

ENGLISH

General Information

3M™ ESPE® Ketac™ Cem Plus Automix is a radiopaque, fluoride-releasing, resin-modified glass ionomer luting cement. It is self-curing with an option for fast light curing of excess cement. **Ketac™ Cem Plus Automix** dental cement consists of a base and catalyst paste packaged in an automix syringe. The automix tip attachment offers convenience since this traditional hand mixed cement system. The cement is available in a white shade. The automix syringe contains 8.5 gm of material.

Indications

Permanent cementation of:
 • Metal and porcelain fused to metal (PFM) crowns and bridges
 • Metal inlays and onlays
 • Crowns made with all-alumina or all-zirconia cores such as 3M™ ESPE® Lava™ or Proceral® AllCeram

• Prefabricated or cast endodontic posts
 • Orthodontic bands and appliances
 • Porcelain fused to metal (PFM), metal-all-alumina or all-zirconia core restorations on implant abutments

Precautionary Information for Patients:
 This product contains substances that may cause an allergic reaction by skin contact in certain individuals. Avoid use of this product in patients with known acrylic allergies, if prolonged contact with oral soft tissue occurs, flush with large amounts of water. If an allergic reaction does occur, discontinue use as needed, remove the product if necessary and consult future use of the product.

This product contains potassium persulfate, which may produce an allergic reaction. Potassium persulfate may trigger an allergic respiratory reaction in certain individuals. This product may not be appropriate for use in those individuals with known sensitivity to sulfites, since a cross-reaction may occur with potassium persulfate.

Precautionary Information for Dental Personnel:
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Wearhinweis für Patienten:
 Dieses Produkt enthält Substanzen, die bei Hautkontakt bei einigen Personen eine allergische Reaktion hervorrufen können. Dieses Produkt nicht bei Patienten mit bekannter Acrylat-Aллерgie verwenden. Bei längerem Kontakt mit oralem Weichgewebe sollte das Produkt mit großem Wasseraufwand abgespült werden. Falls eine allergische Reaktion auftritt, das Produkt aussetzen und die Anwendung beenden.

Dieses Produkt enthält Kaliumpersulfat, das allergische Reaktionen verursachen kann. Kaliumpersulfat kann bei einigen Menschen allergische Atemwegsreaktionen auslösen. Dieses Produkt eignet sich möglicherweise nicht zum Einsatz bei Personen mit bekannter Sulfit-Aллерgie, da es bei diesen Personen zu einer Kreuzreaktion mit Kaliumpersulfat kommen kann.

Warnhinweis für Praxispersonal:
 Dieses Produkt enthält Substanzen, die bei Hautkontakt bei einigen Personen eine allergische Reaktion hervorrufen können. Um das Risiko einer allergischen Reaktion zu reduzieren, minimieren Sie die Kontaktzeit mit diesen Materialien. Insbesondere den Kontakt mit dem nicht ausgehärteten Produkt vermeiden. Bei Hautkontakt die Haut mit Wasser und Seife waschen. Der Gebrauch von Schutzhandschuhen und einer berührungsfreie Technik wird empfohlen, um die Kontaktzeit mit dem Material zu begrenzen. Durchdringen. Wenn das Produkt mit dem Handschuh in Berührung kommt, den Handschuh ausziehen und entsorgen. Hände sofort mit Wasser und Seife waschen und einen neuen Handschuh anziehen. Im Falle einer allergischen Reaktion einen Arzt aufsuchen.

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Instructions for Use

Preparation
 1. Prepare teeth using accepted clinical guidelines.

2. Cover areas in close proximity to the pulp by applying small amounts of hard-setting calcium hydroxide material or resin-modified glass ionomer liner (e.g. 3M™ ESPE® Vitrebond™ Light Cure Glass Ionomer Liner/Base) prior to taking an impression for final seating.

3. Remove the temporary restoration and thoroughly clean the preparation of any temporary cement residue using an oil-free pumice paste.

4. Rinse the prepared tooth or cavity with a water spray, and dry with air, cotton, or a paper tip.

4.1. Leave tooth surface moist. Do not dry.

4.2. The preparation or cavity should be just dry enough that the surface has a slightly glossy appearance. As is the case with any permanent cement, over drying can lead to post-operative sensitivity.

5. Try-in the restoration and check for fit. Adjust if necessary.

6. Sandblast the preparation to be cemented with 30 or 50 micron aluminum oxide at a pressure of 2 Bar (30 psi) to create a matte surface appearance.

7. Thoroughly clean the bonding surface of the restoration.

8. Keep area isolated from blood and saliva contamination during cementation process.

Working and Setting Times

Working Time From Start Of Mixing:	1.5 Minutes	
Excess Cement:		
Tack Light Cure	5 Seconds Per Surface	For Tack Light Curing Use A Conventional Polymerization Device
Self-Curing Gel Phase	2 Minutes After Placement In Mouth	
Self-Curing Setting Time:	5 Minutes After Placement In Mouth	

Directions

1. When Using A New Syringe

1.1. Remove the syringe from the foil pouch and discard foil pouch.
 1.2. Note the date the syringe was removed on the syringe label.
 1.3. Remove the sealing cap from the syringe.
 1.4. Check the syringe openings for blockage. Remove any paste plugs.

1.5. Squeeze out a small quantity of paste onto a pad to equalize the base and catalyst paste in the syringe. Discard the paste which has been squeezed out.

2. During Every Application

2.1. Remove the sealing cap.

2.2. Check the syringe openings for blockage. Remove any paste plugs.

2.3. Attach a new mixing tip and secure it by turning to the right.

2.4. Squeeze out and discard a peppercorn-sized quantity until the first mixed paste in a homogenous color flows out of the tip.

2.4.1. The paste requires a certain amount of time to flow through the mixing tip. The flow through speed cannot be accelerated by increasing the pressure on the plunger.

2.4.2. As soon as the pressure on the plunger decreases, the material flow stops and the paste begins to set up. Do not use force to press out paste that has set as this could damage the syringe.

2.4.3. Apply a thin layer of cement to the inside surface of the restoration. The cement may also be applied directly to the tooth surface for inlay-only restorations.

2.4.5. Keep the opening of the mixing tip immersed in material during the entire application to prevent the inclusion of any air bubbles.

2.4.6. Seal the restoration with firm pressure. Maintain pressure on the restoration to hold proper positioning during cementation.

3. Storing The Unit And Its Dispenser

3.1. For storage, discard the old mixing tip and replace the original sealing cap onto the syringe.

3.2. NOTE: Product must be stored with the sealing cap in place.

4. Clean Up and Finishing

4.1. Excess cement is best removed after brief light exposure (5 seconds per surface with a conventional polymerization device) or during self-curing waxy stage (starting 2 minutes after seating in the mouth). Use a suitable instrument (e.g. scales/explorer) for this process.

4.2. Finish restoration and check occlusion when material has completely set after 5 minutes from placement in the mouth.

Storage and Use

This product is designed to be used at room temperature. If stored in cool conditions, it is advised to reach room temperature prior to use. Shelf life at room temperature is 24 months. Ambient temperatures routinely higher than 27°C/80°F may reduce shelf life. See outer package for expiration date. Keep product in foil package until initial use. Once the foil pouch is opened, the shelf life of the product is 6 months with the original sealing cap.

Do not allow the pastes to dry out. Store with cap securely attached to the automix.

Cleaning and Disinfection

Disinfect the product using an intermediate level disinfection process (liquid contact) as recommended by the Centers for Disease Control (CDC) and endorsed by the American Dental Association.

Centers for Disease Control and Prevention. Summary of Infection Prevention Practices in Dental Settings: Basic Expectations for Safe Care. Atlanta, GA: Centers for Disease Control and Prevention, US Dept of Health and Human Services; October 2016.

References:

CDC Guideline for Disinfection and Sterilization in Healthcare Facilities, 2008. Rutala WA, Weber DJ, Healthcare Infection Control Practices Advisory Committee (HICPAC).

Disposal – See the Safety Data Sheet (available at www.3M.com or through your local subsidiary) for disposal information.

Customer Information

No person is authorized to provide any information which deviates from the information provided in this instruction sheet.

Caution: U.S. Federal Law restricts this device to sale or use on the order of a dental professional.

Warranty

3M ESPE warrants this product will be free from defects in material and manufacture. 3M ESPE MAKES NO OTHER WARRANTIES

INCLUDING ANY IMPLIED WARRANTY OF MERCHANTABILITY OR FITNESS FOR A PARTICULAR PURPOSE. User is responsible for determining the suitability of the product for user's application. If this product is defective within the warranty period, your exclusive remedy and 3M ESPE's sole obligation shall be repair or replacement of the 3M ESPE product.

Limitation of Liability

Except where prohibited by law, 3M ESPE will not be liable for any loss or damage arising from this product, whether direct, indirect, special, incidental or consequential, regardless of the theory asserted, including warranty, contract, negligence or strict liability.

DEUTSCH

Allgemeine Informationen

3M™ ESPE® Ketac™ Cem Plus Automix ist ein röntgenopak, fluoridreleasing, resin-modifizierter glasiger ionomerluting zement. Er ist selbsthärtend und ermöglicht eine einfache Überschusserhöhung durch Lichthärtung. Ketac™ Cem Plus Automix Glasionomerzement wird in einer Automix-Spritze geliefert, die eine Katalysatorpastete enthält. Die Automix-Spritze besteht aus einem automatischen Dosierungssystem, das traditionell von Hand gemischt werden müssen. Der Zement ist in einem weißen Farbton erhältlich. Die Automix-Spritze enthält 8,5 g Material.

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