carevive OPT-IN®

Database & Analytics

Carevive OPT-IN® , the Oncology Pragmatic Trial Investigator Network, is a consortium of clinician- investigators collecting disease-specific, real-world patient experience data, including EHR-derived clinical data and longitudinal patient-reported outcomes (PRO). Our quarterly data deliverables is accompanied by our Cancer Patient Experience Report, combining statistical presentations and expert investigator insights.

Research Objectives

1. Collect and apply symptom prevalence data to support continuous oncology practice improvement
2. Develop evidence for clinical benefit and disease burden across evidence-based treatment options
3. Understanding patient-reported symptoms, burden, physical function, and real-world outcomes to improve patient outcomes

Quarterly Deliverables

1. Data License - De-identified datasets aggregated across all Carevive OPT-IN® sites
2. Cancer Patient Experience Reports - Insights from Carevive's Patient Experience Database (by cancer type)
    - Demographics and compliance metrics validating the feasibility of PRO collection as part of routine care
    - Curated insights from clinicians on:
        - The impact of Carevive's platform on improving clinical practice
        - Hot topics from the cancer center
        - Insights on the Patient Experience via Carevive Grand Rounds (patient profiles)

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Data Dictionary: EHR and PRO

Further tailored to each cancer type through close engagement with our life science partners

EHR Fields	Description
Patient ID	Unique Patient Identifier
Age	Age of the patient
Gender	Patient's sex
Race	Description of patient's race and ethnicity
Cancer type	Cancer diagnosis
Prognostic Factors/Biomarkers	Biomarker or prognostic factor name or description
Stage	Description of overall clinical stage (includes T, N, M)
Pathology	Description of overall pathologic Stage (includes T, N, M)
Drugs	Description of the chemotherapy agent(s) (names, dose, route of administration)
Death	Death, death date and cause of death where available (i.e. if contained in the applicable site's EMR records)
Discontinuation	Reasons for treatment discontinuation (tolerability, clinical progression, disease progression, death)

PRO Measure	Description
Weekly Questions	
PROs Derived from NCI PRO-CTCAE:	Anxious
	Nausea
	Vomiting
	Shortness of breath
	Numbness or tingling
	Pain
	Insomnia
	Fatigue
	Sadness
	Problems with concentration
	Muscle Pain
	Mouth or throat sores
	Constipation
	Diarrhea
PROMIS	Physical Functioning
BPI-SF, single item	24 hour pain
GP5, (FACT-G Single Item)	Treatment bother
Patient Reported Healthcare Resource Utilization	Weekly patient report of hospitalizations and ED visits
Health-related Quality of Life	2 questions from EORTC QLQ C30
Baseline Questions	
Frailty Assessment	Proprietary algorithm for Fit, Intermediate Fit or Frail
Social Determinants of Health	7 question battery

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