

HPLC METHOD DEVELOPMENT FOR LC/MS: TRADITIONAL APPROACHES AND EMERGING TRENDS

1. Introduction and Background of LC/MS/MS in the Bioanalytical Laboratory

- 1.1. Brief History of Coupling HPLC and Mass Spectrometry
 - 1.1.1. Early Pitfalls of LC/MS/MS without Good HPLC Separations
 - 1.1.2. Why is a separation important with the selectivity of MS?
 - 1.1.2.1. Ionization effects in the source of the MS
 - 1.1.2.2. Separation of labile molecules from parent
 - 1.1.3. Separation of isobaric drugs or metabolites

2. Development of LC/MS/MS Methods Based on Structure of Molecule

- 2.1. **What is important in the structure of the analyte?**

3. The Latest Emerging Trends in HPLC/MS/MS Bioanalysis

3.1. The advantages of UPLC

3.2. The advantages of using SPE as a orthogonal technique

3.2.1. Removal of phospholipids

3.2.2. Micro-SPE

3.3. Current automation tools

4. Typical Problems, Solutions and Troubleshooting HPLC for MS Detection

- 4.1. Matrix Interferences and Ion-Suppression in LC/MS
 - 4.1.1. How to determine the occurrence of ion-suppression
 - 4.1.2. How to minimize ion-suppression
- 4.2. How to improve chromatographic peak shape
- 4.3. How to improve chromatographic retention
- 4.4. LC/MS Survival Kit-Tips and Tricks for What you really need to know
 - 4.4.1. Identifying when something goes wrong

5. Dried Blood Spot Analysis: Method Development and Validation Approaches in the LC/MS/MS Bioanalytical Laboratory

6. Tutorial in LC/MS Method Development

- 6.1. Dissecting some case studies for specific problems encountered by attendees
- 6.2. Example method development schemes based on select molecules

7. Questions and Answers