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Preventing neural tube defects in Europe: population based study

Araceli Busby, Lenore Abramsky, Helen Dolk, Ben Armstrong, a Eurocat Folic Acid Working Group

Each year, more than 4500 pregnancies in the European Union are affected by neural tube defects. Unambiguous evidence of the effectiveness of periconceptional folic acid in preventing neural tube defects has been available since 1991, and improving folic status sufficiently could result in the prevention of more than two thirds of all neural tube defects. We report on trends in the prevalence of neural tube defects up to 2001, in the context of a survey in 16 European countries of periconceptional folic acid policies and their implementation.

Participants, methods, and results

Eurocat is a network of population based congenital anomaly registries in Europe (www.eurocat.ulster.ac.uk). A total of 9273212 births were surveyed by 51 registries in 16 countries 1980-2001, including 8913 babies or fetuses with neural tube defects (anecephaly, spina bifida, or encephalocoele); 3298 live births, 844 stillbirths, and 4771 terminations of pregnancy after prenatal diagnosis. In the United Kingdom and Ireland, yearly prevalence of neural tube defects declined, predating any periconceptional folic acid supplementation policy initiatives, from 45 per 10 000 births in 1980 to 10 to 15 per 10 000 in the 1990s (figure). In contrast, in the rest of Europe the prevalence during the 1980s and thereafter was close to 10 per 10 000 births (figure).

The first governments to formulate a periconceptional folic acid supplementation policy were the United Kingdom (1992), Ireland (1993), and the Netherlands (1993); six more (Switzerland 1996, Denmark 1997, Norway 1998, Portugal 1998, France 2000, and Spain 2001) followed. Malta (1994) and Finland (1995) recommended raising folic status by dietary means only. Austria, Belgium, Croatia, Germany, and Italy had no government policy, although in all these countries professional bodies have issued guidelines or recommendations.

For 23 of the registers with data going back to at least 1990, we compared changes in prevalence over time according to policy type by estimating, for each registry, the ratio of prevalence in 1999-2001 to 1989-91 and calculating the mean of such ratios (on a log scale) for all registries in countries with similar policy types (as introduced by 1999), using standard random effects meta-analysis techniques. In the United Kingdom and Ireland we found a significant overall mean reduction in the prevalence of neural tube defects in 1999-2001 compared with 1989-1991 (32% (17% to 44%)), continuing a trend that preceded policies of folic acid supplementation. Registries in other countries with a policy of raising periconceptional folic status (by supplementation or dietary means) introduced by 1999 had a mean reduction in prevalence of 17% (44% reduction to 4% increase), whereas registries from countries with no policy by 1999 showed a mean reduction of 9% (26% reduction to 12% increase).

Comment

The prevalence of neural tube defects in Europe has not declined substantially in the past decade despite national policies of folic acid supplementation in half the countries. The prevalence in the United Kingdom and Ireland fell by 32% continuing a stronger pre-existing trend and remains slightly higher than overall European levels. These data suggest that policies simply recommending periconceptional supplementation of folic acid in planned pregnancies are not effective enough. Many women may not receive or respond to health promotion messages stressing the need to commence supplementation preconceptionally, or may remain unaware that dietary modifications are unlikely to achieve sufficient folate intake, and a large proportion of pregnancies in most countries are unplanned. Folate status of most women of childbearing age could be raised by fortifying a staple food with folic acid, which would also help to reduce socioeconomic inequalities in the prevalence of neural tube defects. Health effects of supplementation and fortification should be monitored, and policies should be reviewed periodically in light of the findings. The potential for preventing neural tube defects in Europe by raising folate status is still far from being fulfilled, and it is unacceptable to continue to rely mainly on prenatal screening and termination to reduce the number of babies born with neural tube defects.
The mystery of the milky effusion

A 59 year old man was admitted to our cardiac surgical intensive care unit after a third operation on the mitral and aortic valves. His previous operations had been six months before. All three surgeries were done via a median sternotomy, and the left pleural space was never opened during the operations. The postoperative recovery was initially uneventful. The day after surgery the patient was extubated, and the chest drains removed. The next day, however, we noticed a reduced air entry in the left lower zone of the chest, and a chest x ray showed some haziness in the left lower zone, consistent with a collection of fluid in the left pleural space. A diagnostic aspiration with a 14 gauge venous cannula in the seventh intercostal space in anterior axillary line produced a whitish fluid that looked uncannily like fresh milk.

Laboratory tests confirmed that the fluid was chyle, and the absence of vegetable particles ruled out stomach contents. Chylothorax due to excessive sternal retraction causing injury to the thoracic duct, though rare, is not unknown. The lab categorically labelled the aspirate as chyle and not as milk from the stomach. However, a vigilant attending nurse noted that the patient had drunk 250 ml of milk only 30 minutes before the aspiration. Meanwhile, the sample in the kidney tray showed curdling after two hours.

Because of persistent haziness on the chest x ray, we took a second aspirate at the sixth intercostal space the next day. This produced serosanguinous fluid rather than chyle, and we inserted an intercostal drain, which drained 600 ml in the first 24 hours. We removed the drain after 48 hours and ruefully concluded that the “chyle” of the first sample must have been due to transthoracic stomach aspiration.

We had all got carried away by the milky aspirate and had considered the exotic possibility of chylothorax from repeat surgeries and extensive retraction of the sternum. Even in a busy cardiac surgical intensive care unit that treats some 1200-1300 cases of open heart surgery a year, a simple situation can be made complex by incorrect interpretation of laboratory results. The milk didn’t curdle in the patient’s stomach immediately because of the high pH of his gastric juice after routine use of H2 blockers. The pathologist didn’t see any vegetable matter in the sample, but the patient had taken only clear fluids for the previous 24 hours. Aspiration was done with the patient in a semi-recumbent position (because of his postoperative status), with a full stomach (250 ml of milk), and during expiration. Clearly, the needle went through the chest wall, pleural space, diaphragm, and peritoneal space and reached the stomach. Some lessons can be learnt from this. Always try to do an aspiration with the patient in the proper position, bending forward against a table, in full inspiration, and, if possible, with an empty stomach. Investigations go hand in hand with clinical inputs and with the patient status. Milk and chyle can have similar physical characteristics and chemical composition, and interpretation can depend on the feedback by the clinician to the laboratory. Chylothorax is a debilitating and serious condition, whereas transthoracic stomach aspiration is a technical error that has no complications but can lead to a vexed clinical situation that can confuse even the most experienced clinician. This was a fantastic learning experience that has made us wiser and much more sober.

Rajnish Duara resident, department of cardiovascular and thoracic surgery, Sri Chitra Tirunal Institute for Medical Sciences and Technology, Thiruvananthapuram, India (rajnishduara@rediffmail.com)