I very much enjoyed CMA7’s 2002 Holiday Review. In particular, Carolyn Brown’s reporting of the IgNobel prizes caught my attention, especially the item about navel-gazing.1

This little report immediately took me back some 36 or 37 years, to my second year of premed studies at the University of Saskatchewan. Our biochemistry lab professor, whose office was just off the lab, had told us to disturb him whenever we had problems or questions. At that particular time, we were studying the hormones found in urine and, among other things, were told about hormone replacement therapy, including the fact that the brand name Premarin (conjugated estrogen) was a short version of “pregnant mare’s urine.”

I had a question about my lab work and proceeded to the professor’s office. He was very busy writing longhand on sheets of foolscap and did not look up for a minute or two. I asked him if he was writing up some research, and he said “Yes, indeed I am” and showed me the title on page 1. It read “On Fuzz-Gathering about the Umbilicus.” I don’t know if he ever completed this aspect of his “research,” but I wish now that I had taken more interest!

As I was about to leave his office after he responded to my query, he asked me to wait a moment and hurriedly penned a couple of lines on a sheet of paper, which he asked me to pass around the lab. If my memory serves me correctly, and I’m sure it does, the lines were as follows:

A permanently pregnant mare piddling perpetually produces more pee than an infinite series of mares peeing into pots periodically.

How I wish we had had more professors with his student rapport and sense of humour.

Joe Golumbia
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Reference

Osteoporosis in children: 2002 guidelines do not apply

The 2002 clinical practice guidelines for the diagnosis and management of osteoporosis in Canada1 cover both primary and secondary osteoporosis, but it is important to remember that these guidelines are based on evidence and experience with adult patients only and hence may not be applicable to younger patients.

Children and adolescents also experience fragility fractures, albeit rarely. In addition to their occurrence in association with genetic diseases (such as osteogenesis imperfecta), pediatric fragility fractures are seen in patients with immobilization (e.g., because of spinal cord injury), inflammatory diseases (e.g., juvenile idiopathic arthritis), glucocorticoid pharmacotherapy and combinations of these factors, sometimes with concomitant nutritional deficiencies of calcium and vitamin D; such fractures may also occur in patients with hypogonadism.

However, the World Health Organization densitometry categories1,4 cannot be applied in these cases, as T-scores for children calculated by standard methods are falsely low because there is no adjustment for their smaller size.4 Although T-scores should be neither computed nor reported for children, interpretation of pediatric densitometry results is possible if one has knowledge of various normal ranges for bone mass that depend on age, sex, bone size, pubertal tempo and pubertal stage. This process is analogous to analyzing children’s growth curves without knowing the parents’ heights.

Also currently lacking are data relating bone mass measurements to fracture risk in these special populations. As a result, it may be advisable to diagnose and consider pharmacotherapy for pediatric osteoporosis in the severe category — children who have already experienced a fragility fracture and who have identifiable risk factors. This definition is conservative but probably appropriate, given the lack of sufficient efficacy and safety data in children for the agents used for preventing fractures in older adults.

As Canadian child health programs develop recommendations for care for osteoporosis in children, it is hoped that diagnostic and clinical trials research will progress to the point that satisfying, evidence-based guidelines on the management of pediatric osteoporosis can one day be included.

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References