Drug eluting and bare metal stents in people with and without diabetes: collaborative network meta-analysis

Christoph Stettler, senior research fellow,1,2,3 Sabin Allemann, research fellow,1,2 Simon Wandel, research fellow,1 Adnan Kastrati, professor of cardiology,4 Marie Claude Morice, professor of cardiology,5 Albert Schömig, professor of medicine,4 Matthias E Pfisterer, professor of cardiology,6 Gregg W Stone, professor of medicine,7 Martin B Leon, professor of medicine,7 José Suárez de Lezo, professor of cardiology,8 Jean-Jacques Goy, professor of interventional cardiology,9 Seung-Jung Park, professor of cardiology,10 Manel Sabaté, associate professor of cardiology,11 Maarten J Suttorp, head of department,12 Henning Kelbaek, associate professor of cardiology,13 Christian Spaulding, professor of cardiology,14 Maurizio Menichelli, interventional cardiologist,15 Paul Vermeersch, interventional cardiologist,16 Maurits T Dirksen, training fellow in cardiology,17 Pavel Cervinka, cardiologist,18 Marco De Carlo, vice director,19 Andreas Erglis, associate professor of cardiology,20 Tania Chechi, interventional cardiologist,21 Paolo Ortolani, interventional cardiologist,22 Martin J Schalij, professor of cardiology,23 Peter Diem, head of division,2 Bernhard Meier, professor of cardiology,24 Stephan Windecker, head of invasive cardiology,24,25 Peter Juni, head of division 24,25

ABSTRACT
Objective To compare the effectiveness and safety of three types of stents (sirolimus eluting, paclitaxel eluting, and bare metal) in people with and without diabetes mellitus. Design Collaborative network meta-analysis. Data sources Electronic databases (Medline, Embase, the Cochrane Central Register of Controlled Trials), relevant websites, reference lists, conference abstracts, reviews, book chapters, and proceedings of advisory panels for the US Food and Drug Administration. Manufacturers and trialists provided additional data. Review methods Network meta-analysis with a mixed treatment comparison method to combine direct within trial comparisons between stents with indirect evidence from other trials while maintaining randomisation. Overall mortality was the primary safety end point, target lesion revascularisation the effectiveness end point. Results 35 trials in 3852 people with diabetes and 10 947 people without diabetes contributed to the analyses. Inconsistency of the network was substantial for overall mortality in people with diabetes and seemed to be related to the duration of dual antiplatelet therapy (P value for interaction 0.02). Restricting the analysis to trials with a duration of dual antiplatelet therapy of six months or more, inconsistency was reduced considerably and hazard ratios for overall mortality were near one for all comparisons in people with diabetes: sirolimus eluting stents compared with bare metal stents 0.88 (95% credibility interval 0.55 to 1.30), paclitaxel eluting stents compared with bare metal stents 0.91 (0.60 to 1.38), and sirolimus eluting stents compared with paclitaxel eluting stents 0.95 (0.63 to 1.43). In people without diabetes, hazard ratios were unaffected by the restriction. Both drug eluting stents were associated with a decrease in revascularisation rates compared with bare metal stents in people both with and without diabetes. Conclusion In trials that specified a duration of dual antiplatelet therapy of six months or more after stent implantation, drug eluting stents seemed safe and effective in people both with and without diabetes.

INTRODUCTION
People with diabetes experience a more generalised form of atherosclerosis than people without diabetes. They are at an increased risk for coronary heart disease and have more restenoses after the implantation of coronary stents. On average sirolimus eluting stents and paclitaxel eluting stents are associated with a noticeable reduction in target lesion revascularisation compared with bare metal stents, whereas the rates of overall mortality and cardiac mortality associated with the three stents are similar.1 Differences in the process and dynamics of restenosis along with variations in metabolic profiles may, however, alter safety or effectiveness profiles of the different stent types, particularly in people with diabetes.

Randomised trials have reported a reduced revascularisation rate with both sirolimus eluting stents and paclitaxel eluting stents compared with bare metal stents in people with diabetes,2-4 but the trials were hampered by small numbers of patients and a limited duration of follow-up. A meta-analysis of four early trials in 428 people with diabetes that compared sirolimus eluting stents with bare metal stents for up to four years suggested a strongly increased risk of mortality (hazard ratio 2.90, 95% confidence interval

1Institute of Social and Preventive Medicine, University of Bern, 3012 Bern, Switzerland
2Division of Endocrinology, Diabetes and Clinical Nutrition, University Hospital, Bern, Switzerland
3International Centre for Circulatory Health, National Heart and Lung Institute, Imperial College, London
4Deutsches Herzzentrum, Technische Universität, Munich, Germany
5Institut Hospitalier Jacques Cartier, Massy, France
6Division of Cardiology, University Hospital, Basel, Switzerland
7Columbia University Medical Center, New York, USA
8Servicio de Cardiología, Hospital Universitario Rona Sofia, Cordoba, Spain
9Service of Cardiology, Clinique Cecil, Lauzanne, Switzerland
10Department of Medicine, ASAN Medical Center, Seoul, Korea
11Department of Interventional Cardiology, Hospital de San Pau, Barcelona, Spain
12Department of Interventional Cardiology, St Antonius Hospital, Nieuwegein, Netherlands
13Cardiac Catheterisation Laboratory, Rigshospitalet, Copenhagen, Denmark
14Cochin Hospital, Assistance Publique Hôpitaux de Paris, Paris
1.38 to 6.10). In view of an average rate of overall mortality of 12% over four years observed in people with diabetes, this hazard ratio would translate into a number needed to harm to cause one death over four years as low as 4 (95% confidence interval 2 to 22). Another meta-analysis, published simultaneously, also included more recent trials with shorter durations of follow-up: pooling 14 trials in 1411 people with diabetes resulted in a hazard ratio of only 1.27, but the 95% confidence interval ranged from 0.83 to 1.95 and was compatible with both moderate benefit and substantial harm.

Network meta-analyses or mixed treatment comparisons allow a unified, coherent analysis of all randomised controlled trials that compared either of the two drug eluting stents with bare metal stents or the two drug eluting stents head to head, while fully respecting randomisation. In a previous network meta-analysis we determined the average benefits and harms of all three stent types and provided preliminary results for overall mortality and the composite of death or myocardial infarction stratified according to the presence or absence of diabetes. Here we extend the network meta-analysis stratified according to diabetes status to include 35 trials in 1799 patients, with data from five additional trials, a longer follow-up in one trial, and data on cardiac death, myocardial infarction, stent thrombosis, and target lesion revascularisation as additional clinical outcomes.

To address earlier concerns we presupposed overall mortality as the primary safety outcome and systematically explored the consistency of mortality data in people with diabetes.

METHODS

We included randomised controlled trials in people with symptoms or signs of myocardial ischaemia as a result of coronary artery disease, that compared the two first generation drug eluting stents approved by the US Food and Drug Administration, a paclitaxel eluting stent (Taxus; Boston Scientific, Natick, MA) and a sirolimus eluting stent (Cypher; Cordis, Miami Lakes, FL) with each other or with a bare metal stent. Trials had to have a clinical follow-up duration of at least six months.

We searched Medline, Embase, the Cochrane Central Register of Controlled Trials (from inception of each database to October 2007), and relevant websites (www.acc.org, www.tctmd.com, www.theheart.org, www.clinicaltrialsresults.org) for studies in any language. We checked reference lists, conference abstracts, relevant reviews, book chapters, and the proceedings of the relevant advisory panels of the Food and Drug Administration, and we contacted manufacturers and trialists. (See web extra appendix 1 for details of the search strategy.)

Data extraction

Two investigators (CSt, SA) extracted data independently, with disagreements resolved in consultation with a third investigator (PJ). We asked the trialists and manufacturers of drug eluting stents to check the extracted information and to provide additional outcome data on an electronic form according to standardised definitions and attempted to obtain outcome data separately for people with and without diabetes.

We specified overall mortality as the primary safety outcome and target lesion revascularisation as the primary effectiveness outcome. Target lesion revascularisation was defined as repeat percutaneous intervention of the target lesion or bypass surgery of the target vessel done for restenosis or other complications of the target lesion (ranging from 5 mm proximal to 5 mm distal to the stent). We recorded the following secondary safety outcomes: cardiac death, defined as any death due to a cardiac cause (for example, myocardial infarction, low output failure, fatal arrhythmia), procedure related deaths, deaths related to concomitant treatment, and death of unknown cause; myocardial infarction, including fatal and non-fatal non-Q wave or Q wave myocardial infarction; a composite of death or myocardial infarction; and stent thrombosis, within the stented segment, confirmed by angiography or post-mortem examination in accordance with the criteria of the Academic Research Consortium, to ensure the inclusion of “secondary” stent thrombosis occurring after a patient had undergone a target lesion revascularisation. In addition, we recorded stent thrombosis according to the definitions used in individual trials (per protocol definition). The numbers of patients experiencing an event and the overall number of patients at risk were recorded separately for years 1 to 4.

To tackle potential sources of inconsistency in the network we determined the month of completion of patient recruitment as a measure of the period when a trial was done, the duration of follow-up in years, and the duration in months of dual antiplatelet therapy after stent implantation and assessed three key domains of internal validity: concealment of allocation, blinding of research staff adjudicating clinical outcomes, and the inclusion of all randomised participants in the analysis according to the intention to treat principle. We considered trials to be of high quality that satisfied all three criteria [see also the criteria for quality assessment in the web extra, appendix 1]. In most trials the precise durations of dual antiplatelet therapy in individual patients were unavailable. Therefore we obtained all relevant auxiliary information available for trials with a duration of dual antiplatelet therapy of less than six months, such as the information stated in the protocol and the percentage of patients reporting the use of dual antiplatelet therapy at relevant time points. On the basis of this information two investigators (CSt, PJ) independently classified trials according to the likely percentage of patients taking clopidogrel at six months, in increments of 10% (<10%, 10% to <20%, and so on).

Statistical analysis

We used a hierarchical random effects model based on piecewise...
constant hazards, with random effects at the levels of trials, adjacent time periods, and comparisons. We simultaneously estimated log hazard ratios for people with and without diabetes and the difference in log hazard ratios between such people. From the posterior distribution of the difference we estimated the P values for interaction between treatment effect and diabetes status. We excluded from the analyses time periods with zero events in both groups. Hazard ratios were estimated from the median, and the accompanying 95% credibility intervals from the 2.5th and 97.5th centiles of the posterior distribution. Because of low event rates we derived relative risks of stent thromboses using a previously described random effects Poisson regression model.1 As in the preceding study,1 we did separate analyses according to time of occurrence of stent thrombosis and an analysis of per protocol definitions of stent thrombosis as used in individual trials (see web extra appendix 1 for details of the models).

We evaluated the inconsistency of the network, defined as the variability of results across different comparisons of the network, by calculating inconsistency factors: the estimated difference between the log hazard ratios from direct comparisons within randomised trials and the log hazard ratios from indirect comparisons between randomised trials with one intervention in common.1 To allow intuitive interpretation we back transformed absolute values of these inconsistency factors to ratios of hazard ratios and expressed inconsistency as percentage difference in hazard ratios between direct randomised comparisons within trials and indirect comparisons between trials. Values can range from 0% to infinity. A value near 0 indicates that all the comparisons in the network are consistent, showing fully coherent estimates of hazard ratios comparing any two types of stent. The more the value deviates from 0% the more inconsistent the network. A value of 25% corresponding to a ratio of hazard ratios of 1.25, may be interpreted to indicate low inconsistency, a value of 50% moderate inconsistency, and a value of 100%, corresponding to a ratio of hazard ratios of 2.00, high inconsistency (see web extra appendix 1). We evaluated heterogeneity between trials, defined as variability of results across trials within comparisons over and above chance, and the goodness of fit of the model to the data (see web extra appendix 1).

To investigate potential sources of variation in the network, we included the characteristics of the trials as covariates in the network meta-analysis of the primary safety outcome. We used unspecified cut-off points of two years for the length of follow-up, January 2004 for completion of patient recruitment, and six months for the duration of dual antiplatelet therapy. In some instances the numbers of trials and events were too low to allow the estimation of random effects at the level of time periods. Therefore we used a random effects Poisson regression model (see web extra appendix 1) for all of these analyses.11 P values for interaction between trial characteristics and treatment effect were derived from the posterior distribution of covariates.

The duration of dual antiplatelet therapy specified in trial protocols was the only variable with a treatment by trial characteristic interaction at P<0.05. Therefore we restricted the dataset to trials with a duration of dual antiplatelet therapy of six months or longer and repeated all analyses. Heterogeneity between trials, defined as variability of results across trials within comparisons over and above chance, and the goodness of fit of the model to the data, were evaluated as previously reported (see web extra appendix 1). All analyses were done in WinBUGS version 1.4.1 and Stata version 9.2.

**RESULTS**

Forty two trials met the inclusion criteria (fig 1). Seven trials, totalling about 900 people with diabetes and 3000 without, were excluded because data stratified by diabetes status were not obtained.w15 w33 w34 w35 w36 w37 w38 w39 w40 w41 w42 The remaining 35 trials w1 w2 w3 w4-w14 w16-w33 w35 w36 w37 w38 w39 w40 were included. Investigators or manufacturers provided additional data for 32 trials. w1 w3 w4-w14 w16-w30 w32 w33 w35 w36 w37 w38 w39 w40 (The characteristics of the 35 included trials are presented in web extra table A.) Four trials w1 w2 w3 w4 included only people with diabetes and one trial only people without diabetes. w33 The trials had randomly allocated 3832 people with diabetes mellitus and 10947 people without diabetes mellitus. Patient recruitment started between August 2000w16 and October 2004w14 and was completed between January 2001w16 and November 2005. w5 Data stratified according to the presence or absence of diabetes were available for all 35 trials on all outcomes, except stent thrombosis (see web extra tables B and C). Twenty nine trials described appropriate methods for allocation concealment, w1 w2 w7 w8 w11-w14 w16-w23 w25 w26 w28-w30 w32 w33 w35 w38 w40 and 28 trials reported blind adjudication for clinical outcomes. w1 w2 w4 w7 w9 w14 w16-w26 w28-w30 w32 w33 w35 w38 w40. For 30 trials all randomised patients could be included in the analyses according to the intention to treat principle.w1 w2 w4 w7 w9 w11 w13 w14 w16-w23 w25 w26 w28-w30 w32 w33 w35 w38 w40 Twenty four trials were considered to be of high quality.w1 w2 w7 w11 w13 w14 w16-w23 w25 w26 w28-w30 w32 w33 w35 w38 w40

**Duration of dual antiplatelet therapy**

The duration of dual antiplatelet therapy specified in the study protocols was two months in five trials, w6 w8 w20 w22 three months in three, w3 w7 w17 six months in 18, w2 w4-7 w9 w14 w22 w24 w26 w29 w32 w33 w35 w38 w40 nine months in one, w1 w4 and 12 months in eight. w1 w6 w8 w22 w25 w27 w30 w31 All eight trials with therapy lasting less than six months compared sirolimus eluting stents with bare metal stents. Auxiliary information for these trials (see web extra appendix 2) indicated that the percentage of patients actually receiving dual antiplatelet therapy of six months or more was likely to be below 10% in five trials. w7 w16-w19 The likely percentage for the other three trials was between 10% and 20%, w24-27 40% and 50%, w20 and 50% and 60%.11 One head to head comparison had specified a minimal duration of dual antiplatelet therapy of two months for sirolimus eluting stents.
and six months for paclitaxel eluting stents in the protocol, but the actual duration was only one month shorter for patients allocated to sirolimus stents than for patients allocated to paclitaxel stents, and about 50% of patients allocated to either stent type were still receiving therapy after eight months.\textsuperscript{w12}

**Network of all trials: overall mortality**

Table 1 presents the results of the network meta-analysis of overall mortality in people with and without diabetes. In people with diabetes the estimated hazard ratio for sirolimus eluting compared with bare metal stents was 1.14 (95% credibility interval 0.74 to 1.60), for paclitaxel eluting versus bare metal stents was 1.09 (0.71 to 1.66), and for sirolimus versus paclitaxel eluting stents was 1.02 (0.70 to 1.57), and compatible with both a substantial harm and a moderate benefit of either eluting stent compared with bare metal stents. The corresponding values for people without diabetes were 1.02 (0.77 to 1.29), 0.90 (0.67 to 1.16), and 1.13 (0.83 to 1.54; table 1). A moderate to high inconsistency of 61% was found among people with diabetes, but none among people without diabetes.

**Exploration of sources of variation**

Table 2 presents an investigation into potential sources of variation in people with diabetes in the network. Estimates of relative risk comparing sirolimus eluting stents with paclitaxel eluting stents depended to some extent on the quality of the trials, the length of follow-up, and the time of completion of patient recruitment (table 2), but 95% credibility intervals were wide and tests for interaction negative (P for interaction ≥0.16). The estimated relative risk of death when sirolimus eluting stents were compared with bare metal stents was greater when the specified duration of dual antiplatelet therapy was less than six months (2.37, 95% credibility interval 1.18 to 5.12) compared with six months or longer (0.89, 0.58 to 1.40, P for interaction 0.02), however. When three trials originally classified to have a short duration of dual antiplatelet therapy\textsuperscript{w13} were reclassified to have a duration of six months or longer, reflecting the auxiliary information indicating that more than 10% of patients in these trials were still receiving dual antiplatelet therapy at six months, differences were maintained.

**Restricted network: overall mortality**

When the network was restricted to trials with dual antiplatelet therapy for six months or longer (table 1), the hazard ratios of death overall among people with diabetes were all below 1: sirolimus eluting stents compared with bare metal stents 0.88 (95% credibility interval 0.55 to 1.30), paclitaxel eluting stents compared with bare metal stents 0.91 (0.60 to 1.38), and sirolimus eluting compared with paclitaxel eluting stents 0.95 (0.63 to 1.43). Compared with the network of all trials the inconsistency decreased to 20% and credibility intervals of hazard ratios became more narrow in the restricted network. Among patients without diabetes results were much the same in the overall and the restricted network (table 1). The hazard ratio for sirolimus eluting stents compared with bare metal stents was 1.05 (0.69 to 1.73), for paclitaxel eluting stents compared with bare metal stents it was 0.89 (0.66 to 1.18), and for sirolimus compared with paclitaxel eluting stents it was 1.23 (0.82 to 1.69). Figure 2 presents corresponding cumulative incidences of death for the three stent types estimated from the restricted network meta-analysis separately for people with and without diabetes. The incidence of death was about twice as high in people with diabetes compared with people without diabetes. Tests for interaction between treatment effect and diabetes status were negative for all comparisons (P for interaction ≥0.28; also see web extra table D).

**Restricted network: secondary safety outcomes**

Table 1 allows a comparison of the results from the network meta-analysis of all trials and the analysis restricted to trials with a dual antiplatelet therapy of six months or more. Among people with diabetes, hazard ratios for drug eluting stents compared with bare metal stents became more beneficial for drug eluting stents for the outcomes of cardiac death, the composite of death or myocardial infarction, and for stent thromboses. The inconsistency decreased mainly for cardiac death and per protocol definitions of stent thromboses. No differences between overall and restricted network meta-analysis were observed for myocardial infarction. Among people without diabetes, results from overall and restricted network meta-analysis were similar. Corresponding cumulative incidences for the three...
target lesion revascularisation: patients with diabetes were included in the comparison of sirolimus eluting stents with bare metal stents, network inconsistency was low, and results were unaffected by the restriction of the analysis to trials with a duration of dual antiplatelet therapy of six months or more in people with and without diabetes (table 1). Differences between sirolimus and paclitaxel

Table 1 | Analyses overall and restricted to trials with dual antiplatelet therapy of at least six months

<table>
<thead>
<tr>
<th>Variable and stent types</th>
<th>People with diabetes</th>
<th>People without diabetes</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>All trials</td>
<td>Trials with dual antiplatelet therapy ≥ 6 months</td>
</tr>
<tr>
<td></td>
<td>Hazard ratio (95% CI)</td>
<td>IC*</td>
</tr>
<tr>
<td>Death overall:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>SES v bare metal stent</td>
<td>1.14 (0.74 to 1.60)</td>
<td>0.88 (0.55 to 1.30)</td>
</tr>
<tr>
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</tr>
<tr>
<td>SES v PES</td>
<td>1.02 (0.70 to 1.57)</td>
<td>0.95 (0.63 to 1.43)</td>
</tr>
</tbody>
</table>

Cardiac death:           |                      |                        |                      |                        |
| SES v bare metal stent   | 1.09 (0.63 to 1.93)  | 0.80 (0.42 to 1.57)    | 0.88 (0.48 to 1.60)  | 0.93 (0.45 to 1.81)    |
| PES v bare metal stent   | 1.08 (0.62 to 2.28)  | 0.94 (0.52 to 1.87)    | 0.90 (0.52 to 1.54)  | 0.89 (0.55 to 1.47)    |
| SES v PES                | 0.98 (0.46 to 2.08)  | 0.85 (0.50 to 1.55)    | 0.96 (0.58 to 1.84)  | 1.04 (0.54 to 1.95)    |

Myocardial infarction:   |                      |                        |                      |                        |
| SES v bare metal stent   | 0.68 (0.44 to 1.05)  | 0.68 (0.43 to 1.12)    | 0.87 (0.64 to 1.20)  | 0.81 (0.55 to 1.14)    |
| PES v bare metal stent   | 0.84 (0.55 to 1.31)  | 0.85 (0.54 to 1.43)    | 1.08 (0.74 to 1.51)  | 1.05 (0.72 to 1.42)    |
| SES v PES                | 0.79 (0.56 to 1.23)  | 0.80 (0.55 to 1.27)    | 0.81 (0.58 to 1.06)  | 0.75 (0.57 to 1.07)    |

Death or myocardial infarction: |                      |                        |                      |                        |
| SES v bare metal stent   | 1.04 (0.75 to 1.61)  | 0.88 (0.57 to 1.27)    | 0.93 (0.71 to 1.13)  | 0.91 (0.69 to 1.13)    |
| PES v bare metal stent   | 1.07 (0.74 to 1.62)  | 0.91 (0.70 to 1.31)    | 1.04 (0.84 to 1.31)  | 1.00 (0.84 to 1.25)    |
| SES v PES                | 0.97 (0.72 to 1.34)  | 0.95 (0.69 to 1.27)    | 0.90 (0.71 to 1.09)  | 0.88 (0.71 to 1.06)    |

Stent thrombosis (ARC definition): |                      |                        |                      |                        |
| SES v bare metal stent   | 0.46 (0.15 to 1.42)  | 0.33 (0.09 to 1.09)    | 1.35 (0.76 to 2.73)  | 1.24 (0.58 to 3.08)    |
| PES v bare metal stent   | 1.05 (0.32 to 4.01)  | 0.82 (0.23 to 3.09)    | 1.56 (0.83 to 3.13)  | 1.48 (0.69 to 3.40)    |
| SES v PES                | 0.44 (0.15 to 1.17)  | 0.40 (0.13 to 1.08)    | 0.87 (0.47 to 1.69)  | 0.84 (0.41 to 1.88)    |

Stent thrombosis (per protocol): |                      |                        |                      |                        |
| SES v bare metal stent   | 0.48 (0.17 to 1.35)  | 0.20 (0.05 to 0.68)    | 1.43 (0.78 to 3.00)  | 1.48 (0.74 to 3.41)    |
| PES v bare metal stent   | 1.27 (0.38 to 4.91)  | 0.73 (0.19 to 2.80)    | 1.73 (0.88 to 3.61)  | 1.80 (0.89 to 3.67)    |
| SES v PES                | 0.38 (0.11 to 1.07)  | 0.27 (0.07 to 0.80)    | 0.82 (0.44 to 1.73)  | 0.82 (0.44 to 1.73)    |

Target lesion revascularisation: |                      |                        |                      |                        |
| SES v bare metal stent   | 0.29 (0.22 to 0.39)  | 0.29 (0.19 to 0.43)    | 0.29 (0.22 to 0.38)  | 0.29 (0.19 to 0.42)    |
| PES v bare metal stent   | 0.38 (0.28 to 0.55)  | 0.38 (0.26 to 0.56)    | 0.46 (0.33 to 0.60)  | 0.46 (0.32 to 0.60)    |
| SES v PES                | 0.76 (0.53 to 1.05)  | 0.78 (0.50 to 1.14)    | 0.63 (0.49 to 0.82)  | 0.64 (0.49 to 0.84)    |

SES=sirolimus eluting stent; PES=paclitaxel eluting stent; ARC=Academic Research Consortium.

*Inconsistency of network expressed as percentage difference in hazard ratios between direct randomised comparisons within trials and indirect comparisons between trials. Values near 0 indicate that all comparisons in network are consistent, showing fully coherent estimates of hazard ratios comparing any two stent types. Values can range from 0% to infinity. The more values deviate from 0%, the more inconsistent the network. A value of 25% may be interpreted to indicate low inconsistency, 50% moderate, and 100% high inconsistency. (Also see table 5 in web extra appendix 3 for 95% credibility intervals and P values of inconsistency estimates.)

Table 3 presents a breakdown of stent thromboses according to time of occurrence. Among people with diabetes little evidence was found for an increased risk of definite or per protocol stent thrombosis associated with sirolimus eluting stents compared with either of the two other stents; all point estimates were below 1 and differences in favour of sirolimus eluting stents became more pronounced with the use of per protocol definitions. For the comparison of paclitaxel eluting stents with bare metal stents all estimates were imprecise for both the Academic Research Consortium definition of definite stent thrombosis and per protocol definitions. Among people without diabetes relative risks were generally higher for both definitions, but tests for interaction between treatment effect and diabetes status were positive only for the comparison of sirolimus eluting stents with bare metal stents on per protocol definitions of stent thrombosis between day 0 and 4 years and between day 30 and 4 years (P for interaction=0.01, see also web extra table D).

Restricted network: target lesion revascularisation: Both drug eluting stents were robustly associated with a decrease in revascularisation rates compared with bare metal stents, network inconsistency was low, and results were unaffected by the restriction of the analysis to trials with a duration of dual antiplatelet therapy of six months or more in people with and without diabetes (table 1). Differences between sirolimus and paclitaxel
**DISCUSSION**

Our collaborative network meta-analysis suggests that previously reported increases in the risk of death associated with sirolimus eluting stents compared with bare metal stents, which translates into a number needed to harm as low as 7 to cause one death over four years. Conversely, trials with dual antiplatelet therapy for six months or more showed no increase in risk from using sirolimus eluting stents compared with bare metal stents. Restricting the network to trials with dual antiplatelet therapy of six months or longer resulted in a smaller evidence base: eight trials of 613 people with diabetes were excluded from the network. Despite this, statistical precision was improved owing to the accompanying decrease in the network’s inconsistency.

Compared with bare metal stents, target lesion revascularisation rates are strongly decreased by use of sirolimus and paclitaxel drug eluting stents in people with and without diabetes. Numbers needed to treat to reduce one event over four years are 6 in people with diabetes and 8 in people without diabetes. Active angiographic follow-up increases the absolute rates of cardiac death and stent thromboses. The restriction of the network resulted in a smaller evidence base: eight trials of 613 people with diabetes were excluded from the analysis. Despite this, statistical precision was improved owing to the accompanying decrease in the network’s inconsistency.
mandatory angiography. The number needed to treat to avoid one revascularisation would therefore be somewhat lower in clinical routine. Assuming revascularisation rates of 12% in people with diabetes and 9% in people without diabetes, as found in the Cardiac Care Network of Ontario at two years, numbers needed to treat were estimated as 13 for people with diabetes and 18 for people without diabetes.

Our study comprises a large body of evidence from randomised controlled trials in people with and without diabetes treated with one of two drug eluting stents or bare metal stents. Investigators and manufacturers provided additional data according to uniform outcome definitions, including a standardised definition for stent thrombosis according to the Academic Research Consortium consensus. This increases comparability between trials and limits bias, such as the censoring of events after intervening revascularisation.

Our previous network meta-analysis was recently updated to include additional trials and new stent types estimated from network meta-analysis for pairwise comparisons in people with and without diabetes and restricted to trials with dual antiplatelet therapy of at least six months.

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duration of dual antiplatelet therapy particularly in early trials.

Our exploration of inconsistency is observational in nature and has the same limitations as other observational studies. Most importantly, earlier trials had specified shorter durations of dual antiplatelet therapy than later trials. The duration of therapy was therefore bound to be negatively correlated with the duration of follow-up, and confounding could exist between the duration of therapy and the length of follow-up. Other potential confounders include changes over time in patient selection and procedural characteristics, such as an under-sizing or under-expansion of stents in early trials, or methodological quality. We addressed this by repeating tests of interaction between treatment effect and components of methodological quality or length of follow-up after the exclusion of trials with a duration of dual antiplatelet therapy of less than six months and found no evidence for an interaction in any of these analyses (data available on request).

We acknowledge that our results could be corroborated by an analysis of the actual duration of dual antiplatelet therapy in individual patients, but precise durations in individual patients are unavailable in most trials and we lacked the resources to retrospectively ascertain and validate usage data. Eight trials had specified a duration of dual antiplatelet therapy of six months or longer, and the P value for interaction between relative risk of death and duration of dual antiplatelet therapy became even smaller. Additionally, strut thickness or type of bare metal stent used in comparison groups might affect clinical outcomes. Even though our results are robust to the adjustment for these characteristics of bare metal stents, we cannot fully exclude the possibility that differences in bare metal stents as comparators contributed to the observed variation in...

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**Figure 3** Cumulative incidence of myocardial infarction and the composite of death or myocardial infarction and corresponding hazard ratios (95% credibility intervals) for three stent types estimated from network meta-analysis for pairwise comparisons in people with and without diabetes and restricted to trials with dual antiplatelet therapy of at least six months.
Table 3 | Stent thromboses in trials with dual antiplatelet therapy of six months or more

<table>
<thead>
<tr>
<th>Variable</th>
<th>Events</th>
<th>Relative risks (95% credibility interval)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>BMS</td>
<td>PES</td>
</tr>
<tr>
<td><strong>ARC definite stent thrombosis</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Patients with diabetes:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>No of patients at risk</td>
<td>557</td>
<td>874</td>
</tr>
<tr>
<td>0–4 days</td>
<td>13</td>
<td>17</td>
</tr>
<tr>
<td>&gt;30 days</td>
<td>2</td>
<td>8</td>
</tr>
<tr>
<td>Patients without diabetes:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>No of patients at risk</td>
<td>2439</td>
<td>3130</td>
</tr>
<tr>
<td>0–4 days</td>
<td>34</td>
<td>56</td>
</tr>
<tr>
<td>&gt;30 days</td>
<td>19</td>
<td>22</td>
</tr>
</tbody>
</table>

**Per protocol definition of stent thrombosis†**

<table>
<thead>
<tr>
<th>Variable</th>
<th>Events</th>
<th>Relative risks (95% credibility interval)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>BMS</td>
<td>PES</td>
</tr>
<tr>
<td>Patients with diabetes:</td>
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<td></td>
</tr>
<tr>
<td>No of patients at risk</td>
<td>723</td>
<td>912</td>
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<tr>
<td>0–4 days</td>
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<td>18</td>
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<tr>
<td>&gt;30 days</td>
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<td>10</td>
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<tr>
<td>Patients without diabetes:</td>
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<td></td>
</tr>
<tr>
<td>No of patients at risk</td>
<td>2577</td>
<td>3382</td>
</tr>
<tr>
<td>0–4 days</td>
<td>29</td>
<td>58</td>
</tr>
<tr>
<td>&gt;30 days</td>
<td>22</td>
<td>24</td>
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</tbody>
</table>

BMS = bare metal stent; PES = paclitaxel eluting stent; SES = sirolimus eluting stent.
*According to Academic Research Consortium criteria.
†According to protocol definitions used in individual trials.

mortality between trials with short and long durations of dual antiplatelet therapy. Four trials included only people with diabetes and one trial only people without diabetes. Performing network meta-analyses separately for people with and without diabetes allowed us to also incorporate these trials in our analysis. An alternative approach would have been to model differences between people with and without diabetes directly within each trial, but at the price of excluding these five trials. A final limitation of our study is that we were unable to record information on specific antidiabetic treatment or on glycaemic control in people with diabetes mellitus and to perform separate analyses for people with diabetes who did or did not use insulin. Although these aspects are related to cardiovascular outcomes, they were per
reduction was 19%. Other mechanisms will therefore need to be balanced against potential risks, such as clinically relevant bleeding. The optimal duration can only be determined in adequately powered large scale randomised controlled trials.

Conclusion

In trials with a duration of dual antiplatelet therapy of six months or longer drug eluting stents were safe and effective in people with and without diabetes. It seems prudent to adhere to a minimal duration of dual antiplatelet therapy of six months in patients undergoing implantation of a drug eluting coronary stent.

The potential benefits of a longer duration of therapy need to be balanced against potential risks, such as clinically relevant bleeding. The optimal duration can only be determined in adequately powered large scale randomised controlled trials.

We thank Boston Scientific and Cordis for the provision of additional data. CTU Bern is supported by the Swiss National Science Foundation.

Contributors: CST and SA contributed equally to the manuscript. PJ and CS conceived the study. PJ, CST, SWa, SA, and SWi were responsible for the conception and design of the study. SWa, CST, SA, and PJ did the analysis and interpreted the analysis in collaboration with SWi. CST, SA, AX, MCM, AS, MEP, GWS, MBL, JSIL, JIG, SIP, MS, MIS, HK, CSg, MM, PV, MTD, PC, ASP, AJN, PD, BM, SWi, and PJ were responsible for the acquisition of data. PJ, CST, SA, and SWi wrote the first draft of the manuscript. All authors critically revised the manuscript for important intellectual content and approved the final version of the manuscript. CST and PJ obtained public funding. PJ, CST, PD, BM, and SWi provided administrative, technical, and logistic support.

Funding: Swiss National Science Foundation (grant Nos 3233BO-115212, 3233-066377, and 3200-066378) to CST and PJ. The implementation and validation of the statistical models used for this study were funded by the Swiss National Science Foundation’s national research programme 53 (grant No 405340-104762) to PJ. CST and PJ are PROSPER (program for social medicine, preventive and epidemiological research) fellows funded by the Swiss National Science Foundation. The funder had no role in study design, data collection, data analysis, data interpretation, or writing of the report.

Competing interests: CST and PJ report receiving unrestricted grants from the Swiss National Science Foundation. AK receives lecture fees from Bristol-Myers Squibb, Cordis, GlaxoSmithKline, Lilly, Medtronic, Novartis, and validation of the statistical models used for this study were funded by the Swiss National Science Foundation’s national research programme 53 (grant No 405340-104762) to PJ. CST and PJ are PROSPER (program for social medicine, preventive and epidemiological research) fellows funded by the Swiss National Science Foundation. The funder had no role in study design, data collection, data analysis, data interpretation, or writing of the report.

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WHAT IS ALREADY KNOWN ON THIS TOPIC

People with diabetes are at an increased risk for coronary heart disease and have more restenoses after the implantation of coronary stents

A meta-analysis suggested a strongly increased risk of death associated with sirolimus eluting stents compared with bare metal stents in people with diabetes

WHAT THIS STUDY ADDS

Reported increases in the risk of death associated with drug eluting stents compared with bare metal stents in people with diabetes were probably due to dual antiplatelet therapy lasting less than six months in early trials

In trials with dual antiplatelet therapy for six months or longer drug eluting stents were safe and effective in people both with and without diabetes

In clinical practice it seems prudent to adhere to a minimal duration of dual antiplatelet therapy than six months in early trials

and Sanofi-Aventis. MCM receives lecture fees from Cordis, Boston Scientific, and Abbott, which go to a research organisation (RCF, Massy, France). AS receives unrestricted grant support for the Department of Cardiology he chairs from Amersham/General Electric, Bayerische Forschungsgstiftung, Bristol-Myers Squibb, Cordis, CryoCath, Guidant, Medtronic, Nycomed, and Schering. MEP receives lecture fees from Medtronic. GWS receives consulting fees from Boston Scientific, Abbott, Guidant, and BMS Imaging, lecture fees from Boston Scientific, Abbott, and Medtronic, has equity interests in Devax and Xent, and is a member of the board of directors of Devax. MBL receives consulting fees from Cordis, Medtronic, Boston Scientific, and OrbisNerich and has equity interests in Conor, Medinol, and OrbusNerich. JG is on the advisory board of Boston Scientific and receives research grant support from Cordis. SJP receives research grant support from Cordis. HK receives unrestricted grant support from Cordis. CSP receives consulting and lecture fees from Cordis, Boston Scientific, Abbott, Lilly, and Pfizer. MTD receives lecture fees from Boston Scientific. BM receives research grant support from various stent companies, including Cordis and Boston Scientific, and is in the speaker bureau for various stent companies, including Cordis and Boston Scientific. SW receives lecture and consulting fees from Abbott, Biotronic, Biosensors, Boston Scientific, Cordis, and Medtronic. GWS and MBL are directors of the Cardiovascular Research Foundation, a public charity affiliated with Columbia University Medical Center, from which they receive no compensation; the Cardiovascular Research Foundation receives research or educational funding from Boston Scientific, Cordis, Sanofi-Aventis, and Biosil-Myers Squibb.

Ethical approval: Not required.

Provenance and peer review: Not commissioned; externally peer reviewed.