

# Metabolic Health

Cardiovascular and Wider Impacts

Dr. Tro Kalayjian, D.O.

April 2026

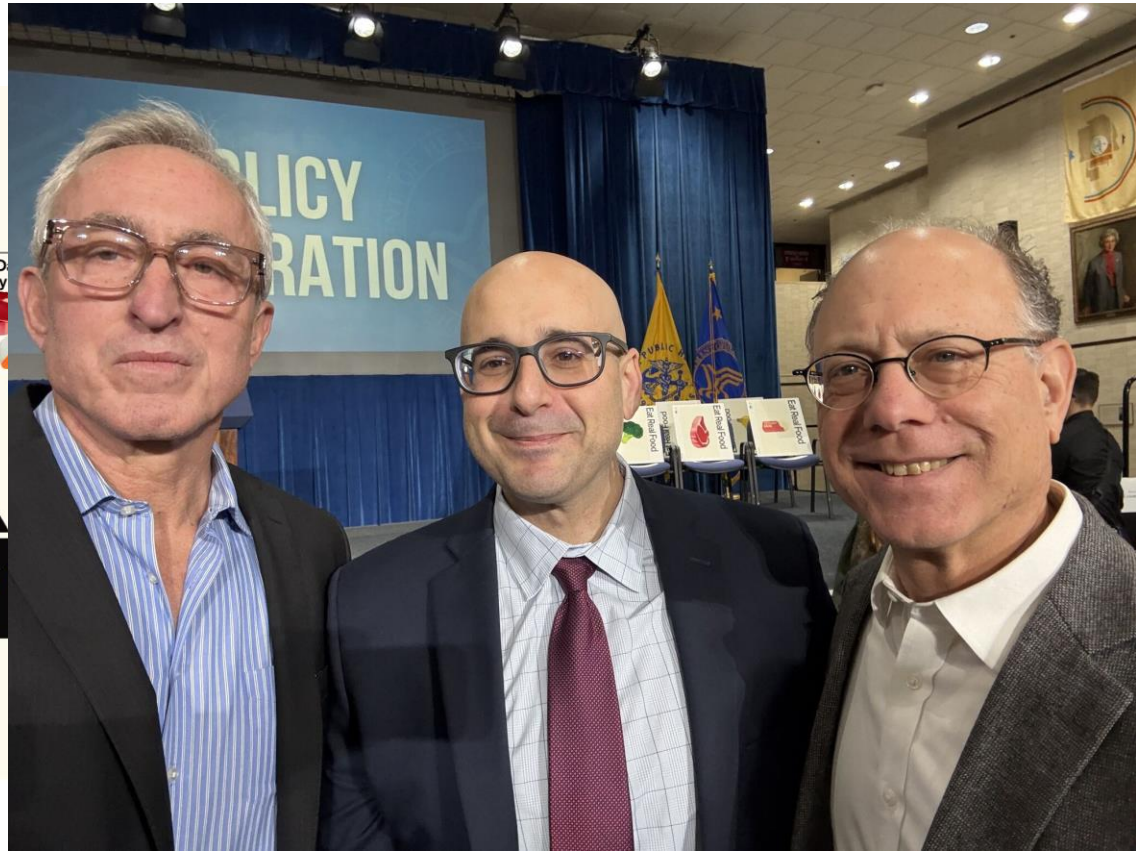
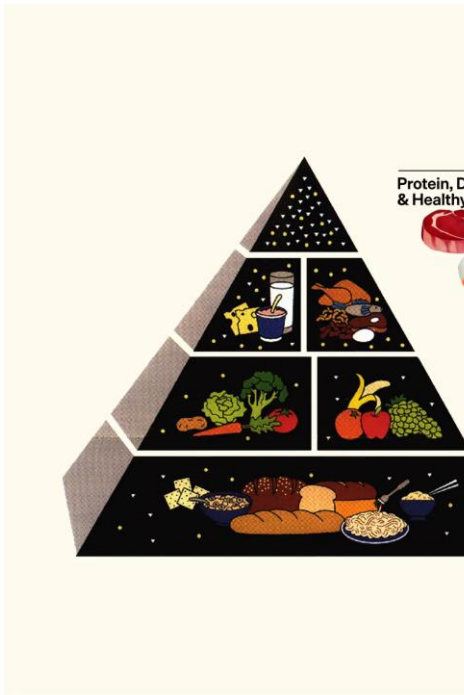


# Who am I?

- Board certified Internal Medicine Doctor
- Board certified Obesity Medicine Doctor
- Certified Personal Trainer, Health Coach
- Nationwide Virtual Obesity & Primary Care Practice
- Board Member of SMHP
- Scientific Publication on Metabolic Health & Regulatory Capture (NLA, APA, AAP, ADA, AHA)
- Personally lost 150lbs
- 2,000 patients, helped lose over 100,000lbs
- Smartphone app with 17k members
- Podcast with 12 million downloads
- 3 ongoing IRB projects with Stanford, Yale New Haven and Wake Forest
- Baszucki Grant Recipient (project to be announced)
- Father of 3 amazing children



**Thanks to the Swiss Re, SMHP, Metabolic Coalition,  
Nina Teicholz, Ben Bikman, and so many others...**



# Controversy with the American Heart Association...

THE WALL STREET JOURNAL.  
Business U.S. Politics Economy Tech Markets & Finance Opinion Free Expression Arts


HEALTH | WELLNESS

## Heart Association Clashes With RFK Jr. Over Red Meat, Dairy and Beef Tallow

The American Heart Association recommends getting protein from plants—rather than meat—and avoiding full-fat dairy

By [Andrea Petersen](#) and [Betsy McKay](#)  
Updated March 31, 2026 6:53 am ET

Share Resize 647 Listen (2 min)




The American Heart Association report cites research showing that diets higher in beans, peas and lentils—and lower in red and processed meat—are associated with a lower risk of heart disease. GETTY IMAGES


the basis for an updated scientific statement reflecting the latest nutrition guidance published today in **Circulation**, the peer-reviewed, flagship journal of the American Heart Association, a global force changing the future of health for all.

The *2026 Dietary Guidance to Improve Cardiovascular Health: A Scientific Statement From the American Heart Association* updates the Association's 2021 guidance with the latest in evidence-based science aligned with reducing cardiovascular disease risk, improving quality of life and saving lives. The statement outlines nine key features of a heart-healthy dietary pattern:

- Adjust energy intake and expenditure to achieve and maintain a healthy body weight:** Try to balance how much you eat with how active you are, to reach and maintain a healthy body weight.
- Eat plenty of vegetables and fruits and choose a wide variety:** Include different colors, textures and types of produce, and remember, even canned and frozen can be nutritious and affordable.
- Choose foods made mostly with whole grains rather than refined grains:** Foods such as whole-wheat bread, brown rice and oatmeal are better choices than refined grains including white bread or white rice.
- Choose healthy sources of protein:** Shift from meat to plant-based sources such as legumes, including beans, peas and lentils, along with nuts and seeds; regularly consume fish and seafood; select low-fat or fat-free dairy products; and if red meat is desired, choose lean cuts, avoid processed forms and limit portion size.
- Choose sources of unsaturated fats in place of sources of saturated fat:** Replace saturated fats with healthy unsaturated fats, including those from nuts, seeds, avocados and nontropical plant oils.
- Choose minimally processed foods instead of ultraprocessed foods:** Go with foods close to their natural state, with minimal added commercial ingredients, rather than those that are highly processed with additives.
- Minimize intake of added sugars in beverages and foods:** Limit the sugar-sweetened beverages you drink and the foods with added sugar you eat.
- Choose foods low in sodium and prepare foods with minimal or no salt:** Be aware of hidden sources of sodium in commercially prepared and packaged foods and season your food with healthier options such as herbs, spices or lemon instead of salt.
- If alcohol is not consumed, do not start; if alcohol is consumed, limit intake:** Alcohol can increase your risk for high blood pressure and other health conditions, so if you don't drink, don't start.



Related Video




Dr. Khera on main updates in new dietary guidance

Amit Khera, M.D., FAHA, volunteer vice-chair of the dietary guidance writing committee and director of preventive cardiology and clinical chief of cardiology at the University of Texas (UT) Southwestern Medical Center.

News release

copyright American Heart Association

Download (73.6 MB)



Failed Food Pyramid vs. New Food Pyramid vs. AHA nutritional guidelines

**My meals look like this... what should I do?**

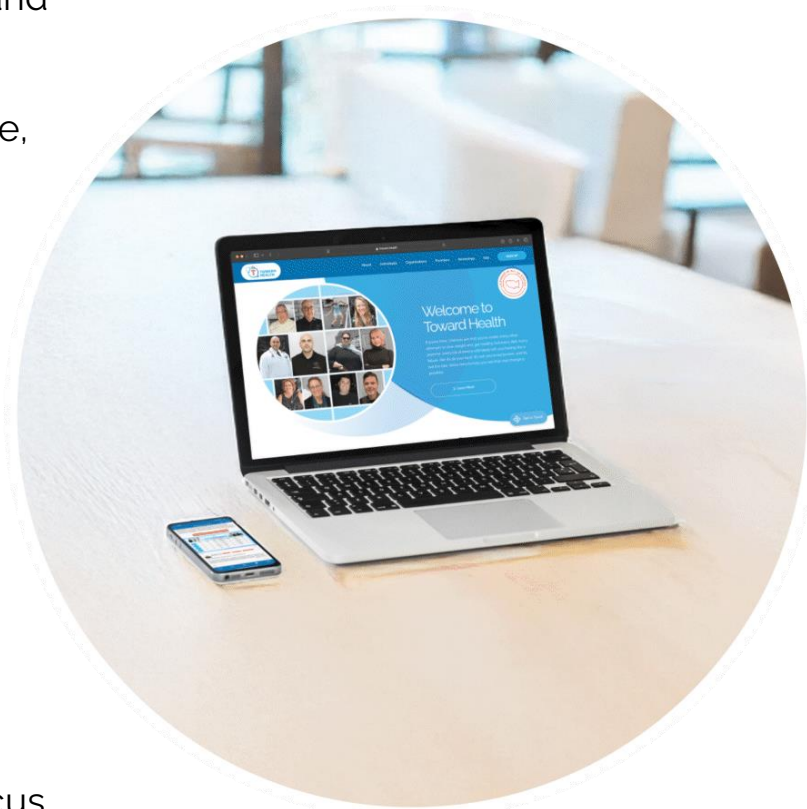
---



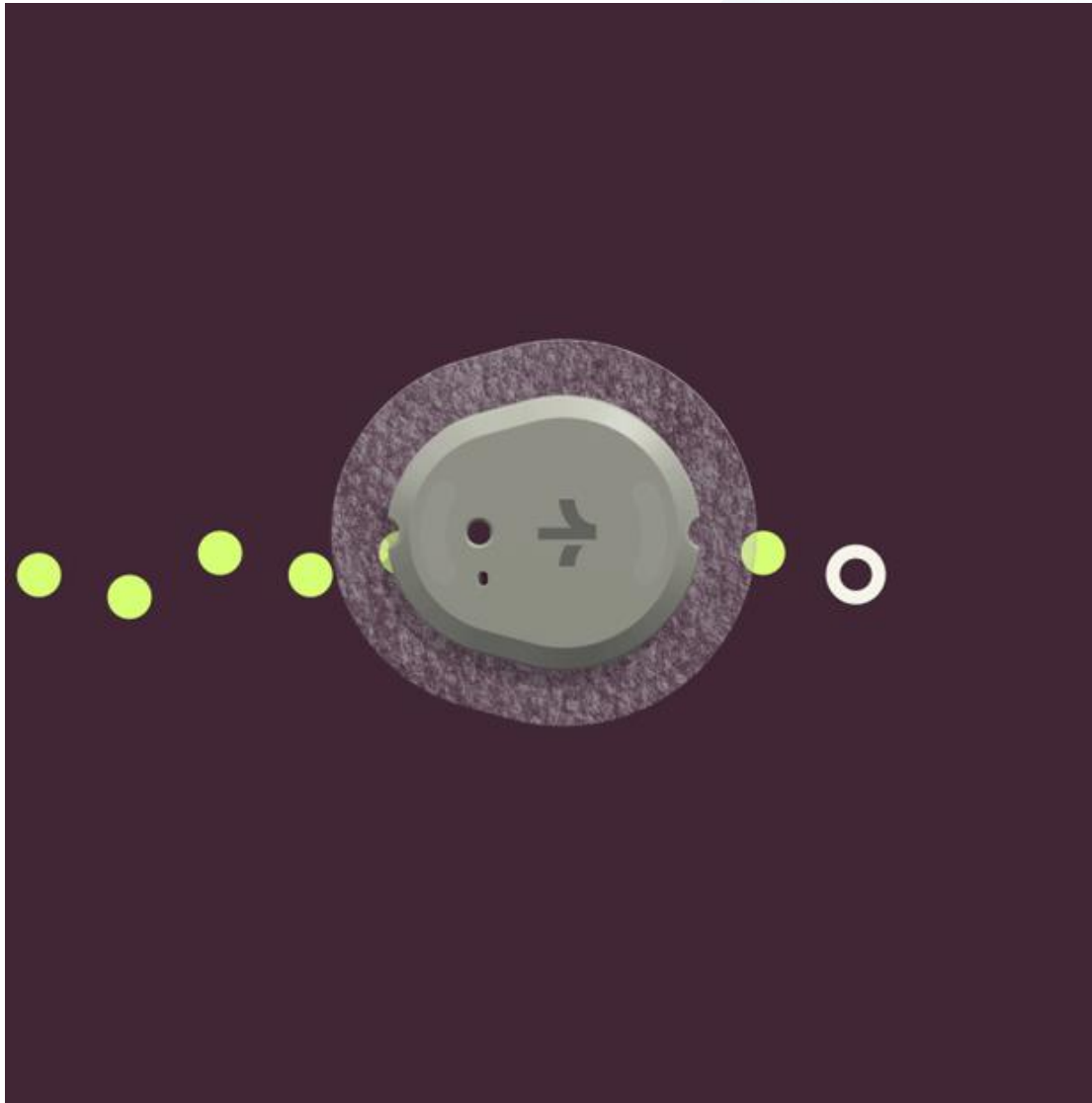
# A Proven Model for Metabolic Health

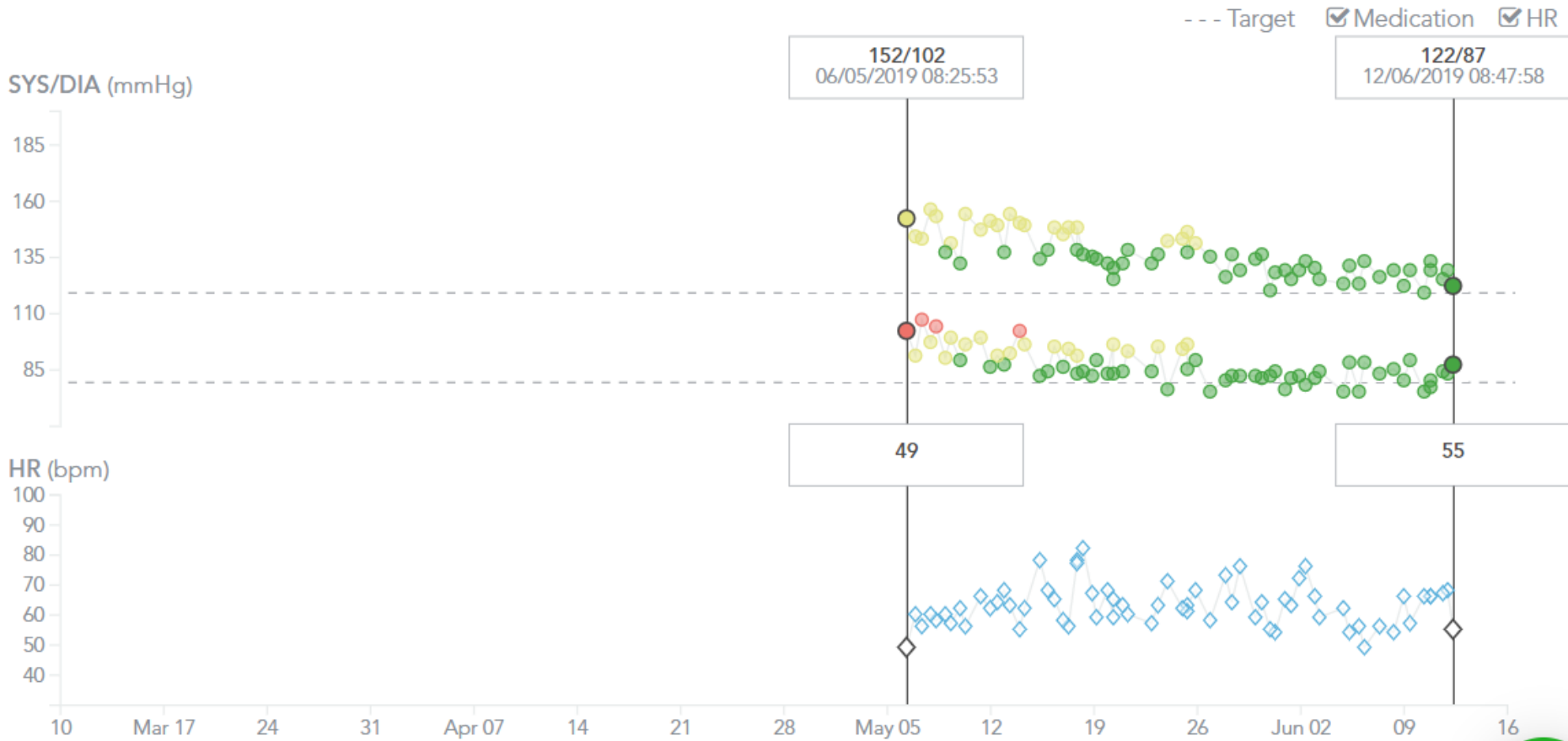
The TOWARD approach prioritizes **personalized, real-time metabolic care** that empowers patients to take control of their health. This model is built on six core principles:

- T Text-Based Communications** – Regular touchpoints and support via messaging keep patients engaged and accountable.
- O Online Interactions** – Virtual visits provide accessible, convenient care without the need for in-person appointments.
- W Wellness Coaching** – Expert health coaches guide patients through sustainable lifestyle changes.
- A Asynchronous Education & Community Support** – Self-paced learning and peer support foster long-term success.
- R Real-Time Biofeedback & Remote Monitoring** – Continuous tracking of metabolic health metrics enables timely adjustments.
- D Dietary Modifications Emphasizing TCR & IF** – A focus on **Therapeutic Carbohydrate Reduction (TCR)** and **Intermittent Fasting (IF)** supports metabolic flexibility and sustainable weight loss.




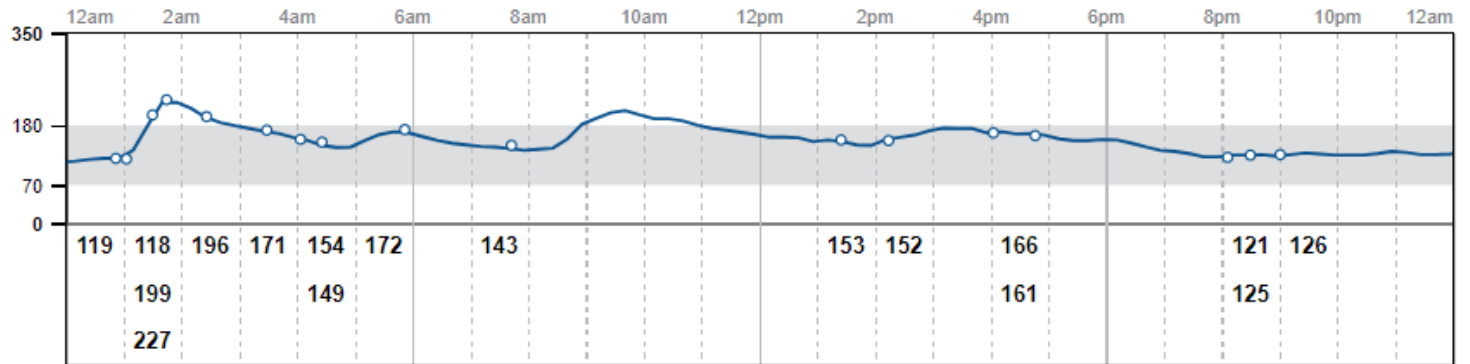
# Leveraging technology to improve medical outcomes






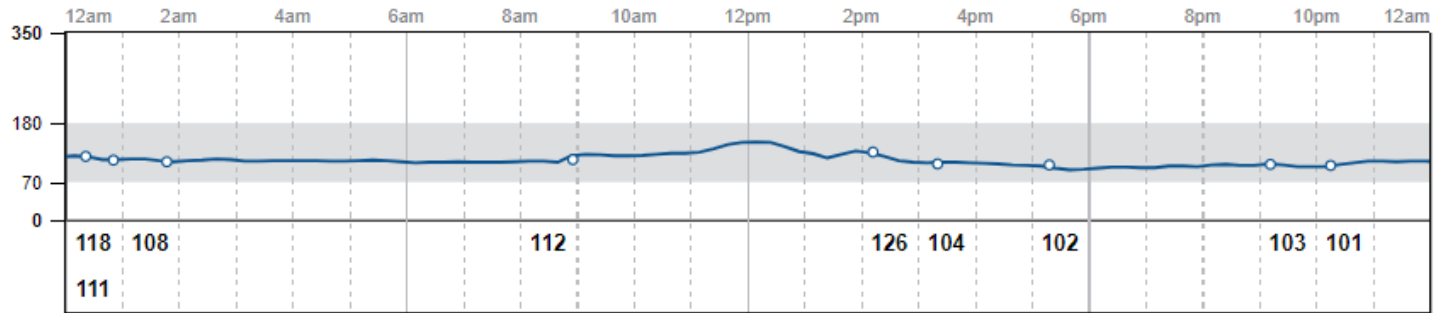
## SUN May 26

 Glucose mg/dL

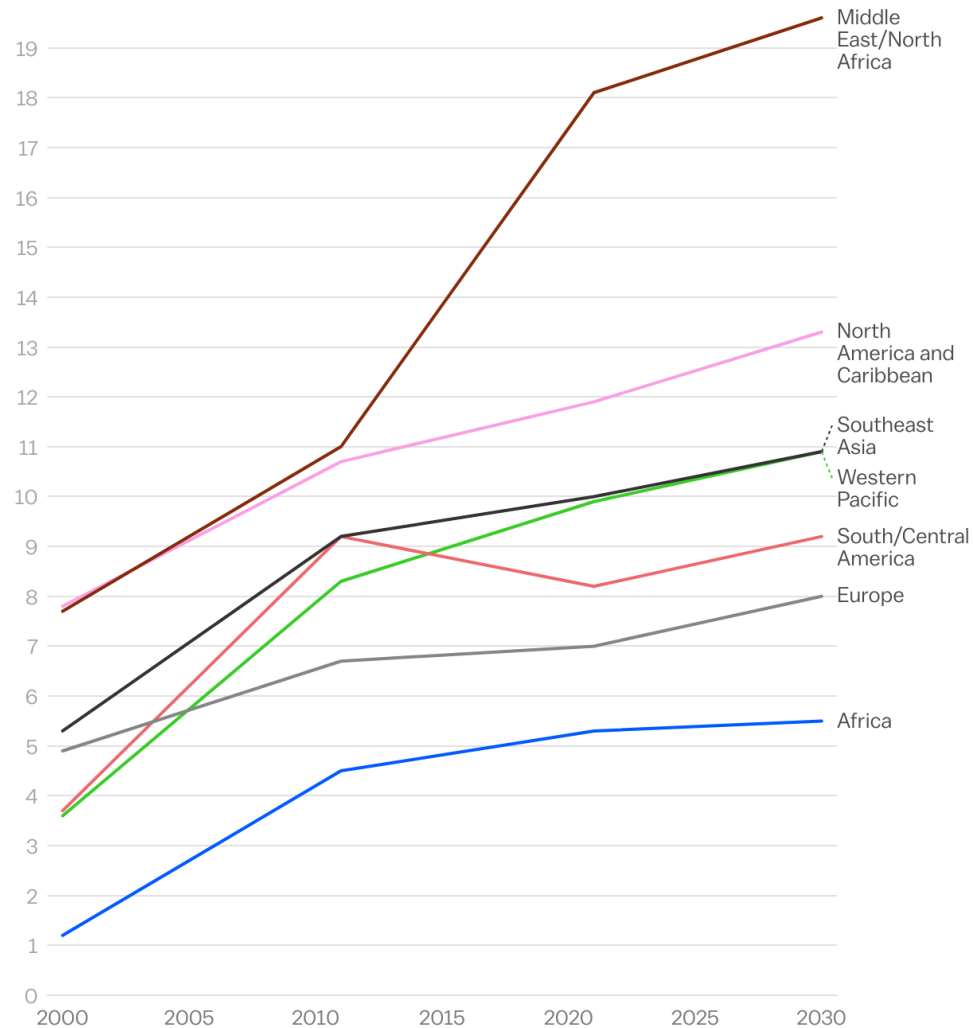


## SUN Jun 2

 Glucose mg/dL



# Ignoring root causes leads to metabolic disease...

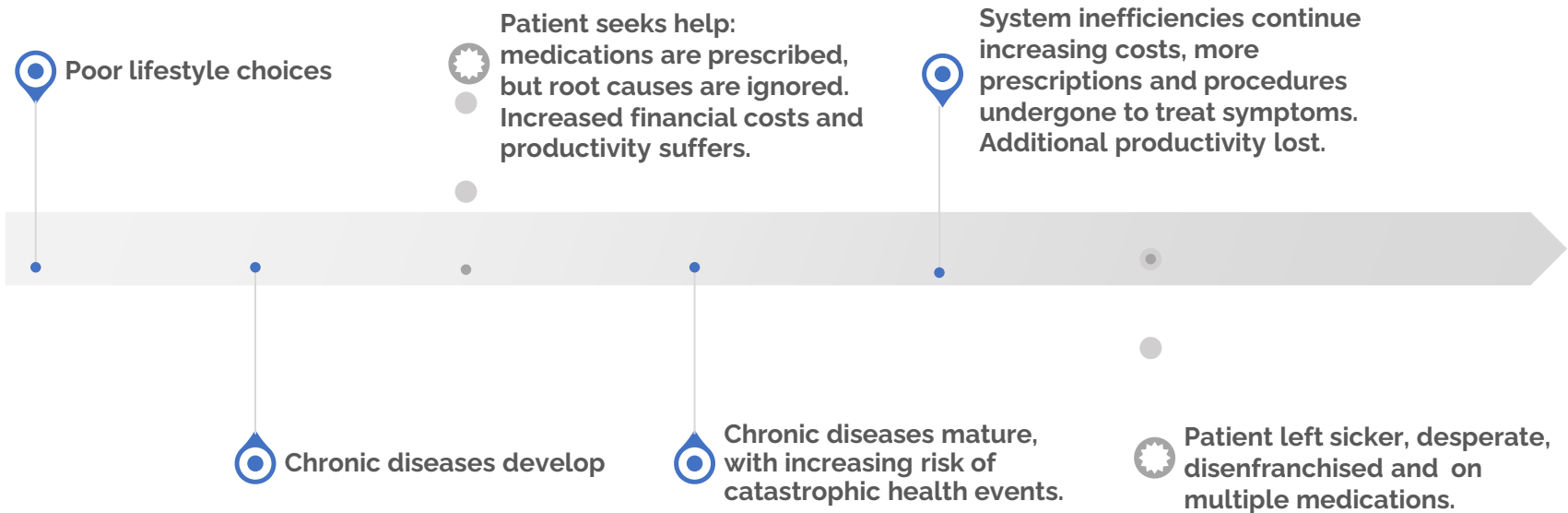


# The Status Quo

---

The current healthcare model is **expensive & does not prevent or reverse chronic disease**

---



# The Problem

My own **case study** to demonstrate the problem:

Poor lifestyle choices



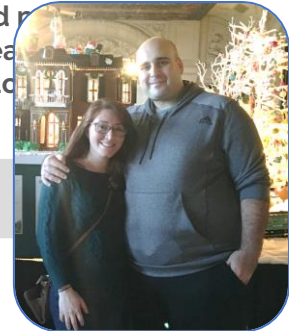
Chronic diseases develop

Patient seeks help:  
medications are prescribed,  
but root causes are ignored.  
Increased  
prod



Chronic diseases mature,  
with increasing risk of  
catastrophic health events.

System inefficiencies continue  
increasing costs, more  
prescriptions and  
undergone to treat  
Additional products



Patient left sicker, desperate,  
disenfranchised and on  
multiple medications.



# Case #1

---



Genetics, the obesogenic environment and thermodynamics didn't change...

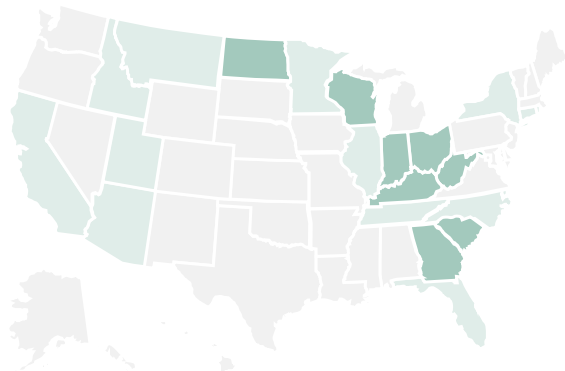
**What changed?**



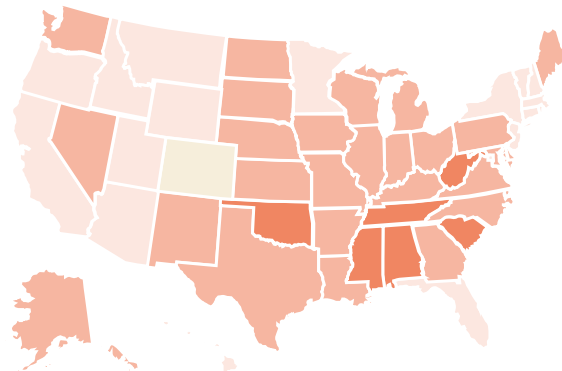
# Let's be detectives together...

## What changed?

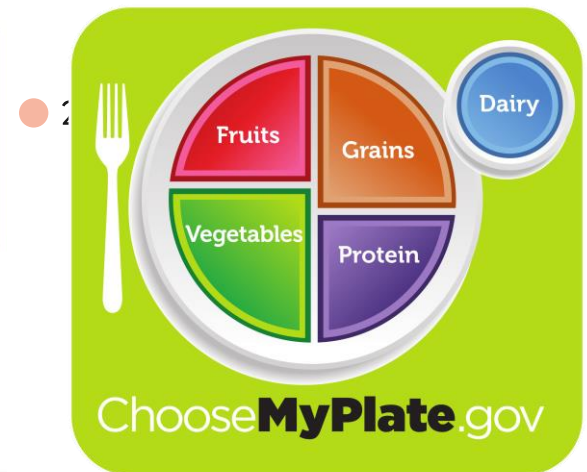
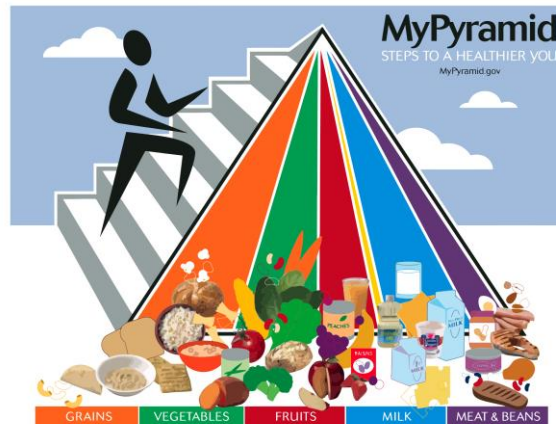
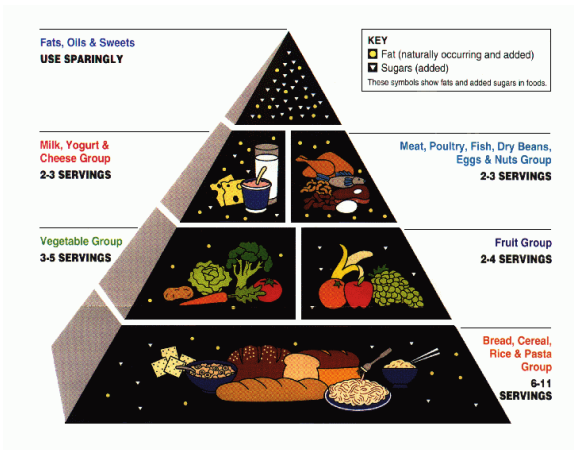
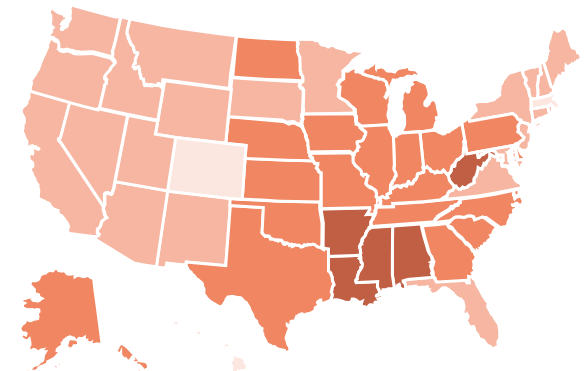
1985



2008

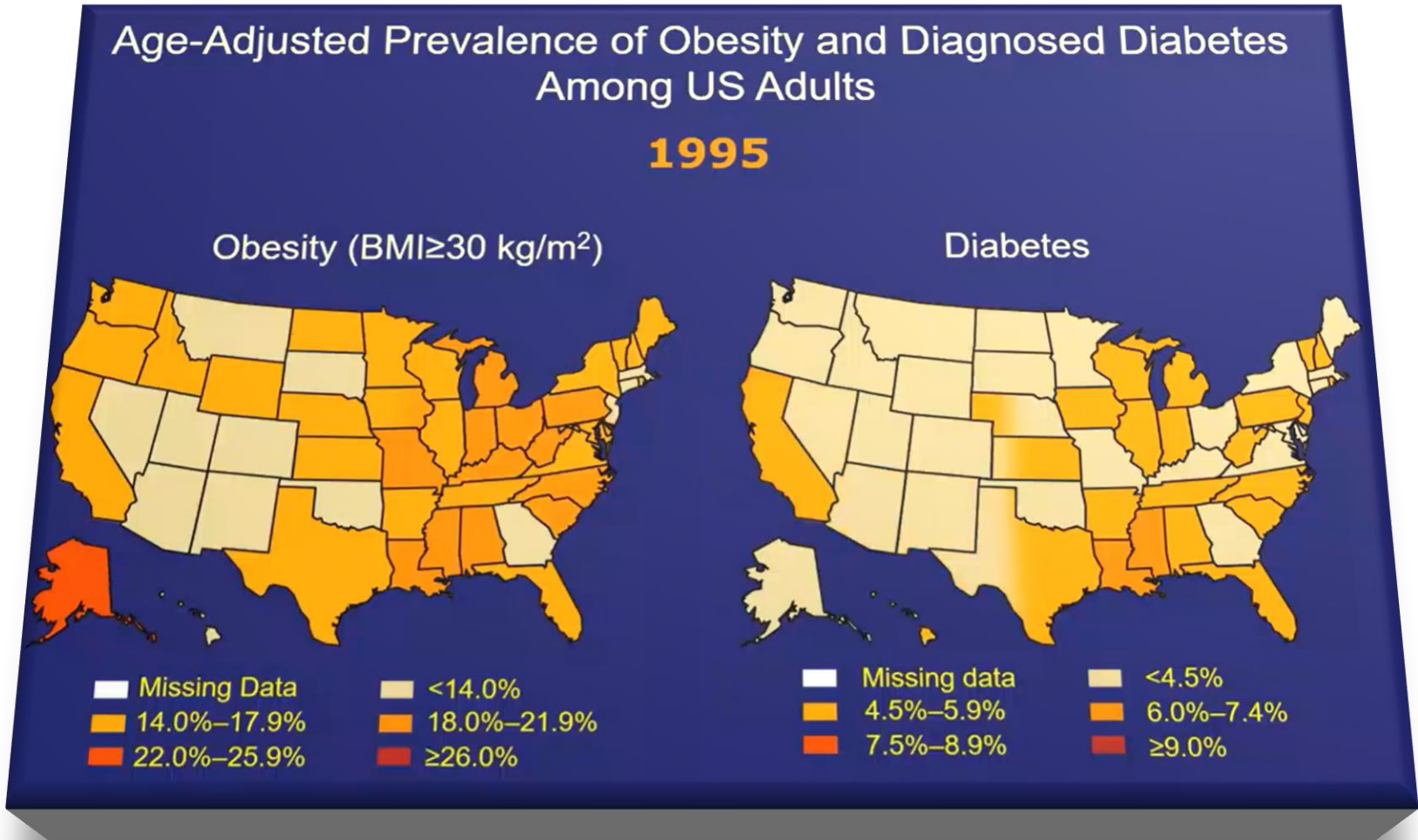


2016



# Let's be detectives together...

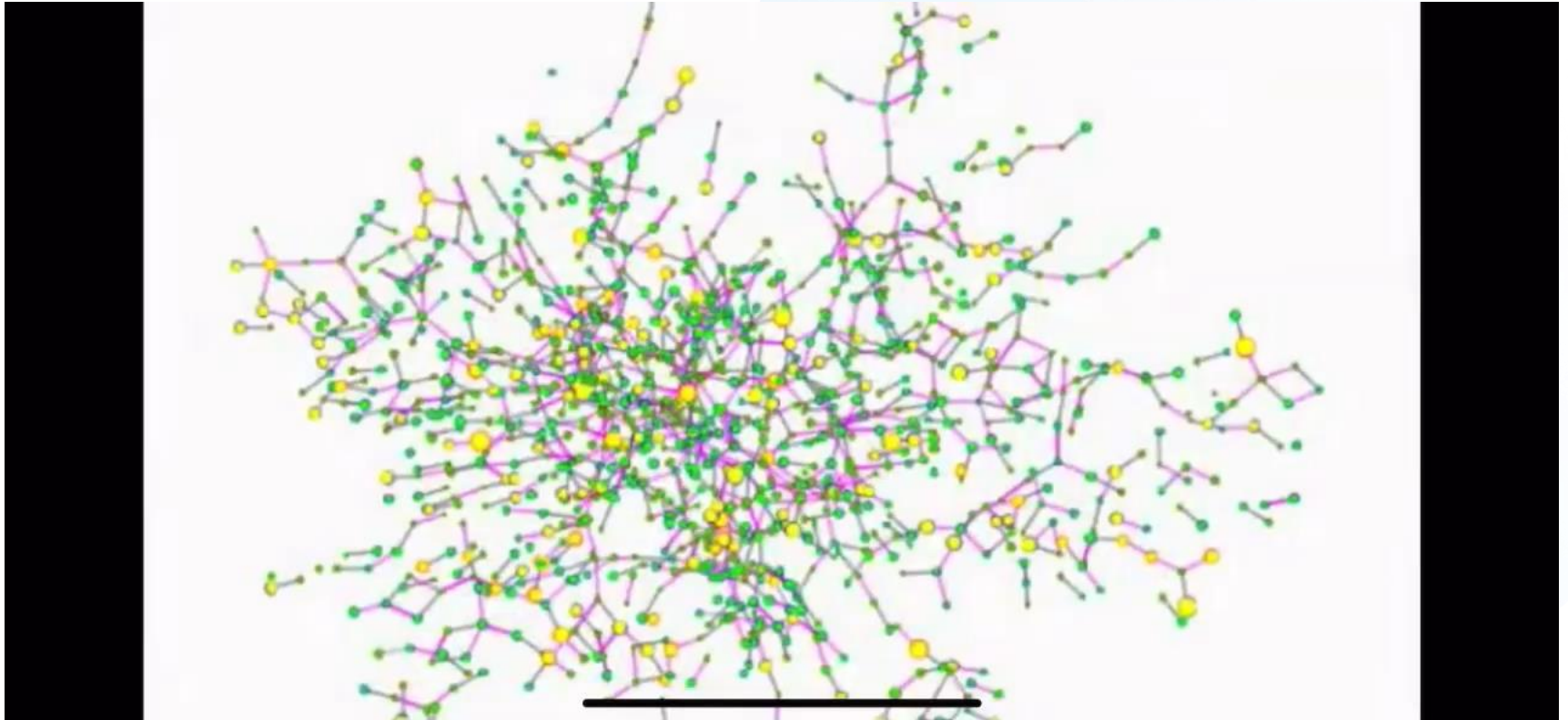
## What changed?



Let's be detectives together...

---

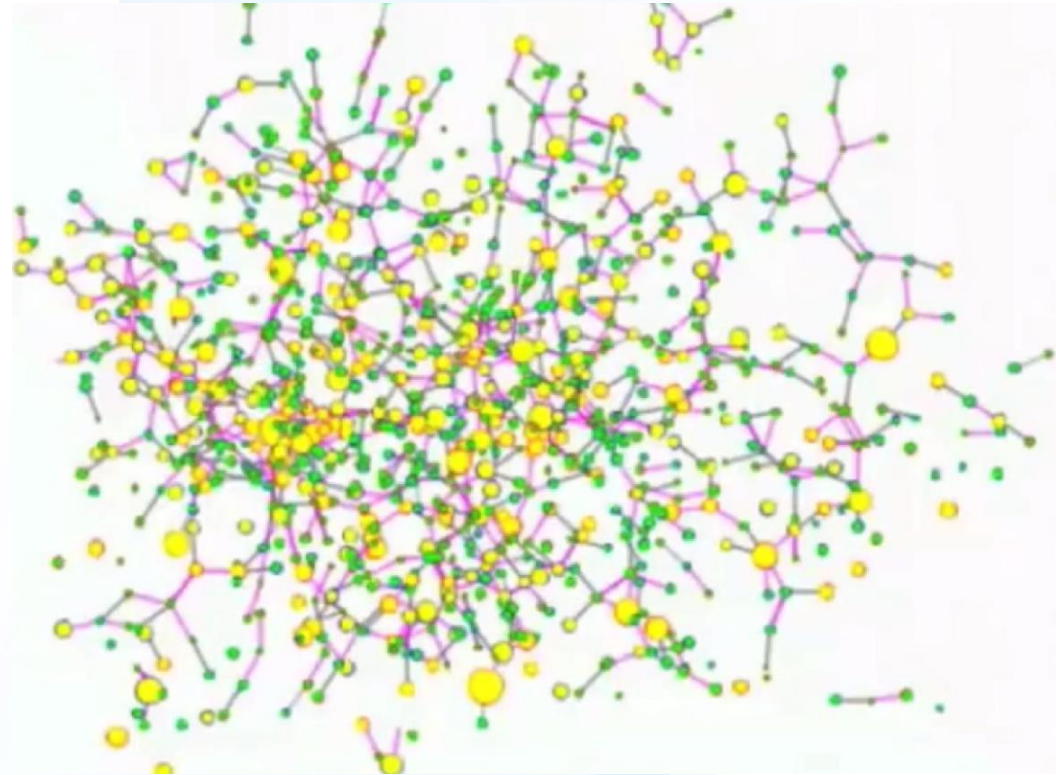
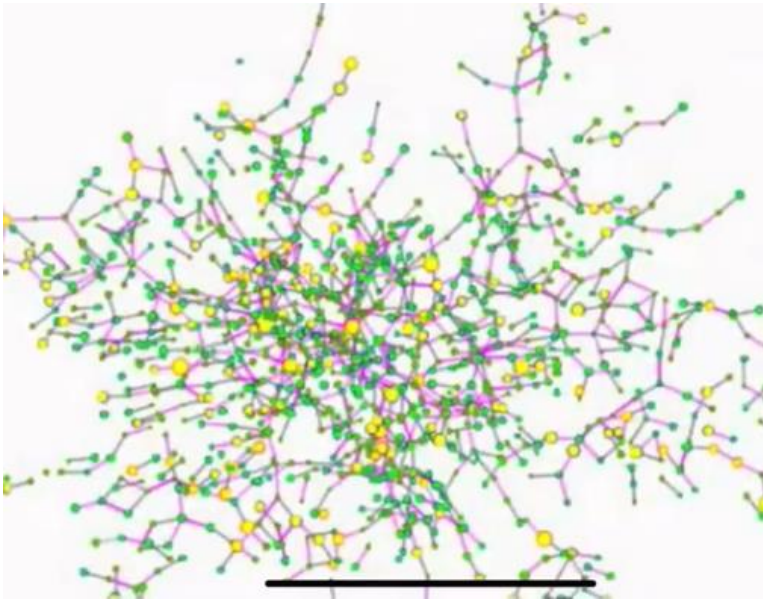
**What changed?**



# Let's be detectives together...

---

## What changed?

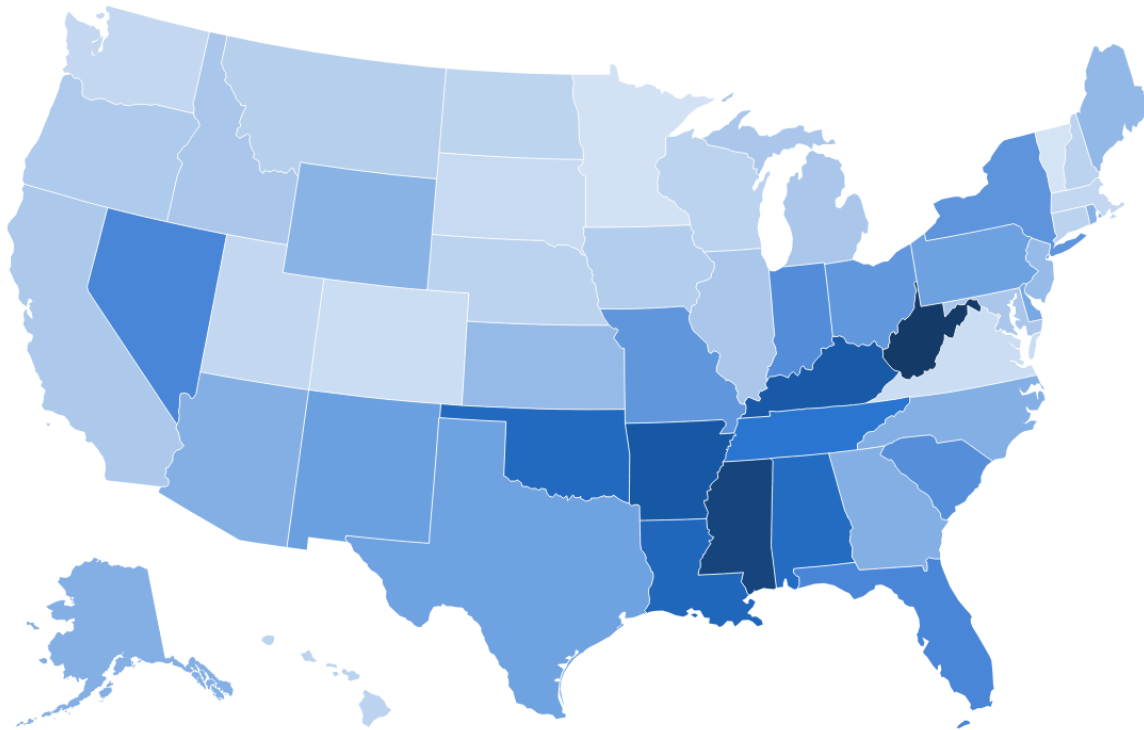


# Let's be detectives together...

---



## **Guess this graph?**



Source: Centers for Disease Control and Prevention



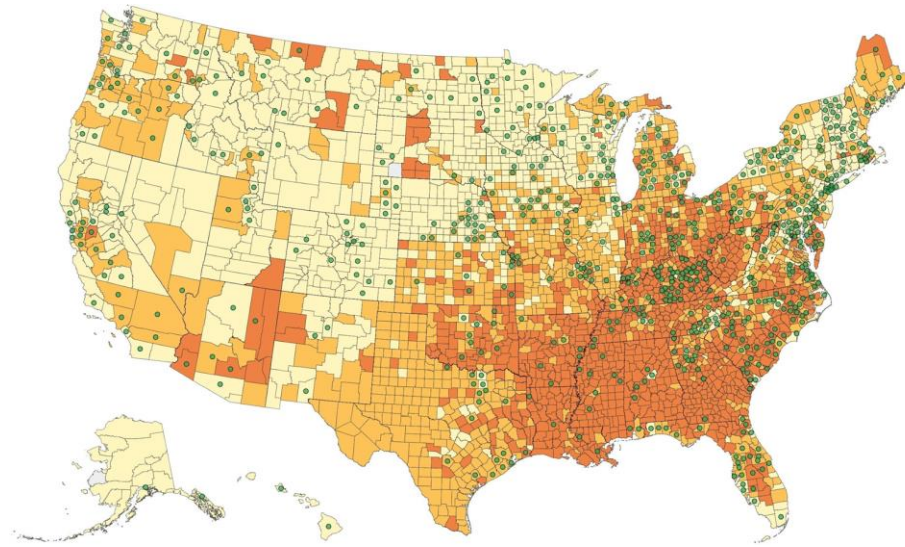
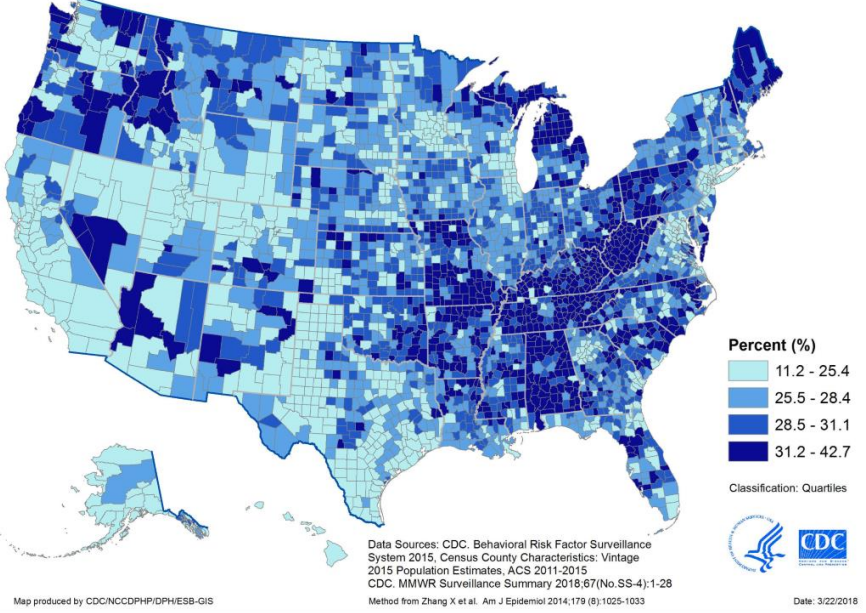
# These maps looks eerily similar...

## Arthritis

## Diabetes

Prevalence by County - 2023

Model-based crude prevalence of arthritis among adults  $\geq 18$  years, by County, United States 2015

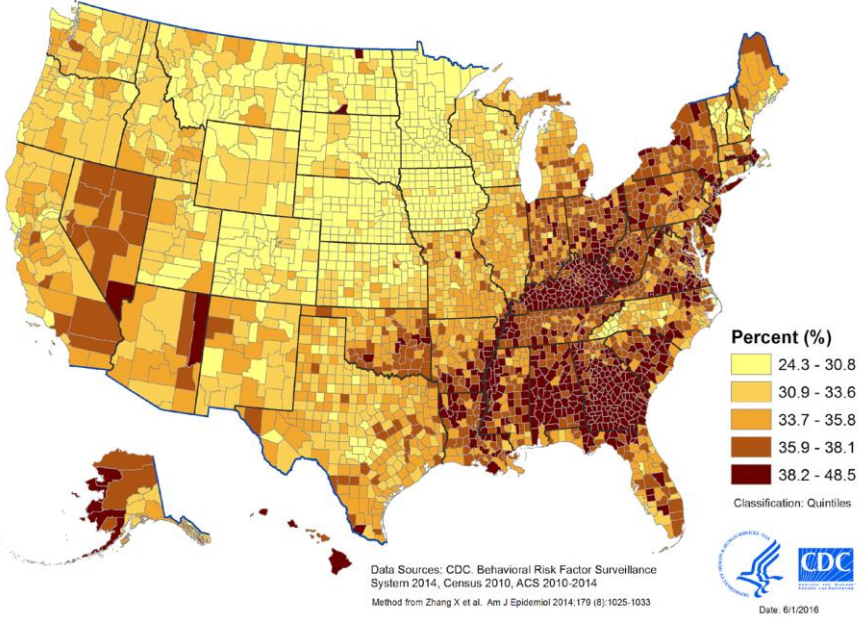


Source: CDC



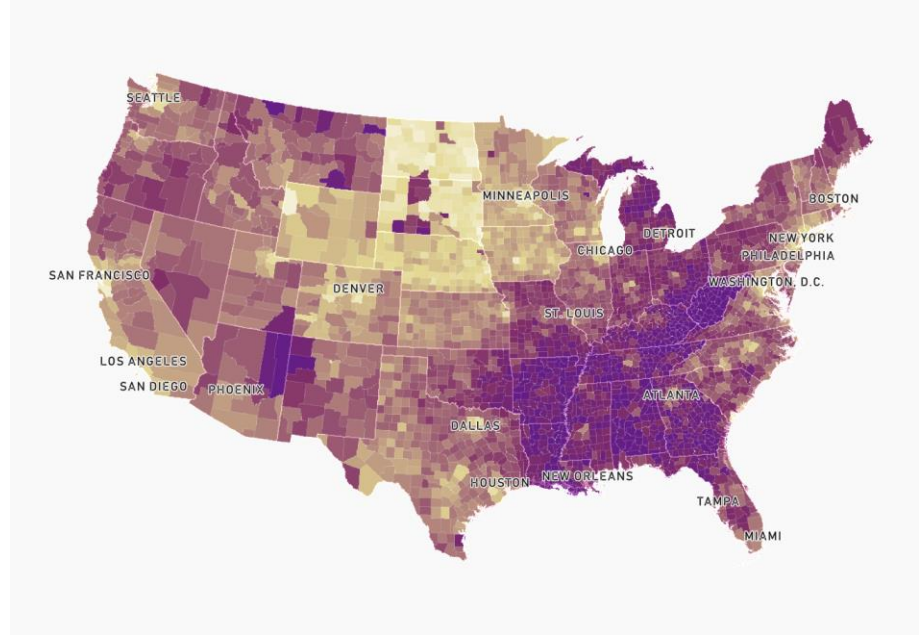
# These maps looks eerily similar...

## Short Sleep Duration 2017



## Mental Distress

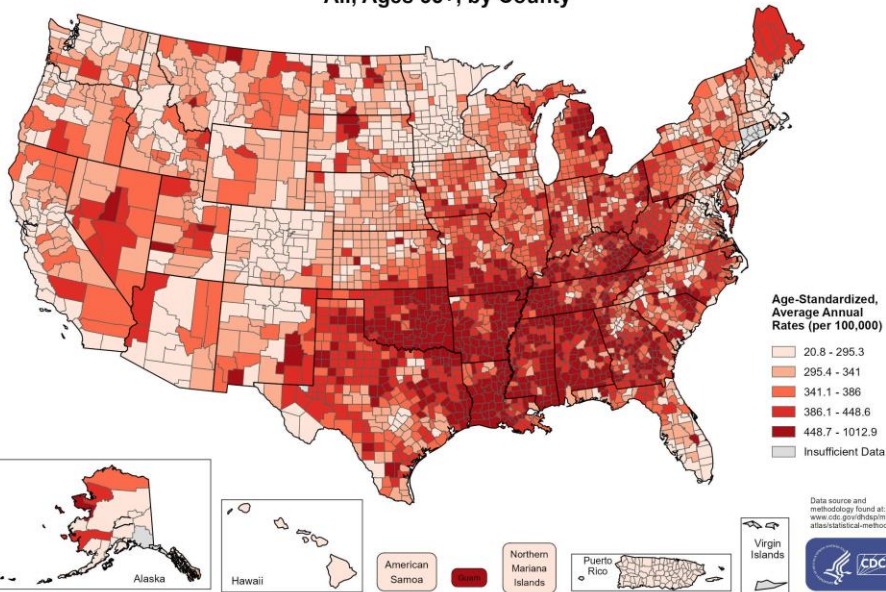
Average number of poor mental health days (2019)



# What is going on here?

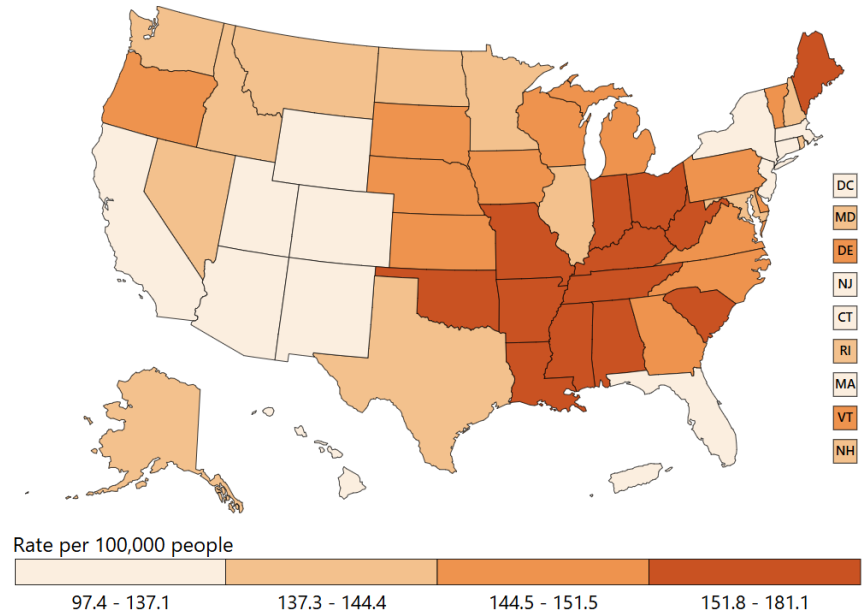
## Heart Disease Death Rates

Heart Disease Death Rates, 2021-2023  
All, Ages 35+, by County



## Cancer Deaths

Rate of Cancer Deaths in the United States, 2023  
All Cancer Sites Combined, Male and Female, All Races and Ethnicities



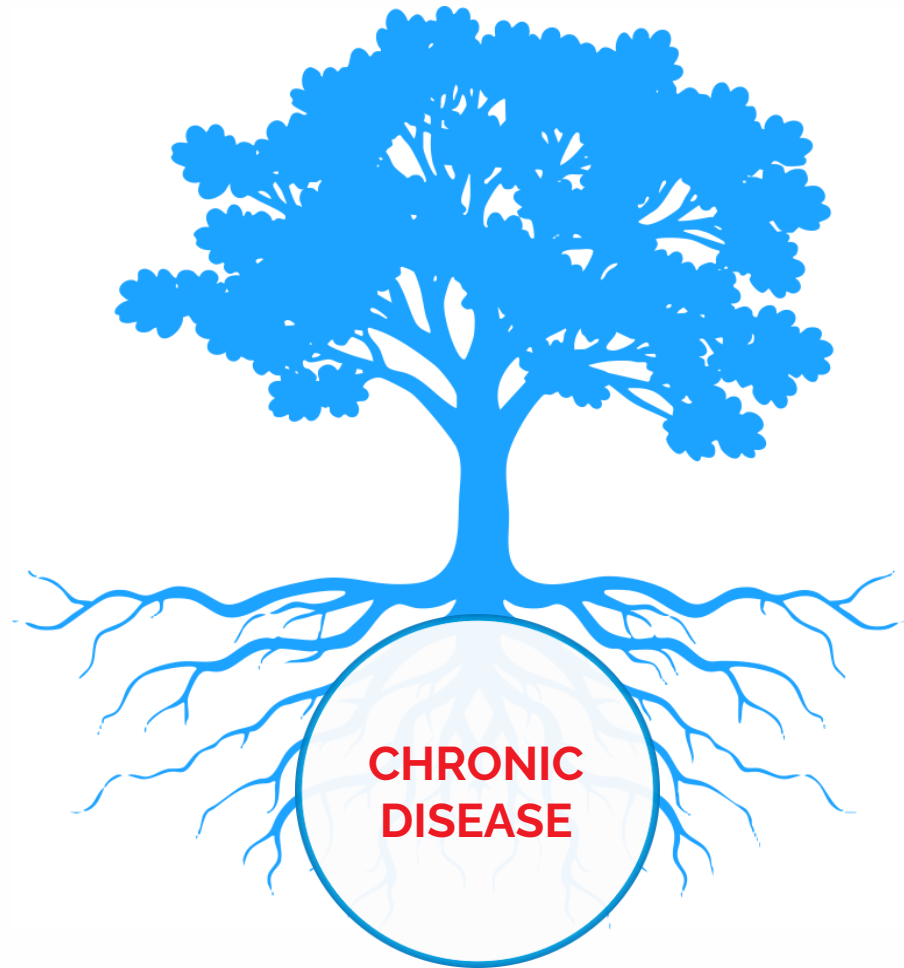
# The Problem

Is it possible  
to explain  
**modern  
chronic  
disease**  
very simply?

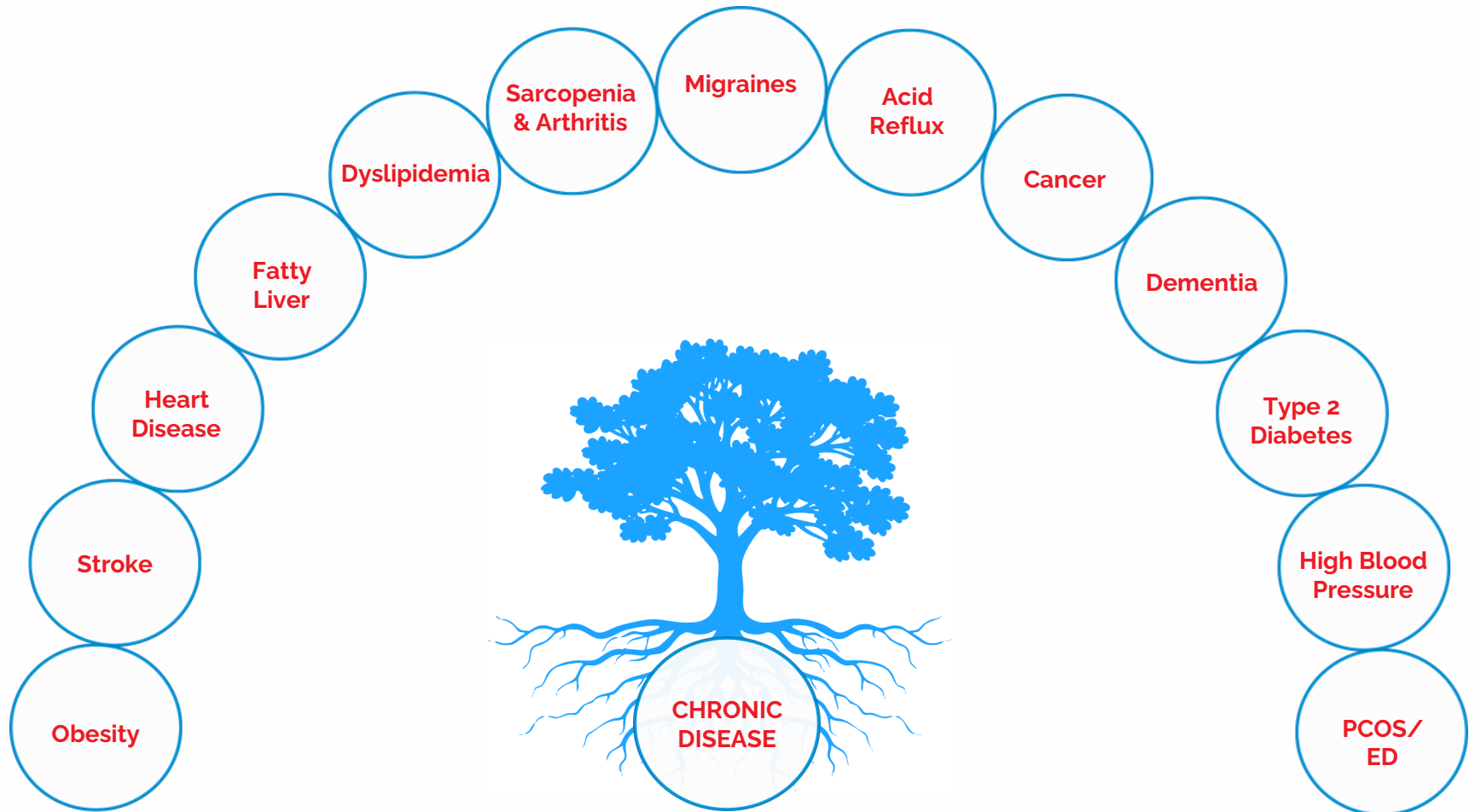


# Ignoring root causes leads to chronic disease...

---



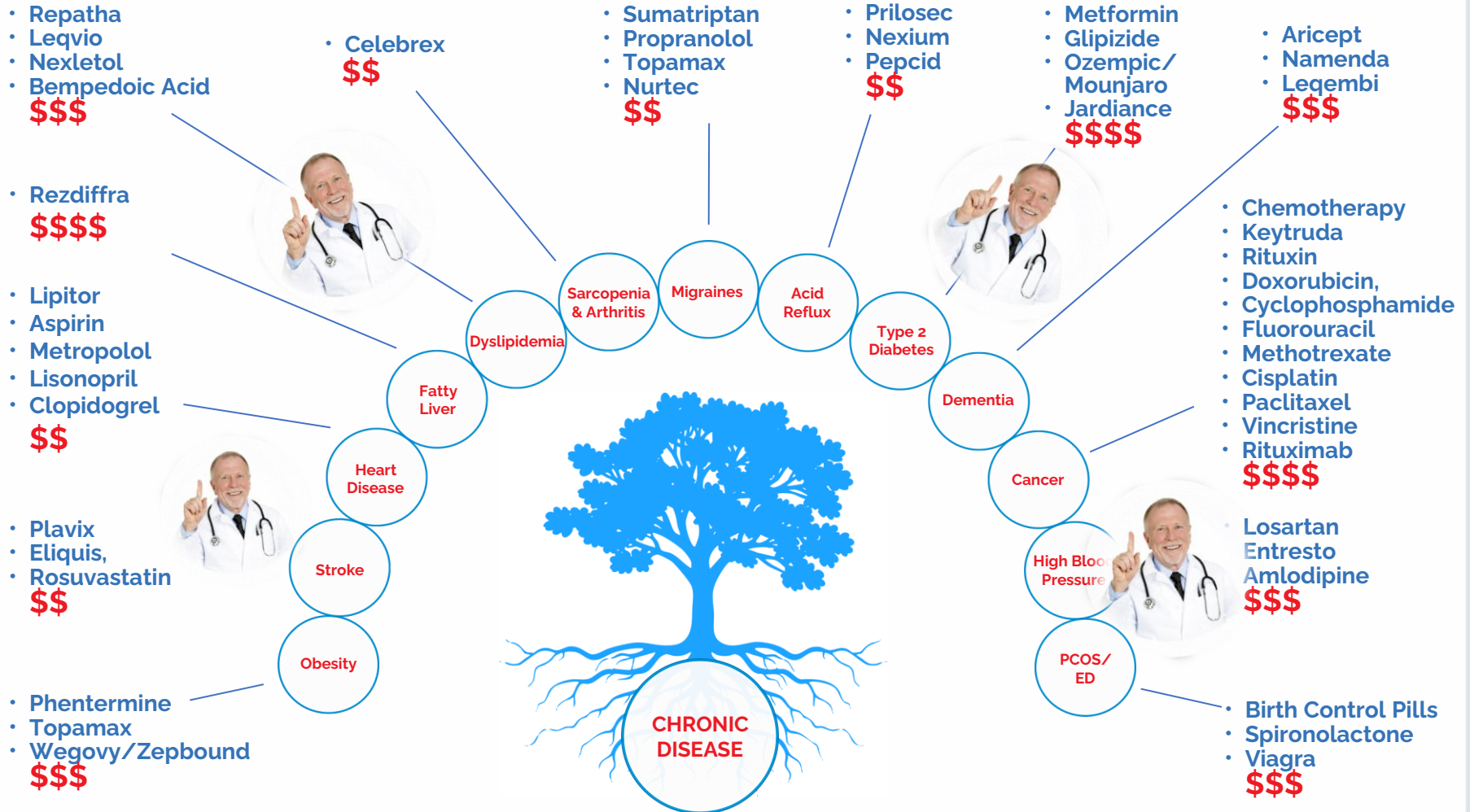
# Ignoring root causes leads to multiple secondary diseases...



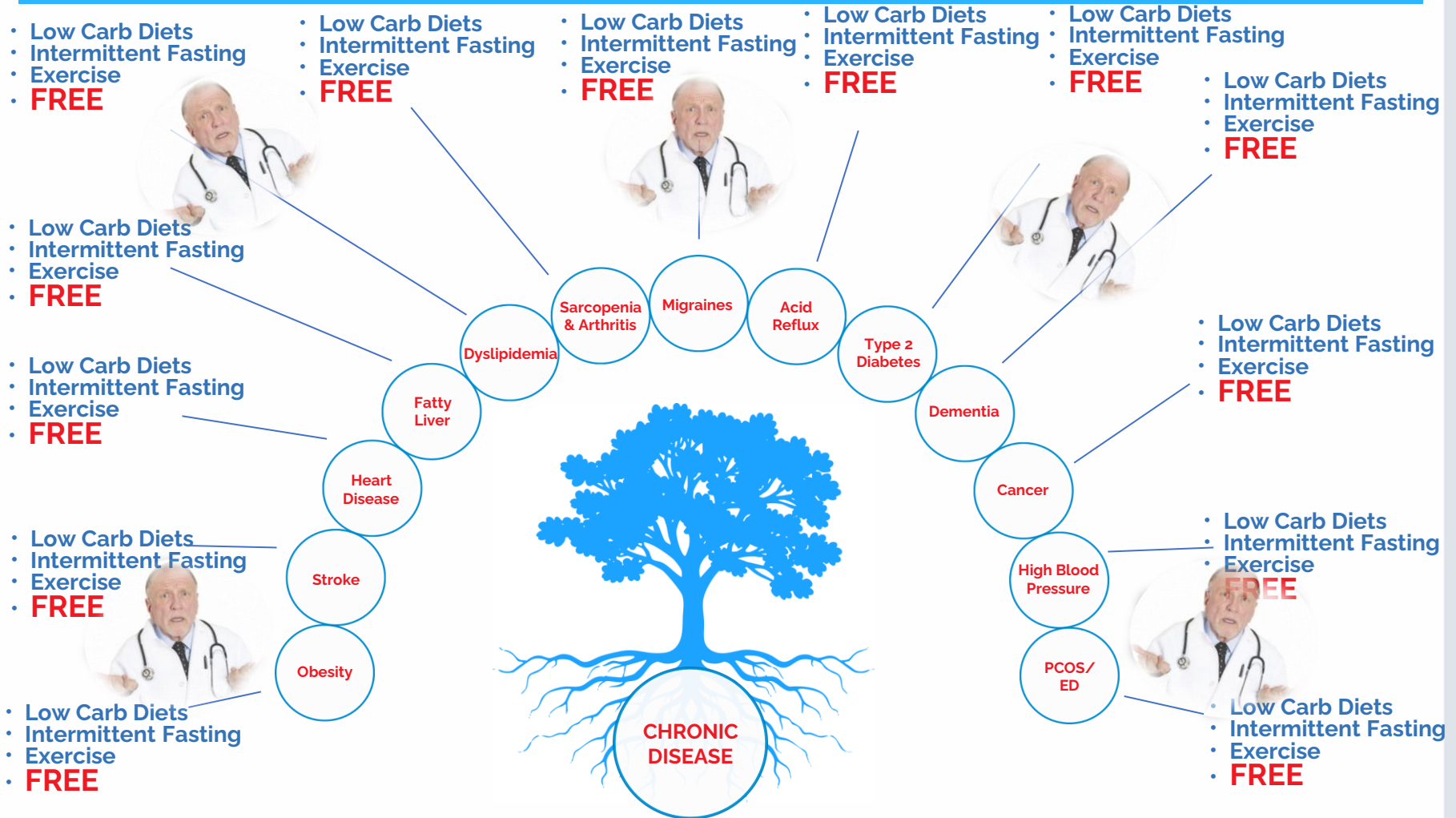
# ...addressing symptoms is lucrative with widespread medical consensus

**KEY:**

- \$ - Less than \$500 per year
- \$\$ - \$500-\$2,000 per year
- \$\$\$ - \$2,000-\$15,000 per year
- \$\$\$\$ - \$15,000+ per year



# Does the data actually support metabolic interventions?



---

OPINION



Case Reports

Case Series

Cohort Studies

Pragmatic Cohort

RCT

Meta-Analysis

Widespread Clinical Use

Type 2 Diabetes

Cardiometabolic



Case Reports

Case Series

Cohort Studies

Pragmatic Cohort

RCT

Meta-Analysis

Widespread Clinical Use

Type 2 Diabetes

Cureus

Open Access Review Article

DOI: 10.7759/cureus.25528

Open Access

Original research

BMJ Open Diabetes Research & Care

# Systematic review and meta-analysis of dietary carbohydrate restriction in patients with type 2 diabetes

Ole Snorgaard,<sup>1</sup> Grith M Poulsen,<sup>2</sup>

**To cite:** Snorgaard O, Poulsen GM, Andersen HK, et al. Systematic review and meta-analysis of dietary carbohydrate restriction in patients with type 2 diabetes. *BMJ Open Diabetes Research and Care* 2017;5:e000354. doi:10.1136/bmjdr-2016-000354

Received 31 October 2016  
Revised 2 February 2017  
Accepted 5 February 2017

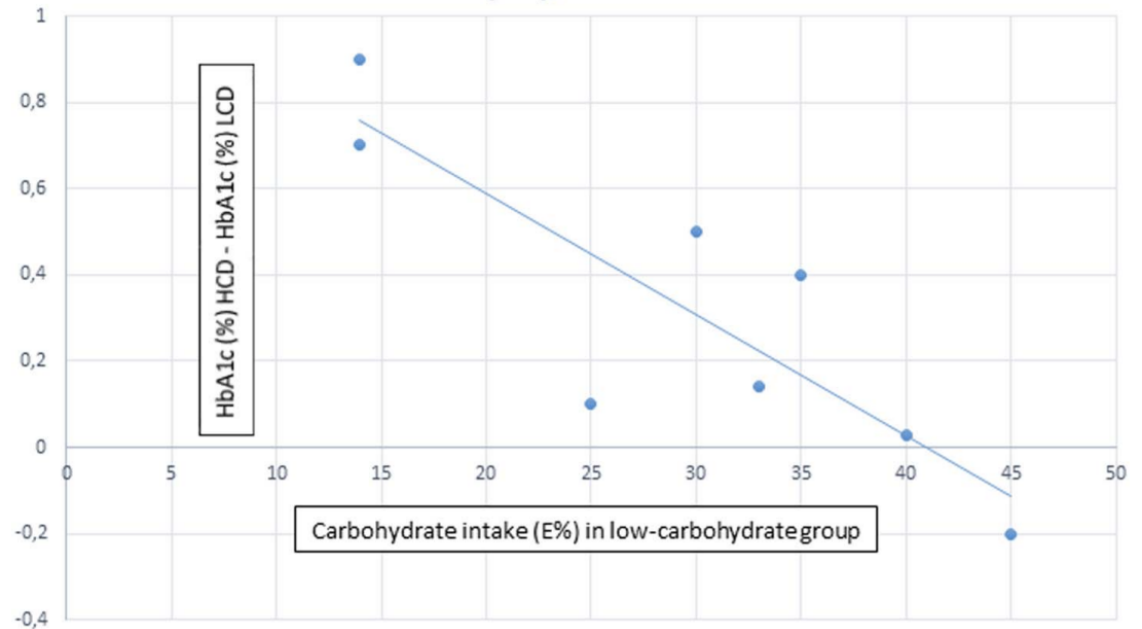
### ABSTRACT

**Objective:** Nutrition therapy is an integral part of management education in patients with type 2 diabetes. Carbohydrates with a low glycemic index are recommended, but the ideal amount of carbohydrate in the diet is unclear. We performed a meta-analysis comparing diets containing low to moderate amounts of carbohydrate (LCD) (energy percentage below 45%) to diets containing high amounts of carbohydrate (HCD) in subjects with type 2 diabetes.

**Research design and methods:** We systematically reviewed Cochrane library databases, EMBASE, MEDLINE in the period 2004–2014 for guideline meta-analyses, and randomized trials assessing outcomes HbA1c, BMI, weight, LDL cholesterol, quality of life (QoL), and attrition.

**Results:** We identified 10 randomized trials comprising 1376 participants in total. In the first 6 months of intervention, LCD was followed by a 0.34% lower HbA1c (3.7 mmol/mol) compared with HCD (95% CI: 0.06 (0.7 mmol/mol), 0.63 (6.9 mmol/mol)). The greater the carbohydrate restriction, the greater the glucose-lowering effect ( $R=-0.85$ ,  $p<0.01$ ). At 6 months or later, however, HbA1c was similar in the 2 diet groups. The effect of the 2 types of diet on BMI, weight, LDL cholesterol, QoL, and attrition rate was similar throughout interventions.

Excess reduction in HbA1c (%) versus carbohydrate intake (E%), 8 RCTs



BMJ Open Diabetes Research and Care 2017;5:e000354. doi:10.1136/bmjdr-2016-000354

Case Reports

Case Series

Cohort Studies

Pragmatic Cohort

RCT

Meta-Analysis

Widespread Clinical Use

Type 2 Diabetes

Obesity

Cardiometabolic



Case Reports

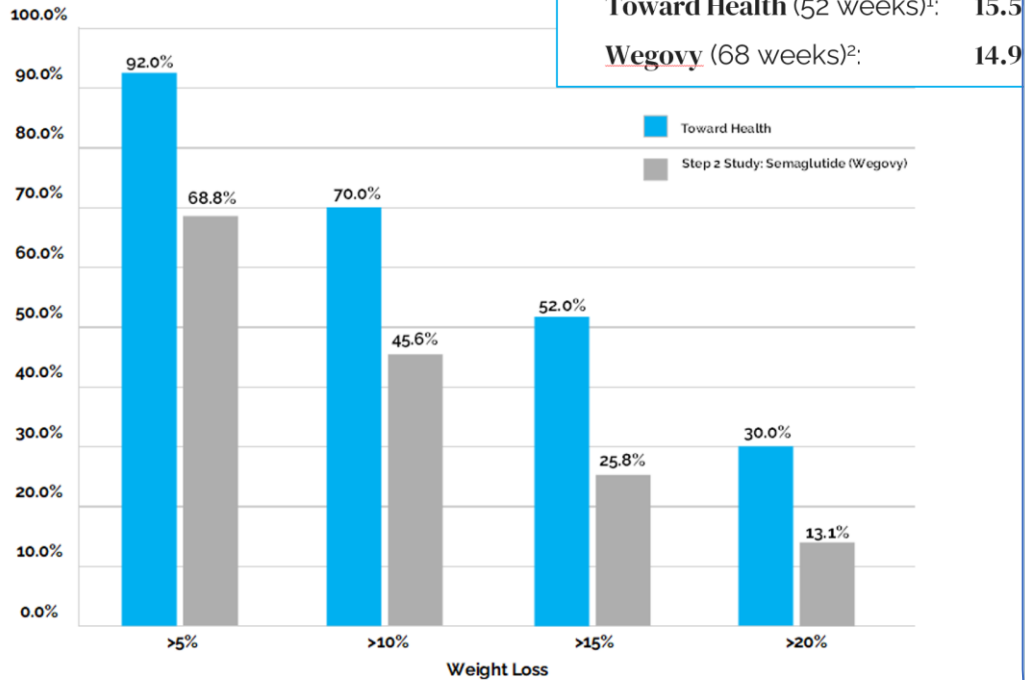
Case Series

Cohort Studies

Pragmatic Cohort

### Published Medical Weight Loss Program Results: **Low-Risk Population**

Comparison of Weight Loss



Change in Body Weight %: Comparison

Toward Health (52 weeks)<sup>1</sup>: 15.5%

Wegovy (68 weeks)<sup>2</sup>: 14.9%

<sup>1</sup>Frontiers in Nutrition

<sup>2</sup>Semaglutide 2.4mg Step 2 Study

Summary WMD

Heterogeneity  $P = 0.000$ ,  $I^2 = 82.2\%$

Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Random, 95% CI
Foster 2010 (39)	30.8	5.8	30	32.8	5.5	31	100.0%	-2.00 [-4.84, 0.84]
<b>Total (95% CI)</b>			<b>30</b>			<b>31</b>	<b>100.0%</b>	<b>-2.00 [-4.84, 0.84]</b>

Heterogeneity: Not applicable  
Test for overall effect:  $Z = 1.38$  ( $P = 0.17$ )

Risk of bias legend

- (A) Random sequence generation (selection bias)
- (B) Allocation concealment (selection bias)
- (C) Blinding of participants and personnel (performance bias)
- (D) Blinding of outcome assessment (detection bias)
- (E) Incomplete outcome data (attrition bias)
- (F) Selective reporting for weight (reporting bias)
- (G) Selective reporting for renal function (reporting bias)
- (H) Other bias

## 12-Month Weight Loss & Deprescription Savings with the TOWARD Approach



ORGANIZATIONAL

# 50

EMPLOYEES



INDIVIDUAL

# 43<sup>lbs</sup>

AVG. WEIGHT LOSS PER PERSON

# 96

total  
DEPRESCRIBED  
MEDICATIONS



# 1.92

DEPRESCRIBED  
MEDICATIONS  
PER PERSON

# TOTAL ANNUALIZED COST SAVINGS \$83,285

# \$1665

ANNUALIZED SAVINGS  
PER PERSON



Case Reports

Case Series

Cohort Studies

Pragmatic Cohort

RCT

Meta-Analysis

Widespread Clinical Use

Type 2 Diabetes

Obesity

Metabolic Syndrome



Frontiers in Nutrition

TYPE Systematic Review

Check for updates

OPEN ACCESS

EDITED BY  
Luciana Neri Nobre,  
Universidade Federal dos Vales do  
Jequitinhonha e Mucuri (UFVJM), Brazil

REVIEWED BY  
Dina Keumala Sari,  
Universitas Sumatera Utara, Indonesia  
Nikola Savic,  
University of Kragujevac, Serbia

\*CORRESPONDENCE  
Ruixi Xi  
✉ xiruixi@163.com  
Ruina Bai  
✉ brntcl@126.com

<sup>†</sup>These authors have contributed equally to this work

RECEIVED 23 October 2024  
ACCEPTED 13 December 2024  
PUBLISHED 06 January 2025

The effect of a low-carbohydrate diet on metabolic health in obese patients with T2DM: a meta-analysis

Wende Tian, Qiyu Liu<sup>1</sup>, J...

<sup>1</sup>National Clinical Center for Diabetes, Institute of Chinese Medicine, Beijing, China, <sup>3</sup>Lia...

## 5 Conclusion

After a thorough meta-analysis of the current evidence and taking into account the potential side effects, low-carbohydrate diets appear to significantly improve glycemic control, TG, HDL-C, weight, BMI, DBP, and waist circumference in overweight or obese patients with T2DM. This study provides valuable dietary guidance for managing T2DM in this patient population. Notably, the significant reduction in waist circumference underscores the effectiveness of low-carbohydrate diets in addressing central obesity, a critical factor in metabolic health and cardiovascular risk. These substantial improvements in both anthropometric and metabolic parameters highlight the potential of low-carbohydrate diets as a comprehensive strategy for managing T2DM. Therefore, adopting low-carbohydrate diets could offer multifaceted benefits in controlling metabolic health and reducing obesity-related risks in individuals with T2DM.



Case Reports

Case Series

Cohort Studies

Pragmatic Cohort

RCT

Meta-Analysis

Widespread Clinical Use

Type 2 Diabetes

Obesity

Metabolic Syndrome

Hypertension

## Comparing Very Low-Carbohydrate vs DASH Diets for Overweight or Obese Adults With Hypertension and Prediabetes or Type 2 Diabetes: A Randomized Trial

Laura R. Saslow, PhD<sup>1</sup>  
 Lenette M. Jones, PhD, RN<sup>1</sup>  
 Ananda Sen, PhD<sup>2,3</sup>  
 Julia A. Wolfson, PhD<sup>4,5</sup>  
 Heidi L. Diez, PharmD<sup>6,7</sup>  
 Alison O'Brien, MPH<sup>1</sup>  
 Cindy W. Leung, ScD<sup>8</sup>  
 Hovig Bayandorian, MA  
 Jennifer Daubenmier, PhD<sup>9</sup>  
 Amanda L. Missel, PhD<sup>1</sup>  
 Caroline Richardson, MD<sup>2</sup>

<sup>1</sup>Department of Health Behavior and Biological Sciences, School of Nursing, University of Michigan, Ann Arbor, Michigan  
<sup>2</sup>Department of Family Medicine, Michigan Medicine, University of Michigan, Ann Arbor, Michigan  
<sup>3</sup>Department of Biostatistics, School of Public Health, University of Michigan, Ann Arbor, Michigan  
<sup>4</sup>Department of International Health, Johns Hopkins Bloomberg School of Public Health, Baltimore, Maryland

### ABSTRACT

**PURPOSE** Adults with a triple multimorbidity (hypertension, prediabetes or type 2 diabetes, but not all) are common. Recommendations for diet, exercise, and medication are often conflicting. We compared the effects of a very low-carbohydrate (VLC) diet to a DASH diet on blood pressure, glycemic control, and weight in this population.

### METHOD

Adults with a triple multimorbidity (hypertension, prediabetes or type 2 diabetes) were randomized to a VLC diet or a DASH diet. Results were compared at baseline and 12 weeks.

### RESULTS

At baseline, VLC diet participants had higher systolic blood pressure (SBP) and HbA<sub>1c</sub> than DASH diet participants. At 12 weeks, SBP and HbA<sub>1c</sub> were lower in the VLC diet group (P = .034).

### CONCLUSIONS

Weight, glycemic control, and blood pressure were improved in the VLC diet group compared with the DASH diet group. These findings suggest that a VLC diet may be a better option for adults with a triple multimorbidity.

**Table 2. Estimated Mean (SE) of Outcomes Across Diet and Time From Linear Mixed Model**

Outcome	VLC Diet				DASH Diet				Difference in Change (VLC Lower)	Between-Group P Value
	Baseline	Post	Change	Within-Group P Value	Baseline	Post	Change	Within-Group P Value		
SBP, mm Hg	133.72 (1.73)	123.95 (1.88)	-9.77 (1.66)	<.001	132.84 (1.69)	127.66 (1.80)	-5.18 (1.59)	.002	-4.59	.046
HbA <sub>1c</sub> , %	6.09 (0.07)	5.74 (0.08)	-0.35 (0.07)	<.001	6.10 (0.07)	5.97 (0.07)	-0.14 (0.07)	.06	-0.21	.034
Weight, lb	219.24 (5.39)	200.10 (5.41)	-19.14 (1.73)	<.001	236.43 (5.1)	226.1 (5.2)	-10.34 (1.73)	<.001	-8.81	.0003

DASH = Dietary Approaches to Stop Hypertension; HbA<sub>1c</sub> = glycated hemoglobin; SBP = systolic blood pressure; VLC = very low-carbohydrate.

Note: Outcomes were analyzed using a linear mixed model including all possible interactions between diet, support, and time and adjusted for age and sex. Results are presented collapsed over all other factors: support allocation, sex, and at the mean value of age. The between-group P value is calculated from a Z-test based on the estimated mean changes and the associated SE values reported in the table.

Case Reports

Case Series

Cohort Studies

Pragmatic Cohort

RCT

Meta-Analysis

Widespread Clinical Use

Type 2 Diabetes

Obesity

Metabolic Syndrome

# Effect of a ketogenic diet on hepatic steatosis and hepatic mitochondrial metabolism in nonalcoholic fatty liver disease

Panu K. Luukkonen<sup>a,b,c</sup>, Sylvie Dufour<sup>a,d</sup>, Kun Lyu<sup>e</sup>, Xian-Man Zhang<sup>a,d</sup>, Antti Hakkarainen<sup>f,g</sup>, Tiina E. Lehtimäki<sup>f</sup>, Gary W. Cline<sup>a,d</sup>, Kitt Falk Petersen<sup>a,d</sup>, Gerald I. Shulman<sup>a,d,e,1,2</sup>, and Hannele Yki-Järvinen<sup>b,c,1,2</sup>

<sup>a</sup>Department of Internal Medicine, Yale School of Medicine, New Haven, CT 06520; <sup>b</sup>Minerva Foundation Institute for Medical Research, Helsinki 00290, Finland; <sup>c</sup>Department of Medicine, University of Helsinki and Helsinki University Hospital, Helsinki 00290, Finland; <sup>d</sup>Yale Diabetes Research Center, Yale School of Medicine, New Haven, CT 06520; <sup>e</sup>Department of Cellular & Molecular Physiology, Yale School of Medicine, New Haven, CT 06520; <sup>f</sup>Department of Radiology, HUS Medical Imaging Center, University of Helsinki and Helsinki University Hospital, Helsinki 00290, Finland; and <sup>g</sup>Department of Neuroscience and Biomedical Engineering, Aalto University School of Science, 00076 Espoo, Finland

Contributed by Gerald I. Shulman, January 31, 2020 (sent for review December 26, 2019; reviewed by Fredrik Karpe and Roy Taylor)

Weight loss by ketogenic diet (KD) has gained popularity in management of nonalcoholic fatty liver disease (NAFLD). KD rapidly reverses NAFLD and insulin resistance despite increasing circulating nonesterified fatty acids (NEFA), the main substrate for synthesis of intrahepatic triglycerides (IHTG). To explore the underlying mechanism, we quantified hepatic mitochondrial fluxes and their regulators in humans by using positional isotopomer NMR tracer analysis. Ten overweight/obese subjects received stable isotope infusions of [D<sub>7</sub>]glucose, [<sup>13</sup>C<sub>4</sub>]β-hydroxybutyrate and [3-<sup>13</sup>C]lactate before and after a 6-d KD. IHTG was determined by proton magnetic resonance spectroscopy (<sup>1</sup>H-MRS). The KD diet decreased IHTG by 31% in the face of a 3% decrease in body weight and decreased hepatic insulin resistance (–58%) despite an increase in NEFA concentrations (+35%). These changes were attributed to increased net hydrolysis of IHTG and partitioning of the resulting fatty acids toward ketogenesis (+232%) due to reductions in serum insulin concentrations (–53%) and hepatic citrate synthase flux (–38%), respectively. The former was attributed to decreased hepatic insulin resistance and the latter to increased hepatic mitochondrial redox state (+167%) and decreased plasma leptin (–45%) and triiodothyronine (–21%) concentrations. These data demonstrate heretofore undescribed adaptations underlying the reversal of NAFLD by KD: That is, markedly altered hepatic mitochondrial fluxes and redox state to promote ketogenesis rather than synthesis of IHTG.

after 48 h of caloric restriction (26). We previously showed that a hypocaloric, KD induces an ~30% reduction in IHTG content in 6 d despite increasing circulating NEFA (27).

While the antisteatotic effect of KD is well-established, the underlying mechanisms by which it does so remain unclear. KD increases plasma NEFA concentrations, the main substrate of IHTG (11). In the liver, NEFA can either be re-esterified into complex lipids, such as TGs, or be transported to the mitochondria to be metabolized by β-oxidation into acetyl-CoA, which in turn can either be irreversibly condensed with oxaloacetate by citrate synthase to form citrate and enter the TCA cycle for terminal oxidation to CO<sub>2</sub> (28, 29) or it can enter the ketogenic pathway, where it is converted into acetoacetate (AcAc) and β-hydroxybutyrate (β-OHB) (28). These mitochondrial fluxes are tightly regulated by substrate availability and product inhibition (29), mitochondrial redox state (30), and hormones, such as leptin (31) and triiodothyronine (T3) (32).

## Significance

Ketogenic diet is an effective treatment for nonalcoholic fatty liver disease (NAFLD). Here, we present evidence that hepatic mitochondrial fluxes and redox state are markedly altered during ketogenic diet-induced reversal of NAFLD in humans.

Cardiometabolic



Case Reports

Case Series

Cohort Studies

Pragmatic Cohort

RCT

Meta-Analysis

Widespread Clinical Use

Type 2 Diabetes

Obesity

## RESULTS OF A METABOLIC WELLNESS PROGRAM

A partnership with a medical clinic, a corporation and employees with obesity, prediabetes or diabetes who voluntarily participated in treatment using carbohydrate-restriction.

**52.9**  
years  
Mean Age

**290.5** lbs  
Mean weight

**44.9** kg/m<sup>2</sup>  
Mean BMI

**56%**  
Female

- Low carbohydrate diet (<30g/day)
- Ongoing Support
- Education
- Continuous Glucose Monitor
- Smartphone App

### Metabolic Results

At 6-months into this ongoing program...



**38** lbs

Avg Weight Loss  
\*All participants lost at least 15lbs



**1.1%**

a1c



**44%**

American College of Cardiology / American Heart Association 10-Year ASCVD Risk



**17** mmHg

Systolic Blood Pressure

### Corporate Savings

\$4,000 per year/patient From Medication Deprescription

### Medications Eliminated

Cholesterol: 🍯🍯

Insulin: 🍯🍯

Reflux: 🍯🍯🍯

Blood Pressure: 🍯🍯🍯🍯

Diabetes: 🍯🍯🍯🍯🍯🍯🍯

This high-risk cohort (nearly 2/3 of patients had diabetes) had an average starting HbA1c of 7.1%:

**Deprescription cost savings:  
\$4,000 per patient, per year**

Metabolites: A Company Is Only as Healthy as Its Workers: A 6-Month Metabolic Health Management Pilot Program Improves Employee Health and Contributes to Cost Savings

# Demonstrated cardiometabolic risk reduction

Study	A1c	Weight	BP (mm Hg)	Deprescriptions
Toward Health*	↓ 7.1% to 6.0% (-1.1%)	↓ 288 lbs to 251 lbs (-38 lbs)	↓ 141/83 to 124/78 (-17/6)	Saved \$4,000 per patient, per year in medication deprescriptions
Toward Health**	↓ 6.34% to 5.5% (-.79%)	↓ 271 lbs to 228 lbs (-43 lbs)	↓ 144/87 to 126/80 (-14/7)	Saved \$1,665 per patient, per year in medication deprescriptions

**Can we be more conclusive?**

The TOWARD Approach provided a **44% reduction in the employees' 10-year risk of heart attack and stroke in an initial pilot**



\* High-risk cohort (6-months): 2/3 with diabetes  
 \*\* Lower-risk cohort (12-months): 1/3 with diabetes

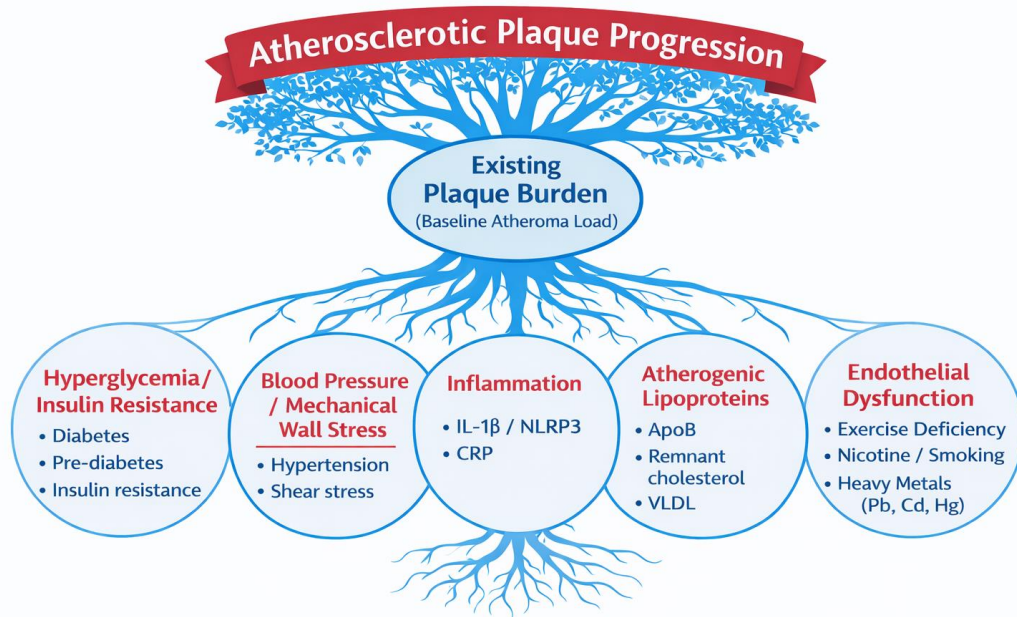
# One-Year PREVENT Data – 120 patients

(in submission, please do not share)



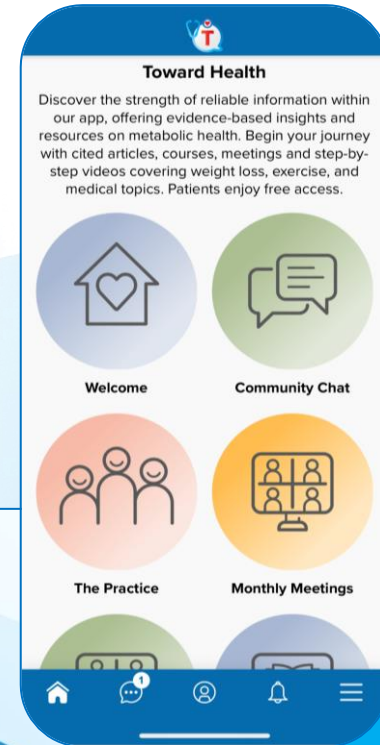
# Reversing Coronary Plaque

## Human Clinical Trial Evidence



**ROOT CAUSES OF CORONARY PLAQUE PROGRESSION**

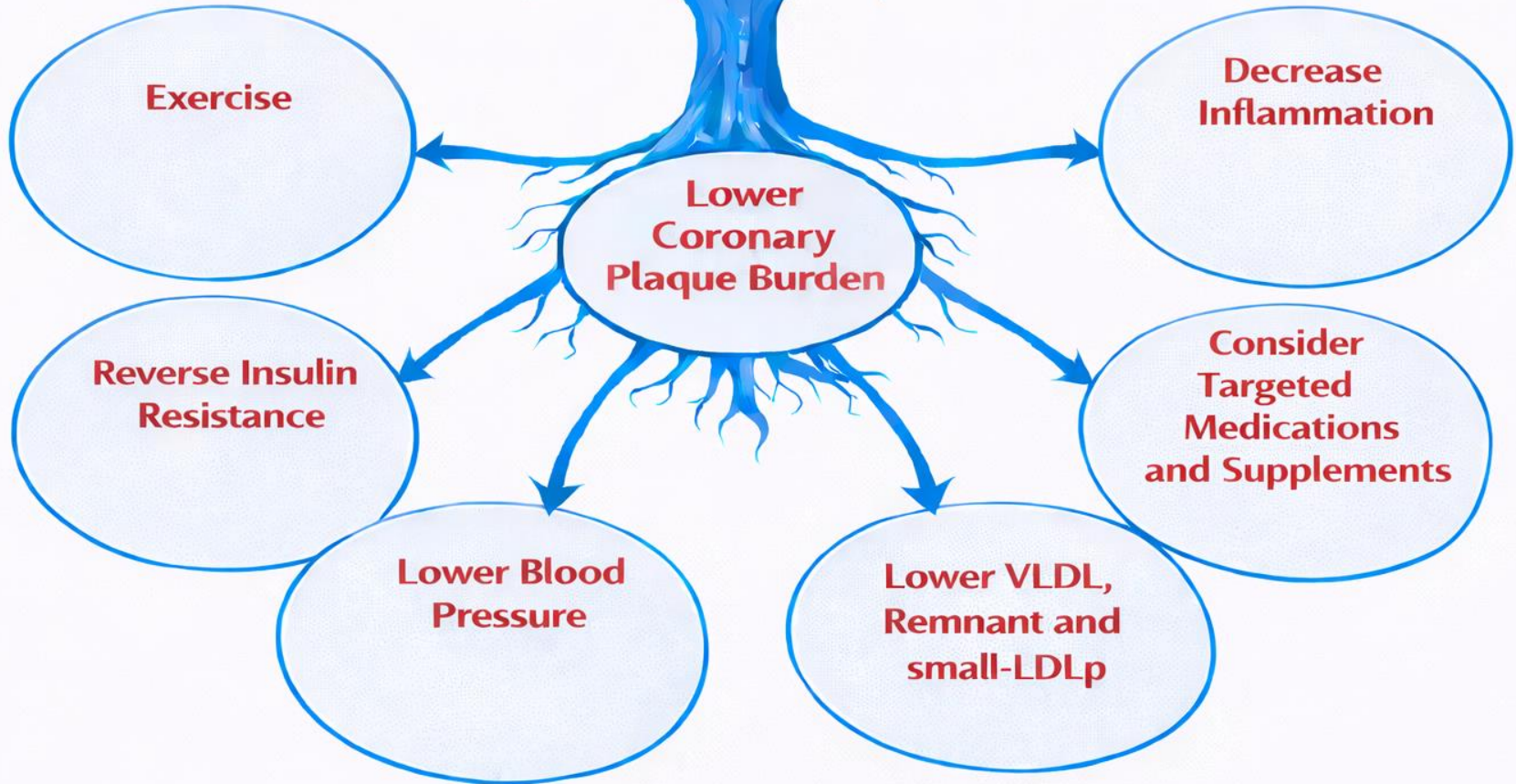
The Toward Health program includes the Toward Health app which has a fully interactive Cardiovascular Disease Reversal and CAC course for all patients.



A metabolic health clinic  
in your pocket



# Plaque Reversal



## PLAQUE REVERSAL STRATEGIES



# Three Diseases, One Risk Calculator

## **Cardiovascular disease**

Heart attacks, strokes, and heart failure

## **Atherosclerotic cardiovascular disease**

Heart attacks, and stroke

## **Heart failure**

Decreased functioning/forward flow of the heart. Can cause leg swelling, shortness of breath, and poor quality of life



**American  
Heart  
Association®**

## **PREVENT Calculator**

Newest risk calculator from the American Heart Association that aims to predict 10-year and 30-year risk of these diseases from metabolic markers



# Three Diseases, One Risk Calculator

CVD

ASCVD

Heart Failure

## Sex\*

Male  Female

## Age (years)\*

30-79

## Total Cholesterol (mg/dL)\*

130-320

## Diabetes

Any history of diabetes.

No  Yes

## Current Smoking

Any cigarette use within the last 30 days

No  Yes

## HDL Cholesterol (mg/dL)\*

20-100

## SBP (mmHg)\*

90-200

## Anti-hypertensive medication

Current use of any medication for hypertension

No  Yes

## Lipid-lowering medication

Current use of statin medication to lower cholesterol

No  Yes

## BMI (kg/m<sup>2</sup>)\*

18.5-39.9

## eGFR (mL/min/1.73m<sup>2</sup>)\*

15-140

The following three predictors are optional for further personalization of risk assessment. When they are clinically indicated or available,

If available or indicated, select "Yes" and enter the value.

## UACR (mg/g)

UACR is clinically indicated for individuals with chronic kidney disease, diabetes, or hypertension

No  Yes

## HbA1C

HbA1c is clinically indicated for individuals with diabetes, prediabetes, overweight, or obesity, or those with history of gestational diabetes

No  Yes

## Zip Code

valid 5-digit zip code is needed to estimate social deprivation index [SDI]

No  Yes



# Cardiovascular Disease Risk

	Risk before Intervention	Risk After Intervention	Absolute Risk Reduction	Relative Risk Reduction
<b>10-year risk</b>	7.3%			
<b>30-year risk</b>	29.3%			

**Cardiovascular Disease = Heart attacks, strokes, heart failure**



# Cardiovascular Disease Risk

	Risk before Intervention	Risk After Intervention	Absolute Risk Reduction	Relative Risk Reduction
<b>10-year risk</b>	7.3%	3.8%		
<b>30-year risk</b>	29.3%	20.3%		

**Cardiovascular Disease = Heart attacks, strokes, heart failure**



# Cardiovascular Disease Risk

	Risk before Intervention	Risk After Intervention	Absolute Risk Reduction	Relative Risk Reduction
<b>10-year risk</b>	7.3%	3.8%	↓ 3.5%	↓ 48%
<b>30-year risk</b>	29.3%	20.3%	↓ 9.0%	↓ 31%

**Cardiovascular Disease = Heart attacks, strokes, heart failure**



# Atherosclerotic Cardiovascular Disease Risk

	Risk before Intervention	Risk After Intervention	Absolute Risk Reduction	Relative Risk Reduction
<b>10-year risk</b>	4.4%			
<b>30-year risk</b>	17.9%			

**Atherosclerotic cardiovascular disease = Heart attacks**



# Atherosclerotic Cardiovascular Disease Risk

	Risk before Intervention	Risk After Intervention	Absolute Risk Reduction	Relative Risk Reduction
<b>10-year risk</b>	4.4%	2.3%		
<b>30-year risk</b>	17.9%	12.0%		

**Atherosclerotic cardiovascular disease = Heart attacks**



# Atherosclerotic Cardiovascular Disease Risk

	Risk before Intervention	Risk After Intervention	Absolute Risk Reduction	Relative Risk Reduction
<b>10-year risk</b>	4.4%	2.3%	↓ 2.1%	↓ 48%
<b>30-year risk</b>	17.9%	12.0%	↓ 5.9%	↓ 33%

**Atherosclerotic cardiovascular disease = Heart attacks and stroke**



# Heart Failure Risk

	<b>Risk before Intervention</b>	<b>Risk After Intervention</b>	<b>Absolute Risk Reduction</b>	<b>Relative Risk Reduction</b>
<b>10-year risk</b>	6.2%			
<b>30-year risk</b>	26.1%			

**Heart failure = Decreased functioning/forward flow of the heart.  
Can cause leg swelling, shortness of breath, and poor quality of life**



# Heart Failure Risk

	Risk before Intervention	Risk After Intervention	Absolute Risk Reduction	Relative Risk Reduction
<b>10-year risk</b>	6.2%	2.3%		
<b>30-year risk</b>	26.1%	14.2%		

**Heart failure = Decreased functioning/forward flow of the heart.  
Can cause leg swelling, shortness of breath, and poor quality of life**



# Heart Failure Risk

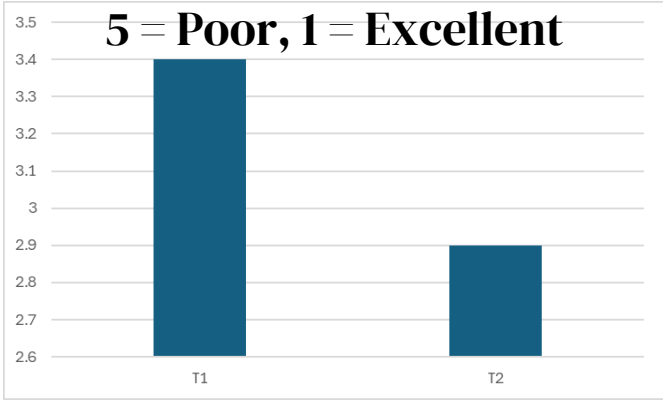
	Risk before Intervention	Risk After Intervention	Absolute Risk Reduction	Relative Risk Reduction
<b>10-year risk</b>	6.2%	2.3%	↓ 3.9%	↓ 63%
<b>30-year risk</b>	26.1%	14.2%	↓ 11.9%	↓ 46%

**Heart failure = Decreased functioning/forward flow of the heart.  
Can cause leg swelling, shortness of breath, and poor quality of life**

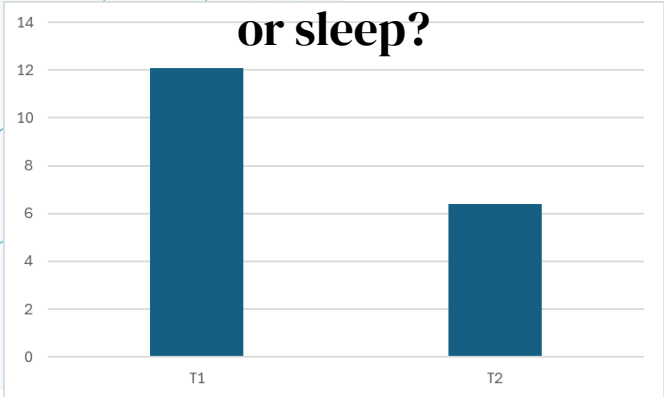


# Quality of Life Improvement Surveys

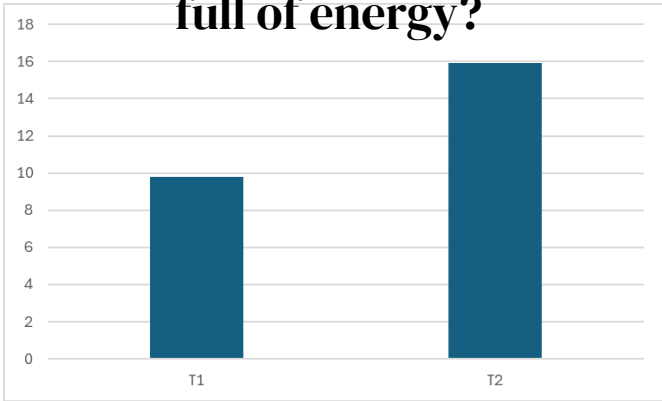
## How Would You Rate Your General Health?



## How many days during the past 30 did you NOT get enough rest or sleep?



## How many days during the past 30 have you felt very healthy and full of energy?



# Effects of the very low-carbohydrate ketogenic diet in women with polycystic ovary syndrome: a systematic review with meta-analysis of clinical trials

Karniza Khalid,  
Syed A. A. Rizvi

<sup>1</sup>Endocrine Unit, Specialised Centre for Endocrinology, 50588 Kuala Lumpur, Malaysia  
<sup>2</sup>Clinical Research Centre, 50588 Kuala Lumpur, Malaysia  
<sup>3</sup>College of Biomedical Sciences, 50588 Kuala Lumpur, Malaysia  
<sup>4</sup>Biochemistry Unit, Specialised Centre for Endocrinology, 50588 Kuala Lumpur, Malaysia  
**Correspondence:** Karniza Khalid, Endocrine Unit, Specialised Centre for Endocrinology, 50588 Kuala Lumpur, Malaysia

## Abstract

**Context:** Ketogenic diet (PCOS). Despite studies derived from small populations.

**Objective:** To pool evidence on the effects of the very low-carbohydrate ketogenic diet on serum progesterone levels.

**Methods:** PubMed, Scopus, and Cochrane were searched for studies on the ketogenic diet in PCOS. The outcome of interest was the effect on serum progesterone levels. Extracted data on study characteristics and outcomes were assessed for bias.

**Results:** Following meta-analysis, the mean difference in reproductive hormone levels was  $-0.223$  (95% CI  $-0.414$  to  $0.168$ ;  $P = 0.002$ ). Significant changes were observed in anthropometric and metabolic markers.

**Conclusion:** Short-term use of the very low-carbohydrate ketogenic diet in women with PCOS is associated with significant improvements in anthropometric and metabolic markers.

**Key Words:** polycystic ovary syndrome, ketogenic diet, progesterone, anthropometric markers, metabolic markers.

**Abbreviations:** BMI, body mass index; LH, luteinising hormone; FSH, follicle-stimulating hormone; T, testosterone; SHBG, sex hormone-binding globulin.

**BJN**

British Journal of Nutrition



British Journal of Nutrition

## Article contents

Abstract

References

# Effects of the very low-carbohydrate ketogenic diet in women with polycystic ovary syndrome: a systematic review with meta-analysis of clinical trials

Published online by Cambridge University Press: 18 November 2025

Jéssica Abdo Gonçalves Tosatti, Fernanda Medeiros Vale Magalhães and Karina Braga Gomes 

Show author details 

Article Supplementary materials Metrics

Get access

Share

Cite

Rights & Permissions

## Abstract

Polycystic ovary syndrome is a disorder characterised by insulin resistance, low-grade inflammation and increased adipose tissue. The very low-carbohydrate ketogenic diet has been suggested to reduce obesity risks in polycystic ovary syndrome. This study aimed to update the evidence on the effects of the very low-carbohydrate ketogenic diet in women with polycystic ovary syndrome. Searches were conducted in electronic databases for randomised clinical trials addressing the research question. The values for the meta-analysis were presented as weighted mean difference (WMD). Twelve studies were included in the qualitative analysis and eleven in the quantitative analysis. Significant reductions were observed in anthropometric outcomes: weight (WMD:  $-9.57$  kg;  $P < 0.0001$ ), waist circumference (WMD:  $-7.75$  cm;  $P < 0.0009$ ), fat body mass (WMD:  $-7.44$  kg;  $P = 0.0008$ ), BMI (WMD:  $-3.45$  kg/m<sup>2</sup>;  $P < 0.0001$ ) and waist-to-hip ratio (WMD:  $-0.02$ ;  $P < 0.0034$ ). Hormonal improvements included free testosterone (WMD:  $-0.31$  ng/dl;  $P < 0.0001$ ), total testosterone (WMD:  $-7.21$  ng/dl;  $P < 0.0001$ ), sex hormone-binding globulin (WMD:  $15.22$  nmol/l;  $P = 0.0035$ ), luteinising hormone (WMD:  $-3.97$  U/L;  $P = 0.0008$ ) and luteinising hormone:follicle-stimulating hormone ratio (WMD:  $-1.04$ ;  $P = 0.0053$ ), but not for follicle-stimulating hormone levels (WMD:  $1.23$  mUI/ml;  $P = 0.12$ ). Significant changes in metabolic markers were seen in blood glucose (WMD:  $-9.65$  mg/dl;  $P = 0.0031$ ), insulin (WMD:  $-2.41$  mg/dl;  $P = 0.0387$ ), homeostatic model assessment for insulin resistance (WMD:  $-2.46$ ;  $P = 0.0123$ ) and TAG (WMD:  $-29.95$  mg/dl;  $P = 0.0188$ ). **The very low-carbohydrate ketogenic diet shows significant benefits in managing body composition, reducing hyperandrogenism, balancing sex hormones and improving glucose metabolism in polycystic ovary syndrome.**



# Effects of dietary fat and fiber on plasma and urine androgens and estrogens in men: a controlled feeding study<sup>1,2</sup>

Joanne F Dorgan, Joseph T Judd, Christopher Longcope, Charles Brown, Arthur Schatzkin, Beverly A Clevidence, William S Campbell, Padmanabhan P Nair, Charlene Franz, Lisa Kahle, and Philip R Taylor

**ABSTRACT** We conducted a controlled feeding study to evaluate the effects of fat and fiber consumption on plasma and urine sex hormones in men. The study had a crossover design and included 43 healthy men aged 19–56 y. Men were initially randomly assigned to either a low-fat, high-fiber or high-fat, low-fiber diet for 10 wk and after a 2-wk washout period crossed over to the other diet. The energy content of diets was varied to maintain constant body weight but averaged  $\approx 13.3$  MJ (3170 kcal)/d on both diets. The low-fat diet provided 18.8% of energy from fat with a ratio of polyunsaturated to saturated fat (P:S) of 1.3, whereas the high-fat diet provided 41.0% of energy from fat with a P:S of 0.6. Total dietary fiber consumption from the low- and high-fat diets averaged 4.6 and 2.0 g  $\cdot$  MJ<sup>-1</sup>  $\cdot$  d<sup>-1</sup>, respectively. Mean plasma concentrations of total and sex-hormone-binding-globulin (SHBG)-bound testosterone were 13% and 15% higher, respectively, on the high-fat, low-fiber diet and the difference from the low-fat, high-fiber diet was significant for the SHBG-bound fraction ( $P = 0.04$ ). Men's daily urinary excretion of testosterone also was 13% higher with the high-fat, low-fiber diet than with the low-fat, high-fiber diet ( $P = 0.01$ ). Conversely, their urinary excretion of estradiol and estrone and their 2-hydroxy metabolites were 12–28% lower with the high-fat, low-fiber diet ( $P \leq 0.01$ ). Results of this study suggest that diet may alter endogenous sex hormone metabolism in men. *Am J Clin Nutr* 1996;64: 850–5.

**KEY WORDS** Diet, dietary fats, dietary fiber, estrogen, androgens

intake, and the association was primarily due to animal fat consumption (8).

Prostate cancer is a hormone-dependent cancer and a current hypothesis is that diet modifies prostate cancer risk through an effect on the sex hormones (10, 11). Male vegetarians have been reported to have lower plasma testosterone and estradiol concentrations than omnivores (12), and dietary fat and fiber have been correlated with sex hormone concentrations in several studies (12–14). Specific diet-hormone relations reported in men include positive correlations of testosterone with polyunsaturated fat (13) and dihydrotestosterone with vegetable fat consumption (14) and inverse correlations of testosterone and estradiol with fiber intake (12). However, comparisons between vegetarians and omnivores and correlations of specific dietary components with hormones from different studies have been inconsistent (12–17). Moreover, in three diet intervention studies (18–20), serum or urine testosterone levels were depressed with the low-fat, high-fiber or vegetarian diet, but findings for other hormones were inconsistent.

In 1986 the National Cancer Institute and Beltsville Human Nutrition Research Center, Agriculture Research Service, conducted a controlled feeding study in men to evaluate the effect of modifying dietary fat and fiber intakes on several indexes potentially related to cancer or atherosclerosis, including plasma lipoproteins (21), prostaglandins (22), fecal mutagens, and hormones. As part of this study, we evaluated the effect of these dietary components on plasma and urine androgens and estrogens.

lowering of circulating androgen levels without changing the



Volume 90, Issue 6  
1 June 2005

Article Content

Subjects and Methods

Results

Discussion

Acknowledgments

Abbreviations:

References

< Previous Next >



Case Reports

Case Series

Cohort Studies

Pragmatic Cohort

RCT

Meta-Analysis

Widespread Clinical Use

Cardiometabolic

Type 2 Diabetes

Obesity

Metabolic Syndrome

Hypertension

Fatty Liver

CVD/Heart Failure

Hormonal

PCOS

Hypotestosteronism

Seizure

Neuropsychiatric

Depression

Binge Eating / Food Addiction

Bipolar

Schizophrenia



NEUROLOGY AND NEUROSURGERY

# The Epilepsy Center

[Home](#) > [Neurology and Neurosurgery](#) > [Specialty Areas](#) > [Epilepsy](#)

## Ketogenic Diet Therapy for Epilepsy

One of the oldest treatments for [epilepsy](#) is the classic **ketogenic diet**, which involves consuming high-fat foods and very few carbohydrates. Johns Hopkins is a longstanding pioneer in this mode of therapy.

In order to be successful, this therapy calls for strict compliance and plenty of patience, especially in the beginning. Most important, patients with epilepsy should only use the diet with the support of a knowledgeable ketogenic diet team, including a doctor and a licensed dietitian who can correctly calculate and monitor the diet for each individual.

## Ketogenic Diet for Epilepsy: Why Choose Johns Hopkins

Cardiometabolic

Hormonal

Neuropsychiatric



[Home](#) | [JAMA Psychiatry](#) | [Vol. 83, No. 1](#)

## Original Investigation

# Ketogenic Diets and Depression and Anxiety A Systematic Review and Meta-Analysis

Reinhard Janssen-Aguilar, MD<sup>1,2</sup>; Tulassi Vije, BHSc<sup>1</sup>; Malika Peera, BHSc<sup>1</sup>; [et al](#)

» [Author Affiliations](#) | [Article Information](#)

☰ RELATED ARTICLES

## Key Points

**Question** What are the associations between ketogenic diets (KDs) and mental health outcomes in adults, particularly depressive and anxiety symptoms?

**Findings** In this systematic review and meta-analysis of 50 studies, KDs were associated with small to moderate improvements in depressive symptoms in randomized clinical trials and with larger improvements in quasi-experimental studies. No significant associations were found for anxiety in randomized clinical trials.

**Meaning** KDs show potential associations with reduced depressive symptoms, but evidence for anxiety is inconclusive; larger, high-quality trials are needed to clarify effectiveness and generalizability.

Cardiometabolic

Hormonal

Neuropsychiatric

JAMA P

RCT: A Ketoge

### POPULATION

26 Men, 61 Women  
1 Nonbinary



Adults with treatment-resistant depression with a Patient Health Questionnaire-9 (PHQ-9) score of 10 or higher. Mean age, 42.1 y

### SETTINGS / LOCATIONS



88 Remissions across trials

Gao M, Kirk M, Knight H, et al. Published online February 1, 2025.



Check for updates

OPEN ACCESS

EDITED BY  
Kwok Kit Tong,  
University of Macau, China

REVIEWED BY  
Samer El Hayek,  
American Center for Psychiatry and  
Neurology, United Arab Emirates  
Georgia Ede,  
Independent Researcher, Northampton,  
MA, United States  
Amy Berger,  
Private Practice Nutritionist, Staunton,  
United States

\*CORRESPONDENCE  
Laura Buchanan  
✉ laura@toward.health

RECEIVED 15 April 2025  
ACCEPTED 02 July 2025  
PUBLISHED 24 July 2025

CITATION  
Saner E, Kalayjian T, Buchanan L,  
Calkins M, Soto-Mota A, Jun D  
and Sethi S (2025) TOWARD: a metabolic  
health intervention that improves food  
addiction and binge eating symptoms.  
*Front. Psychiatry* 16:1612551.  
doi: 10.3389/fpsy.2025.1612551

COPYRIGHT  
© 2025 Saner, Kalayjian, Buchanan, Calkins,  
Soto-Mota, Jun and Sethi. This is an open-  
access article distributed under the terms of  
the Creative Commons Attribution License  
(CC BY). The use, distribution or reproduction  
in other forums is permitted, provided the  
original author(s) and the copyright owner(s)  
are credited and that the original publication  
in this journal is cited, in accordance with  
accepted academic practice. No use,  
distribution or reproduction is permitted  
which does not comply with these terms.

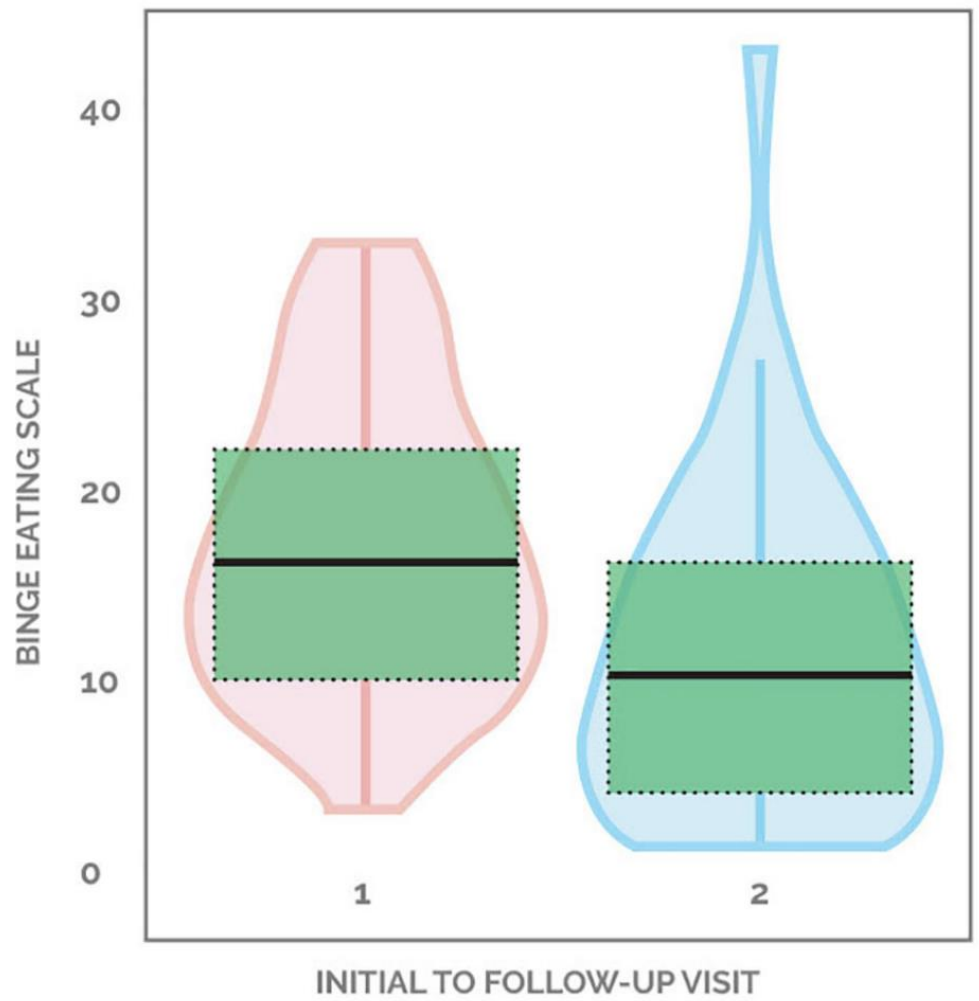


FIGURE 2  
Violin plot of change in BES scores pre- and post-intervention.

Widespread  
Clinical Use

health  
loves

ns

46%

PATIENTS IMPROVED  
FOOD ADDICTION  
SYMPTOMS



ARD



Case Reports

Case Series

Cohort Studies

Pragmatic Cohort

RCT

Meta-Analysis

Widespread Clinical Use

Cardiometabolic

Type 2 Diabetes

Obesity

Metabolic Syndrome

Hypertension

Fatty Liver

CVD/Heart Failure

Hormonal

PCOS

Hypotestosteronism

Seizure

Neuropsychiatric

Depression

Binge Eating / Food Addiction

Bipolar

Schizophrenia

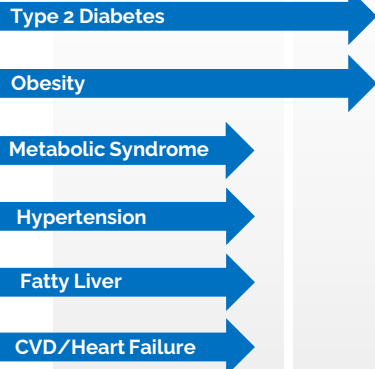




# Percent Offered Metabolic Nutrition Therapy

0%                      <20%

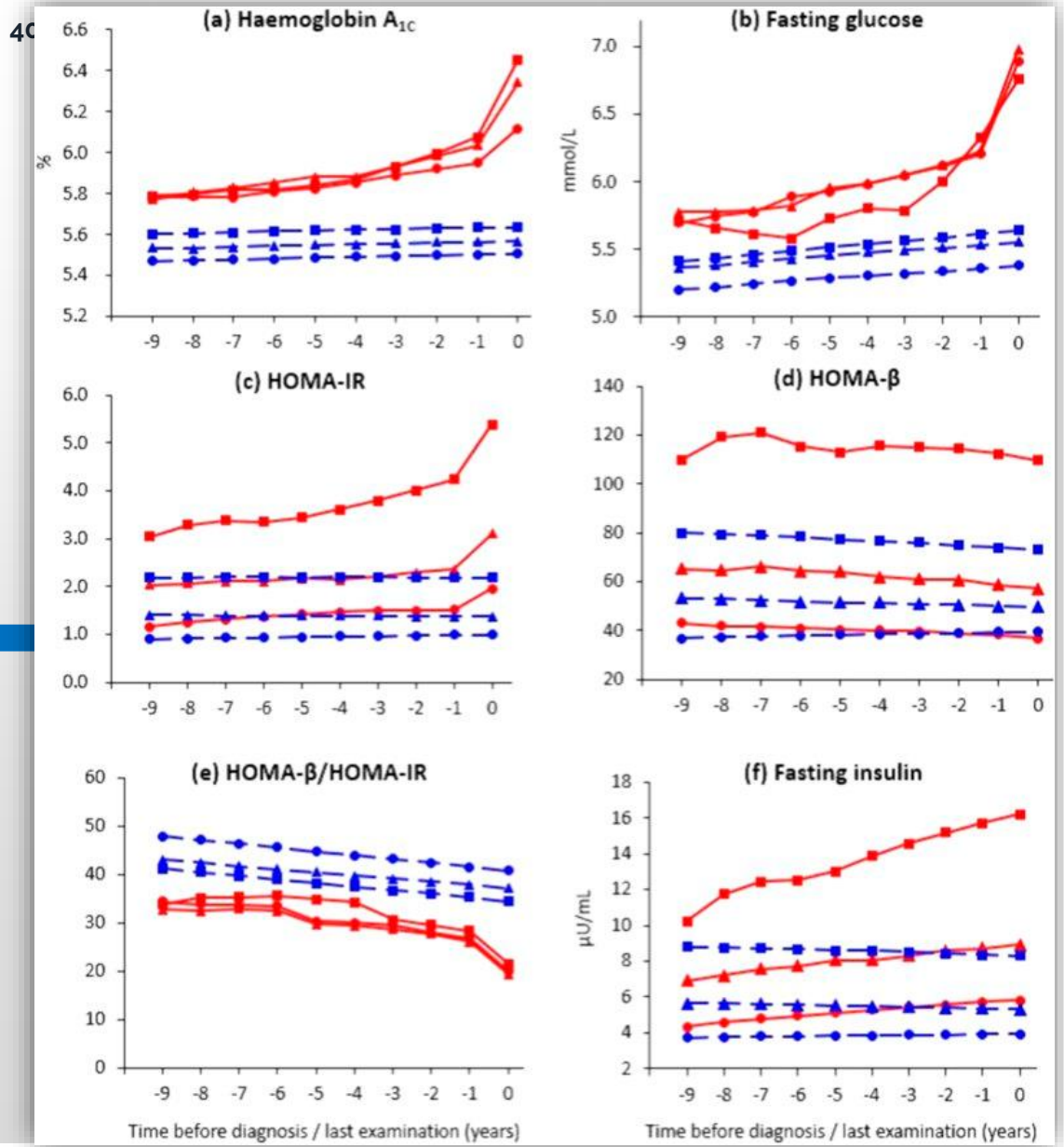
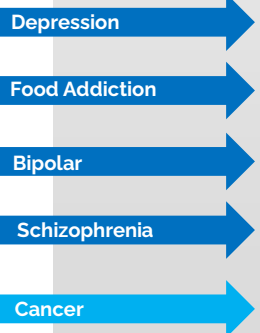
## Cardiometabolic



## Hormonal

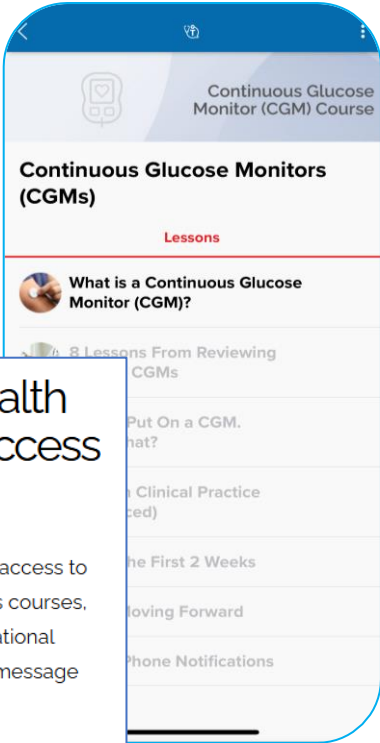


## Neuropsychiatric



# Getting Started

Scan QR code to receive premium access to the Toward Health app with a self-guided CGM course



## Start with the Toward Health App for Free Premium Access

Swiss Re is offering you 12-month Free premium access to the Toward Health app. Premium access includes courses, challenges, live weekly meetings, articles, educational videos, exclusive community & private Swiss Re message boards and more.

Enter Your Email Address

Click the button below to get your FREE App Access.

Get Your FREE Access

Download the App



# Contact

Dr. Tro Kalayjian, D.O.

845-397-2873  
[toward.health](https://toward.health)

