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Mapping the evidence on identity processes and identity-related interventions in the smoking and physical activity domains: a scoping review protocol

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ABSTRACT

Introduction Smoking and insufficient physical activity (PA), independently but especially in conjunction, often lead to disease (and premature) death. For this reason, there is need for effective smoking cessation and PA-increasing interventions. Identity-related interventions which aim to influence how people view themselves offer promising prospects, but an overview of the existing evidence is needed first. This is the protocol for a scoping review aiming to aggregate the evidence on identity processes and identity-related interventions in the smoking and physical activity domains.

Methods The scoping review will be guided by an adaption by Levac et al of the 2005 Arksey and O’Malley methodological framework, the 2020 Preferred Reporting Items for Systematic Reviews and Meta-Analyses: Extension for Scoping Review (PRISMA-ScR) and the 2017 Joanna Briggs Institute guidelines. It will include scientific publications discussing identity (processes) and/or identity-related interventions in the context of smoking cessation and/or physical (in)activity, in individuals aged 12 and over. A systematic search will be carried out in multiple databases (eg, PubMed, Web of Science). Records will be independently screened against pre pilot inclusion/exclusion criteria by two reviewers, using the Active Learning for Systematic Reviews machine learning artificial intelligence and Rayyan QCRI, a screening assistant. A pre pilot charting table will be used to extract data from included full-text articles. Findings will be reported according to the PRISMA-ScR guidelines and include study quality assessment.

Ethics and dissemination Ethical approval is not required for scoping reviews. Findings will aid the development of future identity-related interventions targeting smoking and physical inactivity.

INTRODUCTION

Smoking and insufficient physical activity (PA), individually but especially in conjunction, are key preventable factors of disease (eg, cardiovascular diseases and cancer)1-5 and (premature) death.6-5 It is unlikely that new risk factors will be identified which have a similar or larger impact on health,6 especially when co-occurring.7-9 Despite this, world-wide, one-fifth of the population still smokes regularly10 and one-third to half fails to engage in regular PA.10 The most effective actions to reduce health risks are to quit smoking and increase levels of PA.11-12 While over the past decennia, an impressive number of smoking cessation and/or PA-enhancement interventions have been developed and tested, reviews show that they typically only have small to medium (short-term) effects.13-14 With only mildly successful interventions but impactful health risks, there is need for new and effective strategies to achieve successful smoking cessation and increased PA.

Identity theories and identity-related interventions (ie, which aim to influence how people view themselves) offer promising prospects in facilitating smoking cessation and increased PA. Identity typically starts forming...
This may also be why, to date, only two identity-related interventions are likely to remain difficult to develop effective identity-based interventions for specific target groups. The review will therefore take into consideration demographic characteristics (ie, SEP, health literacy, age, sex), smoking-specific characteristics (ie, age at onset of smoking, heaviness of smoking) and PA-specific characteristics (ie, levels of physical (in)activity) when mapping the available evidence.

In short, the current article describes the protocol for a scoping review aiming to map the available scientific evidence regarding identity processes and identity-related interventions and possible personal characteristics in the contexts of smoking and PA. A scoping review was chosen over other types of syntheses because it allows to uncover and analyse (different types of) evidence about the role of identity in smoking (cessation) and physical (in)activity. Additionally, investigating variations in the role of identity on the two health behaviours depending on personal characteristics will prove useful in developing identity-related interventions for specific target groups.

In sum, smoking and PA-identities and behaviours have been found to vary based on demographic characteristics. Plausibly, this makes personal characteristics highly relevant to consider in a review aiming to synthesise the role of identity in smoking (cessation) and physical (in)activity. Additionally, investigating variations in the role of identity on the two health behaviours depending on personal characteristics will prove useful in developing identity-related interventions for specific target groups.

In a nutshell, the research field is in need of a comprehensive overview encapsulating the role of identity in the contexts of smoking (cessation) and (insufficient) PA to guide the development of identity-based interventions targeting both health risk factors at once. The review to which this protocol belongs aims to provide such an overview.

Personal characteristics have been found to be associated with differences in smoking and PA, and identities. For example, smoking and insufficient PA have been found to be especially prevalent and co-occurring among socioeconomically disadvantaged individuals, men and those with lower levels of health literacy. Also, PA and the likelihood of quitting smoking tend to decline with age. Furthermore, findings show that individuals who started smoking young—around the age of 14–16, who are more nicotine dependent and who are heavy smokers (ie, who smoke 10+ cigarettes per day) are less likely to quit smoking. With regard to personal characteristics and smoking-related and PA-related identities, empirical studies have found that individuals with lower socioeconomic position (SEP), individuals who are more nicotine dependent and older individuals generally identify more with smoking than with quitting.

In short, the current article describes the protocol for a scoping review aiming to map the available scientific evidence regarding identity processes and identity-related interventions and possible personal characteristics in the contexts of smoking and PA. A scoping review was chosen over other types of syntheses because it allows to uncover and analyse (different types of) evidence about the topic and to inform areas for practice and future research. Findings are expected to aid the development of future identity-related interventions aiming to facilitate smoking cessation and increased PA. One such intervention is Perfect Fit (see the Funding statement), a virtual coaching intervention being developed by the authors of this review as part of a large Dutch consortium, which will employ identity-related interventions to motivate people to quit smoking and increase their PA levels.

Next to being the first to provide a comprehensive overview of the evidence on the role of identity, in the domains of smoking and PA, and to directly inform the development of interventions such as Perfect Fit, this scoping review will also be innovative methodologically. It will be the first in the field to make use of Active Learning for Systematic Reviews (ASReview), a machine learning
technology to select relevant literature (see stage 3 of Methods and Analysis).

METHODS AND ANALYSIS
We conducted Preliminary searches of the Open Science Framework (OSF), Cochrane Database of Systematic Reviews, Joanna Briggs Institute (JBI) Evidence Synthesis and PROSPERO in December 2020, at the inception of the study, in February 2021, before preregistering the protocol on OSF (https://osf.io/hkd9c/), and, as additional verification, in September 2021, after finalising the protocol manuscript. No current or underway systematic reviews or scoping reviews on the topic were identified. The current study is expected to run from February 2021 to December 2022.

The present scoping review protocol follows the adapted methodological framework for scoping reviews of Levac et al,57 originally developed by Arksey & O’Malley.58 This protocol as well as the final scoping review also conform to the guidelines published by the JBI59 and the recent Preferred Reporting Items for Systematic Reviews and Meta-Analyses extension for Scoping Review (PRISMA-ScR) guidelines.60 These guidelines were developed as a result of increasing popularity of scoping reviews and as a means to improving their quality.

Stage 1: identifying the research questions
Guided by the PRISMA-ScR Population, Concept, Context (PCC) principles, the research team agreed on articulating the following research questions:

RQ1—What is known about identity (processes) in adolescents and adults, in the contexts of smoking and PA, taking into consideration certain demographic characteristics (ie, SEP, health literacy, age, sex), smoking-specific characteristics (ie, age at onset smoking, heaviness of smoking, number of smoking years) and PA-specific characteristics (ie, levels of physical (in)activity)?

RQ2—What identity-related interventions are being used to influence smoking and PA in adolescents and adults, taking into consideration possible differences in implementation based on demographic characteristics (ie, SEP, health literacy, age, sex), smoking-specific characteristics (ie, age at onset smoking, heaviness of smoking, number of smoking years) and PA-specific characteristics (ie, levels of physical (in)activity)?

Stage 2: identifying relevant studies, eligibility criteria, information sources and search

Eligibility criteria
Inclusion of literature will happen according to the PCC eligibility criteria mentioned hereafter. Chosen criteria for the review are presented here in reverse order, that is, CCP, for readability purposes.

Context
In terms of context, we will include published, peer-reviewed scientific research papers and conference abstracts, written in English, Dutch or French.

Concept
When it comes to concept criteria, we will include literature which describes identity processes relating to smoking (cessation) and/or physical (in)activity (RQ1) and/or describes identity-related interventions implemented in the context of smoking (cessation) and/or physical (in)activity (RQ2). Preferably, the literature also considers demographic characteristics (age, SEP, health literacy, sex), smoking-related characteristics (heaviness of smoking, age at smoking onset) and PA-related characteristics (levels of physical (in)activity), although not considering these factors will not automatically lead to exclusion.

Population
With regard to population criteria, literature will be included when it studies individuals aged 12+ (on average), who smoke or have smoked tobacco (any type) or electronic cigarettes, and/or do or have engage(d) in less than their age-specific recommended levels of PA.47 In line with the explorative nature of a scoping review, no restrictions are formulated in terms of study design or publication year. Literature will be excluded when written in another language than English, Dutch or French, and/or smoking (cessation) or physical (in)activity and related interventions are discussed without relation to identity, and/or the target population is limited to individuals younger than 12 years old (on average).

Identifying relevant studies, information sources and search

In line with JBI guidelines,59 a three-step search strategy will be designed and used with the assistance of an experienced academic librarian from the Leiden University Medical Centre. Step 1 will consist of an initial limited search of PubMed, PsycINFO and Web of Science, using pre-defined keywords extracted from a dozen known key articles, to identify additional relevant keywords and index terms. Step two will include a second search across all relevant databases, that is, PubMed, PsycINFO, Embase, Emscare, Web of Science Core Collection, Wiley Cochrane Library, Psychology, Behavioural Sciences Collection and Academic Search Premier, OpenGrey.eu and British Library EthOS, using all identified keywords and index terms. In step 3, the reference list of the key articles will be handsearched for additional sources (ie, backward reference searching61) and missing search/index terms. A new search will be carried out using updated search and index terms and adapted to all databases (see online supplemental materials 1 for the final search string used for PubMed, which yielded the most records). Search strings for other databases are available on request). Once full-text screening is completed, backward reference searching of included articles61 will be used to identify new and/or missing records. Additionally, once data charting is complete, and before finalising the synthesis, a new search using the search strategy established in step two will be carried out to identify relevant newly published records.
Stage 3: screening and selecting studies
Title and Abstract screening
Following the search, and after removal of duplicates, titles and abstracts of identified records will be collated and uploaded into (1) ASReview, a free and open-access machine learning technology, and (2) Rayyan QCRI, a free title and abstract screening assistant. Following a pilot test of the screening manual, 10% of titles and abstracts will be independently assessed against eligibility criteria by two reviewers. One experienced reviewer (MHHMV) will screen titles and abstracts using ASReview. A second reviewer (KMP) will screen a random portion of titles and abstracts using Rayyan QCRI.

ASReview was chosen as primary screening tool for several reasons, the first being time-efficiency. A study by Ferdinands et al. evaluated the technology using six systematic review datasets from different research fields, and showed that only 8.3%–36.1% of titles and abstracts needed screening to identify the relevant ones. With each inclusion/exclusion decision, the ASReview algorithm learns what the reviewer finds relevant, and it subsequently sorts and presents the most relevant records first. As a result, considerably fewer screening hours are needed to arrive at the final selection. A second reason for choosing ASReview is its ‘human-in-the-loop’ machine learning technique. Through this technique, the reviewer maintains control over the entire screening process by having the final say in the relevance of every record. The reviewer is not dependent on a technology to include the relevant citations, but is aided by it. Finally, ASReview was chosen because it presents only titles and abstracts, and no authors or journal names. This allows the reviewer to judge each citation for its content rather than irrelevant metadata, and thereby removes a potential layer of “authority bias” in the choice of relevance. Despite substantial advantages, ASReview remains a new, undertested technology. This is why, to mitigate possible early technological kinks, we decided to have the second reviewer (KMP) manually perform independent double screening of a portion (ie, 10%) of the titles and abstracts in Rayyan QCRI, a widely used and appreciated screening assistant. Rayyan QCRI facilitates the screening process by highlighting pre-defined keywords in the title and abstract, and makes easy to assign and track reasons for inclusion/exclusion. Additionally, Rayyan QCRI permits to screen from anywhere thanks to its offline functionality, and allows reviewers to collaborate on a project while being blind to others’ screening decisions.

Regardless of the software used, screened titles and abstracts will be marked ‘included (for full-text screening)’ or ‘excluded’ based on assessment against the eligibility criteria. Screening in ASReview will stop after 150 consecutive records marked ‘excluded’. ASReview software developers advised to stop after 100–120 consecutive irrelevant records. However, with very little prior research to guide screen-stop decision in scoping reviews, and none using ASReview, we choose to apply a more conservative heuristic of 150 consecutive irrelevant records.

As previously mentioned, the second reviewer (KMP) will additionally randomly double screen a small portion of the total number of records using Rayyan QCRI. The review team agreed on a double screening amount of 10% of retrieved records. This amount was chosen because screening more would undermine the purpose, time-saving benefits and intelligence of ASReview, as also confirmed by its developers when consulted on the matter. Additionally, the review team expects the search string to yield a substantial amount of records. Therefore, after double screening 10%, we expect the second reviewer (KMP), to have a sufficient feel of the literature to help validate the screening decisions made in ASReview by the first reviewer (MHHMV).

After double screening is complete, a Cohen’s κ inter-rater agreement rate (IRA) will be calculated for the titles and abstracts screened by both reviewers. If IRA reaches at least 80%—level from which IRA is considered strong—double screening will stop. In case IRA is below 80%, a new random 10% of titles and abstracts will be double screened, and so on until 80% IRA is reached or all records have been double screened.

Full-text screening
Full texts of articles included based on title and abstract screening will be managed in Microsoft Excel, and marked ‘included’ or ‘excluded’ based on assessment against eligibility criteria. Reason(s) for exclusion will be recorded and reported in the inclusion flowchart of the scoping review. Where a full-text is unavailable, but the article is assessed as relevant in the title and abstract screening phase, the authors of the paper will be contacted to request a copy of the manuscript. In case key unpublished or missing data remains unobtainable after contact with the authors, or in case the record proves irrelevant to the research after all, the record will be excluded. Full-text screening decisions will be thoroughly documented and reported in the final scoping review. A complete overview of screening decisions will be available on request.

In accordance with PRISMA-ScR, a flow diagram of the search and the study inclusion process will be presented in the final scoping review. Disagreement about screening decisions will be resolved through discussion among the two reviewers, and if necessary coauthors, until consensus is reached.

Stage 4: data charting process and data items
Data will be charted from eligible full-texts by two independent reviewers (KMP and MHHMV) using a data charting tool developed by the entire review team. The data charted will include information about the source (eg, author, year of publication, country of origin) of the record, its methodology, its aims, and findings relevant to the review questions (main outcomes). Where available, information about demographic, smoking-specific characteristics and PA-specific characteristics will also
be charted. Where required, authors of papers will be contacted to request missing or additional data. A draft charting table will be pilot tested on usability prior to starting full-text screening, and updated as necessary. As recommended by Levac et al. in their methodological advice for scoping reviews, the charting table will be a living document, modified and revised as necessary during the process of charting data from each included evidence source. Modifications will be summarised in the scoping review.

In-depth assessment of how research pertaining to our review questions is conducted is, as of yet, lacking. However, such assessment could help synthesise and make sense of the findings in the final scoping review. Consequently, and although not a requirement in scoping reviews, we aim to critically appraise the quality of each included evidence source, using the JBI Critical Appraisal Tools. These tools allow to assess the quality of numerous types of evidence sources, from randomised controlled trials to qualitative studies. Two reviewers (KMP and MHMV) will independently assess the quality of each included full-text record, using the tool appropriate to the study design. Disagreement will be resolved through discussion among assessors, and if necessary coauthors, until consensus is reached.

**Stage 5: collating, summarising and reporting the results**

Results will be summarised into a narrative descriptive synthesis, and may include visual overviews (eg, graph, diagram or table) of the findings (conform the PRISMA-ScR guidelines). Depending on the available data, results will be presented following the PCC principles, that is per health behaviour (smoking, PA), separately for adolescents and adults, and per moderator (heaviness of smoking). Smoking and PA, two key risk factors of numerous diseases, will be charted. Where required, authors of papers will be contacted to request missing or additional data. A draft charting table will be pilot tested on usability prior to starting full-text screening, and updated as necessary. As recommended by Arksey and O’Malley’s framework, we aim to critically appraise the quality of each included evidence source, using the JBI Critical Appraisal Tools. These tools allow to assess the quality of numerous types of evidence sources, from randomised controlled trials to qualitative studies. Two reviewers (KMP and MHMV) will independently assess the quality of each included full-text record, using the tool appropriate to the study design. Disagreement will be resolved through discussion among assessors, and if necessary coauthors, until consensus is reached.

**Stage 6: consultation**

Consistent with Arksey and O’Malley’s framework, we will convene a team of stakeholders to assess the validity of the findings. Stakeholders will consist of clinicians and coaches who counsel individuals with regard to their smoking (cessation) and physical (in)activity levels. They will be asked to reflect on and discuss together the results of the scoping review to facilitate reporting of the findings and to inform future works in the field, including the further development of the Perfect Fit virtual coach. A detailed design of the consultation process will be created after stage 5 of the methodological framework (see above) has been completed.

**Patient and public involvement**

Patients or the public will not be directly involved in the conception, design and planning of this review, but will be involved, with explicit written consent, in the consultation phase (stage 6) of the scoping review.

**Ethics and dissemination**

This study did not require ethical approval as it did not involve human participation. Generally speaking, scoping reviews do not require ethical approval, as they analyse published literature. The final scoping review will be the first in the field to aggregate evidence regarding identity processes and interventions in the contexts of both smoking and PA, two key risk factors of numerous diseases, using a systematic approach. Findings are expected to aid the development of future identity-related interventions targeting smoking and PA, including the Perfect Fit virtual coaching intervention. Review findings will be presented to other key stakeholders, at scientific conferences and published in an open-access peer-reviewed journal.

**Amendment protocol**

In case of amendments, a separate version of the protocol will be maintained showing all modifications with tracked changes. Modifications will be reported and substantiated in the final scoping review.

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**Contributors**

All authors actively contributed to the design of this protocol. KMP, EM and WAG initiated the project. The protocol was drafted by KMP (guarantor) and refined by MHMV, EM and WAG. All authors approved the final protocol.

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**Competing interests**

All authors have completed the ICMJE uniform disclosure form at www.icmje.org/coiDisclosure.pdf and declare no support from any organisation for the submitted work, no financial relationships with any organisations that might have an interest in the submitted work in the previous 3 years, no other relationships or activities that could appear to have influenced the submitted work.

**Patient and public involvement**

Patients and/or the public were not involved in the design, or conduct, or reporting, or dissemination plans of this research.

**Patient consent for publication**

Not applicable.

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**Supplemental material**

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