Published in final edited form as:

Cardiol Young. 2017 July ; 27(5): 905–911. doi:10.1017/S1047951116001591.

Antihypertensive Drug Exposure in Premature Infants from 1997 to 2013

Srikanth Ravisankar, MD¹, Devon Kuehn, MD¹, Reese H. Clark, MD², Rachel G. Greenberg, MD³, P. Brian Smith, MD, MHS, MPH³, and Christoph P. Hornik, MD³

¹Department of Pediatrics, East Carolina University, Greenville, North Carolina, USA

²Pediatric Medical Group, Greenville, Sunrise, Florida, USA

³Duke Clinical Research Institute, Durham, North Carolina, USA

Abstract

Background—Systemic hypertension is increasingly recognized in premature infants. There is limited evidence regarding treatment, and most published treatment recommendations are based solely on expert opinions.

Methods—We identified all infants < 32 weeks and < 1500 g birth weight discharged from one of 348 neonatal intensive care units managed by the Pediatric Medical Group between 1997 and 2013. We defined antihypertensive drugs as vasodilators, angiotensin converting enzyme inhibitors, β receptor blockers, calcium channel blockers and central α_2 receptor agonists. We compared characteristics between infants who were treated with at least one antihypertensive drug during their initial hospitalization and infants who were not prescribed antihypertensive drugs using the Wilcoxon rank sum test or Pearson's χ -square test.

Results—We identified 2504/119,360 (2.1%) infants who required at least one antihypertensive drug. The median postnatal age of first exposure was 48 days (25th, 75th percentile 15, 86), and median length of therapy was 6 days (1, 16). Hydralazine was the most commonly prescribed antihypertensive with 1280/2504 (51.1%) treated infants exposed to the drug. More than 2 antihypertensive drugs were administered in 582/2504 (23.2%) infants, and 199/2097 (9.5%) of the treated infants were discharged home on antihypertensive therapy. Infants who received antihypertensive drugs were of lower gestational age ($p < 0.001$) and birth weight ($p < 0.001$) compared to infants not prescribed antihypertensive drugs.

Address correspondence to: Christoph P. Hornik MD, Assistant Professor of Pediatrics, Duke Clinical Research Institute, 2400 Pratt Street, Durham, NC, 27715; Phone: 919-668-8935; Fax: 919-668-7032; christoph.hornik@duke.edu.

Conflicts of Interest: None

Financial Disclosures: P.B.S. receives salary support for research from the National Institutes of Health (NIH) and the National Center for Advancing Translational Sciences of the NIH (1R21HD080606-01A1 and UL1TR001117), the National Institute of Child Health and Human Development (HHSN275201000003I and 1R01-HD081044-01), and the Food and Drug Administration (1R18-FD005292-01); he also receives research support from Cembra Pharmaceuticals (subaward to HHS0100201300009C) and industry for neonatal and pediatric drug development (www.dcri.duke.edu/research/coi.jsp). C.P.H. receives salary support for research from the National Center for Advancing Translational Sciences of the National Institutes of Health (UL1TR001117). R.G.G. receives salary support for research from the National Institutes of Health training grants (5T32HD043728-10 and 5T32HD043029-13).

Conclusions—Our study is the largest to describe current antihypertensive drug exposure in a cohort of exclusively premature infants <32 weeks gestational age. We found wide variation in practice for treatment of hypertension in premature infants.

Keywords

Pharmacotherapy; hypertension; neonate; prematurity; very low birth weight

INTRODUCTION

Systemic hypertension is increasingly being recognized in infants: current prevalence estimates range between 0.7 to 2%, and is higher in premature infants.^{1, 2} While the leading cause of hypertension in adults is recognized to be essential hypertension, studies have identified a number of etiologies for hypertension in infants.¹ Use of umbilical arterial catheters, certain drugs, and various renal, cardiac, pulmonary, and autonomic problems can interact to cause hypertension in infants.¹ Hypertension may resolve over time, but there are no observational or follow-up studies that describe the morbidities or mortalities associated with this diagnosis in the vulnerable premature neonatal population.³ This lack of data makes it difficult to determine whether either observation or treatment is the correct clinical course.

While drugs are commonly used to treat hypertension in older children and adults, use in infants is less common. This lower use may be due to lack of evidence-based guidance. Most published recommendations are based solely on expert opinions.³ There are no published studies or actively enrolling randomized trials to evaluate the safety and efficacy of antihypertensive drugs in infants. To date, only two studies have attempted to describe the use of antihypertensive drugs in the neonatal intensive care unit (NICU).^{4, 5} Both studies included both term and preterm infants. Further data are needed to describe current antihypertensive prescription practices, particularly in premature infants.

Given the unknown long-term risks of unmanaged hypertension and the limited data regarding drug management, this study aims to define the profile of antihypertensive drugs used and variations in their use in a large cohort of premature infants. Data from this study will support the design, choice of drug, and implementation of future trials.

MATERIALS AND METHODS

Study design and Setting

We used a database derived from the electronic health record (EHR) populated by clinicians of all infants cared for by the Pediatrix Medical Group in 348 neonatal intensive care units (NICU) in North America from 1997 to 2013. Data on multiple aspects of care were entered into a shared EHR to generate admission and daily progress notes and discharge summaries. Information regarding maternal history, demographics, drugs, laboratory results, diagnoses, and procedures were then transferred to the Pediatrix clinical data warehouse for quality improvement and research purposes.⁶

We identified all infants 32 weeks gestation and 1500 g birth weight discharged between 1997 and 2013. We excluded infants with major congenital anomalies. We collected antihypertensive drug exposure, demographic data, laboratory values, diagnoses, presence and duration of umbilical arterial lines, and postnatal systemic steroid exposure. This study was approved by the Duke University Institutional Review Board with a waiver of informed consent.

Definitions

We defined antihypertensive drug exposure as any exposure of any duration to an antihypertensive drug. Antihypertensive drugs were categorized by mechanisms of action and included the following: adrenergic receptor blockers (atenolol, esmolol, labetalol, propranolol), angiotensin converting enzyme (ACE) inhibitors (captopril, enalapril), calcium channel blockers (amlodipine, isradipine, nifedipine), and vasodilators (hydralazine, nitroprusside). Diuretics were excluded due to their frequent use for other indications (such as, prevention and treatment of bronchopulmonary dysplasia). We defined first-line antihypertensive therapy as the first antihypertensive drug received during hospitalization. Combination therapy was defined as use of two antihypertensive drugs on the same day. We defined discharge antihypertensive drug as any antihypertensive drug exposure on the day of discharge or the day prior to discharge. We defined hypertension as a clinical diagnosis of hypertension documented in the medical record by the treating neonatologist. Antenatal steroid exposure was defined as any maternal exposure to steroids prior to delivery, and postnatal steroid exposure was defined as any infant exposure to systemic dexamethasone or hydrocortisone after birth. Umbilical arterial catheter exposure was defined as at least one day with an umbilical arterial catheter in place. An infant was classified as having renal dysfunction if there was at least one serum creatinine level >1.7 mg/dL or renal failure if there was at least one serum creatinine >3.0 mg/dL at any time during hospitalization. To account for the contamination of infant measurements by maternal creatinine, we conducted a sensitivity analysis defining renal dysfunction and renal failure based serum creatinine values obtained at any time after day of life 14.

Statistical Analysis

Standard summary statistics were used to describe demographic characteristics; continuous variables were presented as median (25th and 75th percentiles), and categorical variables were presented as counts (percentages). We used a Wilcoxon rank sum test or Pearson's χ^2 -square test to compare distributions of continuous and categorical variables across groups.

We used univariable logistic regression to evaluate the association between infant factors and discharge on any antihypertensive drug. All covariates significantly associated with the outcome in univariable analysis were included in a full multivariable model. We used backward-stepwise selection of variables with a p value for removal of 0.2 and p value for entry of 0.15. Our final model included the following covariates: gestational age, diagnosis of renal dysfunction, birth by cesarean section, and prenatal steroid exposure. All statistical analyses were performed in STATA 13.1 (StataCorp LP, College Station, TX).

RESULTS

Antihypertensive Drug Exposure

A total of 119,360 infants from 348 neonatal NICUs were included, and 2504 (2.1%) received at least one dose of any antihypertensive drug during their hospitalization. Median postnatal and postmenstrual age at first exposure to any antihypertensive drug was 48 days (15, 86) and 34 weeks (29, 39), respectively. The total number of infant days of exposure to any antihypertensive drug was 30,679 days. Median duration of exposure among infants exposed to antihypertensive drug during their hospitalization was 6 days (1, 16).

Antihypertensive drug exposure decreased slightly over the duration of our study from 2.3% in 1997–2004 to 2.0% in 2005–2013, $p=0.009$ (Figure 1). Infants who received antihypertensive drugs were of lower gestational age and birth weight compared to infants not prescribed antihypertensive drugs, 26 weeks (25, 28) versus 28 weeks (26, 30), $p<0.001$ and 850 g (685, 1080) versus 1030 g (783, 1260), $p<0.001$, respectively (Table 1).

Diagnoses and Interventions

Hypertension was diagnosed in 3961/119,360 (3.3%) of infants, and was more common among infants exposed to antihypertensive drugs (Table 2). Of the infants diagnosed with hypertension, only 1853/3961 (46.8%) received antihypertensive drugs. Renal dysfunction and renal failure occurred in 7807/119,360 (6.5%) and 2326/119,360 (2.0%) of infants, and both were more common among those exposed to antihypertensive drugs (Table 2). When defined based on serum creatinine values obtained after the first 14 days of life only, renal dysfunction and renal failure were less common, occurring in 3569/119,360 (3.0%) and 1242/119,360 (1.0%), but both remained more common among infants exposed to antihypertensive drugs: 8.7% vs. 3.5%, $p<0.001$ for renal dysfunction and 3.0% vs. 1.2%, $p<0.001$ for renal failure. Of the infants diagnosed with renal dysfunction and renal failure, 308/7807 (4.0%) and 97/2326 (4.2%) received antihypertensive drugs. This distribution was similar when renal dysfunction or failure were defined based on creatinine values obtained after 14 days of life only: 188/3569 (5.3%) for renal dysfunction and 65/1242 (5.2%) for renal insufficiency.

Postnatal steroids and umbilical arterial catheters were used in 17,891/119,360 (15.0%) and 70,045/119,360 (58.7%) of infants. Use of postnatal steroids and umbilical arterial catheters were more common among infants exposed to antihypertensive drugs (Table 2). Umbilical arterial catheters also remained in place longer among infants exposed to antihypertensive drugs compared to those not exposed, 11 days (8, 17) versus 1 day (1, 1), $p<0.001$.

Drugs

Hydralazine was the most commonly prescribed antihypertensive drug (1280/2504, 51%), followed by the angiotensin converting enzyme inhibitors captopril (734/2504, 29%) and enalapril (457/2504, 18%) (Table 3). Propranolol was the most commonly used adrenergic receptor blocker (380/2504, 15%) while amlodipine was the most commonly used calcium channel blocker (193/2504, 8%). Postnatal age at the start of therapy and duration of exposure varied based on type of antihypertensive drug. Vasodilators and adrenergic receptor blockers were started at an earlier age compared to calcium channel blocker and angiotensin

converting enzyme inhibitors, and median duration of therapy with atenolol was longer than all other drugs, 19 days (2, 19).

Two or more antihypertensive drugs were administered to 582/2504 (23.2%) of infants, and 11,820/30,679 (38.5%) of infant days of exposure to any antihypertensive drugs. Infants were more likely to be prescribed at least two drugs if they were outborn (27% vs 21%, $p=0.003$) or had a diagnosis of hypertension (87% vs 68%, $p<0.0001$). Gestational age, birth weight, gender, race, postnatal steroid exposure, presence of an umbilical arterial catheter and renal dysfunction or failure were not associated with use of multiple antihypertensive drugs.

Among infants who received two or more antihypertensive drugs, the most common combinations were angiotension converting enzyme inhibitors and calcium channel blockers, 98/582 (16.8%), and adrenergic receptor blockers and vasodilators, 94/582 (16.2%). Median duration of exposure to 2 or more antihypertensive drugs was 1 day (1, 5).

Antihypertensive Drugs at Discharge

Survival among infants exposed to antihypertensive drugs was 92%. Only 199/2097 (9.5%) of infants ever exposed to any antihypertensive drug who survived were discharged on antihypertensive drugs. The most commonly used antihypertensive drugs at discharge among survivors were angiotensin converting enzyme inhibitors, 79/2097 (3.8%), and vasodilators, 62/2097 (3.0%). On multivariable analysis, prematurity was associated with an increased odds of being discharged on an antihypertensive drug, odds ratio=3.14 (95% confidence interval 1.82, 5.44) for gestational age <25 weeks versus gestational age 29–32 weeks, and odds ratio=2.90 (1.84, 4.58) for gestational age 26–28 weeks versus gestational age 29–32 weeks.

DISCUSSION

Current practice for treatment of hypertension in the premature infant is primarily based on expert opinion, with limited evidence-based data for drug use. Our study is the largest to describe current antihypertensive drug exposure in a cohort of hospitalized premature infants <32 weeks gestational age. We found that 2% of hospitalized premature infants were exposed to antihypertensive drugs, with hydralazine, captopril, and enalapril most commonly used. The majority of infants were exposed to only one type of antihypertensive drug during their NICU hospitalization. Duration of treatment was often <1 week and infants were rarely discharged on antihypertensive drugs.

The results of our study are consistent with previous studies that have shown an incidence of 1–3% for neonatal hypertension.^{1, 2, 7–9} The lack of a clear definition for neonatal hypertension makes clinical diagnosis challenging. The American Academy of Pediatrics does not recommend routine blood pressure screening in healthy term infants and there are no recommendations for screening of preterm infants.¹⁰ A systematic review of previous studies defined percentiles of systolic, diastolic and mean blood pressures in premature infants from 26 weeks onward and recommended pharmacotherapy when the infant's blood

pressure was consistently at the 99th percentile.^{311–15} Other expert opinions recommend correction of iatrogenic and reversible causes of hypertension (such as, fluid overload, pain, and use of drugs like steroids) prior to initiation of drug therapy.³ Data from a few published case series reveal a wide variety of agents used by physicians.^{1, 9, 16}

Our results are also consistent with previous observations that only approximately half of infants diagnosed with hypertension are treated with antihypertensive drugs.^{4, 5} However, a key difference between prior studies and ours is the patient population—we focused only on infants < 32 weeks. Though the etiology of hypertension is likely different in premature infants populations compared to term infants, this does not change the prevalence of antihypertensive drug therapy. This finding could be a result of the transient duration of hypertension, lack of clear guidelines in diagnosis and management, or concerns about drug adverse events in premature infants.

We identified several characteristics of premature infants exposed to antihypertensive drugs including lower gestational age at birth, lower birth weight, male gender, a diagnosis of hypertension, laboratory values consistent with renal dysfunction or failure, use of an umbilical arterial catheter, and administration of postnatal steroids. Several of these factors are known risk factors for hypertension.^{1, 7} In our study, it is difficult to know if physicians incorporated awareness of these potential risk factors in their decision of whether or not to treat with antihypertensive drugs.

Hydralazine was the most commonly prescribed antihypertensive drug. Hydralazine is an arterial vasodilator that causes direct smooth muscle relaxation and thereby decreases systemic vascular resistance and improves cardiac output. It is available in oral and intravenous forms. While the intravenous form has better bioavailability, the oral form can be compounded into a stable suspension for 7 days. Onset of action starts at 5–15 minutes after administration and duration lasts for 4–6 hours. Though there is no clear evidence for its use as the first-line agent, experience in using hydralazine in infants with cardiac diseases, its stable oral form and its rapid onset of action, may be the reasons for its preferred use.

Overall, infants were prescribed vasodilators or β blockers earlier than angiotensin converting enzyme inhibitors or calcium channel blockers. The later use of angiotensin converting enzyme inhibitors may be due to concerns about their impairment of the final stages of renal maturation in premature infants.^{3, 18, 19} Specifically, there are concerns about prolonged hypotension and oliguric renal failure with enalapril, even at low doses.^{3, 20, 21} The similar timing of vasodilators and β blockers may be due to the recommendation that β blockers should be used in combination with hydralazine when a single agent has been suboptimal.^{3, 22}

We also report combinations of antihypertensive drugs used in premature infants. In our population, <25% of premature infants receiving antihypertensive therapy receive more than 1 drug. Regardless of whether infants received single or combination therapy, duration of treatment was often <1 week, and infants were rarely discharged home on drugs. Long term follow-up studies of infants discharged home on antihypertensive drug therapy typically

show resolution of idiopathic forms of hypertension except for infants with underlying renal or cardiac disease, or a history of extracorporeal membrane oxygenation.^{7–9, 16, 23, 24} However, these studies are typically limited by small sample sizes. This limited evidence clearly makes it challenging to define appropriate guidelines for duration of therapy and drug choice. The selection of antihypertensive drugs is further complicated by the fact that data on long-term implications of neonatal hypertension and adverse events associated with prolonged pharmacotherapy are lacking.

The key strengths of our study include its large sample size, long study period, and representation of a diverse group of NICUs ranging from academic centers to community hospitals accounting for nearly 25% of all NICU admissions in the United States. Our study was primarily limited by the data source, which is derived from an electronic health record and has not undergone the scrutiny of clinical trial data collection. Antihypertensive drug exposure was extracted from provider documentation and orders, and confirmation of administration and infant tolerance of the drug by the bedside nurse was not available. This limitation extends to the description of combination antihypertensive drug therapy, which may be biased by infants transitioning from one drug class to another, rather than truly requiring two or more drugs to control their hypertension. Data on why individual drugs were chosen were not available, and we did not have access to blood pressure measurements to confirm indication and efficacy. To address this deficiency, we report on the number of infants diagnosed with hypertension, but acknowledge the limitation of such a diagnosis being made by treating providers without standardized diagnostic criteria. We are also unable to comment on the long-term use or impacts of antihypertensive drugs. This limitation may be of particular relevance given that a single center retrospective study of 36 hypertensive preterm infants found that postnatal age at hypertension diagnosis follows a bimodal distribution, with 13/36 infant diagnosed after discharge from the NICU.²⁶ Further, our definition of discharge medication is based on drugs recorded for an infant on the day of discharge or the day prior, and not based on pharmacy instructions recorded in the discharge summary or outpatient prescriptions, as neither were available from our data source. Lastly, our study conclusions apply to a select population of premature infants < 32 weeks gestational age and < 1500 g birth weight without major congenital anomalies. This population was chosen given that the highest prevalence of hypertension in premature infants was previously reported in the very low birth weight population,²⁶ and because of the potential confounding effect of major congenital anomalies on drug therapy. For example, infants with congenital heart disease may be exposed to adrenergic receptor blockers or angiotensin converting enzyme inhibitors for the treatment of arrhythmias or for afterload reduction after cardiac surgery, and not primarily to treat systemic hypertension.^{27,28}

In summary, we found that in practice, premature infants < 32 weeks gestational age are exposed to wide variations in drug use for hypertension. The findings of our study should inform the design and implementation of future randomized trials of antihypertensive drugs in premature infants.

References

1. Seliem WA, Falk MC, Shadbolt B, Kent AL. Antenatal and postnatal risk factors for neonatal hypertension and infant follow-up. *Pediatr Nephrol.* 2007; 22:2081–2087. [PubMed: 17874136]
2. Alagappan A, Malloy MH. Systemic hypertension in very low-birth weight infants with bronchopulmonary dysplasia: incidence and risk factors. 1998; 15:3–8.
3. Dionne JM, Abitbol CL, Flynn JT. Hypertension in infancy: diagnosis, management and outcome. *Pediatr Nephrol.* 2012; 27:17–32. [PubMed: 21258818]
4. Blowey DL, Duda PJ, Stokes P, Hall M. Incidence and treatment of hypertension in the neonatal intensive care unit. *J Am Soc Hypertens.* 2011; 5:478–483. [PubMed: 21925997]
5. Sahu R, Pannu H, Yu R, Shete S, Bricker JT, Gupta-Malhotra M. Systemic Hypertension Requiring Treatment in the Neonatal Intensive Care Unit. *J Pediatr.* 2013; 163:84–88. [PubMed: 23394775]
6. Hsieh EM, Hornik CP, Clark RH, Laughon MM, Benjamin DK, Smith PB. Best Pharmaceuticals for Children Act—Pediatric Trials Network. Medication use in the neonatal intensive care unit. *Am J Perinatol.* 2014; 31:811–822. [PubMed: 24347262]
7. Singh HP, Hurley RM, Myers TF. Neonatal hypertension. Incidence and risk factors. 1992; 5:51–55.
8. Buchi KF, Siegler RL. Hypertension in the first month of life. 1986; 4:525–528.
9. Skalina ME, Kliegman RM, Fanaroff AA. Epidemiology and management of severe symptomatic neonatal hypertension. 1986; 3:235–239.
10. Fetus CO, Newborn. Routine Evaluation of Blood Pressure, Hematocrit, and Glucose in Newborns. *Pediatrics.* 1993; 92:474–476. [PubMed: 8361812]
11. de Swiet M, Fayers P, Shinebourne EA. Systolic blood pressure in a population of infants in the first year of life: the Brompton study. 1980; 65:1028, 1035.
12. Pejovic B, Peco-Antic A, Marinkovic-Eric J. Blood pressure in non-critically ill preterm and full-term neonates. 2006; 22:249–257.
13. Kent AL, Meskell S, Falk MC, Shadbolt B. Normative blood pressure data in non-ventilated premature neonates from 28–36 weeks gestation. 2008; 24:141–146.
14. Lurbe E, Garcia-Vicent C, Torro I, et al. First-year blood pressure increase steepest in low birthweight newborns. 2006; 25:81–86.
15. Zubrow AB, Hulman S, Kushner H, Falkner B. Determinants of blood pressure in infants admitted to neonatal intensive care units: a prospective multicenter study. Philadelphia Neonatal Blood Pressure Study Group. 1995; 15:470–479.
16. Friedman AL, Hustead VA. Hypertension in babies following discharge from a neonatal intensive care unit. A 3-year follow-up. 1987; 1:30–34.
17. Molnar J, Nijland MJM, Howe DC, Nathanielsz PW. Evidence for microvascular dysfunction after prenatal dexamethasone at 0.7, 0.75, and 0.8 gestation in sheep. *Am J Physiol Regul Integr Comp Physiol.* 2002; 283:R561–R567. [PubMed: 12184988]
18. O'Dea RF, Mirkin BL, Alward CT, Sinaiko AR. Treatment of neonatal hypertension with captopril. 1988; 113:403–406.
19. Guron G, Friberg P. An intact renin-angiotensin system is a prerequisite for normal renal development. 2000; 18:123–137.
20. Wells TG, Bunchman TE, Kearns GL. Treatment of neonatal hypertension with enalaprilat. 1990; 117:664–667.
21. Mason T, Polak MJ, Pyles L, Mullett M, Swanke C. Treatment of neonatal renovascular hypertension with intravenous enalapril. 1992; 9:254–257.
22. Flynn JT. Neonatal hypertension: diagnosis and management. 2000; 14:332–341.
23. Sheftel DN, Hustead V, Friedman A. Hypertension screening in the follow-up of premature infants. 1983; 71:763, 766.
24. Arar MY, Hogg RJ, Arant BS, Seikaly MG. Etiology of sustained hypertension in children in the southwestern United States. 1994; 8:186–189.
25. Watkinson M. Hypertension in the newborn baby. 2002; 86:F78, F81.

26. Shah AB, Hashmi S, Sahulee R, Pannu H, Gupta-Malhotra M. Characteristics of systemic hypertension in preterm children. *J Clin Hypertens (Greenwich)*. 2015; 17:364–70. [PubMed: 25775924]
27. Chu PY, Hill KD, Clark RH, Smith PB, Hornik CP. Treatment of supraventricular tachycardia in infants: Analysis of a large multicenter database. *Early Hum Dev*. 2015; 91:345–50. [PubMed: 25933212]
28. Hsu DT, Zak V, Mahony L, Sleeper LA, Atz AM, Levine JC, Barker PC, Ravishankar C, McCrindle BW, Williams RV, Altmann K, Ghanayem NS, Margossian R, Chung WK, Border WL, Pearson GD, Stylianou MP, Mital S. Enalapril in infants with single ventricle: results of a multicenter randomized trial. *Circulation*. 2010; 122:333–40. [PubMed: 20625111]

Figure 1.
Percentage of infants exposed to antihypertensive drugs by discharge year

Table 1

Characteristics of the study population

	No antihypertensive drug N=116,771	Any antihypertensive drug N=2504	P
Gestational age (weeks)			
25	22.1%	34.9%	< 0.001
26–28	37.6%	45.9%	
29–32	40.2%	19.2%	
Birth weight (g)			
< 1000	46.7%	67.3%	< 0.001
1000–1499	53.3%	32.7%	
Small for gestational age	14.9%	17.5%	<0.001
Age at discharge (days) *	52 (33,76)	96 (71,122)	<0.001
Male	51.5%	56.5%	< 0.001
Race/ethnicity			
White	46.7%	49.3%	0.009
Black	27.8%	26.5%	
Hispanic	20.1%	20.0%	
Other	5.4%	4.1%	
Inborn	83.1%	77.8%	<0.001
Cesarean section	70.6%	73.9%	<0.001
5-minute APGAR score			<0.001
0–3	5.5%	6.6%	
4–6	18.1%	23.5%	
7–10	76.5%	69.9%	
Antenatal steroids	74.8%	75.0%	0.75

* median (25th, 75th percentile)

Table 2

Diagnoses and interventions associated with antihypertensive drug use

	No antihypertensive drug N=116,771	Any antihypertensive drug N=2504	P
Diagnosis of hypertension	1.8%	74.0%	< 0.001
Creatinine >1.7 mg/dL	6.4%	12.3%	< 0.001
Creatinine >3.0 mg/dL	1.9%	3.9%	< 0.001
UAC	58.4%	70.7%	< 0.001
Postnatal steroids	14.4%	42.0%	< 0.001

UAC = umbilical arterial catheter

Table 3

Frequency and duration of antihypertensive drug use

	Any antihypertensive drug, N=2504(%)	PNA at first exposure [*]	Duration of use (Days) [*]	Discharged home on drug if exposed during hospitalization ^{**}
Vasodilators	1365 (54.5)	25 (9, 74)	6 (3, 12)	62 (4.5)
Hydralazine	1280 (51.1)	26 (9, 75)	6 (3, 13)	62 (4.8)
Nitroprusside	16 (0.6)	8 (2, 16)	3 (1, 7)	0 (0)
ACE inhibitor	1191 (47.6)	74 (35, 98)	5 (1, 16)	79 (6.6)
Captopril	734 (29.3)	71 (32, 94)	6 (1, 18)	61 (8.3)
Enalapril	457 (18.3)	81 (42, 103)	2 (1, 9)	20 (4.4)
β receptor blocker	460 (18.4)	33 (13, 65)	2 (1, 13)	53 (11.5)
Propanolol	380 (15.2)	36 (16, 68)	2 (1, 13)	53 (13.9)
Labetalol	34 (1.3)	19 (8, 65)	3 (2, 5)	0 (0)
Esmolol	40 (1.6)	11 (3, 23)	3 (1, 10)	0 (0)
Atenolol	5 (0.2)	69 (39, 77)	19 (2, 19)	0 (0)
Metoprolol	1 (<0.1)	9 (9, 9)	1 (1, 1)	0 (0)
Calcium channel blocker	257 (9.9)	84 (67, 101)	1 (1, 10)	16 (6.2)
Amlodipine	193 (7.5)	84 (70, 100)	1 (1, 3)	14 (7.3)
Isradipine	38 (1.5)	95 (78, 131)	7 (1, 14)	0 (0)
Nifedipine	26 (1.0)	30 (6, 84)	7 (5, 15)	2 (7.7)

ACE=angiotension converting enzyme

^{*} median (25th, 75th percentile);^{**} denominator is number of infants ever exposed to each drug during their hospitalization