

Neuroblastoma outcome

For children with low-risk neuroblastoma, the 5-year survival rate is higher than 95%. For children with intermediate-risk neuroblastoma, the 5-year survival rate is between 90% and 95%. For high-risk neuroblastoma, the the-5-year survival rate is around 40% to 50%.

The unfavorable prognosis with a high risk of relapse and death in NB correlates with age over 18 months at diagnosis, advanced disease stage, and established genetic markers. Amplification of the [MYCN](#) oncogene (MNA) is the most robust genetic factor correlated with poor clinical outcome and can be found in about 16–20% of NB cases (and up to 40% in high-risk tumors) ^{1) 2) 3) 4) 5)}.

The regulation of NB [cell death](#) by MYCN represents an important aspect, as it directly contributes to [tumor progression](#) and therapeutic resistance. However, the relationship between MYCN and [cell death](#) remains elusive. [Ferroptosis](#) is a newly identified cell death mode featured by [lipid peroxide](#) accumulation that can be attenuated by [GPX4](#), yet whether and how MYCN regulates ferroptosis are not fully understood.

Lu et al. reported MYCN-amplified NB cells are sensitive to GPX4-targeting ferroptosis inducers. Mechanically, MYCN expression reprograms the cellular [iron metabolism](#) by upregulating the expression of [TFRC](#), which encodes [transferrin receptor 1](#) as a key iron transporter on the cell membrane. Further, the increased [iron](#) uptake promotes the accumulation of labile iron pool, leading to enhanced [lipid peroxide](#) production. Consistently, TFRC overexpression in NB cells also induces selective sensitivity to GPX4 inhibition and ferroptosis. Moreover, they found that MYCN fails to alter the general [lipid metabolism](#) and the amount of [cystine](#) imported by System Xc(-) for [glutathione](#) synthesis, both of which contribute to ferroptosis in alternative contexts. In conclusion, NB cells harboring [MYCN](#) amplification are prone to undergo [ferroptosis](#) conferred by [TFRC](#) upregulation, suggesting that [GPX4](#)-targeting ferroptosis inducers or [TFRC](#) agonists can be potential strategies in treating MYCN-amplified NB ⁶⁾.

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Ambros PF, Ambros IM, Brodeur GM, Haber M, Khan J, Nakagawara A, et al. . International consensus for neuroblastoma molecular diagnostics: Report from the International Neuroblastoma Risk Group (INRG) Biology Committee. Br J Cancer. (2009) 100:1471–82. 10.1038/sj.bjc.6605014

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Canete A, Gerrard M, Rubie H, Castel V, Di Cataldo A, Munzer C, et al. . Poor survival for infants with MYCN amplified metastatic neuroblastoma despite intensified treatment: the International Society of Paediatric Oncology European Neuroblastoma Experience. J Clin Oncol. (2009) 27:1014–9. 10.1200/JCO.2007.14.5839

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Huang M, Weiss WA. Neuroblastoma and MYCN. Cold Spring Harb Perspect Med. (2013) 3:a014415. 10.1101/cshperspect.a014415

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Cohn SL, Pearson AD, London WB, Monclair T, Ambros PF, Brodeur GM, et al. . The International Neuroblastoma Risk Group (INRG) classification system: an INRG task force report. J Clin Oncol. (2009) 27:289–97. 10.1200/JCO.2008.16.6785

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Thompson D, Vo KT, London WB, Fischer M, Ambros PF, Nakagawara A, et al. . Identification of patient subgroups with markedly disparate rates of MYCN amplification in neuroblastoma: a report from the International Neuroblastoma Risk Group project. *Cancer*. (2016) 122:935–45. 10.1002/cncr.29848⁶⁾

Lu Y, Yang Q, Su Y, Ji Y, Li G, Yang X, Xu L, Lu Z, Dong J, Wu Y, Bei JX, Pan C, Gu X, Li B. MYCN mediates TFRC-dependent ferroptosis and reveals vulnerabilities in neuroblastoma. *Cell Death Dis*. 2021 May 19;12(6):511. doi: 10.1038/s41419-021-03790-w. PMID: 34011924.

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