

### Dentist and Patient

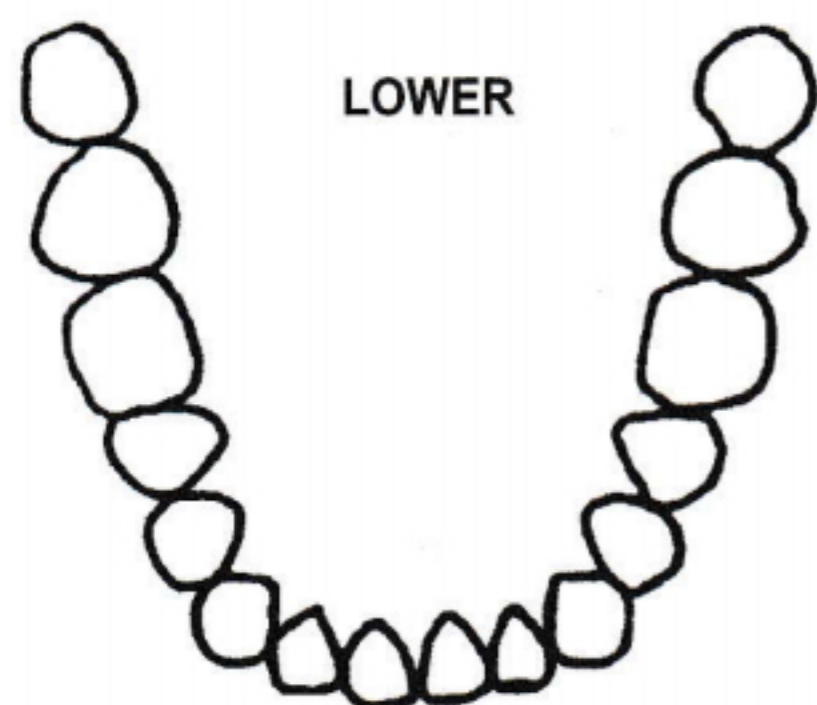
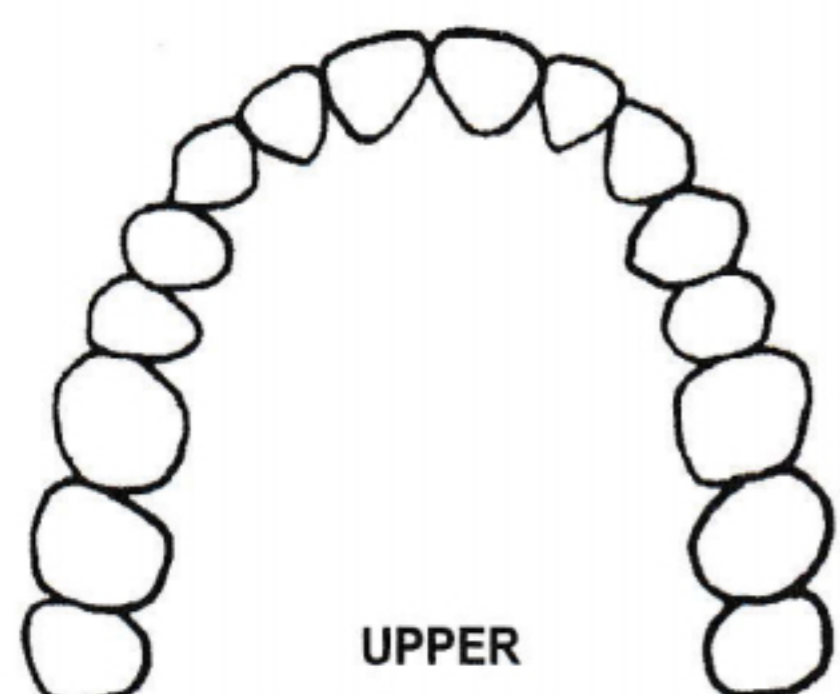
Prescriber \_\_\_\_\_ Patient Name/ID: \_\_\_\_\_  
 Address \_\_\_\_\_ Male / Female \_\_\_\_\_  
 Telephone No \_\_\_\_\_ Age \_\_\_\_\_ Date \_\_\_\_\_

### Case Information

						NHS	Health Plan	Private	Job No (Lab Use)
Type of Appliance Required (please tick below)									
ACRYLIC	CHROME	ORTHO	BLEACH TRAY	SPLINT	MOUTHGUARD	OTHER	MOULD		
DATE REQUIRED		DISINFECTED BY / DATE		APPROVED BY (Lab Use)		COST (Lab Use)		SHADE	
SPECIAL TRAY								LAB USE ONLY	
BITE								IMPS RECD	
CHROME								MODS RECD	
TRY-IN								BITE/FACE BOW	
RE TRY									
FINISH				TOTAL		£			

### Case Instructions

**\*PLEASE DO NOT PUT PRESCRIPTION IN DIRECT CONTACT WITH IMPRESSIONS**  
**MEDICAL DEVICE IS SUPPLIED IN AN UNSTERILISED STATE**



APPROVED FOR MANUFACTURE BY	APPROVED FOR RELEASE BY
DATE	DATE



**Your attention is drawn to the following statement:** This is a custom-made dental appliance 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 1 of the Medical Devices Regulations.

This statement does not apply to Medical Devices that have been repaired and/refurbished for an individual patients 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 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 Medical Device has been wholly manufactured within the EU.

**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.