



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
Fax No.: 91-11-23236973
E-Mail : dci@nic.in

File No. CT/20/000048

To,

M/s. Astrazeneca Pharma India Limited,
Block NO1, 12th Floor Manyata Embassy,
Business Park, Rachenahalli Outer Ring,
Bangalore (India) – 560045.

Sir,

With reference to your application No GCT/CT04/FF/2020/19950 (GCT/50/20) dated 29-05-2020, please find enclosed herewith the permission in Form CT-06 for conduct of clinical trial titled, **“A Phase III Double-Blind, Randomised, Placebo-Controlled Study Assessing the Efficacy and Safety of Capivasertib + Abiraterone Versus Placebo + Abiraterone as Treatment for Patients with De Novo Metastatic Hormone-Sensitive Prostate Cancer (mHSPC) Characterised by PTEN deficiency (CAPItello-281)”**, **Protocol Number: D361BC00001 Version No. 1.0 dated 03/March/2020** 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 Licensing 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;

File No. CT/50/20-DCG(I)

- (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) 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. V. G. Somani)
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 NO1, 12th 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 Number: D361BC00001 Version No. 1.0 dated 03/March/2020** 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 ct, 1940.

Place: New Delhi

Date _____

(Dr. V. G. Somani)
Drugs Controller General (India)
Central Licensing 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	Capivasertib + Abiraterone
Therapeutic class:	Anticancer
Dosage form:	Tablets
Composition:	Capivasertib = 160.0000 milligram (mg) In House Specification Active Capivasertib = 200.0000 milligram (mg) In House Specification Active Abiraterone = 500.0000 milligram (mg) In House Specification Active
Indications:	Treatment for Patients with denovo Metastatic Hormone Sensitive Prostate Cancer (mHSPC) Characterised by PTEN Deficiency

File No. CT/50/20-DCG(I)

Details of clinical trial site:

Names and address of clinical trial site	Ethics committee details	Name of investigator
Department of Medical Oncology, VMMC & Safdarjung Hospital, New Delhi-110029	Institutional Ethics Committee, VMMC & Safdarjung Hospital, New Delhi-110029 ECR/593/ Inst/DL/2014/RR-17	Dr. S P Kataria
Rajiv Gandhi Cancer Institute and Research Centre, Sector-5, Rohini, New Delhi-110085	Institutional Review Board, Rajiv Gandhi Cancer Institute and Research Centre, Sector-5, Rohini, New Delhi-110085 ECR/10/Inst/DC/2013/RR-19	Dr. Sudhir Kumar Rawal
Shetty's Hospital, Plot No. 11 & 12, 12 th F Main Road, Kaveri Nagar, Kodichikkanahalli, Bommanahalli, Bengaluru-560068	Shetty's Hospital Ethics Committee, Shetty's Hospital, Plot No. 11 & 12, 12 th F Main Road, Kaveri Nagar, Kodichikkanahalli, Bommanahalli, Bengaluru-560068, Karnataka, India ECR/918/Inst/KA/2017	Dr. Lokesh K. N
Dr. B L Kapur Memorial Hospital, Pusa Road, New Delhi-110005	Dr. B L Kapur Memorial Hospital Ethics Committee, AARCE, Dr. B L Kapur Memorial Hospital, Pusa Road, New Delhi-110005 ECR/3/BLK/Inst/DL/2013/RR-19	Dr. Amit Agarwal
Asian Institute of Medical Sciences, Badkal Flyover Road, Sec-21A, Faridabad-121001, Haryana, India	Institutional Ethics Committee, Asian Institute of Medical Sciences, Badkal Flyover Road, Sec-21A, Faridabad-121001, Haryana, India ECR/626/Inst/HR/2014/RR-17	Dr. Prashant Mehta
Max Super Specialty Hospital (A Unit of Hometrail Builtech Pvt. Ltd), Near Civil Hospital, Phase-6, Mohali, Punjab-160055, India	Institutional Ethics Committee, Max Super Specialty Hospital (A Unit of Hometrail Builtech Pvt. Ltd), Near Civil Hospital, Phase-6, Mohali, Punjab-160055, India ECR/691/Inst/PB/2014/RR-17	Dr. Gautam Goyal
Valentis Cancer Hospital, Mussoorie, Mawana Road, Meerut-250001, Uttar Pradesh, India	Institutional Ethics Committee, Valentis Cancer Hospital, Mussoorie, Mawana Road, Meerut-250001, Uttar Pradesh, India ECR/1165/Inst/UP/2018	Dr. Amit Jain
Department of Radiotherapy & Oncology, Regional Cancer Centre, PGIMER (Post Graduate Institute of Medical Education and Research), Chandigarh, Sector 12, Pin Code-	Institutional Ethics Committee, Room no 6006, 6th Floor. P. N Chuttani Block PGIMER (Post Graduate Institute of Medical Education and Research), Chandigarh, Sector 12 Pin Code 160012, INDIA ECR/23/Inst/CH/2013/RR-20	Dr. Chandan Krushna Das

File No. CT/50/20-DCG(I)

160012, India		
Amrita Institute of Medical Sciences, AIMS Ponekkara P O, Kochi-682041 Kerala, India	Institutional Ethics Committee, Amrita Institute of Medical Sciences, AIMS Ponekkara P O, Kochi - 682041 Kerala, India ECR/129/Inst/KL/2013/RR-19	Dr Arun Philip
Meenakshi Mission Hospital & Research Centre, Lake Area, Melur Road, Madurai-625107	Institutional Ethics Committee, Room No.6701, 6 th Floor, Meenakshi Mission Hospital & Research Centre, Lake Area, Melur Road, Madurai ECR/398/Inst/TN/2013/RR-19	Dr. Kirushna Kumar Kosanam Subramanian
Regional Cancer Centre, Medical College Campus, Thiruvananthapuram-695011, Kerala, India	Human Ethics Committee, First Floor, Regional Cancer Centre, Medical College Campus, Thiruvananthapuram-695011, Kerala, India ECR/21/Inst/Ker/2013/RR-19	Dr. Francis V James
Department of Medical Oncology, Dr. B.R.A.I.R.C.H, All India Institute of Medical Sciences, Ansari Nagar, East New Delhi-110029	Ethics Committee All India Institute of Medical Sciences, Ansari Nagar, Room No. 102, First Floor, Old, O.T. Block, Ansari Nagar East, New Delhi-110029 ECR/538/Inst/DL/2014/RR-17	Dr. Raja Pramanik
Kasturba Medical College and Hospital, Madhav Nagar, Manipal-576104	MAHE Ethics committee, Mezzanine floor of Old KMC Library Building, MAHE, Manipal-576104 ECR/191/Inst/KL/2013/RR-19	Dr Karthik Udupa