Amelioration of Hyperglycemia with a Sodium-Glucose Cotransporter 2 Inhibitor Prevents Macrophage-Driven Atherosclerosis through Macrophage Foam Cell Formation Suppression in Type 1 and Type 2 Diabetic Mice

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Abstract

Direct associations between hyperglycemia and atherosclerosis remain unclear. We investigated the association between the amelioration of glycemia by sodium-glucose cotransporter 2 inhibitors (SGLT2is) and macrophage-driven atherosclerosis in diabetic mice. We administered dapagliflozin or ipragliflozin (1.0 mg/kg/day) for 4-weeks to apolipoprotein E-null (Apoe−/−) mice, streptozotocin-induced diabetic Apoe−/− mice, and diabetic db/db mice. We then determined aortic atherosclerosis, oxidized low-density lipoprotein (LDL)-induced foam cell formation, and related gene expression in exudate peritoneal macrophages. Dapagliflozin substantially decreased glycated hemoglobin (HbA1c) and glucose tolerance without affecting body weight, blood pressure, plasma insulin, and lipids in diabetic Apoe−/− mice. Aortic atherosclerotic lesions, atheromatous plaque size, and macrophage infiltration in the aortic root increased in diabetic Apoe−/− mice; dapagliflozin attenuated these changes by 33%, 27%, and 20%, respectively. Atherosclerotic lesions or foam cell formation highly correlated with HbA1c. Dapagliflozin did not affect atherosclerosis or plasma parameters in non-diabetic Apoe−/− mice. In db/db mice, foam cell formation increased by 4-fold compared with C57/BL6 mice, whereas ipragliflozin decreased it by 31%. Foam cell formation exhibited a strong correlation with HbA1c. Dapagliflozin did not affect atherosclerosis or plasma parameters in non-diabetic Apoe−/− mice. Aortic atherosclerotic lesions, atheromatous plaque size, and macrophage infiltration in the aortic root increased in diabetic Apoe−/− mice; dapagliflozin attenuated these changes by 33%, 27%, and 20%, respectively. Atherosclerotic lesions or foam cell formation highly correlated with HbA1c. Dapagliflozin did not affect atherosclerosis or plasma parameters in non-diabetic Apoe−/− mice. In db/db mice, foam cell formation increased by 4-fold compared with C57/BL6 mice, whereas ipragliflozin decreased it by 31%. Foam cell formation exhibited a strong correlation with HbA1c. Gene expression of lectin-like ox-LDL receptor-1 and acyl-coenzyme A:cholesterol acyltransferase 1 was upregulated, whereas that of ATP-binding cassette transporter A1 was downregulated in the peritoneal macrophages of both types of diabetic mice. SGLT2i normalized these gene expressions. Our study is the first to demonstrate that SGLT2i exerts anti-atherogenic effects by pure glucose lowering independent of insulin action in diabetic mice through...
suppressing macrophage foam cell formation, suggesting that foam cell formation is highly sensitive to glycemia ex vivo.

**Introduction**

Diabetes accelerates the clinical course of atherosclerosis, a condition associated with arterial endothelial dysfunction and several metabolic abnormalities, can create a pro-inflammatory environment, and can induce foam cell formation of macrophages [1–3]. Because atherosclerosis is accelerated in both type 1 and type 2 diabetes [4–6], it is reasonable to assume that hyperglycemia plays a major role in the pathogenesis of diabetes-induced atherosclerosis. While, patients with metabolic syndrome or pre-diabetes without significant hyperglycemia often develop atherosclerotic cardiovascular events [7–9]. However, it remains unclear whether hyperglycemia contributes to the development of atherosclerosis in diabetes because many risk factors change with the impairment of insulin action. There have been numerous in vitro studies that have examined the toxic effects of high glucose levels on vascular and inflammatory cells related to atherosclerotic lesions [10–12]. However, studies on cultured cells have provided only hypothesis-generating results that need to be evaluated in in vivo studies.

Recently, sodium-glucose cotransporter 2 inhibitors (SGLT2is) have been developed as novel therapeutic agents for the treatment of patients with type 2 diabetes. These drugs inhibit the reabsorption of glucose in the proximal tubules of the kidney, leading to increased urinary glucose excretion and amelioration of hyperglycemia in patients with diabetes [13–16]. This simple mechanism for lowering glucose levels does not directly influence any insulin-related metabolic changes. This agent could be a useful tool to study the direct relationship between glycemia and atherosclerosis independent of insulin secretion and action in vivo. The acceleration of atherosclerosis is influenced by abnormalities in cellular cholesterol homeostasis, which is demonstrated by the subendothelial accumulation of lipid-laden macrophage foam cells. The accumulated foam cells in the subendothelial space create lipid-rich plaque, which is an initial process of atherosclerosis [1–3]. There have been several in vitro studies that have revealed that high glucose levels enhance foam cell formation in cultured macrophages [17, 18]; however, it remains unclear whether foam cell formation measured ex vivo reflects atherosclerotic lesions. Foam cell formation is regulated by several factors: 1) scavenger receptors, such as CD36 and lectin-like ox low-density lipoprotein (LDL) receptor-1 (Lox-1) [1], 2) acyl-coenzyme A:cholesterol acyltransferase 1 (ACAT1), a rate-limiting enzyme for the esterification of cholesterol [1], and 3) free-cholesterol efflux mediated by ATP-binding cassette transporter A1 (ABCA1) and ATP-binding cassette sub-family G member 1 (ABCG1) [1]. Several in vitro studies have revealed that scavenger receptors were upregulated and that either ABCA1 or ABCG1 was downregulated in high glucose conditions [19, 20]. The aim of the present study was to evaluate the effect of the amelioration of hyperglycemia by SGLT2i on the development of aortic atherosclerotic lesions, macrophage foam cell formation, and related molecules in type 1 and type 2 diabetic mice.

**Materials and Methods**

**Chemicals and reagents**

Streptozotocin (STZ) was purchased from Sigma-Aldrich (Saint-Louis, MO, USA). Dapagliflozin and ipragliflozin were kindly gifted from AstraZeneca (Gaithersburg, MD, USA) and Astellas Pharma, Inc. (Tokyo, Japan), respectively.
Animal experiments

This study was conducted in strict accordance with the recommendations in the Guide for the Care and Use of Laboratory Animals of the National Institutes of Health. The protocol was approved by the Institutional Animal Care and Use Committee of Showa University (Permit Number: 04090). All surgeries and sacrifice were performed under general anesthesia using isoflurane, and all efforts were made to minimize suffering. The health condition of all mice was carefully checked every day by us or animal experts in the animal facility of Showa University School of Medicine.

Experiment #1

A total of forty-five male apolipoprotein E-null (Apoe−/−) mice were purchased from Sankyo Labo Service (Tokyo, Japan) at the age of 9 weeks and were kept on a standard rodent chow until the age of 15 weeks. From 15 weeks of age, the animals received intraperitoneal injections of saline or streptozotocin (50 mg/kg/day) for 5 consecutive days to create a type 1 diabetes model, according to the method described by Takeda et al [21]. Adverse effects of STZ injection in mice has been reported as follows: weight loss, respiratory distress, rapid glycemic shifts resulting in life-threatening hypoglycemia, and a generalized poor body condition [22]. In our protocol, a mouse that lost weight by more than 20% from the baseline, or that showed obvious weakness undergoes euthanasia with CO2 exposure to avoid severe pain or distress. Three mice found dead in their cages within 7 days since the last injection of STZ. The other mice did not show any clinical signs including severe weight loss or obvious weakness. The fasting blood glucose (FBG) levels were measured 10 days after the last injection via a tiny cut in the tail vein using the glucose monitoring system (Nipro StatStrip; Nipro Corporation, Osaka, Japan). All survived mice showed fasting blood glucose levels higher than 200 mg/dL, and were enrolled in this experiment as diabetic mice. At the age of 17 weeks, both non-diabetic and diabetic mice were switched to an atherogenic diet containing 30% fat, 20% sucrose, 8% NaCl, and 0.15% cholesterol (Oriental Yeast, Tokyo, Japan) [23–26] and randomly assigned to vehicle or SGLT2i treatment. The SGLT2i (dapagliflozin) was given via drinking water at a dose of 1.0 mg/kg/day. After 4 weeks of this treatment, peritoneal macrophages were harvested, and the mice were sacrificed to collect the aorta under general anesthesia with isoflurane.

Experiment #2

Thirty-one male db/db mice (a mouse model of type 2 diabetes) and seven male C57/BL6 (wild type) mice were purchased from Sankyo Labo Service at the age of 6 weeks and kept on a standard rodent diet. All db/db mice developed diabetes at the age of 8 weeks with FBG levels above 200 mg/dL. Starting from the age of 9 weeks, the diabetic db/db mice were given standard rodent diet with or without diet containing the SGLT2i (ipragliflozin) at the dose of 1.0 mg/kg/day for 4 weeks. The wild-type mice were kept on standard rodent diet; they were used as the control model during the same period. At the end of the treatment period, we collected peritoneal macrophages.

Measurements

Glycated hemoglobin (HbA1c) levels were measured through a cut in the tail vein using the quick test A1C Now Plus (Bayer, Frankfurt, Germany) before sacrificing the animals. The blood samples were collected after a 6-h fast. The plasma levels of glucose, total cholesterol, high-density lipoprotein (HDL) cholesterol, and triglyceride were measured by enzymatic methods (WAKO, Osaka, Japan). We had previously reported that dipeptidyl peptidase-4
inhibitor elevated active glucagon-like peptide-1 (GLP-1) and total glucose-dependent insulinotropic polypeptide (GIP), that prevented the acceleration of atherosclerosis by suppressing foam cell formation in Apoe<sup>−/−</sup> mice [22, 23]. Therefore, we measured the endogenous incretin levels in this study. The plasma levels of insulin, active GLP-1, and total GIP were determined by enzyme-linked immunosorbent assay (ELISA) (Ultra-sensitive mouse insulin ELISA kit from Morinaga, Kanagawa, Japan; GLP-1 (active) ELISA and Rat/Mouse GIP (total) ELISA from EMD Millipore (Billerica, MA, USA).

**Oral glucose tolerance test (OGTT)**

Glucose (0.5 g/kg body weight) was orally administered through a gavage tube after 6 h of fasting, and the blood glucose levels were measured at the specified time points of 0 (pre-glucose/fasting glucose level) and at 15, 30, 60, and 120 min after using the Nipro StatStrip. The area under the curve (AUC) was calculated as the area under the glucose curve from 0 to 120 min multiplied by the minutes at the measured time points.

**Blood pressure measurement**

Blood pressure and pulse were measured on the day of sacrifice in the fasting state using the tail-cuff method (Model MK-2000ST, Muromachi-Kikai, Tokyo, Japan).

**Atherosclerotic lesion assessment**

The whole aorta was washed with perfused phosphate buffered saline (PBS) and fixed with 4% paraformaldehyde (wt/vol). After surrounding connective and adipose tissues were carefully removed, the aorta was longitudinally dissected and excised from the root to the bifurcation of the iliac artery. The entire aorta and cross-sections of the aortic root were stained with oil red O for the assessment of atherosclerotic lesions [23–27]. Macrophage infiltration into the aortic wall was visualized using anti-mouse MOMA-2 antibody staining [23–27]. The en face oil red O positive area was traced by an investigator that was blind to the treatment and measured using an image analyzer (Adobe Photoshop, San Jose, CA, USA; Image J Software, Bethesda, MD, USA). The severity of atheromatous plaques and lesion area of macrophage infiltration were assessed as the sum of stained areas in the aortic root [23–27].

**Cell culture**

Four days after the intraperitoneal injection of 4 mL of aged autoclaved thioglycolate broth, exudate peritoneal cells were isolated from treated non-diabetic or diabetic Apoe<sup>−/−</sup> mice at 21 weeks of age or from diabetic db/db mice at 13 weeks of age using a peritoneal lavage with 8 mL of ice-cold PBS [23–26]. The cells were suspended in a culture medium (RPMI-1640 supplemented with 10% FBS, streptomycin, and penicillin) and seeded onto 3.5-cm dishes (3×10<sup>6</sup> cells/dish) for either a reverse transcription polymerase chain reaction (RT-PCR) or a cholesterol esterification assay. After 1 h of incubation at 37°C in a 5% CO<sub>2</sub> incubator to allow cell adhesion to the dish, the medium was discarded to remove any non-adherent cells, and the adherent cells were identified as peritoneal macrophages [23–26].

**Cholesterol esterification assay**

The adherent macrophages were incubated for 18 h in the RPMI-1640 medium that contained 10 μg/mL human oxidized (ox)-LDL with 0.1 mmol/L [3H]oleate conjugated with bovine serum albumin. The cellular lipids were extracted and the radioactivity of the cholesterol [3H] olate was determined with thin-layer chromatography [23–26].
Gene expression

Total RNA was extracted from peritoneal macrophages using a commercially available kit (QIAGEN; Hilden, Germany). The gene expressions were assessed by real-time RT-PCR using the TaqMan gene expression assay and a sequence detection system (ABI PRISM 7900, Life Technologies, Carlsbad, CA, USA), which used cDNA synthesized from isolated RNA samples to detect the expression of target genes, such as Lox-1; Mm00454586_m1, CD36; Mm01135198_ml, ACAT1; Mm00507463_ml, ABCA1; Mm00442646_ml, and ABCG1; Mm00437390_m1. TaqMan gene expression assays for mouse glyceraldehyde-3-phosphate dehydrogenase (GAPDH) was purchased from Life Technologies. The levels of these gene expressions were normalized using GAPDH as an internal control.

Statistical analysis

All values are expressed as mean ± standard error of mean (SEM). We used one-way analysis of variance (ANOVA) followed by Tukey test for the comparison of more than two groups and the unpaired t-test for the comparison of two groups. The correlation between two valuables was determined using Pearson’s correlation test. Statistical calculations were performed using JMP software (Version 11; SAS Institute Inc., Cary, NC, USA). The significance level was defined as p < 0.05.

Results

Characteristics and laboratory data

Table 1 shows the characteristics and laboratory data from four groups of non-diabetic and diabetic Apoe−/− mice that did or did not receive SGLT2i (dapagliflozin) (Experiment #1). Although the systolic blood pressure (SBP) was slightly decreased in dapagliflozin-treated diabetic Apoe−/− mice, this change was not statistically significant. Non-diabetic Apoe−/− mice that received dapagliflozin, diabetic mice that received vehicle, and diabetic mice that received dapagliflozin consumed 1.4–1.7 times more drinking water and produced 1.7–2.6 times more urine than non-diabetic mice that received vehicle. These diabetic Apoe−/− mice exhibited the classical features of STZ-induced insulin-deficient type 1 diabetes, such as severe hyperglycemia, hyperphagia, low body weight gain, and low insulin concentration. Based on our preliminary experiments with the same protocol used in the present study, we expected a successful induction rate of diabetes in more than 90% of mice, and a mortality rate including euthanasia in less than 10%. In the present study, all the mice except for three dead ones developed diabetes, showing the successful rate of 93% and the mortality rate of 7%. There is one previous study, which used the same administration protocol as ours, reporting the mortality rate of approximately 8% [28], which are comparable with our results. The diabetic mice that received vehicle had slightly elevated levels of total cholesterol compared with their non-diabetic counterparts but this was not statistically significant. Diabetic Apoe−/− mice lost 10% (p < 0.05) of their body weight compared with non-diabetic Apoe−/− mice. In diabetic Apoe−/− mice, dapagliflozin had no effect on insulin levels after 6 h of fasting, but it significantly decreased FBG levels from 417 ± 31 mg/dL to 134 ± 18 mg/dL and HbA1c levels from 8.2% ± 0.3% to 5.8% ± 0.3%. However, there were no changes in the plasma levels of active GLP1, total GIP, total cholesterol, triglyceride, and HDL cholesterol among the groups.

Table 2 shows the metabolic parameters of diabetic db/db mice with or without SGLT2i (ipragliflozin) compared with wild-type mice (Experiment #2). As a result of severe hyperglycemia, diabetic db/db mice consumed approximately 1.3 times more drinking water, had approximately 1.4 times more food intake, and passed 2 times the urine volume compared with wild-
type mice. These diabetic db/db mice exhibited the classical features of type 2 diabetes, such as severe hyperglycemia, high insulin concentrations, and high lipid profiles (total cholesterol, HDL cholesterol, and triglyceride). In diabetic db/db mice, SBP was lower in the ipragliflozin-treated group, but this change was not statistically significant. After ipragliflozin treatment, mice consumed approximately 2.4 times more drinking water and produced approximately 1.6 times more urine than those that received vehicle. These diabetic db/db mice that received ipragliflozin lost 6% ($p < 0.05$) of their body weight compared with those that received vehicle.

Ipragliflozin significantly decreased HbA1c levels from 8.0% ± 0.2% to 6.4% ± 0.3% and FBG levels from 326 ± 32 mg/dL to 180 ± 29 mg/dL. There were no changes in the plasma levels of active GLP1, total GIP, total cholesterol, HDL cholesterol, and triglyceride with or without ipragliflozin treatment in diabetic db/db mice.

Glucose tolerance

In a subset of animals, OGTT was conducted to determine glucose tolerance (Fig 1). Diabetic Apoe−/− mice had a significantly higher glucose loading after OGTT and AUC (975 ± 46 mg/dLxh) than non-diabetic Apoe−/− mice (AUC; 356 ± 20 mg/dLxh). The treatment with dapagliflozin improved glucose loading after OGTT and improved AUC (291 ± 16 mg/dLxh) in diabetic Apoe−/− mice compared with those that received vehicle. In diabetic db/db mice, mice treated with ipragliflozin had significantly lower glucose loading after OGTT and a lower AUC (637 ± 28 mg/dLxh) compared with those that received vehicle (AUC; 1255 ± 108 mg/dLxh) as well.
Atherosclerotic lesions assessment

Atherosclerotic lesions were significantly accelerated in diabetic Apoe<sup>−/−</sup> mice. The surface areas of atherosclerotic lesions, atheromatous plaque size, and macrophage infiltration in the aortic root were increased by 78%, 150%, and 82% compared with those in non-diabetic Apoe<sup>−/−</sup> mice, suggesting the importance of hyperglycemia as an enhancer of atherosclerotic lesion formation. Diabetic Apoe<sup>−/−</sup> mice treated with dapagliflozin showed significant reductions in the surface areas of atherosclerotic lesions (33%, \( p < 0.001 \)), in atheromatous plaque size (27%, \( p < 0.001 \)), and in macrophage infiltration (20%, \( p < 0.01 \)) compared with those that received vehicle. In contrast, dapagliflozin did not show any effect on atherosclerosis in non-diabetic Apoe<sup>−/−</sup> mice (Fig 2).

Foam cell formation in exudate peritoneal macrophages

The number of exudate peritoneal cells and morphological cell characteristics did not differ significantly among the groups. The ox-LDL-induced cholesterol ester accumulation, which indicated the degree of foam cell formation, was approximately 3–4 folds higher in macrophages obtained from both types of diabetic mice than in those obtained from non-diabetic mice \( (p < 0.0001; \text{Fig 3A and 3B}) \). This increased cholesterol ester (CE) accumulation in diabetic Apoe<sup>−/−</sup> mice was significantly suppressed with dapagliflozin treatment (34%, \( p < 0.001; \text{Fig 3A} \)). Similarly, iragliflozin administration significantly reduced cholesterol ester accumulation in macrophages from diabetic db/db mice (31%, \( p < 0.001; \text{Fig 3B} \)).

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### Table 2. Characteristics and laboratory data from three groups of diabetic db/db mice with or without iragliflozin, and wild-type mice that received vehicle.

|                       | C57/BL6 mice | Diabetic db/db mice | Iragliflozin |
|-----------------------|--------------|---------------------|--------------|
|                       | Vehicle      | Diabetic            | Iragliflozin |
|                       |              | db/db               |              |
| Number                | 7            | 18                  | 13           |
| Final body weight (g) | 25.2 ± 0.3   | 49.6 ± 0.6 a        | 46.8 ± 0.7 a,b |
| Food intake (g/day)   | 3.1 ± 0.2    | 4.4 ± 0.3 a         | 4.8 ± 0.4 a  |
| Water intake (mL/day) | 3.2 ± 0.3    | 4.0 ± 0.4           | 9.4 ± 1.1 a,b |
| Urine volume (mL/day) | 3.8 ± 0.4    | 7.6 ± 0.3 a         | 12.5 ± 1.2 a,b |
| SBP (mmHg)            | 102 ± 5      | 109 ± 2             | 104 ± 3      |
| Pulse (/min)          | 546 ± 21     | 609 ± 19            | 633 ± 13     |
| Glucose (mg/dL)       | 101 ± 11     | 326 ± 32 a          | 180 ± 29 a,b |
| HbA1c (%)             | 4.2 ± 0.1    | 8.0 ± 0.2 a         | 6.4 ± 0.3 a,b |
| Insulin (ng/mL)       | 0.63 ± 0.17  | 6.36 ± 0.62 a       | 5.75 ± 0.9 a |
| Total-C (mg/dL)       | 78 ± 8       | 134 ± 8 a           | 171 ± 19 a   |
| HDL-C (mg/dL)         | 30 ± 4       | 63 ± 8 a            | 55 ± 6 a     |
| Triglyceride (mg/dL)  | 20 ± 3       | 67 ± 7 a            | 60 ± 8 a     |
| Active GLP-1 (pmol/L) | 1.67 ± 0.2   | 3.13 ± 0.05 a       | 3.7 ± 0.9 a  |
| Total GIP (pmol/L)    | 24.5 ± 3.9   | 51.2 ± 5.5 a        | 45.4 ± 9.9 a |

Non-diabetic wild-type mice that received vehicle, diabetic db/db mice that received vehicle, diabetic db/db mice that received iragliflozin were measured. The values show mean ± SEM. Iragliflozin, sodium glucose co-transporter 2 inhibitor.

One-way ANOVA followed by Tukey test:
- \( a \), \( p < 0.05 \) vs. wild-type mice that received vehicle;
- \( b \), \( p < 0.05 \) vs. diabetic db/db mice that received iragliflozin.

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Correlation between atherosclerosis and glycemic control

To examine the relationship between the development of atherosclerosis and glycemic control, we tested the correlation between atherosclerotic lesions and HbA1c levels. In diabetic Apoe−/− mice, atherosclerotic lesions highly correlated with HbA1c levels (r = 0.77, p < 0.0001, n = 22). However, no correlation was observed in the non-diabetic group (Fig 4A). In the analysis of the correlation between foam cell formation and atherosclerosis, diabetic Apoe−/− mice showed a strong correlation (r = 0.95, p < 0.0001, n = 15), while non-diabetic mice did not (Fig 4B). We also evaluated the correlation between the degrees of foam cell formation and glycemic control. The test demonstrated a significant correlation between foam cell formation and HbA1c levels only in diabetic Apoe−/− mice (r = 0.91, p < 0.0001, n = 15) and in diabetic db/db mice (r = 0.88, p < 0.0001, n = 31) but not in non-diabetic mice (Fig 5A and 5B). In addition, we examined the correlation of atherosclerosis or foam cell formation with various metabolic parameters other than glucose in diabetic Apoe−/− mice and diabetic db/db mice (Table 3). Body weight, systolic blood pressure, insulin, total cholesterol, or triglyceride did not correlate with the atherosclerotic lesion area or the degree of foam cell formation in either type of diabetic animals.

Gene expressions in exudate peritoneal macrophages

To explore molecular changes related to macrophage foam cell formation in a diabetic state, we analyzed the expression of selected genes in peritoneal macrophages. The macrophages from both diabetic Apoe−/− mice and diabetic db/db mice showed that the expressions of Lox-1, CD36, and ACAT1 were elevated, in contrast the expressions of ABCA1 and ABCG1 were
reduced compared with those in non-diabetic mice (p < 0.05; Figs 6 and 7). Dapagliflozin treatment reversed these changes except for CD36 and ABCG1 (p < 0.05; Fig 6). The gene expressions of Lox-1, CD36, and ACAT1 were upregulated, while those of ABCA1 and ABCG1 were downregulated in db/db diabetic mice. Ipragliflozin treatment normalized these abnormal gene expressions that had facilitated foam cell formation (p < 0.05; Fig 7).

Discussion

This study was the first to demonstrate that SGLT2i prevented the development of atherosclerotic lesions in diabetic Apoe−/− mice. Moreover, the present study revealed a strong relationship between HbA1c levels and atherosclerotic lesions or macrophage foam cell formation. A strikingly close relationship between atherosclerotic lesions and foam cell formation may suggest a causative role for foam cell formation in atherogenesis. Because diabetic db/db mice do not accelerate visible atherosclerotic lesions, macrophage foam cell formation was used as a surrogate marker for atherosclerosis in this study. We also found that macrophage foam cell formation was substantially increased in diabetic db/db mice and that SGLT2i attenuated this in an HbA1c-dependent manner. To the best of our knowledge, the present study was the first to indicate a strong relationship between glycemia and macrophage foam cell formation ex vivo. The assay for foam cell formation usually requires 2–3 mice to obtain enough peritoneal macrophages for completion. We successfully measured foam cell formation in material obtained from one mouse, which enabled us to determine the individual associations of foam cell formation with HbA1c levels or atherosclerotic lesion area.

We used two types of diabetic mice whose insulin secretion and insulin resistance levels were completely in opposition. However, SGLT2i uniformly suppressed macrophage foam cell formation. The data are expressed as mean ± SEM. One way ANOVA followed by Tukey test: †p < 0.01, ‡p < 0.001, §p < 0.0001.

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Fig 2. Suppressive effects of dapagliflozin administration against the development of aortic atherosclerotic lesions in non-diabetic and diabetic Apoe−/− mice. Twenty-two mice at 15 weeks of age were made diabetic with peritoneal injections of STZ (50 mg/kg/day) for 5 consecutive days and twenty mice were treated with saline. The 17-week-old non-diabetic and diabetic Apoe−/− mice were orally given SGLT2i (dapagliflozin) or vehicle for 4 weeks, starting from 17 weeks of age. Representative atherosclerotic lesions in the aortic surface stained with oil red O (a–d) and measured (m). Yellow arrows show notable atherosclerotic lesions. In the aortic root, the atheromatous plaques and monocyte/macrophage accumulations were stained with Oil red O (e–h) or anti-MOMA2 antibody (i–l). Black arrows show notable atheromatous plaques. The severity of atheromatous plaques (n) and degree of monocyte/macrophage accumulation (o) were evaluated. The data are expressed as mean ± SEM. One way ANOVA followed by Tukey test: †p < 0.01, ‡p < 0.001, §p < 0.0001.
formation in both mice. Considering the findings in non-obese STZ-induced diabetic mice and the nature of this agent, it is reasonable to assume that lowering glucose levels can attenuate atherosclerosis independent of any insulin actions. SGLT2i significantly reduced body weight only in obese \textit{db/db} mice but not in STZ-induced diabetic lean mice. Because body weight changes were not correlated with foam cell formation in diabetic \textit{db/db} mice, weight loss itself may not be a major factor for the anti-atherosclerotic effects of SGLT2i. However, there is a possibility that improvement of insulin sensitivity could have indirectly contributed to suppression of atherosclerosis or macrophage foam cell formation. Insulin resistance plays an important role in development and progression of atherosclerosis. Previous studies demonstrate that a SGLT2i improved insulin sensitivity in patients with type 2 diabetes [29]. We conducted oral glucose loading for assessment of glucose tolerability. However, the results did not indicate insulin sensitivity, and the lack of evaluation of insulin sensitivity is one of the limitations in this study. Further study using euglycemic-hyperinsulinemic clamp test with tracer is necessary to clarify change of insulin sensitivity by a SGLT2i and its role in atherosclerosis. Severe hypercholesterolemia may cause atherosclerosis in mice, such as in \textit{Apoe}^{-/-} mice. As in
a previous study [24], plasma cholesterol levels were somewhat higher in diabetic Apoe−/− mice than in non-diabetic Apoe−/− mice and SGLT2i tended to lower these levels. However, there was no significant correlation between cholesterol levels and atherosclerotic lesion area or foam cell formation in either non-diabetic or diabetic Apoe−/− mice that did or did not receive SGLT2i. Therefore, it is unlikely that a mild reduction in cholesterol by SGLT2i can account for a substantial anti-atherosclerotic effect of SGLT2i. Similarly, plasma triglyceride, HDL cholesterol, and blood pressure were unaltered by SGLT2i in Apoe−/− mice and diabetic db/db mice; this indicated that changes in other risk factors for atherosclerosis outside of glycemia could not account for the favorable effect of SGLT2i on atherosclerosis. Thus, the pure glucose lowering effect of SGLT2i may lead to the suppression of atherogenesis in both type 1 and type 2 diabetic animals.
Macrophage CD36 scavenges ox-LDL, which leads to foam cell formation, and appears to be a key proatherogenic molecule [1, 30, 31]. The gene expression of CD36 in macrophages was substantially upregulated in both type 1 and type 2 diabetic mice. There have been supportive data from studies that demonstrated that CD36 expression or protein was highly expressed in diabetic mice [31, 32] and that high glucose induced CD36 expression by stimulating the translational process in cultured macrophages [31–33]. The upregulated gene expression of CD36 was attenuated by SGLT2i treatment in macrophages that were derived from diabetic db/db mice and corresponded to its glucose-lowering effect. However, an insignificant suppressive effect of SGLT2i on CD36 expression was observed in diabetic Apoe−/− mice, although there was a glucose-lowering effect that was similar to the one attained in diabetic db/db mice. On the other hand, another scavenger receptor, Lox-1 expression, was completely attenuated by SGLT2i in both diabetic mice models. Li et al reported that the gene expression

Table 3. Correlation of atherosclerosis or foam cell formation to metabolic parameters

|                | Diabetic Apoe−/− mice |          |          | Diabetic db/db mice |          |          |
|----------------|-----------------------|----------|----------|---------------------|----------|----------|
|                | (n = 22)              | (n = 15) | (n = 31) |                     |          |          |
| Atherosclerotic lesions | r | p value | r | p value | r | p value |
| Body weight    | −0.07                 | 0.74     | −0.09    | 0.74     | 0.28  | 0.14     |
| SBP            | 0.26                  | 0.21     | 0.34     | 0.19     | 0.06  | 0.79     |
| Insulin        | −0.3                  | 0.3      | −0.33    | 0.34     | −0.15 | 0.44     |
| Total-C        | 0.31                  | 0.12     | 0.16     | 0.55     | −0.27 | 0.18     |
| Triglyceride   | 0.32                  | 0.13     | 0.24     | 0.38     | 2.0   | 0.29     |

Total diabetic Apoe−/− mice that received vehicle or dapagliflozin, total diabetic db/db mice that received vehicle or ipragliflozin were measured. SBP, systolic blood pressure; BW, body weight; Total-C, total cholesterol. r values indicate Pearson correlation coefficient, p< 0.05.

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Fig 6. Changes in gene expression related to foam cell formation in the macrophages obtained from Apoe−/− mice. Gene expressions were measured in the peritoneal macrophages obtained from non-diabetic Apoe−/− mice (n = 7), diabetic Apoe−/− mice that received vehicle (n = 6), and those that received dapagliflozin (n = 5). Gene expressions of (A) lectin-like ox-LDL receptor-1 (Lox-1), (B) CD36, (C) acyl-coenzyme A:cholesterol acyltransferase 1 (ACAT1), (D) ATP-binding cassette transporter A1 (ABCA1), and (E) ATP-binding cassette sub-family G member1 (ABCG1) and the association with GAPDH were analyzed by real-time RT-PCR before the addition of ox-LDL. The values show mean ± SEM. One-way ANOVA followed by Tukey test for the comparison of Apoe−/− mice: *p < 0.05, †p < 0.01.

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of Lox-1 in macrophages was elevated by high glucose levels, which was directly linked to foam cell formation [17]. Moheimani et al reported that high glucose levels increased Lox-1 expression but the researchers failed to observe any upregulation of CD36 expression [34]. Lox-1 may be more glucose sensitive than CD36. Nevertheless, the majority of previous studies have supported that high glucose levels induced scavenger receptors in macrophages, which facilitated foam cell formation [17, 18, 31].

Macrophage foam cell formation characterized by cholesterol ester accumulation is modulated by ACAT1 [1], a rate-limiting enzyme for the esterification of free cholesterol from ox-LDL. Our data showed that ACAT1 gene expression was strongly enhanced in macrophages from type 1 and type 2 diabetic mice, and the amelioration of hyperglycemia by SGLT2i attenuated ACAT1 expression. However, because ACAT1 expression is regulated by the substrate (free cholesterol) availability inside cells [35], a change in ACAT1 might simply be a consequence of changes in scavenger receptors. Further studies will be required to clarify the direct relationship between glucose and ACAT1 independent of scavenger receptors.

The cholesterol efflux system plays a pivotal role in the suppression of foam cell formation in macrophages. ABCA1 and ABCG1 are the main molecules for cholesterol efflux from macrophage to apolipoprotein A-1 or HDL particles [1, 36–39]. There have been conflicting results in terms of ABCA1 and ABCG1 expression in macrophages derived from non-diabetic/diabetic mice or in high glucose treatment [19, 20, 40]. Mauldin et al reported that diabetic db/db macrophages had impaired cholesterol efflux to HDL but not to lipid-free apo A-1; this suggested that the increased levels of esterified cholesterol in the diabetic db/db macrophages were due to a selective loss of ABCG1-mediated efflux to HDL [41]. On the other hand, several studies have reported that ABCA1 expression was also downregulated by high glucose [19, 42, 43]. In the present study, a change in glycemia seems to more strongly influence ABCA1 expression than ABCG1 expression. Because we did not compare the actual cholesterol efflux to apolipoprotein A1 or HDL, it remains unknown whether upregulated ABCA1 by SGLT2i contributed to the correction of glucose-induced macrophage foam cell formation.

Polyol pathway flux, advanced glycation end-product (AGE) formation, protein kinase C (PKC) activity, and hexosamine pathway flux have all been shown to be involved in
hyperglycemia-induced vascular damage [44]. In addition, chronic exposure to oxidative stress induces systemic inflammation [45, 46]. Given the fact that a number of studies support the crucial role of inflammation in the pathogenesis of atherosclerosis and the strong association between hyperglycemia and systemic inflammation, amelioration of hyperglycemia by SGLT2i might have also contributed to suppression of systemic inflammation via reduced ROS, subsequently resulting in protection against atherosclerosis. As a limitation of this study, changes in inflammation or ROS were not investigated. Our study should be extended to identify other possible mechanisms for glucose-stimulated macrophage foam cell formation and its related atherogenesis.

Conclusions
Our study is the first to demonstrate that SGLT2i exerts anti-atherogenic effects by the normalization of blood glucose levels in both mouse models of type 1 and type 2 diabetes through suppressing macrophage foam cell formation; this is independent of insulin action. Our results suggest that foam cell formation regulated by Lox-1, ACAT1, and ABCA1 may be highly sensitive to glycemia ex vivo.

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Author Contributions
Conceived and designed the experiments: TH. Performed the experiments: MT MH KK HK. Analyzed the data: MT MH KK HK. Wrote the paper: MT YM TH. Contributed to discussion and revision of the manuscript: MN TW.

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