

Experience with a Clinical Data Repository and Warehouse

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Outline

- History
- Clinical Data Repository
- Clinical Data Warehouse

Clinical Information Systems

- Stage 1: Early computers calculated data in context
- Stage 2: Client applications provided access to ancillary data
- Stage 3: Systems began aggregating data from multiple sources
- Stage 4: Data storage provided historical view
 - And analysis
- Stage 5: Workflow applications formalize processes between clinical roles

Clinical Information System Technology Levels

- Level 1: Departmental applications
- Level 2: Internally-developed integrated systems
- Level 3: Functional vendor-based systems
- Level 4: Comprehensive clinical information systems

Clinical Information Systems at Columbia University

- Began at Stage 3
- Pushing a Level 1 system to Level 2
- Issues
 - Vocabulary
 - Data modeling
 - Interfaces
 - Decision support
 - Data processing
- Recipient of first Nicholas Davies Award

Clinical Information Systems Architecture

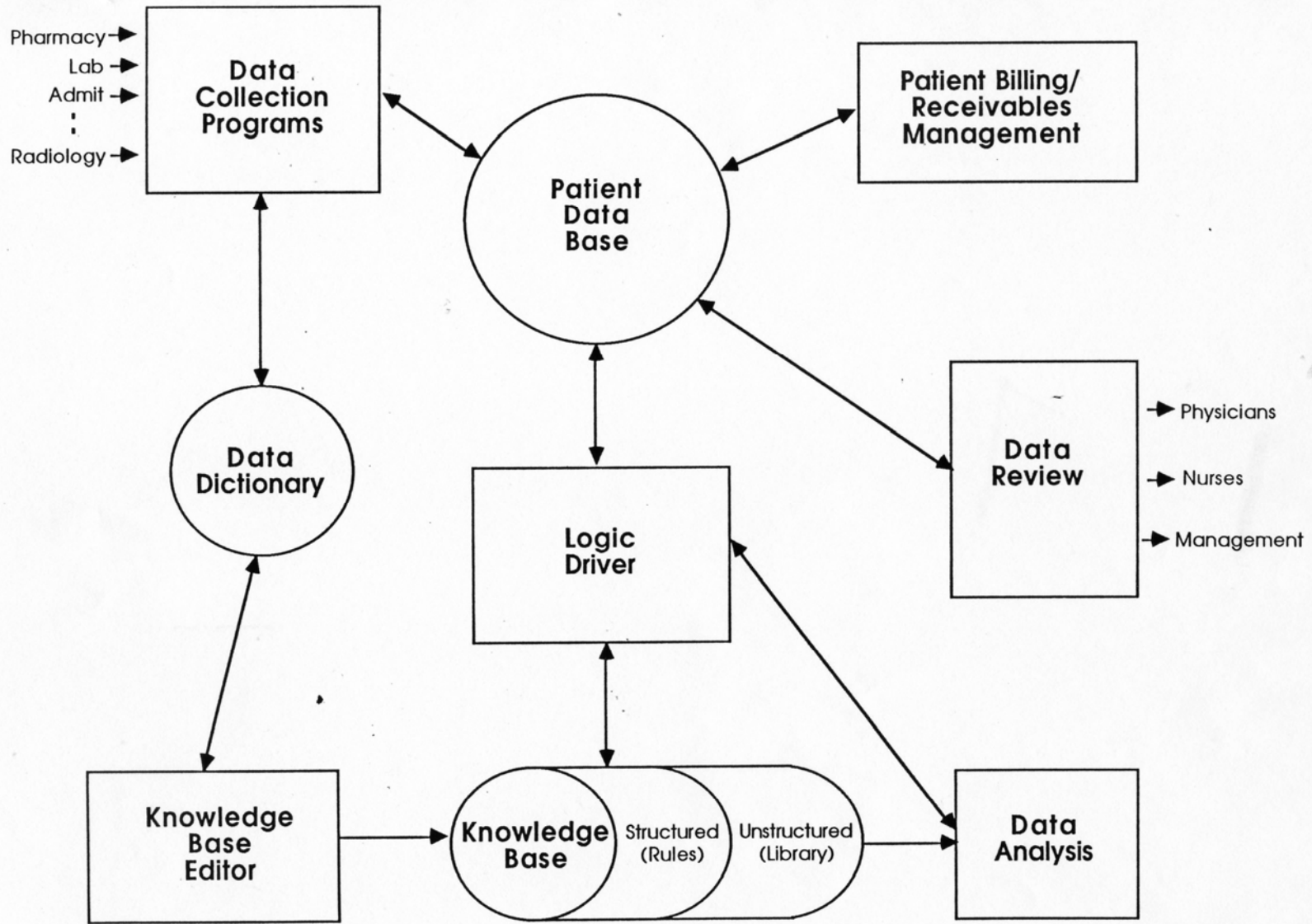
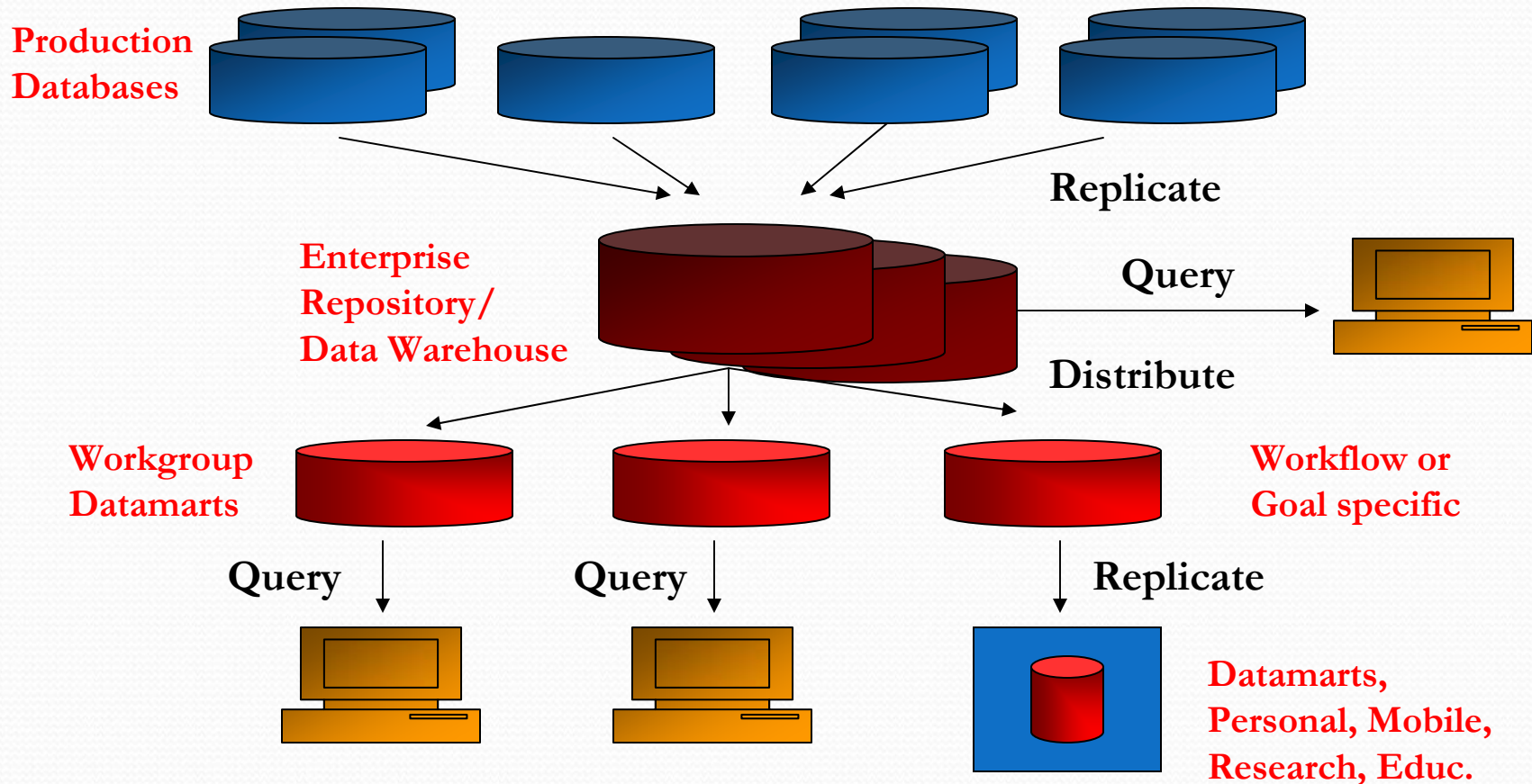
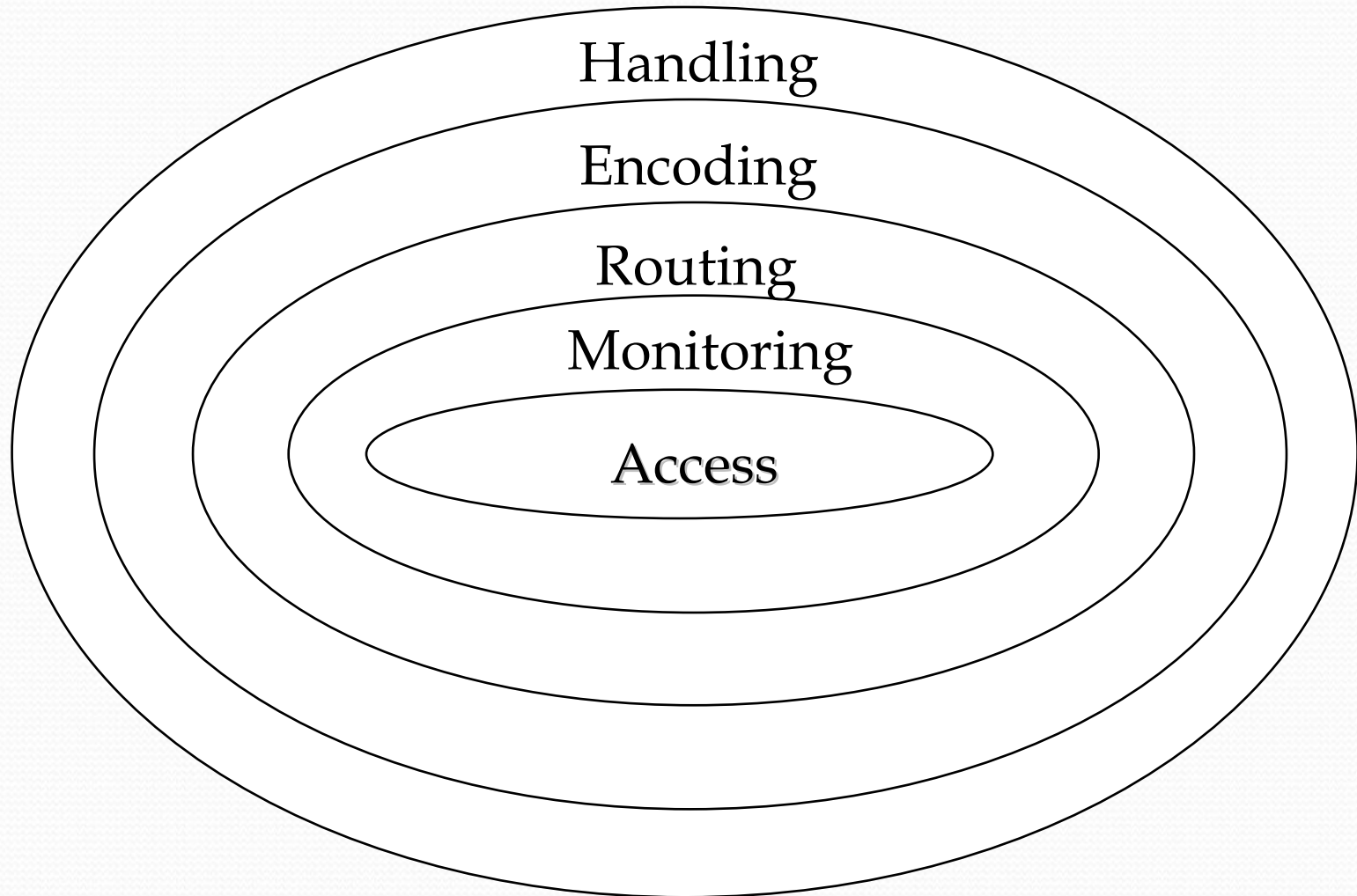


FIGURE B.1

EMR environment



Architecture



Other Level 2 Systems

- Intermountain
- VA
- Partners
- Regenstrief
- Vanderbilt

Level 3 Systems

- Cerner
- Epic
- Eclipsys
- GE
- McKesson

Challenges at Columbia

- Moved from Stage 3 through Stage 4 to Stage 5
- Purchased a vendor system (Level 3)
- How to get to Stage 5 and Level 4?

Challenges at CPMC/CUMC/NYPH/WCMC

- In 1998, merged two academic medical centers into New York Presbyterian Hospital
 - Columbia Presbyterian campus became Columbia University Medical Center
 - New York Hospital became Weill Cornell Medical Center
- Currently 4 different electronic health records
 - Eclipsys (WCMC)
 - Eclipsys (CUMC)
 - Epic (WCMC)
 - Allscripts (CUMC)

NYP Computing Environment
 Sep 2007

NYP

Weill Cornell

ITS
 NYP Email
 NYP Calendar
 WMC Network
 WMC Email, etc.

Core & ITS
 NYP/WMC Network

Information Service
 (Adm), Eagle, BIS,
 (Clin) Eclipsys XA
 WebCIS, Cisyphus.
 (Clin) Amicas, GE PACS
 (Clin) Imnet, Surgical Mgr
 (Adm) Lawson, P'soft, etc.
 (Clin ancil) Cerner, Misys
 MPI, Egate, Data Warehouse
 Network, Desktop, Security

Departments (admin)
 (Clin) My Medical File
 (Clin) Mobile Telemetry (CUMC)
 Pyxis (WMC/CUMC) - OR

Departments (clin)
 Climacs (WMC) - Med
 (Clin) Copath
 Helix (CUMC) - Surgery
 Many others.

LabTest
 Rogosin
 TSI, Eagle
 Psych Inst
 Clearing Houses
 Regulatory

Internet

Service Corp

Cornell PO
 (Adm) IDX
 (Clin) Epic

Research

Departmental

Pub Relations
 www.nyp.org
 Development DB

Core
 CUMC Network
Clin Trial
 Velos
Security

Service Corp

Columbia PO
 (Adm) IDX
 (Clin) Allscripts

CUBHIS
 CUMC Calendar
 Desktop, etc.

Research

Departmental

Internet & CUMS
 CUIT
 CU Email

Sponsored Hospitals
 HSS NYQ NYM CHOB Etc.

Local Clinical and Admin Systems

**Presbyterian
 New York
 MSCHONY
 Westchester
 Allen
 Pavilion**

**Heath Plan,
 etc.**

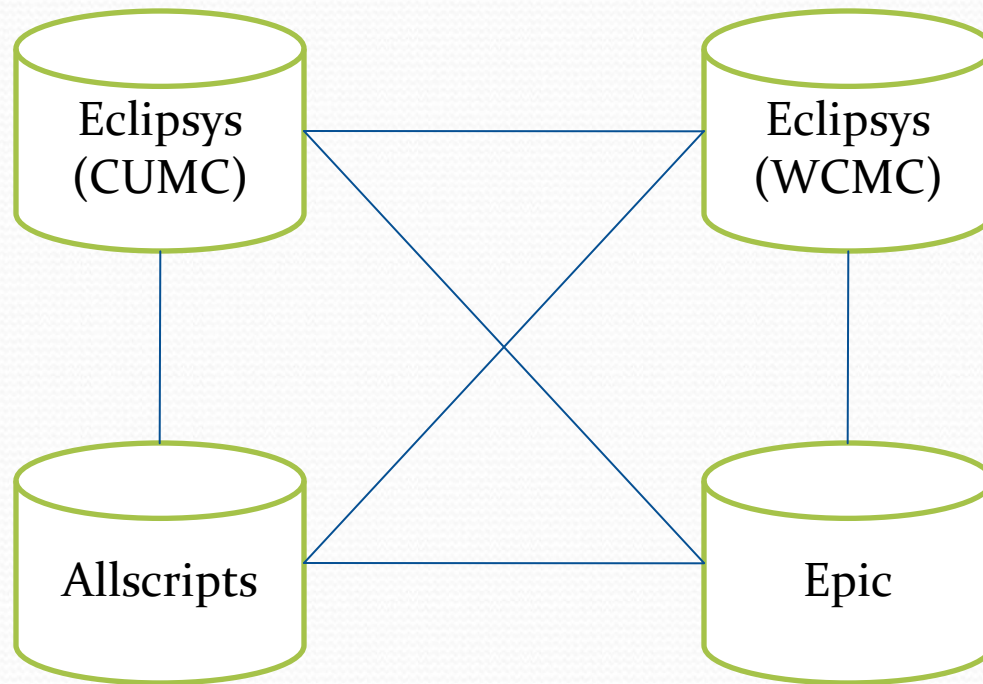
Columbia

St Luke's
 Dept Billing

Harlem

● Also used by some sponsored hospitals
 ● Contractual/Outsourced relationship

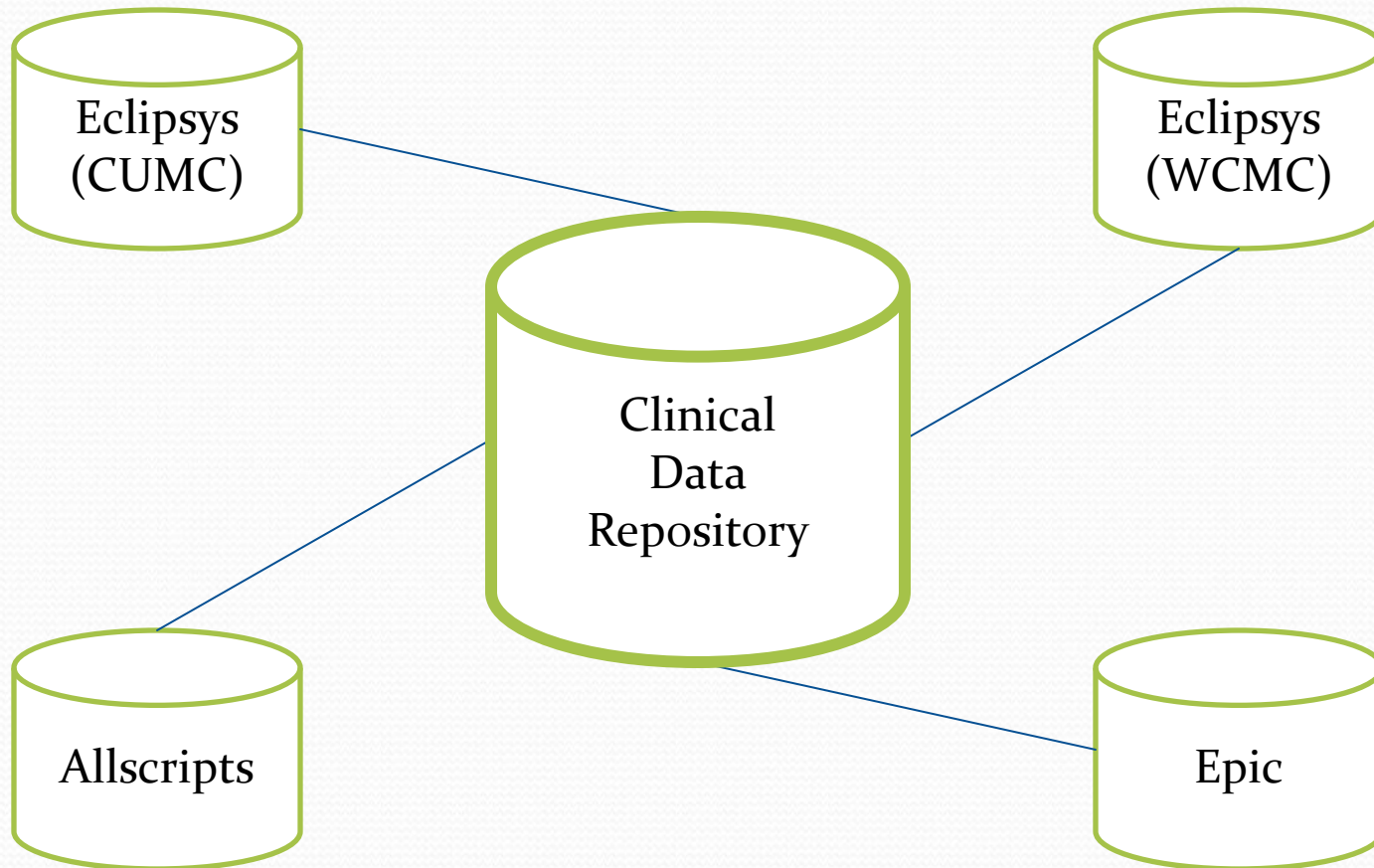
Integrating Among Multiple EHRs



Problems with Integrating to Application Databases

- Must model each system multiple times
 - Increased effort and complexity
- Overloading workflow databases
- Protecting external data consistency (no updates)
- Increased complexity of data protection
- Bringing in data for a new patient
 - When to pull data in
 - Interfaces don't naturally pull in historical data
- Increases complexity as move toward RHIOs

Repository Model



Benefits of CDR

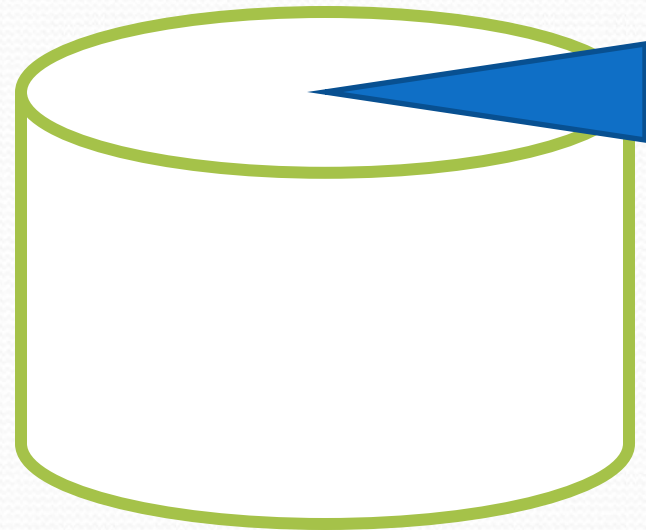
- Only model data from source systems once
- Common data store
- Data are read only
 - Optimized for read
- Historical data included
- Web-based viewer adaptable to multiple applications
- Adaptable to future health information exchange efforts
- Platform of innovation

Optimized for Retrieval

- Relational structure can be difficult to query for both data and context
 - Gathering multiple elements requires multiple table joins
 - Good for data storage
 - Good for aggregating across multiple patients
- Event-based model good for querying across data types
 - Data organized according to patient
 - Not good for querying across patients

Retrieval optimization

- Paradigm shift in how data are used
 - Paper records mainly for primary use
 - Electronic allows secondary use
 - Secondary use can be multiple times



CDR View in Eclipsys

SANDIEGO, CARMEN - Sunrise Clinical Manager

File Registration Edit View GoTo Actions Preferences Tools Help

SANDIEGO, CARMEN 313 13 13 / 000054755 254 75y (03-Jan-1933) Female
 REF-MRAP Joseph, Kathie Ann DOB: 01/03/1933
Allergies:
Current Weight: kg **Height:** cm **BSA:** sq.m **Admit Date:** 23-Jan-2008

Patient List Orders Results Patient Info Summary Documents Flow sheets Clinical Summary CathLab Results OR Manager WebCIS

WebCIS Notes Feb 15
 Signout Nov 26
 Disch Sum 2010
 Operative Feb 13
 OR Notes 2007
 Consult 1997
 Clin Sum 2006
 Laboratory Feb 20
 Radiology 2007
 Pathology 2005
 Neurophys 1997
 Ob/Gyn 2003
 GI Endo 2007
 Cardiology 2007
 HEENT 1997
 Pulmonary 2007
 Endocrinol Oct 09
 Derm Path

Disch Sum (2010-05-18 to 2004-11-05) Newer • Older

Name	Date	Status
Discharge Summary Note	2010-05-18 12:00	F
Discharge Summary Note	2005-03-25 17:35	F
Discharge Summary Note	2005-02-23 10:01	F
Transfer Note	2005-02-23 10:01	P
Transfer Note	2005-02-23 10:01	P
Transfer Note	2005-02-22 08:59	P
Admission Note	2005-02-21 08:56	P
Admission Note	2005-02-20 08:52	P
Admission Note	2005-02-20 08:52	P
Discharge Summary Note	2005-02-20 08:52	P

HISTORY OF PRESENT ILLNESS:
 The patient is a 48 year old woman with a history of heart failure, non-ischemic cardiomyopathy, EF 20% who was evaluated for cardiac transplant. She has had shortness of breath with her ADLs and swelling in her legs and belly with 5 pillow orthopnea and PND, occasional lightheadedness and dizziness, nausea and vomiting with poor appetite. Since November 2004, she has had 3 hospitalizations for heart failure and has progressive symptoms.

PAST MEDICAL HISTORY:
 Dilated cardiomyopathy since 2001, atrial fibrillation, stroke in 2002, BiV/AICD in January 2005.

ALLERGIES:
 None.

MEDICATIONS:
 Home meds: Coreg 6.25 b.i.d., Altace 2.5 a day, Keflex 500 q.6, Insprira 25 a day, Lasix 80 in the morning, 40 at night, Ambien 200 a day, Crestor 10 a day, Dig 0.25 a day.

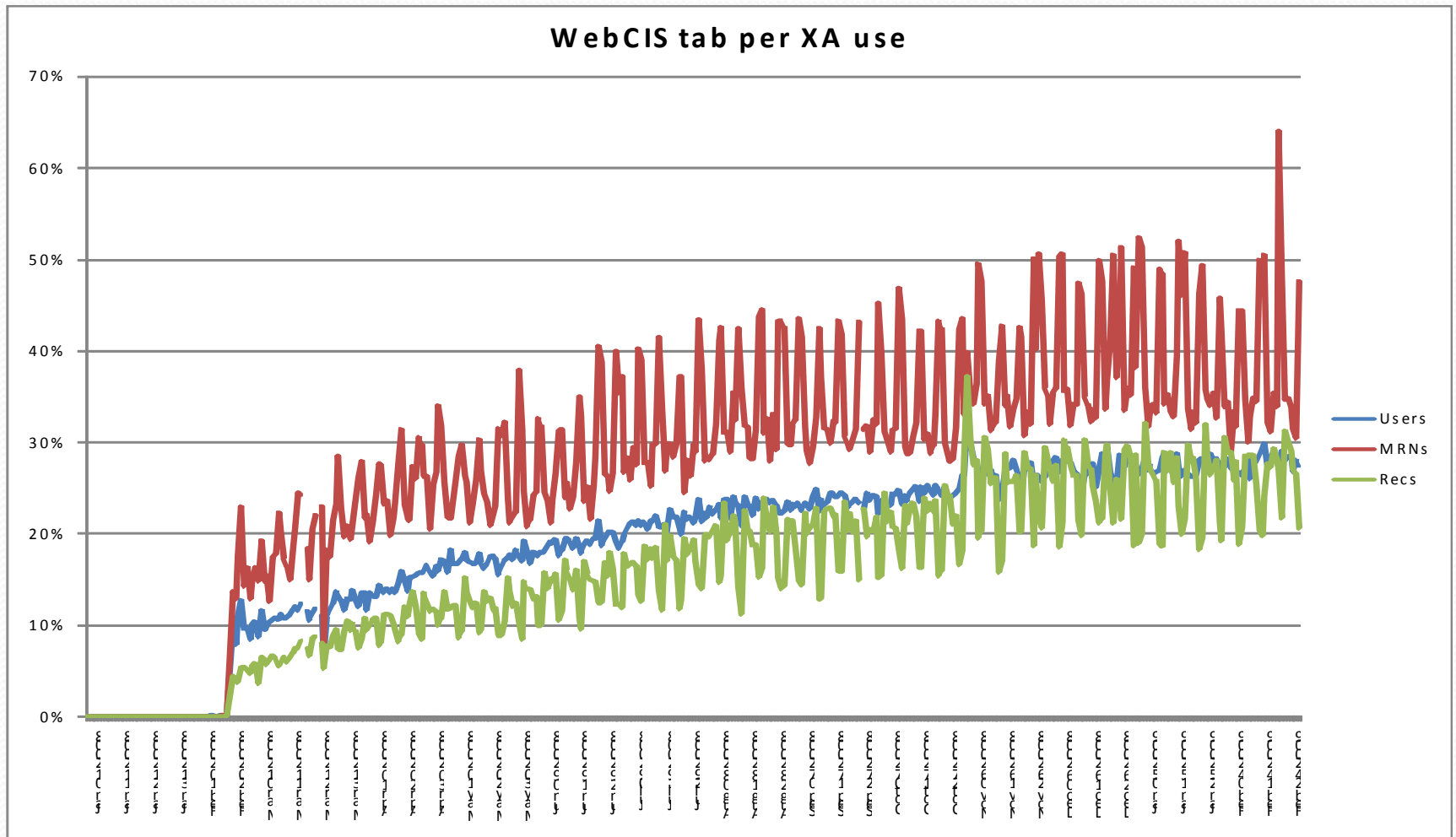
FAMILY HISTORY:
 Brother with cardiomyopathy?

SOCIAL HISTORY:
 No tobacco, alcohol or drugs, divorces. She has 6 children. She had Medicaid and she lives in Long Island.

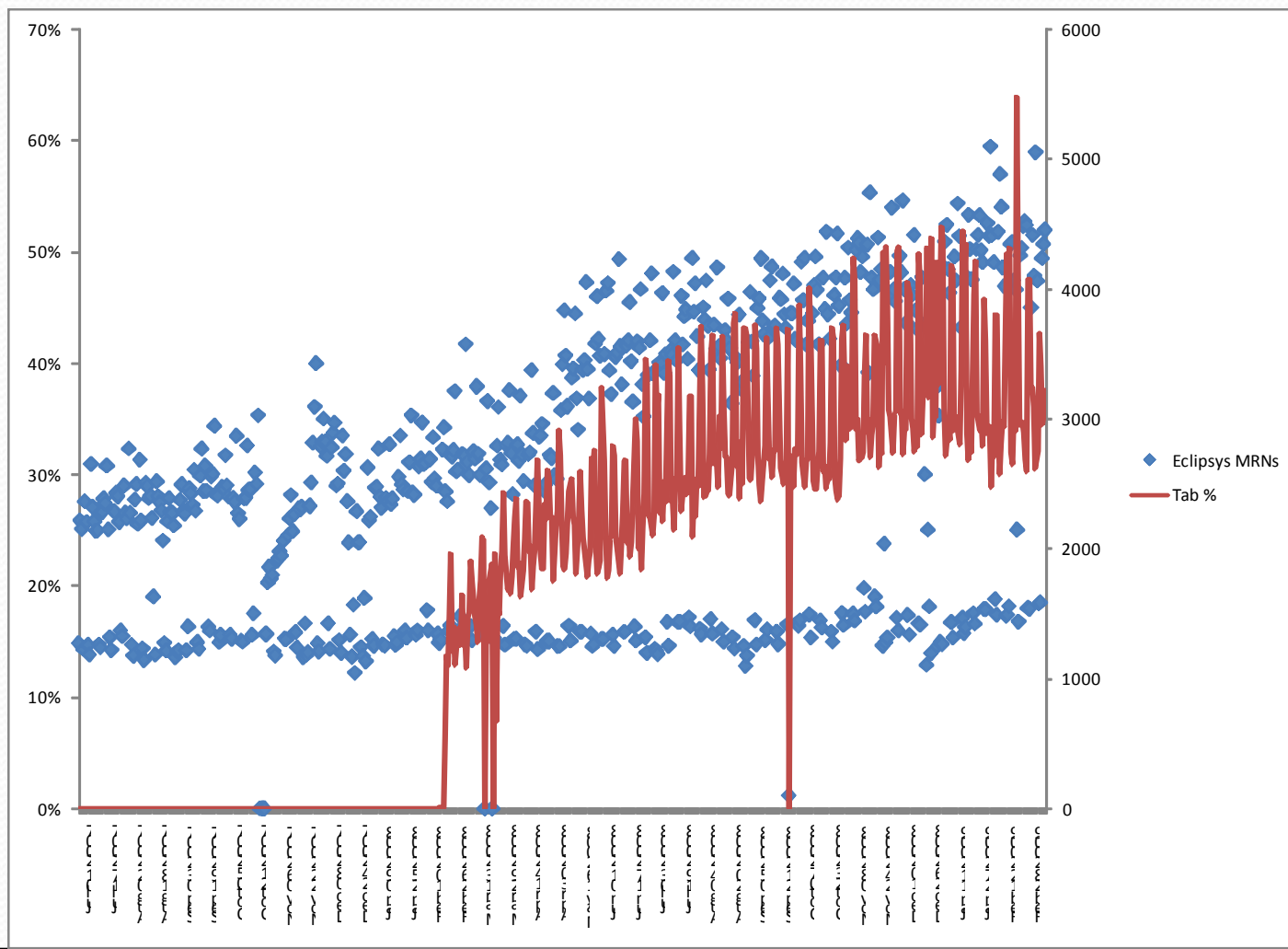
PHYSICAL EXAMINATION:
 She was seen on physical exam on admission: Blood pressure 20/60, both lying and standing with a pulse of 80. She was ill-appearing in moderate respiratory distress. She had JVD to 15 cm, mild crackles at the bases and S3 exam and HJR in her abdomen exam, 1+ edema in lower extremities and nonfocal neuro exam.

Ready

Proportion of CDR Viewer Access



Increase in CDR View Access



CUMC/NYP Clinical Data Warehouse History

- ▶ 1994: Created, sponsored by Columbia University Department of Medical Informatics and Office of Clinical Trials
 - Populated with data from existing clinical data repository
 - Supporting clinical research
- ▶ 1998: Columbia + Cornell = NewYork Presbyterian Hospital
 - Warehouse funded by NYPH
 - Goal to incorporate and provide data across whole system
- ▶ 2004: Formal analysis of CDW user needs by Clinical Quality and Information Technology Committee (CQIT)
 - Creation of Data Warehousing Subgroup
 - Need to bring together disparate clinical data sources
 - Need to manage user requests for data

Uses of the Warehouse

- Clinical research queries
- Management reports
- Clinical trial recruitment

CDW Content Issues

- Began as a copy of the repository
 - Data already gathered
- Mainly for research queries
 - Some data marts built for common queries
- Ability to query rapidly across patients increases security risk

Information Request - Windows Internet Explorer

http://ctcc.cpmc.columbia.edu/rdb/request.html

File Edit View Favorites Tools Help Google G Go Bookmarks Settings

Information Request

Clinical Data Warehouse at CUMC

Mediated Query

Please see **Important Riders, Approval Requirement** and a **Fee Policy** at the bottom of this page.

NAME	<input type="text"/>
DEPARTMENT	<input type="text"/>
TITLE&RANK	<input type="text"/>
ROLE IN PROJECT	<input type="text"/>
OFFICE ADDRESS	<input type="text"/>
TELEPHONE	<input type="text"/>
EMAIL	<input type="text"/>
PROTOCOL (PROJECT) NAME AND A BRIEF DESCRIPTION	<input type="text"/>
	<small>(255 chars max. Use comments if you need more space)</small>
PROJECT SPONSORSHIP/REASON	<input type="text" value="Investigator Initiate"/>
PRINCIPLE INVESTIGATOR	<input type="text"/>
DATE of REQUEST	<input type="text" value="2/7/2007"/>
DATE REPORT IS REQUIRED	<input type="text"/> e.g. 12/31/2005

Done Internet 100%

Information Request - Windows Internet Explorer

http://ctcc.cpmc.columbia.edu/rdb/request.html

File Edit View Favorites Tools Help

Information Request

RIDERS ON PATIENT DATA	APPROVALS	FEE POLICY
<ul style="list-style-type: none">I understand that this data is confidential and I will follow strict procedures to preserve confidentiality in dealing with patient-specific information, in order to protect the privacy rights of individuals as well as the interests of Columbia University and New York Presbyterian Hospital.I further understand that this data is for my own use and those of my direct close co-collaborators only, and I agree not to release or distribute this information, in any form, to any less closely affiliated person or organization, regardless of institutional or organizational affiliation.I understand all requests for data will be reviewed by the Office of Clinical Trials, and final approval for a response to my request rests solely with the Office.I agree that the acquired data will be destroyed once it is no longer required.	<ul style="list-style-type: none">RESEARCH PROJECTS AND PUBLICATIONS: Approval of the CUMC Institutional Review Board (IRB) is required.CLINICIAN ACCESSING DATA ONLY ON HIS OR HER OWN PATIENTS: A signed letter on the clinician's letterhead attesting to this and specifying the desired information.DEPARTMENTAL REVIEW OR GRAND ROUNDS: A signed letter of approval from the sponsoring attending physician specifying the information desired.SUMMARY INFORMATION: No formal letter or specification is required, unless the request becomes resource-intensive or requires extensive analysis.	<ul style="list-style-type: none">Please note that a chargeback methodology is being developed for research-related requests for data from the Clinical Data Warehouse. An important factor in this methodology will be the number of hours of analysts' work involved. We'll estimate and notify you total cost once your Clinical Data Warehouse request is approved.

AGREE CLEAR

Done Internet 100%

Goal of Access Policy

- Provide broader access to data
 - Central control is resource limited
- Allow collection of more data sources
 - Reassure data stewards
 - Three separate institutions
 - Data ownership not completely defined for all data

CDW Structure

- Identifying data
 - Patient identifying information
- Main data
 - Event tables for clinical repository
- Lookup tables
 - Vocabulary translation
 - Contains no patient data
- Specialty data marts

Access Policy

- Identifying data
 - Most restricted
 - Create a research identifier to replace the patient ID
 - Allow access to only ResearchID, sex, birth date (month and year only), marital status, race, death status
- Specialty data
 - Access policy defined by data steward
- Patient clinical data
 - No access to text data
 - Modified dates
- Lookup tables
 - Full access (contain no patient data)

Access Policy

- Specific patient information
 - Sometimes needed to create initial queries
 - Analysts get access only to a randomly selected subset
 - Access request through supervisor
- De-identified patient data
 - Test patients
 - Full access given

CUMC/NYP Clinical Data Warehouse History

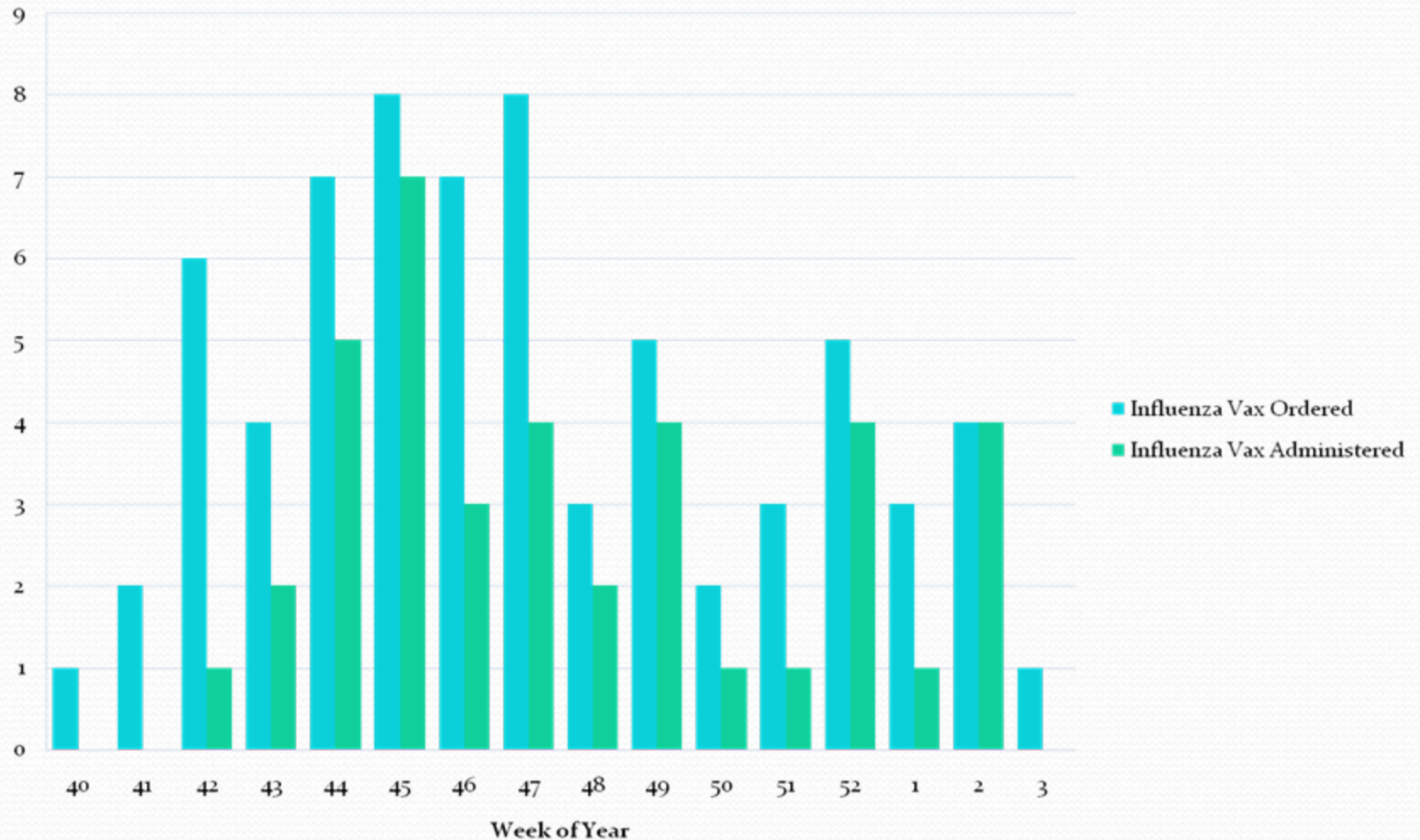
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 - **Need to bring together disparate clinical data sources**
 - **Need to manage user requests for data**

Analysis of Challenges

- Data in vendor-based transactional systems
- Could not query across transactional systems
- Users needed help in defining their needs
- Mature initiatives required more robust data solutions

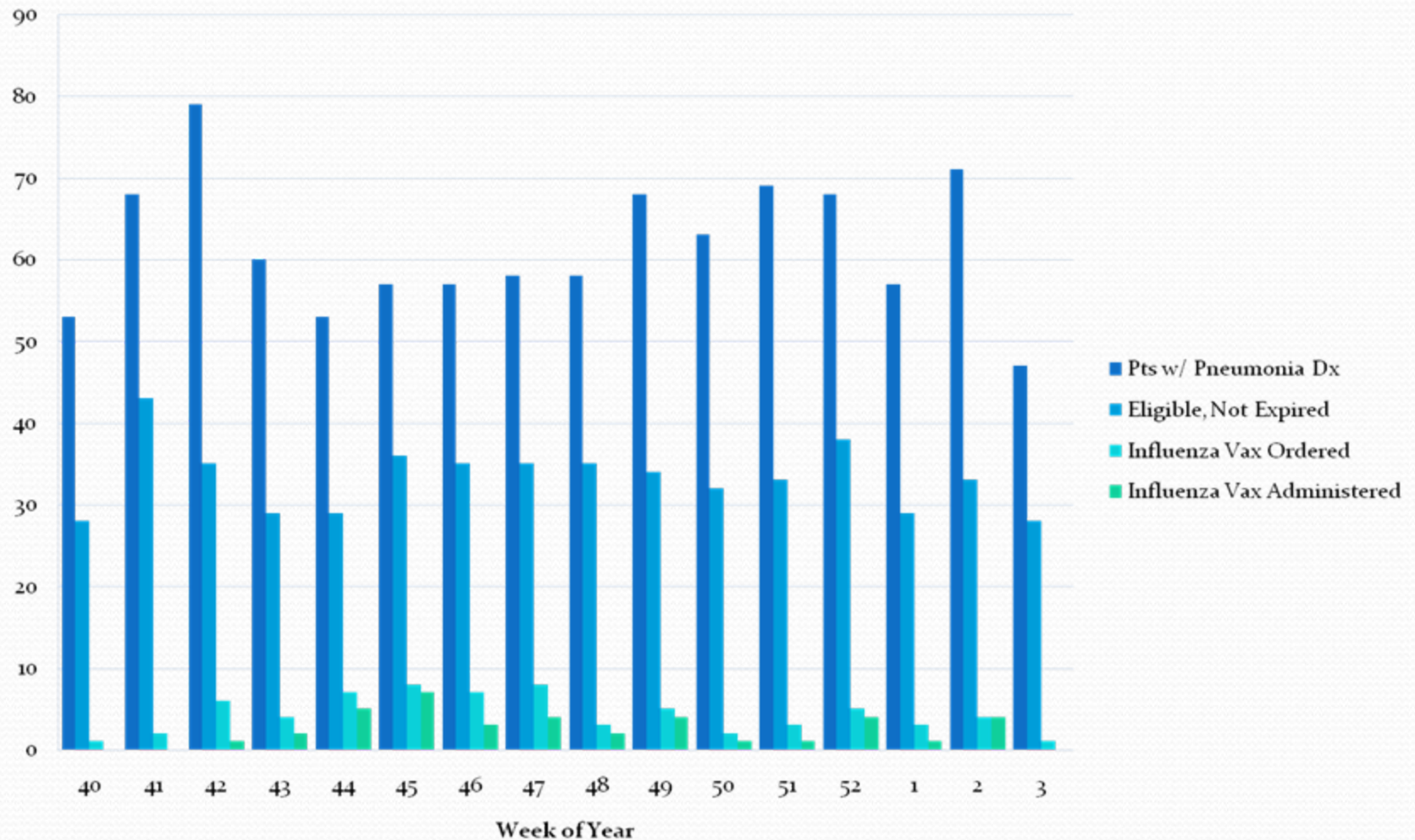
Pneumonia Core Measures

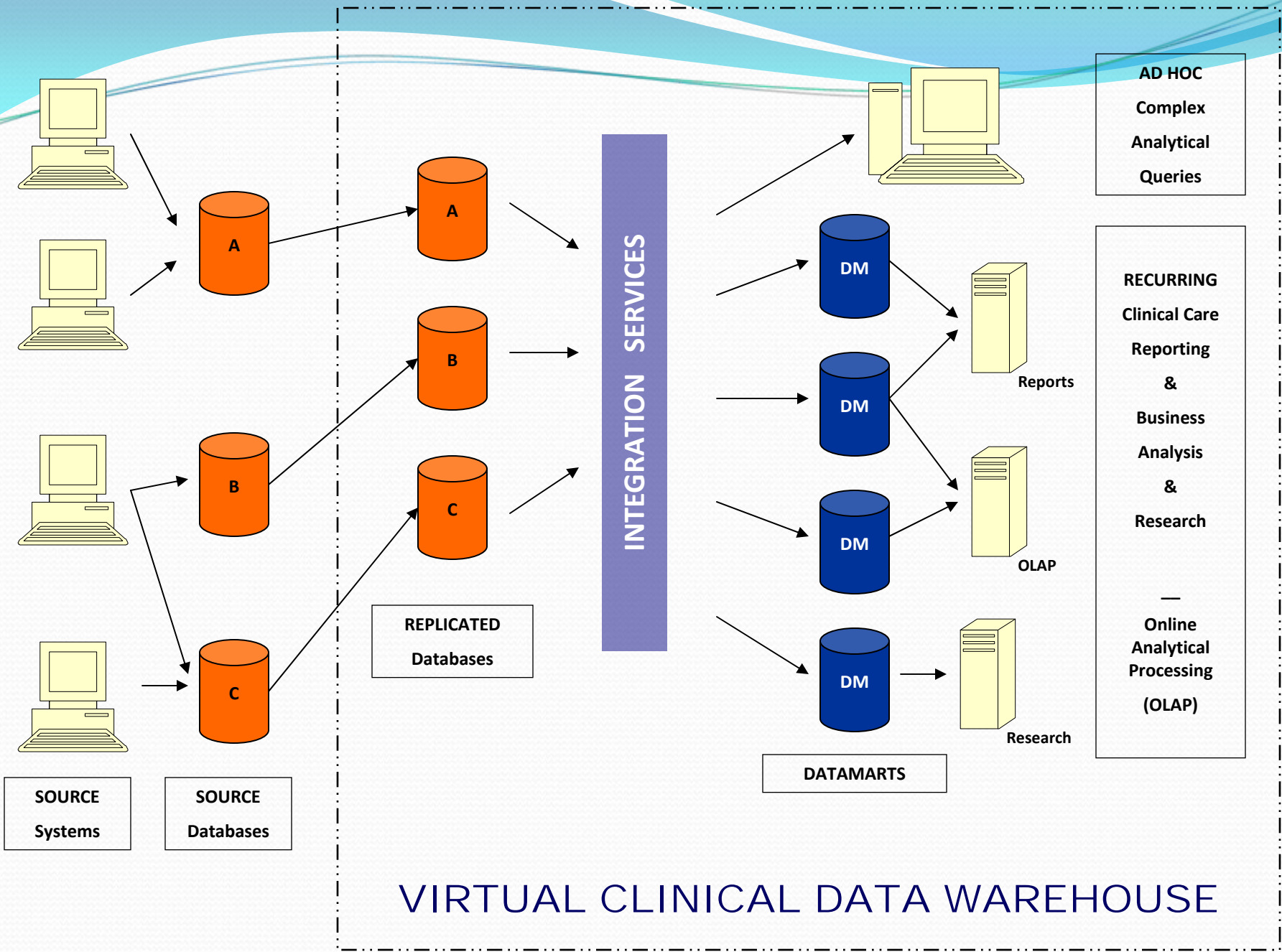
CUMC Influenza Vaccinations Q4 2008 - Q1 2009



Pneumonia Core Measures

CUMC Influenza Vaccinations Q4 2008 - Q1 2009





VIRTUAL CLINICAL DATA WAREHOUSE

SOURCE Systems

SOURCE Databases

REPLICATED Databases

INTEGRATION SERVICES

DATAMARTS

AD HOC Complex Analytical Queries

RECURRING Clinical Care Reporting & Business Analysis & Research

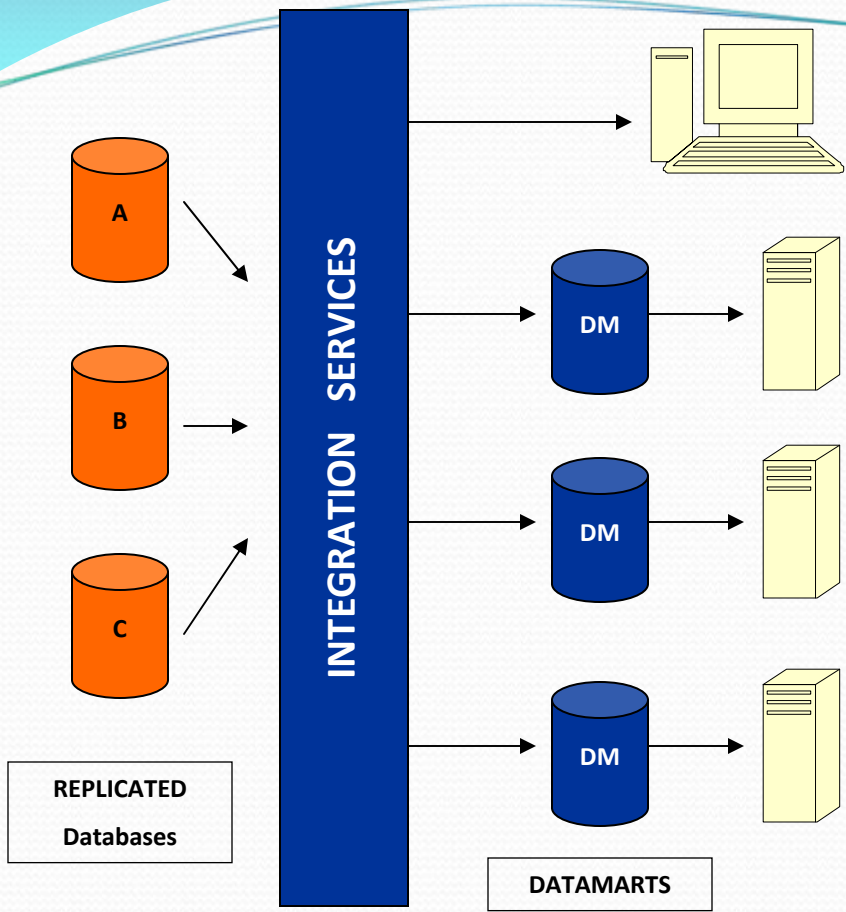
— Online Analytical Processing (OLAP)

Reports

OLAP

Research

Goal	Task	Use	User	Tool	Six Sigma	Cost/ Instance	Instances	Required
Answer a specific question	Ad hoc query	Research	Researcher	SQL	<i>Define</i>			Defined request
Observe trends	Recurring query	Management reports	Manager	Reporting application	<i>Measure</i>			Available owner
Identify dependencies	OLAP	Operational analysis	Analyst	Analytics / Data cubes	<i>Analyze</i>			Content expert/ analyst
Assist decision making	Dashboard display	Point of care	Clinical team	Registries	<i>Improve</i>			Pilot site
Automate processes	Application	Decision support	Clinician/ Role	EMR application	<i>Control</i>			Institutional sponsor



VIRTUAL DATA WAREHOUSE

Ad-Hoc Queries – Questions	Research	Define
Recurring – Automated Queries	Management Reports	Measure
OLAP – Analytics	Operational Reports	Analyze
Dashboards	Point of Care Reporting	Improve
Applications	Decision Support	Control

DATA WAREHOUSE TOOLS

Black Belt Six Sigma Approach

Conclusion

- Integrating clinical data repository view into workflow applications can improve use
- Access policies need to isolate data to reassure data use from different stakeholders
- Data access tools need to account for users' evolving data needs along the quality improvement life cycle