

REVIEW

ScreenPro FH: From the Czech MedPed to International Collaboration. ScreenPro FH Is a Participating Project of the EAS-FHCS

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Summary

This article describes the evolution of our understanding of familial hypercholesterolemia (FH) in the Central, Eastern, and Southern Europe (CESE) region, and the dissemination of this understanding to other countries. Using the ScreenPro FH project as an example, we would like to illustrate the progression from national objectives, to regional networking and, finally, to international collaboration *via* the Familial Hypercholesterolemia Studies Collaboration (FHSC) project under the leadership of the European Atherosclerosis Society (EAS). It is essential to improve our ability to diagnose FH. In this regard, the EAS and its FHSC project must be commended for their educational and organizational activities which, above all, are dedicated to the creation of a global FH patient registry. In the CESE region, FH diagnostics and treatment situation are markedly different than in Western Europe or North America. Since the Czech MedPed project (Make Early Diagnoses to Prevent Early Deaths in Medical Pedigrees) has been so successful (with results not only comparable to, but, for some parameters, even surpassing the results of many Western countries) we decided to apply the Czech experience to the CESE region. Thus, the ScreenPro FH project was created. The aim of ScreenPro FH is to create a specialist network in the CESE region. The primary objective of the ScreenPro FH project was to dramatically reduce the number of premature deaths due to clinical complications of atherosclerosis in FH patients. At present, ScreenPro FH comprises 18 member countries with a total population of

500,000,000; which, in terms of the FH population, represents 1-2 million patients.

Key words

MedPed • ScreenPro FH • FHSC • Familial hypercholesterolemia • Statin intolerance

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Introduction

At present, FH is primarily (but not exclusively) considered to be a receptor disease. It is a monogenic disease transmitted through autosomal dominant inheritance, and stems from either an LDL-receptor defect, apolipoprotein B mutation (Familial defective APOB-FDB), or PCSK9 (Proprotein convertase subtilisin/kexin type 9) mutations (Innerarity *et al.* 1987, Hobbs *et al.* 1992, Goldstein *et al.* 2001).

FH is an example of a disease that, by its very nature, allows us to study the relationship between lipid metabolism, especially low density lipoprotein cholesterol (LDL-C), and atherosclerosis, as well as

premature manifestation of cardiovascular disease. Unfortunately, and most importantly, it is also the most significant clinical syndrome which, through congenitally elevated LDL-C concentrations, leads to the development of coronary artery disease (CAD) and other manifestations of atherosclerotic vascular disease (ASVD). FH is underestimated, underdiagnosed and undertreated (Nordestgaard *et al.* 2013).

Many international projects have sought to create uniform doctrines for the screening, diagnosis, and treatment of FH patients. However, ultimately these efforts have always been highly individualized, and dependent upon the activities of clinicians.

The Czech Republic's (CZ) MedPed (Make Early Diagnoses to Prevent Early Deaths in Medical Pedigrees) project was first launched in the 1990s. Thanks to the efforts of the Czech Society for Atherosclerosis (www.athero.cz) and several outstanding

personalities, the CZ project continued to carry on successfully even when the MedPed project was no longer active at the international level. As a result of this continued diligence and dedication, the CZ is currently positioned at the leading edge of FH screening and diagnostics. Detailed information regarding the Czech MedPed project has been described by Vrablik *et al.* (2017).

The ScreenPro FH constitution

Based on Czech MedPed experience, the international ScreenPro FH project was established at the end of 2014 in an effort to improve FH screening, diagnostics and treatment within the CESE region. Eighteen countries in the CESE region are active project participants (Fig. 1).



Fig. 1. Map of ScreenPro FH countries (Bosnia and Herzegovina, Bulgaria, Croatia, Czech Republic, Greece, Hungary, Kyrgyzstan, Latvia, Lithuania, Poland, Romania, Russia, Serbia, Slovakia, Slovenia, Turkey, Ukraine, Uzbekistan).

Rationale

Despite recent progress in diagnostics and treatments, FH remains a significant problem and a common risk factor for the premature development of CAD. The status of FH programs in each CESE country is quite different. One important project task is the harmonization and unification of the approaches used in these countries. Countries with more advanced FH screening programs can serve as an example for countries that have only just joined the project. The experience gained from the Czech, Slovakian, and Hungarian

projects can be utilized in other countries where FH has not yet been significantly studied and monitored. One objective that could be very advantageous would be to prepare integrated, unified documents for use in all countries (following translation into the local language). Such documents would include general, scientific, and medical information; as well as forms and other documents (e.g. informed consent) and educational materials for patients. Likewise, it would also be beneficial, if permitted by the various healthcare systems, to adopt the same uniform guidelines for diagnostics and treatment (e.g. FH diagnostic criteria).

Table 1. Dutch lipid clinic network diagnostic criteria for FH.

Criteria	Points
Family history	
i. First-degree relative with CAD, M<55 years, F<60 years	1
ii. First-degree relative with LDL-C above 95 th percentile for a given country	1
iii. First-degree relative with tendinous xanthoma and/or arcus lipoides	2
iv. Children under 18 years with LDL-C above 95 th percentile for a given country	2
Personal clinical history	
i. Premature CAD, M<55 years, F<60 years	2
ii. Premature peripheral or cerebral atherosclerosis, M<55, F<60 years	1
Physical exam	
i. Tendinous xanthoma	6
ii. Arcus lipoides in patients under 45 years	4
Biochemical exam	
LDL-C (mmol/l)	
>8.5	8
6.5-8.4	5
5.0-6.4	3
4.0-4.9	1
Molecular biology – diagnostic mutation testing	
i. Mutations in the genes encoding LDL-R, ApoB or PCSK9	8

CAD – coronary artery disease, M – male, F – female, LDL-C – LDL cholesterol, LDL-R – LDL receptor, ApoB – apolipoprotein B100, PCSK9 – proprotein convertase subtilisin/kexin type 9.

Dutch lipid network criteria (Table 1) (Van Aalst-Cohen *et al.* 2006) are generally accepted criteria for FH diagnostics. Therefore these criteria are recommended to all ScreenPro countries. The Dutch criteria are interpreted as follows: For a definite diagnosis of FH, the subject must have more than 8 points; a probable diagnosis of FH requires 6-8 points; and a possible diagnosis of FH is indicated by 3-5 points. Patients who score 0-2 points most likely do not have FH.

Such guidelines would have to be adopted *via* a consensus; at least, with regard to the selection of basic data that is to be monitored and recorded in national databases. This type of unification and harmonization would facilitate consistent, accurate evaluations of patient findings in a given region.

ScreenPro FH Certificate: All ScreenPro FH Project Leaders have received certificates from the National Centre of the ScreenPro FH Project.

The ScreenPro FH Project structure was established as follows: The leadership – Board of advisors/Administrative council – Country leaders. Regional centers are established in individual countries. A functional network exists in some countries (e.g. Czech

Republic, Slovakia, Poland, Hungary).

ScreenPro FH activities

Website

The ScreenPro FH Project website was developed in a basic format to serve as an operational platform both for the project as well as the participating countries. The website, which primarily focuses on FH, provides educational materials pertaining to dyslipidemias and cardiovascular disease prevention. Participating countries (and any interested colleagues) can also access a short summary describing the diagnostic and therapeutic procedures for FH patients (Cuchel *et al.* 2014, Hopkins *et al.* 2011). The website also offers a photo gallery of patients with xanthomas and other FH manifestations (all images are provided courtesy of the Centre for Preventive Cardiology in Prague).

Results from participating countries are under development and they vary from country to country (i.e. very good in Czech Republic and Slovakia; good in Poland and Hungary, and Greece; and some countries are in the early stages of development, e.g. Uzbekistan,

Kazakhstan). Just for basic information we can provide some data, number of centers at some countries as well as number of patients in local databases: Bosnia and Herzegovina 1 center (900 patients), Croatia 1 center (130 patients), Czech Republic 63 centers (7,001 patients), Greece 9 centers (40 patients), Hungary 22 centers, Kyrgyzstan 3 centers, Latvia 1 center (162 patients), Lithuania 5 centers (500 patients), Poland 9 centers (1,884 patients), Russia 17 centers (700 patients), Slovakia 20 centers (2,000 patients), Turkey 1 center (more than 1,000 patients), Ukraine 5 centers (81 patients) and Uzbekistan 13 centers (46 patients). More preliminary results are available at www.screenprofh.com.

Since statin intolerance is a significant issue in FH treatment (particularly treatment of high risk patients) we decided to include a “statin intolerance” paper in the educational section of the website.

The website is also a source of information regarding project structure, contacts, meetings, etc. The practical tools available for the unification and harmonization of FH patient care are quickly becoming (along with the educational content) the most important section of the website. Practical forms and documents such as FH patient cards, informed consent forms, educational information for patients, and information for families of FH patients, have been prepared and made available to all participating centers.

Database

ScreenPro FH has no ambition to own any country's data. Data from each country is the property of its own national network. The decision of whether, and how, data will be made available to the central registry is determined exclusively by national leadership. That said, the project website outlines what needs to be measured and what data should be collected, but other suggestions are “optional”.

A database is also available that could be used by all countries at a later date (financial limitations mean that some countries only maintain paper records, or local electronic records). However, it is necessary to stress one more time, how important it is that participants use a unified data collection process that allows results from different countries to be combined and compared.

The “Familial Hypercholesterolemia” textbook

Since no medical textbooks on FH had ever been available on the local market, the FH textbook, published

in Czech Republic, was translated into English, which made it accessible to all ScreenPro FH Project county leaders, as well as other interested physicians in the CESE region. This English version of the FH textbook was offered to all country leaders for translation. At present, the FH book is available in 6 languages (Czech, English, Ukrainian, Lithuanian, Serbo-Croatian, Russian); the Russian version, which has the potential to improve accessibility to the text in many countries, is also complete.

The project is under the auspices of the IAS and collaborates with other FH groups: EAS-FHSC. Last year, we started collaboration with the 10 Countries Study and we use the same questionnaire for our FH patients. The questionnaire is widely used in Australia and Asia, and it has been made available on our website for ScreenPro FH Project countries.

Future activities: it is time to get practical

Having had several ScreenPro FH Project Member meetings, we are now focusing on the practical aspects of the project. Our goal is to help member countries increase FH awareness in member countries and assist in the development of educational materials both for clinicians and patients (e.g. informed consent forms, educational materials for patients, questionnaires, etc.).

We have translated the FH patient card, which is important both for patients and physicians. It is advantageous for physicians because it allows them to review a patient's previous treatments and find information about cholesterol levels; and it also benefits patients by increasing awareness of their own health status.

Educational materials: A new list of educational materials will be made available, which includes basic materials for everyday clinical practice that have already been translated into English (e.g. Informed Consent, Letter for Relatives, Information for Patients, an FH Questionnaire, a Questionnaire for GPs, an FH patient Card, etc.).

Another important document that will soon be available is a set of Guidelines for the Establishment of a Lipid Clinic/Center, which details the essential steps needed for best practice care for FH patients.

Laboratory collaboration

ScreenPro FH has accepted a joint consensus statement from the EAS and European Federation of Clinical Chemistry and Laboratory Medicine concerning

lipid profile determination. The paper will be available on the project website, and all country leaders will be made aware of this new consensus statement. This may require new agreements regarding laboratory protocols between ScreenPro FH members and the clinical chemistry and laboratory medicine specialists in their individual countries. Patient organizations play a valuable role in the comprehensive care of FH patients. ScreenPro FH is mapping the situation in individual countries; we want to support those patient organizations that already exist, and encourage the development of patient organizations in countries where they do not exist yet. In some cases, it may be possible for patients to join a “regional or international” patient organization, when local organizations are not available; which can provide education and support for FH patients and their families.

Conclusion

To summarize: 1) a basic ScreenPro FH project

framework exists; 2) 18 countries from the CESE region are actively involved in the ScreenPro FH Project; and 3) a network of lipid centers is under development in all participating countries. Moreover, local national databases are growing. The website is now functional and available to project members, as well as other interested medical specialists. ScreenPro FH Project resources (such as the educational content of the website and the availability of the FH textbook in English and other languages), regular country project-leader meetings, and FH awareness itself, are all increasing in the CESE region.

Conflict of Interest

There is no conflict of interest.

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