

# Multi-omics molecular treatment targets for Prostate Cancer

#### What is Prostate Cancer

The **prostate gland** is a male reproductive organ. It secretes prostate fulid, one of the main components of semen. Prostate cancer is a malignant growth in the prostate gland.

Prostate Cancer is the second most commonly diagnosed cancer among men.



#### **Risk factors**

Different factors might influence a man's risk of developing prostate cancer:

- **Age**: Increased risk in men over 50.
- Family history: Increased risk for patients having a family member with prostate cancer.
- Race/ Ethnicity: Black men are at higher risk of developing prostate cancer in comparison to white men.

#### Forms of Prostate Cancer

Patients suffering from Prostate Cancer may experience indolent or aggressive disease.

- Indolent disease slow growing form of Prostate Cancer, unlikely to progress in the absence of treatment.
- **Aggressive disease** life-threatening disease in the absence of treatment.

#### Clinical problem

Accurate discrimination between indolent and aggressive Prostate Cancer is still challenging.

This results in over-treatment of patients with indolent disease and under-treatment of those with lethal disease.

New drugs for advanced Prostate Cancer were recently approved. Low response rate and development of resistance remains a barrier to improve on therapeutic outcomes.

## Why studying Prostate Cancer is important

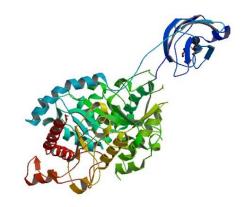
New drugs for Prostate Cancer are urgently needed, linked to the evidence that even though Prostate Cancer is frequently treatable (paradoxically also over-treated) when detected early, once the disease has begun to advance, response to treatment is generally limited and the disease continues to worsen regardless from the means of intervention.



**PCaProTreat** 

#### **Drug target**

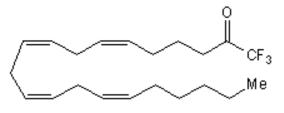
- Usually a protein present in disease-affected tissue
- Associated with disease
- Targeted by a drug to give a curative effect



Selection of the "good" drug target is very important for effective treatment.

## **Drug candidate**

- A compound with therapeutic potential
- It can be antibody, small molecule etc.
- Sufficient selectivity for the target
- Have activity to produce desired effect
- Enough evidence supporting further pre-clinial and clinial development



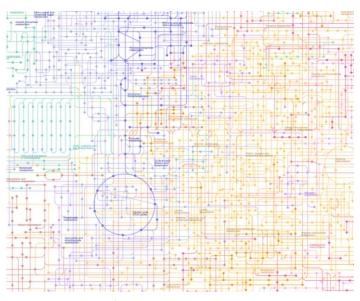
#### Challenge

Molecular changes underlying Prostate Cancer development are complex and differ among patients (heterogeneity), which has an impact on the treatment response.

Therefore, selection of a "good" drug target is very challenging.

Stumbling blocks for selecting the "good" drug target:

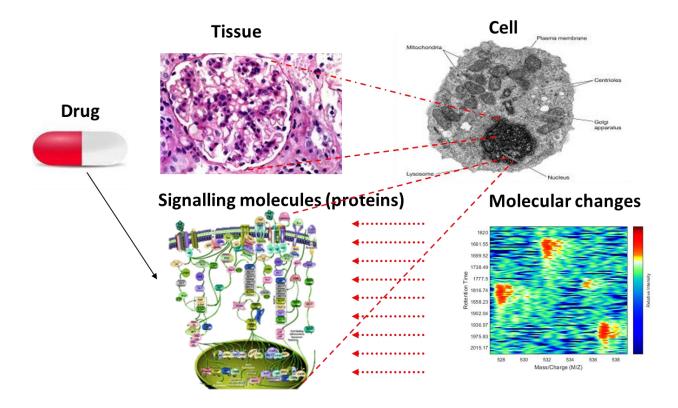
- Insufficient knowledge of the molecular pathophysiology.
- Lack of proper assessment of biological relevance.
- Underestimation of complexity and heterogeneity.



Adapted from: https://www.genome.jp/kegg/

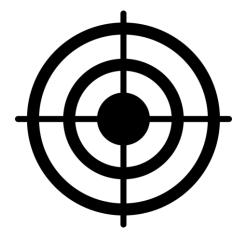
### Make it perfect

We aim to improve our understanding of molecular changes associated with Prostate Cancer progression to define novel drug targets/ drug candidates based on the molecular pathophysiology.

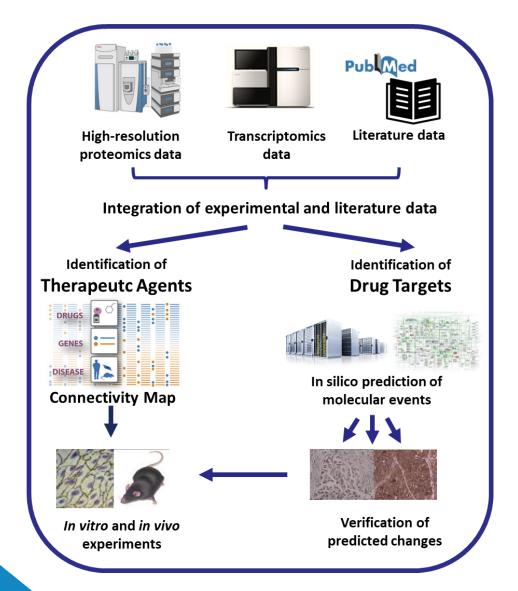


#### **PCaProTreat Objectives**

- 1. Establishment of knowledgebase with molecular features related to Prostate Cancer.
- 2. Characterisation of the molecular changes underlying Prostate Cancer progression through the integration of multisource (tissue, urine, and seminal plasma) and multi-omics data, complemented with literature-mined data.
- 3. Definition of biological processes/ pathways and their regulatory elements, followed by their validation.
- 4. In silico prediction of potential drug candidates.



#### Our Patients' centered approach

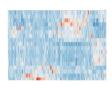


- 1. Identify novel drug targets through linking the molecular changes to the disease etiology.
- 2. Predict new drug candidates able to reverse disease phenotype on the basis of the existence of relationship between disease, proteins, and drugs.

#### Our contribution

# Molecular signature of Prostate Cancer progression

A tissue proteome-based molecular signature comprised of multi-omics and cross-correlated molecular features





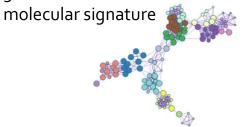
# Prostate Cancer knowledgebase

A knowledgebase of disease-relevant molecular alterations



# Understanding molecular mechanisms of Prostate Cancer Progression

The biological processes and pathways involved in disease progression as revealed based on the



# Potential Drug targets / Drug candidates

Novel drug targets / drug candidates defined on the basis of molecular pathophysiology



# Where you can meet us?

Detailed information on the PCaProTreat Project is provided in our website

https://pcaprotreat.eu/

Join us on Facebook



https://www.facebook.com/PCaProTreat/

#### Who we are

PCaProTreat: Multi-omics molecular treatment targets for Prostate Cancer

Marie Skłodowska-Curie Actions Individual Fellowship H2020-MSCA-IF-2017-800048





#### **Contact Us**

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