

# Older People with Diabetes – Key Aspects that underpin better care and management

Professor Alan Sinclair

Foundation for Diabetes Research in Older People and King's  
College, London



# Foundation for Diabetes Research in Older People (fDROp) *Our mission and vision 2021-24*

Director: Professor Alan Sinclair



National Advisory Panel on Care Home Diabetes



WHO



## Our mission

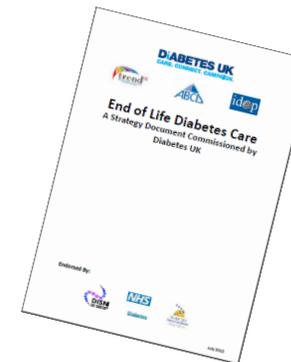
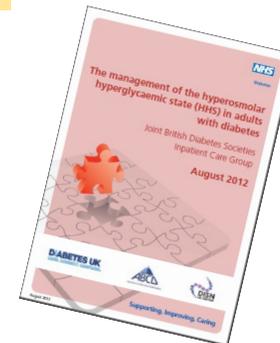
- As a **not for profit research organisation**, to enhance the quality of diabetes care for older people through new initiatives in clinical practice, audit and research
- To provide a **forum for discussion** between health professionals and scientists, and involve people with diabetes, their carers and families, in programmes which promote their health and well-being
- To examine the relationship between diabetes and related metabolic disorders to the development of **frailty and sarcopaenia**

## Our vision

- Establish sustainable academic partnerships
- Ensure policies and strategies are developed to meet the needs of older people with diabetes and related metabolic disease



Innovative Medicines Initiative



# Scope of my talk

- To examine key aspects of diabetes and ageing
- To examine what constitutes the condition of type 2 diabetes in older adults
- To look at those factors that can influence the successful management of type 2 diabetes in older adults



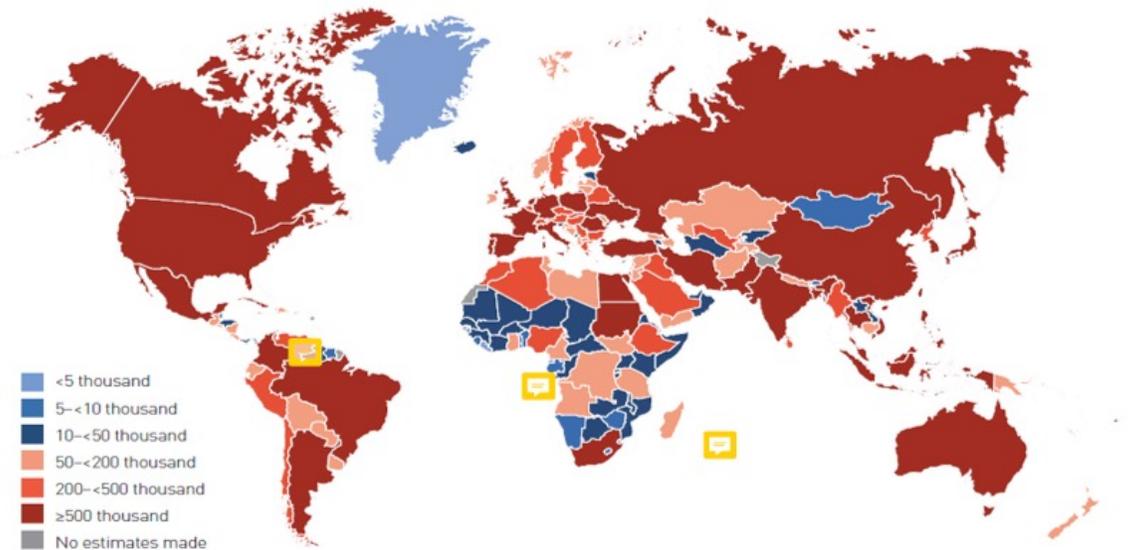
# Global diabetes rates in older people – findings from the IDF Atlas 9<sup>th</sup> edition. *Sinclair AJ et al. Diab Res Clin Pract 2020.*

**Table 1: Global diabetes estimates in people older than 65 years in 2019, 2030 and 2045**

|   | 2017                        | 2019                            | 2030                        | 2045                        |
|---|-----------------------------|---------------------------------|-----------------------------|-----------------------------|
| <b>Adult population (65–99 years)</b>                                   | 652.1 million               | 704.4 million                   | 995.2 million               | 1.4 billion                 |
| <b>Prevalence (65–99 years)</b>   | 9.6% (15.4–23.4%)           | 19.3% (15.3–24.2%) <sup>i</sup> | 19.6% (15.5–24.8%)          | 19.6% (15.2–25.4%)          |
| <b>Number of people older than 65 years with diabetes (65–99 years)</b> | 122.8 million (100.2–152.3) | 135.6 million (107.6–170.6)     | 195.2 million (154.7–247.1) | 276.2 million (214.8–358.9) |

<sup>i</sup>95% confidence intervals are reported in brackets.

**Map 3.3 Number of people older than 65 years with diabetes by country in 2019**



# Diabetes in England



 **3.8m**

Number of people in  
England with diabetes

**£8.8bn**

Current annual cost of  
Type 2 diabetes to the NHS

**940k** 

Number of people with  
undiagnosed diabetes

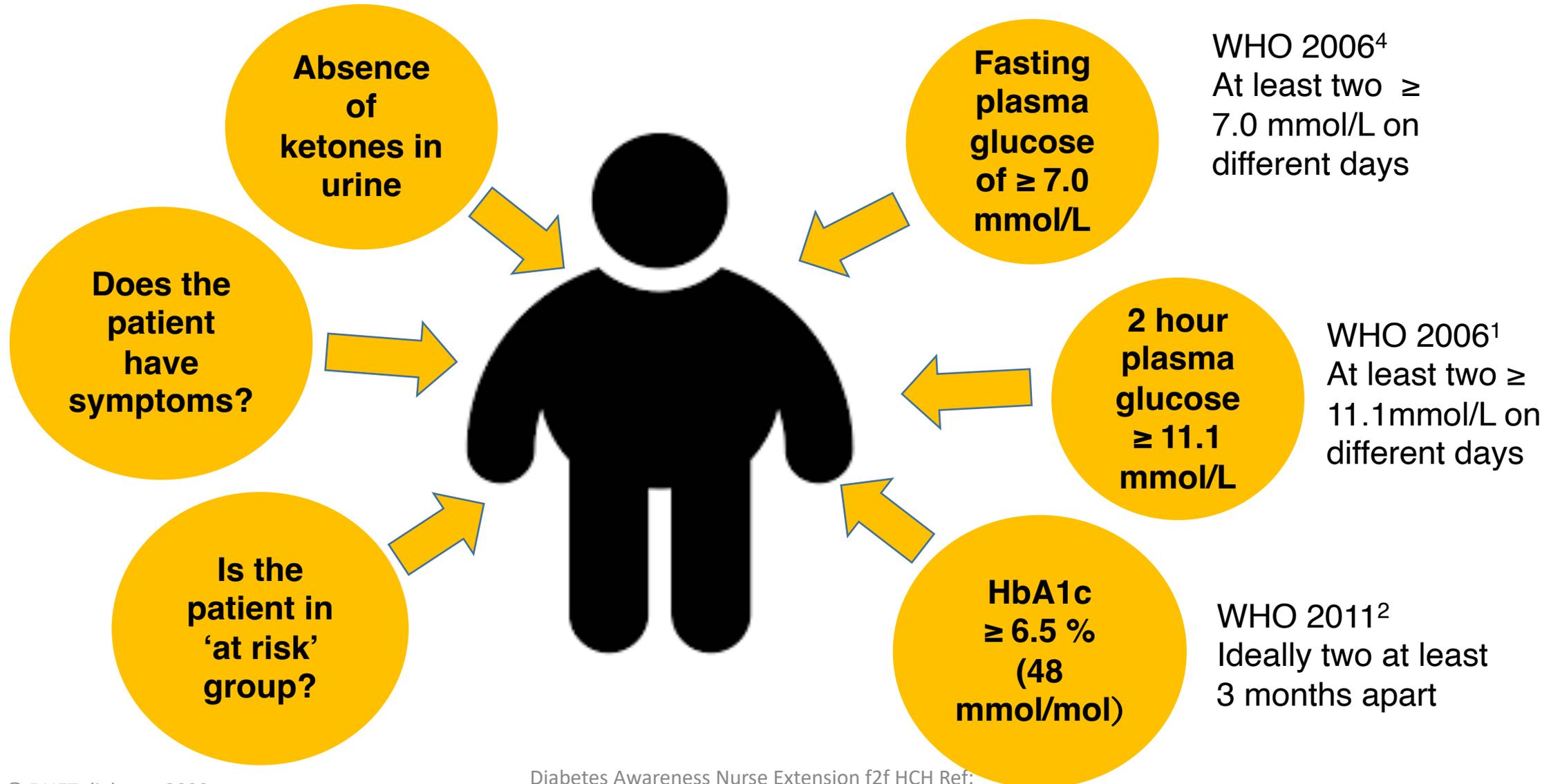
**90%** 

% of diabetes cases  
which are Type 2,  
which is preventable

**4.9m** 

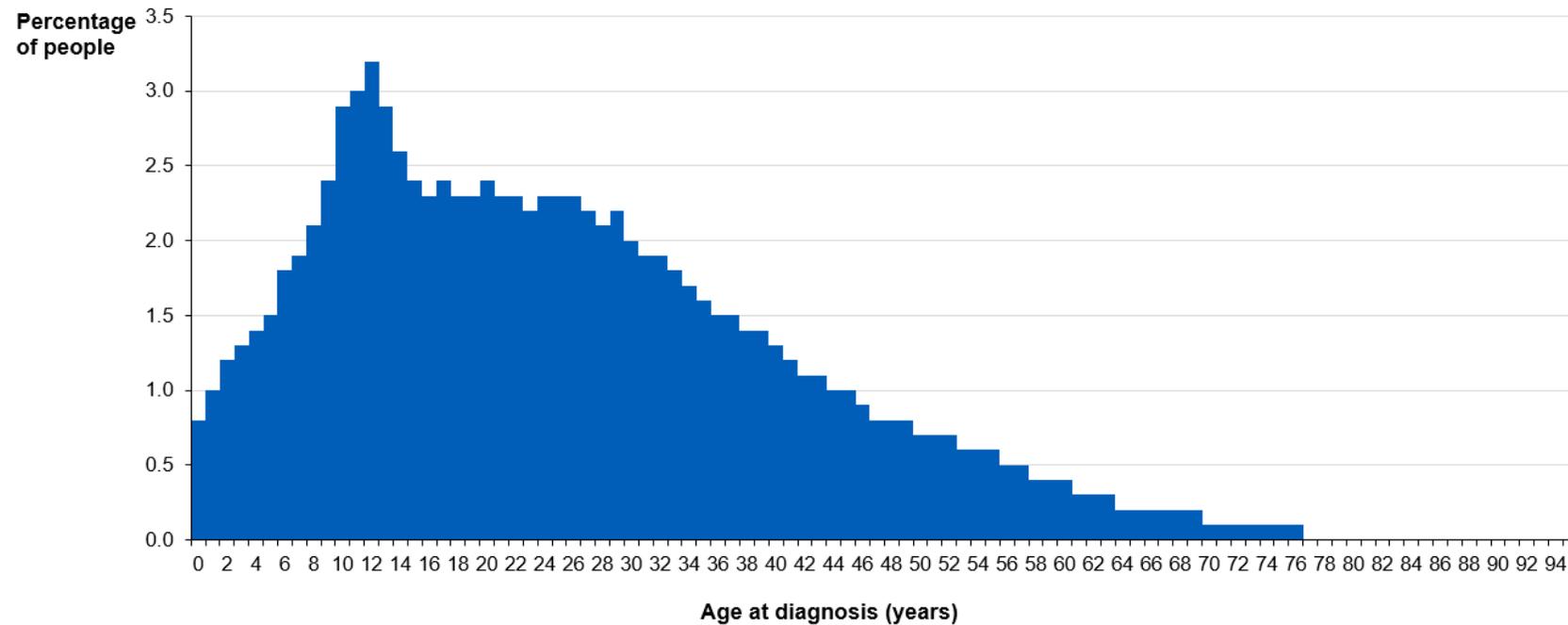
Estimated diabetes  
cases by 2035

# Making a diagnosis of Type 2 Diabetes<sup>1,2,18</sup>



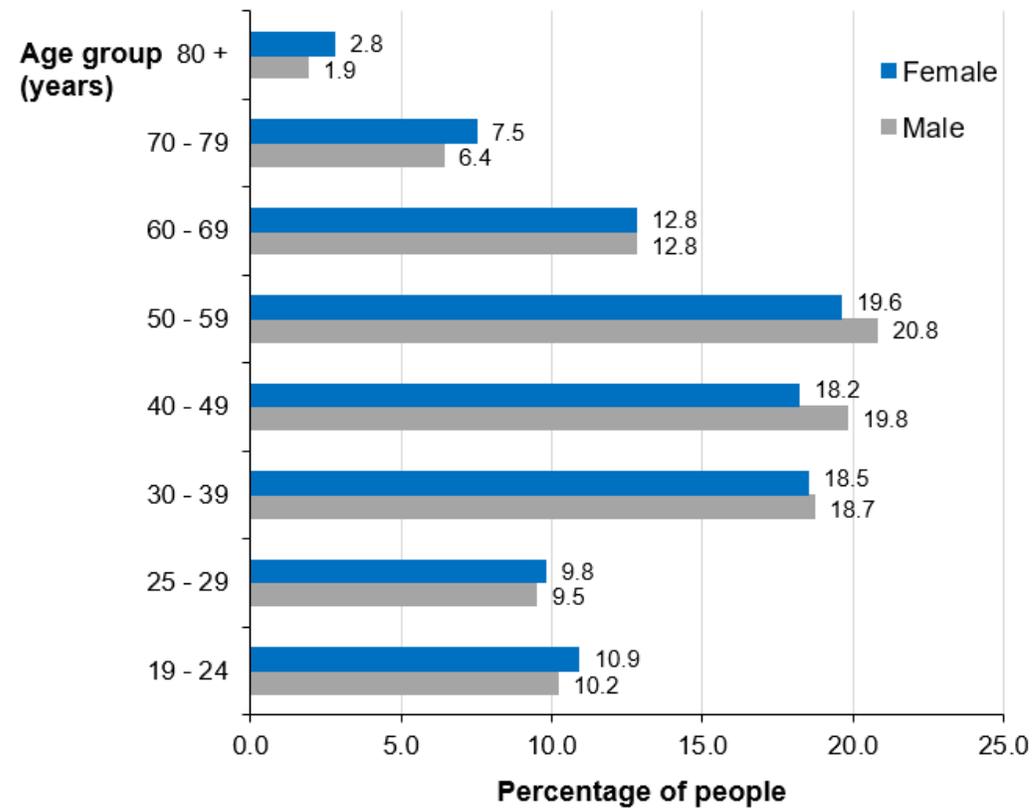
# Type 1 diabetes in England & Wales – 2019-20 NDA Supporting Information

Figure 2: People with type 1 diabetes, by age of diagnosis\*, England and Wales, 2019-20



# Type 1 Diabetes – NDA 2019-20

Figure 1: People with type 1 diabetes, by age group and sex, England and Wales, 2019-20

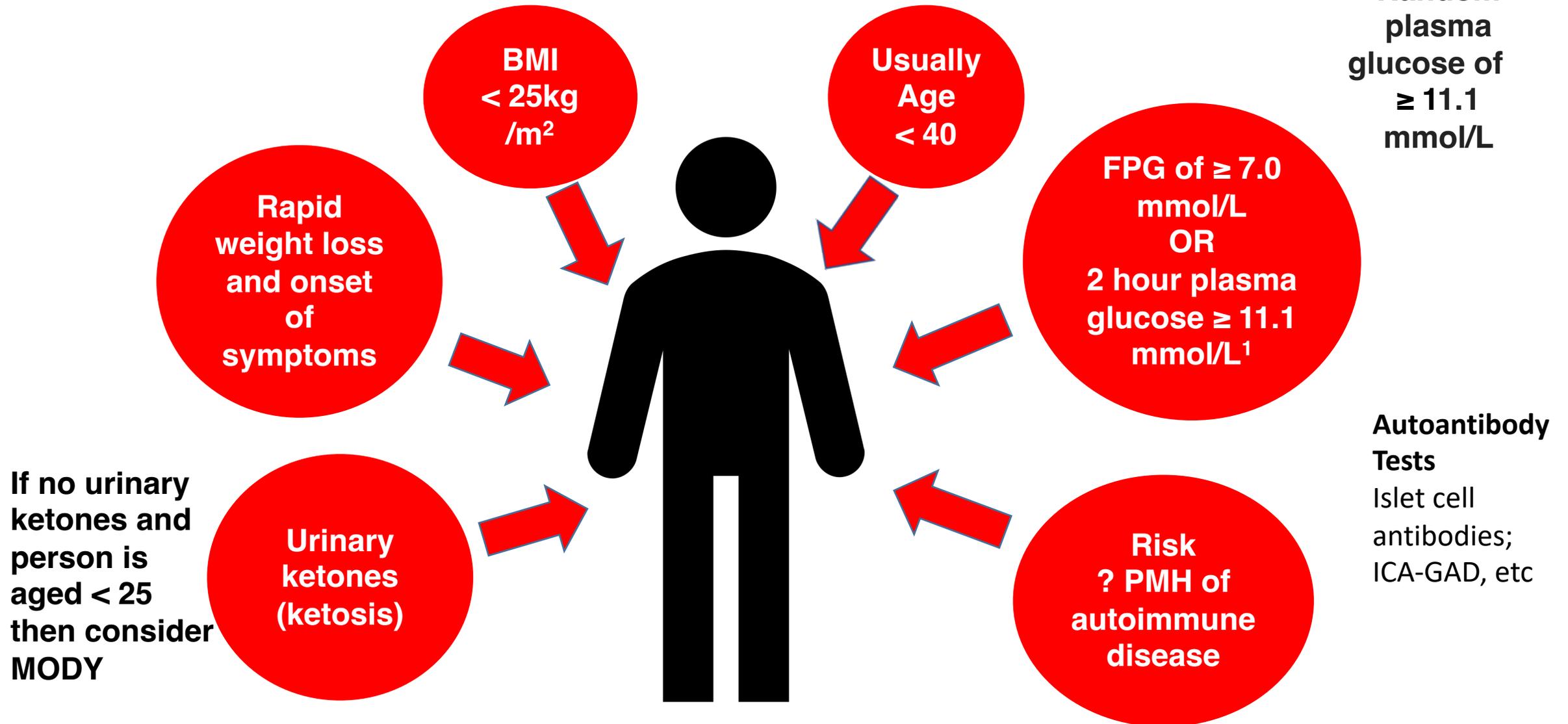


**218, 670 confirmed and unconfirmed cases of type 1 diabetes: 56.9% Male**

**86.2% white  
3.5% Asian  
2.3% Black  
2.1% Mixed/other  
9.5% not known**

*Courtesy of Prof Robert Young; source NHS Digital*

# Making a diagnosis of Type 1 Diabetes<sup>1,2,17</sup>



# Prevalence of Diabetes Mellitus in Care Homes: *the Birmingham and Newcastle Screening Studies*

Sinclair AJ, Gadsby R, Croxson SCM et al, *Diabetes Care* 2001; Aspray et al. *Diabetes Care* 2006, 29 (3):707-8

Little evidence of structured diabetes care

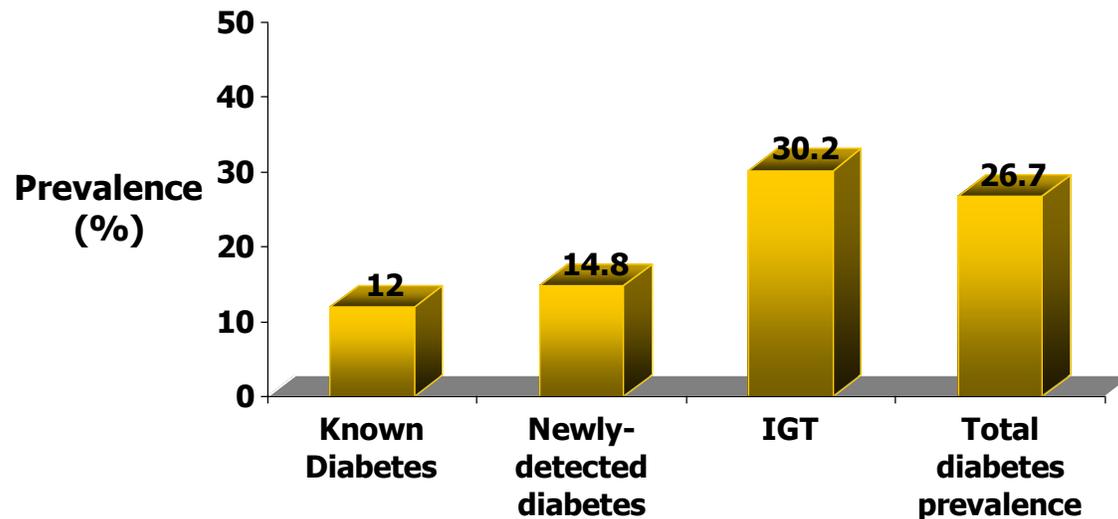
No specialist follow-up

(Reviewed by Sinclair AJ, Aspray TJ, 2009, *Diabetes in Old Age – 3<sup>rd</sup> edition*)

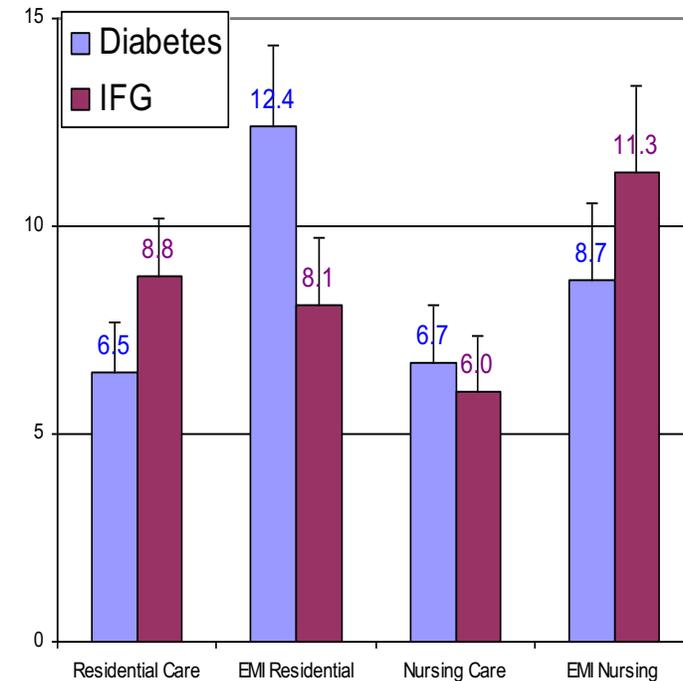
Diabetes is an independent risk factor for admission into a care home

High hospital admission rate with associated high mortality

The Birmingham Study 2001



The Newcastle Study 2006



# Covid-19 and Diabetes in Older People

EClinicalMedicine 22 (2020) 100343



Commentary

## Age, frailty and diabetes – triple jeopardy for vulnerability to COVID-19 infection

A.J. Sinclair<sup>a,\*</sup>, A.H. Abdelhafiz<sup>b</sup>

<sup>a</sup> Foundation for Diabetes Research in Older People, Diabetes Frail Ltd, Droitwich Spa, WR9 0QH, UK and Kings College, London, SE1 9NH, United Kingdom

<sup>b</sup> Department of Geriatric Medicine, Rotherham General Hospital, Moorgate Road, Rotherham S60 2UD, United Kingdom

- Old age, frailty and diabetes are all inter-related and all are risk factors for mortality in covid-19 (a coronavirus disease)
- Frailty worsens prognosis in any severe illness
- Ageing immune system is associated with a low grade and chronic inflammatory state (InflammAgeing) marked by raised inflammatory markers such as IL-6 and C-reactive protein – this creates an exaggerated susceptibility to infection
- Patients infected with covid admitted to ITUs are more likely to have diabetes
- Diabetes is associated with immune dysfunction (impaired macrophage and lymphocyte function) and speeds progression to organ failure and septic shock in severe infections

# UK and European Responses to Covid-19 in Care Homes

*Diabet Med.* 2020 May 5 : 10.1111/dme.14317.  
doi: [10.1111/dme.14317](https://doi.org/10.1111/dme.14317) [Epub ahead of print]

PMCID: PMC7267536  
PMID: [32369634](https://pubmed.ncbi.nlm.nih.gov/32369634/)

## Guidelines for the management of diabetes in care homes during the Covid-19 pandemic

Alan Sinclair,<sup>1</sup> Ketan Dhataria,<sup>2</sup> Olivia Burr,<sup>3</sup> Dinesh Nagi,<sup>4</sup> Kath Higgins,<sup>5</sup> David Hopkins,<sup>3</sup> Mayank Patel,<sup>6</sup> Partha Kar,<sup>7</sup> Catherine Gooday,<sup>2</sup> Dan Howarth,<sup>3</sup> Ahmed Abdelhafiz,<sup>8</sup> Philip Newland-Jones,<sup>6</sup> and Simon O'Neill<sup>3</sup>

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This article has been [cited by](#) other articles in PMC.

### Abstract

The National Diabetes Stakeholders Covid-19 Response Group was formed in early April 2020 as a rapid action by the Joint British Diabetes Societies for Inpatient Care, Diabetes UK, the Association of British Clinical Diabetologists, and Diabetes Frail to address and support the special needs of residents with diabetes in UK care homes during Covid-19. It was becoming obvious that the care home sector was becoming a second wave of Covid-19 infection and that those with diabetes residing in care homes were at increased risk not only of susceptibility to infection but also to poorer outcomes. Its key purposes included minimising the morbidity and mortality associated with Covid-19 and assisting care staff to identify those residents with diabetes at highest risk of Covid-19 infection. The guidance was particularly created for care home managers, other care home staff, and specialist and non-specialist community nursing teams. The guidance covers the management of hyperglycaemia by discussion of various clinical scenarios that could arise, the management of hypoglycaemia, foot care and end of life care. In addition, it outlines the conditions where hospital admission is required. The guidance should be regarded as interim and will be updated as further medical and scientific evidence becomes available.

**Keywords:** Diabetes, care homes, Covid-19, residents, frailty, insulin



*Aging Clin Exp Res.* 2021; 33(4): 895–900.

Published online 2021 Apr 15. doi: [10.1007/s40520-021-01822-1](https://doi.org/10.1007/s40520-021-01822-1)

PMCID: PMC8046642

PMID: [33856663](https://pubmed.ncbi.nlm.nih.gov/33856663/)

## Dexamethasone and oxygen therapy in care home residents with diabetes: a management guide and algorithm for treatment: a rapid response action statement from the European Diabetes Working Party for Older People (EDWPOP) and European Geriatric Medicine Society (EuGMS)

Alan James Sinclair,<sup>1,2</sup> Stefania Maggi,<sup>3,4</sup> Ahmed Hassan Abdelhafiz,<sup>2,5</sup> Nicola Veronese,<sup>4,6</sup> Leocadio Rodriguez-Manas,<sup>2,7</sup> and Isabelle Bourdel-Marchasson<sup>4,8</sup>

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### Abstract

Go to:

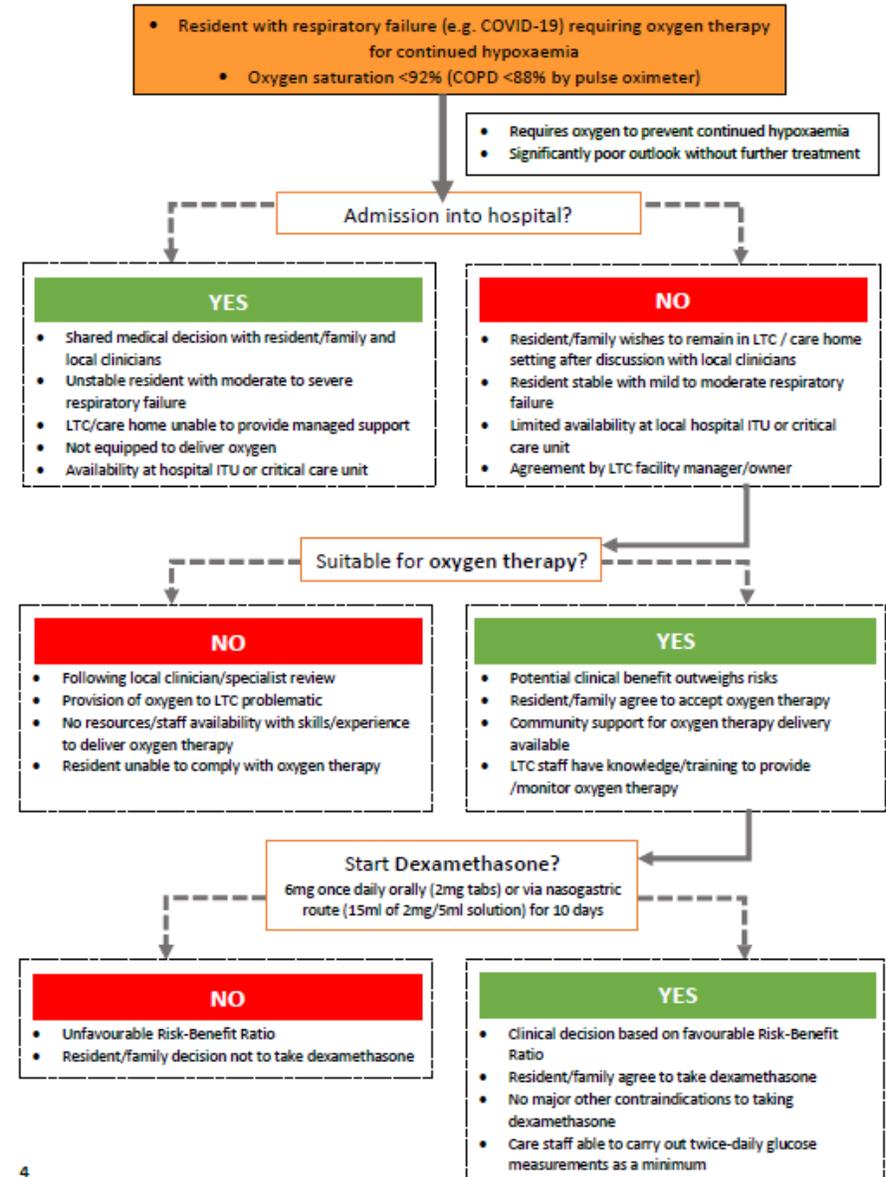
This statement addresses the need to provide clinically relevant and practical guidance for long-term care staff working in care homes and other stakeholders engaged in the care of residents who require consideration for dexamethasone and oxygen therapy. It had been provided following a series of consensus discussions between the EDWPOP and the EuGMS in January and February 2021. Its main aim is to minimise morbidity and mortality from serious acute illnesses including COVID-19 requiring these treatments within the long-term care sector.

**Keywords:** Long-term care, Diabetes, Oxygen, Dexamethasone, COVID-19

# Dexamethasone- Oxygen delivery algorithm<sup>30</sup>



Figure 1: Algorithm for Oxygen and Dexamethasone Therapy in Long-term Care (LTC)



# Stakeholder and Advisory Representatives



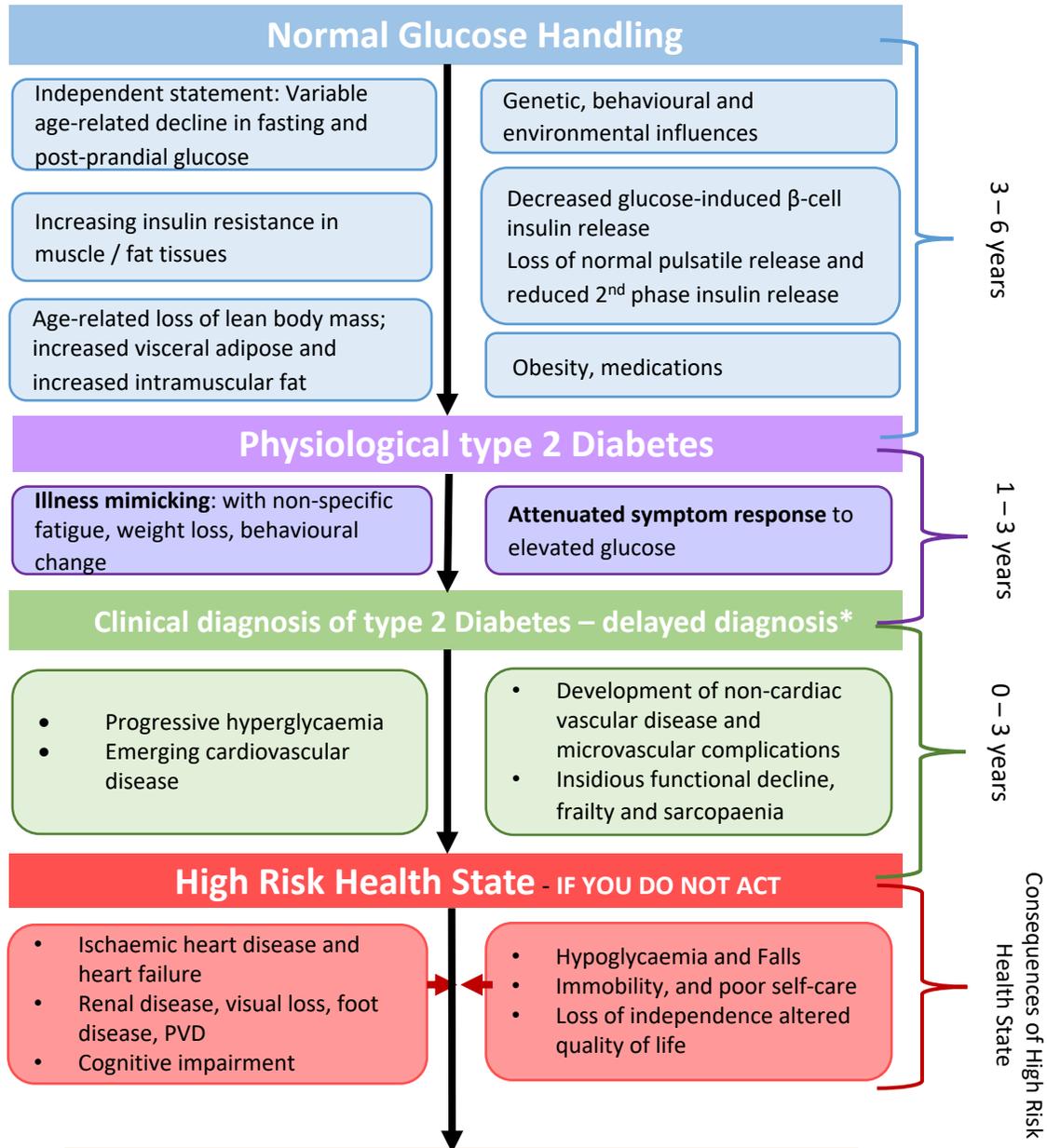
National Advisory Panel on Care Home Diabetes

## Representative Bodies of the NAPCHD

|   |  |  |
|---|--|--|
| <br>DIABETES UK<br>KNOW DIABETES. FIGHT DIABETES.      | <br>KING'S<br>College<br>LONDON                             | <br>The Association for<br>Clinical Biochemistry &<br>Laboratory Medicine<br>Better Science. Better Testing. Better Care. |
| <br>t.rend™<br>DIABETES                                | <br>Directors of<br>adass<br>adult social services          | <br>JBDS-IP<br>Joint British<br>Diabetes Societies<br>for Inpatient care  |
| <br>O.P.D.N.<br>Older People's Diabetes Network        | <br>Aston University<br>BERMINHAM UK                        | <br>CARE ENGLAND<br>Representing independent care providers   |
| <br>BDA<br>The Association<br>of UK Dietitians         | <br>hallmark<br>care homes                                  | <br>DISN<br>UK GROUP<br>WORKING WITH PRACTICE SINCE 2004  |
| <br>RCGP<br>Royal College of<br>General Practitioners | <br>ABCD<br>Association of British Clinical Diabetologists | <br>UKCPA<br>CLINICAL PHARMACY ASSOCIATION   |
| <br>DUETdiabetes                                     | <br>Care Quality<br>Commission                            | <br>fDROPF<br>foundation for Diabetes<br>Research in Older People   |

Available at: <http://fdrop.net/napchd>

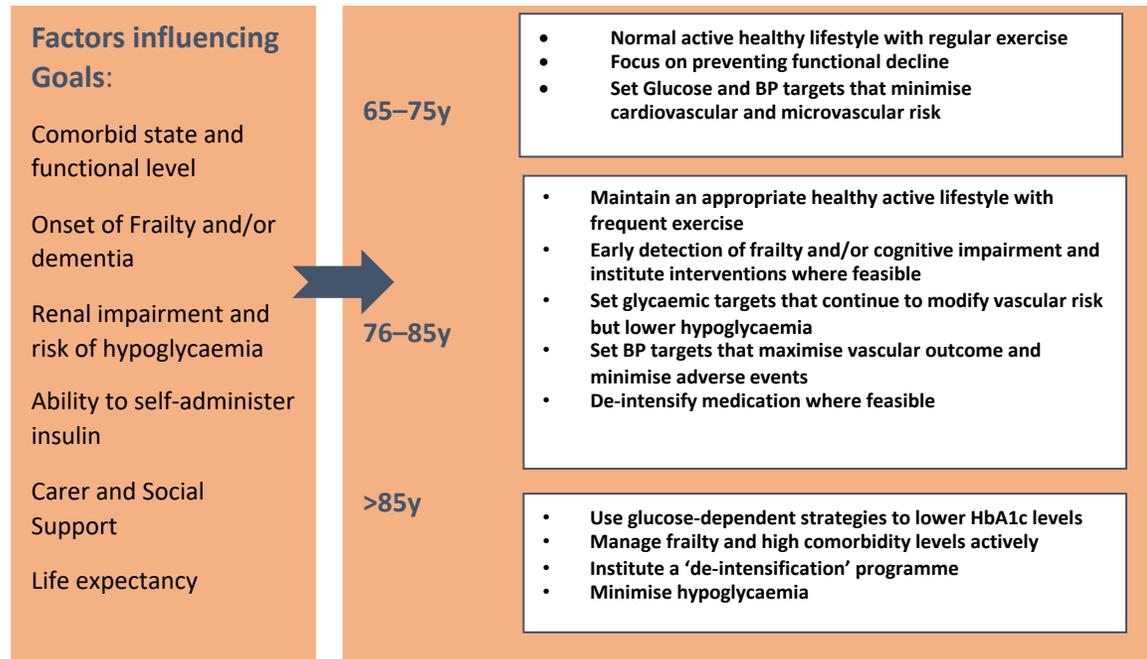
How should we describe type 2 diabetes in older adults ?



## Dynamic Modulation of Goals by decade

# Diabetes in Older Adults – a Complex Illness Model

## Template for Dynamic Modulation of Goals by decade



Sinclair AJ, Abdelhafiz A, Forbes A, Munshi M, Diabetic Med 2018

# An age-related disease in older people where many clinical failures can be prevented?

Diagnosis of Diabetes

- **Delayed diagnosis**

Onset of complications

- **Late detection: particularly eye disease/peripheral nerve damage due to lack of screening/foot disease**

Hypoglycaemia

- **Lack of recognition; use of inappropriate treatment or overtreatment**

Frailty and dementia

- **Lack of recognition plus/minus failure to examine and diagnose**

Care home residents with diabetes

- **Lack of individualized care plans: hypoglycaemia, recurrent infections, inadequate nutrition, pressure sores, lack of monitoring; lack of training and education of care**

# Mortality (*hazard ratio-adjusted*) in Older People with Diabetes (DM) with non-DM as reference

*Forbes A, Murrells T, Sinclair AJ, 2016*

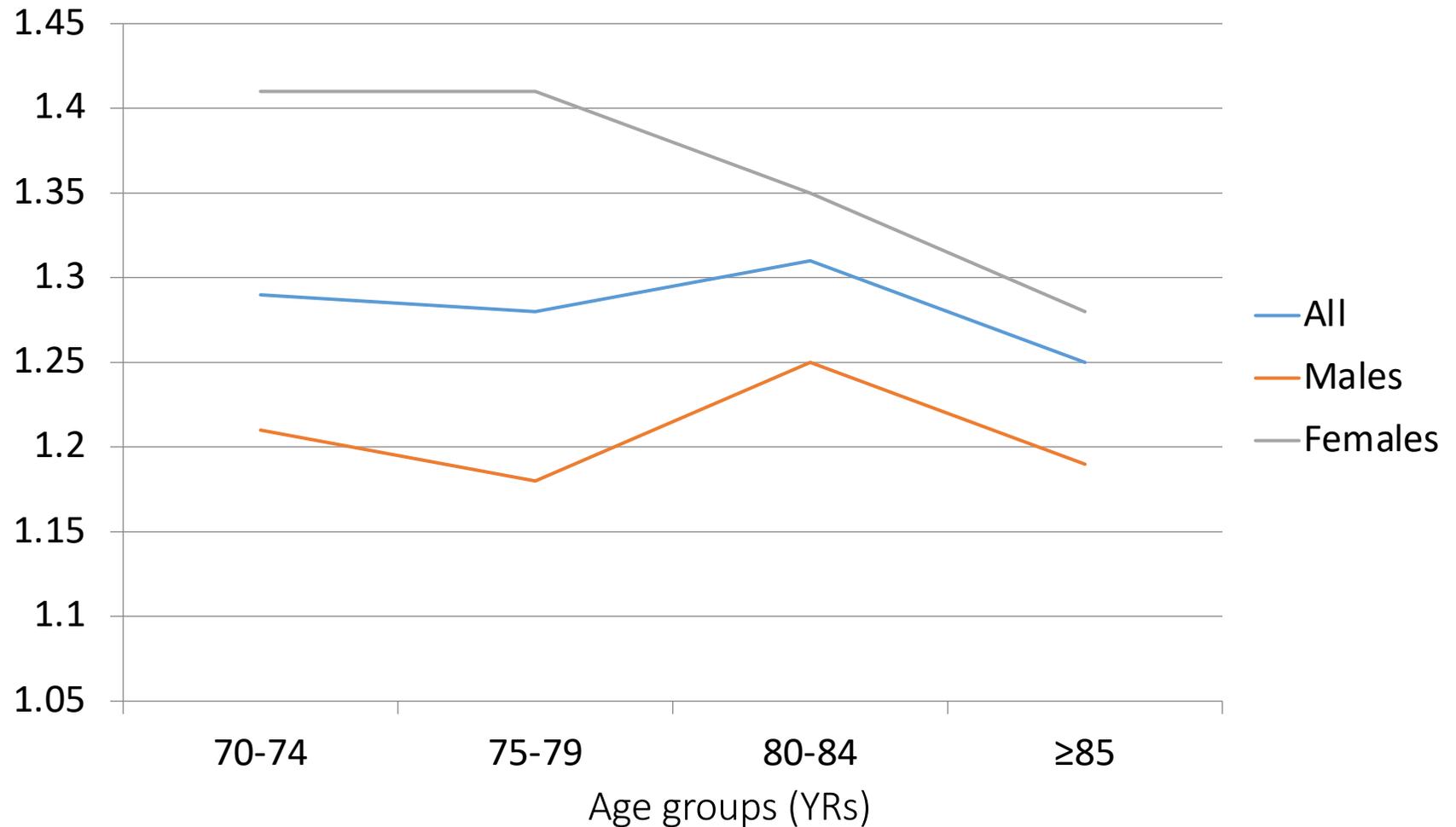
Overall excess mortality risk:

All = 29%

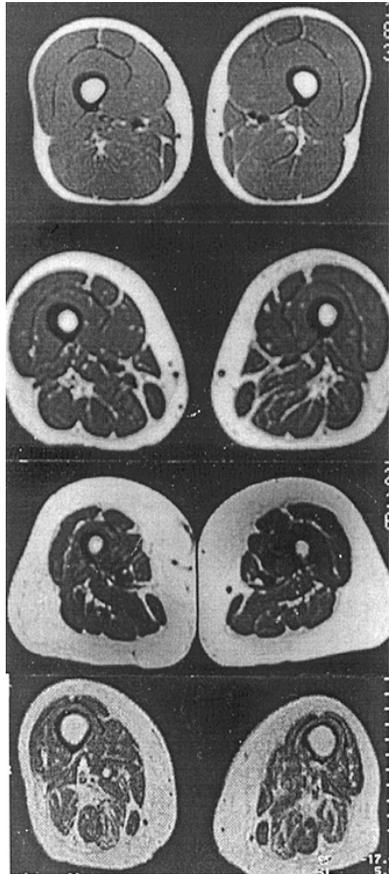
Males = 21%

Females = 36%

The Health Improvement Network (THIN) database: 464 UK general practices: electronic medical records: > 3 million records



# Age-related loss of muscle mass - sarcopaenia



31 yrs (M)

66 yrs (M)

73 yrs (F)

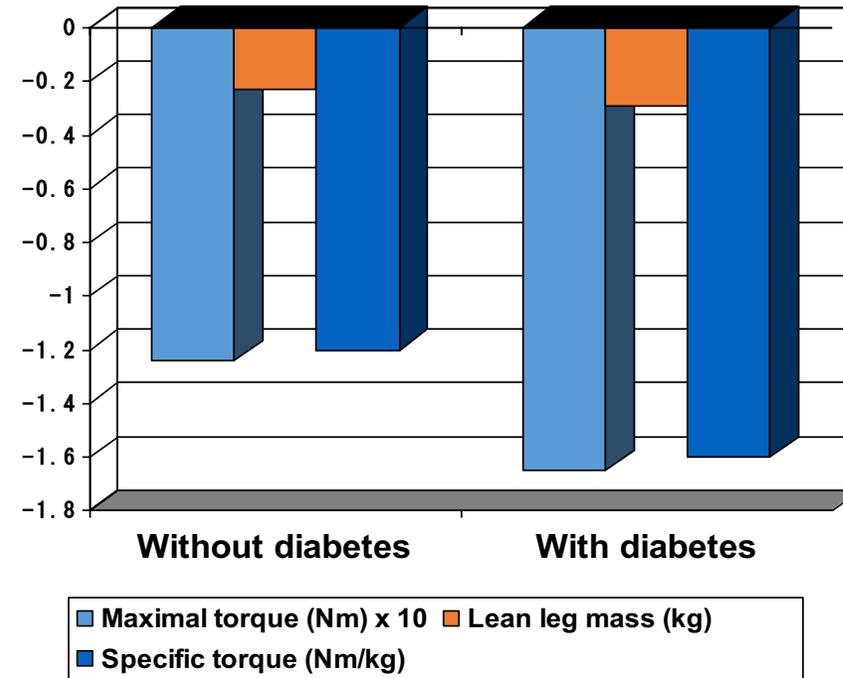
85 yrs (M)

|          |   |
|----------|---|
| <b>A</b> | 33 years old<br>Male<br>BMI: 24.5 kg/m <sup>2</sup><br>Muscle: 588.5 cm <sup>3</sup><br>Subcutaneous fat: 308.4 cm <sup>3</sup><br>Intermuscular fat: 78.7 cm <sup>3</sup>  |
| <b>B</b> | 73 years old<br>Male<br>SPPB: 11<br>Gait speed: 1.15 m/sec<br>BMI: 24.9 kg/m <sup>2</sup><br>Muscle: 461.3 cm <sup>3</sup><br>Subcutaneous fat: 194.7 cm <sup>3</sup><br>Intermuscular fat: 113.8 cm <sup>3</sup> |
| <b>C</b> | 84 years old<br>Male<br>SPPB: 6<br>Gait Speed: 0.49 m/sec<br>BMI: 26.9 kg/m <sup>2</sup><br>Muscle: 364.7 cm <sup>3</sup><br>Subcutaneous fat: 339.3 cm <sup>3</sup><br>Intermuscular fat: 131.7 cm <sup>3</sup>  |

Fig. 1. Representative magnetic resonance images of the femoral region collected at 3T from A) young B) high-functioning older, and C) low-functioning older study groups.

# Diabetes-related accelerated loss of muscle and strength – Park SW et al, Ageing, & Body Composition (ABC) Study 2007

Knee extensor



# Diabetes as a Risk Factor for Dementia and Mild Cognitive Impairment: A Meta-analysis of Longitudinal Studies

G. Cheng et al 2012

- Quantitative meta-analysis of 19 studies from 1996–Dec 2010
- Data from 6184 subjects with diabetes and 38 530 subjects without diabetes
- Main finding – *diabetes was a risk factor for incident dementia (incl AD, VD and any dementia) and MCI*

**Table 3** Summary relative risks of AD, VD and any dementia among subjects with diabetes compared with that without

|                                    | Heterogeneity test |      |         | Random effects |           | Fixed effects |           |
|------------------------------------|--------------------|------|---------|----------------|-----------|---------------|-----------|
|                                    | Chi                | d.f. | P       | RR             | 95%CI     | RR            | 95%CI     |
| Risk for AD                        | 47.3               | 15   | <0.0001 | 1.46           | 1.20–1.77 | 1.54          | 1.40–1.70 |
| Risk for VD                        | 6.3                | 9    | 0.71    | 2.49           | 2.09–2.97 | 2.48          | 2.08–2.96 |
| Risk for any dementia              | 28.9               | 10   | 0.001   | 1.51           | 1.31–1.74 | 1.54          | 1.41–1.67 |
| Risk for mild cognitive impairment | 0.1                | 1    | 0.76    | 1.22           | 1.0–1.45  | 1.21          | 1.02–1.45 |

95%CI, 95% confidence interval; AD, Alzheimer's disease; RR, relative risk; VD, vascular dementia.

# The Mini-Cog Assessment Tool: development of a reliable and quick measure of mental performance in diabetes

## Part A: a three item recall

### Part B:

- Quick and easy to administer
- Participants given a circle (4-10cm in diameter), told that it represents a clock face and instructed to “put in the numbers so that it looks like a clock and the set the time to 10 minutes past 10”
- **Tests executive function and**
  - Auditory comprehension
  - Planning
  - Visual memory and reconstruction
  - Visuo-spatial abilities
  - Motor programming and execution
  - Numerical knowledge
  - Abstract thinking (semantic instruction)
  - Concentration

### Part C – asked to repeat the original three items

- **PILOT study of 207 patients aged 55 – 90 years**
- **Use of Mini-Cog to screen for cognitive impairment in diabetes in primary care settings**

- Use of score of <24 on MMSE, the Mini-Cog had sensitivity of 86%, specificity of 91%, positive predictive value of 54% and negative predictive value of 98%.

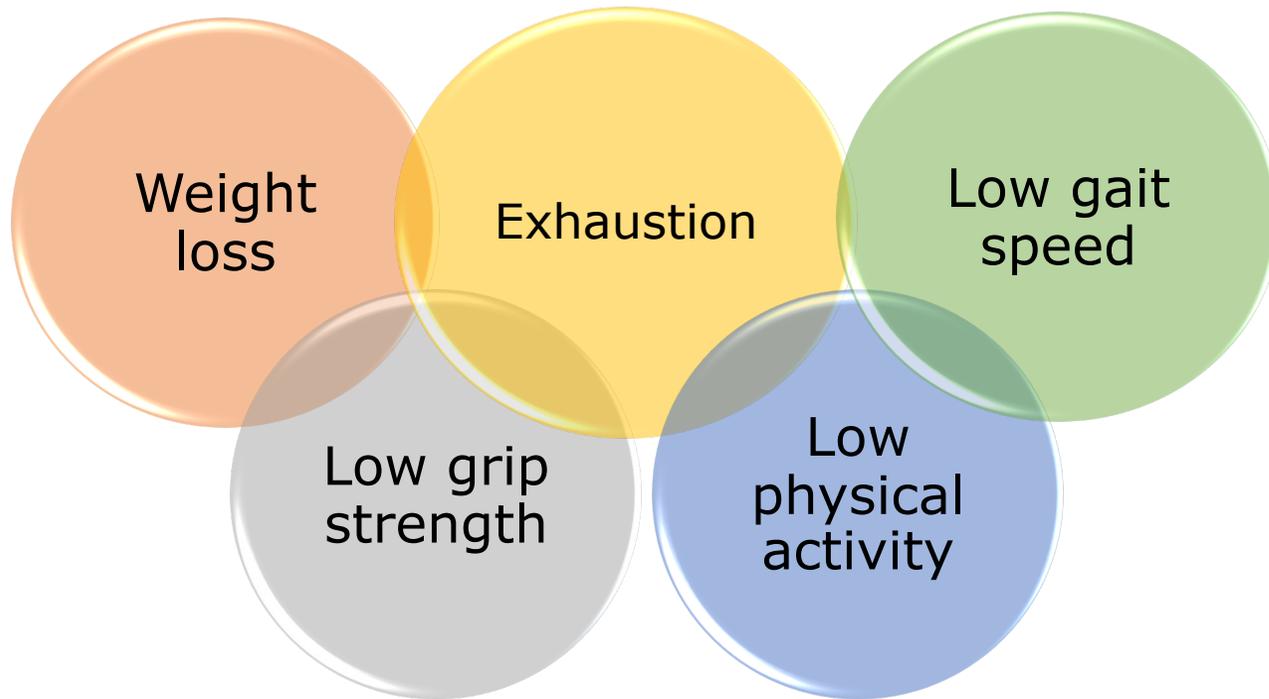
Not influenced by education, culture or language; Performance comparable to MMSE *Borson S. et al, 2000; Shulman, 2000*

| Mini-Cog scores | No. of patients |                                       | No. aged |          |         |        |
|-----------------|-----------------|---------------------------------------|----------|----------|---------|--------|
|                 |                 |                                       | 55-64    | 65-74    | 75-84   | 85+    |
| 0               | 2 (1%)          | <b>35 (17.4%)<br/>screen-positive</b> |          |          | 2(3%)   |        |
| 1               | 4 (2%)          |                                       | 1(2%)    | 1(1%)    | 2(3%)   |        |
| 2               | 9 (4%)          |                                       |          | 4(6%)    | 5(7%)   |        |
| 3               | 20 (10%)        |                                       | 2(4%)    | 8(12%)   | 9(12%)  | 1(11%) |
| 4               | 30 (15%)        |                                       | 7(14%)   | 6(9%)    | 15(20%) | 2(22%) |
| 5               | 136 (68%)       |                                       | 39 (80%) | 48 (72%) | 43(57%) | 6(66%) |



*Sinclair AJ, Gadsby R, Hillson R, Forbes A, Bayer AJ, 2013,*

# Emerging Concepts of Frailty – A multisystem impairment associated with increased vulnerability to stressors



## FRIED Phenotypic Model (Fried L et al, 2001)

Score

- 0–1 = Not frail
- 2 = Pre-frailty
- 3–5 = Frailty

## Cumulative Deficit Model of Frailty: derivation of the Electronic Frailty Index Rockwood K et al, 2007

- The eFI consists of 36 deficits which have been constructed using around 2,000 primary care Read codes
- The eFI calculates a frailty score by dividing the number of deficits present by the total possible: uses 36 validated deficits
- The score is a robust predictor of those who are at greater risk of adverse outcomes: *an eFI > 0.36 have a six-fold increased risk of admission to a care home in the next 12 months and a five-fold increased mortality risk, compared to fit older people*

Clegg A et al, 2016

## Prognostic Outcomes once Frailty is Diagnosed – results from the Cardiovascular Health Study, 2001 – most are preventable with good management

Relative Rates  
Estimated over 3 years

Frail

Incident fall

1.29

Worsening mobility

1.50

Worsening ADL disability

1.98

First hospitalisations

1.29

Death

2.24

# FRAIL TEST – non-invasive frailty screening tool

*Morley JE et al 2012*

## The clinician asks:

**Fatigue:** Are you fatigued?

**Resistance:** Are you unable to walk up one flight of stairs?

**Aerobic:** Are you unable to walk one block?(equivalent of about 200m)

**Illnesses:** Do you have more than 5 illnesses?

**Loss of weight:** Have you lost more than 5% of your weight in the past 6 months?

**Interpretation:** Answers yes to:

≥3: indicates frailty, 1-2: indicates pre-frailty.

## Advantages of Test

- Simple, easy to learn
- Does not require a face to face consultation
- Utilises 4 components of the Cardiovascular Study Index (Fried Criteria) and 1 component from the Rockwood Clinical Frailty Scale
- Correlates well with IADL, gait speed and SPPB
- Valid in late middle age and older adults

*Rosas-Carrasco O et al, 2010 (Mexicans);*

*Li Y et al 2015 (Chinese); Ravindrarajah R et al 2013 (Europeans)*

# Clinical Frailty Scale – *based on the Rockwood Model*

## Clinical Frailty Scale\*



**1 Very Fit** – People who are robust, active, energetic and motivated. These people commonly exercise regularly. They are among the fittest for their age.



**2 Well** – People who have **no active disease symptoms** but are less fit than category 1. Often, they exercise or are very **active occasionally**, e.g. seasonally.



**3 Managing Well** – People whose **medical problems are well controlled**, but are **not regularly active** beyond routine walking.



**4 Vulnerable** – While **not dependent** on others for daily help, often **symptoms limit activities**. A common complaint is being “slowed up”, and/or being tired during the day.



**5 Mildly Frail** – These people often have **more evident slowing**, and need help in **high order IADLs** (finances, transportation, heavy housework, medications). Typically, mild frailty progressively impairs shopping and walking outside alone, meal preparation and housework.



**6 Moderately Frail** – People need help with **all outside activities** and with **keeping house**. Inside, they often have problems with stairs and need **help with bathing** and might need minimal assistance (cuing, standby) with dressing.



**7 Severely Frail** – **Completely dependent for personal care**, from whatever cause (physical or cognitive). Even so, they seem stable and not at high risk of dying (within ~ 6 months).



**8 Very Severely Frail** – Completely dependent, approaching the end of life. Typically, they could not recover even from a minor illness.



**9. Terminally Ill** - Approaching the end of life. This category applies to people with a **life expectancy <6 months**, who are **not otherwise evidently frail**.

### Scoring frailty in people with dementia

The degree of frailty corresponds to the degree of dementia. Common **symptoms in mild dementia** include forgetting the details of a recent event, though still remembering the event itself, repeating the same question/story and social withdrawal.

In **moderate dementia**, recent memory is very impaired, even though they seemingly can remember their past life events well. They can do personal care with prompting.

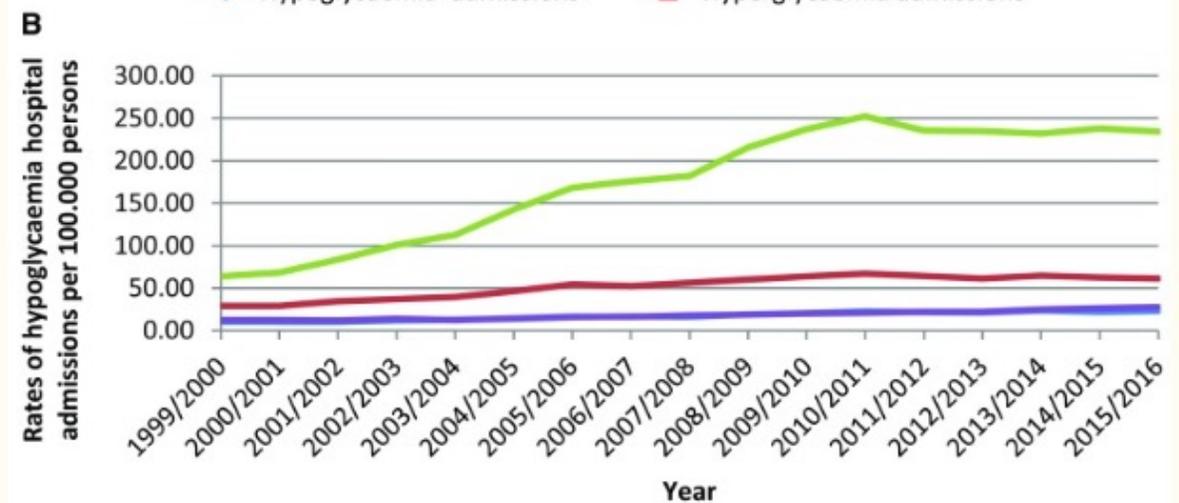
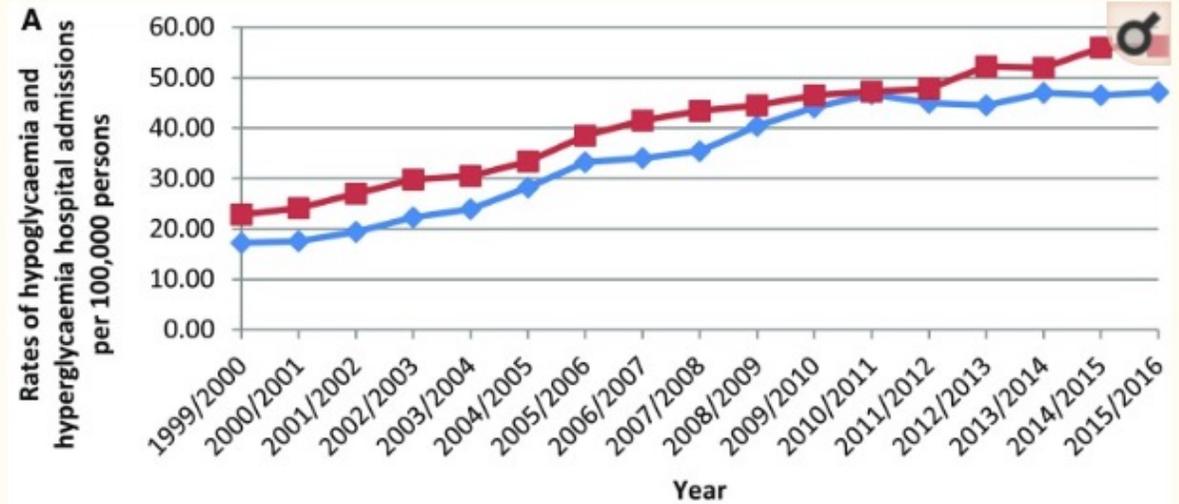
In **severe dementia**, they cannot do personal care without help.

\* 1. Canadian Study on Health & Aging, Revised 2008.  
2. K. Rockwood et al. A global clinical measure of fitness and frailty in elderly people. CMAJ 2005;173:489-495.

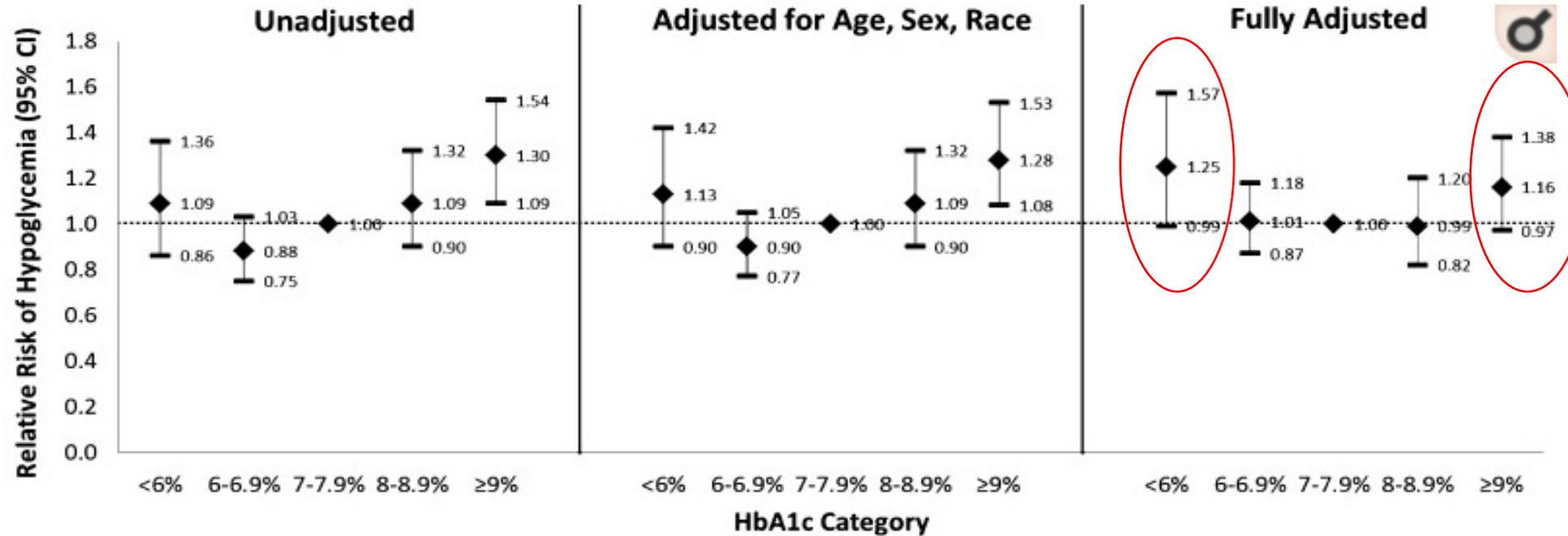
# Hospital Admissions in England and Wales (1999-2016) for Dysglycaemia

Naser AY et al, Diab Ther 2018

- Observational study of all hospital admissions for hypoglycaemia and hyperglycaemia
- Hospital Episode Statistics database in England and the Patient Episode Database for Wales
- **173% increase in hypoglycaemia for the period**
- Strong relationship between hypoglycaemia and parallel increase in glucose-lowering medications



# Hypoglycaemia does not only occur in 'tight control! Data from the Diabetes and Aging Study *Lipska KJ et al, 2013*



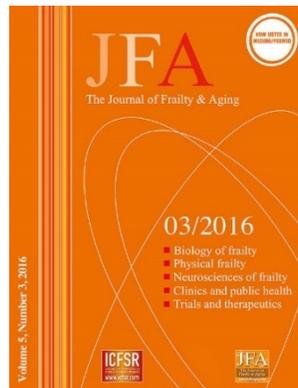
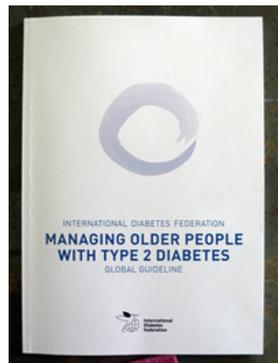
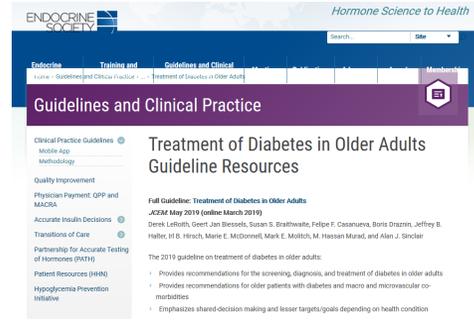
A survey of self-reported severe hypoglycaemia in the past year, n = 9094, aged 60+/-9.8 years; mean HbA1c 7.5% (59 mmol/mol) +/-1.5%.

*Results not altered by age, diabetes duration, or diabetes medication.*

Can we manage type 2 diabetes  
using these perspectives?

# Guidelines DO NOT answer some of the Key Questions in Management

## International Guidelines



## They DO NOT Answer:

- What are the elements of a successful de-escalation approach to management?
- Why a different approach to managing comorbidity and/or frailty is necessary?
- Why a single gluco-centric approach in any case would suit all patient groups and needs of a complex illness model?
- How do we distinguish the various groups of older people with diabetes who are more likely to benefit from specific treatment approaches?
- Should we consider a more precision medicine approach?

# Inherent Difficulties in Managing Older People with Diabetes

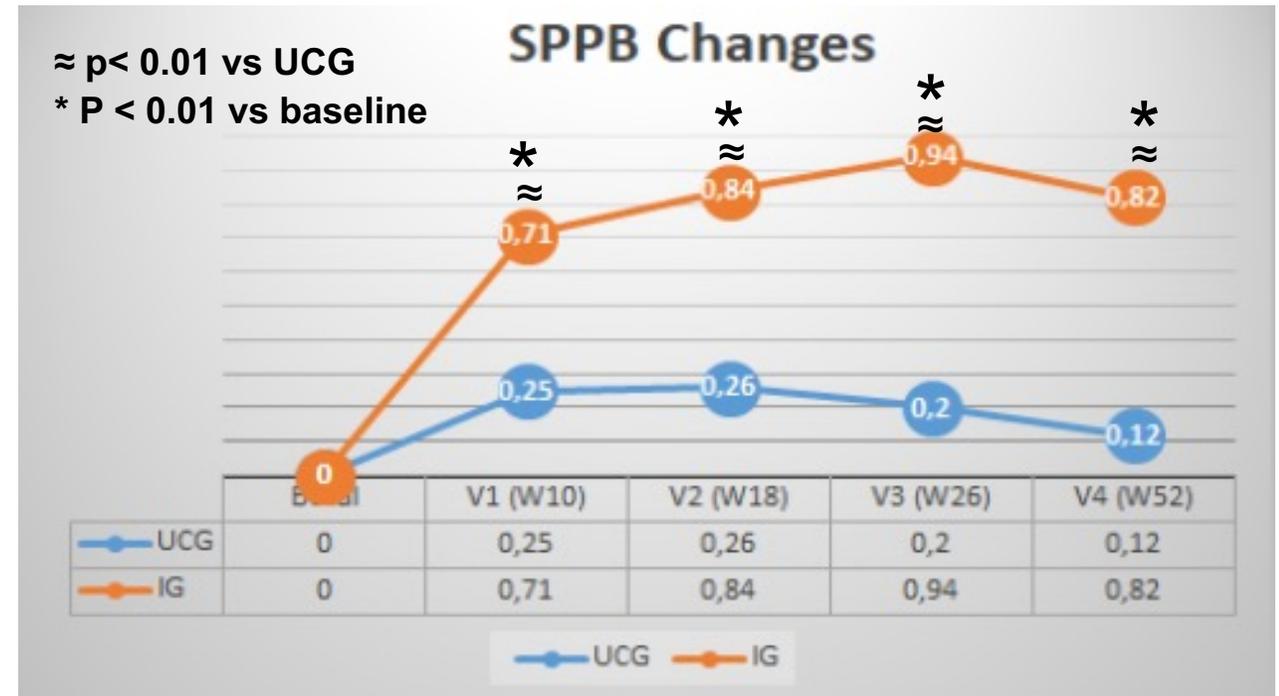
- **The need to consider the significant heterogeneity of an older population of people with diabetes – effects of ageing, changes in renal function, varied susceptibilities to hypoglycaemia, socioeconomic status - see image** →
- **The need to consider the multimorbidity profile and its impact on management**
- **The need to assess the importance and impact of functional loss including frailty, disability and cognitive impairment in setting targets**
- The need to understand better why diabetes self-management can be a challenging prospect in older people
- The need to extrapolate clinical trial evidence from younger populations in the absence of data in older people
- The lack of evaluation in clinical guidelines that target older people with diabetes



# Frailty in Diabetes is reversible



## MAIN RESULTS at 1 YEAR



**J Cachexia Sarcopenia Muscle.** 2019 Apr 23.[Epub ahead of print]  
**Effectiveness of a multimodal intervention in functionally impaired older people with type 2 diabetes mellitus.**

[Rodriguez-Mañas L](#)<sup>1</sup>, [Laosa O](#)<sup>2</sup>, [Vellas B](#)<sup>3</sup>, [Paolisso G](#)<sup>4</sup>, [Topinkova E](#)<sup>5</sup>, [Oliva-Moreno J](#)<sup>6</sup>, [Bourdel-Marchasson I](#)<sup>7</sup>, [Izquierdo M](#)<sup>8</sup>, [Hood K](#)<sup>9</sup>, [Zeyfang A](#)<sup>10</sup>, [Gambassi G](#)<sup>11</sup>, [Petrovic M](#)<sup>12</sup>, [Hardman TC](#)<sup>13</sup>, [Kelson MJ](#)<sup>14</sup>, [Bautmans I](#)<sup>15</sup>, [Abellan G](#)<sup>3</sup>, [Barbieri M](#)<sup>4</sup>, [Peña-Longobardo LM](#)<sup>6</sup>, [Regueme SC](#)<sup>7</sup>, [Calvani R](#)<sup>11</sup>, [De Buyser S](#)<sup>12</sup>, [Sinclair AJ](#)<sup>16</sup>; [European MID-Frail Consortium](#).

# Co-morbid illness attenuates the expected benefits of intensive glucose control in older patients with type 2 diabetes

*Huang ES 2008*

- Use of decision analysis (multiple prediction models)
- Based on UKPDS, NIH Model, and Health and Retirement Study data
- Simulation for patients 40-80y with type 2 diabetes using 5y age
- Compared the projected health benefits of moderate control (HbA1c 7.9%, 63 mmol/mol) with intensive control (HbA1c 7%, 53 mmol/mol)

Similar findings in patients up to 15 years of diabetes duration

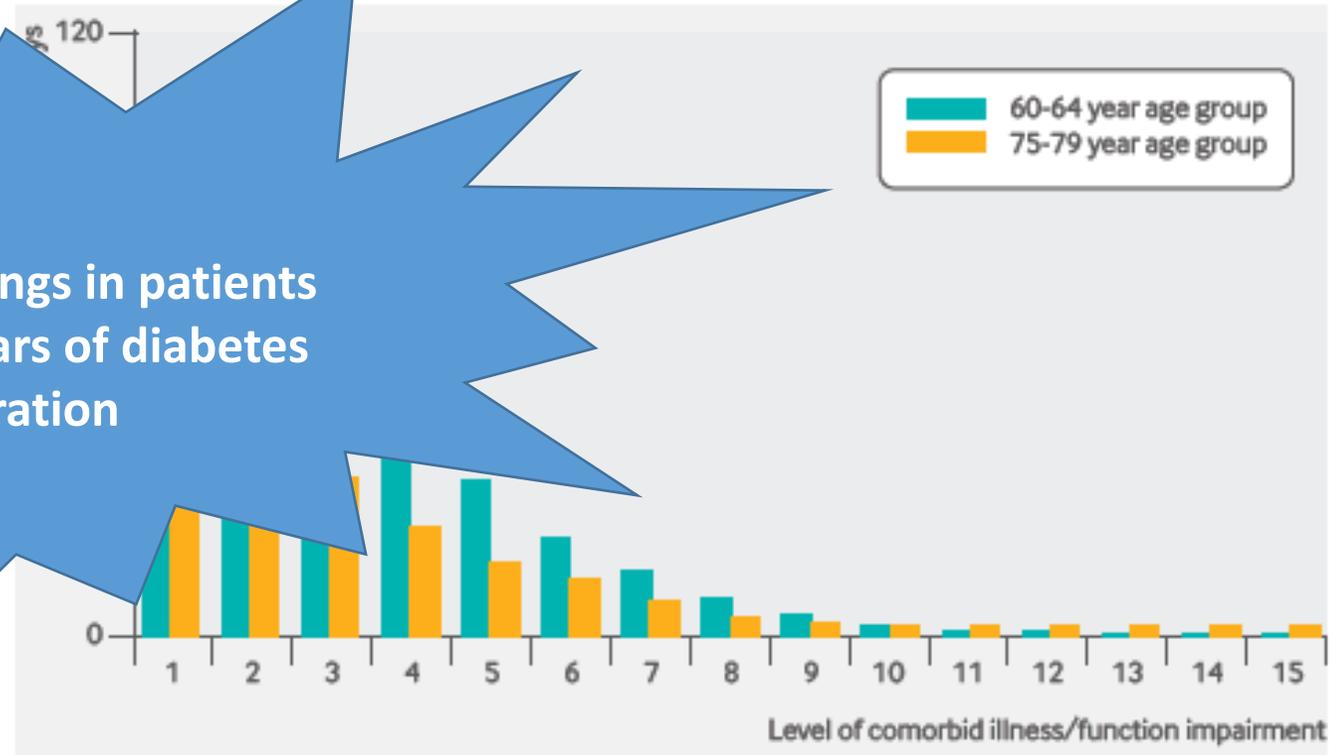
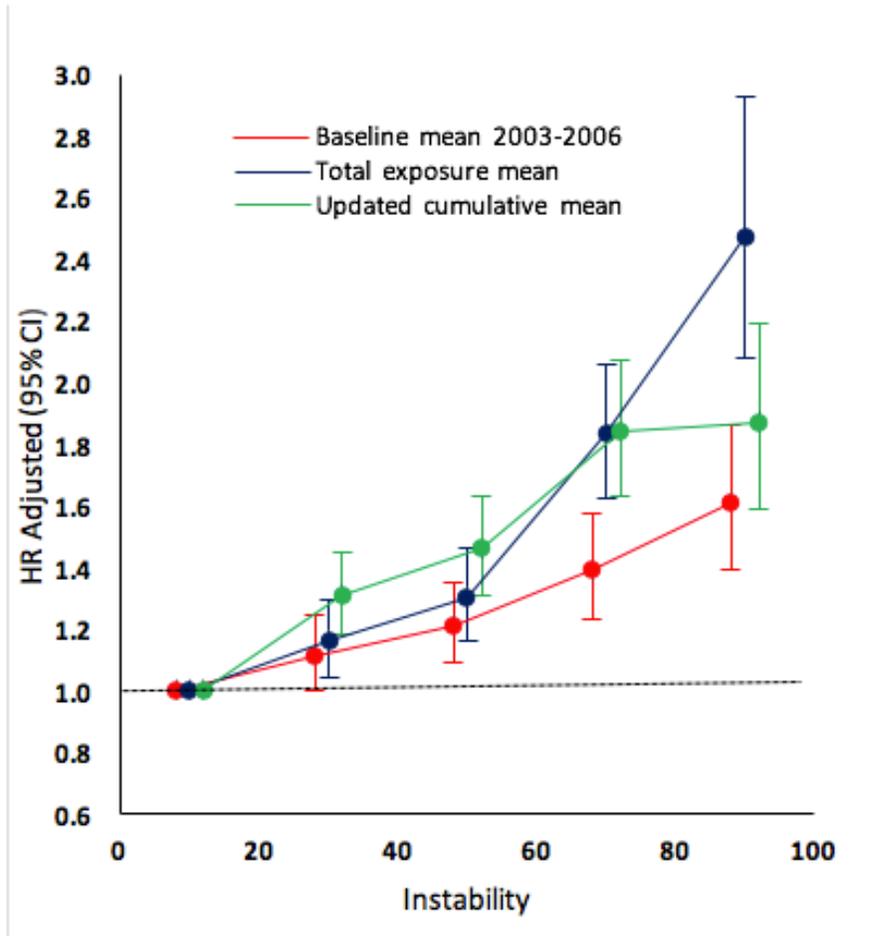


Fig 3 | Expected quality of life benefits of intensive glucose control for 60-64 year old and 75-79 year old patients with newly diagnosed diabetes, with increasing levels of comorbid illness and functional impairment<sup>58</sup>

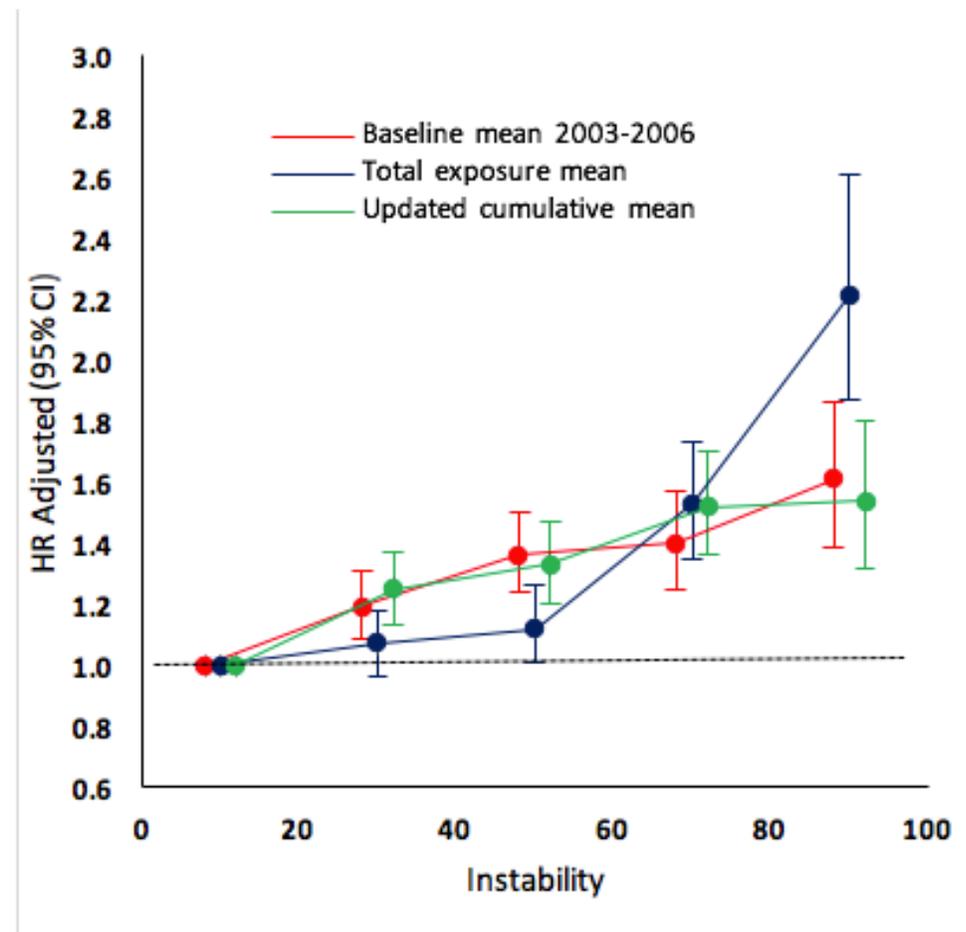
# Adjusted HRs by HbA1c level for all-cause mortality in women (a) and men (b) – importance of a stable level of glycaemia. *Lancet D&E, 2018*

Error bars are 95% CIs. HR=hazard

a) Females, age 70 and over, duration of diabetes five years



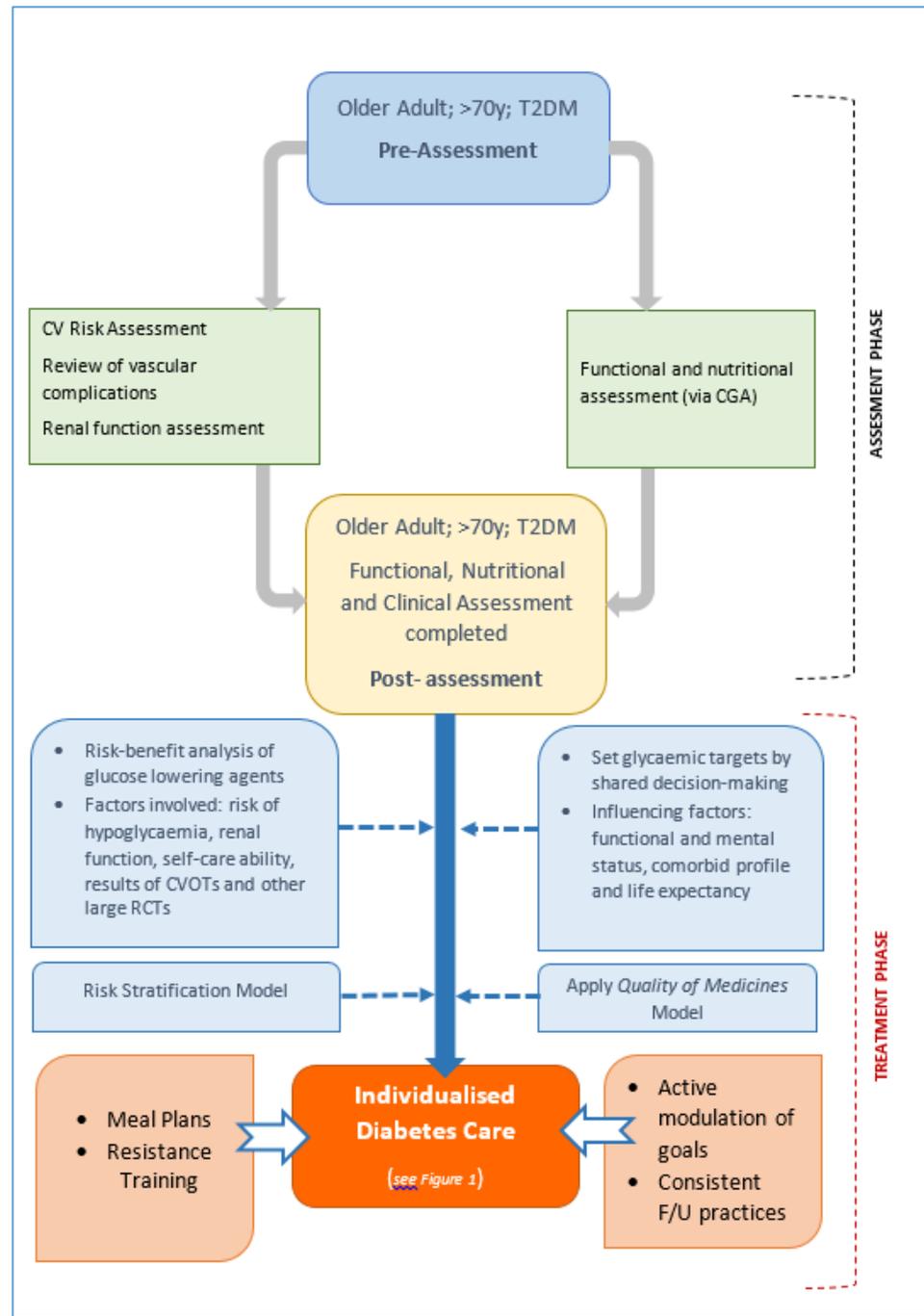
b) Males, age 70 and over, duration of diabetes five years



# Working towards Individualised Care

**Lancet paper**

*Sinclair AJ, Abdelhafiz A, Forbes A, Munshi M, Diabetic Medicine 2018*



# Newer Therapies - Results of Subgroup Analyses in Established CVD—Recent Cardiovascular Outcome Trials – but what do they tell us about older people?

| Trial                 | Agent                     | Outcome | HR, 95% confidence interval |                     |                     |
|-----------------------|---------------------------|---------|-----------------------------|---------------------|---------------------|
|                       |                           |         | Age <65Y                    | Age ≥65-75Y         | Age ≥75Y            |
| EMPA-REG              | Empagliflozin             | 3P-MACE | 0.88, 0.78 to 0.99.         | 0.88, 0.78 to 0.93. | 0.68, 0.46 to 1.00. |
| DECLARE-TIMI          | Dapagliflozin             | 3P-MACE | 0.88, 0.78 to 0.99.         | 0.88, 0.78 to 0.93. | 0.94, 0.65 to 1.36. |
| CANVAS                | Canagliflozin             | 3P-MACE | 0.88, 0.78 to 0.99.         | 0.88, 0.78 to 0.93. | NR                  |
| SGLT-2i meta-analysis | SGLT-2i CV outcome trials | 3P-MACE | 0.88, 0.78 to 0.99.         | 0.88, 0.78 to 0.93. | NR                  |
| LEADER                | Liraglutide               | 3P-MACE | 0.88, 0.78 to 0.99.         | 0.79 to 1.02.       | 0.66, 0.49 to 0.89  |
| SUSTAIN-6             | Semaglutide               | 3P-MACE | 0.74, 0.65 to 0.85.         | 0.71, 0.61 to 1.02  | NR                  |
| EXSCEL                | Exenatide                 | 3P-MACE | 1.05, 0.92 to 1.21.         | 0.80, 0.71 to 0.91. | NR                  |
| GLP-1RA meta-         | GLP-1RA CV outcome        | 3P-MACE | 0.89, 0.76 to 1.03.         | 0.86, 0.80 to 0.92. | NR                  |

- Both SGLT-2i and GLP-1RA reduce 3P-MACE in younger (65Y) and older (≥65y) patients with type 2 diabetes.
- These agents are generally well-tolerated in older people with less risk of hypoglycaemia.
- Older (>75Y) people are less represented in these trials, therefore generalisation of findings is limited by under reporting in this age group.

# Glucose lowering treatment

## (a) A scheme for treating older adults with type 2 diabetes using a 3-step approach \*

### Evidenced-Based Strategies for Glucose Lowering Therapy in Older People

#### Key Steps

#### 1<sup>st</sup> Step:

Metformin is the first line of treatment after lifestyle – caution in severe renal impairment; consider low hypo potential SU or DPP4 inhibitor if MF contraindicated

#### 2<sup>nd</sup> Step:

All other oral agents can be used (e.g. DPP4-I or SU or SGLT2-inhibitor) depending on clinician choice, renal function, frailty status, risk potential for hypoglycaemia, economic considerations (q.v. sulphonylureas – use of gliclazide); if patient is markedly obese (>35) consider GLP-1 agonist; if all OHA are not tolerated consider long-acting basal insulin or GLP-1 agonist

#### 3<sup>rd</sup> Step:

Add in basal insulin or a pre-mixed insulin, or a GLP-1 agonist

Consider the need to prevent worsening of heart or renal function with SGLT2 or GLP1 analogue

#### Key Considerations

- Have a 'risk to benefit' conversation
- Estimate likelihood of worsening renal or hepatic function
- Estimate risk of hypoglycaemia
- Try not to put HbA1c at the heart of your planning – consider quality of life and minimising vascular risk as your main priorities

#### Cautions in Frailty

Consider a **glinide** if eating patterns are irregular (short duration/rapid onset of action) or cognitive impairment;

Consider **avoiding** a SGLT2-Inhibitor in view of weight loss, dehydration, toe amputations;

**Caution** with a GLP-1 agonist (weight loss, anorexia) but as part of a glucose-dependent strategy may reduce hypoglycaemia rate;

**Pioglitazone**: caution with side effects but may be of value in those with high stroke and macrovascular risk

# Individualised Metabolic Targets for Older People – EDWPOP, IDF, AES and ADA Guidelines

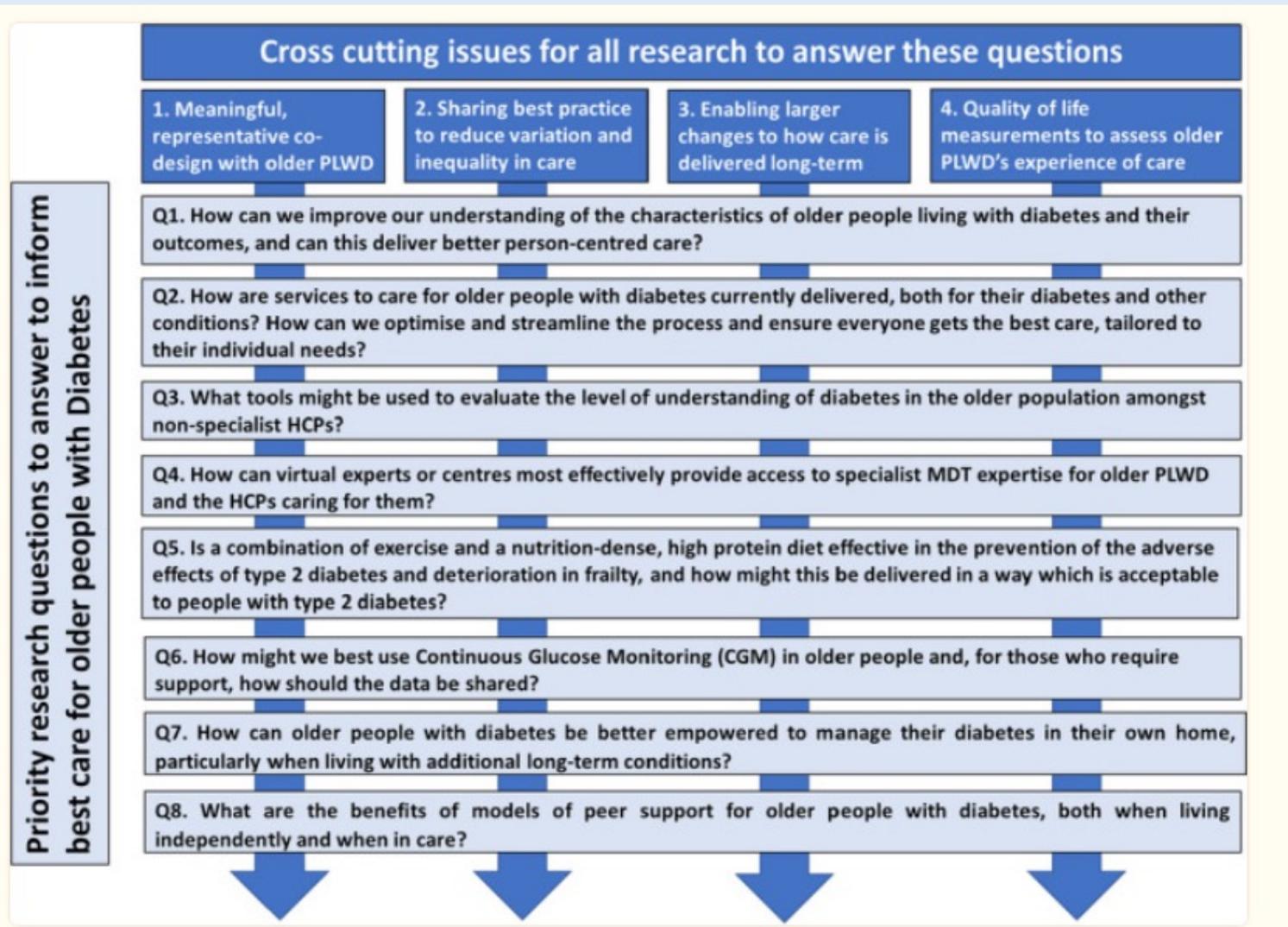
| Target                 | EDWPOP (2011)    |                                | IDF (2013)               |                  | AES (2019)             |  | ADA (2022)                |   |
|------------------------|------------------|--------------------------------|--------------------------|------------------|------------------------|--|---------------------------|---|
|                        | Patient group    | Target                         | Patient group            | Target           | Patient group          | Target                                 | Patient group             | Target  |
| HbA1c mmol/mol (%)     | Non-frail        | 53-59 (7.0-7.5%)               | Functionally independent | 53-59 (7.0-7.5%) | Good health            | < 58 mmol/mol (7.5%)                   | Functionally independent  | A1C <7.0–7.5% (53-58 mmol/mol)                              |
|                        |                  |                                | Functionally dependent   | 53-64 (7.0-8.0%) | Intermediate health    | <64 - ≥58 mmol/mol (<8.0% and ≥7.5%)   |                           |   |
|                        | Frail            | 60-69 (7.6-8.5%)               | Frail                    | <69 (<8.5%)      | Poor health            | <69 and ≥64 mmol/mol (<8.5% and ≥8.0%) | Highly comorbid/dependent | A1C < 8.0% (64 mmol/mol)                                    |
| BP, mmHg               | Non –frail       | <140/80                        | Functionally independent | <140/90          | All aged ≥65 - ≤85 y   | 140/90                                 | Most Older Adults         | <140/90   |
|                        | Frail            | <150/90                        | Frail                    | <150/90          |                        |  |                           |   |
| LDL-cholesterol mmol/l | All older adults | Statins unless contraindicated |                          | <2.0             | All aged 65 y or older | Statins unless contraindicated         | All older adults          | Statins – moderate to intense therapy depending on CVD risk |

# Ageing well with diabetes: A workshop to co-design research recommendations for improving the diabetes care of older people.

*A Diabetes UK initiative (2021). Wylie TAF et al*



The charity for  
people with diabetes

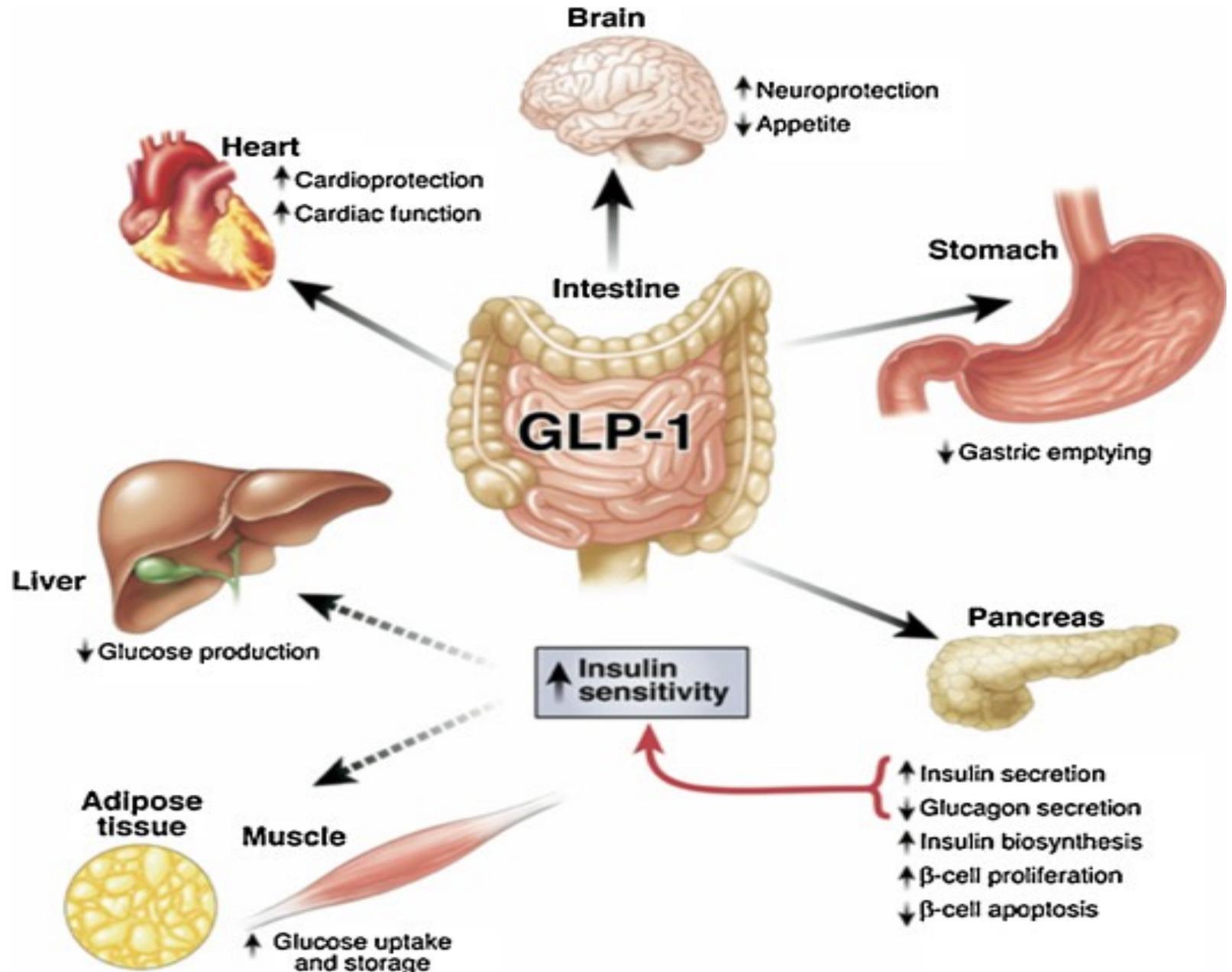


# Conclusions

- Diabetes in older people can OFTEN BE MANAGED SUCCESSFULLY with an individualized management approach
- However, the wide heterogeneity of this condition requires an assessment of key problems that arise to limit effective target setting
- An appreciation of both frailty and other comorbidities is a primary requisite to successful care in diabetes



# How does GLP-1 work?



# How is an injectable GLP-1 agonist given?

- Pen devices are used
- GLP 1 agonists are given as a fixed dose – no titration
- Usually a low dose is given for the first 1 – 2 weeks
- Then a standard dose is given as ongoing therapy
- Can be given with most OGLTs – DPP4s will be stopped
- Can be given once/twice daily or weekly



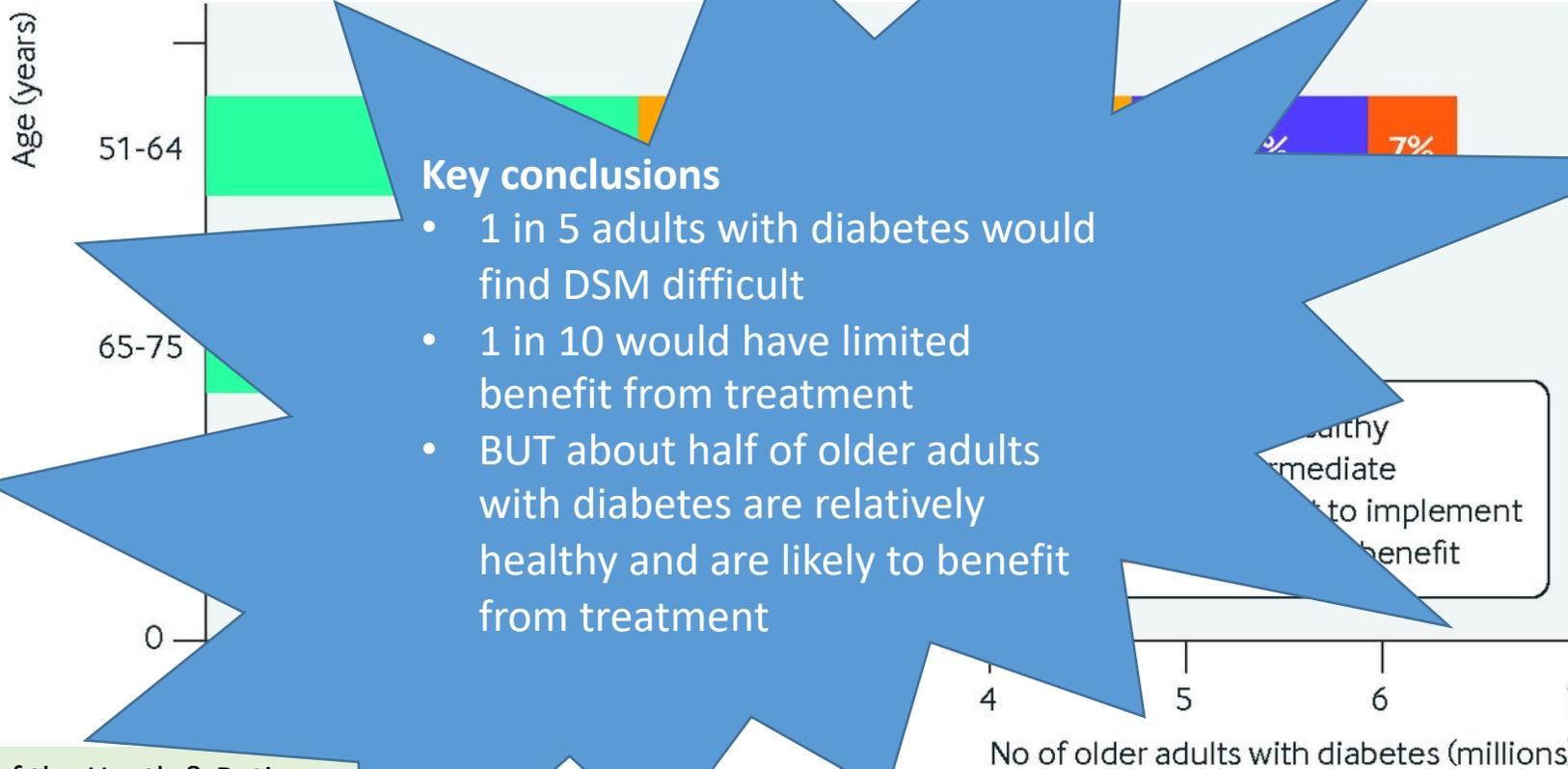
# Frequencies of adults with diabetes in clinical groups by age in United States Health and Retirement Study.

*Blaum C et al, 2010*

**Class 1: relatively healthy – little or no comorbidities**

**Class 2: complex illness profile where self-care may be difficult**

**Class 3: significant multimorbidity profile/functional impairment**



# Overtreatment of hyperglycemia in older people, *Lipska KJ et al, 2015*

Figure 1. Achieved Glycemic Control Among Older US Adults With Diabetes Mellitus Across 3 Health Status Categories

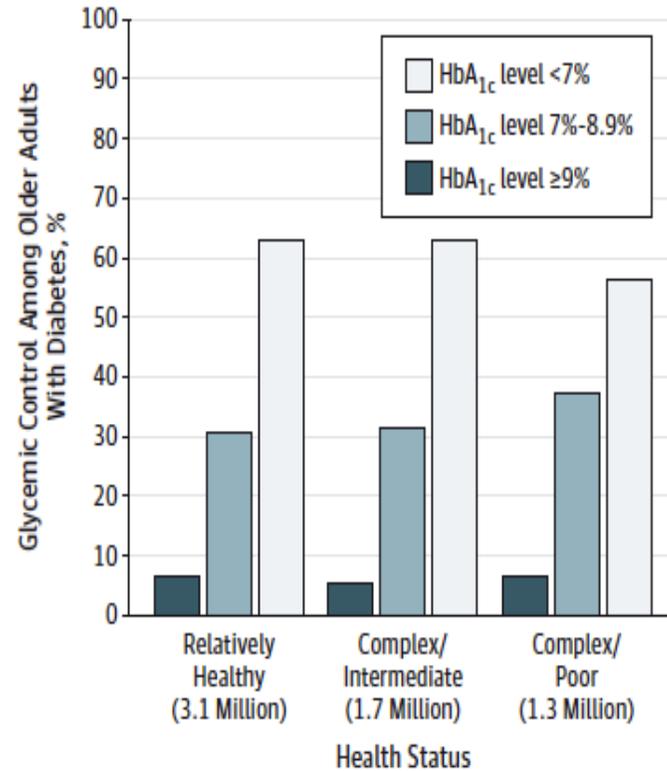
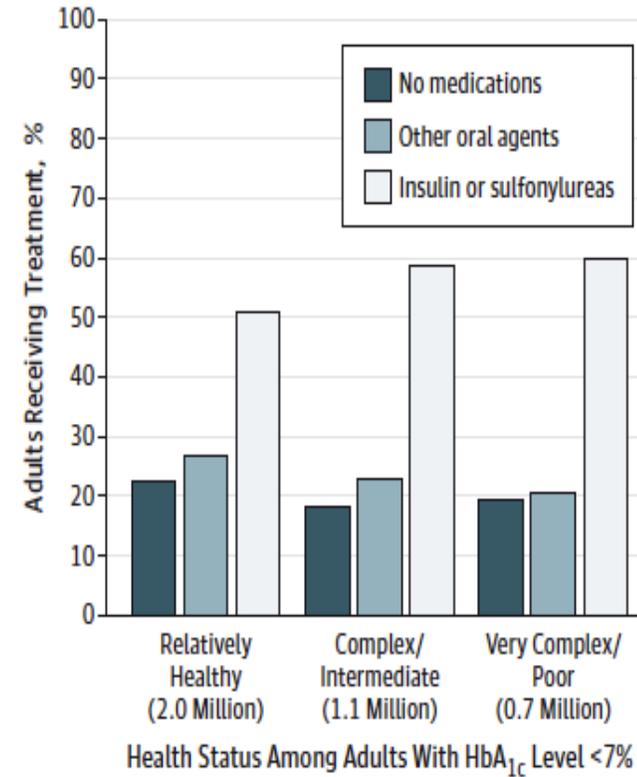
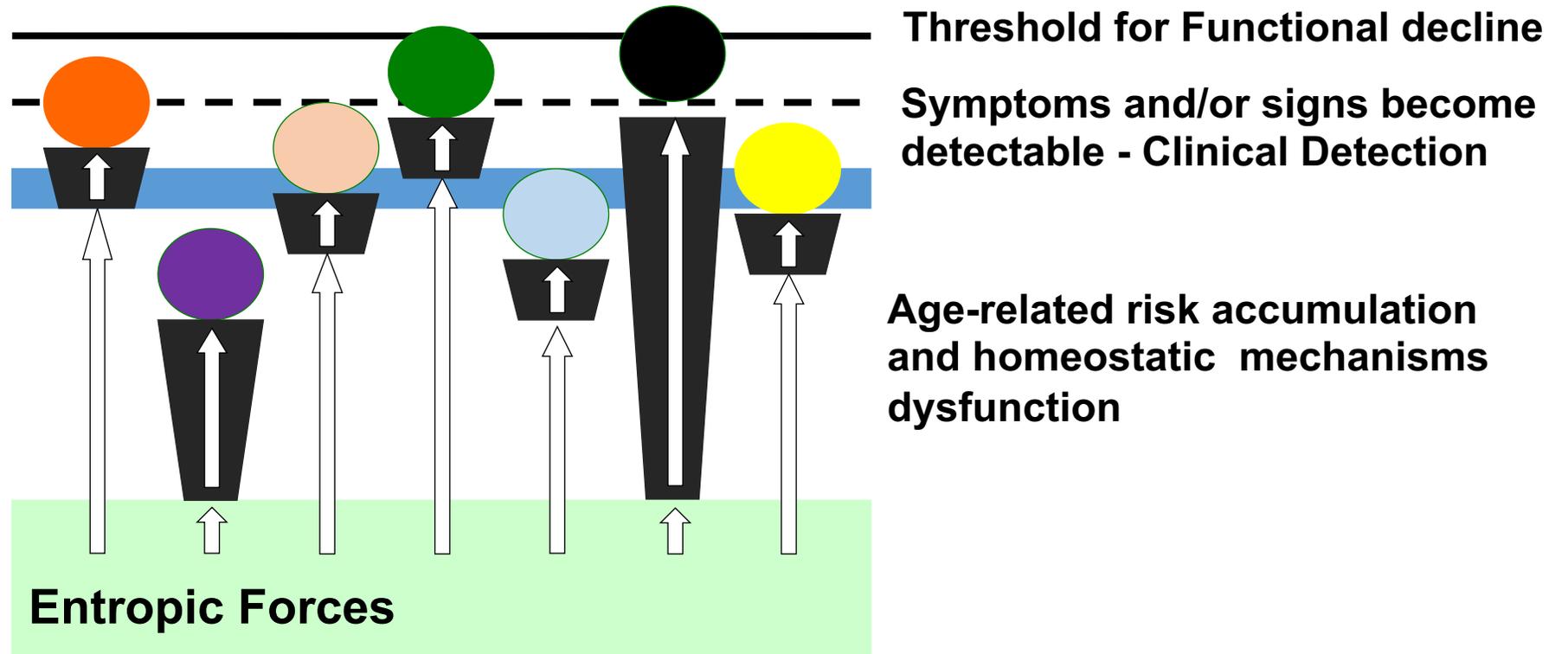


Figure 2. Treatment of Older US Adults With Diabetes Mellitus With an HbA<sub>1c</sub> Level of Less Than 7% Across Health Status Categories



# Age-related Model of Disease and Function



# GPs, new diabetes indicators - Frailty

11 May 2017, Be the First to Comment

## Management of patients with severe frailty under the 2017/18 GP contract

From 1 July 2017 practices will be contractually required to identify and manage patients aged over 65 who are living with moderate to severe frailty. This article has been updated to include links to new guidance.

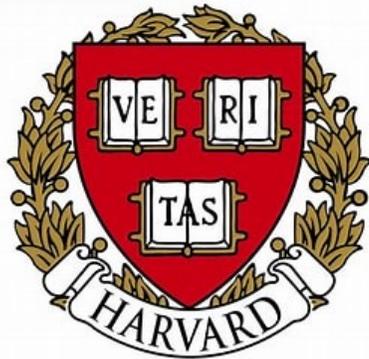


Practices should use an appropriate tool to identify patients that fall into this category, such as the Electronic Frailty Index (eFI). The BMA says these tools should be used as a guide only, and 'the decision to code some as moderately or severely frail should be made by an experienced clinical guided by, but not score.'

## The indicators

- NM157 – The percentage of patients with diabetes without moderate or severe frailty, on the register, in whom the last IFCC-HbA1c is 58mmol/mol or less in the preceding 12 months.
- **NM158 – The percentage of patients with diabetes with moderate or severe frailty, on the register, in whom the last IFCC-HbA1c is 75mmol/mol or less in the preceding 12 months.**
- NM159 – The percentage of patients with diabetes without moderate or severe frailty, on the register, in whom the last blood pressure reading (measured in the preceding 12 months) is 140/80mmHg or less.
- NM160 – The percentage of patients aged 25-84 years, with a diagnosis of type 2 diabetes, without moderate or severe frailty, not currently treated with a statin, who have had a consultation for a cardiovascular risk assessment using a risk assessment tool agreed with the NHS Commissioning Board in the last three years.
- NM161 – The percentage of patients with a diagnosis of type 2 diabetes and a recorded CVD risk assessment score of  $\geq 10\%$  (without moderate or severe frailty), who are currently treated with a statin (unless there is a contraindication or statin therapy is declined).
- NM162 – The percentage of patients with diabetes aged 40 years and over, with no history of CVD and without moderate or severe frailty, who are currently treated with a statin (excluding patients with type 2 diabetes and a CVD risk score of  $< 10\%$  recorded in the preceding 3 years).

# Evidenced-Based Care: Diabetes in Older People



## Invited Review

### Evidence-based diabetes care for older people with Type 2 diabetes: a critical review

A. J. Sinclair<sup>1</sup> , A. H. Abdelhafiz<sup>2</sup>, A. Forbes<sup>3</sup>  and M. Munshi<sup>4</sup>

<sup>1</sup>Foundation for Diabetes Research in Older People, Diabetes Frail Ltd, Droitwich, <sup>2</sup>Rotherham General Hospital, Rotherham, <sup>3</sup>Kings College, London, UK and <sup>4</sup>Harvard Medical School and Joslin Clinic, Boston, MA, USA

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#### Abstract

In our ageing society diabetes imposes a significant burden in terms of the numbers of people with the condition, diabetes-related complications including disability, and health and social care expenditure. Older people with diabetes can represent some of the more complex and difficult challenges facing the clinician working in different settings, and the recognition that we have only a relatively small (but increasing) evidence base to guide us in diabetes management is a limitation of our current approaches. Nevertheless, in this review we attempt to explore what evidence there is to guide us in a comprehensive scheme of treatment for older adults, often in a high-risk clinical state, in terms of glucose lowering, blood pressure and lipid management, frailty care and lifestyle interventions. We strive towards individualized care and make a call for action for more high-quality research using different trial designs.

Diabet. Med. 00: 1–15 (2018)

#### Introduction and background

It is estimated that in 2017 there were 451 million (age 18–99 years) people with diabetes worldwide, and these figures are expected to increase to 693 million by 2045 [1]. A major shift in the epidemiology of diabetes has been to those aged 60–79 years [2]. Apart from this advancing tide of older people with diabetes, the ageing process itself is increasing the number of people living with the sequelae of ill health, chronic diseases, frailty and injuries, all of which enhance disability and functional decline, and pose real clinical challenges and burdens in those with Type 2 diabetes [3]. Older people with diabetes should be a priority target for focused interventions that bring about improved cardiovascular outcomes, enhanced safety and improved survival if the latter has worthwhile disability-free years and associated quality of life [4]. The important area of Type 1 diabetes in older adults is outside the scope of this review but must be addressed in due course.

We recognize that older people with diabetes can span four decades (ages 60–90 years and older), are not a homogeneous group and range from robust adults still in employment to frail residents of nursing homes. Thus, their cognitive and physical status vary widely, and they often have complex health and social care needs [4]. We therefore consider that our review of the literature in general pertains to those aged 70 years and over because

the risks of comorbid illness, frailty and dependency begin to rise after this age, but we accept that other organizations may define being 'old' as less or more than 70 years [5]. It is also important to recognize that to produce valid and evidence-based recommendations for care, it is usually necessary to extrapolate research findings from clinical trials in younger adults, which is a limitation that has implications for developing clinical guidelines [6]. The modern management of older people with diabetes requires an acceptance by clinicians that recommendations of care should be tailored to the individual and take into consideration important factors such as changes in functional status, the comorbid illness profile, whether or not a person is dependent and their estimated life expectancy. These can have a marked influence on management goals, what care model is adapted, and how ongoing and follow-up care is delivered. We call this an 'individualizing care' scheme (Box 1).

Diabetes care for older people is often not straightforward for the reasons cited above, but as advancing age brings about increasing complexity of both the person with diabetes and the management of the illness itself, clinicians face greater challenges to their skills and competence. The different pathway to Type 2 diabetes in older individuals compared with younger individuals reflects changes in body composition, marked changes in insulin resistance in muscle and adipose tissues, a decrease in  $\beta$ -cell capacity and loss of normal insulin pulsatility, and the progressive negative effects on glucose tolerance of comorbid illness, onset of

Correspondence to: Alan Sinclair. E-mail: Sinclair.5@btinternet.com.