

180 μm by placing a small drop of Permount (fresh - not overly viscous from sitting around for ages) and pressing the slide down on top of the cover glass with hand pressure for a few seconds and then heating on a slide warmer for a couple of hours. When I remeasured them at the center point, I got something very close to 1180 μm . My micrometer is marked at 10 μm intervals so greater precision than 5 μm is somewhat dicey.

If I used glass slides that had 8 μm paraffin sections (this was the thickness set on the microtome, which I realize isn't precise) and coverslipped them using the same method, I got a number equal to the thickness of the glass + cover glass + 10 μm (presumably the section thickness + mounting medium). Assuming the paraffin section was close to 8 μm , it would imply the mounting medium added about 2 μm . Although it would have been better if the No. 1 1/2 cover glasses had been closer to 170 μm thick, the percent error in using ones that were 180 μm would be less than starting with No. 1 cover glasses that were 150 μm thick and hoping to get an even 20 μm thick layer of mounting medium.

Finally, let me end with some published comments on cover glass thickness by notable authorities:

"It is therefore best to prepare a microslide with the No. 1 1/2 cover glass." John Gustav Daly in *Photography Through The Microscope* (1988) 9th edition, p. 20; Eastman Kodak Co.

"Standard coverslip are assumed to be 0.17 ± 0.01 mm thick (with a refractive index of 1.515). Number 1 1/2 coverslips are nominally selected for this standard thickness." The author goes on to state that coverslips should be measured for the most critical work. Shinya Inoue 1986) *Video Microscopy* 1st edition. p. 134; Plenum Press, NY.

"No. 1 1/2 generally gives the greatest yield of usable cover glasses." G.P. Berlyn, J. Miksche (1976) *Botanical Microtechnique and Cytochemistry*. pg. 8. Iowa State Univ. Press, Ames.

Regards,
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Dear Editor,

I include in my response to his initial query, with an added example of when No. 1 1/2 cover glasses are suitable (last paragraph) and a new last paragraph that addresses his follow-up post:

Correction: No. 1 cover glasses range 0.13 – 0.16 mm thickness; No. 1 1/2, 0.17-0.19 mm (American Society for Testing Materials. Standard Specification for Cover Glasses and Glass Slides for Use In Microscopy. ASTM Designation (E211-70, Effective 12-24-70).

From a different standard: No. 1 = 0.13 – 0.17 mm; No. 1 1/2 = 0.16 – 0.19 mm (Interim Federal Specification Cover Glass, Microscope. NNN-C-001434A, 01/08/71). No significant difference.

Thickness of mounting medium for tissue sections, 3 sets of 4 slides broken across the section, the broken edges trued up and polished and measured with a micrometer microscope (Aumonier FJ, Settingron R. Some notes on the mounting of histological sections. Proc Roy Micr. Soc. 1967;2:428-9):

Cover glass applied routinely (no pressure) = 10, 51, 63, and 76 μm

Cover glass weighted with 30 g for 2 days = 18, 18, 20, and 30 μm
Cover glass with spring loaded clothespin for 72 hours = 5, 10, 10, and 20 μm

Therefore, the thickness of mounting medium is substantial relative to the difference between the range of thickness for No. 1 cover glass and the tolerance of high dry achromat objectives to

deviations from optimal thickness of 0.17 mm (± 15 μm and more). Ergo, my recommendation to use No. 1 thickness cover glasses. A modest bonus is getting more No. 1 cover glasses per ounce for the same price as for No. 1 1/2.

Fluorite and apochromat objectives have higher NAs power for power than do achromats and so are even more sensitive to cover glass (and mounting medium) thickness. Objectives start to show intolerance to cover glass deviations at approximately 0.6 NA (40X achromat).

Cytologic preparations (e.g., conventional Pap smears, my field) are more problematic than histologic sections. Pap smears can sometimes require up to 12 or more drops of mounting medium to fill in all the valleys of thick preparations.

No. 1 1/2 cover glasses are suitable when there is little or no mounting medium between the specimen and cover glass, such as cells grown in culture on cover glasses, blood films spread on cover glass, cells on Nuclepore filters dissolved on a cover glass, and covered slides that have been "cooked" a la Dr. Ruth Graham in the 1950s (i.e., heated briefly on a hot place to drive off the solvent, which results in a hardened mount that can be handled immediately, beautiful imaging under 40X objectives, and long term stain preservation).

Even if one works with histologic preparations, I still recommend the No. 1 thickness cover glass. As the unique measurements above show, mounting medium thickness is sufficiently great when cover glasses are simply laid on the specimen to bring the combined thickness up to, and into, the 0.17 mm thickness neighborhood. Starting off with a No. 1 1/2 thickness cover glass is likely to result, more often than not, in total thicknesses that exceed the tolerance range of the high numerical aperture objective in use.

And a final repetitive note for emphasis: microscopes that have dirty lenses and are not adjusted to produce Koehler illumination will not reveal the beneficial contribution of correct cover glass/mounting medium. It will be lost in the noise.

Yours truly,
Gary W. Gill,
Diagnostic Cytology
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Protecting Inverted Lenses

Steve Sands, Pfizer Central Research

Inverted microscope objectives are expensive, so it pays to protect them. Unfortunately, being located below little chambers filled with buffers means that these objectives are subject to spills. A trick to help protect your objectives is to obtain small, powder free (latex or otherwise) gloves. Cut off one of the fingers (or thumb, depending on the obj.), cut a small hole in the tip, and then carefully full the glove over the objective such that the objective lens is exposed. It's not perfect, but if fitted properly, most liquid will run over the glove, and not into the objective. A small piece of lint free absorbent material, placed on the nosepiece can then absorb small amounts of runoff, and helps prevent liquid from entering the scope body. ■