

# Square One

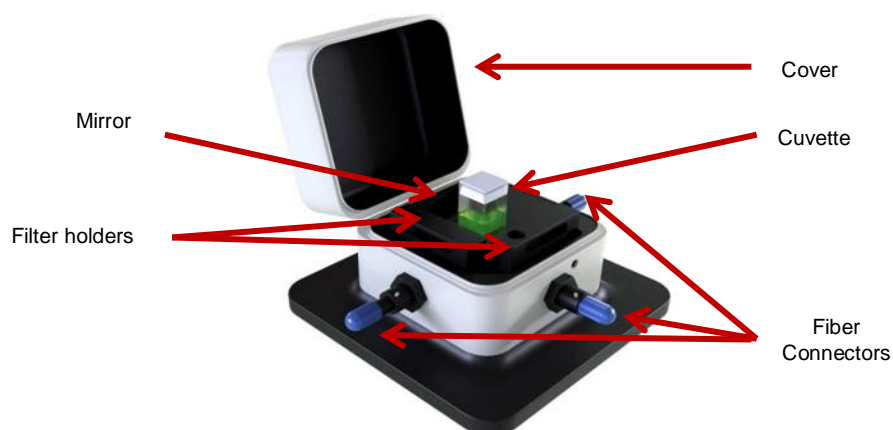
**Cuvette Holder  
Quick Start Guide**

The SQ1-ALL Cuvette Holder for 1-cm pathlength cuvettes couples via SMA-terminated optical fibers to spectrometers and light sources to create small-footprint spectrophotometric systems for absorbance and fluorescence experiments. The SQ1-ALL Cuvette Holder has a fully integrated cover for eliminating ambient light and has 2 filter slots to enable filtering illumination light entering the cuvette holder and/or detected light leaving the cuvette holder. The unit is designed to snugly hold 1-cm square cuvettes without user adjustment, providing high data repeatability.

## Parts Included

- Cuvette holder assembly for 1 cm cuvettes
  - 2 filter holders
  - 1 mirror
- 10 disposable cuvettes
- Allen wrench for adjusting the collimating lenses

## Installation



### Attaching the Fibers

1. Remove the aluminum caps from the desired connectors.
2. Attach one end of a SMA-terminated optical fiber to one of the collimating lenses. Attach the other end of this fiber – the illumination fiber – to a light source.
3. Attach another SMA-terminated optical fiber to the other collimating lens. Attach the other end of this fiber – the read fiber – to the spectrometer.

#### NOTE

The connector on the front of the unit is only used for fluorescence applications.

## Inserting Filters

1. Open the lid of the SQ1-ALL.
2. Remove the desired filter holder(s) from the unit.
3. Place filter on the surface of the holder and clip to secure.
4. Replace filter holder in unit.



## Inserting the Cuvette

1. Open the lid of the SQ1-ALL.
2. Firmly place cuvette in unit. No adjustment is necessary after insertion.
3. Close lid.

## Specifications

<b>Dimensions</b>	3.96" x 4.37" x 2.7" (L x W x H)
<b>Weight</b>	1.4 lb (0.6 kg)
<b>Pathlength</b>	1 cm
<b>Z dimension</b>	15 mm
<b>Collimating lenses (VIS-NIR)</b>	BK 7 glass (~360 nm – 2 μm*), 5 mm diameter, f/2
<b>Collimating lenses (UV-VIS-NIR)</b>	Quartz (200 nm-1.3 μm), 5 mm diameter, f/2
<b>Collimating lens termination</b>	SMA 905
<b>Collimating lenses assembly (sample compartment) dimensions</b>	2.0" x 1.5" (L x W)
<b>Filter slots</b>	2 slots
<b>Optical filters</b>	Accepts optical filters to 12.5 and 25 mm diameters. Filters may not be thicker than 4 mm.
<b>Base material</b>	aluminum
<b>Typical optical fibers specified for optimum performance*</b>	200 μm illumination fiber, 50 μm read fiber
* For all intents, there is no single combination of optical fibers that will satisfy the requirements of every application. As a rule, however, it is best to use a large-diameter (>50 μm) illumination fiber to get the maximum light throughput, and a small-diameter (<50 μm) read fiber to achieve the best optical resolution.	

# FAQ

## **What is the “Z” dimension of the cuvette holder?**

The “Z” dimension, 15 mm, is the optical height for placement of the light beam transmitting through the cuvette and is especially important with small-volume sampling.

## **Can I use the collimating lenses in the cuvette holder assembly for other applications?**

Yes. The 74-UV collimating lenses included with the holder can be unscrewed and used separately in any fixture for applications that require a fiber for free-beam coupling, such as on-line transmission or reflection.

## **How do the 74-MSP mirror screw plugs (accessory item) increase signal in the cuvette holder for fluorescence measurements?**

It's not so much that the screw plugs increase signal as they redirect some of the signal that otherwise is lost as it transmits through the sample compartment. Each plug – you'll need two – screws into a collimating lens port on the cuvette holder to redirect energy back to the sample or back into the collimating lens. The plugs are 0.3" (7.5 mm) UV-enhanced aluminum-coated reflection mirrors designed to collect the fluorescence that otherwise would be lost and to reflect the excitation energy back through the sample.

## **Which of the 3 connectors should I use?**

Light is normally directed toward the sample from one of the side ports. For absorbance measurements, the spectrometer is placed in-line with the light. For fluorescence measurements the spectrometer is connected to the front connector.

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## Questions?

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[info@oceaninsight.com](mailto:info@oceaninsight.com) • **US** +1 727-733-2447

**EUROPE** +31 26-3190500 • **ASIA** +86 21-6295-6600