

Oxygen Saturations and Alarm Limits

Premature infants born at <1000 grams (typically <28 weeks GA) receive respiratory support in the NICU. The recommended **SpO₂ target range is 90–95%** (AAP/most international consensus) or **90–94%** (European Consensus Guidelines), with **alarm limits set at 89% (low) and 95% (high)** per the European guidelines. These recommendations apply regardless of whether the infant is on invasive or non-invasive respiratory support.

Guideline Recommendations by Society Summary with Supporting Evidence

Guideline / Society	SpO ₂ Target Range	Alarm Limits	Notes	References
AAP (2016 Clinical Report)	90–95%	Not explicitly specified	Stated 90–95% "may be safer" than 85–89%; acknowledged ideal target may be individualized by GA, chronological age, disease, and transfusion status	[1] , [2]
European Consensus Guidelines (2022 update)	90–94%	Low: 89%, High: 95%	Most specific guidance on alarm limits; applies to preterm infants on supplemental O ₂	[1]
NICE (UK)	91–95%	Not explicitly specified	For preterm infants <32 weeks GA	[3]
Saugstad (2018) Expert Commentary	90–94%	Low: 89%, High: 95%	Based on NeOProM meta-analysis; from birth to 36 weeks PCA for infants <28 weeks GA	[4]

Evidence Base: The NeOProM Collaboration - The recommendations are grounded in five large RCTs (SUPPORT, BOOST-II UK/Australia/New Zealand, COT) enrolling 4,965 infants <28 weeks GA, collectively analyzed in the NeOProM meta-analysis: [\[2\]](#)[\[5\]](#)

Lower target (85–89%) was associated with **increased mortality** (RR 1.16–1.17) and **increased NEC** (RR 1.24), but **reduced severe ROP** requiring treatment (RR 0.72–0.74). [\[4\]](#)[\[2\]](#)[\[3\]](#)

Higher target (91–95%) was associated with lower mortality but higher rates of treated ROP, though importantly **no difference in blindness** was observed between groups. [\[4\]](#)[\[2\]](#)

The composite outcome of **death or major disability** did not differ significantly between groups in the individual patient data meta-analysis (RR 1.04, 95% CI 0.98–1.09). [\[3\]](#)

The **BOOST-II trials** demonstrated this mortality signal most clearly, as shown in the following forest plot comparing outcomes between the two target ranges:

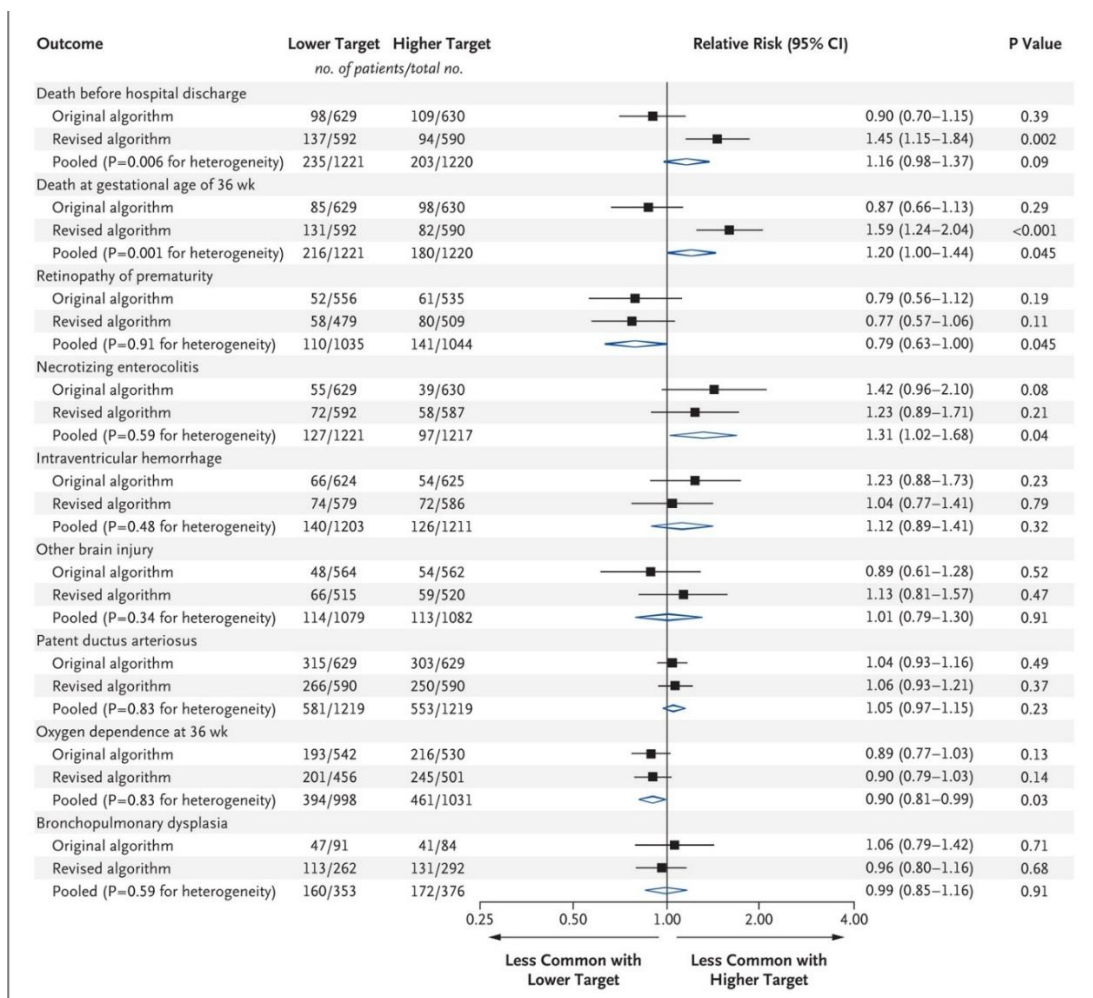


Figure 3 Combined Discharge Outcomes from the Three Trials, According to Oxygen-Saturation Target and Status of the Oximeter-Calibration Algorithm.

Practical Alarm Setting Considerations

The risk of severe hypoxemia ($\text{PaO}_2 \leq 40$ mmHg) becomes marked at $\text{SpO}_2 \leq 85\%$; the risk of severe hyperoxemia ($\text{PaO}_2 \geq 100$ mmHg) becomes marked at $\text{SpO}_2 \geq 98\%$ in infants <33 weeks PMA, supporting alarm limits that trigger before these extremes are reached.^[7]

Maintaining SpO_2 within a narrow target range (e.g., 90–94%) is inherently challenging as the oxyhemoglobin dissociation curve nears vertical around 90% SpO_2 , meaning small PaO_2 changes produce large SpO_2 swings.^[8]

Intermittent hypoxia ($\text{SpO}_2 < 80\%$ for ≥ 1 minute) and oxygen fluctuations are independent risk factors for ROP, emphasizing the importance of minimizing alarm response time and SpO_2 variability.^[3]

Automated closed-loop FiO_2 systems have been shown to increase time spent within the target SpO_2 range compared with manual adjustment, though long-term outcome data remain limited.^{[8][9]}

Signal averaging pulse oximeter time: longer averaging dampens artifact; may mask brief hypoxic episodes ($\text{SpO}_2 < 80\%$).^[9]

Biphasic Targeting (Emerging Concept) - Some retrospective cohort data suggest a **biphasic approach**, targeting lower SpO_2 (e.g., 88–92%) before 30–32 weeks PMA to reduce ROP risk, then increasing the target (e.g., 92–97%) after 32 weeks PMA to reduce mortality, may improve both ROP and survival outcomes. No RCTs have validated this strategy, and the current standard of care remains a **uniform target of 90–95%**.^{[3][10]}

References

1. [Oxygen in the newborn period: Could the oxygen reserve index offer a new perspective?](#). Liz CF, Proença E. Pediatric Pulmonology. 2025;60(1):e27343. doi:10.1002/ppul.27343.
2. [Effects of Targeting Lower Versus Higher Arterial Oxygen Saturations on Death or Disability in Preterm Infants](#). Askie LM, Darlow BA, Davis PG, et al. The Cochrane Database of Systematic Reviews. 2017;4:CD011190. doi:10.1002/14651858.CD011190.pub2.
3. [Retinopathy of Prematurity: A Global Perspective and Recent Developments](#). Sabri K, Ells AL, Lee EY, Dutta S, Vinekar A. Pediatrics. 2022;150(3):e2021053924. doi:10.1542/peds.2021-053924.
4. [Oxygenation of the Immature Infant: A Commentary and Recommendations for Oxygen Saturation Targets and Alarm Limits](#). Saugstad OD. Neonatology. 2018;114(1):69-75. doi:10.1159/000486751.
5. [The Evolution of Modern Respiratory Care for Preterm Infants](#). Owen LS, Manley BJ, Davis PG, Doyle LW. Lancet (London, England). 2017;389(10079):1649-1659. doi:10.1016/S0140-6736(17)30312-4.
6. [Oxygen Saturation and Outcomes in Preterm Infants](#). BOOST II United Kingdom Collaborative Group, BOOST II Australia Collaborative Group, BOOST II New Zealand Collaborative Group, et al. The New England Journal of Medicine. 2013;368(22):2094-104. doi:10.1056/NEJMoa1302298.
7. [Thresholds for Oximetry Alarms and Target Range in the NICU: An Observational Assessment Based on Likely Oxygen Tension and Maturity](#). Bachman TE, Iyer NP, Newth CJL, Ross PA, Khemani RG. BMC Pediatrics. 2020;20(1):317. doi:10.1186/s12887-020-02225-3.
8. [Update on ventilatory management of extremely preterm infants—A Neonatal Intensive Care Unit perspective](#). Schulzke SM, Stoecklin B. Paediatric Anaesthesia. 2022;32(2):363-371. doi:10.1111/pan.14369.
9. [Automated Oxygen Delivery for Preterm Infants With Respiratory Dysfunction](#). Stafford IG, Lai NM, Tan K. The Cochrane Database of Systematic Reviews. 2023;11:CD013294. doi:10.1002/14651858.CD013294.pub2.
10. [Oxygen Saturation Targets in Neonatal Care: A Narrative Review](#). Nguyen TC, Madappa R, Siefkes HM, et al. Early Human Development. 2024;199:106134. doi:10.1016/j.earlhumdev.2024.106134.

Use of Histograms

Oxygen saturation histograms are recommended as a practical, **multidisciplinary quality improvement tool (QI)** for monitoring SpO₂ targeting in preterm infants on respiratory support, though no formal society guideline mandates their use. The evidence supporting their adoption comes primarily from QI studies demonstrating that periodic histogram review **increases time within the target SpO₂ range and is associated with reductions in death or severe ROP**.[\[1\]\[2\]\[3\]](#)

What Are SpO₂ Histograms?

SpO₂ histograms are graphical displays generated by bedside patient monitors (or dedicated software tools) that show the distribution of SpO₂ values over a defined time period. They quantify the percentage of time an infant spends within, above, and below the target SpO₂ range, providing an objective summary that overcomes the limitations of spot checks and manual transcription, which are prone to error and observer fatigue.[\[1\]\[4\]](#)

Evidence Supporting Histogram Use:

Gentle et al. (2020) - In a single-center QI initiative involving 518 infants <29 weeks GA, integration of periodic histogram monitoring into **multidisciplinary rounds** (providers, respiratory therapists, and nurses) increased **time at goal SpO₂ (90–95%) from 48.7% to 57.6%** and was associated with a reduction in the composite outcome of **death or severe ROP from 32.1% to 18.0%**.[\[2\]](#)

Srivatsa et al. (2021) - A retrospective cohort study of 1,532 infants at 23–28 weeks GA across three epochs showed that implementation of automated bedside SpO₂ histograms (Epoch 2) and subsequent software enhancements incorporating simultaneous SpO₂ and FiO₂ monitoring (Epoch 3) were associated with significant reductions in **mortality (aOR 0.48), any ROP (aOR 0.38), ROP requiring treatment (aOR 0.43), and BPD (aOR 0.43)** compared with the pre-histogram era.[\[3\]](#)

How Histograms Are Used in Practice: Sur and Paria (2021) reviewed the technology and evidence behind

histogram analysis and identified several practical applications:[\[1\]](#)

Periodic review during rounds: Histograms are reviewed at regular intervals (e.g., every 4–8 hours or during multidisciplinary rounds) to assess whether the infant is spending adequate time within the target range and to identify patterns of hypoxia or hyperoxia that may not be apparent from real-time monitoring alone.

Guiding respiratory support decisions: Histogram data can be incorporated into algorithms for weaning or escalating respiratory support, providing more objective criteria than point-of-care observations alone.

Nursing-driven monitoring: With appropriate training, nursing staff can use histogram data to reduce the fatigue and error associated with manual recording, enabling more consistent SpO₂ targeting.

Web-Based Oxygenation Dashboards: Poppe et al. (2024) developed a web-based oxygenation dashboard for preterm neonates (<32 weeks GA) integrating real-time SpO₂ histograms with FiO₂ data, SpO₂/FiO₂ ratios, and area under the curve for hypoxic (<80%) and hyperoxic (>95%) episodes. In 65% of patient evaluations, the level of hypoxia was assessed differently when the dashboard was used compared with routine assessment, and 75% of clinicians judged the dashboard to provide added value for clinical decision-making.[\[4\]](#)

Context: Time Within Target Range - The challenge of maintaining SpO₂ within a narrow target range is well documented. Major RCTs reported that patients spend **9–27% of time with SpO₂ <85% and 11–33% of time with SpO₂ >95%** even under study conditions.[\[5\]](#) Automated closed-loop FiO₂ systems represent the most "failsafe" approach to maintain target compliance, but they are not widely available; histogram monitoring serves as a more accessible strategy between manual adjustment alone and full automation.[\[1\]\[5\]](#)

Summary

While no RCTs have been conducted specifically on histogram monitoring as an intervention, the available QI evidence consistently supports its use as a **low-cost, practical tool that improves SpO₂ target adherence and is associated with improved clinical outcomes** including reduced ROP and mortality in extremely preterm infants.[\[1\]\[2\]\[3\]](#) Histogram review is best implemented as part of a multidisciplinary oxygen management bundle that includes standardized SpO₂ targets, alarm limit policies, staff education, and regular audit of achieved saturations.

References

1. [Histogram Analysis for Bedside Respiratory Monitoring in Not Critically Ill Preterm Neonates: A Proposal for a New Way to Look at the Monitoring Data](#). Sur A, Paria A. European Journal of Pediatrics. 2021;180(1):283-289. doi:10.1007/s00431-020-03732-2.
2. [Oxygen Saturation Histogram Monitoring to Reduce Death or Retinopathy of Prematurity: A Quality Improvement Initiative](#). Gentle S, El-Ferzli G, Winter L, Salas AA, Philips Iii JB. Journal of Perinatology : Official Journal of the California Perinatal Association. 2020;40(1):163-169. doi:10.1038/s41372-019-0486-7.
3. [Effect of a Novel Oxygen Saturation Targeting Strategy on Mortality, Retinopathy of Prematurity, and Bronchopulmonary Dysplasia in Neonates Born Extremely Preterm](#). Srivatsa B, Malcolm K, Clark RH, Kupke KG. The Journal of Pediatrics. 2021;234:33-37.e3. doi:10.1016/j.jpeds.2021.03.007.
4. [Development of a Web-Based Oxygenation Dashboard for Preterm Neonates: A Quality Improvement Initiative](#). Poppe JA, Smorenburg RS, Goos TG, et al. Journal of Medical Systems. 2024;48(1):46. doi:10.1007/s10916-024-02064-0.
5. [Automated control of inspired oxygen \(FiO₂\) in preterm infants: Literature review](#). Dani C. Pediatric Pulmonology. 2019;54(3):358-363. doi:10.1002/ppul.24238.