MR. Dental Supplies Limited
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INSTRUCTIONS FOR USE

USES:
Making temporary crowns, bridges and inlays in the dental laboratory. (This material is not suitable for use in the oral cavity.)

PREPARATION:
The restoration is built-up in wax, "flasked" and "boiled-out" in the usual way.

MIXING:
The powder and liquid are mixed to a creamy consistency and is ready for use when the material drips smoothly off the end of the spatula. Care should be taken when mixing to ensure that no air is trapped.

Pour the correct amount of liquid into a mixing vessel, pour in the powder until all the liquid is absorbed. Tap vigorously on the bench and add more powder until all the liquid is taken up. Thoroughly spatulate for one minute and cover.

MIXING RATIOS:	**1 Part Liquid to 2.5 parts Powder by weight**
DOUGHING TIME:	**AT 68 DEG F or 20 DEG C approximately 25 minutes**
WORKING TIME:	**AT 68 DEG F or 20 DEG C approximately 25 minutes.**

PACKING AND PRESING:
When the material comes cleanly away from the side of the mixing pot it is ready for use and should then be packed into a cold flask. Cellophane or Polythene sheets should be used if a trial closure is preferred, although as the flow is extremely good this should not be necessary.

CURING:
After the restoration has been "built up" it may either be left to cure atmospherically or in a "pressure pot".

ALL OTHER INSTRUCTIONS AS PER SUPERCURE HEATCURE MATERIAL

Liquid Used: Universal Liquid

WARNINGS AND CAUTIONS:

- ***IN EXTREME RARE CASES MAY CAUSE SENTISATION BY SKIN CONTACT***
- ***USE ONLY BY TRAINED PROFESSIONALS***
- ***IT IS THE RESPONSIBILITY OF THE TRAINED PROFESSIONAL TO ENSURE THAT THE PRODUCT IS SUITABLE FOR END USER***
- ***STORE BETWEEN 16C TO 23C AWAY FROM DIRECT SUNLIGHT***
- ***DISPOSABLE OF UNCURED LIQUID MUST BE CARRIED OUT IN ACCORDANCE WITH MATERIAL SAFETY DATA SHEET***
- ***AND MIXED POWDER AND LIQUID CAUSES EXOTHERMIC REACTION***
- ***IF THE DEVICE HAS BEEN PLACED IN THE ORAL CAVITY THEN IT MUST BE DISPOSED OF AS POTENTIALLY BIOLOGICAL HAZARD WASTE***
- ***THE USER AND/OR THE PATIENT MUST REPORT ANY SERIOUS INCIDENT THAT HAS OCCURRED IN RELATION TO THE DEVICE TO THE MANUFACTURER AND THE COMPETENT AUTHORITY OF THE MEMBER STATE IN WHICH THE USER AND/OR PATIENT IS ESTABLISHED***

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