## **Prosthetics Lab Ticket**

**BARCODE - LECA OFFICE USE ONLY** 

5 Watt Road, Hillington Park, Glasgow G52 4RY

DATE REQ'D

Phone: 0141 883 6111
Fax: 0141 883 3574
Email: info@lecadental.co

Dental Laboratories Association Registered Member

LECA
an <b>ALS</b> company

				Email: info@lecadental.com Web: www.lecadental.com		
			NH			
				ΤΕ		
DENTIST:						
Surgery:						
Tel:						
PATIENT NAME/ ID:						
Age: Male	J Fem	ale 🖵				
DEVICE REQUIRED						
ACRYLIC		U	L			
FULL		U	L			
PARTIAL		U	L			
VALPLAST U		U	L			
ORTHODONTIC		U	L	EXPRESS SERVICE REQUIRED YES NO		
MAKE OF TEETH SHADE MOULD		NOULD	TEETH TO BE EXTRACTED			
REQUIREMENTS		Delive	ry Date	R L		
Special Trays U 🗆 L 🗖				TEETH REQUIRED ON DENTURE		
Bites U 🗆 L 🗅						
Try-In				R ———— L		
Re-Try						
Finish 🗖						
Finish With: Hi-impact U L Gum Staining U L   Violage V L Ivobase V L Violage V Violage						
DISINFECTED IN SURGERY						
Print Name & Date						
Signature C C C C C						
Approved for Manufactu by GDC Technician	re		FOR LABORATORY US	E ONLY - QUALITY CONTROL		

Your attention is drawn to the following: 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 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 of 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.

PATIENT FEEDBACK

To enable our dental laboratory to comply with the Medical Devices Regulations for Post Market Surveillance patients should direct any queries regarding the fit or performance of this appliance to the prescribing dentist.

PRESCRIBER/DENTIST 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.