



Table S1. PRISMA 2009 Checklist

Section/topic	#	Checklist item	Reported on page #
TITLE			
Title	1	Identify the report as a systematic review, meta-analysis, or both.	1
ABSTRACT			
Structured summary	2	Provide a structured summary including, as applicable: background; objectives; data sources; study eligibility criteria, participants, and interventions; study appraisal and synthesis methods; results; limitations; conclusions and implications of key findings; systematic review registration number.	1
INTRODUCTION			
Rationale	3	Describe the rationale for the review in the context of what is already known.	1-2
Objectives	4	Provide an explicit statement of questions being addressed with reference to participants, interventions, comparisons, outcomes, and study design (PICOS).	3
METHODS			
Protocol and registration	5	Indicate if a review protocol exists, if and where it can be accessed (e.g., Web address), and, if available, provide registration information including registration number.	NA
Eligibility criteria	6	Specify study characteristics (e.g., PICOS, length of follow-up) and report characteristics (e.g., years considered, language, publication status) used as criteria for eligibility, giving rationale.	3
Information sources	7	Describe all information sources (e.g., databases with dates of coverage, contact with study authors to identify additional studies) in the search and date last searched.	2-3
Search	8	Present full electronic search strategy for at least one database, including any limits used, such that it could be repeated.	2
Study selection	9	State the process for selecting studies (i.e., screening, eligibility, included in systematic review, and, if applicable, included in the meta-analysis).	4-5
Data collection process	10	Describe method of data extraction from reports (e.g., piloted forms, independently, in duplicate) and any processes for obtaining and confirming data from investigators.	2
Data items	11	List and define all variables for which data were sought (e.g., PICOS, funding sources) and any assumptions and simplifications made.	3
Risk of bias in individual studies	12	Describe methods used for assessing risk of bias of individual studies (including specification of whether this was done at the study or outcome level), and how this information is to be used in any data synthesis.	2-4
Summary measures	13	State the principal summary measures (e.g., risk ratio, difference in means).	3
Synthesis of results	14	Describe the methods of handling data and combining results of studies, if done, including measures of consistency (e.g., I^2) for each meta-analysis.	3-4



PRISMA 2009 Checklist

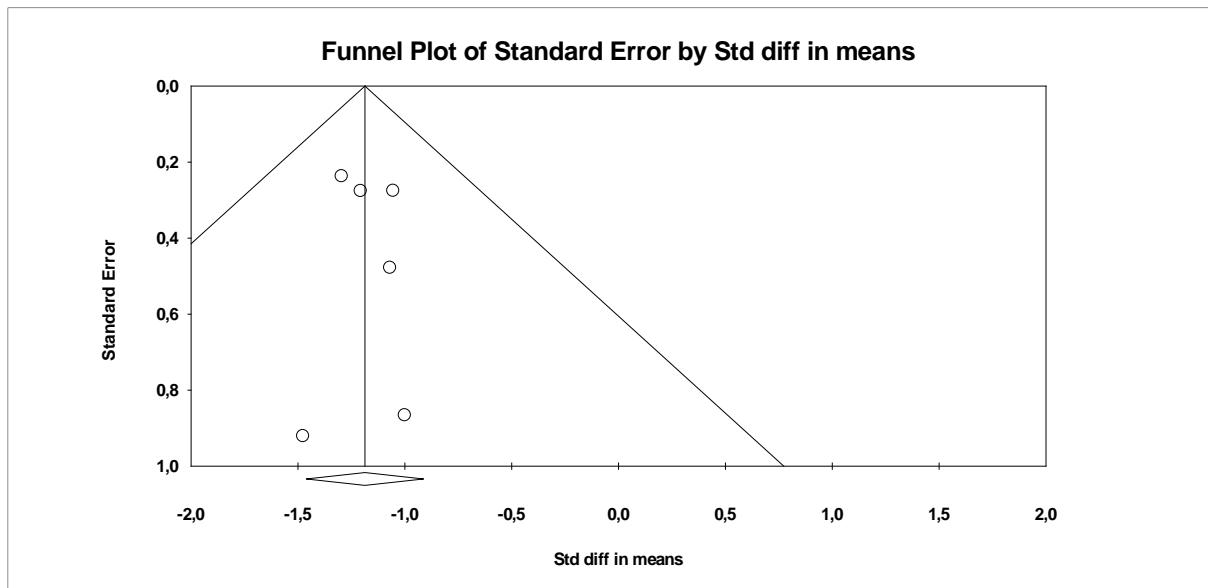
Section/topic	#	Checklist item	Reported on page #
Risk of bias across studies	15	Specify any assessment of risk of bias that may affect the cumulative evidence (e.g., publication bias, selective reporting within studies).	3
Additional analyses	16	Describe methods of additional analyses (e.g., sensitivity or subgroup analyses, meta-regression), if done, indicating which were pre-specified.	4
RESULTS			
Study selection	17	Give numbers of studies screened, assessed for eligibility, and included in the review, with reasons for exclusions at each stage, ideally with a flow diagram.	4-5
Study characteristics	18	For each study, present characteristics for which data were extracted (e.g., study size, PICOS, follow-up period) and provide the citations.	6-7
Risk of bias within studies	19	Present data on risk of bias of each study and, if available, any outcome level assessment (see item 12).	4-17
Results of individual studies	20	For all outcomes considered (benefits or harms), present, for each study: (a) simple summary data for each intervention group (b) effect estimates and confidence intervals, ideally with a forest plot.	4-17
Synthesis of results	21	Present results of each meta-analysis done, including confidence intervals and measures of consistency.	4-17
Risk of bias across studies	22	Present results of any assessment of risk of bias across studies (see Item 15).	4-17
Additional analysis	23	Give results of additional analyses, if done (e.g., sensitivity or subgroup analyses, meta-regression [see Item 16]).	4-17
DISCUSSION			
Summary of evidence	24	Summarize the main findings including the strength of evidence for each main outcome; consider their relevance to key groups (e.g., healthcare providers, users, and policy makers).	17-18
Limitations	25	Discuss limitations at study and outcome level (e.g., risk of bias), and at review-level (e.g., incomplete retrieval of identified research, reporting bias).	19
Conclusions	26	Provide a general interpretation of the results in the context of other evidence, and implications for future research.	19
FUNDING			
Funding	27	Describe sources of funding for the systematic review and other support (e.g., supply of data); role of funders for the systematic review.	20

From: Moher D, Liberati A, Tetzlaff J, Altman DG, The PRISMA Group (2009). Preferred Reporting Items for Systematic Reviews and Meta-Analyses: The PRISMA Statement. PLoS Med 6(7): e1000097. doi:10.1371/journal.pmed.1000097

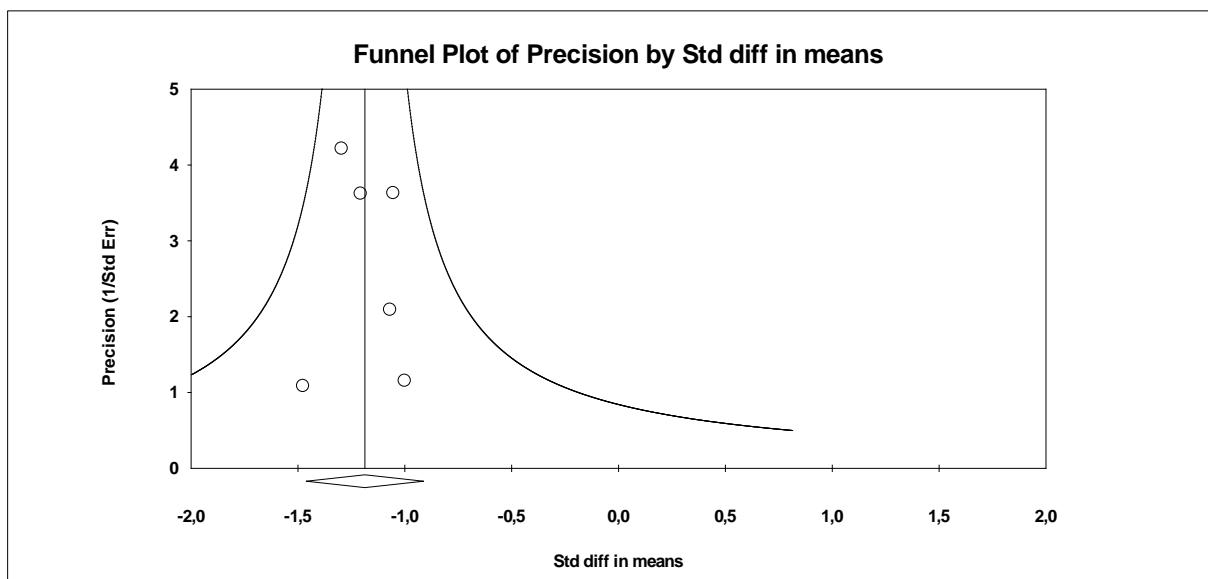
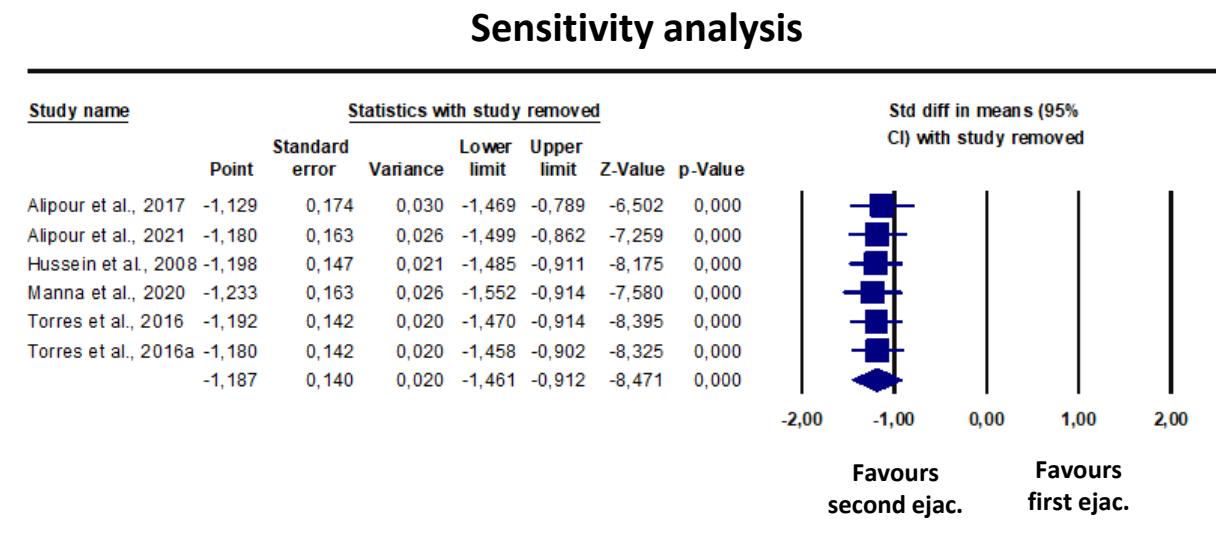
For more information, visit: www.prisma-statement.org.

Supplementary Figure S1. Funnel plot (A) and sensitivity analysis (B) of studies that evaluated the effects of a very short abstinence period on seminal fluid volume in normozoospermic men

A

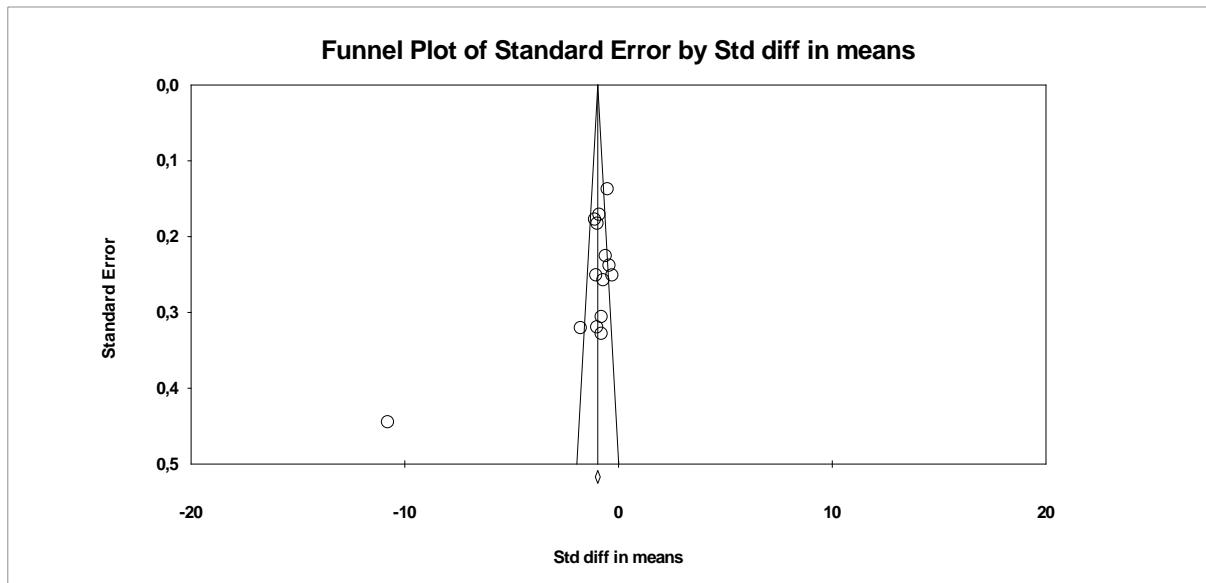


B

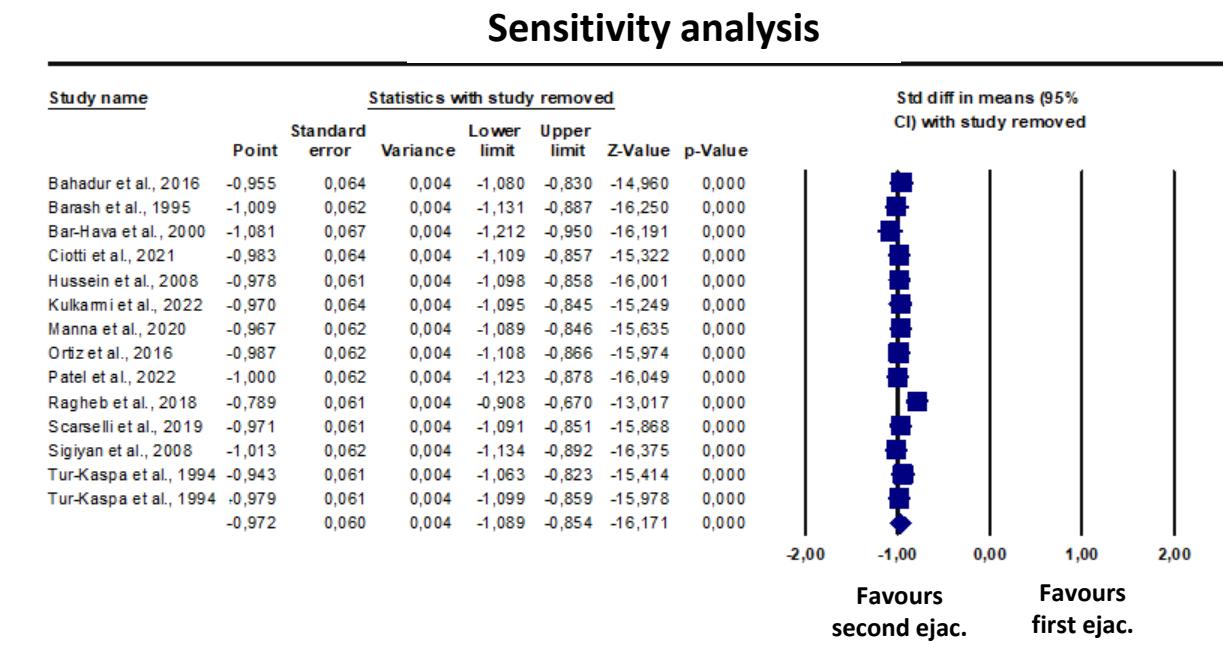


Supplementary Figure S2. Funnel plot (A) and sensitivity analysis (B) of studies that evaluated the effects of a very short abstinence period on seminal fluid volume in Oligo- and/or Astheno- and/or Terato- zoospermic men

A

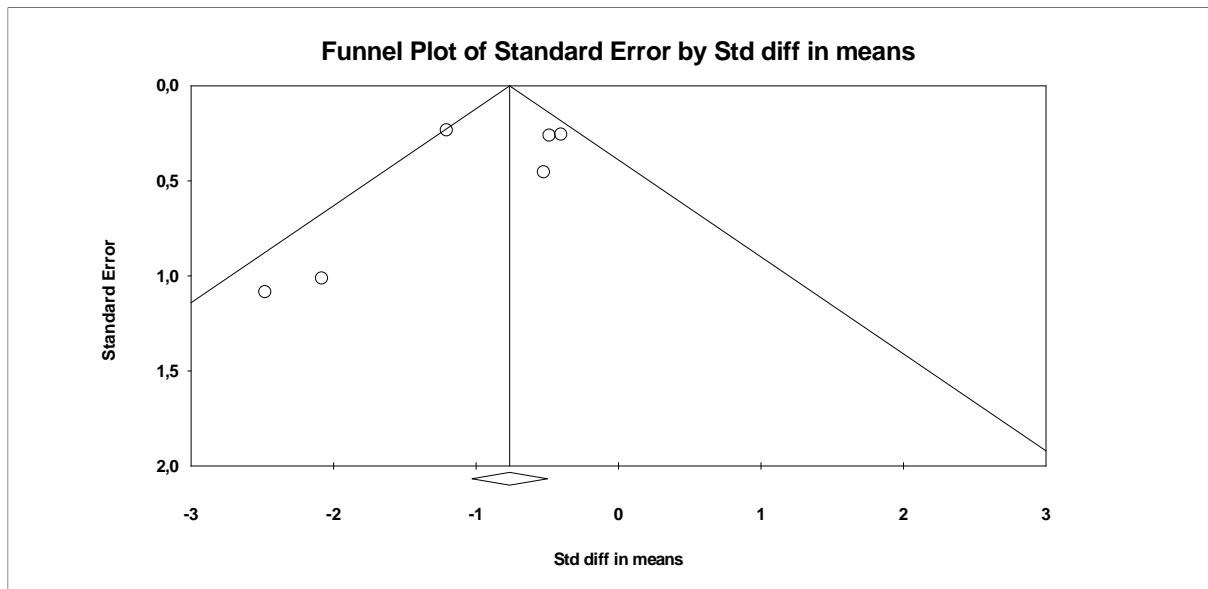


B

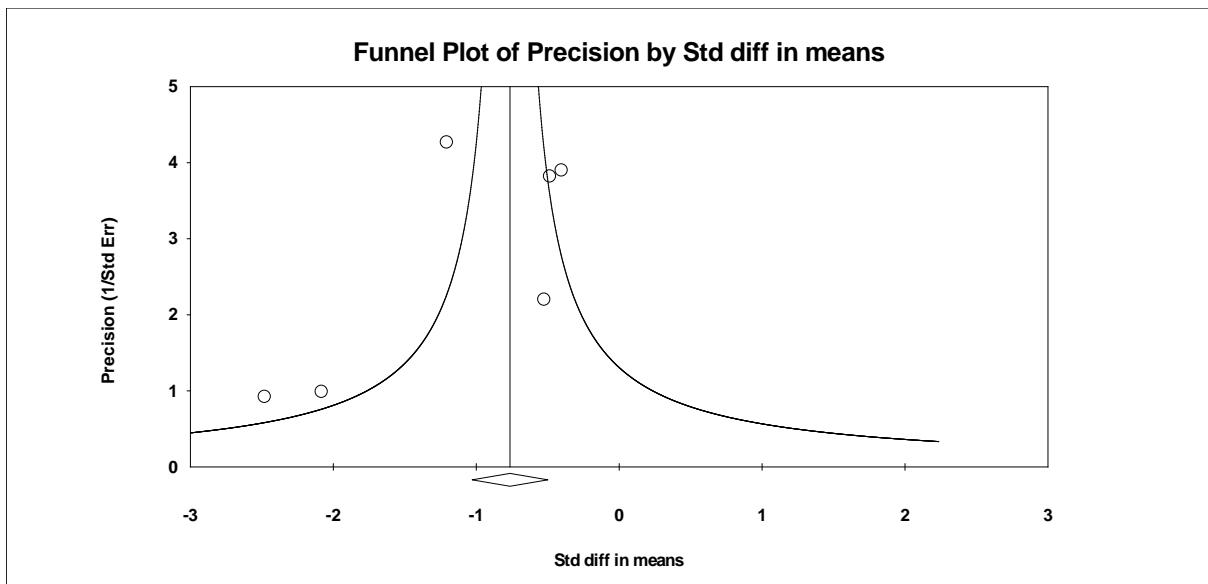
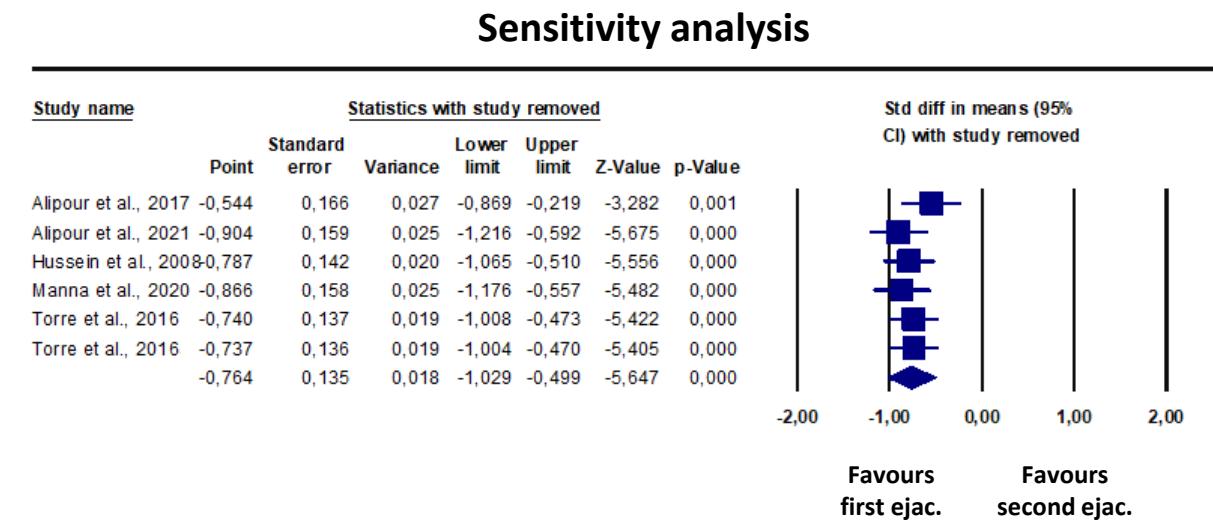


Supplementary Figure S3. Funnel plot (A) and sensitivity analysis (B) of studies that evaluated the effects of a very short abstinence period on sperm concentration in normozoospermic men

A

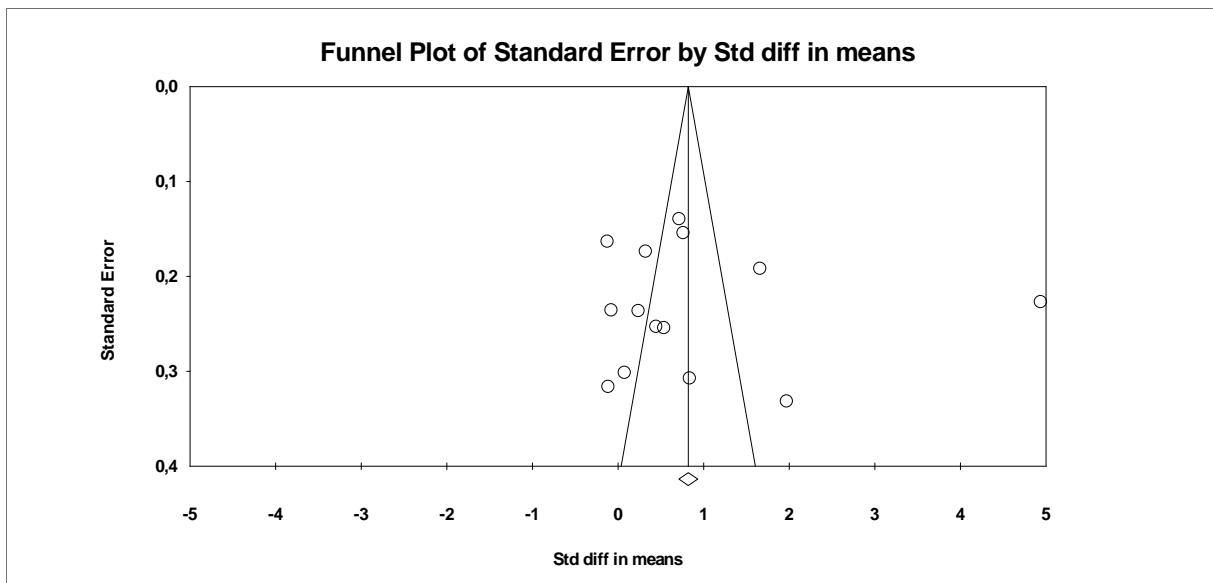


B

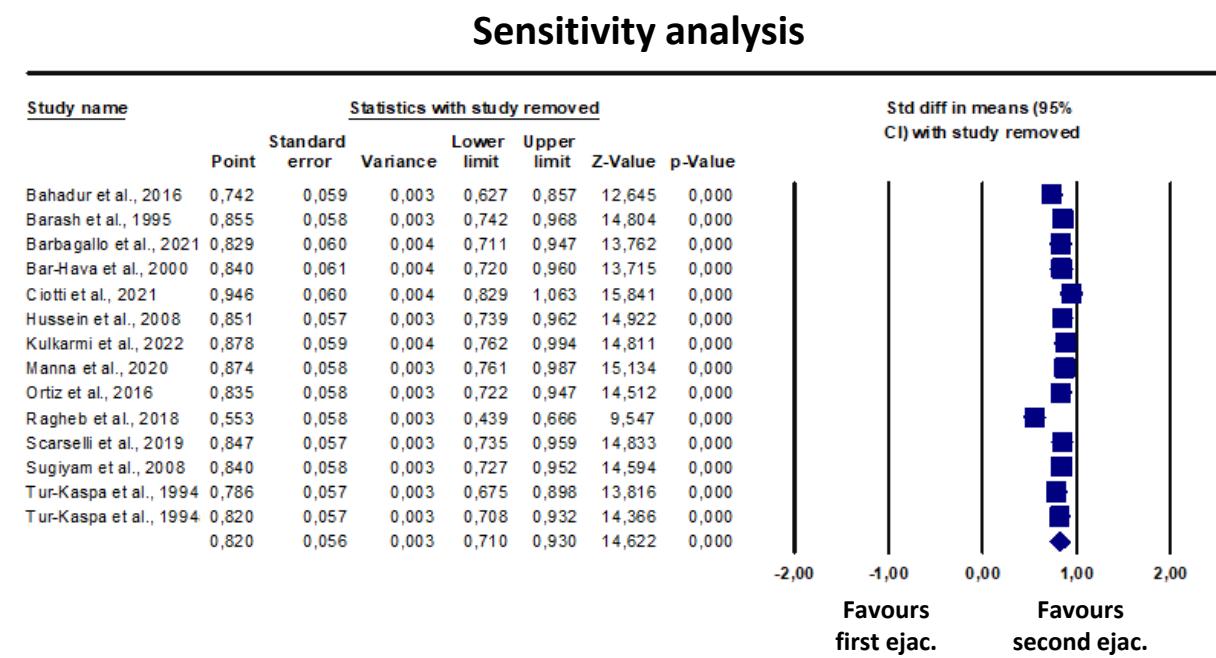


Supplementary Figure S4. Funnel plot (A) and sensitivity analysis (B) of studies that evaluated the effects of a very short abstinence period on sperm concentration in Oligo- and/or Astheno- and/or Terato- zoospermic men

A

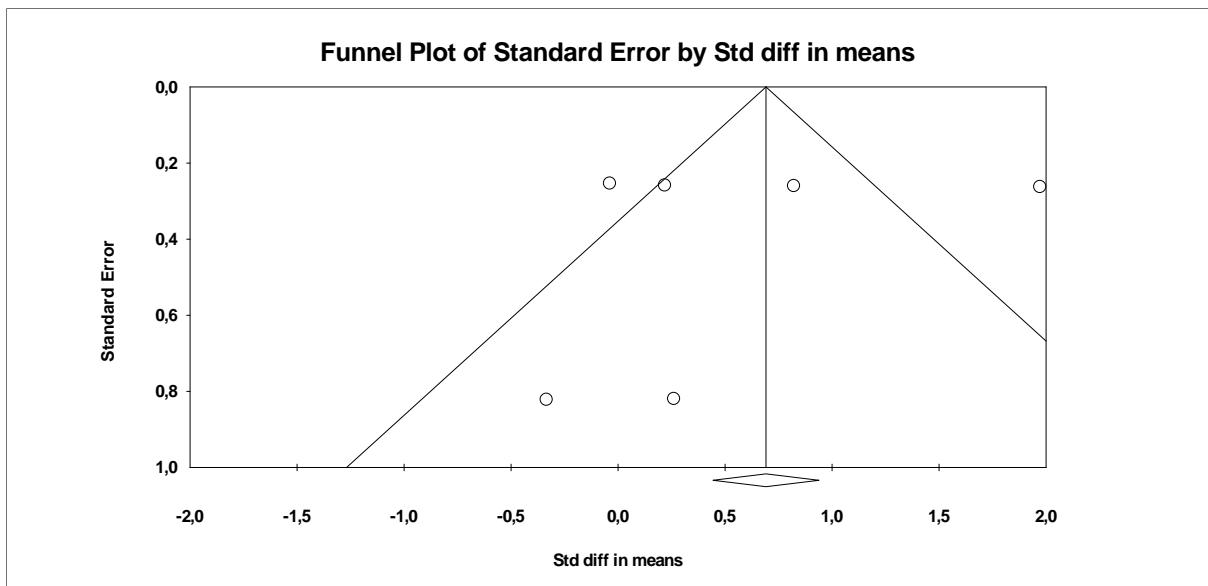


B

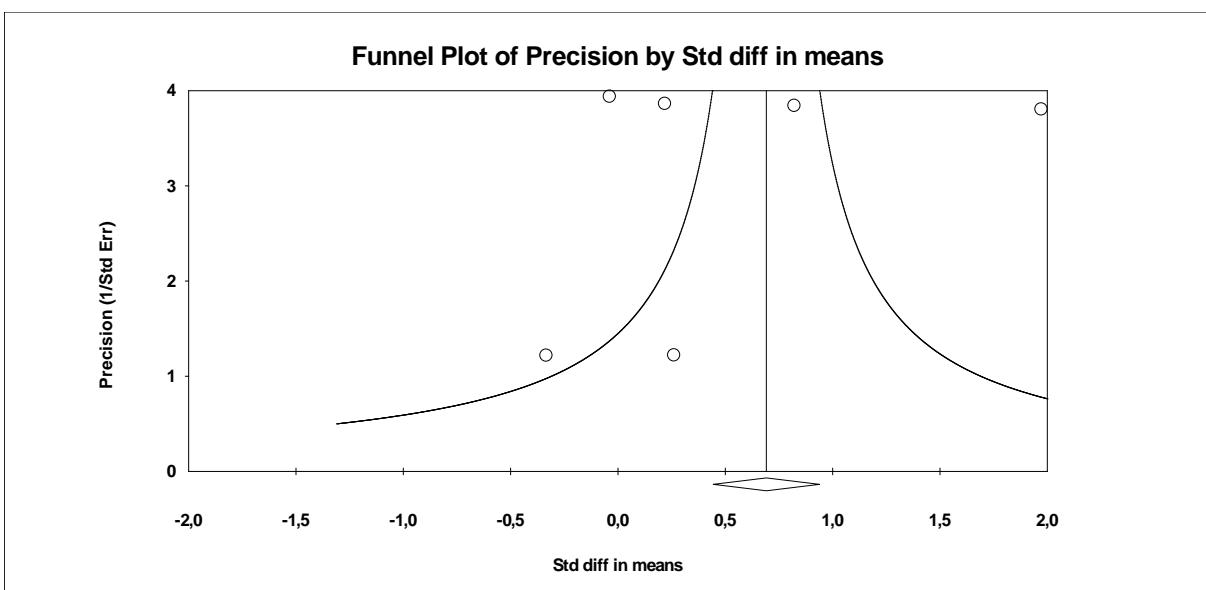
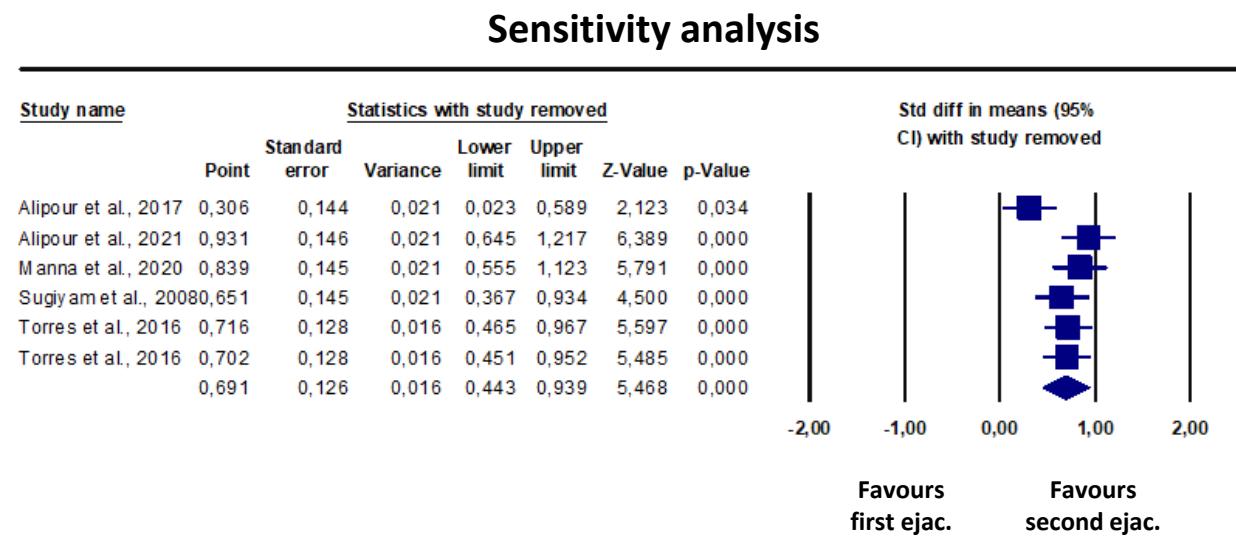


Supplementary Figure S5. Funnel plot (A) and sensitivity analysis (B) of studies that evaluated the effects of a very short abstinence period on sperm total motility in normozoospermic men

A

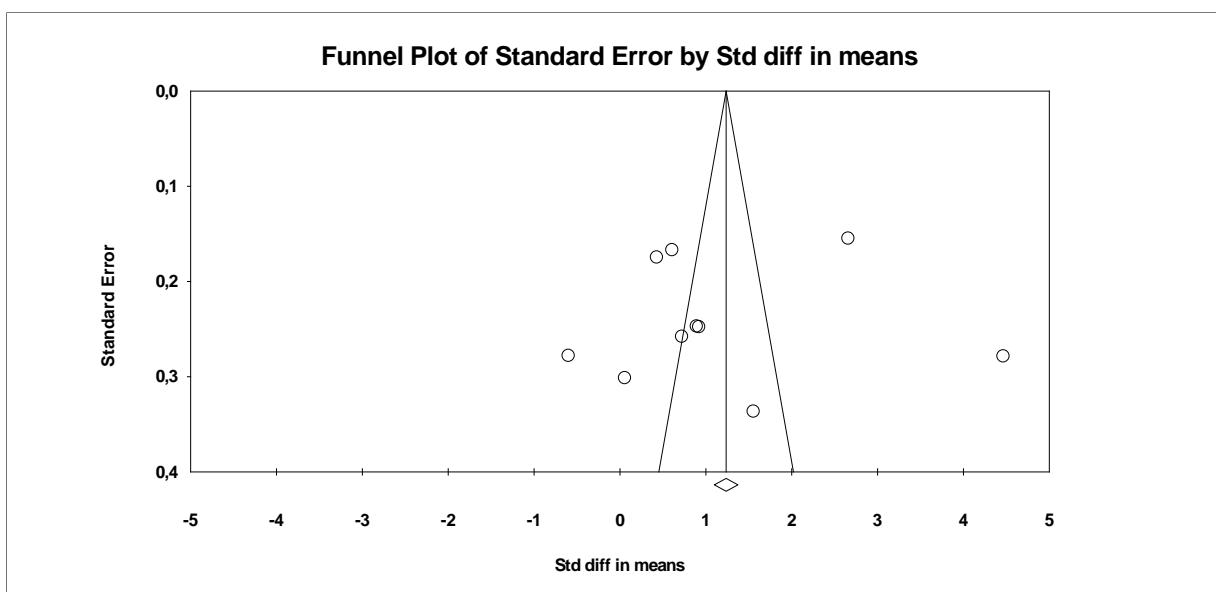


B

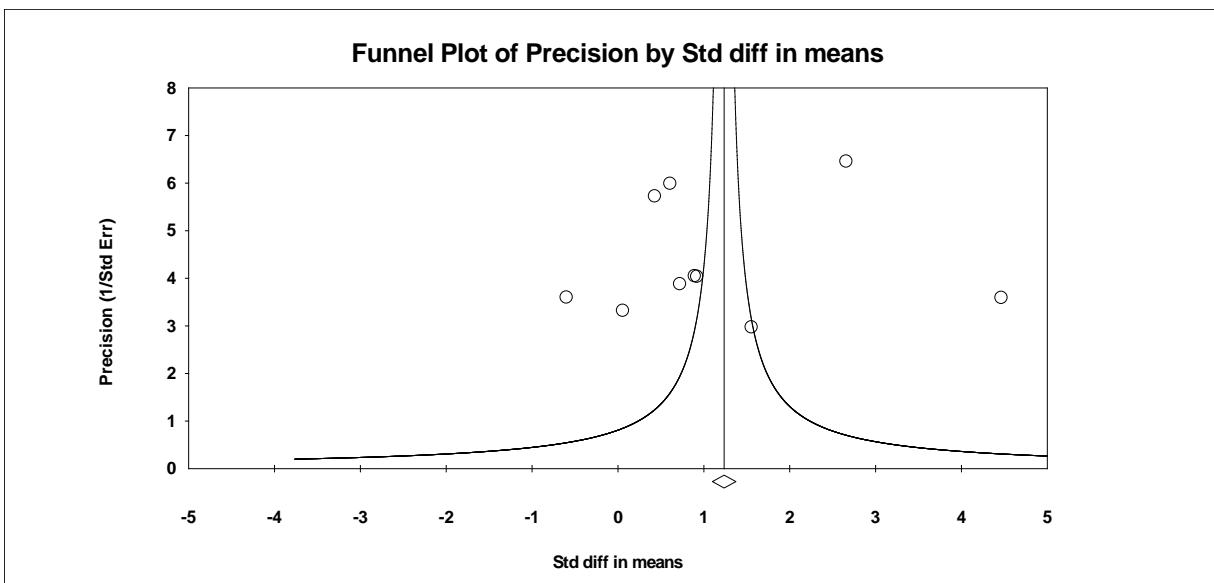
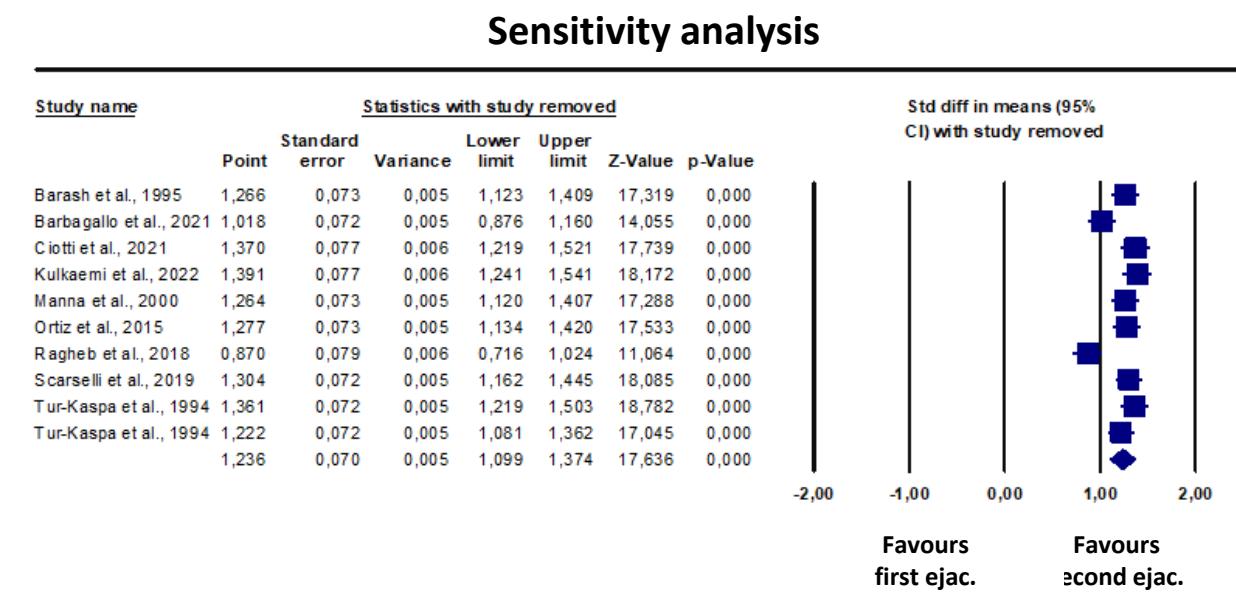


Supplementary Figure S6. Funnel plot (A) and sensitivity analysis (B) of studies that evaluated the effects of a very short abstinence period on sperm total motility in Oligo- and/or Astheno- and/or Terato- zoospermic men

A

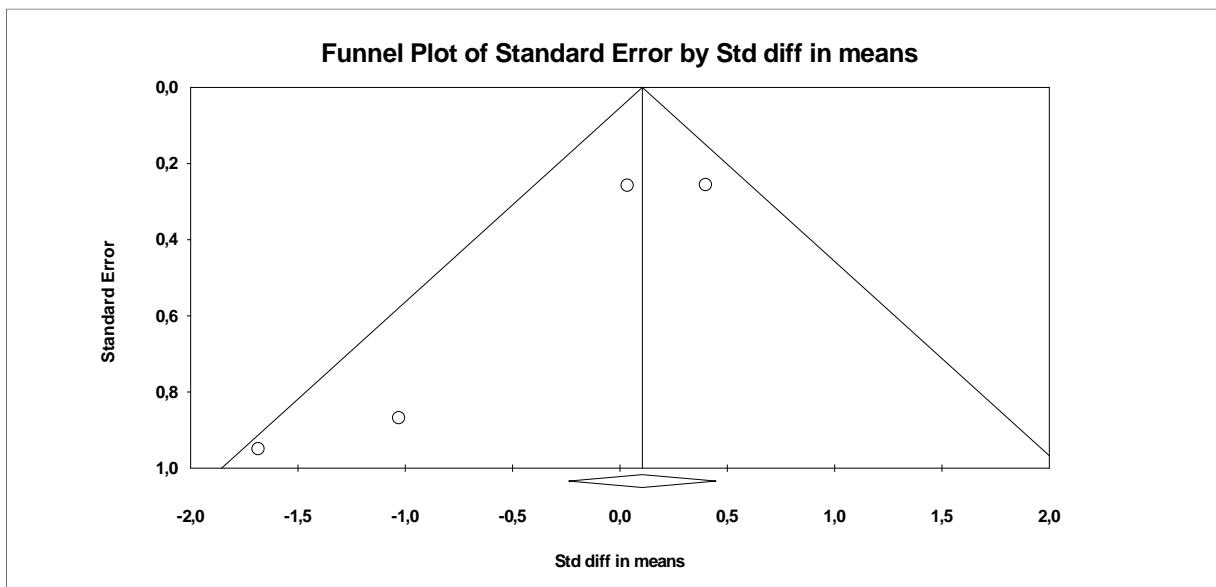


B



Supplementary Figure S7. Funnel plot (A) and sensitivity analysis (B) of studies that evaluated the effects of a very short abstinence period on sperm progressive motility in normozoospermic men

A



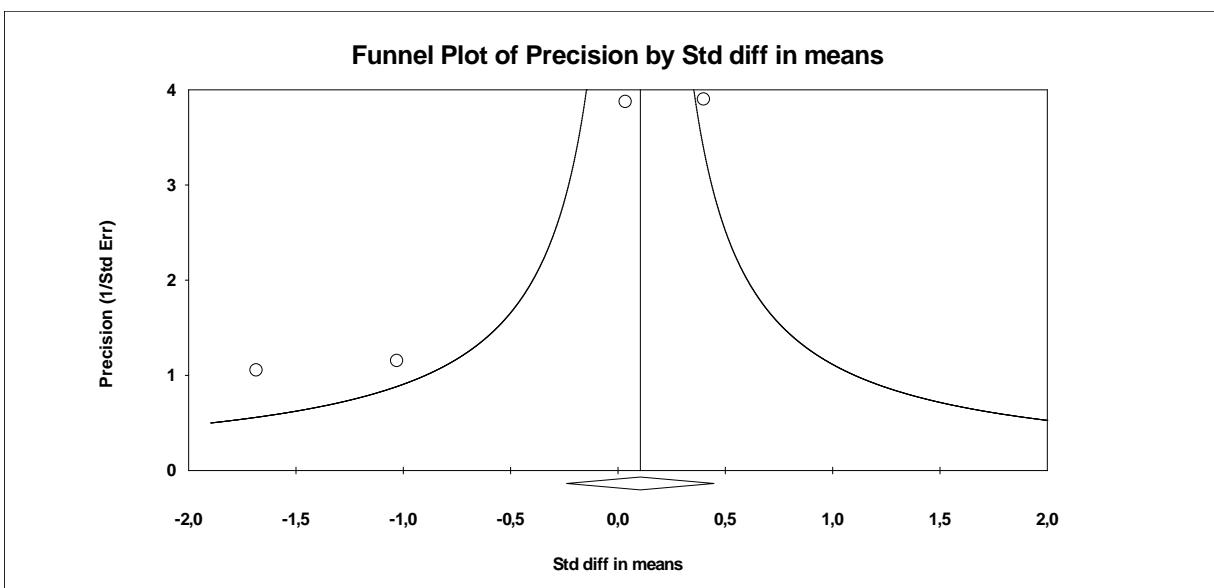
B

Sensitivity analysis

Study name	Statistics with study removed						Std diff in means (95% CI) with study removed
	Point	Standard error	Variance	Lower limit	Upper limit	Z-Value	
Alipour et al., 2021-0,154	0,240	0,057	-0,624	0,315	-0,644	0,520	
Manna et al., 2020 0,162	0,238	0,057	-0,305	0,629	0,681	0,496	
Torres et al., 2016 0,167	0,178	0,032	-0,182	0,516	0,938	0,348	
Torres et al., 2016 0,152	0,179	0,032	-0,198	0,503	0,851	0,394	
	0,104	0,175	0,031	-0,239	0,447	0,595	0,552

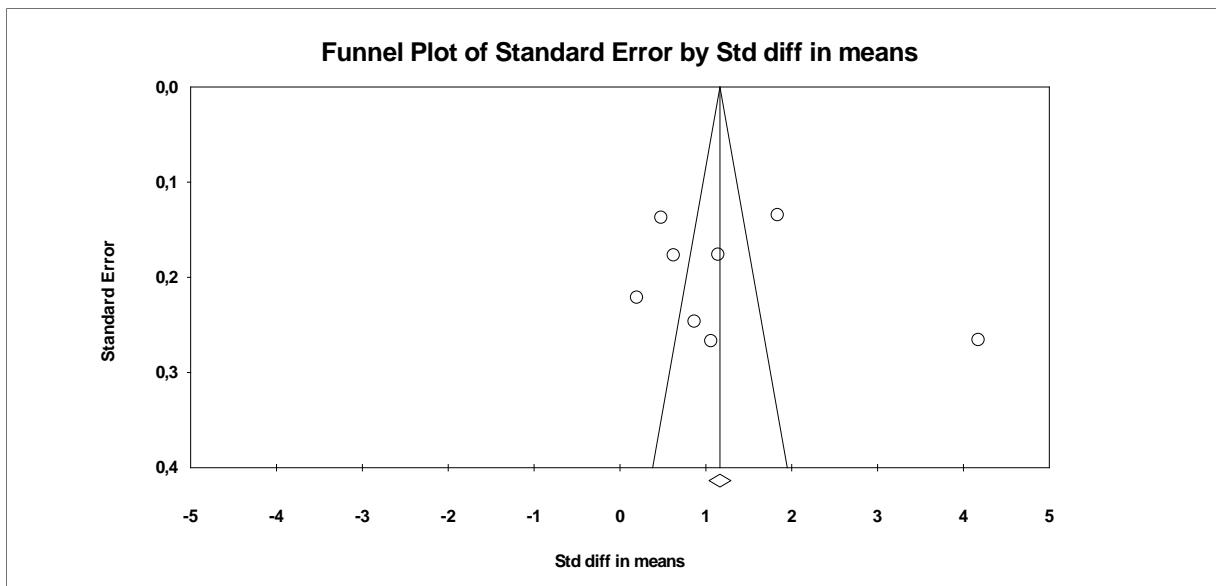
Favours second ejac.

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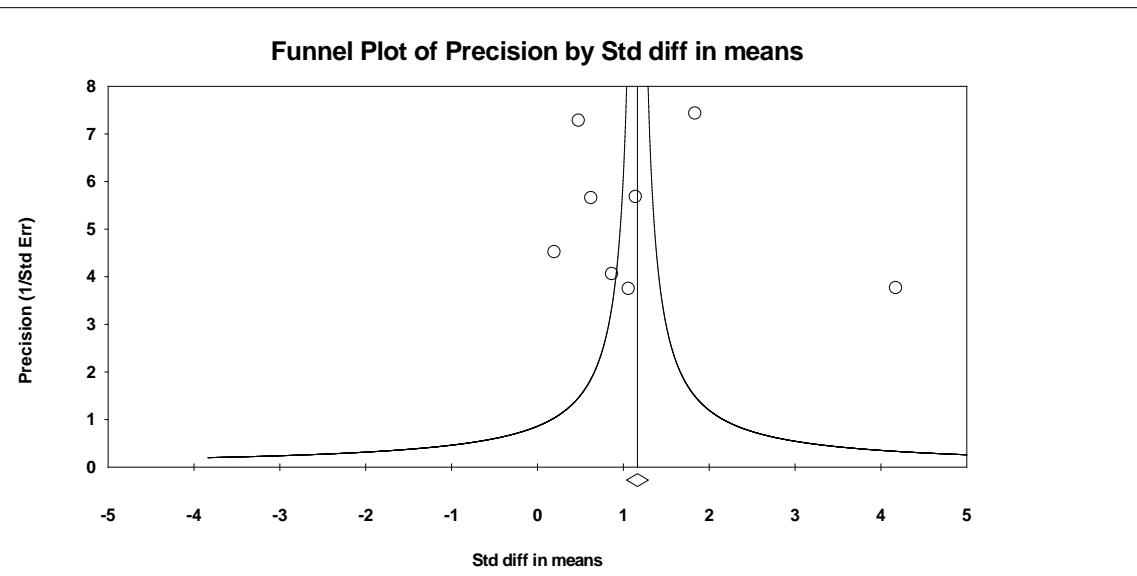
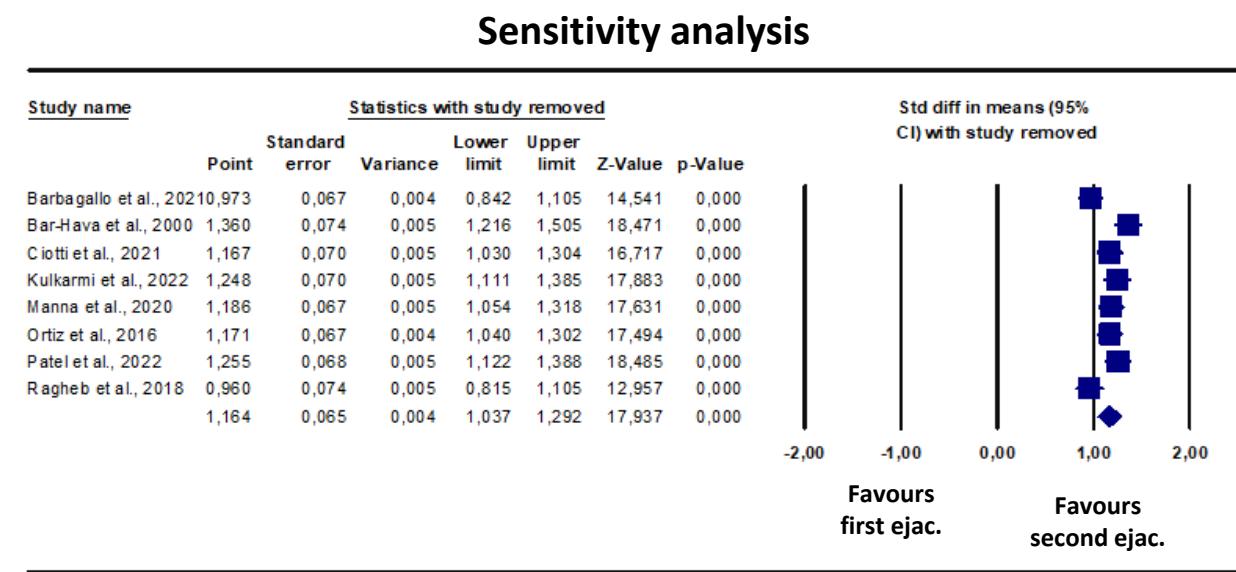


Supplementary Figure S8. Funnel plot (A) and sensitivity analysis (B) of studies that evaluated the effects of a very short abstinence period on sperm progressive motility in Oligo- and/or Astheno- and/or Terato- zoospermic men

A

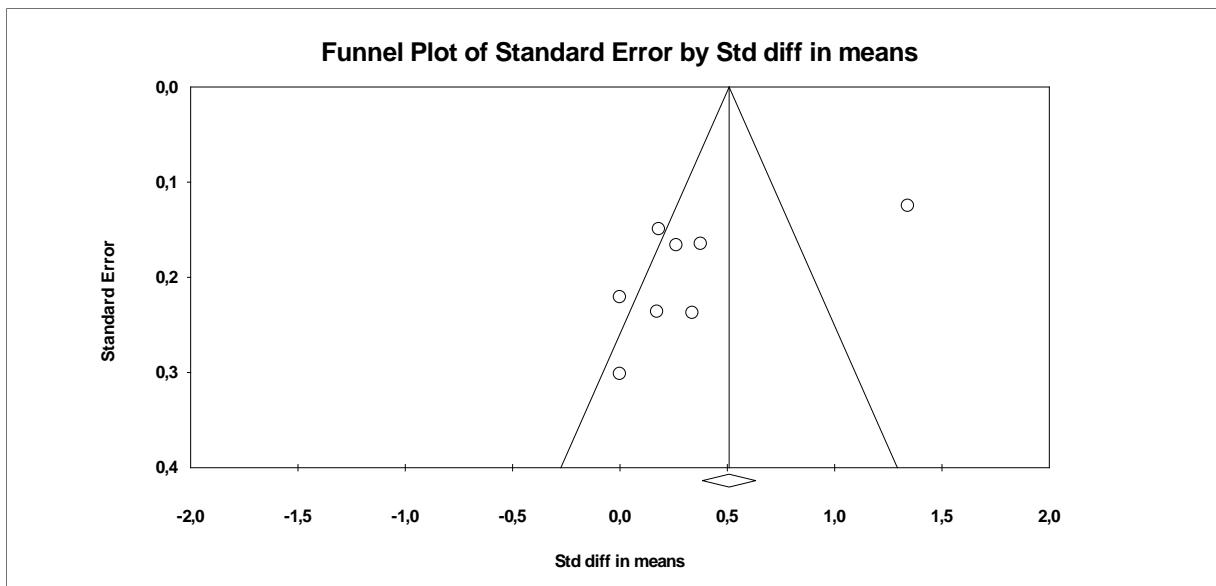


B

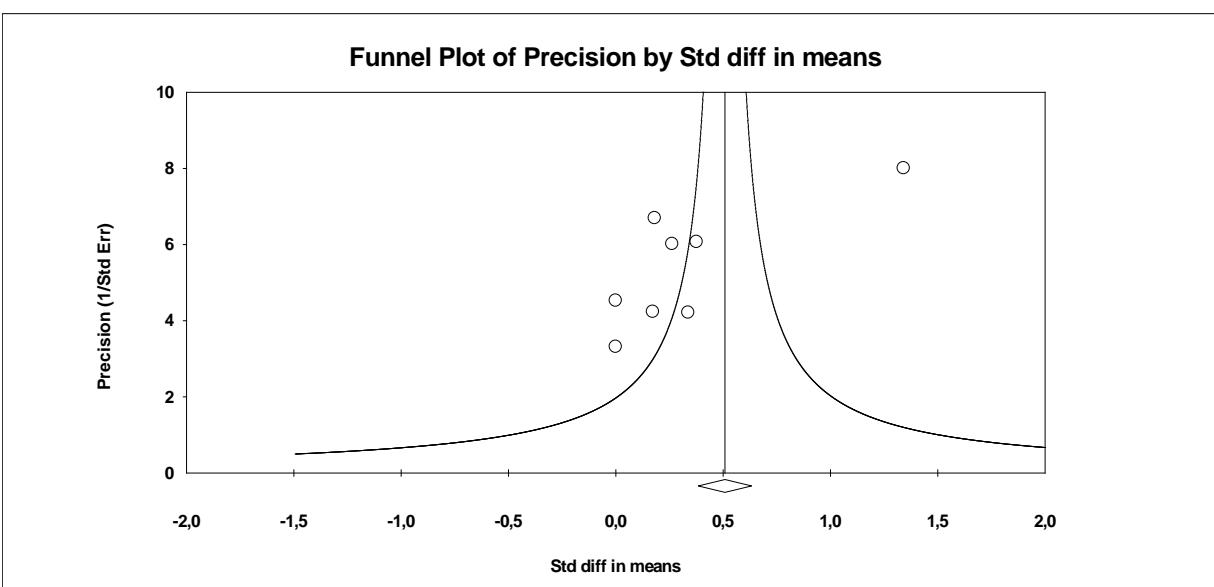
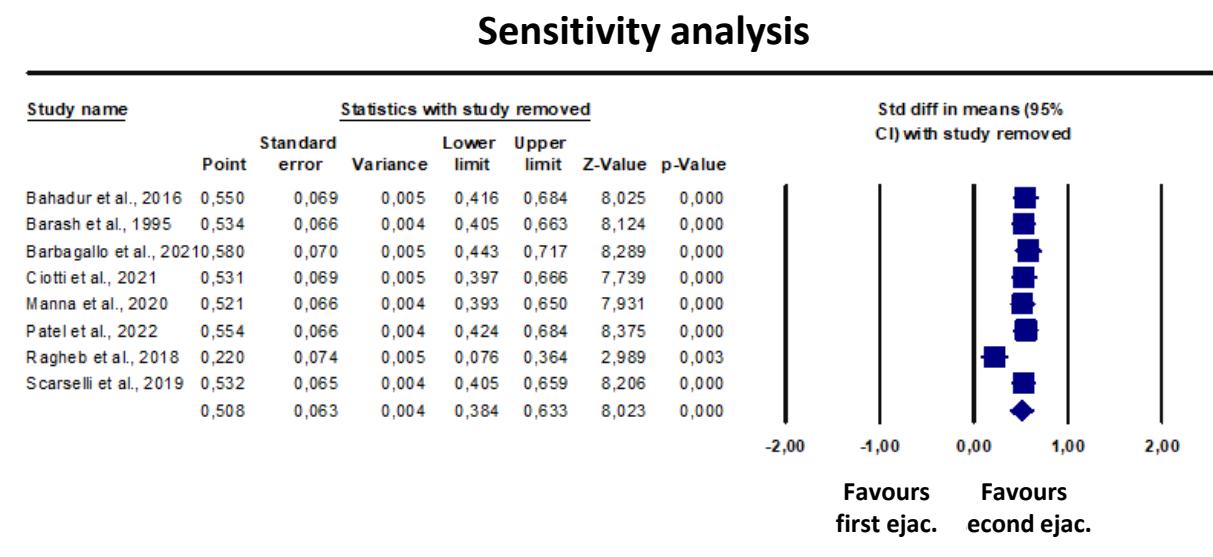


Supplementary Figure S9. Funnel plot (A) and sensitivity analysis (B) of studies that evaluated the effects of a very short abstinence period on sperm morphology in Oligo- and/or Astheno- and/or Terato- zoospermic men

A



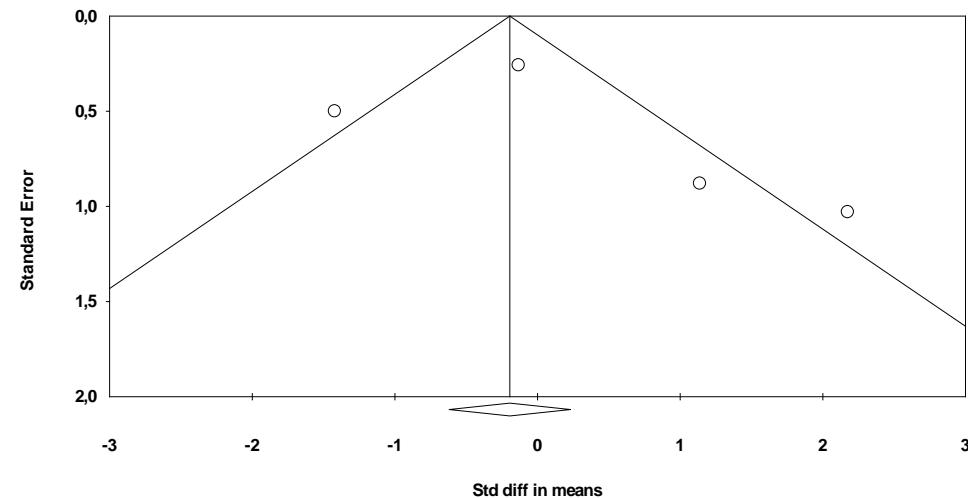
B



Supplementary Figure S10. Funnel plot (A) and sensitivity analysis (B) of studies that evaluated the effects of a very short abstinence period on sperm DNA fragmentation in normozoospermic men

A

Funnel Plot of Standard Error by Std diff in means



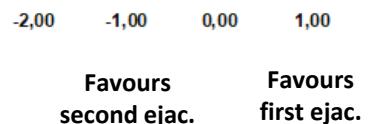
B

Sensitivity analysis

Study name

Study name	Statistics with study removed						Std diff in means (95% CI) with study removed
	Point	Standard error	Variance	Lower limit	Upper limit	Z-Value	
Hussein et al., 2008 0,090	0,241	0,058	-0,382	0,563	0,374	0,709	
Manna et al., 2020 -0,345	0,401	0,161	-1,131	0,440	-0,862	0,389	
Torres et al., 2016 -0,305	0,222	0,049	-0,740	0,131	-1,371	0,170	
Torres et al., 2016 -0,281	0,224	0,050	-0,720	0,158	-1,253	0,210	
	-0,194	0,217	0,047	-0,620	0,231	-0,895	0,371

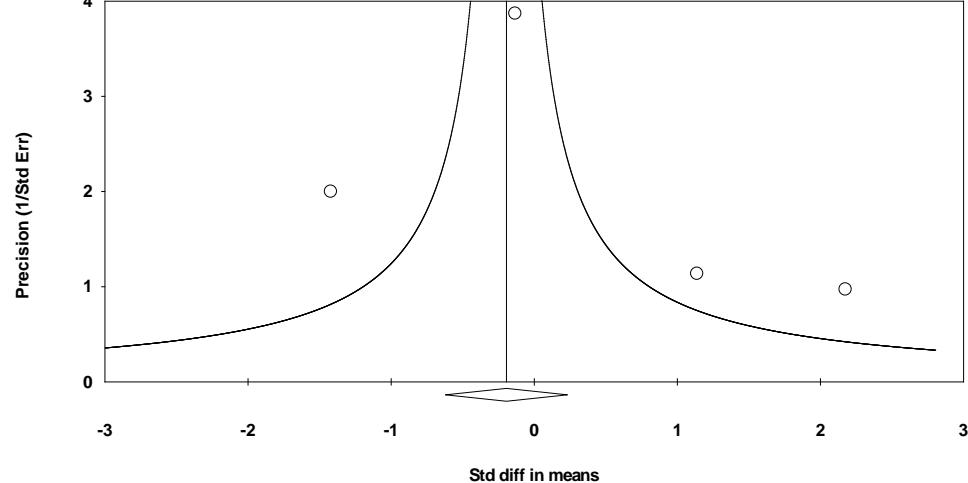
Std diff in means (95% CI) with study removed



Favours second ejac.

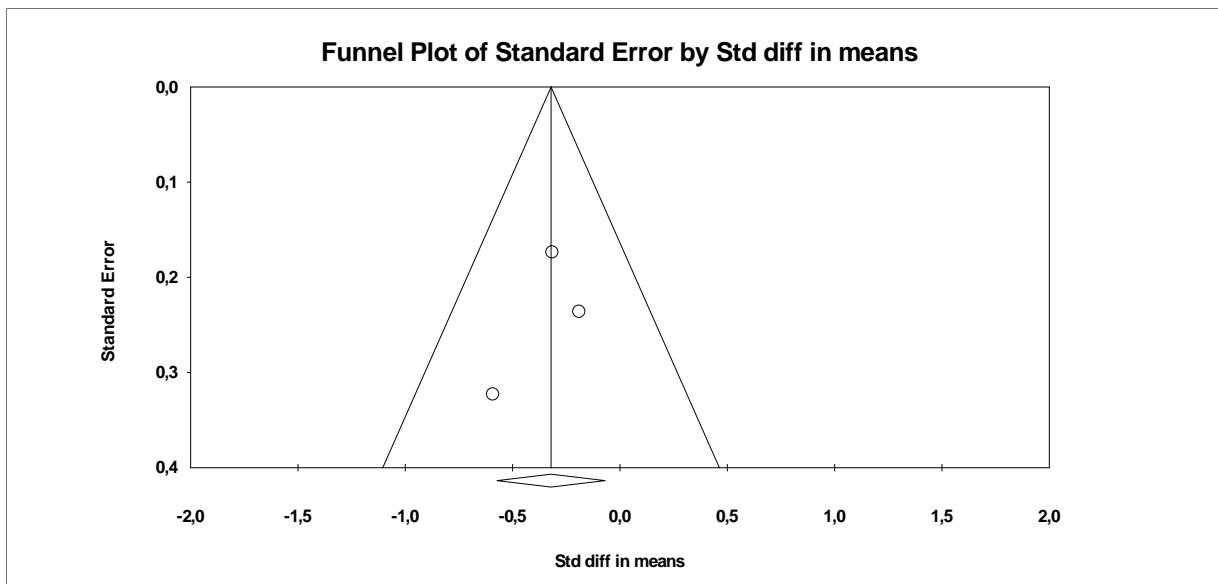
Favours first ejac.

Funnel Plot of Precision by Std diff in means



Supplementary Figure S11. Funnel plot (A) and sensitivity analysis (B) of studies that evaluated the effects of a very short abstinence period on sperm DNA fragmentation in Oligo- and/or Astheno- and/or Terato- zoospermic men

A



B

