# Converting Optum EHR Oncology module into OMOP CDM

ETL logic and concepts mapping overview

#### **INTRO**

- The Optum® Enriched Oncology Data set is a group of tables that can supplement the Optum® de-identified Electronic Health Record dataset.
- It includes specific oncology concepts important for understanding the progression of the disease, which often not available in structured formats, particularly the tumor, node, and metastasis (TNM) values, stage information and biomarkers.
- It is obtained from patient records using NLP methods
- As of 2022, there are approximately 1.9 million patients with at least one solid tumor ICD-9 or ICD-10 diagnosis included in the data set.

### Overall logic

source entry	target domain	target vocabulary
Histology	Condition	SNOMED, ICDO3
Topography	Condition	SNOMED, ICDO3
Laterality	Condition	SNOMED, ICDO3
Behavior (in situ, malignant or benign)	Condition	SNOMED, ICDO3
summary stage	Measurement	Cancer Modifier
metastasis location	Measurement	Cancer Modifier
TNM	Measurement	Cancer Modifier
tumor grade	Measurement	Cancer Modifier
characteristics: advanced, carcinomatosis,		
extensive, infiltrative, invasive, localized, etc	Measurement	Cancer Modifier
		OMOP genomic, LOINC,
Biomarkers	Measurement	SNOMED
evaluation system: Binet Stage,		
Durie/Salmon Stage, ECOG performance		
status, FIGO Stage, Gleason, Gleason score	Measurement	Cancer Modifier
Tumor size	Measurement	Cancer Modifier
treatment regimen	Episode*	HemOnc
Tumor progression	Episode*	Episode
		no mapping - not
		supported by the
treatment response	Observation	vocabulary model

Precoordinated into a single concept

Expression level of immunostaining ("0", "1+", "2+", "3+", 100%, 90%) mapped to 'positive', 'negative', 'equivocal'

Calculated the largest size and mapped to "Largest Dimension of Tumor", others to "Dimension of Tumor"

<sup>\*</sup>will be mapped in the next data refresh

### Tumor progression to Episode

Source data example of a single patient

ABC ptid TI	② note_date	neoplasm_histology_key 🏋 🛊	ABC progression	
	2009-03-24	f868df0ffdc7eeb894f8fca631ebee4b	no recurrence	Pomission Enisodo
	2009-06-16	20929d974a86b70fa3cfcd32169515ac	no recurrence	Remission Episode
	2009-11-06	65a4a22e812708c345fbe5e5c1da6052	no recurrence	2009-03-24 – 2010-09-08
	2010-09-08	575b334a1a30c8ca8373a62d101eb070	recurrence	
	2010-10-01	11d933a33cc0ab888b330b02a39152df	recurrence	Disease Recurrence Episode
	2010-10-10	21c33db9ac0782cb9e0c7a80e64fcae0	recurrence	<u>Disease Hecuiterice</u> Episode
	2010-11-02	62f169f0562235ba9b52bb9a8d877d65	recurrence	
	2010-12-01	17ccdd44a153ad5aa93a0b547c15db51	recurrence	
	2010-12-06	e7b260e330ab00d59513e0ad47f81633	tumor progression	
	2010-12-20	3c5e44007a599c695ad002e6f1b867e6	recurrence	Progression Episode
	2010-12-20	5f0f56a3b15366ea58163aa35001a28f	tumor progression	<u> </u>
	2010-12-20	0dc77a5e3402488e1119e4c7453b7b92	recurrence	
	2010-12-21	38ef251dfa3d2930cf7b27ed1a10de3a	recurrence	
	2010-12-22	0561614da91110eebb1ad90e5688d7b3	recurrence	Disease Recurrence Episode
	2010-12-23	2de13c0d9b0d0b29fe7515095091848e	recurrence	<u> Discase Neculterice</u> Lpisode
	2010-12-24	28753c699e4b5f51126bba1b5baff77c	recurrence	
	2010-12-24	2613b3feffa773cc0e530ab06e8d3a9f	recurrence	
	2010 12 24	400 CELO O 100E0400710007 100E0		

#### Data elements that can't be mapped. Treatment response

#### **Treatment response terms:**

- good therapeutic response
- excellent therapeutic response
- complete therapeutic response
- partial therapeutic response
- complete pathologic therapeutic response
- very good partial response
- minimal residual disease response
- good clinical therapeutic response
- excellent clinical therapeutic response
- fair therapeutic response

#### **Example of the data**

Treatment response in different patients

ABC ptid \\\ އ	② note_date	ABC treatment	RBC treatment_response
	2020-07-30	[NULL]	good therapeutic response
	2022-10-20	[NULL]	good therapeutic response
	2021-02-22	[NULL]	good therapeutic response
	2019-05-13	[NULL]	excellent clinical therapeutic response
	2016-01-28	neoadjuvant chemotherapy	good therapeutic response
	2018-05-23	[NULL]	partial therapeutic response
	2019-11-21	chemotherapy	excellent therapeutic response
	2020-01-31	neoadjuvant chemotherapy	good clinical therapeutic response
	2018-04-09	[NULL]	partial therapeutic response
	2018-05-03	[NULL]	excellent therapeutic response

### Data cleansing

#### Data entries were removed where exist:

- "in situ" and "invasive" at the same day.
- inconsistent numeric and narrative biomarkers results, for example numeric result = "+1", narrative result = "positive mutation" in ERBB2/HER2 measurement.\*
- *More such rules to be applied:* 
  - E.g. Positive and negative biomarker status in the same patient
- Event tables were deduped if at the same date there was the same information
  - condition\_source\_value in Conditions,
  - combination of measurement\_source\_value, value\_as\_number, value\_source\_value in Measurement.

<sup>\*</sup>A score of "1+" suggests that there is a low level of HER2 protein present in the cells. This low level is considered within the normal range, and so the cancer is unlikely to respond to therapies that target HER2. Therefore, a "1+" score is usually interpreted as a negative result for HER2 overexpression.

#### Concept mapping

source_Name	target_concept_name
	ERBB2 (erb-b2 receptor tyrosine kinase 2) gene variant
erb-b2 receptor tyrosine kinase 2 (ERBB2 or HER2/neu)	measurement
estrogen receptor/progesterone receptor (ER/PGR)	ESR1 Protein Expression measurement
estrogen receptor/progesterone receptor (ER/PGR)	PGR (progesterone receptor) gene variant measurement
	MKI67 (marker of proliferation Ki-67) gene variant
marker of proliferation Ki-67 (MKI67 or Ki-67)	measurement
CD274 molecule (CD274 or PD-L1 or PDL1)	CD274 (CD274 molecule) gene variant measurement
adenocarcinoma	Malignant adenomatous neoplasm
carcinoma	Malignant epithelial neoplasm
squamous cell carcinoma	Squamous cell carcinoma
	Neoplasm defined only by histology: Basal cell carcinoma,
basal cell carcinoma	NOS
in situ ductal carcinoma	Intraductal carcinoma in situ of breast
lung non-small cell carcinoma	Non-small cell lung cancer
multiple myeloma	Multiple myeloma
malignant mammary neoplasm	Malignant tumor of breast
prostatic adenocarcinoma	Adenocarcinoma of prostate
lung adenocarcinoma	Adenocarcinoma of lung

**Biomarkers** were mapped mostly to the OMOP Genomic vocabulary, Generic Variation concept class.

77% distinct concepts are mapped to OMOP Genomic, 19% to SNOMED or LOINC,

4% are not mapped, but those have low frequency.

Conditions are mapped well with histology information included, but sometimes it's only histology (in yellow), so you need to define the topography and histology separately when phenotyping.

## Cancer characteristics that can't be mapped

source term	comment
locally advanced	
not metastatic	in theory can be mapped to metastasis+absent, but I afraid people will not use it, + our tools such as CD, doesn't look at values. Is there a concept for 'non-metastatic' – localized or something?
not invasive	
not in situ	
advanced	
localized	
carcinomatosis	there's such Condition, should be measurement
not malignant	
oligometastatic	
multicentric	
extensive	

#### Data evaluation

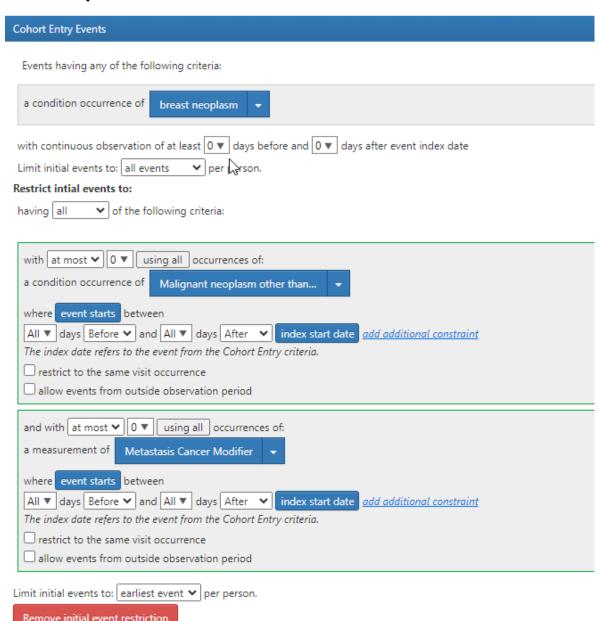
- 1) Conditions and measurements connected grouped
- 2) Create a Cancer cohort and evaluate the distribution of cancer modifiers
- 3) Look for impossible combination of events

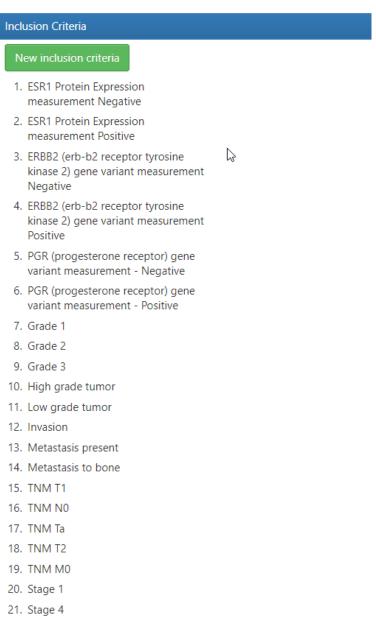
## Top 40 condition-measurement combinations defined using MEASUREMENT modifiers

No topography

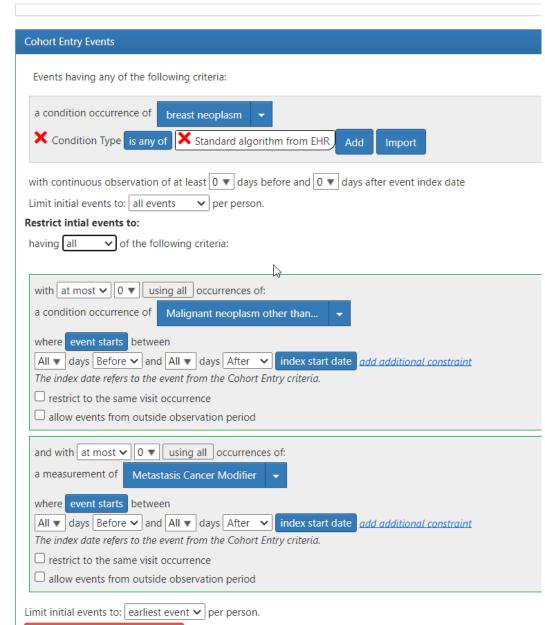
ABC condition_name	123 measuren ∏‡	ABC measurement_name	123 vi 📆 🛊	ABC value_as_concept_ \(\textit{\gamma}\)
Primary malignant neoplasm of breast	[NULL]	[NULL]	[NULL]	[NULL]
Neoplasm of skin	[NULL]	[NULL]	[NULL]	[NULL]
Neoplasm defined only by histology: Basal cell carcinoma, NOS	[NULL]	[NULL]	[NULL]	[NULL]
Primary malignant neoplasm of breast	36,769,449 🗹	Invasion	0	No matching concept
5 Malignant epithelial neoplasm	36,769,449 🗹	Invasion	0	No matching concept
Malignant epithelial neoplasm	36,769,180 🗹	Metastasis	4,181,412	Present
Primary mailignant neoplasm of prostate	[NULL]	[NULL]	[NULL]	[NULL]
Primary malignant neoplasm of breast	35,976,980 🗹	ESR1 Protein Expression measurement	5,884,084	Positive
9 Malignant epithelial neoplasm	1,633,440 🗹	AJCC/UICC N0 Category	[NULL]	[NULL]
10 Malignant adenomatous neoplasm	36,769,180 🗹	Metastasis	4,181,412	Present
11 Primary malignant neoplasm of breast	35,955,862 🗹	ERBB2 (erb-b2 receptor tyrosine kinase 2) gene variant measurement	5,878,583	Negative
12 Squamous cell carcinoma	[NULL]	[NULL]	[NULL]	[NULL]
Primary malignant neoplasm of breast	1,633,440 🗹	AJCC/UICC N0 Category	[NULL]	[NULL]
Primary malignant neoplasm of breast	35,957,667 🗹	PGR (progesterone receptor) gene variant measurement	5,884,084	Positive
15 Neoplasm of colon	[NULL]	[NULL]	[NULL]	[NULL]
16 Malignant epithelial neoplasm	1,635,624 🗹	AJCC/UICC M0 Category	[NULL]	[NULL]
17 Neoplasm of lung	36,769,180 🗹	Metastasis	4,181,412	Present
18 Primary malignant neoplasm of breast	1,635,624 🗹	AJCC/UICC M0 Category	[NULL]	[NULL]
19 Primary malignant neoplasm of prostate	4,272,032 🗹	Prostate specific antigen measurement	1,620,380	Elevated
20 Malignant melanoma	[NULL]	[NULL]	[NULL]	[NULL]
21 Primary malignant neoplasm of breast	36,769,180 🗹	Metastasis	4,181,412	Present
Neoplasm of lung	[NULL]	[NULL]	[NULL]	[NULL]
23 Carcinoma of breast	36,769,449 🗹	Invasion	0	No matching concept
24 Malignant adenomatous neoplasm	36,769,449 🗹	Invasion	0	No matching concept
25 Malignant adenomatous neoplasm	1,633,440 🗹	AJCC/UICC N0 Category	[NULL]	[NULL]
26 Malignant epithelial neoplasm	0 ☑	No matching concept	0	No matching concept
27 Malignant tumor of breast	35,976,980 🗹	ESR1 Protein Expression measurement	5,884,084	Positive
28 Malignant epithelial neoplasm	35,976,980 🗹	ESR1 Protein Expression measurement	5,884,084	Positive
29 Malignant epithelial neoplasm	1,634,752 🗹	Grade 2 tumor	[NULL]	[NULL]
30 Malignant tumor of breast	36,769,449 🗹	Invasion	0	No matching concept
31 Carcinoma of breast	35,976,980 🗹	ESR1 Protein Expression measurement	5,884,084	Positive
Malignant epithelial neoplasm	1,633,749 🗹	Grade 3 tumor	[NULL]	[NULL]
Malignant epithelial neoplasm	35,955,862 🗹	ERBB2 (erb-b2 receptor tyrosine kinase 2) gene variant measurement	5,878,583	Negative
Malignant adenomatous neoplasm	1,633,987 🗹	Stage 4	[NULL]	[NULL]
Malignant epithelial neoplasm	1,633,987 🗹	Stage 4	[NULL]	[NULL]
Primary malignant neoplasm of prostate	1,633,643 🗹	Gleason Primary Pattern Grade 3		[NULL]
Intraductal carcinoma in situ of breast	35,976,980 🗹	ESR1 Protein Expression measurement	5,884,084	Positive
Basal cell carcinoma of skin	[NULL]	[NULL]	[NULL]	[NULL]
Primary malignant neoplasm of breast	1,635,838 🗹	Stage 1	[NULL]	[NULL]
40 Malignant epithelial neoplasm	[NULL]	[NULL]	[NULL]	[NULL]

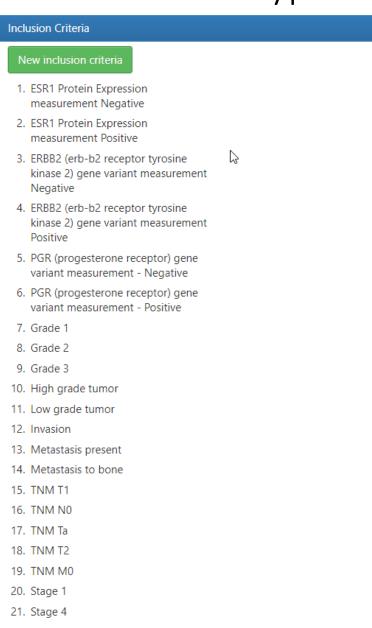
## Cohort definition: Neoplasm of breast excluding other neoplasms, cancer modifiers as inclusion criteria





## Cohort definition: Neoplasm of breast excluding other neoplasms, cancer modifiers as inclusion criteria. Specific condition type





#### Patients with cancer modifiers

Inclu	clusion Report for <b>Optum EHR</b> + <b>Enrich Oncology (v2577)</b> using 1 event per person						Conditions from			
		Match Rate	Matches	Total Events		All		Onco module as		
	Summary Statistics:	0.00%	0	841,688	р	opulation	l į	ndex event		
	Inclusion Rule				N	% Satisfied	N	% Satisfied		
1.	ESR1 Protein Expression measureme	nt Negative			9,400	1.12%	8,115	7.06%		
2.	ESR1 Protein Expression measureme	nt Positive			33,247	3.95%	30,211	26.29%		
3.	ERBB2 (erb-b2 receptor tyrosine kina	se 2) gene variant	measurement Ne	egative	23,505	2.79%	20,834	18.13%		
4.	ERBB2 (erb-b2 receptor tyrosine kina	se 2) gene variant	measurement Po	ositive	5,443	0.65%	4,599	4.00%		
5.	PGR (progesterone receptor) gene va	ariant measuremer	nt - Negative		10,675	1.27%	9,259	8.06%		
6.	PGR (progesterone receptor) gene va	ariant measuremer	nt - Positive		23,523	2.79%	21,344	18.57%		
7.	Grade 1				4,930	0.59%	4,502	3.92%		
8.	Grade 2				9,775	1.16%	8,788	7.65%		
9.	Grade 3				6,642	0.79%	5,678	4.94%		
10.	High grade tumor				4,314	0.51%	3,893	3.39%		
11.	Low grade tumor				2,275	0.27%	2,107	1.83%		
12.	Invasion				32,764	3.89%	29,158	25.38%		
13.	Metastasis present		5,308	0.63%	0	0.00%				
14.	Metastasis to bone				1,848	0.22%	0	0.00%		
15.	TNM T1				19,602	2.33%	17,394	15.14%		
16.	TNM N0				22,074	2.62%	20,201	17.58%		
17.	TNM Ta				621	0.07%	498	0.43%		
18.	TNM T2				8,799	1.05%	7,373	6.42%		
19.	TNM M0				17,208	2.04%	15,440	13.44%		
20.	Stage 1				18,923	2.25%	16,827	14.64%		
21.	Stage 4				3,411	0.41%	1,947	1.69%		

#### Compare with the article results

ncbi.nlm.nih.gov/pmc/articles/PMC7869562/#S1

<u>Journal List</u> > <u>J Clin Med Res</u> > <u>v.13(1); 2021 Jan</u> > PMC7869562

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PMCID: PMC7869562 PMID: <u>33613796</u>

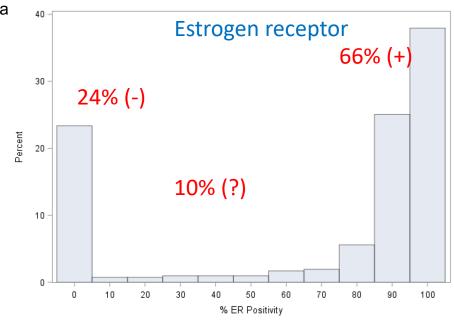
Percentage of Hormone Receptor Positivity in Breast Cancer Provides Prognostic Value: A Single-Institute Study

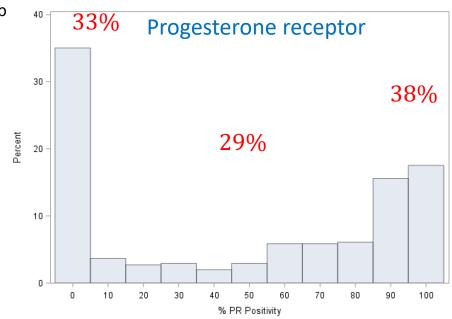
Richard Sleightholm, a,c Beth K. Neilsen, a,c Safwan Elkhatib, Laura Flores, Saihari Dukkipati, Runze Zhao, Songita Choudhury, Bret Gardner, Joey Carmichael, Lynette Smith, Nathan Bennion, Andrew Wahl, and Michael Baine, Michael Baine, Michael Baine, Andrew Wahl, Michael Baine, Michael Baine, Andrew Wahl, Michael Baine, Michael Baine,

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#### in our data: 22%(-) / 78(+)





in our data: 31% (-) / 69(+)

### Future development

- Tumor progression will be mapped to the Episode table in the next iteration
- Line of therapy to be mapped to the HemOnc vocabulary with subsequent run and check of the ARTEMIS
- Data cleansing algorithms to be improved

#### Discussion

- Use cases we can participate in a network study
- Data cleansing approaches
- Data validation algorithms
- Not mapped data elements