Co-repressing immunometabolic processes in atherosclerosis

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Summary

Cardiovascular disease is the primary cause of mortality in the world, and tightly associated with the metabolic syndrome, which is a cluster of interconnected metabolic dysfunctions including insulin resistance, obesity, hypertension and dyslipidaemias. These dysfunctions increase the risk of developing atherosclerosis and consequent cardiovascular diseases, such as myocardial infarction and stroke. Atherosclerosis is primarily triggered by increased plasma cholesterol levels and can be classified as an immunometabolic disease, a chronic disease that is affected by both metabolic and inflammatory triggers and/or mediators. These triggers and mediators activate common downstream pathways, including nuclear receptor signalling. Interestingly, specific cofactors that bind to these complexes act as immunometabolic integrators. This review provides examples of such co-regulator complexes, including nuclear sirtuins, nuclear receptor co-repressor 1 (NCOR1), nuclear receptor interacting protein 1 (NRIP1), and prospero homeobox 1 (PROX1). Their study might provide novel insight into mechanistic regulations and the identification of new targets to treat atherosclerosis.

Keywords: atherosclerosis, immunometabolism, mechanisms of disease, nuclear receptor signalling, nuclear receptor co-repressor

Increased plasma cholesterol triggers atherogenesis

Atherosclerosis is a chronic immunometabolic disease and remains asymptomatic until a plaque becomes large enough to obstruct the lumen to cause ischaemic pain or ruptures causing a myocardial infarction, stroke or peripheral artery disease. Research over the past decades demonstrated that the chronic accumulation of lipids, especially cholesterol, in our circulation is the major trigger of atherogenesis. First, most gene candidates identified in familial atherosclerosis development function by primarily altering the lipid metabolism. Atherosclerosis is primarily triggered by increased plasma cholesterol levels and can be classified as an immunometabolic disease, a chronic disease that is affected by both metabolic and inflammatory triggers and/or mediators. These triggers and mediators activate common downstream pathways, including nuclear receptor signalling. Interestingly, specific cofactors that bind to these complexes act as immunometabolic integrators. This review provides examples of such co-regulator complexes, including nuclear sirtuins, nuclear receptor co-repressor 1 (NCOR1), nuclear receptor interacting protein 1 (NRIP1), and prospero homeobox 1 (PROX1). Their study might provide novel insight into mechanistic regulations and the identification of new targets to treat atherosclerosis.

The liver regulates systemic cholesterol metabolism

The liver plays a crucial role in the development of atherosclerosis by regulating metabolic and inflammatory processes, such as the expression of pro-inflammatory cytokines and acute phase response proteins, the secretion of very low density lipoprotein (VLDL) particles, the uptake of cholesterol from the circulation, and biliary cholesterol excretion. An immunometabolic dysregulation in the liver can promote nonalcoholic fatty liver disease (NAFLD) and the development of atherosclerosis. Importantly, NAFLD leads to adverse cardiovascular functions, such as increased oxidative stress and endothelial dysfunction, hypercoagulability and accelerated development of atherosclerosis. Most of the currently used drugs primarily target hepatic cholesterol synthesis via the inhibition of hydroxymethylglutaryl-coenzyme A (HMG-CA) reductase (statins) or hepatic cholesterol re-uptake from the circulation by decreasing the breakdown of the LDL receptor (e.g., PCSK9 inhibitors or silencing).

Macrophage foam cells promote plaque development

Upon exposure to atherogenic triggers, vascular endothelial cells are activated and start to express adhesion molecules. These adhesion molecules attract and activate blood monocytes, which then transmigrate into the arterial intima. Macrophages are the most abundant population of cells within the plaques and one hallmark of atherogenesis.
is the excessive accumulation of cholesterol in monocyte-derived macrophages, leading to foam cell formation. The uptake of modified – especially oxidised – low-density lipoproteins (oxLDL) is driven by scavenger receptors and these are stored as cholesterol esters in lipid droplets. Moreover, macrophages interact with other immune cells, especially T cells, and this immune response can promote pro- or anti-inflammatory processes (fig. 2) [17, 18].

Clinical evidence of immunometabolic regulation

Several powerful cardiovascular drugs demonstrated that metabolic and inflammatory processes can be targeted to treat atherosclerosis (fig. 3), such as the HMG-CA reductase inhibitors (statins) to lower LDL-cholesterol levels, or interleukin-1β (IL-1β) receptor antagonists or colchicine to block the pro-inflammatory processes in cardiovascular disease and type-2 diabetes [4, 5, 19–21]. Although these drugs primarily target either a metabolic or an inflammatory process, recent research demonstrated that they do also affect inflammatory or metabolic signalling, respectively. For example, besides targeting the HMG-CA reductase, statins exert various pleiotropic effects that are mediated by intermediates of the cholesterol biosynthesis pathway. A recent study demonstrated that, by inhibiting mevalonate synthesis, statins counteract the epigenetic reprogramming that leads to trained immunity in monocyte-derived macrophages [22]. Other metabolites of the cholesterol biosynthesis pathway, such as farnesyl pyrophosphate and geranylgeranyl pyrophosphate, also exhibit important anti-inflammatory effects [23]. Further studies demonstrated that statins also modulate the response of the adaptive immune system, such as the function of Th17 cells in autoimmune disease [24].

Crosstalk of inflammatory and metabolic signalling

One of the first hints for a direct role of inflammatory processes in the development of a metabolic disease came in the 1990s, when it was shown that normal and other...
osclerotic arteries express tumour necrosis factor (TNF) [25]. Two years later, it was demonstrated that the adipose tissue secretes TNF in obese and/or diabetic mice and humans, thereby directly linking an inflammatory cytokine to another chronic metabolic disease [26–28]. Later studies showed that the genetic deletion of Tnf reduces atherosclerosis development [29–31], and it is now known that various cytokines and chemokines regulate atherogenesis [32, 33].

Binding of TNF and other cytokines and chemokines to their corresponding receptors induces the activation of downstream signalling kinases, including IκB kinase (IKK), c-Jun N-terminal kinase (JNK), p38 and Janus kinase (JAK) (fig. 4). Signalling from these kinases leads to the activation of central inflammatory transcription factors, such as nuclear factor-kB (NF-kB) upon IKK signalling, activating protein-1 (AP-1) upon p38/JNK (MAPK) signalling, or signal transducers and activators of transcription (STAT) upon JAK1 signalling. Notably, these inflammatory transcription factors also regulate metabolic processes at different layers. For example, they interact with and coordinate the transcriptional activity of metabolic transcription factors, such as the nuclear receptors peroxisome proliferator-activated receptor γ (PPARγ) and liver X receptors (LXRs), which in turn are activated by specific lipids [34–36]. Furthermore, inflammatory signalling can induce the expression of central metabolic regulators, such as sterol regulatory element-binding proteins (SREBP-1 and SREBP-2), and of a series of metabolic enzymes [37–41].

Lipid-responsive nuclear receptors with immunometabolic functions

The family of nuclear receptors can be subdivided into endocrine receptors that are bound by hormones, adopted orphan nuclear receptors that are activated by different metabolites such as fatty acids and cholesterol derivates, and orphan nuclear receptors that do not (yet) have any known natural ligand [42, 43]. Importantly, most nuclear receptors have a ligand binding pocket and are therefore potential drug targets. The function of several nuclear receptors in atherosclerosis has been reviewed previously [44]. The class of adopted and orphan nuclear receptors is of special interest since several of them are bound and activated by specific ligands, such as PPARγ, LXRs and liver receptor homologue-1 (LRH-1) (fig. 4). Besides regulating various metabolic processes, these lipid-binding nuclear receptors also mediate transrepression of pro-inflammatory molecules in metabolic organs and immune cells [34–36, 45–47], thus acting as immunometabolic regulators.

Transcription cofactors as immunometabolic integrators

Transcriptional co-regulators play a central role in immunometabolic regulation because they are regulated by upstream inflammatory or metabolic mediators and coordinate the transcriptional activity of key immunometabolic transcription factors. Importantly, several cofactors are expressed in a tissue-specific manner, at a specific time point during development or upon specific upstream stimuli, therefore acting under very specific conditions. From an immunometabolic point of view it will be very inter-

Figure 3: Anti-atherosclerotic and/or cardioprotective drugs targeting inflammatory or metabolic pathways. Examples of drugs that target inflammatory or metabolic processes that promote the development of atherosclerosis.
esting to identify and study the co-regulators that are implicated in both inflammatory and metabolic processes. Approximately 300 transcriptional cofactors exist in mammalian cells [48], but the function of most these cofactors is not yet known. The functions of some selected cofactors that regulate atherosogenesis are summarised below.

**Sirtuins – multiple roles in atherosclerosis and cardiovascular diseases**

Sirtuins are nutrient-sensitive protein deacetylases, which play important roles in various molecular and physiological processes [49]. SIRT1, the best studied sirtuin, exerts various protective cardiovascular functions. Several studies demonstrated that SIRT1 diminishes the development of atherosclerosis and exhibits other cardioprotective effects (fig. 5) [51, 52]. By deacetylating and interfering with the activation of the RelA/p65 subunit of NF-κB, SIRT1 inhibits inflammatory signalling, prevents macrophage foam cell formation, diminishes endothelial cell activation and reduces the expression of tissue factor, a key factor in the activation of the coagulation cascade during atherothrombosis [53–56]. Besides exerting these direct anti-inflammatory effects, SIRT1 also regulates the function of specific nuclear receptors. These studies highlight that SIRT1 acts as an immunometabolic regulator. Other sirtuins also exert distinct cardioprotective functions (reviewed in [57]). Several approaches that were tested in order to induce the activity of sirtuins by increasing the levels of its substrate, nicotinamide adenine dinucleotide (NAD⁺), showed protective cardiometabolic effects [58–63].

**NCOR1 – an emerging regulator of immunometabolic processes**

NCOR1 forms a large co-repressor complex containing histone deacetylases that inhibit the transcriptional function of nuclear receptors [64, 65]. Tissue-specific deletions of Ncor1 in metabolic organs showed that it regulates fatty acid metabolism, mitochondrial functions, insulin sensitivity, bile acid composition and intestinal cholesterol absorption [66–69]. Moreover, NCOR1 exerts both pro- and anti-inflammatory functions in macrophages, and its deletion in adipocytes reduces adipose tissue macrophage infiltration and inflammation [35, 67, 70, 71].

Recently published data demonstrated that the deletion of macrophage Ncor1 promotes atherosclerosis development in mice, and their atherosclerotic lesions displayed thinner fibrous caps and larger necrotic cores, suggesting that NCOR1 stabilises atherosclerotic plaques [72]. At the molecular level, it was shown that NCOR1 binds to the Cd36 promoter, a scavenger receptor that mediates the uncontrolled uptake of modified LDL particles [73, 74]. Consequently, NCOR1-deficient peritoneal macrophages displayed increased CD36 expression and enhanced accumulation of oxLDL compared with control macrophages (fig. 5). The increased expression of CD36

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**Figure 4: Immunometabolic signalling at the cellular level.** Simplified overview of important signalling pathways that are activated by inflammatory or metabolic mediators, such as cytokines and lipids, which trigger downstream inflammatory (red labels) or metabolic pathways (blue labels), and activate key transcription factors. Transcription cofactors (yellow) play a central immunometabolic role by directly reacting to upstream immune and metabolic mediators and coordinating the downstream responses at the nuclear level, either by directly driving the expression of target genes or by interfering with the function (e.g., transrepression) of other transcription factors, such as the AP-1 or NF-κB. CHOL = cholesterol; FA = fatty acid; HODE = hydroxyoctadecadienoic acid; DNL = de novo lipogenesis; Mϕ = macrophage; nSIRTs = nuclear Sirtuins; PUFAs = polyunsaturated fatty acids; RCT = reverse cholesterol transport.

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was secondary to a derepression of PPARγ target genes. Treatment of peritoneal macrophages with the PPARγ agonist rosiglitazone led to an enhanced expression of several direct and indirect PPARγ target genes, including both pro- and anti-inflammatory molecules, thus suggesting that NCOR1 exerts atheroprotective functions by repressing detrimental functions of PPARγ in macrophages (fig. 5).

To establish the contribution of macrophage NCOR1 in human atherosclerosis, the authors further explored its expression in publicly available datasets and showed that the expression of Ncor1 was reduced in carotid plaques in comparison to non-atherosclerotic sites from mammary arteries. Moreover, analysis of expression data from laser micro-dissected macrophages revealed that all significantly changed PPARγ target genes were robustly increased in macrophages from ruptured compared with non-ruptured plaques, and targeted protein analyses showed that NCOR1 is reduced in ruptured compared to non-ruptured carotid plaque [72].

Earlier studies demonstrated that NCOR1 can also interact with B cell lymphoma-6 (BCL6), a transcriptional repressor that belongs to a class of zinc-finger transcription factors. This interaction promotes the repression of several processes, including the expression of pro-inflammatory NF-κB target genes [76–78].

In the liver, nuclear corepressors modulate liver energy metabolism during the fasting–feeding transition. The NCOR1-HDAC3 complex regulates both catabolic and anabolic processes in the liver. Upon feeding, the target of rapamycin complex 1 (mTORC1)-AKT signalling pathway is activated by increasing levels of glucose and insulin, which leads to the phosphorylation of serine 1460 of NCOR1 (pS1460 NCOR1). The pS1460 phosphorylation reduces the capacity of NCOR1 to interact with LXRα, hence increasing the transcription of de novo lipogenesis genes [68]. Under fasting conditions, the NCOR1-HDAC3 complex represses the expression of de novo lipogenesis genes (fig. 5) [79, 80].

**PROX1 – a critical regulator of hepatic metabolism**

Prospero-related homeobox 1 (PROX1) is a conserved corepressor that is mainly expressed in liver, lymphatic vessels, lens, dentate gyrus, neuroendocrine cells of the adrenal medulla, megakaryocytes, and platelets [81]. It interacts with several nuclear receptors, including the LRH-1, oestrogen-related receptor (ERR), steroid and xenobiotic receptor (SXR), and retinoic acid-related orphan receptors (ROR), as well as other transcriptional regulators, such as HDAC3 or lysine-specific demethylase 1 (LSD1) [82–84]. The interaction of PROX1 with LRH-1 was first described in Drosophila [85] and human cells [86], and the LRH-1/PROX1 interaction has been reported to regulate liver development and function [86, 87]. Importantly, it was demonstrated that PROX1 is required to promote the beneficial effects of LRH-1 on reverse cholesterol transport and atherogenesis: Atherosclerosis-prone mice carrying a mutation that abolishes the SUMOylation of LRH-1 are significantly protected from atherosclerosis development by promoting reverse cholesterol transport (fig. 5) [88]. This study highlighted that a single posttranslational modification of a nuclear receptor is sufficient to modulate the function of the protein and impact on the development of chronic cardiovascular diseases. Moreover, LRH-1 drives the differentiation of macrophages towards an anti-inflammatory phenotype (fig. 5) [89]. Whether or not the LRH-1-PROX1 complex affects the development of atherosclerosis in humans is not yet known. However, as it is a potent regulator of bile acid synthesis and composition [90], it is tempting to assume that this nuclear receptor-co-repressor complex plays a key role in chronic hepatic diseases or upon liver transplantation. On the other hand, the same LRH-1 mutation promotes the development of NAFLD and early signs of steatohepatitis and Lrh-1-deficient mice are protected against hepatic cancer [91, 92].
RIP140 – a regulator of macrophage metabolism and inflammation

Nuclear receptor interacting protein 1 (NRIPI1; also known as receptor interacting protein 140, RIP140) is a co-receptor that is ubiquitously expressed, interacts with several nuclear receptors and regulates downstream immunometa-

bolic processes [93, 94]. Interestingly, TLR signalling leads to RIP140 ubiquitination and degradation, which in turn reduces pro-inflammatory cytokine production and promotes endotoxin tolerance [95]. Moreover, RIP140 pro-
motes foam cell formation by inhibiting LXR-driven Ab-
cal and Abagl expression, thus impairing HDL-driven cholesterol eflux from macrophages [96, 97]. Consistent-
ly, microRNA-specific silencing of Rip140 in macrophages reduces the development of atherosclerotic lesions in Apoe knockout mice (fig. 5) [96].

In the liver, RIP140 is recruited to a complex of krüppel-like factor 15 (KLF15) and LXR/RRXR on the promoter of genes regulating de novo lipogenesis and thus regulating the feeding-fasting transition (fig. 5) [98, 99]. This RIP140-KLF15-LXR complex likely regulates other processes that affect atherosclerosis development in the liver, macrophages and other tissues. Notably, KLF15 mRNA expression is reduced in human atherosclerotic com-
pared to nonatherosclerotic aortae, and the systemic or smooth muscle cell-specific deletion of Klf15 aggravates atherosclerosis development in Apoe knockout mice [100].

Outlook

Myocardial infarction and stroke are the leading causes of mortality worldwide, and atherosclerotic plaque rupture or erosion are the primary triggers of these two cardiovascular diseases. Currently, the primary approach to reduce the progression of atherosclerosis in humans is to target the dyslipidaemia, especially by reducing increased LDL-cholesterol levels, besides treating hypertension and diabetes mellitus. Targeting the LDL metabolism levels will con-
tinue to be primary intervention in patients with increased total and LDL-cholesterol, which can now be successful-
fully achieved by combining statins with intestinal Nie-
mann-Pick C1-like protein 1 (NPC1L1) or hepatic PCSK9 inhibitors [3–10]. Furthermore, new drugs to treat dyslipi-daemias are emerging, such as apolipoprotein C3 (ApoC3), Apo(a) or angiopoietin like 3 (ANGPTL3) silencing therapeu-
atics [101].

Another potent predictor of major adverse cardiovascular events besides LDL-cholesterol is high-sensitivity C-reactive protein (hsCRP), thus suggesting that targeting central pro-inflammatory processes will probably also be a future therapeutic option. This is also underlined by the success of the recent COLCOT and CANTOS trials [19, 21]. Therefore, there is a clear biomedical interest in identifying, understanding and testing new (anti-)atherogenic pathways as therapeutic targets.

Targeting specific nuclear receptor-cofactor complexes could become a promising approach to promote beneficial metabolic functions and inhibit chronic anti-inflammatory processes. Some examples were discussed in this review, but research over past years identified many more promising targets for cardiovascular diseases [102]. For instance, strategies to stabilise or augment NCOR1 function in macrophages could protect against excessive oxLDL uptake, macrophage foam cell formation and pro-inflammatory gene expression, and thus prevent further growth and destabilisation of atherosclerotic plaques [50, 72], such as in primary prevention for patients with very high risk for atherosclerotic disease or in secondary prevention. In the liver, targeting NCOR1, for example with synthetic siRNAs conjugated to triantennary N-acetylgalactosamine carbohydrates [9, 103], could be tested to prevent fatty acid and cholesterol synthesis in patients with NAFLD, which is a growing patient population globally and exposed to in-
creased cardiovascular risk [50, 68]. However, these ther-
apeutic approaches first have to be tested in pre-clinical models before being translated to humans.

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Potential Competing interests

No conflict of interest relevant to this article was reported.

References

1 Kessler T, Vilne B, Schunkert H. The impact of genome-wide associa-
tion studies on the pathophysiology and therapy of cardiovascular dis-

2 Roberts R. Genetics of coronary artery disease. Circ Res. 2014;114(12):1890–903. doi: http://dx.doi.org/10.1161/CIRCRESA-
HA.114.302992. PubMed.

3 Cannon CP, Blazing MA, Giugliano RP, McCagg A, White JA, Theroux
P, et al.; IMPROVE-IT Investigators. Ezetimibe Added to Statin Thera-
Moa1410489. PubMed.

4 Nissen SE, Tsuzuki EM, Schoenhagen P, Crowe T, Sasaeda W, Tsai J, et al.; Reversal of Atherosclerosis with Aggressive Lipid Lowering (RE-

5 Ridker PM, Cannon CP, Morrow D, Rifai N, Rose LM, McCabe CH, et al.; Pravastatin or Atorvastatin Evaluation and Infection Therapy-
Thrombolysis in Myocardial Infarction 22 (PROVE-IT-TIMI 22) Investi-
Moa042378. PubMed.


Moa1801174. PubMed.


Moa1615758. PubMed.

10 Ray KK, Wright RS, Kallend D, Koenig W, Leiter LA, Ralil FJ, et al.; ORION-10 and ORION-11 Investigators. Two Phase 3 Trials of Inclisis-


Uttermostly, the target of immunomodulation in Th17-mediated autoimmune diseases is to neutralize Th17 cells, which are primarily involved in the induction of inflammation. Several studies have shown that Th17 cells contribute to the progression of autoimmune diseases, such as rheumatoid arthritis, multiple sclerosis, and inflammatory bowel disease. Therefore, the development of immunomodulatory agents that target Th17 cells is of great interest.

The main immunomodulatory mechanisms involved in the regulation of Th17 cells include the induction of apoptosis, the inhibition of Th17 cell differentiation, and the suppression of Th17 cell function. Several agents, such as the anti-inflammatory agents, have been shown to have immunomodulatory effects on Th17 cells.

The treatment of autoimmune diseases has relied primarily on the use of anti-inflammatory and immunosuppressive agents, which are generally effective in controlling the symptoms of these diseases. However, these agents are often associated with significant side effects, such as infections, bleeding, and cancer. Therefore, the development of novel immunomodulatory agents with fewer side effects is of great importance.

The development of immunomodulatory agents has been facilitated by the identification of the key pathways that regulate Th17 cell differentiation and function. These pathways include the NF-κB, PPARγ, and JAK-STAT pathways, which are involved in the regulation of Th17 cell differentiation and function. The development of agents that target these pathways is of great importance, as it has the potential to lead to the development of effective immunomodulatory agents.

In conclusion, the development of immunomodulatory agents that target Th17 cells is of great importance for the treatment of autoimmune diseases. The identification of the key pathways that regulate Th17 cell differentiation and function has facilitated the development of immunomodulatory agents. The development of novel immunomodulatory agents with fewer side effects is of great importance, as it has the potential to lead to the development of effective immunomodulatory agents.


