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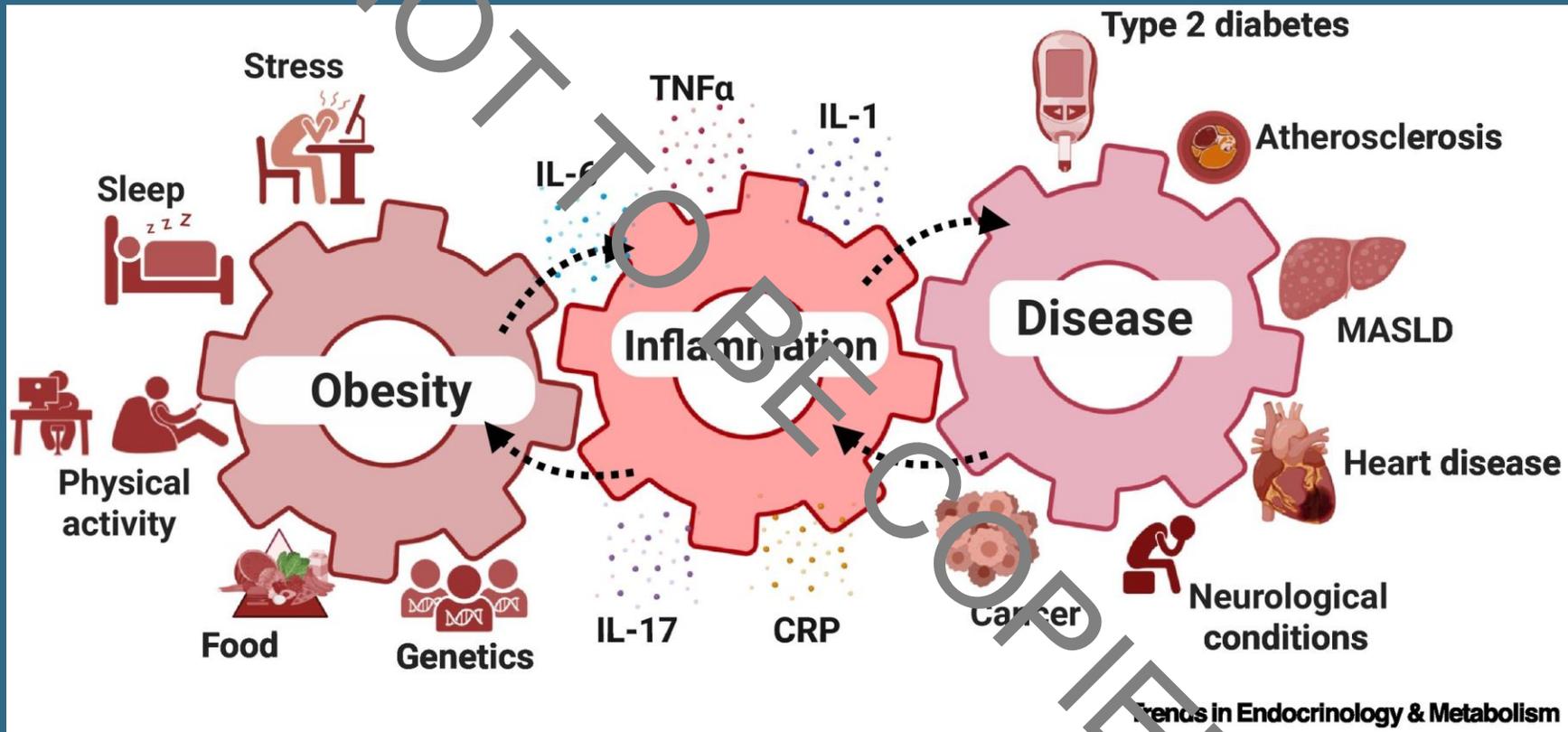
40:00

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Education is not a
singular inoculation



A change of science is a foot.....



...thus a change of GP LTC review infrastructure is a must

CARDIOVASCULAR-KIDNEY-METABOLIC REVIEW

THE NEW LONG TERM CONDITION FRAMEWORK OF CARE FOR OUR HOUSES OF CARE

Northern Ireland Primary Care Diabetes and Obesity Society

25th September 2025

Dr Eimear Darcy

NHS GP Partner

MB BCH BAO DMH MRCGP (2017)

PG Cert Diabetes (Distinction)

SCOPE certified in Obesity Management.

General Practitioner specialising in CKM

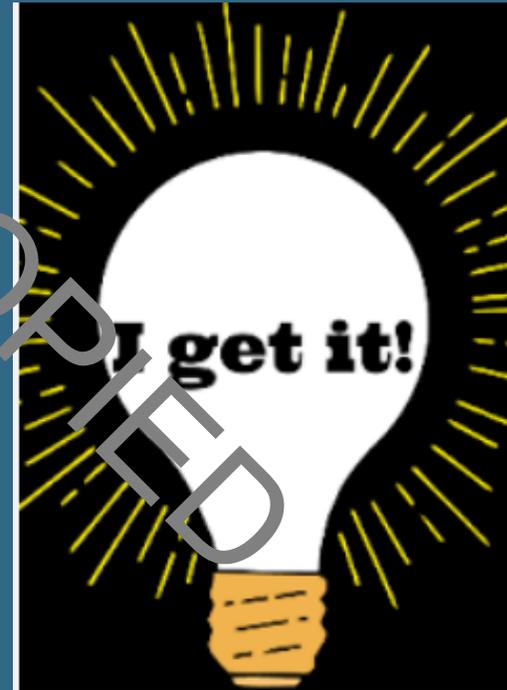


Learning Objectives

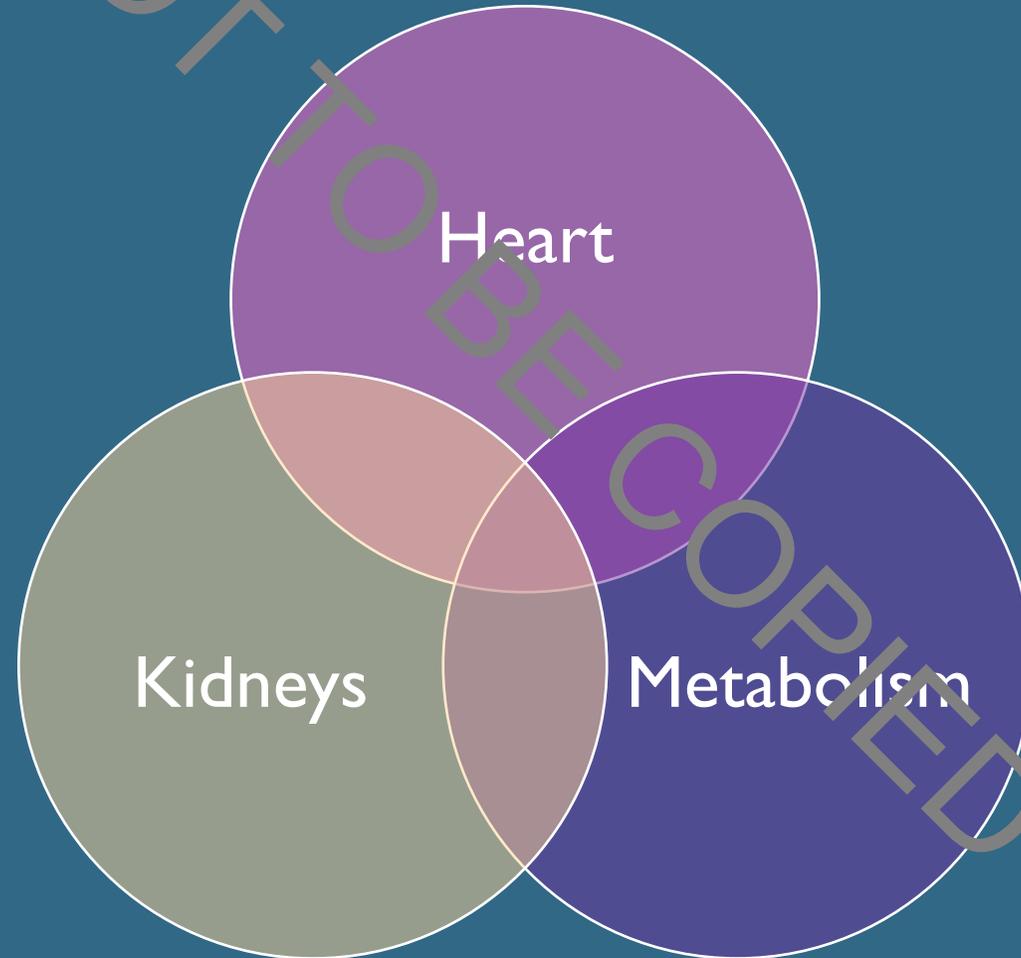
OFFICIALLY..

Insights into modern
management of CKM
Long Term Conditions

PERSONALLY...



CVRM = CKM = CAREME/CRM

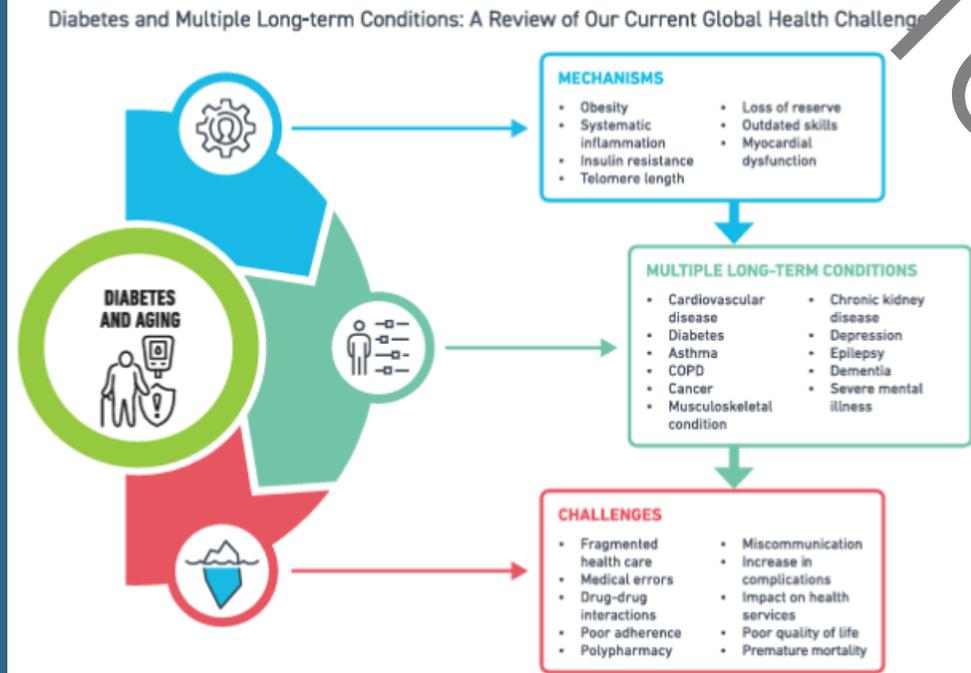


Multiple Long Term Conditions (MTLCs)

Diabetes and Multiple Long-term Conditions: A Review of Our Current Global Health Challenge

Kamlesh Khunti, Yogini V. Chudasama, Edward W. Gregg, Monika Kamkuemah, Shirazni Misra, Jerrisa Suls, Nikhil S. Venkateshmurthy, and Jonathan Valabhji

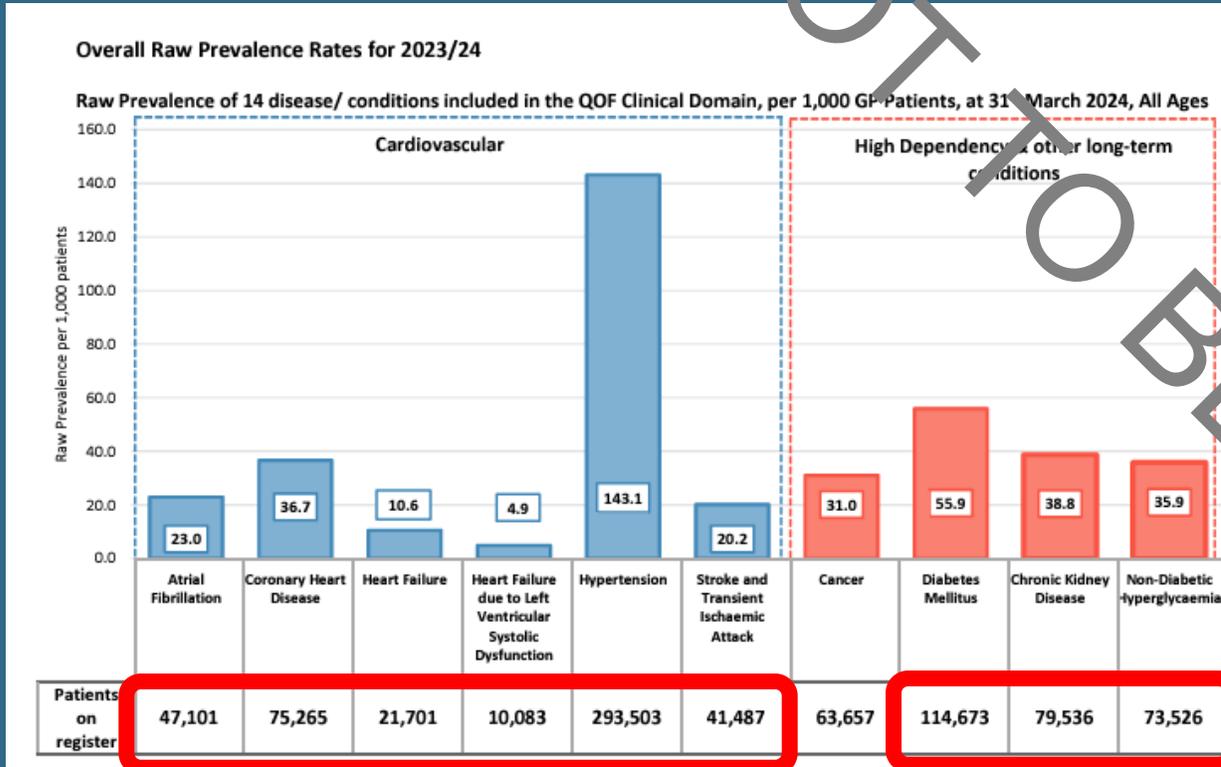
Diabetes Care 2023;46(12):2092–2101 | <https://doi.org/10.2337/dci23-0035>



Heart failure and chronic kidney disease manifestation and mortality risk associations in type 2 diabetes: A large multinational cohort study

Conclusion: In a large multinational study of >750 000 CVD-free patients with T2D, HF and CKD were consistently the most frequent first cardiovascular disease manifestations and were also associated with increased mortality risks. These novel findings show these cardiorenal diseases to be important and serious complications requiring improved preventive strategies.

Northern Ireland overall raw prevalence rates for 2023/24

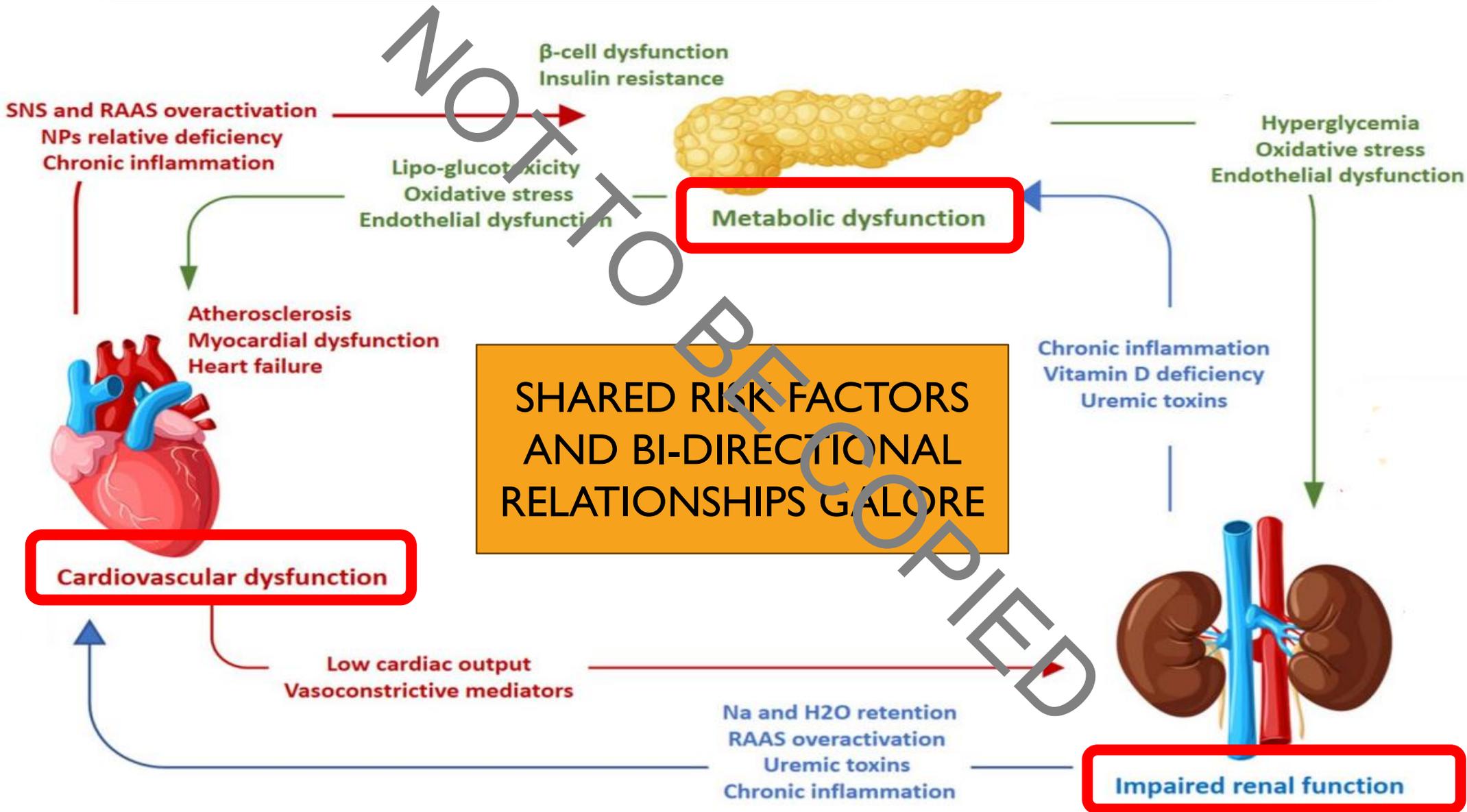


	2013/14	2023/24
Diabetes	81,867	114,673

Difference of **32806** people

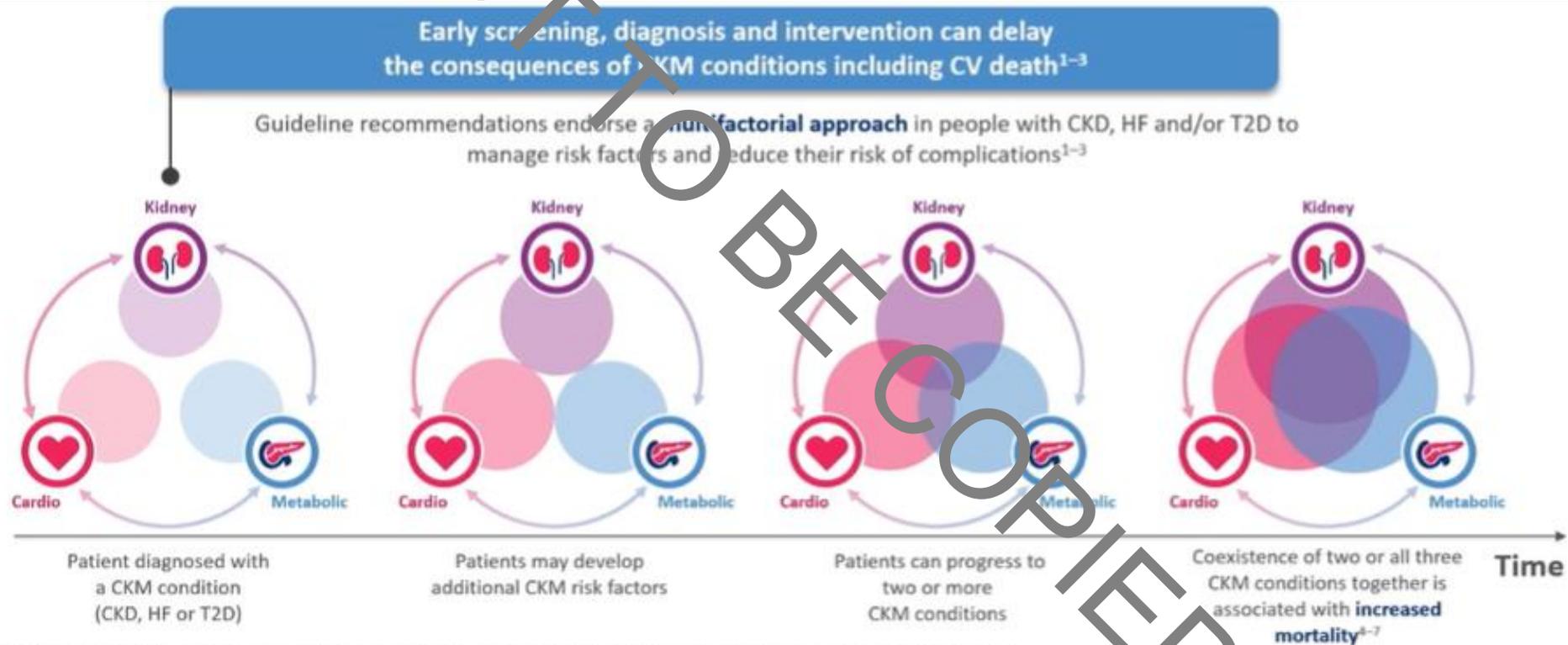
The minority of our population is not living with overweight or obesity (2)

CKM syndrome - Groundhog day for disease



Adapted from Marassi, M., Fadini, G.P. The cardio-renal-metabolic connection: a review of the evidence. *Cardiovasc Diabetol* **22**, 195 (2023) [accessed 2/9/25]

Early and multifactorial intervention is the best way to help protect by reducing the amplifying risks associated with interconnected CKM diseases



CKD, chronic kidney disease; CRM, cardiovascular, renal and metabolic; CV, cardiovascular; HF, heart failure; T2D, type 2 diabetes

1. Kidney Disease: Improving Global Outcomes (KDIGO) CKD Work Group. *Kidney Int* 2022;102:S1; 2. Heidenreich PA et al. *J Am Coll Cardiol* 2022;79:263-81. American Diabetes Association. *Diabetes Care* 2023;46:S1;

4. Birkeland KI et al. *Diabetes Obes Metab* 2020;22:1607; 5. Ather S et al. *J Am Coll Cardiol* 2012;59:998; 6. Afkarian M et al. *J Am Soc Nephrol* 2013;24:302; 7. Lovre D et al. *Endocrinol Metab Clin North Am.* 2018;47:237

CARDIOVASCULAR-KIDNEY-METABOLIC SYNDROME

Health disorder due to connections amount

- Heart disease
- Kidney Disease
- Type Two Diabetes
- Obesity

Leading to poor health outcomes

Circulation

Volume 148, Issue 20, 14 November 2023; Pages 1606-1635

<https://doi.org/10.1161/CIR.0000000000001184>



AHA PRESIDENTIAL ADVISORIES

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

CKM syndrome – medical definition

‘...a systemic disorder characterised by pathophysiological interactions among metabolic risk factors, CKD, and CV system leading to multi-organ dysfunction and a high rate of adverse CV outcomes’.

Includes individuals:

At risk for CVD disease due to metabolic risk factors.

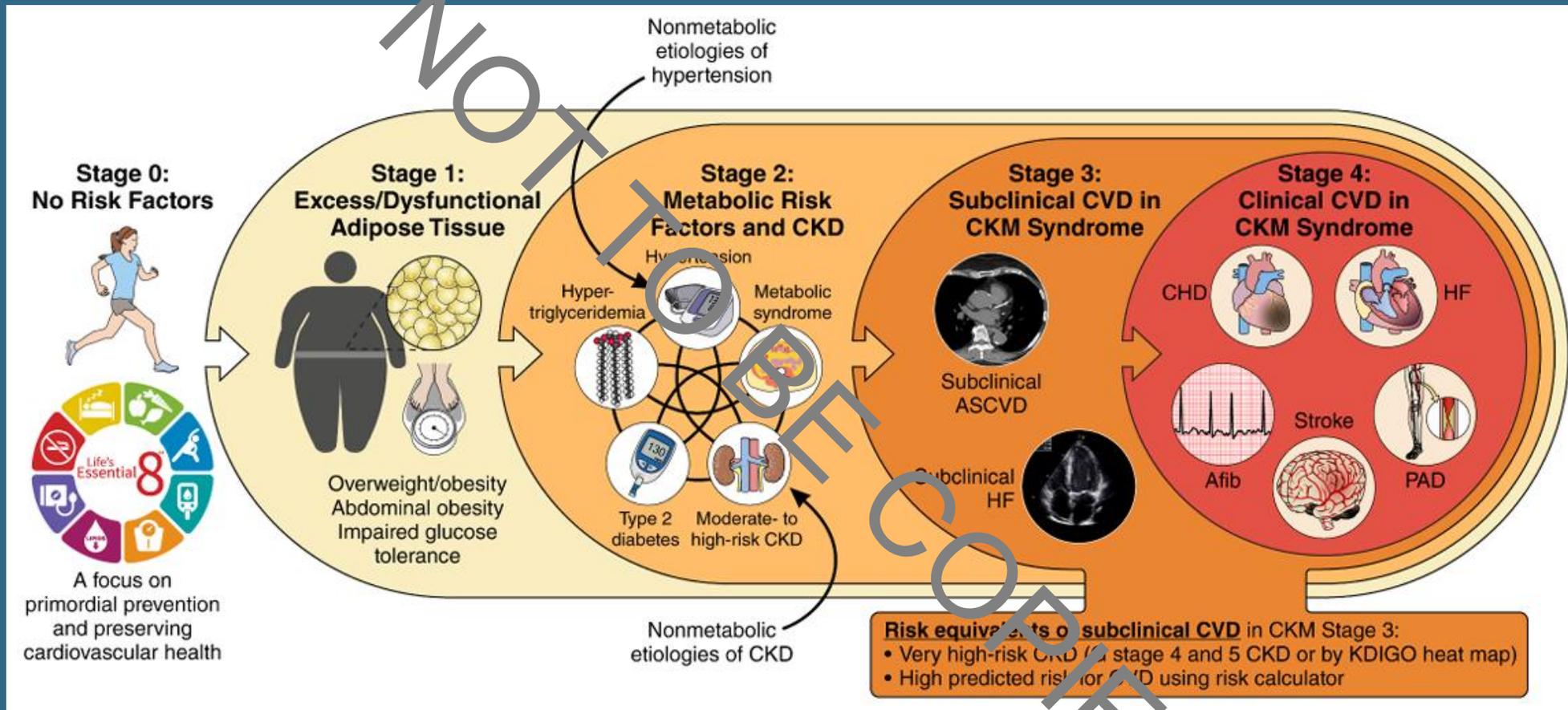
Those with CVD

Those with CKD

General Practice CKM Review along the life journey



..AND IN THE CONTEXT OF THE PERSONS LIFE STORY



Every

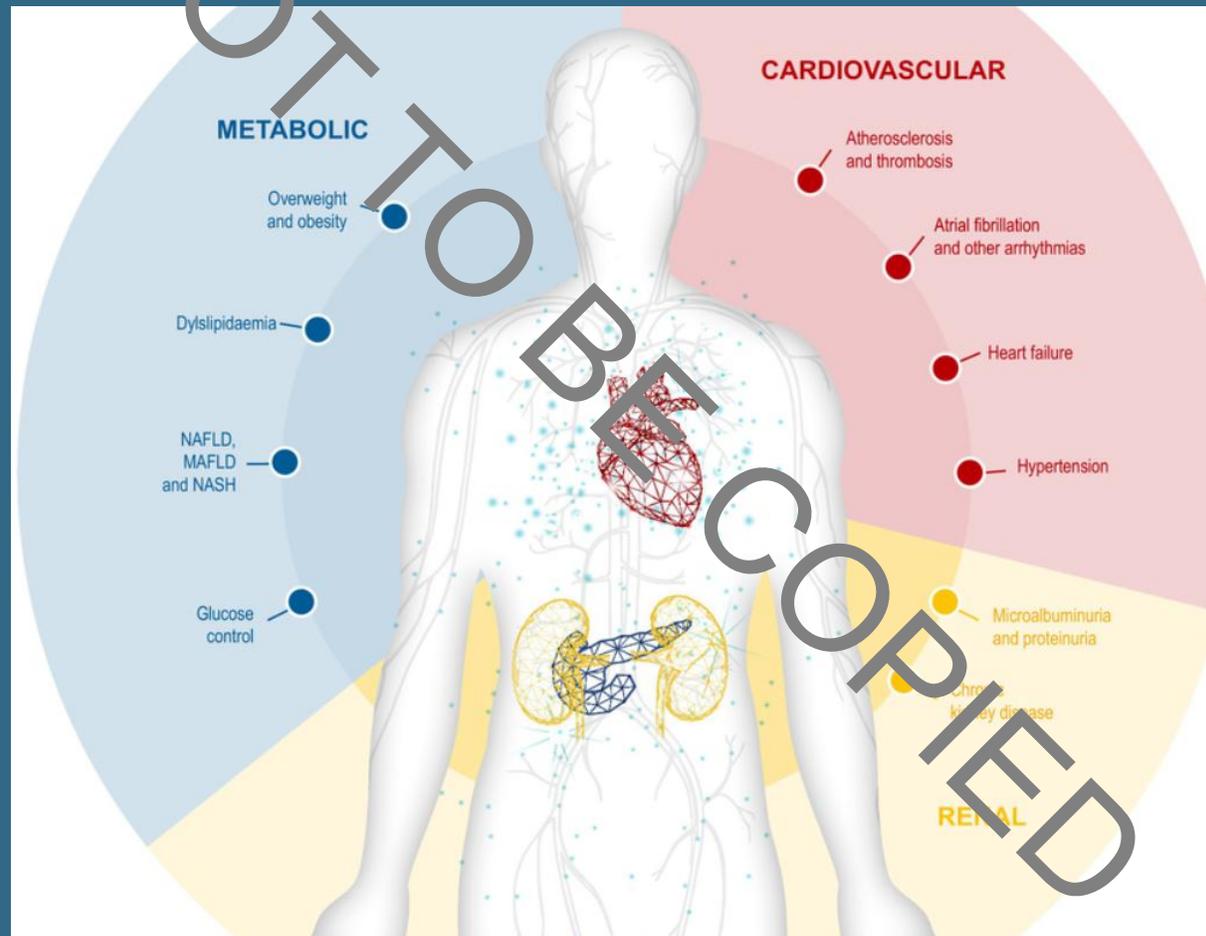
stage

sits

in

General Practice

CKM – CARE OF PEOPLE , NOT JUST PATHOLOGY



HOLISTIC IMPLEMENTATION

holistic

adjective

UK  /hə'ɪs.tɪk/ US  /hoʊl'ɪs.tɪk/

Add to word list 

dealing with or treating the whole of something or someone and not just a part:



Cambridge
Dictionary

Clinical staging to guide management of metabolic disorders and their sequelae: a European Atherosclerosis Society consensus statement

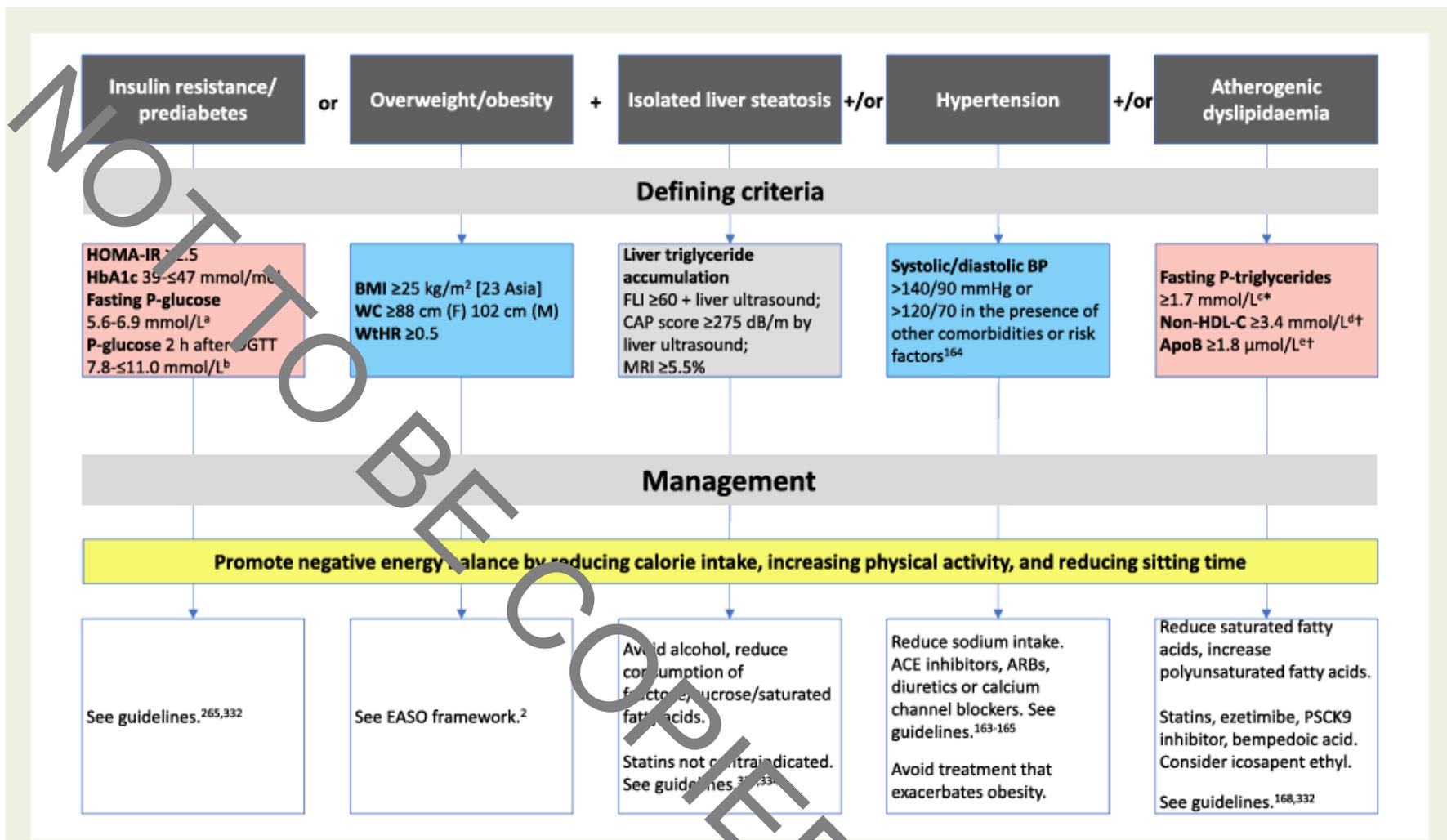


Figure 4 Defining criteria and potential management strategies for systemic metabolic disorder Stage 1. Conventional units: ^a101–124 mg/dL; ^b141 to ≤ 198 mg/dL; ^c150 mg/dL; ^d131 mg/dL; ^e100 mg/dL. *Non-fasting P-triglycerides ≥ 2.0 mmol/L (177 mg/dL). †Values shown for individuals at moderate risk of atherosclerotic cardiovascular disease. For those at high risk of atherosclerotic cardiovascular disease, lower thresholds apply: non-HDL-cholesterol ≥ 2.6 mmol/L (100 mg/dL); apolipoprotein B ≥ 1.5 $\mu\text{mol/L}$ (≥ 80 mg/dL). P, plasma; WC, waist circumference; WtHR, waist-to-height ratio; EASO, European Association for the Study of Obesity; FLI, fatty liver index; CAP, controlled attenuation parameter; MRI, magnetic resonance imaging; BP, blood pressure; C, cholesterol; ACE, angiotensin-converting enzyme; ARB, angiotensin-II receptor blocker



<https://doi.org/10.1038/s43856-025-00742-9>

Multiple long-term conditions as the next transition in the global diabetes epidemic

Updates

Edward W. Gregg^{1,2}, Naomi Holman^{1,2,3}, Marisa Sophiea², Shivani Misra^{4,5}, Jonathan Pearson-Stuttard^{4,6}, Jonathan Valabhji^{7,8,9} & Kamlesh Khunti^{8,9}

Several transitions, or new patterns and dynamics in the contributors and health outcomes, have altered the character and burden of the multi-decade, worldwide growth in prevalence of type 2 diabetes (T2DM). These changes have led to different needs for prevention and care. These dynamics have been driven by diverse demographic, socio-economic, behavioural, and health system response factors. In this Perspective, we describe these transitions and how their attributes have set the stage for multimorbidity, or multiple long-term conditions (MLTCs), to be the next major challenge in the diabetes epidemic. We also describe how the timing and character of these stages differ in high-, middle-, and low-income countries. These challenges call for innovation and a stronger focus on MLTCs across the spectrum of cause, effectiveness, and implementation studies to guide prevention and treatment priorities.

The global pandemic of type 2 diabetes (T2DM) is often viewed as a long-term by-product of the great epidemiologic transition, wherein in the previous century, human patterns of mortality and disease exchanged high rates of mortality due to infections and maternal and child mortality for lower mortality rates driven by chronic conditions such as cardiovascular disease (CVD), cancer, and T2DM^{1,2}. However, there have been differing changes in incidence for specific chronic conditions. For example, deaths due to CVD decreased beginning in the 1970s and 1980s whereas T2DM, lacking effective policy or health care options for prevention, accelerated in the 1990s^{3,4}. Although recent signs of a decrease in T2DM incidence in some countries have emerged^{5,6}, most countries of the world continue to experience unrelenting growth of the problem^{7,8}.

The T2DM pandemic has itself been complex and multi-phasic. Underlying the overall increases in prevalence, have been several transitions across distinct periods of the T2DM epidemic that impacted the burden and approaches to care, prevention, and research^{9,10}. In this Perspective, we explore how these transitions have set the stage for multiple long-term conditions (MLTC, or multimorbidity) impacting morbidity, health services, and approaches to prevent T2DM.

Seeding and steady growth of the problem

The first phase of growth in the T2DM epidemic, evident in the 1970s and 1980s, consisted of steady, gradual, increases in prevalence in high income

countries (HICs) and more rapid increases in selected low- and middle-income countries (LMICs) undergoing rapid urbanization, nutritional and other sociocultural changes associated with westernization¹¹⁻¹³ (Fig. 1). This was associated with a global nutrition transition, with dietary shifts toward refined carbohydrates, added sweeteners and edible oils, and away from legumes and fresh fruits and vegetables. Declining physical activity levels contributed further and collectively they are cited as driving an environment conducive to increases in obesity and T2DM^{14,15}. However, where data were available, diagnosed diabetes prevalence rates were quite low, at around 2–4% in the US, UK, and Northern Europe^{16,17} compared to recent global estimates, which have surpassed 10% of adults¹⁸. Further, this low diabetes prevalence contrasted with very high rates of severe complications, including a 20-fold increased rate of lower extremity amputation, 10 times the rate of end stage renal disease (ESRD), and three times the risk of CVD¹⁹⁻²¹. These high complication risks were attributed to suboptimal diabetes care, risk factors and self-management practices. For example, estimates from the some HICs in the mid-1990s showed that large segments of the population with diagnosed diabetes had HbA1c levels well over 9% (75 mmol/mol) and blood pressure levels > 140/80, levels considered to be a sign of inadequate management by current standards²²⁻²⁵. Thus, perception of the problem centred more on the complications accompanying T2DM rather than the incidence and potential prevention of T2DM.

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Summary of Major Transitions in the Diabetes Epidemic in High Income Countries, 1960-2020

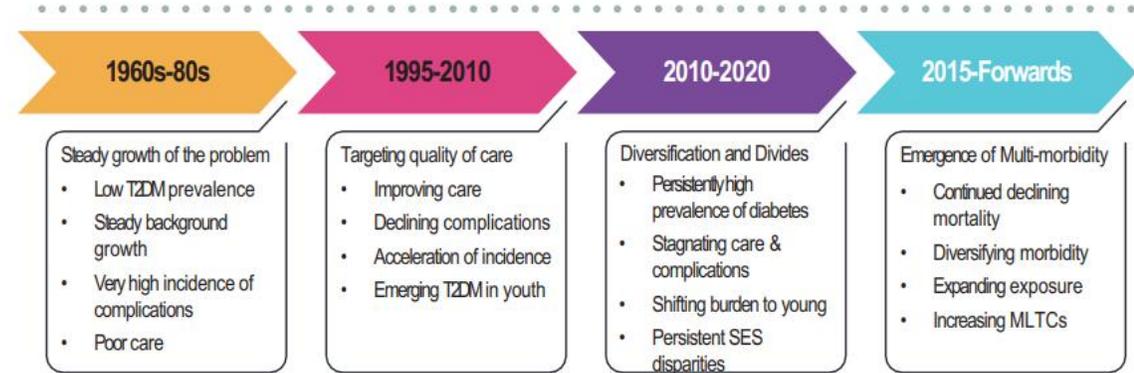


Fig. 1 | Summary of major transitions in the global diabetes epidemic, as experienced by high income countries, 1960–2020. The figure describes key patterns in prevalence, incidence, complications, care, and socio-demographic factors

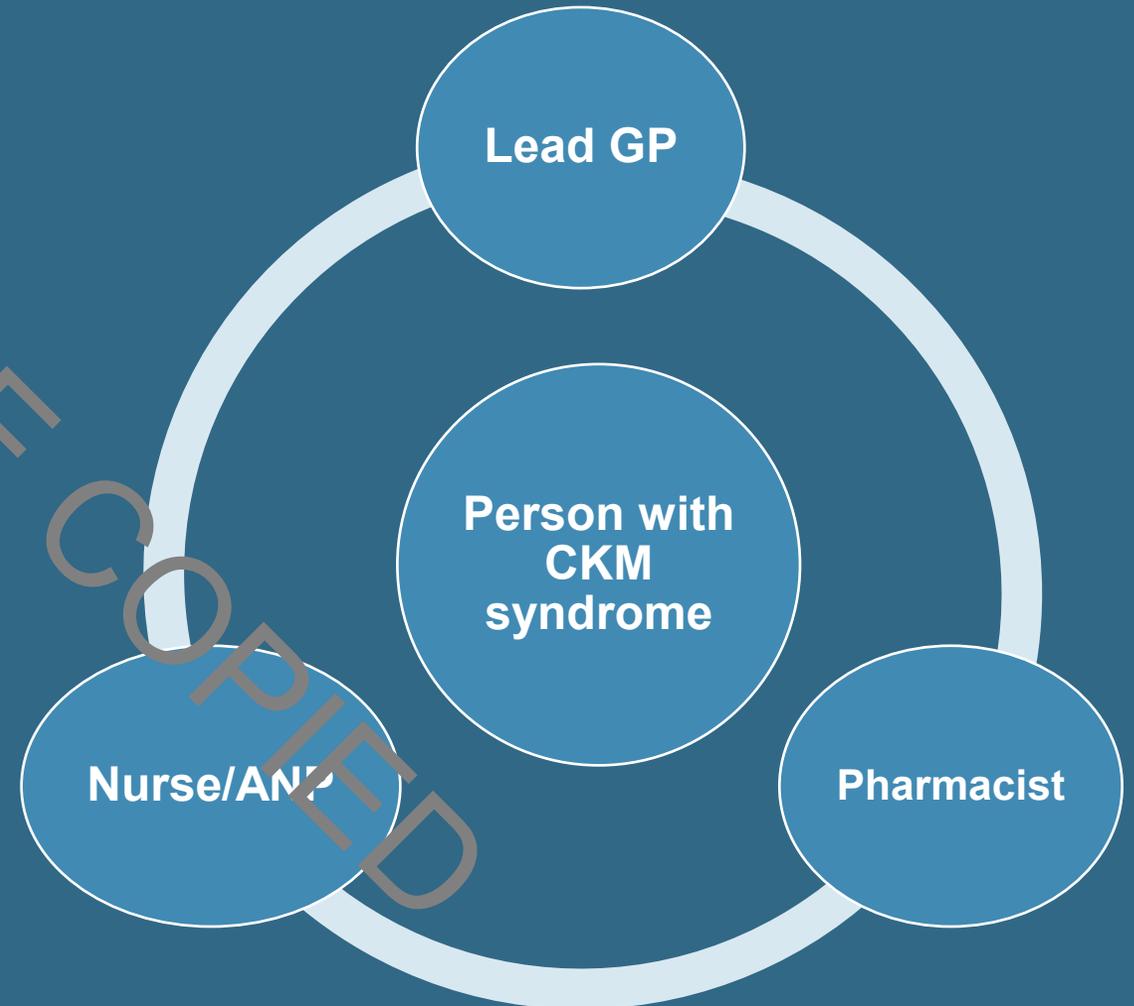
affecting the T2DM pandemic. T2DM: Type 2 diabetes mellitus; DM: diabetes mellitus; SES: socio-economic status; MLTC: multiple long-term conditions

‘if MLTCs are the next major transition in the diabetes epidemic, and the key drivers lie at both ends of the age spectrum as we suggest, the most practical priorities will lie in prevention, ongoing or routine care models and coordination,

i.e GENERAL PRACTICE, THE WIDER PRIMARY CARE.....and secondary care ongoing care.

Grange Family Practice CKM team

#CKM thought processes
are a future cornerstone of
General Practice and the
wider Primary Care Teams
Long Term Condition Care.



Global Effect of Cardiovascular Risk Factors on Lifetime Estimates

The Global Cardiovascular Risk Consortium

ABSTRACT

BACKGROUND

Five risk factors account for approximately 50% of the global burden of cardiovascular disease. How the presence or absence of classic risk factors affects lifetime estimates of cardiovascular disease and death from any cause remains unclear.

METHODS

We harmonized individual-level data from 2,078,948 participants across 133 cohorts, 39 countries, and 6 continents. Lifetime risk of cardiovascular disease and death from any cause was estimated up to 90 years of age according to the presence or absence of arterial hypertension, hyperlipidemia, underweight and overweight or obesity, diabetes, and smoking at 50 years of age. Differences in life span (in terms of additional life-years free of cardiovascular disease or death from any cause) according to the presence or absence of these risk factors were also estimated. Risk-factor trajectories were analyzed to predict lifetime differences according to risk-factor variation.

RESULTS

The lifetime risk of cardiovascular disease was 24% (95% confidence interval [CI], 21 to 30) among women and 38% (95% CI, 30 to 45) among men for whom all five risk factors were present. In the comparison between participants with none of the risk factors and those with all the risk factors, the estimated number of additional life-years free of cardiovascular disease was 13.3 (95% CI, 11.2 to 15.7) for women and 10.6 (95% CI, 9.2 to 12.9) for men; the estimated number of additional life-years free of death was 14.5 (95% CI, 9.1 to 15.3) for women and 11.8 (95% CI, 10.1 to 13.6) for men. As compared with no changes in the presence of all risk factors, modification of hypertension at an age of 55 to less than 60 years was associated with the most additional life-years free of cardiovascular disease, and modification of smoking at an age of 55 to less than 60 years was associated with the most additional life-years free of death.

CONCLUSIONS

The absence of five classic risk factors at 50 years of age was associated with more than a decade greater life expectancy than the presence of all five risk factors, in both sexes. Persons who modified hypertension and smoking in midlife had the most additional life-years free of cardiovascular disease and death from any cause, respectively. (Funded by the German Center for Cardiovascular Research [DZHK]; ClinicalTrials.gov number, NCT05466825.)

LOOK ALIVE – THINK FIVE!

- 5 **modifiable** risk factors account for ~50% global CVD burden.....
- 50% of all CVD cases could potentially be **prevented** through **effective risk-factor management**.
- **Fatal 5 : Smoking ; Blood Pressure ; Cholesterol ; Diabetes ; obesity.**
- If you're 50 and five –less....
- Women – CVD 13 yrs later ; death 14 yrs later
- Men – CVD 10 yrs later ; death 11 years later
- Behavioural change has impact!
 - HTN Mx & smoking cessation in midlife = longer life and more years CVD disease free.

Bottom line: Hope lies in our houses of care ; sustain life locally – look for and act on the five.

BP threshold for treatment

confirm : BP \geq 135/85 mmHg 7-day average HBPM or day-time average ABPM

start : pharmacological therapy on top of diet and lifestyle interventions irrespective of CVD risk.

On-treatment BP target

< 130/80 mmHg (measured by 7-day average HBPM or day-time average ABPM or office BP*)

or ALARA without causing unacceptable side-effects, and within 6-months of initiating treatment **for all adults.**

Exceptions:

frail and/or have limited life expectancy

use clinical judgement

CONSENSUS STATEMENT OPEN Check for updates

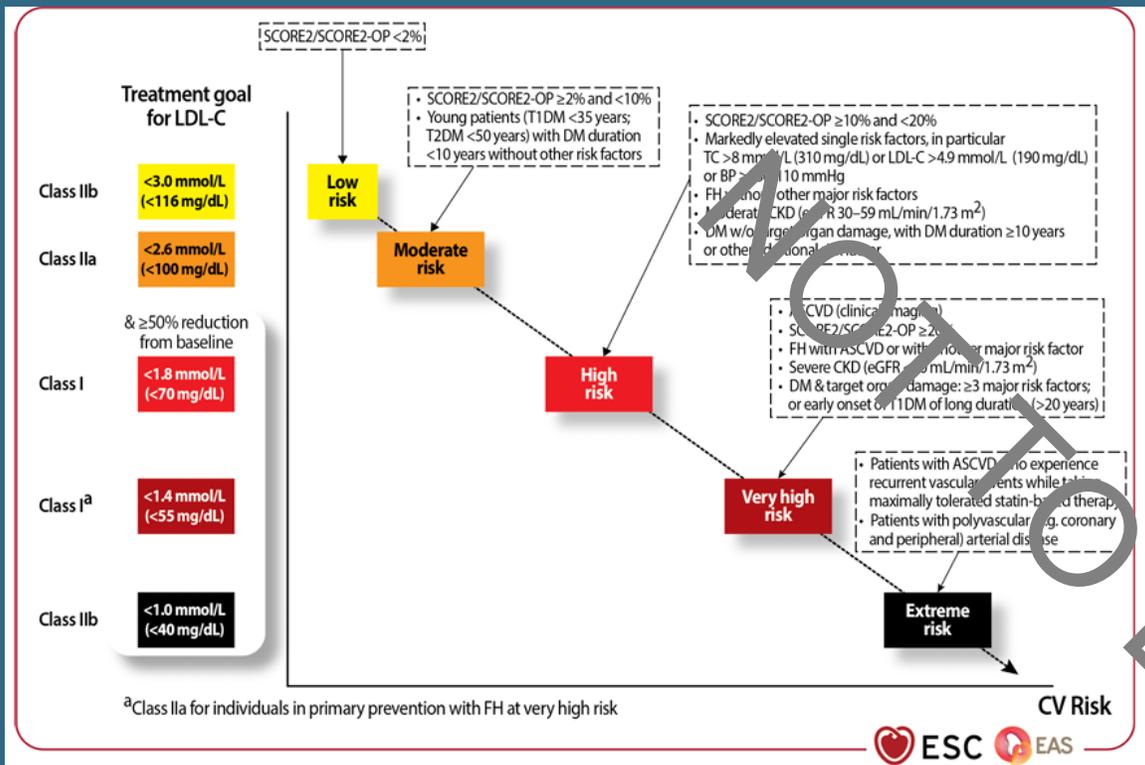
Call to action: British and Irish hypertension society position statement on blood pressure treatment thresholds and targets

Luca Faconti¹, Nayanatar Tantridge², Neil R. Poulter³, Jacob George⁴, Vikas Kapil⁵, Ajay Gupta⁶, Pauline A. Swift⁷, Anthony Heagerty⁸, Eduard Shalchiba⁹, Sarah Partridge¹⁰ and Ian B. Wilkinson²

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In this position statement the British and Irish Hypertension Society (BIHS) present a review of the current evidence for blood pressure (BP) treatment thresholds and targets. The BIHS recommend initiating pharmacological antihypertensive therapy, irrespective of cardiovascular disease risk, following a confirmed diagnosis of hypertension (sustained out-of-office BP \geq 135/85 mmHg despite diet and lifestyle advice). The BIHS recommend an on-treatment BP target < 130/80 mmHg or as low as reasonably achievable without causing unacceptable side-effects, within 6-months of initiating treatment, for all adults. Possible subgroups to whom this may not apply are those who are frail and/or have limited life expectancy where higher targets may be appropriate based on clinical judgement and the individuals' tolerance to treatment. The BIHS believe that this simple 2-step approach will facilitate practitioners deliver evidence-based best practice, discourage therapeutic inertia around BP lowering and improve health outcomes for all adults living with high BP.

Journal of Human Hypertension (2025) 39:537–540; <https://doi.org/10.1038/s41371-025-01055-z>



QRISK MISSES RISK

SCORE2 SAVES LIVES

EXTENT OF LIPID LOWERING WITH AVAILABLE THERAPIES

Approximate reduction in LDL-C

Statin dose mg/day	5	10	20	40	80
Fluvastatin			21%	27%	33%
Pravastatin		20%	24%	29%	
Simvastatin		27%	32%	37%	42%
Atorvastatin		37%	43%	49%	55%
Rosuvastatin	38%	43%	48%	53%	
Atorvastatin + Ezetimibe 10mg		52%	54%	57%	61%

RULE OF 6 SAVES TIME

MARY – TATT AND FLUSHING

- 50M - rare attender
- Last seen 2021 (last bloods 2019).
- Nil reg meds.
- NKDA.
- Nil PMH bar Mirena coil last year
- Non smoker
- Occasional Social Alcohol
- Office worker
- Sedentary lifestyle but tries
- Married 2 kids
- PC: Non-specific TATT ? menopause
- HPC flushing , brain fog , vaginal dryness, more irritable, further weight gain from baseline raised BMI.
- HRT assessment done
- Full systematic questioning and examination yielded **nil TATT red flags**
-
- Asked permission to discuss weight – at current highest weight – feels high hunger , low fullness and cravings. Poor sleep from flushing aggravating snoring at night.
- Pregnancies escalated baseline weight
- Always ‘a heavy teenager’
- Fhx of obesity. No medication culprits.
- Explored ideas , concerns , expectations
- Agreed start HRT , TATT bloods and review with results.

First do no harm

Be aware our weight bias and stigma as health care professionals is harming the patients we are trying to help.

Weight discrimination consequences (I) :

- Reduced health care use
- Avoidance of physical activity
- Low mood/reduced self esteem
- Increased binge eating

Kings Modified Obesity Staging and the 4 Ms Matter in Primary Care

Criterion	Stage 0 Normal health	Stage 1 At risk of disease	Stage 2 Established disease	Stage 3 Advanced disease
A Airway /Apnoea	Normal No snoring Neck circ < 43cm Epworth score <10	Mild sleep apnoea Mild snoring Epworth score ≥ 10 Mild OSA (dip rate<15/hr) Neck circ >43cm (size 18) Mild asthma	Severe CPAP Witnessed apnoea Dip rate ≥ 15/hr Uses CPAP (controlled) Severe asthma	Cor pulmonale Obesity hypoventilation syndrome Uncontrolled OSA
B BMI	<35 kg/m ²	35-50 kg/m ²	50-60 kg/m ²	>60 kg/m ²
C CVD risk	<10% CVD risk <10% over 10 years [JBS coronary risk prediction chart*]	10-20% CVD risk ≥10% over 10 years T2DM	>20% or stable heart disease Stable IHD CCF NYHA I-II, or >20% risk	Severe angina, or CCF NYHA III-IV
D Diabetes	Normal Fasting or random glucose <5.7 mmol/L Normal HbA1c	IFG IFG / IGT, or previous GDM	T2DM Diet, insulin or OHA controlled HbA1c<9%	Uncontrolled T2DM HbA1c>9% Advanced microvascular disease
E Economic complications	Normal Obesity has no financial impact	Financial impact Increased travel cost Increased clothes cost	Workplace disadvantage Earnings limited by obesity Receiving benefits due to obesity	Unemployed due to obesity Financial effect on 3 rd party (e.g. carer required to reduce income)
F Functional status & musculoskeletal	No limitation	Manages one flight of stairs Limitation on work or recreation	Cannot climb stairs (<1 flight) 3 rd party assistance for ADL or for dependents	Housebound Wheel chair user Registered disabled
G Gonadal & reproductive axis	Normal Normal sexual and reproductive function Celibate (not seeking physical relationship)	PCOS / ED PCOS Low testosterone (men) Impaired sexual function/ erectile dysfunction	Subfertility Subfertility or unable to access IVF Marital/ relationship breakdown due to obesity Cessation of all sexual activity	
H Health status (perceived)	Normal Good mental and physical well being	Low mood/poor QoL	Mild-moderate depression Takes treatment for depression	Severe depression Suicidal ideation Unmanaged substance abuse Active self harm
I Body Image & eating behaviour	Minimal or no concern Normal eating pattern	Dislikes mirror appearance Comfort eating Inappropriate eating cues Mild body image dysphoria	Avoids social interaction or mirrors Severe body image dysphoria Controlled eating disorder	Eating disorder Active eating disorder Social phobia
J Oesophago-gastric Junction	Normal, no GORD symptoms	GORD (acid reflux) controlled on standard PPI	Oesophagitis on OGD within 12 months Severe GORD symptoms: requires high dose PPI	Barrett's oesophagus
K Kidney	Normal	Proteinuria	eGFR <60	eGFR <30
L Liver	Normal	Elevated LFTs, NAFLD on ultrasound	NASH	Liver failure

Four M's	Characteristics included
 Mental:	Mood, depression, eating disorders, anxiety, personality, cognition, attention deficit, sleep quality, potentials traumas, addictions.
 Mechanical:	Musculoskeletal disorders, chronic pain, OSA, GERD, urinary incontinence, intertrigo, thrombosis, plantar fasciitis, pseudotumour cerebri.
 Metabolic:	T2D, hyperinsulinemia, dyslipidemia, hypertension, gout, cholelithiasis, MAFLD, PCOS, hypothyroidism, obesity-related cancers.
 Monetary	Medical insurance, impact of obesity on income, education, employment, the ability to afford a healthy diet, access to weight loss programs

- FBP/TFT/B12/coeliac/folate/iron stores all normal
- **Airways** -Snores but no other OSA symps Neck Circumference **41cm** (at risk) STOP BANG 0 EPWORTH 0.
- **BMI** - **36.8** wt104kg WC 141cm wt height ratio **0.66** (highest weight currently ; BMI>30 since at least 2006)
- **CVD:** LDL 3.6 QRISK 14% BP 150/92 at clinic. No S&S HF.
- **Diabetes:** Hba1c **84** (repeat **82**) ; recurrent thrush
- **Economic:** weight not affecting work but menopause is. **She cant play sport with her children.**
- **Functional:** **LBP** – long standing – no red flags - imaging NAD advised supportive mx previously ‘loose weight’.
- **Gonadal** No Infertility/PCOS issues – 2 children
- **Health status** 65 mins /week activity + resistance training x 1
- **Image** No body dysmorphia / no binge eating disorder but **negative perception of self**
- **Gastro** **functional dyspepsia** HB N no nsaid/alcohol h pylori -ve no red flags OTC Rennies
- **Kidney** – GFR: >60 mL/min/1.73m² ACR: Normal
- **Liver** – LFTS N Fib 4 low risk 0.72



Image taken from ECPO bank

DIAGNOSIS FOUND

#CKM Diagnoses Found:

- Obesity
- T2D
- Dyslipidaemia
- Hypertension (24 hr BP cuff confirmation average day 143/92 mmHg) ECG: NSR, no LVH)
- **AND** functional dyspepsia / mechanical lower back ache / fungal infections / relationship with family impacted.

CKM care is comprehensive whole person care

SILO REVIEW CHECKERS



CKM REVIEW CHESS



CHES MOVES MADE



- Food Strategy
- Plate work
- Movement Strategy
- Stress
- Sleep - CBTi

Just start , think SMART

Algorithm 7. Behavioral/lifestyle therapy for people with obesity/adiposity-based chronic disease.

CHESS MOVES MADE



CHESSE MOVES MADE

Non-pharmacological Moves:

DSMES

Podiatry (low risk)

Regional eye screening (none present)

PARS

Rapid Pharmacological Moves:

Metformin 2g

SGLT2i

irbesartan 75 mg (improved systolic to 132)

Atorvastatin 40 mg

Repeat Hba1c at 3 months was 60; **BMI unchanged**. Incretin initiated and titrated to maximum dose

CURRENT STATE OF PLAY

Hba1c 40

U&E N ACR N

LFTS N

Fib 4 N

Systolic 128 (now off ARB)

LDL 1.9 (LLT combo ezetimibe just added in)

BMI 30.1

17 % body weight loss with incretin

11cms of waist circumference

HRT – CVD protection

Taking medications. On routine review.

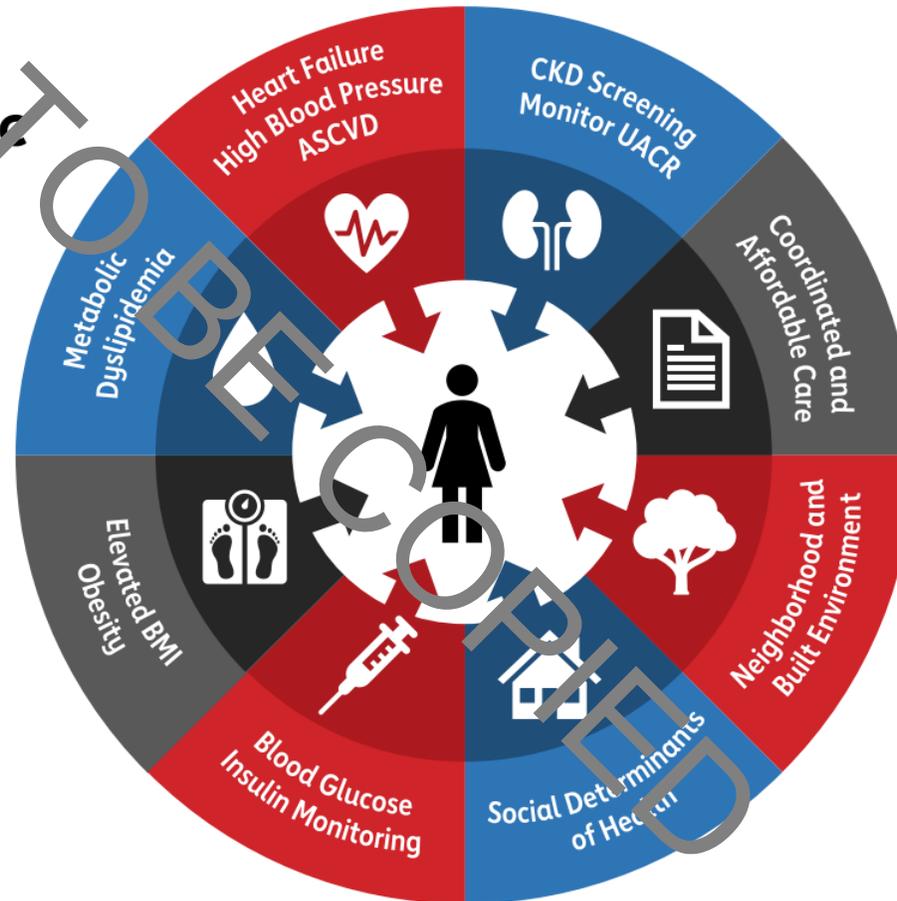


What is she doing currently ? I do not know because she is busy flourishing in her life. I will find out at routine review or if she self represents with evolving clinical need.

CVRM APPROACH ELIMINATES CARE FRAGMENTATION

Cardiovascular-Kidney-Metabolic Syndrome

**Patient-Centered
Implementation
Focus**

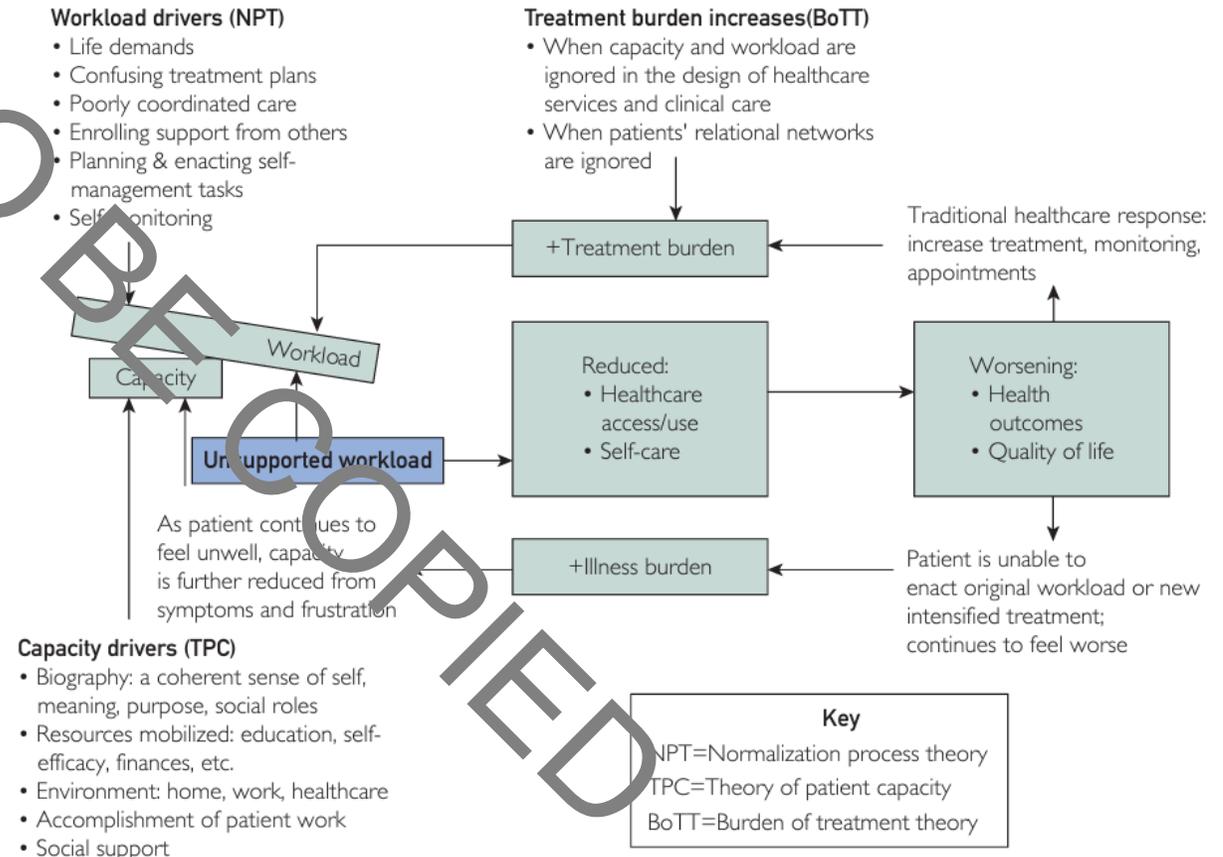


Abbreviations: ASCVD indicates atherosclerotic cardiovascular disease; BMI, body mass index; CKD, chronic kidney disease; and UACR, urine albumin-creatinine ratio.



GENERAL PRACTICE CKM REVIEW IS MINIMALLY DISRUPTIVE MEDICINE

- Multi-organ impact
 - Co-ordinated care
 -  Appointment burden
 - I review facilitating agendas across multiple specialties
 -  Person workload
- ...Polypharmacy is at play

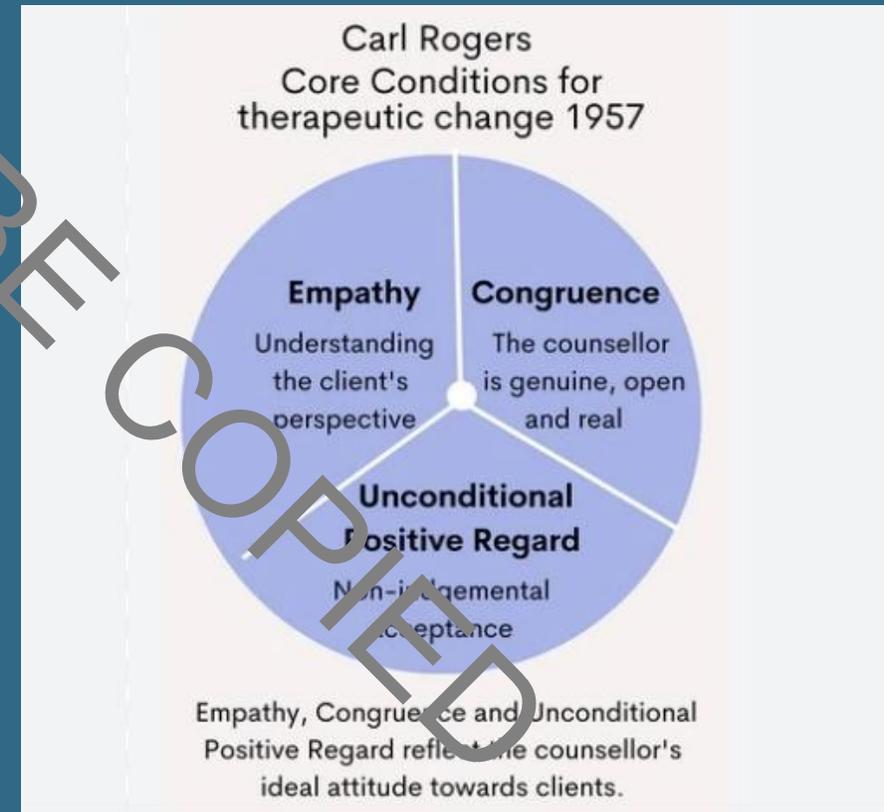


CKM PHILOSOPHY OF CARE

The professional conversation is a tool
– use it.

Behavioural Science

**Anchoring – Frame – Chunking &
Checking – Cognitive ease –
Choice architecture- Status Quo
bias**



A compendium of CKM help for your GP Houses of Care.

Type 2 Diabetes Cardiovascular Renal Metabolic Review Checklist

Medscape UK X Guidelines Primary Care Hooks

Authors: Dr Emma Daly, GP Partner, George Farm Practice, Chislehurst; Dr William Haslegrave, Consultant in Renal and General Medicine, South Tees Health NHS Foundation Trust; Dr Devendra Mehta, Registrar in Renal and General Medicine, South Tees Health NHS Foundation Trust; Dr Devendra Mehta, Registrar in Renal and General Medicine, South Tees Health NHS Foundation Trust; Dr Devendra Mehta, Registrar in Renal and General Medicine, South Tees Health NHS Foundation Trust.

Consider the following during T2D CVM shared decision making

Life Story and Lifestyle Considerations

- Consider lifestyle changes using the Behavioural Change Intervention Theory, and other social determinants of health during shared decision making
- Be aware that around 12% of people with T2D report increasing glycaemic control and an excess risk of microvascular and cardiovascular complications as well as premature death, aggressive renal modification in those individuals is imperative to mitigate this risk
- Assess and weigh up the pros and cons of lifestyle changes to help the diagnostic process of obesity due to its severity as a cardiovascular disease risk marker. Remember to adjust according to ethnicity and discuss individualised weight loss goals as appropriate. Discuss if a referral to a dietitian may be a useful referral
- Discuss if a referral to a dietitian may be a useful referral. Discuss if a referral to a dietitian may be a useful referral.

Individualised HbA_{1c} Goals

- Review the person's current HbA_{1c} and trend, and consider other factors when individualising HbA_{1c} goals, e.g., risks potentially associated with hypoglycaemia and other drug adverse effects, life expectancy, multiple long-term conditions, established vascular complications, and patient preferences, resources, and support systems
- See the expert consensus statement on diabetes and health for individualising management in older adults and adults with frailty and T2D

Primary Care Hooks

- Individualise HbA_{1c} targets in people with diabetic kidney disease. To aware that at SGLT2s have negligible glucose lowering effect once eGFR falls below 45 mL/min/1.73 m², consider adding an additional glucose-lowering medication such as a GLP-1 RA
- eGFR < 60 mL/min/1.73 m² or clinically significant proteinuria (ACR ≥ 30 mg/mol) and/or normally tolerated dose of ACEi, consider adding SGLT2 with next protective step, if appropriate of HbA_{1c}
- See the expert consensus statement on diabetes and health for individualising management in older adults and adults with frailty and T2D

Blood Pressure

There is considerable debate around optimal BP targets for people living with diabetes, with several conflicting guidelines published

- For grade 1 hypertension (people with a clinic SBP 140-159 mmHg and/or a clinic DBP 90-99 mmHg), effective lifestyle changes may delay or prevent the need for pharmacological treatment
- For hypertension on antihypertensive therapy, see the Primary Care Hook, Lifestyle Changes for Managing Hypertension
- First instance on an ACEi or ARB average target of <136/86 mmHg/150/90 mmHg (systolic target in all people)
- Second instance on an ACEi or ARB average target of <130/80 mmHg/145/90 mmHg (systolic target in all people)
- For people living with T2D, start drug treatment for ACEi or ARB, irrespective of age or ethnic background
- Measure renal function (eGFR) in people with hypertension and T2D. If there is a significant postural drop in BP (e.g., >20 mmHg SBP and/or >10 mmHg DBP) that occurs on standing, treat as a BP target based on the standing BP

Lipids

- LDL-C targets for people living with T2D:
 - moderate risk: <2.6 mmol/L
 - high risk: <2.0 mmol/L reduction from baseline and <1.8 mmol/L
 - very high risk: <1.0 mmol/L reduction from baseline and <1.4 mmol/L
- Patients with a 10% or greater increase in LDL-C after primary prevention of CVD in <40 years. Consider using OHS3C (statin) for younger individuals, particularly under the age of 40 years
- LDL-C targets may be adjusted on normally tolerated doses, consider combination lipid lowering therapy, e.g., add in ezetimibe, bempesidic acid, PCSK9 inhibitor, or inclisiran
- Consider cotherapy if the individual has established CVD (secondary prevention) and is on statins, with fasting TG 3.5 mmol/L and LDL-C between 1.04 and 2.60 mmol/L
- For secondary prevention of CVD, offer atorvastatin 80 mg

Heart

- Check pulse, if regular, consider ECG to identify AF
- Consider prevention of CVD in high risk individuals:
 - ORIS2 (NOAC) for age 65 years or age 60 years with an elevated lifetime risk of CVD in the presence of one or more of hypertension, systolic, diastolic, atrial fibrillation, or family history (in a first degree relative) of premature CVD, early onset heart failure with reduced ejection fraction, or atrial fibrillation
 - ASCOV (in individuals with AF) offer early combination therapy with warfarin and an SGLT2, irrespective of HbA_{1c}. Continued overall.

Identification of Chronic Kidney Disease in Primary Care

Medscape UK X Guidelines Primary Care Hooks

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1. Identify Individuals at Risk of CKD

Key Risk Factors:

- Age > 60 years
- Family history of kidney disease
- Structural renal tract disease
- Medication (e.g., NSAIDs, ACEi, ARBs, diuretics)
- Diabetes
- Hypertension
- Long-term treatment with NSAIDs, ACEi, ARBs, diuretics, or other nephrotoxic drugs
- Diabetes
- Hypertension
- Long-term treatment with NSAIDs, ACEi, ARBs, diuretics, or other nephrotoxic drugs

2. Test at Risk Adults for CKD

If a patient is at risk of CKD, test for eGFR and ACR.

- eGFR < 60 mL/min/1.73 m² or ACR ≥ 30 mg/mol
- Refer to renal clinic if eGFR < 45 mL/min/1.73 m² or ACR ≥ 300 mg/mol

3. Diagnose CKD

How CKD is defined:

- eGFR < 60 mL/min/1.73 m² or ACR ≥ 30 mg/mol for 3 months
- Refer to renal clinic if eGFR < 45 mL/min/1.73 m² or ACR ≥ 300 mg/mol

4. Classify and Stage CKD

CKD prognosis is depicted in the RAGWIS table (Figure 1), the recommended frequency for monitoring eGFR is represented by the numbers in boxes (e.g., 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, 41, 42, 43, 44, 45, 46, 47, 48, 49, 50, 51, 52, 53, 54, 55, 56, 57, 58, 59, 60, 61, 62, 63, 64, 65, 66, 67, 68, 69, 70, 71, 72, 73, 74, 75, 76, 77, 78, 79, 80, 81, 82, 83, 84, 85, 86, 87, 88, 89, 90, 91, 92, 93, 94, 95, 96, 97, 98, 99, 100, 101, 102, 103, 104, 105, 106, 107, 108, 109, 110, 111, 112, 113, 114, 115, 116, 117, 118, 119, 120, 121, 122, 123, 124, 125, 126, 127, 128, 129, 130, 131, 132, 133, 134, 135, 136, 137, 138, 139, 140, 141, 142, 143, 144, 145, 146, 147, 148, 149, 150, 151, 152, 153, 154, 155, 156, 157, 158, 159, 160, 161, 162, 163, 164, 165, 166, 167, 168, 169, 170, 171, 172, 173, 174, 175, 176, 177, 178, 179, 180, 181, 182, 183, 184, 185, 186, 187, 188, 189, 190, 191, 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Obesity management as a primary treatment goal for type 2 diabetes: time to reframe the conversation



Ildiko Lingvay, Priya Sumithran, Ricardo V Cohen, Carel W le Roux

Obesity is now recognised as a disease that is associated with serious morbidity and increased mortality. One of its main metabolic complications is type 2 diabetes, as the two conditions share key pathophysiological mechanisms. Weight loss is known to reverse the underlying metabolic abnormalities of type 2 diabetes and, as such, improve glucose control; loss of 15% or more of bodyweight can have a disease-modifying effect in people with type 2 diabetes, an outcome that is not attainable by any other glucose-lowering intervention. Furthermore, weight loss in this population exerts benefits that extend beyond glycaemic control to improve risk factors for cardiometabolic disease and quality of life. We review the evidence supporting the role of weight loss in the management of type 2 diabetes and propose that many patients with type 2 diabetes would benefit from having a primary weight-centric approach to diabetes treatment. We discuss the logistical challenges to implementing a new weight-centric primary treatment goal in people with type 2 diabetes.

Introduction

Over the past decade, management of patients with type 2 diabetes has undergone a major conceptual change, with treatment objectives shifting to include a cardiocentric goal in the subpopulation with high cardiovascular risk, alongside the singular glucocentric goal that has long been held.¹ This advance was driven by studies showing that several glucose-lowering agents, used in addition to standard of care, further lower the risk of myocardial infarction, stroke, and cardiovascular death,^{2,3} largely independently of lowering blood glucose concentration.^{4,5} Yet, even after this landmark evolution, the treatment framework for type 2 diabetes is primarily focused on preventing or treating the downstream metabolic consequences, which tend to occur late in the disease course.

A promising opportunity lies in intervening upstream to address the key pathophysiological driver of type 2 diabetes and its associated metabolic complications: obesity (figure 1). Sustained loss of at least 15% bodyweight has a major effect on progression of type 2 diabetes, inducing remission in a large proportion of patients and markedly improving metabolic status in many others.^{6,7}

Until 2021, the only intervention that could routinely result in maintenance of weight loss of this magnitude was bariatric surgery. However, despite its considerable benefits, a complex surgical procedure is not feasible or scalable as the mainstay for a population-wide intervention. Now, with effective pharmaceuticals to reduce bodyweight in the pipeline, many of which can also directly lower blood glucose concentration, it is time to rethink treatment goals for patients with type 2 diabetes to position obesity management as a principal goal (ie, aiming for substantial weight loss as the primary means to treat patients with type 2 diabetes and reach glycaemic targets). Such a weight-centric intervention would disrupt the underlying pathophysiology of type 2 diabetes, reverse or slow down the disease course, concomitantly benefit other associated cardiovascular

risk factors, and prevent microvascular and macrovascular complications of type 2 diabetes in the long term.

Here, we review the clinical evidence supporting weight loss as a fundamental target, propose a novel therapeutic framework, and explore challenges for the widespread implementation of this approach for people with type 2 diabetes.

Type 2 diabetes and obesity are interconnected, heterogeneous diseases

Obesity and type 2 diabetes are heterogeneous conditions. Not all people who are categorised as having obesity (ie, body-mass index ≥ 30 kg/m²) have excessive adiposity. Moreover, even among people who do have excess adiposity, not all people will have metabolic complications, such as type 2 diabetes.⁸ Conversely, some people with only minimal adiposity develop metabolic complications, prompting the concept that adipose tissue pathology, rather than quantity, might be the primary driver of complications.⁹ Abnormal adipose tissue pathology is characterised by adipocyte hypertrophy, visceral adiposity, and ectopic fat deposition, with resulting systemic inflammation and metabolic dysfunction. This process is not directly proportional to adipose quantity or body-mass

Search strategy and selection criteria

References for this Review were identified by searching MEDLINE, PubMed, and the ClinicalTrials.gov registration site using the search terms: "overweight", "obesity", "weight gain", "weight loss", "weight management", "body weight", "morbid obesity", "obesity pharmacotherapy", "adiposity", and "bariatric surgery" in combination with the term "type 2 diabetes". We included references from relevant articles that were identified during the search. Articles published in English between Jan 1, 1990, and March 30, 2021, were reviewed and included on the basis of originality, relevance to the broad scope of the Review, and impact in the field (ie, number of citations).

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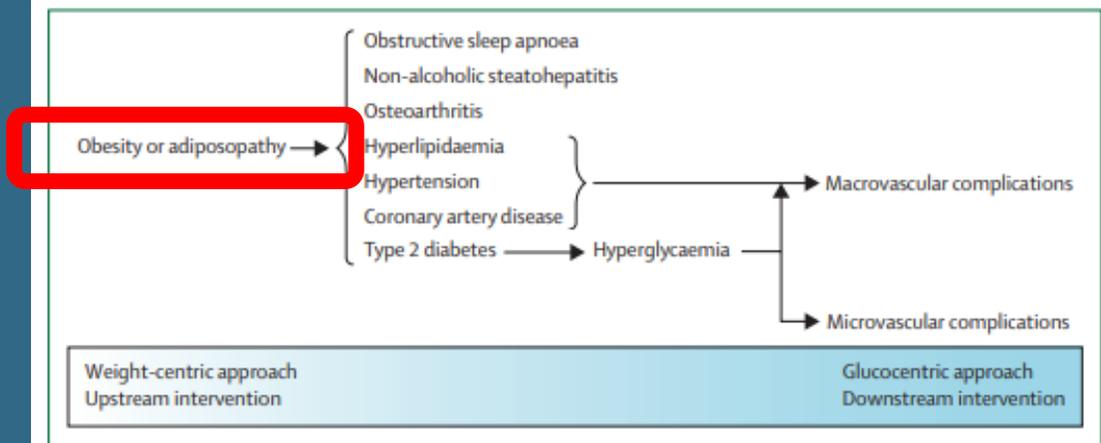


Figure 1: Illustration of the wide-ranging benefits of an upstream weight-centric approach versus a glucocentric management approach

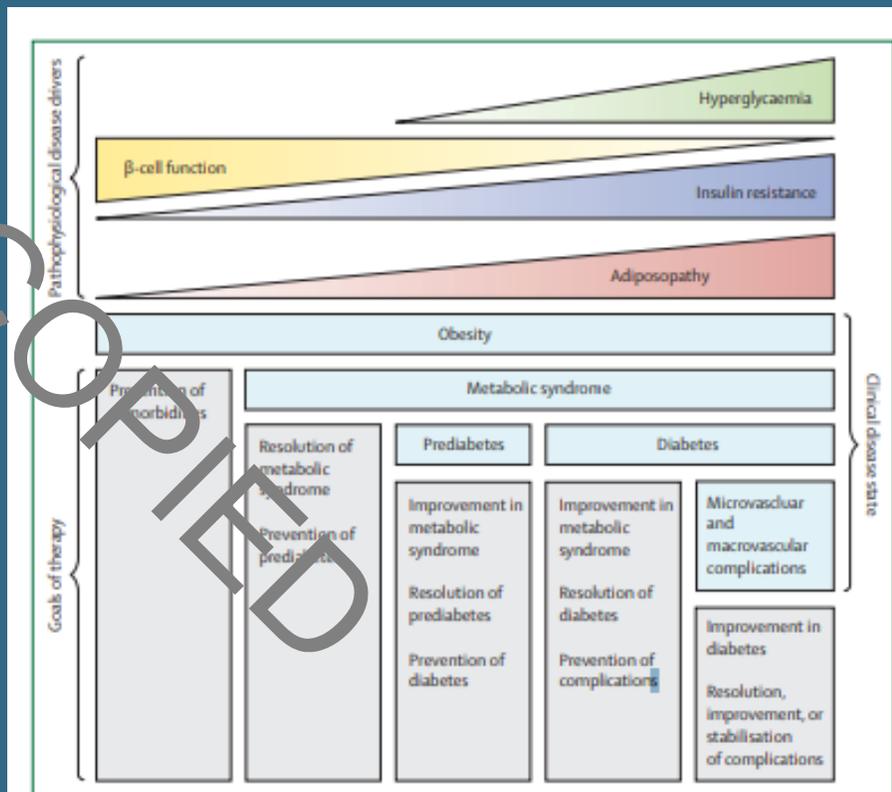


Figure 2: The disease continuum for weight-related type 2 diabetes. Pathophysiology, clinical disease states, and goals of therapy are shown along the continuum.

Striving for early effective glycaemic and weight management in type 2 diabetes: A narrative review

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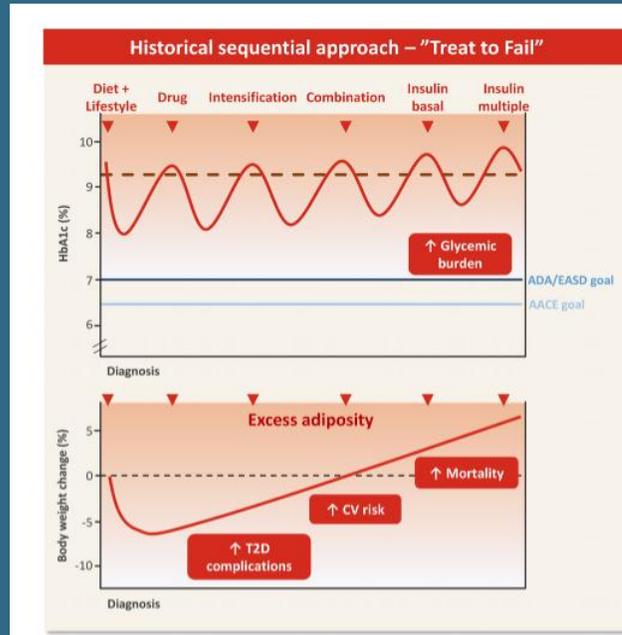
Funding information
Eli Lilly and Company

Abstract

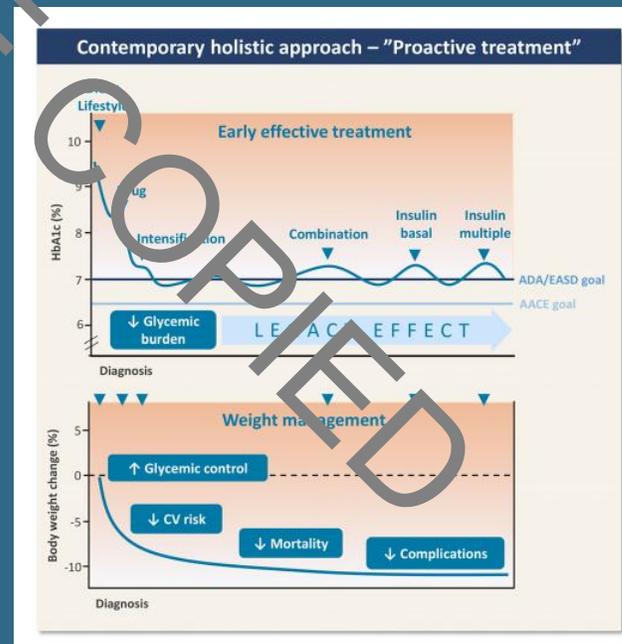
Despite the recognition by key guidelines that achieving early glycaemic control has important benefits in individuals with type 2 diabetes (T2D) and that addressing excess adiposity is one of the central components of comprehensive person-centred T2D care, a substantial proportion of individuals with T2D do not meet their metabolic treatment goals. Prior treatment paradigms were limited by important treatment-associated risks such as hypoglycaemia and body weight gain. Therefore, a more conservative, sequential approach to treatment was typically utilized. One potential consequence of this approach has been a missed opportunity to achieve a 'legacy effect', where early treatment to reach glycaemic targets is associated with enduring long-term benefits in T2D. Additionally, while previous treatment approaches have addressed core defects in T2D, including insulin resistance and β -cell function decline, they have been unable to address one of the underlying causal abnormalities—excess adiposity. Here, we review currently available evidence for the beneficial long-term effects of early glycaemic control and management of body weight in people with T2D and discuss potential next steps.

KEYWORDS

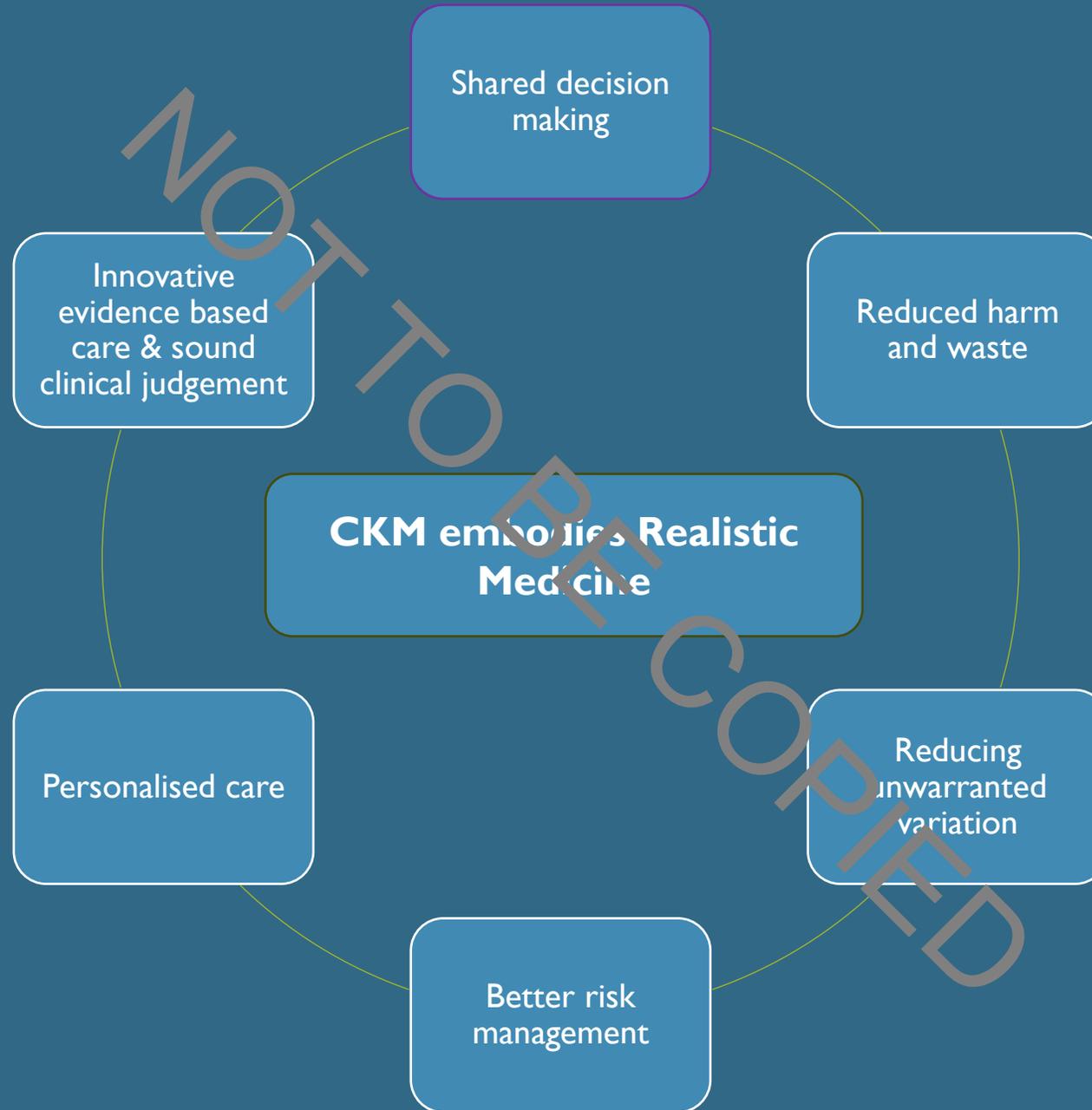
antidiabetic drug, antiobesity drug, diabetes complications, glycaemic control, weight management



Is this really working for us in General Practice?



Advocating for this approach moving forward makes sense.



NHS General Practice CKM care

Cardiology

Nephrology

Hepatology

Person with
CVRM
syndrome
and General
Practice
team

Diabetology &
Obesity physicians

Respiratory

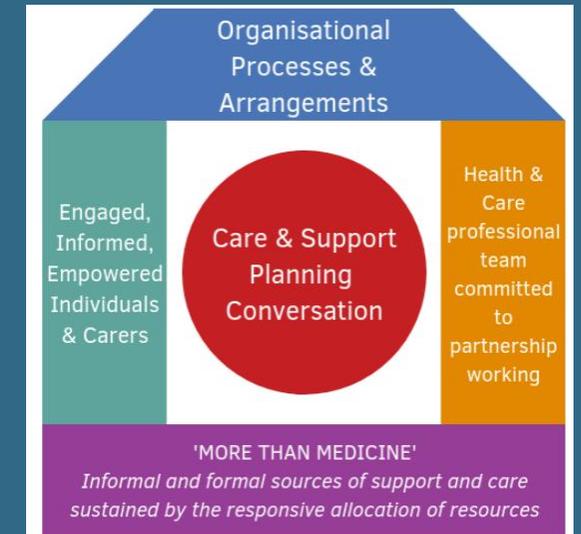
Psychology &
Dieticians

Primary Care Teams –
Community Pharmacy /
Physio/OT/ Social Services
and Voluntary Sector

- **Whole person advanced generalist care :**
 - Co-ordinated
 - Comprehensive
 - Continuous
 - Person centred
- Longitudinal life span care
- MDT care – stronger together
- Agile care – open doors of engagement with and back up from other teams when required.

Take homes from a CKM GP

- CKM syndrome - a dynamic continuum that changes over time.
- Contextualise. Prioritise. Personalise.
- Shared decision making - **with** the person , **not to** the person.
- **Ordering** and **pacing** of interventions – a personalised marathon.
- **Life long treatments and interventions** ; **not** burst therapies or approaches.
- **Combination therapy and interventions** = organ and person protection.
- General Practice MLTC infrastructure **will evolve** from current set up to a **de-siloed** format.
- This will take a **team** – if you are doing this as a lone professional – **stop doing it alone.**



We are stronger together

CODE CODE CODE CODE

NOT
BE
COPIED

General Practice Cardiovascular Kidney Metabolic Review

**VALUE LIES IN A COMPREHENSIVE AND
COMPREHENSIBLE APPROACH TO A
COMPLEX PROBLEM**

The Lightclub



Lets get back to the basics, and get them right.



Journal of
Clinical Medicine



Review

Cardiovascular–Kidney–Metabolic Syndrome: A New Paradigm in Clinical Medicine or Going Back to Basics?

Victoria Mutruc ^{1,2,*}, Cristina Bologa ^{1,2}, Victorița Șorodoc ^{1,2}, Alexandr Ceasovschih ^{1,2},
Bianca Codrina Morărașu ^{1,2}, Laurențiu Șorodoc ^{1,2}, Oana Elena Catar ³ and Cătălina Lionte ^{1,2,*}

Mutruc, V.; Bologa, C.; Șorodoc, V.; Ceasovschih, A.; Morărașu, B.C.; Șorodoc, L.; Catar, O.E.; Lionte, C. Cardiovascular–Kidney–Metabolic Syndrome: A New Paradigm in Clinical Medicine or Going Back to Basics? *J. Clin. Med.* 2025, 14, 2833. <https://doi.org/10.3390/jcm14082833>



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General Practice #CKM care is the enactment of humanity.

Collectivism
Compassion
Care

The responsibility to curate it together lies with us.