

NT-pro BNP—A marker for worsening respiratory status and mortality in infants and young children with pulmonary hypertension

Shahnawaz M. Amdani MBBS, MD¹  | Muhammad Umair M. Mian MD² |
Ron L. Thomas PhD³ | Robert D. Ross MD⁴

¹Division of Pediatric Cardiology, St. Louis Children's Hospital, Washington University School of Medicine, St. Louis, Missouri, USA

²Division of Pediatrics, The Carman and Ann Adams Department of Pediatrics, The Children's Hospital of Michigan, Wayne State University School of Medicine, Detroit, Michigan, USA

³Division of Biostatistics, The Carman and Ann Adams Department of Pediatrics, The Children's Hospital of Michigan, Wayne State University School of Medicine, Detroit, Michigan, USA

⁴Division of Pediatric Cardiology, The Carman and Ann Adams Department of Pediatrics, The Children's Hospital of Michigan, Wayne State University School of Medicine, Detroit, Michigan, USA

Correspondence

Robert D. Ross, Division of Cardiology, Carman and Ann Adams Department of Pediatrics, The Children's Hospital of Michigan, 3901 Beaubien Blvd, Detroit, MI 48201-2119.

Email rross@dmc.org

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Abstract

Aim: To evaluate predictors of morbidity and mortality in pediatric patients with pulmonary hypertension (PH), laboratory and echocardiographic measures of PH were analyzed.

Methods: A retrospective review of all infants and children < 2 years of age with PH from January 2011 to August 2016 was conducted. Correlations were determined using Spearman's rank correlation coefficients. Differences in characteristics between survivors and nonsurvivors were analyzed and Kaplan-Meier survival curves were generated.

Results: Of 56 patients, the majority were extremely premature; of African American ethnicity; and had bronchopulmonary dysplasia. Patients who died were more likely to have underlying congenital heart disease; have a higher increase in the concentration of carbon dioxide in the blood (pCO₂) with a corresponding greater mean percentage decrease in pH and percentage rise in NT-pro BNP during PH exacerbations; more likely to have been on medications for pulmonary hypertension; and have a higher RVSP/SBP (%) ratio and S/D ratio. There were positive correlations between percentage rise in NT-pro BNP and pCO₂; NT-pro BNP and RVSP/SBP (%) ratio; and RVSP/SBP (%) ratio and S/D ratio.

Conclusions: Infants and young children with pulmonary hypertension have increased morbidity and mortality. NT-pro BNP is a useful biomarker for both respiratory exacerbations and mortality, and RVSP/SBP (%) ratio and S/D ratio are echocardiographic identifiers for increased mortality.

KEYWORDS

morbidity, mortality, NT-pro BNP, pCO₂, pulmonary hypertension

1 | INTRODUCTION

Brain natriuretic peptide (BNP) is an amino acid that is secreted in response to volume or pressure overload in the heart.^{1,2} BNP counteracts the renin angiotensin system and causes diuresis, natriuresis, and systemic (arterial and venous) vasodilatation.³ This in turn relieves the pressure and volume load to the ventricles. Both BNP and its precursor N terminal—pro BNP (NT-pro BNP) have been used extensively in the adult population to identify patients with myocardial dysfunction,^{2,4} and to gauge response to anticongestive treatment⁵ and overall outcomes.

Since BNP has a shorter half-life (~ 20 minutes) compared to NT-pro BNP (~ 1–2 hours),⁶ NT-pro BNP may reflect more consistent concentrations to evaluate pressure/volume overload to the ventricles. These cardiac biomarkers have been used in children as well and have correlated with ventricular dysfunction and mortality in patients with heart failure.^{1,7} Moreover, they have been used to predict low cardiac output syndrome (LCOS), and postoperative mortality in patients undergoing cardiac surgery.

At birth, the values of both BNP and NT-pro BNP rise and then decline in early infancy.⁸ It has been found that BNP and NT-pro BNP

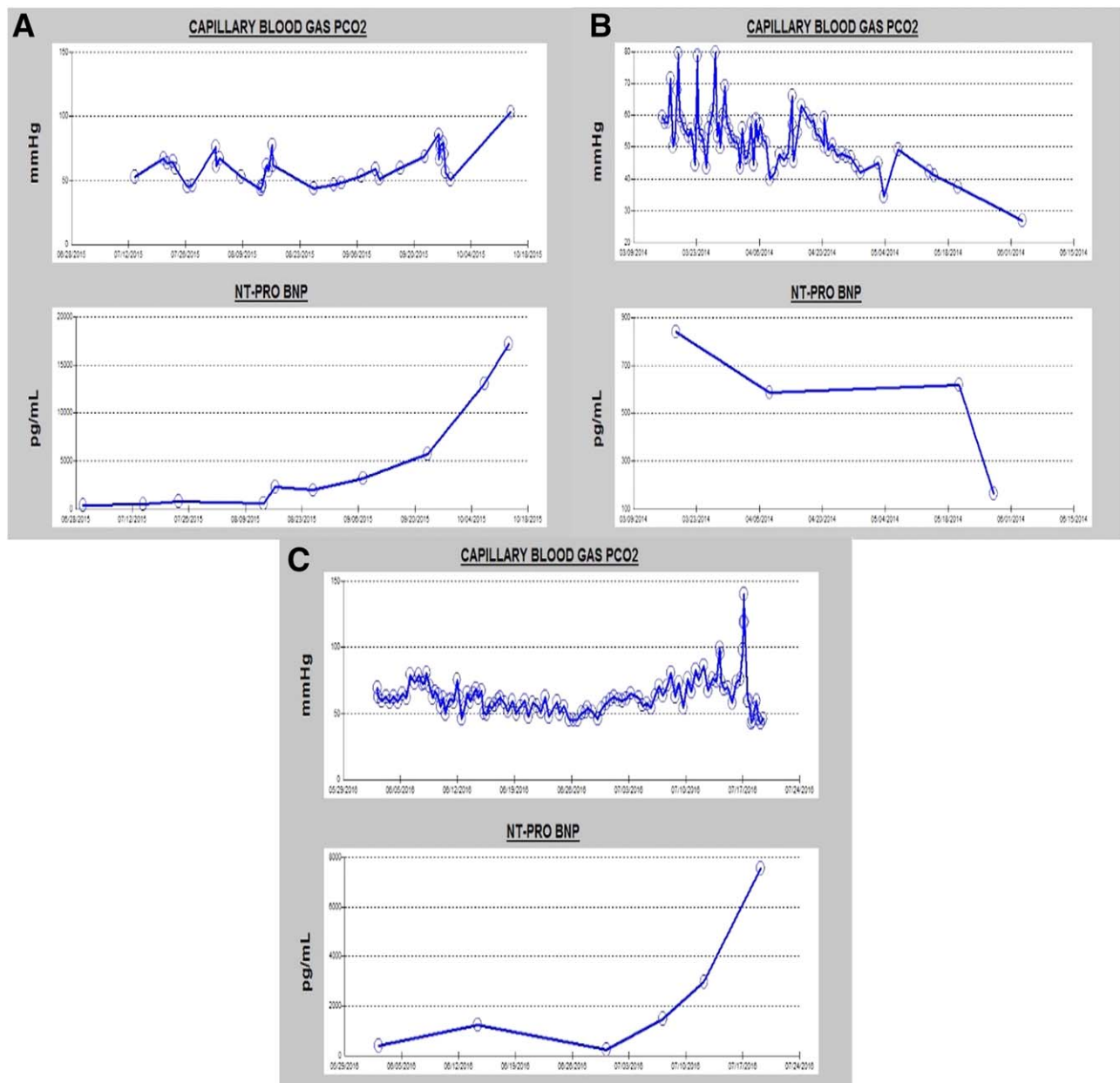


FIGURE 1 (A-C) These are graphs showing the relationship between $p\text{CO}_2$ and NT-pro BNP levels in three pediatric patients with pulmonary hypertension

levels are higher in neonates with a hemodynamically significant patent ductus arteriosus (PDA),⁹ bronchopulmonary dysplasia (BPD),¹⁰ and persistent pulmonary hypertension (PH) of the newborn.¹¹

The purpose of this study was to evaluate predictors of inpatient morbidity and mortality in infants and young children with PH requiring hospitalization. Over the last several years, author RDR observed a trend for an increase in NT-pro BNP concentrations with exacerbations in respiratory status in children with pulmonary hypertension (Figure 1A-C). The hypothesis generated from these observations was that there is a correlation between acute rises in the concentration of carbon dioxide in the blood ($p\text{CO}_2$) as a surrogate for an exacerbation of respiratory status (as in acute worsening of PH, new atelectasis, pneumonia, pneumothorax, or pleural effusion) and NT-pro BNP

concentrations. Therefore, the aim of this study was to assess the relationship between laboratory and echocardiographic measures of pulmonary hypertension and respiratory status as indicated by changes in PCO_2 and pH and to assess these variables as predictors of mortality.

2 | METHODS

2.1 | Study population

The electronic database was queried to identify all infants and children < 2 years from January 1, 2011 to August 31, 2016 who were hospitalized in the neonatal or pediatric intensive care unit with a diagnosis of pulmonary hypertension. A detailed chart review was

conducted to obtain **demographic data** (age, sex, ethnicity); **clinical information** (height, weight, body surface area [BSA], underlying diagnosis, systolic blood pressure, total hospital length of stay, history of intubation, duration of intubation, history of tracheostomy, history of extracorporeal membrane oxygenation [ECMO] use, discharge mortality status), and **medication history** (use of sildenafil, milrinone, nitric oxide). Patients ≥ 2 years of age; those with incomplete documentation on review of medical records or incomplete echocardiographic information were excluded from the study. Approval of the study and waiver of consent were obtained from the Wayne State University Institutional Review Board and Children's Hospital of Michigan.

2.2 | Laboratory data

All NT-pro BNP values that were collected were included for analysis. These were obtained at the discretion of the physician caring for the child or at the request of the PH consultant. To avoid using isolated laboratory values which might not reflect true trends, we collected 48-hour (24 hours before and after NT-pro BNP collection) pH and pCO₂ values, to correlate with respiratory status around the times of NT-pro BNP sampling. The lowest values, highest values and 48-hour averages of pH and pCO₂ were subsequently calculated. The percentage change in NT-pro BNP value during PH exacerbation was calculated. Also, percentage changes in pCO₂ and pH were calculated during the time NT-pro BNP was collected to capture acute changes in respiratory status.

2.3 | Echocardiography data

Transthoracic two-dimensional and Doppler echocardiographic studies were performed using a Philips Sono IE 33 machine (Philips Inc Andover, MA). Only studies completed within 72 hours of the NT-pro BNP collection were included for analysis to achieve best possible correlation with echocardiographic parameters. A single pediatric cardiologist, who was blinded with respect to respiratory status, laboratory results and the patients' prior echocardiogram reports and diagnoses, retrospectively reviewed all echocardiograms. Echocardiographic two-dimensional apical views were used to assess the right ventricle as described by prior studies.^{12,13} In the presence of tricuspid regurgitation (TR) and absence of right ventricular outflow tract obstruction, apical or parasternal long axis views were used to estimate systolic pulmonary artery pressure using modified Bernoulli's equation.¹⁴ The highest estimate of velocity obtained was used for the study. RV systolic function was assessed by utilizing Tricuspid annular plane systolic excursion (TAPSE)¹⁵ and systolic to diastolic time (S/D) ratio was evaluated using methods described previously.^{16,17}

2.4 | Statistical analysis

Descriptive statistics were calculated using mean and 95% confidence intervals or medians and interquartile ranges for continuous variables and percentages for categorical variables. Patients in our study were divided into two groups—survivors (Group 1) and nonsurvivors (Group 2). Student's *t* test was used for normally distributed numeric variables

in the intergroup analysis, Mann-Whitney *U* test was used for non-normally distributed numeric variables, and chi-square test was used for categorical variables in the intergroup analysis. Correlations between various laboratory (NT-pro BNP, pCO₂) and echocardiographic parameters [RVSP/SBP (%) ratio, S/D ratio] were determined using Spearman's rank correlation coefficients. ROC curves were constructed for four variables (highest NT-pro BNP value [pg/mL]; % change in NT-pro BNP value; RVSP/SBP ratio [%]; and S/D ratio), and cutoffs with highest sensitivity and specificity identified. Subsequent Kaplan-Meier analysis with log-rank test was utilized to analyze patient survival for these variables. A *P* value of $< .05$ was considered statistically significant. Statistical analysis was performed using SPSS Version 24.0 (IBM Inc, Chicago, Illinois).

3 | RESULTS

3.1 | Baseline characteristics

Of 151 charts reviewed, 56 met inclusion criteria (80 were > 2 years of age, 10 with missing laboratory values, 5 with echocardiograms not obtained within 72 hours of the laboratory values). The majority of the patients were extremely premature (< 28 weeks gestation) at birth (28/56, 50%); of African American ethnicity (37/56, 66.1%), and there was an equal gender distribution in the study group. (Table 1) The most common underlying diagnosis was bronchopulmonary dysplasia (35/56, 62.5%) followed by congenital heart disease (4/56, 7.1%). In the patients with PH, various forms of support were needed—intubation and mechanical ventilation (49/56, 87.5%); tracheostomy (23/56, 41.1%); sildenafil (42/56, 75%); milrinone (23/56, 41.1%); nitric oxide (32/56, 57%); and ECMO (4/56, 7%). The right ventricular systolic pressure was severely increased in 21/56 (37.5%) of the population. Of the 56 patients in the study, 9 (16.1%) patients died.

3.2 | Comparing pediatric patients with pulmonary hypertension who survived (Group 1) versus nonsurvivors (Group 2)

There was no difference between the 2 groups with regards to gestational age at birth, height, weight, body surface area, or ethnicity (Table 2). Although females with PH had a higher likelihood of dying (77.8 vs 44.7%), this did not achieve statistical significance. There was higher mortality in patients with PH who had congenital heart disease compared to those with structurally normal hearts (33.1 vs 2.1%, $P = .01$). The difference in percentage rise in NT-pro BNP during an acute fall in pH was significantly higher in Group 2 ($U = 35$, Z -score -3.02 , $P < .01$). Moreover, the patients who died had significantly higher mean percentage decrease in pH and higher increase in pCO₂ during an exacerbation in respiratory status (14 ± 32 vs 2.5 ± 2.1 , $P = .01$; 406 ± 247 vs 208 ± 150 , $P < .05$). Patients who died were more likely to have been on pulmonary hypertension medications—sildenafil (100 vs 70.2%, $P < .05$), milrinone (77.8 vs 34%, $P < .05$), and nitric oxide (88.9 vs 51.1%, $P < .05$).

Echocardiographic evaluation revealed that patients who died had a significantly higher RVSP/SBP (%) ratio (76 ± 37 vs 46 ± 43 , $P = .05$)

TABLE 1 Baseline characteristics of study cohort

	n = 56 Mean (95% CI), Median (IQR) or n (%)
Gestational age at birth (weeks)	
< 28	28 (50)
28-< 32	4 (7.1)
32-37	4 (7.1)
> 37	20 (35.7)
Height (cm)	51 (37.8-64.1)
Weight (kg)	3.9 (1.2-6.6)
BSA (m ²)	0.23 (0.13-0.33)
Gender	
M	28 (50)
F	28 (50)
Ethnicity	
African American	37 (66.1)
Caucasian	15 (26.8)
Hispanic/other	4 (7.1)
Underlying Diagnosis	
BPD	35 (62.5)
Congenital diaphragmatic hernia	2 (3.6)
Down syndrome	1 (1.8)
Congenital heart disease	4 (7.1)
Other (skeletal anomalies, omphalocele, other syndromes)	14 (25)
Support needed	
Intubated	49 (87.5)
Tracheostomy	23 (41.1)
Sildenafil	42 (75)
Milrinone	23 (41.1)
Nitric oxide	32 (57)
ECMO	4 (7)
Length of ventilation	70 (41-99)
Length of hospital stay	140 (110-170)
RVSP	
Mildly increased	26 (46.4)
Moderately increased	9 (16.1)
Severely increased	21 (37.5)
Patient status at discharge	
Alive	47 (83.9)
Dead	9 (16.1)

and S/D ratio (2.2 ± 0.9 vs 1.6 ± 0.4 , $P < .01$). Moreover, these Group 2 patients were more likely to have right to left or bidirectional shunting compared to Group 1 ($P < .05$).

3.3 | ROC curves and survival analysis

ROC curve analysis identified the following optimal cut-off points: highest NT-pro BNP value (pg/mL)—8915; % change in NT-pro BNP value during PH exacerbation—1291; RVSP/SBP (%) ratio—64; S/D ratio—1.7.

Patients with NT-pro BNP values > 8915 pg/mL had worse survival compared to patients with NT-pro BNP values ≤ 8915 pg/mL [13

(8.5–17.5) vs 20.3(17.9–22.6) months, $P = .01$] (Figure 2). Patients with increase in NT-pro BNP values during pH exacerbation of $\geq 1291\%$ tended to have worse survival compared to patients with a change $< 1291\%$ ($P = .12$) (Figure 3). Also, patients with RVSP/SBP (%) ratio ≥ 64 tended to have worse survival compared to patients with

TABLE 2 Comparison of survivors to nonsurvivors

	Group 1 Survivors n = 47 Mean \pm SD or n (%)	Group 2 Nonsurvivors n = 9 Mean \pm SD or n (%)	P value
Gestational age at birth (weeks)			
< 28	24 (51.1)	4 (44.4)	NS
28 to < 32	3 (6.4)	1 (11.1)	
32-37	4 (8.5)	0 (0)	
> 37	16 (34)	4 (44.4)	
Height (cm)	51.6 \pm 11.7	54.6 \pm 11	NS
Weight (kg)	4.1 \pm 2.4	4.8 \pm 2.7	NS
BSA (m ²)	0.24 \pm 0.1	0.27 \pm 0.1	NS
Gender			
M	26 (55.3)	2 (22.2)	NS
F	21 (44.7)	7 (77.8)	
Ethnicity			
African American	32 (68.1)	5 (55.6)	NS
Caucasian	13 (27.7)	2 (22.2)	
Hispanic/other	2 (4.2)	2 (22.2)	
Underlying diagnosis			
Congenital heart disease	1 (2.1)	3 (33.3)	
Other (BPD, congenital diaphragmatic hernia, Down syndrome, skeletal anomalies, omphalocele, other syndromes)	46 (97.9)	6 (66.7)	.01
Highest NT-pro BNP during PH exacerbation (pg/ml)	6885 \pm 9204	16148 \pm 9669	< .05
% decrease in pH during PH exacerbation	2.5 \pm 2.1	14 \pm 32	.01
% increase in pCO ₂ during PH exacerbation	208 \pm 150	406 \pm 247	< .05
Length of ventilation	49 \pm 90	116 \pm 74	< .05
Support needed			
Intubated	40 (85.1)	9 (100)	NS
Tracheostomy	19 (40.4)	4 (44.4)	NS
Sildenafil	33 (70.2)	9 (100)	< .1
Milrinone	16 (34)	7 (77.8)	< .05
Nitric oxide	24 (51.1)	8 (88.9)	< .05
ECMO	3 (6.4)	1 (11.1)	NS
Echocardiographic parameters			
S/D ratio	1.6 \pm 0.4	2.2 \pm 0.9	< .01
Maximum RVSP (mm Hg)	56 \pm 21	73 \pm 22	< .1
RVSP/SBP ratio (%)	46 \pm 43	76 \pm 37	.05
Atrial level shunting			
Left to right	36 (76.6)	3 (33.3)	< .05
Right to Left	1 (2.1)	1 (11.1)	
Bidirectional	10 (21.3)	5 (55.6)	

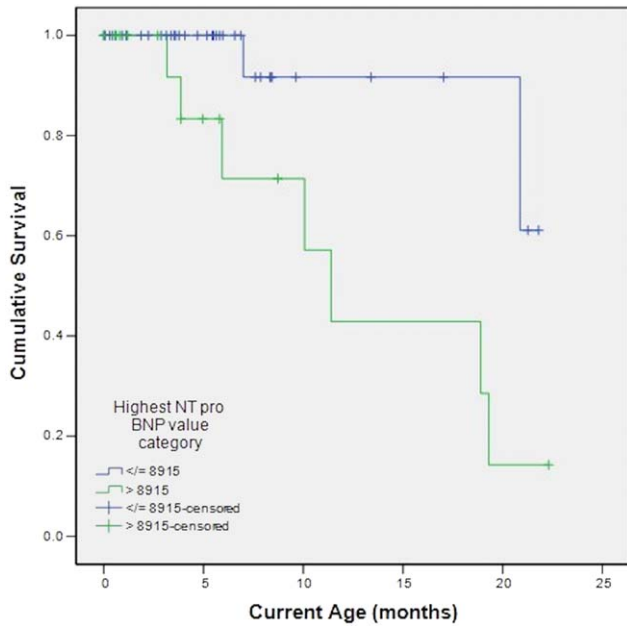


FIGURE 2 Kaplan-Meier survival curve comparing patients with highest NT-pro BNP values > 8915 pg/mL vs ≤ 8915 pg/mL

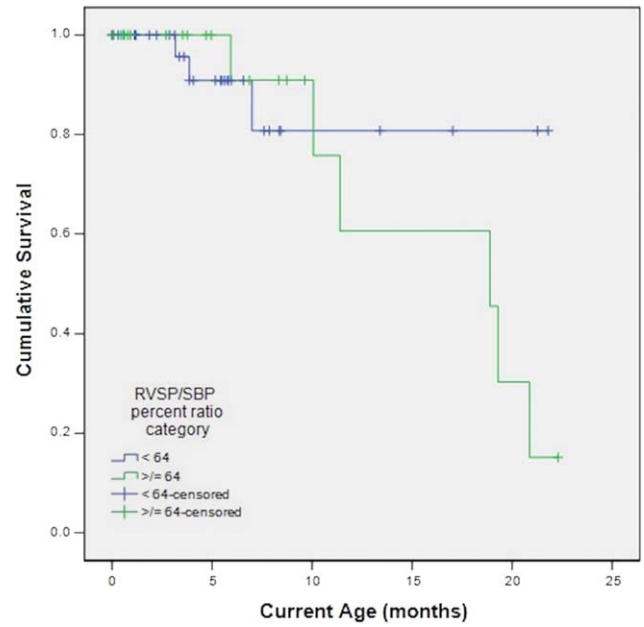


FIGURE 4 Kaplan-Meier survival curve comparing patients with RVSP/SBP ratio $\ge 64\%$ vs $< 64\%$

RVSP/SBP (%) ratio < 64 (16.1 [12.2–20] vs 18.6 [15.2–22.1] months, $P = .38$) (Figure 4). In addition, those with S/D ratio > 1.7 had worse survival compared to patients with S/D ratio ≤ 1.7 (13.4[9.3–17.6] vs 21.5[19.9–23.1] months, $P < .01$) (Figure 5).

There was a significant positive correlation between percentage rise in NT-pro BNP and percentage rise in pCO_2 during exacerbations in respiratory status ($r = 0.40$, $P = .01$). There was a similar correlation between NT-pro BNP value and RVSP/SBP (%) ratio ($r = 0.44$, $P \le .01$); and RVSP/SBP ratio and S/D ratio ($r = 0.42$, $P < .01$).

4 | DISCUSSION

With the growing number of extremely premature births surviving the neonatal period, there has been a steady increase in long-term survivors with bronchopulmonary dysplasia (BPD).^{18,19} This increased incidence of BPD is secondary to prolonged mechanical ventilation required by the premature neonates due to respiratory distress syndrome and deficient surfactant levels. The effect of this is chronic barotrauma, frequent pneumonias and other respiratory complications in these extremely premature infants. In this study, half of the population

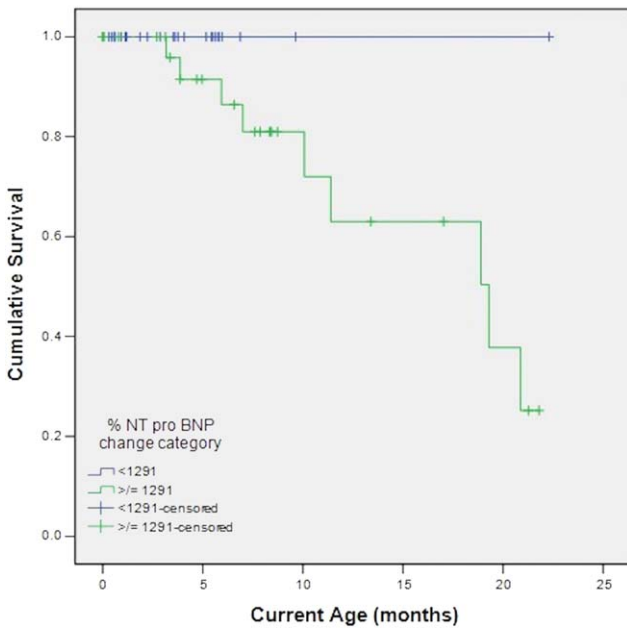


FIGURE 3 Kaplan-Meier survival curve comparing patients with increase in NT-pro BNP value during pH exacerbation $\ge 1291\%$ vs $< 1291\%$

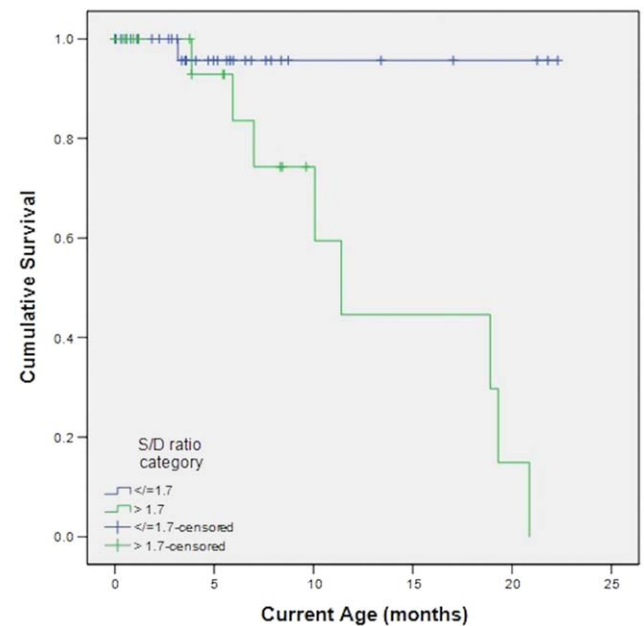


FIGURE 5 Kaplan-Meier survival curve comparing patients with S/D ratio > 1.7 vs ≤ 1.7

was extremely premature who are at higher risk for developing pulmonary hypertension secondary to BPD.^{20,21}

Children with PH have increased morbidity and mortality compared to those without PH.²² Of 56 children with PH in this study, most required some support to help them through the acute phase of PH exacerbation in the form of mechanical ventilation and pulmonary vasodilator medications. The patients who were on these types of support were more likely to die, particularly those who required milrinone and inhaled nitric oxide. The mortality overall was ~16%. Other studies have shown similar high mortality rates in children with pulmonary hypertension, with the highest risk in those with congenital diaphragmatic hernia.^{21,23,24}

BPD was the major reason for PH in this study. Although congenital heart disease (CHD) was much less common in this cohort, its presence was associated with a ~16-fold higher mortality. In the latest classification of pulmonary hypertension in children, CHD has been labeled as an independent diagnosis. Certain types of CHD increase the propensity for developing PH in the neonatal period like those with obstructed total anomalous pulmonary venous return, scimitar syndrome and hypoplastic left heart syndrome with a restrictive atrial septal communication. It has been shown that PH in these patients increases morbidity and mortality.^{25–27}

Patients in this study who died had ~2.5-fold higher peak NT-pro BNP than those who survived. A higher level of this biomarker suggests worsening right ventricular pressure overload as is seen in patients with worsening pulmonary hypertension. Other investigators have corroborated this finding.^{28–30} In this study, patients with a peak NT-pro BNP > 8915 pg/mL had significantly higher mortality. These values are lower than those presented by Baptista et al³¹ in pediatric patients with congenital diaphragmatic hernia. In their study, patients with NT-pro BNP value > 11 500 pg/mL had worse prognosis. Also, the nonsurvivors had higher % change in NT-pro BNP change during PH exacerbation and a higher drop in pH secondary to an acute rise in pCO₂. Moreover, there was a significant positive correlation between peak NT-pro BNP concentration and peak pCO₂. This confirms the observation that NT-pro BNP is influenced by changes in the underlying respiratory status. The correlation here was modest ($r = 0.4$) in part because NT-pro BNP is influenced by other factors including heart failure, blood pressure, heart rate, glomerular filtration rate, and certain medications.³²

Evaluating the echocardiographic parameters nonsurvivors had a higher RVSP/SBP (%) ratio and a higher S/D ratio. Studies evaluating these parameters in patients with PH have shown similar results.^{16,17,33} The S/D ratio is a marker of systolic function and is elevated in patients with pulmonary hypertension.^{16,34} It is particularly useful in patients with PH who do not have a complete tricuspid regurgitation jet. In our study, children with a S/D ratio > 1.7 had significantly higher mortality which is similar to the study by Alkon et al¹⁶ evaluating S/D ratio in 47 children with PH. They found there was worse RV function, 6-minute walk distance, hemodynamic data, and higher mortality in patients with S/D ratio > 1.4.

The results of this study confirm the hypothesis that NT-pro BNP increases with acute worsening of the respiratory status in infants and young children with PH. Moreover, patients that subsequently died

had higher NT-pro BNP concentrations and worse echocardiographic parameters of PH than long-term survivors.

5 | STUDY LIMITATIONS

This was a retrospective study with a relatively small sample size. The NT-pro BNP levels were drawn for clinical indications and thus not at standardized intervals. To minimize this possible confounder, only blood gas and echocardiographic measurements done at specified times around the blood draws were analyzed. Also, this was a single center study, which might limit its generalizability. Due to the limited sample size, we were not able to achieve statistically significant cut-off values for % change in NT-pro BNP and RVSP/SBP (%) despite the fact that there was a trend toward worse survival in patients with %NT-pro BNP change $\geq 1291\%$ and RVSP/SBP (%) ratio ≥ 64 .

6 | CONCLUSIONS

Infants and young children with pulmonary hypertension have increased morbidity and mortality. NT-pro BNP is a useful biomarker for respiratory exacerbations in this population, and the S/D ratio a useful echocardiographic measure for pulmonary hypertension and both are significantly elevated in patients who did not survive. A rise in NT-pro BNP may be in part secondary to worsening respiratory status. Serial measurements may be useful to guide intensification of targeted therapy as an upward trend in NT-pro BNP to >8915 pg/mL carries a higher risk of death.

CONFLICT OF INTEREST

The authors have no conflicts of interest to disclose.

AUTHOR CONTRIBUTIONS

Study design: Amdani, Ross

Data collection: Amdani, Ross, Thomas, Mian


Data analysis: Amdani, Ross, Thomas, Mian

Initial draft of manuscript: Amdani, Ross, Thomas, Mian

Critical revisions: Amdani, Ross, Thomas, Mian

Approval of the final draft of the manuscript: Amdani, Ross, Thomas, Mian

ORCID

Shahnawaz M. Amdani MBBS, MD  <http://orcid.org/0000-0001-6074-3541>

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