

Dairy No.:
ND/CT21/FF/2021/29339
dated 24.12.2021

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F. No. ND/CT/21/000085
Government of India
Directorate General of Health Services
Central Drugs Standard Control Organization
New Drugs Division

FDA Bhawan, Kotla Road,
New Delhi-110002
Dated:

To

Zydus Corporate Park, Scheme No. 63,
Survey No. 536, Khoraj (Gandhinagar),
Nr. Vaishnodevi Circle, S.G. Highway,
Ahmedabad-382481, Gujarat, India

03 FEB 2022
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Subject: Grant of permission to undertake Phase-III Clinical Trial titled "A prospective, randomized, two-arm, double-blind, active controlled, parallel, multicentre, non-inferiority, phase III clinical trial to assess the efficacy and safety of Naftifine hydrochloride cream 2% w/w as compared to Terbinafine hydrochloride cream 1% w/w in patients with superficial fungal infection of the skin" - regarding.


CT NOC No.: CT/ND/04/2022

Sir,

With reference to your application no. ND/MA/21/000211 dated 24.12.2021, please find enclosed herewith the permission in Form CT-06, No. CT/ND/04/2022 to conduct the subject mentioned clinical trial under the provisions of New Drugs and Clinical Trial Rules, 2019.

This permission is subject to the conditions, as mentioned below.

Yours faithfully,


(Dr. V. G. Somani)
Central Licensing Authority

Conditions of Permission

- (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;
- (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 Licensing 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 Licensing 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 Licensing 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 Licensing 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 of the New Drugs and Clinical Trials Rules, 2019;
- (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 the Chapter VI of the said Rules and details of compensation provided in such cases shall be intimated to the Central Licensing Authority within thirty working days of the receipt of order issued by Central Licensing 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 the Chapter VI and details of compensation provided in such cases shall be intimated to the Central Licensing Authority within thirty working days of receipt of the order issued by the Central Licensing 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 Licensing Authority who may be accompanied by officers of the State Licensing Authority or outside experts as authorized by the Central Licensing 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 Licensing 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 utilized 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;
- (xvi) The Central Licensing 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) Informed Consent Documents (ICD) viz. Patient Information Sheet (PIS) and Informed Consent Form (ICF) complete in all respect & must be got approved from the respective Ethics committee and submitted to CDSCO before enrolling first subject at the respective site.
- (xix) It may kindly be noted that merely granting permission to conduct Clinical Trial study with the drug doesn't convey or imply that based Clinical Trial data generated with the drug, permission to market this drug will automatically be granted to you.
- (xx) The firm should submit clinical photograph of the study population.**
- (xxi) Microbial culture should be conducted for initial, end of treatment and follow up for the same study population.**
- (xxii) Accordingly, revised CT protocol should be submitted to CDSCO before initiation of the study.**

FORM CT-06*(See rules 22, 25, 26, 29 and 30)***PERMISSION TO CONDUCT CLINICAL TRIAL OF NEW DRUG OR
INVESTIGATIONAL NEW DRUG**

The Central Licensing Authority hereby permits Zydus Corporate Park, Scheme No. 63, Survey No. 536, Khoraj (Gandhinagar), Nr. Vaishnodevi Circle, S.G. Highway Ahmedabad-382481, Gujarat, India Telephone No.: 079-71800000, E-Mail: mittal.r@zyduscadila.com to conduct clinical trial of the **new drug** as per protocol no. 21-12, version no. 00, dated 23.11.2021 in the below mentioned clinical trial sites.

2. Details of new drug or investigational new drug:

Names of the new drug or investigational new drug:	Naftifine Hydrochloride Cream USP 2% w/w
Therapeutic class:	Antifungal
Dosage form:	Cream
Composition:	Naftifine Hydrochloride USP.....2% w/w Benzyl Alcohol IP.....1% w/w Cream Base.....q.s.
Indications:	For the treatment of: tinea corporis, tinea cruris and interdigital tinea pedis

Details of clinical trial sites-

Sr. No.	Name of Principal Investigator & Trial sites	Ethics Committee Name/ Registration Number
1.	Dr. Raj Kumar Kothiwala Professor & Head, Department of Dermatology, Jawahar Lal Nehru Medical College & Hospital, Kala Bagh, Ajmer – 305001, Rajasthan	Institutional Ethics Committee, Jawahar Lal Nehru Medical College & Hospital, ECR/1156/Inst/RJ/2018
2.	Dr. Bikash Ranjan Kar Professor & Head, Department of Skin & Venereal Diseases, Institute of Medical Sciences (IMS) & SUM Hospital, Kalinga Nagar, Ghatikia, Bhubaneswar –751003, Odisha.	Institutional Ethics Committee, IMS & SUM Hospital ECR/627/Inst/OR/2014/RR-20
3.	Dr. Pinjala Padmaja Professor, Department of Dermatology, Osmania Medical College & General Hospital, Afzalgunj, Hyderabad-500095, Telangana.	Institutional Ethics Committee, Osmania Medical College ECR/300/Inst/AP/2013/RR-19
4.	Dr. Nipul Vara Assistant Professor, Department of Dermatology, Medical College & SSG Hospital, Anandpura, Vadodara -390001, Gujarat.	Institutional Ethics Committee for Human Research, Department of Pharmacology, Medical College & SSG Hospital ECR/85/Inst/GJ/2013/RR-19

5.	Dr. Projna Biswas Assistant Professor, Department of Dermatology, Calcutta School of Tropical Medicine, 108, Chittaranjan Avenue, College Square, Kolkata-700073, West Bengal.	Clinical Research Ethics Committee, Department of Clinical and Experimental Pharmacology, Calcutta School of Tropical Medicine ECR/194/Inst/WB/2013/RR-20
6.	Dr. Swosti Mohanty Assistant Professor, Department of Dermatology, Era's Lucknow Medical College and Hospital, Sarfarazganj, Hardoi Road, Lucknow-226003, Uttar Pradesh.	Institutional Ethics Committee, Era's Lucknow Medical College and Hospital, Sarfarazganj (ECR/717/Inst/UP/2015/RR-18)
7	Dr. Prerna R. Suryatale Consultant Dermatologist, Jeevan Rekha Hospital, Dr. B.R. Ambedkar Road, Opp. Civil Hospital, Veer Chambers, Belagavi (Belgaum) -590002, Karnataka	Institutional Ethics Committee, Jeevan Rekha Hospital, ECR/1242/Inst/KA/2019
8	Dr. Saurabh Jaiswal Consultant Dermatologist, Alexis Multispeciality Hospital Pvt. Ltd., Survey No. 232, House No. 1313, Mankapur Square, Koradi Road, Nagpur-440030, Maharashtra	Institutional Ethics Committee, Alexis Multispeciality Hospital Pvt. Ltd. ECR/1430/Inst/MH/2020
9	Dr. Ipsa Pandya Consultant Dermatologist, Aatman Hospital, 5, Anveshan Row House, Opp. Bopal Garn BRTS, Bopal-Ghuma Road, Bopal, Ahmedabad-380058, Gujarat	Institutional Ethics Committee, Aatman Hospital, ECR/1565/Inst/GJ/2021

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.

(Dr. V. G. Somani)

Central Licensing Authority
Stamp

Dr. V. G. SOMANI
Drugs Controller General (India)
Dte. General of Health Services
Ministry of Health and Family Welfare
FDA Bhawan, Kotla Road, I.T.O.
New Delhi-110002

New Delhi

Date:

03 FEB 2022