

LAB USE ONLY	
Date Received	_____
Job No.	_____
Job Type	_____
Material	_____





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PROSTHETICS

SPECIFICATIONS

CLIENT DETAILS

NAME

SURGERY NAME/AREA

PATIENT DETAILS

NAME/ID

AGE

	DATE REQUIRED	IN SURGERY DECON BY	INDEPENDENT <input type="checkbox"/>	PRIVATE <input type="checkbox"/>	NHS <input type="checkbox"/>
SPECIAL TRAYS			<div>UPPER</div> <div> Full Partial Acrylic Chrome Flexi Splint </div> <div> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> </div> <div>LOWER</div> <div> Full Partial Acrylic Chrome Flexi Splint </div> <div> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> <input type="text"/> </div>		
BITE					
TRY					
RE-TRY					
FINISH					

Mould	SHADE	

AMENDMENTS - LAB USE ONLY	SCAN INFORMATION DIGITAL SCAN SENT <input type="checkbox"/>
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<div>Subcontracted Work Inspection Record</div> <div> Satisfactory YES NO </div> <div> Inspected By: Date </div> <div> Details of unsatisfactory subcontracted work and corrective action taken </div>	ENCLOSURES	<table border="1"> <tr><td>Imps</td><td></td></tr> <tr><td>SP Trays</td><td></td></tr> <tr><td>Final Imp</td><td></td></tr> <tr><td>Reg Block</td><td></td></tr> <tr><td>Try In</td><td></td></tr> <tr><td>Re-Try</td><td></td></tr> <tr><td>Finish</td><td></td></tr> </table>	Imps		SP Trays		Final Imp		Reg Block		Try In		Re-Try		Finish		<div>ORIGIN OF MANUFACTURE DECLARATION</div> <div> This complete appliance has been wholly manufactured within the EU </div> <div> YES <input checked="" type="checkbox"/> NO <input type="checkbox"/> </div>
	Imps																
SP Trays																	
Final Imp																	
Reg Block																	
Try In																	
Re-Try																	
Finish																	
<div>APPROVAL FOR MANUFACTURE</div> <div>DATE</div>	<div> RUBBER UPPER RUBBER LOWER ALG UPPER ALG LOWER BITE PHOTO JIG/FACEBOW OTHER </div>																

*Detailed in amendments

Your attention Is drawn to the following statement: This is a custom-made medical device that has been manufactured to satisfy the design characteristics the prescriber for the above-named patient. This medical device is intended for exclusive use by this patient and conforms to general safety and performance requirements spcified in Annex 1 of the Medical Devices Regulations.

This Statement does not apply to medical devices that have been repaired and or refurbished for a patient's use.

This Device is returned non sterile. Do not expose appliance to extemes of temperatures

Prescriber Feedback

To enable our Dental Laboratory to comply with the Medical Devices Regulation for Post Market Surveillance, please of any feedback or issues the enclosed device (s) as soon as possible.

MHRA N0. 1375