resources required for all aspects of such a program are available.

Sandra A. Farrell, MD, FRCPC, FCCMG
Diane E. Chadwick, PhD
Credit Valley Hospital
Mississauga, Ont.
Anne Marie Summers, MD, FRCPC, FCCMG
Philip R. Wyatt, MD, PhD
North York General Hospital
North York, Ont.

I was excited to read that the apparent sensitivity of triple-marker screening had been improved to the point that it had a rate of false-positive results of only 3.7%. This rate is supported in a table describing the results of four different studies of triple-marker screening. <sup>1-4</sup> Since I have been an opponent of this test, precisely because of its high rate of false-positive results, I decided to critically appraise and present this article to the local journal club.

Since the stated rate of falsepositive results contradicted my own experience, I reviewed the four references that served as the basis for the rate cited. In fact, the rates cited were not the results of the triplemarker test alone but of the triplemarker test in conjunction with some form of confirmation of gestational age, most commonly a subsequent ultrasonographic examination. The rate of false-positive results before further evaluation in each of these studies was approximately double the rate after ultrasonographic confirmation. These findings are more consistent with the previously published data.

Although the article by Dick and the task force was published under the rubric of clinical practice guidelines, the abstract specifically states that "the economic issues involved are complex and were not considered." How can a practice guideline be considered useful if the economic aspects involved are not considered? This would be like suggesting that

every patient with a headache undergo a computed tomography scan so that the risk of missing a brain tumour is reduced. It makes the Ottawa ankle rules practically irrelevant!

I am concerned about the impact of this guideline as published. I believe that it provides false credibility for a test that has serious limitations.

Gary Viner, BSc, MD, CCFP Assistant professor Department of Family Medicine University of Ottawa Ottawa, Ont.

## References

- Haddow JE, Palomaki GE, Knight GJ, et al. Prenatal screening for Down's syndrome with use of maternal serum markers. N Engl J Med 1992;327:588-93.
- 2. Phillips OP, Elias S, Shulman LP, Andersen RN, Morgan CD, Simpson JL. Maternal serum screening for fetal Down syndrome in women less than 35 years of age using α-fetoprotein, hCG, and unconjugated estriol: a prospective 2-year study. Obstet Gynecol 1992;80:353-8.
- 3. Wald NJ, Kennard A, Densem JW, Cuckle HS, Chard T, Butler L. Antenatal maternal serum screening for Down's syndrome: results of a demonstration project. *BMJ* 1992,305:391-4.
- Cheung EY, Luthy DA, Zebelman AM, Williams MA, Lieppman RE, Hickok DE. A prospective evaluation of a second-trimester screening test for fetal Down syndrome using maternal serum α-fetoprotein, hCG, and unconjugated estriol. Obstet Gynecol 1993,81:72-7.

## [The author responds:]

A lthough the letters from Dr. Farrell and associates and Dr. Viner represent different perspectives on screening with the use of maternal serum markers, they both raise questions about the programmatic aspects of prenatal screening for Down syndrome. There also seems to be a misunderstanding of the role of the task force guidelines. Rather than being a practical guide to local prenatal services or programs, the recommendations should be viewed as a guide to the evidence supporting and effectiveness of these interventions according to the literature.

Viner questions the task force's recommendations for triple-marker screening involving maternal serum levels of α-fetoprotein, β human chorionic gonadotropin and unconjugated estriol. He differentiates between the rate of false-positive results when the markers are used with and without confirmation of gestational age. As he notes, earlier studies of maternal serum markers without confirmation of gestational age reported higher rates of false-positive results. However, the task force used only the four recent studies for estimates of screening effectiveness.1-4 These studies met the criteria for level II evidence and constituted the best available evidence.5 The triplemarker screening in these studies included an ultrasonographic examination for confirmation of gestational age. Thus, the estimates cited in the task force recommendations reflect the screening intervention in toto (i.e., maternal serum markers with confirmation of gestational age), as delivered in a comprehensive screening program.

Farrell and associates suggest that the use of the term "triple-marker screening" is inaccurate. They correctly point out that screening with the use of maternal serum markers is evolving. However, they miss the point that triple-marker screening is the most effective combination to have undergone widespread evaluation in clinical trials. In keeping with the task force's emphasis on published evidence, the focus on triplemarker screening was deliberate and the use of the term accurate.

Farrell and associates believe that age should not be used as screening test. It is unclear, however, whether they are suggesting that there is no role for maternal age in counselling women concerning their options. It