

## Costech Elite Dental Laboratory

Unit 18, Lion Business Park, Dering Way,  
Gravesend, Kent, DA12 2DN  
T: 01474 320076 www.costech.co.uk



Dentist Name: \_\_\_\_\_

Surgery Address: \_\_\_\_\_

Patient Name: \_\_\_\_\_

Male  Female  Age  Y / M / O

Vita Shade:

Job Number:

Box Number: \_\_\_\_\_

Date Prepared: \_\_\_\_\_

- | Upper                    | Lower                    |              |
|--------------------------|--------------------------|--------------|
| <input type="checkbox"/> | <input type="checkbox"/> | Acrylic      |
| <input type="checkbox"/> | <input type="checkbox"/> | Chrome       |
| <input type="checkbox"/> | <input type="checkbox"/> | ComFlexin®   |
| <input type="checkbox"/> | <input type="checkbox"/> | NaturElite®  |
| <input type="checkbox"/> | <input type="checkbox"/> | GingiFlex™   |
| <input type="checkbox"/> | <input type="checkbox"/> | Special Tray |

Due Dates

Special Tray:  U  L .....

Bite Block: .....

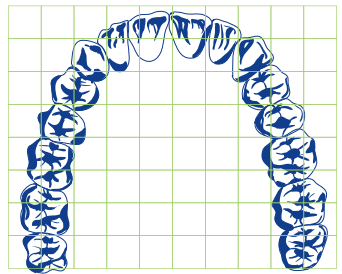
Try-In: .....

Re-Try: .....

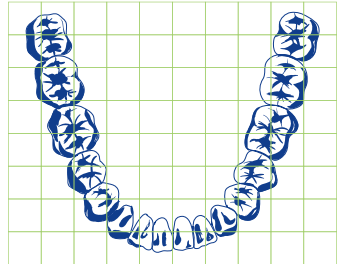
Finish: .....

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 Exclusively.** 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.

**Please note these items are NOT sterile. MHRA: CA000911**

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 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.

Signed ..... Position ..... Date .....