

ReBeatICG: Real-time Low-Complexity Beat-to-beat Impedance Cardiogram Delineation Algorithm

Una Pale, Nathan Müller, Adriana Arza, and David Atienza¹

Abstract—This work presents ReBeatICG, a real-time, low-complexity beat-to-beat impedance cardiography (ICG) delineation algorithm that allows hemodynamic parameters monitoring. The proposed procedure relies only on the ICG signal compared to most algorithms found in the literature that rely on synchronous electrocardiogram signal (ECG) recordings. ReBeatICG was designed with implementation on an ultra-low-power microcontroller (MCU) in mind. The detection accuracy of the developed algorithm is tested against points manually labeled by cardiologists. It achieves a detection Gmean accuracy of 94.9%, 98.6%, 90.3%, and 84.3% for the B, C, X, and O characteristic points, respectively. Furthermore, several hemodynamic parameters were calculated based on annotated characteristic points and compared with values generated from the cardiologists' annotations. ReBeatICG achieved mean error rates of 0.11 ms, 9.72 ms, 8.32 ms, and 3.97% for HR, LVET, IVRT, and relative C-point amplitude, respectively.

I. INTRODUCTION

According to the World Health Organisation (WHO), cardiovascular diseases (CVDs) are globally the highest cause of death. 17.9 million people died from CVDs in 2016, or 31% of all global deaths [1]. Hemodynamic parameters, such as stroke volume or cardiac output, are vital to estimate cardio-respiratory activity and evaluate the subject's condition. Hence, an ambulatory and unobtrusive monitoring of such parameters enables remote CVD monitoring and diagnosis, reduces related hospitalization costs, expands patients' mobility, and improves their quality of life [2].

Widely used noninvasive techniques to obtain the hemodynamic parameters are doppler echocardiography, CO₂ breath analysis, seismocardiography, impedance cardiography (ICG) and phonocardiography [3]. Among them, ICG is a suitable measurement for continuous and real-time monitoring of the hemodynamic function of the cardiovascular system since it is a noninvasive, simple, and low-cost technique. Many useful hemodynamic parameters can be determined from the ICG signal, such as cardiac output (CO), stroke volume (SV), overall vascular resistance, systolic time intervals [4] (e.g., left ventricular ejection time (LVET), pre-ejection period (PEP), systolic time ratio [5]), or thoracic fluid content.

ICG has been used in various applications, including monitoring the cardiac rehabilitation process, monitoring of hemodynamics during hemodialysis, and pharmacological,

physiological, and sleep studies [6], [7]. Moreover, with current advances in wearable technologies and ultra-low-power microcontrollers (MCU), ICG is a promising technique for monitoring hemodynamic parameters and, thus several CVDs [8]. Still, it requires an accurate and real-time detection of the ICG's characteristic points.

However, to the best of our knowledge, no real-time, low-complexity, and beat-to-beat algorithm that can be implemented on an ultra-low-power MCU has been developed yet. Most of the state-of-the-art algorithms are based on the averaging of multiple cardiac cycles technique [4], [9]–[13], which needs the beat-to-beat synchronous reference from the electrocardiogram (ECG) signal. The assemble averaging method allows the removal of movement artifacts, respiratory influences, and stochastic distributed noise, which are the main sources of noise affecting the ICG signal. Although other ICG filtering and delineation approaches have been proposed, they are not suitable to be used in real-time and on a constrained system employing an ultra-low-power MCU.

Therefore, we propose a real-time and low-complexity procedure for the automatic detection of beat-to-beat ICG characteristic points that can be later used on ultra-low-power MCUs. Most of the automatic algorithms found in the literature rely on the synchronously measured ECG signal to detect the prominent peaks of the ICG signal. In contrast, our algorithm relies only on the ICG signal, which has the advantage of reducing the complexity of acquisition and signal processing since no alignment between the two signals is needed. Moreover, our algorithm includes an adaptive filtering stage based on the state-of-the-art Savitzky-Golay filter (SG).

The contributions of this work are therefore as follows:

- ReBeatICG: a new real-time and low-complexity beat-to-beat delineation methodology, relying only on the ICG signal for hemodynamic parameters monitoring on wearable devices.
- An adaptive algorithm for choosing the length of the Savitzky-Golay filter, making our detection more robust to artifacts and baseline shifts.
- Adaptation of an ECG R-peak detection algorithm based on the relative energy method to detect the C points of the ICG, achieving a detection Gmean of 98.6%.
- A novel method to detect the pairs of X and O points, obtaining a detection Gmean of 90.3% and 84.3%, respectively.
- Hemodynamic parameters monitoring, with precision on the level of state-of-the-art algorithms, namely, mean error rates of 0.11 ms, 9.72 ms, 8.32 ms and 3.97% for HR, LVET, IVRT, and relative C-point amplitude,

This work has been partially supported by the ML-Edge Swiss National Science Foundation (NSF) Research project (GA No. 200020182009/1), the PEDESITE Swiss NSF Sinergia project (GA No. SCRSII5 193813/1), and by the ONR-G through the Award Grant No. N62909-20-1-2063.

¹ U. Pale, N. Müller, A. Arza, and D. Atienza are with the Embedded Systems Laboratory (ESL) of Swiss Federal Institute of Technology Lausanne (EPFL), Switzerland. {una.pale, nathan.muller, adriana.arza, david.atienza}@epfl.ch

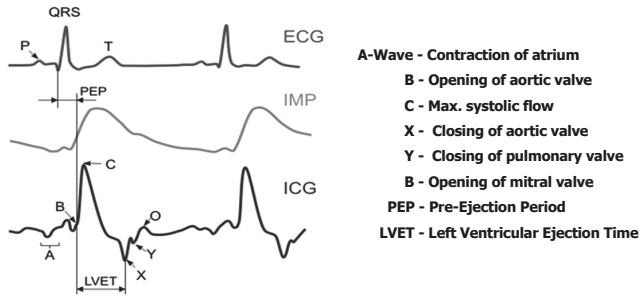


Fig. 1. Left: Electrocardiography (ECG), impedance variation (IMP), and impedance cardiography signal (ICG). Right: Common definition for the ICG characteristic points and hemodynamic parameters [6]

respectively.

- A new open-access database of ICG signals that include 1920 beats fully annotated by a cardiologist used to test our ReBeatICG algorithm.

II. BACKGROUND AND RELATED WORK

A. Impedance Cardiography

Impedance cardiography is a noninvasive technique for measuring changes in the thorax impedance (Z_0) driven by the intrathoracic fluid changes with each heartbeat [14], [15]. Transthoracic impedance is modulated by the cardio-respiratory activity, so that a decrease in impedance is related to an increase in blood flow. ICG is measured by applying a low-amplitude, high-frequency current through two outer electrodes and then acquiring the electrical voltage difference via two inner electrodes. ICG measures dZ/dt , or the first derivative of the impedance signal Z_0 .

1) *Characteristic Points Definition*: The derivative of impedance dZ/dt contains characteristic points related to cardio-dynamic events i.e., A, B, C, X, Y, O [6], [16]. The characteristic points of an ICG signal and its relation with cardio-dynamic events are presented in Fig. 1.

C peak – Defined as the peak with the greatest amplitude in one cardiac cycle that represents the maximum systolic flow.

B point – Indicates the onset of the final rapid upstroke toward the C point [15] that is expressed as the point of significant change in the slope of ICG signal preceding the C point. It is related to the aortic valve opening and is used to calculate the SV and the CO. However, its identification can be difficult due to variations in the ICG signal’s morphology. Thus, different definitions have been proposed [4], [17]–[19].

X point – Often defined as the minimum dZ/dt value in one cardiac cycle. However, this does not always hold true [4] due to variations in the dZ/dt waveform morphology. Thus, the X point is defined as the onset of the steep rise in ICG towards the O point. It represents the aortic valve closing which occurs after the T wave end of the ECG signal.

O point – The highest local maxima in the first half of the C-C interval. It represents the mitral valve opening.

It is important to note that there are many variations in the morphology of the ICG signals [4], [17]. In Fig. 2, locations of B, C and X points on different variations of ICG signal

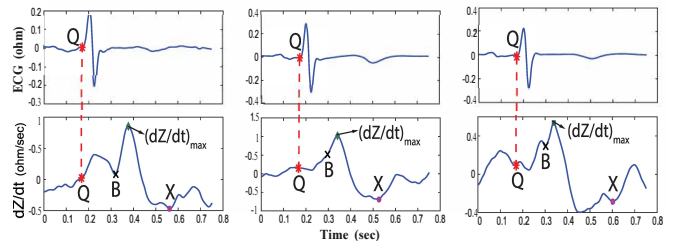


Fig. 2. Three examples of dZ/dt (bottom) with the corresponding ECG signal (top) representing the morphological variations in the dZ/dt signal with its characteristic points presented in [4]. Left case: B and X points occur at a local minimum and global minimum, respectively; Middle case: the B point is a notch; Right case: the X point is only a local minimum, not a global minimum.

morphology can be observed. Thus, defining algorithms and detecting points on ICG signals is quite challenging.

2) *Hemodynamic Parameters*: Various hemodynamic parameters can be estimated from the ICG’s characteristic points. The heart rate (HR) can be measured as the time between two consequent C peaks. The Left Ventricular Ejection Time (LVET) is the time interval measured from the B point to the X point, and can be used to estimate the stroke volume (SV) and the cardiac output (CO). The stroke volume echoes the volume of blood pumped by the left ventricle at each contraction, while the cardiac output is the total volume of blood pumped by the ventricle, usually per minute [20]. The isovolumetric relaxation time (IVRT) is the time interval measured from the X point to the O point. The amplitude of the C peak (denoted dZ/dt_{max}) is commonly defined with respect to the amplitude of the corresponding B point, and is as well necessary to calculate stroke volume and cardiac output.

B. State-of-the-art ICG Preprocessing and Filtering Methods

The most important artifacts affecting the ICG signal are sinusoidal, respiratory, muscle, and electrode artifacts [21]. Different preprocessing and filtering methods have been proposed in literature [13], [16], which vary from simple methods such as ensemble averaging of multiple cardiac cycles [15], [22] to more advanced ones such as adaptive noise cancellation [21], [23] and wavelet analysis [24], [25]. Among them, ensemble averaging is the most commonly used method since it eliminates stochastically distributed noise as well as respiratory influences and movement artifacts [4], [9], [10]. However, averaging several cardiac cycles tends to blur less distinctive events, such as the B point, making its detection more difficult. Besides, it precludes beat-to-beat analysis of cardiac hemodynamics parameters.

More complex techniques but also less suitable for real-time implementation on wearable devices have been proposed. For instance, adaptive filtering techniques that update the filter weights according to the statistical nature of the error signal, such as Least Mean Square and Recursive Least Square algorithms [26], or hybrid algorithms that have been developed to overcome the weight drift and instability of adaptive filtering algorithms [21] were tested.

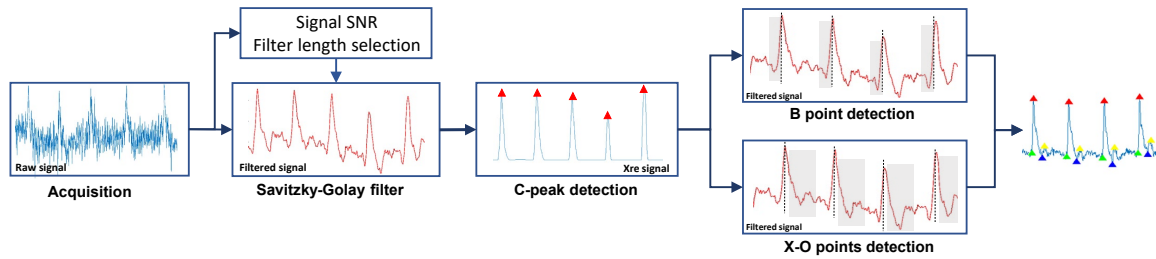


Fig. 3. Workflow of our proposed ReBeatICG algorithm

In [13], common filtering techniques are compared, i.e., the Savitzky-Golay (SG) filter, median filter, wavelet, band-pass filter, and moving average filter. When comparing with a high-pass filter, ensemble average, and spline interpolation, the SG filter is the best option to remove respiration artifacts while preserving the beat-to-beat information [10]. Similarly, [27] shows that the SG filter is the most relevant for denoising ICG signals with better signal shape preservation, particularly the C peak amplitudes. Since the SG filter usage is very simple and efficient [28], we selected it for our methodology.

C. State-of-the-art ICG Delineation Methods

After a filtering preprocessing step, the signal is segmented, usually into beat-to-beat intervals to extract the characteristic points. Most of the available methods, especially those using an assembling averaging step, utilize the R-peak from the ECG signal to define the beat-to-beat time interval in which to search for C peak, defined as the highest point of ICG signal. Many of them apply known R-peak detection algorithms (e.g., the Pan-Tompkins algorithm [28], [29]) to detect the C point due to the similarity between ECG and ICG signals.

For detection of B, X, and O points, and to address its morphological variations (see Fig. 2), several conditions based on zero crossings, minima, and maxima of the dZ/dt signal and its derivatives have been proposed [4], [7], [11], [15], [29], [30]. Other algorithms employ time-frequency [31], wavelet [22], or machine learning-based and data-driven polynomial approaches [12]. However, they are computationally expensive with respect to the condition-based approaches. In addition, particularly time-frequency and wavelet approaches are sensitive to any artifacts that have their frequency content overlapping with the dZ/dt frequency content around the characteristic points.

As the precise identification of the B point is paramount to ensuring accurate computation of SV and CO [8], numerous acquisition methods have been proposed in the literature [29]. The least computationally complex methods rely on derivatives analysis, but they are sensitive to noise and artifacts and may fail when there are several reversals, inflections, and rapid slope changes on the signal [15], [30]. To overcome this, in [12] and [29], a method is presented based on the derivatives, but, applying several conditions and corrections according to the morphology of the signal. For instance, in [29], the B point is detected as the sign change within 20% and 65% of the R–C interval window of the ICG first derivative. On the other hand, in [12], the B point is found

on the most prominent monotonically increasing segments between the A and C points.

X and O points detection methods are less prominent in the literature. They range from complex ones [22], [31] (e.g., time-frequency analysis and wavelet-based) to simple condition checking. The simplest approach to detect the X point is as the first minimum after C wave [13], [28], [32] or the lowest minimum within a given interval (i.e., in the first one-third of the C–C interval) [7]. However, since the X point is defined as the onset of steep rise towards the O point, some algorithms apply different conditions to detect the X–O pair [11], [29].

Although several ICG delineation algorithms have been proposed, there is no complete proposal for low-complex and accurate delineation of B, C, X and O points that can be later implemented in a real-time wearable system. Furthermore, there are no standard evaluation metrics nor databases that allow the assessment and comparison with previously proposed methods.

III. PROPOSED REBEATICG ALGORITHM

Our ReBeatICG ICG delineation algorithm was developed for real-time, beat-to-beat hemodynamics parameters monitoring, which detects B, C, X and O points only relying on the ICG signal. We took advantage of many of the ideas from existing ICG delineation algorithms to improve our algorithm performance while selecting the least complex methods for a later implementation on a lightweight wearable device. The general workflow of ReBeatICG is presented in Fig. 3.

First, during the acquisition, the signal is segmented into windows of 3 seconds. The segments are filtered by applying the SG filter with a length adaptively selected based on the Signal to Noise Ratio (SNRs). Next, C points are detected using a novel procedure based on the R-peak REWARD algorithm [33] and designed to be implemented on ultra-low-power MCUs. Then, within the C–C intervals the B point is detected, followed by the delineation of the X–O pair, employing in both steps a derivative analysis together with condition checking following the signal morphology.

The selected methods provide reliable detection while ensuring the low complexity of our method. Finally, in order to have a real-time, continuous beat-to-beat analysis, the last O point found in the 3s-segmented window processed is used as the beginning of the next segmentation window.

A. ICG Preprocessing using the SG filter

To filter the ICG signal, we use the SG filter, as it is a simple and low complexity filter that uses a polynomial approach by applying a fitting with the least-squares method. However, to properly denoise the signal, the polynomial degree should be selected. Authors in [13], [28] found that the SG filter with a polynomial of order 3 has the lowest error rate for different ICG Signal to Noise Ratios (SNRs). Further, the filter length ("the smoothing window") strongly influences filter effectiveness. Hence, selecting the appropriate filter length is essential for a correct trade-off between reducing noise and retaining important signal details needed to successfully annotate B, C, X and O points.

Adaptive SG filtering has proved its robustness and signal preservation in other applications [34], [35], and could be very useful for ICG's high SNR variability due to inter-subject variability, electrode placement, and various artifact sources. Therefore, we propose to adaptively select the filter length by analyzing its impact on the SNR of each 3s-signal window.

Starting from a filter length of 3, we increase the length in steps of two until signal SNR reaches a target SNR_{thr} , or if the SNR does not improve by more than threshold $SNR_{imprThr}$. By observing how the SNR saturates above a certain filter length, SNR_{thr} and $SNR_{imprThr}$ values of 30 and 1% were chosen. Moreover, these values present a good compromise between reducing noise and maintaining a low filter length, thus reducing complexity, as well as avoiding over-smoothing of the signal (and hence potentially losing valuable details). The SNR is calculated as a ratio between the 2-norm of the high and low signal frequencies considering a 20Hz as cut-off frequency.

B. Detection of the C peaks

The detection of the C peaks is the most important step in the detection procedure, since the other points will be located in time windows before and after the C peaks. We have developed a new C-point detection procedure based on the "relative energy" (Rel-En) preprocessing method [36] and an adaptation of the REWARD algorithm [33] designed to find the R peaks in the ECG.

First, the Rel-En preprocessing method is applied to the filtered dZ/dt signal to enhance the C peaks. The Rel-En method considers the energies of a long sliding window (0.95s) and a short sliding window (0.14s); both centered at sample n . The ratio between the energies of these windows, the coefficient $c(n)$, is multiplied by $signal(n)$ resulting in a signal $X_{re}(n)$, in which the peaks are amplified.

The C peaks are then searched in the X_{re} signal using two adaptive thresholds defined as a percentage of Max_{val} in the window. Max_{val} is defined as the peak amplitude value of the second biggest peak if it is significantly bigger than the mean value of the signal, otherwise the peak value of the biggest peak; hence, providing robustness to the noise and outliers.

Next, segments in which the X_{re} signal goes above the upper threshold, $Thr_{max}=0.2Max_{val}$, and subsequently

Algorithm 1 B Point Detection

Input: Signal Window defined by C points

Output: B point

```

1: def find_B_point( $sig, C_{pos}, C_{Ampl}, A_{frac}, B_{slope1}, B_{slope2}$ ):
2:    $B_{limL} = C_{pos} - 80ms * Fs$ 
3:    $B_{limR} = \mathbf{find} ( sig[i] \leq A_{frac}C_{Ampl} )$ 
4:    $B_{min} = \mathbf{min}(sig)$ 
5:   for  $i = B_{limR}; i \neq B_{limL}; i --$  do
6:     if  $sig[i] == \mathbf{localMin}()$  or  $|sig'[i]| > B_{slope1}$  then
7:       return  $i$ 
8:   for  $i = B_{limR}; i \neq B_{limL}; i --$  do
9:     if  $sig'[i] > B_{slope2}$  then
10:      return  $i$ 
11:  return  $B_{min}$ 

```

below the lower threshold, $Thr_{min}=0.02Max_{val}$, are considered active peak regions, similar to those in the REWARD approach [33]. The maximum of the signal between these two points is considered a C peak candidate. Once the peak candidates have been determined, if the peak to peak interval between two peaks is less than 0.25 seconds, only the highest of the two is kept. Further, the peaks are valid if the C-C time interval is not smaller than $CC_{min}=1.7$ times the average of the five previous C-C intervals. This condition is necessary to prevent the algorithm from detecting an O point as a C peak when, occasionally, the O point has an amplitude comparable to the one of the C peak.

C. Detection of the B points

B point identification can be difficult due to variations in the morphology of the ICG signals (see Fig. 2). Algorithm 1 describes our method of acquiring the B point. First, we define the time window where a B point can possibly be $[B_{limL}, B_{limR}]$ (lines 2-3). B_{limL} is defined as $C_{pos}-80ms$ [4]. B_{limR} is defined as the closest point before a C point where the signal amplitude is less than a fraction of C amplitude ($A_{frac}=0.5$). Next, we search for either the local minimum closest to C_{pos} or the first point at which the slope of the signal exceeds threshold $B_{slope1}=0.11$ (lines 4-7). If B_{limL} is reached before finding a B point, the search is repeated with a less strict slope threshold $B_{slope2}=0.08$ (lines 8-10). If a suitable B point is still not found, a minimum B_{min} is used for the B-point.

Parameters values of A_{frac} , B_{slope1} and B_{slope2} were obtained by performing grid search for optimal parameters. For this, we randomly selected 8 blocks of 10 beats from each subject of the database and annotated by ourselves following the definitions from II-A.

D. Detection of the X and O points

The X point is defined as the onset of the steep rise in the ICG towards the O point, which is the highest local maxima in the first half of the C-C interval. Based on this definition, we follow a new procedure on which we look for possible pairs of X-O points and select the best candidate. Candidates for O points are local maxima in a time window $[C_{pos}+CO_{min}, C_{pos}+CO_{max}]$. Similarly, candidates for X points are local minima in a window $[C_{pos} + CX_{min}, C_{pos} + CX_{max}]$.

Next, we look for the combinations of X-O pair candidates that meet the following criteria:

- O amplitude is higher than X amplitude,
- the X-O time is within [2-15] ms range,
- there are less than 3 local minimums between O and X.

Finally, the O and X pair with the highest amplitude difference is selected as O and X points. Parameters values were also obtained by a grid search as in B point detection. We used $CO_{min}=20$ ms, $CO_{max}=40$ ms, $CX_{min}=15$ ms, $CX_{max}=30$ ms values.

IV. EXPERIMENTAL SETUP

To assess our algorithm, we use a database manually annotated by cardiologists for the purpose of evaluating the ICG annotation algorithms. The database is published as an open-access database [37]. It is described below, together with the used performance metrics.

A. Experimental Database

The database includes 48 recordings of ICG and ECG signals from 24 healthy subjects during an experimental session of a virtual search and rescue mission with drones, described in [38]. Two segments of 5-minute signals are selected from the first day of the reported experiment protocol from each subject. The segments correspond to a baseline state (task BL) and a higher level of cognitive workload (task CW). In total, the recorded database consists of 240 minutes of ICG signals.

In order to assess the performance, a subset of the database was annotated by cardiologists from the Lausanne University Hospital, which served as a test set for our algorithms. The annotation was done using an open access physiological signal labeler software [39]. The subset consists of 4 blocks of randomly chosen signal segments containing 10 beats from BL and CW tasks of each subject. In total, 1920 (80x24) beats were annotated, each containing annotated B, C, X and O positions.

B. Performance Assessment

To assess the performance of our proposed algorithm, the true positives (TPs), false positives (FPs), and false negatives (FNs) detected points by our algorithm were compared against the aforementioned annotations from the cardiologists. We used a tolerance of ± 30 ms from the annotated point [29]. Accordingly, we use the performance metrics of sensitivity (SE), positive prediction value (PPV), detection error rate (DER), geometric mean (Gmean), mean error (me), and its standard deviation (σ).

Moreover, we calculated several hemodynamic parameters, both from annotation points by the cardiologists as well as from our proposed algorithm. Four hemodynamic parameters were calculated:

- 1) CC_{time} interval: Related to the heart rate (HR), calculated as the interval between subsequent C points;
- 2) $LVET$: Left ventricular ejection time, calculated as the time between the B and X point of the same beat;

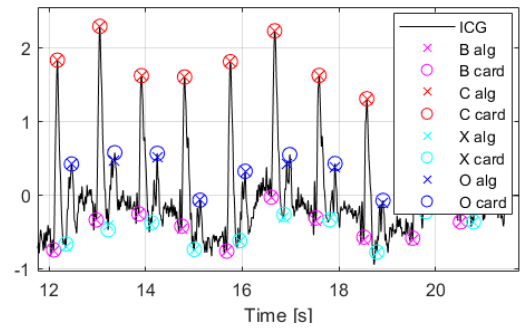


Fig. 4. Example of one block of data including ReBeatICG automatic (X) and manual cardiologists' (O) annotation

- 3) $IVRT$: Isovolumetric relaxation time, calculated as the time between the X and O point of the same beat;
- 4) BC_{ampl} : Relative amplitude of C peak, which is essential for calculating stroke volume and cardiac output.

The precision of hemodynamic parameters was measured by first matching the C peaks between automatic C peak detection and the cardiologists' annotation. Then we calculate the parameters for each of them (e.g. $LVET_{card}$ and $LVET_{alg}$). Next, the absolute error values for all matched beats were calculated and averaged over beats and subjects and parameters. For BC_{ampl} , the relative error was calculated ($abs(BC_{ampl}_{card} - BC_{ampl}_{alg}) / BC_{ampl}_{card}$).

V. EXPERIMENTAL RESULTS

The detection performance results of our automatic ReBeatICG delineation for each B, C, X and O point individually, are shown in Table II. These results represent the performance of ReBeatICG using as reference the beats annotated by cardiologist with ± 30 ms resolution for the matching window. Furthermore, a visual example of ReBeatICG delineation and the reference annotation on one segment of signal is shown in Fig. 4. ReBeatICG C point delineation is highly accurate, while other points delineation are quite precise as well. B point delineation is challenging due to the fact that often the correct location is defined by a slope change and not a minimum, and slope change is very sensitive to noise and filtering. Similarly, X and O points delineation is challenging due to many local minima and maxima sometimes present in the signal. From this reason, several conditions were necessary for identifying the correct X-O pair. Nevertheless, the ReBeatICG algorithm performs very well for the B, C, X, and O points, yielding Gmean performance of 94.9%, 98.6%, 90.3% and 84.3% for each respective point.

A comparison with the existing algorithms for delineating B, C, X or O points is difficult due to different approaches used for validation. For instance, some papers compare hemodynamic parameters calculated from delineated ICG and those from echocardiogram [29], others in reference to ECG R peak detection [30], while others compare annotation positions with manual annotations [4], [19]. Unfortunately, almost no paper clearly mentions the resolution used for matching annotated points window, making the results hardly comparable. Further,

TABLE I
COMPARISON OF REBEATICG WITH STATE-OF-THE-ART METHODS

Ref.	B Performance measures				C Performance measures				X Performance measures			
	SE [%]	PPV [%]	DER [%]	$me \pm \sigma$ [ms]	SE [%]	PPV [%]	DER [%]	$me \pm \sigma$ [ms]	SE [%]	PPV [%]	DER [%]	$me \pm \sigma$ [ms]
[29]	–	–	1.7	2 ± 10.5	–	–	–	5 ± 10	–	–	–	$4 \pm 44^*$
[22]	–	–	–	3.98 ± 2.85	–	–	–	–	–	–	–	23.7 ± 14.9
[19]	94.4	93.9	11.7	–	99.4	98.7	1.8	–	97.0	96.5	6.5	–
[13]	–	93.1	6.9	–	–	100.0	0.0	–	–	99.5	0.5	–
This work	99.04	98.13	2.92	2.03 ± 2.23	99.25	98.28	2.56	0.17 ± 0.73	98.99	98.43	2.66	3.29 ± 4.45

SE-Sensitivity. PPV- Positive Prediction Value, DER- Detection error rate. ME-mean error and SD- standard deviation of error relative to the reference considering ± 150 ms of tolerance, as in [13] and [19], from the annotated point. * Respect to Doppler echocardiogram references because of significant differences across the X point definitions in the literature.

TABLE II
PERFORMANCE OF REBEATICG PER ANNOTATED POINTS

Annot. Points	Performance measures			
	SE [%]	PPV [%]	Gmean [%]	$me \pm \sigma$ [ms]
B	95.30 ± 5.65	94.48 ± 6.96	94.88 ± 6.28	1.75 ± 0.90
C	99.09 ± 1.86	98.13 ± 3.44	98.60 ± 2.50	0.12 ± 0.08
X	90.55 ± 9.51	90.06 ± 9.82	90.30 ± 9.58	1.09 ± 0.35
O	84.58 ± 15.45	84.08 ± 15.45	84.32 ± 15.39	1.31 ± 0.22

The tolerance in respect to the reference values is ± 30 ms

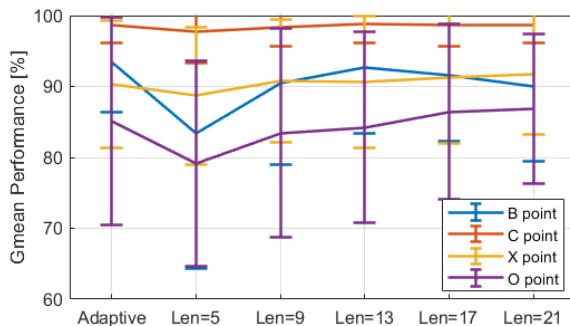


Fig. 5. Performance gain for our adaptive SG filter length versus fixed filter lengths for all annotated points

the performance metrics are also different between papers. Nevertheless, we summarize the results from previous works in Table I. Most of the papers used a resolution of ± 150 ms based on [40] so we also evaluated our results with this resolution and include the results in Table I. It is visible that our ReBeatICG algorithm delineates all characteristic points better or on a comparable level with previous works. Among all performance measures in the table, we consider mean error me [ms] to be the most reliable performance measure for comparison with different papers since it is least influenced by resolution and the one influencing the hemodynamics parameters the most.

Furthermore, in order to assess our adaptive filter length proposal for the SG filter providing a performance gain over a fixed length, we compare the B, C, X and O points detection performance for several fixed filter lengths versus an adaptive one as shown in Fig. 5. The adaptive filter length leads to values from 5 to 25, with the mean length of 12.5 ± 5.1 over all subject data, providing better overall performance compared to equivalently sized fixed length filters.

Finally, we assessed the precision of hemodynamic parameters calculated by ReBeatICG and data annotated by cardiologists. The mean error and standard deviations of absolute and relative values are expressed in Table III. Heart

TABLE III

QUALITY OF AUTOMATIC CALCULATION OF HEMODYNAMIC PARAMETERS

Parameter	HR	LVET	IVRT	BCampl
Mean absolute error [ms]	0.11 ± 0.54	9.7 ± 4.7	8.3 ± 9.4	
Mean relative error [%]	0.01 ± 0.04	3.6 ± 1.7	10.2 ± 11.0	3.9 ± 6.5

rate (HR) is the most precise, with a mean error of 0.11 ms. This result is due to the fact that it depends only on C-points, whose automatic annotation was highly precise. Furthermore, LVET and IVRT are less precise, namely, having a mean error of 9.72 ms and 8.32 ms, respectively, but still well within the 30 ms resolution. Lastly, the relative C peak amplitude was detected with only a 3.97% error with respect to C peak amplitude from annotated data by cardiologists.

We have selected and implemented the least complex methods for a real-time, beat-to-beat hemodynamics parameters monitoring in view of a later implementation on a lightweight wearable device. The above presented results are very promising and motivate future implementation on the low-power microcontroller, but before potential wearable application aspects of latency, energy consumption and memory will have to be analysed when porting to MCU.

VI. CONCLUSION

In this paper, we have presented ReBeatICG, a new real-time and low-complexity algorithm for beat-to-beat delineation based only on the ICG signal. In particular, the algorithm delineates 4 characteristic points, B, C, X and O whose locations are later used for calculation of hemodynamic parameters. To the best of our knowledge, this is first paper including the delineation of all the principal characteristic points of the ICG for hemodynamic parameters monitoring, that allows a later implementation on ultra-low-power MCU.

Our algorithm employed three main state-of-the-art techniques that have both high reliability and low complexity. First, to filter the signal, we employed the Savitzky-Golay filter that is easily implemented as a convolution of the signal with the filter's coefficients, which are integers calculated offline. Second, we used the relative energy method for a highly precise C-peak detection allowing further beat-to-beat delineatio that has been previously implemented on low-power MCU. Finally, for the delineation of B, X and O points we used the derivative analysis with conditioning checking based on time-intervals and signal-amplitude, being a low-complexity method that has also shown high performance.

The automatic detection procedure was compared against the annotated points provided by cardiologists and yielded Gmean performance of 94.9%, 98.6%, 90.3%, and 84.3% for B, C, X, and O points, respectively. Finally, we have assessed the precision of several hemodynamic parameters, obtaining mean errors of 0.11 ms, 9.72 ms, 8.32 ms, and 3.97% for HR, LVET, IVRT, and relative C-point amplitude, respectively. These overall results indicate that our proposed algorithm is highly precise, and could be implemented on a low-power wearable device for hemodynamic parameter monitoring.

ACKNOWLEDGMENT

We especially want to thank Dr. David Meier and Prof. Olivier Muller for all the time and expertise in annotating the ICG database on which we evaluated the ReBeatICG algorithm.

REFERENCES

- [1] "WHO: Cardiovascular diseases, Retrieved from: <https://www.who.int/news-room/fact-sheets/detail/cardiovascular-diseases-cvds>," 2017.
- [2] A. Singhal and M. R. Cowie, "The Role of Wearables in Heart Failure," *Current Heart Failure Reports*, vol. 17, no. 4, pp. 125–132, 2020.
- [3] P. Dehkordi *et al.*, "Comparison of Different Methods for Estimating Cardiac Timings: A Comprehensive Multimodal Echocardiography Investigation," *Frontiers in Physiology*, vol. 10, p. 1057, 8 2019.
- [4] M. Nabian *et al.*, "An open-source feature extraction tool for the analysis of peripheral physiological data," *IEEE Journal of Translational Engineering in Health and Medicine*, vol. 6, 2018.
- [5] A. Hafid *et al.*, "Full Impedance Cardiography Measurement Device Using Raspberry PI3 and System-on-Chip Biomedical Instrumentation Solutions," *IEEE Journal of Biomedical and Health Informatics*, vol. 22, no. 6, pp. 1883–1894, 11 2018.
- [6] H. Yazdani *et al.*, "Design and implementation of a portable impedance cardiography system for noninvasive stroke volume monitoring," *Journal of Medical Signals and Sensors*, vol. 6, no. 1, pp. 47–56, 1 2016.
- [7] M. Forouzanfar *et al.*, "Automatic analysis of pre-ejection period during sleep using impedance cardiogram," *Psychophysiology*, vol. 56, no. 7, p. e13355, 7 2019.
- [8] S. Mansouri *et al.*, "Impedance cardiography: Recent applications and developments," *Biomedical Research (India)*, vol. 29, no. 19, pp. 3542–3552, 2018.
- [9] S. A. Ali Sheikh *et al.*, "An open-source automated algorithm for removal of noisy beats for accurate impedance cardiogram analysis," *Physiological measurement*, vol. 41, no. 7, p. 075002, 8 2020.
- [10] U. Moissl *et al.*, "Filtering respiration in impedance cardiography," *IFAC Proceedings Volumes*, vol. 36, no. 15, pp. 371–376, 8 2003.
- [11] D. Sopic *et al.*, "Touch-Based System for Beat-to-Beat Impedance Cardiogram Acquisition and Hemodynamic Parameters Estimation," in *2016 Design, Automation Test in Europe Conference Exhibition (DATE)*. IEEE, 2016, pp. 726–731.
- [12] M. Forouzanfar *et al.*, "Toward a better noninvasive assessment of preejection period: A novel automatic algorithm for B-point detection and correction on thoracic impedance cardiogram," *Psychophysiology*, vol. 55, no. 8, p. e13072, 8 2018.
- [13] I. B. Salah and K. Ouni, "Denosing of the impedance cardiographic signal (ICG) for a best detection of the characteristic points," in *BioSMART 2017 - Proceedings: 2nd International Conference on Bio-Engineering for Smart Technologies*. Institute of Electrical and Electronics Engineers Inc., 11 2017.
- [14] R. P. Patterson, "Fundamentals of Impedance Cardiography," *IEEE Engineering in Medicine and Biology Magazine*, vol. 8, no. 1, pp. 35–38, 1989.
- [15] A. Sherwood(Chair) *et al.*, "Methodological Guidelines for Impedance Cardiography," *Psychophysiology*, vol. 27, no. 1, pp. 1–23, 1 1990.
- [16] N.-H. David *et al.*, "Fundamentals, recent advances, and future challenges in bioimpedance devices for healthcare applications," 2019.
- [17] S. Benouar *et al.*, "Systematic variability in ICG recordings results in ICG complex subtypes – steps towards the enhancement of ICG characterization," *Journal of Electrical Bioimpedance*, vol. 9, no. 1, pp. 72–82, 2018.
- [18] P. De Carvalho *et al.*, "Robust Characteristic Points for ICG-Definition and Comparative Analysis Proteostasis and mitochondrial function in epicardial adipose tissue View project CONTROLLO 2018-13th APCA International Conference on Control and Soft Computing View project Robust C," Tech. Rep., 2011.
- [19] S. M. Naidu *et al.*, "Automatic detection of characteristic points in impedance cardiogram," in *Computing in Cardiology*, vol. 38, 2011, pp. 497–500.
- [20] M. C. Ipate *et al.*, "The stroke volume and the cardiac output by the impedance cardiography," *UPB Scientific Bulletin, Series C: Electrical Engineering*, vol. 74, no. 3, pp. 219–232, 2012.
- [21] U. R. Zia *et al.*, "Adaptive Noise Cancellation Techniques for Impedance Cardiography Signal Analysis," *International Journal of Innovative Technology and Exploring Engineering (IJITEE)*, vol. 8, 2019.
- [22] L. Y. Shyu *et al.*, "The detection of impedance cardiogram characteristic points using wavelet transform," *Computers in Biology and Medicine*, vol. 34, no. 2, pp. 165–175, 3 2004.
- [23] S. S. Mirza and M. Z. U. Rahman, "Efficient adaptive filtering techniques for thoracic electrical bio-impedance analysis in health care systems," *Journal of Medical Imaging and Health Informatics*, vol. 7, no. 6, pp. 1126–1138, 10 2017.
- [24] R. Stepanov *et al.*, "Wavelet analysis in impedance rheocardiography," in *Bioimpedance in Biomedical Applications and Research*. Springer International Publishing, 3 2018, pp. 257–269.
- [25] S. Liu *et al.*, "Study on cardiac impedance signal feature point extraction," in *Proceedings of 2017 IEEE 3rd Information Technology and Mechatronics Engineering Conference, ITOEC 2017*, vol. 2017-Janua, 11 2017, pp. 790–793.
- [26] O. Dromer *et al.*, "Impedance cardiography filtering using scale fourier linear combiner based on RLS algorithm," in *Proceedings of the 31st Annual International Conference of the IEEE Engineering in Medicine and Biology Society*, 2009, pp. 6930–6933.
- [27] J. Luo *et al.*, "Properties of Savitzky-Golay digital differentiators," *Digital Signal Processing: A Review Journal*, vol. 15, no. 2, pp. 122–136, 3 2005.
- [28] I. B. Salah *et al.*, "Automatic diagnosis of valvular heart diseases by impedance cardiography signal processing," *Biomedical Signal Processing and Control*, vol. 57, p. 101758, 3 2020.
- [29] U. R. Bagal *et al.*, "Detection of opening and closing of the aortic valve using impedance cardiography and its validation by echocardiography," vol. 4, no. 1, 11 2018.
- [30] J. R. Árbol *et al.*, "Mathematical detection of aortic valve opening (B point) in impedance cardiography: A comparison of three popular algorithms," *Psychophysiology*, vol. 54, no. 3, pp. 350–357, 3 2017.
- [31] X. Wang *et al.*, "An Advanced Signal Processing Technique for Impedance Cardiography," *IEEE Transactions on Biomedical Engineering*, vol. 42, no. 2, pp. 224–230, 1995.
- [32] S. Yazdani and J.-M. Vesin, "A Novel Preprocessing Tool to Enhance ECG R-wave Extraction," 2016.
- [33] L. Orlandic *et al.*, "REWARD: Design, Optimization, and Evaluation of a Real-Time Relative-Energy Wearable R-Peak Detection Algorithm," in *Proceedings of the Annual International Conference of the IEEE Engineering in Medicine and Biology Society, EMBS*, 2019, pp. 3341–3347.
- [34] J. Li *et al.*, "Real-time infrared gas detection based on an adaptive Savitzky-Golay algorithm," *Applied Physics B: Lasers and Optics*, vol. 120, no. 2, pp. 207–216, 8 2015.
- [35] D. Acharya *et al.*, "Application of adaptive Savitzky-Golay filter for EEG signal processing," *Perspectives in Science*, vol. 8, pp. 677–679, 9 2016.
- [36] S. Yazdani *et al.*, "A Novel Short-Term Event Extraction Algorithm for Biomedical Signals," *IEEE Transactions on Biomedical Engineering*, vol. 65, no. 4, pp. 754–762, Apr. 2018.
- [37] U. Pale *et al.*, "Rebeat icg database," Apr. 2021.
- [38] F. Dell'Agnola *et al.*, "Cognitive workload monitoring in virtual reality based rescue missions with drones," in *12th International Conference on Virtual, Augmented and Mixed Reality, Copenhagen*, 7 2020.
- [39] S. Zanolli *et al.*, "Physiological signal labeler," Apr. 2021.
- [40] AAMI, *Testing and reporting performance results of cardiac rhythm and ST-segment measurement algorithms*, 2008.