All letters are subject to editing and may be shortened. General letters can be sent to bjgpdisc@rcgp.org.uk (please include your postal address for publication), and letters responding directly to *BJGP* articles can be submitted online via **eLetters**. We regret we cannot notify authors regarding publication. For submission instructions visit: bjgp.org/letters

Targeted holistic review of patients' medicines is well overdue

The authors are right that the evidence base for the long-term benefits of structured medication reviews (SMRs) is not particularly strong.¹ However, with the publication of the Department of Health and Social Care (DHSC) 'overprescribing' report² any primary care clinician or patient/advocate will tell you that there is much to do to support people to optimise treatments. Why? Because in our time-poor, overstretched NHS, with an ever-increasing number of 'single organologists' who invariably just add medicines, never subtracting, therefore multiplying the problems, the dedicated time for a real holistic consultation to find out what matters to a person about their medicines and health/wellbeing should not just be welcomed, but cherished.

Poor adherence to medicines and adverse drug reactions, let alone full-blown problematic polypharmacy, are rife.³ The general overestimation of the benefits of medicines and an underappreciation of their risks, coupled with imprecise systems for maintaining accurate medicines records across care sectors, also contribute to unnecessary follow-up clinician appointments and medicines waste.

Dr Louisa Polak⁴ is also right that clinicians performing SMRs need to be '*competent to interpret evidence-based guidance, but also confident enough to disregard it where they and the patient agree that following it does not serve the patient's agenda*' and that relational continuity can be particularly important. However, as an experienced clinical pharmacist, who regularly teaches GPs about polypharmacy/deprescribing, I would disagree that clinical pharmacists, the new medicines experts in primary care, cannot perform these SMRs. Yes, some less experienced may need additional training and more GP input to perform SMRs in complex patients with multimorbidity. But then many experienced GPs, who have excellent relationships with patients, also lack the confidence to stop medicines and arguably have more appropriate primary care clinical roles that only they can best perform.

I suggest that Primary Care Networks hone their plans to risk stratify which patients are most in need of an SMR, agree how many clinicians have the competency and time to provide them, then set up the maximum appointment sessions they have the capacity to provide and crack on! Our patients will thank us.

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Competing interests

Department of Health and Social Care, *Good for You, Good for Us, Good for Everybody: A Plan to Reduce Overprescribing*, 2021, short-life working group member. NHSE/I Network Contract Direct Enhanced Service SMRs, 2019/2020, short-life working group member.

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Poverty screening in primary care

Many thanks to Gopal and colleagues for providing a balanced article about poverty

screening and for providing guidance to GPs who wish to take this further.¹ However, we must separate our wish to help individuals facing poverty from our concerns about health inequalities: we cannot resolve the latter by doing the former; that just isn't how health inequalities work.

As healthcare providers, we are potentially more able to intervene usefully in healthcare inequalities (unequal access, experiences, and outcomes of health care) than in the health inequalities caused by unfair social conditions. Still, if we are concerned about inequalities in health, and our role in furthering evidence and understanding of these, then we need to acknowledge the social gradient and move on from thinking purely about people 'at the bottom'. If our concerns are to be widely shared, it will be when inequalities are recognised to impact everyone. Using a generic socioeconomic marker, such as occupation or education, and collecting this information from everyone, rather than identifying poverty alone, would not only seem fairer and less stigmatising, but would also provide comparative data.

Finally, there is no obvious reason why primary care should be the best place to identify people experiencing poverty or to signpost sources of support. Presumably HM Revenue and Customs and the Department for Work and Pensions already have a good idea of which people are facing poverty. Could they do the signposting? Or could they share information with the Department of Health and Social Care? Could the Royal College of General Practitioners propose this?

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