Armour House Colthrop Lane Thatcham West Berkshire RG19 4NT

Company Number: 13656153





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marolalabs.com

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hello@marolalabs.com

PRESCRIBING SURGEON'S E	DETAILS	ITEMS ENCLOSE)						
Name:			Date /	Date _/_	Date /_				
Surgery:		Models				C.	000	D_	5
		Rubber IMP						Al.	
PATIENT'S DETAILS		Silicone IMP					Hov	204	
PATIENT 3 DETAILS		Alignate IMP					Upp	jer	
Patient's name:		Bite register				(1)			
Gender:	Age:	Wax set up							(F
PLEASE TICK AS APPROPRIATE	RETURN DATE (NOT FIT DATE)	Photographs					Low	/or	
Chrome	Repair//	Chrome					LOV	761	
Acrylic	Special tray//	Acrylic denture				(2)			NO.
Duraflex Valplast	Bite block//	Flexible denture					3010	Y PC	,
Re-line / Repair	Chrome// Try-in/_/	Other							
Bleaching trays	Re-try / /	Your attention is di	awn to t	he follow	ving state	ement:			
Sports gaurd	Finish //	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.							
Bite raising appliance	VITA GUM/TEETH	This medical device applicable general : Medical Devices Re	safety an	d perforn					
Essix retainer	SHADE MOULD CHARACTERISATION	This statement does not apply to medical devices that have been repaired and/or refurbished for an individual patient's use.							d/or
Other	YES NO								

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.

NOTES / INSTRUCTIONS



This complete appliance has been wholly manufactured within the UK $\&\, \text{EU}$

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.







