

# **Mobile treatment units for medication-based treatments of Opioid Use Disorder**

**An epidemiology study design project**

Thadryan J. Sweeney

1/16/23

# Table of contents

<b>1 Study Description</b>	<b>3</b>
<b>2 Scientific Question (Assn. 1)</b>	<b>4</b>
2.1 Choose one of the scientific questions that you have proposed above and answer the following questions. (Though you may change your study question later) .	4
2.2 Focus in on what is pragmatic or logistically possible to answer the following questions about your scientific question and study design. . . . .	6
2.3 Place your chosen question into the broader context of the existing literature. .	6
2.4 Are there any relevant sources of information bias to consider for your study as designed (consider all potential types of information bias)? How might you prevent these or improve exposure/outcome/covariate data collection to minimize these concerns? . . . . .	7
2.4.1 Recall Bias . . . . .	8
2.4.2 Interviewer Bias . . . . .	8
2.4.3 Loss to follow up . . . . .	8
2.4.4 Misclassification of exposure or outcome . . . . .	8
<b>3 Confounding (Assn. 2)</b>	<b>9</b>
3.1 2. Based on the papers that you reviewed for Questions 3 & 4 in Exercise 1, list the important potential confounders of your exposure-outcome association. Will any of these be particularly challenging to measure? . . . . .	9
3.2 3. How might you integrate prevention or control of confounding into your study design or analysis? . . . . .	9
3.3 4. Based on your answers in part 3 and 4 of Study Design Assignment 1, create a preliminary DAG... . . . .	9
3.3.1 a. Describe/define each individual component in the DAG. . . . .	9
3.3.2 b. Was it difficult to assess directionality of any of the arrows? What additional information would you like to have . . . . .	9
<b>References</b>	<b>10</b>

# 1 Study Description

I propose a study of the impact of mobile treatment vans for methadone delivery to people with opioid use disorder (OUD). A large body of evidence spanning decades supports the use of Methadone (also called Methadone and Dolophine) for treatment of OUD [1]. Despite this, the majority of people suffering from the disorder do not get medication-based treatment [2]. By law Methadone treatment can only be conducted by specially-licensed practitioners and those in treatments are required to report to the clinic every day to get the treatment, at least initially [1]. These policies have been implicated as barrier to treatment [3]. The start of the pandemic caused a wave of logistical restructuring across field, including healthcare, and the use of mobile treatment units (MTUs) that can bring medications to people by van was considered as an option to improve access[4]. A growing line of research assesses the effectiveness and feasibility of this treatment modality[5].

## 2 Scientific Question (Assn. 1)

Study Design Assignment 1: Matching your scientific question to the best study design and preventing information bias. Provide a concise but complete response (At least 2, but not more than 4 pages) to the following questions.

- a. Do mobile treatment units for Methadone increase the access to care for people with OUD? This might be hard to assess but I think it's worth coming up with something because it's really the core of effort to get better treatment to more people. Sub-question: does this result in fewer adverse events?
- b. Do mobile treatment units change the demographics of people receiving methadone? The demographics of the opioid epidemic are complex and have shifted since it began with working class in whites in deindustrialized areas making up the majority of cases and other groups becoming more involved more over time. Monitoring for demographic shifts could help assess if the programs were helping in a fair way.
- c. Do mobile treatment units result in fewer cases of COVID among people with OUD? One thing worth interrogating is if switching to the vans would help at all. If not, it might make sense to divert resources back to the initial clinics. If it reduces infections, it could serve as a treatment model for other diseases.

### 2.1 Choose one of the scientific questions that you have proposed above and answer the following questions. (Though you may change your study question later)

- a. What is your conceptual exposure? Is this exposure rare or common?

My exposure is opioid use disorder in a region where MTUs are being explored as a potential treatment delivery system. It's rare compared to something like cardiovascular disease, and I don't think it's greater than 20% of the population (the rough rule we're using for this class) [double check]. It is increasingly common however, having become the leading cause of accidental death [find where I read this].

- b. What is your conceptual outcome? Is this outcome rare or common?

The outcome would be receiving medication-based treatment for opioid use disorder via a MTU (let's say, >10% of the time)[note: might need to workshop this]. It would also make sense to track adverse events like hospitalizations, overdoses, and deaths.

c. Briefly describe how you might use each of the four major study designs in epidemiology (cohort, case-control, cross-sectional (or ecologic if you like), or randomized trial) to assess this question. For purposes of this exercise, I'd like you to stretch your ideas about study design, so do your best to come up with a way to use every one of the study designs to address your question of interest. Feel free to be a bit creative for this part of the question (you will assess feasibility and logistics in the next part of the question).

- Cohort: In this design, we could enroll people based on their opioid use disorder status and the policy of the closets methadone clinic (if they use vans or not). After the enrollment, we would follow them for a designated period. We could then assess a variety of metrics like adherence, overdoses, or death. The main one would be adherence, defined as people who reported to the van. We could then compare this to the rates in the people who didn't have the van option. This would be the prospective option. For the retrospective version, we could attempt to find data on overdoses, demographics, etc from historical database. We could then see if people in the MTU regions different in terms of endpoints.
- Case control: This one stretches my proposal quite a bit. I need to think on it [return to this]. In this we would need to define a different endpoint, let's say overdose. We could enroll people in a non-MTU region as controls and a MTU region as cases. We could follow them and see if they different on our endpoints. OUD is likely uncommon enough that odds ratio would approximate the risk.
- Cross-sectional: In this method, we might obtain interview or medical records. We would assess the desired information on demographics and outcomes. We could review the records to see if subjects had an overdose, etc. The advantage here is that it's the cheapest and fastest. The downside is that we wouldn't have continuity in terms of time and would have very little control over confounding.
- Randomized trial: In this version of the study we would randomly assign people to treatment regions where MTUs were available and some to those where they weren't. This would be extremely difficult logistically because the point of the MTUs is to address access, and assigning people to facilities other than the closest one would be prohibitive. Perhaps it would be possible to assign the MTU programs randomly as pilot programs instead of using the places where it was already in use.

## **2.2 Focus in on what is pragmatic or logistically possible to answer the following questions about your scientific question and study design.**

- a. Which study design from part 2c seems most feasible? Why does this design seem best for addressing your scientific question?

I think there are two that could work here, the cohort and the cross-sectional study. The cohort would be more ambitious - it would require following people who might be at higher risk for housing instability over a long period of time if the prospective option was used. If the data were available already, it could be approached retrospectively. This might be the most practical option in terms of balancing feasibility with robustness.

The cross-sectional might be the best we could do if the data were only available as a snapshot.

- b. What will you use as your operational exposure and outcome? Or what are some reasonable options for operational exposure and outcome? Note that this should match up to the study design you've identified as most feasible in 3a.

Let's assume we can do a retrospective cohort study. In this case the operational exposure would be a medical record reporting use of illegal opioids. The patients would then be categorized as having used a MTU for medical treatment of OUD or having used a traditional clinic. The outcome would be reporting to the ER with an overdose or a fatality.

- c. What other data will you need to collect for you study (i.e., what are the important covariates for your study)?

Given the complex demographic makeup of the opioid crisis[2], it would be crucial to collect information such as race and sex, and given the variation in risk with employment and education, information on work and schooling. Geographic information may be required as well.

## **2.3 Place your chosen question into the broader context of the existing literature.**

- a. Identify 2-4 relevant papers from the primary literature to provide background and motivation for your proposed study. Provide the citations and a 1-2 sentence summary of the critical background information contained in each study.

Joudrey PJ, Edelman EJ, Wang EA, “Methadone for Opioid Use Disorder—Decades of Effectiveness but Still Miles Away in the US” [1] provides evidence of the effectiveness of methadone as well as background on the restrictions around using it. It also contextualizes status quo by discussing the political and policy factors that created it.

Cerdá M, Krawczyk N, Hamilton L, et al, “A Critical Review of the Social and Behavioral Contributions to the Overdose Epidemic” [2] provides an exhaustive overview of epidemic with a focus on the demographic and policy aspects. It also includes a history of the epidemic and an analysis of economic drivers of supply and demand of drugs.

Jakubowski A, Fox A, “Defining Low-threshold Buprenorphine Treatment” [3] specifically discusses the barriers to entry in pursuing medication-based treatment for OUD. Its main contribution is a definition of and argument for treatment policies that emphasize making treatment more easy to get.

Chan B, Hoffman KA, Bougatsos C, et al “Mobile methadone medication units: A brief history, scoping review and research opportunity” [5] describes the history of mobile treatment units. It also lays out the case for using them as an opportunity to assess drug-based treatments. In particular, because they are becoming legal and being tested, we will now when the change was adopted for comparison.

- b. What knowledge gap does your proposed study address? (i.e., Will it add to our scientific knowledge by answering a completely new question? Will it help us understand a new mechanism to explain a previously observed association? Will it extend the research to a new population?)

My proposed study would assess the demographics of those who seek medication-based treatment for OUD in a mobile setting. This would offer insights into what communities benefit the most, if there is a disparity. This would highlight needs for future work. If the vans are broadly effective and demographics do not show a disparity, it would confirm their value. It could also interrogate the utility in improving outcomes. The demographics would be a new contribution as would the outcomes. The benefit of medication assisted treatment in general is well established and would not be a contribution.

## **2.4 Are there any relevant sources of information bias to consider for your study as designed (consider all potential types of information bias)? How might you prevent these or improve exposure/outcome/covariate data collection to minimize these concerns?**

There are numerous ways this study could be impacted by information bias.

### **2.4.1 Recall Bias**

If we were to pursue the cross-sectional version of the study this would likely be a large issue. Given that opioids influence the mind there might be issues in recalling if, when, and how many non-fatal overdoses occurred. We could restrict to participants with medical records going back a certain amount of time to help with this.

### **2.4.2 Interviewer Bias**

The interviewer bias would be more likely if we chose the cross-sectional version of the study. In this situation an interviewer might be more likely to assume people with certain traits are more likely to be drug users and ask leading questions accordingly (“Are you sure there weren’t more?”, etc). We could address this by automating simple interviews (ie, using some kind of survey tool).

### **2.4.3 Loss to follow up**

If we were to pursue a cohort study, we might have issues if people having moving or transferring care facilities. In particular, given that housing instability is likely higher among people who use drugs, it could be hard to follow up with people with no permanent address. Given that opioid use is criminalized, people could be arrested before the study is complete. I think this would be one of the biggest threats to the study in practice if we did the cohort version. We might be able to ameliorate the address by only enrolling people with a legal address though this would cost us in terms of generalizability. In terms of the legal issues there isn’t a lot we could do.

### **2.4.4 Misclassification of exposure or outcome**

This could manifest in terms of counting overdoses. Someone could take a dangerous amount of a substance and not die but not report to a hospital. This “close call” should, in theory, constitute an overdose in terms of risky behavior, but would not appear on a medical record. This is exactly the kind of thing that gets underestimated in epidemiological studies. Correcting it sully is likely impossible. We could probably not change it but we could consider the effect it would have when interpreting the results. For example, this effect often biases towards the null.



## 3 Confounding (Assn. 2)

Study Design Assignment #2: Creating a strategy for the control of confounding

Provide a concise but complete response (1-2 pages) to the following questions.

**3.1 2. Based on the papers that you reviewed for Questions 3 & 4 in Exercise 1, list the important potential confounders of your exposure-outcome association. Will any of these be particularly challenging to measure?**

**3.2 3. How might you integrate prevention or control of confounding into your study design or analysis?**

**3.3 4. Based on your answers in part 3 and 4 of Study Design Assignment 1, create a preliminary DAG...**

...to define set  $\{S\}$  to describe which confounders you may wish to include in a multivariable model. You may use Daggity, R, Powerpoint, etc., to make your DAG. A clear and legibly hand drawn DAG is also acceptable. (Note: If you chose a randomized trial, please use the DAG to help you describe the confounding structure that will be accounted for via randomization).

**3.3.1 a. Describe/define each individual component in the DAG.**

**3.3.2 b. Was it difficult to assess directionality of any of the arrows? What additional information would you like to have**

(Note: The goal is not necessarily to produce a perfect and final DAG, but to help you gain a

## References

- 1 Joudrey PJ, Edelman EJ, Wang EA. Methadone for Opioid Use Disorder—Decades of Effectiveness but Still Miles Away in the US. *JAMA Psychiatry* 2020;**77**:1105–6. doi:[10.1001/jamapsychiatry.2020.1511](https://doi.org/10.1001/jamapsychiatry.2020.1511)
- 2 Cerdá M, Krawczyk N, Hamilton L, *et al.* A Critical Review of the Social and Behavioral Contributions to the Overdose Epidemic. *Annu Rev Public Health* 2021;**42**:95–114. doi:[10.1146/annurev-publhealth-090419-102727](https://doi.org/10.1146/annurev-publhealth-090419-102727)
- 3 Jakubowski A, Fox A. Defining Low-threshold Buprenorphine Treatment. *J Addict Med* 2020;**14**:95–8. doi:[10.1097/ADM.0000000000000555](https://doi.org/10.1097/ADM.0000000000000555)
- 4 Krawczyk N, Fingerhood MI, Agus D. Lessons from COVID 19: Are we finally ready to make opioid treatment accessible? *J Subst Abuse Treat* 2020;**117**:108074. doi:[10.1016/j.jsat.2020.108074](https://doi.org/10.1016/j.jsat.2020.108074)
- 5 Chan B, Hoffman KA, Bougatsos C, *et al.* Mobile methadone medication units: A brief history, scoping review and research opportunity. *J Subst Abuse Treat* 2021;**129**:108483. doi:[10.1016/j.jsat.2021.108483](https://doi.org/10.1016/j.jsat.2021.108483)