

# Certificate

acc. to **ISO 13485:2016**

Medical Devices - Quality Management Systems - Requirements for Regulatory Purposes

Certificate Registration No.: **19-1605-Q**

TUV USA, Inc. hereby certifies that the quality management system of the company mentioned below is in conformance with **ISO 13485:2016** for Medical Devices-Quality Management Systems-Requirements for Regulatory Purposes.

**Polymap Wireless LLC**  
**310 S. Williams Blvd. Suite 230**  
**Tucson, AZ 85711, USA**

Additional sites covered by QM System: **N/A**

Scope:

**Design, Manufacturing, Service and Distribution of Wireless Data Transmission Systems**

The validity of this certification document can be obtained by contacting the TUV USA, Inc. Office.

**TUV USA, Inc. (a Member of the TÜV NORD Group)**  
215 Main Street, Suite 1, Salem, NH 03079, USA  
Tel: 001-603-870-8023, Fax: 001-603-870-8026, Email: [medical-usa@tuv-nord.com](mailto:medical-usa@tuv-nord.com)



Audit Report Reference No.: **18-8042 SA2-UP**  
Certificate Initial Issue Date: **14-Oct-2016**  
Certificate Revised Date: **29-JAN-2019**

Effective Date:  
**29-JAN-2019 / ed. 2**

Valid Until:  
**13-OCT-2019**



Bradley Chen  
Director, Medical Products Division  
TUV USA, Inc.