Technical University of Denmark



## Laser ablation dynamics and production of thin films of lysozyme

Canulescu, Stela; Schou, Jørgen; Amoruso, S.; Wang, X.; Bruzzese, R.; Matei, A.; Constantinescu, C.; Dinescu, M.

Publication date: 2012

Link back to DTU Orbit

Citation (APA):

Canulescu, S., Schou, J., Amoruso, S., Wang, X., Bruzzese, R., Matei, A., ... Dinescu, M. (2012). Laser ablation dynamics and production of thin films of lysozyme. Abstract from E-MRS 2012 Spring Meeting, Strasbourg, France.

## DTU Library Technical Information Center of Denmark

#### **General rights**

Copyright and moral rights for the publications made accessible in the public portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

• Users may download and print one copy of any publication from the public portal for the purpose of private study or research.

- You may not further distribute the material or use it for any profit-making activity or commercial gain
- You may freely distribute the URL identifying the publication in the public portal

If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

# Abstract: Laser ablation dynamics and production of thin films of lysozyme

S. Canulescu, J. Schou (1), S. Amoruso, X. Wang, R. Bruzzese (2), A. Matei, C. Constantinescu, M. Dinescu (3).

### (1)DTU Fotonik, Risø Campus, DK-4000 Roskilde, Denmark

(2)Dipartimento di Scienze Fisiche & CNR-SPIN, Università degli Studi di Napoli Federico II, I-80126 Napoli, Italy

(3)National Institute for Lasers, Plasma and Radiation Physics, RO-077125 Magurele-Bucharest, Romania Lysozyme is a well-known protein, which is used in food processing because of its bactericidal properties. The mass (14307 amu) is in the range in which it easily can be monitored by mass spectrometric methods, for example by MALDI (Matrix assisted laser desorption ionization). We have recently produced thin films of average thickness up to 300 nm, which not only contained a significant amount of intact molecules, but also maintained the bioactivity. These films were produced by a nanosecond laser in the UV regime at 355 nm with 2 J/cm<sup>2</sup>. The surprising fact that these molecules can be transferred to a substrate as intact molecules by the violent laser impact (~up to 50 mJ/pulse) has not yet been understood. One issue is that up to 150 ng/pulse is removed by the laser, and much of the material is ejected from the target in relatively large chunks.

We have explored as well the excitation mechanics by laser impact. Samples of pressed lysozyme prepared in the same manner as in ns-experiments have been irradiated at 527 nm with 300-fs pulses and at at similar fluence as in ns ablation. Even though the pulse energy was much smaller, there was a considerable ablation weight loss of lysozyme from each shot. This is the first time the ablation by fs-lasers of a protein has been recorded quantitatively. Films of lysozyme produced by fs-laser irradiation were analyzed by MALDI and a significant number of intact molecules in the films with fs-laser deposition was found as well.