

Rational Development of Addiction Pharmacotherapies: Successes, Failures, and Prospects

R. Christopher Pierce¹, Charles P. O'Brien², Paul J. Kenny³, and Louk J. M. J. Vanderschuren^{4,5}

¹Center for Neurobiology and Behavior, Department of Psychiatry, Perelman School of Medicine at the University of Pennsylvania, Philadelphia, Pennsylvania 19104

²Treatment Research Center Department of Psychiatry, University of Pennsylvania School of Medicine, Philadelphia, Pennsylvania 19104

³Department of Molecular Therapeutics, The Scripps Research Institute—Florida, Jupiter, Florida 33458

⁴Rudolf Magnus Institute of Neuroscience, Department of Neuroscience and Pharmacology, University Medical Center Utrecht, Universiteitsweg 100, 3584 CG Utrecht, The Netherlands

⁵Department of Animals in Science and Society, Division of Behavioural Neuroscience, Faculty of Veterinary Medicine, Utrecht University, Yalelaan 2, 3584 CM Utrecht, The Netherlands

Correspondence: rcpierce@mail.med.upenn.edu

There are currently effective, U.S. Food and Drug Administration (FDA)-approved therapies for alcohol, nicotine, and opioid addiction. In some cases these therapeutics were rationally designed and tested using a combination of various animal models of addiction. In many cases, however, effective drug therapies for addiction were derived from the testing of compounds developed for other CNS disorders (e.g., analgesics and antidepressants), which were tested clinically in the absence of prior animal research using addiction models. This article will review the development of eight compounds that are currently most effective in the treatment of alcohol, opioid, and nicotine addiction with an emphasis on pharmacological mechanisms as well as the utility of animal models of addiction in the development of these therapeutics. In contrast to these successes, animal research has identified a number of promising medications for the treatment of psychostimulant addiction, none of which have proven to be effective clinically. This raises questions about the validity of current animal models of psychostimulant addiction. A specific example of an apparently promising pharmacotherapeutic for cocaine addiction (the D1 dopamine receptor antagonist ecapipam) that failed clinically will be examined to determine if this truly represents a challenge to the predictive validity of current models of cocaine addiction. In addition, the development of promising cocaine addiction therapeutics derived from animal research will be reviewed.

R.C. Pierce et al.

The earliest references to animal models of addiction in the literature all referred to work on opioids, mainly morphine. Research using animal models prior to 1960 used the term addiction loosely given that the drug of abuse was experimenter delivered. For example, Plant and Pierce (1928) administered very high doses of morphine to dogs daily for 2–3 months. Indeed, the doses chosen were toxic in some cases as the investigators reported that “two of our animals died in convulsions on doses of 190 and 220 mg. per kilogram.” At that time, the severity of the withdrawal syndrome was thought to be the primary factor contributing to relapse among opioid-dependent humans. Therefore, Plant and Pierce sought to examine and characterize opioid withdrawal in animals. Following cessation of morphine treatment, they noted that “five of our dogs showed marked changes in temperament during the first week of withdrawal, in that they became very cross” and “one animal died in convulsions on the third day of withdrawal” (Plant and Pierce 1928). These authors summed up their observations as follows: “The period of addiction in dogs has given a picture that follows closely the description of addiction in man [including vomiting, constipation, hypersensitivity, scratching, irritability, and decrease in narcotic action of the drug].” Note that Pierce and Plant defined addiction as opioid withdrawal and tolerance (Plant and Pierce 1928).

The first valid animal model of addiction was developed in the early 1960s. As a first step toward developing a model in which animals self-administer drugs of abuse, two water-deprived rhesus monkeys were trained to press a lever to receive intravenous infusions of saline (Clark et al. 1961). The investigators also showed that saline self-administration could be extinguished and brought under stimulus control (i.e., the monkeys would lever press for light cues previously paired with the saline infusions) (Clark et al. 1961). Subsequently, it was shown that rats (Weeks 1962) and monkeys (Thompson and Schuster 1964) would self-administer morphine intravenously.

Although results from the self-administration paradigm have contributed significantly to

the development of more recent addiction pharmacotherapies, the first successes came from drugs that were developed for other purposes. For example, methadone, the first truly successful addiction therapeutic, was originally developed at Hoechst in the 1930s as an analgesic. Methadone was tested for analgesic efficacy by scientists at Lilly and Wellcome in the 1940s (Scott et al. 1947; Thorp et al. 1947). In early papers, methadone was sometimes spelled methadon and also was called dolophine. As an aside, it has been erroneously reported that methadone, which was developed in Germany, was originally named after Hitler (i.e., adolophine). Then as now, a major goal of opioid research was to identify effective analgesics that lacked an addiction liability. Therefore, methadone was tested by a group of prominent behavioral pharmacologists whose findings were summed up as follows: “we believe that unless the manufacture and use of methadon are controlled addiction to it will become a serious public health problem” (Isbell et al. 1947). This conclusion was based in part on reports from their patients (recovering addicts) such as: “That is great stuff. I wouldn’t have believed it possible for a synthetic drug to be so like morphine. Can you get it outside? Will it be put under the narcotic laws? I wish I could get it to kick my next habit.” The investigators also noted that “methadon prevented the appearance of signs of physical dependence in 12 men who had been proved to be addicted to morphine” (Isbell et al. 1947). These observations suggested that methadone might be used to treat opioid addiction. This idea was not tested until the 1960s, with the publication of the landmark study by Dole and Nyswander (1965), which showed that methadone relieved “narcotic hunger” and produced tolerance such that the euphoric effect of heroin was substantially blunted.

Although methadone and morphine are both full μ opioid receptor agonists with substantial addiction liabilities, methadone is a preferable addiction therapeutic because of a substantially longer half-life and higher oral bioavailability. Methadone distribution is restricted to clinics to ensure that the drug is taken orally, which obviates withdrawal and maintains tolerance in the absence of euphoria.

Methadone is not prescribed for home use because of the legitimate fear that, in the absence of monitoring, the drug will be solubilized and administered intravenously, thereby producing a high roughly equivalent to morphine or heroin administered by the same route. In the case of methadone, the clinical trial came before animal studies. Over the years, a number of animal studies have confirmed that methadone prevents opioid withdrawal and blunts relapse in animal models of craving (Goode 1971; Jones and Prada 1977; Negus 2006).

Although there is no question that methadone has proven to be effective in the treatment of opioid addiction, there are several problems with the methadone clinic model. Primarily, the distance to the closest clinic may render daily clinic visits unfeasible. For some patients living in close proximity to a clinic, the stigma associated with daily visits to a methadone clinic reduces compliance. A clever pharmacological strategy was recently developed to produce an opioid addiction therapeutic that could be taken at the convenience of the patient. It was noted that opioid agonists have good oral bioavailability, whereas the opioid receptor antagonist naloxone does not. Thus, if a pill contains both compounds and is taken orally, the opioid receptor agonist effect predominates. In contrast, if the therapeutic is administered intravenously the antagonist would block the agonist effect. This strategy led to the development of Suboxone, which is a combination of buprenorphine and naloxone. Buprenorphine, which was developed by Reckitt and Colman in the 1970s as an analgesic, was chosen over methadone because it is a partial μ opioid receptor agonist which, in contrast to methadone, has a low instance of death associated with overdose (Mendelson and Jones 2003). Suboxone was approved by the FDA for the treatment of opioid addiction in 2002 and rapidly became the first-line treatment for opioid addiction.

ACAMPROSATE FOR ALCOHOL ADDICTION

Acamprosate, a homotaurine derivative, was developed in France in the 1980s. The rationale was that because alcohol activates GABA_A receptors and homotaurine is a GABA receptor

agonist, perhaps homotaurine derivatives might serve as alcohol replacement therapies (Boismare et al. 1984). Indeed, calcium bis acetylhomotaurine (acamprosate) significantly reduced the voluntary intake of alcohol by rats (Boismare et al. 1984). Based on this result, acamprosate was tested in a double-blind placebo-controlled clinical trial with the success criterion defined as alcohol abstinence following three months of outpatient treatment. Results indicated that 20 of 33 patients receiving acamprosate remained alcohol-free compared to 12 of 37 subjects receiving placebo (Lhuintre et al. 1985). The efficacy of acamprosate as an effective alcoholism therapeutic has been repeatedly replicated (Mason and Heyser 2010). Acamprosate has been used for the treatment of alcoholism in Europe since 1989. Surprisingly, the precise mechanism of action of acamprosate remains unclear. Because of the ambiguity of the drug's mechanism of action, the U.S. FDA delayed approval of acamprosate (marketed as Campral) until 2004. In the U.S., acamprosate is currently the first-line pharmacotherapy for alcoholism.

Although acamprosate was targeted for the treatment of alcoholism because of presumed effects on GABA and taurine transmission, the therapeutic effects of this drug appear to be due primarily to effects on glutamate systems. Although initial reports suggested that acamprosate is an NMDA receptor antagonist, subsequent work indicated that acamprosate acts as a partial agonist at spermidine sites on NMDA receptors (Dahchour and De Witte 2000). More recent evidence revealed that acamprosate also is an mGluR5 receptor antagonist (De Witte et al. 2005). Alcohol withdrawal is associated with a number of changes in neurotransmission including, notably, increased glutamate transmission in regions of the CNS (Mason and Heyser 2010). A growing body of evidence is consistent with the notion that acamprosate blunts alcohol craving and withdrawal by normalizing glutamate transmission (Heilig and Egli 2006).

NALTREXONE FOR ALCOHOL ADDICTION

In the early 1970s, receptor binding assays were used to show that "narcotic antagonists" such as

R.C. Pierce et al.

naloxone bind to specific receptors in the brain. Opioid receptor antagonists including naltrexone and naloxone were subsequently tested as pharmacotherapies for opioid addiction (Martin et al. 1973). The discovery of endogenous opioids and their receptors prompted research into the potential role of opioid peptides in the effects of many drugs including alcohol. Initial reports suggested that naltrexone and naloxone reduced alcohol preference in hamsters and rats (Ross et al. 1976) and attenuated alcohol reinforcement in rhesus monkeys (Altshuler et al. 1980). Based on these and similar findings, a placebo-controlled, double blind clinical trial examining the effect of naltrexone on alcohol relapse was performed. Results indicated that naloxone cut the relapse rate approximately in half compared to controls (Volpicelli et al. 1992). These results were rapidly replicated, with the highest rates of abstinence observed in patients who received both naltrexone and supportive therapy (O'Malley et al. 1992). Numerous trials subsequently showed that naltrexone is effective in the treatment of alcoholism.

A single nucleotide polymorphism at A118G (Asn40Asp) in exon I of the μ opioid receptor was identified and shown to triple the potency of β -endorphin at these receptors (Bond et al. 1998). This polymorphism was shown to be associated with alcohol addiction (Bart et al. 2005) and individuals with one or two copies of the Asp40 allele treated with naltrexone had significantly lower rates of relapse than patients homozygous for the Asn40 allele (Oslin et al. 2006). Thus, naltrexone treatment of alcoholism is one of the few examples of a pharmacogenomic therapeutic. Naltrexone, which is marketed as Revia and Depade, has been used in the treatment of alcoholism since 1995. In 2006, an extended-release naltrexone formulation (Vivitrol) was approved by the U.S. FDA for the treatment of alcohol dependence.

Disulfiram also is used in the treatment of alcoholism. However, this drug does not specifically target aspects of addiction or withdrawal. Rather, disulfiram blocks aldehyde dehydrogenase resulting in the accumulation of acetaldehyde after alcohol ingestion, which produces an array of aversive symptoms. Thus, disulfiram

acts as a punishing agent in the event of relapse rather than a therapeutic. Although disulfiram continues to be used in the treatment of alcoholism, there are concerns related both to the safety and effectiveness of this compound (Heilig and Egli 2006).

NICOTINE, VARENICLINE, AND BUPROPION FOR NICOTINE ADDICTION

The history of nicotine replacement therapies dates to a letter Dr. Claes Lundgren, a physiology professor at Lund University, sent to his friend Ove Fernö at Aktiebolaget Leo pharmaceutical company in 1967. Lundgren and his colleague, Stephan Lichtneckert, suggested oral nicotine as a substitute for tobacco based on the observation that sailors sometimes switched from smoking to chewing tobacco without difficulty when assigned to submarine duty. Fernö immediately recognized the promise and commercial potential of nicotine replacement and embarked on a research program to design a means to orally administer nicotine with delayed absorption. The result was nicotine gum. Years later Fernö reflected on his work as follows: "Putting nicotine into chewing gum is not an invention. Fixing the nicotine to an ion exchange resin and putting that into a chewing gum to enable the chewer to control the rate of release—that is an invention" (Fernö 1994). Initial clinical trials performed in Sweden (Fernö 1973) and London (Russell et al. 1976) in the 1970s indicated that nicotine gum was effective in reducing nicotine withdrawal and maintaining smoking abstinence. A landmark randomized double blind placebo controlled clinical trial published in 1982 indicated that smoking abstinence was 47% in the nicotine gum group compared to a 21% success rate among controls (Jarvis et al. 1982). These results led to the approval of nicotine gum, which Aktiebolaget Leo (now McNeil SB) named Nicorette, by the U.S. FDA in 1984. Although nicotine gum remains the most popular form of nicotine replacement therapy, nicotine lozenges, patches, nasal sprays, and inhalers (including so-called electronic cigarettes) also are effective in maintaining abstinence from tobacco use (Polosa and Benowitz 2011).

Animal studies are notably absent in the history of Nicorette. More recently, a rational strategy led to the development of the partial nicotinic receptor agonist varenicline for smoking cessation. Nicotinic acetylcholine receptors (nAChRs) are pentameric ligand-gated ion channels composed of combinations of at least seventeen different subunits (Pierce and Kumaresan 2006). A number of studies indicate that $\alpha 4\beta 2$ nAChRs, which are the most widely expressed subtypes in the brain, play a critical role in nicotine-induced dopamine release and reinforcement (Mineur and Picciotto 2008). The $\alpha 4\beta 2$ nAChR partial agonist varenicline was developed by Pfizer for smoking cessation. The rationale being that varenicline might serve the dual purpose of moderately increasing mesolimbic dopamine levels, which are reduced during nicotine withdrawal, and also blunt nicotine-induced dopamine release in the event of relapse (Coe et al. 2005). Animal studies revealed that varenicline reduced nicotine-induced dopamine release in the nucleus accumbens (Coe et al. 2005) and inhibited nicotine self-administration as well as the reinstatement of nicotine seeking (O'Connor et al. 2010). A randomized double blind clinical trial showed that the smoking abstinence rate with varenicline was 44% compared with nearly 18% for placebo (Gonzales et al. 2006; Jorenby et al. 2006). Notably, measures of nicotine withdrawal and craving also were reduced by varenicline (Gonzales et al. 2006). Based on these results, the U.S. FDA fast tracked approval for varenicline (Chantix) as a smoking cessation drug, which was granted in 2006.

The initial varenicline smoking cessation clinical trials used bupropion as a positive control. The abstinence rate for bupropion was nearly 30%, significantly greater than placebo but substantially lower than varenicline (Gonzales et al. 2006; Jorenby et al. 2006). Bupropion, which was developed by Burroughs Wellcome (now GlaxoSmithKline), was approved for the treatment of depression by the U.S. FDA in 1985. A sustained release formulation of bupropion, marketed as Wellbutrin SR, remains a highly successful antidepressant. Antidepressants were tested as smoking cessation agents because cig-

arette smokers have higher rates of depression that may be exacerbated by nicotine withdrawal (Glassman et al. 1990). Double blind placebo controlled clinical trials revealed that a sustained-release formulation of bupropion significantly increased the rate of smoking cessation (Hurt et al. 1997). Bupropion was approved for the treatment of nicotine addiction in 1997 and is marketed as Zyban. Similar to some other antidepressants, bupropion is a dopamine and norepinephrine reuptake inhibitor. Interestingly, bupropion also is a nAChR antagonist at various subtypes including $\alpha 4\beta 2$ (Slemmer et al. 2000), which may account for the effectiveness of bupropion as a smoking cessation agent relative to other antidepressants.

PSYCHOSTIMULANTS

Despite decades of focused research effort, there are no effective pharmacotherapies for psychostimulant addiction. Indeed, a broad range of drugs targeting multiple CNS transmitter systems have been tested as treatments for psychostimulant dependence without success (Kampman et al. 2005). Drugs that modulate dopaminergic transmission were among the first assessed for the treatment of psychostimulant addiction both in animal studies and clinical trials. Dopamine receptors are classified as either D1-like or D2-like. There was substantial interest in D1-like dopamine receptor antagonists as psychostimulant addiction therapeutics because they lack the sometimes serious extrapyramidal side-effects associated with D2-like dopamine receptor antagonists (Haney and Spealman 2008). Animal and human laboratory studies revealed that acute administration of D1-like dopamine receptor antagonists attenuated the reinforcing effects of cocaine (Romach et al. 1999; Platt et al. 2002). However, clinical use of a D1-like dopamine receptor antagonist requires repeated administrations. Unfortunately, when humans were maintained on the D1-like dopamine receptor antagonist, ecopipam, cocaine self-administration actually increased (Haney et al. 2001). This finding is consistent with results from rhesus monkeys utilizing continuous drug administration (Kleven and

Woolverton 1990) and is likely caused by antagonist-induced increases in D1 dopamine receptor density in the brain (Haney and Spealman 2008). It is important to emphasize that in the case of ecopipam the animal and human data paralleled one another both when the drug was administered acutely and repeatedly.

Fortunately, there are other examples of rationally developed therapeutics for psychostimulant addiction that appear promising. *N*-acetylcysteine (NAC), which is used to treat acetaminophen overdose, has been shown to normalize decreased nucleus accumbens glutamate levels following cocaine self-administration as well as the reinstatement of cocaine-seeking behavior in rats (Baker et al. 2002, 2003). Recent clinical trials show that NAC attenuated cocaine use and decreased desire to use cocaine (LaRowe et al. 2007; Mardikian et al. 2007). Another interesting strategy is the development of cocaine vaccines. Active immunization with a cocaine vaccine attenuated cocaine self-administration as well as the reinstatement of cocaine seeking in rats (Kantak et al. 2000). Clinical data indicate that the cocaine vaccine, TA-CD, produced selective anticocaine antibodies, which blunted the intoxicating effects of cocaine (Haney et al. 2010). In these studies insufficient immune responses to vaccines is a persistent issue, which has led to the development of novel vaccines designed to generate consistently high antibody titers (Wee et al. 2011).

In terms of the validity of animal models of psychostimulant addiction, we note that the preclinical and clinical data are consistent when the animal model is drug self-administration (Haney and Spealman 2008). Moreover, two of the more promising cocaine addiction therapeutics (NAC and cocaine vaccines) were tested primarily with self-administration models of addiction and relapse.

SUMMARY AND CONCLUSIONS

We have reviewed the development of the eight main compounds currently used in the treatment of addictions. As outlined in Table 1, two of these drugs are antagonists, one is a transporter inhibitor, two are full agonists, and three

are partial agonists. Clearly, the most successful treatments for addiction involve receptor stimulation, with the primary goal of obviating drug withdrawal. Agonist therapeutics also reproduce some of the positive aspects of the drug of abuse (e.g., mood elevation), which enhances compliance. Partial agonists are particularly appealing because they blunt the psychoactive effects of the drug of abuse in the event of relapse. Moreover, the risk of overdose is substantially diminished with partial relative to full agonists.

The fact that partial agonists are particularly effective addiction therapeutics raises the question of partial dopamine agonist treatments for psychostimulant addiction (Platt et al. 2002). Surprisingly, there is very little research in this area. Indeed, the only compound tested is the D2-like dopamine receptor partial agonist aripiprazole, which is approved for the treatment of schizophrenia, depression, and bipolar disorder. Aripiprazole reduced cocaine reinstatement in rats (Feltenstein et al. 2007) and decreased the discriminative stimulus properties of amphetamine in a human laboratory study (Lile et al. 2005). Initial clinical trial results have been mixed with one study reporting that aripiprazole reduced cocaine craving and use (Meini et al. 2011). In contrast, another clinical trial showed that aripiprazole increased the self-administration of smoked cocaine, apparently to compensate for decreased subjective effects of cocaine (Haney et al. 2011). These experiments highlight both the promise of partial agonists in the treatment of psychostimulant addiction and the persistent frustration in developing clearly

Table 1. Therapeutics used in the treatment of addiction and their mechanism of action

| Therapeutic | Drug of abuse | Mechanism of action |
|---------------|---------------|---------------------|
| Acamprosate | Alcohol | Partial agonist |
| Buprenorphine | Opioids | Partial agonist |
| Bupropion | Nicotine | Indirect agonist |
| Methadone | Opioids | Agonist |
| Naloxone | Opioids | Antagonist |
| Naltrexone | Alcohol | Antagonist |
| Nicotine | Nicotine | Agonist |
| Varenicline | Nicotine | Partial agonist |

effective therapeutics for stimulant craving and addiction.

REFERENCES

- Altshuler HL, Phillips PE, Feinhandler DA. 1980. Alteration of ethanol self-administration by naltrexone. *Life Sci* **26**: 679–688.
- Baker DA, Shen H, Kalivas PW. 2002. Cystine/glutamate exchange serves as the source for extracellular glutamate: Modifications by repeated cocaine administration. *Amino Acids* **23**: 161–162.
- Baker DA, McFarland K, Lake RW, Shen H, Toda S, Kalivas PW. 2003. N-acetyl cysteine-induced blockade of cocaine-induced reinstatement. *Ann NY Acad Sci* **1003**: 349–351.
- Bart G, Kreek MJ, Ott J, LaForge KS, Proudnikov D, Pollak L, Heilig M. 2005. Increased attributable risk related to a functional μ -opioid receptor gene polymorphism in association with alcohol dependence in central Sweden. *Neuropsychopharmacology* **30**: 417–422.
- Boismare F, Daoust M, Moore N, Saligaut C, Lhuintre JP, Chretien P, Durlach J. 1984. A homotaurine derivative reduces the voluntary intake of ethanol by rats: Are cerebral GABA receptors involved? *Pharmacol Biochem Behav* **21**: 787–789.
- Bond C, LaForge KS, Tian M, Melia D, Zhang S, Borg L, Gong J, Schluger J, Strong JA, Leal SM, et al. 1998. Single-nucleotide polymorphism in the human μ -opioid receptor gene alters β -endorphin binding and activity: Possible implications for opiate addiction. *Proc Natl Acad Sci* **95**: 9608–9613.
- Clark R, Schuster CR, Brady JV. 1961. Instrumental conditioning of jugular self-infusion in the rhesus monkey. *Science* **133**: 1829–1830.
- Coe JW, Brooks PR, Vetelino MG, Wirtz MC, Arnold EP, Huang J, Sands SB, Davis TI, Lebel LA, Fox CB, et al. 2005. Varenicline: An $\alpha 4\beta 2$ nicotinic receptor partial agonist for smoking cessation. *J Med Chem* **48**: 3474–3477.
- Dahchour A, De Witte P. 2000. Ethanol and amino acids in the central nervous system: Assessment of the pharmacological actions of acamprosate. *Prog Neurobiol* **60**: 343–362.
- De Witte P, Littleton J, Parot P, Koob G. 2005. Neuroprotective and abstinence-promoting effects of acamprosate: Elucidating the mechanism of action. *CNS Drugs* **19**: 517–537.
- Dole VB, Nyswander M. 1965. A medical treatment for diacetylmorphine (heroin) addiction: A clinical trial with methadone hydrochloride. *JAMA* **193**: 646–650.
- Feltenstein MW, Altar CA, See RE. 2007. Aripiprazole blocks reinstatement of cocaine seeking in an animal model of relapse. *Biol Psychiatry* **61**: 582–590.
- Ferno O. 1973. A substitute for tobacco smoking. *Psychopharmacologia* **31**: 201–204.
- Ferno O. 1994. Conversation with Ove Ferno. *Addiction* **89**: 1215–1226.
- Glassman AH, Helzer JE, Covey LS, Cottler LB, Stetner F, Tipp JE, Johnson J. 1990. Smoking, smoking cessation, and major depression. *JAMA* **264**: 1546–1549.
- Gonzales D, Rennard SI, Nides M, Oncken C, Azoulay S, Billing CB, Watsky EJ, Gong J, Williams KE, Reeves KR. 2006. Varenicline, an $\alpha 4\beta 2$ nicotinic acetylcholine receptor partial agonist, vs sustained-release bupropion and placebo for smoking cessation: A randomized controlled trial. *JAMA* **296**: 47–55.
- Goode PG. 1971. An implanted reservoir of morphine solution for rapid induction of physical dependence in rats. *Br J Pharmacol* **41**: 558–566.
- Haney M, Spealman R. 2008. Controversies in translational research: Drug self-administration. *Psychopharmacology (Berl)* **199**: 403–419.
- Haney M, Ward AS, Foltin RW, Fischman MW. 2001. Effects of ecopipam, a selective dopamine D1 antagonist, on smoked cocaine self-administration by humans. *Psychopharmacology (Berl)* **155**: 330–337.
- Haney M, Gundersen EW, Jiang H, Collins ED, Foltin RW. 2010. Cocaine-specific antibodies blunt the subjective effects of smoked cocaine in humans. *Biol Psychiatry* **67**: 59–65.
- Haney M, Rubin E, Foltin RW. 2011. Aripiprazole maintenance increases smoked cocaine self-administration in humans. *Psychopharmacology (Berl)* **216**: 379–387.
- Heilig M, Egli M. 2006. Pharmacological treatment of alcohol dependence: Target symptoms and target mechanisms. *Pharmacol Ther* **111**: 855–876.
- Hurt RD, Sachs DP, Glover ED, Offord KP, Johnston JA, Dale LC, Khayrallah MA, Schroeder DR, Glover PN, Sullivan CR, et al. 1997. A comparison of sustained-release bupropion and placebo for smoking cessation. *N Engl J Med* **337**: 1195–1202.
- Isbell H, Wikler A, et al. 1947. Tolerance and addiction liability of 6-dimethylamino-4–4-diphenylheptanone-3 (methadon). *J Am Med Assoc* **135**: 888–894.
- Jarvis MJ, Raw M, Russell MA, Feyerabend C. 1982. Randomised controlled trial of nicotine chewing-gum. *Br Med J (Clin Res Ed)* **285**: 537–540.
- Jones BE, Prada JA. 1977. Effects of methadone and morphine maintenance on drug-seeking behavior in the dog. *Psychopharmacology (Berl)* **54**: 109–112.
- Jorenby DE, Hays JT, Rigotti NA, Azoulay S, Watsky EJ, Williams KE, Billing CB, Gong J, Reeves KR. 2006. Efficacy of varenicline, an $\alpha 4\beta 2$ nicotinic acetylcholine receptor partial agonist, vs placebo or sustained-release bupropion for smoking cessation: A randomized controlled trial. *JAMA* **296**: 56–63.
- Kampman KM, Leiderman D, Holmes T, LoCastro J, Bloch DA, Reid MS, Shoptaw S, Montgomery MA, Winhusen TM, Somoza EC, et al. 2005. Cocaine Rapid Efficacy Screening Trials (CREST): Lessons learned. *Addiction* **100**: 102–110.
- Kantak KM, Collins SL, Lipman EG, Bond J, Giovanoni K, Fox BS. 2000. Evaluation of anti-cocaine antibodies and a cocaine vaccine in a rat self-administration model. *Psychopharmacology (Berl)* **148**: 251–262.
- Kleven MS, Woolverton WL. 1990. Effects of continuous infusions of SCH 23390 on cocaine- or food-maintained behavior in rhesus monkeys. *Behav Pharmacol* **1**: 365–373.
- LaRowe SD, Myrick H, Hedden S, Mardikian P, Saladin M, McRae A, Brady K, Kalivas PW, Malcolm R. 2007. Is

R.C. Pierce et al.

- cocaine desire reduced by N-acetylcysteine? *Am J Psychiatry* **164**: 1115–1117.
- Lhuintre JB, Daooust M, Moore ND, Chretien P, Saligaut C, Tran G, Bosimare F, Hillemand B. 1985. Ability of calcium bis acetyl homotaurine, a GABA agonist, to prevent relapse in weaned alcoholics. *Lancet* **1**: 1014–1016.
- Lile JA, Stoops WW, Vansickel AR, Glaser PE, Hays LR, Rush CR. 2005. Aripiprazole attenuates the discriminative-stimulus and subject-rated effects of D-amphetamine in humans. *Neuropsychopharmacology* **30**: 2103–2114.
- Mardikian PN, LaRowe SD, Hedden S, Kalivas PW, Malcolm RJ. 2007. An open-label trial of N-acetylcysteine for the treatment of cocaine dependence: A pilot study. *Prog Neuropsychopharmacol Biol Psychiatry* **31**: 389–394.
- Martin WR, Jasinski DR, Mansky PA. 1973. Naltrexone, an antagonist for the treatment of heroin dependence: Effects in man. *Arch Gen Psychiatry* **28**: 784–791.
- Mason BJ, Heyser CJ. 2010. Acamprosate: A prototypic neuromodulator in the treatment of alcohol dependence. *CNS Neurol Disord Drug Targets* **9**: 23–32.
- Meini M, Moncini M, Cecconi D, Cellesi V, Biasci L, Simoni G, Ameglio M, Pellegrini M, Forgione RN, Rucci P. 2011. Aripiprazole and ropinirole treatment for cocaine dependence: Evidence from a pilot study. *Curr Pharm Des* **17**: 1376–1383.
- Mendelson J, Jones RT. 2003. Clinical and pharmacological evaluation of buprenorphine and naloxone combinations: Why the 4:1 ratio for treatment? *Drug Alcohol Depend* **70**: S29–37.
- Miner YS, Picciotto MR. 2008. Genetics of nicotinic acetylcholine receptors: Relevance to nicotine addiction. *Biochem Pharmacol* **75**: 323–333.
- Negus SS. 2006. Choice between heroin and food in nondependent and heroin-dependent rhesus monkeys: Effects of naloxone, buprenorphine, and methadone. *J Pharmacol Exp Ther* **317**: 711–723.
- O'Connor EC, Parker D, Rollema H, Mead AN. 2010. The $\alpha 4\beta 2$ nicotinic acetylcholine-receptor partial agonist varenicline inhibits both nicotine self-administration following repeated dosing and reinstatement of nicotine seeking in rats. *Psychopharmacology (Berl)* **208**: 365–376.
- O'Malley SS, Jaffe AJ, Chang G, Schottenfeld RS, Meyer RE, Rounsaville B. 1992. Naltrexone and coping skills therapy for alcohol dependence. A controlled study. *Arch Gen Psychiatry* **49**: 881–887.
- Oslin DW, Berrettini WH, O'Brien CP. 2006. Targeting treatments for alcohol dependence: The pharmacogenetics of naltrexone. *Addict Biol* **11**: 397–403.
- Pierce RC, Kumaresan V. 2006. The mesolimbic dopamine system: The final common pathway for the reinforcing effect of drugs of abuse? *Neurosci Biobehav Rev* **30**: 215–238.
- Plant OH, Pierce IH. 1928. Studies of chronic morphin poisoning in dogs. I. General symptoms and behavior during addiction and withdrawal. *J Pharmacol Exp Therapeutics* **33**: 329–357.
- Platt DM, Rowlett JK, Spealman RD. 2002. Behavioral effects of cocaine and dopaminergic strategies for preclinical medication development. *Psychopharmacology (Berl)* **163**: 265–282.
- Polosa R, Benowitz NL. 2011. Treatment of nicotine addiction: Present therapeutic options and pipeline developments. *Trends Pharmacol Sci* **32**: 281–289.
- Romach MK, Glue P, Kampman K, Kaplan HL, Somer GR, Poole S, Clarke L, Coffin V, Cornish J, O'Brien CP, et al. 1999. Attenuation of the euphoric effects of cocaine by the dopamine D1/D5 antagonist ecopipam (SCH 39166). *Arch Gen Psychiatry* **56**: 1101–1106.
- Ross D, Hartmann RJ, Geller I. 1976. Ethanol preference in the hamster: Effects of morphine sulfate and naltrexone, a long-acting morphine antagonist. *Proc West Pharmacol Soc* **19**: 326–330.
- Russell MA, Wilson C, Feyerabend C, Cole PV. 1976. Effect of nicotine chewing gum on smoking behaviour and as an aid to cigarette withdrawal. *Br Med J* **2**: 391–393.
- Scott CC, Chen KK, et al. 1947. Further observations on the pharmacology of dolophine (methadon, Lilly). *J Pharmacol Exp Ther* **91**: 147–156.
- Slemmer JE, Martin BR, Damaj MI. 2000. Bupropion is a nicotinic antagonist. *J Pharmacol Exp Ther* **295**: 321–327.
- Thompson T, Schuster CR. 1964. Morphine self-administration, food-reinforced, and avoidance behaviors in Rhesus monkeys. *Psychopharmacologia* **5**: 87–94.
- Thorp RH, Walton E, Ofner P. 1947. Analgesic activity in compounds related to amidone. *Nature* **159**: 679.
- Volpicelli JR, Alterman AI, Hayashida M, O'Brien CP. 1992. Naltrexone in the treatment of alcohol dependence. *Arch Gen Psychiatry* **49**: 876–880.
- Wee S, Hicks MJ, De BP, Rosenberg JB, Moreno AY, Kaminisky SM, Janda KD, Crystal RG, Koob GF. 2011. Novel cocaine vaccine linked to a disrupted adenovirus gene transfer vector blocks cocaine psychostimulant and reinforcing effects. *Neuropsychopharmacology* **37**: 1083–1091.
- Weeks JR. 1962. Experimental morphine addiction: Method for automatic intravenous injections in unrestrained rats. *Science* **138**: 143–144.