Cervical cancer: incidence and survival in migrants within Spain

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Abstract

Study objective – This study examined the incidence of cervical cancer and survival rates according to migrant experience of women from different regions of Spain to Girona, Catalonia (Spain).

Design – Using data from the population based cancer registry of Girona for the period 1980–89, crude and age adjusted incidence rates were calculated for local-born and first generation migrants from other Spanish regions. The age standardised rate ratio (SRR) was calculated and Cox’s regression model was used to adjust survival according to migrant status for age and stage at diagnosis.

Main results – The incidence of cervical cancer was significantly higher in first generation Spanish migrants compared with locally born women (SRR: 2.02; 95% CI 1.40:2.92). The stage at diagnosis was more advanced among migrants. Survival probability was significantly associated with stage at diagnosis, but age and region of birth were not.

Conclusions – Migrants from the southern Spanish regions show a twofold excess in the incidence of cervical cancer compared with the Girona-born female population. Cases of cervical cancer in migrants are diagnosed at a more advanced stage and as a consequence have a poorer prognosis.

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Studies of the pattern of cancer incidence or mortality among migrant populations are a useful tool for exploring the aetiology of cancer, especially with regard to the possible role of genetic, lifestyle, and environmental factors.1 2 These studies have generally compared the occurrence of the disease in migrants from very different cultural or even ethnic backgrounds – for instance, Japanese migrants to the USA1 and Brazil,4 or Italian migrants in other countries.5 Different patterns of mortality or incidence are found in these studies in cancers of the stomach, breast, cervix, colon-rectum, and others. Few of these studies, however, have analysed differences in cancer risk in those who migrate within one country.6–9 Cervical cancer shows an important international variation in its incidence.10 Several risk factors have been related to this cancer. Some of the established ones are consistent with the presence of a sexually transmitted aetologic agent; in fact, in recent years, a strong association between human papilloma virus (HPV) and cervical cancer has been established.11 12 Other associated risk factors, such as the number of partners (both of the woman and of her partner(s)) and the age at first intercourse, could be surrogates of HPV infection. Oral contraceptives, smoking, and some dietary factors are also risk factors that have been associated with cervical cancer.

Spain has one of the lowest incidences of cervical cancer, as observed by the country’s various cancer registries.13 14 Intense internal migration has traditionally occurred between regions of Spain. The region of Catalonia and the city of Madrid have been the recipients of most of this migration throughout this century. In fact, 34.1% of people living in Catalonia today were born in another region of Spain.15 This study aimed to examine the incidence and survival associated with cervical cancer in first generation migrants to Girona (Catalonia, Spain) from other Spanish regions using data from a population based cancer registry.

Methods

Data from the population based cancer registry of Girona covering the period 1980–89 were used. Girona is a province of Catalonia in north eastern Spain, with a female population of 246 515 in 1986, of whom 24% were born outside the province. Of the female population, 68.3% were born in Girona, 7.7% in other provinces of Catalonia, 15.2% in Andalusia-Extremadura (southern regions of Spain), 6.6% in other regions of Spain, and 2.2% abroad.16

The population based cancer registry of Girona is restricted to cancer of the breast, uterus, ovary, and other genitals. It actively collects information on all incident cases among the residents in the province of Girona. Quality indices regarding cervical cancer were as follows: 98.3% of histological verification and 1% of cases were identified by death certificate only. Cancer cases are classified according to the ICD-O (9th revision). The cancer registry provided information on the patient’s age, tumour site, stage of the disease (divided into local, regional and disseminated), and follow up status. The registry includes cases since January 1980: all cases were followed up actively and passively (through death certificates) until December 1991. Cancers in situ were included in the registry, but only presented for descriptive purposes. Place of birth is routinely collected, and has been used to identify first generation migrants. The year of migration is not available.

Rates were initially calculated by grouping the cases into two periods 1980–84 and 1985–
89. The results were highly consistent, and results for the entire period (1980–89) are therefore presented. Age adjustment of the rates was performed by the direct method and using the world standard population. Migrants were classified according to region of birth as follows: born in Girona and in other parts of Catalonia, born in the south of Spain (including Andalusia and Extremadura), born in other regions of Spain, and born in other countries.

Comparison of the age standardised incidence rates was done using the standardised rate ratio (SRR) and 95% confidence interval (95% CI).¹⁷ SRR is the ratio between two directly age adjusted rates and is equivalent to the relative risk of disease in migrant population compared to local born population.

It is possible to calculate the statistical significance of the SRR as an indicator of whether the observed ratio is significantly different from unity.

Differences in the stage at diagnosis according to region of birth were analysed by the \( \chi^2 \) statistic. Survival was analysed first by region of birth, using the Kaplan-Meier estimate. To compare the probabilities of survival according to the region of birth, the log rank test was used. To adjust for differences in the stage at diagnosis or age on the relationship between region of birth and survival, Cox's proportional hazard model was used.¹⁸ The following variables were included in the model: age at diagnosis, region of birth, and stage at diagnosis. Age was divided into the following groups: less than 34, 34–45, 45–54, 55–64, and 65 and over. The reference categories used for the analysis were as follows: age group was less than 35 years, stage at diagnosis local, and region of birth Catalonia. Hazards proportionality assumption of Cox's model was checked through including the log survival function against time in the model. Computations were performed using \( \text{EGRET} \) software.¹⁹

### Results

There were 180 women diagnosed with invasive cervical cancer in Girona, Catalonia, during the period 1980–89. The crude annual incidence rate is 8.19 per 100 000 women and the age adjusted rate is 6.21 per 100 000 women. The region of birth was unknown in 10 cases, so 170 cases only are included in the study. Crude, age specific, and age adjusted incidence rates are shown in table 1 according to the region of birth for the period 1980–89. The age adjusted rates in women born in southern regions of Spain and the rest of Spain are approximately twice those of the Girona born women in the study period. The age adjusted incidence rate of women born abroad is the highest, probably as a result of resubregistration in the denominator. Part of the at risk group that was born outside of Spain is still included in the census of their country of origin. This group was consequently excluded from the rest of the analyses. The age specific incidence rates by region of birth (table 1) show a consistently higher incidence rate in migrants of age groups of 45 and older compared with the rates in Catalan born women. In addition, the rate for women born in the rest of Spain and abroad shows a higher rate even in younger age groups, but few cases are involved.

The SRR comparing incidence rates of Spanish migrants versus those born in Girona was 2.02 (95% CI 1.40, 2.92). The SRR was higher in women from southern regions of Spain versus those born in Girona than in other Spanish regions versus Girona (table 2).

The stage distribution at the time of diagnosis, including cervical cancer in situ, is shown in table 3 in relation to the region of birth.
Migrants tend to have a higher proportion of women diagnosed at a later stage than women born in Girona and conversely, the proportion of cancer diagnosed at the "in situ" stage is significantly lower among migrant women.

The probability of survival was significantly related to the stage at diagnosis (p<0.0001) but neither the age (p=0.635) nor the region of birth (p=0.808) were significantly related to the probability of survival in univariate analysis (table 4). Because of the relationship between the stage at diagnosis and region of birth, a Cox's regression analysis was performed to compare survival between migrants and local born women, controlling for stage at diagnosis and age. Taking into account the age distribution and stage at diagnosis, the probability of survival was not significantly different in those born in Girona compared with women born in the southern regions or in the rest of Spain (table 5).

Discussion
A higher risk of developing invasive cervical cancer has been observed in women who migrated from several regions of Spain to Girona. In addition, this increased incidence was shown to be accompanied by a more advanced stage at diagnosis for migrants compared with women born in Girona. Survival was not significantly different according to the region of birth.

Some potential sources of bias should be taken into account when analysing these data. Some women might have migrated to search for medical care or social support after diagnosis, especially mothers or other family members of people who had previously migrated. These women might have become residents only for purposes of treatment. This bias could affect the population denominator because these women may not have been included in the census. It is unlikely, however, that, once included, this very small number of cases in the population denominator could influence the incidence rates in a specific direction. Migration to Girona for search of treatment should be of little relevance if we take into account the proximity (100 km) of Barcelona, where there are five university hospitals and where treatment could be obtained more easily than in Girona.

Ascertainment of cases through histological verification and coverage of the registry with regard to cervical cancer are comparable with other registries.10 It should be noted that in 8.6% of cases it was not possible to ascertain the region of birth. The true distribution of place of birth in cases with an unknown place of birth tends to over-represent migrant patients, who are more likely to escape information retrieval procedures; in this case our findings of a twofold higher incidence among migrants should be viewed as an underestimation of the true incidence excess. If, in contrast, the cases of unknown place of birth were all local born, the analyses presented in tables 1 and 2 would still show a statistically significant excess among migrants. In our study, it was not possible to control for the length of stay in Girona or the age at migration. The former is an important variable in migrant studies,22,23 the latter is relevant in cervical cancer because of the established association with age at first intercourse and number of sexual partners, two variables likely to differ in relation to social class and place of residence.

The higher incidence of invasive cervical cancer among migrant women in Girona could be accounted for by two complementary factors: differential exposure to risk factors for cervical cancer and differential participation rate in screening activities. With regard to the first factor, the higher incidence could be partly explained by low socioeconomic status or lifestyles specific to migrant populations. Migrants are usually young people, and this population has shown different nuptial and fecundity rates but these rates are not independent of age in our country,22,23 so adjusting by age could account for this difference. Cervical cancer has been related to lower socioeconomic status in different countries24 as well as in Spain.25 Migrants to Catalonia tend to have a lower socioeconomic level than the local born population;22 for example, the percentage of residents in Girona who have not completed primary school is 36.6% among Catalan born women and 43.1% among people not born in Girona.28 From this point of view, migration could be considered a surrogate for socioeconomic level, especially in first generation migrants. It has been suggested that social class could have an effect on the risk of cervical cancer independent of sexual behaviour,25 in fact the relationship between sexual behaviour and social class is unclear.29 Our results suggest that internal migration could be related to a different risk of cervical cancer, but the relationship with sexual behaviour is unknown. It is probable that socioeconomic differences are related to an increased risk of cervical cancer.
The second suggested explanation for these results is the lower rate of participation in cervical cancer screening. Screening for cervical cancer effectively reduces the incidence among those who participate for at least 10 years after a negative test. Screening is not centrally organised in Catalonia, it is a byproduct of gynaecological health care, largely private, and in many cases supported by social security. Data from a survey undertaken in the municipality of Girona and its surroundings indicate that the cervical cytology rate of migrants is lower than that of local born women (30%-7% and 23%-4% respectively) and a strong relationship between educational level and self reported cervical screening was observed. Other authors have suggested that women from lower socioeconomic groups are less likely to have a screening test. This is partly supported by the significant difference in the proportions of cancers “in situ” found in relation to the region of birth (table 3). In fact, cancer in situ is largely a surrogate for screening practices. The possibility of uncompleted registration of such cancers makes any conclusion regarding these data provisional. The length of stay of the migrant population, a variable not available in our study, has been related to the observed reduction in the risk of invasive cancer in Israel. Where the trend also differed in relation to the place of birth of the migrants. The suggested explanation for this observation lies on the cumulative exposure to cytological screening in migrants after arriving in Israel. In conclusion, cytological screening could play a major role in explaining the difference in the incidence of invasive cervical cancer among migrants to Girona.

In this context, it is remarkable that differences in the cervical cancer incidence in relation to the region of birth do not suggest a difference in prognosis. This observation could indicate that the main problem of equity in health care detected in our study is one of accessibility to screening and not of the quality of treatment. This finding is consistent with other authors who have found little effect of socioeconomic differences on survival for this type of cancer, and who pointed to equity of access to treatment as the main explanation for this finding.

Migrant studies are a useful tool for describing the epidemiology of a disease in a geographical area. Girona, like other regions of Spain, is an area with a low incidence of cervical cancer and one in which there is a twofold higher incidence rate in migrants than in native born women. The cancer stage at diagnosis is more advanced in migrants but survival is not significantly different in relation to the region of birth. The main reason for the higher incidence seems to lie in limited access to screening among migrants. It is possible that the differences observed in this first generation of migrants will disappear with time and in second generation migrants. Further research is needed to interpret fully these results. In particular, reasons for differences in the distribution of stage at diagnosis and access to screening practices should be addressed.
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