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# Cancer incidence and mortality in children in the Mexican Social Security Institute (1996-2013)

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## Abstract

**Objective.** To identify the cancer incidence and mortality in Mexican Social Security Institute beneficiary (MSSI-B) children during 1996-2013. **Materials and methods.** Both cancer cases (n=4 728) and deaths (n=2 378) were analyzed in MSSI-B children who were registered in five states of the Mexican Republic. The incidence and mortality trends and the incidences (rate x 1 000 000 children / year) of the type of cancer, age, sex, and place of residence were obtained. **Results.** For both indicators (incidence and mortality), there was a downward trend for the period of 1996-2001 and a stable trend for 2002-2013. This occurred in the overall mortality and incidence trends of the Estado de México and Chiapas, and in the leukemia and the acute lymphoid subgroups. The annual overall incidence was 128 cases per 1 000 000 children. Leukemia, lymphomas, and central nervous system tumors were the principal cancer groups. **Conclusions.** Cancer mortality for the period of 2002-2013 did not diminish. Interinstitutional and/or international research should be designed to improve the care of these children.

**Keywords:** cancer; children; trends; incidence; mortality; Mexico

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## Resumen

**Objetivo.** Mostrar la incidencia y mortalidad por cáncer en niños derechohabientes del Instituto Mexicano del Seguro Social (DH-IMSS) (periodo 1996-2013). **Material y métodos.** Se analizaron los casos (n=4 728) y las muertes (n=2 378) por cáncer en niños DH-IMSS registrados en cinco estados de la República mexicana. Se obtuvo la tendencia de la incidencia y mortalidad y la incidencia por tipo de cáncer, edad, sexo y lugar de residencia. **Resultados.** Para ambos indicadores (incidencia y mortalidad) hubo tendencia a la disminución para el periodo 1996-2001 y estabilidad para 2002-2013; esto sucedió para la tendencia global, para el Estado de México y Chiapas, para las leucemias y para el subgrupo de las linfoides agudas. La incidencia global fue de 128 x 1 000 000 niños/año; las leucemias, tumores del sistema nervioso central y linfomas fueron los principales grupos de cáncer. **Conclusiones.** La mortalidad por cáncer para el periodo 2002-2013 no ha disminuido; deben diseñarse investigaciones interinstitucionales e internacionales para mejorar la atención médica de estos niños.

**Palabras clave:** cáncer; niños; tendencias; incidencia; mortalidad; México

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To understand the effect that cancer has had on children (0-14 years old), it is necessary to study both the incidence and the mortality from these causes.<sup>1</sup>

The difficulty in estimating the incidence of cancer lies in the necessity of having a population-based registry of cancer cases, which means recording all cases that occur in a given population and, most importantly, collecting data with quality criteria (indicating that the data should be 1) complete, 2) valid, 3) timely, and 4) following international standards for the coding and comparison of information).<sup>2</sup> These criteria are necessary so that the data will be reliable. Moreover, a registry of a few well-recorded cases is preferred to a registry of many poorly-recorded cases.<sup>3</sup> Accordingly, the incidence of cancer in children in developed countries is well-known.<sup>1</sup>

In reference to the study of cancer mortality in both developed and developing countries, there are well-established policies for building an adequate registry of cancer deaths. Therefore, cancer mortality in children is well-known in most countries worldwide.<sup>4,5</sup>

The interpretation of the mentioned indicators is important because the incidence is related to the risk that a population has of developing a disease.<sup>1,3</sup> For its part, mortality is a fairly accurate reflection of the overall treatment being provided to children with cancer, particularly when the disease incidence is stable.<sup>6,7</sup>

Globally, the overall incidence of cancer in children ranges between 100 and 180 cases per 1 000 000 children / year. Incidences below this range reflect a possible underestimation, and incidences above represent a possible overestimation.<sup>8</sup>

In countries where the survival of patients has increased, the mortality has subsequently decreased; this situation has occurred in developed countries. Furthermore, the mortality rate ranges between 22 and 30 cases per 1 000 000 children / year in these countries.<sup>4,7</sup>

At the *Instituto Mexicano del Seguro Social* (IMSS) [Mexican Social Security Institute (MSSI)], which is located in the *Distrito Federal* (DF) [Federal District, coterminous with Mexico City], a population-based Registry of Cancer in Children (RCC) was initiated in 1996 and has attempted to comply with the aforementioned quality criteria. The Registry contains information regarding the cancer epidemiology of MSSI beneficiary (MSSI-B) children, but only children in some states of the Mexican Republic have been included.<sup>9</sup>

The objective of this study was to analyze the data for both cancer incidence and mortality in

MSSI-B children who were residents of the DF, Estado de México (EM), Morelos (Mor), Guerrero (Gro), and Chiapas (Chis) for the period of 1996-2013 and to discern the tendency that each variable presented.

## Materials and methods

An observational, prospective, descriptive study was conducted. Data from two population registries were analyzed for the period from January 1<sup>st</sup>, 1996 to December 31<sup>st</sup>, 2013.

Data were obtained from two different sources of information.

- a) *Incidence data.* The information for the study of incidence was obtained from the RCC, which is managed at the Clinical Epidemiology Medical Research Unit of the Pediatric Hospital in the National Medical Center XXI Century. It is a population-based registry of cancer cases. The information is collected actively and prospectively by two registrars (nurses) who were trained for the collection, coding (morphology, topography, and staging of cases), and data capture of the different study variables. The coding and staging of the cases are performed in conjunction with the head of the registry (AFG), with a good consistency in the coding of these variables (unweighted Kappa of 0.85).

To code the different cancer cases, topographic and morphologic codes are used. For cases collected from 1996-1999, the second edition of the International Classification of Diseases for Oncology (ICD-O-2) was used, and for cases collected between 2000 and 2013, the third edition (ICD-O-3) was used. The cases were grouped according to the International Classification for Childhood Cancer (ICCC).<sup>10</sup>

This study used data on children with cancer who were treated in the DF at the two hospitals that the MSSI uses for the care of its beneficiaries [the Pediatric Hospital (PH) in the National Medical Center XXI Century and the General Hospital in the National Medical Center La Raza (LR)]. These hospitals contain the necessary infrastructure to establish the precise diagnosis of cancer in patients. These are the reference centers for children with cancer from other states in the Mexican republic. Only the data regarding cancer in children coming from the DF, EM, Mor, Gro, and Chis were included in the analysis because data on cancer cases in children from the mentioned states are sent to the DF for their treatment, and they have been registered there since 1996, the year the RCC was initiated.<sup>11</sup>

The analyzed cases were carefully selected only once to avoid duplicate cases. In 97.5% of the lymphoma and solid tumor cases, the diagnosis was confirmed by means of histopathological study, and in 100% of the leukemia cases, the diagnosis was confirmed through bone marrow aspiration.

- b) *Mortality Data.* Mortality data were obtained from the *Instituto Nacional de Estadística y Geografía* (INEGI) [National Institute of Statistics and Geography].<sup>12</sup> For the study period, two International Classifications of Diseases (ICD) were used for coding the causes of death (ICD-9 for 1990-1997 and ICD-10 for the period 1998-2013); therefore, the table of equivalences for the study of mortality data, in which the equivalence of the ICD-9 to ICD-10 codes is established, was used. The ICD-9 codes 140-240 and the ICD-10 codes C00-97 were used.<sup>13</sup>

The selection criteria for the analysis of the incidence and mortality were as follows: a) children who were diagnosed with cancer of any ICCG group;<sup>10</sup> b) MSSI beneficiaries under the age of 15; and c) residents of the DF, EM, Mor, Gro, and Chis.

The denominators for calculating the incidence and mortality rates corresponded to the population under 15 years of age who were MSSI-B Ascribed to Family Medicine and residents of the selected states; the denominators were obtained from the Coordination of Medical Care of the MSSI.<sup>14</sup>

The absolute frequencies of the different groups and subgroups of cancer were obtained, and the overall and specific incidence was estimated by group and subgroup of neoplasia cases according to the ICCG<sup>10</sup> and was stratified by age, sex, and state of residence. Age was grouped into the following categories: less than 1 year, 1-4, 5-9, and 10-14 years (rates given for 1 000 000 children/year, unless otherwise specified). A rate adjustment was performed by the direct method for the total incidence and the incidence by subgroup. The reference population was the population worldwide.<sup>15</sup>

Furthermore, the annual mortality rates were obtained overall and were stratified by state. To evaluate the trend in both incidence and mortality, the annual percentage of change and the average annual percentage of change during the study period were obtained (APC and AAPC, respectively). These indicators are a weighted average of the annual percentage of increase or decrease that occurs in a population within a short period (APC) or during the entire study period (AAPC). For this analysis, the Joinpoint Regression Analysis program (version 4.0) that provides the Surveillance, Epidemiology, and End Results Program of the United

States of America was used, and a  $p \leq 0.05$  was accepted as the limit of statistical significance.\* The research protocol was approved by the Committee on Research and Ethics of the Helsinki Protocol.

## Results

The overall incidence decreased for the period of 1996-2001 and then stabilized for the period of 2002-2013 [APC = -6.6% (95%CI -11.0, -1.9) and 0.6% (95%CI -0.6, 1.9), respectively]. The same trend occurred in EM and Chis. Both Mor and Gro displayed a decreasing trend for the entire period [AAPC = -2.3% (95%CI -4.1, -0.6) and -2.6% (95%CI -4.9, -0.2), respectively]. The DF displayed a stable trend throughout the period (table I, figure 1).

The same aforementioned phenomenon was found in the overall trend in mortality. There was a decreasing trend for the period of 1996-2002 (APC = -7.0% (95%CI -11.4, -2.4), but mortality remained stable for the following period (2003-2013). EM and Chis displayed a decreasing trend for the entire period [AAPC = -3.3% (95%CI -5.0, -1.5) and -6.2% (95%CI -9.3, -3.1), respectively]. The trends for the DF, Mor, and Gro were stable (table I, figure 1).

Because the trend in overall mortality was stable over the final 11 years (2003-2013), the average exact rate for that period, overall and for each state, was obtained. Overall, the mortality rate was 56.7, and for the DF, EM, Mor, Gro, and Chis, the mortality rate was 61.3, 49.1, 68.9, 68.4, and 85.0, respectively.

For incidence, the same phenomenon was observed in the neoplasia, general leukemia, and acute lymphoid groups. During the period of 1996-2001, there was a decreasing trend, and for the period of 2002-2013, the trend was stable [APC = -7.3% (-17.3, -2.7) and -6.1% (-11.1, -0.5), respectively]. With respect to other groups of neoplasias, myeloid leukemias, lymphomas in general, Hodgkin's lymphoma and non-Hodgkin's lymphoma (NHL) displayed a decreasing trend for the entire study period (table II).

Moreover, the overall incidence was 128.0 cases per 1 000 000 children for all of the different groups of cancer. Leukemias, central nervous system tumors (CNST), and lymphomas were the main groups of neoplasias that were presented, and they displayed incidences of 58.8, 15.3, and 13.0, respectively (table III).

\* Joinpoint Regression Program. 2012. Joinpoint Regression Program, Version 4.0 - December 2012; Statistical Methodology and Applications Branch, Surveillance Research Program, National Cancer Institute. <http://surveillance.cancer.gov/joinpoint/faq/>

**Table I**  
**INCIDENCE\* AND MORTALITY\* TRENDS ACCORDING TO THE STATE OF RESIDENCE**  
**IN THE MEXICAN REPUBLIC OF BENEFICIARY CHILDREN OF THE MSSI. 1996-2013**

State	% Change 1996-2013	Incidence					
		APC <sub>1</sub>		APC <sub>2</sub>		AAPC	
		1996-2001	95%CI	2002-2013	95%CI	1996-2013	95%CI
Overall	-29.8	-6.6 <sup>‡</sup>	-11.0, -1.9	0.6	-0.6; 1.9	-1.5 <sup>‡</sup>	-3.0, -0.0
Distrito Federal	-22.7	—	—	—	—	0.4	-1.0, 0.6
Estado de México	-21.6	-7.1 <sup>‡</sup>	-12.2, -1.8	1.3	-0.2, 2.8	-1.3	-3.8, 0.5
Morelos	-42.5	—	—	—	—	-2.3 <sup>‡</sup>	-4.1, -0.6
Guerrero	-47.4	—	—	—	—	-2.6 <sup>‡</sup>	-4.9, -0.2
Chiapas	-69.2	-11.6 <sup>§‡</sup>	-14.8, -8.3	24.0 <sup>#</sup>	-16.1, 83.4	-6.2	-12.4, 0.5

State	% Change 1996-2013	Mortality					
		APC <sub>1</sub>		APC <sub>2</sub>		AAPC	
		1996-2002	95%CI	2003-2013	95%CI	1996-2013	95%CI
Overall	-37.7	-7.0 <sup>‡</sup>	-11.4, -2.4	-0.5	-2.1, 1.5	-2.8 <sup>‡</sup>	-4.7, -0.9
Distrito Federal	-43.9	—	—	—	—	-1.3	-2.6, 0.1
Estado de México	-43.9	—	—	—	—	-3.3 <sup>‡</sup>	-5.0, -1.5
Morelos	8.0	—	—	—	—	0.4	-4.5, 3.9
Guerrero	65.7	—	—	—	—	1.7	-2.3, 5.9
Chiapas	-70.8	—	—	—	—	-6.2 <sup>‡</sup>	-9.3, -3.1

\* Rates adjusted by age for 1 000 000 children/year; ICCC: International Classification for Childhood Cancer (reference 10); APC: Annual percentage of change; AAPC: Average annual percentage change for the period

<sup>‡</sup> The APC is significantly different from zero at an alpha of 0.05; CI95% = 95% confidence interval

<sup>§</sup> Period 1996-2010

<sup>#</sup> Period 2010-2013

MSSI: Mexican Social Security Institute

The highest incidence within the subgroups of leukemias occurred for acute lymphoid leukemia (48.3); of the CNSTs, astrocytomas occurred at the highest incidence (6.2), and NHL displayed the highest incidence in the lymphoma group (5.6) (table III).

In relation to other neoplasias, germ cell tumors (GCTs) were the 4<sup>th</sup> most common type, with an incidence of 8.6. The remainder of the neoplasias had lower incidences than those noted (table III).

The highest incidence according to age occurred in the 1-4-year-old age group, followed by infants less than 1 year old, children 10-14 years old, and finally children 5-9 years old, with rates of 169.1, 141.1, 113.3, and 98.7, respectively. The incidence according to sex was slightly higher in males (ratio M/F=1.1) (table III).

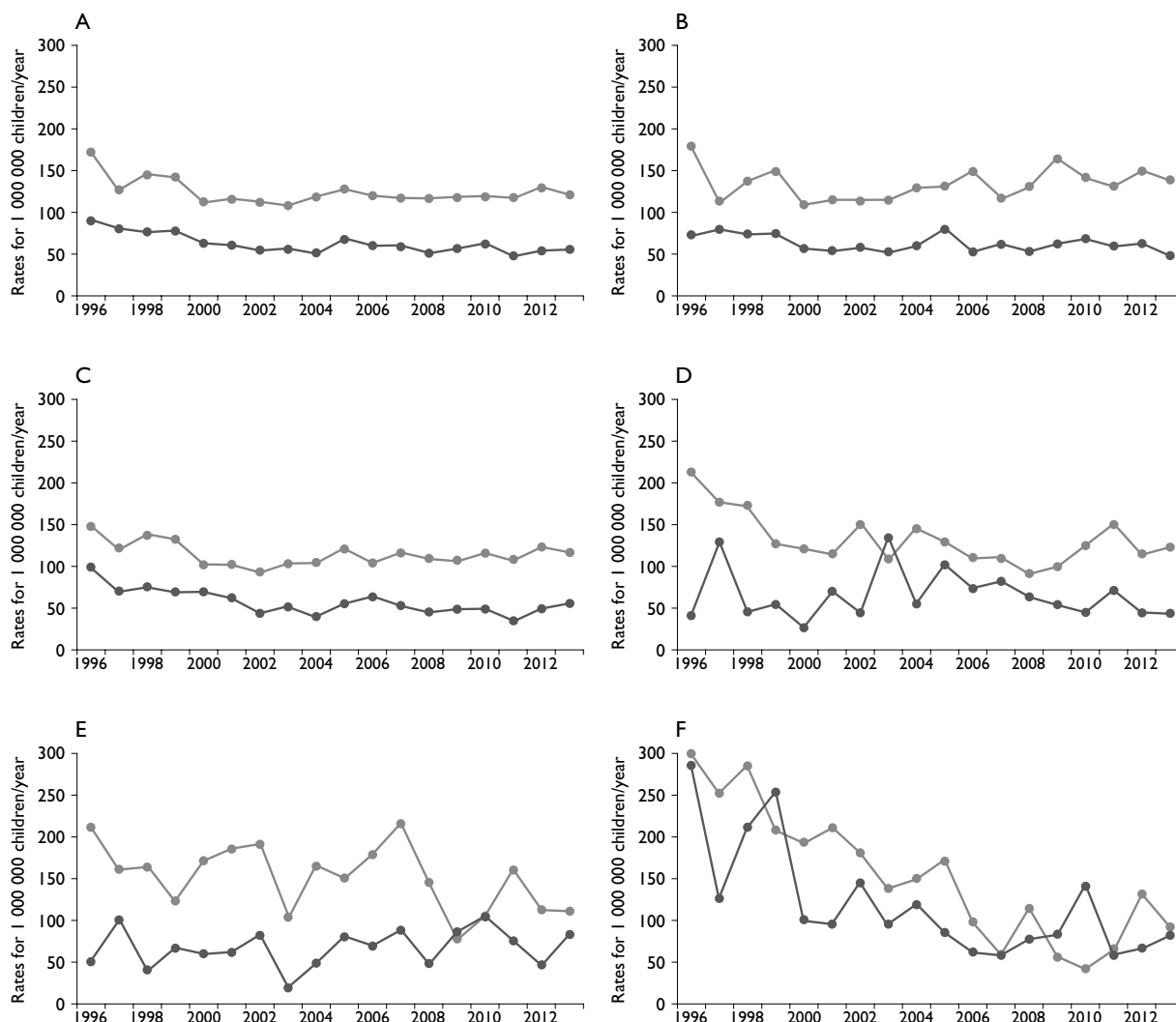
The highest incidence with respect to the state of residence occurred in Gro, followed by Chis, DF, Mor, and EM (rates of 155.1, 152.7, 137.8, 133.8, and 116.2, respectively). In general, the pattern of the main neoplasias was as follows: leukemias, CNST, and lymphomas,

except in EM, wherein both lymphomas and CNSTs had a similar incidence (table IV).

## Discussion

One of the most important aspects identified in the present study is related to the data on the trends of both incidence and mortality. Having already reported the phenomenon, incidence displayed a decrease during the period of 1996-2001,<sup>9</sup> which was spontaneous and subsequently stabilized through 2013. We consider this trend to be a real effect and not the result of poor records in the RCC. The mortality data encountered also lends support to this phenomenon because the observed trend in incidence was also encountered in mortality, which was even presented with a year of delay because the decline occurred during 1996-2002 (table I, figure 1).

We do not know the causes of this phenomenon, but as noted when we first reported the decline, we believe



A. Overall, B. Distrito Federal, C. Estado de México, D. Morelos; E. Guerrero; F. Chiapas

**FIGURE 1. TREND OF CANCER INCIDENCE AND MORTALITY IN BENEFICIARY CHILDREN OF THE MEXICAN SOCIAL SECURITY INSTITUTE WHO WERE RESIDENTS OF FIVE STATES OF THE MEXICAN REPUBLIC (1993-2013)**

that there was a period effect in the study population. That is, prior to 1996, the population of children may have been exposed to a harmful exposure, during which the incidence increased; subsequently, it was eliminated or the exposure was terminated; therefore, (spontaneously) the incidence and mortality decreased, as shown by the data from this study.

In support of this hypothesis, in a study conducted in Mexico, an increase in the mortality rate from 46.8 to 56.3 during the period of 1980-1995 was identified.<sup>16</sup> Moreover, in the evaluation of cancer mortality during

1970-2007 that was performed by L. Chatenoud and colleagues, in countries in America, Asia, and Oceania, an abrupt increase in the trend of mortality was observed in Mexico during the period of 1985-1990, which supports our comments.<sup>7</sup>

Due to the absence of incidence data for periods preceding that recorded (1996-2013) and with the objective of identifying specific points and attempting to explain the phenomenon found, further exploration of the data on cancer mortality in Mexican children in greater detail should be considered.

**Table II**  
**INCIDENCE\* TRENDS ACCORDING TO GROUPS AND SUBGROUPS OF THE ICCC IN BENEFICIARY CHILDREN OF THE MSSI WHO WERE RESIDENTS OF FIVE STATES OF THE MEXICAN REPUBLIC. 1996-2013**

Group and/or subgroup of neoplasias	% Change 1996-2013	APC <sub>1</sub>		APC <sub>2</sub>		AAPC	
		1996-2001	95%CI	2002-2013	95%CI	1996-2013	95%CI
All neoplasias	-29.8	-6.6 <sup>‡</sup>	(-11.0, -1.9)	0.6	(-0.6, 1.9)	-1.5 <sup>‡</sup>	(-3.0, -0.0)
I. Leukemias	-22.1	-7.3 <sup>‡</sup>	(-17.7, -2.7)	1.3	(-0.1, 3.0)	-1.3	(-2.8, 0.2)
Ia. Acute lymphoid leukemia	-12.3	-6.1 <sup>‡</sup>	(-11.1, -0.5)	1.4	(-0.1, 3.0)	-0.9	(-2.7, 1.0)
Ib. Acute myeloid leukemia	-66.4	—	—	—	—	-2.1 <sup>‡</sup>	(-4.6, -0.5)
II. Lymphomas	-57.0	—	—	—	—	-3.7 <sup>‡</sup>	(-5.4, -1.9)
IIa. Hodgkin's lymphomas	-70.1	—	—	—	—	-4.1 <sup>‡</sup>	(-7.2, -0.8)
IIb. Non-Hodgkin's lymphomas	-52.3	—	—	—	—	-3.2 <sup>‡</sup>	(-0.6, -0.3)
III. Neoplasias of the central nervous system	-16.2	—	—	—	—	0.2	(-1.3, 1.7)
IV. Neuroblastomas and other PNS	-73.0	—	—	—	—	-3.8	(-7.7, 0.1)
V. Retinoblastoma	24.0	—	—	—	—	-1.2	(-5.0, 2.8)
VI. Renal tumors	78.0	—	—	—	—	-0.7	(-3.1, 2.2)
VII. Hepatic tumors	-56.1	—	—	—	—	-0.7	(-6.3, 5.3)
VIII. Malignant bone tumors	-14.4	—	—	—	—	-1.5	(-3.6, 0.6)
IX. Soft tissue sarcomas	-46.2	—	—	—	—	-1.0	(-4.2, 2.2)
X. Germ cell tumors	-29.3	—	—	—	—	-0.2	(-2.0, 1.5)
XI. Carcinomas	-25.1	—	—	—	—	1.4	(-2.9, 5.8)

\* Rates adjusted by age for 1 000 000 children/year; ICCC: International Classification for Childhood Cancer (reference 10); APC: Annual percentage of change; AAPC: Average annual percentage change for the period

<sup>‡</sup> The APC is significantly different from zero at an alpha of 0.05; 95%CI= 95% confidence interval

MSSI: Mexican Social Security Institute

Furthermore, relating exclusively to the incidence trend for the period of 2001-2013, the overall trend for all of the states and for the group of leukemias and the acute lymphoid subgroup was stable (tables I and II), which contrasts with that reported in developed countries (the United States of America and European countries), where it has been noted that cancer in children and, above all acute lymphoblastic leukemia, has increased.<sup>13,17</sup> There is controversy in relation to myeloid leukemias. North American children displayed an increasing trend during the period of 1975-1993 and stability during the period of 1994-2012;<sup>13</sup> in contrast, for European children, the trend has remained stable.<sup>18</sup> Our data showed a decreasing trend for myeloid leukemias throughout the study period (table II).

The trend of lymphomas occurred similarly in North American children compared to the previously discussed phenomenon. The trend decreased for the period of 1975-1993, but the period of 1994-2012 displayed an increasing trend.<sup>13</sup> In European children, there was an increasing trend in general, but this only occurred in the southern and western regions of Europe.<sup>19</sup> Data from our study showed a decreasing trend, and in a more detailed study of lymphomas that was performed

by our group, the mixed cellularity subtype decreased in Hodgkin's lymphomas; non-Hodgkin's lymphomas also displayed a decreasing trend, particularly in the unspecified types, which was identified as an improvement in the histopathological diagnosis in recent years.<sup>20</sup>

In relation to mortality exclusively, which, as mentioned, is related to the overall efficacy of treatment that is being administered to the population,<sup>6,21</sup> only the data from the period of 2003-2013 should be considered because during this period, the incidence was stable and it could be verified that the incidence differed from what occurred in developed countries, where the cancer mortality in children decreased.<sup>4,7,13,22</sup> This did not occur for the children in our study and in general for Mexican children because similar results were found when cancer mortality for children throughout the country was studied.<sup>7,23</sup>

Furthermore, the exact mortality rate for some developed countries is between 22 and 30 deaths per 1 000 000 children,<sup>7,13</sup> whereas in our data, it corresponded to 49.3 and 85 (mortality rates for EM and Chiapas, respectively). This situation has been mentioned even by foreign authors.<sup>7,24</sup> Thus, the treatment of children with cancer in our country requires a better organiza-

**Table III**  
**INCIDENCE\* OF MALIGNANT NEOPLASIAS IN BENEFICIARY CHILDREN**  
**OF THE MSSI WHO WERE RESIDENTS OF FIVE STATES OF THE MEXICAN REPUBLIC†, 1996-2013**

Groups and subgroups of neoplasias according to ICCC		Age strata (years)										Sex				Ratio M/F
		<1		1 to 4		5 to 9		10 to 14		0 to 14 <sup>s</sup>		M		F		
		n	rate	n	rate	n	rate	n	rate	n	rate	n	rate	n	rate	
I.	Leukemias	68	35.4	869	85.4	661	49.6	587	46.8	2185	58.8	1177	60.8	1008	54.1	1.1
Ia.	Lymphoid leukemias	48	25.0	741	72.8	566	42.5	437	34.9	1792	48.3	968	50.0	824	44.2	1.1
Ib.	Acute myeloid leukemias	15	7.8	110	10.8	85	6.4	132	10.5	342	9.1	184	9.5	158	8.5	1.1
Ic.	Chronic myeloproliferative diseases	3	1.6	6	0.6	7	0.5	11	0.9	27	0.7	13	0.7	14	0.8	0.9
Id.	Myelodysplastic syndromes and other myeloproliferative processes	1	0.5	4	0.4	0	0.0	1	0.1	6	0.2	3	0.2	3	0.2	1.0
Ie.	Nonspecific leukemias and others	1	0.5	8	0.8	3	0.2	6	0.5	18	0.5	9	0.5	9	0.5	1.0
II.	Lymphomas and reticuloendothelial neoplasias	17	8.9	118	11.6	179	13.4	191	15.2	505	13.0	352	18.2	153	8.2	2.2
IIa.	Hodgkin's lymphomas	0	0.0	16	1.6	60	4.5	88	7.0	164	4.0	119	6.1	45	2.4	2.5
IIb.	Non-Hodgkin's lymphomas (excluding Burkitt)	0	0.0	52	5.1	86	6.5	84	6.7	222	5.6	158	8.2	64	3.4	2.4
IIc.	Burkitt lymphoma	1	0.5	20	2.0	27	2.0	16	1.3	64	1.7	42	2.2	22	1.2	1.8
IId.	Miscellaneous lymphoreticular neoplasias	16	8.3	28	2.8	4	0.3	1	0.1	49	1.3	27	1.4	22	1.2	1.2
IIf.	Nonspecific lymphomas	0	0.0	2	0.2	2	0.2	2	0.2	6	0.2	6	0.3	0	0.0	–
III.	Intracranial and intraspinal CNS neoplasias	21	10.9	196	19.3	195	14.6	161	12.8	573	15.3	290	15.0	283	15.2	1.0
IIIa.	Ependymomas and choroid plexus tumors	3	1.6	38	3.7	25	1.9	23	1.8	89	2.4	39	2.0	50	2.7	0.8
IIIb.	Astrocytomas	8	4.2	64	6.3	84	6.3	80	6.4	236	6.2	115	5.9	121	6.5	0.9
IIIf.	Intracranial and intraspinal embryonal tumors	5	2.6	72	7.1	58	4.4	44	3.5	179	4.8	108	5.6	71	3.8	1.5
IIId.	Other gliomas	0	0.0	10	1.0	15	1.1	5	0.4	30	0.8	12	0.6	18	1.0	0.6
IIIf.	Other specific intracranial and intraspinal neoplasias	5	2.6	10	1.0	11	0.8	9	0.7	35	1.0	14	0.7	21	1.1	0.6
IIIf.	Unspecified intracranial and intraspinal neoplasias	0	0.0	2	0.2	2	0.2	0	0.0	4	0.1	2	0.1	2	0.1	1.0
IV.	Neuroblastoma and other tumors of peripheral nerve cells	36	18.7	50	4.9	13	1.0	5	0.4	104	3.4	49	2.6	55	3.0	0.9
IVa.	Neuroblastoma and ganglioneuroblastoma	36	18.7	49	4.8	12	0.9	3	0.2	100	3.3	46	2.4	54	2.9	0.8
IVb.	Other tumors of peripheral nerve cells	0	0.0	1	0.1	1	0.1	2	0.2	4	0.1	3	0.2	1	0.1	2.9
V.	Retinoblastoma	39	20.3	125	12.3	8	0.6	3	0.2	175	5.6	91	4.7	84	4.5	1.0
VI.	Renal tumors	21	10.9	126	12.4	39	2.9	14	1.1	200	5.9	87	4.5	103	6.0	0.8
VIa.	Nephroblastoma and other non-epithelial renal tumors	21	10.9	125	12.3	39	2.9	11	0.9	196	5.8	87	4.5	109	5.8	0.8
VIb.	Renal carcinoma	0	0.0	1	0.1	0	0.0	3	0.2	4	0.1	0	0.0	4	0.2	–
VII.	Hepatic tumors	18	9.4	46	4.5	9	0.7	13	1.0	86	2.6	48	2.6	38	2.1	1.2
VIIa.	Hepatoblastoma	18	9.4	44	4.3	8	0.6	3	0.2	73	2.3	40	2.1	33	1.8	1.2
VIIb.	Hepatic carcinoma	0	0.0	2	0.2	1	0.1	9	0.7	12	0.3	7	0.4	5	0.3	1.3
VIIc.	Nonspecific malignant hepatic tumors	0	0.0	0	0.0	0	0.0	1	0.1	1	0.0	1	0.1	0	0.0	–
VIII.	Malignant bone tumors	1	0.5	17	1.7	70	5.3	171	13.6	259	6.2	130	6.8	129	7.0	1.0
VIIIa.	Osteosarcomas	0	0.0	9	0.9	55	4.1	131	10.4	195	4.6	97	5.0	98	5.3	1.0
VIIIb.	Chondrosarcoma	0	0.0	1	0.1	0	0.0	4	0.3	5	0.1	0	0.0	5	0.3	0.0
VIIIc.	Ewing tumor and related bone sarcomas	1	0.5	6	0.6	12	0.9	31	2.5	50	1.3	31	1.6	19	1.0	1.6
VIIId.	Other specific malignant bone tumors	0	0.0	1	0.1	2	0.2	3	0.2	6	0.1	1	0.1	5	0.3	0.2
VIIIe.	Nonspecific malignant bone tumors	0	0.0	0	0.0	1	0.1	2	0.2	3	0.1	1	0.1	2	0.1	0.5
IX.	Soft tissue and extraosseous sarcomas	20	10.4	75	7.4	68	5.1	92	7.3	255	6.9	121	6.3	134	7.2	0.9
IXa.	Rhabdomyosarcoma	12	6.2	42	4.1	39	2.9	34	2.7	127	3.5	67	3.5	60	3.2	1.1
IXb.	Fibrosarcomas, peripheral nerves and other fibrous tumors	4	2.1	9	0.9	6	0.5	15	1.2	34	0.9	17	0.9	17	0.9	1.0
IXc.	Kaposi's sarcoma	4	2.1	23	2.3	23	1.7	42	3.3	92	2.4	37	1.9	55	3.0	0.6
IXe.	Nonspecific soft tissue sarcomas	0	0.0	1	0.1	0	0.0	1	0.1	2	0.1	0	0.0	2	0.1	0.0
X.	Germ cell tumors of the gonads and trophoblasts	30	15.6	90	8.8	55	4.1	145	11.6	320	8.6	172	8.9	148	8.0	1.1
Xa.	Intracranial and intraspinal germ cell tumors	2	1.0	5	0.5	24	1.8	37	3.0	68	1.7	50	2.6	18	1.0	2.7
Xb.	Extracranial and extragonadal germ cell tumors	11	5.7	11	1.1	2	0.2	11	0.9	35	1.5	20	1.0	15	0.8	1.3
Xc.	Malignant gonadal germ cell tumors	17	8.9	72	7.1	29	2.2	95	7.6	213	5.8	101	5.2	112	6.0	0.9
Xd.	Other unspecified malignant gonadal tumors	0	0.0	2	0.2	0	0.0	2	0.2	4	0.1	1	0.1	3	0.2	0.3
XI.	Other malignant epithelial neoplasias and melanomas	0	0.0	9	0.9	18	1.4	36	2.9	63	1.5	26	1.4	37	2.1	0.7
XIa.	Adrenocortical carcinoma	0	0.0	6	0.6	2	0.2	1	0.1	9	0.3	4	0.2	5	0.3	0.8
XIb.	Thyroid carcinoma	0	0.0	0	0.0	7	0.5	12	1.0	19	0.4	6	0.3	13	0.7	0.4
XIc.	Nasopharyngeal carcinoma	0	0.0	0	0.0	1	0.1	3	0.2	4	0.1	3	0.2	1	0.1	2.9
XId.	Malignant melanoma	0	0.0	1	0.1	4	0.3	1	0.1	6	0.2	2	0.1	4	0.2	0.5
XIe.	Skin carcinoma	0	0.0	0	0.0	0	0.0	4	0.3	4	0.1	1	0.1	3	0.2	0.3
XIf.	Nonspecific carcinomas and others	0	0.0	2	0.2	4	0.3	15	1.2	21	0.5	10	0.5	11	0.6	0.9
XIIa.	Other nonspecific malignant tumors	0	0.0	1	0.1	0	0.0	2	0.2	3	0.1	0	0.0	3	0.2	0.0
Total		271	141.1	1722	169.1	1315	98.7	1420	113.3	4728	128.0	2543	131.4	2185	117.3	1.1

\* Rates per 1 000 000 children/year

† Distrito Federal, Estado de México, Morelos, Guerrero, and Chiapas; MSSI = Mexican Social Security Institute; ICCC: International Classification for Childhood Cancer (reference 10)

§ Rates adjusted for age; n: number of cases; M: male; F: female; CNS: central nervous system



**Table IV**  
**INCIDENCE\* OF MALIGNANT NEOPLASIAS IN BENEFICIARY CHILDREN OF THE MSSI WHO WERE RESIDENTS OF FIVE STATES OF THE MEXICAN REPUBLIC.† 1996-2013**

Groups of neoplasias according to ICCC	DF		Edo Mex		Morelos		Guerrero		Chiapas		Total	
	n	rate	n	rate	n	rate	n	rate	n	rate	n	rate
I. Leukemias	768	62.2	1 120	58.6	107	52.4	109	58.2	81	43.5	2 185	58.8
II. Lymphomas	198	15.5	216	10.7	26	12.7	38	20.3	27	13.6	505	13.0
III. Intracranial and intraspinal CNS neoplasias	237	18.9	203	10.4	35	18.2	46	23.7	52	30.2	573	15.3
IV. Neuroblastoma and other tumors of peripheral nerve cells	41	4.0	40	2.5	4	2.7	14	9.0	5	3.8	104	3.4
V. Retinoblastoma	47	4.3	83	5.2	12	7.8	11	6.8	22	16.8	175	5.6
VI. Renal tumors	62	5.4	93	5.5	14	7.8	15	8.8	16	9.9	200	5.9
VII. Hepatic tumors	39	3.6	32	1.9	8	4.9	3	1.8	4	2.5	86	2.6
VIII. Malignant bone tumors	90	6.5	116	5.4	15	6.6	18	8.8	20	9.5	259	6.2
IX. Soft tissue sarcomas and other extraosseous sarcomas	92	7.2	121	6.4	14	7.0	11	6.2	17	9.6	255	6.9
X. Germ cell tumors	113	8.8	159	8.6	21	10.7	11	5.8	16	8.6	320	8.6
XI. Other malignant epithelial neoplasias, melanomas	18	1.3	20	0.9	6	2.8	10	5.3	9	4.2	63	1.5
XII. Nonspecific neoplasias	1	0.1	2	0.1	0	0.0	0	0.0	0	0.0	3	0.1
Total	1 706	137.8	2 205	116.2	262	133.8	286	155.1	269	152.7	4 728	128.0

Rates are age adjusted per 1 000 000 children/year

† Distrito Federal, Estado de México, Morelos, Guerrero, and Chiapas; MSSI: Mexican Social Security Institute; ICCC: International Classification for Childhood Cancer (reference 10); n: number of cases; CNS: central nervous system

tion of the health system in charge of offering medical care for Mexican children with cancer, and we believe that research is the key through comprehensive studies and through research in conjunction with international organizations.<sup>1</sup>

With regard to the incidence of the different groups and subgroups of cancer, the data obtained in the present study are consistent with what has previously been reported in MSSI-B children.<sup>25</sup>

In relation to GCT only, it is noteworthy that the incidence found (8.6) is one of the highest worldwide; only children of Gro had a lower incidence (5.6), which was similar to that of North American and German children (5.5 and 4.9, respectively).<sup>13,26</sup> Therefore, we believe that the epidemiology of this group of cancers should be studied in detail worldwide to establish possible etiological hypotheses.

Nationally, some data on the epidemiology of children with cancer who were treated using the strategy of the *Seguro Popular* [People's Insurance] have been published. We could not compare our data with the data of that group because the populations were different. *Seguro Popular* reported on children 0 to 18 years of age, whereas the present study reported on children 0 to 14 years of age. Notably, the ICCC was used to group the different cancer groups, but the groups of neoplasias that were presented are different from that classification,

and the data on the incidence according to age were not calculated according to traditional methods.<sup>27</sup>

## Conclusions

The incidence and mortality rates from cancer in the study population displayed spontaneous declines for the period of 1996-2001 and 1996-2002, respectively. For the additional period (2002-2013; 2003-2013), the trend for both indicators remained stable. According to the mortality data, the efficacy of the treatment that was offered to the children studied was not as effective as that attained by children in developed countries, indicating the need to conduct research, interinstitutional and/or international, to identify the causes of the treatment efficacy and to improve the treatment. The incidence data according to the neoplasm group, age, sex, and place of residence are consistent with those reported previously.

*Declaration of conflict of interests.* The authors declare that they have no conflict of interests.

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