DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration
Silver Spring MD 20993

NDA 205986
NADA 141408

DEEMED GRANTED -
MEDICAL GAS CERTIFICATION REQUEST

Praxair Canada Inc.
Attn: Gordon Edwards, Merchant & Onsite Quality Manager
1 City Centre Drive, Suite 1200
Mississauga, ON L5B 1M2
Canada

Dear Mr. Edwards:

Please refer to your May 27, 2013, request received on June 28, 2013, for certification of Oxygen, USP as a designated medical gas. You have requested to market Oxygen, USP for human and animal drug use.

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 205986) and an approved new animal drug application (NADA 141408) 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you have submitted in connection with your request becomes incomplete or inaccurate, please consult section IV.D of the draft guidance document entitled *Certification Process for Designated Medical Gases* (available at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

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

This is a representation of an electronic record that was signed electronically and this page is the manifestation of the electronic signature.

/s/

MICHAEL M FOLKENDT
09/25/2015

CHARLES J ANDRES
09/28/2015