Return to sports- a risky business? A systematic review with meta-analysis of risk factors for graft rupture following ACL reconstruction, Sports Medicine, Anna Cronström; anna.cronstrom@umu.se, Eva Tengman, Charlotte Häger, Umeå University

Online resource 2.

Table The Quality In Prognosis studies (QUIPS) checklist for risk of bias

Domains	Prompting items for Consideration	Ratings
Study Participation	<ul> <li>a. Adequate participation in the study by eligible persons</li> <li>b. Description of the source population or population of interest</li> </ul>	High bias: The relationship between the prognostic factor (PF) and outcome is very likely to be different for participants and eligible nonparticipants
	<ul> <li>c. Description of the baseline study sample</li> <li>d. Adequate description of the sampling frame and recruitment</li> </ul>	<b>Moderate bias</b> : The relationship between the PF and outcome may be different for participants and eligible nonparticipants
	<ul> <li>e. Adequate description of the period and place of recruitment</li> <li>f. Adequate description of inclusion and exclusion criteria</li> </ul>	Low bias: The relationship between the PF and outcome is unlikely to be different for participants and eligible nonparticipants
Study Attrition	<ul> <li>a. Adequate response rate for study participants</li> <li>b. Description of attempts to collect information on participants who dropped</li> </ul>	<b>High bias</b> : The relationship between the PF and outcome is very likely to be different for completing and non-completing participants
	c. Reasons for loss to follow-up are provided d. Adequate description of participants lost to follow-up	<b>Moderate bias</b> : The relationship between the PF and outcome may be different for completing and non-completing participants
	e. There are no important differences between participants who completed the study and those who did not	Low bias: The relationship between the PF and outcome is unlikely to be different for completing and non-completing participants
Prognostic Factor Measurement	<ul> <li>a. A clear definition or description of the PF is provided</li> <li>b. Method of PF measurement is adequately valid and reliable</li> </ul>	<b>High bias</b> : The measurement of the PF is very likely to be different for different levels of the outcome of interest
	<ul> <li>c. Continuous variables are reported or appropriate cut points are used</li> <li>d. The method and setting of measurement of PF is the same for all study participants</li> </ul>	<b>Moderate bias</b> : The measurement of the PF may be different for different levels of the outcome of interest
	e. Adequate proportion of the study sample has complete data for the PF  f. Appropriate methods of imputation are used.	Low bias: The measurement of the PF is unlikely to be different for different levels of the outcome of interest

	for missing PF data	
Outcome Measurement	a. A clear definition of the outcome is provided     b. Method of outcome measurement used is adequately valid and reliable	<b>High bias</b> : The measurement of the outcome is very likely to be different related to the baseline level of the PF
	c. The method and setting of outcome measurement is the same for all study participants	Moderate bias: The measurement of the outcome may be different related to the baseline level of the PF
		Low bias: The measurement of the outcome is unlikely to be different related to the baseline level of the PF
Study Confounding	a. All important confounders are measured	High bias: The observed effect of the PF
	<ul> <li>b. Clear definitions of the important confounders measured are provided</li> </ul>	on the outcome is very likely to be distorted by another factor related to PF and outcome
	<ul> <li>c. Measurement of all important confounders is adequately valid and reliable</li> </ul>	Moderate bias: The observed effect of the PF on
	<ul> <li>d. The method and setting of confounding measurement are the same for all study participants</li> </ul>	outcome may be distorted by another factor related to PF and outcome
	e. Appropriate methods are used if imputation is used for missing confounder data	<b>Low bias:</b> The observed effect of the PF on outcome is unlikely to be distorted by another
	f. Important potential confounders are accounted for in the study design	factor related to PF and outcome
	g. Important potential confounders are accounted for in the analysis	
Statistical Analysis and Reporting	Sufficient presentation of data to assess the adequacy of the analytic strategy	High bias: The reported results are very likely to be spurious or biased
	<ul> <li>Strategy for model building is appropriate and is based on a conceptual framework or model</li> </ul>	related to analysis or reporting
		Moderate bias: The reported results may be
	<ul> <li>c. The selected statistical model is adequate for the design of the study</li> </ul>	spurious or biased related to analysis or reporting
	d. There is no selective reporting of results	
		<b>Low bias</b> : The reported results are unlikely to be spurious or biased related to analysis or

repor	porting
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**Source:** Hayden JA, van der Windt DA, Cartwright JL, Côté P, Bombardier C. Assessing bias in studies of prognostic factors. Ann Intern Med. 2013;158(4):280-6