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| Altri autori (Persone)  | ReymondJean-Louis  |
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| Nota di contenuto       | Enzyme Assays; Contents; Preface; List of Contributors; Introduction;<br>Enzyme Assays; Part I: High-throughput Screening; Part II: Genetic<br>Selection; Part III: Enzyme Fingerprinting; Enzyme Assays in Other<br>Areas; How to Use this Book; Part I High-throughput Screening; 1<br>Quantitative Assay of Hydrolases for Activity and Selectivity Using Color<br>Changes; 1.1 Overview; 1.2 Direct Assays Using Chromogenic<br>Substrates; 1.3 Indirect Assays Using Coupled Reactions - pH<br>Indicators; 1.3.1 Overview of Quantitative Use of pH Indicator Assay;<br>1.3.2 Applications<br>1.3.2.1 Searching for an Active Hydrolase (Testing Many Hydrolases<br>Toward One Substrate)1.3.2.2 Substrate Mapping of New Hydrolases<br>(Testing Many Substrates Toward Hydrolase); 1.3.3 Comparison with<br>Other Methods; 1.4 Estimating and Measuring Selectivity; 1.4.1<br>Estimating Selectivity without a Reference Compound; 1.4.2<br>Quantitative Measure of Selectivity Using a Reference Compound (Quick<br>E and Related Methods); 1.4.2.1 Chromogenic Substrate; 1.4.2.2 pH<br>Indicators; 1.4.3 Application; 1.4.3.1 Substrate Mapping of Hydrolases;<br>1.4.3.2 Screening of Mutants in Directed Evolution<br>1.4.4 Advantages and DisadvantagesReferences; 2 High-throughput<br>Screening Systems for Assaying the Enantioselectivity of Enzymes; 2.1<br>Introduction; 2.2 UV/Vis Spectroscopy-based Assays; 2.2.1 Assay for |

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|                    | Screening Lipases or Esterases in the Kinetic Resolution of Chiral p-<br>Nitrophenyl Esters; 2.2.2 Enzyme-coupled UV/Vis-based Assay for<br>Lipases and Esterases; 2.2.3 Enzymatic Method for Determining<br>Enantiomeric Excess (EMDee); 2.2.4 UV/Vis-based Enzyme<br>Immunoassay as a Means to Measure Enantiomeric Excess; 2.2.5 Other<br>UV/Vis-based ee-Assays; 2.3 Assays Using Fluorescence<br>2.3.1 Umbelliferone-based Systems2.3.2 Fluorescence-based Assay<br>Using DNA Microarrays; 2.3.3 Other Fluorescence-based ee-Assays; 2.4<br>Assays Based on Mass Spectrometry (MS); 2.4.1 MS-based Assay Using<br>Isotope Labeling; 2.5 Assays Based on Nuclear Magnetic Resonance<br>Spectroscopy; 2.6 Assay Based on Fourier Transform Infrared<br>Spectroscopy for Assaying Lipases or Esterases; 2.7 Assays Based on<br>Gas Chromatography; 2.8 Assays Based on HPLC; 2.9 Assays Based on<br>Capillary Array Electrophoresis; 2.10 Assays Based on Circular<br>Dichroism (CD)<br>2.11 Assay Based on Surface-enhanced Resonance Raman Scattering2.<br>12 Conclusions; References; 3 High-throughput Screening Methods<br>Developed for Oxidoreductases; 3.2.1 Dehydrogenases; 3.2.1.1<br>Colorimetric Screen Based on NAD(P)H Generation; 3.2.1.2 Screens<br>Based on NAD(P)H Depletion; 3.2.2 Oxidases; 3.2.2.1 Galactose<br>Oxidase; 3.2.2.2 D-Amino Acid Oxidase; 3.2.2.3 Peroxidases; 3.2.3<br>Oxygenases; 3.2.3.1 Assays Based on Optical Properties of Substrates<br>and Products |
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|                    | 3.2.3.2 Assays Based on Gibbs' Reagent and 4-Aminoantipyrine   |
| Sommario/riassunto | Edited by one of the leading experts in the field, this book fills the need<br>for a book presenting the most important methods for high-<br>throughput screenings and functional characterization of enzymes. It<br>adopts an interdisciplinary approach, making it indispensable for all<br>those involved in this expanding field, and reflects the major advances<br>made over the past few years.For biochemists, analytical, organic and<br>catalytic chemists, and biotechnologists.  |