

# Is the DRID Key Indicator 'stuck' in the 1990s?



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# Context & issues

Not exhaustive!



Changes in types of psychoactive drugs injected and variation across Europe - Opiates / Poly-drug use / Stimulants.

Ageing cohorts of psychoactive drugs users – particularly PWID from ‘outbreaks’ in 1980s /1990s.

Increasing concern about bacterial infections among PWID, and other infections among PDUs – e.g. STIs and TB.

Increase in use, and injection, of image and performance enhancing drug in some countries.

Link between infections and the responses to them.

Economic crisis – there are the costs of collecting, and also reporting, data – but, more importantly, this could affect patterns of use and resources available for responses.



## a) Hepatitis notification data



Hepatitis 'notification' data also collected by ECDC.

There is a cost associated with is to duplicate reporting.

Q: Does this data need to be collected by two EU bodies?

A: No.

ECDC and EMCDDA need to **work together** to ensure there is a single effective EU system for collecting this data.

The system **must** capture exposure information (i.e. which infections acquired through injecting drug use), **and** be able to provide information on exposure categories by age/gender.

## b) Prevalence data



Currently a lot of 'breakdowns' can be reported through ST9 – the data collection sheet is 'long'.

Collection could [/should] be stream-lined for **each system/study** to **key data** for **highest** geographic area available:- i.e. overall prevalence, with breakdowns by age, gender, years since first injected, and imprisonment, plus NUTS3 breakdown for national or multi-regional studies (?separate optional sub-section/sheet for rest of break downs ?optional reporting for sub-areas **or** for agreed 'sub-groups' – e.g. current injectors only; recent initiates only).

Data comes from a very diverse range of sources.

Ensure collection of good quality data on study methodology **and** sample characteristics.

## c) Behavioural data



Behavioural data collection just revised.

Do these data link well enough with responses – particularly intervention uptake **and** coverage?

When next reviewed:-

- Maintain limit on number of items: ?max 12.
- Review data items collected.
- Collection of data from sources other than those providing prevalence data – ?a separate standard table with method sheet.
- Relationship with responses data collection.
- Would a structured questionnaire be better (for some items)?

# What is not collected / addressed



Should DRID collect data on:

- bacterial infections related to injection [YES]?
- infections in non-injectors (e.g. due to smoking or snorting causing tissue / lung damage) [May be]?
- sexual transmitted infections [May be], TB [YES], etc.?
- also changing patterns of drug use – i.e. Risks [YES]?

Also need to look gathering information on related responses.

Sensitive to ‘evolving’ infection risks – soft data.

A structured questionnaire to collect numerical and descriptive data on evolving risk, infections & on the responses to them?

Finally, current DRID data collection is focused on psychoactive drug users where do image and performance enhancing drug users fit?

Structured questionnaire, or initially a ‘special issue’?

# So is the DRID Key Indicator ‘stuck’ in the 1990s?



In 1990s (and 1980s) focus of work around drug related infectious diseases was driven by the concerns about HIV and hepatitis C among injectors of psychoactive drugs.

These are both still important issues.

During the 2000's, at least in some countries, there has been increasing concern and focus on:

- bacterial infections,
- infections among PDUs more widely,
- impact of changes in the types of psychoactive drugs injected/used on risk, and
- about use and injection of image and performance enhancing drugs.

**May be!**

# Afterward



It is important to remember it is the

European

# Monitoring

Centre for Drugs and Drug Addiction.

**Thank You.**