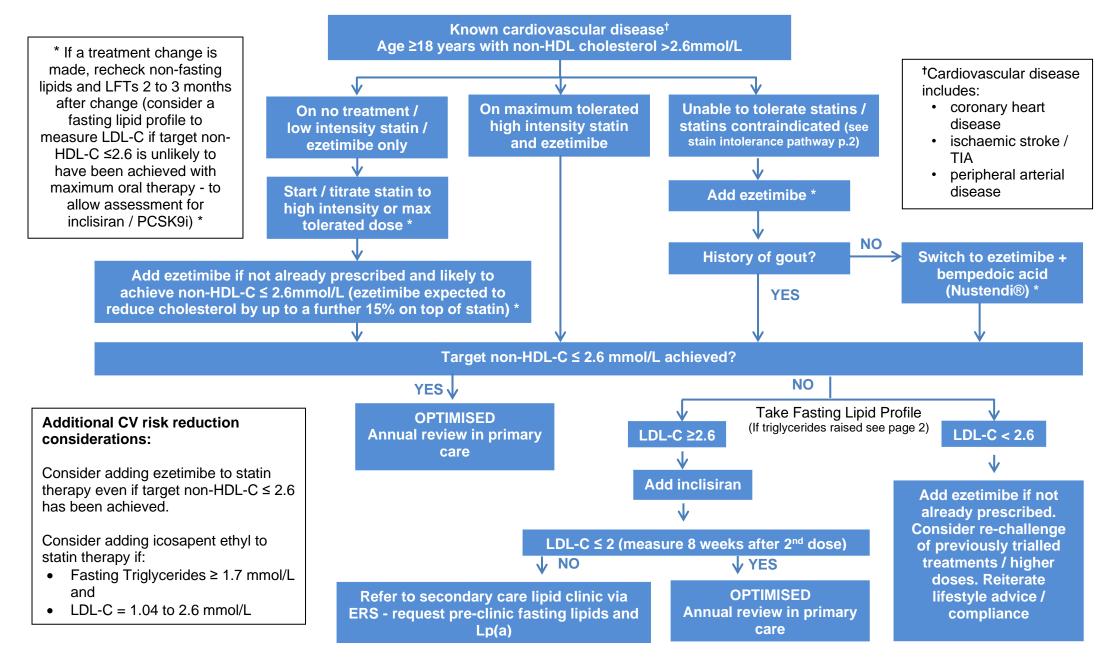


Gloucestershire Secondary Prevention Lipid Management Pathway



Statins

High intensity statins (43 to 55% LDL-C reduction):

- Atorvastatin 20mg-80mg
- Rosuvastatin 10mg-40mg

Start with atorvastatin 80mg (or rosuvastatin 20mg if atorvastatin unsuitable) unless there is a reason to start with a lower dose e.g.:

- Interaction with other drugs
- High risk of adverse effects
- The person would prefer to take a lower dose

If unable to tolerate rosuvastatin:

Start rosuvastatin 5mg once weekly titrating up over 3-4 weeks to twice weekly then alternate days then daily.

If side effects develop then reduce to maximum tolerated dose See also the <u>Statin intolerance pathway</u>

Ezetimibe <u>NICE TA385</u> (further 10 to 15% LDL-C reduction than statin monotherapy) If target not met on maximum tolerated statin (or statins not tolerated): Prescribe ezetimibe 10mg od. Consider adding ezetimibe to statin therapy even if target achieved.

Bempedoic Acid <u>NICE TA694</u> (22 to 33% LDL-C reduction with ezetimibe) Bempedoic acid with ezetimibe is recommended if statins are contraindicated or not tolerated, and ezetimibe alone is insufficient.

Prescribe: Nustendi® (bempedoic acid 180mg + ezetimibe 10mg) 1 tablet od. Treatment with Nustendi® should be discontinued if hyperuricaemia accompanied with symptoms of gout appear.

Inclisiran NICE TA733 (further 48 to 52% LDL-C reduction than maximum tolerated oral therapy)

See local information sheet for practical guidance.

Fibrates

If patient already taking a fibrate (fenofibrate / bezafibrate) apply caution in using statin, use lowest dose atorvastatin (risk of myopathy)

PCSK9 inhibitors <u>NICE TA393</u> / <u>NICE TA394</u> Secondary Care Prescribing only.

Icosapent ethyl <u>NICE TA805</u>

Prescribe: Two 998mg capsules bd (contraindicated in peanut/soya allergy) Higher risk of bleeding in patients taking concomitant antithrombotics. Consider patient preferences, comorbidities, polypharmacy, general frailty and life expectancy.

Also consider principles of shared care decision making <u>HERE</u>. Ensure that modifiable <u>risk factors</u> other than cholesterol are also addressed.

Chronic Kidney Disease:

| L | | |
|---|------------------|--|
| | Atorvastatin: | No dose reduction required |
| | Rosuvastatin: | If eGFR 30-59 mL/min/1.73m ² , max dose = 20mg. |
| | | If eGFR <30 mL/min/1.73m ² , max dose = 5-10mg (<i>Renal</i> |
| | | Handbook, off label). |
| | Ezetimibe: | No dose reduction required |
| | Bempedoic acid: | Limited experience with if eGFR <30 mL/min/1.73m ² |
| | Inclisiran: | No dose reduction required |
| | | (avoid haemodialysis within 72 hours of inclisiran dose) |
| | Icosapent ethyl: | No dose reduction required. |
| | | |

Drug Interactions: Patients prescribed warfarin will need regular INR monitoring with statin dose change. Caution when using statins with amiodarone, verapamil, diltiazem, HIV protease inhibitors – use lowest dose atorvastatin.

Annual Review

- Adherence to medication
- Lifestyle / Diet
- CVD risk factors
- Annual lipids check
- Check LFTs if statin started in previous 12 months
- Patient Resources the Heart UK <u>website</u> contains a number of useful resources for patients

Triglycerides

| Triglyceride concentration | Action |
|----------------------------|---|
| >20 mmol/L | Refer to Lipid Clinic for urgent specialist review |
| 10 – 20 mmol/L | Take fasting TG (if not already done) after 5 to 14 days. |
| | Refer if TG remains >10mmol. Risk of acute pancreatitis. |
| 4.5 – 9.9 mmol/L | Take fasting TG (if not already done). Optimise CVD risk |
| | factors. Refer if TG remains >4.5 and non-HDL-C > 7.5. |

CVD Primary Prevention Guideline click here

Familial Hypercholesterolaemia (FH) Pathway click here