



## FINAL DOCUMENT

**Title:** Quality Management System – Medical Devices –  
Guidance on the Control of Products and Services  
Obtained from Suppliers

**Authoring Group:** GHTF Study Group 3

**Endorsed by:** The Global Harmonization Task Force

**Date:** December 11, 2008

A handwritten signature in black ink, appearing to read 'R. Rotter'.

Dr. Roland Rotter, GHTF Chair

The document herein was produced by the Global Harmonization Task Force, which is comprised of representatives from medical device regulatory agencies and the regulated industry. The document is intended to provide *non-binding* guidance for use in the regulation of medical devices, and has been subject to consultation throughout its development.

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# Ghtf Sg3 Quality Management System Medical Devices

**Jin-Ying Zhang**



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