EC DECLARATION OF CONFORMITY

Company Name: Inogen, Inc. (Legal Manufacturer)
Address: 326 Bollay Drive
Goleta, California 93117
USA

Inogen, Inc.
1125 E. Collins Blvd. Suite 200
Richardson, Texas 75081
USA

Inogen, Inc
1225 Commerce Drive
Richardson, TX 75081
USA

EC Representative Name: Emergo Europe
Address: Prinsessegracht 20
2514 AP The Hague
The Netherlands

Product Information:

Catalog Number/ Part Number	Product Name	Classification	Rule
IO-300	Inogen One G3 Oxygen Concentrator	IIa	11
GS-100	Inogen At Home Oxygen Concentrator	IIa	11
IO-400	Inogen One G4 Oxygen Concentrator	IIa	11
IO-500	Inogen One G5 Oxygen Concentrator	IIa	11

We herewith declare that as of the date of this declaration, the above mentioned medical device(s) meet(s) the requirements of the following EC Council Directives and is supported by the following EC Certificates.

EC Certificate Number: Annex II excluding Section 4, Certificate # CE 670456
BSI Group, The Netherlands B.V.
Notified Body Number: 2797
Address: Say Building
John M. Keynesplein 9
1066 EP Amsterdam

Directives:

- COUNCIL DIRECTIVE 93/42/EEC concerning Medical Devices
- COUNCIL DIRECTIVE 2011/65/EU on the restriction of the use of certain hazardous substances in electrical and electronic equipment (RoHS)

All supporting documentation is retained under the premises of the manufacturer.

Date, Lot Number or Serial Number that CE Mark was first affixed: See RG-01058-05, Inogen CE Marked Products

Signed: _____ Date: October 3, 2019
Title: Inogen Regulatory Affairs and Compliance Director