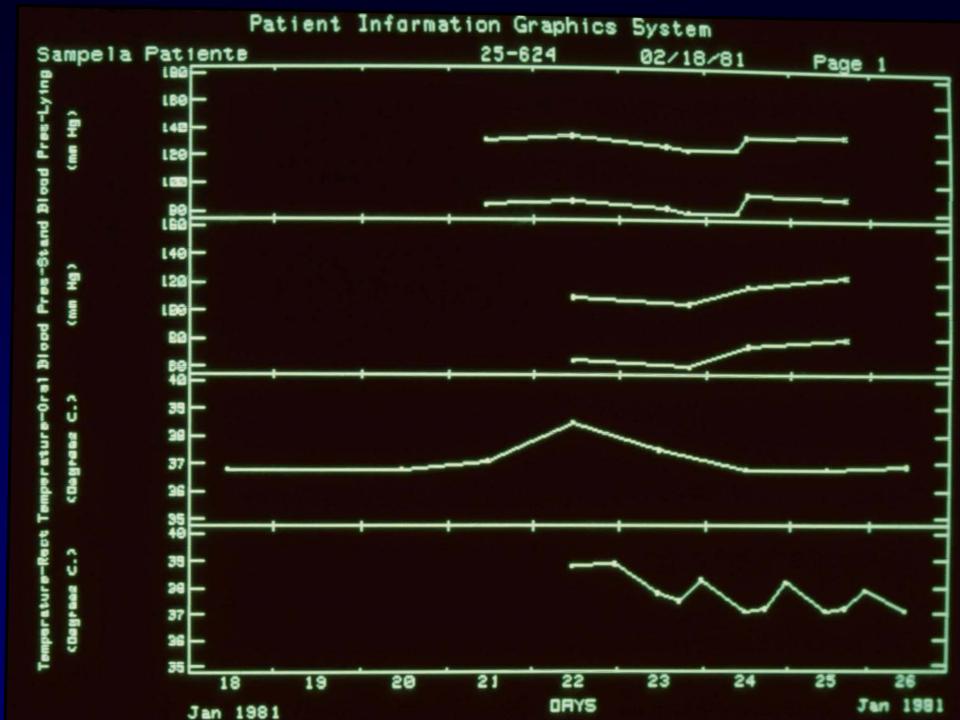
# The NIH Biomedical Translational Research Information System (BTRIS)

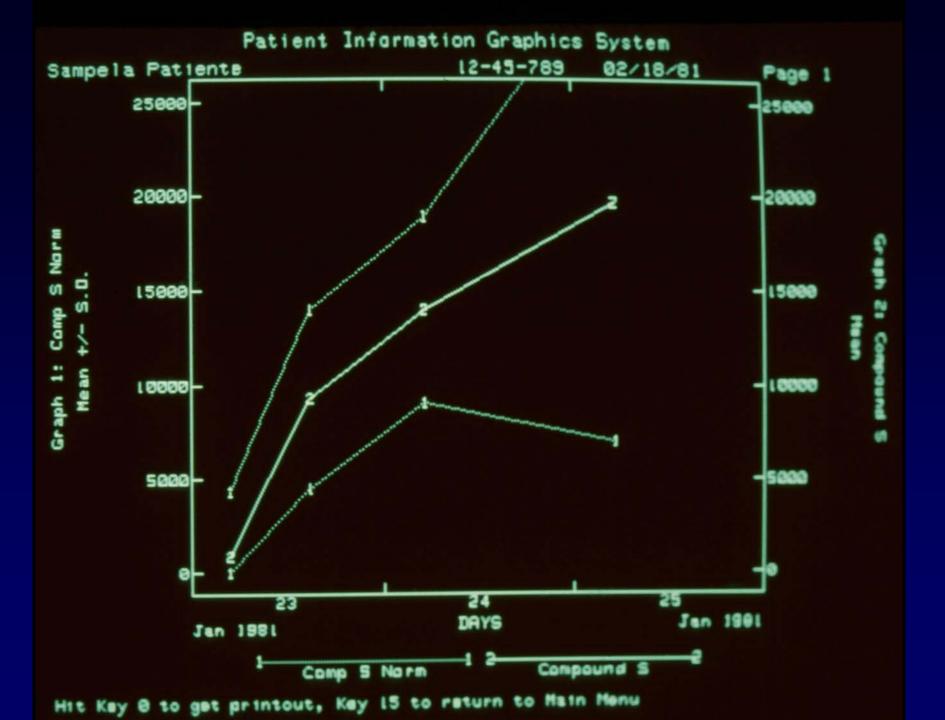
Town Hall Meeting - Information Session February 26, 2008
Lipsett Auditorium

#### The Reuse of Biomedical Data

- Secondary uses of clinical data for:
  - Patient care
  - Research
  - Administrative processes
- Use of patient data for research
- Use of research data for patient care ("translational research")
- Data may require transformation:
  - De-identification and Re-identification
  - Indexing
  - Aggregation by time
  - Abstraction by classification
  - Conversion to relevant concepts







#### **Brief Bio**

- Internal medicine residency (St. Vincent's, NY)
- Medical informatics fellowship (Harvard/MGH)
- 20 years at Columbia
  - Informatics research
  - Building clinical systems
  - Teaching informatics and medicine
  - Clinical practice
- Clinical data repository and warehouse
  - 25+ data sources
  - 2,000,000 patients
  - 20 years of data
  - Innovations in coding and organization
  - Concept-based queries
  - Natural language processing (NLP)

## How often are patient with the diagnosis of myocardial infarction started on beta blockers?

#### Clinical Data Warehouse at CPMC

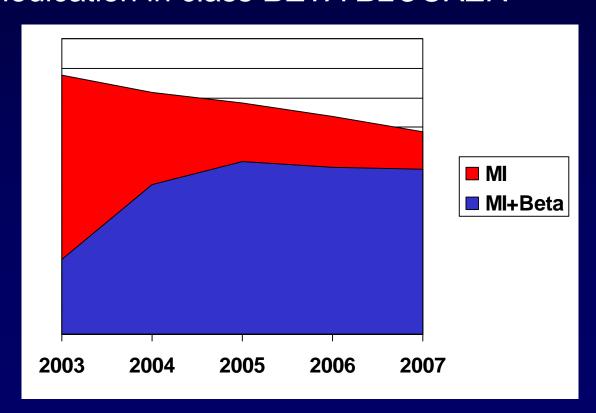
#### Mediated Query Please see Important Riders, Approval Requirement and a Fee Policy at the bottom of this page. James J. Cimino NAME Biomedical Informatics DEPARTMENT Professor TITLE&RANK Principal Investigator ROLE IN PROJECT Vanderbilt Clinic 558 OFFICE ADDRESS 212-305-8127 TELEPHONE ijc7@columbia.edu EMAIL Testing the use of MED Codes in the Warehouse PROTOCOL (PROJECT) NAME AND A BRIEF DESCRIPTION (255 chars max. Use comments if you need more space) Investigator Initiate PROJECT SPONSORSHIP/REASON James J. Cimino PRINCIPLE INVESTIGATOR 3/31/2005 DATE of REQUEST e.g. 12/31/2005 04/05/2005 DATE REPORT IS REQUIRED 1) Number of patients with discharge diagnosis of Myocardial Infarction (MED Code 2618 or descendants, ICD9-CM Code DATA REQUESTED 410.00 to 410.99), by year from 2000-2004. Summary/Non-identify info only (No restriction) APPROVAL LEVEL CYES ONO **HUMAN SUBJECT INVOVLED?** IF YES, IRB #

# How often are patient with the diagnosis of myocardial infarction started on beta blockers?

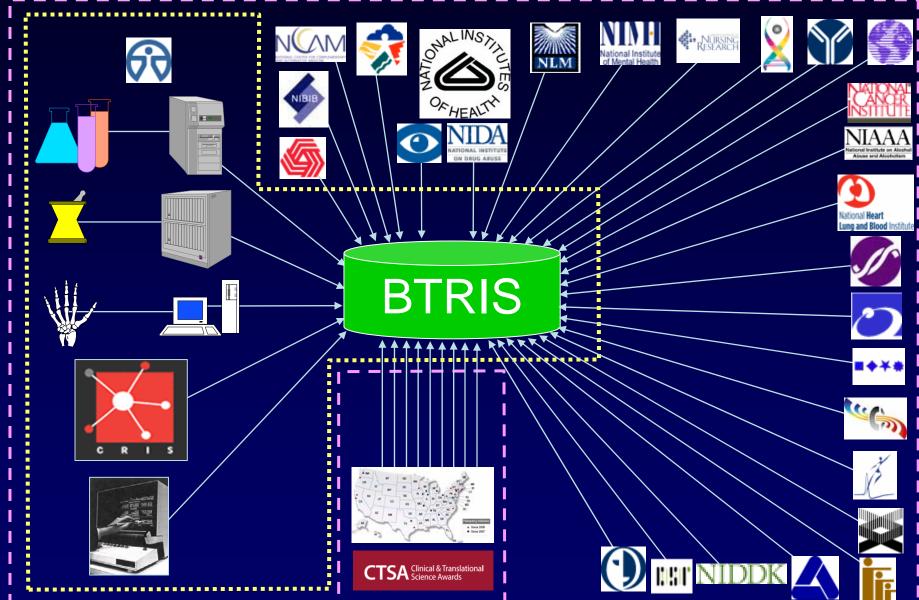
Find all patients with Diagnosis in class MYOCARDIAL INFARCTION

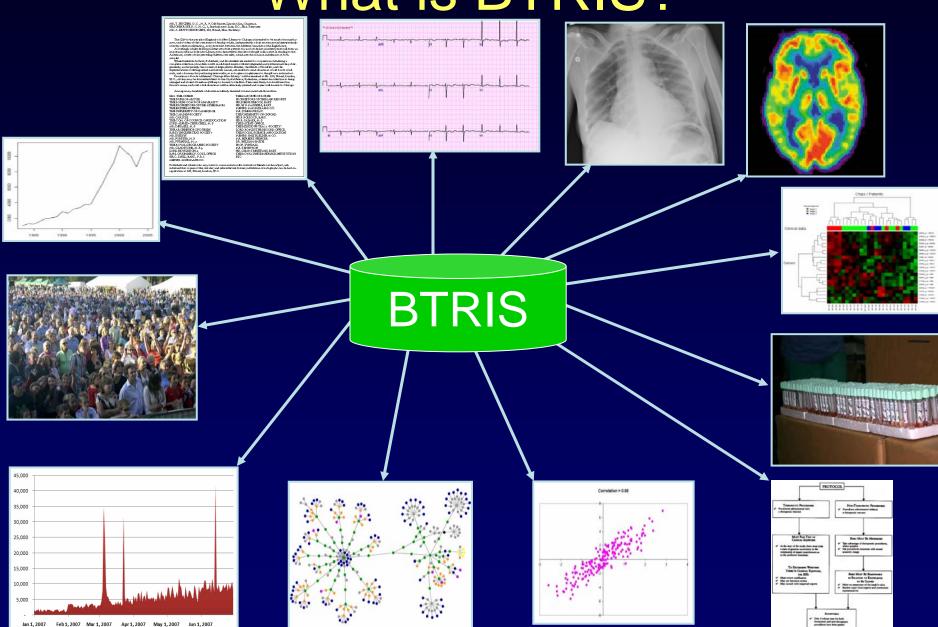
Find all patients with Diagnosis in class *MYOCARDIAL INFARCTION* AND with

Medication in class BETA BLOCKER



- Formerly "CRIS-II"
- Not "Son of CRIS"
- Not just clinical
- Includes focus on translational research
- Hence: Biomedical Translational Research Information System





Subject Recruitment

Data Analysis Tools

**Hypothesis Generation** 

**Hypothesis Testing** 

**Data Retrieval Functions** 

Authorization Subject-Oriented Cross-Subject Re-Identification NLP

#### **Data Repository**

Data Acquisition Processes

Coding Indexing

**De-Identifying** 

Permission Setting

n t o I o g y

O

#### Queries from Requirements-Gathering

- Provide Medication lists at time of patient encounters. Include drug diaries for inpatient, outpatient and in-between encounters in the patient medical record. Include all chemo and non-chemo drugs from CRIS and IC systems
- Provide Medication administration documentation (drug diaries) with times as part of patient record
- Provide ability to compare patient results, Medication Administration Records between dates and/or encounters
- Provide drug randomization info, compliance records and drug accountability info for all investigational, study and prescription drugs
- Provide all Clinical Center lab results with times of specimen draws
- Provide external lab results
- Provide archival images
- Provide demographics data including age, BMI, race, gender, contact info, etc
- Provide access to genomics and bio-markers data
- Provide cumulative blood volumes, research drugs and radiation for subject over a given period
- Provide ability for ICs to feed expanded diagnosis/problem lists
- Provide searching and filtering patients' data by all diagnosis, tests, procedures, protocol & protocol classifications

- Standardize medication and lab test codes.
- Provide integration of adverse events data in the data warehouse,
- Provide integration of protocol census, status and subject accrual tracking data from Protrak in the data warehouse
- Provide ability to attribute different events to protocols, viz., consent signed, protocol activated, orders, observations, adverse events, etc
- Provide integration of staff, patient and user index data across source systems in data warehouse.
- Provide original informed consent, and updated consents for re-contact of patients for research for all protocols. Provide searchable consents and image of consents in database. Provide answers to:
  - Can tissues be used for cancer/genetic research, other research, germ line testing
  - Can patient be re-contacted for questions?
- Provide single patient amendments
- Provide access and track biological specimen data
- Provide access to Appointment Data
- Provide "Review of Systems" info for each patient visit.
- Provide patient de-identification services
- Standardize Units Of Measure

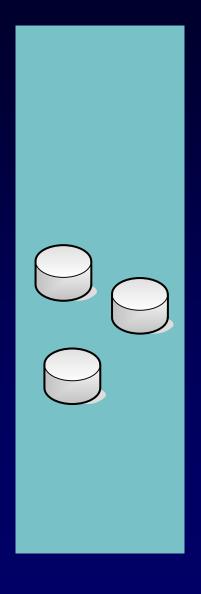
#### Access Control Issues

- Ownership of data
- Authorization for re-use
- Confidentiality and re-Identification of data

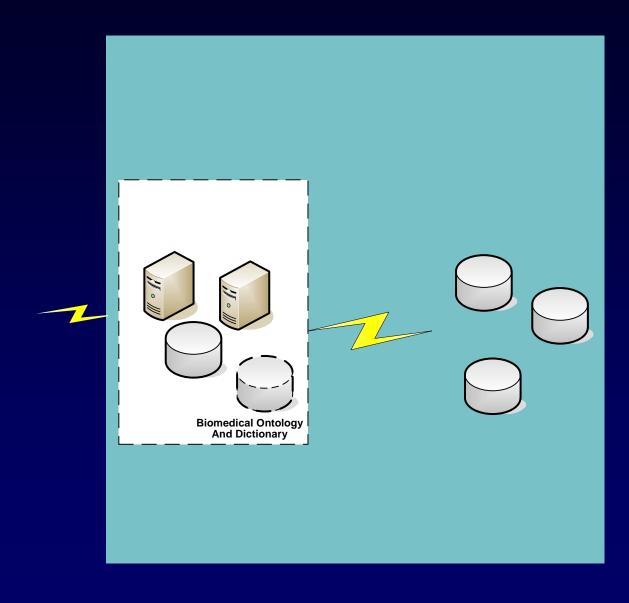
#### The PI User Group

- Volunteers from NIH community
- Most likely to benefit from BTRIS
- Help set requirements and priorities
- Commitment to participate
  - Weekly meetings
  - -Review materials, screen shots, demos
- Beta testers first access to demonstration

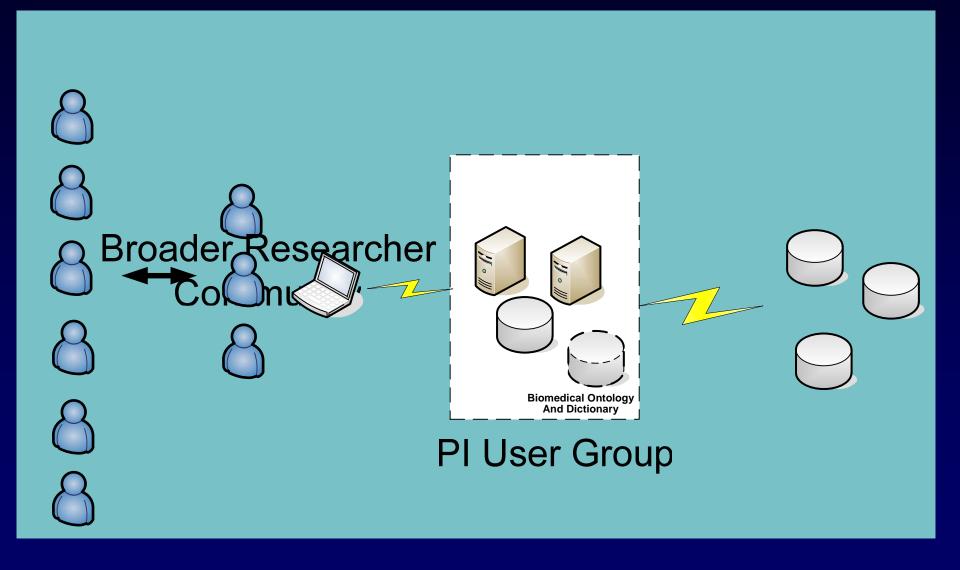
#### BTRIS Demonstration Environment



#### BTRIS Demonstration Environment



#### BTRIS Demonstration Environment



## Demonstration Project

#### Sources:

- Current laboratory data
- Old laboratory data (CDW/CDR or MIS)
- Pharmacy orders (CRIS-I or MIS)
- Terminology Services
  - Code look-up
  - Simple class-based queries
- Data services
  - Data aggregation across sources
  - Data summarization by concept class
  - Patient identification for possible recruitment
  - ...as per PI User Group

#### Timeline for Initial Rollouts

Demonstration Project: July 2008

• BTRIS: July 2009

#### BTRIS Will:

- Be the preferred system to analyze NIH clinical and non-clinical data
- Aggregate and standardize disparate and isolated data sets
- Automate and streamline processes that are traditionally manual and cumbersome
- Prioritize data sources and functionality based on needs of user community

#### Additional Information

www.btris.nih.gov

Questions about BTRIS or to join the PI User Group:

ciminoj@mail.nih.gov