

ICMR-National Institute of Translational Virology and AIDS Research
INDIAN COUNCIL OF MEDICAL RESEARCH
Department of Health Research,
Ministry of Health & Family Welfare, Government of India
Plot No 73, G-Block, Bhosari, Pune-411026 (India)
Ph: 020-27331200

Advt. No:45/NITVAR/Project Cell/2026-27

Dated: 15.06.2026

Sub: Walk-in interview /written test to fill up temporary/contractual project positions under the project entitled "A Phase III, Multicenter, Randomized, Double-Blind, Placebo-Controlled Study to Evaluate the Efficacy, immunogenicity and Safety of Single dose of Dengue Tetravalent Vaccine, Live Attenuated (Recombinant, Lyophilized) – DengiAll of Panacea Biotec Limited in Healthy Indian Adults" PI-Dr. Sheela Godbole – Scientist G, Co-PI – Dr. Abhijit Kadam - Scientist-D at ICMR-NITVAR.

ICMR-National Institute of Translational Virology and AIDS Research Institute, Pune, (formerly known as NARI) is a research Institute under the aegis of Indian Council of Medical Research, Department of Health Research, Ministry of Health & Family Welfare, Government of India, New Delhi.

The institute intends to engage the following Non-Institutional project human resource position(s), purely on a temporary contract basis for our short-term research project referred in the subject above. Required qualifications and other details are given below:-

Sr.No _ 01	
Name of the Post	Consultant (Medical)
Number of vacancy	02 Post – Dengue Vaccine Trial Budget
Age Limit	i) Junior Consultant:- 40 to 50 years ii) Consultant:- 40 to 60 years iii) Senior Consultant:-40 to 70 years
Emoluments	i) Junior Consultant:- Between Rs.60,000-90,000/- ii) Consultant:- Between Rs.80,000-120,000/- iii) Senior Consultant:-Between Rs.1,00,000-1,50,000/- OR For a retired Govt. servant the remuneration shall be fixed in accordance with the Ministry of Finance, Dept. of Expenditure OM No. 3-25/2020-E.IIIA dated 09 Dec 2020.
Essential Qualification	i) Junior Consultant Professionals having proven competency and success in their area of specialization, with at least 6 years of post-qualification experience and should possess minimum MBBS or BDS or equivalent degree in relevant subject. ii) Consultant Professionals having proven competency and success in their area of specialization, with at least 8 years of post-qualification experience and should possess minimum MBBS or equivalent degree in relevant subject. iii) Senior Consultant Professionals having proven competency and success in their area of specialization, with at least 12 years of post-qualification experience and should possess minimum MBBS or BDS or equivalent degree in relevant subject. OR Retired Government employees, with requisite qualification and who were in the regular Pay Level10 and above, having at least 10 years of relevant experience in the required specialization. Note: 1. PhD will be be considered as 4 years' experience (irrespective of the duration taken to complete the degree). 2. MD/MS or equivalent to be considered as 3 years' experience. 3. MPH will not be considered as any experience.

Desirable Qualification	<ul style="list-style-type: none"> • Post Graduate Diploma in Clinical Research. • Three years' Clinical trial experience & Experience in Clinical Research • Good Knowledge of computer applications • Good English writing and communication skills. • Published papers
Scope of Work/ Nature of duties	<ul style="list-style-type: none"> • Develop study protocols, standard operating procedures (SOPs), and other essential documents in line with regulatory guidelines. • Oversee the implementation of study activities, ensuring adherence to timelines, quality standards, and Good Clinical Practice (GCP) Guidelines. • Training and Mentoring of TO and TA on protocol adherence, data collection, and ethical research practices. • To supervise and guide the supportive monitoring activities done by TO and TA • Conduct quality checks, site monitoring visits, and audits to maintain the integrity of data and study processes. • Draft and review scientific manuscripts, project reports, etc. • Review AEs and SAEs
Project Duration	Initially for a period of 6 Months

Sr.No _ 02	
Name of the Post	Consultant (Medical)
Number of vacancy	01 Post – Main Trial Budget
Age Limit	i) Junior Consultant:- 40 to 50 years ii) Consultant:- 40 to 60 years iii) Senior Consultant:-40 to 70 years
Emoluments	i) Junior Consultant:- Between Rs.60,000-90,000/- ii) Consultant:- Between Rs.80,000-120,000/- iii) Senior Consultant:-Between Rs.1,00,000-1,50,000/- OR For a retired Govt. servant the remuneration shall be fixed in accordance with the Ministry of Finance, Dept. of Expenditure OM No. 3-25/2020-E.IIIA dated 09 Dec 2020.
Essential Qualification	i) Junior Consultant Professionals having proven competency and success in their area of specialization, with at least 6 years of post-qualification experience and should possess minimum MBBS or BDS or equivalent degree in relevant subject. ii) Consultant Professionals having proven competency and success in their area of specialization, with at least 8 years of post-qualification experience and should possess minimum MBBS or BDS or equivalent degree in relevant subject. iii) Senior Consultant Professionals having proven competency and success in their area of specialization, with at least 12 years of post-qualification experience and should possess minimum MBBS or BDS or equivalent degree in relevant subject. OR Retired Government employees, with requisite qualification and who were in the regular Pay Level10 and above, having at least 10 years of relevant experience in the required specialization. Note: 1. PhD will be considered as 4 years' experience (irrespective of the duration taken to complete the degree). 2. MD/MS or equivalent to be considered as 3 years' experience. 3. MPH will not be considered as any experience.
Desirable Qualification	<ul style="list-style-type: none"> • Post Graduate Diploma in Clinical Research. • Three years' Clinical trial experience & Experience in Clinical Research • Good Knowledge of computer applications • Good English writing and communication skills. • Published papers

Scope of Work/ Nature of duties	<ul style="list-style-type: none"> • Develop study protocols, standard operating procedures (SOPs), and other essential documents in line with regulatory guidelines. • Oversee the implementation of study activities, ensuring adherence to timelines, quality standards, and Good Clinical Practice (GCP) Guidelines. • Training and Mentoring of TO and TA on protocol adherence, data collection, and ethical research practices. • To supervise and guide the supportive monitoring activities done by TO and TA • Conduct quality checks, site monitoring visits, and audits to maintain the integrity of data and study processes. • Draft and review scientific manuscripts, project reports, etc. • Review AEs and SAEs
Project Duration	Initially for a period of 6 Month

Sr.No _ 03	
Name of the Post	Consultant (Non-Medical) (Statistic)
Number of vacancy	01 Post
Age Limit	<ul style="list-style-type: none"> i) Junior Consultant:- 40 to 50 years ii) Consultant:- 40 to 60 years iii) Senior Consultant:-40 to 70 years
Emoluments	<ul style="list-style-type: none"> i) Junior Consultant:- Between Rs.60,000-90,000/- ii) Consultant:- Between Rs.80,000-120,000/- iii) Senior Consultant:-Between Rs.1,00,000-1,50,000/- <p style="text-align: center;">OR</p> <p>For a retired Govt. servant the remuneration shall be fixed in accordance with the Ministry of Finance, Dept. of Expenditure OM No. 3-25/2020-E.IIIA dated 09 Dec 2020.</p>
Essential Qualification	<ul style="list-style-type: none"> i) Junior Consultant Professionals having proven competency and success in their area of specialization, with at least 6 years of post-qualification experience and should possess minimum M.Sc equivalent qualifications in relevant subject ii) Consultant Professionals having proven competency and success in their area of specialization, with at least 8 years of post-qualification experience and should possess minimum M.Sc/ or equivalent qualifications in relevant subject. iii) Senior Consultant Professionals having proven competency and success in their area of specialization, with at least 12 years of post-qualification experience and should possess minimum M.Sc/ equivalent qualifications in relevant subject <p style="text-align: center;">OR</p> <p>Retired Government employees, with requisite qualification and who were in the regular Pay Level-10 and above and having at least 10 years experience in the required specialization.</p>
Desirable Qualification	<ul style="list-style-type: none"> • Experience in working with clinical trial Data or large databases • Experience with Clinical trial Softwares • Experience in R or any statistical software used globally for clinical trials. • Good Knowledge of computer applications • Good English writing and communication skills.
Scope of Work/ Nature of duties	<ul style="list-style-type: none"> • Monitor and ensure the accuracy, completeness, and consistency of data collected for the ongoing clinical studies. • Conduct regular data quality checks to identify and address discrepancies in collaboration with the study team. • Prepare data monitoring reports, summaries, and documentation for progress reviews and stakeholder updates in compliance with 21 CFR guidelines. • Oversee the process of data Entry, validation, and maintenance in electronic database • Contribute to development of new trial protocols – Sample Size, Quality monitoring, Data analysis plans

	<ul style="list-style-type: none"> • Coordinate with investigators and study staff to resolve data-related issues promptly • Ensure data handling compliance with ethical standards, regulatory guidelines, and confidentiality requirements. • Support statistical analysis as and when required. • Supervision data quality control staff.
Project Duration	Initially for a period of 06 Months

Sr.No _ 04	
Name of the Post	Project Research Scientist I (Medical)
Number of vacancy	01 Post
Age Limit	35 years
Emoluments	Rs.67,000/- + HRA
Essential Qualification	MBBS/ BDS
Desirable Qualification	<ul style="list-style-type: none"> • PG degree including Masters in Public Health • Diploma in Clinical Research from recognized Institute • Experience in clinical trials • Good English writing and communication skills.
Scope of Work/ Nature of duties	<ul style="list-style-type: none"> • Monitor and ensure the accuracy, completeness, and consistency of data collected for the ongoing clinical studies. • Coordinate in the overall implementation of the project for all clinical trial sites. • Preparing training materials and modules for protocol training • Organize, conduct Training of Trainers and coordinate the site level trainings • Develop Study Specific Procedures (SSP) Manual for clinical trial implementation at sites • Required for site training • Assist PI and Co-PI in overall co-ordination of trial activities, study implementation at all sites.
Project Duration	Initially for a period of 06 Months

Place of Interview/Written Test	Auditorium of ICMR-National Institute of Translational Virology and AIDS, Research, Plot No. 73, G-Block, Bhosari, Pune-411026
Date of Walk-in-Interview /Written test and Reporting Time	Date: 30.06.2026 Reporting time 09.00 – 10.00 AM (Candidates arriving late after 10.00 am will not be entertained under any circumstances)
Selection Procedure	Walk-In-Interview and/or a Written test if the number of eligible candidates are more than 30 in numbers for each position

Candidates are invited to attend a walk in interview on 30th June, 2026 at 10.30 am onwards. The Interview will be held at ICMR-National Institute of Translational Virology and AIDS Research, Bhosari Pune.

Candidates are advised to arrive on time and ensure they meet the stipulated date, time, and essential qualifications. Only those fulfilling the required eligibility criteria-pertaining to qualifications, age, experience, etc. and whose original documents are verified by ICMR-NITVAR, Pune will be eligible to participate in the interviews.

Terms and conditions: -

1. Number of positions may vary.
2. The position is meant for temporary basis and is a full time on site position.
3. The Initial term of engagement of Consultants shall be for not more than one year and subsequent

extension, if any, can be considered, on case to case basis, depending upon the job requirements and the frame for its completion, subject to fulfilment of performance evaluation made by the Competent Authority, but should not be more than Five years. In all cases, the duration of engagement of any Consultant, in ICMR (Institutes) will not be more than 05 years.

4. The rates of emoluments/stipend shown in this advertisement are consolidated.
5. Cut-off date for age limit will be as on the date of walk-in-interview.
6. Age relaxation will be as per the guidelines of ICMR.
7. Candidates meeting the age criteria and possessing the required qualifications and experience, may walk in for interview along with the duly filled in the prescribed application attached with this advertisement. All related educational documents, photograph/experience certificates, should be enclosed failing which application shall be rejected.
8. The shortlisted candidates will be allowed to appear before interview Board and candidates may check our website for updates related to the advertisement.
9. Shortlisting of candidates for interview shall be done on basis of educational qualifications and experience. However, if necessary, written test and assignment may also be conducted before the interview if the candidates appearing are more than 30 in numbers. In case of tie at any stage all the candidates who have same marks would be considered to have qualified for further rounds.
10. Selection of Candidates will be based on the performance in the interview.
11. Candidates have to submit the duly self -attested copies of proof of their age, educational qualifications, experiences, testimonials etc. at the time of joining, if selected.
12. Allotment of duties to the successful candidates will be decided by the competent authority at its discretion.
13. Qualification & experience should be in relevant discipline/field and from an Institution of repute. Experience should have been gained after acquiring the minimum essential qualification.
14. Mere fulfilling the essential qualification does not guarantee the selection.
15. Persons already in regular time scale service under any Government Department/ Organizations are not eligible to apply.
16. No TA/DA will be paid to attend interview / personal discussion and candidates have to arrange transport/accommodation themselves.
17. ICMR-NITVAR, Pune reserves rights to consider or reject any application/candidature.
18. Submission of wrong or false information during the process of selection shall disqualify the candidature at any stage.
19. The persons engaged on Human Resource Positions shall not have any claim on a regular post in ICMR or in any of its Institutes/Centers or in any Department of Government of India and their project term with breaks or without breaks in any or multiple projects will not confer any right for further assignment or transfer to any other project or appointment/absorption/regularization of service in funding agency or in ICMR. Benefits of Provident Fund, Pension Scheme, Leave Travel Concession, Medical claim, Staff Quarters and other facilities applicable to the regular staff of ICMR etc. are not admissible to the project human resource positions.
20. ICMR-NITVAR, Pune reserves the right to terminate the positions even during the agreed contract period or extended contract period without assigning any reason.
21. Leave shall be as per the ICMR's policy for the Consultant/Young Professional-II human resource positions.
22. ICMR-NITVAR, Pune reserves the right to cancel/modify the process at any time, at its discretion.
23. The decision of the Competent Authority will be final and binding.
24. Canvassing in any form will be a disqualification.
25. Corrigendum/addendum/further information; if any; in respect of this advertisement, will be published on our website only. Hence, the candidates are advised to see the website of ICMR, New Delhi & ICMR-NITVAR, Pune regularly for further updates related to this advertisement.

Candidate must bring his/her duly filled in application form in the prescribed format with a recent passport size color photograph along with a detailed bio- data/C.V. and all relevant documents; **in original**; with one set of self-attested photocopies, in proof of his/her educational qualifications [all certificates and mark-sheets from 10th Std. onwards], working experience, age, caste and **photo ID** [Aadhar Card/Indian Passport/PAN Card/Driving License] etc., at the time of attending the Walk-in-interview.

Director
ICMR-NITVAR, Pune

ICMR- NATIONAL INSTITUTE OF TRANSLATIONAL VIROLOGY AND AIDS RESEARCH,
PUNE

(Indian Council of Medical Research, New Delhi)

BIO-DATA

Vacancy Notification for 1) Consultant (Medical)

2) Consultant (Non-Medical) 3) Project Research Scientist – I (Medical)

Advertisement No.:- 45/NITVAR/Project Cell/2026-27

1. Name of the Project Post, applied for: _____

2. Name in full (IN BLOCK LETTERS) :

(First Name) (Middle Name) (Surname/ Last Name)

3. Mother's Name : _____

4. Father's Name : _____

5. Husband's Name : _____

6. Address for Correspondence :

7. Tel/Mobile No. _____

8. E-mail ID: _____

9. Permanent Address:

10. Date of Birth: _____ Age as on : _____

DD/MM/YYYY		

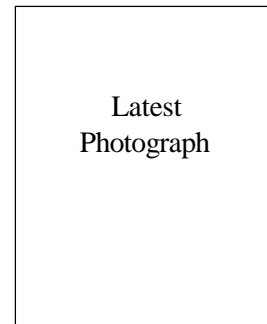
11. Whether SC/ST/OBC/General: _____ Caste: _____

12. Marital Status (Married / Unmarried): _____

13. Sex (Male/Female) : _____

14. Educational Qualifications: (Particulars of all academic examinations and degree obtained. Commencing with the Matriculation or equivalent examination)

Sr.No.	Exam. Passed	Year of Passing	Board/University	% of Marks.



15. Work Experience (in chronological order starting from the present employer)

Sr.No.	From Date	To Date	Scale of pay	Post held	Name of Employer	Nature of Duties

Total relevant experience for suitability to the post applied: _____ Years
 _____ Months

Note: Additional information, if any can be provided on a separate paper or on overleaf of this page.

16. Details of National level exams passed, if any:

Exam Passed	Date of Passing	Valid till

17. Details of research publication (indexed Journal):

Sr No.	Paper published	Title of Paper	Year	Volume & Page No.	First Author or Co-Author	Impact Factor	Citations

18. Fellowship /Awards /Membership (ICMR/CSIR/DEBT/DST/UGC/INSA/Other Academy including International)

S.N.	Name of Fellowship/ Award/ Membership	Name of the sponsoring agency	Month /Year	Amount Salary/Month	Name of the Host Institute	For what Public Health Contribution

19. If selected what period would you require joining the post: _____

DECLARATION

I hereby declare that the information furnished above is true, complete, and correct to the best of my knowledge and belief. I understand that in the event of any of the information provided by me being cancelled/terminated at any stage, my candidature/appointment shall be liable for without notice or any compensation in lieu thereof.

Date:

Name:

Place:
candidate

Signature of the

Enclosures: Self-attested copies of all certificates/ testimonial.