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such virtual clinical trials can leverage digital health technologies for collecting information at each stage of the clinical trial improving trial participant recruitment and retention enabling online based informed consent measuring real time clinical endpoints and continuous tracking of adverse events in this article we outline an oncology specific paradigm for virtual clinical trials identifying the key components for study design and describing the challenges and regulatory considerations for patients providers and policymakers a patient s entire journey in a clinical trial can be made virtual and more inclusive using technology some key benefits include remote screening consent using mobile and web based applications use of electronic patient reported outcome epro diaries for remote recording of patient outcomes through the new nci virtual clinical trials office a centralized team of support staff including research nurses clinical research associates and clinical data specialists will work remotely to assist nci designated comprehensive cancer centers and community practices with their clinical trials activities such virtual clinical trials can leverage digital health technologies for collecting information at each stage of the clinical trial improving trial participant recruitment and retention enabling online based informed consent measuring real time clinical endpoints and continuous tracking of adverse events a virtual clinical trial also known as remote or decentralized clinical trials in the definition refers to digitally empowered clinical trial processes this is a relatively new model of clinical trials in terms of adoption and is therefore the highly underutilized method of conducting clinical research

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