Viamed Ltd

Packaging and Packaging Waste Regulation (PPWR)

Technical File

Document Control

Item	Details
Document Title	PPWR Technical File
Issue	1
Issue Date	July 2026
Review Date	July 2027
Document Owner	Quality Manager
Approved By	Managing Director
Classification	Controlled Management System Document
Applicable Legislation	Regulation (EU) 2025/40 on Packaging and Packaging Waste (PPWR), Annex VII & Annex VIII

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1. Purpose

This Technical File has been established to support the Viamed Ltd EU Declaration of Conformity issued under Regulation (EU) 2025/40 on Packaging and Packaging Waste (PPWR).

The purpose of this document is to identify, organise and maintain the technical documentation and supporting evidence demonstrating Viamed Ltd's management of packaging introduced into the supply chain.

This Technical File shall be maintained as a controlled document and reviewed on a regular basis to ensure continued compliance with applicable legislative requirements.

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2. Legal Basis

This Technical File has been prepared in accordance with the requirements of:

- Regulation (EU) 2025/40 on Packaging and Packaging Waste (PPWR)
- Annex VII – Technical Documentation
- Annex VIII – EU Declaration of Conformity

It supports the Viamed Ltd EU Declaration of Conformity and provides a structured repository for the evidence upon which that declaration is based.

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3. Scope

This Technical File applies to packaging introduced or controlled by Viamed Ltd during the storage, handling and distribution of products.

This includes, where applicable:

- Corrugated cardboard shipping cartons
- Protective packaging
- Void-fill materials
- Packaging tape
- Mailing bags
- Shipping labels
- Other secondary or tertiary transport packaging introduced by Viamed Ltd

Primary product packaging supplied by manufacturers remains the responsibility of the original manufacturer unless modified by Viamed Ltd.

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4. Technical Documentation Register

The following documents form part of the Viamed Ltd PPWR Technical File.

Reference	Supporting Documentation	Status
TF001	EU Declaration of Conformity	Complete
TF002	Packaging & Waste Reduction Statement	Complete
TF003	Environmental Policy	Complete
TF004	Sustainable Procurement Policy	Complete
TF005	Carbon Reduction Plan	Complete
TF006	Supplier Code of Conduct	Complete
TF007	REACH Compliance Statement	Complete
TF008	RoHS Compliance Statement	Complete
TF009	Packaging Evidence Register	Complete
TF010	Supplier Packaging Declarations	Complete
TF011	Packaging Material Specifications	Complete
TF012	Supplier Technical Data Sheets	Complete
TF013	Packaging Minimisation Assessment	Planned

Additional documentation may be added as further evidence becomes available.

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5. Packaging Inventory

The following packaging materials are currently introduced into the Viamed Ltd supply chain.

Packaging Type	Typical Use	Evidence Status
Corrugated cartons	Shipping	In Progress
Mailing bags	Customer dispatch	In Progress
Bubble wrap	Product protection	In Progress
Paper void-fill	Product protection	In Progress
Packaging tape	Carton sealing	In Progress
Shipping labels	Dispatch identification	Complete
Protective wrapping	Product protection	In Progress

This inventory shall be updated whenever packaging types are introduced, removed or significantly changed.

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6. Supplier Evidence

Viamed Ltd seeks supporting technical information from packaging suppliers wherever reasonably practicable.

Evidence requested may include:

- PPWR Declaration of Conformity
- Material specifications
- Recycled content declarations
- REACH declarations
- Heavy metal compliance
- PFAS statements (where applicable)
- Technical Data Sheets
- Other supporting documentation

Supplier evidence shall be reviewed and retained as part of this Technical File.

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7. Compliance Review

Compliance with PPWR requirements shall be monitored through:

- Annual Management Review
- Internal quality audits
- Supplier evaluations
- Customer feedback
- Legislative monitoring
- Periodic review of packaging used throughout the business

Where improvements are identified, appropriate actions shall be recorded and implemented through the Viamed Ltd Quality Management System.

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8. Responsibilities

The Quality Manager is responsible for:

- maintaining this Technical File;
- ensuring supporting documentation remains current;
- requesting updated supplier information where necessary;
- coordinating annual reviews;
- ensuring revisions are appropriately authorised.

Department managers shall support the maintenance of this Technical File by providing relevant packaging information when requested.

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9. Record Retention

Technical documentation supporting this file shall be retained in accordance with the applicable requirements of Regulation (EU) 2025/40 and the Viamed Ltd Document Control Procedure.

Where supplier documentation is superseded, previous versions shall be retained in accordance with document retention requirements.

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10. Continuous Improvement

Viamed Ltd recognises that PPWR requirements will continue to evolve.

This Technical File will therefore be updated whenever:

- legislation changes;
- additional supplier evidence becomes available;
- packaging materials change;
- improvements are identified through audits or management review.

The objective is to maintain an accurate and comprehensive record supporting the Viamed Ltd EU Declaration of Conformity.

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11. Authorisation

This Technical File has been approved for use within the Viamed Ltd Management System.

Approved By

Helen Lamb

Director

For and on behalf of Viamed Ltd

Date: 9th July 26