Fellowship of Postgraduate Medicine Lecture

Apolipoproteins in lipid transport, an impressionist view

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Introduction

The major function of the lipoproteins is to provide the organism with a transport system for triacylglycerol and cholesterol. This system appears to be both highly effective and dynamic in normal subjects. While about 150 g of fat per day are fully absorbed from the gut, the plasma triglyceride pool in the blood is approximately 3 g in the fasting state and does not exceed 12 g in the postabsorptive state. About 1 g/day of cholesterol is produced in the liver and 0.2–0.5 g is absorbed from the gut, but the plasma pool is kept at a steady level under 6 g. It is therefore understandable that minor abnormalities in the proteins that regulate lipid transport lead to grossly abnormal plasma lipid levels. These abnormal lipid levels in turn are associated with lipid accumulations at abnormal sites: the artery wall, macrophages, tendons, skin and various other tissues. This causes the clinical picture doctors observe, such as atherosclerosis, xanthomas, corneal opacifications and lipoaemia retinialis.

Clinicians are used to the description of abnormal lipid transport in terms of the lipids proper, for example hypercholesterolaemia, hypertriglyceridaemia and combined hyperlipidaemia. These parameters can be measured easily and relatively accurately. Cholesterol and triglyceride are, however, innocent bystanders in lipoprotein disorders. We will therefore describe the transport disturbances in terms of the protein abnormalities of apolipoproteins, receptors, transfer proteins and enzymes. This enhances the pathophysiological understanding on the one hand and on the other provides arguments for measuring apolipoproteins in the assessment of atherosclerosis risk.

This article is by no means a complete review of the literature; recently two excellent books have appeared with reviews of the different lipoprotein subjects.1,2 Rather, it is an impressionist picture of a rapidly developing scene.

Protein functions in lipid transport

The description which follows cannot be understood without a list of the lipoproteins involved (Table I). The functions of the apolipoprotein can be described in general as follows. They give a defined structure and size to the emulsified macromolecular complexes, which are lipoproteins [apo B for low density lipoprotein (LDL) apo A for high density lipoprotein (HDL)]. Some of them enable the lipoprotein to move through the cellular membrane from inside to outside [apo B48 for chylomicrons, apo B100 for very low density lipoprotein (VLDL)]. Some act as ligands to specific high affinity receptors and guide the lipoproteins to sites of catabolism (apo B100 to the apo B/E receptor, apo E to the apo E and apo B/E receptor). Some are necessary cofactors for enzymes involved in lipid transport [apo AI for lecithin cholesterol acyltransferase (LCAT), apo CII for lipoprotein lipase (LPL)] or inhibitors of catabolic pathways (apo CIII for triglyceride rich particles such as chylomicrons and VLDL).

The enzymes involved in the intravascular compartment are lipoprotein lipase attached to the endothelial surface, which hydrolyses triacylglycerol from chylomicrons and VLDL, hepatic lipase, which probably hydrolyses triacylglycerol from intermediate density lipoprotein (IDL) and probably helps in converting HDL2 to HDL3 by removing part of the phospholipids and cholesterol,3 and lecithin cholesterol acyl transferase which esterifies cholesterol in HDL particles.4

Proteins which promote neutral lipid exchange and/or transfer are called lipid transfer proteins.5 The liver cells and other tissues contain high affinity receptors for LDL, the B/E receptor or LDL receptor,6

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Table I Lipoproteins

<table>
<thead>
<tr>
<th>Name</th>
<th>Description</th>
<th>Structural apoprotein</th>
<th>Apoproteins attached</th>
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<tbody>
<tr>
<td>chylomicron remnant</td>
<td>carries triglyceride from gut to adipose tissue, muscle, liver chylomicron</td>
<td>B48</td>
<td>AI, AIV, C's</td>
</tr>
<tr>
<td>VLDL</td>
<td>carries triglyceride and cholesterol from liver to periphery</td>
<td>B100</td>
<td>C's, E</td>
</tr>
<tr>
<td>IDL</td>
<td>remnant of VLDL</td>
<td>B100</td>
<td>E</td>
</tr>
<tr>
<td>LDL</td>
<td>carries cholesterol to liver and tissues</td>
<td>B100</td>
<td></td>
</tr>
<tr>
<td>HDL2</td>
<td>carries cholesterol from periphery to liver directly and indirectly via other lipoproteins</td>
<td>AI, II</td>
<td>C, E, LCAT, LTP</td>
</tr>
<tr>
<td>HDL3</td>
<td></td>
<td>AII, I</td>
<td>as HDL2</td>
</tr>
</tbody>
</table>

for remnants, the E receptor,7 and putatively for apo A the HDL receptor.8

Proteins can be described in molecular terms. The cDNA sequence, mRNA and amino acid sequence is known for most apoproteins,9 but is not discussed in this article. In order to better understand their individual role in lipoprotein transport we have coined an impressionist name for the most important ones (Table II). We will describe in the following section their function in more detail, and the consequences of abnormalities of each in man.

Table II Impressionist designation of apolipoproteins

<table>
<thead>
<tr>
<th>Apo</th>
<th>Impressionist designation</th>
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<tbody>
<tr>
<td>apo A</td>
<td>'active alternator'</td>
</tr>
<tr>
<td>apo B</td>
<td>'big bulker'</td>
</tr>
<tr>
<td>apo C</td>
<td>'clever cleaner'</td>
</tr>
<tr>
<td>apo E</td>
<td>'enigmatic envoy'</td>
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Apolipoprotein B, the 'big bulker' (Figure 1)

Apo B is called 'big', because it is one of the largest proteins (512,932 daltons) known to circulate in the blood.10 It transports, as apo B48, the 'bulk' of triglyceride from the gut cells in which chylomicrons are formed,11 via lymph and blood to adipose tissue, muscle and in the form of chylomicron remnants to the liver. Apo B48 appears not to bind to the B/E receptor.12 The removal of remnants therefore is dependant on apo E–E receptor interaction. As apo B100 it carries first triglyceride in VLDL to the same LPL degradation sites as chylomicrons.13 Then the VLDL remnant, or IDL, partly binds to the apo B/E receptor and is transformed to LDL. Apo B100 is the sole protein of this particle. It binds to the apo B/E or LDL receptor and is internalized.

One of its functions can be deduced from an experiment of nature, abetalipoproteinaemia, where either a deficient apo B is produced, or a post-translational defect occurs.14 Triglyceride remains trapped inside the intestinal and liver cells. The blood is devoid of chylomicrons, VLDL and LDL. Only HDL is present to carry cholesterol. The triglyceride level is very low, except in the rare apo B100 deficiency where apo B48 is produced normally,15 or in chylomicron retention disease where chylomicrons are absent, but apo B100 is normal.16 Clinical sequelae of the disease are predominantly due to vitamin E deficiency for which LDL is the principal carrier, and vitamin A deficiency for which chylomicrons are the carriers.

Overproduction of VLDL apo B100, probably due to disregulation of the apo B regulator genes, is the putative causal mechanism for familial combined hyperlipidaemia.17 This entity is the most frequently encountered hyperlipidaemia in myocardial infarction survivors, which underlines the importance of hyperapo-B-aemia in the pathogenesis of atherosclerosis.18 Family members have hyper-apo-B-aemia with either elevated levels of VLDL, LDL, or both.

Impaired removal of apo B100 due to deficiency of the LDL receptor leads to familial hypercholesterolaemia, the entity most feared for its atherogenesis at an early age and for its tendon xanthomas.19 It is possible that in the near future more genetic abnormalities of the apo B molecule itself will

<table>
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<th>Apolipoprotein B function</th>
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<tr>
<td>Liver cell Gut cell</td>
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<tr>
<td>Structure Transport Receptor binding</td>
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Figure 1 Apolipoprotein B, the 'big bulker', facilitates transport of chylomicrons and VLDL through the membrane of the intestinal and liver cells. It gives structure to the chylomicrons, VLDL, IDL and the apo B100 of VLDL, IDL and LDL binds to the apo B/E or LDL receptor for internalization in liver and other cells.
be described, which, if receptor binding domain(s) are included, will lead to the expression as familial hypercholesterolaemia.

In some instances glycoprotein (a) binds to apo B, thereby forming a lipoprotein called Lp(a).30 The concentration of Lp(a) is independent of total LDL levels, but it is probably dependent on the difference in Lp(a) phenotypes.21 The presence of Lp(a) appears to be strongly associated with atherosclerosis. It is not known whether this is due to the similarity of glycoprotein (a) and plasminogen.22

Apo A, the ‘active alternator’ (Figure 2)

Apolipoprotein A-I is active because it is involved in both triglyceride and cholesterol transport and activates the essential enzyme for reverse cholesterol transport LCAT. It is an alternator because it enters the lymph as part of the chylomicron but rapidly switches to HDL. It probably forms new ‘nascent’ HDL particles by taking part of the chylomicron surface phospholipid with it. In doing so it facilitates apo CII transfer from HDL to chylomicrons,22 which in turn facilitates chylomicron triglyceride removal. When the detachment of apo A from chylomicrons is deficient as in Tangier disease, this results in both hypertriglyceridaemia and very low levels of HDL.24 apo A is virtually undetectable in Tangier disease since it rapidly disappears with chylomicrons.

Further down the functional road, apo A in nascent HDL picks up free cholesterol from cell surfaces. It is unknown at the present time whether free cholesterol transport from the cell surface to HDL is a random chemical process or depends on the presence of, for example, a sterol carrier protein.23 The cholesterol thus obtained is esterified by LCAT, which is activated by apo AI. The mature HDL particle carries the cholesterol to the liver, either directly by the action of hepatic lipase,3 or indirectly via chylomicron remnants, IDL, and VLDL.28 apo CII transfers from HDL to these particles is facilitated by one or more of the cholesterol ester transfer proteins (or lipid transfer proteins) and inhibited by a lipid transfer protein inhibitor.27 Thus, the so called reverse cholesterol transport pathway from cell, and perhaps atherosclerotic plaques,28 to the liver is completed. The liver is the major site of excretion of cholesterol into the bile. The pathway is also called the centripetal cholesterol pathway, which assumes the liver to be the centre of the metabolic universe.

Overproduction, or underremoval, of apo A results in hyperalphalipoproteinaemia. This entity is also known as ‘longevity syndrome’ since high HDL (apo A) levels are a strong negative risk factor for atherosclerosis.29 Whether this epidemiological finding is indeed based on the activity of the reverse cholesterol transport pathway and not merely a reflection of a low VLDL state, remains to be established.

Apo C, the ‘clever cleaner’ (Figure 3)

Apo C clears the blood of chylomicron triglyceride, but it is clever in sharing this task with LPL and by only being necessary in minor amounts to do the job.30 Apo C is probably produced in the liver and enters the circulation as part of HDL or VLDL. In the plasma it resides in the HDL pool. When chylomicrons enter the lymph apo C is rapidly transferred to the particles,31 probably in exchange for apo A, which makes space available on the chylomicron surface.32 The apo CII molecule has one part between the phospholipid molecules of the chylomicron surface monolayer and one part which binds to LPL,30 opening up, as it were,
the chylomicron particle to the hydrolytic action of LPL. It is a necessary cofactor for LPL. When apo CII is totally absent, massive hypertriglyceridaemia ensues with prolonged circulating chylomicrons and VLDL.\textsuperscript{34} The clinical picture is not different from LPL deficiency. The very high triglyceride levels are associated with recurrent bouts of pancreatitis, lipaemia retinals and sometimes eruptive xanthomas. The clever apo CII is only necessary in minute amounts to fully activate LPL. Heterozygous family members of apo CII deficient patients are normo-

triglyceridaemic.\textsuperscript{35}

Apo CIII is the clever counterpart of CII. In apo AI-CIII deficiency (both genes on the long arm of chromosome 11) the triglyceride level is very low. This probably is due to a lack of chylomicron remnant clearance inhibition by apo CIII.\textsuperscript{36} Some investigators have found low apo CII/CIII ratios in hypertriglyceridaemia and it may be that a low CII/CIII ratio of transferrable protein predisposes patients to develop secondary hypertriglyceridaemia when acquiring hypothyroidism, diabetes or kidney disease.\textsuperscript{37}

**Apo E the ‘enigmatic envoy’ (Figure 4)**

Apo E seems to be sent out by the liver as an envoy to bring back as much cholesterol as it can from different sources. It retrieves the lost sons of triglyceride rich particles to carry their harvest of cholesterol obtained from the reverse cholesterol transport to the paternastical liver. But it is enigmatic as well. It appears in different isoforms.\textsuperscript{38} Each individual has its phenotype, coded for by three alleles, E2, E3 and E4. The apo E2/E2 phenotype is associated with the relatively rare, atherogenic, familial dysbetalipoproteinaemia when VLDL and therefore IDL production is increased.\textsuperscript{39} In general, however, the apo E2/E2 phenotype protects against atherosclerosis by causing low LDL levels.\textsuperscript{40}

Let me explain. Apo E is excreted from the liver associated with HDL. It transfers to chylomicrons and VLDL during and after the process of LPL-mediated triglyceride hydrolysis. The resulting particles, chylomicron remnants and VLDL remnants (IDL) are thus apo E rich. And apo E is the ligand to the apo E receptor. The affinity of each of the apo E isoforms to the apo E receptor is different. Apo E4 binds best, apo E3 a bit less and apo E2 insufficiently.\textsuperscript{30}

The liver cell aims for homeostasis of its cholesterol level. Let us assume that it can get its cholesterol either from LDL via the LDL receptor pathway or otherwise from remnants via the apo E receptor pathway. When an individual has the apo E4/E4 phenotype the cholesterol will be obtained mainly through the apo E receptor. The LDL receptor will be down-regulated, and relatively high LDL levels are the result. On the other hand when the apo E2/E2 phenotype is present, the apo E receptor pathway will be inactive, the LDL receptor will be up-regulated and LDL will be relatively low. The other phenotypes (E4/E3, E3/E3, E4/E2, E3/E2) will be in between. This is indeed exactly what is found within different populations, E4/E4 is high in the cholesterol distribution and E2/E2 low.\textsuperscript{41} It may well explain genetically determined variations in LDL cholesterol levels, apart from apo B or apo B receptor abnormalities.

Low LDL levels are the rule in apo E2/E2 homozygosity. This is mainly due to inadequate catabolism of IDL to LDL. If an over-production of the IDL precursor VLDL is operative, as in familial combined hyperlipidaemia, alcohol use and diabetes, an accumulation of IDL is the consequence. This is known as type III hyperlipidaemia, broad beta disease, remnant removal disease or familial dysbetalipoproteinaemia. The IDL are detected as combined hyperlipidaemia and increased relative cholesterol levels in the ultracentrifugal VLDL fraction.

Apo E phenotyping is now available by a relatively simple immuno-blotting technique for detection of these patients.\textsuperscript{42} The IDL particles are highly atherogenic and type III patients tend to have early severe atherosclerosis, particularly in the peripheral arteries, and pathognomonic palmar xanthomas. The abnormality rapidly responds to dietary measures or drugs which decrease VLDL production, such as the fibrates.

Recently apo E production has been described in other tissues than the liver, such as macrophages.\textsuperscript{43} The role of apo E in cholesterol transport from these cells remains to be clarified. Total apo E deficiency has been described and adds to the enigmatic character of the protein.\textsuperscript{44} Massive hypertriglyceridaemia in this entity may point to a function of apo E3 in triglyceride removal from chylomicrons and VLDL.

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**Figure 4** Apolipoprotein E, the ‘enigmatic envoy’, binds to chylomicron and VLDL remnants and on the other hand to the apo E receptor. Apo E4 does so with a higher affinity than apo E3, while apo E2 binds poorly to the receptor.
LPL, HL, LCAT, CETP, the ‘magnificent modulators’

They are magnificent since deficiency of each results in grossly abnormal lipoproteins. They are modulators because they strongly change the composition of most lipoproteins.

LPL is a long molecule, attached to the luminal side of the endothelial cell. It keeps away the lipoprotein particle from the cell surface. Does it do so to protect the cell against lipid peroxidation products? It modifies the chylomicron to remnant and the VLDL to IDL. It is hormone dependent, specifically to insulin and probably to thyroxine. Its absence results in massive hyperchylomicronaemia (see apo CII deficiency).

Hepatic lipase (HL) is less well-known, its function is not entirely clear. Some investigators have evidence to show that it not only hydrolyses triglyceride but phospholipids and cholesterol ester as well. It would thus modify HDL2 by removing part of its lipids to HDL3 and complete this arm of the reverse cholesterol transport pathway. Others have evidence that it is operative in the modification of IDL to LDL. We have described above that chylomicron and VLDL remnants are both handled by the liver in the same way. Actually chylomicron remnants containing apo B48 and apo E are rapidly internalized by the liver and not converted to another particle.

VLDL remnants (IDL) containing apo B100 and apo E, on the contrary, are either internalized by the liver or bound to its surface and hence metabolized to LDL containing only apo B100. The choice between these two possibilities may be dependent on the size of the particle, and hepatic lipase may be the enzyme responsible for the conversion of the IDL to LDL.

LCAT was an enzyme without a disease to which relatively little attention was paid, until LCAT deficient patients were described. They have a high level of unesterified cholesterol, a low level of esterified cholesterol and an abnormal composition of all lipoproteins, partly as a consequence of structural changes due to an imbalance in the phospholipid-free cholesterol ratio and partly to modifications in apolipoprotein binding. A large vesicular lipoprotein, LpX, is found in obstructive jaundice.

Cholesteryl ester transfer protein (CETP) is a relatively new member of the lipid transport protein family. It facilitates exchange of neutral lipids between lipoproteins. For example cholesterol ester from HDL is avidly exchanged for triglyceride from chylomicrons and VLDL. This process may be elevating the cholesterol content of remnants. Differences in CETP activity have been described in hyperlipidaemia, high levels being associated with low HDL cholesterol and high VLDL + LDL cholesterol. The clinical significance of these proteins and their inhibitors remains to be established, although differences in lipoproteins between species have been explained in the view of different CETP activity.

Apolipoproteins and atherosclerosis

The atherogenic lipoproteins are LDL, IDL, VLDL and possibly chylomicron remnants. All these lipoproteins have apo B as a common structural apolipoprotein. There is one apo B molecule per lipoprotein particle, while the cholesterol content varies. The total plasma apo B level, therefore, is an indication of the concentration of atherogenic particles. The ‘anti-atherogenic’ lipoprotein is HDL, the major structural apolipoprotein of which is apo A. There are approximately two apo A molecules per HDL particle, so that the apo A level is an indication of the number of anti-atherogenic particles in the circulation. The cholesterol content per particle also varies.

If cholesterol is present in both atherogenic and anti-atherogenic particles and in both in varying quantities, why do we measure cholesterol as a major risk indicator for atherosclerosis? Why do we not measure apolipoproteins B and A instead and calculate the apo B/A ratio? The major reason is probably that cholesterol measurement is and has been widely available by relatively well standardized methods. Large amounts of data can be easily accumulated and compared in prospective epidemiological studies to evaluate the risk predicting potential of the cholesterol level (and HDL cholesterol and triglyceride levels).

There is evidence, however, from a number of cross-sectional studies that apo B is a better indicator than cholesterol or even LDL cholesterol. In familial combined hyperlipidaemia, which is associated with atherosclerosis, apo B is elevated, while in familial hypertriglyceridaemia which is not associated with atherosclerosis, the apo B level is close to reference values.

Since immunological methods for the assessment of both apo B and apo A levels (rocket immuno-electrophoresis, radial immuno-diffusion, immuno-nephelometry, ELISA) have become more widely available, it is to be expected that they will replace cholesterol, HDL cholesterol and triglyceride measurement as assessors of risk for atherosclerosis, both for populations and for the individual. Analysis of apo E isoforms and of Lp(a) will probably be added on a large scale. Proper standardization for the measurement of the apolipoproteins is urgently warranted. We may at some time in the future diagnose hyper-apo-B-aemia or hypo-apo-A-aemia instead of hypercholesterolaemia and hypo-HDL-aemia.

It has been the purpose of this review to enhance the readers understanding of the intriguing faculties of the apolipoproteins and of how intricate a system nature
has designed for lipid transport. The practical advantage of this knowledge may be better advice and
treatment for patients who run a high risk for atherosclerosis.

References


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