of walk-in interview:		**16th December 2026 @09:00 AM at THSTI, NCR Biotech Science Cluster, 3rd Milestone, Faridabad-Gurugram Expressway, Faridabad – 121001**
3.	**पद का नाम/Name of the post**	डाटा एंट्री ऑपरेटर /Data Entry Operator
	पदों की संख्या/Number of the post	01
	परियोजना का नाम/Name of the Project	DREAMEE
	वेतन/Emoluments	Rs. 26,000/-
	उम्र/Age	35 Years
	न्यूनतम शैक्षिक योग्यता और अनुभव/Minimum	• Graduate in any discipline with 1 year experience in a reputed organization. OR

	Educational Qualification and Experience	• Intermediate/ 12th from recognized board with 4 years' experience in a reputed organization. • Basic computer literacy is essential.
	नौकरी का प्रोफ़ाइल/ Job profile	The Data Entry Operator will be responsible for: • The Data Entry Operator will be responsible for error free data entry and management of data within the defined timelines • He / she will be responsible for providing support in maintenance and filing of all data management documents at site • Receiving Case Report Forms (CRFs) from clinical team • Maintain accountability of received CRF pages • Provide support to update the online trackers • Keeping track of reports and provide support in extraction of data • Support to data managers for the maintenance of data management trial master files
	कौशल /Skills	• Willingness to learn clinical/research data management systems and processes. • Excellent reading comprehension and strong written as well as verbal communication skills including good command of English required • Good understanding of needs for project and job responsibilities. • Computer skills including proficiency in use of Microsoft Office applications • Good organizational behavior and problem-solving skills • Well versed in recording the data • Ability to establish and maintain effective working relationships with co-workers, managers, investigators • Good spelling, grammar, and punctuation skills
वॉक-इन साक्षात्कार की तिथि/ Date of walk-in interview:		**09th December 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 रिज़्यूमे, शैक्षिक योग्यता और अनुभव के समर्थन में दस्तावेजों की एक प्रति, मूल दस्तावेज़ 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:

a) 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.

b) All educational, professional and technical qualification should be from a recognized Board/University.

c) 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.

d) Closing date of online application will be the **CRUCIAL DATE** for determining eligibility with regard to age, essential qualification, experience etc.

e) 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.

f) All results/notifications will only be published on our website. Therefore, the candidates should essentially visit THSTI website, regularly.

g) All communications will only be made through email.

h) 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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