

# Brain Tumour: The Deadliest Cancer

\*

Betty Wood

Journal of Tumour Research and Reports, Spain

## ABSTRACT

Brain Tumour is growth of abnormal mass of cells which harms other neighbouring cells in the brain and leads to Brain Cancer in humans. Collection of neoplasms leads to tumour formation. Mostly, brain tumour arises from intracranial tissues and meninges. Tumours which are grown in brain itself are called Primary Brain Tumours whereas those which arouse in other parts of body but got into the brain are called Metastatic Brain Tumour. These Metastatic Tumours are mostly Malignant. Brain Tumour can be Benign (non-cancerous) or can be Malignant (cancerous). Some of the Benign Tumour are as follows: Chordomas, Craniopharyngiomas, Gangliocytomas, Meningiomas, Pineocytomas, Pituitary adenomas, Schwannomas and Acoustic neuromas. Malignant Tumour includes: Gliomas (the most deadly brain tumours).

**Keywords:** Benign Tumour; Gliomas; Metastatic Tumours

## INTRODUCTION

### What leads to Tumour Formation in the Brain?

Brain Tumour can be formed due to Mutations or defects in genes. Mutations in genes leads to formation of uncontrollable abnormal cells which divides into a large number and abruptly takes shape of tumour. Tumour causes self destruction of its own immune cells and leads to production of substances that blocks immune system so that it is unable to differentiate between normal cells and foreign body. Some of the other reasons are: Exposure to heavy amount of Ionizing Radiation which are mostly used in cancer treatment and when some genes in chromosome of cells are damaged. People at Risk: Brain tumour are more common in children and adults (mostly Male).

### Traits and Indicator of brain tumor

Central Nervous System (CNS) of the body is made up of Brain and Spinal column. CNS is responsible for controlling all body functions, thus when brain tumour occurs it affects thought processing capacity, vision and many more.

### Symptoms

Headaches (Occurs Often and worsens in morning and night); Behavior or personality changes; Seizures or convulsions (Uncontrollable Muscle Contractions); Confusion; Difficulty with balance or coordination; Dizziness; Trouble concentrating; Nausea and vomiting; Numbness, Weakness or Tingling (In one part or side of the body or face); Problems with Hearing, Vision or Speech, Seizures; Unusual sleepiness; Trouble with memory, Thinking,

speaking or understanding language; Weakness or paralysis in one part or one side of the body; Gradual loss of sensation or movement in an arm or a leg.

### Diagnosis and Cure

#### Diagnosis

**Neurological examination:** Checking Hearing, Vision, Reflexes as brain tumor has large impact on these body parts.

**Imaging:** Tests like Magnetic Resonance Imaging (MRI), Computerized Tomography (CT) Scan, and Positron Emission Tomography (PET) scan are used to easily diagnose brain tumor.

**Biopsy:** Piece of cells is removed to check whether tumor persist or not.

**Analysis of CSF:** Cerebrospinal fluid is tested to diagnose brain tumor.

**Skull X-Ray:** Skull fractures when brain tumor persist thus X-ray will help in diagnosing the tumor.

**Angiography:** Gives details about blood supply to tumor in the brain.

#### Cure

**Surgery:** Malignant Brain Tumours are removed using Surgery.

**Minimally Invasive Surgery:** Used for removing Cancerous tumours.

**Radiation Therapy:** X-rays are used for damaging tumour present in the brain.

**Correspondence to:** Betty Wood, Journal of Tumour Research and Reports, Spain; E-mail: woodbe@gmail.com

**Received:** February 19, 2021; **Accepted:** February 23, 2021; **Published:** March 10, 2021

**Citation:** Wood B (2021) Brain Tumour: The Deadliest Cancer. J Tumour Res Reports, 6:131.

**Copyright:** © 2021 Wood B. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

**Chemotherapy:** Drugs are injected to kill tumour or cancerous cells.

**Radical surgery:** Gamma Knife and Linear Accelerator is used to treat tumours cells in the brain.

## REFERENCES

1. Collins VP, Jones DT, Giannini C. Pilocytic astrocytoma: Pathology, molecular mechanisms and markers. *Acta Neuropathol.* 2015; 129(6): 775-788.
2. Ellison DW, Hawkins C, Jones DT, Onar-Thomas A, Pfister SM, Reifenberger G, et al. cIMPact-now update 4: Diffuse gliomas characterized by myb, mybl1, or fgfr1 alterations or braf v600e mutation. *Acta neuropathol.* 2019; 137(4): 683-687.
3. Jain SU, Khazaei S, Marchione DM, Lundgren SM, Wang X, Weinberg DN, et al. G34 mutations promote aberrant PRC2 activity and drive tumor progression. *Proc Natl Acad Sci.* 2020; 117(44): 27354-27364.
4. Jones DT, Bandopadhyay P, Jabado N. The power of human cancer genetics as revealed by low-grade gliomas. *Annu Rev Genet.* 2019; 53(1): 483-503.
5. Khuong-Quang DA, Buczkowicz P, Rakopoulos P, Liu XY, Fontebasso AM, Bouffet E, et al. K27M mutation in histone H3. 3 defines biologically distinct subgroups of pediatric diffuse intrinsic pontine gliomas. *Acta Neuropathol.* 2012; 124(3): 439-447.
6. Komori T. Updating the grading criteria for adult diffuse gliomas: Beyond the WHO2016CNS classification. *Brain Tumor Pathol.* 2020; 37(1): 1-4.
7. Komori T, Arai N. Dysembryoplastic neuroepithelial tumor, a pure glial tumor? Immunohistochemical and morphometric studies. *Neuropathology.* 2013; 33(4): 459-468.
8. Krishnatreya R, Zhukova N, Guerreiro Stucklin AS, et al. Clinical and treatment factors determining long-term outcomes for adult survivors of childhood low-grade glioma: A population based study. *Cancer.* 2016; 122: 1261-1269.
9. Ribom D, Eriksson A, Hartman M. Positron emission tomography (11) C-methionine and survival in patients with low-grade gliomas. *Cancer.* 2001; 92: 1541-1549.
10. Louis DN, Perry A, Reifenberger G, Von Deimling A, Figarella-Branger D. The 2016 World Health Organization classification of tumors of the central nervous system: A summary. *Acta Neuropathol.* 2016; 131(6): 803-820.