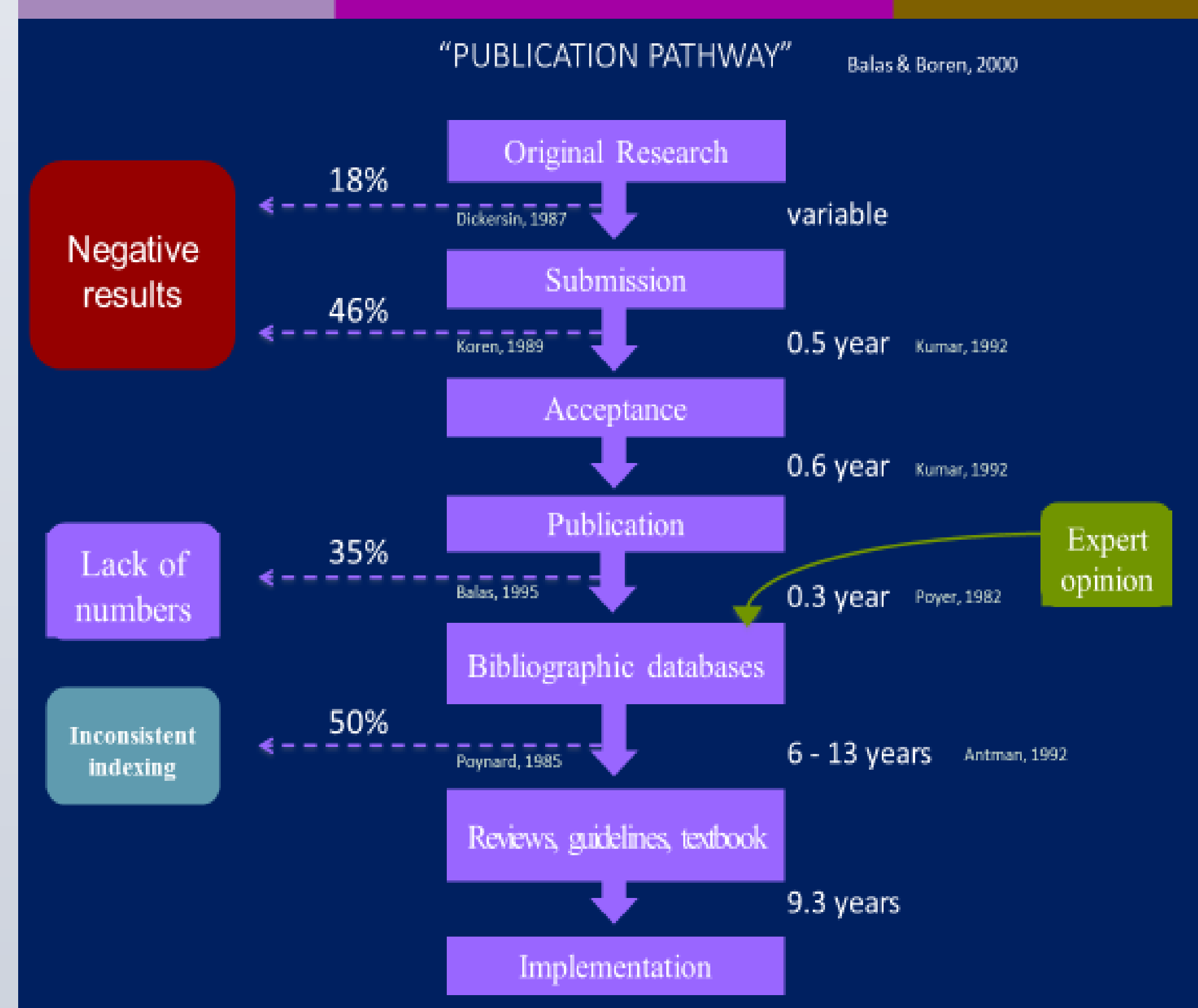


Background

The slow transfer of research evidence to practice is a problem that is widely recognized by researchers, practitioners, policy makers, and patients. According to a review published in 2000 by Balas and Boren, it takes an average of **17 years** for research evidence to reach clinical practice (50% uptake). This finding, published 17 years ago, was based on nine medical procedures in the 1980s. Despite the frequent use of this statistic to highlight the problem of slow research translation, we do not know its relevance to cancer research nor to more recent trends. We explore the amount of time it takes for translation of research to uptake of evidence-based practice in cancer. Our objective was to determine the time it took for evidence from the seminal publication to reach at least 50% uptake in the target population.



Methods

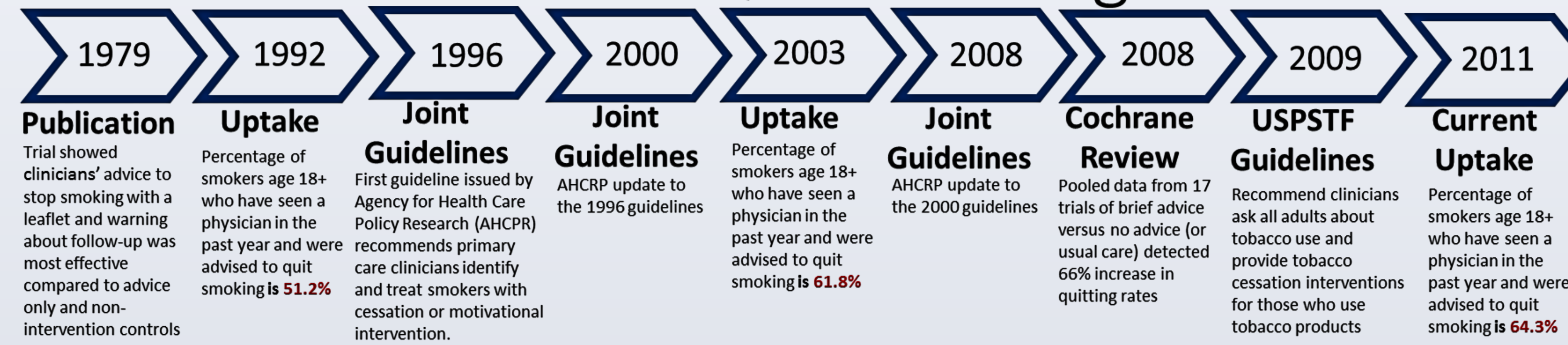
We included evidence-based programs, practices, or interventions (herein referred to as EBPs) in cancer prevention and screening with professional guidelines and population-based data on uptake. We included five EBPs: clinicians' advice to quit smoking, HPV vaccination, HPV testing, colorectal cancer screening, and mammography. To determine the time from research publication to implementation, we identified the seminal study, defined as a published article that provided sufficient evidence for the effectiveness of the EBP. All but one of our EBPs had an RCT as the seminal study. The exception, HPV testing, was an observational study. Next we identified professional guidelines issued wholly or in part by a government agency that were concordant with the findings of the seminal study. We also searched for systematic reviews to determine if the evidence from the seminal study had been incorporated into the review. The data on uptake of the EBP was reviewed for all years where available, to follow any trends.

We calculated the number of years from publication of the seminal study to initial publication of the guideline to implementation, defined as 50% uptake in the population for which that EBP was recommended. We also calculated the average number of years to implementation for all five EBPs. We also traced other important events along the pathway to implementation that occurred after the seminal study and contributed to the development of the full evidence base leading to a guideline or a review.

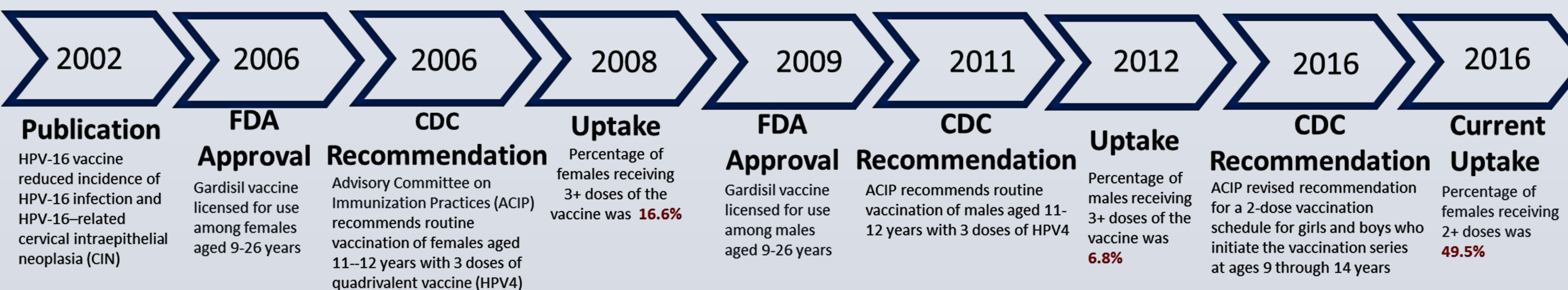
EBP Implementation Timelines

The timelines for each EBP represent different events along the pathway to implementation and the data on uptake that was available either prior to reaching 50% uptake or after reaching 50% uptake.

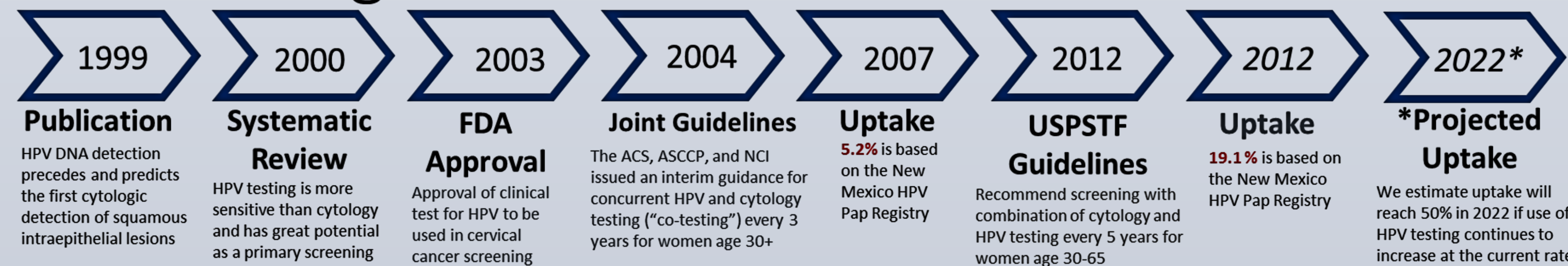
Clinicians' Advice to Quit Smoking



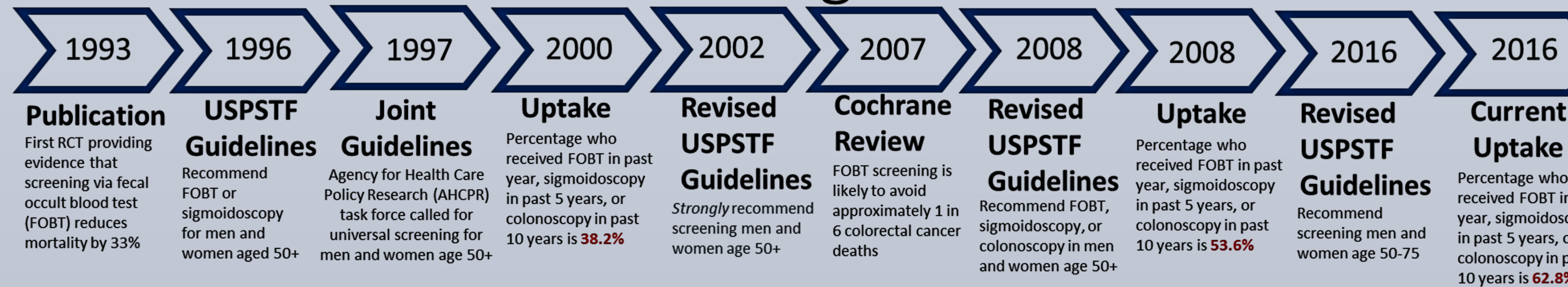
HPV Vaccination



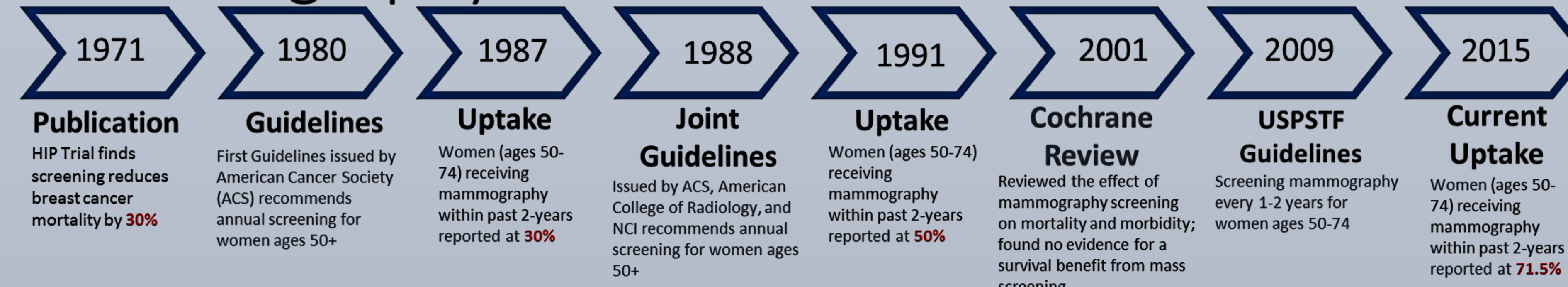
HPV Testing for Cervical Cancer



Colorectal Cancer Screening

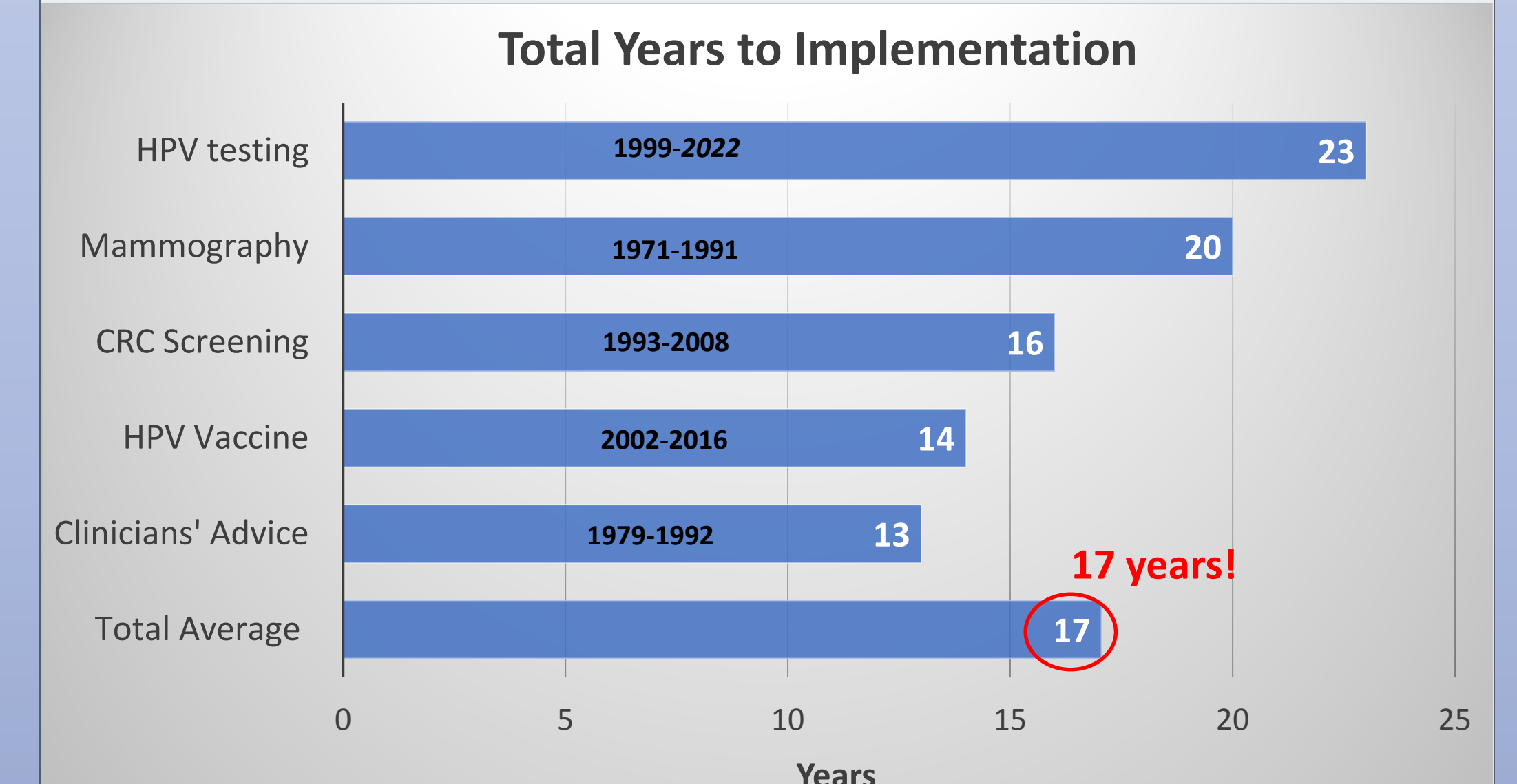


Mammography

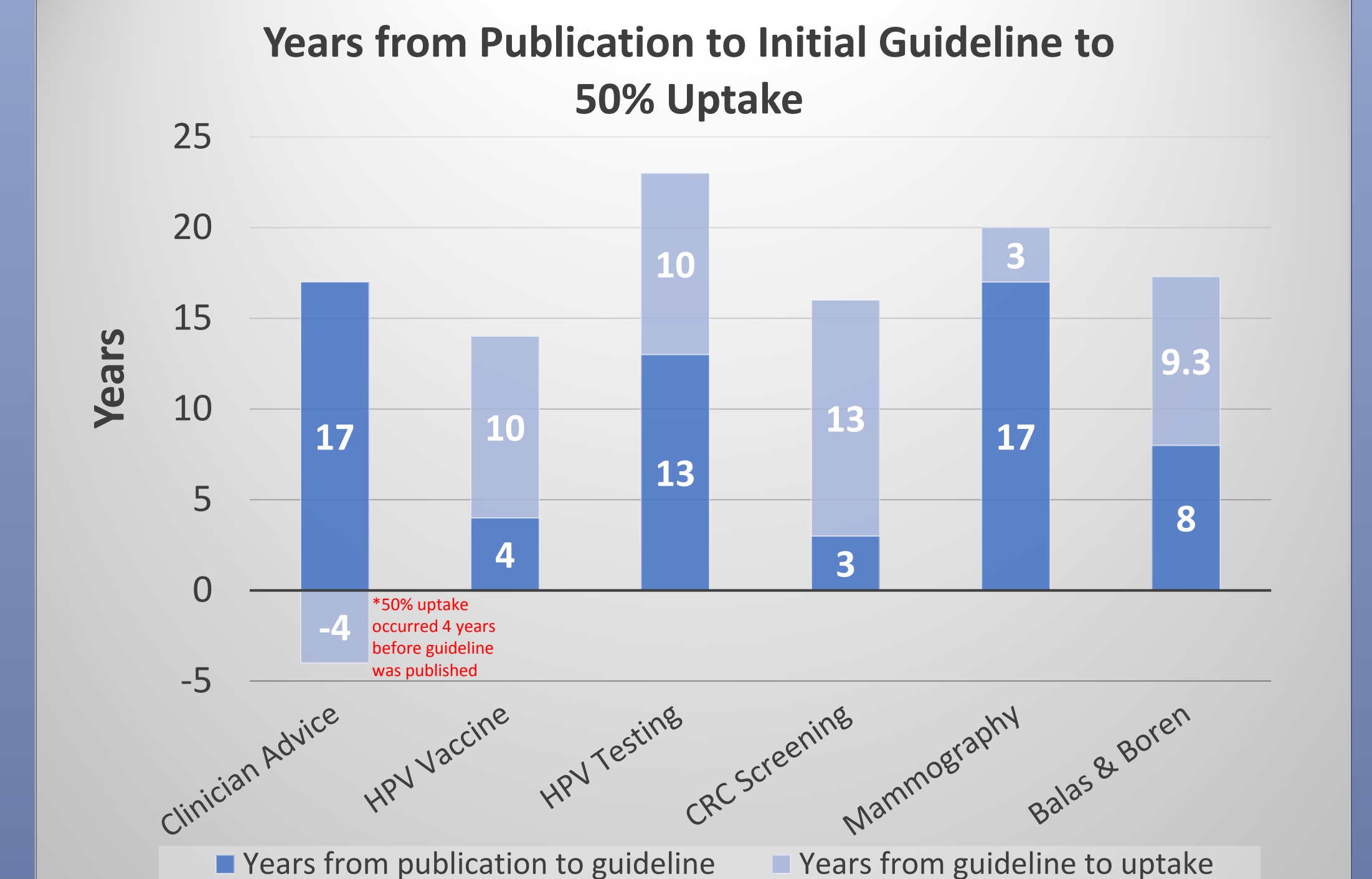


Results

Our findings highlight the complexity of piecing together the pathway from research to implementation to measure time it takes for implementing an EBP in cancer prevention and screening. The total number of years to implementation vary and average 17 years.



Our examples demonstrate that the path to research translation is not linear and each stage builds to inform the next. As more evidence accumulates and contributes to the formation of revised guidelines, we get closer to achieving substantive rates of uptake.



Limitations

We included only five EBPs because data was limited on uptake over time, and for many EBPs, the evidence changed so rapidly that it was impossible to trace uptake before the evidence and subsequent guidelines changed.

Implications and Future Research

We measured the time from publication to implementation, defined as 50% uptake, for select EBPs in cancer control. We found similar results to Balas and Boren's from 2000 that it takes 17 years. All interventions except HPV testing have reached substantial uptake to date. We encourage investigators to explore ways to increase the rate of uptake of HPV testing and other EBPs in cancer control.

Data Sources

- NCI Cancer Trends Progress Report
- State Cancer Profiles
- U.S. Preventive Services Task Force recommendations
- CDC Morbidity and Mortality Weekly Report (MMWR)
- AHCPR (AHRQ) Guidelines
- PubMed, Google Scholar to identify seminal publications and systematic reviews
- Food and Drug Administration (FDA) Pre-Market Approval letter for HPV DNA Test
- NCI and FDA staff expert opinion

Seminal Publications

- Russell MAH, Wilson C, Taylor C, Baker CD. (1979) Effect of general practitioners' advice against smoking. *BMJ*. 2(6184).
- Liao KL, Glass AG, Manos MM, et al. (1999) Detection of human papillomavirus DNA in cytologically normal women and subsequent cervical squamous intraepithelial lesions. *JNCI*. 91(11).
- Koutsky LA, Ault KA, Wheeler CM, et al. (2002) A controlled trial of a human papillomavirus type 16 vaccine. *NEJM*. 347(21).
- Mandel JS, Bond JH, Church TR, et al. (1993) Reducing mortality from colorectal cancer by screening for fecal occult blood. Minnesota Colon Cancer Control Study. *NEJM*. 328(19).
- Shapiro S, Strax P, Venet L. (1971) Periodic Breast Cancer Screening in Reducing Mortality From Breast Cancer. *JAMA*. 215(11).