A (rare disease) Patient's Approach to Innovative Clinical Research

RareKidneyCancer.org

W.G. Paseman
bill@RareKidneyCancer.org

20240206

Agenda

- ***Who am I? - EHR**
- **Critique of Medical Innovation by FDA's Janet Woodcock**
- **A proposed Solution: Hackathons**
  - Using Gamified Tumor Boards to advance Research on particular rare disease patients
- **Next Gen approaches to automate data sharing and research**
  - Digital Twins
  - Research Agents

Who am I? - EHR

- 201311 – Diagnosed DVT – Put on Warfarin
- 201402 – Diagnosed second DVT (while on Warfarin)
  - JHH - Diagnosed with Kidney Cancer and Brain Tumor (1 x 1.2 x 0.8 cm)
- 201403 – Total left Nephrectomy – Pathology indicated papillary Renal Cell Carcinoma
- 201406 – Declined EVEREST (adjuvant) Clinical Trial
  - I asked 13 physicians: Yes: 3; No: 5; Patient must decide: 5
  - Key Opinion: “I do not recommend any adjuvant trial w/ mTOR inhibitors or VEGF targeted agents for papillary RCC. There will be trials with immune checkpoint agents in the near future, but not soon enough to enroll on.”
- 201602 – NIH Pathology indicates p1RCC (indolent – Yaaa!)
- 202303 – Meningioma hits limits for Radiation (1.8 x 1.6 x 1.5 cm - 2.51 cc)
  - I asked 14 physicians – photon guys said use photon; proton guys said use proton
  - Key: One with both said use photon; One said “Ask about the mechanic, not the tools”
  - Did Photon at UCSF
- 202309 - 1.8 x 1.5 x 1.6 cm – 2.43 cc

No SOC or CT for rare disease

https://thepatientstory.com/patient-stories/kidney-cancer/bill-p/

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“It’s Not Working”*

- The current structure of the academic research discourages collaboration, Grant review processes, promotion criteria and even the concept of tenure ought to be reconsidered.
- In addition the process of creating and commercializing a drug has been hindered by corporate secrecy. When the science is valuable and “not to be shared,” drug trials are fragmented and costly.
- The goal isn’t just improving knowledge. The goal isn’t FDA approval. The goal is to improve human health
- As researchers aim to conquer rare diseases, for example, patients and advocacy groups have become more than just cheerleaders for innovation — they’ve become active participants in finding solutions.

* FDA’s Janet Woodcock 20190902 LA Times

Let's try something different

Baseline Approach:

Big teams doing the same things slowly and often secretly without any patient involvement.

Goal: Improve knowledge, FDA approval

Hackathon Approach

Small teams doing different things quickly and openly with patient involvement.

Goal: Improve Health

Small Teams doing different things

Cancer Research is done with **big teams**

Field	% of Papers with <4 Authors
Economics	85%
Astronomy + Astrophysics	41%
Genetics	21%
Cancer	12%

This surprised me because...

- NY Times: Can Science be too Big?
  - Big Teams confirm findings
  - Small Teams generate new ideas
  - Disruption is inversely proportional to author count
- Small teams are more agile
  - Small enables closely space milestones (speed)
  - Closely spaced milestones keep you on track

https://www.natureindex.com/news-blog/paper-authorship-goes-hyper
https://www.nytimes.com/2019/02/13/science/science-research-psychology.html

Researchers and Patients view time differently Quickly

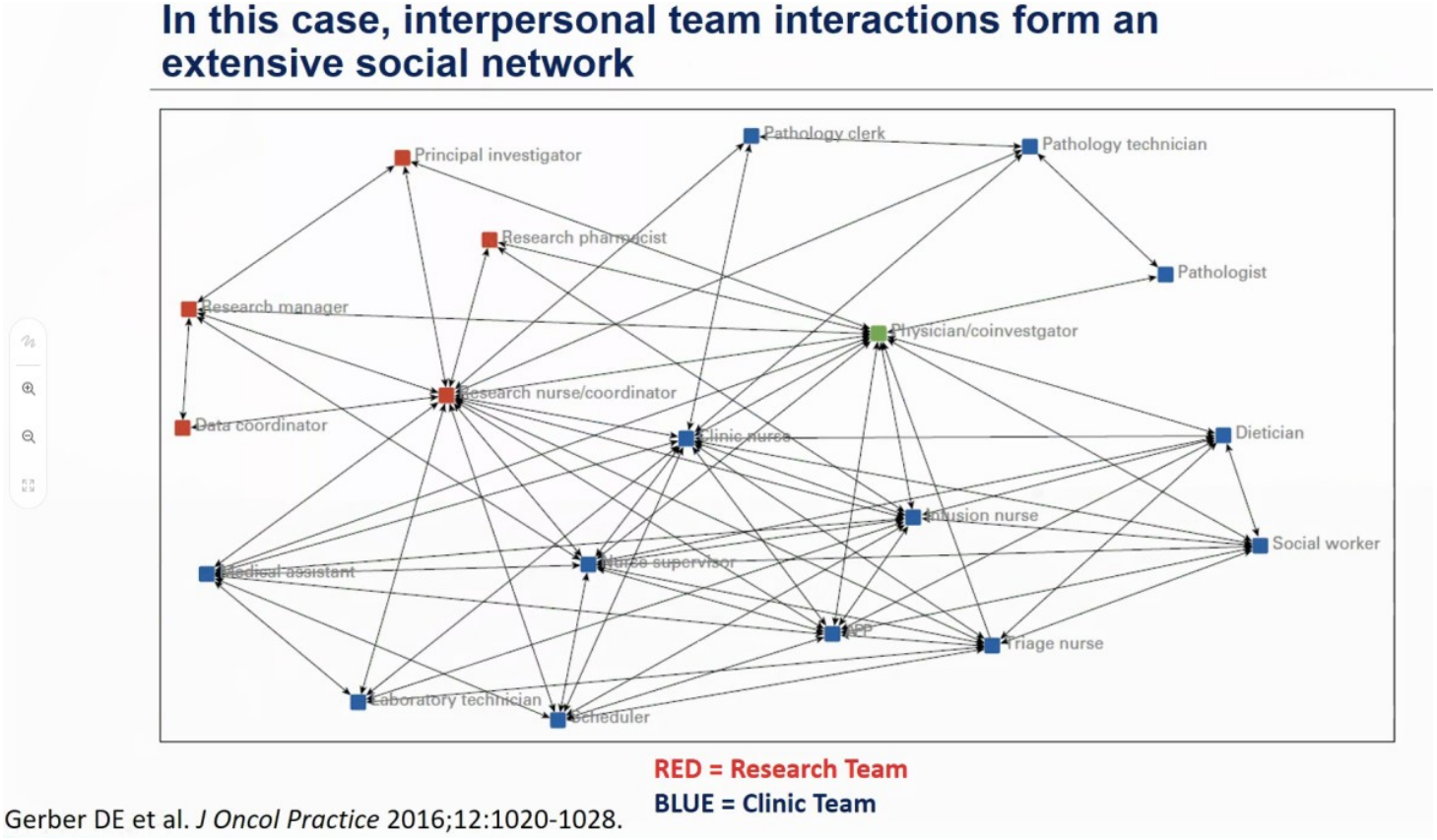
- Different Units of Measure
  - Researchers time measurement: yearly conferences and tenure tracks
  - Patients time measurement: PFS, OS, months between treatments; months until death
- Time pressure affects Motivation (for all you patients out there)
  - Researchers are motivated to win
  - (Some) Patients are desperate to win
    - 49ers Quarterback Steve Young: Everyone wants to win. Steve was desperate to win
  - Question to patients in the audience: Can you do something if your life depended on it?
    - If “yes” and you are not doing it now, it is a motivation issue, not a training issue
    - Hippocrates
      - It's far more important to know what person the disease has than what disease the person has
      - If you are not your own doctor, you are a fool.

Openly

- Now
  - Poor Reproducibility (Bayer) - 25% replication
  - Poor Reporting (alltrials.net)
  - Current problem: e.g. 20240123 Dana Farber retracts 6 studies

With Patient Involvement

In a Cancer Researchers' social network...



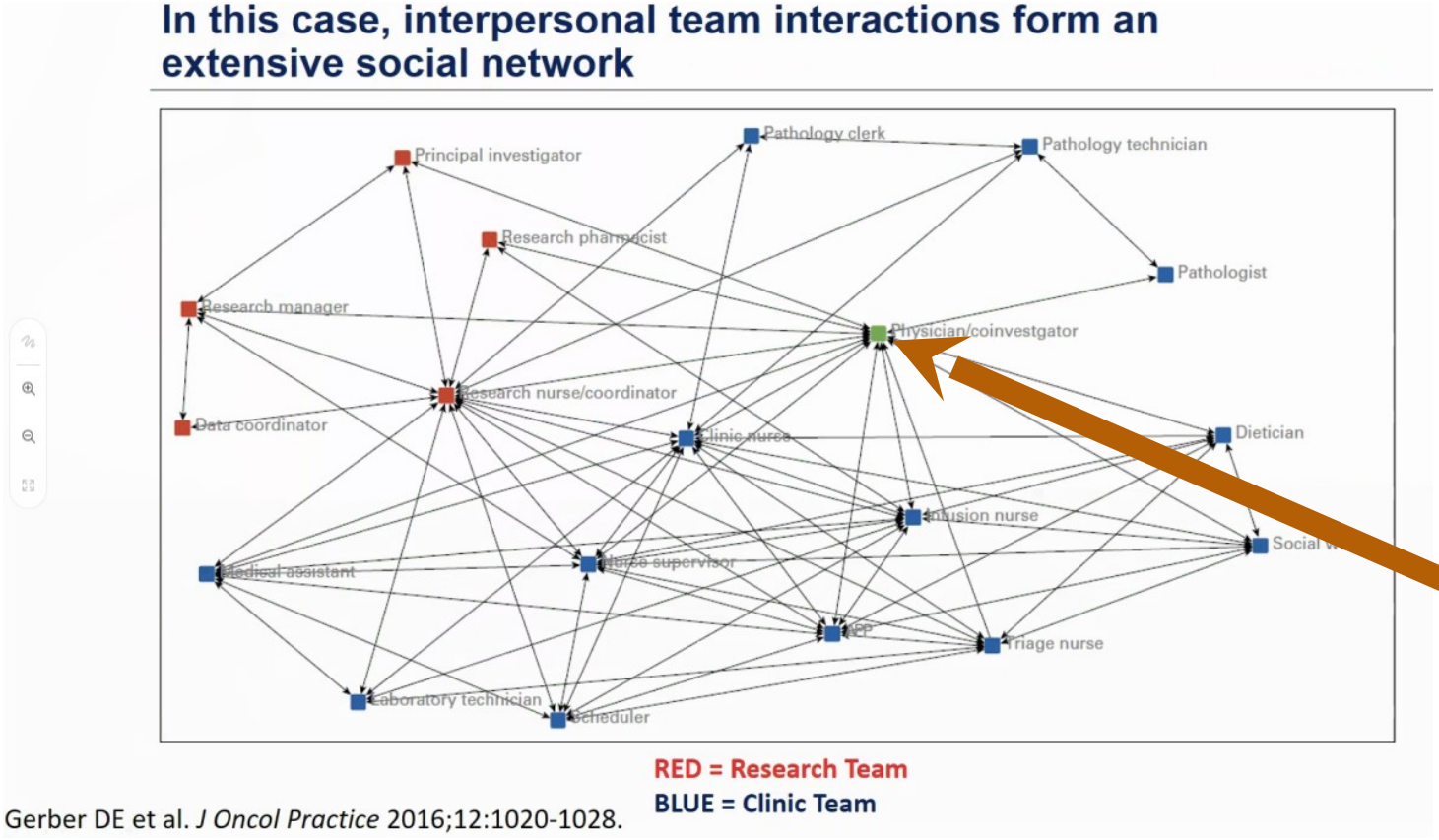
Gerber DE et al. *J Oncol Practice* 2016;12:1020-1028.

Principal investigator
Research Coordinator
Pathologist
Physician
...

With Patient Involvement

… patients don't exist

- Peanuts



Gerber DE et al. *J Oncol Practice* 2016;12:1020-1028.

Principal investigator
Research Coordinator
Pathologist
Physician
...

PATIENT!!

Patient Advocate (e.g. Smart Patients)

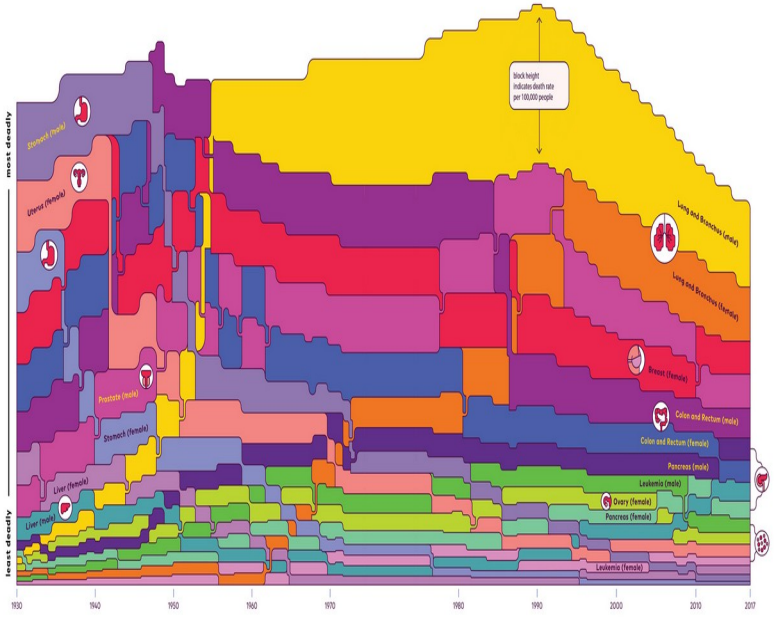
With Patient Involvement

"A "Seat at the Table" when its not your table

- Proposal: Renal Cancer incidence in Native Americans
- My Comments
  - Record if its a rare cancer
    - "Rate of Renal Cell Carcinoma Subtypes in Different Races"
  - Record Occupation
    - My "lived Experience":
      - Lots of Native Americans in Texas, Oklahoma and the Dakotas
      - Lots of Petrochem jobs in Texas, Oklahoma and the Dakotas
- All points rejected, none recorded.
- T-test comment not acted on.

Do something different, or get more of the same

Trends in Age-adjusted Cancer Death Rates by Site, Males



- We spend $150B treating the disease, yet a cancer patient is as likely to die of it today as one was 50 years ago.
- And most new drugs add mere months to ones life at agonizing physical and financial cost.
- The status quo is intolerable.
- The question is: **<u>What can be done?</u>**

*The First Cell – Azra Raza

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A proposed Solution: Hackathons

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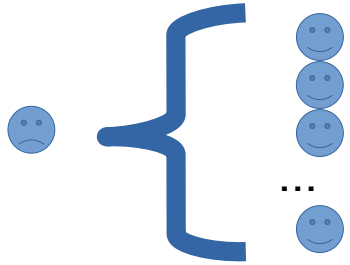
Goal: Improve Health

- If you don't like someone else's party, throw your own

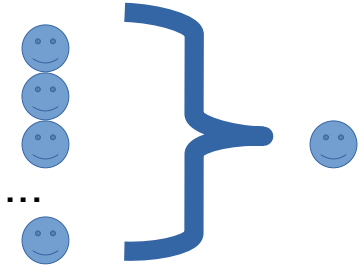
A proposed Solution: Hackathons

Clinical Trial: one Researcher many Patients



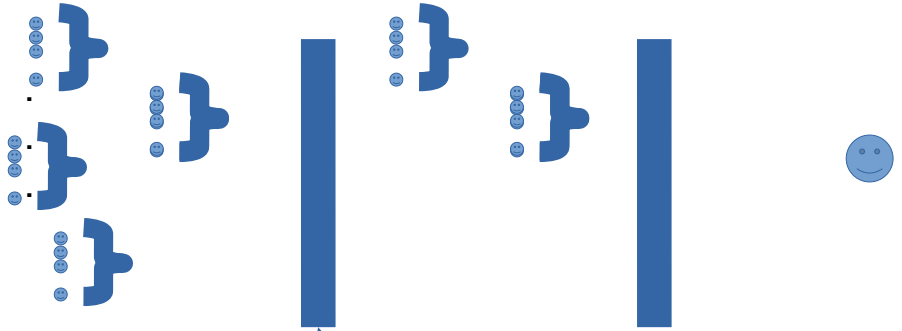
1 Researcher N Patients

Tumor Board: one Patient many Researchers



N Researchers 1 Patient

Hackathon: one Patient many TumorBoards (Teams)



Level 1 (Diagnosis)

Level 2 (Treatment)

1 Patient

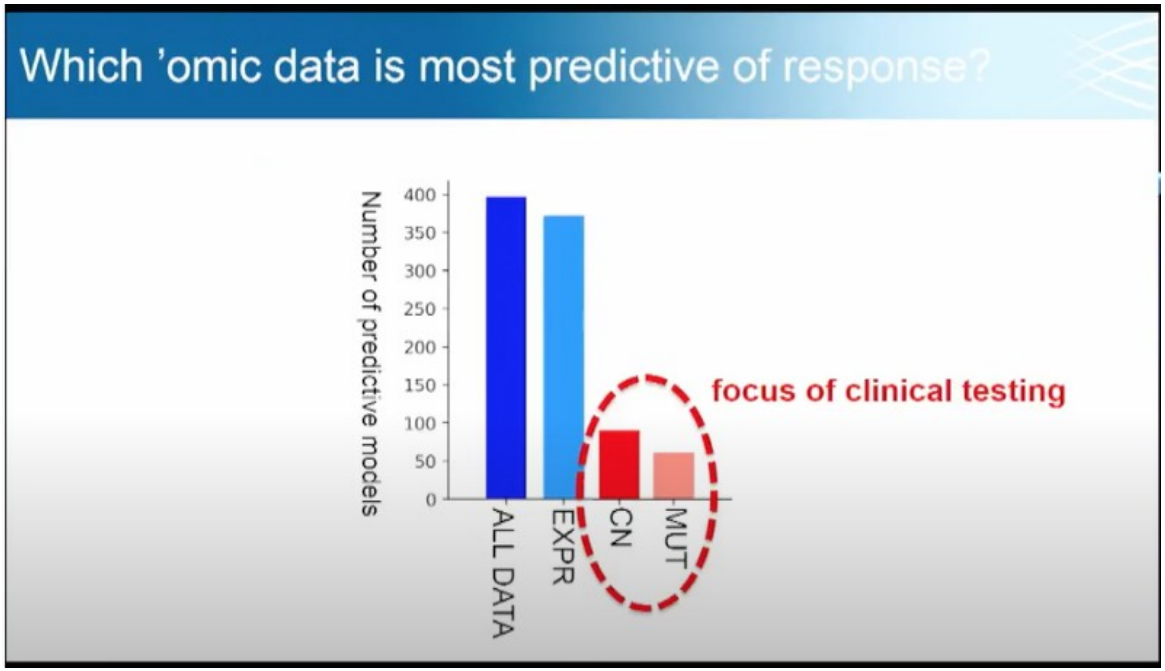
LeaderBoard (Scores)

Ensemble Reasoning (Kaggle)

- “The best way to have a good idea is to have lots of ideas.” - Linus Pauling

2018 p1RCC Hackathon Teams

Clinical vs. Research Data



James Watson: targeting metabolism is a more promising avenue in current cancer research than gene-centered approaches. 20160515 NYT

- Genetics- brother has thyroid cancer
- Genomics - TCGA Thyroid cancer clusters close to p1RCC
- Metabolomics - High Uric Acid - “Thyroid hormones influence kidney function and thereby might alter serum urate levels, a major risk factor for gouty arthritis.”
- Co-morbidities - Bradycardia (Slow Heart Rate): “hypothyroidism results in an insufficient amount of thyroid hormone which leads to a slower heart rate
- Dental Records

2018 p1RCC DNA Hackathon Process

TCGA Data

Bill Data (DNA)

Ensemble of 17 weak classifiers

119 Recommended Genes

cancer-genome-workbench								
causalnucleotidenetwork								
RecausalNucleotideNetworks								
Aizheng	AKR1B10	BASP1P1	CLEC2B	CYP4F11	LINC00621	PLEKHO1	PLEKHO2	...
BioMarkers.ai	DMRT2	FHL1	KNG1	PTGER3	UMOD			
DamTheRiver	AC139425.3	ACSM2A	ANO9	AQP12B	GRIN3B	HEXB	HIVEP3	...
GEViz	NRF2-ARE							
HelloKidney	ITGAM	TNFSF4						
KidneyBean	TUBB8							
studentec	AMPD2	DPP6	FLG2	FTMT	ST6GALNAC5			
trimericOGs	AGBL4	ARIDA1	CUL-2	HPSE2	LAMC-1	SK3	TRABD2B	...
DeeperDrugs	**BARD1**	APOB	CDK9	TTRAP				
GNOME	**BARD1**	**PDE4DIP**	AHNAK	ANAPC1	BCLAF1	DNAJ27	PABPC1	...
HelloKidney2		**PDE4DIP**	FOLH1	GDNF	MTHFR	PFKP	PSMA	...
codeomics						**MTOR**	**PIK3CA**	...
HSIEH	**SETD2**	**NF2**	**BAP1**	**KDM6A**	**PBRM1**	**MTOR**	**PIK3CA**	...
ExpressForce	**SETD2**	**NF2**	**BAP1**	**KDM6A**	**PBRM1**	**FGFR1**	ARID1A	...
HIF1AIsNotAnOncogene						**FGFR1**	CDK4	...

10 Up Weighted Classifications (Genes)

- BARD1
- PDE4DP
- SETD2
- NF2
- BAP1
- KDM6A
- PBRM1
- MTOR
- PIK3CA
- FGFR1

SCIENTIFIC REPORTS

OPEN

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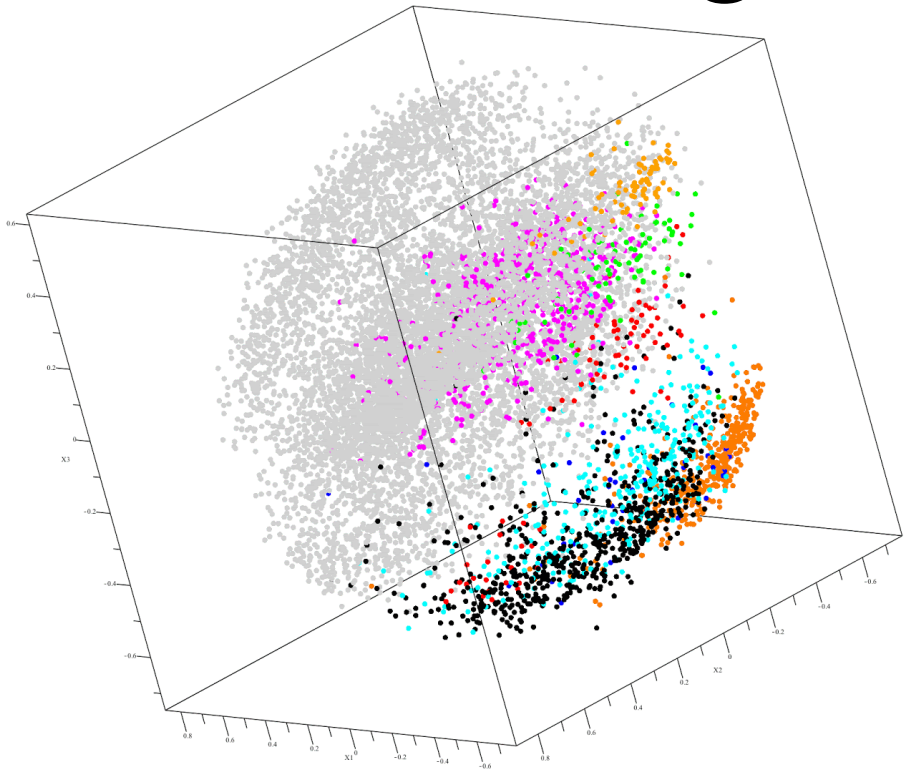
Accepted: 28 January 2019
Published online: 27 February 2019

Linking Binary Gene Relationships to Drivers of Renal Cell Carcinoma Reveals Convergent Function in Alternate Tumor Progression Paths

William L. Poehlman[1], James J. Hsieh[2] & F. Alex Feltus[1]

Renal cell carcinoma (RCC) subtypes are characterized by distinct molecular profiles. Using RNA expression profiles from 1,009 RCC samples, we constructed a condition-annotated gene coexpression network (GCN). The RCC GCN contains binary gene coexpression relationships (edges) specific to conditions including RCC subtype and tumor stage. As an application of this resource, we discovered RCC GCN edges and modules that were associated with genetic lesions in known RCC driver genes, including VHL, a common initiating clear cell RCC (ccRCC) genetic lesion, and PBRM1 and BAP1 which are early genetic lesions in the Braided Cancer River Model (BCRM). Since ccRCC tumors with PBRM1 mutations respond to targeted therapy differently than tumors with BAP1 mutations, we focused on ccRCC-specific edges associated with tumors that exhibit alternate mutation profiles: VHL-PBRM1 or VHL-BAP1. We found specific blends molecular functions associated with these two mutation paths. Despite these mutation-associated edges having unique genes, they were enriched for the same immunological functions suggesting a convergent functional role for alternate gene sets consistent with the BCRM. The condition annotated RCC GCN described herein is a novel data mining resource for the assignment of polygenic biomarkers and their relationships to RCC tumors with specific molecular and mutational profiles.

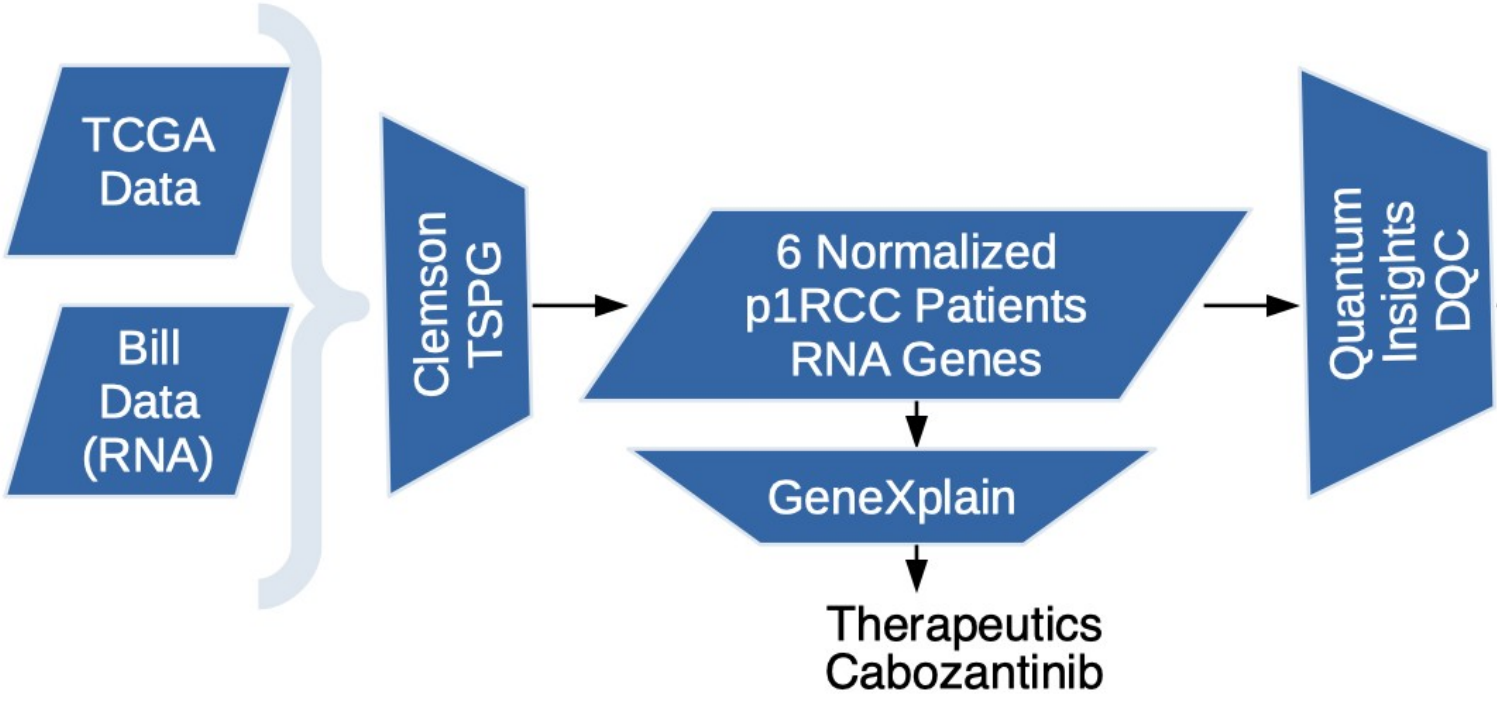
Parents,Siblings,Cohort Genetics



KIRP	Cyan
KIRC	Black
KICH	Red
LIHC	Coral
THCA	Magenta
CHOL	Blue
UVM	Orange
ACC	Lime
Others	Light Gray

2018 QuantumInsights.io DQC

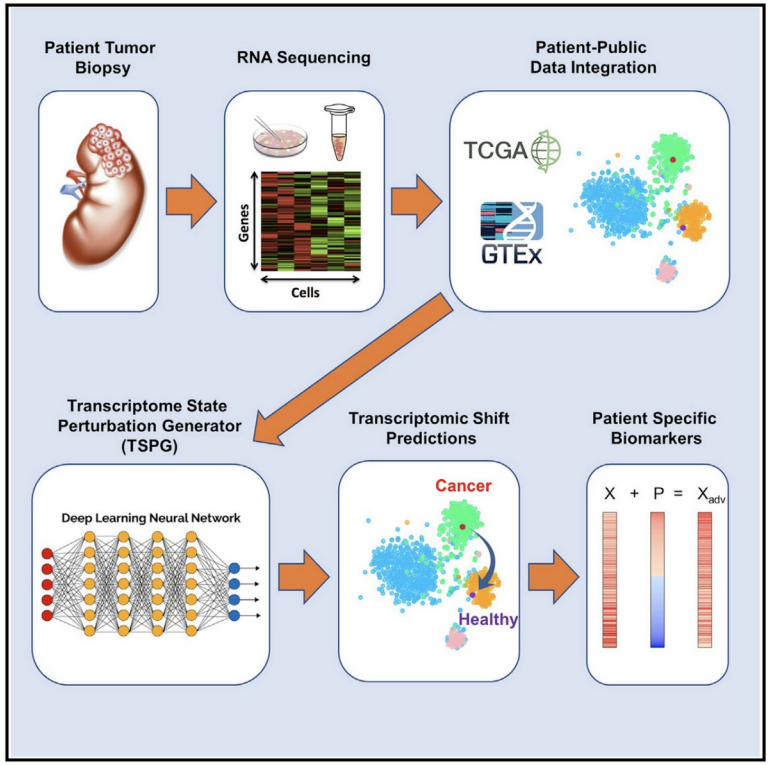
2020 p1RCC RNA Hackathon Process



- Three Teams
  - Clemson
  - GeneXplain
  - QuantumInsights.io

Cellular State Transformations Using Deep Learning for Precision Medicine Applications

Graphical Abstract



Highlights

- We present the Transcriptome State Perturbation Generator (TSPG) application

Authors

Colin Targonski, M. Reed Bender, Benjamin T. Shealy, Benafsh Husain, Bill Paseman, Melissa C. Smith, F. Alex Feltus

Correspondence

ffeltus@clemson.edu

In Brief

Cells express genes in unique combinations that enable distinct functions. Using machine learning, we present an algorithm that takes a source gene expression snapshot and learns how to change it to mimic a target snapshot. We applied the Transcriptome State Perturbation Generator algorithm to learn which genes have changed in a single patient's tumor relative to a normal tissue sample. By knowing which gene expression changes are required to leave a normal state in a single person, it is possible to design therapeutic strategies tailored for that patient.

GAN Generation

Merging 2018 and 2020 Results

Team - 2018	Gene	BP-Tumor -2020
studentec	FLG2	-0.569807
BioMarkers.ai	FHL1	-0.370446
HelloKidney2	TAS2R19	-0.363179
ExpressForce	TERT	-0.358329
HelloKidney2	TYMS	-0.287382
...	...	
trimericOGs	HPSE2	0.567236
BioMarkers.ai	PTGER3	0.59603
BioMarkers.ai	DMRT2	0.621588
BioMarkers.ai	UMOD	0.657959
BioMarkers.ai	KNG1	0.668831

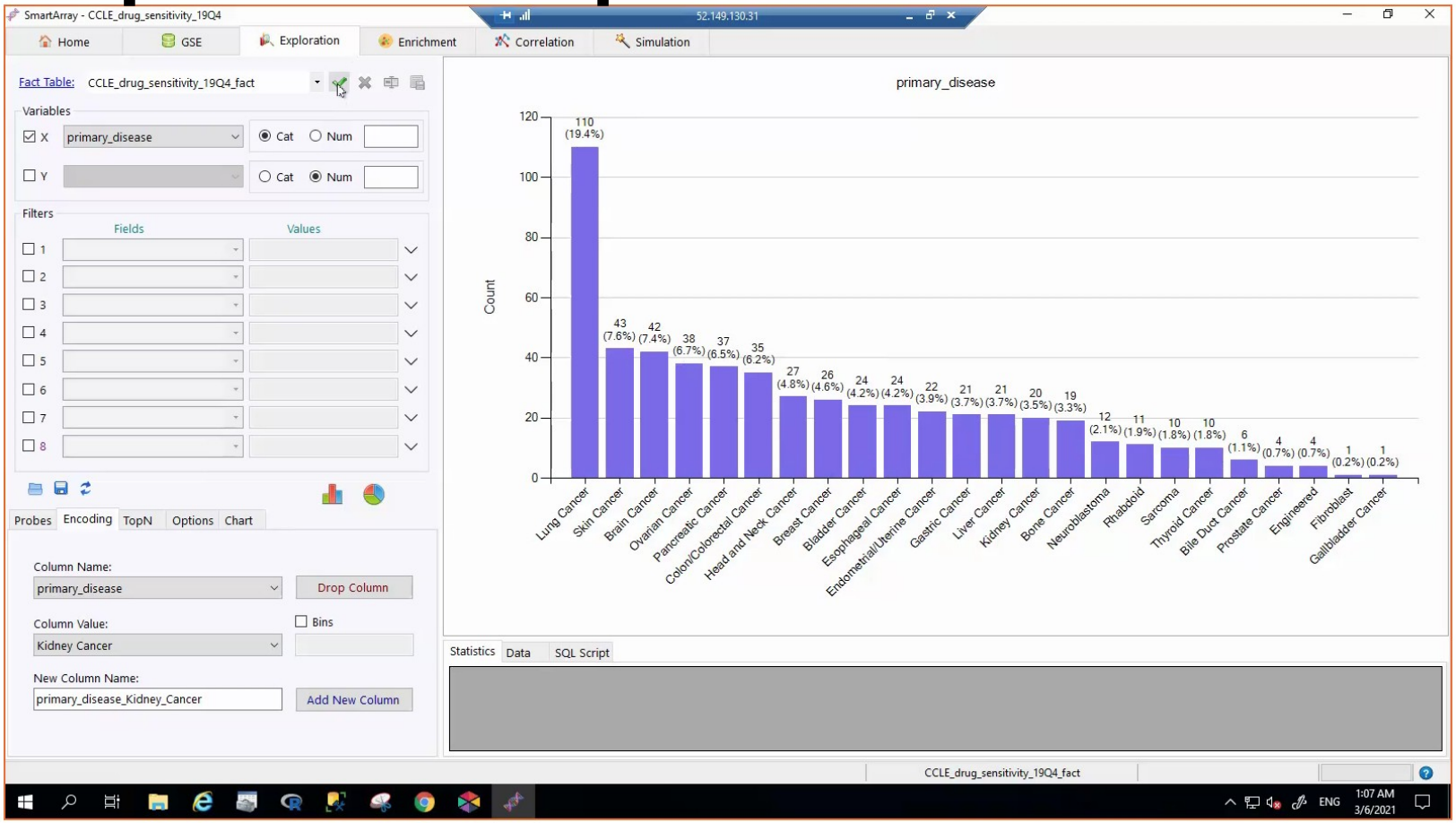
- 2018 p1RCC
  - 119 recommended Genes
- 2020 p1RCC
  - 6 Normalized
  - p1RCC patient's RNA genes

- BioMarkers.ai sorted to either end of the chart.
  - Perhaps diagnostic
  - Likely not therapeutic

Leaderboard Openly

Team - 2018	Gene	BP-Tumor -2020	Approach
studentec	FLG2	-0.569807	https://github.com/SVAI/studentec
BioMarkers.ai	FHL1	-0.370446	https://github.com/SVAI/Biomarkers.AI
HelloKidney2	TAS2R19	-0.363179	https://github.com/SVAI/HelloKidney2
ExpressForce	TERT	-0.358329	https://github.com/SVAI/ExpressForce
HelloKidney2	TYMS	-0.287382	https://github.com/SVAI/HelloKidney2
...	...		
trimericOGs	HPSE2	0.567236	https://github.com/SVAI/trimericOGs
BioMarkers.ai	PTGER3	0.59603	https://github.com/SVAI/Biomarkers.AI
BioMarkers.ai	DMRT2	0.621588	https://github.com/SVAI/Biomarkers.AI
BioMarkers.ai	UMOD	0.657959	https://github.com/SVAI/Biomarkers.AI
BioMarkers.ai	KNG1	0.668831	https://github.com/SVAI/Biomarkers.AI

Therapeutic Options



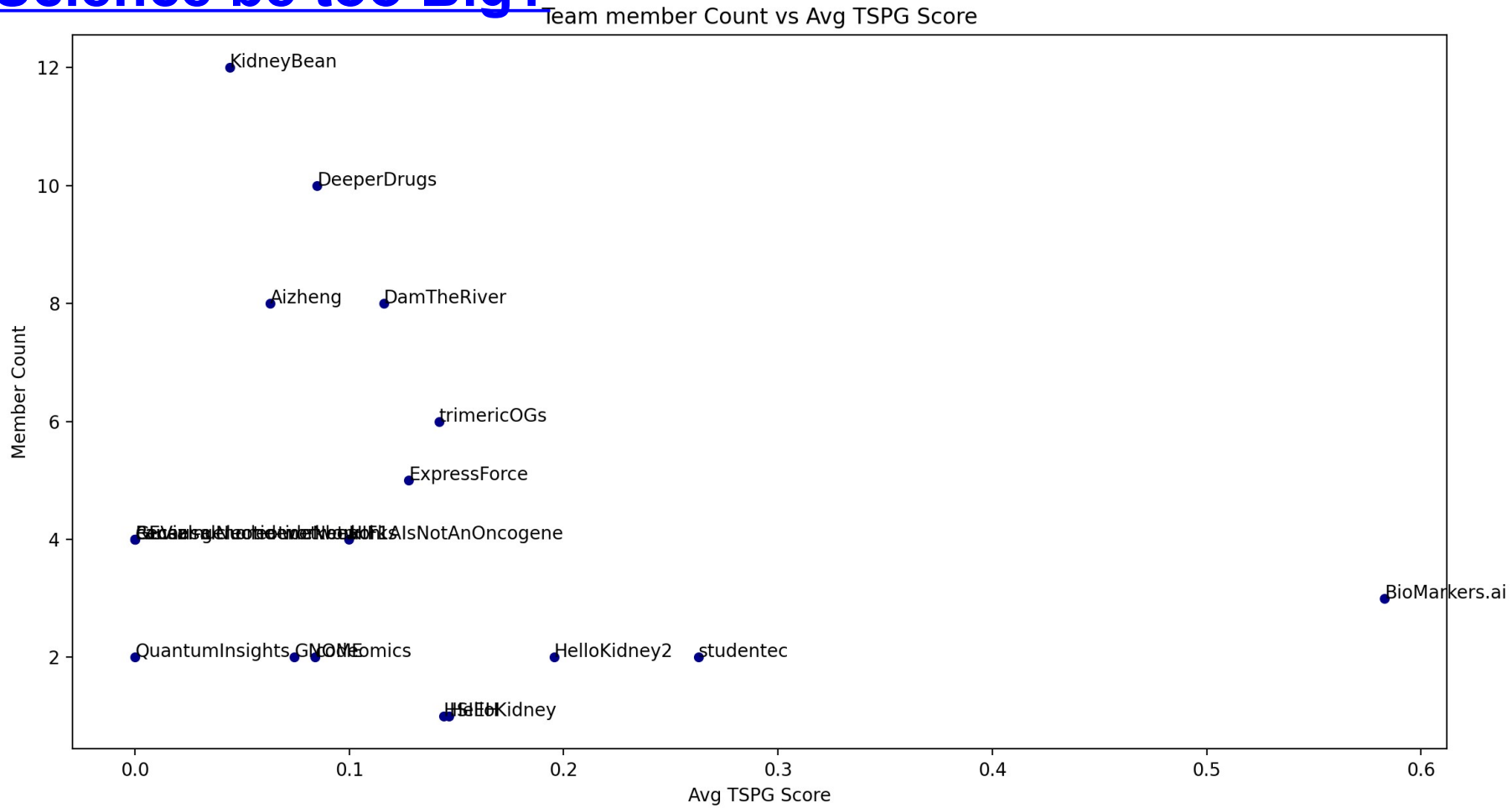
TBD: Wetlab

- Travera
  - 20 wells on a tray
  - Each with fresh tumor
  - And a different Treatment in each well
- Rare Cancer Research Foundation
- https://www.arctoris.com/
  - Cell Line Labs

Biomarkers.ai - 2018

- KNG1 uses alternative splicing to generate two different proteins: High MWt kininogen (HMWK) and MWt kininogen (LMWK). HMWK is essential for blood coagulation and assembly of the kallikrein-kinin system. This might explain my medical history.
  - Got warfarin/coumadin for diagnosis of deep vein thrombosis
  - DVT Symptoms returned. Went back and found: 7 cm mass left kidney, cerebral meningioma and spots in lung.
- Uromodulin (encoded by UMOD; also known as Tamm-Horsfall protein) is the most abundant protein in mammalian urine under normal physiological conditions.
  - UMOD can distinguish Normal Tissue from p1RCC with 100% accuracy.
  - Is UMOD also a good urine-based biomarker for p1RCC?
- FHL1 was an indicator for petrochemical exposure. For a time I worked in chemical refineries and on oil rigs. This might be the source of my somatic mutation.
  - Exposure to benzopyrene and several other agents enhances FHL1 expression

Can Science be too Big?



Patient Centered Game Elements Ensemble Learning

“Patient Centered”

- Patients view themselves as having a “rare disease” that is not served well by cohort analysis. We hope to use sibling and parent genetic data as a "control" in future events.
- Patients themselves host and maintain control of the event and are responsible for providing their own data.
- Data Control allows patients to create a current, longitudinal record over time for each subsequent hackathon as their disease develops.

“Game Elements”

- Hackathon participants are divided up into teams.
- The Game has “levels” which include diagnosis and therapeutic recommendations.
- Team’s results are “scored” which helps the Patient prioritize future research approaches.
- Scores can be posted on a LeaderBoard, which allows sharing of Research Approaches.

“Treat Research Teams as formal computational objects”

- Apply an “Ensemble Learning” technique called "bucket of models".
- For each model m in the bucket:
- Do c times: (where 'c' is some constant)
- Randomly divide the training dataset into two datasets: A, and B.
- Train m with A; Test m with B
- Select the model that obtains the highest average score
-

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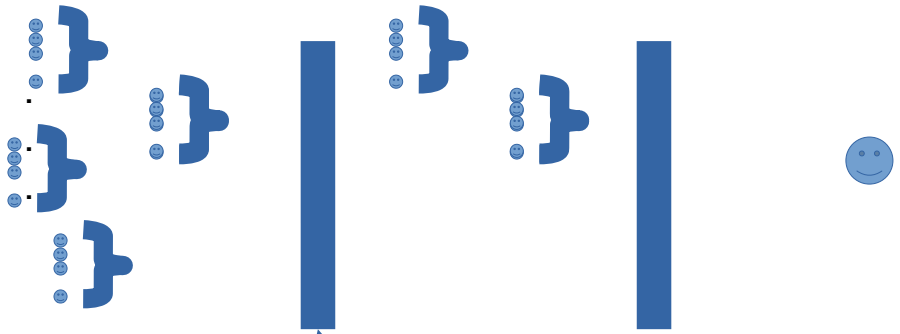
Next Gen approaches to automate data sharing and research

Now: Competition (**Gamification**)



Level 1
(Diagnosis)

Level 2
(Treatment)

1 Patient

LeaderBoard
(Scores)

- Ensemble Reasoning
  - "The best way to have a good idea is to have lots of ideas." - Linus Pauling

Next: Automation

- Process – Basically, Hackathons are multi-armed clinical trials for research processes.
  - Scale Hackathons up and make them faster.
  - Automate creation of Hackathon variants.
- Researcher(s) → LLM Agent team members
  - Chatbot → ResearchBot
- Patient(s) → LLM Agent Digital Twins
  - HIPAA – Not an issue for me
    - Need to do better later
  - EHR → DigitalTwin
    - Need to Represent Time Well
    - Integrate Genomics/Radiology later
    - Diagnosis vs hallucinations
- Data → Genomic GANS for rare diseases

Final Word

An obvious point needs to be made explicitly here. Though I contributed data on the front end of the process and did a few SQL table joins and sorts on the back end of the process, NONE of the biology is my work. It is the work of the many volunteer researchers who spent many hours exploring my data on my behalf. Thank You.

"If you work on frequent cancers, do randomized trials! If you work on rare cancers—find friends!"Olson, TA, Schneider, DT, Brecht, IB, et al. Rare tumors: a different perspective on oncology. In: Schneider, DT, Brecht, IB, Olson, TA, Ferrari, A, eds. Rare Tumors in Children and Adolescents. Berlin: Springer; 2012: 3–15.

Acknowledgements

- Tissue: UCSF's Dr. Max Meng and Tasha Lea
- Sequencing: Yale's Dr. Kaya Bilguvar and Christopher Castaldi and UCLA's Dr. Brian Shuch
- Sequencing Experiment Specification and Validation: Mike D'Amour for specifying the sequencing experiment parameters and fastq Validation Process
- 2018 Venue Donation: Salesforce's Steve Tamm and Lisa Ferrier
- 2018 Hackathon Teams
- 2018 Biomarker.ai Lead: Dr. Saed Sayad
- 2018 and 2020 Hackathon Master of Ceremonies: Ben Busby
- 2020 Hackathon: The TRI-con organizer: Kaitlyn Barago of healthtech
- 2020 Hackathon: Research to the People Organizer: Pete Kane
- 2020 Hackathon: "Clemson's 2020 normalized cohort" creators: Reed Bender, Ben Shealy and Benafsh Hussain from Dr. Alex Feltus' group
- 2020 Hackathon: Therapeutic Recommendations: GeneXplain's Dr. Jeannette Koschmann
- 2018 and 2020 Target Identification: QuantumInsights.io's Bernard Chen and Marvin Weinstein
- 2018 Hackathon: sv.ai volunteers: Ryan Leung, Clayton Melina, Lily Vittayarukskul, Hunter Dunbar, Pete Kane, Bill, Dom Jones, Marguerite, David Schachter, Anabelle Tang. Nina Sardesh, Sean Davis

2018 p1RCC HackathonTeams

Team	Members	Summary
Alzheng	Alex Feltus, Ben Shealy, Colin Targonski, Courtney Shearer, Eddie Weill, Ken Matusow, Sufeng Niu, William Poehlman	Model TCGA-RCC tumors as a “time series” across stage
BioMarkers.ai	Peyman Mirtaheri, Saed Sayad, Usman Qazi	Candidate p1RCC Biomarkers and environmental factors influencing expression
cancer-genome-workbench	Betty, rene lopez, Rui, Sarah	Predict/classify a sample cancer type using genetic data with: Unsupervised clustering, Dimensionality reduction, Somatic SNPs, Data exploration
causalnucleotidenetwork	Arkarachai Fungtammasan, Naina Thangaraj, Ola Zalcman, Steve Osazuwa	Variational Autoencoder and tSNE clustering
codeOmics	Daniel Hornburg, Milena Duerrbaum	Biomarkers to precision drugs
DamTheRiver	Andrew Wallace, Christian Clough, Felix Frayman, Matt Callahan, Nandita Damaraju, Pak Yu, Sebastian Nguyen, William Wright	Identification of neo-antigens present within patient P1RCC sequence data

2018 p1RCC HackathonTeams

DeeperDrugs	Andrew Mills, Biter Bilen, Jeff Lam, Lei Tian, Michael D'Amour, Monika Maleszewska, Prasun Mishra, Tahera Zabuawala, XIAOWEI ZHU	Rigorous variant filtering and target pruning
ExpressForce	Amrit Virdee, Maricris Macabeo, Nikhil Balaji, Sofia Medina Ruiz, Yuri Bendana	Netflix for Genes
geviz	Maytas Monsereenusorn, Natnicha Vanitchanant, Navi Tansaraviput, Thanapat Worasaran	Gene Expression Visualization
GNOME	In-Hee Lee, Sek Won Kong	Prioritizing germline and somatic variants potentially associated with p1RCC
HelloKidney	Terje Norderhaug	Autoimmune Clues to Kidney Cancer
HelloKidney2	Clinton Mielke, Robert Van Spyk	Genetic Markers
HIF1AIsNotAnOncogene	Eric Danziger, Joshua Bloomstein, Stephanie Kinnunen, Wanlin Zheng	A preliminary case study in EGFR

2018 p1RCC HackathonTeams

KidneyBean	Bea Nguy, Eric Kalosa-Kenyon, James (3), Jay (3), Kallen Schwark, Kandy Nachimuthu, Mabel Furutsuki, Maninder Singh, Marcus Strauss, Rahim Hashim, Sam Rapp, Wessam Sonbol	Drug candidates towards personal medicine
RecausalNucleotideNetworks	Andrew Carroll, Jason Chin, Pi-Chuan Chang, Samantha Zarate	How Effective Are Illumina Methods for BGI-SEQ? 20180531 BLOG POST
studentec	Brian Hanley, Rush Tehrani	USING BIGQUERY FOR GENOMIC DATA ANALYSIS
trimericOGs	Christine Kim, Lily Vittayarukskul, Phoebe So, Rohith Krishna, Samson Mataraso, senay yakut	Classifying Tumor Stages based on Structural Variants in Patient Data