K IMPLANT REATORAT	NOI	Case ID:	
accurate + art + aestheric	Impression Date:	Date Required: 1 working day before actual appointment	We Accept All Digital
AAA DENTALAB 1st Floor, Unit 4 Watling Gate, 297-303 Edgware Road, London NW9 6NB. www.aaadentalab.co.uk : infouk@aaadentalab.com	Dentist Name:		Scan
	Practice Name :		
	Postcode (Acc No.):		
	Patient ID / Code :		
©: 0203 6022728 : 07400066241	☐ Male ☐ Female	D.O.B.:	
Restoration Type Cement Retained Screw Retained Single Unit Crown (s)	8 7 6 5 4 3 2 8 7 6 5 4 3 2	 	6 7 8 6 7 8
Multy Unit Bridge Implant Abutment(s)	0 7 0 5 4 5 2	L L	
	cial Instructions:		Shade
Components Enclosed Abutment			Vita Classic
Porcelain Fused Metal Non-Precious Semi-Precious Precious			
Metal Free Ceramics ☐ Composite ☐ Full Zirconia ☐ Zirconia with Emax® Layering ☐ Emax® Press			
If Insufficient Room Metal Occlusion Adjust Opposing			
I have disinfected the impression : With :			

THIS DEVICE IS NON STERILE

(Full Terms & Conditions are attched with our pricelist or visit our web at www.aaadentalab.co.uk) MHRA No. 8496/GDC No. 285519 / DLA MEMBER Your attention is drawn to the following statement: This is a custom-made dental device that has been manufactured to satisfy the design characteristics, properties and detail of what has been prescribed by the above dentist. This dental device is purely manufactured for exclusive use of the patient described above. The product meets the general safety and performance requirements specified in the Annex I of Medical Devices Regulations. This product may have been produced in either or both UK and China by AAA Destalab LTD.

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Instructions for use, storing & handling: It is highly recommended that, this dental device is stored in a clean and safe environment that prevents it from coming in contact with materials, liquids or acids that could cause disfiguration or damage to the dental device. The dental device should not be subjected to extreme heat. Where applicable, you should take care not to damage the dental device(s) when removing from its model.

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.