

Microtome Operation Manual

Mr. A. V. Naigaonkar

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science dealt with in a comprehensive manner tailored to the needs of postgraduate students research scholars for easy understanding The book is profusely illustrated The different chapters described in the book include Introduction Microscopy Plant micro technique Smear Sqush technique Plant tissue culture Herbarium technique Hydrogen ion concentration pH Centrifugation Chromatography Electrophoresis Colorimetry Spectro photometry Radio isotopes in biology and Computers and their application in plant sciences Chapters on Biostatistics Biophysics and Bioinformatics have also been included to help the student in the statistical analysis of the results physical principles involved in the operation of different instruments and basics of bioinformatics We sincerely hope that this book helps to fill up the lacuna and provides what all that is needed about the research methods required for a scholar student in plant sciences to pursue their higher studies

Low-Temperature Microscopy and Analysis Patrick Echlin,2013-11-11 The frozen hydrated specimen is the principal element that unifies the subject of low temperature microscopy and frozen hydrated specimens are what this book is all about Freezing the sample as quickly as possible and then further preparing the specimen for microscopy or microanalysis whether still embedded in ice or not there seem to be as many variations on this theme as there are creative scientists with problems of structure and composition to investigate Yet all share a body of com mon fact and theory upon which their work must be based Low Temperature Micros copy and Analysis provides for the first time a comprehensive treatment of all the elements to which one needs access What is the appeal behind the use of frozen hydrated specimens for biological electron microscopy and why is it so important that such a book should now have been written If one cannot observe dynamic events as they are in progress rapid specimen freezing at least offers the possibility to trap structures organelles macro molecules or ions and other solutes in a form that is identical to what the native structure was like at the moment of trapping The pursuit of this ideal becomes all the more necessary in electron microscopy because of the enormous increase in resolution that is available with electron optical instruments compared to light optical microscopes

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