Declaration of Conformity TO COUNCIL DIRECTIVE 93/42/EEC CONCERNING MEDICAL DEVICES

MANUFACTURER: WEIGAO MEDICAL INTERNATIONAL CO., LTD.

No.1 Weigao Road, High-tech Industrial Development Zone, 264210 Weihai, Shandong Province,

PEOPLE'S REPUBLIC OF CHINA

MEDICAL DEVICE: Insulin syringe

0.3ML \ 0.5ML \ 1ML

CLASSIFICATION - ANNEX IX: CLASS IIA, RULE 6

UMDNS CODES: 13941
CONFORMITY ASSESSMENT ROUTE: ANNEX II.3

We, <u>Weigao Medical International Co., Ltd.</u>, herewith declare that the stated medical devices meet the transposition into national law, the provisions of Council Directive 93/42/EEC of 14 June 1993 concerning medical devices;

INCLUDING, AT 21 MARCH 2010, THE AMENDMENTS BY COUNCIL DIRECTIVE 2007/47/EEC.

ALL SUPPORTING DOCUMENTATION IS RETAINED AT THE PREMISES OF THE MANUFACTURER.

THE MANUFACTURER IS EXCLUSIVELY RESPONSIBLE FOR THE DECLARATION OF CONFORMITY.

NOTIFIED BODY: TÜV SÜD PRODUCT SERVICE GMBH

RIDLERSTR 65, D-80339 MÜNCHEN, GERMANY

(EC) CERTIFICATE(S): G1 087117 0013 Rev.01

EC REP MEDNET EC-REP GMBH

BORKSTRASSE 10, 48163 MUENSTER, GERMANY

VALID UNTIL: 2024.04.16

PLACE, DATE OF DECLARATION: WEIHAI, 2021-09-10

SIGNATURE:

NAME: SONG XINLAN

POSITION: Quality Department Manager