



Sino-Austrian Biomarker Research Center



o.Univ.-Prof. Mag. Dr. Günther Bonn



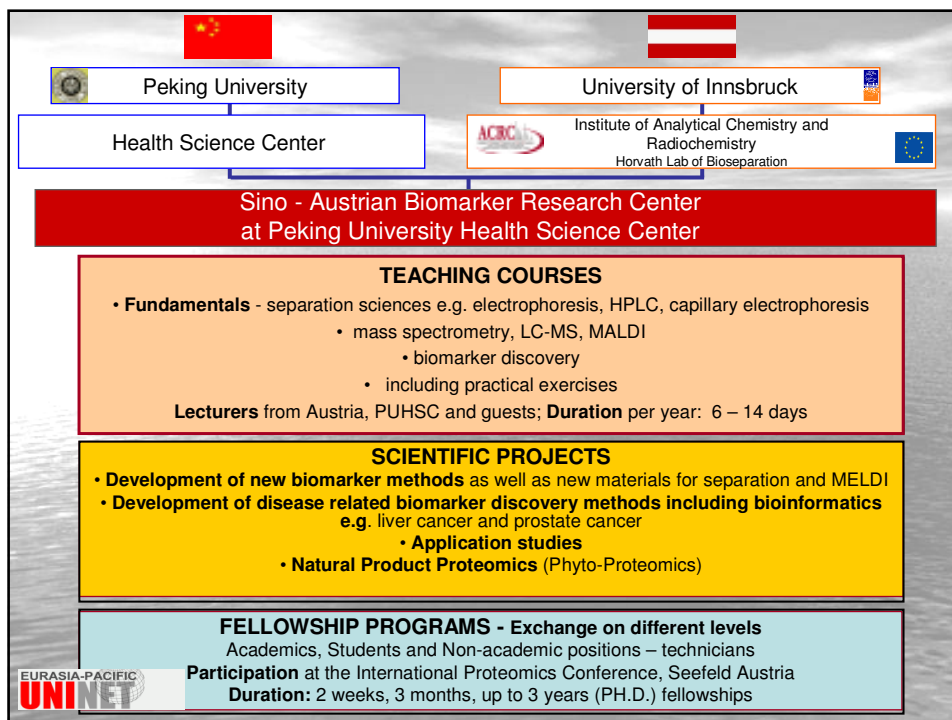
北京大学医学部
Peking University
Health Science Center

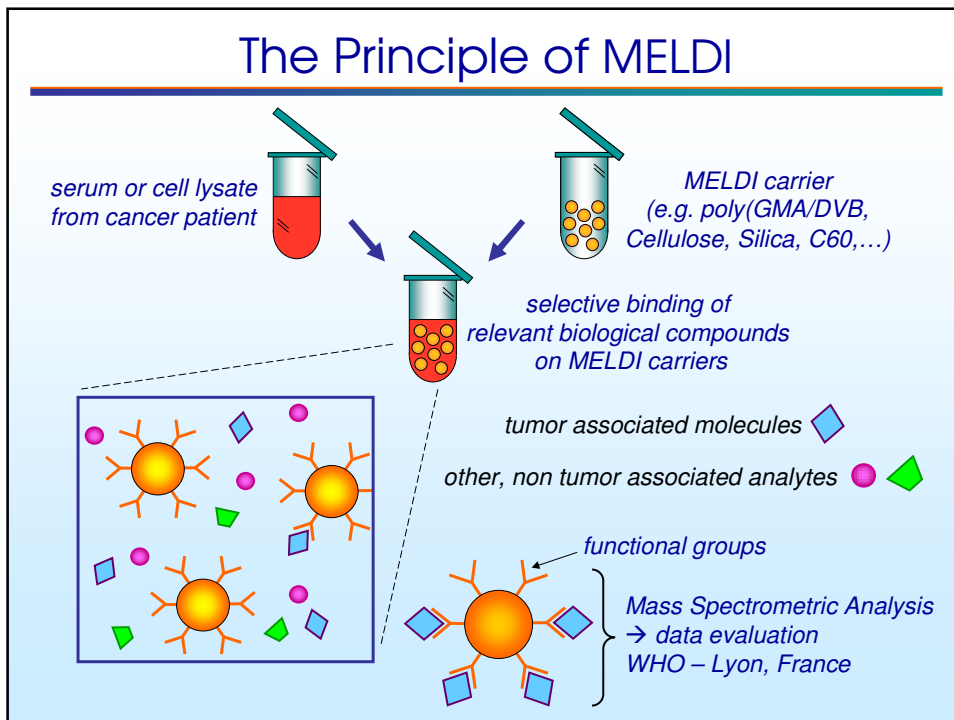
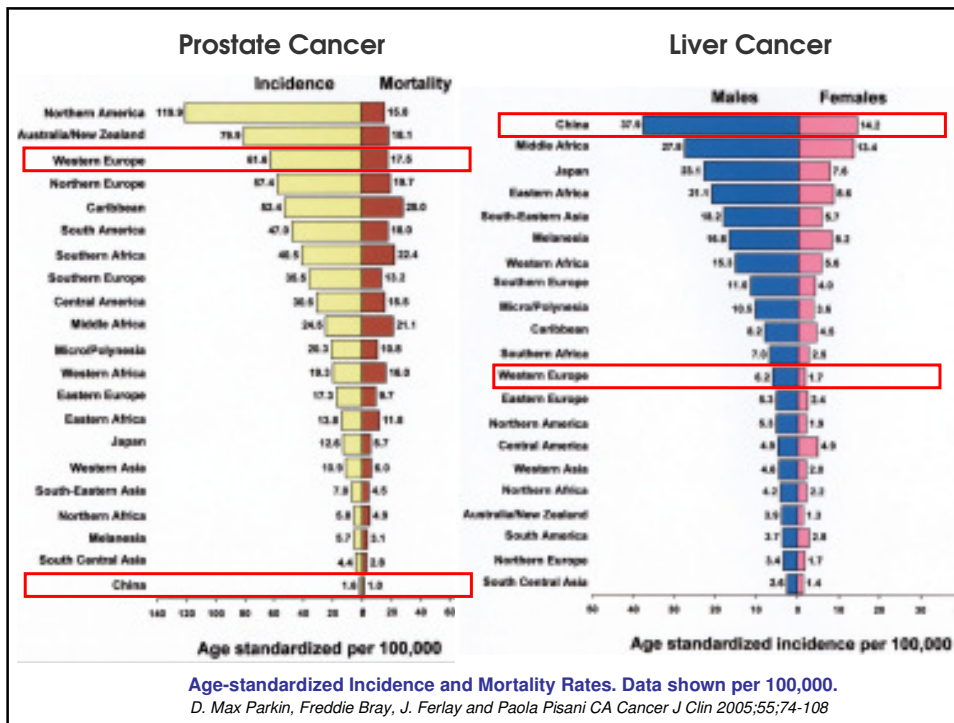


Opening of the
Sino-Austrian Biomarker Research Center
at the 25th May 2006

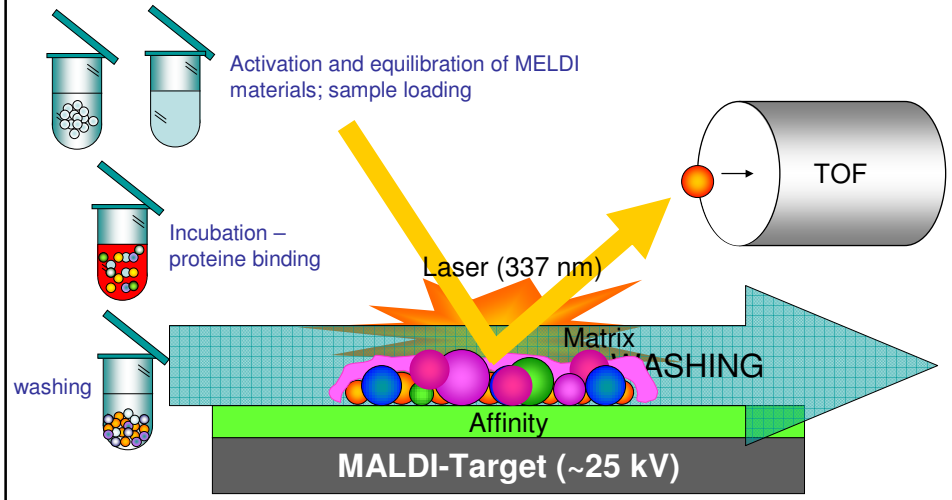


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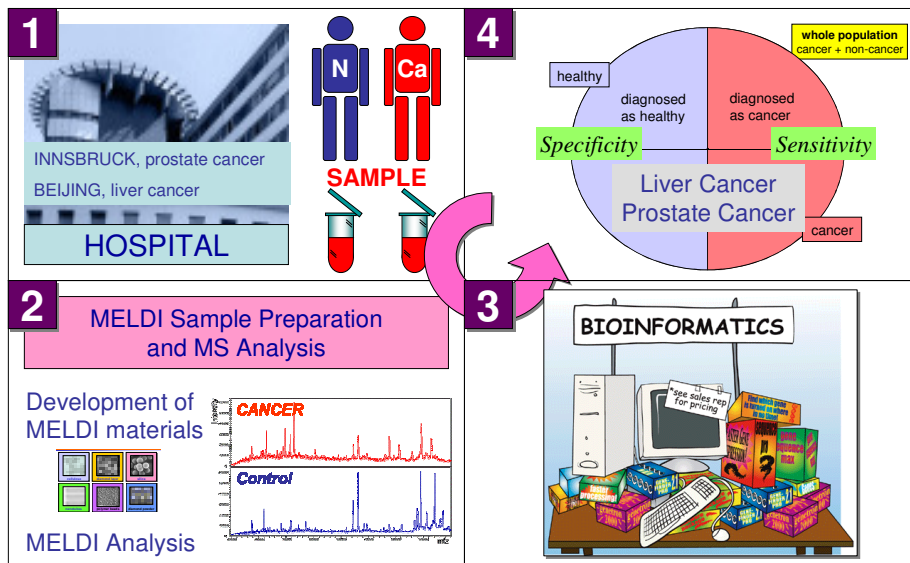


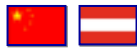
MELDI Sample Preparation



Feuerstein, Isabel; Najam-ul-Haq, Muhammad; Rainer, Matthias; Trojer, Lukas; Bakry, Rania; Aprilita, Nurul Hidayat; Stecher, Guenther; Huck, Christian W.; Bonn, Guenther K.; Klocker, Helmut; Bartsch, Georg; Guttman, Andras. JASMS (2006), 17(9), 1203-1208.

The Principle of MELDI





Synthesis of GMA/DVB Polymers for MELDI-MS



RAPID COMMUNICATIONS IN MASS SPECTROMETRY
Rapid Commun. Mass Spectrom. 2006; 20: 2954–2960
Published online in Wiley InterScience (www.interscience.wiley.com) DOI: 10.1002/rcm.2673

RCM

Ultra-fast mass fingerprinting by high-affinity capture of peptides and proteins on derivatized poly(glycidyl methacrylate/divinylbenzene) for the analysis of serum and cell lysates

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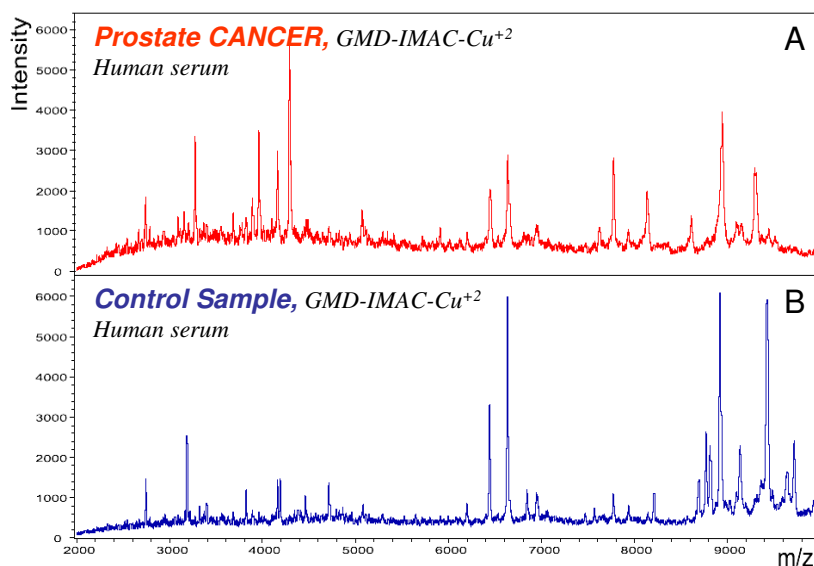
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Received 12 April 2006; Revised 4 July 2006; Accepted 18 July 2006

The development of support materials in mass fingerprinting is an important task required for diagnostic markers in conjunction with matrix-assisted laser desorption/ionization time-of-flight mass spectrometry (MALDI-TOF MS). The material-based approach, which we introduced as material-enhanced laser desorption/ionization (MELDI), focuses not only on different functionalities, but also emphasizes the morphology, i.e. porosity or particle size of the carrier material. As a result, it provides a quick and sensitive platform for effective binding of peptides and proteins out of different biofluids, e.g. serum, spinal fluid, urine or cell lysates, and to subsequently analyze them with MALDI-TOF MS. This approach includes a built-in desalting step for serum protein profiling and is sensitive enough to detect proteins and peptides down to 100 fmol/μL. Here we co-polymerized glycidyl methacrylate (GMA) with divinylbenzene (DVB) using thermal polymerization to yield a

Application of GMA/DVB to screen Prostate Cancer Serum Samples



Biomarker Study on Prostate Cancer

Genetic Algorithm GA
Model 1



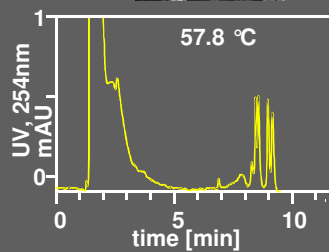
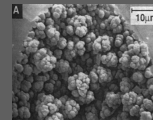
Class	Name	Correct Classified Part of Valid Spectra	1	2	0
1	Control	sensitivity 97.1 %	34	1	0
2	Prostate Cancer	specificity 98.1 %	1	51	0

Mutation Detection by Denaturing HPLC using Packed Columns and Monolithic Capillaries



**PS-DVB C18 stationary phase
HPLC-Equipment**
→ (worldwide ca. 3000 pieces sold)

Monolithic Capillary



column: 50 x 0.2 mm I.D.
flow rate: 3.0 µl/min
injection vol.: 500 nl

Collaborations



北京大学医学部
Peking University
Health Science Center



International Agency for Research on Cancer
Centre International de Recherche sur le Cancer

WHO – Lyon, France