

US EPA ARCHIVE DOCUMENT

Mechanistic Indicators of Childhood Asthma (MICA) -Integrating Environmental, Clinical and Susceptibility Markers to Improve the Impact of Human Air Pollution Studies.

Jane E. Gallagher Ph.D.

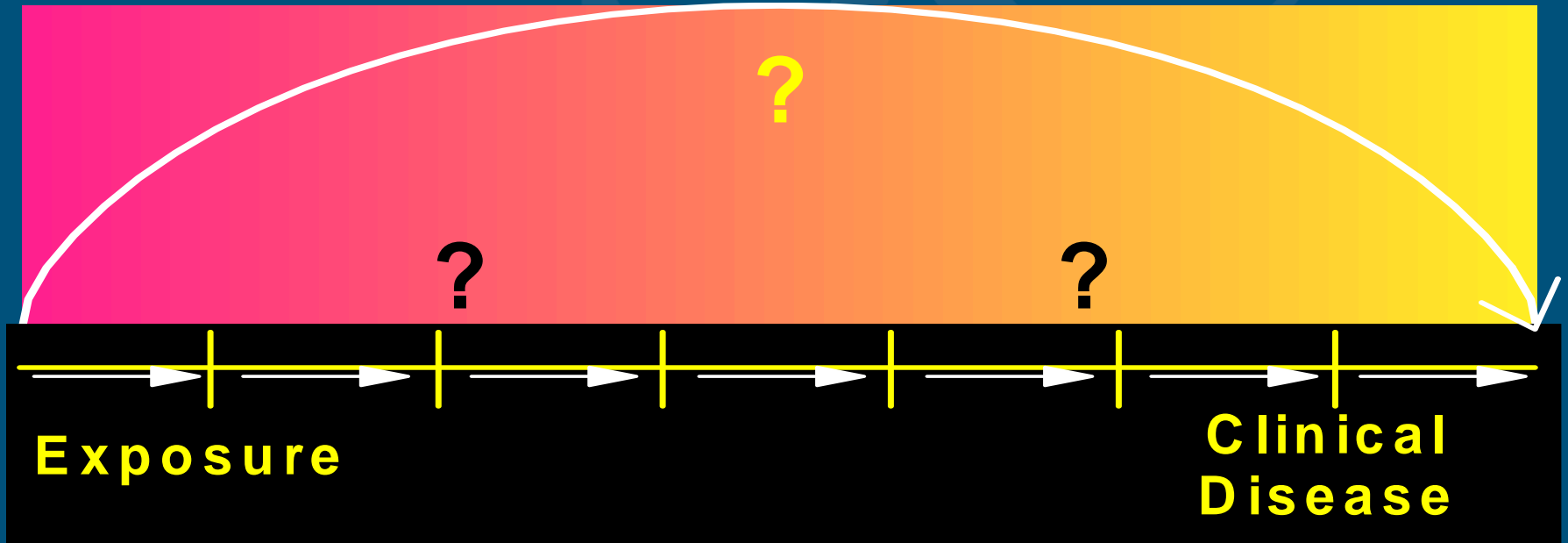
Human Studies Division

ORD NHEERL EPA

September 24-25 2007

Public Health Applications of Human Biomonitoring

Advances in biomarker development have improved our ability to detect early changes at the molecular and cellular level.



Battery of endpoints capturing Net effect—over several mechanisms of action/ classes

- Comet assay
- P53
- FISH
- 1-OH pyrene
- Cholinesterase inhibition
- Cross-linking metals formaldehyde
- HPRT
- DNA adducts methods
- **Mutagen sensitivity**
- **Oxidative damage**
- **Mutagenicity**

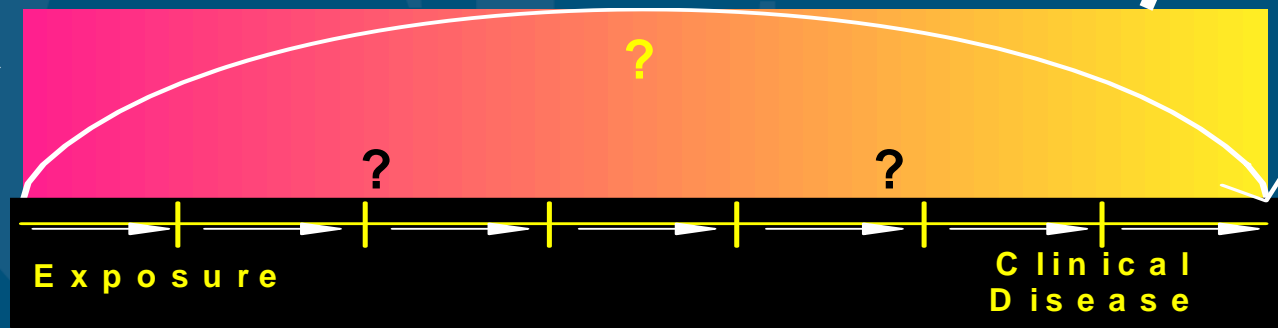
Integrated measure of dose across classes of chemicals

Evolving Technologies



Environment
Health Scientists

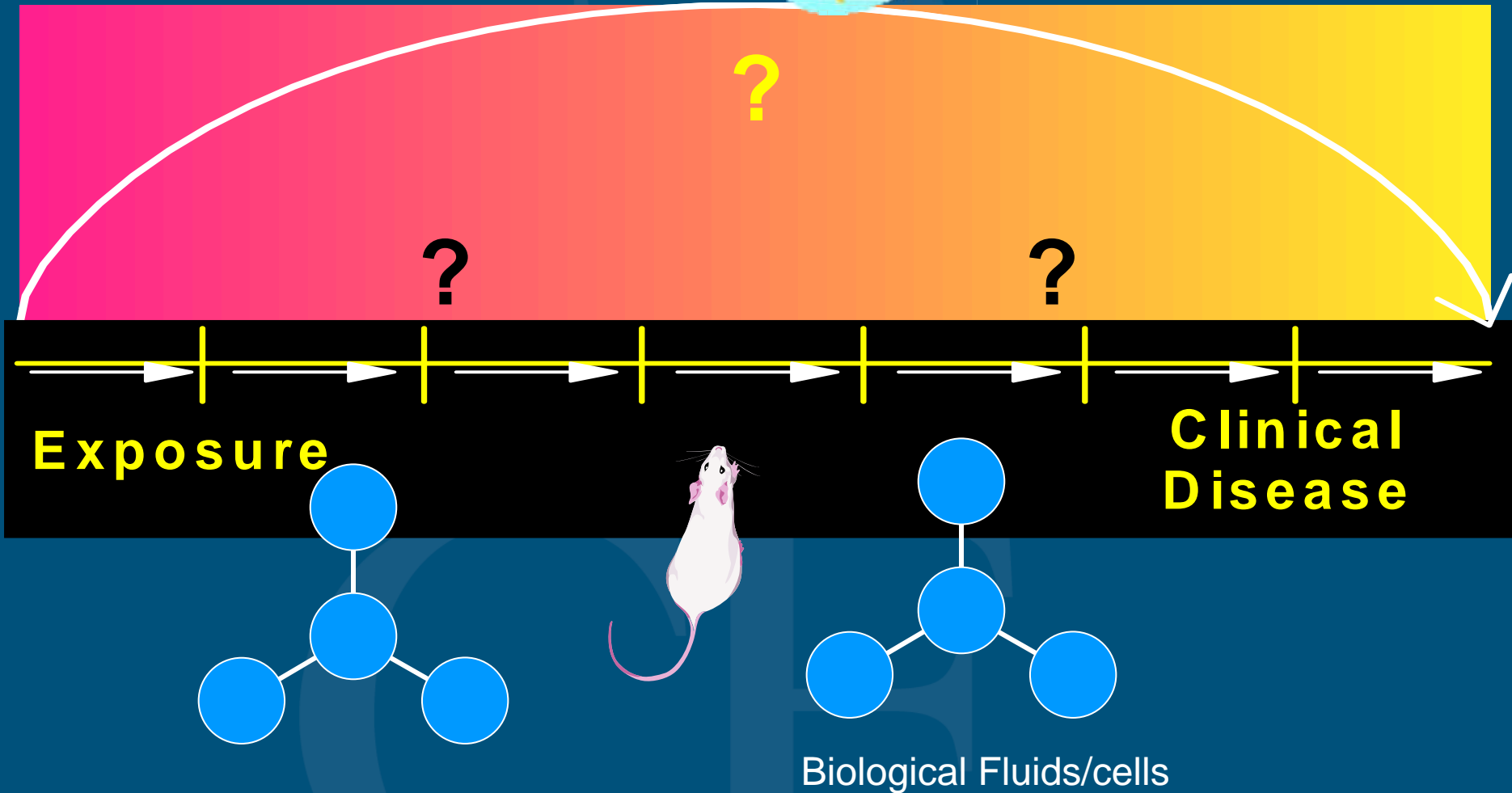
Clinicians



Environmental Studies Benefit from Knowledge Gained from Clinical Disease Studies (and visa versa)

- What factors affect a persons risk for a number of health conditions.
- Early indicators/detection of disease
- Identify Genetic variants that increase susceptibility
- Determine whether the effect of genetic variants that increase risk is different in the presence of environmental exposures

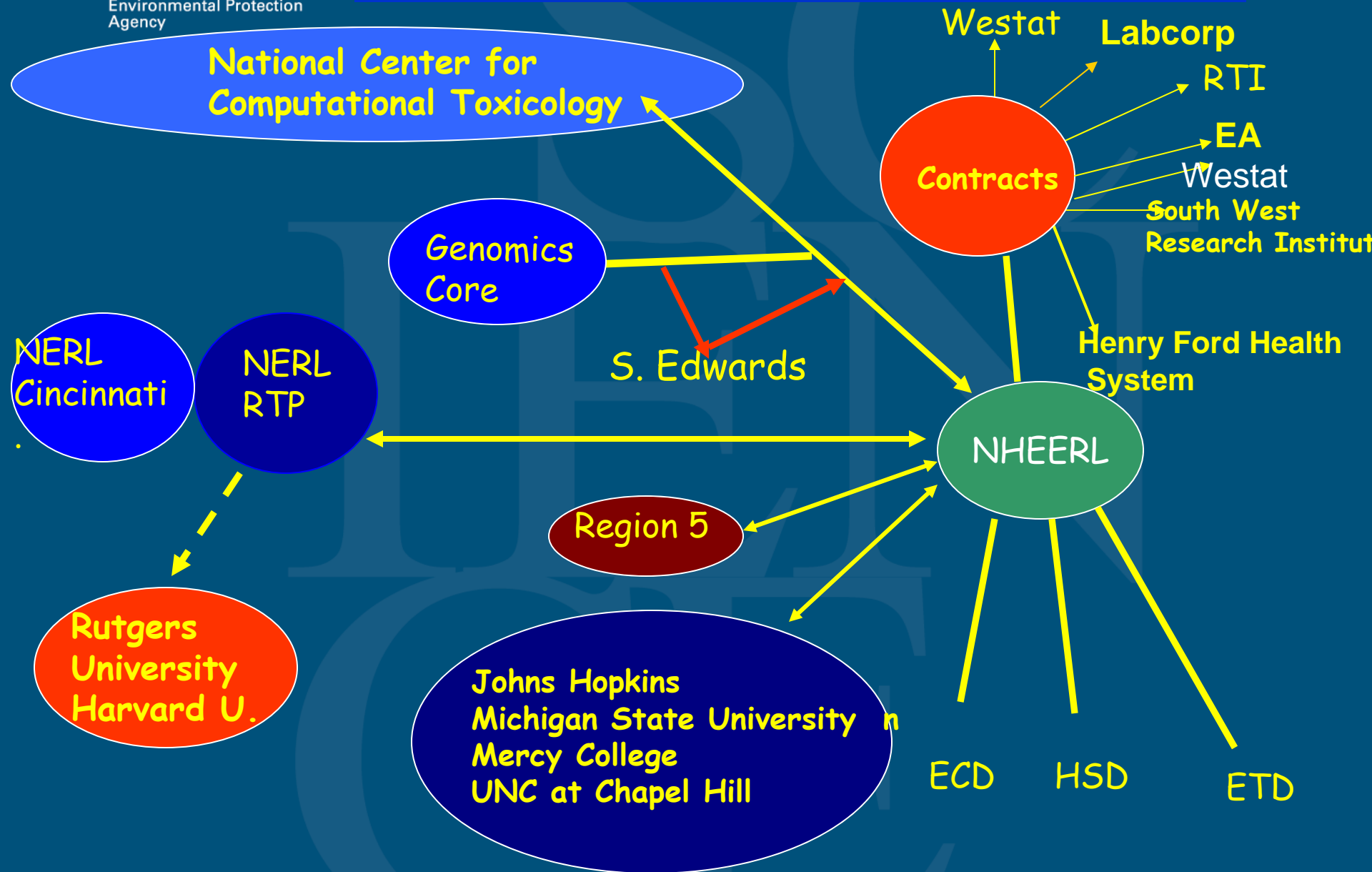
MICA



Change in research paradigm

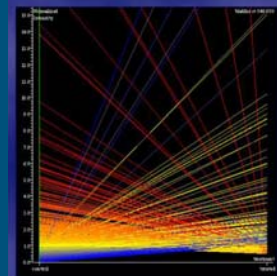
- **Work across disciplines**
- **Give up data for the greater good**
- **communicate**

MICA- Change Research Paradigm



MICA

- **A childhood asthma study and Parallel rodent study**
- **A NHEERL and NCCT Computational Toxicology study**
- **Combines and Integrates biomarkers of exposure effects and susceptibility in the context of clinical measurements and disease (asthma) outcome.**



National Center for Computational Toxicology

Goals

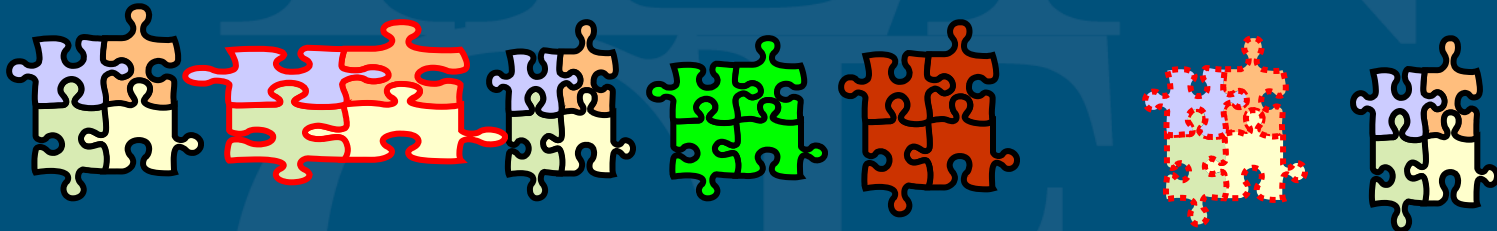
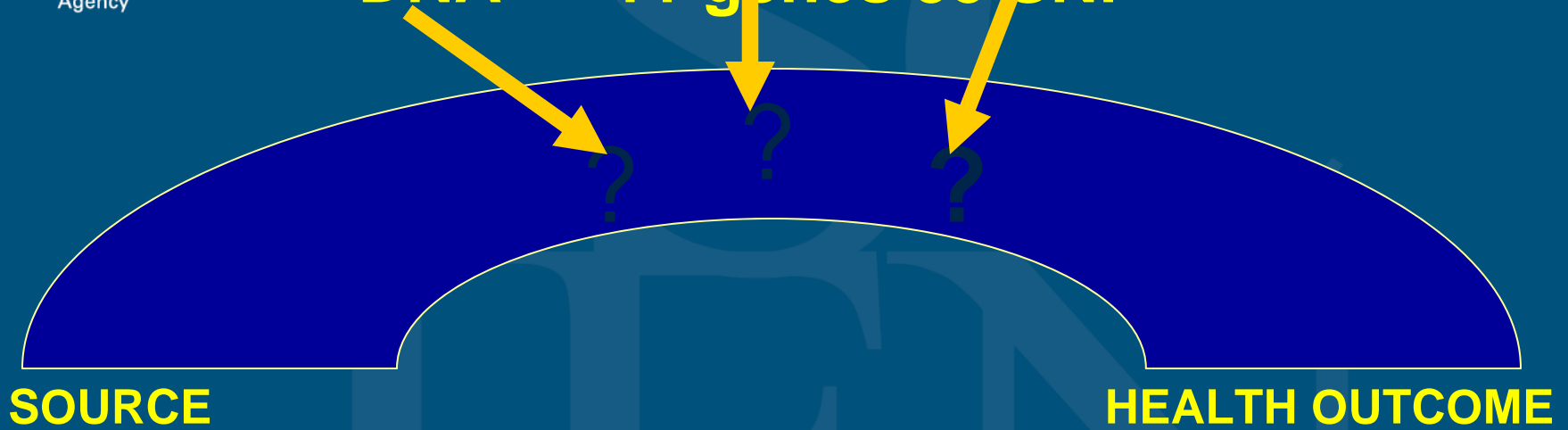
Improve linkages in the source to outcome paradigm

- Provide predictive models for Hazard ID
- Improve Quantitative Risk assessment Dose, species, chemical class

7 New Starts ---MICA

RNA--- Blood Gene expression

DNA ----11 genes 55 SNP



**INTERNAL
DOSE**

**INTEGRATED
DOSE**

**EARLY
RESPONSES**

MICA

Questionnaire
Diet
Time activity



Fingernails



urine



blood



Odor test



Lung function

NO ex



Asthma



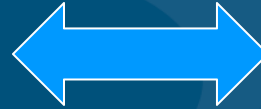
Vacuum dust and
Passive monitoring

BMI
Blood Pressure
O2 Saturation
Cell differentials
Blood Chemistry
Cytokines
Creatinine
cotinine
Medications

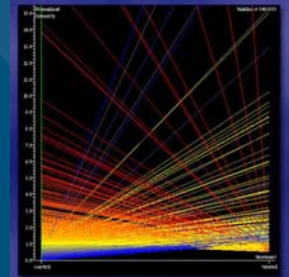
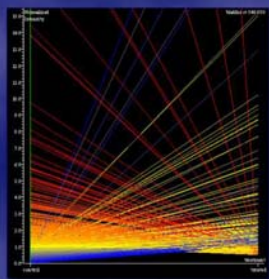
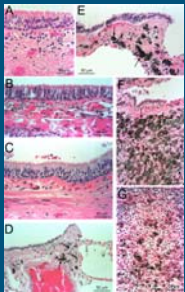
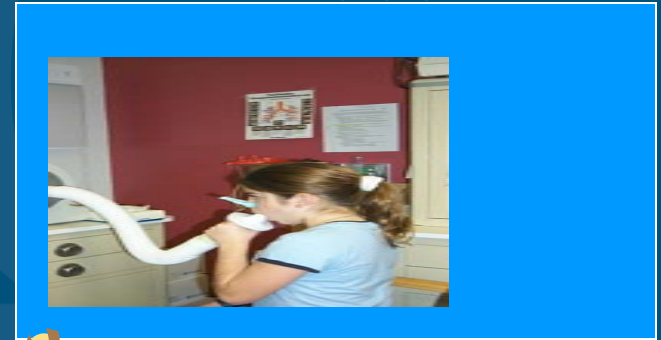
Objective

Increase our understanding of asthma by assessing the complex gene/environmental relationships through the combined use of innovative methods to manage and analyze multifactorial data

MICA RODENT Phase 1



Childhood asthma Phase 2



11 polymorphic
genes
55 SNP

MICA Nested Within Detroit Children's Health Study

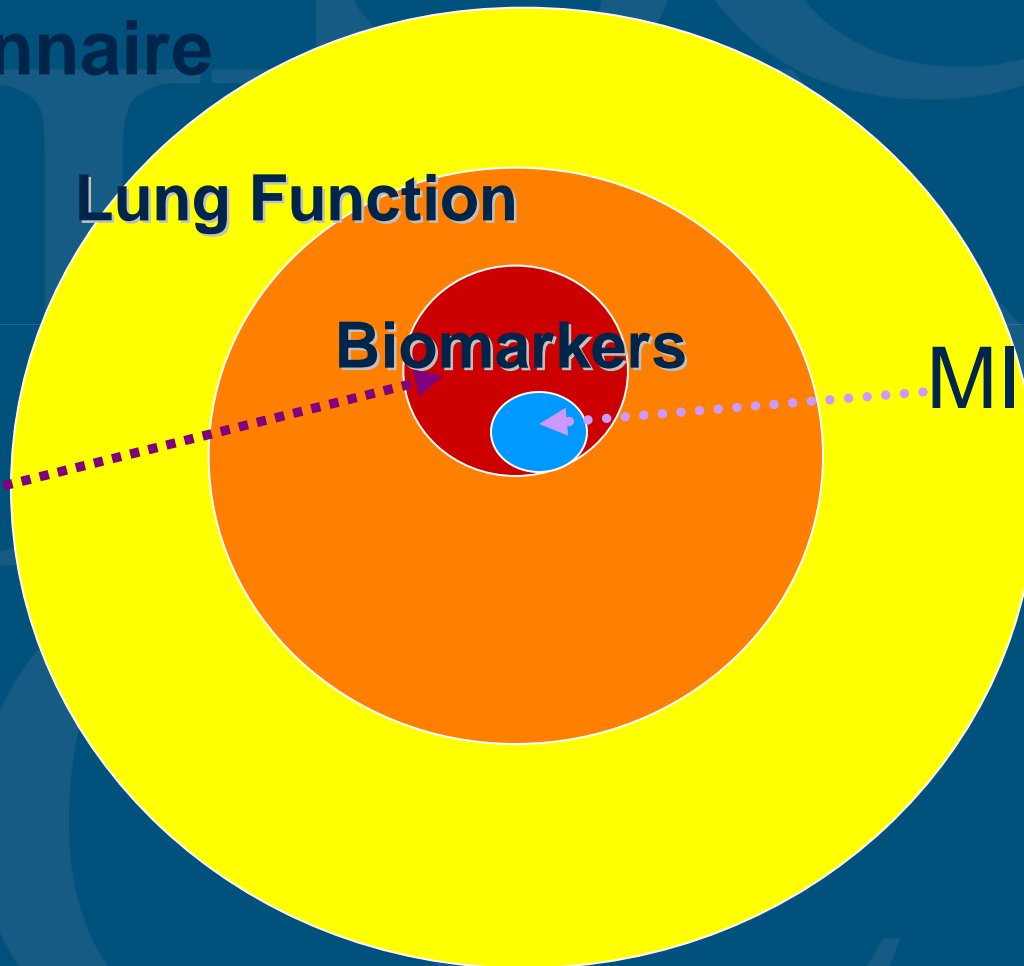
Questionnaire

Lung Function

Biomarkers

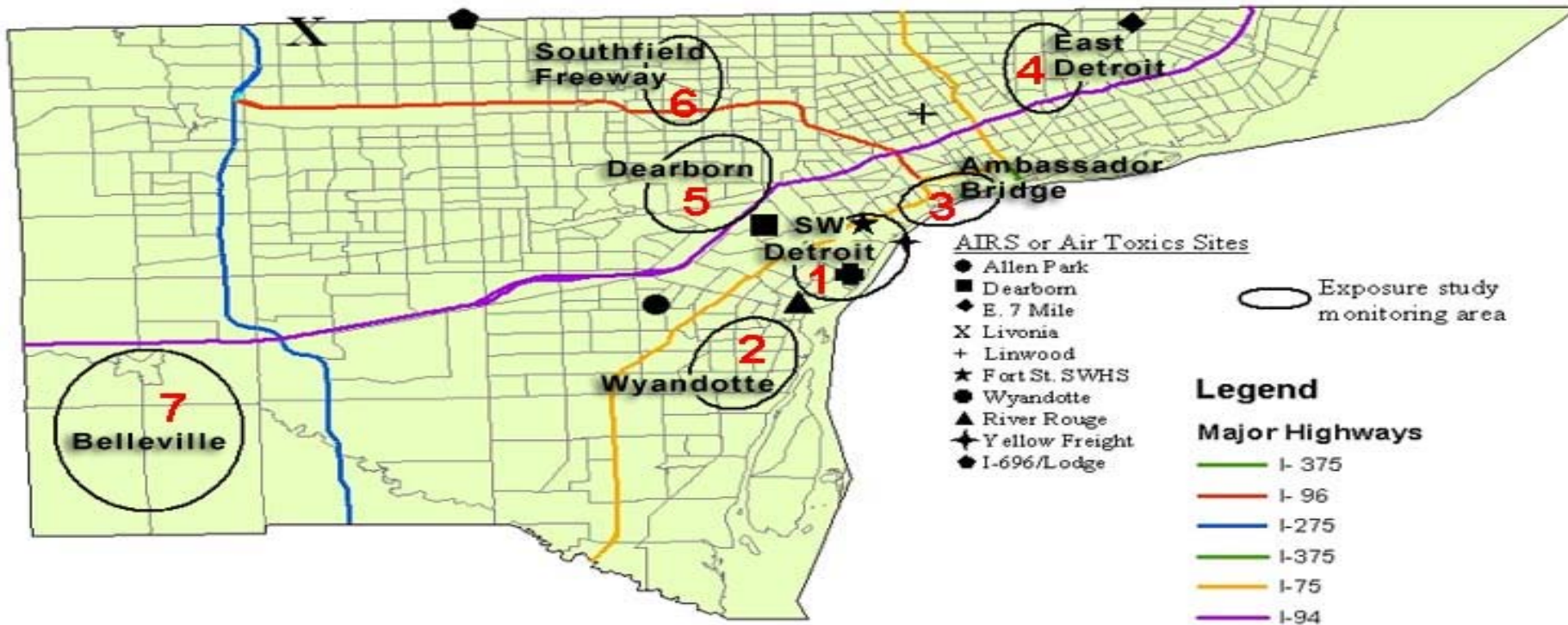
MICA AIR

MICA



National Exposure and Research Laboratory Detroit Exposure Aerosol Research Study "DEARS"

Map of Wayne County (Detroit area), showing Interstate Freeways and AIRS Monitoring Sites. Study participants will be selected for monitoring from the seven circled areas.

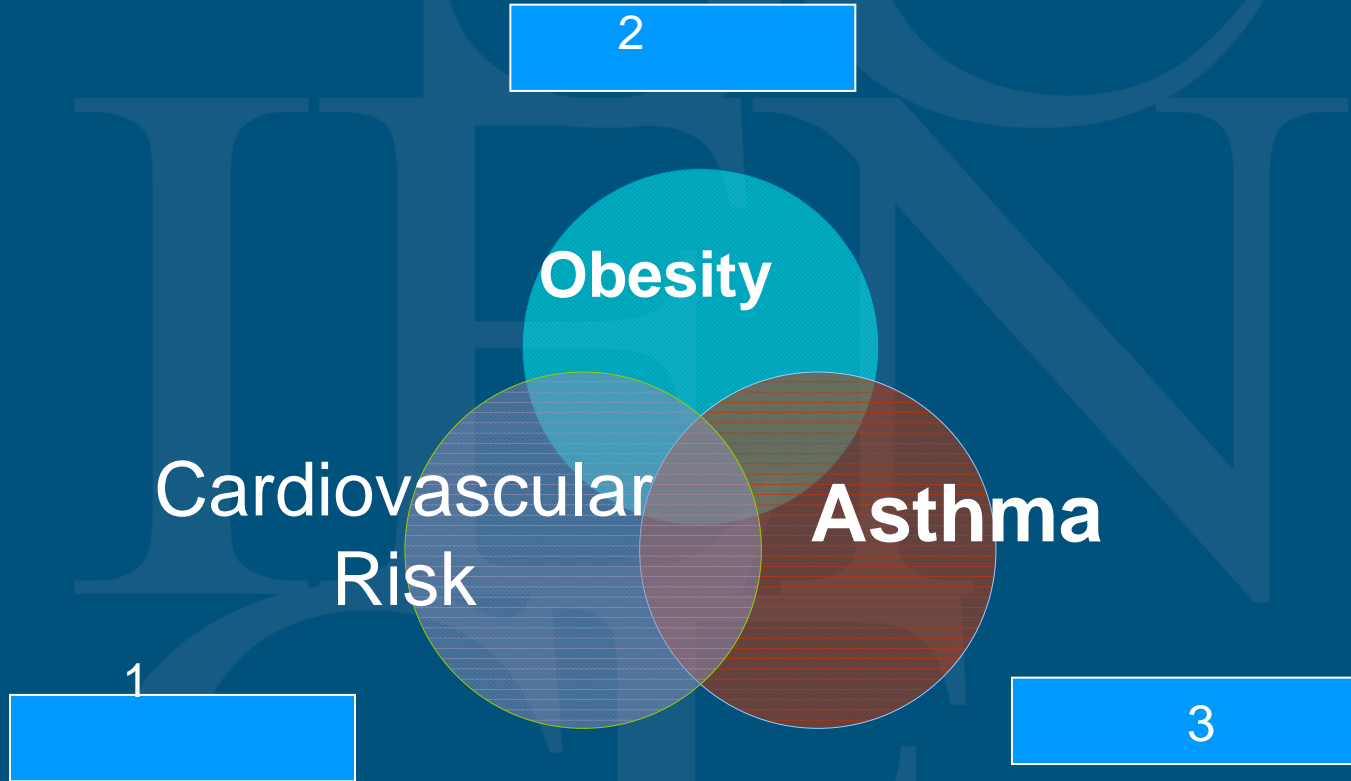


MICA

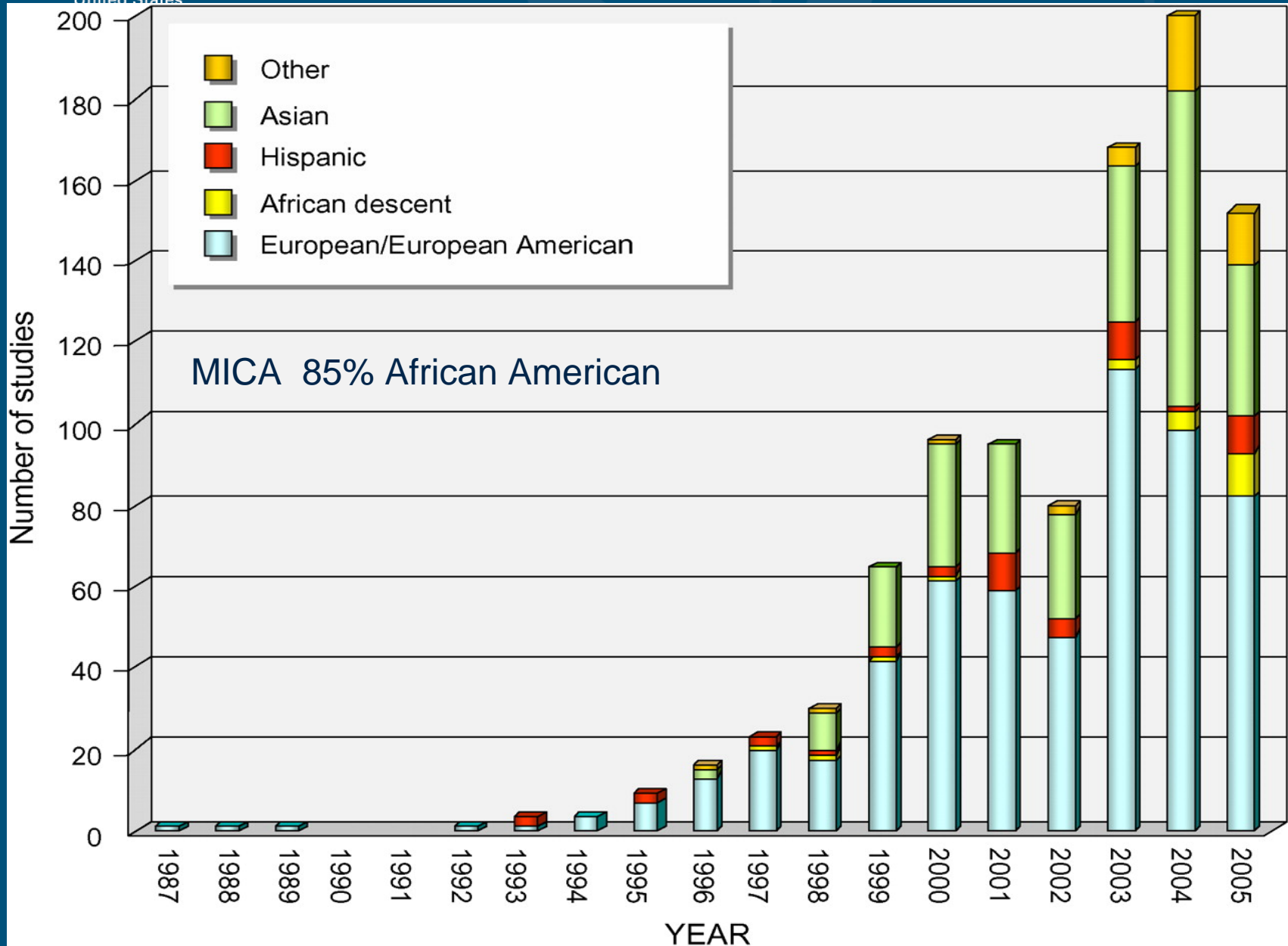
- 200 children (asthma and no asthma) 9-12 years of age
- 100 families participated in self monitoring (indoor outdoor) as part of MICA air
- Vacuum Dust and medication list brought to clinic
- Educational and “station walk through” presentation to provide context to the study
- Consent assent and Questionnaire
- Lung function, NO ex and odor testing
- Blood, urine, fingernails collected
- 90 percent of subjects provided samples at each station.

MICA Childhood Study

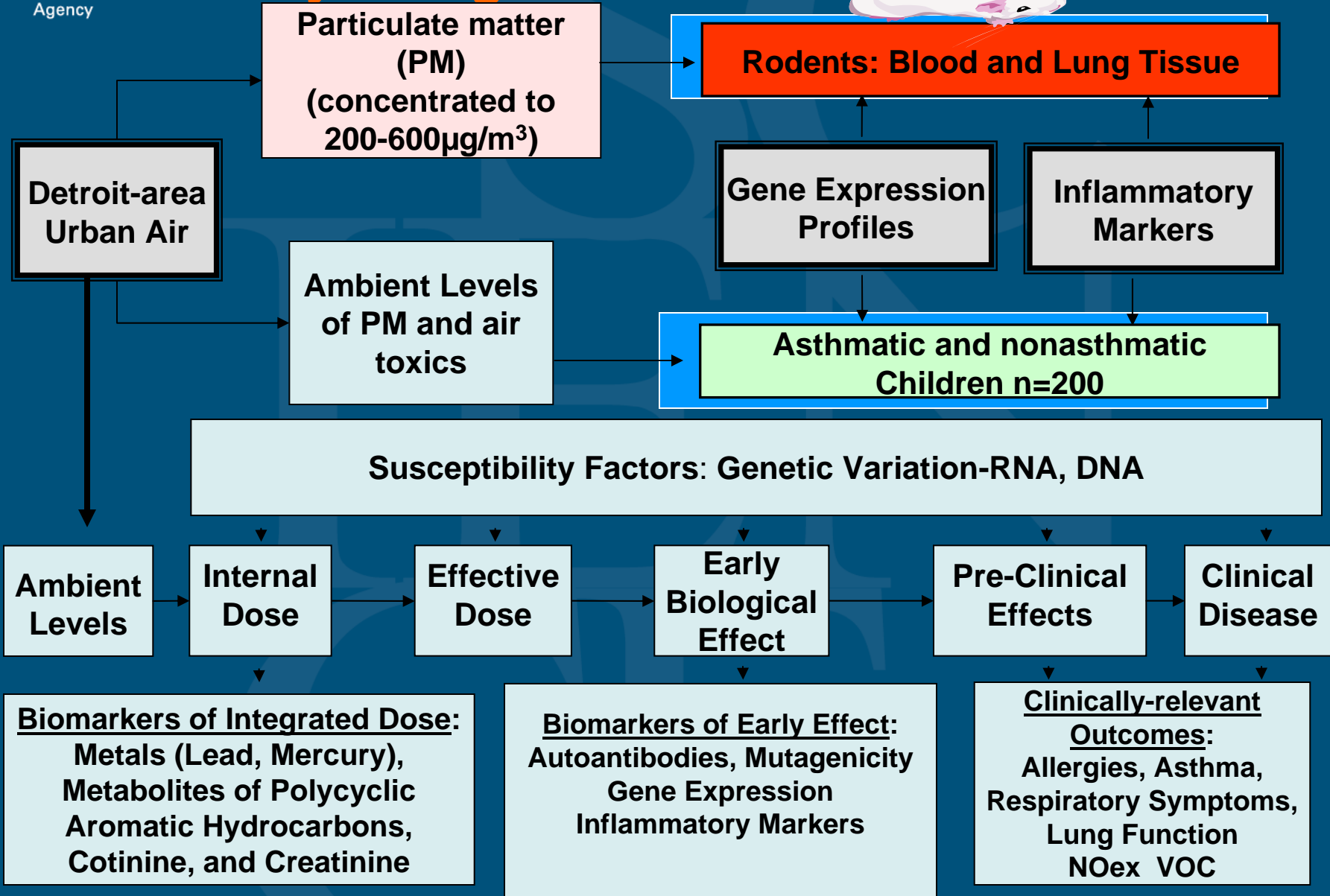
Multiple Risk Factors



Asthma studies by race



Study Design



- **Rodent Study**
- **Detroit AIR**
- **Concentrated
Air Particles
Exposures**



MSU Mobile Air Research Laboratory (*AirCARE 1*)



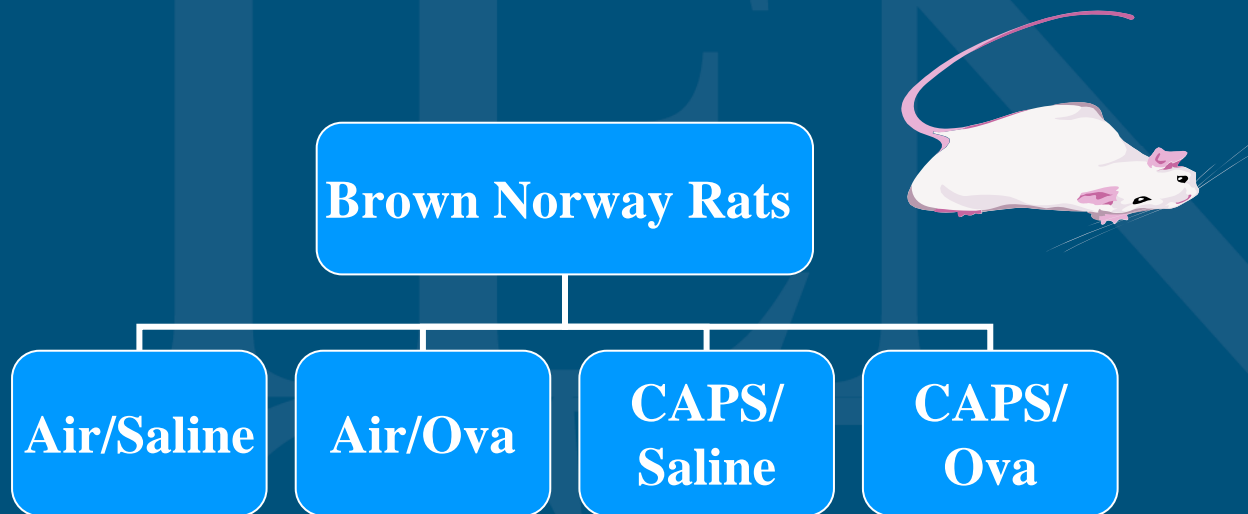


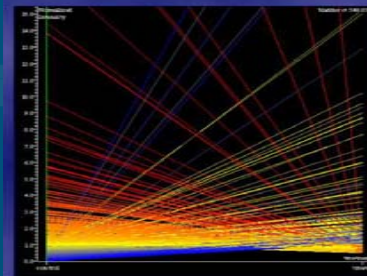
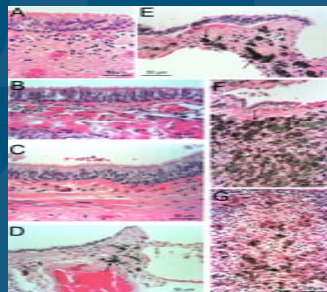
Air Particle
Concentrator
and Inhalation
Exposure System
for Laboratory
Rodents

Animals are
exposed to
concentrated fine
and ultrafine
particles in
specially designed
shoebox cages

MICA (I)

Evaluate Utility of Rodent models for analyzing gene expression data in childhood asthma study



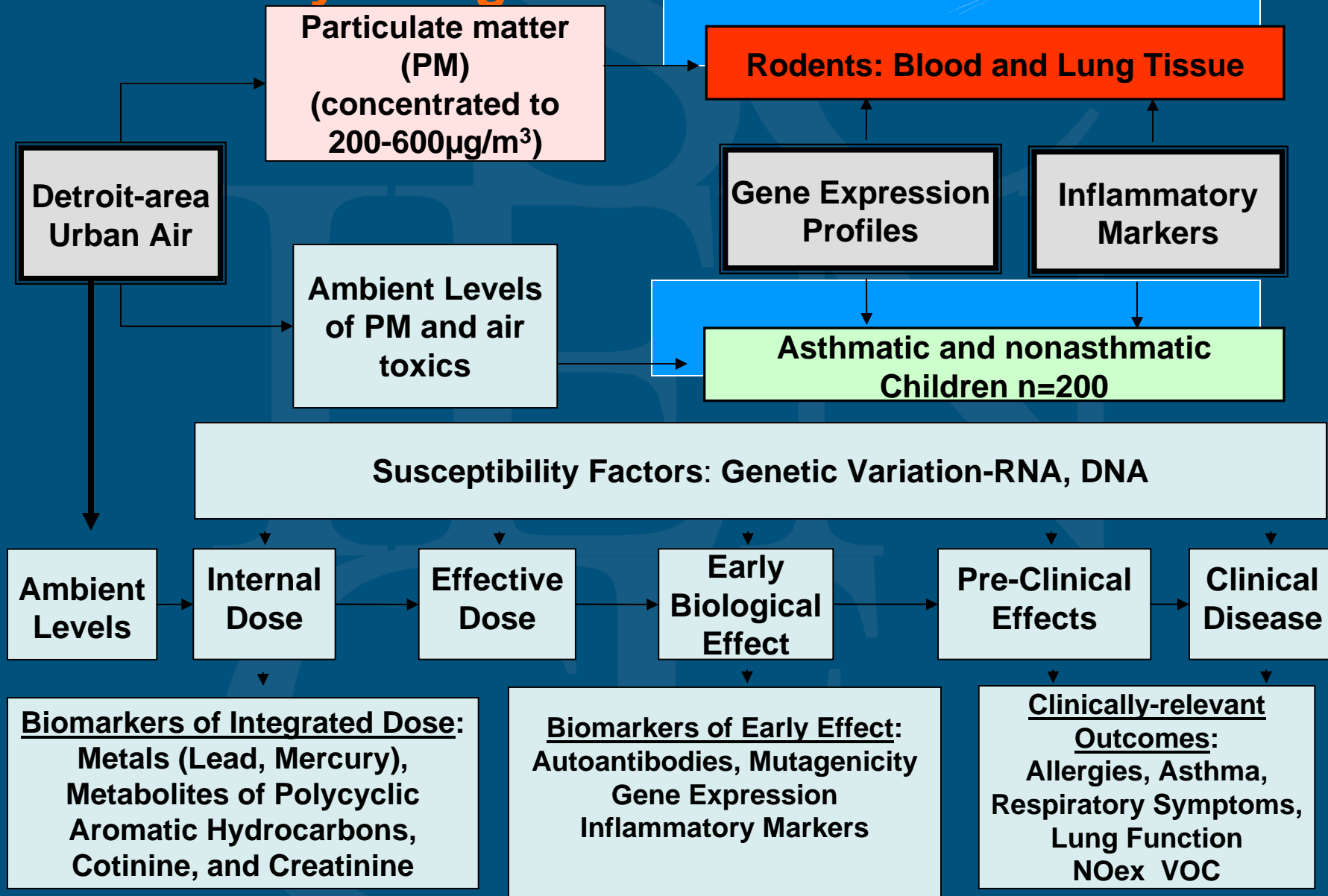
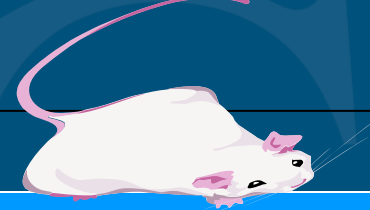




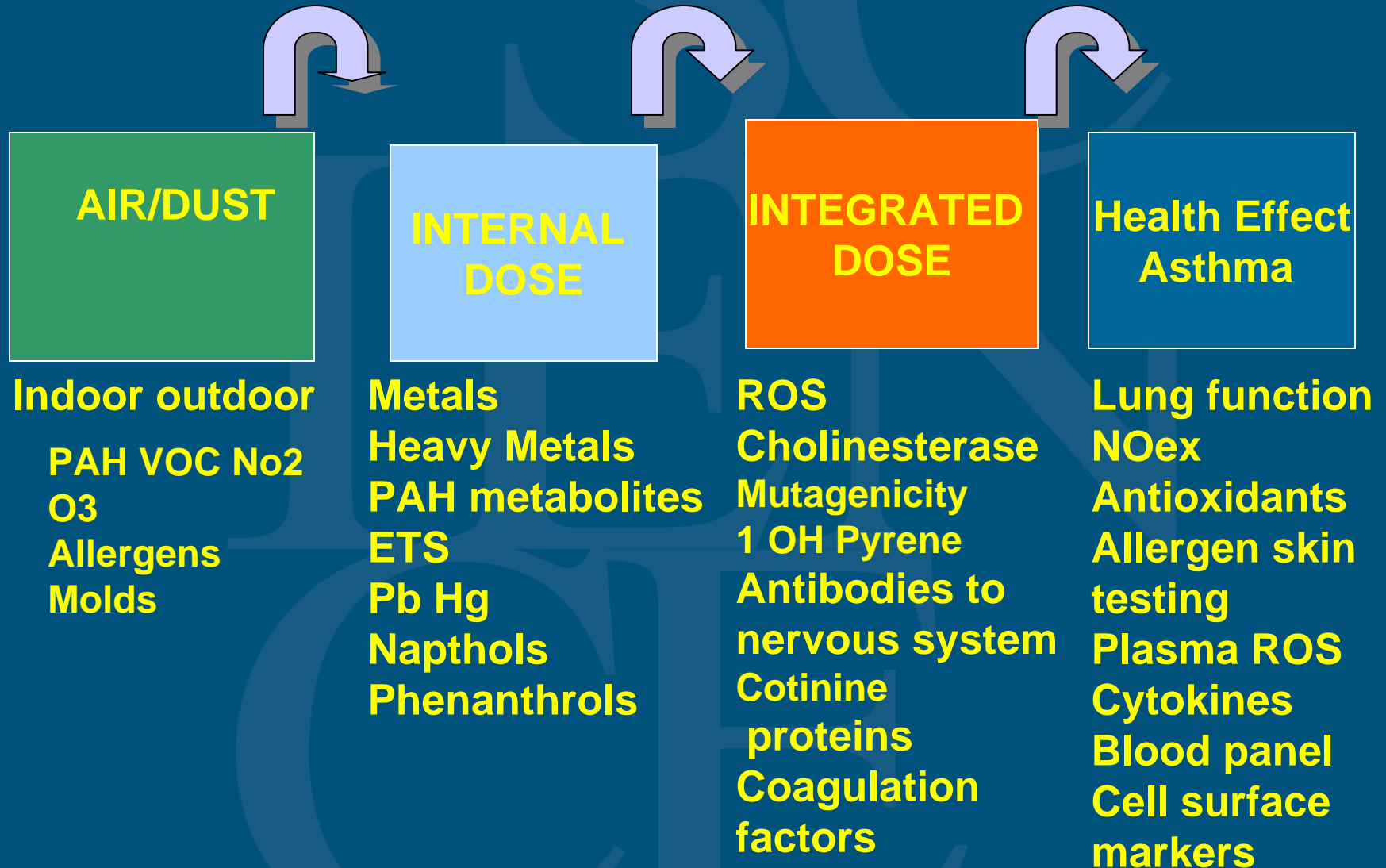
United States
Environmental Protection
Agency

MICA

Study Design



Gene expression - RNA Genotyping - DNA (11 genes 55 SNP)





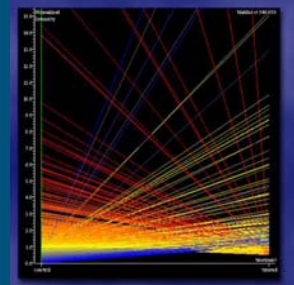
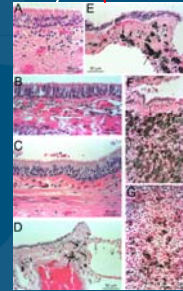
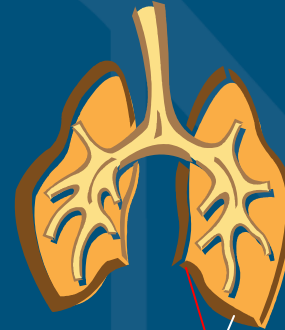
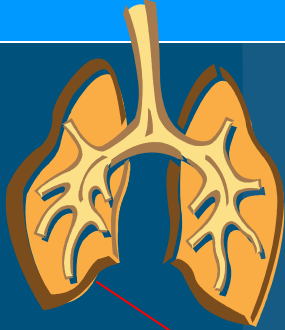
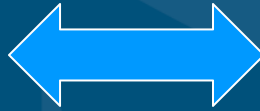
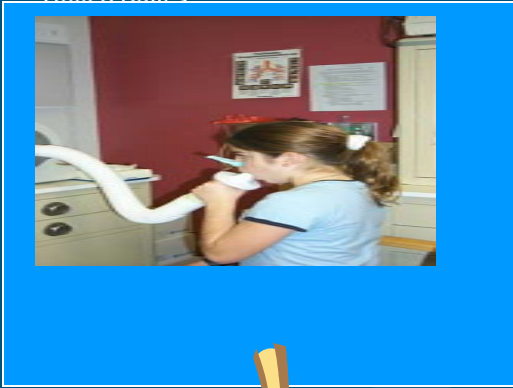
Passive monitoring: NO₂, PAHs VOC, (indoor and outdoors)

Vacuum dust

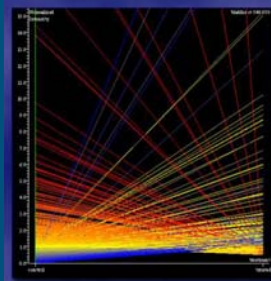
Metals, PAHs, aero-allergens mold, endotoxin)

Biomarkers--- Clinical and Environmental

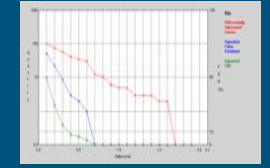
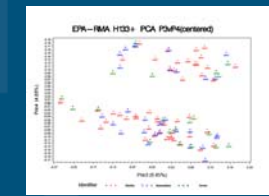
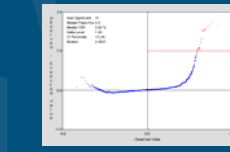
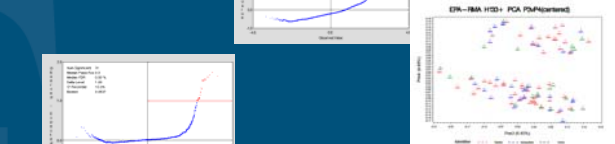
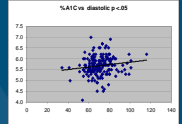
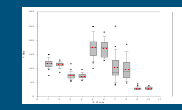
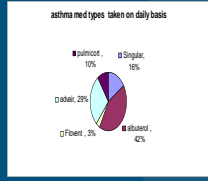
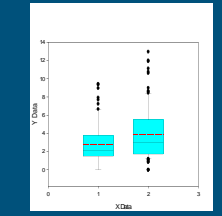
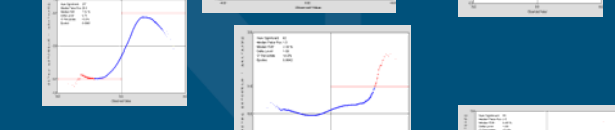
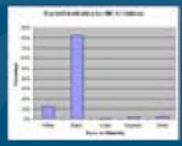
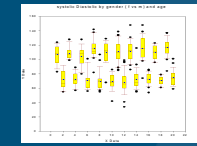
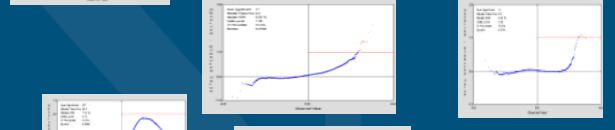
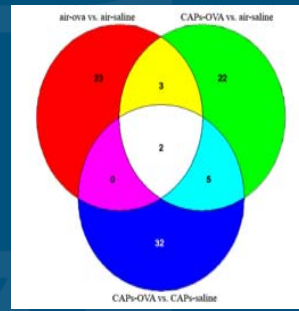
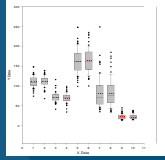
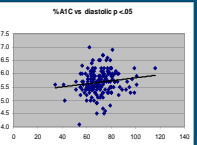
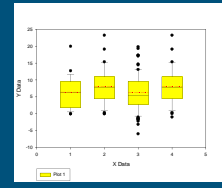
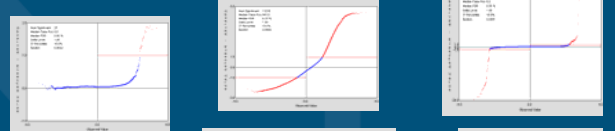
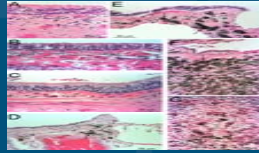
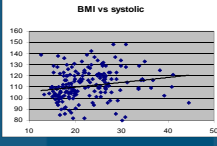
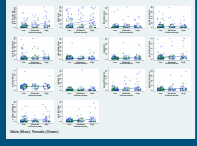
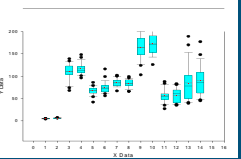
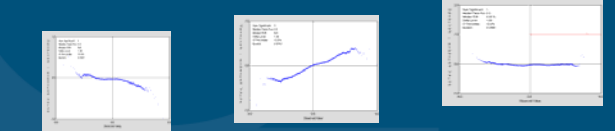
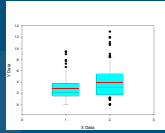
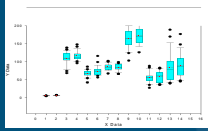
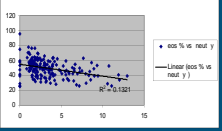
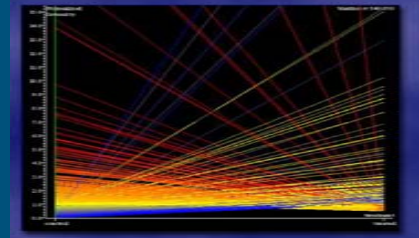
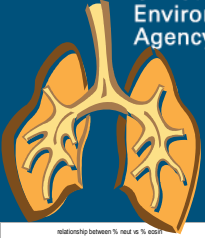
Urine	Cotinine Creatinine
Urine	Metals: Mercury, Cadmium, Arsenic, Chromium, Manganese, Nickel
Urine	1- hydroxypyrene (1-OH pyrene) Naphthols, Phenanthrols, Hydrocarbon Metabolites phthalates
Urine	Mutagenicity Assays
Serum	Autoantibodies for nervous system proteins, Blood Chemistry, Total IgE and specific antibodies to common aero-allergens (multiscreen inhalant and food antibody series)- *dust mite, Cockroach , Mouse, Rat Urine Protein,
Plasma	Reactive Oxygen Species, cytokines (IL4, 6 IL13), tumor necrosis factor-alpha, c-reactive protein, fibrinogen
Whole Blood	hematology panel, lead and mercury, Gene expression (RNA), glycosolated hemoglobin
Nails	Mercury, Cadmium, Arsenic, Chromium, Manganese, Nickel
Serum	IgE-inducing proteins associated with fungal exposures



11 polymorphic
genes
55 SNP



NOW WHAT ???



CINCINNATI CHILDRENS HOSPITAL

J ALLERGY CLIN IMMUNOL
VOLUME 115, NUMBER 2

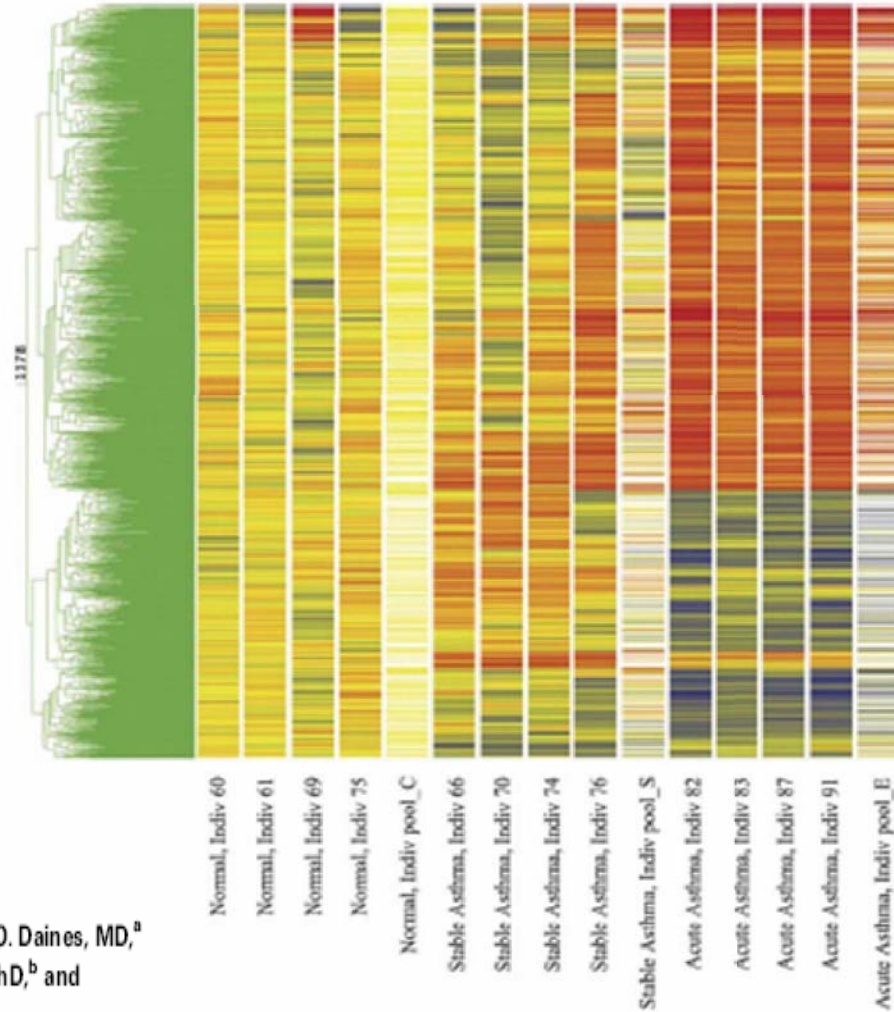
NORMAL

Stable
Asthmatics

UnStable
Asthmatics

Guajardo et al 247

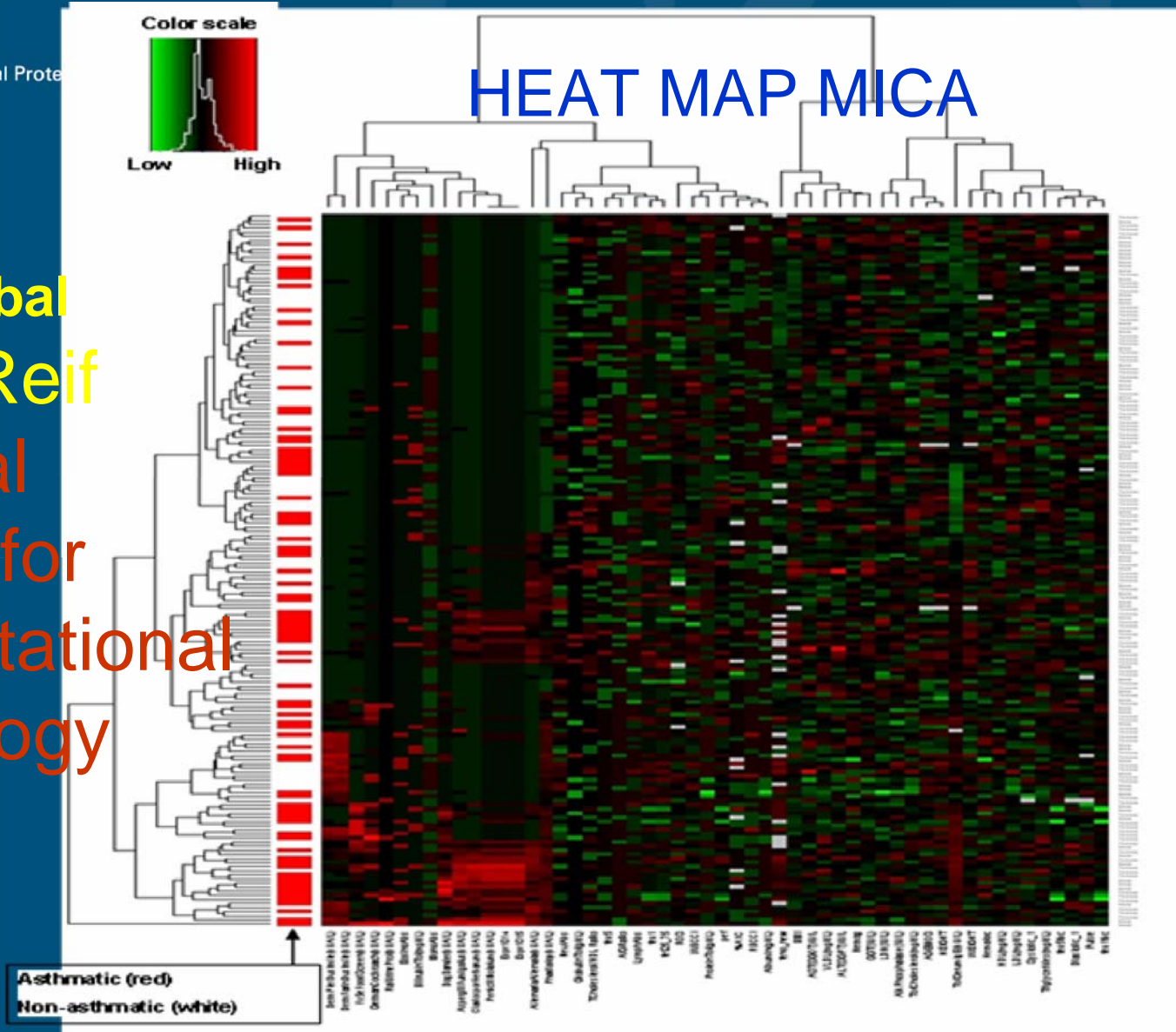
NASAL EPITHELIAL CELLS



Jesus R. Guajardo, MD, MHPE,^a Kathleen W. Schleifer, PhD,^b Michael O. Daines, MD,^a
Richard M. Ruddy, MD,^c Bruce J. Aronow, PhD,^d Marsha Wills-Karp, PhD,^b and
Gurjit K. Khurana Hershey, MD, PhD^a Cincinnati, Ohio

HEAT MAP MICA

Elaine Hubal
David Reif
National
Center for
Computational
Toxicology



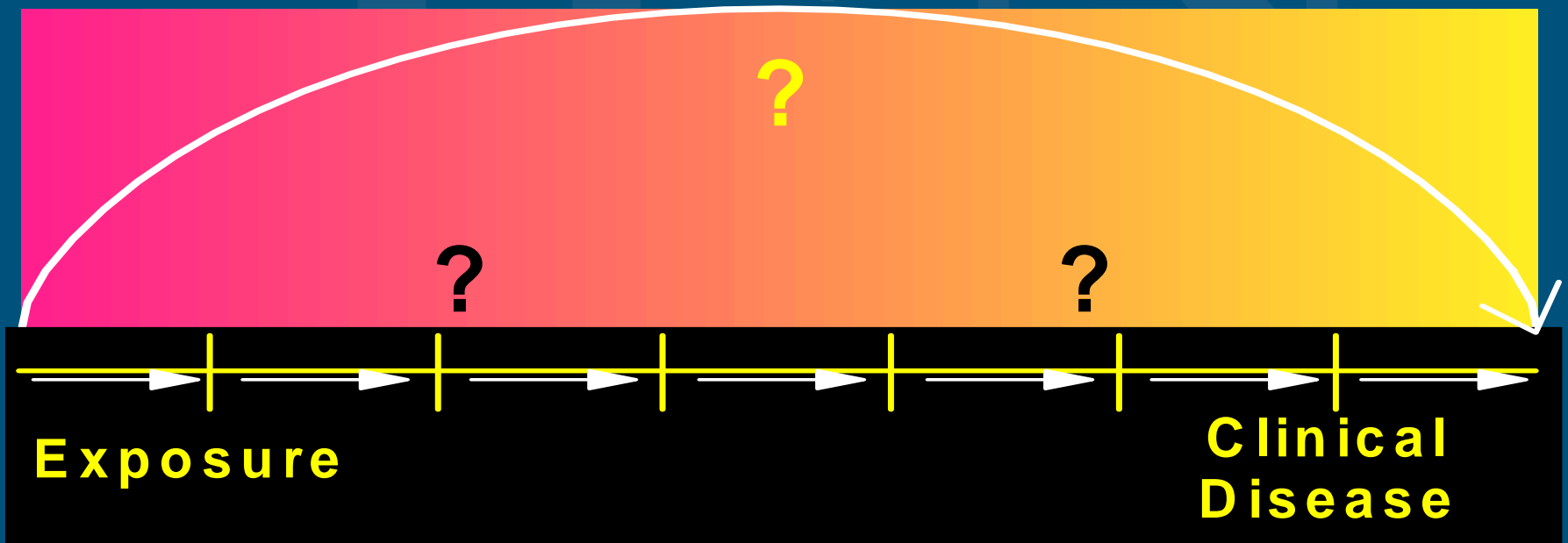
Biomarker Needs

- Exposure biomarkers in the context of clinical health indicators
- Mechanistic information test biological plausibility in rodents
- Validation of surrogate cells with target tissue responses

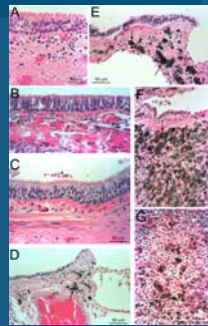
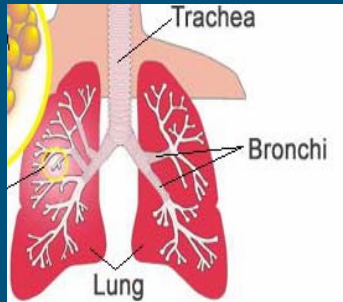
Archiving of biological and environmental samples and measurements as new technologies advance

Summary

- High-data content technologies, elucidating the genetic and environmental basis for toxicity and disease



Integration of Diverse Set of exposure, effects and susceptibility

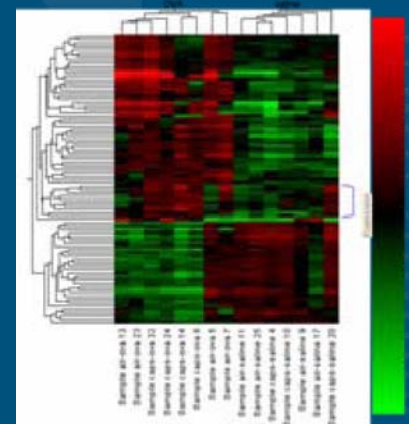
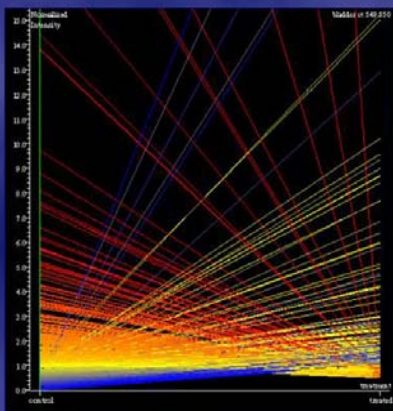


bioinformatic

Gene expression arrays

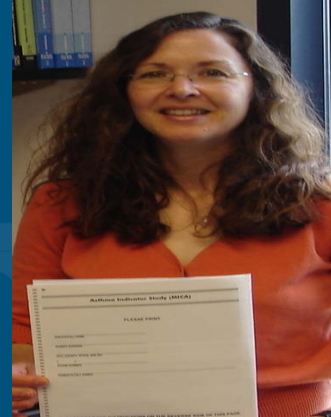
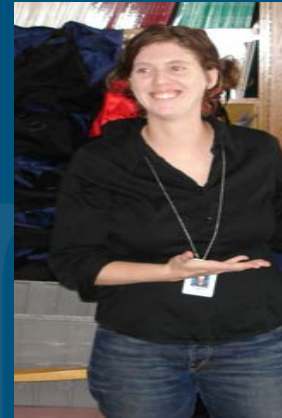
statistical
analysis.

computational



genes, pathways, and networks

Acknowledgements



PHASE II MICA

**National Health Exposure
and Effects Laboratory**

**Stephen Edwards
Elaine Hubal
David Reif**

**National Center for
Computational Toxicology**

Acknowledgements



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Henry Ford Health System Clinic

Henry Ford Health System-lab





Acknowledgements

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Ann Williams

NCCT

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ETD

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Robert Hamilton
John Wiseman
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Frank Weber
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Kurt Patrizi

Henry Ford Health System

Clinical and Lab