

# Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016

This is to certify that:

Tensentric  
2900 Center Green Court  
Boulder  
Colorado  
80301  
USA

Holds Certificate No:

**FM 559260**

and operates a Quality Management System which complies with the requirements of ISO 13485:2016 for the following scope:

Contract Design and Development, Manufacturing, Installation, packaging of Non- Sterile and Sterile product to customer specifications, and Distribution of systems to support diagnostic, preventative, therapeutic, life-sustaining, anatomical, pharmaceutical assisting, infection control, and imaging.

For and on behalf of BSI:



Graeme Tunbridge, Senior Vice President Medical Devices

Original Registration Date: 2010-05-20

Latest Revision Date: 2022-03-07

Effective Date: 2022-05-20

Expiry Date: 2025-05-19

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Certificate No: **FM 559260**

Location	Registered Activities
Tensentric 2900 Center Green Court Boulder Colorado 80301 USA	Contract Design and Development, Manufacturing, Installation, packaging of Non-Sterile and Sterile product to customer specifications, and Distribution of systems to support diagnostic, preventative, therapeutic, life-sustaining, anatomical, pharmaceutical assisting, infection control, and imaging.
Tensentric 474 South Taylor Avenue, Suite C Louisville Colorado 80027 USA	Contract Design and Development, Manufacturing, Installation, packaging of Non- Sterile and Sterile product to customer specifications, and Distribution of systems to support diagnostic, preventative, therapeutic, life-sustaining, anatomical, pharmaceutical assisting, infection control, and imaging.



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To be read in conjunction with the scope above or the attached appendix.

Americas Headquarters: BSI Group America Inc., 12950 Worldgate Drive, Suite 800, Herndon, VA 20170-6007 USA  
A Member of the BSI Group of Companies.