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Applied high-intensity interval cardio yoga improves cardiometabolic fitness, energetic contributions, and metabolic flexibility in healthy adults

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Purpose: Currently, there is no interventional approach to increase the intensity of Surya Namaskar a popular hatha yoga sequence used worldwide. Therefore, this study investigated how tempo-based high-intensity interval cardio yoga (HIICY) and traditional interval hatha yoga (TIHY) affects cardiometabolic fitness in active adults.

Methods: Twenty physically active male and female individuals were randomly separated into HIICY (5 males, 5 females, 1.5 s tempo) and TIHY (5 males, 5 females, 3 s tempo) groups. The intervention included twelve exercise sessions for 4 weeks in both groups. Participants conducted a ramp test to determine their maximal oxygen uptake (\dot{VO}_{2max}), maximal velocity at \dot{VO}_{2max} (\dot{VVO}_{2max}), and maximal heart rate (HR_{max}). Afterward, they performed a 10-min high-intensity cardio yoga test (HICYT) to determine heart rate (HR_{peak} and HR_{mean}), oxygen uptake (\dot{VO}_{2peak} and \dot{VO}_{2mean}), respiratory exchange ratio (RER), blood lactate concentrations (La⁻_{peak} and Δ La⁻), fat and carbohydrate oxidations (FATox, CHOox), and energetic contributions (oxidative; W_{Oxi} , glycolytic; W_{Gly} , and phosphagen; W_{PCr} , total energy demand; W_{Total}).

Results: \dot{VO}_{2max} and \dot{vVO}_{2max} showed time and group x time interactions (p < 0.01, p < 0.001, p < 0.001, respectively). \dot{VO}_{2max} after HIICY was significantly higher than in pre-testing and following TIHY (p < 0.001, p < 0.0001, respectively). $\dot{VO}_{2peak'}$, \dot{VO}_{2mean} , RER, HR_{peak}, and HR_{mean} during the 10-min HICYT showed significant time effects (p < 0.05). ΔLa^- indicated a group x time interaction (p < 0.05). Group x time interaction effects for FATox at the fourth and sixth minute were observed (p < 0.05, respectively). Absolute (kJ) and relative (%) W_{Oxi} , W_{Gly} , and W_{Total} showed time and group x time interaction effects (p < 0.05). Additionally, \dot{VO}_{2max} and \dot{VO}_{2max} were highly correlated with W_{Oxi} in kJ (r = 0.91, 0.80, respectively). Moderate to high correlations were observed among CHOox, FATox, and absolute \dot{VO}_{2max} (r = 0.76, 0.62, respectively).

Conclusion: A 4-week period of HIICY improved cardiometabolic fitness, oxidative capacity, and metabolic flexibility compared with TIHY, in physically active adults. Therefore, HIICY is suitable as HY-specific HIIT and time-efficient approach for relatively healthy individuals.

KEYWORDS

metabolism, physiological adaptation, lactate, aerobic capacity, VO_{2max}

Introduction

The World Health Organization's (WHO) definition of "physical inactivity" refers to the failure to meet recommended levels of weekly physical activity. This guideline consists of at least 150 min of moderate-intensity or 75 min of high-intensity aerobic activity per week (Santos et al., 2023). It is now well appreciated that physical inactivity is the fourth leading cause of death and has been reported to contribute to multiple chronic diseases including insulin resistance, type 2 diabetes mellitus, obesity, metabolic and endocrine diseases (Balakumar et al., 2016; Kerr and Booth, 2022). The global economic burden of non-communicable diseases associated with lack of physical activity was estimated to be between US\$ 53.8 and 67.5 billion as of 2016 (Scheltens et al., 2021; Santos et al., 2023). One increased metabolic equivalent of task (1 MET = $\dot{V}O_2$ 3.5 mL·kg⁻¹·min⁻¹) at the maximal aerobic capacity was related to 13% and 15% risk reductions for all-cause mortality and coronary heart/cardiovascular disease respectively in males and females (Kodama et al., 2009). Furthermore, the prevalence of metabolic diseases was reduced by 6.3-fold in males and 4.9-fold in females.

Cardiorespiratory fitness consists of a wide range of parameters between an aerobic base and maximal aerobic exercise capacity, which represents maximal aerobic performance and the functional capacity of numerous bodily systems. These can be determined by maximal oxygen uptake (VO_{2max}) (Kodama et al., 2009; Meyler et al., 2021; Yang et al., 2022a; Hwang et al., 2022). \dot{VO}_{2max} is a crucial indicator for maximal aerobic performance and cardiorespiratory function in different individuals and athletes, and also provides detailed insights into inter-individual prescriptions in exercise physiology and sports science (Bosquet et al., 2002; Wisløff et al., 2007; Niemeyer et al., 2021). Moreover, WHO considers VO_{2max} as one of the valuable indicators or markers for cardiorespiratory fitness or health, and it is strongly associated with better physical performance. Epidemiological data from previous studies revealed that having a proportionally high VO_{2max} is a potent sign of health and life expectancy in all age groups (Ross et al., 2016; Udhan et al., 2018). Furthermore, exerciseinduced acute cardiorespiratory adaptations enhance the ability of the cardiovascular system to meet the demands of skeletal muscle exercise, via increases in pulmonary ventilation, heart rate, stroke volume, and cardiac output with moderate increases in systolic blood pressure, peripheral vasoconstriction, and vasodilation (Weiner and Baggish, 2012; Predel, 2014).

In terms of improving \dot{VO}_{2max} , the most effective intervention is high-intensity interval training (HIIT) (Buchheit and Laursen, 2013a; Batacan et al., 2017; MacInnis and Gibala, 2017; Sabag et al., 2022; Jacob et al., 2023). It is well known that HIIT protocols can maximally induce the oxygen uptake and utilization system for more than 85% of \dot{VO}_{2max} or peak oxygen uptake (\dot{VO}_{2peak}) and thus provide the most effective stimulation to increase \dot{VO}_{2max} (Buchheit and Laursen, 2013a; Buchheit and Laursen, 2013b; Sheykhlouvand et al., 2018).

HIIT prescription is used as an alternative physical exercise for those who do not have enough time contemporary people and is also consistently ranked in the top 10 fitness trends of the American College of Sports Medicine (ACSM) (Sabag et al., 2022; Kercher et al., 2023). Previous studies have reported that HIIT is efficient at developing the three energy systems utilized in humans in a time-efficient manner, which includes active or passive recovery following repeated rounds of exercise. HIIT protocols are designed to reach an exercise intensity such as >90% of maximal heart rate (HR_{max}), the second ventilatory threshold (>VT₂), over the second lactate threshold (>4 mmol·L⁻¹; zone 3: high-intensity exercise), and >85% of \dot{VO}_{2max} and \dot{VO}_{2peak} (Jamnick et al., 2020; Protzen et al., 2020).

In terms of metabolism, metabolic flexibility is defined as the ability to rapidly convert to generate adenosine triphosphate (ATP) from efficient fat and carbohydrate utilization based on physiological demands (Goodpaster and Sparks, 2017; Galgani et al., 2022). Efficient metabolic flexibility is essential to prevent metabolic diseases directly related to physical inactivity-induced mitochondrial dysfunction (San-Millán and Brooks, 2018; Yang et al., 2022a). HIIT increases metabolic flexibility, which indicates a strong relationship with metabolic health parameters such as insulin sensitivity and mitochondrial respiratory capacity (Aparecido et al., 2022).

In this regard, yoga is a mind-body practice consisting of physical postures (Asana), breathing techniques (Pranayama), and meditation (Dhyana) (Patwardhan, 2017). Notably, yoga is one of the most practiced complementary or alternative exercise interventions to achieve optimal physical and mental health (Udhan et al., 2018; Khoshnaw and Ghadge, 2021). Furthermore, hatha yoga (HY) is becoming increasingly popular in the United States and Europe as an alternative form of physical activity that can support individuals to reach globally recommended levels of physical activity (Larson-Meyer, 2016). The main goals of HY are to enable practitioners to improve body, breath, and spirit states and to prepare a healthy mind and body to immerse oneself in meditation for self-realization" (Schmalzl et al., 2015). A specific set of 19 asanas called Surya Namaskar B (Sun Salutations, SS) includes the vinyasa system and is one of the most basic and representative sequences in many styles of yoga classes worldwide (Papp et al., 2016; Brinsley et al., 2022).

The average metabolic cost during HY focusing on various postures, alignments, and breathing for different times is less than 3 METs and show results similar to walking on a treadmill at $3.2 \ km \cdot h^{-1}$ (Hagins et al., 2007; Larson-Meyer, 2016). However, HY (Surya Namaskar) lasts at least 10 min and can contribute sufficiently intense physical activity to improve an individual's cardiorespiratory fitness with 4-6 METs, corresponding to ACSM's moderate-to-high-intensity physical activity guidelines (Hagins et al., 2007; Mody, 2011; Larson-Meyer, 2016; Potiaumpai et al., 2016). A recent study by Lee et al. (2021) reported that HY, which consisted of 1.5 s for each asana and lasted for 10 min, could be used as HIIT in physically active individuals. The results of this study showed % HR_{peak} of HR_{max}, %HR_{mean} of HR_{max}, METs of VO_{2peak} and VO_{2mean}, and blood lactate concentrations (La⁻) values during high-intensity HY (HIHY), that reached 95.6%, 88.7%, 10.54, and 8.67 METs, and 8.31 mmol·L⁻¹ La⁻, respectively. This HIHY indicated suitable levels for HIIT. However, no interventional approach to HIHY was performed in this study. It was unclear therefore, whether highintensity interval cardio yoga (HIICY) as HIHY could improve VO_{2max}, energetic contributions, and metabolic flexibility in physically active individuals.

Therefore, the aim of the study was to investigate how at least 4week tempo-based HIICY and traditional interval hatha yoga



(TIHY) practices affected cardiometabolic fitness parameters, such as \dot{VO}_{2max} , energetic contributions, and metabolic flexibility.

Materials and methods

Participants

The sample size for the study was calculated using G*Power software version 3.1.9.4 (Heinlich Heine University, Düsseldorf, Germany) which considered: effect size = 0.3, alpha error probability = 0.05, and statistical power = 0.8. The effect size was determined based on previous studies (Astorino et al., 2012; Ormsbee et al., 2015; Lee et al., 2021). The total required sample size was calculated to be twenty participants assuming a 10% dropout rate (n = 20). Participants were randomly separated into two tempo-based HY groups: ten physically active individuals were assigned high-intensity interval cardio yoga (1.5 s tempo HIICY; five males and five females) and another 10 participated in traditional interval hatha yoga (3 s tempo TIHY; five males and five females) (Figure 1). All participants satisfied the minimum physical activity standards per week based on WHO guidelines and had no preexisting cardiovascular, pulmonary, or metabolic diseases or musculoskeletal disorders (Santos et al., 2023). All participants were yoga beginners. Therefore, before the experiment began, they learned the movement sequences with the yoga instructor and quickly became familiar with the movements and sequences through the qualified yoga instructor's demonstrations during the intervention. The participants maintained their previous physical activity levels throughout the 4-week intervention period and performed no additional training. Anthropometric measurements of all participants were taken in the fasting state. The data were as follows (mean \pm standard deviation; SD): age 30 \pm 5 years, height 169.6 \pm 7.4 cm, body mass 66.8 \pm 12.8, body fat 23.4% \pm 7.5%, and BMI of 23.0 \pm 3.3 kg·m⁻². The anthropometric data of HIICY and TIHY groups were not significantly different (Table 1). Participants were instructed not to change their diet during the intervention and did not take any medication before and during pre- and post-tests and abstained from nicotine and alcohol for 24 h prior to the testing. This study was approved by the Institutional Review Board of CHA University (IRB No. 1044308-202206-HR-031-02). The approved protocols followed the ethical standards in the Declaration of Helsinki. All participants signed informed consent forms.

Study procedure

All participants attended two laboratory visits for pre- and posttesting between 4 weeks. Testing was performed at a temperature of 23°C and relative humidity of 50%. Anthropometric data of participants were analyzed using an eight-electrode segmental multi-frequency bioelectrical impedance analyzer (BIA: 20–100 kHz; InBody 270; InBody Co. Ltd., Seoul, Republic of Korea). All participants rested for 2 h after lunchtime and conducted a ramp test for \dot{VO}_{2max} on a treadmill (NR30XA, DRAX Corporation Ltd., Seoul, Republic of Korea). Afterward, a high-intensity cardio yoga test (HICYT) (one

TABLE 1 Anthropometric data.

Parameters	HIICY (n = 10) (Mean ± SD)	TIHY (n = 10) (Mean \pm SD)	All participants (n = 20) (Mean ± SD)
Age (years)	27.6 ± 4.4	32.7 ± 5.2	30.2 ± 5.3
Height (cm)	169.7 ± 7.7	169.5 ± 7.5	169.6 ± 7.4
Body weight (kg)	66.2 ± 12.9	67.3 ± 13.4	66.8 ± 12.8
Body fat (%)	24.5 ± 7.3	22.3 ± 8.0	23.4 ± 7.5
BMI (kg·m ⁻²)	22.9 ± 3.6	23.2 ± 3.2	23.0 ± 3.3

There was no significant difference of anthropometric data between HIICY and TIHY groups. HIICY; high-intensity interval cardio yoga, TIHY; traditional interval hatha yoga. ns; p > 0.05.

set) was performed on the same testing day. \dot{VO}_{2max} and HICYT posttesting were conducted the same as pre-testing (Figure 1).

Maximal oxygen uptake (VO_{2max}) test

The pre- and post-ramp tests were performed via continuous incremental ramp protocols on a treadmill with the breath-bybreath method using a portable gas analyzer (MetaMax 3B; Cortex Biophysik, Leipzig, Germany). The gas analyzer was calibrated before each test with 15% O2 and 5% CO2 (Cortex Biophysik, Leipzig, Germany) and the turbine volume transducer was calibrated using a 3-L syringe (Hans Rudolph, Kansas, United States). An initial warm-up was conducted for 10 min by running at 70% of the estimated HR_{max} (Helgerud et al., 2022). The ramp protocol was applied for $\dot{V}O_{2max}$ determination based on a previous study (Sperlich et al., 2015). The initial running speed was 9.0 km \cdot h⁻¹ at 2% inclination in 2 min. Then, the running speed was increased by 0.72 km·h⁻¹ every 30 s. The investigator verbally encouraged participants to maintain effort for as long as possible to evaluate their maximum aerobic performance seen as reaching a "plateau". The test was stopped when $\dot{V}O_2$ plateau and respiratory exchange ratio (RER) >1.0 were reached, or until volitional exhaustion by the participant (Jurov et al., 2023; Wiecha et al., 2023). The plateau was determined using the method (<2 mL·kg⁻¹·min⁻¹) that has been explained in previous studies (Krustrup et al., 2005; Niemeyer et al., 2021). Furthermore, VO_{2max} was determined as an averaged value of oxygen uptake during the 15-s duration at the end of the plateau (Midgley et al., 2007; Niemeyer et al., 2021; Wiecha et al., 2023). The HR data were recorded using a Polar H10 sensor (Polar Electro, Kemple, Finland). HR_{max} was determined as values through the same section at \dot{VO}_{2max} (Krustrup et al., 2005). Finally, the absolute and relative $\dot{V}O_{2max}$ and velocity at $\dot{V}O_{2max}$ ($v\dot{V}O_{2max}$) were determined (Billat et al., 1996; Riboli et al., 2022).

High-intensity cardio yoga test (HICYT) and 4-week HIICY and TIHY interventions

After \dot{VO}_{2max} testing, the participants were advised to continue low-intensity jogging until blood lactate levels were below 2.0 mmol·L⁻¹ (Schünemann et al., 2023). They wore a portable gas analyzer and performed a high-intensity cardio yoga test (HICYT) (one set) for 10-min to analyze highest oxygen uptake (\dot{VO}_{2peak}), metabolic flexibility (fat and carbohydrate oxidation;

FATox and CHOox) and contributions from the three-energy system. Upon completion of the pre-test, all participants were randomly assigned to either HIICY or TIHY groups. They performed three interventional HIICY and TIHY sessions per week over 4 weeks (a total of 12 sessions) (Costigan et al., 2015; Schmitz et al., 2018) (Figure 1). All training sessions were performed under the supervision of a qualified yoga instructor. For both training sessions and HICYT sequences, the Surya Namaskar B sequence was used which consisted of 19 SS physical exercises (Asanas) (Figure 2). Tempo-based, each movement lasted 1.5 s for HIICY and 3 s for TIHY using a metronome, respectively (Lee et al., 2021). The entire duration for both styles was 30 min, which consisted of 2 sets of 10-min (Sandbakk et al., 2013) HIICY or TIHY and 10-min active recovery between sets. Active recovery was conducted by walking at 40%-45% of the estimated $\mathrm{HR}_{\mathrm{max}}$, as suggested by a previous study (Lee et al., 2021). The post-test was performed within 2 days of the last training session. Furthermore, HR levels were recorded during the 4-week interventions, and were digitally saved by the HR application for monitoring each training session (Table 2).

Calculations of fat and carbohydrate oxidation rate during HICYT

During HICYT, \dot{VO}_2 and carbon dioxide (\dot{VCO}_2) production were used to calculate metabolic flexibility in terms of fat (FATox) and carbohydrate (CHOox) oxidation, using stoichiometric equations according to previous studies (Jeukendrup and Wallis, 2005; San-Millán and Brooks, 2018; Yang et al., 2022a).

 $\begin{aligned} \text{FATox} & (\text{g} \cdot \text{min}^{-1}) = 1.67 \cdot \dot{\text{VO}}_2 \ (\text{L} \cdot \text{min}^{-1}) - 1.67 \cdot \dot{\text{VCO}}_2 \ (\text{L} \cdot \text{min}^{-1}) \\ \text{CHOox} \ (\text{g} \cdot \text{min}^{-1}) = 4.55 \cdot \dot{\text{VCO}}_2 \ (\text{L} \cdot \text{min}^{-1}) - 3.21 \cdot \dot{\text{VO}}_2 \ (\text{L} \cdot \text{min}^{-1}) \end{aligned}$

Calculations of energy system contributions (PCr-La⁻-O₂ method) during HICYT

The contributions of three energy systems (phosphagen $[W_{PCr}]$, glycolytic $[W_{Gly}]$, and oxidative $[W_{Oxi}]$ in kJ and %) were calculated by the PCr-La⁻-O₂ method (Julio et al., 2017; Yang et al., 2018; Yang et al., 2022b; Kaufmann et al., 2022; Yang et al., 2023). Oxygen consumption parameters (resting oxygen consumption, \dot{VO}_{2rest} ; average oxygen consumption during the 10-min HICYT,



TABLE 2 HR_{mean} data of 4-week HIICY and TIHY interventions (HIICY; n = 10 and TIHY; n = 10).

Group	HR _{mean} (beats·min ⁻¹) (Mean ± SD)	% of HR_{max} (mean ± SD)			
HIICY	178 ± 3.7	92.7 ± 2.5			
TIHY	128 ± 16.3	67.2 ± 8.3			

HR_{mean}, mean heart rate; % of HR_{max}, estimated maximal heart rate percentages; HIICY, high-intensity interval cardio yoga; TIHY, traditional interval hatha yoga.

 \dot{VO}_{2mean} ; highest oxygen consumption during the 10-min HICYT, \dot{VO}_{2peak} ; and fast component of excess oxygen consumption $[EPOC_{fast}]$; and off \dot{VO}_2 kinetics) were assessed at 5-min rest, during, and after 10-min HICYT (6 min), by the breath-by-breath method using a mobile gas analyzer. \dot{VO}_{2mean} and \dot{VO}_{2peak} were determined as the average and highest values during the 10-min HICYT, respectively. Capillary blood was collected from the earlobe (20 µL) before, and at 1-min intervals (1st to 10th) after 10-min HICYT to determine resting and maximal blood lactate concentrations (La⁻_{rest} and La⁻_{peak}; the peak value of La⁻ among 10 values) using an enzymaticamperometric sensor chip system (Biosen C-line; EKF diagnostics sales, GmbH, Barleben, Germany).

The W_{Oxi} was calculated by subtracting $\dot{\text{VO}}_{\text{2rest}}$ from $\dot{\text{VO}}_2$ by the trapezoidal method in which the domain under the O₂ data was divided into sections and the summarized trapezoid was utilized to estimate the integral (de Campos Mello et al., 2009; Yang et al., 2022b; Yang et al., 2023). $\dot{\text{VO}}_{\text{2rest}}$ was determined in a standing position on the yoga mat with the last 30 s of a 5 min period applied as a reference (di Prampero and Ferretti, 1999; Beneke et al., 2004; Julio et al., 2017; Yang et al., 2018; Kaufmann et al., 2022; Yang et al., 2022b).

The W_{Gly} analysis was performed by La⁻_{rest} and La⁻_{peak} values, assuming that an accumulation of 1 mmol·L⁻¹ was equivalent to 3 mL O₂·kg⁻¹ of body mass (di Prampero and Ferretti, 1999). Delta La⁻ (Δ La⁻) was calculated as the difference between La⁻_{peak} after 10-min HICYT and La⁻_{rest} before (Beneke et al., 2004; Campos et al., 2012; Yang et al., 2018; Yang et al., 2022b).

The W_{PCr} value was calculated using \dot{VO}_2 after 10-min HICYT and the fast component of excess post-exercise (Gastin, 2001; Beneke et al., 2004; Yang et al., 2022b; Yang et al., 2023). *Off* \dot{VO}_2 kinetics were fitted by mono-exponential and bi-exponential models using OriginPro 2021 statistical software (OriginLab Corp, Northampton, USA). The slow component of the bi-exponential model was negligible. Thus, \dot{VO}_2 values after 10-min HICYT were fitted using a mono-exponential model and W_{PCr} was estimated by calculating the integral of the exponential domain (Beneke et al., 2004; Campos et al., 2012; Julio et al., 2017; Yang et al., 2018; Yang et al., 2022b; Kaufmann et al., 2022).

A caloric quotient of 20.92 kJ was applied in the three energy system calculations (Gastin, 2001). Total energy demand was calculated as the sum of the three energy systems in kJ (Campos



Time and group × time interaction effects and comparisons of absolute and relative VO_{2max} and vVO_{2max} after 4-week interventions. (A) Absolute values of VO_{2max} in HIICY and TIHY groups between pre- and post-tests (B) relative values of VO_{2max} in HIICY and TIHY groups between pre- and post-tests, and (C) velocity at VO_{2max} in HIICY and TIHY groups between pre- and post-tests. ES; effect sizes, VO_{2max} ; maximal oxygen uptake, vVO_{2max} ; velocity at VO_{2max} , HIICY; high-intensity interval cardio yoga, TIHY; traditional interval hatha yoga. ns; p > 0.05, *p < 0.05, **p < 0.001, ****p < 0.0001.

et al., 2012). The relative energy system contributions were calculated in % compared to total energy demand.

Statistical analyses

All parameters were analyzed using GraphPad Prism 9.4.1 (GraphPad Prism Software Inc., La Jolla, CA, USA) and data are presented as mean ± standard deviation (SD). Normality of the data was analyzed using the Shapiro-Wilk test. All physiological variables, energetic contributions, and metabolic flexibility during HICYT were analyzed using a two-way (group x time) repeated-measures analysis of variance (ANOVA) with the Greenhouse-Geisser correction for violation of the sphericity assumption. If the main effect was significant, a Bonferroni post-hoc test was performed to compare among different conditions. The significance level was set at p <0.05. The effect sizes of repeated-measures ANOVA were calculated as partial eta squared $[\eta_p^2]$ and Cohen's [d] was utilized to indicate between different conditions. Thresholds for small, medium and large effects were considered ≥ 0.01 , ≥ 0.06 , and ≥ 0.14 for partial eta squared $[\eta_p^2]$ and ≥ 0.2 , ≥ 0.5 , and ≥ 0.8 for Cohen's [d], respectively (Fritz et al., 2012). Additionally, all data of both groups including pre- and post-tests (n = 40) were analyzed with two-tailed Pearson's correlation among absolute \dot{VO}_{2max} vs absolute W_{Oxi} , $v\dot{VO}_{2max}$ (km·h⁻¹) vs absolute W_{Oxi} , absolute VO_{2max} vs CHOox, and absolute VO_{2max} vs FATox (g·min⁻¹).

Results

VO_{2 max} between HIICY and TIHY

Two-way repeated-measures ANOVA for absolute, relative $\dot{V}O_{2max}$ and $v\dot{V}O_{2max}$ indicated significant time and group × time interaction effects (time effect: p = 0.0014; $[\eta_p^2]$: 0.70, group × time interaction: p < 0.0001; $[\eta_p^2]$: 0.86, time effect: p = 0.0007; $[\eta_p^2]$: 0.74, group × time interaction: p = 0.0003; $[\eta_p^2]$: 0.78, p = 0.0224; $[\eta_p^2]$: 0.46, group × time interaction: p = 0.0011; $[\eta_p^2]$: 0.71, respectively) (Figure 3). The absolute and relative \dot{VO}_{2max} values of the HIICY group in the post-test were significantly higher compared with the pre-test and compared with the post-test of the TIHY group (absolute \dot{VO}_{2max} : p < 0.0001; [d]: 0.33, p <0.0001; [d]: 0.26, relative \dot{VO}_{2max} , p = 0.0002; [d]: 0.67, p = 0.0002; [d]: 0.59, respectively) (Figures 3A, B). Furthermore, $v\dot{VO}_{2max}$ of the HIICY group in the post-test was significantly higher than in the pre-test (p =0.0154; [d]: 0.33) (Figure 3C).

Physiological parameters during HICYT

There was no significant difference in anthropometric data between HIICY and TIHY groups. Table 3 shows the physiological parameters during 10-min HICYT pre- and posttests between different groups. The values of absolute and relative \dot{VO}_{2peak} , \dot{VO}_{2mean} , HR_{peak} , HR_{mean} , and RER indicated significant time effects (p = 0.0263; effect size $[\eta_p^2] : 0.44$, p = 0.0121; $[\eta_p^2] : 0.52$, p = 0.0031; $[\eta_p^2] : 0.64$, p = 0.0053; $[\eta_p^2] : 0.60$, p = 0.0163; $[\eta_p^2] : 0.50$, p = 0.0061; $[\eta_p^2] : 0.59$, p = 0.0013; $[\eta_p^2] : 0.70$, respectively). Furthermore, a significant group × time interaction effect for ΔLa^- (p = 0.0198; $[\eta_p^2] : 0.47$) was observed (Table 3).

Energetic contribution during HICYT

Two-way repeated-measures ANOVA showed significant time effects in absolute (kJ) for W_{Oxi} and W_{Total} , and a group × time interaction effect in absolute W_{Gly} (time effects: p < 0.01; $[\eta_p^2]$: 0.54, p = 0.0173; $[\eta_p^2]$: 0.49, group × time interaction: p = 0.0479; $[\eta_p^2]$: 0.37, respectively). Moreover, significant group × time interaction effects in relative (%) W_{Oxi} and W_{Gly} were observed (p = 0.014; $[\eta_p^2]$: 0.50, p = 0.0084; $[\eta_p^2]$: 0.56, respectively) (Figures 4A, B). The relative W_{Gly} value of the HIICY group in the post-test was

Parameters	HIICY		TIHY		Significance p (ES η_p^2)		
	Pre-test	Post-test	Pre-test	Post-test	Group	Time	Group x Time
VO _{2peak} [L⋅min ⁻¹]	2.73 ± 0.74	2.90 ± 0.82	2.62 ± 0.54	2.77 ± 0.61	ns	*0.0263 (0.44)	ns
VO _{2peak} [mL·kg ^{−1} ·min ^{−1}]	40.97 ± 5.71	43.14 ± 6.37	39.22 ± 6.10	41.60 ± 6.10	ns	*0.0121 (0.52)	ns
VO _{2mean} [L·min ^{−1}]	2.37 ± 0.63	2.55 ± 0.66	2.29 ± 0.44	2.43 ± 0.51	ns	**0.0031 (0.64)	ns
VO _{2mean} [mL·kg ^{−1} ·min ^{−1}]	35.42 ± 4.39	38.00 ± 4.89	34.26 ± 4.76	36.48 ± 5.10	ns	**0.0053 (0.60)	ns
RER [VCO ₂ /VO ₂]	1.10 ± 0.05	1.15 ± 0.04	1.10 ± 0.03	1.10 ± 0.06	ns	*0.0163 (0.50)	ns
HR _{peak} [beats·min ⁻¹]	188.70 ± 5.96	183.80 ± 8.46	182.70 ± 7.69	180.80 ± 7.58	ns	**0.0061 (0.59)	ns
HR _{mean} [beats·min ⁻¹]	177.90 ± 5.60	171.70 ± 8.49	173.80 ± 8.30	168.3 ± 8.12	ns	**0.0013 (0.70)	ns
ΔLa^{-} [mmol·L ⁻¹]	8.57 ± 1.54	7.62 ± 2.98	7.44 ± 2.90	8.51 ± 1.65	ns	ns	*0.0198 (0.47)
La ⁻ _{neak} [mmol·L ⁻¹]	9.48 ± 1.58	8.73 ± 2.98	8.83 ± 2.90	9.59 ± 1.63	ns	ns	ns

TABLE 3 Physiological parameters during the 10-min high-intensity cardio yoga test between different groups (HIICY; n = 10 and TIHY; n = 10).

Data shown as mean \pm standard deviation (n = 20). HIICY, high-intensity interval cardio yoga; TIHY, traditional interval hatha yoga; ES, effect size; \dot{VO}_{2peak} , peak oxygen uptake; \dot{VO}_{2mean} , mean oxygen uptake; RER, respiratory exchange ratio; HR_{peak} , peak heart rate; HR_{mean} , mean heart rate; ΔLa^- , delta lactate; La^- , peak lactate; ns, p > 0.05, *p < 0.05, *p < 0.01.

significantly lower than in the pre-test (p = 0.0376; [d]: 0.68) (Figure 4B).

Metabolic flexibility including FATox and CHOox during HICYT

Two-way repeated-measures ANOVA showed group × time interaction effects at the fourth and time and group × time interaction effects at sixth minute of FATox (p = 0.0312; effect size $[\eta_p^2]$: 0.42, p = 0.0412; $[\eta_p^2]$: 0.39, p = 0.0386; $[\eta_p^2]$: 0.39, respectively) (Figures 5A–D). Otherwise, no significant main effect of FATox and CHOox and between pre- and post-tests were observed.

Relationships between metabolic flexibility, energetic contributions and physiological variables

High positive correlations were found between W_{Oxi} in kJ, and absolute $\dot{\text{VO}}_{2\text{max}}$, as well as $v\dot{\text{VO}}_{2\text{max}}$ (W_{Oxi} vs absolute $\dot{\text{VO}}_{2\text{max}}$: r =0.91, $R^2 = 0.83$, 95%CI: 0.84–0.95; p < 0.0001, W_{Oxi} vs $v\dot{\text{VO}}_{2\text{max}}$: r =0.80, $R^2 = 0.64$, 95%CI: 0.65–0.90; p < 0.0001, respectively) (Figures 6A, B). Furthermore, a high positive correlation between CHOox and absolute value of $\dot{\text{VO}}_{2\text{max}}$ (r = 0.76, $R^2 = 0.57$, 95%CI: 0.58–0.86; p < 0.0001) and moderate positive correlation between FATox and absolute value of $\dot{\text{VO}}_{2\text{max}}$ (r = 0.62; $R^2 = 0.39$; 95%CI: 0.39–0.78; p <0.0001) were found (Figures 6C, D).

Discussion

Training effects of HIIT and HY on health, oxidative capacity, and metabolism have been well reported previously. To date, however, there was no interventional approach to HIICY. It is unclear whether HIICY as HIHY can improve cardiometabolic fitness parameters such as VO_{2max}, energetic contributions, and metabolic flexibility in physically active adults. To the best of our knowledge, this is the first study to investigate how different intensities during 4-week tempo-based HY practices (HIICY and TIHY) affected VO_{2max}, energetic contributions, and metabolic flexibility. Our major findings indicate that time and group \times time interaction effects in absolute and relative VO_{2max} after 4-week HIICY and TIHY training regimens were observed. VO_{2max} was improved between pre- and post-testing in the HIICY but not in TIHY. During the HICYT, time and group × time interaction effects among W_{Oxi} in kJ, W_{Total} , W_{Gly} in kJ, W_{Oxi} in %, and W_{Gly} in % were found. The value of W_{Gly} in % was decreased only between pre- and post-tests in the HIICY group. Furthermore, time and group × time interaction effects in FATox showed at the fourth and sixth minute after HIICY and TIHY, and moderate to high correlations among \dot{VO}_{2max} vs W_{Oxi} , $v\dot{VO}_{2max}$ vs W_{Oxi}, VO_{2max} vs CHOox, and VO_{2max} vs FATox were found.

Regarding maximal aerobic performance, 4-week HIICY improved $\dot{V}O_{2max}$ and $v\dot{V}O_{2max}$ while 10-min TIHY showed no increase in VO_{2max} between pre- and post-tests in physically active individuals. In this regard, at least 10-min of Surya Namaskar B (SS) compared with general HY achieved an intensity that could improve cardiometabolic fitness (Hagins et al., 2007). Furthermore, physiological profiling of Surya Namaskar B between 3 s and 12 s tempos indicated exercise intensities that were moderate-to highintensity and low-intensity (Potiaumpai et al., 2016). A recent study by Lee et al. (2021) showed that Surya Namaskar B at a fast tempo of 1.5 s was suitable for HIIT (>90% of HR_{max}, ≥6 METs; vigorous/ heavy, >4 mmol·L⁻¹ La⁻; zone 3). These outcomes suggest that fast tempo-based HY practice is necessary, if the goal is to increase endurance performance and cardiometabolic fitness. The adjustment of exercise intensity and duration between work and recovery intervals alters the relative demand for specific metabolic pathways within muscle cells, oxygen delivery to the working muscle, and subsequent adaptations. These changes occur at the cellular and systemic levels depending on the specific nature of the training program (Laursen and Jenkins, 2002; Sandbakk et al., 2013; Franchini et al., 2019). HIIT with long-duration (5-10 min) and



shorter intervals of higher intensity such as seen in supramaximal intensity have been shown to be an effective metabolic stimulus for improving $\dot{V}O_{2max}$. Long-duration intervals such as 5–10 min increase $\dot{V}O_{2max}$ and $\dot{V}O_2$ at ventilatory thresholds in national-level junior cross-country skiers (Sandbakk et al., 2013). Therefore, previous study outcomes support our results showing that the long-duration during HIIT is a relevant factor in increasing $\dot{V}O_{2max}$ in well-trained athletes as well as in the general population (Gaskill et al., 1999; Billat, 2001; Laursen and Jenkins, 2002; Seiler, 2010; Seiler et al., 2013).

 $\rm HR_{peak}$ and $\rm HR_{mean}$ tended to improve post-testing in the HIICY group. Significant increases in pulmonary oxygen uptake and skeletal muscle oxygen demand cause HIIT-induced cardiovascular adaptations during intensive and prolonged HIIT sessions. Exercise-induced improvement in $\rm \dot{VO}_{2max}$ is associated with enlarged red blood cell volume, which leads to greater oxygen-rich blood capacity and increased stroke volume. Thus, it can affect an increase in O₂ transport capacity (Predel, 2014). Sabag et al. (2022) indicated in their topical review that systolic and diastolic function, maximum cardiac output, capillary density, and stroke volume were increased after HIIT. Indeed,

the usefulness of HIIT in stimulating significant improvements in cardiometabolic fitness, left ventricular ejection fraction, and pathological left ventricular remodeling of participants was emphasized for those who completed 36 sessions of HIIT over 2–3 months (Hsu et al., 2019). These preliminary findings suggest that increased stroke volume can deliver more oxygen per heartbeat (Kohn et al., 2011). A reduced HR at submaximal intensity after HIIT may be one of the major HIIT-induced physiological adaptations, and one which also associates with increased \dot{VO}_{2max} (Acevedo and Goldfarb, 1989; Kubukeli et al., 2002).

The reduced value of W_{Gly} with increased \dot{VO}_{2max} between pre- and post-testing in the HIICY group (Figure 4B) is likely associated with increased HIIT-induced metabolic responses (Batacan et al., 2017; MacInnis and Gibala, 2017; Chrøis et al., 2020; Sabag et al., 2022). Generally, cellular stress is proportional to exercise intensity and there is strong evidence that higher exercise intensities induce elevated molecular responses compared to moderate intensities (Egan and Zierath, 2013). This may be because higher rates of fuel utilization relies more on carbohydrate oxidation, uses more glycogen, and increases ATP



turnover (Howlett et al., 1998; Van loon et al., 2001). Consequently, it activates signaling pathways involved in mitochondrial biogenesis following intracellular lactate production, creatine kinase, ADP, and AMP accumulation (Howlett et al., 1998; Van loon et al., 2001) phosphorylation of AMP-activated protein kinase, p38 mitogenactivated protein kinase, and Ca2+/calmodulin-dependent protein kinase II, and expression of peroxisome proliferatoractivated receptor gamma coactivator 1-alpha (PGC-1a) mRNA (Gibala et al., 2009; Egan et al., 2010; Little et al., 2011; Kristensen et al., 2015; Metcalfe et al., 2015; Bersiner et al., 2023). Regular and repeated activation of these pathways increases mitochondrial density (Coffey and Hawley, 2007). Greater activation of these specific kinases were induced by highintensity exercise compared to low-intensity exercise resulting in greater expression of mRNA for PGC-1a which is a master regulator of mitochondrial biogenesis (Egan et al., 2010). PGC-1a is responsible for the activation of mitochondrial transcription factors such as nuclear respiratory factors 1 and 2 and the mitochondrial transcription factor A (Knuiman et al., 2015). Downstream of these metabolic signals, mitochondrial protein synthesis has been shown to result in higher mitochondrial biogenesis in response to sustained training performed at higher intensities with a given amount of exercise (Di Donato et al., 2014).

Furthermore, numerous studies have shown an increased density of monocarboxylate transporters (MCT) 1 and 4 after HIIT (Burgomaster et al., 2007; Perry et al., 2008; McGinley and Bishop, 2016). Higher MCT 1 and 4 density increased lactate transport and likely supported a reduction in glycogen breakdown and La⁻ at a given intensity (Perry et al., 2008). Accordingly, group × time interaction effect of energetic contributions was observed and the relative value of W_{Gly} post-testing in the HIICY group was decreased (Figure 4B). Because of the aforementioned physiological adaptations, oxidative capacity might be improved, which would in turn increase the lactate elimination rate and increase ATP re-synthesis during high-intensity aerobic exercise (Burgomaster et al., 2008; MacInnis and Gibala, 2017; Brooks, 2018; Hwang et al., 2022; Yang et al., 2023). Indeed, muscle glycogen is likely conserved, delaying the onset of muscle fatigue and improving oxidative exercise performance (Hearris et al., 2018). Also, high positive correlations among W_{Oxi} in kJ and absolute $\dot{\text{VO}}_{2\text{max}}$ as well as $v\dot{V}O_{2max}$ in our study outcomes, support the conclusion of a HIIT-induced physiological adaptation (Figure 6).

In terms of metabolic flexibility, time and group × time interaction effects of FATox at the fourth and sixth during the HICYT between HIICY and TIHY groups and moderate to high correlations among \dot{VO}_{2max} vs CHOox, and \dot{VO}_{2max} vs FATox were found (Figures 5, 6). These results in FATox may be affected by peripheral improvements at the level of muscle cells, such as in mitochondrial function. Such



FIGURE 6

Relationships between absolute $\dot{V}O_{2max}$, $v\dot{V}O_{2max}$ and absolute W_{Oxi} , and CHOox, FATox and absolute $\dot{V}O_{2max}$. (A) two-tailed Pearson's correlation between absolute $\dot{V}O_{2max}$. (B) two-tailed Pearson's correlation between absolute W_{Oxi} and $v\dot{V}O_{2max}$. (C) two-tailed Pearson's correlation between CHOox and absolute $\dot{V}O_{2max}$, and (D) two-tailed Pearson's correlation between FATox and absolute $\dot{V}O_{2max}$. (C) two-tailed Pearson's correlation between CHOox and absolute $\dot{V}O_{2max}$, and (D) two-tailed Pearson's correlation between FATox and absolute $\dot{V}O_{2max}$. C); confidence interval, $\dot{V}O_{2max}$; maximal oxygen uptake, W_{Oxi} ; oxidative energy demand, $v\dot{V}O_{2max}$; velocity at $\dot{V}O_{2max}$, CHOox; carbohydrate oxidation, FATox, fat oxidation.

changes have been, reported in a number of earlier studies that have detailed improvements in mitochondrial function and insulin sensitivity, and which together represent improved metabolic flexibility (Sandbakk et al., 2013; Chrøis et al., 2020; Yang et al., 2022a; Sabag et al., 2022). Furthermore, a higher percentage of FATox during HIIT than shortinterval training in about half of previous studies was observed (Astorino and Schubert, 2018). In detail, significant increases in β -hydroxyacyl acyl-CoA dehydrogenase, citrate synthase, fatty acid-binding protein, carnitine palmitoyl transferase I or fatty acid translocase/cluster of differentiation 36, and expression of PGC-1a are likely responsible for increased FATox after HIIT (Tunstall et al., 2002; Perry et al., 2008; Talanian et al., 2010; Astorino and Schubert, 2018; Warren et al., 2020). Burgomaster et al. (2008) reported that 18 sessions of HIIT for 6 weeks increased glucose transporter isoform 4 content and promoted glucose uptake during recovery, and greater muscle glycogen levels. Several weeks of HIIT can increase muscle FATox capacity which is associated with more hydroxyacyl-CoA dehydrogenase activity and improved insulin resistance. Also of note, HY has been shown to reduce adipose cell concentrations in the visceral area, diminishing or minimizing the excess free fatty acids released by adipose cells (Khoshnaw and Ghadge, 2021).

In sum, this study has demonstrated increased cardiometabolic fitness, including VO_{2max}, energetic contributions, and metabolic flexibility after a 30-min HIICY regimen (2 sets, 3 times per week) for 4 weeks compared with TIHY in physically active adults. Moreover, we investigated whether HIICY for at least 4 weeks significantly affected the cardiometabolic biomarkers described above. Therefore, we suggest that the positively affected outcomes likely depend on exercise intensity influencing the activation of PGC-1a, the master regulator of mitochondrial biogenesis in human skeletal muscle (Egan et al., 2010; Gibala et al., 2012). Also, the high-intensity level should be maintained as much as possible during repeated sessions, with this being more important than maintaining duration or frequency to induce an increase in VO_{2max} (Hickson and Rosenkoetter, 1981). Therefore, HIICY consisting of a specific sequence of 19 asanas is a fast tempo-based approach, which can provide cardiometabolic health benefits in physically active individuals.

This study does have some limitations. First, it only targeted one cohort of physically active adults and the HIICY approach should be investigated in more diverse populations such as in athletes, sedentary people, and clinical populations in further studies. Second, the small sample size might have influenced our results. Therefore, more participants should be recruited in a future study. Third, experimenting with efficient periodization to progressively increase HY intensity from TIHY to HIICY should be considered. Finally, more direct measurements during training using proteomics and metabolomics, and the measurement of fluorescent protein tools should be investigated to determine molecular responses in the future.

Conclusion

Our findings indicate that 4-week of HIICY compared with TIHY improved cardiometabolic fitness, oxidative capacity, and metabolic flexibility in physically active individuals. Therefore, HIICY is a suitable training regimen for HY-specific HIIT, and is appropriate for relatively healthy individuals who may have HY experience but need timeefficient exercise options. Through our study outcomes, it is once again confirmed that exercise intensity during HY is important to improving VO_{2max}. While HY is an effective practice in many respects, significant and efficient improvements to cardiometabolic fitness require fast-tempo HIICY. Moreover, a proportionately higher VO_{2max} strongly signals greater cardiovascular health and life expectancy at any age. Thus, this substitute system of practice may prevent cardiac/metabolic diseases in the general population. However, HIICY has a very high intensity for exercise beginners. Therefore, it is recommended to start practicing with 3 s tempo-based TIHY during an initial period within the typical linear periodization model for an aerobic base, and then gradually increase the intensity to HIICY.

Data availability statement

The original contributions presented in the study are included in the article/Supplementary material, further inquiries can be directed to the corresponding author.

Ethics statement

The studies involving humans were approved by the Institutional Review Board of CHA University. The studies were conducted in accordance with the local legislation and institutional

References

Acevedo, E. O., and Goldfarb, A. H. (1989). Increased training intensity effects on plasma lactate, ventilatory threshold, and endurance. *Med. Sci. Sports Exerc* 21(5), 563–568. doi:10.1249/00005768-198910000-00011

Aparecido, J. M. L., Marquezi, M. L., Couto, H. L. d. O., Santos, T. M. d. S., Cruz, A. F. C., Lopes, N. B., et al. (2022). Six HIT sessions improve cardiorespiratory fitness and metabolic flexibility in insulin resistant and insulin sensitive adolescents with obesity. *Int. J. Environ. Res. Public Health* 19 (17), 10568. doi:10.3390/ijerph191710568

Astorino, T. A., Allen, R. P., Roberson, D. W., and Jurancich, M. (2012). Effect of high-intensity interval training on cardiovascular function, VO2max, and muscular force. J. Strength Cond. Res. 26 (1), 138–145. doi:10.1519/JSC.0b013e318218dd77

Astorino, T. A., and Schubert, M. M. (2018). Changes in fat oxidation in response to various regimes of high intensity interval training (HIIT). *Eur. J. Appl. Physiol.* 118, 51–63. doi:10.1007/s00421-017-3756-0

Balakumar, P., Maung-U, K., and Jagadeesh, G. (2016). Prevalence and prevention of cardiovascular disease and diabetes mellitus. *Pharmacol. Res.* 113, 600–609. doi:10. 1016/j.phrs.2016.09.040

requirements. The participants provided their written informed consent to participate in this study. Written informed consent was obtained from the individual(s) for the publication of any potentially identifiable images or data included in this article.

Author contributions

S-YP: Conceptualization, Data curation, Formal Analysis, Investigation, Methodology, Visualization, Writing-original draft, Writing-review and editing. W-HY: Conceptualization, Formal Analysis, Methodology, Project administration, Resources, Supervision, Writing-review and editing.

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Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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Batacan, R. B., Jr., Duncan, M. J., Dalbo, V. J., Tucker, P. S., and Fenning, A. S. (2017). Effects of high-intensity interval training on cardiometabolic health: A systematic review and meta-analysis of intervention studies. *Br. J. Sports Med.* 51 (6), 494–503. doi:10.1136/bjsports-2015-095841

Beneke, R., Beyer, T., Jachner, C., Erasmus, J., and Hütler, M. (2004). Energetics of karate kumite. *Eur. J. Appl. Physiol.* 92 (4-5), 518–523. doi:10.1007/s00421-004-1073-x

Bersiner, K., Park, S. Y., Schaaf, K., Yang, W. H., Theis, C., Jacko, D., et al. (2023). Resistance exercise: A mighty tool that adapts, destroys, rebuilds and modulates the molecular and structural environment of skeletal muscle. *Phys. Act. Nutr.* 27 (2), 78–95. doi:10.20463/pan.2023.0021

Billat, L. (2001). Interval training for performance: a scientific and empirical practice. Special recommendations for middle- and long-distance running. Part I: aerobic interval training. *Sports Med.* 31 (1), 13–31. doi:10.2165/00007256-200131010-00002

Billat, V. L., Hill, D. W., Pinoteau, J., Petit, B., and Koralsztein, J. P. (1996). Effect of protocol on determination of velocity at VO2 max and on its time to exhaustion. *Arch. Physiol. Biochem.* 104 (3), 313–321. doi:10.1076/apab.104.3.313.12908

Bosquet, L., Léger, L., and Legros, P. (2002). Methods to determine aerobic endurance. Sports Med. 32 (11), 675–700. doi:10.2165/00007256-200232110-00002

Brinsley, J., Smout, M., Girard, D., and Davison, K. (2022). Acute mood and cardiovascular responses to moderate intensity vinyasa yoga, static yin yoga and aerobic exercise in people with depression and/or anxiety disorders: A 5-arm randomised controlled trial. *Ment. Health Phys. Act.* 22, 100450. doi:10.1016/j.mhpa.2022.100450

Brooks, G. A. (2018). The science and translation of lactate shuttle theory. *Cell. Metab.* 27 (4), 757–785. doi:10.1016/j.cmet.2018.03.008

Buchheit, M., and Laursen, P. B. (2013a). High-intensity interval training, solutions to the programming puzzle: part I: cardiopulmonary emphasis. *Sports Med.* 43 (5), 313–338. doi:10.1007/s40279-013-0029-x

Buchheit, M., and Laursen, P. B. (2013b). High-intensity interval training, solutions to the programming puzzle: part II: anaerobic energy, neuromuscular load and practical applications. *Sports Med.* 43 (10), 927–954. doi:10.1007/s40279-013-0066-5

Burgomaster, K. A., Cermak, N. M., Phillips, S. M., Benton, C. R., Bonen, A., and Gibala, M. J. (2007). Divergent response of metabolite transport proteins in human skeletal muscle after sprint interval training and detraining. *Am. J. Physiol. Regul. Integr. Comp. Physiol.* 292 (5), R1970–R1976. doi:10.1152/ajpregu.00503.2006

Burgomaster, K. A., Howarth, K. R., Phillips, S. M., Rakobowchuk, M., MacDonald, M. J., McGee, S. L., et al. (2008). Similar metabolic adaptations during exercise after low volume sprint interval and traditional endurance training in humans. *J. Physiol.* 586 (1), 151–160. doi:10.1113/jphysiol.2007.142109

Campos, F. A., Bertuzzi, R., Dourado, A. C., Santos, V. G., and Franchini, E. (2012). Energy demands in taekwondo athletes during combat simulation. *Eur. J. Appl. Physiol.* 112 (4), 1221–1228. doi:10.1007/s00421-011-2071-4

Chrøis, K. M., Dohlmann, T. L., Søgaard, D., Hansen, C. V., Dela, F., Helge, J. W., et al. (2020). Mitochondrial adaptations to high intensity interval training in older females and males. *Eur. J. Sport Sci.* 20 (1), 135–145. doi:10.1080/17461391.2019.1615556

Coffey, V. G., and Hawley, J. A. (2007). The molecular bases of training adaptation. *Sports Med.* 37, 737–763. doi:10.2165/00007256-200737090-00001

Costigan, S. A., Eather, N., Plotnikoff, R. C., Taaffe, D. R., and Lubans, D. R. (2015). High-intensity interval training for improving health-related fitness in adolescents: a systematic review and meta-analysis. *Br. J. Sports Med.* 49 (19), 1253–1261. doi:10.1136/ bjsports-2014-094490

de Campos Mello, F., de Moraes Bertuzzi, R. C., Grangeiro, P. M., and Franchini, E. (2009). Energy systems contributions in 2,000 m race simulation: A comparison among rowing ergometers and water. *Eur. J. Appl. Physiol.* 107 (5), 615–619. doi:10.1007/s00421-009-1172-9

Di Donato, D. M., West, D. W., Churchward-Venne, T. A., Breen, L., Baker, S. K., and Phillips, S. M. (2014). Influence of aerobic exercise intensity on myofibrillar and mitochondrial protein synthesis in young men during early and late postexercise recovery. *Am. J. Physiol. Endocrinol. Metab.* 306(9), E1025–E1032. doi:10.1152/ ajpendo.00487.2013

di Prampero, P. E., and Ferretti, G. (1999). The energetics of anaerobic muscle metabolism: A reappraisal of older and recent concepts. *Respir. Physiol.* 118(2-3), 103–115. doi:10.1016/s0034-5687(99)00083-3

Egan, B., Carson, B. P., Garcia-Roves, P. M., Chibalin, A. V., Sarsfield, F. M., Barron, N., et al. (2010). Exercise intensity-dependent regulation of peroxisome proliferatoractivated receptor coactivator-1 mRNA abundance is associated with differential activation of upstream signalling kinases in human skeletal muscle. *J. Physiol.* 588 (10), 1779–1790. doi:10.1113/jphysiol.2010.188011

Egan, B., and Zierath, J. R. (2013). Exercise metabolism and the molecular regulation of skeletal muscle adaptation. *Cell. Metab.* 17 (2), 162–184. doi:10.1016/j.cmet.2012. 12.012

Franchini, E., Cormack, S., and Takito, M. Y. (2019). Effects of high-intensity interval training on olympic combat sports athletes' performance and physiological adaptation: A systematic review. *J. Strength Cond. Res.* 33 (1), 242–252. doi:10.1519/JSC. 000000000002957

Fritz, C. O., Morris, P. E., and Richler, J. J. (2012). Effect size estimates: current use, calculations, and interpretation. J. Exp. Psychol. Gen. 141 (1), 2–18. doi:10.1037/a0024338

Galgani, J. E., Bergouignan, A., Rieusset, J., Moro, C., and Nazare, J.-A. (2022). Editorial: metabolic flexibility. *Front. Nutr.* 1254, 946300. doi:10.3389/fnut.2022.946300

Gaskill, S. E., Serfass, R. C., Bacharach, D. W., and Kelly, J. M. (1999). Responses to training in cross-country skiers. *Med. Sci. Sports Exerc* 31, 1211–1217. doi:10.1097/00005768-199908000-00020

Gastin, P. B. (2001). Energy system interaction and relative contribution during maximal exercise. *Sports Med.* 31 (10), 725-741. doi:10.2165/00007256-200131100-00003

Gibala, M. J., Little, J. P., MacDonald, M. J., and Hawley, J. A. (2012). Physiological adaptations to low-volume, high-intensity interval training in health and disease. *J. Physiol.* 590 (5), 1077–1084. doi:10.1113/jphysiol.2011.224725

Gibala, M. J., McGee, S. L., Garnham, A. P., Howlett, K. F., Snow, R. J., and Hargreaves, M. (2009). Brief intense interval exercise activates AMPK and p38 MAPK signaling and increases the expression of PGC-1alpha in human skeletal muscle. *J. Appl. Physiol.* 106 (3), 929–934. doi:10.1152/japplphysiol.90880.2008

Goodpaster, B. H., and Sparks, L. M. (2017). Metabolic flexibility in health and disease. Cell. Metab. 25 (5), 1027-1036. doi:10.1016/j.cmet.2017.04.015

Hagins, M., Moore, W., and Rundle, A. (2007). Does practicing hatha yoga satisfy recommendations for intensity of physical activity which improves and maintains health and cardiovascular fitness? *BMC Complement. Altern. Med.* 7 (1), 40–49. doi:10. 1186/1472-6882-7-40

Hearris, M. A., Hammond, K. M., Fell, J. M., and Morton, J. P. (2018). Regulation of muscle glycogen metabolism during exercise: implications for endurance performance and training adaptations. *Nutrients* 10 (3), 298. doi:10.3390/nu10030298

Helgerud, J., Haglo, H., and Hoff, J. (2022). Prediction of vo2max from submaximal exercise using the smartphone application myworkout go: validation study of a digital health method. *JMIR Cardio* 6 (2), e38570. doi:10.2196/38570

Hickson, R. C., and Rosenkoetter, M. A. (1981). Reduced training frequencies and maintenance of increased aerobic power. *Med. Sci. Sports Exerc* 13(1), 13–16. doi:10. 1249/00005768-198101000-00011

Howlett, R. A., Parolin, M. L., Dyck, D. J., Hultman, E., Jones, N. L., Heigenhauser, G. J., et al. (1998). Regulation of skeletal muscle glycogen phosphorylase and PDH at varying exercise power outputs. *Am. J. Physiol.* 275 (2), R418–R425. doi:10.1152/ajpregu.1998.275.2.R418

Hsu, C.-C., Fu, T.-C., Yuan, S.-S., Wang, C.-H., Liu, M.-H., Shyu, Y.-C., et al. (2019). High-intensity interval training is associated with improved long-term survival in heart failure patients. *J. Clin. Med.* 8 (3), 409. doi:10.3390/jcm8030409

Hwang, J., Moon, N.-R., Heine, O., and Yang, W.-H. (2022). The ability of energy recovery in professional soccer players is increased by individualized low-intensity exercise. *PLoS One* 17 (6), e0270484. doi:10.1371/journal.pone.0270484

Jacob, N., So, I., Sharma, B., Marzolini, S., Tartaglia, M. C., Oh, P., et al. (2023). Effects of high-intensity interval training protocols on blood lactate levels and cognition in healthy adults: systematic review and meta-regression. *Sports Med.* 53 (5), 977–991. doi:10.1007/s40279-023-01815-2

Jamnick, N. A., Pettitt, R. W., Granata, C., Pyne, D. B., and Bishop, D. J. (2020). An examination and critique of current methods to determine exercise intensity. *Sports Med.* 50 (10), 1729–1756. doi:10.1007/s40279-020-01322-8

Jeukendrup, A., and Wallis, G. (2005). Measurement of substrate oxidation during exercise by means of gas exchange measurements. *Int. J. Sports Med.* 26 (S 1), S28–S37. doi:10.1055/s-2004-830512

Julio, U. F., Panissa, V. L. G., Esteves, J. V., Cury, R. L., Agostinho, M. F., and Franchini, E. (2017). Energy-system contributions to simulated judo matches. *Int. J. Sports Physiol. Perform.* 12 (5), 676–683. doi:10.1123/ijspp.2015-0750

Jurov, I., Cvijić, M., and Toplišek, J. (2023). Predicting VO2max in competitive cyclists: is the FRIEND equation the optimal choice? *Front. Physiol.* 14, 987006. doi:10. 3389/fphys.2023.987006

Kaufmann, S., Latzel, R., Beneke, R., and Hoos, O. (2022). Reliability of the 3component model of aerobic, anaerobic lactic, and anaerobic alactic energy distribution (PCr-la-O2) for energetic profiling of continuous and intermittent exercise. *Int. J. Sports Physiol. Perform.* 17 (11), 1642–1648. doi:10.1123/ijspp.2022-0115

Kercher, V. M., Kercher, K., Levy, P., Bennion, T., Alexander, C., Amaral, P. C., et al. (2023). 2023 fitness trends from around the globe. *ACSM's Health Fit. J.* 27 (1), 19–30. doi:10.1249/fit.00000000000836

Kerr, N. R., and Booth, F. W. (2022). Contributions of physical inactivity and sedentary behavior to metabolic and endocrine diseases. *Trends Endocrinol. Metab.* 33 (12), 817–827. doi:10.1016/j.tem.2022.09.002

Khoshnaw, D. M., and Ghadge, A. A. (2021). Yoga as a complementary therapy for metabolic syndrome: A narrative review. *J. Integr. Med.* 19 (1), 6–12. doi:10.1016/j.joim. 2020.09.002

Knuiman, P., Hopman, M. T., and Mensink, M. (2015). Glycogen availability and skeletal muscle adaptations with endurance and resistance exercise. *Nutr. Metab. (Lond)* 12 (1), 59–11. doi:10.1186/s12986-015-0055-9

Kodama, S., Saito, K., Tanaka, S., Maki, M., Yachi, Y., Asumi, M., et al. (2009). Cardiorespiratory fitness as a quantitative predictor of all-cause mortality and cardiovascular events in healthy men and women: A meta-analysis. *Jama* 301 (19), 2024–2035. doi:10.1001/jama.2009.681

Kohn, T., Essén-Gustavsson, B., and Myburgh, K. (2011). Specific muscle adaptations in type II fibers after high-intensity interval training of well-trained runners. *Scand. J. Med. Sci. Sports* 21 (6), 765–772. doi:10.1111/j.1600-0838.2010.01136.x

Kristensen, D. E., Albers, P. H., Prats, C., Baba, O., Birk, J. B., and Wojtaszewski, J. F. (2015). Human muscle fibre type-specific regulation of AMPK and downstream targets by exercise. *J. Physiol.* 593 (8), 2053–2069. doi:10.1113/ jphysiol.2014.283267

Krustrup, P., Mohr, M., Ellingsgaard, H., and Bangsbo, J. (2005). Physical demands during an elite female soccer game: importance of training status. *Med. Sci. Sports Exerc* 37 (7), 1242–1248. doi:10.1249/01.mss.0000170062.73981.94

Kubukeli, Z. N., Noakes, T. D., and Dennis, S. C. (2002). Training techniques to improve endurance exercise performances. *Sports Med.* 32, 489–509. doi:10.2165/00007256-200232080-00002

Larson-Meyer, D. E. (2016). A systematic review of the energy cost and metabolic intensity of yoga. *Med. Sci. Sports Exerc* 48 (8), 1558–1569. doi:10.1249/MSS. 000000000000922

Laursen, P. B., and Jenkins, D. G. (2002). The scientific basis for high-intensity interval training: optimising training programmes and maximising performance in highly trained endurance athletes. *Sports Med.* 32 (1), 53–73. doi:10.2165/00007256-200232010-00003

Lee, K.-H., Ju, H.-M., and Yang, W.-H. (2021). Metabolic energy contributions during high-intensity hatha yoga and physiological comparisons between active and passive (savasana) recovery. *Front. Physiol.* 12, 743859. doi:10.3389/fphys.2021.743859

Little, J. P., Safdar, A., Bishop, D., Tarnopolsky, M. A., and Gibala, M. J. (2011). An acute bout of high-intensity interval training increases the nuclear abundance of PGC-1 α and activates mitochondrial biogenesis in human skeletal muscle. *Am. J. Physiol. Regul. Integr. Comp. Physiol.* 300, R1303–R1310. R1303-R1310. doi:10.1152/ajpregu.00538.2010

MacInnis, M. J., and Gibala, M. J. (2017). Physiological adaptations to interval training and the role of exercise intensity. J. Physiol. 595 (9), 2915–2930. doi:10.1113/JP273196

McGinley, C., and Bishop, D. J. (2016). Influence of training intensity on adaptations in acid/base transport proteins, muscle buffer capacity, and repeated-sprint ability in active men. J. Appl. Physiol. 121(6), 1290–1305. doi:10.1152/japplphysiol.00630.2016

Metcalfe, R. S., Koumanov, F., Ruffino, J. S., Stokes, K. A., Holman, G. D., Thompson, D., et al. (2015). Physiological and molecular responses to an acute bout of reducedexertion high-intensity interval training (REHIT). *Eur. J. Appl. Physiol.* 115, 2321–2334. doi:10.1007/s00421-015-3217-6

Meyler, S., Bottoms, L., and Muniz-Pumares, D. (2021). Biological and methodological factors affecting VO2max response variability to endurance training and the influence of exercise intensity prescription. *Exp. Physiol.* 106 (7), 1410–1424. doi:10.1113/EP089565

Midgley, A. W., McNaughton, L. R., Polman, R., and Marchant, D. (2007). Criteria for determination of maximal oxygen uptake: A brief critique and recommendations for future research. *Sports Med.* 37, 1019–1028. doi:10.2165/00007256-200737120-00002

Mody, B. S. (2011). Acute effects of Surya Namaskar on the cardiovascular & metabolic system. J. Bodyw. Mov. Ther. 15 (3), 343–347. doi:10.1016/j.jbmt.2010.05.001

Niemeyer, M., Knaier, R., and Beneke, R. (2021). The oxygen uptake plateau-A critical review of the frequently misunderstood phenomenon. *Sports Med.* 51 (9), 1815–1834. doi:10.1007/s40279-021-01471-4

Ormsbee, M. J., Ward, E. G., Bach, C. W., Arciero, P. J., McKune, A. J., and Panton, L. B. (2015). The impact of a pre-loaded multi-ingredient performance supplement on muscle soreness and performance following downhill running. *J. Int. Soc. Sports Nutr.* 12 (1), 2. doi:10.1186/s12970-014-0063-6

Papp, M. E., Lindfors, P., Nygren-Bonnier, M., Gullstrand, L., and Wändell, P. E. (2016). Effects of high-intensity hatha yoga on cardiovascular fitness, adipocytokines, and apolipoproteins in healthy students: A randomized controlled study. *J. Altern. Complement. Med.* 22 (1), 81–87. doi:10.1089/acm.2015.0082

Patwardhan, A. R. (2017). Yoga research and public health: is research aligned with the stakeholders' needs? *J. Prim. Care Community Health* 8 (1), 31–36. doi:10.1177/2150131916664682

Perry, C. G., Heigenhauser, G. J., Bonen, A., and Spriet, L. L. (2008). High-intensity aerobic interval training increases fat and carbohydrate metabolic capacities in human skeletal muscle. *Appl. Physiol. Nutr. Metab.* 33 (6), 1112–1123. doi:10.1139/H08-097

Potiaumpai, M., Martins, M. C. M., Rodriguez, R., Mooney, K., and Signorile, J. F. (2016). Differences in energy expenditure during high-speed versus standard-speed yoga: A randomized sequence crossover trial. *Complement. Ther. Med.* 29, 169–174. doi:10.1016/j.ctim.2016.10.002

Predel, H.-G. (2014). Marathon run: cardiovascular adaptation and cardiovascular risk. *Eur. Heart J.* 35 (44), 3091–3098. doi:10.1093/eurheartj/eht502

Protzen, G. V., Bartel, C., Coswig, V. S., Gentil, P., and Del Vecchio, F. B. (2020). Physiological aspects and energetic contribution in 20s: 10s high-intensity interval exercise at different intensities. *PeerJ* 8, e9791. doi:10.7717/peerj.9791

Riboli, A., Coratella, G., Rampichini, S., Limonta, E., and Esposito, F. (2022). Testing protocol affects the velocity at VO2max in semi-professional soccer players. *Res. Sports Med.* 30 (2), 182–192. doi:10.1080/15438627.2021.1878460

Ross, R., Blair, S. N., Arena, R., Church, T. S., Després, J.-P., Franklin, B. A., et al. (2016). Importance of assessing cardiorespiratory fitness in clinical practice: A case for fitness as a clinical vital sign: A scientific statement from the American heart association. *Circulation* 134 (24), e653–e699. doi:10.1161/CIR.00000000000461

Sabag, A., Little, J. P., and Johnson, N. A. (2022). Low-volume high-intensity interval training for cardiometabolic health. *J. Physiol.* 600 (5), 1013–1026. doi:10.1113/JP281210

San-Millán, I., and Brooks, G. A. (2018). Assessment of metabolic flexibility by means of measuring blood lactate, fat, and carbohydrate oxidation responses to exercise in professional endurance athletes and less-fit individuals. *Sports Med.* 48 (2), 467–479. doi:10.1007/s40279-017-0751-x

Sandbakk, Ø., Sandbakk, S. B., Ettema, G., and Welde, B. (2013). Effects of intensity and duration in aerobic high-intensity interval training in highly trained junior cross-country skiers. *J. Strength Cond. Res.* 27 (7), 1974–1980. doi:10.1519/JSC.0b013e3182752f08

Santos, A. C., Willumsen, J., Meheus, F., Ilbawi, A., and Bull, F. C. (2023). The cost of inaction on physical inactivity to public health-care systems: A population-attributable fraction analysis. *Lancet Glob. Health* 11 (1), e32–e39. doi:10.1016/S2214-109X(22)00464-8

Scheltens, P., De Strooper, B., Kivipelto, M., Holstege, H., Chételat, G., Teunissen, C. E., et al. (2021). Alzheimer's disease. *Lancet* 397 (10284), 1577–1590. doi:10.1016/S0140-6736(20)32205-4

Schmalzl, L., Powers, C., and Henje Blom, E. (2015). Neurophysiological and neurocognitive mechanisms underlying the effects of yoga-based practices: towards a comprehensive theoretical framework. *Front. Hum. Neurosci.* 9, 235. doi:10.3389/ fnhum.2015.00235

Schmitz, B., Rolfes, F., Schelleckes, K., Mewes, M., Thorwesten, L., Krüger, M., et al. (2018). Longer work/rest intervals during high-intensity interval training (HIIT) lead to elevated levels of miR-222 and miR-29c. *Front. Physiol.* 9, 395. doi:10.3389/fphys.2018.00395

Schünemann, F., Park, S. Y., Wawer, C., Theis, C., Yang, W. H., and Gehlert, S. (2023). Diagnostics of ${}^{v}_{La.max}$ and glycolytic energy contribution indicate individual characteristics of anaerobic glycolytic energy metabolism contributing to rowing performance. *Metabolites* 13 (3), 317. doi:10.3390/metabol3030317

Seiler, S., Jøranson, K., Olesen, B. V., and Hetlelid, K. J. (2013). Adaptations to aerobic interval training: interactive effects of exercise intensity and total work duration. *Scand. J. Med. Sci. Sports* 23 (1), 74–83. doi:10.1111/j.1600-0838.2011.01351.x

Seiler, S. (2010). What is best practice for training intensity and duration distribution in endurance athletes? *Int. J. Sports Physiol. Perform.* 5 (3), 276–291. doi:10.1123/ijspp.5.3.276

Sheykhlouvand, M., Gharaat, M., Khalili, E., Agha-Alinejad, H., Rahmaninia, F., and Arazi, H. (2018). Low-volume high-intensity interval versus continuous endurance training: effects on hematological and cardiorespiratory system adaptations in professional canoe polo athletes. *J. Strength Cond. Res.* 32 (7), 1852–1860. doi:10. 1519/JSC.00000000020121

Sperlich, P. F., Holmberg, H.-C., Reed, J. L., Zinner, C., Mester, J., and Sperlich, B. (2015). Individual versus standardized running protocols in the determination of VO2max. *J. Sports Sci. Med.* 14 (2), 386–393.

Talanian, J. L., Holloway, G. P., Snook, L. A., Heigenhauser, G. J., Bonen, A., and Spriet, L. L. (2010). Exercise training increases sarcolemmal and mitochondrial fatty acid transport proteins in human skeletal muscle. *Am. J. Physiol. Endocrinol. Metab.* 299 (2), E180–E188. doi:10.1152/ajpendo.00073.2010

Tunstall, R. J., Mehan, K. A., Wadley, G. D., Collier, G. R., Bonen, A., Hargreaves, M., et al. (2002). Exercise training increases lipid metabolism gene expression in human skeletal muscle. *Am. J. Physiol. Endocrinol. Metab.* 283 (1), E66–E72. doi:10.1152/ ajpendo.00475.2001

Udhan, V. D., WankheDe, S. G., and Phatale, S. R. (2018). Effect of yoga on cardio-respiratory health markers: physical fitness index and maximum oxygen consumption (VO 2 max). *J. Clin. Diagn. Res.* 12 (8), 21–23. doi:10.7860/JCDR/2018/36819.11932

Van loon, L. J., Greenhaff, P. L., Constantin-Teodosiu, D., Saris, W. H., and Wagenmakers, A. J. (2001). The effects of increasing exercise intensity on muscle fuel utilisation in humans. J. Physiol. 536 (1), 295–304. doi:10.1111/j.1469-7793.2001.00295.x

Warren, J. L., Hunter, G. R., Gower, B. A., Bamman, M. M., Windham, S. T., Moellering, D. R., et al. (2020). Exercise effects on mitochondrial function and lipid metabolism during energy balance. *Med. Sci. Sports Exerc* 52 (4), 827–834. doi:10.1249/ MSS.0000000000002190

Weiner, R. B., and Baggish, A. L. (2012). Exercise-induced cardiac remodeling. Prog. Cardiovasc Dis. 54 (5), 380–386. doi:10.1016/j.pcad.2012.01.006

Wiecha, S., Kasiak, P. S., Cieśliński, I., Takken, T., Palka, T., Knechtle, B., et al. (2023). External validation of VO2max prediction models based on recreational and elite endurance athletes. *PLoS One* 18 (1), e0280897. doi:10.1371/journal.pone.0280897

Wisløff, U., Støylen, A., Loennechen, J. P., Bruvold, M., Rognmo, Ø., Haram, P. M., et al. (2007). Superior cardiovascular effect of aerobic interval training versus moderate continuous training in heart failure patients: A randomized study. *Circulation* 115 (24), 3086–3094. doi:10.1161/CIRCULATIONAHA.106. 675041

Yang, W.-H., Heine, O., and Grau, M. (2018). Rapid weight reduction does not impair athletic performance of Taekwondo athletes–A pilot study. *PLoS One* 13 (4), e0196568. doi:10.1371/journal.pone.0196568

Yang, W.-H., Park, J.-H., Park, S.-Y., and Park, Y. (2022a). Energetic contributions including gender differences and metabolic flexibility in the general population and athletes. *Metabolites* 12 (10), 965. doi:10.3390/metabol2100965

Yang, W.-H., Park, S.-Y., Kim, T., Jeon, H.-J., Heine, O., and Gehlert, S. (2023). A modified formula using energy system contributions to calculate pure maximal rate of lactate accumulation during a maximal sprint cycling test. *Front. Physiol.* 14, 1147321. doi:10.3889/fphys.2023.1147321

Yang, W. H., Park, J. H., Shin, Y. C., and Kim, J. (2022b). Physiological profiling and energy system contributions during simulated epée matches in elite fencers. *Int. J. Sports Physiol. Perform.* 17 (6), 943–950. doi:10.1123/ijspp.2021-0497