

EC DECLARATION OF CONFORMITY

Manufacturer: Jiangsu DYNAMIC Medical Technology Co., Ltd
Add: No. 66, ShunXing Road, Lujia Town, Kunshan,
Suzhou, Jiangsu, China
SRN: CN-MF-000020125

European Representative: Shanghai International Holding Corp. Gmbh (Europe)
Add.: Eiffestrasse 80, 20537 Hamburg, Germany
SRN: DE-AR-000000001

Product name: Medical Suction Unit

Model: DS502, DS504, DS506, DS508, DS510, DSG502,
DSG504, DSG506, DSG508, DS01, DS02,
DS04, DSG01, DSG02, DSG04, DS1000, DS2000,
DS3000, DS5000, DS03/06/10, DSG03/06/10,
DS803/805/808/810, DS301, DS302, DSG301,
DSG302, DS2501M, DS3701M, DS7501CS, DAS001,
DAS002, DS910, DS915, DS920, DS930, DS940, DS950,
DAS001D, DAS002D

Introduction for use: It is used for the suction and purification of bacterial
aerosols between medical staff and patients during
dental treatment.

EMDN Code: Z120499

Basic UDI-DI 697141739DS1102B6

Classification: Class I, According to Rule 1, Annex VIII of Regulation

Applied Standards: ISO:15223-1:2021/Amd 1:2025,
ISO 20417:2026, ISO 13485:2016,
ISO 24971:2020, IEC 62366-1:2015+A 1:2020
EN ISO 10993-1:2025,
EN ISO 10993-5:2009/A11:2025,
EN ISO 10993-10:2023,
IEC 60601-1-2020,
MDCG 2019-15.

Conformity assessment procedure:

EU DECLARATION OF CONFORMITY following the
Annex II + Annex III + Article 19 of MDR (EU) 2017/745.

We herewith declare that the above mentioned products meet the transposition into national law, the provisions of the following EU Regulation and Standards. All supporting documentations are retained under the premises of the manufacturer. The manufacturer is exclusively responsible for the declaration of conformity. General applicable regulations, directives: Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices, amending Directive 2001/83/EC, Regulation (EC) No 178/2002 and Regulation (EC) No 1223/2009 and repealing Council Directives 90/385/EEC and 93/42/EEC.

Signature:

Name:

Title: General Manager

Place: Suzhou, China.

Date of Issue: Apr 30, 2026

