High mortality among young crack cocaine users in Brazil – a 5-year follow-up study.

M Ribeiro-Araújo, J Dunn, R Laranjeira, R Sesso

People who misuse licit or illicit substances have higher mortality rates. Most research has been done with other drugs, but little is known about mortality in crack users. In the present study 131 consecutive crack dependent patients, admitted to the detoxification ward of a public hospital with a city-wide catchment area in São Paulo, Brazil between 1992 and 1994, were followed-up 5 years later.

Participants, methods and results
The sample were predominantly young (mean age = 23.6 years SD=6.7), single (67%), white men (75%) of low educational attainment (56%) and high unemployment (69%). All met DSM IV criteria for cocaine dependence.

Between 1998 and 1999, 124 (94.6%) of the original cohort were located (mean follow-up 44.3 months) and were assessed using a structured interview, approved by the local ethics committee. Death certificates were verified from records held at the Municipal Offices. Twenty-three patients (18.5%) had died: 3 were accidental (2 cocaine overdoses and 1 drowning), 7 due to infectious complications of intravenous drug use (6 of AIDS and 1 of hepatitis B) and 13 had been shot. Relatives reported that these latter deaths were related to turf fights, punishment levelled out by drug dealers for unpaid debts or police repression. Mean age at death was 27 years (range: 18 to 40).

Survival analysis showed that the probability of being alive 5 years post-treatment was 0.80 (95% CI = 0.77 to 0.84), see Figure 1. Observed mortality rate, calculated by the direct standardisation method, adjusted for age and sex, was 24.92 per 1000. The expected all cause mortality rate in São Paulo, adjusted for age and
sex, was 3.28 per 1000 inhabitants, giving an excess mortality rate of 21.64 per 1000 – a mortality ratio of 7.60.

Seventeen variables were tested as predictors of mortality using Cox’s proportional hazards regression. These included: demographic data, level of schooling, drug use variables, criminal history, treatment history and pre-admission psychosocial assessment. Three variables were identified as predictors of mortality: history of intravenous drug use (hazard risk = 3.28, 95% CI = 1.42 to 7.59, p = 0.005), unemployment at index admission (hazard risk = 3.48, 95% CI = 1.03 to 11.80, p = 0.045) and premature discharge from index admission (hazard risk = 2.21, 95% CI = 0.94 to 5.18, p = 0.068).

Comment
Follow-up studies of crack/cocaine users are rare and do not have mortality as their main focus\(^1\), making comparisons difficult. The nearest comparable group is opiate addicts for whom more data are available. These follow-up studies usually report death rates from 6 to 22 deaths per 1000 inhabitants, mainly due overdose\(^2\)–\(^4\). As the opiate use is rarely seen in Brazil, the mortality rate related to this substance is not available. In our study, however, homicide was the main cause of death. The age adjusted homicide ratio for our sample was 7.74, suggesting that crack use substantially increased the risk of a violent death. In the USA, which has similar gun laws to Brazil, it has been argued that violence surrounding emerging crack markets led to an increase in the homicide rate\(^5\).

A limitation of our study is that patients came from just one hospital, albeit one that admitted patients from all parts of São Paulo City and one of just 2 hospitals with specialist detoxification units at the time the study began. This limits the generalisability of the results.

This is the first study of mortality in crack users. Regression analysis has identified 3 risk factors, which could inform future treatment policies: (i) enable patients to
stay in treatment longer, (ii) more effective harm reduction interventions for injectors and (iii) social reintegration. However, tighter gun controls would probably have greater impact on mortality for both crack users and the general population.

References

UNIAD, Departamento de Psiquiatria, Escola Paulista de Medicina, UNIFESP, Rua Botucatu 740, São Paulo – SP, Brazil, 04023-900 (M Ribeiro MSc, R Laranjeira PhD)
Department of Psychiatry, Royal Free & University College Medical School, Rowland Hill St, London, NW3 2PF (J Dunn DM)
GRIDEC, Escola Paulista de Medicina (R Sesso DM)