Glioblastoma Bill of Rights

Glioblastoma is an aggressive, deadly brain tumor that currently has no cure. People living with GBM deserve quality care and the ability to make informed decisions.

Now is the time for a GBM Bill of Rights. Developed by patients, alongside our families and advocates, this is the road map to patient power and a brighter future.

**We, the patients, have...**

- **THE RIGHT TO 1** Patient-centered care
  - An equal and collaborative patient/doctor relationship with personalized treatment that includes genomic and genetic testing, and tumor analysis.

- **THE RIGHT TO 2** A second opinion
  - A second opinion offered at appropriate stages of diagnosis and treatments.

- **THE RIGHT TO 3** Clarity of information
  - Up-to-date, accurate information about all potential treatment options; transparency about treatment centers and their outcomes.

- **THE RIGHT TO 4** Specialist care
  - Access to a coordinated medical team — brain tumor specialist, neuro-surgeon, neuro-oncologist, radiation oncologist, and others.

- **THE RIGHT TO 5** Access to trials and experimental treatments
  - Up-to-date, accurate information about new drugs, treatments, and clinical trials, including international options.

- **THE RIGHT TO 6** Psychological support
  - Comprehensive psychological help for us and our carers.

- **THE RIGHT TO 7** Money-blind treatment
  - Best treatment regardless of personal financial situation.

- **THE RIGHT TO 8** Fast-tracked treatments
  - Fast-tracked MRI results, treatments, and ER visits.

- **THE RIGHT TO 9** Patient-owned data
  - Ownership of and access to our health data, and the ability to transport and donate our data and tissue for research.

- **THE RIGHT TO 10** Best end-of-life care
  - Access to pain management, palliative care, and hospice.

- **THE RIGHT TO 11** Brain donation
  - Information provided to us and our families about no-cost brain donation for scientific research.

Thanks to the supporters of this project: