

CUSTOM-MADE DENTAL APPLIANCE PRESCRIPTION

Please complete the appropriate sections of this prescription and return to the address opposite.

8 VICTORIA COURT, BANK SQUARE, MORLEY, LEEDS, LS27 9SE TEL: 07471447391

	LAB REFERENCE	OUTLINE OF DESIGN REQUIRED	PHOTO Y/N
PRESCRIBING CLINICIAN		0.7.5.5.4.2.2.4.5.6.7.0	
		8 7 6 5 4 3 2 1 1 2 3 4 5 6 7 8	
Name:		8 7 6 5 4 3 2 1 1 2 3 4 5 6 7 8	DEVICE(S) REQUIRED
		l	
Address:		INSTRUCTIONS AND AMENDMENTS REC	ORD
PATIENT/ID	Age		
Name:			
	Sex		
ADDOUNTAGENTS			
APPOINTMENTS SHAD)E		
Date:			
Time: AM / PM			
Date:			
Time: AM / PM			
Date:			
Time: AM / PM			
FIELDS BELOW TO BE COMPLETED BY LABORATORY PERSONNEL ONLY			
Approved for manufacture by:		Approved for release by:	Invoice Date:
Sign:		Sign:	Invoice Number:
-		-	. ,2
Details of metaviole steering ind hy muses in a		Data ils of any model annual by massails a	
Details of materials etc supplied by prescribe	ſ	Details of any model approval by prescriber	
Initials:		Initials:	
Your attention is drawn to the following statement: This is a custom-made medical device that has been manufactured to satisfy the design characteristics and			

properties specified by the prescriber for the above-named patient. This medical device that has been manufactured to satisfy the design characteristics and properties specified by the prescriber for the above-named patient. This medical device is intended for exclusive use by this patient and conforms to the general safety and performance requirements specified in Annex I of the Medical Devices Regulations.

This statement does not apply to medical devices that have been repaired and/or refurbished for an individual patient's use.

Storing, handling and instructions for use: It is recommended that before use, this medical device is stored in a clean and safe environment that prevents it from coming into contact with materials, equipment, acids or bleaches that could cause physical or chemical damage to the medical device. The medical device should not be subjected to extremes of temperature during storage. Where applicable, you should take care not to damage the medical device when removing it from its model.

ORIGIN OF MANUFACTURE DECLARATION

This complete appliance has been wholly manufactured within the EU.

PRESCRIBER FEEDBACK:

To enable our dental laboratory to comply with the Medical Devices Regulations for Post Market Surveillance, please inform us of any feedback or issues regarding the enclosed device(s) as soon as possible.

MHRA No: 9645