Analysing falls in coronary heart disease mortality in the West Bank between 1998 and 2009

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ABSTRACT
Objectives: To analyse coronary heart disease (CHD) mortality and risk factor trends in the West Bank, occupied Palestinian territory between 1998 and 2009.

Design: Modelling study using CHD IMPACT model.

Setting: The West Bank, occupied Palestinian territory.

Participants: Data on populations, mortality, patient groups and numbers, treatments and cardiovascular risk factor trends were obtained from national and local surveys, routine national and WHO statistics, and critically appraised. Data were then integrated and analysed using a previously validated CHD model.

Primary and secondary outcome measures: CHD deaths prevented or postponed are the main outcome.

Results: CHD death rates fell by 20% in the West Bank, between 1998 and 2009. Smoking prevalence was initially high in men, 51%, but decreased to 42%. Population blood pressure levels and total cholesterol levels also decreased. Conversely, body mass index rose by 1–2 kg/m² and diabetes increased by 2–8%. Population modelling suggested that more than two-thirds of the mortality fall was attributable to decreases in major risk factors, mainly total cholesterol, blood pressure and smoking. Approximately one-third of the CHD mortality decreases were attributable to treatments, particularly for secondary prevention and heart failure. However, the contributions from statins, surgery and angioplasty were consistently small.

Conclusions: CHD mortality fell by 20% between 1998 and 2009 in the West Bank. More than two-thirds of this fall was due to decreases in major risk factors, particularly total cholesterol and blood pressure. Our results clearly indicate that risk factor reductions in the general population compared save substantially more lives to specific treatments for individual patients. This emphasizes the importance of population-wide primary prevention strategies.

INTRODUCTION
The occupied Palestinian territory (oPt) comprises the Gaza Strip and the West Bank including East Jerusalem. Some 46% of the Palestinian population of 3.8 million are younger than 15 years while only 3% are older than 65 years. However, the number of older people is increasing gradually and the population is slowly ageing. The Palestinians are also undergoing rapid epidemiological transition. Communicable diseases of childhood have already been controlled with effective immunisation programmes, and polio eradication has been achieved. However, non-communicable diseases have now overtaken communicable diseases as the main causes of mortality.1 Thus, cardiovascular disease (CVD) and cancer are now the major causes of morbidity and mortality in the oPt.2 The Ministry of Health (MOH) recently reported that 25% of all deaths are due to cardiovascular diseases in 2010, followed by cerebrovascular diseases (12%), cancer (11%) and diabetes (6%).3 Increasing levels of adverse risk factors such as diabetes, obesity and physical inactivity have been repeatedly documented.1,3

Chronic disease death rates are actually decreasing in the developed world (Western Europe, North America, Australia and New Zealand). However, mortality is increasing in the developing countries. It is predicted that by 2020, CVD deaths will exceed infectious and parasitic disease deaths in all regions except for sub-Saharan Africa.6 Furthermore, the Eastern Mediterranean region has been recognised as a hot spot for diabetes and CVD, yet local data to inform policy is severely limited.

KEY POINTS
- CHD mortality fell by 20% between 1998 and 2009 in the West Bank.
- More than two-third of this fall was due to decreases in major risk factors, particularly total cholesterol, and blood pressure.
The IMPACT coronary heart disease (CHD) model was developed to quantify recent trends in CHD mortality, in order to help maximise the effective use of existing information and resources to develop appropriate policies and strategies. This study aims to adopt the IMPACT CHD model to the Palestinian context, namely the West Bank population, in order to help explain recent changes in CHD mortality.

METHODS
A validated version of the IMPACT CHD mortality model was further modified and updated to suit the countries in the Middle East and specifically the oPt. The IMPACT model was previously validated in many developed countries7–10 and in one middle-income country (China).11

Palestinian data on risk factors levels and current uptake levels of evidence-based treatments were identified by extensive searches for published or unpublished data and complemented with specifically designed surveys. All data sources were critically appraised by the local research team and the results are presented in the appendix. The data needed for the analysis were available for men and women aged 25–75 years in the West Bank, occupied Palestinian territory for the period 1998–2009 with some age limitations as described in the appendix.

The specific data items used to populate the model included: (1) Patient numbers in specific CHD groups (myocardial infarction (MI), congestive heart failure and chronic angina pectoris), (2) uptake of specific medical and surgical treatments, (3) population trends in major cardiovascular risk factors (smoking, total cholesterol, systolic blood pressure, body mass index (BMI), diabetes and physical inactivity).

The main output of the model is the number of deaths prevented or postponed attributed to the changes in specific treatments and/or risk factor levels.

Identification and assessment of relevant data
Information on the West Bank population demographic changes was obtained and validated for the first year from 1997 census-based projections and 2007 census-based projections for the final year by the Palestinian Central Bureau of Statistics (PCBS).12 Numbers of deaths for both years were obtained from the Health Information Management Centre—Palestinian Ministry of Health (MOH). Population risk factors trend data for the year 1998 were obtained from two epidemiological studies conducted in the rural and urban areas of Ramallah governorate in the West Bank. These were the only available published epidemiological studies in the West Bank for that period. They covered a rural and an urban site that were prototypic of many West Bank villages and urban sites.13

CHD numbers of hospital admissions in addition to treatment uptake were obtained from our treatment uptake survey conducted in 2009 which included four hospitals in the north, centre and south of the West Bank. The number of patients undergoing coronary artery bypass grafting and angioplasty were obtained from records in the two hospitals providing this service in the West Bank. The prevalence of angina, heart attack survivors and congestive heart failure in the community were each estimated on the basis of national health surveys and treatment uptake surveys. Information on treatment uptake in the community was also checked by eliciting expert opinion from practising clinicians.

The efficacy of therapeutic interventions was based on recent meta-analyses and randomised controlled trials. The Mant and Hicks14 approach was used to correct for polypharmacy.

Change in CHD deaths
First, the number of CHD deaths expected in 2009 was calculated by indirect age standardisation based on the assumption that 1998 death rates had persisted unchanged until 2009. The number of CHD deaths actually observed in 2009 was then subtracted. The difference between the two represents the fall in CHD deaths (the number of deaths prevented or postponed) that the model needed to explain.

Mortality changes attributed to risk factor trends
The number of deaths prevented or postponed from changes in risk factors was estimated using two approaches. The regression β coefficients approach was used to quantify the population mortality impact of change in those specific risk factors, measured as continuous variables (blood pressure, total cholesterol and BMI). The second approach, population-attributable risk fraction, was employed for categorical variables—diabetes, physical inactivity and smoking:

\[
PAR = \frac{\text{Prevalence} \times (\text{Relative risk} - 1)}{(\text{Prevalence} \times (\text{Relative risk} - 1)) + 1}
\]

PAR calculation was stratified by age and sex. Details of model methodology have been published previously7 and worked examples are shown in the appendix.

Estimating the contribution of medical and surgical treatments
The model aimed to include all medical and surgical treatments in 1998 (the base year) and 2009 (the final year). Treatment uptake data were not available for the year 1998 and thus the data included in the model for this year was estimated after consultation with cardiologists and experts working in hospital and community at that time.

The mortality reduction for each treatment for the number of patients in each group, stratified by age and sex, was calculated as the age-specific case death in that group multiplied by the relative mortality reduction reported in published meta-analyses multiplied by the treatment uptake (the proportion of patients receiving that specific treatment, appendix).
Case-death data were obtained from large, unselected, population-based patient cohorts. The survival benefit over a 1-year time interval was used for all treatments.

**Treatment overlaps**

Potential overlaps between different groups of patients were identified and appropriate adjustments were made. Patients group calculations and assumptions are detailed in the appendix.

**Treatment adherence**

Adherence (defined as the proportion of treated patients actually taking therapeutically effective levels of the prescribed medication) was assumed to be 100% among hospital patients, 70% among all symptomatic community patients and 50% among asymptomatic community patients, based on the literature.15

**Sensitivity analyses**

Because of the uncertainties surrounding some of the values, multiway sensitivity analyses using the Brigg’s analysis of extremes method was used.16

**Model validation: comparison with observed mortality falls**

The model estimate for the changes in deaths attributed to all treatments plus all risk factor changes was summed for men and women in each specific age group. The model fit was then compared with the observed change in mortality for that group.

**RESULTS**

Between 1998 and 2009 CHD mortality in the West Bank fell by 20%. This resulted in approximately 125 fewer CHD deaths compared with the expected number of CHD deaths in 2009 if 1998 death rate had persisted unchanged (table 1).

The reduction in CHD mortality among women (35%) was twice as large as the 19% reduction observed among men. The CHD mortality reduction was seen in all age groups with particularly large reductions being observed among those aged 45–54 years, men (24%) and women (49%).

**Major CHD risk factors**

Changes in cardiovascular risk factors included in the model were together estimated to prevent or postpone approximately 80 deaths in 2009 (minimum estimate 75 and maximum estimate 140), which represented approximately 66% of the total CHD mortality fall (figure 1). Table 2 presents changes in the selected risk factors and the attributed deaths. Changes in risk factors were complex: reduction in total cholesterol (mean reduction 0.34mmol/l in men and 0.22 mmol/l in women), blood pressure (5.27 mm Hg in men and 0.01 mm Hg in women) and in smoking prevalence (11.5% in men and 2.2% in women). These changes together prevented or postponed approximately 125 deaths (table 2). A total fall of 40% in CHD deaths were thus attributable to cholesterol reduction (minimum 38% and maximum 77%), and 36% to blood pressure reduction (minimum 30% and maximum 50%) and 33% to smoking reduction (minimum 22% and maximum 49%). However, an additional 45 deaths were attributable to adverse trends (figure 2). Mainly an increase in diabetes prevalence (8.5% in men and 2% in women) generated approximately 30 additional deaths. The mortality effects of increases in physical inactivity and mean BMI were relatively modest (table 2).

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**Table 1** Population sizes and coronary heart disease death rates in the West Bank, 1998 and 2009

| Age groups | 25–34 | 35–44 | 45–54 | 55–64 | 65–74 | Total |
|------------|-------|-------|-------|-------|-------|-------|
| Men        |       |       |       |       |       |       |
| Population in 1998 | 132084 | 80434 | 41727 | 28653 | 19524 | 302422 |
| Population in 2009 | 177873 | 132945 | 86399 | 41984 | 21619 | 460820 |
| Deaths in 1998 (number) | 9 | 21 | 57 | 101 | 130 | 318 |
| Deaths in 2009 (number) | 11 | 30 | 89 | 131 | 133 | 394 |
| Death rates per 100000 in 1998 | 6.4 | 26.1 | 134.5 | 350.7 | 665.8 | 105.2 |
| Death rates per 100000 in 2009 | 6.2 | 22.8 | 103.4 | 311.2 | 616.7 | 85.5 |
| Percentage of change (crude) | 0 | −12.6 | −23.6 | −11.3 | −7.4 | −19 |

| Women |       |       |       |       |       |       |
|-------|-------|-------|-------|-------|-------|-------|
| Population in 1998 | 125493 | 76747 | 45711 | 36038 | 24445 | 308434 |
| Population in 2009 | 170942 | 127590 | 80883 | 43693 | 28424 | 451532 |
| Deaths in 1998 (number) | 5 | 9 | 22 | 59 | 107 | 202 |
| Deaths in 2009 (number) | 5 | 11 | 20 | 51 | 105 | 192 |
| Death rates per 100000 in 1998 | 3.6 | 11.1 | 48.1 | 162.3 | 437.7 | 65.5 |
| Death rates per 100000 in 2009 | 3.1 | 8.9 | 24.7 | 116.0 | 370.6 | 42.5 |
| Percentage of change (crude) | 0 | −19.8 | −48.6 | −28.6 | −15.3 | −35 |

Abu-Rmeileh NME, Shoaibi A, O’Flaherty M, et al. BMJ Open 2012;2:e001061. doi:10.1136/bmjopen-2012-001061
Medical and surgical treatments

Medical and surgical treatments together prevented or postponed approximately 35 deaths in 2009 (minimum estimate 20 and maximum estimate 108; figure 3). The total treatments thus accounted for approximately 29% (minimum 17% and maximum 82%) of the total CHD mortality reduction (table 3). Secondary prevention following acute MI explained over 7% (minimum 3% and maximum 25%) of deaths prevented or postponed (with ACE inhibitors, aspirin and β-blockers and β-blockers being the main contributors in this group; table 3). Smaller contributions came from chronic angina treatment (6%, minimum 5% and maximum 11%) and heart failure treatment in the community (5%, minimum 3% and maximum 10%) and hypertensive treatment in the community (4%, minimum 1% and maximum 19%). Secondary prevention postrevascularisation had a very modest effect on CHD mortality reduction with a contribution of 2% (minimum 1% and maximum 5%; table 3).

Validation and model fit

In summary, when including all risk factor and treatment data, the model explained approximately 95% of the total CHD mortality reduction observed in the West Bank population between 1998 and 2009. The remaining 5% was unexplained and might reflect other, unmeasured factors. The model estimates of deaths were reasonably consistent with the observed deaths for almost all age groups. Overall, the model fit was better for men than for women. However, in the age group 45–54 the model overestimated deaths in men and women (table 4). Moreover, irrespective of whether best minimum or maximum estimates were used, the relative contributions remained relatively consistent.

DISCUSSION

CHDs mortality decreased by 20% in the West Bank, oPt between 1998 and 2009. Almost two-thirds of this reduction was attributable to changes in major risk factors and approximately one-third was explained by medical and surgical treatments. The recent mortality trends observed in the West Bank were thus characterised by reductions in CHD mortality similar to those observed in the developed rather than developing countries. Other studies have documented similar reductions in CHD mortality in Europe, North America and New Zealand especially since the 1980s.7 8 17 18

The biggest CHD mortality reductions attributable to changes in major risk factors came from declines in total cholesterol and smoking, and also blood pressure in men. Interestingly, the implementation efforts of an antismoking law in 2005 might have contributed to the substantial declines in smoking prevalence.19 20

However, the increasing Westernisation of diet, particularly junk food and soda represent an ominous future threat. Furthermore, diabetes, obesity and physical inactivity increased substantially between 1998 and 2009.21–23 In the year 1998, the prevalence of obesity was 49% for women and 30% for men aged 35–64 years old.24 Since then, the prevalence of obesity has increased mainly among men.25 26 The increase in diabetes prevalence generated approximately 30 additional CHD deaths. Rising diabetes and obesity, especially among men, represents a public health priority.
| Risk factors | Risk factor levels | Absolute change in risk factor 1998–2009 | Relative risk (or β coefficient) | Deaths prevented or postponed | Proportion of overall deaths (%) |
|--------------|--------------------|----------------------------------------|-------------------------------|-------------------------------|---------------------------------|
| Cholesterol (total) (mmol/l) | | | | | |
| Cholesterol (men) | 5.37 | 5.03 | 0.34 | β = 0.65 | 49 | 46 | 94 | −40.3 |
| Cholesterol (women) | 5.25 | 5.03 | 0.22 | β = 0.65 | 40 | 26 | 59 | −32.5 |
| Smoking (total) (%) | | | | | |
| (% men smoking) | 51.5 | 40 | 11.5 | | 5 | −2 | −5 | −2 |
| (% women smoking) | 5.5 | 3.3 | 2.2 | | | | | |
| Body mass index (BMI) (total) (kg/m²) | | | | | |
| BMI (men) | 26.95 | 30.88 | −3.93 | β = 0.02 | −32 | −27 | −60 | 26.2 |
| BMI (women) | 29.12 | 30.99 | −1.87 | β = 0.02 | | | | |
| Diabetes (total) % | | | | | |
| Percentage of diabetes (men) | 9.1 | 17.6 | −8.5 | | 2 | | | |
| Percentage of diabetes (women) | 12.4 | 14.4 | −2 | | 2 | | | |
| Population systolic blood pressure (BP) (total) mm Hg | | | | | |
| Population BP (men) | 125.57 | 120.3 | 5.27 | β = 0.053 | 43 | 39 | 84 | −35.6 |
| Population BP (women) | 118.44 | 118.45 | −0.01 | β = 0.053 | 38 | 37 | 61 | −31.5 |
| Population BP after adjustment for hypertension treatments | | | | | |
| Physical inactivity | 79.3 | 91.3 | 12 | | −12 | −17 | 9.7 | |
| Percentage of physical inactivity men | 87.9 | 93 | 5.1 | | | | | |
| Estimated total risk factor effects | | | | | | 81 | 73 | 138 | −66.5 |
Effective evidence-based interventions exist and should be considered, notably junk-food taxes, labelling and reformulation issues and advertising bans.27 28

The recently developed Palestinian non-communicable diseases (NCDs) strategy adopted an integrated approach encompassing promotion, prevention and control programmes. Population-based multisectoral effective evidence-based interventions were clearly indicated to control diabetes and promote physical activity such as health promotion, fiscal measures, market control and community participation. Yet a clear vision and scope of implementation are still evolving.29

Modern medical treatments accounted for approximately 30% of the CHD mortality reduction. The scale of effect is similar to that reported in studies using the same methodology in Iceland, Sweden and Finland but lower than the 45–50% reported for North America30 and Europe.8 10 The biggest contributions came from aspirin and community-based medications for secondary prevention, chronic angina and heart failure. Treatment uptakes were generally mediocre and need to be improved. However, such treatments create challenges for health providers in terms of identifying patients, providing medications and ensuring their long-term compliance.

Data on patient groups and treatment uptake levels were scarce. Such data were not available at the national or the hospital level. The required information was therefore collected directly from the hospital patients’ records to be utilised in this study. Furthermore, the uptake levels were not consistent among different hospitals and sometimes between physicians. These two findings therefore highlight a lack of standardised healthcare provision and important gaps in the NCD health information system.

**Modelling strengths and limitations**

The modelling approach used in the study was comprehensive, it synthesised all the key risk factors and treatment options to help quantify changes in CHD mortality. Additionally, the model used rigorous sensitivity analyses to systematically examine the potential influence of uncertainties in the data (quality and sources) and model assumptions, and hence quantify the potential maximum and minimum effects of these contributory factors.

This modelling approach also has obvious limitations. Notably, the extent and quantity of available data on CHD risk factor trends and treatment uptake. Furthermore, the model tended to overestimate the number of deaths averted in almost all age groups, especially among women. This may reflect less precision in female data. However, the data used in this model was generally of good quality. Mortality data were obtained from MOH death registry. Death registry data were evaluated in previous studies as medium quality based on the WHO criteria.31 The demographic information was obtained from census data, and the risk factor trends were obtained from well-designed epidemiological studies and national PCBS, MOH and WHO surveys. Treatment uptake and patient group data were obtained from an extensive hospital-based survey conducted in 2009. Very scarce data available on treatment uptake were also obtained. However, researchers had to conduct additional surveys to get the timely, accurate data required by the model. Certain assumptions were needed to fill in the gaps for missing information. For instance, assumptions were made for the small group aged 65–74 years where risk factor information were not available. Assumption on treatment uptake at the starting point was also based on expert opinions working in the system for more than 10 years. Estimates of treatment uptake were collated from international and

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**Figure 2** Coronary heart disease deaths prevented or postponed attributable to specific risk factor changes in Palestine 1998–2009: Sensitivity analysis (▪, best estimate; bars indicate minimum and maximum estimates).

**Figure 3** Coronary heart disease deaths prevented or postponed attributable to specific treatments in Palestine 1998–2009: Sensitivity analysis (▪, best estimate; bars indicate minimum and maximum estimates).
regional literature then validated with the expert opinions. The generalisation of efficiency estimates from meta-analysis of controlled clinical trials to effectiveness in clinical practice in the West Bank setting were clearly optimistic, and may have resulted in an overestimate of the true treatment benefits.

All these assumptions are transparent, being systematically detailed in the appendix, supported by local expert opinions and literature from the region and included in the sensitivity analyses. By good luck, the overall model fit approached 100%. However, it should be noted that fit within specific age groups was much less perfect.

Public health implications
CHD mortality fell by 20% between 1998 and 2009 in the West Bank, oPt. More than two-third of this fall was

Table 3  Deaths prevented or postponed by medical and surgical treatments in the West Bank in 2009

| Treatments                                      | Patients eligible | Treatment uptake (%) | Coronary heart disease deaths prevented or postponed | Proportion of overall deaths prevented or postponed (%) |
|-------------------------------------------------|-------------------|----------------------|------------------------------------------------------|-------------------------------------------------------|
| Acute myocardial infarction                     | 489               | 0.06                 | Best estimate Minimum estimate Maximum estimate      |                                                       |
| Hospital resuscitation                         |                   |                      | 2 1 6                                                | −1.9                                                  |
| Thrombolysis and aspirin                       |                   | 0.90                 | 0 0 2                                                | −0.3                                                  |
| Aspirin alone                                   |                   | 0.39                 | 3 1 6                                                | −2.4                                                  |
| Thrombolysis alone                             |                   | 0.58                 | 1 0 2                                                | −0.6                                                  |
| ACE inhibitors                                  |                   | 0.44                 | 1 0 2                                                | −1                                                   |
| Secondary preventionpostinfarction             | 2266              | 9 4 30               |                                                      |                                                       |
| Aspirin                                         |                   | 0.483                | 4 1 9                                                | −2.9                                                  |
| β-Blockers                                      |                   | 0.388                | 4 1 11                                               | −3.6                                                  |
| Aspirin and β-blockers                         |                   | 0.415                | 4 1 10                                               | −3.4                                                  |
| ACE inhibitors                                  |                   | 0.472                | 5 2 13                                               | −4.2                                                  |
| Statins                                         |                   | 0.046                | 0 0 1                                                | −0.4                                                  |
| Warfarin                                        |                   | 0.100                | 1 0 3                                                | −0.9                                                  |
| Rehabilitation                                  |                   | 0.483                | 1 0 3                                                | −0.9                                                  |
| Secondary preventionpostrevascularisation      | 208               | 2 0 6                |                                                      |                                                       |
| Angina                                           |                   |                      |                                                      |                                                       |
| Coronary Artery Bypass Grafting surgery (2004–2005) | 208              | 1                    | 6 3 10                                               | −4.6                                                  |
| Aspirin in the community                        | 22180             | 0.47                 | 15 13 40                                             | −12.7                                                 |
| Statin in the community                         | 22180             | 0.46                 | 20 11 50                                             | −16.8                                                 |
| Unstable angina                                 | 489               |                      | 1 2 9                                                | −1.1                                                  |
| Heart failure                                   | 2811              |                      |                                                      |                                                       |
| Hospital patients                               | 468               | −2                   | −1 −8                                                | −1.5                                                  |
| Community patients                              | 2342              | −7                   | −4 −4                                                | −5.4                                                  |
| Hypertension treatments                         |                   | −5                   | −2 −23                                               | −4.1                                                  |
| Total treatment effects                         | 35 20 108         |                      |                                                      |                                                       |

Table 4  Model validation: estimated versus observed changes in coronary heart disease (CHD) deaths between 1998 and 2009

| Age groups | 25–34 | 35–44 | 45–54 | 55–64 | 65–74 | Total |
|------------|-------|-------|-------|-------|-------|-------|
| Men        |       |       |       |       |       |       |
| Estimated fall in CHD deaths                    | 11 11 | 117 117| 147 147| 144 144| 454 454|       |
| Observed fall in CHD deaths                     | 11 11 | 89 89 | 131 131 | 133 133 | 394 394 |       |
| Discrepancy                                     | 0 −5 | −28 −28 | −16 −16 | −11 −11 | −60 −60 |       |
| Estimated fall/observed fall in CHD deaths (%)  | 100 100 | 117 117 | 131 131 | 112 112 | 108 108 | 115 115 |
| Women                                              |       |       |       |       |       |       |
| Estimated fall in CHD deaths                    | 6 6 | 39 39 | 71 71 | 124 124 | 254 254 |       |
| Observed fall in CHD deaths                     | 5 5 | 20 20 | 51 51 | 105 105 | 192 192 |       |
| Discrepancy                                     | −1 −1 | −19 −19 | −20 −20 | −19 −19 | −62 −62 |       |
| Estimated fall/observed fall in CHD deaths (%)  | 120 120 | 127 127 | 195 195 | 139 139 | 118 118 | 132 132 |
due to changes in major risk factors mainly total cholesterol, blood pressure and smoking.

Our results clearly indicate that risk factor improvements in the general population saved substantially more lives than specific treatments for individual patients. This emphasizes the importance of population-wide primary prevention strategies. Such strategies should also emphasize the risk factors which had a negative effect on the reduction of CHD in the model especially, diabetes, BMI and increased levels of physical inactivity. The Palestinian policy and strategic plan for NCD had focused on health diet and physical activity as its second and third policy objectives (MOH).52

Acknowledgements The authors would like to thank the Palestinian Ministry of Health, specifically the Palestinian Health Information Centre and Primary Health Care Staff for providing the researchers with the Stepwise Survey 2010 data.

Contributors All authors have fulfilled the following contributions (1) substantial contributions to conception and design, acquisition of data, or analysis and interpretation of data; (2) drafting the article or revising it critically for important intellectual content and (3) final approval of the version to be published.

Funding FP7-HEALTH-2007-B. Project number 223075.

Competing interests None.

Provenance and peer review Not commissioned; externally peer reviewed.

Data sharing statement No additional data are available.

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