

Galectin-1 induces macrophage immunometabolic reprogramming, modulates T cell immunity and attenuates atherosclerotic plaque formation

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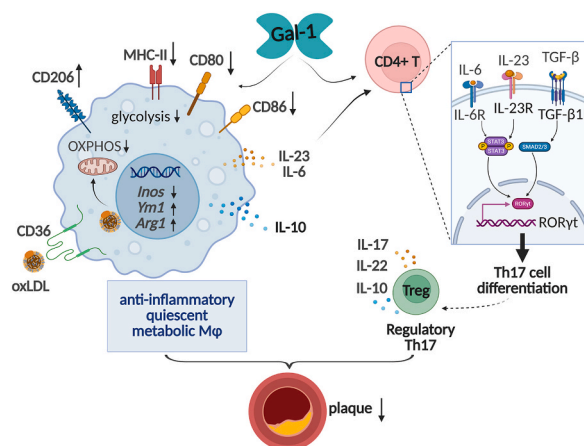
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HIGHLIGHTS

- Gal-1 treatment reduces early atherosclerosis independently of circulating IL-10 signaling.
- Gal-1 reprograms human and mouse macrophages in the plaque towards an anti-inflammatory phenotype.
- Anti-inflammatory effects of Gal-1 are partially independent of its glycan-binding ability.
- Gal-1 modulates T cell immunity by increasing Tregs and IL-10-producing T cells and by promoting a regulatory Th17 response.

GRAPHICAL ABSTRACT



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ABSTRACT

Background and aims: Atherosclerosis is a chronic immunometabolic disease driven by lipid accumulation and immune cell infiltration. Macrophages and T cells play key roles throughout plaque development. Galectin-1 (Gal-1), a glycan-binding protein, modulates immune functions in these cells and has been reported to attenuate atherosclerosis, though its mechanisms remain incompletely understood. Here, we investigated the effects of Gal-1 on macrophages and T cells during plaque formation.

Methods: Effects of Gal-1 on atherosclerosis, macrophages and T cells during lesion formation were studied in *Apoe*^{-/-} mice treated with recombinant Gal-1. Complementary mouse peritoneal foam cell and in vitro macrophage and T cell culture experiments were performed to study T cell differentiation, macrophage function, polarization and energy metabolism. The impact of Gal-1 on human macrophages was further evaluated in endarterectomy specimens.

Results: Gal-1 treatment reduced lesion size and increased circulating IL-10 levels, inversely correlating with plaque burden. Unexpectedly, IL-10 neutralization also mitigated atherosclerosis, indicating that its action is at least partially IL-10-independent. In plaques, Gal-1 promoted anti-inflammatory macrophage phenotypes, mirrored by a quiescent metabolic and anti-inflammatory profile in foamy macrophages ex vivo. The use of the Gal-1^{E71Q} variant revealed that these effects were only partly dependent on glycan binding. Beyond IL-10, Gal-1 reshaped cytokine profiles by increasing IL-17, IL-22, and IL-23, consistent with a macrophage-driven regulatory Th17 response, alongside higher frequencies of IL-10-producing and regulatory T cells.

Conclusion: Gal-1 protects against atherosclerosis associated with reprogramming macrophages and tuning T cell immunity through glycan-dependent and -independent pathways.

1. Introduction

Atherosclerosis, a chronic immunometabolic disease of the arterial wall, is the primary underlying cause of cardiovascular disease (CVD), which remains the leading cause of global death. Despite improvements in life expectancy, particularly in low- and middle-income countries the prevalence of CVD is expected to rise [1]. Central to the development and progression of atherosclerosis is the interplay between dyslipidemia and chronic inflammation, particularly the activation and infiltration of immune cells into the intimal layer of the arterial wall [2]. Macrophages and T cells, in particular, are pivotal players in plaque formation and contribute to the maladaptive immune response that drives disease progression [3].

Traditional therapies for atherosclerosis focus on lipid-lowering and thrombosis prevention, but recent clinical trials—such as the Canakinumab Anti-inflammatory Thrombosis Outcomes Study (CANTOS), the Colchicine Cardiovascular Outcomes Trial (COLCOT), and Low Dose Colchicine (LoDoCo)—have highlighted the therapeutic potential of targeting inflammation to reduce cardiovascular events [4]. However, the cost-benefit ratio of these approaches and substantial side effects

have hindered their widespread clinical adoption, underscoring the need for novel therapeutic strategies.

Among potential anti-inflammatory molecules, Gal-1, an endogenous glycan-binding protein of the galectin family, has emerged as a promising candidate with immunomodulatory effects on various immune cells, including macrophages, T cells, neutrophils and B cells [5–7]. By binding β -galactoside units on glycoproteins on cell surfaces and the extracellular matrix, Gal-1 regulates cell migration, activation, and apoptosis, thereby playing a critical role in all stages of inflammation—from initiation to resolution. Its bi- or multivalent nature enables cis or trans crosslinking of binding partners, further influencing these processes [8]. Beyond glycan binding, galectin-1 and -3 also interact with chemokines and intracellular transcription factors, such as forkhead box P3 (Foxp3), altering leukocyte migration and gene expression [9,10].

Modulation of monocyte/macrophage polarization towards an anti-inflammatory and pro-resolving phenotype has been reported [11]. Additionally, Gal-1 has been shown to inhibit leukocyte-endothelial interactions, further promoting an anti-inflammatory phenotype [12]. Notably, both pro- and anti-inflammatory effects of Gal-1 exist

depending on tissue context, intracellular or extracellular localization, pathologic conditions, and spatiotemporal expression of other regulatory programs [13].

Gal-1 has emerged as an important mediator of cardiovascular inflammation [14]. Circulating levels of Gal-1 are elevated following atherosclerotic events such as large artery stroke, and mouse models suggest that Gal-1 promotes post-stroke healing through interactions with neurons [15]. Moreover, Gal-1-deficient mice exhibit exacerbated cardiac inflammation and worsened vascular remodeling following myocardial infarction. It has been recently reported that Gal-1-deficient mice present with an increased atherosclerotic plaque burden and that exogenous Gal-1 attenuates atherosclerotic lesion development via effects on smooth muscle cells [16,17].

Given the critical roles of macrophages and T cells in atherosclerosis, and the emerging evidence that Gal-1 affects their biology, we sought to additionally investigate how Gal-1 influences immunometabolic reprogramming of macrophages and the immunity of T cells in the context of atherosclerosis.

2. Methods and materials

Comprehensive methods are available in the **Supplementary Information online**.

2.1. Animals and procedures

8-week-old *Apoe*^{-/-} mice (C57BL/6J background, Charles River Laboratories, France) were fed a Western diet (WD; 21 % fat, 0.2 % cholesterol, Ssniff, TD88137) for 8 weeks. They received intraperitoneal (i.p.) injections of either human recombinant Gal-1 (100 µg in 100 µL PBS, three times weekly) or PBS, as established previously [16]. In one setup, mice also received neutralizing IL-10 Ab (InVivoMAb, JES5-2A5, 500 µg i.p., three times a week during the first week and 300 µg twice a week thereafter), while controls received PBS or IL-10 Ab alone [18]. Feces were collected before sacrifice. Mice were euthanized under anesthesia (30 mg/kg xylazine, 150 mg/kg ketamine) following final blood sampling with subsequent perfusion and organ collection.

Immune cell and peritoneal foamy macrophage accumulation were assessed in additional experiments. Female *Apoe*^{-/-} mice were fed a WD for 4 weeks. Starting in week 3, mice received Gal-1 or Gal-1^{E71Q} (100 µg, three times weekly i.p.) or PBS. After the dietary run-in phase, mice were injected with 750 µL of 4 % thioglycolate. Four days post-injection, mice received a final injection of Gal-1, Gal-1^{E71Q}, or PBS, followed by an 8-h fasting period. Peritoneal lavage fluid was collected by washing the abdominal cavity with 10 mL ice-cold PBS containing 0.1 % BSA and 2 mM EDTA. In one setup, female *Apoe*^{-/-} mice were fed a WD for 4 weeks, and then mice were sacrificed and peritoneal lavage was collected.

All experiments were approved by local authorities (Regierung von Oberbayern, Germany; ROB-55.2Vet-2532.Vet_02-17-180, ROB-55.2-2532.Vet_02-22-150) and in compliance with the guidelines of Directive 2010/63/EU of the European Parliament on the protection of animals used for scientific purposes.

2.2. Expression and purification of Gal-1 and Gal-1^{E71Q}

Recombinant galectins were generated using pGEMEX-1 expression plasmid and the pGEX-6p2 as described [19]. A detailed protocol can be found in supplementary Information online.

2.3. Peritoneal macrophage experiment

Peritoneal lavage was collected from mice fed a WD as described above. Erythrocytes were lysed using ACK buffer. Peritoneal cells (1 × 10⁶ cells/well) were seeded onto 12-well cell culture plates and incubated with or without 100 nM Gal-1 for 24 h at 37 °C. Cells were then

collected and polarization of peritoneal macrophages (CD206, CD80, CD86, MHC-II) was assessed by flow cytometry.

2.4. Extracellular flux X24 seahorse measurement

Oxygen consumption rate (OCR) and extracellular acidification rate (ECAR) were measured on an XF-24 Extracellular Flux Analyzer (Agilent) (**detailed in Supplementary Information online**). Briefly, BMDMs or peritoneal macrophages (1 × 10⁵ cells/well) were seeded onto seahorse cell culture plates. Measurements were performed in triplicate. The following compounds were used: oligomycin (1.5 µM), fluoro carbonyl cyanide phenylhydrazone (FCCP, 1.5 µM), and rotenone + antimycin A (Rtn/AA, 2.5 µM) for the Mito Stress Test; glucose (10 mM), oligomycin (1.5 µM), and 2-deoxy-D-glucose (2-DG, 50 µM) for the Glycolysis Stress Test. After the readout, cells were lysed, and protein concentrations were determined using the BCA Protein Assay Kit (Thermo Scientific™ 23225). All metabolic rates were normalized to total protein levels.

2.5. RNA sequencing and analysis

RNA-sequencing was performed from inflammatory peritoneal macrophages using the prime-seq protocol [20]. Details on protocol and analysis can be found in the **Supplementary Information, Methods**.

2.6. Cholesterol and triglycerides measurement

Total cholesterol and triglycerides concentrations were determined using the enzymatic colorimetric method (Cholesterol-CHOD-PAP, Triglycerides-GPO-PAP, Roche Diagnostics) according to the manufacturer's instructions. Lipids were extracted from 100 mg of liver and feces with a chloroform/methanol solution, and dried pellets were dissolved in buffer (0.1 M potassium phosphate, pH 7.4, 0.05 M NaCl, 5 mM cholic acid, 0.1 % Triton X-100).

2.7. Statistical analysis

Statistical analyses were performed using GraphPad Prism version 10 (GraphPad Software, LLC). Data distribution was assessed for normality using the Shapiro-Wilk test, and lognormality was tested via logarithmic transformation. If the data were normally distributed or could be transformed into a normal distribution, parametric tests (paired or unpaired Student's t-test, or ANOVA for comparisons of more than two groups) were applied. For non-normally distributed data, the non-parametric Mann-Whitney test was used. One-way ANOVA was followed by Holm-Sidak's post hoc test for group comparisons. A two-tailed *p* value of <0.05 was considered statistically significant. Data are presented as mean ± standard error of the mean (SEM), unless otherwise specified.

3. Results

3.1. Administration of Gal-1 reduces atherosclerotic plaque area and plasma cholesterol levels

To assess the impact of Gal-1 treatment on atherosclerotic lesions, 8-week-old *Apoe*^{-/-} mice were fed a WD for 8 weeks while receiving three times weekly intraperitoneal injections of either 100 µg of human recombinant Gal-1 or PBS (Fig. 1A). Histological analysis of serial sections from the aortic root, spanning from the proximal to the distal regions, revealed a significant reduction in the absolute plaque area by approximately 40 % in Gal-1-treated mice compared to PBS-treated controls (Fig. 1B), well in line with a previously observed atheroprotective effect of Gal-1 in another model of atherosclerosis [16]. Similarly, the relative plaque area, calculated as the percentage of total plaque area within the three aortic leaflets compared to the total root area, was also

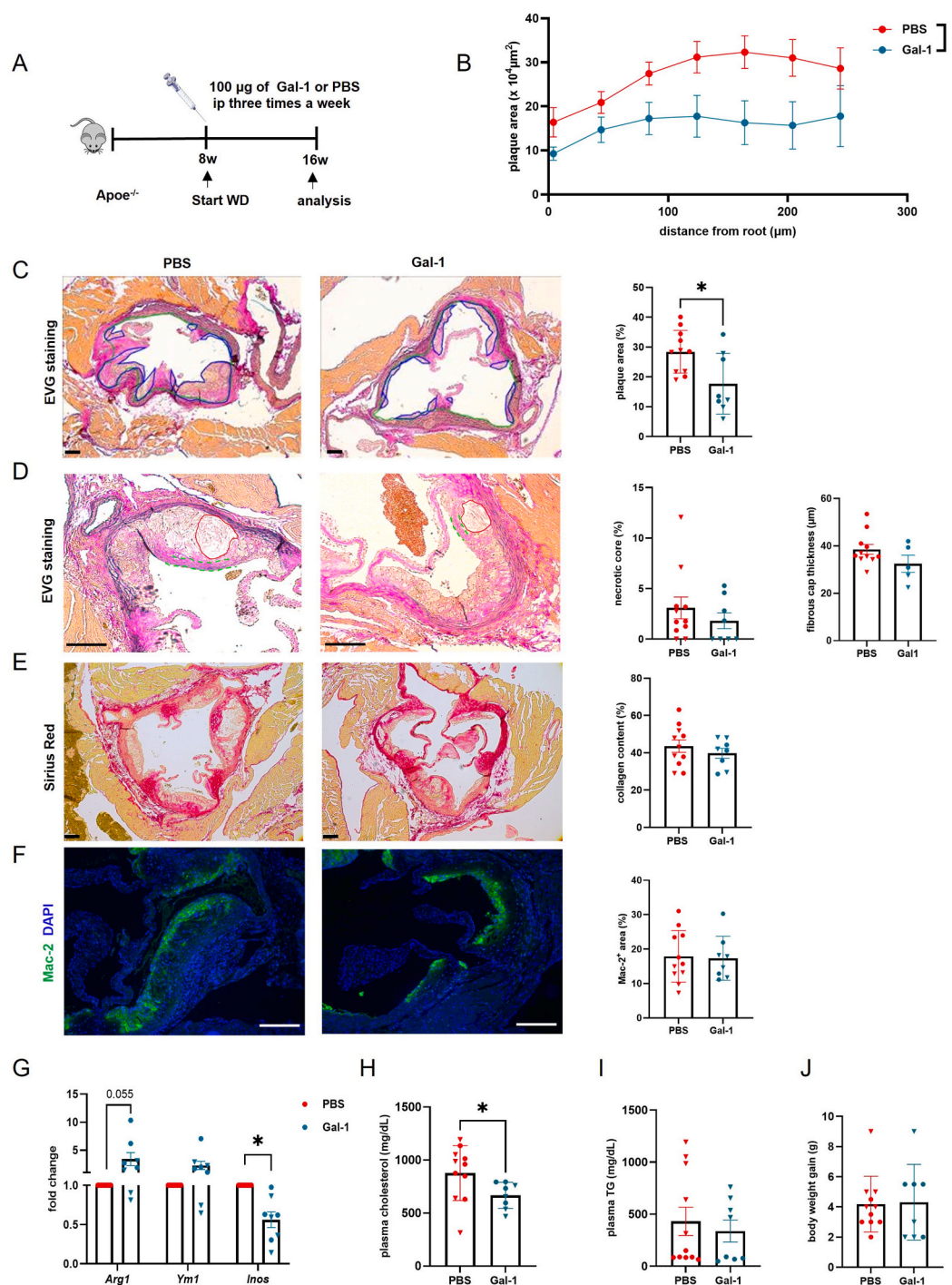


Fig. 1. Administration of Gal-1 reduces atherosclerotic plaque formation and plasma cholesterol levels.

(A) Experimental design. (B) Displayed is the plaque area from the proximal to distal part of the aortic roots from PBS- ($n = 11$, five females) and Gal-1-treated ($n = 8$, four females) $Apoe^{-/-}$ mice after 8 weeks on a Western diet (WD). The area under the curve was used to compare overall plaque burden. (C) Representative images of aortic roots stained with elastin/van Gieson (EVG) showing plaque boundaries (blue outline) and quantification of relative plaque area. (D) Representative images of aortic roots stained with EVG showing necrotic core (red outline), fibrous cap thickness measurements (dashed green outline), and their quantification. (E) Representative images of Sirius Red staining and quantification of collagen deposition in aortic root plaques. (F) Representative immunofluorescence images and quantification of macrophage (Mac-2⁺) infiltration relative to plaque area. (G) Expression of inflammatory genes related to macrophage profiling in the aortic arch of PBS- ($n = 8$) and Gal-1-treated mice ($n = 8$) after 8 weeks of WD. (H) Plasma cholesterol and (I) triglyceride (TG) levels. (J) Body weight gain after 8 weeks of WD. Data are expressed as mean $\pm$ SEM, with individual data points shown. Bullets represent female mice, and triangles represent male mice. * $p < 0.05$, two-tailed Student's t-test. Scale bar: 100 μ m. (For interpretation of the references to color in this figure legend, the reader is referred to the Web version of this article.)

significantly reduced in Gal-1-treated mice (Fig. 1C). Gal-1 did not show any sex-related differences in the plaque reduction; therefore, only female mice were used in the subsequent experiments. Necrotic cores, defined as hypocellular or acellular regions within the plaque, were

generally small at this early stage of atherosclerosis. Neither the size of the necrotic core, nor the thickness of the fibrous cap, the eosinophilic layer overlaying the necrotic core, were altered across groups (Fig. 1D). Furthermore, collagen content of the plaques revealed no changes

following the Gal-1 treatment (Fig. 1E). These findings indicate that Gal-1 treatment reduces atherosclerotic plaque size but does not alter plaque stability.

In terms of cellular composition, the area of lesional macrophages remained unchanged (Fig. 1F). Absolute CD3-positive T cells were reduced in the plaques of Gal-1-treated mice, while their frequency

relative to plaque area remained unaltered (Fig. S1A–C). To further characterize the plaque phenotype, the inflammatory status of the aortic arch was assessed. The anti-inflammatory marker *Arg1* tended to be upregulated, and pro-inflammatory *Inos* was significantly decreased in Gal-1-treated mice compared to PBS-injected mice, indicating a shift toward an anti-inflammatory milieu (Fig. 1G). These findings suggest

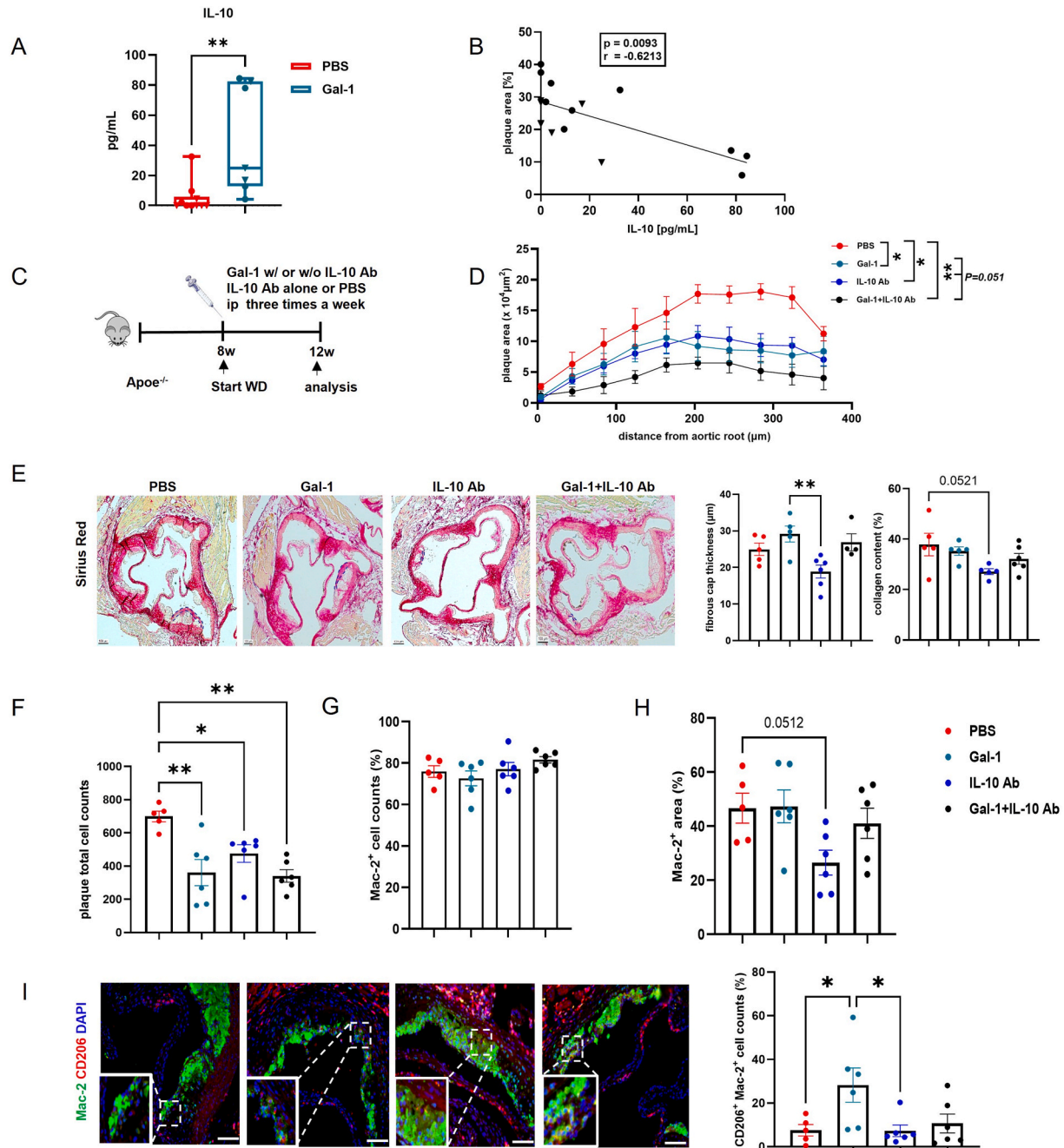


Fig. 2. Additive effects of IL-10 neutralization and Gal-1 in reducing atherosclerosis.

(A) Plasma IL-10 levels from PBS- (n = 10) and Gal-1-treated *Apoe*^{-/-} mice (n = 7) after 8 weeks of WD as shown in Fig. 1A. (B) Spearman's correlation between plasma IL-10 levels and aortic root plaque area. (C) Experimental design. Female *Apoe*^{-/-} mice were treated with PBS, Gal-1, IL-10 Ab (neutralizing IL-10 antibody, clone JES5-2A5), or Gal-1 + IL-10 Ab following 4 weeks of WD (PBS: n = 5, other groups: n = 6). (D) Displayed is the plaque area from the proximal to distal part of the aortic roots resulting in the area under the curve. (E) Stability of the plaque was assessed in Sirius Red-stained aortic root sections showing representative images, fibrous cap (FC) thickness measurements (dashed blue outline) and quantification of collagen content. FC was not quantifiable for two mice in the Gal-1 + IL-10 Ab group. (F) Plaque cellularity was quantified by counting DAPI-positive nuclei. (G–H) Percentage of macrophage numbers (DAPI⁺Mac-2⁺, G) and Mac-2⁺-area (H) is indicated per plaque area. (I) Representative immunofluorescence images and quantification of numbers of CD206⁺Mac-2⁺ macrophages relative to total plaque cells. (A) is expressed as box plots showing the interquartile range, median, and minimum to maximum, analyzed by Mann-Whitney *U* test. (A, B) Bullets represent female mice and triangles represent male mice. (D–I) are presented as mean ± SEM and analyzed by One-way ANOVA followed by Holm-Šidák's post hoc test. **p* < 0.05 and ***p* < 0.01. Scale bar: 100 μm. (For interpretation of the references to color in this figure legend, the reader is referred to the Web version of this article.)

that Gal-1 attenuates plaque development and promotes a less inflammatory plaque environment.

Consistent with a previous study showing that genetic deletion of *Lgals1*, the gene encoding Gal-1, results in elevated blood cholesterol levels, we observed a significant 24 % reduction in plasma cholesterol levels in Gal-1-treated mice (Fig. 1H) [16]. However, no significant differences were detected in triglyceride levels, body weight gain or body weight (Fig. 1I–J, Fig. S1D).

3.2. Additive effects of IL-10 neutralization and Gal-1 in reducing atherosclerosis

Consistent with the anti-inflammatory effects of Gal-1, we observed a significant increase in the potent anti-inflammatory cytokine IL-10 in the plasma of Gal-1-treated mice (Fig. 2A). Notably, plasma IL-10 levels negatively correlated with plaque size, with higher IL-10 levels being associated with smaller plaques (Fig. 2B). In line with this finding, Gal-1 has been reported to mediate immunomodulatory effects through IL-10, and IL-10 injections have also been reported to lower cholesterol levels in humans [18,21]. To determine whether blocking circulating IL-10 would negate the beneficial effects of Gal-1 on atherosclerosis and cholesterol levels, we administered a neutralizing IL-10 antibody, using a previously established experimental protocol, via intraperitoneal injection, combined with the Gal-1 treatment regimen for 4 weeks (Fig. 2C) [18].

After 4 weeks of WD, Gal-1 treatment significantly reduced atherosclerotic lesions, primarily in the aortic root and to a lesser extent in the aortic arch. Unexpectedly, blocking IL-10 did not exacerbate plaque formation; it led to a significant reduction in plaque size, and the combination of *anti*-IL-10 and Gal-1 resulted in a further numerical reduction, though non-significant, compared to either treatment alone (Fig. 2D, Fig. S2A).

There were no significant differences in the necrotic core size, fibrous cap thickness, or collagen content between PBS- and Gal-1-treated groups. In contrast, IL-10 blockade resulted in a thinner fibrous cap and reduced collagen content ($p = 0.05$), suggesting a shift toward a more unstable plaque phenotype (Fig. 2E, Fig. S2B).

Gal-1 treatment reduced plaque cellularity in line with the smaller plaque size, and cellularity was likewise reduced by *anti*-IL-10 alone or in combination with Gal-1 (Fig. 2F). Regarding macrophages, the proportion among total plaque cells remained similar across groups (Fig. 2G). However, the Mac-2 (Gal-3)-positive area was reduced only by *anti*-IL-10 treatment associated with a decrease in circulating monocytes (Fig. 2H, Fig. S2C). A plausible explanation for the decreased Mac-2-positive area in the *anti*-IL-10-treated group, despite unchanged macrophage numbers, is that *anti*-IL-10 treatment may have altered Gal-3 release, expression or reduced macrophage size, in line with a report showing that IL-10 can promote Gal-3 expression in a STAT3-dependent manner [22]. The number of anti-inflammatory macrophages (CD206⁺Mac-2⁺) per total plaque cells was increased following Gal-1 treatment (Fig. 2I) indicating that Gal-1 promotes an anti-inflammatory macrophage phenotype.

T cell numbers and lipid accumulation in the plaque, measured by the intracellular lipid droplet marker perilipin-2, were similar across groups (Fig. S2D and E).

Taken together, these results demonstrate that Gal-1 influences atherosclerosis through mechanisms that are not solely dependent on IL-10, while IL-10 neutralization reduces plaque size and promotes features of instability.

3.3. Gal-1 administration and IL-10 blockade reduce plasma cholesterol without affecting intestinal lipid absorption or liver lipid metabolism

In the 4-week trial, Gal-1 and/or IL-10 antibody treatment reduced plasma cholesterol by ~20 %, consistent with the 8-week results. The combination of Gal-1 and *anti*-IL-10 also reduced cholesterol, but the

effect was not additive (Fig. S3A). Given the association between IL-10 blockade and increased risk of enterocolitis and inflammatory bowel disease (IBD), we considered that the observed reduction could be attributed to decreased dietary cholesterol absorption or enhanced hepatic metabolism [23]. Throughout the trial, all mice showed similar weight gain (Fig. S3B and C), with no signs of diarrhea, bloody stools, or colitis. Stool cholesterol content, and triglyceride levels, were comparable across groups (Fig. S3D–F). Histological analysis of liver sections stained with Oil Red O (ORO) showed no lipid accumulation (Fig. S3G and H), and liver cholesterol and triglyceride levels were similar between groups (Fig. S3I and J). Additionally, hepatic expression of genes related to lipid metabolism showed no significant differences (Fig. S3K). Moreover, Gal-1 has been shown to bind to exosomes, and it is conceivable that it might also bind to lipoproteins, so that cholesterol depletion via an extracellular vesicle (EV)-mediated process is deemed possible [24]. Gal-1 addition to plasma caused precipitation, but the precipitate lacked detectable cholesterol and did not reduce total cholesterol levels compared to controls (Fig. S3L). These findings suggest that neither cholesterol resorption, altered hepatic metabolism, nor binding to extracellular vesicles contribute to the reduced plasma cholesterol levels in Gal-1 treated mice.

3.4. Effects of Gal-1 on energy metabolism and BMDM polarization are independent of oxLDL

We next hypothesized that Gal-1 might modulate lipoprotein uptake by macrophages and thereby influence plaque inflammation. When bone marrow-derived macrophages (BMDMs) were incubated with Gal-1 and oxLDL, spontaneous uptake of oxLDL occurred after 4 and 24 h, which was significantly enhanced by Gal-1 (Fig. 3A and B). Stimulation with oxLDL alone and combined with Gal-1 increased the gene expression of *Cd36*, and combination with Gal-1 significantly upregulated *CD36* surface expression (Fig. 3C and D), suggesting that under conditions where LDL is oxidized, Gal-1 enhances its uptake by macrophages, which may affect foam cell formation and plaque inflammation.

Recent studies indicate that cellular energy metabolism reprogramming is crucial for macrophage activation, influencing their phenotype and function. Glycolysis and mitochondrial respiration cannot be simply defined as pro- or anti-inflammatory [25]. To assess the impact of Gal-1 on energy metabolism, we measured oxygen consumption rate (OCR) and extracellular acidification rate (ECAR) in BMDMs treated for 24 h with Gal-1 and/or oxLDL (Fig. 3E–J). OxLDL reduced ATP-linked basal respiration (Fig. 3E), consistent with previous reports, and Gal-1 also decreased basal respiration to a similar extent [26]. The combination of Gal-1 and oxLDL had an additive effect, suggesting independent mechanisms. The maximal respiration rate, stimulated by FCCP, followed a similar pattern to basal respiration, suggesting that the effects are likely due to impaired substrate supply or oxidation rather than ATP synthesis (Fig. 3E) [27].

ECAR increased in response to electron transport chain inhibitors, a known compensatory response to mitochondrial oxidative phosphorylation (OXPHOS) inhibition (Fig. 3F), but oxLDL independent of Gal-1 attenuated this increase [28]. The OCR/ECAR ratio was reduced by Gal-1 but not by oxLDL, indicating that Gal-1 shifts the metabolic balance toward glycolysis (Fig. 3G). These findings were corroborated by the glycolysis stress test, where both Gal-1 and oxLDL reduced respiration, while inhibition of TCA/ATPase and glycolysis/hexokinase minimized OCR across all groups (Fig. 3H–J). The OCR/ECAR ratio in the glycolysis stress test mirrored the mitostress test, further indicating a shift towards glycolysis in response to both treatments.

Further assessment of polarization markers showed that oxLDL upregulated surface expression of proinflammatory CD86 and CD80, and Gal-1 downregulated CD86 expression in untreated BMDMs. However, Gal-1 did not affect oxLDL-induced CD86 and CD80, suggesting that Gal-1 does not influence oxLDL-induced polarization (Fig. 3K–L). Thus, atheroprotective effects of Gal-1 may occur independently of

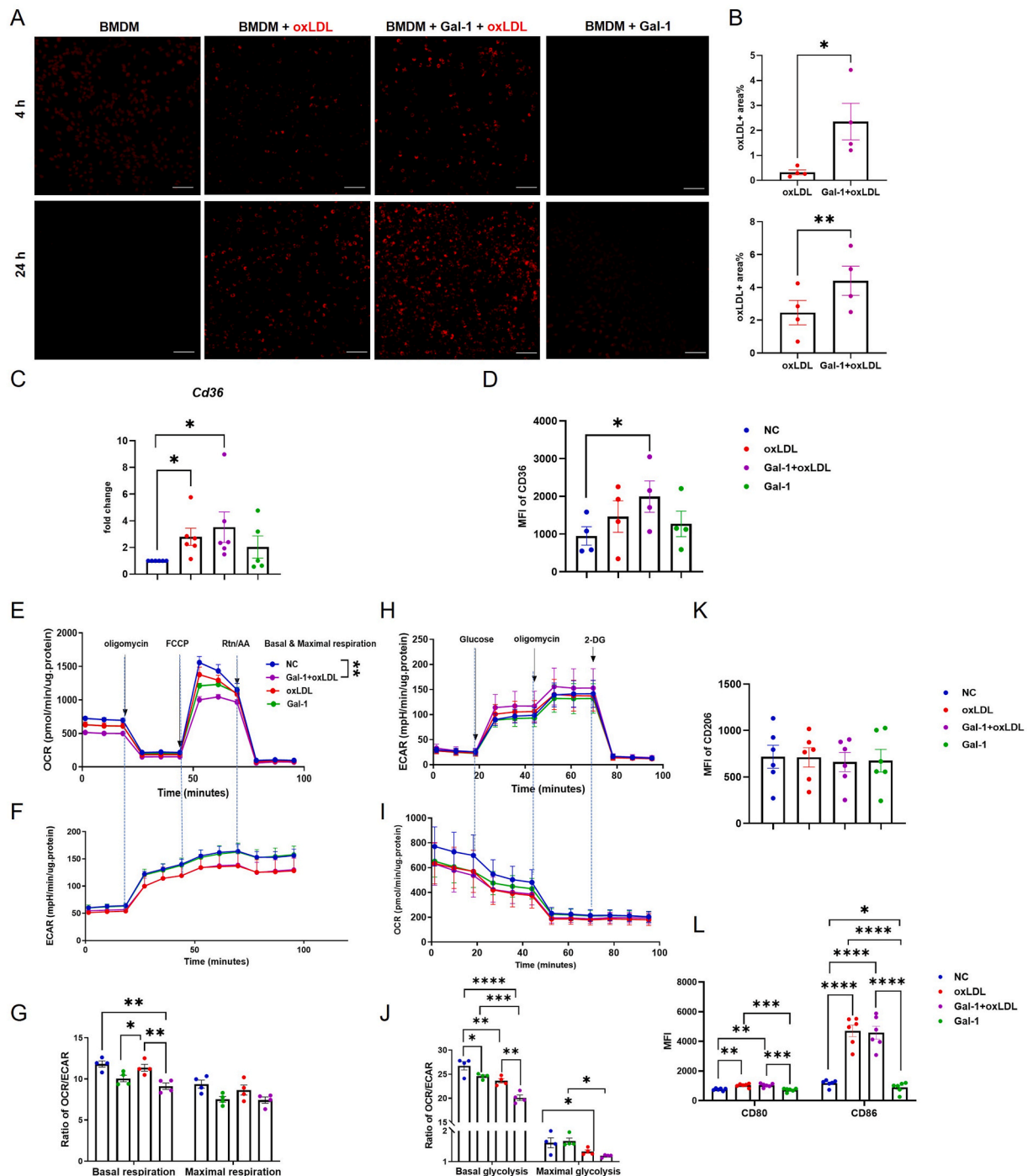


Fig. 3. Effects of Gal-1 and oxLDL on energy metabolism and BMDM polarization.

(A–B) BMDMs from *Apoe*^{-/-} mice were incubated for two distinct time periods (4 and 24 h) with unlabelled oxLDL (50 μ g/mL), with or without Gal-1 (100 nM). The formation of lipid droplets was then visualized through LipidSpotTM 610 staining. (A) The images were captured using a confocal microscope (20 $\times$ magnification), and (B) lipid droplets were quantified as percent positive area of all cells. (C) The expression of *Cd36* in BMDMs was analyzed by quantitative RT-PCR (n = 5–6) and (D) surface expression by flow cytometry. (E–J) Oxygen consumption rate (OCR) and extracellular acidification rate (ECAR) of BMDMs were measured by the Agilent Seahorse XF Flux analyzer. Mitochondrial respiration (E) OCR, (F) ECAR and (G) OCR/ECAR ratio measured at basal conditions and after addition of FCCP to induce maximal respiration. Glycolysis (H) ECAR, (I) OCR, and (J) their ratio measured at basal conditions and after addition of oligomycin (ATP synthase inhibitor) to drive maximal glycolysis. Flow cytometry of (K) CD206, (L) CD80 and CD86 in BMDMs treated as indicated for 24 h (n = 6). Data are presented as the mean $\pm$ SEM. * p < 0.05, ** p < 0.01, *** p < 0.001 and **** p < 0.0001. Statistical analysis: (B–C) two-tailed Student's t-test against NC (negative control), (D–L) one-way ANOVA followed by Holm-Sidak's post hoc test. Scale bar: 100 μ m.

macrophage activation by oxLDL, or BMDMs may not be the ideal model for investigating these effects.

These data together indicate that Gal-1 enhances oxLDL uptake in BMDMs, potentially contributing to plaque foam cell formation and

inflammation. However, Gal-1 failed to reverse the inflammatory profile induced by oxLDL, suggesting that its anti-atherogenic effects may be mediated through distinct inflammatory pathways. In addition, oxLDL-treated BMDMs as an entity in vitro may not fully recapitulate the

phenotype of lesional macrophages in vivo.

3.5. Gal-1 promotes an anti-inflammatory phenotype in peritoneal foamy macrophages

To test the response to Gal-1 in macrophages other than BMDM, we focused on lipid-laden inflammatory macrophages, which more closely resemble the cellular environment in atherosclerotic lesions compared to BMDM. These macrophages were isolated from the peritoneal cavity following intraperitoneal injection of thioglycolate, in the context of WD-induced hypercholesterolemia (Fig. 4A) [29]. In addition, we utilized a Gal-1 variant, Gal-1^{E71Q}, which exhibits a near-complete loss of its ability to bind glycans, thus preventing cell aggregation (Fig. S4A–D) [30]. Mice were initially fed a WD for 2 weeks and subsequently treated with Gal-1, Gal-1^{E71Q} or PBS for an additional 2 weeks. After the entire 4-week treatment period, there were no significant differences in bodyweight gain and triglyceride levels (Fig. S5A and B). Plasma cholesterol levels remained unchanged (Fig. S5C), implying that 2 weeks of treatment was too short to affect cholesterol levels.

In the peritoneum, Gal-1 treatment did not affect cell viability (Fig. 4B), and total peritoneal cell counts were similar across groups (Fig. 4C). However, significant differences in cell composition were observed (Fig. 4D). In controls, approximately half of the peritoneal cells were myeloid non-neutrophil cells (CD11b⁺Ly6G[−]) that predominantly consisted of inflammatory macrophages (F4/80⁺MHC-II^{hi}), with a low number of homeostatic macrophages (F4/80⁺MHC-II^{low}) and few monocytes. Treatment with Gal-1 and Gal-1^{E71Q} reduced the relative number of inflammatory macrophages, while increasing that of homeostatic macrophages (Fig. 4D and E). The Gal-1^{E71Q} mutant induced similar effects, although the changes were less pronounced compared to Gal-1, suggesting that glycan-binding plays only a minor role and is not the sole cause of the observed effects on the macrophages. Furthermore, the frequency of B cells was reduced (Fig. S5D), while those of T cells and neutrophils were increased (Fig. S5D and E). The increase in neutrophils is in line with a previous report showing that Gal-1 mediates neutrophil influx during late stages of peritoneal inflammation [31]. These effects were largely dependent on carbohydrate recognition, as the Gal-1^{E71Q} mutant behaved similarly to PBS control. There were no differences in the numbers of circulating leukocyte subsets between the groups (Fig. S5F and G). Although Gal-1 is known to induce (T cell) apoptosis [32], cell death cannot explain the observed effects given the comparable cell viability across groups, suggesting that Gal-1 may primarily influence macrophage proliferation and chemotactic factors. Collectively, these results indicate that Gal-1 treatment shifts the inflammatory profile of macrophages in an inflammatory environment, supporting an anti-inflammatory effect mediated by both sugar-binding dependent and independent mechanisms.

By accumulating lipids, macrophages exhibited foam cell-like characteristics (Fig. 4F). However, lipid staining and flow cytometric quantification revealed no significant differences in intracellular lipid content between groups (Fig. 4G), suggesting that Gal-1 does not influence peritoneal foam cell formation but rather reduces their inflammatory profile. This contrasts with findings in BMDMs, which may be attributed to differences in gene expression profiles between BMDMs and peritoneal macrophages in relation to foam cell formation [29]. Additionally, extracellular flux analysis showed that peritoneal macrophages from Gal-1-treated mice displayed a notable reduction in both OCR and ECAR but did not affect the OCR/ECAR ratio, suggesting a more quiescent metabolic phenotype, which might contribute to resolving inflammation (Fig. 4H).

3.6. Transcriptomic profiling and polarization effect of Gal-1 on macrophages

To gain a deeper insight into the transcriptomic changes regulated by Gal-1 in macrophages, we isolated peritoneal inflammatory

macrophages (F4/80⁺MHC-II^{hi}) from mice as described in Fig. 4A and conducted transcriptomic analysis by prime-seq [20]. We first investigated the effect of Gal-1 by analyzing differentially expressed genes (DEGs) between PBS- and Gal-1-treated macrophages. Out of 21,797 detected genes, we identified 450 DEGs, with 394 upregulated and 56 downregulated genes in the Gal-1-treated compared to the PBS-treated group (Supplementary Table S1). Among them, Gal-1 treatment upregulated the expression of genes associated with anti-inflammatory response (e.g., *Nlrp12*, *Rcor1*, *Trpc4*), engulfment functionality (e.g., *Trpc4*, *Marco*), and metabolism (e.g., *Suclg2*, *Dhhdh*, *Cpt1b*, *Cpt2*) while downregulating apoptosis-related genes (e.g., *Siah2*, *Birc6*, *Wdr73*) (Fig. 5A). Gene ontology (GO) analysis revealed enrichment in some inflammatory-related terms or pathways related to metabolism, such as Positive Regulation Of Transcription Of Notch Receptor Target (GO:0007221, *P* = 0.0037), SREBP Signaling Pathway (GO:0032933, *P* = 0.0076), Regulation Of Interleukin-18 Production (GO:0032661, *P* = 0.01), Regulation Of Phagocytosis (GO:0050765, *P* = 0.0343), Energy Reserve Metabolic Process (GO:0006112, *P* = 0.0481), or Negative Regulation Of I-kappaB kinase/NF-kB Signaling (GO:0043124, *P* = 0.0495).

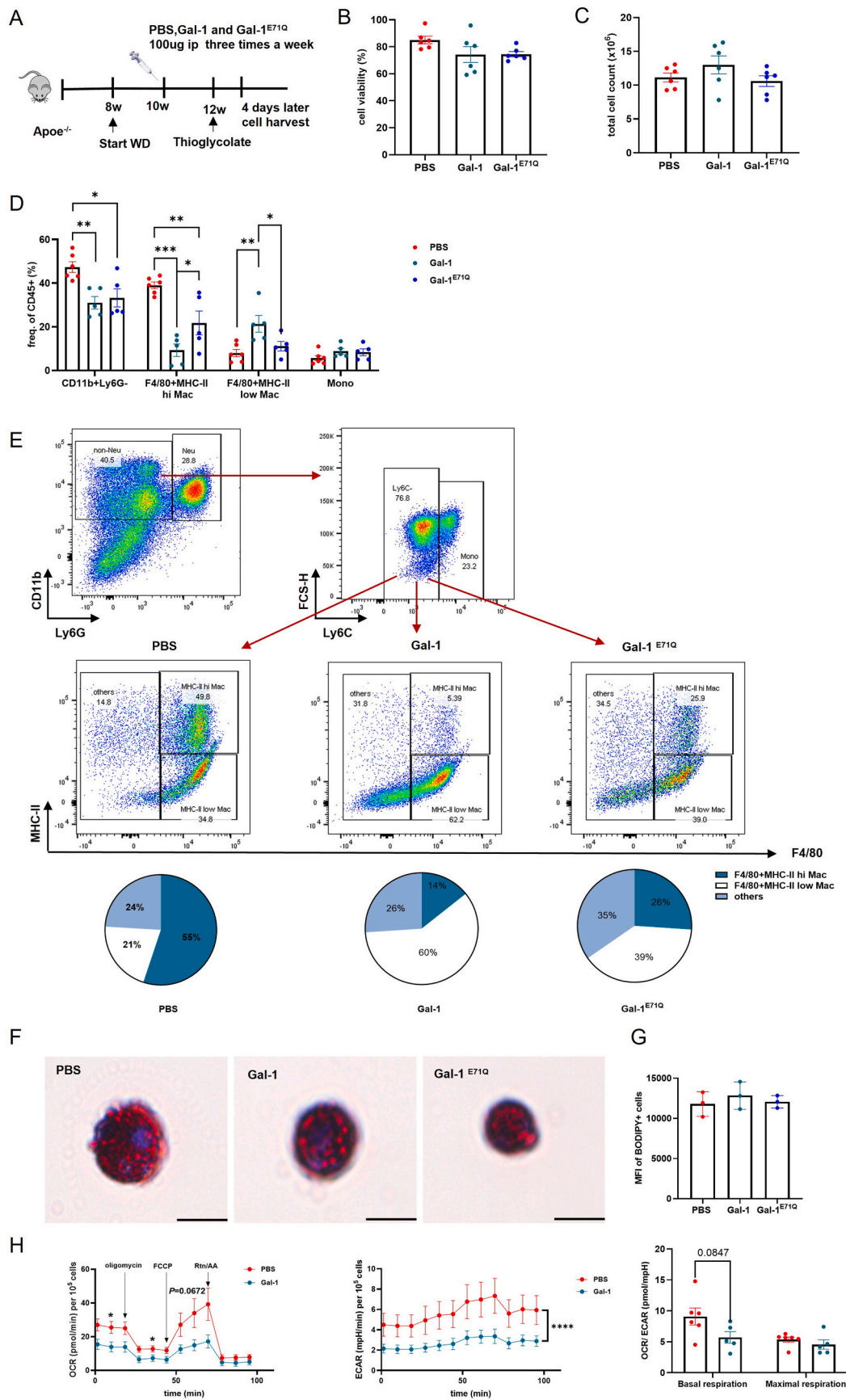
To dissect the role of glycan binding, we compared the DEGs in macrophages of mice treated with Gal-1^{E71Q}, unable to bind glycans. Fewer DEGs (91 upregulated, 10 downregulated) were detected upon Gal-1^{E71Q} compared to PBS treatment (Supplementary Table S2), with 70 genes commonly upregulated and 7 genes commonly downregulated by both Gal-1 and Gal-1^{E71Q} (Fig. 5B and Supplementary Table S3). These included genes relevant to regulating inflammatory response (e.g., *Rcor1*, *Slamf8*, *Pik3r2*) and the Notch signaling pathway (e.g., *Notch1*, *Notch2*), which has been implicated in the regulation of macrophage activation and polarization (Fig. 5B) [33]. This observation may help to explain the shifted pattern of peritoneal inflammatory macrophages observed in Gal-1- and Gal-1^{E71Q}-treated mice.

To further investigate the impact of Gal-1 on macrophage polarization, peritoneal cells were collected from mice after 4 weeks of WD (Fig. 5C) and cultured with Gal-1 or PBS for 24 h. Flow cytometry analysis revealed that Gal-1 treatment significantly reduced the expression of CD86 and CD80 on peritoneal macrophages. In addition, Gal-1 significantly downregulated Notch1 expression on macrophages and had no effect on CD206, MHC-II and Notch2 expression, suggesting an inhibition of pro-inflammatory macrophage phenotype (Fig. 5D–F). Given that Notch1 is implicated in atherogenesis by regulating macrophage polarization and has been shown to interact with Gal-1, we speculate that Gal-1 may facilitate anti-inflammatory macrophage polarization through Notch1 suppression, thereby contributing to its atheroprotective effects [33,34].

To further validate the polarization effect of Gal-1 on macrophages within atherosclerotic plaques, we cultured freshly isolated plaques obtained from femoral or carotid artery endarterectomy samples [35]. Fresh plaque was sliced into 2 mm thick sections and placed on a wetted sponge at the air-medium interface, followed by treatment with or without 100 nM Gal-1 for 24 h (Fig. 5G). To visualize the baseline plaque structure, one slice was further cryosectioned into 10 μm sections, and images were captured prior to treatment (Fig. 5H). After 24 h, the plaques were enzymatically digested and cell populations were analyzed by flow cytometry. While there was no noticeable change in the proportion of total macrophages (Fig. 5I), the expression of CD206 in macrophages was significantly upregulated following Gal-1 stimulation (Fig. 5J). However, the expression of MHC-II and CD163 remained unaltered (Fig. 5K and L). These observations further illustrate the anti-inflammatory action of Gal-1 in modulating macrophage inflammatory reprogramming within atherosclerotic plaques.

3.7. Gal-1 skews T helper balance and boosts Th17-Treg immunity

Gal-1 has been previously reported to modulate T-cell mediated immune responses [18]. To assess the impact of Gal-1 injection on T cell



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Fig. 4. Effects of Gal-1 on peritoneal foam cells.

(A) Experimental scheme of peritoneal foam cell induction. (B) Cell viability of peritoneal cells from PBS-, Gal-1-, and Gal-1^{E71Q}-treated female mice (n = 6 per group) was measured by Trypan blue staining. (C) Total number of cells in the peritoneal lavages per mouse was determined using an automated cell counter. (D) Frequency of myeloid cells were assessed by flow cytometry. (E) Flow cytometry gating strategy and corresponding pie chart depicting the relative abundance of CD11b⁺Ly6G⁺Ly6C⁺ cell subpopulations. (F) Representative images of peritoneal foamy macrophages, stained with Oil Red O (ORO). (G) Quantification of lipid droplets in peritoneal macrophages was obtained by flow cytometry after BODIPY staining (n = 3). (H) Extracellular flux analysis of peritoneal macrophages from PBS- and Gal-1-treated mice after 4 weeks of WD (n = 6). Mitochondrial respiration OCR, ECAR and OCR/ECAR ratio measured at basal conditions and after addition of FCCP to induce maximal respiration. Data are presented as the mean ± SEM. *p < 0.05, **p < 0.01 and ***p < 0.001. (A–G) one-way ANOVA followed by Holm-Šidák's post hoc test, (H) each phase was analyzed using two-tailed Student's t-test. Scale bar: 50 μm. (For interpretation of the references to color in this figure legend, the reader is referred to the Web version of this article.)

profiling in our atherosclerosis model, we conducted plasma cytokine analysis from mice, as shown in Fig. 1. However, no significant differences were observed for Th1- (IFN-γ, TNF-α) or Th2-related cytokines (IL-4, IL-5, IL-9, and IL-13) between PBS- and Gal-1-treated mice. Intriguingly, several cytokines associated with the Th17 immune response (IL-22, IL-23, IL-17, IL-6), which are typically linked to pro-inflammatory responses, were elevated in Gal-1-treated mice (Fig. 6A). This finding suggests that Gal-1 may exert complex immunomodulatory effects, potentially enhancing Th17 signaling pathways. Notably, concurrent IL-17 and IL-10 elevation may indicate a regulatory Th17 response associated with plaque reduction [36].

To investigate the cellular source of IL-10 production, we analyzed leukocyte and T-cell subsets in the blood. Gal-1 treatment significantly increased the percentage of IL-10-producing CD4⁺ T cells in the blood (Fig. 6B), while no changes were observed in inguinal lymph nodes (Fig. S6A). The increase in IL-10-producing CD4⁺ T cells was accompanied by an expansion of CD4⁺FoxP3⁺ regulatory T cells (Tregs) (Fig. 6C) and a significant increase in effector memory CD4⁺ T cells (Tem, CD44⁺CD62L⁺) (Fig. 6D) in the blood of Gal-1-treated mice. As both Tregs and effector memory CD4⁺ T cells can produce IL-10, their expansion likely contributes to the elevated frequency of IL-10-producing CD4⁺ T cells observed in Gal-1-treated mice [37]. However, the frequencies of naïve (CD44⁺CD62L⁺) and central memory (Tcm, CD44⁺CD62L⁺) CD4⁺ T cells, as well as other leukocyte subsets, remained unchanged (Fig. 6E–G). Leukocyte composition and CD4⁺ T-cell apoptosis in the spleen and lymph nodes were also comparable between groups (Fig. S6A and B).

To further elucidate the effects of Gal-1 on T cell differentiation, CD4⁺ T cells were isolated from the spleens of *Apoe*^{−/−} mice and cultured in vitro with or without Gal-1 for 72 h. After three days of culture, Gal-1 treatment significantly skewed T cell differentiation toward the Th17 subset, while the frequencies of Th2, Treg, and Th1 populations remained unchanged (Fig. 6H–L), indicating that Gal-1 preferentially promotes the differentiation of naïve T cells into Th17 cells. Analysis of cytokines in the supernatant revealed a marked increase in IL-10 production, while IL-17 levels remained unchanged in Gal-1-treated cells (Fig. 6M and N). These findings suggest that Gal-1 modulates the equilibrium between Treg and Th17 responses, potentially favoring a regulatory Th17 phenotype that may contribute to the attenuation of atherosclerosis.

4. Discussion

This study offers new insights into the role of Gal-1 in atherosclerosis, revealing its capacity to reduce atherosclerotic burden and promote less inflammatory plaque formation. These effects were associated with changes in macrophage immunometabolism and T cell immunity after Gal-1 treatment (Fig. 7). In macrophages, Gal-1 induced a more quiescent metabolic state, and reduced inflammatory activity. The use of the glycan-binding-deficient variant Gal-1^{E71Q} demonstrated that the anti-inflammatory effects on macrophages are mediated only partially through its carbohydrate-binding capacity. Furthermore, Gal-1 induced a distinct cytokine profile in blood, characterized by elevated levels of Th17 immune response associated cytokines. This was accompanied by an expansion of Tregs and a shift in T-cell differentiation towards the

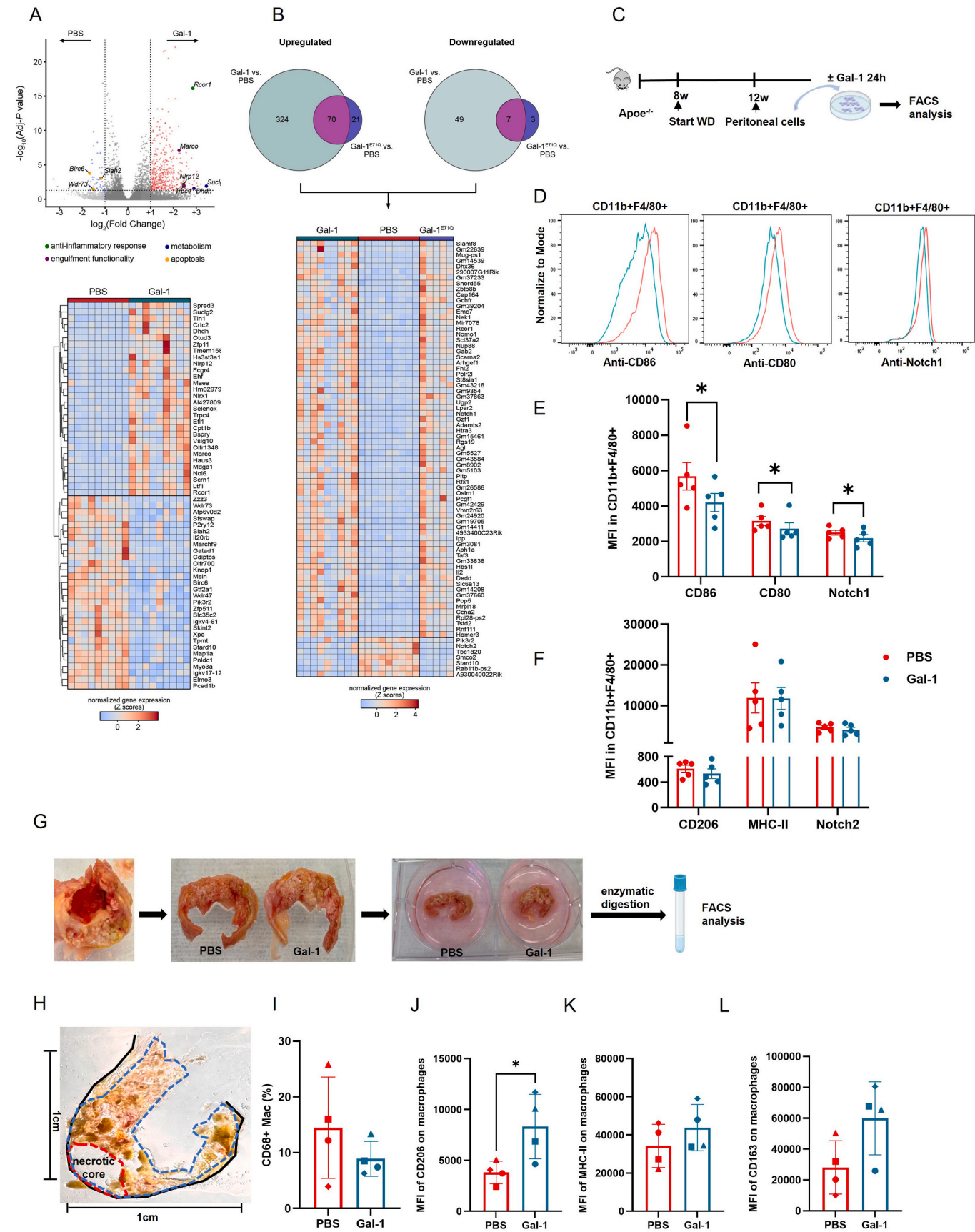
Th17 phenotype.

These findings further support and build upon previous research, demonstrating the atheroprotective effects of Gal-1 associated with smooth muscle cell functions, whereas the atherogenic impact on macrophages and T cells remains less clear [16].

Our study showed that Gal-1 treatment elevated plasma IL-10 levels and increased IL-10-secreting T cells, correlating with reduced plaque size. Gal-1 has previously been shown to enhance IL-10-mediated T cell tolerance and to suppress autoimmune neuroinflammation, with IL-10 widely recognized as a potent anti-inflammatory cytokine [38]. Yet the role of IL-10 in atherosclerosis remains controversial: deleting the IL-10 receptor in myeloid cells reduces lesion size and IL-10 boosts oxLDL uptake and foam cell formation [39,40]. We hypothesized that Gal-1 could reduce atherosclerosis in an IL-10-dependent manner, potentially explaining the reduced cholesterol levels we observed. However, neutralizing IL-10 with an established monoclonal antibody led to a reduction in lesions, and when combined with Gal-1, it failed to restore the reduced atherosclerotic burden. Instead, we observed an additive and thus mechanistically likely independent reduction. One explanation could be that the antibody treatment effects differ from the results of genetic models, as some compartments (e.g., intracellular) might not be antibody accessible. Therefore, if the functions of IL-10 are context-dependent, blocking only a portion of its activity may have led to the opposite effect seen with genetic deletion. Another key factor may be the cholesterol levels, which were reduced by both IL-10 Ab and Gal-1 but showed no additive effects when combined, suggesting a dependency between these two effects. Nevertheless, despite a reduction in plaque size, the neutralization of IL-10 promoted features of plaque instability that are more in line with the anticipated anti-inflammatory effects of IL-10.

Gal-1-treated mice exhibited lower plasma cholesterol levels after 4 weeks, though the underlying mechanisms are complex. Gal-1-deficient mice have been reported to downregulate PPARγ-dependent genes involved in gluconeogenesis and lipogenesis, with reduced liver triglyceride accumulation [41]. However, Gal-1-treatment did not affect cholesterol accumulation in the liver nor change the key lipid metabolism genes, suggesting liver-independent effects on cholesterol. Intestinal cholesterol absorption was not a factor either. Moreover, Gal-1 is known to bind extracellular vesicles [24], and hypothetically also lipoprotein particles but it did not precipitate lipoprotein particles in our study. Another possibility is enhanced oxLDL uptake by macrophages, as observed in BMDMs, which might have partially contributed to reduced plasma cholesterol levels. Possibly, several small factors are the reason which make the identification of a dominant pathway difficult.

Macrophage metabolic reprogramming plays a pivotal role in driving the inflammatory response, with evidence linking this process to oxLDL/CD36 signaling and is essential in the activation and polarization of macrophages, thereby influencing the progression of atherosclerosis [42]. Building on this, we further investigated the metabolic and inflammatory reprogramming of macrophages following Gal-1 treatment. Gal-1 enhanced oxLDL uptake in BMDMs possibly through upregulation of CD36. However, the effects of Gal-1 on energy metabolism were independent of oxLDL. Specifically, Gal-1 modestly reduced OXPHOS in BMDMs, accompanied by a shift toward glycolysis, and more profoundly reduced both respiration and glycolysis in peritoneal foam cells,



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Fig. 5. Transcriptional profiling and polarization effect of Gal-1 on macrophages.

(A) Volcano plot (upper) displaying the differentially expressed genes (DEGs, adjusted p value < 0.05 and $|\log_2FC| > 1$) of peritoneal macrophages from Gal-1-treated mice ($n = 9$) against PBS-treated mice ($n = 9$) as in Fig. 4. Each colored dot represents an upregulated (red) or a downregulated (blue) DEG. Genes relevant to Gal-1 biological processes are highlighted and labeled. The heatmap represents the top 30 upregulated/downregulated genes. The complete list of DEGs is provided in Supplementary Table S1. (B) Venn diagrams and the heatmap show the commonly upregulated and downregulated DEGs by both Gal-1 ($n = 9$) and Gal-1^{E71Q} ($n = 5$) vs. PBS ($n = 9$). The complete list of DEGs between Gal-1^{E71Q} and PBS is provided in Supplementary Table S2, the complete list of commonly regulated DEGs in Gal-1 and Gal-1^{E71Q} is found in Supplementary Table S3. (C–F) Polarization effect of Gal-1 on peritoneal macrophages was assessed by flow cytometry. (C) Scheme of experimental design. (D–F) Surface expression of polarization markers (CD86, CD80, MHC-II, CD206) and Notch receptors (Notch1 and Notch 2) on peritoneal macrophages (CD11b⁺F4/80⁺) treated with 100 nM Gal-1 or PBS for 24 h ($n = 5$). (G–L) Effect of Gal-1 on human plaque macrophages. (G) Experimental work-flow: Atherosclerotic plaques from carotid/femoral endarterectomies were sliced into 2 mm thick sections and cultured for 24 h with or without 100 nM Gal-1 treatment. Samples were then digested and analyzed by flow cytometry. (H) Representative image of human plaque section before culture. (I) The percentage of CD68⁺ macrophages from CD45⁺ cells. (J–L) Surface expression of CD206 (J), MHC-II (K) and CD163 (L) on macrophages was compared between groups. (I–L) Individual patients are represented by distinct symbols (●, ■, ▲, ◆). Data are presented as the mean $\pm$ SEM. * $p < 0.05$, two-tailed paired t -test on matched samples. (For interpretation of the references to color in this figure legend, the reader is referred to the Web version of this article.)

indicating a quiescent phenotype as seen in glioma metabolism [43]. The effect of Gal-1 on the energy metabolism might be essential for anti-inflammatory macrophage activation. Indeed, polarization markers indicated an alternatively activated phenotype mediated by Gal-1 in BMDMs and peritoneal foamy macrophages. Additionally, Gal-1 injection converted the high percentage of pro-inflammatory to anti-inflammatory macrophages, suggesting that Gal-1 inhibits pro-inflammatory macrophage programming. This immunometabolic profile of Gal-1-treated macrophages, together with the increase in anti-inflammatory macrophages in the plaques of Gal-1-treated mice, is consistent with previous research showing that Gal-1 supports a switch to a pro-resolving macrophage phenotype under inflammatory conditions [11,44]. The glycan-binding defect mutant Gal-1^{E71Q} yielded similar anti-inflammatory effects suggesting a partially glycan-independent mechanism [30]. Gal-3 can modulate inflammation via interaction with chemokines in a carbohydrate-binding independent manner [9]. A similar mechanism may apply to Gal-1, explaining the observed effects of Gal-1^{E71Q}. Interactions with chemokines may be one of many possibilities of how galectins can influence inflammation in a carbohydrate-binding independent manner.

The transcriptomic profile of inflammatory peritoneal foam cells from Gal-1 or Gal-1^{E71Q}-treated mice aligns with this. The number of differentially expressed genes was much higher in the Gal-1 group than in the Gal-1^{E71Q} group and comprised most of the genes regulated by Gal-1^{E71Q}. It remains speculative which of the pathways are responsible for the anti-inflammatory and immunometabolic effects on macrophages and atherosclerosis. We observed that some pathways, such as the Notch-signaling pathway, are commonly affected by both variants and are known to play an inflammatory role, further indicating the anti-inflammatory effects of Gal-1 through glycan-independent mechanisms.

On the other hand, T cells are strongly involved in atherogenesis, and Gal-1 is implicated in several T cell responses such as suppressing Th1 responses, promoting a Th2-skewed immune profile, and inducing T cell apoptosis [18,32]. However, our study did not observe significant alterations in Th2/Th1-associated cytokines or T cell apoptosis. Instead, we noted an increase in IL-10 production and a higher frequency of IL-10-secreting CD4⁺ T cells along with an expanded pool of effector memory T cells and Tregs, indicating that Gal-1 might modulate Treg function, which is consistent with studies showing that Gal-1-deficient mice exhibited reduced regulatory activity in CD4⁺CD25⁺ T cells that are endowed with atheroprotective functions [45,46]. The observed cytokine profile indicates that IL-17 release likely results from macrophage activation, which triggers IL-23 and IL-6 production, promoting Th17 differentiation and elevating IL-17 and IL-22 levels [47]. Consistent with this, Gal-1 enhanced the differentiation of in vitro cultured T cells into Th17 cells. These findings suggest that Gal-1 may modulate T cell responses at concentrations below its apoptotic threshold, allowing it to drive specific subsets without inducing cell death. Indeed, it has been reported that Gal-1 does not bind to naïve T cells, Th2 cells, or resting Tregs but strongly binds to Th17, and activated Tregs, where it induces apoptosis only at micromolar concentrations [48]. Direct effects

of Gal-1 on T cell differentiation into Th17 cells that are carbohydrate dependent can be mediated by CD69 have been reported [49].

Our results indicate that Gal-1 enhances a Th17 response. While IL-17 is generally regarded as a pro-inflammatory cytokine, its functional role—whether pro-inflammatory or homeostatic—appears context-dependent [50]. Notably, IL-17 has also been shown to stabilize atherosclerotic plaques and exhibit anti-inflammatory properties when combined with elevated IL-10 levels, as observed in our study [36]. The subset of T helper cells that produce both IL-10 and IL-17 has been described as regulatory Th17 cells, highlighting the balance between homeostatic and pathogenic IL-17-producing cells, as previously reviewed [36,51]. Furthermore, Th17 cells have been reported to transdifferentiate into Tregs during the resolution of inflammation [52]. It is therefore possible that we observed an intermediate stage in which IL-17-producing T cells are transdifferentiating into IL-10-producing T cells under the influence of Gal-1.

Several limitations of this study should be acknowledged. First, although Gal-1 treatment was associated with a reduction in plasma cholesterol, the precise mechanisms driving this systemic lipid-lowering effect remain unclear. However, the modest decrease in cholesterol levels suggests only a limited impact on atherosclerotic lesion development. In addition, while there was no difference in sex-based atherosclerotic plaque reduction after Gal-1 treatment in this study, the small sample size may have limited the statistical power to detect potential sex-related effects. Moreover, while our data indicate that Gal-1 exerts immunomodulatory effects that are at least partly independent of IL-10, and may influence T cell subsets such as Th17 and Treg populations, these observations require further validation. Finally, our investigation focused on immune cells. Other cell types such as smooth muscle cells may also contribute but were not investigated here. Despite these limitations, our findings demonstrate a link between Gal-1 treatment and immunomodulatory effects in the context of atherosclerosis, supporting its potential as an immunoregulatory therapy for this disease.

CRediT authorship contribution statement

Ya Li and Julian Leberzammer: Investigation, Methodology, Writing-original draft. Xavier Blanchet, Runduan Duan, Rui Su, Michael Lacy, Veit Eckardt, Yvonne Jansen and Yvonne Döring: Investigation, Methodology. Vasiliki Triantafyllidou and Donato Santovito: Software, Formal analysis, Visualization, Data Curation. Michael Hristov: Resources, Methodology. Eva Briem and Wolfgang Enard: Resources, Software, Project administration. Anna Simone Jung, Joel Guerra and Alexander Bartelt: Resources, Investigation. Jürgen Bernhagen, Christian Weber and Dorothee Atzler: Resources, Funding acquisition. Herbert Kaltner and Anna-Kristin Ludwig: Resources, Conceptualization. Philipp von Hundelshausen: Conceptualization, Funding acquisition, Supervision, Writing-review & editing.

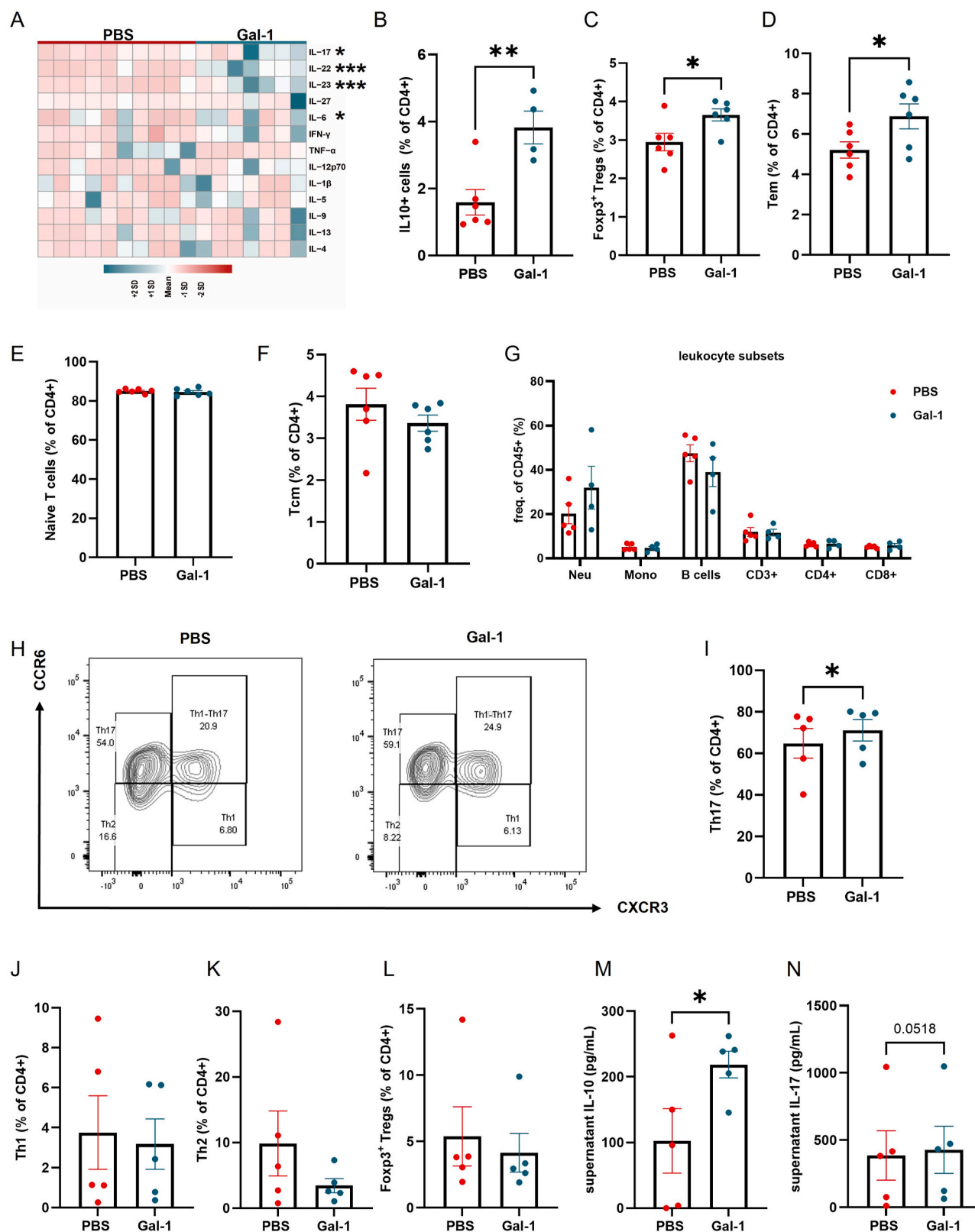


Fig. 6. Plasma cytokine profiling and immune cell analysis.

(A) Heatmap representing plasma multiplex cytokine analysis from PBS- (n = 10) and Gal-1-treated *Apoe*^{-/-} mice (n = 7). (B–F) Flow cytometry was used to quantify the proportions of distinct T cell subsets within the CD4⁺ T cells in blood. (B) IL-10-secreting CD4⁺ T cells, (C) regulatory T cells (Treg: CD4⁺Foxp3⁺), (D) effector memory T (Tem: CD44⁺CD62L⁻), (E) naïve T (CD44⁺CD62L⁺), (F) and central memory T cells (Tcm: CD44⁺CD62L⁺) cells (n = 4–6 per group). (G) Proportions of distinct leukocyte subsets within CD45⁺ cells in blood. (H–N) CD4⁺ T cells were isolated by negative selection from spleens and cultured for 72 h with 50 nM Gal-1 or PBS. (H) Gating strategy for T helper subsets, (I) percentage of Th17 (CCR6⁺CXCR3⁺), (J) Th1 (CCR6⁻CXCR3⁺), (K) Th2 (CCR6⁻CXCR3⁺), (L) Treg (CD4⁺Foxp3⁺). (M) IL-10 and (N) IL-17 levels in the supernatant were measured using Elisa (n = 5 per group). (A) Mann-Whitney U test. Data in (B–N) are shown as mean $\pm$ SEM. **p* < 0.05, ***p* < 0.01 and ****p* < 0.001, (B–G) two-tailed Student's t-test. (I–N) two-tailed paired Student's t-test on matched samples.

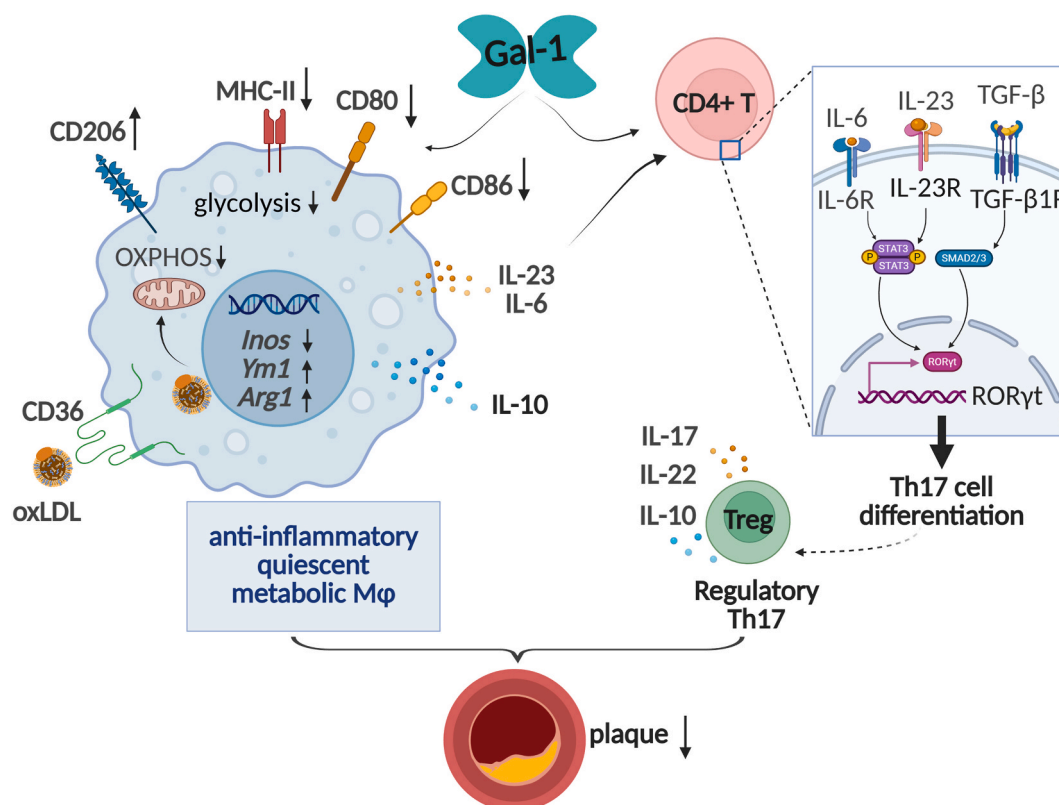


Fig. 7. Gal-1 reprograms macrophage metabolism and shapes T cell responses to reduce atherosclerosis.

Gal-1 acts on macrophages to induce an anti-inflammatory, metabolically quiescent phenotype characterized by increased CD206, reduced MHC-II, CD80, and CD86 expression, decreased glycolysis and OXPHOS, and altered gene expression (↓*Inos*, ↑*Ym1*, ↑*Arg1*). Gal-1 also modulates CD4⁺ T cell responses by shaping cytokine signals that promote a regulatory Th17 profile with increased IL-17, IL-22, and IL-10 supporting a regulatory phenotype. The combined effects of macrophage reprogramming and T cell modulation likely contribute to reduced plaque development.

Ethics approval

All animal procedures were performed conforming to the guidelines from Directive 2010/63/EU, die Regierung von Oberbayern, Germany; (ROB-55.2Vet-2532.Vet_02-17-180, ROB-55.2-2532.Vet_02-22-150), the ARRIVE guidelines and the ethical standards laid down in the 1964 Declaration of Helsinki and its later amendments.

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Conflict of interest

The authors have no financial or non-financial interests to disclose.

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C.W. is a Van de Laar professor of atherosclerosis. The graphical abstract was created with [BioRender.com](https://www.biorender.com/).

Appendix A. Supplementary data

Supplementary data to this article can be found online at <https://doi.org/10.1016/j.atherosclerosis.2025.120608>.

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