

Food and Drug Administration Silver Spring MD 20993

NDA 205865 NADA 141396

DEEMED GRANTED - MEDICAL GAS CERTIFICATION REQUEST

Air Products and Chemicals, Inc. Attn: Edward J. Csira Team Lead, Quality and Regulatory Compliance 7201 Hamilton Blvd Allentown. PA 18195-1501

Dear Mr Csira:

Please refer to your May 21, 2013, request received on May 24, 2013, for certification of Oxygen, USP as a designated medical gas. You have requested to market Oxygen, USP for human and animal drug use.

Please also refer to your amendment regarding this request dated May 23, 2013.

A request for certification of a medical gas as a designated medical gas submitted under section 575(a)(1) of the Federal Food, Drug, and Cosmetic Act (FD&C Act) is deemed to be granted unless, within 60 days of the filing of the request, FDA finds that one or more of the bases for denying the request listed at section 575(a)(2) of the FD&C Act applies. FDA has made no such finding in connection with your request, and 60 days have passed since your request was filed. Accordingly, by operation of section 575(a)(2) of the FD&C Act, your request for certification of Oxygen, USP as a designated medical gas is deemed to be granted, and you now have in effect an approved new drug application (NDA 205865) and an approved new animal drug application (NADA 141396) for this gas.

If any of the information you have submitted in connection with your request becomes incomplete or inaccurate, please consult section IV.D of the draft guidance document entitled *Certification Process for Designated Medical Gases* (available at http://www.fda.gov/downloads/Drugs/GuidanceComplianceRegulatoryInformation/Guidances/UCM332136.pdf) for instructions on providing FDA with complete, up-to-date information. Please address any such communications to:

Food and Drug Administration Center for Drug Evaluation and Research Office of Pharmaceutical Quality 5901-B Ammendale Road Beltsville, MD 20705-1266

Reference ID: 3825201

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Please include the NDA and NADA numbers listed above at the top of the first page of any such communications.

If you have any questions, please contact Michael Folkendt at (301) 796-1670 or by email at michael.folkendt@fda.hhs.gov.

Sincerely,

{See appended electronic signature page}

Charles J. Andres, Ph.D.
Business Process Improvement Manager
Office of New Animal Drug Evaluation
Center for Veterinary Medicine
FDA

Michael Folkendt Associate Director for Regulatory Affairs Office of Program and Regulatory Operations Office of Pharmaceutical Quality Center for Drug Evaluation and Research FDA

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| /s/ |
| MICHAEL M FOLKENDT 09/25/2015 |

CHARLES J ANDRES 09/28/2015

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