

## Laboratory Ticket & Patient Prescription

Please complete & tick relevant boxes/section

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## Registered with the UK competent authority CA009282

If you have any queries regarding			Study models				
	cribing dentist for further inj	ormation	Bite registration rim	Upper	Lower		
Client			Special tray	Upper Upper	Lower	Perforated	Non Perforated
Address - Please print clearly			Chrome Cobalt				
			Complete	Upper	Lower		
			Partial plate	Upper	Lower		
			Skeleton multi bar	Upper	Lower		
			Skeleton single bar	Upper	Lower		
			Acetal resin clasp	Upper	Lower		
Telephone No			Drosthotics			Y DISINFECTED, P	DI FASE TICK ROX
			<u>Prosthetics</u>			POINTMENT DA	
Patient: - Name	Age	M/F	Bite registration		Date	/	Time
			Try-in of metalwork or	nly	Date	/	Time
ob No Shade (	Guide Used Sh	ade Required	Try-in of teeth - metal	work	Date	/	Time
			Try-in of teeth - acryli	c denture	Date	/	Time
	1 =		Process & Finish		Date	/	Time
Contract: All Inclusive:	Staged:		Re-try of teeth		Date	//	Time
nano edp	edp		Re-try and process		Date	//	Time
•			Re-line acrylic		Date	//	Time
idp	Simplex		Soft Lining		Date	<u>//</u>	Time
Florible montial deutima	Implement Page	in a d	Pressure Thermo fo	rming			
Flexible partial denture	Implant Reta	inea —	■ Base plate		Whiten	ing/Fluoride/Me	edical
Design instruction/inforn	nation		Sports mouth guar	rd	Anti-Sn	oring Device	
			Splints/Retainers/	Stents			
	LOWER		Special Instruction (		INT CLEARLY	)	
R L UPPER	R	L (S)					
Lab use only  Imps sent							
☐ Models sent							
☐ Bite sent							
☐ Chrome sent ☐ Try-in sent							
☐ Denture sent							
☐ Other							
Approved for manufacture by	•						
(sign)	(date)						
Approved for release by:							
(sign)	(date)						_
	ANUFACTURE DECLARATION						Request a call
This complete appliance has be Yes No (If no, detail manu	peen wholly manufactured was facturing locations Below)	rithin the EU.	<b>Ø</b> GDC	protecting properties of the protecting prot	patients, ne dental team s	MHRA afeguarding Public Health MDD No: CA009282	
— 140 (11 110, detail fillallu	idetaining locations below)			A11			

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 relevant essential requirements specified in Annex 1 of the Medical Devices Directive and the United Kingdom 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 for 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, alkalies 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. Where applicable, instructions on how to use or clean this medical device may be obtained from the prescriber.