

ABOUT COTA

Bringing clarity to cancer

Founded in 2011 by doctors, engineers and data scientists to create clarity from fragmented and often-inaccessible real-world data. COTA curates fit-for-purpose, regulatory-grade oncology RWD from structured and unstructured EHR data that is used across the drug development lifecycle.

Our data is ready for analysis and contains clear longitudinal patients journeys that support regulatory submissions and research questions including treatment patterns, real-world outcomes, and comparisons.

2M+ Cancers live in our network

1,500+ Oncologists across our network

3 FDA and EMA approvals

OUR REAL-WORLD DATA PRODUCTS



Fit-for-Purpose/ Custom RWD Cohorts



VANTAGE

Standard-Level
Disease Based RWD

 Volume based on disease prevalence and oncology clinical trial enrollment rates in the U.S.



REGULATORY GRADE

COTA RWD in 10+ regulatory filings, including three successful regulatory approvals



UNIQUE DATA MODELS

Data models for each cancer type provide the most clinically relevant data points for that particular disease



DEEP CLINICOGENOMICS

High quality and quantity of clinical and genomics data across solid and liquid cancers



RIGHT SOURCES

50% Academic Networks 50% Community Networks



Geographic representation of patients accessible through our 50/50 mix of academic and community networks

USE CASES

REGULATORY

External control arm support, comparator cohorts

RESEARCH

RWE studies, research publications, scientific innovation, translational epidemiology, early discovery

COMMERCIAL

Comparative effectiveness, HTA

Standard: Data elements defined by our medical team and key industry opinion leaders. Includes demographics, diagnostics, performance status, molecular markers, labs, treatments and outcomes including progression, response, reason for treatment discontinuation, and survival

ONGOING RESEARCH PARTNERSHIPS WITH:

UTSouthwestern Medical Center









LEARN MORE AT: www.cotahealthcare.com