

Rethinking Metabolic Syndrome

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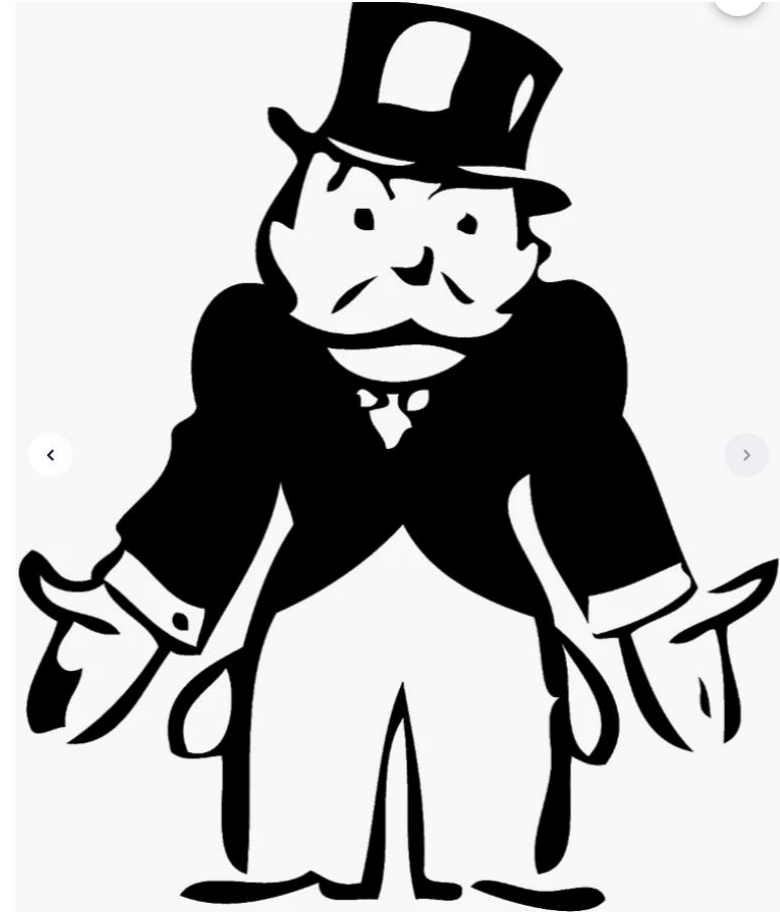
Pacific Northwest University of Health Sciences

College of Osteopathic Medicine

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Disclosures

- None



Educational Objectives

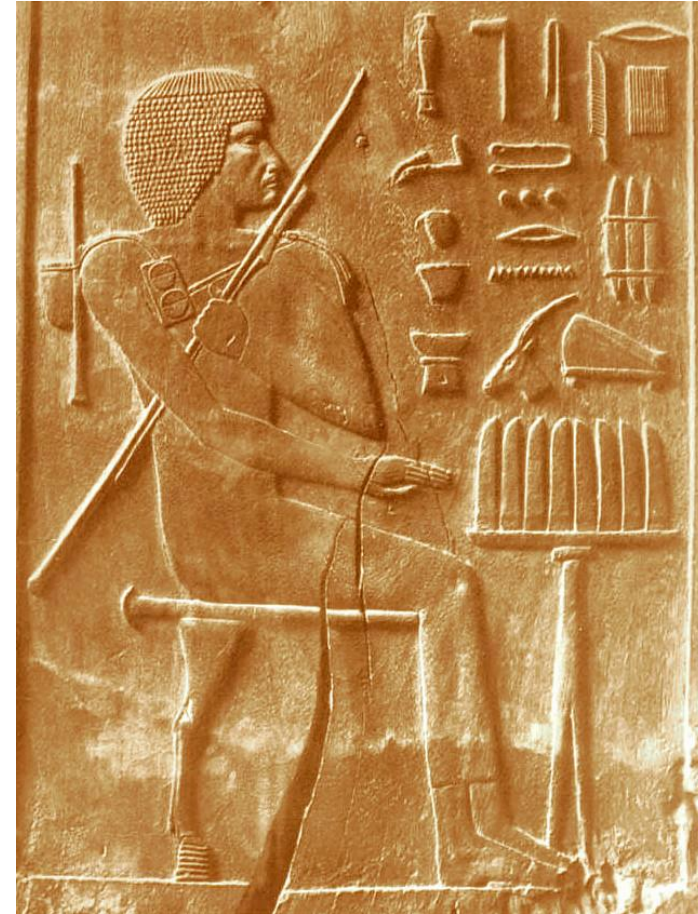
- Have a better understanding of the common underlying mechanisms of CKM Syndrome and their multisystem manifestations
- Describe the individual features and manifestation of specific end organ involvement in CKM
- Be able to classify at risk patients for CKM Syndrome and list risk factors
- Understand and apply evidence based treatment and early intervention to prevent or attenuate the systemic damage

Suggested Readings

- Chiadi E. Ndumele. Circulation. Cardiovascular-Kidney-Metabolic Health: A Presidential Advisory From the American Heart Association, Volume: 148, Issue: 20, Pages: 1606-1635, DOI: (10.1161/CIR.0000000000001184)
- 2023 ADA Standards: [Volume 46 Issue Supplement 1 | Diabetes Care | American Diabetes Association \(diabetesjournals.org\)](#)
- [Winkelmayer KDIGO-Arrhythmia-NKF-NOLA-2020.pdf](#)
- [Blood Pressure in CKD – KDIGO](#)
- [Diabetes in CKD – KDIGO](#)

History of Diabetes Mellitus

- Hesy-Ra ~1,552 B.C.E.
- 1st described treatments for the “passage of too much urine” which attracted ants and was linked to a state of progressive emaciation ants
- First link of kidney to diabetes



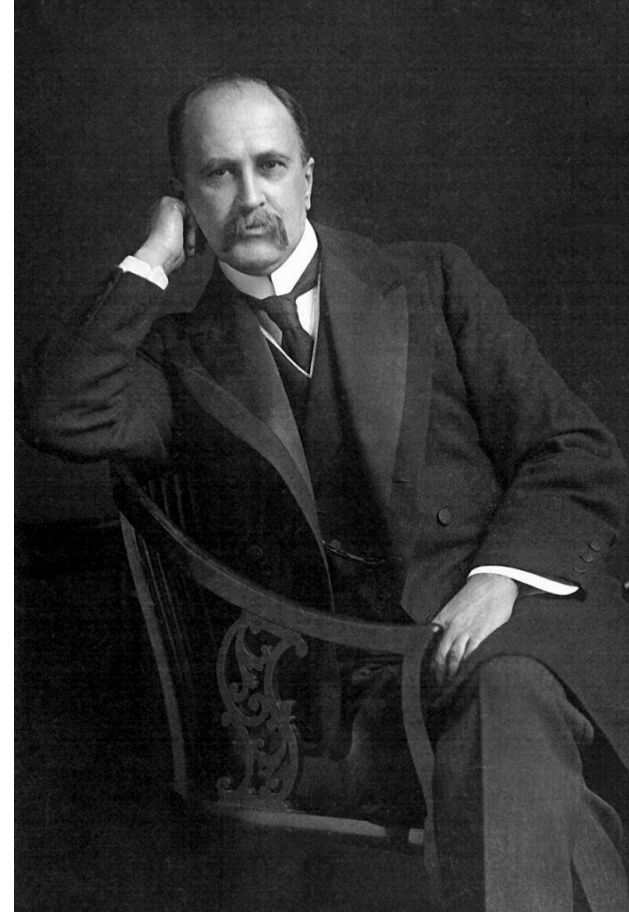
Thomas Willis 1621-1675



- Best known for his brain anatomical work-Circle of Willis
- First to number the cranial nerves (much to the chagrin of 1st year med students)
- Willis 1st used the term *diabetes mellitus*, which he associated with melancholy (depression)
 - Willis's disease is an archaic term for DM

Sir William Osler 1848-1919

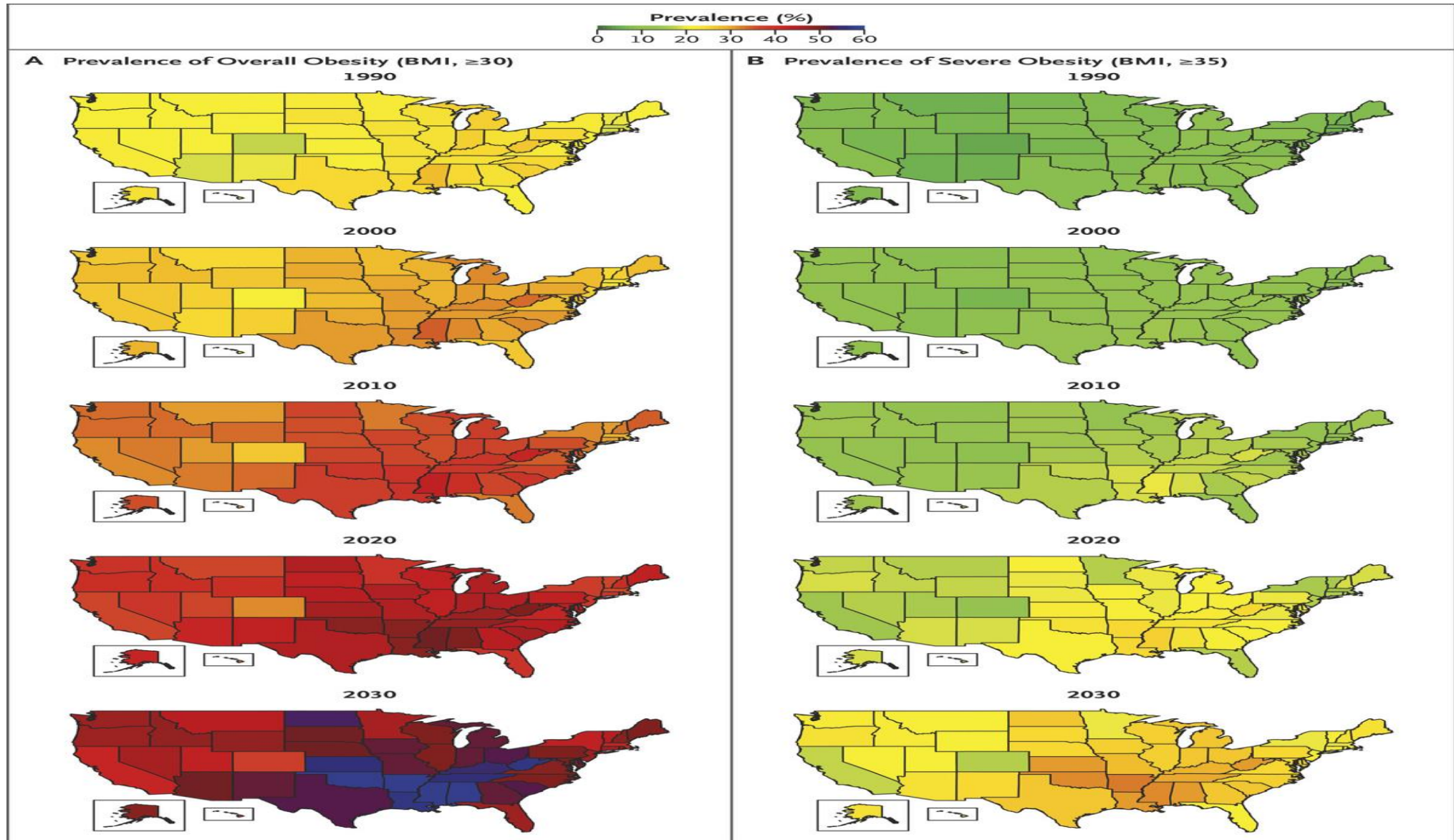
- The platter kills more than the sword



Typical Inpatient Problem circa 1979: 10 Plus Conditions

- 1. Obesity
- 2. Type-DM
 - Retinopathy, peripheral neuropathy, kidney disease
- 3. Hypertension
 - Hypertensive kidney and heart disease
- 4. Hyperlipidemia
- 5. Obstructive sleep apnea
- 6. Cerebral vascular disease
 - Hx TIA
- 7. Coronary heart disease
 - s/p MI
- 8. Peripheral vascular disease
 - s/p below the knee amputation
 - Foot ulcer
- 9. Chronic kidney disease
 - Secondary to DM and HTN
- 10. Congestive heart failure

Estimated Prevalence of Overall Obesity and Severe Obesity by State 1990-2023



Presidential Advisory Cardiovascular-Kidney Metabolic Syndrome (CKM)

Circulation

AHA Journals

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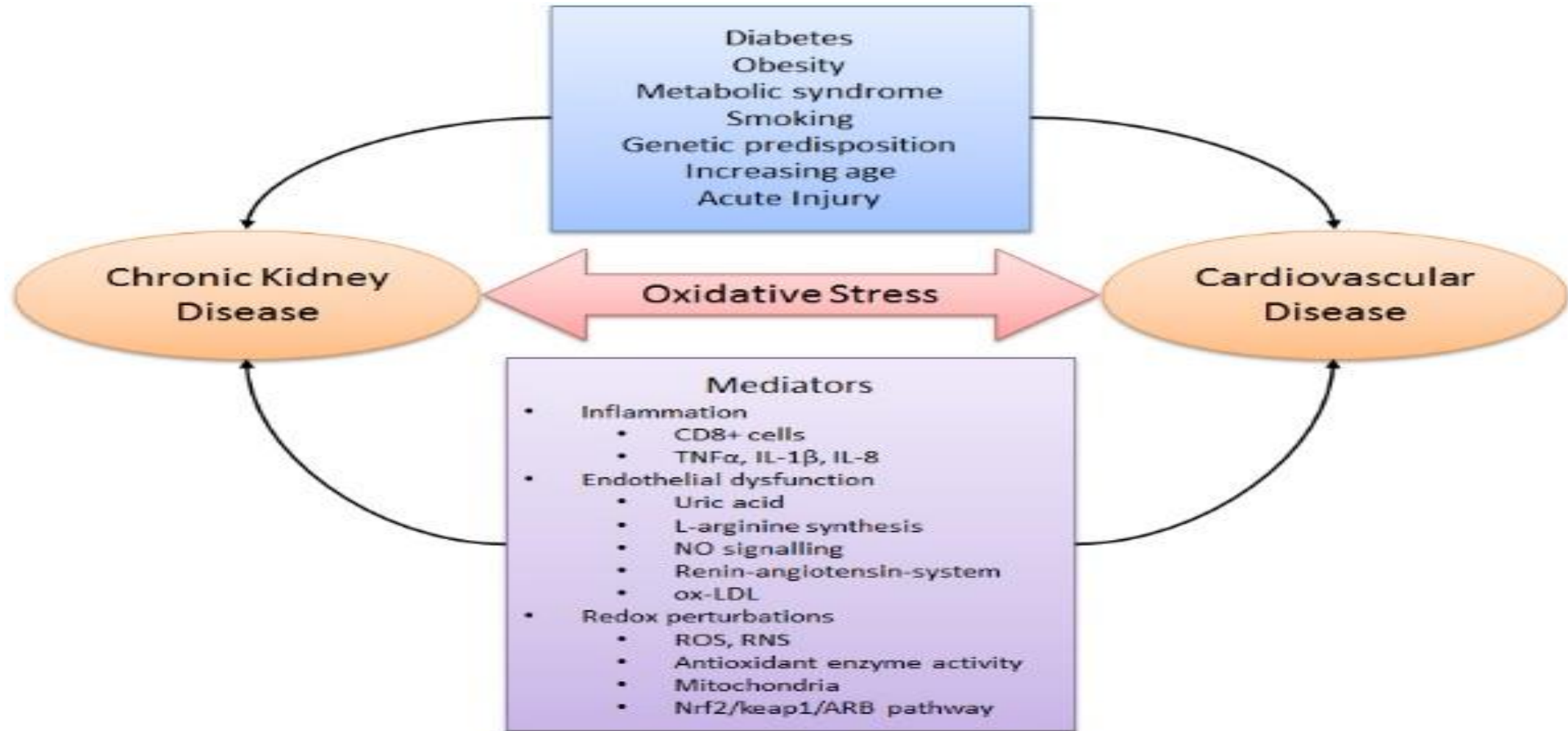
Cardiovascular-Kidney-Metabolic Health: A Presidential Advisory From the American Heart Association

Significance: The first comprehensive incorporation of risk factors, staging, and interventions which include addressing the Social Determinants of Health (SODH). The treatment recommendations are evidence based and incorporate current guidelines from multiple societies.

Definition

- CKM syndrome is a systemic disorder characterized by pathophysiological interactions among metabolic risk factors, CKD, and the cardiovascular system leading to multiorgan dysfunction and a high rate of adverse cardiovascular outcomes.
 - CKM syndrome includes both individuals at risk for CVD due to the presence of metabolic risk factors, CKD, or both and individuals with existing CVD that is potentially related to or complicates metabolic risk factors or CKD.
 - Encompassing multisystem metabolic derangement leading to generation of neuroendocrine and inflammatory mediators leading to endothelial damage.
 - Instead of 10 individual conditions; 1 condition with multiple manifestations
- *Circulation*. 2023;148:4.

Kidney Disease = Cardiovascular Disease



Pathophysiology of Metabolic Syndrome/T2DM

Alteration of Multiple Neuroendocrine Systems

A Perfect Storm of Genetics and Environment

Neuroendocrine mediators

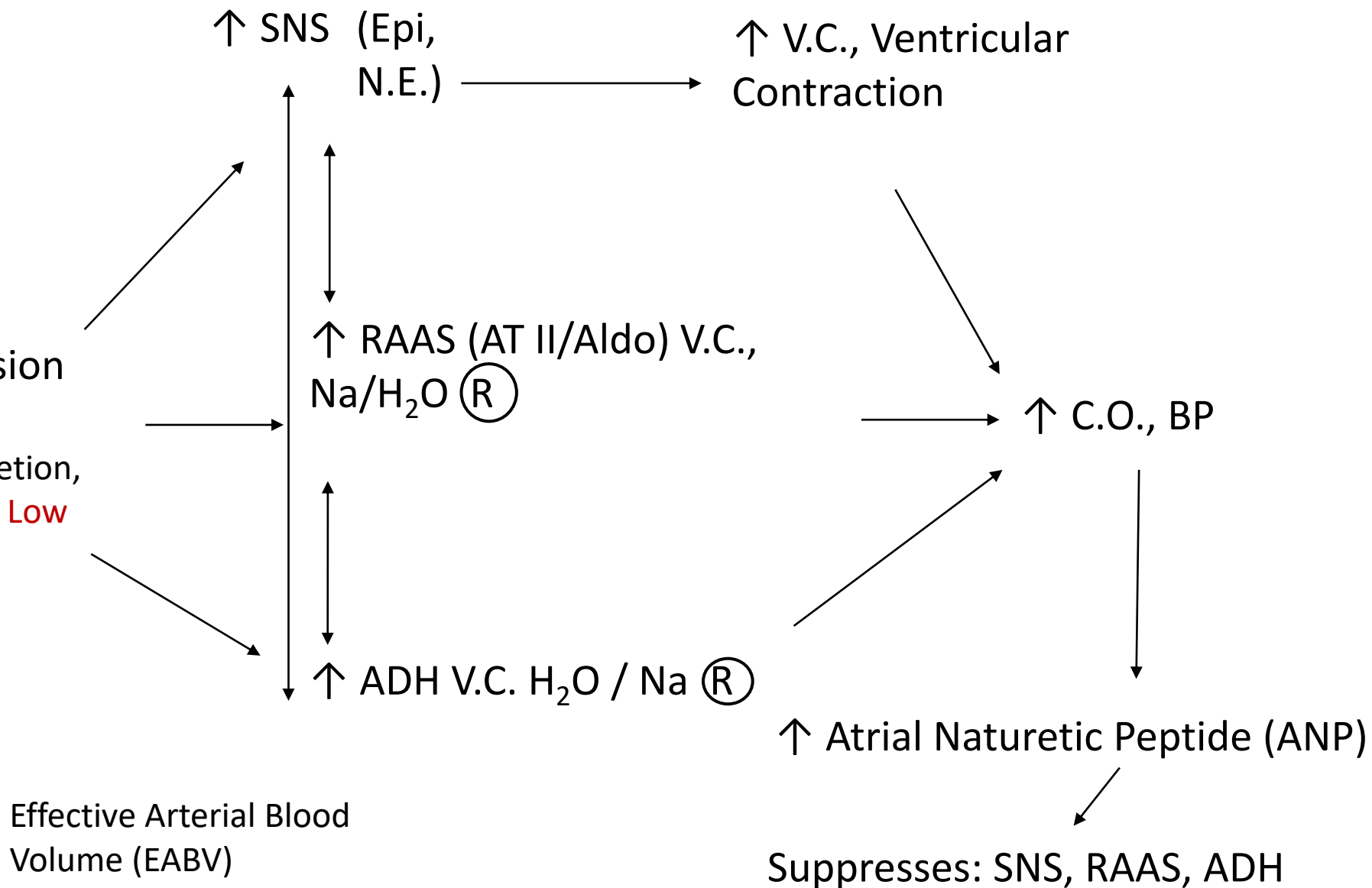
- Renin-angiotensin-aldosterone
- Insulin-glucose-glucagon-cortisol-growth hormone
- Catecholamines
- Glucocorticoids
- Endothelin-1
- Growth hormone
- Endocannabinoids
- Low level of adiponectin

Inflammatory Mediators

- Adipokines
- Leptin
- IL-6
- TNF- α
- Monocyte chemoattractant protein-1 (MCP-1)
- Resistin

How the Body Responds to Volume Depletion & Metabolic Syndrome

Hypoperfusion
& CO/BP
(Volume depletion,
shock, sepsis) **Low**
EABV



Glossary	
SNS	Sympathetic Nervous System
Epi	Epinephrine
N.E.	Norepinephrine
V.C.	Vasoconstriction
R	Reabsorption
RAAS	(Renin Angiotensin Aldosterone System)
ADH	Antidiuretic Hormone
ANP	Atrial Natriuretic Peptide

Effective Arterial Blood
Volume (EABV)

↑ Atrial Natriuretic Peptide (ANP)
Suppresses: SNS, RAAS, ADH

White Adipose Tissue As an Endocrine Organ

- **Adipose tissue** is a source of **inflammatory mediators** leading to **insulin resistance (IR)** and other features of **metabolic syndrome**
- Increased adipose tissue outstrips vasculature leading to local tissue **hypoxia** with **subsequent inflammation** and **angiogenesis**
- The **net effect** is **ACCELERATED** atherosclerosis and increased **CV risk**

Insulin Resistance (IR)

- **Definition:**

- **Inability** of **insulin** levels (endogenous or exogenous) to increase **glucose uptake, metabolism, and utilization** by muscle, adipose, and liver tissues

- **Types of IR:**

- Pre-receptor
- Receptor
- Post-receptor
- Defective insulin signaling:
 - Decrease GLUT-4 receptor

Insulin Resistance (IR)

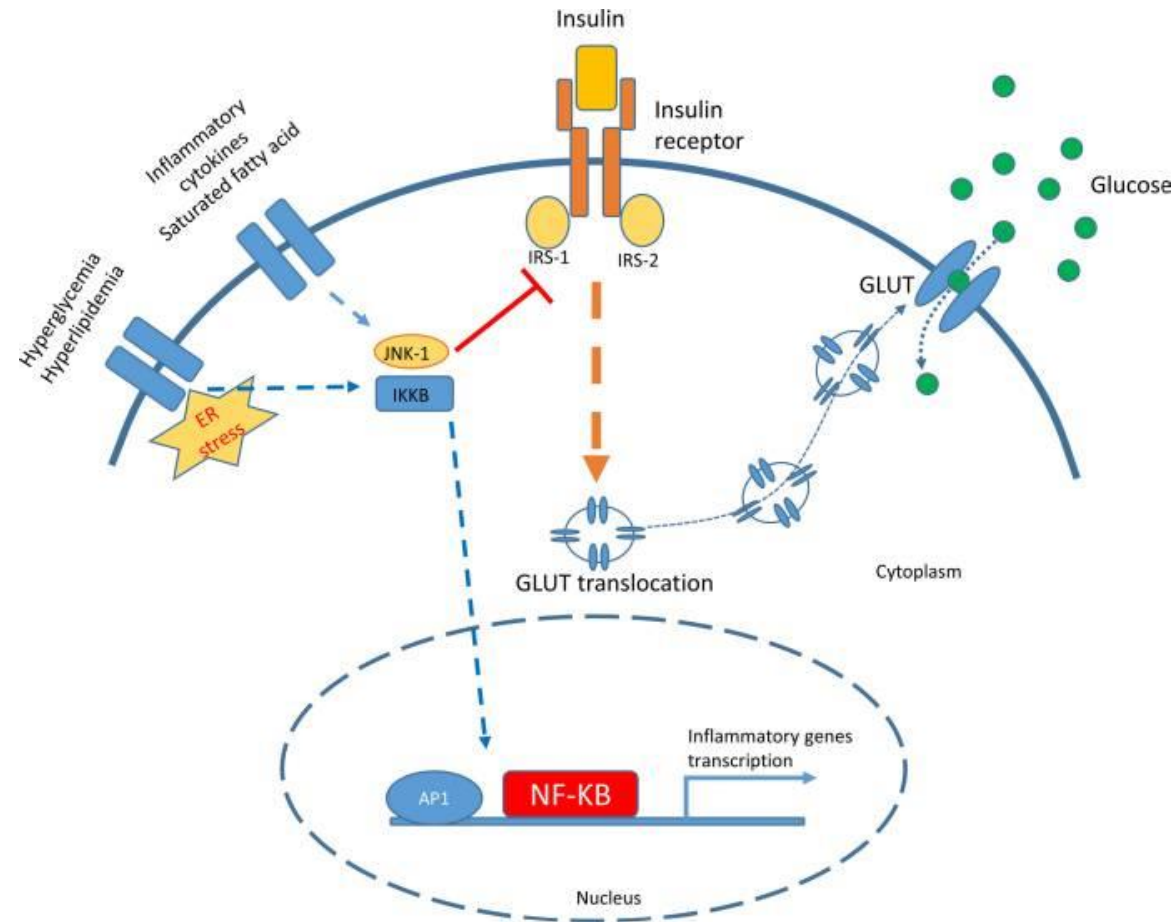
- **Acquired:**

- Excess dysfunctional adipose tissues
- Physical inactivity
- Nutritional imbalance, irregular eating/fructose
- Aging
- High sodium diets
- Glucotoxicity (end organ)
- Lipotoxicity (dietary)
- Medications

- **Genetic:**

- Myotonic dystrophy
- PCOS
- Type A insulin resistance
- Ataxia-telangiectasia
- Alstom syndrome
- Rabson-Mendenhall syndrome
- Werner syndrome
- Lipodystrophy

Inflammation and Insulin Resistance



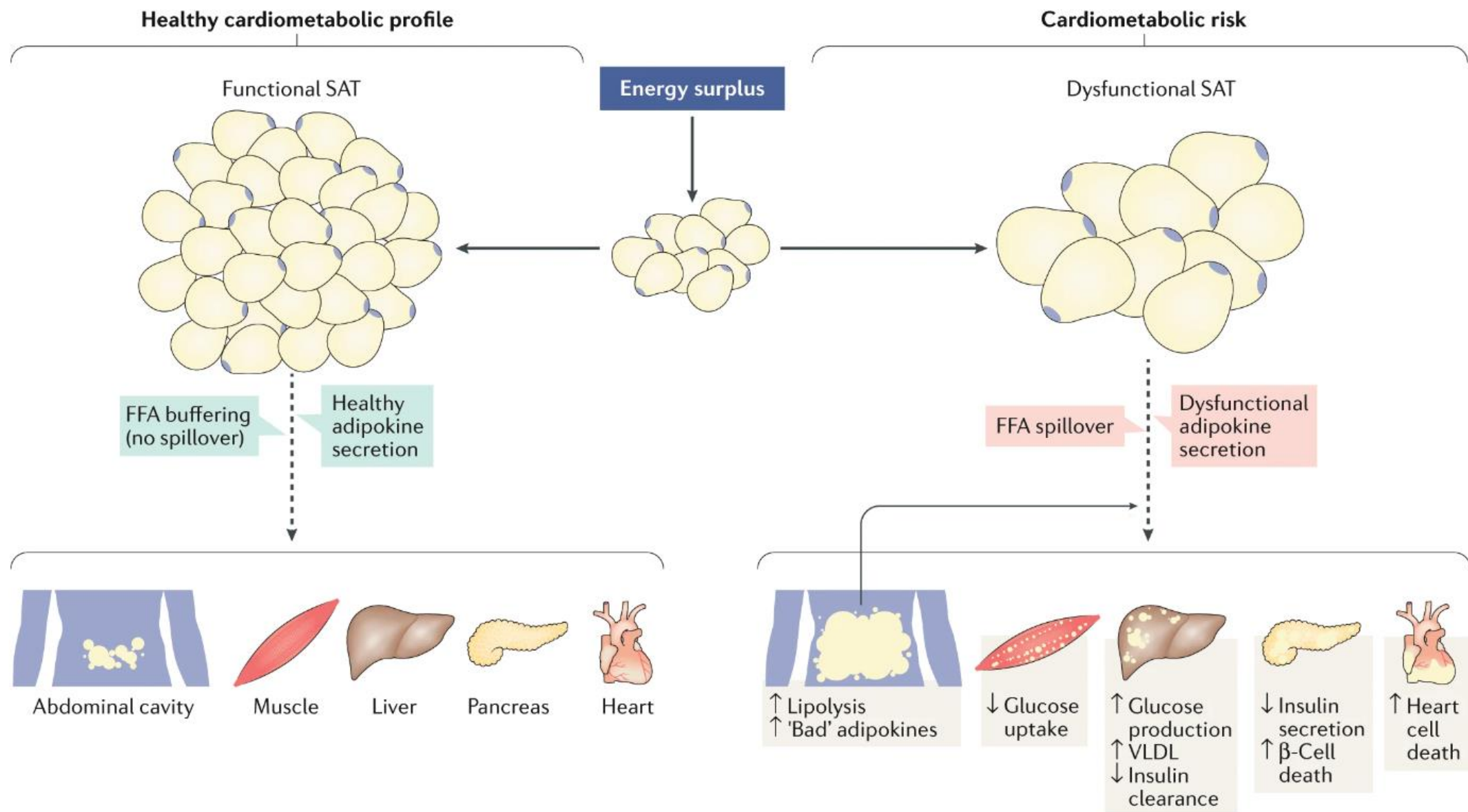
Lipotoxicity

- **Excess/ dysfunctional adipose tissue** secrete **proinflammatory** and **prooxidative factors** and **neuroendocrine mediators** that **promote injury** to **endothelial, cardiac, renal, and hepatic tissues**. These factors also **decrease sensitivity** to the actions of **insulin** leading to **impaired glucose tolerance**

Lipotoxicity

- **Dysfunctional adipose tissue/lipotoxicity:**
 - **Excess dietary glucose/fructose** induced **de novo lipogenesis***
 - Truncal obesity
 - Fatty infiltration of liver, mesentery, heart, liver, etc.
 - NAFLD/MAFLD
 - **Fatty acid induced IR:**
 - De novo lipogenesis
 - **Lipid deposition** in **muscle** tissues (all **3** types)
 - **Lipo-and glucotoxicity** induced **inflammation:**
 - Promotes insulin resistance
 - **Low grade SIRS:**
 - Elevated CRP, Il-6, TNF, etc.

*Schwartz JM, Clearfield M, Mulligan K, *J Osteopathic Medicine* 2017;117:520-527



Effects of Chronically Elevated Glucose Levels

- **Long term effects:**

- Worsening insulin resistance **IR**
- Inflammation, oxidative stress
- Excess glucose leads to de novo fatty acid production and deposition into tissues

- **Genesis of end organ damage:**

- Glucotoxicity
- Insulin resistance
- Lipotoxicity
- AGEs

- **Hyperfiltration of the kidney:**

- RAAS activation
- Subsequent activation of SNS, ADH

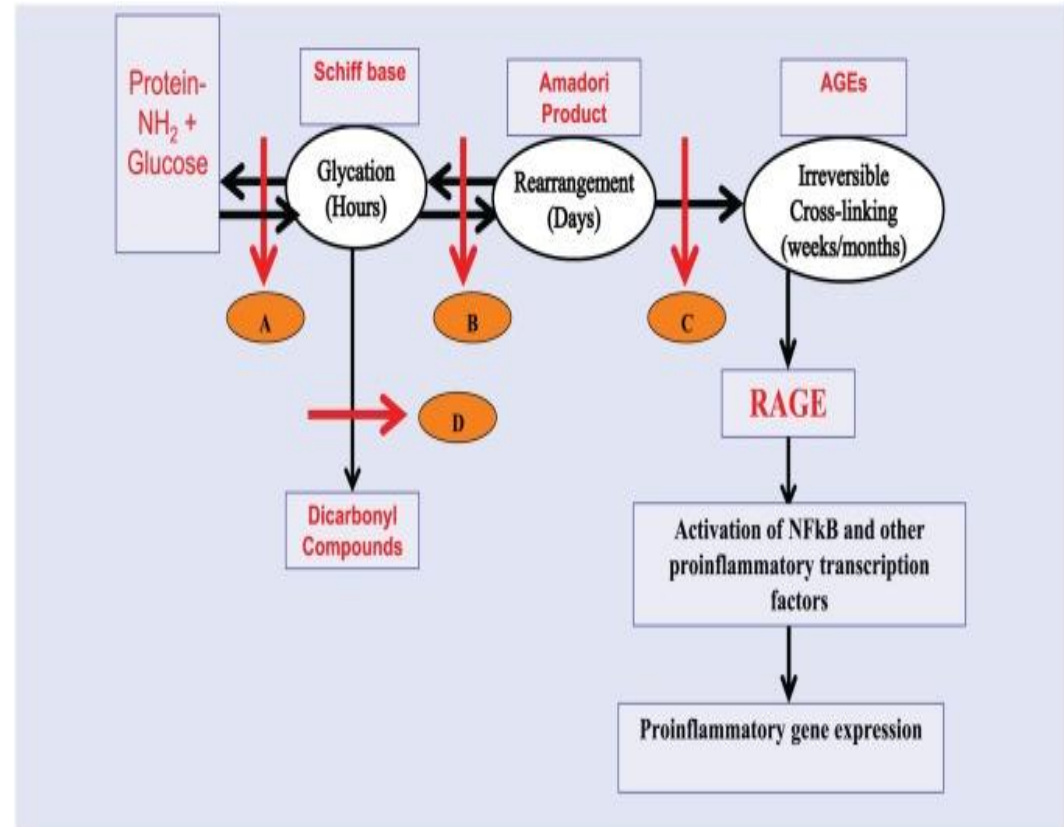
Glucotoxicity

- **Definition:**

- Impaired β -cell function/response during states of elevated glucose
- Mitochondrial stress
- Advanced glycosylated end-products (AGEs)
- End organ damage

Effects of Hyperglycemia: Advanced Glycation/Glycosylation End Products (**AGE**)

- **Elevated glucose levels (glucotoxicity) lead to:**
 - Metabolic interactions with proteins to form **AGE** leading to **tissues injury** and **inflammatory state**



AGEs: Coming to an End Organ Near You Soon



- Kidney
- Retina
- Nerves
- Vasculature

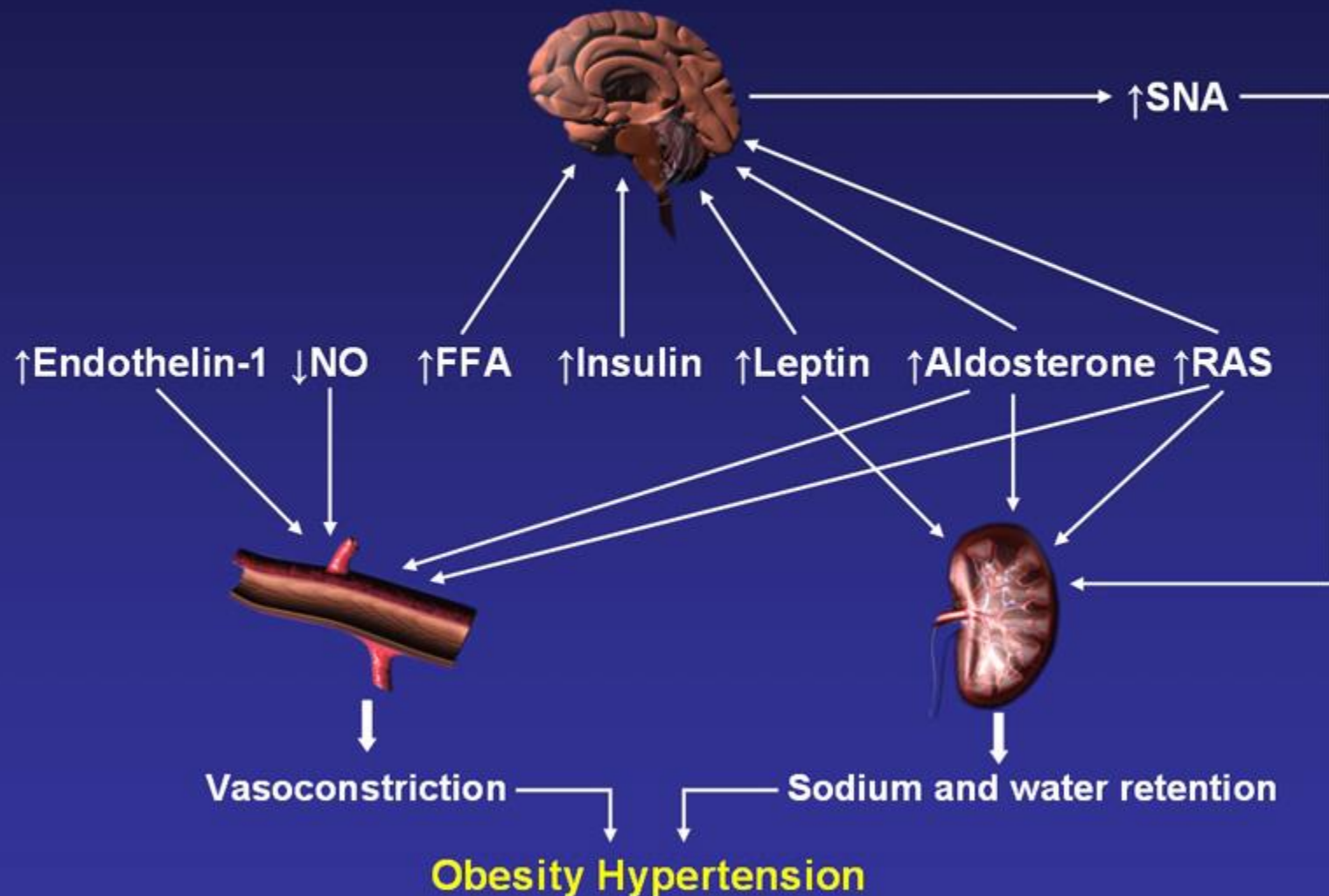
Increased Sympathetic Activity

- Elevated **epinephrine** (Epi) and **norepinephrine** (NE) levels are frequently seen in Type 1 and 2 DM renal disease
- Epi and NE add to **hyperfiltration** by actions on the nephron, peripheral vessels, and positive inotropic and chronotropic effect leading to **elevated BP** and **increased** glomerular capillary pressure (P_{GC})
- Epi and NE **activate** the **RAAS**
- **Increased** all-cause CV risk in part due to increased SNS tone

Aldosterone in Diabetes

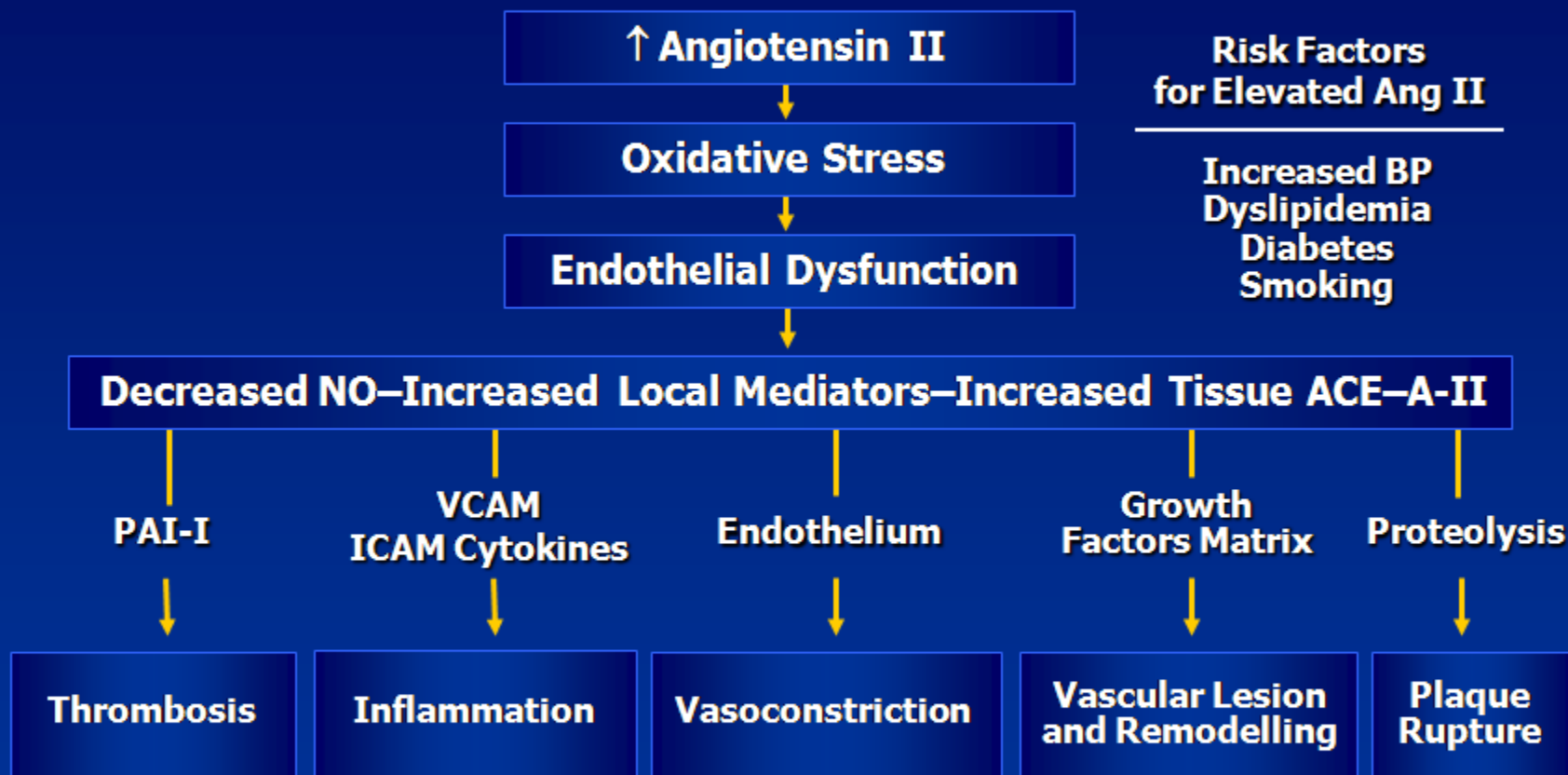
- Increased activity of the **renin-angiotensin-aldosterone system (RAAS)** of seen in and contributes to the pathogenesis of many conditions
 - DM, HTN, cardiovascular, hepatic and kidney diseases
 - Inhibition, blockade are key therapeutic targets
- Patient with **primary hyperaldosteronism** (Conn's syndrome) have a higher incidence of CV events, insulin resistance, and impaired glucose tolerance
- Elevated or upper limit of normal levels of serum aldosterone is strongly associated with **glucose intolerance** (pre-diabetes), **insulin resistance**, and **Type 2 DM**

Mechanisms and Hormonal Systems Involved in Obesity-Associated Hypertension



NO= nitric oxide; FFA= free fatty acids; RAS= renin-angiotensin system; SNA= sympathetic nerve activity.
Rahmouni K et al. *Hypertension*. 2005;45:9–14.

Effect of Tissue-Specific Angiotensin II Upregulation



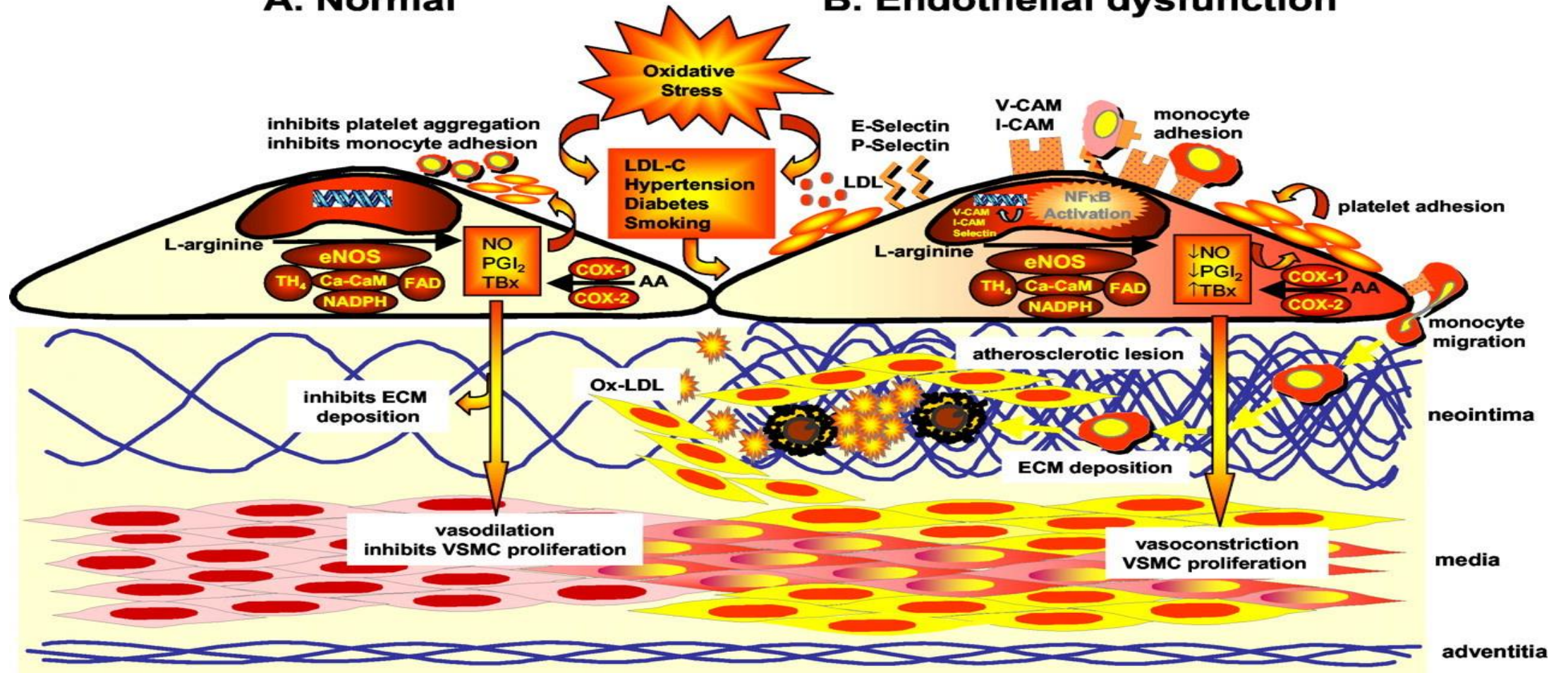
Autocrine or paracrine production of Ang II is not ACE-dependent

Ang II=angiotensin II; NO=nitric oxide; PAI-1=plasminogen activator type 1; VCAM=vascular cell adhesion molecule.

Key Target: Endothelial cells

A. Normal

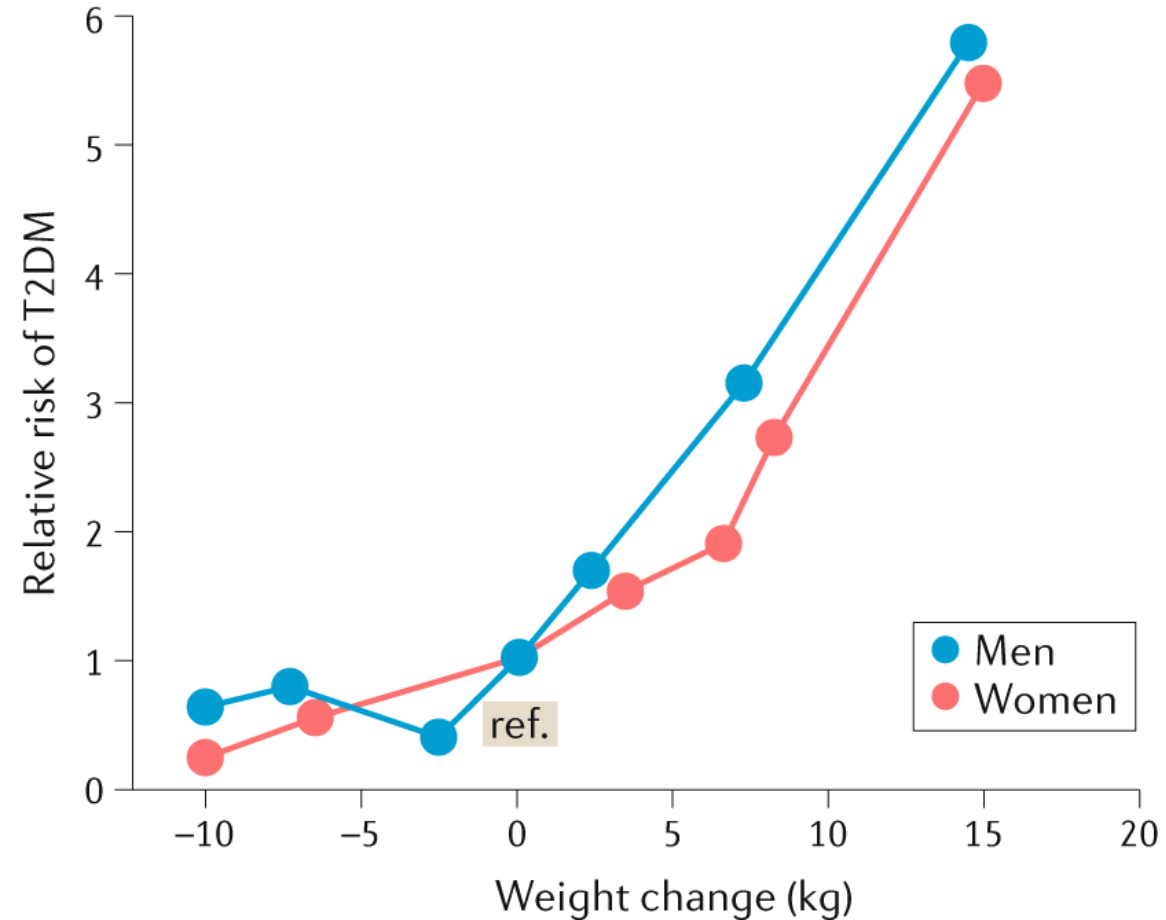
B. Endothelial dysfunction



Obesity to Metabolic Syndrome to T2DM

- Increased **caloric** intake:
 - Simple sugars (fructose), refined, highly processed foods, added antibiotics/steroids all lead to increased insulin requirement
 - Alteration of glucose stimulated insulin secretion (GSIS)
 - Increased renal reabsorption
 - B-cell exhaustion
- Lower intracellular glucose levels (from decreased insulin level or resistance) lead to increased counter regulatory hormones and RAAS
 - **Systemic effects**: Cardiorenal disease, vascular, neuropathic disease, etc.

Role of Weight Gain and Loss of Risk of T2DM



Key Players in CKM Syndrome

- Body mass index (BMI):
 - Metabolic risk factors
- Waist circumference:
 - Metabolic risk factor
 - Circumference should be an independent vital sign 1
- Fasting blood glucose/ HbA1c Metabolic risk factor
- Lipids (LDL, HDL, Triglycerides)
- Blood pressure:
- Chronic kidney disease of any cause (elevated creatinine and/or albuminuria)
- Hepatic steatosis, inflammation, fibrosis/cirrhosis
- Sleep duration/quality:
- Address mental health challenges/stressors
- Address Social Determinants of Health (SODH) which may be impacting the patient's life

Staging of CKM

- Stage 0: Normal weight, glucose, lipids, BP, no CKD or subclinical CVD
- Stage 1: Excess/Dysfunctional Adiposity
- Stage 2: Metabolic Risk Factors and/or CKD
- Stage 3: Subclinical CVD or CKD
- Stage 4: Clinical CVD in CKM

Stage 1: Excess/Dysfunctional Adiposity

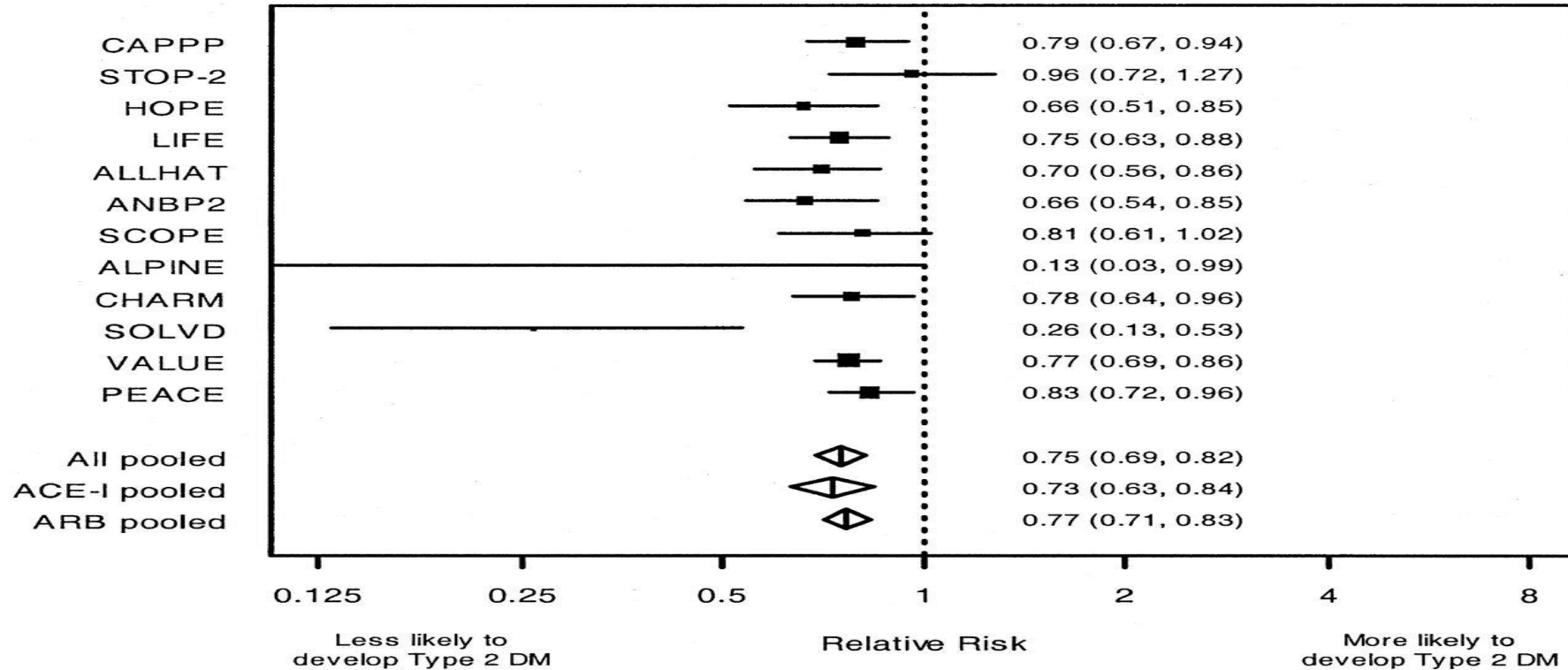
- Overweight, abdominal or dysfunctional obesity
- Waist circumference should be considered as an independent vital sign¹
- No other metabolic, CKD, or CVD risk factors
 - Normal BP, glucose, no albuminuria, normal lipids, non-tobacco use,

1. Waist circumference as a vital sign in clinical practice: a Consensus Statement from the IAS and ICCR Working Group on Visceral Obesity. *Nat Rev Endocrinol* **16**, 177–189 (2020). <https://doi.org/10.1038/s41574-019-0310-7>

Do ACE-I or ARBs Prevent Diabetes?

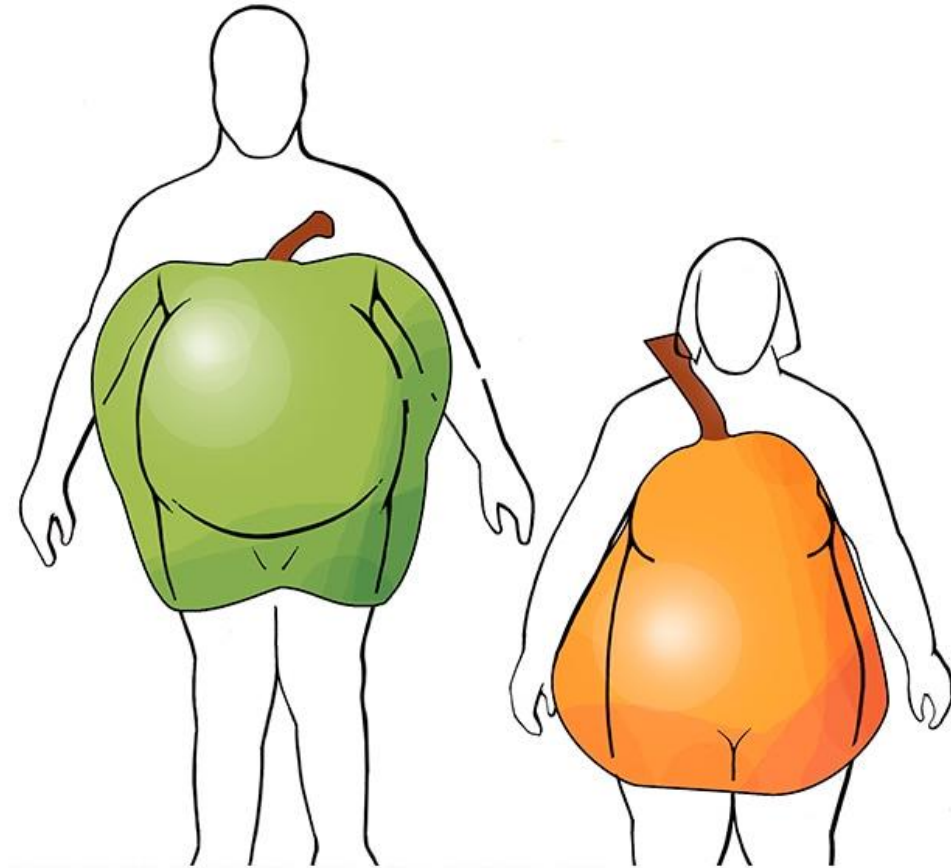
- Meta-analysis of 12 studies involving non-diabetics and the reduction of new onset of T2DM
- 72,333 non-diabetic patients, enrolled in one of 12 trials
 - **ACE-i treated: 27% reduction** in new onset T2DM
 - **ARBs treated: 23% reduction** in new onset T2DM
 - **Pooled: 25% reduction** of new onset T2DM

The Effect of RAAS Blockade on Development of Type 2 Diabetes



Stage 1: Excess/Dysfunctional Adiposity

- Overweight, abdominal or dysfunctional obesity
- **Waist circumference** should be considered as an **independent vital sign**¹
- No other metabolic, CKD, or CVD risk factors
 - Normal BP, glucose, albuminuria, lipids, non-tobacco use



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Metabolic Syndrome/Obesity (3 or more) Cardiovascular-Kidney-Metabolic Risk Factors

- **BMI:**
 - Ideal: 18.5kg/m² -25 kg/m²
 - Overweight: 25kg/m²-29.9 kg/m², Asian ≥ 23 kg/ m²
 - Obese ≥ 30.0 kg/ m²
 - May not be accurate in all patients
- **Waist circumference:**
 - Women: ≥ 88 cm, ≥ 80 cm in Asians
 - Men: ≥ 102 cm, ≥ 90 in Asians
- **Impaired glucose tolerance:**
 - Fasting blood glucose:
 - ≥100-149 mg/dL
 - HemoglobinA1c:
 - 5.7-6.4%
- **Elevated triglycerides:**
 - >150 mg/dL
 - Or currently treated
- **Low HDL:**
 - Men: <40 mg/dL
 - Women: < 50 mg/dL
- **SBP:** ≥130 mm Hg/**DBP** ≥85 mm Hg
 - Or currently treated

Cardiovascular-Kidney-Metabolic Risk Factors

- Evidence of hepatic steatosis
- Obstructive sleep apnea//sleep issues
- Smoking, alcohol

Stage 2: Metabolic Risk Factors and/or CKD

- Metabolic syndrome, Prediabetes, Type-2 DM
- **Elevated triglycerides:**
 - >150 mg/dL
 - Or currently treated
- **Low HDL:**
 - Men: <40 mg/dL
 - Women: < 50 mg/dL
- **SBP:** ≥130 mm Hg/**DBP** ≥85 mm Hg
 - Or currently treated

Stage 2: Metabolic Risk Factors and/or CKD

- **CKD:**

- Presence of **CKD** significantly **accelerates** the course of **CVDz**
- Elevated serum creatinine and/or albuminuria
 - Serum **creatinine** estimated GFR (**eGFR**) non-race based
- Spot urine **albumin/urine creatinine ratio (ACR) \geq 30 mg/dL**
- **Note: Elevated ACR** is the **earliest indicator** of diabetic and other kidney diseases, CVDz and other systemic illnesses:
 - HTN
 - Most types of cardiovascular disease: MI, stroke, PVDz
- ACR is an **early** and **significant risk factors** that should be part of a **cardiovascular risk assessment**

CKD is classified based on:
Cause (C)*
GFR (G)[†]
Albuminuria (A)[†]

			Albuminuria categories Description and range			
			A1	A2	A3	
			Normal to mildly increased	Moderately increased	Severely increased	
			<30 mg/g <3 mg/mmol	30–299 mg/g 3–29 mg/mmol	≥300 mg/g ≥30 mg/mmol	
GFR categories (mL/min per 1.73 m²) Description and range	G1	Normal or high	≥90	Screen 1	Treat 1	Treat and refer 3
	G2	Mildly decreased	60–89	Screen 1	Treat 1	Treat and refer 3
	G3a	Mildly to moderately decreased	45–59	Treat 1	Treat 2	Treat and refer 3
	G3b	Moderately to severely decreased	30–44	Treat 2	Treat and refer 3	Treat and refer 3
	G4	Severely decreased	15–29	Treat and refer [†] 3	Treat and refer [†] 3	Treat and refer 4+
	G5	Kidney failure	<15	Treat and refer 4+	Treat and refer 4+	Treat and refer 4+

Low risk (if no other markers of kidney disease, no CKD)
 High risk

Moderately increased risk

 Very high risk



Stage 3: Subclinical CVD or CKD

- **Subclinical ASCVD** or **subclinical HF** in people with **excess/dysfunctional adiposity:**
- **Subclinical HF:**
- **Subclinical ASCVD Risk Equivalent:**
 - CKD Stages G2 - G3b
 - High risk: G4-G5
 - High predicted all cause CVDz

Stage 3: Subclinical CVD or CKD

- **Subclinical ASCVD** or **subclinical HF** in people with **adiposity**:
 - CT coronary calcium score (CAC)
 - CT angiography
- **Subclinical HF**:
 - **Elevated NT-proBNP**: >125 pg/mL*
 - **Echocardiographic evidence**:
 - Atrial enlargement(s), impaired relaxation, systolic dysfunction
 - **hs-troponin I**:
 - Women: ≥ 10 ng/mL
 - Men: ≥ 12 ng/mL
 - **hs-troponin T**:
 - Women: ≥ 14 ng/mL
 - Men: ≥ 22 ng/mL Hs-troponin I

***Note:**

•CKD, age, female sex, atrial fibrillation, inflammation, hyperthyroidism, sacubitril/valsartan **overestimate** BNP level.

•Obesity, Early MI, pericardial effusion **underestimate** BNP level

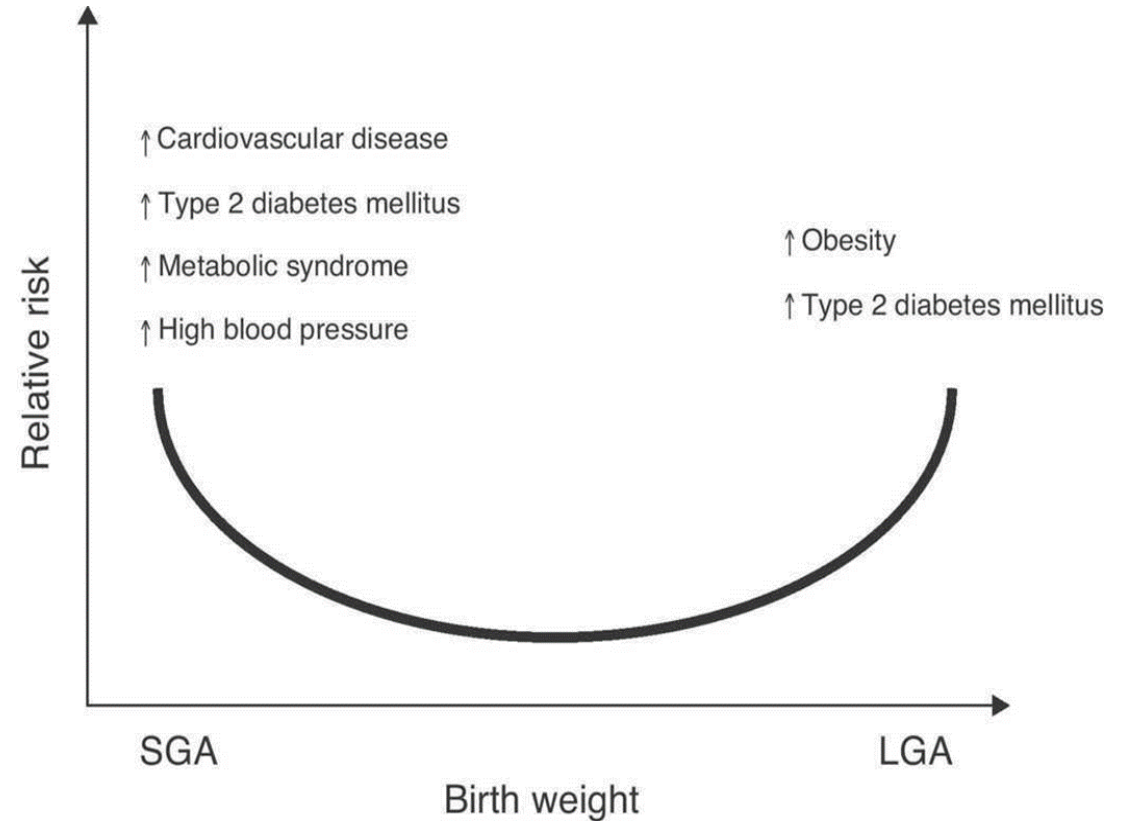
Journal of Cardiology. 2021; 78: 269–274

Stage 4: Clinical CVD in CKM

- **Clinical CVD** in people with **excess/dysfunctional adiposity**
- **Clinical CVD:**
 - Coronary artery disease, HF, stroke, peripheral vascular disease, atrial fibrillation
- **Other CKM risk factors:**
 - **CKM Stage 4a:** no kidney failure
 - **CKM Stage 4b:** kidney failure present (CKD Stage G2-5)

Risk for Type 2 Diabetes Mellitus

- Co-existing **maternal diabetes** (preexisting or gestational)
 - increases risk in child
- **Prematurity**: increased risk
- **SGA**: small for gestational age:
 - <10th percentile for age and weight
- **LGA**: large for gestational age:
 - >4.0 kg



Diabetes Burden: Prevalence by Ethnicity

- **Native American/Alaska Native:** 14.5%
 - Women: 14.8%
- **Black, non-Hispanic:** 12.1%
- **Hispanic:** 11.8 %
 - Mexican 14.4%
 - Puerto Rican: 12.4%
 - Central/South America 8.3%
 - Cuban: 6.5%
- **Asian:** 9.5%
 - Indian subcontinent: 12.6%
 - Filipinos 10.4%
 - Chinese: 5.6%
 - Other Asian: 9.9%
- **White non-Hispanic:** 7.4%
- **Education:**
 - **Less than H.S. education: 13.4%**
 - H.S. education: 9.2%
 - More than H.S. education: 7.1%
- **Family income** (all groups)
 - **Below federal poverty level: 14.1%**

Screening for Prediabetes

- **High risk:**

- Birthweight
- Fam hx/maternal/gestational DM, 1st degree relatives
 - Hx GDM: eval minimum q 3 years, weight
- Age > 35
- BMI ≥ 25 kg/ m²
 - Asian American ≥ 23 kg/ m²
 - Poor diet, sedentary
- Obesity, presence of acanthosis nigricans, HTN, hyperlipidemia
- Polycystic ovary syndrome (PCOS)
- HIV Hx/Tx
- **Ethnicity:** Native American, PI, Alaska native, AA, Asian American
- Role of poverty/education levels
- Medication associated

Screening for Prediabetes/ T2DM in Adolescents and Teens

- Onset of puberty or >10 y.o.
 - Younger?
- High risk groups
- **Overweight: BMI \geq 85th percentile**
- **Obesity: BMI \geq 95th percentile**
- FBS, 2 hour post prandial 75 gm, Hb A1c
- Also get Ab to r/o T1DM and r/o MODY

Criteria for Prediabetes

- **Fasting glucose:**
 - 100-124 mg/dLor
- **2-hour glucose after 75 gm glucose load:**
 - 140-199 mg/dLor
- **HbA1c:**
 - 5.7-6.4%

2023 ADA Standards: [Volume 46 Issue Supplement 1 | Diabetes Care | American Diabetes Association \(diabetesjournals.org\)](https://doi.org/10.2337/141101)

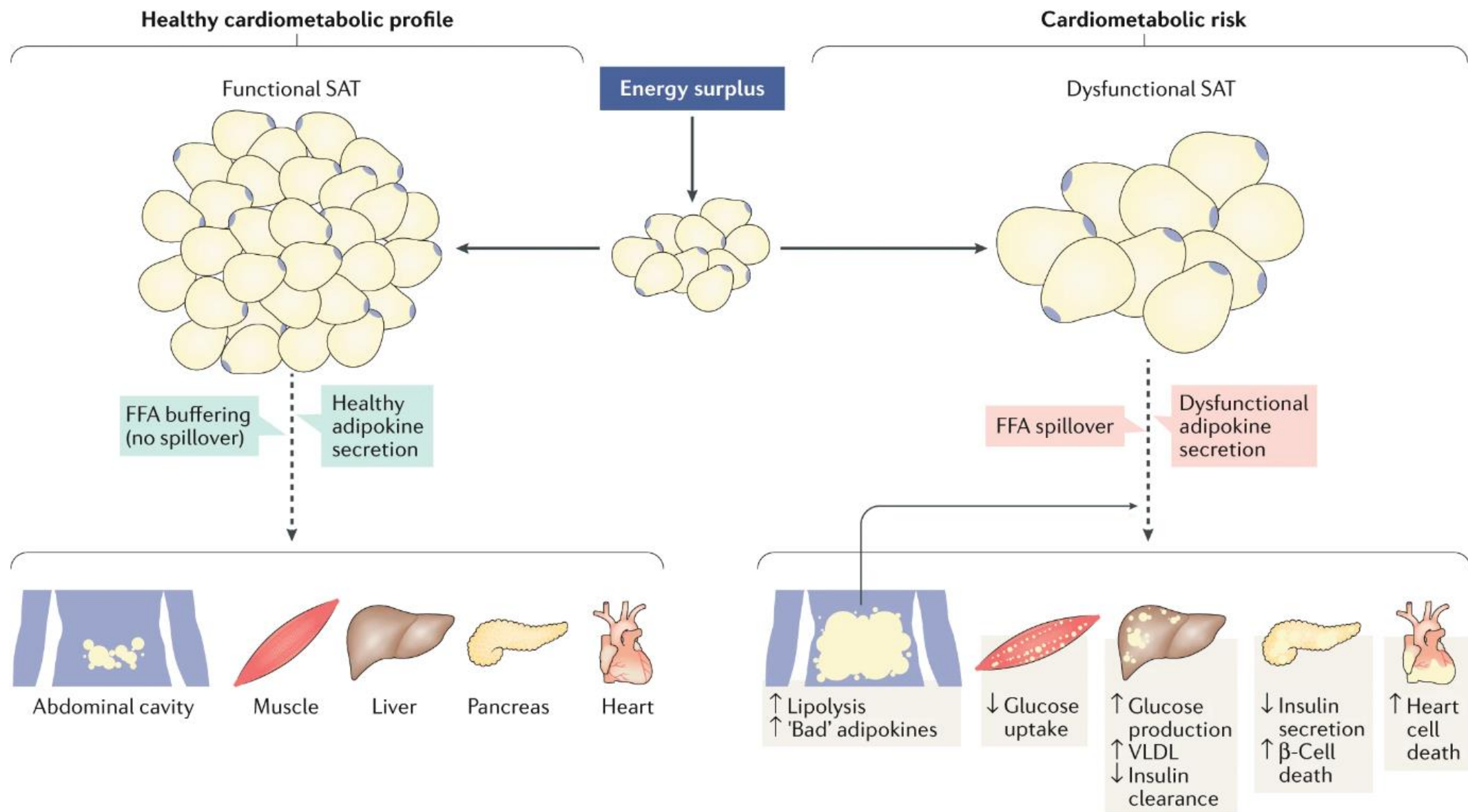
Diagnosis of Type-2 Diabetes Mellitus

- **Fasting plasma glucose:** ≥ 126 mg/dL
- **2-hour plasma glucose:** ≥ 200 mg/dL following a **75 gm oral glucose** tolerance test
- **Hemoglobin A1c:** $\geq 6.5\%$

Type 2 Diabetes

- **Type 2**: insulin resistance
 - normal or elevated insulin levels with elevated glucose
- Directly related to obesity
- Alteration of **multiple neuroendocrine systems**
 - SNS, RAAS, glucagon, cortisol, growth hormone, etc.
- **~77% incidence** of DN when retinopathy present
- **Quiescent** period of 10-15 years may be masked as end organ(s) damage is frequently present at time of presentation of DM

Parving *Kidney Int.* 1992;41:758.



Screening for NAFLD/MAFLD

- **Infiltration of fatty acids** into **liver** leading to **inflammation & fibrosis**
- **Truncal/visceral/abdominal obesity**
- **Most common cause** of **51%** of **chronic liver disease/cirrhosis and liver transplantation**
- **2016 meta-analysis** showed a **prevalence of 25.24%**
- **NAFLD** predicted **risk** of overt **Type-2 DM in prediabetes:**
 - **6.9 fold** increase risk in **men**
 - **5.8 fold** increase risk in **women**

Screening for NAFLD/MAFLD

- Right upper quadrant/hepatic ultrasound
- **AST:ALT** elevated with ratio of **2:1** ($\geq 2X$ normal)
- Elevated gamma glutamyl transferase (**GGT**)
- Occasionally elevated alkaline phosphatase (**AP**)
- **NAFLD/MAFLD & risk of hepatocellular carcinoma:**
 - **2.3%** in **7 years**
 - Up to **12.3 %** in **3 years** (Clin Gastroenterol Hepatol 2012;10:1342.)
- Rule out **Hepatitis B, C**

AFP. 2013;42, 444-447 [Int J Biol Sci](#). 2019; 15(3): 610–616.

Multinodular Hepatic Steatosis



Approach to CKM

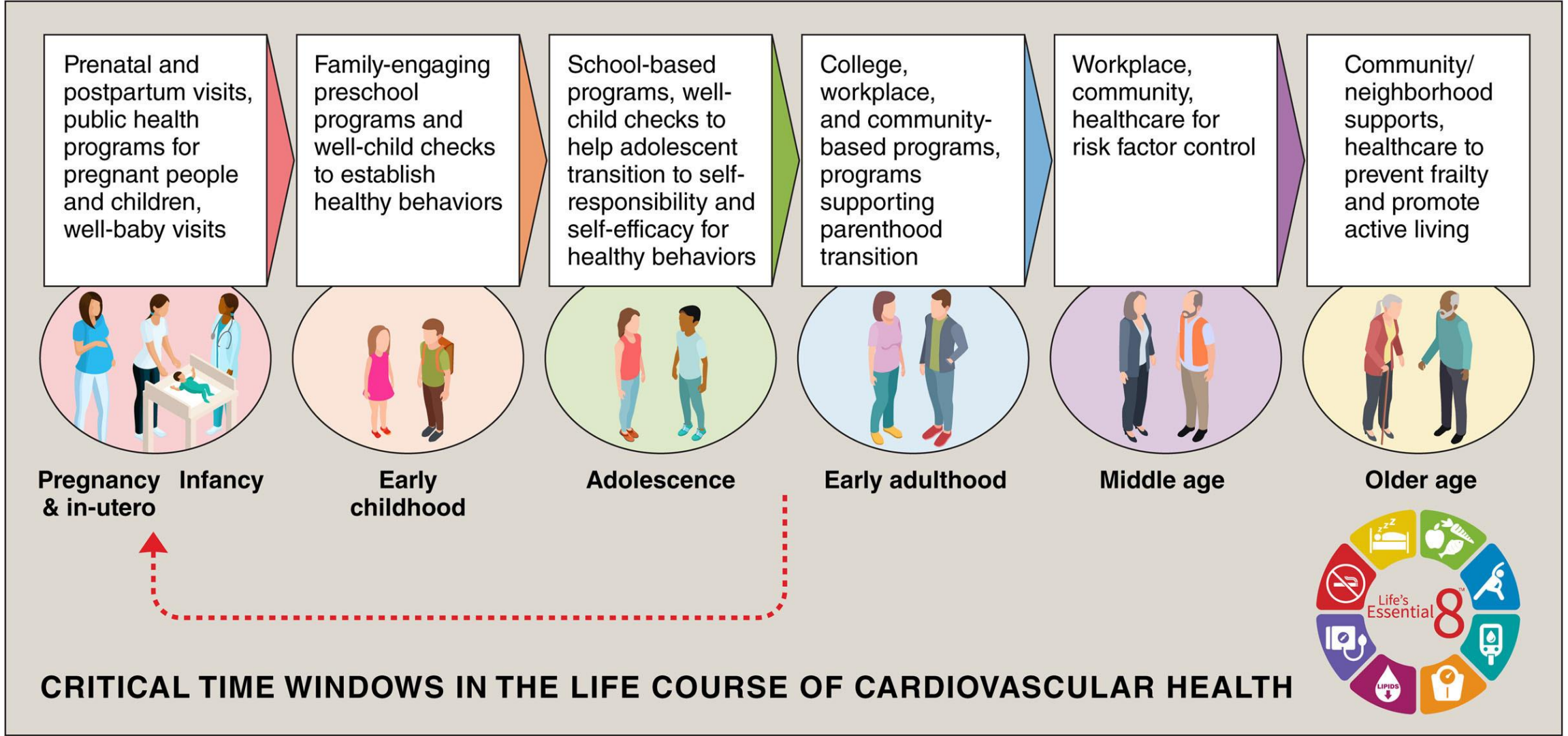
- Early recognition and intervention is key
- Access to health care, education, healthy food, activities

Life's Essential 8

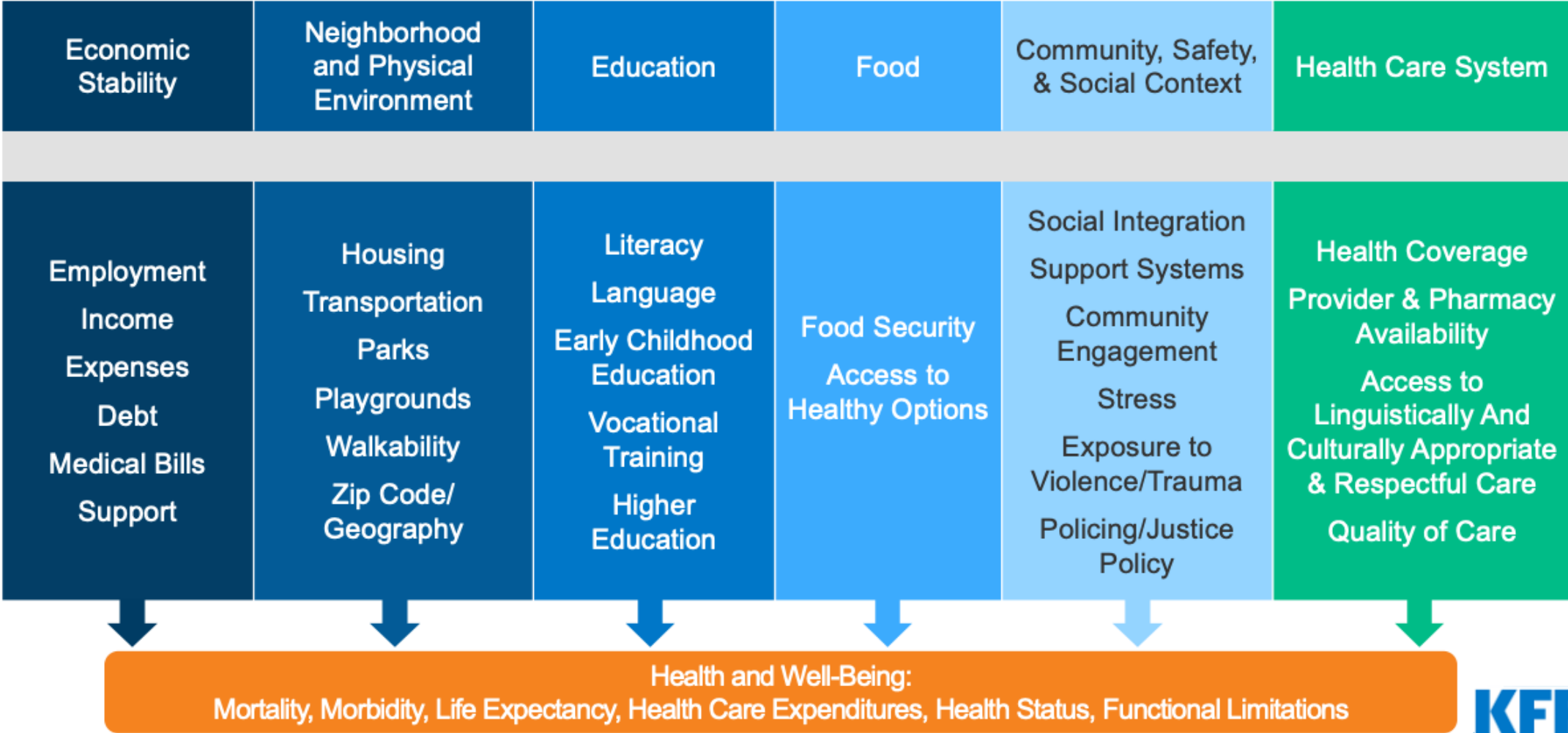
- **Life's Essential 8** includes the 8 components of **cardiovascular health**:
 - Healthy diet
 - Physical activity
 - Avoidance of nicotine (ETOH)
 - Healthy sleep
 - Healthy weight
 - Healthy levels of blood lipids
 - Normal blood glucose
 - Normal blood pressure.



Early (*real early*) Identification and intervention



Social Determinants of Health



Chronic Care Model

- Primary focus on **cardiorenal risk**
- Education
- Addressing “environmental” issues facing the patient
 - Social services/advocacy to address the SDOH
 - Adequate housing, safe neighborhoods
 - Nutritional needs/access
 - Access to healthcare, medication, follow up, healthcare coverage, mental health care

Chronic Care Model

- Comprehensive team based approach
- Physician, CNP, PA-C,
- Nurse manager
- Dietician/Certified diabetic educator
- Social worker
- Mental health facilitator
- Optometrist/ophthalmology
- Podiatry
- Nephrology
- Cardiology
- Endocrinology

Stage 0/1 CKM: Interventions

- Identification of **high risk groups**
- **Promotion of CV health:**
 - BMI/ waist circumference
 - Glucose
 - BP
- **Systematic screening for the SODH/challenges**
- **Intensive lifestyle interventions:**
 - Diet, activity, smoking, ETOH cessation
- **Pharmacotherapies, BMI: ≥ 30 kg/m**
 - Metformin, GLP-1, SGLT-2
- **Bariatric surgery, BMI: ≥ 40 kg/m (w/o comorbidities)**

Stage 2 CKM: Interventions

- **Hyperlipidemia:**

- Maximize statin therapy
- Triglycerides ≥ 500 mg/dL: fibrates
- Glucose control

- **Hypertension:**

- Follow guidelines JNC-8, ACC, KIDIGO, ADA, etc.
- **BP:** $<130/85$ mm Hg (SPRINT $<120/80$ mm Hg)
- Any evidence of **CKD/ albuminuria:** **ACEi/ARB**

Stage 2 CKM: Interventions

- **CKD:**

- Any type
- **Elevated serum creatinine** (n 0.6-1.2 mg/dL)
or
- Urinary albumin/creatinine ratio \geq 30 mg/dL
- ACR $>$ 30 mg/m on ACEi/ARB tx
 - Add spironolactone, eplerenone, finerenone, SGLT-2

- **Diabetes:**

- Intense **statin** therapy
- **Metformin**
- **RAAS inhibition: ACE-i/ARB**
- **SGLT-2**
- **Co-morbidities (CVD/CKD)**
 - **BMI:** $>$ 35 kg/m² -**GLP-1 RA**
 - **HbA1c:** $>$ 9% or high dose insulin-**GLP-1 RA**
 - **CKD/DKD:** **SGLT-2, GLP-1**
- **ACR:** $>$ 30 mg/m² on **ACEi/ARB tx**
 - Spironolactone, eplerenone, finerenone

Stage 3: Interventions

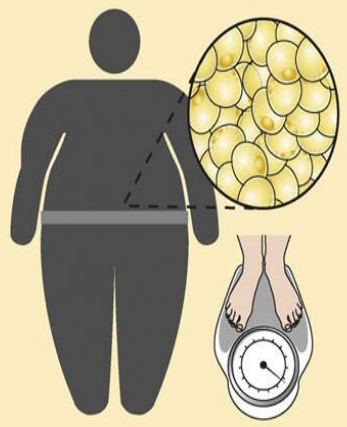
- **Subclinical Atherosclerosis:**
 - **Coronary calcium score (CAC)**
 - **CAC: >0**
 - **CAC: >100**
 - High dose statin, low dose ASA, PCSH9i, GLP-1RA
- **Subclinical HF:**
 - **EF: <40%**
 - ACEi/ARB
 - SGLT-2, esp. in DM
- **CVD risk equivalent for Stage 3 CKM:**
 - **Very high risk: CKD Stage G4, 5 or ESKD**
- **Low dose aspirin:**

**Stage 0:
No Risk Factors**



A focus on primordial prevention and preserving cardiovascular health

**Stage 1:
Excess/Dysfunctional Adipose Tissue**

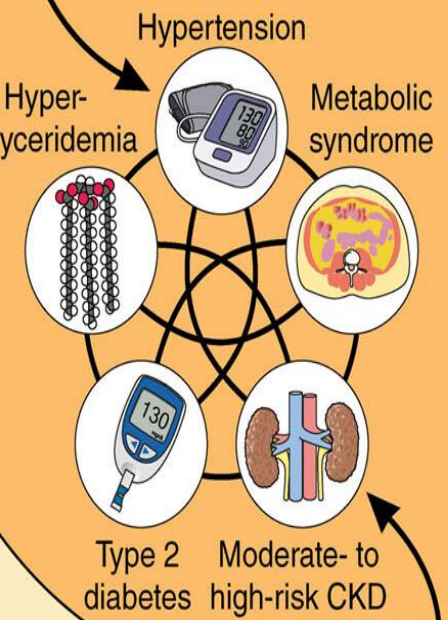


Overweight/obesity
Abdominal obesity
Impaired glucose tolerance

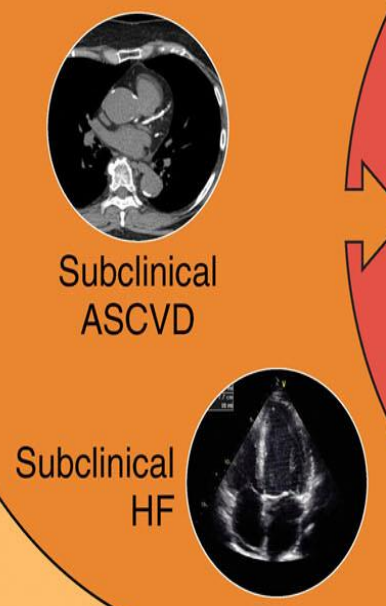
Nonmetabolic etiologies of hypertension

Nonmetabolic etiologies of CKD

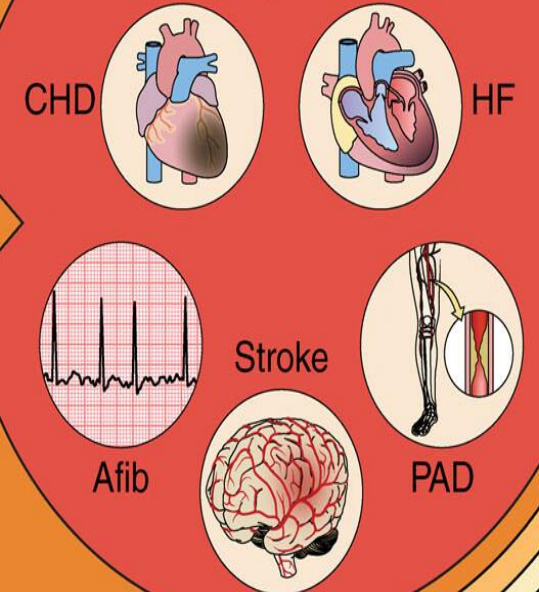
**Stage 2:
Metabolic Risk Factors and CKD**



**Stage 3:
Subclinical CVD in CKM Syndrome**



**Stage 4:
Clinical CVD in CKM Syndrome**



Risk equivalents of subclinical CVD in CKM Stage 3:

- Very high-risk CKD (G stage 4 and 5 CKD or by KDIGO heat map)
- High predicted risk for CVD using risk calculator

Questions?

Thank you

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