Check for updates

OPEN ACCESS

EDITED BY Philipp Maximilian Doldi, LMU Munich University Hospital, Germany

REVIEWED BY Avik Ghose Tata Consultancy Services (India), India

*CORRESPONDENCE Mohamed Elgendi I moe.elgendi@hest.ethz.ch Carlo Menon I carlo.menon@hest.ethz.ch

[†]These authors have contributed equally to this work.

RECEIVED 08 June 2023 ACCEPTED 14 August 2023 PUBLISHED 25 August 2023

CITATION

Elgendi M, Wu W, Guan C and Menon C (2023) Revolutionizing smartphone gyrocardiography for heart rate monitoring: overcoming clinical validation hurdles.

Front. Cardiovasc. Med. 10:1237043. doi: 10.3389/fcvm.2023.1237043

COPYRIGHT

© 2023 Elgendi, Wu, Guan and Menon. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.

Revolutionizing smartphone gyrocardiography for heart rate monitoring: overcoming clinical validation hurdles

Mohamed Elgendi^{1*†}, Wenshan Wu^{1,2†}, Cuntai Guan² and Carlo Menon^{1*}

¹Biomedical and Mobile Health Technology Lab, Department of Health Sciences and Technology, ETH Zurich, Zurich, Switzerland, ²School of Computer Science and Engineering, Nanyang Technological University, Singapore, Singapore

Accurate heart rate (HR) measurement is crucial for optimal cardiac health, and while conventional methods such as electrocardiography and photoplethysmography are widely used for continuous daily monitoring, they may face practical limitations due to their dependence on external sensors and susceptibility to motion artifacts. In recent years, mechanocardiography (MCG)-based technologies, such as gyrocardiography (GCG) and seismocardiography (SCG), have emerged as promising alternatives to address these limitations. GCG has shown enhanced sensitivity and accuracy for HR detection compared to SCG, although its benefits are often overlooked in the context of the widespread use of accelerometers in HR monitoring applications. In this perspective, we aim to explore the potential and challenges of GCG, while recognizing that other technologies, including photoplethysmography and remote photoplethysmography, also have promising applications for HR monitoring. We propose a roadmap for future research to unlock the transformative capabilities of GCG for everyday heart rate monitoring.

KEYWORDS

smartphone gyroscope, heart rate monitoring, gyrocardiography, cardiac waveform, heart rate measurement, public health, digital health, digital medicine

Introduction

The critical importance of heart rate (HR) measurements in detecting irregularities in heart rhythm patterns and cardiovascular health conditions is well established in the literature (1–7). Traditional HR measurement techniques, such as electrocardiograms (ECG) (8), demand sophisticated equipment and specialized knowledge for accurate interpretation, rendering them costly and inaccessible for long-term, everyday monitoring. The widespread adoption of smartphones in recent years (9) has given rise to an array of smartphone-based HR measurement systems employing ECG (10, 11) and photoplethysmography (PPG) (12–14). However, the reliance on smartphone-based ECG systems specialized external hardware sensors poses challenges to precise ambulatory measurements, while the accuracy of smartphone-based PPG recordings is compromised by excessive noise.

Mechanocardiography (MCG) encompasses gyrocardiography (GCG), seismocardiography (SCG), ballistocardiography (BCG), and phonocardiography (15), and increasing numbers of studies attest to its potential for cardiac performance assessment (16). Unlike other methods, MCG directly measures cardiac mechanics induced by heartbeats (17, 18). Despite their

advantages, most smartphone-based HR monitoring algorithms rely on SCG signals, with few considering GCG data for HR estimation. Furthermore, published studies on the subject have been largely authored by a small group of investigators (19–24), highlighting a lack of awareness regarding GCG's potential for HR estimation within the broader research community.

In this perspective article, we aim to analyze the potential and challenges associated with the use of GCG for HR estimation, propose future research directions, and raise awareness of its capabilities. Moreover, we introduce four evaluation metrics to standardize the evaluation process, enabling a fairer comparison between methodologies and fostering further advancements in the field.

Exploring the potential of smartphonebased GCG for heart rate detection

Despite accelerometers being the most commonly used sensors for heart rate monitoring, recent literature suggests that due to the heart's helical shape up to 60% of cardiac vibrational energy is contained in the gyration signal (25, 26). Furthermore, since gyroscope measurements are not affected by gravity (27), GCG signal collection is largely independent of the user's position or posture (27); in addition proper axis selection could even result in GCG signals outperforming a combination of GCG and SCG signals for heart rate estimation (28), raising the possibility that GCG signals could supplant the currently popular SCG signals and shape the future of HR estimation.

The practicality of this application can be significantly enhanced if it can accommodate different measuring locations and postures. Most studies have only conducted HR estimation using MCG signals collected from subjects in a supine position (19–24, 28). However, vertical postures, such as standing or walking, can modulate autonomic regulation of cardiovascular function and capture different HR patterns from supine postures (29–30), owing to an increase in hydrostatic pressure in the thigh (31). Furthermore, placing the smartphone on or near the chest, though reasonable due to its proximity to the heart, could limit the application's usefulness. In most studies, measurements required the phone to be on a bare chest or subjects to wear light clothing, restricting its usage in cold weather conditions.

These constraints may hinder the applicability of these methods in real-life situations where changes in chest pressure may lead to signal variations when subjects assume different postures or engage in physical activities. Methods capable of accurately estimating HR using a smartphone in these contexts would represent a significant advancement in the field. Such models would not only prove valuable in HR estimation but also in identifying signal patterns and characteristics under these conditions. If the models can account for the effects of posture and physical activity on MCG signals, adjusting HR estimations based on these factors to achieve accurate HR predictions becomes feasible.

While some studies have already explored HR estimation using different subject positions (e.g., standing and walking) and measuring locations (e.g., in the hand (32) and pants pocket (33)), these methods utilized either SCG signals (32, 33) or MCG

signals from non-smartphone sources. Given the potential of GCG signals, further investigation into GCG signal collection from various postures and sites for HR estimation is warranted.

Based on the literature (21, 34–39), we proposed a general workflow of using GCG signals from smartphones for HR monitoring, as shown in Figure 1. The smartphones collect GCG data from the user in various locations and orientations. The raw GCG signals are then processed (filtered, feature extraction, and then HR calculation) in the cloud. The computed HRs are sent back to the smartphone, which generates a series of recommendations as feedback to the user. In emergency situations, an alert will be sent to the hospital for faster communication to ensure that medical treatments can be delivered to the user in time.

Obstacles encountered in smartphone GCG-based heart rate monitoring

The application of microelectromechanical systems (MEMS) vibrating structure gyroscopes (VSG) found in smartphones to heart rate monitoring is novel, and although these gyro sensors can potentially detect vital biomarkers such as blood pressure and heart rate, they were not specifically engineered for medical applications. Consequently, additional data cleaning and artifact removal steps must be implemented to achieve the necessary precision and accuracy for reliable heart rate readings.

One of the primary hurdles in employing gyro sensors for heart rate detection is the presence of motion artifacts produced by activities such as walking, running, or even typing. The orientation of the smartphone—for example, held in one's hand versus resting on a stable surface—can also significantly impact the heart rate readings. Additionally, gyro bias error, inherent to MEMS sensors, is a significant factor in gyroscope drift error and can lead to inaccurate readings. Notably, the yaw axis is most susceptible to gyroscope drift (40). As a result, thorough time series analysis of the signals and proper axis selection are crucial for compensating for these factors and generating accurate gyroscope signals (41).

While the use of smartphone gyro sensors offers potential for heart rate detection, it is not without challenges. The accuracy and reliability of the readings can be affected by these obstacles, which must be carefully addressed when using smartphone gyroscopes for dependable heart rate monitoring.

Wider challenges in smartphonebased heart rate monitoring

Lack of clinical data

Before gyroscope data from smartphones can be harnessed for dependable heart rate (HR) monitoring, it is crucial to establish clinical evidence of the proposed methods' accuracy. This demands rigorous testing and validation using a sufficiently sized and diverse participant pool, ensuring that the developed methods are unbiased and universally applicable.



Obtaining medical data, especially in emerging fields such as this, is challenging and time-consuming, as the process often requires costly specialized equipment and expert personnel. Additionally, the sensitive nature of medical data mandates strict adherence to patient privacy and data protection regulations, further complicating data collection, processing, and sharing. Despite these obstacles, concerted efforts are being made to collect and disseminate medical data for research advancement.

Inter- and intra-subject variability

A truly effective model should account for individual variability in physiology and movement patterns. This necessitates incorporating demographic data such as race, gender, and age to create personalized models that adapt to each individual's characteristics and behavior. Additionally, environmental factors like temperature and humidity can influence the signals HR estimation signals, as can electromagnetic interference, which can introduce noise and complicate the extraction of meaningful information.

Moreover, psychological factors, including anxiety, stress, and depression, can impact the cardiovascular system and, consequently, alter heartbeat patterns. Addressing these complexities in heart rate monitoring models ensures their adaptability and applicability across diverse scenarios and individuals, paving the way for more accurate and personalized health monitoring solutions.

Gender effects

The use of smartphone signal data for HR estimation involves non-invasive measurements of vibrations and rotations related to cardiomechanics using accelerometers and gyroscopes placed on the body surface near the chest. However, anatomical differences between men and women can affect the transmission of mechanical vibrations generated by the heart to the surface of the chest, leading to different directions and amplitudes of the MCG signals. Men tend to have larger and broader chests (42), while women tend to have smaller hearts, which can affect the amplitude and timing of the signals resulting from heartbeats. Differences in the shape of the heart and body fat distribution between genders (43) can also alter the mechanical vibrations transmitted to the chest. Wrist-worn wearables for HR estimation have been found to produce higher device errors in males (44), underscoring the importance of gender-specific physiological differences and the need for a balanced gender ratio in subject selection for smartphone-based HR validation studies.

Obesity

Obesity is a well-known cardiovascular risk factor that has been linked to an increased likelihood of heart rhythm disorders, which

can have a significant impact on heart rate patterns (45–47). Anatomical fat distribution can indirectly affect the waveforms produced by the heart, as it can alter the position and orientation of the heart itself. This, in turn, can impact the mechanical forces acting on the chest wall and the signals measured by medical equipment. For example, excess abdominal fat may push the heart upwards and backwards, altering the signal's morphology and amplitude. Additionally, differences in metabolism associated with different fat deposition patterns can also influence the function of the cardiovascular system, leading to changes in cardiac output that can influence the signal.

Although the body mass index (BMI) is widely used as an indicator of relative obesity, it is calculated from height and body weight and does not capture the body fat distribution (48). Therefore, it is generally regarded as a weak indicator of obesity. More effective anthropometric indicators such as chest circumference, waist-hip ratio, waist circumference, hip circumference, and waist stature ratio have been proposed as alternative measures of obesity.

Age distribution

As individuals age, the ability of cardiac muscle cells to divide and regenerate declines, which can lead to a deterioration of cardiac function and an increased risk of heart disease. Aging also causes anatomical and physiological changes within the cardiovascular system that can affect the mechanical forces acting on the chest wall (49). The heart muscle can become stiffer and thicker, especially the left ventricular wall, which can impair the heart's ability to fill with blood and contract effectively. Additionally, the heart may increase in size (50), leading to changes in its shape and position within the chest. These changes in cardiac muscle stiffness, thickness, and size can alter the transmission of mechanical vibrations from the heart to the chest wall, leading to changes in the morphology and amplitude of the signal. Furthermore, the accumulation of lipofuscin, a pigment that results from the buildup of waste products within cells, can impair the function of cardiac muscle cells. The number of pacemaker cells in the heart also decreases with age, which can affect the heart's ability to regulate its own rhythm (51). The alterations in the number and function of pacemaker cells can lead to changes in the timing and frequency of heartbeats, resulting in an irregular rhythm. As in obesity, aging may also lead to changes in total and regional fat distribution, with increased fat deposition often seen in internal organs, including the heart (52), further impacting the signals obtained from the heart. Overall, the interplay between these factors, as well as other comorbidities, can lead to changes in the MCG signals obtained from older adults, potentially requiring adjustments in their interpretation.

Comorbidity

Altered transmission of mechanical signals from the heart to the chest wall can occur with comorbidities such as hypertension and diabetes, which can cause changes in arterial stiffness (53), or respiratory diseases like chronic obstructive pulmonary disease (COPD), which can affect lung mechanics (54). Inflammatory conditions like rheumatoid arthritis and systemic lupus

erythematosus can change joint and muscle mechanics (55, 56), which can affect the position of the chest wall relative to the heart. Other comorbidities that impact the autonomic nervous system, such as heart failure and atrial fibrillation, can lead to changes in heart rate variability and other measures of cardiac autonomic function and to variations in the amplitude and timing of MCG signals (57, 58). Thus, it is important to consider the presence of various comorbidities when interpreting MCG signals, as they can introduce confounding factors affecting the accuracy and reliability of measurements and downstream HR predictions.

Performance evaluation metrics

Many studies have used smartphone MCG data, including GCG signals, to derive HR, but without proper evaluation metrics for HR estimation; however, performance evaluation metrics are essential to evaluate HR estimation methods for accuracy and precision, especially when health and safety decisions are involved. Standardized and consistent metrics for evaluating HR estimation performance can enable objective compare different HR estimation methods, improve reliability, identify effective and efficient methods, and highlight areas for improvement.

There are two aspects to having unified and consistent metrics for evaluating HR estimation performance: a standardized gold standard and evaluation metrics. Some studies have used suboptimal reference methods like HR derived from pulse oximeters or PPG signals, which themselves have varying degrees of error. This can introduce unnecessary measurement errors and potentially undermine findings. Therefore, the gold standard of ECG should ideally be used as the reference signal for HR estimation.

This article proposes four metrics for standardizing the evaluation of HR estimation performance: mean absolute errors (MAE), root mean squared errors (RMSE), Pearson correlation coefficient (r), and equivalence testing. The MAE measures general measurement error, the RMSE detects large prediction errors, the r metric measures the overall statistical relationship between estimated and reference values, and equivalence testing investigates agreements between estimated and reference measurements. A 95% equivalence testing ($\alpha = 0.05$) would be typically used.

Equivalence testing is preferred over the standard two-sample t-test because it confirms statistical equivalence and requires acceptance criteria to be known a priori (59–61). However, it requires the criterion to be based on either empirical evidence or domain knowledge. Researchers could follow previous studies (62, 63) that pre-defined an equivalence zone of $\pm 10\%$ of the reference mean, and perform a statistical test to determine whether the 90% confidence interval from the estimations falls within the equivalence zone with 95% precision ($\alpha = 0.05$).

The lack of standards for evaluating and reporting accuracy reduces transparency and accountability of the method, which could lead to contradictory results, lack of generalizability, false findings, and reproducibility issues. Utilizing a unified and consistent set of standards can avoid these problems, improve the clinical utility of findings in this field, and facilitate objective comparisons of different HR estimation methods.

Conclusion

HR estimation using GCG signals from smartphones is a new and promising field that has gained attention in recent years. However, there are still relatively few studies dedicated to this purpose, which could be attributed to a lack of awareness about the potential of gyroscopes for HR detection. Our work aims to increase awareness of this field and highlight the potential of GCG signals for HR estimation. We also presented some of the challenges that need to be addressed and proposed a gold standard and four evaluation metrics for performance evaluation. By considering these factors and utilizing standardized performance evaluation metrics, comparing results across studies and establishing standard protocols for data analysis can become easier, advancing this field and potentially leading to more robust downstream classification applications.

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 authors.

Author contributions

ME and WW conducted the background and literature search, contributed to the development and refinement of the idea. ME led the study. ME, WW, CG, and CM conceived the study, contributed to the revision and editing of the manuscript, and approved the

References

1. Perret-Guillaume C, Joly L, Benetos A. Heart rate as a risk factor for cardiovascular disease. *Prog Cardiovasc Dis.* (2009) 52:6–10. doi: 10.1016/j.pcad. 2009.05.003

2. Palatini P. Elevated heart rate as a predictor of increased cardiovascular morbidity. Am J Hypertens. (1999) 17:S3-10. doi: 10.1016/S0895-7061(98)00207-6

3. Holmqvist F, Kim S, Steinberg BA, Reiffel JA, Mahaffey KW, Gersh BJ, et al. Heart rate is associated with progression of atrial fibrillation, independent of rhythm. *Heart*. (2015) 101:894–9. doi: 10.1136/heartjnl-2014-307043

4. Indolfi C, Ross J Jr. The role of heart rate in myocardial ischemia, infarction: implications of myocardial perfusion-contraction matching. *Prog Cardiovasc Dis.* (1993) 36:61–74. doi: 10.1016/0033-0620(93)90022-6

5. Palatini P, Julius S. Elevated heart rate: a major risk factor for cardiovascular disease. *Clin Exp Hypertens.* (2004) 26:637–44. doi: 10.1081/CEH-200031959

6. Cook S, Togni M, Schaub MC, Wenaweser P, Hess OM. High heart rate: a cardiovascular risk factor? *Eur Heart J.* (2006) 27:2387–93. doi: 10.1093/eurheartj/ehl259

7. Böhm M, Swedberg K, Komajda M, Borer JS, Ford I, Dubost-Brama A, et al. Heart rate as a risk factor in chronic heart failure (shift): the association between heart rate and outcomes in a randomised placebo-controlled trial. *Lancet.* (2010) 376:886–94. doi: 10.1016/S0140-6736(10)61259-7

8. Berkaya SK, Uysal AK, Gunal ES, Ergin S, Gunal S, Gulmezoglu MB. A survey on ECG analysis. *Biomed Signal Process Control.* (2018) 43:216–35. doi: 10.1016/j.bspc. 2018.03.003

9. [Dataset] GSMA. Global smartphone penetration rate as share of population from 2016 to 2021 (2022) [Online]. Available from: https://www.statista.com/statistics/203734/global-smartphone-penetration-per-capita-since-2005/ (Accessed March 22, 2023).

final manuscript. All authors contributed to the article and approved the submitted version.

Funding

Open access funding by ETH Zurich.

Acknowledgments

This project was a collaboration between ETH Zurich in Switzerland and Nanyang Technical University in Singapore, and it was supported by the Chen-Ning Yang Scholars Programme in Singapore.

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.

Publisher's note

All claims expressed in this article are solely those of the authors and do not necessarily represent those of their affiliated organizations, or those of the publisher, the editors and the reviewers. Any product that may be evaluated in this article, or claim that may be made by its manufacturer, is not guaranteed or endorsed by the publisher.

10. Walker A, Muhlestein J. Smartphone electrocardiogram monitoring: current perspectives. Adv Health Care Technol. (2018) 4:15–24. doi: 10.2147/AHCT.S138445

11. Guzik P, Malik M. ECG by mobile technologies. J Electrocardiol. (2016) 49:894–901. doi: 10.1016/j.jelectrocard.2016.07.030

12. Nemcova A, Jordanova I, Varecka M, Smisek R, Marsanova L, Smital L, et al. Monitoring of heart rate, blood oxygen saturation, blood pressure using a smartphone. *Biomed Signal Process Control.* (2020) 59:101928. doi: 10.1016/j.bspc.2020.101928

13. Matsumura K, Yamakoshi T. iphysiometer: a new approach for measuring heart rate and normalized pulse volume using only a smartphone. *Behav Res Methods*. (2013) 45:1272–8. doi: 10.3758/s13428-012-0312-z

14. Choudhury AD, Misra A, Pal A, Banerjee R, Ghose A, Visvanathan A. Heartsense: Estimating heart rate from smartphone photoplethysmogram using adaptive filter and interpolation. In *Internet of Things. User-Centric IoT: First International Summit, IoT360 2014*; 2014 Oct 27–28; Rome, Italy. Springer (2015). p. 203–9.

15. Sieciński S, Kostka PS, Tkacz EJ. Gyrocardiography: a review of the definition, history, waveform description, and applications. *Sensors.* (2020) 20:6675. doi: 10. 3390/s20226675

16. [Dataset] Faisal A, Gavriel C. Heart monitoring device and method. U.S. Patent No 10,285,650 (2019).

17. Jafari Tadi M, Lehtonen E, Saraste A, Tuominen J, Koskinen J, Teräs M, et al. Gyrocardiography: a new non-invasive monitoring method for the assessment of cardiac mechanics and the estimation of hemodynamic variables. *Sci Rep.* (2017) 7:1–11. doi: 10.1038/s41598-017-07248-y

18. Pourriahi M, Gurman P, Daich J, Cynamon P, Richler A, Elman N, et al. The use of micro-electro mechanical systems in vascular monitoring: implications for clinical use. *Expert Rev Med Devices*. (2016) 13:831–7. doi: 10.1080/17434440.2016.1207520

19. Lahdenoja O, Hurnanen T, Iftikhar Z, Nieminen S, Knuutila T, Saraste A, et al. Atrial fibrillation detection via accelerometer and gyroscope of a smartphone. *IEEE J Biomed Health Inform.* (2017) 22:108–18. doi: 10.1109/JBHI.2017.2688473

20. Jaakkola J, Jaakkola S, Lahdenoja O, Hurnanen T, Koivisto T, Pänkäälä M, et al. Mobile phone detection of atrial fibrillation with mechanocardiography: the MODE-AF study (mobile phone detection of atrial fibrillation). *Circulation*. (2018) 137:1524–7. doi: 10.1161/CIRCULATIONAHA.117.032804

21. Iftikhar Z, Lahdenoja O, Jafari Tadi M, Hurnanen T, Vasankari T, Kiviniemi T, et al. Multiclass classifier based cardiovascular condition detection using smartphone mechanocardiography. *Sci Rep.* (2018) 8:9344. doi: 10.1038/s41598-018-27683-9

22. Tadi MJ, Mehrang S, Kaisti M, Lahdenoja O, Hurnanen T, Jaakkola J, et al. Comprehensive analysis of cardiogenic vibrations for automated detection of atrial fibrillation using smartphone mechanocardiograms. *IEEE Sens J.* (2018) 19:2230–42 doi: 10.1109/JSEN.2018.2882874.

23. Mehrang S, Lahdenoja O, Kaisti M, Tadi MJ, Hurnanen T, Airola A, et al. Classification of atrial fibrillation and acute decompensated heart failure using smartphone mechanocardiography: a multilabel learning approach. *IEEE Sens J.* (2020) 20:7957–68. doi: 10.1109/JSEN.2020.2981334

24. Hurnanen T, Kaisti M, Tadi MJ, Vähä-Heikkilä M, Nieminen S, Iftikhar Z, et al. Heartbeat detection using multidimensional cardiac motion signals, dynamic balancing. In EMBEC & NBC 2017: Joint Conference of the European Medical and Biological Engineering Conference (EMBEC) and the Nordic-Baltic Conference on Biomedical Engineering and Medical Physics (NBC); 2017 Jun 11-15; Tampere, Finland. Springer (2018). p. 896-9.

25. Buckberg GD. Basic science review: the helix and the heart. J Thorac Cardiovasc Surg. (2002) 124:863–83. doi: 10.1067/mtc.2002.122439

26. Romano C, Schena E, Formica D, Massaroni C. Comparison between chestworn accelerometer and gyroscope performance for heart rate and respiratory rate monitoring. *Biosensors*. (2022) 12:834. doi: 10.3390/bios12100834

27. [Dataset] Meriheinä U, Juppo M, Koivisto T, Pänäälä M, Sairanen K, Grönholm M. *Heart monitoring system*. U.S. Patent No 10,178,964 (2019).

 Mohamed R, Youssef M. Heartsense: ubiquitous accurate multi-modal fusionbased heart rate estimation using smartphones. In *Proceedings of the ACM on Interactive, Mobile, Wearable and Ubiquitous Technologies.* Vol. 1. New York, USA: Association for Computing Machinery (2017). p. 1–18.

29. Bera TK, Gore MM, Oak JP.Recovery from stress in two different postures, in Shavasana—a yogic relaxation posture. *Indian J Physiol Pharmacol.* (1998) 42:473–8.

30. Watanabe N, Reece J, Polus BI. Effects of body position on autonomic regulation of cardiovascular function in young, healthy adults. *Chiropr Osteopathy.* (2007) 15:1–8. doi: 10.1186/1746-1340-15-19

31. MacWilliam J. Postural effects on heart-rate, blood-pressure. *Q J Exp Physiol.* (1933) 23:1–33. doi: 10.1113/expphysiol.1933.sp000588

32. Jiokeng K, Jakllari G, Beylot A-L. Handrate: heart rate monitoring while simply holding a smartphone. In 2021 IEEE International Conference on Pervasive Computing, Communications (PerCom); 2021 Mar 22–26; Kassel, Germany. IEEE (2021). p. 1–11.

33. Hernandez J, McDuff D, Quigley K, Maes P, Picard RW. Wearable motion-based heart rate at rest: a workplace evaluation. *IEEE J Biomed Health Inform*. (2018) 23:1920–7. doi: 10.1109/JBHI.2018.2877484

34. Dar R. Effect of real-time monitoring and notification of smoking episodes on smoking reduction: a pilot study of a novel smoking cessation app. *Nicotine Tob Res.* (2018) 20:1515–8. doi: 10.1093/ntr/ntx223

35. Nair MB, Kumar SR, Kishore NA, Mohan N, Anudev J. Instantaneous feedback pedometer with emergency GPS tracker. In 2018 2nd International Conference on I-SMAC (IoT in Social, Mobile, Analytics and Cloud); 2018 Aug 30–31; Palladam, India. IEEE (2018). p. 122–6.

36. Al-Khafajiy M, Baker T, Chalmers C, Asim M, Kolivand H, Fahim M, et al. Remote health monitoring of elderly through wearable sensors. *Multimed Tools Appl.* (2019) 78:24681–706. doi: 10.1007/s11042-018-7134-7

37. Szanto Z, Benko I, Jakab L, Szalai G, Vereczkei A. The use of a smartphone application for fast lung cancer risk assessment. *Eur J Cardiothorac Surg.* (2017) 51:1171-6. doi: 10.1093/ejcts/ezw444

38. Sahoo PK, Thakkar HK, Lee M-Y. A cardiac early warning system with multi channel SCG and ECG monitoring for mobile health. *Sensors.* (2017) 17:711. doi: 10.3390/s17040711

39. Wannenburg J, Malekian R. Body sensor network for mobile health monitoring, a diagnosis and anticipating system. *IEEE Sens J.* (2015) 15:6839–52. doi: 10.1109/JSEN.2015.2464773

40. Hoang ML, Pietrosanto A. Yaw/heading optimization by drift elimination on MEMS gyroscope. *Sens Actuators A Phys.* (2021) 325:112691. doi: 10.1016/j.sna. 2021.112691

41. Diao Z, Quan H, Lan L, Han Y. Analysis and compensation of MEMS gyroscope drift. In 2013 Seventh International Conference on Sensing Technology (ICST); 2013 Dec 3–5; Wellington, New Zealand. IEEE (2013). p. 592–6.

42. St Pierre SR, Peirlinck M, Kuhl E. Sex matters: a comprehensive comparison of female and male hearts. *Front Physiol.* (2022) 13:303. doi: 10.3389/fphys.2022.831179

43. Karastergiou K, Smith SR, Greenberg AS, Fried SK. Sex differences in human adipose tissues-the biology of pear shape. *Biol Sex Differ*. (2012) 3:1-12. doi: 10. 1186/2042-6410-3-13

44. Shcherbina A, Mattsson CM, Waggott D, Salisbury H, Christle JW, Hastie T, et al. Accuracy in wrist-worn, sensor-based measurements of heart rate and energy expenditure in a diverse cohort. *J Pers Med.* (2017) 7:3. doi: 10.3390/jpm7020003

45. Powell-Wiley TM, Poirier P, Burke LE, Després J-P, Gordon-Larsen P, Lavie CJ, et al. Obesity and cardiovascular disease: a scientific statement from the American Heart Association. *Circulation*. (2021) 143:984–1010. doi: 10.1161/CIR.00000000000973

46. Pouwels S, Topal B, Knook MT, Celik A, Sundbom M, Ribeiro R, et al. Interaction of obesity and atrial fibrillation: an overview of pathophysiology and clinical management. *Expert Rev Cardiovasc Ther.* (2019) 17:209–23. doi: 10.1080/14779072.2019.1581064

47. Yadav RL, Yadav PK, Yadav LK, Agrawal K, Sah SK, Islam MN. Association between obesity and heart rate variability indices: an intuition toward cardiac autonomic alteration—a risk of CVD. *Diabetes Metab Syndr Obes.* (2017) 10:57–64. doi: 10.2147/DMSO.S123935

48. Nuttall FQ. Body mass index: obesity, BMI, and health: a critical review. Nutr Today. (2015) 50:117. doi: 10.1097/NT.00000000000092

49. Agrawal T, Nagueh SF. Changes in cardiac structure and function with aging. J Cardiovasc Aging. (2022) 2:13. doi: 10.20517/jca.2021.40

50. Schenk K, Heinze G. Age-dependent changes of heart valves and heart size. Recent Adv Stud Card Struct Metab. (1975) 10:617–24.

51. Geovanini GR, Vasques ER, de Oliveira Alvim R, Mill JG, Andreão RV, Vasques BK, et al. Age and sex differences in heart rate variability and vagal specific patterns— Baependi Heart Study. *Glob Heart.* (2020) 15(1):71. doi: 10.5334/gh.873

52. Kuk JL, Saunders TJ, Davidson LE, Ross R. Age-related changes in total and regional fat distribution. Ageing Res Rev. (2009) 8:339-48. doi: 10.1016/j.arr.2009.06.001

53. de Oliveira Alvim R, Santos PCJL, Musso MM, de Sá Cunha R, Krieger JE, Mill JG, et al. Impact of diabetes mellitus on arterial stiffness in a representative sample of an urban Brazilian population. *Diabetol Metab Syndr*. (2013). 5, 1–8. doi: 10.1186/1758-5996-5-45

54. Loring SH, Garcia-Jacques M, Malhotra A. Pulmonary characteristics in COPD and mechanisms of increased work of breathing. *J Appl Physiol Respir Environ Exerc Physiol.* (2009) 107:309–14. doi: 10.1152/japplphysiol.00008.2009

55. Guo Q, Wang Y, Xu D, Nossent J, Pavlos NJ, Xu J. Rheumatoid arthritis: pathological mechanisms and modern pharmacologic therapies. *Bone Res.* (2018) 6:15. doi: 10.1038/s41413-018-0016-9

56. Di Bartolomeo S, Alunno A, Carubbi F. Respiratory manifestations in systemic lupus erythematosus. *Pharmaceuticals.* (2021) 14:276. doi: 10.3390/ph14030276

57. Kim SH, Lim KR, Seo J-H, Ryu DR, Lee B-K, Cho B-R, et al. Higher heart rate variability as a predictor of atrial fibrillation in patients with hypertension. *Sci Rep.* (2022) 12:3702. doi: 10.1038/s41598-022-07783-3

58. Guzzetti S, Magatelli R, Borroni E, Mezzetti S. Heart rate variability in chronic heart failure. *Auton Neurosci.* (2001) 90(1-2):102-05. doi: 10.1016/S1566-0702(01) 00274-0

59. Limentani GB, Ringo MC, Ye F, Bergquist ML, McSorley EO. Beyond the *t*-test: statistical equivalence testing. *Anal Chem.* (2005) 77:A221–6. doi: 10.1021/ac053390m

60. Wellek S. Testing statistical hypotheses of equivalence and noninferiority. New York: CRC Press (2010).

61. Dixon PM, Saint-Maurice PF, Kim Y, Hibbing P, Bai Y, Welk GJ. A primer on the use of equivalence testing for evaluating measurement agreement. *Med Sci Sports Exerc.* (2018) 50:837. doi: 10.1249/MSS.00000000001481

62. Bai Y, Welk GJ, Nam YH, Lee JA, Lee J-M, Kim Y, et al. Comparison of consumer and research monitors under semistructured settings. *Med Sci Sports Exerc.* (2016) 48:151–8. doi: 10.1249/MSS.00000000000727

63. Lee J, Kim Y, Welk G. Validity of consumer-based physical activity monitors. *Med Sci Sports Exerc.* (2014) 46:1840–8. doi: 10.1249/MSS.00000000000287