Leveraging Data from the Illinois Prescription Monitoring Program to Address the Opioid Epidemic Through Academic Detailing

June 26, 2019

Sarah Pointer, PharmD Clinical Director of the Illinois Prescription Monitoring Program Bureau of Pharmacy & Clinical Support Services

Sarah.Pointer@illinois.gov

Christopher D. Saffore, PharmD Department of Pharmacy Systems, Outcomes and Policy College of Pharmacy, University of Illinois at Chicago

<u>csaffo3@uic.edu</u>



## **Disclosure & Funding Source**

- Disclosure
  - The presenters have nothing to disclose.
- Funding source
  - The Centers for Disease Control and Prevention. Grant #1U17CE002739-01.





### <u>Outline</u>

- Illinois Prescription Monitoring Program Initiatives
- Overview of Academic Detailing Initiatives
- Preliminary Evaluations and Outcomes
- Implications of Academic Detailing Outcomes





## **Acknowledgements**

- IL PMP
  - Craig Berberet
  - Stan Murzynski
  - Ed Dowllar
  - Andrew Hollo
- IDPH
  - Dejan Jovanov
- CDC
  - Jamie Mells, PhD
  - Wes Sargent, EdD
- AMITA Health
  - Darin Jordan, MD
  - Ankur Dave, MD
  - Reinhold Llerena, MD
- SIUE
  - Chris Herndon, PharmD
- SIHF
  - Theodore Ross, MD

- University of Illinois at Chicago
  - Evaluation Activities
    - A. Simon Pickard, PhD
    - Todd A. Lee, PharmD, PhD
    - Shan Xing, PharmD, PhD
    - Inyoung Lee
    - Mary Schiff, MPH
    - Andrea Monteiro, MS
    - Maja Kuharic, MS
  - Academic Detailing
    - Christopher D. Saffore, PharmD
    - Mary Smart, PharmD, MS
    - Farah Khan
    - Sarette Tilton
    - Aleksandrina Ruseva
    - Dayna Redini
    - Esther Lee
    - Nevena Varagic
    - Shannon Menard
    - Victoria Kulbokas
    - Ammarah Nadeem
  - Medication Review and Academic Detailing (MRAD) group
    - Mary Lynn Moody, BSPharm



SYSTEMS OUTCOMES AND POLICY COLLEGE OF PHARMACY

PHARMACY

# Illinois Prescription Monitoring Program (IL PMP)

- IL PMP one of oldest PMPs
- Home-grown
   system
- Captures data
- from pharmacies



on all controlled substance prescriptions as well as naloxone



#### State of Illinois Opioid Action Plan



#### **IL PMP Initiatives**

Focus in four key areas:

- 1. Identify High Risk Behaviors
- 2. Provide Education
- 3. Increase Utilization of the PMP
- 4. Prevent Overdose





### Strategies to Achieve Initiatives

- Academic detailing (AD) may be used as a strategy to achieve IL PMP initiatives
- AD is a method of educational outreach<sup>1,2</sup>
  - One-on-one, face-to-face, encounters with clinicians
- Utilizes trained academic detailers to provide current, unbiased evidence-based information
   Circle of Trust
- Aims to improve prescribing behavior
- Most effective when trusting relationship between provider and detailer

1. Avorn J, Soumerai SB. Improving drug-therapy decisions through educational outreach. A randomized controlled trial of academically based "detailing". N Engl J Med. 1983;308(24):1457-63.

Soumerai SB, Avorn J. Principles of educational outreach ('academic detailing') to improve clinical decision making. JAMA. 1990;263(4):549-56.









## Academic Detailing is Not

- Didactic lecture in healthcare provider's office
- Written materials or emails sent directly to providers
- Focused solely on cost savings or limiting industry influence
- Punitive in nature







#### Importance of Tailoring Academic Detailing Programs

Challenges when developing and implementing AD programs

□ Variations in prescribing patterns

Establishing partnerships

Educational messages





#### Establishing Partnerships

- Essential when developing and implementing AD programs
  - State-based prescription monitoring programs (PMP)
  - State departments of health and human services
  - Local academic institutions
  - Provider groups & healthcare systems
  - National Resource Center for Academic Detailing (NaRCAD)







#### **Illinois Opioid AD Program Implementation**





#### AD Program Summary

- Complete 2 visits with primary care providers (MD, DO, NP, PA)
  - Visit length between 15 and 30 minutes
  - 2 visits separated by 6 to 8 weeks
- Content development
  - Focused on CDC prescribing guidelines
  - Tailored to needs of providers
  - Prescriber-specific data
- Detailer training
  - NaRCAD train-the-trainer model
  - Quality assurance and troubleshooting
- Evaluation
  - Effect of the AD
  - Development of AD tools



#### **CDC Guidelines Key Messages**

#### 1. Opioids are not first-line therapy

- 2. Establish goals for pain and function
- 3. Discuss risks and benefits
- 4. Use immediate-release opioids when starting
- 5. Use the lowest effective dose
- 6. Prescribe short durations for acute pain
- 7. Evaluate benefits and harms frequently
- 8. Use strategies to mitigate risk
- 9. Review PDMP Data
- 10. Use urine drug testing
- 11. Avoid opioids and benzodiazepine co-prescribing
- 12. Offer treatment for opioid use disorder

**Red** = Key messages covered

#### EFFECTIVELY AND RESPONSIBLY MANAGE CHRONIC PAIN

GUIDELINE FOR PRESCRIBING OPIOIDS FOR CHRONIC PAIN





GUIDELINES FOR PRIMARY CARE PROVIDERS	REVIEW DDMP (Recommendation #9)	
Primary care providers occurit for opproximately 50% of prescription opicids of prescription o	er drunic     and report     er drunic     and report	te prescription chigh risk for
	sin a	alan a the
WITH US IRVIN	USE NON-OPIOID TREATMENT (Recommendation #1)	STRATEGIES TO MITIGATE RISK (Recommendation #8)
Opioids are effective long-term evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic pain effectively over the long to evidence that capadid control chronic	Non-pharmacologic therapy and non-opioid pharmacologic therapy are preferred for chronic pain.     Consider opioid therapy only if expected benefits for both pain and function are anticipated to outwaigh risks to the nation.	<ul> <li>Before starting and periodically after, evaluate risk factors for opioid-related harms.</li> <li>Consider offering naloxone when there is an increased risk for opioid overdoses (i.e. history of overdose and/or substance use disorder, higher opioid dosages (≥50 MME/day), or concurrent benzodiazepine use).</li> </ul>
WHAT CAN PROVIDERS DO?  First, do no harm. Long-term opioid use has uncertain benefits  First, do no harm. Long-term opioid u	<ul> <li>If opioids are used, they should be combined with non-pharmacologic therapy and non-opioid pharmacologic therapy, as appropriate.</li> <li>Non-opioids available over the counter for mild pain: <sup>24</sup></li> </ul>	NALOXONE Rx <sup>5</sup> NARCAN Nasal Spray 4mg Nasal Spray #1 (two-pack) Directions: PRN for opioid overdose (Pace and hold tip of nozzle in either nostril. Press plunger rimit to release doe into patient's nose. Report with second dreige into other nozifil drez 2-3 minutes if no or
PRACTICES AND ACTIONS  PRACTICES AND ACTIONS  PRACTICES AND ACTIONS  Provide preserving request  Provide preservin	IBUPROFEN (Advil, Motrin): 400 mg every 4-6 hours, as needed for pain ACETAMINOPHEN (Tylenol): 325 - 650 mg every 4-6 hours, as needed for pain (do not exceed 4,000 mg in a day; or 3,000 mg if over 65 years old)	PVZIO Auto-Injector     Z mg Auto-Injector     Z mg Auto-Injector     PVZIO Auto-Injector     PUZIO Auto-Injector     PUZ
USE NON-OPIOID TREATMENT Coloids are not first-line or routine therapy for dronic pain (Riccommondation #1) USE NON-OPIOID TREATMENT The lowest effective does (Riccommondation #5)	START LOW AND GO SLOW (Recommendation #5)      Prescribe the lowest effective dosage when starting opioids.	OFFER TREATMENT FOR OPIOID USE DISORDER (Recommendation #12)     As many as     Offer or arrange evidence-based treatment     (northear lifeting evidence-based treatment)
STRATEGIES TO MITIGATE RISK     Incorporte strategies to mitigate rule     Source of the strategies to mitigate     source of the strategies     source of the strategies to mitigate     source of the strat	<ul> <li>Reassess individual benefits and risks at dosages ≥ 50 MME/day.</li> <li>Avoid increasing dosage to ≥90 MME/day.</li> <li>HOW MUCH IS 50 OR 90 MME/DAY FOR</li> </ul>	patients receiving long-term opioid therapy in primary care settings
Alther recommendations GRADEA Indicating that most recommended course of action	COMMONLY PRESCRIBED OPIOIDS? Dosages ≥ 50 MME/day increase risks for overdose by at least 2 X · · · · · · · · · · · · · · · · · ·	Identify treatment resources for opioid use disorder in the community and ensure sufficient treatment capacity for opioid use disorder at the practice level.
LEARN MORE   www.cdc.gov/drugoverdose/prescribing/gu and profession	90 MME/DAY 90 mg of hydrocodone (9 tablets of 10/325) • 60 mg of oxycodone (2 tablets of sustained release 30 mg)	Strugger wire oppreid Use disorder.  Iowed DH, Dave KCK, Galdacter Proceing-Opioid En Chronic Pain – United States, 2015, MMMR Researce Rg 2016, 2016;555(1):14-9.  Anorchoot Structure RT 27. The Information Canada States, 2016, Pain 2016,
	<ul> <li>20 mg of methadone (4 tablets of methadone 5 mg)</li> </ul>	For more information please visit



PHARMACY SYSTEMS OUTCOMES AND POLICY COLLEGE OF PHARMACY

UIC

**Provider-specific Information** 

- Audit and feedback is a widely used strategy to motivate behavior change
- Feedback on provider clinical performance was provided via opioid prescribing information
- Provider-specific opioid prescribing information was obtained from the IL PMP
- Detailers shared this information with providers at each visit







#### Illinois Prescription Monitoring Program Dashboard Metrics

Below is a 6-month comparison (Nov 2017-Apr 2018) of your number of opioid prescriptions ( morphine milligram equivalents (MME) per day, average number of monthly opioid prescripti number of monthly PMP queries along with the average for all AMITA Health primary care priproviders from Cook County.

	You		AMITA Health	
<50 MME/day (%) <sup>1a</sup>	27	(56%)	12,903	(76%)
50 - 89 MME/day (%)	14	(29%)	3,668	(22%)
≥90 MME/day (%)	7	(15%)	312	(2%)
Average number of monthly opioid prescriptions	8	3.0	13.5	
Average number of monthly PMP queries	0.0		3.	6

1. % = proportion of total opioid prescriptions over the 6-month period

a. i.e. Your MME/day <50 = 10%, meaning 10% of your total opioid prescriptions over 6 months v



Email: info@ilpmp.org | Website: www.ilpmp.org



PHARMACY SYSTEMS OUTCOMES AND POLICY COLLEGE OF PHARMACY

UIC

#### Quality Assurance Process

- Detailers documented visits in field notes
- Field notes reviewed by program coordinators
- Weekly detailer phone calls
- Provider satisfaction measure





#### Second Visit Differences in Delivery

- Key difference in delivery of second visit
- In-person vs. technology-based









#### Providers Visited in Urban and Rural Sites

#### Phase I: Urban Providers

Provider characteristics		
Total Providers, n	186	
Sex, n (%)		
Female	103	(55.4)
Male	83	(44.6)
Years of Practice, mean (SD)		
Mean	14.6	(12.0)
Provider Type, n (%)		
MD/DO	160	(86.0)
PA/NP	26	(14.0)

#### Phase II: Rural Providers

#### **Provider characteristics** 119 Total Providers, n Sex, n (%) 56 (47.0)Female 63 (53.0)Male Years of Practice, mean (SD) 13.8 (11.0)Mean Provider Type, n (%) 76 (63.9)MD/DO 43 (36.1)PA/NP



#### Provider Satisfaction Measure Results

<u>ltem*</u>	<u>Urban</u>	<u>Rural</u>
This is an important topic	97%	100%
The detailer was knowledgeable	93%	100%
The detailer was an effective communicator	96%	100%
The key messages are feasible to implement in my practice	89%	94%
My practice is likely to change as a result of this visit	49%	69%
I would be receptive to future visits	78%	94%

\*Response options: "not at all", "slightly", "moderately", "very", or "extremely". The results reported are for "very" or "extremely" responses



UIC

#### Preliminary Evaluations

- Change in mean monthly number of:
  - Total opioid prescriptions
  - High dose opioid prescriptions (>90 MME/day)
  - Patients co-prescribed opioids and benzodiazepines
- Outcomes measured at six months post-AD program implementation (September 2018 to February 2019)
- Comparison groups: Academic detailing vs. No academic detailing
- Used Difference-in-Difference approach to compare two groups before and after AD visits



## **Preliminary Evaluations**

Table 1. Baseline demographics comparison between AD-Exposed and AD-Unexposed providers in the Urban region

	Ov	Overall		AD-Exposed		AD-Unexposed	
n (%)	5	550		(27.5%)	399	(72.5%)	
Sex							
Female	286	(52.0%)	88	(58.3%)	198	(49.6%)	
Male	264	(48.0%)	63	(41.7%)	201	(50.4%)	
Years of Practice							
Median (interquartile range)	19	(17)	18	(15)	19	(17)	
Provider Type							
MD	423	(76.9%)	87	(57.6%)	336	(84.2%)	
DO	74	(13.5%)	38	(25.2%)	36	(9.0%)	
NP	34	(6.2%)	18	(11.9%)	16	(4.0%)	
PA	19	(3.5%)	8	(5.3%)	11	(2.8%)	
Provider Specialty							
Family Medicine	228	(41.5%)	115	(76.2%)	113	(28.3%)	
Internal Medicine	322	(58.5%)	36	(23.8%)	286	(71.7%)	



UIC

#### **Preliminary Outcomes**

Table 2. Difference-in- Difference Estimates for Mean Monthly TotalOpioid Prescriptions per Provider

	Pre-AD Mean	Post-AD Mean	D-I-D Estimator	95% CI	P-value
AD-exposed	15.22	15.51	0.85	(126 0 22)	0.001
AD-unexposed	13.86	15.00	-0.85	(-1.30, -0.33)	0.001

#### Interpretation:

- On average, nearly 1 less opioid prescription per month per provider were dispensed among AD-exposed providers relative to AD-unexposed providers
- This translates to ~1,500 fewer opioid prescriptions dispensed annually (Ex: -0.85 opioid prescriptions x 151 AD-exposed providers x 12 months = ~1,500 fewer opioid prescriptions)



## Preliminary Outcomes (Cont'd)

Table 3. Difference-in- Difference Estimates for Mean Monthly High-doseOpioid Prescriptions per Provider

	Pre-AD Mean	Post-AD Mean	D-I-D Estimator	95% CI	P-value
AD-exposed	0.86	0.55	-0.11	(-0.24, 0.01)	0.08
AD-unexposed	1.10	0.90			

#### Interpretation:

- On average, 0.11 fewer high-dose opioid prescriptions per month per provider were dispensed among AD-exposed providers relative to ADunexposed providers
- This translates to ~200 fewer high-dose opioid prescriptions dispensed annually (Ex: -0.11 opioid prescriptions x 151 AD-exposed providers x 12 months = ~ 200 fewer high-dose opioid prescriptions)



### Preliminary Outcomes (Cont'd)

Table 4. Difference-in- Difference Estimates for Mean MonthlyPatients Co-Prescribed Opioids and Benzodiazepines

	Pre-AD Mean	Post-AD Mean	D-I-D Estimator	95% CI	P-value
AD-exposed	3.68	3.36	-0.22	(-0.41, -0.04)	0.02
<b>AD-unexposed</b>	3.31	3.21			

#### Interpretation:

- On average, 0.22 fewer patients were co-prescribed benzodiazepines and opioids per month per provider among AD-exposed providers relative to AD-unexposed providers
- This translates to ~400 fewer patients co-prescribed benzodiazepines and opioids annually (Ex: -0.22 patients co-prescribed benzodiazepines and opioids x 151 AD-exposed providers x 12 months = ~ 400 fewer patients co-prescribed benzodiazepines and opioids)



#### **Implications**

- <u>Establishing partnerships are crucial</u> for implementation of strategies to achieve initiatives that address the opioid epidemic
- AD was effective at reducing <u>the number of opioid prescriptions</u> and patients co-prescribed benzodiazepines and opioids among AD-exposed providers relative to AD-unexposed providers
- Future efforts should include <u>scaling-up of opioid-related AD</u> <u>programs</u> for delivery to other relevant providers (surgeons, dentists, etc.) across the state



#### Next Steps

- Evaluate AD program in southern Illinois
- Continue evaluating the impact of the AD initiative on changes in opioid prescribing rates, duration of days supply, and accessing the PMP
- Explore opportunities for continuation and expansion of our AD initiatives
- Evaluate additional impacts of AD through endpoints such as naloxone prescribing, opioid-related hospitalizations, opioid-related deaths





PHARMACY SYSTEMS OUTCOMES AND POLICY COLLEGE OF PHARMACY

UIC

