

NOTIZEN

Electron-Beam Induced Conformational Changes in Polypeptide Layers: an Infrared Study

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Langmuir-Blodgett type polypeptide monolayers have been irradiated under conditions simulating the hazards of electron microscopic imaging. The damaging effect, *i. e.* the randomization of the α -helical layers, which occurs in the dose range between 50 and 200 e^-/nm^2 , has been monitored by infrared dichroism.

The beam of an electron microscope is besides being a flux of short wavelength radiation suitable for high resolution structure determination, also an intense flux of ionizing radiation which inevitably

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deteriorates the structure under investigation. For organic and biomolecules radiation damage is the most serious obstacle in approaching resolution levels which allow to deduce their molecular structure. Meaningful high resolution information is only to be expected if the damaging effects can be minimized. This in turn requires a thorough knowledge of the damaging effects involved, which at present does not exist.

In an attempt to elucidate the especially intriguing response of proteins to irradiation^{1–3} we have concentrated our efforts onto relatively simple polypeptide systems. It has been shown in the pioneering work of Malcolm^{4,5} and Loeb⁶ that some high molecular weight synthetic homopolypeptides can be spread as monolayers at water-air interfaces and deposited onto solid supports by conventional Langmuir-Blodgett-techniques. The possibility to manipulate the conformation assumed at the interface by changing the spreading solvent and the liquid subphase⁷ as well as the high degree of long range order in these monolayers make them attractive for studying the effects of radiation damage on the three basic secondary structures of proteins, *i. e.* to measure the relative sensitivities of α -helices, β -structures and random coil structures. Infrared spectroscopy is diagnostic for transitions between these conformations and has the advantage that it can be performed under solid state conditions, *i. e.* under conditions relevant to electron microscopy. With the ATR-IR-technique⁸ radiation

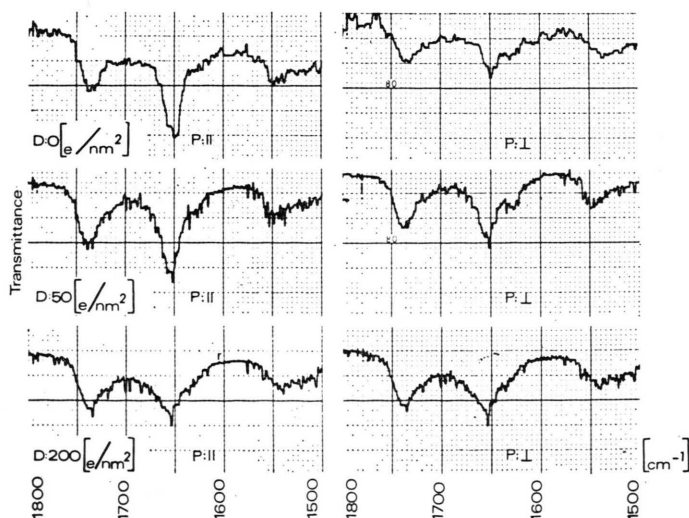


Fig. 1. Polarized ATR-IR spectra of poly- α -methyl-L-glutamate bilayers (P: || parallel polarized light; P: $\perp$ perpendicular polarized light). The strong polarization of the amide I band ($\sim 1650 \text{ cm}^{-1}$) in the non-irradiated control reflects the long range order in the α -helical layers. An electron dose of 50 e^-/nm^2 significantly diminishes the dichroic ratio and with 200 e^-/nm^2 absorption is almost isotropic, *i. e.* the layers are completely randomized.

damage effects can be measured in extremely thin (~ 2 nm) layers⁹.

The polypeptide chosen for this study was poly- α -methyl-L-glutamate which was spread from chloroform/dichloroacetic acid (10:1) on water and transferred to germanium internal reflection plates at film pressures closely below the collapse pressure ($18 \text{ dyn}\cdot\text{cm}^{-1}$). The pronounced infrared dichroism reflects the high degree of long-range order of the α -helices. These layers were irradiated with electron doses between $0-400 \text{ e}^-/\text{nm}^2$ simulating the imaging conditions of electron microscopy. The degradation of order was followed by measuring the decrease of the dichroic ratio of the initially sharp amide I ($\sim 1650 \text{ cm}^{-1}$) band. Though we have not yet attempted a quantitative band-shape analysis it becomes immediately apparent from the polarized spectra (Fig. 1) that the polypeptide layers are randomized by doses as low as $50 \text{ e}^-/\text{nm}^2$, which is comparable to the critical dose for lipid layers⁹. It should be kept in mind that this dose is approximately 2 orders of magnitude lower than the minimum dose needed for single heavy atom detection in monolayers¹⁰ and at least 3 orders of magnitude

lower than the doses conventionally applied to obtain high resolution pictures of isolated protein molecules¹¹. This stresses the importance of matching the e.m. resolution to a level governed by the radiation sensitivity of the specimen¹¹.

It should be mentioned in this context that the investigation of the structure of polypeptide layers has still another impact upon electron microscopy: The high degree of long-range order in polypeptide films built up at interfaces lets seem it feasible to design spatially defined charge patterns at interfaces, by spreading polypeptides with charged side chains. These charge patterns could serve as templates for adsorbing proteins in a regular fashion which would allow to perform their structural analysis at significantly lower doses as if they were in a random orientation.

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