

# DIABETES MELLITUS

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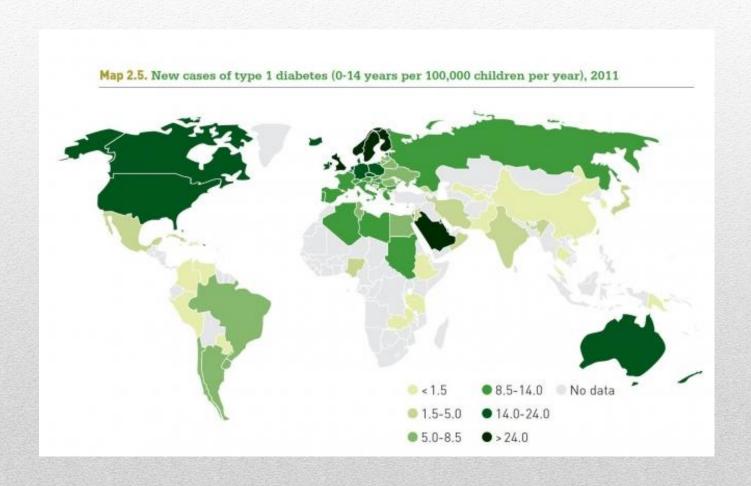
### **Outline**

- Epidemiology
- Pathophysiology
- Clinical Context
- Complications
- Treatment Plan
- Prognosis

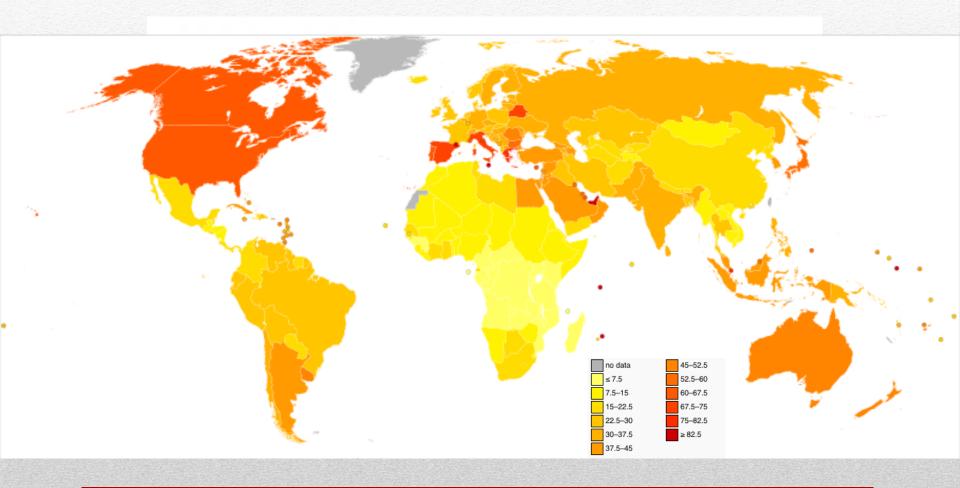
### DM I vs. DM II

|                          | TYPE I                              | TYPE II                                |  |
|--------------------------|-------------------------------------|--|--|
| Clinical<br>Presentation | Juvenile<br>Thin/Normal Body<br>DKA | Adult/Elderly Overweight Ethnic Groups |  |
| Cause                    |                                     |  |  |
| Prevalence               | 5%                                  | 95%                                    |  |
| Onset/Progression        | Abrupt                              | Gradual                                |  |
| Endogenous Insulin       | Absent                              | Normal, Low/High                       |  |
| Concordance              | 50%                                 | 90%                                    |  |
| Symptoms                 | Severe                              | Less severe                            |  |
| Treatment                | Insulin                             | Weight loss<br>Hypoglycaemics          |  |

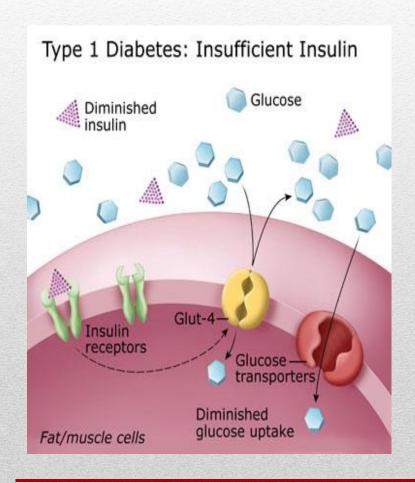
# **Epidemiology**

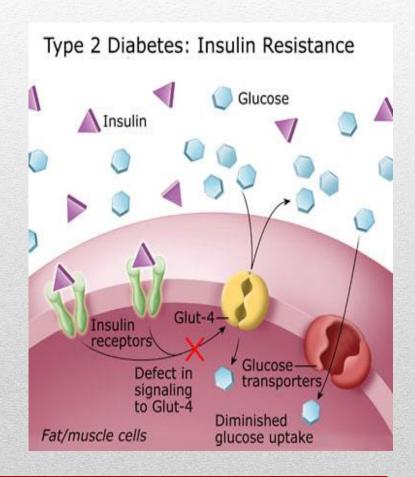


# **Epidemiology**



### **Pathophysiology**





## **Secondary Causes**

- Acromegaly,
- Cushing syndrome,
- Thyrotoxicosis,
- Pheochromocytoma
- Chronic pancreatitis,
- Cancer
- Drug induced:
  - Atypical Antipsychotics
  - Beta-blockers
  - Calcium Channel Blockers
  - Corticosteroids
  - Thiazide Diuretics

# Signs & Symptoms



### Investigations

- Random Plasma Glucose (>11.1 mmol/L or >200mg/dL)
- Fasting Plasma Glucose (>7.0 mmol/L or >126mg/dL)
- Oral Glucose Tolerance Test (>11.1mmol/L)
- Antibodies
- Insulin/C-peptide Levels
- Glycated Haemoglobin A1c (6.5%)
- Urinary Glucose/Albumin
- Serum Lipids
- Urea/Electrolytes

## Diagnosis

- Symptoms of diabetes (polyuria, polydipsia and unexplained weight loss) with:
  - A random venous plasma glucose of > 11.1 mmoI/L
  - or fasting plasma glucose is greater than or equal to 7.0 mmoI/L
  - or 2 h post 75g oral glucose load of greater than or equal to 11.1 mmoI/L
- With **no symptoms**, diagnosis should **not** be based on a single glucose value, but requires confirmation with another value in the diabetic range, ideally collected fasting on more than one occasion.



## **Multidisciplinary Team**

### **HOSPITAL TEAM**

- Endocrinologist
- Diabetes Specialist Nurse
- Dietician
- Podiatrist
- Optometrist
- Clinical Psychologist

### **COMMUNITY TEAM**

- General Practitioner
- Practise Nurse
- Community Dietician
- Social Worker

SPECIALIST CLINICS

## Monitoring

### CLINICAL ASSESSMENT

- •Weight/BMI
- •Symptoms of hyperglycaemia
- •Hypoglycaemic attacks if any
- •Problems with medication
- •Problems with eyesight, parasthesia, impotence
- •Any symptoms of angina or claudication
- Patient monitoring records

### **RETINOPATHY**

Ensure patient has follow up either with their optometrist or at hospital retinal screening clinic

### RENAL DISEASE

- •Check sitting BP
- •Send urine for microalbumin screening

### **FOOT CARE**

(Annual review by podiatrist)

- Assessment of peripheral pulses
- •Assessment of sensations with 10g monofilament
- •Foot care education
- •Plan appropriate package of care relative to risk

### **GLYCAEMIC CONTROL**

- •Patient's own monitoring at home
- •Check HbA1c

### DIETARY AND LIFE STYLE REVIEW

- •Review diet- dietician
- •Smoking/alcohol
- •Advise on appropriate exercise
- •Advise on Diabetes (UK) membership benefits

### **BIOCHEMICAL TESTS**

- •HbA1c
- •Lipid profile, creatinine, TFT
- •Urine for albumin creatinine ratio (ACR)

### INTERIM REVIEWS AS NECESSARY

- •Dietary/lifestyle review
- Continuing education
- •BP measurement
- •Patient monitoring record
- •Check HbA1c (minimum interval between two tests at least 3 months)
- •Creatinine/TFTs if indicated

# Complications

- Diabetic Ketoacidosis
- Hyperosmolar Hyperglycaemic Nonketotic Syndrome
- Coronary Artery Disease
- Hypertension
- Diabetic Nephropathy
- Diabetic Retinopathy
- Peripheral Vascular Disease

### **Diabetic Ketoacidosis**

- Diabetes Type I
- Breakdown of fat and overproduction of ketones by the liver and loss of bicarbonate
- Occurs when Diabetes Type 1 is undiagnosed or known diabetic has increased energy needs (Physical/emotional stress)
- Signs & Symptoms
  - Kussmaul Breathing
  - Fruity Breath
- Treatment
  - Insulin
  - Monitor K<sup>+</sup> levels
  - Restore fluid balance
  - Treat Underlying Condition

### HHNS

- Diabetes Type 2
- Enough insulin is secreted to prevent ketosis, but not enough to prevent hyperglycemia
  - Marked hyperglycemia (>50mmol) with no ketones
- High blood sugar causes an extreme diuresis with severe electrolyte and fluid loss
- Signs & Symptoms
  - Altered mental state, focal signs
  - Hyperviscosity
  - Polydipsia
- Treatment
  - Same as DKA (less insulin, more cautious fluid replacement and MUST use heparin)

### **Chronic Diabetic Complications**

- Microvascular
  - Nephropathy
  - Retinopathy
  - Neuropathy
- Macrovascular
  - Heart disease
  - Stroke
  - Peripheral vascular disease
  - Hypertension

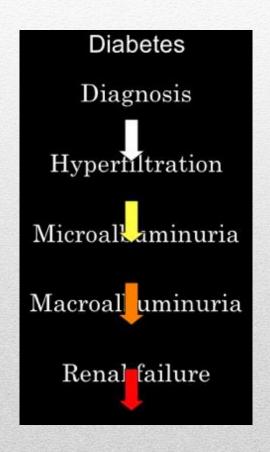
### **RISK FACTORS**

- Hyperglycaemia
- Hypertension
- Dyslipidaemia
- Excess weight
- Smoking
- Disease duration
- Family history

### **Diabetic Nephropathy**

- Definition: glomerular changes in kidneys of diabetics leading to impaired renal function
- Diabetics without treatment go on to develop hypertension, edema, progressive renal insufficiency
- In type 1 diabetics, 10 15 years
- May occur soon after diagnosis with type 2 diabetes since many are undiagnosed for years
- Most common cause of end-stage renal failure
- Kimmelstiel-Wilson syndrome: glomerulosclerosis associated with diabetes

# **Diabetic Nephropathy**



| Stage | Description  | GFR               |
|-------|--|-------------------|
| 1     | Kidney damage with normal or increase in GFR       | >90               |
| 2     | Kidney damage with normal or mild reduction in GFR | 60-89             |
| 3a    | Moderate reduction of GFR                          | 45-59             |
| 3b    |  | 30-44             |
| 4     | Severe reduction in GFR                            | 15-29             |
| 5     | Renal failure                                      | <15 (or dialysis) |

### **Treatment Plan**

 Diet and Exercise Oral hypoglycaemic therapy Insulin Therapy

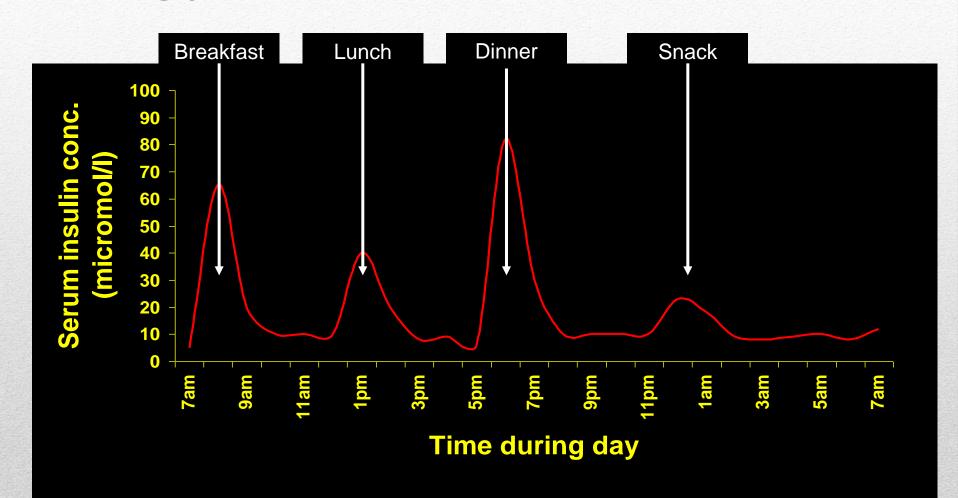
## Lifestyle Changes

- Physical Activity
  - Promotes weight loss and insulin sensitivity
  - Moderate-intensity recommended (2.5 hours walking/week)
- Diet
  - Well-balanced and scheduled meals
  - Carbohydrates 60-70%, Protein 15-20%, Fats < 10%
  - Limited alcohol/sodium intake
  - Stay hydrated

# Medication

| Drug Class                            | Drug Name                             | Brand Name  | Mechanism of Action  |
|---------------------------------------|---------------------------------------|---|--|
| Biguanides                            | Metformin                             | Glucophage®   | Inhibit glucose<br>production by the liver                   |
| Sulfonylure as<br>(second-generation) | Glimepiride<br>Glipizide<br>Glyburide | Amaryl®<br>Glucotrol®<br>Diabeta®,<br>Glynase PresTab®,<br>Micronase® | Increase insulin<br>secretion by pancreatic<br>beta cells    |
| Meglitinides                          | Repaglinide<br>Nateglinide            | Prandin®<br>Starlix®  | Increase insulin secretion<br>by pancreatic beta cells       |
| Thiazolidinediones<br>(TZDs)          | Pioglitazone<br>Rosiglitazone         | Actos®<br>Avandia®  | Increase glucose uptake<br>by skeletal muscle                |
| Alpha-glucosidase<br>inhibitors       | Acarbose<br>Miglitol                  | Precose®<br>Glyset®   | Inhibit carbohydrate<br>absorption in the small<br>intestine |

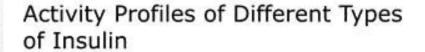
### Insulin

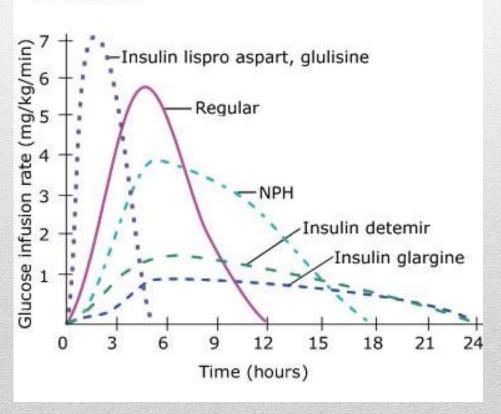


# Insulin

| Types of insulin  |  |   |   |  |  |  |
|---|--|---|---|--|--|--|
| Insulin type/action<br>(appearance)   | Brand names<br>(generic name<br>in brackets)   | Basal/bolus                                   | Dosing schedule   |  |  |  |
| Rapid-acting analogue (clear) Onset: 10–15 minutes Peak: 60–90 minutes Duration: 4–5 hours  | Humalog®<br>(insulin lispro)<br>NovoRapid®<br>(insulin aspart)   | Bolus   | Usually taken right<br>before eating<br>or to lower<br>high blood glucose |  |  |  |
| Short-acting (clear) Onset: 0.5–I hour Peak: 2–4 hours Duration: 5–8 hours  | Humulin®-R<br>Novolin®ge Toronto   | Bolus   | Taken about 30 minutes before eating, or to lower high blood glucose      |  |  |  |
| Intermediate-acting (cloudy) Onset: I-3 hours Peak: 5-8 hours Duration: up to 18 hours  | Humulin®-N<br>Novolin®ge NPH   | Basal   | Often taken at<br>bedtime, or twice<br>a day (morning<br>and bedtime)     |  |  |  |
| Extended long-acting analogue<br>(Clear and colourless)<br>Onset: 90 minutes<br>Peak: none<br>Duration: 24 hours  | Lantus®<br>(insulin glargine)<br>Levemir®<br>(insulin detemir)   | Basal   | Usually taken<br>once or twice<br>a day                                   |  |  |  |
| Premixed (cloudy) A single vial contains a fixed ratio of insulins (the numbers refer to the ratio of rapid- or fast-acting to intermediate-acting insulin in the vial) | Humalog® Mix 25 <sup>™</sup><br>Humulin® (20/80,<br>30/70)<br>Novolin®ge (10/90,<br>20/80, 30/70,<br>40/60, 50/50) | Combination<br>of basal and<br>bolus insulins | Depends on the combination  |  |  |  |

### Insulin





## **Prognosis**

- Lifelong disease
- No cure
- Usual cause of death CVD/Stroke
- Prognosis dependent on:
  - Time of diagnosis
  - Progress of diabetic complications
  - Compliance with treatment
- Life Expectancy
  - Type I 20 years less
  - Type II 10 years less