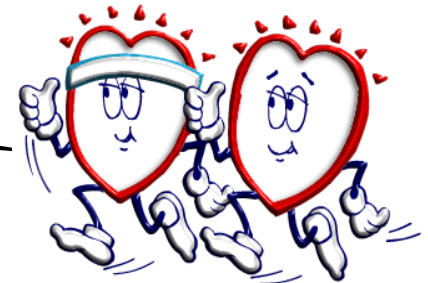


BI 358 Lecture 11

Next session, Dr. Padgett, Medical Director OHVI! Hip! Hip! Hooray!!



I. Announcements Presentations Group I today!
Tight time-frame + immediate feedback? Q?

II. CVD-Atherosclerosis Connections: Lecture 10 slides \geq # 38

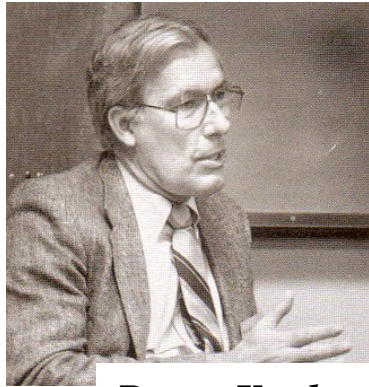
III. Endocrinology Overview G&H ch 74+75, LS, Norris, Fox...

- A. Endocrine vignette: Cushing's Syndrome LS
- B. What's an endocrine? Hormone criteria & classifications?
- C. Mechanisms of hormonal action fig 74-6, 74-2, 74-7,...
- D. Endocrinology focuses on the relationship between the Hypothalamus - Controller \rightarrow Pituitary - Subcontroller
- E. Endocrine organ & hormonal overview fig 74-1, tab 74-1
- F. Hypothalamus-Post & Ant Pituitary fig 75-9, 75-4, 75-2
- G. Anterior pituitary hormone functions tab 75-1, Fox + LS
- H. Negative feedback loops G&H p 885 + LS
- I. Growth Hormone (GH/STH) fig 75-5, 75-6, tab 75-3
Body builder's dream or fountain of youth? Neither!

IV. Peripheral Endocrine Organs G&H ch 76, 77, 78

- A. Pancreas: insulin vs. glucagon, diabetes, G&H ch 78 + Fox
- B. Thyroid: T3 & T4 G&H fig 76-2 thru fig 76-9 + DC
- C. Adrenal cortices G&H fig 77-1 & 77-2 + DC

Bruce Kottke's Bathtub Analogy



Bruce Kottke

5 forms of cholesterol:

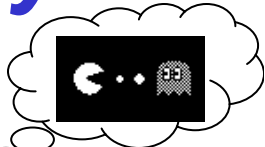
Chylomicrons, VLDL, LDL, IDL, HDL

β



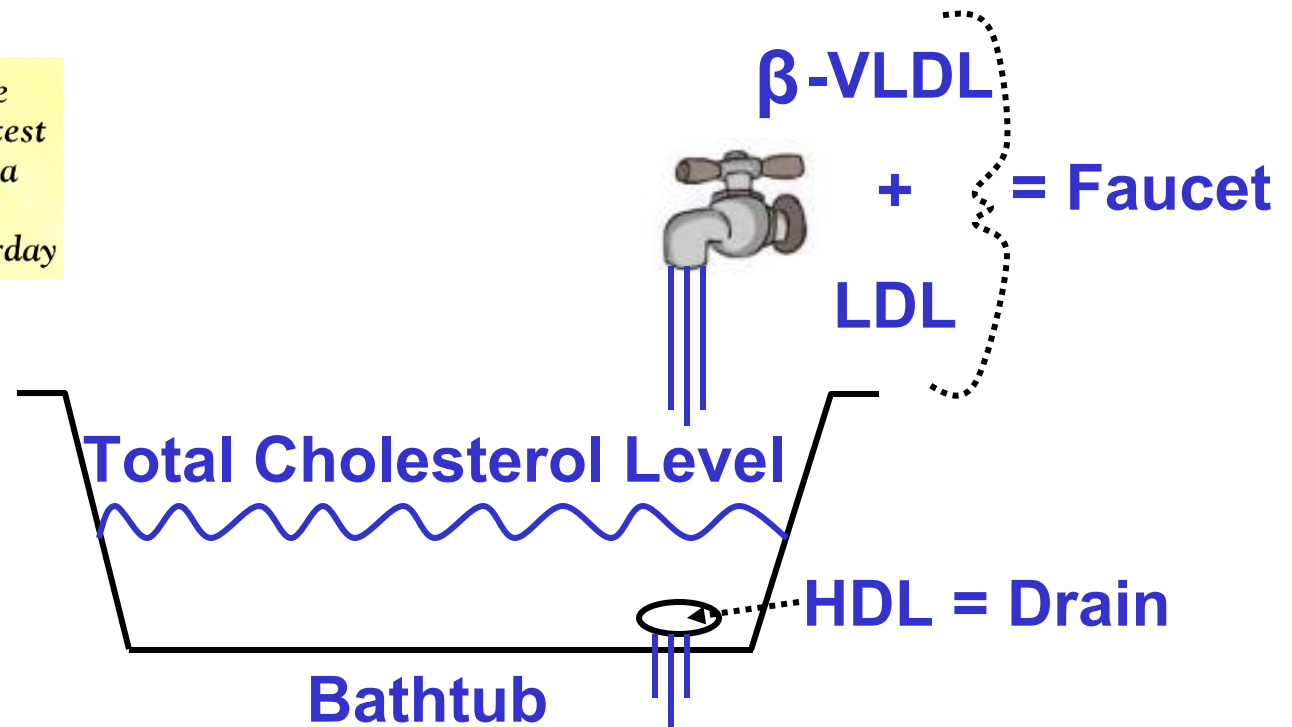
Atherogenic

Anti-Atherogenic



Biological Artifact!?

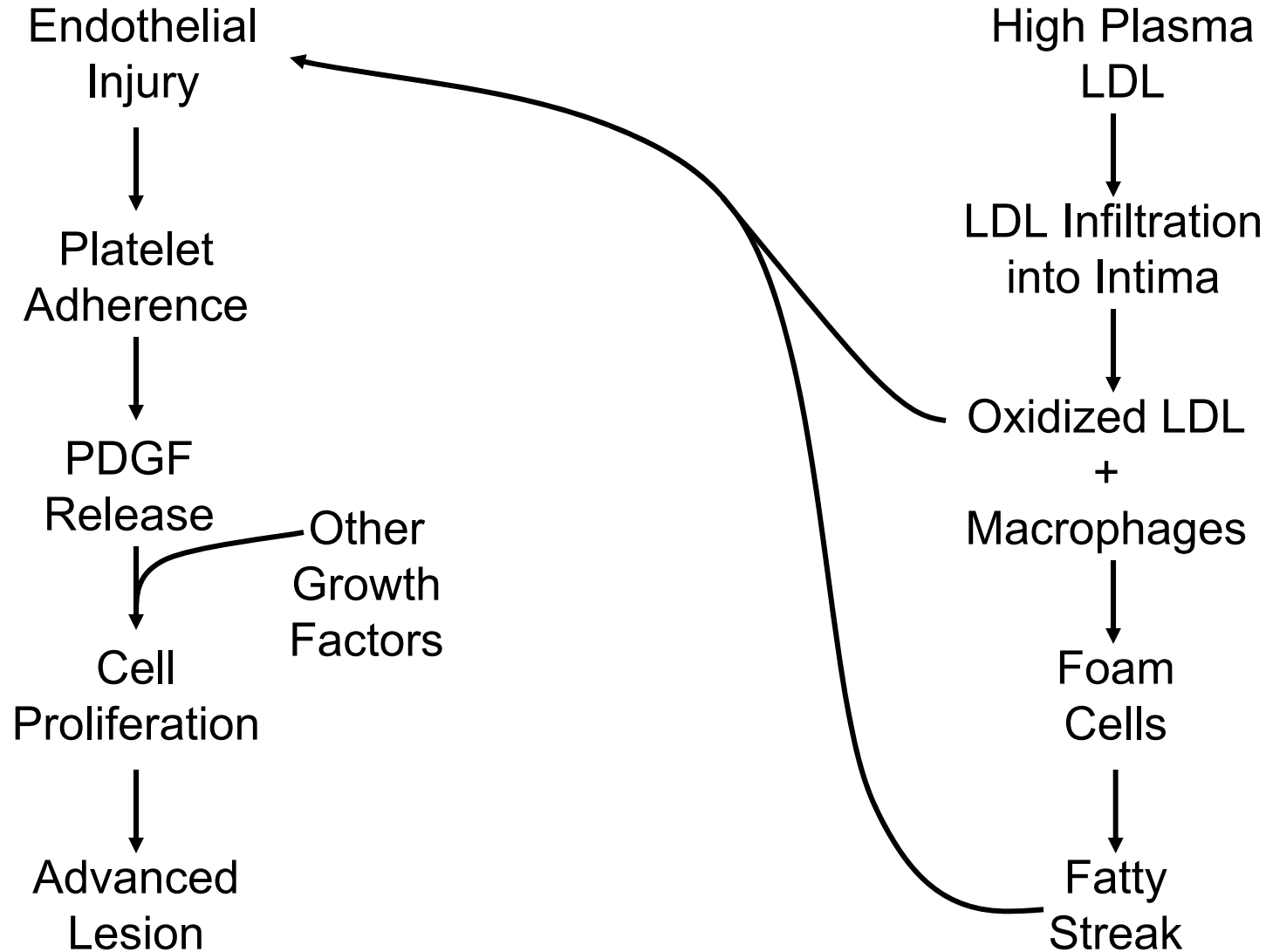
"I don't think the total cholesterol test by itself is worth a damn."
—Eliot Corday



Historical Hypotheses for Atherosclerosis Development

Ross & Glomset

Steinberg & Witztum



How Inflammation Attacks the Heart

1 LDL Oxidized

Oxidized LDL cholesterol creates the "injury" by burrowing into the artery wall. Cigarette smoking, high blood pressure, and high blood sugar make the injury worse.

2 Monocytes Migrate 3 Monocytes → Macrophages

In response to the injury, the immune system sends in a team of inflammatory cells, including white blood cells called monocytes.

Monocytes migrate into the artery wall, where they turn into macrophages. The macrophages' job: gobble up the LDL cholesterol.

4 Fatty Streak

The macrophages, now stuffed with LDL cholesterol, form a "fatty streak" in the artery wall.

5 Fibrous Plaque

Over the decades, more cholesterol, connective and elastic tissue, calcium, and cell debris accumulate and turn the fatty streak into plaque. As the artery tries to heal itself, smooth muscle cells migrate in to cover the plaque, forming a fibrous cap around it.

6 Cap Breakdown

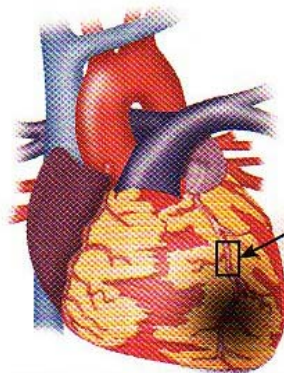
Macrophages kill the smooth muscle cells and release enzymes that break down the fibrous cap.

7 Cap Rupture

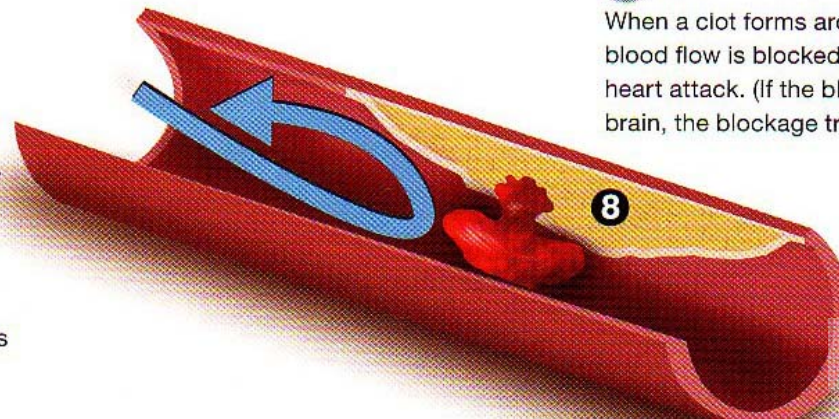
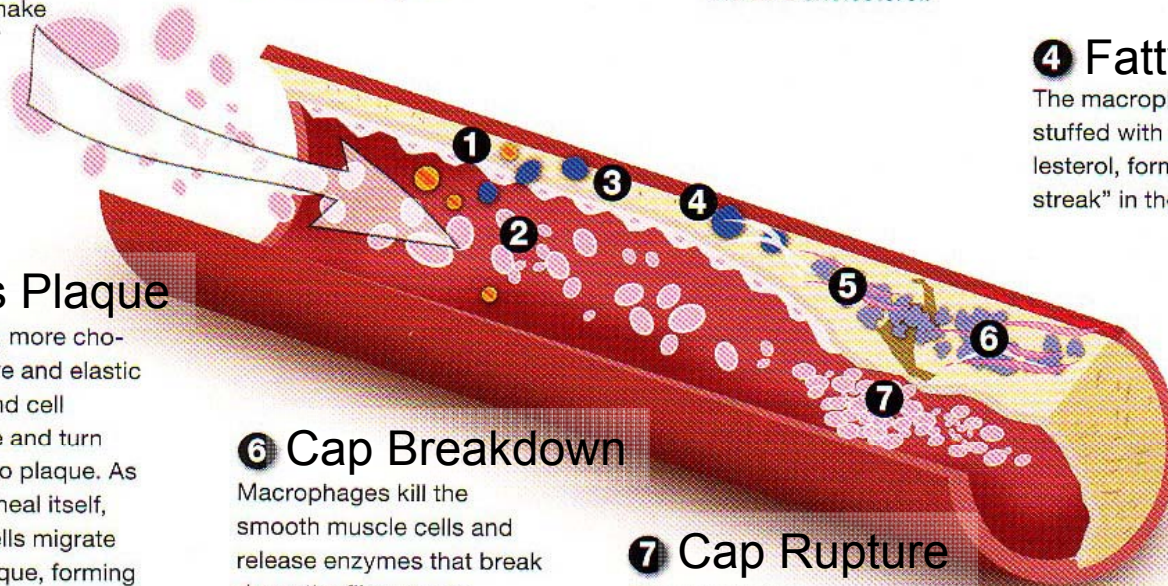
The cap ruptures.

8 Clot Formation

When a clot forms around the rupture, blood flow is blocked, which triggers a heart attack. (If the blocked artery feeds the brain, the blockage triggers a stroke.)



Coronary artery (supplies blood and oxygen to the heart muscle).





TROUBLE AHEAD?

HOW TO KEEP YOUR BRAIN SHARP

So far, no one has found a magic bullet to stop Alzheimer's disease, which gums up the brain with protein clumps and tangles. But it's not just clumps and tangles.

Brain Basics

Plaques and tangles. Those are the classic hallmarks of Alzheimer's disease.

The plaques are clumps of a protein fragment called beta-amyloid. The tangles are clusters of misshapen "tau" proteins that show up later in the disease.

But plaques and tangles alone don't explain what happens to many aging brains.

"Thirty percent of people over the age of 70 have elevated beta-amyloid and are cognitively normal," says David Knopman, professor of neurology at the Mayo Clinic in Minnesota.

Scientists aren't sure why.

"The most prevalent idea is that amyloid deposits are only the initiating step

Damage to the brain's blood vessels—often due to high blood pressure, smoking, or diabetes—can also play a role, not just in dementia but in milder memory loss as well.

Here's how to keep a clear head for as long as possible.

often assume that it's just Alzheimer's," notes Reed. "But it's uncommon to find people with dementia who just have a single pathology. Very often, it's mixed pathology."

The most common other problem: damaged blood vessels in the brain.^{1,2}

"The arteries become stiffened, narrowed, and sort of tortuous," says Reed. "It's much harder for the blood flow to occur normally."

That can lead to a stroke that's obvious, or to one that's never noticed. "Around

"In fact, some of the symptoms we think of as normal brain aging may be due to injury to the brain's blood vessels," he notes.

Researchers know the major threats. "The big risks for

vascular brain injury are smoking, high blood pressure, and diabetes," says Reed.

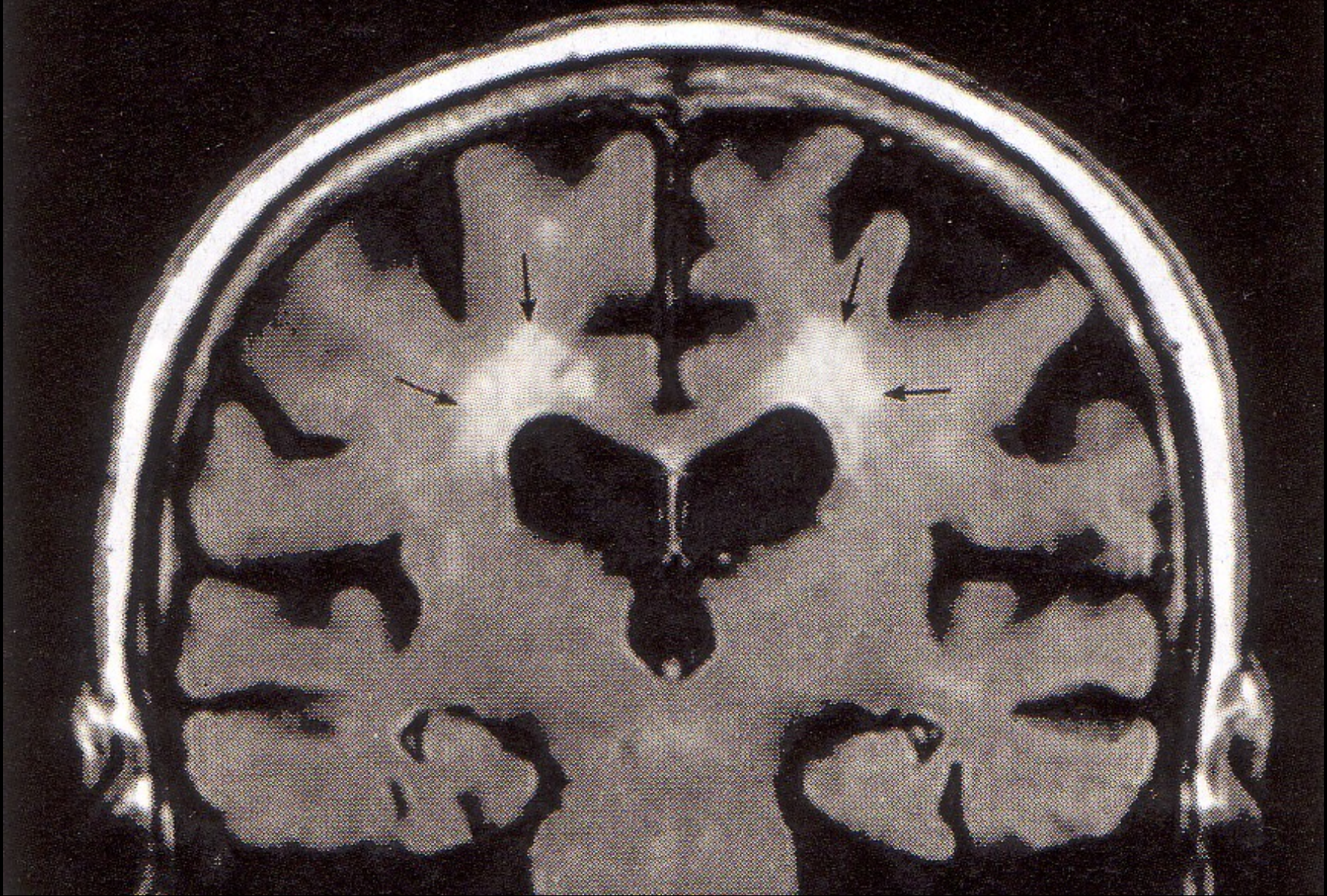
The causes of Alzheimer's pathology are more murky. But new evidence suggests that insulin may play a role.

Here's how to keep your brain in good working order.

1. Watch your blood pressure

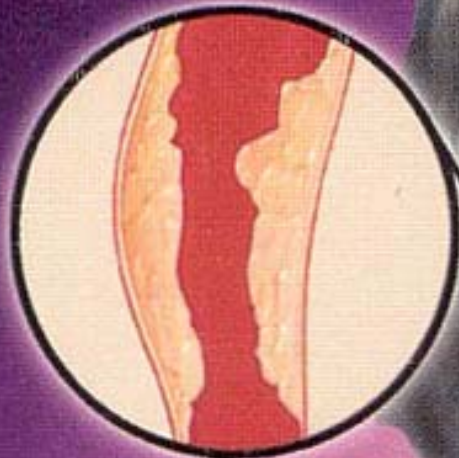
"There's a wealth of evidence that high blood pressure is a risk factor for late-life cognitive impairment," says Knopman.

MRI Hyperintensities, Hypertension & Dementia



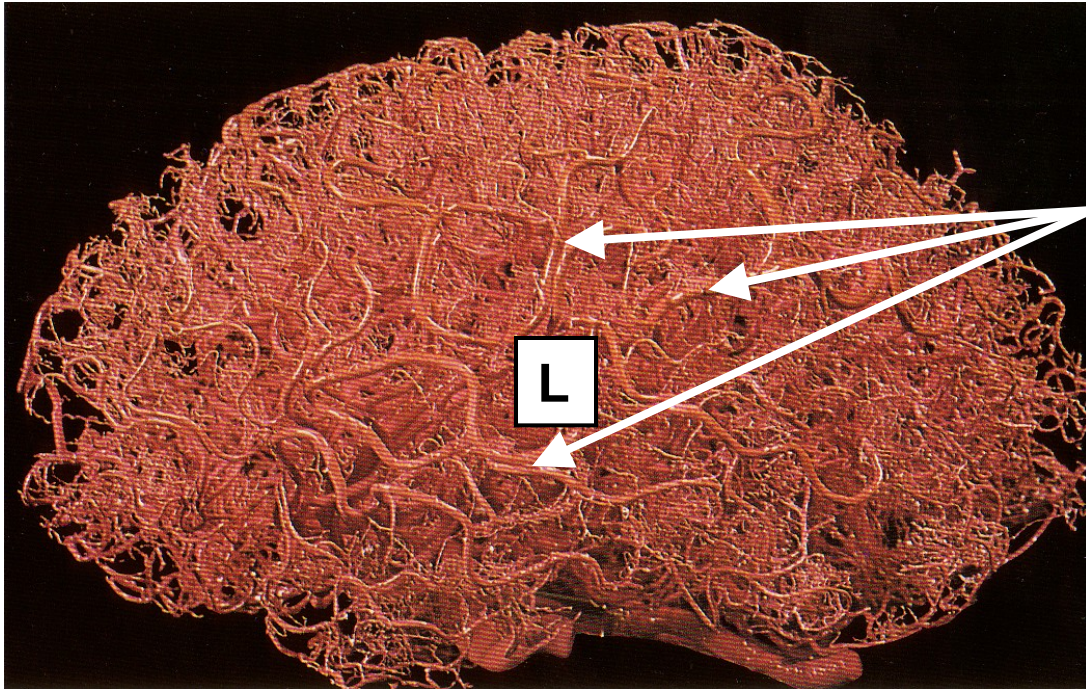
NAHL CSPI, Jan-Feb 2014

DISEASED CAROTID ARTERY



HEALTHY CAROTID ARTERY

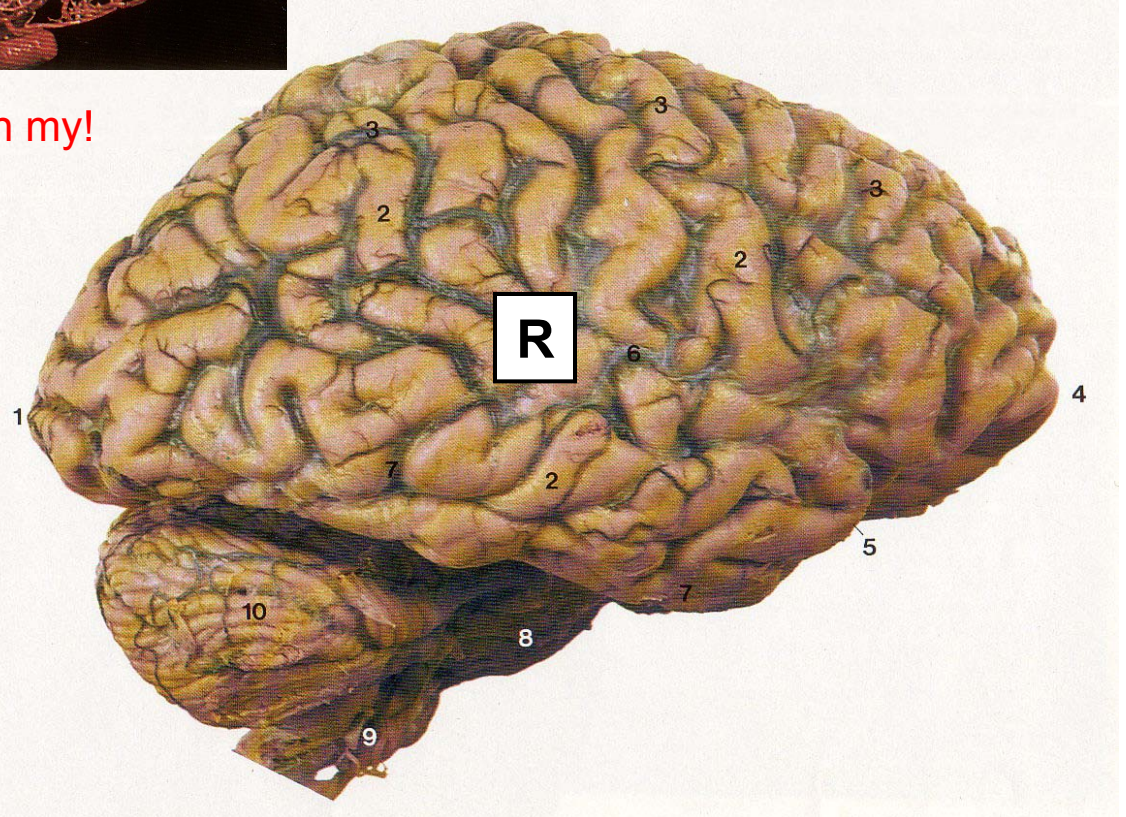
SOURCE: Lifeline Screening, 2007



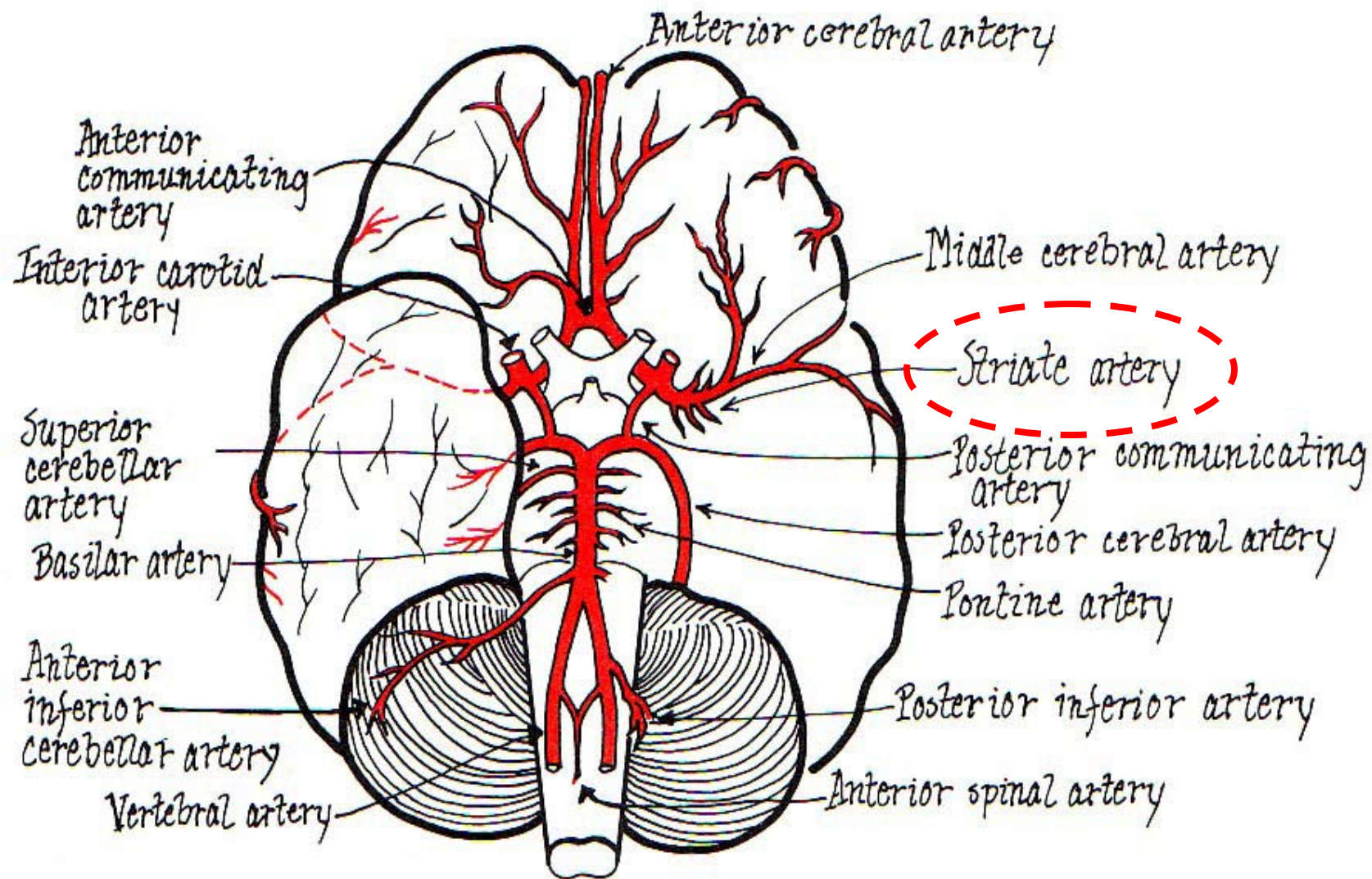
Middle Cerebral Artery Branches



...Cerebral vasculature! Oh my!



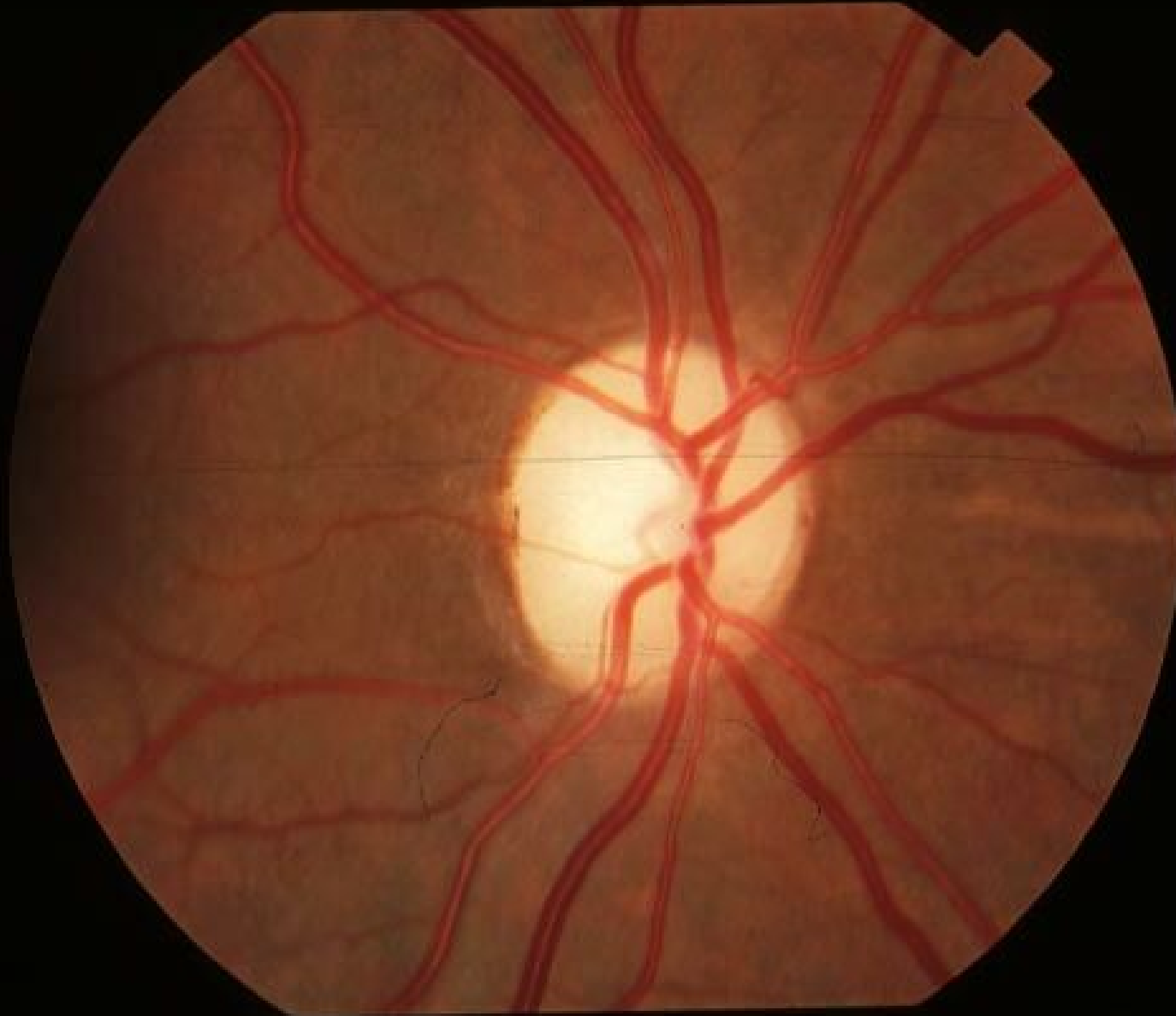
SOURCE: McMinn & Hutchins, 1977.



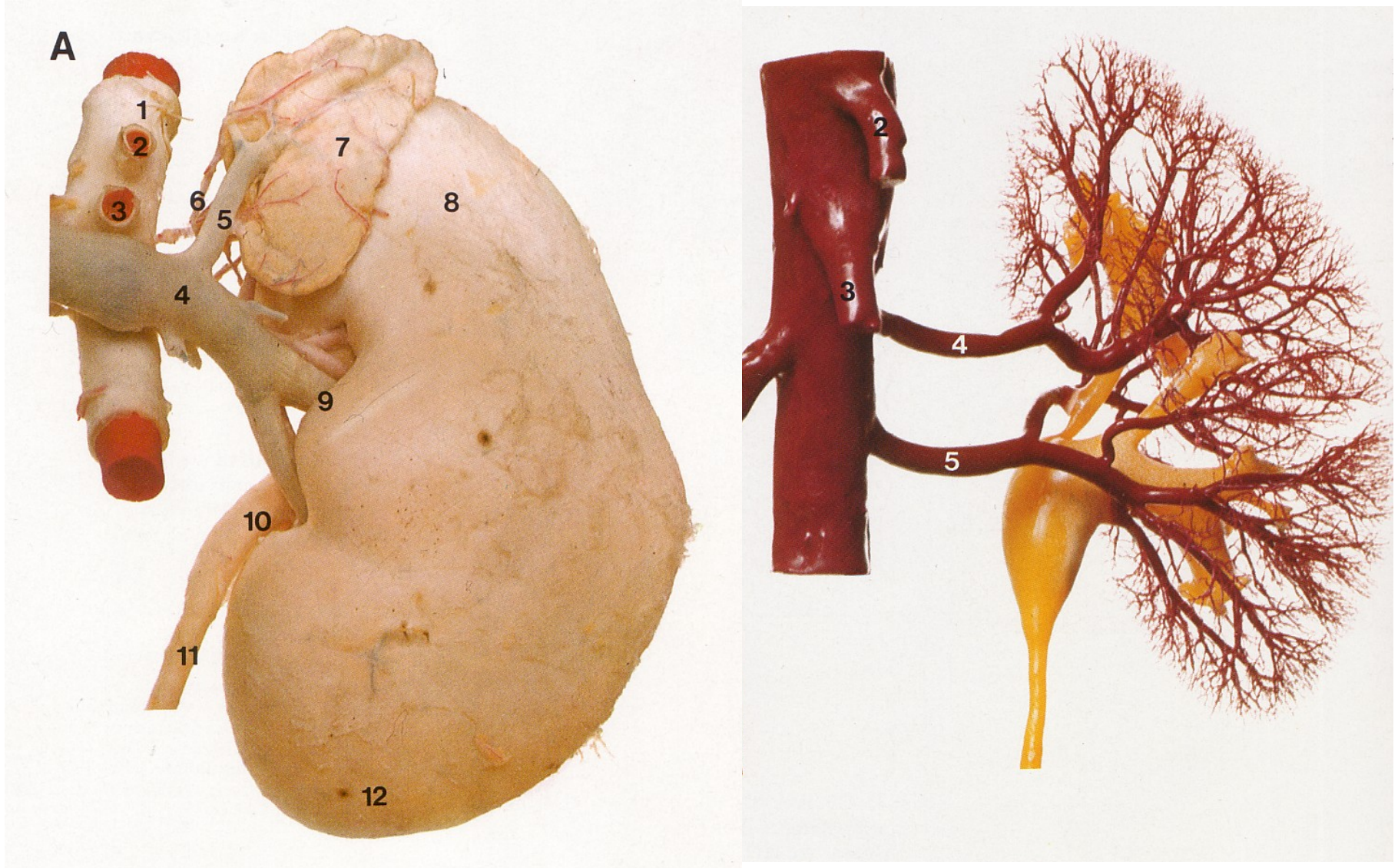
() = Artery of Stroke

SOURCE: Liebman, 1979.

The Window to the CV System?



Renal Vasculature



SOURCE: McMinn & Hutchins, 1977.

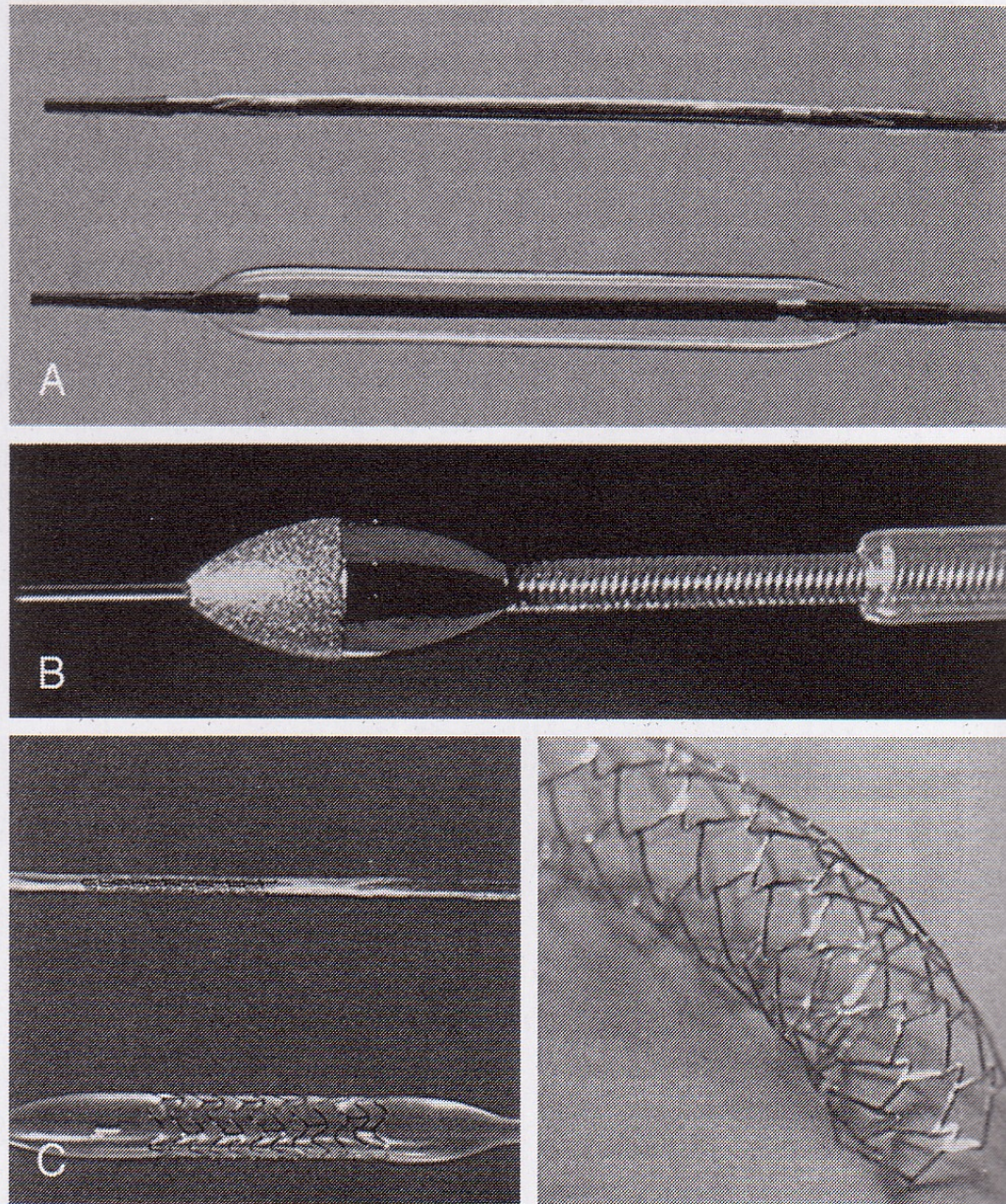
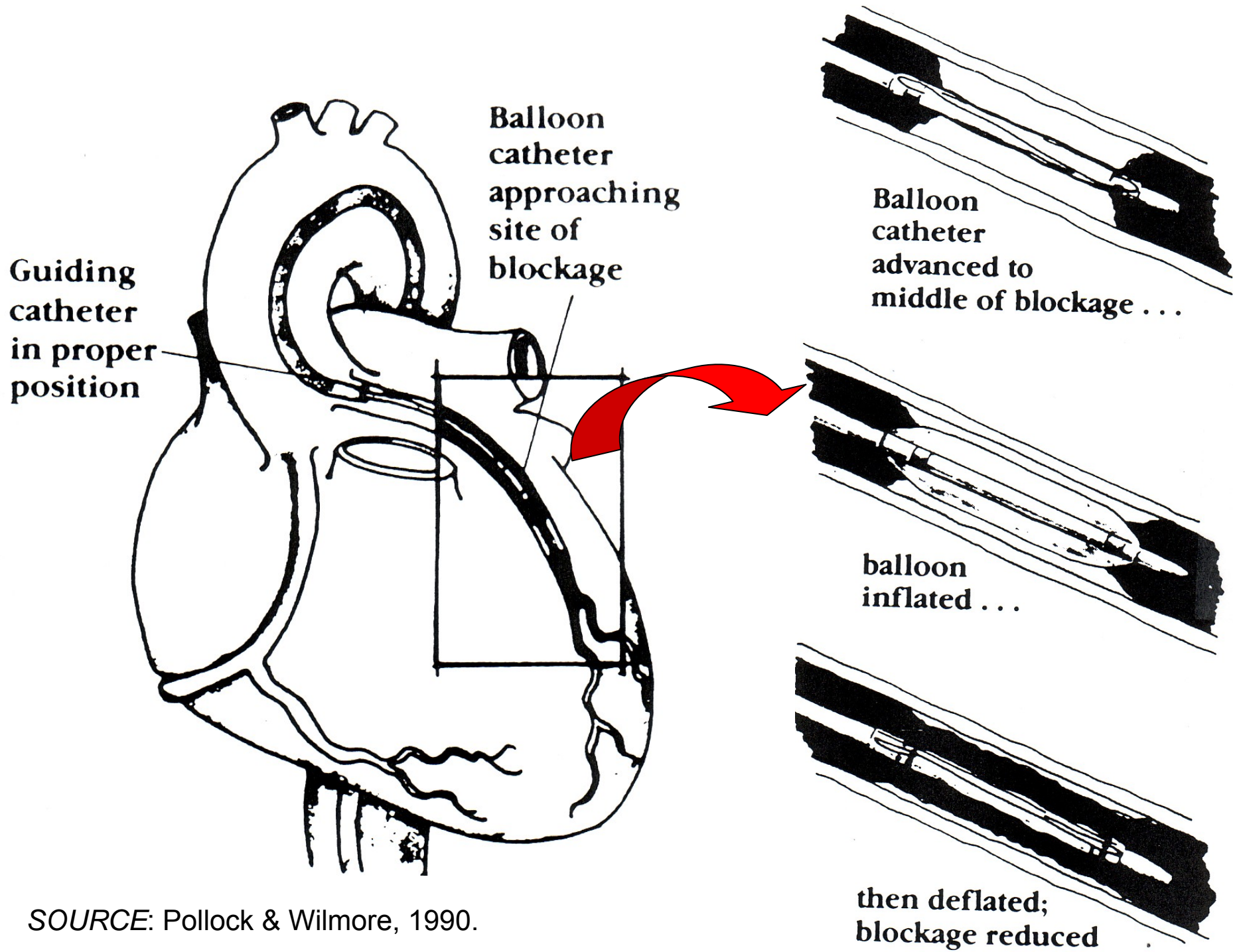
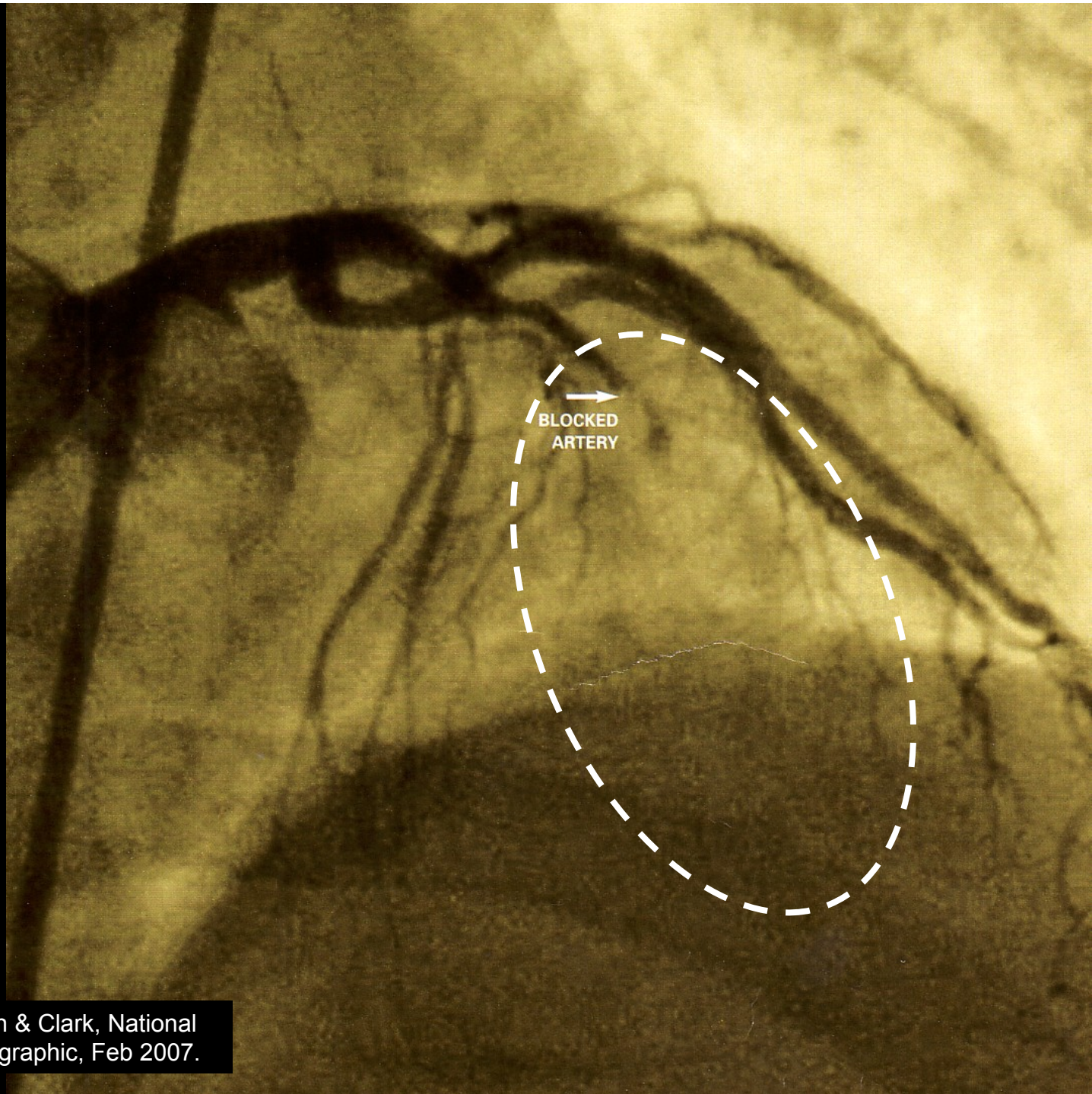


FIGURE 37-1 Devices for percutaneous transluminal coronary interventions. **A**, Coronary balloon. **B**, Rotational atherectomy burr (Rotablator). **C**, Coronary stent.

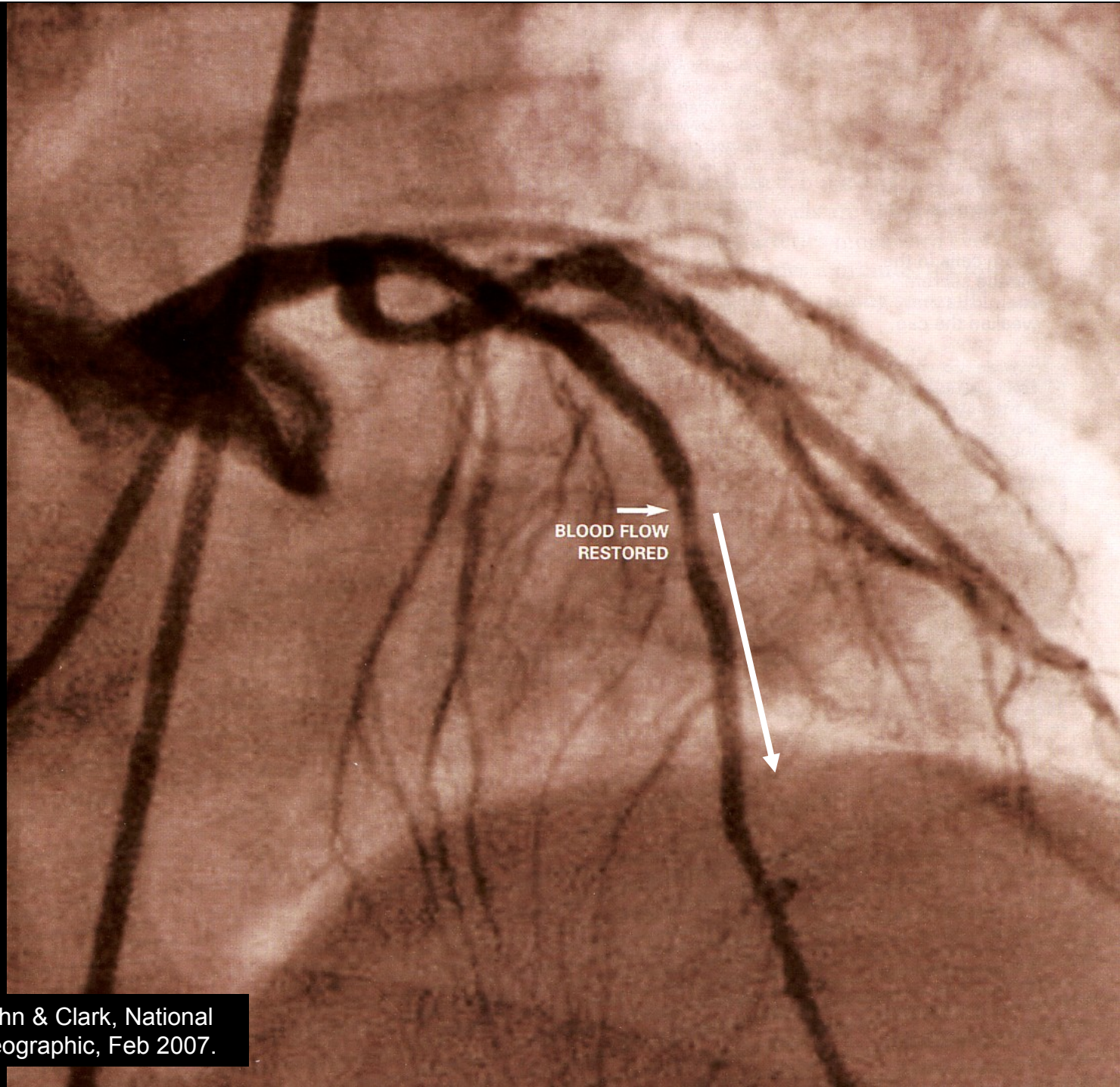
SOURCE: Willerson & Cohn, 2000.



SOURCE: Pollock & Wilmore, 1990.

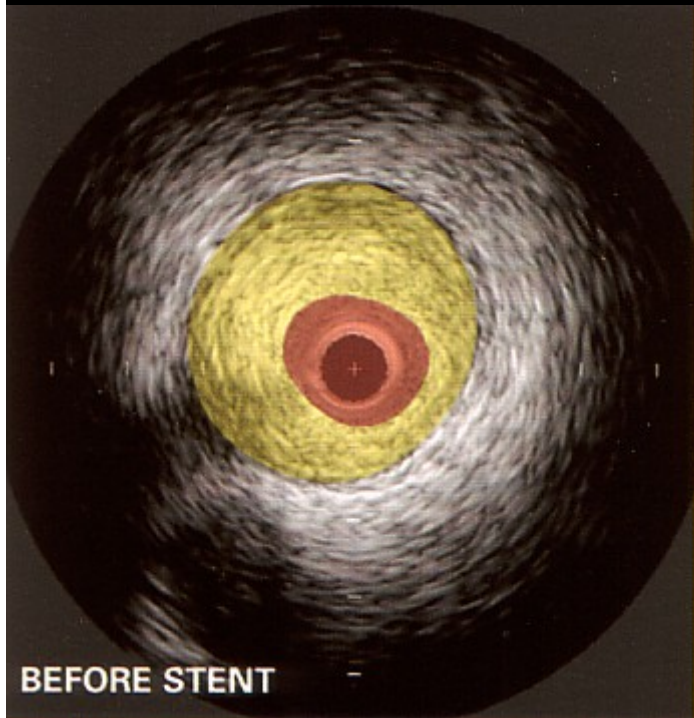


SOURCE: Kahn & Clark, National Geographic, Feb 2007.

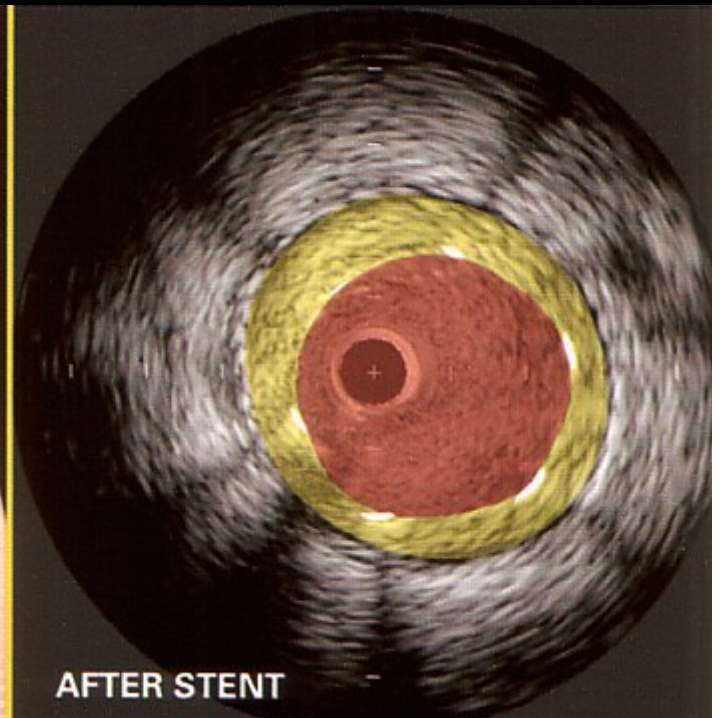


→
BLOOD FLOW
RESTORED

SOURCE: Kahn & Clark, National Geographic, Feb 2007.



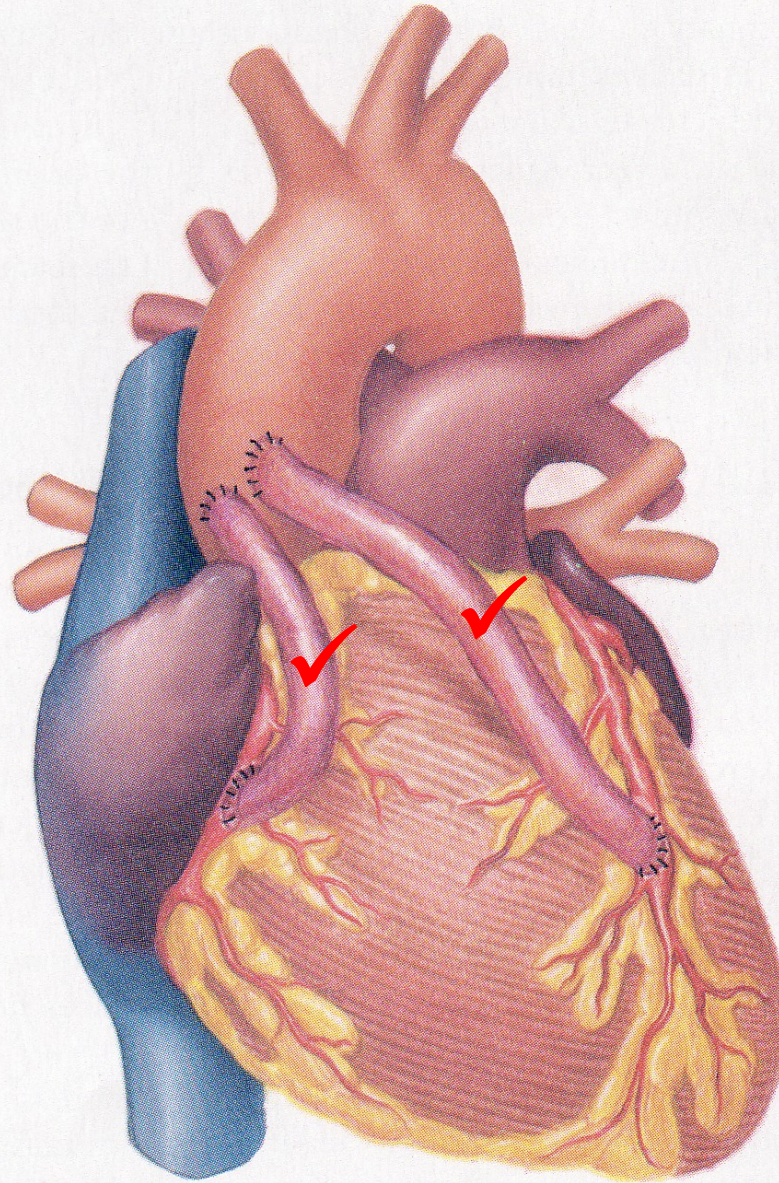
BEFORE STENT



AFTER STENT

SOURCE: Kahn & Clark, National Geographic, Feb 2007.

CABG = Coronary Artery Bypass Graft

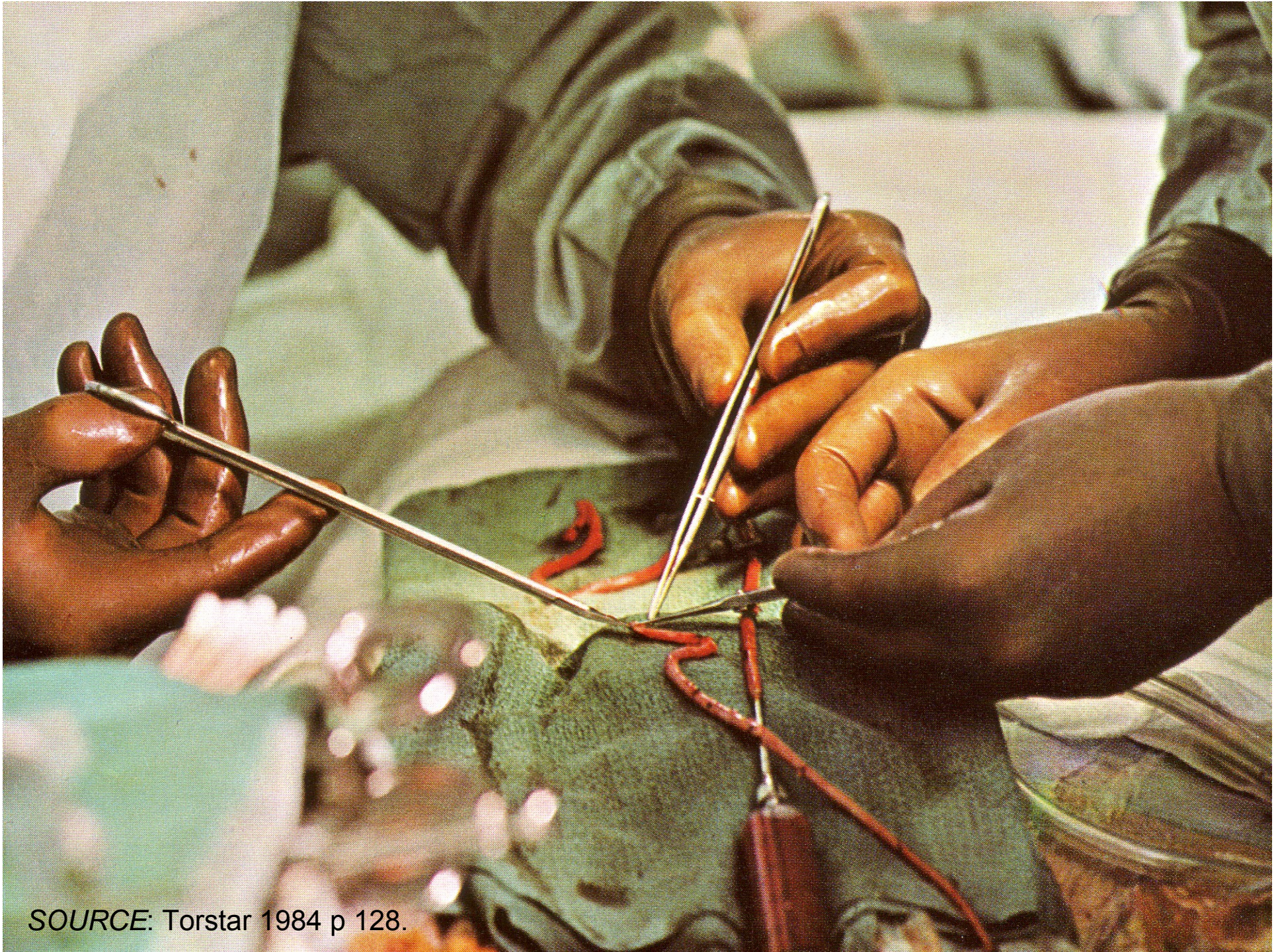


Double?

Triple?

Quadruple?

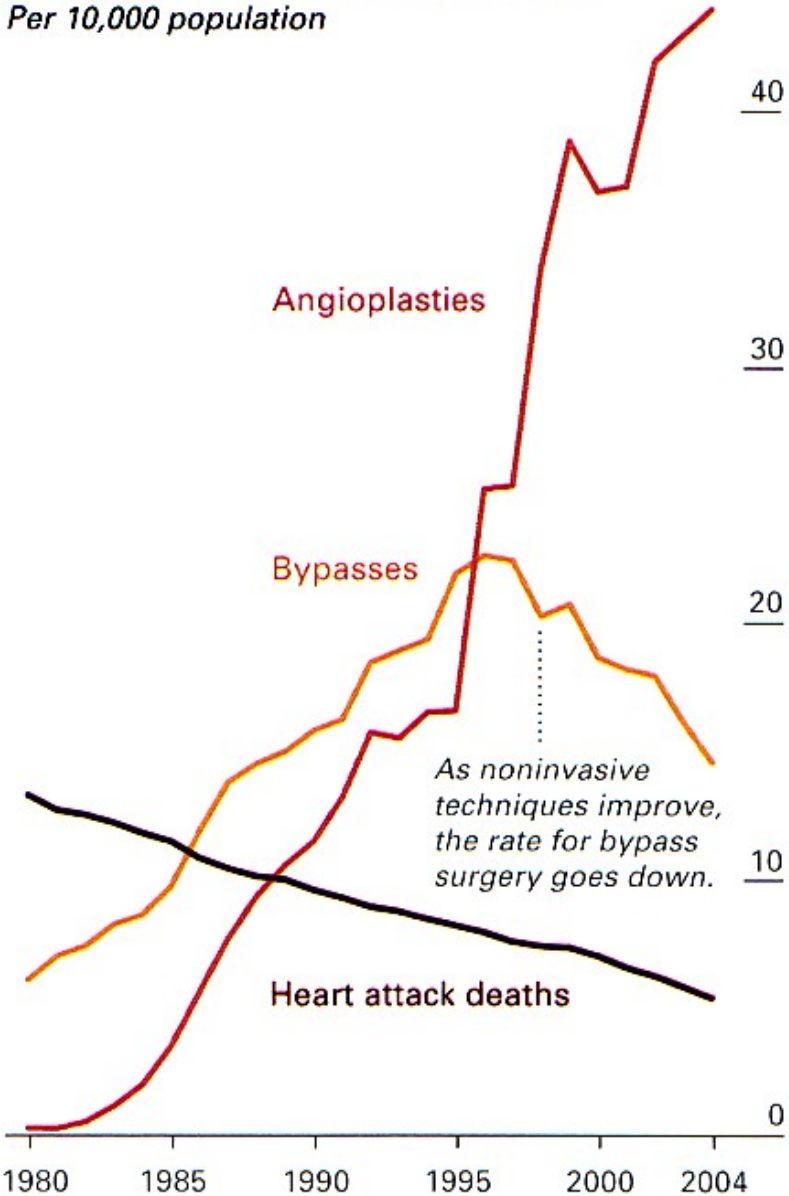
Quintuple?



SOURCE: Torstar 1984 p 128.

Procedures and heart attack deaths

Per 10,000 population

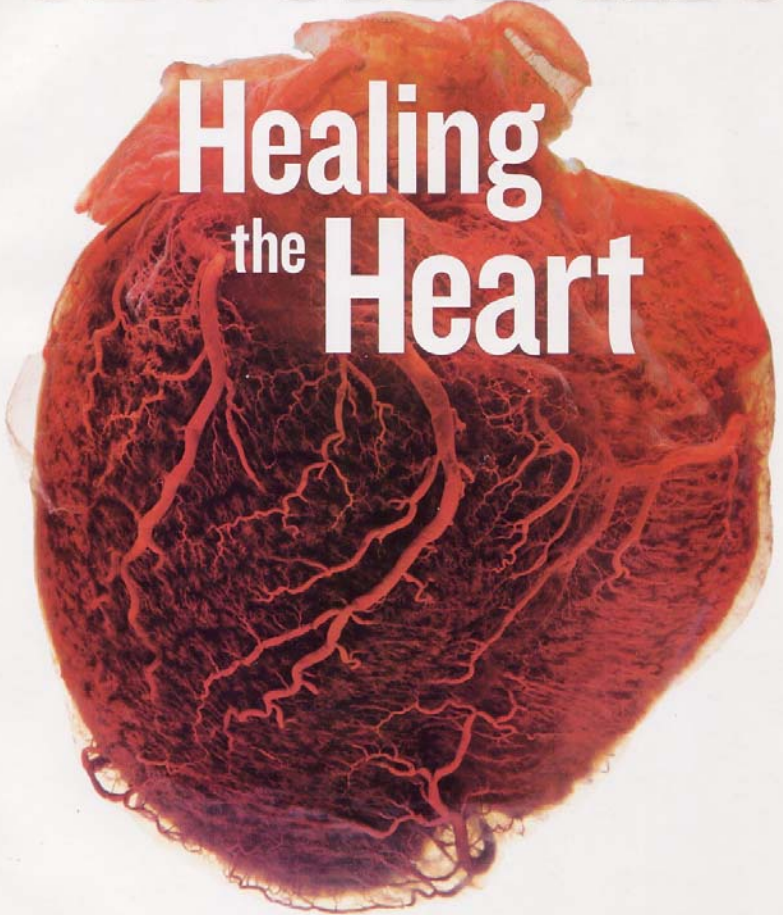


SOURCES: THOMAS THOM, NATIONAL HEART, LUNG, AND BLOOD INSTITUTE; GAUTAM GOWRISANKARAN, WASHINGTON UNIVERSITY IN ST. LOUIS; SALIM YUSUF, McMASTER UNIVERSITY, THE INTERHEART STUDY

NATIONALGEOGRAPHIC.COM/MAGAZINE

FEBRUARY 2007

NATIONAL GEOGRAPHIC



Healing the Heart

Beauty on the Border 66 Curse of Nigerian Oil 86
Hawaii's Unearthly Worms 118 Forests of the Tide 132



**CardioWest artificial heart = \$106,000!
3000 await transplants, but only 2100
donors are available...**

<http://ngm.nationalgeographic.com/2007/02/hearts/hearts-text.html>



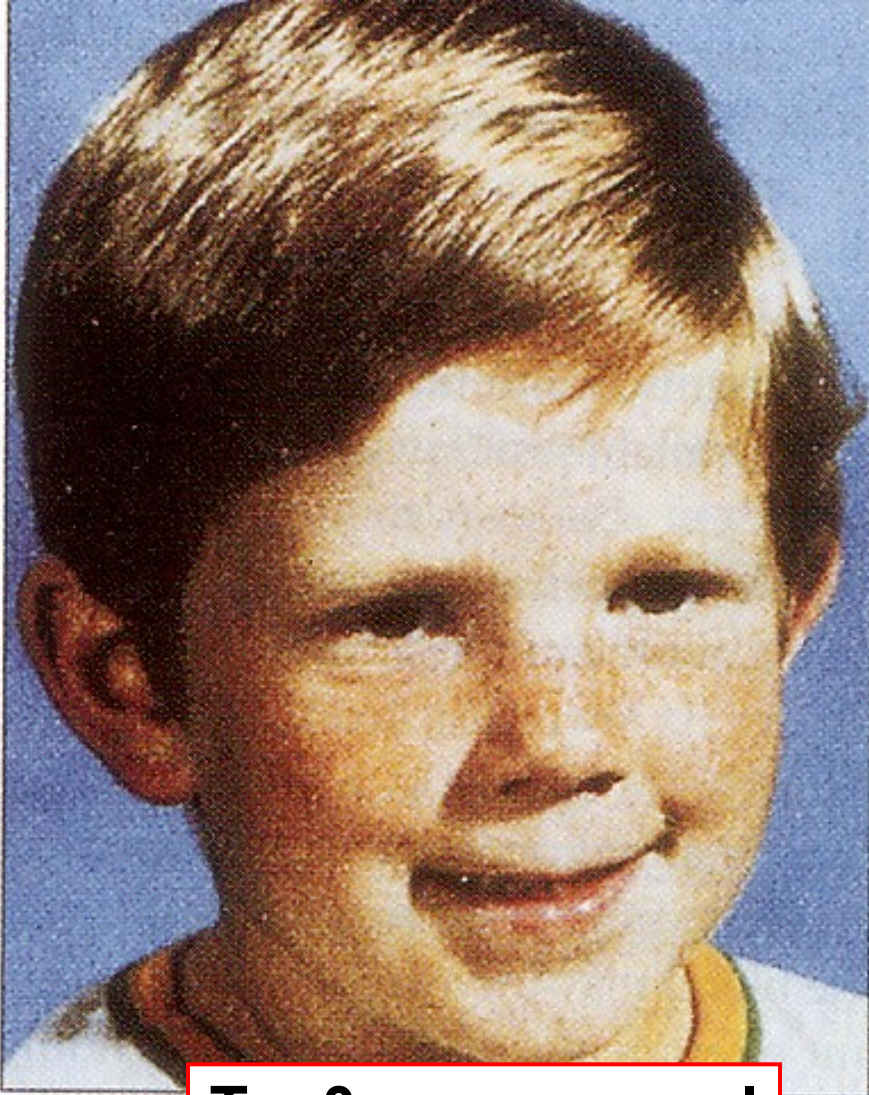
Photograph by Robert Clark

Discussion

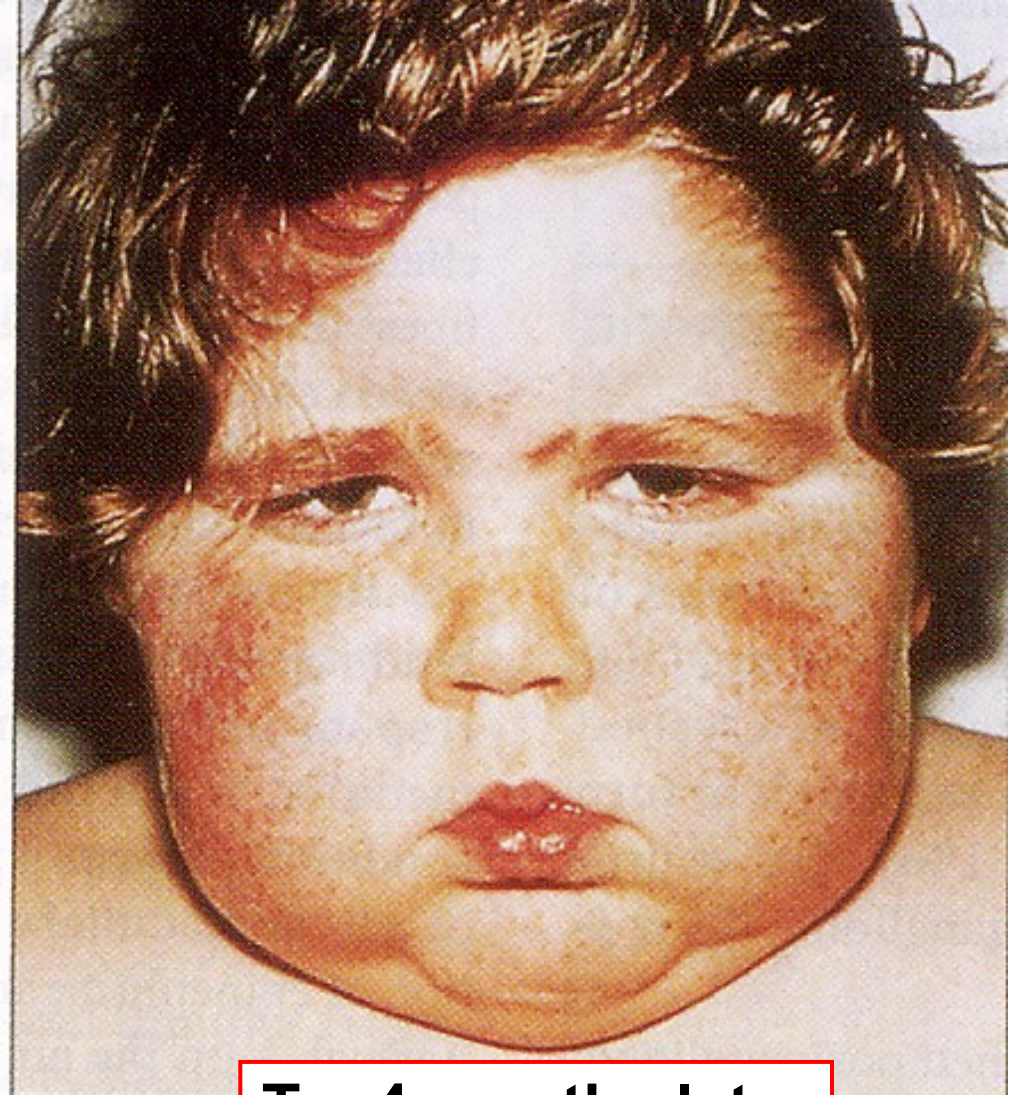
Comments

Q?

***Cushing's Syndrome = Hypersecretion
of Cortisol: Hypothalamic (CRH),
Pituitary (ACTH), or Adrenal (Cortisol)***



T = 0, near normal



T = 4 months later

Endocrine/Hormone?

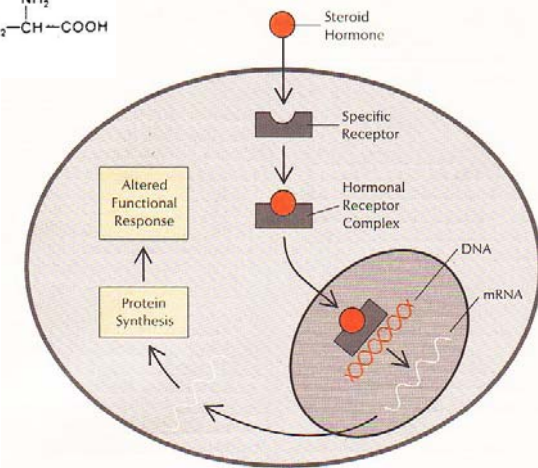
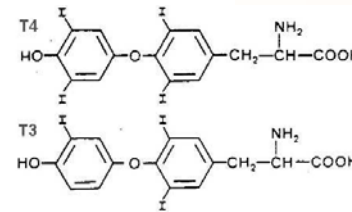
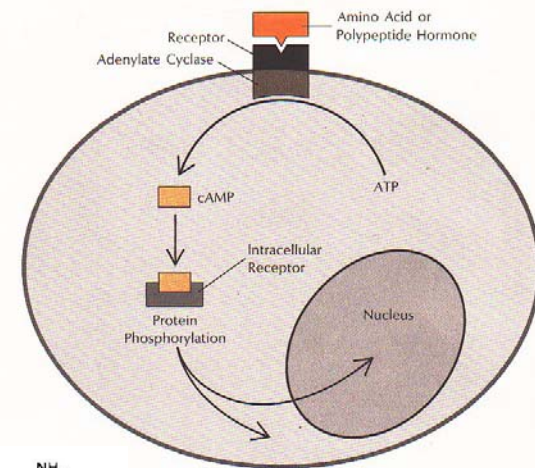
- ① *Made by gland?*
- ② *Secreted into blood?*
- ③ *Acts on target?*

Hormone/Endocrine Classifications

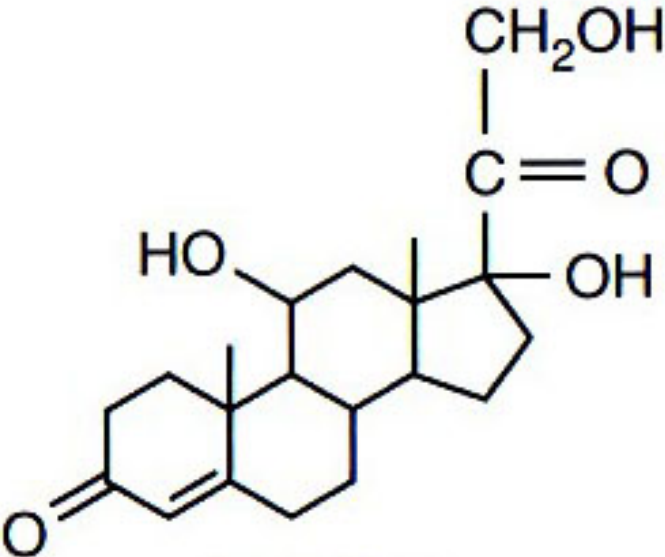
Exogenous



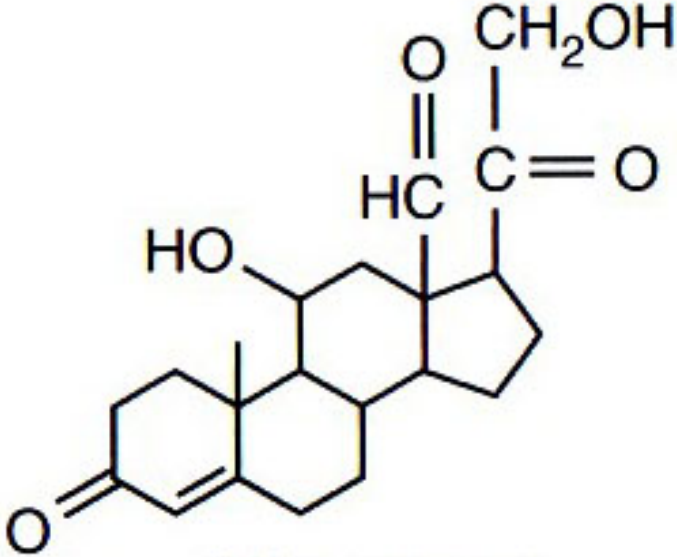
Endogenous



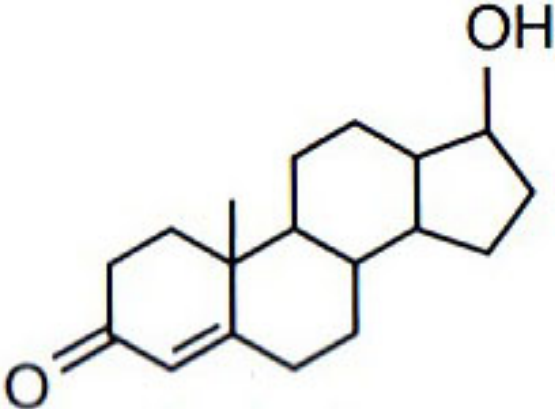
Steroid Hormone Structure: Cholesterol Backbone



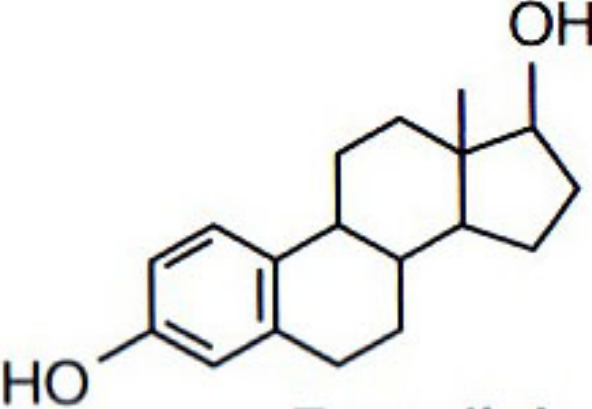
Cortisol



Aldosterone



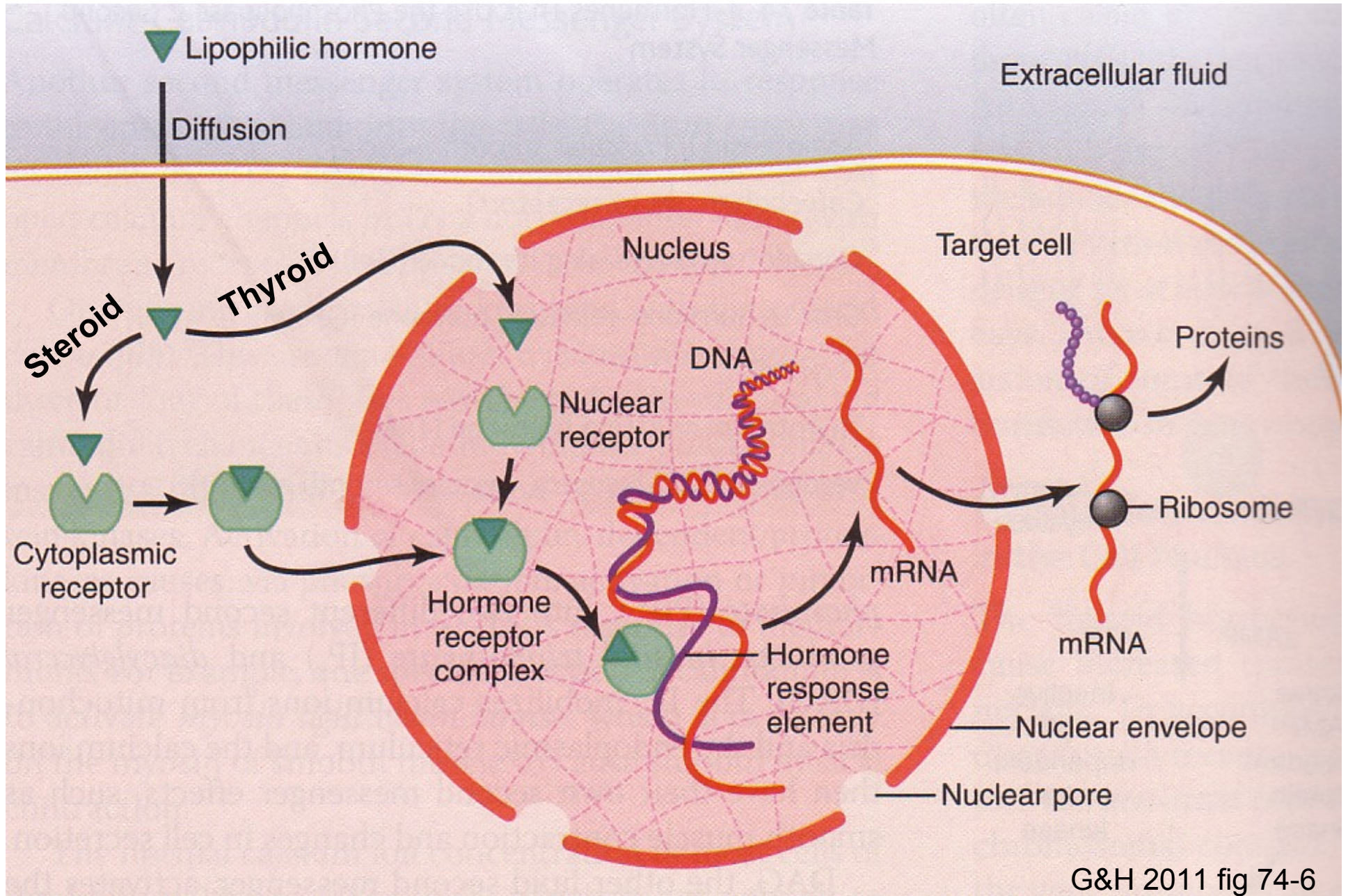
Testosterone



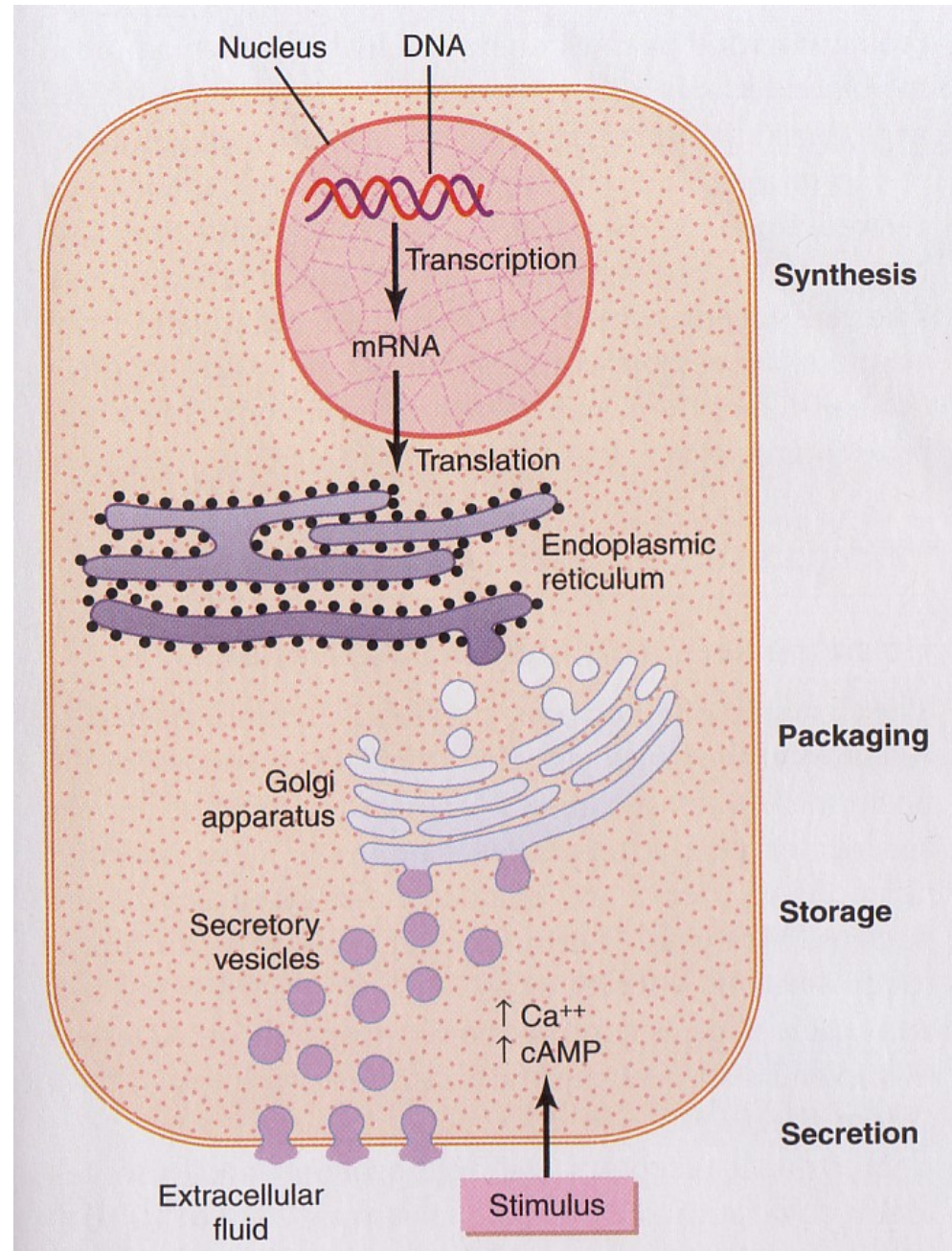
Estradiol

G&H 2011 fig 74-3; cf: fig 77-2

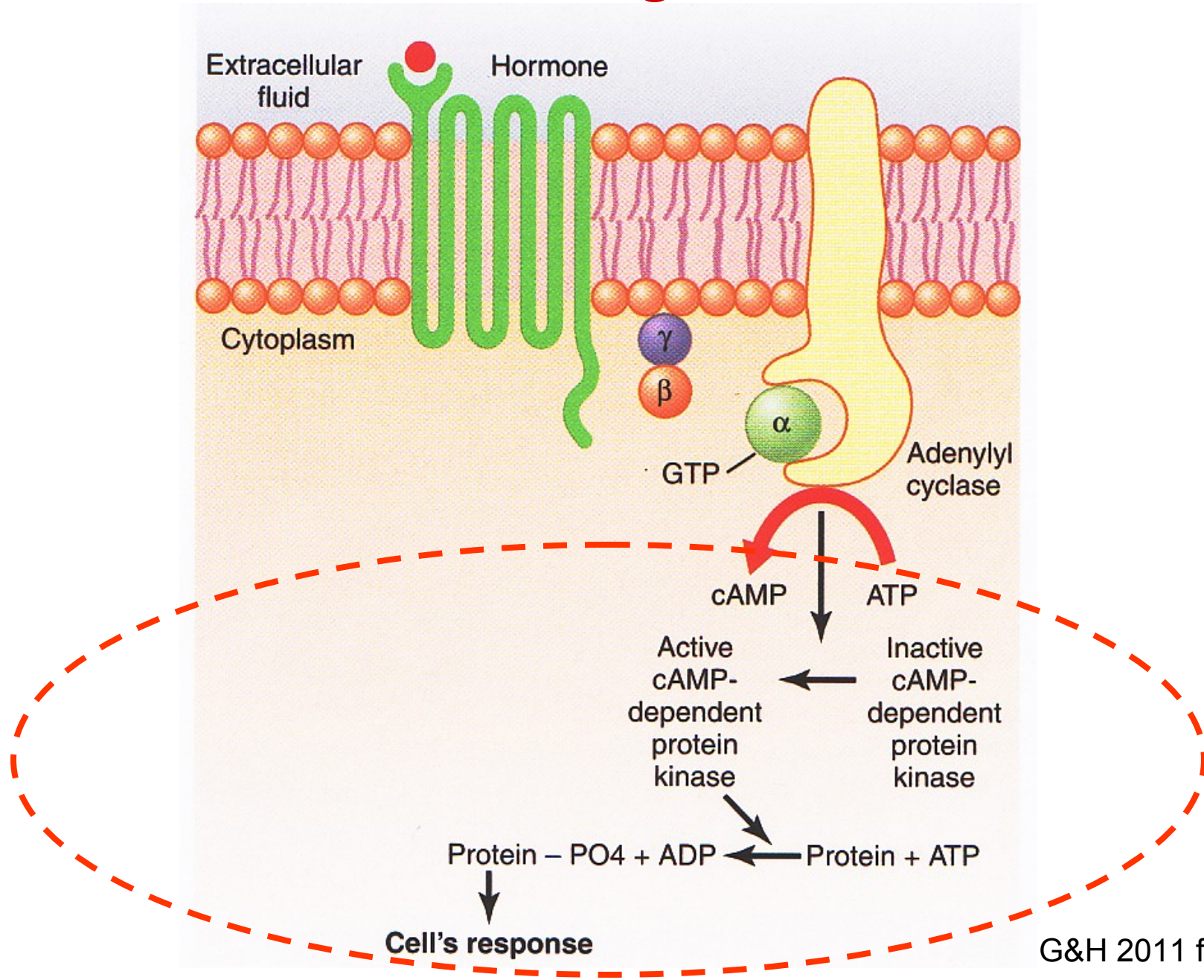
Lipophilic (Steroid+Thyroid) Hormone Mechanisms



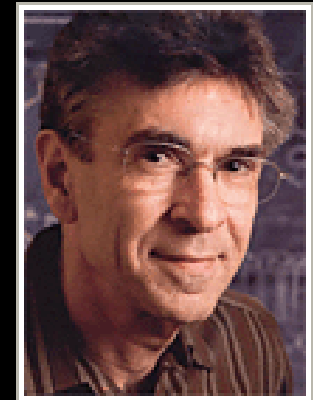
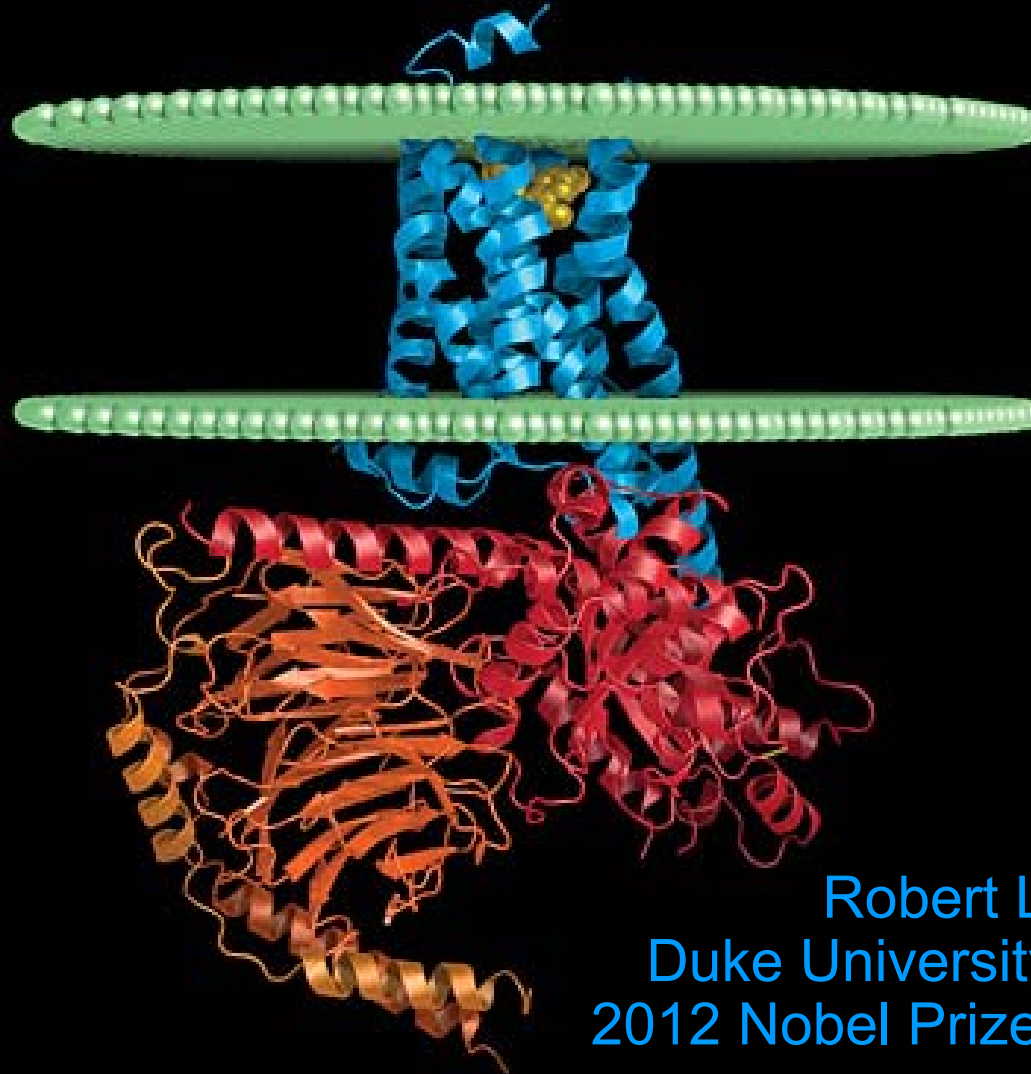
Peptide Hormone Synthesis & Secretion



cAMP 2nd Messenger Mechanism



G-Protein Coupled Receptor (blue) sits within lipid bilayer (green) to respond to hormone (yellow)



Robert Lefkowitz, MD
Duke University Cardiologist
2012 Nobel Prize in Chemistry

<http://www.hhmi.org/bulletin/winter2013/features/index.html>

Image by Wayne Decatur

Table 74-2 Hormones That Use the Adenylyl Cyclase – Cyclic AMP Second Messenger System

Adrenocorticotrophic hormone (ACTH)
Angiotensin II (ANG II, epithelial cells)
Calcitonin
Catecholamines (β receptors)
Corticotropin-releasing hormone (CRH)
Follicle-stimulating hormone (FSH)
Glucagon
Human chorionic gonadotropin (hCG)
Luteinizing hormone (LH)
Parathyroid hormone (PTH)
Secretin
Somatostatin (SS, GH RIH)
Thyroid-stimulating hormone (TSH)
Vasopressin (ADH, VP, V_2 receptor, epithelial cells)

Phospholipase C 2nd Messenger Mechanism

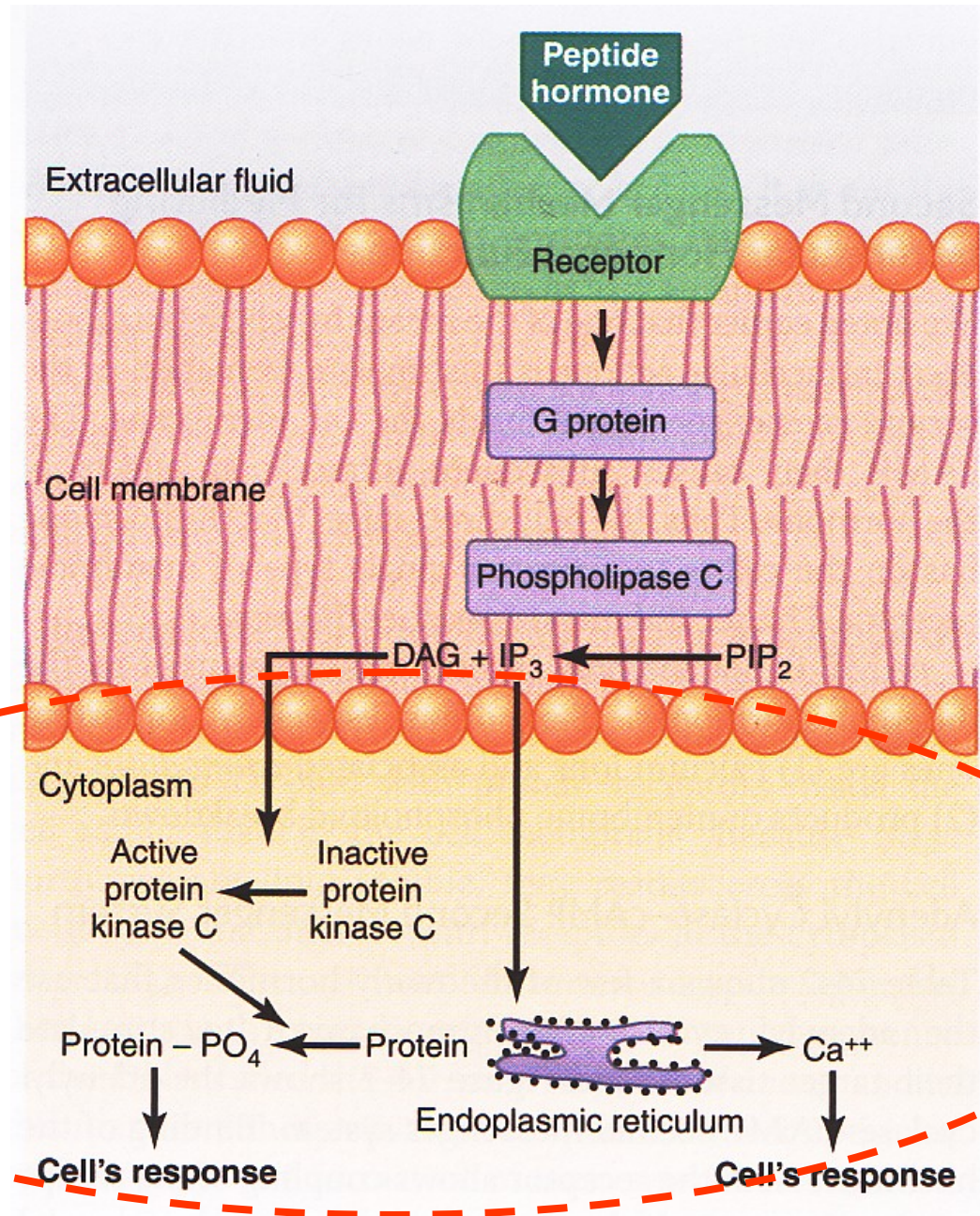


Table 74-3 Hormones That Use the Phospholipase C Second Messenger System

Angiotensin II (ANG II, vascular smooth muscle)

Catecholamines (α receptors)

Gonatotropin-releasing hormone (GnRH)

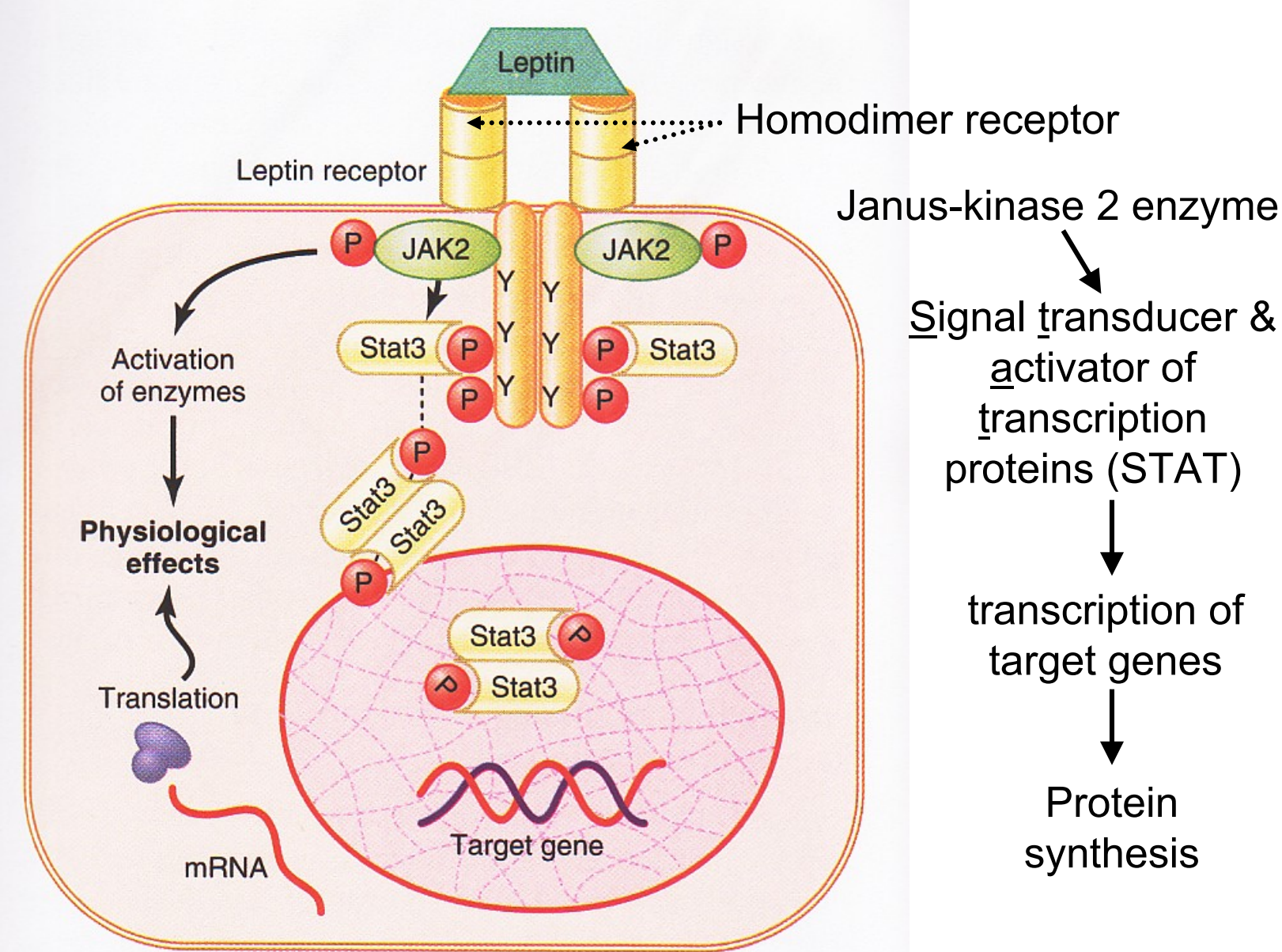
Growth-hormone-releasing hormone (GHRH)

Oxytocin (OXY, hypothalamus production, posterior pituitary storage)

Thyrotropin releasing hormone TRH)

Vasopressin (ADH, VP, V_1 receptor, vascular smooth muscle)

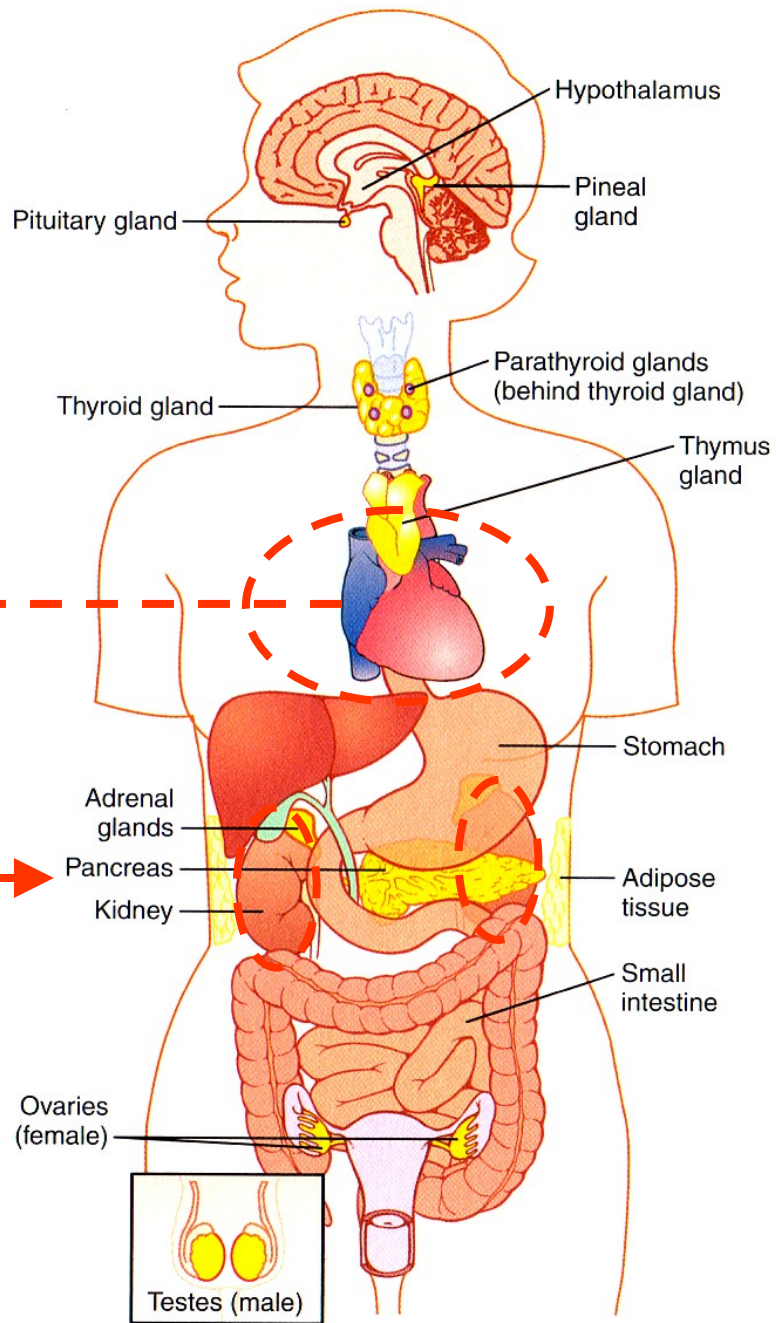
Leptin: Enzyme-Linked Hormone Receptor



<http://www.ncbi.nlm.nih.gov/pubmed/22249808>

G&H 2011 fig 74-5

**ANP =
Atrial
Natriuretic
Polypeptide**



2 Vasodilation

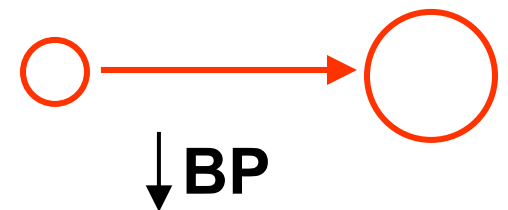
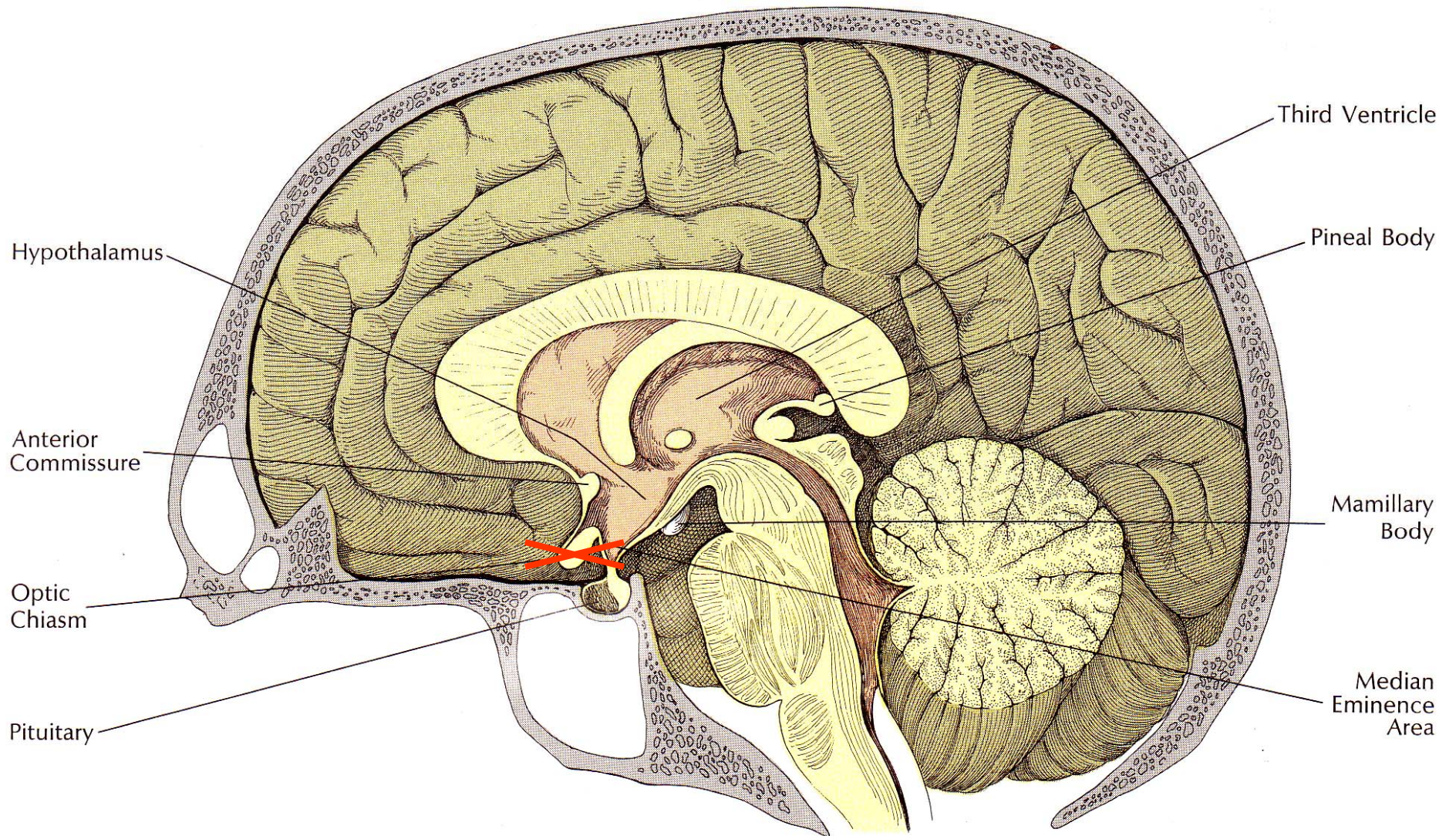
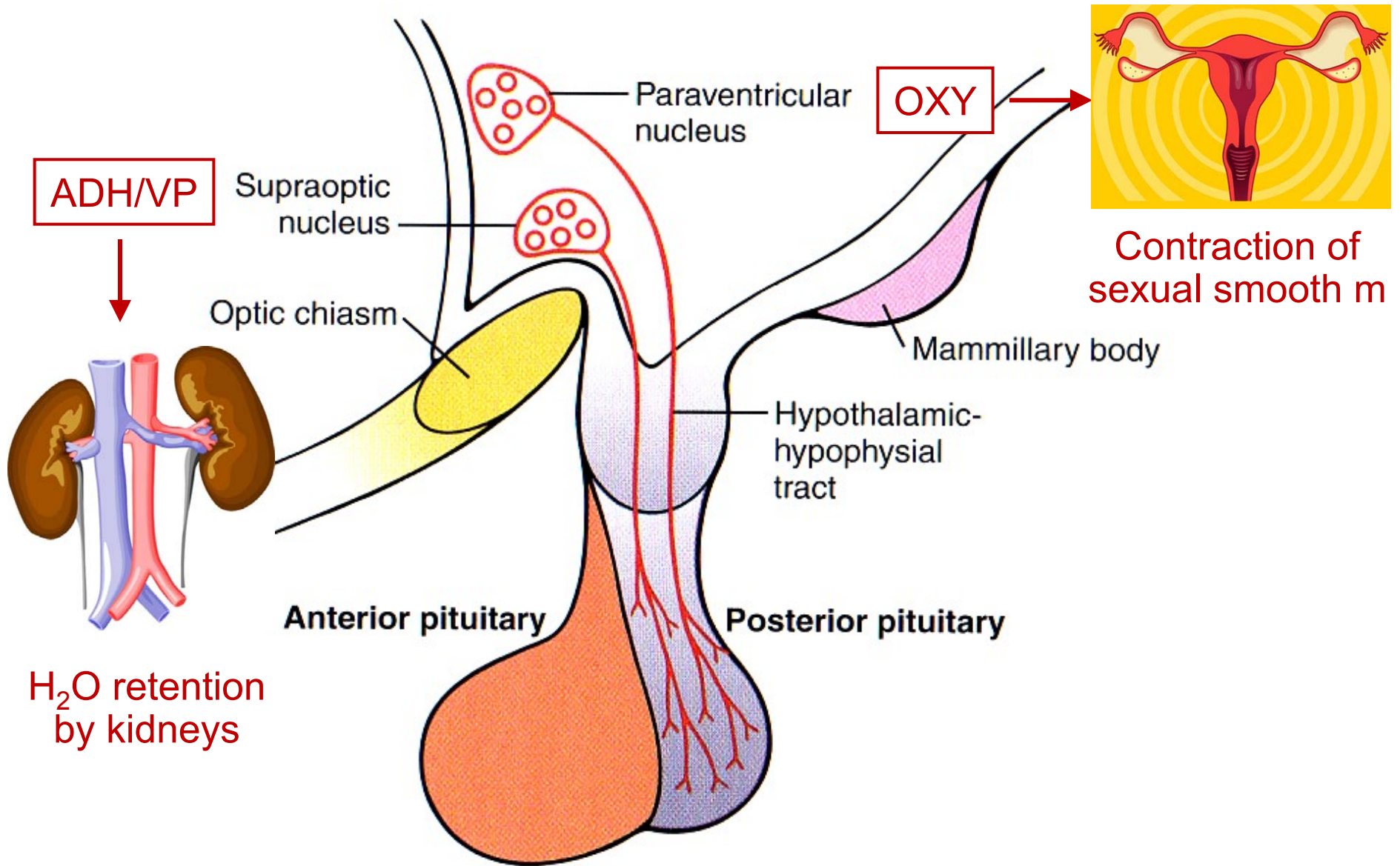


Figure 74-1 Anatomical loci of the principal endocrine glands and tissues of the body.

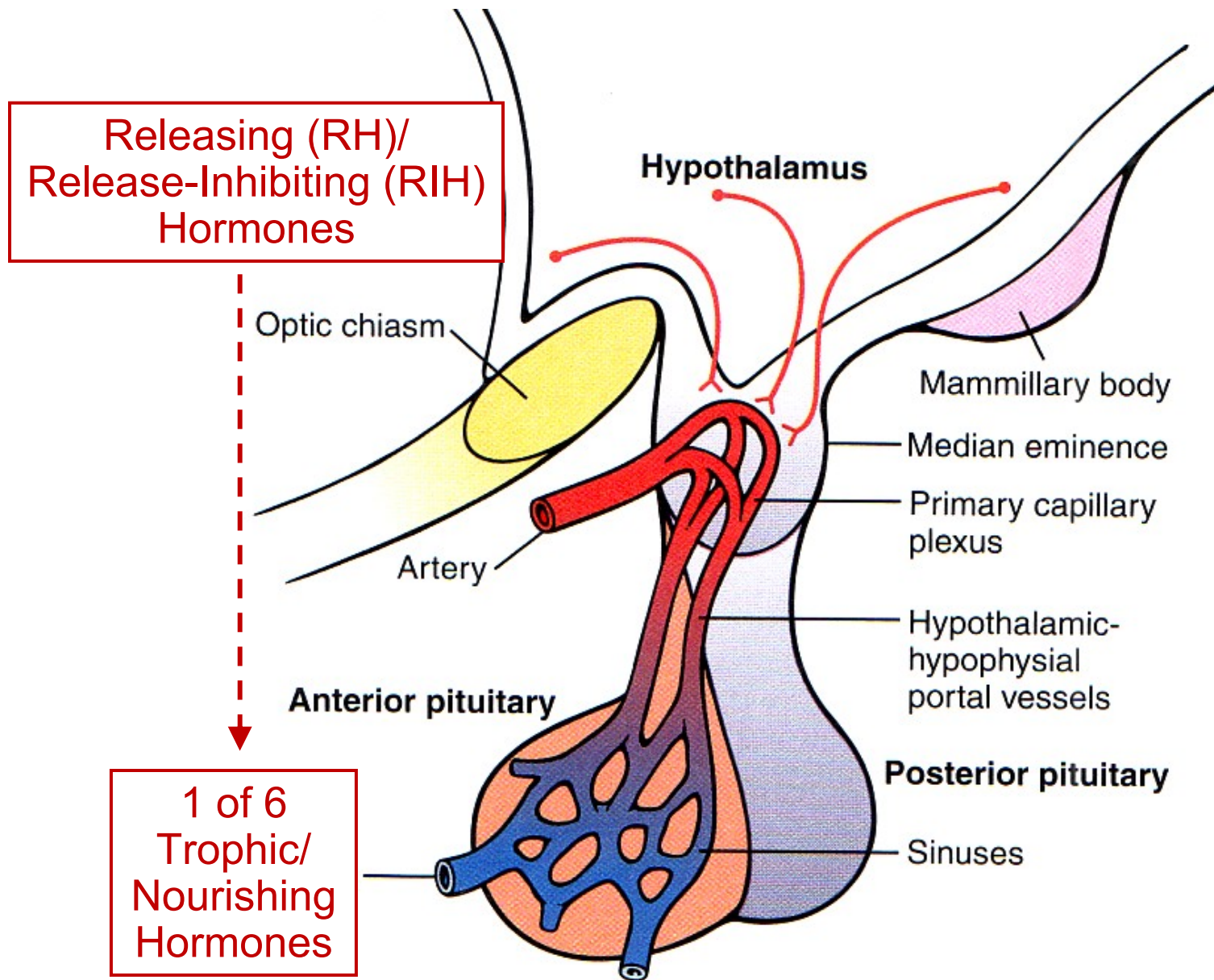
Lateral View Showing Relationship of the Pituitary Gland to the Hypothalamus



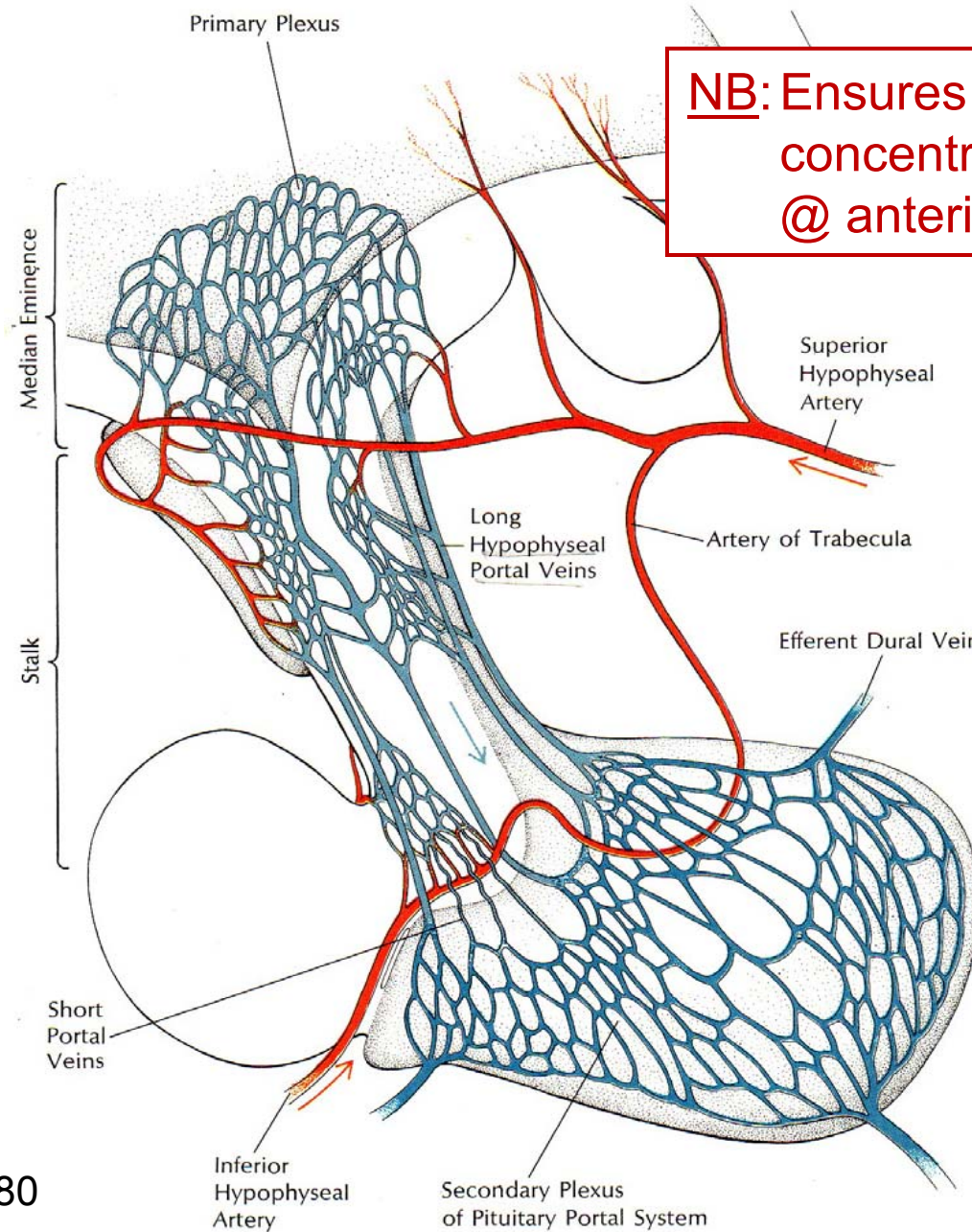
Hypothalamus – Posterior Pituitary Nervous Connection



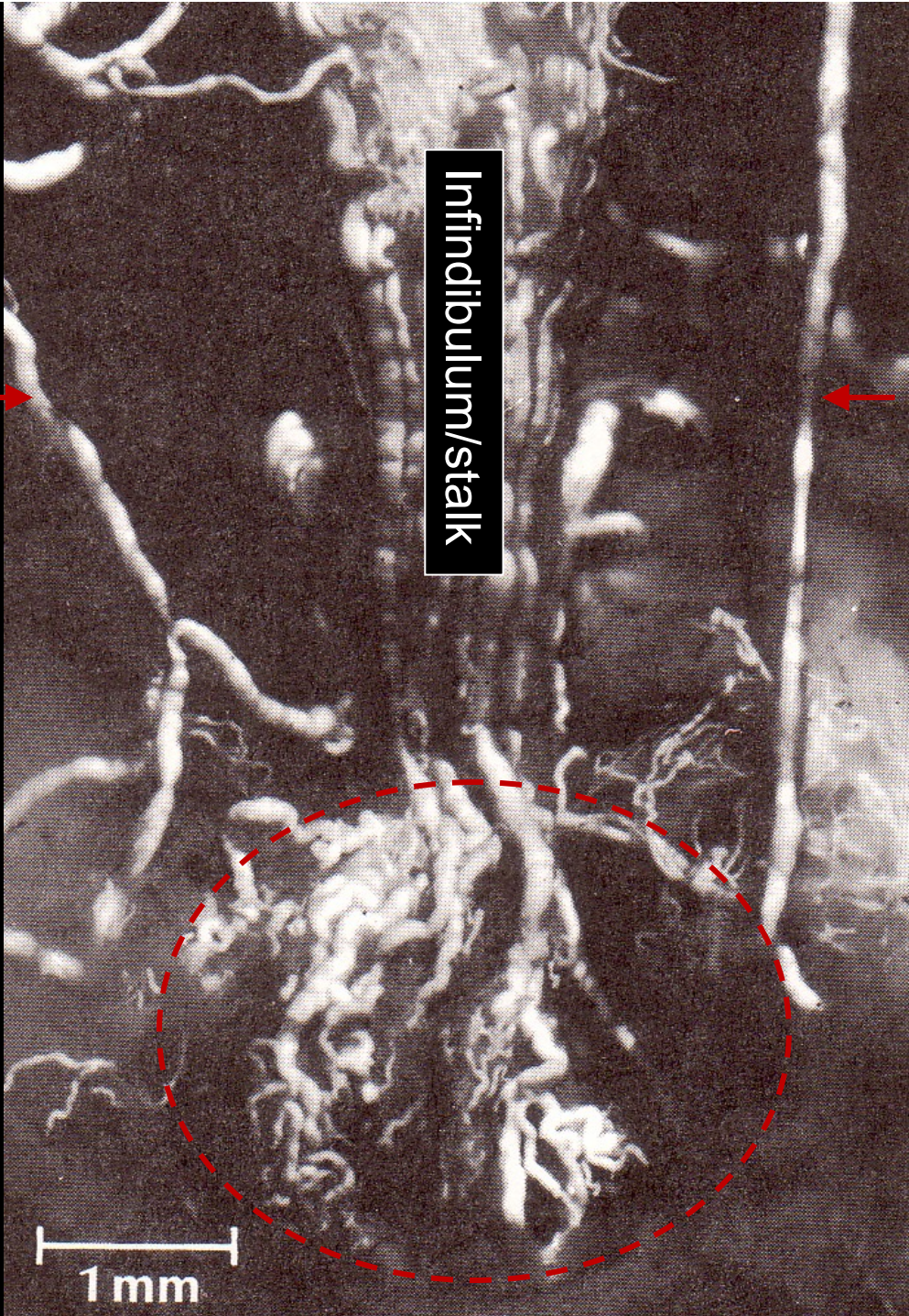
Hypothalamus – Anterior Pituitary Vascular Connection



Capillary-Venule-Capillary Circulation



NB: Ensures RH/RIH super-concentrated upon arrival @ anterior pituitary!



Infundibulum/stalk

Long hypophyseal-portal veins

Pituitary removed!

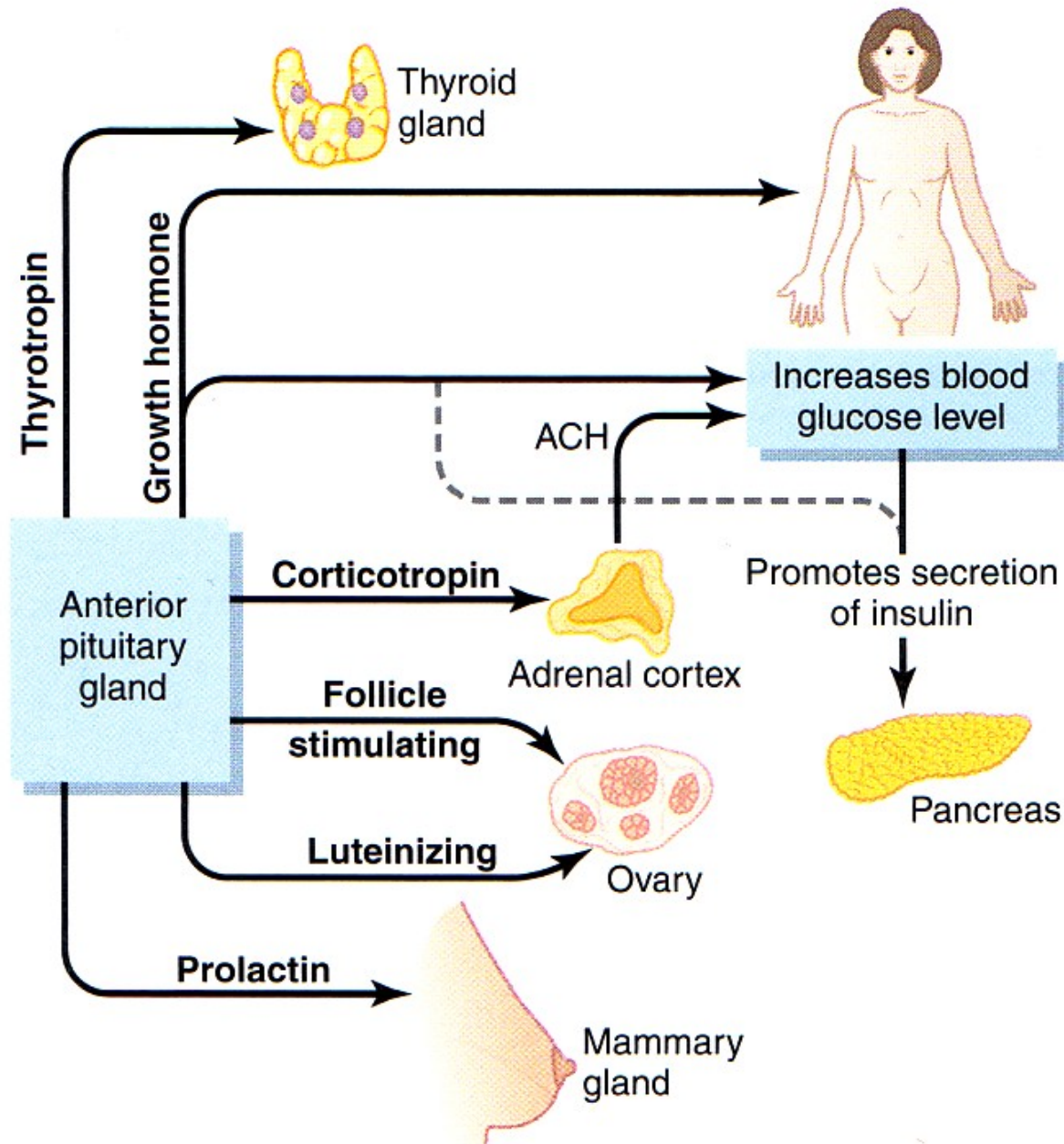
Krieger & Hughes
1980

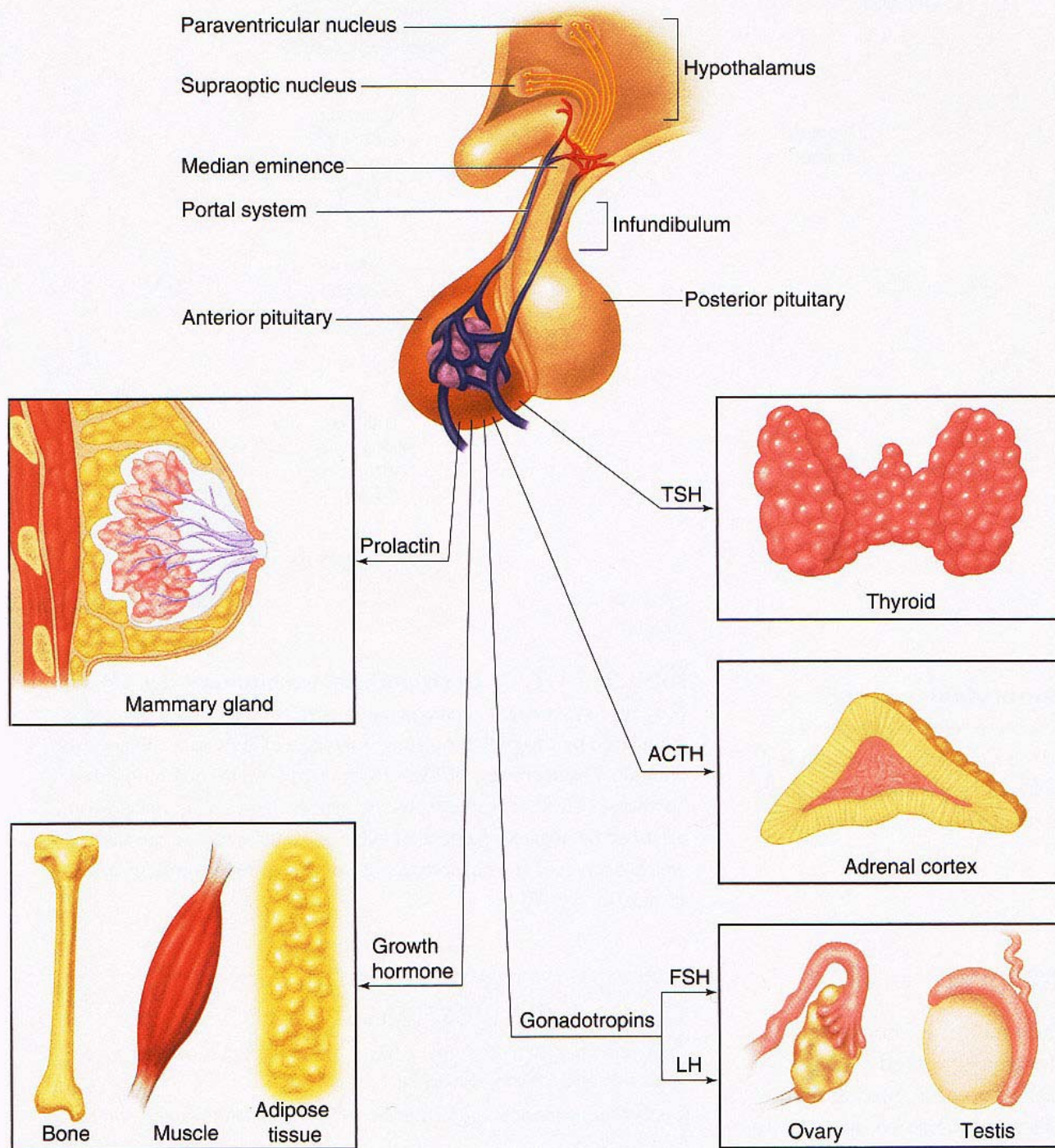
1 mm

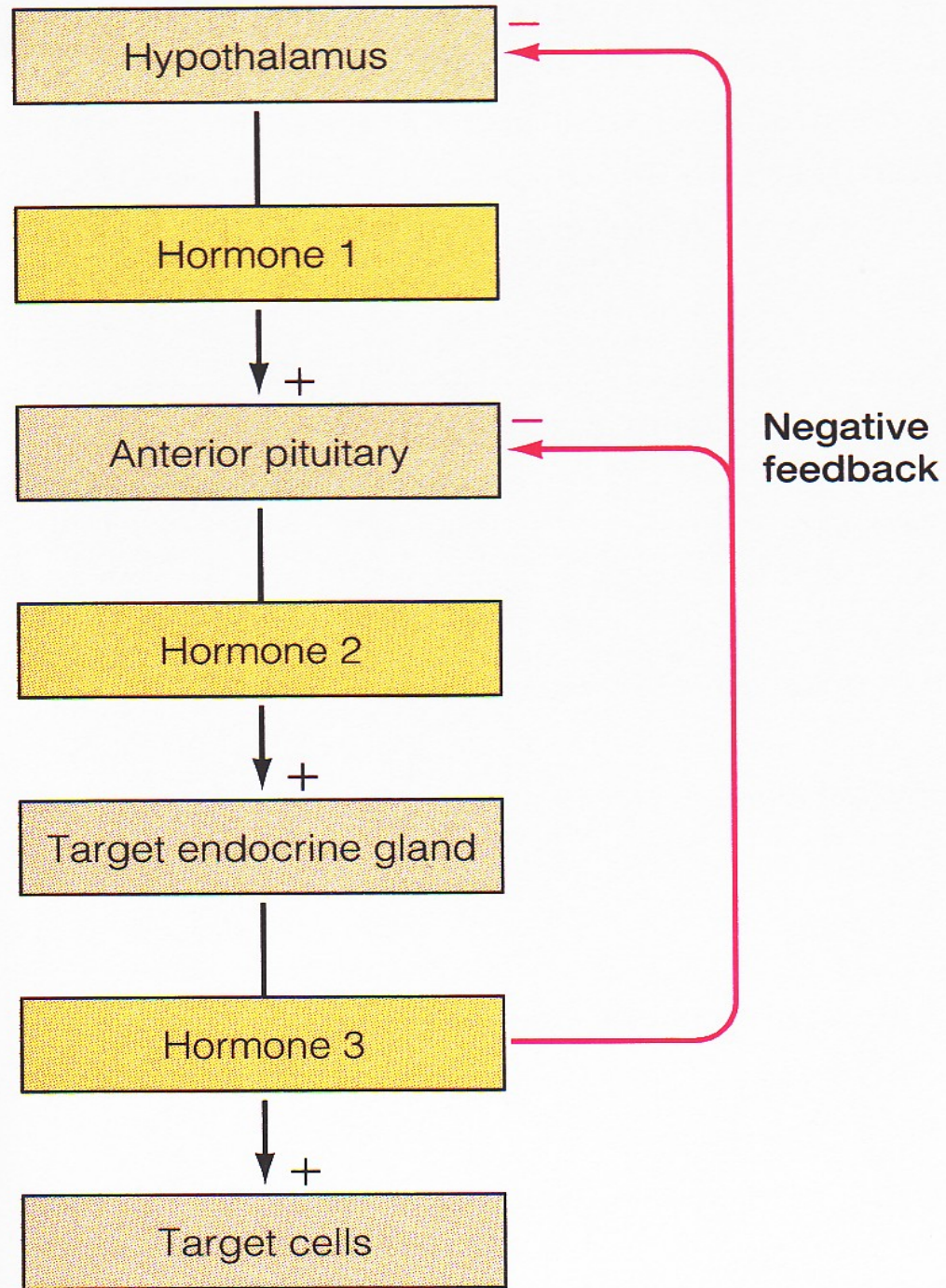
Table 74-1 Endocrine Glands, Hormones, and Their Functions and Structure

Gland/Tissue	Hormones	Major Functions	Chemical Structure
Hypothalamus (Chapter 75)	Thyrotropin-releasing hormone (TRH)	Stimulates secretion of thyroid-stimulating hormone (TSH) and prolactin	Peptide
	Corticotropin-releasing hormone (CRH)	Causes release of adrenocorticotrophic hormone (ACTH)	Peptide
	Growth hormone–releasing hormone (GHRH)	Causes release of growth hormone	Peptide
	Growth hormone inhibitory hormone (GHIH) (somatostatin)	Inhibits release of growth hormone	Peptide
	Gonadotropin-releasing hormone (GnRH)	Causes release of luteinizing hormone (LH) and follicle-stimulating hormone (FSH)	
	Dopamine or prolactin-inhibiting factor (PIF)	Inhibits release of prolactin	Amine
Anterior pituitary (Chapter 75)	Growth hormone	Stimulates protein synthesis and overall growth of most cells and tissues	Peptide
	TSH	Stimulates synthesis and secretion of thyroid hormones (thyroxine and triiodothyronine)	Peptide
	ACTH	Stimulates synthesis and secretion of adrenocortical hormones (cortisol, androgens, and aldosterone)	Peptide
	Prolactin	Promotes development of the female breasts and secretion of milk	Peptide
	FSH	Causes growth of follicles in the ovaries and sperm maturation in Sertoli cells of testes	Peptide
	LH	Stimulates testosterone synthesis in Leydig cells of testes; stimulates ovulation, formation of corpus luteum, and estrogen and progesterone synthesis in ovaries	Peptide

Anterior Pituitary Metabolic Functions







GH, a Protein Hormone (191 AA)

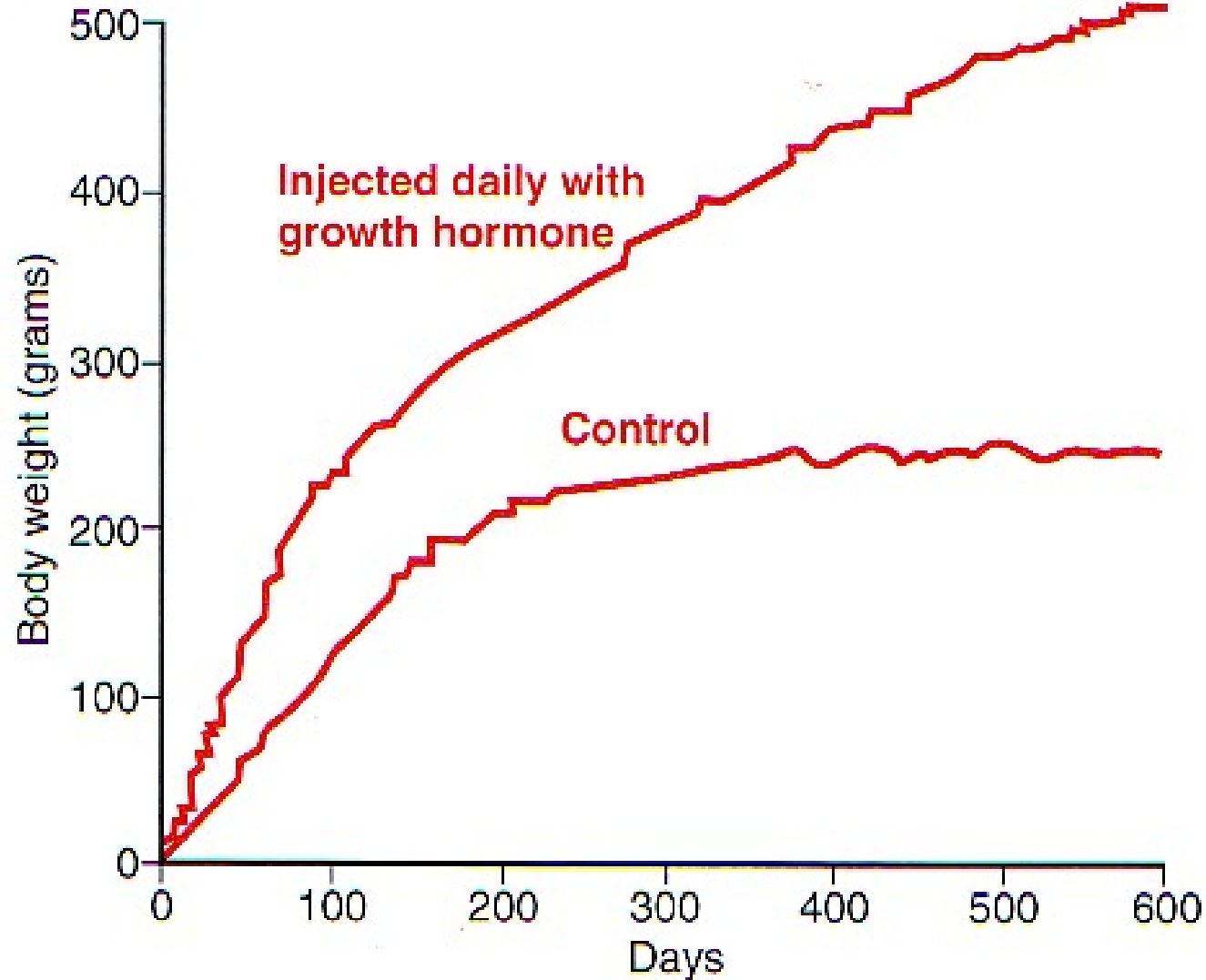


FIGURE 75-5

Comparison of weight gain of a rat injected daily with growth hormone with that of a normal littermate.



LS2 2006

Progression & Development of Acromegaly

Age 13



Age 21



Age 35



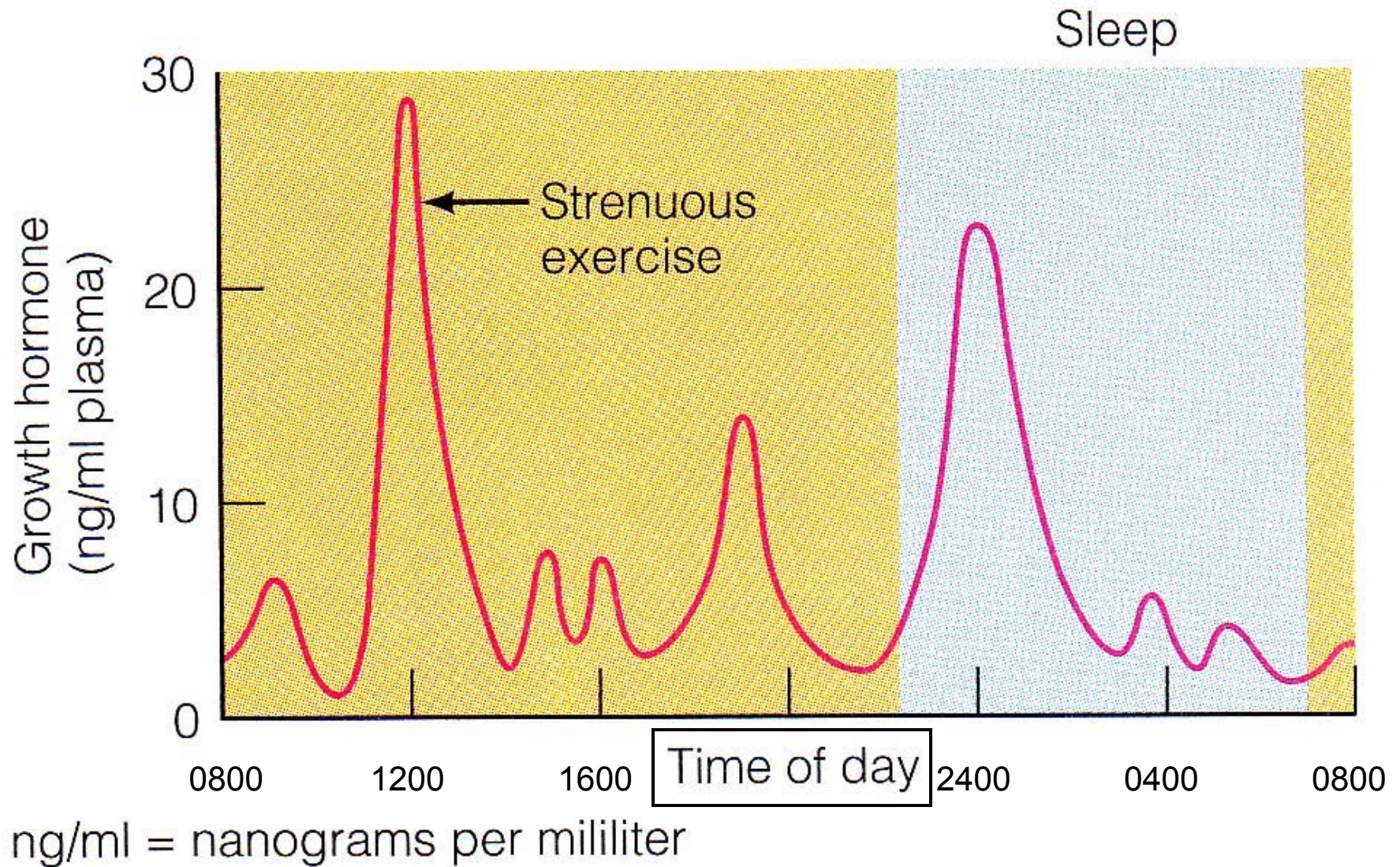


Growth Hormone ≡
Somatotrophic Hormone
Body Builder's Dream?

GH/STH Effects: Insulin Resistance/Type II Diabetes?

- ↑ Amino acid uptake & protein synthesis
- ↑ Lipolysis & fatty acid mobilization
- ↓ Glucose uptake
(skeletal muscle & adipocytes)
- ↑ Glucose production
(liver glycogenolysis)
- ↑ Insulin secretion

Increase GH naturally with exercise & sleep!!



cf: G&H 2011 fig 75-6

Proinsulin with C-Connecting Peptide

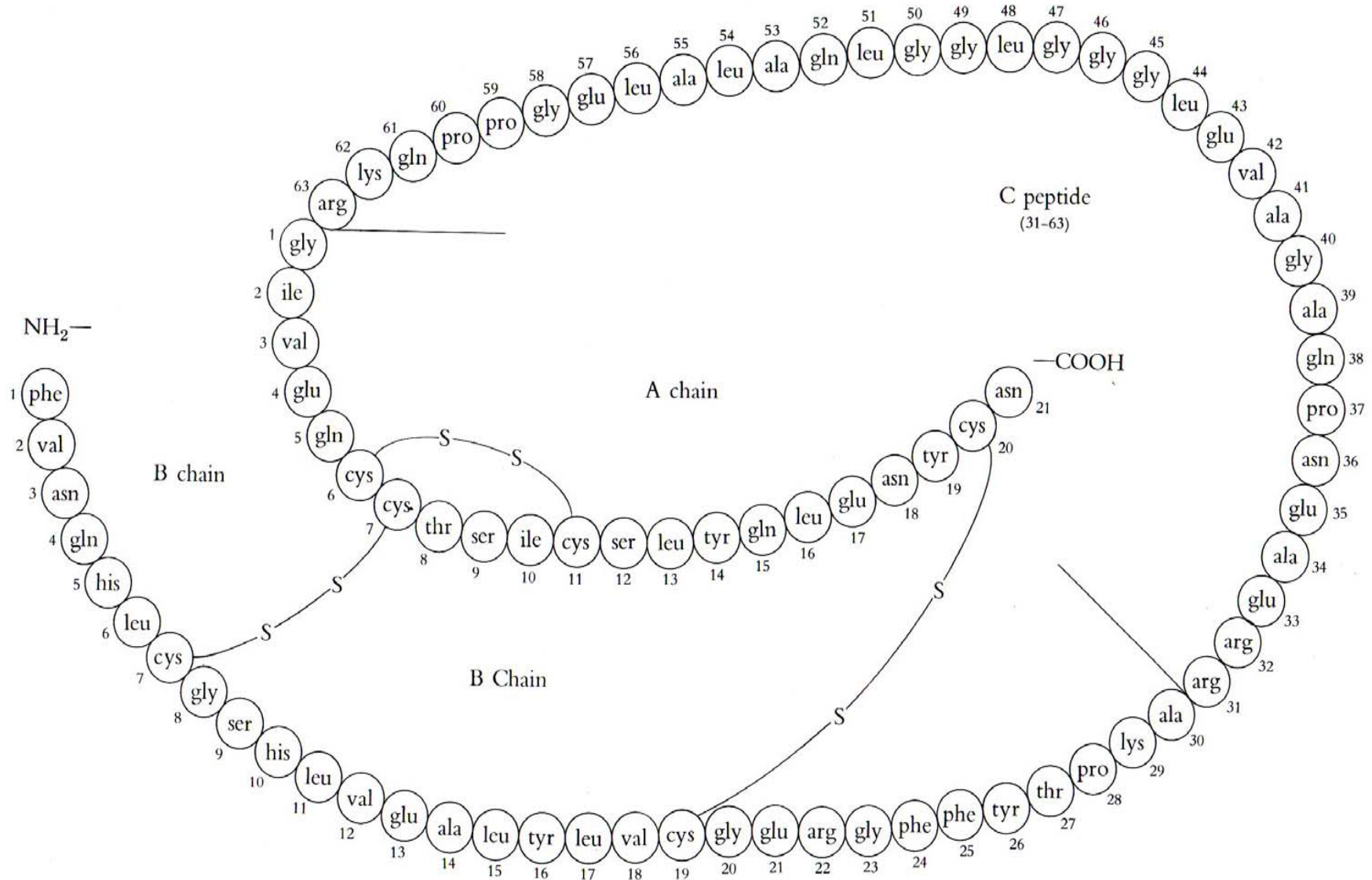
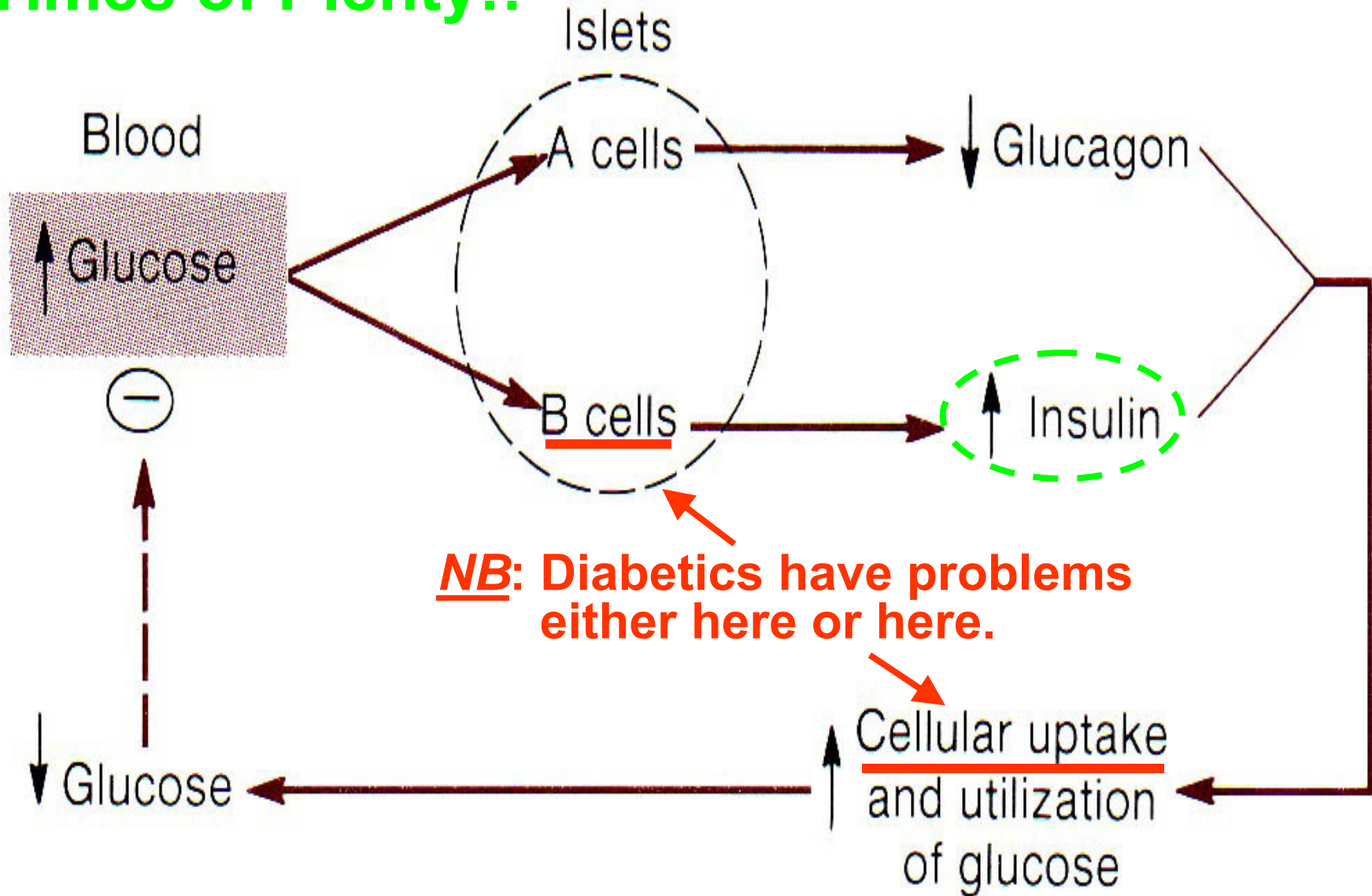


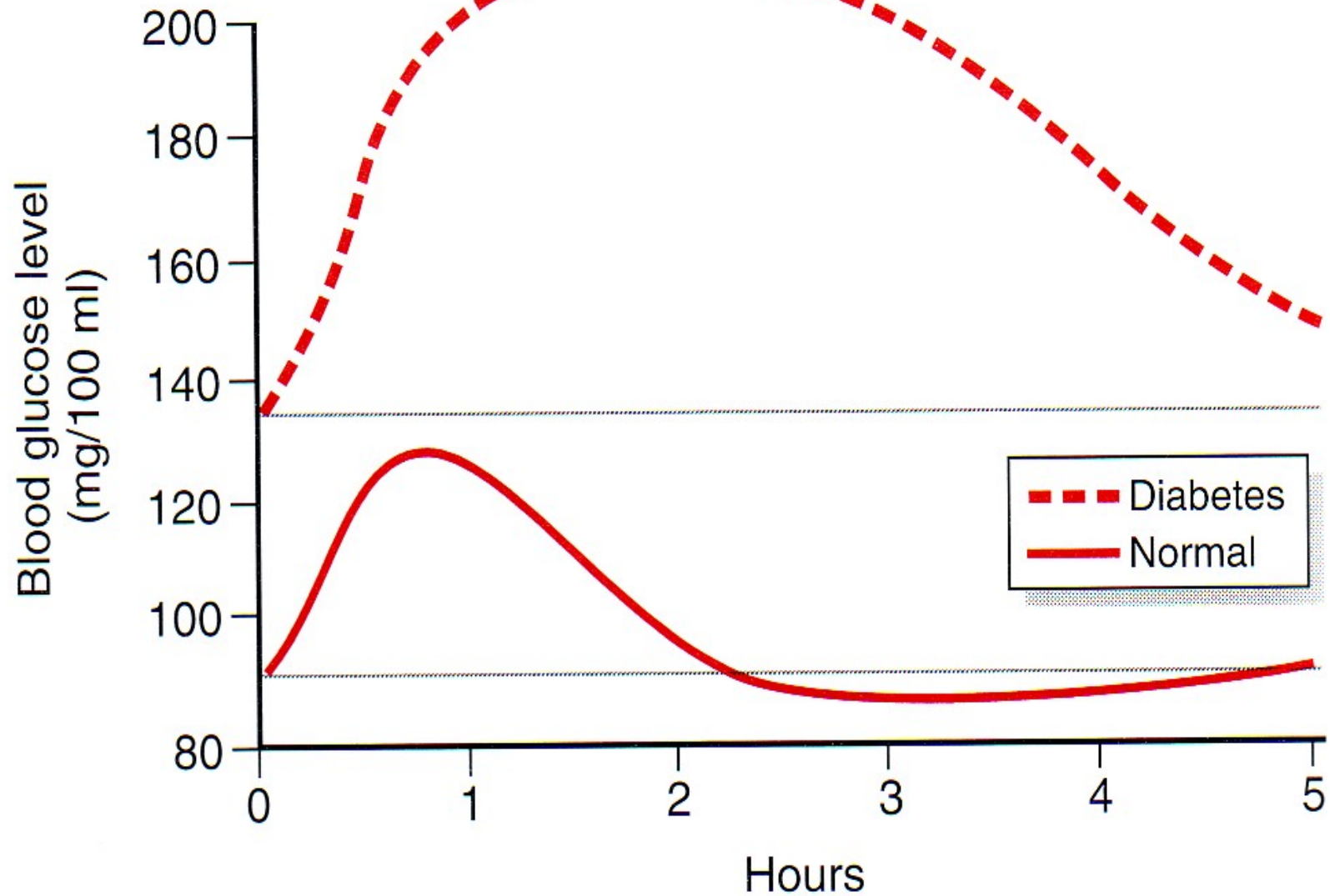
FIG. 10-4. Amino acid sequence of a mammalian proinsulin molecule. Note how the insulin molecule can be formed by cleaving this polypeptide chain at two locations to liberate the C peptide.

Times of Plenty!!



Store!

Diabetic & Normal Response to Glucose Load



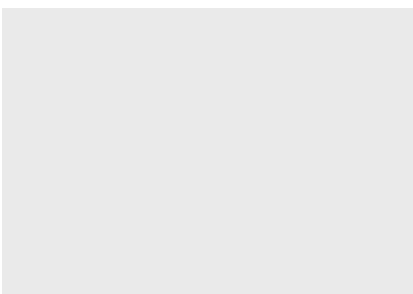
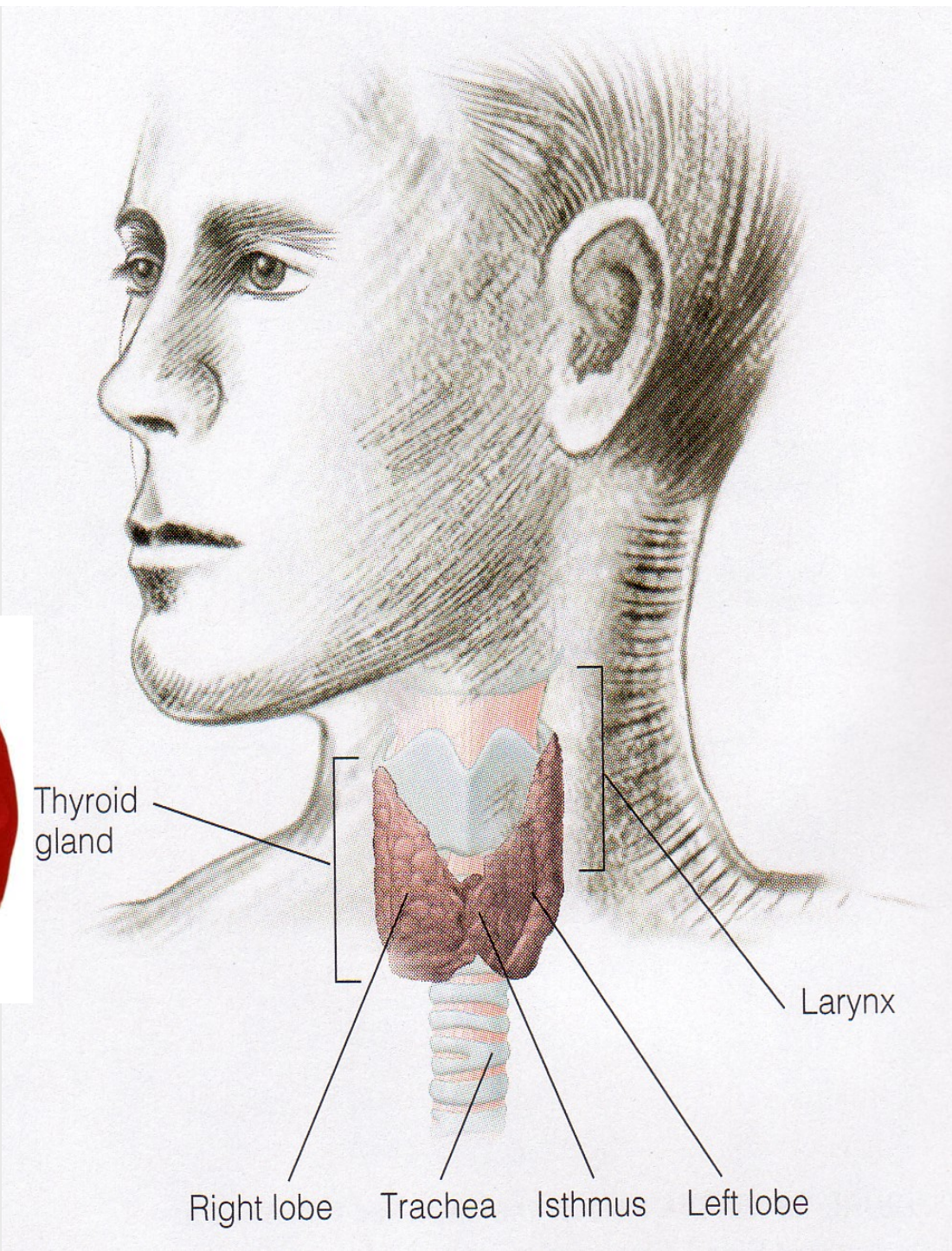
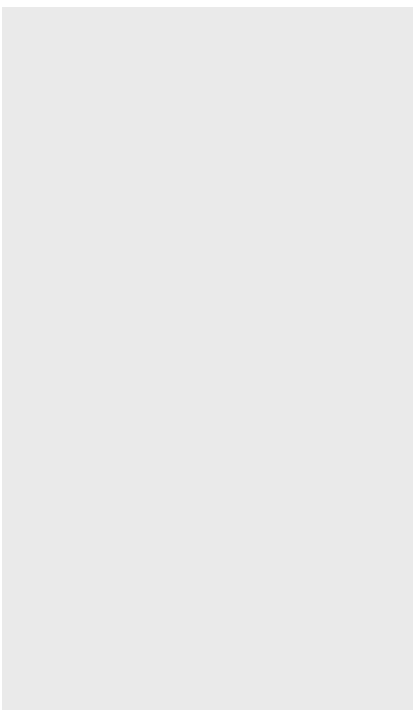
Glucose:
Sugar in Blood

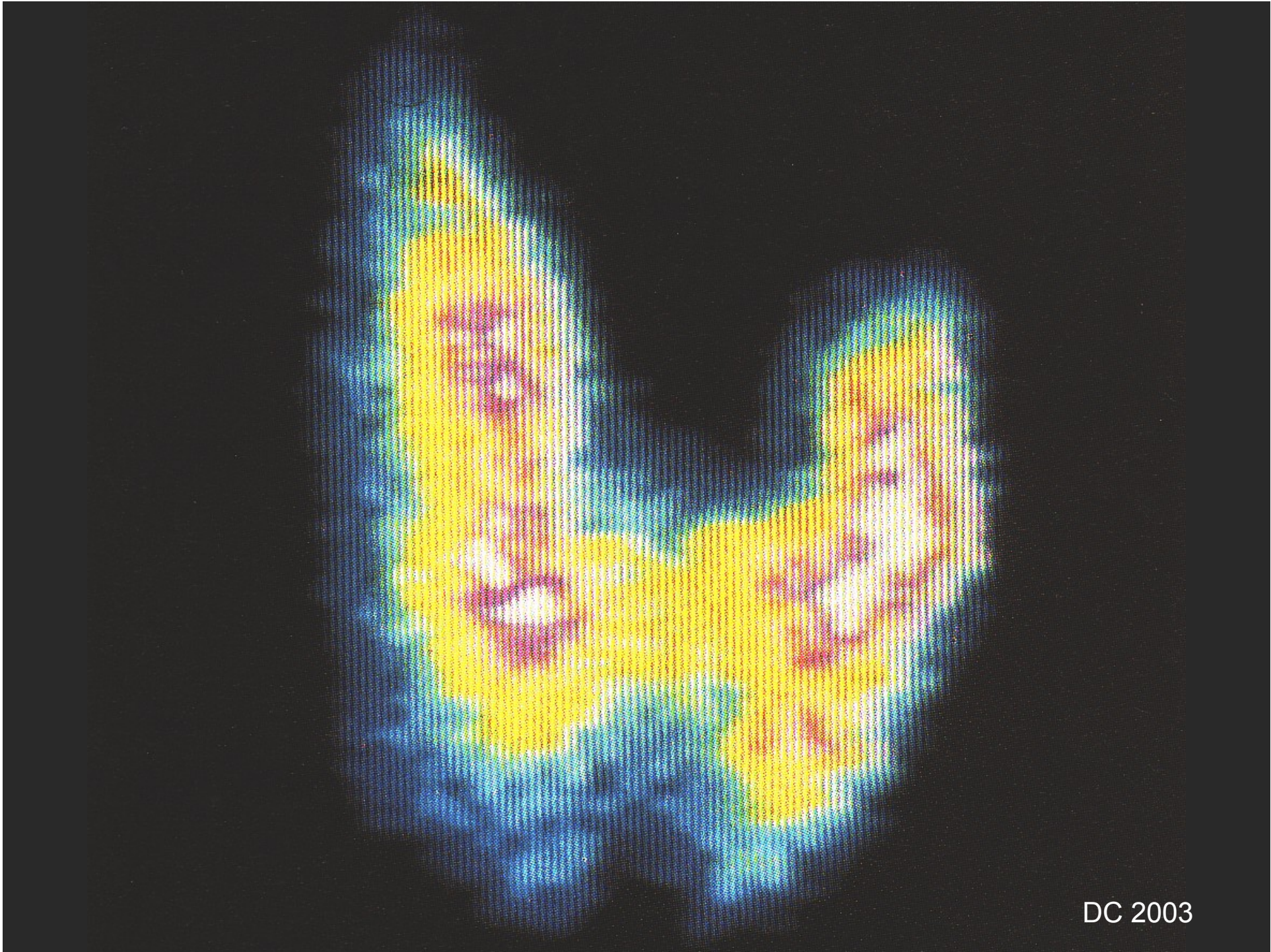


Normal: 70-99

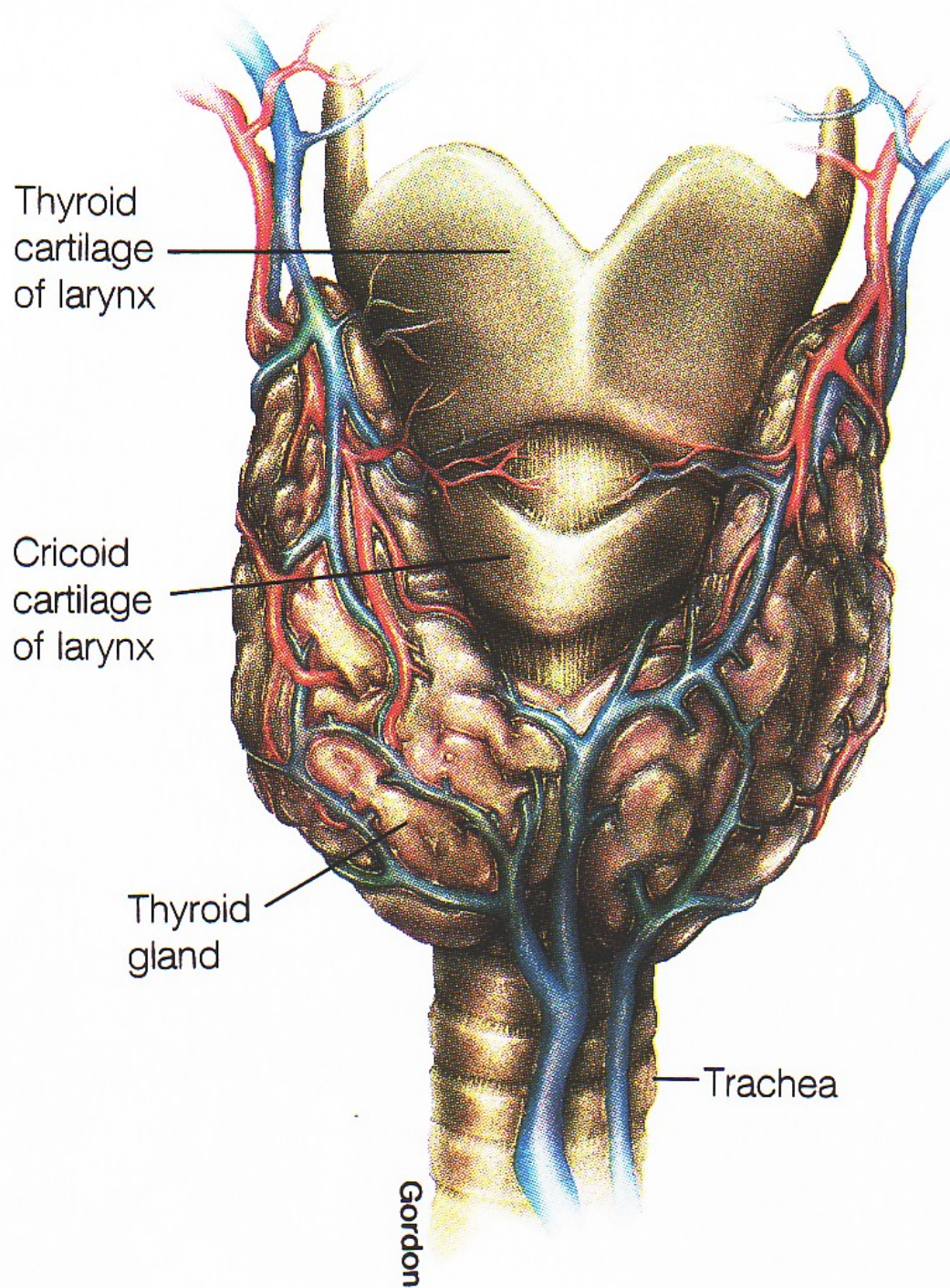
Pre-Diabetes: 100-125

Diabetes: ≥ 126 mg/dL



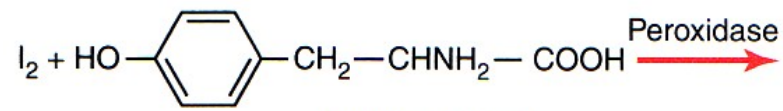


DC 2003

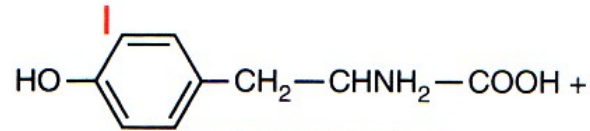


(a)

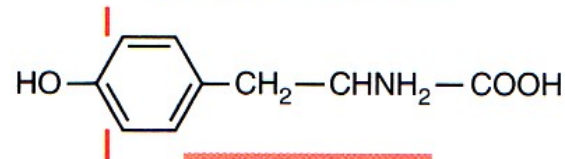
DC 2003



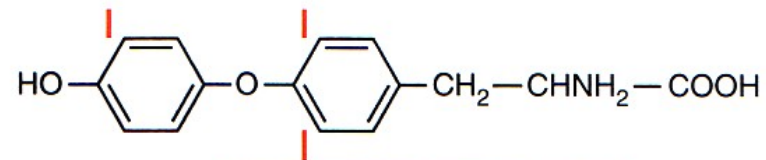
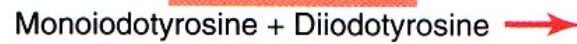
Tyrosine



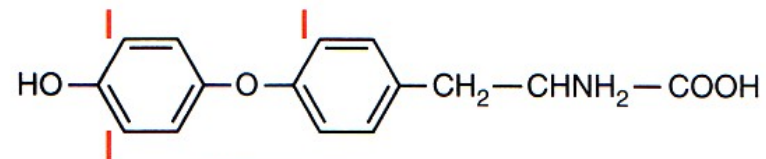
Monoiodotyrosine



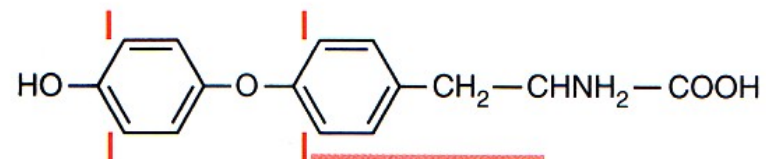
Diiodotyrosine



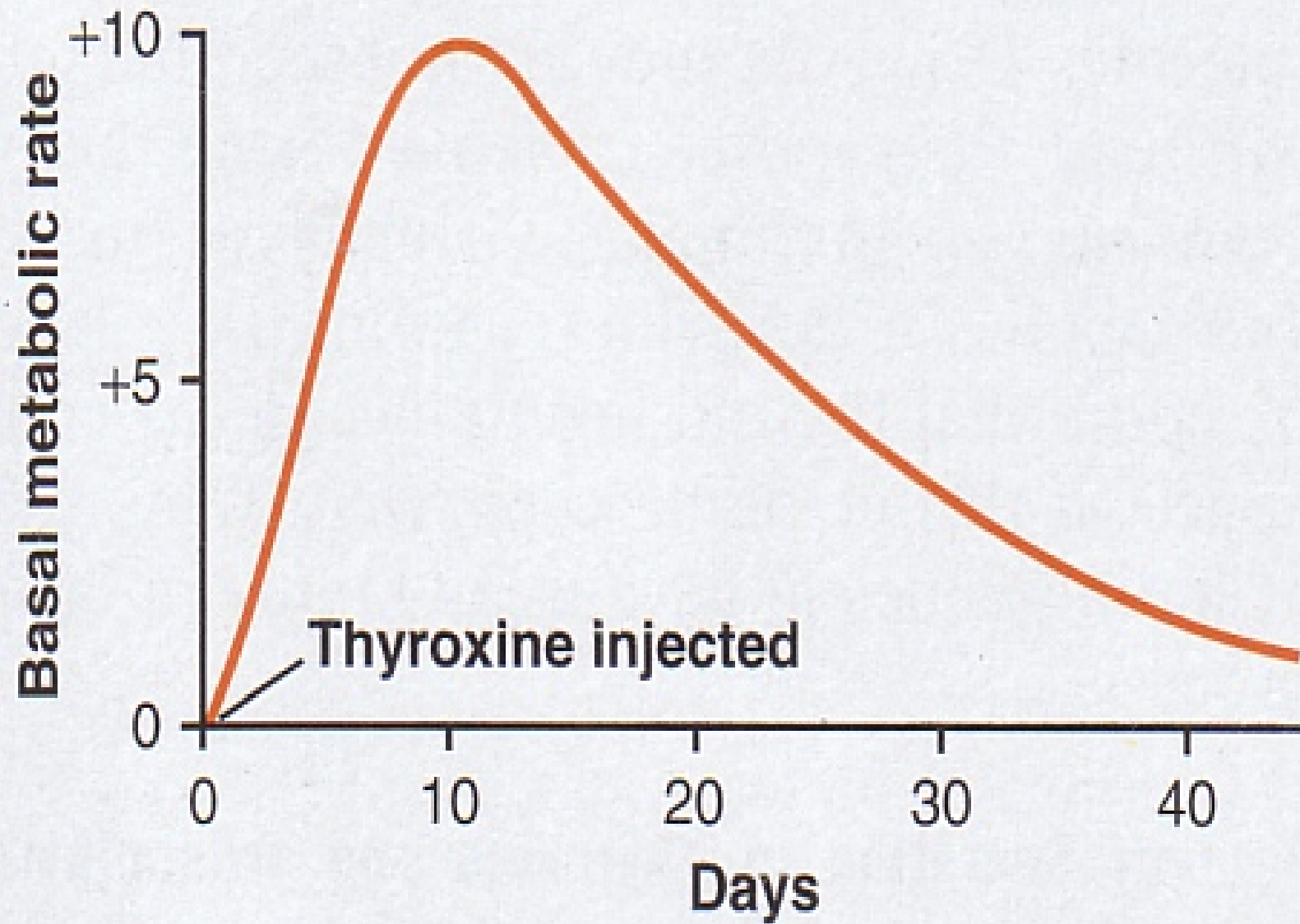
3,5,3'-Triiodothyronine (T₃)



3,3',5-Triiodothyronine (RT₃)



Thyroxine (T₄)



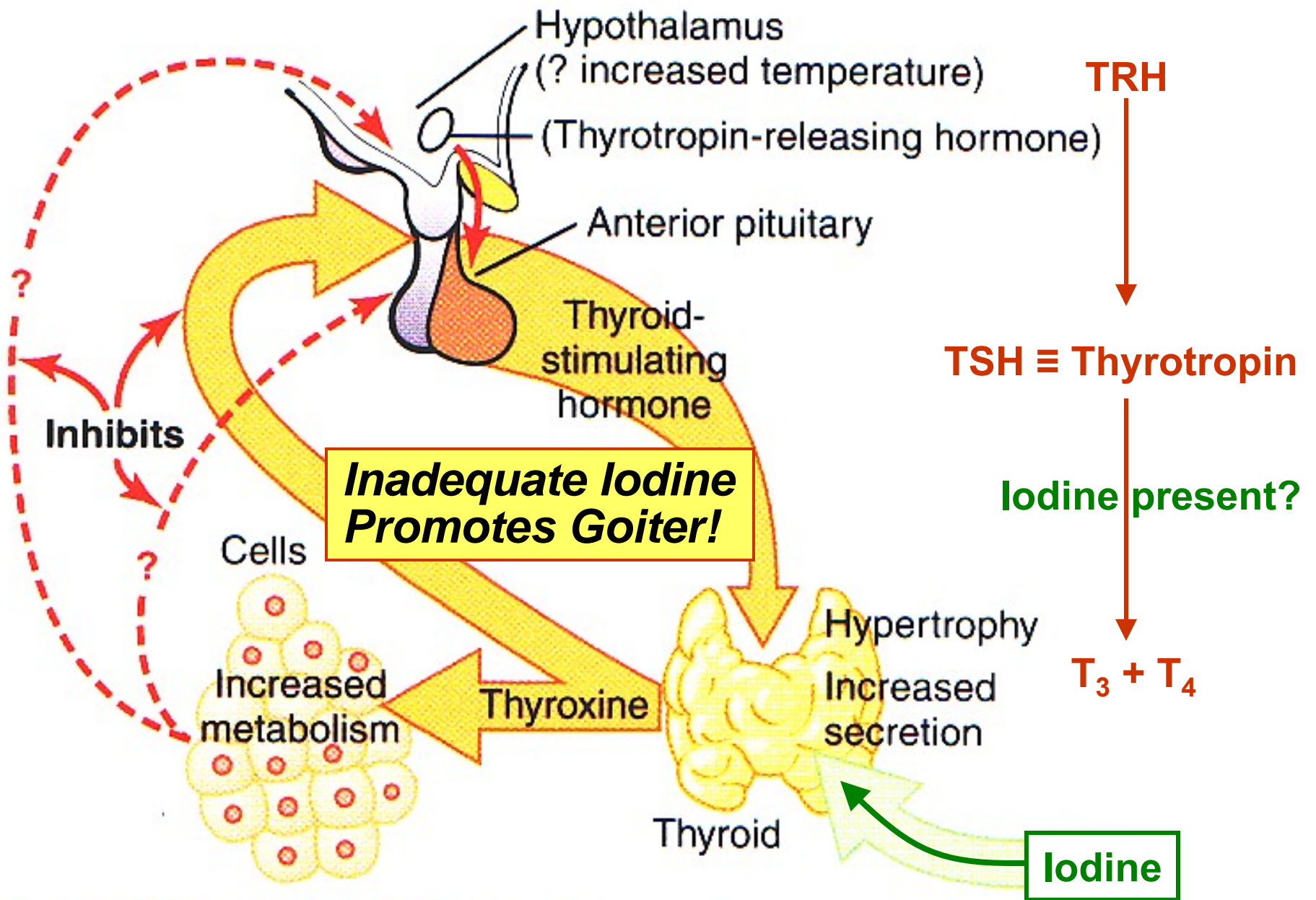


Figure 76-7 Regulation of thyroid secretion.



DC 2003

cf: G&H
2011
fig 76-8



DC
2003

Near
absence of
thyroid-
hormone
function +
myxedema



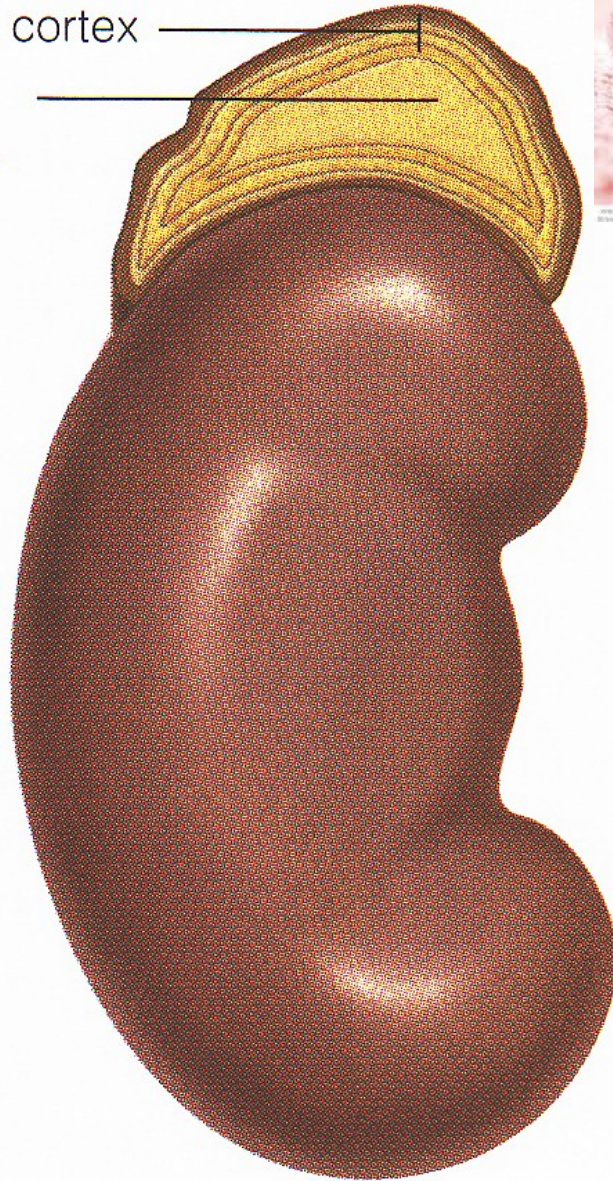
Figure 76-8. Patient with myxedema. (Courtesy of Dr. Herbert Langford.)

Adrenal gland

Adrenal cortex

Adrenal medulla

Kidney

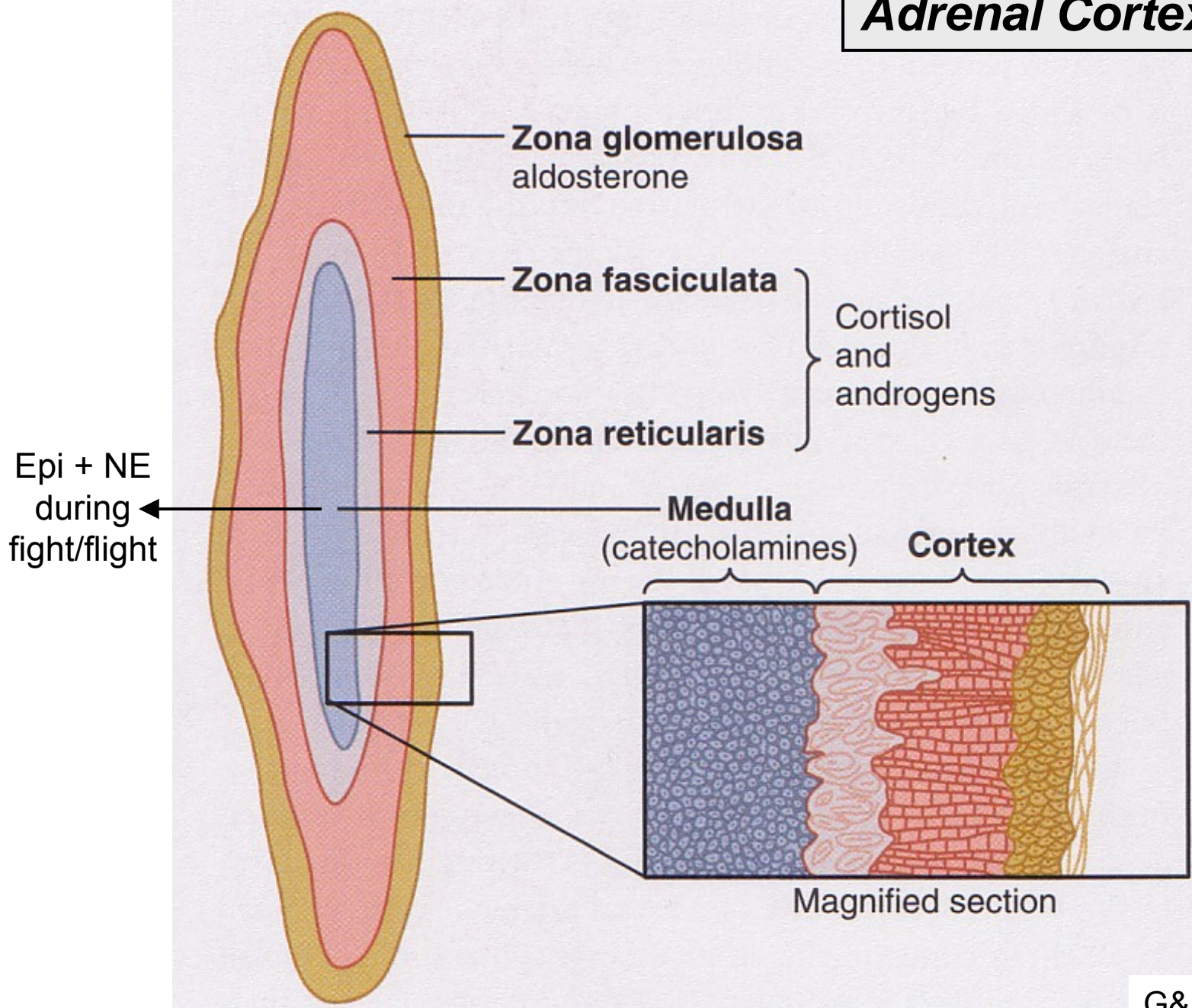


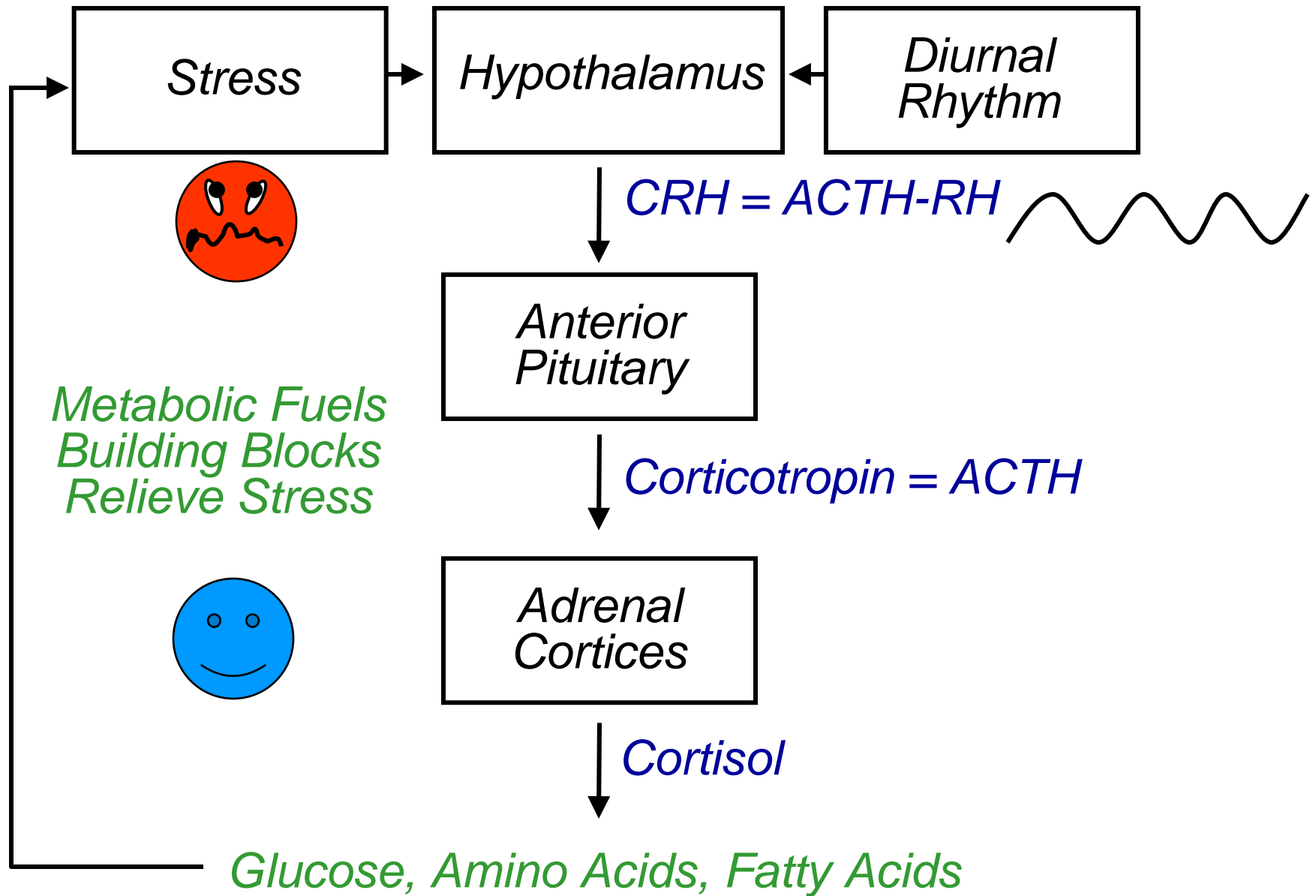
Scoop of
ice cream on
North pole!

FIGURE 13-12

Adrenal Gland The adrenal glands sit atop the kidney and consist of an outer zone of cells, the adrenal cortex, which produces a variety of steroid hormones, and an inner zone, the adrenal medulla. The adrenal medulla produces adrenalin and noradrenalin.

Adrenal Cortex Zones





SOURCE: Modified after D Chiras 2003