

The Active and Healthy Aging Study (Dutch: Actief en Gezond Oud Studie).

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Internet-based coaching combined with accelerometry of inactive individuals aged 60-70 years is effective in increasing physical activity, and this increase in physical activity associates with enhanced 24-hour glucose regulation.

Ethische beoordeling	Positief advies
Status	Werving gestopt
Type aandoening	-
Onderzoekstype	Interventie onderzoek

Samenvatting

ID

NL-OMON27923

Bron

Nationaal Trial Register

Verkorte titel

AGO Study

Aandoening

Inactive individuals aged 60 to 70 years, without contra-indications to increase their physical activity.

Keywords: glucose metabolism, aging, physical activity, lifestyle intervention, web-based, self-monitoring

Dutch: glucoseregulatie, veroudering, fysieke activiteit, lifestyle-interventie, thuismonitoring

Ondersteuning

Primaire sponsor: Leiden University Medical Center

Overige ondersteuning: Netherlands Consortium for Healthy Ageing
Philips Health Care

Onderzoeksproduct en/of interventie

Uitkomstmaten

Primaire uitkomstmaten

Level of daily activity, as measured using independent tri-axial accelerometers. These accelerometers have been validated against energy expenditure measured using VO₂-max. We will use this independent measure of activity, because:

- A. The DirectLife activity monitor is part of the intervention therefore not independent not suitable as measure of end point;
- B. The Genea accelerometer yields more detailed data on physical activity.

Toelichting onderzoek

Achtergrond van het onderzoek

With an increasingly ageing population, there is a need for lifestyle interventions that can be implemented at a population level to improve health in old age and increase healthy lifespan in the general population. Healthy longevity has been associated with enhanced glucose tolerance and insulin sensitivity, characteristics that can be modified by increasing physical activity. The Philips Directlife program is a web-based lifestyle intervention directed at increasing daily physical activity consisting of 1) an activity monitor, 2) a personal website, and 3) a personal e-coach. Glucose metabolism can be measured with minimal invasive measures and sensitively using a 24-hour glucose sensor (continuous glucose measurement, CGM) placed subcutaneously in the abdomen. This study will test the following hypotheses:

Primary: The DirectLife program is effective in increasing the level of physical activity in an ageing population.

Secondary:

1. The level of physical activity associates with parameters of 24-hour glucose rhythms in an ageing population;
2. An increase in physical activity associates with beneficial changes in 24-hour glucose rhythms as measured with CGM, and with changes in metabolic parameters, anthropometry, well-being and sleep quality.

We will perform a randomized controlled trial in 316 participants aged 60-70 year with sedentary lifestyle. We will apply the Philips Directlife program as intervention vs. no intervention in the control group. The primary outcome is change in daily physical activity, as measured by independent tri-axial accelerometers. Secondary outcome parameters include parameters measured during 24-hour glucose rhythms, such as mean glucose, standard deviation of the mean, postprandial glucose rhythms etc., and metabolic parameters, well-being and sleep quality. The overview of the study is shown in figure 1. Shortly, participants

are recruited from the general population and screened for physical activity level using questionnaire. The study outline comprises two visits to the study center. The primary visit is at $t=0$ weeks during which baseline anthropometric measurements and well-being are assessed and the CGM sensor applied which is worn for 4 days. The second visit is at $t=14$ weeks, after which outcome parameters are assessed and CGM applied for the second time to be worn for four days. After $t=14$ weeks, the control group will get access to the Philips DirectLife program after the study has ended.

Doel van het onderzoek

Internet-based coaching combined with accelerometry of inactive individuals aged 60-70 years is effective in increasing physical activity, and this increase in physical activity associates with enhanced 24-hour glucose regulation.

Onderzoeksopzet

T=0: Baseline visit for assessment of physical activity, glucose monitoring, blood sampling and anthropometrics;

T=1 wk to T=13 wk: Intervention or control period;

T = 13 wks: End visit for assessment of PA, glucose monitoring, blood sampling and anthropometrics.

Onderzoeksproduct en/of interventie

The study is an open, randomized controlled intervention trial. Participants will be randomized to DirectLife intervention or control group. The control group will be offered use of DirectLife after the end of the study, i.e. the second "assessment period", such that all participants will receive an equal amount of study visits, and can both benefit from the DirectLife program.

Philips DirectLife has developed a lifestyle activity intervention program, DirectLife, to increase people's daily life physical activity. Based on health behavior change models (Prochaska et al., 1994; Vries and Mudde, 1998), DirectLife takes into account the current individual activity level, and subsequently provides a personal goal. Very briefly, DirectLife consists of three elements, namely:

1. An activity monitor;
2. A personal website;
3. A personal e-coach, who provides regular updates by e-mail of the physical activity status and gives advice to increase physical activities.

By means of these elements, DirectLife increases awareness about one's own physical activity behavior, gives feedback, and provides support to make sustainable changes in physical activity behavior.

DirectLife has a scientific base. The activity monitor of DirectLife is based on the Tracmor, i.e. a tri-axial accelerometer, which has been reviewed as the most accurate device for the estimation of total daily life energy expenditure (Bonomi et al., 2010; Plasqui et al., 2005; Plasqui and Westerterp, 2007). Moreover, DirectLife combines high-tech with human elements to help people to change to a more active lifestyle (Goris and Holmes, 2008). Participants of the DirectLife program continuously wear an accelerometer throughout the day to measure daily activity. These data are uploaded through the internet. After an initial "assessment period" in which the current level of daily activity is measured, a target is set to increase the level of daily activity during a twelve week web based interactive coaching program. Participants are given a target for daily activity which increases weekly, and data from the accelerometer are used for regularly feedback.

Contactpersonen

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Deelname eisen

Belangrijkste voorwaarden om deel te mogen nemen (Inclusiecriteria)

1. Age 60 to 70 years;
2. Sedentary lifestyle;
3. Motivated to increase physical activity;
4. Familiar with use of and access to personal computer.

Belangrijkste redenen om niet deel te kunnen nemen (Exclusiecriteria)

1. Physical condition that inhibits increase of physical activity;
2. Diabetes mellitus type I or II;
3. Use of glucose lowering medication.

Onderzoeksopzet

Opzet

Type:	Interventie onderzoek
Onderzoeksmodel:	Parallel
Toewijzing:	Gerandomiseerd
Blinding:	Open / niet geblindeerd
Controle:	Geneesmiddel

Deelname

Nederland	
Status:	Werving gestopt
(Verwachte) startdatum:	01-11-2011
Aantal proefpersonen:	232
Type:	Werkelijke startdatum

Ethische beoordeling

Positief advies

Datum: 24-08-2011

Soort: Eerste indiening

Registraties

Opgevolgd door onderstaande (mogelijk meer actuele) registratie

Geen registraties gevonden.

Andere (mogelijk minder actuele) registraties in dit register

Geen registraties gevonden.

In overige registers

Register	ID
NTR-new	NL2899
NTR-old	NTR3045
Ander register	METC LUMC : P10.220
ISRCTN	ISRCTN wordt niet meer aangevraagd.

Resultaten

Samenvatting resultaten

N/A