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Hebe Agustina Mena, Matthew Spite

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### Commentary

Nonresolving inflammation contributes to the progression of atherosclerosis, a chronic disease characterized by the accumulation of lipid-rich arterial plaques infiltrated with immune cells. In this issue of the *JCI*, Arnardottir and Thul et al. report that GPR32, a receptor for proresolving lipid mediators including resolvin D1, was decreased in human atherosclerotic lesions and that overexpression of this human receptor in mice reduced lesion area and necrosis of atherosclerotic plaques. Mechanistically, GPR32 signaling blunted the production of proinflammatory cytokines, enhanced macrophage phagocytosis, and reduced leukocyte accumulation. These results suggest that therapeutic targeting of GPR32 could be an approach to resolving chronic inflammation in atherosclerosis.

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# Proresolving receptor tames inflammation in atherosclerosis

Hebe Agustina Mena and Matthew Spite

Center for Experimental Therapeutics and Reperfusion Injury, Department of Anesthesiology, Perioperative and Pain Medicine, Brigham and Women's Hospital and Harvard Medical School, Boston, Massachusetts, USA.

**Nonresolving inflammation contributes to the progression of atherosclerosis, a chronic disease characterized by the accumulation of lipid-rich arterial plaques infiltrated with immune cells. In this issue of the *JCI*, Arnardottir and Thul et al. report that GPR32, a receptor for proresolving lipid mediators including resolvin D1, was decreased in human atherosclerotic lesions and that overexpression of this human receptor in mice reduced lesion area and necrosis of atherosclerotic plaques. Mechanistically, GPR32 signaling blunted the production of proinflammatory cytokines, enhanced macrophage phagocytosis, and reduced leukocyte accumulation. These results suggest that therapeutic targeting of GPR32 could be an approach to resolving chronic inflammation in atherosclerosis.**

## Proresolving receptors actively promote the resolution of inflammation

The acute inflammatory response protects the host, yet is actively resolved to prevent tissue injury and facilitate repair/regeneration. During resolution, polymorphonuclear neutrophil (PMN) recruitment is dampened and phagocyte-mediated removal of apoptotic (i.e., efferocytosis) and necrotic cells ensues. Macrophages are key expert phagocytes in resolution that are temporally reprogrammed to facilitate tissue repair, which associates with decreased proinflammatory mediators and a reciprocal increase in antiinflammatory mediators and growth factors. These key functions in resolution are regulated in part by a superfamily of lipid autacoids termed specialized proresolving mediators (SPMs), the discovery of which helped to define resolution as an active biological process (1).

The SPMs comprise distinct chemical mediators enzymatically biosynthesized

from polyunsaturated fatty acids. These mediators include the E-series and D-series resolvins, derived from omega-3 fatty acids eicosapentaenoic acid and docosahexaenoic acid, respectively, and lipoxins, derived from omega-6 fatty acid arachidonic acid. The biological actions of SPMs are largely mediated by GPCRs expressed in leukocytes and vascular cells (2). For example, resolvin D1 (RvD1; 7S,8R,17S-trihydroxy-4Z,9E,11E,13Z,15E,19Z-docosahexaenoic acid) and its aspirin-triggered (AT) 17R epimer are ligands for two distinct human GPCRs termed ALX/FPR2 and GPR32 (also known as DRV1; ref. 3). Authentic RvD1 binds and activates these receptors with high potency ( $EC_{50}$  is ~1–10 pM) and initiates downstream signaling, leading to regulation of proinflammatory cytokines and microRNAs, reduced PMN–endothelial cell (PMN-EC) interactions, and enhanced macrophage phagocytosis (3–5). In vivo, the proresolving and tissue-reparative actions of RvD1 are lost in mice with *Alx/Fpr2* deficiency,

while overexpression of human ALX/FPR2 enhances resolution (2, 4–6). Nonetheless, the biological roles of ALX/FPR2 in vivo are time and context dependent, given that both proinflammatory (e.g., serum amyloid A) and other proresolving mediators (e.g., lipoxin  $A_4$ ) are ligands. In contrast, there is no murine homolog of GPR32, and thus its in vivo role in resolution has been uncharacterized to date. In this issue of the *JCI*, Arnardottir and Thul et al. address this gap by reporting on their generation of mice with Tg overexpression of human GPR32 on an *Apoe*- and *Alx/Fpr2*-deficient background and demonstrate a proresolving role of GPR32 in acute inflammation as well as chronic unresolved inflammation associated with atherosclerosis (7).

## Nonresolving inflammation plays a role in atherosclerosis progression

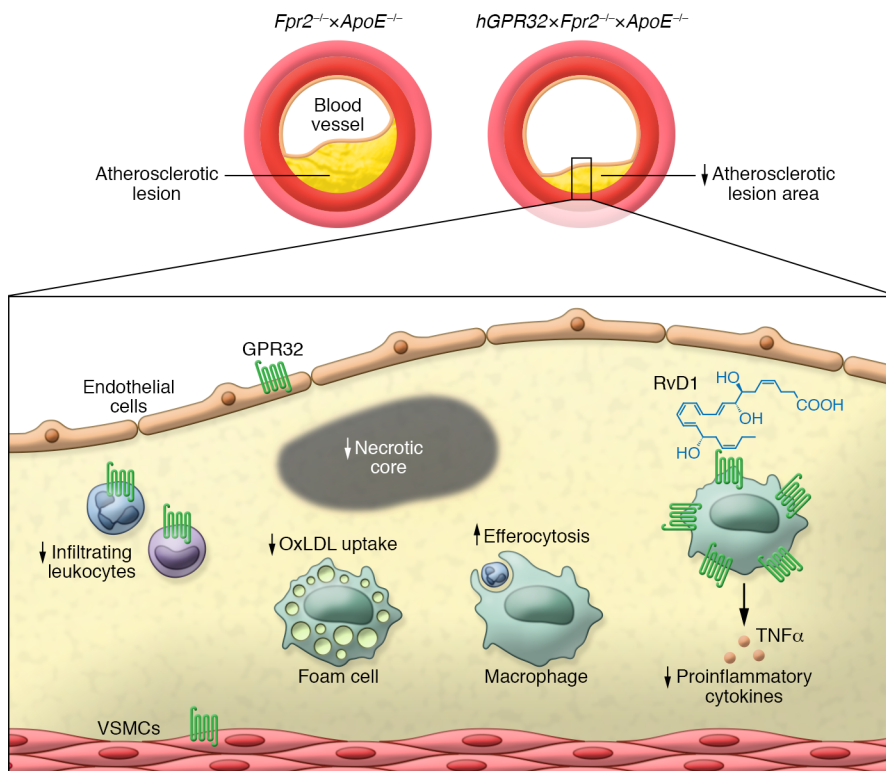
Atherosclerosis is a chronic vascular disease in which subendothelial retention of modified LDLs initiates maladaptive inflammation. Atherosclerotic plaques are characterized by the accumulation of innate and adaptive immune cells, macrophage and vascular smooth muscle cell-derived (VSMC-derived) foam cells, and production of proinflammatory cytokines, chemokines, and lipid mediators (8). The formation of a necrotic core and thin fibrous cap are characteristics of advanced lesions prone to rupture, which can initiate thrombosis and lead to myocardial infarction and stroke. Defective macrophage efferocytosis is a causal factor in the development of advanced plaques, suggestive of impaired resolution mechanisms (8). Indeed, the ratio of SPMs (e.g., resolvins) to proinflammatory lipid mediators (e.g., leukotrienes) is decreased in vulnerable human plaques, which is also reflected in the saliva of individuals with cardiovascular disease (9–11). The ratio of SPMs to proinflammatory lipid mediators is positively associated with angiographically defined atherosclerosis regression in patients with stable coronary

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**Conflict of interest:** MS is an inventor on an awarded patent assigned to Brigham and Women's Hospital titled "Anti-inflammatory particles" (US patent 10,154,977).

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**Figure 1. The role of human GPR32 in atherosclerosis.** Tg overexpression of *hGPR32* on an *ApoE*<sup>-/-</sup> and *Fpr2*<sup>-/-</sup> background reduces atherosclerotic lesion area and necrotic cores as compared with that of non-Tg *Fpr2*<sup>-/-</sup> × *ApoE*<sup>-/-</sup> mice. Lesions of *hGPR32* Tg mice are characterized by a lower number of monocytes and neutrophils, reduced macrophages, and decreased levels of the proinflammatory cytokine TNF- $\alpha$ . Of note, overexpression of *hGPR32* also increases macrophage efferocytosis in peritonitis, and RvD1 enhances macrophage phagocytosis and lowers the uptake of oxidized LDL by macrophages in vitro in an *hGPR32*-dependent manner. These results suggest that GPR32 has multiple proresolving and atheroprotective roles. Similarly, RvD1 binds GPR32, enhances phagocytosis, and lowers the uptake of oxidized LDL.

artery disease (12). Mechanistic studies in animal models of atherosclerosis collectively demonstrate that increasing SPMs resolves plaque inflammation, improves efferocytosis, decreases necrotic cores, and increases markers of lesion stability (i.e., fibrous cap thickening; refs. 8, 9, 11, 13). SPMs also diminish the extent of injury following vascular and endovascular surgical procedures, in part by targeting proliferative VSMCs and inflammation (14). Arnardottir and Thul et al. add to this growing body of literature and establish the proresolving and atheroprotective roles of GPR32 (7).

### Proresolving actions of GPR32 in atherosclerosis

Proresolving receptors, such as ALX/FPR2 and ERV1/ChemR23 (a receptor for resolvin E1; RvE1), are expressed in both immune cells and vascular cells of human atherosclerotic plaques. Further, ALX/FPR2 levels are increased in atherosclerosis (8, 15). In contrast, Arnardottir and Thul et al. observed that human atherosclerotic plaques show decreased *GPR32* mRNA levels compared with control arteries and showed that *GPR32* mRNA levels positively correlated with expression of genes encoding M2 macrophage marker *ARG1*, Treg marker *FOXP3*, and Th2 cyto-

kine *IL4* as well as M1 macrophage marker *NOS2*. At the protein level, GPR32 was expressed in *ARG1*<sup>+</sup> macrophages and to a lower extent in T cells, VSMCs, and ECs (Figure 1). The authors hypothesized that increasing its expression may have atheroprotective roles in vivo, acting by resolving plaque inflammation. To this end, they generated a mouse strain in which human *GPR32* (*hGPR32*) was expressed in atherosclerosis-prone *ApoE*-deficient mice that also lacked murine *Alx/Fpr2* (7). This approach was needed because SPMs that activate GPR32 also target ALX/FPR2 (2). After high-fat diet feeding, *hGPR32* Tg mice had smaller atherosclerotic lesions as compared with nontransgenic littermates. Features of advanced atherosclerosis, such as necrotic cores, were reduced in *hGPR32* Tg mice, and this reduction coincided with reduction in leukocytes and the proinflammatory cytokine TNF- $\alpha$ . These atheroprotective effects occurred independently of changes in circulating immune cells.

The results of the study by Arnardottir and Thul et al. are congruent with prior studies demonstrating that loss of *Erv1/ChemR23* in atherosclerosis-prone mice worsens lesion development and that increasing RvE1 reduces atherosclerosis in both mice and rabbits (16, 17). Moreover,

RvD1 treatment in *Ldlr*<sup>-/-</sup> mice diminishes necrotic cores, which is similar to what is observed in *hGPR32* Tg mice and indicative of enhanced efferocytosis (9, 18). However, it should be noted that, while RvD1 increases lesional efferocytosis and clearance of necrotic cells, these processes were not evaluated in the study by Arnardottir and Thul et al. (9, 18). Moreover, SPMs increase fibrous cap thickness in advanced murine lesions, although these changes in plaque morphology were not specifically assessed in the study by Arnardottir and Thul et al. (9, 11). Moreover, the *hGPR32* Tg × *ApoE*<sup>-/-</sup> × *Fpr2*<sup>-/-</sup> mice were only compared with *ApoE*<sup>-/-</sup> × *Fpr2*<sup>-/-</sup> littermates and loss of *Fpr2* alone is known to affect atherosclerotic lesion area and stability (8, 15). Nonetheless, all aforementioned studies demonstrate atheroprotective roles for SPMs and their receptors and these effects occur independently of changes in cholesterol, indicating that activating proresolving receptors may offer an approach for reducing residual risk in patients already on lipid-lowering therapy.

### Regulation of proresolving immune cell subsets by GPR32

RvD1 engages GPR32 to promote macrophage efferocytosis and phagocytosis,

key functions in resolution (3). Moreover, the RvD1/GPR32 axis modulates leukocyte-EC interactions in response to proinflammatory stimuli (5, 19). To obtain additional evidence for a proresolving role of GPR32 in vivo, Arnardottir and Thul et al. utilized a well-characterized model of self-limited peritoneal inflammation. Here, overexpression of hGPR32 in *Apoe*<sup>-/-</sup> × *Fpr2*<sup>-/-</sup> mice reduced PMN infiltration in response to zymosan as compared with that in control mice, although the impact of hGPR32 on inflammation resolution in the absence of *Apoe* deficiency-induced hypercholesterolemia was not specifically evaluated. Notably, AT-RvD1 administration prior to zymosan similarly reduced PMN levels and AT-RvD1 had no effect in *Apoe*<sup>-/-</sup> × *Alx/Fpr2*<sup>-/-</sup> mice, demonstrating receptor specificity. Overexpression of hGPR32 markedly increased the macrophage/PMN ratio and increased efferocytosis in vivo, providing definitive evidence that GPR32 is a proresolving receptor. These results are congruent with overexpression of hALX/FPR2 in mice, which increases resolution in response to RvD1 (4). Interestingly, hGPR32 Tg mice had higher levels of *Alox15*, a resolution macrophage marker encoding an enzyme that promotes SPM biosynthesis, as well as lower levels of proinflammatory cytokines *Il1β* and *Tnfa*. These results are consistent with GPR32-dependent roles of RvD1 in regulating cytokine production in human macrophages (20). In macrophages isolated from hGPR32 Tg mice, AT-RvD1 markedly enhanced phagocytosis of zymosan and bacteria while blunting the uptake of atherogenic oxidized LDL. Functional coupling of the hGPR32 to downstream signaling was further evidenced by phosphorylation of ERK1/2 in response to AT-RvD1 in these macrophages. Although the role of the RvD1/GPR32 axis on T cell subsets was not investigated, the authors note that GPR32 was expressed on T cells in human atherosclerotic lesions and correlated with FOXP3, a Treg marker. Importantly, RvD1 prevents naive CD4<sup>+</sup> T cell differentiation into Th1 and Th17 cells and enhances de novo generation and function of FOXP3<sup>+</sup> Tregs in a GPR32-dependent manner (21). Given that regression of atherosclerosis requires Tregs to modulate macrophage phenotype and efferocytosis, it will be important to further define the cell-specific roles of GPR32 in future studies (22).

## Summary and future directions

Arnardottir and Thul et al. provide evidence of the proresolution and atheroprotective actions of GPR32. Targeting GPR32 with a stable analog, such as benzo-diacetylenic-17R-RvD1-methyl ester, could also be considered, as it is more resistant to metabolic inactivation than native RvD1, yet has similar biological functions (23). Moreover, a recent chemical library screen identified additional GPR32 small molecule chemotype agonists that may offer advantages in terms of stability and scalability (24).

Building on decades of animal studies that show a causal role for inflammation in the progression of atherosclerosis, recent clinical trials demonstrate a reduction in cardiovascular events in patients treated with inhibitors of inflammation (25). Nonetheless, traditional antiinflammatory approaches may impair host defense. Unlike antagonists, SPMs serve as agonists to actively dampen inflammation, promote apoptotic and necrotic cell clearance, engage tissue repair programs, and enhance immune cell-mediated containment of pathogens (1). Thus, proresolving therapeutics have the potential to dampen inflammation and stabilize lesions without compromising host defense. In pursuit of that goal, Arnardottir and Thul et al. provide evidence that GPR32 can potentially serve as a therapeutic target for resolving chronic atherosclerotic inflammation (7).

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Address correspondence to: Matthew Spite, Hale Building for Transformative Medicine, 60 Fenwood Rd., Room 3016K, Boston, Massachusetts 02115, USA. Phone: 617.525.5133; Email: mspite@bwh.harvard.edu.

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