Biomedical Informatics at the University of Chicago: A Multidisciplinary Approach

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Research Associate - Associate Professor



The Biomedical Research "Cloudscape"



The science machine



UC Research IS structures



The CBIS Academic and Research Applications Group





Current Situation





Clinical Research Data Integration



The BSD/BRI Initiative Data exchange architecture



Infra-Structure demands

• Storage

Networking



Growth of Storage by Fiscal Year



Researcher



Data Acquisition: User entered, LIMS, Collected by Instrumentation, etc.

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Application Server



Integrated Solution Workflow





Velos Usage

The University of Chicago is currently using Velos in many of their operational processes.

Daily Use	Database	Reports
Clinical Trials Regulatory Management	Clinical Trials Review Committee	Clinical Data Upload System (CDUS)
Clinical Trials Data Management	Scientific and Accrual Monitoring Committee	Grant Reporting
Listing of Open Protocols on the Intranet	Regulatory Manager Dashboard	Study IRB Expiration Report
Consent Assess For Clinics	Supervisory Dashboard	NCI Summary 4 Report



Velos Current Stats

- Patients ~ 26,000
- Active Studies (2,849)
 - Surgery 213; Neurology 31; Medicine 930; Pediatrics 15; Hem/Onc 1670 (1280 - open to accrual, 390 -closed to accrual)
- Users ~ 270
 - Suregery 21, Neurology 7, Medicine 65, Hem/Onc 143, Biostats 4, Investigational Pharmacy 3, OCR 5, CBIS 2



Current use of Velos at UofC

Departments	
Health Studies	
CBIS	
Cancer Research Cent	ter
General Clinical Resea	arch
Gynecology/Oncology	Ý
Hematology/Oncolog	У
Pulmonary/Critical Ca	ire
Medicine/Oncology	
Office of Shared Rese	arch Facilities
Clinical Research Supp	port Office
Neurology	
Gynecology/Obstetric	2
Office of Clinical Rese	arch
Psychology	
Pediatrics	
Pediatrics Oncology	
Radiology	
Radiology/Oncology	
Surgery	
Surgery/Oncology	
University of Chicago	Pharmacy
Medicine	
Neurology/Oncology	
Cancer Clinical Trials (Office

Office of Clinical Research: Faculty and Staff



Office of Clinical Research: Faculty and Staff



http://clinicalresearch.bsd.uchicago.edu/faculty_staff/velos/velos-comm.shtml (1 of 2) [5/11/2009 12:46:18 AM]

Forms Samples

$\frac{C H I C A G O}{CANCER RESEARCH CENTER}$	
Surgery date: Surgery type:	☐ None ☐ Highest mediastinal ☐ Upper paratracheal
Histology: If other: Path size:	Pre-vascular and retrotracheal Lower paratracheal (incl. Azygo Subaortic (A-P window) Para-aortic (ascending aorta or p
Lymph node dissection? C Yes C No Lymph nodes removed: Lymph nodes positive: Positive lymph node location(s): @	I Subcarinal Paraesophageal (below carina) Pulmonary ligament Hilar Interlobar Lobar
Grade: TTFI: C Not Done T Path: C Negative N Path: C Positive M Path:	□ Segmental □ Subsegmental ➤ Cancel ≏ Reset ▶ Done



Specimen Submission

Instructions: Send blocks or slides and blood for each patient enrolled to the University of Chicago. Include original copy of this form with sample then place one copy in study chart. Please refer to section 8.0 of the protocol for shipping address and number of samples required. I APPE ALL SPECIMENS WITH PATIENT STUDY NUMBER AND INITIALS

ue Block Samp	les	
Tumor	Surgical procedure date: Total number received:	<u>Specimen Bar Codes</u> (max: 5) 1
Normal	Surgical procedure date: Total number received:	
Lymph Node	Surgical procedure date: Total number received:	Specimen Bar Codes (max: 5) 1

	Surgical procedure date:		
Tumor	20µm unstained slides: (qty) <u>Specimen Bar Codes</u> (<i>mex:</i> 5) 1	5µm unstained slides:	
	Surgical procedure date:		
Vormal	20µm unstained slides: (qty) <u>Specimen Bar Codes</u> (<i>max:</i> 5) 1♣	5µm unstained slides:	<u>+</u>
	Surgical procedure date:		
lormal	20µm unstained slides:	5µm unstained slides:	÷
	Surgical procedure date:		
Lymph Node	20µm unstained slides: (qty) <u>Specimen Bar Codes</u> (mex: 5) 1.	5µm unstained slides: (qty) Specimen Bar Codes (max: 20) 1.	*

		Bar Code	Amount	Storage Location
	1.			-80° freezer in E203 💌
Plasma	2.			-80° freezer in E203 💌
C	1.			-80° freezer in E203 💌
Serum	2.			-80° freezer in E203 💌
DNA	1.			-80° freezer in E203 💌
DNA	2.			-80° freezer in E203 🐱

Examples of Forms Adverse Events

Preview

Form Name: Internal AE - Serious and Unexpected

INSTITUTIONAL REVIEW BOARD

OFFICE OF RESEARCH SERVICES McGiffert Hall, 2nd Floor 5751 S. Woodlawn Ave., Chicago, Illinois 60637 Phone: / Fax: 773-834-0659

INTERNAL ADVERSE EVENT FORM: SERIOUS AND UNEXPECTED EVENTS

For adverse events occuring at the U of C Hospitals or a hospital affiliated with the University of Chicago

An adverse event is an undesirable and unintended, although not necessarily unexpected, result of therapy or other intervention.

Serious adverse events include: death, a threat to life, hospitalization, prolongation of existing hospitalization, persistent or significant anomaly or birth defect, causes cancer, is an overdose, any medical event which requires treatment to prevent one of the medical outcomes listed. (Please note: Do not report death or life-threatening events on this form.)

Unexpected adverse events include: those for which the specificity or severity is not consistent with any of the following: the investigator brochure, protocol, risk information in the consent form, the reasonably expected natural history and progression of the underlying disease or condition.

For drug/biologic studies - Any adverse event that occurs that is **both** serious and unexpected must be reported to the IRB. In addition, adverse events occur that are moderate in severity and are not necessarily serious and unexpected, but in the investigator's opinion, should be considered by the IRB due to a possible relationship with the drug/biologic being studied. In both of these cases the adverse event should be reported to the IRB within 10 working days of the investigator's knowledge of the adverse event.

For device trials, any unexpected adverse events must be reported to the IRB within 10 days of the investigator's knowledge of the event.

A. GENERAL INFORMATION	
Protocol Number:* Find Numb	xer/Title
Principal Investigator Name:* <u>Find</u>	Principal Investigator Email:*
Treating Physician Name:*Find Phy	rsician Treating Physician Email:*
Quality Assurance Coordinator:*	Find Coordinator Quality Assurance Coordinator Email:*
Protocol Title:	

B. EVENT INFORMATION

https://velos.bsd.uchicago.edu/eres/jsp/formpreview.jsp?formId=428 (1 of 3) [5/11/2009 12:19:46 AM]

Examples of Forms **BP Form**



Preview

Form Name: 12-Hour Blood Pressure Collection

12-Hour Blood Pressure Collection

Visit:	Average systolic BP:	
Date:	Average diastolic BP:	

	BP Index	Time	Systolic	Diastolic	Mean	Heart Rate
1						
2						
3						
4						
5						
6						
7						
8						
9						
10						
11						
12						
13						
14						
15						
16						







Clinical Trials at The University of Chicago Cancer Research Center

Web Application **Protocol Listing**





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CANCER RESEARCH CENTER

UCCRC Home

Open Protocols

Select Disease Site: Advanced Disease (Phase I) Bladder Bone Brain and Nervous System Breast Cancer Control Cancer Risk Cervix Colorectal Esophagus Gastrointestinal (GI) Other Gastrointestinal Stroma (GIST Sarcoma) Head and Neck Hodgkins Lymphoma Kidney/ Renal Leukemia Liver/ Hepatobiliary Lung/ Chest Melanoma (skin) Mesothelioma Multiple Myeloma Myelodysplastic Syndrome (MDS)

https://med-www02.bsd.uchicago.edu/crc_web/SR/internet/DiseaseSites.aspx (1 of 2) [5/11/2009 12:31:30 AM]

Web Application Protocol Listing



UCCRC Home

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Open Protocols > Protocols by Disease Site

Clinical Trials at The University of Chicago Cancer Research Center

Leukemia Cancer Studies Currently Recruiting Participants at the University of Chicago Cancer Research Center

Protocol Title	IRB #	Physician
CALGB C10001/SWOG C10001: A Phase II Trial of Sequential Chemotherapy, Imatinib Mesylate (Geeevec,STI571) &" Transplantation for Adults with Newly Diagnosed PH+ Acute Lympoblastic Leukemia by the CALGB and SWOG	11700A	Stock, Wendy
CALGB 10002: A Phase II Study of Rituximab and Short Duration, High Intensity Chemotherapy with G-CSF Support in Previously Untreated Patients with Burkitt Lymphoma/Leukemia	11752B	Stock, Wendy
High-dose Cytarabine/Mitoxantrone followed by Autotransplantation for t-MDS/ t-AML	11884A	Godley, Lucy
CTSU E2902: A Phase III Randomized Study of Farnesyl Transferase Inhibitor R115777 in Acute Myeloid Leukemia (AML) Patients in Second or Subsequent Remission or in Remission after Primary Induction Failure	13686B	Rich, Elizabeth
An open-Label Extended-Use Study of Oral CEP-701 in Patients with Hematologic and Non-Hematologic Malignancies Who Have completed a Clinical Study of CEP-701	14054B	Godley, Lucy
Phase I Study of PXD101 in Combination with 5-Azacitidine (5-Aza) for Advanced Hematologic Malignancies	14510A	Odenike, Olatoyosi
	14649A	Stock, Wendy
A Randomized, Double Blind, Placebo Controlled Study Evaluating the Efficacy and Safety of AMG 531 Treatment of Subjects with Low or Intermediate-1 Risk Myelodysplastic Syndrome (MDS) Receiving Lenalidomide.	15070B	Larson, Richard



Web Application **Protocol/ Consents**



Protocols by Program

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HE UNIVERSITY OF CHICAGO

CANCER RESEARCH CENTER

UCCRC Home

Programs > Protocols by Program

Logout

CRC_Breast Program

Title	Consent	Protocol	I.Brochure	Schema	IRB#	PI
Multicenter Phase III						
Randomized Trial						
Comparing						
Doxorubicin and						
Cyclophosphamide						
Followed by						
Docetaxel with						
Docetaxel and						
Cyclophosphamide						
Followed by						
Docetaxel and						
Trastuzumab and						
with Docetaxel,					10964B	Olopade
Platinum Salt and						
Trastuzumab (TCH)						
in the Treatment of						
Node Positive and						
High Risk Node						
Negative Adjuvant						
Patients with						
Operable Breast						
Cancer Containing						

https://med-www02.bsd.uchicago.edu/crc_web/SR...spx?tarea_id=231&prog_name=CRC_Breast Program (1 of 9) [5/11/2009 12:41:12 AM]

Regulatory Dashboard Application

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THI	EUNIVERSITYOF					
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	CANCER RESEARCH CENTER					
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Welcome	sbapat					
Digital D	ashboard:					
Digitai D						
Reco	rd Complete. Click for further details.					
	rd Complete. Click for further details. Ite Required. Click for further details.					
Recc	rd Complete. Click for further details. ite Required. Click for further details. ie - action may be required.					
Recc Upda Notic	rd Complete. Click for further details. Ite Required. Click for further details. Ite - action may be required. not applicable (non-clickable)					
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Recc Upda Noti Item Study Number	rd Complete. Click for further details. te Required. Click for further details. te - action may be required. not applicable (non-clickable)	PI Olopade, Olufunmilayo	IRB A	Affiliate	Study Accrual	eVelos Inconsistencies
Recc Upda Notic Terr Study Number 10964B	rd Complete. Click for further details. te Required. Click for further details. te - action may be required. not applicable (non-clickable) Study Title Phase III Doxorubicin + Cyclophosphamide + Docetaxel Followed by Docetaxel and Trastuzumab + Platinum Salt in the Treatment of Node + and High Risk Node - Adjuvant Patients with Operable Breast Cancer Phase I Study of R(+)XK469 in Patients with Advanced Solid Tumors and Lymphoma	PI Olopade, Olufunmilayo Undevia, Samir	IRB A	sffiliate	Study Accrual	eVelos Inconsistencies
Recc Upda Notiv Tterr Study Number 10964B 11108B 11249A	rd Complete. Click for further details. te Required. Click for further details. te - action may be required. not applicable (non-clickable) Study Title Phase III Doxorubicin + Cyclophosphamide + Docetaxel Followed by Docetaxel and Trastuzumab + Platinum Salt in the Treatment of Node + and High Risk Node - Adjuvant Patients with Operable Breast Cancer Phase I Study of R(+)XK469 in Patients with Advanced Solid Tumors and Lymphoma A Phase I Escalating Multiple Dose Study with Continuous Dosing of ABT-751 in Patients with Advanced Cancer	PI Olopade, Olufunmilayo Undevia, Samir Fleming, Gini	IRB A	affiliate	Study Accrual	eVelos Inconsistencies
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Regulatory Dashboard Application

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Study Number V			Exposure and		to Accrual	0600 2007				Study	eVelos Inconsistencies
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SPFORAL45 A	Admini										
SPFORAL45 A	A 12-W Admini	No-Accrual or Accrual below 10% of stated goal									
Mock - A	A 12-W										
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Mock Study 1 T	Fest sti]		000000000000000000000000000000000000000	100000000000000000000000000000000000000		Omberto				
Mock Study 2 A Phase II trial of fictional drug QTISZAJM in patient with Non-Hodgkins Lymphoma Tachinardi, Umberto											
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The Velos "Core Team" at UofC

UCCRC

Marcy List Amber Burnett Jia Cheng Eugene Limb Swapna Bapat Arthur Christoph Consuelo Skosey

OCR

Dionisia Saner Bethany Martell

Department of Medicine

Don Saner Nick Shank Tim Holper

Department of Pediatrics

Eneida Mendonca Michael Rose

CBIS

Seigmund Johnson Ryan Crist Clyde Danganan

& many more....



Translational Data Mart (TraM): What Is TraM and What Can TraM Do?

Xiaoming Wang April 22, 2009

What Is TraM and What Does It Do?

- TraM is a healthcare data integration warehouse with a web-based application interface: https://tram.uchicago.edu
- TraM is a "generic" system that can be used with a broad range of healthcare data and has been customized for all types of cancer data, not just breast cancer
- TraM can provide an integration environment for various biomedical domain databases
- TraM is scalable and configurable but still a work in progress

Data Unification and Integration Workflows at UCMC



Dynamic Statistics (not yet synced to source data)

Cancer Cases in TraM

Cancer Site	Cases	Percentage of Total
BLOOD, BONE MARROW, & HEMATOPOIETIC SYS	2199	6.22
BONES & JOINTS	163	0.46
BRAIN, & CRANIAL NERVES, & SPINAL CORD, (EXCL. VENTRICLE, CEREBELLUM)	659	1.86
BREAST	5199	14.70
CERVIX UTERI	800	2.26
CONNECTIVE & SOFT TISSUE	775	2.19
CORPUS UTERI	944	2.67
ESOPHAGUS	469	1.33
GALLBLADDER & EXTRAHEPATIC BILE DUCTS	224	0.63
GUM, FLOOR OF MOUTH, & OTHER MOUTH	265	0.75
HYPOPHARYNX	104	0.29
KIDNEY	1530	4.33
LARGE INTESTINE, (EXCL. APPENDIX)	2004	5.67
LARYNX	382	1.08
LIVER	452	1.28
LUNG & BRONCHUS	3375	9.54
LYMPH NODES	1256	3.55
MENINGES (CEREBRAL, SPINAL)	215	0.61
OROPHARNYX	223	0.63
OVARY	761	2.15
PANCREAS	917	2.59
PLEURA	277	0.78
PROSTATE GLAND	5088	14.38
RECTUM	684	1.93
RETROPERITONEUM & PERITONEUM	138	0.39
SALIVARY GLAND	116	0.33
SKIN	859	2.43
SMALL INTESTINE	184	0.52
STOMACH	662	1.87
TESTIS	195	0.55
THYROID GLAND	807	2.28
TONGUE	411	1.16
UNKNOWN	648	1.83
URINARY BLADDER	985	2.78
UTERUS, NOS	92	0.26
UTVAGINA & LABIA	135	0.38
VULVA, NOS	203	0.57
OTHER	973	2.75
TOTAL	35373	100.00

Support Data Mining (Longitudinal)

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Pub ID	Ethnicity	Age	Gender	IRB	Clinical Diagnosis	Treatment	Pathology		Specimen	Biomarker
TR_0002179	BLACK (AMERICA) AFRICAN AMERICAN	46*	FEMALE	10760B	BREAST - NOS DUCT CARCINOMA 01/31/2000@AGE 40	TOTAL (SIMPLE) MASTECTOMY : 02/21/2000	MALIG-PRIMARY (INVA DIFFERENTIATED :DU	SIVE):POORLY ICT CARCINOMA	BREAST:RIGHT	PR:POSITIVE ER:POSITIVE
TR_0002279	BLACK (AMERICA) AFRICAN AMERICAN	56	FEMALE	10760B	BREAST - NOS DUCT CARCINOMA 01/28/2000@AGE 47	UNKNOWN : LUMPECTOMY OR EXCISIONAL BIOPSY : 01/28/2000	MALIG-PRIMARY (INVA DIFFERENTIATED :DU	SIVE):POORLY ICT CARCINOMA	BREAST:RIGHT	PR:NEGATIVE ER:NEGATIVE
TR_0002337	UKNOWN	59*	FEMALE	10760B	BREAST - NOS DUCT CARCINOMA 07/15/2000@AGE 54	LUMPECTOMY OR EXCISIONAL BIOPSY : 07/15/2000 RADIATION : 08/15/2000 ADRIAMYCIN (DOXORUBICIN)::CYCLOPHOSPHAMIDE (CYTOXAN) : 08/15/2000	MALIG-PRIMARY (INVA DIFFERENTIATED :DU	SIVE):POORLY ICT CARCINOMA	BREAST:RIGHT	PR:NEGATIVE ER:POSITIVE
TR_0002458	BLACK (AMERICA) AFRICAN AMERICAN	81*	FEMALE	10760B	BREAST - UPPER-OUTER DUCT CARCINOMA 04/06/2000@AGE 78	UNKNOWN : MODIFIED RADICAL MASTECTOMY : 07/07/2000	MALIG-PRIMARY (INVA DIFFERENTIATED :DU	SIVE):POORLY ICT CARCINOMA	BREAST:RIGHT	PR:NEGATIVE ER:NEGATIVE
TR_0002731	WHITE	43*	FEMALE	10760B	BREAST - OVERLAP LES DUCT CARCINOMA 05/26/2000@AGE 42	ADRIAMYCIN (DOXORUBICIN)::CYCLOPHOSPHAMIDE (CYTOXAN)::PACLITAXEL (TAXOL): 07/28/2000 REEXCISION OF BIOPSY: 05/26/2000	MALIG-PRIMARY (INVA DIFFERENTIATED :DU	SIVE):POORLY ICT CARCINOMA	BREAST:LEFT	PR:NEGATIVE ER:NEGATIVE
TR_0003005	BLACK (AMERICA) AFRICAN AMERICAN	73*	FEMALE	10760B	BREAST - OVERLAP LES DUCT CARCINOMA 01/04/2000@AGE 72 LUNG - UPPER LOBE CARCINOMA/EPITHELIAL	LUMPECTOMY OR EXCISIONAL BIOPSY : 01/04/2000 RADIATION : 03/29/2000	MALIG-PRIMARY (INVA DIFFERENTIATED :DU MALIG-PRIMARY (INVA DIFFERENTIATED :CA TUMOR	SIVE):POORLY ICT CARCINOMA SIVE):POORLY RCINOMA/EPITHELIAL	BREAST:LEFT	PR:NEGATIVE ER:NEGATIVE
Support Data Mining (cross-sectional)



Records Foun	nd 43 D	isplay	20	Rec	ords Per Page	<prev 1="" next="" page=""></prev>	Go To Page 1 Fie	elds Export		
Pub ID	Ethnicity	Age	Gender	IRB	Clinical Diagnosis	Treatment	Pathology	Radiology	Specimen	Biomarker
TR_0001972	WHITE	82	FEMALE	10760B	BREAST - UPPER- OUTER DUCT CARCINOMA 03/20/2006@AGE 79 BREAST - OVERLAP LES SOLID CARCINOMA 03/20/2006@AGE 79	LUMPECTOMY OR EXCISIONAL BIOPSY : 06/09/2006 LUMPECTOMY OR EXCISIONAL BIOPSY : 06/09/2006	MALIG-PRIMARY (INVASIVE):WELL DIFFERENTIATED :DUCT CARCINOMA CA IN SITU:MODERATELY DIFFERENTIATED:SOLID CARCINOMA	MRI : 05/01/2006 :MASS : 0VAL MRI : 05/01/2006 :NON- MASS : LINEAR MAMMOGRAM :	BREAST:LEFT BREAST:RIGHT	PR:POSITIVE ER:POSITIVE PR:POSITIVE ER:POSITIVE
TR_0002284	BLACK (AMERICA) -AFRICAN AMERICAN	64 	FEMALE	10760B	BREAST - NOS DUCT CARCINOMA 09/05/2001@AGE 57	RADIATION : 11/05/2001 LUMPECTOMY OR EXCISIONAL BIOPSY : 10/05/2001	CAIN SITU:WELL DIFFERENTIATED :DUCT CARCINOMA	MRI : 03/22/2004 :MASS : 0VAL MAMMOGRAM : 03/16/2004 :	BREAST:LEFT	PR:POSITIVE ER:POSITIVE
TR_0002344	UKNOWN	51	FEMALE	10760B	BREAST - OVERLAP LES CRIBRIFORM CARCINOMA 01/24/2005@AGE 47 BREAST - UPPER- OUTER LOBULAR CARCINOMA 02/18/2005@AGE 47	MODIFIED RADICAL MASTECTOMY WITH REMOVAL OF UNINVOLVED CONTRALATERAL BREAST : 02/18/2005 MODIFIED RADICAL MASTECTOMY WITH REMOVAL OF UNINVOLVED CONTRALATERAL BREAST : 02/18/2005	CA IN SITU:MODERATELY DIFFERENTIATED:CRIBRIFORM CARCINOMA MALIG-PRIMARY (INVASIVE):MODERATELY DIFFERENTIATED:LOBULAR CARCINOMA	MRI : 02/07/2005 :MASS : ROUND MRI : 02/07/2005 :NON- MASS : DIFFUSE MAMMOGRAM : 02/07/2005 :	BREAST:RIGHT BREAST:LEFT	PR:POSITIVE ER:POSITIVE PR:POSITIVE ER:POSITIVE
TR_0002351	WHITE	57	FEMALE	10760B	BREAST - UPPER- OUTER LOBULAR CARCINOMA 03/14/2003@AGE 51	LUMPECTOMY OR EXCISIONAL BIOPSY : 03/14/2003	CA IN SITU:NOT DETERMINED; N/A; UNK :LOBULAR CARCINOMA	MRI : 04/14/2003 :NON- MASS : FOCAL MRI : 04/25/2005 : MAMMOGRAM : 04/10/2003 :	BREAST:RIGHT	PR:POSITIVE ER:POSITIVE
TR_0002360	WHITE	91	FEMALE	10760B	BREAST - UPPER- INNER	TOTAL (SIMPLE) MASTECTOMY : 12/22/2003	CA IN SITU:POORLY DIFFERENTIATED :COMEDOCARCINOMA	MRI : 02/09/2004 :NON-	BREAST:LEFT	PR:NEGATIVE ER:POSITIVE

Support Data Mining (zone in study)

currently underdevelopment



Locate Sample for Basic Research

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ib ID	Ethnicity	Age	Gender	IRB	Clinical Diagnosis	Treatment	Pathology	Specimen	Sample
_0000685	BLACK (AFRICA) BINI	209	FEMALE	13304B	BREAST - NOS BREAST CANCER	DUODENECT WITH PARTIAL GASTRECTOMY (WHIPPLE) : 02	& UNKNOWN)2/28/2007	BLOOD:	DNA:977684
_0001980	BLACK (AMERICA)- -AFRICAN AMERICAN	76	FEMALE	8962	BREAST - UPPER-INNER LOBULAR CARCINOMA 10/04/2006@AGE 74	TOTAL (SIMPLE) MASTECTOMY 11/01/2006	Y : MALIG-PRIMARY (INVAS DIFFERENTIATED :LOB CARCINOMA	SIVE):WELL BLOOD: BULAR BREAST:RIGHT	CONCEPTUAL_TISSUE:TUMOR:2 SERUM:6016779 DNA:6027910 SERUM:6016781 SERUM:6016780 DNA:6022715
_0002002	WHITE	62	FEMALE	8962 10760B	BREAST - UPPER-OUTER LOBULAR CARCINOMA 09/21/2006@AGE 60	RADIATION : 12/05/2006 LUMPECTOMY OR EXCISIONAL BIOPSY : 10/20/2006	MALIG-PRIMARY (INVAS L DIFFERENTIATED :LOB CARCINOMA	SIVE):WELL BLOOD:NOS SULAR BLOOD:NOS BREAST:LEFT	EBV CELLS:7042801 SERUM:7043860 CONCEPTUAL_TISSUE:TUMOR:2 EBV CELLS:7042804 DNA:7068300 EBV CELLS:7042802 EBV CELLS:7042803 SERUM:7043859 EBV CELLS:7042805 SERUM:7043858
_0002018	BLACK (AMERICA)- -AFRICAN AMERICAN	71	FEMALE	10760B	BREAST - NOS DUCT CARCINOMA 09/16/2003@AGE 66	MODIFIED RADICAL MASTECT 10/15/2003 ADRIAMYCIN (DOXORUBICIN)::CYCLOPHOS (CYTOXAN) : 11/21/2003	OMY : MALIG-PRIMARY (INVAS DIFFERENTIATED :DUC CARCINOMA SPHAMIDE	SIVE):POORLY BREAST:LEFT CT BREAST:RIGHT	CONCEPTUAL_TISSUE:TUMOR:1 TISSUE FROZEN:Diseased:80164 TISSUE FROZEN:Normal:8016421 TISSUE PARAFFIN:Diseased:8011 TISSUE FROZEN:Normal:8016421 TISSUE FROZEN:Diseased:80164 TISSUE PARAFFIN:Normal:80164
_0002029	BLACK (AMERICA)- -AFRICAN AMERICAN	43	FEMALE	13344A	BREAST - NOS DUCT CARCINOMA 10/16/2006@AGE 40	ADRIAMYCIN (DOXORUBICIN)::CYCLOPHOS (CYTOXAN)::PACLITAXEL (TAXC 11/16/2006 LUMPECTOMY OR EXCISIONAL BIOPSY : 03/16/2007 RADIATION : 04/26/2007	MALIG-PRIMARY 3PHAMIDE (INVASIVE):MODERATEI OL): DIFFERENTIATED:DUC CARCINOMA IL	OVARY:LEFT LY BREAST: T FALLOPIAN TUBE:LEFT FALLOPIAN TUBE:RIGHT BLOOD: BREAST:RIGHT	SERUM:6028866 TISSUE PARAFFIN:Normal:80188 TISSUE PARAFFIN:Normal:80188 TISSUE FROZEN:Normal:801881 TISSUE FROZEN:TUMOR:600606 TISSUE FROZEN:Normal:801881- TISSUE FROZEN:NORMAL:60060 TISSUE FROZEN:Normal:801881!

Support Full Scale Curation (editing survey questionnaire)

Add Question-Category	
Branch Name	Reproductive System (Section G-J) 💌
Branch Name	Reproductive System (Section G-J)
Category	Pregnancy history (Section H)
	Update Delete Cancel

Updat	te Order Add Item			
Order	Item Name	Description	Unit of Measurement	Options
1	ever_pregnant	H1. Have you ever been pregnant?		Yes; No; Don't know
2	num_pregnan	H2. How many pregnancies have you had	Pregnancies	
3	num_livebirths	H3. How many live births have you had	Live births	
4	age_first_birth	H4. Age you had your first live birth	Years	
5	age_last_birth	H5. Age you had your last live birth	Years	
6	ever_breastfeed_one_month	H6. Ever breast-feed for at least one month		Yes; No; Don't know
7	pregancy_id (Enter pregnancy number)			
8	preg_outcome	H7. What was the outcome of this pregnancy?		Currently pregnant; Single live birth; Multiple birth; Still birth; Miscarriage; Tubal or ectopic pregnancy; Induced abortion; Don't know
9	date_baby_born	H8. When was your baby born for this pregnancy? (format mm/YYYY)	mm/YYYY	
10	preg_duration	H9. How long did this pregnancy last?	Months	<=3; 4-6; >=7; Don't know
11	num_boys_preg	H.10 How many boys did you have in this pregnancy?		
12	num_girls_preg	H10. How many girls did you have in this pregnancy?		
13	ever_breastfeed	H.11 Did you breast feed this baby?		Yes; No; Don't know
14	mths_breastfeed	H12. How many months did you breastfeed this baby?	Months	<1; 1-5; 6-11; 12-24; >24; Don't know

Multiple Level of Data Privacy Control (per IRB, project, and user role)

Breast Cancer SF	ORE
User Project Departme	ent IRB List Project List Logs Emails Default Consent
Update Profile	
User Name	·~@medicine.bsd.uchicago.edu
First Name	C
Last Name	C · · ·
Phone Number	773-834-1945
Project Group	Center for Clinical Cancer Genetics 🛩
User Role	user
Account Status	group admin curator
Pl's E-mail	user
Application Type	administrator
Department	Medicine
Are you a PI?	Yes No
Secret Question	What is your father's middle name?
	Update
Change to Temporary F	Password
Password	
Confirm Password	
	Assign
Profile Deletion	
	Delete User

Acknowledgements

Computation Institute

Xiaoming Wang Lili Liu Greg Cross Ian Foster

Dept. of Radiology

Paul Chang Gilliam Newstead Sunny Arkani Janson

Center for Cancer Genetics

Funmi Olopade Kisha Hope Shelly Porcellino Jim Fackenthal Others

Other

Breast Cancer SPORE Cancer Registry (Cassie Simon) Specimen Bank (Leslie Martin)

Department of Medicine

T.R.I.D.O.M.

Total patients asked:	6943
Patients consented:	5370
Patients refused :	1499
Percent consented :	77.34%
Samples received :	3446

Department	Consented	Samples Received
CARDIOLOGY	155	118
DERMATOLOGY	2	1
EMERGENCY MEDICINE	2	2
ENDOCRINOLOGY	214	166
GASTROENTEROLOGY	1312	932
GENERAL MEDICINE	800	527
HEMATOLOGY/ONCOLOGY	310	225
HOSPITALIST MEDICINE	5	5
INPATIENT	168	64
NEPHROLOGY	52	48
NEUROPHYSIOLOGY	10	4
PULMONARY/CRITICAL CARE	1335	521
RHEUMATOLOGY	975	842
Unknown	30	17

Be a Part of Something Bigger



TRANSLATIONAL RESEARCH INITIATIVE AT THE DEPARTMENT OF MEDICINE





Genomic research – The future for everyday healthcare

What is genomic research? It is the study of uniquely inherited information, which helps to determine the make up and functioning of our bodies. Through this research project, we will link genomic information with various diseases and how they are treated in the clinics. We hope that our efforts will generate new patient care technologies and therapies.



White:	1,839	53%
Black or African American:	1,319	38%
Unknown or Not Reported:	170	5%
Asian:	65	2%
More Than One Race:	38	1%
American Indian/Alaska Native :	10	0%
Native Hawaiian or Other Pacific Islander:	4	0%

Tridom



The Chicago BioMedicine Information Services (CBIS) Service Oriented Architecture (SOA)

Translational Research Support Services:

Patient Data Services (Laboratory, Pathology, Radiology) Electronic Honest Broker DICOM iBroker

Paul J. Chang, M.D., FSIIM Professor & Vice-Chairman, Radiology Informatics Medical Director, Pathology Informatics University of Chicago School of Medicine

> Medical Director, Enterprise Imaging Architect, CBIS SOA Initiative University of Chicago Medical Center

Typical Legacy-based IT Environment: "Using humans to integrate workflow"



Integration using a "Single Vendor Solution"



Using "Edge" Protocols (DICOM & HL7) to Orchestrate Integrated Workflow: IHE PACS – RIS Integration Workflow Model



Modality Worklist Modality Performed Procedure Storage Commit Integration Engine Approach Using Edge Protocols (HL-7, DICOM) to Transfer State Example: PACS – RIS Integration Workflow Model







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NJ community grieves for Chile tourists Seattle Post Intelligencer - all 800 related »	Vive La Wireless	Gmail Inbox Hide preview	edit 🗙
Update 1: Family Members of ETA's Victims Hopeful Forbes - all 799 related »	Fortune.comedit IXBiggest Biz Stories of 2005Why Google Will Falter in 2006	nadiya@dreamtown.com - I NADIYA - NADIYA YAKOV nadiya@dreamtown.com - I	4:00am Mar 22 Mar 21
CNN edit 🛛	Two Losers and a Winner	nadiya@dreamtown.com - I Tickets - Schedule Change	Mar 12 Mar 11
Blaze breaks out aboard US cruise ship, 1 dead Tour bus plummets into canyon, kills	Slashdot edit X	Weather Pittsburgh, PA	edit 🛛
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Enterprise Integration Model: Towards a Service Oriented Architecture



Middle ("Business Logic") Layer (Agents, ORBs, Web Services, etc.)



Patient Data Services (Laboratory, Pathology, Radiology)

Electronic Honest Broker

Process Overview



High-Level Logical Architecture



Technologies Overview

- Java SE6, EE5, JAX-WS 2.1
- SOAP Service and Client JAX-WS generated from WSDL
- Database agnostic; Oracle implementation
- LDAP agnostic; AD implementation
- Servlet Container agnostic; Tomcat implementation

Login Page



Electronic Honest Broker

Log in

Username: Password:	Please Log In	Welcome to the University of Chicago Medical Center Electronic Honest Broker application. The purpose of this application is to provide medical researchers access to anonymous, clinical data. The Electronic Honest Broker does not provide any actual clinical data; rather, it serves as a tool for use by human
	Log in	honest brokers whose job it is to make sure research is conducted in a manner consistent with HIPAA regulations.
		Human honest brokers and administrators from the Office of Clinical Research should be the only users accessing this application. Use of this application constitutes agreement on your part to adhere to the University of Chicago Medical Center Security Policy and Internet Use and Privacy Policy. (Please click the links below to view the policies.)

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Electronic Honest Broker

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Active	Studies		
Study	# Title	Expiration	

Administrative Functions

- Enter a New Study
- User Administration
- Role Administration
- View Logs

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Electronic Honest Broker: Add Study



Electronic Honest Broker

Home | Log Out kleiding

Study Information	
Study Number:	11111
Title:	New Study
Description:	s a new study for demo purposes.
IRB Number:	12-34567
Expiration (YYYY-MM-DD):	2009-12-21
Principal Investigator:	Duke
Create Study	

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Electronic Honest Broker: New Study Added



Electronic Honest Broker

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New study added successfully.

Active St						
Study #	Title	Expiration	Expiration			
11111	New Study	2009-12-21	View			

Administrative Functions

- Enter a New Study
- User Administration
- Role Administration
- View Logs

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Electronic Honest Broker: Adding a New Patient



Electronic Honest Broker

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	Allee		Study Number:	11111
Anonymous ID •	Allas	Phi	Title:	New Study
			Description:	This is a new study for demo purposes.
			IRB Number:	12-34567
			Expiration:	2009-12-21
			Principal Investigator:	Duke
			Edit Study	
			Add a Study Patient	
			Patient MRN:	12345
			Alias (optional):	Patient #1
				Add
			Study Users	
			Username	Role
			Assign Study Role to	User
			Username	Role
				Honest Broker 🔹 🔿

Electronic Honest Broker: Verify Patient

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	MEDICAL CENTER
	MEDICAL CENTER

Electronic Honest Broker

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Study Patients No patients in study. Anonymous ID Alias	PHI	Study Information Study Number: Title:	11111 New Study
	Patient Information MRN First Name Last Name Is this the correct patie	nt? es, Continue	a new study for demo es. i67 2-21 i4567 ent #1 id
		Username	Role
		Assign Study Rol	e to User
		Username	Role Honest Broker 🛟 (Assign)

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Electronic Honest Broker

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Study Patients Patients 1-10 of 10 Anonymous ID 11524183 17688601 29271315 31740714 38268833 50370420 52245218 52488833 69148400 90812355	Alias PH Pa Patient Information Pa MRN Pa Anonymous ID Pa Alias Pa First Name Pa Last Name Pa Date of Birth Pa Close Patient Information	Study Information Study Number: Title:	11111 New Study a new study for demo es. i67 2-21
		Username cgoodman Assign Study Rol Username	Role Honest Broker Remove

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Honest Broker Study View



Electronic Honest Broker

Home | Log Out kleiding

Study Patients

Anonymous ID 💌	Alias		PHI
11524183	Patient #10	Change	View
17688601	Patient #3	Change	View
29271315	Patient #5	Change	View
31740714	Patient #1	Change	View
38268833	Patient #9	Change	View
50370420	Patient #2	Change	View
52245218	Patient #8	Change	View
52488833	Patient #7	Change	View
69148400	Patient #4	Change	View
90812355	Patient #6	Change	View

Study Information							
Study Number:	11111						
Title:	New Study						
Description:	This is a new study for demo purposes.						
IRB Number:	12-34567						
Expiration:	2009-12-21						
Principal Investigator:	Duke						
Edit Study							

Add a Study Patient		
Patient MRN:		
Alias (optional):		
	Add	

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- A portal and related services that provide end-users simple and powerful HIPAA compliant access to clinical image datasets
- Leverages Electronic Honest Broker



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ID	Date	Name	Value	Low	High	Norm	Unit
966	2/4/2009	FERI-FERRITIN	8	10	220	10-220	ng/mL
958	2/4/2009	FE-IRON, SERUM	30	40	160	40-160	mcg/dL
2383	2/4/2009	TIBC-TOT, IRON-BINDING CAP.	466	230	430	230-430	mcg/dL
1888	2/4/2009	PSAT-% SATURATION	6.4	14	50	14-50	%
2586	2/4/2009	WBC-WBC	5.2	3.5	11	3.5-11	K/uL
1941	2/4/2009	RBC-RBC	4.48	3.88	5.26	3.88-5.26	M/uL
1200	2/4/2009	HGB-HEMOGLOBIN	12.3	11.5	15.5	11.5-15.5	g/dL
1156	2/4/2009	HCT-HEMATOCRIT	36.6	36	47	36-47	%
1537	2/4/2009	MCV-MCV	81.6	81	99	81-99	fL
1532	2/4/2009	MCH-MCH	27.4	26	33	26-33	pg
1534	2/4/2009	MCHC-MCHC	33.6	32	35	32-35	g/dL
1955	2/4/2009	RDW-RBC DIST WIDTH	17.4			<15.0	%
1830	2/4/2009	PLAT-PLATELET COUNT	211	150	450	150-450	K/uL
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Date	Reports
4/2/2009	 FINAL PATHOLOGIC DIAGNOSIS A. Biopsy, cervix at 6 o' clock: - Benign endocervical mucosa. B. Biopsy, cervix at 12 o'clock: - Cervical mucosa with HPV effect and squamous metaplasia. C. Curettage, endocervix: - Strips of benign endocervical glands. Interpretation performed by the Attending Pathologist and reviewed with the Resident or Fellow. Electronically Signed Out by Clinical History: The patient is a with abnormal Pap smear showing ASC-H. Specimens Received: A: Cx Bx @ 6 o clock B: Cx Bx @ 12 o'clock C: ECC Gross Description: The specimens are received in three containers each labeled with the patient's name and medical record number. A. The first container is additionally identified as, "1 cervix biopsy at 6 o' clock". Received in formalin is one soft, pink-white tissue fragment, measuring 0.4 cm, which are submitted in toto as A1. B. The second container is additionally identified as, "2 cervix biopsy at 12 o' clock". Received in formalin is one soft, pink-white tissue fragment, measuring 0.4 cm, which are submitted in toto as B1. C. The third container is additionally identified as, "2 cervix biopsy at 12 o' clock". Received in formalin is one soft, pink-white tissue fragment, measuring 0.4 cm, which are submitted in toto as B1. C. The third container is additionally identified as, "2 cervix biopsy at 12 o' clock". Received in formalin is one soft, pink-white tissue fragment, measuring 0.4 cm, which are submitted in toto as B1. C. The third container is additionally identified as, "5 cervix biopsy at 12 o' clock". Received in formalin is one soft, pink-white tissue fragment, measuring 0.4 cm, which are submitted in toto as B1. C. The third container is additionally identified as, "5 cervix biopsy at 12 o' clock".

First integrated use of SOA



caBIG

- caBIG uses caGrid as the foundation which uses various components of Globus Toolkit
 - WS-RF Framework
 - Security Framework
 - Index Service
- We lead the Workflow Project of caGrid
- We are part of the Architecture workspace

Acknowledgement: Ravi Madduri

UCCRC-caBIG

- We have a caGrid node up and running on caBIG
- We evaluated caArray and we deployed it in UCCRC
- We are working with Velos in an effort to do CDUS submissions using a caGrid Service

caGrid Environment



*All Services Register with the Index Service

IRIS

Integrated Research Information Services



Current IT Research Environment

- □ ARG Academic and Research Group (PSMTrak,
- □ CI Computational Institute (TRAM)
- $\Box \quad CRC Cancer Research Center (Velos)$
- □ DID Development, Integration & Databases (Clinical SOA)
- □ HSD Health Studies Department (Sandstone)
- □ iBi Institute for BioInformatics
- □ MAD Medicine Application Developers (TRIDOM)
- □ NSIT Networking Services and Information Technologies

Functional Academic & Research Domains



5/26/2009

Chicago Biomedicine Information Services

Current ARG Applications & Systems











Chicago Biomedicine Information Services

Current Databases and Feeds





BSD



ARG Ecosystem



Chicago Biomedicine Information Services

SOA Layers

(Service Consumer) Clinical, Researchers, External Affiliates

(Access Layer) Identity Server, Portal, SOAP Client

(Process Layer) (Business Process, Workflows)







University of Chicago Initiative in Biomedical Informatics



Genome Science and Biomedical Informatics increase the "experimental space" and "clinical space", respectively one can "search" for biomedical solutions



In "genetic" studies we seek to correlate heritable changes in DNA to Disease Phenotype. In "genomic" studies we seek to correlate DNA changes with other changes in "molecular state" to predict Disease Phenotype. In **Biomedical Informatics** we seek to characterize "disease" by its component parts with the goal of predicting disease susceptibility by correlation to individual differences in "molecular states".



As of Jan 2008, *GenBank* houses just under 1 Terabyte (10¹²) of DNA sequence information. Though huge by pre-genome sequencing standards, the data storage challenge is really a consequence of the new "experimental space" that genome science and genome technology enable.

Shorter than Moore's law (computer power doubling every 20 months!)

Automated literature mining to generate a Molecular Interaction Network



Discovering Pathogenic Pathways: why do we need to integrate information?



All currently stored in iBi repository.

New "experimental space"

PhenoGO : A Resource for the Multiscale Integration of Clinical and Biological Data



Lee T. Sam, Eneida A. Mendonça, Carol Friedman,

Yves A. Lussier, M.D. (Corresponding Author) Director, Dept. of Medicine Center for Biomedical Informatics Associate Director for Informatics, Cancer Research Center Co-Director, Clinical Translational Science Award (CTSA) Informatics Core Associate Professor of Medicine, Biological Science Division

Judith Blake, Jackson Lab

The PhenoGO Encoding Pipeline



Lussier Y, Borlawsky T, Rappaport D, Liu Y, Friedman C. PhenoGO: assigning phenotypic context to gene ontology annotations with natural language processing. *Pac Symp Biocomput*. 2006;:64-75.

Diseases and Disorders

- 78,947 total annotations associated with diseases and disorders
 - 69,899 Human and mouse annotations
 - 3,209 distinct
 disease and
 disorders of 6,084
 distinct contexts



Web Query: Basic

 Runs a query analogous to an SQL OR for the search terms

🟉 PhenoGO DB Query - Internet E	xplorer provided by Dell			
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🔶 💠 🌈 PhenoGO DB Quer	у		🏠 🔹 🗟 👻 🗟	Page ▼ () Tools ▼ [≫]
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About: (Databa	se updated on November 1st 2008)			E
PhenoGO is : existing asso (GOA). Conte Unified Medic taxonomy, an Note: Some u	a multiorganism database that provides pheno iciations between gene products and Gene Or xt to identifiers are mapped to general biologic cal Language System (UMLS), species from Ta d some specialized ontologies such as Mamr users of Internet Explorer 7.0 may enounter sp a speciety	otypic context, such as the cell ty itology (GO) terms as specified cal ontologies, including the Cel axonomy of the National Center malian Phenotype Ontology (MP) oradic issues with the website. 1	ype, disease, and tissue and organ to in the Gene Ontology Annotations II Ontology (CO), phenotypes from the for Biotechnology Information (NCBI)) and Mouse Anatomy (MA). We recommend using an alternative	
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Web Query: Advanced

Runs a query analogous to an SQL AND for the search terms

🟉 PhenoGO: Advanced Query - Internet Explorer pr	ovided by Dell						
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Conclusions

- Substantial additions to a high-quality, wide-raging gene-GO-phenotype resource drawn from the biomedical literature and existing knowledge bases
- It's a fantastic resource and I urge everyone to make use of it
- http://www.phenogo.org

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 - Jianrong Li
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 - NLM, R01 LM007659-01 (CF)
 - NLM 1U54 CA121852-01A1 National Center: Multi-Scale Study of Cellular Networks

Discovery of Protein Interaction Networks Shared by Diseases

Lee Sam, Yang Liu, Jianrong Li, Carol Friedman, Yves A. Lussier



PhenoGO.org

Lussier Y, Borlawsky T, Rappaport D, Liu Y, Friedman C. Pac Symp Biocomput. 2006;:64-75.

- Database of Phenotypic context for GO annotations
 - Gene-phenotype-GO
 - NLP (BioMedLEE) and computational terminologyderived
- 3,102 distinct diseases and phenotypes (# human)
- 32,911 distinct proteins (7,016 human)
- 532,407 total records (# human)
- Precision: 85% [n=120]
- Recall: 76% [n=120]

Overall process

o Protein-Protein Interaction network

- Reactome
- o Phenotype-genotype association





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- Joanna Amburger, OMIM

Grants

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- NLM, R01 LM007659-01 (CF)
- NLM 1U54 CA121852-01A1 National Center: Multi-Scale Study of Cellular Networks (YL/Phenotypes & CF/NLP Team Leaders)

Probing genetic overlap among complex human phenotypes

Andrey Rzhetsky*+*, David Wajngurt*, Naeun Park*, and Tian Zheng⁵

*Department of Biomedical Informatics, Center for Computational Biology and Bioinformatics and Joint Centers for Systems Biology, and *Judith P. Sulzberger, M.D., Columbia Genome Center, Columbia University, New York, NY 10032; and *Department of Statistics, Columbia University, New York, NY 10027

11694-11699 | PNAS | July 10, 2007 | vol. 104 | no. 28





David Wajngurt

Naeun Park Tian Zheng

AR

SANG

Data

$\sum_{=1.5 \times 10^6}$ patient records



27|U|M^079.9:8|493.90:10 ...

Data: ICD9 codes in Columbia University clinical database
Results: Autism



Results: Schizophrenia











CIQR ("Seeker"): Context-Initiated Question and Response

Eneida Mendonca, MD, PhD

History

- □ Linking medical records to information resources
- □ Making retrieval specific to a particular patients
- Using statistical and semantic models to identify relevant information in patient's records
- Re-ranking retrieval of scientific literature based on the patient's map
- Problems: not a good time for retrieval, questions not always related to the part of the record the physician is looking at.

CIQR: Objectives

- Capture information needs when they occur, without disrupting workflow
 - Deploy mobile devices using data and voice input
- □ Translate high-level information needs into search strategies adapted to user needs capabilities of resources
 - Develop models of search strategies using human search expertise (reference librarians)
 - Study how librarians classify, clarify and refine questions
 - Automate using speech processing and natural language parsing
- Deliver materials relevant to information needs in an accessible format and in a timely manner
 - Organize materials retrieved by librarians and transmit them to clinicians for display on mobile devices

Where we are going?

- □ Improved training
- □ Testing new devices
- □ Text input vs. voice input
- □ Full text analysis
- Delivery of information





From Bench to Bedside: Lessons Learned in the University of Chicago CTSA

Re-Engineering Translational Research at The University of Chicago Clinical and Translational Science Award U54 RR023560

> Julian Solway, MD University of Chicago

s sans hangle

a transfer

< 26/16-

South Side of Chicago



#	Community
1	McKinley Park
2	Bridgeport
3	Armour Square
4	Douglas
5	Oakland
6	New City
7	Fuller Park
8	Grand Boulevard
9	Kenwood
10	West Englewood
11	Englewood
12	Washington Park
13	Hyde Park
14	Woodlawn
15	Greater Grand Crossing
16	South Shore
17	Auburn Gresham
18	Chatham
19	Avalon Park
20	South Chicago
21	Burnside
22	Calumet Heights
23	Beverly
24	Washington Heights
25	Roseland
26	Pullman
27	South Deering
28	East Side
29	Morgan Park
30	West Pullman
31	Riverdale
32	Hegewisch

The health status of the 1.1 M residents of the UCMC PSA is extremely poor, as reflected in extraordinarily high rates of common complex diseases and of infant mortality. Furthermore, 10-15% of UHI adult residents are disabled.

Description	PSA	Illinois	PSA / IL
HEART FAILURE & SHOCK	10.5	5.3	199%
DIABETES AGE >35	3.4	1.2	285%
RENAL FAILURE	3.1	1.5	213%
BRONCHITIS & ASTHMA	1.6	0.7	219%
HYPERTENSION	1.6	0.6	262%
All Adult Med/Surg DRGs	169.0	125.2	135%

It is in this context that the University of Chicago CTSA program seeks to translate research discoveries into real, effective therapies.

		% Infant		
Community	Area	Mortality		
Grand Boulevard	PSA	3.1%		
Riverdale	PSA	2.2%		
Clearing	Other	1.9%		
West Englewood	PSA	1.9%		
Douglas	PSA	1.9%		
Washington Park	PSA	1.8%		
Chatham	PSA	1.8%		
West Garfield Park	Other	1.7%		
North Lawndale	Other	1.6%		
Humboldt Park	Other	1.6%		
East Garfield Park	Other	1.6%		
Hyde Park	PSA	1.6%		
Englewood	PSA	1.6%		
Austin	Other	1.5%		
Pullman	PSA	1.5%		
Roseland	PSA	1.4%		
Ashburn	Other	1.4%		
Loop	Other	1.4%		
West Pullman	PSA	1.3%		
Kenwood	PSA	1.3%		
South Shore	PSA	1.2%		
Morgan Park	1.2%			
Chicago Average	0.9%			
Illinois Average	0.73%			

A comprehensive approach to the challenges:

- Institute for Translational Medicine (ITM) a new University structure to collect, integrate, and disseminate the intellectual, organizational, and resource infrastructure needed to reduce barriers to translational research interactions among UC investigators. The ITM:
 - provides <u>new modes of "research navigation" assistance</u> to help UC and Community faculty and trainees identify and contact collaborators
 - will <u>incentivize multidisciplinary collaborations</u> with pilot and collaborative seed grant funding
 - <u>support the actual research</u> with a wide spectrum of core and intellectual capabilities
- 2) <u>Urban Health Initiative (UHI)</u> principal goals are:
 - fully engage our Community in <u>developing clinical and community</u> <u>research agendas</u>
 - involve Community organizations and practices in the <u>conduct</u>, <u>interpretation and dissemination of that research</u>
 - <u>accelerate the translation of health knowledge</u> into and out of our Community
 - most importantly, <u>reduce the marked health disparities</u> in our Community through a <u>combination of service, education, and research initiatives</u>.
 CTSA Community Translational Science Cluster <u>is</u> the principal research infrastructure component of the UHI.

Three 3 bold steps that will overcome these obstacles.

3) <u>Training in Clinical and Translational Science</u>

- Greatly expanded our training for careers in clinical and translational research, now providing research training and career development opportunities for physician and non-physician scientists, allied health professionals, undergraduates, and high school students, both at UC and in our Community.
- Previous barrier to such research training goal as one of "activation energy", in that we have had the essential capability for such training and career development, but failed to realize our full potential due to inadequate mentor incentives, inadequate course offerings, and inadequate recognition of the need for comprehensive training for careers throughout the translational research spectrum.
- <u>ITM Training Cluster</u> broad research training/career development mission, enhanced faculty/mentor incentives, more clearly defined training pathways, and wide buy-in throughout the CTSA community, operating across divisional, departmental, professional school, and university lines to bring together CTSA faculty and students from UC, our allied institutions, and the Community.
- <u>Committee on Clinical and Translational Science</u> new academic entity to advance multidisciplinary PhD/MS training in clinical and translational research.

The University of Chicago CTSA



The University of Chicago ITM



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