



Improving Molecular-Based Clinical Trial Search

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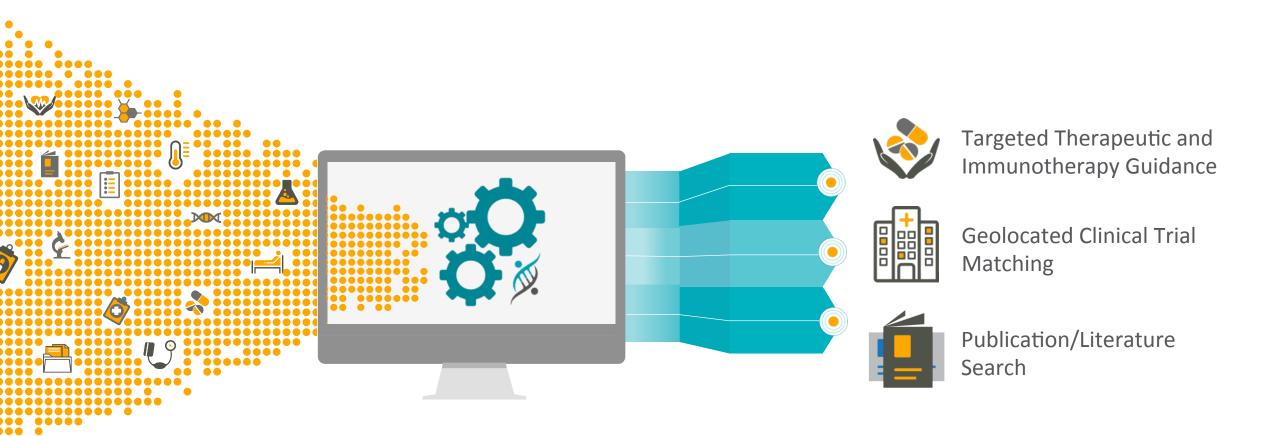
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Molecular Match Products



Trial Matching Pain Points

Problem #1 – Users' search terms can match anywhere in the trial record, leading to lists of trials where search terms are found in the exclusion criteria.

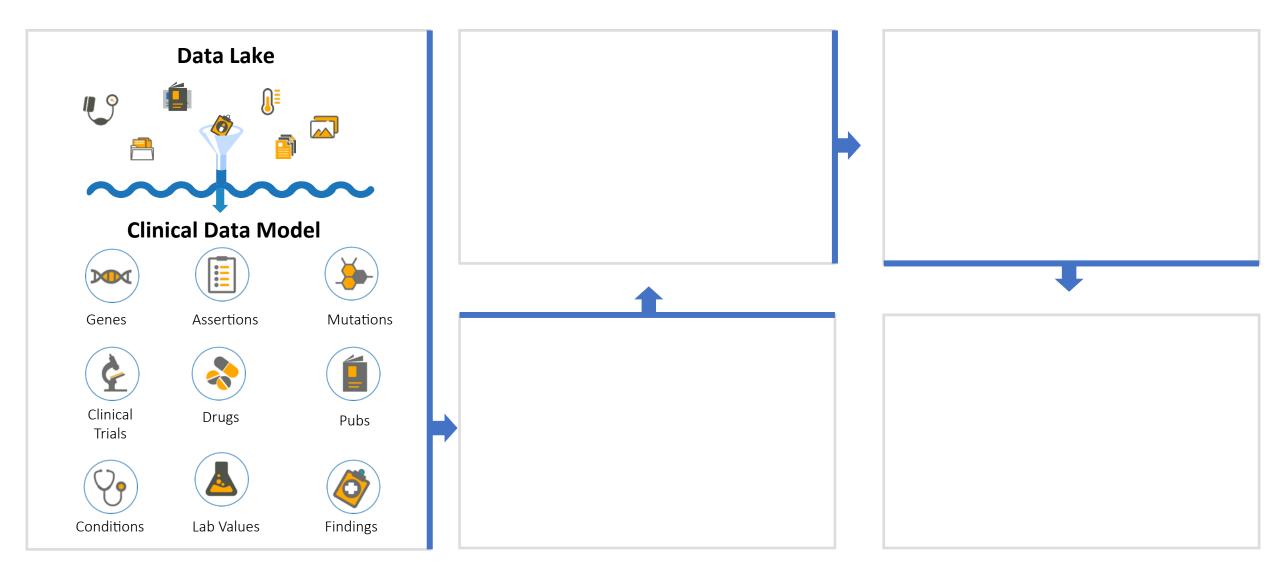
How do we efficiently prioritize relevant trials and eliminate long lists of irrelevant trials?

Problem #2 – Many trials do not specify molecular biomarkers as inclusion criteria, even when targeted drugs are part of the trial intervention.

How do we improve matching of molecular patient profiles to these non-molecular trials?

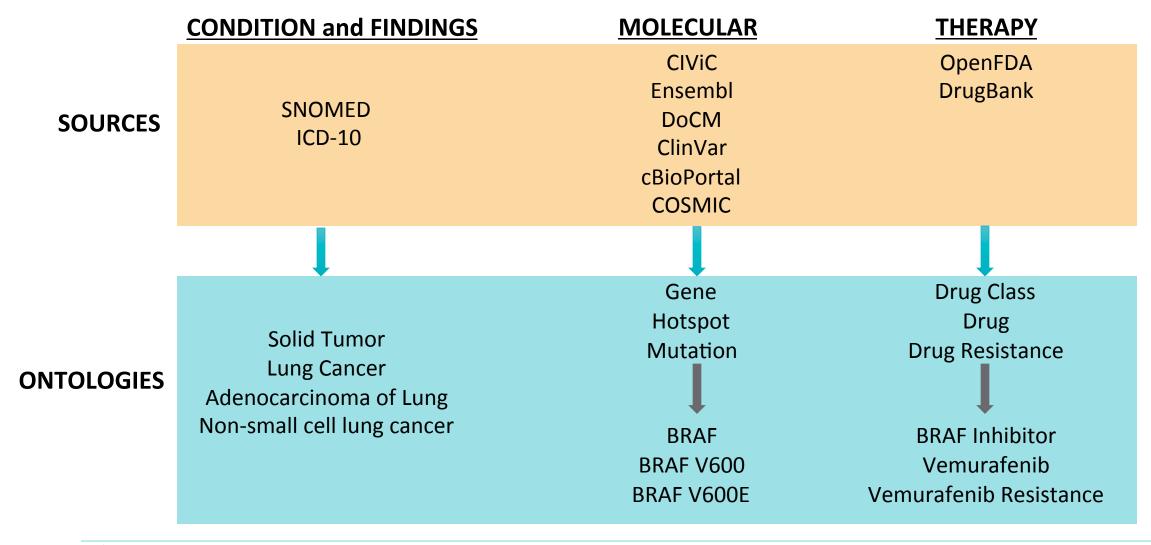


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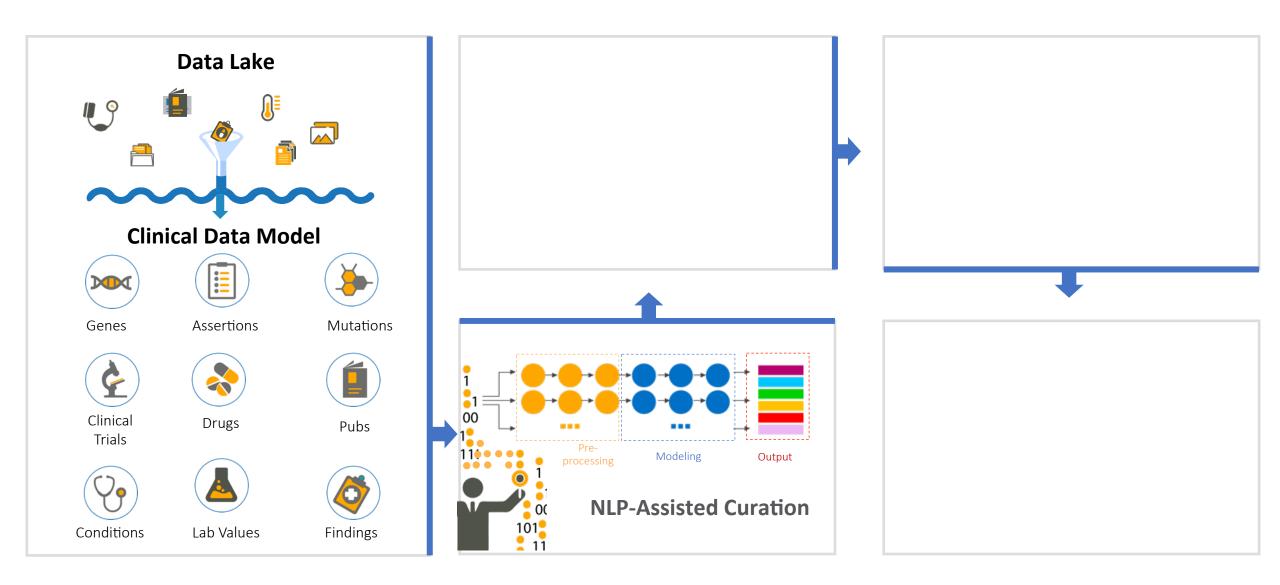


Data Sources and Ontologies Data relationships





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Distinguishing Eligibility Criteria

Search: NSCLC EGFR T790M NIH U.S. National Library of Medicine Find Studies ▼ About Studies ▼ Submit Studies ▼ Resources ▼ About Site ▼ Clinical Trials.gov Search Results > Study Record Detail Save this student
 Save this student Ningetinib (CT053PTSA) Plus Gefitinib in Stage IIIB or IV NSCLC Patients With EGFR Mutation and T790M Negative ClinicalTrials.gov Identifier: NCT03758287 The safety and scientific validity of this study is the responsibility of the Recruitment Status 1 : Recruiting study sponsor and investigators. Listing a study does not mean it has First Posted 1: November 29, 2018 ▲ been evaluated by the U.S. Federal Government. Know the risks and Last Update Posted 1 : December 3, 2018 potential benefits of clinical studies and talk to your health care **See Contacts and Locations** provider before participating. Read our disclaimer for details.

Sponsor:

Sunshine Lake Pharma Co., Ltd.

Information provided by (Responsible Party):

Sunshine Lake Pharma Co., Ltd.



Distinguishing Eligibility Criteria

Criteria

Inclusion Criteria:

- Histologically or cytologically confirmed Stage IIIB or IV NSCLC
- Acquired resistance to EGFR TKI (1st, 2nd or 3rd generation)
- · Histological or cytological evidence of EGFR mutation and T790M negative after progression on last EGFR TKI therapy
- Measurable disease according to Response Evaluation Criteria in Solid Tumors version 1.1 (RECIST 1.1)
- Toxicity recovered to NCI CTCAE v.4.03 Grade ≤1 from previous treatments (except alopecia)
- ECOG performance status (PS) 0 or 1
- Life expectancy of ≥ 12 weeks
- Adequate organ function

Exclusion Criteria:

- · Prior treatments
 - o Chemotherapy, targeted therapy (except EGFR TKI), immunotherapy, radiotherapy, or major surgery within 4 weeks prior to study treatment
 - Nitrosourea and mitomycin chemotherapy within 6 weeks prior to study treatment
 - EGFR TKI treatment within 2 weeks prior to study treatment
 - Had received live vaccine within 4 weeks prior to study treatment
 - Had received any investigational agent from other clinical study within 4 weeks prior to study treatment or are currently participating in other clinical trials
 - Previous treatment with any other c-MET inhibitor or Axl inhibitor (eq, crizotinib, cabozantinib, volitinib, INC280)
- · Symptomatic, untreated or unstable central nervous system metastases

Include:

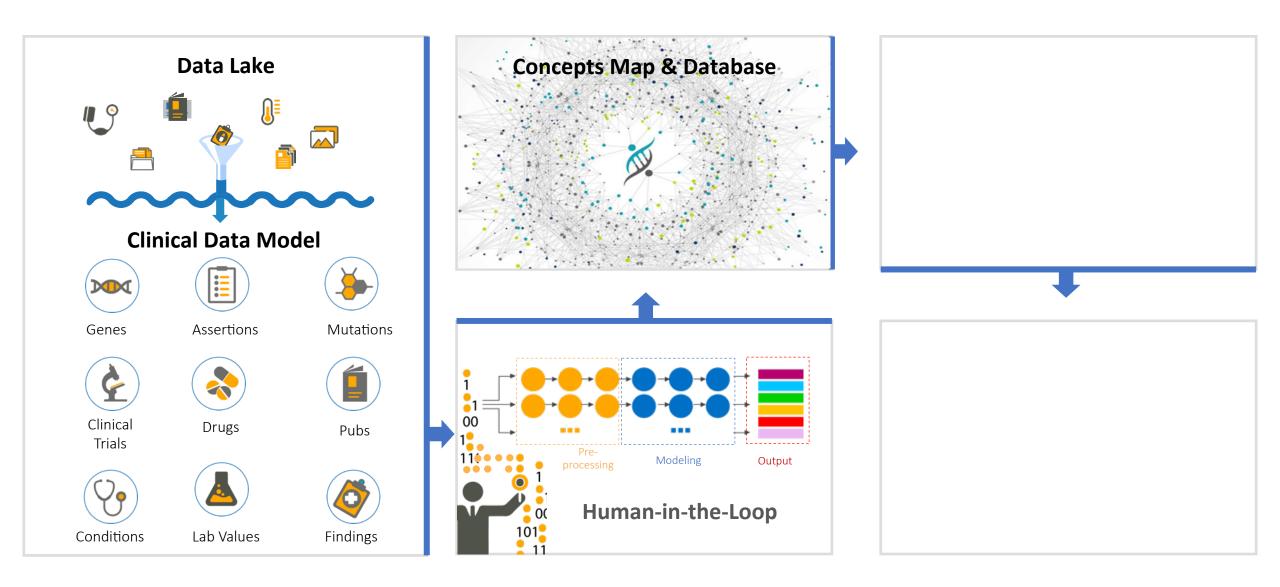
Erlotinib resistance Afatinib resistance Gefitinib resistance Etc.

Exclude:

Crizotinib resistance
Cabozantinib resistance
Volitinib resistance
Etc.



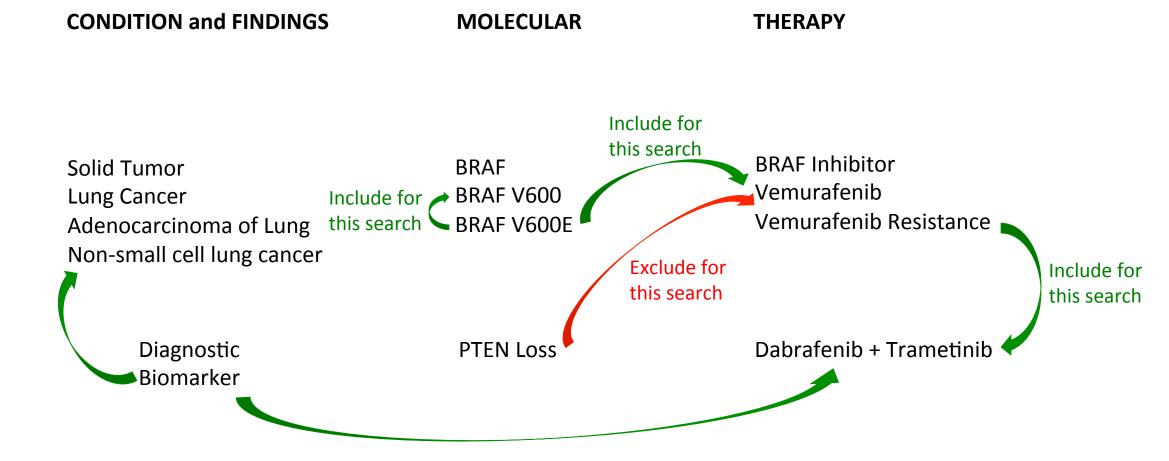
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Concept Association and Inference

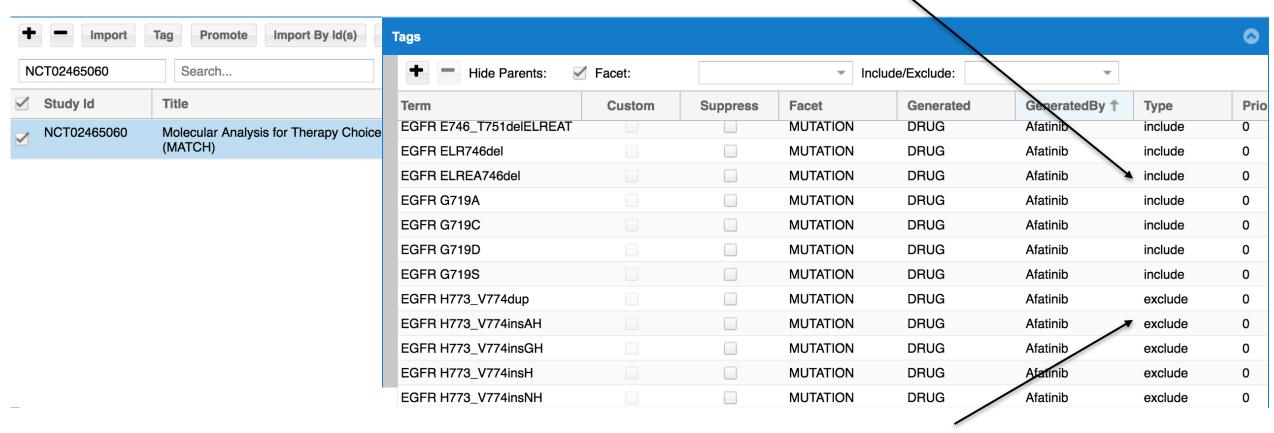
Connecting the data



Concept Association and Inference

Molecular association with drug sensitivity

Include variants that confer sensitivity to drug



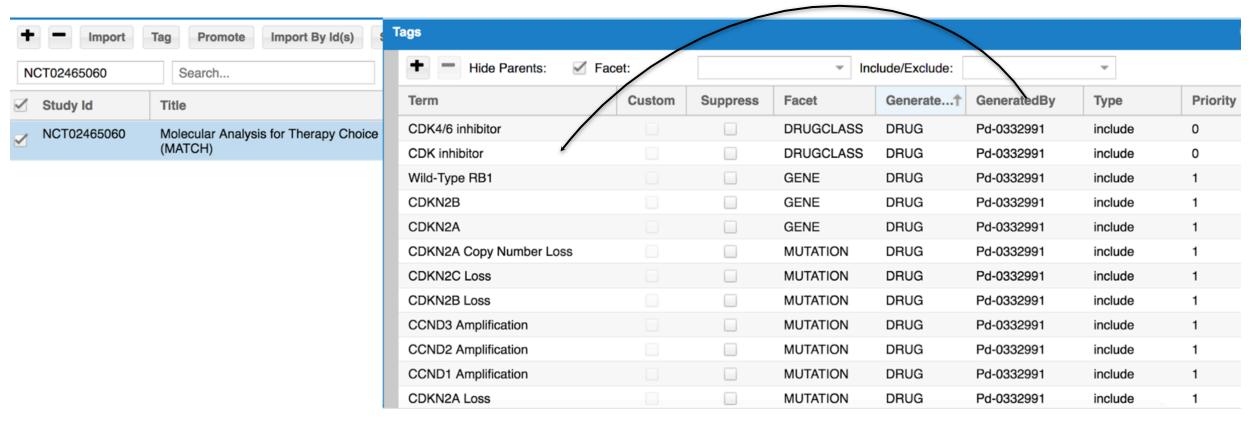
Exclude variants that confer resistance to drug



Concept Association and Inference

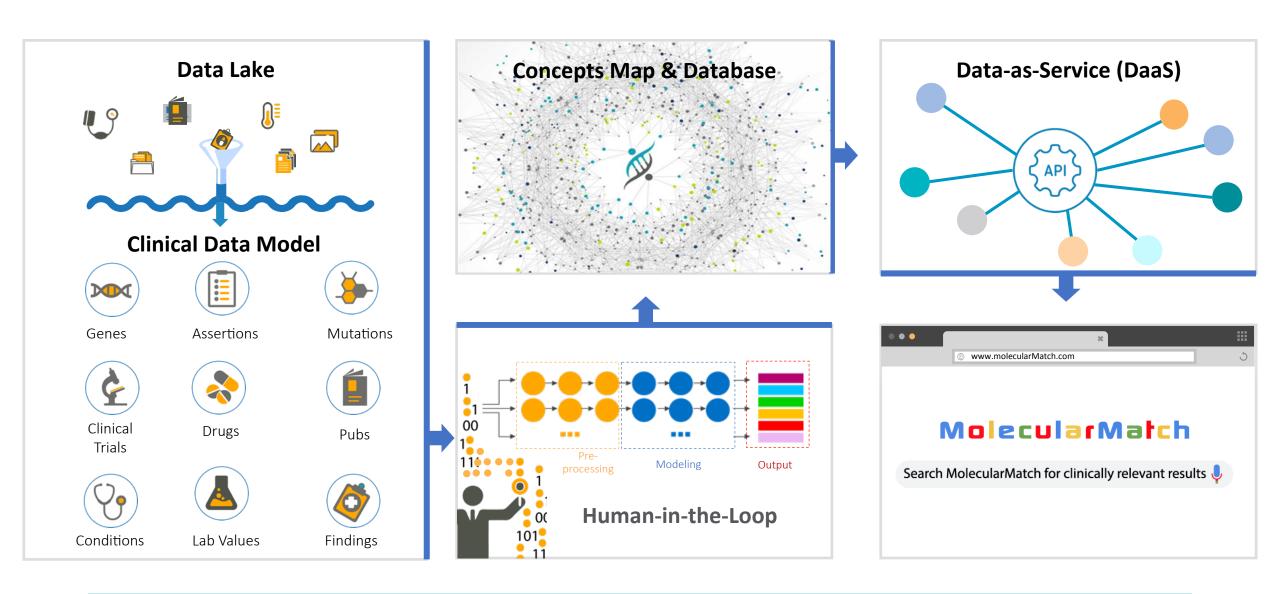
Expansion of relevant pathway mutations to therapy

Palbociclib infers genes and variants within CDK4/6 pathway



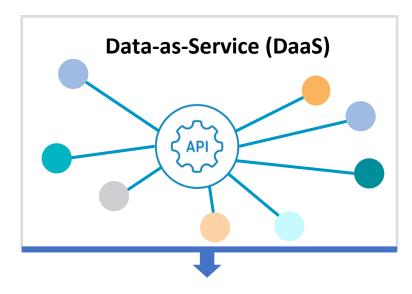


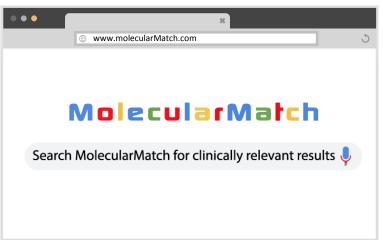
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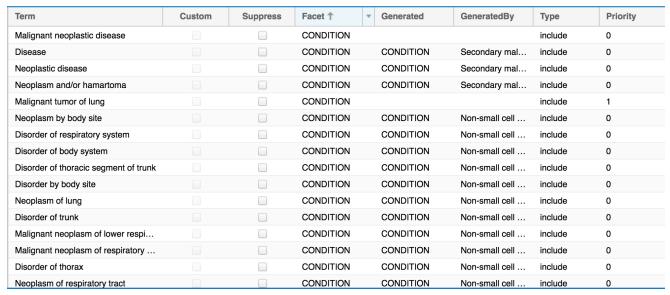


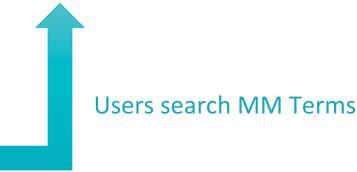


MolecularMatch is a Concept-based Search Engine











MolecularMatch Trials Data Global Depth of Coverage

> 150

Countries covered

> 72,000

Oncology-specific clinical trials

> 52,000,000

Medical terms in cancer ontology

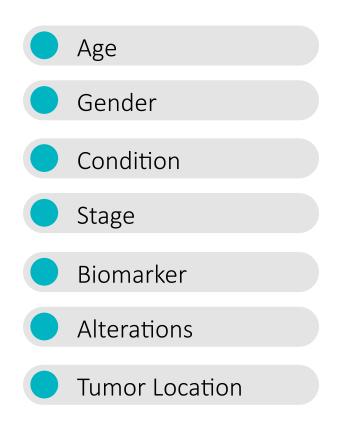


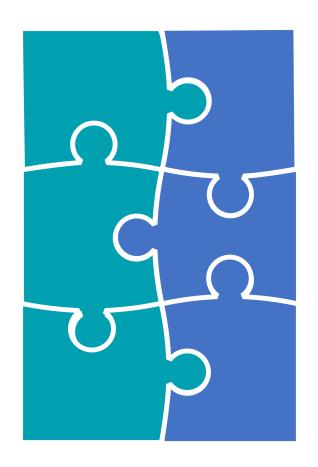
All Searchable by MolecularMatch
API or Web Portal



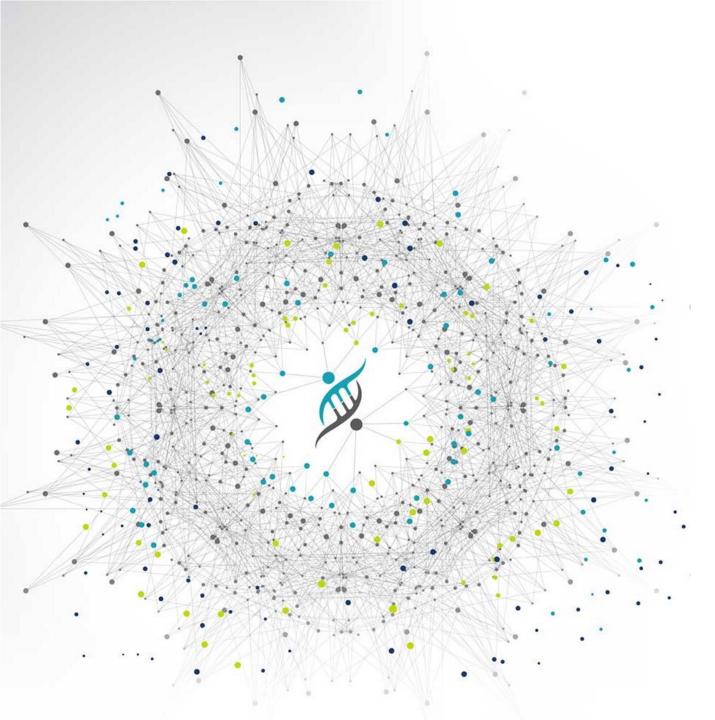
The Solution for Precision Medicine

Aggregated Clinically Relevant Metadata











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13th Nov 2019 Room 2004, TRI, Princess Alexandra Hospital, Woolloongabba, QLD

CASE PRESENTATION

Dr POH SEE CHOO

Medical Oncologist

Greenslopes Private Hospital

Mater Cancer Care Centre

Sunnybank Private Hospital

OVARIAN CARCINOMA

FIRST PRESENTATION

35 year old female

R oophorectomy in Fiji 2009

Borderline tumour

CA125 continue to be elevated post-surgery

6 cycles of Cisplatin chemotherapy

CA125 remained elevated post-chemotherapy

RECURRENT OVARIAN CANCER

Completion hysterectomy and L oophorectomy, resection of R adnexal mass in 2011 in New Zealand

-Review of original histology- confirm clear cell ca

Regular follow up- no recurrence with normal CA125

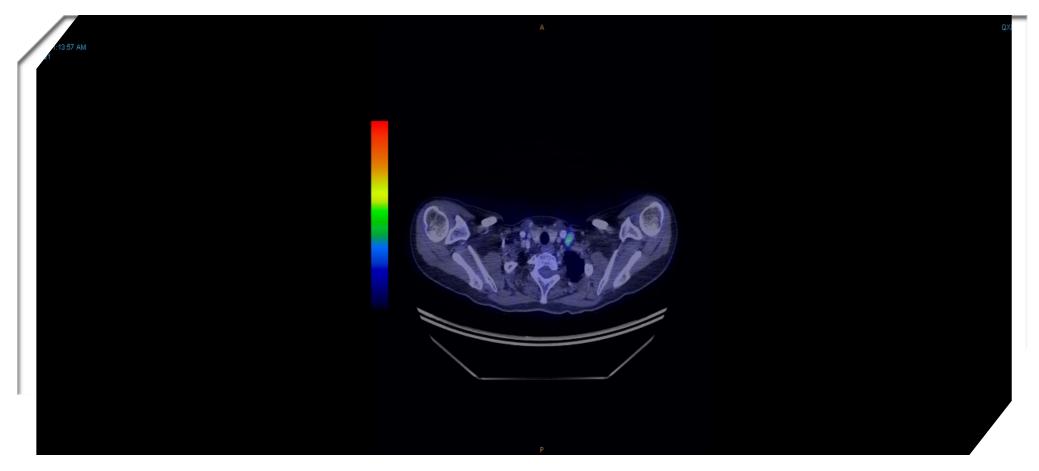
SECOND RECURRENCE

December 2016

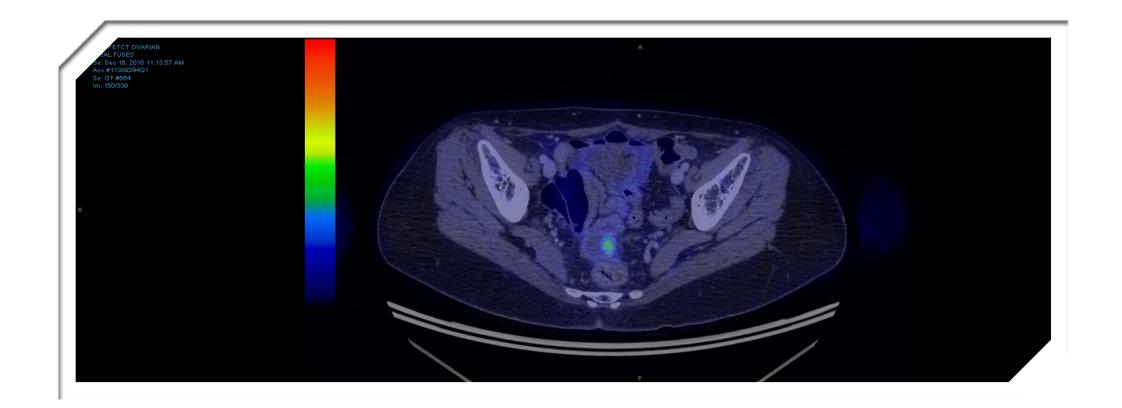
Elevated CA125 and L supraclavicular LN and pelvic recurrence on PET/CT

FNA confirmed recurrent disease in supraclavicular LN

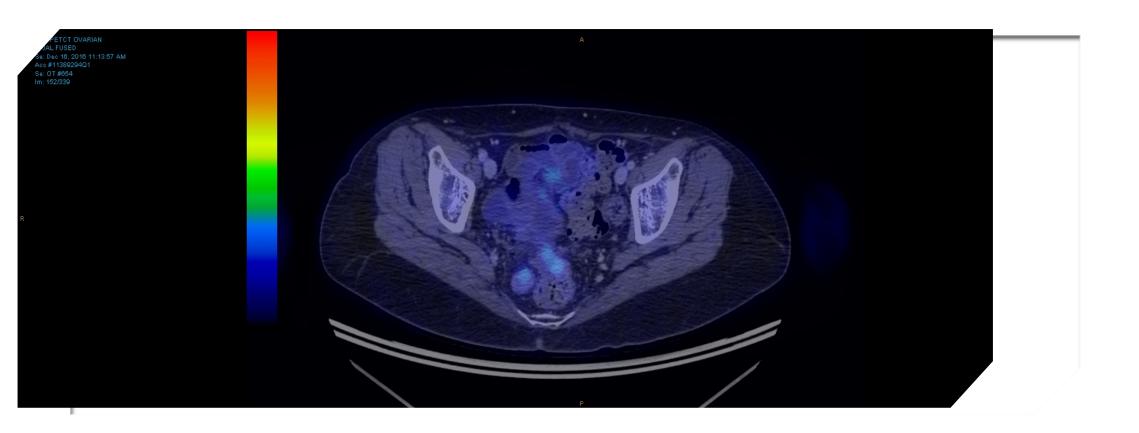
Received 3 cycles of weekly Carboplatin/Paclitaxel



DEC 2016



PELVIC RECURRENCE



PELVIC RECURRENCE

DEBULKING SURGERY

March 2017

laparoscopic resection of the pelvic mass and peritoneal metastasis with residual disease fixed to the sacrum and rectosigmoid mesentery

Colonoscopy- necrotic tumour invading into rectum

JULY 2017

Ultralow anterior resection and loop ileostomy

POST OPERATIVE CHEMOTHERAPY

April 2017 to June 2017

weekly carboplatin/paclitaxel

September 2017

Carboplatin/Paclitaxel with Avastin

Continue with 12 month Avastin

THIRD RECURRENCE

November 2018

Recurrence in the L supraclavicular and mediastinal LN

Commence on Caelyx in Jan 2019

Progression with cerebellar metastasis

Treated with stereotactic XRT

Changed to Avatisn/weekly Paclitaxel

So far responded to treatment. CA125- normal; 17