BioPad

Textile Engineering

the functional textile connection

PhotoBonding Technology

Key features for biomolecule immobilization by light

- No surface functionalization required
- No probe molecule functionalization required
- Biomolecule surface density solely depends on irradiation intensity (dose and time)
- Co-immobilization of stabilizing agents possible
- Cost-effective single-step process
- Probe molecule orientation: 50 % right-side up (statistical)

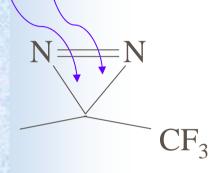
PhotoBonding Technology

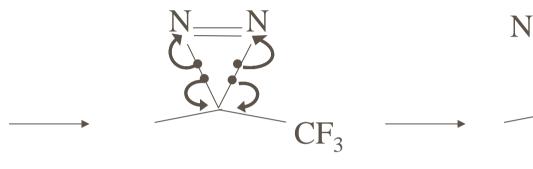
Key features for dextran-based photolink-polymer

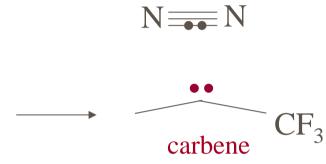
- Hydrophilic
- Biocompatible
- Surface passivation (suppress non-specific binding, suppression of bacterial deposition)
- Bi-functional polymeric cross-linker (heterofunctions for secondary modifications)

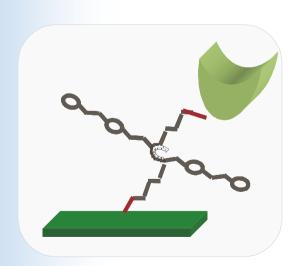
$\lambda = 350 \text{ nm}$

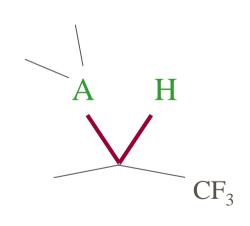
PhotoBonding: Carbene - Chemistry

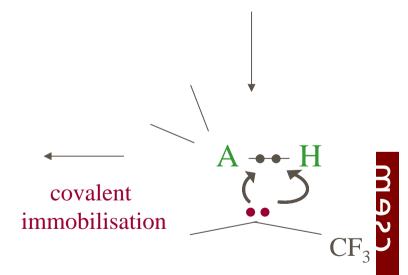






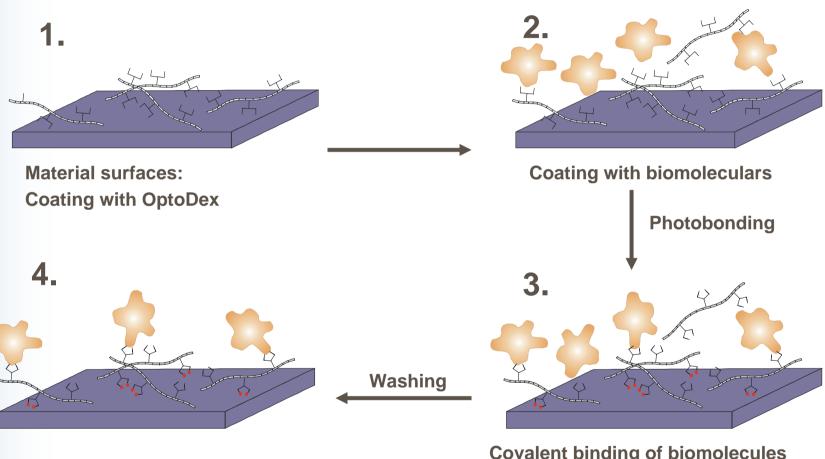






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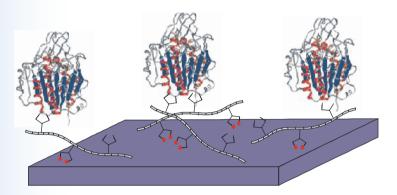
PhotoBonding: Proceeding



Covalent binding of biomolecules on matrial surfaces

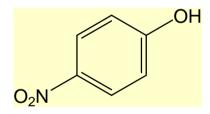
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Photobonding of Alk. Phosphatase on Tissue



Photobonded Alk. Phosphatase on tissue

pNPP: a substrate for detecting Alk. Phosphatase (colorless) A referenced experiment



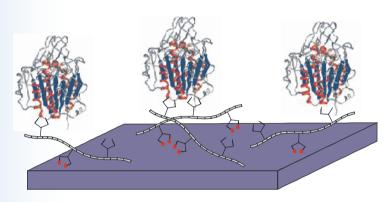
When the Alk. Phosphatase and pNPP reacted, a yellow-water-soluble reaction product is formed, it absorbs light at 405 nm

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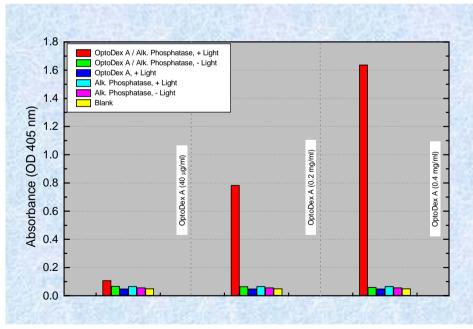
Photobonding of Alk. Phosphatase on Tissue

A referenced experiment



- OptoDex / Alk, Phosphatase, + Light
- OptoDex / Alk, Phosphatase, Light
- OptoDex, + Light
- Alk, Phosphatase, + Light
- Alk, Phosphatase, Light
- Background

Photobonded Alk. Phosphatase on tissue (polyester)



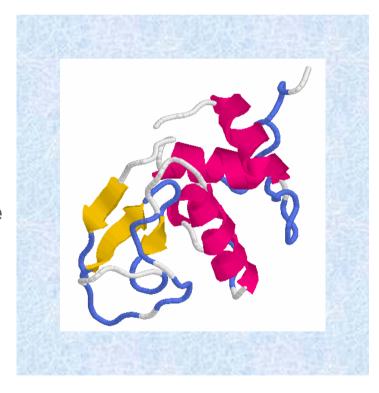
Lysozyme:

a ubiqutous enzyme present in human serum, urine, seminal fluid and milk.

widespead distribution in aminals and plants

a 'natural antibiotic', attacks gram-positive bacteria (by breaking sugar backbone of peptidoglycan components in cell wall)

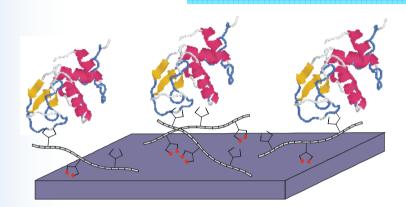
suppresses inflammation



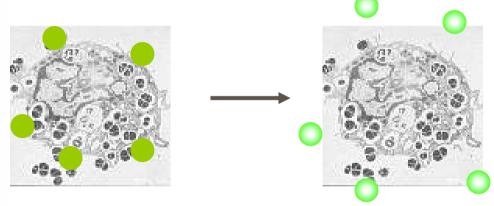
Lysozyme activity assay

Lysozyme cleaves the cell wall and relieves the quenching, yielding a dramatic increase in fluorescence that is proportional to lysozyme activity.

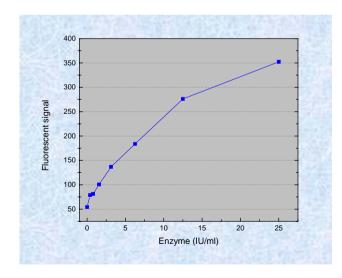
The fluorescence increase can be measured with the spectrofluorometer.



Photobonded lysozyme on material surface



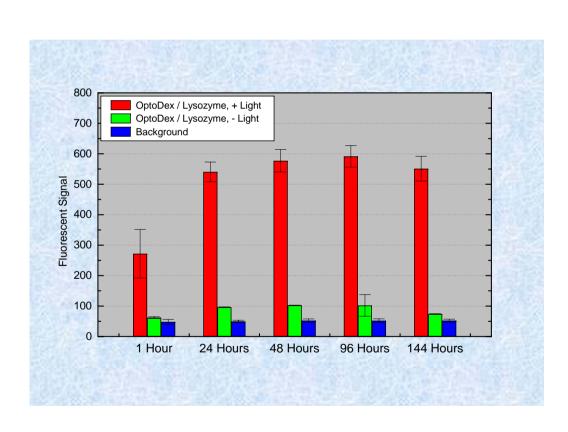
Micrococcus lysodeikticus: are labeled to such a degree that the fluorescence is quenched.



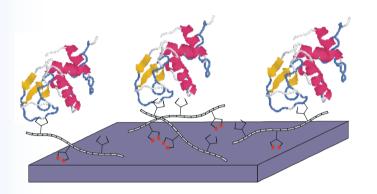
A mimic experiment A

Photobinding of lysozyme on microtiter plate (polystyrene)

Detection of remained enzymatic activity with fluorescent-*Micrococcus lysodeikticus*

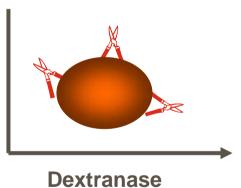


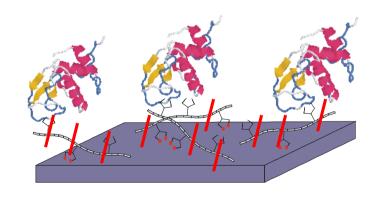
A mimic experiment B



Dextranase degradates photolinker polymer by catalyzing the endohydrolysis of 1,6- α -glucosidic linkages in dextran.

Then photoboned lysozyme is released



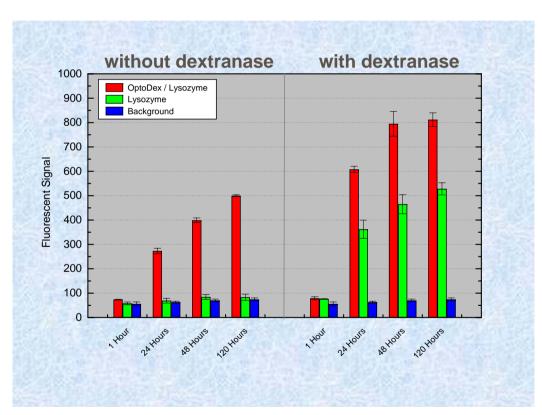


A mimic experiment B

Photobinding of lysozyme on microtiter plate (polystyrene)

Degradation of OptoDex by dextranase

Detection of remained enzymatic activity with fluorescent-*Micrococcus lysodeikticus*



Preliminary experiment:

photobonding of lysozyme on the tissue (polyester)

