

# Prioritizing targeted cancer therapies with CDGnet

Panel - The National Cancer Institute's Informatics Technology for Cancer Research Program: Building a Community of Practice in Cancer Informatics

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I and my spouse/partner have no relevant relationships with commercial interests to disclose.

# Acknowledgments

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# Project goal

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Build a tool – **CDGnet** (Cancer Drug Gene networks) – to help researchers and eventually clinicians **expand the number of targeted therapies** available for individuals with cancer who have specific mutations (DNA changes) or genes/proteins that are overexpressed.

# Precision oncology framework

Precision oncology refers to tailoring interventions to patients in ways that go beyond traditional characteristics of age, sex, disease, symptoms etc by considering **biomarkers**.

Biomarkers may be:

- **genetic characteristics**: can be either *germline* (inherited, in normal tissue) or *somatic* (in cancer cells but not normal tissue)
- **mRNA or protein expression values**: refer to expression in tumors, either in comparison to other tumors or to adjacent normal tissues

# Tumor molecular profiling and targeted therapies

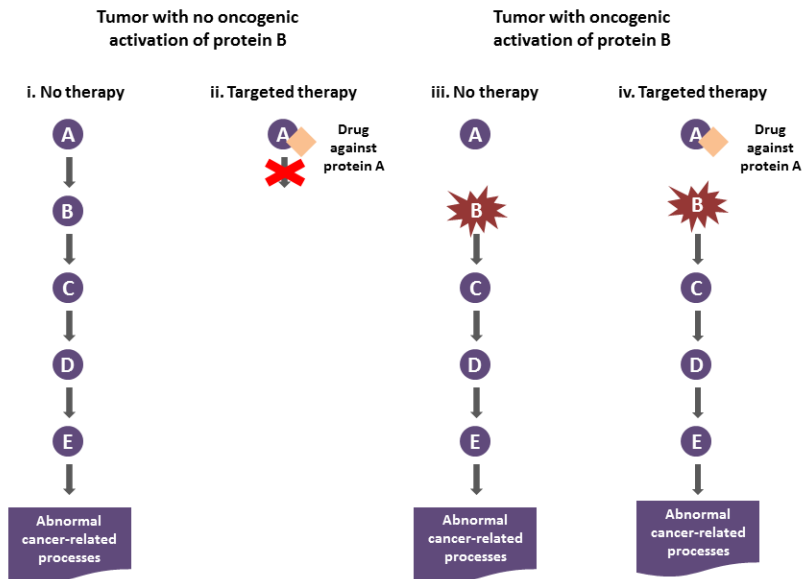
It is now routine to perform molecular profiling in certain tumor types to check for specific molecular features at diagnosis to decide on a targeted treatment plan (e.g. HER2+ breast cancer and trastuzumab)

In many cases tumor molecular profiling is used after a patient has progressed on multiple lines of therapy and/or has few/no therapy options left.

- Patient may then receive an off-label therapy that is prescribed for their alteration in another tumor type
- Our goal with CDGnet is to expand the number of possible targeted therapies and prioritize them using drug-gene network approaches

# How do we include pathway information?

## Look at downstream targets of oncogenes



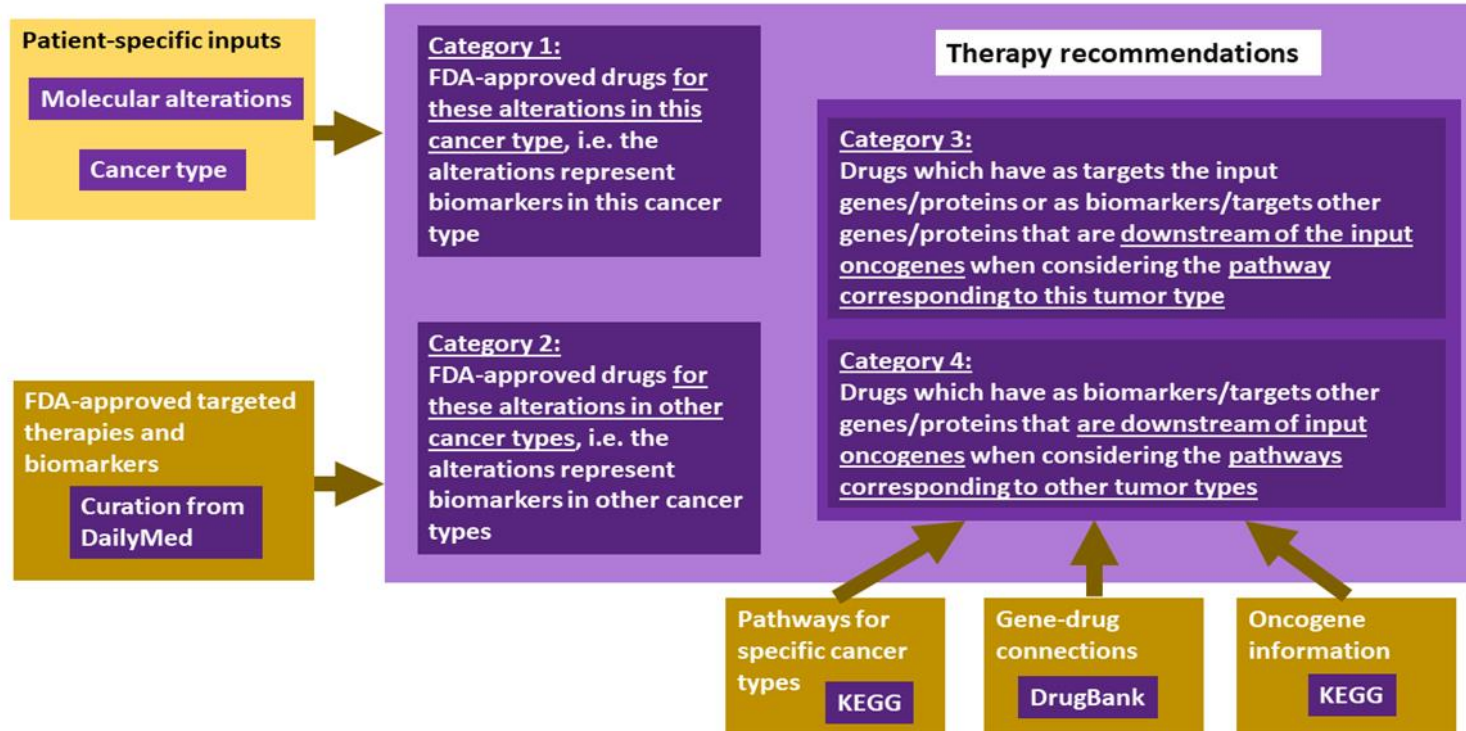
# Therapy prioritization: 4 categories

1. FDA-approved drugs for which the patient's alterations/genes/proteins are biomarkers in their tumor type
2. FDA-approved drugs for which the patient's alterations/genes/proteins are biomarkers in other tumor types
3. Drugs which have as targets these alterations/genes/proteins or as biomarkers/targets others that are downstream of input oncogenes when considering the pathway corresponding to this tumor type.
4. Drugs which have as targets/biomarkers either these alterations/genes/proteins or as biomarkers/targets others that are downstream of input oncogenes when considering the pathways corresponding to other tumor types.

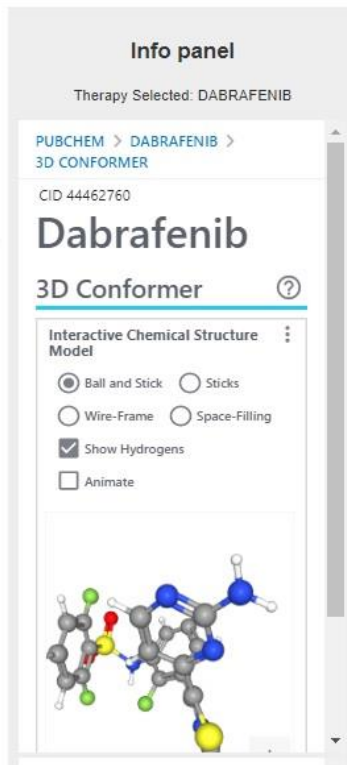
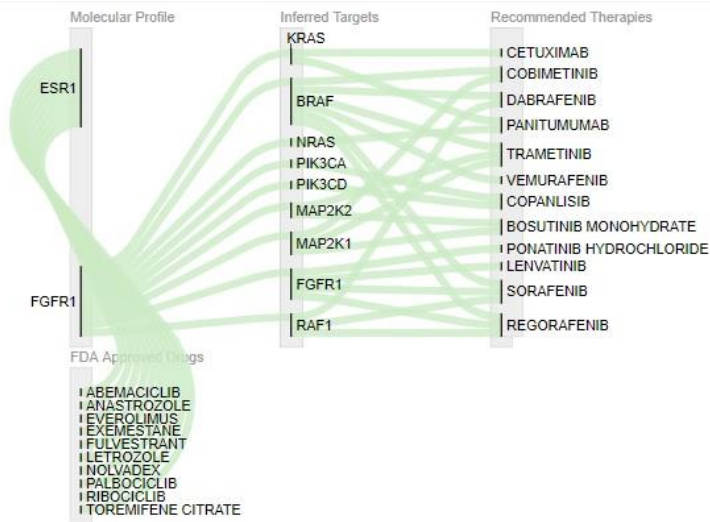




# General approach and data sources



# CDGnet flow diagram example for category 3 recommendations



[epiviz.cbcb.umd.edu/shiny/CDGnet/](http://epiviz.cbcb.umd.edu/shiny/CDGnet/)

# How did we choose the pathways?

- We are currently using KEGG pathways but considering alternatives
- **Benefits:**
  - Well-known, include pathways for many individual cancers
  - Expert-curated pathways
- **Issues:**
  - May not always agree with papers chosen by curators
  - Not all cancer types included
  - Due to KEGG's policy on data use, cannot easily download their pathways and a complicated approach to pulling in their data
    - This approach makes it difficult to perform updates and harder for others to reproduce our work

# How did we choose the drug targets?

- We considered the DrugBank database, due to its comprehensiveness, expert-curation, and ease-of-use
- Note that DrugBank is not tissue-specific

The screenshot shows the DrugBank website interface. The browser address bar displays the URL <https://www.drugbank.ca/drugs/DB00188>. The DrugBank logo is visible in the top left of the page header, along with navigation links for 'Browse', 'Search', and 'Downloads'. The 'TARGETS' tab is selected, showing a list of targets. The first target is '1. Proteasome subunit beta type-5'. Below this, a table provides details about the target:

Kind	Protein	General Function	Threonine
Organism	Humans	Specific Function	The proteasome is a large protein complex that degrades proteins.
Pharmacological action	Yes		
Actions	Inhibitor		
		Gene Name	PSMB5
		Uniprot ID	P28074
		Uniprot Name	Proteasome subunit beta type-5
		Molecular Weight	28480.0

The second target listed is '2. Proteasome subunit beta type-1'.

# How has being part of ITCR helped CDGnet development?

- Initial funding!
- Learning about other tools and collaborating with other investigators
  - Supplement to R21 to work with Tim Spicer and Louis Scampavia at Scripps – who was funded through the IMAT program – to integrate CDGnet into 3D spheroid/organoid high-throughput drug screening
  - Working with NDEx team to consider different pathways, share our pathways in NDEx, eventually integrate with Cytoscape
  - Exploring opportunities with CIViC and OncoMX
- Learning about best practices in reproducibility and training

# Managing CDGnet as an open source software project

- Open-source projects are more than a code base!
- We have a code repository at <https://github.com/SiminaB/CDGnet> and a web application at <http://epiviz.cbcb.umd.edu/shiny/CDGnet/>
  - Currently do not have automatic updating
- Github repository includes README file, as well as:
  - Code of conduct – Many large projects and societies/scientific meetings now have this, to reduce possible harassment and bullying
  - Contribution guidelines and list of current contributors
  - Documentation that includes a step-by-step analysis for a patient use case scenario
  - R markdown notebook for a specific example
  - Example input files

# Relevant links

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- Shiny-based web application available at <http://epiviz.cbcb.umd.edu/shiny/CDGnet/>
- Code available at <https://github.com/SiminaB/CDGnet> and <https://github.com/jkanche/nfpmShinyComponent> (package for interactive visualization)
- Preprint available at <https://www.biorxiv.org/content/10.1101/605261v1> (Kancherla, Rao et al)

# Thank you!

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