

PubMed

**Format:** Abstract

J Clin Oncol. 2019 Mar 20;37(9):731-740. doi: 10.1200/JCO.18.00969. Epub 2019 Feb 7.

## Late Morbidity and Mortality Among Medulloblastoma Survivors Diagnosed Across Three Decades: A Report From the Childhood Cancer Survivor Study.

Salloum R<sup>1</sup>, Chen Y<sup>2</sup>, Yasui Y<sup>2,3</sup>, Packer R<sup>4</sup>, Leisenring W<sup>5</sup>, Wells E<sup>4</sup>, King A<sup>6</sup>, Howell R<sup>7</sup>, Gibson TM<sup>3</sup>, Krull KR<sup>3</sup>, Robison LL<sup>3</sup>, Oeffinger KC<sup>8</sup>, Fouladi M<sup>1</sup>, Armstrong GT<sup>3</sup>.

1 Cincinnati Children's Hospital Medical Center, Cincinnati, OH.

2 University of Alberta, Edmonton, Alberta, Canada.

3 St. Jude Children's Research Hospital, Memphis, TN.

4 Children's National Health System, Washington, DC.

5 Fred Hutchinson Cancer Research Center, Seattle, WA.

6 Washington University in St Louis, St Louis, MO.

7 The University of Texas MD Anderson Cancer Center, Houston, TX.

8 Duke University, Durham, NC.

**PURPOSE:** Treatment of medulloblastoma has evolved from surgery and radiotherapy to contemporary multimodal regimens. However, the impact on long-term health outcomes remains unknown.

**METHODS:** Cumulative incidence of late mortality (5 or more years from diagnosis), subsequent neoplasms (SNs), and chronic health conditions were evaluated in the Childhood Cancer Survivor Study among 5-year survivors of medulloblastoma diagnosed between 1970 and 1999. Outcomes were evaluated by treatment exposure, including historical therapy (craniospinal irradiation [CSI]  $\geq$  30 Gy, no chemotherapy), high risk (CSI  $\geq$  30 Gy + chemotherapy), standard risk (CSI  $<$  30 Gy + chemotherapy), and by treatment decade (1970s, 1980s, 1990s). Rate ratios (RRs) and 95% CIs estimated long-term outcomes using multivariable piecewise exponential models.

**RESULTS:** Among 1,311 eligible survivors (median age, 29 years [range, 6 to 60 years]; median time from diagnosis, 21 years [range, 5 to 44 years]), the 15-year cumulative incidence rate of all-cause late mortality was 23.2% (diagnosed 1970s) versus 12.8% (1990s;  $P = .002$ ), with a recurrence-related mortality rate of 17.7% versus 9.6% ( $P = .008$ ). Lower late mortality rates as a result of other health-related causes were not observed. Among 997 survivors who completed a baseline survey, the 15-year cumulative incidence of SNs was higher among survivors with multimodal therapy (standard risk, 9.5%; historical, 2.8%;  $P = .03$ ). Survivors treated in the 1990s had a higher cumulative incidence of severe, disabling, life-threatening, and fatal chronic health conditions (56.5% in 1990s v 39.9% in 1970s;  $P < .001$ ) and were more likely to develop multiple

conditions (RR, 2.89; 95% CI, 1.31 to 6.38). However, survivors of standard-risk therapy were less likely to use special education services than high-risk therapy survivors (RR, 0.84; 95% CI, 0.75 to 0.93).

**CONCLUSION:** Historical changes in medulloblastoma therapy that improved 5-year survival have increased the risk for SNs and debilitating health conditions for survivors yet reduced the need for special education services.

PMID: 30730781 DOI: [10.1200/JCO.18.00969](https://doi.org/10.1200/JCO.18.00969)

---