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- 1 Connolly RM1, Carducci MA, Antonarakis ES. Use of androgen deprivation therapy in prostate cancer: Indications and prevalence. *Asian J Androl* 2012; **14**: 177–86.
- 2 Hearn JWD, AbuAli G, Reichard CA, et al. *HSD3B1* and resistance to androgen-deprivation therapy in prostate cancer: a retrospective, multicohort study. *Lancet Oncol* 2016; published online Aug 26. [http://dx.doi.org/10.1016/S1470-2045\(16\)30227-3](http://dx.doi.org/10.1016/S1470-2045(16)30227-3).
- 3 Li R, Evaul K, Sharma KK, et al. Abiraterone inhibits 3 β -hydroxysteroid dehydrogenase: A rationale for increasing drug exposure in castration-resistant prostate cancer. *Clin Cancer Res* 2012; **18**: 3571–79.
- 4 Chang KH, Li R, Kuri B, et al. A gain-of-function mutation in DHT synthesis in castration-resistant prostate cancer. *Cell* 2013; **154**: 1074–84.
- 5 Sweeney CJ, Chen YH, Carducci M, et al. Chemohormonal therapy in metastatic hormone-sensitive prostate cancer. *N Engl J Med* 2015; **373**: 737–46.
- 6 Ryan CJ, Smith MR, de Bono JS, et al. Abiraterone in metastatic prostate cancer without previous chemotherapy. *N Engl J Med* 2013; **368**: 138–48.
- 7 Reichard C, Almassi N, Russell C, et al. *HSD3B1* and resistance to CYP17A1 inhibition in prostate cancer (abstract MP07–04). *J Urol* 2016; **195** (suppl 4S): e77.
- 8 Li Z, Bishop AC, Alyamani M, et al. Conversion of abiraterone to D4A drives anti-tumour activity in prostate cancer. *Nature* 2015; **523**: 347–51.
- 9 Antonarakis ES, Lu C, Wang H, et al. AR-V7 and resistance to enzalutamide and abiraterone in prostate cancer. *N Engl J Med* 2014; **371**: 1028–38.
- 10 Mateo J, Carreira S, Sandhu S, et al. DNA-repair defects and olaparib in metastatic prostate cancer. *N Engl J Med* 2015; **373**: 1697–708.

Improving cervical cancer screening in Baltic, central, and eastern European countries



Cervical cancer is a disease with a high social and psychological burden, for which screening has shown efficacy and cost effectiveness.¹ However, implementation of cervical cancer screening has been proving difficult in several European countries (eg, Bulgaria and Romania), resulting in delayed adoption of effective treatment. After a 2010 European Parliament resolution called for the enactment of cancer prevention programmes (mainly cervical, breast, and colon cancer), some European Union member states set up screening programmes, while others began to implement organised cervical cancer screening.² Notably, organised screening is more effective than spontaneous screening in reducing cervical cancer incidence and mortality.³ However, even in some European countries with organised screening, coverage is inadequate. The incidence of cervical cancer is higher in women who have not been invited to screening or have not accepted the invitation to be screened.⁴ These considerations suggest a close connection between scientific evidence (ie, incidence and mortality rates) and the implementation of a sound screening programme.

In *The Lancet Oncology*, Salvatore Vaccarella and colleagues⁵ provide comprehensive information about cervical cancer incidence in six Baltic and central and eastern European countries (BCEE)—Estonia, Latvia, Lithuania, Belarus, Bulgaria, and Russia—as well as

projections for cervical cancer rates for 2017–40 in these countries. A previous article⁶ reported high cervical cancer incidence in BCEE states, without estimating future projections, and had suggested that preventive actions should be urgently implemented. The latest study by Vaccarella and colleagues therefore completes the information about cervical cancer in BCEE states, and its findings are interesting for many reasons. The authors used population-based registry data and an age-period-cohort model approach to develop cervical cancer incidence projections in two scenarios: no change (ie, continued absence of screening; scenario A) and introduction of screening from 2017 onwards (scenario B). Without effective screening programmes, cervical cancer incidence is expected to continue to rise in nearly all six BCEE countries studied, whereas the adoption of effective prevention and screening would achieve a gradual reduction of up to 50–60% in cervical cancer by 2040.⁵ In this age-period-cohort model, the period effect could be viewed as the result of the enactment of effective screening programmes that would reduce the cervical cancer incidence trend and the cohort effect as the action exerted by human papillomavirus (HPV) vaccination on the cohorts that would subsequently undergo screening. The period effect will also correlate with screening coverage.

These findings portray a dismal scenario, and should stimulate further consideration because they



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emphasise the need for urgent adoption of health policy actions and initiatives tailored to each country and area. Such policies should aim, on the one hand, to improve cervical cancer prevention and screening, and on the other, to manage the large number of cervical cancer cases that are expected in clinical practice. The findings of the study by Vaccarella and colleagues suggest the value of studying neighbouring countries such as Romania, which is characterised by high mortality and incidence rates and low screening coverage.¹ Additionally, all the countries reviewed by the authors have a high incidence of cervical cancer, largely opportunistic screening, low screening coverage, and use of the Pap smear as the primary test (except in the St Petersburg area of Russia, where the HPV test is in use). Screening in BCEE countries could be re-organised by replacing the Pap smear with the more sensitive HPV test,² thus creating a new era for cervical cancer screening in these countries. A further strategy to enhance participation is self-sampling, which has shown adequate accuracy in HPV testing.⁷ Indeed, according to a 2015 study,⁸ home mailing of a self-sampling kit proved effective in increasing screening participation, even in programmes that used the HPV test as the primary test. This approach would be especially useful for young women aged younger than 39 years, for whom screening is a high priority, but might also make testing more practical and acceptable to older women (aged older than 59 years). In addition to increasing participation directly, this approach might also ensure high rates of compliance in subsequent rounds of screening.

Although well designed, the study by Vaccarella and colleagues has some minor limitations, especially with regards to the data from Russia. These data must be considered with caution, both because they come from local registries (rather than the population-based cancer registries used for the other countries) and because the International Classification Disease, 9th revision, codes are not always accurate.⁶ Indeed, several deaths from uterine cancer are sometimes simply coded as tumour of the uterus, which provides no information about whether they were caused by cervical cancer or tumours of the body of the uterus.⁴ According to Mathers and colleagues,⁹ the quality level of data coming from Russia is D2 (ie, suboptimum). This low quality might result in

skewed projections and strategies. Nonetheless, the projections estimated by Vaccarella and colleagues provide a useful scenario that could be reassessed if higher quality data from Russia become available. The continuous improvement in the quality of epidemiological data collection provides crucial support to public health decision-makers. Russia is actually a high-income country¹⁰ and could therefore potentially afford to increase health-care spending and improve the quality of its epidemiological data.

In conclusion, cervical cancer screening programmes in BCEE countries should be based on existing European guidelines and harness the facilities, personnel, and economic structures available in each country and area. However—and crucially—this is not merely a health-related issue, because prioritisation of cancer prevention will also go a long way towards reducing social inequalities in these countries.

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I declare no competing interests.

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- 1 Ronco G, Dillner J, Elfström KM, et al. International HPV screening working group. Efficacy of HPV-based screening for prevention of invasive cervical cancer: follow-up of four European randomised controlled trials. *Lancet* 2014; **383**: 524–32.
- 2 Altobelli E, Lattanzi A. Cervical carcinoma in the European Union: an update on disease burden, screening program state of activation, and coverage as of March 2014. *Int J Gynecol Cancer* 2015; **25**: 474–83.
- 3 Arbyn M, Rebolj M, De Kok IM, et al. The challenges of organising cervical screening programmes in the 15 old member states of the European Union. *Eur J Cancer* 2009; **45**: 2671–78.
- 4 Zucchetto A, Ronco G, Giorgi Rossi P, et al. IMPATTO CERVIC Working Group. Screening patterns within organized programs and survival of Italian women with invasive cervical cancer. *Prev Med* 2013; **57**: 220–26.
- 5 Vaccarella S, Franceschi S, Zaridze D, et al. Preventable fractions of cervical cancer via effective screening in six Baltic, central, and eastern European countries 2017–40: a population-based study. *Lancet Oncol* 2016; published online Aug 22. [http://dx.doi.org/10.1016/S1470-2045\(16\)30275-3](http://dx.doi.org/10.1016/S1470-2045(16)30275-3).
- 6 Arbyn M, Antoine J, Magi M, et al. Trends in cervical cancer incidence and mortality in the Baltic countries, Bulgari and Romania. *Int J Cancer* 2011; **128**: 1899–907.
- 7 Arbyn M, Verdoodt F, Snijders PJ, et al. Accuracy of human papillomavirus testing on self-collected versus clinician-collected samples: a meta-analysis. *Lancet Oncol* 2014; **15**: 172–83.
- 8 Giorgi Rossi P, Fortunato C, Barbarino P, et al. HPV Self-sampling Italian Working Group. Self-sampling to increase participation in cervical cancer screening: an RCT comparing home mailing, distribution in pharmacies, and recall letter. *Br J Cancer* 2015; **112**: 667–75.
- 9 Mathers CD, Fat DM, Inoue M, Rao C, Lopez AD. Counting the dead and what they died from: an assessment of the global status of cause of death data. *Bull World Health Organ* 2005; **83**: 171–77.
- 10 The World Bank. Countries and economies. <http://data.worldbank.org/country> (accessed July 4, 2016).