



सत्यमेव जयते

GOVERNMENT OF INDIA
CENTRAL DRUGS STANDARD CONTROL
ORGANISATION (Headquarter)
(Directorate General of Health Services)
Ministry of Health & Family Welfare
FDA Bhavan
ITO, Kotla Road
New Delhi - 110002 (Delhi)
Phone No.: 91-11-23216367
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File No. CT/23/000145

To,

M/s. AstraZeneca Pharma India Limited,
Block N1, 12th Floor Manyata Embassy,
Business Park, Rachenahalli Outer Ring Road,
Bangalore, Karnataka (India) – 560045.

Sir,

With reference to your application No. GCT/CT04/FF/2023/40513 (GCT/145/23) dated 16.11.2023, please find enclosed herewith the permission in Form CT-06 for conduct of clinical trial titled, **“A Phase III, Randomized, Open-Label, Multi-Center, Global Study of Volrustomig (MEDI5752) as Sequential Therapy Versus Observation in Participants with Unresected Locally Advanced Head and Neck Squamous Cell Carcinoma, Who Have Not Progressed Following Definitive Concurrent Chemoradiotherapy (eVOLVE-HNSCC)” Protocol No. D798EC00001, CSP Version 1.0 dated 08.08.2023 with a total of up-to 32 subjects from India** under the provisions of New Drugs and Clinical Trial Rules, 2019

The permission granted by the Central Licensing Authority to conduct clinical trial shall be subject to following conditions, namely:-

- (i) Clinical trial at each site shall be initiated after approval of the clinical trial protocol and other related documents by the Ethics Committee of that site, registered with the Central Licencing Authority under rule 8;
- (ii) where a clinical trial site does not have its own Ethics Committee, clinical trial at that site may be initiated after obtaining approval of the protocol from the Ethics Committee of another trial site; or an independent Ethics Committee for clinical trial constituted in accordance with the provisions of rule 7:

Provided that the approving Ethics Committee for clinical trial shall in such case be responsible for the study at the trial site or the centre, as the case may be:

Provided further that the approving Ethics Committee and the clinical trial site or the bioavailability and bioequivalence centre, as the case may be, shall be located within the same city or within a radius of 50 kms of the clinical trial site;

- (iii) in case an ethics committee of a clinical trial site rejects the approval of the protocol, the details of the same shall be submitted to the Central Licensing Authority prior to seeking approval of another Ethics Committee for the protocol for conduct of the clinical trial at the same site;

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- (iv)** the Central Licencing Authority shall be informed about the approval granted by the Ethics Committee within a period of fifteen working days of the grant of such approval;
- (v)** clinical trial shall be registered with the Clinical Trial Registry of India maintained by the Indian Council of Medical Research before enrolling the first subject for the trial;
- (vi)** clinical trial shall be conducted in accordance with the approved clinical trial protocol and other related documents and as per requirements of Good Clinical Practices Guidelines and the provisions of these rules;
- (vii)** status of enrolment of the trial subjects shall be submitted to the Central Licencing Authority on quarterly basis or as appropriate as per the duration of treatment in accordance with the approved clinical trial protocol, whichever is earlier;
- (viii)** six monthly status report of each clinical trial, as to whether it is ongoing, completed or terminated, shall be submitted to the Central Licencing Authority electronically in the SUGAM portal;
- (ix)** in case of termination of any clinical trial the detailed reasons for such termination shall be communicated to the Central Licencing Authority within thirty working days of such termination;
- (x)** any report of serious adverse event occurring during clinical trial to a subject of clinical trial, shall, after due analysis, be forwarded to the Central Licencing Authority, the chairperson of the Ethics Committee and the institute where the trial has been conducted within fourteen days of its occurrence as per Table 5 of the Third Schedule and in compliance with the procedures as specified in Chapter VI;
- (xi)** in case of injury during clinical trial to the subject of such trial, complete medical management and compensation shall be provided in accordance with Chapter VI and details of compensation provided in such cases shall be intimated to the Central Licencing Authority within thirty working days of the receipt of order issued by Central Licencing Authority in accordance with the provisions of the said Chapter;
- (xii)** in case of clinical trial related death or permanent disability of any subject of such trial during the trial, compensation shall be provided in accordance with Chapter VI and details of compensation provided in such cases shall be intimated to the Central Licencing Authority within thirty working days of receipt of the order issued by the Central Licencing Authority in accordance with the provisions of the said Chapter;
- (xiii)** the premises of the sponsor including his representatives and clinical trial sites, shall be open for inspection by officers of the Central Licencing Authority who may be accompanied by officers of the State Licencing Authority or outside experts as authorised by the Central Licencing Authority, to verify compliance of the requirements of these rules and Good Clinical Practices Guidelines, to inspect, search and seize any record, result, document, investigational product, related to clinical trial and furnish reply to query raised by the said officer in relation to clinical trial;
- (xiv)** where the new drug or investigational new drug is found to be useful in clinical development, the sponsor shall submit an application to the Central Licencing Authority for permission to import or manufacture for sale or for distribution of new drug in India, in accordance with Chapter X of these rules, unless otherwise justified;
- (xv)** the laboratory owned by any person or a company or any other legal entity and utilised by that person to whom permission for clinical trial has been granted used for research and development, shall be deemed to be registered with the Central Licencing Authority and may be used for test or analysis of any drug for and on behalf of Central Licencing Authority;
- (xvi)** the Central Licencing Authority may, if considered necessary, impose any other condition in writing with justification, in respect of specific clinical trials, regarding the objective, design, subject population, subject eligibility, assessment, conduct and treatment of such specific clinical trial;
- (xvii)** the sponsor and the investigator shall maintain the data integrity of the data generated during clinical trial.

- (xviii) Merely granting permission to conduct the clinical trial with the Investigational Drug Product does not convey or imply that, based on the clinical trial study data generated with the investigational drug, permission to market this drug in the country will automatically be granted to you;
- (xix) The permission to initiate clinical trial granted under rule 22 in form CT-06 or automatic approval under rule 23 in Form CT 4A shall remain valid for a period of **two years** from the date of its issue, unless extended by the Central Licencing Authority.

Yours faithfully,

(Dr. Rajeev Singh Raghuvanshi)
Drugs Controller General (India) &
Central Licencing Authority
Stamp

FORM CT-06

(See rules 22,25,26,29 and 30)

**PERMISSION TO CONDUCT CLINICAL TRIAL OF NEW DRUG OR
INVESTIGATIONAL NEW DRUG**

1. The Central Licensing Authority hereby permits **M/s. AstraZeneca Pharma India Limited, Block NO 1, 12th Floor, Manyata Embassy Business Park, Rachenahalli, Outer Ring Road, Bangalore, Karnataka (India) - 560045** to conduct clinical trial of the new drug or investigational new drug as per **Protocol No. D798EC00001, CSP Version 1.0 dated 08.08.2023** in the below mentioned clinical trial sites [As per Annexure].
2. Details of new drug or investigational new drug and clinical trial site [As per Annexure].
3. This permission is subject to the conditions prescribed in part A of Chapter V of the New Drugs and Clinical Trials Rules, 2019 under the Drugs and Cosmetics Act, 1940.

Place: New Delhi

Date _____

(Dr. Rajeev Singh Raghuvanshi)
Drugs Controller General (India) &
Central Licencing Authority
Stamp

Note: The permission to initiate clinical trial granted under rule 22 in form CT-06 shall remain valid for a period of **two years** from the date of its issue, unless extended by the Central Licencing Authority.

Annexure:

Details of new drug or investigational new drug:

Names of the new drug or investigational new drug	Volrustomig (MEDI5752)
Therapeutic class:	Biologic Human IgG1 mAb
Dosage form:	Lyophilized product For concentrate for solution for infusion
Composition:	Volrustomig (MEDI5752)=250.00 milligram (mg); In House Specification; Active; L-Histidine =6.6 milligram (mg) U.S.P,J.P.,Ph.Eur; Inactive; L-Histidine hydrochloride monohydrate =12.10 milligram (mg) J.P.,Ph.Eur; Inactive; Sucrose =410.80 milligram (mg)

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	J.P.,Ph.Eur, NF; Inactive; Polysorbate 80 =1.00 milligram (mg) J.P.,Ph.Eur, NF; Inactive. Water for injection = q.s. U.S.P.,Ph.Eur; Inactive
Indications:	Unresected locally advanced head and neck squamous cell carcinoma (LA - HNSCC) after receiving definitive concurrent chemoradiotherapy (cCRT)

Annexure:

Details of clinical trial site:

Sr. No.	Names and address of clinical trial site	Ethics committee details	Name of investigator
1.	HCG - Helath Care Global Enterprises Ltd, HCG Tower, No. 8, P. Kalinga Rao Road, Sampangi Rama Nagar, Bangalore - 560027, Karnataka	HCG Central Ethics Committee, HCG Towers, #8, P.kalinga Rao Road, Sampangi Ram Nagar, Bangalore -560027 ECR/386/Inst/KA/2013/RR-19	Dr Srinivasa B J
2.	Meenakshi Mission Hospital and Research Centre, Lake Area, Melur Road, Madurai - 625107, Tamil Nadu	Institutional Ethics Committee, Room No 6701/6th Floor, Meenakshi Mission Hospital & Research Centre, Lake Area, Melur Road, Madurai - 625107, Tamil Nadu ECR/398/Inst/TN/2013/RR-19	Dr Kirushna Kumar Kosanam Subramanian
3.	Sri Ram cancer & Superspeciality Centre, Mahatma Gandhi Medical College & Hospital, Ricco Industrial Area, Sitapura, Tonk Road, Jaipur - 302022, Rajasthan	Institutional Ethics Committee, Mahatma Gandhi Medical College & Hospital, RIICO Institutional Area, Sitapura, Jaipur- 302022, Rajasthan ECR/125/Inst/RJ/2013/RR-19	Dr Lalit Mohan Sharma
4.	Kailash Cancer Hospital & Research Center, Muni Seva Ashram Goraj, Waghodia - 391760, Vadodara, Gujarat	IEC-Kailash Cancer Hospital & Research Center, Muni Seva Ashram Goraj, Waghodia - 391760, Vadodara, Gujarat ECR/49/Inst/GJ/2013/RR	Dr. Santhosh Van danasetti

		-19	
5.	Mahamana Pandit Madan Mohan Malaviya Cancer Centre, Sundar Bagiya colony, Sundarpur, Near Nariya Gate, Banaras Hindu University Campus, Varanasi - 221005, Uttar Pradesh	IEC - MPMMCC and HBCH Varanasi, Mahamana Pandit Madan Mohan Malviya Cancer Centre, Sundar Bagiya , Near Nariya Gate, Banaras Hindu University, Campus, Varanasi - 221005, Uttar Pradesh ECR/1501/Inst/UP/2021	Dr Akhil Kapoor
6.	NH - Rabindranath Tagore International Institute of Cardiac Sciences, 1489 (124) Eastern Metropolitan Bypass Premises No: 1489, Mukundapur, Kolkata - 700099, West Bengal	NHRTIICS Ethics Committee 124, Mukundpur, E.M, Bypass Kolkata -700099, West Bengal ECR/316/Inst/WB/2013/RR -19	Dr Chandrakanth MV
7.	Care Hospitals, Old Mumbai Highway, Near Cyberabad Police Commissionerate, Jayabheri Pine Valley, HITEC City, Hyderabad, Telangana - 500032	Care Hospitals, Institutional Ethics Committee, Care Hospital, 6 -3- 248/2, Care Hospital, In -Patient building, 4th Floor Room No. 401, Road No - 1, Banjara Hills, Hyderabad, Telangana 500034 ECR/94/Inst/AP/2013/RR -21	Dr Deepak Kopakka
8.	Aster Medcity, Kuttisahib Road Cheranelloor, South Chittoor, Kochi - 682027, Kerala	Institutional Ethics Committee, Aster DM Health Care Limited, Aster Medcity, Kuttisahib Road Cheranelloor, South Chittoor, Kochi - 682027, Kerala ECR/737/Inst/KL/2015/RR -21	Dr Arun R Warriar
9.	Regional Cancer Centre, Medical	Human Ethics Committee, Regional Cancer	Dr Rejnish Kumar R

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	College Campus, Thiruvananthapuram - 695011, Kerala	Centre, Medical College Campus, Thiruvananthapuram - 695011, Kerala ECR/21/Inst/Ker/2013/RR -19	
10.	King Georges Medical University, Shahmina Road, Lucknow - 226003, Uttar Pradesh	Institutional Ethics Committee, King Georges Medical University, Shahmina Road, Lucknow - 226003, Uttar Pradesh ECR/262/Inst/UP/2013/RR - 19	Dr Vijay Kuma
11.	Max Super Speciality Hospital, Near Civil Hospital, Phase 6, Mohali, Punjab - 160055 India Manglore Karnataka - 575002	Institutional Ethics Committee, Max Super Speciality Hospital, Mohali, Near Civil Hospital, Chandigarh Road, Phase VI, Shahibzada Ajit Singh Nagar Kharar, Punjab Mohali SAS Nagar Punjab – 160055 ECR/691/Inst/PB/2014/RR -21	Dr Sachin Gupta
12.	Indraprastha Apollo Hospitals, Sarita Vihar, Delhi-Mathura Road, New Delhi-110076	Institutional Ethics Committee - Clinical Studies, Indraprastha Apollo Hospitals, Sarita Vihar, Delhi - Mathura Road, New Delhi - 110076, India ECR/5/Inst/DL/2013/RR -19	Dr Manish Kumar Singha
13.	Regency Hospital Ltd. Tower-2 Day Care 1st floor Regency Hospital Ltd. Sarvodaya nagar Kanpur 208005, Uttar Pradesh	Regency Hospital Ethics Committee, Regency Hospital Ltd., A -2, Sarvodaya nagar Kanpur 208005, Uttar Pradesh ECR/825/Inst /UP/2016/RR -19	Dr Vikas T Talreja

