

RESEARCH ARTICLE

Epidemiological Trends of Histopathologically WHO Classified CNS Tumors in Developing Countries: Systematic Review

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Abstract

Background: Many developing countries are lagging behind in reporting epidemiological data for individual central nervous system (CNS) tumors. This paper aimed to elicit patterns for the epidemiology of individual World Health Organization (WHO) classified CNS tumors in countries registered by WHO as “developing”. **Materials and Methods:** Cyber search was carried out through 66 cancer networks/registries and 181 PubMed published papers that reported counts of CNS tumors for the period of 2009-2012. The relationship between the natural log of incidence Age Standardized Rate (ASR) reported by Globocan and Latitude/ Longitude was investigated. **Results:** Registries for 21 countries displayed information related to CNS tumors. In contrast trends for classified CNS tumor cases were identified for 38 countries via 181 PubMed publications. Extracted data showed a majority of unclassified reported cases [PubMed (38 countries, 45.7%), registries (21 countries, 96.1%)]. For classified tumors, astrocytic tumors were the most frequently reported type [PubMed (38 countries, 1,245 cases, 15.7%), registries (21 countries, 627 cases, 1.99%)]. A significant linear regression relationship emerged between latitudes and reported cases of CNS tumors. **Conclusions:** Previously unreported trends of frequencies for individually classified CNS tumors were elucidated and a possible link of CNS tumors occurrence with geographical location emerged.

Keywords: CNS tumors - epidemiology - developing countries - cancer registries - latitude - risk factors

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Introduction

Eliciting patterns for the distribution of disease between different populations over time is important for the development and the realization of health policies (Chauvin et al., 2012). For tumors originating in the Central Nervous System (CNS), there is a collection of a large spectrum of neoplasms classified by World Health Organization (WHO), with each tumor having its own features, location, morphology, prognosis and treatment (DeAngelis, 2001; Louis et al., 2007; Katchy et al., 2011).

The latest Globocan report for 2012 estimated a total incidence ASR of 3.4 per 100,000 and a mortality ASR of 2.5 per 100 000 worldwide (Parkin et al., 2001; GLOBOCAN, 2014). Incidence ASR of CNS tumors in the developing region was estimated to be 3.0 per 100,000, and for mortality ASR the estimation was 2.2 per 100,000. In addition, Globocan described gender-related data and showed that men have a higher CNS tumors incidence ASR 3.9 compared with women 3.0 worldwide. Unfortunately, however, trends for individually classified types of CNS tumors are not available from this database.

Many developed countries across the globe have developed cancer registries, which record and provide data of CNS tumors' specific incidences and mortalities (Surawicz et al., 1999; CancerResearchUK, 2013a; Villano et al., 2013; Woehrer, 2013). Recent reports show that Astrocytomas were the most common in the UK (CancerResearchUK, 2013b), Meningiomas were most reported in the USA (Dolecek et al., 2012) and glioblastoma multiform (GBM) tumors had the highest frequencies in Canada (BrainTumourFoundation, 2013).

Developing countries are lagging behind in reporting epidemiological data for cancer (Ferlay et al., 2010; Qaddoumi et al., 2011). In addition, although there are cancer registries for some developing countries, they inadequately report data for different types of CNS tumors (AFCRN, 2013). To obtain an estimated perspective of the current status of CNS tumors in developing countries, detailed and global epidemiological studies are required (Katchy et al., 2011). This work aimed to elicit patterns for the epidemiology of individually reported CNS tumors subtypes in countries registered by the WHO as “developing”.

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Materials and Methods*Search Strategy and Data Extraction*

Cyber methods of search strategies were implemented

to retrieve data for the incidence of CNS tumors between the years 2009-2012 in 149 countries registered by WHO as “developing” (Figure 1). Briefly, for cancer registries/networks search, Google website was searched to find

Table 1. A list of Countries who Reported CNS Tumor Cases Via Cancer Registries or PubMed and the Predominant Tumor Types

Region Country	Reported in Cancer Registry			Reported in PubMed		
	No. of Cases Total (%)	Predominant Tumors Type	No. of cases (%)	No. of Cases Total (%)	Predominant Tumors Type	No. of cases (%)
EAP						
China	NR	NR	NR	685 (8.66)	Meningeal	180 (26.3)
Malaysia	NR	NR	NR	2 (0.03)	None	None
Nepal	77 (0.24)	Unclassified	77 (100)	NR	NR	NR
Thailand	117 (0.37)	Astrocytic	63 (53.8)	88 (1.11)	Gliomas	61 (69.3)
ECA						
Albania	381 (1.21)	Unclassified	381 (100)	NR	NR	NR
Belarus	501 (1.59)	Unclassified	501 (100)	NR	NR	NR
Bosnia & Herzegovina	299 (0.95)	Unclassified	299 (100)	30 (0.38)	Astrocytic	16 (53.3)
Bulgaria	2123 (6.73)	Unclassified	2123 (100)	2781 (35.1)	Unclassified	2508 (90.2)
Croatia	1627 (5.16)	Unclassified	1627 (100)	57 (0.72)	Astrocytic	40 (70.2)
Georgia	NR	NR	NR	495 (6.26)	Unclassified	190 (38.4)
Hungary	595 (1.89)	Unclassified	595 (100)	33 (0.42)	Astrocytic	14 (42.4)
Latvia	256 (0.81)	Unclassified	256 (100)	45 (0.57)	Gliomas	45 (100)
Lithuania	308 (0.98)	Unclassified	308 (100)	348 (4.4)	Meningeal	117 (33.6)
Moldova	174 (0.55)	Unclassified	174 (100)	NR	NR	NR
Montenegro	66 (0.21)	Unclassified	66 (100)	NR	None	None
Poland	4467 (14.1)	Unclassified	4467 (100)	73 (0.92)	Ependymal	39 (53.4)
Romania	1895 (6.01)	Unclassified	1808 (95.4)	168 (2.12)	Pituitary	142 (84.5)
Russian Federation	7377 (23.4)	Unclassified	7377 (100)	254 (3.21)	Unclassified	231 (90.9)
Serbia	2454 (7.78)	Unclassified	2454 (100)	6 (0.08)	None	None
Turkey	NR	NR	NR	242 (3.06)	Pituitary	146 (60.3)
Ukraine	7361 (23.3)	Unclassified	7361 (100)	NR	NR	NR
LAC						
Argentina	NR	NR	NR	29 (0.37)	Unclassified	22 (75.9)
Brazil	NR	NR	NR	38 (0.48)	Embryonal	37 (97.4)
Chile	NR	NR	NR	6 (0.08)	Pituitary	3 (50)
Colombia	NR	NR	NR	59 (0.75)	Pituitary	47 (79.7)
Mexico	NR	NR	NR	121 (1.53)	Metastatic	55 (45.5)
MENA						
Algeria	NR	NR	NR	2 (0.03)	None	None
Egypt	481 (1.53)	Unclassified	397 (82.5)	166 (2.1)	Gliomas	60 (36.1)
Iran	NR	NR	NR	1582 (20)	Unclassified	615 (38.9)
Jordan	172 (0.55)	Astrocytic	98 (57)	4 (0.05)	None	None
Lebanon	NR	NR	NR	4 (0.05)	None	None
Libya	NR	NR	NR	1 (0.01)	None	None
Morocco	NR	NR	NR	79 (1)	Astrocytic	34 (43)
Oman	NR	NR	NR	2 (0.03)	None	None
Qatar	NR	NR	NR	2 (0.03)	None	None
Saudi Arabia	281 (0.89)	Astrocytic	116 (41.3)	7 (0.09)	None	None
Tunisia	NR	NR	NR	7 (0.09)	None	None
UAE	NR	NR	NR	4 (0.05)	Cranial & Paraspinal Nerves	2 (50)
SA						
Bangladesh	NR	NR	NR	9 (0.11)	Meningeal	8 (88.9)
India	NR	NR	NR	70 (0.89)	Astrocytic	66 (94.3)
Pakistan	518 (1.64)	Astrocytic	277 (53.5)	376 (4.75)	Pituitary	282 (75)
SSA						
Kenya	NR	NR	NR	26 (0.33)	Unclassified	17 (65.4)
Nigeria	NR	NR	NR	5 (0.06)	None	None
South Africa	NR	NR	NR	4 (0.05)	Embryonal	2 (50)
Total Reported Cases	ALL	31530			7910	
Total Reporting Countries	ALL	21			38	

official websites of cancer registries networks, such as Globocan, AFRN (African Cancer-Registry Network) and EUCAN, or cancer registries for 149 individual countries. For PubMed search, four main limitation steps were applied. The first included year, tumor, incidence and country name, the second involved scanning retrieved abstracts for a list of key words, the third involved checking for authenticity of names (Sometimes a name of country overlapped with other elements for example the name Georgia stands for a European developing country and an American state) and finally a check on the date of the reported cases was applied (some papers were published in 2010 but were reporting cases prior to 2009).

Data analysis

Identified reported cases were fitted within 17 WHO classified groups (Louis et al., 2007) based on descriptive names by source. Grouped tumors for reported names included: astrocytic tumors [reported names were pilocytic astrocytoma, pilomyxoid astrocytoma, diffuse astrocytoma, fibrillary astrocytoma, anaplastic astrocytoma, gbm, gliosarcoma, and sega], meningeal tumors [reported names were meningioma, atypical meningioma, haemangiopericytoma and haemangioblastoma], oligodendroglial tumors [reported names were oligodendroglioma and anaplastic oligodendroglioma], embryonal tumors [reported names were embryonal, medulloblastoma, melanotic neuroectodermal tumour of infancy, and atypical teratoid/rhabdoid tumor], oligoastrocytic tumors [reported names were oligoastrocytoma and anaplastic oligoastrocytoma], ependymal tumors [reported names were ependymoma and myxopapillary ependymoma], tumors of cranial and paraspinal nerves [reported names were schwannoma, neurinoma, acoustic neuroma, nerve sheath tumors and spinal neurofibromatosis], other neuroepithelial tumors [reported names were astroblastoma and chordoid glioma], mesenchymal tumors [reported names were chondroma, lipoma, haemangioma, angiolipoma, rhabdomyosarcoma

and ewing sarcoma – pnet], neuronal and mixed neuronal-glial tumors [reported names were ganglioglioma, neurocytoma and papillary glioneuronal tumors], pineal tumors [reported names were pineoblastoma and other pineal tumors], germ cell tumors [reported names were germ cell, germinoma, yolk sac tumors and teratoma], and choroid plexus tumors [reported names were choroid plexus and choroid plexus papilloma]. tumors reported as “low grade gliomas” or “high grade gliomas” were included as an independent “gliomas” group. Unclassified tumors were also considered as an independent group.

Identified countries were grouped within six regions: Europe and Central Asia (ECA), Latin America and the Caribbean (LAC), East Asia and Pacific (EAP), Middle East and North Africa (MENA), South Asia (SA), and Sub-Saharan Africa (SSA), as per WHO cataloging.

Linear regression analyses for the relationship of Globocan reported incidence ASR (Ln) for all CNS tumors reported from 169 WHO registered countries, or 126 developing countries, verses latitude or longitude were run using SPSS Graduate Pack 16.0. A similar process was applied to test the significance of the relationship between PubMed reported Astrocytic tumors counts (Ln) for 25 countries with latitude or longitude. Natural logs of CNS tumors counts were chosen since values for counts had skewed distributions. For each country, latitude/longitude coordinates were taken from Google Earth 6.0.2. (Google, 2014).

Results

Countries reporting CNS tumors

Out of 66 investigated registries, registries for 21 countries provided information related to CNS tumors (Table 1). Unfortunately, this number was even lower for those registries that provided detailed WHO classifications of CNS tumors (Thailand, Egypt, Jordan, Saudi Arabia, Pakistan, and Romania). The total reported cases identified in cancer registries for the years 2009–2012 was 31530, with Russian Federation (7377 cases, 23.4%), Ukraine (7361 cases, 23.3%) and Poland (4467 cases, 14.1%) being the top contributors (Batut, 2009 and 2010; Chiricuta, 2010; NCREgypt, 2010; WHO, 2010; KFSHRC, 2011; CroatianNCR, 2012; EUCAN, 2012; IARC, 2012; NCIThailand, 2012; Piya and Acharya, 2012; SHATOBulgaria, 2012; AFRN, 2013; AMAAC, 2013; MOHJordan, 2013; NCIUkraine, 2013; ShaukatKhanum, 2013). In contrast, 38 countries have reported CNS tumor cases in 181 published papers (Ahmad et al., 2009; Akhaddar et al., 2009; Al-Dahri et al., 2009; Al-Hussain and Dababo, 2009; Ali et al., 2009; Alimohamadi et al., 2009; Alpizar-Aguirre et al., 2009; Awad et al., 2009; Bien et al., 2009; Boongird et al., 2009; Bozic et al., 2009; Charfi et al., 2009; Ciobanu et al., 2009; Conca et al., 2009; Enchev et al., 2009; Ghitou et al., 2009; Gomori et al., 2009; Grahovac et al., 2009; Jamjoom et al., 2009; Joo et al., 2009; Komolafe et al., 2009; Limaem et al., 2009; Liu et al., 2009; Mallea-Gil et al., 2009; Marinovic et al., 2009; Mikati et al., 2009; Mosqueda-Taylor et al., 2009; Naydenov et al., 2009; Olufemi Adeleye and Balogun, 2009; Pronin et al., 2009;

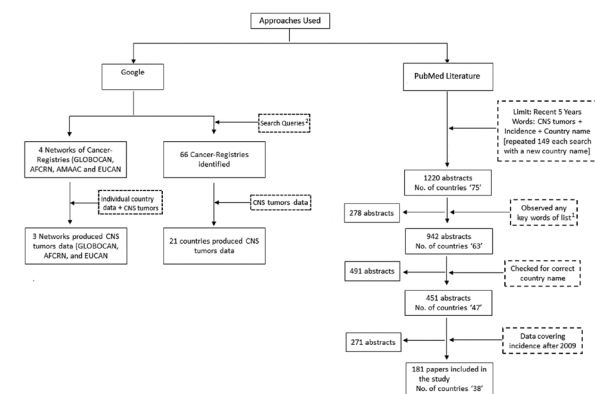


Figure 1. A Flow Chart for Search Strategy and Data Extraction Approaches Used in this Study. Dashed-Box (---) represents limitations used for the search. Keywords list1: CNS tumors in [Country name] 149 developing countries, Cancer, Registry, Data, Statistics, Incidence rate, Mortality/death rate, Prevalence, Epidemiology, Registration, Quantitative study, Clinical profile, or ASR. Search Queries2: [Country name] 149 developing countries and Cancer registry or Cancer Society or Cancer report or Cancer statistics

Salih et al., 2009; Sarwat and Al-Salam, 2009; Till et al., 2009; Vega et al., 2009; Alzarar et al., 2010; Arshad et al., 2010; Azarpira et al., 2010; Bankole et al., 2010; Benzagmout et al., 2010; Campos and Linhares, 2010; Chansriwong and Sirisinha, 2010; Dezmalj-Grbelja et al., 2010; Diez et al., 2010; El-Bilbeisi et al., 2010; El Beltagy et al., 2010; Fatehi et al., 2010; Gryngarten et al., 2010; Ishaqi et al., 2010; Jabri et al., 2010; Khawam et al., 2010; Khoddami and Gholampour Shahaboddini, 2010; Lakicevic et al., 2010; Las Heras and Pritzker, 2010; Masroujeh et al., 2010; Mondok et al., 2010; Nasir et al., 2010; Nicolae et al., 2010; Niculescu et al., 2010; Ortiz-Plata et al., 2010; Rizvi et al., 2010; Shamov et al., 2010; Vuckovic et al., 2010; Yaneva et al., 2010; Adeleye and Rabi, 2011; Ajani et al., 2011; Al-Hussaini et al., 2011; Al-Masri et al., 2011; Amirjamshidi et al., 2011; Asmoniene et al., 2011; Ayoub, 2011; Balinisteanu et al., 2011; Binesh et al., 2011; Garcia-Quintanilla et al., 2011; Ghosal et al., 2011; Jakab et al., 2011; Jamebozorgi et al., 2011; Khalatbari et al., 2011a; Khalatbari et al., 2011b; Khan et al., 2011; Kovacs et al., 2011; Maheshwari et al., 2011; Moona and Mehdi, 2011; Muller et al., 2011; Pruksakorn et al., 2011; Qiao et al., 2011; Quadri et al., 2011; Soliman et al., 2011; Soritau et al., 2011; Tomac et al., 2011; Tsekov et al., 2011; Tunon-Pitalua et al., 2011; Vulcani-Freitas et al., 2011; Witoonpanich et al., 2011; Zakrzewska et al., 2011; Zawrocki et al., 2011; Abbasi et al., 2012; Adeleye et al., 2012; Ajler et al., 2012; Akhtar et al., 2012; Albright and Okechi, 2012; Antic et al., 2012; Berindan-Neagoe et al., 2012; Bieza and Krumina, 2012; Binesh et al., 2012; Bochev et al., 2012; Carrasco et al., 2012; Chentli et al., 2012; Chiun et al., 2012; Choudhury et al., 2012; Chowdhury et al., 2012; Cortes-Medina and Guerrero-Rascon, 2012; Coulibaly et al., 2012; Cruz-Roa et al., 2012; Dawood and Cristofanilli, 2012; Deltuva et al., 2012; Dipro et al., 2012; Dizdarevic et al., 2012; Ech-Cherif El Kettani et al., 2012; Eshraghi et al., 2012;

Fedoul and Souirti, 2012; Gamboa-Vignolle et al., 2012; Gee et al., 2012; Gopalakrishnan et al., 2012; Hantous-Zannad et al., 2012; Herrak et al., 2012; Ilker et al., 2012; Khalatbari et al., 2012; Khoshnevisan et al., 2012; Lavrnjic et al., 2012; Macvanski et al., 2012; Malik et al., 2012; Mnejja et al., 2012; Moguchaia et al., 2012; Naydenov et al., 2012a; Naydenov et al., 2012b; Nunez-Farias et al., 2012; Okechi and Albright, 2012; Ortiz et al., 2012a; Ortiz et al., 2012b; Ouni et al., 2012; Pigac et al., 2012; Qi et al., 2012; Ragab et al., 2012; Ren et al., 2012; Romero-Rojas et al., 2012; Saeedinia et al., 2012; Streletz et al., 2012; Svistov et al., 2012; Tanboon et al., 2012; Thota et al., 2012; Todua and Chedia, 2012; Topkan et al., 2012; Zhai et al., 2012; Zhang et al., 2012; Anis et al., 2013; Beizig et al., 2013; Buglyo et al., 2013; Bunevicius et al., 2013a; Bunevicius et al., 2013b; Casas Parera et al., 2013; Cortez-Conradis et al., 2013; Gigineishvili et al., 2013; Hafiz et al., 2013; Hernandez-Martinez et al., 2013; Hilmani et al., 2013; Jaimovich et al., 2013; Jain et al., 2013; Jazayeri et al., 2013; Khalatbari et al., 2013; Khamlichi et al., 2013; Khan et al., 2013; Naidoo et al., 2013; Piyapittayanan et al., 2013; Rabah et al., 2013; Rahman et al., 2013; Romero-Rojas et al., 2013a; Romero-Rojas et al., 2013b; Romero-Rojas et al., 2013c; Saglam et al., 2013; Tuntiyatorn et al., 2013; Yildirim et al., 2013), all of which had papers classifying CNS tumors into WHO classification. However, a fewer cases (7910 cases) were reported in PubMed during the corresponding period. Bulgaria (2781 cases, 35.1%), Iran (1582 cases, 20.0%) and China (685 cases, 8.6%) were the top contributing countries identified in the PubMed search.

Predominant tumor types reported in individual countries

The most common reported tumor types for each country included in this study were identified, (Table 1). As expected, the majority of countries reported CNS tumors mainly as unclassified in their cancer registries.

Table 2. Individually Classified CNS Tumor Cases Reported Via Cancer Registries and PubMed Publications

Types of Reported CNS Tumors	Reported Cases in Cancer Registries		Reported Cases in PubMed	
	Total (%)	Average (STD DEV (+/-))	Total (%)	Average (STD DEV (+/-))
Unclassified CNS Tumors	30311 (96.1)	1443.3 (493.9)	3612 (45.6)	95.0 (67.5)
Astrocytic Tumors	627 (1.99)	29.8 (14.4)	1245 (15.7)	32.7 (15.8)
Pituitary Tumors	NR	NR	792 (10)	20.8 (9)
Meningeal Tumors	100 (0.32)	4.76 (2.51)	544 (6.88)	14.3 (6.88)
Gliomas (Not WHO Classified)	154 (0.49)	7.33 (4.13)	420 (5.31)	11.0 (4.55)
Oligodendroglial Tumors	78 (0.25)	3.71 (2)	355 (4.49)	9.34 (5.74)
Embryonal Tumors	107 (0.34)	5.1 (2.64)	198 (2.5)	5.21 (2.51)
Oligoastrocytic Tumors	NR	NR	191 (2.41)	5.03 (4.47)
Ependymal Tumors	91 (0.29)	4.33 (2.18)	138 (1.74)	3.63 (2.26)
Metastatic Tumors	NR	NR	117 (1.48)	3.08 (1.85)
Tumors of Cranial & Paraspinal Nerves	1 (0)	0.05 (0.05)	82 (1.04)	2.16 (1.19)
Tumors of the Sellar Region	NR	NR	64 (0.81)	1.68 (0.86)
Other Neuroepithelial Tumors	36 (0.11)	1.71 (0.74)	43 (0.54)	1.13 (1.08)
CNS Lymphoma	14 (0.04)	0.67 (0.48)	41 (0.52)	1.08 (0.87)
Mesenchymal Tumors	5 (0.02)	0.24 (0.14)	25 (0.32)	0.66 (0.27)
Neuronal and Mixed Neuronal-glial Tumors	2 (0.01)	0.1 (0.1)	21 (0.27)	0.55 (0.24)
Pineal Tumors	2 (0.01)	0.1 (0.1)	11 (0.14)	0.29 (0.12)
Germ Cell Tumors	NR	NR	8 (0.1)	0.21 (0.08)
Choroid Plexus Tumors	2 (0.01)	0.1 (0.1)	3 (0.04)	0.08 (0.04)
Total	31530 (100)	1501.4 (487)	7910 (100)	208.15 (83.48)

The exceptions for this were Thailand (63 cases, 53.8%), Jordan (98 cases, 57%), Saudi Arabia (116 cases, 41.3%) and Pakistan (277 cases, 53.5%), who reported Astrocytic tumors as the most frequent type. For data recovered from PubMed, Astrocytic tumors predominated Bosnia and Herzegovina (16 cases, 53.3%), Croatia (40 cases, 70.2%), Hungary (14 cases, 42.4%), Morocco (34 cases, 43%) and India (66 cases, 94.3%). Meningeal tumors predominated China (180 cases, 26.3%), Lithuania (117 cases, 33.6%) and Bangladesh (8 cases, 88.9%). Gliomas (Not WHO Classified) tumors were most frequent in Thailand (61 cases, 69.3%), Latvia (45 cases, 100%), and Egypt (60 cases, 36.1%). Pituitary Tumors were the most reported type in Romania (142 cases, 84.5%), Turkey (146 cases, 60.3%), Chile (3 cases, 50%), Colombia (47 cases, 79.7%) and Pakistan (282 cases, 75.0%), while Embryonal tumors were the most published type in Brazil (37 cases, 97.4%) and South Africa (2 cases, 50%) and Ependymal tumors were most common in Poland (39 cases, 53.4%).

CNS tumor cases reported via cancer registries

Percentage ordered cases for all reported histopathologically distinguished types of CNS tumors identified through cancer registries showed an overwhelming majority of reported cases to be WHO unclassified (30311 cases, 96.1%), (Table 2). The data for unclassified tumor cases showed a huge variation of reported cases between countries, ranging between a 25-percentile of 77 to a 75-percentile of 2123 and a standard deviation of ± 493.0 . The top three reported types were astrocytic tumors (627 cases, 1.99%), gliomas (not who classified) (154 cases, 0.49%) and embryonal tumors (107 cases, 0.34%). the least reported types were tumors of cranial and paraspinal nerves (1 case, 0.05%). Only 12 WHO classified categories were noted.

CNS tumor cases reported via pubmed

For CNS tumor cases identified via PubMed publications 17 WHO CNS tumor categories were noted, (Table 2). Collected data for all tumor cases showed a tighter variation of reported cases between countries ranging between a 25-percentile of 5.3 to a 75-percentile of 155 and a standard deviation of ± 83.48 . Under half of cases (3612 cases, 45.6%) were unclassified and were labeled as brain and/or CNS tumors. The top three reported types were astrocytic tumors (1245 cases, 15.7%), pituitary tumors (792 cases, 10.0%) and meningeal tumors (544 cases, 6.88%). the least reported types were choroid plexus tumors (3 cases, 0.04%).

Latitude as a possible risk factor for CNS tumors in developing countries

Several environmental risk factors have been associated with the epidemiology of CNS tumors including ionizing radiation (dental X-rays, CT-scans) (Braganza et al., 2012; Claus et al., 2012; Isaacs, 2013), bisphenol (Zhu et al., 2010), N-nitroso compounds (Huncharek, 2010), pesticides (Searles Nielsen et al., 2010), reproductive hormones (Michaud et al., 2010), JC-virus (Noch et al., 2012), human herpes-viruses (Kofman et al., 2011), and ELF-EMF exposure (Kheifets et al., 2010; Baldi et al.,

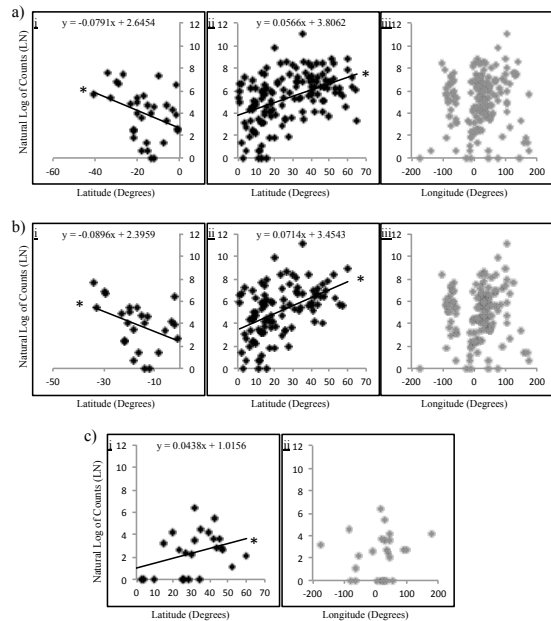


Figure 2. Scatter Blots for the Natural Logs of CNS Tumors Counts Verses Geographical Coordinates of WHO Registered Countries. a) The natural logs of Globocan reported incidence ASR (Ln) for all CNS tumors from 169 countries versus i) Latitudes of countries south the equator, (-40 to 0°), ii) Latitudes of countries north the equator, (0 to 70°), or iii) Longitudes. b) The natural logs of Globocan reported incidence ASR (Ln) for all CNS tumors from 126 developing countries versus i) Latitudes of countries south the equator, (-40 to 0°), ii) Latitudes of countries north the equator, (0 to 70°), or iii) Longitudes. c) The natural logs counts of Astrocytic tumor collected from 25 developing countries versus i) Latitudes or ii) Longitudes. Equations for linear regression curves are shown above corresponding blots. The (*) indicates a significant linear regression relationship

2011). Previous work had shown an association for the occurrence of melanoma (Crocetti et al., 2012), colorectal cancers (Cuomo et al., 2013), lymphoid neoplasms (van Leeuwen et al., 2013) and others (Grant, 2012) with geographical parameters. A linear regression analysis showed a significant relationship between the natural logs of CNS tumors ASR data sourced by Globocan for 169 WHO registered countries and latitudes ranging from -40 till 0 degrees ($F(2, 32)=4.66, p=0.03$) (Figure 2ai), and for latitudes ranging from 0 till 70 degrees ($F(2, 137)=33.7, p=0.00$) (Figure 2a ii). The analysis suggested that the counts of CNS tumor cases increased as the coordinates move away from the equator (Degree 0) in both directions. In contrast, there was no significant relationship observed between the natural logs of CNS tumor incidence ASR of all countries and longitude ($F(2, 169)=1.04, p=0.30$) (Figure 2a iii). A similar pattern was observed for data representing developing countries, where a significant linear regression relationship was observed between the natural logs of CNS tumors ASR data sourced by Globocan for 126 WHO registered developing countries and latitudes ranging from -40 till 0 degrees ($F(2, 25)=4.40, p=0.04$) (Figure 2bi), and for latitudes ranging from 0 till 70 degrees ($F(2, 101)=27.5, p=0.00$) (Figure 2b ii). No significant relationship was observed between the natural logs of CNS tumors ASR

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data and longitudes of developing countries (Figure 2biii). Analysis of Astrocytic tumors counts sourced by PubMed showed a comparable pattern (Figure 2c), where natural logs of counts had a significant linear regression relationship with latitudes ranging from 0 till 70 degrees ($F(2, 25)=5.69, p=0.02$), (Figure 2ci). There was no significant relationship between the natural logs of Astrocytic tumor counts and longitude ($F(2, 25)=0.51, p=0.48$) (Figure 2cii).

Discussion

This study aimed to show epidemiological trends of individually reported CNS tumors subtypes in developing countries. Through a cyber-search approach, it was possible to elucidate trends of frequencies for individually classified CNS tumors and show patterns that were otherwise unavailable. Our data shows that the majority of reported cases from PubMed and cancer registries were for unclassified tumors. For classified tumors, Astrocytic tumors appear to be the most reported type. Interestingly, a relationship between geographical location and the occurrence of CNS tumors may exist.

Unfortunately, out of 149 developing countries only 66 registries representing 21 countries reported any information related to CNS tumors, confirming previous observations for the lack of reporting for CNS tumors in the developing region (Ferlay et al., 2010; Qaddoumi et al., 2011). The degree of reporting for tumors maybe linked to the level of socio-economic development for any particular country. The quality of data collected from the identified registries was poor compared to data retrieved thought published resources. We ensured that included PubMed publications were sourced by clearly individual hospitals or health sectors, and did not produce repetition or overlap of cases. Importantly, data produced from PubMed showed a larger range of tumor classification that was not detected in cancer registries. One problem was the lack of gender or age categorization for declared tumors in publications. Another caveat of retrieving data from such source is the hypothetical bias of results towards research interest in a particular type of tumors. However, our method of selecting published papers was intended to focus on country related data and not any particular tumor classification.

Alarmingly, the majority of reported cases were unclassified (PubMed (45.6%), registries (96.1%)) consistent with the notion that reporting unclassified cancer variants is a growing phenomenon (Tavtigian et al., 2008). This could be a reflection of routed miscommunication between clinicians and statistical departments. It is possible that the epidemiological context for each of these bodies is different. While clinicians and pathologists are more concerned with understanding disease classifications to make crucial decisions involving management of treatment (Tavtigian et al., 2008), reporting statistical departments may be more concerned with “number of cases” in general. In addition, reporting classified tumors is more complex and involves extra documentations, such as pathological reports for classified tumors (Jensen and Storm, 1991). Unfortunately, such vague reporting

systems hamper sharing of detailed medical knowledge and perhaps hinder the development of effective targeted medical management programs.

The retrieved data from PubMed shows that the top three reported types of CNS tumors were Astrocytic tumors, Pituitary tumors and Meningeal tumors. Reasons for the high frequency of Astrocytic tumors are generally not explained (Karipidis et al., 2007). It is possible that since Astrocytic tumors have devastating outcomes with poor survival, more researchers are interested to study them and thus there is more reporting in PubMed, however, Astrocytic tumors were also the most frequently reported type in cancer registries. Another consideration is the complexity of classification for astrocytic tumors, as several different histological categories of variants exist (Wen and Kesari, 2008; Adesina et al., 2010; Karsy et al., 2012) raising the possibility of effect due to increased size of grouping.

Interestingly, when analyzing the relationship between the natural logs of CNS tumors incidence ASR for all WHO registered countries including developing countries verses latitude, or for the natural logs of counts for Astrocytic tumors verses latitude, positive correlations emerged. A similar association has been observed for other tumors (Crocetti et al., 2012; Grant, 2012; Cuomo et al., 2013; van Leeuwen et al., 2013). Latitude effect may be associated with several factors such as UV irradiation, Vitamin D biosynthesis or geographical differences for regional health systems.

In conclusion, reporting epidemiological data for unclassified CNS tumors is a growing phenomenon in the developing world that contradicts the attitude of health professionals to deal with CNS tumors as a collection of independent tumors, of which each requires a targeted health management plan. Unfortunately, Globocan reports treated epidemiological data for CNS tumors within a single category, and only a few cancer registries for developing countries provided information related to the incidences of defined histological CNS tumors types. Through the retrieval of CNS tumor cases using ISI PubMed publications, overall frequencies for individually classified CNS tumors in 38 different countries were elucidated. Patterns associated with frequencies and ranking of classified tumors were deduced as well as novel data implicating an association between individual CNS tumors, such as astrocytic tumors, with geographical location. A consensus for future reporting of CNS tumors incidences need to be established in order to help progress prospects of CNS tumors' health management.

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