

# MASS EFFECT AND MIDLINE SHIFT IN GLIOBLASTOMA-WORSE PROGNOSIS?

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#### Introduction:

- Midline shift caused by glioblastoma mass effect has been associated with a worse prognosis
- This evidence is not consistent in the literature

### Objective:

 To analyse relationship between midline shift (MLS), progression free survival (PFS) and overall survival (OS) in a series of glioblastoma patients submitted to resective surgery at our centre

#### Methods:

- Retrospective study of pre-operative MRI scans (axial contrast-enhanced T1-weighted or T2-weighted sequences) of 163 glioblastoma patients submitted to resective surgery
- MLS measurements were performed according to the c=a/
  2-b method described in the literature
- PFS and OS calculated with Kaplan-Meier method (log-rank test SPSS version 23)

### Results:

Table 1 - Patient data and results

No. patients	163
Sex (male/female)	98/65
Median Age (years; range)	59 (26-79)
ECOG 0-1	129 (79.1%)
ECOG >1	34 (20.9%)
Gross Total Resection	91 (55.8%)
MLS (mm; median; range)	3.5 (0-15.2)
MLS < 5mm	107 (65.6%)
MLS ≥ 10mm	14 (8.6%)

Global PFS (months; median; range) 8 (1-66)

Global OS (months; median; range) 17 (3-87)

Figure 1 - PFS Kaplan-Meier (KM) curves for groups MLS < 5mm vs. MLS  $\geq$  5 mm (left image) and MLS < 10mm vs. MLS  $\geq$  10 mm (right image).

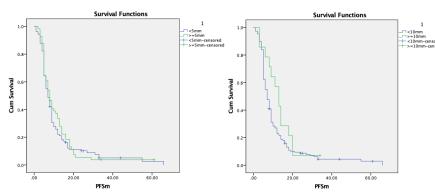
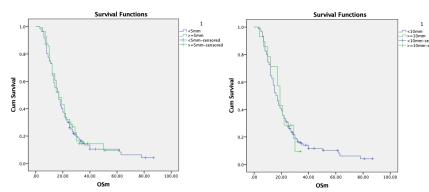


Figure 2 - OS KM curves for groups MLS < 5mm vs. MLS  $\geq$  5 mm (left image) and MLS < 10mm vs. MLS  $\geq$  10 mm (right image).



Median PFS (Fig. 1)

MLS < 5mm vs.  $\geq$  5 mm =  $\frac{7 \text{ vs. 8}}{10 \text{ months}}$  months MLS <  $\frac{10 \text{ mm}}{10 \text{ vs.}} = \frac{7 \text{ vs. 13}}{10 \text{ months}}$  p = 0.591p = 0.147

Median OS (Fig. 2) MLS < 5mm vs.  $\geq$  5 mm =  $\frac{17 \text{ vs. } 17}{17 \text{ months}}$ MLS <  $\frac{10}{17 \text{ months}}$  p = 0.875p = 0.921

## Conclusion:

- Despite some previous evidence that MLS may have an impact on PFS or OS, our series does not replicate these results
- Therefore, we believe MLS should not be considered as a prognostic marker in glioblastoma, nor should it be used to influence the choice of treatment in these patients

## References:

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