



Division of Bacterial, Parasitic and Allergenic Products

Jay E. Slater, MD

Director, DBPAP

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Division of Bacterial, Parasitic and Allergenic Products (DBPAP)

- Merger (1999) of
 - Division of Bacterial Products (DBP) and
 - Division of Allergenic Products and Parasitology (DAPP)
- Leadership
 - Drusilla Burns, PhD (Acting) 1999-2000
 - Richard Walker, PhD 2000-2008
 - Milan Blake, PhD 2008-2010
 - Jay Slater, MD 2010-present



Division of Bacterial, Parasitic and Allergenic Products (DBPAP)

Immediate Office of the Director
Jay E. Slater, MD – Director
Drusilla Burns, PhD – Deputy Director
8 FTE

Laboratory of Bacterial Polysaccharides
Willie Vann, PhD – Chief
Mustafa Akkoyunlu, MD, PhD
Margaret Bash, MD
Wei Wang, PhD
John Cipollo, PhD
Daron Freedberg, PhD
19 FTE, 9 ORISE

Laboratory of Respiratory and Special Pathogens
Michael Schmitt, PhD – Chief
Drusilla Burns, PhD
James Keller, PhD
Tod Merkel, PhD
18 FTE, 1 ORISE

Laboratory of Immunobiochemistry
Ron Rabin, M.D. – Chief
Jay E. Slater, MD
10 FTE, 7 ORISE

Laboratory of Mucosal Pathogens & Cellular Immunology
Scott Stibitz, Ph.D. – Chief
Paul Carlson, PhD
Siobhán Cowley, PhD
Karen Elkins, PhD
19 FTE, 6 ORISE

DBPAP regulatory/research portfolio

Non-invasive, toxin producers

- *Bacillus anthracis*
- (*Bordetella parapertussis*)
- *Bordetella pertussis*
- (*Clostridium botulinum*)
- *Clostridium tetani*
- *Corynebacterium diphtheriae*
- (*Clostridium difficile*)

Invasive, protective responses to polysaccharides

- *Haemophilus influenzae*
- *Neisseria meningitidis*
- *Streptococcus pneumoniae*

Intracellular

- (*Francisella tularensis*)
- *Mycobacterium tuberculosis*
- (*Mycobacterium bovis*)

Enteric

- (*Campylobacter jejuni*)
- *Salmonella typhi*
- (*Salmonella typhimurium*)
- (*Shigella dysenteriae*)

Parasite

- (*Plasmodium spp*)
- (*Leishmania donovani*)

Other/emerging

- (*Yersinia pestis*)
- (*Staphylococcus aureus*)
- Allergenic products
- (Probiotics)
- (Fecal microbiota)
- (Bacteriophage)

DBPAP regulatory/research portfolio

Bacterial Polysaccharides (LBP)

Non-invasive, toxin producers

- *Bacillus anthracis*
- (*Bordetella parapertussis*)
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Bacterial polysaccharides (LBP)

- Akkoyunlu
 - Regulation of humoral immune response against polysaccharide vaccines by bacterial products
- Bash
 - To present protein targets as alternatives based on molecular epidemiology of the pathogen where polysaccharide vaccines have lacked success
- Cipollo
 - Mass spectrometry of carbohydrates as a tool for characterization of bacterial vaccines and pathogens
- Freedberg
 - Improvement of biological product quality by application of new technologies to characterize of vaccines and blood products: NMR spectroscopy and light scattering
- Vann
 - To determine the metabolic pathway for the synthesis of polysaccharides in gram negative pathogens and to use this knowledge to development methods for preparing glycoconjugate vaccines using metabolic engineering
 - To improve and understand the manufacture of current conjugate vaccines through the investigation of conjugation chemistry
- Wang
 - To investigate the genetics of pathogenesis of gram negative bacteria. *M. catarrhalis* is a model organism for these studies.
 - To study the regulation of *M. catarrhalis* virulence gene expression and identify biomarkers for developing vaccine against *M. catarrhalis*.

DBPAP regulatory/research portfolio

Immunobiochemistry (LIB)

Non-invasive, toxin producers

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- *Corynebacterium diphtheriae*
- (*Clostridium difficile*)

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- *Neisseria meningitidis*
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- (*Mycobacterium bovis*)

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Parasite

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- (*Leishmania donovani*)

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- (*Staphylococcus aureus*)
- **Allergenic products**
- (Probiotics)
- (Fecal microbiota)
- (Bacteriophage)

DBPAP regulatory/research portfolio

Mucosal Pathogens and Cellular Immunology (LMPCI)

Non-invasive, toxin producers

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- *Bordetella pertussis*
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- *Clostridium tetani*
- *Corynebacterium diphtheriae*
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- *Mycobacterium tuberculosis*
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- Probiotics
- Fecal microbiota
- Bacteriophage

Mucosal Pathogens and Cellular Immunology (LMPCI)

- Stibitz
 - Molecular genetics of virulence and virulence-gene regulation in bacterial respiratory pathogens
- Elkins
 - Define mechanisms by which vaccine-induced immune T cells provide protection against intracellular bacteria, and thus establish practical immune correlates
- Cowley
 - Characterization of unique aspects of mucosal immune responses to intracellular pathogens.
- Carlson
 - Molecular aspects of *C difficile* pathogenesis
- Malaria program – collaboration with OBRR

DBPAP regulatory/research portfolio

Respiratory and Special Pathogens (LRSP)

Non-invasive, toxin producers

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- *Bordetella parapertussis*
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- *Clostridium tetani*
- *Corynebacterium diphtheriae*
- (*Clostridium difficile*)

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Respiratory and Special Pathogens (LRSP)

- Schmitt
 - Characterization of iron transport systems of *Corynebacterium diphtheriae*
 - Analysis of gene regulation in *C. diphtheriae*
- Burns
 - Bacterial toxin structure/function relationships
 - Study of host response to immunization with anthrax and *Staphylococcus aureus* vaccines
- Keller
 - Clostridial neurotoxin vaccine and antitoxin research
- Merkel
 - Development of animal aerosol challenge models for pertussis, anthrax and disease caused by *Staphylococcus aureus*.
 - Analysis of gene regulation in *Bordetella pertussis*

LIB presenters

- Ronald L. Rabin, MD, Lab Chief, Supervisory Medical Officer (PI)
- Philippa Hillyer, PhD, Staff Fellow
- Nikunj Sharma, PhD, Staff Fellow
- Jay E. Slater, MD, Supervisory Medical Officer (PI)