

Preventing potential patient harm through clinical content interventions during oncology clinical trial implementation.

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Ensuring the <u>safety</u> of clinical trial <u>participants</u> is paramount to successful, meaningful clinical research. In this study, researchers examined 585 clinical trial documents and found that 17% included potential patient safety interventions (e.g., resolving medication dosing discrepancies). The authors suggest that clinical specialists' review of study protocol documents could enhance patient safety during clinical trial conduct.