



**Dentist Account Details:**

# Prosthetic

## Patient & job Details

Patient's Name: \_\_\_\_\_

Job ID: \_\_\_\_\_ Date Started: \_\_\_\_\_

Male

Chrome

UPPER LOWER

Reline

UPPER LOWER

Additions

UPPER LOWER

Repair

UPPER LOWER

Acrylic

UPPER LOWER

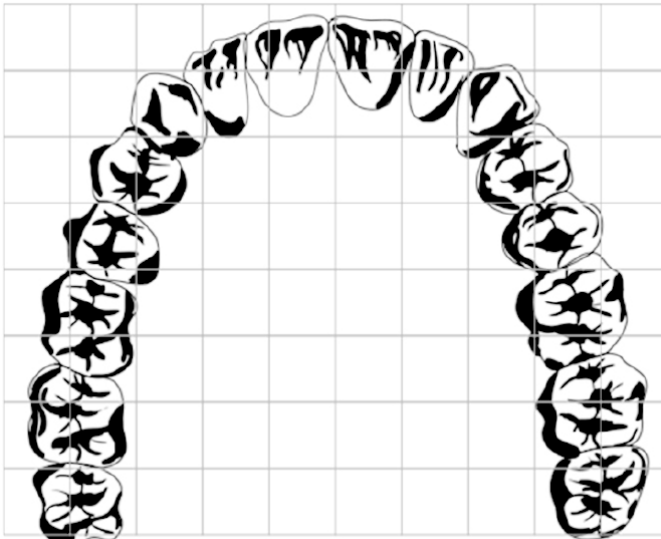
ComFlexin®

UPPER LOWER

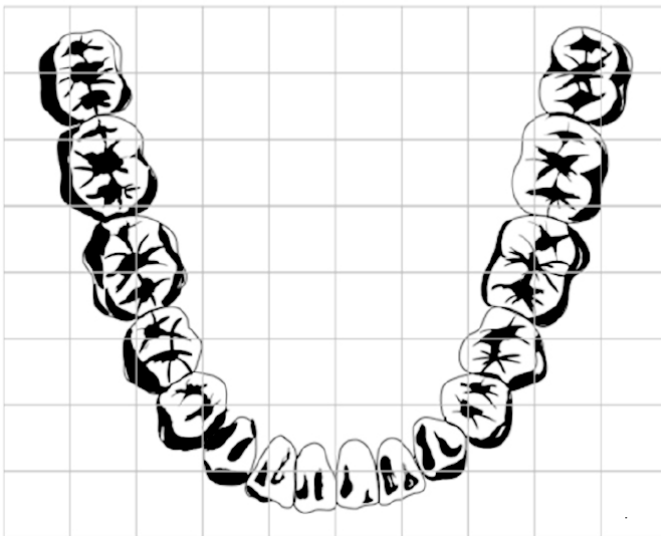
Female

**Vita Shade:**

**Please circle teeth to be included**



Design



**Due Dates**

Special Tray: \_\_\_\_\_  U  L

Bite Block: \_\_\_\_\_

Try-In: \_\_\_\_\_

Re-Try: \_\_\_\_\_

Finish: \_\_\_\_\_

**Notes**

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**CUSTOM MADE DEVICE Certified as conforming to the Medical Device Directive, manufactured and supplied by Royal Dental lab.**

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 I 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.

For full terms and conditions please see our website on [www.royal-dental.co.uk](http://www.royal-dental.co.uk)

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. **6 months Warranty**

Q.C APPROVAL

Please note these items are NOT sterile. MHRA Registration: 8999