

ago, Chile. Data were obtained in a stratified sample ($N = 1500$) on demographic characteristics of subjects, benzodiazepine use, and long-term use. Past-year prevalence of use was 31.4% (higher in females, the elderly and 13% long-term users). Forty-five percent of use was over-the-counter. Long-term users were older than users, predominantly females over 40 years. Results show that unrestricted availability of benzodiazepines leads to high use, but long-term use seems to be within rates described for other populations.

THE EFFECTS OF OXAZEPAM ON ANXIETY IN A NONCLINICAL POPULATION. Jennifer F. Landon* and Kenneth J. Sher.† *Kansas City Department of Veterans Affairs Medical Center, Kansas City, MO, and †University of Missouri, Columbia, MO.

The present study investigated the effects of a benzodiazepine (oxazepam) on anxiety as measured by autonomic and self-report indices in a nonclinical sample. These effects were assessed during a resting state and stressed state. Psychophysiological and self-report measures were recorded during an initial baseline (before drug administration), postdrug baseline (after drug administration when subjects were unaware of upcoming stressor), countdown, stressor, and poststressor baseline phases. Anxiolytic effects were found during stressed state as measured by skin conductance level. The benzodiazepine did not significantly affect heart rate or self-reported anxiety. Implications are discussed.

ANXIETY AND DRINKING BEHAVIOR: MODERATING EFFECTS OF ALCOHOL-RELATED EXPECTANCIES. Matt G. Kushner* and Kenneth J. Sher.† *University of Minnesota, Minneapolis, MN, and †University of Missouri, Columbia, MO.

We evaluated whether alcohol-related expectancies moderate the association between measures of anxiety and alcohol use/abuse. As predicted, student subjects with stronger expectations for alcohol-induced tension reduction showed the strongest positive correlations between measures of anxiety and drinking behavior; however, this finding held for male subjects only. These findings are consistent with past studies showing gender differences in the alcohol/anxiety connection, and also highlight the need to understand better the processes underlying the development of alcohol-related expectancies. We suggest that these issues potentially relate to the etiology of alcohol problems for some individuals.

IMPLICIT PRIMING OF AN ALCOHOL EXPECTANCY MEMORY NETWORK. Genevieve M. Chenier and Mark S. Goldman. University of South Florida, Tampa, FL.

This study was designed to test the theory that alcohol expectancies can be usefully conceived as a semantic network that contains information about alcohol effects and that can operate on an implicit level with no awareness or conscious deliberation. An implicit memory, word fragment completion task was primed using either an alcohol-related or neutral context. Word fragments were derived from alcohol expectancies and neutral words.

Subjects were not told that the context was pertinent to the purpose of the experiment. A repeated measures ANOVA indicated a significantly elevated number of expectancy fragments completed in the alcohol context, as compared to other conditions, $F(1, 113) = 12.46$, $p < .001$. The study offers support for operation of expectancies on an automatic memory level.

PRIMING THE PUMP: ALCOHOL EXPECTANCY ACTIVATION INCREASES DRINKING BEHAVIOR. Laurie Roehrich* and Mark S. Goldman.† *University of California, San Francisco, CA, and †University of South Florida, Tampa, FL.

If expectations about alcohol exist as stored memories, priming this construct may influence drinking patterns. Eighty undergraduate women ($n = 20$ per cell), participated in two, supposedly unrelated studies. A 2×2 factorial design simultaneously varied contextual primes (bar or control video) with construct primes (expectancy or neutral words). Beer consumption during a subsequent taste-rating task served as the primary dependent variable. Women exposed to unobtrusive alcohol cues drank significantly greater amounts ($p < .001$) of beer, compared with subjects who received control primes. The priming techniques appeared to have additive effects. Alcohol primes may also have differential effects for heavier versus lighter female social drinkers.

EFFECT OF MINIMUM DRINKING AGE ON ALCOHOL CONSUMPTION AND IMPAIRMENT. Vincent J. Adesso, Eric D. Devine, Constantine Ioannidis and Bertrand D. Berger. University of Wisconsin, Milwaukee, WI.

Research has found no change in consumption and mixed results in impairment with changes in the minimum legal drinking age. Alcohol consumption and impairment due to alcohol were compared in 1,016 male and female college students 6 months younger ($n = 516$) and 6 months older ($n = 500$) than the minimum legal drinking age (MLDA). The sample was selected from each of four semesters across a 2-year period while the new MLDA took full effect. Results of the $2 \times 2 \times 4$ (Gender \times Age \times Time) MANOVA revealed a Gender by Time interaction for impairment with both men and women increasing in impairment across time and a main effect for gender—men consumed more than women.

GENDER AND AGE EFFECTS ON ALCOHOL EXPECTANCIES. Leslie H. Lundahl, Tania M. Davis, Vincent J. Adesso, Bertrand D. Berger and Celeste O. Milligan. University of Wisconsin, Milwaukee, WI.

Little work has investigated the possible relations among gender, age, and alcohol expectancies. Using multivariate analysis of variance, the Alcohol Expectancy Questionnaire (AEQ) was used to examine the differences among these variables. Results revealed that expectancies of increased physical pleasure, more global positive changes, and tendencies toward stronger tension reduction differentiated the men from the women. In addition, older subjects generally were less likely to report global, positive expectancies, increased assertiveness,

and arousal with feelings of power than were younger subjects. Results failed to indicate a significant interaction between gender and age.

DEFEAT AND THREAT OF ATTACK OCCASION PENTYLENETETRAZOLE-APPROPRIATE RESPONDING. Jeffrey A. Vivian, E. M. Weerts and Klaus A. Miczek. Tufts University, Medford, MA.

Being threatened by an attacking conspecific prompts defensive and submissive reactions and may induce anxiety. Male Long-Evans rats were trained to discriminate 20 mg/kg pentylenetetrazol (PTZ) or 0.4 mg/kg midazolam (MDZ) from saline in a two-choice drug-discrimination task. After brief defeats (Ss serving as intruders), a) administration of saline engendered PTZ-appropriate responding in the intruder but b) did not alter MDZ-appropriate responding after flumazenil pretreatment. During four exposures to the threat of attack, saline generated 55, 42, 56, and 33% PTZ-appropriate responding. These results suggest that an anxiety-like state occurs during defensive behavior and does not appear to be associated with long-term changes at the benzodiazepine receptor.

REPEATED ADMINISTRATION OF PCP AND AMPHETAMINE: EFFECTS ON SOCIAL BEHAVIOR. Rhea E. Steinpreis, J. D. Sokolowski, A. Papanikolaou and J. D. Salamone. The University of Connecticut, Storrs, CT.

Both phencyclidine (PCP) and amphetamine produce psychotic reactions in humans that resemble different aspects of schizophrenia. Our laboratory has previously demonstrated that acute administration of these drugs also affects social behavior in rats. The present study was used to determine the effects of repeated administration of these drugs on rat social behavior. Rats were given repeated daily IP injections of either PCP (4.0 mg/kg), amphetamine (4.0 mg/kg) or 0.9% saline for 7 days. On days 1, 4, and 7 rats were placed in a stable home colony of three other rats and observed for social behavior in 30-min sessions. On the first day of administration both drugs reduced various social behaviors. With repeated injections, the effects of PCP showed tolerance and the effects of amphetamine demonstrated sensitization.

STRAIN-DEPENDENT EFFECTS IN BEHAVIORAL PHARMACOLOGY. John R. Glowa. National Institutes of Health, Bethesda, MD.

Several behavioral differences between the histocompatible LEW/N and F344/N rat are described, including data on the behavioral effects of corticotropin-releasing hormone (CRH) and the effects of alprazolam and buspirone on the startle response. Since these strains have known differences in their stress response, GABAergic and serotonergic function, alcohol intake, and preference for cocaine and morphine, these strain-dependent differences are discussed in light of possible relationships between the stress response and individual susceptibility to stimulant abuse. It appears that the degree of sensitivity of central components of the stress response may be better than the magnitude of the glucocorticoid response in predicting abuse potential.

OPIOID SENSITIVITY AS MEASURED BY OPERANT SCHEDULE-CONTROLLED RESPONDING. Gregory I. Elmer,* J. O. Pieper,* Steven R. Goldberg* and Frank R. George.† *National Institute on Drug Abuse Addiction Research Center, Baltimore, MD, and †University of New Mexico, Albuquerque, NM.

The purpose of this study was to determine the effect of an opioid on schedule-controlled behavior in genotypes that differ significantly in innate opiate receptor concentration. The effects of etonitazene on fixed ratio responding for water was examined in CXBK/ByJ, CXBH/ByJ, C57BL/6J, and DBA/2J mice; the ED₅₀ for etonitazenes rate-depressant effects were 38.4, 16.6, 13.1, and 8.8 µg/kg, respectively. There appears to be no relationship between sensitivity to the rate-depressant effects of etonitazene and previously reported differences of the analgesic, stimulant or reinforcing properties of etonitazene.

NICOTINE INCREASES PRE-PULSE INHIBITION OF ACOUSTIC STARTLE REFLEX IN RATS. Jane B. Acri,* David E. Morse† and Neil E. Grunberg.* *Uniformed Services University of the Health Sciences, Bethesda, MD, and †Food and Drug Administration, Rockville, MD.

Nicotine can increase acoustic startle amplitude and may enhance attention, but effects of nicotine on pre-pulse inhibition (PPI) of acoustic startle are not known. PPI reflects gating of sensory stimuli related to attention. In the present study, acutely administered nicotine had a biphasic dose effect on startle amplitude in rats, with increases at lower doses (0.01 mg/kg) and decreases at higher doses (0.5-5.0 mg/kg) compared to controls. Lower doses of nicotine also increased amount and percentage of PPI, whereas higher doses reduced amount but not percentage of PPI. Results are consistent with nicotine's reported effects on attention and biphasic dose effects.

CONSUMPTION OF CONCURRENTLY AVAILABLE MONEY AND CIGARETTES: RESPONSE REQUIREMENT EFFECTS. Richard J. DeGrandpre, Warren K. Bickel, Stephen T. Higgins and John R. Hughes. University of Vermont, Burlington, VT.

In behavioral economics, consumption of a reinforcer is determined by its price and by the price of other available reinforcers. This study examined the effects of price manipulations on the consumption of concurrently available money and cigarettes. During approximately fifteen 4-h sessions, money and cigarettes were concurrently available according to fixed-ratio (FR) schedules of reinforcement. After consumption stabilized under a FR 100 for both reinforcers, the response requirement for each reinforcer was varied separately (FR 100, 1,000, 2,500), while the other reinforcer was held constant at FR 100. Increasing the FR value decreased money and cigarette consumption to a similar degree and in a positively decelerating fashion. Also, as the price for one reinforcer increased, consumption of the other generally increased. These results indicate that the demand for these two commodities can be significantly altered by own-price and other-price manipulations.