Endothelial lipase and reverse cholesterol transport in type 2 diabetes mellitus
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ABSTRACT
Aims/Introduction: Endothelial lipase (EL) plays an important role in high-density lipoprotein (HDL) metabolism and experimental data suggest that EL might be proatherogenic. We have investigated whether serum EL concentration is associated with changes in serum capacity to induce cholesterol efflux and arterial stiffness in type 2 diabetes.

Materials and Methods: Serum EL was assayed by ELISA in 172 diabetic patients and 175 controls. The ability of serum to induce cholesterol efflux was measured using a cell culture system and arterial stiffness was determined by measuring pulse wave velocity (PWV) between carotid and femoral arteries.

Results: Diabetic patients had significantly higher C-reactive protein (CRP) and EL (27.7 ± 16.6 ng/mL vs 24.0 ± 11.3, P < 0.05). Cholesterol efflux to serum mediated through scavenger receptor class B type I was impaired (15.1 ± 2.5% vs 16.7 ± 3.1, respectively, P < 0.01). In controls, serum EL correlated with cholesterol efflux to serum (r = −0.16, P = 0.025), but only a trend was seen in the diabetic patients. Linear regression showed that in controls, HDL, serum EL and waist circumference were major independent determinants of cholesterol efflux; whereas in the diabetic cohort, the major independent determinants of cholesterol efflux were HDL, CRP and age. PWV was increased in the diabetic patients (P < 0.01), but no association between serum EL and PWV was seen in either groups.

Conclusions: Serum EL was increased in diabetic patients, but impaired serum capacity to induce cholesterol efflux in these patients was mainly related to low HDL and subclinical inflammation. (J Diabetes Invest, doi: 10.1111/j.2040-1124.2010.00016.x, 2010)

KEY WORDS: Endothelial lipase, Reverse cholesterol transport, Cholesterol efflux
63%. ATP-binding cassette transporter A1 mainly mediates cholesterol efflux to APOA1 and pre-β-HDL, whereas SR-BI mainly mediates cholesterol efflux to mature HDL. Because serum EL concentration is increased in patients with type 2 diabetes, we have investigated whether increased EL is associated with changes in the serum capacity to induce cholesterol efflux and cardiovascular risk in diabetic patients. Cardiovascular risk was assessed non-invasively by measuring arterial stiffness, which provides useful information on vascular health and serves as a surrogate for cardiovascular morbidity and mortality risk\textsuperscript{13,16}.

**METHODS**

Patients with type 2 diabetes mellitus according to World Health Organization criteria were recruited from the diabetes clinics at Queen Mary Hospital. Because we have previously shown that cellular cholesterol efflux to serum was impaired in diabetic patients with incipient or overt nephropathy\textsuperscript{17}, only diabetic patients with normoalbuminuria (urinary albumin excretion rate <30 mg/day) were recruited for the present study to avoid the potential confounding effect of nephropathy. Patients on insulin therapy or lipid lowering agents, or had a history of cardiovascular disease were also excluded. All subjects must have had stable glycaemic control with no change in anti-diabetic therapy for the preceding 3 months. Healthy age-matched controls were recruited from the community. Fasting blood samples were taken for the measurement of glucose, glycated hemoglobin (HbA\textsubscript{1c}), lipids, high sensitivity C-reactive protein (CRP), cholesterol efflux and EL. Arterial stiffness was assessed non-invasively by measuring aortic pulse wave velocity (PWV) using SphygmoCor Vx PWV system (version 7.0; AtCor Medical, Sydney, Australia). Pulse wave velocity was determined by measuring the velocity of the blood pressure waveform between the carotid and femoral artery sites using a single-lead electrocardiogram and a tonometer to measure the pressure pulse waveform sequentially in the two artery sites. The present study was approved by the Ethics Committee of the University of Hong Kong and informed consent was obtained from all subjects.

Plasma total cholesterol and triglyceride were determined enzymatically on a Hitachi 912 analyzer (Roche Diagnostics, Mannheim, Germany). HDL cholesterol was measured using a homogenous method with polyethylene glycol-modified enzymes and alpha-cyclodextrin. LDL cholesterol was calculated by the Friedewald equation or measured directly if plasma triglyceride was >4.5 mmol/L. Plasma APOA1 and APOB were measured by rate nephelometry using the Beckman Array System (Beckman Instruments, Fullerton, CA, USA). HbA\textsubscript{1c} was measured in whole blood using ion-exchange high performance liquid chromatography with the Bio-Rad Variant Haemoglobin Testing System (Bio-Rad Laboratories, Hercules, CA, USA). Plasma high sensitivity CRP was measured by a particle-enhanced immunoturbidimetric assay (Roche Diagnostics) using anti-CRP mouse monoclonal antibodies coupled to latex micro-particles.

Serum EL was measured by competitive ELISA using a rabbit polyclonal antibody specific to human EL (Novus Biologicals, Littleton, CO, USA) as described\textsuperscript{9}. The immunogen used for raising the antibody was an N-terminal synthetic peptide made to the human EL protein sequence. The anti-EL antibody was highly specific and there was no cross-reactivity with hepatic lipase or lipoprotein lipase. Briefly, a 96-well EIA microtiter plate (Costar, Corning, NY, USA) was coated with antigen, the N-terminal synthetic peptide of the human endothelial lipase (Novus Biologicals) in coating buffer overnight at 4°C. The wells were then washed extensively five times and incubated with blocking reagent at room temperature for 2 h. After rinsing five times, equal volume of one-quarter diluted serum samples plus anti-EL antibody (Novus Biologicals) was added. The plate was then incubated for 3 h at room temperature. After another five washes, horseradish peroxidase-conjugated goat-anti-rabbit secondary antibody was added and incubated for 1 h. The plate was finally measured at 450 nm by an ELISA reader. The inter-assay precision was 7.3%.

Fu5AH rat hepatoma cells (a generous gift from Dr GH Rothblat) were used to measure the capacity of serum to induce cholesterol efflux by SR-BI and ABCA1. Fu5AH rat hepatoma cells have a high expression level of SR-BI, but they lack functional ATP-binding cassette transporters. We have previously shown that ABCA1 expression can be specifically induced in Fu5AH cells by stimulating the cells with 22(R)-hydroxycholesterol (22R-HC)/9-cis-retinoic acid (9cRA) and can be used to assay ABCA1-mediated cholesterol efflux, and both SR-BI and ABCA1 expression was not affected by incubation with serum\textsuperscript{17}. The magnitude of ABCA1-mediated cholesterol efflux in our assay is similar to the magnitude of ABCA1-mediated cholesterol efflux reported in efflux assays using RAW264.7 cells stimulated with cyclic adenosine monophosphate (cAMP) to induce ABCA1 expression\textsuperscript{18}. To measure cellular cholesterol efflux to serum, Fu5AH cells were cultured in minimal essential medium (MEM) containing 5% calf serum and labeled with \textsuperscript{3}H]cholesterol for 48 h (1 μCi/well; GE Healthcare Biosciences, Piscataway, NJ, USA). ABCA1-mediated cholesterol efflux was measured using \textsuperscript{3}H]cholesterol-labeled Fu5AH cells treated with 22R-HC (5 μg/mL) and 9cRA (10 μmol/L) and SR-BI-mediated cholesterol efflux was measured in untreated cells. 5% diluted serum in MEM was added and incubated at 37°C for 4 h. Radioactivity was measured in both medium and cells, and the radioactivity released to the medium was expressed as the fraction of the total radioactive cholesterol present in the well. Cholesterol efflux mediated by SR-BI, expressed as percent, was calculated as the amount of label recovered in the medium divided by the total label in each well in untreated cells. Cholesterol efflux mediated by ABCA1 was calculated as fractional cholesterol efflux from 22R-HC/9cRA-treated cells minus fractional cholesterol efflux from untreated control cells. Each assay of cholesterol efflux was carried out in triplicate. The interassay coefficient of variations for SR-BI mediated and ABCA1-mediated cholesterol efflux were 8.0% and 4.5%, respectively.
Results are expressed as means ± SD or as median and interquartile range if the data were not normally distributed. Comparisons between two different groups were carried out using an independent samples t-test, and skewed data were logarithmically transformed before analysis. Pearson’s correlations were used to test the relationship among variables. Multiple linear regression analysis was used to simultaneously assess the relationship between EL and various variables.

RESULTS

The clinical characteristics of the subjects are shown in Table 1. Age, sex and smoking status were not significantly different between the two groups, but body mass index (BMI) and waist circumference were significantly higher in the diabetic patients than controls. In the present study, 32 patients were taking metformin, 23 were taking sulphonylurea and the rest were taking a combination of metformin and sulphonylurea. The mean duration of diabetes was 9.7 ± 4.9 years. Forty-nine percent of patients were on anti-hypertensive agents and 27% had evidence of retinopathy. Fasting glucose, HbA1c and CRP were significantly higher in the diabetic patients than the controls. Pulse wave velocity was also elevated in the diabetic patients.

Table 1 | Clinical characteristic of controls and diabetic patients

| Clinical parameter          | Controls (n = 175) | DM (n = 172) |
|-----------------------------|-------------------|-------------|
| Male/female (%)             | 51/49             | 53/47       |
| Age (years)                 | 49.6 ± 6.6        | 51.3 ± 8.4  |
| Smokers (%)                 | 14                | 128         |
| BMI (kg/m²)                 | 24.5 ± 3.1        | 26.1 ± 4.2**|
| Waist circumference (cm)    | 80.9 ± 9.2        | 87.6 ± 11.9**|
| Systolic BP (mmHg)          | 114 ± 17          | 128 ± 18**  |
| Diastolic BP (mmHg)         | 74 ± 10           | 77 ± 9*     |
| PWV (m/s)                   | 7.1 ± 1.1         | 90 ± 15**   |
| Fasting glucose (mmol/L)    | 4.84 ± 0.82       | 8.60 ± 2.22**|
| HbA1c (%)                   | 6.76 ± 0.80       | 8.21 ± 1.35**|
| CRP (mg/L)                  | 0.75 (0.41–1.70)  | 1.35 (0.63–2.49)**|

Values are mean ± SD or median (interquartile range). *P < 0.05, **P < 0.01 vs controls. BMI, body mass index; BP, blood pressure; CRP, C-reactive protein; DM, diabetes mellitus; HbA1c, glycated hemoglobin; PWV, pulse wave velocity.

Plasma levels of lipids and apolipoproteins are shown in Table 2. Plasma HDL cholesterol and APOA1 were reduced and triglyceride was higher in the diabetic group. SR-BI mediated cholesterol efflux to serum was impaired in the diabetic patients, whereas no significant difference in ABCA1 mediated cholesterol efflux to serum was found. Because plasma HDL levels differed significantly between diabetic patients and controls, data were also analyzed after adjusting for HDL cholesterol levels, and SR-BI mediated cholesterol efflux to serum remained impaired in the diabetic patients (P < 0.01). Serum EL was significantly increased in the diabetic patients (Table 2). There was no significant correlation between serum EL and HDL in controls (r = −0.08) or diabetic patients (r = −0.05), whilst there was a weak trend between serum EL and logCRP (controls r = 0.13, P = 0.08; diabetics r = 0.14, P = 0.07). No association between serum EL and PWV was observed in either the diabetic patients or controls. In the diabetic patients, PWV correlated with logCRP (r = 0.20, P < 0.05) and SR-BI mediated cholesterol efflux (r = −0.17, P < 0.05) in addition to its strong correlation with systolic BP (r = 0.54, P < 0.01). Neither the association between PWV and logCRP nor SR-BI mediated cholesterol efflux remained significant after adjusting for blood pressure.

Univariate analysis was carried out to investigate which clinical parameters were associated with cholesterol efflux. In the control subjects, serum EL correlated inversely with SR-BI mediated cholesterol efflux (Figure 1a) but not with ABCA1 mediated cholesterol efflux. SR-BI mediated cholesterol efflux to serum also correlated with HDL (r = 0.59, P < 0.001), APOAI (r = 0.60, P < 0.001), waist circumference (r = −0.16, P < 0.05), but not with logCRP. However, in the diabetic patients, there was only a trend towards an association between serum EL and SR-BI mediated cholesterol efflux (Figure 1b) and there was no correlation with ABCA1 mediated cholesterol efflux. In the diabetic cohort, SR-BI mediated cholesterol efflux to serum correlated with HDL (r = 0.62, P < 0.001), APOAI (r = 0.51, P < 0.001), waist circumference (r = −0.19, P < 0.05) and logCRP (r = −0.32, P < 0.001), whereas ABCA1 mediated cholesterol efflux to serum correlated only with HDL (r = 0.23, P < 0.01).

To determine whether serum EL was an independent determinant of SR-BI mediated cholesterol efflux to serum, multiple linear regression analysis was carried out including variables that had significant associations. In controls, age, sex, waist circumference, smoking status, HDL and serum EL were entered into the model, and the results are shown in Table 3. Plasma HDL,
serum EL and waist circumference were the independent determinants of SR-BI mediated cholesterol efflux to serum in controls, accounting for 24%, 10% and 2% of the variability of cholesterol efflux respectively. In contrast, the major independent determinants of SR-BI mediated cholesterol efflux in the diabetic patients were HDL, logCRP and age (Table 4), accounting for 26%, 6% and 2% of the variability of cholesterol efflux, respectively. Forcing serum EL and/or HbA1c into the model did not change the results.

DISCUSSION

Efflux of free cholesterol from cells is an early step of reverse cholesterol transport and the efficiency of cellular cholesterol efflux is influenced by the concentrations of lipoproteins that act as cholesterol acceptors and the activities of serum proteins, such as lipases and lipid transfer proteins involved in the remodeling of lipoproteins19. The capacity of whole serum or plasma to stimulate cholesterol efflux from cells has been used by a number of laboratories as a means to investigate reverse cholesterol transport, as whole serum provides integrated information on lipoproteins and serum components involved in promoting cholesterol efflux from cells20–24. The capacity of serum to induce cholesterol efflux has been shown to be an independent predictor of coronary artery atherosclerosis in clinical studies22,23. Serum capacity to induce SR-BI-mediated cellular cholesterol efflux is reduced in type 2 diabetic patients with or without diabetic complications17,24 and ABCA1-mediated cholesterol efflux is also impaired in patients with nephropathy17. Similar to previous studies, we have shown that serum capacity to induce SR-BI-mediated cellular cholesterol efflux is decreased in diabetic patients, whereas no significant reduction was seen in serum capacity to induce ABCA1-mediated cholesterol efflux.

In the present study, we have determined whether serum capacity to induce cellular cholesterol efflux was related to serum EL level. As expected, plasma HDL level was the major determinant of serum capacity to induce cellular cholesterol efflux, be it mediated by SR-BI or ABCA1. In healthy control subjects, there was a weak but significant inverse correlation

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**Table 3** | Multiple regression analysis with scavenger receptor class B type I mediated cholesterol efflux to serum as the dependent variable in control subjects

| Regression coefficient | SE of regression coefficient | P-value |
|------------------------|-----------------------------|---------|
| Intercept              | 6.103                       | 3.142   | <0.001 |
| Age                    | 0.005                       | 0.028   | 0.054  |
| Sex                    | 0.099                       | 0.015   | 0.849  |
| Waist circumference     | 0.063                       | 0.029   | 0.031  |
| Smoker, yes/no         | -0.021                      | 0.040   | 0.960  |
| HDL                    | 5.222                       | 0.647   | <0.001 |
| EL                     | -0.196                      | 0.022   | <0.001 |

Adjusted $R^2$ of model = 36%. EL, endothelial lipase; HDL, high-density lipoprotein.

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**Table 4** | Multiple regression analysis with scavenger receptor class B type I mediated cholesterol efflux to serum as the dependent variable in diabetic subjects

| Regression coefficient | SE of regression coefficient | P-value |
|------------------------|-----------------------------|---------|
| Intercept              | 12.188                      | 1.910   | <0.001 |
| Age                    | -0.049                      | 0.019   | 0.012  |
| Sex                    | 0.295                       | 0.353   | 0.404  |
| Waist circumference     | 0.005                       | 0.015   | 0.736  |
| Smoker, yes/no         | 0.431                       | 0.266   | 0.108  |
| HDL                    | 3.690                       | 0.526   | <0.001 |
| Log (CRP)              | -1.061                      | 0.412   | 0.010  |

Adjusted $R^2$ of model = 34%. CRP, C-reactive protein; HDL, high-density lipoprotein.

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**Figure 1** | Correlation between serum endothelial lipase and scavenger receptor class B type I mediated cholesterol efflux to serum in (a) controls and (b) diabetic patients. SR-BI, scavenger receptor class B type I.
between EL and serum capacity to induce SR-BI-mediated cellular cholesterol efflux. This is consistent with animal studies showing that increased EL was associated with a decrease in efflux potential of serum through SR-BI$^{12,14}$. EL can reduce SR-BI-dependent cholesterol efflux partly by altering the chemical composition and physical properties of HDL$^{12}$. EL has both catalytic and non-catalytic functions and can mediate binding of lipoproteins independent of its enzymatic activity$^{25}$. Because the association between EL and serum capacity to induce SR-BI-mediated cellular cholesterol efflux remains independent of HDL on analysis, whether this might be related to the non-catalytic function of EL warrants further investigation.

In contrast to the findings in healthy controls, only a trend towards an association between serum EL and serum capacity to induce cellular cholesterol efflux was seen in type 2 diabetic patients. In these patients, CRP turned out to be an important determinant of serum capacity to induce SR-BI-mediated cellular cholesterol efflux. Inflammation has a major effect on reverse cholesterol transport$^{26}$ and our data would suggest that the influence of inflammation on serum capacity to induce SR-BI-mediated cellular cholesterol efflux might override that of EL in diabetic subjects. Chronic subclinical inflammation has been shown in patients with type 2 diabetes mellitus$^{27}$. Although inflammation has been shown to activate EL$^{28}$, this is unlikely to be the main mechanism whereby inflammation impairs serum capacity to induce cholesterol efflux in our diabetic patients, because we did not find any significant correlation between EL and cholesterol efflux. Inflammation can influence serum capacity to induce cholesterol efflux by a number of mechanisms independent of EL. Inflammation also activates secretory phospholipase A2 and inflammatory remodeling of HDL impairs its capacity to serve as cholesterol acceptor ex vivo and in vivo$^{26,29-30}$. In addition, inflammation also affects serum proteins that are involved in promoting cholesterol efflux from cells. In human subjects, attenuation of lecithin cholesterol acyltransferase and cholesterol ester transfer protein activity has been reported during infection and inflammation$^{31,32}$. Taken together, subclinical inflammation might play a role in reducing serum capacity to induce cellular cholesterol efflux in type 2 diabetes mellitus. However, the present study was limited by its cross-sectional nature, and we could only show an association and not a causal relationship.

Because EL might be proatherogenic and plasma EL has been associated with coronary artery calcification score in healthy individuals with a family history of premature coronary heart disease$^{6}$, we have also determined in the present study whether plasma EL is correlated with arterial stiffness in diabetic patients. Vascular stiffness reflects both functional and structural changes in the artery wall that precede and accompany atherosclerosis. Pulse wave velocity reflects arterial stiffness and is regarded as a surrogate marker of severity of atherosclerosis$^{33}$. Experimental studies have shown that aortic PWV increases with the development of atherosclerosis in primates$^{34}$ and there is a strong association between aortic PWV with intima-media thickness and severity of plaques evaluated by ultrasonographic images$^{15}$. We have shown that arterial stiffness is increased in diabetic patients without overt cardiovascular disease, but we did not find an association between EL and PWV in our diabetic patients.

In conclusion, in patients with type 2 diabetes, serum EL concentration was increased, but impaired serum capacity to induce cholesterol efflux in these patients was mainly related to low HDL and subclinical inflammation.

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The authors declare no conflict of interest.

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