

Drug misuse prevention

Review protocol for evidence review 2

	Details	Additional comments
Review question 2a	How acceptable are drug misuse prevention interventions that people currently receive?	None.
Review question 2b	What drug misuse prevention interventions and support do people feel might be more effective?	None.
Language	English only	None.
Study design	All types of qualitative primary studies including surveys, interviews, case studies, or ethnographic or action research. Relevant data may also be extracted from quantitative studies including studies which met the inclusion criteria for evidence review 1.	None.
Setting	<p>Social environments where drugs may be available such as nightclubs, pubs, festivals and music venues.</p> <p>Fitness environments such as gyms and sporting events.</p> <p>Environments where drugs may be used in a sexual context (for example, 'chemsex' parties).</p> <p>Online and 'virtual' environments, including social media.</p> <p>Youth clubs and youth organisations.</p> <p>Schools, colleges and universities.</p> <p>Health, social care and other environments where interventions may be delivered, for example, primary health care services, sexual health services and custody suites.</p> <p>Interventions in prisons and young offender institutions</p>	Countries: as described, the reviews will only include studies from countries in the European Union, Switzerland, Iceland, Norway, USA, Canada, New Zealand and Australia. All other countries are considered to have significant cultural differences to the UK and are therefore excluded.

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	<p>will be excluded.</p> <p>Universal school based interventions (i.e. those not targeted at any of the population groups described) will be excluded.</p> <p>Interventions set in the workplace will be excluded (other NICE guidance covers workplace interventions).</p> <p>Included countries are: Australia, Austria, Belgium, Bulgaria, Canada, Croatia, Cyprus, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Israel, Italy, Latvia, Lithuania, Luxembourg, Malta, Netherlands, New Zealand, Norway, Poland, Portugal, Romania, Slovakia, Slovenia, Spain, Sweden, Switzerland, Turkey, UK, USA.</p>	
Population	<p>Following discussion with Liverpool HE team on 25/06/15, it has been agreed that only at risk population groups will be considered for the purposes of targeting. Only the 10 following population groups will be included:</p> <ul style="list-style-type: none"> people who have mental health problems people involved in commercial sex work or are being sexually exploited people who are lesbian, gay, bisexual or transgender people not in employment, education or training (including children and young people who are excluded from school or are regular truants) children and young people whose parents use drugs looked after children and young people children and young people who are in contact with young offender team but not in secure environments (prisons 	<p>It is not possible to look at all populations who misuse drugs. The list identified represents the children, young people and adults who are most likely to start misusing drugs or who are already experimenting or who misuse drugs occasionally or who are risk of moving onto other drugs.</p> <p>The NICE team also considered if any particular ethnic groups should be included in the target population group list above. Based on 2013/14 statistics in Section 5 of the Crime Survey for England and Wales it was decided on 30/06/15 that ethnicity would not be covered in the population target group list.</p>

	Details	Additional comments
	and young offender institutions) people who are considered homeless people who attend nightclubs and festivals people who are known to use drugs occasionally / recreationally	
Intervention	Interventions will be included that have a stated and measured aim of enhancing personal and social skills, improving their self-confidence, increasing knowledge and awareness about the risks of drug use and/or increasing knowledge and awareness about how to reduce the risks and harms of drug use: <hr/> group-based skills training or information provision using lessons, talks and activities <hr/> one-to-one skills training, information provision and advice given as part of planned outreach activities <hr/> one-to-one skills training, advice and information provided using peer education initiatives <hr/> opportunistic skills training, advice and information provision <hr/> using targeted print and new media for different groups at risk of drug misuse to influence social norms or enhance skills and provide information and advice <hr/> family-based programmes providing structured support for children and young people at risk of drug misuse group-based behaviour therapy for children and young people who are at risk of drug misuse	<hr/> For example, targeted refusal skills training in schools and colleges. <hr/> For example, for young people at festivals. <hr/> For example, with gay men in nightclubs. <hr/> For example, provided by youth workers. <hr/> For example, magazines, websites, social media, text messages. <hr/> For example, motivational interviewing for parents or carers and parental skills training. <hr/> Focusing on coping mechanisms, problem-solving and goal setting.

	Details	Additional comments
	parental skills training for parents or carers of children who are at risk of drug misuse	Focusing on stress management, communication skills, helping children develop problem-solving skills and setting behavioural targets.
Outcomes	<p>Acceptability of interventions including barriers and facilitators.</p> <p>Details of others interventions / support that study participants feel might be more effective in preventing drug misuse (with rationales).</p>	None.
<p>Searches</p> <p>[See Appendix 2A for the search for each step]</p>	<p>Searches will follow a iterative, step-wise approach. Individual search steps are outlined below. The same set of results will be sifted for reviews 1 and 2. Note that steps 1-6 represent the original search outline. Subsequent steps (7-11) were added to the process after the amount of time available for sifting was revised. Where a search step was dependent on prior steps the decisions made as to which direction to adopt are either noted below or in the relevant section of appendix 2A.</p>	<p>An initial search using a 'classical' search approach indicated that in excess of 36,000 papers would be retrieved in Medline alone, even with a relatively small set of search terms. The team decided it would not be appropriate to use this type of approach for this guideline. The iterative, step-wise approach outlined below aims to balance precision and sensitivity without producing an unmanageable volume of results. This is in accordance with the guidance set out in the NICE guidelines development manual, section 5.1. In particular it should be noted that preventative programmes (as described in the NICE scope) are not always flagged as such in the relevant literature.</p>
	<p>Step 1: Search for systematic reviews only from 1995 to 2015 using the following databases:</p> <p>Cochrane Database of Systematic Reviews (CDSR)</p> <p>Database of Abstracts of Reviews of Effectiveness (DARE)</p> <p>HealthEvidence.org</p> <p>Campbell library</p>	<p>Some sources will be browsed rather than searched where this approach is considered more time-efficient than searching. Reviewer A and B will sift all the search results (100% double sifting) for step 1.</p>

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	<p>DoPHER NIHR systematic reviews programme website</p> <p>Guidelines: Guidelines.gov will be searched and any guidelines identified during the scoping exercise for this work will also be factored in. Any systematic reviews identified in any relevant guidelines will be ordered.</p>	
	<p>Step 2: All relevant systematic reviews identified in step 1 will be entered into Web of Science and a backward citation search will be performed to produce a database of included studies.</p> <p>Timeline: search results available 7th July to review team.</p>	<p>Note that only those references which can be automatically downloaded from the Web of Science Core Collection (including Medline) will be downloaded. This is therefore not “pure” citation searching, though it is expected to be far more efficient in terms of time. We expect subsequent protocol steps will help to address some of the deficit in retrieval compared to pure citation searching.</p>
	<p>Step 3: Given the time lag between the publication of a primary study and its potential inclusion in a published systematic review, the following sources will be searched from 2010 in order to identify recent primary evidence.</p> <p>Cochrane Central Register of Controlled Trials (CENTRAL) HTA database Embase MEDLINE/MEDLINE in Process Social Policy and Practice Social Care Online</p>	<p>The search strategy at this stage is likely to be broad but not comprehensive. In particular, we may choose to run less comprehensive searches on Embase and PsychInfo as the incremental yields from these databases are likely to be low once other sources are factored in.</p>

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	PsychInfo TrOPHI	
	Step 4: A more focused search of the databases in step 3, ranging back to 1995. This step is carried out as a back-stop in the event that we identify gaps in coverage for the evidence retrieved steps 1-3.	Following sifting of evidence from previous search steps the review team decided to focus on identifying additional evidence relating to evidence review 2 (acceptability of interventions) at this stage. Material retrieved was still sifted for potential inclusion in either review 1 or review 2.
	Step 5: Web searching In addition, the following websites will be searched: NIHR Public Health Research Programme Advisory Council on the Misuse of Drugs European Monitoring Centre for Drugs and Drug Addiction UN Office on Drugs & Crime Organization of American States Relevant material identified by the review team via other routes (for example during the scoping exercise for this review) will also be included at this stage.	None.
	Step 6: Named programme search. A list of specific programmes (for example the “good life approach”) will be compiled by the review team during steps 1-5. An additional search of Medline will be carried out using the names of these programmes as keywords in order to identify any additional, named articles.	None.
	Steps 7-10: Additional citation searching in Web of	None.

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	<p>Science, based on included references or selected, topic-relevant but non-includable material identified during previous search steps.</p>	
	<p>Step 11: Any additional references included in NICE guideline PH4 or the subsequent Evidence Update document, which also meet the date limits for the present guideline, will also be submitted for sifting.</p>	None.
Data screening	<p>All references from the database searches in each step will be downloaded, deduplicated and screened on title and abstract against the criteria above. Where no abstract is available, a web search will be used to locate one; if none is found, references will be screened on title alone.</p> <p>For all search steps described previously, except step 1, a randomly selected sample of 10% of records at title and abstract level will be screened by two reviewers (A and B) independently. The rate of agreement will be recorded. Disagreement will be resolved through discussion and with the arbiter as necessary. All records will be screened at title and abstract level by reviewer A.</p> <p>Where abstracts meet all the inclusion criteria, or if it is unclear from the study abstract whether it does, the full text will be retrieved and re-screened. Full-text screening will be carried out by two reviewers (A and B) independently and any differences resolved by discussion and with the arbiter if necessary. Inter-rater reliability will be recorded.</p>	None.

	Details	Additional comments
	Studies that are excluded at the full paper stage will be recorded along with the reason for their exclusion.	
Data extraction and QA	Quality assessment and data extraction for all included studies will be conducted using the tools in Developing NICE guidelines: the manual . All studies will be quality assessed and data extracted by one reviewer, with all data checked in detail by a second reviewer. Details of all extracted data will be entered into comprehensive evidence tables.	None.
Data synthesis	Data will be synthesised narratively in the first instance. If sufficiently homogeneous and high-quality data are located, meta-analysis may be considered, although this is unlikely.	None.
Subgroup analysis	Where possible, the effectiveness of interventions for subgroups will be disaggregated and reported, along with any differential effect on different subgroups.	None.
Other information/criteria	The review will report on any unintended consequences or adverse outcomes.	None.