

MALDI Ion Imaging and Biological Ion Imaging with a new Scanning UV-Laser Microprobe

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LAMMA 2000 is a new scanning laser ion microprobe, developed in our laboratory, for inorganic and organic mass spectrometrical analysis of e.g. biological or technical samples.

An area of $100 \times 100 \mu\text{m}$ is scanned by a high-frequency pulsed UV laser with a lateral resolution of $\approx 0.5 \mu\text{m}$ ($\approx 1.0 \mu\text{m}$ for MALDI samples). Time-of-flight mass spectra of each pixel are evaluated with respect to several ion signals and are transformed into two-dimensional ion distribution plots by the data acquisition program (ULISSES 7.3). For detailed technical description see other paper on this conference.

Ion images obtained so far demonstrate the instrumental performances with respect to imaging lateral distributions of ion concentrations from various technical and biological samples. For non-flat samples, signal intensities are not a direct measure of substance concentrations, but are convoluted with a variation of the total ion current. This is due to the fact that the focus depth is in the μm range, which additionally allows to develop three-dimensional mass spectrometry techniques.

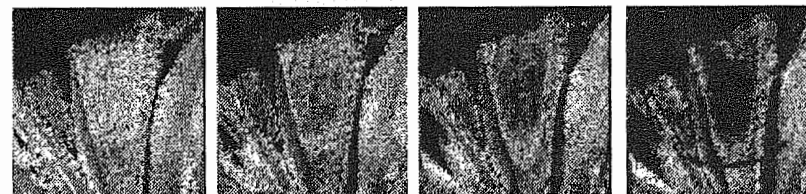
Samples prepared for MALDI (matrix assisted laser desorption ionization) MS analysis of peptides have been investigated by LAMMA 2000 ion imaging. The goal of this study was the development of a method of correlating preparational protocols with microscopical sample topology and mass spectrometrical results. In MALDI MS of biopolymers the preparation protocol plays a major role for analytical success, achievable sensitivity and topological homogeneity of the sample with respect to analyte ion formation.

Ion images obtained from MALDI samples demonstrate that

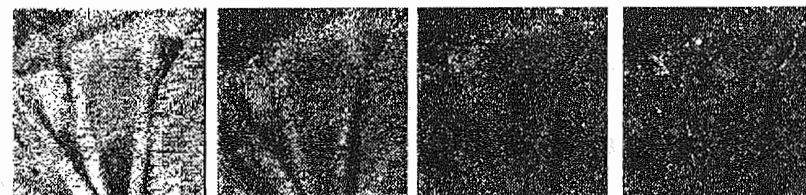
- MALDI-MS is possible even under strongly focused conditions (focus diameter $\approx 1 \mu\text{m}$), suggesting the development of sensitivity-enhanced micro-preparation procedures,
- analyte ion intensities basically image the physical structure of matrix crystals,
- analyte ion intensities and alkali ion intensities are in general mutually exclusive,
- alkali ions are mainly located between larger crystals and (homogeneously dispersed) in the inner part of the sample,
- alkali ions are not incorporated into matrix crystals,
- analyte ions are incorporated into matrix crystals,
- analyte ion images usually look less smooth from the first laser shot per pixel, compared to the following shots,
- the method allows to investigate dynamical sample erosion, preparational effects, influences of e.g. impurities and adducts.

Acknowledgement:

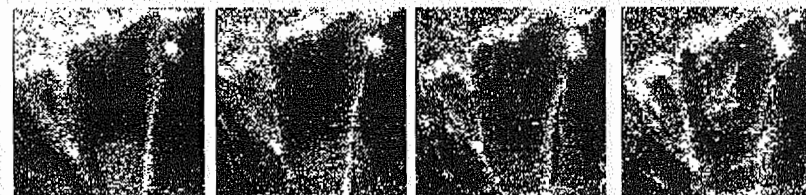
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Substance P

100 μm 

DHB



K

scan 1

scan 2

scan 3

scan 4

Ion images of MALDI samples of substance p (MW=1348 u) in 2,5-dihydroxybenzoic acid as matrix. Ion signals of analyte, matrix and potassium are imaged with one laser shot per pixel. Four consecutive scans are shown.

The area scanned is part of the crystallized rim of a dried droplet, with the inner area of the droplet on top.

White = high ion intensity; Black = low ion intensity