FORMAT FOR BIO-DATA

1. **Name:** Mr. Vinod Bhagwat Joshi

2. **Address (full work address):** Central Administrative Tribunal Gulistan Building, 4th Floor, Prescott Road, Fort, Mumbai – 400001

3. **Telephone number:** 9869362880 & 9892218108 **E-mail-ID:** joshivb@yahoo.co.in

4. **Present affiliation (Job title, department, and organization):** Senior Counsel for Union of India, Central Administrative Tribunal, Bombay High Court, Mumbai

5. **Affiliation with host institute:** No

6. **Qualification (starting from basic, add additional rows if needed):**

COURSES/SUBJECT	INSTITUTE/ORGANIZATION	YEAR
B.Com.	Nagpur University, Nagpur	1985
LL.B.	Amravati University, Amravati	1989
Doctor of Philosophy (Honoris Causa)	Commonwealth Vocational University, Makaunga, Hahake, Tongatapu, Kingdom of Tonga	2021

7. **Previous and other affiliations (add additional rows if needed):**

AFFILIATION	DESIGNATION	DURATION
Central Administrative Tribunal	Senior Counsel for Union of India	15.12.2000 – Present
High Court of Judicature, Bombay	Senior Counsel for Govt. of India	22.10.02 – 30.08.2018
Bombay High Court	Additional Public Prosecutor for Govt. of India	18.07.05 – 18.07.2009
Maharashtra Administrative Tribunal	Government Pleader : Presenting Officer	01.11.99 – 29.03.01
Government of Maharashtra	Special Counsel	2010
UPSC for High Court Bombay and Central Administrative Tribunal	Nodal Counsel	Since 2015
AG Office Maharashtra at Mumbai, Nagpur, and Aurangabad	Representing	Since 2000
National Institute of Fashion Technology, Mumbai	Representing	Since 2015
Indian Institute of Population Science	Representing	Since 2015

8. **Role in proposed Ethics Committee (also add dual Role if any):** Member

9. **Suitability of the member in the assigned role, Elaborate (less than 100 words):** Dr. Vinod Bhagwat Joshi, as a Senior Counsel, reviews the legal aspects of research, ensuring compliance with both national and international laws. His legal expertise helps verify that research protocols, respect regulations and guidelines, privacy laws, and data protection frameworks. He also ensures the proper use of informed consent, protection of participant rights, and ethical compliance, reducing legal risks and aligning the research with ethical standards required by the IRB for human subject protection.

10. **Previous EC experience: YES/NO, if yes add role/ duration with name of EC: (previous EC experience is mandatory for the Chairperson):** No

NAME OF ETHICS COMMITTEE	DESIGNATION/ ROLE	DURATION	
		FROM	TO

11. Relevant research training/experience in the area*:(add additional rows if needed):

NAME OF ETHICS COURSE/ TRAINING	ORGANIZED BY	DATE	DURATION OF TIMIMG	ATTACH AGENDA/ TOPICS COVERED

12. Relevant publications (any 5) and additional information (if any): Not applicable

Signature: 

Date: 25-09-2024

* ***Details must primarily include training in ICMR National Ethical Guidelines for Biomedical and Health Research Involving Human Participants, 2017, GCP Guidelines (if applicable), New Drugs and Clinical Trials (NDCT) Rules, 2019, EC Functions & SOPs and relevant regulations of the country.***