## **Online Supplemental Materials**

**Supplementary Table 1.** Comparisons of HDL cholesterol (HDL-C) and ApoAI levels between diabetics and non-diabetics subjects, with and without adjustment for sex. Statistics presented as Pearson's correlation (95% CI).

Measure	Unadjusted Mean Difference (95% CI)	Unadjusted P-value	Adjusted Mean Difference (95% CI)	Adjusted P-value	
HDL-C	-3.69(-16.91,9.52)	0.56	-3.76(-17.53,10.00)	0.57	
ApoAl	4.42(-14.87,23.70)	0.63	4.04(-15.70,23.78)	0.67	

**Supplementary Table 2.** Correlation of total and ABCA-independent cholesterol efflux with hyperglycemia (HbA<sub>1c</sub>) and ApoAI glycation at Lys-12 and Lys-133 residues. Statistics presented as Pearson's correlation (95% CI).

Cholesterol Efflux	HbA1c		Lys-12 (	Slycation	Lys-133 Glycation		
	r	P	r	P	r	P	
Total efflux	-0.28	0.03	-0.53	0.001	-0.13	0.16	
ABCA1-independent	-0.07	0.31	-0.22	0.06	-0.01	0.70	

HbA<sub>1c</sub>: glycated hemoglobin, ABCA1: ATP-binding cassette (ABC) family transporter.

**Supplementary Table 3.** Comparisons of the fractional catabolic rates (FCRs) of ApoAI, ApoAI glycation and HDL functions between T2DM patients and healthy controls with adjustment for HbA<sub>1c</sub>. The parameters that were not different after the adjustment for HbA<sub>1c</sub> are highlighted in bold.

Measure	Unadjusted Mean Difference (95% CI)	Unadjusted P- Value	Adjusted Mean Difference (95% CI)	Adjusted P- Value					
Fractional catabolic rate of ApoAl									
ApoAl	0.01 (0.00,0.01)	0.01) <b>&lt;0.001</b> 0.01 (0.00,0.01)		0.031					
Glycation measures at different lysine sites of ApoAl									
logApoA1K133	0.25 (-0.00,0.50)	0.054	0.39 (-0.09,0.88)	0.100					
logApoA1K12	0.25 (0.15,0.36)	<0.001	0.10 (-0.08,0.27)	0.240					
logApoA1K205	0.71 (0.13,1.30)	0.021	0.88 (-0.26,2.01)	0.120					
logApoA1K96	0.01 (-0.29,0.30)	0.960	-0.04 (-0.61,0.53)	0.880					
	HDL function								
PON1	-97.14 (-159.02,-35.27)	0.005	-55.76 (-173.58,62.06)	0.33					
ABCA1- independent	-1.15 (-2.43,0.13)	0.074	-2.20 (-4.65,0.26)	0.075					
Total efflux	-2.31 (-3.66,-0.95)	0.002	-2.98 (-5.64,-0.32)	0.031					
ABCA1-dependent	-1.16 (-1.90,-0.43)	0.004	-0.81 (-2.26,0.64)	0.250					

FCR: fractional catabolic rate, ApoA1K12: ApoAI glycation at lysine 12 site, ApoA1K96: ApoAI glycation at lysine 96 site ApoA1K133: ApoAI glycation at lysine 133 site, ApoA1K205: ApoAI glycation at lysine 205 site HbA<sub>1c</sub>: glycated hemoglobin; HDL: high density lipoprotein, PON1: paraoxonase 1.

**Supplementary Table 4.** List of the proteins associated with ApoAI. Two identical sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGEs) were performed for ApoAI-specific Western blotting and proteomics analysis. Based on the molecular weight markers and Western blot analysis matching, corresponding silver-stained gel bands containing ApoAI were analyzed by mass spectrometry. Proteins identified in each band are check marked.

A	Red.		Molecular	Control			T2DM		
Accession Number			Weight (kDa)	27 kDa	50 kDa	75 kDa	27 kDa	50 kDa	75 kDa
P01009	A1AT	Alpha-1-antitrypsin	47		$\checkmark$			$\checkmark$	$\sqrt{}$
P01008	ANT3	Antithrombin-III	53		$\sqrt{}$		<b>V</b>	$\sqrt{}$	$\checkmark$
P02647	APOA1	Apolipoprotein A-I	31	$\sqrt{}$	$\sqrt{}$	$\sqrt{}$	$\sqrt{}$	$\sqrt{}$	$\sqrt{}$
P04114	APOB	Apolipoprotein B-100	516		$\sqrt{}$		$\sqrt{}$	$\checkmark$	$\sqrt{}$
P02649	APOE	Apolipoprotein E	36			$\sqrt{}$		$\sqrt{}$	
P01857	IGHG1	Immunoglobulin heavy constant gamma 1	36		√			<b>√</b>	<b>V</b>
P01871	IGHM	Immunoglobulin heavy constant mu	49		$\sqrt{}$		$\sqrt{}$	$\sqrt{}$	V
Q14624	ITIH4	Inter-alpha-trypsin inhibitor heavy chain H4	103			√	√	<b>V</b>	√
P02768	ALBU	Serum albumin	69		√	√	√	√	√
P04004	VTNC	Vitronectin	54		√		√	√	$\sqrt{}$
P02765	FETUA	Alpha-2-HS-glycoprotein	39		√				
P06727	APOA4	Apolipoprotein A-IV	45			V			
P05090	APOD	Apolipoprotein D	21			V			
O95445	APOM	Apolipoprotein M	21			$\sqrt{}$			
P04217	A1BG	Alpha-1B-glycoprotein	54						V
P08697	A2AP	Alpha-2-antiplasmin	55					√	V
P01023	A2MG	Alpha-2-macroglobulin	163					√	V
P04003	C4BPA	C4b-binding protein alpha chain	67				$\sqrt{}$	$\sqrt{}$	$\sqrt{}$
P09871	C1S	Complement C1s subcomponent	77					<b>V</b>	
P01024	CO3	Complement C3	187				√	√	V
P0C0L4	CO4A	Complement C4-A	193				√	√	$\sqrt{}$
P06396	GELS	Gelsolin	86					√	
P05546	HEP2	Heparin cofactor 2	57				√	√	$\sqrt{}$
P04196	HRG	Histidine-rich glycoprotein	60						V
P19823	ITIH2	Inter-alpha-trypsin inhibitor heavy chain H2	106						√
P55058	PLTP	Phospholipid transfer protein	55				√	$\sqrt{}$	$\sqrt{}$
P00747	PLMN	Plasminogen	91					√	
P00734	THRB	Prothrombin	70				<b>V</b>	$\sqrt{}$	$\sqrt{}$
P02787	TRFE	Serotransferrin	77					$\sqrt{}$	$\sqrt{}$

**Supplementary Figure 1.** <sup>2</sup>H-labeling of native and glycated ApoAI analyzed by mass spectrometry. ApoB100-depleted plasma samples from a patient with type 2 diabetes at the baseline and after 4 days of <sup>2</sup>H<sub>2</sub>O labeling experiment were digested with trypsin and analyzed by high resolution mass spectrometry. To illustrate the changes in isotope distribution, the heavy isotopomers (M<sub>1</sub>-M<sub>3</sub>) were normalized relative to M<sub>0</sub>. After 4 days of <sup>2</sup>H<sub>2</sub>O labeling, the intensities of heavy isotopomers, in particular M1, were increased relative to monoisotopic peak in both non-glycated VSFLSALEEYTK (**A**) and glycated Q<sup>Gly</sup>**K**LHELQEK (**B**) ApoAI peptides. However, the extent of the increase in the glycated peptide was substantially higher than in the non-glycated peptide, indicating that the glycated peptide has higher turnover rate and shorter half-life.

