## Instructions μ-Slide 8 Well



The ibidi product family is comprised of a variety of  $\mu$ –Slides and  $\mu$ –Dishes, which have all been designed for high–end microscopic analysis of fixed or living cells. The high optical quality of the material is similar to that of glass, so you can perform all kinds of fluorescence experiments with uncompromised resolution and choice of wavelength. The  $\mu$ –Slide 8 Well is an array of 8

square fields where cells can be cultivated and, subsequently, investigated with microscopical methods. This open µ–Slide (chambered coverslip) with 8 wells is intended for immunofluorescence, live cell imaging, and high-end microscopy.

#### **Material**

ibidi  $\mu\text{-Slides}$ ,  $\mu\text{-Dishes}$ , and  $\mu\text{-Plates}$  are made of a plastic that has the highest optical quality. The polymer coverslip on the bottom exhibits extremely low birefringence and autofluorescence, similar to that of glass. Also, it is not possible to detach the bottom from the upper part. The  $\mu\text{-Slides}$ ,  $\mu\text{-Dishes}$ , and  $\mu\text{-Plates}$  are not autoclavable, since they are only temperature–stable up to  $80^{\circ}\text{C}/175^{\circ}\text{F}$ . Please note that gas exchange between the medium and incubator's atmosphere occurs partially through the polymer coverslip, which should not be covered.

Optical Properties ibidi Polymer Covers		
Refractive index n <sub>D</sub> (589 nm)	1.52	

Thickness No. 1.5 (180 μm)

Material polymer coverslip

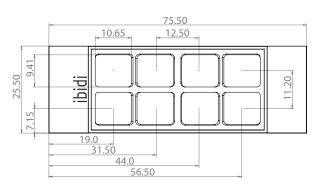
Please note! The ibidi Polymer Coverslip is compatible with certain types of immersion oil only. A list of suitable oils can be found on page 3.

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### Geometry

Abbe number

The  $\mu$ -Slide 8 Well provides a standard slide format according to ISO 8037/1.



Geometry of µ-Slide 8 Well		
Outer dimensions in mm $(w \times l)$	25.5×75.5	
Number of wells	8	
Dimensions of wells in mm $(w \times l \times h)$	$9.4 \times 10.7 \times 6.8$	
Volume per well	$300\mu l$	
Total height with lid	8 mm	
Growth area per well	$1.0 \text{ cm}^2$	
Coating area per well	$2.2\mathrm{cm}^2$	
Bottom	ibidi Polymer Coverslip	

### **Shipping and Storage**

The  $\mu$ -Slides,  $\mu$ -Dishes and  $\mu$ -Plates are sterilized and welded in a gas-permeable packaging. The shelf life under proper storage conditions (in a dry place, no direct sunlight) is listed in the following table.

Conditions		
Shipping conditions	Ambient	
Storage conditions	RT (15-25°C)	
		_
Shelf Life		
ibiTreat, Uncoated	36 months	
Collagen IV, Poly-L-Lysine	18 months	

#### **Surface**

The tissue culture treated ibiTreat surface is a physical surface modification and optimized for adhesion of most cell types. The uncoated surface is a very hydrophobic surface and allows no direct cell growth. It is suitable for specific coatings or suspension cells.

If you like to establish a particular coating for your demands we recommend testing your coating procedure on uncoated and ibiTreat surfaces, since some proteins and

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biomolecules adhere differently to hydrophobic or hydrophilic polymer surfaces.

### Coating

Specific coatings are possible following this protocol:

- 1. Prepare your coating solution according to the manufacturer's specifications or reference.
- 2. Apply 300 µl and leave at room temperature for at least 30 minutes.
- 3. Aspirate the solution and wash with the recommended protein dilution buffer.
- 4. Optionally let dry at room temperature. Attention, some coating proteins might degenerate when drying!

Detailed information about coatings is provided in Application Note 08 Cell culture coating.

### **Seeding Cells**

- Trypsinize and count cells as usual. Dilute the cell suspension to the desired concentration. Depending on your cell type, application of a  $5-11 \times 10^4$  cells/ml suspension should result in a confluent layer within 2–3 days.
- Apply 300 µl cell suspension into each well of the slide. Avoid shaking as this will result in inhomogeneous distribution of the cells.

• Cover the slide with the supplied lid. Incubate at 37°C and 5 % CO<sub>2</sub> as usual.

Undemanding cells can be left in their seeding medium for up to three days and grow to confluence there. However, best results might be achieved when the medium is changed every 1–2 days. Carefully aspirate the old medium and replace it by 300 µl fresh medium per well.

### Tip:

As you may know from 96 well plates, the bent meniscus at the air–liquid interphase in small open wells destroys the phase contrast effect of your microscope image. To avoid this problem, we recommend using our channel Slides such as the  $\mu$ –Slides I Luer and  $\mu$ –Slide VI  $^{0.4}$  or a Ph+ Slide.

### **Cell Microscopy and Solvents for Fixation**

To analyze your cells, no special preparations are necessary. Cells can be observed live, or fixed directly in the  $\mu\text{-Slide 8}$  Well, preferably on an inverted microscope. Due to the thin bottom, high resolution microscopy is possible. The material is compatible to most fixatives, like acidic acid, alcohols and PFA. For a full list of compatible solvents and more information on chemical compatibility, please visit the FAQ section on www.ibidi.com. For optimal results in fluorescence microscopy and storage of stained probes ibidi provides a mounting medium (50001) optimized for  $\mu\text{-Dishes}$  and  $\mu\text{-Slides}.$ 



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#### **Immersion Oil**

When using oil immersion objectives with the ibidi Polymer Coverslip, use only the immersion oils specified in the table below. The use of any non–recommended oil could damage the ibidi Polymer Coverslip. The resulting leakage may harm objectives and microscope components. All immersion oils that are not listed in the table below should be considered as non–compatible.

Company	Product	Ordering No.	Lot Number	Test Date
ibidi	ibidi Immersion Oil	50101	16-12-27	01/2017
Zeiss	Immersol 518 F	444960	160706	01/2017
Zeiss	Immersol W 2010	444969	101122	04/2012
Leica	Immersion Liquid	11513859	n.a.	03/2011
Cargille	Type A	16482	100592	01/2017
Cargille	Type HF	16245	92192	01/2017
Olympus	Silicone Immersion Oil	SIL300CS-30CC	N4190800	01/2017
Carl Roth	Immersion oil	X899.1	414220338	01/2017



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### **Ordering Information**

The  $\mu$ –Slide 8 Well family comprises Slides with different surfaces and bottom characteristics. See table below for choosing your  $\mu$ –Slide 8 Well.

### μ-Slide 8 Well



Cat. No.	Description
80826	μ–Slide 8 Well ibiTreat: #1.5 polymer coverslip, tissue culture treated, sterilized
80826-90	$\mu$ –Slide 8 Well ibiTreat, Bulk Pack: #1.5 polymer coverslip, tissue culture treated, sterilized
80822	μ-Slide 8 Well Collagen IV: #1.5 polymer coverslip, sterilized
80824	μ–Slide 8 Well Poly-L-Lysine: #1.5 polymer coverslip, sterilized
80821	μ–Slide 8 Well Uncoated: #1.5 polymer coverslip, hydrophobic, sterilized
80827	$\mu$ –Slide 8 Well Glass Bottom: #1.5 (170 $\mu$ m ±5 $\mu$ m) D 263 M Schott glass, sterilized
80827-90	$\mu\text{Slide 8 Well Glass Bottom, Bulk Pack: } \#1.5~(170~\mu\text{m}~\pm 5~\mu\text{m})~D~263~M~Schott~glass,$ sterilized

#### μ-Slide 8 Well Grid-500



Cat. No.	Description
80826-G500	$\mu$ –Slide 8 Well ibiTreat Grid–500: #1.5 polymer coverslip, tissue culture treated, grid repeat distance 500 $\mu$ m, sterilized
80821-G500	$\mu\text{-Slide 8 Well Uncoated Grid-500}\colon$ #1.5 polymer coverslip, hydrophobic, grid repeat distance 500 $\mu m$ , sterilized

### For research use only!

Further technical specifications can be found at www.ibidi.com. For questions and suggestions please contact us by e-mail *info@ibidi.de* or by telephone +49 (0)89/520 4617 0. All products are developed and produced in Germany. © ibidi GmbH, Am Klopferspitz 19, 82152 Martinsried, Germany.