TOP 8 REASONS CLINICAL TRIAL BIOSPECIMENS PUT YOUR STUDY AT RISK

1. INTERIM ANALYSIS PREPARATION

2. UNSCHEDULED MONITORING VISITS

3. CONSENT EXPIRATION RESULTING IN IMPROPER STORAGE OR UTILIZATION

4. OUTDATED CONSENT RESULTING IN IMPROPER USE OF PATIENT SAMPLES

5. PATIENTS DROPPED DUE TO MISHANDLED OR LOST SAMPLES

6. PATIENT ENROLLMENT OR FIRST TREATMENT DELAYED DUE TO MISHANDLED SAMPLES

7. DATABASE LOCK DELAYED DUE TO SAMPLE ISSUES

8. MISSING SITE QUALITY METRICS AND KRI'S ON SAMPLES IN A RISK-BASED APPROACH

See how you can reduce risk of clinical trial delays caused by biospecimen issues and ensure regulatory compliance to patient consent for future-use samples.

BIOFORTIS

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