

# Diabetes and Thrombosis

Enrico Cagliero, MD  
Associate Professor of Medicine,  
Harvard Medical School  
Diabetes Center,  
Massachusetts General Hospital

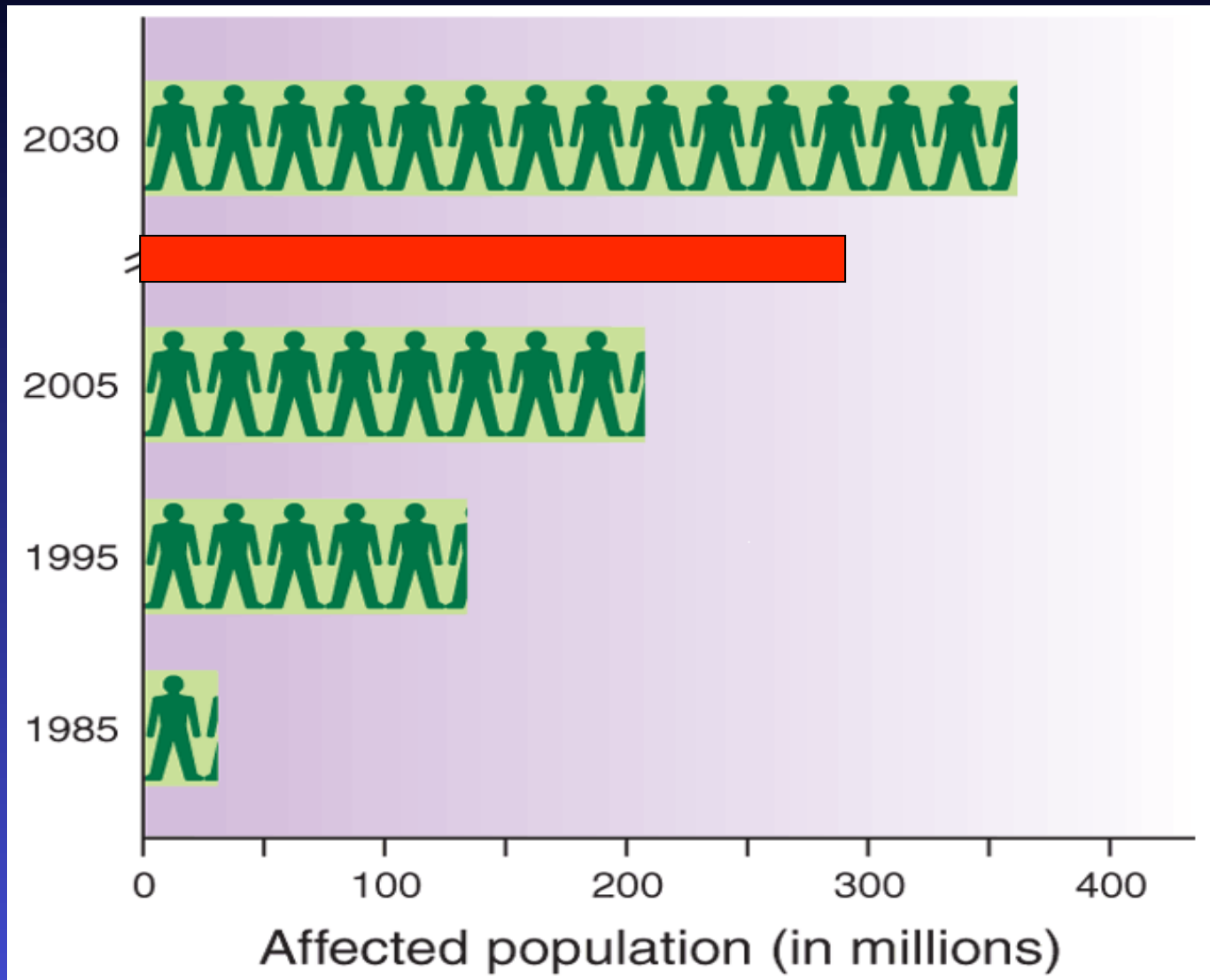


# Overview

- Background and epidemiology
- Blood glucose and cardiovascular disease
- Diabetes as a prothrombotic condition
- Role of glycemic control in CVD prevention
- Can we prevent CVD in diabetic patients?

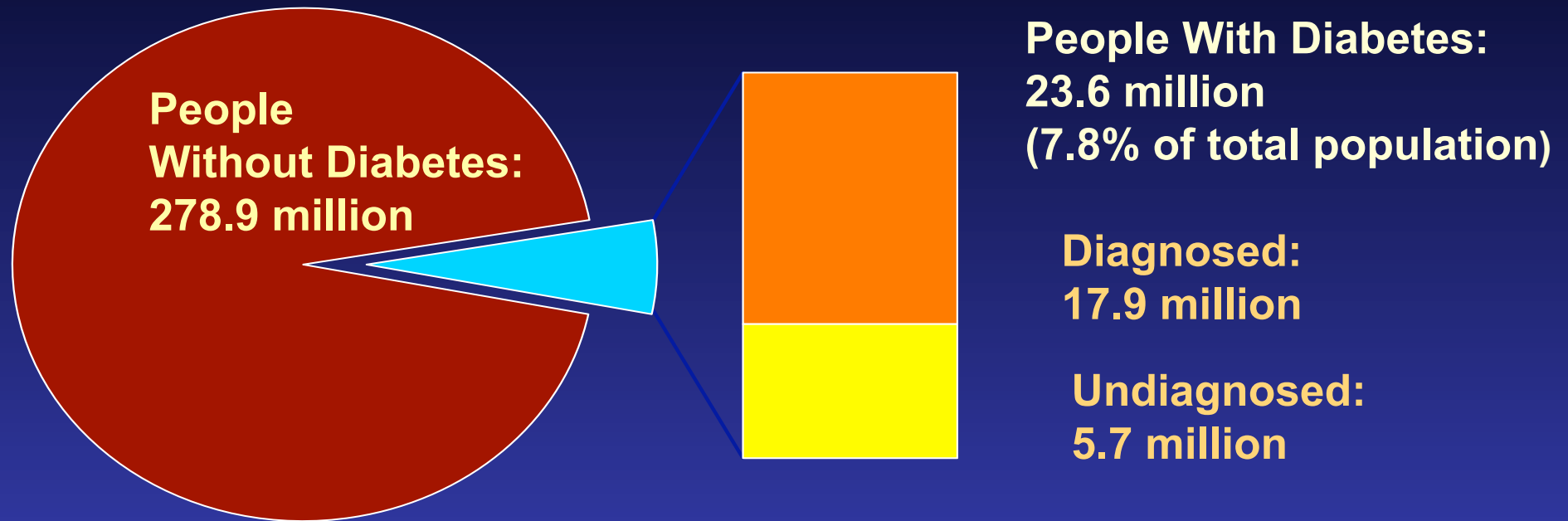
# Background and epidemiology

# Worldwide Diabetic Population



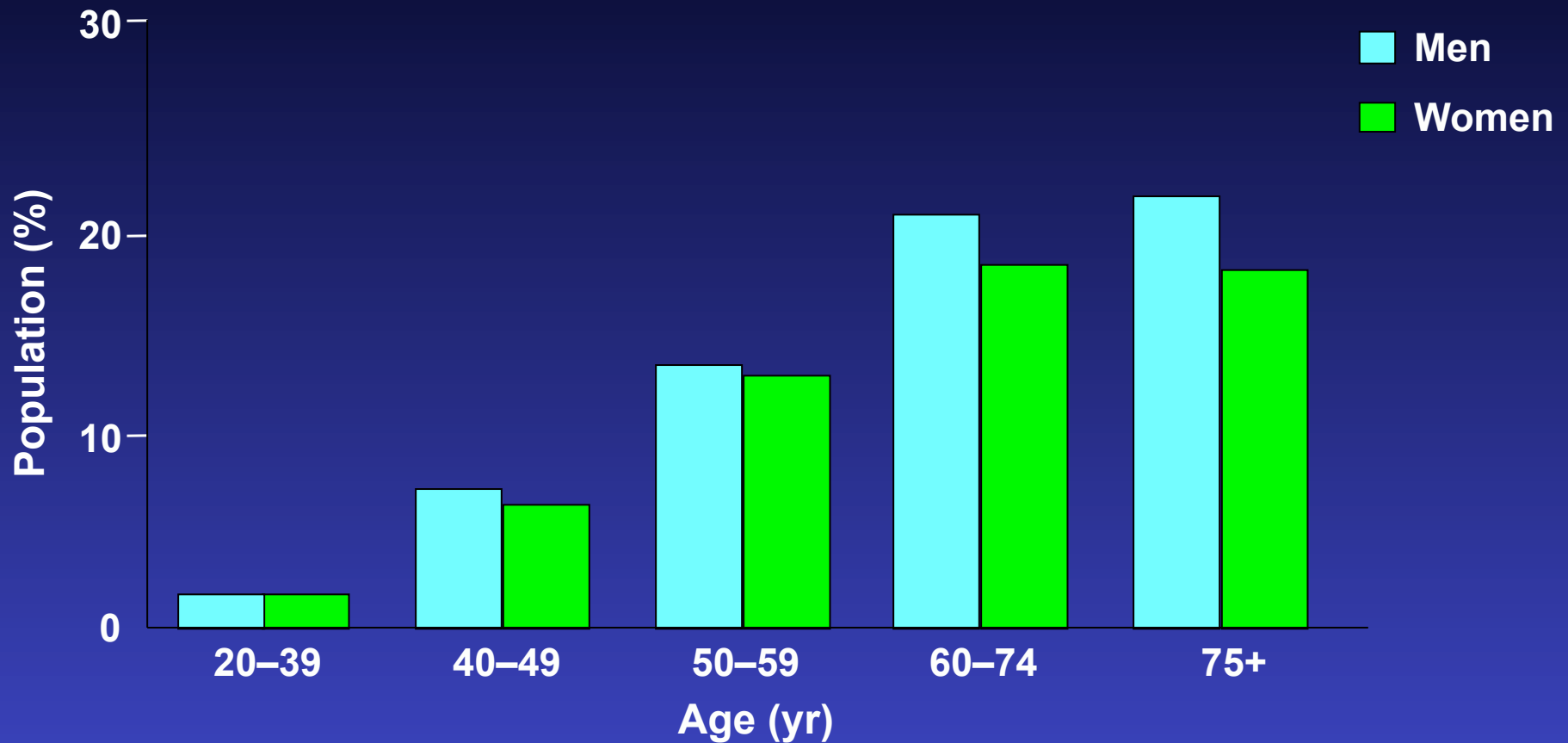
*Smyth and Heron: Nature Medicine 12:75-80, 2005*

# Overview of Diabetes in the United States



- Vast majority (90%-95%) of cases are type 2 diabetes
- One out of 3 children born in 2000 will develop diabetes during its life
- One out of 2 minority children born in 2000 will develop diabetes during its life

# Estimated Prevalence of Diabetes in US: Adult Men and Women



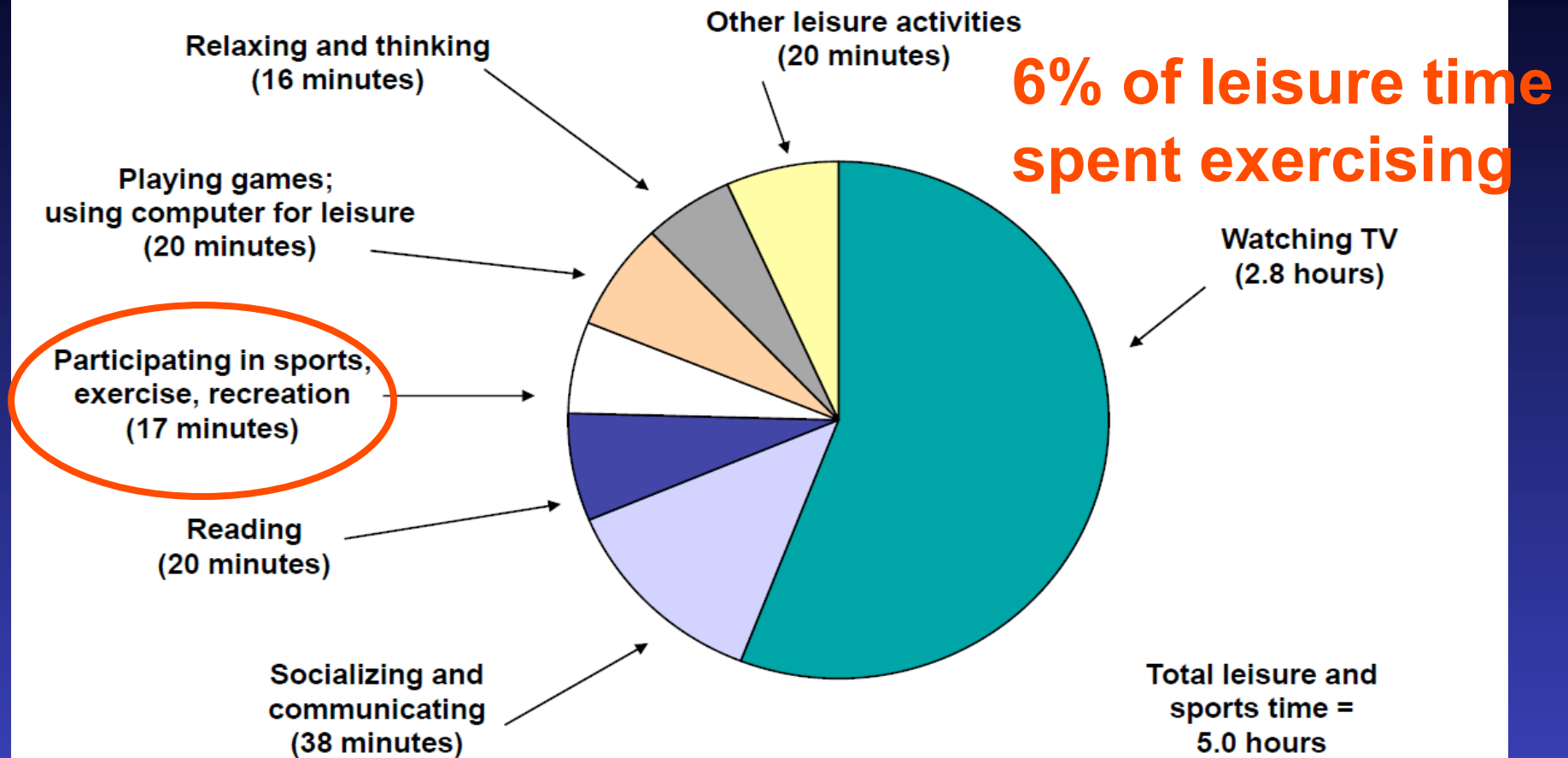


68% of US adults are overweight or obese



- 40% of US adults report no physical activity at all
- 31% of US adults reports at least 20' of exercise 5 days/week
- 5% of adults, obtain 30' of moderate activity 5 days/week

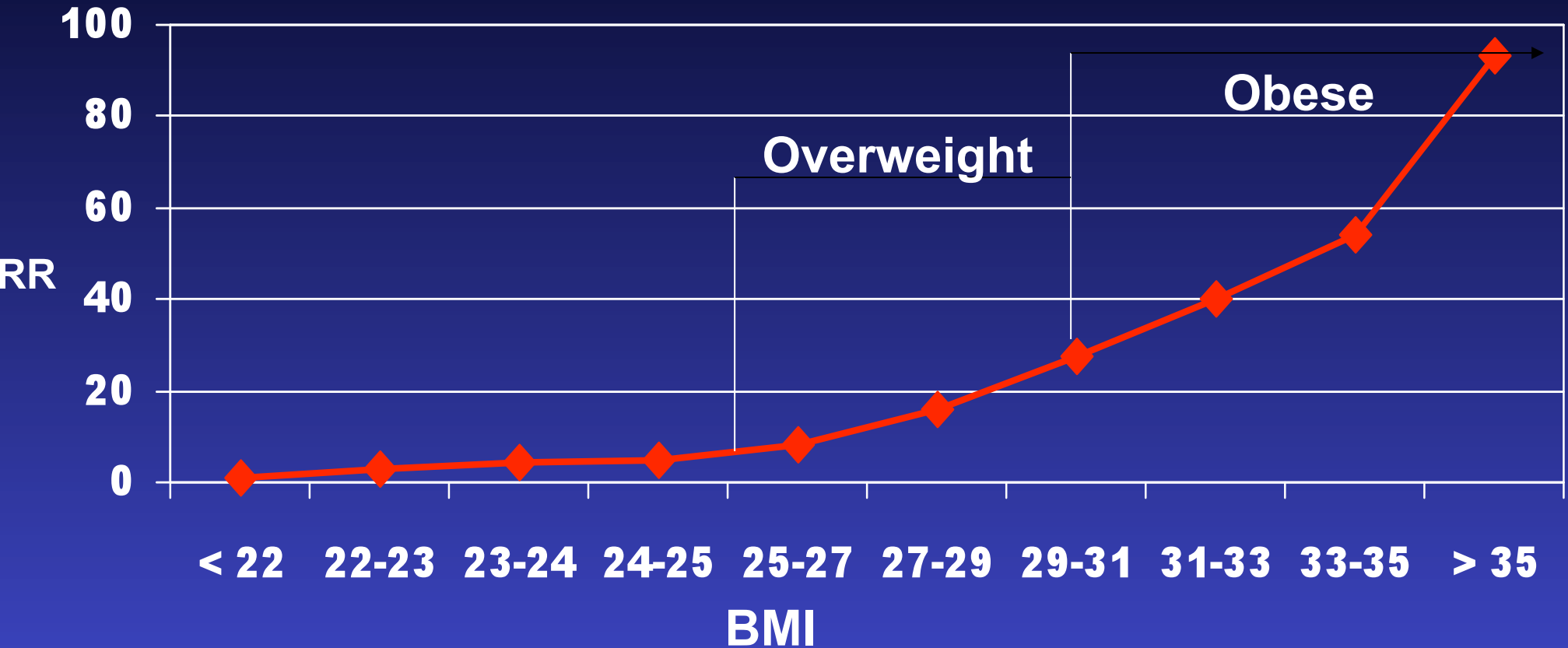
# Leisure time on an average day



NOTE: Data include all persons age 15 and over. Data include all days of the week and are annual averages for 2008.

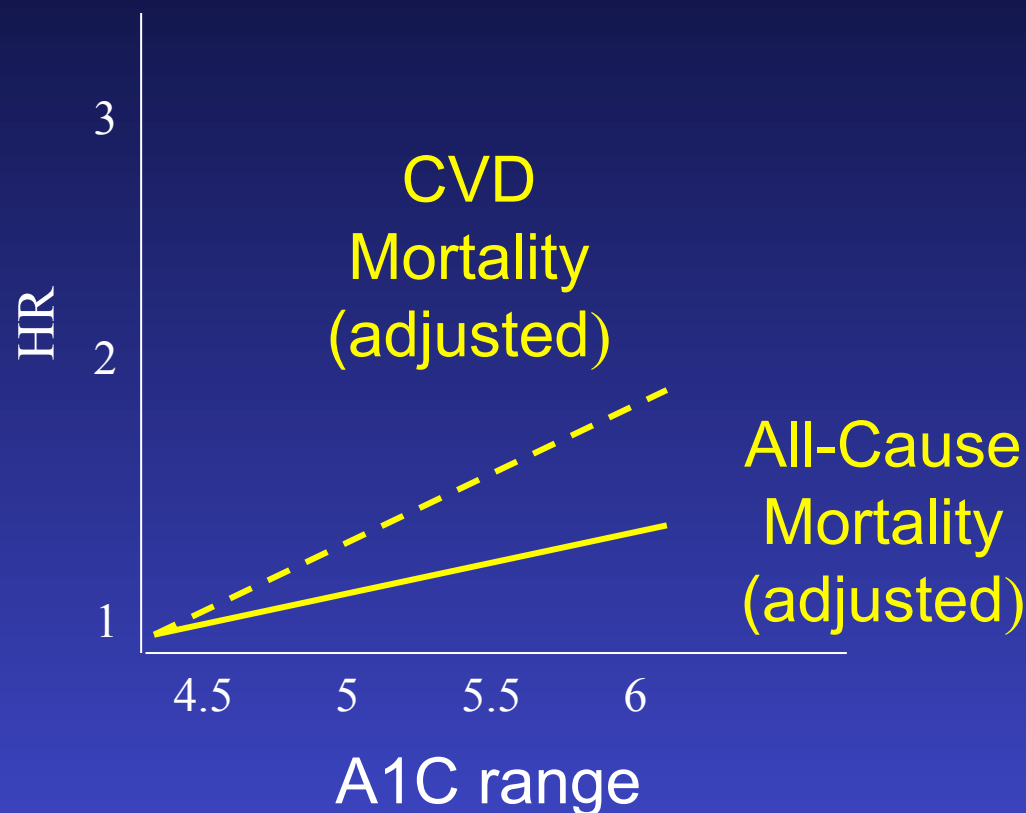
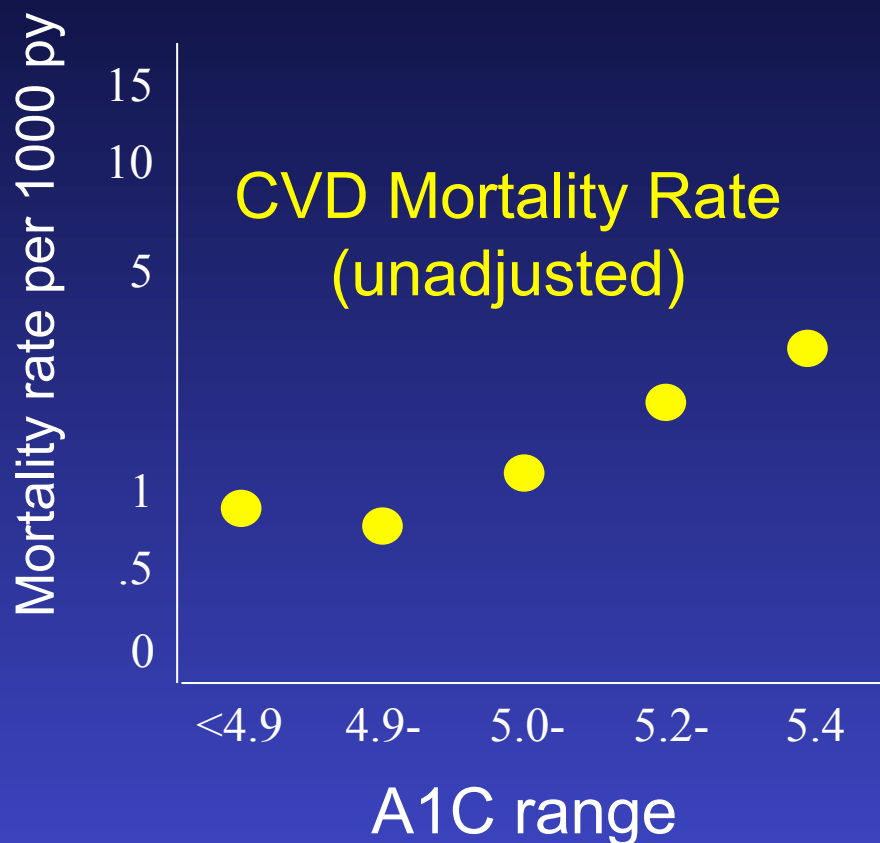
SOURCE: Bureau of Labor Statistics

# Risk of Development of Type 2 Diabetes Nurses Health Study

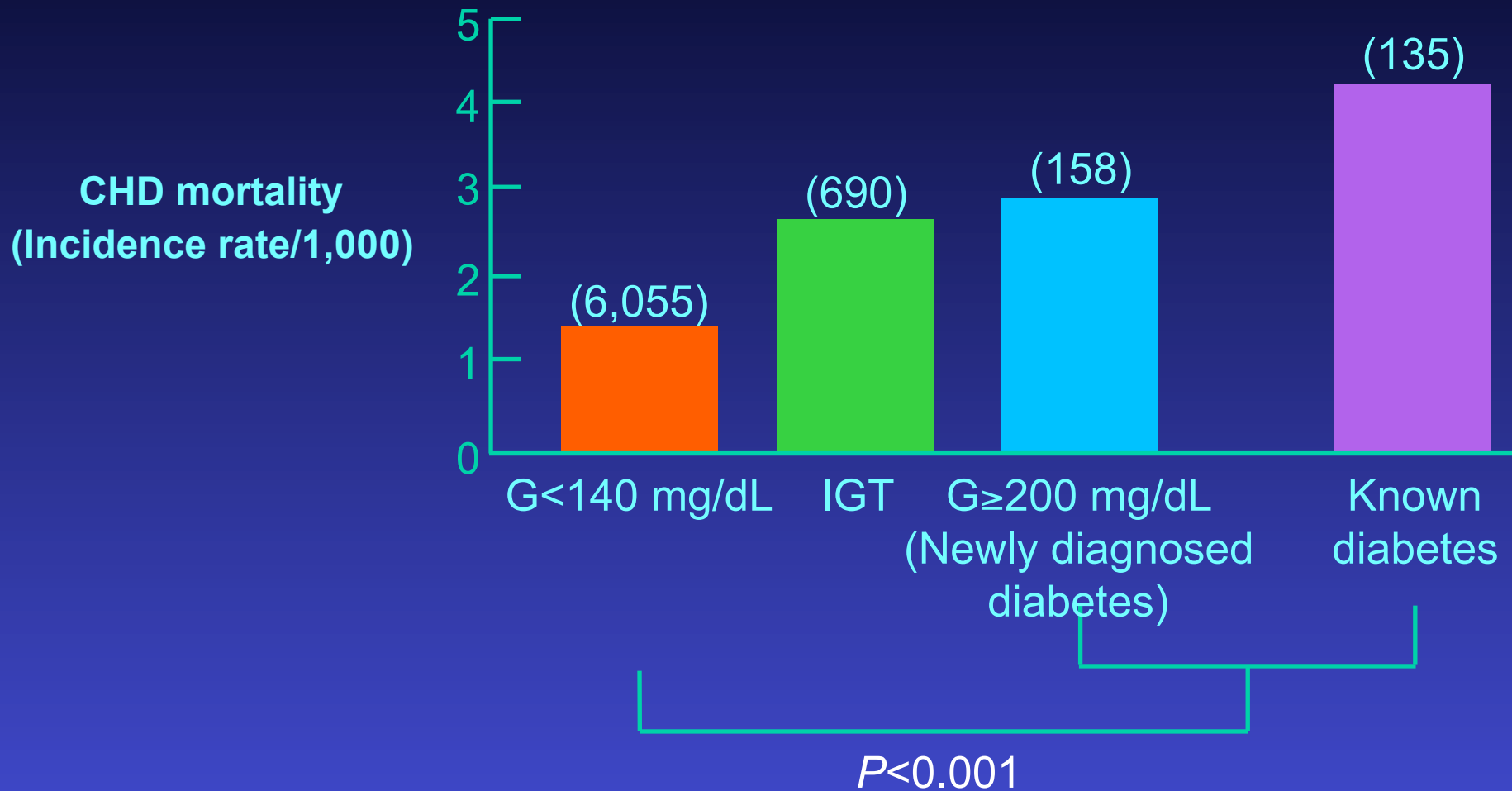


# Blood Glucose and Cardiovascular Disease

# CVD Risk Increases As Glucose Levels (in Normal Range) Increase



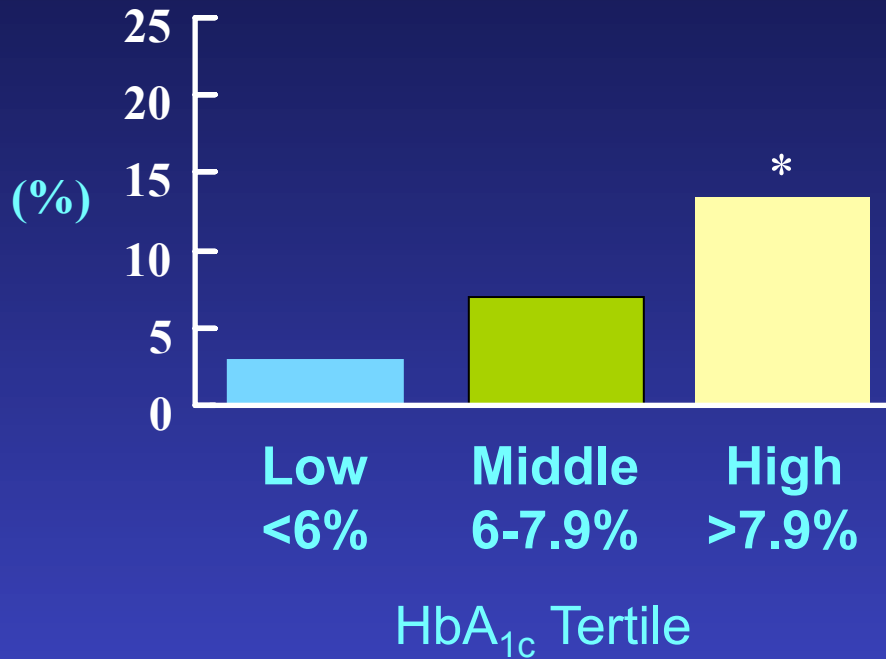
# IGT Progressively Increases Risk of CHD Mortality: Paris Prospective Study



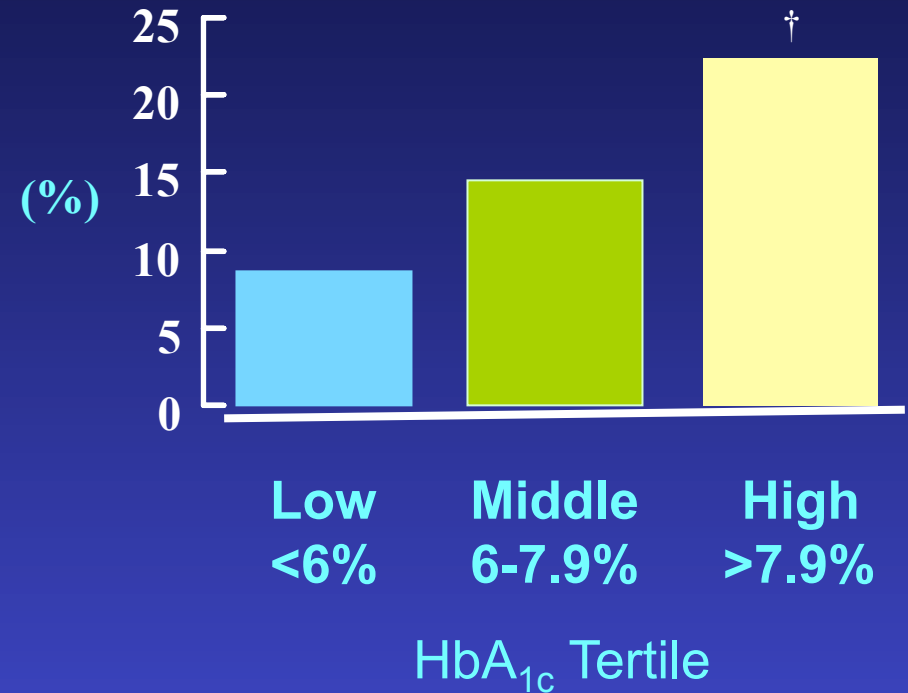
Eschwège E et al. *Horm Metab Res.* 1985;15(suppl):41-46.

# HbA<sub>1c</sub> Predicts CHD in Type 2 Diabetes

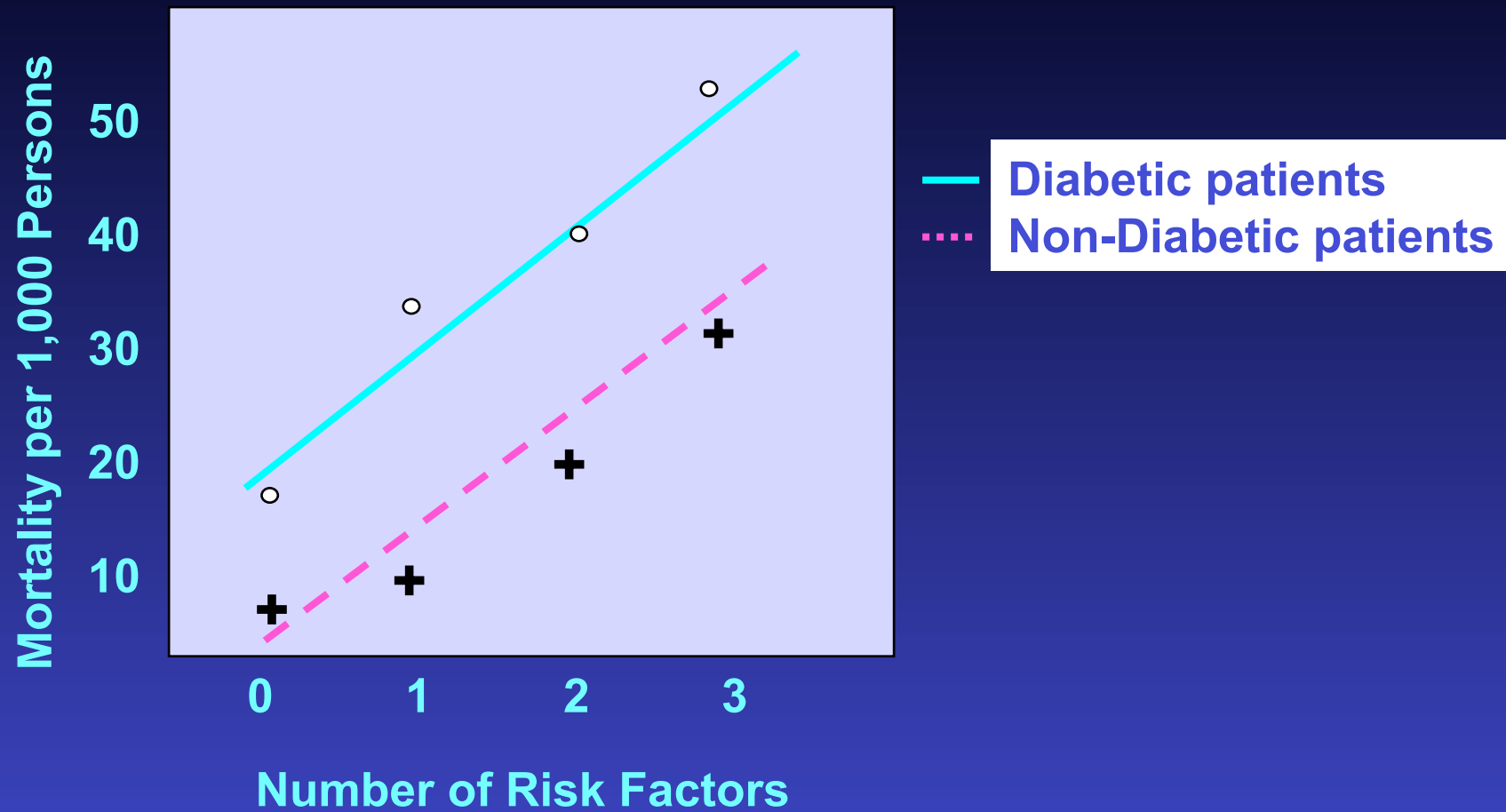
**CHD Mortality**  
Incidence (%) in 3.5 years



**All CHD Events**  
Incidence (%) in 3.5 years



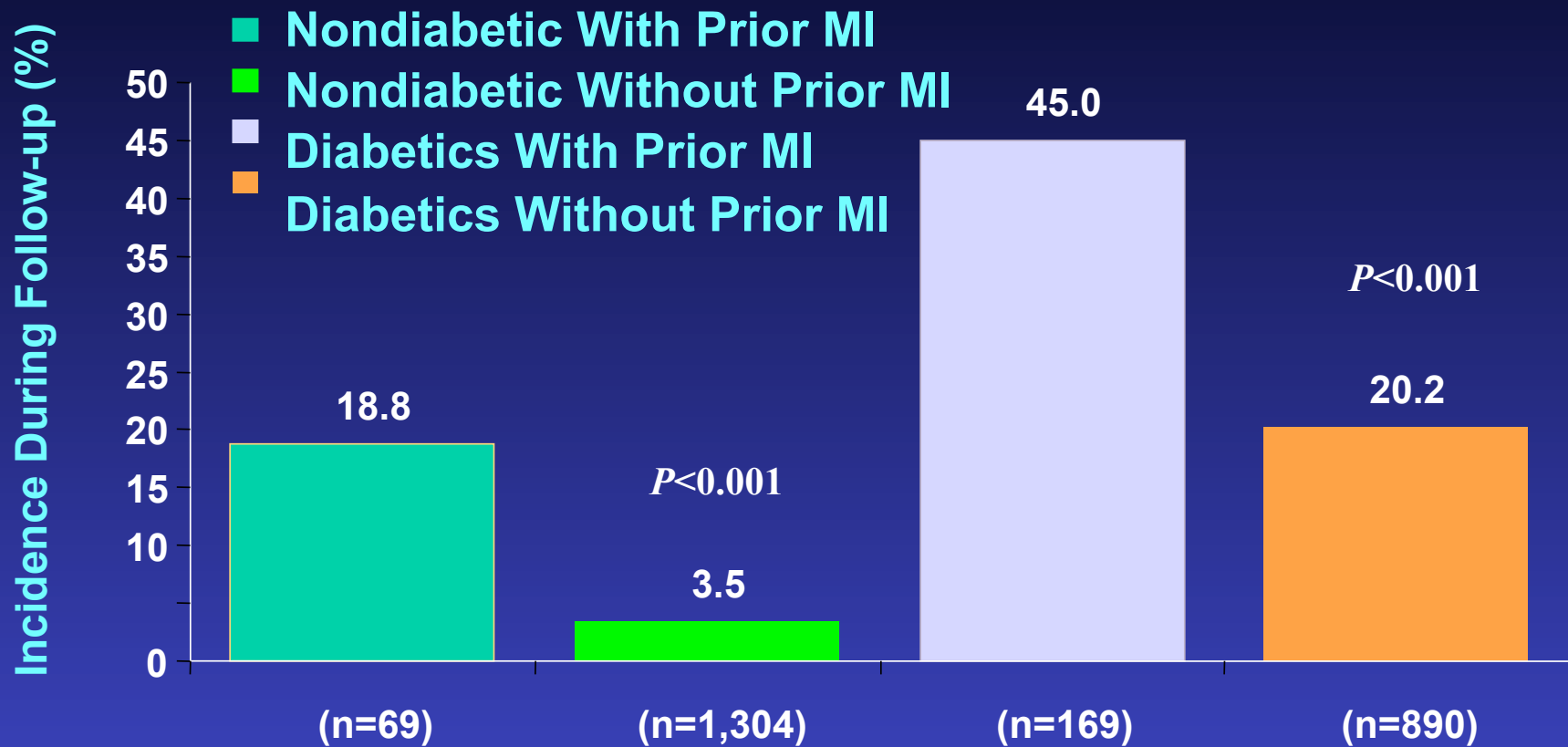
# Impact of Diabetes on Cardiovascular Mortality



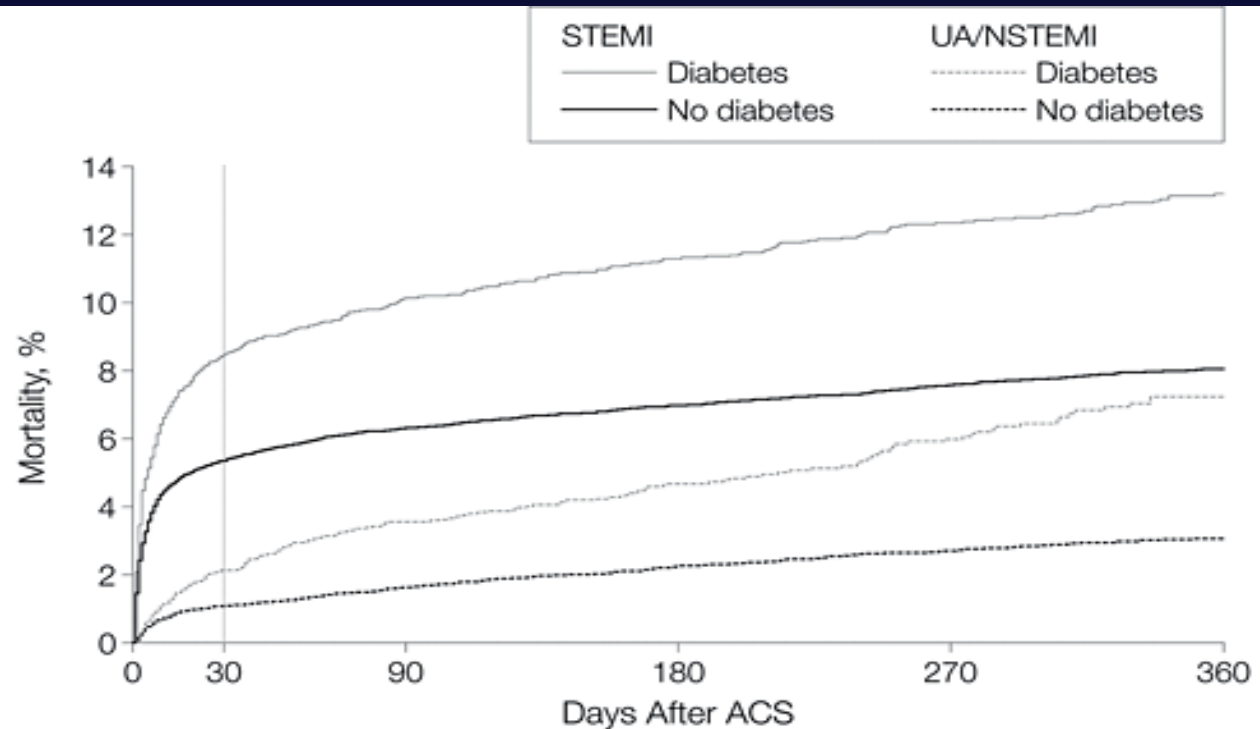
Stamler J, et al. *Diab Care* 1979;2:142

Stamler J, et al. *Diab Care* 1993;16:434-444.

# Incidence of Fatal or Nonfatal MI in Relation to History of MI in Nondiabetic vs Diabetic Subjects: 7-Year Follow-up



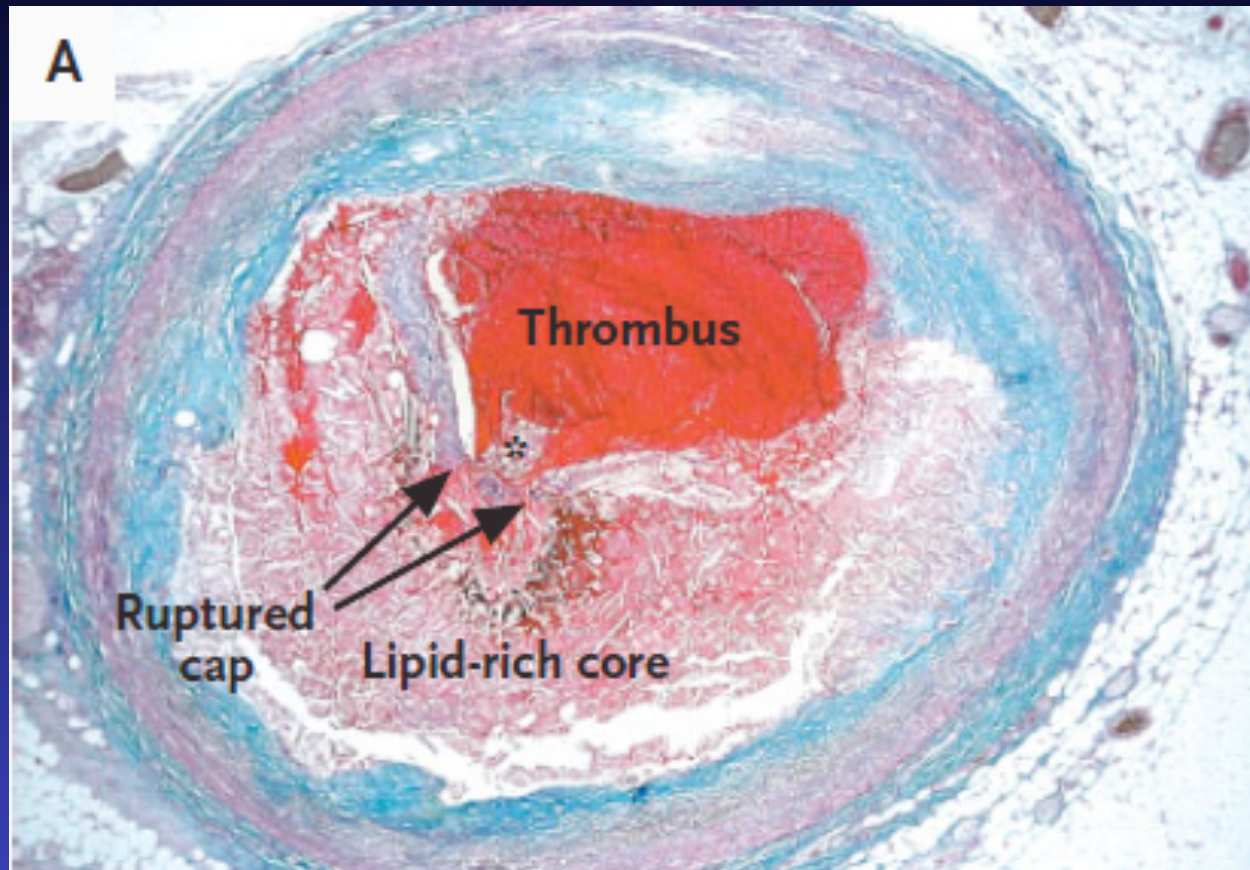
# Diabetes and Mortality Following Acute Coronary Syndromes



| No. at Risk |  |       |       |       |       |       |      |
|-------------|--|-------|-------|-------|-------|-------|------|
| STEMI       |  |       |       |       |       |       |      |
| Diabetes    |  | 7156  | 6508  | 2947  | 2653  | 2118  | 1610 |
| No diabetes |  | 39421 | 37136 | 16685 | 15274 | 12276 | 9351 |
| UA/NSTEMI   |  |       |       |       |       |       |      |
| Diabetes    |  | 3457  | 3313  | 2923  | 2339  | 1317  | 924  |
| No diabetes |  | 12002 | 11658 | 10505 | 8191  | 5141  | 4008 |

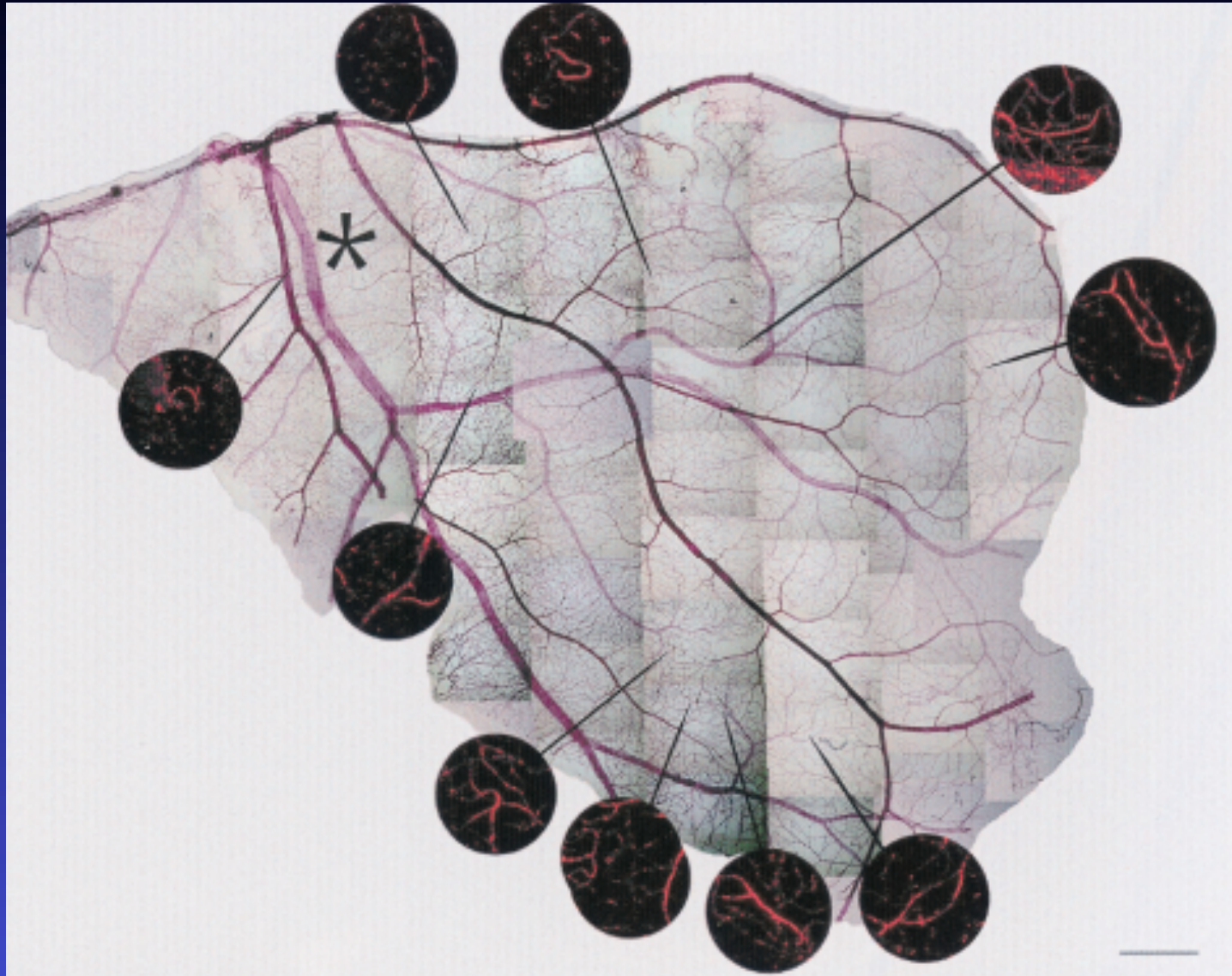
# Diabetes as a prothrombotic condition

# Thrombosed Coronary Artery



*N Engl J Med* 2005; 352:1685-95

# Large Thrombi in the Retina of a Diabetic Patient



*Diabetes* 2001; 50:1432-39

# DIABETES

## Decreased Fibrinolysis

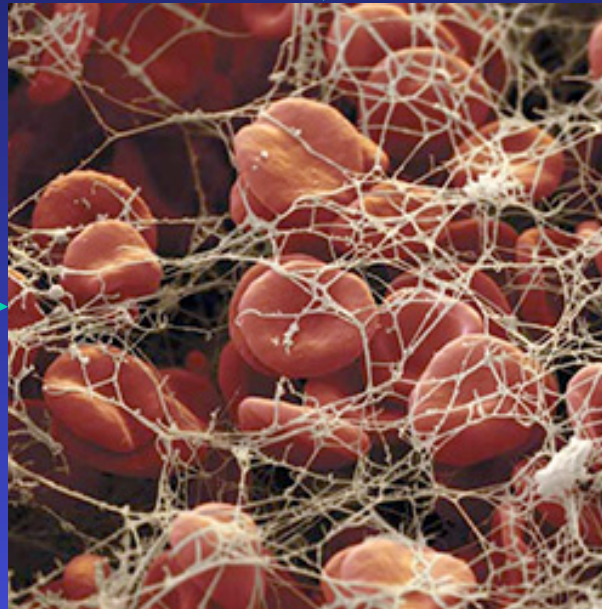
*(elevated levels of PAI-1,  
decreased levels of tPA)*

## Hypercoagulability

*(increased levels of  
fibrinogen, factor VII)*

## Endothelial Dysfunction

*(increased levels of  
von Willebrand factor,  
ICAM-1, leucocyte  
adhesion molecules)*



## Platelet Abnormalities

*(increased  
aggregation and  
decreased  
sensitivity to anti-  
aggregants)*

# **Role of glycemic control in CVD prevention**

# Will treatment of hyperglycemia reduce diabetes complications?

- Microvascular
  - Overwhelming consensus is, “yes”
- Macrovascular
  - Emerging data suggest
    - “Yes” in patients with short duration of disease and low CVD risk
    - “No” in patients with longstanding diabetes and significant atherosclerotic disease

# Intensive Glycemic Control in Patients with Lower CVD Risk

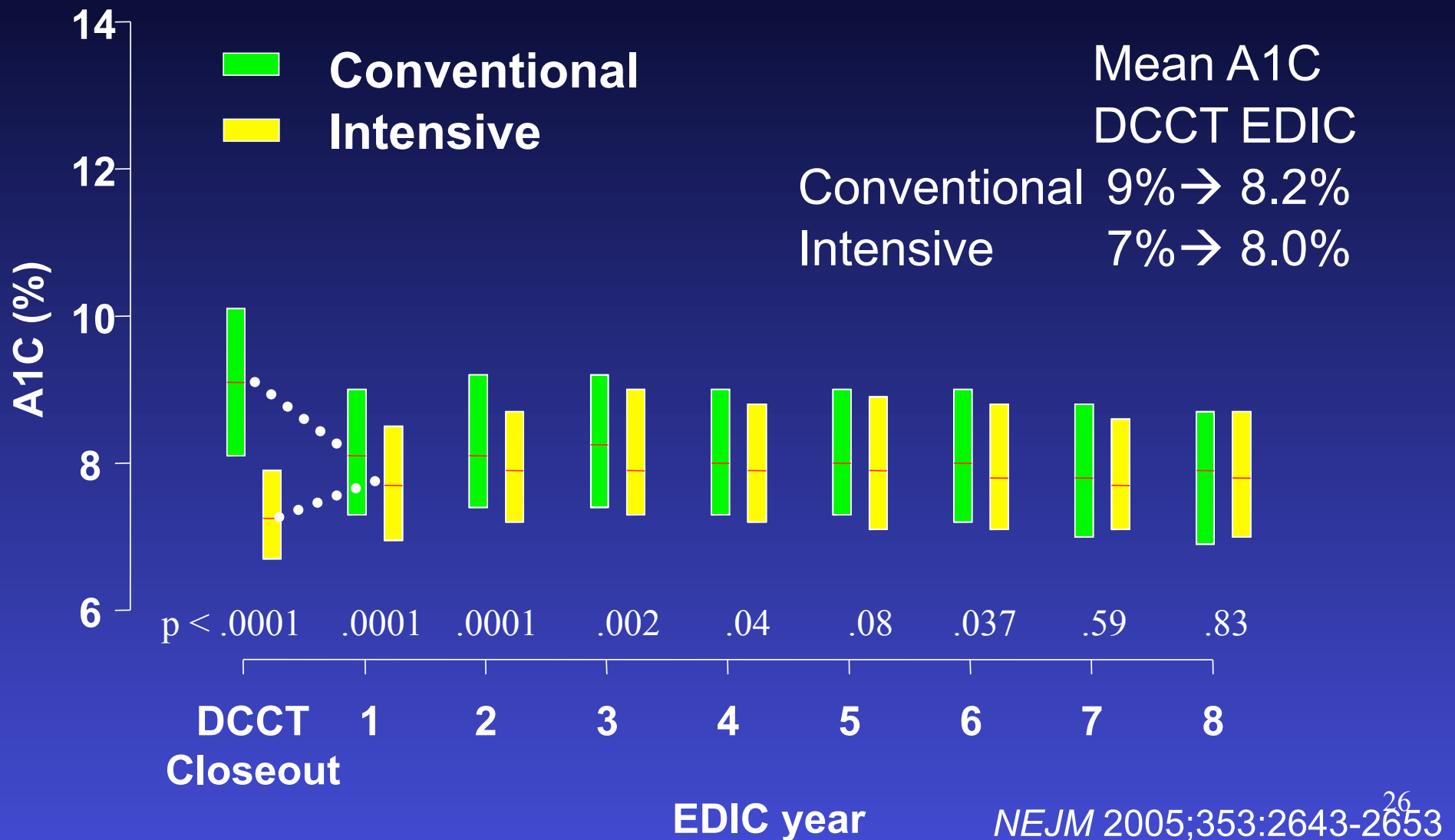
| Randomized controlled trials | DCCT (T1DM)       | UKPDS (T2DM) |
|------------------------------|-------------------|--------------|
| A1C (%)                      | 9 vs. 7           | 8 vs. 7      |
| Average follow-up (yrs)      | 6.5               | 10           |
| Microvascular complications  | Decreased         |              |
| <b>CVD events</b>            | <b>No benefit</b> |              |

DCCT. NEJM, 329(14), 1993; UKPDS Group. Lancet, 352:837-853, 1998 <sup>24</sup>

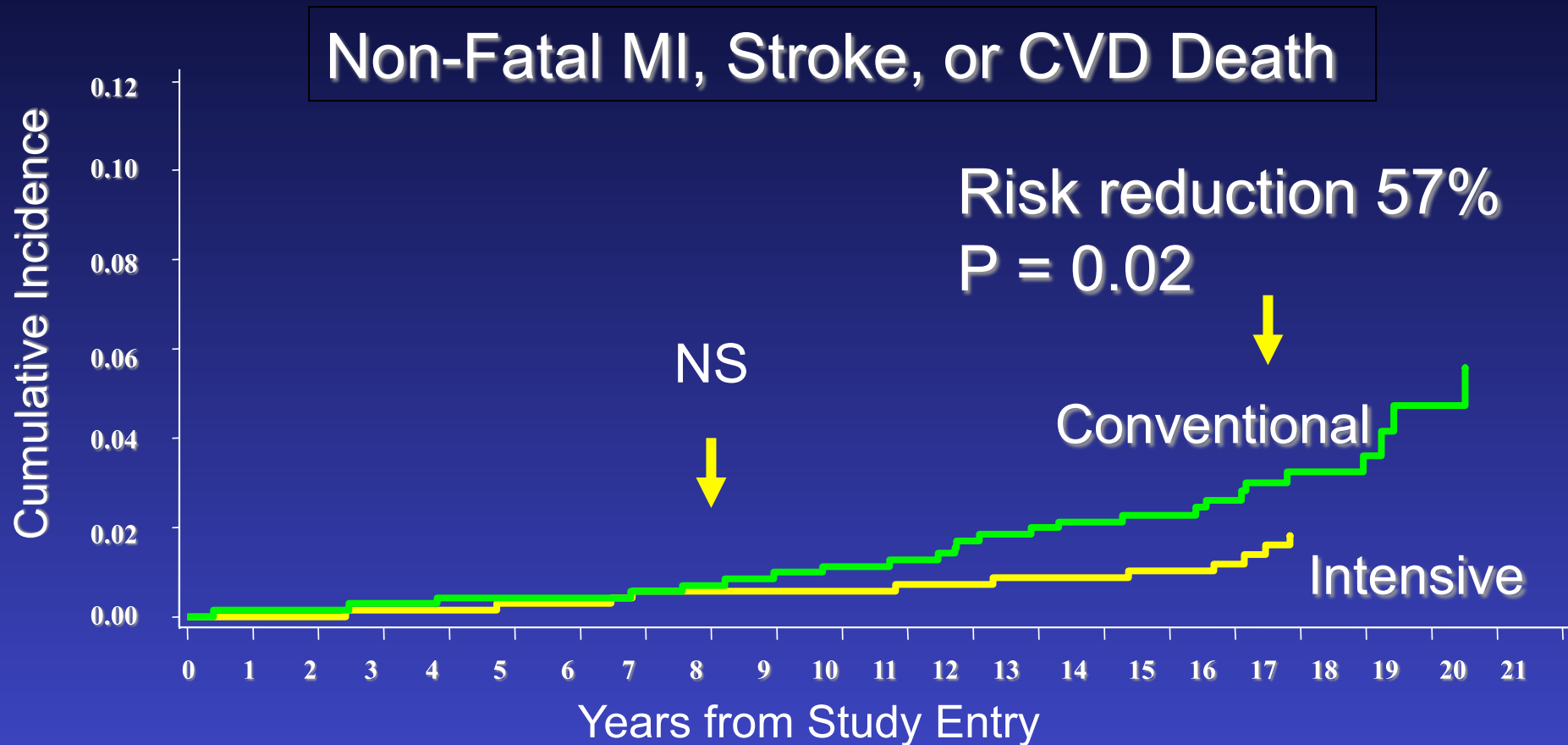
# Delayed Benefits (“Legacy Effect”) of Intensive Glucose Control

| Observational Monitoring Following RCT | DCCT (T1DM)          | UKPDS (T2DM)          |
|--|----------------------|-----------------------|
| A1C (%)                                | ~ 8                  |                       |
| Average follow-up (yrs)                | RCT 6.5<br>Monitor 9 | RCT 10<br>Monitor 8.5 |
| Microvascular complications            | Decreased            |                       |
| <b>CVD events</b>                      | <b>Decreased</b>     |                       |

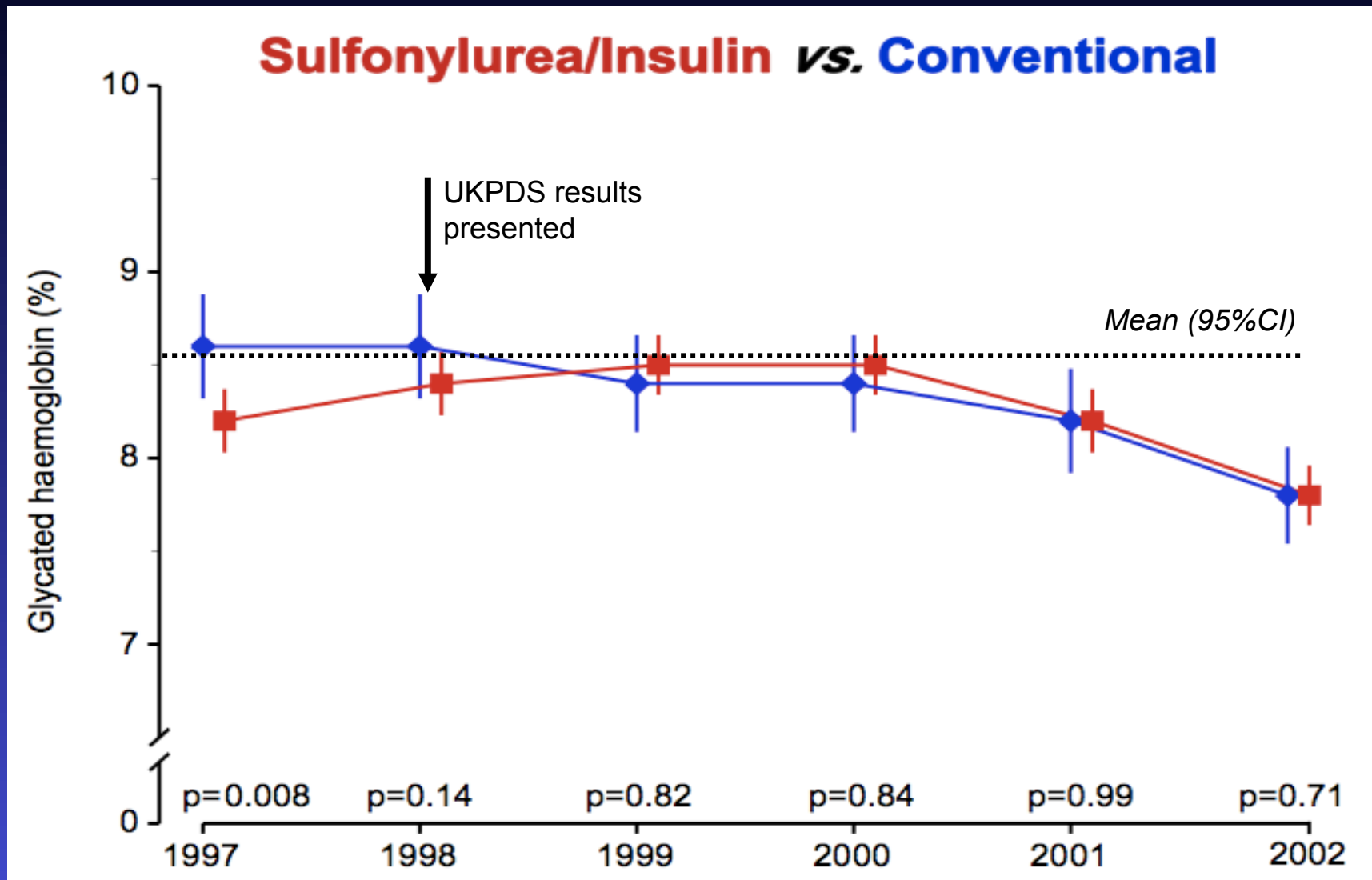
# DCCT/EDIC: No Differences in Post Trial A1C Levels



# EDIC: Decreased CVD Events Eight Years After DCCT Closeout



# UKPDS: Post-Trial Changes in A1C



# Legacy Effect of Earlier Glucose Control

After median 8.5 years post-trial follow-up

|                               |             | 1997         | 2007         |
|-------------------------------|-------------|--------------|--------------|
| Any diabetes related endpoint | <i>RRR:</i> | 12%          | 9%           |
|                               | <i>P:</i>   | 0.029        | <b>0.040</b> |
| Microvascular disease         | <i>RRR:</i> | 25%          | 24%          |
|                               | <i>P:</i>   | 0.0099       | <b>0.001</b> |
| Myocardial infarction         | <i>RRR:</i> | 16%          | 15%          |
|                               | <i>P:</i>   | <b>0.052</b> | <b>0.014</b> |
| All-cause mortality           | <i>RRR:</i> | 6%           | 13%          |
|                               | <i>P:</i>   | <b>0.44</b>  | <b>0.007</b> |

# Intensive vs. Standard Glucose Control in T2DM With High CVD Risk

|                         | <b>ACCORD<sup>1</sup></b> | <b>ADVANCE<sup>2</sup></b> | <b>VADT<sup>3</sup></b> |
|-------------------------|---------------------------|----------------------------|-------------------------|
| N                       | 10,251                    | 11,140                     | 1,791                   |
| Age (yrs)               | 62                        | 66                         | 60                      |
| Duration diabetes (yrs) | 10                        | 8                          | 11.5                    |
| Baseline A1C (%)        | 8.1                       | 7.2                        | 9.4                     |
| Intense A1C goal (%)    | <6                        | <u>≤</u> 6.5               | action >6.5             |
| CVD risk                | High                      |                            |                         |
| Follow-up (yrs)         | 3.5 (stopped early)       | 5                          | 5.6                     |

# Intensive vs. Standard Glucose Control in T2DM With High CVD Risk

|                     | <b>ACCORD<sup>1</sup></b>        | <b>ADVANCE<sup>2</sup></b> | <b>VADT<sup>3</sup></b> |
|---------------------|----------------------------------|----------------------------|-------------------------|
| A1C standard (%)    | 7.5                              | 7.0                        | 8.5                     |
| A1C intensive (%)   | 6.4                              | 6.3                        | 6.9                     |
| Hypoglycemia        | Greater with intensive treatment |                            |                         |
| Weight gain         |                                  |                            |                         |
| <b>CVD benefit*</b> | <b>None</b>                      |                            |                         |
| <b>Death*</b>       | <b>Increased</b>                 | No difference              |                         |

1. NEJM 2008;358:2545-59, 2. NEJM 2008;358:2560-72, 3. NEJM 2009;360:129-139

\*intensive vs. standard

# Intensive Glucose Control Summary 1

- Benefits on microvascular and neuropathic complications is well established
- Evidence for CVD benefit is strongest in T1DM
  - There may be a “legacy effect” in which benefits of glucose control do not appear until years following intensive glucose control

# Intensive Glucose Control Summary 2

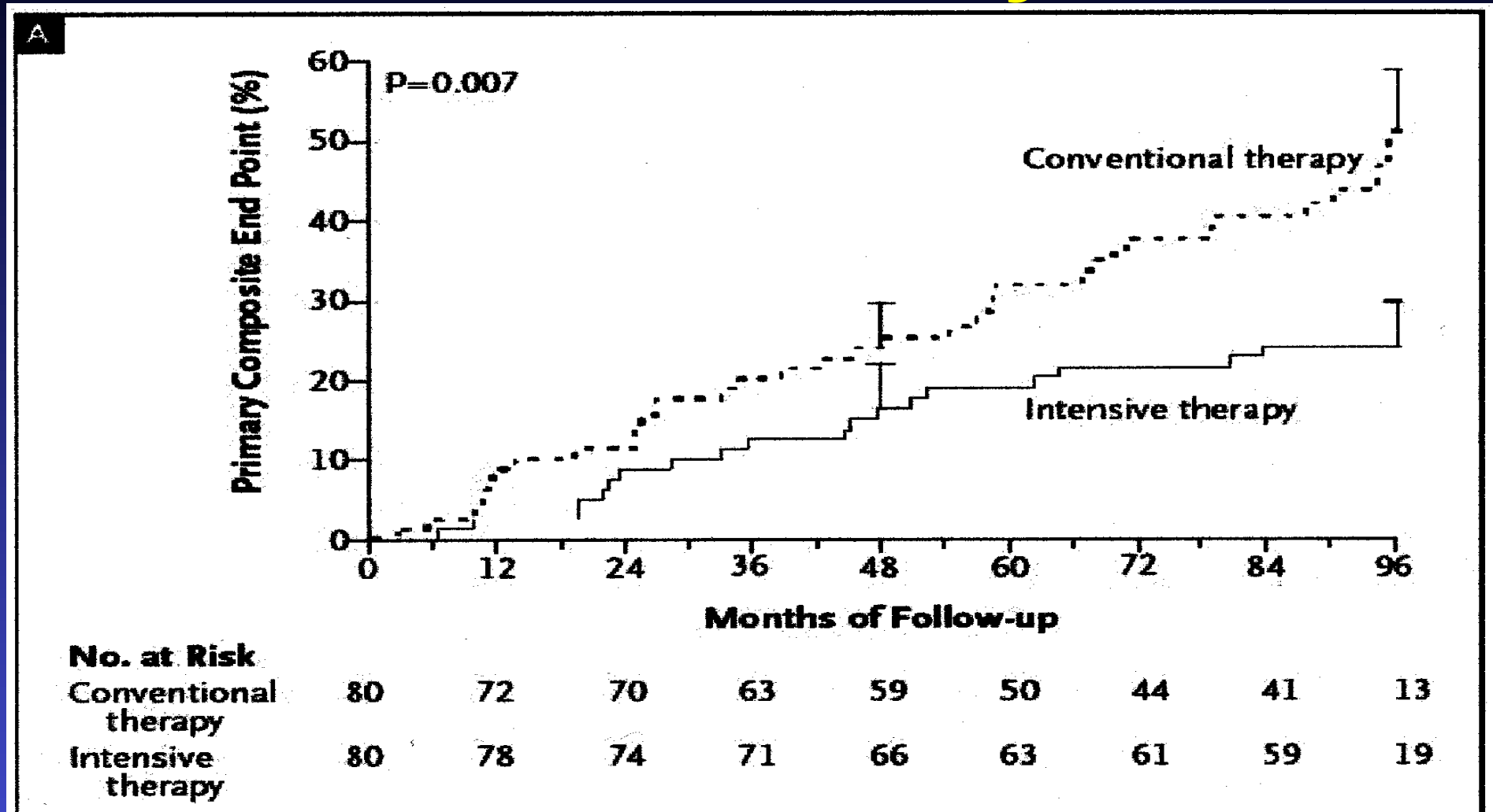
- Possible CVD benefits in patients with shorter duration of diabetes and without established atherosclerotic disease
- Potential risks may outweigh benefits when
  - Long duration of diabetes
  - Known history of severe hypoglycemia
  - Advanced atherosclerosis,
  - Advanced age/frailty

**Can we prevent CVD in diabetic patients?**

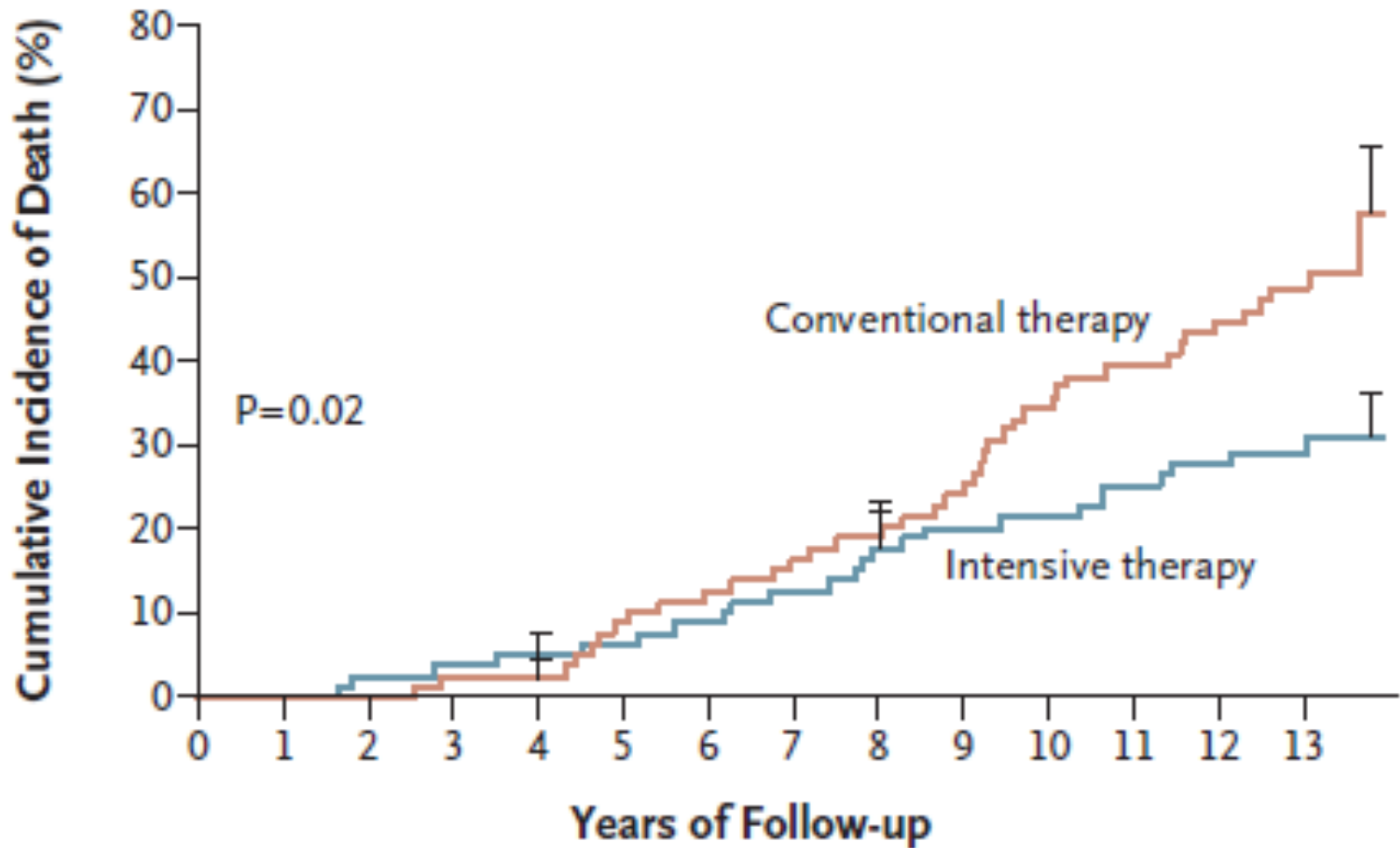
# Steno-2 Study

- 160 type 2 diabetic patients randomized to conventional or intensive treatment
- Mean follow-up of 7.8 years
- Treatment goals (changed over time) included A1c, BP and lipids
- Interventions in intensive therapy group:
  - Diet (total fat < 30%, saturated fats < 10%)
  - Exercise (30 minutes 3-5 days/week)
  - Aspirin 150 mg
  - Captopril 50 mg bid
  - Vitamin C, Folic Acid, D-a-tocopherol, Chromium<sup>35</sup>

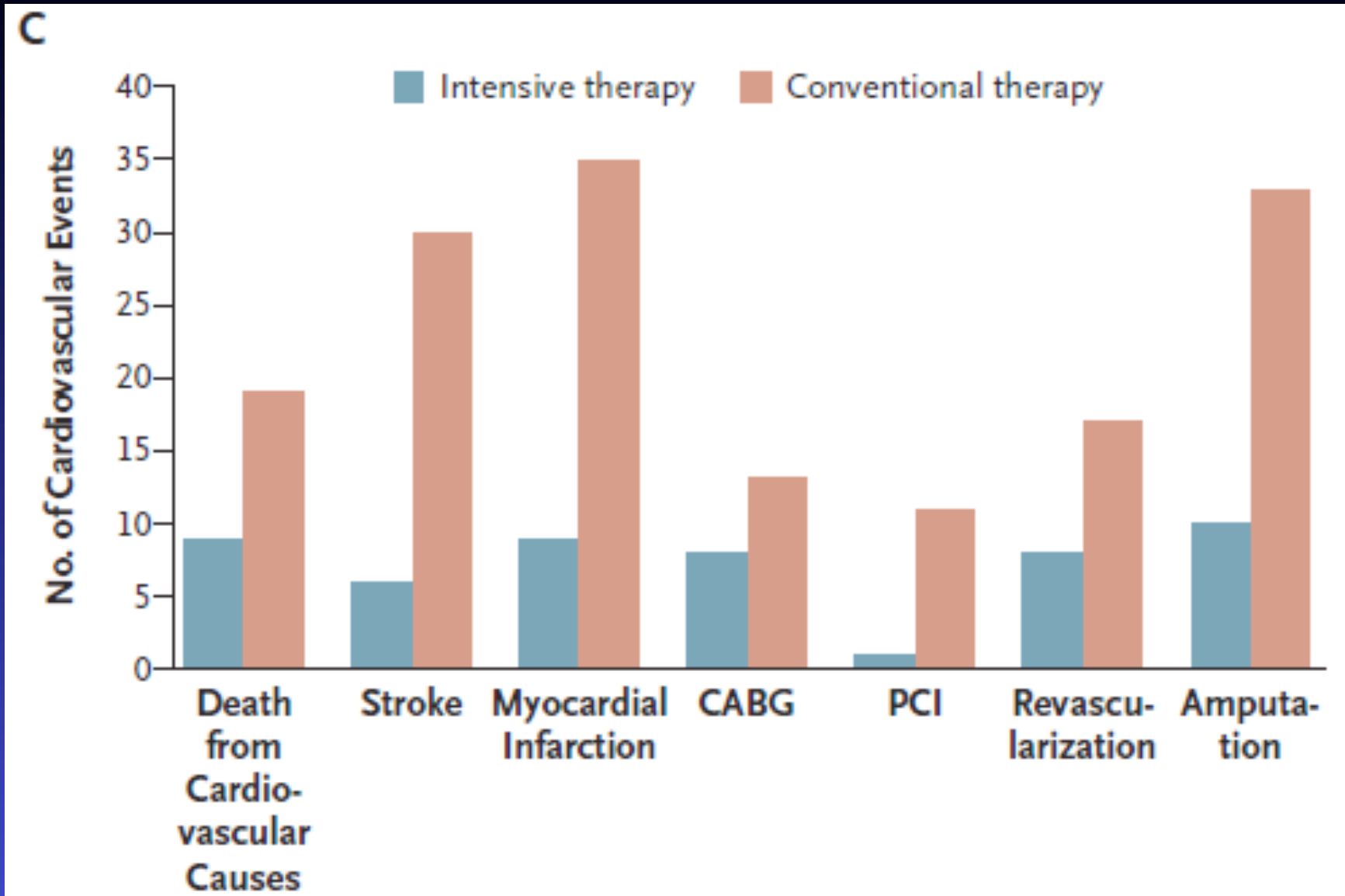
# Steno-2 Study



# Steno-2 Study



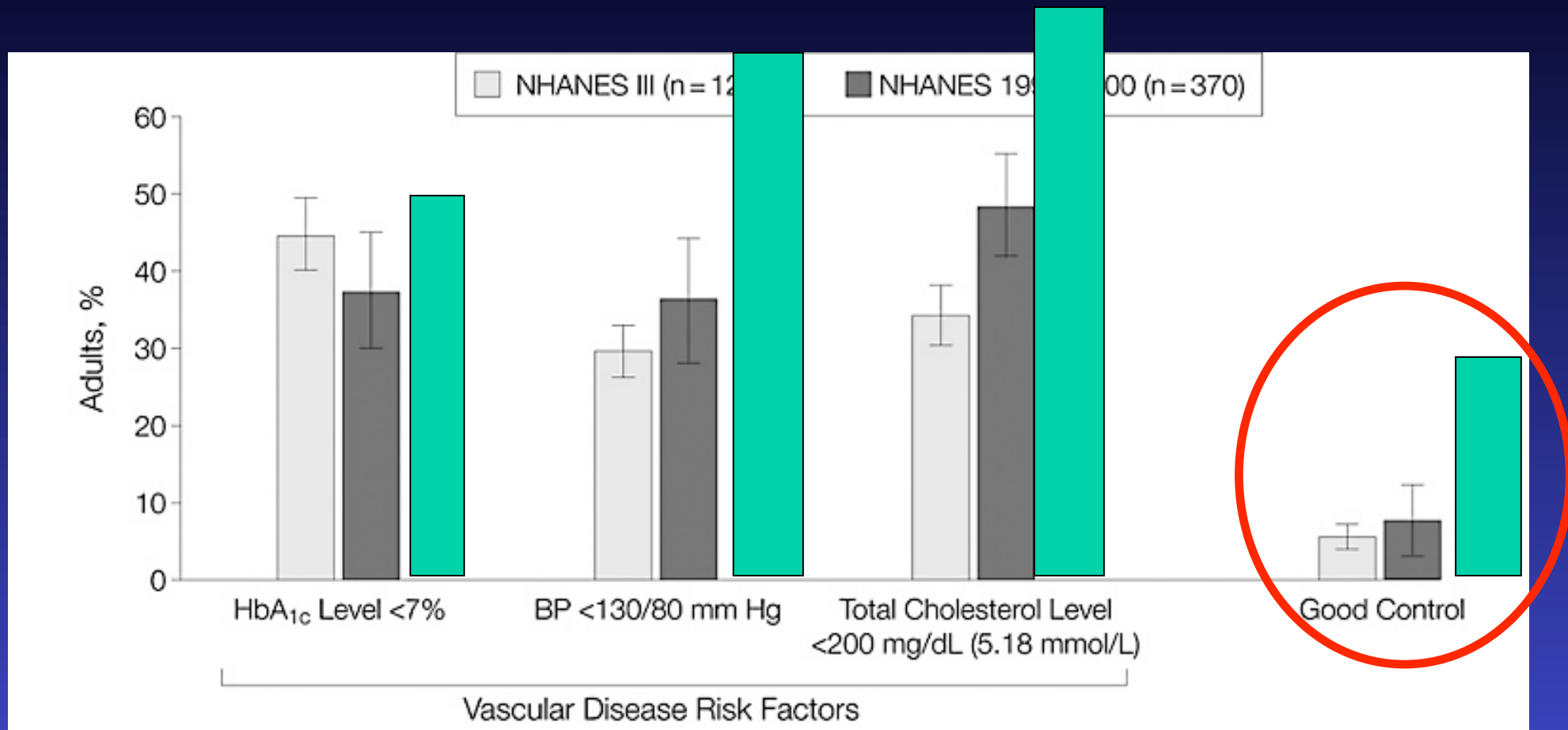
# Steno-2 Study



**Are we achieving these goals?**

***Not really!***

# Percentage of Adults with Recommended Levels of Risk Factors



# Comments or Questions?

