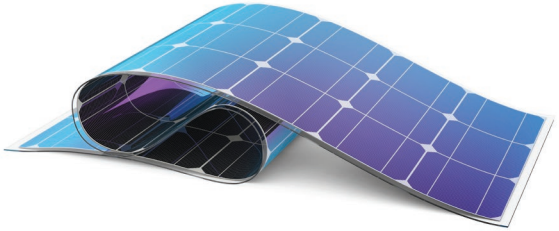
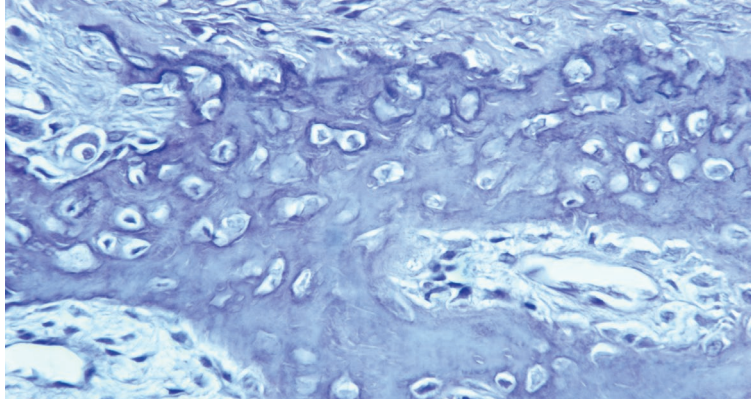
PHYSICAL ELECTRONICS
A DIVISION OF ULVAC-PHI

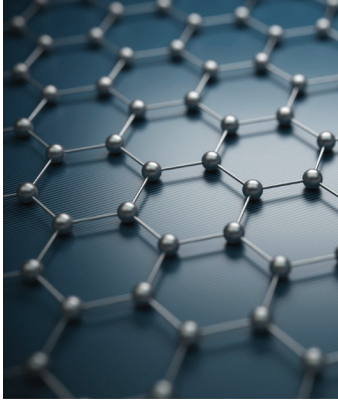
PHI

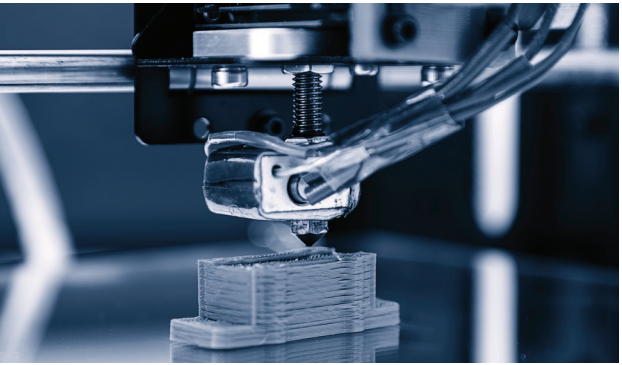
nanoTOF II

TOF-SIMS Parallel Imaging MS/MS

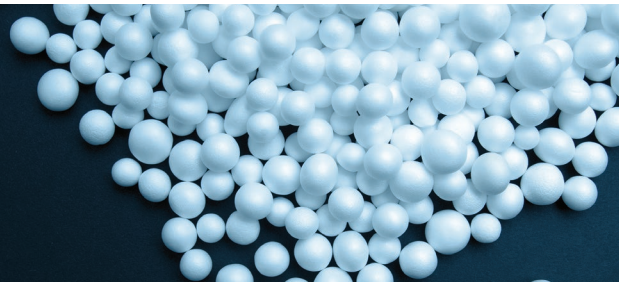


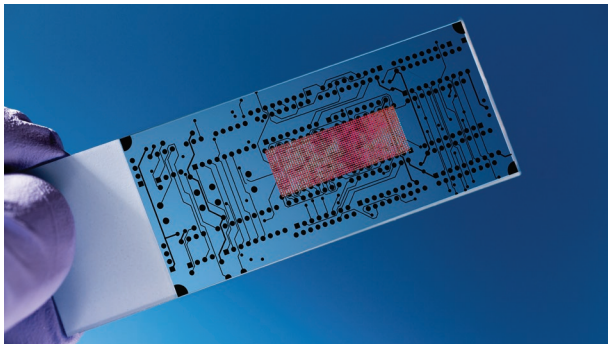







10:55





PHI *nanoTOF* II

Designed for Confident Molecular Identification and Superior Imaging

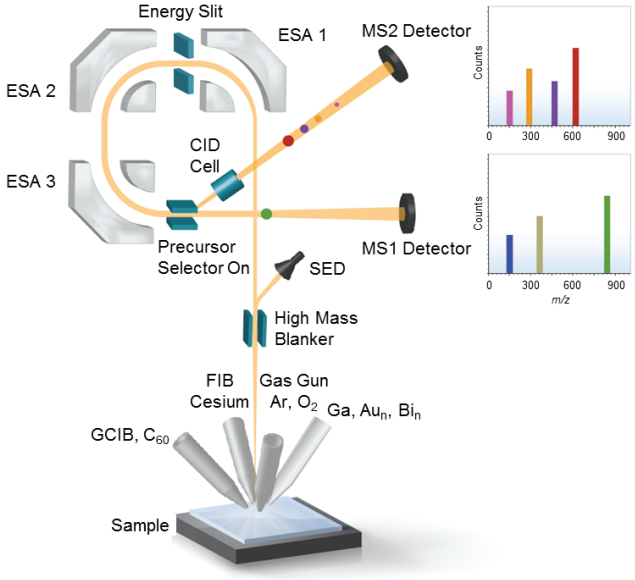
Time-of-flight secondary ion mass spectrometry (TOF-SIMS) is a key analytical technique for detecting, identifying, and imaging the distribution of both elements and molecules on the surface of materials. TOF-SIMS is the only mass spectrometry imaging technique that provides less than 70 nm spatial resolution with full mass range detection. The PHI *nanoTOF* II Parallel Imaging MS/MS platform has established itself as uniquely capable of providing superior analytical data, even on the most challenging samples. The *nanoTOF* II incorporates PHI's revolutionary Parallel Imaging MS/MS mass spectrometer technology into the base system. This patented spectrometer dramatically simplifies TOF-SIMS data interpretation and peak identification without compromising spatial resolution and speed. For the first time, confident identification is possible for high mass ions, transforming peak identification from ***"I think"*** to ***"I know!"***

The PHI *nanoTOF* II Parallel Imaging MS/MS instrument was designed with ***"No Compromise"*** in mind ensuring that there are no compromises in analytical performance. The synergistic design optimizes and combines multiple performance metrics into straight forward modes of analysis. It provides high lateral resolution and high mass resolution simultaneously in the HR^2 mode of analysis. The patented mass spectrometer provides superior imaging of highly topographic samples and high mass resolution spectra with excellent mass accuracy over the entire mass range. PHI's integrated MS/MS enables confident peak identification and delivers high speed TOF-TOF mode imaging at > 8 kHz. The sample stage provides full 5-axis sample motion and active temperature control from sample introduction through analysis. Productivity is maximized with a high level of automation and intuitive software.

TANDEM MASS SPECTROMETERS

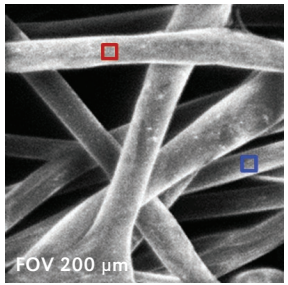
Maximum 2D/3D Information Content

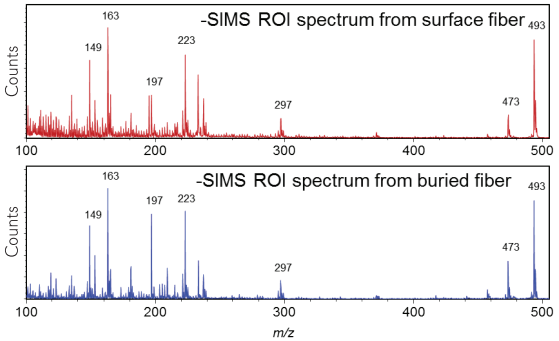
No compromise provides the capability for parallel and synchronous TOF-SIMS (MS^1) and tandem MS (MS^2) imaging. Never before has tandem MS imaging been available at the < 70 nm spatial resolving power provided by TOF-SIMS. The revolutionary and patented PHI Parallel Imaging MS/MS provides the maximum mass spectrometry imaging information, with unequivocal molecular identification, from any specified analytical region. Since the TOF-SIMS and tandem MS data are collected simultaneously and from the same volume, the MS^1 and MS^2 images are always in perfect registry. The elegant and fully integrated Parallel Imaging MS/MS spectrometer provides peak identification and imaging in a single analysis.



The schematic highlights the triple electrostatic analyzer (ESA) design of the *nanoTOF* II's Parallel Imaging MS/MS spectrometer. Selected precursor ions are deflected at keV energy into a collision-induced dissociation (CID) cell for fragmentation. The product ions are collected at the MS2 detector. Any ions not within the monoisotopic precursor selection window continue on their original flight path to the MS1 detector.

UNMATCHED PERFORMANCE





A gray-scale total ion image of a non-woven fiber mat. A region-of-interest (ROI) of a surface fiber is highlighted in red, and an ROI of a buried fiber is highlighted in blue. Irrespective of the sample topography and height, the spectral performance is almost identical owing to the exceptional depth-of-field of the mass spectrometer.

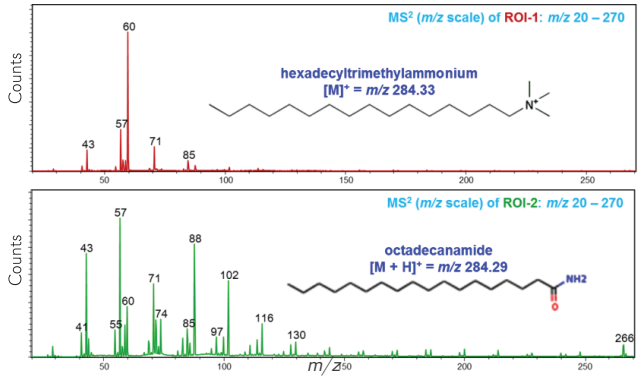
Confident Chemical Analysis of All Samples

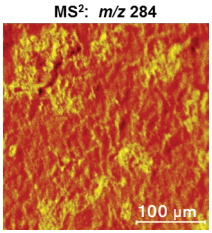
No compromise is apparent in the uncompromising performance of PHI's Parallel Imaging MS/MS spectrometer. The mass spectrometer provides high collection efficiency, high mass resolution, high mass accuracy and high signal-to-background spectra, simultaneously, for chemical imaging of flat and rough samples alike. Accurate elemental and molecular imaging of highly topographic, textured and FIB-sectioned samples is a unique capability due to the TOF-TOF spectrometer's optics design.

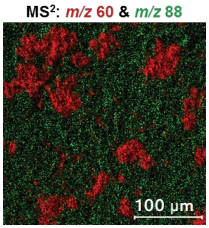
PARALLEL IMAGING WITH IDENTIFICATION

High Speed Imaging and Identification

No compromise delivers ion identification, structural elucidation, and high resolutoin imaging within a tandem MS analysis. PHI's TOF-TOF spectrometer enables the generation of pure, single-molecule spectra from the complex mixture spectrum which greatly simplifies data interpretation and peak identification. Molecular identification and imaging are achieved at the highest sensitivity; therefore, one-of-a-kind samples may be probed repeatedly for identification of numerous molecular precursors. PHI's Parallel Imaging MS/MS takes TOF-SIMS peak identification from *"I think" to "I know!"*

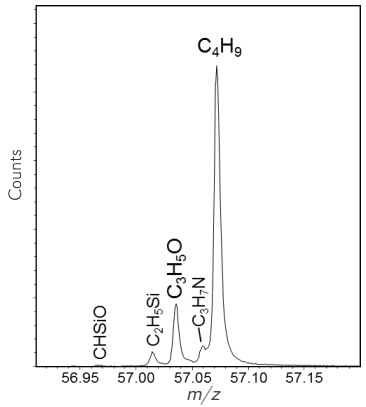


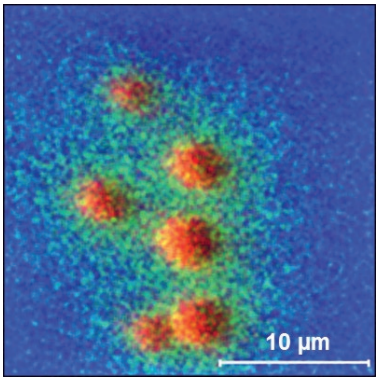




The power of Parallel Imaging MS/MS for molecular identification via tandem MS imaging is demonstrated in the chemical separation of two precursor ions both having a nominal *m/z* of 284. In only a 10 minute acquisition, each molecular component is easily observed in the MS^2 overlay shown in the lower right image. Mass spectra are generated from each region-of-interest (ROI), comprised of either the red or the green pixels, and the corresponding spectra are shown to the left. The composition and identification of each molecule is made by reference library matching.

HR² IMAGING CLUSTER LMIG





Low MW HC Oil, Aromatic HCs, Substrate

The power and utility of HR² mode imaging is exemplified in the surface analysis of micro droplets. In only a 6 minute acquisition the micron-sized droplets are spatially resolved, and the high mass resolution spectrum allows molecular identification.

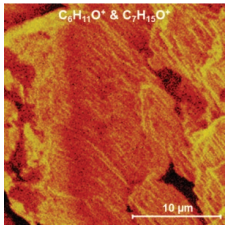
Simultaneous Spatial and Spectral Resolution of Chemistry

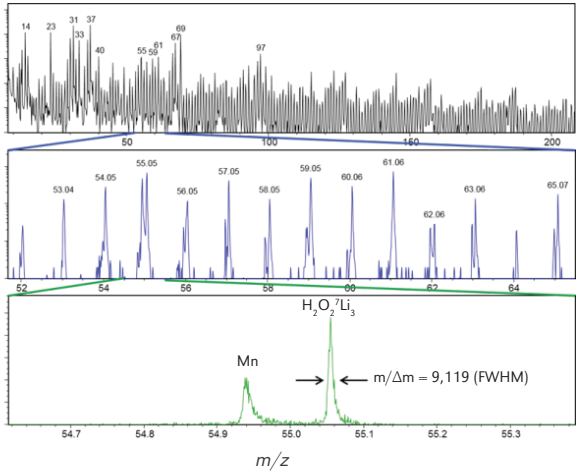
No compromise is best illustrated by the PHI cluster liquid metal ion gun (LMIG) that is designed to optimize performance in the HR² mode of analysis. This analysis mode provides high spatial resolution imaging and high mass resolution spectra in a single analysis. What's more, the HR² mode is realized using a high analytical beam current so that a typical analysis is conducted in only a few minutes. Now it is not necessary to choose between optimized imaging or spectroscopy; PHI's unique HR² mode of operation gives you both in a single rapid analysis.

FIB-TOF IMAGING

2D/3D Characterization of Challenging Specimens

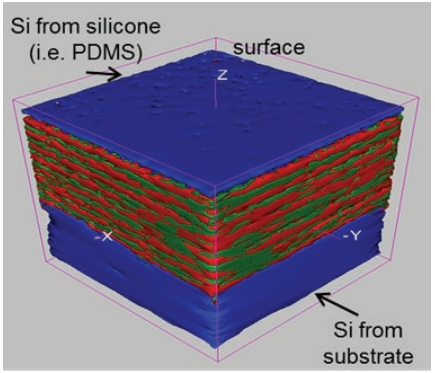
No compromise spectrometer design ensures that *in situ* sectioning and 3D tomography, via the dedicated PHI FIB option, results in superior TOF-SIMS chemical imaging. Porous and multi-phase materials which are not amenable to traditional sputter depth profiling are readily characterized by FIB-TOF analysis. Full range mass spectra, collected at high mass resolution, high mass accuracy and with excellent signal-to-background (S/B), enable robust chemical imaging of the FIB sidewall. The quality of the spectrum over the entire analytical mass range, combined with uniform collection efficiency over the depth of the FIB section, provides an unmatched capability for high resolution 2D and 3D imaging.

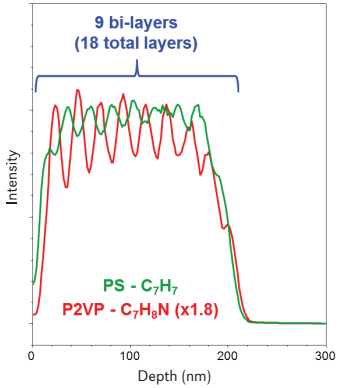




A TOF-SIMS image, collected in 5 minutes using the HR^2 mode of analysis, of a FIB-sectioned lithium ion battery anode showing the distribution of $C_6H_{11}O^+$ (*m/z* 99) and $C_7H_{15}O^+$ (*m/z* 115) ions. The lateral resolution was measured to be ≤ 150 nm in both ion polarities. The spectra demonstrate excellent signal-to-background and high mass resolution for full chemical and isotopic characterization.

GCIB CLUSTER ION BEAM





A depth profile of a poly(styrene)/poly(2-vinyl pyridine) (PS/P2VP) copolymer blend approximately 200 nm-thick on a Si substrate. The profiles and 3D image data expose the 3-dimensional structure and composition.

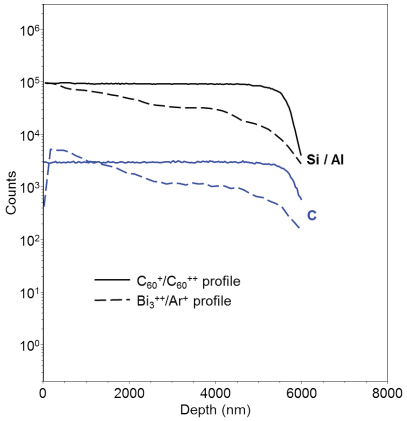
Non-Destructive Organic Depth Profiling

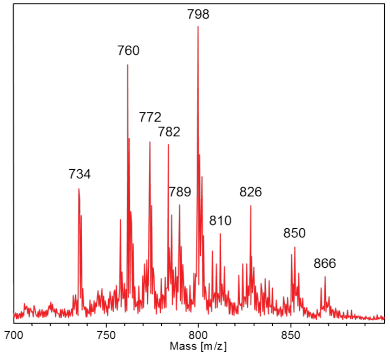
No compromise equips the analyst with tools for in-depth, non-destructive characterization of organic and molecular specimens without sacrificing resolution or molecular specificity. The PHI **gas cluster ion beam (GCIB)** option may be used for surface cleaning prior to analysis and for rapid 3D characterization to tens of microns in depth. For the characterization of multi-layer materials, the ultimate layer resolution may be realized while also making full use of the HR^2 mode for high resolution molecular imaging.

C_{60} CLUSTER ION BEAM

Profiling and Imaging with Minimum Artifacts

No compromise engineering provides advanced ion beams to address complex analytical problems. The PHI C_{60} gun option is a uniquely effective beam for sputter depth profiling of inorganic and mixed composition materials and may be used in an interleaved fashion as a beam for analysis. Interface transients and differential sputtering are greatly reduced with C_{60} compared to monatomic sputter ion beams. Additionally, the use of C_{60} for analysis provides 50x – 100x higher signal compared to Au_3 and Bi_3 due to reduced beam-induced damage.

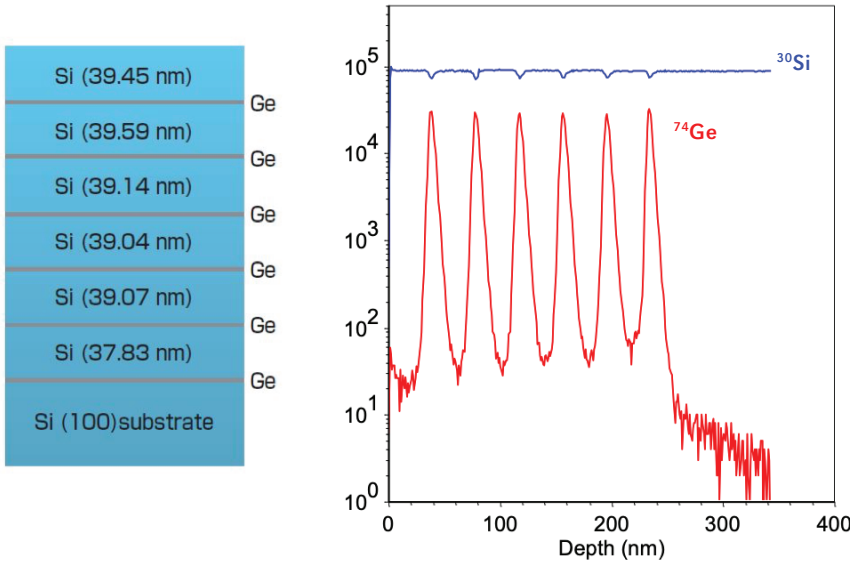




(LEFT) Depth profiles of a 6 µm-thick aluminosilicate sol-gel on copper by Ar and C_{60} sputtering. The C_{60} ion beam does not produce the differential sputtering artifacts produced by the Ar ion beam. Additionally, the C_{60} analysis beam does not produce the surface transient artifact that is produced by the Bi analysis beam.

(RIGHT) The C_{60} analysis beam delivers the greatest sensitivity for molecular imaging as demonstrated by the spectrum and maps of molecular lipids in a coronal rat brain cross-section.

DEPTH PROFILING WITH Ar^+/O_2^+ AND Cs^+ BEAMS



A depth profile of a multi delta layer (MDL) sample of Ge in Si was acquired using a 30 keV Bi^+ primary ion beam (50 µm square raster) and a 1 keV Cs^+ sputter ion beam (200 µm square raster). The Ge delta layers are 0.4 nm thick.

High Current Density with Rapid Low-Voltage Sputtering

No compromise is demonstrated by PHI's 2 kV Cs^+ sputter beam column. The column is designed and optimized to deliver maximum current at low beam voltage into a small beam diameter. The increased current density improves the sputter efficiency for rapid depth profiling even at low beam voltage. The low accelerating voltage is required for accurate characterization of today's electronic designs which incorporate shallow junctions and dopant implants because at low voltage the interlayer mixing is minimized. What's more, the improved current density enables profiling of single features, such as bond pads, without eroding nearby features.

PHYSICAL ELECTRONICS
A DIVISION OF ULVAC-PHI

Physical Electronics - USA

Phone: 952-828-6100
Email: sales@phi.com
Web: www.phi.com

ULVAC-PHI Inc. - Japan

Phone: 81-467-85-4220
Email: webmasterjp@phi.com
Web: www.ulvac-phi.co.jp

Standard Features

- Parallel Imaging MS/MS mass analyzer
- 30 kV LMIG with Bi, Au, or Ga emitter
- Dual beam charge neutralization
- 5 axis sample stage
- *In situ* optical viewing
- Secondary electron detector
- SmartSoft-TOF instrument control and TOF-DR data reduction software packages
- Analysis chamber with four primary ion gun ports
- High throughput turbo molecular pump
- Integrated bakeout facilities

Optional Accessories

- 20 kV C_{60} pulsed ion gun
- 20 kV Ar_{2500} gas cluster ion gun
- 2 kV Cs ion gun
- 5 kV gas gun (Ar/O_2)
- Oxygen flood module
- 30 kV Ga FIB gun
- Hot/Cold sample stage module
- Flash cooling for sample intro chamber
- High temperature sample stage module
- Fast sample rotation module
- Sample transfer vessel
- Intro chamber glove box
- Voltage cycling sample stage module
- Sample preparation chambers
- XPS/AES Transfer Adapter