

Modeling the Heroin Epidemic

Suzanne Lenhart, Tricia Phillips, Christopher Strickland

April 26, 2019

Blue font: contains questions I have, things to discuss.

Red font: things I am working on/need to do.

Reminder for me: use present tense, use powerful/direct language.

Abstract

Fill in after write paper.

Introduction

Want to “funnel in”: start with general epidemic information and then narrow in on the drugs involved.

General opioid epidemic

Re-order to better fit funnel with “all” stats first, then prescription opioids, then heroin, then fentanyl (follows flow of people better, too)

Fill in stats from various sources in below paragraphs.

Find motor vehicle vs. drug overdose stats

Start of stats about opioid epidemic at large to begin paragraph: The misuse of opioids, including prescription pain relievers, synthetic opioids, and the illegal drug heroin, is rampant in today’s society [18]. The opioid crisis was declared a public health emergency in October 2017 by the United States Department of Health and Human Sciences [62]. It was estimated that the total economic cost of prescription opioid dependence, abuse and overdoses in 2013 alone was \$78.5 billion [24]. Put in idea about how fentanyl has become leading cause of overdoses, etc. Due to the health risks of the addicted individuals, public health concerns, the number of overdose deaths and the economic burden, this is an issue worth paying attention to.

Statistics regarding heroin and why we should care:

A national survey estimated that out of the United States national population of individuals of 12 years of age or older, 948,000 were heroin users in 2016, although this number may include under-reporting [56]. There were an estimated 13,219 heroin deaths in 2016, a more than six-fold increase from the year 2002, despite there being roughly half as many heroin users in 2002 [55]. This, in part, is due to the recent trend of lacing heroin with fentanyl (put in background on fentanyl), a surgical-grade opioid that is up to fifty times more potent than heroin alone, and therefore, users are unaware of the purity of the heroin they obtain [10, 18, 65]. Find source regarding purity of heroin trends/graphs for past few decades. The effects of heroin, however, extend further than just the specific individual using the drug. Due to the sharing of needles and other equipment involved in the injection of heroin, the human immunodeficiency virus (HIV) and the hepatitis C virus (HCV) are more easily contracted as transmittance occurs through bodily fluids. In

addition, risky sexual behavior is heightened under the influence, so addicts have an increased risk of transmitting and contracting these viruses. Women who use heroin during pregnancy put their babies at risk for neonatal abstinence syndrome in which the drug is passed along to the baby, resulting in dependency and thus, withdrawal symptoms upon birth [40].

Prescription opioid information

About prescription opioids and why we should care:

Another large portion of opioids consist of prescription pain relievers which are either natural (morphine and codeine), semi-synthetic (oxycodone, hydrocodone, oxycodone), or synthetic, (fentanyl, tramadol) [12, 58]. From 1991 to 2011, there was a near tripling of prescriptions that pharmacies distributed [43]. This was in part due to a number of new opioids that were approved by the FDA for use, such as OxyContin, Actiq, Fentora, and Onsolis (fentanyl), in addition to other unapproved opioid products for pain management [63]. Moreover, in the early 2000s, drug manufacturers funded publications and physicians to support opioid use for pain control [37]. A national survey estimated there were 11.5 million individuals of 12 years of age or older in the United States that experienced pain reliever misuse in the past year, referring to the year 2016 [56]. In this survey, misuse was defined as taking the prescription at a higher dose, more frequently or longer than prescribed, taking someone else's medication, or any other way not directed by a doctor [57]. Misuse of prescription pain relievers occurs for a variety of reasons and the same survey asked individuals who reported prescription pain reliever misuse to give the reason and the source for their most recent misuse. The most prominent responses for reasons of misuse were to relieve physical pain, to feel good or get high, and to relax or relieve tension. The largest source was from friends/relatives or from a healthcare provider, followed by given a prescription or stolen from a health care provider [56]. Options for treatment of opioid addiction involve medications such as buprenorphine, methadone, and naltrexone, in combination with counseling and behavioral therapies [53]. In addition, pregnant women who take prescription opioids also put their babies at risk of neonatal abstinence syndrome, similar to heroin [13].

Connection between opioid and heroin abuse

Transitioning from prescription opioids to heroin:

The misuse of prescription pain relievers leads some individuals to start heroin. According to the National Survey of Drug Use and Health (NSDUH) survey information from 2002-2011, nearly 80% of heroin users reported non-medical prescription pain reliever use prior to their heroin use. Here, non-medical prescription use is defined as taking prescriptions that were not prescribed to the user directly or used only for the feelings it causes. In fact, those who had prior non-medical prescription pain reliever use were 19 times more likely to initiate heroin use than those without prior use [38]. This could in part be due to the higher availability of heroin in recent years at a lower cost than alternative opioids [43]. Moreover, approximately 3.6 percent of non-medical prescription pain reliever users began using heroin within 5 years of their first opioid; although a small percentage, this is a significant number of individuals given the magnitude of opioid addiction [38]. In 2010, an abuse-deterrent formulation of the commonly abused prescription opioid OxyContin was released that made abuse through injection and inhalation more challenging.

Although done with the intent of reducing opioid abuse, studies showed that many individuals switched to heroin use instead [15, 16]. One study of young adults ages 18-25 concluded that among many factors including race, education status, marital status, other drug use, as well as many others, the use of non-prescribed opioid pain relievers in the past year was the biggest indicator of an individual using heroin in the past month, past year or in their lifetime [33]. Comparing NSDUH data from 2002-2004 and 2008-2010, the average yearly rates of past year heroin use increased among non-medical opioid users, but heroin use stayed static for those who reported no non-medical use of opioids; the highest rate of heroin use was among individuals with past year non-medical use of opioids ranging between 100 and 365 days of use [34]. In the 1960's, heroin users were composed mainly of younger, nonwhite men in urban areas with their initial opioid being heroin, but in recent decades, this trend has shifted to older, white, rural and suburban men and women with their initial opioid being a prescription [17]. Opioids are of no shortage in society today and since opioid addiction is driven largely by legal prescription medication availability, this has made a significant proportion of society susceptible to misuse and addiction to opioids, including heroin.

Heroin information

About the drug

Heroin is an illicit drug classified as an opioid and comes in the form of a white or brown powder or as a black substance resembling roofing tar. The drug is injected, sniffed, snorted or smoked and quickly enters the brain to bind to opioid receptors. It provides the user with feelings of euphoria, in addition to physical effects such as heavy feelings in the arms and legs, dry mouth, and sometimes nausea and vomiting [41, 40]. There are short and long term negative effects on the body for using the drug, and consequently, it is currently considered a schedule I drug, meaning that there is no approved medical use of heroin and there is a high likelihood for abuse. [60, 41]. Addicted individuals have withdrawal symptoms such as restlessness, diarrhea, vomiting, and cold flashes, along with cravings for the drug which makes stopping use of the drug very difficult [41]. Treatment options for heroin use are the same as for prescription opioids [53, 41]. Heroin users build up a tolerance to the drug with repeated use and can overdose on the drug, in which their heart rate and breathing is slowed to a dangerous level without medical assistance [40, 41]. Due to built-up tolerance, the risk of overdose is high when individuals stop their use of the drug for a period of time (i.e. while in recovery or hospitalized) and return to use. This is because they may return to the previous amount they were taking before, without knowing what their body can currently tolerate [18].

Other models

Previous models involving heroin

“Group” other models after 4 sentences of White and Comiskey (for paper). Say “other models include...” or “Approaches also include...agent-based modeling, etc.” Or “of interest is...” to cut down on length significantly (cut out all details for paper, keep for dissertation)

Look for fentanyl use/addiction models, if any

There have been several models formulated focusing on heroin addiction. Most of them

were motivated by and extensions of the White and Comiskey ordinary differential equation model with three compartments each representing a different stage as a drug-user: the susceptible class including individuals aged 15-64, the drug user class composed of individuals not in treatment, and finally, the drug users in treatment. In this model, individuals in treatment for drug use were only able to die, relapse to drug use, or complete treatment and be immune to drug use for the remainder of the modeling time period. The basic reproduction number, \mathcal{R}_0 , was calculated and deemed most sensitive to the rate of individuals in the susceptible class becoming drug users; therefore, prevention is more important than treatment for reducing drug use [70]. Wang, Yang and Li did analysis of this model, changing the assumption that the population was not constant [67].

The White and Comiskey model was modified later on by Liu and Zhang in order to incorporate a relapse distribution in which there was a non-constant time to relapse. They formulated a delay-differential equation, and also took away the assumption that the population was constant. Their conclusions about the sensitivity of \mathcal{R}_0 to certain parameters were in line with those from White and Comiskey [35]. Using the same three compartments, another model was formulated with the idea of the relapse rate relying on the length of time the individual has been undergoing treatment. This coupled ODE-PDE model assumed that susceptibles could only become addicted via interaction with heroin users not undergoing treatment. Prevention was deemed more important than treatment, yet again [22]. Fang, Li, Martcheva and Caiaiso (2015) also modeled the heroin epidemic allowing susceptibility of becoming a drug user to depend on age, again resulting in a coupled ODE-PDE model [23]. A non-autonomous version of the White and Comiskey model included a distributed time delay for becoming a drug user along with both the parameters and the total high-risk population size being time-dependent [48]. From this, an autonomous model was developed incorporating the distributed time delay, which was converted into a discrete model with the time step-size being one [2]. Finally, a three compartment model consisting of susceptibles, users not in treatment, and those in treatment, assumed that those not in treatment could not recover and return to being susceptible, and that it took two contacts with a drug user to become addicted, resulting in a nonlinear incidence rate; results showed that perhaps the heroin prevalence undergoes periodic changes depending on conditions [36].

A deviation from the three compartmental model was done that included a fourth compartment which consisted of individuals who successfully recovered from their heroin use, whether through deciding to stop use by themselves or via treatment measures [68]. The treatment of heroin users is limited by the availability of treatment facilities and resources and therefore, a saturated treatment function is incorporated; the saturation parameter was concluded as having the most vital role in the persistence of the epidemic. This implies that intervention is important early on before heroin addicts accumulate in the community and thus, prompt treatment is necessary. Another model consisting of 84 ODE's captured the dynamics between prescription opioid users with acute pain, those with chronic pain, illicit opioid users, heroin users and those who overdose. Results showed that increasing addiction treatment and reducing prescriptions for individuals with chronic pain seemed to have a greater impact on reducing both heroin overdose deaths and the number of opioid abusers compared to reducing prescriptions for acute pain without treatment increases [1]. This was from a conference preceding...in future, look for published model

from Benneyan.

Others have taken a completely different approach, specifically using agent-based modeling. In addition to their ODE model, Benneyan, Garrahan, Ilies, and Duan (2017) formulated two agent-based models: the first being a cellular automata model consisting of a grid with neighboring cells updated depending on state variables adjacent to the cells; the second being a network model with nodes that represent individuals/subpopulations and weighted pathways dependent on relational strength and nearness. **Not all variables are defined, such as $P(x,y)$, so look for full model from Benneyan when published.** The former was to better understand regional spread and the latter to investigate spread among individuals; their results suggested that with various regulation practices for opioids, heroin death rates would decrease, but the authors recognize the reality of unintended consequences of these legislations [1]. Agar (2001) created an agent-based model, consisting of a world with 2,500 patches and 100 agents each assigned an experimentation value which defines how likely that individual is to experiment with heroin, depending on their interest in trying something new, their attitude toward illicit drug use, and their desire to follow societal rules. The model was able to track the number of heroin experimenting individuals over time with the agent moving randomly, one patch selected at random to contain heroin in the beginning, and the ability for individuals to obtain heroin from their original patch, from others who have experimented, or from another patch where the transferring of heroin between agents occurred [3]. **Put in brief explanation of “Reducing the complexity of an agent-based local heroin market model” paper by Heard.**

Summary of Christopher’s opioid paper and why formulating another model:

Need to update with new revisions/any new results from final version that’s published!

The motivation for our heroin model comes from a previous model focusing on opioid addicts through prescriptions and via the black market [6]. In order for the model to have an addiction-free equilibrium, both addictions that come from prescriptions and addictions from accessibility to excess prescription drugs must be eliminated, which equates to the need for close administration and monitoring of those prescribed. Furthermore, near the addiction-free state, the prevention of prescription opioid users becoming addicted is more important for staying near the addiction-free equilibrium than reducing the number of prescriptions getting into the hands of non-prescribed users. **Fix wording:** However, away from the addiction-free state, a realistic situation, the most important factors in reducing the number of addicted individuals include increasing the prescription completion rate, increasing entry into treatment (even among low success rates), and decreasing the prescription rate. The purpose of formulating a separate model from this prescription opioid model is to be able to understand the more complicated dynamics that arise among opioid addiction with the addition of heroin/fentanyl use. **Make the case for the new model clearer: want to emphasize how we need new model because of fentanyl being the biggest killer now and with putting in a heroin/fentanyl class, this could lead to different management strategies. If need any more general information for Tennessee, great information in PDO 2018 report, and has fentanyl information, especially in executive summary!**

As exemplified previously, heroin and fentanyl play a significant role in the process of opi-

oid addiction and recovery and so it's inclusion provides a more accurate overall picture of the epidemic. **Read clarifying definitions again:** From 2000-2015, there were fewer heroin overdoses compared to prescription opioid overdoses but around 2010, a sharp increase of heroin overdoses occurred primarily due to the involvement of fentanyl. Therefore, it is crucial to take a more detailed look at this increasing problem [11]. **Regarding overdoses: study directly.** Moreover, we alter how the recovery process is viewed, by thinking about those in recovery still as addicts, and not having the ability to become a susceptible individual again.

How our model is different than previous models:

Include details on how my model different than previous ones.

Took out "heroin history," so need to incorporate argument regarding why we constructed a data-driven model versus a theoretical model as done in the past (i.e. use recent data changes as an argument, especially with increase in fentanyl. Previous models have not incorporated the connection between prescription opioid misuse and heroin or fentanyl use at all, instead focusing solely on heroin addiction and recovery. Although there are many factors that play a role in population size, susceptibility to drug use, relapse time and other parts of the drug-using process, we consider individuals in each of the classes to be homogeneous, for simplicity of a starting model. **discussion of how our model differs from the prescription opioid model: best to differentiate throughout the paper**

Provide argument/connect to info before for why the problem is so large now that we need data-driven models compared to just a theoretical heroin addiction model as done for decades.

Goal paragraph

Goals for our model/questions trying to answer:

To address this apparent problem in today's society, we model the prescription opioid/heroin/fentanyl epidemic in order to understand the dynamics behind the epidemic and predict the trajectory of the epidemic. This model analyzes the epidemic since it has been established, how the number of prescriptions have dropped each year after 2013, resulting in fewer pills lying around but leading to the use of other drugs such as heroin and fentanyl due to their lower cost and availability. This contrasts the opioid-only model in [6] in which the focus was on the dynamics of the epidemic as it was ramping up; we care about the sustained dynamics and investigate what do we do now that the epidemic is established. We identify important conditions relating to the reduction of opioid/heroin/fentanyl addiction. To do that, we have formulated a population level system of ordinary differential equations model consisting of classes of individuals taking prescription opioids, addicted to opioids, using heroin or fentanyl, and successfully recovered from addiction to opioids, heroin and/or fentanyl, and analyzed it. Our overall goal is to explore how different management strategies may alter the epidemic trajectory; specifically, we would like to investigate management strategies for optimally treating pain with prescriptions while reducing opioid, heroin, and fentanyl addiction.

Model Formulation

To the extent of our knowledge, there have been no mathematical models incorporating

both prescription opioid addiction and heroin/fentanyl addiction. Although terminology is not always clearly defined in the literature, such as opioid misuse, abuse, dependency, addiction and use disorder, we aim to focus on addiction only, where addiction to opioids is defined as having a pattern of continued non-medical use that is already, or could be, harmful [66]. We take opioid use disorder to fall in this categorization of addiction, due to the definition presented in [54] which includes sustained use regardless of interference with life obligations. Our model consists of five subgroups of a population, in which they are all proportions of the entire population:

1. Susceptibles S : This portion of the population consists of individuals who are not taking prescription opioids of any kind, nor using heroin or fentanyl.
2. Prescription opioid users P : This class of individuals consists of individuals who are prescribed opioids by a health care provider and take the opioids at a level that is not considered addicted. We note that individuals in this class could be misusing prescription opioids, but not at the level of addiction.
3. Opioid addicts A : This group of individuals are addicted to opioids, but not using heroin or fentanyl, or are actively in treatment for opioid addiction. For individuals who go to treatment for opioid addiction, they remain in this class for at least 4 weeks *after* being discharged from treatment; if they relapse within 4 weeks after treatment, they are again considered addicted and can remain in this class for longer. (Although the opioid class of drug includes heroin and fentanyl, here we take opioids to mean non-heroin and non-fentanyl.)
4. Heroin/fentanyl users H : This class is composed of individuals who are addicted to heroin or fentanyl, or are actively in treatment for heroin/fentanyl addiction. For individuals who go to treatment for heroin/fentanyl addiction, they remain in this class for at least 4 weeks *after* being discharged from treatment; if they relapse within 4 weeks after treatment, they are again considered addicted and can remain in this class for longer. (We note that individuals in this class could be using other drugs or are addicted to opioids in addition to heroin or fentanyl, but they are at least addicted to one of these drugs).
5. Recovered individuals R : This class consists of individuals who completed treatment for opioid or heroin/fentanyl addiction and did not relapse within 4 weeks after treatment, and therefore are considered in a “stable” recovery state.

Clarifications on definitions: We include illicit fentanyl users in the class of heroin users, because of the potency of the drug, which is what those addicted to prescription opioids seek—lower cost and a better high. In addition, a portion of those who overdose from the very powerful drug fentanyl are individuals who are using heroin that is laced with it; therefore, the use of fentanyl is oftentimes intertwined with the use of heroin. There is much variability in the purity of heroin due to the lacing of the drug with fentanyl, and 1 in 5 overdose deaths have multiple drugs present so it is difficult to know the actual cause of death [11]. We believe it is difficult to separate the use of heroin and fentanyl, so putting them in the same class is the most natural choice. Finally, we remark that use of heroin is implicitly understood to be addictive due to the highly addictive nature of the

drug, and we assume no casual use of the drug [41]. If an individual uses heroin at a level that it would be easy to stop using, they are better classified as a susceptible individual rather than a heroin addict, based on the definition of addiction we are using. Similarly, individuals who use fentanyl outside of short-term, carefully monitored settings with a health provider is assumed to be addicted. **Source for fentanyl addictive nature, length of time of prescription short?** We try to take into account these complexities as best as possible with the information in literature.

We denote the initial conditions as $S(0) = S_0$, $P(0) = P_0$, $A(0) = A_0$, $H(0) = H_0$, and $R(0) = R_0$, and we assume all of these values are positive. The initial values for each of the classes are proportions of the entire population **and therefore are dimension-less...add this to parameter/unit table in appendix.**

Here, we present our ordinary differential equation model:

$$\frac{dS}{dt} = -\alpha S - \beta_A SA - \beta_P SP - \theta_1 SH + \epsilon P + \mu(S + P + R + A + H) + \mu_A A + \mu_H H - \mu S \quad (1)$$

$$\frac{dP}{dt} = \alpha S - \epsilon P - \gamma P - \theta_2 PH - \mu P \quad (2)$$

$$\frac{dA}{dt} = \gamma P + \sigma R \frac{A}{A + H + \omega} + \beta_A SA + \beta_P SP - \zeta A - \theta_3 AH - (\mu + \mu_A) A \quad (3)$$

$$\frac{dH}{dt} = \theta_1 SH + \theta_2 PH + \theta_3 AH + \sigma R \frac{H}{A + H + \omega} - \nu H - (\mu + \mu_H) H \quad (4)$$

$$\frac{dR}{dt} = \zeta A + \nu H - \sigma R \frac{A}{A + H + \omega} - \sigma R \frac{H}{A + H + \omega} - \mu R \quad (5)$$

$$S + P + A + H + R = 1 \quad (6).$$

See if best way to write this last equation?

Why can we assume this sum is 1?...short time frame, population constant; we are imposing because population not changing so much (i.e. we made the model to be this way/work out this way mathematically) We assume that these five compartments sum to one, so that the population is of constant size, i.e. the total death rate is equal to the incoming rate for the susceptible class. Since these rates are the same, μS , equation (1) simplifies to:

$$\frac{dS}{dt} = -\alpha S - \beta_A SA - \beta_P SP - \theta_1 SH + \epsilon P + \mu(P + R) + (\mu + \mu_A) A + (\mu + \mu_H) H \quad (1)$$

The parameters involved in this model represent transition rates from one class to another and are per capita yearly rates:

- αS : rate at which susceptible individuals are prescribed opioids (1/year) **May change to be clearer in each of these parameters about the units (i.e. per prescription user**

year or whatever it is) or may just leave that for the parameter table where units are listed

- $\beta_A SA$: rate at which susceptible individuals become addicted to opioids by black market drugs or interaction with other addicts (1/year)
- $\beta_P SP$: rate at which susceptible population obtains extra prescription opioids and becomes addicted per year
- $\theta_1 SH$: rate at which susceptible population becomes addicted to heroin or fentanyl by black market availability or interaction with other heroin or fentanyl users (1/year)
- ϵP : rate at which individuals return to the susceptible class after being prescribed opioids and did not develop an addiction (1/year)
- $\mu S, \mu P, \mu A, \mu H, \mu R$: natural death rates (1/year)
- $\mu_A A$: enhanced death rate for opioid addicts; overdose rate which results in death (1/year)
- $\mu_H H$: enhanced death rate for heroin or fentanyl addicts; overdose rate which results in death (1/year)
- γP : rate at which prescribed opioid users become addicted to opioids (1/year)
- $\theta_2 PH$: rate at which prescribed opioid users become addicted to heroin or fentanyl (1/year)
- $\sigma R \frac{A}{A+H+\omega}$: rate at which individuals transition from the recovered class into the opioid addicted class which is proportional to the approximate proportion of individuals in the recovered class that came from A that are relapsing back to that class. We include a perturbation term ω here in the case that A and H are both 0, such as in the addiction-free case. (note: even though we do not know where individuals came from originally, we make the assumption that individuals go back to their primary drug of choice/the reason they entered treatment, so relapse back approximately proportional to where they came from **Source/explanation below..also, fix wording when take care of this depending on what we go with. Also mention that this is an approximation because it's based on the "current time" class proportions (not the time step when they went to R). We are doing this because the percentages/stats we are using are "relapse rates" so that means individuals are going back where they started.**) (1/year)
- ζA : rate at which addicted opioid users enter treatment/rehabilitation (1/year)
- $\theta_3 AH$: rate at which the opioid addicted population becomes addicted to heroin or fentanyl (1/year)
- $\sigma R \frac{H}{A+H+\omega}$: rate at which individuals transition from the recovered class into the heroin/fentanyl addicted class which is proportional to the approximate proportion of individuals in the recovered class that came from A that are relapsing back to

that class. We include a perturbation term ω here in the case that A and H are both 0, such as in the addiction-free case. (note: even though we do not know where individuals came from originally, we make the assumption that individuals go back to their primary drug of choice/the reason they entered treatment, so relapse back approximately proportional to where they came from **Source below..also, fix wording when take care of this.**) (1/year)

- νH : rate at which heroin or fentanyl users enter treatment/rehabilitation

Clarification of some aspects of the model:

We note that although sellers are not directly involved in the model nor have to be addicts themselves, addicted individuals act as a proxy for sellers of the drug and general availability of the drug, whether it is opioids or heroin. This is because the higher number of addicted individuals there are, the more sellers there needs to be which results in an increased exposure to the drug and therefore, increased chance of addiction. Although treatments for addictions are an enormous problem in and of themselves since there is limited success as shown by the parameter values for the recovery rates, we are acknowledging treatments as a part of recovery in our model, but do not go into specifics for this first model. We are classifying overdoses to mean opioid overdose-related deaths (i.e. not from secondary factors or long term effects). Although overdose death data includes both accidental and non-accidental deaths (i.e. suicide, assault), we assume here that if an individual died from an overdose-related death, they were addicted **see calculation in appendix; I don't think we assume this anymore from [7] information** (i.e. the number of intentional deaths is negligible). **This is CDC definition (from data packet) is for prescription opioids and fentanyl; see if also same for heroin.** We also note the fact that Naloxone, an overdose-reversing medication, has saved individuals from overdoses, but **[40] make some argument that the number saved is negligible compared to the number of overdoses, so we assume results in death most of the time.**

Explanation of different R class, how different from Christopher's opioid paper:

Fix: don't want to use "negative" language such as "we did something to *avoid* something else," etc. Reword so saying something like "Other choices for dealing with recovery include..., two recovery classes, etc..."

We note that the terms $\sigma R_{\frac{A}{A+H+\omega}}$ and $\sigma R_{\frac{H}{A+H+\omega}}$ were included in order to avoid having two recovered classes, which would prevent the transition of individuals who recovered from one drug to initiate use of the other (or if we included those pathways, would become too complicated for a first model). One study suggested that 10% of the prescription opioid addicts they followed up with post-treatment had used heroin at least five times in the previous year, which demonstrates the importance of keeping these transitions as possibilities [69]. We decided to define a recovered class instead of a recovery class as the model presented in [6] did, because the recovery class would have consisted of two very different groups of individuals: those with a high chance of relapse (e.g. recently finished treatment) and those with a much lower chance of relapse (e.g. successfully finished treatment in the past and has remained free of addiction for a significant amount of time). This definition of the recovered class creates a compartment to place individuals who have been addicted in the past and have finished treatment but are not considered actively

addicted anymore and should be dealt with differently than susceptible individuals, those in treatment, and those recently out of treatment.

We note here that the recovered class and relapse from that class is one of the key features that distinguishes our model from that of [6]. In the opioid-only model, as the focus was on the dynamics of the epidemic as it was ramping up, at beginning of the epidemic, relapse did not depend on the number that were addicted. In contrast, since the epidemic has been established, there are high numbers of opioid and heroin/fentanyl addicts; the proportion in each category affects where people relapse to. **Be clearer with words here, depending on if interaction term or not; may need to alter argument about primary drug/source-driven?**

Support/data/arguments for why choosing 4 weeks/acute withdrawal period

Summarize these multiple supporting sources into a few sentences and only go into this much detail in disseration? We choose to use 4 weeks after treatment with no relapse as the mark of when people are “successfully recovered” and can move to R. We chose this time frame due to several studies exhibiting a high level of relapse within weeks of discharge from treatment, which suggests individuals are not stably recovered. One study following up with opiate dependent individuals (of which 88% of the study population were heroin users) showed that 59% of the individuals discharged from treatment relapsed within 1 week and 71% within 4 weeks [49]. Moreover, 80% of individuals who relapsed after treatment did so within 4 weeks and that 92% of individuals who relapsed after discharge had gone back to treatment prior to their follow-up interview (which was between 18-42 months). These statistics convey the unstable nature of an addicts’ recovery, particularly within a few weeks of post-treatment (since that’s when a majority of them relapsed). In another opiate dependent group study evaluating their relapse risk, 27% had relapsed on the discharge day of their most recent treatment program, 41% had relapsed within 1 week and 65% within 4 weeks [5]. The only study found specifically for prescription opioid addicts suggested that 91% of them who completed a two-phase treatment program relapsed back into addiction within 8 weeks post-treatment [69]. On the lower side, in another study an estimated 32% of the 68 opiate addicts interviewed returned to addiction within 4 weeks and 65% lapsed, which we note has potential to lead to dependence in time; at the 6 month follow-up, 50% of the 60 individuals interviewed were dependent [9]. An older study following up with opiate users post-treatment for 6 months showed that 32% had lapsed within 1 week post-discharge when interviewed; 71% had lapsed by the 6th week; and 44% had returned to the daily use of opiates (considered addicted) at the 2 month mark [25]. Furthermore, the study stated that several individuals who completed the withdrawal portion of the program lapsed while in the second, in-patient part of treatment (although not necessarily equating to a full relapse back into addiction) which supports our claim that those who entered treatment addicted should be still be considered addicted while in treatment. Later analysis of this study regarding relapse factors stated that 25% of those who lapsed mentioned withdrawal symptoms as playing a role in their lapses [8]. These statistics suggest that opiate-abstinence can be difficult to achieve in this initial time period post-treatment: “...the period immediately after leaving a residential treatment is of massively high risk: the great majority of lapses occurred with the first few weeks after discharge...after the first four weeks, there were few additional lapses...first few weeks after discharge as a *critical period* in the process of recovery” [25].

This suggests four weeks can be viewed as an important marker in recovery, and that individuals who bounce back to addiction/frequently relapse in this time frame should simply remain in their respective addicted classes. Other studies suggest significant lapse and relapse rates within weeks and months for heroin addicts and opiate addicts, as well [32, 26].

Therefore, since 71% of heroin addicts relapse within 4 weeks post-treatment and an estimated 91% of prescription opioid addicts in recovery relapse back to addiction within 8 weeks post-treatment (the majority most likely within four weeks based on these other studies), we keep these individuals in the addiction class up for 4 weeks post-treatment since they have not stably “recovered,” i.e. at a point where they are less likely to fall back into addiction [49, 69].

Information on relapse coming from primary drug and not source-driven, argument for form of our transition terms being proportional to number in each addicted class

In one study, 63% of initial lapses occurred around other opioid users, 48% in another addicts’ home, and 31% in ones own home [25]. This may lead one to believe that interactions with other addicts would play a large role with an initial lapse/relapse; however, following this study, the authors explored the circumstances that led to the lapses of these study subjects and results proved differently. The top factors that led to these initial lapses were cognitive factors (planned to use), mood states (angry, sad, lonely, etc.), and external influences (situations unrelated to drug use that led to use). This was taking into consideration the number of people who mentioned the factors, the total number of times the factors were mentioned, the number of individuals who deemed the factor most important, and finally, the number of individuals that indicated the factor was the initiator of a sequence of other factors, as lapse often is a result of multiple factors occurring simultaneously. These were also the factors that were primarily contributed to the continued use of opiates. It seems clear that source-driven factors, specifically social pressure (offered drugs), drug availability (in another users’ home or in the area), drug-related clues (objects used for drug-taking/observing others under the influence), and interpersonal influences (seeing or thinking about certain people) had relatively little importance compared to the others mentioned above [8]. It is because of this that we argue individuals return to their primary drug of choice compared to what is simply available; although we are interested specifically in a return to addiction, that is umbrellaed under the broader category of lapses.

We mention that another study specifically on heroin users enabled the study subjects to choose categories they deemed as most important in their relapse according to a phrase attached to the category and this was compared to the results of independent judges selecting the categories based on individuals’ responses. The highest mean ratings for the categories that the subjects chose were for the giving into temptations/urges in the presence of substance cues and direct social pressure, whereas the judges deemed negative emotional states and indirect social pressure as the most frequent reasons. However, given that subjects were provided a phrase that correlated with a certain category, one could argue that if an individual did not agree with the exact phrase, they may have chosen an alternative option that they related to more (e.g. “I felt bored” represented the entire category of “negative emotional states other than frustration and/or anger.”). Moreover, the previous study applies both to opioid addicts and heroin addicts, rather than just heroin addicts [28].

Explanation of pathways omitted:

We omitted an interaction term between prescribed users and addicted individuals moving into the addicted class under the assumption that their own prescription is the primary source of their addiction, and that an illicit source or interaction with addicts would be secondary. Moreover, we did not include an interaction term RA to go from the recovered class to the opioid addicted class because we assume that individuals who are in a stable recovery state are not influenced by addicts or by the general availability of the drug to fall back into addiction themselves; similarly for RH. **If keep this, can refer to opioid paper.** **Currently working on these terms/ideas, so will most likely be updated.** We do not have a pathway from R to P; although individuals have the potential of obtaining a legal prescription from a doctor since doctors are not necessarily informed that their patient was in recovery in the past **find resource that talks about doctors being “case-history blind”**, we believe it is not realistic that an individual who has recovered from an opioid or heroin addiction would be able to successfully take prescription opioids in a non-addictive manner, and therefore would be in the addicted class almost immediately. Although there may be a few exceptions to this, we make this simplifying assumption for a first model.

We omitted the pathway from R to S since individuals who have been to treatment and recovered are much more likely to fall into addiction than a susceptible individual, and therefore, must be treated differently than a susceptible individual who has not been to recovery (i.e. cannot be homogeneously mixed in S). We think of individuals in the recovered class as in a permanent state of recovery for the rest of their lives, although they are not undergoing active recovery, and we think of this class as an absorbing state. **Find source talking about addicted individuals who underwent recovery much more likely to fall back into addiction than susceptible individual...addiction being a disease. This is a change from the opioid model; discuss dynamical difference.** Along these same lines, there is no pathway from A or H to S because we are under the assumption that an addicted individual either remains addicted or goes to recovery to help their addiction and will never be characterized as a susceptible individual again. We recognize that there may be additional pathways among the classes, but the ones we have chosen to include in this first model are a result of balancing simplicity and robustness.

This compartmental model can be represented by the following flow diagram with each arrow representing either the transition rate between one class to another or birth/death:



Figure 1: Schematic diagram for heroin model

Analysis of the Model

Addiction-Free Equilibrium

To find the addiction-free equilibrium, we set equations (1)-(5) equal to zero and require that $A = H = 0$ since these two compartments consist of individuals who are addicted to opioids or heroin. We are left with the system:

$$0 = -\alpha S^* - \beta_P S^* P^* + \epsilon P^* + \mu(P^* + R^*) \quad (7)$$

$$0 = \alpha S^* - \epsilon P^* - \gamma P^* - \mu P^* \quad (8)$$

$$0 = \gamma P^* + \beta_P S^* P^*. \quad (9)$$

$$0 = \mu R^* \quad (10)$$

$$1 = S^* + P^* + A^* + H^* + R^* \quad (11)$$

Equation (10) forces either $\mu = 0$ or $R^* = 0$; however, $\mu > 0$ since this is the natural death rate, so we require $R^* = 0$. If $P^* \neq 0$, this forces $\gamma + \beta_P S^* = 0$ from equation (9), and since all of our parameters and variables are non-negative, then it must be $\gamma = 0$ and $\beta_P = 0$. We note that $\gamma = 0$ means that individuals who are prescribed opioids cannot become addicted to opioids, and $\beta_P = 0$ means that only black market opioids are available for susceptibles to become addicted to opioids and there are no excess prescription drugs available. Under the assumption that $\gamma = 0 = \beta_P$ to ensure the existence of our addiction-free equilibrium, we calculate the addiction-free equilibrium to be:

$$\begin{aligned}
S^* &= \frac{\epsilon + \mu}{\alpha + \epsilon + \mu} \\
P^* &= \frac{\alpha}{\alpha + \epsilon + \mu} \\
A^* &= 0 \\
H^* &= 0 \\
R^* &= 0
\end{aligned}$$

If $P^* = 0$, then equation (8) forces either $S^* = 0$ or $\alpha = 0$. If $S^* = 0$, then the solution $S^* = P^* = A^* = H^* = R^* = 0$ contradicts equation (11). If $\alpha = 0$, then we use equation (11) to solve for the addiction-free equilibrium of (1,0,0,0,0); this solution is included in the addiction-free equilibrium above when $\alpha = 0$, as well.

Basic Reproduction Number, \mathcal{R}_0

The basic reproduction number, denoted \mathcal{R}_0 , is a term used in epidemiological models that gives the expected number of secondary disease cases that result from the introduction of a disease to a susceptible population. The value of \mathcal{R}_0 represents how successful the spread of the disease is expected to be; if $\mathcal{R}_0 < 1$, then the disease is expected to die out and the disease-free equilibrium will be locally stable; conversely, if $\mathcal{R}_0 > 1$, then the disease is expected to spread and the disease-free equilibrium will be unstable [64].

This idea may be applied to the context of our model since for the addiction-free equilibrium to occur, γ and β_P must both be equal to 0, which means individuals can become addicted only with interactions with opioid addicted individuals or heroin users so this takes the form of an infectious disease. Thus, in this case in which the prescription part is essentially taken out of the model so that it is a black-market-only driven model, \mathcal{R}_0 can be calculated. This value will be the ratio of the number of new addictions in the next year compared to the current year. At the addiction-free equilibrium, the population is completely susceptible, as needed for \mathcal{R}_0 to be an accurate measurement of addiction potential. We note that our goal in this section is to explore the structure of the model and not obtain a result for the full model. The disease compartments are those that contain infected individuals, or in our case, addicts [64]. Our model incorporates two addiction compartments, A and H, since these both consist of opioid or heroin/fentanyl addicted individuals. We will utilize the Next Generation Matrix Method in order to calculate \mathcal{R}_0 .

For the purposes of calculating \mathcal{R}_0 , we will assume $\gamma = 0$ and $\beta_P = 0$ in order to ensure the existence of the addiction-free equilibrium. This results in the following system:

$$\begin{aligned}
\frac{dS}{dt} &= -\alpha S - \beta_A SA - \theta_1 SH + \epsilon P + \mu(P + R) + (\mu + \mu_A)A + (\mu + \mu_H)H \\
\frac{dP}{dt} &= \alpha S - \epsilon P - \theta_2 PH - \mu P \\
\frac{dA}{dt} &= \sigma R \frac{A}{A + H + \omega} + \beta_A SA - \zeta A - \theta_3 AH - (\mu + \mu_A)A
\end{aligned}$$

$$\begin{aligned}\frac{dH}{dt} &= \theta_1 SH + \theta_2 PH + \theta_3 AH + \sigma R \frac{H}{A+H+\omega} - \nu H - (\mu + \mu_H)H \\ \frac{dR}{dt} &= \zeta A + \nu H - \sigma R \frac{A}{A+H+\omega} - \sigma R \frac{H}{A+H+\omega} - \mu R.\end{aligned}$$

In general, the differential equations of the n disease compartments, x'_i , may be written as:

$$x'_i = \mathcal{F}_i(x, y) - \mathcal{V}_i(x, y), \quad i = 1, \dots, n$$

where x is a vector with the disease compartments as components, y a vector with the non-disease compartments as components, \mathcal{F}_i represents the rate that secondary infections contribute to disease compartment i and \mathcal{V}_i represents the rate of transitions, i.e. rate at which the disease compartment i is decreased by means of death, recovery and progression of the disease for $i = 1, 2$ [64].

Thus, for our two addicted compartments, we may write:

$$\begin{aligned}\frac{dA}{dt} &= \mathcal{F}_1(x, y) - \mathcal{V}_1(x, y) \\ \frac{dH}{dt} &= \mathcal{F}_2(x, y) - \mathcal{V}_2(x, y)\end{aligned}$$

where $x = [A \quad H]^T$ and $y = [S \quad P \quad R]^T$.

Thus, under the assumption that A and H are the addicted compartments and abiding by the parameter restrictions stated above, the assumptions of the Next Generation Method are satisfied for matrices \mathcal{F} and \mathcal{V} formulated here:

$$\begin{aligned}\mathcal{F} &= \begin{pmatrix} \sigma R \frac{A}{A+H+\omega} + \beta_A S A \\ \theta_1 S H + \theta_2 P H + \sigma R \frac{H}{A+H+\omega} \end{pmatrix} \\ \mathcal{V} &= \begin{pmatrix} \zeta A + \theta_3 A H + (\mu + \mu_A) A \\ -\theta_3 A H + \nu H + (\mu + \mu_H) H \end{pmatrix}.\end{aligned}$$

These assumptions include the following:

- $\mathcal{F}_i(0, y) = 0$ and $\mathcal{V}_i(0, y) = 0$, $\forall y \geq 0$, $i = 1, 2$, which ensures that new addictions arise only from interacting with those currently addicted, and there is no immigration into the addicted compartments; these guarantee that the population free of addiction remains that way.

- $\mathcal{F}_i(x, y) \geq 0, \forall x, y \geq 0, i = 1, 2$ since these represent new additions.
- $\mathcal{V}_i(x, y) \leq 0$ whenever $x_i = 0, i = 1, 2$ since there must be inflow only when the respective compartment is empty.
- $\sum_{i=1}^2 \mathcal{V}_i(x, y) \geq 0, \forall x, y \geq 0$ since this represents the total outflow from all addicted compartments.
- The addiction-free system, $\left. \frac{dS}{dt} \right|_{A,H=0}, \left. \frac{dP}{dt} \right|_{A,H=0}$, and $\left. \frac{dR}{dt} \right|_{A,H=0}$ has a unique equilibrium, $(\frac{\epsilon+\mu}{\alpha+\epsilon+\mu}, \frac{\alpha}{\alpha+\epsilon+\mu}, 0)$ that is asymptotically stable.

Taking $F = \frac{\partial \mathcal{F}_i}{\partial x_j}(0, y_0)$ and $V = \frac{\partial \mathcal{V}_i}{\partial x_j}(0, y_0), i, j = 1, 2$, where $(0, y_0) = (\frac{\epsilon+\mu}{\alpha+\epsilon+\mu}, \frac{\alpha}{\alpha+\epsilon+\mu}, 0, 0, 0)$ is the addiction-free equilibrium, we calculated the following [64]:

$$F = \begin{pmatrix} \beta_A \frac{\epsilon+\mu}{\alpha+\epsilon+\mu} & 0 \\ 0 & \theta_1 \frac{\epsilon+\mu}{\alpha+\epsilon+\mu} + \theta_2 \frac{\alpha}{\alpha+\epsilon+\mu} \end{pmatrix}$$

$$V = \begin{pmatrix} \zeta + \mu + \mu_A & 0 \\ 0 & \nu + \mu + \mu_H \end{pmatrix}.$$

The eigenvalues of FV^{-1} are calculated to be:

$$\sigma(FV^{-1}) = \left\{ \frac{\beta_A(\epsilon+\mu)}{(\alpha+\epsilon+\mu)(\zeta+\mu+\mu_A)}, \frac{\theta_1(\epsilon+\mu)+\theta_2\alpha}{(\alpha+\epsilon+\mu)(\nu+\mu+\mu_H)} \right\}$$

\mathcal{R}_0 may then be determined as the spectral radius of FV^{-1} , in which the (i, j) entry is the expected number of secondary additions in compartment i produced by individuals initially in compartment j :

$$\mathcal{R}_0 = \text{WHICHEVER IS LARGEST WHEN KNOW VALUES!}$$

Include proof of non-negative IC implies non-negative solutions?
What other analysis should be done?

Numerical Results (literature information and parameter estimation)

Data used within code

Applying model to Tennessee population

This is where start to talk about Tennessee

We apply our model to the Tennessee population specifically in order to analyze the opioid epidemic on a smaller scale than the national level. Although there is still heterogeneity across Tennessee, it is lower than the amount one would find on a national scale. Below we present a data table consisting of estimates of the number of individuals in Tennessee in various categories for the years 2013-2018 found either in literature or that we estimated; explanations of each of the categories follows.

Leave data table/information here or put in appendix? Put in information of US prescription rate per capita at it's peak was 81.3/100 people and how TN has been far above that every year.

Table 1: Number of individuals in each category, 2013-2018

	2013	2014	2015	2016	2017	2018
Total population	6,490,795	6,540,007	6,590,726	6,649,404	6,715,984	6,770,010
Population 12 and older	<i>5,517,176</i>	<i>5,559,006</i>	<i>5,602,117</i>	<i>5,651,993</i>	<i>5,708,586</i>	<i>5,759,999</i>
Heroin users	-	14,000	14,000	19,000	-	-
Heroin addicts	-	7,560	7,560	10,260	-	-
Prescription opioid addicts (includes heroin addicts)	-	-	48,000	42,000	-	-
Prescription opioid addicts (excludes heroin)	<i>43,418</i>	<i>42,928</i>	42,816	37,464	<i>34,816</i>	<i>32,816</i>
Prescribed opioid users (includes those addicted)	1,845,144	1,824,342	1,819,581	1,761,363	1,636,374	1,517,374
Prescribed opioid users (excludes those addicted)	<i>1,825,910</i>	<i>1,805,325</i>	1,800,613	1,744,766	<i>1,620,951</i>	<i>1,500,951</i>
Prescription opioid overdose deaths	-	-	679	-	-	-
Heroin/fentanyl overdose deaths	-	-	374	-	-	-
Prescription opioid treatment admissions	4,485	4,530	4,326	-	-	-
Heroin treatment admissions	555	743	1,083	-	-	-

Explanations of the estimates/where they come from in literature/assumptions made

Some of these are data points that are used directly in parameter estimation code, but some are used for calculation of μ , μ_A , μ_H , IC,...leave these explanations here or put them in the appendix?

We note that *italicized* values are numbers that we estimated by extrapolating from a different year, **bolded** values are numbers that we estimated by information within the same year, the remaining numbers are actual data and the numbers in blue are data used directly in parameter estimation.

The first row provides the **total population estimates in Tennessee** each year [59]. However, since the remainder of the data we use is specific to individuals 12 and older, we estimate the number in this age group. There were an estimated 1,073,214 individuals in Tennessee in 2018 aged 12 and under. (Note: we could not find the estimate for this age group for the years 2013-2017, so this is the best estimate we have for this age group). To figure out those who are 12 years old, we take approximately 1/8th of the individuals that are in the age group 5-12, which is approximately 83,175 individuals [19]. Thus, an estimated 990,039 individuals are *under* the age of 12 in Tennessee. Given the total population estimate for 2018 being 6,770,010 from row 1, this means that approximately 15% of the population is under the age of 12. Since we do not see a reason for this percentage to be significantly different from year to year, we assume that this percentage is constant throughout the time period we are looking

at. Then, we are able to consider the **Tennessee population estimates for individuals 12 and older** in order to align with the rest of the data that is in this age range by taking off 15% of the above total population estimates; this is shown in the second row.

The **total number of individuals taking prescription opioids for pain (prescribed)** is given in the third row [44]. Although this number does not explicitly state it is for individuals 12 and older, we assume it is since it comes from the Tennessee Department of Health; if it does include individuals under 12, we assume that number is negligible.

The 2015/2016 average number of individuals 12 and older with “Pain Reliever Use Disorder” (**prescription opioid addicts**) is 48,000 and the 2016/2017 average number was 42,000 [51, 52]. For these estimates, we choose the lower of the two years to place the data points; for example, for the 2015/2016 average, 48,000 is placed in the year 2015. *May need to fill in more details about specific data such as collection process, self-reported probably equates to underreported, etc.* We note that their definition of pain reliever use disorder includes those who meet the American Psychiatric Association criteria for dependence or abuse. Here, opioid dependence is classified as having “signs and symptoms that reflect compulsive, prolonged self-administration of opioid substances that are used for no legitimate medical purpose or...are used in doses that are greatly in excess of the amount needed for pain relief...regular patterns of compulsive drug use that daily activities are typically planned around obtaining and administering opioids.” This definition falls under our definition of opioid addiction. Opioid abuse, on the other hand, they consider to be less severe than dependence, and would not lead to the development of withdrawal symptoms. This latter definition does not fall under our characterization of addiction, but we make a note of this to say that this estimate for those with a pain reliever use disorder may be overestimated for what we are concerned with, but is an acceptable approximation [4]. In addition, we assume very few, if any, individuals are addicted to opioids outside of prescription opioids, but even if they are, we assume they are taking prescription opioids in some capacity, so we take pain reliever use disorder to include those with other non-prescription opioid addiction. We also remind the reader that these numbers include both individuals who are personally prescribed opioids or may obtain them illicitly. *May find a better place for this later: Finally, we note that those who are addicted to opioids are strictly in the A class, as individuals in R are not considered addicted in our model.*

The 2014/2015 average number of individuals 12 and older with “Past Year Heroin Use” (**heroin/fentanyl addicts**) was 14,000, the 2015/2016 average number was 14,000, and the 2016/2017 average number was 19,000 [50, 51, 52]. This number includes those who may have used heroin once or twice in the past year; however, it is estimated that 54% of heroin users are dependent on the drug, which gives the resulting **bold** estimates in the table [39]. In addition, the number of heroin users does not include fentanyl users explicitly, but we are under the assumption that those who take fentanyl are a subset of those who use heroin, and therefore, would mostly be included in these numbers. We admit the values may be slightly too low, for the cases of individuals who do fentanyl and not heroin, but data has not been found for fentanyl addicts only. Therefore, we are working under the assumption that it would be a negligible population that takes

fentanyl without heroin. Again, for these estimates, we choose the lower of the two years to place the data points; for example, 10,260 is placed in the year 2016. The values **bolded** in Table 1 are because numbers were based on the data of individuals from the same year, and did not rely on any extrapolated data from a different year.

In 2010, on the national level it was estimated that 14% of individuals who abused or were dependent on pain medications also used heroin [39]. It was right around this year that the number of overdose deaths due to heroin increased and continued to increase until 2016 [44, 11]. Although fentanyl began being laced with heroin in this time frame, the overdose deaths due to fentanyl were recorded separately and heroin overdose deaths were still increasing despite that; therefore, it seems likely that an increase in overdose deaths would equate to a larger number of heroin addicts in each of those years. Therefore, we assume a higher percentage that 20% of prescription opioid addicts used heroin each of the years 2013-2017. However, only 54% of heroin users are addicts, which means that 10.8% of prescription opioid addicts are also heroin addicts ($.54 \times .2 = .108$) [39]. We subtract these heroin addicts from the prescription opioid addict category each year; this results in the non-heroin addicted prescription opioid addicts.

From 2013 to 2017, the number of prescribed opioid users was decreasing, likely due to more awareness of the problem as suggested in [14] which resulted in a decreasing prescribing rate [45]; thus, we expect the number of addicts to also be decreasing in that time period, especially due to the movement of individuals to heroin use. For 2015, we see there were 42,816 (non-heroin addicted) prescription opioid addicts compared to 1,819,581 prescribed opioid users, which is an approximate ratio of 1:42.5; thus, we use this exact ratio of addicts to prescribed opioids users for 2013 and 2014, as well, to estimate the number of opioid addicts (these values are in *italics* in Table 1 since they are extrapolated from a different year). Although the numbers are most likely higher than this, this is an estimate. We apply a similar idea for the year 2017, utilizing data from 2016 that there are 37,464 (non-heroin addicted) prescription opioid addicts out of 1,761,363 prescription users, which is an approximate ratio of 1:47 (also *italicized* in Table 1).

The total number of prescribed opioid users each year, however, includes individuals addicted to their prescriptions. In a 2015 study done among adults with prescription opioid use disorder, 44.3% of them had obtained prescription opioids for their most recent episode of misuse from 1 or more physicians [27]. We make the assumptions that individuals who are addicted to heroin are not using *prescribed* opioids, since they are more expensive and provide less of a high; although this study consisted of individuals with prescription opioid use disorder, regardless of their heroin addiction status, we will extrapolate this percentage to those who are non-heroin addicted[43].. We recognize this percentage may be on the low side (i.e. if heroin addicts were taken out of the study, this percentage may be higher since those non-heroin addicted would possibly obtain more from a physician source), but we take this as our best estimate. Therefore, we assume that 44.3 out of 100 non-heroin addicted prescription opioid addicts are personally prescribed opioids and subtract 44.3% of the number of non-heroin prescription opioid addicts in each year from the prescribed opioid class in the corresponding year to obtain the number of non-addicted prescribed opioid users for 2013-2017.

We use data on the number of prescription opioid overdose deaths which include natural, semi-synthetic, and synthetic opioids; however, we subtract out the number of fentanyl overdoses (fentanyl is classified as a synthetic prescription opioid), since those overdoses are counted for in their own category, listed below. This results in $(848-169=)$ 679 as the **total number of prescription opioid overdose deaths** for 2015. [45]. Although this number does not explicitly state it is for individuals 12 and older, we assume it is since it comes from the Tennessee Department of Health; if it does include individuals under 12, we assume that number is negligible.

We add together the heroin and fentanyl overdoses for the state of Tennessee to obtain the **total number of heroin and fentanyl overdoses** in 2015 to be $(205+169=)$ 374 [45]. Although this number does not explicitly state it is for individuals 12 and older, we assume it is since it comes from the Tennessee Department of Health; if it does include individuals under 12, we assume that number is negligible.

For Tennessee individuals 12 and older, **the number of treatment admissions for non-heroin opiates/synthetics** as the primary substance of abuse to facilities that receive state/public funding (generally referring to funding by the state substance abuse agency) is found in row eight of Table 1 [20]. Here, we take non-heroin opiates/synthetics to mean prescription opioids. We are under the assumption that if one were addicted to heroin in addition to prescription opioids, their heroin problem would be the primary reason for going to treatment and would be included in the following numbers. Finally, row nine of Table 1 gives the **number of treatment admissions** for heroin as the primary substance of abuse to facilities that receive state/public funding (generally referring to funding by the state substance abuse agency) for Tennessee individuals 12 and older [20]. Again, these numbers do not include fentanyl users explicitly, but we are under the assumption that those who take fentanyl are a subset of those who use heroin, and therefore, would mostly be included in these numbers. We admit the values may be slightly too low, for the cases of individuals who do go to treatment with the primary substance of abuse being fentanyl, but there is not data available for those numbers currently. Calculations of parameters and estimates of initial conditions from this data is found in Appendix A.

Parameter estimation process

[Edit from/update from /combine with information from data_using_parameter_estimation file in heroin model multistart folder](#) The age-adjusted death rate (μ) and overdose death rates (μ_A and μ_H) are calculated and assumptions on θ_2 , θ_3 , β_A , and β_P are stated in Appendix A. In addition, although we have an estimate for the total number of prescription opioid users and opioid addicts in 2013, we do not know these numbers at the start of 2013; similarly for the other states. Thus, we approximate the initial conditions of our states for 2013 in Appendix A using information from Table 1.

Parameter (rate)	Value Assumed	Units
β_A		
β_P		
θ_2		
θ_3		
P_0		
A_0		
H_0		
R_0		

In order to estimate the remaining parameters for our model, we utilized the ordinary least squares method and formulated an objective function to minimize, which consists of the squared differences between data and model simulations. We have data on the total number of non-addicted prescription users, opioid addicts, and heroin/fentanyl addicts each year, and therefore on their corresponding proportions out of the entire population in a given year. We have data on the proportion of individuals in P at some point during each of the years 2013-2017, the proportion of individuals in A at some point during each of the years 2014-2015, and the proportion of individuals in H at some point during each of the years 2014-2016, resulting in a total of 10 data points.

[Include these next couple of descriptions in paper or only in dissertation?](#)

We simulate the proportions in each of these classes throughout the year in the following way. In order to count the proportion of individuals in P, A, or H at some point throughout a certain year, we need to count those who are in each of the classes AT ALL during the year, even if they leave or come back at some point. We note that we neglect higher order terms, i.e. people who go into the P class multiple times, since we can't keep track of individuals in model; the data we use is about total proportion of people who are in the class at some point throughout the year but they are kept track of and only counted once.

To get the output from the model of the proportion of non-addicted prescription opioid users in each year, we take the proportion of non-addicted prescription opioid users at the beginning of each of the years and add on the proportion of individuals that enter the P class at any point during the year. This latter part comes from integrating over those who enter the P class: $\int_0^t X'(t)dt = \int_0^t \alpha S(t)dt$ which is equal to $X(t) - X(0)$; here, we take $X(0) = 0$. For every year except 2013, a subtraction must be done since integrating gives the proportion from time 0 to time t , but we only want the proportion from time $t - 1$ to time t (i.e. the new cases in a given year). Thus, the calculation is as follows for each year, with the last equation in each line what was calculated in the code:

$$2013: P_0 + \int_0^1 \alpha S = y(1, 2) + y(2, 6) - y(1, 6)$$

$$2014: y(2, 2) + \int_1^2 \alpha S = y(2, 2) + \int_0^2 \alpha S - \int_0^1 \alpha S = y(2, 2) + y(3, 6) - y(2, 6)$$

$$2015: y(3, 2) + \int_2^3 \alpha S = y(3, 2) + \int_0^3 \alpha S - \int_0^2 \alpha S = y(3, 2) + y(4, 6) - y(3, 6)$$

$$2016: y(4, 2) + \int_3^4 \alpha S = y(4, 2) + \int_0^4 \alpha S - \int_0^3 \alpha S = y(4, 2) + y(5, 6) - y(4, 6)$$

$$2017: y(5, 2) + \int_4^5 \alpha S = y(5, 2) + \int_0^5 \alpha S - \int_0^4 \alpha S = y(5, 2) + y(6, 6) - y(5, 6)$$

To get the output from the model of the proportion of opioid addicts in 2014 and

2015, we take the initial proportion of opioid addicts in each of these years and add the proportion of individuals that enter the A class at any point during the year 2014 or 2015. This latter part comes from integrating over those who enter the A class: $\int_0^t L'(t)dt = \int_0^t (\gamma P + \sigma R \frac{A}{A+H+\omega} + \beta_A SA + \beta_P SP)dt$ which is equal to $L(t) - L(0)$; here, we take $L(0) = 0$. Similar to above with the subtraction of integrals to calculate for a specific year, we have:

$$2014: y(2, 3) + \int_1^2 (\gamma P + \sigma R \frac{A}{A+H+\omega} + \beta_A SA + \beta_P SP)dt = y(2, 3) + y(3, 7) - y(2, 7)$$

$$2015: y(3, 3) + \int_2^3 (\gamma P + \sigma R \frac{A}{A+H+\omega} + \beta_A SA + \beta_P SP)dt = y(3, 3) + y(4, 7) - y(3, 7)$$

Finally, to obtain the output from the model of the proportion of heroin/fentanyl addicts for each year 2014-2016, we take the initial proportion of heroin/fentanyl addicts in each of these years and add the proportion of individuals that enter the H class at any point during the year. This latter part comes from integrating over those who enter the H class: $\int_0^t M'(t)dt = \int_0^t (\theta_1 SH + \theta_2 PH + \theta_3 AH + \sigma R \frac{H}{A+H+\omega})dt$ which is equal to $M(t) - M(0)$; here, we take $M(0) = 0$. Similar to above with the subtraction of integrals to calculate for a specific year, we have:

$$2014: y(2, 4) + \int_1^2 (\theta_1 SH + \theta_2 PH + \theta_3 AH + \sigma R \frac{H}{A+H+\omega})dt = y(2, 4) + y(3, 8) - y(2, 8)$$

$$2015: y(3, 4) + \int_2^3 (\theta_1 SH + \theta_2 PH + \theta_3 AH + \sigma R \frac{H}{A+H+\omega})dt = y(3, 4) + y(4, 8) - y(3, 8)$$

$$2016: y(4, 4) + \int_3^4 (\theta_1 SH + \theta_2 PH + \theta_3 AH + \sigma R \frac{H}{A+H+\omega})dt = y(4, 4) + y(5, 8) - y(4, 8)$$

We use the Global Optimization Toolbox in MATLAB, specifically utilizing the MultiStart algorithm and fmincon local solver. Since we wish to know the global minimum, we use **fill in number** starting points as this is a local solver. We give lower bounds of 0.00001 for all of the parameters and an upper bound of 2 for all of the parameters except ϵ with an upper bound of 4 since this parameter can easily be greater than 1 (i.e. in a given year, a prescription user may end prescription use more than once). We run our model from 2013-2018, keeping a short time frame so that dynamics are relatively the same in the time period with the time step in years.

How many details to give for paper? Put in anything needed for paper from Mat-Lab_descriptions.pdf if applicable.

The following table displays the 7 parameters that were estimated with this process, along with their resulting estimated values:

Parameter	Estimated Value	Units
α		
θ_1		
ϵ		
γ		
σ		
ζ		
ν		

Sensitivity Analysis There are two ways in which we analyze how the uncertainty in our outputs can be apportioned to the uncertainty in each of the inputs of the model [30?].

[Sobol sensitivity analysis](#)

In order to explore the sensitivity of each of the population classes to the inputs (param-

eters and initial conditions), we implemented Sobol sensitivity analysis. First, the Saltelli sampler is utilized to generate $N(2D+2)$ samples where N =fill in is the number of sample points and $D=20$, the number of inputs in the model for a total of fill in samples [30]. Saltelli are the statistically good points chosen in the parameter space to test and the outputs are the values of S, P, A, H and R at the final time (year 2018) for the different parameter and initial condition samples. Cannot figure out why $N(2D+2)$ when including second order interactions. One method as described in [47] creates an $N \times 2k$ matrix of random numbers, and then splits this into two $N \times k$ matrices containing half of the samples each (call them J and L). From there, a third matrix is created in (M_i) which the i^{th} column is substituted by the i^{th} column of matrix L. The model output is then computed for all parameter samples (all of the rows of J, L, and M_i), which gives a total of $N+N+D \times N$ evaluations, or $N(D+2)$. From here, the first-order sensitivity index is measured as $S_i = \frac{V[E(Y|X_i)]}{V(Y)}$

should these have subscripts for V (subscript X_i) and E (subscript X_i)?

If so, does the mean come from only functions evaluated from M_i matrix? But that wouldn't match what's in [46] because equivalent of M_i there is $B_A^{(i)}$ which is not what they take the mean/variance over, it's X_i . So the first order index utilizes output from matrices A and B that are completely random as well and not focused on only varying that one parameter X_i ? This measures how much the output variance could be reduced on average if the i^{th} input were fixed and the sum of the S_i 's is equal to 1 if the model is additive(? a stats thing, so ours is not?) and less than 1 otherwise; the sum of the S_{T_i} values is only equal to 1 for additive models and otherwise greater than 1.

In [46], instead, the matrix X_i represents the $N \times (k-1)$ matrix of all inputs except input i . While fixing input i to some value, each row of X_i is used to evaluate the model; this is done for each generated value of the i^{th} input and then the mean of all of the outputs is taken. The variance of this mean is taken for this fixed input i , and then normalized over the variance of all of the outputs resulting in the first-order sensitivity index $S_i = \frac{V_{X_i}[E_{X_i}(Y|X_i)]}{V(Y)}$ (this is normalized since the numerator can take values all the way from 0 to $V(Y)$ based on the identity $V_{X_i}[E_{X_i}(Y|X_i)] + E_{X_i}[V_{X_i}(Y|X_i)] = V(Y)$). The total effect index is given by $S_{T_i} = 1 - \frac{V_{X_i}[E_{X_i}(Y|X_i)]}{V(Y)}$. Understand why.

How know which one the sobol analysis is using in SALib package, can't find anywhere how many evaluations it does; does it matter?

Read Saltelli paper and other reference total indices equation 44: term+complement+interaction terms

For first-order indices, only one parameter is varied and the rest are held constant; for second-order indices, two parameters are varied and the rest are held constant; and for total-order indices, every combination of parameters is varied, from a single parameter varying alone to all higher-order interactions between parameters. The length of the colored bars, with each color representing a specific class, measures the contribution of a certain parameter to the variance of each of the classes in the model. The longer the colored bar, the higher the effect the parameter has on that class of individuals; the sum of each color adds to 1, so that for each parameter, the relative contribution to each of the classes is represented. We are able to retrieve the confidence intervals on the sensitivity of each of the populations to a respective change in the parameter(s) for first order changes, second order changes and total order changes.

Below are the ranges we chose; most are $\pm 50\%$ of the baseline values found from parameter estimation or calculations of the parameters except for...FILL IN. We note that $\alpha = m * t + b$.

Parameter	Range of values
m	
β_A	
β_P	
θ_1	
ϵ	
γ	
σ	
μ	
μ_A	
μ_H	
θ_2	
ζ	
θ_3	
ν	
ω	
b	
P_0	
A_0	
H_0	
R_0	

Latin Hypercube Sampling/PRCC

Conclusions

What did we show that we didn't know before?

Think of questions that we can confront the model with regarding heroin and fentanyl deaths and connect to data on deaths; give results that give the reader an answer to these questions; think about analysis can do on model.

Extensions

Need to edit

In our model we assume homogeneous mixing of the individuals in each of the compartments and therefore, each individual has the same probability of transitioning from one stage to another, interacting with individuals from other stages, or leaving the system via death. Although done for the purpose of simplification, this is not an accurate representation of reality since there are factors that affect these probabilities, such as race, gender, geographical location and age. For example, there has been a shift in heroin use from urban areas to some suburban and rural areas and there is an increasing number of individuals ages 18-25 using the drug [40]. West Virginia, New Hampshire and Ohio had the highest rates of opioid-related overdose deaths per 100,000 people in 2016 and

Tennessee and Arkansas were among the states with the highest prescription rates per 100 people the same year [42]. Men have a higher likelihood of overdosing on prescription pain relievers compared to women, but since 1999, there has been a steeper increase in the number of women overdoses than men. In addition, women may form a quicker dependence on opioids compared to men, and have a higher probability of being prescribed higher doses and for longer periods of time [13]. Specific to heroin, using data from the 2008-2010 NSDUH studies, past year heroin use was twice as likely for men compared to women, and highest in the 18-25 year old age group [34]. Incorporating these ideas into the model and parameter information would be possible future extensions. Although there is data on the misuse of opioids, we decided to focus specifically on addiction rather than misuse, due to the harmful consequences this behavior can have for an individual. However, the behavior and actions of individuals are not always very clearly defined, and there may be a benefit to expanding the spectrum of classes that individuals can fall into.

Appendix A

Calculations of parameters from literature

The **age-adjusted death rate** for Tennessee in 2016 was calculated to be 886.3 out of 100,000 individuals, or approximately 50,094 people out of a total population 12 and older of 5,651,993 [29]. Subtracting off the number of people who died from a prescription opioid or heroin/fentanyl overdose in 2016 results in 48,825 people who died that year. This implies that $(5,651,993 - 48,825) / 5,651,993 \approx 0.991$ is the proportion of the population that remains by the beginning of 2017. If we consider T_0 to be the total population in 2016, we can find the continuous-time rate at which individuals die naturally from the equation $0.991T_0 = T_0 e^{-\mu t}$, which results in the natural death rate $\mu \approx 0.00868$.

We calculate the overdose death rate in the year 2015, since we know the number of prescription opioid addicted individuals in that year. To find the continuous-time rate at which individuals are dying from the addicted class, we consider the equation $kA_0 = A_0 e^{-\mu_A t}$, where A_0 is the number of individuals addicted to prescription opioids in 2015 and k is the proportion of these individuals in the addicted class at the start of 2016 (when $t = 1$). In 2015, there were 679 individuals out of the entire Tennessee population 12 and older that overdosed on prescription opioids [45]. However, it is estimated that only 54.6% of these individuals were actually at an increased risk for an opioid-related overdose death; we assume that if an individual met the criteria for at least one high-risk factor, that they were considered addicted to prescription opioids [7]. Therefore, we assume 54.6% of the 679 individuals that overdosed were addicted. With a total of 42,816 prescription opioid addicts (non-heroin addicted) in 2015, this means that $(42,816 - 0.546 \cdot 679) / 42,816 \approx 0.992$ is the proportion of addicted individuals that remain by the beginning of the next year. This implies $0.991A_0 = A_0 e^{-\mu_A(1)}$, and solving results in **$\mu_A \approx 0.00870$** . We note here that since we want a rate of overdose deaths out of the addicted class, we use the number of non-heroin addicted prescription opioid addicts, although the overdose deaths are out of all prescription opioids addicts, whether they are addicted to heroin or not. However, since we do not know the number of overdoses specifically for non-heroin addicted prescription opioid addicts, we use this simplification that all of them are for this group.

Similarly, we may calculate μ_H . There were 7,560 heroin/fentanyl addicts in 2015 and 374 heroin-related overdoses. We make the simplifying assumption that if an individual died of a heroin overdose they were addicted to heroin. This means that $(7,560 - 374) / 7,560 \approx$

0.951 is the proportion of heroin users that remain at $t = 1$, which implies $0.951H_0 = H_0e^{-\mu_H(1)}$, which results in $\mu_H \approx \mathbf{0.0507}$, the continuous-time rate at which individuals are dying from the heroin class.

We make a note that individuals that do not have an opioid use disorder and die because of an opioid overdose are counted in the “natural mortality rate.”

Estimates of parameters from literature

Since there is no information for values of θ_1 , θ_2 , or θ_3 for Tennessee in the literature, we consider a national study of individuals 12 and older to establish a relationship among these three rates. For a national study consisting of 609,000 participants, “the recent heroin incidence rate was 19 times higher among those who reported prior non-medical pain reliever (NMPR) use (0.39%) than among those who did not report NMPR use (0.02%) [38]. NMPR use can occur within the prescription class (i.e. from misuse that’s not considered addiction), or in the addiction class. Thus, we will extrapolate this information to say that the rate that prescription opioid users and opioid addicts move to heroin use is 19 times greater than the rate at which susceptibles move to heroin use (i.e. $\theta_2 + \theta_3 > 19\theta_1$), where both θ_2 and θ_3 are greater than θ_1 . We will make an assumption here that $\theta_2 + \theta_3$ is exactly $19\theta_1$, and that there is somewhat of an elevated risk of moving to heroin use from the prescription class versus the susceptible class, taking $\theta_2 = 3\theta_1$. This leaves $\theta_3 = 16\theta_1$.

Okay to use even though opioid model is not a sub-model anymore (since we’ve changed the recovery class and terms? We argue that β_A is a reasonable parameter to estimate because the opioid-only model was insensitive to this parameter and although less so, β_P was relatively insignificant compared to other parameters explored [6]. We were not able to find values in the literature specific to Tennessee regarding an illicit-induced addiction rate; however, we use information from the national level, as we do not have any evidence that this national average rate would be significantly different in Tennessee. Have this work checked/all units okay? In 2015, there were an estimated 2.1 million individuals 12 and older that initiated pain reliever misuse recently. Using this source, we take pain relievers to be equivalent to prescription opioids [31]. To calculate approximately how many of these would end up having a use disorder in 2015, this same study states that 12.5 million people misused prescription opioids in the past year total, and that there were 2.0 million with a prescription opioid disorder; using these number, we make an estimate that

$$\frac{2 \text{ million people with pain reliever use disorder in 2015}}{12.5 \text{ million people who misused pain relievers in 2015}} = 0.16.$$

We extrapolate this to estimate that 16% of the 2.1 million recent initiates are those who should be counted in the use disorder category (although we do not know the timeline of developing the use disorder and it may not happen within the year, it gives a rough idea). This results in an estimate of 336,000 individuals that were recent initiates of pain reliever misuse that we will consider having a use disorder within the year 2015. In the same year, a study was done regarding the source of the most recent episode of misuse of prescription opioids among adults reporting an prescription opioid use disorder in the past year [27]. (Note: this study was for individuals age 18 and older whereas the study done in [31] was for those 12 and older; we will assume for simplicity that the information for those 18 and older will be similar to those between 12 and 18, as well.) Here, 52.8% of the prescription opioids were from a non-physician source, which included being given

to/bought from a friend/relative, or bought from a drug dealer/stranger. For simplicity, let IPUD indicate individuals with a prescription use disorder. Thus,

$$\begin{aligned} & \frac{336,000 \text{ IPUD}}{320 \text{ million people}} \cdot \frac{0.528 \text{ opioid source for IPUD is illicit}}{\text{opioid sources} \cdot \text{year}} \\ &= 0.00105 \frac{\text{IPUD obtain opioids illicitly}}{\text{opioid sources} \cdot \text{year}}. \end{aligned}$$

From [27], we see that

$$\frac{.138 \text{ opioids obtained from drug dealers/strangers by IPUD}}{.528 \text{ opioids obtained illicitly by IPUD}} = .26$$

relates to individuals obtaining opioids by the black market or interaction with other addicts and

$$\frac{.218 + .141 + .031 \text{ opioids given/bought from a friend/relative or other for IPUD}}{.528 \text{ opioids obtained illicitly by IPUD}} = .74$$

relates to individuals obtaining opioids from extra prescriptions that are available. This means that 74% of illicit opioids come from a friend/relative/other source, whereas 26% of illicit opioids are from the black market for opioid addicts. In our model, we consider two cases in which individuals become addicted (friend/relative versus black market) and we make the assumption that these percentages also apply to the two routes that one would become an addict in the first place via illicit opioids.

Thus,

$$\begin{aligned} & \frac{0.26 \text{ opioids from drug dealers/strangers}}{\text{opioids obtained illicitly}} \cdot \frac{0.00105 \text{ opioids obtained illicitly}}{\text{opioid sources} \cdot \text{year}} \\ &= \frac{0.000273 \text{ opioids from drug dealers/strangers}}{\text{opioid sources} \cdot \text{year}} \end{aligned}$$

is the rate that individuals illicitly obtain their prescription opioids from the black market/other addict interaction and become addicted

$$\implies \beta_A = \frac{0.000273}{\text{year}}.$$

Similarly,

$$\begin{aligned} & \frac{0.74 \text{ opioids from friend/relative or other}}{\text{opioids obtained illicitly}} \cdot \frac{0.00105 \text{ opioids obtained illicitly}}{\text{opioid sources} \cdot \text{year}} \\ &= \frac{0.000777 \text{ opioids from friend/relative or other}}{\text{opioid sources} \cdot \text{year}} \end{aligned}$$

is the rate that individuals illicitly obtain their prescription opioids from extra pills that are available and become addicted

$$\implies \beta_P = \frac{0.000777}{\text{year}}.$$

As mentioned before, 71% of heroin users in 4 weeks [49]. Although these are on a national

level, we will assume the rates do not differ significantly for Tennessee, as we were not able to find any relapse statistics specifically for the state. It is estimated that for every 10 people with substance use disorder, only 1 receives treatment for it [61]. We assume this is an approximation for specifically opioid and heroin/fentanyl addicts, as well. In 2015, there were 13,110 discharges total in the state of Tennessee for substance abuse programs in facilities that receive state/public funding (generally referring to funding by the state substance abuse agency); out of those, 7,003 were due to completion of treatment, which is 53.4% of all discharges [21]. Realistically, an individual cannot go to a stably recovered class without going to treatment, completing treatment and remaining clean for 4 weeks post-treatment, since addiction is a disease. Using HA to represent heroin addict, we calculate

$$\begin{aligned}
& \frac{1 \text{ HA in treatment}}{10 \text{ HA} \cdot \text{year}} \cdot \frac{53.4 \text{ HA complete treatment}}{100 \text{ HA in treatment}} \cdot \frac{29 \text{ HA do not relapse within 4 weeks post-treatment}}{100 \text{ HA completed treatment}} \\
&= \frac{0.015486 \text{ HA do not relapse within 4 weeks post-treatment}}{\text{HA} \cdot \text{year}} \\
&\implies \nu \approx \frac{0.0155}{\text{year}}.
\end{aligned}$$

Studies done among opiate addicts (not just heroin) in recent years ([5, 9]) suggest lower relapse rates at 4 weeks compared to the study consisting mostly of heroin users in [49] in which 71% of the addicts relapsed within 4 weeks. Thus, we assume the relapse rate for prescription opioids will be slightly lower than that for heroin. We know from a study done in [69] that at the 8 week mark, 91% of prescription opioids users had relapsed, so this is an upper limit, and we know from studies that higher rates of relapse occur in the first few weeks following treatment, as discussed before. Therefore, we make an estimate that 60% of prescription opioids users relapse within 4 weeks after treatment. Letting OA represent opioid addicts, we calculate

$$\begin{aligned}
& \frac{1 \text{ OA in treatment}}{10 \text{ OA} \cdot \text{year}} \cdot \frac{53.4 \text{ OA complete treatment}}{100 \text{ OA in treatment}} \cdot \frac{40 \text{ OA do not relapse within 4 weeks post-treatment}}{100 \text{ OA completed treatment}} \\
&= \frac{0.02136 \text{ OA do not relapse within 4 weeks post-treatment}}{\text{OA} \cdot \text{year}} \\
&\implies \zeta \approx \frac{0.0214}{\text{year}}.
\end{aligned}$$

Since the H class is homogeneously mixed with all addicts, whether they have been in treatment or not, the calculation for ν takes that into consideration to produce an average rate among all addicts within the class and therefore, gives a rate at which any individual from that class would transition to the recovered class; similarly for the A class.

Estimates of initial conditions from literature

In the first quarter of 2013, there were 856,000 individuals prescribed opioids in Tennessee, and a total of 2,000,000 prescriptions in the same quarter; this is approximately 2.34 prescriptions per prescribed user in this quarter [45]. In addition, the average number

of days that prescribed users in Tennessee actively took prescription opioids in 2013 was 66.9 days; extrapolating this to the first quarter gives

$$\frac{x \text{ average prescription days}}{90 \text{ days}} = \frac{66.9 \text{ average prescription days}}{365 \text{ days}}$$

$$\implies x \approx 16.5$$

as the average number of prescription days in the first quarter of 2013 for a prescribed user. Since we do not know the number of individuals using prescription opioids at the beginning of the year, we use

$$\begin{aligned} & \# \text{ of prescribed users at beginning of 2013} \\ & + \frac{2,000,000 \text{ prescriptions}}{90 \text{ prescription days}} \cdot \frac{16.5 \text{ average prescription days}}{2.34 \text{ average number of prescriptions}} \\ & = 856,000 \text{ prescription users in entire first quarter of 2013} \end{aligned}$$

gives an estimate that

$$\# \text{ of prescribed users at beginning of 2013} = 699,304.$$

However, as before, approximately 44.3% of these prescription opioid users are addicted to their prescriptions, which means

$$\# \text{ of non-addicted prescribed users at beginning of 2013} = (1 - .443) * 699,304 = 389,512.$$

Although we do not know the total population at the beginning of 2013, we will take it to be approximately the total population from 2012 before any births or deaths occur, which was 6,450,632. Adjusting this to the population 12 and older as done in Table 1, we take 15% off of the total population to arrive at 5,483,037 [59].

Thus, we estimate

$$P_0 = \frac{389,512}{5,483,037} = 0.0710$$

of the population was using opioid prescriptions at the beginning of 2013. In 2013, we observe a ratio of approximately 1 non-heroin addicted prescription opioid addict to 42.1 non-addicted prescription opioid user, so we assume this is the ratio at the beginning of the year, as well. This gives

$$A_0 = \frac{P_0}{42.1} = 0.00169.$$

WILL MOST LIKELY DELETE BECAUSE NOW ESTIMATING H₀ and R₀ WITH PARAMETER ESTIMATION/MULTISTART.

We do not have information on the number of heroin users in 2013, so we extrapolate from information in 2015. We note first that although opioid overdose deaths increase from 2013 to 2015, that does not necessarily equate to more heroin users in 2015, due to the rise in fentanyl contamination during this time period...HOWEVER, IT SEEMS THAT if heroin were laced with heroin, it would be counted as fentanyl overdose, so the fact that heroin overdoses are also still increasing through 2016 suggests more heroin addicts

overall [11]. In order to keep a proportion approximately equivalent to the number of opioid addicts to heroin users in 2015, we assume the same ratio of approximately 1:3.43 heroin users to opioid addicts in 2013, as well. In 2013, addicts make up 0.00148 of the population at the start of 2013, so we assume $H_0 \approx 0.000431$. (Note: the proportion of prescription opioid users to heroin users is also similar, particularly regarding the order of magnitude expected.)

Before, we estimated 48,674 opioid addicts in 2013; this higher number is supported by an overall higher prescribing rate in 2013 compared to the next few years [14]. We will assume the number of heroin users is the same in 2013 as in 2014 and 2015. There were 4,485 prescription opioid treatment admissions and 555 heroin treatment admissions in 2013, which is 0.057 of the opioid addict and heroin user population combined that are in recovery. Only 35% of those in treatment do not relapse within 4 weeks after treatment, and thus have the ability to be in R, which is approximately .0804 of the entire A+H class [5]. Since 90% of individuals relapse within one year post-treatment, that means that an additional 55% relapse after the 4 week mark and move out of R; therefore, we make a simplifying assumption that 10% of those that can be in R actually are at a specific point in time, resulting in a proportion of .00804 of addicted and heroin users total [5]. This equates to approximately 504 individuals, which out of a total population 12 years and older in 2013 of 5,517,716, we conclude $R_0 = 0.000091$.

CS-unit conversions from data to parameters is what I've done above already right?

Table of parameter values and units of parameters

Fill in

Parameter	Description	Units
μ	natural mortality rate	$\frac{1}{\text{year}}$
μ_A	opioid addict overdose death rate	$\frac{1}{\text{year}}$
μ_H	heroin addict overdose death rate	$\frac{1}{\text{year}}$
α	prescription rate	$\frac{1}{\text{year}}$
β_A	illicit addiction rate from the black market	$\frac{1}{\text{year}}$
β_P	illicit addiction rate from availability of excess pills	$\frac{1}{\text{year}}$
θ_1	heroin addiction rate for susceptible individuals	$\frac{1}{\text{year}}$
ϵ	rate of finishing prescription addiction-free	$\frac{1}{\text{year}}$
γ	opioid addiction rate from prescription	$\frac{1}{\text{year}}$
θ_2	heroin addiction rate for prescription opioids users	$\frac{1}{\text{year}}$
σ	relapse to addiction	$\frac{1}{\text{year}}$
ζ	rate of stable recovery for opioid addict	$\frac{1}{\text{year}}$
θ_3	heroin addiction rate for opioid addicts	$\frac{1}{\text{year}}$
ν	rate of stable recovery for heroin addict	$\frac{1}{\text{year}}$
ω	perturbation term	dimensionless
S	proportion of susceptible individuals	dimensionless
P	proportion of susceptible individuals	dimensionless
A	proportion of susceptible individuals	dimensionless
H	proportion of susceptible individuals	dimensionless
R	proportion of susceptible individuals	dimensionless

References

- [1] (2017). *Modeling approaches, challenges, and preliminary results for the opioid and heroin co-epidemic crisis*.
- [2] Abdurahman, X., Shang, L., and Teng, Z. (2014). Global dynamics of a discretized heroin epidemic model with time delay. *Abstract and Applied Analysis*, 2014.
- [3] Agar, M. H. (2001). Another complex step: A model of heroin experimentation. *Field Methods*, 13(4):353–369.
- [4] American Psychiatric Association (2000). American psychiatric association: *Diagnostic and Statistical Manual of Mental Disorders Fourth Edition*. Available at <https://dsm.psychiatryonline.org/doi/pdf/10.1176/appi.books.9780890420249.dsm-iv-tr> (accessed 10/10/2018).
- [5] Bailey, G. L., Herman, D. S., and Stein, M. D. (2013). Perceived relapse risk and desire for medication assisted treatment among persons seeking inpatient opiate detoxification. *Journal of Substance Abuse Treatment*, 45:302–305.
- [6] Battista, N., Percy, L., and Strickland, W. C. (2018). Modeling the prescription opioid epidemic. *NEED TO UPDATE*.
- [7] Baumblatt, J. A. G., Wiedeman, C., Dunn, J. R., Schaffner, W., Paulozzi, L., and Jones, T. F. (2014). High-risk use by patients prescribed opioids for pain and its role in overdose deaths. *JAMA Internal Medicine*, 175(5):796–801.
- [8] Bradley, B., Phillips, G., Green, L., and Gossop, M. (1989). Circumstances surrounding the initial lapse to opiate use following detoxification. *British Journal of Psychiatry*, 154:354–359.
- [9] Broers, B., Giner, F., Dumont, P., and Mino, A. (2000). Inpatient opiate detoxification in geneva: follow-up at 1 and 6 months. *Drug and Alcohol Dependence*, 58:85–92.
- [10] Centers for Disease Control and Prevention (2016). Synthetic opioid overdose data. Available at <https://www.cdc.gov/drugoverdose/data/fentanyl1.html> (accessed 05/01/2018).
- [11] Centers for Disease Control and Prevention (2017a). Opioid data analysis. Available at <https://www.cdc.gov/drugoverdose/data/analysis.html> (accessed 05/01/2018).
- [12] Centers for Disease Control and Prevention (2017b). Opioid overdose. Available at www.cdc.gov/drugoverdose/opioids/index.html (accessed 07/16/2018).
- [13] Centers for Disease Control and Prevention (2017c). Prescription painkiller overdoses. Available at <https://www.cdc.gov/vitalsigns/PrescriptionPainkillerOverdoses/index.html> (accessed 7/19/2018).
- [14] Centers for Disease Control and Prevention (2017d). Vital signs: Changes in opioid prescribing in the united states 2006-2015. Available at <https://www.cdc.gov/mmwr/volumes/66/wr/mm6626a4.htm> (accessed 3/5/19).

- [15] Cicero, T. J., , Ellis, M. S., and Surratt, H. L. (2012). Effect of abuse-deterrent formulation of oxycontin. *New England Journal of Medicine*, 367(2):187–189. PMID: 22784140.
- [16] Cicero, T. J. and Ellis, M. S. (2015). Abuse-deterrent formulations and the prescription opioid abuse epidemic in the United States: Lessons learned from oxycontin. *JAMA Psychiatry*, 72(5):424–430.
- [17] Cicero, T. J., Ellis, M. S., Surratt, H. L., and Kurtz, S. P. (2014). The changing face of heroin use in the United States: A retrospective analysis of the past 50 years. *JAMA Psychiatry*, 71(7):821–826.
- [18] Compton, W. M. (2017). Research on the use and misuse of fentanyl and other synthetic opioids. Available at <https://www.drugabuse.gov/about-nida/legislative-activities/testimony-to-congress/2017/research-use-misuse-fentanyl-other-synthetic-opioids> (accessed 07/18/2018).
- [19] Department of Health and Human Services: Administration for Children and Families, Office of Child Care (2018). State/territory profile-tennessee. Available at: <https://childcareta.acf.hhs.gov/state-profiles/profiles/TN/pdf> (accessed 11/5/18).
- [20] Department of Health and Human Services: Substance Abuse and Mental Health Services Administration (2017). Treatment episode data set (teds) 2005-2015 state admissions to substance abuse treatment services. Available at: https://www.dasis.samhsa.gov/dasis2/teds_pubs/2015_teds_rpt_st.pdf (accessed 9/24/18).
- [21] Department of Health and Human Services: Substance Abuse and Mental Health Services Administration (2018). Treatment episode data set (teds) 2015: Discharges from substance use treatment services: Tables. Available at https://www.dasis.samhsa.gov/dasis2/teds_pubs/2015_teds_rpt_d.pdf.
- [22] Fang, B., Li, X.-Z., Martcheva, M., and Cai, L.-M. (2015a). Global asymptotic properties of a heroin epidemic model with treat-age. *Applied Mathematics and Computation*, 263:315–331.
- [23] Fang, B., Li, X.-Z., Martcheva, M., and Cai, L.-M. (2015b). Global stability for a heroin model with age-dependent susceptibility. *Journal of Systems Science and Complexity*, 28(6):1243–1257.
- [24] Florence, C. S., Zhou, C., Luo, F., and Xu, L. (2016). The economic burden of prescription opioid overdose, abuse and dependence in the United States, 2013. *Medical Care*, 54(10):901–906.
- [25] Gossop, M., Green, L., Phillips, G., and Bradley, B. (1989). Lapse, relapse and survival among opiate addicts after treatment: A prospective follow-up study. *British Journal of Psychiatry*, 154:348–353.
- [26] Gossop, M., Stewart, D., Browne, N., and Marsden, J. (2002). Factors associated with abstinence, lapse or relapse to heroin use after residential treatment: protective effect of coping responses. *Addiction*, 97:1259–1267.

- [27] Han, B., Compton, W. M., Blanco, C., Crane, E., Lee, J., and Jones, C. M. (2017). Prescription opioid use, misuse, and use disorders in u.s. adults: 2015 national survey on drug use and health. *Annals of Internal Medicine*, 167(5):293–302.
- [28] Heather, N., Stallard, A., and Tebbutt, J. (1991). Importance of substance cues in relapse among heroin users: comparison of two methods of investigation. *Addictive Behaviors*, 16:41–49.
- [29] Henry J Kaiser Family Foundation (2018). Number of deaths per 100,000 population. Available at [https://www.kff.org/other/state-indicator/death-rate-per-100000/?currentTimeframe=0&sortModel=\(accessed 11/5/18\)](https://www.kff.org/other/state-indicator/death-rate-per-100000/?currentTimeframe=0&sortModel=(accessed 11/5/18)).
- [30] Herman, J. and Usher, W. (2017). Salib-sensitivity analysis library in python. Available at <https://salib.readthedocs.io/en/latest/> (accessed 05/01/2018).
- [31] Hughes, A., Williams, M. R., Lipari, R. N., Bose, J., Copello, E. A., and Kroutil, L. (2016). Prescription drug use and misuse in the united states: Results from the 2015 national survey on drug use and health. Available at <https://www.samhsa.gov/data/sites/default/files/NSDUH-FFR2-2015/NSDUH-FFR2-2015.htm> (accessed 3/14/19).
- [32] Hunt, W. A., Barnett, L. W., and Branch, L. G. (1971). Relapse rates in addiction programs. *Journal of Clinical Psychology*, 27(4):455–456.
- [33] Ihongbe, T. O. and Masho, S. W. (2016). Prevalence, correlates and patterns of heroin use among young adults in the United States. *Addictive Behaviors*, 63:74–81.
- [34] Jones, C. M. (2013). Heroin use and heroin use risk behaviors among nonmedical users of prescription opioid pain relievers – united states, 2002–2004 and 2008–2010. *Drug and Alcohol Dependence*, 132(1):95 – 100.
- [35] Liu, J. and Zhang, T. (2011). Global behaviour of a heroin epidemic model with distributed delays. *Applied Mathematics Letters*, 24:1685–1692.
- [36] Ma, M., Liu, S., and Li, J. (2017). Bifurcation of a heroin model with nonlinear incidence rate. *Nonlinear Dynamics*, 88(1):555–565.
- [37] Mandell, B. (June 2016). The fifth vital sign: A complex story of politics and patient care. *Cleveland Clinic Journal of Medicine*, 83(6).
- [38] Muhuri, P. K., Gfroerer, J. C., and Davies, M. C. (2013). Associations of nonmedical pain reliever use and initiation of heroin use in the United States.
- [39] National Institute on Drug Abuse. Abuse of prescription pain medications risks heroin use. Available at https://www.drugabuse.gov/sites/default/files/infographic_presc_heroin.pdf (accessed 3/27/19).
- [40] National Institute on Drug Abuse (2018a). Heroin. Available at <https://d14rmgtrwzf5a.cloudfront.net/sites/default/files/828-heroin.pdf> (accessed 7/18/2018).

- [41] National Institute on Drug Abuse (2018b). Heroin. Available at <https://www.drugabuse.gov/publications/drugfacts/heroin#ref>(accessed 07/17/2018).
- [42] National Institute on Drug Abuse (2018c). Opioid summaries by state. Available at <https://www.drugabuse.gov/drugs-abuse/opioids/opioid-summaries-by-state> (accessed 7/18/2018).
- [43] National Institute on Drug Abuse (2018d). Prescription opioids and heroin. Available at <https://d14rmgtrwzf5a.cloudfront.net/sites/default/files/19774-prescription-opioids-and-heroin.pdf> (accessed 04/16/2018).
- [44] Office of Information and Analytics Tennessee Department of Health. Data dashboard. Available at: <https://www.tn.gov/health/health-program-areas/pdo/pdo/data-dashboard.html> (accessed 8/14/18).
- [45] Office of Information and Analytics: Tennessee Department of Health (2018). Prescription drug overdose program 2018 report. Available at https://www.tn.gov/content/dam/tn/health/documents/pdo/PDO_2018_Report_02.06.18.pdf (accessed).
- [46] Saltelli, A., Annoni, P., Azzini, I., Campolongo, F., Ratto, M., and Tarantola, S. (2010). Variance based sensitivity analysis of model output. design and estimator for the total sensitivity index. *Computer Physics Communications*, 181:259–270.
- [47] Saltelli, A., Ratto, M., Andres, T., Campolongo, F., Cariboni, J., Gatelli, D., Saisana, M., and Tarantola, S. (2008). *Global Sensitivity Analysis: The Primer*. Wiley.
- [48] Samanta, G. (2011). Dynamic behaviour for a nonautonomous heroin epidemic model with time delay. *Journal of Applied Mathematics and Computing*, 35(1-2):161–178.
- [49] Smyth, B., Barry, J., and E. Keenan, K. D. (2010). Laspse and relapse following inpatient treatment of opiate dependence. *Irish Medical Journal*, 103(6):176–179.
- [50] Substance Abuse and Mental Health Services Administration. Table 95 selected drug use, past year alcohol use disorder, and past year mental health measures in tennessee, by age group: Estimated numbers (in thousands), annual averages based on 2014-2015 nsduhs. Available at <https://www.samhsa.gov/data/report/2014-2015-nsduh-state-specific-tables> (accessed 2/7/19).
- [51] Substance Abuse and Mental Health Services Administration. Table 95 selected drug use, perceptions of great risk, past year substance use disorder and treatment, and past year mental health measures in tennessee, by age group: Estimated numbers (in thousands), annual averages based on 2015-2016 nsduhs. Available at <https://www.samhsa.gov/data/report/2015-2016-nsduh-state-specific-tables> (accessed 11/27/17).
- [52] Substance Abuse and Mental Health Services Administration. Table 95 selected drug use, perceptions of great risk, past year substance use disorder and treatment, and past year mental health measures in tennessee, by age group: Estimated numbers (in thousands), annual averages based on 2016-2017 nsduhs. Available at

- <https://www.samhsa.gov/data/report/2016-2017-nsduh-state-specific-tables> (accessed 2/7/19).
- [53] Substance Abuse and Mental Health Services Administration (2015a). Medication and counseling treatment. Available at <https://www.samhsa.gov/medication-assisted-treatment/treatment> (accessed 07/16/2018).
 - [54] Substance Abuse and Mental Health Services Administration (2015b). Substance use disorders. Available at <https://www.samhsa.gov/disorders/substance-use> (accessed 07/16/2018).
 - [55] Substance Abuse and Mental Health Services Administration (2017a). 2016 NSDUH report: America's behavioral health changes and challenges. Available at https://www.samhsa.gov/sites/default/files/topics/data_outcomes_quality/nsduh-ppt-09-2017.pdf (accessed 06/26/2018).
 - [56] Substance Abuse and Mental Health Services Administration (2017b). Key substance use and mental health indicators in the United States: Results from the 2016 national survey on drug use and health. Available at <https://www.samhsa.gov/data/sites/default/files/NSDUH-FFR1-2016/NSDUH-FFR1-2016.pdf> (accessed 06/14/2018).
 - [57] Substance Abuse and Mental Health Services Administration and Center for Behavioral Health Statistics and Quality (2015). 2016 national survey on drug use and health (NSDUH): Final approved CAI specifications for programming (english version). Available at <https://www.samhsa.gov/data/sites/default/files/NSDUHmrbCAIquex2016v2.pdf> (accessed 06/14/2018).
 - [58] Tennessee Department of Mental Health and Substance Abuse Services (2017). Trends in drug overdose deaths involving opioids: Tennessee and the united states 2014 and 2015. Available at https://www.tn.gov/content/dam/tn/mentalhealth/documents/Drug_Overdose_Deaths_CDC_2.14.2017 (accessed 9/25/18).
 - [59] United States Census Bureau (2018). National, state, and puerto rico commonwealth totals datasets: Population, population change, and estimated components of population change: April 1, 2010 to july 1, 2017. Available at: <https://www.census.gov/data/datasets/2017/demo/popest/state-total.html#partextimage500989927> (accessed 9/6/18).
 - [60] United States Drug Enforcement Administration. Drug scheduling. Available at <https://www.dea.gov/druginfo/ds.shtml> (accessed 07/17/2018).
 - [61] U.S. Department of Health and Human Services (2016). Facing addiction in america: The surgeon general's report on alcohol, drugs, and health. Available at <https://addiction.surgeongeneral.gov/key-findings/early-intervention> (accessed 3/24/19).

- [62] U.S. Department of Health and Human Services (2017). HHS acting secretary declares public health emergency to address national opioid crisis. Available at <https://www.hhs.gov/about/news/2017/10/26/hhs-acting-secretary-declares-public-health-emergency-address-national-opioid-crisis.html> (accessed 04/16/2018).
- [63] U.S. Food and Drug Administration (2018). Timeline of selected FDA activities and significant events addressing opioid misuse and abuse. Available at <https://www.fda.gov/downloads/Drugs/DrugSafety/InformationbyDrugClass/UCM566985.pdf> (accessed 06/13/2018).
- [64] van den Driessche, P. and Watmough, J. (2008). *Mathematical Epidemiology*, chapter 6: Further Notes on the Basic Reproduction Number, pages 159–178. Springer, Berlin, Heidelberg.
- [65] Volkow, N. D. (2016). What science tells us about opioid abuse and addiction. Available at <https://www.drugabuse.gov/about-nida/legislative-activities/testimony-to-congress/2018/what-science-tells-us-about-opioid-abuse-addiction> (accessed 07/17/2018).
- [66] Vowles, K. E., McEntee, M. L., Julnesa, P. S., Frohe, T., Neyb, J. P., and van der Goes, D. N. (2015). Rates of opioid misuse, abuse, and addiction in chronic pain: a systematic review and data synthesis. *PAIN*, 156(4):569–576.
- [67] Wang, X., Yang, J., and Li, X. (2011). Dynamics of a heroin epidemic model with very population. *Applied Mathematics*, 2(6):732–738.
- [68] Wangari, I. M. and Stone, L. (2017). Analysis of a heroin epidemic model with saturated treatment function. *Journal of Applied Mathematics*, 2017.
- [69] Weiss, R. D. and Rao, V. (2017). The prescription opioid addiction treatment study: What have we learned. *Drug and Alcohol Dependence*, 173:48–54.
- [70] White, E. and Comiskey, C. (2007). Heroin epidemics, treatment and ODE modelling. *Mathematical Biosciences*, 208:312–324.