

everX Flow™

SHORT-FIBRE REINFORCED FLOWABLE COMPOSITE FOR DENTIN REPLACEMENT

For use only by a dental professional in the indications for use.

INDICATIONS FOR USE

everX Flow is a reinforcing dentin replacement material suitable for: All direct composite restorations including large posterior cavity, deep class II inlays, cavities with anatomical form preparation after amalgam removal and cavities where inlays and amalgams would also be used.

2. Core buildups

Note:

Although everX Flow features a high strength and fracture toughness, it should always be fully covered with a layer of light-cured universal restorative composite to ensure sufficient wear resistance.

CONTRAINDICATIONS

1. Direct pulp capping.

2. Composite buildups on the pulpa.

3. Composite buildups on dentin and surface composite layer. Always cover with a layer of light-cured universal composite.

PRODUCT DESCRIPTION

everX Flow is a light-cure, radio-opaque, flowable restorative material to be used intra-orally and classified as a Type 2 and Class 2 (Group 1) per ISO standard 4049.

This material has a radiopacity equivalent to 2.0-2.5 mm of aluminum oxide.

The particle size of inorganic filers is 0.02-1.3 μm.

The average length of fibers is 140 µm and the diameter is 6 µm.

The total amount of inorganic filler is approximately 50% weight.

COMPOSITION

Base resin, dimethacrylate, glass fiber, initiator, pigment, silicon dioxide, stabilizer

DIRECTIONS FOR USE

1. Preparation

a) Turn the syringe upright and remove the cap by turning counter-clockwise.

b) Remove and secure the tip by attaching the dispensing tip to the syringe by turning clockwise.

c) Place the light protective cap onto the tip.

Note:

Take care to attach the tip to the tip to the tip. This may damage the screw. In order to ensure a tight connection, make sure that the threads are free of residual material.

2. Dispensing shade

Clean the tooth with pulp and water. Select the appropriate shade of everX Flow. Bulk shade or Dentin shade. Refer to Fig. 6. Light-curing for the shade guide of everX Flow is available on the GC Website.

3. Light-curing

Prepare the curing using standard techniques. Dry by gently blowing with air or flame. The use of a rubber dam is recommended to isolate the shade from contamination with saliva, blood or sucralose fluid.

Note:

Shade selection of the final composite layer should be made prior to initiation.

The shade guide of everX Flow is available on the GC Website.

4. Bonding

Prepare the curing using standard techniques. Dry by gently blowing with air or flame. For bonding everX Flow onto dentin, use GC-BOND G-Bond, G-enamel G-BOND or G-BOND (Fig. 2) Follow manufacturer's instructions.

5. Placement

After curing everX Flow – see note for Class 2 and large cavities.

a) Remove the light protective cap from the tip.

b) Prior to initiating the material into the cavity, make sure that the tip is tightly attached to the tip. Check the following:

1) The dispensing tip is securely attached to the tip.

2) There is no residual material at the opening of the tip.

3) A slight amount of residue is visible from the dispensing tip.

4) Place the dispensing tip as close as possible to the cavity edge.

5) Place the dispensing tip as close to the edge as possible to the extra-matrix (Fig. 4). When placing everX Flow in the cavity, ensure that there is enough space for the extra-matrix to bond to the extra-matrix and the surface of the restoration (refer Fig. 7 Placement of the dispensing tip and opening of the composite).

6. Large cavities

Large cavities – refer to note for Class 2 and large cavities.

7. Remove the dispensing tip and remove it from the syringe.

8. After a slight exposure to ambient light, Ambient light Note:

After use, immediately remove and dispose of the dispensing tip and tightly close the syringe with the tip.

9. Cleaning

Light-curing

Light-cure everX Flow using a light curing unit. Keep the light curing tip close to the surface of the restorative (Fig. 6). Refer to the following chart for irradiation Time and Energy required for curing.

Irradiation Time and Energy required for curing

Note:

The effective wavelength range of each dental curing unit must cover 450-480 nm.

Note:

The only Bulk shade can be placed using a bulk application. The Dentin shade should be placed and light-cured in layers as described in the note for shade selection.

10. Lower intensity may cause insufficient curing and discoloration of the shade.

Clinical Hint:

In order to inject everX Flow, use the surface tension of the material to draw it towards you. Do not force the surface tension of the restoration during build up. Once the required amount has been injected, release the pressure on the tip and the dispensing tip will move in a direction perpendicular to the surface. This will allow the material to separate from the dispensing tip and remain on the restoration.

6. Light-curing

Light-cure everX Flow using a light curing unit. Keep the light curing tip close to the surface of the restorative (Fig. 6). Refer to the following chart for irradiation Time and Energy required for curing.

Irradiation Time and Energy required for curing

Note:

The effective wavelength range of each dental curing unit must cover 450-480 nm.

Note:

The only Bulk shade can be placed using a bulk application. The Dentin shade should be placed and light-cured in layers as described in the note for shade selection.

11. Removal of the dispensing composite

Fill the remaining cavity space with a wear-resistant and polished restorative composite, such as G-enamel or Essentia. The composite layer should have a thickness of 1-2 mm on the occlusal surface (Fig. 7).

12. Finishing and Polishing

Check and adjust the occlusion.

Finishing and polish using standard techniques.

SHADE

everX Flow is available in two shades: Bulk shade and Dentin shade.

STORAGE

Recommended for optimal performance, store in a cool and dark place (4-25°C / 39-77°F) away from high temperatures or direct sunlight.

PACKAGES

1. Dispensing tip 0.10 mL x 1, DISPENSING TIP ™ Plastic x 20.

1. Light-protective cap x 1

Dispensing tip reflex x 1

2. DISPENSING TIP ™ Plastic x 20, Light-protective cap x 2

CAUTION

1. Prior to using on clinical tissue or skin, remove immediately with air or pressure if any residual material remains.

2. Wear protective eye glasses during light-curing.

3. Avoid getting material on the skin and wear a mask if there is any chance of inhalation of dust.

4. Do not mix with other materials.

5. Do not use on skin or sensitive areas. To prevent cross-contamination between patients, do not reuse the dispensing tip.

6. Do not use everX Flow in combination with ethyl acetate.

7. Avoid getting material on the skin or sensitive areas with air or pressure if any residual material remains.

8. Some products produced in the present IFU may be classified as hazardous according to GHS. Always familiarise yourself with the safety data sheet of the product.

See the Summary of Safety and Clinical Performance (SSCP) page on the website for further information: <http://www.europeor.com>

They can also be obtained from your supplier.

CLEANING AND DISINFECTION

MULTI-FIBER SYSTEM: To avoid cross-contamination between patients this device requires mid-level disinfection. Immediately after use, please refer to the relevant device and label for disinfection. Discard device if damaged.

DISINFECTANT

The multi-fiber system consists of a dental restorative composite material, which is extremely similar to the dental restorative composites manufactured by the manufacturer.

To prevent cross-contamination between patients, do not reuse the dispensing tip.

DISINFECTANT

Aerosolized spray (Aerosol Disinfectant) such as gloves, face masks and safety eyewear should always be worn.

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Regulatory go.europeor.com

Understands effects - Report if you are experiencing any kind of undesired effect, reaction or similar events experienced by this product, including those not listed in this instruction and label for deactivation. Discard device if damaged.

DO NOT USE

everX Flow is a dental restorative composite, which is designed for the accumulation of contaminants. Discard with a mid-level registered healthcare infection control product according to regional / national guidelines.

For the Summary of Safety and Clinical Performance (SSCP) please see EUADAMED database (<https://ec.europa.eu/tools/euadamed>) or contact us at Regulatory.go@europeor.com

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