#### Summer Institute in Statistics for Clinical Research

## Exploratory Analyses: Why Do We Need Particular Caution?

July 28, 2016

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- \* Fleming TR "Clinical Trials: Discerning Hype from Substance"
- *Annals of Internal Medicine* 2010; 153:400-406

### Data Driven Hypothesis for the Cancer Risk with Vytorin in Aortic-Valve Stenosis

•	<b>SEAS Trial</b>	<u>N</u>	CA. Incidence	CA. Deaths
	Vytorin	944	101	37
	Placebo	929	65	20
	Relative Risk:		1.55	1.78
	95% C.I.:		(1.13, 2.12)	(1.03.3.11)

### • IMPROVE-IT & SHARP Trials N CA. Incidence CA. Deaths

 Vytorin
 10,391
 313
 97

 Control
 10,298
 326
 72

Relative Risk: **0.96 1.34** 

95% C.I.: (0.82, 1.12) (0.98, 1.84)

#### Interest in "Positive" Results in Clinical Trials

- Industry Sponsors
  - ~ Company profits, ↑ value of stock options, promotion
- Government Sponsors
  - Claims of success in advancing health care
  - ~ Leverage for ↑ in federal funding
- Journal Editors (Publication bias)
- Academic Investigators / Caregivers
  - Increased ability to publish results
     † professional stature, earlier promotion, † salary
  - Desire to offer more therapeutic options to patients
- ....Result: Wide Spread & Significant Conflicts of Interest

#### Bias for "Positive" Results in Clinical Trials

- ~ What is the definition of a successful clinical trial?
- A very common response:
   "A clinical trial that achieves a positive result'
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#### Bias for "Positive" Results in Clinical Trials

- ~ What is the definition of a successful clinical trial?
- A very common response:

  "A clinical trial that achieves a *positive* result"
- > The proper scientific response:
  - "A clinical trial that
    addresses a clinically important issue,
    and that *reliably answers* the questions
    it was designed to address"

- Hyp. Confirmation vs. Hyp. Generation
  - Post-hoc analyses & Random High Bias (new endpoints, new analyses, interim analyses subgroup analyses, covariate adjustments)

- Clinical Endpoints in Pulmonary Arterial Hypertension
  - ~ Overall survival
  - ~ Quality of Life: SF-36 (8 domains), Borg Dyspnea Score
- ~ NYHA Functional Class
  - ~ 6MWT: @18 wk, 24 wk, 48 wk, etc.
  - ~ Time to Clinical Worsening
    - ✓ Death, PAH Hosp, L.T., (NYHA↑ & 6MWT↓)
- Analysis Methods
  - ~ Normally distributed: **T-test**, ANCOVA, Wilcoxon
  - ~ Time to event: Log-rank, Cox Regression
  - ~ Dichotomous: Fisher's Exact Test, Pearson  $\chi^2$

- Biomarker Endpoints (Hemodynamic parameters)
  - Pulmonary Arterial Pressure
  - ~ Systolic & Diastolic Systemic Arterial Pressure
  - ~ Systemic & Pulmonary Vascular Resistance
  - ~ Heart Rate & Cardiac Output
- Analyses over Calendar Time
- Normally distributed: T-test, ANOVA, Wilcoxon
- ~ Time to event: Log-rank, Cox Regression
  - ~ Dichotomous: Fisher's Exact Test, Pearson χ2

- Subgroup Analysis & Prognostic Covariate Adjustment
  - ~ WHO PAH Functional Class: I v II v III v IV
  - ~ Etiology: Idiopathic PAH, Assoc w CTD, SLE, Other
  - ∼ Baseline Walking Distance: < 325 v > **325 meters**
  - ~ Gender: male v female Epoprostenol +/— Sildenafil
  - ~ Age: By decade
  - ~ Ethnicity: White v Black v Asian v Other
  - $\sim$  mean PAP: <50 v >50

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Illustrations and Motivation:

- Hyp. Confirmation vs. Hyp. Generation
  - Post-hoc analyses & Random High Bias (new endpoints, new analyses, interim analyses subgroup analyses, covariate adjustments)

#### Illustrations and Motivation:

Maternity Wards, Baseball & Clinical Research

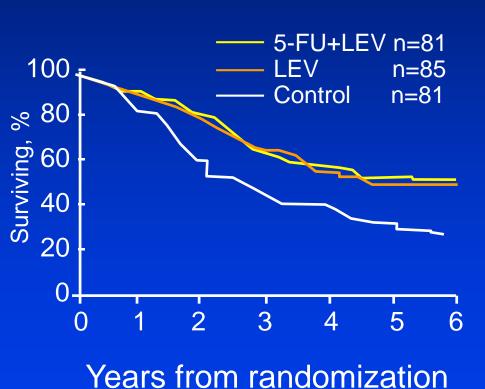
20 vs 2: (.71, .99), 2p = 0.0001

# An Illustration of Exploratory Analyses: Post-hoc Subgroup Analyses

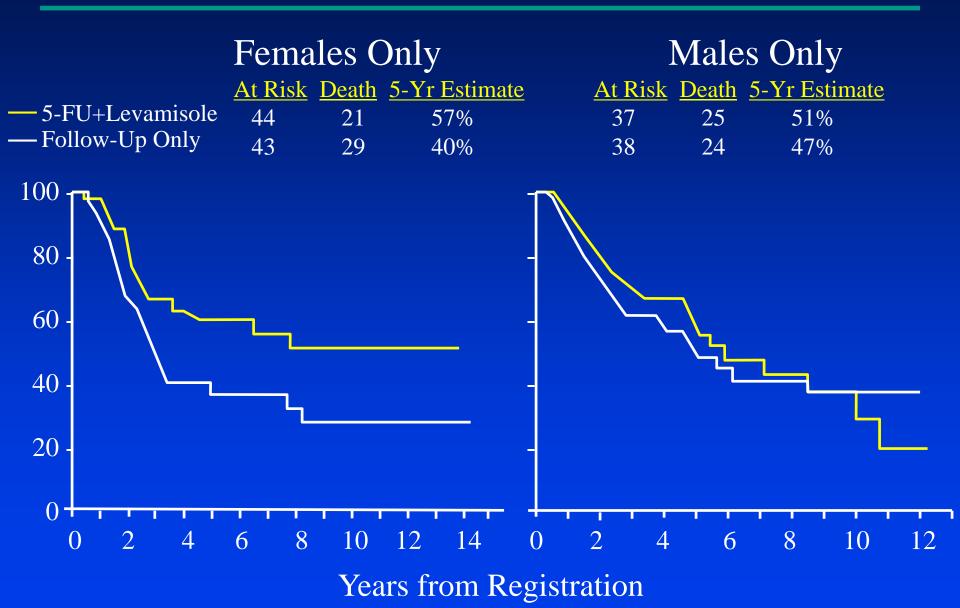
Surgical Adjuvant Therapy of Colorectal Cancer

### Surgical Adjuvant Therapy: Colorectal Cancer

#### **NCCTG** Trial

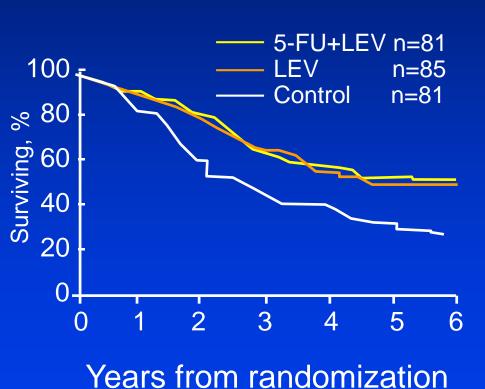


### NORTH CENTRAL TREATMENT GROUP STUDY Looking at Treatment Effect on Overall Survival

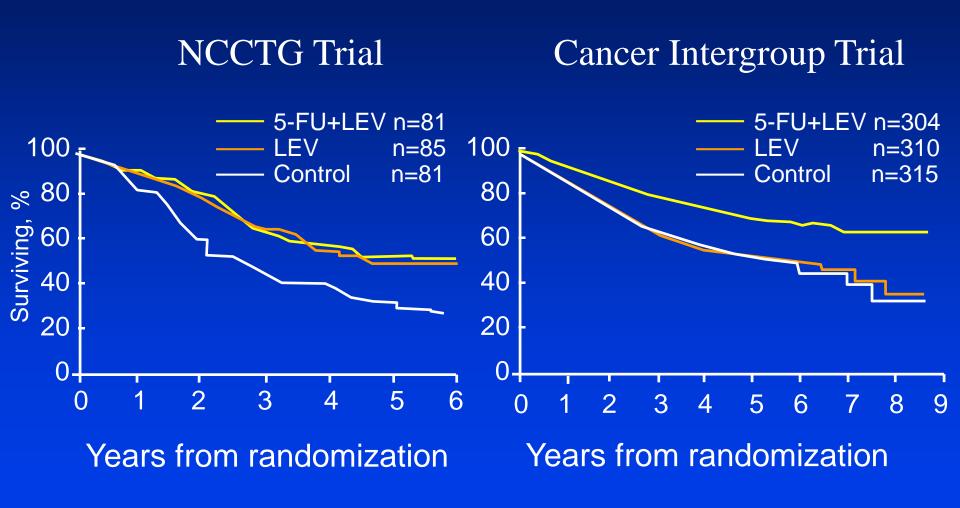


### Surgical Adjuvant Therapy: Colorectal Cancer

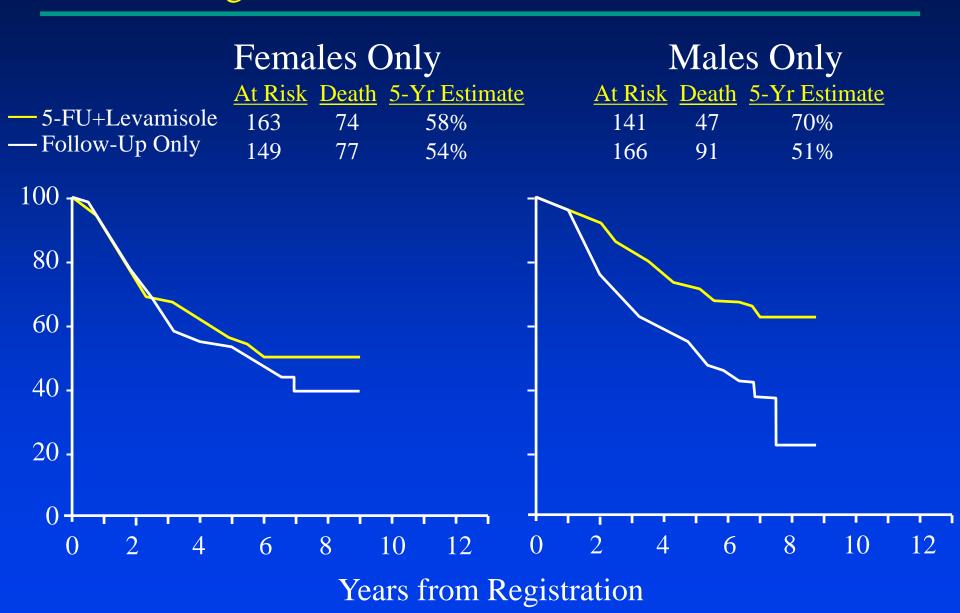
#### **NCCTG** Trial



### Surgical Adjuvant Therapy: Colorectal Cancer



### INTERGROUP STUDY 0035 Looking at Treatment Effect on Overall Survival



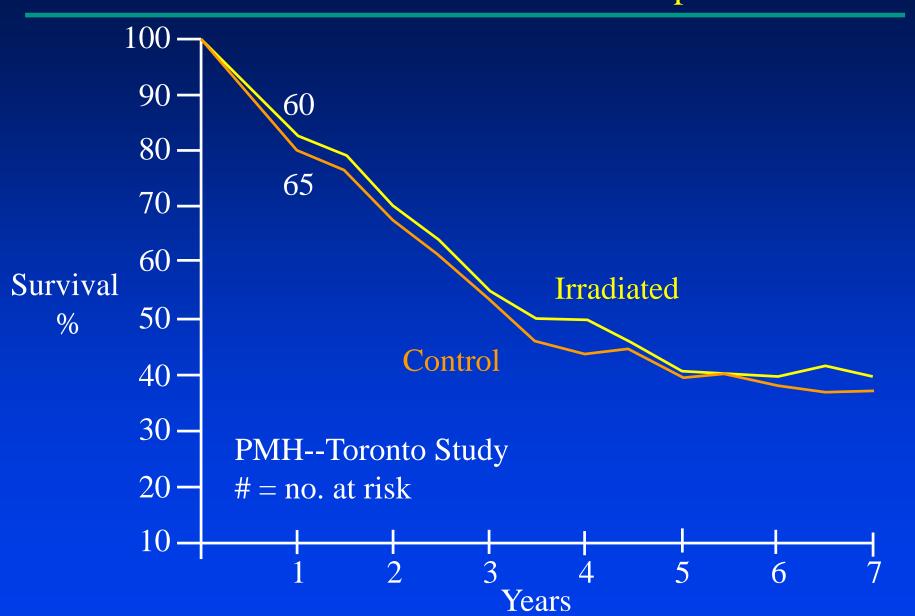
### Duke's C Colon Cancer Adjuvant

Percent \ in \	Death Rate:	5-FU + Levamisole Control
Analysis Group	North Cent Treatmen Group Stud (n = 162)	t Study 1
All patients	28%	33%
Female Male	43% 9%	15% 50%
Young Old	40% 13%	23% 41%

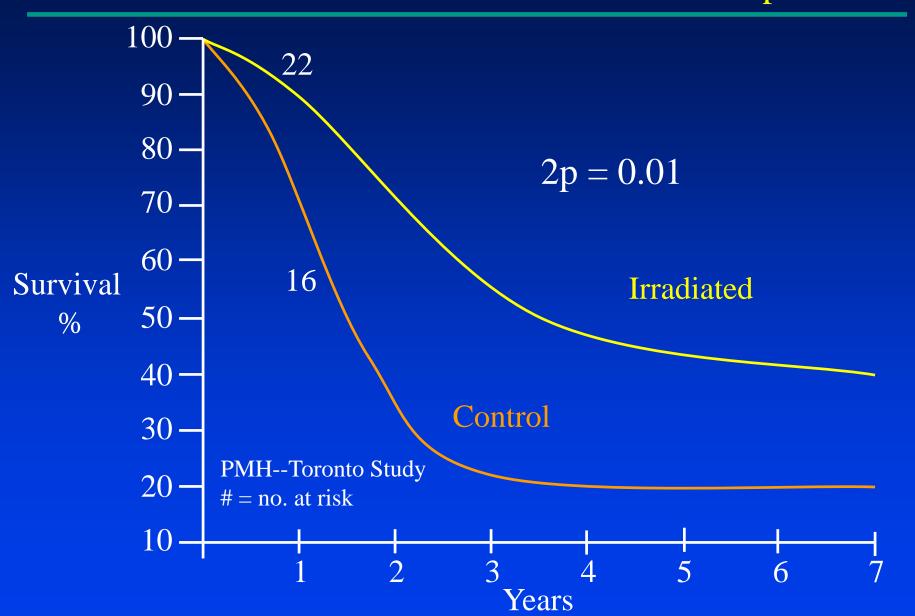
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Radiation Treatment in Rectal Cancer Princess Margaret Hospital

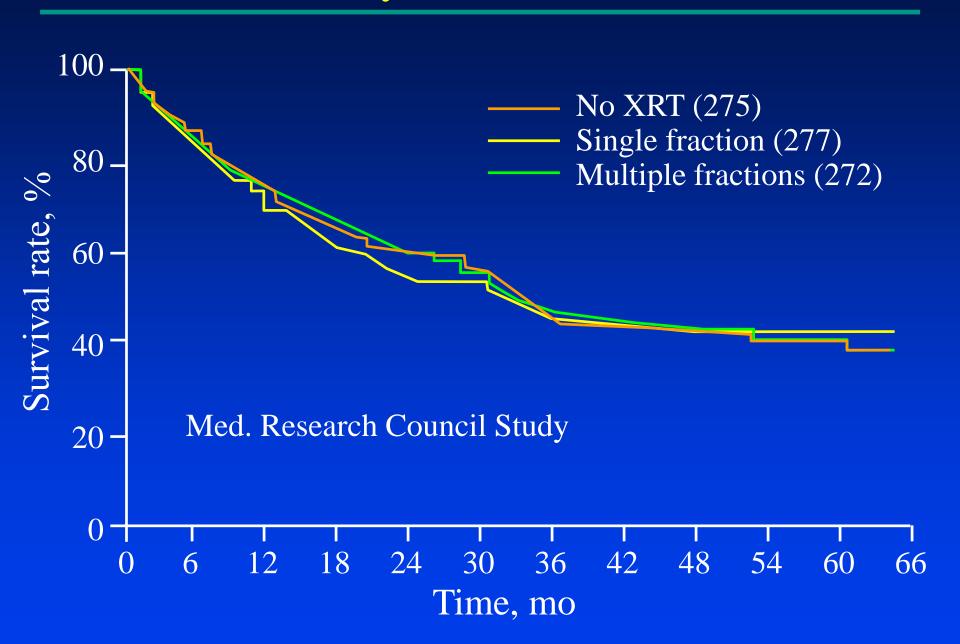
### Survival of Patients with Rectal Carcinoma in Control and Irradiated Groups



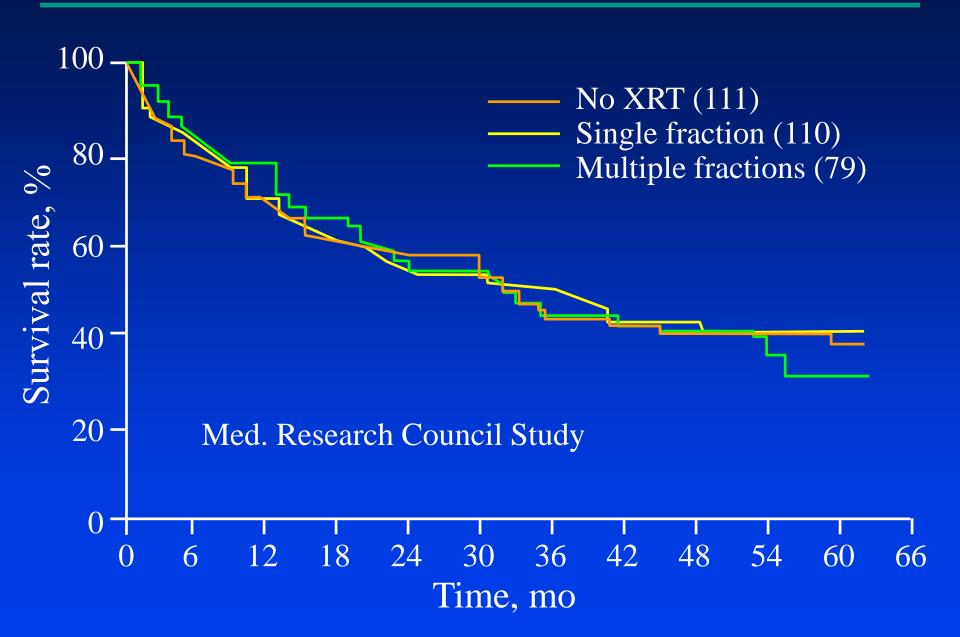
### Survival of Patients with Dukes' Stage C Rectal Carcinoma in Control and Irradiated Groups



### Survival by Treatment Allocated



### Survival by Treatment for Dukes' C Cases

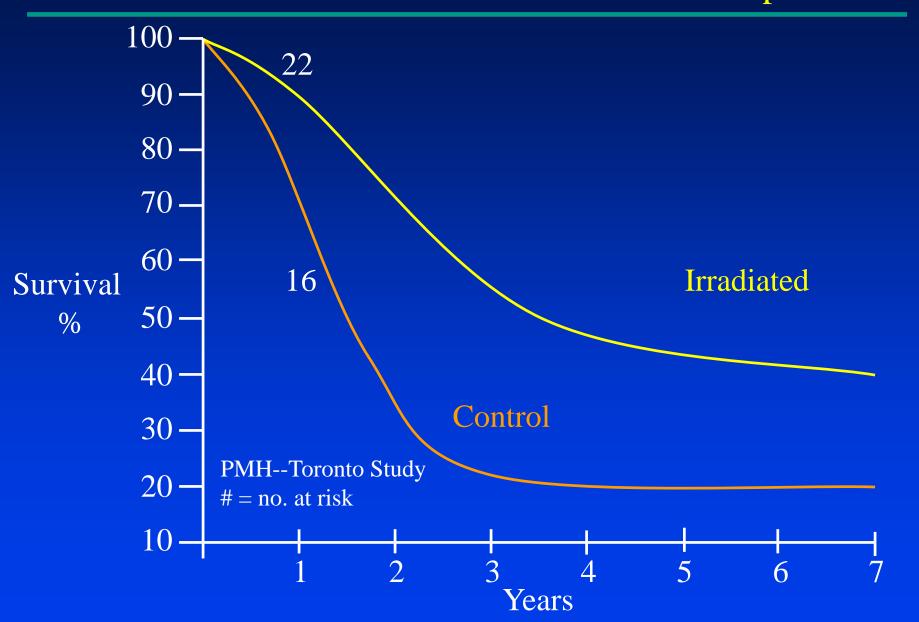


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  - Post-hoc analyses & Random High Bias (new endpoints, new analyses, interim analyses subgroup analyses, covariate adjustments)

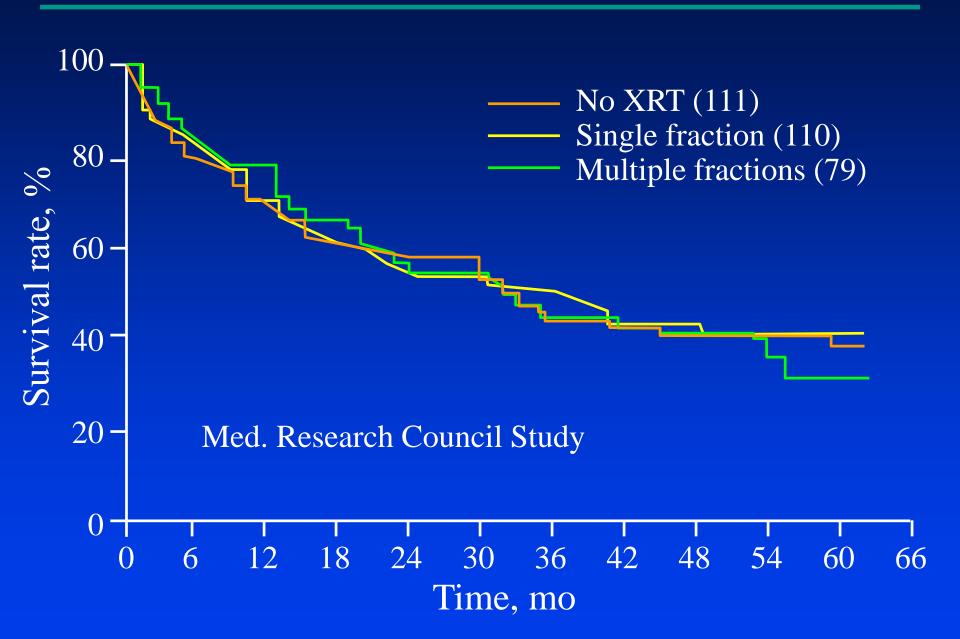
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#### Survival by Treatment for Dukes' C Cases



- GISSI (Lancet '86)
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    - < **65** years
    - < 6 hours from symptom onset

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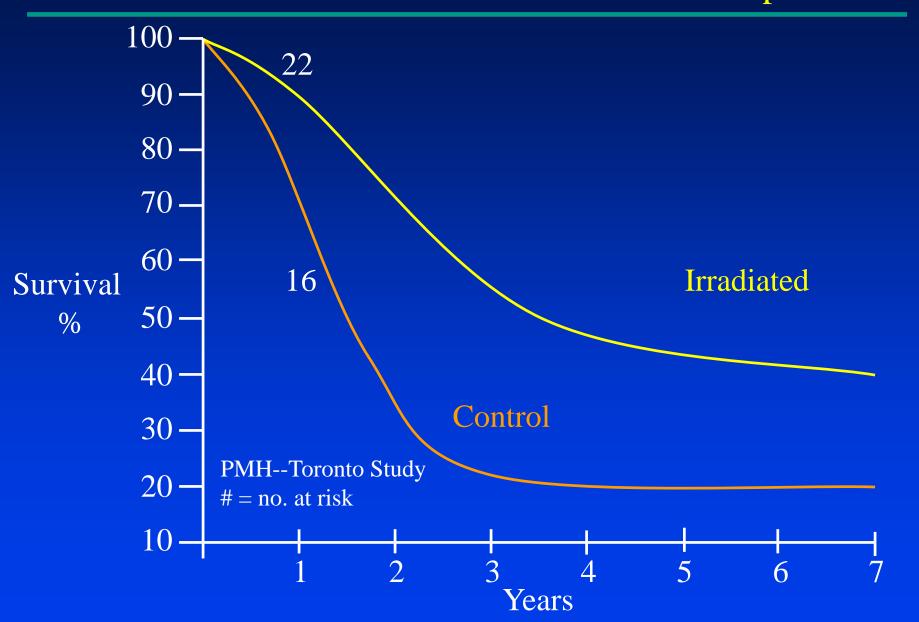
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  - While in ISIS-2:
    - Aspirin beneficial overall...
    - ... yet harmful to patients with
      - astrological signs Libra and Gemini

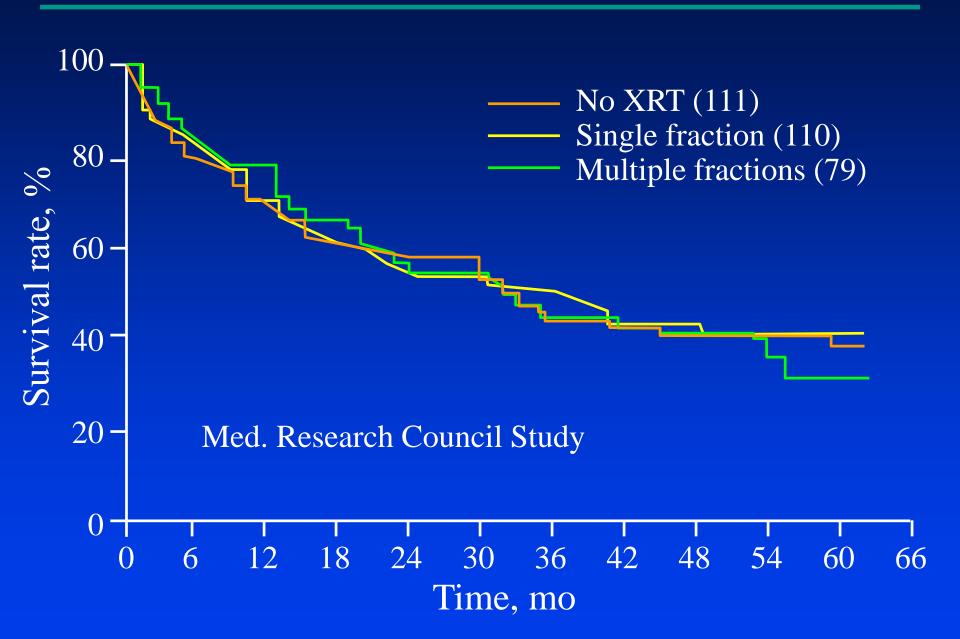
## Can Efficacy or Safety Signals Discovered in Exploratory Analyses Be Viewed to be Reliable Results?

- Criteria to be simultaneously satisfied:
- ✓ < P-values (e.g., Natalizumab & PML & Carvedilol in Heart Failure)
- ✓ Biologically plausible effect
  - White Paper Illustration
- Confirmed by external results

### Survival of Patients with Dukes' Stage C Rectal Carcinoma in Control and Irradiated Groups

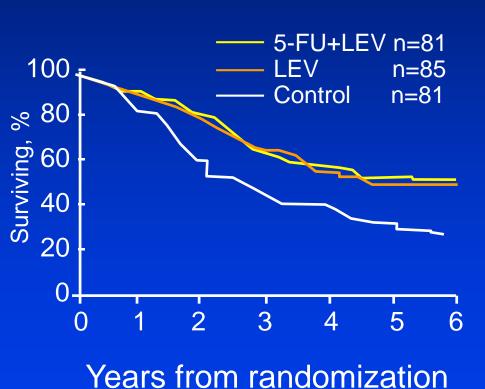


#### Survival by Treatment for Dukes' C Cases



### Surgical Adjuvant Therapy: Colorectal Cancer

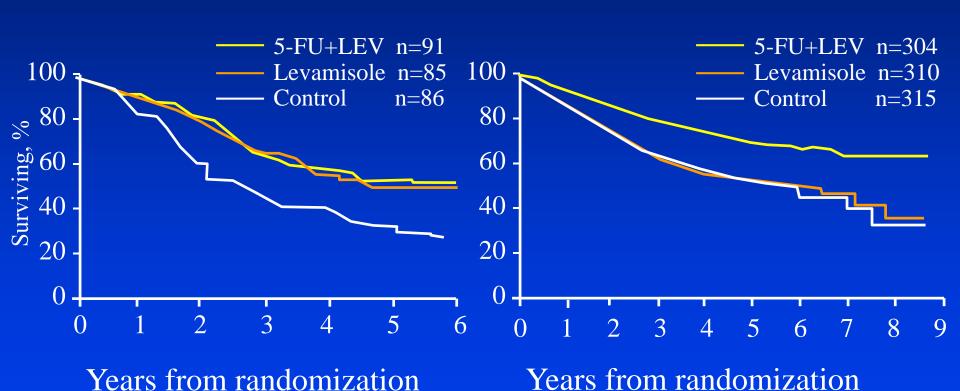
#### **NCCTG** Trial



### Surgical Adjuvant Therapy Of Colorectal Cancer



#### Cancer Intergroup Trial



Of all experimental interventions studied in colon adjuvant, suppose only 4% are truly positive & 96% are truly negative.

Suppose the "false negative error rate" is  $\beta = 0.10$  (so the "statistical power" is  $1-\beta = 0.90$ ) & Suppose the "false positive error rate" is  $\alpha = 0.025$ 

Then, the probability a trial positive will be a true positive is 36/60 = 0.60

RESULT OF EXPERIMENT	TRUTH Positive Negative		
Positive Negative	36 4	24 936	60 940
	40	960	1000

Of all experimental interventions studied, suppose 60% are truly positive & 40% are truly negative

Suppose the "false negative error rate" is  $\beta = 0.10$  (so the "statistical power" is  $1-\beta = 0.90$ ) & Suppose the "false positive error rate" is  $\alpha = 0.025$ 

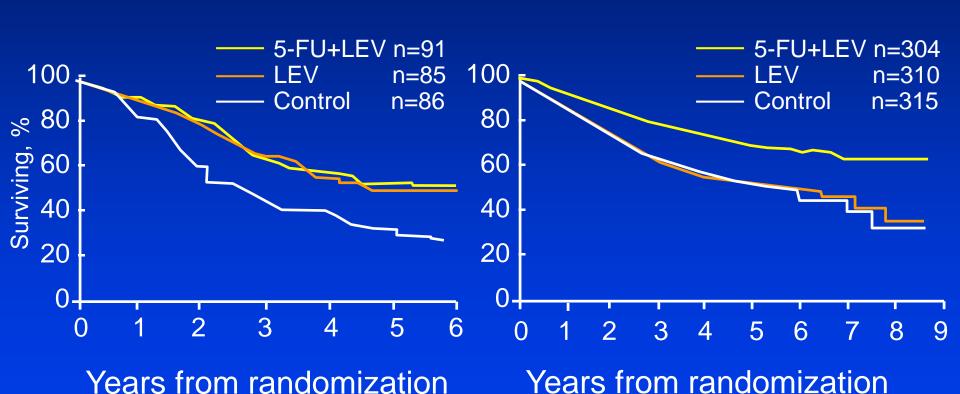
Then, the probability a trial positive will be a true positive is 540/550 = 0.98

RESULT OF EXPERIMENT	TRUTH Positive Negative		
Positive Negative	540 60	10 390	550 450
	600	400	1000

# Surgical Adjuvant Therapy Of Colorectal Cancer



#### Cancer Intergroup Trial



"It isn't so much the things we don't know that get us in trouble.

It's the things we know that aren't so".

—Artemus Ward (1834-1867)

#### **Some Conclusions**

- P-values are only interpretable when you understand the sampling context from which they were derived
- Random High bias is real
- Exploratory Analyses usually should be viewed to be "Hypothesis Generating"
- Confirmatory Trials
   greatly enhance the reliability of conclusions

# Confirmatory vs. Exploratory Analyses

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  - Post-hoc analyses & Random High Bias (new endpoints, new analyses, interim analyses subgroup analyses, covariate adjustments)

#### Illustrations and Motivation:

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20 vs 2: (.71, .99), 2p = 0.0001

Meta-Analysis: 31 vs 13: (.55, .83), 2p = 0.0096

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- Protocol Specified Primary Objective of the Clinical trial:
- Very frequent wording:
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- Protocol Specified Primary Objective of the Clinical trial:
- Very frequent wording:
  - ~ "To *establish* that the experimental regimen is safe and effective"
- Scientifically unbiased wording:
  - ~ "To determine whether the experimental regimen is safe and effective"
    - ...building a story with supportive analyses...

### Bias for "Positive" Results in Clinical Trials

...Andrew Fleming's insight from Psychology...

"Cognitive Dissonance"

...The Harvard Professor's Course...

...The Apparent Lack of Benefit in Males...

- Abetimus Sodium: Reducing Renal Flare Rate in Lupus
- Trial #1: Time to renal flare: Minimal effect, (2p = 0.51)

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- Trial #2 conducted in high affinity subgroup: Time to renal flare:

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- <u>Trial #3</u> conducted in high affinity subgroup with prespecified truncation at 12 months follow-up:

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- Trial #2 conducted in high affinity subgroup:
  Time to renal flare: Minimal non-significant effect
  ...exploratory truncation at 12 months is favorable
- Trial #3 conducted in high affinity subgroup with prespecified truncation at 12 months follow-up: ...early termination by DMC for futility.

# "If you Torture Data Long Enough, They will Confess"

\* Fleming TR "Clinical Trials: Discerning Hype from Substance" Annals of Internal Medicine 2010; 153:400-406

# Principles & Insights

"The Goal of Clinical Research:

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