





**IPEDs** 

### **WHY COLLECT DATA?**

To monitor service needs (e.g. equipment and staffing), identify gaps in service provision, support funding bids and to help provide continuity of care with clients

## WHEN AND HOW TO COLLECT DATA?

Date collection should only occur when the client is comfortable with it, to prevent possible barriers to engagement. Where possible, collect data electronically.

#### MINIMUM DATA TO COLLECT

At minimum aim to collect:

- Date of contact
- Initials
- Date of hirth
- Gender
- Drug(s) used
- Equipment and/or intervention provided

Initials, date of birth and gender can be combined to create a unique anonymised client identifier commonly used in healthcare data collection.



#### **ADDITIONAL DATA**

Demographic and training-related data can help determine client needs and gauge their understanding of drug use and training, which can help with client engagement.



- Occupation
- Physical training history
- Diet
- Patterns of IPED use (history, doses, cycles, drugs used)

#### **HEALTH-HISTORY**

Health-history data helps track client progress and initiate discussions around wellbeing.

Availability of specific resources is required for some data (i.e. bloodwork), but collecting such data can be a valuable tool to increase engagement.



- Psychoactive drug use
- Medication (prescribed)
- Bloodwork (e.g. liver function and hormone levels)
- Injection sites and technique
- BBV testing/sexual health
- Self-reported side effects of IPED use
- Mental health assessment and history











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