

Fraunhofer TESTED[®] DEVICE

hawo GmbH HM 5000 PRINTPAK MED Report No. HA 2401-1488

Single product Hygienic Design

This document only applies to the named product in its original state and is valid for a period of 5 years from the date the first document was issued. The document can be verified under www.tested-device.com.

Detailed information and parameters of the test environment can be found in the Fraunhofer IPA test report.

Qualification Certificate

This is to certify that the product mentioned above, provided by

hawo GmbH Obrigheim, Germany

has been awarded a Fraunhofer certificate TESTED DEVICE bearing the report number HA 2401-1488.

The system HM 5000 DC-VI PRINTPAK MED SEALING EQUIPMENT + INKJET (V2A stainless steel) is principally suitable for use in hygienic areas up to the following GMP Classes according to EU GMP Annex 1:

System component	Suitability
HM 5000 C INKJET	up to GMP-Klasse D
HM 5000 DC-VI PRINTPAK MED	up to GMP-Klasse C
Overall result	up to GMP Class D

However, this recommendation only pertains to the operating utility when in a resting state. An overall assessment of the can only be made after its installation in the production environment.

HA 2401-1488 Report No. first document Stuttgart, March 1, 2024 Place, date of first document issued

Report No. current document

Place, current date on behalf of Dr. Burn Dr.-Ing. Frank Bürger, Project Manager Fraunhofer IPA

