## **INSIDE Dyslipidemia Management**

INspiring System Improvement with Data-Driven Excellence

ers



## **INSIDE Dyslipidemia Management** Program Goals

Benchmark current lipid management and goal attainment for high-risk CV patients

Identify and prioritize specific clinical process/management challenges that affect goal attainment

Systematically make clinical changes to improve goal attainment processes and outcomes

Measure the results and share best practices via educational workshops, online CME and publications

## **Clinical Sites & QI Team Leaders**

University of Chicago Lipid Clinic	•Michael H. Davidson, MD (PI) •Karen Wickens, RN
University of Chicago Section of Cardiology	•Amita Singh, MD •Luke Laffin, MD •Lane Benes, MD
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NorthShore Department of Medicine	<ul> <li>Divisions of Cardiology and Endocrinology</li> <li>David Davidson, MD (PI)</li> </ul>
ADA Quality Improvement Services	•Roy Furman, MD, PhD •Elise Furman, RN, MEd, MBA

#### **INSIDE Program Features**





## **Program Design at a Glance**

INSIDE Dyslipidemia Management is based on ADA's successful Diabetes INSIDE program that has improved diabetes care processes at health systems across the US since 2012



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# Quality Improvement Arm at University of Chicago

# **Goals of Quality Improvement in Population Health**

# Reducing clinical variation

- In processes over time
- In <u>outcomes</u> between rational subgroups

Time/groups

# Achieving clinical guideline standards

- For <u>all patients</u>
- <u>Safely and appropriately</u>

Standards

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#### **Baseline Data – Demographics**

Total N=20,871; Mean Age=64		Ν	%
Gender			
	Male	10,658	51.1%
	Female	10,213	48.9%
Race			
	African American	9,978	46.9%
	White	9,750	46.7%
	Asian/Mideast Indian	643	3.1%
	Other	419	2.0%
	American Indian or Alaskan Native	81	0.4%
CVD/C\	/ Risk Diagnoses		
	Hypertension	10,781	49.0%
	Dyslipidemia	7,632	34.7%
	Coronary Artery Disease	6,681	30.4%
	Diabetes/prediabetes	1,658	7.5%
	Peripheral Vascular Disease	955	4.3%

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Patient selection is visit-level data for all adults following ICD-9 or ICD-10 codes for major cardiovascular disease, dyslipidemia, diabetes, hypertension, or smoking history

Adults	18-89 yo	
ICD 9 code	Condition	ICD 10 code
250	diabetes mellitus	E10-E14
272	lipid metabolism disorders	E78
305.1	nondependent tobacco use disorder	F17
401-405	hypertensive disease	110-115
410-414	ischemic heart disease	120-125
415-417	diseases of pulmonary circulation	126-128
424	other diseases of endocardium	134-136
425	cardiomyopathy	142
426	conduction disorders	144
427.5	cardiac arrest	146
429	ill-defined complications of heart disease	147-151
430-438	cerebrovascular disease	161-167, 169, G45
440-448	diseases of arteries, arterioles, and capillaries	170-177
451	phlebitis and thrombophlebitis	180
V12.5x	personal history of unspecified circulatory disease	Z86.7
V15.1	personal history of surgery to heart and great vesse	els
V15.82	history of tobacco use	Z87.891
V17.3	family history of ischemic heart disease	Z82.49
V17.4x	family history of other cardiovascular diseases	Z82.41, Z82.49
V18.0	family history of diabetes mellitus	Z83.3

Joshy G, Korda R, Abhayaratna W, Soga K, Banks E. Categorising major cardiovascular disease hospitalisations from routinely collected data. *Public Health Research & Practice*. 2015;25(3). doi:10.17061/phrp2531532





### Visit Distribution by Location, Department, Provider



#### 608,517 visits for 21,986 patients with 177 providers over 5.2 years; mean age=64



Treemap displaying hierarchical (tree-structured) patient data as a set of nested rectangles. Each rectangle has an area proportional to the specified dimension of the data.



#### **Distribution of ICD 9/10 Diagnoses**



#### 608,517 visits for 21,986 patients with 177 providers over 5.2 years; mean age=64

		/ 1		•		-	•	
Other Forms Of Hea	art Disease	Hypertensive Disease	Other Metabolic And Immunity Disorders	Other Form	s Of Heart D	isease	Hypertensive Diseases	Metabolic Disorders
				Heart failure	Atrial fibr	illation and flutter		
Cardiac dysrhythmias Hea	art failure	_	Endocrine, Nutritional					
Diseases Of	The Circula	atory System	And Metabolic Diseases,	Other Diseas	ses of the	circulato	ry system	Endocrine, nutritional
Ill-defined			And Immunity Disorders Disorders of	armythmas	Cardiomyopathy	Other		lipoprotein metabolism
descriptions andOther	Cardiomyopathy		lipoid metabolism	Complications	A	Atrioventricular diseases of pericardium		And other lipidemias
complications of diseases of	Conduction	Essential hypertension	Disorders of fluid, electrolyte, and	and ill-defined	Barayyamal	oundle-branch block	Essential (primary)	Diabetes Mellitus
heart disease  endocardium	disorders	Hypertensive heart disease Secondary hypertension	Overweight, Gout	heart disease	tachycardia	Other conduction arrest		
Ischemic Heart Disc	ease	Arteries, Disease	hyperalimentation	Ischemic H	eart Disease	S Diseases	Of Cerebrovascular Diseases	-
	0 marine	Arterioles, And-cerebral	Diseases Of Other			Arteries, And Cap	illaries infarction	Type 2 diabetes
	pectoris	Capillaries	Endocrine Glands			Angina	Occls and stenosis of precerb art, not rstt in cereb infrc	mellitus
		And dissection And Lymphatics, And				and dissec	ION Atherosclerosis Rheumatic Venous	Obesity And Of
		Diseases Of Circulatory System				STEMI Pulmonary	leart Disease Diseases and	Hyperalimentation Thyroid
Other forms of chronic		Pulmonary Chronic Rheumatic	Diabetes mellitus	Observation in a la servation la		mocard Pulmonary	Circulation triouspid thrombos	Overweight
Nephritis, Other Diseases	Neurotic Disorders, Personality	Diseases Of The	Secondary diabetes melitus	Acute Kidney	Teart disease Diseases Diseases Diseases Diseases Diseases Diseases	Oth and uns	Other Mental And Disea	es of the blood and Hepatitic
Nephrotic Urinary Disease	es Of Disorders, And Other Nonpsychotic Mental	Diseases Of The Respiratory	eoplasms Of The Injury Blood And And	Failure And	Senital Paroxysmal	Diseases of the	Diseases Disorders Due blood- certain	forming organs and disorders involving Neoplasms
Genitourinary System System	And Disorders	System And Connective	Organs	denitourinary syst	em nervous syste	system and	of the Substance Use Dis	seases of
disease (CKD)	rgans	Tissue Diseases Of The Digestive System A	nomalies And Parasitic	Bisease	disorders	tissue	system [Affective] Disorders the	digestive
	Psychoses		Diseases	disease (CKD)		Disorders		system
			ICD 9					ICD 10

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ICD 10 INSIDE Dyslipidemia Management | 10



# Cardiovascular Risk Factor Diagnoses (ICD 10)

#### 21,983 patients; mean age=64

#### **Obesity, prediabetes, diabetes underdiagnosed as risks**





#### **Baseline Data - Rx Frequencies**



#### 953,675 prescriptions for 20,230 patients using 1,427 drugs over 5.2 years; mean age=64



Treemap displaying hierarchical (tree-structured) patient data as a set of nested rectangles. Each rectangle has an area proportional to the specified dimension of the data.

American Diabetes Association.

## **Baseline Data – Demographics**



# 953,675 prescriptions for 20,230 patients using 1,427 drugs over 5.2 years; mean age=64 EHR data gaps in structured fields

Category	Patients	With Known Risks	And LDLC
On any Rx	20,219	16,464	8,939
On lipid Rx	15,262	11,169	5,659
On statin Rx	14,096	10,911	5,521
On PCSK9 Rx	112	55	25



## **LDLC Levels vs Rates of Coronary Events**



1. Raymond C, Cho L, Rocco M, Hazen SL. New cholesterol guidelines: Worth the wait? *Cleveland Clinic Journal of Medicine*. 2014;81(1):11-19. doi:<u>10.3949/ccjm.81a.13161</u> 2. Cholesterol Treatment Trialists' (CTT) Collaboration. Efficacy and safety of more intensive lowering of LDL cholesterol: a meta-analysis of data from 170 000 participants in 26 randomised trials. *Lancet*. 2010;376(9753):1670-1681. doi:<u>10.1016/S0140-6736(10)61350-5</u>

2013 ACC/AHA Guideline on the Treatment of Blood Cholesterol to Reduce Atherosclerotic Cardiovascular Risk in Adults

# **Dyslipidemia Guidelines**

2013 ACC/AHA Guideline on the Treatment of Blood Cholesterol to Reduce Atherosclerotic Cardiovascular Risk in Adults



## Summary of Statin Initiation Recommendations to Reduce ASCVD Risk (Revised Figure)



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## Summary of Statin Initiation Recommendations to Reduce ASCVD Risk (Revised Figure)



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# **Clinical Outcomes**

2013 ACC/AHA Guideline on the Treatment of Blood Cholesterol to Reduce Atherosclerotic Cardiovascular Risk in Adults





## LDLC and Statin Prescribing by AHA/ACC Guidelines Only includes patients on lipid-lowering drugs







## LDLC and Statin Prescribing by AHA/ACC Guidelines Patients with any prescriptions







## LDLC and Statin Prescribing by AHA/ACC Guidelines Only includes patients with diabetes



### **Baseline Data – LDLC Distribution by # of Risk Factors** N=12,823 patients





# **Dyslipidemia Guidelines**

AACE 2017 Guidelines for Management of Dyslipidemia



#### Major Atherosclerotic Cardiovascular Disease Risk Factors

Major risk factors	Additional risk factors	Nontraditional risk factors
Advancing age	Obesity, abdominal obesity	Lipoprotein (a)
Total serum cholesterol	Family history of hyperlipidemia	Clotting factors
Non-HDL-C	Small, dense LDL-C	Inflammation markers (hsCRP; Lp- PLA2)
LDL-C	Аро В	Homocysteine levels
Low HDL-C	LDL particle concentration	Apo B4 isoform
Diabetes mellitus	Fasting/post-prandial hypertriglyceridemia	Uric acid
Hypertension	PCOS	TG-rich remnants
Chronic kidney disease	Dyslipidemic triad	
Cigarette smoking		
Family history of ASCVD		

#### Atherosclerotic Cardiovascular Disease Risk Categories and LDL-C Treatment Goals

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Risk category	Risk factors/10-year risk	LDL-C (mg/dL)	Non-HDL-C (mg/dL)	Apo B (mg/dL)	
Extreme risk	<ul> <li>Progressive ASCVD including angina in patients after achieving LCL-C &lt;70 mg/dL</li> <li>Established clinical cardiovascular disease in patients with DM, CKD 3/4, or HeFH</li> <li>History of premature ASCVD (&lt;55 male, &lt;65 female)</li> </ul>	<55	<80	<70	
Very high risk	<ul> <li>Established or recent hospitalization for ACS, coronary, carotid or peripheral vascular disease, 10-year risk &gt;20%</li> <li>Diabetes or CKD 3/4 with 1 or more risk factor(s)</li> <li>HeFH</li> </ul>	<70	<100	<80	
High risk	<ul> <li>≥2 risk factors and 10-year risk 10-20%</li> <li>Diabetes or CKD 3/4 with no other risk factors</li> </ul>	<100	<130	<90	
Moderate risk	• ≤2 risk factors and 10-year risk <10%	<100	<130	<90	
Low risk	O risk factors	<130	<160	NR	

#### American Diabetes Association.

Treatment Goals

Lipid Goals for Patients at Risk for Atherosclerotic Cardiovascular Disease

Lipid parameter	Goal (mg/dL)
ТС	<200
LDL-C	<130 (low risk) <100 (moderate risk) <100 (high risk) <70 (very high risk) <55 (extreme risk)
Non-HDL-C	30 above LDL-C goal; 25 above LDL-C goal (extreme risk patients)
TG	<150
Аро В	<90 (patients at high risk of ASCVD, including those with diabetes) <80 (patients at very high risk with established ASCVD or diabetes plus ≥1 additional risk factor) <70 (patients at extreme risk)

# **Clinical Outcomes**

AACE 2017 Guidelines for Management of Dyslipidemia



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# **Baseline Data – Cholesterol and Triglycerides Stratified**

#### Most recent labs for 12,921 patients grouped by AACE risk category and gender







## **Baseline Data – LDLC Stratified**

#### Most recent labs for 12,750 patients grouped by AACE risk category and gender





## **Baseline Data – Non-HDLC Stratified**

#### N=5,183 patients, mean interval = 2.1 years (6 days to 5.1 years)



## **Baseline Data – Composite AACE Lipid Goals**



#### N=5,183 patients, mean interval = 2.1 years (6 days to 5.1 years)









## **Baseline Data – LDLC Guideline Outcomes Compared**

#### Most recent labs for 12,750 patients grouped by guideline risk category and gender





## **Baseline Data** – **Evolving Individual LDLC Change** N=5,125 patients, mean interval = 2.1 ± 1.3 years (6 days to 5.1 years)





## **Baseline Data – Evolving Individual LDLC Change** Stratified by AACE Risk Categories: Low and Moderate




#### Baseline Data – Evolving Individual LDLC Change Stratified by AACE Risk Categories: High and Very High





# Baseline Data – Evolving Individual LDLC Change

#### Stratified by AACE Risk Categories: Extreme



# **Quality Improvement**

Reducing clinical variation and achieving guideline standards

Standards

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Time/groups

Quality Management in Population Health Planning, assuring, controlling, improving

#### **Clinical variation**

- Are population clinical outcomes stable and predictable?
- Do patient subpopulations have similar outcomes?

#### **Clinical goals**

• Do process and clinical outcomes achieve clinical standards?

#### Clinical change

- Have prior interventions affected outcomes?
- Has patient population changed over time?

Standards

#### **Process Behavior (Control) Charts Illustrated**



A reliable, repeatable process is "predictable" and randomly varies minimally about the mean over time within natural process limits.

# **Statistical Significance with Control Charts**

#### Simple visual alternative to T-tests, p-values, and other analytical tests

#### By Time

- Within control limits
  - Mean and control limits analogous to statistical error of the mean
  - Runs ≥7 points above/below mean statistically significant
  - Patterns and monotonic trends may be statistically significant
  - Otherwise due to random variation
- Outside control limits
  - Any point is statistically significant

#### Rational Subgroup

- Within control limits
  - Groups do not have statistically significant between-group variation
- Outside control limits
  - Group has statistically significant variation

# **Population Lipid Outcomes**

Are population lipid profiles stable over time?

Have prior awareness, education, or quality programs varied lipid outcomes? Does diagnosis affect lipid levels?





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#### How Does Population LDLC Vary Over Time at U of C?







#### **Population LDLC Standard Deviation Consistent Over Time**



12,828 patients with 22,529 LDLC tests over 5.1 years



### Mean LDLC Variation Over Time in ICHD/CVA/TIA



Stage A: LDLC in Ischemic CHD/CVA/TIA (N=8,719) (4,907 patients)



# **Population Lipid Outcomes**

How does medication choice and intensity affect lipid levels?



# Statin Intensity Does Not Affect Population Mean LDL

Stage A: LDLC by Statin Intensity (N=17,197) (8,127 patients)



# LDLC Varies with Selection of Statin and Intensity



Stage A: LDLC by Statin Drug Intensity (N=18,669) (8,127 patients)



Statins used: atorvastatin, fluvastatin, lovastatin, pitavastatin, pravastatin, rosuvastatin, simvastatin

# LDLC Varies with Drug Class Combinations

Stage A: LDLC by Lipid Class Combos (N=13,189) (8,318 patients)



Classes: Statins, PCSK9s, cholesterol inhibitors, PPAR-alpha agonists, nicotinc acids, bile acid sequestrants, omega-3 fatty acids

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### **LDLC Varies with Drug Class**



Stage A: LDLC by Lipid Drug Classes (N=14,330) (8,318 patients)



Lipid Drug Classes

Classes: Statins, PCSK9s, cholesterol inhibitors, PPAR-alpha agonists, nicotinc acids, bile acid sequestrants, omega-3 fatty acids

### LDLC Varies with Drug Combinations

Stage A: LDLC by Lipid Drug Combos (N=13,189) (8,318 patients)





### **LDLC Varies with Drug Treatment**



Stage A: LDLC by Lipid Drugs (N=17,724) (8,318 patients)



# Diabetes

How is diabetes and cardiovascular risk being managed?



#### **Mean HbA1c Variation Over Time**



Stage A: HbA1c (N=16,949) (11,211 patients)



Months

# **Proportion with Poor HbA1c Control on Insulin**



Stage A: Poor Control on Insulin (N=15,725) (11,211 patients)





#### **LDLC Variation Over Time in Diabetes**



Stage A: LDLC in DM (N=2,507) (1,812 patients)



Months

#### HbA1c Varies with Drug Class Combos





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### HbA1c Varies with Drug Classes





DM Drug Classes

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# HbA1c Varies with DM Drug Combinations







### HbA1c Varies with DM Drug Class





Drug Classes

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# **Population Risk Profiles**

How do lipid profiles compare among different risk categories?



# Mean LDLC Stratified by # of Major Risk Factors



#### Age, Cholesterol, non-HDLC, LDLC, HDLC, DM, CKD III/IV, Smoking

Stage A: LDLC by # Major Risk Factors (N=22,529) (12,828 patients)



# Mean LDLC Stratified by AACE ASCVD Risk Category

Stage A: LDLC by AACE Risk Category (N=22,529) (12,828 patients)



AACE Risk Category

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#### Mean LDLC Stratified by ACC ASCVD Risk Category

Stage A: LDLC by ACC Risk Category (N=22,529) (12,828 patients)



#### **LDLC Stratified by Diagnosis**

Stage A: LDLC by Dx Category (N=15,124) (12,493 patients)



# **Population Weight**

How does body mass effect lipid outcomes?



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#### **Population BMI Variation Over Time**

Stage A: BMI (N=75,498) (20,787 patients)



Months



#### LDLC vs BMI

Stage A: LDLC by BMI (N=20,084) (12,010 patients)



BMI

#### **Triglycerides vs BMI**

Stage A: Triglycerides by BMI (N=20,363) (12,114 patients)



BMI

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UCL

### **Population BMI Variation With Age**

Stage A: BMI by Age (N=75,498) (12,010 patients)



Age
#### **Population Triglyceride Variation With Age**

Stage A: Triglycerides by Age (N=22,830) (12,931 patients)



Age

LCL

# LDLC by AACE Risk Categories

Do AACE risk categories have different clinical variation?



### LDLC By Age Compared to AACE Low Risk Goal





### LDLC By Age Compared to AACE Moderate Risk Goal



LDLC By Age Compared to AACE High Risk Goal



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### LDLC By Age Compared to AACE Very High Risk Goal



#### LDLC By Age Compared to AACE Extreme Risk Goal



# **Treatment Disparities**

Do population outcomes differ with race or gender?



#### Mean LDLC Stratified by Race and Gender



Stage A: LDLC by Race/Gender (N=22,529) (12,828 patients)



Race/Gender



#### Mean LDLC Stratified by Age and Gender



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#### Mean HDLC Stratified by Age and Gender



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#### Mean Triglycerides Stratified by Age and Gender





# **Providers and Risk Profiles**

Do providers achieve different outcomes?

Is population risk profile changing over time?

#### **LDLC Stratified by Provider**



Stage A: LDLC by Provider ID (N=22,529) (120 providers managing 12,828 patients)



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### LDLC Stratified by PCE 10-Year Risk Estimates



Stage A: PCE 10-Year Risk (N=13,782) (8,287 patients)



Months

### LDLC Stratified by Number of CV Risk Factors



Stage A: Number of Major CV Risks (N=87,762) (21,950 patients)



Months

### **Potential Quality Issues to Assess and Address**



# Structural

•EHR lab and data gaps for managing and monitoring population health

## Process

Compare and select among current dyslipidemia guidelines
Undercoding for ASCVD risk factors of obesity, prediabetes, diabetes
Better assessment of diabetes population

# Outcome

• Dyslipidemia undertreatment with progressive increase in ASCVD risk

- •Decrease proportion of patients with poor diabetes control
- •Gender disparity in dyslipidemia management
- Poorer dyslipidemia control in younger, at-risk patients

# **Dyslipidemia Comparison**

University of Chicago and NorthShore











#### **Baseline Data – Demographics**

Total N=20,871; Mean Age=64	N	%	Total N=84,463; Mean Age=64	N	%
Gender			Gender		
Male	10,658	51.1%	Male	43,772	51.8%
Female	10,213	48.9%	Female	40,681	48.2%
Race			Race		
African American	9,978	46.9%	White	58,778	69.6%
White	9,750	46.7%	Other	17,562	20.8%
Asian/Mideast Indian	643	3.1%	Asian	4,090	4.8%
Other	419	2.0%	African American	3,598	4.3%
American Indian or Alaskan Native	81	0.4%	American Indian or Alaskan Native	192	0.2%
CVD/CV Risk Diagnoses			CVD/CV Risk Diagnoses		
Hypertension	10,781	49.0%	Hypertension	44,338	52.5%
Dyslipidemia	7,632	34.7%	Diabetes	18,510	21.9%
Coronary Artery Disease	6,681	30.4%	Dyslipidemia	14,530	17.2%
Diabetes/prediabetes	1,658	7.5%	Coronary Artery Disease	12,908	15.3%
Peripheral Vascular Disease	955	4.3%	Peripheral Vascular Disease	2,821	3.3%

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### Cardiovascular Risk Factor Diagnoses (ICD 10)



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### **Baseline Data – Evolving Individual LDLC Change**



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### Statin Rx for ACC Risk Categories

#### Only includes patients on lipid-lowering drugs









#### **Statin Rx for ACC Risk Categories**

#### Only includes patients with diabetes on lipid-lowering drugs









#### **Baseline Data – LDLC Stratified**



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#### **Baseline Data – Patients Not at AACE Goals**



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