INCIDENCE OF NON-FATAL AND FATAL OPIOID OVERDOSE FOLLOWING RELEASE FROM PRISON AMONG MEN WHO REGULARLY INJECTED DRUGS PRE-IMPRISONMENT: A PROSPECTIVE COHORT STUDY

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Background:

Opioid overdose is a leading cause of preventable morbidity and mortality among people released from prison. Previous Australian estimates of non-fatal and fatal opioid overdose post-prison release potentially underestimate overdose risk among people engaging in injecting drug use (IDU) preimprisonment, as these cohorts included people who may not have drug use histories (i.e., at low or no risk of overdose). Among a cohort of men who reported regular IDU pre-imprisonment, we determined non-fatal and fatal opioid overdose incidence post-release.

Methods:

Participant data were linked to ambulance and National Death Index data. We defined non-fatal opioid overdose as an ambulance contact including naloxone administration and subsequent improvement in Glasgow Coma Scale. Fatal opioid overdose was a poisoning death (using ICD-10 codes) with opioids recorded as a contributing cause. Person-years (PY) at-risk commenced from prison release (Sept2014–Dec2016) and ended at death or 31 December 2018. Separate crude incidence rates (IR) for multiple-event non-fatal opioid overdoses and fatal opioid overdoses were calculated overall, and at 30, 180, 365 days post-release, reported per 1000PY with 95% confidence intervals (95%CI).

Results:

Participants (N=400) contributed 1222.3PY. Participants had 70 non-fatal (range: 0–4, IR: 57.3/1000PY, 95%CI: 45.3–72.4) and 14 fatal opioid overdoses (IR: 11.5/1000PY, 95%CI: 6.8–19.3). Incidence was approximately seven (non-fatal opioid overdose IR: 396.8, 95%CI: 230.4–683.4) and five (fatal opioid overdose IR: 61.1, 95%CI: 15.3–244.1) times greater in the first 30 days post-release, tapering off and stabilising thereafter.

Conclusion:

Among men reporting regular IDU pre-imprisonment, we found high rates of non-fatal and fatal opioid overdose, particularly in the first 30 days. Expanded access to evidence-based interventions such as opioid agonist treatment and take-home-naloxone may reduce overdose risk. Ongoing surveillance of overdose incidence post-release is required to monitor overdose trends and measure the effectiveness of such interventions.

Disclosure of Interest Statement:

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