Supplementary Table 3A	. Description and	decision criteria	for each domain	n in ROBINS-I
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Bias domain	Explanation	Judgments		
Bias due to	1. Is there potential for confounding of the	1. Low risk of bias: No bias expected due to		
confounding	effect of intervention in this study?	confounding, including time-varying confounding.		
	2. Did the authors use a	2. Moderate risk of bias: Confounding is expected:		
	multivariable-adjusted analysis method that	including at least 5 factors of the following factors:		
	controlled at least for the important	age, body mass index, etiology, location of the		
	confounding domains (age, body mass index,	stricture, length of the stricture, prior intervention		
	etiology, location of the stricture, length of the	management, others (i.e. comorbidities,		
	stricture, prior intervention management,	socio-economic status) and have been appropriately		
	others)?	controlled for in a multivariable-adjusted analysis.		
	3. Were confounding domains that were	3. Serious risk of bias: 3-4 above-mentioned factors		
	controlled for measured validly and reliably	were measured or appropriately controlled for.		
	by the variables available in this study?	4. Critical risk of bias: less than 3 above-mentioned		
	4. Did the authors control for any post-	factors were measured or appropriately controlled		
	intervention variables that could have been	for.		
	affected by the intervention?	5. No information: No information on which		
	5. Did the authors use an appropriate analysis	confounders have been controlled for.		
	method that controlled for all the important			
	confounding domains and for time-varying			
	confounding?			
	6. Were confounding domains that were			
	controlled for measured validly and reliably			
	by the variables available in this study?			
Bias in selection	1. Was selection of participants into the study	1. Low risk of bias: All participants who would have		
of participants	based on participant characteristics observed	been eligible for the target study were included in		
into the study	after the start of intervention?	the study.		
	2. Were the post-intervention variables that	2. Moderate risk of bias: Selection into the study		
	influenced selection likely to be associated	may have been related to exposure and outcome and		
	with intervention?	the authors used appropriate methods to correct for		
	3. Were the postintervention variables that	the selection bias.		
	influenced selection likely to be influenced by	3. Serious risk of bias: Selection into the study was		
	the outcome or a cause of the outcome?	related to intervention and outcome and this could		
	4. Do start of follow-up and start of	not be corrected for in the analyses; or the start of		
	intervention coincide for most participants?	follow-up and start of exposure do not coincide and		
	5. Were adjustment techniques used that are	the rate ratio is not constant over time.		
	history to correct for the presence of selection	4. Unucal risk of blas: Selection into the study was		
		and this could not be corrected for in the analysis of		
		and this could not be confected for in the analyses; or		
		a substantial amount of follow-up time is likely to be missing from analyses 3 the rate ratio is not constant		
		over time		
		5. No information: No information is reported about		
	5. Were adjustment techniques used that are likely to correct for the presence of selection biases?	 the rate ratio is not constant over time. 4. Critical risk of bias: Selection into the study was very strongly related to intervention and outcome and this could not be corrected for in the analyses; or a substantial amount of follow-up time is likely to be missing from analyses 3.the rate ratio is not constant over time. 5. No information: No information is reported about 		

		selection of participants into the study.
Bias in 1	1. Were intervention groups clearly defined?	1. Low risk of bias: The patient clearly underwent
classification of 2	2. Was the information used to define	urethral balloon dilation, and no measurement error
interventions i	intervention groups recorded at the start of the	is expected in its assessment.
i	intervention?	2. Moderate risk of bias: Intervention status is well
	3. Could classification of intervention status	defined and some aspects of the assignments of
1	have been affected by knowledge of the	intervention status were determined retrospectively.
	outcome or risk of the outcome?	3. Serious risk of bias: Intervention status is not well
		defined; or major aspects of the assignments of
		intervention status were determined in a way that
		could have been affected by knowledge of the
		outcome.
		4. Critical risk of bias: An extremely high amount of
		misclassification of intervention status (i.e. because
		of unusually strong recall biases).
		5. No information: No definition of the intervention
		or no explanation of the source of information about
		intervention status is reported.
Bias due to	1. Were there deviations from the intended	1. Low risk of bias: Patients did not receive other
deviations from i	intervention beyond what would be expected	invasive urethral stricture treatments between the
intended i	in usual practice?	time they underwent balloon dilatation and the
interventions 2	2. Were these deviations from intended	follow-up period to assess success.
i	intervention unbalanced between groups and	2. Moderate risk of bias: There were deviations
1	likely to have affected the outcome?	from usual practice, but their impact on the outcome
		is expected to be slight.
		3. Serious or critical risk of bias: There were
		deviations from usual practice that were unbalanced
		between the intervention groups and likely to have
		affected the outcome.
		4. Critical risk of bias: There were substantial
		deviations from usual practice that were unbalanced
		between the intervention groups and likely to have
		affected the outcome.
		5. No information: No information on deviations
		from the intervention is reported.
Bias due to	1. Were outcome data available for all, or	1. Low risk of bias: Little loss-to-follow-up and data
missing data 1	nearly all, participants?	on intervention and other variables were reasonably
2	2. Were participants excluded due to missing	complete (<10% missing data) and was unlikely to
	data on intervention status?	introduce bias; or the analysis addressed missing
3	3. Were participants excluded due to missing	data and is likely to have removed any risk of bias.
	data on other variables needed for the	2. Moderate risk of bias: There is a proportion of
2	analysis?	missing data in the original cohort or a high
2	4. Are the proportion of participants and	proportion of loss-to-follow-up; and the analysis is
1	reasons for missing data similar across	unlikely to have removed the risk of bias arising

	interventions?	from the missing data (i.e. using logistic regression).
	5. Is there evidence that results were robust to	3. Serious risk of bias: High proportions (>50%) of
	the presence of missing data?	missing data; and the analysis is unlikely to have
		removed the risk of bias arising from the missing
		data; or missing data were addressed inappropriately
		in the analysis; or the nature of the missing data
		means that the risk of bias cannot be removed
		through appropriate analysis.
		4. Critical risk of bias: There were critical
		differences between interventions in participants
		with missing data; and missing data were not, or
		could not, be addressed through appropriate analysis.
		5. No information: No information is reported about
		missing data or the potential for data to be missing.
Bias in	1. Could the outcome measure have been	1. Low risk of bias: The methods of outcome
measurement of	influenced by knowledge of the intervention	assessment were comparable across intervention
outcomes	received?	groups; and the outcome measure was unlikely to be
	2. Were outcome assessors aware of the	influenced by knowledge of the intervention status
	intervention received by study participants?	of study participants; and any error in measuring the
	3. Were the methods of outcome assessment	outcome is unrelated to intervention status (i.e.,
	comparable across intervention groups?	objective measures such as confirmed medical
	4. Were any systematic errors in measurement	records, record linkage).
	of the outcome related to intervention	2. Moderate risk of bias: The methods of outcome
	received?	assessment were comparable across intervention
		groups; and any error in measuring the outcome may
		be minimally related to intervention status or if the
		outcome measure was not reliable measured (i.e.
		confirmed records are not available for the whole
		study population).
		3. Serious risk of bias: The methods of outcome
		assessment were not comparable across intervention
		groups; or the outcome measure was subjective (i.e.
		vulnerable to influence by knowledge of the
		intervention received by study participants); and
		error in measuring the outcome was related to
		intervention status.
		4. Critical risk of bias: The methods of outcome
		assessment were so different that they cannot
		reasonably be compared across intervention groups.
		5. No information: No information is reported about
		the methods of outcome assessment.
Bias in selection	1. Is the reported effect estimate likely to be	1. Low risk of bias: There is a clear description of
of the reported	selected from multiple analyses of the	all analyses and the analyses are consistent and all
result	intervention-outcome relationship?	reported results correspond to all intended outcomes,

2. Is the reported effect estimate likely to be selected from different subgroups?analyses and sub-cohorts.2. Moderate risk of bias: The analyses are clearly defined; and there is an indication of selection of the reported analysis from among multiple analyses; and there is an indication of selection of the cohort or subgroups for analysis and reporting on the basis of the results (i.e. estimates not shown for all analyses).3. Serious risk of bias: There is a high risk of selective reporting from among multiple analyses; or the cohort or subgroup is selected from a larger study for analysis and appears to be reported based on the results.4. Critical risk of bias: There is evidence or strong supicion of selective reporting of results; and the unreported results are likely to be substantially different from the reported results.5. No information: There is too little information to make a judgment.	· · · · · · · · · · · · · · · · · · ·	
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 subgroups for analysis and reporting on the basis of the results (i.e. estimates not shown for all analyses). Serious risk of bias: There is a high risk of selective reporting from among multiple analyses; or the cohort or subgroup is selected from a larger study for analysis and appears to be reported based on the results. Critical risk of bias: There is evidence or strong suspicion of selective reporting of results; and the unreported results are likely to be substantially different from the reported results. No information: There is too little information to make a judgment. 		there is an indication of selection of the cohort or
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 4. Critical risk of bias: There is evidence or strong suspicion of selective reporting of results; and the unreported results are likely to be substantially different from the reported results. 5. No information: There is too little information to make a judgment. 		on the results.
suspicion of selective reporting of results; and the unreported results are likely to be substantially different from the reported results. 5. No information: There is too little information to make a judgment.		4. Critical risk of bias: There is evidence or strong
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different from the reported results.5. No information: There is too little information to make a judgment.		unreported results are likely to be substantially
5. No information: There is too little information to make a judgment.		different from the reported results.
make a judgment.		5. No information: There is too little information to
		make a judgment.

Overall judgment

1. Low risk of bias

The study is judged to be at a low risk of bias for all domains.

2. Moderate risk of bias

The study is judged to be at low or moderate risk of bias for all domains.

3. Serious risk of bias

The study is judged to be at serious risk of bias in at least one domain, but not at critical risk in any domain.

4. Critical risk of bias

The study is judged to be at critical risk of bias in at least one domain.

Supplementary Table 3B. Quality assessment results using the ROBINS-I tool

Study	Bias due to	Bias in	Bias in	Bias due to	Bias due	Bias in	Bias in	Overall
	confounding	selection of	classification	deviations	to	measurement	selection	judgment
		participants	of	from	missing	of outcomes	of the	
		into the	interventions	intended	data		reported	
		study		interventions			result	
Virasoro,	Moderate	Moderate	Low	Low	Moderate	Moderate	Moderate	Serious
Ramon et al.								
2022								
Beeder, L. A.	Moderate	Serious	Moderate	Low	Low	Serious	Moderate	Serious
et al. 2022								
Alibekov, M.	Serious	Serious	Moderate	Low	Low	Serious	Serious	Serious
M. et al. 2022								
Yi, Y. A. et al.	Moderate	Serious	Moderate	Low	Low	Moderate	Moderate	Serious
2020								
Kumano, Y.	Serious	Serious	Moderate	Low	Low	Serious	Moderate	Serious
et al. 2019								
Zhou, Y. et al.	Moderate	Serious	Moderate	Low	Low	Moderate	Moderate	Serious
2016								
Yu, S. C. et al.	Moderate	Serious	Moderate	Low	Low	Moderate	Moderate	Serious
2016								
Chhabra, J. S.	Moderate	Serious	Moderate	Low	Low	Moderate	Moderate	Serious
et al. 2016								
Ishii, Gen et	Serious	Critical	Moderate	Low	Low	Serious	Moderate	Critical
al. 2015								
Mao, D. et al.	Moderate	Serious	Moderate	Low	Low	Moderate	Moderate	Serious
2014								
Vyas, J. B. et	Serious	Serious	Moderate	Low	Low	Moderate	Moderate	Serious
al. 2013								
Alguersuari,	Serious	Serious	Moderate	Low	Low	Serious	Serious	Serious
A. et al. 2012								
MacDiarmid,	Serious	Serious	Moderate	Low	Moderate	Serious	Moderate	Serious
S. A. et al.								
2000								
Mohammed,	Critical	Serious	Low	Low	Moderate	Serious	Serious	Critical
S. H. et al.								
1988								