



2021

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Highlight: Continuing to drive more transparent reporting in clinical trials



Research and development

- [Evidence that reporting of harms is still sub-optimal](#)
- Finalising the CONSORT Harms update and working with senior CONSORT Executive so that enhanced harms reporting is incorporated into the main checklist



Best practice and guidance

- [Utilization of the evidence from studies with no events in meta-analyses of adverse events](#)
- [Framework to guide evidence synthesis practice for meta-analysis with zero-events studies](#)



Methods implementation

- Provided feedback on Cochrane's new Editorial Manager system
- Available to support Cochrane Reviews