

# **Hammersmith Hospitals Trust**

## **Guidelines for Management of Diabetes Mellitus**

# Hammersmith Hospitals Trust

## Guidelines for Management of Diabetes Mellitus

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# Glossary

AAA	Abdominal aortic aneurysm
ABPIs	ankle brachial pulse index
Ang2	Angiotensin 2
ACE	Angiotensin converting enzyme
BM	Finger prick glucose measurement
BMI	Body Mass Index
BP	Blood pressure
CHD	Coronary heart disease
CI	Contra-indication
C-peptide	Insulin C-peptide
CSII	Continuous subcutaneous insulin infusion
CT	Computer tomography
CVA	Cerebrovascular disease
CVD	Cardiovascular disease
CXH	Charing Cross Hospital
CXR	Chest X-ray
DM	Diabetes mellitus
DI	Diabetes insipidus
DKA	Diabetic ketoacidosis
DNS	Diabetic nurse specialist
DP	Dorsalis pedis
DVLC/DVLA	Driving and vehicle licensing authority
DVT	Deep venous thrombosis
ECG	Electrocardiogram
ETT	Exercise Tolerance Test
HGV	Heavy goods vehicle
HH	Hammersmith Hospital
HONK	Hyperosmolar non-ketotic state
GDM	Gestational diabetes mellitus
GI	Gastrointestinal
GP	General Practitioner
IFG	Impaired fasting glucose
IGT	Impaired Glucose Tolerance
IHD	Ischaemic heart disease
IM	Intra muscular
IRMAs	Intra retinal
IV	Intravenous
JPS	Joint position sense
LP	Lumbar puncture
LT	Light touch

MELAS	Mitochondrial myopathy, encephalopathy, lactic acidosis and stroke
MI	Myocardial infarction
MODY	Maturity onset Diabetes on the young
MSU	Mid stream urine
NVD	Neovascularisation of the disk
NVE	Neovascularisation elsewhere
NVI	Neovascularisation of the iris
OGD	Oesophago-gastroduodenoscopy
OGTT	Oral glucose tolerance test
OM	Osteomyelitis
PGR	Prandial glucose regulator
PP	Pin prick sensation
PSV	Public service vehicle
PT	Posterior tibial
PVD	Peripheral vascular disease
SC	Sub Cutaneous
SRD	State registered dietician
SU	Sulphonylurea
TIA	Transient ischaemic attack
UKITC	UK islet cell transplantation consortium
VS	Vibration sense
Wt	Weight

# Referrals and Contacts

## Urgent referrals:

**Telephone Endocrine Registrar and Diabetes specialist nurses and Fax referral letter**

1. Likely to need insulin
  - Significant ketonuria at any age
  - Young, non-obese, ill, vomiting or rapid wt loss
2. Risk of Non-ketotic hyperosmolar coma
  - ,Elderly, intercurrent illness, glucose >25
3. Sudden deterioration in visual acuity
4. Foot ulceration
5. Pregnancy

## Routine referrals:

**Supply the following information to assess priority**

1. Diagnostic criteria for DM (ie fasting, random or OGTT results)
2. Latest HbA1c
3. Body Mass Index and weight change
4. Symptoms (severity, duration, rapidity of onset)
5. Blood pressure
6. Evidence of existing complications
7. All present medications
8. Past medical history
9. Occupation and contact number (working hours)

## Contact Numbers

### Hammersmith Hospital

Endocrinology/Diabetes Secretaries	Tel: 020 8383 4828 Fax: 020 8383 3360
Diabetic Nurse Specialists	Tel: 020 8383 4693 Fax: 020 8383 2348
Podiatry	Tel: 020 8383 4616
Dietetics	Tel: 020 8383 3048
Appointments	Tel: 020 8383 5000 Fax: 020 8383 8383
24h emergency contact number	Tel: 020 8383 1000 (Ask for endocrine SpR)

### Charing Cross Hospital

Endocrinology/Diabetes Secretaries	Tel:020 8846 1065	Fax: 020 8846 1862
Diabetic Nurse Specialists	Tel: 020 8846 1062	Fax: 020 8846 1080
Podiatry	Tel: 020 8846 1621	
Dietetics	Tel: 020 8846 1445	
Appointments	Tel: 020 8383 5000	Fax: 8346 7564
24h emergency contact number	Tel: 020 8383 1000	(Ask for endocrine SpR)



# **Background and Introduction**

## **Background**

Since the first version of the diabetic handbook in 1993 the DCCT and UKPDS studies have greatly increased the evidence base for the management of Diabetes and these revised guidelines have been prepared over 2001-2002 to document in a readily accessible form our standard practice in both the in and out patient settings of secondary care.

## **Introduction**

These guidelines have been primarily developed in a highly visual flow chart format for the use of medical staff in secondary care but may also be of use to primary care.

The guidelines contain information on departmental policy as regards patient referrals, discharges and patient DNAs together with copies of standard letters and proformas.

The Hammersmith Hospitals trust diabetic clinics has around 5,000 patients under follow up and received more than 900 referrals per year and these guidelines aim to help deliver a consistently high quality service to our patients, to improve communication between primary and secondary care and to reduce waiting times for both new and follow up appointments.

# Diabetes Diagnosis and Classification

## A) Presenting and associated features

Polyuria, Polydipsia, Nocturia, Weight loss, Blurred vision  
Recurrent cutaneous sepsis, Balanitis, Pruritis vulvae  
Foot ulcers, Neuropathy  
Ischemic heart disease, Cerebrovascular disease  
Peripheral vascular disease

## B) Risk Factors

Ethnicity, Indian-Asian, Afro-Caribbean, Middle -Eastern  
First Degree Relative  
Obesity  
Recurrent stillbirths; babies > 4.5Kg

## C) Diagnostic Criteria

### **HBA1c CAN NOT BE USED FOR DIAGNOSIS**

#### **2x Random Glucose Measurement (1x if symptomatic)**

=11.1            Diabetes Mellitus  
<11.1           Normal if =6.1 do 2x fasting

#### **2x Fasting Glucose Measurement (1x if symptomatic)**

=7.0            Diabetes Mellitus  
=6.1 but <7.0   Impaired Fasting Glucose do OGTT

#### **Oral Glucose Tolerance Test 2h Glucose**

=11.1            Diabetes Mellitus  
=7.8 but <11.1   Impaired Glucose Tolerance  
<7.8            Normal

## D) Aetiology of Diabetes Mellitus

**Type I diabetes:** Idiopathic or autoimmune immune mediated  $\beta$  cell destruction leading to insulin deficiency

**Type II diabetes:** Insulin resistance with relative insulin deficiency associated with obesity, hypertension and dyslipidaemia

## E) Other Causes

### **Genetic defects of $\beta$ cell function**

Mitochondrial: maternal inheritance (nt3243 of Leu tRNA identical to MELAS mutation)

Wolfram's syndrome: AR lack of  $\beta$ -cells (DM,DI,hypogonadism,deafness,optic atrophy)

Maturity-Onset Diabetes of the Young (MODY): <25yr. dominant, impaired insulin secretion

MODY-1 : (20q) hepatic nuclear factor-4 $\alpha$ ?

MODY-2 : (7p) Defective glucokinase

MODY-3 : (12q24) hepatic nuclear factor-1 $\alpha$ ? most common

? MODY-4 insulin promotor factor 1(IPF-1) and hepatic nuclear factor-1 $\beta$ ?

### **Genetic defects in insulin action**

Type A insulin resistance: associated with acanthosis nigricans and PCOS

Leprechaunism : insulin receptor mutation abnormal facial features

Rabson-Mendenhall syndrome : insulin receptor mutation abnormal teeth, nails, pineal

Lipodystrophy : error in post insulin receptor signalling

### **Pancreatic Destruction**

Cystic fibrosis, pancreatitis, pancreatic surgery, haemachromatosis

### **Endocrinopathies**

Hyperthyroidism, Cushings, Acromegaly, Glucagonoma, Pheochromocytoma

**Drug induced**

Glucocorticoids; Thyroxine, thiazides,  $\beta$ -blockers, Nicotinic acid, phenytoin, proteinase inhibitors

**Gestational Diabetes**

# Impaired Glucose Tolerance

## Definition of IGT

Oral glucose tolerance test 2h sample greater than or equal to 7.8 but <11.1

## Incidence and progression of IGT

Present in 10% to 30% of >65 year olds

On repeat testing

30% revert to normal on repeat testing

50% continue to show IGT

20% are diabetic

(Progression to overt diabetes occurs at 5-10% pre year)

Patient with IGT have increased risk coronary artery, peripheral vascular and cerebrovascular disease.

## Management

Annual fasting glucose : proceed to OGTT if greater than or equal to 6.1 but <7.0

Annual blood pressure and lipids

Avoid thiazides,  $\beta$ -blockers and oral steroids when possible

## Life style advice

Aim for BMI of 25

If BMI>30 consider

1. Orlistat 120mg with meals  
(2.5Kg loss in 4 weeks before starting and stop if <5% wt loss in 3 months)
2. Sibutramine 10mg od (15mg after 1 month if <2kg loss)  
(Stop if wt loss <5% in 3 months)
3. Metformin 500mg bd

Regular exercise average of 30min/day

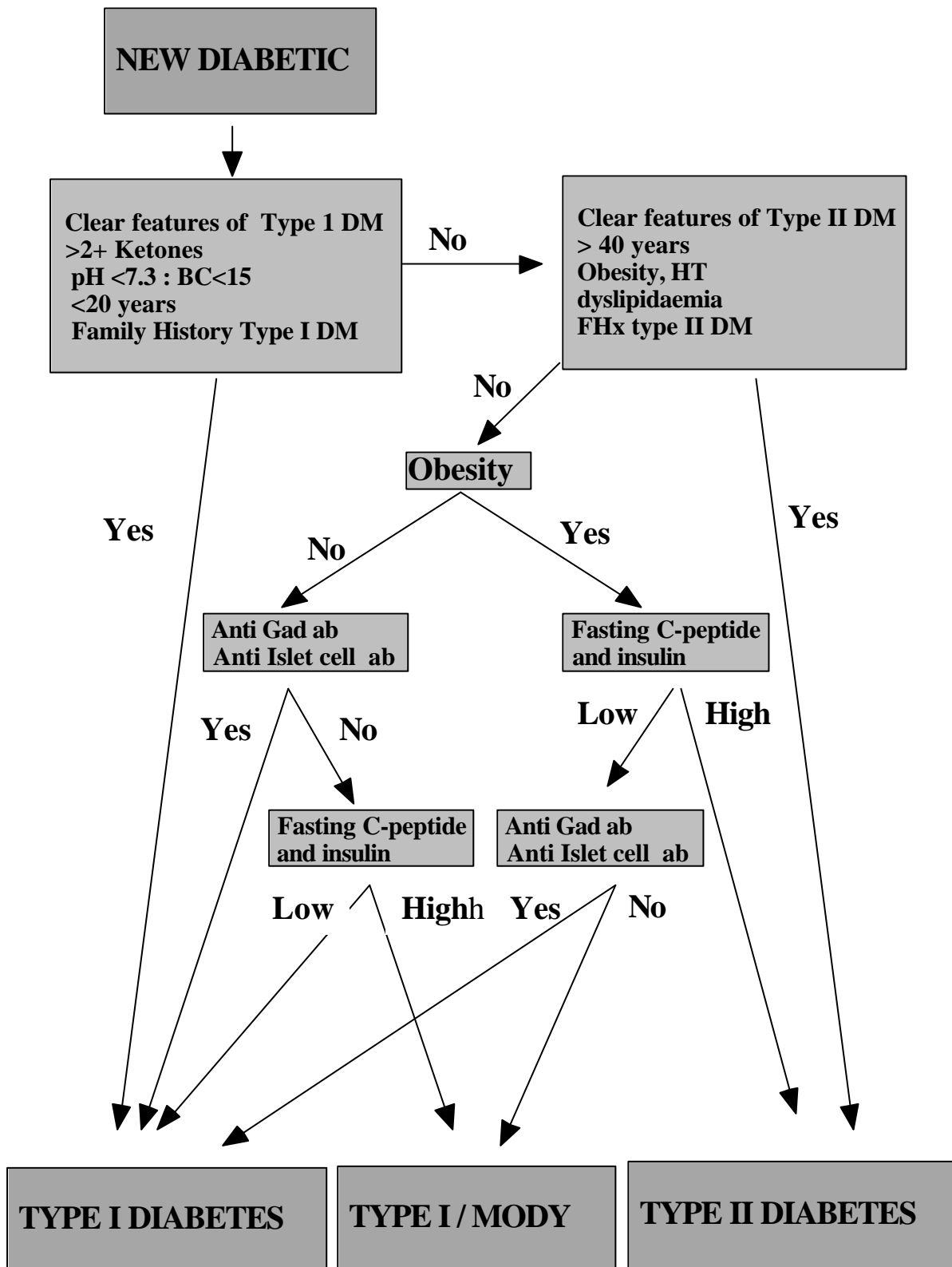
Stop smoking

Moderate alcohol intake may be beneficial

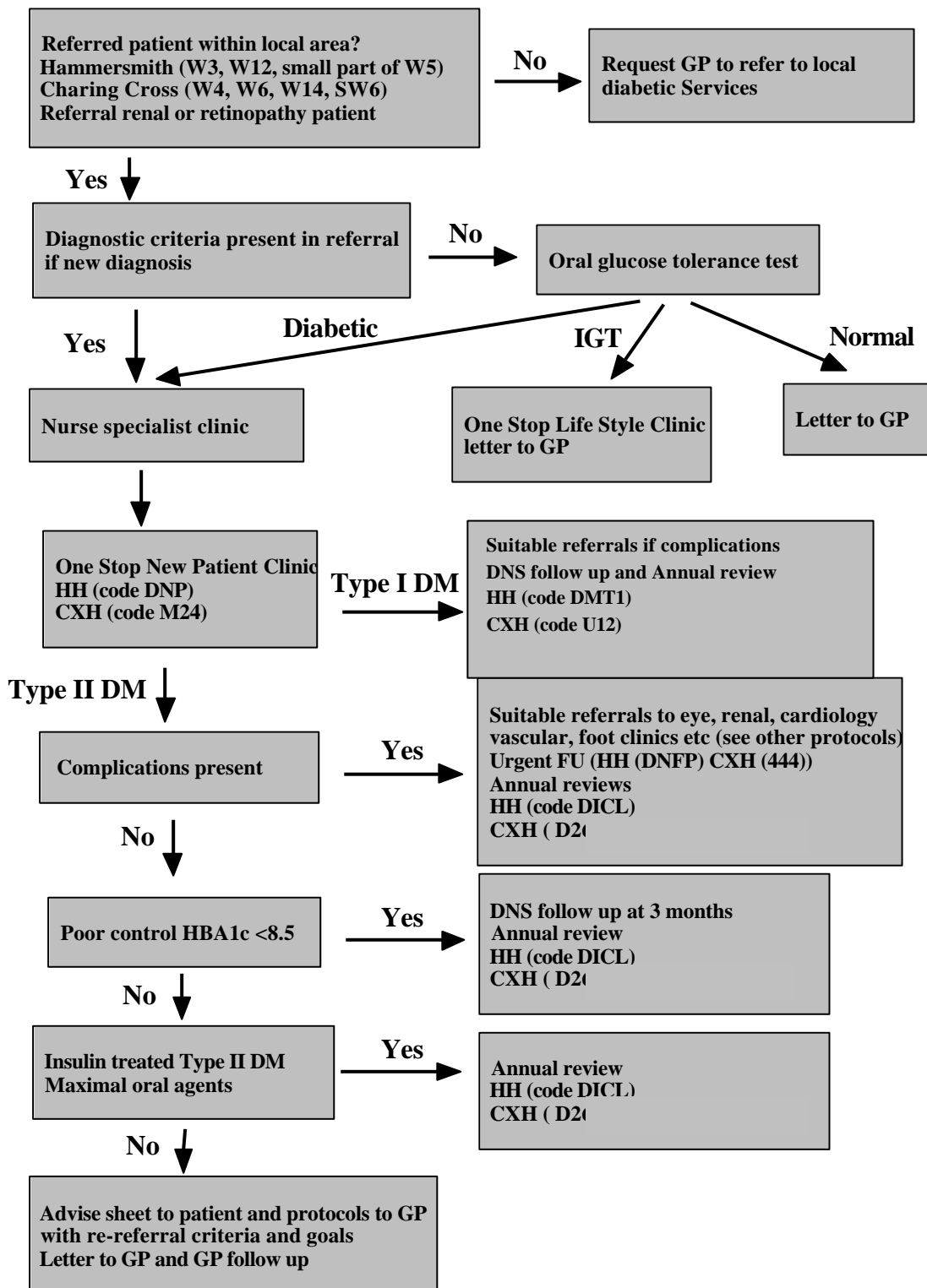
Dietary advise on cholesterol lowering (treatment if 10y CHD risk >30%)

Minimising these risk factors can reduce the incidence of diabetes by 50%

# Types of Diabetes Mellitus



# Management of New Diabetic Referrals



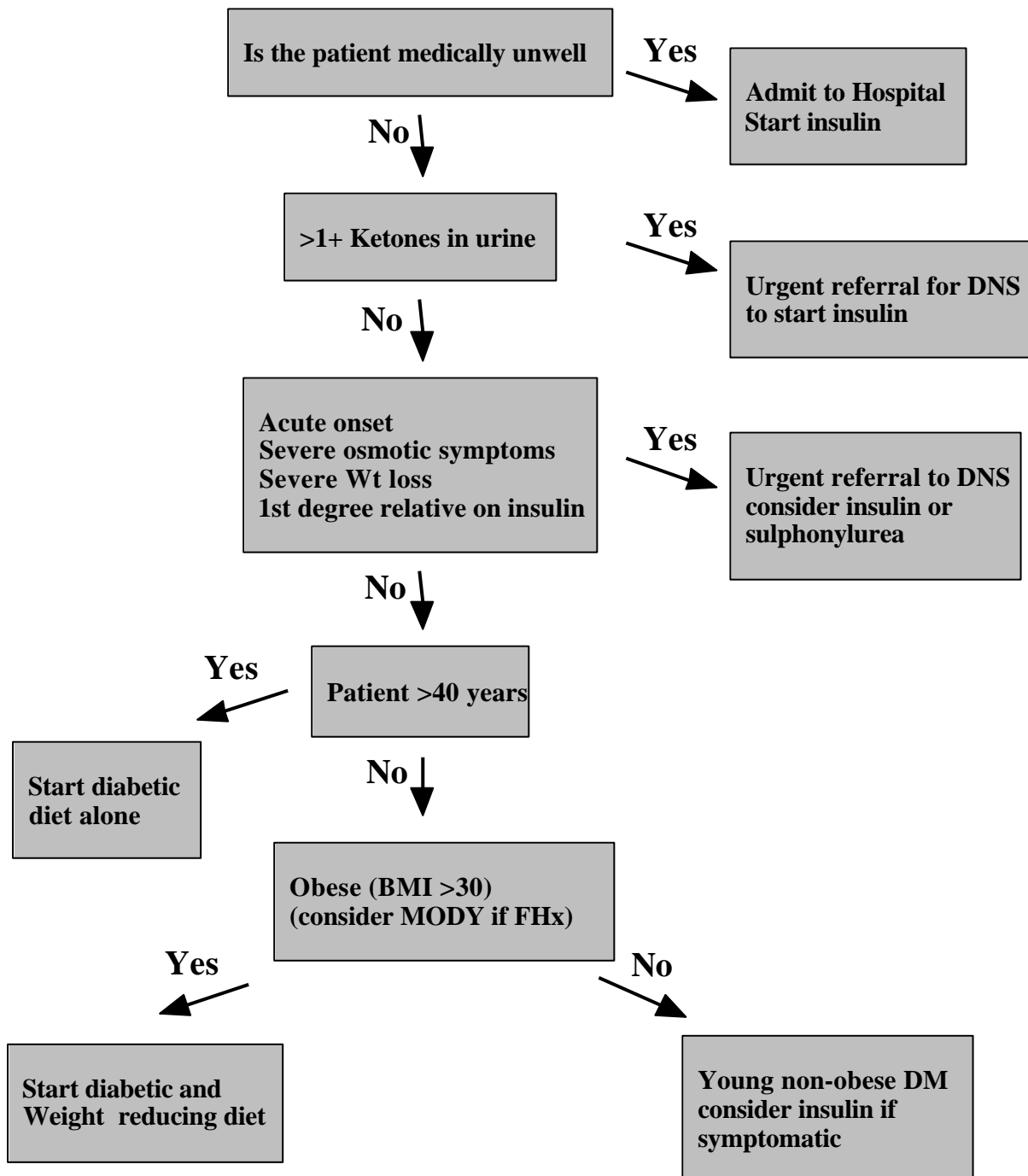
# Nurse Specialist Assessment

AIM : To enable the patient to understand and accept their diabetes, and to try and minimise the long-term complications.

Education is tailored to the patient's need topics include

- 1) Explanation of diabetes and complications
- 2) Management of diabetes
- 3) Learning skills: Blood glucose monitoring, urine testing, insulin injection
- 4) Discussion of pre conceived ideas and patients questions
- 5) Emergency contact number
- 6) Information on Diabetes UK
- 7) Institute initial management

# Initial Assessment of Treatment





# New Patient Medical Assessment

See Proforma 1

**RECORD** : Wt ; height ; random glucose ; urine dipstick and pin-hole acuity

## **History to include**

### **Patient's occupation and ethnic origin**

### **Record date of diagnosis and diagnostic criteria**

2x random glu =11.1 or 2x fasting glu =7.1 or OGTT 2h glu >11.1  
Presumed type of diabetes mellitus

### **Presentation**

Hospital admission with DKA or HONK  
Polyuria, polydipsia, visual disturbance, weight loss  
Infections : cutaneous, foot ,urinary tract, balanitis, pruritis vulvae

### **Record known previous complications**

IHD, PVD, CVD, retinopathy/cataracts, neuropathy, nephropathy, ulcers, impotence

### **Directly question regarding**

Exercise related chest pain  
Intermittent claudication  
Foot infections and symptoms of peripheral neuropathy  
Erectile dysfunction  
Visual disturbances

### **Assess present control**

HBGM results: hypo and hyperglycaemic episodes  
Dietary compliance, exercise regime  
Has the patient seen the diabetic nurse specialist and dieticians?

### **Obstetric history**

Number of pregnancies; miscarriages; stillbirths, birth weights and mode of delivery  
History of gestational diabetes, future pregnancy plans and contraception

### **Family history**

Diabetes mellitus, thyroid disease, hypertension, renal disease  
IHD, CVD, PVD with age of onset

### **Full list of medication and drug allergies**

### **Social history**

Smoking history, alcohol intake  
Driving (check informed DVLC and insurance company)  
Assess occupational risks (driving or machinery etc)

### **Physical Examination to include**

General, cardiovascular, respiratory and abdominal systems

### **Specifically record**

Evidence of hyperlipidaemia (arcus, tendon xanthomata etc.)  
Pulse, blood pressure lying and standing, heart sounds, carotid bruits  
Peripheral pulses: abdominal, femoral, dorsalis pedis and posterior tibials  
Reflexes at the knee and ankle  
Lower limb sensation; vibration, 10g microfilament, pinprick  
Feet : General skin, nails, ?fungal infection and deformity, ?Charcot's  
Visual acuity, presence of cataracts and dilated funduscopy

### **Investigations to include**

FBC  
Renal function, Liver function and GGT, Calcium, (Vitamin D if Asian origin)  
Thyroid function,  
Fasting Cholesterol, Triglycerides, HDL, LDL, Chol/HDL ratio  
Glucose and HBA1c  
Random urinary albumin /creatinine ratio  
ECG and CXR if indicated  
B<sub>12</sub>, folate and tissue specific autoantibody screen in Type I (including anti GAD and anti-islet cell)

### **Repeat fasting blood test 2 weeks before next clinic**

U&E, LFT, TFT, Chol, Tg ,HDL, LDL, Glu, HBA1c and (CK, GGT if on a statin/glitazone)

# New Patient Dietetic Assessment

## NEW DIABETIC PATIENT DIETETIC REVIEW



### St VINCENT DECLARATION

Set standards for care of people with diabetes  
“People with diabetes should have access to expert nutritional and dietary advice  
Should see state registered dietitian within 4 weeks of diagnosis and then annually  
National Service Framework for Diabetes  
Proposes a pathway of care where dietary management has a role from prevention to the management of complications



### INITIAL APPOINTMENT WITH SRD

Explanation of diabetes, risk of complications, role of diet in treatment  
Collection of baseline information co-morbidities  
Record parameters  
    Weight  
    Height  
    BMI  
    Waist circumference,  
    Glucose and Biochemistry  
    Drugs  
Review of diet diary, 24h diet history, activity levels  
Set and agree dietary, activity and weight targets  
Aimed at prevention or reduction in risk factors  
    Optimal diet  
    Lipid lowering  
    Weight loss  
    Improving glycaemic control  
    Blood pressure reduction  
    Improved nutritional status



### SET GOALS

Specify the dietary goals set  
    Encouragement  
    Motivation  
    Support  
Arrange follow up appointments as necessary

# Driving and Diabetes Mellitus

Adapted from DVLA Medical standards of fitness to drive *Feb 2002* (App.I)

All patients must inform the DVLA of diagnosis and will be sent detailed information

## **Insulin treated diabetes**

Car drivers: Must recognise warning signs of hypoglycaemia and meet visual standards

HGV/PSV: Applicants since 1/4/91 barred from driving.

Drivers licensed before 1/4/91 require annual consultant certificate.

Since April 2001 allow 'exceptional cases' to apply for or retain their entitlement to drive class C1 vehicles (3500kg – 7500kg) subject to an annual medical

## **Temporary insulin treatment** (Gestational DM or Post MI)

Car drivers: Notify DVLA. May drive but must stop if severe hypos.

Must notify DVLA if continues insulin treatment continues for > 3 months

HGV/PSV: May not drive, reapply for licence when not on insulin

## **Oral hypoglycaemics**

Car drivers: Retain licence until 70y unless visual acuity or fields affected

HGV/PSV: Licensed unless acuity or fields affected

## **Diet only**

Car Drivers: To notify DVLA if visual acuity or fields affected

HGV/PSV: Licensed unless visual acuity or fields affected

## **Diabetic Complications**

### **Frequent Hypoglycaemia or hypoglycaemic unawareness**

Car drivers: Stop driving until control back to normal (medical report to DVLA)

HGV/PSV: May not drive

### **Visual Problems**

Car drivers: Must be better than 6/9 to 6/12 corrected and fields >120° horizontal

HGV/PSV: Must be 6/9 or better in best eye (corrected)

Must be 6/12 or better in worst eye (corrected)

Uncorrected better than 3/60 in both

Normal visual fields

### **Renal Problems**

Car drivers: No restriction unless dizziness, fainting or cognitive problems

HGV/PSV: Individual assessment by DVLA

# The Annual Diabetic Review

See Appendix III

**RECORD :** Wt, Wt change; BMI ; random glucose ; urine dipstick result and acuity

## History Examination and Investigation

### **Record accurately**

- Present age, type of diabetes, age of onset/duration, complications
- Other clinical diagnoses
- Smoking status
- Full list of present medications

### **List Recent Results (from 2 weeks prior to clinic)**

HBA1c, Chol, Tg ,HDL, LDL, Chol/HDL, Cre, Alb/Cre ratio, FT4, TSH, LFT  
(GGT and CK if on a statin)

### **Assess present control**

- Record home blood glucose monitoring
- Record hypo and hyperglycaemic episodes
- Dietary compliance
- Level of regular exercise

### **Directly question regarding**

- Exercise related chest pain
- Intermittent claudication
- Foot infections
- Symptoms of peripheral neuropathy
- Visual disturbance
- Erectile dysfunction
- Pregnancy or plan for pregnancy and contraception
- Chiropody

### **Physical Examination**

#### **Specifically record**

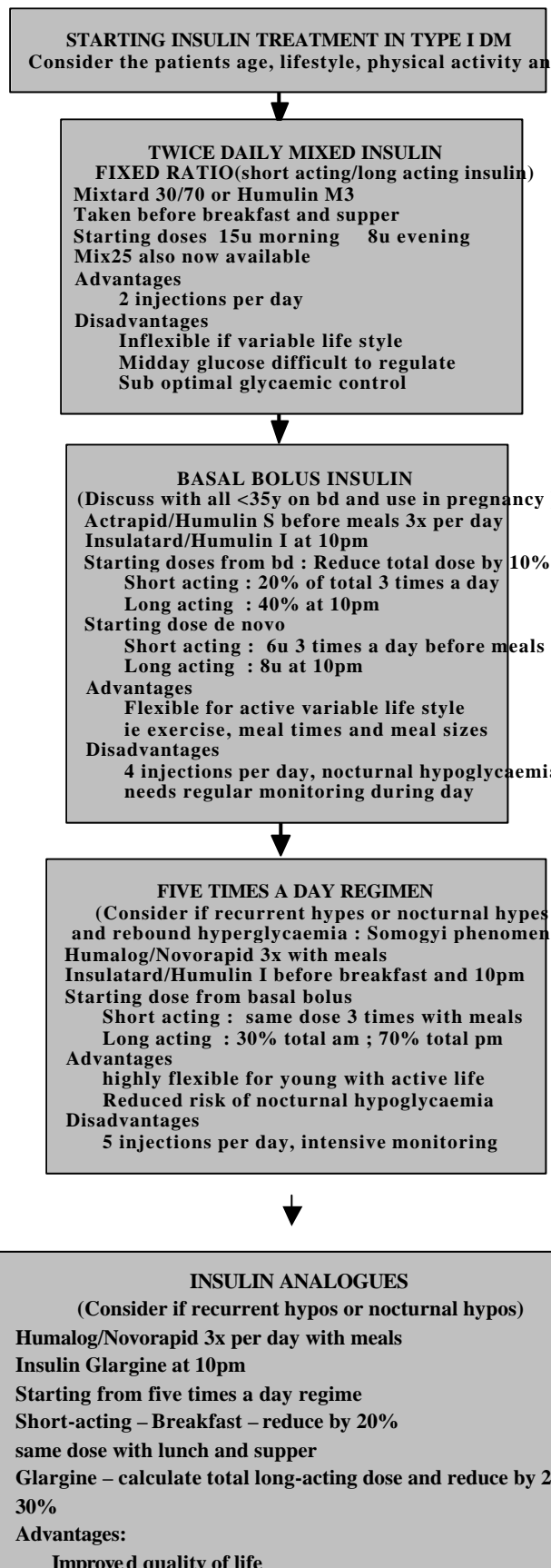
- Blood pressure lying and standing
- Peripheral pulses: dorsalis pedis and posterior tibials
- Reflexes at the knee and ankle
- Lower limb sensation : vibration, 10g microfilament and pinprick
- Feet : General skin, nails, fungal infections, deformity/Charcot's, Ulcers
- Visual acuity, presence of cataracts and dilated funduscopy

### **Repeat fasting blood test 2 weeks before next clinic**

U&E, LFT, TFT, Chol, Tg ,HDL, LDL, Glu, HBA1c and (CK, GGT if on a statin/glitazone)  
Random albumin/creatinine ratio

**Dietetics Review** at least every 2 years

# Insulin Treatment in Type I DM



# Commonly Used Human Insulins

Most insulin is used with 3ml re-loadable cartridge pens. 1.5ml cartridges are no longer available

Humapen and Novopen

Disposable pens are also available for ease of use

Special pens are available for partially sighted and elderly

Long acting

Insulin Glargine                      flat profile    duration 24h

Intermediate acting

Insulatard                              peak 5h        duration 10h

Humulin I                                peak 5h        duration 10h

Fast Acting (Taken 30 minutes before meals)

Actrapid                                peak 2h        duration 7h

Humulin S                                peak 2h        duration 7h

Very Fast Acting (Taken with or after start of meals)

Humalog (Lispro)                      peak 1h        duration 4h

Novorapid                                peak 1h        duration 4h

Mixtures of Human Insulin

Human Mixtard 30/70 (30% Actrapid : 70% Insulatard)

also available 10/90, 20/80, 40/60, 50/50

Humulin M3 (30% Humulin S and 70% Humulin I)

Also available M2, M5

Mix25 (25% Humalog : 75% Humulin I)

# INSULIN GLARGINE

**Insulin Glargine** is a long acting human insulin analogue prepared by modifying the structure of insulin to allow more consistent release during the day and thereby mimicking natural basal insulin release.

Glargine maintains a basal concentration of insulin in the blood which can then be increased by injections of short-acting insulin analogues as required. It is used as the basal component of a basal-bolus regime. The prolonged action of glargine, without pronounced peaks over 24 hours, make it ideal for this purpose.

Glargine does not require resuspension before use and this reduces intra and inter-user variability.

In type I diabetes, 3 studies have shown a significant improvement in fasting blood glucose. One study reported a significant improvement in HbA1c though the study was only 4 weeks duration. The manufacturers report a 1.7% reduction in HbA1c. One study reported a significant reduction in nocturnal hypoglycaemic episodes, though this was not supported by 2 other studies. The manufacturers report a 70% reduction in hypoglycaemic episodes.

In type II diabetes, glargine produces no change in fasting blood glucose or HbA1c according to 2 randomised controlled trials but the manufacturers report a 1.4% reduction in HbA1c. Both RCTs report a significant reduction in nocturnal hypoglycaemic episodes.

## **NICE guidelines**

Glargine is a treatment option for **all** patients with type I diabetes

Glargine should only be considered in type II diabetes under the following circumstances

- a) The individual requires assistance from a healthcare professional to administer insulin.
- b) The individual's lifestyle is severely restricted by recurrent symptomatic hypoglycaemic episodes.
- c) The individual would otherwise use twice daily basal insulin in combination with oral hypoglycaemic agents.

# Alternatives to s/c Insulin in Type I Diabetes

## BRIEF REVIEW OF ALTERNATIVE TREATMENT OF TYPE 1 DM



### INSULIN PUMP THERAPY (In UK 1:1000 Type I DM) Continuous Subcutaneous Insulin Infusion (CSII)

Pump driven infusion of soluble insulin subcutaneously via a cannula into the abdomen  
Continuous background dose and boluses before meals

#### Candidates for consideration of insulin pump

- Very motivated to take control of own DM and do 4 blood sugars every day
- Good understanding of Type 1 DM and effects of insulin, exercise and food
- Young type I DM unstable control (hypos/hypers) on 5 times a day basal bolus
- Young type I DM poor control and early reversible microvascular complications
- Poor control on basal bolus in pregnancy
- Sufficient financial resources or health authority funding (initial £2000 & annual £

#### Advantages

- Improved glycaemic control over basal bolus and quality of life
- Reduce risk of complications
- Help in restoration of hypoglycaemic warning symptoms

#### Disadvantages

- Motivated patients with good knowledge only
- Expensive initial equipment £2000 and then £600 per year paid by patient
- Experience required by health professionals
- Needs 24h help line
- Pump failure can lead to rapid DKA
- Infections at infusion site



### ISLET CELL TRANSPLANTATION

Islet cell transplantation in Type I DM is a research trial only at present  
UKITC c/o Research and Information Dept 10 Queen Anne Street London W1G 9LH  
UK Islet Cell Transplantation Consortium : planned facilities  
King's College London, Royal Free Hospital London  
Addenbrookes Hospital Cambridge, John Radcliffe Hospital Oxford  
Worcester Acute Hospital, University College Leicester, Southmead Hospital Bristol

#### Patient Selection Criteria

- Type I DM between 18 and 65 years
- Failure of control and severe hypoglycaemia on optimal 5x per/day regime
- early microvascular complications

#### Exclusion Criteria

- Renal Disease
- Insulin resistance
- repeated episodes DKA

#### EDMONTON PROTOCOL (Shapiro et al NEJM 2000 343 : 230-238)

##### Islets

- Isolated from human pancreas (11,000 islets per Kg of recipients weight )
- Cross matched for blood type and lymphocytotoxic ab but not HLA

##### Transplant

- Percutaneous transhepatic access to portal vein under sedation
- Islets infused over 5 min in 120ml of medium

##### Immunosuppression

- Sirolimus od, Tacrolimus bd and daclizumab iv 5x 2 weekly

##### Prophylaxis

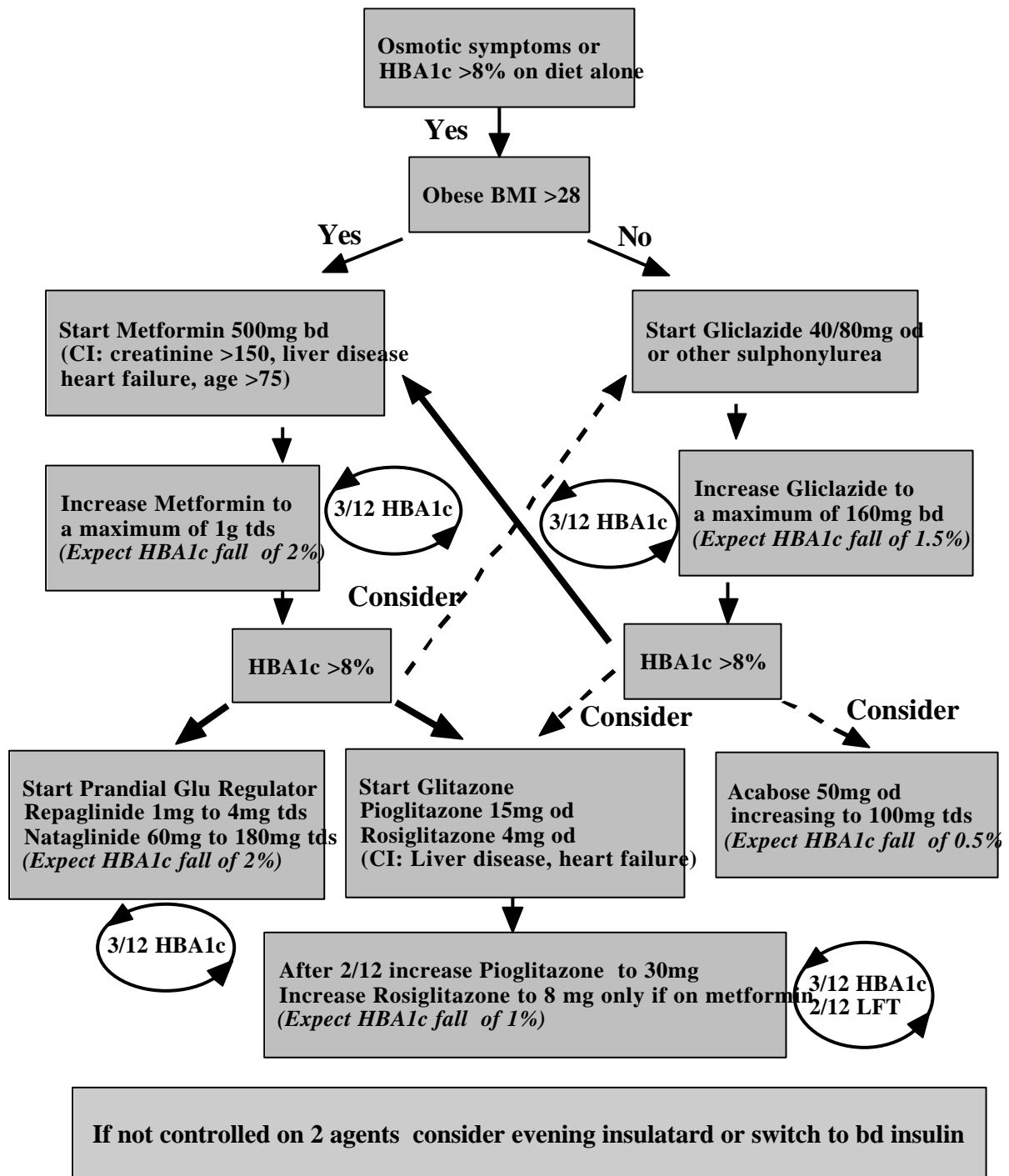
- IV vancomycin, Imipenem, VitE/B/A, inhaled Pentamidime monthly
- oral gancyclovir tds for 14 weeks post transplant

##### Results

- All patients required islets from at least 2 pancreases
- 7/7 had insulin independence and normal HBA1c at 1y FU



# Oral Hypoglycaemics Type II DM



# Oral Hypoglycaemic Agents

## **Sulphonylureas** (Expect reduction in HBA1c of 1 to 1.5%)

Action : Stimulate pancreatic insulin secretion

(Closes  $K^+$  channels opens  $Ca^{2+}$  channels resulting in incised insulin secretion)

Gliclazide initial dose 40mg od before meals titrate monthly to max of 160mg bd

Risk of hypoglycaemia and weight gain average 5Kg

Avoid long acting sulphonylureas in patients >70y,

Avoid sulphonylureas in severe renal impairment (Cre>350) and liver disease

Gliclazide (40mg od to 160mgbd) short acting

Tolbutamide (0.5g od to 1g bd) short acting low incidence of hypoglycaemia

Glimepiride (1mg od to 4mg od) long acting avoid in >70y

Glibenclamide (5mg od to 15mg od long acting avoid in >70y

## **Biguanides** (Expect reduction in HBA1c of 0.8 to 2%)

Action : Increases muscle glucose uptake and decreases hepatic gluconeogenesis

does not result in wt loss or gain

Metformin initial dose 500mg bd with meals titrate monthly to maximum 850mg tds

Risk of GI disturbance, lactic acidosis, renal impairment with radiographic contrast

Avoid in hospital inpatients, renal impairment (Cre>150), liver disease, heart failure and pregnancy

IV contrast: Do not take metformin 48h before or after IV contrast

## **Acarbose** (Expect reduction in HBA1c of 0.5%)

Actions :  $\alpha$ -glucosidase inhibitor delayed glucose absorption

Acabose initial dose 50mg od increasing slowly to 100mg tds with meals

Risk of GI disturbance frequent poor compliance

## **Prandial Glucose Regulators** (Expect reduction in HBA1c of 1% to 2%)

Actions : Rapid short action via the sulphonylurea receptor stimulating insulin release

Repaglinide 0.5 to 4mg 15 minutes before meals (max 16mg/d)

Nateglinide 60 to 180mg 15 minutes before meals (maximum 540mg/d)

Possible benefits: less wt gain and hypoglycaemia, can be used with metformin

Repaglinide may be used as monotherapy and both in renal impairment

Neither to be used with sulphonylureas or in pregnancy

Risks GI disturbance : avoid in severe hepatic or renal impairment and pregnancy

## **Thiazolidinediones** (Expect reduction in HBA1c of 1% )

Actions : PPAR $\gamma$  nuclear receptor activator (peripheral insulin sensitisation)

Indicated for use in inadequately controlled diabetics on

- sulphonylurea but intolerant of metformin
- obese patients on metformin.

Not licensed currently for use as monotherapy or with insulin

Pioglitazone 15mg od increasing to 30mg od

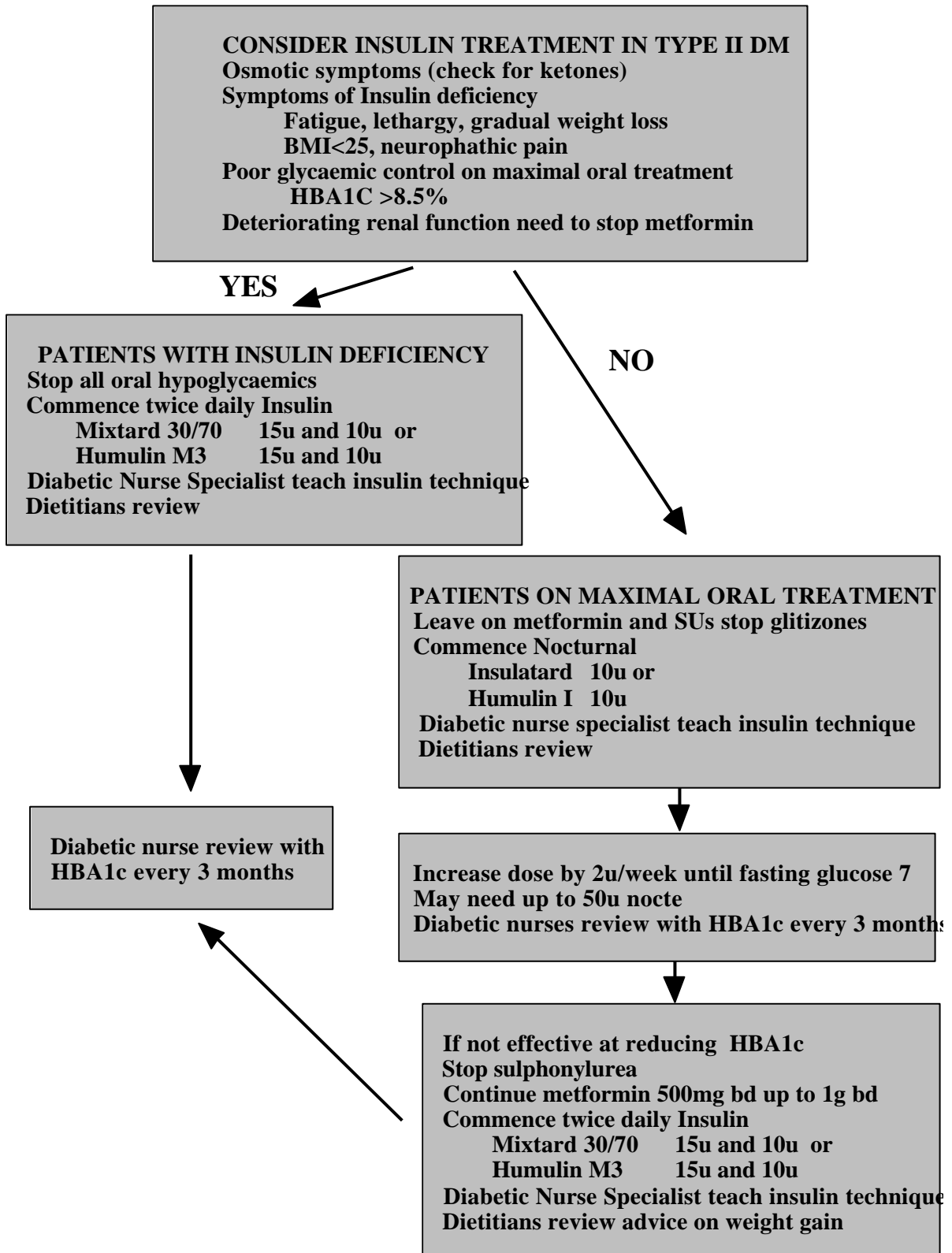
Rosiglitazone 4mg od increasing to 8mg od after minimum of 2 months (only with metformin)

Risks GI disturbance, wt gain and oedema

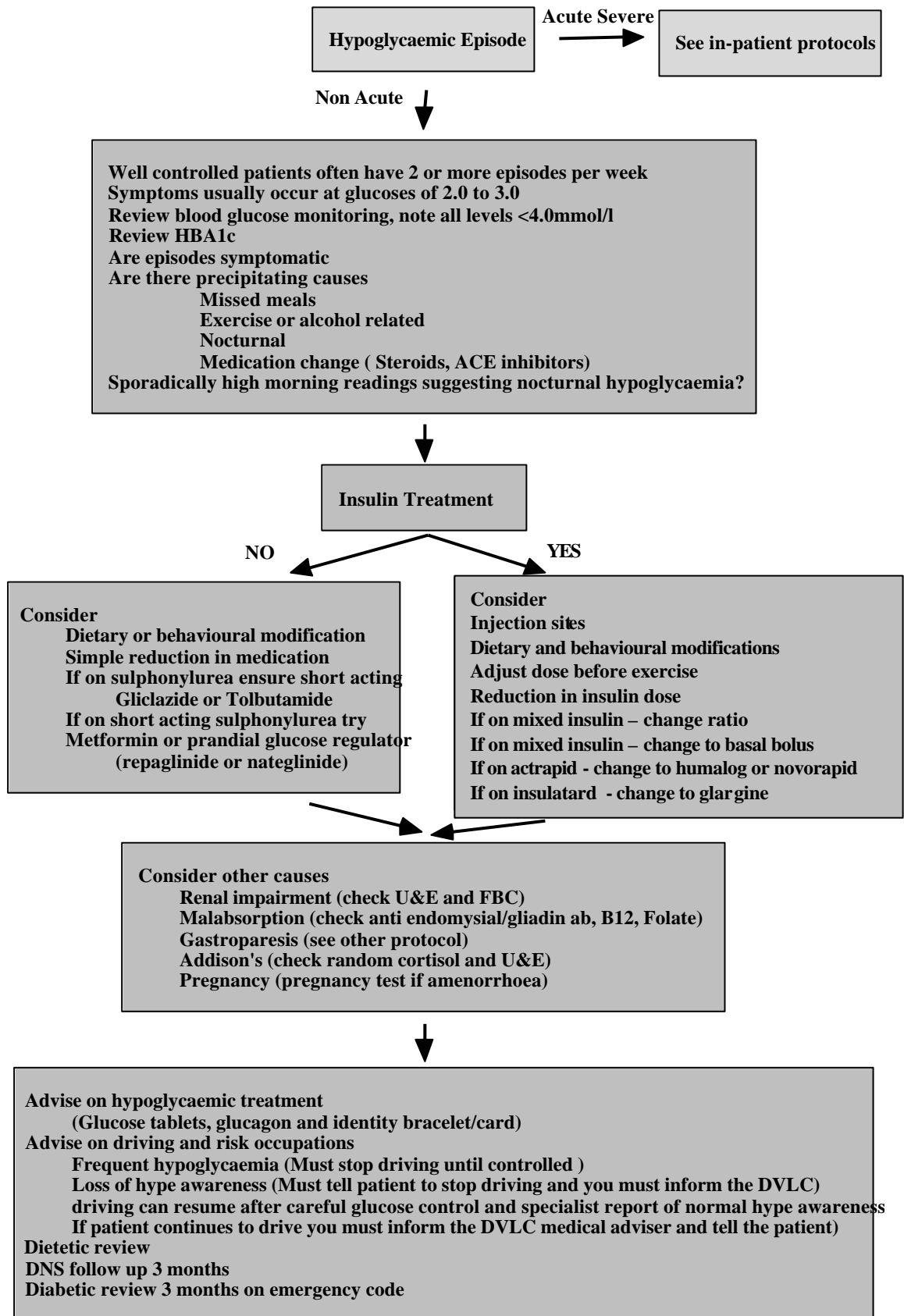
Avoid in hepatic impairment, heart failure and pregnancy

Monitor liver function at baseline and every 2 months for first year (stop if ALT 3x normal upper limit)

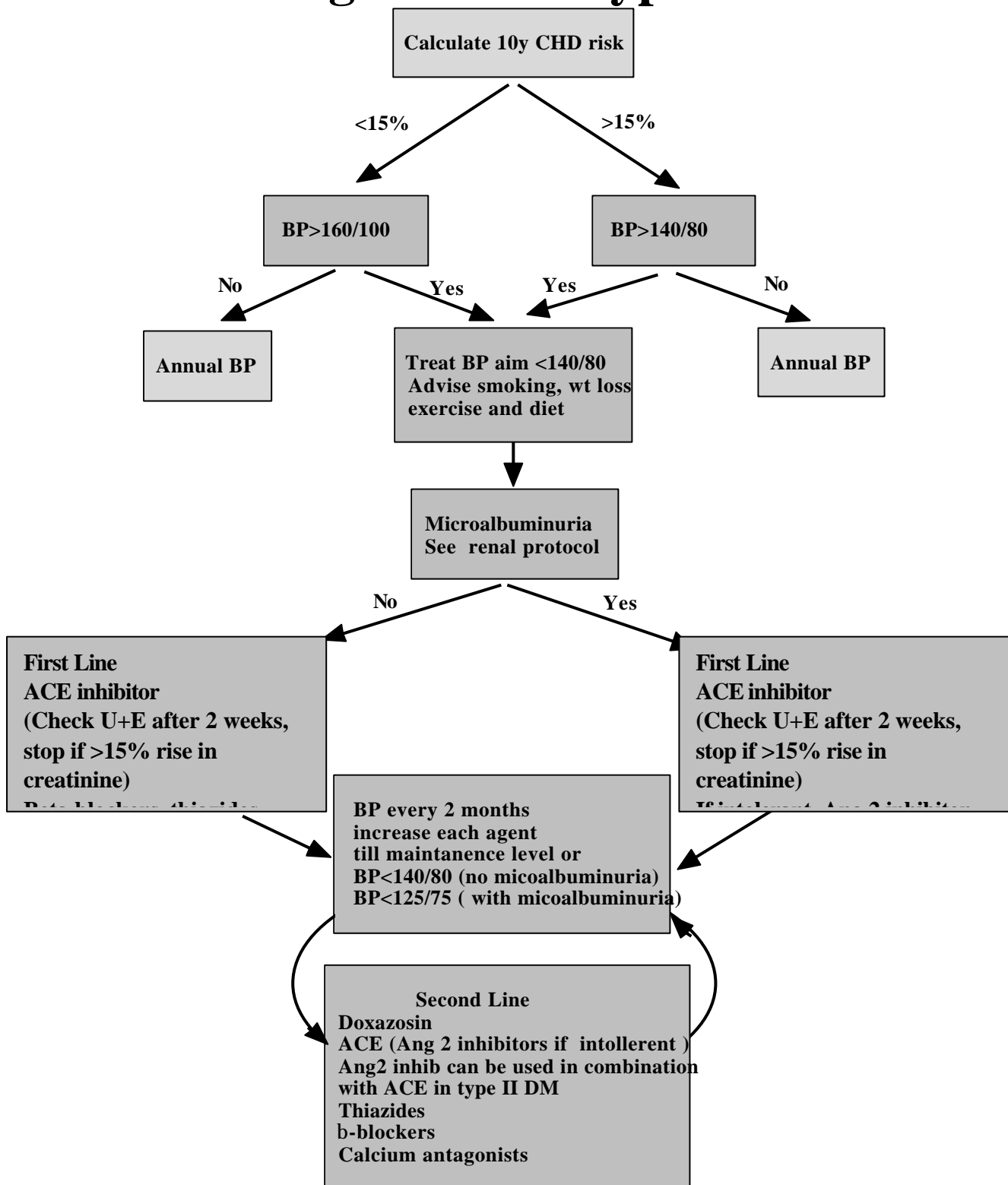
# Insulin Treatment in Type II Diabetes



# Recurrent Hypoglycaemia



# Management of Hypertension



## Guidelines for starting ACEI or Ang2 inhibitors:

- Check U+E after 1-2 weeks – stop if Cr rises by > 15%
- Stop thiazides whilst initiating treatment to avoid volume depletion
- Discontinue NSAIDs
- If significant RVD, perform MRA to exclude renal artery stenosis

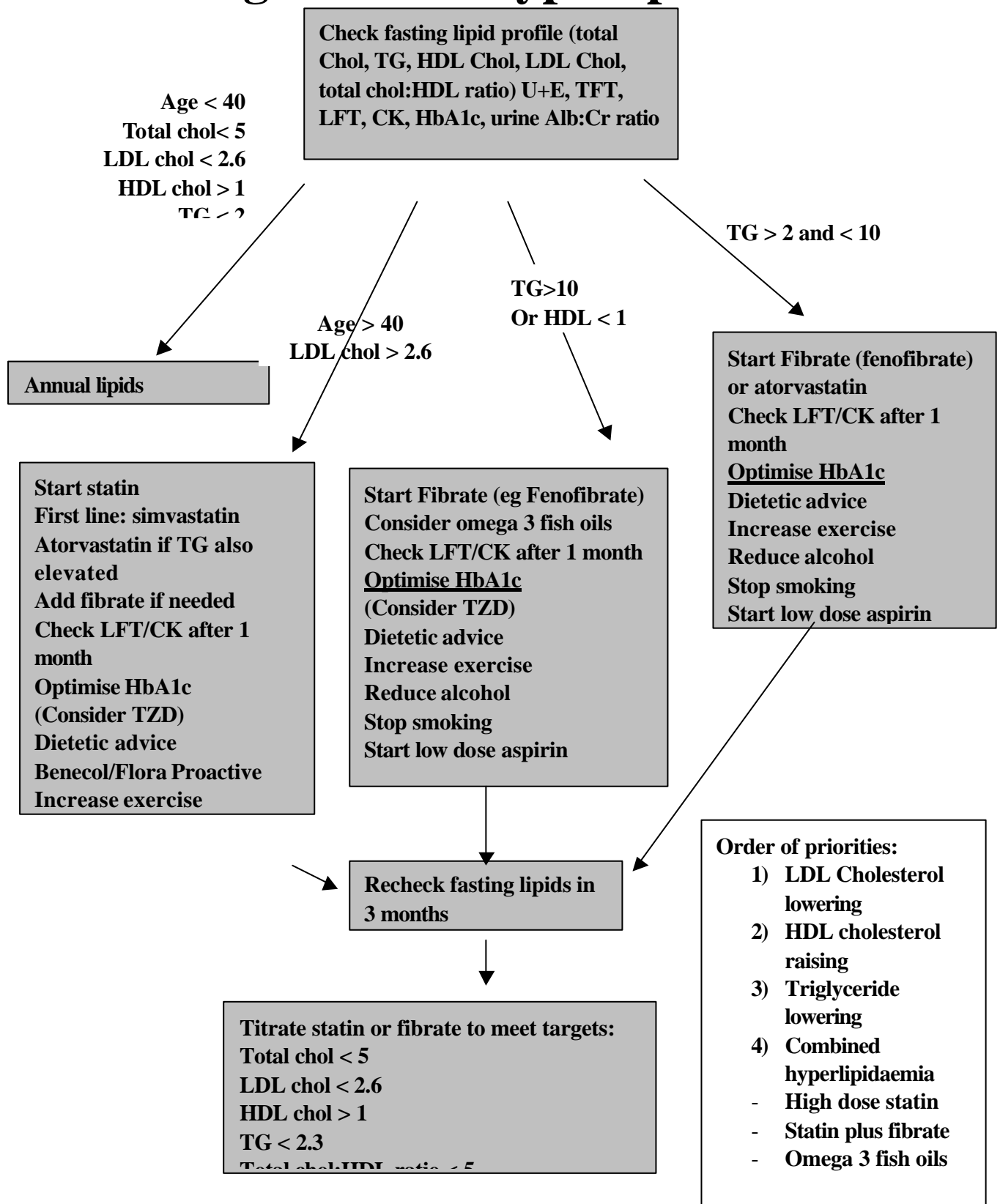
# Hypertension Data

- Hypertension occurs more frequently in patients with diabetes than in the general population. 40% of newly diagnosed subjects enrolled in the UKPDS had were hypertension and this increased during the study to being 48% of all men and 54% of all women
- In type 1 diabetes the incidence of hypertension increases with increasing duration of diabetes and frequently has a renal origin
- In type 2 diabetes hypertension is frequently part of the insulin resistance syndrome occurring with other cardiovascular risk factors.
- Hypertension is significant contributor to micro and macrovascular disease in both type 1 and type 2 diabetes
- In the type 2 diabetic subjects enrolled in the UKPDS for each 10 mm Hg decrease in mean systolic blood pressure there was associated with reductions in risk of 12% for any complication related to diabetes
- The third working party of the British Hypertension Society based the decision to treat hypertension in non-diabetics subjects with a sustained systolic BP of 140 - 159 mm Hg or sustained diastolic BP of 90-99 mm Hg on the presence of end organ damage, cardiovascular disease or a 10-year coronary heart disease risk of =15%. As the majority of diabetic patients will have a coronary heart disease risk of =15% and microvascular and macrovascular complications increase with increasing blood pressure the report recommended initiating antihypertensive drug therapy in diabetic subjects with a sustained systolic BP =140 mm Hg or sustained diastolic BP is =90 mm.
- Angiotensin-converting enzyme (ACE) inhibitors are the first line hypertensive agent for patients with type 1 diabetes as they has proven renal protective properties. All diabetic subjects with micoalbuminures should receive an ACE-inhibitor or an A-II receptor antagonists if intolerant to the former.
- A-II receptor antagonists provide some theoretical advantages over ACE inhibitors in that they directly inhibit A-II by binding to the AT(1) receptor subtype thereby blocking all A-II action at this receptor even when generated through the non classical renin-angiotensin pathways. A-II receptor antagonists have also been shown to significantly reduce the progression of diabetic renal disease in patients with type 2 diabetes.
- Many hypertensive diabetic patients require three of more antihypertensive agents to control blood pressure. Thiazides and  $\beta$ -blockers remain effective and useful first treatment options given alone or in combination. In the UKPDS atenolol was equally effective as the ACE-inhibitor captopril however was significantly cheaper.
- Doxacacin and long acting dihydropyridine calcium antagonists are effective second line agents ideally suited for combination with ACE-inhibitors, Thiazides and  $\beta$ -blockers if blood pressure target are not met.

## Key References

- Association of systolic blood pressure with macrovascular and microvascular complications of type 2 diabetes (UKPDS 36): Adler AI. Stratton IM. Neil HA. Yudkin JS. Matthews DR. Cull CA. Wright AD. Turner RC. Holman RR. BMJ. 321(7258):412-9, 2000
- Guidelines for management of hypertension: report of the third working party of the British Hypertension Society. Ramsay L. Williams B. Johnston G. MacGregor G. Poston L. Potter J. Poulter N. Russell G. Journal of Human Hypertension. 13(9):569-92, 1999

# Management of Hyperlipidaemia



**Stop statin or fibrate if transaminases over 3 x upper limit of normal or CK greater than 1000**

## Hyperlipidaemia Data

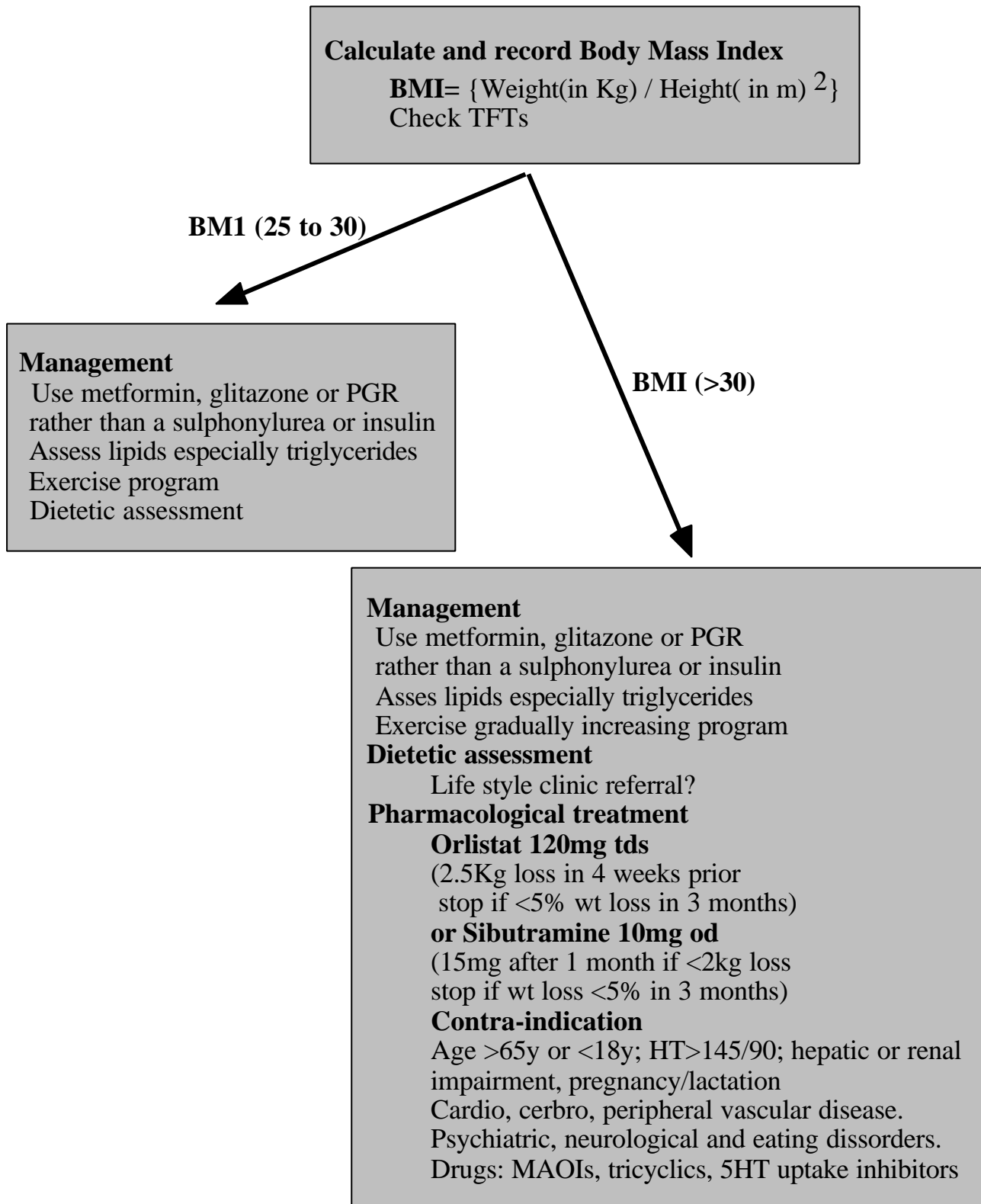
- All diabetic patients should receive dietary advice on reducing their dietary fat intake, replacing saturated fat with monounsaturated rich fats and oils and on the use of low fat spreads and fat substitutes, ie the use of Benecol and other plant stanols and sterols. Patients should be made aware of how to lower their CVD risk through life style changes.
- Total cholesterol and serum triglycerides in the newly diagnosed patients enrolled in the UKPDS were significantly higher than for the general population. Although the HDL concentration in the diabetic women in the UKPDS were higher than the men by 7% this sex differential was considerably less than for the general population in which it was 22%.
- The Multiple Risk Factor Intervention Trial (MRFIT) demonstrated for or any given level of cholesterol, the incidence of coronary artery disease in diabetes is increased 2-4 times.
- The Scandinavian Simvastatin Survival Study (4S) and Cholesterol and Recurrent Events Trial (CARE) were secondary intervention trials using a statins (simvastatin, and pravastatin) that performed post hoc analyses of the diabetic subgroups. Despite the limitations of such subgroup analyses, these studies strongly suggest that treating hypercholesterolemia in diabetes will reduce the risk of recurrent cardiac events in individuals with pre-existing CAD.
- West of Scotland Coronary Prevention Study Group.(WOSCOP) showed evidence of a statin (pravastatin) in the primary prevention of CVD specific studies on the use of statins for the primary prevention of CVD in diabetics are awaited.
- Statins also provide protection against stroke and peripheral vascular disease
- In diabetic patients with a low HDL cholesterol and low or normal LDL concentrations fibrates offer good secondary protection against further CVD and stroke. In the Department of Veterans Affairs Intervention Trial (VA-HIT) subjects with established CVD and low HDL cholesterol but normal LDL concentrations were randomized gemfibrozil (1,200 mg/day) or placebo for 5 years. Of the 627 diabetics (25% of the study population) gemfibrozil resulted in a 24% relative risk reduction in CVD end points (CHD death, nonfatal myocardial infarction, and definite stroke) compared with placebo.
- Fibrates are first line management in diabetic patients with significant hypertriglyceridaemia. A meta-analysis of 17 population-based studies ( 46 000 men and > 10 000 women) showed that risk of CVD increased by ~ 30% in men and by ~ 75% in women for every 1 mmol/l increase in triglycerides. A recent trial on progression of coronary atherosclerosis in type 2 diabetes over 3 years assessed by angiographic criteria showed in a placebo controlled trial micronised fenofibrate (200 mg/day) a showed a significantly smaller increase in percentage diameter stenosis with the fibrate than the placebo group.

## Key References

- Rubins HB, Robins SJ, Collins D. The Veterans Affairs High-Density Lipoprotein Intervention Trial: baseline characteristics of normocholesterolemic men with coronary artery disease and low levels of high-density lipoprotein cholesterol. *Am J Cardiol* 1996;78:572-575.
- Hokanson JE, Austin MA. Plasma triglyceride level is a risk factor for cardiovascular disease independent of high-density lipoprotein cholesterol level: a meta-analysis of population-based prospective studies. *Journal of Cardiovascular Risk* 1996; 3:213-219.
- U.K. Prospective Diabetes Study Group: U.K. Prospective Diabetes Study 27: plasma lipids and lipoproteins at diagnosis of NIDDM by age and sex. *Diabetes Care* 20:1683-1687, 1997
- Scandinavian Simvastatin Survival Study Group: Randomized trial of cholesterol lowering in 4,444 patients with coronary heart disease: the Scandinavian Simvastatin Survival Study (4S). *Lancet* 344:1383-1389, 1994



# Obese Diabetic Patients



# Proteinuria and Microalbuminuria

Refer all patients with diabetes AND increasing microalbuminuria or proteinuria and/or renal impairment, at least for initial assessment, or any patients in whom the diagnosis of urine or renal abnormalities is unclear

Early morning Alb:Cr ratio >2



Repeat in clinic  
Check MSU

Type I  
Alb:Cr ratio >2

Type II  
Alb:Cr ratio >5



24hour urine for protein, serum albumin  
Microalbuminuria = 30-300mg/L

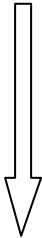
**Progression of renal failure in diabetes is preventable and requires impeccable BP control and reduction of proteinuria**

**STOP SMOKING  
REGULAR EXERCISE  
AVOID WEGHT GAIN**

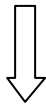
**DIABETES  
Without  
MICROALBUMINURIA**



**DIABETES  
with**



**DIABETES  
with PROTEINURIA  
(at least positive on 200mg)**



**DIABETES  
with RENAL IMPAIRMENT  
(creatinine > 150 micromol/l)**

**ASPIRIN,  
STATIN if LDL Chol >2.6  
MAINTAIN HbA1c < 8%  
BP < 140/80 mmHg  
6 MONTHLY URINE  
ALBUMIN:CREATININ  
E RATIO**

**AIIRA or ACE  
INHIBITOR in all  
BP < 130/80 mmHg  
6 MONTHLY URINE  
ALBUMIN:CREATININ  
E RATIO  
Titrate ACEI/AIIRA to**

**AIIRA or ACE  
INHIBITOR in all  
BP < 125/75 mmHg  
Ensure under regular**

**Antihypertensive Therapy**

- **Angiotensin receptor antagonists (AIIRA) or ACE inhibitors** should be used in all patients with microalbuminuria or proteinuria to reduce rate of progression to renal impairment, and non-renal vascular morbidity
- AIIRA may be better than ACEI
- Monitor serum creatinine after initiation (at 2 weeks) – creatinine will rise in MOST patients – stop ACEI/AIIRA ONLY if creatinine increases >15% above baseline
- Add diuretic (thiazide or indapamide) or calcium channel blockers (use once daily preparations)
- Beta and Alpha blockers useful
- Patients usually need multiple drugs

# Proteinuria and Microalbuminuria

- Diabetic nephropathy is the leading cause (25-44%) of end-stage renal failure in Europe, the United States and Japan. Microalbuminuria precedes the development of proteinuria and progressive renal impairment. The progression is of microalbuminuria to persistent proteinuria and renal disease is highly dependent on blood pressure, glycaemic control and duration of diabetes. In type 1 diabetes there is also a genetic susceptibility to diabetic. The proportion of renal replacement patients with diabetes is expected to double within the next 15 years, due to the increased prevalence of type 2 diabetes, the younger age of onset of diabetes and patient living longer.
- Microalbuminuria is associated with an increased risk of diabetic the other microvascular complications, namely retinopathy and neuropathy, it is also associated with a fourfold increased risk of macrovascular disease: disease in type II DM.
- Primary prevention of diabetic nephropathy is improved glycaemic control and blood pressure lowering. In the Diabetes Control and Complication Trial, showed intensified glycaemic therapy in type 1 diabetic patients reduced the occurrence of microalbuminuria by 39%, and that of albuminuria by 54%. In the UKPDS the impact of blood pressure control was earlier and more dramatic than the effect of glycaemic control on the development and progression of microalbuminuria.
- Early ACE inhibition, even when blood pressure is completely normal in type 1 diabetic patients with microalbuminuria can normalise albumin excretion rates and reduce the progression of microalbuminuria to persistent proteinuria.
- Angiotensin II type I receptor antagonists in type 2 hypertensive diabetics indicate that these agents also reduce microalbumin excretion rates, an effect that can be further increased with the addition of an ACE inhibitor.

## Key References

- EUrodiab Controlled trial of Lisinopril in Insulin dependent Diabetes mellitusThe EUCLID Study Group. Randomised placebo-controlled trial of lisinopril in normotensive patients with insulin -dependent diabetes and normoalbuminuria or microalbuminuria. *Lancet*,**349**:1787-92, 1997
- Brenner BM. Cooper ME. de Zeeuw D. Keane WF. Mitch WE. Parving HH. Remuzzi G. Snapinn SM. Zhang Z. Shahinfar S. RENAAL Study Investigators. Effects of losartan on renal and cardiovascular outcomes in patients with type 2 diabetes and nephropathy. *New England Journal of Medicine*. 345:861-9, 2001
- Mogensen CE, Neldam S, Tikkanen I, *et al*. Randomised controlled trial of dual blockade of the renin-angiotensin system in hypertensive microalbuminuric, non-insulin dependent diabetes: the candesartan and lisinopril microalbuminuria (CALM) study. *BMJ*. 321(7274):1440-4, 2000

# Blood Pressure in Renal Replacement

**Blood pressure in renal; dialysis patients**  
**Haemodialysis aim for <140/90 pre dialysis**  
Calcium channel blockers can be used daily  
No ACE or b-blockers or a-blockers on dialysis days  
**Peritoneal Dialysis aim for <140/90 pre dialysis**  
All antihypertensives may be used

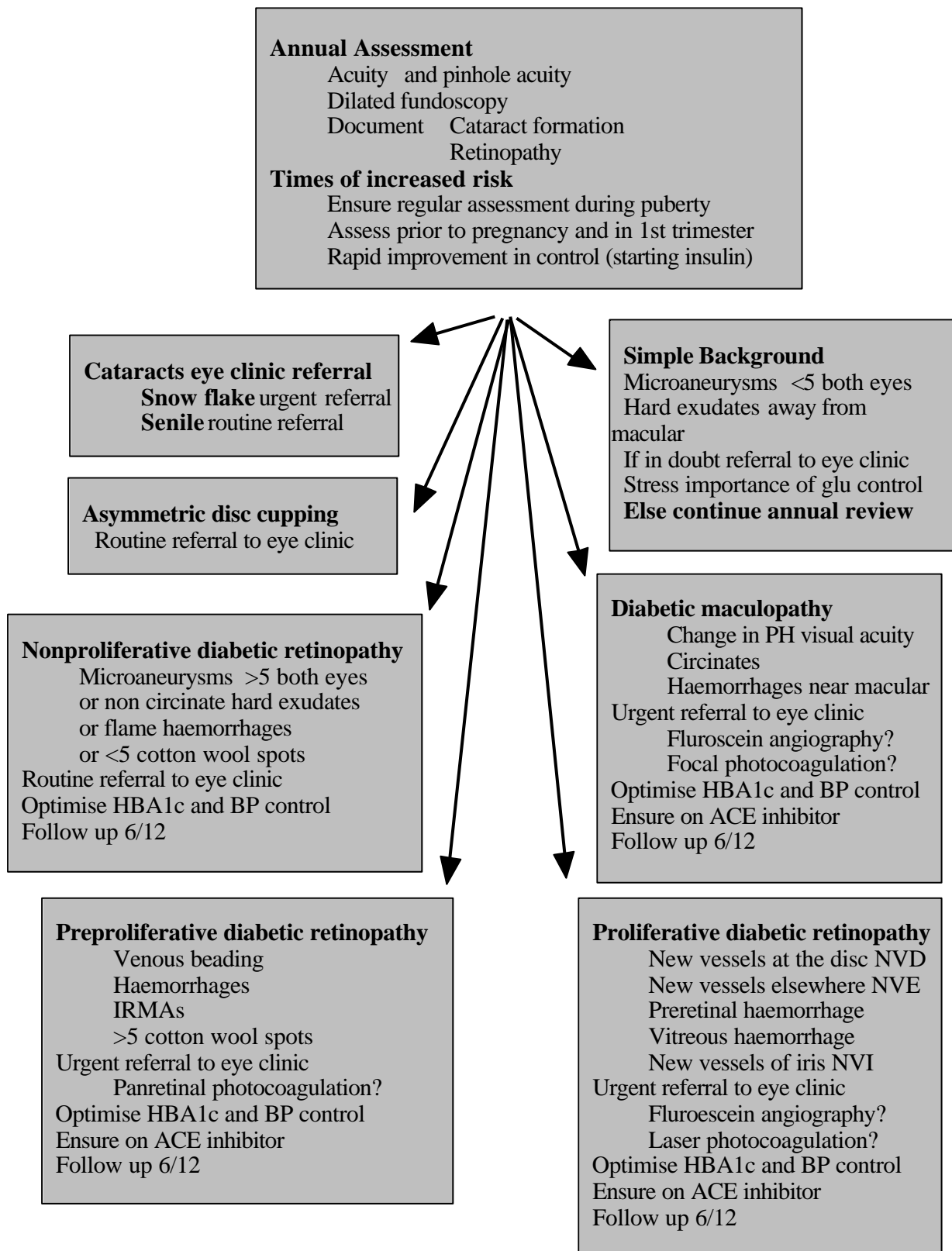
**Blood pressure in renal transplant patients**  
Calcium channel blockers can interact with  
(cyclosporin,tacrolimus,rapamicin)  
Amlodipine can be used safely however  
If use ACE inhibitors reduce diuretics and check U&E at 1 week  
>20% rise in K<sup>=</sup> or creatinine stop ACE

# Lipids in Renal Replacement

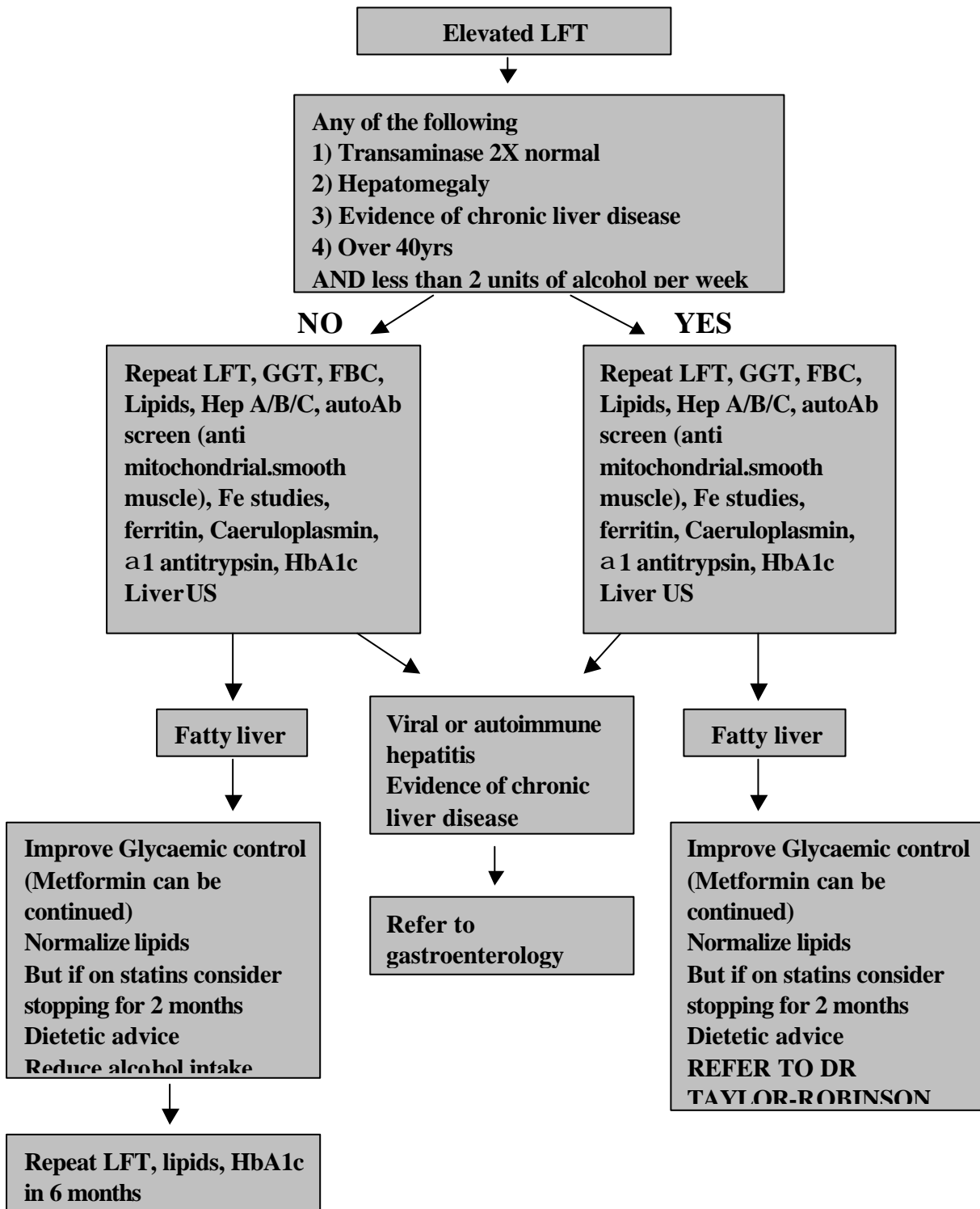
**Lipids in renal dialysis patients**  
Aim for Cholesterol <5 and LDL <3  
All current statins may be used  
Fibrates should only be used at low dose (Gemfibrozil)  
Statin and Fibrate together should not be used  
unless discuss with renal team

**Lipids in renal transplant patients**  
Aim for Cholesterol <5 and LDL <3  
Consider interactions with cyclosporin, tacrolimus and rapamicin  
Use Pravastatin and Fluvastatin (different elimination to cyclosporin)  
Statin and Fibrate together should not be used

# Diabetic Retinopathy



# Steatosis in Diabetic Patients



# Peripheral Neuropathy

**History**  
 Paraesthesia, hyperaesthesia, allodynia  
 Numbness and contact sensitivity  
 Burning pain in feet, worse at night,  
 Poor sleep and restless legs  
 (May be precipitated by poor control or  
 rapid improvement in control)

**Examination**  
 Evidence of motor/sensory/autonomic neuropathy?  
 Ascent reflexes, anhydrosis  
 Warm hairless legs frequently with strong pulses  
 Glove stocking neuropathy  
 Reduced LT, PP, VS, JPS and Temp sensation  
 Assess 10g microfilament sensation

**Investigations**  
 HBA1c, U&E, LFTs, Igs and PEP  
 FBC, ESR and B12, Intrinsic factor ab  
 Syphilis serology  
 Consider Nerve Conduction Studies?

**Initial Management**  
 Educate about foot problems  
 Formal podiatry assessment  
 Doppler studies if no palpable pulse  
 Regular 3/12 chiropody

**Mild Symptoms**

**Severe Symptoms**

**Management**  
 Optimise glycaemic control  
 HBA1c <7.5  
 Simple advice tights under cloths  
**Simple analgesia**  
 (aspirin/paracetamol/coproximol)

**Reassurance**  
 Severe pain usually improves in <2y  
 Regular 6/12 follow up in diabetic clinic  
 Optimise glucose control (HBA1c <7.5%)  
 Low threshold to switch to insulin

**Physical methods**  
 Bed cradle (to remove covers from feet)  
 Semipermeable plastic foot wrap (Opsite)  
 TENS

**Pharmacological**  
 Gabapentin (300mg od, 300mg bd, 300mg tds  
 increase to maximum of 600mg tds)  
 Amitriptyline (25mg nocte to 100mg)  
 Carbamazepine (100mg od to 600-800mg/d)  
 Capsaicin ( 4x/day releases substance P)

**Referral to Professor Anand Hammersmith**

# Diabetic Amyotrophy

## History

Motor Neuropathy (lumber-sacral plexus)  
Frequently male, Type II DM, 50-60y,  
Gradual onset  
Thigh weakness, severe pain and wt loss  
Anorexic



## Investigations

U&E, Calcium, LFTs, CRP  
FBC, ESR  
HbA1c  
PSA  
Xray Chest, Thoracolumber spine  
Nerve Conduction Studies



## Management

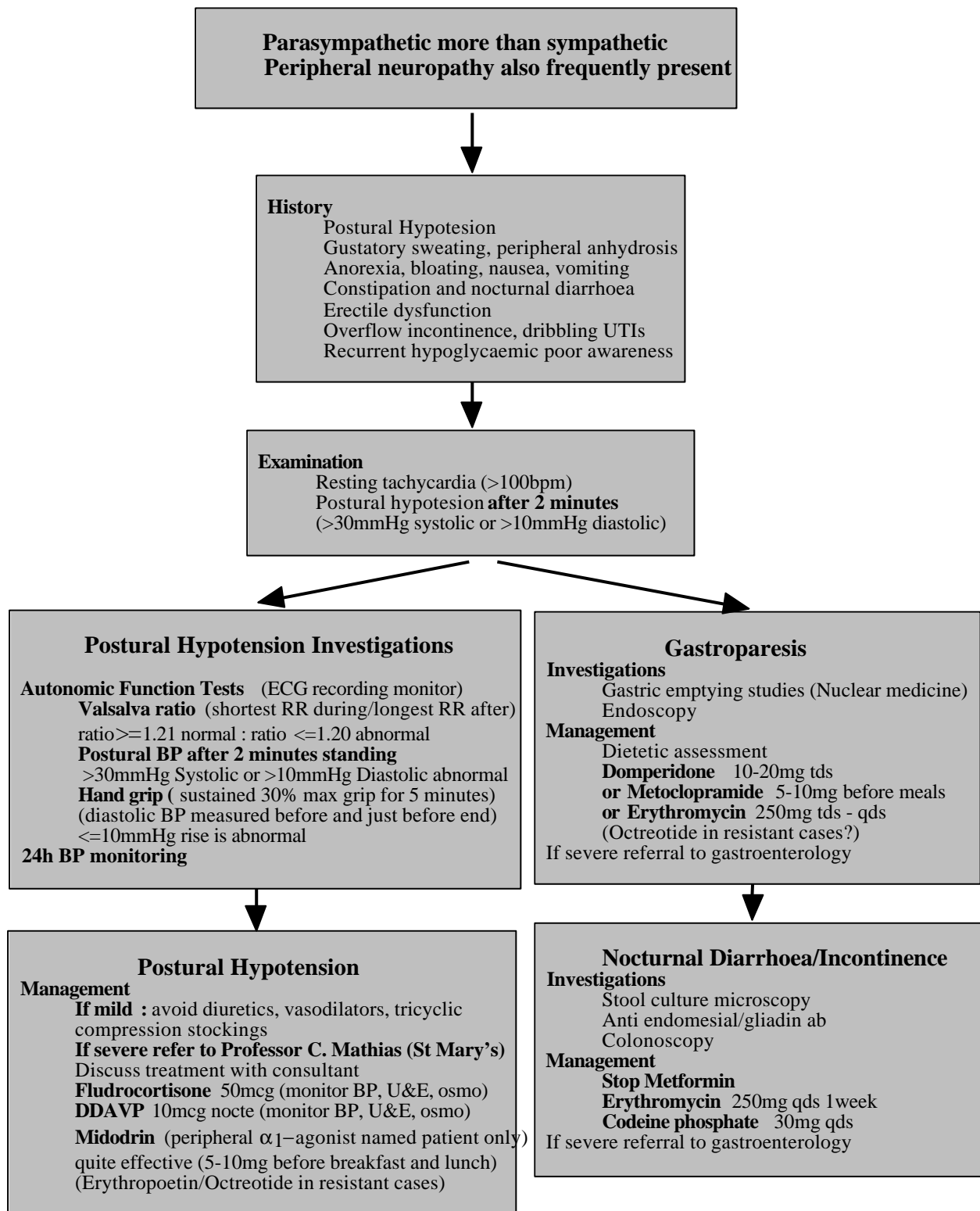
Admit to hospital  
Switch to qds insulin  
Dietetic assessment  
Optimise HbA1c  
Analgesia (may need opiates, tricyclics )  
Physiotherapy  
IV steroids used in trials



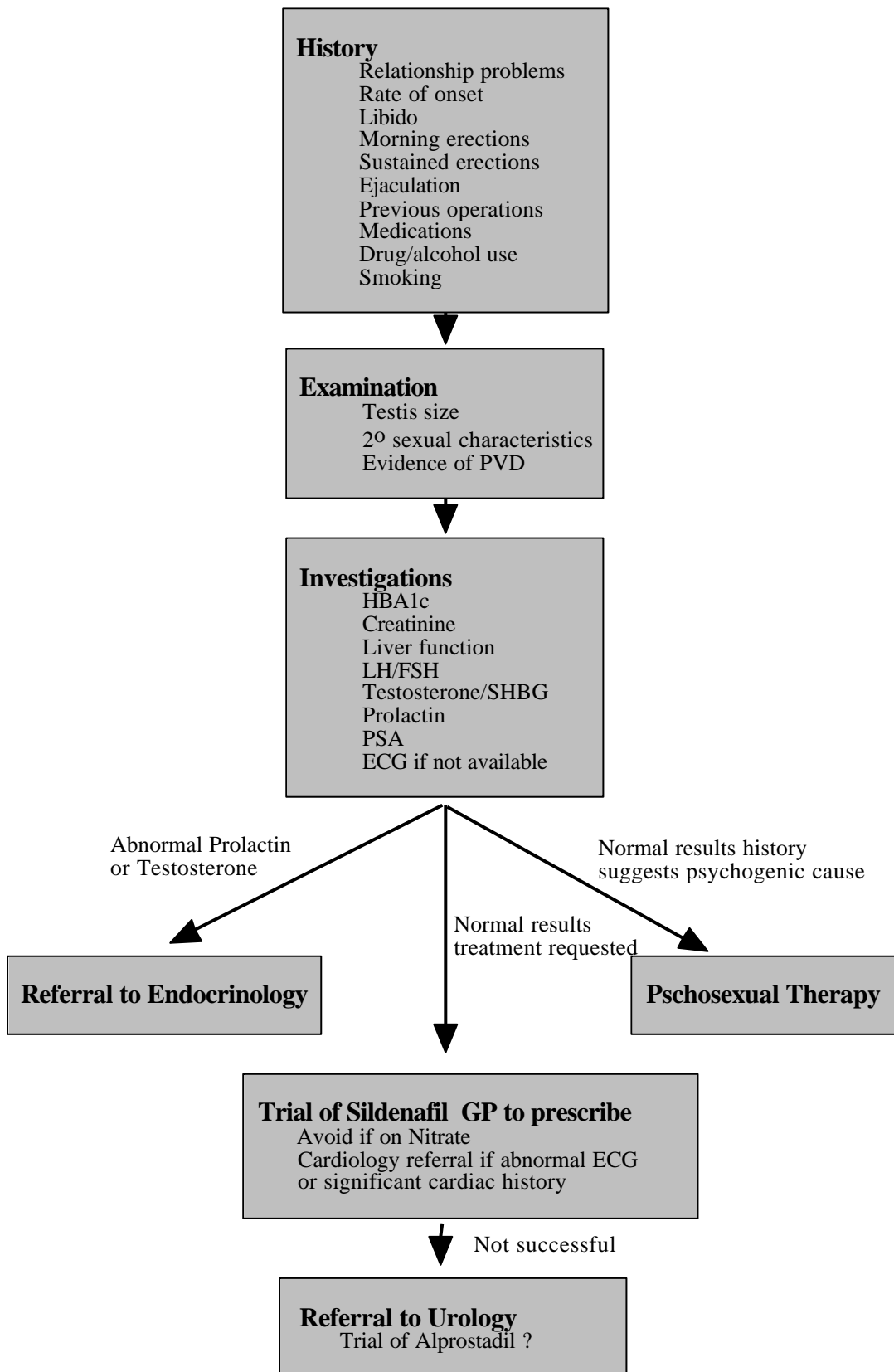
**Expect gradual improvement over 3/12**



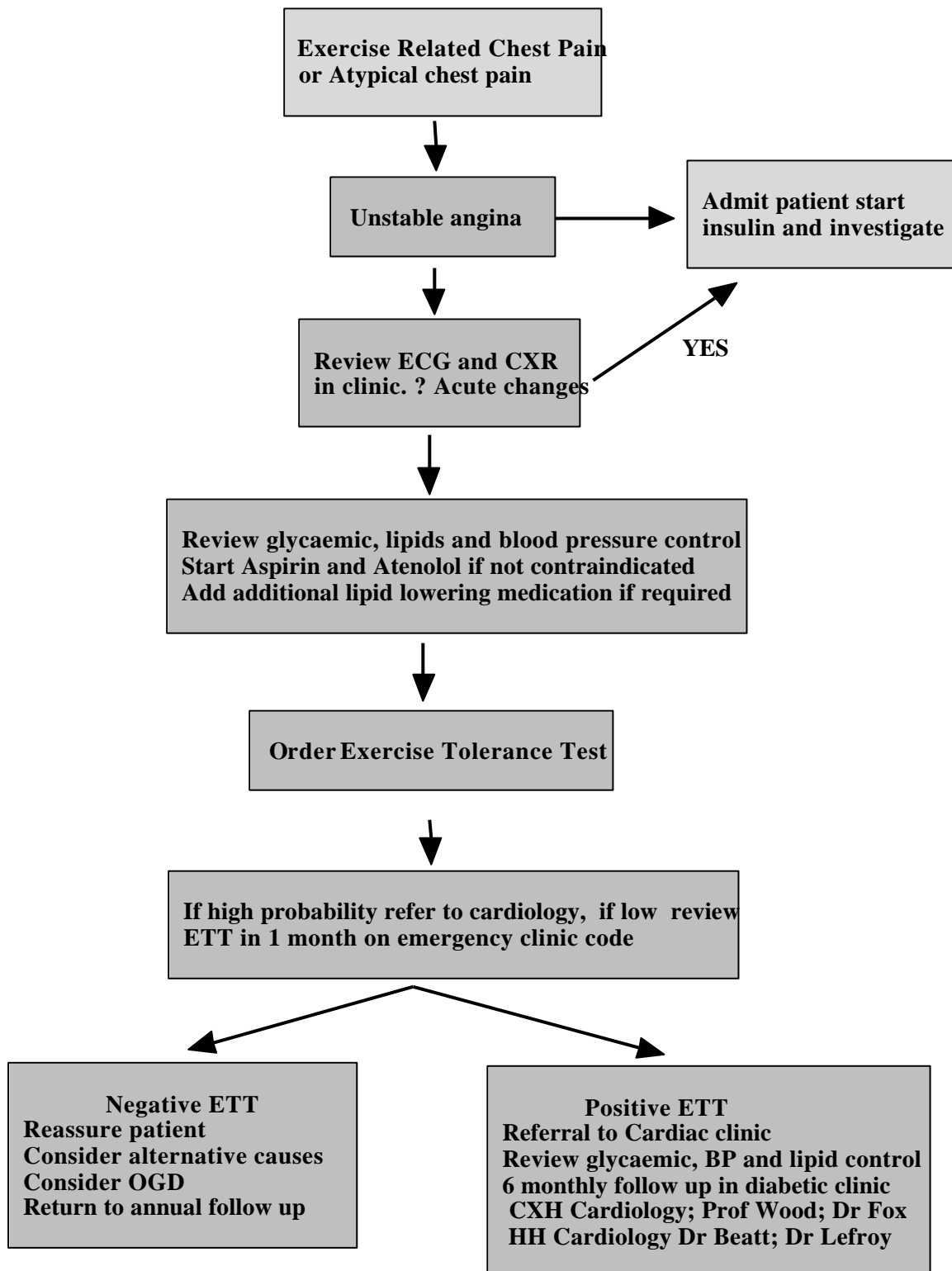
# Autonomic Neuropathy



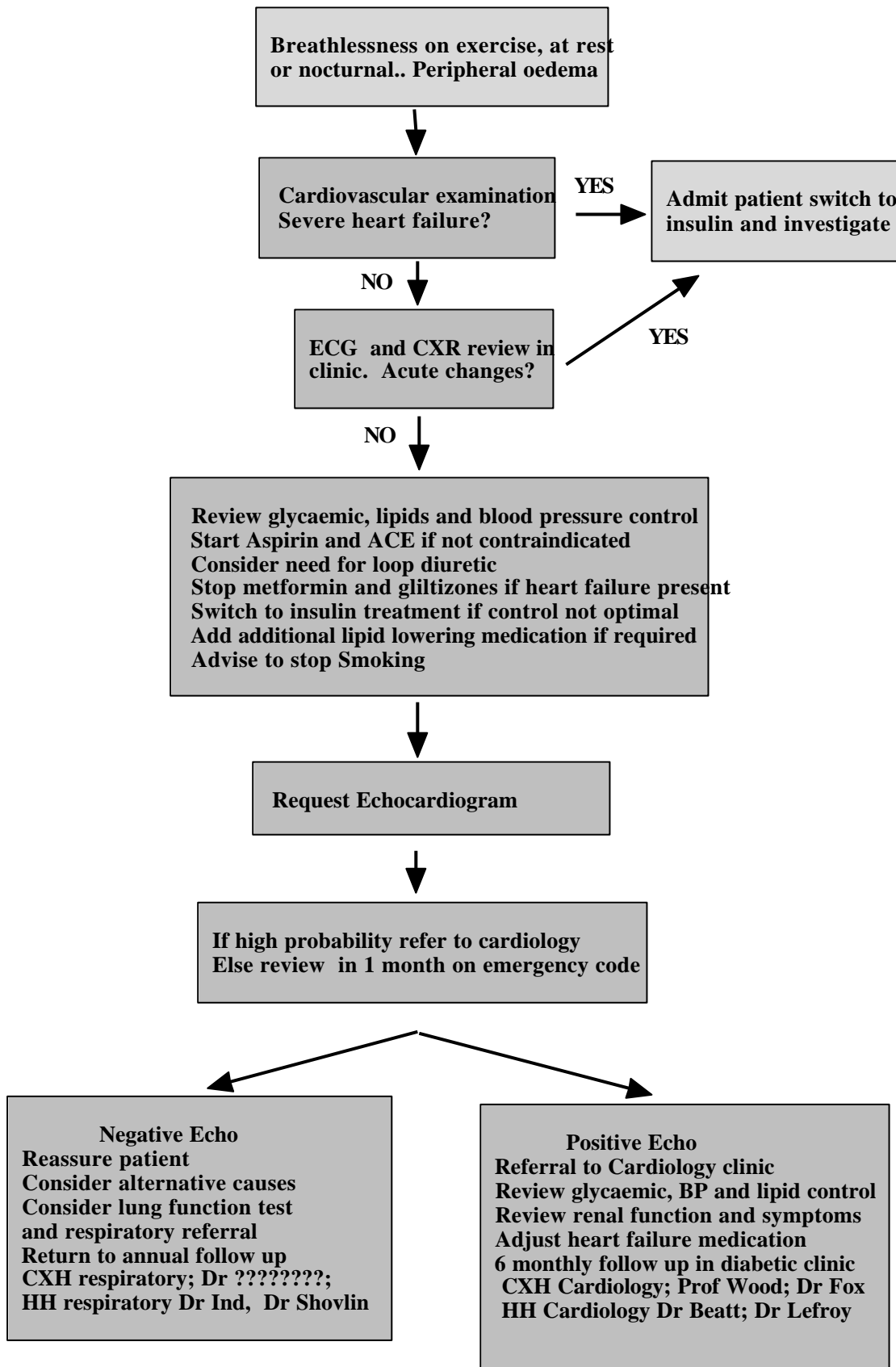
# Erectile Dysfunction



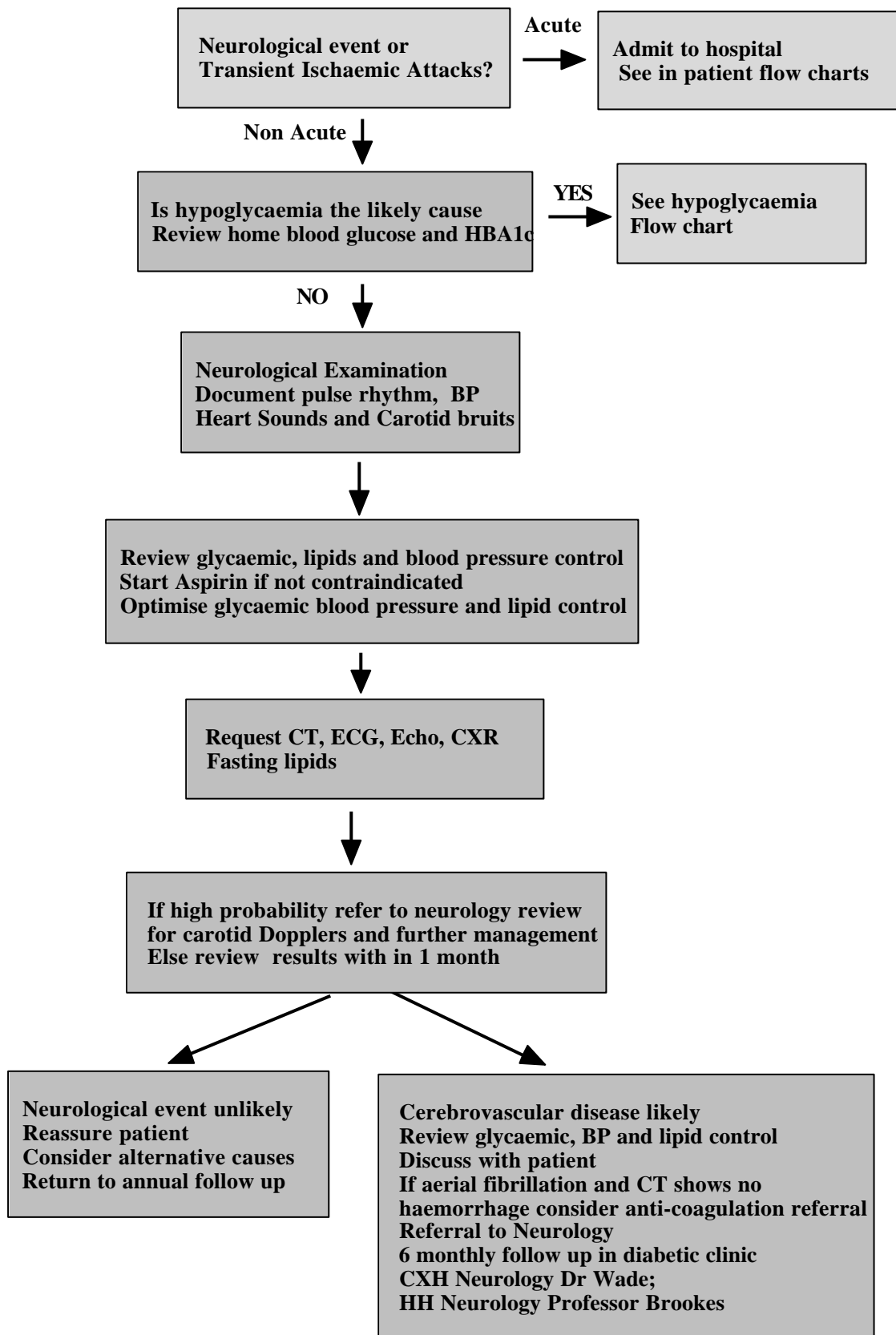
# Coronary Heart Disease



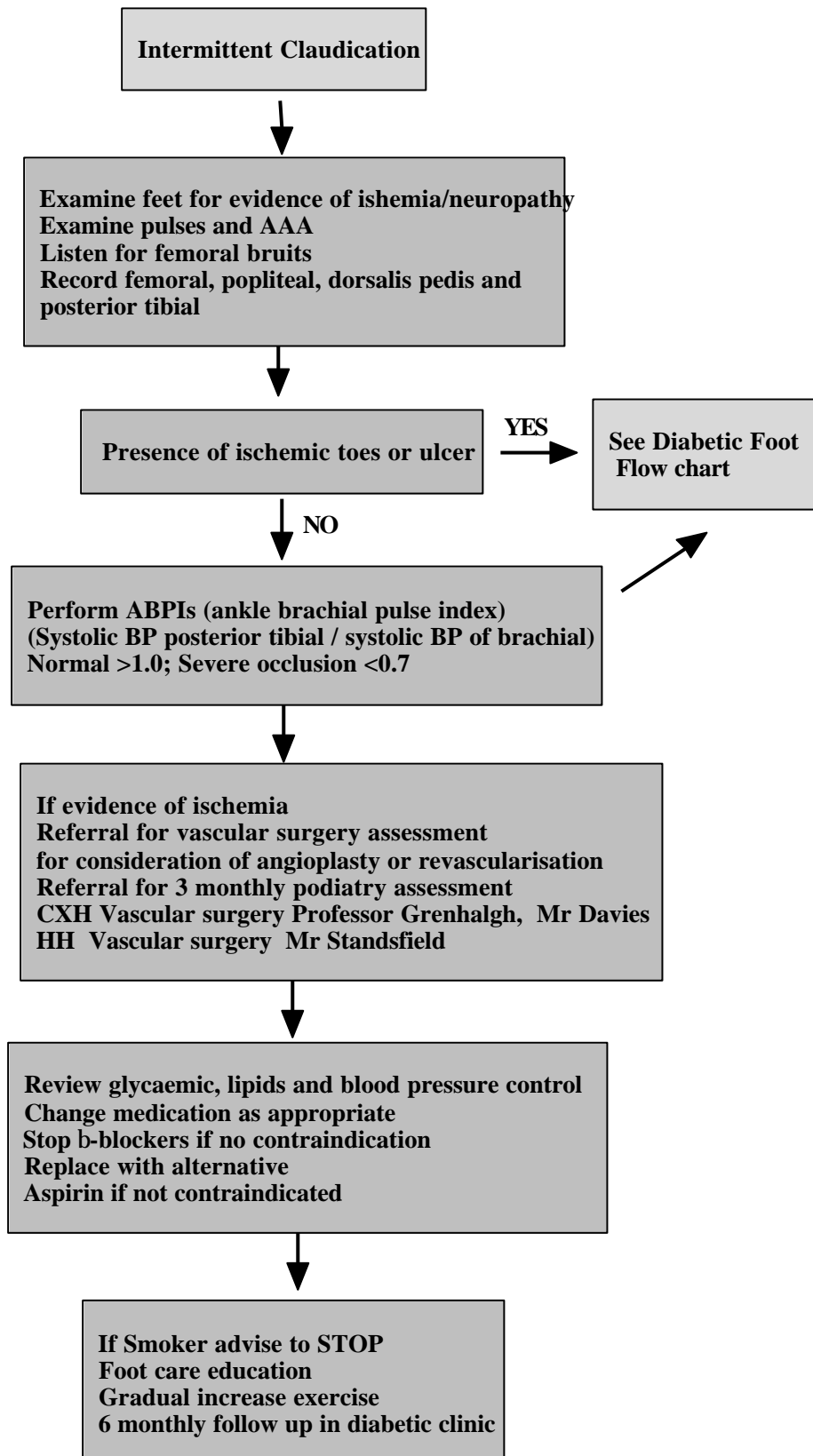
# Heart Failure



# Cerebrovascular Disease



# Peripheral Vascular Disease



# Diabetic Foot Assessment

	Right			Left		
Pulses	Palpable	Weak	Absent	Palpable	Weak	Absent
D Pedis						
Posterior Tibial						
ABPI	D pedis			Post Tibial		
Cap. Filling time				sec		
Colour	Cyanosed	Normal	Erythema	Cyanosed	Normal	Erythema
Toes						
Foot						
Temperature						
Toes						
Foot						
Vascular Signs & Symptoms	Present		Absent	Present		Absent
Intermittent Claudication						
Rest Pain						
Oedema						
Neurological	Present		Absent	Present		Absent
Pin prick						
10g Neurofilament						
Vibration perception						
	Hallux					
	Malleoli					
Reflexes	Present	Reduced	Absent	Present	Reduced	Absent
Knee						
Ankle						
Footwear	Suitable	<input type="checkbox"/>	Unsuitable	<input type="checkbox"/>	Specialist	<input type="checkbox"/>
						Orthotic
CLASSIFICATION	R		L	R		L
Stage One : Normal foot				Stage Four: Infection/ Necrosis		
Stage Two : High Risk				Stage Five : Amputation		
Stage Three : Ulceration						
Neuropathic				Neuroischaemic		

Clinicians name:	Date for review:
------------------	------------------



# Diabetic Foot

## Risk Factors

Previous Ulcer,  
 Poor Foot care, deformity  
 Retinopathy  
 Peripheral neuropathy (10g microfilament)  
 Autonomic neuropathy  
 Nephropathy  
 Elderly

## History

May be no history of pain, swelling warmth  
 Trauma or new footwear

## Examination

Document site and description of cellulitis/ulcer/deformity  
 Reflexes, LT, VS, PP, JPS, 10g micofilament  
 Pulses femoral, popliteal, DP and PT  
 Neuropathic ulcer: Warm clean punched out at pressure points  
 Neuroischemic ulcer: Cold pulseless foot, gangrenous heel/toes  
 Probe full depth of ulcer, if to bone then assume osteomyelitis  
 Always consider the possibility of Charcot Joints

## Investigations

U&E, CRP, HBA1c  
 FBC, ESR  
 Ulcer Swab (probe full depth of ulcer)  
 Foot X-ray  
 Bone scan if suspect OM/Charcot  
**ABPI** (Ankle-brachial pressure index)  
 (systolic BP at post tibial/ systolic BP at brachial)  
 Normal >1.0 <0.7 significant occlusion

**Minor superficial infection**

**Severe infection**

## Management

Optimise glycaemic control  
 Podiatry review (today and weekly)  
 Weekly foot swabs  
 Foot care education  
 Footwear assessment  
 Start oral antibiotics

**Amoxycillin 500mg tds**  
**Flucloxacillin 50mg qds**  
**Metronidazole 400 mg tds**

**Or Augmentin 625mg tds**  
 Penicillin allergic  
**Ciprofloxacin 500mg bd**  
 Regular U+E, FBC, CRP, LFT  
 Regular chiropody review when recovered

**No Improvement  
 after 1 week**

## Management

Podiatry review TODAY  
 Admit patient  
 Start insulin (basal bolus)  
 Start iv antibiotics

**Penicillin G 1.2g tds**  
**Flucloxacillin 500mg qds**  
**Metronidazole 400mg tds**

Osteomyelitis suspected  
**Ciprofloxacin 500mg bd**  
**Clindamycin 300mg qds**  
 Vascular surgical opinion  
 Tissue viability nurse assessment  
 Education regarding foot care  
 Footwear assessment  
 Regular U+E, FBC, CRP, LFT  
 Regular chiropody 3/12 when normal

# Charcot Joint

## History

Acute onset unilateral erythema, oedema and warmth  
Pain may or may not be present  
history of mild trauma  
Early diagnosis and treatment is essential  
Charcots usually involves midfoot  
Osteomyellitis usually preceded by an ulcer and affects metatarsal/calcaneum  
Differential diagnosis  
Cellulitis, Osteomyelitis, DVT, Gout

## Examination

Document site and description of cellulitis/ulcer/deformity  
Reflexes, LT, VS, PP, JPS, 10g micofilament  
Pulses femoral, popliteal, DP and PT  
Document lying and standing BP

## Investigations

U&E, AlkP, CRP, Urate , HBA1c  
FBC, ESR  
Foot X-ray  
Bone scan and then White Cell Scan

## Management

Podiatry review TODAY  
Admit patient  
Antibiotics if osteomyelitis is possible continue till excluded  
Optimise glycaemic control  
Radiology review +/- MRI scan  
Orthopaedic surgery review  
Immobilisation non wt bearing cast for 1 month  
Then total contact cast or aircast with very gradual mobilisation  
Regular CRP and AlkP

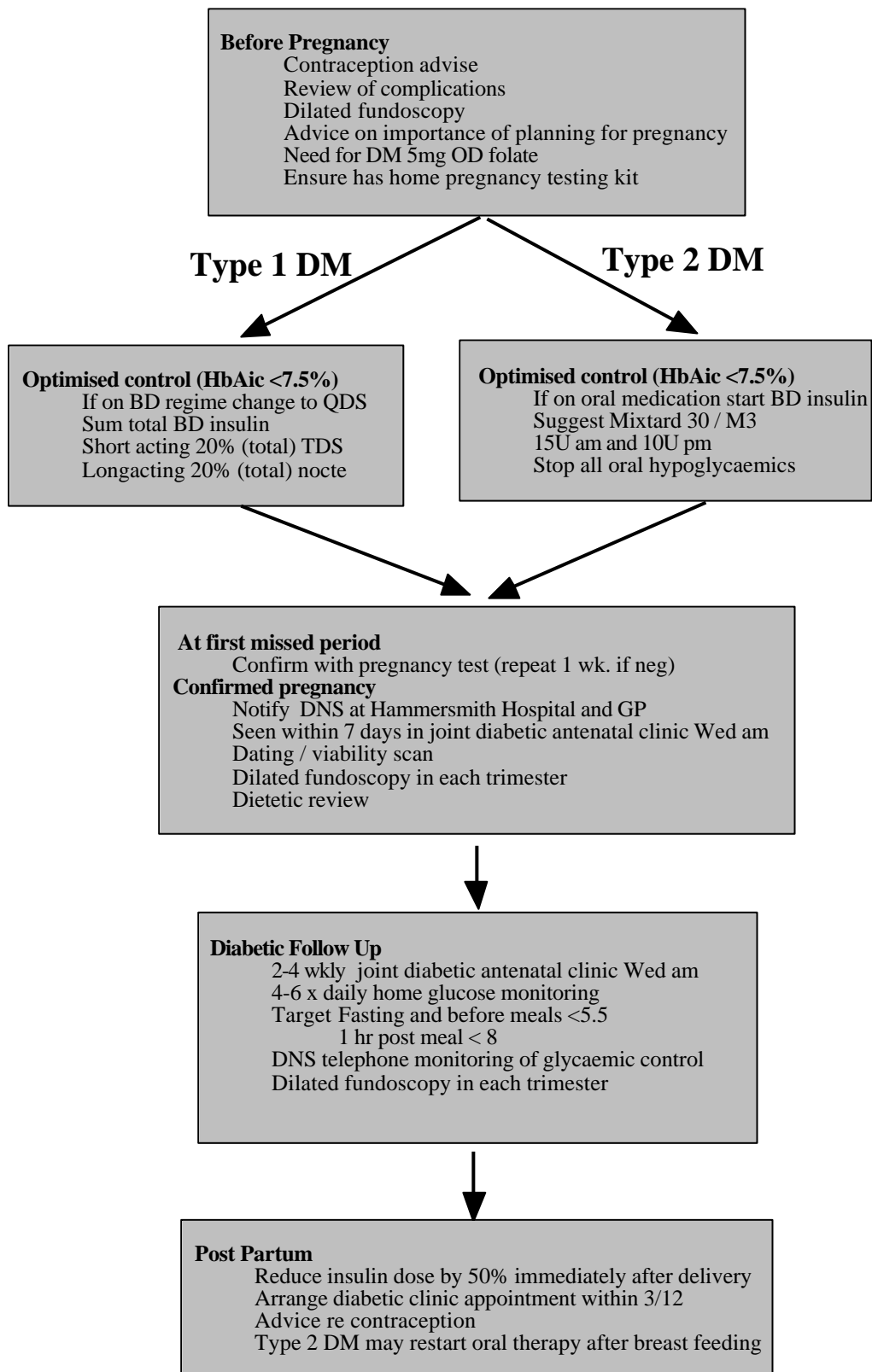
## Pharmacological

Bisphosphonates may help :  
( ? Pamidronate 90mg iv, or oral bisphosphonates)

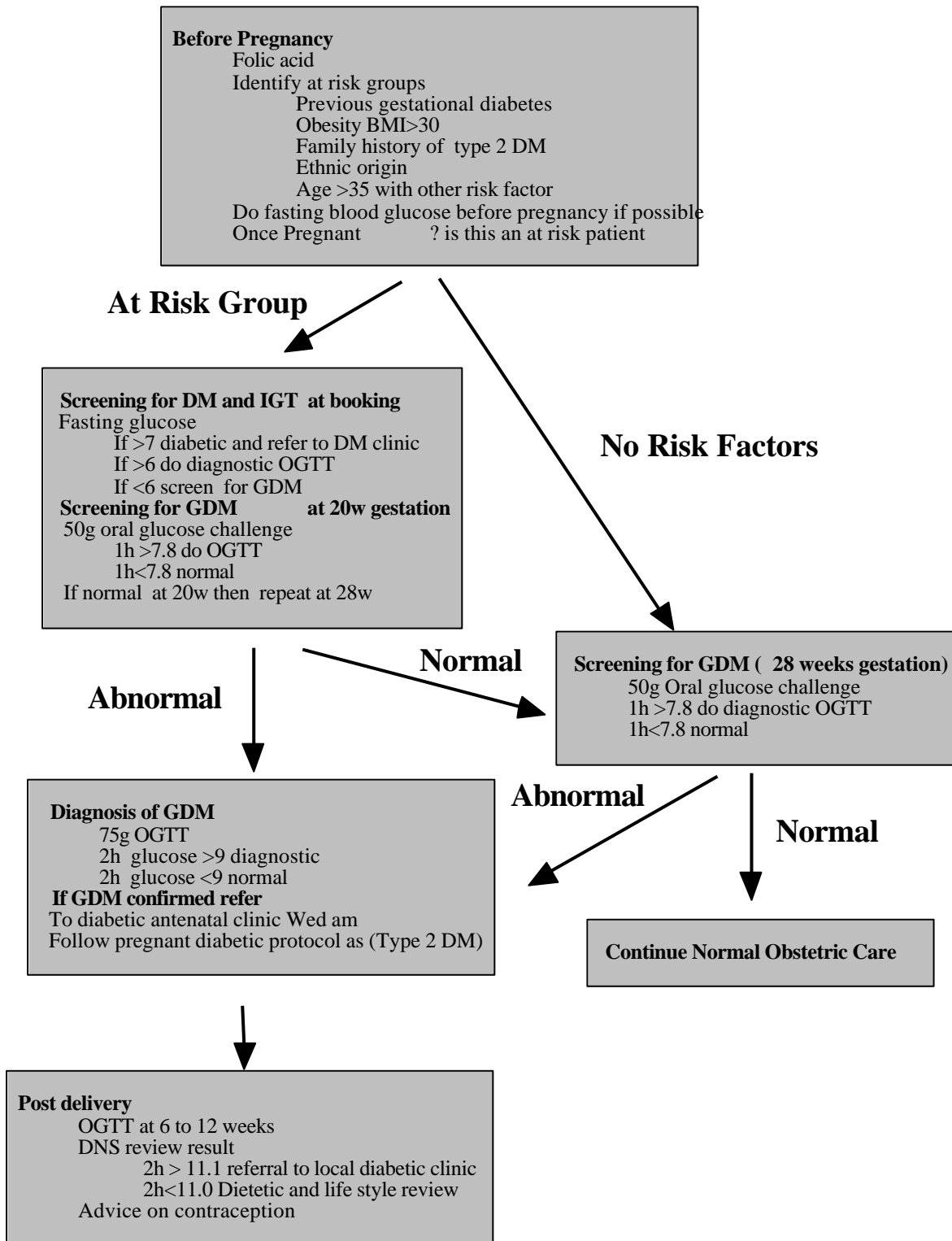
## Follow up

Footwear assessment moulded inserts etc  
Regular podiatry 3/12 when discharged  
Orthopaedic surgery follow up

# Pregnancy in Known Diabetics



# Gestational Diabetics



# Diabetic Ketoacidosis

( See also Hammersmith Hospitals Guidelines )

## DIABETIC KETOACIDOSIS OCCURS IN TYPE I DM

### Presentation (Usually young and thin)

Dehydration, tachypnoea, tachycardia, ketosis, ketonuria,  
Vomiting, abdo pain, drowsiness and confusion

### Diagnosis

Urinary ketones +2+, bicarbonate  $\leq 15$ mmol/l, pH <7.35

### Precipitating factors

New diagnosis type I, Infection, missed insulin, steroids  
MI, CVA, trauma, hyperthyroidism, pheochromocytoma

### Complications

Cerebral oedema, RDS, thromboembolism, mortality <4%

**AIM TO CORRECT ACIDOSIS WITH INSULIN AND IV FLUID AND RESTORE ELECTROLYTE BALANCE REGARDLESS OF GLUCOSE**

### Initial Investigations

Finger prick blood glucose (record in notes)  
Urine dipstick (record level of ketones in notes)  
FBC, U&E, Bicarbonate, Osmolality, Amylase  
Glucose, CK, Blood Gas  
Blood cultures X2 and Urine culture  
ECG, CXR  
?Pregnancy test, CT LP (if suspect meningitis)

### Initial Management

**ECG monitor, Oxygen if  $pO_2 < 10.5$  kPa**

**Urinary Catheter** (If elderly, impaired consciousness, no urine in 1h)

**CVP line** (If very unwell, BP < 90, pH < 7.0, Cardiovascular disease, or > 65y)

**NG tube** if impaired consciousness or vomiting (aspirate hourly)

**Consider ITU** (impaired consciousness, severe hypotension, pH < 7.0)

**Bicarbonate** if pH < 7.0 consider 100ml of 1.26% NaHCO<sub>3</sub> (CVP) hourly until pH > 7.0

Give 10-30mmol KCl in separate infusion

**IV fluids** (average deficit 4-6l; postural BP or CVP)

BM > 12 give 0.9% saline (usually at least first 4-5l)

BM < 12 give and pH < 7.3 give 10% glucose

Volume (1.5l in 1h; 1l in 1h; 1l in 2h; 1l in 4h; 1l in 8h)

### KCl supplements

Add KCl from the second litre of fluid (Stop if K > 6 or anuric)

KCl (K < 3 40mmol/h; K 3-4 30mmol/h; K 4-5 20mmol/h; K 5-6 10mmol/h)

### Insulin

Give Actrapid 20u SC stat; If dehydrated give 10u IV stat in addition

Put 50u actrapid in 50ml 0.9% saline in syringe pump

Give at 6ml/h until pH > 7.3 (Start 10% glucose if BM < 12)

When pH > 7.3 use 5% glucose if BM < 12, 0.9% saline if BM > 12

Use scale (BM 0-5 1u/h; BM 5-12 3u/h; BM > 12 4u/h)

Do not stop insulin if glucose low give IV glucose

Heparin 5000u 8h sc

Antibiotic if evidence of infections

### Monitor

Temp, P, BP, JVP (postural BP) GCS

Urine output and ketones; BM hourly, (>20 do glucose)

K<sup>+</sup>, pH and Bicarb 2-4 hourly all on blood gas

A venous sample (blood gas syringe) once pH > 7.2 is OK

### Continued Management

Contact endocrine registrar, diabetic nurse specialist and dietitian

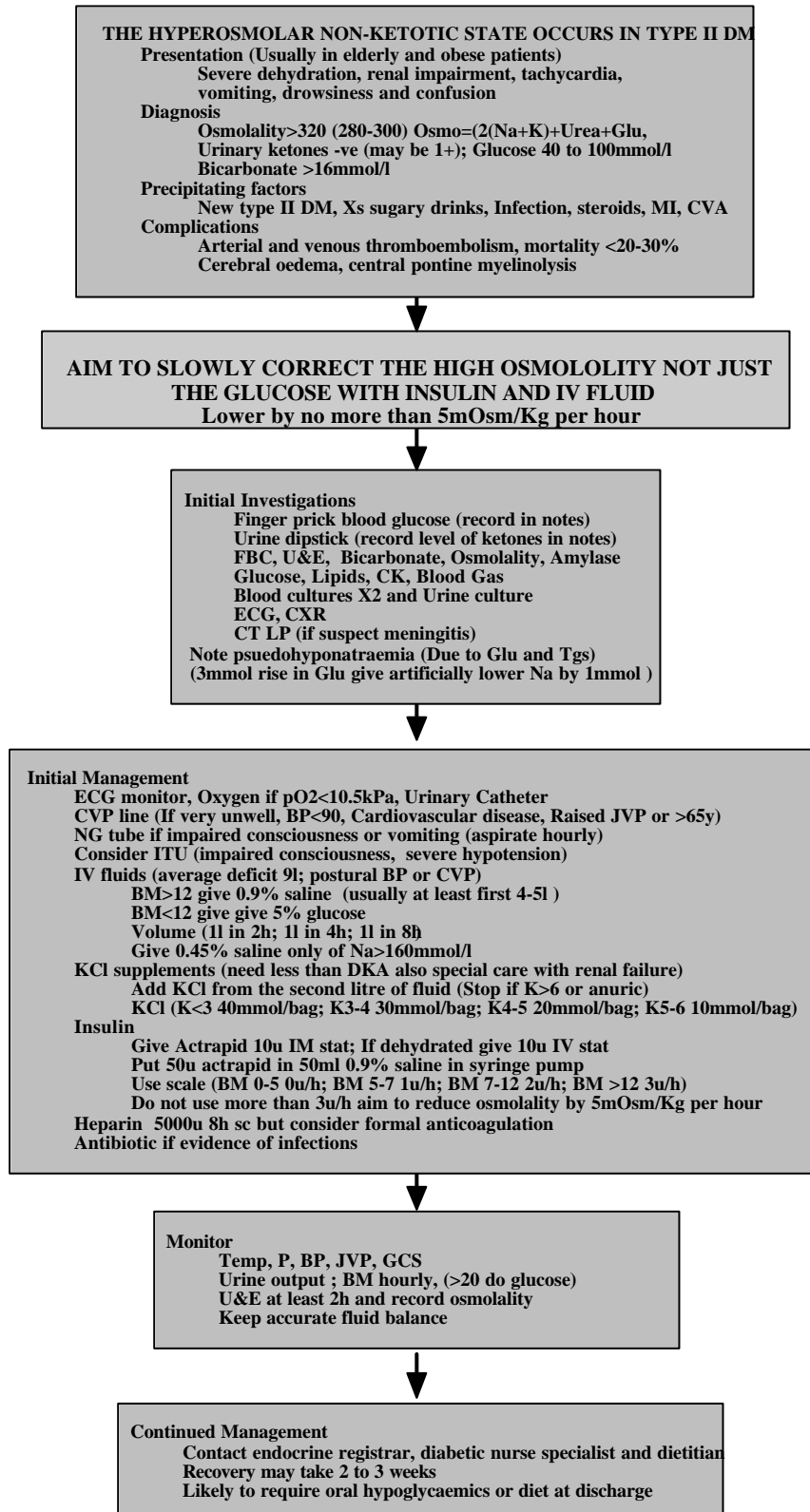
If no ketones and patient can eat and drink convert to s/c insulin

May need 10-20% higher than normal dose (acidotic resistance)

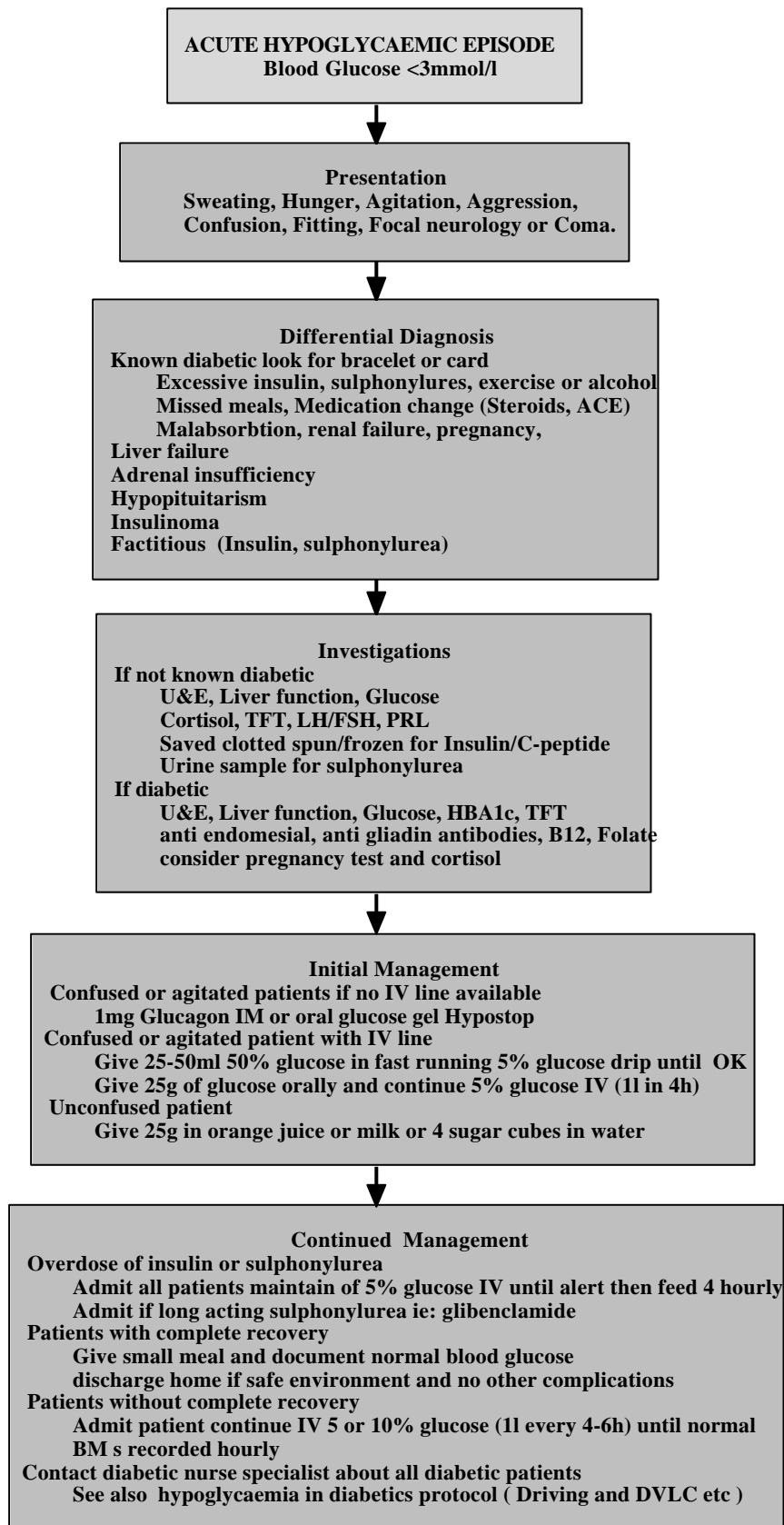
give sc insulin before meal, 1h after meal stop iv insulin

# Hyperosmolar Non-Ketotic State

( See also Hammersmith Hospitals Guidelines )



# Acute Hypoglycaemia



# Diabetics and MI (DIGAMI Protocol)

**MYOCARDIAL INFARCTION IN DIABETICS**



**DIGAMI 1997**  
Diabetics and patients admitted with a Glu>11  
Treated for 24h by IV insulin/Glucose and  
Then 3/12 qds basal bolus insulin  
**REDUCED 3Y MORTALITY BY 11%**



Manage as per Myocardial Infarction except  
Check formal Lab Glucose at admission  
HBA1c (exclude stress hyperglycaemia)  
STOP all oral hypoglycaemics  
Check for proliferative retinopathy or haemorrhage  
before thrombolysis



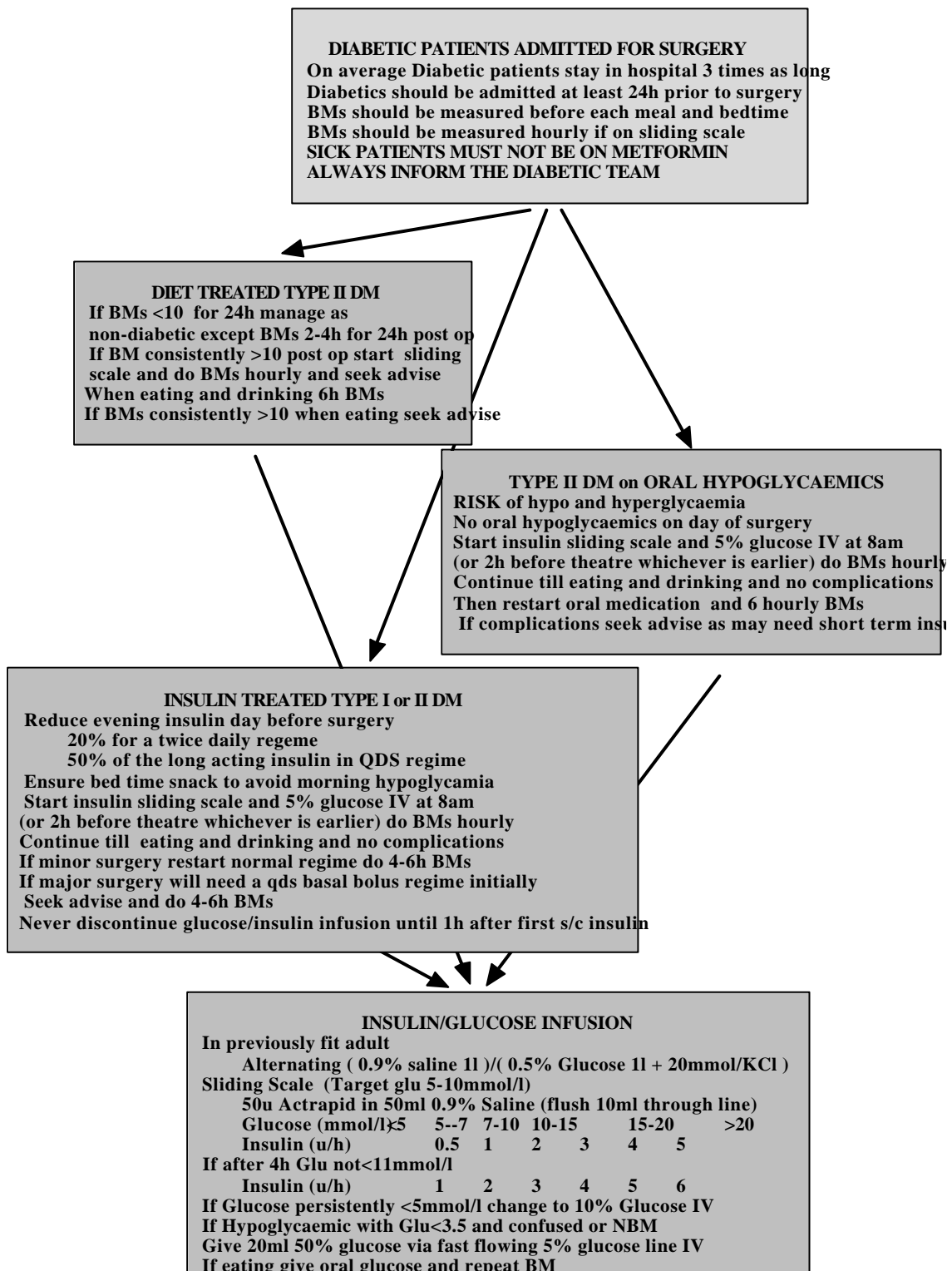
**INSULIN/GLUCOSE INFUSION**  
Place 80u Actrapid insulin in 500ml of 5% glucose ( 1u/6ml )  
Start at 30ml/h check BM at 1h  
BM >15 Give 8u Actrapid IV increase rate by 6ml/h  
BM 11-15 Increase infusion by 3ml/h  
BM 7-11 No change in infusion rate  
BM 4-7 Decrease infusion by 6ml/h  
BM <4 Stop infusion 15 min retest every 15 min till BM 7  
If rate changed check BMs after 1 h  
If BM>11 and decrease >30% no change  
If BM 7-11 and decrease >30% reduce by 6ml/hr  
Then check BMs 2h  
After 10pm if glucose stable and  $\leq 11$  reduce rate by 50% over night



**S/C INSULIN**  
After minimum of 24h convert s/c insulin  
If previously well controlled on bd insulin and HBA1c <7.5 restart this  
Else Actrapid 4u tds : Insulatard 6u  
Contact Endocrine Registrar, diabetic nurse specialist and Dietitian



# Surgical Patients with Diabetes



# Incidental Diabetic Diagnosis

## INCIDENTAL DIAGNOSIS OF DIABETES IN CASUALTY

### Presentation

Polyuria, polydipsia, blurred vision, weight loss, cutaneous sepsis,

### Diagnosis

Random glucose  $\cdot 11.1$



### Examination must document

Blood pressure lying and standing

Peripheral pulses

Lower limb reflexes, peripheral sensation

Examination of feet for infection, ulcers, Charcots

Visual acuity and fundoscopy



### Initial Investigations

Finger prick blood glucose (record in notes)

Urine dipstick (record level of ketones in notes)

FBC, U&E, LFTs, Lipids, HBA1c, TFTs, Glucose

Bicarbonate, Osmolality, Amylase, Blood Gas

Consider ECG, CXR



### Patient Admission

Admission is necessary if

Not eating and drinking normally

Nausea or Vomiting

Ketoacidosis (pH  $< 7.35$ )

Other complicating problems

Admission is not necessary if

Patient is otherwise well, no sepsis/ulcers

Eating and drinking normally



### Further Management

#### Type I DM

Patients will need to see the diabetic nurse specialist to start insulin on that day or at next possible opportunity in working hours

Referral to diabetic consultant for out patient clinic

#### Type II DM

Patients can be seen routinely but contact diabetic nurse specialists

If osmotic symptoms consider starting Gliclazide 80mg od

If no osmotic symptoms, obese, normal renal/liver function

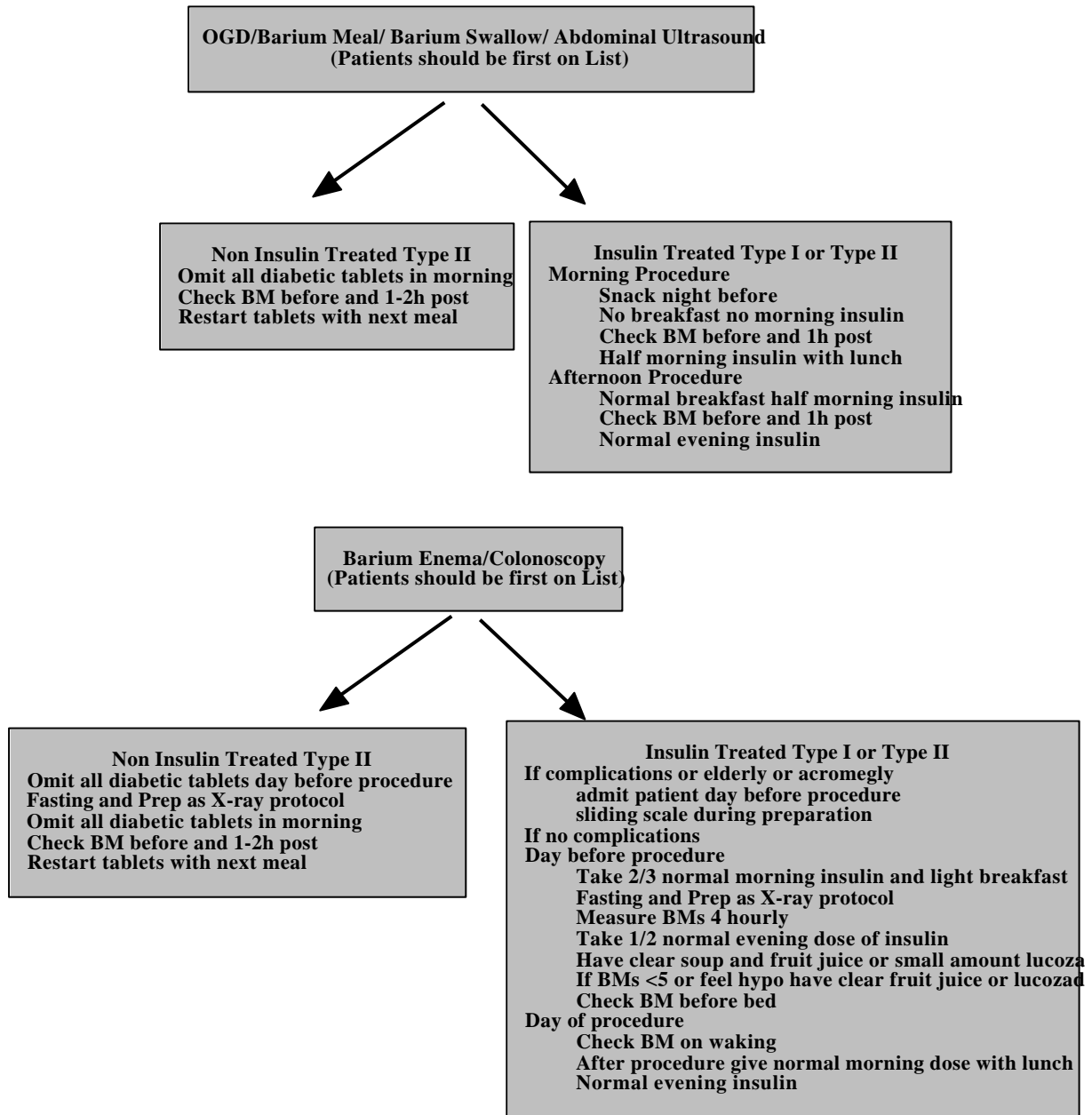
consider commencing metformin 500mg bd

Letter to GP patient to see GP within 2 days

Explain diagnosis and give dietary advice regarding sugar and sugary drinks

To return to A&E if unwell or not eating or drinking

# Diabetics and Out Patient Procedures



# METFORMIN AND IV CONTRAST

The following protocol is used by the radiology departments at Hammersmith and Charing Cross Hospitals.

## **Protocol for diabetic patients on Metformin having IV contrast.**

- 1) The patient should be given the information sheet when the investigation is booked. Details of the requesting doctor and their contact number should be added to the sheet.
  
- 2) An information sheet should be sent to the doctor requesting the investigation.
  
- 3) The patient should not take metformin on the day of the contrast examination. Alternative medication does not need to be prescribed.
  
- 4) Before leaving the department the patient should be given a completed U+E form and told to have the blood taken 48 hours later (either in the hospital or by the GP). They will also be given the name and contact number of the doctor who requested the investigation.
  
- 5) The day after the blood test, the patient should contact the doctor who has ordered the investigation. The doctor will advise them whether or not metformin can be restarted. (Metformin should not be restarted unless the creatinine is less than 150).

## **Information for diabetic patients on Metformin having IV contrast medium**

Dear patient

You have been given an appointment to have an investigation which uses an injection of 'contrast medium'. Very occasionally, contrast medium with metformin can result in illness. In order to prevent this, you must follow the guidelines below;

- 1) Do not take your metformin on the day of the contrast test. You do **not** need to have a different medicine whilst the metformin is stopped.
- 2) Before you leave the X-ray department, please make sure you have a blood test form and the name and contact number of the doctor who requested your test.
- 3) 48 hours after your contrast test, you must have the blood test requested (either in the hospital, at clinic 6+7 on the 1<sup>st</sup> floor or from your GP).
- 4) The day after the blood test is taken, you must ring the doctor below who requested the test. They will tell you if you can restart the metformin.

**The doctor to contact is** \_\_\_\_\_

**Their contact number is** \_\_\_\_\_

## **Information for doctors requesting investigations with IV contrast medium in diabetic patients on Metformin**

Dear Doctor

RE:

You have requested an investigation involving the use of IV contrast medium on a patient taking metformin. There are case reports of lactic acidosis in patients on metformin due to renal impairment following IV contrast. Therefore your patient has been given the following information

- 1) The patient should not take metformin on the day of the contrast examination. Alternative medication does not need to be prescribed.
- 2) Before leaving the X-ray department the patient will be given a completed U+E form and told to have the blood taken 48 hours later (either in the hospital or by the GP). **They will also be given the name and contact number of the doctor who requested the investigation.**
- 3) The day after the blood test, **the patient should contact the doctor who has ordered the investigation.** The doctor will advise them whether or not metformin can be restarted.

Your patient will contact **you** 24 hours after their renal function has been checked for advice on whether or not to restart metformin.

Metformin should not be restarted unless the creatinine is less than 150.



# The Initial Diagnosis of Diabetes

## SYMPTOMS AT DIAGNOSIS MAY INCLUDE

Thirst  
Frequent passing of urine  
Weight loss  
Blurred Vision  
Recurrent infections

## DIAGNOSIS OF DIABETES

Must be made in one of 3 ways

- 1) A random blood sugar greater than 11
- 2) 2 fasting blood sugars greater than 7
- 3) A blood sugar of  $>11$ , 2 hours after drinking 75g of sugar in water

## DIABETES MELLITUS

Insulin is the important hormone in sugar (Glucose) control  
Diabetes is a failure of the body to control its sugar correctly  
There are 2 ways this can happen and so 2 types of diabetes

### Type I diabetes

Caused by destruction of the cells that make insulin in the pancreas  
There is therefore a total lack of insulin  
Usually occurs under the age of 30 years  
Always requiring insulin treatment

### Type II diabetes

Caused by an increased resistance to the effects of insulin  
Usually occurring over 30 years of age  
Can usually initially be treated by diet or tablets but may require insulin later  
Diet, weight loss and exercise increase sensitivity to insulin and may therefore reduce the amount of treatment required



# Complication of Diabetes

## PROBLEMS ASSOCIATED WITH DIABETES

The associated problems or complications of diabetes may be present at the time of diagnosis or may occur at any time after the diagnosis

The reason for careful follow up of diabetic patients is to significantly reduce the chance of these problems occurring.

The complications of diabetes are due to the long term damage of the large and small blood vessels in the body due to the effects of a high blood sugar, blood pressure and cholesterol.

## DAMAGE TO LARGE BLOOD VESSELS INCREASED RISK OF

- Heart attacks due to poor blood supply to the heart
- Strokes due to poor blood supply to the brain
- Calf pain on walking due to poor blood supply to the legs
- Foot infections that may lead to amputation

## DAMAGE TO SMALL BLOOD VESSELS INCREASED RISK OF

- Damage to the eye that may result in poor vision or blindness
- Damage to the kidney that may result in kidney failure and dialysis
- Damage to nerves that may result in numbness or pain, damage to the feet and ankles and problems with erections in men.

## THE FOLLOWING REDUCE THE CHANCES OF THESE COMPLICATION

- Not Smoking
- Good control of blood sugar levels (HbA1c <7.5%)
- Good control of blood pressure <140/80
- Good control of cholesterol Cholesterol/HDL <5.0
- Normal body weight Body Mass Index <30
- Healthy life style (Diet and exercise) At least 30 minutes exercise 3 times a week
- Good foot care

**Please ask your Doctor about your results each time you are seen**

# New Diabetic Outpatient Assessment

## ATTENDING APPOINTMENTS

**It is essential that you attend all you appointments with the members of the diabetic team. If you fail to attend an appointment and have not cancel it in advance with appointments department you will be discharged from the clinic.**

## NEW REFERRAL RECEIVED AND REVIEWED BY CONSULTANT

Priority decided by consultant  
Nurse and dietician clinic appointment booked  
Medical outpatient appointment booked  
(Diabetic diagnosis confirmed if not stated in referral)

## NEW PATIENT DIETICIAN CLINIC

Dietary history and advise on diabetic diet  
Advise on low fat and weight loss diets as appropriate  
Life style advise

## NEW PATIENT DIABETIC NURSE SPECIALIST CLINIC

Initial assessment and history  
Education about Diabetes and life style advise  
Initial treatment decided (diet or tablets or insulin)  
Initial treatment started  
Blood tests performed  
    Full blood count  
    Kidney function, Liver function, Lipid Profile, Thyroid function  
    Glucose and HBA1c (long term sugar control measurement)  
Initial urine tests  
ECG in type II diabetics  
Follow up appointment made if necessary

## NEW PATIENT MEDICAL OUTPATIENT CLINIC

Medical history  
Physical examination  
Assessment of home glucose monitoring  
Assessment of blood test results (Organise further tests if required)  
Further diabetic education  
Adjustment of medication for diabetes  
If needed start medication for high blood pressure and high cholesterol  
If needed referral to other specialists  
Decide on Hospital or GP follow up

# Follow up for diabetic patients

## **CONTINUED HOSPITAL OUT PATIENT FOLLOW UP**

- Diabetics treated with Insulin
- Diabetics with complications affecting their
  - Eyes (Retinopathy)
  - Kidneys (Nephropathy)
  - Nerves (Neuropathy)
  - Heart
  - Feet
- Diabetics with poorly controlled
  - Sugar
  - Blood pressure
  - Cholesterol
  - Severe obesity
- Other complex medical problems

## **LOCAL FOLLOW UP WITH GENERAL PRACTITIONER**

- Diabetics controlled with diet only
- Diabetics with no complications on tablets and
  - Good sugar control
  - Good blood pressure control
  - Cholesterol within guidelines

## **DIABETIC PATIENT FOLLOW UP SHOULD INCLUDE**

- Diabetics with complications should be seen at 6 monthly
- Diabetics with no complications should be seen at least yearly
- At each yearly review the patient should have their
  - Blood pressure checked
  - Feet examined
  - Vision tested
  - Eyes dilated and examined
- Urine tested for microalbuminuria
- Blood tested should include
  - Kidney function, Liver function
  - Lipids (Cholesterol)
  - HbA1c (long term sugar control)

# Pregnancy and Diabetes

## **BEFORE PREGNANCY**

Use contraception until planning pregnancy

If planning pregnancy consult your doctor as soon as possible

It is very important that you have excellent glucose control before pregnancy

You should

- Have all your medications reviewed

- Commence Folate 5mg once a day

- Have your eyes dilated and assessed

- Obtain a pregnancy testing kit

## **In Type 1 Diabetes**

Your insulin should be switched to 4 times a day regime

## **In Type 2 Diabetes**

You should be treated with a twice daily insulin regime

Your diabetic tablets will be stopped

You should do regular blood glucose monitoring

Insulin doses will be adjusted so your HbA1c is <7.5%

## **DURING PREGNANCY**

Check a pregnancy test if you miss a period

If this is negative repeat test after 1 week

If Pregnancy test is positive

Notify GP and Diabetic nurse specialists at Hammersmith Hospital

You will be seen within 7 days at the diabetic antenatal clinic

You will then have 2 to 4 week follow up appointments at this clinic

Your eyes should be dilated and assessed 3 times during pregnancy

## **Glucose control during pregnancy**

You should do glucose monitoring 4 to 6 times per day

Glucose targets are

- Before meals glucose <5.5

- 1 hour after meals glucose <8

The diabetic nurse specialists will advise you about your control by phone

# Appendix I

# Appointment Policy

## **NEW REFERRAL POLICY**

### **GP in Hammersmith Area (W12/W3/(small part W5))**

Appointment with DNS, Dietitians and out patients HH (DNP)

### **GP in Charing Cross Area (W4/W6/W14/SW6)**

Appointment with DNS, Dietitians and out patients CXH (M24)

### **GP outside these areas**

Send out of area letter (Appendix I)

## **NEW PATIENT POLICY**

### **DNA the DNS appointment**

See patient at the medical outpatient appointment

### **Patient seen in medical outpatient**

Follow up DICL (HH) or D26/U12(CXH)

### **DNA new patient appointment**

Discharge patient letter to GP re referral (appendix I)

## **FOLLOWUP PATIENT POLICY**

### **Patient seen in medical outpatient**

Follow up DICL(HH)or D26/U12(CXH)

discuss with consultant if <1 year follow up

### **DNA followup appointment**

make next routine follow up DICL (HH) or D26/U12(CXH)

if urgent discuss with consultant

### **2XDNA followup appointment**

Discharge patient letter to GP re referral (appendix I)

# Diabetic Clinic Codes

## Hammersmith Hospital

### DNS clinics

Tuesday am new patients	DBNC(6)
Tuesday pm follow up	BDN2(6)
Thursday am follow up	DTNC(14)
Thursday pm follow up	DTN2(5)

### Medical outpatient clinics

Diabetic new patient list	DNP(10)
Diabetic Type II follow up	DICL(40)
Diabetic Type I follow up (1 <sup>st</sup> week even months)	DMT1(40)
Diabetic Emergency follow up (within 2/12)	DEM(5)
Diabetic Foot clinic (1 <sup>st</sup> week of month)	DMFT(5)

### Medical retinopathy clinics

Diabetic retinopathy	DOC1(30)
Diabetic retinopathy screening	DRFC

## Charing Cross Hospital

### DNS clinics

Monday am follow up review clinic	D28(14)
Tuesday pm new type II patients	D24(3)
Thursday am new patient clinic	D105(5)
Friday am review clinic	D106(14)

### Medical outpatient clinics

Diabetic new patient clinic	M24(10)
Diabetic follow up Type II	D26(40)
Diabetic follow up Type I (1 <sup>st</sup> week even months)	U12(40)
Diabetic Emergency follow up (within 2/12)	444(5)
Eyes only review (within 2/12)	K74(3)
Diabetic Foot clinic	FTDM(6)

# **Standard Letters for** **Hammersmith Hospital**

**Area for diabetic clinic W3, W12 small part W5**

<b>New referral out of area letter</b>	<b>68</b>
<b>No diagnostic criteria letter</b>	<b>69</b>
<b>OGTT result letter</b>	<b>70</b>
<b>New patient DNA letter</b>	<b>71</b>
<b>One time DNA and out of area letter</b>	<b>72</b>
<b>One time DNA and in area letter</b>	<b>73</b>
<b>Two times DNA in area letter</b>	<b>74</b>
<b>Discharge letter</b>	<b>75</b>
<b>Letter to all clinic patients attending clinic</b>	<b>76</b>
<b>Appointment request from patient</b>	<b>77</b>
<b>Information sheet to discharged patients</b>	<b>78</b>



Direct Line: 020 8383 4823

Internal Ext: 34823

Fax: 020 8283 3360

Appointments 020 8383 5000

Ref:

Date:

Clinic Date:

**HAMMERSMITH HOSPITAL DIABETIC CLINIC**

GP Address

Dear Dr

RE:

Thank you for your referral letter for this patient with Diabetes. Unfortunately due to the high rate of new local referrals and our follow up clinics being fully booked for more than 1 year, it is difficult for us to see patients from outside our local area. I would be grateful therefore if you could, in the first instance refer this patient to your local Diabetes services.

Our local area includes W3, W12 and a small part of W5.

Kind regards.

Yours sincerely,

**Anne Dornhorst DM FRCP/ Duncan Bassett MA PhD MRCP  
Consultants in Endocrinology and Diabetes**

Direct Line: 020 8383 4823

Internal Ext: 34823

Fax: 020 8283 3360

Appointments 020 8383 5000

Ref:

Date:

Clinic Date:

**HAMMERSMITH HOSPITAL DIABETIC CLINIC**

GP Address

Dear Dr

RE:

Thank you for your referral letter for this patient. Unfortunately your referral contains no diagnostic criteria for diabetes. Due to the high frequency of new diabetic referrals would be grateful if you would confirm the diagnosis prior to referring the patient.

**Diagnostic Criteria (HBA1c CAN NOT BE USED FOR DIAGNOSIS)**

**2x Random Glucose Measurement (1x if symptomatic)**

=11.1 Diabetes Mellitus  
<11.1 Normal if =6.1 do 2x fasting

**2x Fasting Glucose Measurement (1x if symptomatic)**

=7.0 Diabetes Mellitus  
=6.1 but <7.0 Impaired Fasting Glucose do OGTT  
<6.1 Normal

**Oral Glucose Tolerance Test 2h Glucose**

=11.1 Diabetes Mellitus  
=7.8 but <11.1 Impaired Glucose Tolerance  
<7.8 Normal

We are most grateful for your help with this and will make a new patient appointment as soon as we receive the re-referral

Kind regards.

Yours sincerely,

Anne Dornhorst DM FRCP/ Duncan Bassett MA PhD MRCP  
Consultants in Endocrinology and Diabetes

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Fax: 020 8283 3360

Appointments: 020 8383 5000

DATE

**HAMMERSMITH HOSPITAL DIABETIC CLINIC**

GP Address:

GP Address Sticker

Re patient:

Patient Sticker

Your patient had an oral glucose tolerance test on .....

The results demonstrate that the patient.....

WHO Diagnostic Criteria

**Fasting Glucose**

=7.0	Diabetes Mellitus
=6.1 but <7.0	Impaired Fasting Glucose
<6.1	Normal

**2h Glucose during Oral Glucose Tolerance Test**

=11.1	Diabetes Mellitus
=7.8 but <11.1	Impaired Glucose Tolerance
<7.8	Normal

If the patient has impaired fasting glucose or impaired glucose tolerance we would recommend life style advice and at least yearly a fasting glucose.

We have not made any further follow up appointment.

Kind regards.

Yours sincerely,

Anne Dornhorst DM FRCP/ Duncan Bassett MA PhD MRCP  
Consultants in Endocrinology and Diabetes

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Internal Ext: 34823

Fax: 020 8283 3360

Appointments: 020 8383 5000

Clinic Date:

**HAMMERSMITH HOSPITAL DIABETIC CLINIC**

GP Address:

GP Address Sticker

Re patient:

Patient Sticker

We are sorry that your patient did not attend their appointment at the new diabetic clinic.

Due to the high frequency of new patient referrals we have not made a follow up appointment for your patient. If you wish to re-refer the patient we would be grateful if you could confirm that they are prepared to will attend the hospital diabetic clinic. n addition, we would be grateful if you could and ask them to rearrange the appointment date if they are unable to attend as their non-attendance prevents other urgent patients from being seen.  
Kind regards.

Yours sincerely,

**Anne Dornhorst DM FRCP/ Duncan Bassett MA PhD MRCP**

**Consultants in Endocrinology and Diabetes**

cc Patient

Direct Line: 020 8383 4823

Internal Ext: 34823

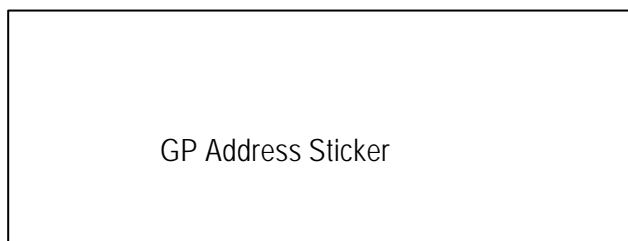
Fax: 020 8283 3360

Appointments: 020 8383 5000

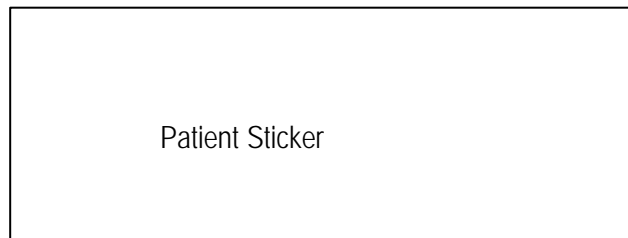
Clinic Date:

**HAMMERSMITH HOSPITAL DIABETIC CLINIC**

GP Address:



Re patient:



We are sorry to say that your patient has not attended today's clinic appointment and it may be because of transport difficulties to the Hammersmith Hospital.

If you would like diabetic clinic follow-up for this patient, the patient may be more compliant if you referred them to a more local hospital.

We have not arranged another appointment.

Yours sincerely,

**Anne Dornhorst DM FRCP/ Duncan Bassett MA PhD MRCP**



**Consultants in Endocrinology and Diabetes**

cc Patient

Direct Line: 020 8383 4823

Internal Ext: 34823

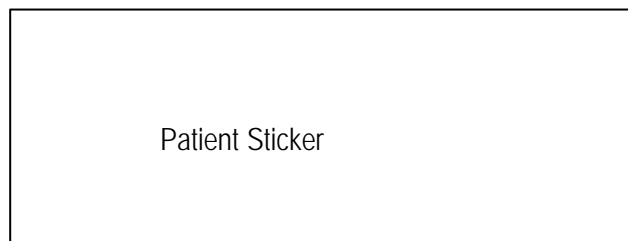
Fax: 020 8283 3360

Appointments: 020 8383 5000

Clinic Date:

### HAMMERSMITH HOSPITAL DIABETIC CLINIC

Patient:



We are sorry that you did not attend the diabetic clinic today

Because of the large number of diabetic patients already under follow up and the high rate of new referrals the clinic is fully booked for the next year and there are no routine appointments available in the next 12 months. This means if you miss a diabetic clinic appointment you will not receive a new appointment for another year. **It is therefore very important that you attend all your diabetic appointments and have a blood test and urine test done at Hammersmith Hospital 2 weeks before the appointment.**

If you fail to attend 2 appointments in succession you will be discharged from the clinic and will require a new referral from your GP to be seen again.

Of course should you develop a new problem your GP can write an emergency referral and we may see you sooner if this is necessary.

We have given you the next available routine appointment, which will be in about 1 year. The details of which will be sent to you in the next few weeks. During the next few months we would advise you to see your GP to check your blood pressure, vision, renal function, glucose control (HbA1c) and cholesterol.

Yours sincerely,

**Anne Dornhorst DM FRCP/ Duncan Bassett MA PhD MRCP**  
**Consultants in Endocrinology and Diabetes**

cc GP

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Internal Ext: 34823

Fax: 020 8283 3360

Appointments: 020 8383 5000

Clinic Date:

**HAMMERSMITH HOSPITAL DIABETIC CLINIC**

GP Address:

GP Address Sticker

Re patient:

Patient Sticker

We are sorry to say that your patient has not attended the last 2 diabetic clinic appointments.

It may be that the patient has moved address. If you have a different address and you would still like diabetic clinic follow-up for this patient, please send us another referral with the correct address.

We have not arranged another appointment.

**Anne Dornhorst DM FRCP/ Duncan Bassett MA PhD MRCP**

**Consultants in Endocrinology and Diabetes**

cc Patient

Direct Line: 020 8383 4823  
Internal Ext: 34823  
Fax: 020 8283 3360  
Appointments 020 8383 5000

Ref:  
Date:  
Clinic Date:

**HAMMERSMITH HOSPITAL DIABETIC CLINIC**

GP Address

GP Address

RE:

Patient  
Sticker

Your patient was reviewed in the diabetic clinic today and is considered suitable for discharge from this clinic as they have type II diabetes, are well controlled on diet / sub-maximal oral therapy and have no significant diabetic complications.

**Present Medication**

**Results Date:**

Blood Pressure (<145/85)  
BMI (<35)  
HbA1c (<8.5%)  
Creatinine (<125mmol/l)  
Chol/HDL ratio (<5.5)  
Urine Alb/Cre ratio (<3.0)  
Acuity Right: Left:

We have advised your patient that they should be seen at least annually in the practice. Annual review should include a minimum of blood pressure measurement, examination of the feet for peripheral pulses and evidence of peripheral neuropathy. Visual Acuity and dilated funduscopy should be performed annually either in the practice, by an optician or at the Hammersmith retinal screening clinic. Blood test should include renal function, full fasting lipid profile including HDL and LDL, HbA1c and liver function and CK where appropriate. Urine testing should include urine dipstick analysis and urine albumin/creatinine ratio.

We will be happy to see your patient again in the Hammersmith diabetic clinic should they develop diabetic complications or their control fall outside the above ranges.

Kind regards.

Yours sincerely,

**Anne Dornhorst DM FRCP/ Duncan Bassett MA PhD MRCP**  
**Consultants in Endocrinology and Diabetes**

Cc Dr Andrew Western

Cc The Patient

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Internal Ext: 34823  
Fax: 020 8283 3360  
Appointments 020 8383 5000

Ref:  
Date:  
Clinic Date:

## HAMMERSMITH HOSPITAL DIABETIC CLINIC

### To all patients attending the Hammersmith diabetic follow up clinic

Dear Sir/Madam,

This letter is just to inform you about out patient appointments in the Hammersmith diabetic clinic and to help us give a better service to you and all our diabetic patients.

With the present number of staff we are only able to see diabetic patients once per year unless they have diabetic complication affecting their eyes or kidneys when they will be seen every six months.

Of course should you develop a new problem your GP can write an emergency referral and we may see you sooner if this is necessary.

Because of the large number of diabetic patients already under follow up and the high rate of new referrals the clinic is fully booked for the next year and there are no routine appointments available in the next 12 months. This means if you miss your diabetic clinic appointment you will not receive a new appointment for another year. **It is therefore very important that you attend all your diabetic appointments and have a blood test and urine test done at Hammersmith Hospital 2 weeks before the appointment.**

If you fail to attend 2 appointments in succession you will be discharged from the clinic and will require a new referral from your GP to be seen again.

If you have any questions please discuss them when you see the doctor in clinic today.

**Anne Dornhorst DM FRCP/ Duncan Bassett MA PhD MRCP**  
**Consultants in Endocrinology and Diabetes**

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Internal Ext: 34823  
Fax: 020 8283 3360  
Appointments 020 8383 5000

Date:  
Ref:

## **HAMMERSMITH HOSPITAL DIABETIC CLINIC**

Dear

### **Thank you for your request for an earlier Diabetic Clinic appointment**

We are very sorry but we are unable to make an earlier appointment. Unfortunately because of the large number of diabetic patients already under follow up and the high rate of new referrals the clinic is fully booked for the next year and there are no routine appointments available in the next 12 months. This means if you have missed a diabetic clinic appointment there is no appointment available until next year.

**It is therefore very important that you attend all your diabetic appointments and have a blood test and urine test done at Hammersmith Hospital 2 weeks before the appointment.**

Of course should you develop an urgent new problem your GP can write an emergency referral and we will see you sooner if this is necessary.

If you have any concerns please discuss these with your GP. During the next few months we would also advise you to see your GP to check your blood pressure, vision, renal function, glucose control (HbA1c) and cholesterol.

We apologise for the inconvenience.

Best wishes,

**Anne Dornhorst DM FRCP/ Duncan Bassett MA PhD MRCP  
Consultants in Endocrinology and Diabetes**

cc GP



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## **Advise to patient discharged for Hammersmith Diabetic Clinic**

**You have been discharged from the diabetic clinic as your diabetes is well controlled.**

**To continue this good control you should be seen by your general practitioner at least once or twice a year.**

**At least once a year you should have your**

**blood pressure checked  
vision tested  
eyes examined  
feet examined  
your weight measured**

**You should have a urine test to check for protein**

**You should have blood tests for**

**Kidney function  
Liver function  
Lipids (Cholesterol)  
HbA1c (long term sugar control)**

**In the future your diabetes may become less well controlled and your GP can of course refer you back to our clinic**

# **Standard Letters for Charing Cross**

## **Area for diabetic clinic W4, W6, W14 and SW6**

<b>New referral out of area letter</b>	<b>80</b>
<b>No diagnostic criteria letter</b>	<b>81</b>
<b>OGTT result letter</b>	<b>82</b>
<b>New patient DNA letter</b>	<b>83</b>
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<b>Discharge letter</b>	<b>87</b>
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Direct Line: 020 8846 1065/1067

Internal Ext: 1065/1067

Fax: 020 8846 1862

Appointments 020 8383 5000

Ref:

Date:

Clinic Date:

### **CHARING CROSS HOSPITAL DIABETIC CLINIC**

GP Address

Dear Dr

RE:

Thank you for your referral letter for this patient with Diabetes. Unfortunately due to the high rate of new local referrals and our follow up clinics being fully booked for more than 1 year, it is difficult for us to see patients from outside our local area. I would be grateful therefore if you could, in the first instance refer this patient to your local Diabetes services.

Our local area includes W4, W6, W14 and SW6.

Kind regards.

Yours sincerely,

**Anne Dornhorst DM FRCP/ Duncan Bassett MA PhD MRCP  
Consultants in Endocrinology and Diabetes**

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Fax: 020 8846 1862

Appointments 020 8383 5000

Ref:

Date:

Clinic Date:

## CHARING CROSS HOSPITAL DIABETIC CLINIC

GP Address

Dear Dr

RE:

Thank you for your referral letter for this patient. Unfortunately your referral contains no diagnostic criteria for diabetes. Due to the high frequency of new diabetic referrals would be grateful if you would confirm the diagnosis prior to referring the patient.

### **Diagnostic Criteria (HBA1c CAN NOT BE USED FOR DIAGNOSIS)**

#### **2x Random Glucose Measurement (1x if symptomatic)**

=11.1 Diabetes Mellitus  
<11.1 Normal if =6.1 do 2x fasting

#### **2x Fasting Glucose Measurement (1x if symptomatic)**

=7.0 Diabetes Mellitus  
=6.1 but <7.0 Impaired Fasting Glucose do OGTT

#### **Oral Glucose Tolerance Test 2h Glucose**

=11.1 Diabetes Mellitus  
=7.8 but <11.1 Impaired Glucose Tolerance  
<7.8 Normal

We are most grateful for your help with this and will make a new patient appointment as soon as we receive the re-referral

Kind regards.

Yours sincerely,

Anne Dornhorst DM FRCP/ Duncan Bassett MA PhD MRCP  
Consultants in Endocrinology and Diabetes

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DATE

**CHARING CROSS HOSPITAL DIABETIC CLINIC**

GP Address:

GP Address Sticker

Re patient:

Patient Sticker

Your patient had an oral glucose tolerance test on .....

The results demonstrate that the patient.....

WHO Diagnostic Criteria

**Fasting Glucose**

=7.0	Diabetes Mellitus
=6.1 but <7.0	Impaired Fasting Glucose
<6.1	Normal

**2h Glucose during Oral Glucose Tolerance Test**

=11.1	Diabetes Mellitus
=7.8 but <11.1	Impaired Glucose Tolerance
<7.8	Normal

If the patient has impaired fasting glucose or impaired glucose tolerance we would recommend life style advice and at least yearly a fasting glucose.

We have not made any further follow up appointment.

Kind regards.

Yours sincerely,

Anne Dornhorst DM FRCP/ Duncan Bassett MA PhD MRCP  
Consultants in Endocrinology and Diabetes

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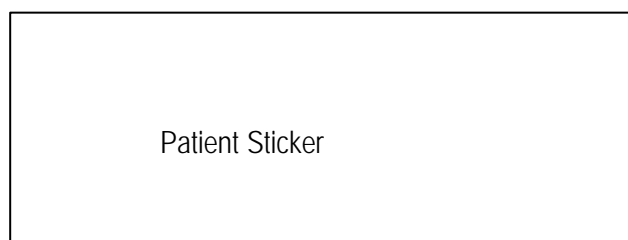
Clinic Date:

### CHARING CROSS HOSPITAL DIABETIC CLINIC

GP Address:



Re patient:



We are sorry that your patient did not attend their appointment at the new diabetic clinic.

Due to the high frequency of new patient referrals we have not made a follow up appointment for your patient. If you wish to re-refer the patient we would be grateful if you could confirm that they are prepared to will attend the hospital diabetic clinic. In addition, we would be grateful if you could and ask them to rearrange the appointment date if they are unable to attend as their non-attendance prevents other urgent patients from being seen.

Kind regards.

Yours sincerely,

**Anne Dornhorst DM FRCP/ Duncan Bassett MA PhD MRCP**  
**Consultants in Endocrinology and Diabetes**



cc Patient

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Internal Ext: 1065/1067

Fax: 020 8846 1862

Appointments 020 8383 5000

Clinic Date:

**CHARING CROSS HOSPITAL DIABETIC CLINIC**

GP Address:



Re patient:



We are sorry to say that your patient has not attended today's clinic appointment and it may be because of transport difficulties to the Charing Cross Hospital diabetic clinic.

If you would like diabetic clinic follow-up for this patient, the patient may be more compliant if you referred them to a more local hospital.

We have not arranged another appointment.

Yours sincerely,

**Anne Dornhorst DM FRCP/ Duncan Bassett MA PhD MRCP  
Consultants in Endocrinology and Diabetes**

cc Patient

Direct Line: 020 8846 1065/1067

Internal Ext: 1065/1067

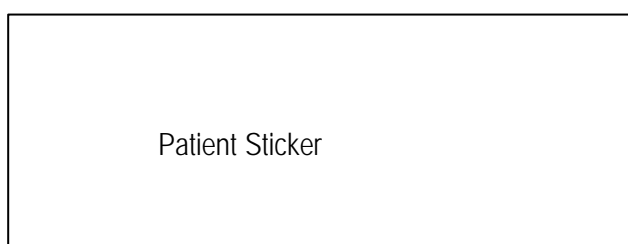
Fax: 020 8846 1862

Appointments 020 8383 5000

Clinic Date:

## CHARING CROSS HOSPITAL DIABETIC CLINIC

Patient:



We are sorry that you did not attend the diabetic clinic today

Because of the large number of diabetic patients already under follow up and the high rate of new referrals the clinic is fully booked for the next year and there are no routine appointments available in the next 12 months. This means if you miss a diabetic clinic appointment you will not receive a new appointment for another year. **It is therefore very important that you attend all your diabetic appointments and have a blood test and urine test done at Charing Cross Hospital 2 weeks before the appointment.**

If you fail to attend 2 appointments in succession you will be discharged from the clinic and will require a new referral from your GP to be seen again.

Of course should you develop a new problem your GP can write an emergency referral and we may see you sooner if this is necessary.

We have given you the next available routine appointment, which will be in about 1 year. The details of which will be sent to you in the next few weeks. During the next few months we would advise you to see your GP to check your blood pressure, vision, renal function, glucose control (HbA1c) and cholesterol.

Yours sincerely,

**Anne Dornhorst DM FRCP/ Duncan Bassett MA PhD MRCP**

## Consultants in Endocrinology and Diabetes

cc GP

Direct Line: 020 8846 1065/1067

Internal Ext: 1065/1067

Fax: 020 8846 1862

Appointments 020 8383 5000

Clinic Date:

**CHARING CROSS HOSPITAL DIABETIC CLINIC**

GP Address:

GP Address Sticker

Re patient:

Patient Sticker

We are sorry to say that your patient has not attended the last 2 diabetic clinic appointments.

It may be that the patient has moved address. If you have a different address and you would still like diabetic clinic follow-up for this patient, please send us another referral with the correct address.

We have not arranged another appointment.

Yours sincerely,

**Anne Dornhorst DM FRCP/ Duncan Bassett MA PhD MRCP**

**Consultants in Endocrinology and Diabetes**

Cc Patient

Direct Line: 020 8846 1065/1067  
Internal Ext: 1065/1067  
Fax: 020 8846 1862  
Appointments 020 8383 5000

Ref:  
Date:  
Clinic Date:

**CHARING CROSS HOSPITAL DIABETIC CLINIC**

GP Address

GP Address

RE:

Patient  
Sticker

Your patient was reviewed in the diabetic clinic today and is considered suitable for discharge from this clinic as they have type II diabetes, are well controlled on diet / sub-maximal oral therapy and have no significant diabetic complications.

**Present Medication**

**Results Date:**

Blood Pressure	(<145/85)
BMI	(<35)
HbA1c	(<8.5%)
Creatinine	(<125mmol/l)
Chol/HDL ratio	(<5.5)
Urine Alb/Cre ratio	(<3.0)
Acuity	Right: Left:

We have advised your patient that they should be seen at least annually in the practice. Annual review should include a minimum of blood pressure measurement, examination of the feet for peripheral pulses and evidence of peripheral neuropathy. Visual Acuity and dilated funduscopy should be performed annually either in the practice, by an optician or at the Hammersmith retinal screening clinic. Blood test should include renal function, full fasting lipid profile including HDL and LDL, HbA1c and liver function and CK where appropriate. Urine testing should include urine dipstick analysis and urine albumin/creatinine ratio.

We will be happy to see your patient again in the Hammersmith diabetic clinic should they develop diabetic complications or their control fall outside the above ranges.

Kind regards.

Yours sincerely,

**Anne Dornhorst DM FRCP/ Duncan Bassett MA PhD MRCP**



## **Consultants in Endocrinology and Diabetes**

Cc Dr Andrew Western

Cc The Patient

Direct Line: 020 8846 1065/1067  
Internal Ext: 1065/1067  
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Appointments 020 8383 5000

## CHARING CROSS HOSPITAL DIABETIC CLINIC

### To all patients attending Charing Cross diabetic follow up clinic

Dear Sir/Madam,

This letter is just to inform you about our patient appointments in the Hammersmith diabetic clinic and to help us give a better service to you and all our diabetic patients.

With the present number of staff we are only able to see diabetic patients once per year unless they have diabetic complication affecting their eyes or kidneys when they will be seen every six months.

Of course should you develop a new problem your GP can write an emergency referral and we may see you sooner if this is necessary.

Because of the large number of diabetic patients already under follow up and the high rate of new referrals the clinic is fully booked for the next year and there are no routine appointments available in the next 12 months. This means if you miss your diabetic clinic appointment you will not receive a new appointment for another year. **It is therefore very important that you attend all your diabetic appointments and have a blood test and urine test done at Charing Cross Hospital 2 weeks before the appointment.**

If you fail to attend 2 appointments in succession you will be discharged from the clinic and will require a new referral from your GP to be seen again.

If you have any questions please discuss them when you see the doctor in clinic today.

**Anne Dornhorst DM FRCP/ Duncan Bassett MA PhD MRCP**  
Consultants in Endocrinology and Diabetes

Direct Line: 020 8846 1065/1067

Internal Ext: 1065/1067

Fax: 020 8846 1862

Appointments 020 8383 5000

Date:

Ref:

## CHARING CROSS HOSPITAL DIABETIC CLINIC

Dear

### **Thank you for your request for an earlier Diabetic Clinic appointment**

We are very sorry but we are unable to make an earlier appointment. Unfortunately because of the large number of diabetic patients already under follow up and the high rate of new referrals the clinic is fully booked for the next year and there are no routine appointments available in the next 12 months. This means if you have missed a diabetic clinic appointment there is no appointment available until next year.

**It is therefore very important that you attend all your diabetic appointments and have a blood test and urine test done at Hammersmith Hospital 2 weeks before the appointment.**

Of course should you develop an urgent new problem your GP can write an emergency referral and we will see you sooner if this is necessary.

If you have any concerns please discuss these with your GP. During the next few months we would also advise you to see your GP to check your blood pressure, vision, renal function, glucose control (HbA1c) and cholesterol.

We apologise for the inconvenience.

Best wishes,

**Anne Dornhorst DM FRCP/ Duncan Bassett MA PhD MRCP  
Consultants in Endocrinology and Diabetes**

cc GP

Direct Line: 020 8846 1065/1067

Internal Ext: 1065/1067

Fax: 020 8846 1862

Appointments 020 8383 5000

## **Advise to patient discharged for Charing Cross diabetic clinic**

**You have been discharged from the diabetic clinic as your diabetes is well controlled.**

**To continue this good control you should be seen by your general practitioner at least once or twice a year.**

**At least once a year you should have your**

**blood pressure checked**

**vision tested**

**eyes examined**

**feet examined**

**your weight measured**

**You should have a urine test to check for protein**

**You should have blood tests for**

**Kidney function**

**Liver function**

**Lipids (Cholesterol)**

**HbA1c (long term sugar control)**

**In the future your diabetes may become less well controlled and your GP can of course refer you back to our clinic**

# Appendix II

# HbA<sub>1c</sub>: Method & Interferences

HbA<sub>1c</sub> is formed by non-enzymatic glycosylation of the N-terminal valine of the  $\beta$  chain of haemoglobin.

HbA<sub>1c</sub> is measured in the Hammersmith by **HPLC-boronate affinity chromatography**.

This method is **not** affected by:

- Fetal haemoglobin (HbF)
- Haemoglobin variants (HbS, HbC, HbD)
- Carbamylated Hb (such as in uraemia)

However, it must be remembered that if any of these conditions result in increased red cell turnover, then although there is not direct analytical interference, the shortened red cell (Hb) half life will result in a reduction in HbA<sub>1c</sub>.

This method has been shown to have good within-laboratory precision, although it may be less satisfactory between laboratories.

I don't have specific information on the thalassemia traits, although as the rate of red cell turnover in these patients is pretty constant then it is presumably reasonable to compare within patient HbA<sub>1c</sub> results.

# **Appendix IV**



# THE HAMMERSMITH DIABETIC CLINIC

**CONSULTANTS**  
 Professor Steve Bloom  
 Dr Anne Dornhorst  
 Dr Duncan Bassett  
 Dr Dominic Withers  
 Dr Karim Meeran  
 Dr Jeannie Todd

**CLINICAL ASSISTANT**  
 Dr Shahenaz Walji

**DIABETIC NURSES**  
 Debbie Lake  
 Marie O'Conner  
 Jane Ormond  
 Ursula Kirwan  
 Molly Nanka-Bruce

**DIETITIANS**  
 Dr Gary Frost  
 Linda Carter  
**PODIATRISTS**  
 Trusha Patel  
 Annabel Trimble



Endocrine Secretaries Tel: 020

Appointments Office Fax: 020 8383 3360  
 Tel: 020 8383 5000  
 Fax: 020 8383 8383

Diabetic Nurse Specialists Tel: 020 8383 4693  
 24h Emergency Service Tel: 020 8383 1000  
 Ask for Endocrine SpR

Dear Dr \_\_\_\_\_

DATE \_\_\_\_\_ 2002

Your patient was reviewed in the diabetic clinic today.

DIAGNOSES	AGE:
1. Type ___ DM;	Years diagnosed ___ y
2.	
3.	
4.	
5.	
6.	
SMOKING ( YES NO ):	

PRESENT MEDICATIONS	
1.	8.
2.	9.
3.	10.
4.	11.
5.	12.
6.	13.
7.	14.

RESULTS (date):	BM Glucose	Kg	Kg/m <sup>2</sup>
Wt change and BMI (<27)			
HbA1c (<8.0%)			%
Creatinine (<120mmol/l)			mmol/l
Urine Alb/Cre (2)			
Urine Dipstick (NAD)			
Cholesterol (<5.0mmol/l)			mmol/l
Triglycerides (<2.3 mmol/l)			mmol/l
HDL (>1.0mmol/l)			mmol/l
LDL (<3.5mmol/l)			mmol/l
Chol/HDL ratio (<5.0)			
LFTs (if on statin)			
CK (statin/fibrate) (<200)			U/l
FT4 (9-26pmol/l)			pmol/l
TSH (0.3-4.2mU/l)			mU/l

EXAMINATION	Right	Left
BP (<140/80)		Standing
Tropicamide		
Pin Hole Acuity		
Retinopathy		
Eye - Other		
Reflexes KJ:		
AJ:		
Sensation VS:		
10g microfilament:		
PP:		
Pulses DP:		
PT:		
Feet		
Injection sites		

**NEW EVENTS**

- HBGM/Hypoglycaemia
- Diet / Exercise
- Chest pain / Claudication
- Change in vision
- Neuropathic pain
- Foot Infection/Ulcer
- Pregnancy plans
- Erectile dysfunction

Yours sincerely

Dr \_\_\_\_\_  
 Dr

**PLAN OF ACTION**

( SHO / SpR / Associate Specialist / Consultant )      Next Review      Weeks/Month      see DSN comment

# THE CHARING CROSS DIABETIC CLINIC



CONSULTANTS

Professor Steve Bloom  
 Dr Anne Dornhorst  
 Dr Duncan Bassett  
 Dr Dominic Withers  
 Dr Karim Meeran  
 Dr Jeannie Todd

CONSULTANTS

Dr Stuart McHardy Young

DIABETIC NURSES

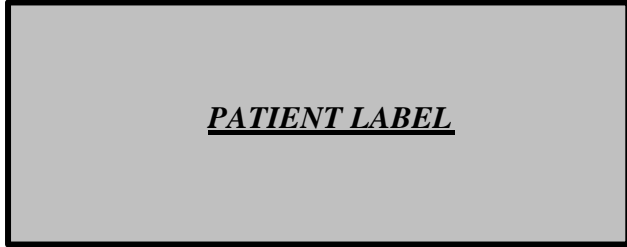
Debbie Lake  
 Debbie Hutchins  
 Brenda Lawrence  
 Ursula Kirwan

DIETITIANS

Dr Gary Frost  
 Linda Carter

PODIATRISTS

Trusha Patel  
 Annabel Trimble



PATIENT LABEL

Endocrine Secretaries      Tel: 020 8846 1065  
    Fax: 020 8846 1080  
 Appointments Office      Tel: 020 8383 5000  
    Fax: 020 8346 7564  
 Diabetic Nurse Specialists      Tel: 020 8846 1062  
 24h Emergency Service      Tel: 020 8383 1000  
 Ask for Endocrine SpR

DATE \_\_\_\_\_ 2002

Dear Dr \_\_\_\_\_  
 Your patient was reviewed in the Diabetic Clinic today.

**DIAGNOSES      AGE:**

1. Type ___ DM;      Years diagnosed ____ y
2.
3.
4.
5.
6.
SMOKING YES NO

**MEDICATION**

1.	8.
2.	9.
3.	10.
4.	11.
5.	12.
6.	13.
7.	14.

**RESULTS (date):      BM Glucose**

Wt change and BMI (<27)	Kg	Kg/m <sup>2</sup>
HbA1c (<8.0%)		%
Creatinine (<120mmol/l)		mmol/l
Urine Alb/Cre (2)		
Urine Dipstick (NAD)		
Cholesterol (<5.0mmol/l)		mmol/l
Triglycerides (<2.3 mmol/l)		mmol/l
HDL (>1.0mmol/l)		mmol/l
LDL (<3.5mmol/l)		mmol/l
Chol/HDL ratio (<5.0)		
LFTs (if on statin)		
CK (statin/fibrate) (<200)		U/l
FT4 (9-26pmol/l)		pmol/l
TSH (0.3-4.2mU/l)		mU/l

EXAMINATION	Right	Left
BP (<140/80)		Standing
Tropicamide		
PH Acuity		
Retinopathy		
Eye - Other		
Reflexes      KJ:		
AJ:		
Sensation      VS:		
10g microfilament:		
PP:		
Pulses      DP:		
PT:		
Feet		
Injection sites		

NEW EVENTS

HBGM / Hypoglycaemia  
 Diet / Exercise  
 Chest pain / Claudication  
 Change in vision  
 Neuropathic pain / Foot Infection/Ulcer  
 Pregnancy plans  
 Erectile dysfunction

Yours sincerely

Dr \_\_\_\_\_  
 Dr \_\_\_\_\_

PLAN OF ACTION

Next Review \_\_\_\_\_ Weeks/Month \_\_\_\_\_  
 ( SHO / SpR / Associate Specialist / Consultant )      see DSN comment

