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Approaches"; ""1.3. Mass Spectrometry in Metabolomics Research"; ""2. STAGES IN AN LC-MS BASED METABOLOMICS ANALYSIS"; ""2.1. Sample Collection and Sample Storage"; ""2.2. Sample Homogenisation/Extraction/Deproteinisation"; ""2.3. Liquid Chromatography (LC)"; ""2.4. The Ionisation Process as an Interface between LC and MS: Focus on Matrix Effect"; ""2.5. Mass Spectrometry"; ""2.6. Data Handling"; ""2.6.1. Data processing"; ""2.6.1.1. Filtering""
""2.6.1.2. Feature/peak detection""""2.6.1.3. Alignment""; ""2.6.1.4. Normalisation"; ""2.6.2. Data analysis"; ""2.6.2.1. Principal component analysis (PCA) ""; ""2.6.2.2 Partial least-squares projections to latents structures (PLS)"; ""2.6.2.3. O-PLS"; ""2.7. Metabolite Identification"";
""3. CONCLUSION""; ""REFERENCES""; ""BIOMARKER DISCOVERY FOR CANCER DIAGNOSIS USING SERUM PROTEOMIC ANALYSIS:FROM BASIC RESEARCH TO CLINICAL APPLICATION""; ""ABSTRACT"";
""INTRODUCTION""; ""1. PROTEOMIC ANALYSIS OF SERUM/PLASMA IS EFFECTIVE TO SEARCH FOR CANCER DIAGNOSTIC MARKERS""
""1.1. Characteristics of Global Analyses and Diagnostic Availability of Biomarkers""""1.2. Advantages of Proteomic Analysis in the Search for Diagnostic Biomarkers""; ""1.3. Specimen""; ""2. SEPARATION TECHNOLOGIES IN PROTEOMIC ANALYSIS""; ""2.1. 2D-DIGE""; ""2.2. Protein Chip ArrayA®""; ""2.3. ClinProtA®""; ""2.4. Shotgun Proteomics Using LC""; ""3. MASS SPECTROMETRY""; ""3.1. Fundamentals of MS"";
""3.2. Ionization Methods""; ""3.3. Mass Analyzers""; ""3.4. Types of MS Used in Proteomic Analysis""; ""4. IDENTIFICATION OF PROTEINS/PEPTIDES USING MS""
""4.1. Protein Identification by Peptide Mass Fingerprinting""
