

CERTIFICATE No. GIF-IW-400/0061\_05\_01/04/81/14

*Main Pharmaceutical Inspector***CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER****Part 1**

Issued following an inspection in accordance with Art. 111(5) of Directive 2001/83/EC as amended

**Main Pharmaceutical Inspector***/the Competent Authority of Poland/*

confirms the following:

the manufacturer

**Air Products Sp. z o.o.**  
59 Pory Str., 02-757 Warsaw, POLAND

site address

**Air Products Sp. z o.o.**  
7 Waryńskiego Str., 47-220 Kędzierzyn-Koźle, POLAND

has been inspected under the national inspection programme in connection with manufacturing authorisation No. **GIF-IW-400/0061/01/175/ZW72/14** in accordance with Art. 40 of Directive 2001/83/EC transposed in pharmaceutical law of 6<sup>th</sup> of September 2001 (Dz. U. z 2008 r. Nr 45, poz. 271, z późn. zm.).

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on **06-07/05/2014**, it is considered that it complies with the Good Manufacturing Practice requirements laid down in Directive 2003/94/EC.

This certificate reflects the status of the manufacturing site at the time of the inspection noted above and should not be relied upon to reflect the compliance status if more than three years have elapsed since the date of that inspection, after which time the issuing authority should be consulted.

This certificate is valid only when presented with all pages and both Parts 1 and 2.

The authenticity of this certificate may be verified in EudraGMP. If it does not appear, please contact the issuing authority.



date: 2014 -08- 07

Main Pharmaceutical Inspectorate  
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Zofia Ulz  
Main Pharmaceutical Inspector

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Part 2

Human Medicinal Products

**1 MANUFACTURING OPERATIONS**

**1.2 Non-sterile products**

**1.2.1 Non-sterile products (list of dosage forms)**

1.2.1.7 Medicinal gases

**1.6 Quality control testing**

**1.6.3 Chemical/Physical**



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