

EU DECLARATION OF CONFORMITY

Sensidyne, LP

1000 112th Circle North, Suite 100 St. Petersburg, Florida 33716

Certificate No: G5000 Issue 8 November 23, 2021

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

Product Type: Portable Air Sampling Pump

Product designation: Gilian 5000, Gilian 3500, Gilian 800i, Gilian 10i

Manufacturer: Sensidyne, LP Intended Use: Air Sampling

Notified Body: FM Approvals Ltd. Notified body Number: CE 1725

1 Windsor Dials Windsor Berkshire UK SL4 1RS

Intrinsically Safe:

US
Class I, II, III Division 1, Groups: A, B, C, D, E, F, and G
Class I, II, III Divis

Class I, II, III Division 1, Groups: A, B, C, D, E, F, and G
IS/I,II,III /1/ ABCDEFG / T4 ta = 45°C

Class I, II, III Division 1, Groups: A, B, C, D, E, F, and G
Class I, Zone 0, Group: IIC

CSA/CAN C22.2 No. 142 Class 3600 2011 2000 Class 3610 2007 CAN C22.2 No. 157 1992 Class 3810 2005 CAN/CSA-E60079-0-02 2002 ANSI/ISA-60079-0 CAN/CSA-E60079-11 2005 2002 ANSI/ISA-60079-11 2002 CAN/CSA C22.2 61010-1 2004

EC FM07ATEX0018X

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Conforming to the following standards:

EN60079-0:2006 EN60079-11:2007 EN60079-26:2004 Report: 3039791EC

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

EMC: Emissions and Immunity Standards

EN 55011: 1998/A1 Group 1 Class B

EN 61326:1997/A1

Reference Product Safety Engineering Report 07F164C / 07F164I

Additional Standards:

Personal Sampling Pump Performance Requirements: ISO/EN 13137

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.