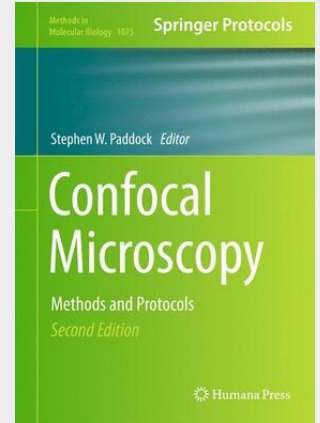


Paddock

Confocal Microscopy

Methods and Protocols

Confocal Microscopy: Methods and Protocols, Second Edition takes the researcher from the bench top through the imaging process, to the page. Protocols for the preparation of tissues from many model organisms including worms, flies and mice have been included as well as chapters on confocal imaging of living cells, three dimensional analysis, and the measurement and presentation of confocal images for publication. Emphasis has been placed on the laser scanning confocal microscope since this is still the instrument used for most routine applications. The current generation of modern confocal instruments produces optical sections of cells and tissues that are free of out-of-focus fluorescence with reduced chances of artifacts from the techniques of specimen preparation. This allows the imaging of living specimens and measurements of physiological events within cells. Confocal microscopy has become essential in many fields of contemporary biomedical research where a light microscope is required for imaging fluorescently labeled cells and tissues, especially cell biology, developmental biology, neurobiology, and pathology. Written in the successful Methods in Molecular Biology series format, chapters include introductions to their respective topics, lists of the necessary materials and reagents, step-by-step, readily reproducible protocols, and notes on troubleshooting and avoiding known pitfalls. Authoritative and easily accessible, Confocal Microscopy: Methods and Protocols, Second Edition is aimed primarily, but not exclusively, at the novice user with pointers to more advanced techniques.



160,49 €

149,99 € (zzgl. MwSt.)

Lieferfrist: bis zu 10 Tage

Artikelnummer: 9781588293510

Medium: Buch

ISBN: 978-1-58829-351-0

Verlag: Springer

Erscheinungstermin: 20.09.2013

Sprache(n): Englisch

Auflage: 2. Auflage 2014

Serie: Methods in Molecular Biology

Produktform: Gebunden

Gewicht: 8928 g

Seiten: 381

Format (B x H): 183 x 260 mm

