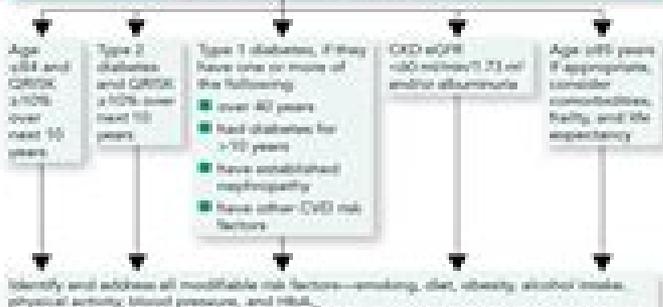


INITIAL CONSIDERATIONS

Measure non-fasting full lipid profile (TC, HDL-C, non-HDL-C, triglycerides and HbA_{1c}) as part of an initial baseline assessment. Consider secondary causes of hyperlipidaemia and manage as needed. Ensure appropriate baseline and follow-up tests as detailed in the guideline. Measure BMI. Identify and exclude people with contraindications/drug interactions. If non-fasting triglyceride above 4.5 mmol/l see the guideline text.

PRIMARY PREVENTION

Consider statin therapy for adults who do not have established CVD but fit one of the categories below. Use QRISK risk assessment tool where appropriate (see Primary prevention risk assessment in the guideline).



Consider additional risk factors, if present, together with QRISK score (raised for FH, severe mental illness, taking medicines that cause dyslipidaemia, systemic inflammatory disorder (e.g. SLE), impaired fasting glycaemia, recent change in risk factors).

PRIMARY PREVENTION

Consider statin therapy in individuals at intermediate or lower risk (see below): **Atorvastatin 20 mg OD**

- Measure full lipid profile again after 3 months (non-fasting)
- High-intensity statin treatment should achieve reduction of non-HDL-C >40% from baseline. If not achieved after 3 months:
 - discuss treatment adherence, timing of dose, diet, and lifestyle
 - If at higher risk based on comorbidities, risk score or clinical judgement—see Additional risk factors—consider increasing the dose every 2–3 months up to a maximum dose of atorvastatin 80 mg OD^a
 - for how to increase in people with CKD, see the section Special patient populations

- If patients on a high-intensity statin have side effects, offer a lower dose or an alternative statin (see Statin intensity table)
- If maximum tolerated dose of statin does not achieve non-HDL-C reduction >40% of baseline value after 3 months, consider adding ezetimibe 10 mg OD
- If recommended statin treatment is contraindicated or not tolerated:
 - see AAC (Statin Intolerance Algorithm)^a for advice regarding adverse effects
 - ezetimibe 10-mg monotherapy may be considered. Assess response after 3 months
 - ezetimibe 10-mg/rosuvastatin acid 180-mg combination may be considered when ezetimibe alone does not control non-HDL-C/DL-C well enough (NICE TA696)

If non-HDL-C reduction remains <40% of baseline (despite maximal tolerated lipid-lowering therapy (including people with intolerance and contraindications)) consider referral to specialist lipid management clinic according to local arrangements

SEVERE HYPERLIPIDAEMIA

If TC >7.5 mmol/l and/or LDL-C >4.9 mmol/l and/or non-HDL-C >5.9 mmol/l, a personal or/and family history of (premature) CVD (with/without and with no secondary causes, suggest FH (possible heterozygous FH). Do not use QRISK risk assessment tool.

DIAGNOSIS AND REFERRAL

Take fasting blood for repeat lipid profile to measure LDL-C. Use the **Simon Broome or Dutch Lipid Clinic Network** criteria to make a clinical diagnosis of FH. Refer to lipid clinic for further assessment if clinical diagnosis of FH or if TC >10 mmol/l and/or LDL-C >4.9 mmol/l and/or non-HDL-C >7.5 mmol/l or fasting triglycerides >10 mmol/l (regardless of family history).

TREATMENT TARGETS IN FH

If clinical diagnosis of FH and/or other risk factors present, follow the recommended treatment management pathway for primary or secondary prevention as for non-FH, **BUT** aim to achieve at least a 50% reduction of LDL-C (or non-fasting non-HDL-C) from baseline.

Consider specialist referral for further treatment and/or consideration of PCSK9 therapy if:

- they are assessed to be at very high risk of a coronary event^a
 - all therapy is not tolerated
 - or LDL-C remains >5 mmol/l (primary prevention)
 - or LDL-C remains >2.5 mmol/l (secondary prevention) despite maximal tolerated statin and ezetimibe therapy.
- ^adefined as any of the following:
- established CVD
 - two or more other CVR risk factors

SECONDARY PREVENTION

Offer statin therapy to adults with CVD, including angina, previous MI, heart failure, stroke, or TIA, atherosclerotic peripheral arterial disease, to not doing so will increase risk of a person has acute coronary syndrome. Take a lipid profile on admission (within 24 hours).

Identify and address all modifiable risk factors—smoking, diet, obesity, alcohol intake, physical activity, blood pressure, and HbA_{1c}.

SECONDARY PREVENTION

On secondary prevention in secondary prevention when managing modifiable risk factors. Prescribe a high-intensity statin: **Atorvastatin 80 mg OD**. Use a lower dose of atorvastatin if there is a previous drug interaction, high risk of an exacerbating adverse effect, or patient preference. (see assessment of drug-drug interactions with CVD and comorbidities).

- Measure full lipid profile again after 3 months (non-fasting)
 - High-intensity statin treatment should achieve reduction of non-HDL-C >40% from baseline. If not achieved after 3 months:
 - discuss treatment adherence, timing of dose, diet, and lifestyle measures
 - If started on less than atorvastatin 80 mg and the person is judged to be at higher risk (based on comorbidities, risk score or clinical judgement—see Additional risk factors), consider increasing to 80 mg atorvastatin. For how to increase in people with CKD see Special patient populations
 - If non-HDL-C baseline value is not available^a, consider target non-HDL-C <2.5 mmol/l (approximately equivalent to LDL-C <1.8 mmol/l) as recommended by Joint British Societies
 - If patients on a high-intensity statin have side effects, offer a lower dose or an alternative statin (see Table of lipid lowering with available therapies)
- ^athe extent is not covered by NICE OD10^a

If maximum tolerated dose of statin does not control non-HDL-C/DL-C well enough after 3 months, confirm statin adherence then consider the following options (based on shared decision-making^a with the patient).

^aSee the full guideline for information to support shared decision-making

If recommended statin treatment is contraindicated or not tolerated, follow the AAC (Statin Intolerance Algorithm)^a for advice regarding adverse effects

- If statin intolerance is confirmed, consider:
 - ezetimibe 10-mg monotherapy. Assess response after 3 months (NICE TA696)
 - ezetimibe 10-mg/rosuvastatin acid 180-mg combination when ezetimibe alone does not control non-HDL-C sufficiently (NICE TA696)

Ezetimibe 10-mg daily (NICE TA696) reassess after 3 months. If non-HDL-C remains >2.5 mmol/l, consider injectable therapies, arrange a fasting blood test, and assess eligibility (see below)

If non-HDL-C remains >2.5 mmol/l despite either lipid lowering therapies, consider injectable therapies—arrange a fasting blood test and assess eligibility (see below) (TA696, TA719)

Injectable therapies^a
If non-HDL-C >2.5 mmol/l, arrange fasting blood test to measure LDL-C to assess eligibility:

- inclisiran**—if fasting LDL-C <0.6 mmol/l, despite maximum tolerated lipid lowering therapy (TA718)
- or **PCSK9i**—see below (if extent of lipid lowering with available therapies for LDL-C threshold (TA697-9)

If eligibility criteria are not met, consider **ezetimibe 10-mg daily** (if not previously considered). ^ainclisiran and PCSK9i should not be prescribed concurrently.

[A] England: this advice publication relates to the intolerance pathway

BMI=body mass index; CVD=cardiovascular disease; CKD=chronic kidney disease; CVD=cardiovascular disease; eGFR=estimated glomerular filtration rate; FH=familial hypercholesterolaemia; HbA_{1c}=haemoglobin A_{1c}; HDL-C=high-density lipoprotein cholesterol; LDL-C=low-density lipoprotein cholesterol; MI=acute myocardial infarction; OD=once daily; PCSK9i=proprotein convertase subtilisin/kexin 1 inhibitor; SLE=systemic lupus erythematosus; TC=total cholesterol; TIA=transient ischaemic attack

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Sujoy Ghosh, Banshi Saboo



New Lipid Guidelines Summary:

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