units that will be effective in approximately 90% of the population, these dosage units are excessive for many young patients and may be inappropriate for frail elderly people. Most monographs in the *Compendium of Pharmaceuticals and Specialties* list the number of fixed-strength tablets or capsules that may be given in a 24-hour period. If an elderly 50-kg woman and a 100-kg man each consume one capsule they are certainly not getting the same dose. The presentation of dosage should include a measure of body weight or body surface area.

The Ontario Drug Benefit Formulary has taken on the role of paymaster for the pharmaceutical industry. Pharmacists are discouraged from finding creative ways to tailor medications to the specific needs of patients.

Recent advances in pharmacogenomics have produced much excitement concerning the future of personalized medicine. However, customized doses for elderly patients are needed today. The technology to deliver personalized medications is available now, but unfortunately it is seldom used by pharmacists or requested by physicians.

Véronique Koo

Pharmacy Manager Pharmacy.ca Toronto, Ont.

Reference

 Fischbach MS, Gold JL, Lee M, Dergal JM, Litner GM, Rochon PA. Pill-splitting in a longterm care facility. CMAJ 2001;164(6):785-6.

[Two of the authors respond:]

In an ideal world and using existing technology, every hospital, pharmacy and physician's office would be equipped with a database capable of computing suitable starting and maintenance doses for each patient's medications on the basis of the patient's age, body weight, surface area and creatinine clearance rate. The doses could subsequently be modified on the basis of therapeutic effect. This would allow physicians to prescribe and pharmacists to dispense essential therapies in a truly

personalized and standardized manner. Effective disease management would thereby be maximized and adverse events would be curtailed. Pharmacogenomics may promise even further advances, but its practical applications will likely not be implemented in the near future.

Until the pharmaceutical industry manufactures medicines in formulations that allow for such customized dosing (especially very small doses); until hospitals, pharmacies and physicians' offices invest in the infrastructure and information systems required to implement such an undertaking; and until hospital and provincial drug formularies and funding guidelines are revamped to account for variations in dosing, pill-splitting will remain an unfortunate reality.1 This is particularly true among community-dwelling and institutionalized elderly people who so often require the "start low, go slow" strategy.

Michelle Litner (née Fischbach)

Baycrest Centre for Geriatric Care Toronto, Ont.

Jennifer Gold

Baycrest Centre for Geriatric Care Toronto, Ont.

Reference

 Fischbach MS, Gold JL, Lee M, Dergal JM, Litner GM, Rochon PA. Pill-splitting in a longterm care facility. CMA7 2001;164(6):785-6.

Give us clear, not convoluted, clinical practice guidelines

The recent article on chemoprevention of breast cancer has left me a confused general practitioner. The authors refer to assessment of a woman's risk of breast cancer using the Gail index and make recommendations regarding the prescription of tamoxifen to women who have a Gail index that is greater than or equal to 1.66% over 5 years. But they point out that the Gail index has not been validated and has not been evaluated for use as a routine screening or case-finding instrument. Nowhere in the article can I find satis-

factory reconciliation of these conflicting notions.

Because the Gail index has not been evaluated and validated it does not seem to me that there are sufficient grounds for publication of a high-profile article setting out official guidelines for all Canadian physicians.

As a result of the publication of this article many patients will no doubt visit their physician's office to discuss chemoprevention of breast cancer with tamoxifen. When I am faced with such patients I will be at a loss as to how to proceed, not knowing whether the advice given in the article is valid or not.

Michael R. Lawrence

Physician and surgeon Vancouver, BC

Reference

 Levine M, Moutquin J-M, Walton R, Feightner J. Chemoprevention of breast cancer. CMAJ 2001;164(12):1681-90.

fter reading the guideline on **1** chemoprevention of breast cancer,¹ I feel compelled to vent my frustration at the publication of yet another verbose, convoluted and impractical guideline for those of us in clinical practice to follow. The appendix entitled "Questions and answers on chemoprevention and breast cancer: a guide for women and their physicians" also seems totally impractical. The woman and her physician are advised to obtain the Gail index from a Web site but told that it will only be useful in determining "whether to further discuss the benefits and harms of taking tamoxifen." A woman is supposed to decide whether she feels "a tamoxifen-induced stroke would be far worse than breast cancer" or "breast cancer would be far worse than a stroke." She is then advised, "You will have to determine the value you place on the possible consequences of taking or not taking tamoxifen after a full discussion with your doctor." Like so many other CMA7 guidelines, this provides little assistance in the decisionmaking process for the physician or the patient. Am I supposed to ask my patients if they would prefer to die of