




EU DECLARATION OF CONFORMITY

According to ANNEX IV of Regulation (EU) 2017/745 on Medical Devices

Manufacturer:	BQ Plus Medical Co., Ltd. Add.: No. 18, Che Ye Road, Che Dun Town, Songjiang, 201611 Shanghai, P.R. China
SRN:	CN-MF-000010259
Trademark:	
Product Name:	Extension Sets
Model:	Extension Set; Extension Line; IV Extension Line; Secondary Infusion Set; S-ES-000; S-ES-001; S-ES-002; S-ES-003
European Representative:	Prolinx GmbH Add.: Brehmstr.56, 40239 Dusseldorf, Germany
SRN:	DE-AR-000005129
EMDN Code:	A030201 EXTENSIONS
Intended Purpose:	This device is intended for direct injection, intermittent infusion, continuous infusion or aspiration.
Basic UDI:	697262011BQES1DB: Extension Set, S-ES-002, S-ES-003; 697262011BQES2DD: Extension Line; 697262011BQES3DF: IV Extension Line; 697262011BQES4DH: Secondary Infusion Set; 697262011BQES5DK: S-ES-000; 697262011BQES6DM: S-ES-001
Classification acc. to MDR	Is
Ax. VIII:	
Conformity Assessment	Annex IX Chapter 1, Section 2 and 3, MDR (2017/745)
Procedure:	
CE Certificate No.:	HZ 2087599-1
Name and ID of the Notified	Name: TÜV Rheinland LGA Products GmbH
Body:	Address: Tillystraße 2, 90431, Nürnberg, Germany (Notified Body number: 0197)

We, the manufacturer, herewith declare under our sole responsibility that the above-mentioned products meet the provisions of the Regulation (EU) 2017/745 on Medical Devices (MDR). All

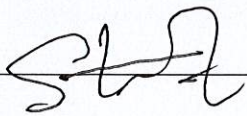


supporting documentations are retained under the premises of the manufacturer.

Place: Shanghai

Position: Management Representative

Name: Steven Zhang

Signature:  2025.06.03

Date: 2025.06.03