

Associations of childhood-to-adulthood body size trajectories and genetic susceptibility with the risks of osteoarthritis: a population-based cohort study of UK Biobank data



Muhui Zeng, Shibo Chen, Tianxiang Fan, Hao Yang, Peihua Cao, Zhiqiang Wang, Yujie Zhang, Yan Zhang, David Hunter, Qian Yang, Changhai Ding, Zhaohua Zhu

Abstract

Lancet Glob Health 2023

Published Online March 2, 2023

This online publication has been corrected. The corrected version first appeared at thelancet.com/lancetgh on March 17, 2023

Clinical Research Centre (M Zeng MBBS, S Chen PhD, H Yang MBBS, P Cao PhD. Z Wang PhD, Y Zhang PhD, D Hunter MD PhD, C Ding PhD, Z Zhu PhD) and Department of Orthopedics (Z Zhu), Zhuijang Hospital, and Department of Epidemiology, School of Public Health (Y Zhang), Southern Medical University, Guangzhou, China; Department of Orthopedics, General Hospital of Southern Theater Command of PLA, Guangzhou, China (M Zeng): Department of Rehabilitation Sciences, The Hong Kong Polytechnic University, Hong Kong Special Administrative Region, China (T Fan PhD); Department of Rheumatology, Royal North Shore Hospital and Sydney Musculoskeletal Health, Kolling Institute, University of Sydney, Sydney, NSW, Australia (D Hunter); MRC Integrative Epidemiology Unit

Department of Rheumatology, Guangzhou First People's Hospital, School of Medicine, South China University of Technology, Guangzhou, China (C Ding); Department of Orthopaedics, Affiliated Hospital of Youjiang Medical University for Nationalities. Baise, China (C Ding)

at University of Bristol, Bristol,

UK (Q Yang PhD); Population Health Sciences, Bristol Medical School, University of Bristol,

Bristol, UK (Q Yang);

Correspondence to: Dr Zhaohua Zhu, Clinical Research Centre, Zhujiang Hospital of Southern Medical University, Guangzhou 510280, China zhaohua.zhu@utas.edu.au Background Large adulthood body size was associated with increased risk of osteoarthritis. We aimed to examine the association between body size trajectories from childhood to adulthood and potential interactions with genetic susceptibility on osteoarthritis risk.

Methods We included participants from the UK Biobank aged 38-73 years in 2006-10. Childhood body size information was collected by questionnaire. Adulthood BMI was assessed and transformed into three categories (<25 kg/m² for normal, 25-29 · 9 kg/m² for overweight, and >30 kg/m² for obesity). A Cox proportional hazards regression model was applied to assess the association between body size trajectories and osteoarthritis incidence. Osteoarthritis-related polygenic risk score (PRS) was constructed to evaluate its interactions with body size trajectories on osteoarthritis risk.

Findings For the 466 292 participants included, we identified nine body size trajectories [thinner to normal (11.6%), overweight (17·2%), or obesity (26·9%); average to normal (11·8%), overweight (16·2%), or obesity (23·7%); and plumper to normal (12.3%), overweight (16.2%), or obesity (23.6%)]. Compared with individuals in the average-tonormal group, all other trajectory groups had higher risks of osteoarthritis, after adjustment for demographic, socialeconomic and lifestyle covariates (hazard ratios [HRs] 1.05-2.41; all p<0.01). Among them, thinner-to-obesity (HR 2·41; 95% CI 2·23-2·49) had the most prominent association with increased osteoarthritis risk. A high PRS was significantly associated with an increased risk of osteoarthritis (1·14; 1·11–1·16), whereas no interaction between childhood-to-adulthood body size trajectories and PRS on osteoarthritis risks was observed. The population attributable fraction suggested that body size towards normal in adulthood could eliminate osteoarthritis cases by 18.67% for thinner-to-overweight to 38.74% for plumper-to-obesity.

Interpretation Average-to-normal body size seems to be the healthiest childhood-to-adulthood trajectory for osteoarthritis risk, whereas a trajectory of increased body size from thinner to obesity has the highest risk for osteoarthritis. These associations are independent of osteoarthritis genetic susceptibility.

Funding The National Natural Science Foundation of China (32000925) and Guangzhou Science and Technology Program (202002030481).

Copyright © 2023 The Author(s). Published by Elsevier Ltd. This is an Open Access article under the CC BY 4.0 license.

ZZ, MZ, QY, SC and YZ designed the study. MZ, SC, TF, HY, PC, YZ, and YQ managed and analyzed the data. MZ wrote the first draft of the article. All authors contributed to data interpretation and preparation of the report. All authors provided intellectual content to the manuscript and approved the submission.

Declaration of interests

We declare no competing interests.