Incidence Trends in the Anatomic Location of Primary Malignant Brain Tumors in the United States: 1992–2006

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Key words

- Astrocytoma
- Brain neoplasm
- Epidemiology
- Glioma
- Glioblastoma multiforme
- Incidence
- Location

Abbreviations and Acronyms

AA: Anaplastic astrocytoma
AAIR: Age-adjusted incidence rates
APC: Annual percent changes
CCR: California Cancer Registry
CNS: Central nervous system
GBM: Glioblastoma multiforme
ICD0-3: International Classification of Disease for
Oncology, Third Edition
LAC: Los Angeles County Cancer Surveillance
Program
SEER: National Cancer Institute's Surveillance,
Epidemiology, and End Results
WHO: World Health Organization

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INTRODUCTION

Over the last 3 decades, several populationbased studies have reported an overall increase in the incidence of malignant primary brain tumors (I, 4, 6-9, II, I4). Although it has been generally accepted that this phenomenon is at least in part accounted for by higher detection rates associated with the increasing frequency and sensitivity of diagnostic imaging, it remains to be determined whether the true incidence of primary central nervous system (CNS) tumors is independently increasing as a result of environmental factors (I, II, I8). BACKGROUND: This study sought to determine incidence trends of the anatomical origin of primary malignant brain tumors.

METHODS: Incidence data for histologically confirmed brain tumors were obtained from the Los Angeles County Cancer Surveillance Program (LAC), the California Cancer Registry (CCR), and the National Cancer Institute's Surveillance, Epidemiology, and End Results (SEER) program for 1992 to 2006. Ageadjusted incidence rates (AAIR) and annual percent changes (APC) were calculated by histologic subtypes and anatomic subsites. Statistical analyses were performed using the SEER*Stat analytic software and SAS statistical software.

■ RESULTS: Increased AAIRs of frontal (APC +2.4% to +3.0%, $P \le 0.001$) and temporal (APC +1.3% to +2.3%, $P \le 0.027$) lobe glioblastoma multiforme (GBM) tumors were observed across all registries, accompanied by decreased AAIRs in overlapping region GBMs (-2.0% to -2.8% APC, $P \le 0.015$). The AAIRs of GBMs in the parietal and occipital lobes remained stable. The AAIR of cerebellar GBMs increased according to CCR (APC +11.9%, P < 0.001). The AAIR of all gliomas, which includes all anatomical subsites, decreased (-0.5% to -0.8% APC, $P \le$ 0.034). Low-grade and anaplastic astrocytomas demonstrated decreased AAIRs in the majority of brain regions.

CONCLUSIONS: Data from 3 major cancer registries demonstrate increased incidences of GBMs in the frontal lobe, temporal lobe, and cerebellum, despite decreased incidences in other brain regions. Although this may represent an effect of diagnostic bias, the incidence of both large and small tumors increased in these regions. The cause of these observed trends is unknown.

Although many previous reports have analyzed trends in the overall incidence of gliomas and various glioma subtypes, few recent studies have examined these trends according to the anatomical subsites of primary malignant brain tumors over the last several decades (2, 10). Furthermore, no recent studies have analyzed populationbased incidence trends by both tumor grade subtype and anatomic location. Given the increasing trends of primary malignant brain tumors, we sought to determine whether any notable trends in the anatomical topography of primary CNS tumors have occurred. Data from 3 major populationbased cancer registries were reviewed to identify any trends in the incidence of primary malignant brain tumors, their location of origin, and various demographic risk factors. The current study is the first to analyze population-based incidence trends of malignant brain tumors according to anatomical parameters.

CLINICAL MATERIALS AND METHODS

Data used in our analysis were obtained from 3 sources, the largest of which is the National Cancer Institute's Surveillance, Epidemiology, and End Results (SEER) Program (16). A signed limited-use data agreement was obtained to access these data. This program includes incidence and population data associated by age, gender, race/ ethnicity, and year of diagnosis. Thirteen GABRIEL ZADA ET AL.

ANATOMIC LOCATION OF PRIMARY MALIGNANT BRAIN TUMORS

Table 1. Demographic Characteristics of Patients Harboring Gliomas in 3 Major Tumor Registries, 1992 to 2006						
	LA County		CCR		SEER 12 Registries	
	Number	%	Number	%	Number	%
Total number	5736	100.0	10,412	100.0	22,419	100.0
Sex						
Male	3154	55.0	5864	56.3	12,651	56.4
Female	2582	45.0	4548	43.7	9768	43.6
Age						
0–19	476	8.3	711	6.8	1477	6.6
20–64	3141	54.8	5880	56.5	12,386	55.2
65+	2119	36.9	3821	36.7	8556	38.2
Ethnicity						
Non-Hispanic white	3410	59.4	6776	65.1	18,658	83.2
Hispanic white	1514	26.4	2057	19.8	1288	5.7
Black	402	7.0	603	5.8	1161	5.2
Asian/other	410	7.1	976	9.4	1312	5.9
Location						
Frontal lobe	1434	25.0	2653	25.5	5781	25.8
Temporal lobe	1045	18.2	1938	18.6	4548	20.3
Parietal lobe	766	13.4	1449	13.9	3125	13.9
Occipital lobe	147	2.6	291	2.8	762	3.4
Overlapping	1238	21.6	2238	21.5	3897	17.4
Cerebellum	124	2.2	203	1.9	344	1.5
Brainstem	249	4.3	437	4.2	934	4.2
Ventricle	97	1.7	151	1.5	263	1.2
Cerebrum	277	4.8	470	4.5	1129	5.0
Brain NOS	359	6.3	582	5.6	1636	7.3
Histology						
Astrocytoma, NOS (WHO I and II)	706	12.3	1091	10.5	2260	10.1
Anaplastic astrocytoma (WHO III)	599	10.4	993	9.5	1792	8.0
GBM (WHO IV)	3094	53.9	5868	56.4	12,714	56.7
Protoplasmic/fibrillary Astrocytoma	87	1.5	137	1.3	421	1.9
Unique astrocytoma variants	34	0.6	48	0.5	96	0.4
Ependymoma	172	3.0	298	2.9	563	2.5
Mixed glioma	226	3.9	412	4.0	720	3.2
Glioma, NOS	369	6.4	662	6.4	1570	7.0
Oligodendroglioma	325	5.7	653	6.3	1714	7.6
Anaplastic oligodendroglioma	124	2.2	250	2.4	569	2.5
Laterality						
lpsilateral	5673	98.9	10325	99.2	22,291	99.4
Bilateral	35	0.6	48	0.5	64	0.3
Paired, unknown laterality	28	0.5	39	0.4	64	0.3
CCR, California Cancer Registry; GBM, glioblastoma multiforme; LA, Los Angeles County; NOS, not otherwise specified; SEER, National Cancer Institute's Surveillance, Epidemiology, and End Results; WHO, World Health Organization.						

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