

JOB NO.

MHRA No. 24892



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PRESCRIBER _____

SURGERY _____

PATIENT _____

DATE _____

SPECIAL TRAYS ↑ ↓ **Date Required** _____

BITE BLOCKS ↑ ↓ **Date Required** _____

F/ /F P/ /P

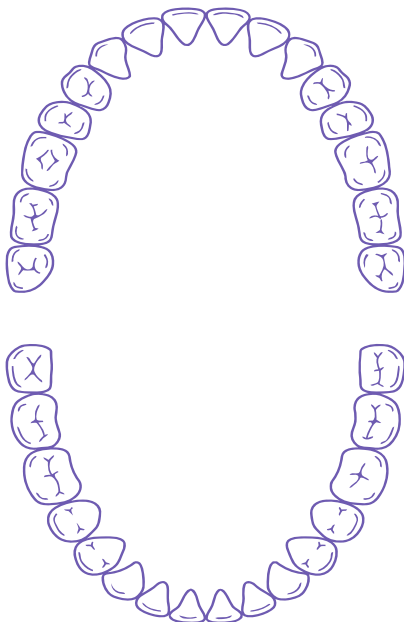
TRY-IN _____ **Date Required** _____

RE-TRY _____ **Date Required** _____

FINISH _____ **Date Required** _____

SHADE _____

APPROVED FOR MANUFACTURE



R _____ **L**

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.

OTHER INSTRUCTIONS

DENTURE TYPES

ACRYLIC

CHROME

FLEXI

HYBRID

DURACETAL

RESIN PRINTED

KARADENT

iFLEX

CLASPS

STEEL

FLEXI

AESTHETIC

EXTRA FEATURES

GUM CONTOURING

CLEAR PALATE

FIBER FORCE

OTHER

RELINE

SOFT LINING

ADDITION

REPAIR

STUDY MODEL ↑ ↓

ESSIX RETAINER ↑ ↓

NIGHT APPLIANCE ↑ ↓

SOFT SEMI-RIGID

BLEACHING TRAY ↑ ↓

RESERVOIRS Y N

MOUTH GUARD

COLOUR(S) _____

THICKNESS: LT / MED

MED / HVY

SPORT _____

DATE REQUIRED _____ **APPROVED FOR MANUFACTURE**

This device has been approved for release by a GDC registered technician. It is supplied in a non-sterile state.

SIGNED FOR RELEASE _____ **£** _____

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 is intended for exclusive use by this patient and conforms to the applicable general safety and performance requirements specified in the UK & EU Medical Devices Regulations. This statement does not apply to medical devices that have been repaired and/or refurbished for an individual patient's use.