TABLE 1 | GRADE summary of findings.

TEAS group compared to Control group for the prevention of the incidence of perioperative neurocognitive disorders (PND).

Patient or population: patients undergoing surgical procedures

Settings: Inpatients

Intervention: TEAS

Comparison: Sham intervention

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect	No of Participants	Quality of the	Importance
	Assumed risk	Corresponding risk	(95% CI)	(studies)	evidence	
					(GRADE)	
	Control intervention	TEAS				
PND response rate	198 per 1000	85 per 1000	RR 0.43	999	$\oplus \oplus \ominus \ominus$	CRITICAL
		(63 to 121)	(0.32 to 0.61)	(13 studies)	0000	
					low ^{1,2}	
POD response rate	242 per 1000	94 per 1000	RR 0.39	579		CRITICAL
Follow-up: within 7 days		(62 to 143)	(0.26 to 0.59)	(7 studies)	$\oplus \oplus \ominus \ominus$	
					low ^{1,2}	
CAM scores	-	The mean CAM scores in the TEAS groups was	Not applicable	251		IMPORTANT
		-1.30 lower		(3 studies)	$\oplus \oplus \ominus \ominus$	
Follow-up: within 7 days		(-2.14 to -0.46 lower)			low ^{1,2}	
Dosage of anestheticsremifentanil	-	The mean dosage of remifentanil in the TEAS	Not applicable	180		IMPORTANT
		groups was		(3 studies)	$\oplus \Theta \Theta \Theta$	
		-1.58 standard deviations lower			very low ^{1,2,3}	
		(-2.54 to -0.63 lower)				

Dosage of anestheticspropofol	-	The mean dosage of propofol in the TEAS groups	Not applicable	180	$\oplus \oplus \ominus \ominus$	IMPORTANT
		was		(3 studies)		
		-0.42 standard deviations lower			low ^{1,2}	
		(-1.33 to 0.50 lower)				

*The basis for the assumed risk (e.g. the median control group risk across studies) is provided in footnotes. The corresponding risk (and its 95% confidence interval) is based on the assumed risk in

the comparison group and the relative effect of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

¹ Downgraded one level owing to high risk of performance bias across some included studies.

² Downgraded one level owing to small sample size for detecting publication bias.

³ Downgraded one level owing to substantial heterogeneity (I²>50%).

TABLE 2 | GRADE summary of findings.

TEAS group compared to Control group for the prevention of the incidence of perioperative neurocognitive disorders (PND).

Patient or population: patients undergoing surgical procedures

Settings: Inpatients

Intervention: TEAS

Comparison: Control intervention (no treatment/ sham intervention)

Outcomes	Illustrative comparative risks* (95% CI)		Relative	No of	Quality of the	Importance
	Assumed risk	Corresponding risk	effect	Participants	evidence	
			(95% CI)	(studies)	(GRADE)	
	Control	TEAS				
	intervention					
DNR response rate	180 per 1000	92 per 1000	RR 0.51	569		CRITICAL
		(59 to 140)	(0.33 to 0.78)	(8 studies)	$\oplus \oplus \ominus \ominus$	
Follow-up: within 30 days (postoperative day 7;					low ^{1,2}	
postoperative day 30)						
MMSE scores	-	The mean MMSE scores in TEAS groups was	Not applicable	350	$\oplus \Theta \Theta \Theta$	IMPORTANT
		0 higher		(5 studies)	0000	
Follow-up: postoperative day 7		(-0.46 to 0.46 higher)			very low ^{1,2,3}	
Biochemical indicator(S100β)	-	The mean biochemical indicator(s100 β) in the	Not applicable	324	$\oplus \ominus \ominus \ominus$	IMPORTANT
		TEAS groups was		(4 studies)		
Follow-up: immediate postoperative period		-1.08 standard deviations lower			very low ^{1,2,4}	
		(-1.67 to -0.49 lower)				

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% confidence interval) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

¹ Downgraded one level owing to high risk of performance bias across some included studies.

² Downgraded one level owing to small sample size for detecting publication bias.

³ Downgraded one level for imprecision (CI included the null effect).

⁴ Downgraded one level for inconsistency ($l^2 > 50\%$).