Analysis of spatial-temporal clusters of childhood cancer incidence in the province of Córdoba, Argentina (2004-2013)

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ABSTRACT

Introduction. In spite of its low incidence, childhood cancer is becoming increasingly more relevant in Argentina. More advances have been made in cancer treatment than in the study of its etiology or determining factors. There are no investigations that analyze its spatial and temporal distribution or potential clustering.

Objective. To perform exploratory spatial and temporal analyses based on the database of the Registry of Tumors of the Province of Córdoba (2004-2013) to determine the clustering of childhood cancer incidence in Córdoba (Argentina).

Populations and methods. Epidemiological, retrospective, ecological study. Data from 1098 patients with malignancies aged 0-14 years old from the Registry of Tumors of the Province of Córdoba (2004-2013) were used. A geographic information system model was developed. The presence of spatial, temporal, and spatial-temporal clusters was analyzed in the districts of Córdoba using the SaTScan software.

Results. Spatial clusters were detected, with a high number of cases, for total tumors ($p=0.01$), leukemias ($p=0.02$), malignant neoplasms of lymphoid, hematopoietic and related tissue ($p=0.03$), central nervous system tumors ($p=0.03$), and a high level of indicators of risk for renal tumors ($p=0.01$). In addition, a temporal cluster ($p=0.01$) and a spatial-temporal cluster ($p=0.02$) for neuroblastoma and other peripheral nervous cell tumors were also observed.

Conclusions. Significant clusters were determined, with important associated indicators observed in several districts of Córdoba. This is the first methodological step towards the development of new investigations on the risk factors for childhood cancer and its etiology.

Key words: cancer, children, cluster analysis, incidence, geographic information systems.

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INTRODUCTION

Advances made in pediatric health care have led to an epidemiological transition in the 0-14 year old group in Argentina. Thus, in the face of a reduction in the relative burden of infectious diseases and malnutrition, the importance of childhood cancer is becoming increasingly more evident, in spite of its low incidence. 1,2 Childhood cancer refers to a group of diseases of multifactorial origin, for which more advances have been made in terms of treatment than in the study of their etiology or determining factors. 3-7 Although the origin or influence factors of these diseases consider both genetic and environmental factors (and their interaction), many investigations suggest that the environment carries far more weight. 4,5,8-10 Environmental factors refer to anything in interaction with the child, that is not related to a hereditary trait, and that may influence the development of cancer in the pediatric or adult stage. 4,5 Given such complex scenario, i.e. the multifactorial origin of cancer and the lack of knowledge on its etiology or associated factors in children, it is critical to have information provided by official registries based on analytical tools that enable the overlapping and interaction of various data in a cohesive and objective manner in the search for new forms of data stratification and visualization. 11 Geographic information systems (GIS), together with cluster analysis tools, establish the first approach to delimiting research on specific population groups and/or areas and make it effective, helping to the future knowledge of childhood cancer etiology. 7,11

The objective of this research project is to perform exploratory spatial-temporal analyses based on the database of the Registry of Tumors of the Province of Córdoba (2004-2013) to determine the clustering of childhood cancer incidence in Córdoba (Argentina).
POPPULATION AND METHODS

This was an epidemiological, retrospective, ecological study. The study region is made up of the 26 districts of the province of Córdoba (spatial analysis units). As per the National Census of 2010, the province has 3,304,825 inhabitants, and 805,512 of them are younger than 15 years of age.12

Data were provided by the Registry of Tumors of the Province of Córdoba, which depends on the Provincial Ministry of Health. It is a hospital-based registry arranged as a network of reporting sites.13

For the proposed study, data from 1098 patients with tumors aged 0-14 years old, corresponding to the 2004-2013 period, were used; these patients had a registered address in the province of Córdoba. The database was classified as per the International Classification of Diseases, 10th revision (ICD-10), and the International Classification of Childhood Cancer, 3rd edition (ICCC-3), thus allowing data grouping. In order to compare these data with those of other registries, frequency indexes, crude rates, and standardized rates were estimated for the overall group and for childhood cancer subgroups (standardized rates were estimated using the direct method of the “old standard world population” defined by Doll and Waterhouse and recommended by the World Health Organization).14

Once data were classified, a geo-referenced database was developed using GIS for the representation of data and the development of new data for the subsequent cluster analyses. Polygonal layer files were used to represent the province and the different districts, provided by the Geoportal of the government of the province of Córdoba.15

Spatial, temporal, and spatial-temporal analyses of childhood cancer were done using the SaTScan software, which determines the presence of spatial and temporal clusters.16

SaTScan is a three-dimensional analytical tool based on the geographic and temporal location of cases. This is an exploratory methodology, and its purpose is to detect and locate clusters with a higher frequency of cancer occurrence than that expected for the studied region. The software analyzes “spaces” or “windows” shaped like a cylinder, with a centroid and definite radius. The general software procedure consists in analyzing the risk for the occurrence of an event of interest within each “window” and comparing it to the rest of the studied territory. This methodology is based on the assumption that cases have a Poisson-like distribution, which is the probability distribution used to analyze discrete outcome measures that represent rare events.16 The tested null hypothesis is that the risk for the epidemiological event of interest remains constant over space and time, i.e., there is no clustering. The alternative hypothesis proposes that, for at least one of the cylinders, the risk is different inside and outside it, i.e., such cylinder is a cluster in itself. In addition, its statistical significance is verified and a relative risk of disease is attributed in relation to the areas not included in the cluster.

The software requires the following data to perform the analyses: geographic location of each analysis unit (in our study, district centroids), number of cases per temporal analysis unit (number of childhood cancer cases in children aged 0-14 years old per district, per year), the population at risk per unit, per year (total number of children aged 0-14 years old per district, per year). To establish the population at risk per year of analysis, population data provided by the National Censuses of Population and Housing of 2001 and 201012,17 were used. These data were extrapolated using the AGEINT template, which is part of the Population Analysis Spreadsheets developed by the U.S. Census Bureau.18

Analyses were done considering the total tumors group (total tumors) and subgroups included in the database as per the two classifications mentioned above. In addition, age and sex were introduced as covariates to assess whether adding data shaped or influenced clusters.

RESULTS

The Registry of Tumors of the Province of Córdoba provided data on 1098 cases of children aged 0-14 years old with malignant tumors diagnosed in Córdoba between 2004 and 2013. Cases were classified based on two international nomenclatures.

Table 1 shows the number of cases per year and the annual crude incidence rate of the different cancer groups classified as per the ICCC-3. The indicators from this table were compared to those of the Argentine Hospital Oncopediatric Registry; some differences were observed that may not be statistically verified, and that we believe are rather related to data management than to their origin.19

Table 2 shows the frequency of cases per year, the crude rate and the standardized rate of childhood cancer in Córdoba by district (2004-2013). It is worth noting the low frequency of
childhood cancer in some districts, which is even null for some or several years (e.g., no cases were observed in the Pocho district over the entire study period). In addition, the size of the child population in some districts is small, which makes estimated rates fluctuate due to data variability among and within geographic units in different years.

Table 3 and Figure 1 describe each spatial cluster and associated parameters observed in the province of Córdoba over the studied period. The introduction of unclassified data led to the detection of a significant spatial cluster \( (p = 0.01) \) for total tumors in the Capital and Colón districts. The following results were obtained using the data classified as per the two international nomenclatures: a significant spatial cluster for leukemias \( (p = 0.02) \) in the Capital, Río Primero, Río Segundo, and Tercero Arriba districts; a significant spatial cluster for renal tumors \( (p = 0.01) \) in the Cruz del Eje, Minas, Pocho, Punilla, San Alberto, San Javier, and Santa María districts; a significant spatial cluster for central nervous system tumors \( (p = 0.03) \) in the Capital and Colón districts; and a significant spatial cluster for primary or presumably primary malignant neoplasms of lymphoid, hematopoietic and related tissue \( (p = 0.03) \) in the Capital, Río Primero, Río Segundo, and Tercero Arriba districts (consistent, in terms of geographic distribution, with the leukemia cluster shown in Figure 1).

The spatial-temporal analysis detected a significant cluster for neuroblastoma and other peripheral nervous cell tumors \( (p = 0.02) \) in the Capital, Colón, and Santa María districts in the 2009-2010 period (Figure 2). The software used a crude rate of 9 cases per one million children aged 0-14 years old as a global parameter. Twenty cases were observed within this cluster, although as per the software estimations, 7.5 cases were expected (a 168% difference). The crude rate estimated for this cluster was 25 cases per one million children aged 0-14 years old, with a relative risk of 3.29.

Finally, the temporal analysis showed only one significant result \( (p = 0.01) \) for the neuroblastoma and other peripheral nervous cell tumors group in the 2009-2010 period. The software used a crude rate of 9 cases per one million children aged 0-14 years old as a global parameter; 27 cases were found, although as per the software estimations,

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ICCC: International Classification of Childhood Cancer.

* Average incidence in children younger than 15 years old per 1 000 000 children.

Source: prepared by the author based on data provided by the Registry of Tumors of the Province of Córdoba and the National Censuses of Population and Housing of 2001 and 2010.12,15,17
### Table 2. Frequency of cases per year, crude rate and standardized rate by district. Province of Córdoba, Argentina, 2004-2013 period

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<td>135.2</td>
</tr>
<tr>
<td>Total</td>
<td>807,377</td>
<td>109</td>
<td>114</td>
<td>108</td>
<td>104</td>
<td>115</td>
<td>107</td>
<td>132</td>
<td>110</td>
<td>106</td>
<td>93</td>
<td>1098</td>
<td>136</td>
<td>139.4</td>
</tr>
</tbody>
</table>

* Average population of children aged 0-14 years old for the 2004-2013 period (source: National Censuses of 2001 and 2010).12,17
** Annual crude rate per one million children aged 0-14 years old.
*** Standardized rate using the direct method of the “old standard world population” defined by Doll and Waterhouse and recommended by the World Health Organization.14
Source: prepared by the author based on data provided by the Registry of Tumors of the Province of Córdoba and the National Censuses of Population and Housing of 2001 and 2010.12,13,17

### Table 3. Significant spatial clusters of childhood cancer incidence observed in the province of Córdoba, Argentina, 2004-2013 period

<table>
<thead>
<tr>
<th>Group defining the cluster</th>
<th>Districts included in the cluster</th>
<th>Observed cases*</th>
<th>Expected cases**</th>
<th>Obs. cases/ exp. cases</th>
<th>Relative risk</th>
<th>Crude incidence rate***</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total tumors</td>
<td>Capital and Colón</td>
<td>572</td>
<td>513.7</td>
<td>1.11</td>
<td>1.24</td>
<td>151</td>
</tr>
<tr>
<td>Leukemias (ICCC I)</td>
<td>Capital, Río Primero, Río Segundo and Tercero Arriba</td>
<td>188</td>
<td>157.7</td>
<td>1.19</td>
<td>1.44</td>
<td>49</td>
</tr>
<tr>
<td>Renal tumors (ICCC VI)</td>
<td>Cruz del Eje, Minas, Pocho, Punilla, San Alberto, San Javier and Santa María</td>
<td>18</td>
<td>7.6</td>
<td>2.35</td>
<td>2.99</td>
<td>16</td>
</tr>
<tr>
<td>Tumors of the central nervous system (ICD-10 C69-C72)</td>
<td>Capital and Colón, Río Segundo and Tercero Arriba</td>
<td>262</td>
<td>226.6</td>
<td>1.16</td>
<td>1.34</td>
<td>69</td>
</tr>
</tbody>
</table>

ICCC: International Classification of Childhood Cancer; ICD: International Classification of Diseases.
* Cases observed within the cluster as per the database.
** Cases expected within the cluster as per the SaTScan software estimation.
*** Per 1 000 000 inhabitants, within the cluster as estimated by the SaTScan software.
**** Primary or presumably primary malignant neoplasms of lymphoid, hematopoietic and related tissue (ICD-10 C81-C96).
Source: prepared by the author based on data provided by the Registry of Tumors of the Province of Córdoba and the National Censuses of Population and Housing of 2001 and 2010.12,13,17
Figure 1. Maps of some significant spatial clusters of childhood cancer incidence in the province of Córdoba, Argentina, 2004-2013 period

Source: estimated based on data provided by the Registry of Tumors of the Province of Córdoba (2004-2013) and the National Censuses of Population and Housing of 2001 and 2010.12,13,17
14.9 cases were expected (an 80% difference), with a relative risk of 2.26.

The introduction of the age and sex covariates into the software provided the same results as without them. This suggests that such covariates do not shape or influence data aggregation into clusters.

**DISCUSSION**

We believe our research provides information on the geographic and temporal distribution of childhood cancer in the province of Córdoba as a first methodological step to advance in the epidemiological study of this type of cancer. This will allow for further, more specific research aimed at studying the risk factors of childhood cancer and its etiology. One of the disadvantages of working with registries of childhood cancer is its low incidence, which results in methodological challenges in terms of data consistency. Although we were able to work with data corresponding to a ten-year period provided by the Registry of Tumors of the Province of Córdoba, it is important to continue corroborating results in the following years considering that the registry is rather recent and is under constant review. In the future, the plan involves working with data from children and youth aged 0-19 years old to widen case reporting and include an age group whose characteristics is more related to child and adolescent development than to the adult stage.

The cluster-associated indicators observed in our research stress the importance of further deepening analyses in such geographic areas. In addition, it is worth noting the following characteristics of some of the clusters observed in our study: the spatial cluster for total tumors is especially important because it covers 52% of the database (572 out of 1098 cases), and is concentrated in two out of the 26 provincial districts (Capital and Colón), the place of residence of 47% of the child population. The spatial cluster for leukemias (ICCC I) and for primary or presumably primary malignant neoplasms of lymphoid, hematopoietic and related tissue (ICD-10, C81-C96) also stands out because of the large child population in the region included in the cluster and, at the same time, because it encompasses the largest childhood cancer group (leukemias). The spatial cluster for renal tumors (ICCC VI) is particularly striking because it includes a type of cancer that is uncommon in children but evidences very high associated indicators (observed/expected ratio, relative risk, and rate within the cluster). In addition, it is located in a very definite and distinct geographic area compared to the rest of the clusters that show geographic overlapping. Finally, the analysis of results from the spatial-temporal cluster for neuroblastoma and other peripheral nervous cell tumors (ICCC IV) emphasizes the specificity of the analysis to detect the cluster in two particular years across the period, with high associated indicators. The result of the temporal cluster for the same group indicates that the incidence peaked across the province in the 2009-2010 period. This finding is probably the result of including the incidence of the three districts (Capital, Colón, and Santa María) in the total for the province for this period.

Although there are other studies conducted in Córdoba that analyze geographic distribution patterns, they do not always include data...
on incidence. In addition, those investigations were done in a global age group or in adults only. A separate analysis is critical in the case of childhood cancer because its dynamics is absolutely different from that of adults.3,7

The lack of studies on childhood cancer conducted in Argentina using the same methodology or tools as in this study may be considered a disadvantage because it is not possible to compare results that would certainly improve the approach to this problem. In turn, it is critical to create interdisciplinary research and working groups to establish monitoring, research and prevention systems targeted at childhood cancer and its risk factors in Argentina.

At an international level, there are several studies that applied different methodologies to analyze the spatial-temporal distribution of childhood cancer to establish clusters. Although they cannot be directly compared, those studies found, as in our study, clusters within the large groups of total tumors, leukemias, tumors of the central nervous system, and lymphomas.7,24-31 However, those studies failed to establish a conclusive relationship between results and a socioeconomic factor, even though the presence of clusters is probably an evidence of the fact that environmental factors are related to the presence and development of cancer.32-34 Apart from ionizing radiation and some congenital genetic syndromes, there is little evidence —and lots of suspicion— regarding other factors that may influence the clustering of childhood cancer. For example, unidentified infectious agents,29,30 agrochemicals and pesticides,29,30 pollutants in the air or the water,34,35 polluting industrial products or waste (due to parental occupational exposure) or direct environmental exposure.29,32,34

It is worth noting that most of those studies, as our study, highlight the power of the methodologies that assess spatial-temporal patterns as statistical tools that make it possible to focus on areas or populations where the rates of childhood cancer are unusually high and, as a result, to study its risk factors and etiology.7,25,28,31 There are several disease spatial-temporal analysis software programs, but SaTScan stands out because it is designed for rare statistical events (and uses case individual count in its analyses). This feature makes it optimal for the study of childhood cancer, which is a very rare disease. Among cluster-associated indicators, the relative risk estimated by SaTScan is specific to this software, so it makes it possible to know the risk for an event (presence of disease) within the cluster compared to the rest of the studied region. It also enables the introduction of covariates, which makes it possible to test factors that may influence their distribution. Finally, an approach using other statistical software methodologies would be more difficult because they use data summarized into similar indicators or rates that assume a loss of resources due to the small size of populations.7

There are few studies conducted in Argentina that analyze the spatial-temporal distribution of cancer, let alone in children. Results obtained in our study open the way for new, future research hypotheses, with the advantage of limiting to particular geographic areas and thus start understanding more of the epidemiological dynamics of childhood cancer. In addition, this study has demonstrated the usefulness of a tool that may be extrapolated to other age groups and used in different geographic scales to monitor population health.

CONCLUSIONS

Significant clusters were determined, with important associated indicators observed in several districts of Córdoba. This is the first methodological step towards the development of new investigations on the risk factors for childhood cancer and its etiology.

Acknowledgments

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REFERENCES


