

Drug repurposing for SARS-CoV-2 infection using machine learning and mechanistic models of the COVID-19 Disease Maps

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The problem

Learn potential **relations** between proteins (**targets of drugs**) and the **COVID-19 Disease Map**

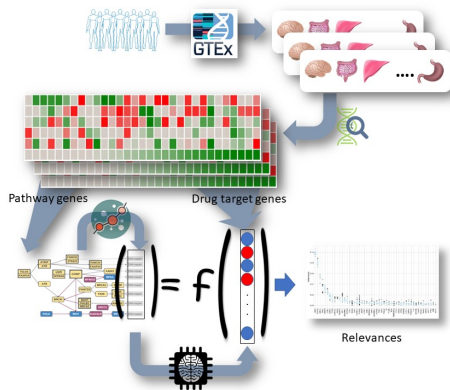
Mechanistic modeling of the MAP

Connect the **Drug** (Protein) Space with the MAP **Signalization** Space

Infer what is important to **regulate** the MAP

Suggest potential *repurposable* **drugs**.

Multi-output Supervised Learning



Multi-output Supervised Learning

Build the Signalization Space

>10k samples across 53 tissue sites
The COVID-19 Disease Map
277 circuits in 47 sig. pathways.

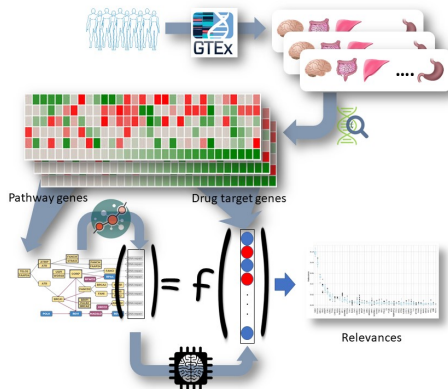
Hipathia Signal Transduction

$$S_n = v_n (1 - \prod_{s_a \in A} (1 - s_a)) \prod_{s_i \in I} (1 - s_i)$$

Connect the Dots

The Map results in 10k x 277 activities.
Over 2k KDT targeted by 1.7k drugs.
Most KDT lie outside the Disease Map.

Multi-output Supervised Learning



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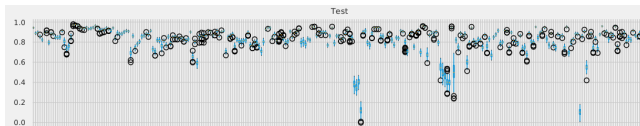
Data-driven Validation

Performance

Model: Multi-Output Random Forest
Hyper-parameter optimization with TPE
Repeated Nested 10-fold CV
Use SHAP values for explanations
Summarize by MeanAbs and take 1st decil

Explanation stability

Robustness Vs Stochastic, Noise, Sampling
100 Holdout splits of half the sample size
Split *training* into *learning* and *validation*
Check unbiased performance over *test*
Compute Nogueiras statistic test and CI

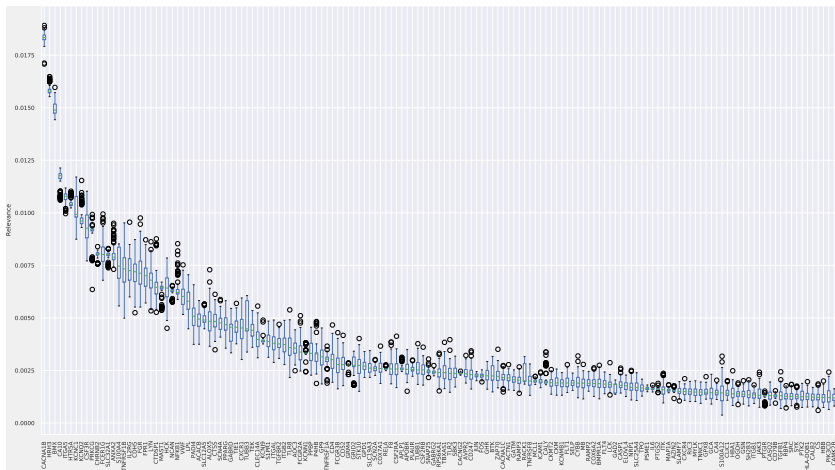


Mean R^2 : 0.82 ± 0.02
N-Stat CI: (0.729, 0.735)

Figure: R^2 score distribution over the test (R10-foldCV).

Classical Model Explanations

Useful but too broad. Only speak about the whole map. Misleading?



SHapley Additive exPlanations

Fair feature responsibility attribution.

Dis-aggregated by circuit by construction.

Additive (use biologically-relevant groups).

Template patterns

Conform a series of templates, e.g.:

Affect massively almost all MAP

Affect a few very specific circuits

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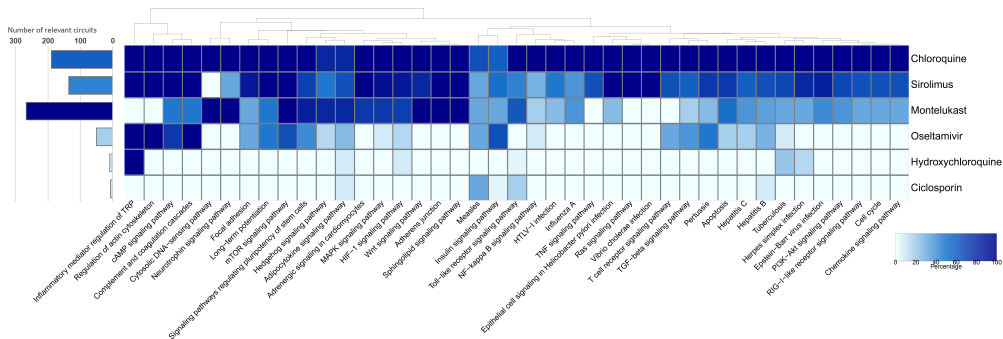
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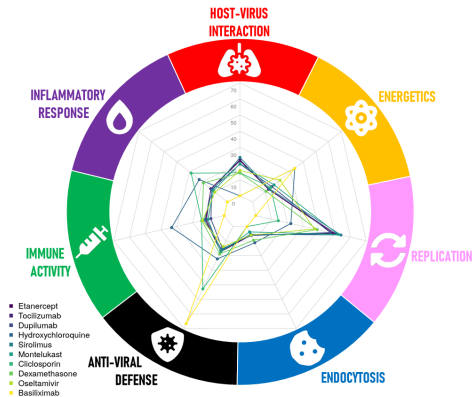
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Conform a series of templates, e.g.:

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Affect a few very specific circuits





380 KDTs (targeted by 679 drugs) have direct influence over the whole or partial parts of the map.

The GO biological processes enriched are mostly related to immune activity (T-cell, inflammatory response)

The COVID-19 Hallmarks are represented.

Future works

Speed the computation (GPU extensions).

Develop *sign* like SHAP aggregations.

Aggregations in *task*-space.

Use directly the COVID-19 DISEASE MAP (near done! see Kinza's talk, and our ML framework is DB-agnostic (**modular!**))

Machine learning software and other tools

python	ML coded in python 3.7 (and metadata parsing)
scikit-learn	ML pipelines, experimental design, etc
seaborn	ML plots
click	Command line interface
other	\LaTeX , bash, slurm, jupyter lab, Steven's \LaTeX styles, OBS studio ...

Accepted for publication in Signal Transduction & Targeted Therapy!

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Signal Transduction and Targeted Therapy

Drug repurposing for COVID-19 using machine learning and mechanistic models of signal transduction circuits related to SARS-CoV-2 infection

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and the BiER (CIBERER Network of Centers for Research in Rare Diseases)



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