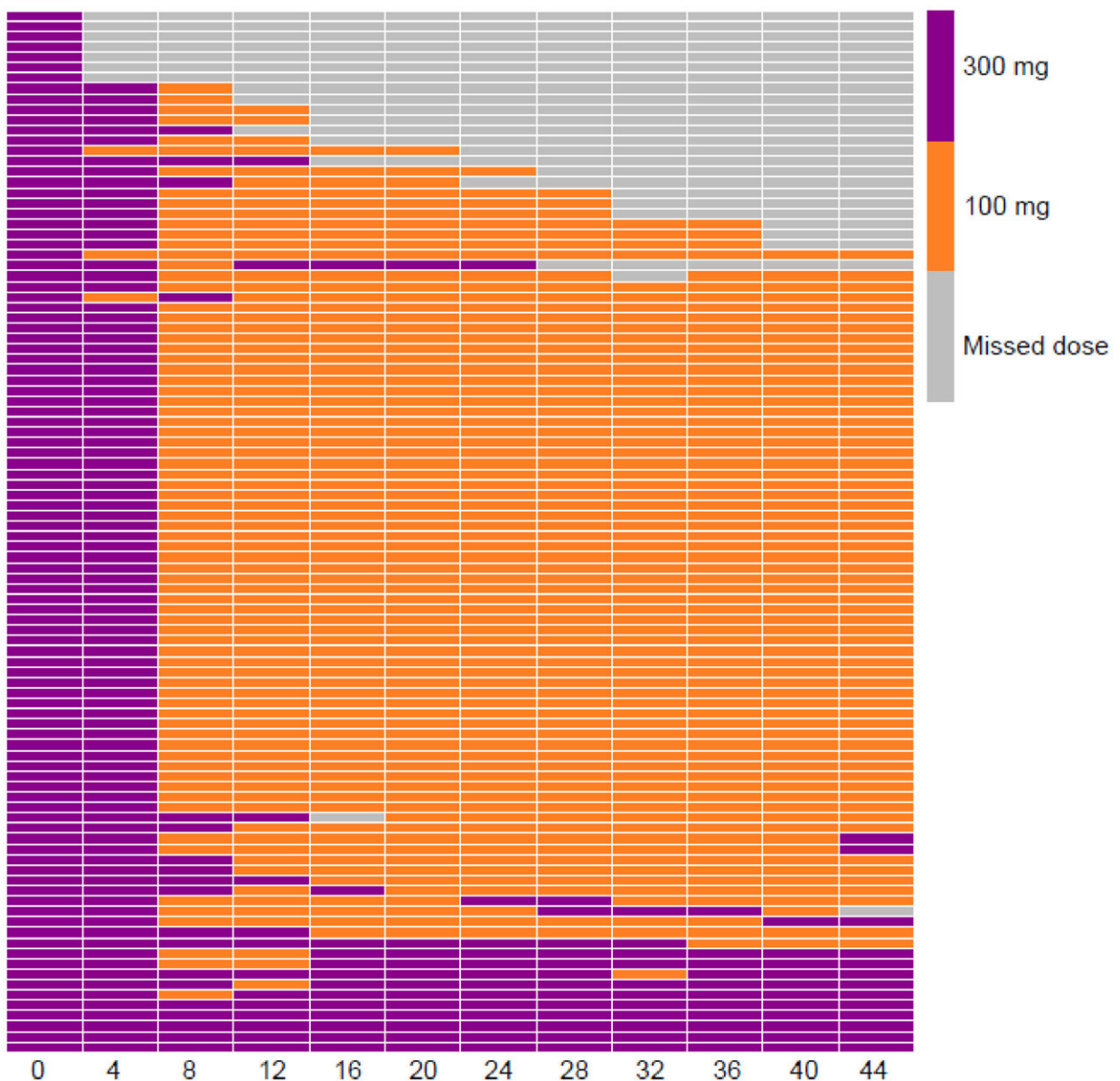


Results of clinical trial will impact treatment for opioid dependence in Australia and internationally

November 5 2021



BUP-XR utilization for each participant across the study period. Monthly adherence to treatment and dosing schedule at an individual level over the entire 48 weeks of treatment. Each row represents an individual participant and the column represents the treatment timepoint (in weeks). Gray boxes represent missed doses, orange (100 mg BUP-XR) and purple (300 mg BUP-XR). Failure to receive a BUP-XR for more than 56 days between injections would mean non-adherence to protocol defined dosing regimen and would result in automatic discontinuation from treatment. Credit: DOI: 10.1016/j.drugpo.2021.103492

The results of a clinical trial of extended-release buprenorphine injections (BUP-XR), an opioid agonist treatment (OAT), demonstrate high retention and treatment satisfaction.

Led by researchers at the National Drug and Alcohol Research Centre (NDARC), UNSW Sydney, the study had 100 participants with an average age of 44 years, mainly men (72 percent) and a long and a long OAT history and were recruited from across a network of specialist public drug [treatment](#) services, [primary care](#) and some private practices in three states, all receiving monthly subcutaneous BUP-XR injections for the treatment of opioid dependence.

The dosing standard schedule involved two doses of 300 mg BUP-XR at baseline and month one. Thereafter, doses were flexible with 100 mg or 300 mg every 28 days, based on clinical decisions between participants and treating doctors.

Treatment retention at 24 and 48 weeks was 86 percent and 75 percent, respectively.

Director of NDARC, Professor Michael Farrell said, "A major strength of this study is that study sites reflected 'real-world' community-based

services, primarily consisting of publicly funded drug treatment services."

"The high retention and treatment satisfaction rates in this study demonstrate that further expansion of BUP-XR treatment can be incorporated well within the existing [service](#) system."

The [study](#) found that during treatment there were declines in heroin use, non-prescribed opioid use, and injecting drug use. Improvements in quality of life, participation in employment, and treatment satisfaction measures were also observed.

"Although modest, these improvements are consistent with ongoing retention and stabilization within a treatment program among a cohort of people already very treatment-experienced," said Professor Farrell.

The other benefits of BUP-XR are outlined in the findings and include not having to travel daily for supervised (often daily) dosing, reducing experience of stigma, and reducing negative rituals and habits associated with dosing.

"Nonetheless it is crucial that OAT choice remains," said Professor Farrell.

"For some, particularly those who already have flexibility in their OAT, this formulation may be less attractive. It is important to ensure that patient choice around which medications people receive in OAT are retained, to ensure that the impact of choice on treatment outcomes is given full consideration in future planning and service change."

"These results may impact future Australian and international guidelines and [health policy](#), and give more treatment options to people with opioid dependence."

More information: Michael Farrell et al, Outcomes of a single-arm implementation trial of extended-release subcutaneous buprenorphine depot injections in people with opioid dependence, *International Journal of Drug Policy* (2021). [DOI: 10.1016/j.drugpo.2021.103492](https://doi.org/10.1016/j.drugpo.2021.103492)

Provided by International Journal of Drug Policy

Citation: Results of clinical trial will impact treatment for opioid dependence in Australia and internationally (2021, November 5) retrieved 26 June 2024 from <https://medicalxpress.com/news/2021-11-results-clinical-trial-impact-treatment.html>

This document is subject to copyright. Apart from any fair dealing for the purpose of private study or research, no part may be reproduced without the written permission. The content is provided for information purposes only.