Supplementary Material

Plasma high density lipoprotein cargo is altered in Alzheimer's disease and is associated with regional brain volume

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Supplementary Table 1

	НС	HC-Conv	AD	р
µgHDL-assApoA-I/ml PL	493 ± 144	460 ± 114	431 ± 94	0.0091
µgHDL-assApoA-II/ml PL	138 ± 54	123 ± 49	106 ± 32	0.0014
µgHDL-assApoC-I/ml PL	138 ± 46	130 ± 40	116 ± 37	0.0356
µgHDL-assApoC-III/ml PL	28.8 ± 13.0	26.5 ± 12.0	24.4 ± 9.7	0.2180
µgHDL-assApoD/ml PL	21.1 ± 4.5	20.0 ± 5.2	21.1 ± 5.0	0.9650
µgHDL-assApoE/ml PL	6.75 ± 3.34	5.78 ± 2.39	4.49 ± 2.23	0.0072
µgHDL-assApoH/ml PL	248 ± 69	231 ± 57	208 ± 67	0.0046
µgHDL-assApoJ/ml PL	22.3 ± 6.1	19.7 ± 4.3	19.1 ± 4.6	0.0008
µgHDL-assCRP/ml PL	2.47 ± 1.03	2.12 ± .51	1.95 ± .99	0.0009
µgHDL-assSAA/ml PL	5.92 ± 5.91	4.15 ± 4.27	8.22 ± 8.88	0.7730

Comparison of HDL-associated cargo (ApoA-I, ApoA-II, ApoC-I, ApoC-III, ApoD, ApoE, ApoH, ApoJ, CRP and SAA) per ml of plasma in stable HC, HC-Converters and AD.

Supplementary Table 1. Values are presented as mean \pm S.D. Generalized Linear Model analyses were performed unadjusted. Data are presented as raw values, but statistical analysis was performed on log-transformed data to better approximate normal distribution. Analysis was considered significant with p < 0.05 (bold).

Supplementary Table 2

Partial correlation of HDL-cholesterol and HDL-cargo (ApoA-II, ApoC-I, ApoC-III, ApoD, ApoE, ApoH, ApoJ, CRP and SAA) expressed as ratio to ApoA-I with MMSE (at t0) in individuals with AD.

	MMSE t0	
	r ^a	p ^a
µgCholesterol/µgApoA-I	.023	.838
ngApoA-II/µgApoA-I	102	.376
ngApoC-I/µgApoA-I	104	.364
ngApoC-III/µgApoA-I	.078	.498
ngApoD/µgApoA-I	022	.852
ngApoE/µgApoA-I	.001	.995
ngApoH/µgApoA-I	.134	.244
ngApoJ/µgApoA-I	.018	.876
ngCRP/µgApoA-I	010	.933
ngSAA/µgApoA-I	050	.665

Supplementary Table 2. Pearson's correlation adjusted for age, gender, and ApoE ϵ 4-carrier status was performed between HDL-cargo and MMSE. Analysis was considered significant with p < 0.05. ^a Data were adjusted for age, gender, and ApoE ϵ 4-carrier status.

Supplementary Figure 1



Supplementary Figure 1. Graphical representation of individual distribution across clinical classification groups.