



North Central London  
Joint Formulary Committee

## Guidance for the management of hypertriglyceridaemia

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NCL JFC is funded by and provides advice to Acute Trusts and Clinical Commissioning Groups in NCL.

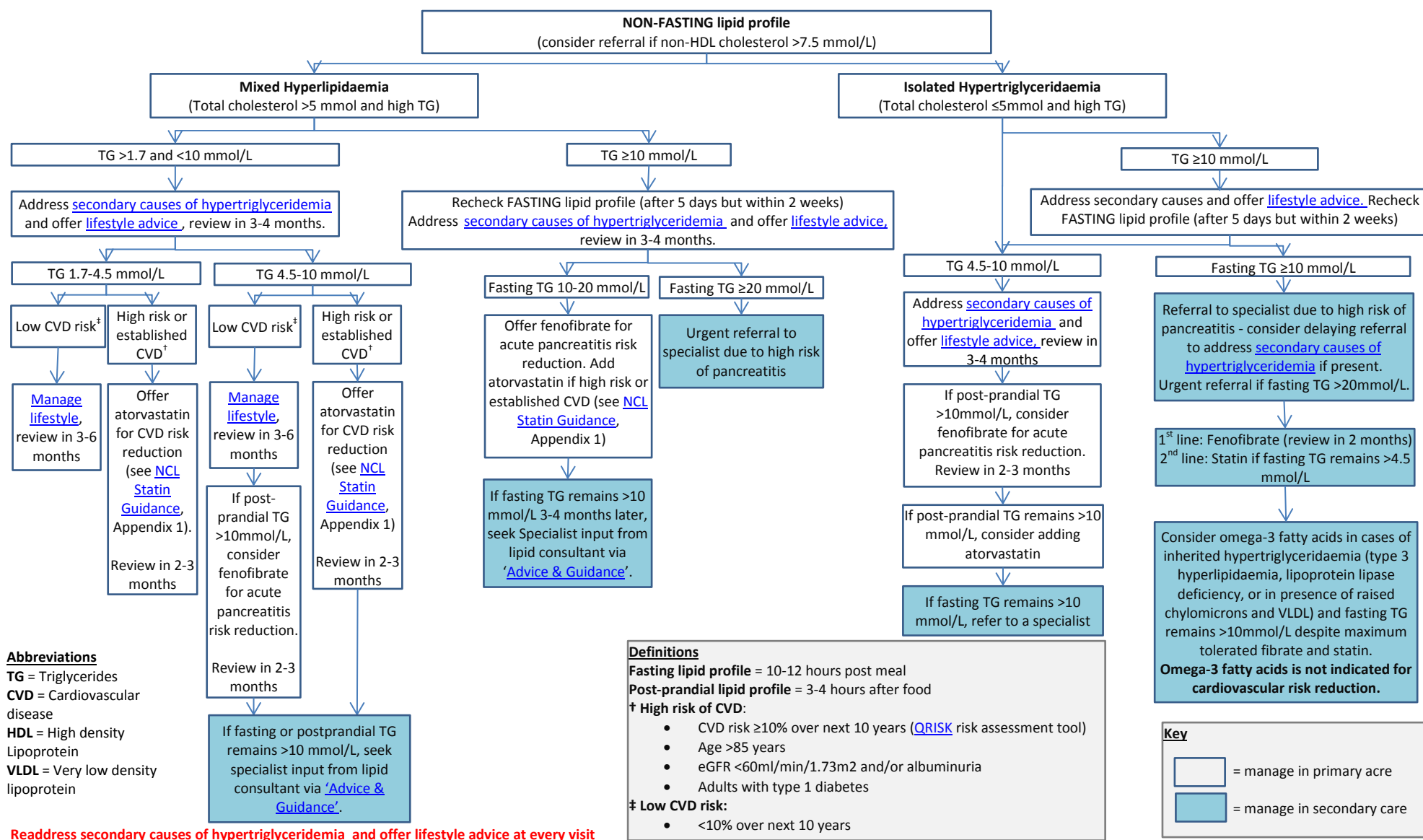
## Document control

Date	Version	Amendments
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## Document management

Groups / Individuals who have overseen the development of this guidance:	Dr Devika Nair, Royal Free London NCL JFC Support Pharmacist
Groups which were consulted and have given approval:	Royal Free London Drug and Therapeutics Committee
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# Management of raised triglyceride concentration



<p><u>Secondary causes of hypertriglyceridemia<sup>1</sup></u></p> <ul style="list-style-type: none"> <li>• Obesity, often in association with an elevation in serum cholesterol</li> <li>• Diabetes mellitus, where there is a relationship to poor glycaemic control and, in type 2 diabetes, obesity</li> <li>• Nephrotic syndrome, often in association with hypercholesterolemia, and renal failure</li> <li>• Hypothyroidism, often in association with hypercholesterolemia</li> <li>• Serum total cholesterol and triglyceride concentrations normally increase markedly during pregnancy</li> <li>• Medicines: <ul style="list-style-type: none"> <li>○ Oestrogen replacement administered orally</li> <li>○ Tamoxifen can cause marked hypertriglyceridemia in a minority of women</li> <li>○ Beta blockers, with the exception of carvedilol</li> <li>○ Immunosuppressive medications, such as glucocorticoids and cyclosporine</li> <li>○ HIV antiretroviral regimens</li> <li>○ Oral retinoids (e.g. isotretinoin)</li> </ul> </li> </ul>	<p><u>Lifestyle advice</u></p> <p>Lifestyle modifications to reduce triglyceride levels are similar to those recommended for individuals at high risk of cardiovascular disease<sup>1</sup> (full lifestyle advice published in <a href="#">NICE CG181<sup>2</sup></a>)</p> <ul style="list-style-type: none"> <li>• Cardioprotective diet including: <ul style="list-style-type: none"> <li>○ Restrict consumption of high glycaemic index/load foods as well as refined sugars, fruit juices, and high fructose beverages<sup>1</sup></li> <li>○ Increased consumption of oily fish<sup>1,2</sup> (pregnant women to limit their oily fish to no more than 2 portions per week and to avoid marlin, shark and swordfish<sup>2</sup>)</li> <li>○ People with very high triglycerides (&gt;10mmol/L) may benefit from the specialist advice from a lipid clinic regarding a very low fat diet</li> </ul> </li> <li>• Physical activity (at least 150 minutes of moderate intensity aerobic activity or 75 minutes of vigorous intensity aerobic activity)</li> <li>• Weight management for those who are overweight or obese</li> <li>• Avoid binge drinking and limit alcohol intake to national recommended limits</li> <li>• Smoking cessation (primarily CV protection)</li> </ul>
<p><u>Investigations for causes of hypertriglyceridemia</u></p> <ul style="list-style-type: none"> <li>• Urine dipstick (nephrotic syndrome)</li> <li>• Blood tests: <ul style="list-style-type: none"> <li>○ Lipid profile (total cholesterol, HDL, non-HDL and triglycerides)</li> <li>○ Fasting glucose or HbA1c</li> <li>○ Renal function</li> <li>○ Thyroid function tests (TFTs)</li> <li>○ Liver function (LFTs)</li> </ul> </li> </ul>	<p><u>Monitoring fibrate therapy</u></p> <ul style="list-style-type: none"> <li>• Recheck lipid levels within 3 months of initiation, aiming for a triglyceride level &lt;4.5 mmol/L</li> <li>• Check serum creatinine at baseline, within 3 months of initiation of treatment and at least annually thereafter (more frequently if clinical indicated). <ul style="list-style-type: none"> <li>○ Hold treatment if creatinine levels &gt;50% ULN (upper limit of normal)</li> <li>○ Consider dose reduction if renal function declines in line with the SPC / BNF</li> </ul> </li> <li>• Monitor liver transaminase levels every 3 months during the first 12 months of treatment and thereafter periodically. <ul style="list-style-type: none"> <li>○ Discontinue therapy if AST or ALT levels increase to more than 3x ULN.</li> <li>○ If symptoms indicative of hepatitis occur (e.g. jaundice, pruritus), and diagnosis is confirmed by laboratory testing, fenofibrate therapy should be discontinued</li> </ul> </li> <li>• Baseline CK should only be checked in those who may already be taking a medicine that will increase the risk of myopathy when used concomitantly with fibrate, such as statin therapy.</li> <li>• Routine CK monitoring for asymptomatic individuals is not recommended. Monitor CK for patients with muscle weakness/pain to assess severity of muscle damage and aid the decision to continue treatment</li> </ul>

For information on prescribing statins and lipid modification for the prevention of CVD see:

- North Central London Statin Prescribing & Lipid Modification Guideline for the Prevention of Cardiovascular Disease
  - [https://www.ncl-mon.nhs.uk/wp-content/uploads/Guidelines/2\\_Lipid\\_modification\\_prevention\\_cardiovascular\\_disease.pdf](https://www.ncl-mon.nhs.uk/wp-content/uploads/Guidelines/2_Lipid_modification_prevention_cardiovascular_disease.pdf)

**Advice and Guidance:** Accessible via eRS: <https://nww.ebs.ncrs.nhs.uk/>. The Royal Free Lipid Centre supports Advice and Guidance and can be identified on eRS as “Lipid Management Service-Cardiology-Royal Free Hospital-RAL”

**Acknowledgement:** Sections of this guideline were taken, with permission, from the South East London APC ‘Guidance for the Management of Hypertriglyceridaemia’ (July 2018)

**Expert opinion:** There are no national guidelines available for the management of hypertriglyceridaemia therefore the pathway structure is based on expert opinion from Royal Free London NHS Foundation Trust Lipid Clinic.

**References:**

1. UpToDate. Hypertriglyceridemia. (2019).
2. National Institute for Health and Care Excellence. CG181: Cardiovascular disease: risk assessment and reduction, including lipid modification. (2016). Available at: <https://www.nice.org.uk/guidance/cg181/chapter/1-Recommendations>. (Accessed: 16th January 2019)