



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/20/000074

To,

M/s. Novo Nordisk India Pvt. Ltd.,
Plot No. 32, 47-50, EPIP Area, Whitefield,
Bangalore, Karnataka (India) – 560066.

Sir,

With reference to your application No GCT/CT04/FF/2020/20993 (GCT/74/20) dated 11-08-2020, please find enclosed herewith the permission in Form CT-06 for conduct of clinical trial titled, **“A 78-week trial comparing the effect and safety of once weekly insulin icodec and once daily insulin glargine 100 units/mL, both in combination with noninsulin anti-diabetic treatment, in insulin naïve subjects with type 2 diabetes.”, Protocol No NN1436-4477 Version No. 1.0 dated 19/June/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) **The cutoff value for “No reduction in HbA1c” should be objectively defined and to provide guidelines to investigators with regard to safety of patients.**
- (ii) 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;
- (iii) 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;
- (iv) 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;
- (v) 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;
- (vi) 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;
- (vii) 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;

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- (viii) 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;
- (ix) 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;
- (x) 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;
- (xi) 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;
- (xii) 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;
- (xiii) 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;
- (xiv) 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;
- (xv) 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;
- (xvi) 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;
- (xvii) 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;
- (xviii) the sponsor and the investigator shall maintain the data integrity of the data generated during clinical trial.
- (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. 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. Novo Nordisk India Pvt. Ltd., Plot No. 32, 47-50, EPIP Area, Whitefield, Bangalore, Karnataka (India) – 560066** to conduct clinical trial of the new drug or investigational new drug as per **Protocol No NN1436-4477 Version No. 1.0 dated 19/June/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 Act, 1940.

Place: New Delhi

Date _____

(Dr. V. G. Somani)
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	Insulin icodec
Therapeutic class:	Antidiabetic Drug
Dosage form:	Injection
Composition:	Water for injection = 1.0000 ml U.S.P.,E.P.,J.P. Inactive Hydrochloric acid = 0.0000 q.s. U.S.P.,E.P.,J.P. Inactive Sodium hydroxide = 0.0000 q.s. U.S.P.,E.P.,J.P. Inactive Sodium chloride = 1.1700 mg/ml U.S.P.,E.P.,J.P. Inactive Metacresol = 1.0800 mg/ml U.S.P.,E.P. Inactive Phenol = 5.6500 mg/ml U.S.P.,E.P.,J.P. Inactive Glycerol = 15.0000 mg/ml U.S.P.,E.P.,J.P. Inactive Insulin icodec = 700.0000 IU/ml In House Specification Active Zinc (as zinc acetate) = 101.0000 µg/ml I.P.,U.S.P.,J.P. Inactive
Indications:	Type 2 diabetes

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Details of clinical trial site:

Sr. No.	Name and address of clinical trial site	Ethics committee details	Name of Investigator
1.	TOTALL Diabetes Hormone Institute, A Unit of Diabetes Thyroid Hormone Research Institute Pvt. Ltd., BCM Health Island, PU 4, Scheme 54, Behind Prestige Management Institute, Near Bombay Hospital, Indore-452010 (Madhya Pradesh), India	Ethics Committee of Diabetes Thyroid Hormone Research Institute, BCM Health Island, PU 4, Scheme 54, Behind Prestige Management Institute, Near Bombay Hospital, Indore-452010 (Madhya Pradesh), India ECR/409/Inst/MP/2013/RR-19	Dr. Sunil M. Jain
2.	Madras Diabetes Research Foundation, No. 4, Conran Smith Road, Gopalapuram, Chennai, Tamil Nadu	Institutional Ethics Committee of Madras Diabetes Research Foundation, No. 4, Conran Smith Road, Gopalapuram, Chennai, Tamil Nadu ECR/194/Inst/TN/2013/RR-19	Dr. V. Mohan
3.	810, Dept. Of Endocrinology Diabetes and Metabolism, Christian medical College, Vellore-632004	Ethics Committee-Silver, Carman Block, Christian Medical College, Bagayam, Vellore-632002, TamilNadu, India ECR/326/Inst/TN/2013/RR-19	Dr. Nihal Thomas
4.	Department of Endocrinology, 4 th Floor, Room No. 419, College Building, T.N., Medical College and BYL Nair Ch. Hospital, Dr. A.L. Nair Road, Mumbai - 400008	Institutional Ethics Committee, 'G' building, Ground Floor, B Y L Nair Ch. Hospital, Dr. A.L. Nair Road, Mumbai - 400008 ECR/22/Inst/Maha/2013/RR-19	Dr. Nikhil Madhusudan Bhagwat
5.	Ward Block, 6 th Floor, AB6 ward seminar room, Department of Endocrinology & Metabolism, AIIMS, Ansari Nagar, New Delhi-110029	Institute Ethics Committee, Room No. 102, 1 st Floor, Old O.T. Block, AIIMS, Ansari Nagar, New Delhi-110029 ECR/538/Inst/DL/2014/RR-17	Dr. Nikhil Tondon
6.	Kumudini Devi Diabetes Research Center, Ramdevrao Hospital (A Unit of Sivananda Rehabilitation Home), Kukatpally, Hyderabad-500072, Telangana, India	Institutional Ethics Committee, Ramdevrao Hospital (A Unit of Sivananda Rehabilitation Home), Kukatpally, Hyderabad-500072, Telangana, India ECR/709/Inst/TG/2015/RR-18	Dr. Paturi Vishnupriya Rao

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7.	Department of Endocrinology, Pt. B.D. Sharma Post Graduate Institute of Medical Science, Rohtak, Haryana-124001	Institutional Ethics Committee, Pt. B.D. Sharma Post Graduate Institute of Medical Science, Rohtak, Haryana-124001 ECR/293/Inst/HR/2013/RR-19	Dr. Rajesh Rajput
8.	Department of Endocrinology, 2 nd Floor, OP Building, Golden Jubilee Block, Osmania General Hospital, Osmania Medical College, Afzalgunj, Hyderabad, Telangana-500012, India	Institutional Ethics Committee, Osmania Medical College, Koti, Hyderabad, Telangana-500095, India ECR/300/Inst/AP/2013/R-19	Dr. Rakesh Kumar Sahay
9.	Department of Endocrinology, Seth G.S. Medical College and KEM Hospital, Parel, Mumbai-400012	Institutional Ethics Committee-I, Seth G.S. Medical College and KEM Hospital, Parel, Mumbai-400012 ECR/229/Inst/2013/RR-19	Dr. Bandgar Tushar Ramkrishna
10.	Department of Endocrinology, 3 rd Floor, Main Building, Gandhi Medical College/Gandhi Hospital-Secunderabad-500003	Institutional Ethics Committee, Principal Office, Gandhi Medical College, Secunderabad-500003 ECR/180/Inst/AP/2013/RR-19	Dr. D. Vijay Sheker Reddy,
11.	Department of Endocrinology, IPGME&R & SSKM Hospital, Ronald Ross Building, 4 th Floor, Room 8, 244, A.J.C Bose Road, Kolkata-700020	IPGME&R Research Oversight Committee, Institute of Post Graduate Medical Education & Research, Office of the Dean, College Building, 5 th Floor, 244, A.J.C Bose Road, Kolkata-700020 ECR/35/Inst/WB/2013/RR-19	Dr. Subhankar Chowdhury
12.	Chellaram Diabetes Institute, "Lalani Quantum", Pune - Bangalore National Highway no. 4, opp. Calsoft Building, Bavdhan (Budruk), Pune, Maharashtra - 411021	Chellaram Diabetes Institute- Institutional Ethics Committee, "Lalani Quantum", Pune- Bangalore National Highway no. 4, opp. Calsoft Building, Bavdhan (Budruk), Pune, Maharashtra - 411021	Dr. Umnikrishnan A.G.