Follicular regulatory T cell in atherosclerosis

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Abstract
Atherosclerosis is a chronic inflammatory disease involving the infiltration of immune cells, such as monocytes/macrophages, neutrophils, T cells, and B cells, into the inner layer of vessel walls. T and B cell functions in the process of atherogenesis, as well as their mutual regulation, have been investigated but several aspects remain to be clarified. In the present review, we give a brief overview of the functions of follicular regulatory T cell (Tfr) on follicular T (Tfh) and B cell regulation related to atherosclerosis pathogenesis, including their influence on lymphangiogenesis and lipoprotein metabolism. We will also discuss their potential therapeutics properties in the resolution of established atherosclerotic lesions.

KEYWORDS
Atherosclerosis, Regulatory B cells, B cell, Follicular T cell

1  |  INTRODUCTION
Cardiovascular diseases (CVD) are the leading cause of mortality worldwide.1 Atherosclerosis is the most studied CVD. Atherosclerosis is a chronic inflammatory disease characterized by the accumulation of lipids in the artery walls and involving both the innate and adaptive immune systems.2 The infiltration of proinflammatory monocytes in the artery walls in presence of elevated concentrations of circulating lipids triggers atherogenesis.3–5 During this complex process, circulating monocytes infiltrate the inner layer of vessel walls and differentiate into macrophages. This phenotypical switch allows the integration of lipids in excess in the artery walls leading to inflammation, foam cell formation, and establishment of plaques composed by the infiltrated elements, connective-tissue components, but also other particles.6,7 Over time, plaques enlarge and can break, causing thrombus formation, leading to ischemia, and myocardial infarction.8 Although the specific roles of T and B cell subtypes are not yet totally understood, their importance in the establishment and development of atherosclerosis has been clearly established.9–13 In this review, we will try to give an overview of the functions of immune cells in the regulation of atherosclerosis with particular emphasis on regulatory B cells (Breg), follicular T cell (Tfh), and follicular regulatory T cell (Tfr).

2  |  T CELLS IN ATHEROSCLEROSIS
During the development of atherosclerotic lesions, T cells are recruited at a very early stage.14 In the presence of low-density lipoprotein (LDL), adhesion molecules and chemoattractants located and expressed in the intima lead to the infiltration of T cells into the plaques.15,16 Once in the artery wall, antigens such as LDL and/or oxidized LDL activate the lymphocytes.17,18 T cells’ activation increases inflammation through proinflammatory cytokines release and creates an amplification loop between T cells and innate-immunity cells, such as macrophages.19

To study atherosclerosis, transgenic mice depleted of Apoe or ldlr genes are the most currently used models.20 In this context, Kyaw et al. determined that the majority of T cells present in atherosclerotic plaques are CD4+ T cells, although CD8+ T cells are present mainly in human and mouse advanced plaques21 with functions of atherosclerotic plaque destabilization.22 CD8+ T cells indeed trigger the apoptosis of macrophages, smooth muscle cells, and endothelial cells by its cytotoxic and inflammatory ability, leading to necrotic cores creation.22 CD8+ are also able to regulate Tfh23,24 known to be proatherogenic.25–27 CD4+ T cells have also been shown to have a proatherogenic function.28,29 Immunosuppressed Apoe−/− mice are atheroprotected, but adoptive transfer of CD4+ cells abolishes this
atheroprotective effect. Although T cells are crucial in the generation and sustenance of atherosclerosis, a strong heterogeneity is present in T cell subtypes (Table 1).

CD4+ effector T helper (Th) cells are divided in various subsets: Th1, Th2, Foxp3+ regulatory T (Treg) cells, Th17 cells, and Tfh cells. A new Treg cell subtype named Tfr has recently been described and provides a better understanding of Treg functions, particularly in autoimmunity. Similar to the Tfh population, this subset expresses the receptor CXCR5, which is important for the migration and the homing of Tfh. CXCR5 thus drives the localization of Tfr in the germinal-center (GC) in response to CXCL13. The characteristics of Tfr are also close to the Treg population. Tfr indeed expresses FoxP3 and CD25. By their regulatory function and their localization, Tfr cells negatively regulate Tfh population. To determine the exact function of T cells in atherogenesis, each subset needs to be thoroughly investigated.

Th1 cell population is widely present in plaques. Their ability to secrete proatherogenic cytokines certainly inhibits the atheroprotective action of other subsets. Th2 and Th17 cell populations are also found but their functions are still unclear. The functions of Treg cells in atherosclerosis have been largely investigated and show an overall atheroprotective role. It is also important to consider the possible atherogenic effect of T cell populations led by their potential ability to regulate cholesterol levels through the lymphatic system. It has been thus shown that areas with higher density of lymphatic vessels show less atherosclerotic plaques probably by their ability to stimulate a process called reverse cholesterol transport (RCT). In the context of cholesterol homeostasis, Treg cells seem to have opposite functions to other T cell subsets. In Apoe−/− mice and in humans with coronary artery disease, Treg cells are impaired, although they are known to initially increase in numbers in early lesions. Over the longer term, atherosclerosis is associated with a reduction of Treg cells. Furthermore, depletion or reduction of this subset leads to an aggravation of atherosclerosis: bigger lesion size, diminution of plaque stability, and change in lipid profile. The atheroprotective effects of Treg cells are, however, not only due to the modulation of cholesterol. Treg cells have proven to have a direct effect on endothelial cell activation, lymphocyte infiltration, foam cell formation, and on macrophage differentiation. During atherosclerosis, Treg cells can lose FoxP3 and mutate into memory, IFN-γ+ or Tfh cells. During this process, cells lose their ability to produce TGF-β1, a cytokine known to be atheroprotective and for its ability to modulate Tfh cell population. Lymphangiogenesis is also inhibited through Tfh-dependent IL-4 secretion. Diminishing Tfh cells and, in turn, IL-4 expression in mice increases the lymphangiogenesis. Due to their close phenotypical characteristics, there is no study in our knowledge in which Tfh cell population is investigated independently of Tfr cells. Gaddis et al. as well as our laboratory studied the effect induced by the inhibition of Bcl6, a transcription factor essential for Tfr and Tfh cells with disparate results. The different approaches used could, however, explain the discrepancy, Gaddis et al. have generated Bcl6-specific CD4+ knockout Apoe−/− mice. In this model, they observed a slight reduction of atherosclerotic plaques in mice under western diet whereas our study, based on a Bcl6 specific inhibitor, show an increase of atherosclerotic plaques. As Bcl6 is essential for the formation of GCs, the absence of GCs in Bcl6-depleted mice leads to a depletion of GC B cells, a cell type known to influence atherosclerosis. By suppressing Bcl6 with an inhibitor, our data are not influenced by the absence of GCs in Apoe−/− mice and results in an up-regulation of cholesterol levels and, thus, of atherosclerotic lesions. These findings are consistent with published data showing an increase of cholesterol level when Bcl6 is inhibited in mice and could be explained by the disruption of the lymphatic system during atherosclerosis. Bcl6-inhibitor used in our experiments is not strictly specific to Tfh cells and influence directly Tfr cells and other cell types such as macrophages. In this context, Bcl6 influences directly macrophages by inhibiting their proliferation and thus plaque enlargement could be due to an increased number of this cell subset. Our data show, however, no increase of CD68+ cells in atherosclerotic lesions. When Tfr cells are transferred into Apoe−/− mice, inversely, the number of macrophages present in the plaques and their size diminish. These results indicate that the atheroprotective effect of Tfr is certainly not only explained by its control of Tfh expansion. In fact, our data demonstrate that Tfr population stimulates directly lymphangiogenesis and Breg cells proliferation. They suggest also that Tfr cell population seems to have more significant functions in

| Cells types          | Breg | Tfh | Tfr | Refs |
|----------------------|------|-----|-----|------|
| Markers              | B220+ IgM+ | CD4+ PD-1+ | CD4+ PD-1+ | 33,35,61,69,84-87 |
|                      | CD43− CD1d+ | ICOS+ CXCR5+ | ICOS+ CXCR5+ |
|                      |       | CD25− FoxP3− | CD25+ FoxP3+ |
| Cytokines            | IL-10; IL-35; TGF-β | IL-4; IL-21 | IL-10; TGF-β | 61,86,88-91 |
| Localization         |       |       | 34,35,69 |
| Artery wall          | +    | ++   | +    |
| Spleen               | + +  | + +   | + +  |
| Lymph nodes          | + +  | Co-localized in the GCs |
| Effect on atherosclerosis | Atheroprotector | Atherogenic | Atheroprotector | 25-27,80 |
atherosclerosis than Tfh cell population. Finally, adoptive transfer of Tfr cells in mice with established atherosclerotic plaques is unable to decrease atherosclerosis despite the reduction of Tfh cell population. Thus, Tfr and Tfh cell populations are involved in atherogenesis but not in the sustainment or resolution of established atherosclerotic lesions.27

3 | B CELLS IN ATHEROSCLEROSIS

In regard of previous data on the functions of Tfh and Tfr cell populations, their particular role in B cell regulation have to be investigated. It is now well established that B cell dysfunction contributes to the development of autoimmune diseases through diverse mechanisms, such as autoantibody production or immune modulations. Although the exact function of B cell subsets in atherogenesis is still controversial and remains to be fully defined, B cell subsets are now more finely characterized. B cells are separated in three subtypes: B1, B2, and Breg. The B1 subset is composed of two populations: B1a (B220lowIgMhighCD43+CD5+) and B1b (B220lowIgMhighCD43−CD5−). Meanwhile, the B2 subset is composed by Marginal zone B2 (MZB) cells (B220IgMhighCD21highCD43+CD23−) and Follicular B2 (FOB) cells (B220IgMlowCD21+CD43−CD23+). The third subset is Breg (B220+CD43−IgMhighCD1dhigh).69,70 In mouse models, B cell populations have been identified at different stages of atherosclerotic lesions whereas, in humans, their presence is restricted to advanced plaques.11,71,72 Furthermore, splenectomy in mice has shown to enhance the development of atherosclerotic lesions, a process that is reversed by B cells adoptive transfer, suggesting an atheroprotective function of B cells.73,74 Similar to T cells, B cell populations seem, however, having both atheroprotective and atherogenic properties.73 Thus, the exact functions of independent B cell subsets required to be further investigated in atherogenesis. In this context, B1a and B1b cell populations have been shown to possess atheroprotective properties through IgM secretion.75,76 The effects of B2 cell population on atherosclerosis are unclear. It has been indeed demonstrated that B2 depletion in mice decreases atherosclerotic plaques whereas adoptive transfer aggravates the lesions.77 Inversely, MZB cells have been recently proven to protect against atherosclerosis plaque formation.24 Thus, within B2 cell populations, a distinction should be made between atherogenic FOB cells and atheroprotective MZB cells.78 Concerning Breg cells, their role in atherosclerosis is still debated. The controversy is mainly based on the function of IL-10 derived from Breg cell population in atherosclerosis.79,80 Although one of the phenotypical characteristics of Breg cells is their ability to produce IL-10, spontaneous ex vivo IL-10 production by B cells is difficult to observe. Identification of Breg population based on IL-10 production in vitro thus may potentially lead to misinterpretation.

4 | T AND B CELLS AND THEIR MUTUAL REGULATION IN ATHEROSCLEROSIS

The recent studies from our group indicate that Tfr cells exert part of their atheroprotective functions through the regulation of Breg cells differentiation.27 Our findings indicate also that Tfr cells control Breg cell population through cell-cell interaction. It thereby seems that Breg and Tfr cell populations are mutually self-regulated, leading to an amplification loop of regulation dampening Tfh cells frequency and, in turn, atherogenesis (Fig. 1). Consistently, whereas Tfh cell are the main producers of IL-4 in GCs, IL-4 is able to indirectly inhibit Breg cells expansion.81 The lack of Tfh cells regulation leads to nonantigen-specific B cells expansion, leading to autoimmune pathologies.35,82 The association of atherosclerosis and autoimmune diseases is correlated with the increase of the Tfh cell population during atherosclerotic plaque development.23,27 The number of Tfh cells present in secondary lymphoid organs of Apoe−/− mice is, however, reduced after the establishment of the lesions via a B cell-dependent process.24 In this context, we have demonstrated that MZB, Breg and Tfr cell populations increase during lesions development.27 FOB and Breg expansions are Tfr-dependent but not MZB, as demonstrated by adoptive transfer of Tfr cells.27 Consistently, Achour et al. have further demonstrated that human Breg cells control Tfh cells maturation, expand Tfr cells, and inhibit Tfh cell-mediated antibody secretion.53 These observations highlight the pleiotropic functions of both Tfr and Tfh cells in the modulation of anti- and proatherosclerotic immune processes in an Apoe−/− mouse model.
5 | CONCLUSION

The role of immune cells in atherosclerosis has already been well studied, particularly T and B cells. Recent discoveries of new lymphocyte subsets, however, require constant updating of our knowledge to have a better understanding of the disease. The ability of Tfr and Tfh cells to modulate atherosclerosis permitted to highlight the mutual regulation between B and T cells. Although further investigations are required in the field of immunomodulation of atherosclerosis, the atheroprotective properties of Tfr population through the control of Tfh cells, Breg cells, lymphangiogenesis, and lipid metabolism make Tfr cell population a key player of atherogenesis (Fig. 1). Moreover, the appearance of new T cell subsets spotlights the importance to further consider T and B cells subsets interactions, their localization and their functions, not only in the artery wall but also on the lymphatic system, all promising lines of research. More largely, the study of immune system and its potential outcomes will certainly participate to unravel the complicated multifactorial physiopathology of atherosclerosis.

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