

A Deep Learning Approach for Gait Event Detection from a Single Shank-Worn IMU: Validation in Healthy and Neurological Cohorts

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Abstract: Many algorithms use 3D accelerometer and/or gyroscope data from IMU sensors to detect gait events (i.e., initial and final contact). However, these algorithms often require knowledge about sensor orientation and use empirically derived thresholds. As alignment cannot always be controlled for in ambulatory assessments, methods are needed that require little knowledge on sensor location and orientation, e.g., a convolutional neural network-based deep learning model. Therefore, 160 participants from healthy and neurologically diseased cohorts walked 5 meter distances at slow, preferred, and fast walking speed, while data were collected from IMUs on the left and right ankle and shank. Gait events were detected and stride parameters were extracted using a deep learning model and an optoelectronic motion capture (OMC) system for reference. Results showed a high detection rate for both initial and final contact detection across the IMUs (recall $\geq 92\%$, precision $\geq 97\%$). Time agreement was excellent, as witnessed median time error (0.005 s) and corresponding inter-quartile range (0.020 s). The extracted stride-specific parameters were in good agreement with parameters derived from the OMC system (maximum mean difference 0.003 s and corresponding maximum limits of agreement (-0.049, 0.051) s for 95% confidence level). Thus, the deep learning approach was considered a valid approach for detecting gait events and extracting stride-specific parameters with little knowledge on exact IMU location and orientation.

Keywords: gait; inertial measurement units; deep learning

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1. Introduction

Gait deficits are common in older adults and possibly reflect the presence of an underlying neurodegenerative disease [1,2]. For example, conversion to Parkinson's Disease [3] or from mild cognitive impairment to Alzheimer's Disease [4,5] are linked with changes in spatiotemporal gait parameters. Similarly, temporal gait parameters are different for stroke patients [6,7] and patients with multiple sclerosis [8,9] when compared to healthy controls. To objectively quantify gait deficits, stride-specific parameters such as stride time or stride length are often used [10], where the beginning and end of a stride are determined from two successive initial contacts (ICs) of the same foot [11,12]. The IC is when the foot contacts the ground and together with the instant at which the foot leaves the ground (final contact, FC), each stride can be divided in a stance and swing phase [13,14]. The events of IC and FC, also referred to as *gait events*, are commonly determined using force or pressure measuring devices [14], or passive, retro-reflective marker-based optoelectronic motion capture systems (henceforth referred to as the *marker-based* system or method) [15,16]. These systems are relatively expensive, and restricted to usage in expertise laboratories [17,18]. As there is increasing evidence that gait measured in the lab does not reflect daily-life gait [19–21], there is more and more interest in measurement systems that

allow for continuous gait analysis in ambulatory settings. Therefore, the use of inertial measurement units (IMUs) is especially attractive, as these can be used to measure gait in ecologically valid environments, such as the home environment, thereby painting a more complete picture of health status [22,23] and providing clinical information that is complementary to standardized lab-based assessments [20,21,24,25].

Previous research suggests that gait event detection is more accurate using an IMU worn on a lower limb (e.g., shank or foot) compared to an IMU worn on the low back [26–28]. Now, in order to get from abstract IMU sensor readings to clinically relevant gait parameters (e.g., from accelerations and angular velocities to stride times) [10], different algorithmic approaches have been developed in the last twenty years of clinical gait research. A recent study has evaluated a cross-section of these algorithms for different sensor locations on the lower leg and foot [29]. The algorithms were categorized according to which signals were analyzed, for example the angular velocity about the medio-lateral axis, or the accelerations along vertical and antero-posterior axis. This means the sensor readings need to be linked with the anatomical axes, that is, one needs to know which sensor axis aligns with for example the medio-lateral axis. In most approaches, it is simply assumed that due to sensor attachment the sensor axis aligns roughly with the anatomical axis of interest ([30–36]) or an additional calibration procedure (e.g., [37]) is required ([29,38]). In ambulatory assessments however, study participants often attach the sensor themselves, and therefore the sensor location and alignment cannot be controlled for. Furthermore, it is unlikely that each time the sensor is (re-)attached study participants, especially those with gait deficits, perform a calibration procedure that usually consists of holding a pre-defined pose and performing some known movement sequences (TODO: reference?! Kong? Seel?).

Taken together, this drives the need for an approach that is invariant to sensor orientation, and is applicable across a variety of pathological gait patterns. As pointed out by [10], in the field of image analysis similar requirements have been successfully addressed by algorithms that share a common underlying methodology referred to as deep learning [39,40]. Recent applications of deep learning algorithms have already shown improved performance in detecting gait events from marker-based motion capture when compared to conventional, often heuristics-based, algorithms [41–43]. Another study used a deep learning approach to detect gait events from either three IMUs (worn on the low back, and both ankles) or a single IMU (worn on the low back), and showed that the time error was considerably smaller for the deep learning algorithm than for a commonly applied wavelet-based approach [44].

In this study, we further these works by validating this deep learning approach for detecting gait events in a heterogeneous cohort of healthy and neurologically diseased adults considering a single IMU setup, worn on the lower leg.

2. Materials and Methods

2.1. Data Collection

Gait analyses were performed in the Universitätsklinikum Schleswig-Holstein (UKSH) campus Kiel, Germany. The study [45] was approved by the ethical committee of the medical faculty at the UKSH (no: D438/18). In total, data from 160 participants were included for the current analysis, including data from young adults (YA; age: 18 - 60 years), older adults (OA; age: >60 years), people with Parkinson's Disease (PD; according to the UK Brain Bank criteria [46]), people with a recent (<4 weeks) symptomatic stroke (stroke), people with multiple sclerosis (MS; according to the McDonalds criteria [47]), people with chronic low back pain (cLBP), and people with other diagnoses that were assumed not to affect mobility (Table 1). Inclusion criteria were an age of 18 years or older, and the ability to walk independently without a walking aid. Participants were excluded from the study with a Montreal Cognitive Assessment [48] score <15 and other movement disorders that affected mobility, as noticed by the clinical assessor.

Table 1. Demographics data of the study participants. Age, height, and weight are presented as mean (standard deviation).

Group	Gender	Number of subjects	Age years	Height cm	Weight kg
YA	F	21	27 (7)	173 (5)	67 (8)
	M	22	30 (9)	185 (8)	81 (13)
OA	F	11	70 (6)	167 (6)	72 (17)
	M	11	73 (6)	180 (6)	83 (11)
PD	F	12	67 (6)	168 (6)	70 (14)
	M	20	62 (11)	178 (6)	87 (13)
MS	F	12	37 (9)	174 (9)	75 (9)
	M	9	42 (15)	189 (8)	96 (30)
stroke	F	4	66 (10)	160 (6)	65 (11)
	M	17	67 (17)	178 (7)	84 (14)
cLBP	F	3	64 (10)	166 (5)	65 (5)
	M	7	63 (16)	178 (7)	90 (16)
other	F	3	60 (13)	166 (3)	79 (15)
	M	8	68 (18)	182 (7)	85 (13)

Group: YA: younger adults, OA: older adults, PD: Parkinson's Disease, MS: multiple sclerosis, cLBP: chronic low back pain; **Gender:** F: female, M: male.

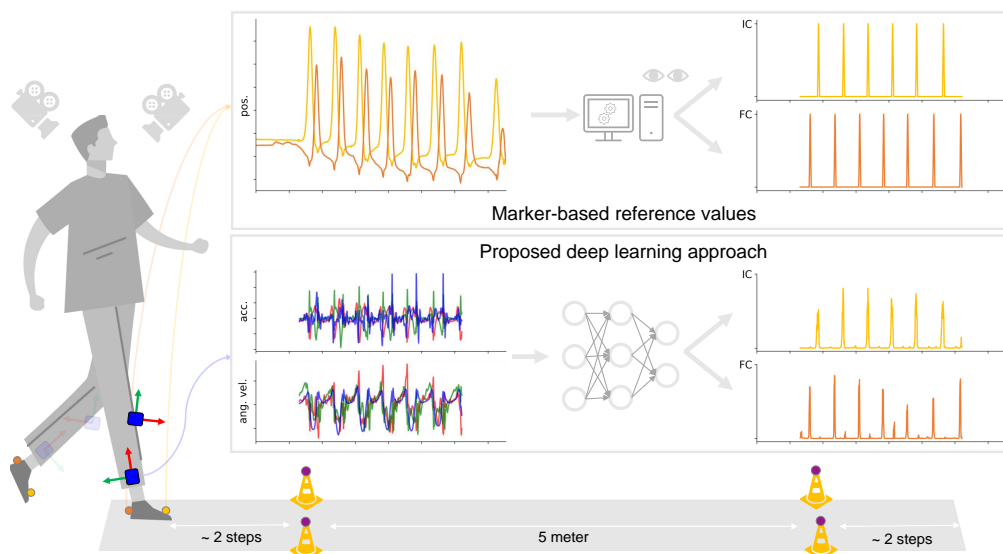


Figure 1. Schematic depiction¹ of the current study. Study participants wore IMUs on the ankle and shanks, and reflective markers were adhered on the heel and toe of usual footwear (shown on the left). Marker data were used to obtain reference values for the timings of initial and final contacts (top), where accelerometer and gyroscope data from each tracked point were inputted to a neural network that predicted timings of the same initial and final contacts (bottom).

¹ Picture from: <https://www.vecteezy.com/free-vector/man-walking>.

Participants performed three walking trials consisting of walking 5 meter at either (1) preferred speed ("Please walk at your normal walking speed."), (2) slow speed ("Please walk half of your normal walking speed."), or (3) fast speed ("Please walk as fast as possible, without running or falling."). The 5 meter distance was marked with two cones on both ends, and participants were asked to start walking approximately two steps before the cones on one end, and stop walking approximately two steps after passing the cones on the other end.

For the current analysis data from four IMUs (Noraxon USA Inc., myoMOTION, Scottsdale, AZ, USA) were considered, namely those that were attached laterally above the left and right ankle joint and those attached proximally at the left and right shank. IMUs were secured to participants using elastic bands with a special hold for the IMU. Furthermore, reflective markers were attached on top of the usual foot wear at the heel and toe of both feet (Figure 1). Marker data were recorded using a twelve-camera optoelectronic motion capture system (Qualisys AB, Göteborg, Sweden) at a sampling frequency of 200 Hz. IMU data were recorded at the same sampling frequency, and both systems were synchronized using a TTL signal [45]. For some recordings, the sampling frequency of the IMUs was mistakenly set at 100 Hz, and therefore data from these trials were upsampled before further analysis.

2.2. Data Pre-processing

2.2.1. Marker Data

The reflective marker data were first used to determine start and end of each trial. The start of the trial was defined as the time for which the first toe marker (left or right) crossed the *virtual* starting line. The end was defined as the time for which the last heel marker crossed the end line. For both marker and IMU systems, data were then cropped from start to the end of the trial.

Any remaining gaps in the marker data were filled by interpolation making use of inter-correlations between markers [49,50]. The data were then low-pass filtered using a double pass 6th order Butterworth filter with a cut-off frequency of 20 Hz [51,52]. The filtered data were differentiated to get velocity signals, and timings of ICs and FCs were determined from local maxima and minima in the heel and toe vertical velocity signals [53,54]. Like in [34,72], all identified ICs and FCs were manually checked, and corrected if necessary, using Qualisys Track Manager 2018.1 software (Qualisys AB, Göteborg, Sweden). The resulting annotated ICs and FCs were considered the *true* events (also *labels* or *targets*), and were used as reference timings to derive stride-specific gait parameters.

2.2.2. IMU Data

The idea behind the deep learning approach was that a *model* was trained to predict the likelihood of an IC and FC, given accelerometer and gyroscope data from a single IMU. The data from a single sensor channel, e.g., the acceleration in forward direction, or the angular velocity about the vertical direction, was denoted by $\mathbf{x}_d = [x_d[1] \ x_d[2] \ \cdots \ x_d[N]]^T$, with d referring to the d th sensor channel (i.e., $d = 1, \dots, D$) and n referring to the n th sample or time step (i.e., $n = 1, \dots, N$). Data from all D sensor channels together was then denoted by:

$$\mathbf{X} = \begin{bmatrix} | & | & \cdots & | \\ \mathbf{x}_1 & \mathbf{x}_2 & \cdots & \mathbf{x}_D \\ | & | & \cdots & | \end{bmatrix} = \begin{bmatrix} x_1[1] & x_2[1] & \cdots & x_D[1] \\ x_1[2] & x_2[2] & \cdots & x_D[2] \\ \vdots & \vdots & \cdots & \vdots \\ x_1[N] & x_2[N] & \cdots & x_D[N] \end{bmatrix}, \quad \mathbf{X} \in \mathbb{R}^{N \times D} \quad (1)$$

Likewise, the *labels* were denoted by:

$$\mathbf{y}_{\text{IC}} = \begin{bmatrix} y_{\text{IC}}[1] \\ y_{\text{IC}}[2] \\ \vdots \\ y_{\text{IC}}[N] \end{bmatrix}, \quad \mathbf{y}_{\text{FC}} = \begin{bmatrix} y_{\text{FC}}[1] \\ y_{\text{FC}}[2] \\ \vdots \\ y_{\text{FC}}[N] \end{bmatrix}, \quad y_{\text{IC/FC}}[n] \in [0, 1] \quad (2)$$

The model was then iteratively trained to learn a mapping $h_{\Theta}(\mathbf{X}) : \mathbf{X} \rightarrow \mathbf{y}$, where h_{Θ} was also referred to as the *hypothesis* which was parameterized by the weights and biases, collectively denoted by Θ .

All participant data were split in three independent data sets, namely a training set, a validation set, and a test set. Each set contained data from approximately one-third of the participants. Participants were randomly assigned to one of the sets, stratified by both group (i.e., diagnosis) and gender (Table 1).

Accelerometer and gyroscope data were normalized by subtracting the channel-wise mean, and dividing by the channel-wise standard deviation. Then, for the training and validation data set, the data were partitioned into equal length time windows [43] of 400 samples, with an overlap of 50% between successive windows (corresponding to 2 s windows, and 1 s overlap, respectively).

2.3. Model

2.3.1. Model Architecture

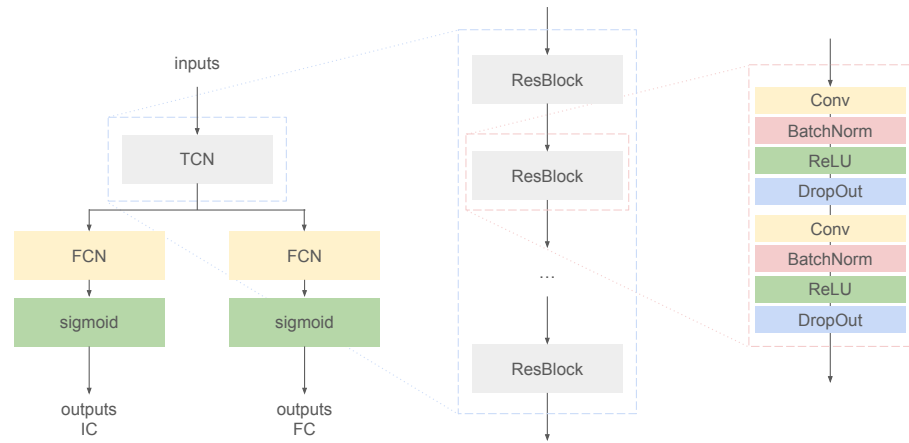


Figure 2. The generic model architecture of the deep learning model to predict initial contacts (ICs) and final contacts (FCs). The inputs are the accelerometer and gyroscope data from a single inertial measurement unit, and are fed to a temporal convolutional network (TCN) (left). The TCN consisted of repeating residual blocks (ResBlock) with exponentially increasing dilation factor (middle). Each ResBlock was built from two sequences of a convolutional layer (Conv), batch normalization layer (BatchNorm), a rectified linear unit activation layer (ReLU), and a dropout layer (DropOut) (right).

The generic architecture for the deep learning model was based on a temporal convolutional network (TCN) [43,57,58]. The TCN consisted of a sequence of residual blocks with exponentially increasing dilation factor [56,57]. Each residual block was built from two sequences of a dilated convolutional layer [56], a batch normalization layer [60], a rectified linear unit (ReLU) activation layer, and dropout layer [61] (Figure 2). The model was built in Python [62] using the high-level TensorFlow API Keras [58,63].

For the current analysis only convolutions of the "same" type were considered [58], i.e., the model was non-causal and zero-padded to account for edge effects, and the likelihood of an IC or FC was based on input data both before and after the current sample, n :

$$\hat{y}_i[n] = f(\cdots, \mathbf{X}[n-1], \mathbf{X}[n], \mathbf{X}[n+1], \cdots), \quad i \in \{\text{IC}, \text{FC}\} \quad (3)$$

The number of samples that the predictions at time n "sees", was referred to as the *receptive field* [64] and was a function of the kernel size and the dilation factors [58]. Dilation factors were always given as a sequence of increasing powers of 2 [56,57,65].

The outputs of the TCN block were fed to two separate fully-connected (FCN, or dense) layers, that both were followed by a sigmoid activation layer. Outputs were then

predicted separately for ICs and FCs. The mean squared error (MSE) was used as loss function, and a gradient descent-based optimization algorithm with adaptive moment (Adam) optimizer was used to iteratively learn the weights [66,67].

2.3.2. Hyperparameter Optimization

In order to find the best model architecture, hyperparameter tuning was performed using KerasTuner [68]. Here, the number of filters, the kernel size, and the maximum dilation factor (Table 2) were optimized using a random search strategy [69].

Table 2. Model hyperparameters that were optimized for, and the corresponding sets of possible values.

Description	Possible values
Number of filters	8, 16, 32, 64, 128
Kernel size	3, 5, 7
Dilations	[1, 2], [1, 2, 4], [1, 2, 4, 8]

The model architecture that resulted from the hyperparameter optimization was then trained on the combined set of training and validation data. The trained model was used to predict the likelihoods of ICs and FCs events on the hold-out test set.

2.4. Analysis

The predictions of the model on the test set data were compared with the labels from the test set. The model performance was evaluated for (1) overall detection performance, (2) time agreement between the predicted events and the (marker-based) annotated events, (3) agreement between subsequently derived gait parameters.

2.4.1. Overall detection performance

The overall detection performance quantified how many of the annotated events were detected by the model (true positives, TP), how many of the annotated events were not detected (false negatives, FN), and how many event that were detected, were actually not annotated (false positives, FP). An annotated event was considered to be detected if any detected event was within 0.075 s. A detected event were defined as peaks in the predicted likelihoods with a minimum height of 0.4, i.e. 40% likely that it corresponded to an event.

From these metrics, the recall (or sensitivity), precision and F_1 score were calculated as:

$$\text{recall} = \frac{TP}{TP + FN}$$

$$\text{precision} = \frac{TP}{TP + FP}$$

$$F_1 \text{ score} = 2 \cdot \frac{\text{recall} \cdot \text{precision}}{\text{recall} + \text{precision}}$$

2.4.2. Time Agreement

For all correctly detected gait events (TP, Section 2.4.1), the time agreement between the annotated and detect gait event was given by the time error:

$$\text{time error} = t_{\text{ref}} - t_{\text{pred}} \quad (4)$$

with t_{ref} the gait event time from the marker-based annotations, and t_{pred} the gait event time from the model predictions. As a robust measure for the average time error and its spread, the median time error and the inter-quartile range (IQR) were reported [70].

2.4.3. Stride Parameters

For those trials for which all gait events were detected, and no false positives were detected, the stride time, stance time and swing were calculated. Stride time was the time between two successive ICs of the same foot. Stance time was the time between a FC and the preceding IC of the same foot. Swing time was the time between the IC following the last FC of the same foot.

3. Results

3.1. Overall Detection Performance

The performance of detecting ICs and FCs was objectively quantified by the number of TPs, the number of FNs, and the number of FPs. From these, recall, precision and F_1 score were calculated (Table 3).

Table 3. Overall detection performance for initial contacts and final contacts as quantified by recall, precision and F_1 score.

Tracked point	Initial contacts						Final contacts					
	TP	FN	FP	recall	precision	F1	TP	FN	FP	recall	precision	F1
left ankle	624	19	5	97%	99%	98%	606	32	10	95%	98%	97%
right ankle	599	42	8	93%	99%	96%	614	17	12	97%	98%	98%
left shank	605	38	15	94%	98%	96%	585	53	18	92%	97%	94%
right shank	603	36	15	94%	98%	96%	595	30	9	95%	99%	97%

TP: true positives, FN: false negatives, FP: false positives, F1: F_1 score.

For both ICs and FCs, recall is high for each of the tracked points (i.e., $\geq 92\%$), and so is precision (i.e., $\geq 97\%$). Differences between the tracked points are small, i.e. the minimum recall is 92% and the maximum recall is 97%, and the minimum precision is 97% and the maximum precision is 99%.

3.2. Time Agreement

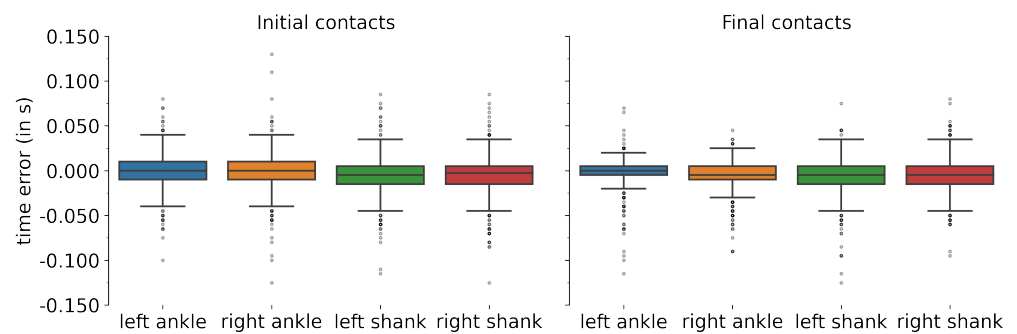


Figure 3. Time errors for initial (left) and final (right) contacts detection, for each of the different tracked points.

Time agreement between the annotated and detected events was quantified for the TPs for each of the tracked points (Figure 3). The median time error for each of the tracked points, and for both ICs and FCs was close to zero (Table 4), and the largest deviation was -0.005 s, corresponding to one sample period. The IQR was at most 0.020 s, corresponding to 4 sample periods.

Table 4. Time errors for the correctly detected gait events.

Tracked point	Initial contacts		Final contacts	
	median s	IQR s	median s	IQR s
left ankle	0.000	0.020	0.000	0.010
right ankle	0.000	0.020	-0.005	0.015
left shank	-0.005	0.020	-0.005	0.020
right shank	-0.003	0.020	-0.005	0.020

IQR: inter-quartile range.

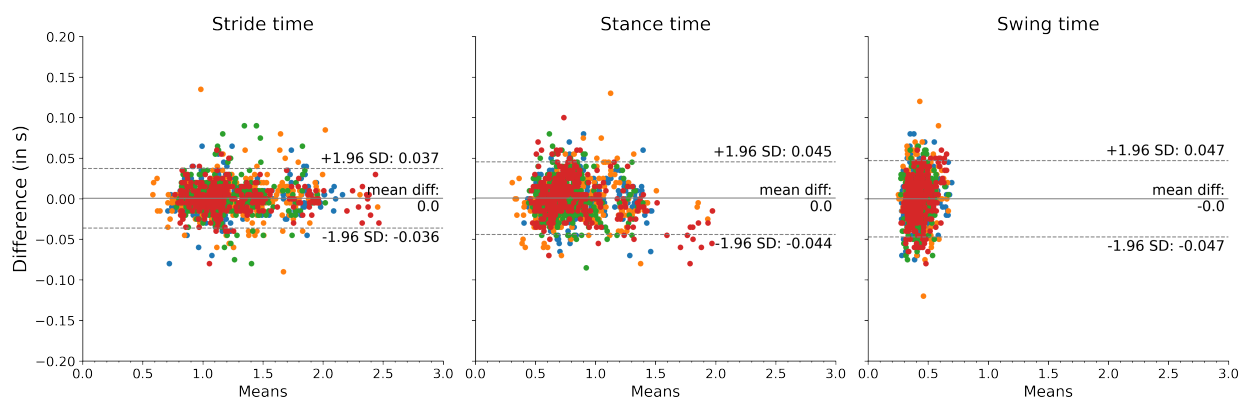
3.3. Stride Parameters

For those trials for which all gait events were correctly detected (and no false positives were detected), stride time, stance time and swing time were calculated. The mean difference and the limits of agreement between the marker-based annotations and the model-based detections were calculated.

For all stride-specific gait parameters, and for all tracked points, the mean difference was close to zero, i.e., the maximum mean difference was 0.003 s, namely for the calculated swing time of the right ankle (Table 5). Furthermore, for all gait parameters and for all tracked points the limits of agreement, based on a 95% confidence interval were distributed around a zero mean difference with the overall limits of agreement at -0.049 s and 0.051 s (Figure 4).

Table 5. Time agreement between the stride-specific parameters.

Tracked point	Parameters	Mean difference ms	Limits of Agreement (ms, ms)
left ankle	stride time	0.001	(-0.035, 0.036)
	stance time	0.002	(-0.039, 0.042)
	swing time	-0.001	(-0.045, 0.043)
right ankle	stride time	0.000	(-0.039, 0.040)
	stance time	-0.002	(-0.048, 0.044)
	swing time	0.003	(-0.046, 0.051)
left shank	stride time	0.001	(-0.039, 0.041)
	stance time	0.002	(-0.043, 0.046)
	swing time	-0.001	(-0.049, 0.047)
right shank	stride time	-0.000	(-0.031, 0.031)
	stance time	0.002	(-0.046, 0.049)
	swing time	-0.002	(-0.049, 0.046)

**Figure 4.** The agreement of extracted gait parameters between the sensor-based and marker-based methods.

4. Discussion

The current study aimed to validate a deep learning approach for detecting gait events from a single IMU worn on the lower leg. Data from a left and right ankle- and shank-worn IMU were used for training a deep learning model to detect gait events from walking trials performed by healthy younger and older adults, participants diagnosed with Parkinson's Disease, multiple sclerosis, or chronic low back pain, participants who had a recent symptomatic stroke, and participants diagnosed with other neurological diseases. Participants walked a 5 meter distance at three different self-selected walking speeds. The gait event timings that were predicted by the deep learning model were compared to a common reference method, i.e., optoelectronic motion capture, and clinically relevant stride-specific gait parameters were extracted.

A first measure for the model performance was given by the recall, precision and F_1 score. For both ICs and FCs, a high recall ($\geq 95\%$) and high precision ($\geq 98\%$) were observed, meaning that most events can be detected and most detected events were actually true events. There was little difference in recall and precision between tracked points (Table 3) confirming that the deep learning approach is relatively invariant to exact sensor location. These values for recall and precision were slightly lower than our previously reported values [34] that used a heuristics-based approach [30], but in the current study more trials were included and from a more heterogeneous study population. The F_1 score was comparable to the F_1 score obtained by [43] who compared marker-based gait event detection to manual annotations, and was also comparable to the F_1 score reported by [71] for gait event detection from a continuous wavelet-based approach.

Next, the time agreement, expressed by the time error the annotated event and the detected event, was of interest. For both ICs and FCs, and for all tracked points, the observed time error was small, and the middle 50% of the time errors were within a range of $[-0.015, 0.010]$ s (Table 4, Figure 3). These data showed that the deep learning-based approach is precise in detecting ICs and FCs. Time errors were slightly smaller than our previously reported results [34] that used a heuristics-based approach [30]. The heuristics-based approach determined ICs and FCs as local minima in the medio-lateral angular velocity [30] and it could be that these minima do not exactly coincide with the referent event timing as determined from the optoelectronic motion capture systems. The median and IQR of the time errors were in the same range as previously found by [44] that used a similar model architecture. The time errors were lower than previously reported time errors from a continuous wavelet-based approach [71].

From the correctly detected gait events, stride-specific gait parameters were derived. These are probably of greatest clinical relevance, as changes in stride-specific gait parameters have been linked directly with disease onset and progression [3–9]. Therefore, stride time, stance time, and swing time were calculated, and the differences between the deep learning-based approach and the marker-based reference method were quantified (Table 5, Figure 4). The limits of agreement for a 95% confidence interval were calculated, and for all metrics the zero mean difference was enclosed in the limit of agreement. These data provided evidence that the deep learning-based model was able to derive stride-specific gait parameters. The differences between the deep learning model-based stride parameters and the marker-based stride parameters were in a similar range as a recent study that compared IMU-derived stride parameters against stride parameters obtained with a pressure sensing walkway [29,44], and were also in the same range as a study that compared IMU-derived stride parameters with stride parameters obtained with an optoelectronic motion capture systems [72].

The main limitations of the current study were that only walking trials involving straight-line walking were considered, and the walking distance was relatively short. The observed gait patterns from these walking trials may therefore not fully representative of gait patterns observed in daily-life [19–21]. However, as the deep learning-based approach does not rely on fixed thresholds or assumptions regarding to which sensor axis is used, it is theoretically transferable and scalable to other conditions if input data and corresponding

labels can be provided. For example, in [44] a similar deep learning model was tested for a setup with three IMUs and for a setup with a single IMU, and observed differences in time errors compared to a pressure sensing walkway were negligible.

5. Conclusions

In this study we have validated a deep learning-based approach to detect gait events and subsequently extract clinically relevant stride parameters from a single inertial measurement unit worn either laterally above the ankle joint or proximally below the knee joint. Performance analysis showed an excellent detection rate, and low time errors in both event detection and stride parameter calculation for different walking speeds and across both healthy and neurological cohorts. Our next step is to validate these methods with real-life walking sequences.

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Institutional Review Board Statement: The study was conducted in accordance with the Declaration of Helsinki, and approved by the Ethics Committee of the medical faculty of the Christian-Albrechts-Universität zu Kiel (D438/18, approved on 8 May 2018).

Informed Consent Statement: Informed consent was obtained from all subjects involved in the study.

Data Availability Statement: Data from the first 10 participant are available online at <https://github.com/neurogeriatricskiel/Validation-dataset>. Additionally, we are preparing the open-source release of all data from the healthy younger and older adults. Data from patient groups can be shared upon reasonable request. The full algorithm pipeline is going to be made publicly available online at <https://github.com/rmndrs89/my-gait-events-tnn>.

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Conflicts of Interest: The authors declare no conflict of interest.

Abbreviations

The following abbreviations are used in this manuscript:

cLBP	chronic low back pain
CNN	convolutional neural network
IMU	inertial measurement unit
MS	multiple sclerosis
OA	older adults
PD	Parkinson's Disease
TCN	temporal convolutional network
YA	younger adults

References

1. Snijders, A.H.; van de Warrenburg, B.P.; Giladi, N.; Bloem, B.R. Neurological gait disorders in elderly people: clinical approach and classification. *Lancet Neurol.* **2007**, *6*(1), 63–74.
2. Hodgins, D. The importance of measuring human gait. *Med. Device Technol.* **2008**, *19*(5):42, 44–7.

3. Del Din, S.; Elshehabi, M.; Galna, B.; Hobert, M.A.; Warmerdam, E.; Suenkel, U.; Brockmann, K.; Metzger, F.; Hansen, C.; Berg, D.; Rochester, L.; Maetzler, W. Gait analysis with wearables predicts conversion to Parkinson disease. *Ann. Neurol.* **2019**, *86*(3), 357–367. 299
4. König, A.; Klaming, L.; Pijl, M.; Demeurraux, A.; Davis, R.; Robert, P. Objective measurement of gait parameters in healthy and cognitively impaired elderly using the dual-task paradigm. *Aging. Clin. Exp. Res.* **2017**, *29*(6), 1181–1189. 300
5. Bertoli, M.; Cereatti, A.; Trojaniello, D.; Avanzino, L.; Pelosin, E.; Del Din, S.; Rochester, L.; Ginis, P.; Bekkers, E.M.J.; Mirelman, A.; Hausdorff, J.M.; Della Croce, U. Estimation of spatio-temporal parameters of gait from magneto-inertial measurement units: multicenter validation among Parkinson, mildly cognitively impaired and healthy older adults. *Biomed. Eng. Online* **2018**, *17*(58). 301
6. von Schroeder, H.P.; Coutts, R.D.; Lyden, P.D.; Billings, E., Jr; Nickel, V. L. Gait parameters following stroke: a practical assessment. *J. Rehabil. Res. Dev.* **1995**, *32*(1), 25–31. 302
7. Mohan, D.M.; Khandoker, A.H.; Wasti, S.A.; Ismail Ibrahim Ismail Alali, S.; Jelinek, H.F.; Khalaf, K. Assessment methods of post-stroke gait: A scoping review of technology-driven approaches to gait characterization and analysis. *Front. Neurol.* **2021**, *12*. 303
8. Griškevičius, J.; Apanskienė, V.; Žižienė, J.; Daunoravičienė, K.; Ovčnikova, A.; Kizlaitienė, R.; Sereikė, I.; Kaubrys, G.; Pauk, J.; Idźkowski, A. Estimation of temporal gait parameters of multiple sclerosis patients in clinical setting using inertial sensors. In Proceedings of the 11th Int Conf BIOMDLORE 2016, Druskininkai, Lithuania, 20–22 Oct 2016; 80–82. 304
9. Flachenecker, F.; Gaßner, H.; Hannik, J.; Lee, D.H.; Flachenecker, P.; Winkler, J.; Eskofier, B.; Linker, R.A.; Klucken, J. Objective sensor-based gait measures reflect motor impairment in multiple sclerosis patients: Reliability and clinical validation of a wearable sensor device. *Mult. Scler. Relat. Dis.* **2019**, *39*, 101903. 305
10. Hannink, J.; Kautz, T.; Pasluosta, C.F.; Gaßmann, K.-G.; Klucken, J.; Eskofier, B.M. Sensor-Based Gait Parameter Extraction With Deep Convolutional Neural Networks. *IEEE J. Biomed. Health* **2017**, *21*(1), 85–93. 306
11. Perry, J.; Burnfield, J.M. *Gait analysis: normal and pathological gait*, 2nd ed.; SLACK Inc.: Thorofare, NJ, USA, 2010. 307
12. Richards, J.; Levine, D.; Whittle, M. *Whittle's Gait Analysis*, 5th ed.; Churchill Livingstone: London, UK, 2012. 308
13. Rueterbories, J.; Spaich, E.G.; Larsen, B.; Andersen, O.K. Methods for gait event detection and analysis in ambulatory systems. *Med. Eng. Phys.* **2010**, *32*(6), 545–552. 309
14. Bruening, D.A.; Ridge, S.T. Automated event detection algorithms in pathological gait. *Gait Posture* **2014**, *39*(1), 472–477. 310
15. Chiari, L.; Della Croce, U.; Leardini, A.; Cappozzo, A. Human movement analysis using stereophotogrammetry: Part 2: Instrumental errors *Gait Posture* **2005**, *21*(2), 197–211. 311
16. Topley, M.; Richards, J.G. A comparison of currently available optoelectronic motion capture systems. *J. Biomech.* **2020**, *106*:109820. 312
17. Iosa, M.; Picerno, P.; Paolucci, S.; Morone, G. Wearable inertial sensors for human movement analysis. *Expert Rev. Med. Devic.* **2016**, *13*(7), 641–659. 313
18. Jarchi, D.; Pope, J.; Lee, T.K.M.; Tamjidi, L.; Mirzaei, A.; Sanei, S. A Review on Accelerometry-Based Gait Analysis and Emerging Clinical Applications. *IEEE Rev. Biomed. Eng.* **2018**, *11*, 177–194. 314
19. Hillel, I.; Gazit, E.; Nieuwboer, A.; Avanzino, L.; Rochester, L.; Cereatti, A.; Della Croce, U.; Rikkert, M.O.; Bloem, B.R.; Pelosin, E.; Del Din, S.; Ginis, P.; Giladi, N.; Mirelman, A.; Hausdorff, J.M. Is every-day walking in older adults more analogous to dual-task walking or to usual walking? Elucidating the gaps between gait performance in the lab and during 24/7 monitoring. *Eur. Rev. Aging Phys. A.* **2019**, *16*(6). 315
20. Warmerdam, E.; Hausdorff, J.M.; Atrsaiei, A.; Zhou, Y.; Mirelman, A.; Aminian, K.; Espay, A.J.; Hansen, C.; Evers, L.J.W.; Keller, A.; Lamothe, C.; Pilotto, A.; Rochester, L.; Schmidt, G.; Bloem, B.R.; Maetzler, W. Long-term unsupervised mobility assessment in movement disorders. *Lancet Neurol.* **2020**, *19*(5), 462–470. 316
21. Atrsaiei, A.; Corrá, M.F.; Dadashi, F.; Vila-Chã, N.; Maia, L.; Mariani, B.; Maetzler, W.; Aminian, K. Gait speed in clinical and daily living assessments in Parkinson's disease patients: performance versus capacity. *npj Parkinsons Dis.* **2021**, *7*(1):24. 317
22. Del Din, S.; Godfrey, A.; Mazzà, C.; Lord, S.; Rochester, L. Free-living monitoring of Parkinson's disease: Lessons from the field. *Movement Disord.* **2016**, *31*(9), 1293–1313. 318
23. Shah, V.V.; McNames, J.; Mancini, M.; Carlson-Kuhta, P.; Nutt, J.G.; El-Gohary, M.; Lapidus, J.A.; Horak, F.B.; Curtze, C. Digital Biomarkers of Mobility in Parkinson's Disease During Daily Living. *J. Parkinson Dis.* **2020**, *10*(3), 1099–1111. 319
24. Fasano, A.; Mancini, M. Wearable-based mobility monitoring: the long road ahead. *Lancet Neurol.* **2020**, *19*(5), 378–379. 320
25. Corrá, M.F.; Atrsaiei, A.; Sardoreira, A.; Hansen, C.; Aminian, K.; Correia, M.; Vila-Chã, N.; Maetzler, W.; Maia, L. Comparison of Laboratory and Daily-Life Gait Speed Assessment during ON and OFF States in Parkinson's Disease. *Sensors* **2021**, *21*(12):3974. 321
26. Ben Mansour, K.; Rezzoug, N.; Gorce, P. Analysis of several methods and inertial sensors locations to assess gait parameters in able-bodied subjects. *Gait Posture* **2015**, *42*, 409–414. 322
27. Storm, F.A.; Buckley, C.J.; Mazzà, C. Gait event detection in laboratory and real life settings: Accuracy of ankle and waist sensor based methods. *Gait Posture* **2016**, *50*, 42–46. 323
28. Panebianco, G.P.; Bisi, M.C.; Stagni, R.; Fantozzi, S. Analysis of the performance of 17 algorithms from a systematic review: Influence of sensor position, analysed variable and computational approach in gait timing estimation from IMU measurements. *Gait Posture* **2018**, *66*, 76–82. 324
29. Niswander, W.; Kontson, K. Evaluating the Impact of IMU Sensor Location and Walking Task on Accuracy of Gait Event Detection Algorithms. *Sensors* **2021**, *21*(12):3989. 325

30. Salarian, A.; Russmann, H.; Vingerhoets, F.J.; Dehollain, C.; Blanc, Y.; Burkhard, P.R.; Aminian, K. Gait assessment in Parkinson's disease: Toward an ambulatory system for long-term monitoring. *IEEE Trans. Biomed. Eng.* **2004**, *51*(8), 1434–1443. 357
31. Catalfamo, P.; Ghoussayni, S.; Ewins, D. Gait Event Detection on Level Ground and Incline Walking Using a Rate Gyroscope. *Sensors* **2010**, *10*(6), 5683–5702. 358
32. Sabatini, A.; Martelloni, C.; Scapellato, S.; Cavallo, F. Assessment of Walking Features from Foot Inertial Sensing. *IEEE Trans. Biomed. Eng.* **2005**, *52*(3), 486–494. 359
33. Maqbool, H.F.; Husman, M.A.B.; Awad, M.; Abouhossein, A.; Mehryar, P.; Iqbal, N.; Dehghani-Sanij, A.A. Real-time gait event detection for lower limb amputees using a single wearable sensor. In Proceedings of the 2016 38th Ann. Int. Conf. of the IEEE Eng. in Med. Biol. Soc. (EMBC), Orlando, FL, USA, 16–20 Aug. 2016; IEEE: Piscataway, NJ, USA, 2016. 360
34. Romijnders, R.; Warmerdam, E.; Hansen, C.; Welzel, J.; Schmidt, G.; Maetzler, W. Validation of IMU-based gait event detection during curved walking and turning in older adults and Parkinson's Disease patients. *J. Neuroeng. Rehabil.* **2021**, *18*:28. 361
35. Jasiewicz, J.M.; Allum, J.H.; Middleton, J.W.; Barriskill, A.; Condie, P.; Purcell, B.; Li, R.C.T. Gait event detection using linear accelerometers or angular velocity transducers in able-bodied and spinal-cord injured individuals. *Gait Posture* **2006**, *24*(4), 502–509. 362
36. Trojaniello, D.; Cereatti, A.; Pelosin, E.; Avanzino, L.; Mirelman, A.; Hausdorff, J.M.; Della Croce, U. Estimation of step-by-step spatio-temporal parameters of normal and impaired gait using shank-mounted magneto-inertial sensors: Application to elderly, hemiparetic, parkinsonian and choreic gait. *J. Neuroeng. Rehabil.* **2014**, *11*:152. 363
37. Ferraris, F.; Grimaldi, U.; Parvis, M. Procedure for effortless in-field calibration of three-axis rate gyros and accelerometers. *Sensor. Mater.* **1995**, *7*(5), 311–330. 364
38. Greene, B.R.; McGrath, D.; O'Neill, R.; O'Donovan, K.J.; Burns, A.; Caulfield, B. An adaptive gyroscope-based algorithm for temporal gait analysis. *Med. Biol. Eng. Comput.* **2010**, *48*(12), 1251–1260. 365
39. LeCun, Y.; Bengio, Y.; Hinton, G. Deep learning. *Nature* **2015**, *521*, 436–444. 366
40. Ter Haar Romeny, B.M. A Deeper Understanding of Deep Learning. In *Artificial Intelligence in Medical Imaging: Opportunities, Applications and Risks*; Ranschaert, E.R., Morozov, S., Algra, P.R., Eds.; Springer Int. Publish.: Cham, Switzerland, 2019; pp. 25–38. 367
41. Kidziński, L.; Delp, S.; Schwartz, M. Automatic real-time gait event detection in children using deep neural networks. *PLoS ONE* **2019**, *14*(1):e0211466. 368
42. Lempereur, M.; Rousseau, F.; Rémy-Néris, O.; Pons, C.; Houx, L.; Quéllec, G.; Brochard, S. A new deep learning-based method for the detection of gait events in children with gait disorders: Proof-of-concept and concurrent validity. *J. Biomech.* **2020**, *98*, 109490. 369
43. Filtjens, B.; Nieuwboer, A.; D'cruz, N.; Spildooren, J.; Slaets, P.; Vanrumste, B. A data-driven approach for detecting gait events during turning in people with Parkinson's disease and freezing of gait. *Gait Posture* **2020**, *80*, 130–136. 370
44. Gadaleta, M.; Cisotto, G.; Rossi, M.; Ur Rehman, R.Z.; Rochester, L.; Del Din, S. Deep Learning Techniques for Improving Digital Gait Segmentation. In Proceedings of the 2019 41st Ann. Int. Conf. of the IEEE Eng. in Med. Biol. Soc. (EMBC), Berlin, Germany, 23–27 Jul. 2019; IEEE: Piscataway, NJ, USA, 2019. 371
45. Warmerdam, E.; Romijnders, R.; Geritz, J.; Elshehabi, M.; Maetzler, C.; Otto, J.C.; Reimer, M.; Stuermer, K.; Baron, R.; Paschen, S.; Beyer, T.; Dopcke, D.; Eiken, T.; Ortmann, H.; Peters, F.; von der Recke, F.; Riesen, M.; Rohwedder, G.; Schaade, A.; Schumacher, M.; Sondermann, A.; Maetzler, W.; Hansen, C. Proposed Mobility Assessments with Simultaneous Full-Body Inertial Measurement Units and Optical Motion Capture in Healthy Adults and Neurological Patients for Future Validation Studies: Study Protocol. *Sensors* **2021**, *21*(17):5833. 372
46. Gibb, W.R.; Lees, A.J. The relevance of the Lewy body to the pathogenesis of idiopathic Parkinson's disease. *J. Neurol. Neurosurg. Ps.* **1988**, *51*(6), 745–752. 373
47. Thompson, A.J.; Banwell, B.L.; Barkhof, F.; Carroll, W.M.; Coetsee, T.; Comi, G.; Correale, J.; Fazekas, F.; Filippi, M.; Freedman, M.S.; Fujihara, K.; Galetta, S.L.; Hartung, H.P.; Kappos, L.; Lublin, F.D.; Marrie, R.A.; Miller, A.E.; Miller, D.H.; Montalban, X.; Mowry, E.M.; Soelberg Sorensen, P.; Tintoré, M.; Traboulsee, A.L.; Trojano, M.; Uitdehaag, B.M.J.; Vukusic, S.; Waubant, E.; Weinshenker, B.G.; Reingold, S.C.; Cohen, J.A. Diagnosis of multiple sclerosis: 2017 revisions of the McDonald criteria. *Lancet Neurol.* **2018**, *17*(2), 162–173. 374
48. Nasreddine, Z.S.; Phillips, N.A.; Bédirian, V.; Charbonneau, S.; Whitehead, V.; Collin, I.; Cummings, J.L.; Chertkow, H. The Montreal Cognitive Assessment, MoCA: A Brief Screening Tool For Mild Cognitive Impairment. *J. Am. Geriatr. Soc.* **2005**, *53*(4), 695–699. 375
49. Federolf, P.A. A Novel Approach to Solve the "Missing Marker Problem" in Marker-Based Motion Analysis That Exploits the Segment Coordination Patterns in Multi-Limb Motion Data. *PLoS ONE* **2013**, *8*(10), 1–13. 376
50. Gløersen, Ø.; Federolf, P. Predicting Missing Marker Trajectories in Human Motion Data Using Marker Intercorrelations. *PLoS ONE* **2016**, *11*(3), 1–14. 377
51. Kormylo, J.; Jain, V. Two-pass recursive digital filter with zero phase shift. *IEEE T. Acoust. Speech* **1974**, *22*(5), 384–387. 378
52. Rácz, K.; Rita, M.K. Marker displacement data filtering in gait analysis: A technical note. *Biomed. Signal Proces.* **2021**, *70*:102974. 379
53. Pijnappels, M.; Bobbert, M.F.; Van Dieën, J.H. Changes in walking pattern caused by the possibility of a tripping reaction. *Gait Posture* **2001**, *14*(1), 11–18. 380
54. O'Connor, C.M.; Thorpe, S.K.; O'Malley, M.J.; Vaughan, C.L. Automatic detection of gait events using kinematic data. *Gait Posture* **2007**, *25*(3), 469–474. 381

72. Carcreff, L.; Gerber, C.; Paraschiv-Ionescu, A.; De Coulon, G.; Newman, C.; Armand, S.; Aminian, K. What is the best configuration of wearable sensors to measure spatiotemporal gait parameters in children with cerebral palsy? *Sensors* **2018**, *18*(2):394. 415
56. Yu, F.; Koltun, V. Multi-Scale Context Aggregation by Dilated Convolutions. In Proceedings of the 4th Int. Conf. on Learning Representations (ICLR), San Juan, Puerto Rico, 2–4 May 2016, Conference Track Proceedings, 2016. 416
57. Bai, S.; Kolter, J.Z.; Koltun, V. An Empirical Evaluation of Generic Convolutional and Recurrent Networks for Sequence Modeling. *arXiv:1803.01271* **2018**. 417
58. Rémy, P. Temporal Convolutional Networks for Keras. *GitHub repository* **2020**, *GitHub*, <https://github.com/philipperemy/keras-tcn>. 418
59. LeCun, Y.; Boser, B.; Denker, J.S.; Henderson, D.; Howard, R.E.; Hubbard, W.; Jackel, L.D. Backpropagation Applied to Handwritten Zip Code Recognition. *Neural Comput.* **1989**, *1*(4), 541–551. 419
60. Ioffe, S.; Szegedy, C. Batch Normalization: Accelerating Deep Network Training by Reducing Internal Covariate Shift. *arXiv:1502.03167* **2015**. 420
61. Srivastava, N.; Hinton, G.; Krizhevsky, A.; Sutskever, I.; Salakhutdinov, R. Dropout: a simple way to prevent neural networks from overfitting. *J. Mach. Learn. Res.* **2014**, *15*, 1929–1958. 421
62. Van Rossum, G.; Drake, F.L. *Python 3 Reference Manual*; CreateSpace, Scotts Valley, CA, USA, 2009. 422
63. Chollet, F. and others. Keras. **2015**. <https://keras.io>. 423
64. Lea, C.; Flynn, M.D.; Vidal, R.; Reiter, A.; Hager, G.D. Temporal convolutional networks for action segmentation and detection. In Proceedings of the IEEE Conference on Computer Vision and Pattern Recognition, Honolulu, HI, USA, 22–25 Jul. 2017; pp. 156–165. 424
65. van den Oord, A.; Dieleman, S.; Zen, H.; Simonyan, K.; Vinyals, O.; Graves, A.; Kalchbrenner, N.; Senior, A.W.; Kavukcuoglu, K. WaveNet: A generative model for raw audio. *arXiv:1609.03499* **2016**. 425
66. Kingma, D.P.; Ba, J. Adam: A Method for Stochastic Optimization *arXiv:1412.6980v9* **2014**. 426
67. Schmidt, R.M.; Schneider, F.; Henning, P. Descending through a crowded valley-benchmarking deep learning optimizers. In Proceedings of the 38th Int. Conf. on Machine Learning (ICML), online, 18–24 Jul. 2021, pp. 9367–9376. 427
68. O'Malley, T.; Bursztein, E.; Long, J.; Chollet, F.; Jin, H.; Invernizzi, L.; and others; KerasTuner. *GitHub repository* **2019**, *GitHub*, <https://github.com/keras-team/keras-tuner>. 428
69. Bergstra, J.; Bengio, Y. Random Search for Hyper-Parameter Optimization. *J. Mach. Learn. Res.* **2012**, *13*(10), 281–305. 429
70. Diez, D.; Çetinkaya-Rundel, M.; Barr, C.D. *OpenIntro Statistics*, 4th ed.; 2019. <https://www.openintro.org/book/os/> 430
71. Ji, N.; Zhou, H.; Guo, K.; Samuel, O.W.; Huang, Z.; Xu, L.; Li, G. Appropriate Mother Wavelets for Continuous Gait Event Detection Based on Time-Frequency Analysis for Hemiplegic and Healthy Individuals. *Sensors* **2019**, *19*(16):3462. 431
72. Carcreff, L.; Gerber, C.N.; Paraschiv-Ionescu, A.; De Coulon, G.; Newman, C.J.; Armand, S.; Aminian, K. What is the Best Configuration of Wearable Sensors to Measure Spatiotemporal Gait Parameters in Children with Cerebral Palsy? *Sensors* **2018**, *18*(2):394. 432