

## **EU DECLARATION OF CONFORMITY**

## Sensidyne, LP

1000 112<sup>th</sup> Circle North, Suite 100 St. Petersburg, Florida 33716

The undersigned declares that the products named in this certificate meet the applicable provisions of the relevant Union harmonization legislation: Directive 2014/34/EU concerning equipment and protective systems intended for use in potentially explosive atmospheres. This declaration is issued in the sole responsibility of Sensidyne, LP.

Product Type: Air Sampling Pump

Product designation: GilAir 3, Gilair 5

800485-7 GilAir 3 Basic 610-0201-01 GilAir 3 Basic - ATEX 800508-7 GilAir 3 Clock 610-0202-01 GilAir 3 Clock - ATEX

800510-7 GilAir 3 Programmable 610-0203-01 GilAir 3 Programmable - ATEX

800883-7 GilAir 5 Basic - ATEX

800885-7 GilAir 5 Clock 610-0102-01 GilAir 5 Clock - ATEX

800884-7 GilAir 5 Programmable 610-0103-01 GilAir 5 Programmable – ATEX

Manufacturer: Sensidyne, LP

Intended Use: Air Sampling

Personal Sampling Pump Performance Requirements: ISO/EN 13137

Intrinsically Safe:

UL

Class 1, Div 1, Groups A, B, C, D;

**Class 2, E, F, G**;

Class 3

**ATEX** 

→ II 2G

EEx ia IIC T4 (Ta = -20°C to +40°C)

EN 50014:1997 (amendments A1 to A2) EN 50020:2002

ATEX Quality Assurance Notification: SIRA Certification Notified body Number: 0518

EMC: Emissions Standards

EN 55011:2009/A1:2010 Group 1, Class A <u>Industrial, scientific and medical equipment. Radio-frequency disturbance characteristics. Limits and methods of measurement</u>

ICES-003 –Issue 2 Class A Information Technology Equipment (ITE) — Limits and Methods of Measurement FCC Part 15 (per ANSI C63.4:2014)) Class A Verification

EMI: Immunity Standards

EN 61326-1:2013 <u>Electrical equipment for measurement, control and laboratory use. EMC requirements. General requirements</u>

IEC 61000-4-2:2008 / EN 61000-4-2:2009 Electrostatic Discharge Immunity Test

IEC 61000-4-3:2006 / EN 61000-4-3:2006/A1:2008/A2:2010 Radiated, radio-frequency, electromagnetic field immunity

Reference Product Safety Engineering Report 15F172I

Signed: Date: 11/23/2021

Name: Sean Shannon

Title: Manager: Quality Assurance /Regulatory Affairs Sensidyne, LP

Who is the natural and legal person with responsibility for the design, manufacture, packaging and labeling before the device is placed on the market under his own name, regardless of whether these operations are carried out by the Manufacturer or on his behalf by a third party.

