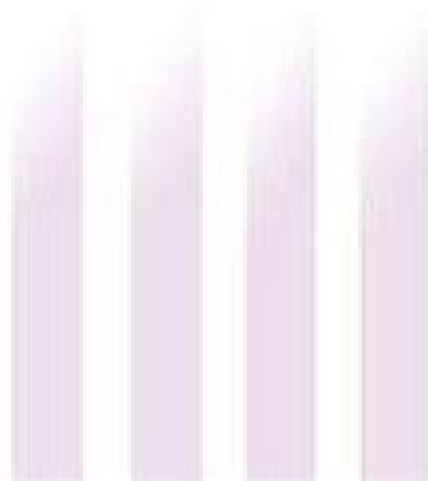


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PCR: Clinical Diagnostics and Research



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Biochemical Technology, Part A L. Bulow, 1997-05-06 In December 1992 the Department of Pure and Applied Biochemistry at the Chemical Center in Lund Sweden organized an international meeting the Mosbach Symposium on Biochemical Technology to celebrate the 60th birthday of professor Klaus Mosbach one of the founders of modern biotechnology The history of Pure and Applied Biochemistry had its start in 1970 a couple of years after the foundation of the Chemical Center Klaus Mosbach has been its professor and head of Pure and Applied Biochemistry since its start During the 1980 s he also maintained a professorship at the ETH in Zurich Switzerland Professor Mosbach is internationally well known and he has world leading position within the field of immobilization of bioactive substances and cells as well as affinity chromatography In 1990 Professor Mosbach was awarded the gold medal by the Royal Swedish Academy of Engineering Sciences for his contributions to biotechnology especially on the immobilization of bioactive substances The research activities of the Department of Pure and Applied Biochemistry cover a broad area such as affinity and separation techniques bioprocess control biosensors development of new carriers and new immobilization procedures for small molecules as well as proteins and cells including animal and plant cells gene technology processes based on immobilized biocatalysts and construction of organic polymers with enzyme like properties The hallmark of the department is its diversified research that generates considerable synergistic effects that are manifested by many new techniques and concepts emanating from the laboratory during the last 20 years Several of these are marketed by various biotechnology companies At this meeting we therefore arranged for some of the world's leading experts in biochemistry and biotechnology to give lectures The topics covered comprise enzyme technology immobilization of enzymes and cells abzymes metabolic engineering biosensors and molecular recognition The official gift from the symposium committee and the participants is this Festschrift which covers several important fields of research within the area of biochemical technology We have made a very unusual approach and have let the hero of the occasion present the history of his research

PCR: Clinical Diagnostics and Research Arndt Rolfs, Irmela Schuller, Ulrich Finckh, Ines Weber-Rolfs, 2011-12-23 In 1985 Kary Mullis was driving late at night through the flowering buckeye to the ancient California redwood forest cogitating upon new ways to sequence DNA Instead he came upon a way to double the number of specific DNA modules and to repeat the process essentially indefinitely 1 He thought of using two oligonucleotide sequences oppositely oriented and a DNA polymerase enzyme to double the number of DNA targets Each product would then become the target for the next reaction effectively yielding a product which doubled in quantity with each repeated cycle Like the chain reaction leading to nuclear fission with each cycle event each initial reactant neutron or DNA molecule yields two similar products each of which can serve as the initial reactant The invention of this exponentially increasing amplification system quickly became known as the polymerase chain reaction PCR The tremendous sensitivity of PCR ultimately resides in the necessity for each of two specific oligonucleotide annealing reactions

to occur at the same time in the proper orientation The DNA annealing reaction is a very specific reaction Single genes have been detected by hybridization of a DNA probe to chromosome preparations together with sensitive fluorescence microscopy This is the equivalent to detecting a gene present in a single copy per cell genome It is the combination of two such specific annealing reactions which makes possible the amplification needed to detect a single molecule with a specific DNA sequence in over 100 000 cell genomes

Nonisotopic Probing, Blotting, and Sequencing Larry J. Kricka,1995-06-01 Since the publication of Nonisotopic DNA Probe Techniques in 1992 the move away from radioactive materials for research and diagnostics has continued This is due in part to public awareness of the hazards of radioactive waste and laws making radioactive disposal more difficult and costly and to improvement in both the sensitivity and convenience of nonisotopic techniques Several new nonisotopic techniques have been developed and substantial improvements made to existing nonisotopic methods since 1992 and these are now included in Nonisotopic Probing Blotting and Sequencing Nonisotopic Probing Blotting and Sequencing is an updated expanded edition of the bestseller Nonisotopic DNA Probe Techniques It has been thoroughly revised to include the latest improvements in nonisotopic tagging techniques for macromolecules Like its predecessor it enables researchers to select the best nonisotopic method for their needs and maximize success by following its straightforward protocols Provides strategies and detailed procedures for labeling blotting and probing specific nucleic acid sequences and with this edition protein molecules Gives protocols for nonisotopic DNA sequencing new in this edition Gives extensive practical information Presents background information for each method Provides expert accounts from the inventor or developer of each method Contains seven entirely new chapters Covers all major types of nonisotopic procedures for labeling and detection

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