## Risk assessment and health promotion of the healthy patient in primary care

## PART 1

Giuseppe Parisi

## Evidence and recommendation

## Classes of recommendations

| Level of evidence |  | Classes of recommendations | Deffiliton | Suggested wording to use |
| :---: | :---: | :---: | :---: | :---: |
| Level of evidence A | Data derived from multiple randomized clinical trials or meta-analyses. | Class 1 | Evidence and/or general agreement that a given treatment or procedure is beneficial, useful, effective. | Is recommended/is indicated |
| Level of evidence B | Data derived from a single randomized clinical trial or large non-randomized studies. | Class II | Conflicting evidence and/or a divergence of opinion about the usefulness/efficacy of the given treatment or procedure. |  |
| Level of evidence C | Consensus of opinion of the experts and/ | Class Ila | Weight of evidence/opinion is in favour of usefulness/efficacy. | Should be considered |
|  | or small studies, retrospective studies, registries. | Class IIb | Usefulness/efficacy is less well established by evidence/opinion. | May be considered |
|  |  | Class III | Evidence or general agreement that the given treatment or procedure is not usefulleffective, and in some cases may be harmful. | Is not recommended |

## CV risk factors

## - Genetic/epigenetic <br> - Family history <br> Non modifiable <br> - Age <br> Risk factors <br> - Sex <br> - Smoking <br> - Cholesterol LDL/Lipid levels <br> - BP <br> Modifiable <br> Risk factors <br> - BMI <br> - DM <br> - Psycho-social factors

## CV risk estimation systems

Table 2 Current cardiovascular disease risk estimation systems for use in apparently healthy persons, updated from ${ }^{59,60}$

|  | Framingham" | SCORE ${ }^{30}$ | ASSIGN - SCORE ${ }^{\text {s/ }}$ | QRISKI ${ }^{\text {¢ }}$ \& QRISK2 ${ }^{\text {" }}$ | PROCAM ${ }^{\text {19 }}$ | Pooled Cohort Studies Equations ${ }^{50}$ | CUORE* | Globorisk ${ }^{\text {52 }}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Data | Prospective studies: Framingham Heart Study and Framingham offspring study. Latest version includes both | 12 pooled prospective studies | SHHEC Prospective study | QRESEARCH database | Prospective study | 4 Pooled prospective studies <br> ARIC <br> CHS <br> CARDIA <br> Framingham (original and offspring studies) | CUORE | Derivation cohort: 8 pooled prospective studies - Atherosclerosis Risk in Communities, Cardiovascular Health Study, Framingham Heart Study original cohort and offspring cohort, Honolulu Program, Multiple Risk Factor Intervention Trial, Puerto Rico Heart Health Program, and Women's Health Initative Clinical Trial |
| Population | General population, Framingham, <br> Massachusetts, USA. <br> Baselines: 1968-1971, <br> 1971-1975, 1984-1987 | 12 prospective studies from II European countries. Baselines: 1972-199\| | Random sample from general population in Scotland, baseline: 1984-1987 | Data collected from 1993-2008 from GP databases - imputation of missing data | Healthy employees. Baseline: <br> 1978-1995 | Baselines 1987-89 (ARIC), 1990 and 1992-3 (CHS), 1985-6 (CARDIA), 1968-1971, 1971-1975, 1984-1987 (Framingham) | 1980s and 1990s | 8 prospective studies from North America. <br> Baselines: 1948-1993 |
| Sample size | 3969 men and 4522 women | 117098 men and 88080 women | 6540 men and 6757 women | 1.28 million (QRISKI) <br> 2.29 million (QRISK2) | 18460 men and 8515 women | II 240 white women, 9098 white men, 264 African-American women and 1647 African-American men | $\begin{aligned} & 7520 \text { men and } 13127 \\ & \text { women } \end{aligned}$ | 33323 men and 16806 women |
| Calculates | 10-year risk of CAD eventsoriginally. Latest version: 10 -year risk of CVD events NCEP ATP III version: 10-year risk of hard coronary events | 10-year risk of CVD mortality | 10-year risk of CVD events | 10-year risk of CVD events. <br> Lifetime risk | Two separate scores calculate 10 -year risks of major coronary events and cerebral ischaemic events | 10-year risk for a first atherosclerotic CVD event. <br> Lifetime risk | 10-year probability of developing a first major CV event (myocardial infarction or stroke) | 10 year risk of fatal cardiovascular disease |
| Age range (years) | 30-75 | 40-65 | 30-74 | 35-74 | 20-75 | 20-79 | 35-69 | 40-84 |



Figure 2 SCORE chart: 10 -year risk of fetal cardiovascular dsease in populations of countries at low cardiowscuar risk based on the following risk factors: age, sex smokng sysolic blood pressure, total cholesterol CVD $=$ cardiovascular disesse: SCORE $=$ Systematic Coronay Risk Estimation

## Risk categories

| Very high-risk | Subjects with any of the following: <br> - Documented CVD, clinical or unequivocal on imaging. Documented clinical CVD includes previous $\mathrm{AMI}, \mathrm{ACS}$, coronary revascularization and other arterial revascularization procedures, stroke and TIA, aortic aneurysm and PAD. Unequivocally documented CVD on imaging includes significant plaque on coronary angiography or carotid ultrasound. It does NOT include some increase in continuous imaging parameters such as intima-media thickness of the carotid artery. <br> - DM with target organ damage such as proteinuria or with a major risk factor such as smoking or marked hypercholesterolaemia or marked hypertension. <br> - Severe CKD (GFR <30 mL/min/I. 73 m 2 ). <br> - A calculated SCORE $\geq 10 \%$. |
| :---: | :---: |
| High-risk | Subjects with: <br> - Markedly elevated single risk factors, in particular cholesterol $>8 \mathrm{mmol} / \mathrm{L}(>310 \mathrm{mg} / \mathrm{dL})$ (e.g. in familial hypercholesterolaemia) or $B P \geq 180 / 110 \mathrm{mmHg}$. <br> - Most other people with DM (with the exception of young people with type I DM and without major risk factors that may be at low or moderate risk). <br> - Moderate CKD (GFR $30-59 \mathrm{~mL} / \mathrm{min} / \mathrm{I} .73 \mathrm{~m}^{2}$ ). <br> - A calculated SCORE $\geq 5 \%$ and $<10 \%$. |
| Moderate risk | SCORE is $\geq 1 \%$ and $<5 \%$ at 10 years. Many middleaged subjects belong to this category. |
| Low-risk | SCORE <1\%. |

ACS $=$ acute coronary syndrome; $\mathrm{AMI}=$ acute myocardial infarction; $\mathrm{BP}=$ blood pressure; $\mathrm{CKD}=$ chronic kidney disease; $\mathrm{DM}=$ diabetes mellitus; $\mathrm{GFR}=$ glomerular filtration rate; PAD $=$ peripheral artery disease; $\mathrm{SCORE}=$ systematic

## High and low risk populations





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## The chart we use: low risk

## - Low risk

- Andorra, Austria, Belgium, Cyprus, Denmark, Finland, France, Germany, Greece, Iceland, Irelar Israel, Italy, Luxembourg, Malta, Monaco, The Neth- erlands, Norway, Portugal, San Marino, Slovenia, Spain, Sweden, Switzerland and the United Kingdom.
- High risk
- Bosnia and Herzegovina, Croatia, Czech Re- publ Estonia, Hungary, Lithuania, Montenegro, Morocco, Poland, Romania, Serbia, Slovakia Tunisia and Turkey.
- Albania, Algeria, Armenia, Azerbaijan, Belarus, Bulgaria, Egypt, Georgia, Kazakhstan, Kyrgyzstan Latvia, former Yugoslav Republic of Macedonia, Moldova, Russian Fed- eration, Syrian Arab Republic, Tajikistan, Turkmenistan, Ukraine and Uzbekistan.



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