Graphical Summaries of Trial Conduct in DMC Reports
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For many clinical trials, Data Monitoring Committees (DMCs) are charged with monitoring not only the safety and efficacy of an intervention, but also the conduct of the trial itself. A study with low accrual, high dropout, or an unacceptable lag in data collection, adverse event coding or endpoint adjudication may not have information of sufficient quality for monitoring, and may ultimately prove unable to answer the clinical questions of interest.

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