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## Introduction

- **Rapid progression and vague symptoms** of pancreatic cancer contribute to its predominantly late diagnosis and high mortality rate
- In Germany, it is projected to become the **second leading cause of cancer-related deaths** and the **fifth most common cancer** by 2030 (Quante et al. 2016)
- **Early detection** enables surgical and other treatment options which can **improve survival rates** and **patient outcomes**
- The **societal impact of a blood-based pancreatic cancer test**, which is being developed by the PANCAID project, will be evaluated



What is the impact of a pancreatic cancer screening policy on patient survival for screened vs. not screened patients?

## Results

- Annual screening of a hypothetical population with 100,000 individuals at 1.9% pancreatic cancer lifetime risk starting at age 55 for 4 rounds resulted in an average life-year gain of **1.6 years per true-positively screened individual**
- 44 individuals had a true-positive result and out of these patients, 36 received an early diagnosis at stage 1 or 2
- The share of patients surviving at least 5 years after a pancreatic cancer diagnosis was **29% vs. 5% when true-positively screened versus not screened**
- A specificity of 90% or 95% would lead to 30,000 or 17,500 false-positive results
- Sensitivity analysis: starting screening later than 55 increases the average life-year gain up until age 67 as start age, from which this gain decreases

## Conclusion

- Screening may result in a **life-year gain** in true-positive patients compared to the no-screening alternative
- Pancreatic cancer screening in the normal population at about 2% lifetime risk would likely result in an unacceptable number of false-positive and few true-positive patients  
→ Early detection strategies for pancreatic cancer should target high-risk individuals instead of the general population
- Further modifications of the model will investigate which threshold risk, test parameters, and optimal screening protocols, could enable early detection under certain circumstances
- Evaluating the cost-effectiveness of blood-based testing for pancreatic cancer will be the ultimate purpose of the model

## Methods

- **Microsimulation** of screening effectiveness compared to no screening
- Generating **life histories** of male and female individuals at risk (i.e. time of birth and death)
- Simulating **pancreatic cancer histories** of individuals given a lifetime risk
- Applying a hypothetical **screening policy** with assumptions (e.g. start/ stop age, frequency) and data specifications (e.g. sensitivity, specificity)
- Survival data was sourced from the literature (Huang et al. 2018)
- Analysis of **differences in survival**: successfully screened vs if no screening would have taken place

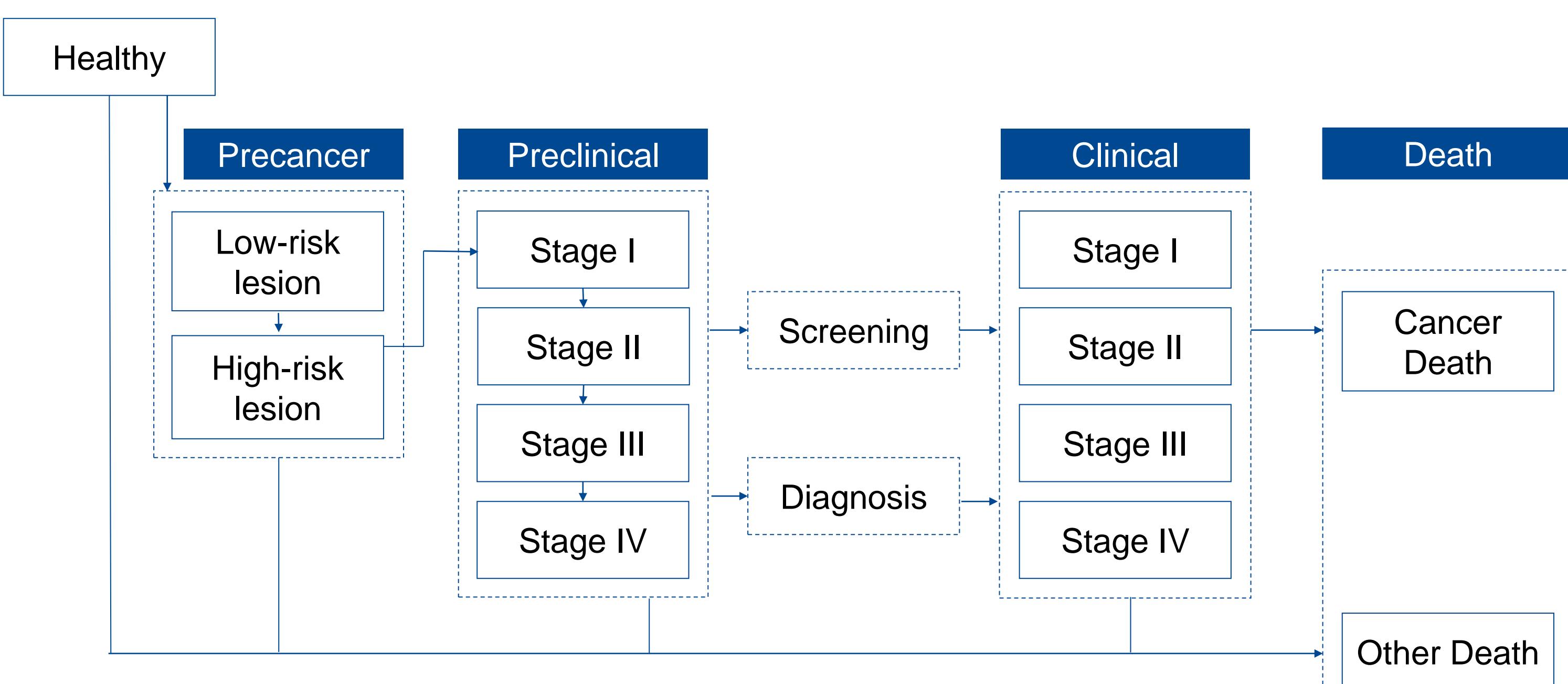


Figure 1: State transition diagram

**Population:**  
100,000 individuals

**Duration:**  
Lifetime

**Test sensitivity parameters:**  
from 65% (Stage I) to 95% (Stage IV)

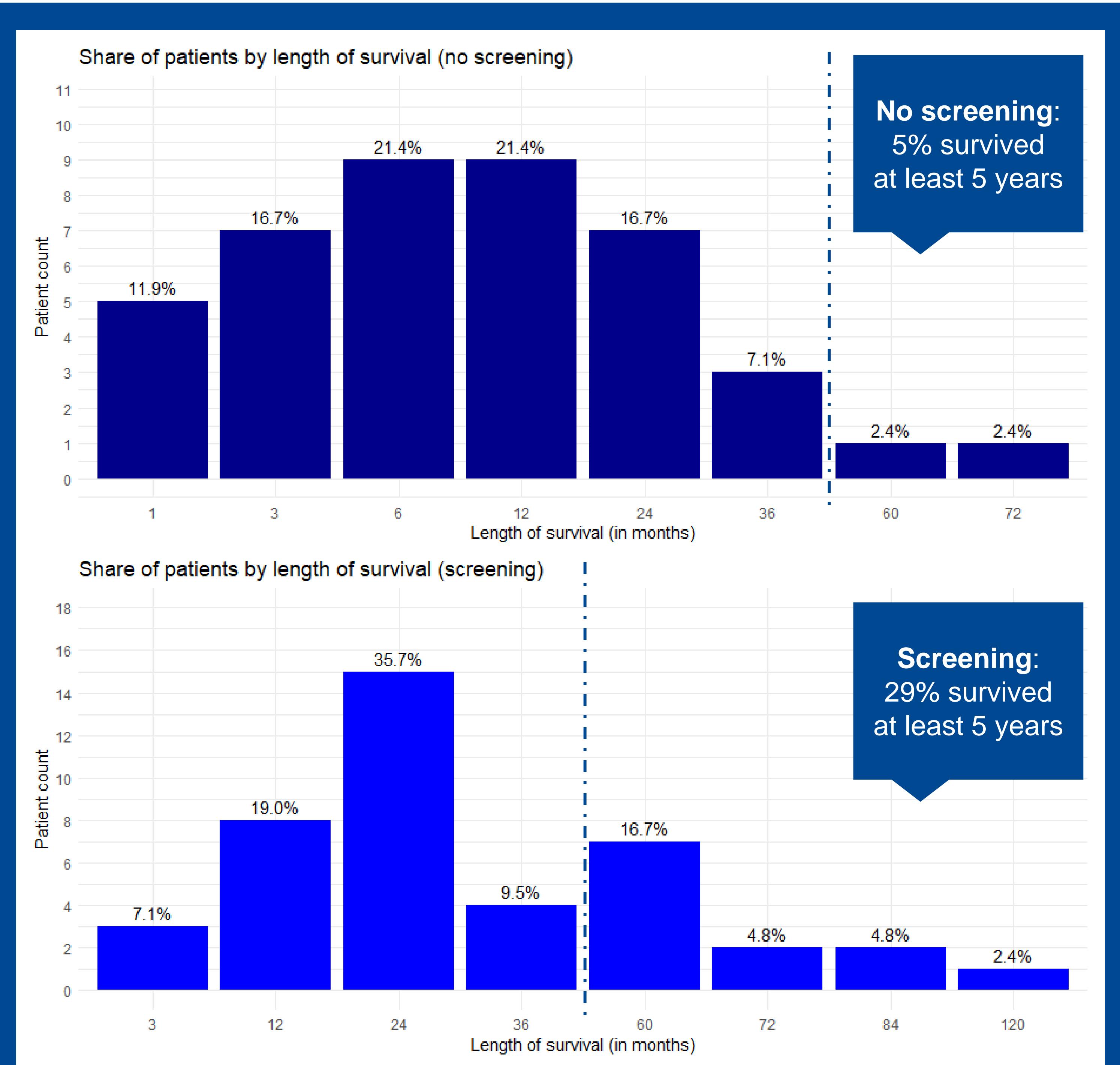


Figure 2: Share of survival lengths after pancreatic cancer diagnosis in presence and absence of screening

## References

Huang, L., Jansen, L., Balavarca, Y. et al. Stratified survival of resected and overall pancreatic cancer patients in Europe and the USA in the early twenty-first century: a large, international population-based study. *BMC Med* 16, 125 (2018).

Quante AS, Ming C, Rottmann M, Engel J, Boeck S, Heinemann V, Westphalen CB, Strauch K. Projections of cancer incidence and cancer-related deaths in Germany by 2020 and 2030. *Cancer Med*. 2016 Sep;5(9):2649-56.

## PANcreatic CAncer Initial Detection via liquid biopsy



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