Liraglutide Attenuates Preestablished Atherosclerosis in Apolipoprotein E–Deficient Mice via Regulation of Immune Cell Phenotypes and Proinflammatory Mediators


School of Biomolecular and Biomedical Science (R.B., O.B.) and School of Medicine (S.C., S.K., E.P.B., F.C.M.), Diabetes Complications Research Centre, and Mass Spectrometry Resource (E.T.D.), University College Dublin (UCD) Conway Institute, UCD, Belfield, School of Public Health, Physiotherapy and Sports Science, UCD, Belfield (G.L., M.E.O’R.), and Vascular Surgery, St. Vincent’s University Hospital (M.B., S.S.), Dublin, Ireland

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ABSTRACT

We have shown that the glucagon-like peptide-1 receptor agonist (GLP-1RA) liraglutide (Lir) inhibits development of early atherosclerosis in vivo by modulating immune cell function. We hypothesized that Lir could attenuate pre-established disease by modulating monocyte or macrophage phenotype to induce atheroprotective responses. Human atherosclerotic plaques obtained postendarterectomy and human peripheral blood macrophages were treated ex vivo with Lir. In parallel, apolipoprotein E–deficient (ApoE–/–) mice received a high-fat, high-cholesterol diet to induce atherosclerosis for 8 weeks, after which ApoE–/– mice received 300 µg/kg of Lir daily or vehicle control for a further 4 weeks to investigate the attenuation of atherosclerosis. Lir inhibited proinflammatory monocyte chemotactant protein-1 secretion from human endarterectomy samples and monocyte chemoattractant protein-1, tumor necrosis factor-α, and interleukin (IL)-1β secretion from human macrophages after ex vivo treatment. An increase in CD206 mRNA and IL-10 secretion was also detected, which implies resolution of inflammation. Importantly, Lir significantly attenuated pre-established atherosclerosis in ApoE–/– mice in the whole aorta and aortic root. Proteomic analysis of ApoE–/– bone marrow cells showed that Lir upregulated the proinflammatory cathepsin protein family, which was abolished in differentiated macrophages. In addition, flow cytometry analysis of bone marrow cells induced a shift toward reduced proinflammatory and increased anti-inflammatory macrophages. We concluded that Lir attenuates pre-established atherosclerosis in vivo by altering proinflammatory mediators. This is the first study to describe a mechanism through which Lir attenuates atherosclerosis by increasing bone marrow proinflammatory protein expression, which is lost in differentiated bone marrow–derived macrophages. This study contributes to our understanding of the anti-inflammatory and cardioprotective role of GLP-1RAs.

SIGNIFICANCE STATEMENT

It is critical to understand the mechanisms through which liraglutide (Lir) mediates a cardioprotective effect as many type 2 diabetic medications increase the risk of myocardial infarction and stroke. We have identified that Lir reduces proinflammatory immune cell populations and mediators from plaque-burdened murine aortas in vivo and augments proresolving bone marrow–derived macrophages in attenuation of atherosclerotic disease, which provides further insight into the atheroprotective effect of Lir.

Introduction

Atherosclerosis is a chronic progressive disease that is characterized by the accumulation and deposition of lipids and fibrous elements coupled with an inflammatory response resulting in the development of an arterial atherosclerotic lesion (Ross, 1993). The earliest clinical hallmark of a developing lesion is the accumulation of lipid-laden macrophages known as foam cells, which aggregate to form the “fatty streak” (Gerhard and Duell, 1999). Fatty streaks are precursors for advanced lesions, where clinical events stem from lesion rupture or erosion and acute occlusions from thrombus formation, which clinically manifest as a myocardial infarction or stroke (Lusis, 2000).

In addition to established risk factors, such as hyperlipidemia and hypertension, it is now accepted that diabetes
mellitus–associated hyperglycemia and oxidative stress contribute to accelerated-atherosclerosis by modulating monocyte and macrophage function (Bornfeldt and Tabas, 2011). Work over the past decade has identified the importance of monocyte and macrophage cells in atherosclerotic plaque formation. In a simplified model, macrophages are classified as proinflammatory (M1) or anti-inflammatory (M2). M1 macrophages are induced by inflammatory cytokines and sustain the ongoing inflammatory response via generation of tumor necrosis factor-α (TNF-α). M2 macrophages are induced by interleukin (IL)-10 and IL-4/IL-13 and promote tissue repair and healing (Fujisaka et al., 2009). M1 and M2 macrophages show plasticity in response to stimuli from the microenvironment and have functional roles in other disorders of chronic low-grade inflammation such as obesity and atherosclerosis (Fujisaka et al., 2009). We have previously shown there is an M2-to-M1 switch during atherosclerotic plaque progression in human patients (de Gaetano et al., 2016), and it is now known that type 2 diabetes mellitus is characterized by a reduction in M2 populations, which is supported by in vivo studies, showing that a shift in M1:M2 ratio directly correlates to the development of insulin resistance (Ye et al., 2016). Stabilizing the progression of cardiovascular disease (CVD) remains a major target in people with type 2 diabetes mellitus. Indeed, recently, there has been increased emphasis on identifying the CVD safety and benefit of recently developed glucose-lowering agents.

Increasing evidence suggests that glucagon-like peptide-1 (GLP-1) therapies, including liraglutide (Lir), a GLP-1 receptor agonist (GLP-1RA), may simultaneously impact the pathogenesis of both type 2 diabetes mellitus and CVD (Arakawa et al., 2010; Gaspari et al., 2011; Nagashima et al., 2011; Rizzo et al., 2015). Dipeptidyl peptidase-4 inhibitors and GLP-1RAs inhibit monocyte and macrophage inflammatory responses and accumulation resulting in the attenuation of atherosclerotic lesion progression in apolipoprotein E knockout (Apoe deficient) mice (Arakawa et al., 2010; Nagashima et al., 2011) and improve endothelial cell function and reduce adhesion marker expression (Gaspari et al., 2011). Lir also modulates inflammation and reduces oxidative stress in diabetic patients (Rizzo et al., 2015). In addition, GLP-1RA decrease proinflammatory macrophages in obese type 2 diabetic patients (Hogan et al., 2014) and promote M2 polarization in human monocytes from healthy volunteers (Shiraishi et al., 2012).

We have previously reported that regression of pre-established atherosclerosis in vivo is associated with increased M2 macrophage polarization and IL-10 secretion in the murine aorta (McCarthy et al., 2013). Recently, we have shown Lir can modulate the development of early atherosclerosis in Apoe deficient mice by promoting a proresolving macrophage phenotype (Bruen et al., 2017), which implies that Lir may protect against macrovascular complications of diabetes. To date, a limited number of studies have examined the effect of Lir on immune cell phenotypes in the context of regression of pre-established atherosclerosis. This is important because most patients who have established disease at diagnosis of atherosclerosis, with evident cellular and clinical plaques, and the therapeutic goal is to attenuate progression or induce resolution of the disease. This present study is the first to show that Lir stabilizes plaque progression and may induce regression of pre-established atherosclerosis in vivo. In addition, through phenotypic and proteomic analysis of immune cell populations, it provides mechanistic insight into the atheroprotective effect of the GLP-1RA (i.e., Lir) in established atherosclerotic disease.

### Materials and Methods

#### Human Studies

Studies were approved by St. Vincent’s University Hospital, Dublin Ethics Committee, and adhered to international guidelines and the Declaration of Helsinki principles as revised in 2008. All participants provided informed written consent. Human endarterectomy samples were obtained from consenting patients of mixed sexes postvascular surgery and sectioned into relatively disease-free (RDF) and diseased plaque (DP) portions. Plaque portions were cultured in RPMI medium, supplemented with 10% fetal bovine serum, 1% L-glutamine, and 100 U of penicillin-streptomycin (P-S) (Bio-sciences, Dublin, Ireland) as described previously (Érbel et al., 2014; Brennan et al., 2018). Plaque sections were treated ex vivo with 1 μM Lir (Novo Nordisk, Dublin, Ireland) or phosphate-buffered saline (PBS) (Bio-sciences) as a vehicle control (VC) for 28 hours. Peripheral blood mononuclear cells (PBMCs) were isolated from whole blood by layering diluted blood in PBS (1:1) onto lymphoprep and centrifuging for 25 minutes at 400g. The PBMC layer was removed and centrifuged for 5 minutes at 800g to remove platelets. Cells were resuspended in 10 μL of M199 medium (Anablu, Dublin, Ireland), supplemented with 10% human serum (Sigma-Aldrich, Wicklow, Ireland) and 100 U of P-S and plated on 10-cm Petri dishes for 2 hours. Adherent cells after 2 hours were considered monocytes, and these cells were counted and reseduced in 12-well plates at a density of 1 × 10⁶ cells/ml M199 medium containing 100 ng/ml of monocyte-colony stimulating factor (Biologend, London, UK). Cells were treated with 250 nM Lir or PBS VC for 6 hours in the presence or absence of 100 ng/ml of lipopolysaccharide (LPS) (InvivoGen, Ireland) for 4 hours and 5 mM ATP (Sigma-Aldrich) for the final hour of treatments for inflammasome activation. Enzyme-linked immunosorbent assay (ELISA) was carried out on plaque and cell supernatants, and quantitative real-time polymerase chain reaction (qRT-PCR) was performed on differentiated PBMCs.

#### Animals

The Animal Research and Ethics Committee, University College Dublin, and the Health Products Regulatory Agency of Ireland approved all protocols; these protocols adhered to institutional and international guidelines. Eight-week-old C57BL/6J ApoE−/− male mice (C57BL6J-ApoE<sup>−/−</sup>, 002052; RRID:IMSR_TAC:apoe; Charles River, Margate, UK) were housed in specific pathogen-free conditions in 12-hour light/dark cycles. ApoE<sup>−/−</sup> mice were randomized to receive a high-fat (60% kcal from fat), high-cholesterol (1%) diet (HFFCVD) (Research Diets Inc, New Brunswick, NJ) for 8 weeks (n = 10 per group)

### Abbreviations

- ApoE<sup>−/−</sup>: apolipoprotein E knockout
- BMDM: bone marrow-derived macrophage
- Cat, cathepsin
- CCR2, CC chemokine receptor 2
- Ct, cycle threshold
- CVD, cardiovascular disease
- DC, dendritic cell
- DP, diseased plaque
- ELISA, enzyme-linked immunosorbent assay
- GAPDH, glyceraldehyde-3-phosphate dehydrogenase
- GLP-1, glucagon-like peptide-1
- GLP-1RA, GLP-1 receptor agonist
- HFFCVD, high-fat, high-cholesterol diet
- IL, interleukin
- INOS, inducible nitric oxide synthase
- LFD, low-fat diet
- LFD, low-fat diet
- LFQ, label-free quantification
- Lir, liraglutide
- LN, lymph node
- LPS, lipopolysaccharide
- MCP-1, monocyte chemoattractant protein-1
- MS, mass spectrometry
- MS/MS, tandem mass spectrometry
- PBMC, peripheral blood mononuclear cell
- P-S, phosphate-buffered saline
- P-S, penicillin-streptomycin
- qRT-PCR, quantitative real-time polymerase chain reaction
- RDF, relatively disease free
- TNF-α, tumor necrosis factor-α
- VC, vehicle control
to establish disease. For analysis of established disease, 8-week-old ApoE<sup>−/−</sup> mice were fed a low-fat (10% kcal from fat), 0% cholesterol diet (LFD) or HFD for a total of 12 weeks, in which mice were randomized to receive the HFD and received once daily Lir subcutaneous injections from weeks 9 to 12, titrated upward for 10 days (1, 3, 10, 30, 50, 100, 150, 200, 250, and 300 µg/kg) and maintained at 300 µg/kg (n = 21 per group) or VC PBS from weeks 9 to 12. Mice fed LFD received daily subcutaneous PBS injections from weeks 9 to 12 as a control. Mice were scored daily and euthanized under isoflurane (Duggan Veterinary, Tipperary, Ireland) by a terminal retro-orbital bleed and cervical dislocation. Aortae, lumbar, and inguinal lymph nodes (LN), spleen, and bone marrow were harvested.

The unit of analysis was a single animal or organ. Aortae were analyzed by en face staining (n = 8 per group) or cultured ex vivo for 6 hours with aortic tissue analyzed by qRT-PCR (n = 5–7 per group), and supernatants were analyzed by ELISA (n = 7 per group). Hearts were sectioned for aortic root analysis of atherosclerotic lesions by Oil Red O staining and for F4/80<sup>+</sup> macrophages (n = 3–5 per group) as detailed in the Supplemental Methods. LN, spleen, and bone marrow cells were analyzed by flow cytometry (n = 5 per group), and bone marrow cells were also analyzed by mass spectrometry (n = 3 per group).

**En Face Whole Aorta Sudan IV and Aortic Root Oil Red O Staining**

En face staining was carried out on whole ApoE<sup>−/−</sup> aortae, perfused with PBS in vivo, fixed in 10% neutral buffered formalin, and stained using Sudan IV as in Bruen et al. (2017). ApoE<sup>−/−</sup> hearts were perfused with PBS before freezing in OCT medium over dry ice and were stored at −80°C until cryosectioned at −25°C. When the aortic valves were clearly visible, eight 8 µm sections per mouse were collected onto Superfrost slides spanning a region of ∼800 µm. Cryosections were dried at room temperature and frozen at −80°C. For Oil Red O aortic root staining, frozen slides were thawed at 37°C for 1 minute, air-dried for 1 hour at room temperature, and fixed with 4% paraformaldehyde for 5 minutes, followed by 60% isopropanol for 1 minute, air-dried for 1 hour at room temperature, and fixed with 4% paraformaldehyde for 5 minutes, and 60% isopropanol for 5 minutes. Sections were stained with Oil Red O working solution for 10,000 rpm for 5 minutes. Peptides were acidified using 50% acetonitrile and 0.1% trifluoroacetic acid and then centrifuged at 10,000 rpm for 5 minutes and repeated twice. Samples were eluted using 50% acetonitrile (Fisher Scientific) and 0.1% trifluoroacetic acid through a StageTip. Evaporation of samples was carried out for 20-30 minutes at 60°C using a CentriVap Concentrator. Samples were resuspended in 2.5% acetonitrile and 0.5% acetic acid. Protein concentration and confirmation of peptide bonds was evaluated using the NanoDrop2000 at 218 nm. Samples were centrifuged at 13,000 rpm for 5 minutes and transferred to a mass spectrometry vial.

**Mass Spectrometry.** Peptide samples were analyzed on a quadrupole Orbitrap (X-Exact; Thermo Scientific, Dublin, Ireland) mass spectrometer. The Orbitrap was equipped with a reversed-phase NANO LC UltiMate 3000 high-performance liquid chromatography system (Dionex LC Packings, now Thermo Scientific). Peptide fractions were loaded onto C18 reversed-phase columns (5 cm length, 75 µm inner diameter) and eluted with a linear gradient from 5% to 40% acetonitrile with 0.5% trifluoroacetic acid in 60 minutes at a flow rate of 3 µL/min. Five microliters was used as the injection volume. The Orbitrap was operated in a data-dependent mode, automatically switching between mass spectrometry (MS) and MS2 acquisition. Survey full-scan MS spectra (m/z 350–1600) were with a resolution of 70,000, MS2 spectra with a resolution of 17,500. The 12 most intense ions were sequentially isolated and fragmented by higher-energy C-trap dissociation (O’Reilly et al., 2016).

**Protein Identification.** Raw data from the Orbitrap were processed using MaxQuant version 1.5.6.0 (Max Planck Institute of Biochemistry, Munich, Germany). To identify peptides and proteins, tandem mass spectrometry (MS/MS) spectra were matched to the UniProt mouse database. Tryptic specificity allowing two missed cleavages was used for all searches. The database searches were performed with carbamidomethyl as fixed modification and acetylation (protein N terminus) and oxidation (M) as variable modifications. Mass spectra were searched using the default setting of MaxQuant 1.5.6.0, namely, a false discovery rate of 1% on the peptide and protein level. Label-free quantitative (LFQ) ion intensities for protein profiles were generated using signals of corresponding peptides in different nano-high-performance liquid chromatography. MS/MS runs were matched by MaxQuant 1.5.6.0 applying a mass accuracy of at least 20 ppm and a maximum time window of 1 minute.

**Proteomic Data Analysis.** MaxQuant 1.5.6.0 identified proteins and generated LFQ intensities, which were analyzed using Perseus version 1.4.1.3 (Max Planck Institute of Biochemistry). Protein identifications were filtered to eliminate the reverse database and common contaminants. Data were filtered with stringent inclusion to obtain peptides in which the peptide was present in all biologic replicates in all samples. Data were log-transformed, and unpaired t test comparisons of fractions were carried out. For heat-map visualization, missing values were imputed with values from the normal distribution. The data set was normalized by z-score. Proteins were analyzed for pathways using Ingenuity Pathway Analysis (2018 Qiagen, Manchester, UK).

For trypsin digestion, dithiothreitol (Fisher Scientific, Dublin, Ireland) was used to reduce disulphide bonds of plasma protein samples, followed by 30 minutes of incubation at 60°C on a thermostirer without shaking. Samples were briefly centrifuged, and 200 mM iodoacetamide (Sigma-Aldrich) was added to alkylate the samples. dithiothreitol, iodoacetamide, and urea concentrations were diluted using 50 mM ammonium bicarbonate (Sigma-Aldrich) to ensure that 6 M of urea was diluted to 2 M urea before the addition of trypsin, and 0.5 µg/µl of trypsin (Sigma-Aldrich) was added to each sample with a working ratio of protein to trypsin 50:1. Digestion was carried out overnight at 37°C on a thermostirer set to 350 rpm. To stop digestion, acetic acid (1% final volume) (Fisher Scientific) was added. Peptides were washed twice with 1% trifluoroacetic acid (Fisher Scientific) in deionized water using a StageTip and centrifuged at 10,000 rpm for 5 minutes. Peptides were acidified using 50% acetonitrile and 0.1% trifluoroacetic acid and then centrifuged at 10,000 rpm for 5 minutes and repeated twice. Samples were eluted using 50% acetonitrile (Fisher Scientific) and 0.1% trifluoroacetic acid through a StageTip. Evaporation of samples was carried out for 20-30 minutes at 60°C using a CentriVap Concentrator. Samples were resuspended in 2.5% acetonitrile and 0.5% acetic acid. Protein concentration and confirmation of peptide bonds was evaluated using the NanoDrop2000 at 218 nm. Samples were centrifuged at 13,000 rpm for 5 minutes and transferred to a mass spectrometry vial.

**Bone Marrow-Derived Macrophages**

ApoE<sup>−/−</sup> bone marrow-derived macrophages (BMDMs) were cultured as previously described (Bruen et al., 2017). Briefly, bone marrow was flushed from femurs and tibiae for proteomics or cultured in 25% L929-conditioned medium for 7 days. Suspension cells on day 7 of culture represented monocytes, and adherent cells represented macrophages. Both monocytes and BMDMs were analyzed by flow cytometry and BMDMs by qRT-PCR.

**Proteomics**

Mass spectrometry analysis was carried out on bone marrow cells taken directly from femurs and tibiae from ApoE<sup>−/−</sup> mice. Bone marrow cells were lysed in 6 M urea (Sigma-Aldrich), and the protein concentration was determined using a Bradford protein assay (Bio-Rad, Fannin Ltd, Dublin, Ireland) with 40 µg of protein stored in 40 µl of 6 M urea at −20°C. Proteins were trypsin-digested with peptides, acidified, and washed.
Flow Cytometry Analysis

Day 7 BMDMs, LNs, and spleens, which were homogenized and flushed with PBS, were stained with antibodies described in Table 1. All flow cytometry antibodies were purchased from BD Biosciences (Oxford, UK) (Supplemental Table 1). Flow cytometry controls included single stains consisting of a single antibody mixed with BMDMs, LN, or splenic tissue. For gating strategies, fluorescence minus one control was used, where all antibodies excluding one were incubated with BMDMs, LN, or splenic tissue. Samples were run on the Beckman Coulter CyanADP flow cytometer (Beckman Coulter, Brea, CA) and analyzed using FlowLogic software (Miltenyi Biotec Ltd, Surrey, UK).

ELISA

ELISA was performed for human TNF-α, monocyte chemoattractant protein-1 (MCP-1), and IL-1β (Thermo Fisher, Dublin, Ireland) and murine TNF-α (R&D systems, Oxon, UK) according to the manufacturer’s instructions.

qRT-PCR

Cellular RNA was extracted from PBMC-derived human macrophages and ApoE−/− BMDMs using RNeasy Qiagen kits (Qiagen Ltd) per the manufacturer’s instructions. Murine aortas were homogenized using the Qiagen Tissue Lyser II and steel beads (Qiagen Ltd) for 10 minutes at 30 Hz, twice before RNA extraction, and extracted using Trizol reagent (Bio-Sciences). Briefly, samples were incubated on ice (5 minutes) with the addition of 1:1 chloroform:isoamyl alcohol (Sigma-Aldrich), vortexed, incubated on ice (8 minute), centrifuged (18,500 g, 30 minutes) with the RNA pellet washed twice in 75% cold ethanol (Sigma-Aldrich), resuspended in RNase-free water (Qiagen Ltd.), and quantified using the NanoDrop2000; 100–1000 ng RNA was reverse-transcribed to cDNA and analyzed on a Thermo Fisher Quantstudio 7 qRT-PCR machine. SYBR Green human TNF-α (NC_000096), murine TNF-α (Y00467.1), and murine inducible nitric oxide synthase (iNOS) (NC_000017) (Table 2) (Eurofins BPT Ireland Ltd, Waterford, Ireland) sequences are available in GenBank with associated accession numbers, murine Tgamman Arg1 (Mm00475988_m1), cathepsin (Cat)B (Mm01310506_m1) and CatZ (Mm00517697_m1) were analyzed using glyceraldehyde-3-phosphate dehydrogenase (GAPDH) (human-Hs02786624_g1, murine-Mm99999915_g1), and 18S rRNA (Hs9999991_1) (Bio-sciences) as reference genes. Cycle threshold (Ct) values were analyzed by the ΔΔCt method.

<p>| TABLE 1 |
| Flow cytometry antibodies used for flow cytometry analysis of bone marrow cells, spleens and lymph nodes (LNs). |</p>
<table>
<thead>
<tr>
<th>Cell</th>
<th>Antibodies</th>
</tr>
</thead>
<tbody>
<tr>
<td>Inflammatory bone marrow monocytes</td>
<td>CD45+CD115+ F4/80+ Ly6C+</td>
</tr>
<tr>
<td>Resident bone marrow monocytes</td>
<td>CD11c+</td>
</tr>
<tr>
<td>M1-like bone marrow–derived macrophages (BMDMs)</td>
<td>CD45+CD115+CD11b+F4/80+80+Ly6C+</td>
</tr>
<tr>
<td>M2-like BMDMs</td>
<td>CD45+CD115+CD11b+F4/80+80-Ly6C+</td>
</tr>
<tr>
<td>Monocytes and dendritic cells (DCs) in LNs</td>
<td>CD45+CD11b+Ly6C+80+80+CD11c+</td>
</tr>
<tr>
<td>DCs in LNs</td>
<td>CD45+CD11b+Ly6C+80+80+CD11c+</td>
</tr>
<tr>
<td>Splenic M1 macrophages</td>
<td>CD45+CD11b+F4/80+CD11c+</td>
</tr>
<tr>
<td>Splenic M2 macrophages</td>
<td>CD45+CD11b+F4/80+CD11c+</td>
</tr>
</tbody>
</table>

SYBR green primer sequences

<table>
<thead>
<tr>
<th>SYBR Green Primer</th>
<th>Gene Sequence</th>
</tr>
</thead>
<tbody>
<tr>
<td>Human TNF-α forward</td>
<td>5'-CTCGAACCCTCCATGTTGACAAA-3'</td>
</tr>
<tr>
<td>Human TNF-α reverse</td>
<td>5'-GCTGCCCCTCAGCTTGAG-3'</td>
</tr>
<tr>
<td>Murine iNOS forward</td>
<td>5'-CCCTCCTGATCTTTGGTGGTA-3'</td>
</tr>
<tr>
<td>Murine iNOS reverse</td>
<td>5'-CCACCCAGGAGCTCCTGGAAAC-3'</td>
</tr>
<tr>
<td>Murine TNF-α forward</td>
<td>5'-GGCAGGGTCTACATTGGAGTACTTGC-3'</td>
</tr>
<tr>
<td>Murine TNF-α reverse</td>
<td>5'-ACATTGGAGGTCAGTGAATTCG-3'</td>
</tr>
</tbody>
</table>

iNOS, inducible nitric oxide synthase; TNF, tumor necrosis factor.

Statistical Analysis

All data were analyzed using GraphPad Prism 5.0c (GraphPad Software Inc, San Diego, CA) and are expressed as the mean ± S.E.M. Shapiro-Wilk normality tests were carried out to determine whether the data were parametric or nonparametric. For comparison of human endarterectomy, DP PBS versus DP Lir and PBMC-derived macrophages VC + LPS versus Lir + LPS paired t tests or Wilcoxon-matched pairs signed rank tests were used to analyze parametric or nonparametric data, respectively. For comparison of LFD versus HFHCD and HFHCD versus LFD, Mann-Whitney tests were used. For multiple comparisons of animal weight, a two-way analysis of variance with Bonferroni’s post-test was used. Statistical significance comparing DP PBS versus DP VC, VC + LPS versus Lir + LPS and LFD versus HFHCD was considered when *P < 0.05, **P < 0.01, and ***P < 0.001, and P > 0.05 was NS and similarly considered when HFHCD versus HFHCD + Lir with *P < 0.05, **P < 0.01, and ***P < 0.001.

Results

Lir Alters Inflammatory Chemokine Secretion from Human Atherosclerotic Plaque. Inflammatory chemokine secretion of MCP-1 from human atherosclerotic plaques cultured postendarterectomy and treated ex vivo with 1 μM Lir was analyzed by ELISA. Plaques were sectioned into DP and RFD portions, which served as intraindividual controls. Ex vivo Lir treatment significantly decreased MCP-1 secretion from human DP atherosclerotic plaques (DP VC 79.32 ± 48.53 ng/ml vs. Lir 58.00 ± 35.31 ng/ml, n = 11, P < 0.01), whereas MCP-1 from RFD plaques was not altered compared with VC after Lir treatment (Fig. 1A).

Lir Modulates Monocyte and Macrophage Phenotype. To assess whether Lir can modulate monocyte or macrophage phenotype, we next analyzed proinflammatory cytokines in human PBMC-derived macrophages. Initially, MCP-1 secretion was analyzed similarly to the human endarterectomy samples, where a significant reduction in MCP-1 secretion from human PBMC-derived macrophages treated with Lir + LPS compared with the VC + LPS was detected (VC + LPS 3046 ± 892.9 pg/ml vs. Lir + LPS 2398 ± 761.6 pg/ml, P < 0.05) (Fig. 1B). Next, gene expression and secretion of the proinflammatory cytokine TNF-α were analyzed in healthy PBMC-derived macrophages. TNF-α mRNA expression was significantly reduced in LPS-stimulated human macrophages treated with Lir compared with VC (VC + LPS 154.4 ± 36.39-fold change vs. Lir + LPS 106.4 ± 33.32-fold change, P < 0.05) (Fig. 1C), and TNF-α secretion was significantly decreased in LPS-stimulated PBMC-derived macrophages treated with Lir compared with VC (VC + LPS 801.5 ± 201.5 pg/ml vs. Lir + LPS 223.0 ± 72 pg/ml, n = 12, P < 0.001) (Fig. 1D). In addition, Lir significantly reduced IL-1β secretion after LPS stimulation (VC + LPS 72.9 ± 17.5 pg/ml vs. Lir + LPS 35.5 ± 15.3 pg/ml, **P < 0.01 vs. VC + LPS).
Thus, the effect of Lir on inhibiting MCP-1 secretion from human endarterectomy plaques and human macrophages and TNF-α and IL-1β secretion from human macrophages suggests that Lir inhibition of proinflammatory cytokine secretion may alter the atherosclerotic plaque microenvironment to increase a proresolving or anti-inflammatory response. To investigate whether Lir could promote an anti-inflammatory response, IL-10 secretion and CD206 mRNA gene expression were analyzed in human macrophages, where a trend toward increased IL-10 secretion was detected with a significant induction of CD206 mRNA in LPS-stimulated human macrophages treated with Lir (VC 1.33 ± 0.21-fold change, Lir 1.86 ± 0.33-fold change, \( P < 0.05 \)) (Fig. 1, F and G). To address this issue comprehensively, we used an in vivo model of atherosclerosis regression.

Lir Inhibits Progression of Established Atherosclerosis in ApoE\(^{-/-}\) Mice Despite an HFHCD Challenge.

After 8 weeks of HFHCD administration to establish extensive aortic lesion burden (Supplemental Fig. 1), ApoE\(^{-/-}\) mice were continued on HFHCDs and treated with either 300 \( \mu \)g/kg of Lir or VC for a further 4 weeks to examine the effect of Lir on established disease. ApoE\(^{-/-}\) mice fed LFD for 12 weeks were used as controls. No significant differences in weight, glucose, food intake, or water intake were noted between HFHCD and HFHCD + Lir–treated mice at study completion (Supplemental Fig. 2). Therefore, all data are representative of weight-independent effects, which were anticipated given the dose of Lir used (Gaspari et al., 2013). En face analysis of aortae from the aortic arch to the iliac bifurcation showed HFHCD + Lir–treated ApoE\(^{-/-}\) mice had a marked decrease in whole aorta atherosclerotic lesion burden (HFHCD 6.8% ± 0.3% vs. HFHCD + Lir 4.0% ± 0.4%, \( P < 0.001 \)) and significant decrease in lesion area in all subsections of the aorta (Fig. 2, A–F) compared with the HFHCD-fed mice. Aortic root lesions were also quantified, and a significant lesion reduction was found in the HFHCD + Lir–treated mice compared with the HFHCD-treated mice (Fig. 2, G and H). Simultaneously, F4/80\(^{+}\) macrophage cells were quantified in the aortic root, where a reduction in macrophage content was found in HFHCD + Lir–treated ApoE\(^{-/-}\) mice compared with the HFHCD control, although this result was not significant (Supplemental Fig. 3). This finding clearly demonstrates that 4 weeks of Lir treatment significantly attenuates established disease and may induce resolution of pre-established disease.
In parallel, aortae were harvested from HFHCD- and HFHCD + Lir-treated ApoE\(^{-/-}\) mice and analyzed for inflammatory cytokine secretion. Although there was a trend toward decreased TNF-\(\alpha\) mRNA expression in HFHCD + Lir-treated aortae, Lir significantly decreased aortic TNF-\(\alpha\) secretion (HFHCD 36.50 \(\pm\) 14.9 pg/ml vs. HFHCD + Lir 2.21 \(\pm\) 2.2 pg/ml, \(P < 0.001\)) (Fig. 3, A and B). In addition, HFHCD administration induced a significant increase in mRNA expression of the M1 marker iNOS in ApoE\(^{-/-}\) aortae, which was blunted in HFHCD + Lir–treated animals (Fig. 3C). Finally, there was a trend toward increased aortic expression of the M2 marker Arg-1, although this trend was not significant (Fig. 3D). Together, these data suggest that Lir may alter the plaque microenvironment to attenuate atherosclerosis.

Proteomic Analysis of Bone Marrow Cells from Lir-Treated ApoE\(^{-/-}\) Mice Highlights Enrichment of Proinflammatory Pathways Essential for Regression. To gain insight into the mechanisms through which Lir mediates the atheroprotective effect in attenuating the progression of established atherosclerosis, we performed comprehensive proteomic analysis of bone marrow cells from ApoE\(^{-/-}\) mice, fed LFD for 12 weeks or HFHCD for 12 weeks with either Lir or PBS daily subcutaneous injections for the final 4 weeks of feeding. We found that 561 proteins were significantly altered with HFHCD + Lir (286 upregulated and 275 downregulated) compared with the HFHCD control. Further refinement of the data and elimination of the proteins that were significantly altered between the HFHCD and LFD groups were performed using Perseus version 1.4.1.3 (Supplemental Fig. 4A), resulting in the identification of 338 and 187 significantly altered proteins in the HFHCD and HFHCD + Lir groups, respectively, compared with the LFD control. Importantly, approximately 20% of the proteins detected were unique to the HFHCD + Lir–treated group (Fig. 4A). Statistical analysis using an unpaired \(t\) test identified a unique signature of proteins associated with Lir-mediated inhibition of atherosclerosis (Fig. 4B). The top 25 significantly altered proteins are listed in Supplemental Table 2. Comprehensive bioinformatic analysis using Ingenuity Pathway Analysis was performed on proteins significantly and uniquely regulated by Lir to elucidate potential mechanisms through which Lir mediates its effect.
Cathepsins are proteolytic macrophage enzymes documented to play a role in protein degradation and atherosclerotic lesion development. For example, CatB degrades the plaque extracellular matrix, and CatZ facilitates proinflammatory cytokine release (Zhao and Herrington, 2016). Surprisingly, proteomic analysis identified CatB and CatZ as significantly enriched proteins in bone marrow cells from HFHCD + Lir-treated mice, despite that these mice having a significantly reduced lesion burden. CatB was identified in a network linked to the insulin receptor and interferon-γ, both of which are also implicated in atherosclerosis pathogenesis (Fig. 4C). Other proteins significantly upregulated by HFHCD + Lir in this pathway are presented in Supplemental Fig. 4, B–E. Recently, proinflammatory monocytes were shown to be required for regression of atherosclerosis (Rahman et al., 2017), suggesting that enrichment of proinflammatory proteins in monocyte precursor cells may be associated with the inhibition of atherosclerosis observed with Lir-treated animals. Validation of the LFQ proteomic data confirmed a significant increase in bone marrow cell expression of the proatherogenic mediators CatB and CatZ in mice receiving daily subcutaneous Lir or VC and were analyzed by flow cytometry for proinflammatory and proresolving monocytes from HFHCD + Lir–treated ApoE−/− mice. Interestingly, CatB mRNA expression was significantly reduced, and CatZ mRNA expression was blunted in HFHCD + Lir–treated mice aortae compared with HFHCD-fed animals (Fig. 5, E and F). Thus, it is feasible to hypothesize that Lir mediates its effect by recruiting inflammatory bone marrow cells to differentiate into proresolving macrophages; in accord with recent publications, this effect mediates regression of atherosclerosis (Rahman et al., 2017). Therefore, the next series of experiments were designed to investigate monocyte and macrophage populations in bone marrow and lymphoid tissues during Lir-induced attenuation of established atherosclerosis.

**BMMCs from ApoE−/− Mice are Atheroprotective with In Vivo Lir Treatment.** ApoE−/− bone marrow monocytes and BMMCs were harvested from mice fed a LFD for 12 weeks or a HFHCD for 12 weeks; during the final 4 weeks, mice received daily subcutaneous Lir or VC and were analyzed by flow cytometry for proinflammatory and proresolving phenotypes. Although inflammatory monocytes in HFHCD-fed mice decreased significantly compared with LFD controls, monocytes from HFHCD + Lir–treated mice were not significantly different from those in the HFHCD control, and both HFHCD− and HFHCD + Lir–treated mice had significantly more resident monocytes compared with LFD controls (Fig. 6A). The proteomic data also suggest that the inflammatory status of monocytes is increased with Lir, which is in keeping with recent studies, which show inflammatory monocytes convert to proresolving macrophages in atheroprotection (Rahman et al., 2017). Thus, the analysis was extended to investigate the effect of Lir on macrophage populations. As
expected, the HFHCD-fed mice had significantly increased proinflammatory and decreased anti-inflammatory macrophage populations compared with the LFD control (LFD M1-like 7.3% ± 1.1% vs. HFHCD M1-like 18.1% ± 3.3%; LFD M2-like 92.7% ± 1.1% vs. HFHCD M2-like 71.3% ± 13.0%, P < 0.01). Importantly, Lir significantly increased M2-like and decreased M1-like macrophages compared with HFHCD-treated mice (HFHCD M1-like 18.1% ± 3.3% vs. HFHCD + Lir M1-like 5.8% ± 1.7%; HFHCD M2-like 71.3% ± 13.0% vs. HFHCD + Lir M2-like 94.2% ± 1.7%, P < 0.01). Indeed, Lir normalized the M2-like macrophage populations to those comparable with LFD-treated mice (LFD M2-like 92.7% ± 1.1% vs. HFHCD + Lir M2-like 94.24% ± 1.7%) (Fig. 6B). Finally, similar numbers of monocyte/macrophage ratio in the bone marrow from all animal groups (Fig. 6C), suggest that Lir does not influence reduction in monocyte to macrophage number, but it induces significant alterations in M2 macrophage phenotypes.

Lir Alters the Phenotype of Lymphoid Cells. Anti-inflammatory macrophages are depleted during atherosclerotic disease progression, with proinflammatory plaque macrophages dominating in the developing lesion (de Gaetano et al., 2016). Further flow cytometry analysis showed that splenic M2 macrophages were depleted after HFHCD administration compared with LFD controls as determined by a significant inhibition of M2 phenotype (LFD M2 4.3 × 10⁷ ± 2.3 × 10⁷ cells/mg vs. HFHCD M2 4.7 × 10⁷ ± 2.5 × 10⁷ cells/mg, P < 0.05). Interestingly there was no significant reduction in M2 splenic macrophages in HFHCD + Lir–treated mice compared with LFD controls, suggesting that Lir blocks the depletion of M2 macrophages observed in the HFHCD mice (Fig. 6D). Monocytes and dendritic cells (DCs) are normally recruited into advanced atherosclerotic plaques, and egress of macrophages and DCs to regional and systemic LNs occurs in the regression of atherosclerosis (de Gaetano et al., 2015). Flow cytometry analysis of LNs showed a reduction of immature monocytes and DCs and mature DCs in lymphoid tissue from HFHCD compared with LFD controls. Importantly, Lir rescued mature DCs within LNs as seen by an increase in population numbers (Fig. 6E), suggesting emigration of DCs from the resolving plaque. As there is a trend toward increased DC numbers in the draining LN with Lir, these data suggest that Lir may promote DC egress from atherosclerotic lesions, which is necessary for the induction of plaque regression.
Discussion

Chronic low-grade inflammation is associated with both diabetes and atherosclerosis; patients with type 2 diabetes mellitus and atherosclerosis have increased proinflammatory monocyte and macrophage populations (Nikiforov et al., 2017). Indeed, chronic inflammatory diseases are associated with a shift in the M1:M2 ratio toward increased proinflammatory M1 (de Gaetano et al., 2016; Ye et al., 2016). This shift in macrophage ratio has also been linked to the development of insulin resistance in adipose tissue (Ye et al., 2016) and in the progression of atherosclerosis from an asymptomatic to symptomatic disease state (de Gaetano et al., 2015).

It was previously thought that inflammatory and resident monocytes give rise to M1 and M2 macrophages, respectively (Nikiforov et al., 2017; Patel et al., 2017). Fadini et al. (2013) have reported that in type 2 diabetes, patients have markedly reduced anti-inflammatory monocytes through dysregulation in bone marrow function, which may have a negative impact on microangiopathy (Fadini et al., 2013); however, recent evidence has identified that inflammatory monocytes are required to induce proresolving macrophages to mediate regression of atherosclerosis (Rahman et al., 2017). The monocyte-derived macrophage cells within atherosclerotic plaques are thought to originate from bone marrow cells as opposed to recruitment of new circulating monocytes (Patel et al., 2017). It has been previously shown that Lir mediates an anti-inflammatory effect in humans as observed by decreased TNF-α, IL-1β, IL-6, and C-reactive protein inhibition (Shiraishi et al., 2012; Hogan et al., 2014). Hence, this may also be of clinical importance in the context of atherosclerosis and diabetes-accelerated atherosclerosis. Recently, the Liraglutide Effect and Action in Diabetes: Evaluation of CV Outcomes Results (LEADER) trial reported that after 3.8 years, the primary outcome of first occurrence of CVD death, nonfatal myocardial infarction, or nonfatal stroke was significantly reduced in Lir-treated patients (Marso et al., 2016).

Our previously published work provided mechanistic insight through which Lir may mediate atheroprotection, where Lir dictated macrophage cell fate toward an M2 pro-resolving macrophage and halted the development of early atherosclerosis in ApoE−/− mice (Bruen et al., 2017); however, because a significant portion of type 2 diabetes mellitus patients already have established atherosclerotic disease, a therapeutic goal would be to reduce and stabilize disease by blunting progression or even inducing regression of atherosclerosis while maintaining glycemic control. Our hypothesis was that, in established atherosclerosis, Lir would increase the inflammatory status of monocytes, which, when in the plaque milieu would promote a proresolving M2 macrophage that would dominate, reducing proinflammatory responses associated

![Fig. 5. Increased Cat protein expression in bone marrow cells is reversed after differentiation to macrophages and in ApoE−/− aortae. LFQ protein expression of (A) CatB and (B) CatZ from Perseus version 1.4.1.3 analysis. The mRNA expression of (C) BMDM CatB, (D) BMDM CatZ, (E) aortic CatB, and (F) aortic CatZ from LFD, HFHCD, and HFHCD + Lir–treated ApoE−/− mice were analyzed by qRT-PCR using GAPDH and 18S rRNA as reference genes. Error bars are representative of (A and B) n = 3, (C and D) n = 5, and (E and F) n = 8. Statistical analysis was performed carrying out Mann-Whitney tests between LFD vs. HFHCD where statistical significance was considered, with *P < 0.05, **P < 0.01, ***P < 0.001 and HFHCD vs. HFHCD + Lir, where $P < 0.05; $$$P < 0.01 was considered significant.](https://jpet.aspetjournals.org/doi/10.1124/jpet.117.248217)
with disease regression and stabilization. The work presented here shows that the inflammatory microenvironment of human atherosclerotic plaques is altered in response to ex vivo Lir treatment, where Lir reduces secretion of MCP-1 from human endarterectomy samples. MCP-1 regulates monocyte chemotaxis and infiltration by binding to the CC chemokine receptor 2 (CCR2) (Ding et al., 2015). Both murine and human studies have shown an association with increased CCR2/MCP-1 expression and atherosclerosis progression (Ding et al., 2015). Indeed, increased circulating MCP-1 levels are related to increased risk of CVD and CVD mortality in patients with coronary artery disease (Ding et al., 2015). As such, strategies for inhibiting MCP-1/CCR2 are currently being explored as a therapeutic goal for the treatment of vascular disorders.

In addition, further analysis on cultured LPS-stimulated inflammatory human PBMC-macrophages showed that Lir significantly decreased secretion of the M1 proinflammatory mediators, MCP-1, TNF-α, and IL-1β. We also showed Lir could promote anti-inflammatory responses in human macrophages by increasing IL-10 secretion and CD206 gene expression. As Lir altered the inflammatory environment of human atherosclerotic plaques and altered the inflammatory response of macrophage phenotypes in vitro, we hypothesized that Lir may impact the progression of pre-established atherosclerosis. Previous studies investigating GLP-1RAs and their effect on atherosclerosis in mice have been studied in early disease development and progression (Bruen et al., 2017). Here, ApoE⁻/⁻ mice were fed HFHCD for 8 weeks to establish atherosclerotic disease and maintained on the diet for a further 4 weeks with daily subcutaneous Lir treatment. This is a strength of our study and is more reflective of clinical presentation of established atherosclerosis. Mice were injected with incremental doses of Lir for 10 days to minimize adverse effects and to ensure a dosing regimen similar to that used clinically in human patients. On study completion, aortae were quantified for lesion burden. Lir significantly inhibited lesion burden throughout the aortic root and whole aorta, although this possibility warrants further investigation. Therefore, this is the first report to show that Lir blocks progression of established disease and thus presents us with a unique opportunity to interrogate the potential cellular mechanisms through which Lir mediates its effect on established disease.

Previous studies have shown that lixisenatide alters the plaque milieu by decreasing macrophage content and increasing M2 macrophages in ApoE⁻/⁻ mice (Vinué et al., 2017), although in low-density lipoprotein receptor knockout mice, Lir did not alter macrophage content or aortic mRNA expression of MCP-1 or iNOS (Bisgaard et al., 2016); however, both studies were performed in the context of early disease progression. Here, we analyzed the plaque microenvironment in established disease and have shown that there is increased TNF-α secretion and aortic iNOS mRNA expression after a 12-week HFHCD challenge. Importantly, Lir administration inhibits aortic TNF-α secretion during attenuation of disease progression.

In atherosclerotic disease progression, monocytes are recruited to the developing lesion and differentiate into macrophages, which sustain the inflammatory response and drive disease progression (Tabas and Lichtman, 2017). As bone marrow cells are recruited into lesions and are precursors for macrophage cells, we characterized these cells in the...
context of attenuated progression of established disease, using proteomic analysis of bone marrow cells from ApoE−/− mice. Bioinformatic analysis showed an upregulation of proinflammatory signaling pathways. This finding is in keeping with recent data showing that inflammatory monocyte precursor cells differentiate into anti-inflammatory macrophages during attenuation of atherosclerosis (Rahman et al., 2017).

Cathepsin proteases are involved in destabilizing atherosclerosis and driving disease progression (Zhao and Herrington, 2016). In bone marrow cells from HFHCD + Lir–treated mice, there was an enrichment of cathepsin proteases despite reduced lesion burden in these mice. Upregulation of this protein family suggests Lir increases the inflammatory status of bone marrow cells, which are precursor cells to monocytes. To address the hypothesis that an increased inflammatory status in bone marrow cells may convert into M2 macrophages and the aorta.

Indeed, M2 macrophages have been identified in numerous murine atherosclerosis regression models (Feig et al., 2012). To address whether the loss of the proinflammatory signal in bone marrow cells impacts on macrophase phenotype, we investigated immune cell phenotypes in bone marrow, spleen, and lymph nodes. The data clearly show, as expected, that Lir administration increased proinflammatory macrophage proteins in the bone marrow of ApoE−/− mice, suggesting a proinflammatory signature and that this resulted in increased M2-like macrophage populations upon differentiation. Furthermore, further flow cytometry analysis showed that the proresolving M2 macrophage phenotype prevails both in the bone marrow and in splenic tissue, suggesting that Lir blocks the depletion of M2 macrophages. Finally, we showed a trend toward increased numbers of migratory cells in aortic draining LNs, suggesting increased migratory ability out of lesions, and implies that this may contribute to the ability of Lir to attenuate progression and induce regression of atherosclerosis, although further investigation is warranted.

The data presented here have elucidated an important mechanism in Lir-mediated resolution of inflammation in established atherosclerosis, where proinflammatory bone marrow cells differentiate into proresolving macrophages and likely induce plaque egress of immune cells, promoting repair and inhibiting lesion burden.

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Authorship Contributions
Participated in research design: Bruen, Belton, McGillicuddy.
Conducted experiments: Bruen, Curley, Kajani, Lynch, O’Reilly, Dillon.
Contributed new reagents or analytic tools: Barry, Sheehan.
Performed data analysis: Bruen, Dillon, Brennan.
Wrote or contributed to the writing of the manuscript: Bruen, Belton.

References


Address correspondence to: Orina Belton, Diabetes Complications Research Centre, School of Biomolecular and Biomedical Science, UCD Conway Institute, University College Dublin, Belfield, Dublin 4, Ireland. E-mail: orina.belton@ucd.ie