

IN THE NAME OF GOD

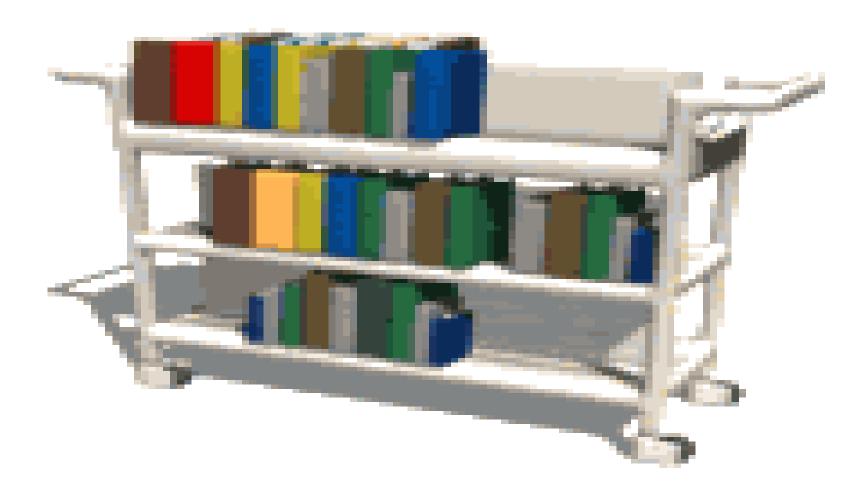
Evidence – Based Medicine

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What do you think about EBM?





Minimum reading to keep up-todate with pediatrics

- Pediatrics 40 articles x 12 months
- New England Journal of Medicine 5 articles x 52 weeks
- Lancet 6 articles x 52 weeks
- Journal of Pediatrics 18 articles x 12 months
- Pediatric Infectious Disease Journal 15 articles x 12 months
- JAMA 8 articles x 12 months
- BMJ 10 articles x 52 months
- Archives of Pediatric and Adolescent Medicine 10 articles x 12 months
- 1694 article per year= 5 articles per day



- MIDDEL 19 CENTURY IN FRANCE MEDICAL SCHOOL
- EBM WAS CREATED M.C MASTER UNIVERSITY 1980
- It was initially proposed by <u>Dr. David</u> <u>Sackett</u> and colleagues at McMasters University in Ontario, Canada.

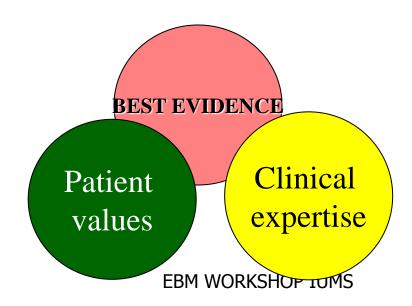


Evidence-based medicine (EBM) is an important change in the way physicians practice, teach, and do research.

DEFINITION

INTEGRATION OF CLINICAL EXPERIENCE WITH THE BEST EVIDENCE PROVIDED BY SYSTEMATIC AND OBJECTIVE — ORIENTED RESEARCH

EBM MODEL







DEFINITION

CONSCIENTIOUS, EXPLICIT
&JUDICIOUS USE OF CURRENT BEST
EVIDENCE IN MAKING DECISIONS
ABOUT CARE OF INDIVIDUAL
PATIENTS OR THE DELIVERY OF
HEALTH SERVICES DAVID SACKETT.



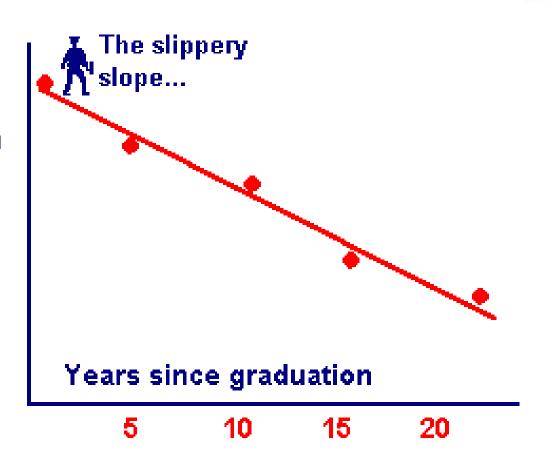
EBM OBJECTIVES

- KEEPING YOUR SKILLS UP TO DATE
 - -MEMORY DECREASE
 - -NEW TREATMENT METHODS
- SAVING TIME
- SAVING LIVES
- SUPPLEMENTING CLINICAL JUDGEMENT(EBM MODEL)



Knowledge of best hypertension care

Shiri et al, CMAJ, 1993





- قابل آموزش به پزشکان در سطوح مختلف
- پر کردن شکاف بین تحقیقات بالینی و بکار گیری نتایج آنها
 - تقویت آموزش مستقل و خود محور
 - تقویت بحث گروهی
 - روز آمد کردن اطلاعات پزشکان
 - درك عميق روش تحقيق توسط متخصصين باليني



- افزایش اعتماد به نفس پزشکان بالینی در اخذ تصمیم بالینی
 - افزایش توانائی پزشکان در جستجوی اطلاعات
 - عادت به مطالعه را در پزشکان می افزاید
- امکان توجیه منطقی تصمیمات درمانی را برای بیماران فراهم می کند.
 - طراحی دستور العمل مشتر ک برای تصمیمات بالینی توسط متخصصین محلی



مضرات EBM

■ آموزش و بكارگيرى EBM در بالين وقت گير است

■ هزینه فراهم سازی امکانات زیاد است

■ کاهش اعتماد به نفس پزشکان در مقابل اطلاعات جدید و اقدامات فعلی آنها

EBM Method

Assess your patient



Ask clinical questions

Acquire the best evidence



Appraise the evidence



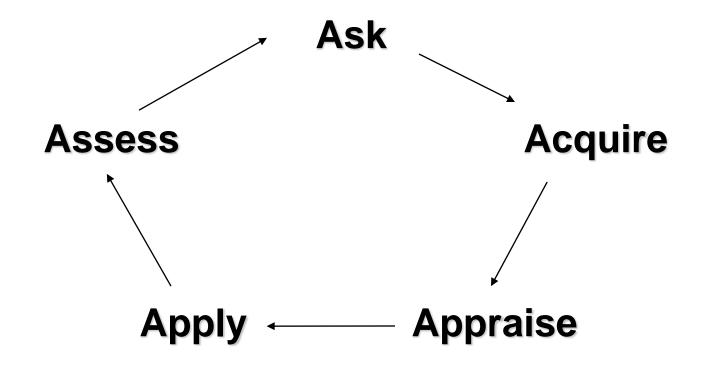
Apply evidence to patient care

EBM PROCESS



- 1. PATIENT PROBLEM
- 2. CLINICAL QUESTION
- 3. SEARCH FOR EVIDENCE
- 4. CRITICAL APPRAISAL OF THE EVIDENCE
- 5. APPLYING THE RESULTS INTO PRACTICE (CURRENT PATIENT)

Evidence-based Practice

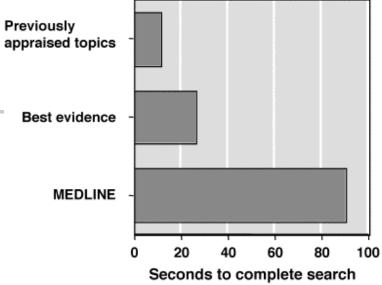


The Practice of EBM

- Step 1: Asking an answerable question
- Step 2: Tracking down the best evidence to answer that question
- Step 3: Critically appraise the evidence for validity, size of the effect, and utility of the findings
- Step 4: Incorporate the clinical appraisal into our clinical expertise and patient's individual issues
- Step 5: Evaluate and improve steps 1-4 with each new opportunity to apply these principles

Time to complete searches on the evidence cart

Evidence Cart





Dave Sackett 3/11/2017



EBM WORKSHOP IUMS



Domains of EBM

- TREATMENT
- PROGNOSIS
- DIAGNOSIS
- ETIOLOGY/CAUSATION/HARM

Types of Clinical Questions

By Content

- Diagnosis
- Therapy
- Etiology
- Prognosis

By Format

Background

Foreground

Good clinical questions

"Background" Questions

- General knowledge
- Two components
 - Root (who, what, when, where, why)
 - A disorder or aspect of a disorder
- E.g., "What is the typical age of onset of bipolar disorder?"
- "How do I decide to use a typical vs. atypical antipsychotic for agitation?"



Good clinical questions

"Foreground" Questions

- These ask for specific information about managing a patient with a disorder
- They have 3-4 essential components

COMPONENTS OF CLINICAL QUESTIONS

P - patient and problem(population)

I - intervention(treatment, test, prognosis...)

C - comparison

O - outcome



Diagnosis

"In patients with suspected pulmonary fibrosis, how does high-resolution CT compare with lung biopsy for establishing the diagnosis?"

P = Pulmonary fibrosis

= High-resolution CT

C = Lung biopsy

O = Sensitivity/specificity/PVs/LRs



Etiology

"Do obstetrical complications during pregnancy increase the likelihood of schizophrenia in the child?"

- P = Pregnant females
- I = Obstetrical complications
- **C** = No obstetrical complications
- O = Childhood schizophrenia

Prognosis

"In patients with acute leukemia, is a normal white cell count at the time of diagnosis an independent predictor of disease-free survival?"

- P = Acute leukemia
- = Normal white cell count
- C = Abnormal white cell count
- O = Disease-free survival

Ask Clinical Questions

Components of Clinical Questions

Patient/
Population

Intervention/ Exposure

Comparison

Outcome

In patients with acute MI

In women with suspected coronary disease

In postmenopausal women does early treatment with a statin

what is the accuracy of exercise ECHO

does hormone replacement therapy

compared to placebo

compared to exercise ECG

compared to no HRaT

decrease cardiovascular mortality?

for diagnosing significant CAD?

increase the risk of breast cancer?

Clinical question(scenario) for treatment

- P –in a child with frequent febrile seizures
- I would anticonvulsant therapy

C – compared to no treatment

O – results in seizure reduction

Question for diagnosis



- P in an otherwise healthy 15 yrs old boy with sore throat
- I- how does the clinical exam

C- compare to throat culture

O- In diagnosing GAS infection ?



Question Prognosis

P- In children with Down syndrome

I - Is IQ an important prognostic factor

C

In predicting Alzheimer's later in life



Etiology/Harm

- P -controlling for confounding factors, do otherwise healthy children
- I -exposed in utero to cocaine
- C compared to children not exposed
- O have increased incidence of learning disabilities at age six years?

Type of Question

Suggested best type of Study



Prevention	RCT>cohort study > case control > case series
Prognosis	cohort study > case control > case series
Etiology/Harm	RCT > cohort > case control > case series
Diagnosis	prospective, blind comparison to a gold standard
Therapy	RCT>cohort > case control > case series

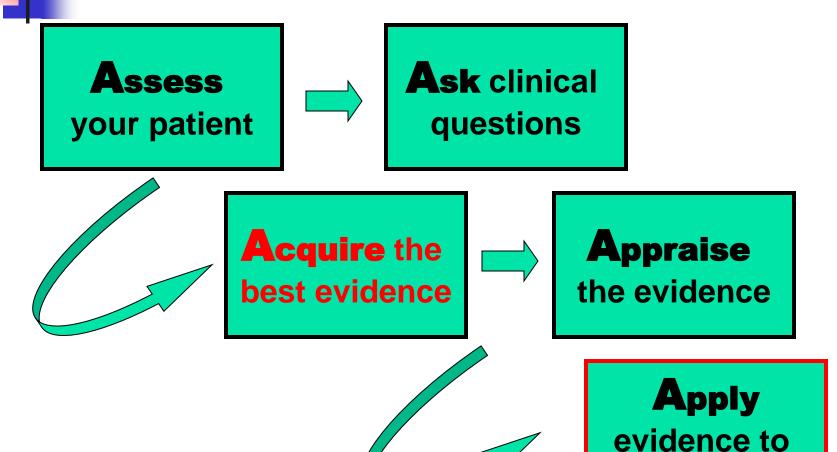




THANK YOU ANY QUESTIONS?

EBM Method

WORKSHOP IUMS



patient care



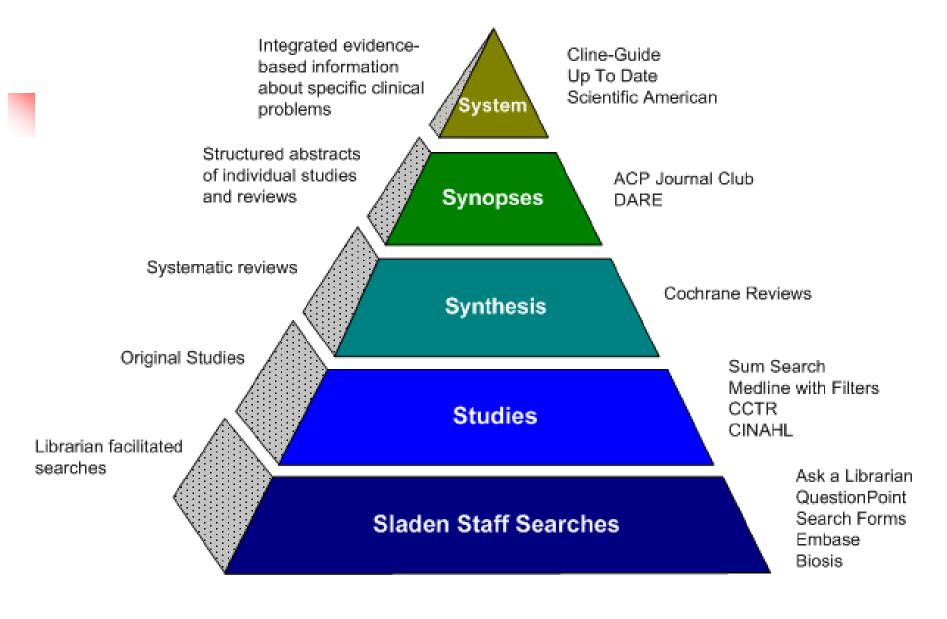
How to Learn About Best Information Resources?

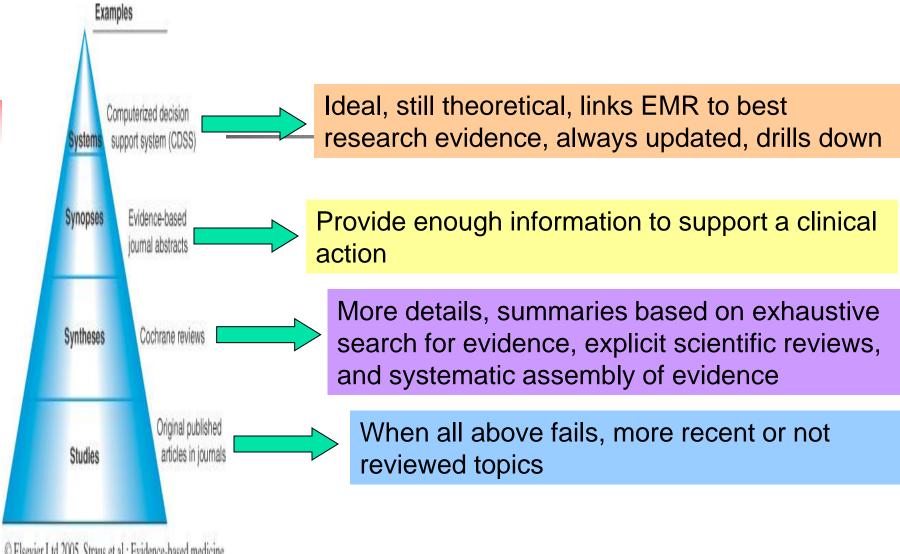
- From librarians (hands-on training)
- From experts in medical informatics
- Courses/ Tutorials

Searching for Answers: The "4S" Approach of Haynes

Haynes RB: EBMH 2001;4:47 and ACP Journal Club 2001;134:A11

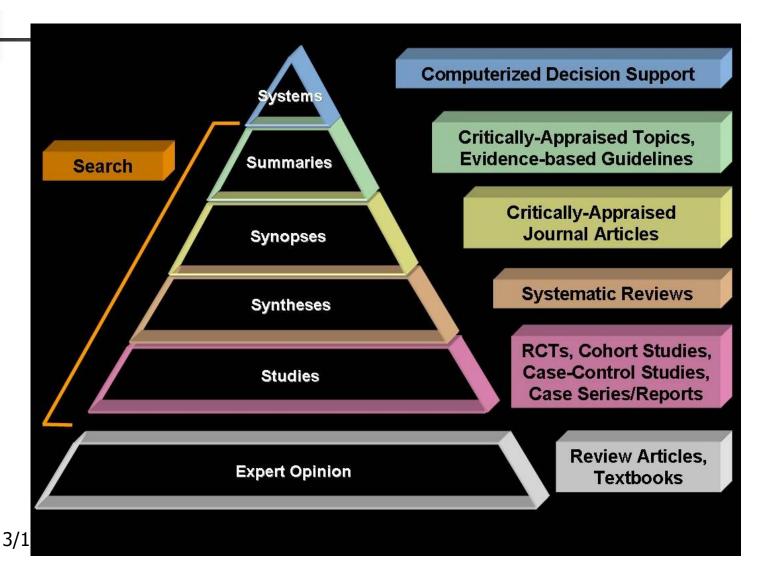
- Systems (comprehensive resources)
 - Clinical Evidence (<u>www.clinicalevidence.com</u>)
 - Collection of evidence-based guidelines
- Synopses (structured abstracts)
 - Evidence-Based Mental Health (http://ebmh.bmjjournals.com/)
 - ACP Journal Club (<u>www.acpjc.org</u>)
- Syntheses (systematic reviews)
 - Cochrane Database (OVID)
 - DARE (http://agatha.york.ac.uk/darehp.htm)
- Studies (original research)





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EBM hierarchy Haynes 5S pyramid



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Systems

- Clinical Evidence (BMJ)
 - URL: http://www.clinicalevidence.com
 - Contains limited range of clinical questions
 - PIER (the Physician's Information and Education Resource) by ACP
 - URL: http://pier.acponline.org
 - Only for members
 - UpToDate®
 - URL: http://www.uptodate.com
 - Updated quarterly
 - Extensively referenced
 - ACP Medicine (Formerly Scientific American Medicine)
 - URL: http://www.acpmedicine.com

Systems (Cont'ed)

- Harrison's Principles of Internal Medicine
 - URL: http://www.harrisonsmed.com
 - Only updated every 3 years
- Evidence Based on Call
 - URL: http://www.eboncall.org/content.jsp.htm
- Evidence-Based Pediatrics and Child Health
 - URL: http://www.evidbasedpediatrics.com
- Evidence Based Cardiology
 - URL: http://www.evidencebasedcardiology.com/

OVID includes and links EBMR (Cochrane, ACP Journal Club, the Database of Abstracts of Reviews of Exidences (DARE), and Medline

Criteria to evaluate systems

Look for Systems that:

- Are revised at least once a year: Date of revision should be listed
- Select and appraise the evidence in an explicit way (Introduction)
- Site evidence to support clinical care declarations

Synopses

- Published in secondary journals
 - Select only high-quality original research and review articles
 - Use explicit quality criteria for selection
 - Appraise for validity
 - Prepare structured, "value-added" abstract
 - Accompanying commentary
 - Declarative title that gives "bottom line"



- ACP Journal Club http://www.acpjc.org/
- Give you the summary and links you to the evidence
- Ex: "Low Molecular Weight Heparin is Effective and Safe in the Acute Coronary Syndromes"

Syntheses: Systematic Reviews

- What makes a review systematic?
 - Comprehensive search
 - Use only high-quality studies
 - Summarize results
- Sources of systematic reviews
 - Cochrane Library (available through OVID)
 - Database of Abstracts of Systematic Reviews
 (DARE): http://agatha.york.ac.uk/darehp.htm



Cochrane Library

URL: http://www.cochranelibrary.com/

OVID's EBMR

 (Includes ACP Journal Club, Cochrane Database of Systematic Reviews (CDSR), and DARE)

Studies

Specialized

ACP Journal Club:

www.acpjc.org

Evidence Based Medicine:

> <u>www.ebm.bmjjournals.co</u> <u>m</u>

Evidence Based Nursing:

> www.ebn.bmjjournals.co m

Evidence Based Mental Health:

www.ebmh.bmjjournals.c

General

- Cochrane Central Register of Controlled Trials (Therapy)
- MEDLINE:

http://www.ncbi.nlm.ni h.gov/PubMed/

- Using the Clinical Queries Search
- ASK MEDLINE

http://askmedline.nlm.n ih.gov/ask/ask.php



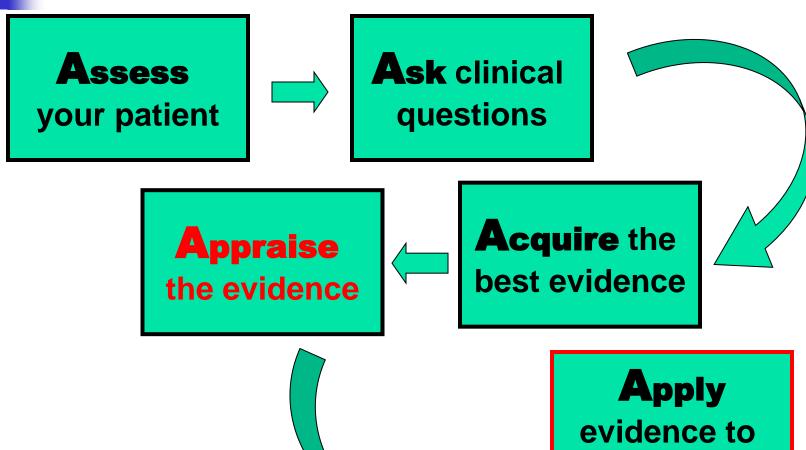
Textbooks are only useful for "background questions"

(Pathophysiology of clinical problems)

Alternatives to the "4S" Search Approach

- TRIP database (<u>www.tripdatabase.com</u>)
 - Searches Cochrane, DARE, collections of systematic reviews and guidelines, and some online journals
 - Links to PubMed clinical queries
- SUMSearch (http://sumsearch.uthsca.edu)
 - Searches MEDLINE, DARE, National Guidelines Clearinghouse
 - Takes longer than TRIP

EBM Method



3/11/2017

EBM WORKSHOP

patient care

CRITICAL APPRAISAL



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EDC, Iran University of Medical Sciences
3/11/2017 EBM WORKSHOP IUMS



CRITICAL APPRAISAL is the process of assessing and interpreting evidence, by systematically considering its validity, results and relevance to your own work

Critical Appraisal of Literature

Intended to enhance the clinician's skill to determine whether the results reported in an article were likely to be

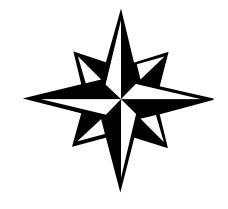


- ... true
- ... important
- ... applicable to their patients!



VALIDITY

RELIABILITY



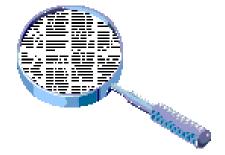
IMPORTANCE

Tools for Critical Appraisal

EBM "simplified" approach:

What are the results?

Are the results valid?



Will the results help me in patient care?



3 Important Questions

Are the results of the study valid?

What are the results?

Will the result help locally?

INTRODUCTION

In concise statement of the problem

Inadequate review of the literature

Weak study rationale

METHODS

- Inadequate sample size, nonrepresentative sample, or biases in subject selection or recruitment
- Inadequate controls (random assignment, or well-matched controls?)
- Measurement biases (valid tools? blinded? timing appropriate? follow-up?)



RESULTS

Selection and/or number of statistical tests performed

Selection of variables for inclusion

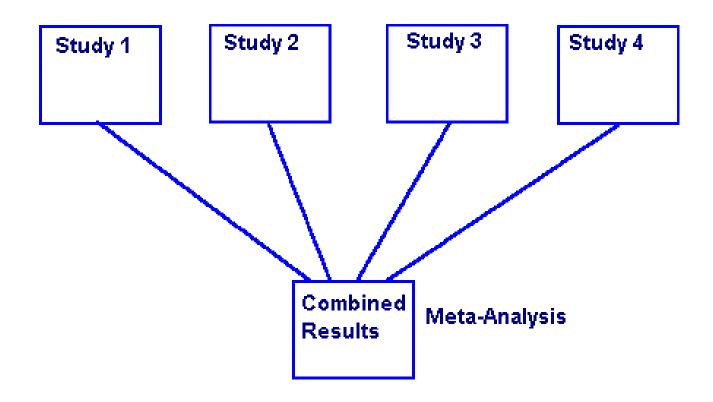
DISCUSSION

- Failure to link findings to current literature
- Inappropriate inferences
- Failure to critique own work
- Little insight or direction provided

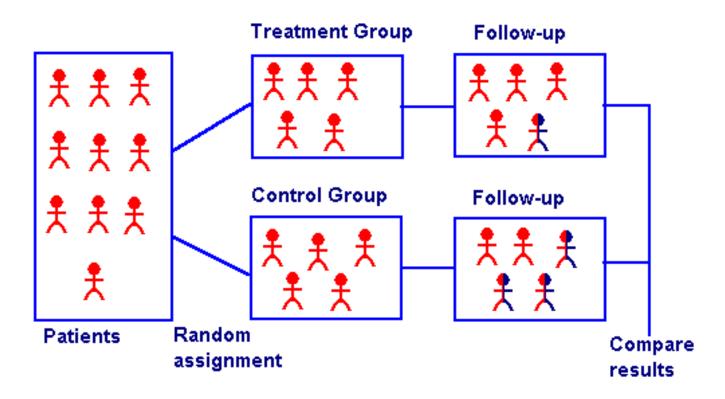




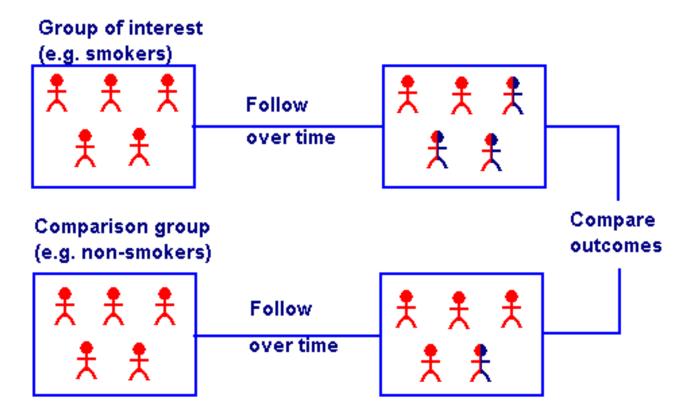
Systematic Reviews and Meta-Analyses



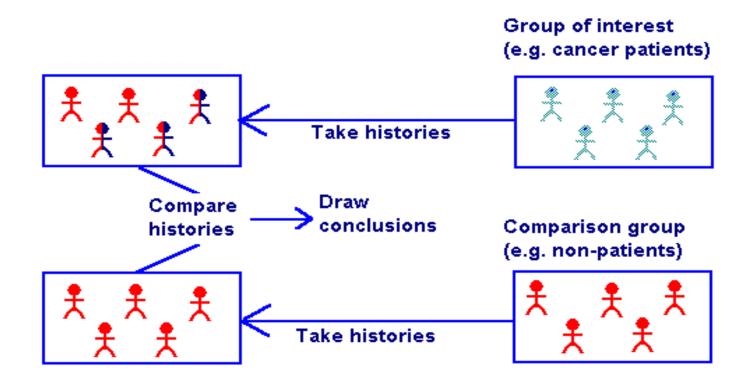
Randomized Controlled Studies



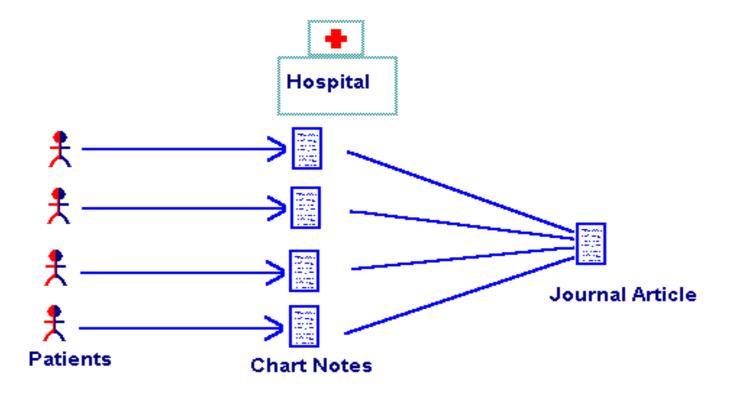
Cohort Studies



Case Control Studies



Case Series and Case Reports





Thank You! Any Question?

DIAGNOSIS: SEN,SPE,PPV.NPV.LR

PROGNOSIS: CI, SURVIVAL ANALYSIS

■ THERAPY: CER,EER,RRR,ARR,NNT

HARM: OR,RR,NNH

DIAGNOSIS

Sensitivity and Specificity

Positive and Negative Predictive Values

Likelihood Ratios

Sensitivity: the proportion of patients <u>with</u> the disease who have a <u>positive</u> test result

$$Se = P(T+ \mid D+)$$

Specificity: the proportion of patients <u>without</u> the disease who have a <u>negative</u> test result

$$Sp = P(T-|D-)$$

Information for a dichotomous test

Disease

Present

Absent

Test Result **Positive**

Negative

True positive

Α

False negative

C

False positive

В

True negative

D

Sensitivity = A / (A+C)

Specificity = D / (B+D)

A+C

B+D



Information for a dichotomous test

Disease

Present

Absent

Test Result **Positive**

Negative

True positive

A = 103

False negative

C = 12

False positive

B = 16

True negative

D = 211

Predictive values

PPV: the proportion of patients with a <u>positive</u> test result who <u>have</u> the disease

$$PPV=P(D+/T+)$$

NPV: the proportion of patients with a negative test result who do not have the disease

$$NPV=P(D-/T-)$$

Present Absent

Positive Test Result Negative Present Absent

True positive False positive B B A+B C+D C D

PPV = A / (A+B)

NPV = D / (C+D)

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Disease

Present

Absent

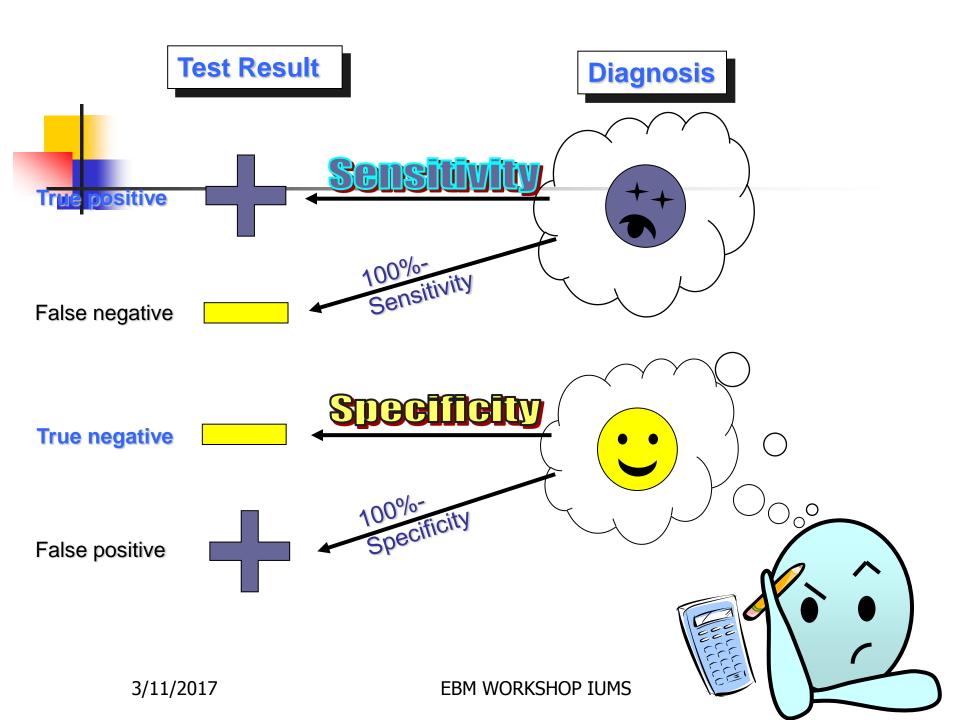
Test Result **Positive**

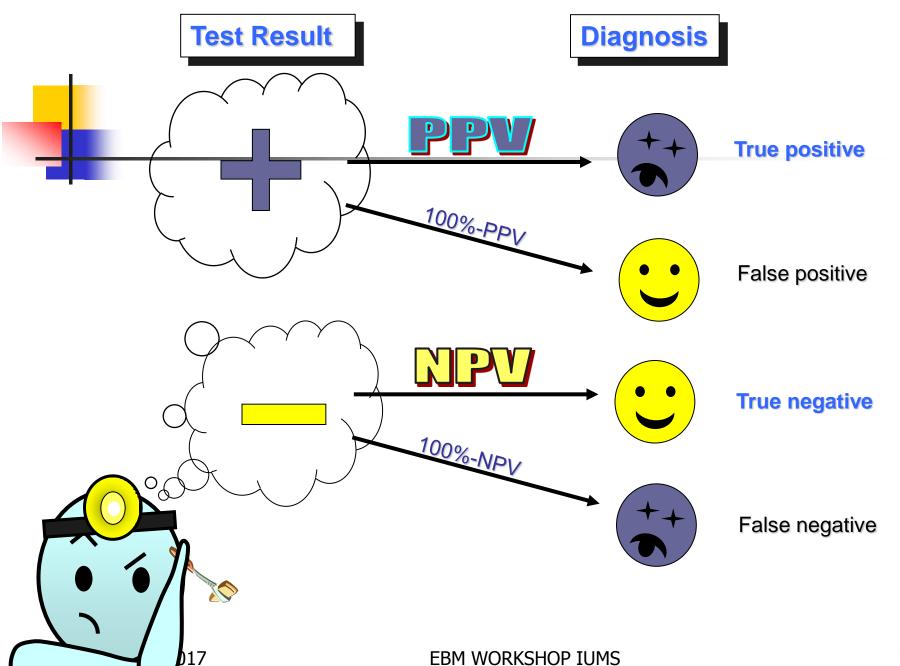
Negative

True positive	False positive
A = 103	B = 16
False negative	True negative
C = 12	D = 211

Sensitivity=103/(103+12)=89%

Specificity=211/(16+211)=93%







Likelihood ratio

Likelihood ratio =the likelihood of a test result in patients with the disease / the likelihood of a test result in patients without the disease

- LR(+) = sensitivity/(1-specificity)
- LR(-) = (1-sensitivity)/specificity



Likelihood Ratio

When ordering a test, which tests will best help us rule in or rule out disease?

- Initial assessment of likelihood of disease = pre-test probability
- Final assessment of likelihood of disease = post-test probability



Likelihood Ratio

Probability of patient with disease having a given test result

Probability of patient without disease having a given test result



Positive Likelihood Ratio (LR+)

Probability of patient with disease having a positive test result

Probability of patient without disease having a positive test result

legative Likelihood Ratio (LR-)

Probability of patient with disease having a negative test result

Probability of patient without disease having a negative test result



Likelihood Ratios

LR+

LR-

sensitivity

1-sensitivity

1 - specificity

specificity

Disease

Present

Absent

Test Result **Positive**

False positive	True positive
В	Α
True negative	False negative
D	С

Negative

Sensitivity = A / (A+C)

Specificity = D / (B+D)

PPV = A / (A+B)

NPV = D / (C+D)

$$LR(+) = \frac{A/(A+C)}{B/(B+D)} = sn/(1-sp)$$

LR(-) =
$$\frac{C/(A+C)}{D/(B+D)}$$
 = (1-sn) / sp

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Disease

Present

Absent

Test Result **Positive**

Negative

True positive	False positive
A = 103	B = 16
False negative	True negative
C = 12	D = 211

Sensitivity=103/(103+12)=89%

Specificity=211/(16+211)=93%

$$LR(+) = \frac{A/(A+C)}{B/(B+D)} = sn/(1-sp)=12.7$$

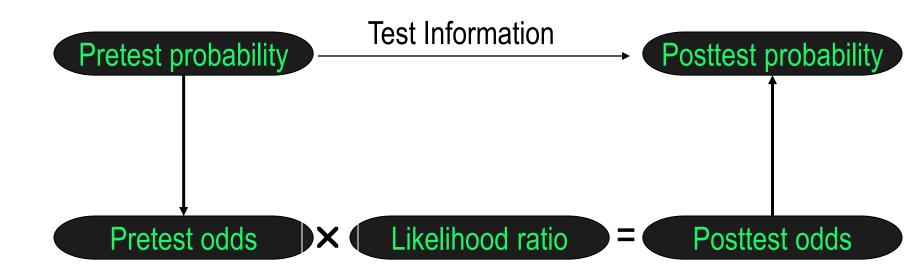
LR(-) =
$$\frac{C/(A+C)}{D/(B+D)}$$
 = (1-sn) / sp=0.11

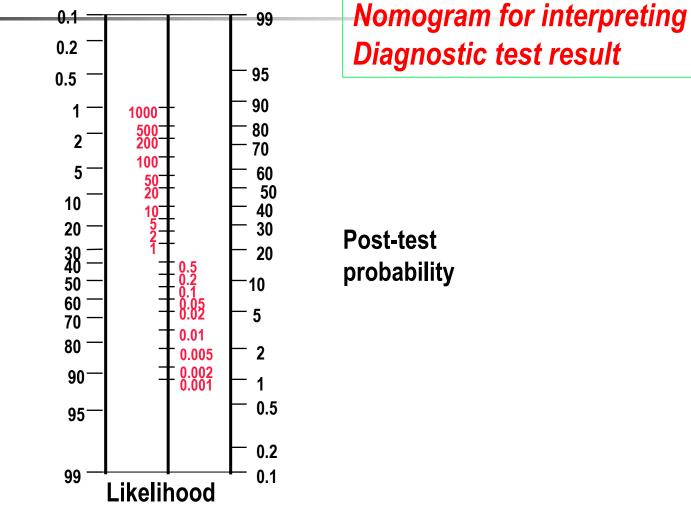
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Likelihood ratio

- LR can be derived for diagnostic tests that have multiple levels or categories of results
- LR from different, independent tests can be used together sequentially to easily calculate a single estimate of a patient's post test probability of disease

Calculating posttest probability





ratio

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Pre-test

probability

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- You are consulted to visit a 62-year-old man with 3 months history of severe back pain. His weight remained stable. CBC and routine biochemistry were normal. ESR was 52 mm / hour. An x-ray of the lumbar and thoracic spine was reported to showing degenerative changes.
- what is your approach to this patient?

Clinical findings predicting cancer as a cause of back pain

LR 2.7 2.7 14.7 3.0 Persistent pai 2.6 Dura 1.6 2.4 19.2 55.5

Finding

- Age > 50 years
- Unexplained weight loss •
- Previous history of cancer
- Persistent pain despite 1 month of treatment
 - Duration of this episode > 1 month
 - Severe pain
 - ESR > 20
 - ESR > 50
 - ESR > 100
 - Hematocrit < 30%
 - Lytic or blastic lesion on spine x-ray

15.2

-120



Given that the probability of malignancy as the cause of persistent back pain in the general population is about 0.3%, what is the effect of patient's ESR on the probability of malignancy in this patient?

Clinical findings predicting cancer as a cause of back pain

LR

2.7

2.7

14.7

3.0

2.6

-1.6

2.4

-19.2

-55.5

-15.2

-120

Finding

Age > 50 years

Unexplained weight loss

Previous history of cancer

Persistent pain despite 1 month of treatment

Duration of this episode > 1 month

Severe pain

ESR > 20

ESR > 50

ESR > 100

Hematocrit < 30% ■

Lytic or blastic lesion on spine x-ray

Calculating posttest probability



Pretest oddsxlikelihood ratio=posttest odds

Consider that x-ray of spine in this patient shows a lytic lesion then what will be the probability of malignancy in this patient considering also patients age and ESR?

Clinical findings predicting cancer as a cause of back pain

Finding

- Age > 50 years
- Unexplained weight loss •
- Previous history of cancer
- Persistent pain despite 1 month of treatment
 - Duration of this episode > 1 month
 - Severe pain
 - ESR > 20
 - ESR > 50
 - ESR > 100
 - Hematocrit < 30% ■
 - Lytic or blastic lesion on spine x-ray



2.7

14.7

3.0

-2.6

1.6

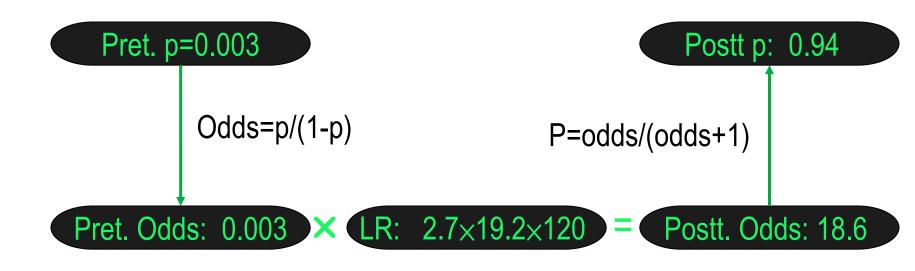
2.4

-19.2

55.5

15.2

-120



Pretest odds \times LR1 \times LR2 \times LR3=posttest odds



Thank You! Any Question?



PROGNOSIS

CONFIDENCE INTERVAL

a range of values that includes the true population value

Expressed with a given degree of expected certainty such as 95%

$$X + /- SE$$

• For example, Frequency of lung cancer =4.1% could have 95% CI of -1.0 to 9.2

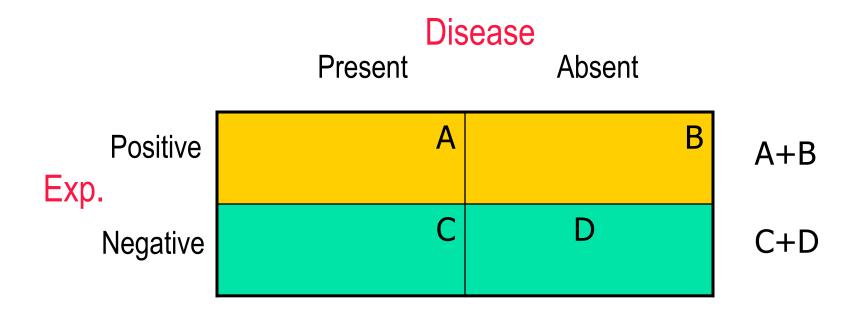


Thank You! Any Question?



THERAPY

- Relative Risk (RR)
- Relative Risk Reduction (RRR)
- Absolute Risk Reduction (ARR)
- Number Needed to Treat (NNT)

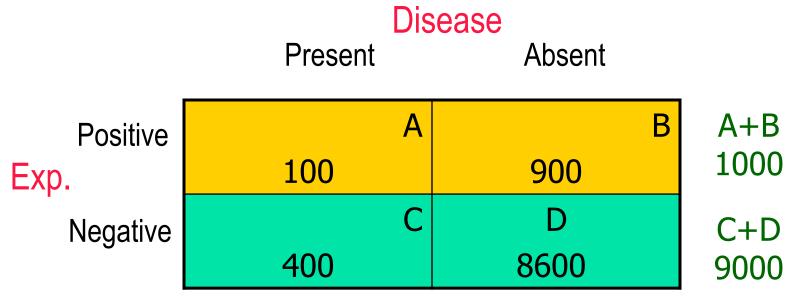


EER = A/(A+B)

CER = C/(C+D)

EER = 100/1000

CER = 400/9000



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Absolute Risk Reduction

ARR = CER-EER

ARR = C/(C+D) -A/(A+B)

ARR= 200/1000 - 600/1000 Disease

Present

Absent

	Positive
Exp	
	Negative

Α	В
600	400
C	D
200	800

A+B 1000

C+D 1000

Relative Risk

 Risk Ratio is the ratio of risk of the outcome event in the experimental (intervention or treated group) to the risk in control group

RR = EER/CER = [A/(A+B) / C/(C+D)]

RR=600/1000 / 200/1000

Disease

Present Absent

Positive Exp.

Negative

Α	В
600	400
С	D
200	800

A+B 1000 C+D

1.0

Relative Risk Reduction

RRR = [(CER-EER)/CER]

RRR = 1-RR

RRR=[(200/1000 - 600/1000)] / 200/1000 X 100

Disease

Present

Absent

	Positive
Exp.	

Negative 3/11/2017

Α	В
600	400
С	D
200	800

A+B

1000

C+D

1.7

Number Needed to Treat

- NNT is particularly useful to clinicians who want to know whether the probable benefits of some treatments or intervention will be worthwhile in their patients
- NNT = 1/ARR
- \blacksquare NNT = 1/0.041 = 24



Thank You! Any Question?



In horse racing terms, 10 horses running you bet on 1 horse

Odds of winning are 1:9 (you Vs. the rest)

Risk of winning is 1:10

(you Vs. all the whole field)

1.9



HARM

OR NNH

Odds ratios

- Cannot use relative risk unless we are looking forward in time (cohort study, RCT)
- For case-control study, can calculate the odds ratio (OR) which tells us the odds of having had a certain exposure in diseased versus not diseased (dead or alive)
- Note, in rare diseases (a situation where you are likely to perform case-control study) OR approximates RR pretty well

Odds ratio = odds of exposure for cases odds of exposure for controls

	Controls	Cases	
1643	984 (b)	659 (a)	Smokers
373	348 (d)	25 (c)	Non- smokers
2016	1332	684	

3/11/2017

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- The odds of lung cancer patient having smoked is the ratio of the number of cases who smoked to those who did not (659/25 = a/c)
- The odds of a controls having smoked is the ratio of the number of controls who smoked to those who did not (984/348 = b/d)

```
Odds ratio = a/c
b/d

= ad/bc (cross product)
= 9.32
```

Interpretation ???

NNH

Rates of adverse events due to treatment (\mathbf{R}) number needed to harm (NNH)

$$NNH = \frac{1}{R}$$

$$NNH = \frac{1}{R_1 - R_2}$$

= the reciprocal of the actual difference in rates of bad adverse events between experimental (R, R1) and control (R2)

 $NNH = \frac{1}{R_1 - R_2}$ = group. = the number of patients who must be treated with the experimental treatment in order for with the experimental treatment in order for one to experience a harmful event

CER =

EER =

RR=

RRR=

ARR=

NNT=

Disease

Present

Absent

Positive

Exp.

Negative

В
1200
О
900

C+D 1000

A+B

1500

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Odds ratio =

	Controls	Cases	
1400	800 (b)	600 (a)	Smokers
450	400 (d)	50(c)	Non- smokers
1850	1200	650	

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Thank You! Any Question?



What is a Decision?

A decision is an irreversible choice among alternatives to allocate valuable resources



Decision Making Strategies

- Group Strategies
 - Brainstorming
 - Delphi Method
 - Nominal Group Technique
- Individual Strategies
 - Implicit favorite model
 - Satisfying ("administrative") model
 - Maximizing ("rational-economic") model
 - Markov model



Decision Analysis

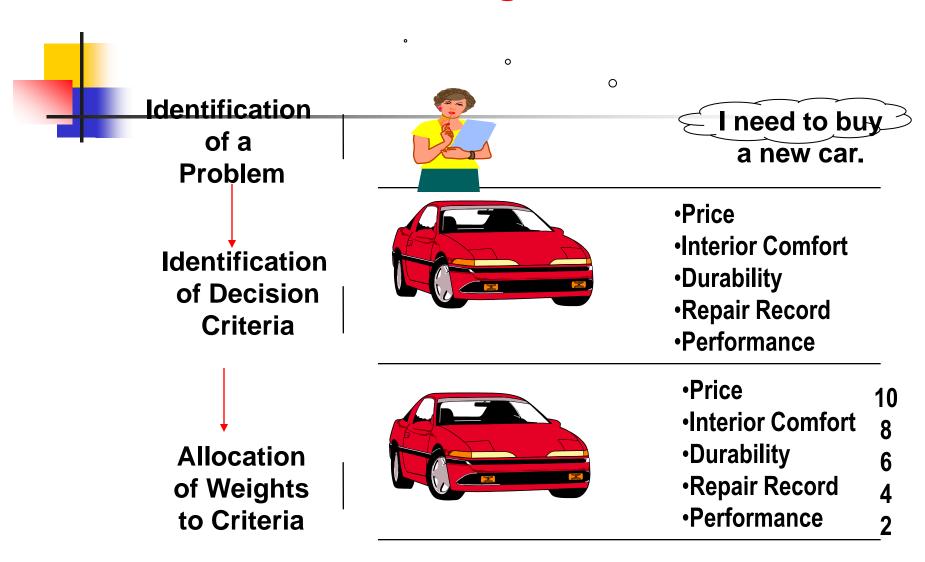
- A systematic, structured approach to decision making when consequences are uncertain.
- Decision analysis is a formalization of the medical decision-making process.

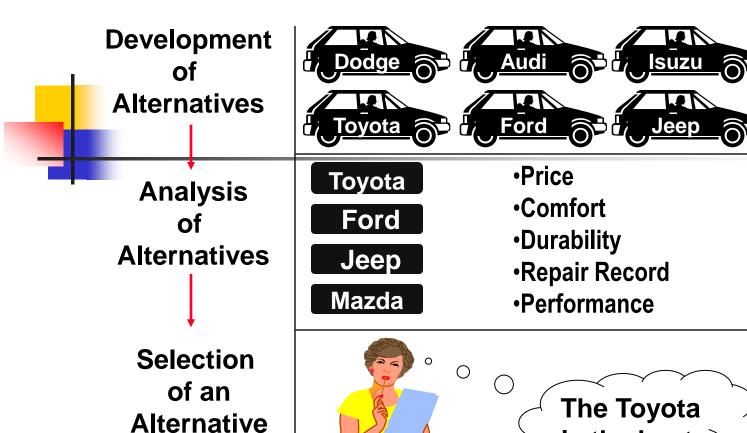


Uses of Decision Analysis

- identify available options when faced with a decision
- predict the consequences or outcomes of each option
- assess the *probability* of occurrence for each outcome
- determine the *value* of each outcome
- select the option that will yield the best "pay-off"

The Decision-Making Process





The Toyota is the best.

Implementation of the Alternative

Appraisal of Decision Results

Dodge

Audi

Isuzu

Chevy

MS



Steps in Decision Analysis

- Formulate an explicit question
- Create a decision tree
- Calculate the expected value of each decision alternative
- Choose the decision alternative with the highest expected value
- Use sensitivity analysis to test the conclusions of the analysis



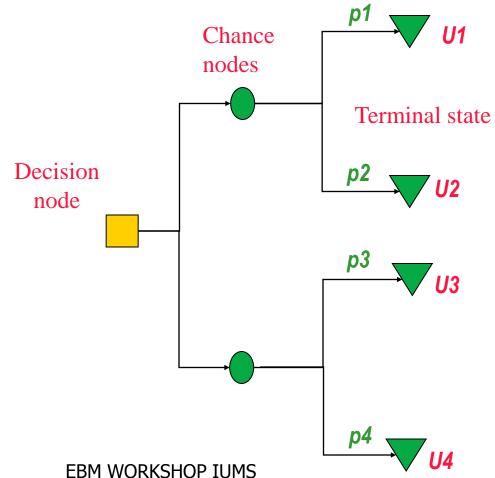
Decision tree

Decision node: represented on the tree as a **square**, is a crossroads in clinical medicine at which the physician must choose an action or strategy.

Chance nodes: which appear as **circles** on the decision tree represent events that are beyond our control; they are the uncertainty in clinical medicine.

Terminal state: which appears as **triangles** on decision tree represents one of the final outcomes





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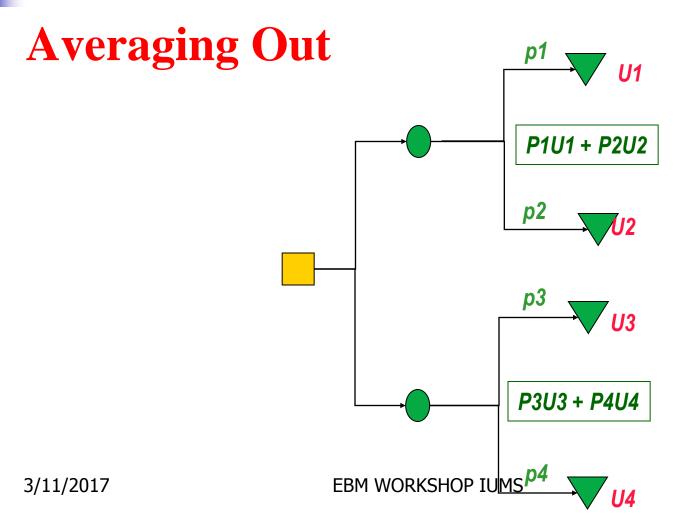


Averaging Out

 Process of calculating an event from several conditional probabilities

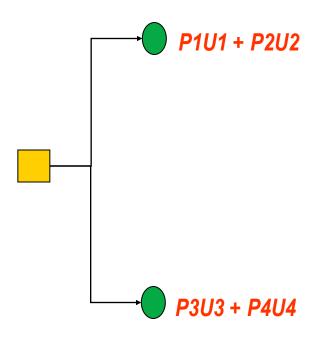
 Multiply the probability of each branch by the value attached to it and sum the values of all branches of the node



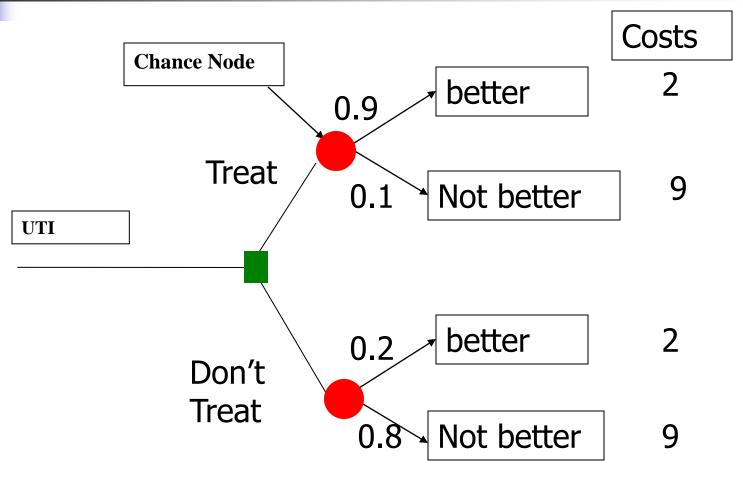




Folding Back



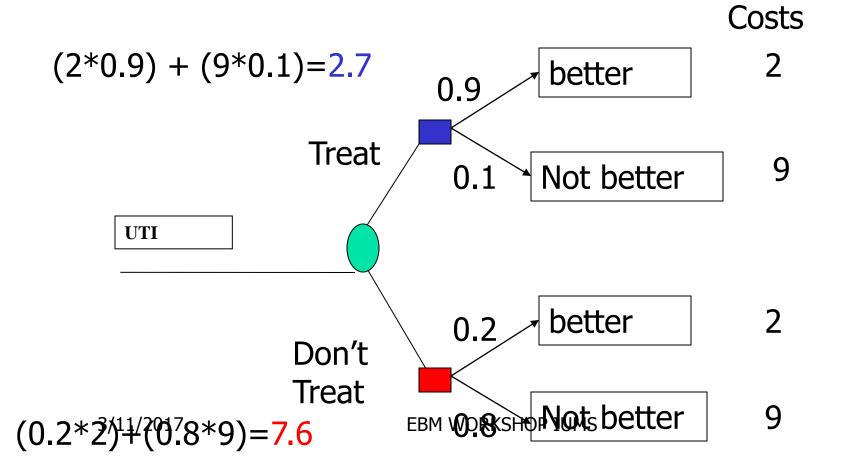




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Rollback Costs



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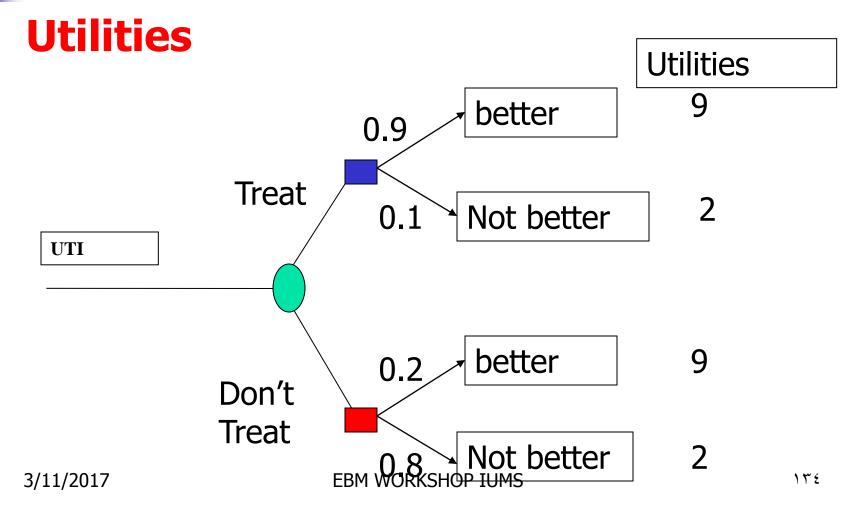
What does the patient think?

■ Utilities (e.g QALYs)

Three common methods for calculating personal utilities

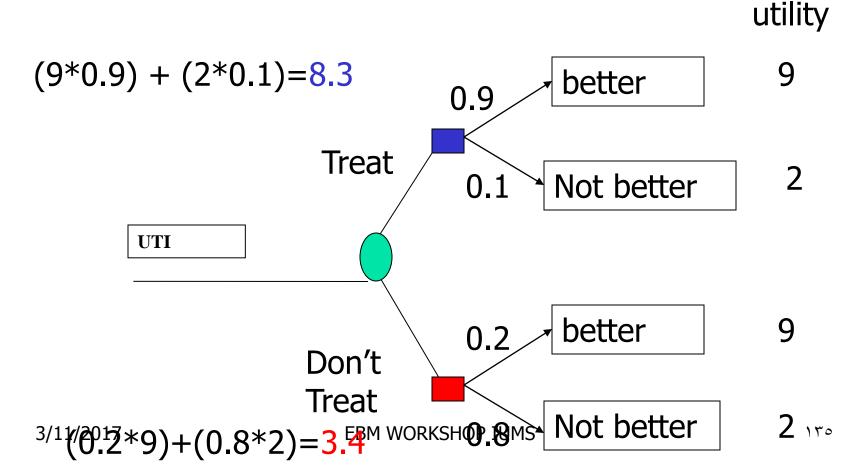
- Visual analog scale
- Time trade-off
- Standard gamble







Rollback





Results

- More people get better (90 % vs 20%)
- It is cheaper (2.70 vs 7.60)
- The utilities are better (8.3 vs 3.4)

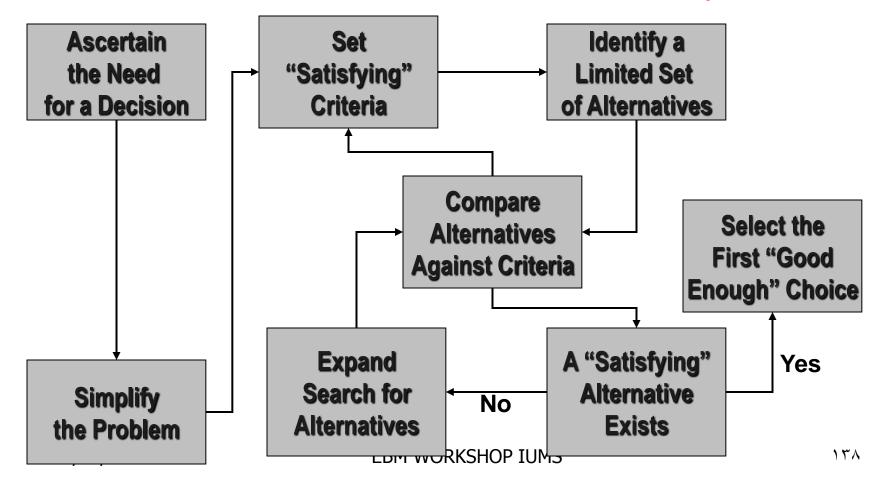
TREATMENT OPTION IS MUCH BETER THAN NO TRETMENT



Individual Decision Making Models

- Implicit favorite
- Satisfied ("administrative")
- Maximizing ("rational-economic")
- Markov

A Model of Bounded Rationality





Sensitivity Analysis

Sensitivity analysis tests the stability of an analysis over a range of probability estimates and value judgments

One-way sensitivity analysis Two-way sensitivity analysis



Thank You! Any Question?