

CVD Prevention National Picture & Strategy

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SRO CVD Prevention.

Long Term Plan – the NHS in 10 years 2019-2029





The NHS Long Term Plan

Prevention and Health Inequalities

CVD Prevention

- Prevent 150,000 heart attacks, strokes and dementia cases
- Provide education and exercise programmes to tens of thousands more patients with heart problems, preventing up to 14,000 premature deaths



CVDP Strategy



To do the basics correctly -

Good Processes
Good Guidelines



Prevention Everywhere



The Role of the NHS in CVD Prevention?

Chris Whitty (Chief Medical Officer)

Government should be putting more money into prevention (https://www.youtube.com/watch?v=1hj9yBzC2-0)

NHS delivers CVD -- Prevention?

- Tobacco support
- Weight Management
- Alcohol
- CVD : ABC Atrial Fibrillation/ Blood pressure / Cholesterol





CVD Prevention - Strategy Clinical Leadership is Key



Development of CVD Prevention Clinical Leadership at all levels



Data driven decisions



CVD Prevention
Communications Strategy
(Public and Professional)



CVD Prevention Meetings and Educational Events (Webinars, Stroke Network Education Event, wider events) Good Guidelines



Bringing together departments & organisations to embed the prevention agenda

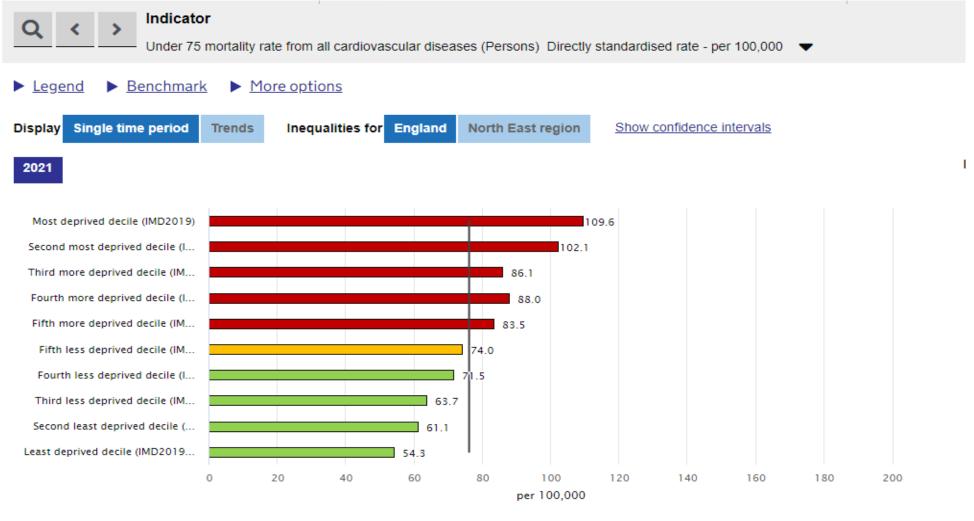
NHS & local authorities working as one



Embracing new technology and new models of care Embracing innovation



Health Inequalities





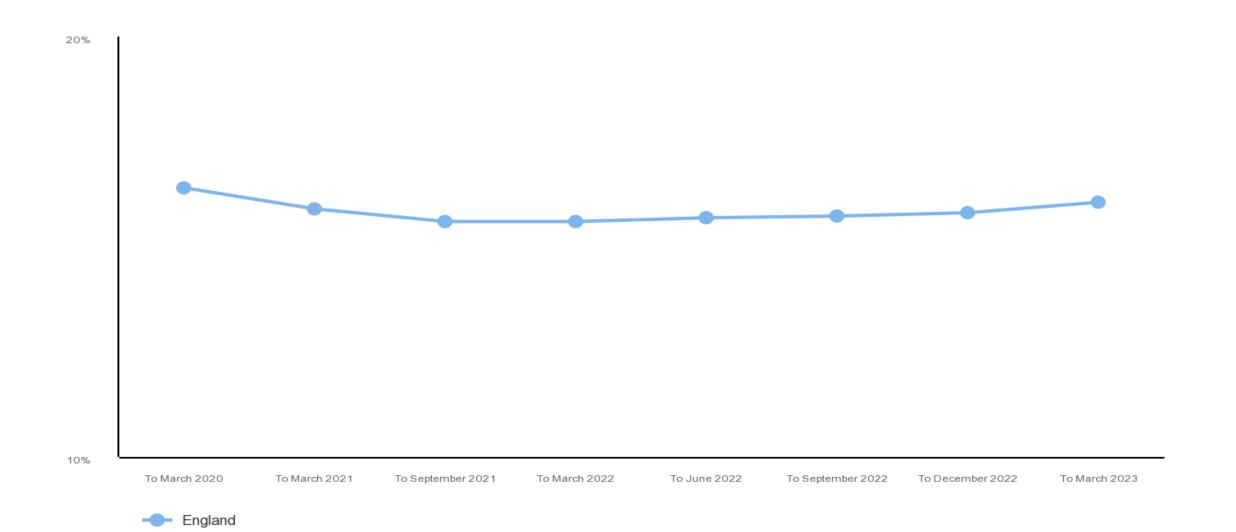
Blood Pressure --

 HYP008. The percentage of patients aged 79 years or under with hypertension in whom the last blood pressure reading (measured in the preceding 12 months) is 140/90 mmHg or less (or equivalent home blood pressure reading)

• HYP009. The percentage of patients aged 80 years or over, with hypertension, in whom the last blood pressure reading (measured in the preceding 12 months) is 150/90 mmHg or less, (or equivalent home blood pressure reading)

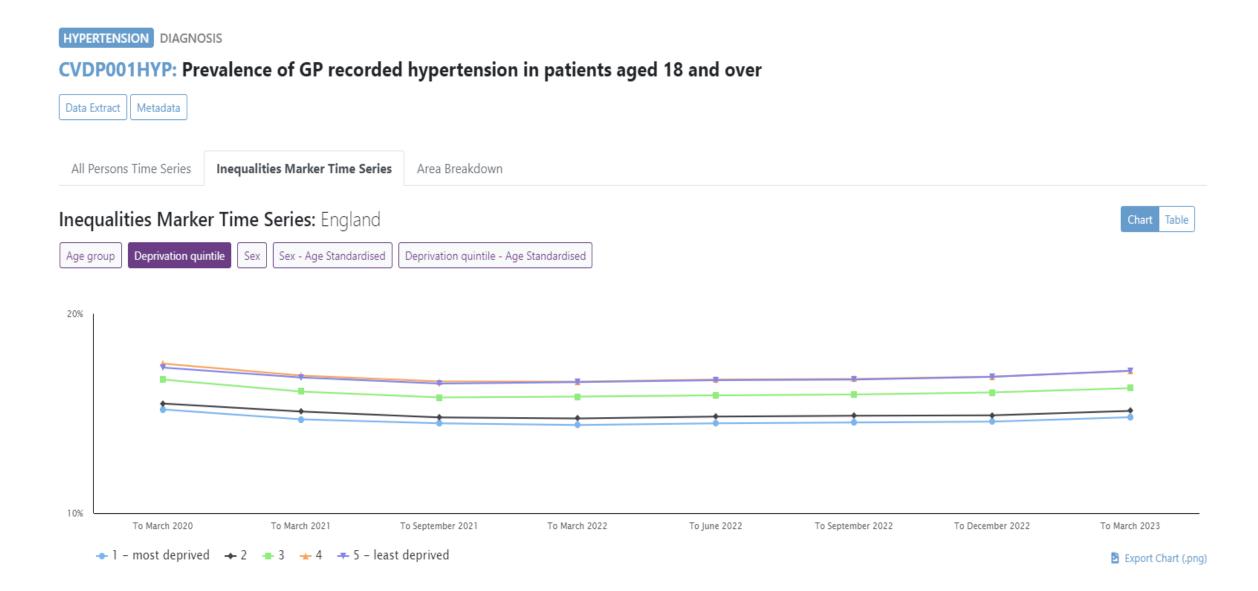
Prevalence of Hypertension – 16.6%





More hypertension in Least Deprived



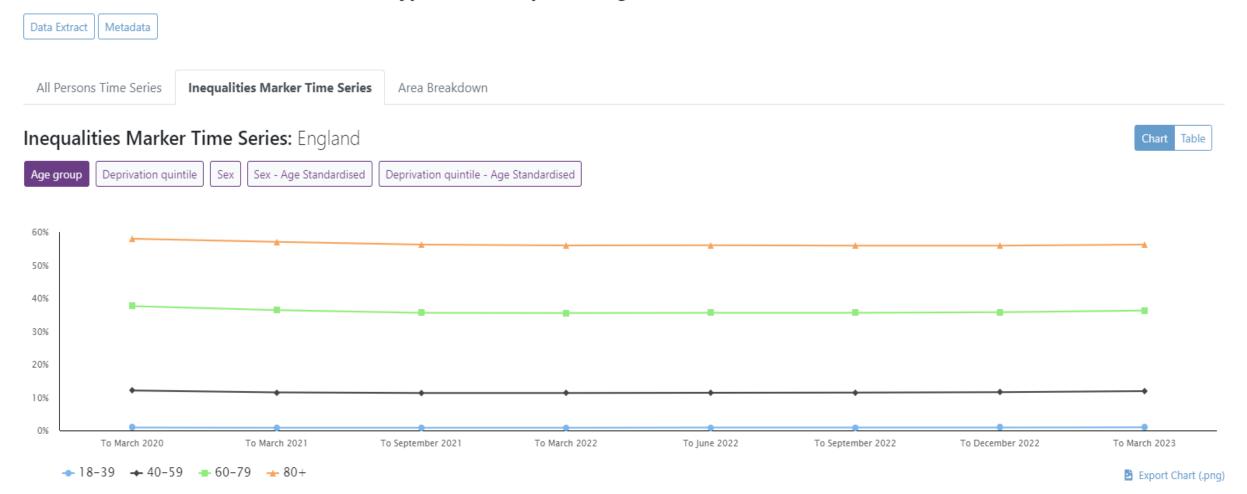




More Hypertension in Elderly Groups

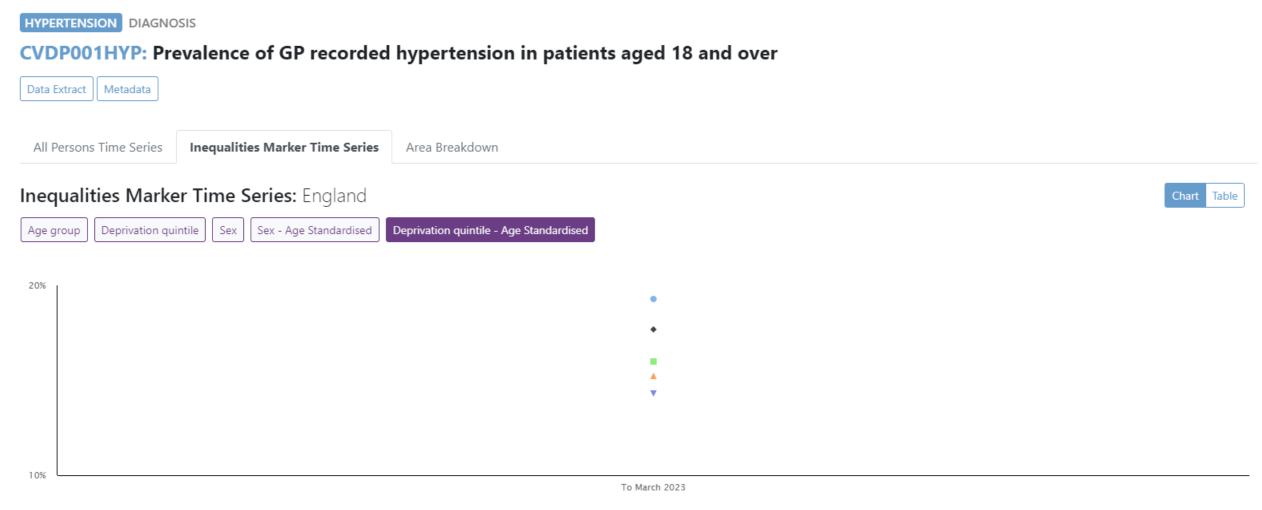
HYPERTENSION DIAGNOSIS

CVDP001HYP: Prevalence of GP recorded hypertension in patients aged 18 and over



Age Standardised – More disease in the deprived areas

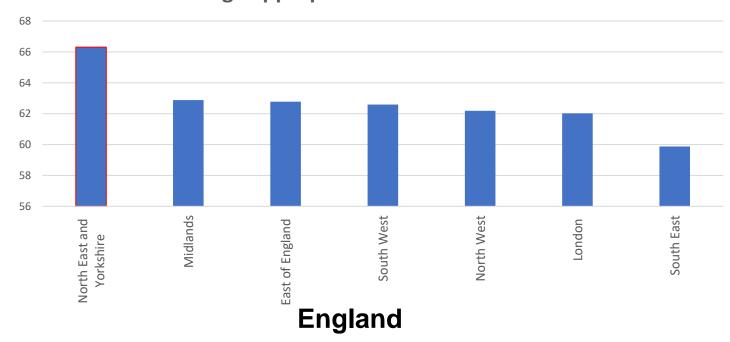




Current % of patients with hypertension treated to target



Percentage of patients aged 18 and over, with GP recorded hypertension, in whom the last blood pressure reading (measured in the preceding 12 months) is below the age appropriate treatment threshold





NEY Breakdown

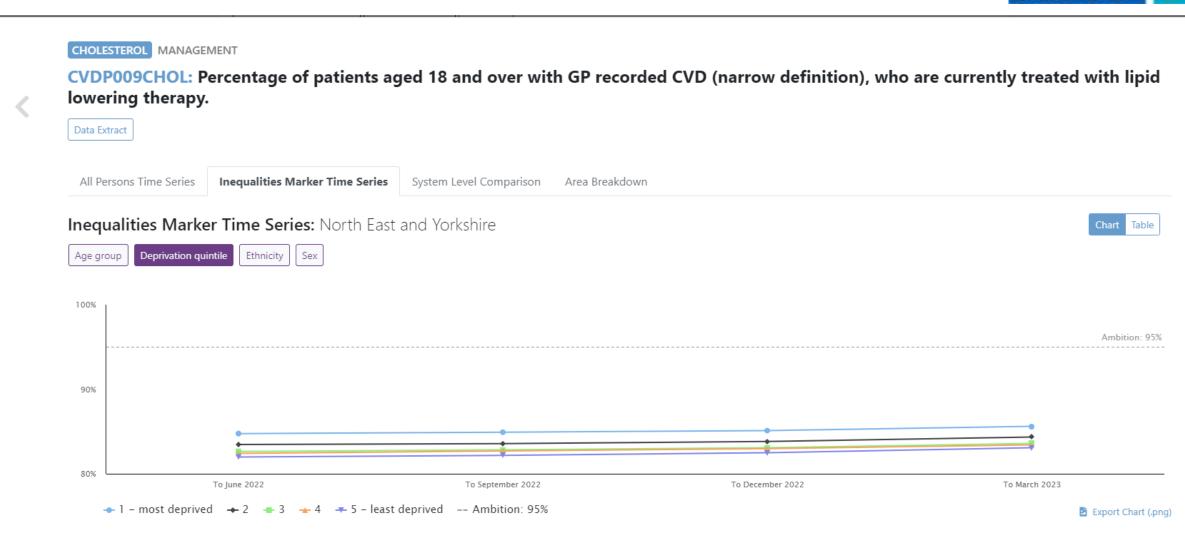


QOF – Lipids

- CHOL001. Percentage of patients on the QOF Coronary Heart Disease, Peripheral Arterial Disease, Stroke/TIA or Chronic Kidney Disease Register who are currently prescribed a statin, or where a statin is declined or clinically unsuitable, another lipid-lowering therapy 70-95% (14)
- CHOL002. Percentage of patients on the QOF Coronary Heart Disease, Peripheral Arterial Disease, or Stroke/TIA Register, who have a recording of non-HDL cholesterol in the preceding 12 months that is lower than 2.5 mmol/L, or where non-HDL cholesterol is not recorded a recording of LDL cholesterol in the preceding 12 months that is lower than 1.8 mmol/L 20-35% (16)

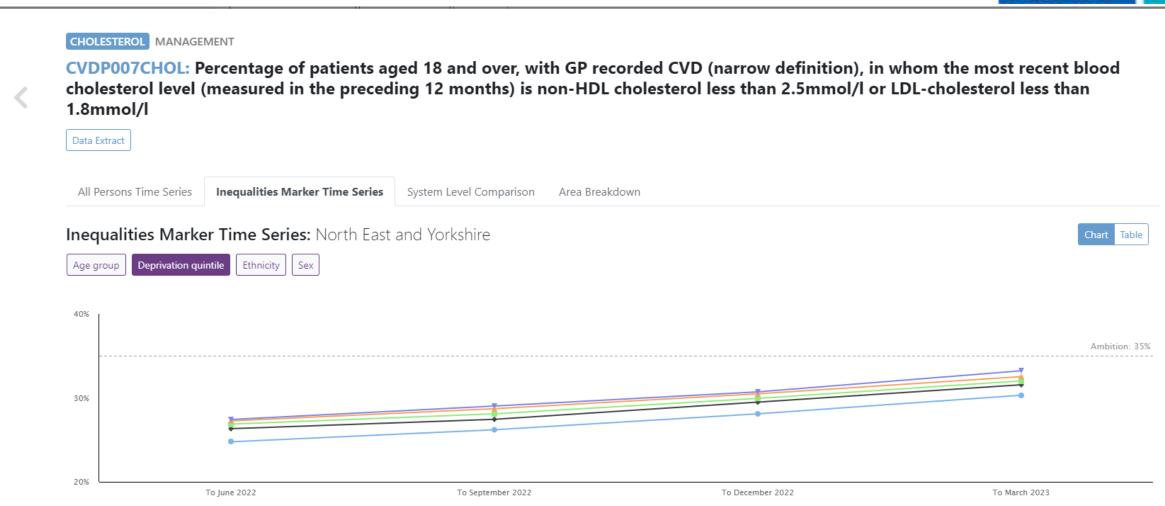


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Export Chart (.png)



→ 1 - most deprived → 2 → 3 → 4 → 5 - least deprived -- Ambition: 35%

Summary of National Guidance for Lipid Management for Primary and Secondary Prevention of CVD



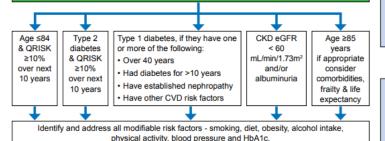


INITIAL CONSIDERATIONS:

- Measure non-fasting full lipid profile (total cholesterol, HDL-C, non-HDL-C, triglycerides) and HbA1c 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 on page 2. Measure BMI. Identify and exclude people with contraindications/drug interactions If non-fasting triglyceride above 4.5mmol/L see page 2.

PRIMARY PREVENTION

Consider statin therapy for adults who do not have established CVD but fall into the categories below. Use QRISK risk assessment tool where appropriate (see page 2, 'Primary Prevention Risk Assessment')



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

PRIMARY PREVENTION

If lifestyle modification is ineffective or inappropriate offer statin treatment. **Atorvastatin 20mg daily**

- · 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 page 2
 'Additional Risk Factors') consider increasing the dose every 2-3 months up to a maximum
 dose of atorvastatin 80mg daily.
- For how to increase in people with CKD see 'Special Patient Populations' (page 2).
- If patients on a high-intensity statin have side effects, offer a lower dose or an alternative statin (see page 2 'Extent of lipid lowering with available therapies')
- If maximum tolerated dose of statin does not achieve non-HDL-C reduction > 40% of baseline value after 3 months consider adding Ezetimibe 10mg daily (NICE TA385)
- If statin treatment is contraindicated or not tolerated;
- See AAC Statin Intolerance Algorithm for advice regarding adverse effects (click here)
- Ezetimibe 10mg monotherapy may be considered. Assess response after 3 months.
- Ezetimibe 10mg/bempedoic acid 180 mg combination may be considered when ezetimibe alone does not control non-HDL-C/LDL-C well enough (NICE TA694).

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

SEVERE HYPERLIPIDAEMIA

If TC>7.5mmol/L and/or LDL-C >4.9mmol/L and/or non-HDL-C >5.9mmol/L, a personal and/or family history of confirmed CHD (<60 years) and with no secondary causes: suspect familial hypercholesterolaemia (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 (DLCN) criteria to make a clinical diagnosis of FH.

Refer to Lipid Clinic for further assessment if clinical diagnosis of FH or if TC>9.0mmol/L and/or LDL-C >6.5mmol/L and/or non-HDL-C >7.5mmol/L or Fasting triglycerides > 10mmol/L (regardless of family history) (page 2)

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 PCSK9i therapy IF

- they are assessed to be at very high risk of a coronary event**
- OR therapy is not tolerated
- OR LDL-C remains >5mmol/L (primary prevention)
- OR LDL-C remains >3.5mmol/L (secondary prevention)

despite maximal tolerated statin and ezetimibe therapy.

- *defined as any of the following:
- Established coronary heart disease
 Two or more other CVD risk factors

SECONDARY PREVENTION

Offer statin therapy to adults with CVD, this includes CHD, angina, Acute Coronary Syndrome (MI or unstable angina), revascularisation, stroke or TIA, or symptomatic peripheral arterial disease. Do not delay statin treatment if a person has acute coronary syndrome. Take a lipid sample on admission (within 24 hours).

Identify and address all modifiable risk factors - smoking, diet, obesity, alcohol intake, physical activity, blood pressure and HbA1c.

SECONDARY PREVENTION

Do not delay statin treatment in secondary prevention while managing modifiable risk factors.

Prescribe a high intensity statin:

Atorvastatin 80mg daily

Use a lower dose of atorvastatin if there is a potential drug interaction, high risk of or experiencing adverse effects, or patient preference.

Offer atorvastatin 20mg if CKD (people with GFR< 60 mL/min/1.73m²).

- 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 80mg and the person is judged to be at higher risk (based on comorbidities, risk score or clinical judgement - see page 2 'Additional Risk Factors'), consider increasing to 80mg atorvastatin. For how to increase in people with CKD see 'Special Patient Populations' (page 2).
- If non-HDL-C baseline value is not available*, consider target non-HDL-C < 2.5mmol/L (approximately
 equivalent to LDL-C < 1.8mmol/L) as recommended by Joint British Societies (JBS3).
 "this scenario is not currently covered by NICE CG181. NICE will consider this as part of the guideline
 update with publication currently expected December 2023
- If patients on a high-intensity statin have side effects, offer a lower dose or an alternative statin (see page 2 'Extent of lipid lowering with available therapies')

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

If recommended statin treatment is contraindicated or not tolerated - follow AAC Statin Intolerance Algorithm for advice regarding adverse effects (click here).

If statin intolerance is confirmed, consider:

- Ezetimibe 10mg monotherapy. Assess response after 3 months (TA385)
- Ezetimibe 10mg/bempedoic acid 180 mg combination when ezetimibe alone does not control non-HDL-C sufficiently. (NICE TA694)

If non HDL-C remains > 2.5mmol/L despite other lipid lowering therapies consider njectable therapies - arrange a fasting blood test and assess eligibility criteria (TA393/394, TA733)

Ezetimibe 10mg daily (NICE TA385). Reassess after three months. If non-HDL-C remains > 2.5mmol/L; consider injectable therapies arrange a fasting blood test and

a and Inclisiran - if fasting LDL-C
≥ 2.6mmol/L despite maximum tolerated lipid lowering therapy (TA733)

eligibility:

* See overleaf for information to support shared decision making

assess eligibility

** Inclisiran and PCSK9i should not be prescribed concurrently • PCSK9i - see overleaf for LDL-C thresholds. (TA393/4) f eligibility criteria not met,

Injectable therapies**

If non-HDL-C > 2.5mmol/L;

Arrange fasting blood test to

measure LDL-C to assess

If eligibility criteria not met, consider ezetimibe 10mg daily (if not previously considered)

Additional CV risk reduction considerations - check fasting triglycerides levels and consider icosapent ethyl. See triglycerides section overleaf.

Summary of National Guidance for Lipid Management for **∧CCELERATED ∧CCESS Primary and Secor** PRIMARY PREVENTION **INITIAL CONSIDERATIONS:** Measure non-fasting full lipid profile (to Ensure appropriate baseline and follow u Consider statin therapy for adults who do not have established CVD but fall into the categories below. Use QRISK risk assessment tool where appropriate (see page 2, 'Primary Prevention Risk Assessment') below. Use QRISK risk assessment tool whe Type 1 diabetes, if Age ≤84 Type 2 & QRISK diabetes or more of the follo & QRISK Over 40 years Age ≤84 Type 2 Type 1 diabetes, if they have one CKD eGFR Age ≥85 over next ≥10% Had diabetes for 10 years over next & QRISK diabetes or more of the following: < 60 vears Have established 10 years Have other CVD & QRISK ≥10% mL/min/1.73m² if appropriate · Over 40 years and/or ≥10% consider over next Identify and address all modifiable risk fa Had diabetes for >10 years physical activity, blo 10 years over next comorbidities. albuminuria Have established nephropathy 10 years frailty & life Consider additional risk factors, if presen Have other CVD risk factors severe mental illness, taking medicines that ca expectancy (e.g. SLE), impaired fasting glyc PRIMARY If lifestyle modification is ineffect Identify and address all modifiable risk factors - smoking, diet, obesity, alcohol intake, Atorvast physical activity, blood pressure and HbA1c. Measure full lipid profile again after 3 months High intensity statin treatment should achieve nfirm not achieved after 3 months: - discuss treatment adherence, timing of do Consider additional risk factors, if present, together with QRISK score (treated for HIV, - If at higher risk (based on comorbidities, ri 'Additional Risk Factors') consider increas severe mental illness, taking medicines that cause dyslipidaemia, systemic inflammatory disorder I/L; dose of atorvastatin 80mg daily. st to For how to increase in people with CKD se (e.g. SLE), impaired fasting glycaemia, recent change in risk factors) SS If patients on a high-intensity statin have side DL-C (see page 2 'Extent of lipid lowering with avail If maximum tolerated dose of statin does not 33) value after 3 months consider adding Ezetimi PRIMARY PREVENTION If statin treatment is contraindicated or not tol - See AAC Statin Intolerance Algorithm for ac If lifestyle modification is ineffective or inappropriate offer statin treatment. 93/4) Ezetimibe 10mg monotherapy may be cons Ezetimibe 10mg/bempedoic acid 180 mg co et, Atorvastatin 20mg daily alone does not control non-HDL-C/LDL-C v ηg

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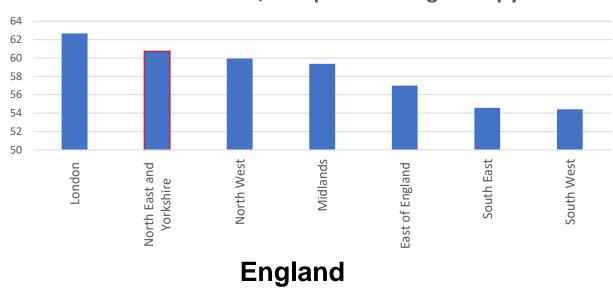
If non-HDL-C reduction remains < 40% of b

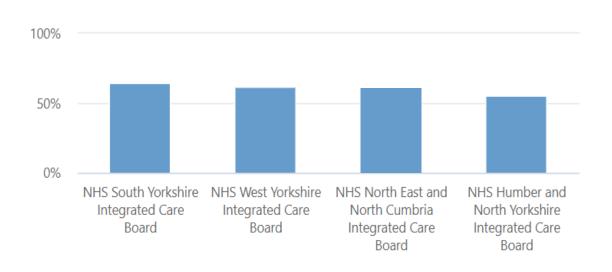
therapy (including people with intolerances ar



Current % of Patients with a Qrisk >20% on Lipid Lowering Therapy

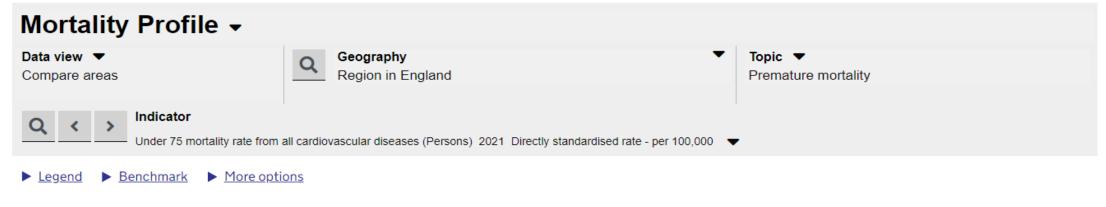
Percentage of patients aged 18 and over with no GP recorded CVD and a GP recorded QRISK score of 20% or more, on lipid lowering therapy





NEY Breakdown





Show 99.8% CI values

Areas All in England

Area ▲▼	Recent Trend	Count ▲▼	Value ▲▼		95% Lower Cl	95% Upper CI
England	_	37,669	76.0	H	75.3	76.8
North West region	_	6,175	92.8	H	90.5	95.1
Yorkshire and the Humber region	_	4,288	86.8	H	84.2	89.4
North East region	_	2,166	84.5	Н	81.0	88.1
West Midlands region	_	4,339	83.5	Н	81.1	86.1
East Midlands region	_	3,674	81.9	H	79.2	84.6
London region	_	4,394	74.3	H	72.1	76.6
East of England region	_	3,733	65.0	H	62.9	67.1
South West region	_	3,622	64.6	Н	62.5	66.8
South East region	-	5,278	63.1	H	61.4	64.8

Source: Office for Health Improvement and Disparities (based on Office for National Statistics source data)

Display Table

Table and chart

Indicator Definitions and Supporting Information

Why invest in cardiovascular disease prevention

PHE estimates that optimising detection of risk factors for CVD and the uptake of anticoagulants, antihypertensives and statins in line with the ambitions, could prevent:





Summary

- We are in a better place than some thank you
- Big challenges ahead
- Still need more work in deprived communities
- Do the basics well
 - Clinical leadership is key
 - Use data
 - Prevention everywhere
 - Simple Guidelines and Professional & Public Education
 - Work together across organisations
 - Identify the high risk individuals early.



Thank you

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