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/22/2025 **दिनांक: 12 नवंबर 2025**

RECRUITMENT NOTICE NO.: THS-C/RN/22/2025 **Dated: 12 November 2025**

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**	वरिष्ठ परियोजना सहयोगी /Sr. Project Associate
	पदों की संख्या/Number of the post	01
	परियोजना का नाम/Name of the Project	INDIGO Effective and Affordable flu Vaccine for the world
	वेतन/Emoluments	Rs. 42,000/- + HRA
	उम्र/Age	35 years
	न्यूनतम शैक्षिक योग्यता और अनुभव/Minimum Educational Qualification and Experience	• Graduate degree (3-year course) in Life Sciences with a minimum of two years of experience in clinical research, **OR** • Professional degree in MBBS, BDS, BVSc, BAMS, BHMS, BUMS, BSMS, BNYS, B.Sc. Nursing, BPT, B.Pharm, with a minimum of one year of experience in clinical research **OR** • Postgraduate or PhD in Life Sciences with a minimum of one year of experience in clinical research. **Good Clinical Practice (GCP) certification is mandatory.**
	नौकरी का प्रोफ़ाइल/ Job profile	The **Senior Project Associate** plays a pivotal role in supporting the execution and management of clinical trials across all stages. Working under the direction of the Project Manager and study monitors, they ensure the smooth operation of daily trial activities, maintain regulatory and trial documentation, and coordinate communication and logistics across study stakeholders. This role is expected to work independently on assigned tasks, proactively identify potential issues, and propose process improvements. **<u>Key Responsibilities:</u>** **Clinical Trial Support & Documentation** • Coordinate and track the distribution and reconciliation of clinical trial supplies, laboratory kits, and investigational products to investigational sites. • Ensure timely delivery and tracking of essential study documents and materials in accordance with study timelines and site activation plans. • Maintain and regularly update trial tracking tools (e.g., enrollment logs, regulatory document trackers, training logs).

		Regulatory & Site Start-up Support • Support CRAs and site staff in the collection, review, and tracking of essential regulatory documents for ethics committee and regulatory authority submissions. • Assist in preparing site initiation packages and supporting site readiness for activation. • Liaise with regulatory, legal, and contracts departments to ensure timely processing of site contracts and confidentiality agreements. **Trial Master File (TMF) Oversight** • Lead TMF set-up and ongoing maintenance, ensuring completeness, accuracy, and audit-readiness of clinical documentation. • Perform periodic TMF quality control (QC) checks and contribute to TMF metrics and reconciliation activities. • Support the development and implementation of TMF filing plans and oversight reports. **Meeting Coordination & Communication** • Schedule, coordinate, and document clinical team meetings, site communications, and teleconferences; maintain meeting agendas and minutes. • Assist with the organization and execution of investigator meetings, including logistics, preparation of materials, and follow-up documentation. **Data & Site Management** • Support CRAs with clinical data flow, Case Report Form (CRF) tracking, and resolution of data queries with investigational sites. • May accompany CRAs on monitoring visits to gain on-site experience and provide additional support during critical phases of the trial. **Cross-functional & Operational Support** • Serve as the central point of contact for the clinical team regarding project-specific communications and documentation. • Collaborate with Clinical Portfolio Management, Regulatory Affairs, Data Management, and Quality Assurance to ensure alignment of deliverables. • Support budget tracking, invoice verification, and financial documentation coordination related to trial expenses and vendor contracts. **Quality & Compliance** • Ensure adherence to ICH-GCP, applicable regulatory requirements, and internal Standard Operating Procedures (SOPs). • Participate in internal and external audits and inspections as needed; support audit readiness and CAPA (Corrective and Preventive Action) implementation.

		• Contribute to internal quality initiatives and process optimization efforts.
	कौशल /Skills	• Strong understanding of ICH-GCP guidelines and clinical trial lifecycle. • Experience working with electronic Trial Master File (eTMF), Clinical Trial Management Systems (CTMS), and document management platforms. • Excellent communication, organizational, and problem-solving skills. • Detail-oriented with the ability to manage multiple tasks and prioritize effectively. • Proficient in MS Office Suite (Word, Excel, PowerPoint, Outlook).
वॉक-इन साक्षात्कार की तिथि/ Date of walk-in interview:		**01st December 2025 @09:00 AM at THSTI, NCR Biotech Science Cluster, 3rd Milestone, Faridabad-Gurugram Expressway, Faridabad – 121001.**
2.	**पद का नाम/Name of the post**	प्रधान परियोजना सहयोगी /Principal Project Associate
	पदों की संख्या/Number of the post	01
	परियोजना का नाम/Name of the Project	INDIGO Effective and Affordable flu Vaccine for the world
	वेतन/Emoluments	Rs. 49,000/- + HRA
	उम्र/Age	35 years (relaxation as per Government of India rules)
	कार्य स्थल/Job Location	**Christian Medical College (CMC), Vellore**
	न्यूनतम शैक्षिक योग्यता और अनुभव/Minimum Educational Qualification and Experience	• Bachelor's in medical sciences with minimum three years of relevant clinical trial monitoring experience or clinical trial/study coordinator, or clinical trial/study associate experience. OR • Master's degree/ diploma, life sciences, pharmacy, public health, healthcare or other related discipline with minimum 2 years of relevant clinical trial monitoring experience or clinical trial/study coordinator, or clinical trial/study associate experience. OR • MBBS/ BDS/ BHMS/ BAMS/ BPT with a minimum of 2 years of relevant clinical trial monitoring or clinical trial/study coordinator, or clinical trial/study associate experience.
	नौकरी का प्रोफ़ाइल/ Job profile	• The Principal Project Associate/ Clinical Research Associate conducts monitoring visits for the assigned trial protocol and trial sites. Overall, the responsibilities are to ensure that the trial is conducted in accordance with the protocol, standard operating procedures, good clinical practice, and applicable regulatory requirements. • Performs site monitoring throughout the trial, which involves visiting the trial sites regularly (site initiation to site closeout) in accordance with the 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 center 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 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 onto 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. • Evaluates 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 Clinical Portfolio Management on other projects as directed and with other internal departments on their requirements as and when required. **Skills:** • Computer skills, including proficiency in the use of Microsoft Office applications • Basic knowledge and ability to apply GCP and applicable regulatory guidelines. • Strong written and verbal communication skills, including a good command of English, are required. • Excellent organizational and problem-solving skills. • Effective time management skills and ability to manage competing priorities.
वॉक-इन साक्षात्कार की तिथि/ Date of walk-in interview:		**01st December 2025 @09:00 AM at THSTI, NCR Biotech Science Cluster, 3rd Milestone, Faridabad-Gurugram Expressway, Faridabad – 121001.**
3.	**पद का नाम/Name of the post**	**वरिष्ठ क्लिनिकल अनुसंधान सहयोगी/ Senior Clinical Research Associate**
	पदों की संख्या/Number of the post	03

	परियोजना का नाम/Name of the Project	**Effect of Immediate Kangaroo Mother Care (iKMC) on neonatal mortality and culture-positive sepsis in low-birth-weight neonates in district hospitals in Chhattisgarh, India: a stepped-wedge cluster randomized trial**
	वेतन/Emoluments	INR 75,000 per month
	उम्र/Age	40 years
	स्थान/Location	**एम्स रायपुर/Based at AIIMS Raipur and travel to 10 District Hospitals in Chhattisgarh**
	न्यूनतम शैक्षिक योग्यता और अनुभव/Minimum Educational Qualification and Experience	• Bachelor's in Life Sciences with a minimum of three years of relevant clinical trial monitoring or clinical trial/study coordinator, or clinical trial/study associate experience. OR • Master's degree/ diploma, life sciences, pharmacy, public health, healthcare or other related discipline with a minimum of 2 years of relevant clinical trial monitoring or clinical trial/study coordinator, or clinical trial/study associate experience. OR • MBBS/ BDS/ BHMS/ BAMS/ BPT with a minimum of 2 years of relevant clinical trial monitoring or clinical trial/study coordinator, or clinical trial/study associate experience.
	नौकरी का प्रोफ़ाइल/ Job profile	• The Sr. 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 monitoring visits (on-site and remote), including initiation, routine monitoring, and closeout. • Ensure trial sites comply with regulatory, protocol, and GCP requirements. • Conduct risk-based monitoring and escalate site issues and protocol deviations appropriately. • Verify informed consent and subject safety in alignment with ethical standards. • Monitor AE/SAE reporting timelines to ensure compliance with regulatory requirements and escalate delayed submissions to the pharmacovigilance team. • Review source documents and CRFs to verify data accuracy and consistency (SDV). • Ensure appropriate management and documentation of investigational product (IP). • Maintain essential trial documents in accordance with ICH GCP and local regulations. • Prepare detailed monitoring visit reports and manage action items. • Support regulatory and ethics submissions, patient recruitment, and resolution of data queries. • Provide training to site personnel on study protocols, GCP, and SOPs.

		• Ensure timely delivery and proper handling of study supplies and investigational product. • Monitor quality metrics and assist with CAPA implementation. • Ensure site readiness for audits and regulatory inspections. • Use clinical trial systems (EDC, CTMS, eTMF) for tracking, documentation, and communication. • 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. • 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:		**02nd December 2025 @09:00 AM at THSTI, NCR Biotech Science Cluster, 3rd Milestone, Faridabad-Gurugram Expressway, Faridabad – 121001.**
4.	**पद का नाम/Name of the post**	डाटा प्रबंधक/Data Manager
	पदों की संख्या/Number of the post	01
	परियोजना का नाम/Name of the Project	INDIGO Effective and Affordable flu Vaccine for the world
	वेतन/Emoluments	67,000 + HRA
	उम्र/Age	40 years
	न्यूनतम शैक्षिक योग्यता और अनुभव/Minimum Educational Qualification and Experience	**Essential:** • Master's degree in any field preferably in science, with 6 years of post-qualification experience in clinical data management/ clinical research/ operations/MIS/data analysis/IT/computer science/ healthcare field OR • Graduation degree in any field preferably in science, with 8 years of post-qualification experience in clinical data management/ clinical research/operations/MIS/data analysis/IT/computer science/ healthcare field **Desirable:** • 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 प्रोफ़ाइल/ Job profile	**Responsibilities** • 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 Guidelines etc. • Prepare data transfer guidelines for external data loads and self-evident correction charts. • Manage change requests and coordinate approvals from sponsors and stakeholders. • Design and review Case Report Forms (CRFs/eCRFs) aligned with protocol requirements. • Oversee and quality check of clinical database setup, validation programming, annotated CRFs, data extract views, and final data listings. • Ensure adherence to standards such as CDISC, SDTM, and institutional SOPs. • Reconcile adverse event (AE/SAE) data and integrate medical coding dictionaries (e.g., MedDRA, WHODrug). • 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. • Maintain strict compliance with GCP, ICH, and data protection regulations. • Ensure participant confidentiality and secure handling of all clinical trial data. • Train and supervise site staff in data entry and protocol compliance. • Sound knowledge of Clinical Database Development tools, logics and techniques and GCDMP • Generate interim reports and review of listings of data for clinical trial status and data extraction in collaboration with the statistician • Generating ad-hoc reports as needed to support project oversight and decision-making. • Maintain oversight of data status reports for internal and external communication. • Maintaining and archiving of clinical study related documents • Participates in cross functional team meetings & external client meetings as DM representative • 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 Biorepository Management Systems (BMS) for tracking specimen status and turnaround time. • 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. • Development of Standard Operation Procedures and training to the study team • Supervise DM activities at the clinical site.
	कौशल /Skills	• Good management & leadership skills • Familiarity with GCP, US-FDA 21 CFR 11, regulatory requirements and data standardization guidelines. • IT literate (experience with Microsoft based applications and other CDMS applications) • Must have experience in handling EDC tools • Validation programming • 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:		**26th November 2025 @09:00 AM at THSTI, NCR Biotech Science Cluster, 3rd Milestone, Faridabad-Gurugram Expressway, Faridabad – 121001.**

नोट:1) उम्मीदवारों क्रम संख्या 1, 2, 3 ओर 4 के पदों के लिए आवेदनको अपना 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 post of S.No. 1,2, 3 & 4 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) 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.

f) 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.

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

h) All communications will only be made through email.

i) 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.

j) The no. of vacancy indicated above may change subjected to the actual requirement at the time of Written test/skill test/interview.

k) With regard to any provisions not covered in this notification, the bye laws of THSTI / Govt. of India rules/ guidelines shall prevail.

l) 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

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