# Exposure Sensitivity to Biofunctionalized Polymer-Based Nanoparticles

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#### **Bacterial Binding to Host is Mediated by Adhesins**





Transmission electron micrograph of *E. coli* adhering to epithelium in the intestine of a pig.

Moon, H.W. 1997. Comparative histopathology of intestinal infections. In: Mechanisms in the pathogenesis of enteric diseases (P.S. Paul, D.H. Francis and D.A. Benfield, eds.) Adv. Exptl. Med. Biol. 412:1. Plenum Press, New York.

### **Bacterial Cell Binding Strategies**



High NP Concentration: Bacterial Isolation Intermediate NP Concentration: Bacterial Agglutination Low NP Concentration: Bacterial Tagging

### Nanoparticle Chemical Structure: Mannose Functionalization



### **Nanoparticle Design Strategy**



Functionalized PEG side chains extending from hydrophobic polymer backbone chain.

Diagram illustrates the self assembly into the nanoparticles followed by photochemical curing.

Nanoparticle

# *E. coli* - NP Interaction

TEM images (dark-field) showing the agglutination of *E. coli* ORN178 mediated by D-mannose-tethered nanoparticles

(a,b) Lower magnification and (c,d) higher magnification

(e) *E. coli* ORN178 only (similarly with bare nanoparticles)

(f) *E. coli* ORN208 with the same D-mannose-tethered polymeric nanoparticles.



Acute Nanoparticle Exposure Sensitivity Studies

- In vitro studies

   cell toxicity studies
- In vivo studies
  - -Skin (rabbit)
  - Ocular (rabbit)
  - Inhalation (rat)
  - Ingestion (rat)
- In vivo studies: poultry

### **In Vitro Results: Dermal Fibroblasts**

1 ml cells + medium / 50 μl 2wt% np solution (core-PEG np)P = proliferating cells;NonP = nonproliferating cellsnp = with nanoparticles;C = control (w/o np)

	Total Cell Count			
<u>Trial</u>	<u>P( C)</u>	<u>P(np)</u>	<u>NonP(C)</u>	<u>NonP(np)</u>
Mean (N=4):	95,625	95,000	316,875	281,875
95%CI:	29,476	28,865	86,619	35,779
p value:	0.963 (not significant)		0.300 (not significant)	

## Dermal Test: Mannan Nanoparticles



#### Site preparation



Applying gauze



#### Application of dose (1 mL, 2.0 wt.%)



Overview after procedure

### Results: Dermal Test (48 hrs)







### **Ocular Test: Mannan Nanoparticles**





Right and left eye before procedure



Application of dose (0.1mL at 2.0 wt.%)



Right eye 1 min. after dose

### Results: Ocular Test



### 48 hr.





### Inhalation Studies: FITC-labeled Mannan-NP



### Inhalation Study: Lung Tissue (fluorescence) 72 hr.

Alveolar Sac / Alveolar duct



Control (200x)

**Test (200x)** 

### Inhalation study: Lung Tissue (H&E stain)

#### Alveolar Sac / Alveolar duct



#### Control (1000x)

**Test (1000x)** 

Dark spots are nuclei of endothelial and connective tissue cells. Red spots are red blood cells. No detectable difference.

### Ingestion Studies: FITC-labeled Mannan-NP



#### **Oral Ingestion:** Small Intestine Tissue (H&E stain) 72 hr.

#### **Transverse sections**







#### Test (400x)

No apparent difference.

### **Oral Ingestion: Kidney (H&E stain) 72 hr.**



#### Control (400x)

Glomerulus

**Test (400x)** 

No apparent difference.

### **Oral Ingestion:** Liver (H&E stain) 72 hr.





#### Control (200x)

#### **Test (200x)**

#### No apparent difference.

# **Poultry Studies**

- 1-2 poults/pen gavaged with 0.1, 0.5 or 1.0 mL per day of core-PEG nanoparticles, 2wt.%.
- 3 control poults/pen gavaged with distilled water
- Body weights at 1, 3 and 6 wk; observation to 14 wk
- Commercial feed and water *ad libitum*

### Poult Performance: 6-week Body Weight



No significant effect of nanoparticles on poult body weight.

# **Concluding Remarks**

- In vitro & in vivo studies conducted with polystyrene-based nanoparticles.
- No adverse cellular response for dermal fibroblast cells.
- No apparent adverse tissue response from dermal, ocular, inhalation, or ingestion routes of exposure.
- No adverse growth response from poultry studies.
- Further in vitro and in vivo studies planned.

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