## Supplemental File 1: Diagnostic parameters estimated based on published studies

	Sensitivity 95% CI	Specificity 95% CI	+ predictive value 95% CI	- predictive value 95 CI
RCTs (assuming all trial failures as true negatives)	100%	53% (48%- 57%)	65% (61%-69%)	100%
RCTs (assuming 50% trial failures as true negatives)	77% (73%-81%)	35% (30%- 41%)	65% (61%-69%)	50% (43%-56%)
Observational studies (assuming all trial failures as true negatives)	100%	36% (23%- 49%)	61% (50%-72%)	100%
Observational studies (assuming all trial failures as true negatives	85% (76%-94%)	22% (9%-35%)	61% (50%-72%)	50% (27%-73%)

CI, confidence interval; RCT, randomized controlled trial

## Supplemental File 2: Differences between permanent and temporary lead trials

Factors considered	Permanent lead trial*	Temporary lead trial		
Approach	Cylindrical leads anchored and tunneled during the trial, inserted with the intent to continue for therapy after a successful trial by connecting to an IPG	Cylindrical leads inserted with an intent to discard after the trial, and followed by insertion of another set of leads connected to an IPG for therapy		
Difference in trial success rate	No relative difference	No relative difference		
Time for insertion	Relatively higher	Relatively lower		
Infection risk	Potentially higher	Lower		
Patient discomfort or pain post procedure	Potentially higher	Lower		
Lead migration risk	Lower	Potentially higher		
Conduct of the trial	Requires operating room	Can be considered outside the operating room		
Overall radiation exposure	Lower	Higher (considering additional final phase implant)		
* It does not apply to wireless trials in which there is no separate IPG; IPG: implantable pulse				

<sup>\*</sup> It does not apply to wireless trials in which there is no separate IPG; IPG: implantable pulse generator