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PRINS and In Situ PCR Protocols (Second Edition)
Primed In Situ Hybridization (PRINS) was introduced about the same time as Fluorescent In Situ Hybridization (FISH). PRINS offers a faster detection method by making use of small unlabeled DNA sequences (oligonucleotides) that specifically bind to repeat sequences on the chromosomes (alpha satellite, Alu repeats, or telomeres) and then multiply exponentially through annealing and in situ elongation of the chromosomes. Visualization can take place by incorporating one labeled nucleotide into PRINS and In Situ PCR Protocols: Franck Pellestor ... Books Advanced Search Today's Deals New Releases Best Sellers The Globe & Mail Best Sellers New York Times Best Sellers Best Books of the Month Children's Books Textbooks Kindle Books Livres en fran ais PRINS and In Situ PCR Protocols: Franck Pellestor ... Books Advanced Search Today's Deals New Releases Amazon Charts Best Sellers The Globe & Mail Best Sellers New York Times Best Sellers Best Books of the Month Children's Books Textbooks Kindle Books Livres en fran ais

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In Situ PCR to Cells and to Wax Sections Christiane Mougin, Jean Marc Didier, Dominique Bettinger, Laurence Madoz, Sylviane Coumes-Marquet, Michel Lab Pages 77-97

PRINS and in situ PCR protocols. - cabdirect.org
Abstract : This multiauthor text contains 6 chapters on protocols relating to PRINS (oligonucleotide primed in situ synthesis), one on applying PRINS to extended chromatin preparations, 2 on combining PRINS with immunocytochemical analysis of cellular and chromosomal proteins, and a further 6 on applying polymerase chain reaction (PCR) in situ to different problems.

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The in situ hybridization and PCR technologies are now well-established molecular techniques for studying chromosomal aneuploidy and rearran- ments, gene localization and expression, and genomic organization. Over the last decade, we have seen increasing applications in these fields. By combining

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in situ, and it is to this novel field that PRINS and In Situ PCR Protocols is devoted. Conventional in situ hybridization uses a DNA or RNA probe, labeled with a detectable moiety (most commonly a hapten, such as biotin or digoxigenin) that is annealed to the DNA or RNA of a target tissue or cyto-logical preparation. Imperfectly annealed probe is removed by stringent washing, and the

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Abstract. This chapter provides detailed methods and material descriptions of in situ polymerase chain reaction protocols. It describes all the essential components of the technique, various protocols suitable for different kinds of tissues and cell preparations, and also gives various instructions for troubleshooting.

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In the first edition of PRINS and In Situ PCR Protocols, edited by John R. Gosden, experts in the field presented in detail a variety of applications of PRINS and in situ PCR techniques, in a wide range of clinical conditions. Since the publication of this successful reference book, there have been significant improvements in in situ detection techniques. This completely revised and updated

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The in situ hybridization and PCR technologies are now well-established molecular techniques for studying chromosomal aneuploidy and rearran- ments, gene localization and expression, and genomic organization. Over the last decade, we have seen increasing applications in these fields. By combining

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