

Outcome and Prognosis of Metastatic Brain Tumour: A Study of 35 Cases

HARADHAN DEB NATH¹, KANAK KANTI BARUA², MOHAMMAD AFZAL HOSSAIN³,
MD ABUL KHAIR⁴, MD AMINUL ISLAM⁵

Abstract

Thirty five patients were selected to evaluate the outcome & prognosis of metastatic brain tumour in the department of Neurosurgery, Bangabandhu Sheikh Mujib Medical University, (BSMMU) Dhaka from February 2002 to December 2005.

A structured questionnaire was made. Data were collected after the patient admission. All patients were evaluated with detailed history and clinical examination. MRI of Brain of these patients revealed intracranial lesions highly suspicious of metastatic brain tumour. Investigations like USG of the whole abdomen and x-ray chest were done to locate any primary site. Some of the primary lesions were confirmed by FNAC and histological examinations. Solitary intracranial space occupying lesion (ICSOL) and sizable lesion among the multiple lesions were treated by surgery, and then histopathological examination were done followed by radiotherapy and / or chemotherapy.

This was a prospective study.

This study showed the highest age of incidence of tumours were above 60 years. Male predominated than female (60.00%). Among the clinical features the most

common sign was hemiparesis (34.28%). The commonest site of lesion was in frontal region (34.28%). Histopathological reports showed adenocarcinoma 40.00%, small cell carcinoma of lungs 28.57%, squamous cell carcinoma 22.85%, follicular carcinoma of thyroid 5.71%. Treatment options were surgery, radiotherapy and chemotherapy. Sixty percent patients were improved after treatment. Highest survival rate was 3 to 6 months with treatment (31.42%). This study revealed that commonest type of brain metastases was adenocarcinoma and primary site of lesion was lungs. Best option of treatment was surgery plus radiotherapy and longest survival rate of 1 year was in 20.00% cases.

Introduction:

The incidence of brain metastases and the spectrum of metastasizing primary cancers vary with patient's age¹. Brain metastases occur more frequently in adults than in children². Among adults, the highest incidence is observed in the fifth to seventh decades of life³. The most common sources of brain metastases in this group of patients were cancers of the lungs, breasts, and skin in descending order. In children, the most common cause of brain

1. Assistant Professor, Department of Neurosurgery, Chittagong Medical College, Chittagong.
2. Professor, Department of Neurosurgery, Bangabandhu Sheikh Mujib Medical University (BSMMU), Dhaka, Bangladesh.
3. Professor and Chairman, Department of Neurosurgery, BSMMU, Dhaka, Bangladesh.
4. Associate Professor, Department of Neurosurgery, BSMMU, Dhaka, Bangladesh.
5. Consultant Neurosurgeon, Combined Military Hospital, Dhaka, Bangladesh.

metastases is leukaemia, followed by lymphoma⁴.

The overall incidence of brain metastases is not affected by patient's gender, nor the incidence of brain metastases from a given primary cancer. The only apparent exception to this is melanoma, which is more likely to spread to the brain in male patients⁵. Lung cancer is the most common source of brain metastases in men where as breast cancer is the most common source in women⁶.

The histological type of the primary tumour appears to be the major dictator of the frequency and pattern of intracranial spread. Lungs, breast cancer, melanoma, renal and colon cancers account for the majority of all brain metastases in order of decreasing relative frequency. Primary lung tumours account for 30 to 60% of all brain metastases cases⁷. Breast cancer ranks second contributing 10 to 30% of all brain metastases among women. Approximately 5 to 21% of patients with brain metastases have melanoma as the primary tumour. However, virtually any malignancy can metastasize to the brain. Patients with symptoms caused by a brain metastases varies⁸.

Treatment modalities of brain metastases are: (i) radiation therapy a. Dose-fractionation schemes for whole brain radiation therapy b. altered fractionation schemes c. radiosensitizers d. prophylactic cranial irradiation for small cell lung cancer (ii) surgical resection (iii) postoperative whole brain radiation therapy (iv) stereotactic radiosurgery and (v) chemotherapy. Sometimes metastatic brain tumours are encountered where the primary escapes detection despite meticulous search by clinical and investigation method. Sometimes radiotherapy and

chemotherapy are used in regulated combination⁹.

Materials and Methods:

This was study a prospective study carried out from February 2002 to December 2005. A structured questionnaire was made. Data were collected after patient's admission. MRI of brain with contrast revealed intracranial tumour, whose radiological features were strongly suspicious of being metastatic tumour, histopathological diagnosis were made from material harvested at brain biopsy or by FNAC of obvious primaries.

Results:

Table-I

Distribution of patients by age (n=35)

Age in years	Number	%
>20	01	2.86
20-39	02	5.71
40-59	11	31.43
<60	21	60
Total	35	100

Table I shows that major proportion (60.00%) of the patients belonged to the age group of <60 years and the second height (31.43%) age group was 40-49 years. Only 5.7% was in the age of 20-39 years and 2.85% were in the age group of less than 20 years.

Table: II

Distribution of patients by sex (n=35)

Sex	Number	Percentage
Male	21	60.00
Female	14	40.00
Total	35	100.00

Table II shows that among the 35 patients 60% were males & 40% were females.

Table-III
Distribution of patients by occupations (n=35)

Occupations	Number	Percentage
Day labourer	11	31.43
Service holder	08	22.86
Farmer	07	20.00
Professional	04	11.43
Others	05	14.28
Total	35	100.00

Table III shows that the highest occupational group were day laborers (31.43%) followed by service-holders (22.85%) and farmers (20%).

Table-IV
Distribution of patients by presenting symptoms (n=35)

Presenting Symptoms	Number	Percentage
Headache	21	60.00
Vomiting	14	40.00
Convulsion	7	20.00
Altered Consciousness	5	14.29
Blurring of Vision	5	14.29

Table III shows that the commonest symptoms were headache (60%) followed by vomiting (40%) and convulsion (20%).

Table-V
Distribution of patients by clinical features (n=35)

Clinical features	Number	Percentage
Hemiplegia & hemiparesis	27	77.14
monoparesis	08	22.86
cranial nerve palsy	05	14.29
Impaired heigher psychic function	06	17.14

Table V shows that highest group of clinical presentation was one sided weakness (71.19%) followed by monoparesis (22.86%).

Table-VI
Distribution of patients by site of lesion (n=35)

Site	Number	Percentage
frontal lobe	12	34.28
Parietal lobe	10	28.57
Fronto-parietal	05	14.28
Temporal lobe	01	2.85
Occipital lobe	01	2.85
Cerebellum	04	11.42
Total	35	100

Table-VI shows that the commonest site of lesion was in frontal tobe followed by parietal lobe (28.57%) and frontoparietal lobe (14.28%).

Table-VII
Distribution of patients by histological types of the tumour (n=35)

Histological types of the tumuors	Number	Percentage
Adenocarcinoma	14	40.00
Small cell carcinoma of lungs	10	28.57
Squamous cell carcinoma	08	22.86
Follicular carcinoma of thyroid	03	8.57
Total	35	100.00

Table – VII shows that the heighest group of patients had adenocarcinoma (40.00%) followed by small cell carcinoma of lungs (28.57%) and squamous cell carcinoma (22.86%)

Table-VIII*Distribution of patients by primary site of lesions.*

Site	Number	Percentage
Lungs	21	60.00
GIT	03	8.57
Breasts	02	5.71
Thyroid	02	5.71
Kidneys	01	2.86
Scalp	01	2.86
Prostate	01	2.86
Unknown	04	11.42
Total	35	100.00

Table VIII shows that highest incidence of primary site was lungs (60%) followed by GIT (8.57%), breasts (5.7%) & thyroid (5.7%) respectively.

Table-IX*Distribution of patients by options of treatment (n=35)*

Treatment options of patients	Number	Percentage
Surgery + Radiotherapy	15	42.86
Surgery+Chemotherapy+ Radiotherapy	14	40.00
Radiotherapy	05	14.29
Radioiodine ablation	01	2.85
Total	35	100

Table IX shows that highest group of patients (42.86%) were treated by surgery+ radiotherapy followed by surgery + chemotherapy + Radiotherapy (40%) and Radiotherapy (14.29%) alone.

Table-X*Distribution of patients by outcome after treatment (n=35)*

Outcome	Number	Percentage
Improved	21	60.00
Same as before	08	22.86
Deteriorated	03	8.57
Others	03	8.57
Total	35	100

Table-X shows that highest group (60%) of patients were improved after treatment and 22.85% of patients remained same as before & 8.57% patients deteriorated.

Table: XI*Distribution of patients survival rate after treatment (n=35).*

Months	Number	Percentage
> 3 months	04	11.42
3 to 6 months	11	31.42
> 6 to 9 months	10	28.57
> 9 to 12 months	07	20.00
12 months and more	03	8.57

Table – XI shows that highest survival rate was 3 to 6 months (31.42%) followed by 6-9 months (28.57) and 9-12 months (20.00%).

**Fig.-1:** *Small cell carcinoma metastases from lungs.*

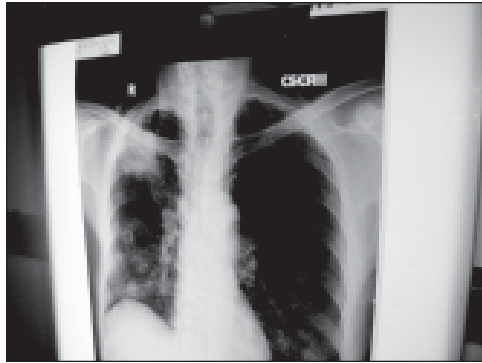


Fig.-2: *Small cell carcinoma of lungs of same patient.*

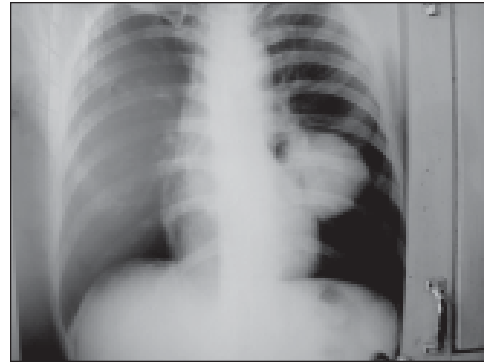


Fig.-5: *Squamous cell carcinoma of lungs of the same patient.*

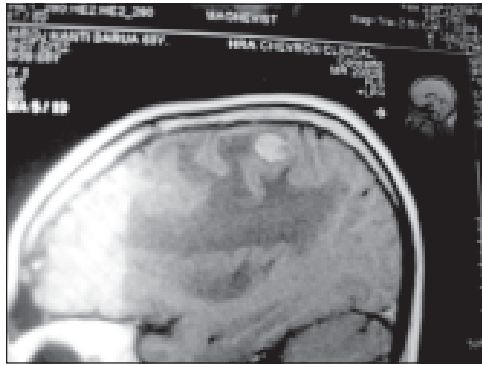


Fig.-3: *Adenocarcinoma metastases from lungs.*

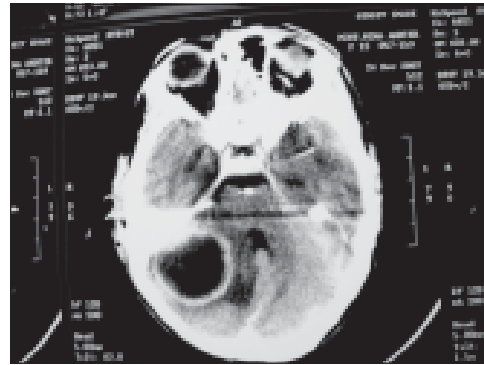


Fig.-6: *Adenocarcinoma metastases from Kidney.*

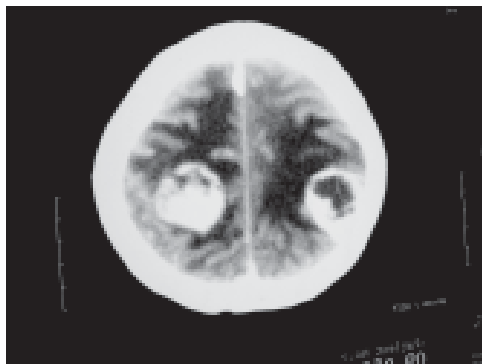


Fig.-4: *Squamous cell carcinoma metastases from lungs.*

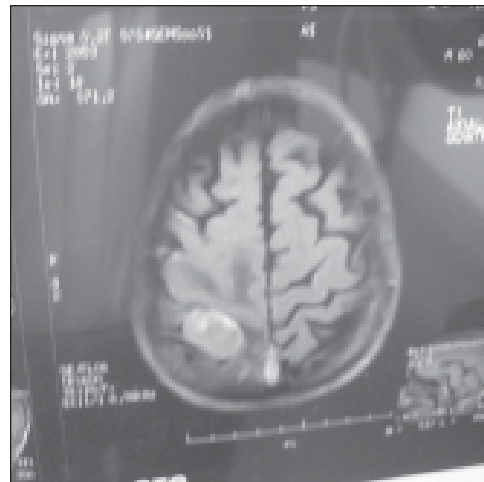


Fig.-7: *Follicular carcinoma metastases from thyroid.*

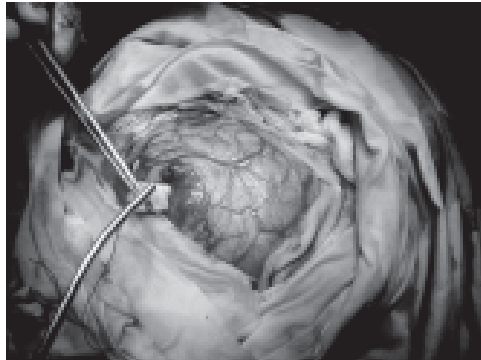


Fig.-8: *Per-operative picture of a metastatic brain tumor.*



Fig.-9: *Post-operative picture of a metastatic brain tumor patient.*

Discussion:

This was a prospective study. In this study the highest age group was <60 years. In a previous study the highest age groups were from fifth to seventh decade¹⁰. Among the clinical features most common was one sided weakness. Most of the primary sites were from lungs (60%). In a previous study the most common sources of brain metastases were cancers of the lung, breast and skin, in descending order¹⁰. In our study the commonest site of metastasis was frontal lobe (34.28%) and the most common histopathological type was

adenocarcinoma (40.00%). Of patients with lung cancer, 18 to 65% developed brain metastasis and more than 40% of the patients had adenocarcinoma in a previous study¹¹. In this study, treatment options were surgery, radiotherapy, chemotherapy, radioiodine-ablation etc. Sixty percent of the patients improve after treatment and survival rate was 3 to 6 months in 31.42% cases, 6 to 9 months in 28.57% cases, 9 to 12 months in 20.00% cases and more than 12 months were 8.57% cases. In a previous study median survival time of 1 year was in 30% of patients¹².

Conclusion:

This study showed that commonest type of brain metastases was adenocarcinoma and primary site of lesion was lungs. Best option of treatment was surgery plus radiotherapy and median survival rate of 1 year was in 20.00% of cases.

References:

1. Lindsay KW, Bone I. Spinal cord and root compression. In: Lindsay KW, Bone I, Editors, Neurology and Neurosurgery Illustrated, 3rd ed. London: Churchill Livingstone, 1997; p 383.
2. Gowers WR. Syringal haemorrhage into the spinal cord.. In: Churchill, Editor. Lectures on diseases of the nervous system. 2nd edition. Churchill livingstone, 1904; p 200.
3. MacEwen W. Meningitis, abscess of the brain, infective sinus thrombosis. In: J Maclose. editor. Pyogenic infective diseases of the brain and spinal cord. 4th ed. London: Churchill Livingstone, 1883; p 354.

4. Elseberg CA, Beer E. The operability of intramedullary tumors of the spinal cord . Remarks on extrusion of intraspinal tumors. *Am J Med sci* 1911; 142: 636.
5. Kopelson G, Dinggood RM, Kleinman GM. Management of Intramedullary Spinal Cord Tumors. *Radiology* 1980; 135: 437-9.
6. Adams RD, Victor M: Intraspinal Tumors. In: Adams RD, Victor M editors. *Principles of Neurology*. 2nd edition . New York, Mcgraw Hill, 1981; pp 638-41.
7. Stein B. Surgery of Intramedullary Spinal Cord Tumors. *Clin Neurosurg* 1979; 26: 473-9.
8. Chin HW, Hazel JJ, Kum TH. Oligodendrogliomas. I. A Clinical Study of Cerebral Oligodendrogliomas. *Cancer* 1998; 45: 1458-66.
9. Nittner K. Spinal tumor. In: Olivecrona H and Tonnis W, editors, *Handbuch der Neurochirurgie*, 7th edition, New York, Springer Verlag, 1972 ; pp 1-606.
10. Lyons MK, O'Neill BP, Kurtin PJ. Diagnosis and Management of Primary Spinal Epidural Non-Hodgkin's Lymphoma. *Mayo Clin Proc* 1996; 71: 453-7.
11. Go RCP, Lamiell JM, Hsia YE. Segregation and Linkage Analysis to von Hippel-Lindau Disease Among 220 Descendants from one Kindred. *Am J Human Genet* 1984; 36: 131-42.
12. Glenn GM, Linehan WM, Hosoe S. Screening for von Hipel-Lindau Disease by DNA Polymorphism Analysis. *JAMA* 1992; 267: 1226-31.