



ब्रिक-ट्रान्सलेशनल 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, 3<sup>rd</sup> Milestone, Faridabad – Gurugram Expressway,  
P.O. Box No. 04, Faridabad – 121001

**भर्ती नोटिस सं. : टीएचएस/आरएन/04/2026**

**दिनांक: 17 अप्रैल 2026**

**RECRUITMENT NOTICE NO.: THS/RN/04/2026**

**Dated: 17<sup>th</sup> April 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 ट्रान्सलेशनल अनुसंधान करने और मानव स्वास्थ्य में सुधार के लिए अवधारणाओं को उत्पादों में ट्रान्सलेट करने के लिए विषयों और व्यवसायों में अनुसंधान सहयोग विकसित करता 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 conducts 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, चिकित्सा अनुसंधान केंद्र, ट्रान्सलेशनल अनुसंधान सुविधा 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, Medical Research Centre, Translational Research Facility 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
<b>प्रोजेक्ट/Project: First-in-India Sham-Controlled Human Challenge Model Using Influenza Vaccines: A Catalyst for Early Clinical Validation of Novel Vaccines and Immunotherapies</b>			
1.	परियोजना वैज्ञानिक (फार्मासिस्ट) / <b>Project Scientist (Pharmacist)</b>  एक पद/ One post  Rs. 75,000/-  35 वर्ष/ 35 years  वॉक-इन इंटरव्यू की तिथि/ <b>Date of walk-in interview: 5<sup>th</sup> May 2026</b>	M.Pharma from a recognized university with atleast one (1) years' of post-qualification experience in clinical trial pharmacy. <b>OR</b> B.Pharma from a recognized university with atleast three (3) years' of post-qualification experience in clinical trial pharmacy.  The candidate must have a valid registration certificate from the State Pharmacy Council.  The candidate must have a knowledge of drug storage and dispensing regulations.	<ul style="list-style-type: none"> <li>Understanding of establishing a sterile compounding pharmacy setting and GCP style documentation and traceability to support regulated Phase I human clinical trials.</li> <li>Manage the complex preparation of the Investigational Product (IP), often involving "Extemporaneous Preparation"</li> <li>Maintain the "Blinding" integrity of the study</li> <li>Work directly with the Dose Escalation Committee (DEC). Once a safety cohort is cleared, the pharmacist is responsible for calculating and preparing the next (higher) dose level for the subsequent cohort</li> <li>Ensure that high-precision analytical balances and laminar flow hoods are qualified (IQ/OQ/PQ) and calibrated</li> </ul>

			<ul style="list-style-type: none"> <li>Investigational product management (receipt, storage, dispensing, accountability)</li> <li>Temperature monitoring and maintenance of pharmacy logs</li> <li>Drug accountability and reconciliation</li> <li>Preparation and labeling as per protocol requirements</li> <li>Compliance with pharmacy regulations and GCP guidelines</li> <li>Coordination with sponsor for drug supplies</li> <li>Support in sample processing and laboratory activities.</li> </ul>
<p><b>प्रोजेक्ट/Project: Development of monovalent and multivalent poultry vaccine using THSTI Multivalent self-assembled Nanocage Vaccine platform (MSN platform)</b></p> <p><b>पीआई/ PI : Dr. Sweety Samal</b></p>			
2.	<p><b>परियोजना सहायक / Project Assistant</b></p> <p>एक पद/ One post</p> <p>Rs. 35,000/-</p> <p>35 वर्ष/ 35 years</p> <p><b>वॉक-इन इंटरव्यू की तिथि/ Date of walk-in interview: 8<sup>th</sup> May 2026</b></p>	<p>Masters in any branch of Life Sciences from a recognized university with two (2) years' of post-qualification experience in virology, molecular biology and immunology.</p>	<ul style="list-style-type: none"> <li>Knowledge of virology, biochemistry, immunology and molecular biology related to cell culture, virus characterization, RT-PCR.</li> <li>Preference will be given to candidate having experience of handling human or animal viruses.</li> <li>Ability to perform physical tasks within a laboratory workspace and working experience in ABSL2 / ABSL3.</li> </ul>
<p><b>प्रोजेक्ट/Project: A Study of Etiopathogenesis of Systemic Inflammation in Acute Pancreatitis, its Functional Consequences and Progression to Chronic Pancreatitis</b></p> <p><b>पीआई/ PI : Dr. Nitya Wadhwa</b></p>			
3.	<p><b>डाटा प्रबंधक / Data Manager</b></p> <p>एक पद/ One post</p> <p>Rs. 56,000/- plus HRA</p> <p>35 वर्ष/ 35 years</p> <p><b>वॉक-इन इंटरव्यू की तिथि/ Date of walk-in</b></p>	<p>Masters' degree in any field preferably in science, with six (6) years' of post-qualification experience in clinical data management/ clinical research/ operations/ MIS/ data analysis/IT/ computer science/ healthcare field.</p> <p><b>OR</b></p> <p>Graduation degree in any field preferably in science, with eight (8) years' of post-qualification experience in clinical data</p>	<ul style="list-style-type: none"> <li>Overall responsible for all data management activities of assigned project.</li> <li>Support the setting-up, maintenance, and closure of the data management aspects of the project.</li> <li>Perform database testing and database audit as part of Quality Control.</li> <li>Prepare or perform review of Data Management documents - DMP (Data Management Plan),</li> </ul>

	<p><b>interview: 7<sup>th</sup> May 2026</b></p>	<p>management/ clinical research/ operations/ MIS/ data analysis/IT/ computer science/ healthcare field.</p> <p><b>Desirable:</b></p> <ul style="list-style-type: none"> <li>• Diploma in IT/ computer applications/ clinical data management.</li> <li>• Familiarity with industry standard CDMS and some programming skills.</li> <li>• Demonstrated experience of developing clinical study data management documents.</li> <li>• Exposure to programming languages (Python, R) and data visualization tools (Tableau) will be an added advantage.</li> </ul>	<p>DVP (Data Validation Plan/ Edit Checks Document), Annotated CRF, Data Entry Guidelines etc.</p> <ul style="list-style-type: none"> <li>• Working knowledge of query management, data cleaning, data freezing and data archival.</li> <li>• Ensure accuracy, consistency, and completeness of the data for analysis.</li> <li>• Prepare interim/ad-hoc reports and participating in data extraction collaboration with the stakeholders.</li> <li>• Create data visuals (charts, graphs, dashboards, and other visual representations) in Tableau, Power BI, as required to present data.</li> <li>• Coordinate with intra- or inter-departments to ensure all required, vital information and documentation is acquired in a timely manner.</li> <li>• Participate in team meetings &amp; external meetings as DM representative.</li> <li>• Prepare data transfer guidelines for external data load and self-evident correction chart.</li> <li>• Ensure that data integrity and confidentiality of participants are protected</li> <li>• Coordinate the archiving of study databases and related documents.</li> <li>• Prepare interim reports for clinical study status and participating in data extraction in collaboration with the sponsor and coordinating site.</li> <li>• Generate ad-hoc reports as needed</li> <li>• Sound knowledge of logics and data management techniques</li> <li>• Coordinate the archiving of study databases and related documents</li> <li>• Develop standard operation procedures (SOPs) and working</li> </ul>
--	--	--	--

			<p>practice documents including the training of staff.</p> <ul style="list-style-type: none"> <li>• Supervision of Data Management activities at sites.</li> </ul>
<p><b>प्रोजेक्ट/Project : Development of an investigational combo pneumo vaccine (Industry Collaboration)</b></p> <p><b>पीआई/ PI : Dr. Sweety Samal</b></p>			
4.	<p>एग्जीक्यूटिव असिस्टेंट (मानव संसाधन) / <b>Executive Assistant (HR)</b></p> <p>एक पद/ One post</p> <p>Rs. 45,000/-</p> <p>35 वर्ष/ 35 years</p> <p>वॉक-इन लिखित परीक्षा/ इंटरव्यू की तिथि/<b>Date of walk-in written test/ interview: 4<sup>th</sup> May 2026</b></p>	<p>Post-Graduation degree in any discipline from a recognized university.</p>	<ul style="list-style-type: none"> <li>• Will be responsible to handle and manage work in the HR and Administration department.</li> <li>• Strong interpersonal and written communication skills in English &amp; Hindi.</li> <li>• Multitasking under tight deadlines while maintaining discretion and professionalism.</li> <li>• Supporting HR meetings, training programs, and employee data management.</li> </ul>
<p><b>प्रोजेक्ट/Project : Optimized salicylamide derivatives and prodrugs for the treatment of influenza infections</b></p> <p><b>को-पीआई/ Co-PI : Dr. Dinesh Mahajan</b></p>			
5.	<p>परियोजना अनुसंधान वैज्ञानिक-II (संश्लेषणात्मक कार्बनिक रसायनज्ञ) / <b>Project Research Scientist-II (Synthetic Organic Chemist)</b></p> <p>एक पद/ One post</p> <p>Rs. 67,000/- plus HRA</p> <p>40 वर्ष/ 40 years</p> <p>वॉक-इन इंटरव्यू की तिथि/ <b>Date of walk-in interview: 4<sup>th</sup> May 2026</b></p>	<p>Masters' in Organic Chemistry from a recognised university/ Institute with four (4) years' of post-qualification experience in synthetic organic chemistry.</p> <p><b>Desirable:-</b> A candidate with relevant experience in an industrial/ Pharma/ CRO/ biotech setup shall be preferred.</p>	<p>The incumbent is expected to work on synthetic organic chemistry projects in a time-bound manner at the Centre for Drug Discovery (CDD) at THSTI. The candidate with prior experience in synthetic organic chemistry, especially in photochemistry, electrochemistry, and/or organometallic chemistry, is preferred.</p>

**प्रोजेक्ट/Project: Graded Risk-based Assessment for Screening and Prevention of Placental disorders (GRASPP) - A Randomized Controlled Trial Evaluating a Two-Step Contingent Screening Strategy for Great Obstetric Syndrome**

**पीआई/ PI : Dr. Nitya Wadhwa**

<p>6.</p>	<p>परियोजना अनुसंधान वैज्ञानिक-III (चिकित्सा)/ <b>Project Research Scientist-III (Medical)</b></p> <p>एक पद/ One post</p> <p>Rs. 93,000/- plus HRA</p> <p>45 वर्ष/ 45 years</p> <p><b>वॉक-इन इंटरव्यू की तिथि/ Date of walk-in interview: 7<sup>th</sup> May 2026</b></p>	<p>M.D./ MDS/ MPH/ Ph.D. or equivalent degree with basic degree as MBBS/ BDS from a recognized university, including the integrated PG degrees, with three (3) years' of post qualification experience in clinical research.</p> <p><b>Desirable:</b></p> <ul style="list-style-type: none"> <li>• Experience in clinical research or related field.</li> <li>• Good writing skills as well as verbal communication skills.</li> <li>• Computer skills including proficiency in use of Microsoft Office applications.</li> <li>• Good organizational and problem-solving skills.</li> <li>• Effective time management and ability to manage competing priorities.</li> </ul>	<p>The candidate selected for the said post will act as Clinical Research Coordinator (CRC) and 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 startup to completion of study, ensuring compliance with GCP and applicable guidance. CRC will be the primary link between study coordination unit and study investigators.</p> <p>CRC will be responsible for:</p> <ul style="list-style-type: none"> <li>• Supervising all study site activities such as enrollment, consenting, administration of tests, CRF filling and training.</li> <li>• Ensuring that study is conducted in accordance with study protocol, standard operating procedures, Good Clinical Practice, and applicable guidelines.</li> <li>• Providing input 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</li> <li>• Supporting the submissions for relevant government/ ethics approvals</li> <li>• Developing training module and planning the initial and retraining sessions for the research study staff along with the site CROs.</li> <li>• Structuring and supervising compliance for the study management plans</li> <li>• Ensuring compliance with the project requirements and cascading the issues/ updates to the relevant stakeholders</li> </ul>
-----------	---	--	---

			<ul style="list-style-type: none"><li>• Supervising the site preparation, study implementation at site and ongoing study and QC activities to ensure good quality of study data</li><li>• Reviewing protocol deviations and loss to follow up to ensure quality data is delivered</li><li>• Communicating with investigators at THSTI and site investigator for tracking patient recruitment and progress to study timelines; maintaining and reporting metrics for clinical site performance</li><li>• Providing input and support to maintain appropriate documentation for adverse event safety monitoring, and collaborating in submission of safety reports to sponsor, Ethics Committees and other applicable authorities</li><li>• Willing to undergo training, conduct training and monitor study team performing of Neuro developmental assessment of infants</li><li>• Liasoning with all stakeholders and IECs of all sites</li><li>• Providing support to site team to prepare for clinical audits and to respond to audit findings conducted by internal QA and external agencies.</li><li>• Supervising the data management progress with data manager and the DM team</li><li>• Work with coordinating PI 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</li><li>• Keeping stakeholders informed on study progress, risks and accomplishments, share reports, etc.</li></ul>
--	--	--	---

			<ul style="list-style-type: none"> <li>• Knowledge of adverse medical event investigation, analysis, and reporting procedures and standards</li> <li>• Any other work as assigned by PI.</li> <li>• The selected candidate may be posted at any of the following sites: AIIMS Delhi, VMMC &amp; Safdarjung Hospital Delhi, ESIC Medical College Faridabad, or THSTI Faridabad, with occasional visits to the other sites as required.</li> </ul>
7.	<p>परियोजना तकनीकी सहायक-III/ <b>Project Technical Support-III</b></p> <p>एक पद/ One post</p> <p>Rs. 28,000/- plus HRA</p> <p>35 वर्ष/ 35 years</p> <p>वॉक-इन इंटरव्यू की तिथि/ <b>Date of walk-in interview: 30<sup>th</sup> April 2026</b></p>	<p>Graduate degree in Economics / Public Health / Health Economics/ or related social science field from a recognized university with atleast three (3) years' of post-qualification experience in a relevant field.</p> <p><b>OR</b></p> <p>Post-Graduate degree in Economics / Public Health / Health Economics/ or related social science field.</p> <p><b>Desirable:</b></p> <ul style="list-style-type: none"> <li>• Diploma in any health-related Experience in health research.</li> <li>• Well-versed with MS Office tools, particularly Microsoft Excel.</li> </ul> <p>Basic computer skills.</p>	<ul style="list-style-type: none"> <li>• Assist in data collection for health economics studies, including field visits where required.</li> <li>• Support conducting structured interviews with clinicians and healthcare providers for data collection.</li> <li>• Perform data entry, cleaning, and basic analysis using Excel or similar tools.</li> <li>• Assist in literature searches and summarizing relevant evidence.</li> <li>• Assist in developing and formatting data collection tools (questionnaires, interview guides, templates).</li> <li>• Coordinate with team members and stakeholders for scheduling field activities and interviews.</li> </ul> <p>Provide support in day-to-day project posted at THSTI with travel to AIIMS-New Delhi, ESIC-Faridabad, VMMC &amp; SJH Hospital.</p>
8.	<p>परियोजना तकनीकी सहायक-I/ <b>Project Technical Support-I</b></p> <p>एक पद/ One post</p> <p>Rs. 18,000/- plus HRA</p> <p>28 वर्ष/ 28 years</p>	<p>Standard 10<sup>th</sup> with DMLT or equivalent with two (2) years' of experience in relevant subject or field.</p> <p><b>OR</b></p> <p>Three years graduate degree in relevant subject/ field with one (1) year experience in relevant subject or field.</p> <p><b>Desirable:</b></p>	<ul style="list-style-type: none"> <li>• The candidate selected for the said post will act as Field Worker/ Multi-Tasking Staff and he/ she will be responsible for:</li> <li>• Making telephone calls to participants before their visits to remind them of their next visit and confirm their arrival.</li> <li>• Enquire about the participants queries, clarify their doubts regarding the schedule, counsel them and report to seniors for</li> </ul>

	<p>वॉक-इन इंटरव्यू की तिथि/ <b>Date of walk-in interview: 30<sup>th</sup> April 2026</b></p>	<ul style="list-style-type: none"> <li>• Diploma in any health-related discipline from a recognized University.</li> <li>• Graduation from recognized university/ Medical Social Worker (MSW)</li> <li>• Experience in clinical/health related research projects</li> <li>• Effective communication skills to provide timely and accurate information to study participants.</li> <li>• Basic computer skills</li> </ul>	<p>other queries for proper guidance.</p> <ul style="list-style-type: none"> <li>• Transportation of blood samples, sample kit arrangements, receiving and transportation storage process of the clinical samples, and submission to AIIMS Lab from all the hospital sites maintaining appropriate guidelines.</li> <li>• Maintain the call log, visit log and sample transportation log, appointment log with reason for not attending, if any.</li> <li>• Assist Technician and Senior Technician in lab processes.</li> <li>• Have a two wheeler and valid driving license.</li> <li>• Any other as assigned by PI</li> <li>• Selected candidate will have to coordinate between AIIMS Delhi, VMMC &amp; Safdarjung Hospital Delhi, ESIC Medical College Faridabad and will be posted at any of these sites as per work demand.</li> </ul>
<p><b>प्रोजेक्ट/Project: A multi-country, multi-centre, three-arm, parallel group, double-blind, placebo-controlled, randomized trial of two doses of antenatal corticosteroids for women with a high probability of birth in the late preterm period in hospitals in low-resource countries to improve newborn outcomes (Action-III Trial)</b></p> <p><b>पीआई/ PI : Dr. Nitya Wadhwa</b></p>			
<p>9.</p>	<p>वरिष्ठ अनुसंधान वैज्ञानिक (क्लीनिकल)/ <b>Senior Research Scientist (Clinical)</b></p> <p>एक पद/ One post</p> <p>Rs. 1,25,000/-</p> <p>45 वर्ष/ 45 years</p> <p>वॉक-इन इंटरव्यू की तिथि/ <b>Date of walk-in interview: 7<sup>th</sup> May 2026</b></p>	<p>MD/ MS/ DNB or equivalent degree in Obstetrics and Gynecology / Pediatrics / Community Medicine from a MCI recognized university with atleast one (1) year of post-qualification clinical research experience in the domain of maternal and child health</p> <p><b>OR</b></p> <p>MDS or equivalent degree from a recognized university with atleast two (2) years of post-qualification clinical research experience in the domain of maternal and child health.</p> <p><b>OR</b></p>	<p>The selected candidate will be responsible for:</p> <ul style="list-style-type: none"> <li>• Clinical oversight for participant screening, enrolment, intervention administration, and outcome assessments, ensuring compliance with the protocol, SOPs, GCP, and regulatory guidelines.</li> <li>• Supervise safety monitoring and timely reporting of AEs/SAEs to all stakeholders.</li> <li>• Coordinate across investigators, project staff, data management, and monitoring teams to ensure smooth trial conduct and deliverables.</li> </ul>

		<p>MBBS/ BDS or equivalent degree from a recognized university with atleast five (5) years of post-qualification clinical research experience in the domain of maternal and child health.</p> <p><b>Desirable:</b></p> <ul style="list-style-type: none"> <li>• Demonstrated leadership in clinical research or trials, particularly maternal-child health or multi-site RCTs.</li> <li>• Experience in developing and implementing trial monitoring, safety, and quality assurance plans.</li> <li>• Familiarity with IRB/ethics submissions, regulatory requirements, and amendment processes.</li> <li>• Working knowledge of ICH-GCP and national guidelines</li> <li>• Authorship of peer-reviewed publications; experience in manuscript preparation.</li> <li>• Strong organizational, problem-solving, and time management skills.</li> <li>• Ability to lead and motivate multi-disciplinary teams and manage competing priorities.</li> </ul>	<ul style="list-style-type: none"> <li>• Develop SOPs or documents (as per requirement) to facilitate operations of the trial</li> <li>• Plan and deliver site trainings, capacity-building activities, and refresher sessions for clinical and research staff.</li> <li>• Contribute to progress reports, regulatory submissions, and scientific manuscripts.</li> <li>• Provide oversight of IMP management, including receipt, storage, accountability, reconciliation, and destruction, ensuring adherence to protocol, GCP, and regulatory requirements.</li> <li>• Lead site-level interactions, troubleshooting, and problem resolution; ensure adherence to project timelines and quality standards.</li> <li>• Facilitate stakeholder meetings, site visits, and regulatory engagements as required.</li> <li>• The candidate will be based at THSTI Faridabad with regular visits to hospital site(s) in Delhi.</li> </ul>
--	--	---	--

उपर्युक्त मानदंडों को पूरा करने वाले इच्छुक उम्मीदवार, उपर्युक्त पदों के लिए उल्लिखित तिथि पर, सुबह 9:00 बजे लिखित परीक्षा / कौशल परीक्षण / साक्षात्कार के लिए टीएचएसटीआई, एनसीआर बायोटेक साइंस क्लस्टर, तीसरा माइलस्टोन, फरीदाबाद-गुरुग्राम एक्सप्रेसवे, फरीदाबाद - 121001 में आ सकते हैं।

For posts mentioned above, interested candidates fulfilling the criteria as mentioned above may walk-in for written test/skill test/interview at 9:00 am on the date mentioned against each positions at THSTI, NCR Biotech Science Cluster, 3<sup>rd</sup> Milestone, Faridabad-Gurugram Expressway, Faridabad – 121001.

**नोट/ NOTE:-**

**1) The candidates must bring their latest CV/resume, one set of photocopy of documents in support of their educational qualification and experience along with their originals and a valid ID card 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 at THSTI are mandatorily required to deposit their mobile phone along with 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 qualifications 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 eligibility criteria.
- d) The date of Interview of the respective post will be the **CRUCIAL DATE** for determining eligibility with regard to age, experience, essential qualification 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 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 defense 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 many candidates appear for the walk-in interview, 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”**

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

=====**End of the document**=====