



User Guidelines for





# **General Information**

### Storage

DEGRAD INX<sup>©</sup> X100 should be stored at 4 °C. Protect it from light. Expiry dates of the kit components are indicated on the vials. The products can be stored for a maximum of 3 months after opening and should be consumed before the expiry date. Always re-seal the resin vial with parafilm after use.

### **Intended Use**

Research use only. This product is not intended for use in diagnostic or therapeutic procedures.

### **Safety Information**

Work in a ventilated area and use suitable personal protective equipment. For more information, please refer to the safety data sheet.

## **User Guidelines**

### **Preparation**

- 1. Warm up the resin vial by placing it into a pre-heated water bath (50 °C) using the supplied floater for 10 min.
- 2. Open the screw cap of the resin vial and gently pipette back and forth.
- **3.** Draw ± 0.1 ml resin and gently pipette onto a pre-silanized<sup>b</sup> glass substrate. Avoid air bubbles.
- 4. Place the substrate onto a heated plate (70 °C) for 20 min for the removal of sacrificial solvent.
- **5.** After heating, the resin is ready for printing process.

#### **Processing**



DEGRAD INX X100 is only suitable for conventional printing methods.



1 The printing process must be completed within 6h after the resin is placed in the printer. The resin is not suitable for longer processing times.

<sup>&</sup>lt;sup>a</sup> After use, promptly re-close the vial with the screw cap and seal with parafilm.

**b** Recommended silanization protocol: Immerse the glass substrates in 3-(TrimethoxysilyI) propyl methacrylate (CAS: 2530-85-0) solution (1 v/v % in ethanol) for 45 min. Rinse thoroughly in ethanol and dry via a lens blower.



The recommended processing parameters for 10x/0.4NA objective:

Pulse Duration	90 fs
Repetition Rate	80 MHz
Center Wavelength	780 nm
Hatching	0.5 μm
Layer Spacing	5 μm
Writing Speed	600 mm/s
Average Laser Power	> 70 mW

## **Developing**

If present, wipe off the large part of the immersion oil from the bottom of the slide, clean remaining oil residue with a tissue and isopropanol.

Put the sample in a beaker filled with the supplied developer until full dissolution.



Work in a fume hood with ventilation and use suitable personal protective equipment.



Cover the beaker to prevent evaporation of the developer.

After developing, the sample can be stored at room temperature.

### **Imaging**

The printed samples can be imaged using a scanning electron microscope or an inverted microscope using an excitation wavelength of 488 nm and a fluorescence emission wavelength of 507 nm.