

DECLARATION OF CONFORMITY



MANUFACTURER:

ZHEJIANG RUNQIANG MEDICAL INSTRUMENTS CO., LTD.
NO.618 DADE ROAD, JIAXING XIUZHOU DISTRICT JIAXING CITY,
ZHEJIANG PROVINCE, P.R. CHINA
TEL: +86-573-88107358, +86-573-88106708,
FAX: +86-573-88110618

MEDICAL DEVICE:

DISPOSABLE ANAESTHESIA NEEDLE AND ANAESTHESIA KIT

CLASSIFICATION - ANNEX IX:

CLASS III, RULE 7 OF ANNEX IX

CONFORMITY ASSESSMENT ROUTE:

ANNEX II .3+ II .4

WE, ZHEJIANG RUNQIANG MEDICAL INSTRUMENTS CO., LTD., HEREWITH DECLARE THAT THE STATED MEDICAL DEVICES MEET THE TRANSPOSITION INTO NATIONAL LAW, THE PROVISIONS OF COUNCIL DIRECTIVE 93/42/EEC CONCERNING MEDICAL DEVICES, AS AMENDED BY 2007/47/EEC; ALL SUPPORTING DOCUMENTATION IS RETAINED AT THE PREMISES OF THE MANUFACTURER.

STANDARDS APPLIED: *RALATED APPLICABLE HARMONIZED STANDARDS(PUBLISHED IN THE OFFICIAL JOURNAL OF THE EUROPEAN COMMUNITIES)*

NOTIFIED BODY:

TÜV SÜD PRODUCT SERVICE GMBH
RIDLERSTR 65, D-80339 MÜNCHEN, GERMANY

IDENTIFICATION NUMBER

CE 0123

(EC) CERTIFICATE(S):

G1 12 09 82107 003

EC REP

EUROPEAN REPRESENTATIVE:

Shanghai International Holding Corp. GmbH (Europe)
Eiffestrasse, 80, 20537, Hamburg, Germany Tel: 0049-40-2513175, Fax: 0049-40-255726

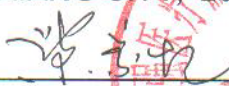
START OF CE-MARKING:

2013-05-07

PLACE, DATE OF DECLARATION:

JIAXIANG CITY, 2013-05-16

SIGNATURE:


NAME: MRS CAI JIA YI
POSITION: CAIJIAIYI DIRECTOR