## Screening in Exchanges: Some Facts and Findings from Geruso, Layton, Prinz (2016)

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# Despite RA, Concerns about Screening in Exchanges

Thinking here about selection influencing not risk pool, but plan design



# Despite RA, Concerns about Screening in Exchanges

- Even in the absence of direct discrimination via premiums or coverage denials, possibility of dissuading consumers from joining plans via benefit design
- Anecdotes point to limiting access to entire classes of drugs as a backdoor discrimination. (Undoes intended protections for pre-existing conditions.)
- In November 2015, the National Multiple Sclerosis Society filed a comment with HHS's Office for Civil Rights explaining that "common health insurance practices that can discriminate against people with MS are formularies that place all covered therapies in specialty tiers."
- Separately, HHS has noted that one method indicating discrimination is to place "most or all drugs that treat a specific condition on the highest cost tiers."

# Drug Tiering in Exchanges/Marketplaces

- We study selection-related formulary design in 2015 in the ACA Exchanges
- Investigate whether drugs treating chronic conditions are a plausible screen
  - Prices are relatively transparent
  - Patient needs are predictable, and coverage may be salient at enrollment

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- First, examine whether there is scope for selection: Does drug use predict profits net of risk adjustment?
- Second, ask whether formularies of Exchange plans track the incentive

## Part 1: How Well is Payment System Performing in Neutralizing Screening Incentives?

# Selection Incentive - Data

- Marketscan administrative health insurance claims data (mostly self-insured employers) for about 12M people
- For each individual we observe
  - Demographics
  - Total spending
  - Prescription drug claims
  - All diagnoses appearing in claims
- Use HHS formulas/software to simulate person-specific plan revenues
  - Premiums
  - Risk adjustment transfer
  - Reinsurance
- Note that this is not Exchange data: Instead, we use it to produce out-of-sample predictions of which drugs insurers are incentivized to ration due to selection

### Selection Incentive - Aggregating up to Therapeutic Classes

- We group into standard therapeutic classes using REDBOOK e.g., *Anticoagulants* (blood thinners), *Antihyperlipidemics* (statins); *Oral Contraceptives*; *Antidiabetic Agents, Insulins*
- 220 mutually exclusive drug classes c
- Goal is to avoid conflating screening with steering patients to lower cost alternatives among classes of substitutes.
- From patient-specific costs,  $C_i$ , and revenues,  $R_i$ , calculate means  $\overline{C_c}$  and  $\overline{R_c}$  among consumers who fill a prescription for a drug in class c

Fact 1: For most classes, selection incentives neutralized



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Geruso (Various)

#### Part 1: Incentives

Fact 2: For some outliers, drug consumption signal of profitability



*biological response modifiers* (treat multiple sclerosis, others)

~\$61,000 in costs ~\$47,000 in revenue =

\$4,200 in premiums, \$34,420 in RA, and \$8,648 in reinsurance

Geruso (Various)

### Fact 3: No overall correlation between profitability and cost



- No correlation btwn cost and implied profit
- Implies RA + Reinsurance succeed in decoupling profitability from patient costs on avg
- Implies that if plan designs track these incentives, some sophistication on part of insurers

▶ zoom in ) ▶ zoom out

# Why the 'Errors' in the Payment System?

- Possible technological change in the intervening period between calibration and now (Carey 2016)
- HHS-HCC system based on Medicare Advantage's CMS-HCC system; in fact, does a good job compensating diabetes and heart disease.
- More generally, no reason to believe that predictors (drug utilization) that were not included in the RA algorithm are orthogonal to profitability

Fact 4: Reinsurance affects predictable profitability



- For the low cost groups (triangles on left) there is a small increase in profitability
- For the high cost groups (red lines on right) there is a large decrease in profitability

#### Part 2: Does Formulary Design Track the Incentive?

## Data

- Question: Are drugs that predict unprofitable patients covered ungenerously?
  - If an unprofitable group of consumers uses a cheap drug, an insurer will want to inefficiently distort coverage to be poor for that cheap drug
- Unit of analysis: drug class × plan, because class captures the set of substitutable therapies.
- We require data on formulary restrictiveness by drug class
  - Formulary tiering for the universe of state and federal exchanges in 2015 from MMIT

## Restrictiveness - Measure

- To measure restrictiveness we use harmonized tiers
  - 1. Generic Preferred
  - 2. Generic
  - 3. Preferred
  - 4. Covered/ Non-preferred Brand
  - 5. Specialty
  - 6. Not listed
  - 7. Medical
  - 8. Prior authorization/Step therapy
  - 9. Not covered
- We draw a line below "covered" and call tiers below the line "restrictive" and tiers above the line "non-restrictive"
- For each REDBOOK drug class, we define formulary restrictiveness as the % of drugs in the class on a restrictive tier

#### Fact 5: Drug Predicting Unprofitable Patients Are Restricted



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Fact 6: Drugs are a small share of spending even among groups whose drug use flags them as unprofitable. Indicates sophistication.



#### Screening in Exchanges

# What Are Insurers Responding To? Not Costs!



Already controlling for drug class FEs, but perhaps HIX plans are *differentially* attentive to high cost consumers...

Look within vertical slices: Equally costly but differentially profitable

Indicates sophistication

Geruso (Various)

# Concluding Observations

- 1. Risk adjustment + reinsurance do a good job overall in neutralizing screening incentives. *But some very unprofitable outliers exist*
- 2. Reinsurance important in reducing the incentive to avoid high-cost types
- 3. This is not about plans nudging consumers to lower cost or generic options
- 4. Both cost-sharing and utilization management are margins of distortion
- 5. It is not high drug costs that determine high cost sharing. It is drugs that are unprofitable, net of RA/Reinsurance. We see plans making it hard/expensive to access even cheap drugs.
- 6. EHB cannot solve this problem. Too many hard to measure and hard to regulate plan features (prior-authorization, requirement to use in-house mail-in pharmacy)
- 7. Problems may be solveable with fairly minor reforms
  - Incorporating diagnoses X drug utilization into RA scheme; currently considered

Geruso (Various)

# APPENDIX

Fact 5: HIX Formularies More Restrictive on Price and Non-Price

Figure : Frequency of Assignment to Restrictive Tier



Fact 5: HIX Formularies More Restrictive on Price and Non-Price

Figure : Frequency of Non-price Hurdles to Access



#### Selection Incentives - Top Drug Classes

#### Here limiting to classes with > 0.01% takeup

	March Hand Dava	Ora differen Ten eta di bu Marat	No.41 and
	Most Used Drug	Conditions Treated by Most	Net Loss:
Class	in Class	Used Drug	Cost - Revenue
(1)	(2)	(3)	(4)
Largest Incentives to Avoid			
Gonadotropins, NEC	Ovidrel	infertility in women	\$15,326
Biological Response Modifiers	Copaxone	relapsing multiple sclerosis	\$13,977
Opiate Antagonists, NEC	naltrexone	substance abuse disorders	\$5,977
Ovulation Stimulants, NEC	clomiphene citrate	infertility in women	\$5,304
Pituitary Hormones, NEC	desmopressin	diabetes insip., hemophilia A	\$4,633
Vitamin A and Derivatives, NEC	Claravis	severe nodular acne	\$4,428
Analg/Antipyr, Opiate Agonists	hydrocodone-acetamin.	moderate to severe pain nerve pain; fibromyalgia;	\$3,001
CNS Agents, Misc.	Lyrica	seizure poisonings; pre-surgical	\$2,965
Mydriatics EENT, NEC	atropine	preparations	\$2,877
Androgens and Comb, NEC	AndroGel	low testosterone	\$2,688

#### Selection Incentives - Top Drug Classes

#### Largest Incentives to Attract

methotrexate sodium	autoimmune diseases	-\$2,885
Folbic	vitamin deficiency	-\$3,058
warfarin	blood clots; stroke prevention primary biliary cirrhosis:	-\$4,328
ursodiol	gallstones	-\$4,751
	edema due to heart, liver, kidney disease; high blood	
furosemide	pressure	-\$5,813
lactulose	complications of liver disease seziures; heart arrhythmias;	-\$7,181
phenytoin sodium ext.	neuropathic pain	-\$7,275
amiodarone	heart arrhythmias chronic pancreatitis; cystic	-\$7,942
Creon	fibrosis; pancreatic cancer heart arrhythmias; heart	-\$12,350
Digox	failure	-\$12,857
	methotrexate sodium Folbic warfarin ursodiol furosemide lactulose phenytoin sodium ext. amiodarone Creon Digox	waturous calibers, valuous   methotrexate sodium   Folbic   warfarin   blood clots; stroke prevention   primary biliary cirrhosis;   ursodiol   gallstones   edema due to heart, liver, kidney disease; high blood   furosemide   pressure   lactulose   complications of liver disease seziures; heart arrhythmias;   phenytoin sodium ext.   amiodarone   heart arrhythmias chronic pancreatitis; cystic   Creon   bigox   failure

various cancers: various

Fact 1: For most classes, selection incentives neutralized **Pack** 



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Fact 3: No overall correlation between profitability and cost **Pack** 



Fact 3: No overall correlation between profitability and cost • Back



Most classes are clustered very near neutral

Ratio Measure

Ellis-McGuire Measure



#### Residuals: Difference Measure

Residuals from  $Y_{jc} = \gamma_c + \alpha_j + \epsilon_{cj}$ Grouping classes into 20 bins by selection incentive (Difference).



#### Residuals: Ratio Measure

Residuals from  $Y_{jc} = \gamma_c + \alpha_j + \epsilon_{cj}$ Grouping classes into 20 bins by selection incentive (Ratio).



Moral Hazard? We recode data to be matchable to Einav, Finkelstein, and Polyakova (2016)



# Moral Hazard? No: Selection Incentive Uncorrelated with Elasticity • Back

