European Health Examination Survey –
towards a sustainable monitoring system

Hanna Tolonen¹, Päivikki Koponen², Jennifer Mindell³, Satu Männistö¹, Kari Kuklasmaa¹

¹ Department of Chronic Disease Prevention, National Institute for Health and Welfare (THL), Helsinki, Finland

² Department of Health, Functional Capacity and Welfare, National Institute for Health and Welfare (THL), Helsinki, Finland

³ Research Department of Epidemiology and Public Health, University College London (UCL), London, United Kingdom

Corresponding author:
Hanna Tolonen (hanna.tolonen@thl.fi)
National Institute for Health and Welfare (THL)
Department of Chronic Disease Prevention
P.O. Box 30
00271 Helsinki
Finland
Abstract

**Background.** Health examination surveys (HES) including both questionnaire and physical measurements, and in most cases also collection of biological samples, can provide objective health indicators. This information complements data from health interview surveys and administrative registers, and is important for evidence based planning of health policies and prevention activities. HESs are valuable data sources for research. The first national HESs in Europe were conducted in the late 1950’s and early 1960’s. They have recently been carried out in an increasing number of countries, but there has been no joint standardization between the countries.

**Methods.** The European Health Examination Survey (EHES) Pilot Project was conducted in 2009-2012. The EHES Pilot Reference Centre was established and pilot surveys were conducted in twelve countries.

**Results.** European standardized protocols for key measurements on main chronic disease risk factors (height, weight, waist circumference, blood pressure, blood lipids, and fasting glucose or HbA1c) were prepared. European level training and external quality assessment were organized. Although the level of earlier experience, infrastructures, economic status and cultural settings varied between the pilot countries, it was possible to standardize measurements of HESs across the populations. Obtaining high participation rates was challenging.

**Conclusion.** HESs provide high quality and representative population data to support policy decisions and research. For future national HESs, centralized coordination, training and external quality assessment is needed to ensure comparability of the results. Further studies on
effects of different survey methods on comparability of the results and on recruitment and
motivation of survey participants are needed.

**Key words:** health examination survey, measurements, Europe, questionnaire, standardization, risk factors
Introduction

Health examination surveys (HES), including both questionnaire and physical measurements, and in most cases also collection of biological samples such as blood, urine and saliva, are data sources for health indicators. HESs can provide objective measures for many health indicators which are not available from other data sources, including health interview surveys (HIS) or administrative registers. A HIS can be used to measure health behaviours, health status and diseases which are known to the person. It cannot provide indicators on undiagnosed conditions, is subject to reporting bias and cultural based preferences in reporting of health problems. Administrative registers, like hospital records, are limited to those who have been diagnosed with the disease. Administrative registers often have rather limited information on individuals’ background variables such as lifestyle, socio-economic status and family composition. Comparability of data from health care utilization records is low due to differences in the structure of health care systems and recording practices.

The first national HESs on adults in Europe were conducted in the late 1950’s and early 1960’s (Figure 1).(1, 2) Since then, nationally representative HESs have been carried out in at least 16 EU Member States and EFTA/EEA countries. In additional to these national HESs, in most countries at least some regional, disease-specific or age-specific HESs have been conducted(2). Finland, Germany, Ireland, Italy, Netherlands, Poland, Romania, Slovakia and UK (England and Scotland) have carried out national HESs repeatedly.(2) In the USA, a standardized nationally representative HES has been conducted since the 1960’s (NHANES)(3). Also countries such as Canada(4), Mexico(5) and Korea(6) have conducted national HESs repeatedly.
Comparability of the results from national HESs has been limited due to differences in survey methods used in these surveys. For example, for weight measurement the removal of clothing differs between surveys. In some surveys participants have been measured on their underwear and in other surveys they are asked to remove only heavy outer garments. Accurate and comparable HES results require proper standardization of the measurement protocols and devices used, thorough training of the survey personnel and adequate quality control. For reporting, common definition of indicators is also required.

Standardized measurement protocols for cardiovascular disease risk factors were introduced by G. Rose in 1968(7); they have been widely used by individual surveys. In the 1980’s and 1990’s, the World Health Organization (WHO) MONICA Project coordinated standardized cross-sectional cardiovascular disease risk factor surveys in 38 populations in 21 countries.(8) These surveys were regional and therefore not nationally representative. Since the mid 1990’s, when the WHO MONICA Project ended, there has been no cross-national standardization for HESs in Europe. The WHO has prepared the Stepwise Approach for Risk Factor Surveillance (STEPS), a standardized HES protocol which has been mainly targeted for low and middle income countries.(9)

The Feasibility of the European Health Examination Survey (FEHES) Project was conducted in 2006-2008 to assess the feasibility of conducting a standardized HES in all EU Member States and EFTA/EEA countries.(2, 10, 11) A strong demand for international standardization of the HES methods and for international coordination to ensure comparability of the national results was expressed by national experts of the FEHES network.

The FEHES Project concluded that it is feasible to conduct standardized national HESs in Europe and recommended setting up the European HES in two phases:(10)
• Phase I: setting up the central coordination, and the planning and preparation of national HESs, including pilot surveys, in the first 8-12 countries.

• Phase II: full-size national HESs in those first countries, while other countries start planning and preparing for their national HES.

This paper describes how the European Health Examination Survey (EHES) Pilot Project was established, and tools developed to facilitate the standardization of HESs. Basic results from the EHES pilot surveys, relating to the organization and contents of the surveys, are also presented.

**Establishing the EHES Pilot Project**

Following the recommendations from the FEHES project, the setting up the European HES started in autumn 2009 as the EHES Pilot Project. The first step was to establish the EHES Pilot Reference Centre (EHES RC). It was a joint operation of the National Institute for Health and Welfare (THL), Finland; Statistics Norway (SSB), Norway; and Istituto Superiore di Sanità (ISS), Italy. It received funding from the European Commission/DG Sanco for 2009-2012.

The EHES RC is a coordinating body for the EHES. It is responsible for

• the development of the European level standardized protocols for HESs and publication of those in the EHES Manual;

• providing professional support to the countries planning a national HES;

• developing a training programme, preparing training materials and organizing European level training seminars;
• coordinating external quality control and preparing guidelines for internal quality control; and
• establishing a centralized data management and reporting system for basic survey data.

In the beginning of 2010, 14 countries preparing for a national HES joined the EHES Pilot Project. The aim of the EHES Pilot Project was to prepare for a full-scale HES in these 14 countries and to pilot the fieldwork, data collection and assessment, and reporting. Twelve of them completed the requirements of the Project, including successful pilot surveys (Figure 2). In four of the pilot countries (Italy, Germany, Netherlands, and UK/England), a full-size national HES started already before or early in the Pilot Project. In these countries, the purpose of pilot survey was to evaluate the feasibility of incorporating EHES standardized protocols into the existing HES system. In remaining eight other countries, the feasibility of the EHES standards was tested in small pilot surveys. The pilot surveys received funding from the European Commission/DG Sanco for periods in 2010-2011.(12)

**Tools developed for the EHES**

**The EHES Manual**

To minimize variation due to differences in measurement protocols, and also more widely in the survey processes, standardized measurement protocols and guidelines for planning the survey, implementation of the fieldwork, and reporting were prepared and documented in the EHES Manual.(13-15) The EHES Manual includes information on sampling, ethical issues, organization of the fieldwork, protocols of the measurements, definition of indicators for reporting, and other details relating to the planning and conducting of a national HES.
EHES standardized protocols and recommendations are based on review of existing international guidelines, protocols used and experience from the previous national HESs, and scientific literature. Final versions were updated based on experience gained during the EHES pilot project.

Detailed standardized measurement procedures should be followed without national modifications. However, for topics such as sampling and ethical conduct, national circumstances, legislation and regulations vary between countries. Therefore, the EHES Manual cannot provide only one solution but it lists the critical issues and provides guidelines and potential solutions.

*Recommended measurements*

In the EHES Pilot Project, standardization of the physical measurements and collection of biological samples was limited to key measurements of major chronic disease risk factors. For countries without previous experience of HES, it is recommended to start with few important measurements and do them well, then expand the set of measurements after they have acquired more experience in conducting basic HESs.

The EHES Manual has a core set of measurements: height, weight, waist circumference, blood pressure, blood lipids, fasting blood glucose or glycated haemoglobin (HbA1c) and a questionnaire which should be included to all national HESs. The inclusion of these measurements is justified by their role as major risk factors for several chronic diseases and limited availability through other data sources. The accuracy of these EHES core measurements is known to be sensitive to deviations in the measurement procedures (16-24), which also supports the need for standardization. The EHES Manual (13) provides detailed measurement protocols for all core measurements.
A set of core questions was selected to be included in all national HESs (13), including questions on socio-economic position, health status, health care use, and smoking. The questionnaire covers items which are important to know from the same individuals on whom the physical measurements are made and whose biological samples are collected. Whenever feasible, the questions were selected from the European Health Interview Survey (EHIS) (25).

*Training programme*

For fieldwork personnel conducting the measurements, learning the standardized measurement protocols is the first step. Training should also include teaching practical skills to be able to follow the protocols correctly.

A training programme was developed in the EHES Pilot Project (14). It includes European level training and outlines the issues which should be included in the training of national survey personnel. A set of training material was prepared and made publicly available (26). The training material includes guidelines for selection of the population sample, recruitment of survey participants, informed consents, measurement protocols for anthropometric measurements and blood pressure, blood sample collection and handling, and questionnaire administration. Questionnaire administration includes guidelines how to check self-administered questionnaire and conduct an interview.

The aim of the European level training is to train the trainers for each country. These national trainers would then be responsible for training within their own country. The main tools for the European level training are training seminars. During the EHES Pilot Project, two European level training seminars were organized: one on planning and preparation for the EHES at the national level, and another one on fieldwork for the national HES. For full-scale
national HESs, the need for the third training seminar on data analysis, reporting and dissemination of the survey results was identified.

**Quality control**

In EHES, special attention is given to quality control. Instructions on how to organize quality control are given in the EHES Manual (14). They include guidelines for organizing internal quality control and external quality assessment by the EHES RC. The external quality assessment includes review of the national HES manuals, site visits, assessment of collected individual level data and external laboratory quality assessment.

**Results of the pilot surveys**

During the EHES Pilot Project, the countries planned their national HES and prepared their own national HES manuals. A national HES manual template was prepared by the EHES RC to support the preparation of national HES manuals (27). The pilot surveys tested local survey protocols in a small fieldwork pilot of people aged 25-64 years. A minimum sample size (number of persons examined) for the pilot surveys was 200 individuals. The average participation rate for the pilot surveys was 45% varying from 25% to 63%. Participation rates were lower in men than in women in all but one survey.

The way the health examinations were organized varied between surveys. In one survey, all the examinations were conducted at the home of the participants and in another survey they took place at a mobile examination unit (a specially equipped bus). The other ten surveys used a series of fixed local settings for the health examinations. These were either clinical settings in local health care centres or examination centres especially set up for this purpose in other premises such as community houses, and office spaces. In three surveys, using fixed
examination centres, home visits were offered for those not able or not willing to come to the examination centre. The frequency of home visits in these surveys varied. In two countries home visits were not considered feasible, based on previous experience that people were reluctant to let survey fieldwork personnel into their home, or access to the apartment buildings in the cities would be limited due to locked front doors. (Table 1)

In each pilot country, training was organized nationally for the fieldwork personnel involved in the pilot survey. Those responsible for the national training had participated in the training seminars organized by the EHES RC. The duration of the national training varied from one day to three weeks, depending on the previous experience of the fieldwork personnel and number and type of measurements included in the survey. (Table 1.) In all countries, the fieldwork personnel went through the survey protocol, measurement techniques were demonstrated by trainers, and fieldwork personnel also practiced measurements on volunteers during the training sessions. Training was seen as an important part of the standardization and quality assurance for the surveys. In several countries it was noted that during the training more emphasis should be given for obtaining informed consent, the scope and objectives of the survey, and the content of the questionnaire. Practical hands-on training with volunteers was also considered important. For small pilot surveys, the duration of training was felt to be adequate but for future full-size HESs, several countries felt that longer training periods would be needed.

Of the recommended EHES core measurement, height, weight, waist circumference, blood pressure, and total and HDL cholesterol were included in all EHES pilot surveys. Fasting glucose was measured in seven surveys, in one survey HbA1c was measured, and in four surveys both fasting glucose and HbA1c measurements were included.
At least one additional measurement was included in eleven pilot surveys. Additional measurements varied a lot, including hip circumference, spirometry, body fat measurement by bioimpedance, visual acuity, electrocardiography (ECG), several functional capacity tests, oral health examination, and cognitive function tests. The most common additional measurement was hip circumference which was included in seven surveys. Body fat measurement and spirometry were also undertaken in more than one survey. (Table 2.)

Urine (24 hour or spot) samples were collected in more than one survey and saliva samples in one survey. Most surveys also collected more blood than was required for EHES core measurements of lipids and glucose. From these additional blood samples triglycerides, c-reactive protein, gamma-glutamyl transferase (GGT) and number of other biological markers were analysed. In nine surveys, additional blood samples were taken for storage for future analysis. (Table 2.)

In two pilot surveys questionnaires were filled-in during the interview, in five surveys questionnaires were self-administered, and in five surveys both interview and self-administration was used. In most surveys, self-administered questionnaires were sent to the survey invitees together with the invitation and they were asked to fill them in before coming to the examination. Questionnaires were checked at the examination centre for completeness. (Table 3.)

Questionnaires used in the pilot surveys were not identical. In the countries with an ongoing national HES or existing HES systems there was a need to maintain questions used in previous surveys to ensure valid trend estimates. Also the development of the EHES core questionnaire was not finished by the time that the first pilot surveys got started, resulting in small deviations between the EHES core questionnaire and those used in the surveys.
Deviations were usually related to specifications in the questions such as ‘diagnosed by a medical doctor’ or ‘during the past two weeks’.

Background items of sex, date of birth, education, labour status, household size and income; and health status and smoking questions were included in all surveys except for occasional questions which were missing from one or two surveys. Health care questions were missing from about half the surveys.

Each survey included some additional questions. Most commonly sets of additional questions were related to diet and alcohol use, physical activity and/or medications used. Questions about social support were also included in several surveys. (Table 3.)

During the EHES Pilot Project, the EHES RC conducted site visits to evaluate the pilot surveys. The local fieldwork personnel were observed taking the measurements (with the consent of the person being examined). The staff involved in the planning and organising of the survey were interviewed. Feedback was provided immediately on site and all observations were documented in the site visit reports. Some minor deviations from the EHES measurement protocols were observed during the site visits, and were usually corrected immediately after feedback. Deviations were usually related to the posture of the subject during the blood pressure measurement (legs crossed or unsupported arm) or failure to use a step when the height was measured of a participant who was taller than person making the measurement. The site visits also provided valuable information for the EHES RC on cultural norms affecting the measurement protocols, such as the extent of undressing for the anthropometric measurements.

Discussion
The major chronic diseases are leading causes of death and disability in Europe. Most of these have common risk factors: obesity, hypertension, elevated blood lipids and smoking. These risk factors are modifiable and therefore a large proportion of premature deaths and complications causing work disability could be prevented. Nevertheless, Organisation for Economic Co-operation and Development (OECD) data shows that 97% of the annual health expenditure goes for healthcare and treatment of diseases, and only 3% for prevention in the EU Member States.(28) Appropriate and cost-effective targeting of prevention activities and their evaluation require health information, much of which is available only through HES.

The World Health Assembly (WHA65(8)) has adopted a global target of a 25% reduction in premature mortality from non-communicable diseases by 2025.(29) To promote this target, a set of voluntary global targets for the prevention and control of non-communicable diseases have been defined. These targets are based on monitoring a set of 25 health indicators including mean population intake of salt per day, prevalence of raised blood glucose, blood pressure, and total cholesterol, and prevalence of overweight and obesity.(30) National HESs would be needed to fulfil these WHO voluntary global targets. Within Europe, this could be done through the EHES framework.

The EHES Pilot Project has shown that it is feasible to conduct standardized national HESs in countries with different infrastructures, economic status and cultural settings. National HESs with representative samples, through EHES framework can provide reliable and comparable health information on European populations to be used for planning of prevention activities. Currently, the health information provided by EHES is focused on most common chronic disease risk factors but surveys could easily be expanded to cover other health indicators and determinants of health.
The vision of EHES is to become a sustainable system of national HESs, covering all EU Member States and EFTA/EEA countries. It would provide nationally representative, high quality, comparable information to support the planning and evaluation of health policies and prevention activities. Data and biological samples collected through national HESs under the framework of EHES would also support a wide range of research. This is needed for translating the data into interpretable information for the benefit of public health. Further studies are also needed to improving the data collection methods, especially recruitment and motivation of survey participants.

After the EHES pilot phase ended, the established EHES network exists but without centralized coordination. The EHES RC has no funding to continue the coordination and cross-national standardization of national HESs. However, several European countries are now conducting their national HESs and more countries are planning to start their national HES within the next few years. The continuation of the EHES RC activities is needed to ensure that data collected in future HESs in Europe is comparable and can provide best possible evidence-base for health policy decisions.

**Acknowledgement**

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The EHES Pilot Project was made possible through collaboration of the dedicated personnel in the EHES Reference Centre and the EHES Pilot countries (see Annex I).
Conflict of interest

None declared.

Key points (3-5 bullets)

- Reliable and comparable health information is needed for evidence based planning and evaluation of health policies and prevention activities as well as for research.
- Health examination surveys can provide key health data which are not available from other data sources.
- EHES has set up European level support and standardization for national health examination surveys.
- The EHES Pilot Project demonstrated that it is feasible to conduct standardized national HESs in countries with different infrastructures, economic status and cultural settings.
- The sustainability of the EHES Reference Centre is not secured, and therefore the standardization of forthcoming national health examination surveys is endangered.
References

Annex I. Sites and key personnel contributing to the EHES Pilot Project

Czech Republic

National Institute of Public Health, Prague: Ruzena Kubinova, Nada Capkova, Jana Kratenova and Michala Lustigova

Finland


Germany

Robert Koch Institute, Berlin. For the DEGS Study Team: Antje Gösswald, Cornelia Lange, Panagiotis Kamtsiuris.

Greece


Italy

Malta

Department of Health Information & Research, Gwardamangia: Neville Calleja, Dorothy Gauci.

The Netherlands

National Institute of Public Health and the Environment (RIVM), Bilthoven: W.M.Monique Verschuren.

Norway

Norwegian Institute of Public Health: Grethe S. Tell, Patricia Schreuder, Sidsel Graff-Iversen, Nina Hovland;

University of Bergen: Kristin Klock;


Poland


Portugal
Instituto Nacional de Saúde Dr. Ricardo Jorge, Lisbon: Carlos Dias, Ana Paola Gil.

**Slovakia**

Regional Authority of Public Health, Banská Bystrica. Maria Avdicova, Katarina Francisciova, Jana Namesna, Silvia Kontrosova.

**UK**

UCL (University College London), London: Jennifer Mindell, Nicola Shelton, Barbara Carter-Szatynska, Alison Moody;

NatCen Social Research, London: Rachel Craig, Susan Nunn, Deanna Pickup, Chloe Robinson;

The Health and Social Care Information Centre, Leeds: Steve Webster, Victoria Cooper.
Table 1. Examination site, and duration and contents of the training

<table>
<thead>
<tr>
<th>Pilot survey</th>
<th>Examination site</th>
<th>Possibility for home visit offered</th>
<th>Duration of training of the fieldwork staff</th>
<th>Personnel had previous experience on HES</th>
</tr>
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<tbody>
<tr>
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<td>Fixed location</td>
<td>No</td>
<td>1 day</td>
<td>Yes</td>
</tr>
<tr>
<td>DE</td>
<td>Fixed location</td>
<td>No</td>
<td>3 weeks</td>
<td>No</td>
</tr>
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<td>EL</td>
<td>Fixed location</td>
<td>Yes</td>
<td>2 days</td>
<td>Yes, partly</td>
</tr>
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<td>Fixed location</td>
<td>No</td>
<td>2.5 days</td>
<td>Yes</td>
</tr>
<tr>
<td>IT</td>
<td>Fixed location</td>
<td>No</td>
<td>5 days</td>
<td>No</td>
</tr>
<tr>
<td>MT</td>
<td>Fixed location</td>
<td>Yes</td>
<td>5 days</td>
<td>No</td>
</tr>
<tr>
<td>NL</td>
<td>Fixed location</td>
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<td>1 day</td>
<td>No</td>
</tr>
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<td>NO</td>
<td>Mobile unit</td>
<td>No</td>
<td>3 days</td>
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<td>Fixed location</td>
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<td>6 days</td>
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</tr>
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<td>PT</td>
<td>Fixed location</td>
<td>No</td>
<td>4 days</td>
<td>No</td>
</tr>
<tr>
<td>SK</td>
<td>Fixed location</td>
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<td>UK/England</td>
<td>Home</td>
<td>Not relevant</td>
<td>1 day for those with previous experience</td>
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</table>
Table 2. Additional physical measurements, biological samples and laboratory analysis

<table>
<thead>
<tr>
<th>Pilot survey</th>
<th>Additional physical measurements</th>
<th>Additional blood samples stored for future use</th>
<th>Additional biological samples</th>
<th>Additional laboratory analysis*</th>
</tr>
</thead>
<tbody>
<tr>
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<td>None</td>
<td>No</td>
<td>None</td>
<td>None</td>
</tr>
<tr>
<td>DE</td>
<td>Hip circumference, Functional capacity tests, Cognitive function tests, SD-sonography</td>
<td>Yes</td>
<td>Spot urine</td>
<td>Allergic sensitization, blood count, calcium, CRP, GGT, GOT (ASAT), GPT (ALAT), Hb, kidney functions, nutrients, triglycerides, rheumatic diseases</td>
</tr>
<tr>
<td>EL</td>
<td>Hip circumference, Body fat</td>
<td>Yes</td>
<td>None</td>
<td>ALT, AST, creatinine, total protein, totals CA²⁺⁺, urine acid</td>
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<tr>
<td>FI</td>
<td>Hip circumference, Body fat</td>
<td>Yes</td>
<td>None</td>
<td>Apo-A1, Apo-B, blood count, CRP, DNA, GGT, RNA, triglycerides</td>
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<td>IT</td>
<td>ECG, Spirometry, Carbon monoxide, Bone density</td>
<td>Yes</td>
<td>24-hour urine</td>
<td>Creatinine (blood), hemocrome, sodium and potassium excretion, triglycerides, urine creatinine and albumin</td>
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<td>Spirometry, Visual acuity</td>
<td>Yes</td>
<td>None</td>
<td>Lead</td>
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<tr>
<td>NL</td>
<td>Hip circumference</td>
<td>Yes</td>
<td>None</td>
<td>Triglycerides</td>
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<tr>
<td>NO</td>
<td>Dental examination</td>
<td>No</td>
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<td>None</td>
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<td>PL</td>
<td>Hip circumference, ECG</td>
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<td>None</td>
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<tr>
<td>Pilot survey</td>
<td>Additional physical measurements</td>
<td>Additional blood samples stored for future use</td>
<td>Additional biological samples</td>
<td>Additional laboratory analysis*</td>
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<td>---------------------------------</td>
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<td>-------------------------------</td>
<td>--------------------------------</td>
</tr>
<tr>
<td>PT</td>
<td>Hip circumference</td>
<td>Yes</td>
<td>None</td>
<td>ALT, AST, blood count, creatinine, CRP, DNA, GGT, triglycerides</td>
</tr>
<tr>
<td>SK</td>
<td>Body fat</td>
<td>No</td>
<td>None</td>
<td>Triglycerides</td>
</tr>
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<td>UK/England</td>
<td>Hip circumference</td>
<td>Yes</td>
<td>Saliva</td>
<td>Cotinine</td>
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</table>

*CRP=c-reactive protein, GGT=gamma-glutamyl transferase, ALT=alanine aminotransferase, AST=aspartate aminotransferase
<table>
<thead>
<tr>
<th>Pilot survey</th>
<th>Questionnaire administration</th>
<th>Additional questions</th>
</tr>
</thead>
<tbody>
<tr>
<td>CZ</td>
<td>Self-administered, checked at the examination</td>
<td>Diet</td>
</tr>
<tr>
<td>DE</td>
<td>Interview and self-administered, checked at the examination</td>
<td>Diet and alcohol, injuries, living environment, mood, pain, sleeping habits/problems, social support, used medications, vision and hearing</td>
</tr>
<tr>
<td>EL</td>
<td>Interview</td>
<td>Diet and alcohol, physical activity</td>
</tr>
<tr>
<td>FI</td>
<td>Self-administered, checked at the examination</td>
<td>Diet and alcohol, family history of chronic conditions, physical activity, quality of life, sleeping habits/problems, used medications</td>
</tr>
<tr>
<td>IT</td>
<td>Interview and self-administered</td>
<td>ADL-IADL, diet and alcohol, family history of cardiovascular diseases, physical activity, self-rated health, used medications</td>
</tr>
<tr>
<td>MT</td>
<td>Interview</td>
<td>Passive smoking i.e. exposure to tobacco smoke indoors</td>
</tr>
<tr>
<td>NL</td>
<td>Self-administered, checked at the examination</td>
<td>Diet and food supplements, mood, physical activity, sedentary activities, social support</td>
</tr>
<tr>
<td>NO</td>
<td>Self-administered, checked at the examination</td>
<td>Alcohol, physical activity</td>
</tr>
<tr>
<td>PL</td>
<td>Interview and self-administered, checked at the examination</td>
<td>Diet and alcohol, depression, physical activity, quality of life, social support, used medications</td>
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<tr>
<td>PT</td>
<td>Interview and self-administered, checked at the examination</td>
<td>Diet and alcohol, exposure to sun, mental health, physical activity, quality of life, social support, used medications</td>
</tr>
<tr>
<td>SK</td>
<td>Self-administered, checked at the examination</td>
<td>Diet, physical activity, stress</td>
</tr>
<tr>
<td>UK/England</td>
<td>Interview and self-administered</td>
<td>Attitudes to personal health and lifestyle, chronic pain, dental health, diet and alcohol, ethnicity, religion, sexual orientation, self-care, social care, used medications, well-being</td>
</tr>
</tbody>
</table>
Figure 1. Time line of national HESs in EU Member States (RO = Romania, FI = Finland, DE = Germany, NL = Netherlands, UK/E = UK/England, NO = Norway, UK/S = UK/Scotland, IT = Italy, CY = Cyprus, IE = Ireland, SK = Slovakia, CZ = Czech Republic, PL = Poland, FR = France, DK = Denmark, ES = Spain)
Figure 2. Components of standardization of HES measurements
Figure 3. Countries completing the fieldwork for the EHES Pilot Project