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The Association of Level of Care With NICU Quality

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BACKGROUND: Regionalized care delivery purportedly optimizes care to vulnerable very low birth weight (VLBW; <1500 g) infants. However, a comprehensive assessment of quality of care delivery across different levels of NICUs has not been done.

METHODS: We conducted a cross-sectional analysis of 21 051 VLBW infants in 134 California NICUs. NICUs designated their level of care according to 2012 American Academy of Pediatrics guidelines. We assessed quality of care delivery via the Baby-MONITOR, a composite indicator, which combines 9 risk-adjusted measures of quality. Baby-MONITOR scores are measured as observed minus expected performance, expressed in standard units with a mean of 0 and an SD of 1.

RESULTS: Wide variation in Baby-MONITOR scores exists across California (mean [SD] 0.18 (1.14), range –2.26 to 3.39). However, level of care was not associated with overall quality scores. Subcomponent analysis revealed trends for higher performance of Level IV NICUs on several process measures, including antenatal steroids and any human milk feeding at discharge, but lower scores for several outcomes including any health care associated infection, pneumothorax, and growth velocity. No other health system or organizational factors including hospital ownership, neonatologist coverage, urban or rural location, and hospital teaching status, were significantly associated with Baby-MONITOR scores.

CONCLUSIONS: The comprehensive assessment of the effect of level of care on quality reveals differential opportunities for improvement and allows monitoring of efforts to ensure that fragile VLBW infants receive care in appropriate facilities.



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Dr Profit had full access to all of the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis; he acquired funding for the study, conceptualized and designed the study, selected data for inclusion in analyses, analyzed the data, interpreted the results, and drafted the initial manuscript; Drs Gould, Goldstein, and Phibbs helped design the analysis and interpret the results and revised the manuscript; Dr Bennett executed the analysis, helped to interpret the results, and revised the manuscript; Dr Draper helped design the analysis and interpret the results, and revised the manuscript; Dr Lee helped design the analysis and interpret the results, analyzed the data, and revised the manuscript; Dr Lee helped design the study, assisted with interpretation of the results, and revised the manuscript; and all authors approved the final manuscript as submitted.

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Address correspondence to Jochen Profit, MD, MPH, Perinatal Epidemiology and Health Outcomes Research Unit, Division of Neonatology, Department of Pediatrics, Stanford University School of Medicine, MSOB Room x115, 1265 Welch Rd, Stanford, CA 94305. E-mail: profit@stanford.edu WHAT'S KNOWN ON THIS SUBJECT: Regionalized NICU care delivery and birth at perinatal centers minimizes mortality in very low birth weight infants. There is a lack of a more comprehensive assessment of quality and outcomes of care across different levels of care.

WHAT THIS STUDY ADDS: Using the Baby-MONITOR, we found wide differences in quality of care provided to very low birth weight infants across NICUS. Level of care was not associated with Baby-MONITOR scores, but subcomponents highlighted opportunities for improvement at all levels.

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Delivery of neonatal intensive care in regionalized systems has long been regarded as critical to providing high-quality health care to vulnerable very low birth weight (VLBW; <1500 g) infants. However, over the past decades the regionalized care systems for sick newborns may have been weakened by financial rewards under fee-for-service arrangements and demand from community hospitals and families seeking to deliver close to home.¹

Lower mortality of VLBW infants has been observed after birth in a perinatal center.² Phibbs and colleagues showed higher mortality in lower-level and lower-volume NICUs.^{3,4} A meta-analysis indicated 62% higher odds of mortality during the birth hospitalization with birth outside a high-level NICU.⁵ NICU volume may be an even more important predictor of mortality.^{6,7}

These studies imply that quality of care delivery for vulnerable VLBW infants at lower-level NICUs may be suboptimal. However, mortality as a sole measure of quality is limited. In isolation, it provides little information about the care provided to the 85% of infants⁸ who survive to discharge.⁹ Yet a comprehensive assessment of care and outcomes across different levels of NICUs does not exist.

Neonatal intensive care is a complex and multidimensional activity, and the measurement of its quality should reflect this fact. Although individual measures contain important information, there is also value in summarizing performance by combining the information from multiple measures because such a summary can convey quality from many different perspectives.¹⁰ In previous work, we created a composite indicator, the Baby-MONITOR, as a comprehensive measure of the care and outcomes for VLBW infants.^{11,12} In this article, we used the Baby-MONITOR and its individual components to examine

whether care and outcomes differ between different NICU levels.

METHODS

Overview

We conducted a cross-sectional population-based analysis of clinical data obtained from the California Perinatal Quality Care Collaborative (CPQCC).¹³ More than 90% of California NICUs are members of the CPQCC. Data for this study are derived from the CPQCC clinical data sets, which include several quality assurance mechanisms. Annual training sessions for local NICU personnel help to promote accuracy and uniformity in data abstraction. In addition, each record has range and logic checks, both at the time of data collection and data closeout, with auditing of records with excessive missing data.

The sample included live-born infants with a birth weight of 401 to 1500 g or a gestational age between 25 0/7 and 31 6/7 weeks. We used multiyear analyses (January 1, 2008, to December 31, 2012) because of the small number of VLBW infants cared for in some institutions.

We used previously published selection criteria aimed at creating a relatively homogenous sample of VLBW infants.¹¹ To ensure that patient outcomes reflected NICU quality of care, we excluded infants who died before 12 hours of life and those with severe congenital anomalies (see Supplemental Information). We also excluded infants born before 25 weeks of gestation to minimize bias at the threshold of viability.¹⁴

Data for individual infants are linked such that they can be followed if transferred between CPQCC NICUs. Because patient transfers may bias NICU performance assessments, we developed detailed algorithms to avoid unduly crediting or penalizing NICUs for care delivered elsewhere. Guiding principles for these algorithms were as follows:

- Only infants with at most 3 admission records from 2 hospitals are included.
- 2. If the birth hospital transferred an infant by 3 days of age (day 1 being the day of birth), subsequent relevant outcomes (eg, chronic lung disease) accrue to the receiving hospital (counted as missing for the birth hospital).
- 3. If the birth hospital transferred an infant after 3 days of age, subsequent relevant outcomes accrue to the birth hospital (counted as missing for the receiving hospital).

Measures

See also Supplemental Table 3.

Outcome Variable

Baby-MONITOR: measures for the composite scale were selected by an expert panel¹⁵ and affirmed by practicing neonatologists.¹⁶ Measure definitions used standard CPQCC algorithms. The measures were expressed as binary variables at the patient level and as proportions at the unit level. They include (1) any antenatal steroid administration; (2) moderate hypothermia (<36°C) on admission; (3) non-surgically induced pneumothorax; (4) hospital-acquired bacterial or fungal infection; (5) oxygen requirement at 36 weeks' gestational age; (6) retinopathy of prematurity screening at the age recommended by the American Academy of Pediatrics (AAP); (7) discharge on any human milk; (8) mortality during the birth hospitalization; and (9) growth velocity (less or more than the median of 12.9 g/kg/day) calculated by using a logarithmic function.¹⁷

Variable of Interest: Level of Care

NICU level of care was derived as a self-reported variable derived from the 2012 Vermont Oxford Network Survey of NICU directors. Designations follow the 2012 definitions set forth by the AAP.¹⁸ This study included Level (L) II through IV NICUs.

Missing AAP levels and discrepancies were checked and confirmed with the NICUs. Four centers only provided the older AAP levels (eg, IIA, IIB), in which case we determined the new AAP level based on the ventilation duration, the number of cardiac surgeries, and care levels as designated by the California Children's Services.¹⁹

Additional Covariates

Organizational variables: hospital ownership (government, notfor-profit, for-profit, other) and neonatologist coverage (in-house or at home) were obtained from the 2012 Vermont Oxford Network Annual Survey of NICU directors. Hospital volume was obtained from the eligible infants from the study cohort in the CPQCC data. Hospital teaching status was derived from the Regional Perinatal Programs of California.²⁰

Clinical variables: these data were obtained from the CPQCC data set and included prenatal care, gender, weight for gestational age below the 10th percentile, outborn, multiple birth, 5-minute Apgar score, and Cesarean delivery. Gestational age at birth was categorized into 25 weeks to 27 weeks 6 days, 28 weeks to 29 weeks 6 days, and \geq 30 weeks gestation groups, based on similar patient numbers among groups. Apgar score was categorized as \leq 3, between 4 and 6, and >6. Prenatal care was defined as receipt of any prenatal obstetrical care before the admission during which birth occurred.

Analyses

Baby-MONITOR Scores

Computation of Baby-MONITOR scores requires that its subcomponents are aligned according to valence (higher score = better performance), risk adjusted, and standardized using the Draper-Gittoes method.^{12,21} With this method, a standardized observed minus expected *z* score was computed, with an expected mean of 0 and a SD of 1. Each *z* score was equally weighted and averaged to derive a Baby-MONITOR score for each NICU. We used bootstrapping (a simulation in which each NICU's patients were resampled with replacement 500 times²²) to compute 95% confidence intervals.

Association of Baby-MONITOR Scores With Level of NICU Care

We grouped NICUs according to their level of care and calculated Baby-MONITOR scores for each level weighted by number of infants. We used the *F* and *t* tests to assess differences in composite scores between NICU levels. To examine the effect of patient volume on quality of care delivery, we stratified the analyses according to VLBW volume using the cutoffs for high- and low-volume based on median annual volumes, achieving balance of NICUs within high- and lowvolume groups (ie, L II: 1-6 = low, >6= high; L III: 1–29 = low, >29 = high; L IV: 1–61 = low, >61 = high). These cut points are broadly consistent with those used in the literature, which had an empirical basis.^{5,8}

Controlling for the Effects of Organizational Variables

We performed a multivariate analysis regressing Baby-MONITOR score onto NICU level, controlling for other covariates. To choose the covariates for the final model, we used backward selection with a *P* value criterion of <.15.

Differences in Baby-MONITOR Subcomponents by Level of Care

We used analysis of variance to test for differences in performance on risk adjusted Baby-MONITOR subcomponent scores across levels of care. We used Bonferroni adjustment to correct for multiple testing.

Human Subjects Compliance

This study was approved by the Stanford Internal Review Board.

RESULTS

Sample Characteristics

The sample included 21 051 VLBW infants with 22 984 hospital records (transfers included) in 134 NICUs born between January 1, 2008, and December 31, 2012 who met the inclusion criteria. Of these NICUs, 25 are designated as L II, 89 as L III, and 20 as L IV.¹⁸ Approximately 4% of infants were born at L 1 hospitals, other outpatient setting, out of state, or military hospitals. Excluded from the analysis were 1194 infants (~5%) who were transferred to ≥ 3 institutions. Of these, nearly 70% received cared at L IV NICUs. Table 1 shows the unadjusted population and NICU characteristics for the combined sample. Approximately 5% of infants were born at an L II NICU (1012 of 21 051). L IV NICUs cared for a higher proportion of high-risk infants. On average, infants in L IV NICUs were of lower gestational age and their mothers were more likely to be of advanced maternal age and carrying multiples. In unadjusted analyses, L II NICUs exhibited significant opportunities for process improvement. They had lower rates of antenatal steroid administration, eye examinations, and any human milk feeding at discharge, and higher rates of hypothermia on admission. On the other hand, they exhibited lower rates among several outcome measures including rates of pneumothoraces, health care associated infections, chronic lung disease, and mortality (P < .05 for all comparisons).

Baby-MONITOR Scores Across NICUs

We found significant variation in Baby-MONITOR scores across California (mean [SD] 0.18 [1.14], range –2.26 to 3.39). Figure 1 shows a caterpillar plot of the Baby-MONITOR scores with NICUs

TABLE 1 Sample Characteristics

	All Admissions ($N = 22984$)		Level II ($N = 1012$)		Level III ($N = 15618$)		Level IV ($N = 6354$)	
Characteristics	n/N	%	n/N	%	n/N	%	n/N	%
Gestational age, wk	8927/22980	39	288/1012	28	5756/15615	37	2883/6353	45
≤27	6448/22980	28	253/1012	25	4528/15615	29	1667/6353	26
28–29	7605/22980	33	471/1012	47	5331/15615	34	1803/6353	28
30+								
Male gender	11987/22980	52	533/1012	53	8158/15617	52	3296/6351	52
Prenatal care	22 082/22 856	97	953/1006	95	15022/15546	97	6107/6304	97
Multiple gestation	6283/22984	27	208/1012	21	4237/15618	27	1838/6354	29
Cesarean delivery	16960/22983	74	717/1012	71	11624/15618	74	4619/6353	73
Small for gestational age	6094/22973	27	324/1012	32	4192/15615	27	1578/6346	25
Maternal age, y	2150/22962	9	127/1011	13	1458/15609	9	565/6342	9
Under 20								
20–29	9578/22962	42	435/1011	43	6549/15609	42	2594/6342	41
30—39	9748/22962	42	405/1011	40	6639/15609	43	2704/6342	43
40+	1486/22962	6	44/1011	4	963/15609	6	479/6342	8
Apgar 5 min								
≤3	1098/22843	5	49/1005	5	650/15547	4	399/6291	6
4–6	3654/22843	16	117/1005	12	2268/15547	15	1269/6291	20
≥7	18091/22843	79	839/1005	83	12629/15547	81	4623/6291	73
Outborn	3883/22984	17	206/1012	20	1541/15618	10	2136/6354	34
Baby-MONITOR Measures								
Antenatal corticosteroid								
administration	17757/21062	84	619/839	74	12 106/14 348	84	5032/5875	86
Moderate hypothermia on								
admission	3125/22682	14	213/996	21	2092/15392	14	820/6294	13
Pneumothorax	743/22973	3	13/1008	1	441/15612	3	289/6353	5
Any health care—associated								
infection	2431/21944	11	61/753	8	1497/14992	10	873/6199	14
Chronic lung disease at 36 wk								
gestational age	4379/20031	22	98/698	14	2851/13784	21	1430/5549	26
Timely eye exam	14 164/15 043	94	360/418	86	9677/10250	94	4127/4375	94
Any human milk at discharge	13 300/22 970	58	510/1010	50	9057/15611	58	3733/6349	59
In-hospital mortality	1452/22966	6	15/1012	1	889/15618	6	548/6336	9
High growth velocity	9996/19993	50	369/701	53	7175/13733	52	2452/5559	44

ordered with regard to ascending composite score for the clinical measures. We show both a figure based on the standard units (Fig 1A) and a conversion to percentiles (Fig 1B). The variation in performance between these NICUs was highly significant in practical terms (indicated by the 5.65 standard units of difference between the top and bottom providers). These results were robust with regard to changing the transfer cutoff days from a baseline of day 3 to scenarios including transfer on days 2 and 4 of age, as well as assigning outcomes for all transfers to the birth hospital. Finally, we included all deaths before 12 hours of age in the analysis. The correlation in Baby-MONITOR scores between these scenarios was high, ranging from 0.94 to 0.99, consistent

with our previous work²³ (see online Supplemental Table 4).

Level of Care and Baby-MONITOR Scores

On average, L III NICUs achieved the highest Baby-MONITOR scores (L III mean [SD (range)] 0.43 [1.35 (-2.26 to 2.64)], L IV 0.37 [1.39 (-1.61 to 3.39)], L II -0.22 [0.89 (-1.82 to 1.23]), but these differences were not statistically significant (P = .53). Stratification (Fig 2) revealed a VLBW volume effect that widened with increasing level of care (L II low 0.15 [0.5] versus high -0.3 [0.93]; L III low 0.15 [1.02] versus high 0.52 [1.43]; L IV low -0.08 [1.08] versus high 0.52 [1.45]). Neither these differences nor any of the associations of Baby-MONITOR scores with organizational variables,

including hospital ownership, neonatologist coverage, and hospital teaching status, reached statistical significance (see Supplemental Information, Sensitivity Analysis).

Level of Care and Baby-MONITOR Subcomponents

Figure 3 shows significant differences across levels of care for several subcomponents with L IV NICUs scoring higher on several process measures of care, including antenatal steroids (P = .002) and any human milk at discharge (P = .092), but lower on other outcomes such as health care-associated infections (P = .040), pneumothorax (P < .001), and growth velocity (P = .006).

Table 2 also shows pairwise comparisons using L IV NICUs as

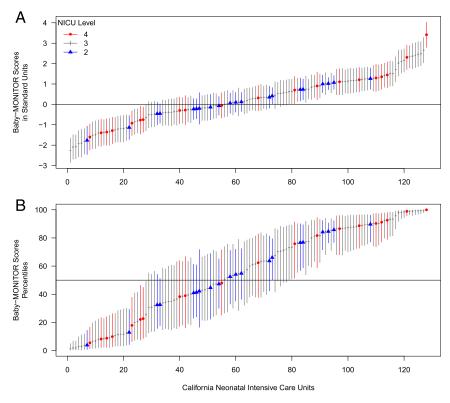
a reference. Compared with L III NICUs, they had higher rates of antenatal steroids (P = .040), any human milk at discharge (P = .030), and survival (P = .045), but also of health care-associated infections (P = .014) and poor growth (P = .002). After Bonferroni adjustment, survival and human milk at discharge were no longer significant.

Compared with L II NICUs, we found higher rates of antenatal steroids (P = .036), pneumothorax (P = .012) and a trend toward higher retinopathy of prematurity examinations (P = .099). After Bonferroni adjustment, only pneumothorax remained significant.

DISCUSSION

Using population-based data, we present a multidimensional, nuanced assessment of the relation between quality of NICU care provided to VLBW infants and NICU level of care. We found significant variation in Baby-MONITOR scores across NICUs but no statistically significant association with level of care. Subcomponent analysis revealed interesting differences, with L IV NICUs performing better on process measures, as well as marginally on survival, and L II NICUs better on other outcome measures.

We consider 4 potential causes to explain our findings. First, previous literature and general advances in high-risk maternal care, including greater use of antenatal corticosteroids²⁴ and imaging, may have fostered more appropriate utilization and regionalization patterns. Compared with previous studies,^{3,4} we found an inconsistent association between level of care and survival. The proportion of infants born in L II NICUs is low (5%), and case mix is favorable to survival. Thus, selection bias may have impeded our ability to demonstrate significant differences in survival of infants in L II compared with L IV NICUs. Consistent with previous





Baby-MONITOR scores for 132 California NICUs. The different colors and symbols designate different levels of NICUs, and the stars designate low VLBW infant volume L IV NICUs. A, Mean (95% confidence interval) expressed in observed minus expected *z* scores, measured in standard units. B, Information expressed as percentiles of the distribution of ranking of the NICUs against each other. This illustration highlights the relative uncertainty in NICU rankings. For instance, the lowest-ranking NICU has all of its vertical line close to the 0th percentile, meaning that we are confident that its Baby-MONITOR scores are much lower than those of NICUs whose performance is near the 50th percentile. The uncertainty regarding NICU performance is much greater in the middle. Only NICUs with a minimum of 10 infants are shown.

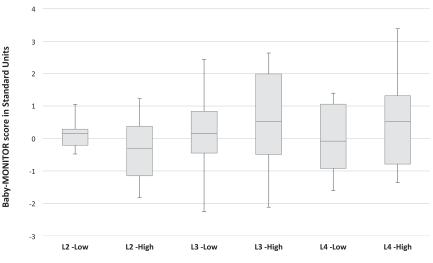


FIGURE 2

Box-and-whisker plot of Baby-MONITOR scores by NICU level, stratified by high and low volume according to the 50th percentile for each NICU level (Level II: 1-3 = 1 ow, >3 = high; Level III: 1-23 = 1 ow, >23 = high; Level IV: 1-48 = 1 ow, >48 = high). Horizontal line in a boxplot indicates a weighted mean of the sample.

^a Statistical significance after Bonferroni adjustment for multiple testing.

Outcome Measure	Level	Ν	Mean	SD	Range	Р	Type III P
Antenatal steroid	2	7	-1.12	1.54	-2.49 to 1.54	.036	.002
	3	86	0.32	2.91	-10.9 to 5.76	.0008ª	
	4	15	2.7	3.4	-2.39 to 7.46	Ref	
	Overall	108	0.85	3.19	-10.9 to 7.46		
No hypothermia on admission	2	7	-2.86	3.23	-7.34 to 2.61	.124	.286
	3	87	1.2	4.66	-13.02 to 9.49	.466	
	4	20	2.03	6.76	-8.2 to 11.02	Ref	
	Overall	114	1.32	5.37	-13.02 to 11.02		
Any human milk at discharge	2	7	-0.46	2.42	-4.2 to 4.41	.430	.092
	3	87	-0.44	4.06	-12.8 to 6.78	.030	
	4	20	1.6	5.26	-7.75 to 10.98	Ref	
	Overall	114	0.13	4.49	-12.79 to 10.98		
Timely eye examination	2	7	-1.71	1.88	-4.63 to 1.14	.099	.248
	3	87	0.71	2.44	-8.21 to 5.32	.853	
	4	20	0.81	2.88	-3.39 to 5.12	Ref	
	Overall	114	0.67	2.59	-8.21 to 5.32		
Survival	2	7	1.1	1.38	-1.19 to 2.75	.659	.083
	3	87	-0.11	1.78	-3.61 to 4.33	.045	
	4	20	0.64	1.73	-1.64 to 4.12	Ref	
	Overall	114	0.13	1.8	-3.61 to 4.33		
No chronic lung disease	2	7	-0.84	1.81	-3.51 to 2.17	.583	.792
-	3	87	0.35	3.08	-5.96 to 8.11	.800	
	4	20	0.18	3.32	-6.86 to 4.76	Ref	
	Overall	114	0.27	3.13	-6.86 to 8.11		
No health care—associated infection	2	7	-0.67	2.79	-5.67 to 2.22	.967	.040
	3	87	0.65	2.58	-6.42 to 6.81	.014ª	
	4	20	-0.73	2.65	-5.86 to 4.13	Ref	
	Overall	114	0.22	2.68	-6.42 to 6.81		
No pneumothorax	2	7	1.16	0.68	-0.39 to 2.19	.012	<.0001ª
	3	87	0.48	1.56	-3.15 to 3.74	<.001ª	
	4	20	-1.14	1.51	-3.15 to 2.8	Ref	
	Overall	114	0.05	1.7	-3.15 to 3.74		
High growth velocity	2	7	0.93	2.67	-2.3 to 4.47	.215	.006
	3	87	0.71	4.23	-10.92 to 10.28	.002ª	
	4	20	-2.12	4.0	-9.1 to 6.28	Ref	
	Overall	114	-0.08	4.32	-10.92 to 10.28		

TABLE 2 Baby-MONITOR Subcomponents by AAP Level Outcome Measure

3

2

0

FIGURE 3

analysis of variance.

Baby-MONITOR Scores in Standard Units -1 ** -2 ** -3 High growth Antenatal No Any breast milk at Timely eye Survival No chronic No health care No associated peumothorax steroids hypothermia examination lung disease velocity discharge infection

Baby-MONITOR subcomponents by NICU AAP level of care. Statistical significance is derived by

Subcomponents by AAP Level

** P < .05 * P < .1

■ III ■ IV

research, we found a borderline survival benefit of L IV compared with L III NICUs. However, this difference was not significant after adjustment for multiple comparisons. Given differing biases of providers or parents for life-sustaining treatments, survival may not accurately reflect actual quality of care delivery. We think our results should be viewed as supporting continued national efforts to limit VLBW births in L II NICUs and for regionalized care delivery.²⁵

Second, the current approach to defining level of care, as well as selfdesignation of this variable, may lead to misclassification and dilute the association with measures of quality.

However, using California-specific NICU designations assigned by the state also did not result in significant associations with Baby-MONITOR scores (see Supplemental Information).

Third, L II NICUs did achieve lower scores on many process measures, indicating opportunity for quality improvement, yet they also achieved higher scores for many outcome measures. These findings might be the result of selection bias not adequately mitigated by risk adjustment. For example, growth velocity is difficult to predict using patient characteristics from the immediate peripartum time period. Future ability to extract additional data from the electronic record may help refine risk models. In addition, pseudo-randomization methods, such as an instrumental variable approach, may address some of the unobserved selection bias. This requires additional study, but previous applications of these methods to NICU outcomes have demonstrated that the benefits of care at higher-volume and/or higher-level NICUs are larger than with traditional risk-adjustment methods such as those that we used.26

Fourth, transfer bias may have depressed scores for higher-level NICUs. Outcomes for L II NICUs are measured not according to birth at such a facility but according to intent to keep such infants at a L II for treatment. However, we were careful to mitigate transfer bias by including inborn status in risk adjustment models and by assigning negative outcomes of care for infants transferred after day of age 3 to the sending NICU (outcome is missing for receiving NICU, yet positive outcome is assigned to both NICUs). We did assign negative outcomes of infants transferred before or on day of age 3 to the receiving NICU. In addition, assigning all outcomes of transferred infants back to the birth hospital also did not have significant influence on our results. Finally, there is a known inverse relation between the volume

of high-risk deliveries and in-hospital fetal death rates that may be associated with the ability to perform rapid cesarean deliveries.³ This can cause a bias because fetal deaths are not included in our data and many of the cases in which the fetal death is "averted" in the high-volume hospitals will have elevated risks not captured by our data.

This study provides a good example for the usefulness of composite indicators. The composite provides a global picture of differences in quality of care and of the association with important predictors of quality. Conversely, drawing inferences on overall care based on a single measure, such as mortality, is hazardous because individual measures contain biases, making them nonrepresentative. In addition, we have previously shown that NICUs that perform well in 1 area of care may not perform well in others.

Equally important is the process of drilling down into individual subcomponents of the composite because averaging across the measures may hide important differences. This study exemplifies this by revealing important and modifiable differences between NICUS.^{27,28}

This study must be viewed within the context of its design. Observational studies allow for the establishment of associations and the generation of hypotheses but not causal inference. In addition, as mentioned above, incomplete risk adjustment and transfer bias and confounding from unobserved variables (eg, patient-tonurse ratios) might have affected our findings. Nevertheless, these methods have been previously published, and inclusion of institutional confounding variables may not be appropriate for quality of care comparisons. This study included nearly all of the NICUs in California, the country's most populous state with diverse geography. Given our objective to study the effect of care organization on quality, our findings may have

broad relevance to other regions in the United States and abroad. Finally, we used only a 1-time designation of NICUs in 2012 of their level of care and applied this designation to the entire study period. Because the AAP designation changed in 2012, we do not have earlier designations based on this classification scheme. However, examining changes in classification over previous years, we found them to be highly stable. Because changes in level of care usually occur toward a higher level, this limitation would bias our results toward the null.

CONCLUSIONS

In this population-based study, we found wide variation in overall quality of care provided to VLBW infants by using the Baby-MONITOR, but no significant associations with NICU level of care. We did, however, find important associations with its subcomponents, with L IV NICUs receiving higher-quality scores for measures of care process, and L II NICUs receiving higher scores for several care outcomes. These findings highlight opportunities for further improvements that can be addressed through targeted interventions.

ACKNOWLEDGMENTS

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ABBREVIATIONS

AAP: American Academy of Pediatrics CPQCC: California Perinatal Quality Care Collaborative VLBW: very low birth weight PEDIATRICS (ISSN Numbers: Print, 0031-4005; Online, 1098-4275).

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