



DECLARATION OF CONFORMITY
ACCORDING TO (EU) 2017/745 MEDICAL DEVICE REGULATION

EU Representative

Name: Shanghai International Holding
Corp. GmbH(Europe)
Address: Eiffestrasse 80, 20537 Hamburg,
Germany
SRN: DE-AR-000000001

Conformity Assessment

Conformity Assessment Procedure
Annex II+III of Regulation (EU) 2017/745

Applicable Standards

ISO 14971: 2019, ISO 15223-1:
2021+A1:2025
ISO 20417: 2021, EN ISO 10993-1: 2020
ISO 10993-5: 2009, ISO 10993-10: 2021,
ISO 10993-11: 2017,BS EN1644:1997,
BS EN1644-2: 2000, EN ISO
13485:2016+A11:2021

Remark

The declaration of conformity is valid in
connection with the release ISO 13485
Certificate by TUV SUD with Notified Body
identification no.0123:

ISO 13485:2016 Certificate: Q5 002037
0008 Rev. 06

Expire date of the Certificate: 2028-08-31

Place, Date of Issue: Hubei province,
P.R.C,2025-08-11

Manufacturer

Name: Allmed Medical Products Co.,Ltd
ADDRESS: No.18 Qixing Road, Majiadian Town,
443200Zhijiang City, Hubei province, PEOPLE'S
REPUBLIC OF CHINA
SRN: CN-MF-000007970

Product Information

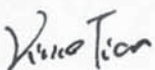
Name: Non-sterile Adhesive Plasters
Model: 500081, 500082, 500083, 500084
EMDN: M040101
Basic UDI: 69415580adhesivedrNSSJ
Classification: Class I, According to Rule 4, Annex
VIII, Regulation (EU) 2017/745

Declaration

We herewith declare in our own responsibility that
the above mentioned products meet the
Regulation (EU) 2017/745 and, if applicable, with
any other relevant Union legislation that provides for
the issuing of an EU declaration of conformity. All
supporting documentations are retained under the
premises of the manufacturer. We are exclusively
responsible for the declaration of conformity.

Position: General Manager of Quality

Name: Vince Tian Place: Zhijiang City

Signature:  Date:2025-11-04