

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

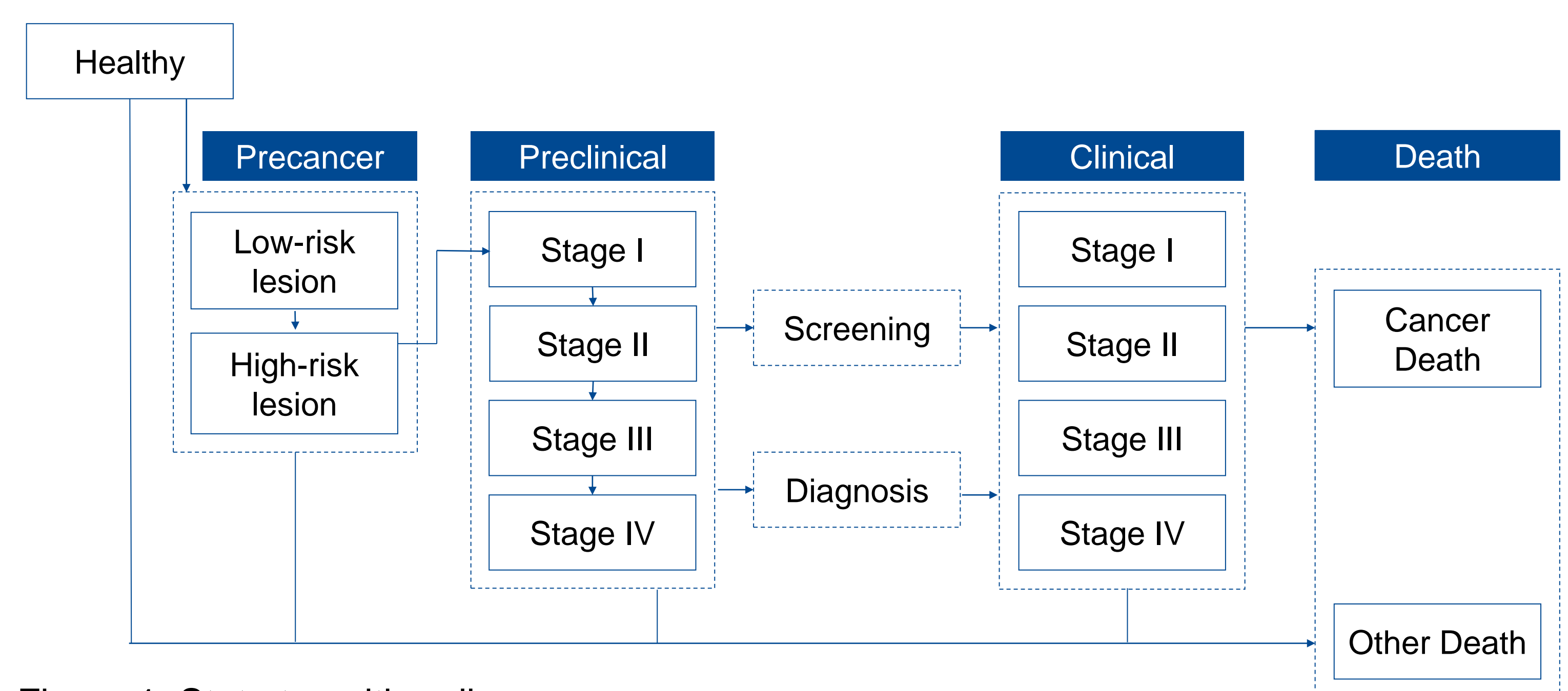


Figure 1: State transition diagram



Population:
100,000
individuals



Duration:
Lifetime



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

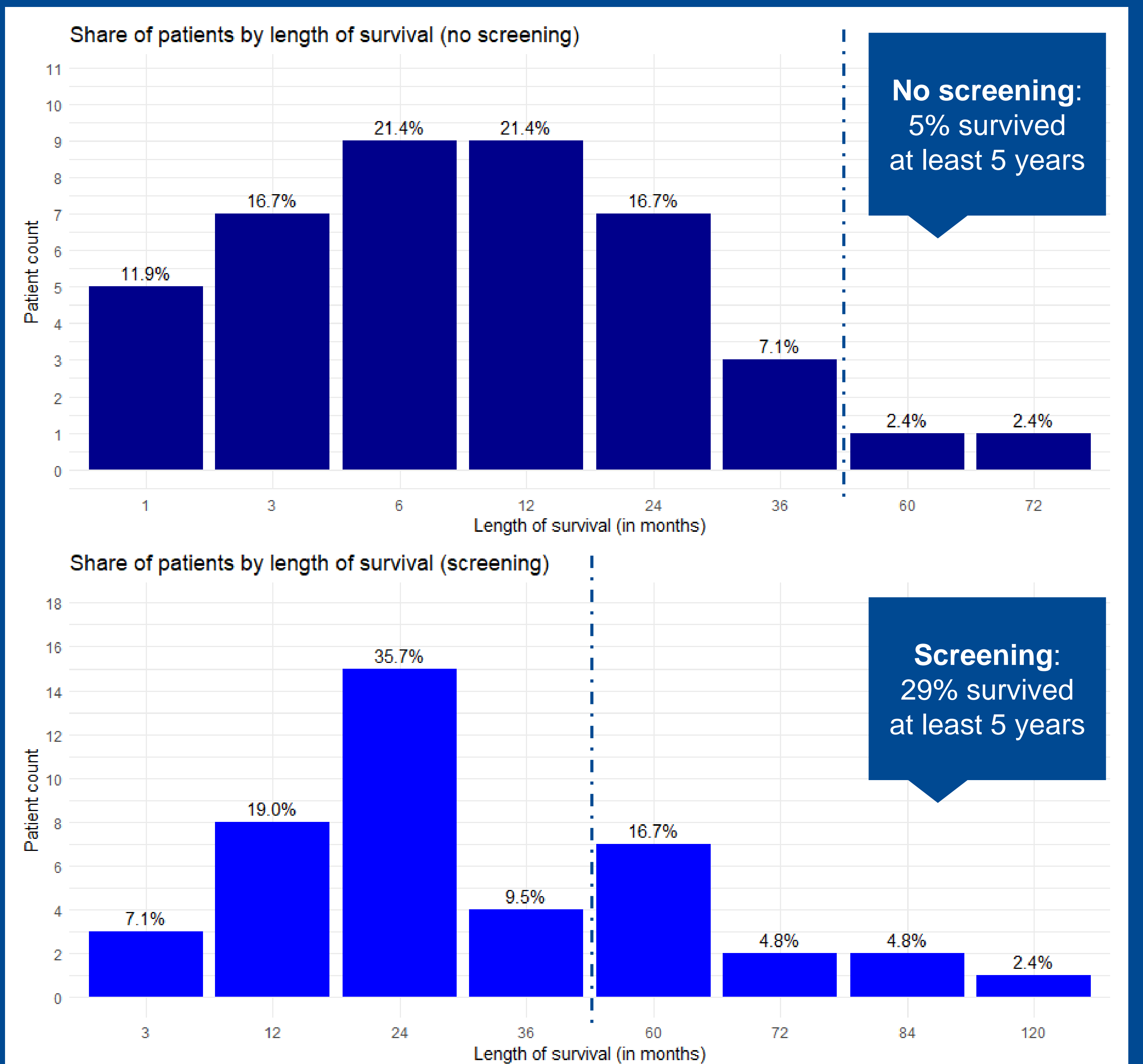


Figure 2: Share of survival lengths after pancreatic cancer diagnosis in presence and absence of screening

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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