



## Accumulation of saturated intramyocellular lipid is associated with insulin resistance<sup>S</sup>

David B. Savage,\* Laura Watson,† Katie Carr,† Claire Adams,\* Soren Brage,§ Krishna K. Chatterjee,\*† Leanne Hodson,\*\* Chris Boesch,†† Graham J. Kemp,§§ and Alison Sleigh<sup>1,\*†,\*\*\*</sup>

Metabolic Research Laboratories, Wellcome Trust-MRC Institute of Metabolic Science,\* and National Institute for Health Research/Wellcome Trust Clinical Research Facility, Cambridge University Hospitals NHS Foundation Trust,† Cambridge Biomedical Campus, Cambridge, United Kingdom; Oxford Centre for Diabetes, Endocrinology and Metabolism,\*\* Radcliffe Department of Medicine, University of Oxford, Oxford, United Kingdom; Department of Clinical Research and Radiology,†† AMSM, University Bern, Bern, Switzerland; Department of Musculoskeletal Biology,§§ University of Liverpool and MRC–Arthritis Research UK Centre for Integrated Research into Musculoskeletal Ageing, Liverpool, United Kingdom; Wolfson Brain Imaging Centre\*\*\* and MRC Epidemiology Unit,§ University of Cambridge School of Clinical Medicine, Cambridge, United Kingdom

**Abstract** Intramyocellular lipid (IMCL) accumulation has been linked to both insulin-resistant and insulin-sensitive (athletes) states. Biochemical analysis of intramuscular triglyceride composition is confounded by extramyocellular triglycerides in biopsy samples, and hence the specific composition of IMCLs is unknown in these states. <sup>1</sup>H magnetic resonance spectroscopy (MRS) can be used to overcome this problem. Thus, we used a recently validated <sup>1</sup>H MRS method to compare the compositional saturation index (CH<sub>2</sub>:CH<sub>3</sub>) and concentration independent of the composition (CH<sub>3</sub>) of IMCLs in the soleus and tibialis anterior muscles of 16 female insulin-resistant lipodystrophic subjects with that of age- and gender-matched athletes (*n* = 14) and healthy controls (*n* = 41). The IMCL CH<sub>2</sub>:CH<sub>3</sub> ratio was significantly higher in both muscles of the lipodystrophic subjects compared with controls but was similar in athletes and controls. IMCL CH<sub>2</sub>:CH<sub>3</sub> was dependent on the IMCL concentration in the controls and, after adjusting the compositional index for quantity (CH<sub>2</sub>:CH<sub>3adj</sub>), could distinguish lipodystrophics from athletes. This CH<sub>2</sub>:CH<sub>3adj</sub> marker had a stronger relationship with insulin resistance than IMCL concentration alone and was inversely related to VO<sub>2max</sub>.<sup>1</sup> The association of insulin resistance with the accumulation of saturated IMCLs is consistent with a potential pathogenic role for saturated fat and the reported benefits of exercise and diet in insulin-resistant states.—Savage, D. B., L. Watson, K. Carr, C. Adams, S. Brage, K. K. Chatterjee, L. Hodson, C. Boesch, G. J. Kemp, and A. Sleigh. **Accumulation of saturated**

**intramyocellular lipid is associated with insulin resistance.** *J. Lipid Res.* 2019. 60: 1323–1332.

**Supplementary key words** triglycerides • fatty acids • lipodystrophies • spectroscopy • lipid composition • muscle • exercise • in vivo

After it was demonstrated that <sup>1</sup>H magnetic resonance spectroscopy (MRS) can noninvasively distinguish intramyocellular lipids (IMCLs) from extramyocellular lipids (EMCLs) (1, 2), associations were reported between soleus (SOL) IMCL accumulation and insulin resistance independent of fat mass (3–5). Given that skeletal muscle represents the primary site for postprandial glucose disposal (6), these findings were of considerable physiological interest. Furthermore, these data strongly supported the link between ectopic fat accumulation and insulin resistance (7, 8). Although it soon became clear that triglycerides themselves were unlikely to be involved in causing insulin resistance, intramuscular triglyceride content does seem to correlate with insulin resistance in some states (5, 9–12). One particularly striking and surprising exception was reported in athletes, in which histological studies suggested that neutral lipid accumulation was a feature of skeletal muscle in insulin-sensitive, endurance-trained athletes (13, 14), and this finding has led to the now widely cited

This work was supported by grants from the UK National Institute for Health Research (NIHR) Cambridge Biomedical Research Centre; the UK Medical Research Council Centre for Obesity and Related Metabolic Diseases; the Clinical Research Infrastructure Grant; and Wellcome Trust Grant 107064 (D.B.S.). A.S. was supported by the NIHR via an award to the NIHR Cambridge Clinical Research Facility. The views expressed are those of the authors and not necessarily those of the NHS, NIHR, or Department of Health and Social Care.

\*Author's Choice—Final version open access under the terms of the Creative Commons CC-BY license.

Manuscript received 19 December 2018 and in revised form 11 March 2019.

Published, JLR Papers in Press, May 2, 2019

DOI <https://doi.org/10.1194/jlr.M091942>

Copyright © 2019 Savage et al. Published by The American Society for Biochemistry and Molecular Biology, Inc.

This article is available online at <http://www.jlr.org>

Abbreviations: AGLD, acquired generalized lipodystrophy; EMCL, extramyocellular lipid; FPLD, familial partial lipodystrophy; HOMA-IR, Homeostatic Model Assessment of Insulin Resistance; IMCL, intramyocellular lipid; LD, lipodystrophic; MRS, magnetic resonance spectroscopy; NIHR, National Institute for Health Research; SOL, soleus; TA, tibialis anterior.

<sup>1</sup>To whom correspondence should be addressed.

e-mail: [as626@cam.ac.uk](mailto:as626@cam.ac.uk)

<sup>S</sup>The online version of this article (available at <http://www.jlr.org>) contains a supplement.

notion of an “athlete’s paradox” (13). This concept is consistent with the idea that triglyceride content itself is not casually involved in insulin resistance and has prompted several efforts to identify the lipid intermediates responsible for causing insulin resistance or preserving the insulin sensitivity of athletes.

Saturated fat has been implicated in the pathogenesis of metabolic disease (15, 16), and we have recently described and validated (using IMCL/EMCL-simulated phantoms of known composition) a <sup>1</sup>H MRS method that provides an in vivo compositional marker of IMCLs that primarily reflects the degree of saturation of the FA chains within triglycerides (17). This marker, which we call the IMCL saturation index (CH<sub>2</sub>:CH<sub>3</sub>), utilizes good-quality spectra acquired at 3T with a short echo time and compares the CH<sub>2</sub> resonance located at 1.3 ppm (which is influenced by both concentration and composition) with that of the CH<sub>3</sub> resonance at 0.9 ppm (which is independent of triglyceride composition); this is illustrated in Fig. 1. Figure 1 also shows that using a concentration of hydrogen that resonates at 1.3 ppm (CH<sub>2</sub>) to represent the concentration of lipids without knowing the underlying composition, as has been the practice in virtually all published <sup>1</sup>H MRS studies of IMCLs so far, confounds the contributions of both the concentration of lipids and their composition. This can potentially lead to a significant error in estimating the concentration: the composition would contribute as much as 50% to the observed signal (equivalent to a 100% theoretical increase in the signal) if the pool were stearic acid instead of linoleic acid. Therefore, we used the IMCL CH<sub>3</sub> peak at 0.9 ppm to estimate the total concentration of IMCLs, as this is independent of the degree of saturation of the FA chains within triglycerides (i.e., composition). We call this the composition-

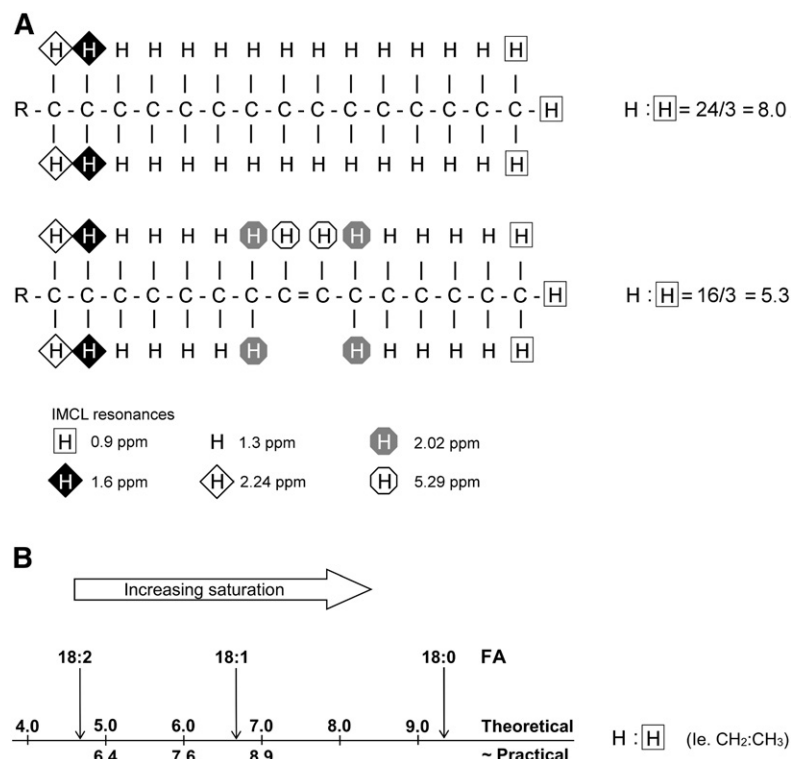
independent IMCL concentration estimate to distinguish it from the conventional estimate using CH<sub>2</sub>.

Lipodystrophy is a rare cause of severe insulin resistance and is typically characterized by prominent ectopic fat accumulation due to both the reduction in adipocyte lipid storage capacity and the associated hyperphagia induced by leptin deficiency. To ascertain whether IMCL composition is altered in lipodystrophic (LD) subjects and if such changes in lipid composition might help to elucidate the athlete’s paradox, we determined the compositional saturation index (CH<sub>2</sub>:CH<sub>3</sub> ratio) and composition-independent concentration (from CH<sub>3</sub>) of IMCLs in the SOL and tibialis anterior (TA) muscles of female insulin-resistant LD subjects as well as age- and gender-matched athletes and non-athlete controls.

## MATERIALS AND METHODS

### Participants

Sixteen female subjects with lipodystrophy were identified as part of a long-standing study of human insulin-resistant syndromes, while age- and gender-matched controls (*n* = 41) and athletes (*n* = 14) were recruited by advertisement. SOL IMCL data from five of the subjects were included in a previously published study (18). Control and athlete exclusion criteria included current smoking; drug or alcohol addiction; any current or past medical disorder or medications that could affect measurements, including supplements such as creatine; and standard MRI contraindications. Controls were recruited who exercised less than three times per week for 1 h each time, while the athletes, some of whom competed in international events, were part of a running club and regularly ran distances between 10 and 40 km. Subjects with lipodystrophy were recruited if they could perform an overnight fast without insulin or a rapid-acting insulin analogue. The



**Fig. 1.** The ratio of CH<sub>2</sub>:CH<sub>3</sub> is influenced primarily by the degree of saturation of the FAs within triglycerides. This figure shows the principle of the CH<sub>2</sub>:CH<sub>3</sub> ratio as a compositional saturation index of IMCLs. A: In this example, the palmitoleic acid component of triglyceride has a theoretical ratio of CH<sub>2</sub> (at 1.3 ppm) to CH<sub>3</sub> (at 0.9 ppm) of 16/3 = 5.3, which is lower than the equivalent ratio of palmitic acid (8.0). This is due not only to the desaturation of two CH<sub>2</sub> groups but also to the alteration in the chemical environment of the neighboring CH<sub>2</sub> groups. For FA chains with an equal number of double bonds, the CH<sub>2</sub>:CH<sub>3</sub> ratio will also be scaled by chain length, although this will have a proportionally smaller effect (17). B: Theoretical CH<sub>2</sub>:CH<sub>3</sub> values for stearic, oleic, and linoleic FAs and approximate practical values, which are systematically shifted with respect to the theoretical values, as discussed in Thankamony et al. (17). Part of this figure has been reproduced from Thankamony et al. (17), which is licensed under a Creative Commons Attribution 4.0 International License.

studies relating to lipodystrophy were approved by the NHS Research Ethics Committee, and the healthy volunteer studies were approved by the East of England Cambridge Central Ethics Committee. Studies were conducted in accordance with the Declaration of Helsinki, and all participants provided written informed consent.

### Protocol

Volunteers were instructed to follow normal dietary habits for 3 days before arriving at the National Institute for Health Research (NIHR)/Wellcome Trust Clinical Research Facility. Participants provided fasting blood samples and were given a light breakfast of either toast or cereal immediately prior to  $^1\text{H}$  MRS. Athletes and controls were instructed to refrain from vigorous exercise for at least 24 and 19 h, respectively, prior to  $^1\text{H}$  MRS.

HOMA-IR was calculated as fasting insulin (pmol/l)  $\times$  fasting glucose ( $\mu\text{U}/\text{ml}$ )/22.5. Body composition was assessed by dual-energy X-ray absorptiometry (Lunar Prodigy enCORE version 12.5 for controls and Lunar iDXA enCORE version 16 for athletes; GE Healthcare, Madison, WI).

### $^1\text{H}$ MRS

$^1\text{H}$  MRS studies were performed on a 3T scanner (Siemens; Erlangen, Germany) using the point-resolved spectroscopy sequence with a short echo time of 35 ms. A water-suppressed  $^1\text{H}$  spectrum was acquired from a voxel with a cube length of 1.3 cm positioned to avoid visible fat on  $T_1$ -weighted images within TA and SOL using a 5 s repetition time and 64 averages (4 averages for the nonwater-suppressed spectrum). Data were analyzed in jMRUI (19, 20) and fitted with the AMARES (21) algorithm using identical prior-knowledge parameters: Gaussian line shapes (except water: Lorentzian), soft constraints on EMCL/IMCL  $\text{CH}_2$  frequencies and line widths,  $\text{CH}_3$  resonant frequencies and line widths determined from known and inferred prior knowledge relative to the  $\text{CH}_2$  resonance (22), and with all amplitudes estimated. Because the  $\text{CH}_3$  resonance is small and may be subject to spectral overlap, the results were later checked for robustness by reanalyzing the data using different fitting parameters, as outlined in supplemental Table S1. IMCL  $\text{CH}_2$  and  $\text{CH}_3$  are quantified relative to the methyl group of creatine plus phosphocreatine at 3.0 ppm. Because this resonance exhibits different line-shape characteristics in the TA and SOL muscles (23), comparable quantification between muscles using a nominal concentration of muscle creatine is not valid; instead, a scaling factor of creatine to water for each muscle was established from a subset of participants who had nonwater-suppressed data sets, yielding a calculated water signal. Absolute composition-independent IMCL concentrations in mmol/kg muscle wet weight were calculated from the compositionally invariant  $\text{CH}_3$  IMCL resonance, with standard assumptions regarding muscle water content and correction for  $T_2$  relaxation effects, J coupling, and proton density as outlined below.

Absolute IMCL concentrations in mmol/kg muscle wet weight were calculated using the  $\text{CH}_3$  IMCL resonance (which is compositionally invariant) using the following equation:

$$[\text{IMCL}] = \left( S_{\text{o IMCL CH}_3} / S_{\text{o water-calc}} \right) \cdot [\text{water}]$$

where  $S_{\text{o}}$  is the corrected signal intensity of the resonance, water-calc is the calculated water signal from the internal standard (creatine and phosphocreatine), and [water] is the concentration of water in skeletal muscle [calculated using a pure water concentration of 55,342 mmol/l and assuming a relative tissue water

content in human skeletal muscle of 0.81 (kg/kg) and tissue density of 1.05 g/ml (24)]. Because the IMCL  $\text{CH}_3$  resonance is subject to J-coupling effects and has an unknown  $T_2$  relaxation time, we utilized the theoretical-to-measured IMCL  $\text{CH}_3:\text{CH}_2$  ratio that would take into effect both J-coupling and  $T_2$  effects at this echo time as well as any bias of constrained fitting prior knowledge of the  $\text{CH}_3$  resonance.

$$\left( S_{\text{o IMCL CH}_3} / S_{\text{o water-calc}} \right) = \left( S_{\text{o IMCL CH}_3} / S_{\text{o IMCL CH}_2} \right) \cdot \left( S_{\text{o IMCL CH}_2} / S_{\text{o water-calc}} \right)$$

Therefore,

$$\left( S_{\text{o IMCL CH}_3} / S_{\text{o water-calc}} \right) = \left( S_{\text{IMCL CH}_3} / S_{\text{water-calc}} \right) \cdot \left( G_{\text{TMCH}_3:\text{CH}_2} \right) \cdot \left( T_{2\text{corr CH}_2/\text{water}} \right) \cdot \left( n_{\text{water:IMCL CH}_3} \right)$$

where  $S$  is the uncorrected signal intensity of the resonance. ( $G_{\text{TM CH}_3:\text{CH}_2}$ ) = 1.1966 and is the gradient of the line of best fit through the origin of the graph of theoretical-to-measured  $\text{CH}_3:\text{CH}_2$  in vitro in IMCL- and EMCL-simulated phantoms using the point-resolved spectroscopy sequence at 3T with an echo time of 35 ms with the same fitting (17), assuming that J-coupling and  $T_2$  relaxation effects would be similar in vivo. ( $T_{2\text{corr CH}_2/\text{water}}$ ) is the correction factor for  $T_2$  effects of  $\text{CH}_2$  and water that was calculated using accepted  $T_2$  values at 3T for each muscle (25), and ( $n_{\text{water:IMCL CH}_3}$ ) is the correction for proton density.

The IMCL saturation index ( $\text{CH}_2:\text{CH}_3$ ) was calculated as IMCL  $\text{CH}_2:\text{CH}_3$ , and the IMCL saturation index adjusted for quantity ( $\text{CH}_2:\text{CH}_{3\text{adj}} = \text{CH}_2 - (m\text{CH}_3 + c)$ , where  $m$  and  $c$  are the gradient and intercept, respectively, of the regression line through the control data points of  $\text{CH}_2$  versus  $\text{CH}_3$ . Investigators were blind to the insulin-resistance status of the participants during  $^1\text{H}$  MRS analysis.

### Assessment of $\text{VO}_{2\text{max}}$

Participants underwent continuous incremental exercise testing to an 85% age-predicted maximum heart rate (controls) or volitional exhaustion (athletes) on a Trackmaster TMX425 treadmill (Med-Electronics, Beltsville, MD). LD subjects did not perform an exercise test. Oxygen consumption was measured using a spiroergometer (Medical Graphics UK Ltd, Gloucester, UK) and BreezeSuite gas-exchange software. For the control participants, a standard incremental protocol was performed (26), whereas the athletes undertook a protocol that began with a 10 min warm-up period at each participant's preferred warm-up running speed, after which the test was initiated at 9 km/h and increased steadily (0.74 km/h/min), with a ramp at 5 min (increasing 0.5% every 15 s) until exhaustion or a plateau in  $\text{VO}_2$  was apparent.  $\text{VO}_{2\text{max}}$  was calculated in the control participants by extrapolating the submaximal heart rate –  $\text{VO}_2$  relationship to the age-predicted maximum heart rate (27).

### Statistics

All statistics were performed in SPSS Statistics 24 (IBM, Armonk, NY) with significance set at  $P < 0.05$ . Normality was assessed by the Shapiro-Wilk test, and nonnormally distributed data were log-transformed prior to statistical testing. ANOVA with Games-Howell post hoc analysis was used to compare means between groups, and Pearson's correlation coefficient was used for analyzing associations. Due to the nonnormality of  $\ln(\text{HOMA-IR})$ , IMCL associations with HOMA-IR were assessed by Spearman's rank correlation coefficient. Data are presented as means  $\pm$  SEMs.

## RESULTS

### Participants

Of the insulin-resistant subjects with lipodystrophy, 13 had partial forms [8 subjects with familial partial lipodystrophy (FPLD) type 2 due to *LMNA* mutations, 5 subjects with FPLD3 due to *PPARG* mutations], and 3 had generalized lipodystrophy [2 subjects with acquired generalized lipodystrophy (AGLD) and 1 due to mutations in the *PCYT1A* gene (28)]. Of the LD subjects, two were taking no medication at all, eight were prescribed metformin, three were taking statins, five were taking fibrates, and five were taking long-acting insulin analogues. The age- and gender-matched controls had a wide BMI range (19.6–35.6 kg/m<sup>2</sup>) and HOMA-IR (0.3–4.9). As a group, insulin and HOMA-IR were significantly higher in the LD subjects and lower in the athletes (Table 1) compared with controls, as expected. Fat mass and percentage body fat were similar between LD subjects and athletes, which were both lower compared with controls (Table 1). Serum triglycerides were higher and HDL-cholesterol concentrations were lower in the LD subjects (Table 1) compared with either controls or athletes. EMCLs were absent in the two subjects with AGLD (Fig. 2), but those with partial forms of lipodystrophy had an EMCL concentration such that overall LD subjects' EMCL concentration was similar to both controls and athletes (Table 1).

### <sup>1</sup>H MRS analysis of IMCL concentration

In the SOL muscle, IMCL concentrations derived from the IMCL CH<sub>3</sub> peak (0.9 ppm) (composition-independent IMCL concentrations) were not significantly increased

( $P = 0.477$ ) in the LD subjects compared with controls but were higher compared with the lean athletes ( $P = 0.003$ ) (Fig. 3A). In the more glycolytic TA muscle, composition-independent IMCL concentrations were similar in all three groups (Fig. 3B). We also observed linear inverse correlations of VO<sub>2max</sub> and the IMCL concentration in the subset of controls who underwent VO<sub>2max</sub> testing and athletes together (Fig. 3C, D). SOL IMCL concentration was significantly lower in the athletes compared with the controls ( $P = 0.004$ ; Fig. 3A), and this remained significant ( $P = 0.025$ ) compared with a subset of the controls matched for percentage body fat (controls: body fat = 21.5 ± 2.3%,  $n = 10$ ; athletes: 21.1 ± 1.8%,  $n = 14$ ; see supplemental Fig. S1).

The conventional estimate of IMCL concentration using the CH<sub>2</sub> resonance uncorrected for composition, CH<sub>2</sub>/water, showed similar trends to the composition-independent estimate using the CH<sub>3</sub> peak with the exception that the LD subjects' SOL IMCL CH<sub>2</sub> was significantly increased compared with controls (Table 1). This could be regarded as an artifact of the effects of compositional differences, which we consider next.

### <sup>1</sup>H MRS analysis of IMCL composition

IMCLs had a significantly higher saturation index (CH<sub>2</sub>:CH<sub>3</sub>) in both muscles of the LD subjects compared with controls (SOL  $P = 0.008$ ; TA  $P = 0.024$ ) but not athletes (Fig. 4A, B). In the control group, smaller IMCL pools were associated with a higher saturation index, as shown by the linear regression line (dotted line in Fig. 3E, F) having a gradient ( $\Delta\text{CH}_2/\Delta\text{CH}_3$ ) that was less than the mean CH<sub>2</sub>:CH<sub>3</sub> (e.g., gradient SOL = 6.7 vs. mean CH<sub>2</sub>:CH<sub>3</sub> = 8.8;

TABLE 1. Participant characteristics and conventional muscle lipid estimates

	LD Females ( $n = 16$ )	Female Controls ( $n = 41$ )	Female Athletes ( $n = 14$ )	<i>P</i>			
				ANOVA	LD—Controls	LD—Athletes	Controls—Athletes
Participant characteristics							
Age (years)	38.9 ± 3.9	35.5 ± 2.0	36.5 ± 2.9	0.694			
BMI (kg/m <sup>2</sup> )	24.2 ± 0.7	24.4 ± 0.6	20.3 ± 0.6	<0.001	0.999	0.001	<0.001
Mass (kg)	66.1 ± 2.8	65.2 ± 2.3	54.6 ± 1.3	0.013	0.916	0.006	0.001
Fat mass (kg)	10.6 ± 1.4	23.1 ± 1.6	11.6 ± 1.1	<0.001	0.001	0.631	<0.001
Fat-free mass (kg)	55.4 ± 1.7	42.1 ± 0.9	43.0 ± 1.2	<0.001	<0.001	<0.001	0.705
Body fat (%)	16.1 ± 1.7	34.2 ± 1.2	21.1 ± 1.8	<0.001	<0.001	0.129	<0.001
Triglycerides (mmol/l)	3.54 ± 0.67	0.92 ± 0.07 <sup>a</sup>	0.84 ± 0.08	<0.001	<0.001	<0.001	0.920
HDL-cholesterol (mmol/l)	0.97 ± 0.71	1.64 ± 0.06 <sup>a</sup>	2.20 ± 0.12	<0.001	<0.001	<0.001	0.001
Glucose (mmol/l)	6.00 ± 0.58	4.56 ± 0.06 <sup>a</sup>	4.46 ± 0.13	<0.001	0.063	0.002	0.764
Insulin (pmol/l) <sup>b</sup>	145.6 ± 22.0	38.6 ± 4.2 <sup>a</sup>	22.6 ± 4.7	<0.001	<0.001	<0.001	0.025
HOMA-IR	5.53 ± 1.0	1.14 ± 0.13 <sup>a</sup>	0.66 ± 0.15	<0.001	<0.001	<0.001	0.025
HbA1c (%)	6.6 ± 0.4 <sup>c</sup>	ND	5.3 ± 0.1	ND		0.008	
VO <sub>2max</sub> (ml/kg/min)	ND	36.1 ± 1.5 <sup>d</sup>	46.9 ± 1.4	ND			<0.001
Conventional estimates <sup>e</sup>							
SOL IMCLs	1.90 ± 0.21	1.22 ± 0.07	0.79 ± 0.10	<0.001	0.027	<0.001	0.007
TA IMCLs	0.89 ± 0.16 <sup>f</sup>	0.61 ± 0.04	0.45 ± 0.04	0.035	0.562	0.115	0.078
SOL EMCLs	2.05 ± 0.34	2.22 ± 0.19	1.81 ± 0.20	0.493			
TA EMCLs	1.43 ± 0.28 <sup>f</sup>	2.30 ± 0.19	1.24 ± 0.15	0.002	0.163	0.785	0.005

Data are presented as mean ± SEM unless otherwise stated. Nonnormally distributed variables were log-transformed prior to performing ANOVA and Games-Howell post hoc analysis; bold *P* values are statistically significant.

<sup>a</sup> $n = 38$ .

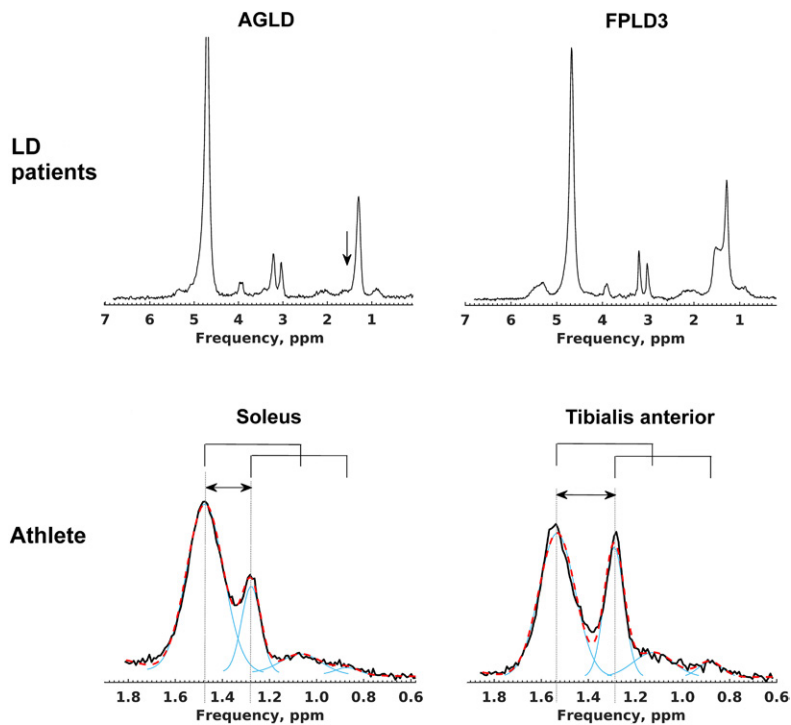
<sup>b</sup>To convert to  $\mu\text{U}/\text{ml}$  divide by 6.945.

<sup>c</sup> $n = 13$ .

<sup>d</sup> $n = 24$ .

<sup>e</sup>Expressed as methylene protons resonating at 1.3 ppm quantified as a percentage of the uncorrected calculated water resonance.

<sup>f</sup> $n = 12$ .



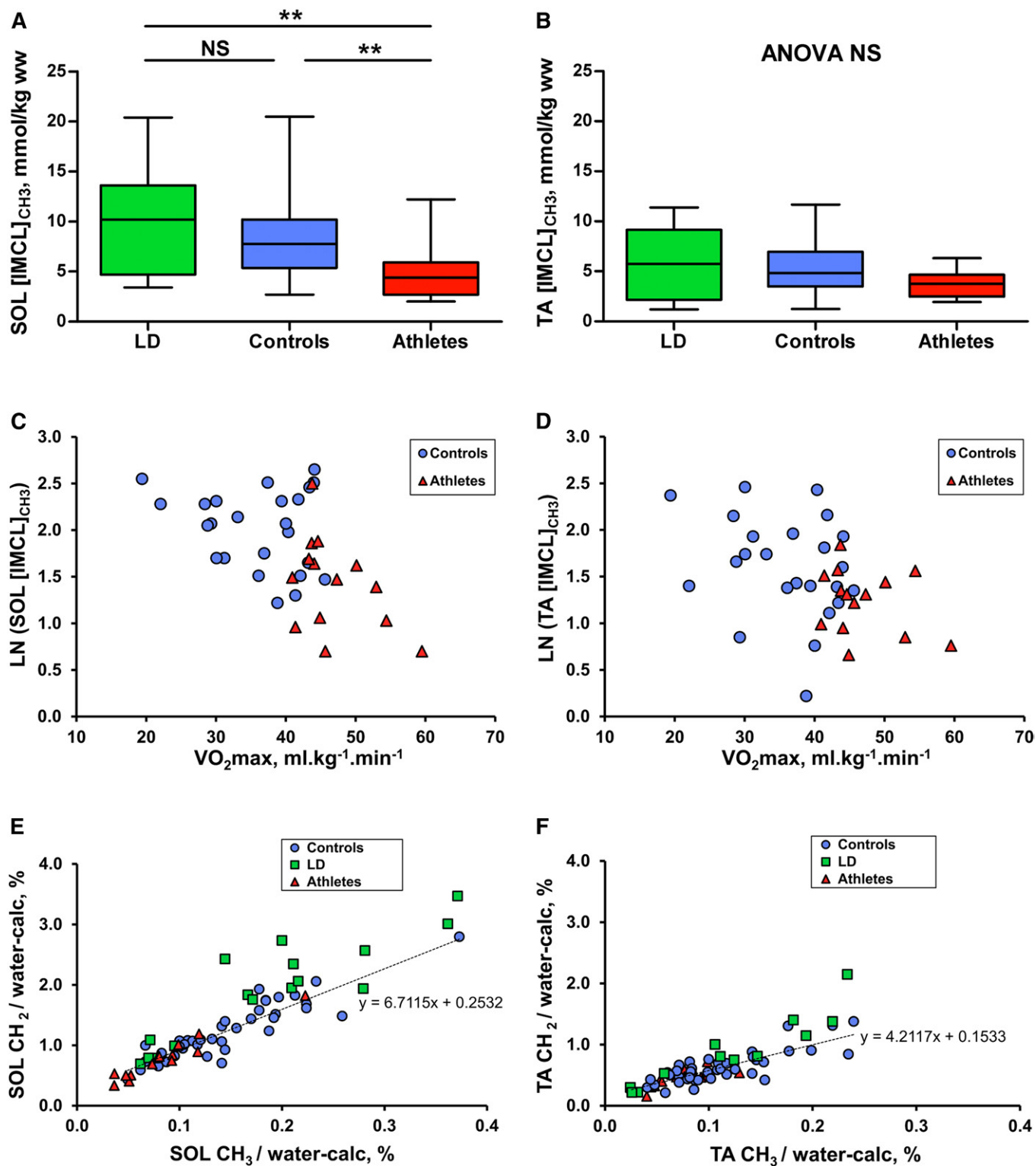
**Fig. 2.** Representative  $^1\text{H}$  MRS spectra from LD subjects and an athlete. Water-suppressed spectra from the SOL muscle of a subject with AGLD (upper left) and FPLD3 (upper right); the absence of EMCLs in AGLD is highlighted by the vertical arrow. Spectra from an athlete's SOL (lower left) and TA (lower right) muscles illustrate the raw data (solid black) and overall fit (red dashed) and individual fit components (solid blue) in the frequency range that contains the EMCL  $\text{CH}_2$  ( $\sim 1.5$  ppm),  $\text{CH}_3$  ( $\sim 1.1$  ppm), IMCL  $\text{CH}_2$  (1.3 ppm), and  $\text{CH}_3$  (0.9 ppm) resonances. The horizontal arrows indicate that the EMCL resonances are systematically shifted very slightly upfield in the SOL muscle compared with the TA due to fiber-orientation effects. The  $\text{CH}_3$  resonant frequencies are linked to the  $\text{CH}_2$  frequencies (solid bridge lines above spectra) and are also shifted. The fitting procedure fixes the relative  $\text{CH}_2$  to the  $\text{CH}_3$  frequency shift for both EMCLs and IMCLs and the  $\text{CH}_3$  line width relative to the corresponding  $\text{CH}_2$  line width but permits soft constraints on the  $\text{CH}_2$  frequencies.

gradient TA = 4.2 vs. mean  $\text{CH}_2:\text{CH}_3 = 6.0$ ). This phenomenon seemed to be independent of insulin sensitivity in the controls (**Table 2**; there was no relation of HOMA-IR with IMCL concentration). To generate a pathophysiologically meaningful measure of composition that is independent of IMCL quantity, the vertical ( $\text{CH}_2$ ) deviation from this regression line was measured and taken as a marker of the saturation of the pool that is adjusted for quantity, which we term the adjusted saturation index ( $\text{CH}_2:\text{CH}_{3\text{adj}}$ ). This adjusted compositional marker was significantly higher in LD subjects compared with athletes (SOL  $P = 0.001$ ; TA  $P = 0.046$ ) and controls (SOL  $P = 0.003$ ), with a tendency in the TA that falls just short of conventional statistical significance ( $P = 0.06$ ) (Fig. 4C, D).

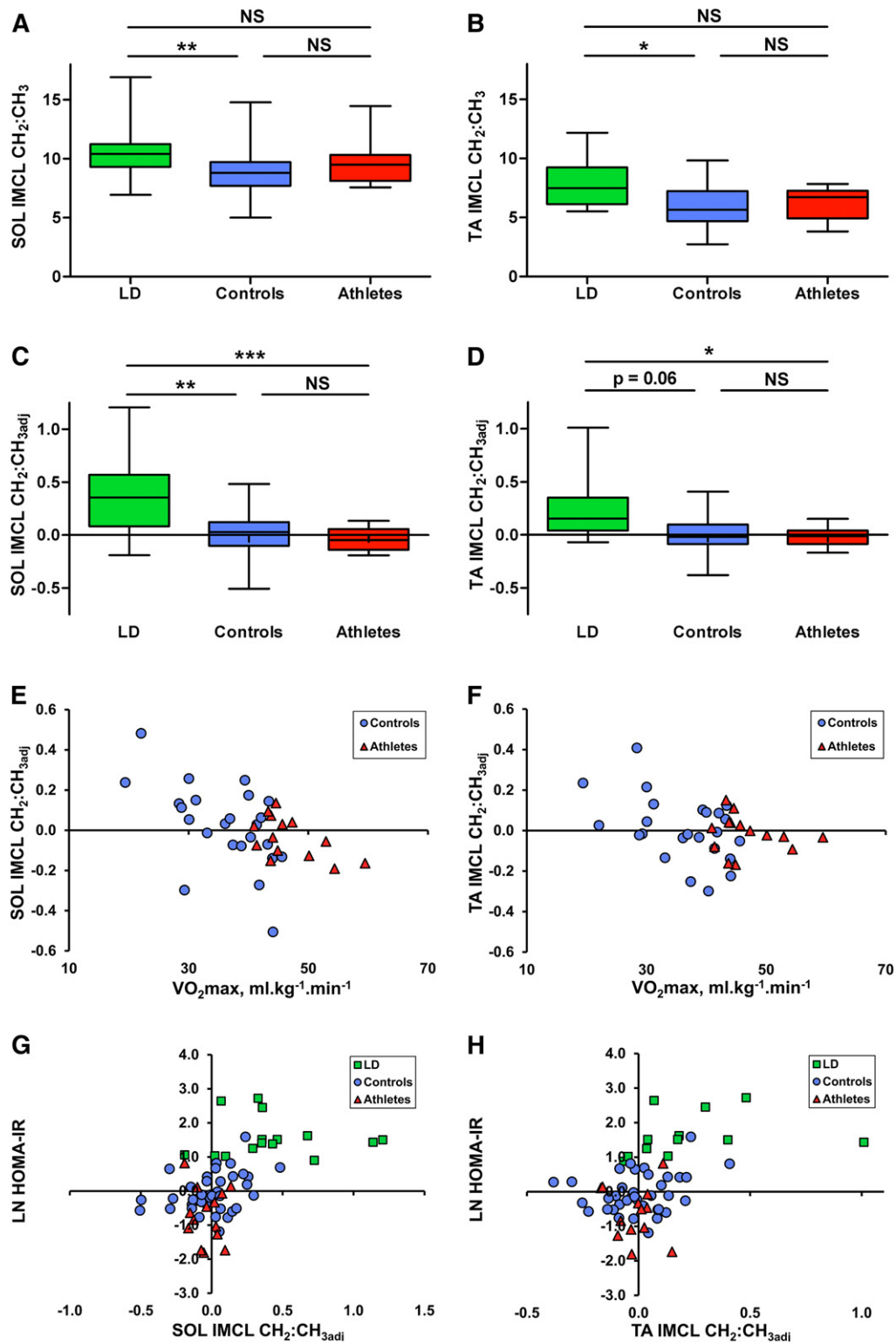
Unlike the uncorrected measure of composition, this adjusted composition also had a significant relation to  $\text{VO}_{2\text{max}}$  (Fig. 4E, F) within the control subset alone, athletes alone (SOL), and control and athletes combined, such that fitter individuals had less saturated IMCLs for the same absolute quantity of IMCLs.  $\text{VO}_{2\text{max}}$  was significantly correlated with HOMA-IR in the control subset ( $r = -0.59$ ,  $P = 0.003$ ,  $n = 23$ ) and in controls and athletes together ( $r = -0.53$ ,  $P = 0.001$ ,  $n = 37$ ). Figure 4G and H show the relation of the adjusted composition to HOMA-IR. Table 2 shows relations of IMCL concentration and composition with insulin sensitivity. The saturation index was higher in the SOL muscle compared with the TA in all three groups. This was still the case in the two EMCL-deficient AGLD subjects (subject 1: SOL  $\text{CH}_2:\text{CH}_3 = 9.5$  and TA  $\text{CH}_2:\text{CH}_3 = 6.3$ ; subject 2: SOL  $\text{CH}_2:\text{CH}_3 = 11.2$  and TA  $\text{CH}_2:\text{CH}_3 = 7.1$ ). The IMCL relations with HOMA-IR were robust to differing fitting parameters, as shown in supplemental Table S1.

## DISCUSSION

Using a recently validated  $^1\text{H}$  MRS method we compared a compositional saturation index ( $\text{CH}_2:\text{CH}_3$  ratio) of IMCLs in the SOL and TA muscles of female insulin-resistant LD subjects with that of age- and gender-matched athletes and healthy controls and showed it to be significantly higher in both muscles compared with controls but not athletes. The finding that smaller IMCL pools in the control group had a relatively higher saturation index than larger pools irrespective of insulin sensitivity could possibly explain why the athletes studied here, who had small IMCL pools, had a composition that was statistically similar to LD subjects. This observed concentration-composition relationship seems physiologically plausible given that more unsaturated and shorter-chain FAs are preferentially mobilized (17). A similar trend was also visible in a previous data set from both TA and SOL muscles of 19 healthy males after an 8 h fast (17). We hypothesize that a person may “move” along a line such as this while performing daily activities and that a measurement of deviation from this relationship may therefore be a more sensitive and specific measure of muscle metabolic physiology or pathophysiology. To take this concentration-composition dependence into consideration we adjusted the compositional saturation index for concentration ( $\text{CH}_2:\text{CH}_{3\text{adj}}$ ), and this marker was able to distinguish between athletes and LD subjects in both muscles. The strong inverse relation of  $\text{CH}_2:\text{CH}_{3\text{adj}}$  with  $\text{VO}_{2\text{max}}$  indicates that fitness is associated with a relatively lower saturation of IMCLs, although overall athletes and controls were not statistically different; this is similar to results from a biopsy study (29) that demonstrated a comparable percentage of saturated intramuscular triglycerides in male controls and athletes. Interestingly, previous



**Fig. 3.** Composition-independent concentrations of IMCLs in LD subjects (green), controls (blue), and athletes (red). **A, B:** Box and whisker plots showing SOL and TA composition-independent IMCL concentration from the <sup>1</sup>H MRS of methyl protons, as assessed by ANOVA and Games-Howell post hoc analyses. **C, D:** The relationship of SOL and TA composition-independent IMCL concentration with VO<sub>2</sub>max in a subset of participants who underwent VO<sub>2</sub>max testing, as assessed by Pearson's correlation coefficient (controls: blue circles, *n* = 24; athletes: red triangles, *n* = 14). This was only significant when controls and athletes were combined (SOL: *r* = -0.52, *P* = 0.001; TA: *r* = -0.42, *P* = 0.009). **E, F:** SOL and TA IMCL CH<sub>2</sub> and CH<sub>3</sub> components. The values are expressed relative to the calculated water signal (water-calc), as described in Materials and Methods. The dotted line represents the linear regression line of the control data points. \**P* < 0.05, \*\**P* < 0.01, and \*\*\**P* < 0.001.



**Fig. 4.** <sup>1</sup>H MRS measures of IMCL composition in LD subjects (green), controls (blue), and athletes (red). A, B: Box and whisker plots of (A) SOL and (B) TA IMCL compositional saturation index (CH<sub>2</sub>:CH<sub>3</sub> ratio), as assessed by ANOVA and Games-Howell post hoc analyses. C, D: SOL and TA IMCL compositional saturation index adjusted for quantity. E, F: Relation of SOL and TA IMCL compositional adjusted saturation index with VO<sub>2max</sub> in the subset of participants who underwent VO<sub>2max</sub> testing, as assessed by Pearson's correlation coefficient (controls: blue circles, *n* = 24; athletes: red triangles, *n* = 14). There was a significant correlation in the SOL (*r* = -0.546, *P* = 0.006) and TA (*r* = -0.453, *P* = 0.026) for controls alone, in the SOL (*r* = -0.558, *P* = 0.038) for athletes alone, and in the SOL (*r* = -0.520, *P* = 0.001) and TA (*r* = -0.362, *P* = 0.025) for controls and athletes combined. G, H: Relation of HOMA-IR with SOL and TA IMCL compositional adjusted saturation index. Correlation coefficients are shown in Table 2. \**P* < 0.05, \*\**P* < 0.01, and \*\*\**P* < 0.001.

TABLE 2. Correlation coefficients of whole-body insulin resistance with IMCLs

IMCL Measure	HOMA-IR			
	Controls ( <i>n</i> = 38)	Controls and Athletes ( <i>n</i> = 52)	Controls and LD Subjects ( <i>n</i> = 54) <sup>a</sup>	Controls, LD Subjects, and Athletes ( <i>n</i> = 68) <sup>b</sup>
<b>SOL</b>				
Concentration (CH <sub>3</sub> )	-0.016	0.248	0.149	0.312*
Concentration and composition (CH <sub>2</sub> )	0.153	0.315*	0.394**	0.477***
Composition (CH <sub>2</sub> :CH <sub>3</sub> )	0.217	-0.048	0.439***	0.241*
Composition adjusted for quantity (CH <sub>2</sub> :CH <sub>3adj</sub> )	0.320*	0.271*	0.583***	0.532***
<b>TA</b>				
Concentration (CH <sub>3</sub> )	0.200	0.251	0.205	0.258*
Concentration and composition (CH <sub>2</sub> )	0.224	0.247	0.338*	0.344**
Composition (CH <sub>2</sub> :CH <sub>3</sub> )	0.100	-0.046	0.337*	0.202
Composition adjusted for quantity (CH <sub>2</sub> :CH <sub>3adj</sub> )	0.218	0.127	0.445***	0.364**

\**P* < 0.05, \*\**P* < 0.01, and \*\*\**P* ≤ 0.001.

<sup>a</sup>*n* = 50 for TA.

<sup>b</sup>*n* = 64 for TA.

reports of intramuscular triglyceride composition in insulin-resistant states (30, 31) yielded no difference in saturation percentage but did reveal a difference in linoleate (31). In these studies, the intramuscular triglyceride content was higher in insulin-resistant states; therefore, a higher saturation percentage may not be apparent given the composition-concentration relation we found. This lack of a difference in saturation could also equally be explained by the difference in sampling intramuscular versus intramyocellular pools, as EMCLs generally make up a significant proportion of intramuscular triglycerides (approximately twice the IMCL pool in our healthy cohorts even when voxels were placed to avoid visible marbling on T<sub>1</sub>-weighted images; Table 1).

By our measure of composition-independent IMCLs (using the CH<sub>3</sub> instead of the uncorrected CH<sub>2</sub> resonance), we found that the LD subjects' IMCL concentration was not significantly higher in either muscle compared with age-, gender-, and BMI-matched controls but was higher in the SOL relative to athletes. There are few reports of IMCLs in LD subjects, but our findings are in agreement with Peterson et al. (32), who in three subjects with generalized lipodystrophy (two congenital and one acquired) found SOL IMCL concentrations that were similar to six age-, BMI-, and weight-matched controls. Calf IMCL concentration was also lower in a case report of acquired generalized lipodystrophy (33) compared with controls, while a study of four subjects with congenital generalized lipodystrophy suggested that IMCL concentration was higher (24). Using the CH<sub>2</sub> resonance, as has been conventional in previous studies, we found SOL IMCL content to be significantly higher in our LD subjects compared with controls (Table 1), demonstrating, we argue, the influence of composition on measures of concentration.

### The athlete's paradox

We found that the athletes' IMCL concentration was significantly lower (SOL) or similar (TA) to controls. Although this seems to contradict well-known reports of an athlete's paradox using both biopsy methods (13, 29, 34–36) and <sup>1</sup>H MRS (37, 38), this finding is in agreement with

studies that reported no such paradox compared with old or young controls (39), obese individuals (40), or in certain fiber types (14). It is known that athletes can have a large depletion-repletion range of IMCLs, and it is possible that the IMCL concentration had not fully recovered since the last training session (24–48 h prior), as IMCLs can still increase significantly after these intervals (41). In fact, our results demonstrate an inverse relation of IMCL content and VO<sub>2max</sub> that is consistent with a study by Boesch et al. (42), in which a combination of daily training at 60% of the VO<sub>2</sub> peak with a low-fat (10% to 15% fat) diet depleted IMCL levels in both the vastus lateralis and TA muscles to a consistent level that correlated with the VO<sub>2</sub> peak, suggesting that our elite female athletes were nearly "empty" of IMCLs; we did not control for diet in our study.

### Relationship of whole-body insulin resistance to IMCLs

Within the controls, only the compositional saturation index adjusted for quantity (CH<sub>2</sub>:CH<sub>3adj</sub>) in the SOL was significantly correlated with whole-body insulin resistance (Table 2). The inclusion of insulin-resistant LD subjects increases the statistical significance of this relation and yields associations with other composition-influenced markers, but not IMCL concentration, in either muscle. In our study, the addition of athletes predictively reduced the associations with composition, as their pools were small and therefore had a tendency for a higher saturation index, but relations remained with measures that reflect large saturated pools (i.e., CH<sub>2</sub>:CH<sub>3adj</sub> and CH<sub>2</sub>). These striking results suggest that the accumulation of saturated IMCLs and not concentration alone relates to early-stage insulin resistance.

Supporting our findings, the concentration of the most abundant saturated fat, palmitic acid, within muscle triglycerides has been shown to be related to insulin sensitivity (43). Palmitic acid is known to increase ceramide concentrations, which are thought to engage stress-responsive serine kinases that impede insulin activation of its cell-surface receptor, as well as downstream signaling molecules such as insulin receptor substrate 1 and protein kinase B/Akt (44). Our study was of course not able to probe these mechanisms.




## Limitations

Unlike previous  $^1\text{H}$  MRS studies that have conventionally reported IMCL concentrations using the predominant  $\text{CH}_2$  resonance while assuming a notional normal composition, here we utilized the smaller  $\text{CH}_3$  resonance, which has the advantage of composition independence and, with comprehensive prior-knowledge constraints, including line-width constraints relative to the  $\text{CH}_2$  resonance (22), fitted the IMCL  $\text{CH}_3$  resonance from the overlapping EMCL/IMCL  $\text{CH}_3$  signals. Due to fiber-orientation differences between the SOL and TA muscles, the EMCL resonances are very slightly systematically shifted between muscle groups. Despite this potential for a systematic difference in the  $\text{CH}_2$ : $\text{CH}_3$  ratio between muscles, this ratio was still higher in the SOL compared with the TA muscle in both EMCL-deficient AGLD subjects, consistent with findings in our other participants in this study and a previous study (17) suggesting a compositional difference between muscles. Our LD subjects, who mainly had partial forms of lipodystrophy, had overall similar quantities of EMCLs to those of our controls and athletes that has helped to reduce potential intergroup influence of this overlapping resonance. In addition, the relations of IMCLs with HOMA-IR were robust to differing fitting routines, including constraining the EMCL  $\text{CH}_3$  amplitude and accounting for asymmetric line shapes. These findings, together with the finding that neither the IMCL  $\text{CH}_2$ : $\text{CH}_3$  nor IMCL  $\text{CH}_2$ : $\text{CH}_{3\text{adj}}$  markers related to either EMCL  $\text{CH}_2$  or  $\text{CH}_3$ , suggests a lack of EMCL influence in our data sets; however, it is possible that in other insulin-resistant cohorts large overlapping EMCL resonances may be a confounding factor.

## Summary

The use of our recently validated and potentially widely applicable  $^1\text{H}$  MRS approach to determine both the IMCL composition and concentration independent of composition within the SOL and TA muscles of female individuals covering a wide range of insulin sensitivities has revealed that markers of the accumulation of saturated triglycerides in the IMCL pool are more strongly associated with whole-body insulin resistance than IMCL concentration alone. Differences in associations of insulin resistance with IMCL concentration when using the  $\text{CH}_3$  and conventional  $\text{CH}_2$  peaks for quantification highlights the need for awareness of the potential influence of composition on previously reported  $^1\text{H}$  MRS measures of concentration.

Our finding of a strong relationship between  $\text{VO}_{2\text{max}}$  and relatively unsaturated IMCL pools in controls and athletes points to a role of exercise in decreasing the amount of saturated fat within the IMCL store. The association of insulin resistance with the accumulation of saturated IMCLs, even within a healthy control population, could suggest an early involvement in its pathogenesis and provide a reason why combined exercise and diet are effective therapeutic options in the early stages of insulin resistance. 

The authors thank the participants and staff at both the NIHR/Wellcome Trust Cambridge Clinical Research Facility and the Wolfson Brain Imaging Centre. The authors also thank the

NIHR Core Biochemistry Assay Laboratory, Cambridge Biomedical Research Centre, for providing the insulin analysis and Wiktor Olszowy for statistical advice.

## REFERENCES

- Schick, F., B. Eismann, W-I. Jung, H. Bongers, M. Bunse, and O. Lutz. 1993. Comparison of localized proton NMR signals of skeletal muscle and fat tissue in vivo: two lipid compartments in muscle tissue. *Magn. Reson. Med.* **29**: 158–167.
- Boesch, C., J. Slotboom, H. Hoppeler, and R. Kreis. 1997. In vivo determination of intra-myocellular lipids in human muscle by means of localized  $^1\text{H}$ -MR-spectroscopy. *Magn. Reson. Med.* **37**: 484–493.
- Perseghin, G., P. Scifo, F. De Cobelli, E. Pagliato, A. Battezzati, C. Arcelloni, A. Vanzulli, G. Testolin, G. Pozza, A. Del Maschio, et al. 1999. Intramyocellular triglyceride content is a determinant of in vivo insulin resistance in humans: a  $^1\text{H}$ - $^{13}\text{C}$  nuclear magnetic resonance spectroscopy assessment in offspring of type 2 diabetic parents. *Diabetes.* **48**: 1600–1606.
- Jacob, S., J. Machann, K. Rett, K. Brechtel, A. Volk, W. Renn, E. Maerker, S. Matthaei, F. Schick, C. D. Claussen, et al. 1999. Association of increased intramyocellular lipid content with insulin resistance in lean nondiabetic offspring of type 2 diabetic subjects. *Diabetes.* **48**: 1113–1119.
- Krssak, M., K. Falk Petersen, A. Dresner, L. DiPietro, S. M. Vogel, D. L. Rothman, G. I. Shulman, and M. Roden. 1999. Intramyocellular lipid concentrations are correlated with insulin sensitivity in humans: a  $^1\text{H}$  NMR spectroscopy study. *Diabetologia.* **42**: 113–116.
- Shulman, G. I., D. L. Rothman, T. Jue, P. Stein, R. A. DeFronzo, and R. G. Shulman. 1990. Quantitation of muscle glycogen synthesis in normal subjects and subjects with non-insulin-dependent diabetes by  $^{13}\text{C}$  nuclear magnetic resonance spectroscopy. *N. Engl. J. Med.* **322**: 223–228.
- Petersen, K. F., and G. I. Shulman. 2002. Pathogenesis of skeletal muscle insulin resistance in type 2 diabetes mellitus. *Am. J. Cardiol.* **90**: 11–18.
- Savage, D. B., K. F. Petersen, and G. I. Shulman. 2005. Mechanisms of insulin resistance in humans and possible links with inflammation. *Hypertension.* **45**: 828–833.
- Sinha, R., S. Dufour, K. F. Petersen, V. LeBon, S. Enoksson, Y-Z. Ma, M. Savoye, D. L. Rothman, G. I. Shulman, and S. Caprio. 2002. Assessment of skeletal muscle triglyceride content by  $^1\text{H}$  nuclear magnetic resonance spectroscopy in lean and obese adolescents: relationships to insulin sensitivity, total body fat, and central adiposity. *Diabetes.* **51**: 1022–1027.
- Phillips, D. I. W., S. Caddy, V. Ilic, B. A. Fielding, K. N. Frayn, A. C. Borthwick, and R. Taylor. 1996. Intramuscular triglyceride and muscle insulin sensitivity: evidence for a relationship in nondiabetic subjects. *Metabolism.* **45**: 947–950.
- Pan, D. A., S. Lillioja, A. D. Kriketos, M. R. Milner, L. A. Baur, C. Bogardus, A. B. Jenkins, and L. H. Storlien. 1997. Skeletal muscle triglyceride levels are inversely related to insulin action. *Diabetes.* **46**: 983–988.
- Forouhi, N. G., G. Jenkinson, E. L. Thomas, S. Mullick, S. Mierisova, U. Bhonsle, P. M. McKeigue, and J. D. Bell. 1999. Relation of triglyceride stores in skeletal muscle cells to central obesity and insulin sensitivity in European and South Asian men. *Diabetologia.* **42**: 932–935.
- Goodpaster, B. H., J. He, S. Watkins, and D. E. Kelley. 2001. Skeletal muscle lipid content and insulin resistance: evidence for a paradox in endurance-trained athletes. *J. Clin. Endocrinol. Metab.* **86**: 5755–5761.
- van Loon, L. J. C., R. Koopman, R. Manders, W. van der Weegen, G. P. van Kranenburg, and H. A. Keizer. 2004. Intramyocellular lipid content in type 2 diabetes patients compared with overweight sedentary men and highly trained endurance athletes. *Am. J. Physiol. Endocrinol. Metab.* **287**: E558–E565.
- Hernández, E. Á., S. Kahl, A. Seelig, P. Begovatz, M. Irmeler, Y. Kupriyanova, B. Nowotny, P. Nowotny, C. Herder, C. Barosa, et al. 2017. Acute dietary fat intake initiates alterations in energy metabolism and insulin resistance. *J. Clin. Invest.* **127**: 695–708.
- Luukkonen, P. K., S. Sädevirta, Y. Zhou, B. Kayser, A. Ali, L. Ahonen, S. Lallukka, V. Pelloux, M. Gaggini, C. Jian, et al. 2018. Saturated fat

is more metabolically harmful for the human liver than unsaturated fat or simple sugars. *Diabetes Care*. **41**: 1732–1739.

17. Thankamony, A., G. J. Kemp, A. Koulman, V. Bokii, D. B. Savage, C. Boesch, L. Hodson, D. B. Dunger, and A. Sleight. 2018. Compositional marker in vivo reveals intramyocellular lipid turnover during fasting-induced lipolysis. *Sci. Rep.* **8**: 2750.
18. Sleight, A., A. Stears, K. Thackray, L. Watson, A. Gambineri, S. Nag, V. I. Campi, N. Schoenmakers, S. Brage, T. A. Carpenter, et al. 2012. Mitochondrial oxidative phosphorylation is impaired in patients with congenital lipodystrophy. *J. Clin. Endocrinol. Metab.* **97**: E438–E442.
19. Naressi, A., C. Couturier, J. M. Devos, M. Janssen, C. Mangeat, R. de Beer, and D. Graveron-Demilly. 2001. Java-based graphical user interface for the MRUI quantitation package. *MAGMA*. **12**: 141–152.
20. Stefan, D., F. Di Cesare, A. Andrasescu, E. Popa, A. Lazariev, E. Vescovo, O. Strbak, S. Williams, Z. Starcuk, M. Cabanas, et al. 2009. Quantitation of magnetic resonance spectroscopy signals: the jMRUI software package. *Meas. Sci. Technol.* **20**: 104035.
21. Vanhamme, L., A. Van Den Boogaart, and S. Van Huffel. 1997. Improved method for accurate and efficient quantification of MRS data with use of prior knowledge. *J. Magn. Reson.* **129**: 35–43.
22. Boesch, C., J. Machann, P. Vermathen, and F. Schick. 2006. Role of proton MR for the study of muscle lipid metabolism. *NMR Biomed.* **19**: 968–988.
23. Vermathen, P., C. Boesch, and R. Kreis. 2003. Mapping fiber orientation in human muscle by proton MR spectroscopic imaging. *Magn. Reson. Med.* **49**: 424–432.
24. Szczepaniak, L. S., E. E. Babcock, F. Schick, R. L. Dobbins, A. Garg, D. K. Burns, J. Denis McGarry, D. T. Stein, and J. Denis McGarry. 1999. Measurement of intracellular triglyceride stores by <sup>1</sup>H spectroscopy: validation in vivo. *Am. J. Physiol. Endocrinol. Metab.* **276**: E977–E989.
25. Krssák, M., M. Roden, V. Mlynárik, M. Meyerspeer, and E. Moser. 2004. <sup>1</sup>H NMR relaxation times of skeletal muscle metabolites at 3 T. *MAGMA*. **16**: 155–159.
26. Brage, S., N. Brage, U. Ekelund, J. Luan, P. W. Franks, K. Froberg, and N. J. Wareham. 2006. Effect of combined movement and heart rate monitor placement on physical activity estimates during treadmill locomotion and free-living. *Eur. J. Appl. Physiol.* **96**: 517–524.
27. Tanaka, H., K. D. Monahan, and D. R. Seals. 2001. Age-predicted maximal heart rate revisited. *J. Am. Coll. Cardiol.* **37**: 153–156.
28. Payne, F., K. Lim, A. Gironde, R. J. Brown, N. Kory, A. Robbins, Y. Xue, A. Sleight, E. Cochran, C. Adams, et al. 2014. Mutations disrupting the Kennedy phosphatidylcholine pathway in humans with congenital lipodystrophy and fatty liver disease. *Proc. Natl. Acad. Sci. USA*. **111**: 8901–8906.
29. Bergman, B. C., L. Perreault, D. M. Hunerdosse, M. C. Koehler, A. M. Samek, and R. H. Eckel. 2010. Increased intramuscular lipid synthesis and low saturation relate to insulin sensitivity in endurance-trained athletes. *J. Appl. Physiol.* **108**: 1134–1141.
30. van Hees, A. M. J., A. Jans, G. B. Hul, H. M. Roche, W. H. M. Saris, and E. E. Blaak. 2011. Skeletal muscle fatty acid handling in insulin resistant men. *Obesity (Silver Spring)*. **19**: 1350–1359.
31. Perreault, L., B. C. Bergman, D. M. Hunerdosse, and R. H. Eckel. 2010. Altered intramuscular lipid metabolism relates to diminished insulin action in men, but not women, in progression to diabetes. *Obesity (Silver Spring)*. **18**: 2093–2100.
32. Petersen, K. F., E. A. Oral, S. Dufour, D. Befroy, C. Ariyan, C. Yu, G. W. Cline, A. M. DePaoli, S. I. Taylor, P. Gorden, et al. 2002. Leptin reverses insulin resistance and hepatic steatosis in patients with severe lipodystrophy. *J. Clin. Invest.* **109**: 1345–1350.
33. Brechtel, K., S. Jacob, J. Machann, B. Hauer, M. Nielsen, H. P. Meissner, S. Matthaei, H. U. Haering, C. D. Claussen, and F. Schick. 2000. Acquired generalized lipotrophy (AGL): highly selective MR lipid imaging and localized <sup>1</sup>H-MRS. *J. Magn. Reson. Imaging*. **12**: 306–310.
34. Gemmink, A., S. Daemen, B. Brouwers, P. R. Huntjens, G. Schaart, E. Moonen-Kornips, J. Jörgensen, J. Hoeks, P. Schrauwen, and M. K. C. Hesselink. 2018. Dissociation of intramyocellular lipid storage and insulin resistance in trained athletes and type 2 diabetes patients; involvement of perilipin 5? *J. Physiol.* **596**: 857–868.
35. Amati, F., J. J. Dubé, E. Alvarez-Carnero, M. M. Edreira, P. Chomentowski, P. M. Coen, G. E. Switzer, P. E. Bickel, M. Stefanovic-Racic, F. G. S. Toledo, et al. 2011. Skeletal muscle triglycerides, diacylglycerols, and ceramides in insulin resistance: another paradox in endurance-trained athletes? *Diabetes*. **60**: 2588–2597.
36. Russell, A. P., G. Gastaldi, E. Bobbioni-Harsch, P. Arboit, C. Gobelet, O. Dériaz, A. Golay, J. L. Witztum, and J-P. Giacobino. 2003. Lipid peroxidation in skeletal muscle of obese as compared to endurance-trained humans: a case of good vs. bad lipids? *FEBS Lett.* **551**: 104–106.
37. Décombaz, J., B. Schmitt, M. Ith, B. Decarli, P. Diem, R. Kreis, H. Hoppeler, and C. Boesch. 2001. Postexercise fat intake repletes intramyocellular lipids but no faster in trained than in sedentary subjects. *Am. J. Physiol. Regul. Integr. Comp. Physiol.* **281**: R760–R769.
38. Klepochová, R., L. Valkovič, T. Hochwartner, C. Triska, N. Bachl, H. Tschan, S. Trattng, M. Krebs, and M. Krššák. 2018. Differences in muscle metabolism between triathletes and normally active volunteers investigated using multinuclear magnetic resonance spectroscopy at 7T. *Front. Physiol.* **9**: 300.
39. Bruce, C. R., M. J. Anderson, A. L. Carey, D. G. Newman, A. Bonen, A. D. Kriketos, G. J. Cooney, and J. A. Hawley. 2003. Muscle oxidative capacity is a better predictor of insulin sensitivity than lipid status. *J. Clin. Endocrinol. Metab.* **88**: 5444–5451.
40. Bergman, B. C., L. Perreault, A. Strauss, S. Bacon, A. Kerege, K. Harrison, J. T. Brozinick, D. M. Hunerdosse, M. C. Playdon, W. Holmes, et al. 2018. Intramuscular triglyceride synthesis: importance in muscle lipid partitioning in humans. *Am. J. Physiol. Endocrinol. Metab.* **314**: E152–E164.
41. Howald, H., C. Boesch, R. Kreis, S. Matter, R. Billeter, B. Essen-Gustavsson, and H. Hoppeler. 2002. Content of intramyocellular lipids derived by electron microscopy, biochemical assays, and <sup>1</sup>H-MR spectroscopy. *J. Appl. Physiol.* **92**: 2264–2272.
42. Ith, M., P. M. Huber, A. Egger, J-P. Schmid, R. Kreis, E. Christ, and C. Boesch. 2010. Standardized protocol for a depletion of intramyocellular lipids (IMCL). *NMR Biomed.* **23**: 532–538.
43. Manco, M., G. Mingrone, A. V. Greco, E. Capristo, D. Gniuli, A. De Gaetano, and G. Gasbarrini. 2000. Insulin resistance directly correlates with increased saturated fatty acids in skeletal muscle triglycerides. *Metabolism*. **49**: 220–224.
44. Muoio, D. M. 2010. Intramuscular triacylglycerol and insulin resistance: Guilty as charged or wrongly accused? *Biochim. Biophys. Acta*. **1801**: 281–288.