Use Incentives to Stop Inmates' Substance Abuse

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CORONADO, CALIF. — Treatment and continuing care are two key components to a chronic care approach to effective recovery for patients with a substance abuse problem.

But in a correctional setting, that basic model faces several challenges and is sometimes impossible to employ, Dr. Jack Kuo said at the annual meeting of the American Academy of Addiction Psychiatry.

Frequent lockdowns, lack of communication between mental health and substance abuse staff, and access to drugs by inmates are just a few obstacles he faces as a staff psychiatrist for the California Department of Corrections and Rehabilitation.

"Many people think that prisoners with substance abuse problems are abstinent

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because they are in prison, he said. "Unfortunately, that's not always the case. They have access through various types of smugglings, sometimes through visitors to the inmates, sometimes through guards, s o m e t i m e s through med-

ical personnel. You do have a number of illegal drugs that make their way into the system."

Abuse of prescription medications is common, he said, and inmates "will manufacture complaints to get their hands on these products." A popular drug of abuse is Wellbutrin, "which a lot of them will crush and snort as a cheap stimulant. A lot of them will also use Seroquel or other types of sedating medications."

Interventions that have been demonstrated in research studies to be effective for drug-abusing offenders include residential substance abuse treatment, cognitive-behavioral therapy, contingency management, and medications. However, treatment must last an average of 90 days to produce stable behavior change, Dr. Kuo said. That's difficult to achieve in a state prison system like California's, where lockdowns because of infighting or gang violence shut down prison yards for months at a time and medications such as methadone, naltrexone, and buprenorphine cannot be used.

In addition, some correctional officers may frown upon efforts to rehabilitate inmates with a history of substance abuse. "They figure, 'hey these people did something that's illegal. That's why they're locked up. Why should they get treatment?' Overcoming those types of attitudes can be fundamental to providing treatment."

The California Department of Corrections and Rehabilitation provides services to 9,200 inmates in 22 prisons. All programs use the therapeutic community model and are operated by private companies. A report issued in February 2007

by the California Office of Inspector General found that despite an annual cost of \$36 million, the state's in-prison substance abuse treatment programs have little or no effect on recidivism.

To improve the current system, Dr. Kuo recommends integrated public health and safety strategies that involve research-based treatment under close supervision, the opportunity to avoid incarceration or a criminal record when possible, and consequences for noncompliance. Treatment

that emphasizes contingency management holds the most promise, he said.

"Treatment does not need to be voluntary to be effective," added Dr. Kuo, who is also a psychiatrist with Promises Treatment Centers in Malibu, Calif. "Strong motivation can facilitate the treatment process. Sanctions or incentives related to family, employment, or the criminal justice system can significantly increase treatment entry and retention rates and the success of drug treatment interventions. It is important to

use rewards and sanctions to encourage prosocial behavior and treatment progress."

Research has shown that using rewards to recognize progress is an effective way to change behavior. "Rewards can take many forms, including certificates of achievement or verbal praise from an authority figure such as a judge," he said. "Establishing an attitude of 'catching people doing things right' creates a positive environment for fostering and maintaining behavior change."



GEODON is indicated for the treatment of acute manic or mixed episodes associated with bipolar disorder and for the treatment of schizophrenia.

Elderly patients with dementia-related psychosis treated with atypical antipsychotic drugs are at an increased risk of death compared to placebo. GEODON is not approved for the treatment of patients with dementia-related psychosis.

GEODON is contraindicated in patients with a known history of QT prolongation, recent acute myocardial infarction, or uncompensated heart failure, and should not be used with other QT-prolonging drugs. GEODON has a greater capacity to prolong the QT $_{\!\!\!\! c}$ interval than several antipsychotics. In some drugs, QT prolongation has been associated with torsade de pointes, a potentially fatal arrhythmia. In many cases this would lead to the conclusion that other drugs should be tried first.

As with all antipsychotic medications, a rare and potentially fatal condition known as neuroleptic malignant syndrome (NMS) has been reported with GEODON. NMS can cause hyperpyrexia, muscle rigidity, diaphoresis, tachycardia, irregular pulse or blood pressure, cardiac dysrhythmia, and altered mental status. If signs and symptoms appear, immediate discontinuation, treatment, and monitoring are recommended.

Please see brief summary of prescribing information on adjacent page.

Prescribing should be consistent with the need to minimize tardive dyskinesia (TD), a potentially irreversible dose- and duration-dependent syndrome. If signs and symptoms appear, discontinuation should be considered since TD may remit partially or completely.

Hyperglycemia-related adverse events, sometimes serious, have been reported in patients treated with atypical antipsychotics. There have been few reports of hyperglycemia or diabetes in patients treated with GEODON, and it is not known if GEODON is associated with these events. Patients treated with an atypical antipsychotic should be monitored for symptoms of hyperglycemia.

Precautions include the risk of rash, orthostatic hypotension, and seizures

The most common adverse events associated with GEODON in bipolar mania were somnolence, extrapyramidal symptoms, dizziness, akathisia, and abnormal vision.

In short-term schizophrenia trials, the most commonly observed adverse events associated with GEODON at an incidence of ${\geq}5\%$ and at least twice the rate of placebo were somnolence and respiratory tract infection.

In short-term schizophrenia clinical trials, 10% of GEODON-treated patients experienced a weight gain of ≥7% of body weight vs 4% for placebo.