



Costech Dental Laboratory

Unit 17-18, Lion Business Park, Dering Way,
Gravesend, Kent, DA12 2DN
email: dentist@costech.co.uk



Dentist Name: _____

Surgery Address: _____

Patient Name: _____

Male

Female

Vita Shade

Job Number:

Date Started: _____

- | | | |
|--------------------------|--------------------------|---------------|
| Upper | Lower | |
| <input type="checkbox"/> | <input type="checkbox"/> | Acrylic |
| <input type="checkbox"/> | <input type="checkbox"/> | Chrome |
| <input type="checkbox"/> | <input type="checkbox"/> | Full |
| <input type="checkbox"/> | <input type="checkbox"/> | 8 + Partial |
| <input type="checkbox"/> | <input type="checkbox"/> | 4 - 7 Partial |
| <input type="checkbox"/> | <input type="checkbox"/> | 1 - 3 Partial |
| <input type="checkbox"/> | <input type="checkbox"/> | Reline |
| <input type="checkbox"/> | <input type="checkbox"/> | Repair |
| <input type="checkbox"/> | <input type="checkbox"/> | Additions |

Due Dates

Special Tray: U L

Bite Block:

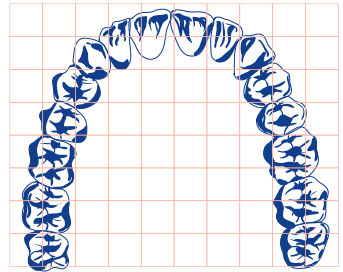
Try-In:

Re-Try:

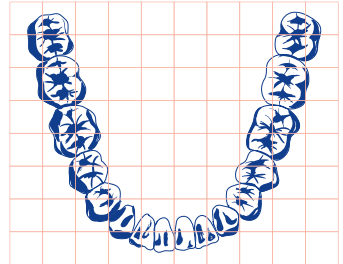
Finish:

Please circle teeth to be included

8	7	6	5	4	3	2	1		1	2	3	4	5	6	7	8
8	7	6	5	4	3	2	1		1	2	3	4	5	6	7	8



Design



Notes:

CUSTOM MADE DEVICE Certified as conforming to the Medical Device Directive, manufactured and supplied by Costech Elite.

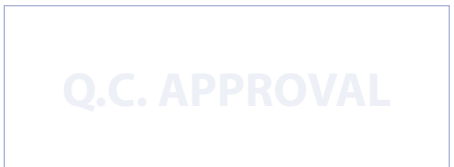
The product here packed is a custom made device for the named patient stated above and conforms to the essential requirements set out in Annex 1 of the EC Medical Device Directive 93/42/EEC dated June 1993 and if any of these requirements are not fully met the details are documented on reverse or attached and despatched to the user.

In the event that the prescriber has supplied some of the materials etc for incorporation in a particular custom made dental appliance then this appliance cannot be guaranteed to fully meet with the applicable relevant essential requirements.

** The Dr named above takes responsibility for prescribing the correct material/ alloy in accordance to current regulations in their region.*

For your patient statement please go to:
www.costech.co.uk/patientstatement

The grounds for placing such a device on the market is that the risk of compromising the patient's health and safety by using materials etc supplied by the prescriber is assessed as minimal. This risk assessment relies upon a duly qualified medical practitioner's competence and duty to supply materials etc that is either from a CE marked source or from an appropriate European Competent Authority registered manufacturer of custom-made medical devices.



Please note these items are NOT sterile. MHRA Registration: 5759