Supplementary material

Supplementary Table 1 Preferred Reporting Items for Systematic Review and Meta-Analysis (PRISMA) checklist [26]

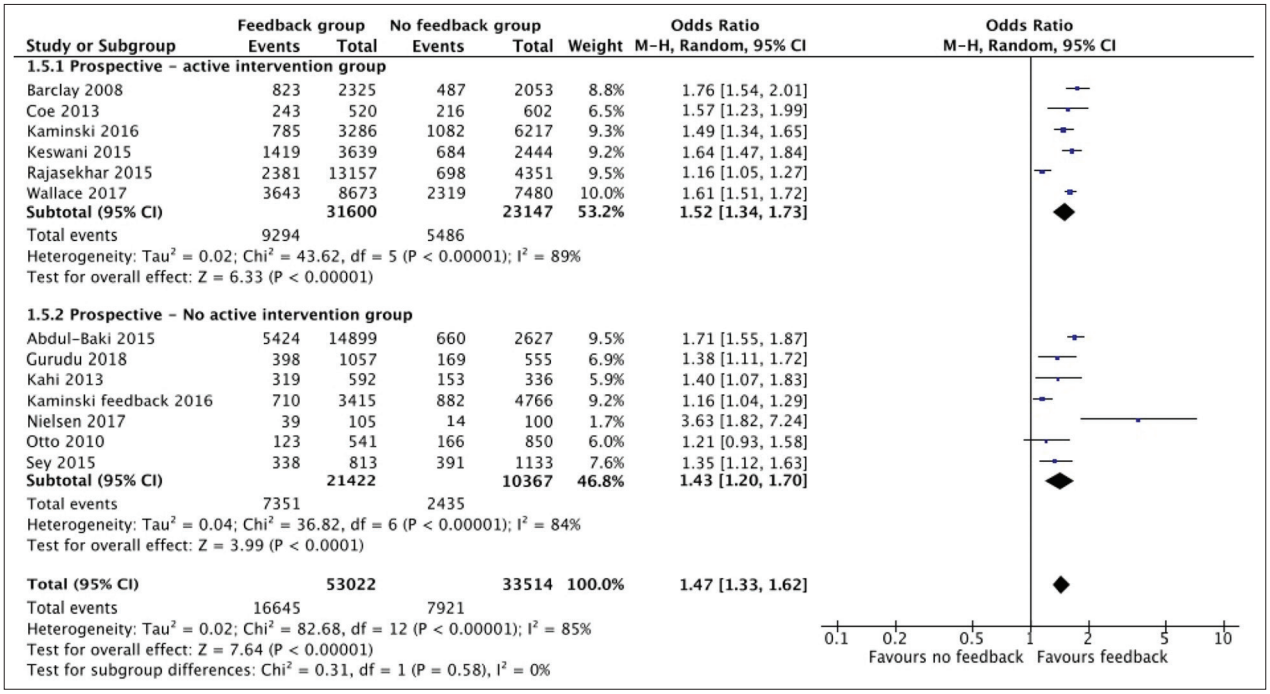
Section/topic	#	Checklist item	Reported on page #
TITLE			
Title	1	Identify the report as a systematic review, meta-analysis, or both	1
ABSTRACT			
Structured summary	2	Provide a structured summary including, as applicable: background; objectives; data sources; study eligibility criteria, participants, and interventions; study appraisal and synthesis methods; results; limitations; conclusions and implications of key findings; systematic review registration number	3-4
INTRODUCTION			
Rationale	3	Describe the rationale for the review in the context of what is already known	5-6
Objectives	4	Provide an explicit statement of questions being addressed with reference to participants, interventions, comparisons, outcomes, and study design (PICOS)	5
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	7
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	7
Search	8	Present full electronic search strategy for at least one database, including any limits used, such that it could be repeated	7
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)	7-8
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	7
Data items	11	List and define all variables for which data were sought (e.g., PICOS, funding sources) and any assumptions and simplifications made	7
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	9
Summary measures	13	State the principal summary measures (e.g., risk ratio, difference in means)	8-9
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	9
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)	9
Additional analyses	16	Describe methods of additional analyses (e.g., sensitivity or subgroup analyses, meta-regression), if done, indicating which were pre-specified	8-9
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	10
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	10

(Contd...)

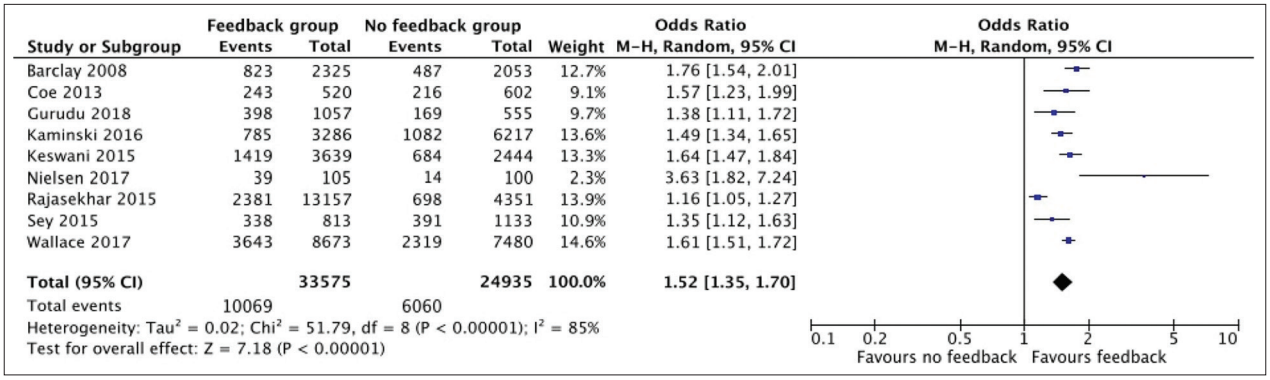
Supplementary Table 1 *(Continued)*

Section/topic	#	Checklist item	Reported on page #
TITLE			
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)	
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 (b) confidence intervals, ideally with a forest plot	22
Synthesis of results	21	Present results of each meta-analysis done, including confidence intervals and measures of consistency	10-13
Risk of bias across studies	22	Present results of any assessment of risk of bias across studies (see Item 15).	14
Additional analysis	23	Give results of additional analyses, if done (e.g., sensitivity or subgroup analyses, meta-regression [see Item 16])	13
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)	14-15
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)	16
Conclusions	26	Provide a general interpretation of the results in the context of other evidence, and implications for future research	17
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	None

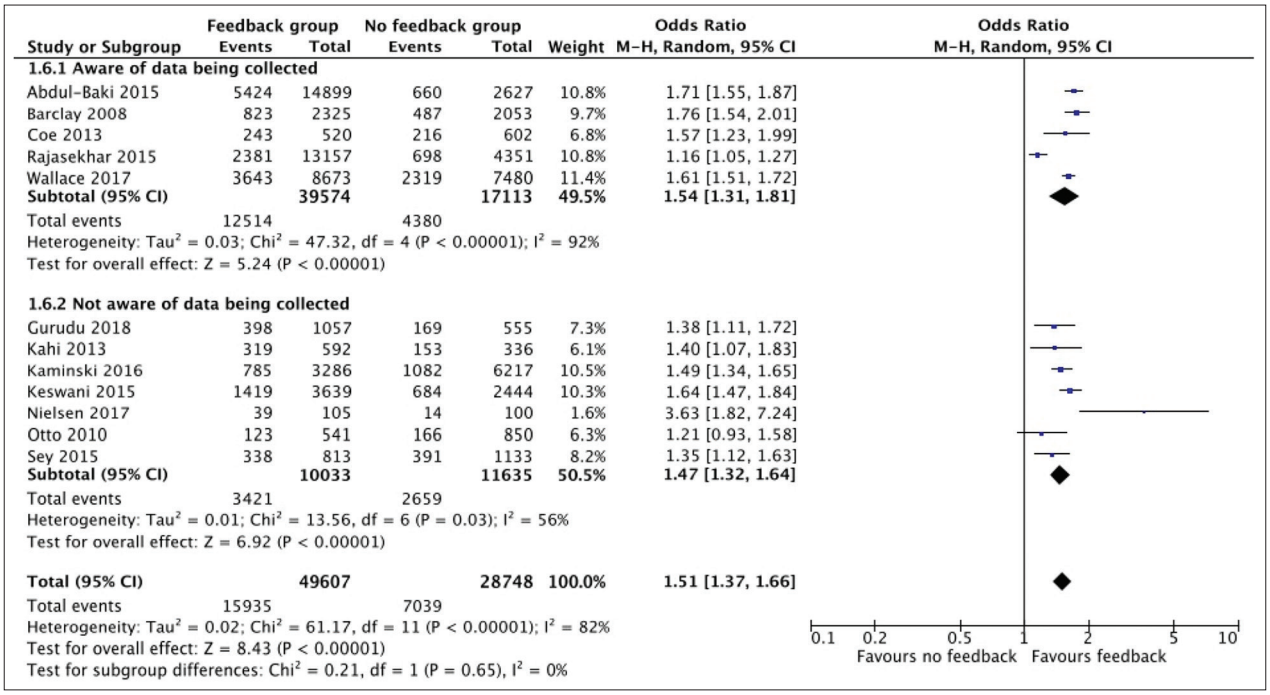
NA, not available



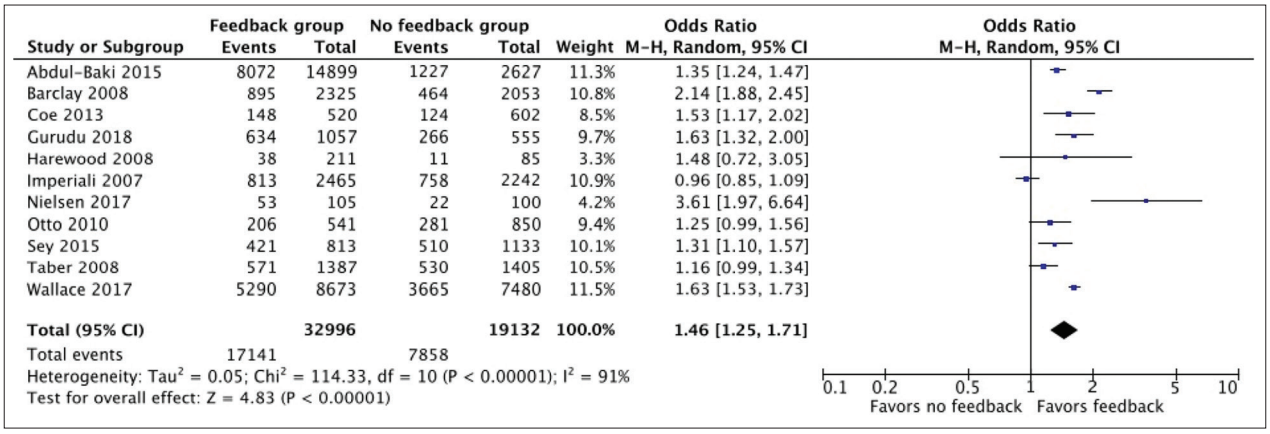
Supplementary Figure 1 Forest plot for pooled analysis of adenoma detection rate based on the type of feedback (active vs. passive)
CI, confidence interval



Supplementary Figure 2 Forest plot analysis for adenoma detection rate in prospective studies
CI, confidence interval



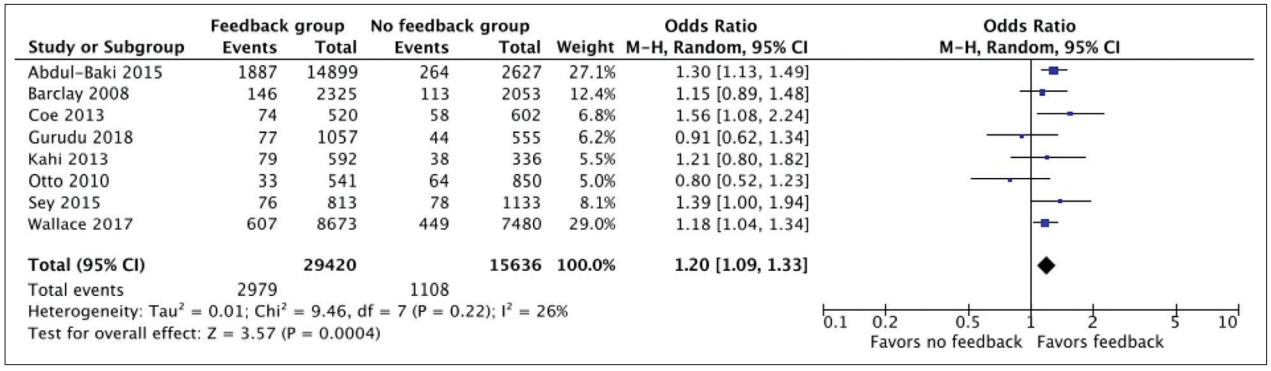
Supplementary Figure 3 Forest plot analyzing the effect of awareness of data being collected prior to feedback on adenoma detection rate
CI, confidence interval



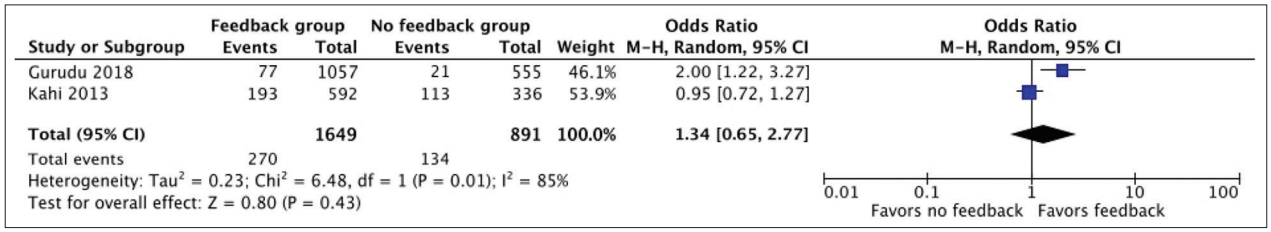
Supplementary Figure 4 Forrest plot for pooled analysis of polyp detection rate
CI, confidence interval

Supplementary Table 2 Quality assessment of included studies based on the Newcastle-Ottawa scale for cohort studies

Criteria	Abdul-Baki *et al* [19]	Gurudu *et al* [16]	Kahi *et al* [20]	Nielsen *et al* [22]	Lin *et al* [23]	*Sey et al* [17]	Coe *et al* [25]	Kaminski *et al* [27]	Barclay *et al* [7]	Taber *et al* [31]	Imperiali *et al* [29]	Coe *et al* [25]	Kaminski *et al* [27]	Wallace *et al* [18]	Harewood *et al* [30]
Selection															
Representativeness of the exposed cohort	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*
Selection of the non-exposed cohort	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*
Ascertainment of exposure	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*
Demonstration that outcome of interest was not present at onset of study	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*
Comparability of cohorts															
Study controls for age/sex?			*							*		*	*	*	*
Study controls for at least 3 additional risk factors?					*		*	*				*	*	*	*
Outcome															
Assessment of outcome	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*
Was follow up long enough for outcomes to occur?	*	*	*	*	*	*	*	*	*	*	*	*	*	*	*
Total score	6	6	7	6	7	6	7	7	6	7	6	8	8	8	8



Supplementary Figure 5 Forrest plot for advanced adenoma detection rate pooled analysis
CI, confidence interval



Supplementary Figure 6 Forrest plot for serrated sessile adenoma detection rate pooled analysis
CI, confidence interval