



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/24/000150**

To,

M/s Johnson and Johnson Private Limited,  
HIGI House, L. B. S. Marg, Mulund (West),  
Maharashtra (India) - 400080

Sir,

With reference to your application No. GCT/CT04/FF/2024/46840 dated 14-Dec-2024, please find enclosed herewith the permission in Form CT-06 for conduct of clinical trial titled, **“A Phase 3, Multicenter, Randomized, Double-blind, Placebo-controlled Study Evaluating the Efficacy and Safety of JNJ-77242113 for the Treatment of Biologic-naïve Participants with Active Psoriatic Arthritis” Protocol no: 77242113PSA3001 Amendment 1 dated 18 Nov 2024 with a total of up-to 72 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:-

- 1) **Human biological samples i.e. Human blood (whole) and serum related to clinical trial shall be permitted to be exported for analysis subject to the clearance by port offices of CDSCO. Further, ICMR/DHR permission shall be obtained for export of optional biological samples as per DGFT notification number 72/2023 dated 11.03.2024.**
- 2) 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;
- 3) 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;
- 4) 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;

- 5) 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;
- 6) 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;
- 7) 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;
- 8) 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;
- 9) 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;
- 10) 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;
- 11) 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;
- 12) 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;
- 13) 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;
- 14) 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;
- 15) 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;
- 16) 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 Licensing Authority and may be used for test or analysis of any drug for and on behalf of Central Licensing Authority;
- 17) 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;
- 18) the sponsor and the investigator shall maintain the data integrity of the data generated during clinical trial.
- 19) 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;

- 20)** 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 Johnson and Johnson Private Limited, HIGI House, L. B. S. Marg, Mulund (West), Maharashtra (India) - 400080** to conduct clinical trial of the new drug or investigational new drug as per **Protocol no: 77242113PSA3001 Amendment 1 dated 18 Nov 2024** 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>	JNJ-77242113
<b>Therapeutic class:</b>	IL 23 inhibitor peptide
<b>Dosage form:</b>	Tablets
<b>Composition:</b>	JNJ-77242113-AAC =200.0000 milligram (mg) In House Specification Active
<b>Indications:</b>	Biologic naive Participants with Active Psoriatic Arthritis

**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>
1	SIDS Hospital and Research Centre A unit of SIDS Health Care Private Limited, Off Ring Road, Near Shell Petrol Pump, Ring Road Sosyo Circle Lane, Surat Gujarat - 395002	Surat Institute of digestive sciences EC, SIDS Hospital and Research Centre, Off Ring Road, Near Shell Petrol Pump, Ring Road Sosyo Circle Lane, Surat Gujarat - 395002  ECR/813/Inst/GJ/2016/RR-19	Dr. Alpana Alpesh kumar Parmar
2	Apollo Hospitals Enterprise Limited, Plot No-251, Sainik School Road, Unit-15, Bhubaneswar Orissa - 751005	Institutional Ethics Committee-Clinical Studies Apollo Hospital Bhubaneswar, Plot No 251, Sainik School Road, Unit 15, Odisha, Bhubaneswar- 751005  ECR/246/INST/OR/2013/RR- 19	Dr. Ajit Kumar Surin
3	Indraprastha Apollo Hospital, Sarita Vihar, Delhi Mathura Road New delhi Delhi -110076	Institutional Ethics Committee-Clinical Studies Hostel basement complex, Indraprastha Apollo hospital, Sarita Vihar, Delhi Mathura Road, Delhi-110076  ECR/5/Inst/DL/2013/RR-19	Dr. Sundeep Kumar Upadhyaya
4	Kokilaben Dhirubhai Ambani Hospital and Medical Research Institute,, Rao Saheb, Rao Saheb Achutrao Patwardhan Marg, Four Bungalows, Andheri West Mumbai Maharashtra - 400053	INSTITUTIONAL ETHICS COMMITTEE Second floor, Kokilabaen Dhirubhai ambani hospital and medical research institute, Rao Saheb Achutrao patwardhan marg, Four bungalows, Andheri West, Mumbai, 400053  ECR/141/Inst/MH/2013/RR-24	Dr. Jyotsna Oak

5	Nizams Institute of Medical Sciences, Punjagutta Rd, Punjagutta Hyderabad Telangana - 500082	NIMS Institutional Ethics Committee Nizams Institute of Medical Sciences, Punjagutta Hyderabad Telangana - 500082 India  ECR/303/Inst/AP/2013/RR-19	Dr. Amirtha Gopalan
6	Fortis Hospital Noida, B-22, Rasoolpur Nawada, D Block, Sector 62 ,Noida Uttar Pradesh-201301	FORTIS HOSPITAL INSTITUTIONAL ETHICS COMMITTEE Room no –1341, IPD 3rd Floor, Fortis Hospital, B-22, Sector 62, Noida, Uttar Pradesh  ECR/69/Inst/UP/2013/RR-20	Dr. Bimlesh Dhar Pandey
7	All India Institute of Medical Sciences, Sri Aurobindo Marg, Ansari Nagar, New Delhi Delhi- 110029	Institute Ethics Committee All India Institute of Medical Sciences Old OT Block, Room No. 102, AIIMS Hospital Ansari Nagar, New Delhi-29 Delhi – 110029  ECR/538/Inst/DL/2014/RR-20	Dr. Uma Kumar
8	JSS Hospital, Department Clinical Immunology and Rheumatology, Mahatma Gandhi Road, Fort Mohalla Mysuru Karnataka - 570004	Institutional Ethics Committee, JSS Medical College, Sri Shivarathreeshwara Nagara, Mysuru Mysore Karnataka - 570015 India  ECR/387/Inst/KA/2013/RR-22	Dr. Shiva Prashad BN
9	Apollo Multispeciality Hospitals Ltd, 58, Canal Circular Rd, Kadapara, Phool Bagan, Kankurgachi, Kolkata, West Bengal -700054	Institutional Ethics Committee Apollo Multispeciality Hospitals Ltd, 58 Canal Circular Road kolkata 700054 West Bengal.  ECR/373/Inst/WB/2013/RR-19	Dr. Syamasis Bandyopadhyay
10	Government Medical College, Kozhikode, Medical college PO Kozhikode Kerala - 673008	Institutional Ethics Committee Govt Medical College Kozhikode 4th Floor, Golden Jubilee Annex Institute of Maternal and Child Health Kozhikode, Kerala – 673008  ECR/395/Inst/KL/2013/RR-20	Dr. Neeraj Manikath

\*\*\*\*\*