DECLARATION OF CONFORMITY TO COUNCIL DIRECTIVE 93/42/EEC CONCERNING MEDICAL DEVICES

MANUFACTURER:	*ZERO-PLUS INTERNATIONAL LTD.* *ROOM 1004, 10/F, JOIN-IN HANG SING CENTRE,* *71-75 CONTAINER PORT ROAD,* *KWAI CHUNG, NEW TERRITORIES, HONG KONG*
MEDICAL DEVICE:	*VENENWALKER BASIC – HNS018839*
CLASSIFICATION - *ANNEX IX*:	*CLASS IIA, RULE 9*
UMDNS CODE:	*10969*
CONFORMITY ASSESSMENT ROUTE:	*ANNEX II EXCLUDING (4)*

WE, ZERO-PLUS INTERNATIONAL LTD., 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 THIS DOC

NOTIFIED BODY:	TÜV SÜD PRODUCT SERVICE GMBH RIDLERSTR 65, D-80339 MÜNCHEN, GERMANY
IDENTIFICATION NUMBER	CE 0123
(EC) CERTIFICATE(S):	G1 070522 0014 REV. 01
VALID UNTIL:	*2024-05-26*
EC REP EUROPEAN REPRESENTATIVE:	GLOBALMIND CONSUMER ELECTRONICS GMBH ERNST-MANTIUS-STR.11 21029 HAMBURG, GERMANY

START OF CE-MARKING: *2019-08-21*

PLACE, DATE OF DECLARATION: HONG KONG, *2020-03-25*

SIGNATURE:

NAME: ESTHER LAU
POSITION: GENERAL MANAGER