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Computational Pharmacology: New Avenues for COVID-19 Therapeutics Search and Better Preparedness for Future Pandemic Crises

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Abstract

The COVID-19 pandemic created an unprecedented global healthcare emergency prompting the exploration of new therapeutic avenues, including drug repurposing. A large number of ongoing studies revealed pervasive issues in clinical research, such as the lack of accessible and organised data. Moreover, current shortcomings in clinical studies highlighted the need for a multi-faceted approach to tackle this health crisis. Thus, we set out to explore and develop new strategies for drug repositioning by employing computational pharmacology, data mining, systems biology, and computational chemistry to advance shared efforts in identifying key targets, affected networks, and potential pharmaceutical intervention options. Our study revealed that formulating pharmacological strategies should rely on both therapeutic targets and their networks. We showed how data mining can reveal regulatory patterns, capture novel targets, alert about side-effects, and help identify new therapeutic avenues. We also highlighted the importance of the miRNA regulatory layer and how this information could be used to monitor disease progression or devise treatment strategies. Importantly, our work bridged the interactome with the chemical compound space to better understand the complex landscape of COVID-19 drugs. Machine and deep learning allowed us to showcase limitations in current chemical libraries for COVID-19 suggesting that both *in silico* and experimental analyses should be combined to retrieve therapeutically valuable compounds. Based on the gathered data, we strongly advocate for taking this opportunity to establish robust practices for treating today's and future infectious diseases by preparing solid analytical frameworks.

Introduction

A global healthcare crisis created by the COVID-19 pandemic led to an unprecedented challenge in healthcare. This infectious disease caused by the SARS-CoV-2 virus first emerged in December 2019 in Wuhan, China and rapidly spread around the world affecting multiple countries. While the initial response of the World Health Organisation (WHO) was restrained to a 'health emergency', the status was soon recategorised as a 'pandemic' (1,2). COVID-19 patients presented with varying symptoms which were classified into three categories, namely mild, moderate, and severe. The disease progression, associated complications, and mortality were identified to show age and gender dependent differences with comorbidities also playing a significant role (3,4). The initial therapeutic management of COVID-19 was limited and the urgent nature of the pandemic prompted clinicians to try different approaches (3). A variety of protocols were developed to provide treatment and aid

patient recovery where the therapeutic options included antiviral drugs, anti-inflammatory drugs, immunomodulators and other types of intervention (3,5,6). As more information was collected and emergency approvals for clinical trials produced data, the clinical strategies were further refined (3,5,6). This also presented an opportunity to explore drug repurposing and search for effective clinical management options capturing both infection prevention and the alleviation of symptoms (2,7,8).

Repurposing drugs offers the advantage of reduced clinical development time and working with an already well-established pharmacological profile (9,10). Moreover, many researchers took full advantage of various *in silico* approaches, such as machine learning, bioinformatics or computational chemistry, to test new candidate molecules against various COVID-19 targets (11–13). While computational methods undoubtedly help to refine the complex chemical space, the field still requires a structured approach and practice establishment to ensure that outcomes are comparable and properly benchmarked (14–16). Thus, unsurprisingly most of the molecules that were selected for repurposing were not effective in treating severe COVID-19 cases leading to various controversies (10,17,18). The less-than-optimal results may also indicate that the research in drug repurposing focuses too narrowly on either the targets or the chemical entities without considering broader systemic implications (10,13,19–21).

Seeing the challenges and urgent need for new research and discovery strategies, we set out to explore the COVID-19 infection from the perspectives of computational pharmacology, systems biology, and cheminformatics bridging the system-wide effects with compound chemical features. Mining the data of available COVID-19 clinical trials allowed us to capture the links between compounds under investigation and their targets spanning both direct and expanded interactions (i.e., additional degrees of separation between interacting proteins). Since in-depth information on the COVID-19 clinical profile is limited and study-dependent (10,22), we used tested chemical entities as a proxy to understand the disease network and what pathways might be relevant if a drug proves to be beneficial. We employed several different techniques ranging from a direct to an extended interactome recreation which enabled us to identify a broader scope of affected signalling networks under a single drug influence (23,24). Our analysis of COVID-19 drug targets demonstrated that there is an overlap across certain target groups and the diversity of pharmaceuticals provides more options to adjust medications based on their systemic effects. Intriguingly, looking into the extended networks of drug targets, we found rich clusters of shared features. This allowed us to hypothesise that a direct drug-target interaction will have an effect on other associated interactors and this information could be used to assess drugs under investigation that share a predicted network (5,6,10,23–25). Moreover, this approach could be used to select

combination therapy regimens. Under these assumptions, we extracted the most noticeable clusters of drugs that have multiple shared targets through their signalling networks and used the ontology-based enrichment analysis to identify cellular processes, functional signalling networks, and linked pathways. Indeed, we were able to verify that compounds modulate a number of shared processes that are engaged through different network nodes. These findings have several important implications for future repurposing studies highlighting that focusing on a single target can lead to potentially missing a number of other relevant systemic effects (26). Selecting treatment options based on a network-centric perspective can provide insights into short- and long-term effects, and gathering such data could significantly improve how medical practitioners prescribe therapeutics and/or mitigate unwanted outcomes (5,27,28). Such methods could help establish robust practices to collect and organise data for drug repositioning and clinical studies (10,22,29).

We also explored the regulatory miRNA space and found that many target genes/proteins showed a staggering number of shared regulatory miRNAs. These findings also agree with the clinical miRNA observations in COVID-19; however, such data is still very limited (30,31). Our identified enrichment highlights that processes affected by the drugs used to treat viral infections have a complex regulatory interplay and analysing them further could help refine treatment options from biomarkers to RNA interference therapeutics (30,32,33).

While understanding the biological space is extremely important when optimising drug selection or identifying new antiviral and/or anti-inflammatory treatment methods, computational pharmacology can aid in bridging compound chemical features with observed effects (34–36). Thus, we explored the chemical profiles of the drugs that are undergoing or have been undergoing clinical trials for COVID-19. Such information can be a valuable guide when designing a screening library either by narrowing down or expanding drugs to diversify the compound set (34–36). In our study, we demonstrated that drugs used to treat COVID-19 do not have clear structural patterns, aside from a steroid subgroup and certain smaller subclusters. Chemical space mining (> 2.1 M compounds) and our built quantitative structure-activity relationship (QSAR) models via machine learning (ML) and deep learning (DL) enabled us to extract the most relevant features of drugs predicted to have antiviral properties against the SARS-CoV-2 virus. We applied the models to investigate which chemical entities in the COVID-19 clinical trial set could have antiviral properties. Both models had similar accuracy (ML- 96.77%; DL- 96.48%) differentiating between antiviral and control compounds when they were tested on the said drug set. Some of the identified drugs were already known antivirals tested for COVID-19 (e.g., ritonavir). To our surprise, when we used the trained models on 95 small fragment-like molecules that were experimentally verified to bind the M^{pro} protein on the SARS-CoV-2 virus (37), none were selected as antivirals. These results underscored

that current libraries used to test compounds might be too narrow or biased. Thus, experimental activity data on targets of interest is crucial in establishing QSAR relationships (38).

Drug repurposing and new compound development need to be built on diverse compound screening libraries with a strong understanding of their interactome and regulome. Such integrative approaches can prevent early failures in discovery pipelines or ineffective treatment regimens in repurposing studies (36,39). As can be seen from our in-depth analysis, evaluating compounds for COVID-19 should expand beyond direct drug-target interactions and consider a more complex space of affected networks in order to develop more robust combination therapies (Fig. 1). Thus, we should take this opportunity to establish research and discovery practices for today's and future infectious diseases by preparing solid analytical frameworks.

Methods

Data collection and mining

Data for COVID-19 associated clinical trials and drugs involved in treatment and/or clinical investigation protocols were primarily retrieved from the Open Targets platform that curates information on clinical testing, known targets, and compound information (40,41). Mining (November, 2021) returned 1,375 target-drug pairs which constituted 230 unique drugs and 356 unique targets (i.e., some drugs have multiple main targets or different drug formulations). In addition, Open Targets were searched for compound and known target associations to extract the relevant chemical data (e.g., SMILES, InchiKey, etc.) - this provided information on 18,376 compounds. To expand and verify the data sets, the information was cross-referenced against PubChem COVID-19 records (1,625 compound data) (42,43) and the STITCH database containing compound-protein interaction data (15,473,939 interaction points) (44). Additional interactome data was retrieved mining the STRING database (135,660 interactions, 5922 new targets for the expanded interactome network) (45,46). Reactome database was used to extract information on relevant pathways (47). miRNA database was used from the OmicInt package associated repository to mine non-coding RNA interactions (48). ChEMBL compound database (>2.1 M chemical entities) was used to search for similar and control compounds when investigating COVID-19 clinical trial drugs (49). COVID-19 CAS and Diamond/Xchem M^{pro} compound repositories were used to extract predicted and experimentally tested antiviral drugs (37,50).

Computational pharmacology and bioinformatics analysis

COVID-19 clinical trial data mining, cleaning, and analysis was performed in R programming environment (v4.1.2) with RStudio (51). Specific libraries used for enrichment, clustering, and ontology analyses include STRINGdb (v2.6.0), ClusterProfiler (v4.2.0), EnrichGO (v4.2.0), EnrichPathway (v4.2.0), and BioMart (v2.50.1) (52–56).

Cheminformatics analysis

Python programming environment (v3.9.7) (57) was used for chemical descriptor extraction, Morgan fingerprinting, Mol2vec fingerprinting (58), compound similarity assessment, substructure search, and image generation. Used packages and analytical frameworks include Rdkit (v2021.9.4), NumPy (v1.22.1), Pandas (v1.3.5), Seaborn (v0.11.2), Matplotlib (v3.5.1), and Chemexpy (v1.0.10) (59–64). Custom algorithmic assessments, comparative analyses, and data mining were performed using Rdkit (v2021.9.4) (59).

Machine and deep learning

Python programming environment (v3.9.7) (57) was used for machine and deep learning. The machine learning framework was implemented via Scikit-learn library (65) where LGBMClassifier (66) was used as a classifier with default parameters, train-test split at 0.2, where features comprised vectorised and normalised Morgan fingerprints (radius=3, nBits=2048). Deep learning neural networks were built for Mol2vec (58) encoded chemical features using the following set-up facilitated by TensorFlow and Keras libraries (67,68): sequential addition of layers starting with a Dense layer (hidden units=200, activation='relu', and input shape=(300,)), followed by Dense layers with hidden units: 128, 100, 50 and a dropout of 0.25 after each. All layers except the last were activated with 'relu' function, the last dense layer had only 1 hidden unit and sigmoid activation. Binary cross-entropy with adam optimiser and metrics for accuracy were used for the model compilation. The analysis was run for 200 epochs using 256 units for batch size with 0.2 split of the original data for validation. Deep learning was performed using Python 3 Google Compute Engine backend (Tensor processing units, TPU), RAM 12 GB, and HDD 107 GB.

Results

COVID-19 clinical trials represent a broad spectrum investigation of potential therapeutics to modify the disease course

We opted to use mined data from the Open Targets platform on referenced COVID-19 clinical trials (230 unique drugs with a known target status) so that our analyses were focused on a consistent set of compounds (40,41). Assessing clinical phase distributions for drugs that are undergoing or underwent clinical trials revealed that the majority of the therapeutics are in phase 2 (46.52%) with phase 3 and 4 being the other predominant categories at 28.26% and 20.43%, respectively (Fig. 2, A). Based on the available information, the largest proportion of clinical studies (50.87%) are still recruiting patients while other categories, such as 'Not yet recruiting' or 'Completed' distribute below the 20% mark (Fig. 2, B). Supplementary Figure 1 captures the clinical phase and status of every drug profile with a clear shift towards advanced clinical phases underscoring the emergency status of the disease and a large number of therapeutics under investigation to capture population-

based effects (40,41). Drug targets and types of drugs used to combat the COVID-19 infection and the associated complications (Fig. 2, C) further exemplify a broad therapeutic engagement. Most of the drugs in clinical trials are inhibitors (nearly half of all treatment options) with agonists (20.87%) and antagonists (17.83%) comprising the other two main categories (Fig. 2, D). About two thirds of the pharmacological intervention options belong to a small-molecule category (Fig. 2, F).

Antibodies (14.35%) and other proteins (10%) are two additional drug classes that are important in treating this viral infection (Fig. 2, F). However, it is necessary to note that an important evidence gap has been reported for the safety of the drugs tested for COVID-19. As reported in January, 2021, 40.4% of completed trials did not post results on ClinicalTrials.gov or in the academic literature (22).

COVID-19 treatment strategies highlight the need for a more in-depth understanding of drug pharmacological action

The original set of compounds (Open Targets, 230 unique drugs with a known target status) used to treat or investigated for the treatment of COVID-19 was searched for any shared proteins across the main known drug targets (Suppl. Fig. 2). Since a compound might be listed to engage one or more proteins, we aimed to cross-reference drugs and their targets to get insights into shared action (23). This analysis returned several smaller clusters with denser sets in the top left corner of the heatmap (Suppl. Fig. 2). For example, dipyridamole and pentoxifylline engage a broad spectrum of phosphodiesterases (PDEs), such as PDE3B, PDE1A, or PDE5A. Comparing dipyridamole and pentoxifylline action allowed identifying some additional main targets where these compounds differ. Specifically, dipyridamole inhibits the equilibrative nucleoside transporter-1 (ENT1 or SLC29A1) which serves a sodium-independent transporter for purine and pyrimidine nucleosides and pentoxifylline is believed to downregulate adenosine A2A receptor (A2AR)-mediated pathways (69–71). Propofol, sevoflurane, and isoflurane also form a noticeable group in the heatmap (Suppl. Fig. 2). These drugs are used in anaesthesia protocols for patients requiring mechanical ventilation and prolonged, deep sedation to optimize oxygenation and ventilation during respiratory failure from COVID-19 (72–75). Analgesia drugs also form a cluster through their shared action on PTGS1 and PTGS2, also known as cyclooxygenase 1 and cyclooxygenase 2 (76). The corticosteroid section (bottom left of the heatmap, Suppl. Fig. 2) have one shared target that stands out – nuclear receptor subfamily 3 group C member 1 (NR3C1). This receptor was implicated in the progression of the COVID-19 infection where a single cell transcriptome study revealed that the NR3C1-CXCL8-Neutrophil axis determines the severity of the COVID-19 disease (77).

Based on the drug-target network analysis, we concluded that in order to better understand pharmacological interaction processes, we needed to expand the drug interactome space.

Interactome investigation revealed new opportunities for drug repurposing and combination therapies

Taking into consideration the fact that targets are part of complex signalling networks prompted us to explore what the overlap size of the interactome is for each drug and if it could be used to find alternative therapies, repurpose existing ones, or help develop new drugs faster. In other words, the main known targets for each drug were expanded by mining protein-protein interaction networks to extract an estimated network that has links to the drug through its main target. These networks were searched for overlaps in pairwise drug comparisons where the overall protein set consisted of 5922 unique targets. On average each main drug target had about 97.6 interactors (Suppl. Fig. 3) ranging from no known interactors to a maximum number of 558 other proteins. This search allowed us to identify emerging patterns in data on a systemic level (Fig. 3) where three larger and two smaller clusters with relatively large, shared networks were found for different drugs. Clusters 1, 2, and 3 were found to be the least diverse considering the number of seed proteins (or the main drug targets) (Fig. 3; Suppl. Table 1 and Suppl. Fig. 4). By contrast, clusters 4 and 5 showed the most diversity with 17 and 29 unique seed proteins, respectively.

Exploring the two most diverse clusters, clusters 4 and 5, we can see that cluster 4 contains drugs used to treat hypertension, namely losartan, valsartan, telmisartan, candesartan, and ambrisentan. Losartan is an angiotensin II receptor blocker (ARB) used to treat hypertension and it has been proposed that this drug acting as a selective antagonist of the angiotensin II type 1 (AT1) receptor may offer some protection from lung damage induced by COVID-19 (78–80). As can be seen, cluster 4 drugs are predicted to mostly engage the same size networks; however, comparing and exploring drug-specific networks can help identify diverging biochemical processes that could be therapeutically relevant. One example of such an approach can be found in cluster 4 for ambrisentan - a selective type A endothelin (ET-A) receptor antagonist. This selective antagonist is used to primarily treat pulmonary arterial hypertension and has been applied in COVID-19 combination therapy with dapagliflozin which inhibits sodium glucose co-transporter-2 (SGLT-2) (81). While dapagliflozin has not been listed in the Open Targets COVID-19 clinical trial data (November 2021), we explored the extended interactome for SGLT-2 using the same principles as for other compounds. We then compared dapagliflozin with ambrisentan for any overlapping targets in their networks. The dapagliflozin network is relatively small – only 11 targets, while ambrisentan's network size is 183 proteins. The networks shared only two targets, namely adenylate cyclase 7 (ADCY7) and glucagon (GCG) highlighting the different cellular process engagement with a two-gene convergence point. Similarly, cluster 5 combines several different drug classes, e.g., aviptadil, prasugrel, chlorpromazine, and naltrexone, that converge through a shared interactor network (Fig.

3; Suppl. Table 1 and Suppl. Fig. 4). Ticagrelor and prasugrel (P2Y12 platelet inhibitors) have been employed to manage acute coronary syndrome (ACS) and as a thromboprophylaxis in patients with COVID-19. Numerous studies explored combinations of an enhanced prophylactic doses to correct the parameters of viscoelastic coagulation (82–84). Other drugs, such as chlorpromazine (phenothiazine antipsychotic), naltrexone (opioid receptor antagonist), and fingolimod (a sphingosine 1-phosphate receptor modulator), have been recognised for their multi-modulation potential and have also been tested in various clinical settings (85–88). Considering our findings, it is important to highlight the need for the consolidation and further exploration of the pleiotropic effects of the drugs used to treat COVID-19. Consequently, we selected cluster 5, the most diverse cluster, to explore the occurring interactions more in-depth.

Interactome analyses accentuated diverse process networks that could be used to advance therapeutics development

Rather than exploring genes in isolation we examined what signalling networks and functional processes can be enriched for the identified clusters. Figure 4 highlights that cluster 5 identified during the interactome analysis (Fig. 3, Suppl. Table 1 and 2) also shows varied functional enrichment patterns ranging from calcium ion homeostasis to neutrophil migration. A proportion of genes from cluster 5 also belongs to the extracellular signal-regulated protein kinase (ERK) cascade which has previously been reported as a potential therapeutic target in coronavirus infections where cascade inhibition was observed to lead to infection resolution (89). As can be seen other clusters had different profiles where clusters 3 and 4 had several shared functional themes involving haemostasis (Suppl. Fig. 5). Haemostatic aspects of COVID-19 have been reported as a serious concern in stabilising patients and reducing tissue/organ damage (82,90,91). Furthermore, clusters 1 and 2 are good examples demonstrating how even a few genes/proteins (i.e., cluster drug main targets) can impact multiple different cellular functions through the extended network; such considerations can always be useful when predicting drug effects or exploring alternative uses (Suppl. Fig. 5) (5,7,10). To better understand specific functions in a gene cluster, it is helpful to explore if any of the genes in the over-represented functional groups belong to the same pathway. Cluster 5 has several clear themes where some genes are shown to play a role in multiple pathways and others are much more pathway specific (Fig. 5). As an example we can inspect one drug - cenicriviroc from cluster 5 (experimental drug, inhibitor of C-C motif chemokine receptors, namely CCR2 and CCR5) which has several targets, e.g., interleukin 10 (IL-10), CCR5, C-C motif chemokine ligand 20 (CCL20), C-X-C motif chemokine ligand 10 (CXCL10), CD86, and formyl peptide receptor 1 (FPR1), belonging to the IL-10 signalling pathway (Fig. 5; Suppl. Table 1 and 2). Due to its apparent broad engagement spectrum, cenicriviroc has been included in clinical trials to

assess its anti-inflammatory and immunomodulatory effects (92–95). Drugs with different targets, such as icatibant (bradykinin 2 receptor antagonist) or ozanimod (sphingosine 1-phosphate receptor agonist), show a potential pathway overlap through shared network targets and this understanding could be useful in managing their COVID-19 clinical trials (Fig. 5; Suppl. Table 2) (96,97).

The identified enrichment clusters as well as individual drugs could be used to compare and match different combination therapy regimens, such as haemodynamics modulating and anti-inflammatory. In the case of anti-inflammatory action, it is possible to compare drugs having immediate vs long-term effect in order to reduce tissue damage occurring in acute and chronic disease progression.

miRNAs represent a potentially new biomarker and therapeutic modulation space linking the drug interactome

Seeing the complexity of the drug networks, we also analysed the non-coding regulatory layer for the most diverse cluster 5 (Fig. 3). We used the mined data of validated miRNAs and their regulome genes to explore the dynamics of miRNAs in the selected gene cluster (48). We found a rich network of miRNAs known to be involved in the regulation of multiple genes (Suppl. Fig. 6). We identified that some genes from cluster 5 are linked to miR-320 family which downregulation has been associated with the progression of disease severity and miR-320 have been suggested as potential biomarkers for SARS-CoV-2 (98,99). Other miRNAs from cluster 5 have also been reported in other studies as prognostic markers. For example, circulating miRNAs from ten COVID-19 patients (sampling done longitudinally with ten age and gender matched healthy donors) allowed to profile the alteration of 55 miRNAs in COVID-19 patients during early-stage disease (30). Our enrichment recovered miR-31-5p (marked upregulation in COVID-19 patients) and other strongly associated biomarker miRNAs, namely miR-423-5p and miR-23a-3p (30).

Compound chemical profiles capture certain interactome features and also reveal a highly heterogeneous chemical space

While various drugs showed interesting interactome overlaps, we assessed the chemical and pharmacological characteristics and investigated if network analyses can be superimposed to chemical data. Only 158 compounds out of 230 unique drugs used in COVID-19 investigational studies were selected for chemical profiling after filtering out similar drugs (i.e., only differences in formulation) and antibodies or small peptides. To analyse the chemical feature space, we employed chemical descriptors, structural analysis, and fingerprint-based approaches. We started compound analysis from a medicinal chemistry perspective (e.g., calculated partition coefficient - CLogP, molecular weight - MW, topological polar surface area - TSPA, etc.) to gain important insights about any biases in the data, such as lipophilicity or hydrophilicity. As can be seen, compounds

showed diverse characteristics (Fig. 6); however, no specific correlation patterns could be identified for pairwise comparisons aside from the expected physicochemical relationships, e.g., MW vs C atom count. Such analyses provide initial glimpses into any emerging patterns that could be explained by linear dependencies and also help to evaluate any outliers or composition biases. We continued this analysis by performing cross-compound similarity evaluation using Morgan fingerprints (nBits=2048, radius=2) and Tanimoto similarity scores (Fig. 7) (58,100). Surprisingly, most of the compounds showed only borderline similarity fluctuating around 0.2 and just the steroid group stood out with higher similarity scores forming a cluster. Other smaller groups can also be identified, e.g., angiotensin receptor blockers (Fig. 7). However, only cluster 2 (Fig. 3) drugs show clear links between network and chemical features (Fig. 7). Other categories are not only more dispersed (with partitioned clusters) but also show very little overall similarity. Overall, the assessment underlined specific fragments or structural elements, such as heterocycles, fused ring structures, and/or amphiphatic groups, as a few chemical features influencing the observed similarity across the analysed compounds.

The network-based representation of the SARS-CoV-2 infection has also found support in other studies searching for a framework to evaluate specific clinical outcomes (101–103). We, however, add to this proposition by bridging biological, pharmacological, and chemical spaces where searching for privileged structures to treat COVID-19 might involve a significantly larger chemical space and more variation than currently is considered (101–103). We used cluster 5 as our case study to further evaluate what chemical features exist for these compounds and if we could use that information to search for new drugs or design alternative compounds.

To explore cluster 5 compounds (Fig. 3, Suppl. Table 1 and 2) that were split into smaller groups based on chemical similarity in depth (Fig. 7), we isolated these compounds and assessed them as a separate set (Suppl. Fig. 7). Cluster 5 compounds share little similarity with the highest similarity score being 0.43 (Tanimoto similarity) between prasugrel and cenicriviroc. These drugs have several features contributing to the similarity where, for example, heterocycles and substituted benzenes, are among major elements linking these two groups. On the other hand, most other compounds from the cluster have marginal similarity. Another example of low similarity and different chemical structures can be found in the comparison between dexmedetomidine (used as an anxiolytic, sedative, and pain management drug with ability to provide sedation without risk of respiratory depression) and melatonin (sleep-wake cycle regulating hormone) (Tanimoto similarity=0.09) (Suppl. Fig. 7) (104,105). These observations underline why selecting pharmacological management options cannot rely on the chemical similarity of drugs alone or just main known targets because pharmacological engagement depends on multiple direct and indirect effects contributing to the cellular and organism level response (Suppl. Table 2).

Machine and deep learning based QSAR models underlined the limitations in current chemical libraries used for drug design

We began building a QSAR model by first mining the ChEMBL database (>2.1 M compounds) to investigate if cluster 5 compounds had any matches based on similarity (49). To mine the database, we set the Tanimoto similarity threshold for >0.4 based on earlier observations of the existing heterogeneity within COVID-19 compounds (Fig. 7). This assessment returned various compounds that matched specific reference drugs from cluster 5 (Suppl. Table 3). Such findings demonstrate that while similarity-based search might lead to identifying more drugs, this may not be the most optimal strategy as more complex chemical relationships need to be established when searching for active compounds. Specifically, the earlier chemical space analysis motivated us to explore compound chemical characteristics not limited to similarity but relying on the drug's 'architecture' features (58,100,106).

We first built a machine learning model (LightGBM) with a gradient boosting framework to take advantage of the tree-based learning algorithm for complex classification tasks (7,21,106–108). In order to develop this model, we needed a balanced dataset representing active and inactive compounds. A curated set of known antivirals and/or compounds resembling antivirals (COVID-19 CAS) was used to build a reference compound set for the expected actives (n=48,876) (50). To prepare the inactives (n=50,000), we opted to randomly search ChEMBL database and select the least similar compounds when compared to the actives (<0.2 Tanimoto similarity score) (49). Chemical characteristics analysis (Suppl. Fig. 8 and Suppl. Fig. 9) confirmed that the selected groups are diverse without any noticeable biases in composition. We set aside 20% of combined data for the model evaluation. Each compound was prepared as a vectorised representation of Morgan fingerprints (nBits=2048, radius=3) to represent the chemical features which we reasoned captured both structural and composition elements. The model showed 96.77% accuracy without any marked overfitting and successfully classified the test compounds as active and inactive. Applying the model to the original dataset of 158 compounds in COVID-19 clinical trials, 13 drugs were predicted to have antiviral activity (Suppl. Fig. 10). Some known antiviral drugs, such as ritonavir or maraviroc, were also included in the returned group. Other interesting therapeutic options included menthol which has been suggested to have anti-inflammatory and antiviral properties (109) and amantadine which was shown to block the ion channel activity of Protein E from SARS-CoV-2 (a conserved viroporin among coronaviruses) (110). Surprisingly, when we used the developed QSAR model to test experimentally validated antivirals against the COVID-19 M^{pro} protein (37), none of the compounds were classified to possess any activity against COVID-19.

We developed a deep learning neural net to perform the same classification using a different type of compound feature encoding (Mol2vec) to mitigate any effects stemming from the model itself or compound preparation (58). The training set-up was again a randomised selection of compounds setting aside 20% to monitor the performance of the neural network and inspect for under-/over-fitting. The model reached 96.48% accuracy after 200 epochs and returned 10 compounds from the COVID-19 clinical trial drug set as active against the virus. The compounds matched the ML model for decitabine, maraviroc, ritonavir, ribavirin, amantadine, baricitinib, hydroxychloroquine, etoposide, and cobicistat (Suppl. Fig. 11). Again, the M^{pro} experimental dataset was returned as inactive.

These interesting findings point to the fact that compounds expected to be effective against the SARS-CoV-2 virus based on searching known antivirals and/or similar compounds do not represent a complete and relevant chemical space. This might also explain why identifying new therapeutic strategies for COVID-19 drug repurposing has been difficult (111). Such observations highlight the need to combine both *in silico* analytics and experimental data to retrieve valuable compounds.

Finally, we also investigated the M^{pro} experimental dataset which included a range of compounds with varying similarity (Suppl. Fig. 12). We mined the ChEMBL database to search for compounds that showed a Tanimoto similarity higher than 0.6 for M^{pro} molecules based on fingerprint comparisons. This returned several compounds and we then inspected their maximum common substructures (MCS) to evaluate any shared features (Suppl. Fig. 12). Interestingly, the experimental compound, namely Z1220452176, contained a functionalised indole ring which matched Melatonin. Melatonin has been included in COVID-19 clinical trials and while two different QSAR models did not predict it to be antiviral compound, it clearly shares a large substructure with the experimental compound (112).

Discussion

The global spread of the SARS-CoV-2 virus resulted in a fast-evolving pandemic which prompted researchers and clinicians to investigate many different therapeutic avenues to combat the emerging healthcare crisis. While this created an opportunity to take advantage of drug repurposing strategies, the large number of ongoing studies revealed pervasive issues in clinical research (1,10,18,22). Specifically, the lack of accessible and organised data to effectively compare clinical protocols and experimental studies as well as missing updates on clinical trial outcomes create transparency problems. Consequently, this may negatively impact the research and meta-analyses in this field (22). Such trends also highlight that we need to have a better preparedness for future infectious diseases where rapid information sharing might be critical.

With our study we aimed not only to assess the existing clinical trial data but also to expand the available information so that new perspectives could be applied in clinical studies and therapeutic decision making. We based our analyses on computational pharmacology and systems biology principles to understand the networks that drugs modulate beyond a single main target (Fig. 1) (24,113). Moreover, we reasoned that focusing on a single target limits our understanding about broader systemic effects and how to effectively select combination therapeutic regimens. We also wanted to bridge the interactome with the chemical compound space and explore shared similarities and emerging patterns as combined information could prove to be useful when developing repurposing strategies further (6,10).

Clinical study and data organisation issues complicate drug repurposing

We began our analysis by exploring the current status of the COVID-19 clinical trial data based on drug clinical phases, drug mode of action, and known main targets. COVID-19 infection management options span multiple clinical phases and drug pharmacological action (Fig. 2). In combination, the results suggest that many different therapeutic avenues have been and are currently being explored to combat the infection as well as the associated complications. However, the overall clinical trial organisation and monitoring come with evidence gaps as there are no systematic data collection and verification, with some reports missing or not harmonised across different databases (22). Moreover, across academic reports and various databases, drug profiling might reflect different formulations, while in other cases this information is not exclusively reported. Consequently, the lack of an organised approach to monitor clinical trials and systemically collect data hinders repurposing and/or new compound development. These shortcomings not only reflect the current challenges but also call to take action and improve our data collection approaches. Specifically, by creating a unified method to combine clinical and academic data, we can significantly improve our forecasting and analytical capabilities which might be very important in various clinical areas, such as drug repurposing, new drug development, and preparing for other epidemics (5,6,10,13).

From a single target to network-centric pharmacology

As we focused on a selected set of compounds (Open Targets, 230 unique drugs with a known target status) used to treat or investigated for the treatment of COVID-19, we explored how their known main targets can be used to get insights into shared action and/or help predict side effects (Suppl. Fig. 2) (6,9). This led to several interesting findings.

First, our target-focused comparison revealed that while some drugs, e.g., dipyridamole and pentoxifylline, share a number of key targets, their differences in other targets might point to

potential side effects. We used dipyridamole and pentoxifylline as a case study since these pharmaceuticals engage a broad spectrum of PDEs. Moreover, multiple studies have demonstrated the importance of phosphodiesterases in regulating various cellular processes (25,69,71,114–116). PDE1A stands out from the rest of the PDE family with earlier research suggesting that this enzyme plays a role in myofibroblasts formation (117). Since PDE1A preferentially hydrolyses cyclic guanosine monophosphate (cGMP), which regulates a variety of cellular responses, including proliferation, transformation, extracellular matrix expression, apoptosis, and vascular tone, it makes sense that under a severe immune challenge curtailing abnormal pro-fibrotic processes might help preserve multiple tissue functions (117). Support for this also comes from a PDE5A inhibition study (with sildenafil) where the inhibition has been shown to reduce cardiac hypertrophy, adverse remodelling, as well as cardiac inflammation and apoptosis in the hypertensive heart (118). When comparing dipyridamole and pentoxifylline action there are some additional main targets where these compounds differ and what could be used as a therapeutic guidance. Dipyridamole inhibits the ENT1 that serves as a sodium-independent transporter for purine and pyrimidine nucleosides (70,115). Since adenosine is known to contribute to the pathophysiology of respiratory disease where adenosine challenge can lead to bronchospasm and dyspnoea, adenosine clearance can be therapeutically beneficial (115). ENT1 facilitates the removal of this nucleoside from the extracellular environment, thus terminating its action. Consequently, inhibition of ENT1 can lead to a rapid spike in extracellular adenosine concentration and increased adenosine receptor signalling. While dipyridamole has been suggested as a therapeutic option for COVID-19 with multiple beneficial properties and clinical trials are on-going (70,71), it might prove to be advantageous to consider the side effects when selecting broad spectrum therapeutics. Specifically, higher adenosine concentration has been documented to lead to bronchoconstriction and dyspnoea in asthmatic or chronic pulmonary disease (COPD) patients (119). Pentoxifylline is also known to act as an immunomodulator with anti-inflammatory properties. The 5'-nucleotidase inhibition of pentoxifylline leads to the reduced production of adenosine and inosine from their monophosphate forms. Through a nonselective phosphodiesterase inhibition as well as A2AR-mediated pathways, this drug downregulates the expression of tumor necrosis factor alpha (TNF α), IL-1, IL-6 and interferon gamma (IFN γ) (25,120). Current evidence also suggests that pentoxifylline downregulates the A2AR pathway where this therapeutic multi-modulatory action could protect from adenosine receptor overactivity (25,119,120). Thus, pentoxifylline has been suggested as a repurposing candidate to reduce tissue damage during the cytokine storm resulting from the SARS-CoV-2 infection (25). Considering the clinical data and based on the target-centric analysis, this drug appears to have more clinical benefits in comparison to dipyridamole. Overall, PDEs modulators can prove to be useful in regulating multiple cellular functions and minimising tissue

damage during uncontrolled or prolonged immune responses. Varying drug specificity towards PDEs also allows more flexibility in clinical approaches; for example, in contrast to other PDE inhibitors, apremilast offers a specific inhibition of PDE4 (114)(Suppl. Fig. 2). The lessons learnt could be applied to other similar infections, especially employing network-based assessments for drug actions.

Another key observation was that drugs with limited sets of known main targets, such as propofol, sevoflurane, isoflurane, cyclooxygenase inhibitors, or corticosteroids, reduce our ability to infer therapeutic and off-target effects. This likely explains why it has been difficult to draw conclusions about the efficacy of some of these drugs in COVID-19 treatment, e.g., in the case of corticosteroid or acetaminophen (paracetamol) use (121,122). The majority of the studies for non-steroidal anti-inflammatory agents (NSAIDs) did not indicate any associations between their use and increased mortality rates or an increased risk for respiratory failure during COVID-19 and thus, NSAID use is supported to manage COVID-19 symptoms, such as fever or muscle pain (121,123,124). There are reports where acetaminophen (paracetamol) was found to be linked with worse outcomes; yet other case studies do not report any significant differences between clinical outcomes for paracetamol or ibuprofen users (125,126). Corticosteroid treatment has also been linked to IL-6 levels where one study showed that alveolar macrophages, endothelial cells, and smooth muscle cells co-express NR3C1 and IL-6, implicating a potential corticosteroid role in cytokine release storm (127). However, corticosteroid use has diverging support as systemic studies are lacking and data collected from meta-analyses does not allow to account for all patient subgroups (122,128).

We appreciate that the more information we have about known drug targets, the better differentiation and selection of drugs can be achieved (24). Thus, until the compounds used in treating COVID-19 have extensive studies on their other potential targets, we must to rely on data mining to understand the larger modulation potential of each and every drug.

To compensate for these limitations, we propose to use an extended interactome analysis where drug main target or targets are used to search for close interactors and establish network-centric characteristics of a drug. For drugs in COVID-19 clinical trials, we identified several clusters that have multiple interactions where clusters 1, 2, and 3 were found to be the least diverse and two remaining clusters had the most diversity considering the number of seed proteins (or the main drug targets) (Fig. 3; Suppl. Table 1 and Suppl. Fig. 4). One of the most diverse clusters – cluster 4, contains drugs used to treat hypertension where clustered pharmaceuticals mostly belong to the networks of the same size. One drug from this cluster, namely losartan, acts through angiotensin-converting enzyme (ACE), angiotensin II (Ang II), and AT1 or ACE–Ang II–AT1 axis in the renin–angiotensin system (RAS) which is a known molecular pathway for end-organ fibrosis (78).

However, clinical studies focusing on another blocker, namely valsartan, did not report any significant benefits (80). Thus, comparing drugs based on their network similarity could aid in planning clinical trials. A good case example of identifying diverging biochemical processes can also be found analysing the extended network targets for ARBs and cetirizine (an antihistamine drug) which has an interesting gene uniquely belonging to a cetirizine cluster. Histidine decarboxylase (HDC) is a member of the group II decarboxylase family and converts L-histidine to histamine in a pyridoxal phosphate dependent manner. This histamine producing enzyme is known to be induced at inflammatory sites of both allergic and non-allergic reactions. Since histamine regulates various physiologic processes, including neurotransmission, gastric acid secretion, smooth muscle tone, and inflammation, its regulatory pathways can prove to be therapeutically valuable (129,130). In addition, cetirizine and other antihistamines have been tested in COVID-19 clinical trials and showed beneficial outcomes (129,131). Consequently, exploring therapeutics use from a network-centric perspective can reveal new therapeutic targets based on known drug efficacy and comparative studies. This information could be used either to repurpose or develop new drugs.

Integrating clinical studies, patient evaluation, and advanced *omics* analyses can enable pooling and assessing clinical readouts which could make not only mono-therapies but also combination treatment more effective (5–7). Moreover, the latter therapeutic strategy might offer a better pharmacological intervention ensuring a targeted pathway modulation across multiple effectors to avoid disbalanced responses. Such views are gaining more support; that is, even though certain therapies have shown benefit in subsets of the treatment population, the complexity of the viral infection underscores combination therapy usefulness in increasing treatment efficacy (10). We discussed an example of ambrisentan - a selective ET-A receptor antagonist that has been used in combination with SGLT-2 inhibition to treat COVID-19 (81). Since endothelin is a potent vasoconstrictor with pro-inflammatory and atherosclerotic action, selectively inhibiting ET-A receptor can be expected to improve pulmonary haemodynamics and oxygenation as well as reduce tissue injury. Ambrisentan's high potency and selectivity towards ET-A (4000 times greater affinity for the ET-A versus ET-B receptors) have been hypothesised to mitigate adverse effects through ET-A receptors, while preserving the potential beneficial vasodilatation of NO and prostacyclin which release is mediated by ET-B receptors on vascular endothelial cells (81). To attenuate the injurious effects of COVID-19, concomitant SGLT-2 inhibition with dapagliflozin may also prove effective to reduce inflammatory cytokines and improve endothelial function as well as cardiovascular haemodynamics (81). As a drug class, SGLT-2 inhibitors depend on blood glucose concentration and kidney function since their action of lowering blood glucose levels is achieved via the kidney independently of insulin secretion and sensitivity status. Thus, SGLT-2 inhibition would be expected to reduce inflammation and improve in glucose homeostasis, cellular

metabolism, endothelial function, and cardiovascular haemodynamics. Dapagliflozin and ambrisentan networks demonstrate well that while classical pharmacological assessment depends on searching protocols and comparing data on a single target (or several known main targets), we can benefit from computational pharmacology-oriented data mining and algorithmic evaluations. In other words, network-centric approaches allow to simultaneously explore multiple targets and their associations across the network of interest (19,24,103). Seeing the value in the extended network analyses encouraged us to explore further if we could associate specific clusters with cellular processes and pathways as a way to capture specific cluster features and enrich the analytical space.

Enrichment analysis offers new pharmacological insights

Cluster enrichment analysis not only revealed specific cellular processes based on the cluster's gene composition but also uncovered shared similarities (Fig.4; Suppl. Fig. 5). For example, some genes from cluster 5 are linked to the ERK cascade which has been suggested as a potential therapeutic target in coronavirus infections (89). Calcium homeostasis also appears to be affected by the extended network genes in cluster 5 (Fig. 4) and since calcium-linked cellular processes have been implicated in various COVID-19 outcomes, this could be helpful in narrowing down specific clinical strategies (132,133). Less diverse clusters in terms of their seed proteins, namely clusters 1 and 2, also help to illustrate that even a small number of main targets can be very important in their process modulation because of their interactome size (Fig. 3). We investigated whether the ontology-based process exploration can be mapped onto pathways. Using cluster 5 as a case study we identified diverse patterns for genes in the cluster. Some genes were mostly shared between several pathways, while others dominated across many signalling events. Cenicriviroc served as a good example for a broad modulator where several members of the extended network belong to the IL-10 signalling pathway. The extensive regulatory potential of cenicriviroc has been taken advantage of in clinical management of COVID-19 (93). A different drug from the same cluster, namely HuMax-IL8 (experimental antibody inhibiting human IL-8), shares quite a few network members with cenicriviroc. IL-8 is a pro-inflammatory cytokine involved in neutrophil activation and has been linked to the COVID-19 pathogenesis. In addition, as SARS CoV-2 has led to an increase in complications including Acute Respiratory Distress Syndrome (ARDS), the crucial role of IL-8 in lung inflammation has been suggested as a possible new therapeutic target to modulate the hyper-inflammatory response in ARDS (134,135). Expanding this analysis with computational pharmacology could greatly increase research translational potential and help identify new therapeutic regimens, especially since ARDS has limited therapeutic options (135). Furthermore, some drugs, e.g., icatibant or ozanimod, appear to show a potential pathway overlap through shared network targets and such information could also be very helpful in clinical decisions. Specifically,

such comparative analyses can help with not only off-target prediction but also finding new therapeutic combinations to manage acute and chronic disease progression. As a result, our exploration of the drug associated interactome reveals how critical it is to understand the broader pharmacological network of a drug. Such integrative analyses could help prioritise therapeutic repurposing and even predict unwanted outcomes. For example, hydroxychloroquine, after various clinical trials, was found not to show beneficial action and strong recommendations were issued against the drug's inclusion in clinical protocols (136). Our analysis revealed that hydroxychloroquine did not have many shared targets in the extended network (Fig. 3) and exploring common targets could have helped predict some off-target effects or optimise treatment for the most suitable patient groups. For example, hydroxychloroquine shares several extended network nodes with celecoxib (a NSAID) which has known cardiotoxic effects and similar issues were found for hydroxychloroquine in clinical trials (137). Understanding drug combination use is also integral in the intensive care settings where patients are treated with many pharmaceuticals at once. It has been reported that between 46 and 90% of patients admitted to the intensive care unit (ICU) are exposed to potential drug-drug interactions (138).

miRNAs open new possibilities for therapeutics investigation

The interactome is only one aspect of the complex cellular features. We suggest a new clinically valuable avenue of miRNAs as biomarkers or even therapeutic targets in COVID-19. We used the most diverse cluster 5 (Fig. 3) to explore the non-coding regulatory layer by mining the data of validated miRNAs and their regulated genes. We identified a rich network of miRNAs known to be involved in the regulation of multiple genes from cluster 5 (Suppl. Fig. 6). Evidently, miRNAs have multiple pleiotropic effects and could be used as therapeutic targets. Specifically, a drug or combination therapy could be used to influence this regulome layer by inducing expression or suppression of miRNAs to achieve a clinical effect (139). While reports are limited on the miRNA role in COVID-19, it is possible to appreciate their potentially significant function (31,99,140). As a result, using compound target and/or interactome data we can extrapolate miRNA involvement and use that to guide therapeutic decisions or disease monitoring. It is also necessary to stress that seeking only significantly changed expression levels of miRNAs might lead to missing important clues for the overall pathways and processes that may be linked through miRNAs. This is also exemplified by miR-150 which lacks noticeable changes in people with COVID-19 but still regulates and interlinks a number of genes (Suppl. Fig. 6) (141).

Computational pharmacology allows bridging the chemical and biological space

Computational pharmacology analyses and disease-associated process modelling/investigation can be further enriched by exploring the medicinal chemistry space and linking functional parameters

with drug chemical characteristics. Such analyses could significantly improve our therapeutic evaluation strategies and also aid in uncovering broader trends leading to specific pharmaceutical action. As we profiled chemical features of compounds in COVID-19 investigational studies, we immediately identified a high heterogeneity across all pharmaceuticals (Fig. 7). Moreover, there was very little overlap between pharmacological action and physicochemical features with the exceptions being the steroid and angiotensin receptor blocker groups. This prompted us to theorise that a high similarity is not necessary for the COVID-19 drugs as the modulation of signalling pathways and/or biological processes is most likely achieved through different interactome nodes. This not only means that a broader chemical space can be considered as a therapeutic option but also that repurposing strategies should take into account the interactome of a drug (101–103). Importantly, such comparative analysis could offer a better combination therapy selection and a more robust approach towards clinical and repurposing studies (5,6).

Machine and deep learning have been shown to be highly effective when identifying lead compounds. Compound fingerprinting and graph convolution principles are employed to build neural networks; however, these methods do come with shortcomings depending on the neural architecture itself and the available data (6,12). We also saw the dependence on chemical training data in our QSAR model designed to predict antiviral compound properties that could guide the selection of COVID-19 antivirals (142). While we built two different QSAR models (machine learning: LightGBM and a deep learning neural network) both of which relied on different compound feature encoding, none of the experimentally tested antivirals against COVID-19 M^{pro} protein were identified to have antiviral properties. This contrasted with the results for the COVID-19 clinical trial dataset as both QSAR models were successful in selecting drugs with predicted antiviral properties against COVID-19. Some of these identified drugs are known antiviral therapeutics. We reasoned that these discrepancies underscore the still limited chemical space for COVID-19. As modelling required a large number of molecules to train, we relied on COVID-19 CAS data (containing known and/or predicted antivirals) (50). Consequently, it is likely that enriching the compound set and also including experimental data would significantly improve our ability to identify new compounds and avoid biases. Moreover, exploring a wider chemical feature space (not limited to similar compounds target- and structure-wise) could potentially lead to discovering broad action compounds where multi-action profile could help in managing more aspects of inflammation and reduce tissue damage.

Overall, compared to *de novo* drug discovery, repurposing can be an attractive option. This is because of the significantly lower development risks since drugs have established safety and pharmacological profiles allowing a direct entry into phase II clinical trials. Accelerated therapeutic

translation, cost reduction, and possibility to explore combination therapies make therapeutic repurposing especially desirable (21,27). Despite the obvious advantages of drug repurposing, there are several major pitfalls. For example, the target population might differ significantly from the one that was involved in the original drug clinical trials. In addition, our understanding about targets and the interactome might be limited which further complicates repurposing (6). Seeing this and some recent controversies, e.g., hydroxychloroquine use, we propose an integrative approach that relies on computational pharmacology, systems biology, medicinal as well as computational chemistry to efficiently evaluate potential therapeutic candidates. A comprehensive strategy is critical for the identification of COVID-19 therapeutic solutions and future drug repurposing (20,143).

Chakraborty and colleagues provide an excellent summary of the conflicting results in COVID-19 clinical trials which accentuates the need for better global strategies in clinical trials and epidemiological meta-analyses (10,22). Finally, the issue is not just COVID-19, we need to have a better preparedness for future pandemics and also learn how to use the existing data to advance treatments for other diseases.

Conclusion

New compound development and drug repurposing need to incorporate diverse compound screening libraries with a strong understanding of their interactome and regulome. Importantly, employing computational pharmacology, data mining, systems biology methods, and computational chemistry can greatly advance our efforts in identifying the key targets and their affected networks. Our study revealed that formulating optimal pharmacological intervention options should rely on integrative approaches. We explored not only the current trends and shortcomings in COVID-19 drug repurposing, but also demonstrated the value of new perspectives using computational pharmacology and cheminformatics principles. The introduced in-depth analysis revealed the importance of expanding clinical studies beyond direct drug-target interactions and considering a more complex space of the affected networks. We also showed a number of interactions and pathways that could be exploited when considering combination therapies. The findings of miRNAs networks offer a new strategy to search for valuable biomarkers or therapeutic management options. Even though computational modelling is a powerful tool in prioritising compounds, it is important to include their biological action, experimental results, and extended network data to build better predictors. We demonstrated that the chemical space for COVID-19 investigational compounds might not be broad enough and could benefit from additional experimental evidence to create more robust models. Despite limitations in available data, it is still possible to extract valuable information that could potentially save time and resources by helping to better prioritise compounds for *in vitro* screening. Finally, we strongly advocate for taking this opportunity to establish

comprehensive practices for today's and future infectious diseases by preparing solid analytical frameworks.

Bibliography

1. Chauhan S. Comprehensive review of coronavirus disease 2019 (COVID-19). *Biomed J* [Internet]. 2020 Aug 1 [cited 2022 Jan 3];43(4):334–40. Available from: <https://pubmed.ncbi.nlm.nih.gov/32788071/>
2. Serafin MB, Bottega A, Foletto VS, da Rosa TF, Hörner A, Hörner R. Drug repositioning is an alternative for the treatment of coronavirus COVID-19. *Int J Antimicrob Agents* [Internet]. 2020 Jun 1 [cited 2022 Jan 22];55(6):105969. Available from: </pmc/articles/PMC7194941/>
3. Cascella M, Rajnik M, Cuomo A, Dulebohn SC, Di Napoli R. Features, Evaluation, and Treatment of Coronavirus (COVID-19). *StatPearls* [Internet]. 2021 Sep 2 [cited 2022 Jan 3]; Available from: <https://www.ncbi.nlm.nih.gov/books/NBK554776/>
4. Gebhard C, Regitz-Zagrosek V, Neuhauser HK, Morgan R, Klein SL. Impact of sex and gender on COVID-19 outcomes in Europe. *Biol Sex Differ* [Internet]. 2020 May 25 [cited 2022 Jan 3];11(1). Available from: <https://pubmed.ncbi.nlm.nih.gov/32450906/>
5. Akinbolade S, Diarmuid Coughlan |, Fairbairn R, Mcconkey G, Powell H, Ogunbayo D, et al. Combination therapies for COVID-19: An overview of the clinical trials landscape. *Br J Clin Pharmacol* [Internet]. 2021 Oct 17 [cited 2022 Jan 16]; Available from: <https://onlinelibrary.wiley.com/doi/full/10.1111/bcp.15089>
6. Ng YL, Salim CK, Chu JJH. Drug repurposing for COVID-19: Approaches, challenges and promising candidates. *Pharmacol Ther* [Internet]. 2021 Dec 1 [cited 2022 Feb 6];228:107930. Available from: </pmc/articles/PMC8220862/>
7. Jarada TN, Rokne JG, Alhaji R. A review of computational drug repositioning: Strategies, approaches, opportunities, challenges, and directions. *J Cheminform* [Internet]. 2020 Jul 22 [cited 2022 Jan 16];12(1):1–23. Available from: <https://jcheminf.biomedcentral.com/articles/10.1186/s13321-020-00450-7>
8. Ledford H. Dozens of coronavirus drugs are in development - what happens next? *Nature*. 2020 May 1;581(7808):247–8.
9. Pushpakom S, Iorio F, Eyers PA, Escott KJ, Hopper S, Wells A, et al. Drug repurposing: progress, challenges and recommendations. *Nat Rev Drug Discov* 2018 181 [Internet]. 2018 Oct 12 [cited 2022 Jan 4];18(1):41–58. Available from: <https://www.nature.com/articles/nrd.2018.168>
10. Chakraborty C, Sharma AR, Bhattacharya M, Agoramoorthy G, Lee S-S. The Drug Repurposing for COVID-19 Clinical Trials Provide Very Effective Therapeutic

- Combinations: Lessons Learned From Major Clinical Studies. *Front Pharmacol* [Internet]. 2021 Nov 18 [cited 2022 Jan 4];12. Available from: [/pmc/articles/PMC8636940/](#)
11. Choudhury C. Fragment tailoring strategy to design novel chemical entities as potential binders of novel corona virus main protease. *J Biomol Struct Dyn* [Internet]. 2020 [cited 2021 Apr 13];1. Available from: [/pmc/articles/PMC7284137/](#)
 12. Yu R, Chen L, Lan R, Shen R, Li P. Computational screening of antagonists against the SARS-CoV-2 (COVID-19) coronavirus by molecular docking. *Int J Antimicrob Agents*. 2020 Aug 1;56(2):106012.
 13. Bharti R, Shukla SK. Molecules against Covid-19: An in silico approach for drug development. *J Electron Sci Technol*. 2021 Mar 1;19(1):100095.
 14. Mao J, Akhtar J, Zhang X, Sun L, Guan S, Li X, et al. Comprehensive strategies of machine-learning-based quantitative structure-activity relationship models. *iScience*. 2021 Sep 24;24(9):103052.
 15. Tsou LK, Yeh SH, Ueng SH, Chang CP, Song JS, Wu MH, et al. Comparative study between deep learning and QSAR classifications for TNBC inhibitors and novel GPCR agonist discovery. *Sci Reports* 2020 101 [Internet]. 2020 Oct 8 [cited 2021 Dec 30];10(1):1–11. Available from: <https://www.nature.com/articles/s41598-020-73681-1>
 16. Gligorijević V, Pržulj N. Methods for biological data integration: perspectives and challenges. *J R Soc Interface* [Internet]. 2015 Nov 6 [cited 2021 Sep 6];12(112). Available from: <https://royalsocietypublishing.org/doi/abs/10.1098/rsif.2015.0571>
 17. McCreary EK, Meyer NJ. Covid-19 controversies: the tocilizumab chapter. *BMJ* [Internet]. 2021 Jan 27 [cited 2022 Jan 4];372. Available from: <https://pubmed.ncbi.nlm.nih.gov/33504502/>
 18. Gupta A, Malviya A. Chloroquine and hydroxychloroquine for COVID-19: time to close the chapter. *Postgrad Med J* [Internet]. 2021 Oct 1 [cited 2022 Jan 4];97(1152):676–7. Available from: <https://pubmed.ncbi.nlm.nih.gov/32788309/>
 19. Gordon DE, Jang GM, Bouhaddou M, Xu J, Obernier K, White KM, et al. A SARS-CoV-2 protein interaction map reveals targets for drug repurposing. *Nat* 2020 5837816 [Internet]. 2020 Apr 30 [cited 2021 Nov 1];583(7816):459–68. Available from: <https://www.nature.com/articles/s41586-020-2286-9>
 20. Lavecchia A, Cerchia C. In silico methods to address polypharmacology: Current status, applications and future perspectives. Vol. 21, *Drug Discovery Today*. Elsevier Ltd; 2016. p. 288–98.
 21. March-Vila E, Pinzi L, Sturm N, Tinivella A, Engkvist O, Chen H, et al. On the Integration of In Silico Drug Design Methods for Drug Repurposing. *Front Pharmacol* [Internet]. 2017 May 23 [cited 2021 Jul 10];8(MAY):298. Available from: www.frontiersin.org
 22. Rodgers F, Pepperrell T, Keestra S, Pilkington V. Missing clinical trial data: the evidence gap in primary data for potential COVID-19 drugs. *Trials* [Internet]. 2021 Dec 1 [cited 2022 Jan 4];22(1). Available from: <https://pubmed.ncbi.nlm.nih.gov/33451350/>

23. Yildirim MA, Goh K Il, Cusick ME, Barabási AL, Vidal M. Drug-target network. *Nat Biotechnol.* 2007;25(10):1119–26.
24. Hopkins, L A. Network Pharmacology. *Netw Pharmacol.* 2017;25(10):127–64.
25. Hendry BM, Stafford N, Arnold AD, Sangwaiya A, Manglam V, Rosen SD, et al. Hypothesis: Pentoxifylline is a potential cytokine modulator therapeutic in COVID-19 patients. *Pharmacol Res Perspect [Internet].* 2020 Aug 1 [cited 2022 Jan 15];8(4). Available from: /pmc/articles/PMC7383088/
26. Kanapeckaitė A, Burokienė N. Insights into therapeutic targets and biomarkers using integrated multi-'omics' approaches for dilated and ischemic cardiomyopathies. *Integr Biol (Camb) [Internet].* 2021 May 1 [cited 2021 Sep 21];13(5):121–37. Available from: <https://pubmed.ncbi.nlm.nih.gov/33969404/>
27. Zhou Y, Wang F, Tang J, Nussinov R, Cheng F. Artificial intelligence in COVID-19 drug repurposing. *Lancet Digit Heal [Internet].* 2020 Dec 1 [cited 2022 Jan 16];2(12):e667–76. Available from: <https://pubmed.ncbi.nlm.nih.gov/32984792/>
28. Malik MA, Wani MY, Hashmi AA. Combination therapy: Current status and future perspectives [Internet]. *Combination Therapy Against Multidrug Resistance.* Elsevier Inc.; 2020. 1–38 p. Available from: <https://doi.org/10.1016/B978-0-12-820576-1.00001-1>
29. Flockhart D, Bies RR, Gastonguay MR, Schwartz SL. Big Data: Challenges and opportunities for clinical pharmacology. Vol. 81, *British Journal of Clinical Pharmacology.* Blackwell Publishing Ltd; 2016. p. 804–6.
30. Farr RJ, Rootes CL, Rowntree LC, Nguyen THO, Hensen L, Kedzierski L, et al. Altered microRNA expression in COVID-19 patients enables identification of SARS-CoV-2 infection. *PLOS Pathog [Internet].* 2021 Jul 1 [cited 2022 Jan 11];17(7):e1009759. Available from: <https://journals.plos.org/plospathogens/article?id=10.1371/journal.ppat.1009759>
31. Fani M, Zandi M, Ebrahimi S, Soltani S, Abbasi S. The role of miRNAs in COVID-19 disease. *Future Virol [Internet].* 2021 Apr 1 [cited 2022 Jan 11];16(4):301–6. Available from: /pmc/articles/PMC7989380/
32. Hanna J, Hossain GS, Kocerha J. The potential for microRNA therapeutics and clinical research. *Front Genet.* 2019;10(MAY):478.
33. Bofill-De Ros X, Gu S. Guidelines for the optimal design of miRNA-based shRNAs. *Methods.* 2016 Jul 1;103:157–66.
34. Schneider HC, Klabunde T. Understanding drugs and diseases by systems biology? *Bioorg Med Chem Lett.* 2013 Mar 1;23(5):1168–76.
35. Brogi S, Ramalho TC, Kuca K, Medina-Franco JL, Valko M. Editorial: In silico Methods for Drug Design and Discovery. *Front Chem [Internet].* 2020 Aug 7 [cited 2021 Jul 10];8:612. Available from: <http://faerun.gdb.tools>
36. Du J, Guo J, Kang D, Li Z, Wang G, Wu J, et al. New techniques and strategies in drug discovery. *Chinese Chem Lett.* 2020 Jul 1;31(7):1695–708.

37. Downloads - - Diamond Light Source [Internet]. [cited 2022 Jan 11]. Available from: <https://www.diamond.ac.uk/covid-19/for-scientists/Main-protease-structure-and-XChem/Downloads.html>
38. Ekins S, Mestres J, Testa B. In silico pharmacology for drug discovery: methods for virtual ligand screening and profiling. *Br J Pharmacol* [Internet]. 2007 Sep [cited 2021 Jul 5];152(1):9. Available from: [/pmc/articles/PMC1978274/](https://pubmed.ncbi.nlm.nih.gov/1978274/)
39. Abi Hussein H, Geneix C, Petitjean M, Borrel A, Flatters D, Camproux AC. Global vision of druggability issues: applications and perspectives. Vol. 22, *Drug Discovery Today*. Elsevier Ltd; 2017. p. 404–15.
40. Koscielny G, An P, Carvalho-Silva D, Cham JA, Fumis L, Gasparyan R, et al. Open Targets: A platform for therapeutic target identification and Validation. *Nucleic Acids Res* [Internet]. 2017 Jan 1 [cited 2020 Dec 11];45(D1):D985–94. Available from: <https://www.>
41. Home - Open Targets [Internet]. [cited 2020 Dec 7]. Available from: <https://www.opentargets.org/>
42. PubChem [Internet]. [cited 2020 Dec 16]. Available from: <https://pubchem.ncbi.nlm.nih.gov/>
43. PubChem Covid-19 clinical trials [Internet]. [cited 2021 Nov 1]. Available from: [https://pubchem.ncbi.nlm.nih.gov/#tab=compound&query=covid-19 clinicaltrials](https://pubchem.ncbi.nlm.nih.gov/#tab=compound&query=covid-19%20clinicaltrials)
44. STITCH: chemical association networks [Internet]. [cited 2020 Dec 7]. Available from: <http://stitch.embl.de/>
45. Szklarczyk D, Gable AL, Lyon D, Junge A, Wyder S, Huerta-Cepas J, et al. STRING v11: Protein-protein association networks with increased coverage, supporting functional discovery in genome-wide experimental datasets. *Nucleic Acids Res*. 2019 Jan 8;47(D1):D607–13.
46. STRING: functional protein association networks [Internet]. [cited 2021 Nov 1]. Available from: <https://string-db.org/>
47. Home - Reactome Pathway Database [Internet]. [cited 2020 Dec 7]. Available from: <https://reactome.org/>
48. Kanapeckaite A. OmicInt package: Exploring omics data and regulatory networks using integrative analyses and machine learning. *Artif Intell Life Sci* [Internet]. 2021 Dec 1 [cited 2022 Jan 15];1:100025. Available from: <https://linkinghub.elsevier.com/retrieve/pii/S2667318521000258>
49. ChEMBL Database [Internet]. [cited 2021 Dec 14]. Available from: <https://www.ebi.ac.uk/chembl/>
50. Download CAS COVID-19 Antiviral Candidate Compounds Dataset | CAS [Internet]. [cited 2022 Jan 11]. Available from: <https://www.cas.org/covid-19-antiviral-compounds-dataset>
51. RStudio | Open source & professional software for data science teams - RStudio [Internet]. [cited 2020 Oct 26]. Available from: <https://rstudio.com/>

52. Bioconductor - STRINGdb [Internet]. [cited 2022 Jan 15]. Available from: <https://www.bioconductor.org/packages/release/bioc/html/STRINGdb.html>
53. Bioconductor - clusterProfiler [Internet]. [cited 2020 Dec 7]. Available from: <https://bioconductor.org/packages/release/bioc/html/clusterProfiler.html>
54. enrichGO function - RDocumentation [Internet]. [cited 2022 Aug 22]. Available from: <https://www.rdocumentation.org/packages/clusterProfiler/versions/3.0.4/topics/enrichGO>
55. enrichPathway function - RDocumentation [Internet]. [cited 2022 Aug 22]. Available from: <https://www.rdocumentation.org/packages/ReactomePA/versions/1.16.2/topics/enrichPathway>
56. Bioconductor - biomaRt [Internet]. [cited 2022 Aug 22]. Available from: <https://bioconductor.org/packages/release/bioc/html/biomaRt.html>
57. Welcome to Python.org [Internet]. [cited 2022 Aug 22]. Available from: <https://www.python.org/>
58. Jaeger S, Fulle S, Turk S. Mol2vec: Unsupervised Machine Learning Approach with Chemical Intuition. *J Chem Inf Model*. 2018;58(1):27–35.
59. RDKit [Internet]. [cited 2022 Jan 15]. Available from: <https://www.rdkit.org/>
60. NumPy [Internet]. [cited 2022 Jan 15]. Available from: <https://numpy.org/>
61. pandas - Python Data Analysis Library [Internet]. [cited 2022 Jan 15]. Available from: <https://pandas.pydata.org/>
62. Waskom M. seaborn: statistical data visualization. *J Open Source Softw*. 2021 Apr 6;6(60):3021.
63. Matplotlib — Visualization with Python [Internet]. [cited 2022 Jan 15]. Available from: <https://matplotlib.org/>
64. chemexpy · PyPI [Internet]. [cited 2022 Jan 15]. Available from: <https://pypi.org/project/chemexpy/>
65. scikit-learn: machine learning in Python — scikit-learn 1.0.2 documentation [Internet]. [cited 2022 Jan 15]. Available from: <https://scikit-learn.org/stable/>
66. lightgbm.LGBMClassifier — LightGBM 3.3.2.99 documentation [Internet]. [cited 2022 Aug 22]. Available from: <https://lightgbm.readthedocs.io/en/latest/pythonapi/lightgbm.LGBMClassifier.html>
67. TensorFlow [Internet]. [cited 2022 Jan 15]. Available from: <https://www.tensorflow.org/>
68. Keras: the Python deep learning API [Internet]. [cited 2022 Aug 22]. Available from: <https://keras.io/>
69. Nosengo N. Can you teach old drugs new tricks? *Nature* [Internet]. 2016 Jun 14 [cited 2022 Feb 6];534(7607):314–6. Available from: <https://pubmed.ncbi.nlm.nih.gov/27306171/>

70. Wang C, Lin W, Playa H, Sun S, Cameron K, Buolamwini JK. Dipyridamole analogues as pharmacological inhibitors of equilibrative nucleoside transporters. Identification of novel potent and selective inhibitors of the adenosine transporter function of human equilibrative nucleoside transporter 4 (hENT4). *Biochem Pharmacol* [Internet]. 2013 [cited 2022 Feb 7];86(11):1531–40. Available from: [/pmc/articles/PMC3866046/](#)
71. Aliter KF, Al-Horani RA. Potential Therapeutic Benefits of Dipyridamole in COVID-19 Patients. *Curr Pharm Des* [Internet]. 2021 Oct 1 [cited 2022 Jan 24];27(6):866–75. Available from: <https://pubmed.ncbi.nlm.nih.gov/33001004/>
72. Hanidziar D, Baldyga K, Ji CS, Lu J, Zheng H, Wiener-Kronish J, et al. Standard Sedation and Sedation With Isoflurane in Mechanically Ventilated Patients With Coronavirus Disease 2019. *Crit Care Explor* [Internet]. 2021 Mar 5 [cited 2022 Jan 15];3(3):e0370. Available from: https://journals.lww.com/ccejournal/Fulltext/2021/03000/Standard_Sedation_and_Sedation_With_Isoflurane_in.7.aspx
73. Witenko CJ, Littlefield AJ, Abedian S, An A, Barie PS, Berger K. The Safety of Continuous Infusion Propofol in Mechanically Ventilated Adults With Coronavirus Disease 2019. *Ann Pharmacother* [Internet]. 2022 Jan 1 [cited 2022 Jan 15];56(1):5. Available from: [/pmc/articles/PMC8127019/](#)
74. Nieuwenhuijs-Moeke GJ, Jainandunsing JS, Struys MMRF. Sevoflurane, a sigh of relief in COVID-19? *BJA Br J Anaesth* [Internet]. 2020 Aug 1 [cited 2022 Jan 15];125(2):118. Available from: [/pmc/articles/PMC7252148/](#)
75. Kaura V, Hopkins PM. Sevoflurane may not be a complete sigh of relief in COVID-19. *Br J Anaesth* [Internet]. 2020 Dec 1 [cited 2022 Jan 15];125(6):e487–8. Available from: <http://www.bjanaesthesia.org.uk/article/S0007091220307625/fulltext>
76. Agúndez JAG, Blanca M, Cornejo-García JA, García-Martín E. Pharmacogenomics of cyclooxygenases. *Pharmacogenomics* [Internet]. 2015 Apr 1 [cited 2022 Aug 25];16(5):501–22. Available from: <https://pubmed.ncbi.nlm.nih.gov/25916522/>
77. Park JH, Lee HK. Re-analysis of Single Cell Transcriptome Reveals That the NR3C1-CXCL8-Neutrophil Axis Determines the Severity of COVID-19. *Front Immunol* [Internet]. 2020 Aug 28 [cited 2021 Dec 12];11. Available from: <https://pubmed.ncbi.nlm.nih.gov/32983174/>
78. Zeinalian M, Salari-Jazi A, Jannesari A, Khanahmad H. A potential protective role of losartan against coronavirus-induced lung damage. *Infect Control Hosp Epidemiol* [Internet]. 2020 Jun 1 [cited 2022 Jan 16];41(6):1. Available from: [/pmc/articles/PMC7137531/](#)
79. Puskarich MA, Cummins NW, Ingraham NE, Wacker DA, Reilkoff RA, Driver BE, et al. A multi-center phase II randomized clinical trial of losartan on symptomatic outpatients with COVID-19. *eClinicalMedicine* [Internet]. 2021 Jul 1 [cited 2022 Jan 29];37:100957. Available from: <http://www.thelancet.com/article/S2589537021002376/fulltext>

80. de Ligt M, Hesselink MKC, Jorgensen J, Jocken JWE, Blaak EE, Goossens GH. The angiotensin II α 1 receptor blocker valsartan in the battle against COVID-19. *Obesity (Silver Spring)* [Internet]. 2021 Sep 1 [cited 2022 Jan 16];29(9):1423–6. Available from: <https://pubmed.ncbi.nlm.nih.gov/33955183/>
81. Fisk M, Althage M, Moosmang S, Greasley PJ, Cope AP, Jayne DR, et al. Endothelin antagonism and sodium glucose Co-transporter 2 inhibition. A potential combination therapeutic strategy for COVID-19. *Pulm Pharmacol Ther* [Internet]. 2021 Aug 1 [cited 2022 Jan 16];69:102035. Available from: </pmc/articles/PMC8084922/>
82. Ranucci M, Ballotta A, Di Dedda U, Bayshnikova E, Dei Poli M, Resta M, et al. The procoagulant pattern of patients with COVID-19 acute respiratory distress syndrome. *J Thromb Haemost* [Internet]. 2020 Jul 1 [cited 2022 Jan 16];18(7):1747–51. Available from: <https://pubmed.ncbi.nlm.nih.gov/32302448/>
83. Akşit E, Kırılmaz B, Gazi E, Aydın F. Ticagrelor Can Be an Important Agent in the Treatment of Severe COVID-19 Patients with Myocardial Infarction. *Balkan Med J* [Internet]. 2020 [cited 2022 Jan 16];37(4):233. Available from: </pmc/articles/PMC7285674/>
84. Kow CS, Hasan SS. The use of antiplatelet agents for arterial thromboprophylaxis in COVID-19. *Rev Esp Cardiol (Engl Ed)* [Internet]. 2021 Jan [cited 2022 Jan 16];74(1):114. Available from: </pmc/articles/PMC7455174/>
85. Choubey A, Dehury B, Kumar S, Medhi B, Mondal P. Naltrexone a potential therapeutic candidate for COVID-19. *J Biomol Struct Dyn* [Internet]. 2020 [cited 2022 Jan 16]; Available from: <https://pubmed.ncbi.nlm.nih.gov/32930058/>
86. Sullivan R, Kilaru A, Hemmer B, Campbell Cree BA, Greenberg BM, Kundu U, et al. COVID-19 Infection in Fingolimod- or Siponimod-Treated Patients: Case Series. *Neurol Neuroimmunol neuroinflammation* [Internet]. 2021 Jan [cited 2022 Jan 16];9(1):e1092. Available from: <https://pubmed.ncbi.nlm.nih.gov/34848501/>
87. Gomez-Mayordomo V, Montero-Escribano P, Matías-Guiu JA, González-García N, Porta-Etessam J, Matías-Guiu J. Clinical exacerbation of SARS-CoV2 infection after fingolimod withdrawal. *J Med Virol* [Internet]. 2021 Jan 1 [cited 2022 Jan 16];93(1):546–9. Available from: <https://pubmed.ncbi.nlm.nih.gov/32644205/>
88. Plaze M, Attali D, Petit A-C, Blatzer M, Simon-Loriere E, Vinckier F, et al. Repurposing chlorpromazine to treat COVID-19: The reCoVery study. *Encephale* [Internet]. 2020 Jun 1 [cited 2022 Jan 16];46(3):169–72. Available from: <http://www.ncbi.nlm.nih.gov/pubmed/32425222>
89. Kindrachuk J, Ork B, Hart BJ, Mazur S, Holbrook MR, Frieman MB, et al. Antiviral potential of ERK/MAPK and PI3K/AKT/mTOR signaling modulation for Middle East respiratory syndrome coronavirus infection as identified by temporal kinome analysis. *Antimicrob Agents Chemother* [Internet]. 2015 Feb 1 [cited 2021 Dec 12];59(2):1088–99. Available from: <https://pubmed.ncbi.nlm.nih.gov/25487801/>

90. Iba T, Levy JH, Levi M, Thachil J. Coagulopathy in COVID-19. *J Thromb Haemost* [Internet]. 2020 Sep 1 [cited 2022 Jan 16];18(9):2103–9. Available from: <https://pubmed.ncbi.nlm.nih.gov/32558075/>
91. Aggarwal M, Dass J, Mahapatra M. Hemostatic Abnormalities in COVID-19: An Update. *Indian J Hematol Blood Transfus* [Internet]. 2020 Oct 1 [cited 2022 Jan 16];36(4):616–26. Available from: <https://pubmed.ncbi.nlm.nih.gov/32837053/>
92. NIH begins large clinical trial to test immune modulators for treatment of COVID-19 | National Institutes of Health (NIH) [Internet]. [cited 2022 Jan 16]. Available from: <https://www.nih.gov/news-events/news-releases/nih-begins-large-clinical-trial-test-immune-modulators-treatment-covid-19>
93. Files DC, Tacke F, O’Sullivan A, Dorr P, Ferguson WG, Powderly WG. Rationale of using the dual chemokine receptor CCR2/CCR5 inhibitor cenicriviroc for the treatment of COVID-19. *PLoS Pathog* [Internet]. 2022 Jun 24 [cited 2022 Aug 26];18(6):e1010547. Available from: <https://pubmed.ncbi.nlm.nih.gov/35749425/>
94. Shaw RJ, Bradbury C, Abrams ST, Wang G, Toh CH. COVID-19 and immunothrombosis: emerging understanding and clinical management. *Br J Haematol* [Internet]. 2021 Aug 1 [cited 2022 Aug 28];194(3):518–29. Available from: <https://onlinelibrary.wiley.com/doi/full/10.1111/bjh.17664>
95. COVID-19 Therapeutics Prioritized for Testing in Clinical Trials | National Institutes of Health (NIH) [Internet]. [cited 2022 Aug 28]. Available from: <https://www.nih.gov/research-training/medical-research-initiatives/activ/covid-19-therapeutics-prioritized-testing-clinical-trials>
96. Mansour E, Bueno FF, de Lima-Júnior JC, Palma A, Monfort-Pires M, Bombassaro B, et al. Evaluation of the efficacy and safety of icatibant and C1 esterase/kallikrein inhibitor in severe COVID-19: study protocol for a three-armed randomized controlled trial. *Trials* [Internet]. 2021 Dec 1 [cited 2022 Jan 16];22(1). Available from: </pmc/articles/PMC7816150/>
97. Pérez-Jeldres T, Alvarez-Lobos M, Rivera-Nieves J. Targeting Sphingosine-1-Phosphate Signaling in Immune-Mediated Diseases: Beyond Multiple Sclerosis. *Drugs* [Internet]. 2021 Jun 1 [cited 2022 Jan 16];81(9):985–1002. Available from: <https://pubmed.ncbi.nlm.nih.gov/33983615/>
98. Duecker RP, Adam EH, Wirtz S, Gronau L, Khodamoradi Y, Eberhardt FJ, et al. The mir-320 family is strongly downregulated in patients with covid-19 induced severe respiratory failure. *Int J Mol Sci* [Internet]. 2021 Oct 1 [cited 2022 Jan 16];22(19). Available from: </pmc/articles/PMC8508658/>
99. Paul S, Bravo Vázquez LA, Reyes-Pérez PR, Estrada-Meza C, Aponte Albuquerque RA, Pathak S, et al. The role of microRNAs in solving COVID-19 puzzle from infection to therapeutics: A mini-review. *Virus Res*. 2022 Jan 15;308:198631.

100. Alvarsson J, Eklund M, Engkvist O, Spjuth O, Carlsson L, Wikberg JES, et al. Ligand-based target prediction with signature fingerprints. *J Chem Inf Model*. 2014 Oct 27;54(10):2647–53.
101. Messina F, Giombini E, Montaldo C, Sharma AA, Zoccoli A, Sekaly RP, et al. Looking for pathways related to COVID-19: confirmation of pathogenic mechanisms by SARS-CoV-2–host interactome. *Cell Death Dis* 2021 128 [Internet]. 2021 Aug 12 [cited 2022 Jan 16];12(8):1–10. Available from: <https://www.nature.com/articles/s41419-021-03881-8>
102. Farahani M, Niknam Z, Mohammadi Amirabad L, Amiri-Dashatan N, Koushki M, Nemati M, et al. Molecular pathways involved in COVID-19 and potential pathway-based therapeutic targets. *Biomed Pharmacother*. 2022 Jan 1;145:112420.
103. Nashiry A, Sarmin Sumi S, Islam S, Quinn JMW, Moni MA. Bioinformatics and system biology approach to identify the influences of COVID-19 on cardiovascular and hypertensive comorbidities. *Brief Bioinform* [Internet]. 2021 Mar 1 [cited 2022 Jan 16];22(2):1387–401. Available from: <https://pubmed.ncbi.nlm.nih.gov/33458761/>
104. Dexmedetomidine: Uses, Interactions, Mechanism of Action | DrugBank Online [Internet]. [cited 2022 Jan 16]. Available from: <https://go.drugbank.com/drugs/DB00633>
105. Melatonin: Uses, Interactions, Mechanism of Action | DrugBank Online [Internet]. [cited 2022 Jan 16]. Available from: <https://go.drugbank.com/drugs/DB01065>
106. Bajusz D, Rácz A, Héberger K. Why is Tanimoto index an appropriate choice for fingerprint-based similarity calculations? *J Cheminform* [Internet]. 2015 Dec 8 [cited 2020 Oct 19];7(1):20. Available from: <https://jcheminf.biomedcentral.com/articles/10.1186/s13321-015-0069-3>
107. Lavecchia A. Machine-learning approaches in drug discovery: Methods and applications. Vol. 20, *Drug Discovery Today*. Elsevier Ltd; 2015. p. 318–31.
108. Zhang J, Mucs D, Norinder U, Svensson F. LightGBM: An Effective and Scalable Algorithm for Prediction of Chemical Toxicity-Application to the Tox21 and Mutagenicity Data Sets. *J Chem Inf Model* [Internet]. 2019 [cited 2022 Jan 16];59(10). Available from: <https://pubmed.ncbi.nlm.nih.gov/31560206/>
109. Asif M, Saleem M, Saadullah M, Yaseen HS, Al Zarzour R. COVID-19 and therapy with essential oils having antiviral, anti-inflammatory, and immunomodulatory properties. *Inflammopharmacology* [Internet]. 2020 Oct 1 [cited 2022 Jan 16];28(5):1. Available from: <https://pubmed.ncbi.nlm.nih.gov/34853399/>
110. Toft-Bertelsen TL, Jeppesen MG, Tzortzini E, Xue K, Giller K, Becker S, et al. Amantadine has potential for the treatment of COVID-19 because it inhibits known and novel ion channels encoded by SARS-CoV-2. *Commun Biol* [Internet]. 2021 Dec [cited 2022 Jan 16];4(1). Available from: <https://pubmed.ncbi.nlm.nih.gov/34853399/>
111. Lipinski CF, Maltarollo VG, Oliveira PR, da Silva ABF, Honorio KM. Advances and Perspectives in Applying Deep Learning for Drug Design and Discovery. *Front Robot AI*. 2019 Nov 5;6:108.

112. Camp OG, Bai D, Gonullu DC, Nayak N, Abu-Soud HM. Melatonin interferes with COVID-19 at several distinct ROS-related steps. *J Inorg Biochem*. 2021 Oct 1;223:111546.
113. Earm K, Earm YE. Integrative approach in the era of failing drug discovery and development. *Integr Med Res* [Internet]. 2014 Dec [cited 2021 Jul 28];3(4):211. Available from: [/pmc/articles/PMC5481768/](https://pubmed.ncbi.nlm.nih.gov/3481768/)
114. Schett G, Sloan VS, Stevens RM, Schafer P. Apremilast: A Novel PDE4 Inhibitor in the Treatment of Autoimmune and Inflammatory Diseases. *Ther Adv Musculoskelet Dis* [Internet]. 2010 [cited 2022 Jan 24];2(5):271. Available from: [/pmc/articles/PMC3383510/](https://pubmed.ncbi.nlm.nih.gov/3383510/)
115. Rosenbrier Ribeiro L, Ian Storer R. A semi-quantitative translational pharmacology analysis to understand the relationship between in vitro ENT1 inhibition and the clinical incidence of dyspnoea and bronchospasm. *Toxicol Appl Pharmacol* [Internet]. 2017 Feb 15 [cited 2022 Jan 24];317:41–50. Available from: <https://pubmed.ncbi.nlm.nih.gov/28041785/>
116. Dipyridamole to Prevent Coronavirus Exacerbation of Respiratory Status (DICER) in COVID-19 - Full Text View - ClinicalTrials.gov [Internet]. [cited 2022 Jan 24]. Available from: <https://clinicaltrials.gov/ct2/show/NCT04391179>
117. Zhou HY, Chen WD, Zhu DL, Wu LY, Zhang J, Han WQ, et al. The PDE1A-PKC α Signaling Pathway Is Involved in the Upregulation of α -Smooth Muscle Actin by TGF- β 1 in Adventitial Fibroblasts. *J Vasc Res* [Internet]. 2010 Jan [cited 2021 Dec 12];47(1):9. Available from: [/pmc/articles/PMC2855283/](https://pubmed.ncbi.nlm.nih.gov/2855283/)
118. Westermann D, Becher PM, Lindner D, Savvatis K, Xia Y, Fröhlich M, et al. Selective PDE5A inhibition with sildenafil rescues left ventricular dysfunction, inflammatory immune response and cardiac remodeling in angiotensin II-induced heart failure in vivo. *Basic Res Cardiol* [Internet]. 2012 Nov 1 [cited 2021 Dec 12];107(6). Available from: <https://pubmed.ncbi.nlm.nih.gov/23117837/>
119. Brown RA, Spina D, Page CP. Adenosine receptors and asthma. *Br J Pharmacol* [Internet]. 2008 Mar 1 [cited 2022 Jan 24];153(S1):S446–56. Available from: <https://onlinelibrary.wiley.com/doi/full/10.1038/bjp.2008.22>
120. Konrad FM, Neudeck G, Vollmer I, Ngamsri KC, Thiel M, Reutershan J. Protective effects of pentoxifylline in pulmonary inflammation are adenosine receptor A2A dependent. *FASEB J* [Internet]. 2013 Sep [cited 2022 Jan 24];27(9):3524–35. Available from: <https://pubmed.ncbi.nlm.nih.gov/23699177/>
121. Pergolizzi J V., Varrassi G, Magnusson P, LeQuang JA, Paladini A, Taylor R, et al. COVID-19 and NSAIDs: A Narrative Review of Knowns and Unknowns. *Pain Ther* [Internet]. 2020 Dec 1 [cited 2022 Jan 15];9(2):353–8. Available from: <https://pubmed.ncbi.nlm.nih.gov/32447629/>
122. Wagner C, Griesel M, Mikolajewska A, Mueller A, Nothacker M, Kley K, et al. Systemic corticosteroids for the treatment of COVID-19. *Cochrane database Syst Rev* [Internet]. 2021 Aug 16 [cited 2022 Jan 16];8(8). Available from: <https://pubmed.ncbi.nlm.nih.gov/34396514/>

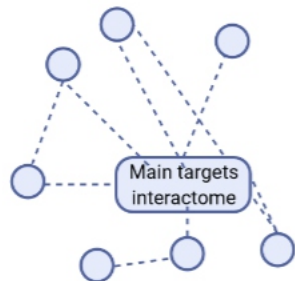
123. Park J, Lee SH, You SC, Kim J, Yang K. Non-steroidal anti-inflammatory agent use may not be associated with mortality of coronavirus disease 19. *Sci Reports* 2021 111 [Internet]. 2021 Mar 3 [cited 2022 Jan 15];11(1):1–7. Available from: <https://www.nature.com/articles/s41598-021-84539-5>
124. Poutoglidou F, Saitis A, Kouvelas D. Ibuprofen and COVID-19 disease: separating the myths from facts. *Expert Rev Respir Med* [Internet]. 2021 [cited 2022 Jan 15];15(8):979–83. Available from: <https://pubmed.ncbi.nlm.nih.gov/34196258/>
125. Rinott E, Kozer E, Shapira Y, Bar-Haim A, Youngster I. Ibuprofen use and clinical outcomes in COVID-19 patients. *Clin Microbiol Infect* [Internet]. 2020 Sep 1 [cited 2022 Jan 15];26(9):1259.e5-1259.e7. Available from: <https://pubmed.ncbi.nlm.nih.gov/32535147/>
126. Manjani L, Desai N, Kohli A, Arya R, Woods C, Desale S. Effects of Acetaminophen on Outcomes in Patients Hospitalized with COVID-19. *Chest* [Internet]. 2021 Oct [cited 2022 Jan 15];160(4):A1072. Available from: </pmc/articles/PMC8503320/>
127. Awasthi S, Wagner T, Venkatakrishnan AJ, Puranik A, Hurchik M, Agarwal V, et al. Plasma IL-6 levels following corticosteroid therapy as an indicator of ICU length of stay in critically ill COVID-19 patients. *Cell death Discov* [Internet]. 2021 Jun 1 [cited 2022 Jan 16];7(1). Available from: <https://pubmed.ncbi.nlm.nih.gov/33723251/>
128. van Paassen J, Vos JS, Hoekstra EM, Neumann KMI, Boot PC, Arbous SM. Corticosteroid use in COVID-19 patients: a systematic review and meta-analysis on clinical outcomes. *Crit Care* [Internet]. 2020 Dec 1 [cited 2022 Jan 16];24(1). Available from: <https://pubmed.ncbi.nlm.nih.gov/33317589/>
129. Morán Blanco JI, Alvarenga Bonilla JA, Homma S, Suzuki K, Fremont-Smith P, Villar Gómez de las Heras K. Antihistamines and azithromycin as a treatment for COVID-19 on primary health care – A retrospective observational study in elderly patients. *Pulm Pharmacol Ther* [Internet]. 2021 Apr 1 [cited 2022 Jan 29];67:101989. Available from: </pmc/articles/PMC7833340/>
130. Hirasawa N. Expression of Histidine Decarboxylase and Its Roles in Inflammation. *Int J Mol Sci* [Internet]. 2019 Jan 2 [cited 2022 Jan 29];20(2). Available from: <https://pubmed.ncbi.nlm.nih.gov/30654600/>
131. Hogan RB, Cannon T, Rappai M, Studdard J, Paul D, et al. Dual-histamine receptor blockade with cetirizine - famotidine reduces pulmonary symptoms in COVID-19 patients. *Pulm Pharmacol Ther* [Internet]. 2020 Aug 1 [cited 2022 Jan 29];63. Available from: <https://pubmed.ncbi.nlm.nih.gov/32871242/>
132. Crespi B, Alcock J. Conflicts over calcium and the treatment of COVID-19. *Evol Med Public Heal* [Internet]. 2021 [cited 2022 Jan 16];9(1):149. Available from: </pmc/articles/PMC7717197/>
133. Jiang B, Liang S, Liang G, Wei H. Could dantrolene be explored as a repurposed drug to treat COVID-19 patients by restoring intracellular calcium homeostasis? *Eur Rev Med Pharmacol*

- Sci [Internet]. 2020 [cited 2022 Jan 16];24(19):10228–38. Available from: <https://pubmed.ncbi.nlm.nih.gov/33090434/>
134. Cesta MC, Zippoli M, Marsiglia C, Gavioli EM, Mantelli F, Allegretti M, et al. The Role of Interleukin-8 in Lung Inflammation and Injury: Implications for the Management of COVID-19 and Hyperinflammatory Acute Respiratory Distress Syndrome. *Front Pharmacol* [Internet]. 2022 Jan 12 [cited 2022 Jan 16];0:3931. Available from: <https://www.frontiersin.org/articles/10.3389/fphar.2021.808797/full>
 135. Chiang CC, Korinek M, Cheng WJ, Hwang TL. Targeting Neutrophils to Treat Acute Respiratory Distress Syndrome in Coronavirus Disease. *Front Pharmacol*. 2020 Oct 9;11:1576.
 136. WHO expert panel strongly advises against use of hydroxychloroquine to prevent covid-19 | *BMJ* [Internet]. [cited 2022 Feb 6]. Available from: <https://www.bmj.com/company/newsroom/who-expert-panel-strongly-advises-against-use-of-hydroxychloroquine-to-prevent-covid-19/>
 137. Caldwell B, Aldington S, Weatherall M, Shirtcliffe P, Beasley R. Risk of cardiovascular events and celecoxib: a systematic review and meta-analysis. *J R Soc Med* [Internet]. 2006 Mar [cited 2022 Feb 6];99(3):132. Available from: </pmc/articles/PMC1383759/>
 138. Bakker T, Klopotoska JE, Eslami S, De Lange DW, Van Marum R, Van Der Sijs H, et al. The effect of ICU-tailored drug-drug interaction alerts on medication prescribing and monitoring: Protocol for a cluster randomized stepped-wedge trial. *BMC Med Inform Decis Mak* [Internet]. 2019 Aug 13 [cited 2022 Apr 28];19(1):1–10. Available from: <https://bmcmmedinformdecismak.biomedcentral.com/articles/10.1186/s12911-019-0888-7>
 139. Alshalalfa M, Alhadj R. Using context-specific effect of miRNAs to identify functional associations between miRNAs and gene signatures. *BMC Bioinformatics*. 2013 Sep 24;14(SUPPL12).
 140. Zhang S, Amahong K, Sun X, Lian X, Liu J, Sun H, et al. The miRNA: a small but powerful RNA for COVID-19. *Brief Bioinform* [Internet]. 2021 Mar 1 [cited 2022 Jan 16];22(2):1137–49. Available from: </pmc/articles/PMC7989616/>
 141. Gutmann C, Khamina K, Theofilatos K, Diendorfer AB, Burnap SA, Nabeebaccus A, et al. Association of cardiometabolic microRNAs with COVID-19 severity and mortality. *Cardiovasc Res* [Internet]. 2021 Nov 10 [cited 2022 Jan 16]; Available from: </pmc/articles/PMC8689968/>
 142. Hirohara M, Saito Y, Koda Y, Sato K, Sakakibara Y. Convolutional neural network based on SMILES representation of compounds for detecting chemical motif. *BMC Bioinformatics* [Internet]. 2018 Dec 31 [cited 2022 Jan 12];19(19):83–94. Available from: <https://bmcbioinformatics.biomedcentral.com/articles/10.1186/s12859-018-2523-5>
 143. Wu F, Zhou Y, Li L, Shen X, Chen G, Wang X, et al. Computational Approaches in Preclinical Studies on Drug Discovery and Development. *Front Chem*. 2020 Sep 11;0:726.

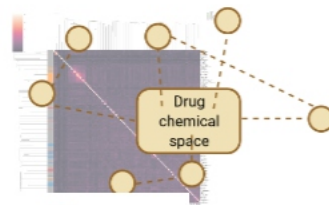
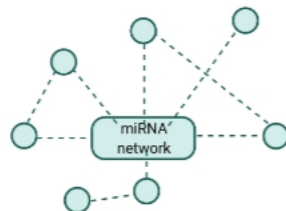
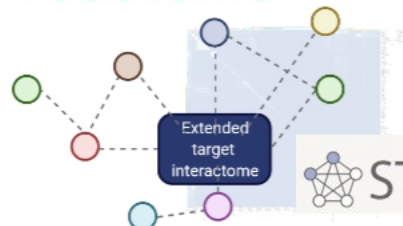
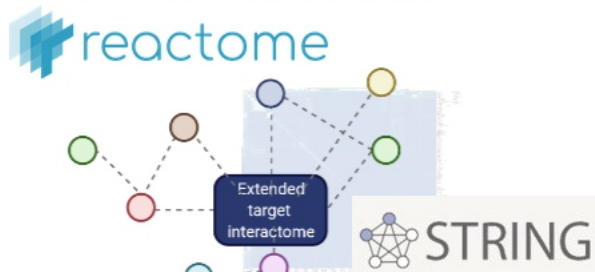
Covid-19 clinical trial data



Compound and drug data

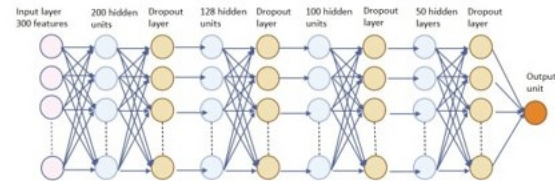
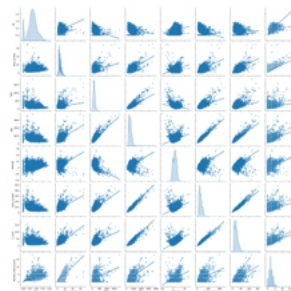
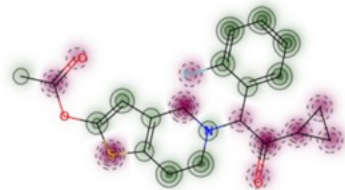


In-depth drug interactome analyses

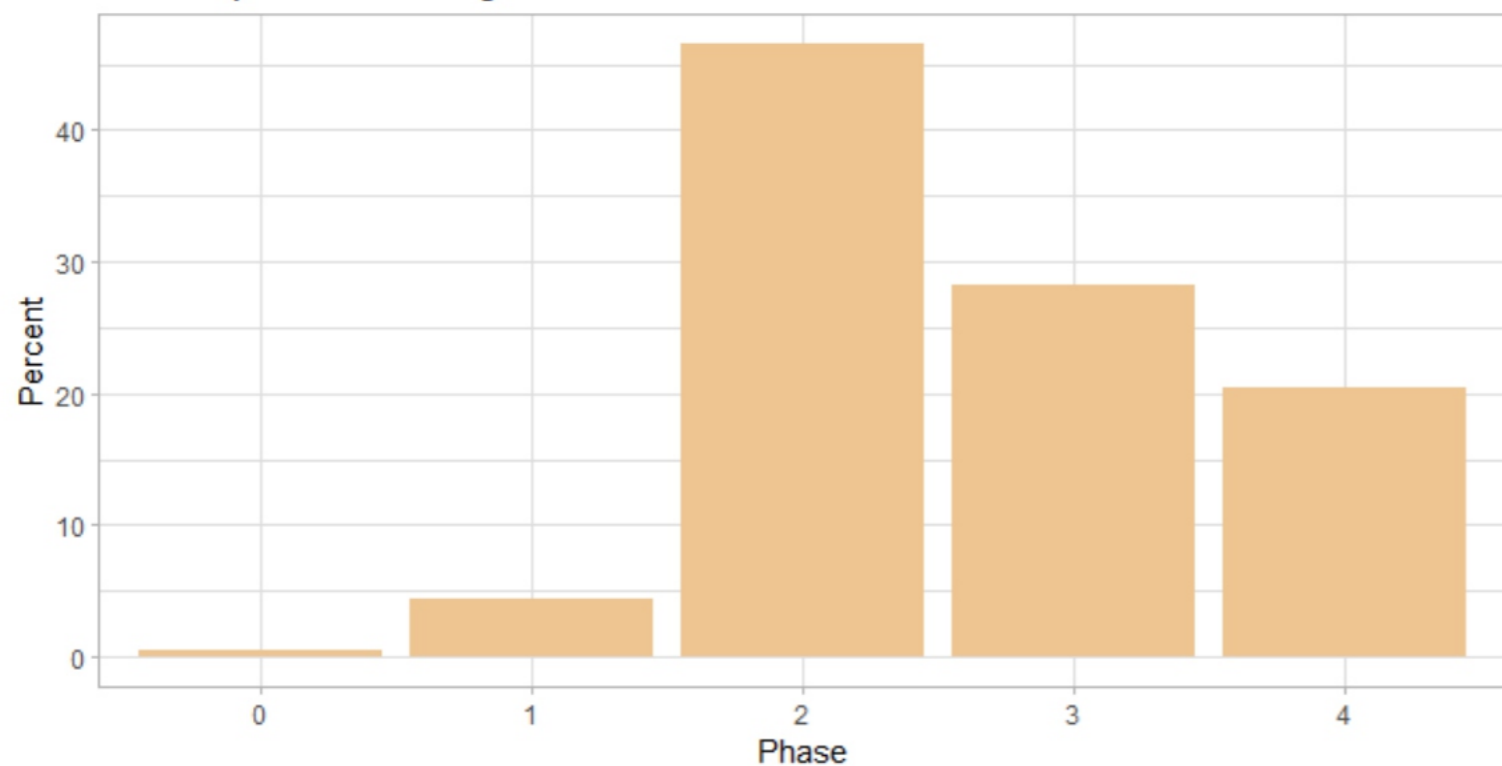


PubChem

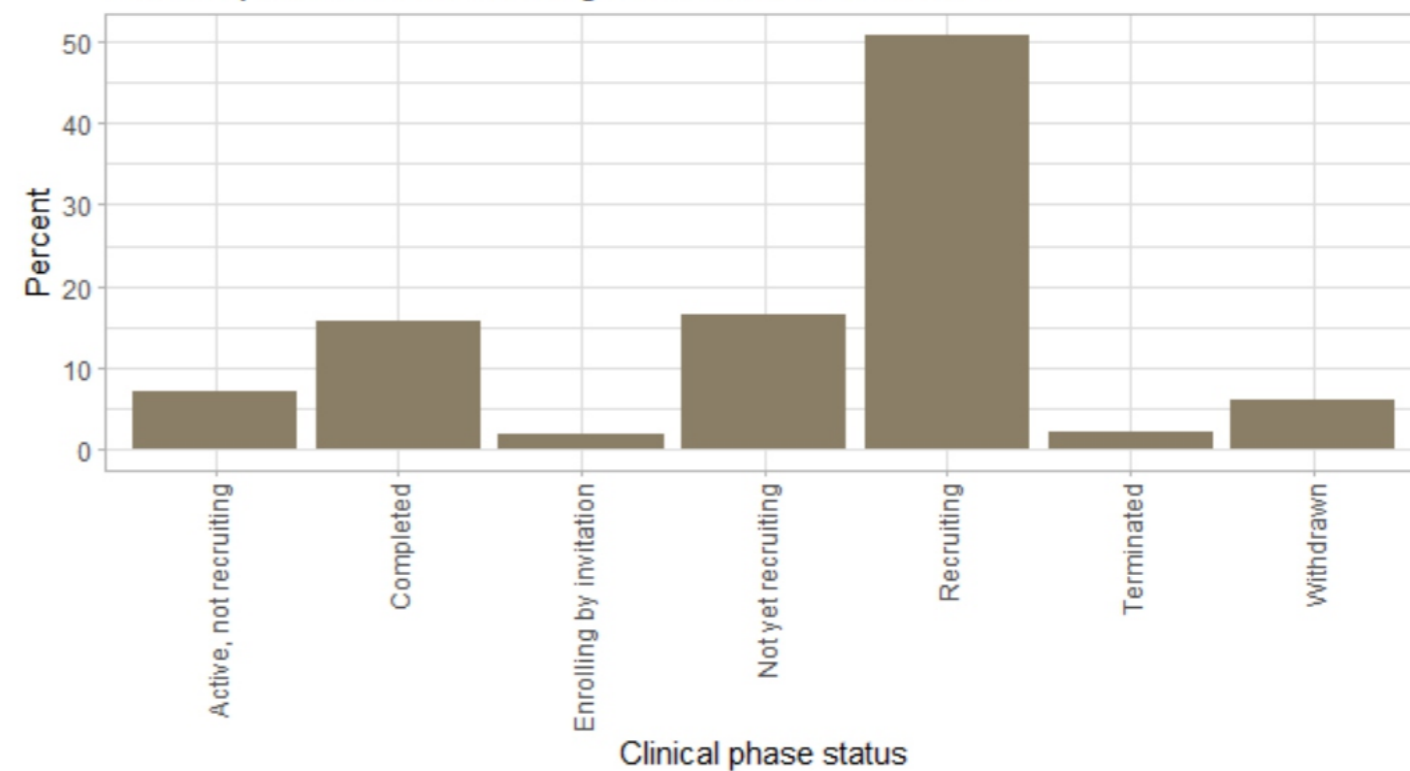
Chemical space analysis and modelling



Clinical phases for drugs used to treat COVID19

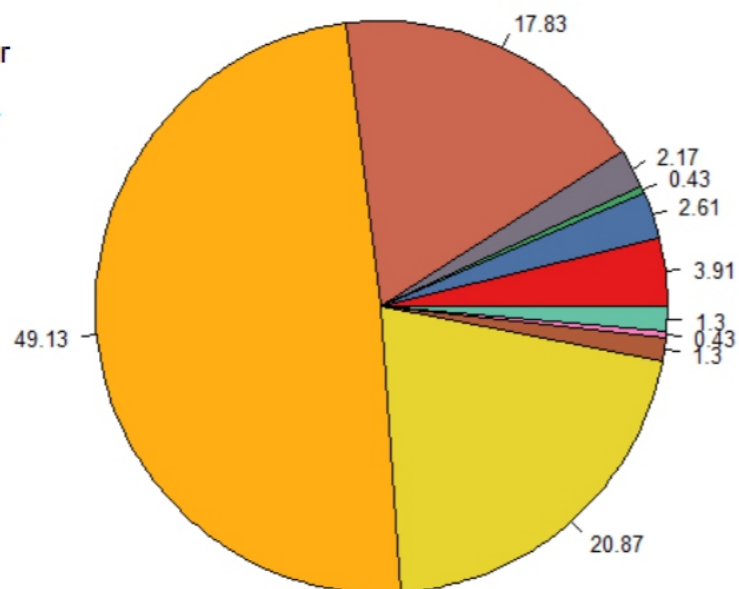


Clinical phase status for drugs used to treat COVID19



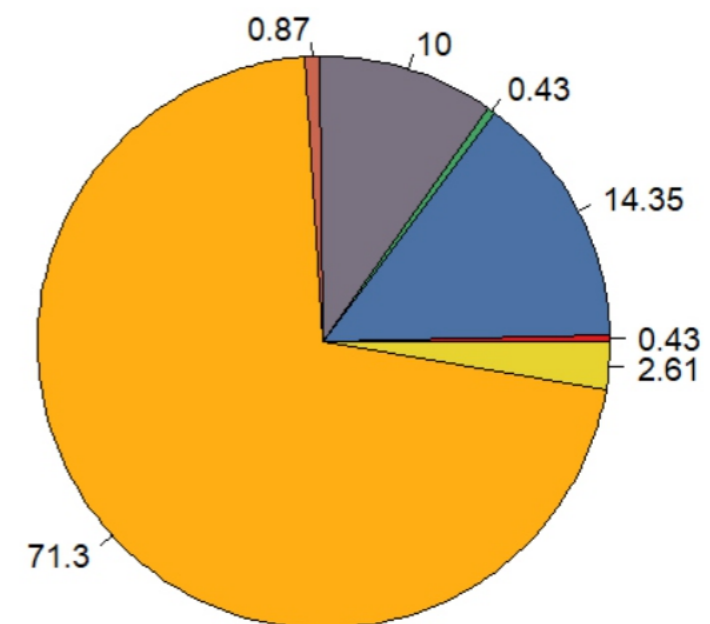
Drug activity types

- Activator
- Blocker
- Binding agent
- Modulator
- Antagonist
- Inhibitor
- Agonist
- Negative allosteric modulator
- Opener
- Positive allosteric modulator



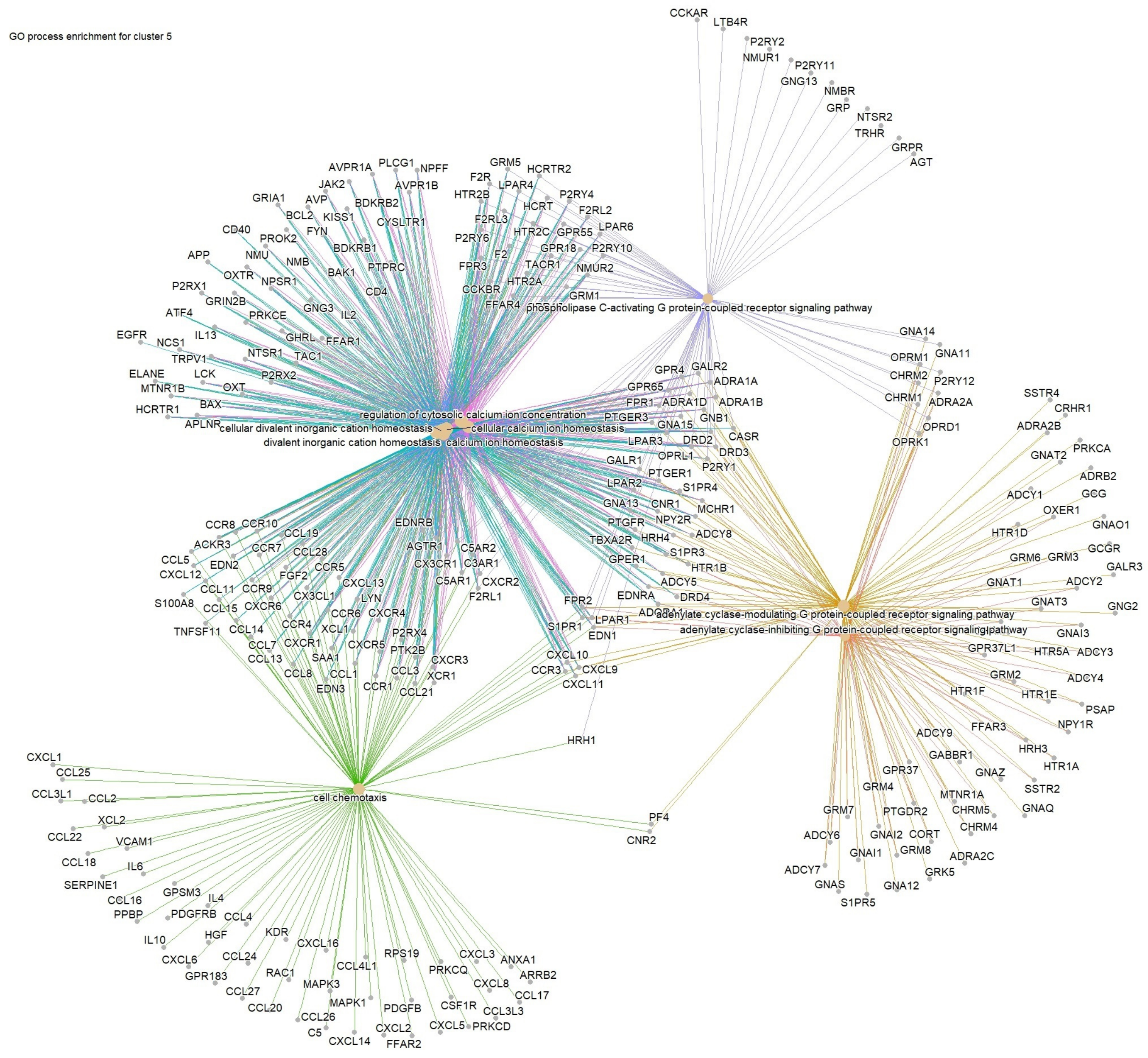
Drug molecule types

- Enzyme
- Antibody
- Oligonucleotide
- Protein
- Unknown
- Small molecule
- Oligosaccharide

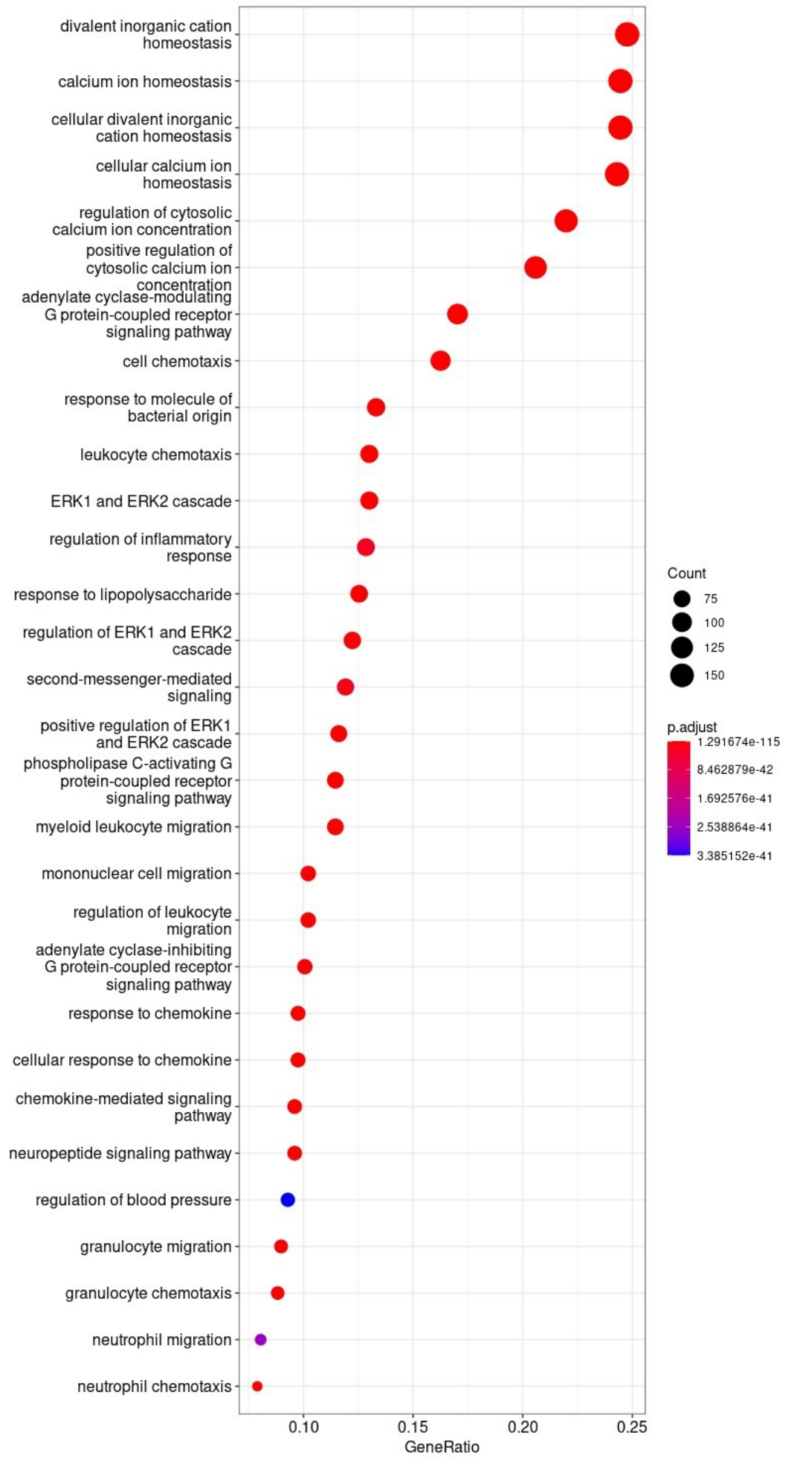


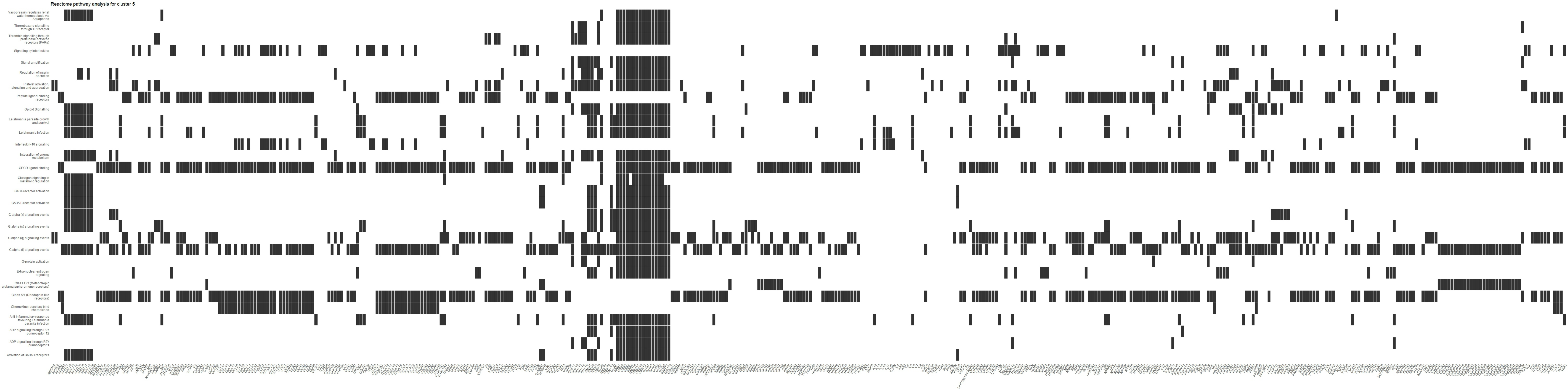
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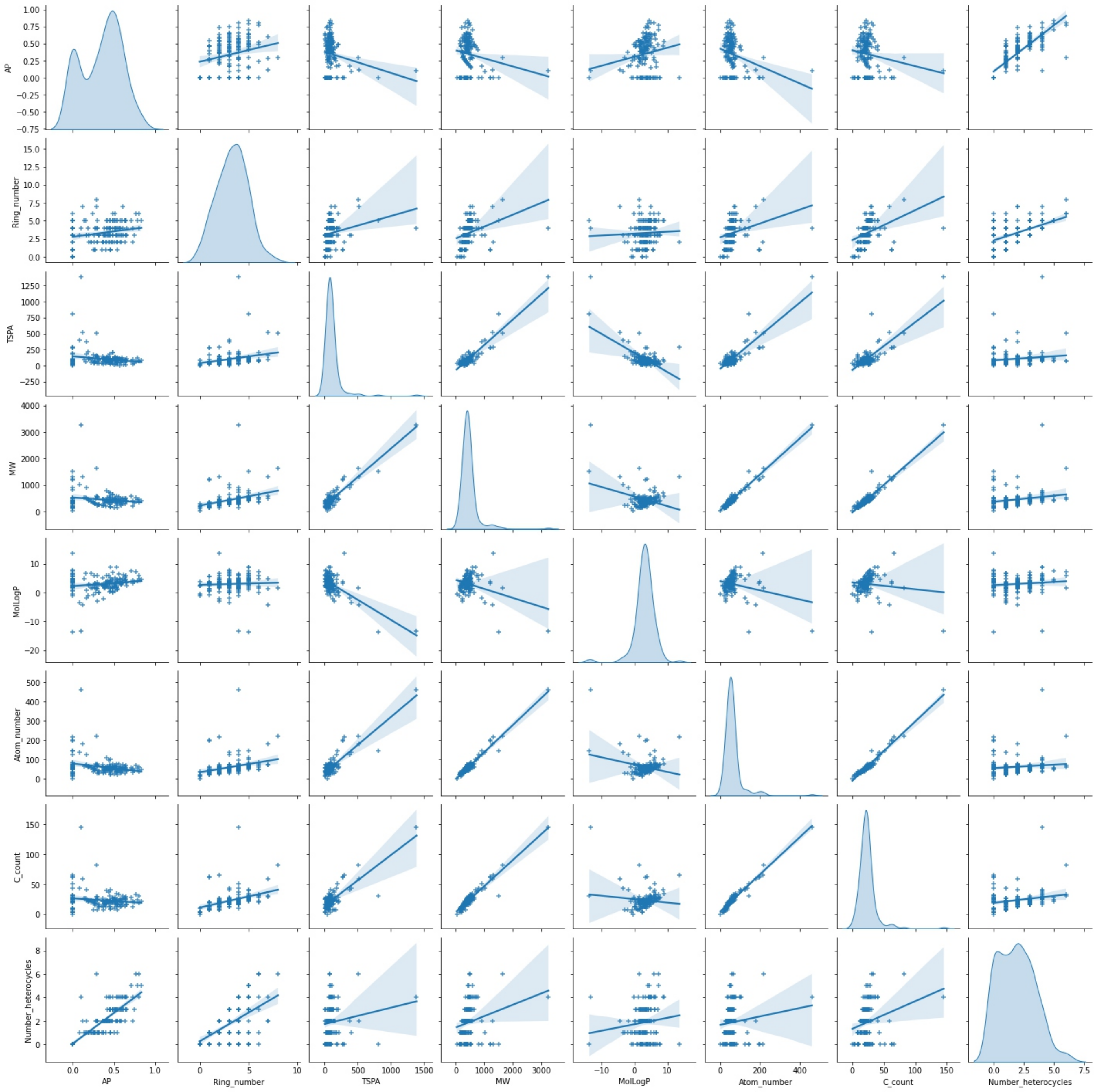
GO process enrichment for cluster 5

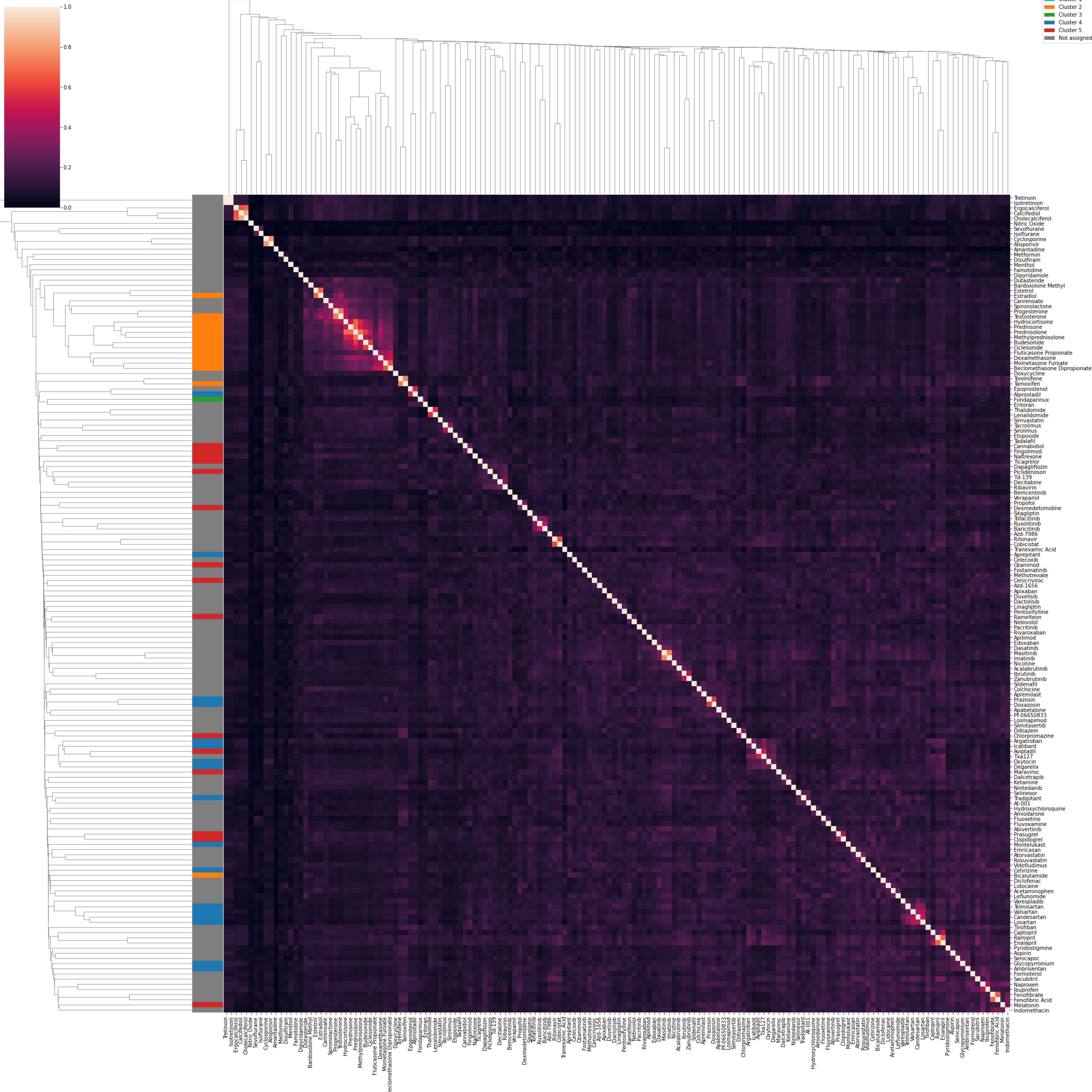
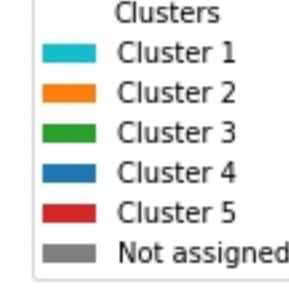
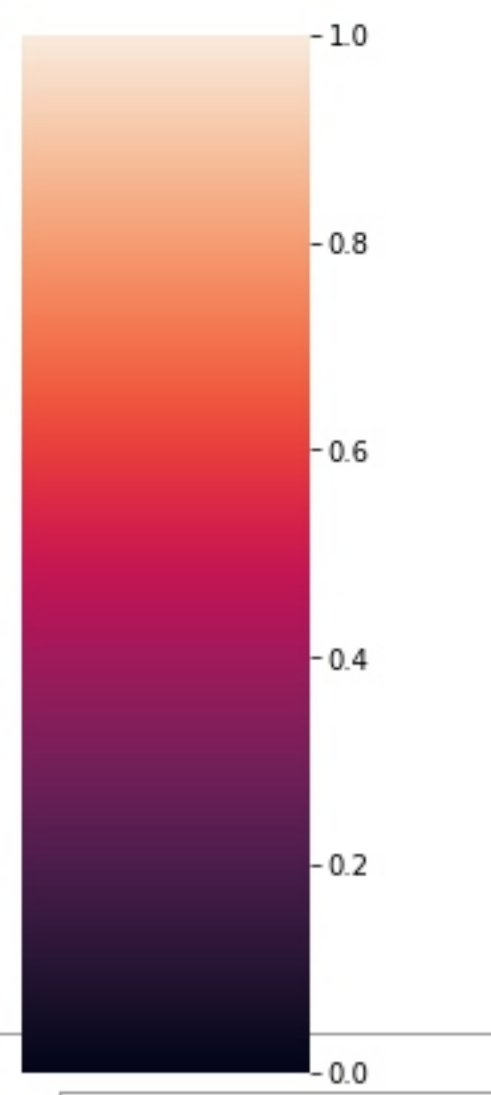
**B**

Functional enrichment for cluster 5









Tretinoin
Isotretinoin
Ergocalciferol
Calcifediol
Cholecalciferol
Nitric Oxide
Sevoflurane
Isoflurane
Cyclosporine
Alisporvir
Amantadine
Metformin
Disulfiram
Menthol
Famotidine
Dipyridamole
Dulasteride
Bardoxolone Methyl
Estradiol
Canrenate
Spironolactone
Progesterone
Testosterone
Hydrocortisone
Prednisone
Prednisolone
Methylprednisolone
Budesonide
Ciclesonide
Fluticasone Propionate
Dexamethasone
Mometasone Furoate
Beclomethasone Dipropionate
Doxycycline
Tremifene
Tamoxifen
Epoprostenol
Alprostadil
Fondaparinux
Ertoran
Thalidomide
Lenalidomide
Sunitinib
Tacrolimus
Sirolimus
Etoposide
Ecdalafi
Cannabidiol
Fingolimod
Nafarexone
Ticagrelor
Dapagliflozin
Piclidenoson
Td-129
Decitabine
Ribavirin
Bemcentinib
Verapamil
Propofol
Dexmedetomidine
Stagglipitin
Ticagrelor
Ruxolitinib
Baricitinib
Acd-7986
Ritonavir
Cobicistat
Tanexamic Acid
Aprepitant
Celecoxib
Ozanimod
Fostamatinib
Methotrexate
Cenicriviroc
Azd-1656
Apixaban
Duvelisib
Dactolisib
Linagliptin
Pentoxifylline
Ranmelteon
Nebivolol
Pacritinib
Rivaroxaban
Aplilmod
Edoxaban
Dasatinib
Masitinib
Imatinib
Nicotine
Acalabrutinib
Ibrutinib
Zanubrutinib
Sildenafil
Colchicine
Apremilast
Prazosin
Doxazosin
Apabetalone
PF-06650833
Losmapimod
Siltuximab
Diltiazem
Chlorpromazine
Argatroban
Icatibant
Aviptadil
Tx127
Oxytocin
Degarelix
Maraviroc
Ketamine
Dalcetrapib
Ketamine
Nintedanib
Sildenafil
Tadipitant
AZ-001
Hydroxychloroquine
Amiodarone
Fluoxetine
Fluvoxamine
Abivertinib
Prasugrel
Clopidogrel
Montelukast
Etrimeric
Atorvastatin
Rosuvastatin
Vidofluvidimus
Cetirizine
Bicalutamide
Diclofenac
Lidocaine
Acetaminophen
Leflunomide
Varespladib
Sildenafil
Valsartan
Candesartan
Losartan
Tirofiban
Captopril
Ramipril
Enalapril
Pyridostigmine
Aspirin
Senicapoc
Glycopyrronium
Ambrisentan
Formoterol
Sacubitril
Nagroxen
Ibuprofen
Fenofibrate
Fenofibric Acid
Melatonin
Indomethacin