

The background of the slide features a large, stylized DNA double helix structure in shades of blue. A globe is positioned in the center, partially obscured by the DNA helix. To the left of the main text, there is a smaller, wireframe globe showing the continents of Europe and Africa.

ESPT

European Society of Pharmacogenomics and Personalised Therapy
A Scientific Society for Individualised Medicine

European **S**ociety of
Pharmacogenomics and
personalised **T**herapy



www.ESPTnet.eu

ESPT creation & development

5th Santorini Conference October 2010: first plans



Prof Dr Gerard Siest
First ESPT President
(2011 – 2016)

ESPT board - 2011

Gerard Siest (FRA) (president)
Janja Marc (SLO) (secretary)
Adriano Henney (UK) (treasurer)
Sophia Siest (FRA)

Ron van Schaik (NL)
Vangelis Manolopoulos (GRE)
Adrian Llerena (ESP)
George Patrinos (GRE)
Peter Jacobs (BEL)

“It was proposed that a scientific European organization be developed to realize independent studies and clinical trials and to participate in educational efforts”

Vangelis G Manolopoulos, Bryan Dechairo, Alain Huriez, Alexander Kühn, Adrián Llerena, Ron H van Schaik, Kiang-Teck J Yeo, Georgia Ragia & Gerard Siest. Pharmacogenomics and Personalised medicine in clinical practice *Pharmacogenomics* (2011) **12**(5)



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ESPT creation & development



*Prof Dr Ron van Schaik
Second ESPT President
(2016 – 2020)*

ESPT board - 2017

*Ron van Schaik (NL) - president
Vangelis Manolopoulos (GRE) - VP
Sanja Stankovic (SER) - secretary
Ingolf Cascobi (GER) - treasurer
Sophia Siest (FRA)*

*Maurizio Simmaco (ITA)
Csilla Sipekey (FIN)
Adrian Llerena (ESP)
Janja Marc (SLO)
Marc Ansari (CH)*

Aims of ESPT

To extend knowledge and facilitate the clinical implementation of Pharmacogenomics and Personalised Medicine in Europe

- To improve understanding of **clinicians and patients** on use of PGx
- To promote, inform and offer an independent view of PGx and Personalised medicine to **regulators**.
- To be a partner for **industry** in strategical discussions related to the clinically implementation of Personalised Medicine
- To offer a platform for **researchers** on PGx for interaction, exchange of ideas, sharing research outcomes and facilitating formation of relevant research consortia and research proposals



ESPT

European Society of Pharmacogenomics and Personalised Therapy
A Scientific Society for Individualised Medicine

Organisation



National PGx Societies linked to ESPT

Country	Name of the Society	Contact
Italy	SIMeP – Italian Society of Personalised Medicine	Maurizio Simmaco
Turkey	Turkish Pharmacogenetics Society	Belgin Susleyici
Greece	Greek Pharmacogenomics and Personalised Society	Drakoulis / Haliassos
Hungary	Hungarian Society of Personalised Medicine	Gyorgy Németh
Spain	Spanish Society of Pharmacogenetics and Pharmacogenomics	Adrian LLerena
Serbia	Serbian Society of Pharmacogenomics and Theranostics	Sanja Stankovic
Israel	Israel Society of Clinical Pharmacology	Mati Berkovich
France	Personalised Therapy and Pharmacogenomics Group	Frederic Libert
Netherlands	Dutch Clinical Pharmacogenetics Network	Ron van Schaik
Finland	Finnish Phamacogenetics Society	Mikko Niemi
Switzerland	Swiss Network Pharmacogenomics and Personalised Medicine	Marc Ansari
Denmark	Danish Network Pharmacogenomics and Personalised Medicine	Ivan Brandslund

ESPT Advisory Board

Philippe Beaune, France
Juergen Brockmöller, Germany
Roland Bühlmann, Switzerland
Pierre Chambon, France
Anne K.Daly , UK
Panos Deloukas, UK
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Paolo Fortina, USA
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Munir Pirmohamed , UK
Mary V. Relling, USA
Wolfgang Sadee, USA
Julia Stingl, Germany
Giulio Superti Furga, Austria
Roland Wolf, UK
Ulrich Zanger, Germany

ESPT international activities

6 international congresses (200-350 participants)
3 international summerschools (80- 120 participants)

- 2011:** 1st ESPT conference in Bled – Slovenia
- 2012:** 1st summerschool in Ljubljana - Slovenia (August)
1st ESPT/Santorini combined meeting Thira – Greece (Oct)
- 2013:** 2nd ESPT conference in Lisbon – Portugal
- 2014:** 2nd ESPT summerschool Rome – Italy (August)
2nd ESPT/Santorini combined meeting Thira – Greece (Oct)
- 2015:** 3rd ESPT conference in Budapest - Hungary (Oct 7 – 10)
- 2016:** 3rd ESPT Summerschool Belgrade – Serbia (Aug)
3rd ESPT/Santorini combined meeting Thira – Greece (Oct)

ESPT - Divisions



ESPT Organisation

European Society of Pharmacogenomics and Personalised Therapy
A Scientific Society for Individualised Medicine

**Clinical
Implementation**

Ron van Schaik (NL)

**Scientific
Research**

Marc Ansari (CH)

Education

Janja Marc (SLO)
Vangelis Manolopoulos

**Communications &
External relations**

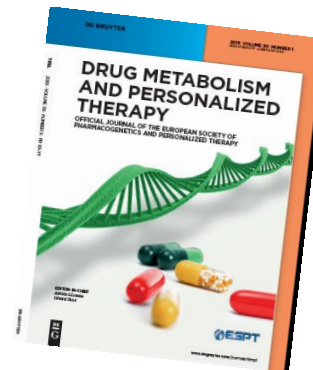
Maurizio Simmaco (ITA)

**Congress
& Meetings**

Sofia Siest (FRA)



Working groups:
Pediatric Oncology
Drug Transporters
Transcription Factors
Drug interactions
PGx for Generic drugs



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European Pharmacogenetics Implementation Consortium

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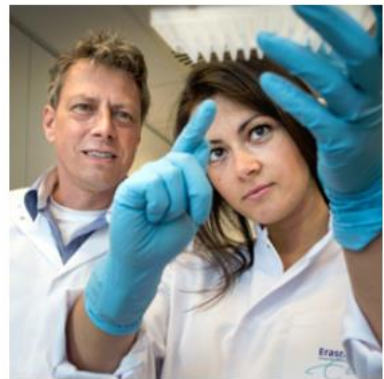
CONTACT

Mission

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Horizon2020



Our mission

To improve patient care in Europe by integrating Pharmacogenetic information in existing health care, personalising drug therapy.



4th ESPT Conference

Catania, Italy
October 4-5-6-7 (2017)

[More info](#)



Eu-PIC symposium May 27, Rotterdam

Eu-PIC successfully organized, together with Golden Helix Foundation, Erasmus MC Rotterdam and the CMBD/NVKG, the 17th Golden Helix Pharmacogenomics Day in Rotterdam, the Netherlands.



Personalised medicine

has the potential to respond to, amongst others, the increasing burden of chronic disease and the complexity of co-morbidities and in doing so contribute to the sustainability of health and care systems. [More info](#)

EU **quality** control program
Ensuring high quality genotyping

EU Network for
exchanging information

**Reimbursement and Regulatory
Aspects Working Group**

Development of **new
genotyping/multiplex
platforms** for cheaper
and faster analyses

Creating **Pharmacogenetic testing
facilities** in 18 EU countries: integration
Of **pre-emptive genotyping** for Cardiology,
Internal Medicine, Psychiatry and Oncology

Dissemination of knowledge
through IFCC, ESPT and national
Pgx networks

Development of a **Decision
Support Tool** and a **safe IT
environment** for reliable and
rapid translation of PGx results
into the clinic

**Evidence-based dosing
recommendations,**
EU-wide, and up-to-date

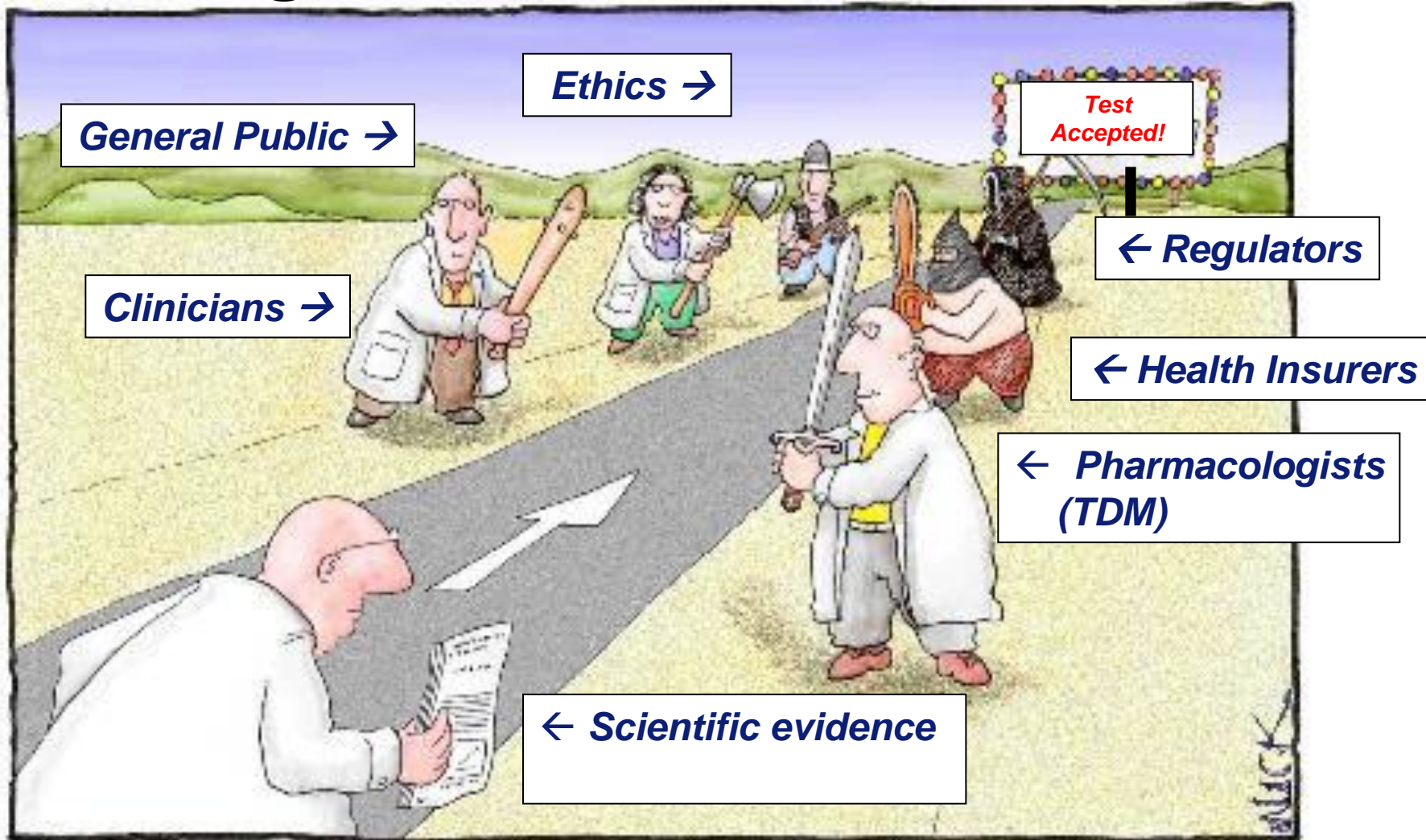
Scientific Trials
for identification of scientific
gaps and design of new projects

**Health Technology Assessment
and Economical Evaluation**
For evaluation of cost/benefit

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Challenges



Pharmacogenomics | **Can We Make It Happen?**

Clopidogrel (Plavix): needs activation by **CYP2C19**

N ENGL J MED 360;4 NEJM.ORG JANUARY 22, 2009

The NEW ENGLAND JOURNAL of MEDICINE

ORIGINAL ARTICLE

Genetic Determinants of Response to Clopidogrel and Cardiovascular Events

Tabassome Simon, M.D., Ph.D., Céline Verstuyft, Pharm.D., Ph.D., Murielle Mary-Krause, Ph.D., Lina Quteineh, M.D., Elodie Drouet, M.Sc., Nicolas Méneveau, M.D., P. Gabriel Steg, M.D., Ph.D., Jean Ferrières, M.D., Nicolas Danchin, M.D., Ph.D., and Laurent Becquemont, M.D., Ph.D., for the French Registry of Acute ST-Elevation and Non-ST-Elevation Myocardial Infarction (FAST-MI) Investigators

CONCLUSIONS

Among patients with an acute myocardial infarction who were receiving clopidogrel, those carrying *CYP2C19* loss-of-function alleles had a higher rate of subsequent cardiovascular events than those who were not. This effect was particularly marked among the patients undergoing percutaneous coronary intervention. (ClinicalTrials.gov number, NCT00673036.)

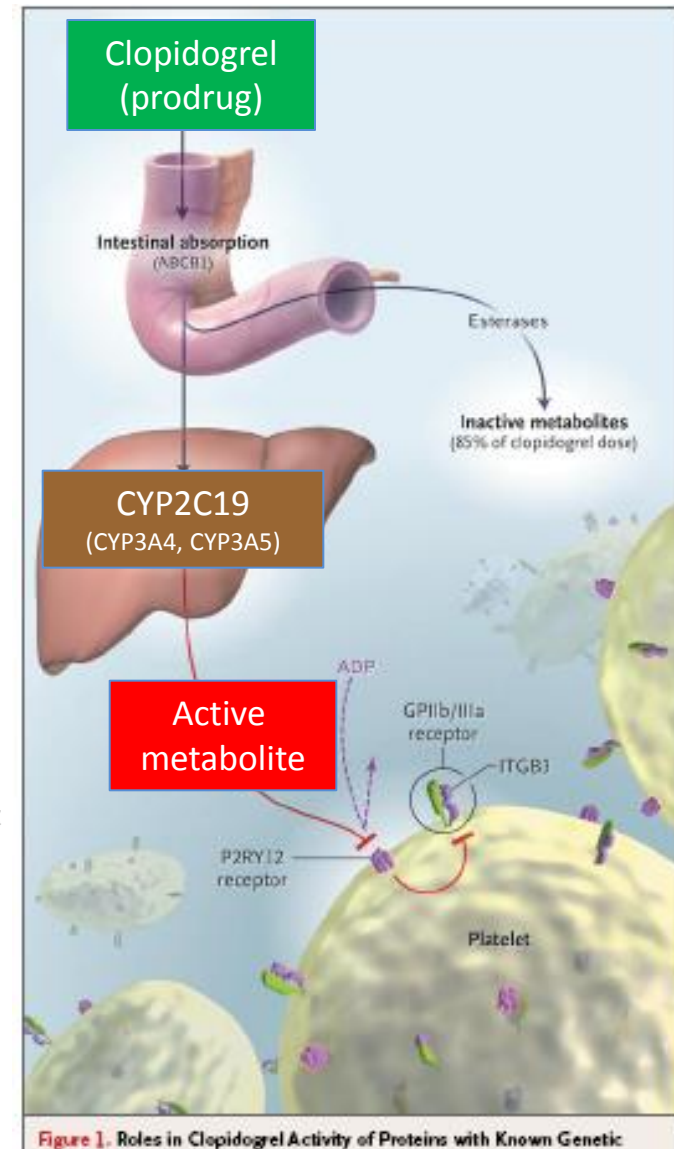


Figure 1. Roles in Clopidogrel Activity of Proteins with Known Genetic

Clopidogrel (Plavix): needs activation by **CYP2C19**

Caucasians: 3% PMs, 26% IMs; Asian: 30% PM, 50% IM

Test for CYP2C19 variants:

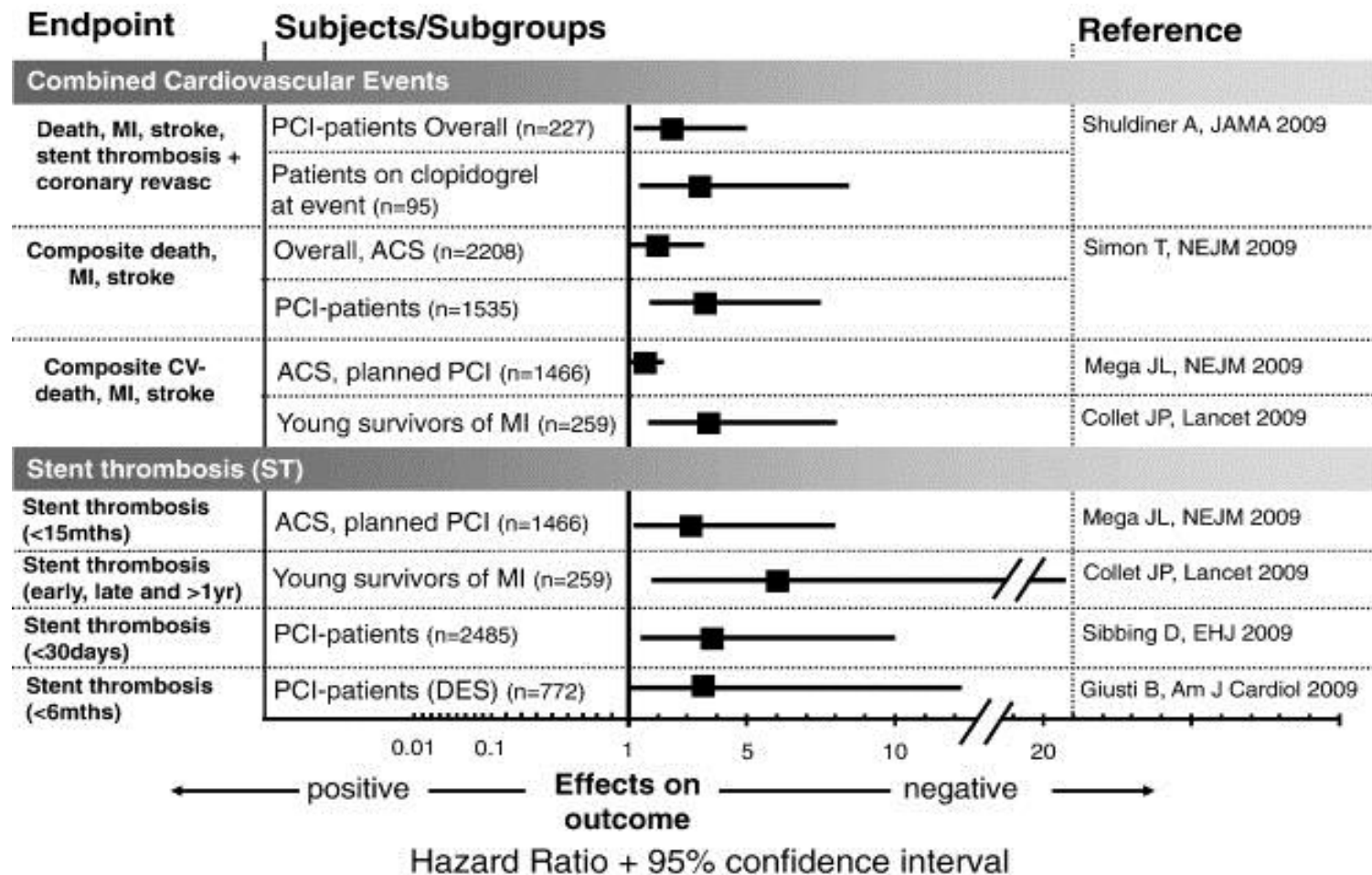
Negative →

clopidogrel (€)

Positive →

prasugrel/ticagrelor (€€€)

CYP2C19*2 carriers are at risk



FDA Boxed Warning on Clopidogrel

Warning: Diminished Effectiveness in Poor Metabolizers

- Effectiveness of clopidogrel depends on activation of active metabolite by the cytochrome P450 (CYP) system, specifically CYP2C19
- Poor metabolizers treated with clopidogrel at a standard dose exhibit higher cardiovascular event rates following acute coronary syndrome (ACS) or percutaneous coronary intervention in patients with normal CYP2C19 function
- Tests are available to identify a patient's CYP2C19 genotype and can be used as an aid in determining therapeutic approach
- Consider alternative treatment or treatment modification for patients identified as CYP2C19 poor metabolizers

action.¹⁷⁵ At present, genetic testing cannot be recommended in routine clinical practice due to insufficient prospective data. In con-



European Heart Journal
doi:10.1093/eurheartj/ehv320

August 2015

ESC GUIDELINES

2015 ESC Guidelines for the management of acute coronary syndromes in patients presenting without persistent ST-segment elevation – Web Addenda

Task Force for the Management of Acute Coronary Syndromes in Patients Presenting without Persistent ST-Segment Elevation of the European Society of Cardiology (ESC)

Authors/Task Force Members: Marco Roffi* (Chair), Carlo Patrono* (co-Chairperson), Christian Mueller

Document Reviewers: Helmut Baumgartner (CPG Review Coordinator) (Germany), Oliver Giermerli (CPG Review Coordinator) (Switzerland), Stefan Agewall (Norway), Lisa Badimon (Spain), Colin Baigent (UK), Hector Bueno (Spain), Raffaele Bugiardini (Italy), Scipione Careri (Italy), Filip Caselmann (Belgium), Thomas Cuisset (France), Cetin Erol (Turkey), Donna Fitzsimons (UK), Martin Halle (Germany), Christian Hamm (Germany), David Hildick-Smith (UK), Kurt Huber (Austria), Efstathios Iliodromidis (Greece), Stefan James (Sweden), Basil S. Lewis (Israel), Gregory Y. H. Lip (UK), Massimo F. Piepoli (Italy), Dimitrios



“Here is my genotype...”

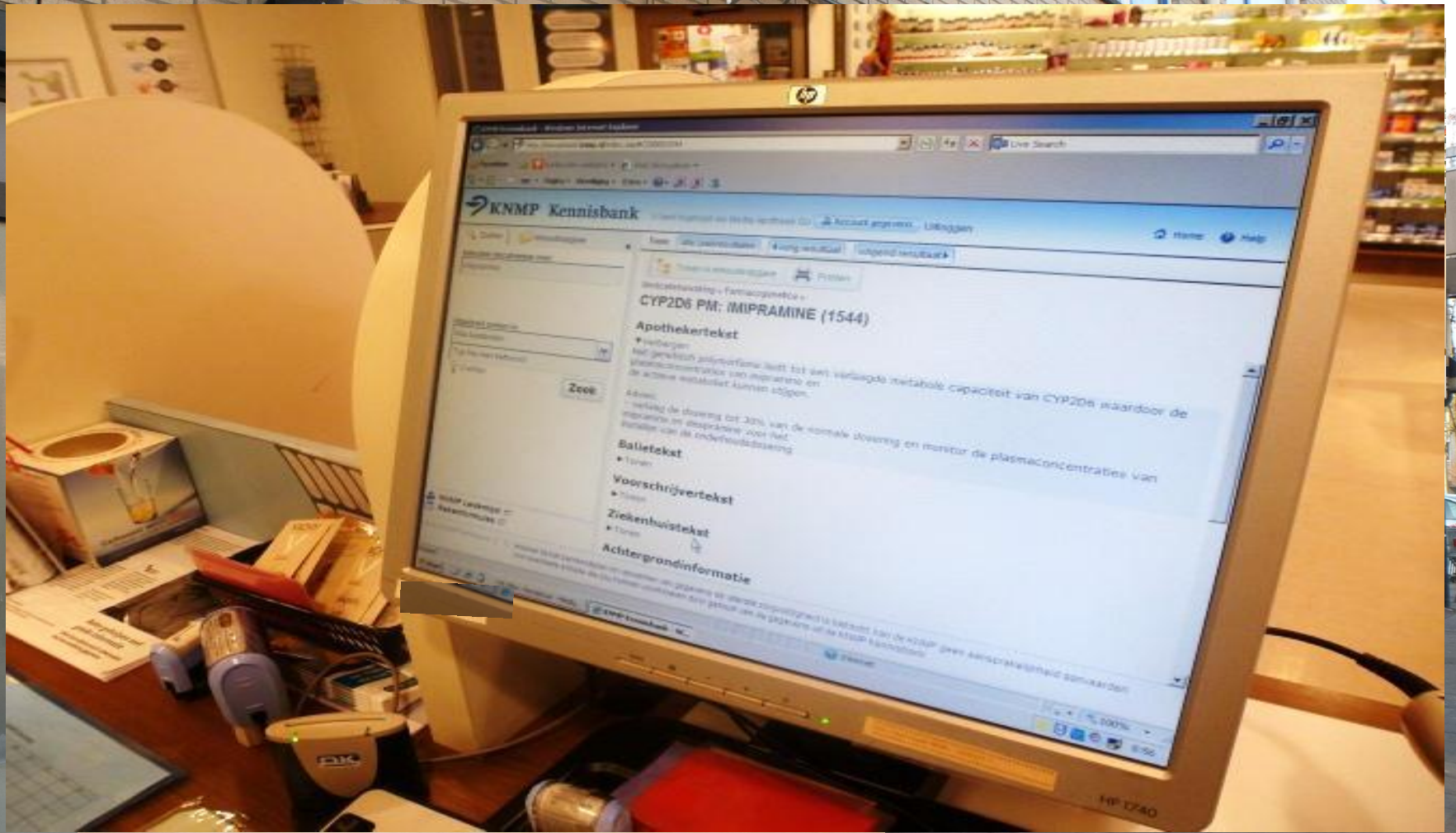
(The New Yorker (2000))

Dutch Pharmacogenetics
Working Group (DPWG):

Evidence based guidelines for
dosing on genotype for > 80 drugs

www.pharmgkb.org

Daily practice...



Farmacogenetica Profiel



Contact: farmacogenetica@erasmusmc.nl
 Telefoon: 010-7033119
www.erasmusmc.nl/farmacogenetica
www.farmacogenetica.nl













Naam: Test Erasmus MC/RvS

Geb. datum: 01/01/1980

BSN: 12345678

Uitgifte kaart: 14/07/2014

! geneesmiddelen mogelijk een uitsluitend ter beoordeling van or uw apotheker te vinden in de

Gen:	Uitslag:	Metabolisme	Prev.: ¹	Getest op:
 CYP1A2	*1/*1	Normaal	45%	*1C, *1F, *1K
 CYP2B6	*4/*6	Intermediair	25%	*4, 5, 6, 7, 8, 9, 13, 16, 18
 CYP2C9	*1/*2	Intermediair	17%	*2, 3
 CYP2C19	*1/*1	Normaal	80%	*2, 3, 17
 CYP2D6	*1/*2xN	Ultrasnel	3%	25 varianten (AmpliChip)
 CYP3A4	*1/*1	Normaal	80%	*1B,1G,3-6,10,12,17,18,20,22
 CYP3A5	*3/*3	Nonexpressor	80%	*3, *6
 BChE	U/S	Normaal	99%	A, K, F1, F2, H, J, Sc, Silent
 DPYD	*1/*2A	Intermediair	2%	*2A
 HLA-B*5701	NEG	Normaal	96%	
 TPMT	*1/*1	Normaal	89%	*2, 3A, 3B, 3C
 VKORC1	AA	Gevoelig	20%	-1639G>A

¹ In blanke bevolking. Kan afwijken bij andere ethniciteiten



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Pharmacogenomics
and Personalised
Medicine: research
progress and clinical
implementation

www.2017ESPTcongress.eu

ESPT 2017

FOURTH
CONFERENCE

OCTOBER 4-5-6-7

CATANIA, ITALY, Monastero dei Benedettini

