

Crown & Bridge Lab Ticket

MHRA: 2949

5 Watt Road,
Hillington Park,
Glasgow G52 4RY

 Dental Laboratories Association
Registered Member

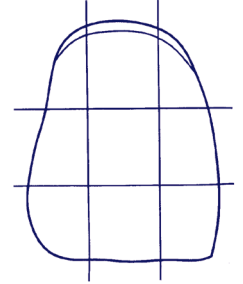


LECA
an **ALS** company

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BARCODE - LECA OFFICE USE ONLY	DATE REQ'D
	NHS
	PRIVATE

SPECIAL INSTRUCTIONS



DENTIST:

Surgery:

Tel:

PATIENT NAME/ ID:

Age: Male Female

PREPARED TOOTH COLOUR (STUMP SHADE)	SHADE REQUIRED
18 17 16 15 14 13 12 11	21 22 23 24 25 26 27 28
48 47 46 45 44 43 42 41	31 32 33 34 35 36 37 38

MATERIALS PROVIDED

BITE REG PARAPOST

ALIGNATE RUBBER BASE

STUDY MODELS PHOTOS

FACEBOW REG OTHER

EXPRESS SERVICE REQUIRED YES NO

NON PRECIOUS POST & CORE

PRECIOUS POST & CORE (ALLOY EXTRA)

NP SHELL

SILVER GOLD

40% PRECIOUS SHELL (PALE GOLD)

60% GOLD INLAY/ONLAY/SHELL (ALLOY EXTRA)

PRECIOUS BONDED CROWN (ALLOY EXTRA)

PRECIOUS BONDED BRIDGE (ALLOY EXTRA)

NON PRECIOUS BONDED CROWN

NON PRECIOUS BRIDGE

MARYLAND BRIDGE

DIAGNOSTIC WAX UP

ALL CERAMIC RESTORATIONS

FULL CONTOUR ZIRCONIA CROWN

ZIRCONIA CROWN

ZIRCONIA BRIDGE

IPS E.MAX CROWN

IPS E.MAX BRIDGE

IPS E.MAX VENEER

IPS E.MAX INLAY

IPS E.MAX ZIRPRESS CROWN (HIGH STRENGTH)

IPS E.MAX ZIRPRESS BRIDGE (HIGH STRENGTH)

PORCELAIN INLAY

FELDSPATIC VENEER

FOR LABORATORY USE ONLY - QUALITY CONTROL

DATE	Approved for Manufacture by GDC Technician						

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.