Image displays RAMS IRB extended smartform fields (pre-toolkit go-live on July 14, 2023) next to the reduced RAMS IRB smartform effective July 14, 2023. The extended fields are initial setup: study identification, federal regulations, IRB panel setup, review setup, and initial setup complete. Background rationale and goals: background, rationale, and goals, study population, background rationale section complete. Research plan: study procedures, project details, bio-medical drug/supplement/other compound details, sample collection details, secondary data/specimen details, costs to participants, compensation, contingency plan, research plan complete. Consent plan: consent process, waiver of some or all elements of consent, consent plan complete. Risk, privacy and confidentiality: risks, discomforts, potential harms and monitoring, privacy, data confidentiality and storage, data retention, sharing plan, pertinent results and incidental findings, risk benefits complete. Populations with special considerations: populations with special considerations, populations complete. Institutional requirements: study funding, types of sites, personnel, conflict of interest, other VCU requirements, HIPAA, partial waiver of authorization, institutional requirements complete. Summary: documents, document complete.