



Comparing European countries with 'high' or 'increasing' drug-related death (DRD) rates

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"Scotland May Now Have Highest Rate of Drug-Related Deaths in the EU"

Drug-related Deaths (per 1 million pop.)



Figures rounded to closest whole number. Data based on latest available year.

200

"Comparing data on drug-related deaths is <u>difficult</u> because there are differences in definitions, toxicology and coroner processes, under-reporting and delays in reporting."



Figure 27: Rate of drug related deaths per million population, by country and European average (which includes England)

Source: Public Health England (2017)

"Although national differences in coding and reporting practices, as well as possible under-reporting, <u>make it difficult to compare countries</u>, analysing trends over time <u>within individual countries</u> is valuable" (EMCDDA, European Drug Report, 2015).



NB: these are general population DRD rates and do not account for variation in the size (and rate) of the populations at risk of DRD.

Aim

- To explore why DRD rates in European countries are *high* or *increasing*.
 - Sweden
 - Norway
 - Scotland
 - Finland
 - Denmark
 - Estonia
 - Ireland

NB: selected countries pre-determined by EMCDDA

Methods

- Country profiles, developed with national experts, and relevant, available EMCDDA indicators
- All seven countries: opioids implicated in 80%-90% of DRD; therefore, opioid-related DRD were the primary focus
- Considered:
 - Differences (and trends) in the number of drug users at risk of DRD
 - Differences (and trends) in factors that may influence the risk of DRD (among those at such risk)
 - Differences in / changes to mechanisms to record DRD

Drivers of the extent of DRD:

- The size of the population specifically at risk of DRD
 - Available (albeit flawed) estimates indicate a 30x difference in prevalence rates across EU countries for the main 'at risk' group (EMCDDA, 2016): so, we should expect to observe differences in DRD general population rates!
- The level of risk experienced by the 'at risk' group
 - Are users in one place or time more or less likely to suffer a fatal overdose? A much more interesting question.... <u>and much more difficult to answer</u>

Comparison of DRD rates for two hypothetical countries:

Country A:

100 opioid DRD p.a. General Population=1,000,000 persons Estimated POU prevalence 5,000 persons (POU population rate=50 per 10,000)

Country B:

500 opioid DRD p.a. General Population=5,000,000 persons Estimated POU prevalence 60,000 persons (POU population rate=120 per 10,000)

General population DRD rate=1 per 10,000 POU DRD rate=200 per 10,000

- = ≠
- General population DRD rate=1 per 10,000 POU DRD rate=83 per 10,000





Mean annual number of opioid related deaths (2009-13) vs. 'best (gu)estimates' of problem opioid prevalence (or proxy):



Mean annual number of opioid related deaths (most recent 5-year period) vs. 'best (gu)estimates' of problem opioid prevalence (or proxy):



Cohort studies:

- 23 drug user mortality studies identified for the 7 countries
- 16 excluded did not report a DRD rate
- Additional 3 excluded lack of case definition comprising active drug use during observation
- 4 remaining studies, 2 countries, based on 2 cohorts
- Scotland (opiate users, observation 1996-2006): DRD rate during & post-treatment 4.4 (95% CI: 4.1–4.6) per 1,000 PY (Merrall et al., 2012)
- Norway (opiate users, observation 1997-2003): DRD rate duringtreatment 4 (95% C.I. 0-8), post-treatment 21 (17–25), circa 6.7 (derived) per 1,000 PY combined (Clausen et al., 2008): note wide C.I.

Trends in the number of Drug-Related Deaths involving opioids: 2004-2015:



Trends in the number of Drug-Related Deaths involving opioids: 2004-2015:



Available HRDU/PDU prevalence trend estimates

Country 🔶	Method 🔶	2014 🝦	2013 🔶	2012 🔶	2011 🝦	2010 🔶	2009 🔶	2008 🝦	2007
AT		ţ			1				
BE			3						3
BG	1		:				1		
СҮ	TP	2.7	2	1.77	2.09	1.54	2.5	2	3.66
cz	ТМ	6.73	6.28	5.71	5.51	5.3	5.04	4.39	4.2
DE	TM	3	4.42	4.4	4.65	4.06	3.71	3.96	3.3
DK	2	2	1	3	1	5	3	e -	4
EE	i.	1	4	:	ī.	ī.	Ŧ	÷	1
ES	:	3	:	3	(T)	1	:	:	1
FI	1	1	4	<u>a</u>	-10	2	1	1	4
FR	1	2	:	:	1	1	1	1	1
GR	1			:			1	:	
HR	MM	3	:	3.48	3.3	:	:	:	3
HU	1	- 2	4	3	10	1	1	1	d
IE	2	2	1	4	1	2	:	:	1
Π	TM						9.95	9.8	
LT		1			1				
LU	ОТ	2	4	3		20	6.16	1	7.7
LV	TM		:		9.37	13.33	1	:	1
МТ	ž.			3			1	:	
NL	5	3	:	3	1	2	:	:	1
NO	1	2	d		1	2	1	1	d
PL	:		:	4			:	:	1
РТ									
RO	:	:	1	3	12	2	:	:	1
SE	1	1	4		-1×	2	1	1	4
SI	1	2	:	:	1	1	1	1	1
SK	от		:	:		1	÷,	2.68	4.7
TR	2	3	:	3	12	2	:	:	4
UK	СМ	4	3	9.16	9.19	9.38	9.79	1	10.1

http://www.emcdda.europa.eu/data/stats2016

Drivers of risk:

- Demographic
- Behavioural
- Contextual/environmental setting
- Set of (non-exhaustive) hypotheses about potential drivers focus on drivers where (trend) data *may* be available
- We are looking at a moving target
- Upward and downward drivers will co-occur (and may operate simultaneously with changing prevalence)
- Likely complex set of interactions between some drivers
- No simple answers



Demographic risk: trend in mean age at DRD (all DRD):



Variation in behavioural risk?

- Do the at risk populations vary wrt injecting, type of opioid use, poly drug use, etc?
- Injecting: there is substantial variation in rate of injecting; Scotland is somewhere in the middle/lower end of the distribution.
- Type of opioid: there is variation; fentanyl in Estonia likely to put users at higher risk; buprenorphine (with alcohol & benzos) in Finland; Scotland (& Ireland) unusual re dominance of heroin.
- Polydrug use: toxicology suggests that polydrug use is common, perhaps more common in Scotland (but perhaps superior screening?)

Polydrug use:

Figure A.6.1: Crude numbers of annual drug-induced deaths recorded in Scotland, 2000-2015 (source: National Records of Scotland, 2016)



Prevalence of BBV among PWID (2010-16):

Country	HCV	HBV	HIV	
Denmark	75%	35%	<5%	
Estonia	76-90%	3-22%	~50%	
Finland	74%	1.2%	-	
Ireland	68%	-	-	
Norway	64%	(Oslo) 35%	2.4%	
Scotland	58%	-	1.9%	
Sweden	60-80%	-	-	

OST coverage:

- Variations between countries
- Absence of trend data on the size of the atrisk population, it is not possible to assess the potential effect that changes in the size of the OST group exert on DRD trends
- Lack of information on the delivery of treatment, dimensions of which are likely to modify a treatment's protective effect with regard to DRD

Availability of Harm Reduction Interventions:

Country	Methadone Maintenance Treatment	Buprenorphine Treatment	Buprenorphine /Naloxone	Needle & Syringe exchange	Supervised Injecting Facilities	Heroin Assisted Treatment	Take-Home Naloxone
Denmark	1	1	×	✓	1	1	1
Estonia	\checkmark	✓	×	✓	×	×	1
Finland	\checkmark	1	✓	1	×	×	×
Ireland	\checkmark	×	✓	✓	×	×	1
Norway	\checkmark	1	1	1	1	×	1
Scotland	\checkmark	1	1	1	×	×	1
Sweden	\checkmark	1	✓	1	×	×	×

Summary:

- Scotland's DRD rate (per person at risk) is broadly equivalent to (perhaps less than?) those of the other countries considered
- There is little clear evidence of elevated behavioural, demographic, or environmental (Tx, HR, BBV) risk in Scotland, vs. other countries
- Demographic risk (age) has increased
- Scotland has a reasonably comprehensive set of interventions, incl OST, to reduce risk – at least maintain them & improve them – <u>clear gap is</u> <u>SIF/HAT</u>

Thank you

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