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# STATISTICAL ANALYSIS OF MOBILE RADIATION AND IMPACT ON HEALTH: A CASE STUDY ON OCCURRENCE OF BRAIN TUMORS

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#### Objective

To analyze whether mobile phone radiation is one of the source for brain tumor. This study is carried out by computing odd ratios.

### Abstract

The mobile radiation seems to be severe among children than the adults because their skulls are tiny and thinner. Mobile phones emit radiations and RF-EMF is considered as a Group 2B (carcinogen) and the survey suggests that 10-year latency period is practically enough for the development of tumors. The studies from various groups involve cases of 35487 and controls about 82609. The results for meta- analysis of glioma gave an odd ratio OR=1.10, 95% CI=0.79-1.54 and for the latency period  $\geq 10$  years, OR=1.38, 95% CI=0.70-2.73. The highest risk was found in the age group below 20 years from the Hardell Group.

Keywords:- exposure, mobiles, radiation, risk

#### INTRODUCTION

Intense use of wireless communication has raised a concern over the human health, an increased risk for brain tumors. Mobile phones and cordless phones emit radiofrequency (RF) radiation when in use. The minimum period was  $\geq 10$  years or  $\geq 1640$  hours for the development of tumors using wireless phones with the higher the exposure on the same side of the brain (ipsilateral) whereas the contralateral side is exposed less [1].

In May 2011, an approach for the scientific evidence of the risk for brain tumor was completed by the International Agency for Research on Cancer (IARC) at the World Health Organization (WHO). IARC is independently funded and has categorized the extremely low frequency (ELF) as a Group 2B carcinogen.

The radio frequency energy emitted by mobile phones can be absorbed by brain tissues near to the mobile; is a form of non-ionizing electromagnetic radiation. The brand of mobile phone, the distance between the phone and user and user's distance from cell phone towers are the key factors in the amount of radio frequency energy a mobile user is exposed. In 1998, the International Commission on Non-Ionizing Radiation Protection (ICNIRP) has established the exposure guidelines and short-term thermal effect from RF radiation [2].

#### **Child's Brain Endanger**

A child is in danger when the child holds the mobile phone, I-pad or any radiating device, as a child's brain can absorb higher rates than adults due to the smaller head size, thinner skull bones and higher brain conductivity. The study reveals that most of the children below two years are left with mobiles, i-pad or any other electronic gadget because the parents are busy in their jobs and regular routine work. Henceforth, children are at higher risk of exposure to carcinogens than adults, and the smaller the child the greater the risk [3] [4]. There are several studies that prove that children absorb more Micro Wave Radiation (MWR) than adults. In 1996, one of the study concluded that the penetration of absorbed MWR was deeper into the children's brain of age 5 to 10 years [5] [6]. In 2008, a French senior researcher named Joe Wiart, pointed that the child's brain tissue absorbed two times more MWR than adults [7]. In 2009, the study stated that the Central Nervous System absorption in children's is higher ( $\sim$ 2x) as the MWR source is closer and skin and bone layers are thinner and bone marrow exposure ( $\sim$ 10x) varies with the age [8]. In 2010, Andreas Christ and his team described that the child's hippocampus and hypothalamus absorbs 1.6–3.1 times greater and the cerebellum absorbs 2.5 times higher MWR compared to adults; the bone marrow of child absorbs 10 times more MWR radiation than adults [9] [10].

#### **Mobile Phones SAR Evaluation**

The Specific Absorption Rate (SAR; W kg<sup>-1</sup>) depends on the type of phone used; one can calculate the SAR per number of hours exposed [11]. Based on the type of phone manufacturer, few phones give peak SAR value above the ear, some beneath the ear and few even below the ear. The SAR value will affect the human ear and later damage the human brain. And whose ears are affected with SAR cannot hear properly. Jonna Wilen has calculated mean values of SAR<sub>1g</sub> for a certain number of people who had a particular symptom (W–S) and those who do not have the symptom (WO–S). Jonna's study from her observation concluded that 30 out of 44 comparisons (W–S<sub>mean</sub>/WO–S<sub>mean</sub>) are greater than 1. The symptom discomfort is known from W–S=95, SAR<sub>1g</sub>=0.67 (0.23) and WO–S=2094,

 $SAR_{1g}=0.59$  (0.27), (W-S<sub>mean</sub> / WO-S<sub>mean</sub>) =1.14. The symptom concentration W-S =180,

 $SAR_{1g} = 0.65 (0.25)$  and WO-S = 2009,  $SAR_{1g} = 0.59 (0.27)$  and  $(W-S_{mean} / WO-S_{mean}) = 1.10$  below the ear (volume 3), the equivalent differences were larger than 10% [12]. The initial increase for brain tumor risk associated with mobile phones was issued before 10 years and was found for ipsilateral mobile phone use [14] [15].

#### Statistical Methodology

For data analysis, odds ratio was calculated using Review Manager 5.3. The random effects model was used to measure summary odds ratio, based on chi-square test, tau squared and I-squared statistics for heterogeneity as shown in figure 1 and figure 4 [19]. Latency is defined as the first year of use of wireless to the diagnosis year. Latency was analyzed using periods groups > 10 years, 10-15 years, 15-20 years and >25 years.

Based on Hardell group and Interphone group studies the meta-analyses were performed with the mobile phones use. The model was chosen based on the latency and the number of hours ( $\geq 10$  years and  $\geq 1640$  h) a mobile is used to test for heterogeneity. The total number cases 35487 and control 82609 evaluated for all 23 studies considered from the Hardell group and Interphone study group in the analysis of patients with glioma yielded OR =1.10 95% CI = 0.79-1.54 (p = 0.57). The relationship between the study qualities and odd ratios are shown in graphs in figure 2 and figure 3.

#### Odd Ratios for mobile use over 10 years +

Lonn et al and group, Karolinska Institute in Sweden conducted a study on glioma and meningioma cases. The mobile use of  $\geq$  10 years showed for ipsilateral glioma OR=1.6, 95% CI=0.8-3.4 for 15 cases and for contralateral glioma OR=0.7 95% CI=0.3-1.5 for 11 cases. Shoemaker et al (2010) of the INTERPHONE study showed the results for acoustic neuroma from six regions, the results for 678 cases of lifetime use ( $\geq$ 10 years) showed OR =1.8, 95% CI=1.1-3.1 for ipsilateral acoustic neuroma, and OR=0.9 95% CI=0.5-1.8 for contralateral tumor. The Danish of the INTERPHONE study for  $\geq$  10 years produced OR= 1.0 (0.3 to 3.2) for meningioma, low–grade glioma OR=0.5, 95% CI=0.2-1.3 and highgrade glioma OR=0.5, 95% CI=0.2-1.3 for 252 glioma cases, 175 meningioma cases with 822 controls. Hepworth et al, England showed results on glioma as a part of the

INTERPHONE study for ≥10 years, ipsilateral provided OR=1.6, 95% CI=0.9-2.8 and

	Cas	es	Cont	trol		Odds Ratio	Odds Ratio
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Random, 95% CI	M-H, Random, 95% CI
1.1.1 Study Quality 1							5
Aydin 2011	546	688	975	1375	4.4%	1.58 [1.27, 1.96]	
schuz 2006	242	747	517	1494	4.4%	0.91 [0.75, 1.09]	+
INTERPHONE STUDY 2011	1121	1361	7658	14354	4.4%	4.08 [3.54, 4.71]	
INTERPHONE STUDY 2010	5190	7416	7658	14354	4.5%	2.04 [1.92, 2.16]	
Subtotal (95% CI)		10212		31577	17.7%	1.86 [1.12, 3.11]	◆
Total events	7099		16808				
Heterogeneity: Tau <sup>2</sup> = 0.27; Chi <sup>2</sup> = 168.66, df = 3 (P < 0 Test for overall effect: Z = 2.38 (P = 0.02)	0.00001);	l² = 98%					
1.1.2 study Quality 2							
Hardell 2007	101	120	339	647	4.0%	4.83 [2.89, 8.07]	
Hardell 2009	460	531	1266	1920	4.4%	3.35 [2.56, 4.37]	
Hardell 2009 (Incidence of brain Tumor)	1254	1425	2162	2429	4.4%	0.91 [0.74, 1.11]	+
Hardell 2011	1251	1472	2438	2900	4.4%	1.07 10.90, 1.281	+
Subtotal (95% CI)		3548		7896	17.2%	1.94 [0.98, 3.83]	-
Total events	3066		6205				
Heterogeneity Tau <sup>2</sup> = 0.46: Chi <sup>2</sup> = 90.56 df = 3 (P < 0.	000011: 12	= 97%	0200				
Test for overall effect: $Z = 1.90$ (P = 0.06)		0. %					
1.1.3 Study Quality 3							
Hardell 2014	1498	1691	3530	4038	4.4%	1.12 (0.94, 1.33)	+
Carlberg 2015	2068	2349	3530	4038	4 4 %	1 06 0 91 1 241	+
M Carlberg 2017 (Evaluation of Glioma Risk)	732	2708	1279	2972	4 5%	0 49 10 44 0 551	÷
M Carlberg 2017	1346	1939	3485	5404	4 5%	1 25 [1 12 1 40]	-
Subtotal (95% CI)	1010	8687	0.00	16452	17.8%	0.92 [0.57, 1.49]	-
Total events	5644		11874				1 100 <b>-</b>
Heterogeneity: Tau <sup>2</sup> = 0.24; Chi <sup>2</sup> = 155.71, df = 3 (P < 0 Test for overall effect: $Z = 0.33$ (P = 0.74)	0.00001);	l² = 98%					
A 11							
1.1.4 Study Quality 4							0
Hardell 2002	588	649	581	649	4.3%	1.13 [0.78, 1.62]	
Hardell 1999	78	209	161	425	4.3%	0.98 [0.69, 1.37]	
Hardell 2010	346	464	619	927	4.4%	1.46 [1.14, 1.87]	
coureau 2014	222	447	443	892	4.4%	1.00 [0.80, 1.26]	
Hardell 2005	2437	3729	2162	2437	4.4%	0.24 [0.21, 0.28]	
Subtotal (95% CI)		5498		5330	21.7%	0.82 [0.36, 1.90]	
Total events	3671		3966				
Heterogeneity: Tau <sup>2</sup> = 0.90; Chi <sup>2</sup> = 242.00, df = 4 (P < 0 Test for overall effect: $Z = 0.46$ (P = 0.65)	0.00001);	I <sup>2</sup> = 98%					
1.1.5 Study Quality 5							
Warren 2003	21	51	53	141	3.8%	1 16 10 60, 2 231	· · · · ·
Muscat 2000	66	469	76	422	4.3%	0.75 (0.52, 1.07)	
Inskin 2001	308	782	358	798	4 4 %	0.80 (0.65, 0.98)	
Subtotal (95% CI)		1302		1361	12.5%	0.81 [0.68, 0.96]	•
Total events	395		487				
Heterogeneity: Tau <sup>2</sup> = 0.00; Chi <sup>2</sup> = 1.39; df = 2 (P = 0.5	0): $ ^2 = 0\%$						
Test for overall effect: Z = 2.49 (P = 0.01)							
1.1.6 Study Quality 6							
Hardell 2013	316	338	3530	4038	4.2%	2.07 [1.33, 3.22]	· · · ·
Hardell 2013 (Brain tumor between 2007 and 2009)	593	1601	1368	1601	4 4 %	0 10 10 08 0 1 21	<u>+</u>
Hardell 2012	2708	4301	7658	14354	4.5%	1 49 [1 39 1 59]	· · · · · · · · · · · · · · · · · · ·
Subtotal (95% CI)		6240		19993	13.0%	0.67 [0.09, 5.13]	
Total events	3617		12556				
Heterogeneity Tau <sup>2</sup> = 3.21; Chi <sup>2</sup> = 825.13; df = 2 /P < 1	1 000011	P= 1009	6				
Test for overall effect: $Z = 0.38$ (P = 0.70)							
Total (95% CI)		35487		82609	100.0%	1.10 [0.79, 1.54]	•
Total events	23492		51846				
Heterogeneity: Tau <sup>2</sup> = 0.64; Chi <sup>2</sup> = 2307,45; df = 22 (P	< 0.0000	1);   <sup>2</sup> = 99	196				
Test for overall effect: Z = 0.57 (P = 0.57)							U.U1 U.1 1 10 100
Test for subgroup differences: Chi <sup>2</sup> = 14.34, df = 5 (P =	= 0.01), P	= 65.1%					RISK DECIERSES RISK INCRESSES

Fig 1. Odds Ratio for the Brain Tumor risk is calculated using Review Manager 5.3

Contralateral OR=0.8, 95% CI=0.4- 1.4 for 966 cases [1]. The INTERPHONE study group (2011) pointed that the time a mobile used for  $\geq$ 10 years OR=0.83, 95% CI=0.58–1.19 for 10 years after regular use and for the hours  $\geq$ 1640 h of cumulative call time it was 2.79 (1.51–5.16) [20]. Hardell (2013), the study on the cumulative use of wireless phones in different quartiles, the quartile > 1486 h gave OR=2.6, 95% CI=1.5-4.4 for mobile phone and wireless phone use produced OR=2.2, 95% CI=1.5-3.4 [22]. Hardell (2012), analyzed that using a cordless phone gave an increased highest risk in diagnosing of glioma in the latency group >10 years OR =3.8, 95% CI=1.8-8.1 and for glioma with the mobile use  $\geq$ 1640 h; the increased risk in the temporal lobe giving OR =1.87, 95% CI=1.09-3.22 from Table 1 [23].

Carlberg (2017) concluded that the longer latency of 10+ years OR = 1.62, 95% CI=1.20–2.19. The three studies revealed a consistent increase of glioma risk with latency. The highest OR with longest latency of 10+ years with 732 exposed cases and 1,279 exposed controls [24]. Carlberg (2015) analyzed that the cumulative use > 3358 hours relates to >90<sup>th</sup> percentile. An increased risk for mobile phone (2G and 3G) with OR=1.5, 95% CI=1.0005-2.3(p-trend=0.045) and cordless phone showed OR=2.1, 95% CI=0.7-6.3 (ptrend=0.65) [25].

Hardell (2011), the peak cumulative use of mobile phones  $\geq$  1640 h for glioma OR=1.40 95% CI=1.03-1.89 was calculated. The risk increased for ipsilateral use OR=1.96, 95% CI =1.22-3.16. In temporal lobe, the highest risk was found with more exposure in the anatomical area [26].

#### Table 1: Summary of studies of wireless phones usage and glioma risk, Hardell (2012)

Study	Cases/events	OR 95%CI	Comments
Hardell 2013 (1997- 2003)	1148/57	2.9(1.8-4.7)	Latency period > 10 years, mobile phone (ipsilateral)
	1148/20	3.8(1.8-8.1)	Latency period >10 years, cordless phone (ipsilateral)
Interphone study group 2010 (2000- 2004)	2708/210	1.40(1.03- 1.89)	Mobile phone Cumulative hours ≥ 1640 h
	2708/78	1.87(1.09- 3.22)	MobilephonecumulativeHours $\geq$ 1640 h, tumor in thetemporal lobe
	2708/100	1.96(1.22- 3.16)	Ipsilateral phone usage cumulative hours $\geq$ 1640h



## Fig: The relationship graph between logs odd ratio and study quality



Fig. 4 Output using mobile phones or cordless phones for more than 10 years

#### Findings

Carlberg (2017) examined a largest case-control study on brain tumors and occupational exposure to ELF-EMF. The INTEROCC study proved a relation between exposure to ELFEMF and glioma and finally concluded that the final phases of astrocytoma grade IV for occupational ELF-EMF exposure had an increased risk [18]. Carlberg (2015) concluded a fact that the occupational ELF-EMF exposure has an increased risk of glioma. Bradford Hill's viewpoints on association on RF radiation and glioma risk. Hence concluded that glioma is caused by RF radiation [24].

Hardell (2014), the questionnaire was answered by 1498 of 1691 cases, of whom 879 were men and 619 women, of 4038 controls, 1492 men and 2038 women participated to give the total of 3530. The glioma risk at various age groups for the wireless phone use was found to be increasing. The risk increased in both mobile and cordless phones and OR is high before the age of 20 years. Children are more exposed to RF-EMF than adults as higher conductivity in the brain tissue and a smaller head [17]. Hardell (2013) concluded that the volume of tumor increased for ipsilateral use of mobile phones of the digital 2G type and for cordless phones per year of latency. Interphone Study (2010) the glioma cases diagnosed were 23% of 2708 cases before the age of 40.

Hardell (2009) suggested that a consistent association between use of mobile or cordless phones and astrocytoma grade I - IV and acoustic neuroma. For latency period >10 years, the risk was highest for ipsilateral exposure to microwaves. The greater risk for persons who started to use mobiles before the age of 20 years were identified in Sweden during 2000 - 2007 and the results were supported by the increase of incidence of astrocytoma [27]. Hardell (1999) revealed a non – significant increase risk for brain tumors located in the temporal or occipital lobe was identified for those who used cell phones on the same side of the head. Acoustic neuroma develops with higher exposure to microwave radiation from a mobile phone [28].

	Cases		Contr	ol	Odds Ratio
Study or Subgroup	Events	Total	Events	Total	M-H, Fixed, 95% Cl
1.1.1 Analogue					
Total 1 year	86	316	558	3530	1.99 [1.53, 2.59]
Total 1-5 year	16	316	87	3530	2.11 [1.22, 3.64]
Total 10-15 years	16	316	113	3530	1.61 [0.94, 2.76]
Total 15-20 years	9	316	107	3530	0.94 [0.47, 1.87]
Total 5-10 year	33	316	137	3530	2.89 [1.94, 4.30]
Total above 20 years	12	316	114	3530	1.18 [0.65, 2.17]
1.1.2 Mobile Phone					
Total 1 year	200	316	2148	3530	1.11 [0.87, 1.41]
Total 1-5 year	65	316	674	3530	1.10 [0.82, 1.46]
Total 10-15 years	34	316	476	3530	0.77 [0.54, 1.12]
Total 15-20 years	12	316	196	3530	0.67 [0.37, 1.22]
Total 5-10 year	77	316	688	3530	1.33 [1.02, 1.74]
Total above 20 years	12	316	114	3530	1.18 [0.65, 2.17]
1.1.3 Cordless Phone					
Total 1 year	156	316	1724	3530	1.02 [0.81, 1.29]
Total 1-5 year	72	316	653	3530	1.30 [0.99, 1.71]
Total 10-15 years	19	316	294	3530	0.70 [0.44, 1.14]
Total 15-20 years	2	316	109	3530	0.20 [0.05, 0.81]
Total 5-10 year	60	316	655	3530	1.03 [0.77, 1.38]
Total above 20 years	3	316	13	3530	2.59 [0.74, 9.15]
1.1.4 Digital Type					
Total 1 year	216	316	2393	3530	1.03 [0.80, 1.31]
Total 1-5 year	93	316	796	3530	1.43 [1.11, 1.85]
Total 10-15 years	38	316	584	3530	0.69 [0.49, 0.98]
Total 15-20 years	9	316	242	3530	0.40 [0.20, 0.78]
Total 5-10 year	73	316	758	3530	1.10 [0.84, 1.44]
Total above 20 years	3	316	13	3530	2.59 [0.74, 9.15]
1.1.5 Wireless Phone					
Total 1 year	227	316	2472	3530	1.09 [0.85, 1.41]
Total 1-5 year	72	316	748	3530	1.10 [0.83, 1.44]
Total 10-15 years	44	316	578	3530	0.83 [0.59, 1.15]
Total 15-20 years	13	316	253	3530	0.56 [0.31, 0.98]
Total 5-10 year	84	316	767	3530	1.30 [1.00, 1.70]
Total above 20 years	14	316	126	3530	1.25 [0.71, 2.20]

#### **Comparative Analysis**

The comparison were done using the results obtained by the calculation carried out using review manager and the respective groups results as follows Inskip 2001, reported that the relative risk of gliomas RR=0.9, 95% CI=0.7-1.1 with the cases=308 and controls=358. According to figure 1, Inskip 2001, the relative risk RR=0.88, 95% CI=0.78- 0.99 data was not shown in the figure [21]. Hardell (2014), the evaluated results of glioma (n=1380) and use of mobile and cordless phones in different latency group as >10 years, > 15-20 years, > 20-25 years, and > 25 years.

The cordless phone use gave OR=1.65, 95% CI=1.32-2.05 in the latency group > 15-20 years. The digital type 2G, 3G and cordless showed OR=1.54, 95% CI=1.34-1.78 in the latency group >10 years for wireless phones. The OR=1.35, 95% CI=0.51-3.58 for cordless for latency >20-25 years. The increased risk for wireless phones is given by OR=1.032, 95% CI=1.019-1.046 [17].

Fig 5.ORs for acoustic without any adjustments for age/gender/year of diagnosis

Hardell (2012) pointed that the OR=0.81, 95% CI= 0.70 - 0.94 for mobile phone use > 1 year for glioma and further risk increased for glioma in the temporal lobe yielding OR=1.87, 95% CI=1.09-3.22, the results from figure 1 reflects OR=1.49, 95% CI=1.39-1.59. The regular use of mobile phone and cordless phone for latency period>10 years yielded OR=2.9, 95% CI=1.8-4.7 for ipsilateral use. It was found doubling of glioma risk for total wireless phone use with OR=2.1, 95% CI=1.6-2.8. The results from review manager showed OR=2.60, 95% CI=1.71-3.94 for more than 10 years [23]. Hardell (2007) showed for acoustics neuroma over the latency period >10 years OR=1.3, 95% CI=0.6-2.8 whereas from figure 1, the obtained OR=4.83, 95% CI=2.89-8.07[1]. The result of mobile phones, cordless phone and wireless phone with the OR result analysis for 316 cases and 3530 controls by Hardell (2013). From figure 5, mobile phone use for 5 to 10 years OR=1.33 95% CI=1.021.74, cordless phone OR=1.03, 95% CI=0.77-1.38 and wireless phone use shows OR=1.30, 95% CI=1.00-1.70.

Hardell group has shown with an adjustment of age, gender and year of diagnosis and the latency>5-10 years the OR=2.3, 95% CI=1.6-3.3 for mobile phone, for cordless phone OR=1.6 95% CI=1.1-2.5 and wireless phone OR=1.9, 95% CI=1.3-2.7 for the same number of cases and controls.

The output of odds ratio using Review Manager 5.3 evaluated for all the studies for the total number of cases 35487 and control 82609 from the Hardell group and Interphone study group for the analysis of patients with glioma yielded OR =1.10, 95% CI=0.79-1.54 (p=0.57) which on comparing to the odds ratio OR=1.03, 95% CI=0.92-1.14 (p=0.64) calculated by the group from AIIMS (Delhi) using Review Manager with total cases 12426 and controls 19334. It indicates significant increase of OR=0.07.

## Conclusions

Results from the above various case- control studies on brain tumors and mobile phone usage for above 10 years suggests that there is an increased risk in glioma. The adults are to be aware when their children are left with mobile or any electronic gadget for entertainment, that there is risk for the child's health. The overall result of meta-analysis showed a significant increase of 1.38 times OR in brain tumor risk.

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## **Conflicts of Interest**

The authors declare that there are no conflicts of interest regarding the publication of this paper.

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