

CellPath Ltd

80 Mochdre Enterprise Park,
Newtown, Powys,
SY16 4LF Mid Wales, UK

Tel: +44 (0) 1686 611 333
Fax: +44 (0) 1686 622 946
Email: sales@cellpath.com
Website: www.cellpath.com

EU DECLARATION OF CONFORMITY – TISSUEWRAP

Date: 27/07/2021

This Declaration of Conformity has been issued solely by CellPath Limited (Company Number 01831261) in accordance with the definitions and regulations as described in the EU Regulation 2017/746 of The European Parliament and of the Council on In-Vitro Diagnostic Medical Devices.

General Information

Manufacturer CellPath Limited

Address of Production Facility CellPath Ltd

Unit 80, Mochdre Enterprise Park

Newtown, Powys Mid Wales, UK SY16 4LE

EMERGO EUROPE (NL-AR-000000116)

EU Authorised Representative Prinsessegracht 20

2514 AP The Hague The Netherlands

Product Name Histological tissue cassette biopsy insert IVD

Product Description A flexible and porous material designed to securely hold a biopsy specimen within a

tissue cassette, to prevent loss of the specimen during specimen processing (e.g., fixation, dehydration, infiltration) in preparation for subsequent cytological or histological examination. The flexible and porous material (e.g., paper, synthetic polymer) is intended to allow infiltration of reagents and maximize fluid-exchange during tissue processing. It is available in various forms (e.g., sheet, bag, foam pad) and

is placed in a tissue processing cassette, with the biopsy specimen wrapped or

contained inside it. This is a single-use device.

Product Classification (Classified in accordance with EU Regulation 2017/746) CLASS A





CellPath Ltd 80 Mochdre Enterprise Park, Newtown, Powys,

+44 (0) 1686 611 333 +44 (0) 1686 622 946 Email: sales@cellpath.com Website: www.cellpath.com

DEVICE(S) INFORMATION				
PART NUMBER	GTIN	UDI-DI	DESCRIPTION	YEAR OF CE MARKING
EBB-0102-10A	05060470564476	05060470EBB001RK	TISSUEWRAP (60mm x 60mm) - WHITE	2006

Conformity to the aforementioned regulation is demonstrated by the company's compliance to the following international manufacturing standard:

EN ISO 13485:2016 - Medical Devices, Quality Management Systems - Requirements for Regulatory Purposes

Signed on behalf of the Manufacturer CellPath Limited

Address of Issue CellPath Ltd

Unit 80, Mochdre Enterprise Park

Newtown, Powys Mid Wales, UK **SY16 4LE**

27/07/2021 Date of Issue

Name of Signatory Hannah Moore

Position of Signatory Product Manager

Signature H Moore

Tissuewrap - Declaration of Conformity - EU

Final Audit Report 2021-08-05

Created: 2021-07-27

By: Rosa Moore (rosa.moore@cellpath.com)

Status: Signed

Transaction ID: CBJCHBCAABAAcEkVmkfvOsEgn_Q8OwO86EwRKCFrO_Jw

"Tissuewrap - Declaration of Conformity - EU" History

Document created by Rosa Moore (rosa.moore@cellpath.com) 2021-07-27 - 12:50:46 PM GMT- IP address: 62.232.237.234

Document emailed to H Moore (hannah.moore@cellpath.co.uk) for signature 2021-07-27 - 12:51:09 PM GMT

Email viewed by H Moore (hannah.moore@cellpath.co.uk)

Document e-signed by H Moore (hannah.moore@cellpath.co.uk)

Signature Date: 2021-08-05 - 8:42:23 AM GMT - Time Source: server- IP address: 94.193.189.30

Agreement completed. 2021-08-05 - 8:42:23 AM GMT