

Viamed Ltd

Packaging and Packaging Waste Regulation (PPWR)

EU Declaration of Conformity

Regulation (EU) 2025/40

Document Control

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Applicable Legislation	Regulation (EU) 2025/40 on Packaging and Packaging Waste (PPWR), Annex VIII

1. Purpose

This Declaration of Conformity has been prepared in accordance with Annex VIII of Regulation (EU) 2025/40 on Packaging and Packaging Waste (PPWR).

Its purpose is to demonstrate Viamed Ltd's commitment to meeting applicable packaging legislation and to provide customers, competent authorities and other interested parties with a formal declaration regarding packaging introduced into the supply chain by Viamed Ltd.

This declaration is supported by the Viamed Ltd PPWR Technical File (PPWR-002) and associated supporting documentation.

2. Legal Basis

This Declaration of Conformity has been prepared in accordance with Annex VIII of Regulation (EU) 2025/40 on Packaging and Packaging Waste (PPWR). It shall be read in conjunction with the supporting technical documentation maintained by Viamed Ltd as part of its PPWR Technical File.

3. Company Details

Company Name

Viamed Ltd

Registered Address

15 Station Road

Cross Hills

Keighley

West Yorkshire

BD20 7DT

United Kingdom

VAT Number:

GB287389593

EORI Number:

GB287389593000

4. Scope

This declaration applies to packaging introduced by Viamed Ltd during the supply of products, including secondary and tertiary packaging used for storage, handling and transport.

Examples include:

- Corrugated cardboard cartons
- Shipping boxes
- Mailing bags
- Protective packaging
- Void-fill materials
- Packaging tape

- Shipping labels
- Other transport packaging introduced by Viamed Ltd

Primary product packaging supplied by original equipment manufacturers remains the responsibility of the original manufacturer.

Where manufacturer packaging is supplied unchanged, Viamed Ltd does not alter its design or regulatory characteristics.

5. Declaration

Viamed Ltd declares, under its sole responsibility, that the packaging introduced by the company into the supply chain has been reviewed against the applicable requirements of Regulation (EU) 2025/40 based upon the technical information currently available to Viamed Ltd.

This declaration is made on the basis of:

- information currently available from approved suppliers;
- supplier declarations and technical documentation;
- Viamed Ltd's Packaging & Waste Reduction Statement;
- Viamed Ltd's Environmental Management processes;
- continual review of packaging used within the business.

Where evidence from suppliers is still being collected as part of the phased implementation of PPWR, the associated Technical File will be updated accordingly.

6. Compliance Statement

Viamed Ltd confirms that:

- packaging introduced by Viamed Ltd is reviewed to minimise unnecessary packaging whilst maintaining product protection;
 - packaging is selected to support safe transportation of medical, veterinary and specialist electronic equipment;
 - reusable packaging is utilised wherever practical;
 - recyclable packaging is used wherever reasonably practicable;
 - packaging suppliers are encouraged to reduce environmental impact through continual improvement;
 - supporting technical documentation is maintained as part of the PPWR Technical File.
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7. Limitations

Viamed Ltd is primarily a distributor of medical, veterinary and specialist electronic products.

The company does not generally manufacture packaging nor alter manufacturer packaging supplied with products.

Responsibility for compliance of original product packaging remains with the original manufacturer unless Viamed Ltd has introduced or modified the packaging.

This declaration therefore applies only to packaging introduced or controlled by Viamed Ltd.

8. Supporting Documentation

This declaration is supported by, but not limited to:

- PPWR-002 Technical File Index
 - PPWR-003 Packaging Evidence Register
 - Packaging & Waste Reduction Statement
 - Environmental Policy
 - Sustainable Procurement Policy
 - Carbon Reduction Plan
 - Supplier Code of Conduct
 - Supplier Packaging Declarations
 - REACH Statements
 - RoHS Statements
 - Technical Data Sheets where applicable
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9. Document Maintenance

This declaration shall be reviewed:

- annually;
- following significant legislative changes;
- following material changes to packaging used by the business;
- following significant supplier changes.

Supporting evidence shall be maintained within the PPWR Technical File.

10. Authorised Declaration

I declare that the information contained within this declaration is accurate to the best of my knowledge and that Viamed Ltd will continue to maintain appropriate technical documentation supporting this declaration.

Signed



Helen Lamb

Director

Viamed Ltd

Date: 9th July 2026