

# Predicting the activity of protein-ligand complexes

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# Declaration

I hereby declare and confirm that this thesis is entirely the result of my own original work. Where other sources of information have been used, they have been indicated as such and properly acknowledged. I further declare that this or similar work has not been submitted for credit elsewhere. This printed copy is identical to the submitted electronic version.

Hagenberg, June 27, 2023

Lukas Fallmann

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# Preface

# Abstract

This should be a 1-page (maximum) summary of your work in English.



# Kurzfassung

An dieser Stelle steht eine Zusammenfassung der Arbeit, Umfang max. 1 Seite. ...

# Chapter 1

## Introduction

The discovery of new drugs or any chemically active compounds for that matter is an expensive and time-consuming process. It has been estimated, that it takes about 14 Years from the initial discovery of a promising new compound to the release of a marketable drug[1]. In addition to that the price of this drug-discovery circle ranges up to 800 Million Dollars[2]. All techniques which aim to improve the efficiency of drug discovery can be generalized as one of two methods. These two are called High-throughput-screening (HTS) and virtual screening (VS)[1].

When using an HTS-approach there are many compounds which are tested against some type of target protein. During testing, it is measured whether a certain compound biochemically interacts with a protein. Those interacting combinations are considered active and are marked by researchers as hits. To improve the performance of HTS there are a number of factors to consider. Through miniaturization, it is possible to investigate more compounds at the same time. With a higher throughput quality-control is more time-consuming and leads to an overall more expensive process. For this reason HTS is most efficient, when analyzing a small set of compounds as the technology is not suitable for large datasets[3].

In contrast to the in vitro approach of HTS, VS is a theoretical in silico approach. To save resources in the laboratory the activity of certain compounds is predicted using a preexisting library of small molecules. The activity can be predicted using the ligands of a compound and their respective binding sites or the 3D structure of a compound. The Key idea behind the ligand based approach (LBVS) is that similar compounds have similar chemical properties. Therefore, the goal of LBVS is to find molecules which have similar or identical chemical properties as the sample compound[4]. Structure-based VS uses the 3D structure of a compound to predict which molecules from the dataset will bind to the provided sample. Each molecule of a certain database subset is fitted (docked) to the sample. Hereby it is important to differentiate between rigid and flexible docking.

In rigid docking the dataset sample is rotated and translated in a six-dimensional space in order to fit the sample protein. For each fitted molecule a score is calculated based on how well the molecule fits to the sample[5]. Although this algorithm often predicts actual possible binding sites and bound proteins, there is no guarantee that this compound will actually bind in vitro. Therefore, predicted interactions should be seen as a hypothesis. Still rigid docking provides a great baseline at a comparatively low cost[4]. The low

accuracy of rigid docking is due to the nature of biochemical substances as samples in a database can only provide a snapshot of a sample. With flexible docking it is possible to simulate moving binding sites. The flexibility can be introduced at different stages. Implicit flexibility is achieved by smoothing protein surfaces and therefore allowing room for interpretation when docking. Cross- or Ensemble docking can be done by repeating the docking process with different conformations. Explicit flexibility is reached through allowing side-chain flexibility. Most commonly utilized is the approach where the ligand is flexible, and the receptor is rigid. Even though this approach does provide better more accurate results it takes considerably longer to compute[6].

Regardless of the docking type the score decides which pose between a protein and a ligand is most likely to exist. In addition to that, the score also determines whether a protein-ligand complex is considered active. There are a lot of different scoring functions which can be grouped into four categories: physics-based, empirical, knowledge-based, and machine learning-based[7].

The focus of this work is on implementing a machine-learning based scoring approach. Machine-learning based scoring functions work by training on pre-classified data and finding the best model for predicting future data. To accurately and efficiently train a model crucial binding sites need to be identified beforehand. The basis of this thesis is the master thesis of Birklbauer Micha[8]. In his thesis a selection of eleven proteins from the directory of useful decoys have been selected to be analyzed. For the selected proteins all possible interactions have been analyzed. Based on the interaction-data a few basic scoring functions have been implemented. The direct result of this thesis are proteins and the frequency of their interactions.

Since this work aims to implement different machine learning algorithms for use in scoring functions the state of the art is described in the following.

## 1.1 Machine Learning in drug design and activity prediction

The following chapter summarizes the recent developments in drug design using various machine learning techniques.

Today there exist a multitude of machine learning approaches in the field of drug design and activity prediction. As a result of various AI breakthroughs in recent years there have been numerous research projects regarding the usability of artificial intelligence in various bioinformatic domains. MILCDock uses the Output of five traditional Scoring Functions as input for a neural Network. This technique has a slight performance benefit when compared to traditional scoring functions [9].

## 1.2 Goals

The goals of this thesis are twofold:

1. Evaluate common machine learning approaches for activity prediction and compare results with current literature.
2. Fine tune the provided datasets using a multitude of feature-engineering practices, to increase performance of the ML approaches.

The second goal can be viewed as an extension of the first one since its primary aim is to improve the results achieved while pursuing the first goal.

## Chapter 2

# Methods

### 2.1 Data description

Explain where data came from and how it is structured.

#### 2.1.1 Proteins

Overview of the proteins that were used for this thesis.

#### 2.1.2 Interactions

Overview of the interactions between protein and ligands used in this thesis.

### 2.2 Data partitioning

Explain concept of train-test split, cross-evaluation ?

### 2.3 Machine Learning approaches

Introduction to the ML Approaches used for the thesis.

### 2.4 quality metrics

How to measure the quality of a machine learning system.

### 2.5 hyperparameter search

Introduction to hyperparameter optimization.

### 2.6 feature engineering

Explain concept of feature engineering and possible implications for thesis.

## Chapter 3

# Results

### 3.1 Performance per Protein-Complex

Evaluate models tuned for selected proteins.

### 3.2 Performance Overview – Comparing ML-approaches

Compare performance of overall ml approaches.

## Chapter 4

# Discussion

### 4.1 Conclusion

Recap findings of thesis.

### 4.2 Improvements and outlook

Explain possible improvements to used technique. Provide general outlook on topic.

## Appendix A

# Technical Details



## Appendix B

# Supplementary Materials

List of supplementary data submitted to the degree-granting institution for archival storage (in ZIP format).

### B.1 PDF Files

Path: /

thesis.pdf . . . . . Master/Bachelor thesis (complete document)

### B.2 Media Files

Path: /media

\*.ai, \*.pdf . . . . . Adobe Illustrator files  
\*.jpg, \*.png . . . . . raster images  
\*.mp3 . . . . . audio files  
\*.mp4 . . . . . video files

### B.3 Online Sources (PDF Captures)

Path: /online-sources

Reliquienschrein-Wikipedia.pdf [10]

Appendix C

Questionnaire

Appendix D

LaTeX Source Code

# References

## Literature

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- [9] Connor Morris et al. “MILCDock: Machine Learning Enhanced Consensus Docking for Virtual Screening in Drug Discovery”. *Journal of chemical information and modeling* 62 (Nov. 2022). DOI: 10.1021/acs.jcim.2c00705 (cit. on p. 3).

## Online sources

- [10] *Reliquienschrein*. Aug. 29, 2022. URL: <https://de.wikipedia.org/wiki/Reliquienschrein> (visited on 02/11/2023).

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