File No. CT/128/22-DCG(I)

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

सत्यमेव जयते

File No. CT/22/000128

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/2022/34468 (GCT/128/22) dated 19-10-2022, please find enclosed herewith the permission in Form CT-06 for conduct of Phase 3a clinical trial titled, "**Efficacy and safety of cagrilintide s.c. 2.4 mg in combination with semaglutide s.c. 2.4 mg (CagriSema s.c. 2.4 mg/2.4 mg) once-weekly in participants with overweight or obesity and type 2 diabetes", Protocol Number: NN9838-4609, Protocol Version 5.0 dated 19-August-2022 with a total of up-to 120 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) 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;

2) 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;

3) 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;

4) 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;

5) 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;

6) 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;

7) 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;

8) 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;

9) 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;

10) 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;

11) 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;

12) 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;

13) 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;

14) 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;

15) 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;

16) 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;

17) the sponsor and the investigator shall maintain the data integrity of the data generated during clinical trial.

18) 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;

19) 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,

RAJEEV SINGH RAGHUVANSHI

Digitally signed by RAJEEV SINGH RAGHUVANSHI
DN: c=IN, o=CENTRAL DRUGS STANDARD CONTROL ORGANIZATION, ou=RAJEEV SINGH RAGHUVANSHI, 2.5.4.20=80c62f6a23e4eafbe8a239774cdeb03c2769041015a06564fe67f54d765db1cb, postalCode=600034, st=TAMIL NADU, serialNumber=657F5E47D94C985D8F03BDC902D0E1FE73CFA12A1A126EA94FA5701124A19013, cn=RAJEEV SINGH RAGHUVANSHI
Date: 2023.03.31 11:00:37 +05'30'

(Dr. Rajeev Singh Raghuvanshi)
Drugs Controller General (India) &
Central Licencing Authority
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File No. CT/128/22-DCG(I)

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 (India) - 560066** to conduct clinical trial of the new drug or investigational new drug as per **Protocol Number: NN9838-4609, Protocol Version 5.0 dated 19-August-2022** 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

RAJEEV SINGH RAGHUVANSHI

Digitally signed by RAJEEV SINGH RAGHUVANSHI
Date: 2023.03.31 11:00:46 +05'30'

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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	CagriSema (0.25 mg/0.25 mg) / placebo CagriSema (0.5 mg/0.5 mg) / placebo CagriSema (1.0 mg/1.0 mg) / placebo CagriSema (1.7 mg/1.7 mg) / placebo CagriSema (0.25 mg/0.25 mg) / placebo
Therapeutic class:	Anti-obesity and Antidiabetic
Dosage form:	Solution for injection
Composition:	**Active substance:** Cagrilintide **Excipients:** L-Glutamic Acid Glycerol Hydrochloric Acid Sodium Hydroxide WFI
Indications:	Overweight or Obesity and Type 2 diabetes

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Annexure:

Details of clinical trial site:

Sr. No.	Names and address of clinical trial site	Ethics committee details	Name of investigator
1.	6-1-1070 1 to 4 Gleneagles Global Hospitals, Lakdikapool, Hyderabad-500004, Telangana, India	Institutional Ethics Committee Gleneagles Global Hospitals 6-1-1070 1 to 4, Lakdikapool, Hyderabad-500004, Telangana, India ECR/158/Inst/AP/2013/RR-19	Dr. Lakshmi Lavanya Alapati
2.	Room No. 108, department of Medicine, Academic Block, First Floor, Lady Hardinge Medical College, C-604, Shaheed Bhagat Singh Road, DIZ Area, Connaught Place, New Delhi- 110001	Institutional Ethics Committee Lady Hardinge Medical College & Associated Hospital, Room No. 625, Academic Block, Floor, New Delhi 110001 ECR/435/Inst/DL/2013/RR-20	Dr. Anupam Prakash
3.	Noble Hospital Pvt. Ltd., 153 A, Magarpatta city road, Hadapsar, Pune -411013, Maharashtra, India.	Noble Hospital Institutional Ethics Committee, Noble hospital Pvt. Ltd. Room No. 5, Clinical Research Department, Noble Annex 153 A, Magarpatta city Road, Hadapsar, Pune 411013, Maharashtra India ECR/259/Inst/MH/2013/RR-19	Dr. Reema Kashiva
4.	Indian Institute of Diabetes, Pulayanarkotta, Thiruvananthapuram-695301, Kerala, India	Institutional Ethics Committee Indian Institute of Diabetes, Pulayanarkotta, Thiruvananthapuram-695301, Kerala, India ECR/688/Inst/KL/2014/RR-20	Dr. P.K Jabbar
5.	Apollo Multispecialty Hospitals Limited, 58, Canal Circular Road Kolkata-700054	Institutional Ethics Committee Apollo Multispecialty Hospitals Limited, 58, Canal Circular Road Kolkata-700054 ECR/373/Inst/WB/2013/RR-19	Dr. Tirthankar Chaudhury
6.	Department of Endocrinology, Seth G.S. Medical College & KEM Hospital, Parel, Mumbai — 400012	Institutional Ethics Committee-I, New UG/PG Hostel, 20 storey Hostel Building, Ground Floor, KEM Hospital Campus. Near Main boys hostel, Dr. S.S. Rao Marg, Parel, Mumbai — 400012, Maharashtra. India	Dr. Anurag Ranjan Lila

		ECR/229/Inst/MH/2013/RR-19	
7.	All India Institute Of Medical Sciences, Ansari Nagar, New Dethi-110029	Institute Ethics Committee, Room No.102, 1st Floor, Old OT Block, AIIMS, Ansari Nagar New Delhi-110029 ECR/538/Inst/DL/2014/RR-17	Dr. Yashdeep Gupta
8.	HCG Hospitals, Mithakhali, Ellisbridge, Ahmedabad-380006, Gujarat, India	HCG Multi Specialty Ethics Committee, HCG Hospitals, Mithakhali, Ellisbridge, Ahmedabad-380006, Gujarat, India ECR/92/inst/GJ/2013/RR-19	Dr. Tiven Marwah
9.	Diabetes Care n' Research Centre, 42, Lendra Park, Ramdaspeth, Nagpur, Maharashtra - 440010	Arneja's Institutional Ethics Committee, Arneja's Heart and Multispecialty Hospital, Floor,123, Ramdaspeth behind Somalwar High School Nagpur, Maharashtra 440010 ECR/726/Inst/MH/2015/RR-21	Dr. Sunil Surajprasad Gupta
10.	Excel Endocrine Centre, Diabetes Corner, 1758, E-Ward, 4th, Lane Rajarampuri, Kolhapur-416008, India	Excel Endocrine Centre Institutional Ethics Committee, Excel Endocrine Centre, Diabetes Corner, 1758, E-Ward, 4th, Lane Rajarampuri, Kolhapur-416008, India ECR/1406/Inst/MH/2020	Dr. Sharvil Suresh Gadve
11.	Government Medical College, Kozhikode, Medical College Junction 17, Mavoor Road, Near Police station, Kozhikode, 673008-Kerala, India	Institutional Ethics Committee Government Medical College, Kozhikode, Room No. 1 & 2, Ground Floor, Lecture Theatre Complex, Medical College Campus, Medical College P.O, Calicut-673008, Kerala ECR/395/Inst/KL/2013/RR-20	Dr. Chandni R
12.	Madras Diabetes Research Foundation, No: 4 Conran Smith Road, Gopalapuram, Chennai -600086	Institutional Ethics Committee of Madras Diabetes Research Foundation, No: 4 Conran Smith Road, Gopalapuram, Chennai -600086 ECR/194/Inst/TN/2013/RR-19	Dr. V. Mohan
13.	M. S. Ramaiah Medical College & Hospitals, M. S. Ramaiah Nagar, MSRIT Post, Bangalore-560054, Karnataka, India	Ethics Committee M. S. Ramaiah Medical College & Hospital, M. S. Ramaiah Nagar, MSRIT Post, Bangalore-560054, Karnataka, India	Dr. Chitra Selven

		ECR/215/Inst/KA/2013/RR-22	
14.	PGIMER, Dept. of Endocrinology, Nehru Extension Block, Ground Floor, Room number 008 Chandigarh, 160012	Institutional Ethics Committee, PGIMER, Room No. 6006, IEC Office, 6th Floor, P N Chuttani Block, Chandigarh, 160012 ECR/25/Inst/CH/2013/RR-20	Dr. Sanjay Kumar Bhadada
15.	Department of Endocrinology, Room No- 43 A Forth Floor, Dhanvantri OPD Block SMS Hospital Jaipur-302004	Ethics Committee SMS Medical College & Attached Hospitals Jaipur-302004 ECR/26/Inst/RJ/2013/RR-19	Dr. Bal Ram Sharma
16.	Apollo Hospitals, No 21 Greams Lane, Off Greams Road, Chennai, Tamil Nadu-600006, India	Institutional Ethics Committee-Clinical Studies, Apollo Hospital Enterprises Limited, No 21, Greams Lane, Off Greams Road, Chennai-600006, India ECR/37/Inst/TN/2013/RR-19	Dr. Krishna G Seshadri
17.	Department of Endocrinology, 4" Floor, Room No.419, College building, T.N. Medical College and BYL Nair Ch, Hospital, Dr. A.L. Nair Road, Mumbai - 400 008	Institutional Ethics Committee, 'G' Building, Ground Floor, B.Y.L. Nair Ch, Hospital, Dr. A.L. Nair Road, Mumbai - 400 008 ECR/22/Inst/Maha/2013/RR-19	Dr. Nikhil Madhusudan Bhagwat

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