

	Male		Female		Total
	Low BF%	High BF%	Low BF%	High BF%	
<i>n</i>	11	7	7	7	32
<u>Race (<i>n</i>; %):</u>					
White	7; 64%	7; 100%	6; 86%	4; 57%	24; 75%
Black	1; 9%	0; 0%	0; 0%	1; 14%	2; 6%
Asian	2; 18%	0; 0%	1; 14%	1; 14%	4; 13%
Other	1; 9%	0; 0%	0; 0%	1; 14%	2; 6%
Age (yr)	29.4 ± 6.9	31.3 ± 6.8	25.0 ± 3.9	31.2 ± 5.1	29.2 ± 6.2
Height (cm)	178.9 ± 7.9	181.2 ± 4.1	167.4 ± 1.8	162.7 ± 6.5	173.3 ± 9.5
Weight (kg)	78.6 ± 10.0	107.2 ± 16.4	73.3 ± 6.7	88.0 ± 20.4	85.7 ± 18.2
BMI (kg/m ²)	24.6 ± 2.8	32.6 ± 4.3	24.7 ± 4.1	33.1 ± 6.4	28.2 ± 5.9
Body Fat (%)	21.7 ± 6.3	36.8 ± 4.2	32.2 ± 4.4	44.3 ± 3.2	32.2 ± 10.0

Supplemental Table 1: *Participant characteristics.* BF% = body fat percentage.

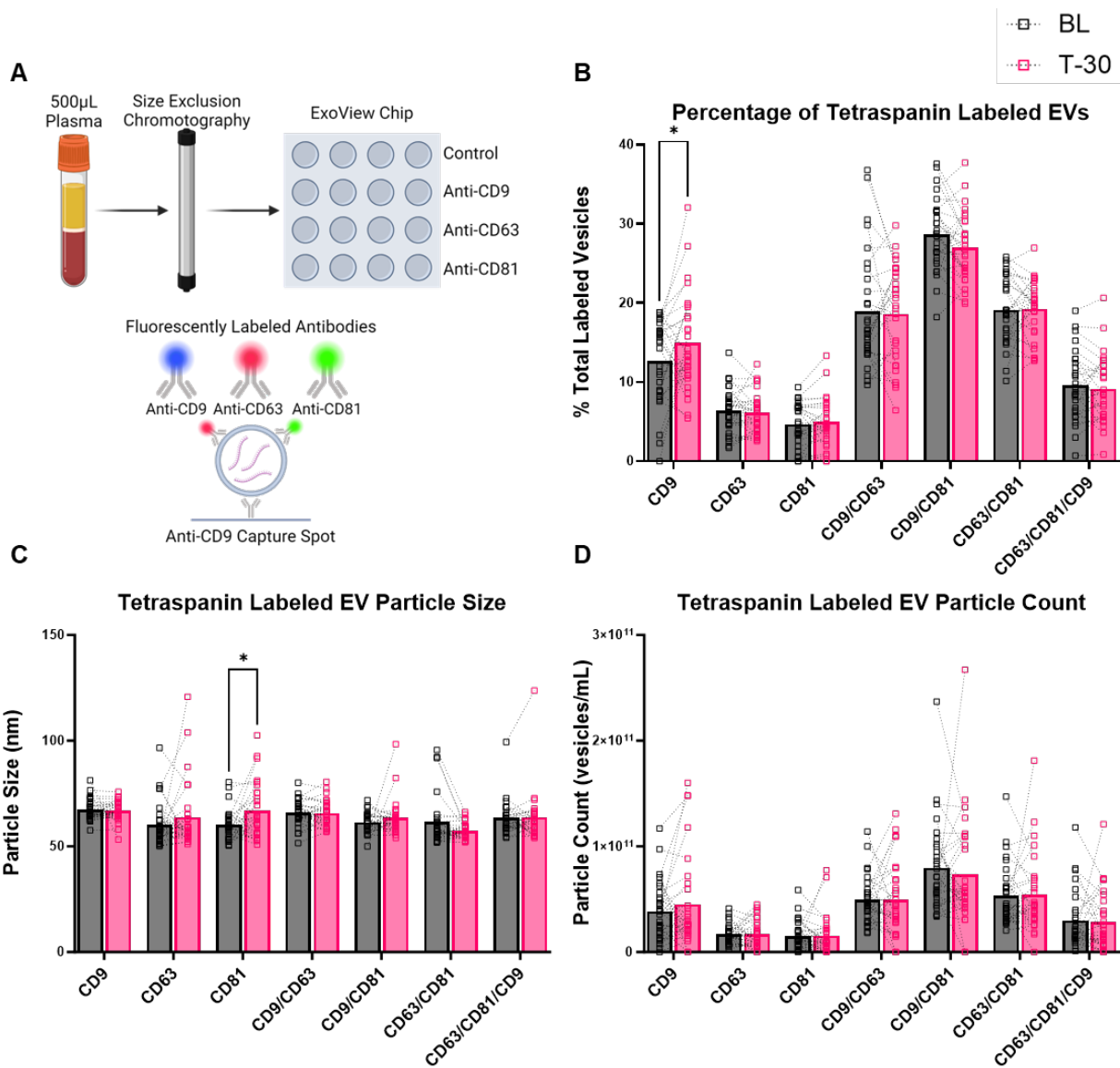
miR-1 Relative Abundance (Fold Change from BL)	Predictors	β -Coefficient	p -value
Muscle	Age	-0.041	0.081
T-0 EV	Body Mass Index	0.118	0.079
T-30 EV	Bilateral Thigh Muscle Mass	0.000	0.540
T-60 EV	Bilateral Thigh Muscle Mass	0.000	0.262
T-90 EV	Estimated Visceral Adipose Mass	0.418	0.070
Adipose	Sex	0.954	0.257

Supplemental Table 2: *Low sample size precludes significant findings for predictors of changes in muscle, serum EV, and adipose miR-1. EV = extracellular vesicle; T-0 = 0-minute timepoint; T-30 = 30-minute timepoint; T-60 = 60-minute timepoint; T-90 = 90-minute timepoint.*

Gene	log(Fold Change)	p-value
CAV2	-0.399	0.049
CYB5B	-0.282	0.046
E2F5	-1.507	0.025
FAM169A	-2.622	0.038
GCLC	-0.420	0.023
GLCCI1	-0.733	0.013
KANK4	-1.279	0.049
MAL2	-0.917	0.039
PGD	-0.432	0.039
PPP2R5A	-0.308	0.045
RTN4IP1	-0.763	0.018
TMEM120B	-0.603	0.016
TRIM6	-1.207	0.012
ZNF112	-1.080	0.041
ZNF566	-0.397	0.045
ZNF677	-0.578	0.022

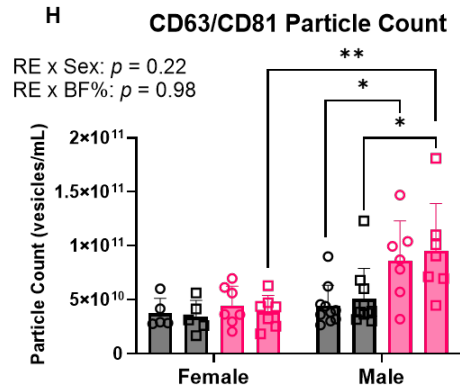
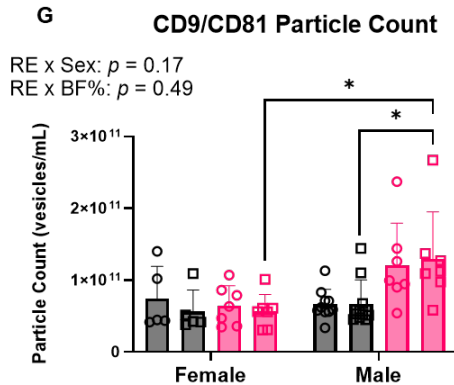
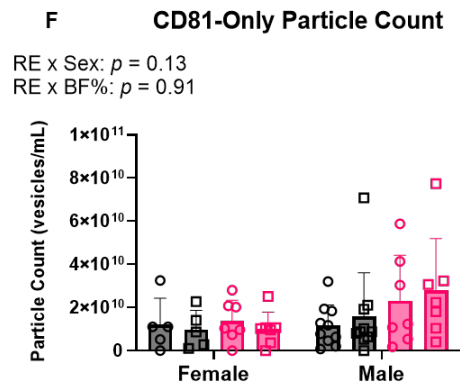
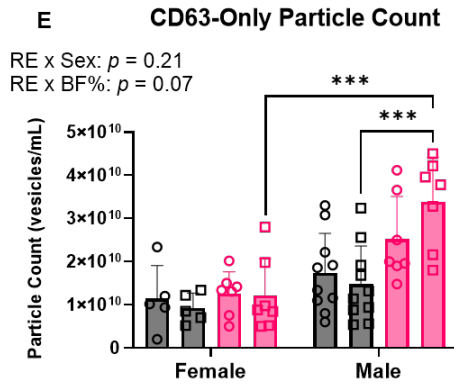
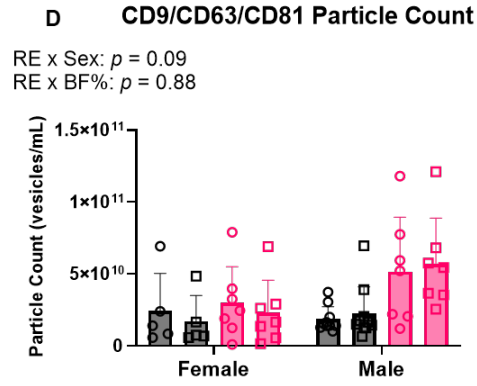
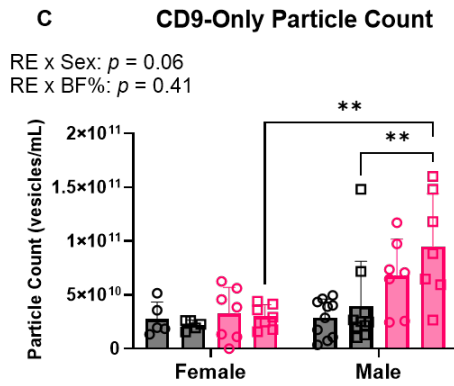
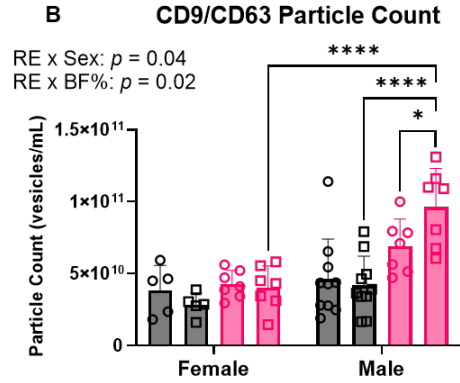
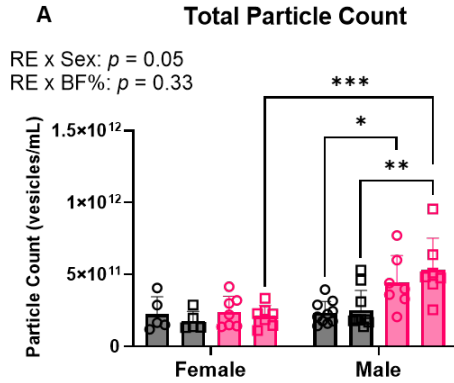
Supplemental Table 3: *Predicted miR-1 targets identified as differentially downregulated*

through RNA-sequencing analyses. CAV2 = caveolin 2; CYB5B = cytochrome b5 type B; E2F5 = E2F transcription factor 5; FAM169A = family with sequence similarity 169 member A; GCLC = glutamate-cysteine ligase catalytic subunit; GLCCI1 = glucocorticoid induced 1; KANK4 = KN motif and ankyrin repeat domains 4; MAL2 = T cell differentiation protein 2; PGD = phosphogluconate dehydrogenase; PPP2R5A = protein phosphatase 2 regulatory subunit B'alpha; RTN4IP1 = reticulon 4 interacting protein 1; TMEM120B = transmembrane protein 120B; TRIM6 = tripartite motif-containing protein 6; ZNF112 = zinc finger protein 112; ZNF566 = zinc finger protein 566; ZNF677 = zinc finger protein 677.

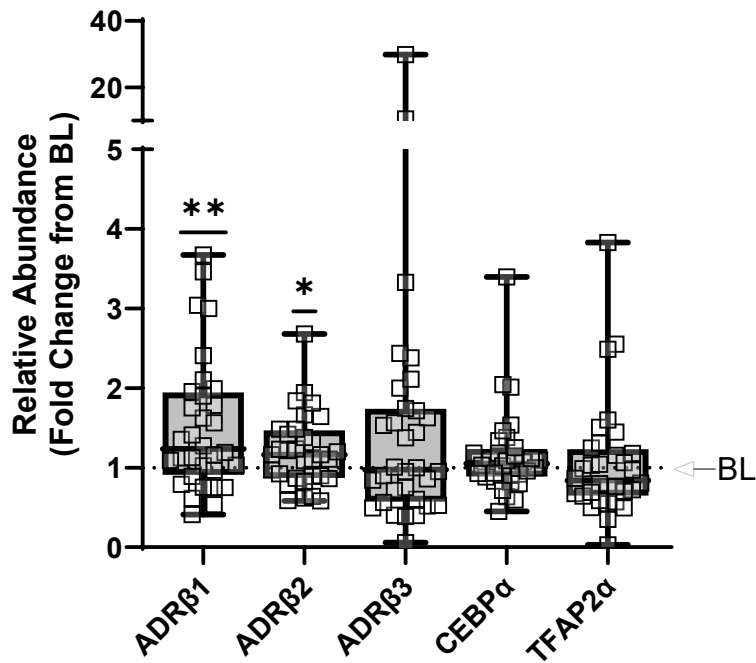


Supplemental Figure 1: Acute resistance exercise alters extracellular vesicle (EV) size and population composition in humans. (A) Schematic diagram of the workflow for the protein-capture ExoView platform. Changes in the (B) proportion of labeled EVs, (C) EV particle size, and (D) EV particle count in response to exercise in different populations of tetraspanin-labeled EVs ($n = 29$). Data were analyzed using mixed-effects analyses with Šidák corrections for multiple comparisons. * = $p < 0.05$. BL = baseline; T-30 = 30-minute timepoint.

Low BF% BL High BF% BL
 Low BF% T-30 High BF% T-30



Supplemental Figure 2: Sex and adiposity alter extracellular vesicle response to acute resistance exercise. Participants were grouped by sex into high (male $n = 7$, female $n = 7$) and low (male $n = 10$, female $n = 5$) body fat percentage (BF%) groups. Interactions of sex and BF% with the effects of an acute bout of resistance exercise (RE) on (A) CD63-only, (B) CD9/CD63 double positive, (C) CD9-only, (D) CD9/CD63/CD81 triple positive, (E) total, (F) CD81-only, (G) CD9/CD81 double positive, and (H) CD63/CD81 double positive extracellular vesicles. Data were analyzed using three-way ANOVAs with Šidák corrections for multiple comparisons. * = $p < 0.05$; ** = $p < 0.01$; *** = $p < 0.001$; **** = $p < 0.0001$. BL = baseline; T-30 = 30-minute timepoint.



Supplemental Figure 3: Resistance exercise increases the expression of ADRβ1 and ADRβ2, but not ADRβ3, CEBPα, or TFAP2α. Transcript abundance of β-adrenergic receptor (*ADRβ*) 1, *ADRβ2*, *ADRβ3*, CCAAT/enhancer-binding protein alpha (*CEBPα*), and transcription factor AP-2 alpha (*TFAP2α*) in adipose tissue relative to baseline (BL; denoted by the dotted line). Data are expressed with min-to-max box plots and were compared using one-sample Wilcoxon t-tests. * = $p < 0.05$; ** = $p < 0.01$.