### ROBINS-I tool (Stage I): At protocol stage

Specify the review question	Bør patienter anbefales bevægerestriktioner efter total hoftealloplastik operation?
Participants	THA post op
Experimental intervention	Hip precautions/restrictions
Comparator	No restrictions
Outcomes	Hofteluksation, tidlig (Dislocations, early)
List the confounding domains relevant to all or most studies	Ledhovedstørrelse i studier med kohorte i tidsperioder før/efter praksisændring fra <32mm ledhoveder til >32mm

ledhoveder.

List co-interventions that could be different between intervention Operation procedure (e.g. posterior, lateral, anterior) groups and that could impact on outcomes

### ROBINS-I tool (Stage II): For each study

	Allen2018	Lightfoot2020	Mikkelsen2014	vanderWeegen2019
Design	Retrospective cohort, before/after	Prospective cohort, before/aft	er non-randomized controlled	Prospective cohort + control group of consecutive pts
Participants				
	THA post op	THA post op	THA post op	THA post op
Experimental intervention	Hip precautions/restrictions	Hip precautions/restrictions	Hip precautions/restrictions	Hip precautions/restrictions
Comparator	No restrictions	No restrictions	No restrictions	No restrictions
Confounders	Ledhovedstørrelse	Ledhovedstørrelse	No confounders	Ledhovedstørrelse
Co-interventions				
Is your aim for this study?				
Specify which outcome is being assessed for risk of bias				
Specify the numerical result being assessed				

ROBINS-I tool (Stage II): For each study

Specify target trial							Preliminary considerations of confounders			Additional confounding domains				Preliminary considerations of confounders			Additional co-interventions		
									OPTIONAL:				OPTIONAL:						
								is there	is failure to adjust				is failure to adjust			is the presence of this			is the presence of this
								evidence is the	for this variable				for this variable			co-intervention likely to			co-intervention likely to
								for this confounding				is the	(alone) expected to			favour outcomes in the			favour outcomes in the
								variable domain measur	e favour the			confounding domail			this co-			evidence that	
									experimental		evidence for this	measure validly and	d experimental		Intervention	intervention or		controlling for this	
					Specify which outcome is being assessed for		Confounding	Measured unnecess reliably by this	intervention or the	Confounding		reliably by this	intervention or the		was	thecomparator		co-intervention	thecomparator
Reference	Design	Participants Experimental Intervention	Comparator	is your aim for this study_?	risk of blas	Specify the numerical result being assessed	domain	variables any? variable?	comparator?	domain	variables unnecessary?	variable?	comparator?	Co-interventions	unnecessary?		Co-interventions	was unnecessary?	
Allen2018	Retrospective cohort, before/after	673/2551 Hip precautions/restrictions	Minimal postoperative restrictions	To evaluate the risk of early dislocations after removal of early hip resictions after THA	Hofteluksation, tidlig (Dislocations, early)														
Lightfoot2020	Prospective cohort, before/after	119/118 Hip precautions/restrictions	Minimal postoperative restrictions	To evaluate the risk of early dislocations after removal of early hip resictions after THA	Hofteluksation, tidlig (Dislocations, early)														
Mikkelser2014	non-randomized controlled	219/145 Hip precautions/restrictions	Minimal postoperative restrictions	To evaluate the risk of early dislocations after removal of early hip resictions after THA	Hofteluksation, tidlig (Dislocations, early)														
vanderWeegen2019	Prospective cohort + control group of consecutive pts	1049/1102 Hip precautions/restrictions	Minimal postoperative restrictions	To evaluate the risk of early dislocations after removal of early hip resictions after THA	Hofteluksation, tidlig (Dislocations, early)														

1. Bias due to confounding

Study	Outcome	1.1	L 1	.2	1.3	1.4	1	.5	1.6	1.7	7 1	.8	Risk of bias judgement	Comment and supporting quote	additional comment	Comment to ROBINS-I
Allen2018	All outcomes	N		-	-	-		-	-	-		-	Low	Majority of patients operated with >32mm (92.28 % og 97.33 %.)	3 Hip luxations reported for patients operated with posterior approach (desired)	
Lightfoot2020	All outcomes	PN	۱	-	-	_		-	-	-		-	Moderate	No information about head size	Hip luxations reported for patients operated with posterior approach (desired)	
Mikkelsen2014	All outcomes	N		-	-	-		-	-	-		-	Low	No confounding suspected. Majority of patients operated with >32mm (95.9%)	Posterior surgical approach	
vanderWeegen2019	All outcomes	N		-	-	-		-	-	-		-	Low	Hip luxations reported for patients operated with >32mm head size.	Posterior surgical approach	

#### 2. Bias in selection of participants into the study

Study	Outcome	2.	1 2.2	2.3	3 2.4	2.5	Risk of bias judgement	Comment and supporting quote	additional comment	Comment to ROBINS-I
Allen2018	All outcomes	Y	ΡΥ	Y	Y	N	Serious	2.1. Excluding due to post intervention exclusion of participants due to no precautions in 2009-2012 and due to a specific surgeon performing the procedure (argument of reducing bias).	Participants selected retrospectively from registry from two time periods with/without restrictions	2.1 excluding on post intervention factors such as revision surgey, conversion surgey and resurfacing operations. 2.2. Precations might be associated with re-operation.
Lightfoot2020	All outcomes	PN	-	-	Y	-	Low	No reasons to suspect selection bias.	Participants selected retrospectively from registry from two time periods with/without restrictions. Participants selected prospectively from two time	
Mikkelsen2014	All outcomes	Ν	-	-	ΡΥ	-	Low	No reasons to suspect selection bias.	periods with/without restrictions Participants in one group selected prospectively and	
vanderWeegen2019	All outcomes	PN	-	-	ΡΥ	-	Low	No reasons to suspect selection bias.	the other retrospectively.	1

3. Bias in classification o	of interventions
-----------------------------	------------------

Study	Outcome	2 1	3.2	2 2	Risk of bias	Comment and supporting	additional	Comment to ROBINS-I
Study	Outcome	5.1	5.2	5.5	judgement	quote	comment	Comment to KOBINS-I
						3.1. No informaton		
Allen2018	All outcomes	PN	Υ	Ν	Low	about restrictions		
Lightfoot2020	All outcomes	PN	Υ	Ν	Low			
Mikkelsen2014	All outcomes	Υ	Υ	Ν	Low			
						3.1. reference to earlier		
vanderWeegen2019	All outcomes	NI	Υ	Ν	Low	publication		

## 4. Bias due to departures from intended interventions

	Study	Outcome	4.1	4.2	4.3	4.4	4.5 4	1.6	Risk of bias udgement	Comment and supporting quote	additional comment	Comment to ROBINS-I
Allen2018		All outcomes	NI	NI	-	-		N		NI on deviations nor patient adherence to protocol		
Lightfoot2020		All outcomes	NI	-	-	-		N	<b>JI</b>	4.1 same reason for dropout between groups	NI on patient adherence to protocol	
Mikkelsen2014		All outcomes	NI	NI	-	-		N	JI	NI on deviations nor patient adherence to protocol		
vanderWeegen2019		All outcomes	NI		-	-		N		NI on deviations nor patient adherence to protocol	NI on deviations nor patient adherence to protocol	

# 5. Bias due to missind data

Study	Outcome	5.1	5.2	5.3	5.4	5.5	Risk of bias judgement		 Comment to ROBINS-I
								5.1 exclusion due to missing	
Allen2018	All outcomes	PN	PN	NI	NI	Ν	Serious	outcome data 5.1 unable to	
Lightfoot2020	All outcomes	N	PN	PN	Υ	Ν	Moderate	contact proportion for missing data differs across	
Mikkelsen2014 vanderWeegen2019	All outcomes All outcomes	PN Y	PN N	Y N	N -	N -	Moderate low	groups	

## 6. Bias in measurement of outcomes

Study	Outcome	6.1	6.2	2 6.3	6.4	Risk of bias judgement	Comment and supporting quote	additional comment	Comment to ROBINS-I
Allen2018	All outcomes	Ν	Υ	Υ	PN	Low			
Lightfoot2020	All outcomes	Ν	Υ	Υ	PN	Low			
Mikkelsen2014	All outcomes	Ν	Υ	Υ	PN	Low			
vanderWeegen2019	All outcomes	Ν	Υ	Υ	PN	Low			

## 7. Bias in selection of the reported result

Study	Outcome	7.1	7.2	7.3	Risk of bias judgement	Comment and supporting quote	additional comment	to ROBINS-
Allen2018	All outcomes	PN	PN	PN	Moderate	No protocol pre-registered No deviations from published		
Lightfoot2020	All outcomes	ΡN	PN	PN	Low	protocol.		
Mikkelsen2014 vanderWeegen2019	All outcomes All outcomes				Moderate Moderate	No protocol pre-registered No protocol pre-registered		

Commont