

**TO WHOM IT MAY CONCERN**

**UNDERSIGNED AUTHORITY**

**SLOVENSKÁ OBCHODNÁ A PRIEMYSELNÁ KOMORA /SOPK/**  
**(SLOVAK CHAMBER OF COMMERCE AND INDUSTRY, /SCCI/),**  
a public legal institution from the Law No.9 / 1992 Coll. as amended and revised,  
and on the basis of submitted documents  
**HEREBY CERTIFIES THAT:**

- 1) A company **OXYWISE, s.r.o.**, /hereinafter referred as "**Manufacturer**"/, headquartered in Hurbanova 21, Piešťany, Postcode 921 01, Slovak Republic, ID no. 44 651 465, is the limited liability company established under the laws of the Slovak Republic on March 5, 2009 and registered in the Business Register of the District Court Trnava, Section: Sro, Insert No.: 23387/T, with the subject matter: manufacture of machinery and equipment, wholesale, retail, repair and servicing of medical equipment.
- 2) Manufacturer has its Manufacturing Plant in Piešťany (SK), which has been granted authorisation (license), No. RUVZ/2016/01555/4/Ra-PPL for manufacturing of oxygen and nitrogen generators, under Articles 6(3)(g) and 13(4)(a) of the Act no. 355/2007 Coll., as amended, by the Regional Public Health Authority Trnava, (the state administration for health care and health protection in the Slovak Republic, EU), on September 20, 2016..
- 3) The MANUFACTURER and Medical Devices (*hereinafter referred as "MDs"*) made in the Manufacturing Plant are registered in accordance with the Act no. 362/2011 Coll. with the State Institute for Drug Control (*SIDC*) - the Competent Authority for MDs in the Slovak Republic, EU.
- 4) The MDs which are listed in the Annex, comply with the requirements and standards for the safety of medical devices as required by applicable legislation of the Slovak Republic and the European Union, i.e.:
  - Act no. 355/2007 Coll. on Protection, Support and Development of Public Health and on Amendments and Supplements to Certain Acts, as amended
  - Act no. 362/2011 Coll. on medicines and medical devices and on amendments to certain laws, as amended;
  - Act no. 56/2018 Coll. on product conformity assessment, making a determined product on the market and on the amendment of certain laws,
  - Gov. Regulation no. 582/2008 Coll. as amended by 2015/2013 Coll. laying down details on European technical requirements and conformity assessment procedures for medical devices (MDs) Class IIb under Annex II of the Medical Device Directive no. 93/42/EEC and Directive no. 2011/65/EU;
- 5) The MANUFACTURER is the holder of the EC Certificate of Quality Management System (No.260979-2018-AQ-CZS-RvA), according to the standard ISO 9001:2015, (valid by April 25, 2021), issued by DNV GL - Business Assurance Czech Republic s.r.o. Prague, CZ (AB no. RiA C 024, NL) and the Management System Certificate for design and development, sale, production and installation, service and maintenance of medical oxygen generating systems and filling/reduction ramps (No. 258652-2018-AQ-CZS-NA-Ps Rev.0.0), according to the standard EN ISO 13485:2016 (valid by April 7, 2021), issued by DNV GL & NEMKO PRESAFE AS, Høvik (NO), (CB no.MSD-CO-078 by NO-Accreditation MSYS 018).
- 6) The MANUFACTURER has prepared technical documentation and affixed the CE mark with the MDs specified in the Annex according to Annex II of the MDD no. 93/42/EEC, as amended, therefore the MDs maybe freely sold in all member states of the European Economic Area including the Slovak Republic.
- 7) This Manufacturing & Free Sales Certificate is issued to the MANUFACTURER on its request. Export of the MDs outside EU is not prohibited.

Bratislava, 14 -06- 2018

(Place and date)



Juraj Knopp

(Name, signature of competent officer of SCCI)

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**Annex to M&FSC:**

**LIST OF MEDICAL DEVICES**  
**OXYWISE, s.r.o., Piešťany, Slovakia (SK) - /MANUFACTURER/**

Item:	List of medical devices (MDs)		SIDC Reg. no.	MD Class	EC Declaration of Conformity (EC DoC) / datum of issue
	Product description	Product Name/ Model			
1	<b>Medical Oxygen generator</b> (UMDNS 12883) in 29 standard models with capacity ranging from 0, 4 to 150 Nm <sup>3</sup> /hour up to 95% purity.	Oxygen generator O1	P 98143	II b	EC DoC according the Medical Device Directive no. 93/42/EEC as amended, on May 28, 2018 and EC DoC No. 1700755/OXY/2017 according the Pressure Equipment Directive 2014/68/EU, Modules B+D, on 16/06/2017 both issued by OXYWISE, s.r.o. Piešťany (SK) – the Manufacturer
		Oxygen generator O2			
		Oxygen generator O4			
		Oxygen generator O6			
		Oxygen generator O9			
		Oxygen generator O12			
		Oxygen generator O15			
		Oxygen generator O20			
		Oxygen generator O20+			
		Oxygen generator O27			
		Oxygen generator O27+			
		Oxygen generator O35			
		Oxygen generator O35+			
		Oxygen generator O50			
		Oxygen generator O50+			
		Oxygen generator O65			
		Oxygen generator O65+			
		Oxygen generator O80			
		Oxygen generator O80+			
		Oxygen generator O100			
		Oxygen generator O100+			
Oxygen generator O125					
Oxygen generator O125+					
Oxygen generator O150					
Oxygen generator O150+					
Oxygen generator O80T					
Oxygen generator O100T					
Oxygen generator O125T					
Oxygen generator O150T					