BUREAU VERITAS
Certification

Qatari German Company for Medical Devices

Head Office and Operative Site:
Building 136, Street 54 Abu Hamour, P.O. Box 22556 – DOHA, QATAR

Bureau Veritas Italia S.p.A. certify that the Management System of the above organisation has been audited and found to be in accordance with the requirements of the management system standards detailed below

Standard

EN/ISO 13485:2012

Scope of certification

Manufacture of sterile single use medical devices such as: hypodermic syringes with or without needles, safety syringes with or without needles, insulin syringes with and without needles, IV cannula with and without ports, hypodermic needles.

Certification awarded in conformity with the requirements of ACCREDIA DT 02-DC REV.00

Certification cycle start date: **01 May 2016**
Subject to the continued satisfactory operation of the organization's Management System, this certificate expires on**: 30 April 2019**

Original certification date: **01 May 2007**

Certificate No. IT267753 Version N.**1** Revision date: **01 May 2016**

ANDREA FILIPPI – Local Technical Manager

Certification body address:

Bureau Veritas Italia spa, Via Miramare, 15, 20126 Milano, Italia
Further clarifications regarding the scope of this certificate and the applicability of the management system requirements may be obtained by consulting the organisation.
To check this certificate validity please refer to the website www.bureauveritas.it

ACCREDIA
L'ENTE ITALIANO DI ACCREDITAMENTO

SGQ	N° 009A	PRS	N° 076C
SGA	N° 008D	SGE	N° 009M
PRD	N° 009B	EMAS	N° 004P
SCR	N° 008F	GHG	N° 008O
FSMS	N° 003I	ISP	N° 006E

Membro degli Accordi di Mutuo Riconoscimento EA e IAF
Signatory of EA and IAF mutual Recognition Agreements