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In Situ Hybridization Screening of a Lambda Library for Chromosome Specific DNA: The Isolation of a Rabbit Chromosome Number 9 Probe

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1 IN SITU HYBRIDIZATION SCREENING OF A LAMBDA LIBRARY FOR
CHROMOSOME SPECIFIC DNA: THE ISOLATION OF A RABBIT
CHROMOSOME NUMBER 9 PROBE

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3 Down's syndrome, a disease which affects 1 of every 783 newborns, is the result of aneuploidy of human chromosome 21. The closely related rabbit chromosome 20 can be used as a model for the study of the cause of the misdistribution of this chromosome during meiosis. The procedure, called karyotyping, requires long hours behind the microscope; however, it can be replaced with the more accurate, time-efficient restriction fragment length polymorphism (RFLP) assay. The purpose of this study was to test the feasibility of finding a chromosome specific probe by searching a lambda library using IN SITU hybridization. IN SITU hybridization involves annealing a radioactive DNA probe directly to metaphase chromosomes fixed to a microscope slide. A photographic emulsion is spread over the slides. Photographic grains appear over the chromosome to which the DNA has annealed thus identifying the chromosome from which the probe DNA originated (FIG. 1). Probe DNA was phenol extracted from lambda phage virus clones that carry rabbit DNA inserts. Inserts were purified by gel electrophoresis (FIG. 2). Using this procedure, a chromosome 9 probe was discovered in the 86 phage DNA clones isolated (FIG. 3). This suggests that a larger search would be capable of finding the chromosome 20 probe needed to develop a rabbit model for Down's syndrome.