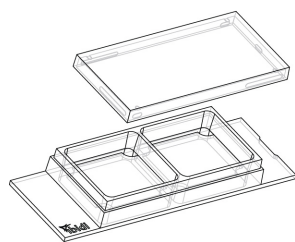


## Instructions

## $\mu$ -Slide 2 Well <sup>Ph+</sup> Glass Bottom



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 glass bottom versions of the  $\mu$ -Slides and  $\mu$ -Dishes are especially designed for TIRF, super resolution and single molecule applications. The  $\mu$ -Slide 2 Well <sup>Ph+</sup> Glass Bottom (Phase contrast plus) is a convenient chambered coverslip with 2 wells for cell culture, immunofluorescence, and high-end microscopy. The  $\mu$ -Slide 2 Well <sup>Ph+</sup> Glass Bottom improves the optical quality of phase contrast microscopy. In contrast to the classic  $\mu$ -Slide 2 Well, the Ph+ version provides a special plate in the center of the wells. This plate sup-

presses the meniscus which is disturbing the phase contrast effect in normal open wells. Openings near the corners provide access to the wells for filling and aspirating liquids easily.

## Material

The  $\mu$ -Slide 2 Well <sup>Ph+</sup> Glass Bottom is made with a glass coverslip bottom. It is not possible to detach the bottom. The  $\mu$ -Slide 2 Well <sup>Ph+</sup> Glass Bottom is not autoclavable since it is temperature stable only up to 80°C/175°F.

### Optical Properties ibidi Glass Bottom

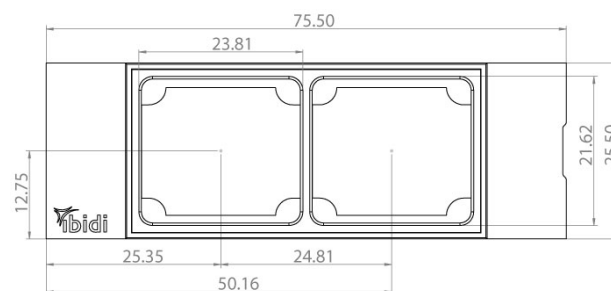
Refractive index $n_D$	1.523
Abbe number	55
Thickness	No. 1.5H (selected quality 170 $\mu$ m, $\pm$ 5 $\mu$ m)
Material	Schott borosilicate glass, D 263M

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	
Glass Bottom	36 months

## Geometry

The  $\mu$ -Slide 2 Well <sup>Ph+</sup> Glass Bottom provides a standard slide format according to ISO 8037/1.



### Geometry of the $\mu$ -Slide 2 Well <sup>Ph+</sup> Glass Bottom

Outer dimensions in mm (w × l)	25.5×75.5
Number of wells	2
Dimensions of wells in mm (w × l × h)	21.6 × 23.8 × 3.0
Volume per well	1.5 ml
Liquid height	3.0 mm
Total height with lid	10.8 mm
Growth area per well	5.1 cm <sup>2</sup>
Coating area per well	11.4 cm <sup>2</sup>
Bottom	Glass Bottom

## Attention!

Be cautious when handling ibidi labware products with glass bottom! The glass coverslip or glass slide is very fragile and might break easily. Handle with care to avoid physical injury and damage to devices through leakage of the medium.

## Surface

The  $\mu$ -Slide 2 Well <sup>Ph+</sup> Glass Bottom is manufactured with an uncoated glass coverslip. Washing steps (e.g. with PBS) before cell seeding can remove glass dust which is advantageous for direct cell growth on the surface.

## Shipping and Storage

The  $\mu$ -Slides,  $\mu$ -Dishes and  $\mu$ -Plates are sterilized and welded in a gas-permeable packaging. The shelf life under

### Coating

Detailed information about coatings is provided in [Application Note 08: Coating protocols for ibidi labware products](#).

In short, specific coatings are possible following this protocol:

1. Prepare your coating solution according to the manufacturer's specifications or reference.
2. Apply 1.5 ml and leave at room temperature for at least 30 minutes.
3. Aspirate the solution and wash with the recommended protein dilution buffer.
4. The μ-Slide 2 Well Ph+ Glass Bottom is ready to be used. Optionally let dry at room temperature. Attention, some coating proteins might degenerate when drying!

### 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 1.5 ml 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 1.5 ml fresh medium per well.

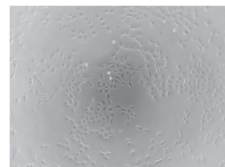
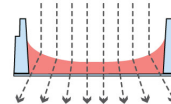
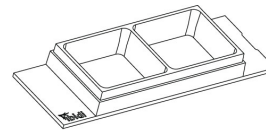
#### Tip:

The day before seeding the cells we recommend placing the cell medium and the μ-Slide into the incubator for equilibration. This will prevent the liquid inside from emerging air bubbles over the incubation time.

### μ-Slide 2 Well Selection Guide

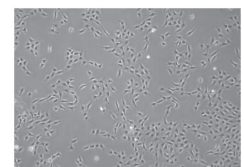
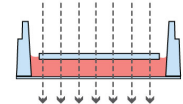
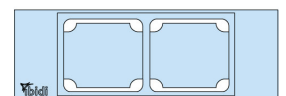
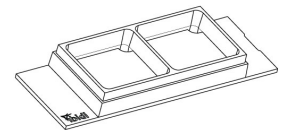
#### μ-Slide 2 Well

Standard open wells for maximum sample access. Meniscus disturbs the beam path. Good phase contrast quality only in the center of each well.



#### μ-Slide 2 Well Ph+

Special plate in the center of the wells suppresses meniscus formation. No meniscus – parallel beam path. For excellent phase contrast microscopy all over the wells.



### Immersion Oil

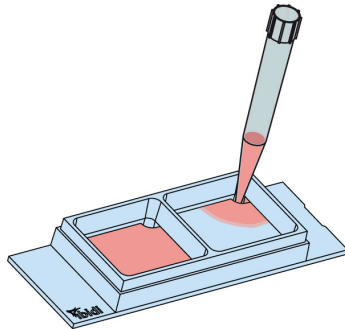
When using ibidi Glass Bottom products with oil immersion objectives, there is no known incompatibility with any immersion oil on the market. All types of immersion oils can be used.

### Microscopy

To analyze your cells, no special preparations are necessary. Cells can be directly observed live or fixed, preferably on an inverted microscope. The bottom cannot be removed. For optimal results in fluorescence microscopy and storage of fixed and stained samples, ibidi provides a mounting medium (50001) optimized for μ-Dishes, μ-Slides, and μ-Plates.

### Filling and Handling

Fill the wells by using a standard pipet. Inject the cell suspension directly into one of the openings. Medium exchange is easily done by aspirating the entire volume and refilling using 1.5 ml per well.



### Chemical Compatibility

The following table provides some basic information on the chemical and solvent compatibility of the μ-Slide 2 Well <sup>Ph+</sup> Glass Bottom. For a full list of compatible solvents and more information on chemical compatibility, please visit the FAQ section on [ibidi.com](https://www.ibidi.com).

Chemical / Solvent	Compatibility
Methanol	yes
Ethanol	yes
Formaldehyde	yes
Acetone	no
Mineral oil	yes
Silicone oil	yes
Immersion oil	See <b>Immersion Oil</b> on page 2.

## Instructions

## $\mu$ -Slide 2 Well <sup>Ph+</sup> Glass Bottom

### Ordering Information

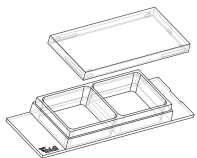
The  $\mu$ -Slide 2 Well is available as open well and as a Ph+ version, as well as in a glass bottom version. See the table below for choosing your  $\mu$ -Slide 2 Well.

#### $\mu$ -Slide 2 Well



Cat. No.	Description
80286	$\mu$ -Slide 2 Well ibiTreat: #1.5 polymer coverslip, tissue culture treated, sterilized
80282	$\mu$ -Slide 2 Well Collagen IV: #1.5 polymer coverslip, sterilized
80284	$\mu$ -Slide 2 Well Poly-L-Lysine: #1.5 polymer coverslip, sterilized
80281	$\mu$ -Slide 2 Well Uncoated: #1.5 polymer coverslip, hydrophobic, sterilized
80287	$\mu$ -Slide 2 Well Glass Bottom: 1.5H (170 $\mu$ m $\pm$ 5 $\mu$ m) D 263 M Schott glass, sterilized

#### $\mu$ -Slide 2 Well <sup>Ph+</sup>



Cat. No.	Description
80296	$\mu$ -Slide 2 Well <sup>Ph+</sup> ibiTreat: #1.5 polymer coverslip, tissue culture treated, sterilized
80292	$\mu$ -Slide 2 Well <sup>Ph+</sup> Collagen IV: #1.5 polymer coverslip, sterilized
80294	$\mu$ -Slide 2 Well <sup>Ph+</sup> Poly-L-Lysine: #1.5 polymer coverslip, sterilized
80297	$\mu$ -Slide 2 Well <sup>Ph+</sup> Glass Bottom: 1.5H (170 $\mu$ m $\pm$ 5 $\mu$ m) D 263 M Schott glass, sterilized

### For research use only!

Further information can be found at [www.ibidi.com](http://www.ibidi.com). For questions and suggestions please contact us by e-mail [info@ibidi.de](mailto:info@ibidi.de) or by telephone +49 (0)89/520 4617 0.

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