Triomed

PRESS RELEASE

Triomed strengthens commitment to quality with ISO 13485:2012 –

Quality Management System Certification, April 22, 2015.

LUND, SWEDEN – Triomed, a manufacturer of medical devices for the treatment of congestive heart failure, is proud to announce it has received ISO 13485:2004 certification for the design, development, production, service, sales and distribution of active and non-active medical devices for the ultrafiltration of congestive heart failure patients with fluid overload.

Certification to ISO 13485:2012 requires a thorough review of the company's internal quality management system processes by an accredited third-party auditing organization to ensure the company is capable of consistently delivering products and services that meet both customer needs and expectations and regulatory requirements for the industry. Triomed's quality management system has been registered by TÜV SÜD Product Service GmbH, one of the world's leading providers of management system certification, recognized for their expert auditors and thorough assessments.

"As an organization we take pride in ensuring customer satisfaction by providing medical devices that fulfil our customer needs. Receiving this certification emphasizes our commitment to quality throughout our business," says Anders Wallenås, CEO.

Quality Management System – Quality of Life – inside and out

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Triomed is a Swedish medical engineering company developing wearable dialysis system for Cardiac and Renal patients. Triomed was founded in 2005 and has today 15 employees and the head quarter is located in Lund, Sweden.