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লোকপ্ৰিয় গোপীনাথ বৰদলৈ ক্ষেত্ৰীয় মানসিক স্বাস্থ্য সংস্থান

তেজপুৰ: অসম: পিন: 784001

LGB REGIONAL INSTITUTE OF MENTAL HEALTH

(An Autonomous body under Ministry of Health and Family Welfare, Govt. of India)

Website: www.lgbrimh.gov.in, e - Mail: mail@lgbrimh.gov.in

Post Box No. 15:: FAX No. (03712) 233623

TEZPUR:: 784001 :: ASSAM

LGB/NAZ/4060/26/5195

Date: 07.09.2026

Part 1- NOTICE INVITING REQUEST FOR QUALIFICATION (RFQ)

The Director, Lokopriya Gopinath Bordoloi Regional Institute of Mental Health (LGBRIMH), Tezpur, invites Requests for Qualification (RFQ) from experienced and financially sound agencies/firms/companies/consortia for establishment, operation and maintenance of a CT Scan facility within the Institute premises on Public-Private Partnership (PPP) basis, with patient charges not exceeding the applicable Central Government Health Scheme (CGHS) rates. Submissions will be received online only through the CPPP e-procurement portal (<https://eprocure.gov.in>). The RFQ document may be downloaded free of cost from the Institute website and the portal.

RFQ Information Summary

Particulars	Details
Name of work	Establishment, Operation and Maintenance of CT Scan Services on PPP basis at LGBRIMH, Tezpur
Facility	One CT Scan facility with a brand-new whole-body multi-slice CT scanner, minimum 32 slices, operated 24x7x365 within the Institute premises
Place of execution	LGBRIMH campus, Tezpur, District Sonitpur, Assam - 784001
Term (indicative, for the concession to be awarded at the RFP stage)	Seven (7) years from the Commercial Operation Date, extendable by up to three (3) years subject to satisfactory performance
Publication	CPPP (https://eprocure.gov.in and Institute website www.lgbrimh.gov.in)
Mode of submission	Online only, through the stated portal, with a valid Class-III Digital Signature Certificate of the authorised signatory. Physical submissions will not be accepted.
RFQ fee / EMD / application fee	Nil. No fee, deposit or security of any kind is payable at the RFQ stage.
Date of issue	07.09.2026

Particulars	Details
Last date for pre-RFQ queries	14.09.2026, by e-mail/post to the Nodal Officer
Pre-RFQ meeting (optional)	16.09.2026 at 11:00 hrs, MS Conference Room, Administrative Building LGBRIMH, Tezpur and simultaneously by video-conference; attendance is optional and not an eligibility condition
Last date and time of submission	28.09.2026 up to 16:00 hrs
Opening of submissions	29.09.2026 at 16:00 hrs
Validity	180 days from the last date of submission of the RFQ
Financial bid at this stage	None. No price, rate, revenue share or other financial information shall be submitted or evaluated at the RFQ stage.
Nodal Officer	Dr. Hemanta Dutta, OSD & Deputy Director, LGBRIMH, Tezpur Phone no.8473062148, Email :- drhd1984@gmail.com

Nature of this RFQ: This RFQ is a preliminary step only, for identifying and shortlisting capable agencies for a subsequent Request for Proposal (RFP). It is not a tender, auction, rate quotation or bid; it involves no price evaluation; and it creates no right in any respondent to be shortlisted, invited to the RFP, selected or paid, and no commitment by the Institute to proceed with the project. The RFP will be a separate document issued to shortlisted agencies only.

Sd/-

DIRECTOR,

Lokopriya Gopinath Bordoloi Regional Institute of Mental Health,
Tezpur, District- Sonitpur, Assam.

Part 2 - Background, Objective and Intended RFQ Outcome

2.1 Lokopriya Gopinath Bordoloi Regional Institute of Mental Health (LGBRIMH), Tezpur, is an autonomous institute under the Ministry of Health & Family Welfare,

Government of India, and the apex mental-health care, teaching, training and research institution of the North Eastern Region, serving the eight North Eastern States.

- 2.1.1 This RFQ and the subsequent RFP are being issued in compliance with GFR 2017 (Chapter 6: Procurement of Non-Consulting Services) and the DEA/Ministry of Finance Guidelines for two-stage bidding in PPP projects, as applicable to zero-CAPEX concession arrangements where the private partner bears the entire capital expenditure.
- 2.2 A substantial proportion of the Institute's in-patients and out-patients require CT-based neuroimaging for differential diagnosis of organic brain syndromes, dementia, seizure disorders, head injury, alcohol- and substance-related neurological complications, pre-ECT evaluation, treatment planning and medico-legal certification, besides general body CT examinations for co-morbid physical conditions.
- 2.3 In the absence of an in-house CT facility, patients are presently referred to external centres, involving escorted transport of psychiatric in-patients with attendant safety and custodial risk, higher out-of-pocket expenditure, and avoidable diagnostic delay. Establishment of an on-campus CT facility on PPP mode will secure affordable, quality-assured imaging within the Institute without capital expenditure from public funds.
- 2.4 Objective: to identify capable, experienced and financially sound agencies interested in establishing, operating and maintaining the CT facility on PPP basis, and to shortlist them for the subsequent RFP, in which the concession will be competitively awarded.
- 2.5 Intended RFQ outcome: (a) a published shortlist of agencies meeting the minimum eligibility criteria in Part 4; (b) invitation of only the shortlisted agencies to the RFP; and (c) no selection, award, reservation of scope or commitment of any kind at this stage. The Institute may, in accordance with applicable rules and portal procedure, modify the requirement, re-advertise, or discontinue the process at any stage.
- 2.6 The prospective arrangement: the selected agency will finance, supply, install, test, commission, operate and maintain the facility at its own cost and risk for the concession term; the Institute will provide licensed space and supporting utilities; patient charges will be capped at applicable CGHS rates; and the Institute's consideration will be a share of the gross revenue, the exact financial structure being disclosed only in the RFP.

Part 3 - Scope and Information Required from Respondents

- 3.1 Indicative scope of the prospective concession (the complete scope will be issued with the RFP):
- (a) By the Institute: licensed built-up space of approximately **48 sq. m.** at the Ground floor, Hospital Block, free of rent; electricity and water through a sub-meter installed by the agency at its cost and payable at actuals; captive patient flow from

OPD, IPD, emergency, forensic and outreach services; facilitation with the Institute's cash-collection counter, HMIS/patient-registration integration and general supervision through a Nodal Officer and Monitoring Committee.

- (b) By the selected agency (at the RFP/contract stage): site preparation at its cost, including civil, electrical, air-conditioning, fire-safety, networking and radiation shielding conforming to the AERB-approved room layout; supply, installation, testing and commissioning of a brand-new, current-production whole-body multi-slice CT scanner, minimum 32 slices, with dual-head pressure injector, post-processing workstation, DICOM printer, online UPS, emergency accessories and CCTV; all statutory licences and approvals including AERB eLORA registration and type approval, Bio-Medical Waste Management authorisation, clinical establishment registration, fire NOC, trade licence and labour registrations; 24x7x365 operation with qualified staff (radiologist MD/DNB registered with NMC/State Medical Council; AERB-certified Radiation Safety Officer/technologists; staff trained in contrast-reaction management and basic life support; attendants sensitised to the care of psychiatric patients); all plain and contrast CT examinations within machine capability at CGHS-capped rates; free examinations for notified categories (BPL, EWS, destitute and undertrial/court-referred patients) as specified in the RFP; digital (DICOM) images and signed reports with turnaround of 60 minutes for emergency and 24 hours for routine studies; equipment uptime of not less than 95% per month; comprehensive maintenance with committed on-site response; biomedical waste disposal; records, registers and monthly MIS; patient-data confidentiality and applicable radiology safeguards; and decommissioning/handover on expiry or termination.

3.2 Information required from respondents (capability information only; no financial information):

- (a) Profile of the agency: legal status, year of establishment, ownership, and particulars of key personnel;
- (b) Experience of establishment and/or operation of CT scan or comparable diagnostic imaging services, with client details and completion/performance certificates;
- (c) Financial capacity summary: turnover from healthcare/diagnostic operations and net worth, with audited statements and CA certificate (with UDIN);
- (d) Statutory registrations: PAN, GSTIN, EPFO/ESIC (or undertaking to obtain before the Commercial Operation Date);
- (e) AERB eLORA licences for existing CT installation(s), or an undertaking to obtain registration before the Commercial Operation Date;
- (f) Proposed equipment configuration (indicative) with the OEM data sheet; final technical specifications will be issued with the RFP;

- (g) Proposed manpower deployment (position-wise with qualifications) and an outline implementation schedule from award to commercial operation;
- (h) Declarations at Part 7, and any other information specifically sought in the prescribed forms.

3.3 Respondents must not submit price, rate, revenue-share, premium or any other financial bid with the RFQ. Any such information included in a submission will be ignored, and its inclusion may render the submission liable to rejection.

Part 4 - Minimum Eligibility and Submission Requirements

4.1 Submissions will be considered only from applicants satisfying all of the following minimum criteria, supported by self-attested documentary evidence:

Sl.	Criterion	Requirement	Supporting documents
1	Legal status	Company, LLP, society, trust, partnership firm or proprietorship registered under Indian law. Consortium of not more than two members permitted; the Lead Member must meet the experience and financial criteria in full and hold not less than 51% participation.	Certificate of incorporation/registration; MoA/AoA, partnership or trust deed
2	Experience	Minimum three (3) completed years of establishment and/or operation of CT scan or comparable diagnostic imaging services in the last seven (7) years, in at least one (1) facility. Sector-neutral: Government and private sector experience will be counted equally.	Work orders/agreements with completion or satisfactory-performance certificates from clients
3	Financial capacity	Average annual turnover of not less than Rs. 2.00 crore from healthcare/diagnostic operations in the last three audited financial years, and positive net worth in the latest audited financial year.	Audited financial statements; CA certificate with UDIN
4	Statutory registrations	Valid PAN and GSTIN; EPFO and ESIC registrations, or undertaking to obtain them before the Commercial Operation Date.	Registration certificates / undertaking on letterhead
5	AERB capability	Valid eLORA licence(s) for existing CT installation(s), or undertaking to obtain AERB licence before the Commercial Operation Date.	Licence copies / undertaking

Sl.	Criterion	Requirement	Supporting documents
6	Non-blacklisting	Not blacklisted, debarred or suspended by any Central/State authority, PSU, autonomous body or local authority as on the date of submission.	Self-attested declaration (Form 2)
7	Bidders from countries sharing a land border with India	Declaration of compliance with the restrictions applicable to bidders from countries sharing a land border with India, including registration with the Competent Authority where applicable, and beneficial-ownership particulars as required under applicable instructions.	Declaration (Form 3)

Clarification on Make in India (MII): As this project operates on a 100% private CAPEX concession model where the Institute does not purchase or fund the capital equipment, mandatory public procurement Make in India (MII) purchase preference quotas do not apply to the concessionaire's capital assets. However, domestic sourcing of equipment is encouraged where clinically and commercially viable.

4.2 Only the documents listed against each criterion (and the forms at Part 7) are required.

Self-attested copies are acceptable; original documents will be verified, if required, only in respect of shortlisted agencies. No notarised affidavit, stamp-paper declaration, tender fee, earnest money deposit, performance security or Integrity Pact is required at the RFQ stage.

4.3 Submission must be made online through the stated portal, uploaded as a single technical submission in the prescribed forms, digitally signed by the authorised signatory. The financial bid will be sought only at the RFP stage.

4.4 Relaxations: prior turnover and prior experience may be relaxed for DPIIT-recognised Startups, and Micro & Small Enterprises registered on Udyam enjoy all exemptions applicable under Government policy, including from any fee or deposit that may be prescribed at a subsequent stage. Technical, clinical, radiation-safety and quality requirements will not be relaxed for any category of applicant.

Part 5 - Evaluation, Shortlisting and Next Procurement Stage

5.1 Evaluation will be document-based and confined to a responsive/not-responsive (pass/fail) assessment against the published minimum criteria in Part 4. No marks, scoring, ranking, presentation, demonstration or site visit will be used at this stage; if any such step is later considered indispensable, it will be disclosed by corrigendum before the submission date. The Institute may verify any claim with the issuing authority.

- 5.2 All applicants found responsive against every criterion will be shortlisted. The shortlist will be published on the portal/Institute website. No criterion or sub-criterion not stated in this document will be applied.
- 5.3 Next stage: only shortlisted agencies will be invited to a separately issued Request for Proposal (RFP). The RFP will contain the complete scope, line-by-line technical specifications, CGHS-based financial model, performance standards, service levels, contract conditions and the evaluation method, including the financial bid parameter - the percentage of gross monthly revenue offered to the Institute, evaluated as the highest revenue share subject to a notified reserve and a sustainability check. These parameters are disclosed in advance only for transparency; they will be sought and evaluated exclusively under the RFP.
- 5.3.1 Clarification on RFP Financial Evaluation Method: It is expressly clarified that the award of the concession at the RFP stage will NOT be based on L1 (Lowest Bidder) or QCBS (Quality and Cost Based Selection) parameters. The award will be determined strictly on an “H1 (Highest Bidder)” basis, defined as the agency offering the highest percentage of gross monthly revenue to the Institute, subject to a notified reserve floor and a financial sustainability/viability check of the bid. This H1 mechanism is standard for zero-CAPEX PPP concessions under DEA guidelines, where the concessionaire’s revenue-share offer is the sole financial evaluation parameter at the RFP stage.
- 5.4 Participation in this RFQ confers no right to be invited to the RFP beyond the published shortlisting criteria, and no right to award, reimbursement or exclusive dealing of any kind.

Part 6 - Essential Terms and Conditions

- 6.1 This RFQ is not a tender, contract, work order, purchase order or assurance of award. It does not create a contract, any commitment to procure, or any right to award; no payment, reimbursement or compensation of any kind is due to any respondent in any circumstance.
- 6.2 All costs of preparing and submitting the RFQ, including portal charges if any, are borne by the respondent.
- 6.3 The Institute reserves the right to accept or reject any or all submissions, in whole or in part, to modify the requirement, or to annul the process at any stage, for reasons

recorded in writing, without liability and without assigning reasons to respondents, in accordance with applicable rules and portal procedure.

- 6.4 All information submitted must be true, complete and capable of verification. Furnishing false or fabricated information may lead to rejection of the submission and to such further action as is warranted under law, including debarment under applicable Government instructions at the RFP or contract stage.
- 6.5 Confidential business information will be handled in accordance with applicable law and Government procedure; however, submissions are subject to statutory disclosure obligations. Claimed confidentiality must be specifically identified with reasons; blanket confidentiality claims will not be accepted.
- 6.6 No price evaluation, commercial ranking or selection will take place at the RFQ stage. Any indicative cost or volume information volunteered by a respondent will not be evaluated and will create no entitlement or preference.
- 6.7 Canvassing in any form, or any attempt to influence any officer or employee of the Institute, will render the submission liable to summary rejection.
- 6.8 No person in the service of the Institute or of Government shall have any direct or indirect interest in an applicant agency or in any resulting arrangement.
- 6.9 Corrigenda and addenda, if any, will be published only on the portal and the Institute website; applicants should check these regularly; no individual communication will be issued.
- 6.10 The Institute may verify any claim with the issuing authority and may inspect the applicant's existing facilities, on reasonable notice, for the limited purpose of confirming eligibility claims.
- 6.11 Terms governing the concession itself - including the concession term and extension, security deposit and performance security, payment mechanics, revenue-share and minimum guarantee (if any), insurance, indemnity, termination, penalties and dispute resolution - will be issued with the RFP and are not part of this RFQ.
- 6.12 Disputes relating to this RFQ are subject to the jurisdiction of the courts at Tezpur, District Sonitpur, Assam, and to Indian law.
- 6.13 Clinical, radiation and service safeguards: nothing in this RFQ dilutes any statutory requirement. The facility, when established, will be required to comply with the Atomic Energy (Radiation Protection) Rules, 2004 and AERB requirements (including eLORA registration, type approval and a certified Radiation Safety Officer), the Bio-Medical Waste Management Rules, 2016, applicable clinical establishment and fire-safety requirements, the Mental Healthcare Act, 2017 in the care of persons with mental illness, the Pre-conception and Pre-natal Diagnostic Techniques (PC-PNDT) Act, 1994, the Digital Personal Data Protection Act, 2023, and applicable labour laws; and to maintain the service standards (24x7 operation, uptime, turnaround,

free/subsidised categories, CGHS rate cap) indicated in Part 3. Full obligations will be specified in the RFP and the Agreement.

6.14 Code of Integrity: The Code of Integrity under Rule 175 of GFR 2017 shall strictly apply to this procurement. Any bidder found to have engaged in any corrupt, fraudulent, collusive, coercive, or obstructive practice will be disqualified and may face debarment action.

Part 7 - Minimum Forms / Declarations

7.1 Submissions must use only the following forms; all declarations are self-attested.

Form	Contents	Execution
Form 1	Covering letter and applicant profile summary (identity, constitution, authorised signatory with Board Resolution/Power of Attorney)	Self-attested
Form 2	Declaration of truthfulness of information and non-blacklisting/debarment/suspension status of the agency and its proprietors/partners/directors	On the applicant's letterhead, self-attested
Form 3	Declaration of compliance with the restrictions applicable to bidders from countries sharing a land border with India (including beneficial-ownership particulars, where required)	Prescribed Government format, self-attested
Form 4	Undertaking to obtain AERB eLORA registration, EPFO/ESIC and other statutory registrations before the Commercial Operation Date (only where the applicant does not already hold them)	Self-attested
Form 5	Document checklist (consolidated list of enclosures)	Signed by the authorised signatory

7.2 All pages of the submission will be serially numbered and self-attested by the authorised signatory with the seal of the agency.

7.3 The formats of Forms 1 to 5, to be used for submission without alteration of substance, are set out in the Annexures following this Part.

Annexures - Prescribed Forms (Forms 1 to 5)

The following forms constitute the minimum and exclusive formats for the RFQ submission referred to in Part 7. Submissions must use these formats without alteration of substance; additional sheets may be appended only where expressly permitted. All declarations are self-attested; no notarised affidavit, stamp-paper declaration, tender fee or earnest money deposit is required. Where a field is not applicable to an applicant, "Not Applicable" shall be entered against it; no field shall be left blank.

Form 1 - Covering Letter and Applicant Profile Summary

(To be submitted on the applicant's letterhead, self-attested)

To,

The Director,

Lokopriya Gopinath Bordoloi Regional Institute of Mental Health (LGBRIMH),

Tezpur - 784001, District Sonitpur, Assam.

Subject: Submission of Request for Qualification for establishment, operation and maintenance of CT Scan services on Public-Private Partnership (PPP) basis at LGBRIMH, Tezpur - RFQ No. LGB/NAZ/4060/26/5195 Dated: 07-09-2026

Reference: (i) RFQ No **LGB/NAZ/4060/26/5195** **Dated: 07-09-2026** published on the CPPP e-procurement portal (<https://eprocure.gov.in>); and (ii) corrigenda and addenda thereto, if any.

Sir/Madam,

Having examined the RFQ document, including all corrigenda and addenda published on the portal, we, [name of the agency], hereby submit our Request for Qualification for establishment, operation and maintenance of CT Scan services on PPP basis at LGBRIMH, Tezpur. We confirm that: (i) the information furnished and the documents annexed are true, complete and capable of verification; (ii) we meet each of the minimum eligibility criteria at Part 4 of the RFQ document; (iii) we unconditionally accept the terms and conditions at Part 6 of the RFQ document; (iv) we understand that this RFQ is not a tender and that participation confers no right to shortlisting, invitation to the Request for Proposal or award; (v) no price, rate, revenue-share or other financial bid is included in this submission; and (vi) this submission, and the eligibility status claimed therein, shall remain valid for one hundred eighty (180) days from the last date of submission of RFQs.

Applicant Profile

Sl.	Particulars	Details
1	Name of the agency	[_____]
2	Constitution / legal status	[Company / LLP / Society / Trust / Partnership firm / Proprietorship]

Sl.	Particulars	Details
3	Constitutional documents (CIN/LLPIN/registration number, as applicable)	[_____]
4	Consortium (Yes / No)	[No / Yes - lead member: [_____]; participation of lead member: [_____]%]
5	Year of establishment	[_____]
6	Registered office address	[_____]
7	PAN	[_____]
8	GSTIN	[_____]
9	EPFO registration number	[_____ / to be obtained before the Commercial Operation Date - see Form 4]
10	ESIC registration number	[_____ / to be obtained before the Commercial Operation Date - see Form 4]
11	AERB eLORA licence number(s) held for existing CT installation(s)	[_____ / not held - see Form 4]
12	Authorised signatory (name, designation, e-mail, telephone)	[_____]
13	Authorisation for the signatory (Board Resolution / Power of Attorney)	[Enclosed at page no. _____]

Financial Capacity Summary (last three audited financial years)

Financial year	Total annual turnover (Rs. lakh)	Turnover from healthcare / diagnostic operations (Rs. lakh)	Audited statements and CA certificate with UDIN (page nos.)
FY [____-____]	[_____]	[_____]	[_____]
FY [____-____]	[_____]	[_____]	[_____]
FY [____-____]	[_____]	[_____]	[_____]

Experience particulars are to be furnished only through the work orders and completion/performance certificates listed against Criterion 2 in **Form 5**.

Place: [_____]

Signature of the authorised signatory

Date: [DD.MM.2026]

(with the seal of the agency)

Name: [_____], Designation: [_____]

Form 2 - Declaration of Truthfulness and Non-Blacklisting

(On the applicant's letterhead, self-attested)

RFQ No. LGB/NAZ/4060/26/5195

Dated:07-09-2026

Establishment, Operation and Maintenance of CT Scan Services on PPP basis at LGBRIMH, Tezpur.

1. We, [name of the agency], having our registered office at [address], in respect of the above-referenced RFQ, declare that the information furnished in the RFQ submission and its annexures is true, complete, accurate and capable of verification, and that all documents enclosed are authentic copies of originals.
2. We declare that neither the agency nor any of its proprietors, partners, directors or key managerial personnel has been blacklisted, debarred, suspended or placed on any caution/restriction list by any Central or State Government department, public sector undertaking, autonomous body, statutory authority or local authority as on the date of this declaration, and that no proceedings leading to any such action are pending.
3. We declare that no conflict of interest exists: no person in the service of the Institute or of Government has any direct or indirect interest in the agency or in the submission made by it.
4. We understand that the submission of false, misleading or fabricated information may lead to rejection of the submission, forfeiture of eligibility at the RFP stage, and such further action as is warranted under applicable law and Government instructions, including debarment.
5. This declaration is made for the sole purpose of the referenced RFQ, is made by the authorised signatory of the agency, and is self-attested.

Place: [_____]

Date: [DD.MM.2026]

Signature of the authorised signatory

(with the seal of the agency)

Name: [_____], Designation: [_____]

Form 3 - Declaration in Respect of Bidders from Countries Sharing a Land Border with India

(Self-attested; to be executed in the format and manner prescribed under the applicable instructions of the Central Government where such instructions prescribe a specific format)

RFQ No. LGB/NAZ/4060/26/5195

Dated:07-09-2026,

Establishment, Operation and Maintenance of CT Scan Services on PPP basis at LGBRIMH, Tezpur.

1. We, [name of the agency], declare whether the applicant, any of its consortium members, its beneficial owners, or the OEM of the equipment proposed to be deployed is a bidder from a country which shares a land border with India, within the meaning of the applicable instructions of the Central Government: [Yes / No].
2. If Yes, the country(ies) concerned and full particulars of the ownership and control structure (beneficial ownership) are: [_____]; supporting particulars are annexed at page no. [_____].
3. Where registration with the Competent Authority designated under the applicable instructions is required, we declare that such registration has been obtained, the registration number and date being [_____] / [DD.MM.____], and a copy is enclosed at page no. [_____].
4. We undertake to comply with all applicable instructions on this subject and confirm that no award in our favour is permissible at any subsequent stage of procurement without full compliance with the said instructions.

Place: [_____]

Date: [DD.MM.2026]

Signature of the authorised signatory

(with the seal of the agency)

Name: [_____], Designation: [_____]

Form 4 - Undertaking to Obtain Statutory Registrations and Licences

(Self-attested; required only where the applicant does not already hold the registration / licence concerned - see Part 4, Criteria 4 and 5)

RFQ No. LGB/NAZ/4060/26/5195**Dated:07-09-2026****Establishment, Operation and Maintenance of CT Scan Services on PPP basis at LGBRIMH, Tezpur.**

Where any of the registrations, licences or approvals listed below is not held by us, [name of the agency], as on the date of this RFQ, we undertake that the same shall be obtained at our cost before the Commercial Operation Date of the facility, and that failure to obtain the same before that date may entitle the Institute to such action as is specified in the RFP and the Agreement.

Sl.	Registration / licence or approval	If held: registration number, date and validity	If not held: confirmation of undertaking
1	AERB eLORA registration and equipment type approval for the proposed CT installation	[_____]	[Undertaking given]
2	Bio-Medical Waste Management authorisation	[_____]	[_____]
3	Clinical establishment registration (as applicable)	[_____]	[_____]
4	Fire safety NOC	[_____]	[_____]
5	Trade licence and labour-law registrations (as applicable)	[_____]	[_____]
6	EPFO and ESIC registration	[_____]	[_____]
7	Any other statutory approval notified in the RFP	[_____]	[_____]

It is clarified that this undertaking relates only to the timing of obtaining the listed registrations and licences; the underlying substantive requirements, including the technical, clinical and radiation-safety requirements of the RFQ, are not relaxed (Part 4.4).

Place: [_____]

Date: [DD.MM.2026]

**Signature of the authorised signatory
(with the seal of the agency)**

Name: [_____], Designation: [_____]

Form 5 - Document Checklist

(To be enclosed as the first page of the submission, signed by the authorised signatory; against each enclosure, insert the serial page numbers of the submission)

Sl.	Document / enclosure	Reference	Page numbers
1	Form 1 - covering letter and applicant profile summary, with Board Resolution / Power of Attorney authorising the signatory	Form 1	[____]
2	Certificate of incorporation / registration, with MoA / AoA or partnership / trust deed	Part 4, Criterion 1	[____]
3	Consortium agreement and particulars of lead-member participation (where applicable)	Part 4, Criterion 1	[____]
4	Work orders / agreements with completion or satisfactory-performance certificates	Part 4, Criterion 2	[____]
5	Audited financial statements for the last three financial years	Part 4, Criterion 3	[____]
6	CA certificate with UDIN (turnover and net worth)	Part 4, Criterion 3	[____]
7	PAN and GSTIN registration certificates	Part 4, Criterion 4	[____]
8	EPFO / ESIC registration certificates or undertaking	Part 4, Criterion 4	[____]
9	AERB eLORA licence(s), or Form 4 undertaking	Part 4, Criterion 5	[____]
10	Form 2 - declaration of truthfulness and non-blacklisting	Form 2	[____]
11	Form 3 - declaration regarding bidders from countries sharing a land border with India	Form 3	[____]
12	Form 4 - undertaking to obtain statutory registrations (if applicable)	Form 4	[____]
13	OEM data sheet of the proposed CT scanner (indicative configuration)	Part 3.2(f)	[____]
14	Proposed manpower deployment and outline implementation schedule	Part 3.2(g)	[____]

Documents other than those listed above are not required at the RFQ stage; additional material, if any, may be sought only from shortlisted agencies.