

Additional file 2: Risk of bias schema

Major risk of bias domains*	Risk	Criteria	Hints/ notes
<p>1. Recruitment procedure & follow-up (in cohort studies):</p> <p>For cohort studies</p> <p><i>HINT: We are looking for selection bias:</i></p> <ul style="list-style-type: none"> - Was the cohort representative of a defined population? # - Was everybody included who should have been included? # - If response rate is slightly <50% but does not indicate selection bias, it will be listed as a demerit in extraction table. <p><i>PRELIMINARY RULING:</i></p> <ul style="list-style-type: none"> - If the cohort recruitment is based on a convenient/ self-reported sampling OR if response is <10% or not reported, the study will be excluded from analysis. 	low	<ul style="list-style-type: none"> <input type="checkbox"/> Cohort recruitment was acceptable.# <input type="checkbox"/> Baseline response is acceptable (50% or more) OR is <50% and >30%, but substantial differential selection could be excluded. <input type="checkbox"/> Loss to follow-up is below 20% in total and not different between the two groups (up to 10% difference).* 	
	high	<ul style="list-style-type: none"> <input type="checkbox"/> Cohort recruitment was not acceptable.# <input type="checkbox"/> Response not reported/ not calculable. <input type="checkbox"/> Total loss to follow-up is larger than acceptable (20% or more)* OR drop out differs between the groups by more than 10%* OR the reasons for drop out considerably differ between exposed and non-exposed groups.* 	
<p>For case-control studies</p> <p><i>HINT: We are looking for selection bias:</i></p> <ul style="list-style-type: none"> - Were the cases and control subjects representative of the same defined population ("study base"; geographically and/or temporally)? # - Was there an established reliable system for selecting all the cases? # - The same exclusion criteria are used for both cases and controls. # - Comparison is made between participants and non-participants to establish their similarities or differences. # - If response rate is slightly <50% but does not indicate selection bias, it will be listed as a demerit in extraction table. <p><i>PRELIMINARY RULING:</i></p> <ul style="list-style-type: none"> - If the recruitment of the study population is based on a convenient/ self-reported sampling OR if response is <10% or not reported, the study will be excluded from analysis. 	low	<ul style="list-style-type: none"> <input type="checkbox"/> Case selection and recruitment was acceptable.# <input type="checkbox"/> Control subjects' selection and recruitment was acceptable.# <input type="checkbox"/> Non-response was less than 50% for cases and/or control subjects OR it was >50% and <70%, but substantial differential selection of cases and control subjects could be excluded* 	
	high	<ul style="list-style-type: none"> <input type="checkbox"/> Case selection and recruitment was not acceptable.# <input type="checkbox"/> Control subjects' selection and recruitment was not acceptable.# <input type="checkbox"/> Non-response was >70% for cases or control subjects OR it was >50% and <70%, but substantial differential selection of cases and control subjects could not be excluded.* <input type="checkbox"/> Response not reported/ not calculable 	

*according to Ijaz et al. (2013), with modifications; # SIGN/CASP § Shamliyan

Major risk of bias domains*	Risk	Criteria	Hints/ notes
For cross-sectional studies <i>HINT: We are looking for selection bias:</i> <ul style="list-style-type: none"> - Was the study population representative of a defined population? # - Was everybody included who should have been included? # - If response rate is slightly <50% but does not indicate selection bias, it will be listed as a demerit in extraction table. <i>PRELIMINARY RULING:</i> <ul style="list-style-type: none"> - If the recruitment of the study population is based on a convenient/ self-reported sampling OR if response is <10% or not reported, the study will be excluded from analysis. 	low	<input type="checkbox"/> Recruitment of the study population was acceptable.# <input type="checkbox"/> Non-response was less than 50% OR it was >50% and <70%, but substantial differential selection of the study population could be excluded.*	
	high	<input type="checkbox"/> Recruitment of the study population was not acceptable.# <input type="checkbox"/> Non-response was >70% OR it was >50% and <70%, but substantial differential selection of the study population could not be excluded.* <input type="checkbox"/> Response not reported/ not calculable.	
2. Exposure definition and measurement	low	<input type="checkbox"/> Exposure was defined adequately covering more than one aspect of exposure (duration, frequency, intensity) and was assessed objectively: direct measurement or systematic observations or using a questionnaire that is validated.*	
	high	<input type="checkbox"/> Exposure was not defined adequately covering only one aspect of exposure (duration, frequency, intensity) and/or was assessed subjectively (self-report, questionnaire, interview) or using a proxy used to allocate exposure status (job matrix, job title).* <input type="checkbox"/> Different methods were used to measure exposure in different groups/ cases and control subjects (<i>in case-control studies</i>).§	
	unclear	<input type="checkbox"/> Not reported.	
3. Outcome “rate of/ risk to develop meniscal lesions”. Source and validation	low	<input type="checkbox"/> Outcome was accurately/ objectively measured to minimize bias (e.g. arthroscopically, MRI, open surgery)# <input type="checkbox"/> Measurement methods were similar in the different groups.#	
	high	<input type="checkbox"/> Outcome was not accurately or subjectively measured (self-reported, clinical examination).# <input type="checkbox"/> Measurement methods were different in the groups.#	
	unclear	<input type="checkbox"/> Not reported.	
4. Confounding and effect modification	low	<input type="checkbox"/> If risk estimators were calculated, major confounding factors (age, sex) were considered. <input type="checkbox"/> If only prevalence or incidence was assessed, at least sex and age are described.	
	high	<input type="checkbox"/> Major confounding factors (age, sex) were not considered.	
	unclear	<input type="checkbox"/> Not reported.	

Major risk of bias domains*	Risk	Criteria	Hints/ notes
5. Analysis method: methods to reduce research specific bias	low	<input type="checkbox"/> Authors used adequate statistical models to reduce bias (e.g. standardization, matching, adjustment in multivariate model, stratification, propensity scoring). [§]	
	high	<input type="checkbox"/> Authors did not use adequate statistical models to reduce bias.	
	unclear	<input type="checkbox"/> Not reported.	
6. Chronology	low	<input type="checkbox"/> Incident diseases were included. [#] <input type="checkbox"/> Temporal relation may be established (exposure precedes the outcome). [#] <input type="checkbox"/> No meniscal damage known at baseline (<i>in cohort and case-control-studies</i>).	
	high	<input type="checkbox"/> People with prevalent meniscal damage were included OR people with prevalent meniscal damage of baseline were not excluded (<i>in cohort studies</i>). [#] <input type="checkbox"/> Temporal relation cannot be established. <input type="checkbox"/> Meniscal status is unknown at baseline.	
	unclear	<input type="checkbox"/> Not reported.	

Minor risk of bias domains*	Risk	Criteria	Hints/ notes
7. Blinding of assessors	low	<input type="checkbox"/> Assessors were reported or indicated to be blind for individual exposure-status in cohort and cross-sectional studies and to case status in case-control and cross-sectional studies	
	high	<input type="checkbox"/> Assessors were reported or indicated <u>not</u> to be blind for individual exposure-status in cohort and cross-sectional studies and to case status in case-control and cross-sectional studies	
	unclear	<input type="checkbox"/> Not reported.	
8. Funding	low	<input type="checkbox"/> Grant/ non-profit-organizations* <input type="checkbox"/> Study was clearly not affected by sponsors.*	
	high	<input type="checkbox"/> Sponsoring organization participated in data analysis. <input type="checkbox"/> Study was probably affected by sponsors.	
	unclear	<input type="checkbox"/> Industry, combined industry+grant*, unclear if study was affected by sponsors. <input type="checkbox"/> Not reported.	
9. Conflict of interest	low	<input type="checkbox"/> Reported not having conflict of interest or clear from report/ communication that study was not affected by author(s) affiliation.*	
	high	<input type="checkbox"/> Conflict of interest exists (at least one author).*	
	unclear	<input type="checkbox"/> Not reported.	

Overall risk of bias assessment		Low Risk	High Risk	Unclear Risk
Major domains	1. Recruitment procedure & follow-up (in cohort studies)			
	2. Exposure definition and measurement			
	3. Outcome “rate of/ risk to develop meniscal lesions”. Source and validation			
	4. Confounding and effect modification			
	5. Analysis method: methods to reduce research specific bias			
	6. Chronology			
Minor domains	7. Blinding of assessors			
	8. Funding			
	9. Conflict of interest			
General rule for rating: Low risk of bias: low risk in all major domains High risk of bias: if not low risk		Overall assessment:		