Brief Communication

Adulterants in crack cocaine in Brazil

Adulterantes no crack vendido na “Cracolândia”

Introduction: Brazil is the world’s biggest consumer of crack cocaine, and dependence is a major public health issue. This is the first study to investigate the prevalence of potentially harmful adulterants present in hair samples from Brazilian patients with crack cocaine dependence.

Method: We evaluated adulterants in hair samples extracted by convenience from 100 patients admitted at the 48 hour-observation unit of Centro de Referência de Álcool, Tabaco e Outras Drogas (CRATOD), Brazil’s largest center for addiction treatment. A cross-sectional analysis was performed with the data obtained.

Results: Adulterants were found in 97% of the analyzed hair samples. The most prevalent adulterant was lidocaine (92%), followed by phenacetin (69%) and levamisole (31%).

Conclusion: Adulterants were widely prevalent in hair samples from crack users treated at CRATOD: at least one adulterant was present in virtually all the hair samples collected. This points to a need to monitor adverse effects in the clinical setting in order to provide this high-risk group of patients with prompt and effective care related to the acute and chronic complications associated with these adulterants.

Keywords: Cocaine, crack cocaine, adulterants, substance use disorders, toxicology.
Introduction

Crack cocaine dependence is a significant public health issue in Brazil, which is the world’s largest consumer of crack cocaine; in 2013, the lifetime and one-year prevalence of crack cocaine dependence were estimated at 1.5 and 0.8%, respectively. Most of the cocaine and crack cocaine traded contains adulterants and diluents in variable proportions. Some adulterants have similar effects to cocaine or are used to enhance the effect of the drug and reduce the amount of cocaine used, thus reducing production costs and maximizing profit.

The most popular adulterants present in crack cocaine are caffeine, lidocaine, phenacetin, levamisole, benzocaine, procaine and hydroxyzine. Here we present an evaluation of adulterants present in hair samples from patients admitted to Centro de Referência de Álcool, Tabaco e Outras Drogas (CRATOD), affiliated with the State Department of Health of the State of São Paulo. CRATOD is the largest center for addiction treatment in Brazil. It is located in the surroundings of “Cracolândia,” an area in the city of São Paulo, Brazil, in which crack cocaine is openly traded and consumed. Although other studies have looked into adulterants present in samples of cocaine confiscated in Brazil, this is the first study to evaluate the absorption of such substances by Brazilian crack cocaine users.

Methods

We used a cross-sectional design in a convenience sample. Hair samples were collected from patients admitted to CRATOD, considering the following eligibility criteria: being 18 years old or older; having been brought to the service from “Cracolândia” between June and December 2014 (n=100); and having signed an informed consent form as determined by the institutional review board at CRATOD.

Hair samples were stored and analyzed at Laboratório Chromatox. Samples were collected from the posterior vertex region during the first 48 hours of observation. All samples were analyzed for cocaine, benzoylecgonine, cocaethylene, hydroxyzine, lidocaine, procaine, benzocaine, tetrahydrocannabinol (THC), phenacetin and levamisole, following the methods previously described by Tsanaclis & Wicks, using 10 to 20 mg of hair, alkaline hydrolysis, and solid phase extraction. The analyses were performed using liquid chromatography-mass spectrometry (Agilent, Santa Clara, U.S.).

Results

The sample comprised 100 patients admitted to CRATOD for the treatment of crack cocaine addiction. Adulterants were present in 97% of the hair samples. All the samples resulted positive for cocaine and benzoylecgonine, 94% for cocaethylene and 23% for THC. The most prevalent adulterant was lidocaine (92%), followed by phenacetin (69%) and levamisole (31%). Other adulterants found were benzocaine (19%), procaine (5%) and hydroxyzine (2%). The sample could be classified into 13 groups on the basis of adulterants present in the hair samples (Figure 1). The most common combination was phenacetin and lidocaine, followed by lidocaine only.

Table 1 - Sociodemographic profile of crack cocaine users (n=100) submitted to hair analysis at CRATOD, São Paulo, Brazil, from May to December 2018

<table>
<thead>
<tr>
<th>Variable</th>
<th>n (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean age (years), mean (SD)</td>
<td>29.3 (8.56)</td>
</tr>
<tr>
<td>Age range (years)</td>
<td></td>
</tr>
<tr>
<td>≤ 20</td>
<td>16 (16)</td>
</tr>
<tr>
<td>21-30</td>
<td>43 (43)</td>
</tr>
<tr>
<td>31-40</td>
<td>31 (31)</td>
</tr>
<tr>
<td>≥ 41</td>
<td>10 (10)</td>
</tr>
<tr>
<td>Gender</td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>49 (49)</td>
</tr>
<tr>
<td>Female</td>
<td>36 (36)</td>
</tr>
<tr>
<td>Transgender</td>
<td>15 (15)</td>
</tr>
<tr>
<td>Race</td>
<td></td>
</tr>
<tr>
<td>White</td>
<td>36 (36)</td>
</tr>
<tr>
<td>Afro-Brazilian</td>
<td>64 (64)</td>
</tr>
<tr>
<td>Civil status (n=96)</td>
<td></td>
</tr>
<tr>
<td>Single</td>
<td>53 (55.2)</td>
</tr>
<tr>
<td>Married</td>
<td>15 (15.6)</td>
</tr>
<tr>
<td>Divorced</td>
<td>26 (27.1)</td>
</tr>
<tr>
<td>Widow</td>
<td>2 (2.1)</td>
</tr>
<tr>
<td>Schooling (n=92)</td>
<td></td>
</tr>
<tr>
<td>&lt; 8 years</td>
<td>70 (76)</td>
</tr>
<tr>
<td>≥ 8 years</td>
<td>22 (24)</td>
</tr>
<tr>
<td>Homeless</td>
<td></td>
</tr>
<tr>
<td>Yes</td>
<td>73 (73)</td>
</tr>
<tr>
<td>No</td>
<td>27 (27)</td>
</tr>
<tr>
<td>Currently working (n=94)</td>
<td></td>
</tr>
<tr>
<td>Yes</td>
<td>34 (36.2)</td>
</tr>
<tr>
<td>No</td>
<td>60 (63.8)</td>
</tr>
</tbody>
</table>

Data presented as n (%), unless otherwise specified.
CRATOD = Centro de Referência de Álcool, Tabaco e Outras Drogas.
Discussion

The large body of research available on adulterants and diluents present in cocaine has provided robust evidence of their use and its clinical consequences. A study carried out in the Netherlands found that 40.6% of their cocaine samples contained adulterants, and the most common were benzocaine, caffeine, lidocaine, levamisole, diltiazem, procaine, atropine and phenacetin. In 9,681 samples collected in Spain between 1985 and 1987, lidocaine was the main adulterant, present in 69.6%. Conversely, lidocaine was absent from samples collected between 1992 and 1993, with caffeine being the most prevalent adulterant. Fucci & De Giovanni investigated the composition of 156 samples collected in Rome between 1996 and 1997 and found 14 different adulterants, including lidocaine (16.5% in 1996 and 12.3% in 1997), caffeine (11% in 1997 and 6.1% in 1997) and phenacetin (6.7% in 1997) in the illicit cocaine market.

Following these initial studies, a collaboration called the Trans-European Drug Information (TEDI) project was launched. This combined the consumer-targeted drug testing systems of the European Union with data from Spain, Switzerland, Austria, the Netherlands, Brussels, and Portugal and found differences in terms of the purity and composition of drugs. Several adulterants were found in illicit cocaine, with levamisole being the most commonly detected adulterant in 2013, followed by phenacetin and caffeine.

In the U.S., the Drug Enforcement Administration (DEA) reported that in 2004 2-20% of cocaine samples contained hydroxyzine and 8-20% contained diltiazem. The first report of levamisole as an adulterant in the U.S. was by the DEA in 2005, and it is now routinely found in cocaine samples. Since the early studies, the reported prevalence of adulteration of cocaine with levamisole, diltiazem or hydroxyzine has increased from 6.5% in 1999 to 53.6% in 2007.

Levamisole was originally developed in the 1960s as an antihelminthic agent; it has immuno-stimulant properties that increase endogenous opiate levels and alter monoaminergic function – these effects may be responsible for its popularity as a cocaine adulterant. In vivo research using the conditioned place preference paradigm to investigate the combination of cocaine and levamisole showed a marked synergistic effect. Levamisole is also very cheap, so using it as an adulterant substantially increases profit. It has also been postulated that levamisole acts as a monoamine oxidase inhibitor and increases dopamine transmission. Aminorex, a psychoactive metabolite of levamisole, has a potent amphetamine-like effect on dopamine and norepinephrine transporters. Because the half-life of aminorex far exceeds that of cocaine, combining levamisole with cocaine may prolong the duration of the stimulant effects.

Because of its immunomodulatory effects, levamisole has been used to treat immune-mediated and inflammatory disorders, such as rheumatoid arthritis, nephritic syndrome, inflammatory bowel disease, aphthous ulcers and colon cancer. As a result, one of the most severe consequences of the adulteration of cocaine with levamisole is leukoencephalopathy. Xu et al. speculated that levamisole could cause acute and recurrent white matter lesions. There are previous reports of cases related to the presence of levamisole.

In Brazil, studies performed in the state of Minas Gerais reported that 72% of samples were less than 20% pure, with caffeine, lidocaine, benzocaine, and prilocaine being the most common adulterants in cocaine. In the state of São Paulo, Carvalho & Mídio reported that lidocaine, procaine, and caffeine were the main adulterants in 389 samples of cocaine confiscated in 1997. Maldaner et al. reported that phenacetin was presented in 30% of 210 samples seized in various Brazilian states; they also found levamisole, caffeine and lidocaine.

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Leukoencephalopathy is similar to other spongiform encephalopathies in which fluid is trapped in small vacuoles within the myelin lamellae. Findings from magnetic resonance imaging (MRI) include hyper-intense T2 and FLAIR signals, abnormality in the diffusion-weighted signal, gadolinium enhancement and/or surrounding edema. White matter lesions are typically found in subcortical and periventricular white matter, but can involve the brainstem and cerebellum. A brain biopsy can demonstrate loss of myelin and accumulation of perivascular lymphocytes. Other clinical consequences, such as neutropenia, agranulocytosis, and dermatological disorders, have also been reported.

Since the first case of agranulocytosis was described by Buchanan in 2010, there have been multiple case reports of severe agranulocytosis after chronic abuse of cocaine and crack cocaine. Cutaneous and thrombotic vasculopathy associated with the use of levamisole-adulterated cocaine is more prevalent in female users and is characterized by pronounced retiform purpura with cutaneous necrosis, which can lead to serious deformations and may require amputation. It has previously been hypothesized that there is a synergistic relationship, or cross-sensitization, between cocaine and levamisole, and cocaine alone has been linked to necrotizing granulomatous vasculitis and Churg-Strauss vasculitis.

Another severe complication of chronic cocaine abuse, related to adulteration of cocaine with phenacetin and local anesthetics, is methemoglobinemia, which can lead to severe hypoxemia, seizures, coma, and death. This blood disorder is characterized by the presence of circulating methemoglobin in erythrocytes, resulting in reduced tissue oxygen due to a reduction of the oxygen-carrying ability of erythrocytes, and an increased affinity for oxygen in non-affected heme molecules, which impairs the off-loading of oxygen to the tissues.

At least one adulterant was present in 97% of the hair samples analyzed in the present study, most often local anesthetics, phenacetin and levamisole. The widespread presence of adulterants in hair samples may be due to the use of crack cocaine from multiple sources, although our sample usually consumed the drug in the vicinity of “Cracolândia.”

Our findings should be considered in light of the limitations of the study, such as its cross-sectional design, reliance on a single center and the small sample size. Further prospective studies of the consequences of the widespread presence of crack cocaine adulterants in hair samples should be performed, in order to provide a clearer picture of their public health impact.

This is the first study of the presence of potentially harmful adulterants in hair samples from Brazilian patients with crack cocaine dependence. On the basis of the prevalence of adulterants in the hair samples of users from “Cracolândia,” we conclude that there is a need to monitor adverse effects in the clinical setting in order to provide this high-risk group of patients with prompt and effective care.

 Disclosure
No conflicts of interest declared concerning the publication of this article.

 References
8. Tanner-Smith EE. Pharmacological content of tablets sold as "ecstasy": results from an online testing service. Drug Alcohol Depend. 2006;83:247-54.


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