

of first-generation DES did—show a benefit with respect to a 12-month clinical endpoint compared with BMS. On the basis of these discrepant findings and the different pathophysiology of SVG and native coronary vessels, the ideal stent type for SVG disease could be hypothesised to be different from that for native vessel disease. However, this conclusion is speculative. Further research comparing the effect of different stent types in SVG disease is warranted.

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- 1 Palmerini T, Benedetto U, Biondi-Zoccai G, et al. Long-term safety of drug-eluting and bare-metal stents: evidence from a comprehensive network meta-analysis. *J Am Coll Cardiol* 2015; **65**: 2496–2507.
- 2 Windecker S, Kolh P, Alfonso F, et al. 2014 ESC/EACTS guidelines on myocardial revascularization: the task force on myocardial revascularization of the European Society of Cardiology (ESC) and the European Association for Cardio-Thoracic Surgery (EACTS). Developed with the special contribution of the European Association of Percutaneous Cardiovascular Interventions (EAPCI). *Eur Heart J* 2014; **35**: 2541–619.
- 3 Vermeersch P, Agostoni P, Verhey S, et al. Randomized double-blind comparison of sirolimus-eluting stent versus bare-metal stent implantation in diseased saphenous vein grafts: six-month angiographic, intravascular ultrasound, and clinical follow-up of the RRISC trial. *J Am Coll Cardiol* 2006; **48**: 2423–31.

- 4 Brilakis ES, Lichtenwalter C, de Lemos JA, et al. A randomized controlled trial of a paclitaxel-eluting stent versus a similar bare-metal stent in saphenous vein graft lesions the SOS (Stenting of Saphenous Vein Grafts) trial. *J Am Coll Cardiol* 2009; **53**: 919–28.
- 5 Mehilli J, Pache J, Abdel-Wahab M, et al. Drug-eluting versus bare-metal stents in saphenous vein graft lesions (ISAR-CABG): a randomised controlled superiority trial. *Lancet* 2011; **378**: 1071–78.
- 6 Jeger R. Drug-eluting vs. bare metal stents in saphenous vein grafts: the prospective randomized BASKET-SAVAGE trial. European Society of Cardiology Congress; Rome, Italy; Aug 27–31, 2016.
- 7 Brilakis ES, Edson R, Bhatt DL, et al. Drug-eluting stents versus bare-metal stents in saphenous vein grafts: a double-blind, randomised trial. *Lancet* 2018; published online May 11. [http://dx.doi.org/10.1016/S0140-6736\(18\)30801-8](http://dx.doi.org/10.1016/S0140-6736(18)30801-8).
- 8 Sarjeant JM, Rabinovitch M. Understanding and treating vein graft atherosclerosis. *Cardiovasc Pathol* 2002; **11**: 263–71.
- 9 de Feyter PJ. Percutaneous treatment of saphenous vein bypass graft obstructions: a continuing obstinate problem. *Circulation* 2003; **107**: 2284–86.
- 10 Jonas M, Stone GW, Mehran R, et al. Platelet glycoprotein IIb/IIIa receptor inhibition as adjunctive treatment during saphenous vein graft stenting: differential effects after randomization to occlusion or filter-based embolic protection. *Eur Heart J* 2006; **27**: 920–28.
- 11 Banerjee S, Xu H, Fuh E, et al. Endothelial progenitor cell response to antiproliferative drug exposure. *Atherosclerosis* 2012; **225**: 91–98.
- 12 Vermeersch P, Agostoni P, Verhey S, et al. Increased late mortality after sirolimus-eluting stents versus bare-metal stents in diseased saphenous vein grafts: results from the randomized DELAYED RRISC trial. *J Am Coll Cardiol* 2007; **50**: 261–67.
- 13 Colleran R, Kufner S, Mehilli J, et al. Long-term outcomes of a randomized comparison of drug-eluting stents versus bare metal stents in saphenous vein graft lesions—5-year follow-up of the ISAR-CABG trial. *Heart* 2017; **103** (suppl 6): A23–24.
- 14 Brilakis ES, Lichtenwalter C, Abdel-karim AR, et al. Continued benefit from paclitaxel-eluting compared with bare-metal stent implantation in saphenous vein graft lesions during long-term follow-up of the SOS (Stenting of Saphenous Vein Grafts) trial. *JACC Cardiovasc Interv* 2011; **4**: 176–82.



Reducing child mortality in high-income countries: where to from here?



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In 2000, the UN envisioned a path of global action to eradicate inequalities and fight the many dimensions of poverty with the Millennium Development Goals (MDGs), which have served as an overarching framework for health, social, and economic development.¹ During the MDG era, the global under-5 child mortality rate declined by an impressive 53%,² but still fell short of the targeted two-thirds reduction from 1990 to 2015.³ Progress in improving child survival has been uneven and inequalities persist across both developing and high-income countries.

In their birth cohort study of under-4 child mortality in Sweden and England, Ania Zylbersztejn and colleagues⁴ highlight the disparity in mortality among infants and children under the age of 4 years between these two countries. Among 3 932 886 births in England there were 11 392 deaths, and among 1 013 360 births

in Sweden there were 1927 deaths. The unadjusted hazard ratios (HRs) for England versus Sweden were 1.66 (95% CI 1.53–1.81) at 2–27 days, 1.59 (1.47–1.71) at 28–364 days, and 1.27 (1.15–1.40) at 1–4 years. Zylbersztejn and colleagues sought to establish the drivers of these differences. Using adverse birth characteristics (birthweight, gestational age, and congenital anomalies) and sex as proxies for risk factors occurring before birth, and socioeconomic factors (maternal age and socioeconomic status) as measures of risk factors after birth, they report that the most important driver of the excess mortality in England relative to Sweden is the prevalence of adverse birth characteristics. In the first month of life, 77% of the excess risk of death in England compared with Sweden was explained by birth characteristics and a further 3% by socioeconomic

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factors. From the second month to the end of the first year of life, 68% of the excess risk was explained by birth characteristics and a further 11% by socioeconomic factors. At 1–4 years, there was no significant difference between countries.⁴

The findings of this study are important because they suggest efforts made to improve maternal health before and during pregnancy offer the most promise for reducing child mortality in England.⁴ Such efforts could support chronic disease management and mental health and addiction treatment, improve pre-pregnancy and antenatal nutrition, encourage smoking cessation, and ensure pregnant women receive adequate prenatal care.⁵ But although the authors report that prenatal factors have a larger role in child mortality than socioeconomic factors present after birth,⁴ the results do not discount the important influence of social factors. The authors point to the extensive literature connecting adverse birth characteristics to socioeconomic factors, highlighting the importance of efforts to reduce poverty. Poverty reduction strategies for women during the prenatal period have shown great promise in reducing adverse birth characteristics⁶ and increasing maternal health-care-seeking behaviours during pregnancy.⁷

The authors also emphasise the benefits that efforts to reduce adverse birth characteristics could have for child health and wellbeing, and that these benefits extend far beyond reducing child mortality.⁴ A large body of evidence, particularly around low birthweight and preterm birth, shows that adverse birth outcomes can continue to have impacts throughout childhood and into adulthood, including increased risks of poorer physical and mental health,^{8–11} lower educational attainment,^{8,10,11} and lower income.⁸ What this study⁴ could not ascertain from the analysis was whether adverse birth characteristics remain the most important driver of suboptimal health and wellbeing for those children who survive. Indeed, other research suggests that social factors operating after birth pose a similar threat to child health and education outcomes as preterm birth and low birthweight; however, because these social factors are far more prevalent, their role in child outcomes at the population level far outweighs the role of adverse birth characteristics.^{10,12}

Although it is clear from this study that there is room for improvement in reducing child mortality in high-income countries,⁴ more research is needed on other predictors of optimal child health and development. Routinely collected

person-level administrative data from multiple sources are useful for conducting population-based studies. Linking electronic health records and data from other sectors at the individual level adds further value for quantifying inequalities, allowing identification of the best periods in which to intervene to find out which risk factors are driving disparities between populations. The methods adopted by Zylbersztejn and colleagues⁴ can be used to compare child mortality across countries and to explore other health and wellbeing outcomes, across and within countries. This research aligns with the shift from the MDGs to the Sustainable Development Goals, which consolidate global efforts to end all forms of poverty and fight inequalities, driving the change from improving childhood survival to ensuring that children live, grow, and thrive.¹³

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MB has collaborated and co-authored papers with one of the authors (Ruth Gilbert) of the Article discussed in this Comment. We declare no other competing interests.

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- 1 UN. The Millennium Development Goals Report, 2015. New York: United Nations, 2015.
- 2 You D, Hug L, Ejdemyr S, et al. Global, regional, and national levels and trends in under-5 mortality between 1990 and 2015, with scenario-based projections to 2030: a systematic analysis by the UN Inter-agency Group for Child Mortality Estimation. *Lancet* 2015; **386**: 2275–86.
- 3 Wang H, Liddell CA, Coates MM, et al. Global, regional, and national levels of neonatal, infant, and under-5 mortality during 1990–2013: a systematic analysis for the Global Burden of Disease Study 2013. *Lancet* 2014; **384**: 957–79.
- 4 Zylbersztejn A, Gilbert R, Hjern A, Wijlaars L, Hardeid P. Child mortality in England compared with Sweden: a birth cohort study. *Lancet* 2018; published online May 3. [http://dx.doi.org/10.1016/S0140-6736\(18\)30670-6](http://dx.doi.org/10.1016/S0140-6736(18)30670-6).
- 5 Bhutta ZA, Das JK, Bahl R, et al. Can available interventions end preventable deaths in mothers, newborn babies, and stillbirths, and at what cost? *Lancet* 2014; **384**: 347–70.
- 6 Brownell MD, Chartier MJ, Nickel NC, et al. Unconditional prenatal income supplement and birth outcomes. *Pediatrics* 2016; **137**: e20152992.
- 7 Leyland AH, Ouédraogo S, Nam J, et al. Evaluation of health in pregnancy grants in Scotland: a natural experiment using routine data. Southampton: NIHR Journals Library, 2017.
- 8 Moster D, Lie RT, Markestad T. Long-term medical and social consequences of preterm birth. *N Engl J Med* 2008; **359**: 262–73.
- 9 Saigal S, Doyle LW. An overview of mortality and sequelae of preterm birth from infancy to adulthood. *Lancet* 2008; **371**: 261–69.
- 10 Jutte DP, Brownell M, Roos NP, Schippers C, Boyce WT, Syme SL. Rethinking what is important: biologic versus social predictors of childhood health and educational outcomes. *Epidemiology* 2010; **21**: 314–23.
- 11 Hack M, Flannery DJ, Schluchter M, Cartar L, Borawski E, Klein N. Outcomes in young adulthood for very-low-birth-weight infants. *N Engl J Med* 2002; **346**: 149–57.
- 12 Brownell MD, Ekuma O, Nickel N, Chartier M, Koseva I, Santos RG. A population-based analysis of factors that predict early language and cognitive development. *Early Child Res Q* 2016; **35**: 6–18.
- 13 Sidebotham P, Fraser J, Covington T, et al. Understanding why children die in high-income countries. *Lancet* 2014; **384**: 915–27.