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Overcoming the “Valley of Death” in Medications Development for Alcohol Use Disorder

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Abstract

Background: As the development of novel pharmacotherapies for alcohol use disorder (AUD) has been slow, the discovery and testing of more efficacious pharmacotherapies for AUD represents a high priority research area. In fact, the transition from pre-clinical to clinical testing of novel compounds has been termed the “valley of death” in medications development. One key obstacle consists of the lack of an articulated set of goals for each stage of medications development. Specifically, the knowledge outputs required to make the transition from safety testing, to early efficacy detection, to confirming clinical efficacy remain unclear, and this is despite a great deal of interest and substantial financial investment in developing novel therapeutics for AUD.

Method: This qualitative review manuscript seeks to draw parallels and lessons from the well-established stage model for behavioral therapies research with alcohol and other substance use disorders, and to apply these insights to AUD pharmacotherapy development.

Results: We argue that human laboratory models and/or pilot randomized controlled trials should serve as intermediaries in the transition from pre-clinical studies to large, and costly, randomized controlled efficacy trials. The relative strengths and weaknesses of pilot clinical trials versus human laboratory studies for bridging the “valley of death” are discussed and explored via a Monte Carlo data simulation study.

Conclusions: Multiple permutations of suitable research designs informed by the behavioral therapies development model are discussed with the overall goal of promoting consilience and maximizing efficiency across all phases of clinical testing of novel AUD pharmacotherapies.

Keywords

medications development; pharmacotherapy; behavioral therapies research; alcohol use disorder; treatment

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Introduction

Pharmacotherapies for alcohol use disorder (AUD) are used less often than behavioral therapies (Fuller and Hiller-Sturmhofel, 1999). The limited use of pharmacotherapy for AUD is due, in part, to a very limited number of approved options. Currently, the only pharmacotherapies approved by the Food and Drug Administration (FDA) for the treatment of AUD are disulfiram (Antabuse®), acamprosate, oral naltrexone, and Vivitrol (i.e., an injectable extended-release formulation of naltrexone). Given the dearth of available pharmacotherapies, as well as their limited-to-moderate efficacy (Rosner et al., 2010, Jonas et al., 2014, Blodgett et al., 2014, Donoghue et al., 2015), medications development to treat AUD remains a top research priority (Litten et al., 2012a, Litten et al., 2016). In fact, current foci in medications development emphasize the identification of new molecular targets and the development of novel compounds for these targets (Litten et al., 2012a, Ray et al., 2014). This is consistent with the phenotypic complexity of AUD, which calls for multiple molecular targets that can effectively address multiple pathways of risk (Litten et al., 2015). Despite rigorous efforts in medications development for AUD over the past two decades, it is apparent that there are numerous obstacles to an efficient process of screening and developing novel compounds.

One primary obstacle is the transition from pre-clinical to clinical testing, which has been termed the “valley of death” in medications development (Litten et al., 2012). Specifically, the process of shepherding novel compounds from promising preclinical results in animal models through Investigational New Drug (IND) application requirements and human testing has been historically fraught; several promising compounds never advanced from preclinical to human subjects testing due to a variety of issues. A number of structural and regulatory issues are at play and NIH-level initiatives have recently been put in place to facilitate this transition, including incentives for industry-academia collaborations (Litten et al., 2016). Although outside of the scope of the present review, excellent reviews and commentaries from NIAAA scientists have outlined the necessity of industry-academia collaborations in medication development for AUD (Litten et al., 2012a, Litten et al., 2014a, Litten et al., 2014b, Litten et al., 2016).

One of the largest issues contributing to the “valley of death” is the lack of infrastructure to support the transition from animal to human testing in academia. Further, even if a compound makes it past this valley, most researchers lack the financial resources, regulatory expertise, and marketing resources to move the medication through costly clinical trials and into the post-approval market. Partnerships with industry remove, or at least mitigate, many of these issues, and for the foreseeable future it is highly unlikely that any medication will receive FDA approval without collaboration with pharmaceutical companies. However, even with industry support, a remaining barrier is a minimally articulated set of goals and methods for each stage of human testing. Specifically, the requirements to successfully transition from safety testing in humans, to early efficacy detection in clinical samples, and finally to confirming clinical efficacy remain unclear, despite a great deal of interest and substantial investments in medications development for AUD. This article seeks to draw parallels and lessons from the well-established stage model of behavioral therapies research for alcohol and drug abuse and to discuss its application to research on pharmacological

treatment development. In this paper, we provide theoretical and empirical support for the standard medication development practices that human laboratory models and/or pilot randomized controlled trials should serve as intermediaries in the transition from pre-clinical studies to large, and costly, randomized controlled efficacy trials. The relative strengths and weaknesses of pilot clinical trials versus human laboratory studies for bridging the “valley of death” are discussed and explored via a Monte Carlo data simulation study. The overarching goal of this manuscript is to shed light on the optimal approach to efficiently and reliably translate preclinical findings on promising medications for AUD to human clinical populations suffering from the disorder.

Stages of Medications Development

In an influential manuscript setting an agenda for medications development for AUD, Litten and colleagues (2012) outlined a set of objectives for the next decade of research (Litten et al., 2012a). They describe stages for both preclinical testing and clinical testing in drug development. Litten *et al* particularly emphasize the “valley of death” where the majority of medications that show promise in animal models never make the transition to human subjects testing. This obstacle to medications development persists despite many well-validated methods for preclinical testing of AUD pharmacotherapies (Egli, 2005). Next, we address why medications fail to be shepherded across this valley and propose potential solutions informed by behavioral therapies research.

Clinical testing for a pharmacological treatment consists of multiple stages. Phase 1 is comprised of safety, pharmacokinetics (pK), dose selection, and alcohol interaction studies in non-treatment seeking samples. This early stage of human testing is sometimes broken down into Phase 1a and 1b where Phase 1a focuses on safety, pK, and alcohol interaction studies and Phase 1b tests initial efficacy in the human laboratory with either single or multiple medication doses. Phase 2 is comprised of early efficacy studies in the human laboratory as well as safety and dose finding. Like Phase 1, Phase 2 efficacy testing can be described as early Phase 2 (referring to human laboratory testing in non-treatment seeking samples) and late Phase 2 (referring to initial randomized controlled trials in treatment seeking samples). Notably, Phase 1b and early Phase 2 overlap considerably in terms of goals and methods. Late Phase 2 is specifically defined as an efficacy determination phase through initial randomized controlled trials with treatment-seeking individuals. Phase 3 consists of verifying efficacy and safety through placebo-controlled randomized clinical trials with treatment-seeking individuals and possibly through large scale multi-site trials that support a new drug application to the FDA. Finally, Phase 4 is defined by monitoring safety, implementation, and adoption of a novel treatment after FDA approval during post-market safety monitoring. Regardless of the preferred nomenclature for the stage model of medications development, there is a general recognition that clinical testing should progress from first determining safety, to confirmation of both clinical efficacy and safety, to subsequent regulatory approval.

While the clinical trial stages of medication development are fairly well defined, the outcomes needed to advance a medication through early phases are ambiguous, and the optimal primary treatment endpoint for late stage trials has been debated for decades. For

pilot and laboratory studies conducted in the context of Phase 1 and early Phase 2 trials, the question of “What outcomes determine safety and efficacy?” has no precise, well-defined answer at this time. Instead, researchers are left to decide which outcomes are most likely to detect safety and efficacy signals in order to increase the probability of moving a medication to late phase trials. The problems with this approach to medication development are obvious: there is little consistency and consistency between studies in both the measures being administered and variables being assessed. Conversely, the outcome measures for assessing efficacy in Phase 3 trials are well defined. The FDA accepts either complete abstinence or no heavy drinking days as clinical trial endpoints in order for a medication to be considered for approval. Although both abstinence and low-risk drinking (i.e., consuming alcohol without experiencing a heavy drinking event) after treatment are stable behaviors and associated with positive long-term health outcomes (Kline-Simon et al., 2017, Kline-Simon et al., 2013, Witkiewitz et al., 2017b, Witkiewitz et al., 2017a, Kline-Simon et al., 2014), recent evidence suggests that both endpoints may be overly strict and may not capture all individuals who respond to a medication.

The Alcohol Clinical Trials Initiative (ACTIVE) Workgroup and affiliated researchers have re-examined several of the largest AUD clinical trials in order to address key issues associated with medication development in this field (Anton et al., 2012). Most relevant to this review, several of such studies have evaluated potential alternate Phase 3 trial endpoints or at least extensions of the current FDA-recommended endpoints. Individuals who are able to reduce their alcohol consumption to low-risk levels during treatment do not substantially differ from abstainers in terms of health care utilization or medical costs and are able to sustain this reduction in alcohol consumption over several years (Kline-Simon et al., 2017, Kline-Simon et al., 2014, Witkiewitz et al., 2017a, Witkiewitz et al., 2017b). Furthermore, in a re-analysis of the COMBINE study, individuals who received naltrexone and still reported some heavy drinking days by the end of the study had fewer heavy drinking days than those who received placebo (Falk et al., 2014), consistent with the notion that non-abstainers benefited from naltrexone in COMBINE (Ray et al., 2010b). Similarly, other studies have found that individuals who report limited heavy drinking during treatment have comparable psychosocial functioning to those who report abstinence or non-heavy drinking (Witkiewitz et al., 2017a, Witkiewitz et al., 2017b, Wilson et al., 2016). As the individuals described above could be, and often are, categorized as treatment failures by the current FDA-recommended outcomes despite showing beneficial responses to pharmacotherapy, less-stringent drinking outcomes and endpoints that account for improvements in quality of life should be considered when evaluating medications for AUD.

One measure of alcohol consumption that has received recent attention as a promising alternative trial endpoint is the World Health Organization (WHO, International Guide for Monitoring Alcohol Consumption and Related Harm. Geneva, Switzerland, 2000) risk levels of alcohol use (very high risk, high risk, moderate risk, and low risk defined by mean ethanol consumption in grams per day). The European Medicines Agency (EMA), which is Europe’s equivalent of the FDA, allows for a two-level categorical shift in WHO risk levels (e.g., high risk to low risk) as an endpoint for medication evaluation in addition to complete abstinence, heavy drinking days, and total alcohol consumption (2010, Guideline on the Development of Medicinal Products for the Treatment of Alcohol Dependence). For

example, WHO risk level reduction was used by the EMA to evaluate, and eventually approve, nalmefene as a treatment for AUD (EMA, Selincro Assessment Report, 2012). Recent studies from the ACTIVE workgroup has supported the validity of WHO risk levels as endpoint in US studies as well. In another re-analysis of the COMBINE study, reduction in WHO risk level during treatment was predictive of a lower level of alcohol-related consequences and mental health problems at the end of one year after treatment (Witkiewitz et al., 2017a). These findings were supported by an examination of National Epidemiologic Survey on Alcohol and Related Conditions cohort data which found that a reduction in WHO risk level was associated with an attenuated risk of alcohol dependence (Hasin et al., 2017). As others have discussed, the addition of WHO drinking risk levels as an endpoint in US clinical trials could improve medications development not only by assessing another valid and ecologically important marker of medication efficacy but also by facilitating medication development for AUD globally through the creation of common assessment instruments (Johnson, 2017b, Johnson, 2017a, Litten et al., 2017).

While improving our ability to detect medication effects via implementing additional drinking endpoints in late stage (e.g., Phase 3) clinical trials would certainly increase the likelihood of developing new and better pharmacotherapies for AUD, this step would not necessarily affect the issues presently interfering with the success of Phase 1 and 2 trials. Thus, the current review focuses on clinical testing and ways to optimize the process of establishing safety and initial efficacy using both human laboratory and randomized clinical trial designs. In fact, one of the priorities identified by Litten *et al.* (2012) is to develop and implement efficient screening models for both animal and human laboratory paradigms. In our own work, we have written about the role of human laboratory models of AUD in medications development (Ray et al., 2010a, Yardley and Ray, 2016, Plebani et al., 2012). *Simply put, the central question to be addressed in early efficacy testing is whether a given pharmacotherapy shows sufficient evidence of early efficacy to warrant further clinical testing through randomized controlled clinical trials.* The extent to which the human laboratory paradigms currently available can provide the needed evidence to make these important go/no-go decisions about pharmacotherapies for AUD remains unestablished. Our recent qualitative review of the literature showed little consilience between findings from human laboratory models and findings from clinical trials (Yardley and Ray, 2016). Furthermore, selective reporting biases for laboratory trials which often measure dozens of outcomes likely result in highly inflated estimates of laboratory efficacy (Simmons et al., 2011) and undermines the translational validity of human laboratory methods. Nevertheless, quantitative approaches, such as a meta-analysis, are required to directly address the question of whether human laboratory findings predict clinical trial outcomes; although to our knowledge, valid meta-analytic approaches for testing this translational question have yet to be articulated in the literature. Indeed, this question is particularly complicated and compounded by the fact that the predictive ability of laboratory measures may be specific to a mechanism of action and/or pharmacological target of the compound being tested. Questions about the translational validity of human laboratory methods are amplified for preclinical research where methodological differences are compounded by large biological differences in the test subjects between animal models and clinical subjects (Egli, 2018). While data from different sources and methods using different medications do not lend

themselves neatly to integration with current meta-analytic methods, additional efforts to quantify the predictive utility of these models are clearly warranted to test whether laboratory outcomes predict clinical success.

In the field of AUD medications development, we have seen compelling examples of proof-of-concept human laboratory studies that were positive and were followed by a positive result in a randomized clinical trial (RCT). For instance, McKee et al. (2009) utilized a laboratory-based self-administration paradigm which found that varenicline significantly reduced alcohol craving, subjective response to alcohol, and alcohol consumption in heavy drinking smokers (McKee et al., 2009). This initial evidence of efficacy for face-valid laboratory phenotypes was later followed by several small clinical trials (Fucito et al., 2011, Mitchell et al., 2012, Plebani et al., 2013) and a successful RCT which advanced varenicline as an efficacious treatment for alcohol dependence (Litten et al., 2013). However, the field has also seen promising medications based on human laboratory (Ray et al., 2011) and initial clinical trials (Kampman et al., 2007) fail in larger scale confirmatory trials (Litten et al., 2012b). These examples highlight the inconsistent fashion with which results from early efficacy trials, mostly conducted in the human laboratory, relate to clinical outcomes in RCTs, the gold-standard for evaluating and ultimately approving novel treatments for AUD.

In recognizing the role of early efficacy studies for medications development, and in particular, of human laboratory studies of AUD pharmacotherapies, the next step is to clearly articulate a set of goals for early efficacy phases of research. To that end, we contend that a useful approach would be to review the literature on the mature field of behavioral therapies development for Substance Use Disorders (SUD). While both fields incorporate a stage model of development, the behavioral therapies approach has been established and utilized for two decades (Rounsaville et al., 2001). Therefore, a critical comparison of these models may yield important insights that can, in turn, inform a more efficient and effective course for novel compounds in medications development for AUD.

Stages of Behavioral Therapies Development

The field of behavioral treatment was revolutionized by the advancement of the technology model of research implemented in the National Institute of Mental Health (NIMH) Treatment of Depression Collaborative Research Program (Elkin et al., 1985). This framework provided prescriptive methods for how behavioral techniques should be evaluated, including issues of therapist fidelity and trial monitoring. Building on the technology model of behavioral therapy development and recognizing the large burden on investigators to conduct full-scale efficacy trials, Onken et al. (1997) proposed a Stage Model of Behavioral Therapies research (Onken et al., 1997). Of note, the stage model of behavioral therapies development was preceded by the stage model of pharmacotherapy development, and intended to parallel NIDA's medications development program (<https://grants.nih.gov/grants/guide/pa-files/PA-94-078.html>). In this model, Stage 1 consists of pilot/feasibility testing, which includes research activities such as manual writing, training therapists, and adherence measurement of the novel treatment. Stage 1 is further divided into two substages, Stage 1a is described as therapy development and manual writing while Stage 1b consists of a pilot trial with the goal of estimating a treatment effect size (Rounsaville et

al., 2001). It is not uncommon for Stage 1a to be associated with a small open label trial to assess the acceptability and feasibility of the novel treatment, prior to enrolling participants into the pilot randomized controlled trial (Stage 1b). In a more recent description of the stage model of behavioral therapy development, Carroll and Onken (2005) emphasize the notion that Stage 1 research provides the opportunity for creativity and innovation in clinical behavioral science by allowing researchers to develop novel behavioral therapies and/or improve existing treatments (Carroll and Onken, 2005). Further, treatment research at this stage has the potential for a high yield with regard to evaluating clinical strategies that have not yet undergone empirical testing. This is also an opportunity to integrate advances in basic neuroscience and basic research on affective, cognitive, and social factors, including brain imaging research into Stage 1a and 1b research (Carroll and Onken, 2005). Stage 2 consists of RCTs to evaluate the efficacy of the pilot-tested treatments and to address mechanistic questions, particularly regarding the mechanisms of action of treatments with established efficacy. Stage 3 consists of studies to evaluate transportability of treatments, including opportunities to make these treatments more usable in community treatment settings (Carroll and Onken, 2005). One notable difference between behavioral treatment development and medications development is that all stages of behavioral treatment development are typically conducted with treatment-seeking samples, likely because fear of adverse events in a vulnerable population are relatively minimal.

The stage model of behavioral therapy research is the ‘gold standard’ in the field of mental health, including addiction, and this has been facilitated by a funding mechanism within the National Institutes of Health dedicated to behavioral therapies development. Specifically, the R34 mechanism supports those research activities proposed for Stage 1 of the behavioral treatment development model. An added advantage of the R34 mechanism is that it allows for 3 years of research support to conduct the proposed research, typically incorporating aspects of Phase 1a and 1b. We are especially interested in the Stage 1 of behavioral therapies research given that it parallels Phase 1 for medications development, namely safety and initial efficacy testing in human subjects (also termed Phases 1a and 1b). To that end, behavioral therapies research has provided concrete guidelines as to what elements are required at each stage of treatment development and how those steps can be integrated into research proposals (Rounsaville et al., 2001). This approach is highly consistent with the technology model of behavioral therapy and has implications for how these therapies are later evaluated for the strength of the evidence of their efficacy (i.e., evidence-based treatments approach) (Chambless, 2015).

In summary, a review of the stage model of behavioral treatment development suggests a prescriptive approach for Stage 1 of treatment development, informed largely by the technology model, associated with a discrete mechanism of NIH support (i.e., R34) and that accommodates a range of clinical research methods and seeks to nurture creativity and innovation. While this model was preceded and influenced by the medications development stage model, the flexibility and clear delineation of research activities at each stage afforded by the behavioral therapies model may in turn be used to refine its predecessor, namely the stage model of medications development. Next, we consider how the behavioral therapies stage model of treatment development could inform an actionable model for bridging the “valley of death” in AUD medications development.

Recommendations for AUD Medications Development

The goal of this article is to draw parallels from the well-established stage model of behavioral therapies research for SUD and to discuss its application to research on medications development for AUD. Given the description of each separate model for medications and behavioral therapies development, we take this opportunity to envision how aspects of the behavioral therapies treatment development model may be applied to medications development.

One aspect of the Stage 1 research in behavioral therapies development that is particularly intriguing is the notion of dividing early development functions into Stage 1a and 1b. The medications development model parallels this distinction with targets of safety (1a) and initial efficacy (1b). However, since establishing a safe medication $\times$ alcohol interaction is a necessary component of demonstrating that a medication is safe for AUD, laboratory-based alcohol administration studies will often assess safety concurrently with analogues of clinical efficacy such as alcohol craving and subjective responses to alcohol (e.g., Ray et al., 2017a, Ray et al., 2011, Roche et al., 2016, Hutchison et al., 2001). Whether this is a better approach than separating safety and early efficacy testing remains up for debate. The behavioral therapies model would argue that there should be a sequence to the 1a and 1b phases, with one informing the other. In fact, if the acceptability of a behavioral treatment is found to be problematic during the open label pilot, changes are made to the manual and to the treatment delivery to address those issues (e.g. dosing schedule, compounding, etc. in medications development). Separating safety and early efficacy may have an important benefit of allowing the safety trial to generate hypotheses for testing under optimal conditions during Stage 1b activities.

Pilot Clinical Trials versus Laboratory Studies

Another distinction between the development of behavioral therapies compared to medications is the fact that in behavioral therapies development, pilot randomized controlled trials (RCTs) are the gold-standard for early efficacy testing, whereas in medications development, human laboratory models are often the preferred approach to screening novel therapeutics. As noted previously, there is a rich literature supporting the use of experimental psychopharmacology paradigms in AUD etiology and treatment development (Bujarski and Ray, 2016). An important distinction between behavioral pharmacology studies of AUD medication and clinical trials for AUD is the nature of the sample, with the former being comprised of non-treatment seekers and the latter comprised of treatment seeking individuals. Our group (Ray et al., 2017b) and others (Rohn et al., 2017) have recently highlighted differences between treatment seeking and non-treatment seeking samples on a host of relevant demographic and clinical features. In addition, we have argued that one of the ways to promote consilience between human laboratory models and clinical trials may be to engage a similar population of treatment-seeking individuals (Ray et al., 2017b). To that end, using pilot randomized controlled trials with participants who are actively seeking treatment for AUD to screen medications for early efficacy would address concerns associated with sample discrepancies as well as with outcome discrepancies. Concerns about safety with testing medications in affected and thus potentially vulnerable populations represent a central ethical concern in clinical research. To promote efficient medications

development, we thus recommend that researchers carefully weigh the two opposing considerations of external validity and safety in designing early clinical trials of novel medication.

One important consideration with pilot RCTs concerns the small sample sizes inherent to such studies and particularly whether it is appropriate to use pilot studies to generate effect size estimates for novel treatments. Recent discussion about this issue has suggested that estimating effect sizes from pilot studies is unsupported from a methodological viewpoint (Leon et al., 2011, Kraemer et al., 2006) and that stable estimation of effect size requires hundreds of participants to be enrolled (Schönbrodt and Perugini, 2013). Since laboratory studies are typically able to enroll more participants and minimize statistical noise through greater experimental control, the methodological concerns about estimating efficacy are somewhat mitigated. With this caveat, however, it is clear that pilot studies have a lot to offer the literature, including evaluating the feasibility of recruitment, randomization, retention, assessment procedures and implementation of the novel intervention. Pilot RCTs can also identify modifications needed in the design of a larger hypothesis-testing trial (Leon et al., 2011). In summary, the pilot RCT approach may be better suited at setting the stage for the larger scale efficacy trial (Phase 2) than to establish the initial efficacy of the novel compound. Insofar as grant reviewers and stakeholders have proper expectations for the results of pilot studies, they can play a crucial role in medications development for AUD.

Another important consideration is the fact that investigators may continue to use human laboratory methods to conduct Phase 1b activities in medications development, namely gauging initial efficacy. To the extent that human laboratory models of AUD are suited for medication screening, they can continue to have a meaningful role in medications development. The use of human laboratory models in medications development for AUD can be bolstered by a set of standardized procedures with common methods and endpoints that can be implemented reliably across research groups. To that end, NIAAA has established the NIAAA Human Laboratory Program (HLAB), designed to screen compounds for effectiveness using human laboratory paradigms. The experimental paradigms derived from this important initiative can in turn be validated and disseminated as standards for screening of novel compounds.

A critical direction in using human laboratory models in medications development relates to the identification and refinement of paradigms that have high external validity for predicting clinically-relevant (and clinical trials-related) outcomes. In other words, there is a pressing need to develop valid and efficient methods for shepherding novel compounds from initial discovery and safety testing through RCTs and ultimately approval/dissemination. While traditional human laboratory methods have been proposed to serve these aims, at present no human laboratory method has demonstrated translational validity. The degree to which human laboratory outcomes predict AUD clinical trial outcomes remains hotly debated yet mostly untested (Yardley and Ray, 2016, Litten et al., 2016, Ray et al., 2016). In short, human laboratory studies have been measuring the same core of efficacy outcomes (e.g., how a medication affects subjective response to alcohol, alcohol cue-induced craving, alcohol self-administration, etc.), albeit with improving measurement precision, for decades with minimal advancement or variation. The outcomes from Phase 1 and 2 medication

testing should provide confidence in the success of Phase 3 trials FDA-recommended clinical trial endpoints. Despite the lengthy list of experimental psychopharmacology paradigms used in AUD research (Bujarski and Ray, 2016), as a field we have no evidence that a medication's effect on any of these measures is predictive of that medication's success in Phase 3 trials. Let us examine subjective response to alcohol as an example. The subjective stimulating/rewarding and sedative/aversive effects of alcohol are predictive of AUD development and drinking behavior (e.g., King et al., 2011). Thus, as a logical extension, it has been widely and implicitly accepted without explicit evidence that a medication's ability to block the rewarding effects or potentiate the aversive effects of alcohol in the laboratory transitively confers the ability to reduce hazardous drinking in the real world. Yet, in medication development for other addictive substances, the utility of subjective response to acute drug administration as an outcome measure appears to be limited (Comer et al., 2008, Haney, 2009). For example, the dose at which a medication effectively reduces positive subjective responses to cocaine is not necessarily sufficient to reduce self-administration in the laboratory or drug use in a clinical-trial setting (Haney and Spealman, 2008, Comer et al., 2008). Given the obstacles to medication development discussed in this review, we believe it is imperative that future AUD research employ combined laboratory-longitudinal approaches under the umbrella of late Phase 2 or Phase 3 trials in order to establish the predictive utility of pharmacological manipulation of laboratory-based methods as a robust and reliable indicator of medication efficacy in clinical settings. Additionally, we should seek to validate new human laboratory paradigms that increase confidence in future success in Phase 3 trials. A model seeking to bolster external validity of medication screening in the laboratory has been tested in the smoking literature with early success (Perkins and Lerman, 2014, Perkins et al., 2010, Perkins et al., 2008, Perkins et al., 2006). This model consists of a "practice quit attempt" in which smokers are asked to abstain for one week and to come to the laboratory daily for bioverification of nicotine abstinence and would seemingly translate quite well to AUD samples (Perkins and Lerman, 2014). In brief, the more that the human laboratory paradigm can approximate drinking outcomes, the more likely it is to provide valuable information regarding the initial efficacy of a novel compound.

Simulation Comparing Pilot Trials to Laboratory Studies

To further explore the relative efficiency of pilot trials versus laboratory studies in screening medications for AUD, we conducted a Monte Carlo simulation study. For this simulation study we tested the following parameters: (1) average medication effect size (Cohen's $d = 0.2, 0.5, \text{ and } 0.8$ representing small, medium, and large effects), pilot study sample size (N_{pilot} range 6 to 36), the multiplicative increase in sample size associated with typically less expensive and quicker laboratory studies (Lab Multiple = 1, 2, and 4), and lastly the correlation between clinical and laboratory effect sizes ($\rho_{\text{Lab-Clinic}} = 0.3, 0.6, 0.9$, representing relatively poor, moderate, and strong correlation between clinical and laboratory effects). For each of these parameter combinations we simulated 10,000 clinical effect sizes, half from a null distribution $N(0, 0.2)$ and half from the specified mean effect size $N(d, 0.2)$. Laboratory effect sizes were then simulated based on population correlation $\rho_{\text{Lab-Clinic}}$ with equivalent distributions to the clinical effect sizes. We then calculated the probability of a significant and positive trial (i.e. statistical power with $\alpha = 0.025$, one-tailed)

for each of the simulated effect sizes in both a pilot study of sample size N_{pilot} , and a laboratory study of sample size $N_{\text{pilot}} \times \text{Lab Multiple}$. By summing these powers we calculated the expected number of positive trials for medications with true and false effects (i.e. drawn from distribution $N(d, 0.2)$ or $N(0, 0.2)$ respectively). We then calculated the sensitivity and positive predictive value for each parameter combination. All Monte Carlo simulations were conducted in R by SB. Full R code and all figures are available at www.github.com/sbujarski

Since early efficacy trials are primarily interested in testing whether a treatment shows sufficient evidence of efficacy to warrant a full-scale Phase 3 RCT, we focused primarily on sensitivity or the probability that a medication with a true effect would screen positive. As expected, results showed that sensitivity increased with greater sample size, greater average effect size, and greater correlations between laboratory and clinic effect sizes (See **Figure 1**).

These simulations suggest that a pilot study is superior in terms of sensitivity only when a laboratory study confers no advantage in terms of sample size (i.e. Lab Multiple = 1). However, when a laboratory study has twice the sample size of a pilot study, even a mediocre laboratory paradigm (i.e. $\rho_{\text{LabClinic}} = 0.6$) exhibited greater sensitivity than a pilot trial across the range of sample sizes and medication effect sizes tested. For laboratory studies enrolling four times the sample size, sensitivity was superior to pilot studies even with a poor laboratory analogue ($\rho_{\text{LabClinic}} = 0.3$). However, while laboratory studies had generally greater sensitivity, pilot studies were generally superior to laboratory studies in terms of positive predictive value or the probability that a positive trial is a true positive (**Figure 2**). Since the probability of a statistically significant pilot trial is so low, a significant result is more predictive of a true effect as compared to more highly powered laboratory study which measures a more distal outcome. It is worth noting however, that the effect sizes calculated in small, yet statistically significant studies are likely to be dramatically inflated, and thus should not be used to determine sample sizes for large scale RCTs (Gelman and Carlin, 2014).

In sum, this Monte Carlo simulation found that a laboratory study with a paradigm well-calibrated to capture meaningful clinical effects and do so with a larger sample size is better able to detect true positive medication effects. However, since the concordance between laboratory and clinical effects is not perfect, positive signals are less likely to be predictive of a true positive medication effect than the effects obtained in pilot trials. Furthermore, these simulations reinforce the conclusions of Leon et al. (2011) and Kraemer et al. (2006) that small pilot trials are ill-equipped to estimate effect sizes, or even engage in hypothesis testing of treatment efficacy. Laboratory studies that enroll more subjects may be better positioned to engage in hypothesis testing, but this assumes that the laboratory outcomes correlate with clinical efficacy, an assertion that is currently unproven in the field. Lastly, it should be noted that results of this simulation, while intended to inform medication testing for AUD, would be generalizable and applicable to medication development for other psychiatric and medical conditions that use the FDA Clinical Research Phases.

Structural Support for Early Phase Testing

Upon reading the series of stages in behavioral treatment development, pharmacology researchers may quickly raise issues of time and feasibility. Phase 1 activities are very involved and time consuming in the context of medications development, although still less in comparison than Phase 2 and 3 testing. The recognition of the burden on investigators to implement clinical trials within the framework of the technology model led NIMH to recognize manual development and initial open label testing as important activities that require time and support. To that end, the 3-year scope of the R34 mechanism may provide a unique set of benefits to the behavioral therapies approach, which are currently unavailable to medications development. This is particularly relevant with increasing recognition of the regulatory burden associated with medications development, including tasks such as obtaining and managing an Investigational New Drug (IND) application to the FDA, implementing good clinical practices across all study procedures, and establishing and maintaining regular and productive communications with various stakeholders such as pharmaceutical companies, IRBs, university entities, clinical research centers, and research participants. Taken together, these regulatory and research activities argue for increased support for Phase I activities in medications development as a way to overcome the “valley of death” and to more effectively move promising compounds from preclinical to clinical studies.

Conclusions

Medications development for AUD is a high research priority area with great potential to have a meaningful public health impact. As with other fields of science, developing novel therapeutics for AUD is not without a host of obstacles. A notable obstacle in medications development for AUD has been the transition from pre-clinical testing to human testing, which has been termed the “valley of death,” given that very few promising therapeutics from pre-clinical studies are actually tested in humans. In order to overcome this notable obstacle, this review focuses on the first steps in human testing, particularly studies of safety and initial efficacy of a novel compound. Regulatory issues aside, we discuss ways in which initial efficacy testing can take place and draw upon the literature on behavioral therapies development, which more clearly prescribes a pilot randomized controlled trial as an intermediate step between the development of a novel behavioral treatment and an efficacy trial (i.e., RCT). In the behavioral pharmacology tradition, however, human laboratory endpoints (e.g., alcohol craving and subjective response to alcohol) are often used as indicators of early efficacy. This manuscript contrasts the human laboratory and pilot RCT approaches both in theoretical terms and using a Monte Carlo data simulation approach. The pilot RCT approach is advantageous over the human laboratory approach in terms of ecological validity as well as for determining treatment tolerability and acceptability whereas the human laboratory approach appears to be more cost effective and time-efficient. A major limitation of the pilot RCT approach is that outcomes from small trials should not be used to estimate effect sizes whereas a major limitation of the human laboratory approach is that their outcomes may not be robustly associated with the clinical outcomes we are most interested in. Results of our Monte Carlo simulation found that a laboratory study with a paradigm well-calibrated to capture meaningful clinical effects is better able to detect true

positive medication effects than pilot RCTs. Together, these results argue for the careful “calibration” of human laboratory paradigms as a critical step in overcoming the “valley of death” in medications development for AUD and ultimately promoting the translation of novel compounds to clinical populations. However, structural and funding changes are necessary to address barriers to early stage medication development. In particular, we have highlighted the importance of the R34 mechanism in behavioral treatment development research in supporting scientists in the early stages of human subjects testing which for medications development carries a high regulatory burden including IND applications, developing practice guidelines, coordination with various stakeholders, etc. By providing grant support for these laborious yet necessary steps in medications development the NIH could encourage scientists to shepherd a medication across the “valley of death.”

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Figure 1:

Sensitivity results for pilot versus laboratory screening of novel medications. These sensitivity analyses are the result of a Monte Carlo simulation study ($n = 10,000$ per specification) examining the effect of pilot study sample size, mean medication effect size (Cohen's d), multiplicative increase in laboratory versus pilot clinical trial (Lab Multiple), and correlation between laboratory and clinic effect sizes. Sensitivity was defined as the expected number of positive screens as a proportion of the total number of true positive medications.

Figure 2:

Positive predictive value results for pilot versus laboratory screening of novel medications. These positive predictive values are the result of a Monte Carlo simulation study ($n = 10,000$ per specification) examining the effect of pilot study sample size, mean medication effect size (Cohen's d), multiplicative increase in laboratory versus pilot clinical trial (Lab Multiple), and correlation between laboratory and clinic effect sizes. Positive predictive value was defined as the proportion of positive screens that represent true positives.