



**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/23/000108**

To,

M/s. AstraZeneca Pharma India Limited,  
Block NO 1, 12th Floor Manyata Embassy,  
Business Park, Rachenahalli Outer Ring Road,  
Bangalore, Karnataka (India) – 560045.

Sir,

With reference to your application No. GCT/CT04/FF/2023/39490 (GCT/108/23) dated 04-09-2023, please find enclosed herewith the permission in Form CT-06 for conduct of clinical trial titled, **“A Phase III, Randomised, Double-blind, Placebo-controlled, Multi-Centre, Global study of Volrustomig in Women with High Risk Locally Advanced Cervical Cancer Who Have Not Progressed Following Platinum-based, Concurrent Chemoradiation Therapy (eVOLVE-Cervical)” Protocol No: D7984C00002: Version 2.0 dated 07/Aug/2023, CSP Protocol-Addendum IND-1, Version 1.0, dated 21/Aug/2023 with a total of up-to 70 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 Licensing 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;

- (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;

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- (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

**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, 12<sup>th</sup> Floor, Manyata Embassy Business Park, Rachenahalli, Outer Ring Bangalore (India) - 560045** to conduct clinical trial of the new drug or investigational new drug as per **Protocol No: D7984C00002: Version 2.0 dated 07/Aug/2023, CSP Protocol-Addendum IND-1, Version 1.0, dated 21/Aug/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:

|                                                          |                                                                                                                                                                                                                                                                                                                                                                                                                               |
|----------------------------------------------------------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| <b>Names of the new drug or investigational new drug</b> | Volrustomig( MEDI5752) Lyophilized Powder for solution for infusion 250 mg/vial                                                                                                                                                                                                                                                                                                                                               |
| <b>Therapeutic class:</b>                                | Human IgG1 monoclonal antibody                                                                                                                                                                                                                                                                                                                                                                                                |
| <b>Dosage form:</b>                                      | Lyophilized Powder for solution for infusion                                                                                                                                                                                                                                                                                                                                                                                  |
| <b>Composition:</b>                                      | Volrustomig (MEDI5752)=250.0000 milligram (mg) In House Specification Active<br>L-Histidine = 6.6000 milligram (mg) U.S.P,J.P, Ph. Eur Inactive<br>L-Histidine hydrochloride monohydrate =12.1000 milligram (mg) J.P, Ph.Eur Inactive<br>Sucrose = 410.8000 milligram (mg) J.P, Ph. Eur Inactive<br>Polysorbate 80 = 1.0000 milligram (mg) J.P, Ph. Eur Inactive<br>Water for injection = 1.0000 q.s. U.S.P, Ph. Eur Inactive |

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| <b>Indications:</b> | Women with high risk locally advanced cervical cancer (LACC) who have not progressed following platinum-based, Concurrent Chemoradiation Therapy (CCRT) |
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**Annexure:**

Details of clinical trial site:

| <b>Sr. No.</b> | <b>Names and address of clinical trial site</b>                                                                            | <b>Ethics committee details</b>                                                                                                                                                         | <b>Name of investigator</b>            |
|----------------|----------------------------------------------------------------------------------------------------------------------------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------------------------|
| <b>1.</b>      | HCG Manavata Cancer Centre, Behind Shivang Auto, Mumbai Naka, Nashik, 422002, Maharashtra, India                           | Manavata Clinical Research Institute Ethics Committee, HCG Manavata Cancer Centre, Behind Shivang Auto, Mumbai Naka, Nashik, 422002, Maharashtra<br><br>ECR/500/Inst/MH/2013/RR-20      | Dr. Shruti Deepak Kate                 |
| <b>2.</b>      | Meenakshi Mission Hospital & Research Centre, Lake area, Melur Road, Madurai, Tamil Nadu, 625107                           | Institutional Ethics Committee Meenakshi Mission Hospital & Research Centre, Lake Area, Melur Road, Madurai, Tamil Nadu, 615107<br><br>ECR/398/Inst/TN/2013/RR-19                       | Dr. Kirushna Kumar Kosaram Subramanian |
| <b>3.</b>      | Tata Medical Center, 14, Major Arterial Road (EW), New Town Rajarhat, Kolkata, West Bengal, 700160                         | Institutional Review Board, Tata Medical Center, 14, Major Arterial Road (EW), New Town, Rajarhat, Kolkata, West Bengal –700160<br><br>ECR/269/Inst/WB/2013/RR-19                       | Dr. Joydeep Ghosh                      |
| <b>4.</b>      | Apex Wellness Hospital, Survey no. 799, plot no. 187, Behind Praksh Petrol Pump, Govind Nagar, Nashik, Maharashtra, 422009 | Apex Wellness Ethics Committee, Apex Wellness Hospital, Survey no. 799, plot no. 187, Behind Praksh Petrol Pump, Govind Nagar, Nashik, Maharashtra, 422009<br><br>ECR/1500/Inst/MH/2021 | Dr. Shailesh Bondarde                  |
| <b>5.</b>      | MVR Cancer Centre & Research Institute, CP 13/516, B.C. Vellalasseri REC (via) Poolacode, Kozhikode, Kerala, 673601, India | Institutional Ethics Committee, MVR Cancer Centre & Research Institute, CP 13/516, B.C. Vellalasseri, REC (via) Poolacode, Kozhikode, Kerala, 673601<br><br>ECR/1259/Inst/KL/2019       | Dr. Narayanankutty Warriar             |

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| <b>6.</b>  | KIMS -Kingsway Hospitals, 44, Parwana Bhavan, Kingsway Road, Nagpur, Maharashtra, 440001                                                                                            | Kingsway Hospitals Ethics Committee, Kingsway Hospital, 44, Parwana Bhavan, Kingsway Road, Nagpur, Maharashtra, 440001<br><br>ECR/1269/Inst/MH/2019                                                               | Dr. Saurabh Prasad       |
| <b>7.</b>  | Kailash Cancer Hospital & Research Centre, Muni Seva Ashram, Goraj, Waghodia, Vadodara, 391760, Gujarat, India                                                                      | IEC -KCHRC, Kailash Cancer Hospital and Research Centre, First Floor, Department of Clinical Research, Goraj, Waghodia, Vadodara, Gujarat, 391760<br><br>ECR/49/Inst/GJ/2013/RR-19                                | Dr. Santosh Vandanasetti |
| <b>8.</b>  | Department of Medical Oncology , Room No. 160A, All India Institute of Medical Sciences (AIIMS), Dr. B.R Ambedkar Institute Rotary Cancer Hospital, Ansari Nagar, New Delhi, 110029 | Institute Ethics Committee, All India Institute of Medical Science, Room No. 102 First Floor, Old T Block, Ansari Nagar, New Delhi, 110029<br><br>ECR/538/Inst/2014/RR-20                                         | Dr. Sachin Khurana       |
| <b>9.</b>  | Clinical Trial & Research Department, Bhagwan Mahaveer Cancer Hospital and Research Center, Jawahar Lal Nehru Marg, Jaipur, Rajasthan, 302017, India                                | Institutional Ethics Committee, Bhagwan Mahaveer Cancer Hospital and Research Centre, Jawahar Lal Nehru Marg, Jaipur, Rajasthan, 302017(Raj), India<br><br>ECR/19/Inst/RJ/2013/RR-20                              | Dr. Deepak Gupta         |
| <b>10.</b> | Max Super Speciality Hospital, Sector 56, Near Civil Hospital, Phase 6, Mohali, Punjab, 160055, India                                                                               | Institutional Ethics Committee, Max Super Speciality Hospital, Near Civil Hospital, Chandigarh Road, Phase 6, Shahibzada Ajit Singh Nagar Kharar, Mohali, Punjab, 160055, India<br><br>ECR/691/Inst/PB/2014/RR-21 | Dr. Gautam Goyal         |
| <b>11.</b> | King Georges Medical University, Department of surgical oncology, 3rd floor, Shatabdi Phase 2, King Georges Medical University, Lucknow, Uttar Pradesh, 226003, India               | Institutional ethics committee, King Georges Medical University, Shahmina road, Lucknow, 226003, Uttar Pradesh, India<br><br>ECR/262/Inst/UP/2013/RR-19                                                           | Dr. Vijay Kumar          |

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