

Economic Analysis In Clinical Research

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Introduction to the Principles and Practice of
Clinical Research
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Overview

- * Introduction
- * Concepts
- * Technical and Statistical Issues
- * Examples
 - Value of Health
 - Cost of Illness
 - Cost-Benefit Analysis
 - Marginal Analysis
 - Cost Effectiveness Analysis
 - * Clinical Trial plus Economic Follow-up
 - * Clinical Trial plus Modeling
 - * CEA Along Side a Clinical Trial

INTRODUCTION

Purpose of Economic Analysis in Health Care Research

- * Policy and program evaluation
- * Health care resource allocation decisions
- * For understanding determinants of technical and organization efficiency of health care delivery
- * For understanding determinants of distribution, access and equity issues in health care delivery



Feature Story: Overlooked and Undertreated: Mental Health Care Is a Major Unmet Need in Countries Around the World

The last half of the 20th century saw enormous progress in the development of treatments for mental disorders, and many of them are both effective and cost-effective even in low-income countries

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About DCPD

The Disease Control Priorities Project (DCPP) is an ongoing effort to assess disease control priorities and produce evidence-based analysis and resource materials to inform health policymaking in developing countries. DCPP has produced three volumes providing technical resources that can assist developing countries in improving their health systems and ultimately, the health of their people. [More](#)

DCPP Publications

Disease Control Priorities in Developing Countries

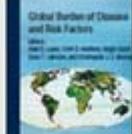
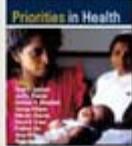
The second edition of this seminal work (DCP2), written by more than 350 specialists in diverse fields from around the world, provides the results of indepth research, offers insightful analyses, and proposes context-sensitive policy recommendations to significantly reduce the burden of disease in developing countries and to improve the quality of life for all people.

Priorities in Health

This companion volume distills the essence of DCP2 into a succinct and readable format, providing information on how to devise better strategies, policies, and choices among health interventions; how to put those decisions into practice; and how to allocate scarce resources to implement them.

Global Burden of Disease and Risk Factors

This book summarizes the concepts and estimates of the burden of disease and the attribution of this burden to several major risk factors.



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Region

Country

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Topic

EXPERT Q&A

DEAN T. JAMISON bio

The Senior Editor of *Disease Control Priorities in Developing Countries*, second edition, is pleased to answer your questions about the history of the Disease Control Priorities Project, including the scope of the subject areas covered, and the methodological foundation underlying the burden of disease estimates and cost-effective analyses. In addition, Dr. Jamison will respond to questions about the main messages of the effort, including intervention selection and neglected opportunities where investment of modest sums might have positive impact on global health.

[Submit Questions](#)

EXPERT ESSAYS

Essays from world health leaders on the value of DCPD information.

[Sir George Alleyne, Director Emeritus, Pan American Health Organization](#)

[Mr. Rajiv Mera, former Secretary of Health, India](#)

Upcoming Events

Oct 25, 2006 - Oct 27, 2006
[The World Congress on Communication for Development](#)

Oct 27, 2006
[USAID Global Health Mini-University](#)

Oct 29, 2006 - Nov 2, 2006
[Global Health Forum for Health Research](#)

[More Events](#)

Latest News

Sep 29, 2006
[DCPP - Inaugural Newsletter](#)

Related Research

- [World Development Report 1993: Investing in Health](#) (PDF | 6.1MB)
- [Global and Regional Burden of Disease and Risk Factors, 2001: Systematic Analysis of Population Health Data \(The Lancet 2006; 367:1747-1757\)](#)
- [Vaccine Preventable Diseases Still Take Toll in the Developing World](#)
- [Advancement of Global Health: Key Messages from the Disease Control Priorities Project \(The Lancet 2006; 367:1193-1205\)](#)
- [DCP2 Tuberculosis Supplementary Material \(Chapter 16 Annex\)](#) (PDF | 564.2k)

Relationship of Economic Studies to Clinical Trials

- * Health Services Research: observational data in community settings or quasi-controlled experimental settings
- * Health Economics Modeling: economic modeling combined with results of randomized clinical trials through modeling
- * Economic Studies Along-side Clinical Trials: used especially in Pharmaco-economics

CONCEPTS

Some Types of Economic Analysis in Health Care Studies

* Descriptive Studies

- Economic Value of Health
- Cost-of-Illness Studies

* Cost - efficiency Studies - What should it cost?

* Cost Evaluation Studies

- Cost Benefit Analysis (CBA)
- Cost Effectiveness Analysis (CEA)
- Cost Utility Analysis (CUA)

Principles of Cost Evaluation Studies

- * All relevant costs and benefits should be counted
- * Purpose of evaluation is to compare alternative used of resources
- * Measurement is incremental * All relevant costs and benefits should be counted
- * Purpose of evaluation is to compare alternative used of resources
- * Measurement is incremental

Cost Benefit Analysis

- * All costs and health effects are expressed in monetary terms (i.e., must put a \$ value on a year of life)
- * Cost Benefit -> all benefits minus all costs, sometimes call Social Return on Investment
- * Cost Benefit Ratio -> All benefits divided by all costs, sometimes called Social Rate of Return

Cost Effectiveness Analysis

- * Costs are expressed in monetary terms
- * Benefits are expressed in "natural units," e.g., life-years
- * Cost Effectiveness Ratio \rightarrow Cost divided by life-years (or other measure of benefit)

Cost Utility Analysis

- * Costs are expressed in monetary terms
- * Benefits are expressed in quality-adjusted "natural units," e.g., quality adjusted life-years
- * Cost Utility Ratio \rightarrow Cost divided by Quality Adjusted Life Years

Incremental Cost Effectiveness (or Utility) Ratios

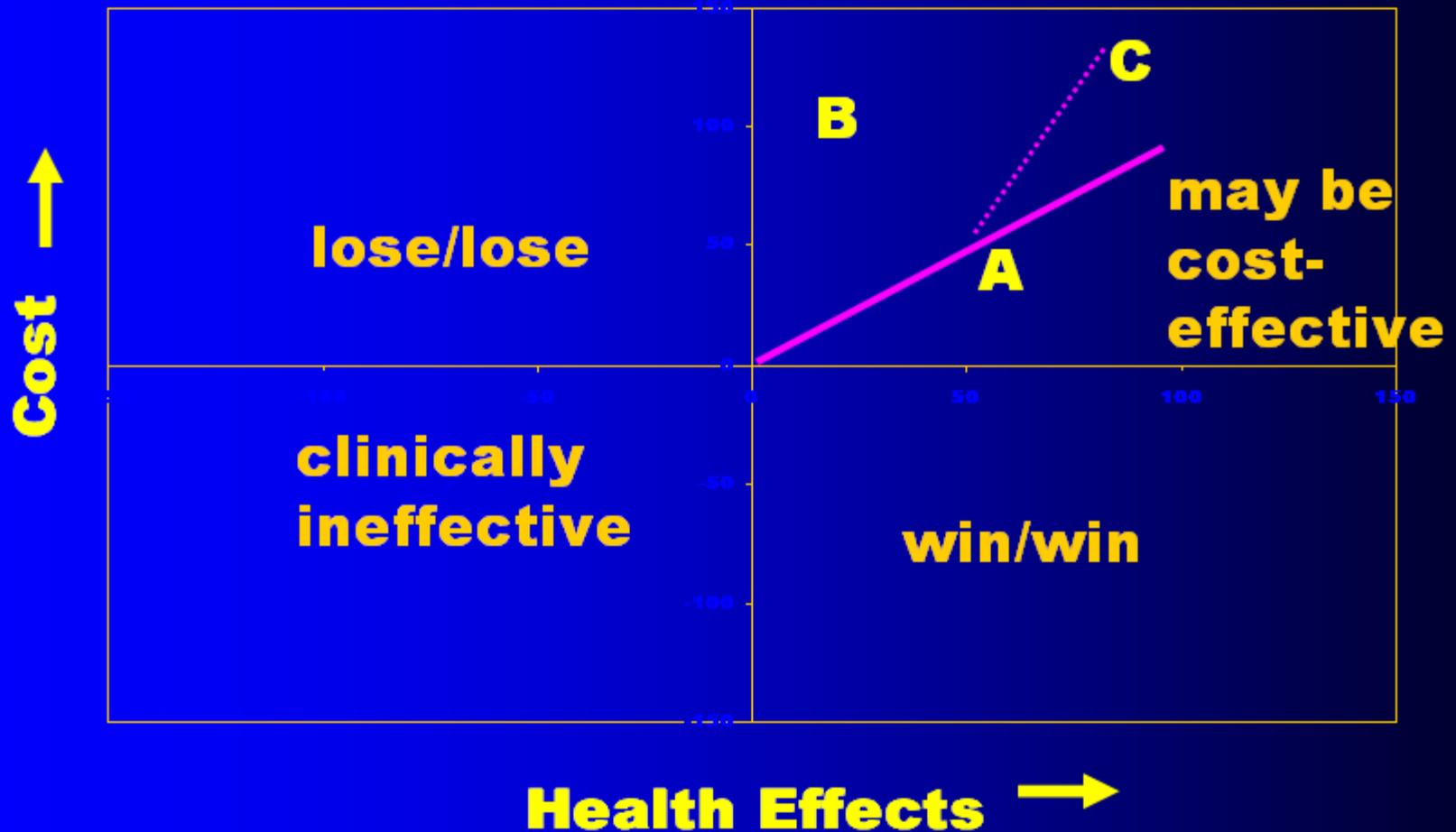
- * Let C_a and C_b be the costs of Intervention a and Intervention b;
- * Let E_a and E_b be the health effects of Intervention a and Intervention b;
- * Intervention a is often defined as status quo or standard treatment.

Incremental Cost Effectiveness Ratio (ICER)

$$\mathbf{ICER = [C_b - C_a] / [E_b - E_a]}$$

Note: This is the equation for the slope of a line when C is the vertical axis and E is the horizontal axis

The Cost-Effectiveness Plane



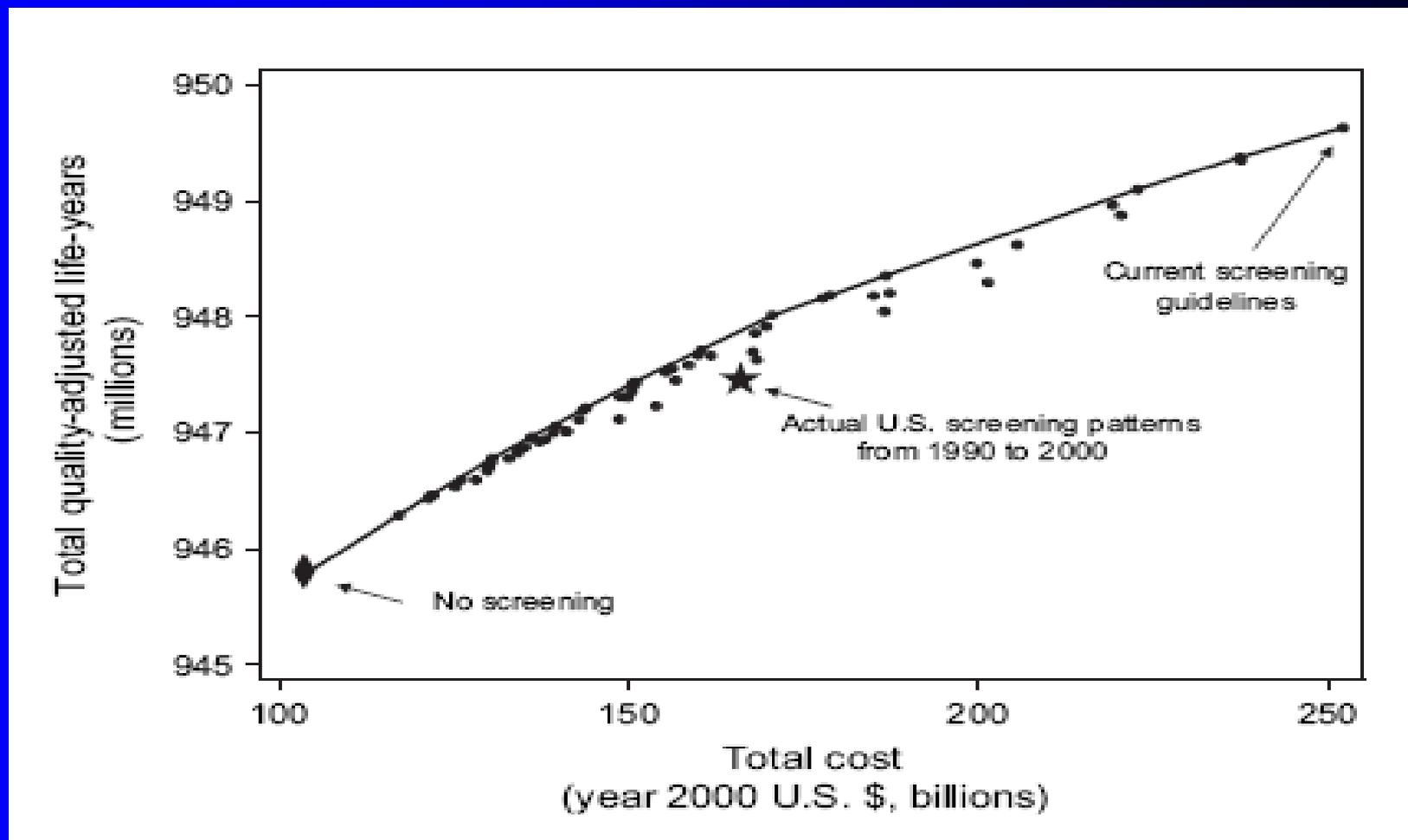
Dominance

- * Intervention A produces more health benefits at lower cost than intervention B: A “dominates” B.
- * Intervention C produces more health benefits than intervention A, but at higher cost: the ICER of A relative to C can be computed.
- * ICER of A relative to C is the slope of the dotted line

Caution

- * Sometimes health effects are shown on the vertical axis and costs on the horizontal axis
- * Example: Stout et al. examined 64 different breast cancer screening scenarios (starting/stopping ages and screening frequency) for the U.S.
- * There are possible scenarios that dominate current U.S. screening practices

Cost-Effectiveness of Mammography Screening in the U.S



Source: Stout et al. J Natl Cancer Inst 2

Transformations of ICER

- * ICER may be transformed to:
 - Net Benefit (NB) = $E - (C * 1/V_s)$
 - Net Monetary Benefit (NMB) = $(V_s * E) - C$
- * V_s = the Social Value of health (e.g., \$100,000 per life-year, but this can also be varied)
- * Can be used to synthesize CEA and CBA through "Acceptance Curves"

The Social Value of Health

* Two Interpretations

- Value = "Willingness to Pay"

* \$7 Million per Death Averted (in U.S.)

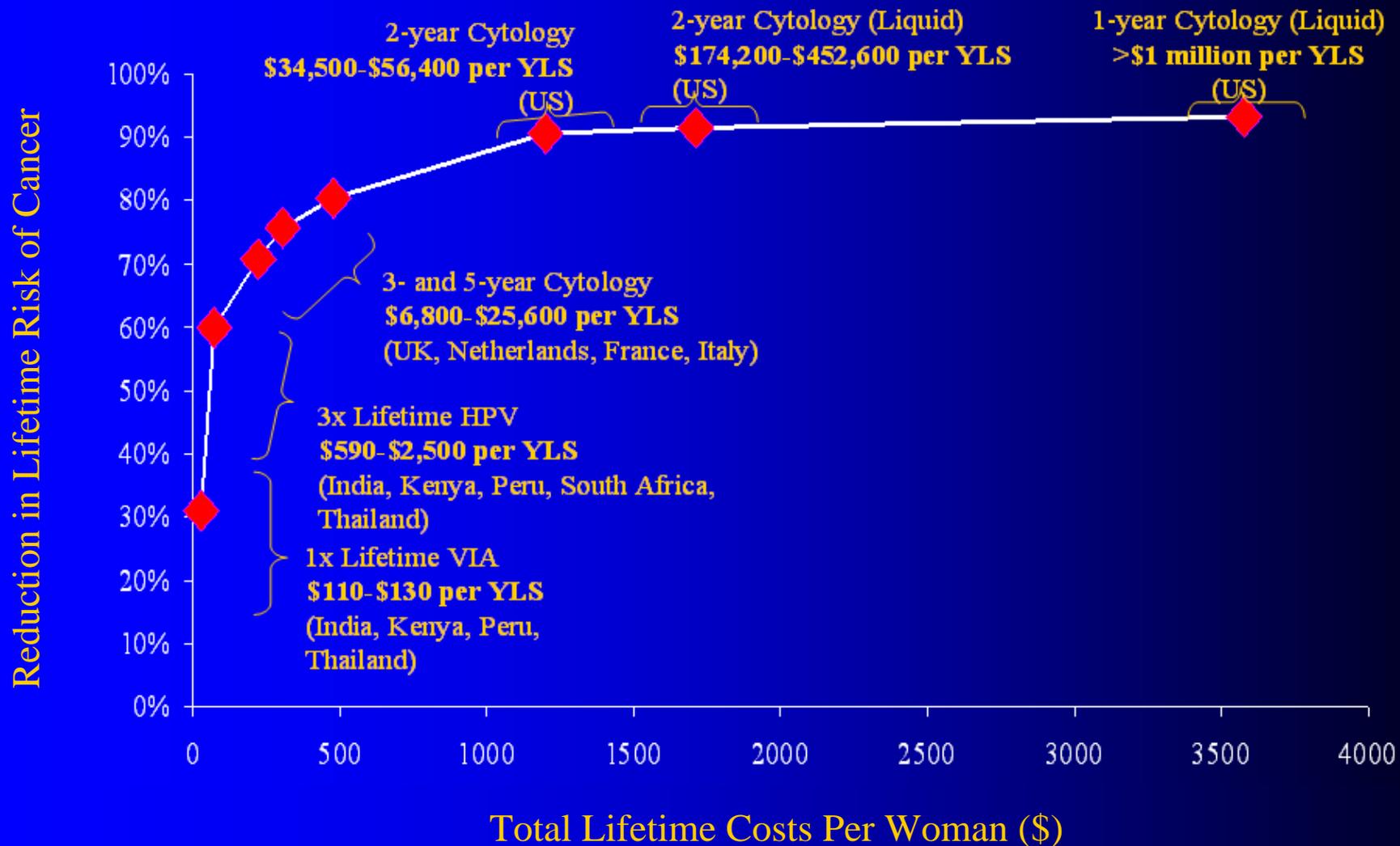
* \$150,000 per Life-Year (in U.S.)

- Value Reflects Nation's Healthcare Budget

* In UK, $V_s = \$50,000$ per Life-Year

* WHO Commission on Macroeconomics:
 $V_s = 1X - 3X$ per capita Gross Domestic Product per Life-Year

Cost Effectiveness of Cervical Cancer Screening Across National Settings



Comparison of Interventions

- * Micro-analysis – comparison is no treatment or status quo treatment
- * Marginal analysis – comparisons are different intensities of the same intervention
- * Macro-analysis
 - Comprehensive “league table,” e.g. Disease Control Priorities Project
 - Comparison against some value of V_s

Practical Considerations in Cost Evaluation Studies

- * Sources of Cost Data
- * Technical Economic Considerations
- * Statistical Considerations

Potential Sources of Economic Data

- * Clinical trial forms/medical record abstraction
- * Hospital bills
- * Health system cost-accounting systems (e.g. HMOs)
- * Administrative claims data (e.g. Medicare, Medstat)
- * Patient/provider survey (e.g. MEPS)
- * Cost scenario
- * Time-motion study
- * Engineering study

Some Technical Economic Issues

- * Adjusting for price (unit cost) differences
 - For different years
 - For different settings/locations
 - For different countries (currencies) (e.g. DCPP)
- * Discounting
- * Pricing non-market goods

Some Statistical Issues

- * Economic data are complex
- * Economic data tend to be highly skewed and censored
 - special estimation techniques have been developed
- * Trials designed for clinical end-points may be under-powered for economic and/or cost-effectiveness results
- * Cost-effectiveness or Cost-utility ratio estimates pose specific problems for analyzing and presenting confidence intervals (regions)

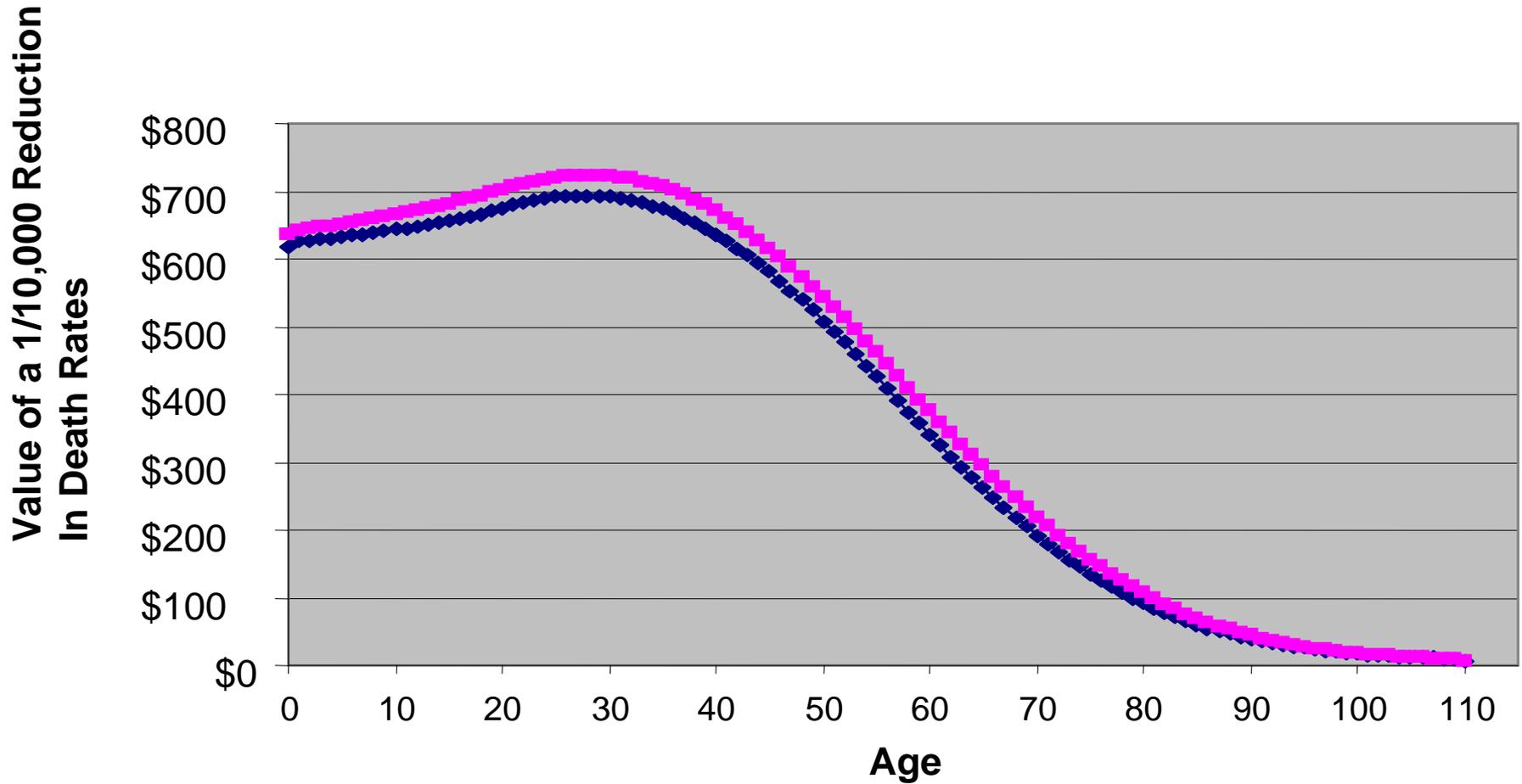
EXAMPLES

DESCRIPTIVE STUDIES

Economic Value of Health

- * The benefit side of the Cost-Benefit equation
- * Includes all aspects of health in monetary terms
- * One way to think about V_s

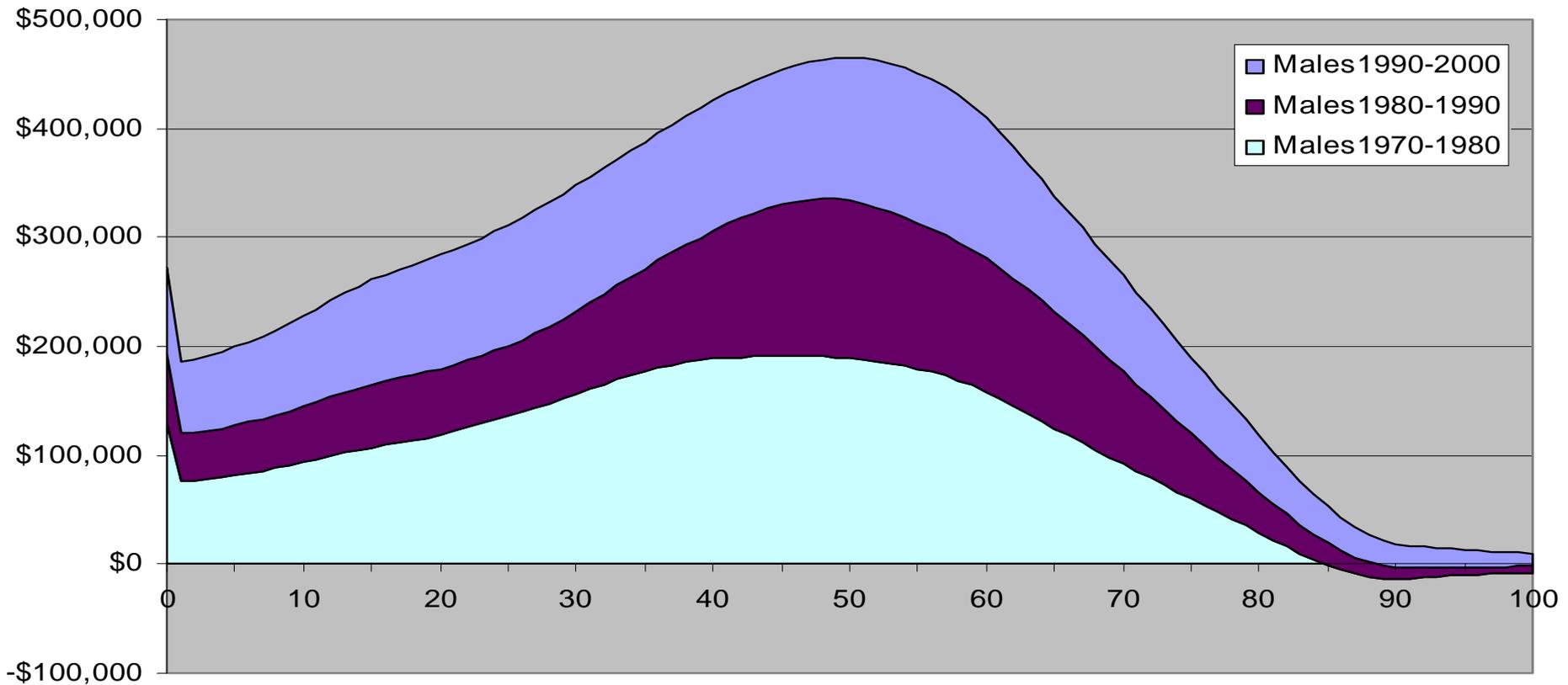
Economic Value of Health



Source: K.M. Murphy

Economic Gains From Increased Longevity - Males

Figure 6a. Gains from Increased Longevity for Males 1970-2000



Source: K.M. Murphy

Economic Value of Health Compared to Healthcare Expenditures: 1970 - 2000

- * Economic value of health improvements:
\$95,345 Billion
- * Healthcare expenditures:
\$34,725 Billion
- * Implies a favorable Cost-Benefit Ratio

BUT:

- * How much of health improvements are due to healthcare?

Cost Domains

- * Cost domains refers to categories of costs according to whether they are directly or indirectly related to the provision of marketed health care services.
- * The cost domain may also determine whether accessible and/or high quality cost data is available to the researcher and what degree of effort is required to obtain data.

Examples of Cost Domains

- * Direct health care costs (e.g. Medicare payments)
- * Direct non-health care costs (e.g., paid child care)
- * Patient time costs (e.g., value of time to attend treatment)
- * Morbidity costs (e.g., lost productivity due to work disability)
- * Mortality costs (e.g., lost productivity due to premature death)

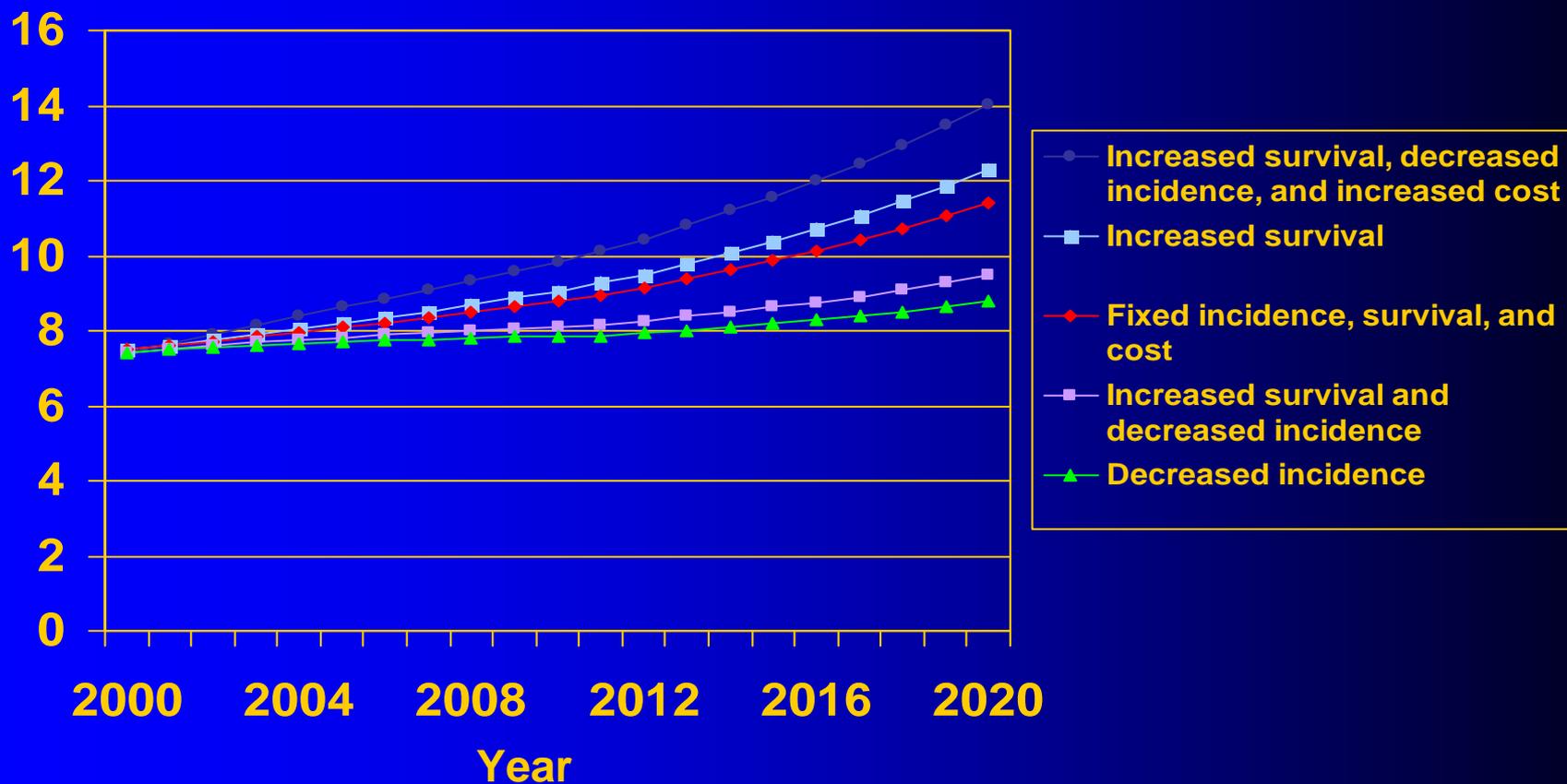
NIH Cost of Illness Report

Disease/Condition	Total Costs	Direct Costs	Indirect Costs
Alzheimer's Disease	\$87.9 billion	\$13.3 billion	\$74.6 billion
Atherosclerosis	\$5 billion	\$4.4 billion	\$0.6 billion
Cancer	\$96.1 billion	\$27.5 billion	\$68.7 billion
Stroke	\$43.3 billion	\$28.3 billion	\$15 billion
Liver Disease	\$3.2 billion	\$1.2 billion	\$2.1 billion
Pulmonary Disease	\$37.3 billion	\$21.6 billion	\$16.2 billion
Diabetes	\$98.2 billion	\$44.1 billion	\$54.1 billion
Heart Disease	\$175.3 billion	\$97.9 billion	\$77.4 billion
HIV/AIDS	NA	\$10.3 billion	NA
Homicide	\$33.7 billion	\$10.4 billion	\$23.3 billion
Injury	\$338 billion	\$89 billion	\$248 billion
Kidney Disease	\$40.3 billion	\$26.2 billion	\$14.1 billion
Pneumonia/Influenza	\$22.9 billion	\$17.5 billion	\$5.4 billion
Septicemia	\$7.2 billion	\$4.9 billion	\$2.3 billion
Suicide	NA	NA	\$10.2 billion

Source: H. Varmus, *Disease Specific Estimates of Direct and Indirect Costs of Illness and NIH Support*

Projected Treatment Costs for Colorectal Cancer 2000-2020

Dollars, in billions



Source: Yabroff KR, et al. Health Economics 2007.

Time Cost: Initial Treatment for Colorectal Cancer

	Category of Service	Visits		Time (hours)	
		Cases	Controls	Cases	Controls
		Initial Phase	Office visits	16.95	5.98
	Emergency room visits	0.62	0.35	2.17	1.22
	Chemotherapy	6.61	0.05	22.78	0.17
	Radiation therapy	1.43	0.04	2.34	0.06
	Hospitalization LOS	17.96	1.89	294.90	37.82
	Out-patient surgery	1.17	0.25	6.18	1.30
	Initial Phase Total**			355.02	49.26

Source: Yabroff et al. Medical Care, 2005.

How do time costs compare to direct costs?

- * For colorectal cancer, time costs (valued by average wage rates) during initial treatment were \$4655, 20% of direct medical expenditures in that period.

COST BENEFIT ANALYSIS

Using Cost of Illness in a Cost-Benefit Analysis

- * COI of Neural Tube Defects at Birth
- * Cost-Benefit Analysis of Folic Acid Fortification

COI of Neural Tube Defects

* Cost Domains Included in COI Estimate:

- Medical care
- Developmental services
- Special education
- Morbidity cost

* COI per case:

- Spina bifida: \$349,133
- Anencephaly: \$485,016

Cost-Benefit Analysis of Folic Acid Fortification

- * Cost of low level folic acid fortification = \$27.94 million per yr
- * Note: folic acid fortification can "mask" vitamin B12 deficiency
- * Cost of surveillance of those with undiagnosed vitamin B12 deficiency- \$5 million per year

Benefits of Folic Acid Fortification

- * Proportion of target population with inadequate folate intake - 66%
- * Cases of birth defects averted: 191 spina bifida, 113 anencephaly

Folic Acid Fortification: Cost-Benefit Analysis

- * Economic benefit of birth defects averted: \$121.5 million
- * Net benefit of fortification program = \$93.6 million
- * Benefit/Cost Ratio = 4.3
- * Is supplementation a better policy?

Supplementation vs. Fortification

- * Cost-effectiveness analysis of folic acid supplementation vs. fortification
- * Both found to be cost-savings compared to doing nothing
- * Fortification (dominantly) cost-effective relative to supplementation

Did the Policy Work?

- * Studies of NTD prevalence pre- and post-fortification
- * More fortification than anticipated
- * Original estimate did not take dose-response into account
- * NTDs averted:
 - 520 Spina Bifida
 - 92 Anencephaly
- * Net Benefit = \$143 million (in 2002\$)

Additional Benefits of Folic Acid?

- * New evidence that folic acid prevents cardiovascular disease and colon cancer [but next slide!]
- * Cost-effectiveness estimates of fortification at 140, 350 and 700 mcg per 100 grams of grain
- * 700 mcg most cost effective – 322,940 QALYs gained, \$4.4 billion saved per year

Economic Evaluation (Like All Good Science) Never Ends!

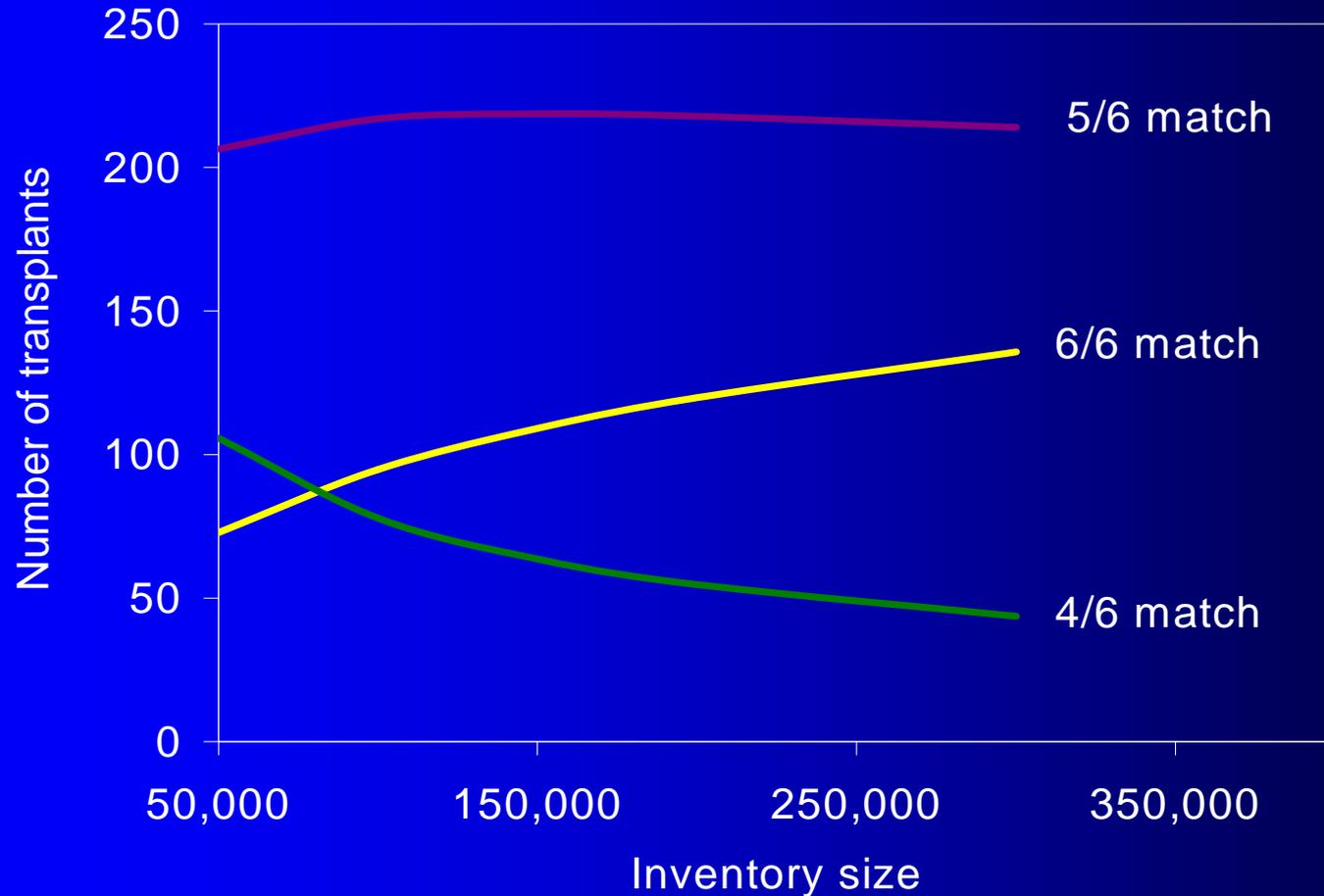
- * Are there un-discovered harms from folic acid fortification?
 - Negative results of folic acid prevention trial for adenomas (Cole BF, et. al. JAMA 2007;297:2351-9)
 - Association of increased colon cancer incidence (above trend) with start of folic acid fortification (Mason, et. Al. Ca Epi Bio Prev 2007;16:1325-9)

MARGINAL COST ANALYSIS

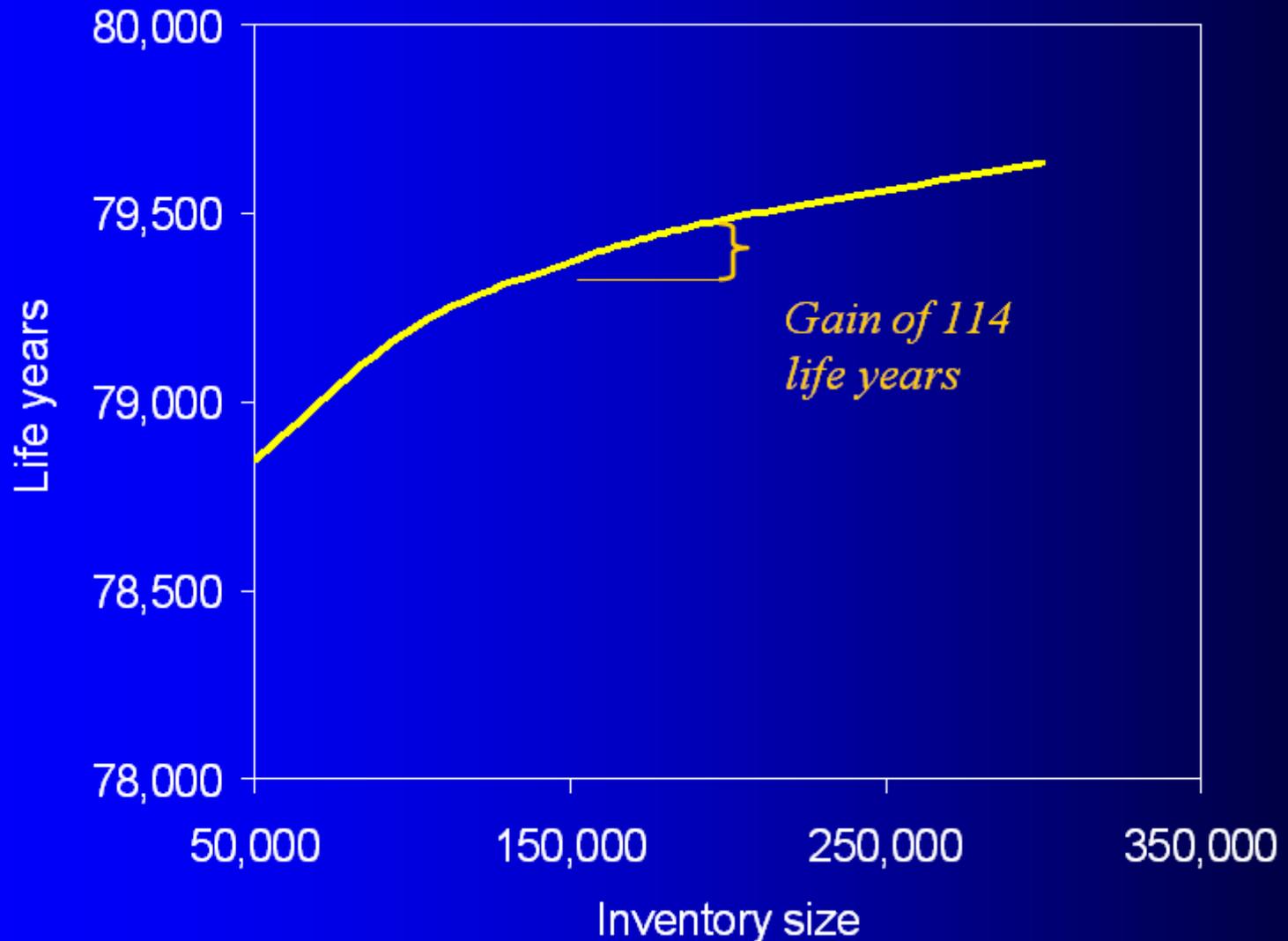
Marginal Analysis National Cord Blood Bank

- * How large should National Cord Blood Bank be (beyond 50,000 inventory initially proposed)?
- * Human leukocyte antigen match rate (and therefore transplant benefit) increases with size
- * Cost increases with size

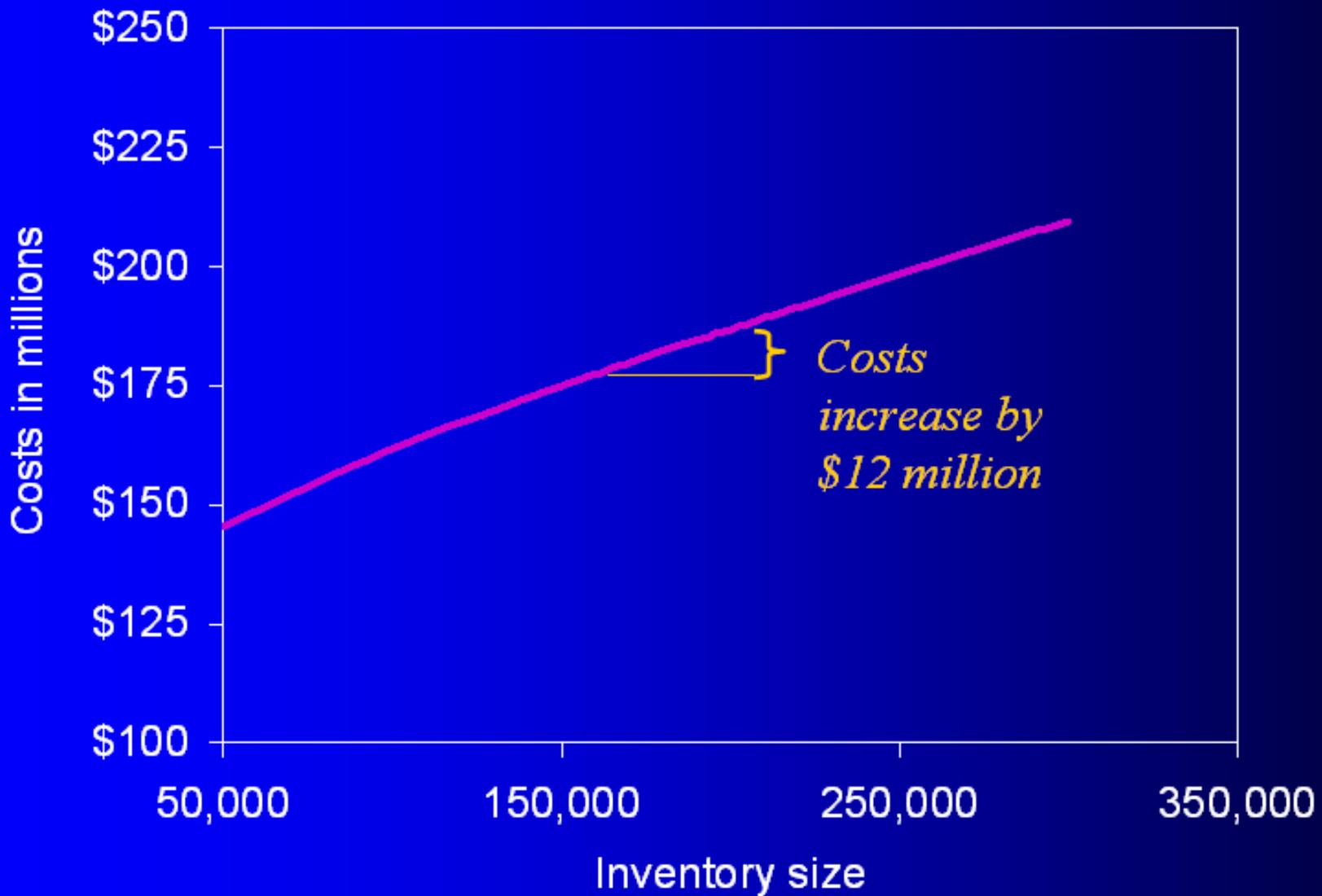
Cord transplants by match level for patients age <20 as a function of inventory



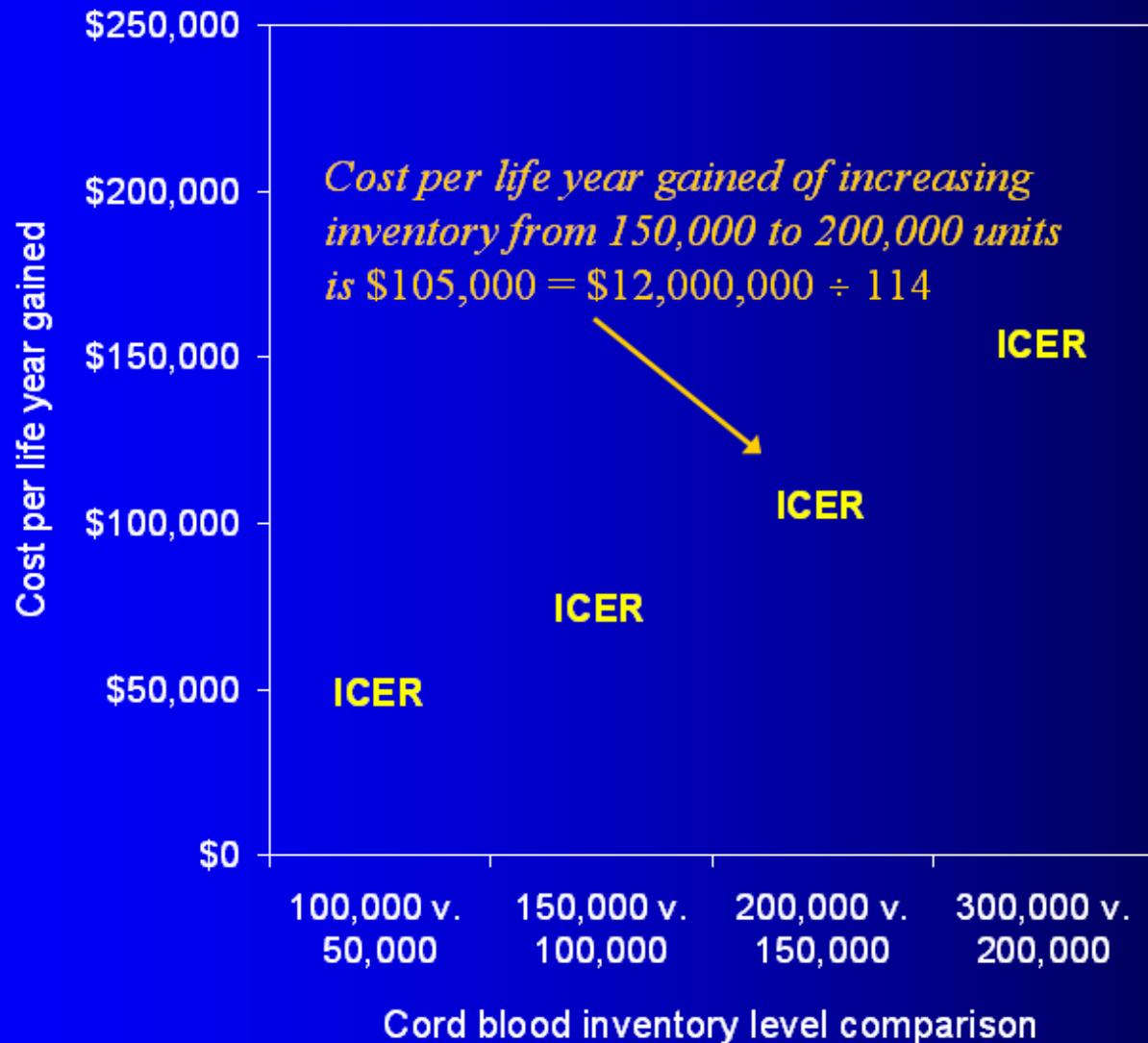
Patient life years as a function of inventory



Total bank costs as a function of inventory



Incremental cost-effectiveness ratios (ICER)



Cost Evaluation Studies and Clinical Trials

- * Clinical Trial with Economic Follow-up
- * Economic Modeling Combined with Clinical Trial Results
- * Linking to HMO Computerized Data on Patient Care Costs
- * Analyzing the Confidence Region for the Cost-Effectiveness Ratio

Clinical Trial with Economic Follow-up

- * Hlatky MA, et al. Medical costs and quality of life 10 to 12 years after randomization to angioplasty or bypass surgery for multivessel coronary artery disease. *Circulation* 2004;110:1960-1966.
- * Bypass Angioplasty Revascularization Investigation (BARI) trial
- * Study of Economics and Quality of Life (SEQOL) follow-up study

BARI / SEQOL Study

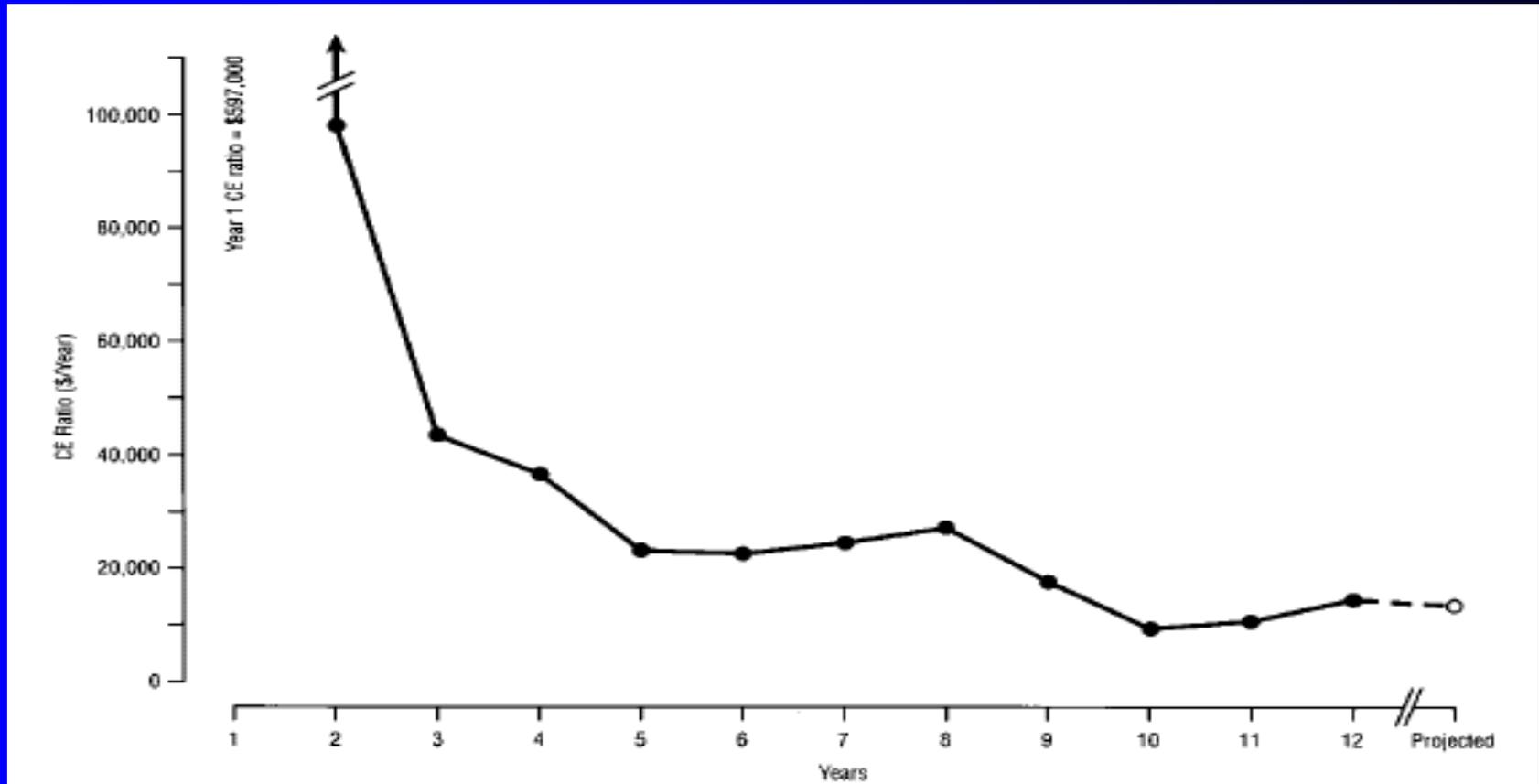
* BARI Study

- Randomization: 1988-1991
- Follow-up through 1996

* SEQOL Study

- Sub-sample follow-up through 2001
- Medical costs
- Quality of life measures

Cost-Effectiveness of CABG vs PTCA by Year of Follow-up



Source: Hlatky et al. *Circulation* 2004

Lesson of the BARI/SEQOL Study

“The improvement in the cost-effectiveness ratio over time was largely because of the narrowing of the cost differential between the 2 procedures, with the remainder resulting from a small survival advantage among CABG patients. These observations underscore the importance of a long-term perspective in economic evaluation, as an initially costly procedure may prove cost-effective over the long term if it either provides extended clinical benefits or the initially higher cost can be offset by preventing subsequent hospitalizations.”

Economic Modeling Combined With Clinical Trial Results

- * Berthelot JM et al. Decision framework for chemotherapeutic interventions for metastatic non-small-cell lung cancer. *Journal of the National Cancer Institute*, 2000 Aug 16;92(16):1321-9
- * Combining trial data with a modeling approach to costs and longer term outcomes

CEA of Lung Cancer Treatment

- * CEA of chemotherapy vs. best supportive care for advanced stage lung cancer
- * Survival benefits of treatment based on survival curves modeled (out to 48 months using Weibull survival function) from RCT results and community survival data
- * Costs based on Canadian cost scenarios (POHEM model)

Results for stage IV NSCLC

- * Total cost of best supportive care = \$25,904
- * Total cost of chemotherapy = \$25,105 - \$41,576 depending on regimen
- * Hospital/Clinic costs are higher for best supportive care compared to chemotherapy – e.g. intervention results in down-stream cost savings
- * ICER of chemotherapy ranges from cost-savings to \$37,800 / quality adjusted life year

CEA From Patient Level Data in a Clinical Trial

- * Statistical modeling of longterm outcomes and costs
- * Accounting for stochastic uncertainty in the measurment of health outcomes and costs – Bootstrap Confidence Interval
- * Assessing CEA as a function of the Social Value of health – the Cost Effective Acceptability Curve

CEA of Hormonal Treatment for Prostate Cancer

- * Cost-effectiveness analysis of adding early hormonal therapy to radiotherapy for locally advanced prostate cancer
- * Clinical Trial – EORTC 22863

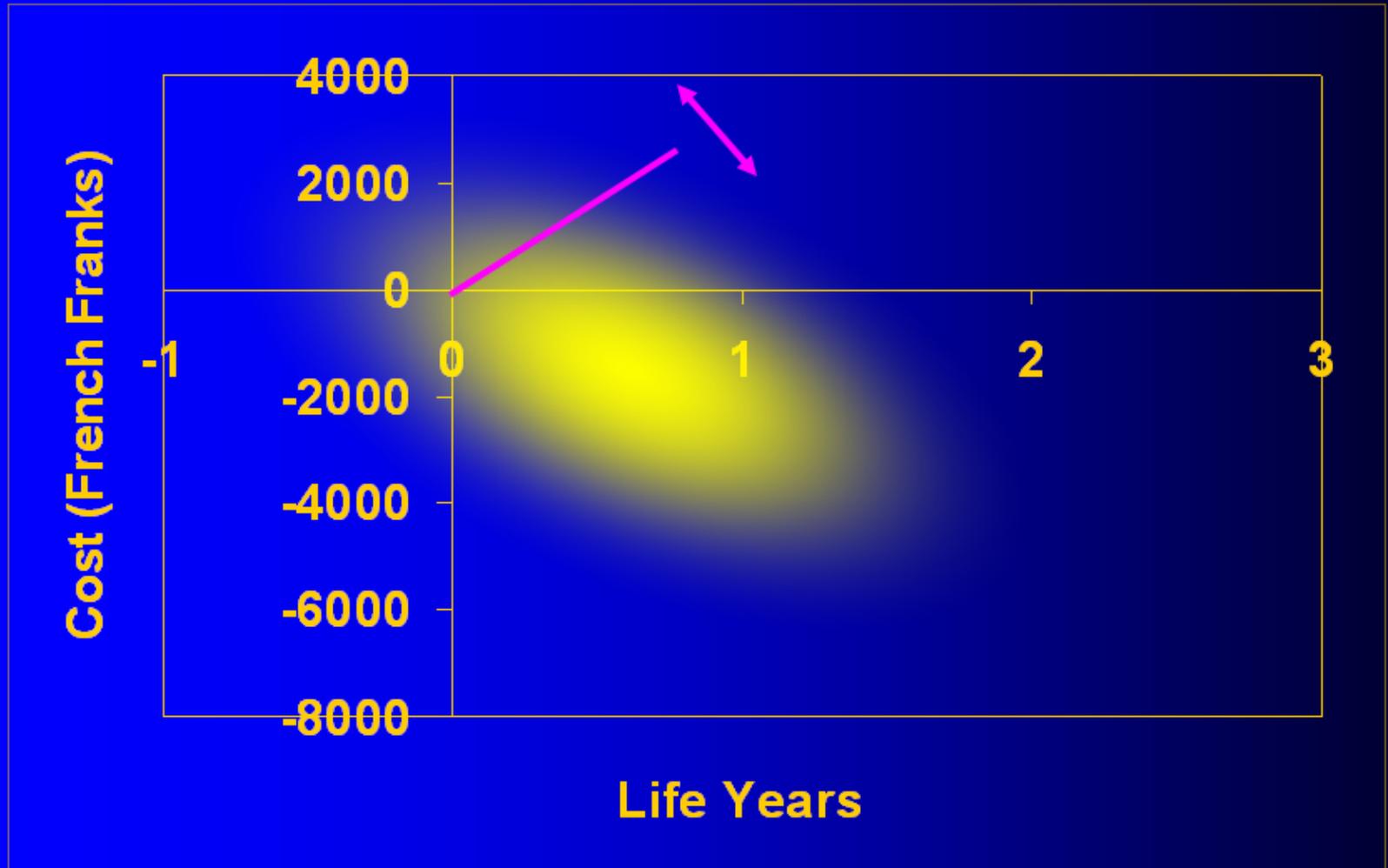
Methods

- * Direct medical resource use obtained of 90 subjects for up to 11 years of follow-up
- * Unit costs based on based on standard national French tariffs
- * Method of Lin et al. used to adjust for censoring in longitudinal cost data
- * Mean survival estimated using the restricted means method

Construction of 95% Confidence Region

- * Using standard Monte Carlo simulation methods, (sampling with replacement) 5000 replications of the Incremental Cost Effectiveness Ratio, were calculated

CEA of Hormonal Treatment Confidence Region



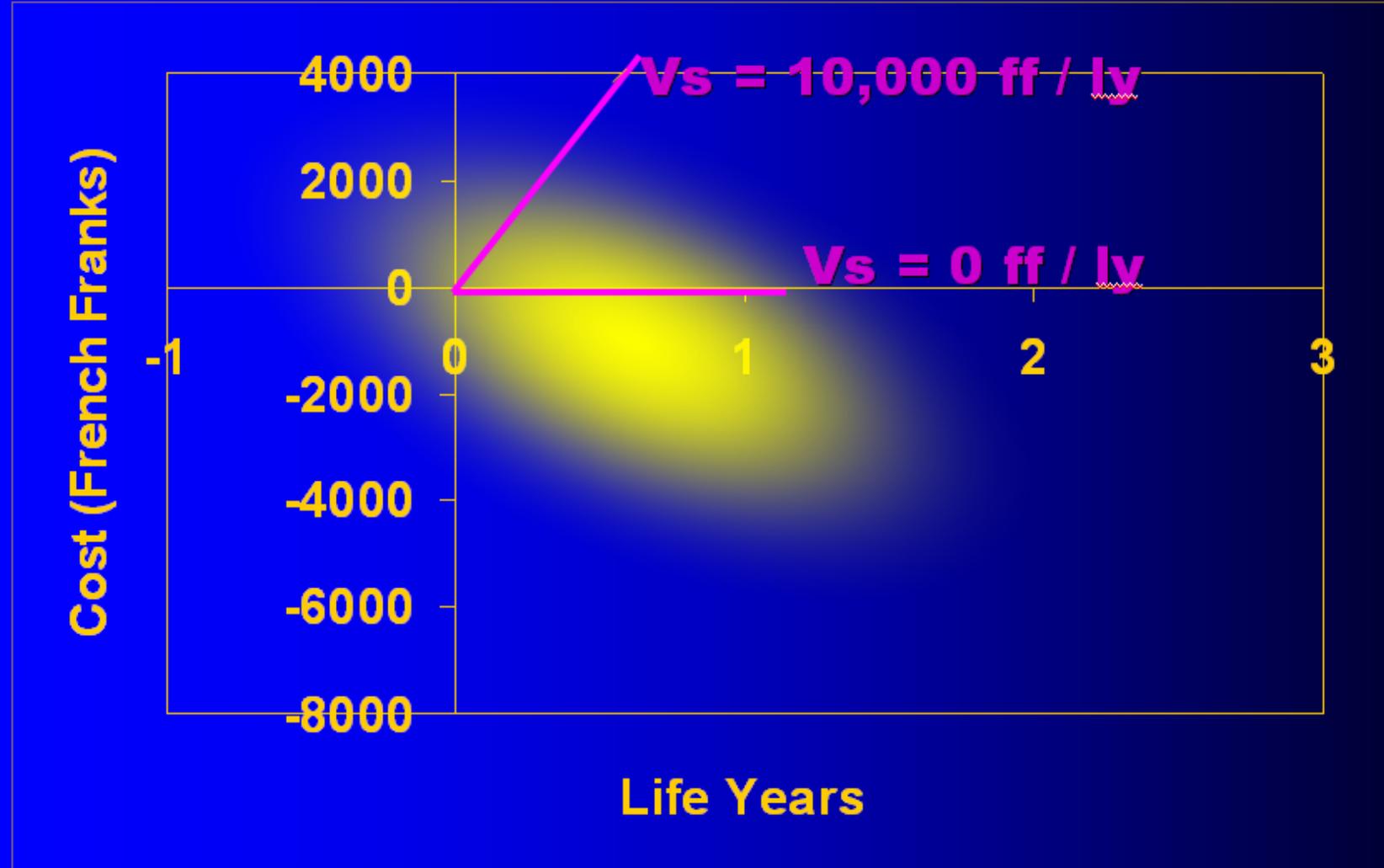
Cost Effectiveness Acceptability Curves

- * Conduct bootstrap simulation
- * Examine all results that fall within 95% confidence intervals for the cost effectiveness ratio
- * Compare to reference values for social value of health (V_s)
- * Calculate probability that:
 $CER < V_s$

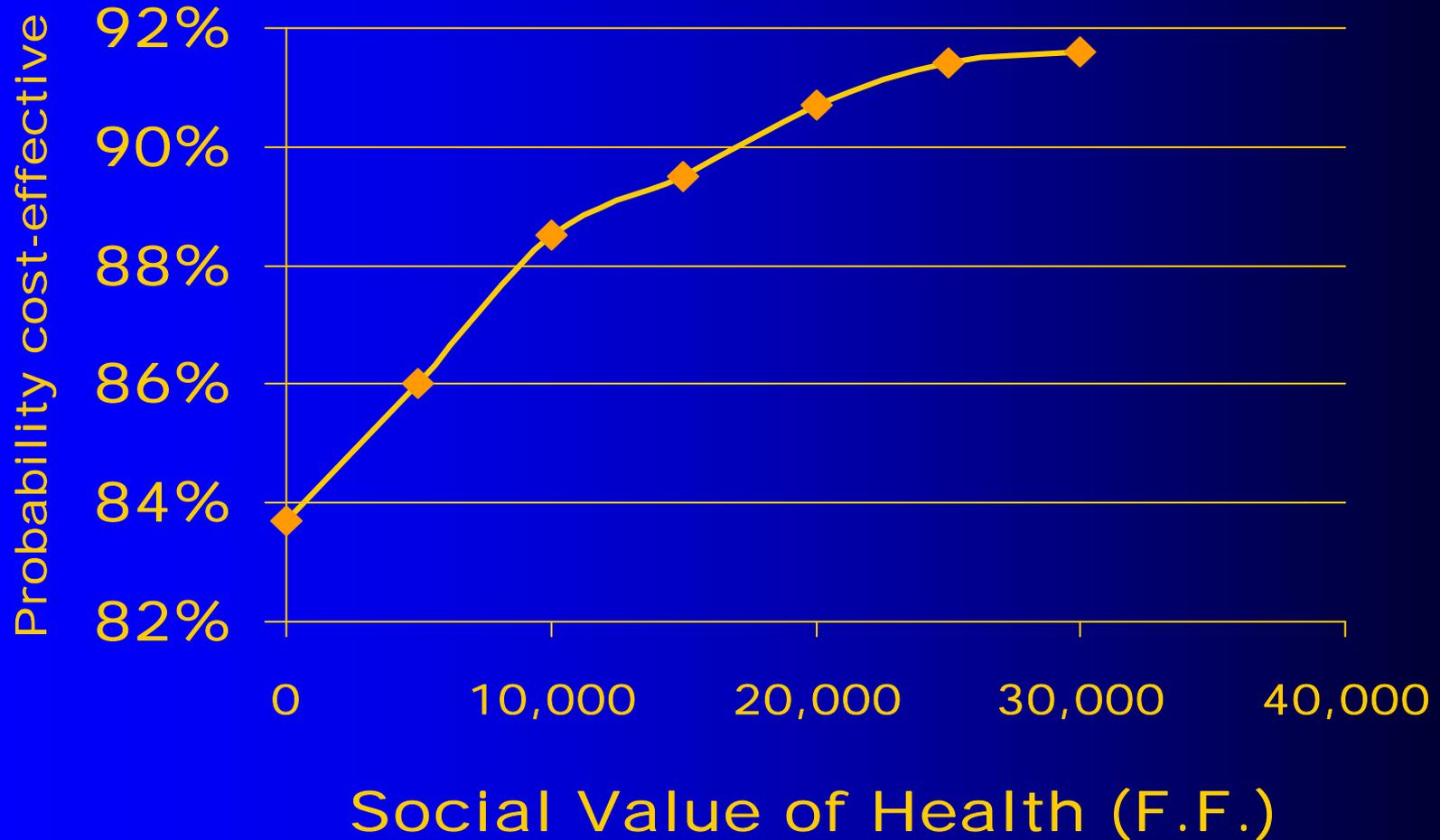
Graphical Analysis

- * E.g., Rotate pink cost-effectiveness line – this is equivalent to varying the value of V_s
- * Assess what proportion of confidence ellipse lies beneath the line

CEA of Hormonal Treatment Confidence Region



Cost-Effectiveness Acceptability Curve for Hormonal Treatment of Prostate Cancer



Sources of Uncertainty in Economic Evaluation

- * Parameter Uncertainty
 - biological, demographic, epidemiological, medical and economic parameters (as in example just shown)
- * Model Uncertainty
 - Choice of model type and structure
- * Methodological Uncertainty
 - Choice of CBA, CEA, CUA, perspective, time horizon, etc.