

Research on physical activity variability and changes of metabolic profile in patients with prediabetes using Fitbit activity trackers data

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Abstract.

BACKGROUND: Monitoring physical activity with consumers wearables is one of the possibilities to control a patient's self-care and adherence to recommendations. However, clinically approved methods, software, and data analysis technologies to collect data and make it suitable for practical use for patient care are still lacking.

OBJECTIVE: This study aimed to analyze the potential of patient physical activity monitoring using Fitbit physical activity trackers and find solutions for possible implementation in the health care routine.

METHODS: Thirty patients with impaired fasting glycemia were randomly selected and participated for 6 months. Physical activity variability was evaluated and parameters were calculated using data from Fitbit Inspire devices.

RESULTS: Changes in parameters were found and correlation between clinical data (HbA1c, lipids) and physical activity variability were assessed. Better correlation with variability than with body composition changes shows the potential to include nonlinear variability parameters analysing physical activity using mobile devices. Less expressed variability shows better relationship with control of prediabetic and lipid parameters.

CONCLUSIONS: Evaluation of physical activity variability is essential for patient health, and these methods used to calculate it is an effective way to analyze big data from wearable devices in future trials.

Keywords: Fitbit, Poincaré plot, variability, physical activity monitoring, pre-diabetes

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1. Introduction

Lack of physical activity and obesity belong to the biggest health problems we face. This leads to increased mortality related to cardiovascular diseases and diabetes. One of the conditions caused by lack of physical activity and malnutrition is prediabetes. Prediabetes is a transitional hyperglycaemic condition when glucose levels are elevated but not high enough to meet the diagnostic criteria for diabetes [1,2].

However, it is a reversible condition, and prompt diagnosis and preventive action can stop or delay development or progression of its complications [3]. One of the actions to prevent these chronic diseases is increasing and promoting of physical activity.

The World Health Organization (WHO) 2010 guidelines state that an adult without significant mobility impairments should assign at least 150 minutes per week to moderate-intensity aerobic exercise or at least 75 minutes to high-intensity physical exercise or an equivalent combination of physical activity. To improve health and prevent chronic diseases, it is recommended at least to double time indicated in the basic guidelines for physical activity [4–6].

Healthy lifestyle professionals and disease prevention program developers are actively looking for ways to adapt and introduce more active lifestyle habits to the general public. Walking is considered a form of moderate physical activity and is a great way to maintain the required amount of physical activity [7–10]. It not only facilitates easy access from point A to point B but also contributes to better health. The American College of Sports Medicine has systematized WHO recommendations and compared the usual recommended physical activity with 7,000–9,000 steps per day [11–13]. Recent years went through enormous mobile devices breakthroughs, and now we can count our steps using smartphones, smartwatches, and activity trackers [14]. Smart bracelets measure the daily number of steps with sufficient accuracy and are suitable for long-term physical activity assessment [15]. Analysis of the data collected by these smart devices has shown that, on average, people walk 4,000 to 6,000 steps per day, which is only two-thirds of the recommended physical activity [16].

Now we have a simple way to test the physical activity change in free-living conditions using consumer wearables, such as smartwatches and physical activity trackers. Patients have the opportunity to perform self-monitoring of their physical activity. Some studies show us how much time people spend sedentary or physically active. Studies also show that physical activity during the week varies and can differ significantly [17–20].

The aim of this study was to evaluate how physical activity variability, in the long run, impacts people's health in free-living conditions and to analyse what impact physical activity variability in 6 months has on their glycated hemoglobin, cholesterol, and body composition. Our study included people with prediabetes who have a higher risk of developing diabetes.

During the literature review, we haven't found the same or similar research looking for physical activity variability impact to people's health in free-living conditions, so we have got some intriguing results for future research.

The aims of this study are: 1) to analyze the potential of patient physical activity monitoring using Fitbit devices; 2) to find variables associated with lipid profile and HbA1c changes in 6 months; and 3) to find solutions for possible implementation in the health care routine and health information system.

2. Methods

2.1. Subjects

Vilnius University Hospital Santaros Klinikos (VUHKS) Family Medicine Centre patients visiting their family doctor were included in our study. The selected study population had increased fasting glucose

levels (5.6 to 6.9 mmol/l), did not have skeletal or muscular diseases, and conditions that compromise patients' ability to move. The participants had to have a smartphone with the latest operating system, such as Apple iOS 13 or later and Android OS 8.0 or later, an email address and be able to use it.

30 subjects were selected. Subjects were informed about the study, its possibilities and benefits for their health. Informed consent form has been signed.

Blood tests were performed during the first consultation: total cholesterol, triglycerides, low-density lipoproteins, high-density lipoproteins, glycated hemoglobin. We also performed body mass composition analysis with the medical bioimpedance device X-Contact 356. Also, we provided physical activity recommendations and consulted on a healthy diet.

Subjects were given Fitbit Inspire physical activity trackers (PAT) and they were asked to connect the bracelet to their smartphones through the Fitbit application. If the patient failed to perform these actions, the PAT was connected by a family doctor.

After 6 months, during a consultation at the family medicine center, the subject was asked to transfer data from his online Fitbit account about his physical activity during the study period to the principal researcher's main computer.

During this visit, we repeated blood tests: total cholesterol, triglycerides, low-density lipoproteins, high-density lipoproteins, glycated hemoglobin, and bodyweight analysis. The collected information was analysed and discussed with patients in order to suggest lifestyle changes.

The study was approved by the Vilnius Regional Biomedical Research Ethics Committee (approval no. 2019/6-1143-634).

2.2. Physical activity monitoring using Fitbit

Participants received Fitbit Inspire PAT (free of charge) during the consultation with the family doctor and agreed to allow researchers to collect data from their accounts. Subjects installed the Fitbit app on their smartphones and created their own user accounts with a personal email address. Then using a "Bluetooth" connection on the smartphones they connected their smart bracelets and synchronized with their personal Fitbit account. If the participant had difficulty with creating a personal account or connecting a bracelet, we helped him to do this and we taught him how to use the device [21]. Subjects had to periodically synchronise data with their phone throughout the study period by activating the Fitbit application on their smartphone and regularly charge their bracelet every 6–7 days.

After 6 months, during the routine visit at the family doctor's office, we asked patients to log in to their accounts at <https://accounts.fitbit.com/login>. Then we used the Fitbit data export platform to download the data covering 30 days interval from start to finish of the study (Figs 1 and 2).

Data obtained from the smart bracelets were: physical activity recording data, steps per day, distance, minutes sedentary, minutes weakly active, active, minutes fairly active, minutes very active.

2.3. Poincaré plot and physical activity variability parameters

In this study, the Poincaré plot was used to analyze physical activity variability. The Poincaré plot is a two-dimensional scatterplot of paired values of time series data points, where the x-axis represents the current value at data point $n(x_n)$ and the y-axis represent value at data point with a time delay of $i(x_{n+i})$ (Fig. 3).

This method is widely used to study physiological signals [22], especially in the analysis of heart rate variability [23–25]. but there are also promising applications in other areas such as measuring

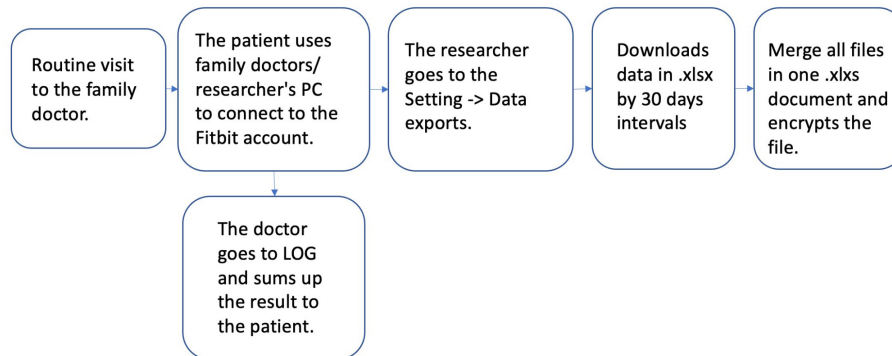


Fig. 1. Fitbit data extraction process.

The screenshot shows the 'Your Fitbit Data' export interface. On the left is a navigation menu with sections for 'DEVICES' (Inspire), 'SETTINGS' (Personal Info, Notifications, Privacy, Data Export, Manage Data, Manage Account Access, Applications, Help), and a 'Logout' button. The main content area is titled 'Your Fitbit Data' and includes the instruction: 'Export your Fitbit data in Microsoft Excel or CSV format, which can be opened in any spreadsheet application.' Below this, the 'Time Period' section allows selection of 'This Week', 'This Month', 'Last Week', 'Last Month', or 'Custom' (selected). The 'Start Date' is set to 2021-01-01 and the 'End Date' is 2021-01-31. The 'Include Data' section has checkboxes for 'Body', 'Foods', 'Activities' (checked), and 'Sleep'. The 'File Format' is set to 'Microsoft Excel' and a 'Download' button is visible at the bottom.

Fig. 2. Fitbit data export platform.

hand grip strength [26], assessing blood pressure variability [27], monitoring temporal blood glucose fluctuations [28–30], studying reaction time during repeated testing [31], etc.

To analyze physical activity variability, the Poincaré plot method with a time delay of one day ($i = 1$) was applied and additional measures for selected physical activity time series accessed from patients' Fitbit data were calculated. An average value (AVG), standard deviation (SD) and some standard Poincaré plot parameters representing short-term variability (SD1), long-term variability (SD2), the ratio of SD1 and SD2 (SD12) and area of fitting ellipse (AFE) were selected to analyze variability, as defined in other studies [22–31]:

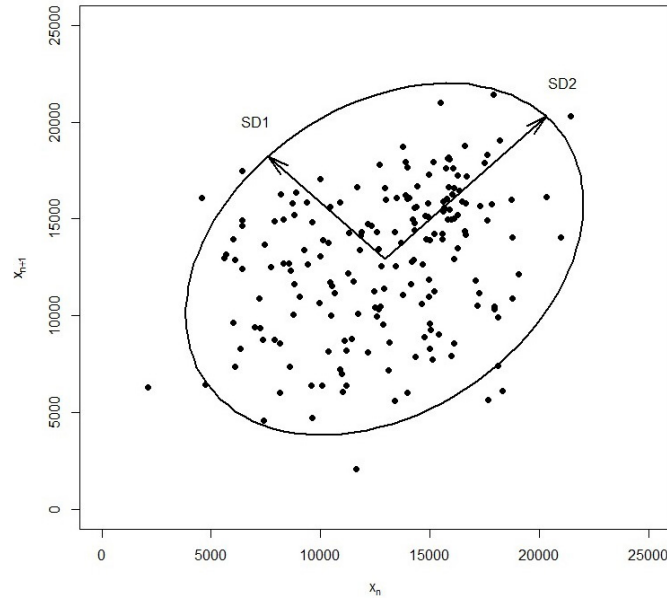


Fig. 3. Example of Poincaré plot for Step counts per day.

$$SD1 = \frac{\sqrt{2}}{2} * SD(x_n - x_{n+1})$$

$$SD2 = \sqrt{2SD(x_n)^2 - \frac{1}{2}SD(x_n - x_{n+1})^2}$$

$$SD12 = \frac{SD1}{SD2}$$

$$AFE = \pi * SD1 * SD2$$

2.4. Statistical analysis

The Shapiro-Wilk test was used for normality testing. Data was presented as mean and standard deviation (mean \pm SD) or median and quartiles (median [Q1–Q3]) when appropriate. Paired *t* test or Wilcoxon's matched pairs signed rank test was used for baseline and follow-up data comparison. Pearson's or Spearman's correlation was performed to assess association between changes of selected variables and physical activity variability parameters. Analyses were conducted using jamovi 1.6 [32] and R 4.1.0 [33], and R packages RHRV [34], corrplot [35]. The list of abbreviations used in this paper is presented in Table 1.

3. Results

3.1. Subjects

Mean age of study subjects was 53.8 ± 9.1 (median [Q1–Q3]: 54 [49.5–61]), there were 9 males and 21 females. After intervention, significant differences in patient weight, body composition and self-reported physical activity were observed (Table 2).

Table 1
The list of abbreviations

Shortening	Full text	Units
HgbA1c %	Glycated hemoglobin, percentage	%
HgbA1c mmol/l	Glycated hemoglobin, millimoles per liter	mmol/l
TC	Total cholesterol	mmol/l
TG	Triglycerides	mmol/l
HDL	High-density lipoprotein	mmol/l
LDL	Low-density lipoprotein	mmol/l
<i>Bio-electrical impedance analysis</i>		
	Height	cm
	Weight	kg
BMI	Body Mass Index	
LBM	Lean body mass	kg
MBF	Mass of body fat	kg
SLM	Soft lean mass	kg
SMM	Skeletal muscle mass	kg
Mineral	Body minerals	kg
Protein	Body proteins	kg
TBW	Total body water	kg
PBF	Percent body fat	%
VFL	Visceral fat level	units
VFA	Visceral fat area	cm ²
WHR	Waist to hip ratio	Ratio
AC	Abdominal circumference	cm
SLM Lt. ARM	Left arm soft lean mass	kg
SLM Rt.ARM	Right arm soft lean mass	kg
SLM Lt.LEG	Left leg soft lean mass	kg
SLM Rt.LEG	Right leg soft lean mass	kg
SLM Trunk	Trunk soft lean mass	kg
MBF Lt. ARM	Left arm mass of body fat	kg
MBF Rt.ARM	Right arm mass of body fat	kg
MBF Lt.LEG	Left leg mass of body fat	kg
MBF Rt.LEG	Right leg mass of body fat	kg
MBF Trunk	Trunk mass of body fat	kg
E.C.W/T.B.W	Extra-cellular water to total body water ratio	Ratio
<i>Fitbit activity tracker data</i>		
Data	Day of physical activity registration	DD-MM-YYYY
S	Steps per day	
D	Distance per day	km
MS	Minutes sedentary	minutes
MLA	Minutes lightly active	minutes
MFA	Minutes fairly active	minutes
MVA	Minutes very active	minutes

3.2. Physical activity monitoring using Fitbit

In order to analyze physical activity variability, the Poincaré plots (Fig. 4) were represented and corresponding parameters were calculated for the exported Fitbit data (Table 3). Different patterns of physical activity variability were observed in different patients and different Fitbit data variables. SD1/SD2 ratio varied between 0.697 and 0.870 indicating higher long-term than short-term variability.

3.3. Poincaré plot and physical activity variability (PHAV) parameters

To measure association between physical activity variability (PHAV) parameters accessed from the Poincaré plot, Pearson's and Spearman's correlation coefficients were calculated (Fig. 5).

Table 2
Paired samples tests

	Baseline (n = 30)	Follow-up (n = 30)	P value	Effect size
HgbA1c [%]#	5.61 ± 0.352	5.65 ± 0.395	0.367	0.1674
HgbA1c [mmol/l]#	37.7 ± 3.74	38.2 ± 4.36	0.331	0.1804
TC [mmol/l]#	5.70 ± 1.14	5.90 ± 1.25	0.338	0.1779
TG [mmol/l]\$	1.81 [1.19–2.50]	1.16 [1.16–2.14]	0.271	−0.2344
HDL [mmol/l]\$	1.29 [1.12–1.52]	1.11 [1.11–1.61]	0.593	0.1140
LDL [mmol/l]#	3.44 ± 1.04	3.67 ± 1.23	0.247	0.2156
Weight [kg]#	87.6 ± 16.9	86.1 ± 16.4	0.022*	−0.4406
BMI [kg/m2]\$	32.0 [26.4–34.6]	27.0 [27.0–35.0]	0.054	−0.4115
LBM [kg]#	54.2 ± 12.1	56.0 ± 11.1	0.181	0.2505
MBF [kg]#	32.0 ± 9.72	30.2 ± 9.05	< 0.001*	−0.7617
SLM [kg]#	50.7 ± 10.1	51.2 ± 10.3	0.080	0.3312
SMM [kg]#	30.4 ± 6.07	30.7 ± 6.21	0.076	0.3357
Mineral [kg]#	4.93 ± 0.937	4.86 ± 0.926	0.055	−0.3651
Protein [kg]\$	10.0 [8.67–12.0]	8.90 [8.90–12.4]	0.026*	0.4759
TBW [kg]#	40.0 ± 7.87	40.4 ± 8.03	0.141	0.2764
PBF [%]#	36.1 ± 6.86	34.7 ± 6.58	< 0.001*	−0.8971
VFL [Units]\$	16.0 [13.0–18.0]	12.3 [12.3–17.0]	< 0.001*	−0.87619
VFA [cm2]#	169 ± 79.1	148 ± 69.1	< 0.001*	−0.9162
WHR#	0.948 ± 0.0912	0.926 ± 0.0864	< 0.001*	−0.7798
AC [cm]#	98.1 ± 13.0	95.9 ± 12.2	< 0.001*	−0.7618
SLM Lt. ARM [kg]#	3.50 ± 0.718	3.51 ± 0.728	0.751	0.0584
SLM Rt. ARM [kg]#	3.52 ± 0.736	3.53 ± 0.745	0.795	0.0478
SLM Lt. LEG [kg]#	9.07 ± 2.17	9.41 ± 1.97	0.095	0.3155
SLM Rt. LEG [kg]#	9.32 ± 1.90	9.51 ± 1.98	0.008*	0.5158
SLM Trunk [kg]#	25.1 ± 4.96	25.2 ± 5.02	0.258	0.2105
MBF Lt. ARM [kg]#	1.93 ± 0.612	1.86 ± 0.586	0.008*	−0.5227
MBF Rt. ARM [kg]#	1.91 ± 0.625	1.82 ± 0.603	0.005*	−0.5518
MBF Lt. LEG [kg]#	5.84 ± 1.75	5.50 ± 1.62	< 0.001*	−0.7810
MBF Rt. LEG [kg]#	5.83 ± 1.76	5.47 ± 1.62	< 0.001*	−0.8146
MBF Trunk [kg]#	16.5 ± 4.98	15.5 ± 4.65	< 0.001*	−0.7666
E.C.W/T.B.W#	0.401 ± 0.008	0.399 ± 0.00839	0.052	−0.3700

Values are presented as mean ± SD or median [Q1–Q3]. *Two-sided P value < 0.05.

Paired t-test, Cohen's d (effect size). \$ Wilcoxon's matched pairs signed rank test, Rank biserial correlation (effect size).

Statistically significant negative correlations between changes (delta = follow-up – baseline) of lipid values and variability of Steps count, distance and minutes of very active physical activity were found, showing a decrease of lipid levels with higher long-term variability and area of an ellipse, but not short-term variability. Additionally, decreased body composition measurements were associated with minutes being fairly or very active. However, the SD1/SD2 ratio showed an opposite relationship with body composition measures indicating that more balanced long-term and short-term variability and possibly higher short-term variability could be associated with positive changes. Also, a similar association between HbA1c mmol/l and Steps and Distance SD1/SD2 ratio was observed.

3.4. Vision of remote patient physical monitoring system

In previous paragraphs, we have described in detail how data were collected and analysed. However, many data operations (extraction, integration and analysis) were done manually. We have a vision to improve the proposed remote patient physical monitoring system in order the system could be used for

Table 3
Fitbit parameters and calculated corresponding physical activity variability parameters

Fitbit variable	Parameter	Mean \pm SD	Median [Q1–Q3]	<i>P</i> value [#]
Steps per day	SD1	3457 \pm 1120	3375 [2601–4157]	0.631
	SD2	4441 \pm 1462	4176 [3568–5342]	0.063
	SD1/SD2	0.791 \pm 0.164	0.707 [0.680–0.933]	0.002
	AFE	5.22e+7 \pm 3.33e+7	4.68e+7 [2.92e+7–6.40e+7]	0.001
	AVG	8924 \pm 3050	9388 [6548–11410]	0.087
	SD	3997 \pm 1245	3875 [3149–4575]	0.218
Distance [km/day]	SD1	2.37 \pm 0.842	2.33 [1.77–2.90]	0.348
	SD2	3.05 \pm 1.10	2.92 [2.21–3.70]	0.118
	SD1/SD2	0.790 \pm 0.163	0.709 [0.680–0.932]	0.002
	AFE	25.0 \pm 17.4	20.6 [13.0–32.7]	0.001
	AVG	6.18 \pm 2.09	6.68 [4.57–8.06]	0.117
	SD	2.74 \pm 0.946	2.60 [2.06–3.35]	0.272
Minutes lightly active [min/day]	SD1	70.0 \pm 29.1	59.6 [50.1–75.0]	< 0.001
	SD2	108 \pm 48.2	88.1 [72.7–137]	0.001
	SD1/SD2	0.697 \pm 0.209	0.696 [0.593–0.797]	0.691
	AFE	26158 \pm 20444	16300 [12568–33848]	< 0.001
	AVG	245 \pm 95.9	234 [191–315]	0.889
	SD	92.0 \pm 37.1	75.0 [65.2–129]	< 0.001
Minutes fairly active [min/day]	SD1	18.6 \pm 7.53	17.1 [14.8–20.6]	0.086
	SD2	23.2 \pm 9.69	21.8 [18.1–25.5]	0.027
	SD1/SD2	0.819 \pm 0.141	0.811 [0.768–0.890]	0.200
	AFE	1558 \pm 1291	1156 [782–1597]	< 0.001
	AVG	19.0 \pm 11.3	17.3 [9.42–24.8]	0.212
	SD	21.1 \pm 8.52	19.4 [16.0–23.1]	0.026
Minutes very active [min/day]	SD1	14.5 \pm 8.27	12.0 [9.24–17.7]	0.002
	SD2	16.7 \pm 8.55	14.1 [11.4–23.3]	0.013
	SD1/SD2	0.870 \pm 0.151	0.873 [0.768–0.917]	0.065
	AFE	957 \pm 954	507 [338–1472]	< 0.001
	AVG	13.0 \pm 9.86	11.3 [6.43–13.5]	< 0.001
	SD	15.7 \pm 8.28	12.9 [10.5–21.7]	0.004

[#]Shapiro-Wilk test.

all patients and we could remotely monitor patient health on a day-to-day basis. In this section, we will describe a vision of an integrated remote patient physical monitoring system compatible with existing SANTA-HIS and Lithuanian national eHealth framework.

VUHSK has developed hospital information system (SANTA-HIS), which integrates electronic health record, laboratory, picture archiving and communication system (PACS), registry of informed consent, staff and resource management, document management, quality management system, adverse events management system and many other systems that are necessary for effective health care services. SANTA-HIS system is integrated with Lithuanian national health information system which is called Electronic Health Services and Cooperation Infrastructure Information System (ESPBI IS). ESPBI IS was launched for patients and health care practitioners from 2015 and already has huge amounts of clinical data.

Figure 6 shows the vision of a remote patient physical monitoring system. The elements of the systems, which are shown in green, are already developed and used on a daily basis by health care practitioners.

Remote patient physical monitoring system and external medical devices integration system (shown in red) should be implemented and integrated into existing SANTA-HIS in order to automate data operations.

We strongly believe that we can benefit from the systems integrations because we will have all needed data in one place, and we can make detailed analyses and have a clear picture of the patient's health.

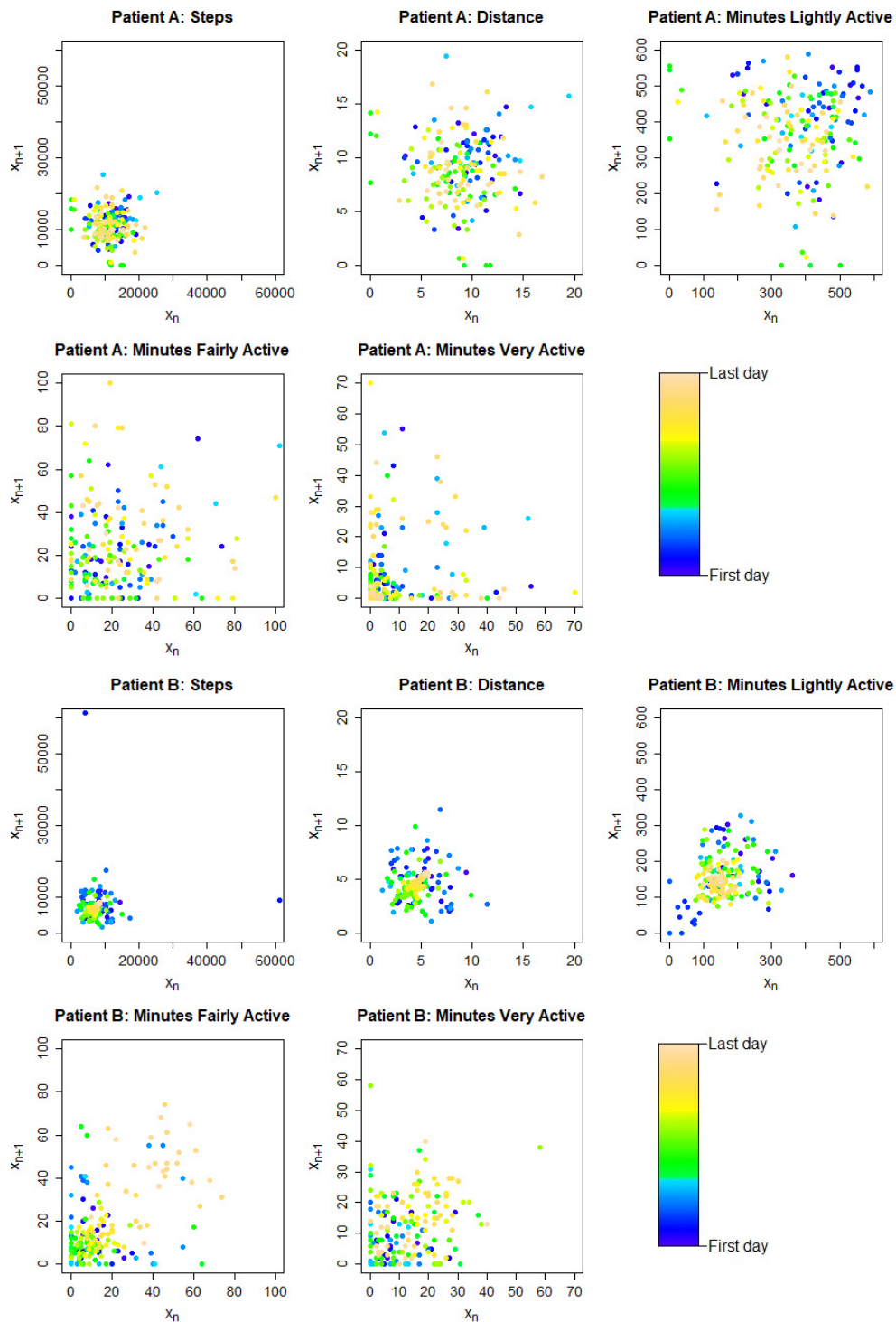


Fig. 4. Examples of the Poincaré plot for Fitbit parameters. Patient A – male, age 42 years and patient B – female, age 65 years.

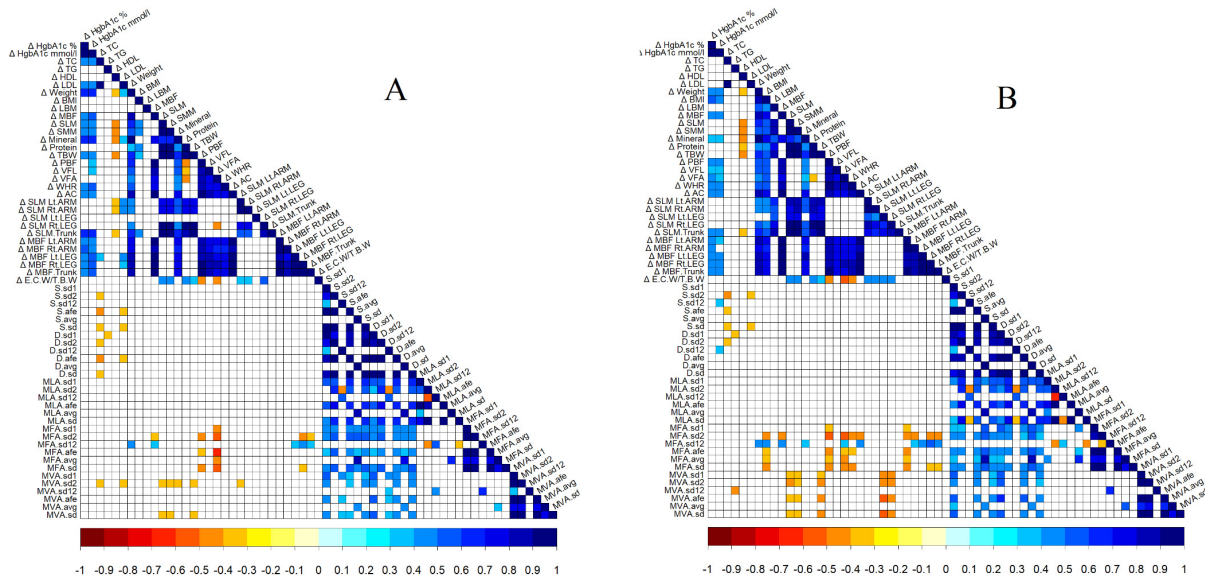


Fig. 5. Heatmap of Pearson's (A) and Spearman's correlation (B) coefficients. Non-significant associations were shown as white spaces.

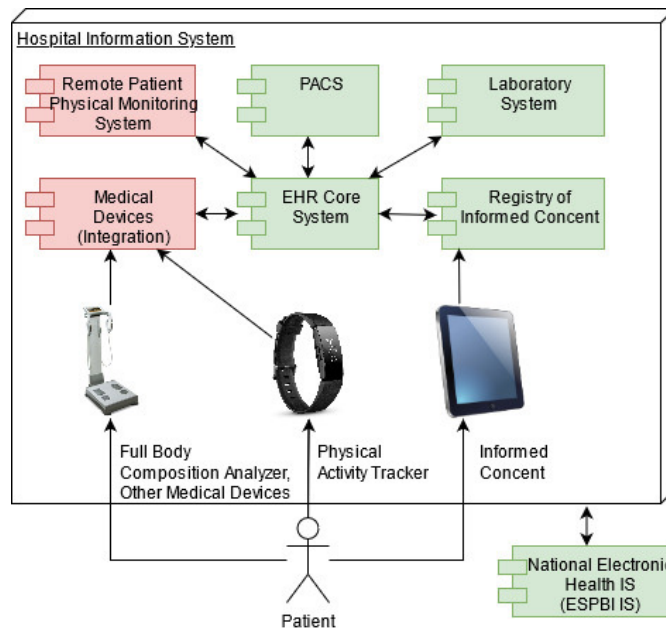


Fig. 6. Vision of remote patient physical monitoring system.

Systems integration will let health care practitioners to review and use patient data collected in other clinics and laboratories. Objective patient physical activity data and other parameters are essential in clinical work because it helps us know how the patient feels under real-life conditions and adheres doctor's recommendations. We can find out why a patient can't reach those goals and help solve the problem faster and more effectively.

4. Conclusions

Nonlinear physical activity variability parameters showed potential to include variability measurement in monitoring patients with prediabetic status. Using these parameters, we can visualize physical activity variability (Poincaré plot). Implementation architecture suggested in our paper can be used for implementation of these parameters into clinical practice and improve telemonitoring availability for physicians. In future research, the target value for variability in daily steps and other physical activity parameters should be evaluated not only in the larger group of prediabetic patients but also in healthy subjects to understand the impact on clinical parameters and health improvement.

Conflict of interest

None to report.

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