Very preterm birth is a risk factor for increased systolic blood pressure at a young adult age

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Abstract Children born very prematurely who show intrauterine growth retardation (IUGR) are suggested to be at risk of developing high blood pressure as adults. Renal function may already be impaired by young adult age. To study whether very preterm birth affects blood pressure in young adults, we measured 24-h ambulatory blood pressure (Spacelabs™ 90207 device) and renin concentration in 50 very premature individuals (<32 weeks of gestation), either small (SGA) or appropriate (AGA) for gestational age (21 SGA, 29 AGA), and 30 full-term controls who all were aged 20 years at time of measurement. The mean (standard deviation) daytime systolic blood pressure in SGA and AGA prematurely born individuals, respectively, was 122.7 (8.7) and 123.1 (8.5) mmHg. These values were, respectively, 3.6 mmHg [95% confidence interval (CI) −0.9 to 8.0] and 4.2 mmHg (95% CI 0.4–8.0) higher than in controls [119.6 (7.6)]. Daytime diastolic blood pressure and nighttime blood pressure did not differ between groups. We conclude that individuals born very preterm have higher daytime systolic blood pressure and higher risk of hypertension at a young adult age.

Keywords Hypertension · Intrauterine growth retardation · Premature birth · Developmental origins of health and disease · Ambulatory blood pressure monitor · Blood pressure

Abbreviations
IUGR Intrauterine growth restriction
SGA Small for gestational age
AGA Appropriate for gestational age
POPS Project of Preterms and Small for Gestational Age Infants
ABPM Ambulatory blood pressure monitor
SBP Systolic blood pressure
DBP Diastolic blood pressure
MAP Mean arterial pressure
SD Standard deviation
SDS Standard deviation score

Introduction

Over the past 20 years, evidence has accumulated indicating that intrauterine growth restriction (IUGR) affects normal development of the kidneys and vascular system, thereby increasing the likelihood of hypertension and/or cardiovascular diseases [1–4]. Impaired fetal kidney development leading to nephron deficit is considered an important pathway in the development of hypertension after IUGR birth [5–10]. Apart from IUGR, premature birth may contribute to the development of disease at adult age [11–15]. We demonstrated poorer renal function in very preterm individuals at age 19 years born with IUGR compared with very preterm individuals born with appropriate birth weight. Blood pressure did not differ between these groups [16, 17]. The findings from that study seemed to point to higher-than-normal prevalence of hypertension in both groups, but evidence was inconclusive for lack of controls.
In addition we performed a prospective follow-up study at age 20 years of individuals born very prematurely and in controls born after full gestation. Data obtained included inulin and para-amino hippuric acid clearance before and after renal stimulation, renal functional reserve capacity, microalbuminuria, renal size, blood pressure during renal function tests, 24-h ambulatory blood pressures and renin concentration. The glomerular filtration rate (GFR) did not differ between groups. The effective renal plasma flow was impaired, however, in preterm individuals with IUGR. These also showed increased microalbuminuria and higher systolic blood pressure (SBP) compared with controls. Renal size was related to GFR and effective renal plasma flow and was stunted, both regarding length and volume. In this article we report more detailed results of ambulatory blood pressure monitoring (ABPM) in these individuals.

Methods

Study population

We recruited three groups of 20-year-olds: (1) Born very prematurely (<32 weeks) and small for gestational age (SGA). (2) Born very prematurely and appropriate for gestational age (AGA). (3) Born full term (37–42 weeks) and appropriate for gestational age (controls). Individuals in the first two groups participate in a follow-up study of a Dutch birth cohort (all born in 1983) including 94% (N=1,338) of all live-born individuals in The Netherlands with gestational age <32 weeks and/or birth weight <1,500 g. This is the POPS cohort: Project on Preterms and Small for Gestational Age Infants. The so-called POPS 19 study included 596 (62.1%) of the 959 survivors until age 19 years. Four hundred and twenty-two of those were born with a gestational age <32 weeks and, 174 with a gestational age ≥32 weeks but with a birth weight <1,500 g. This is the POPS cohort: Project on Preterms and Small for Gestational Age Infants. The so-called POPS 19 study included 596 (62.1%) of the 959 survivors until age 19 years. Four hundred and twenty-two of those were born with a gestational age <32 weeks and, 174 with a gestational age ≥32 weeks but with a birth weight <1,500 g. Nonresponders of this study differed from responders by male gender, non-Dutch origin, low maternal education, severe handicap, and special education. SGA status did not differ between responders and nonresponders.

Participants in our study were recruited from the 422 individuals born with a gestational age <32 weeks who participated in the POPS 19 study (see also Fig. 1). Recruitment of SGA individuals started with the lowest birth weight adjusted for gestational age [≤−2.0 standard deviation scores (SDS)] of AGA individuals with the highest birth weight adjusted for gestational age (0 to +2.0 SDS, N=205). This strategy was aimed at gaining great contrast between groups. For the sake of convenience, those living closest to the Erasmus MC University Medical Center, Rotterdam, The Netherlands, were recruited first. Controls were term born (37–42 weeks of gestation) with normal birth weight (−2 to +2 SDS) between 1 January 1982 and 31 December 1984. They were all medical students at Erasmus MC and had been recruited by distribution of flyers in the medical school that mentioned as inclusion criteria normal gestational age, normal birth weight, and birth years 1982–1984.

Data collection

Patient characteristics Present weight, height, and body mass index (BMI) were measured and converted to SDS. Birth weights, gestational ages, obstetric, and perinatal parameters of the prematurely born individuals were derived from the POPS database (TNO Quality of Life, Leiden, The Netherlands). Birth weight was converted to SDS adjusted for gestational age using Swedish reference standards and was considered a measure of IUGR. Birth weights and gestational ages of controls were preferably obtained from birth records, midwives’ administration, birth announcement cards, or diaries. Otherwise, they were derived from mother’s recall. If reliable data on birth weight and gestational age could not be obtained, controls were not included in the study.

Blood pressure measurement

All participants underwent three blood pressure measurements at a minimum interval of 1 min in sitting position.
after at least 15 min of rest. Measurements were at the nondominant arm using a calibrated automatic blood pressure device (Hewlett Packard). The mean of these three measurements was used in the statistical analysis (baseline blood pressure). Thereafter, SBP and diastolic blood pressure (DBP) were measured with a Spacelabs™ 90027 ambulatory blood pressure monitor (ABPM) at the nondominant upper arm over 24 h. Cuff size was adjusted for upper arm length [24]. Reading frequency was programmed for every 20 min from 7 a.m. to 11 p.m. and for every hour between 11 p.m. and 7 a.m. Participants reported exercise, bedtime, and wake-up time in a diary. Mean daytime (awake), night-time (sleeping), and overall blood pressure values were calculated, and absolute and relative differences in day- and night-time blood pressure were derived. Hypertension criteria were those suggested by the European Society of Hypertension [25]. In 24-h measurements, normal blood pressure is defined as ≤130/80 mmHg and hypertension as >135/85 mmHg. Normal daytime blood pressure is defined as ≤135/85 mmHg and daytime hypertension as >140/90 mmHg. Normal nocturnal blood pressure is defined as <120/70 and nocturnal hypertension as >125/75. Blood pressure load is defined as the percentage of measurements exceeding the 95th percentile, adjusted for awakeness and sleep [26].

Blood was drawn at rest after baseline blood pressure measurement. Renin (ng Al/mg per hour) was quantitated by its capacity to generate angiotensin I from excess substrate angiotensinogen measured by an in-house radioimmunoassay, as described previously [27]. All participants also underwent a renal function test and renal ultrasonography by a method described elsewhere [18]. Data on plasma creatinine levels (μmol/liter), creatinine clearance (ml/min/1.73 m²), and total kidney length (cm) and volume (cm³) were also recorded.

Informed consent and institutional approval

Informed consent was obtained after oral and written information. All participants were able to provide consent with understanding and in writing. The Erasmus MC review board approved the study protocol.

Statistics

Statistical analysis was performed with SPSS 15.0 software. Multiple regression analysis served to analyze group differences adjusted for gender. Two group comparisons were performed to analyze the effect of IUGR (SGA vs AGA) and gestational age (AGA vs controls). Results are presented as mean, standard deviation (SD) and P value for the differences between groups. Multiple regression analysis was also performed to analyze relations between renin concentration and blood pressure adjusted for gender. Pearson’s chi-square tests were performed to analyze differences in perinatal factors between AGA and SGA groups and to compare prevalences of current hypertension between all groups. Statistical significance was considered at the level of 5%.

Results

Eighty-two individuals participated: 23 SGA, 29 AGA, and 30 controls. Thirty-five individuals registered with the research team as possible controls. Five were excluded because birth weight or gestational age were uncertain. For 12 of the 30 controls (40%), birth weight or gestational age were derived from maternal recall. Data of two of the SGA individuals were excluded from data analysis. In one, blood pressure measurements may have been unreliable because of frequent premature heartbeats. The second suffered an unrelated allergic episode prior to ABPM measurement. These two participants did not undergo ABPM. Table 1 shows baseline characteristics of the remaining 80 participants by study group. SGA individuals had significantly lower weight and height than AGA individuals and controls. BMI at age 20 did not differ between the three groups. Table 2 shows that proportions of mothers with hypertension during pregnancy, and cardiotocographic deterioration in the SGA group were significantly higher (Pearson χ² test: P<0.05) than those in the AGA groups. Infant respiratory distress syndrome also occurred more frequently in AGA individuals. Other prenatal, perinatal, and postnatal parameters were equally distributed (Table 2). There were no gender differences for birth weight (g), birth weight SDS and gestational age. At age 20, men were taller (mean height difference 13.4 cm 95% CI 10.0–16.9) and heavier (mean weight difference 12.3 kg 95% CI 7.7–17.0) than women, but BMI (mean difference 0.6 kg/m² 95% CI –0.7 to 1.9) did not differ significantly.

Baseline and ambulatory blood pressure monitor measurements

Men had significantly higher SBP and significantly lower heart rate than women (P<0.05). All regression analyses were adjusted for gender (Table 3).

Prematurity Baseline SBP in AGA individuals was 8.4 mmHg higher than in controls (95% CI 4.1;12.7, P<0.001). Ambulatory daytime SBP in AGA individuals was 4.2 mmHg higher (95% CI 0.4–8.0, P=0.03) than that in controls. Baseline DBP, ambulatory DBP, heart frequency (HF), mean arterial pressure (MAP), and nocturnal SBP dip
did not differ between these groups. Night-time SBP was slightly, though not significantly, higher in AGA individuals (AGA 3.6 mmHg higher with 95% CI −0.9 to 8.1, \(P=0.12\)). SBP load in AGA individuals was 5.5% higher than that in controls (\(P=0.049\)). DBP load did not differ between groups. Baseline MAP in AGA individuals was higher than that in controls.

Renal ultrasonography revealed subclinical renal anomalies in eight prematurely born individuals (three SGA and five AGA), i.e. nephrocalcinosis (\(N=1\)), unilateral pyelocaliceal dilatation (\(N=3\)), ureteropelvic junction obstruction (\(N=1\)), ureter dilatation (\(N=1\)), extrarenal pelvis (\(N=1\)), ureter duplication and ureterocele (\(N=1\)). In a separate analysis without these individuals, night-time SBP was also significantly higher in AGA individuals compared with controls (\(P=0.035\)), and SBP load was no longer significantly different (\(P=0.062\)). Baseline SBP, 24-h SBP, and daytime SBP all remained significantly higher in AGA individuals compared with controls (\(P<0.05\)).

**Table 2** Obstetric and perinatal parameters in the prematurely born groups

| Parameter                                | SGA (21) |   | AGA (29) |   |
|------------------------------------------|----------|---|----------|---|
| Hypertension during pregnancy*           | 13       | 62% | 2        | 7%
| Diabetes gravidarum                      | 1        | 5%  | 2        | 7%
| Blood group Antagonism                   | 0        | 0%  | 0        | 0%
| Pre-existing maternal disease            | 2        | 10% | 1        | 3%
| Premature rupture of membranes           | 2        | 10% | 8        | 28%
| Amnionitis                               | 0        | 0%  | 3        | 10%
| Cardiotocographic deterioration*         | 15 (3 unknown) | 71% | 4 (7 unknown) | 14%
| Use of corticosteroids                    | 2        | 10% | 8        | 28%
| Apgar score <7 after 5 minutes           | 1 (2 unknown) | 5%  | 1 (2 unknown) | 3%
| Infant respiratory distress syndrome*    | 5        | 24% | 16       | 55%
| Sepsis                                   | 8        | 38% | 13       | 45%
| Convulsions                              | 0        | 0%  | 0        | 0%
Hypertension prevalence

Table 4 shows equal distribution of normal blood pressure, borderline hypertension, and hypertension prevalences ($\chi^2=6.135; P=0.197$) between groups. No difference was found in daytime or night-time prevalences between groups.

Nocturnal dipping (decrease in blood pressure >10%) was observed in two of 21 SGA, four of 29 AGA, and four of 30 controls. In this regard, no significant difference was reached between groups ($\chi^2=0.233, P=0.556$). Renin concentration (ng Al/ml per hour) was available in 79 of the 80 individuals (Table 3). Renin concentration in SGA individuals was 0.42 lower than that in AGA individuals ($95\% \text{ CI } -0.08 \text{ to } 0.92, P=0.097$) and 0.48 lower than that in controls ($95\% \text{ CI } -0.09 \text{ to } 1.05, P=0.10$). No relation between renin concentration and SBP or DBP was found. In a linear regression analysis, renin concentration was not related to birth weight as a continuous variable (both SDS and g) or to gestational age.

Renal function and size

Renal size in controls was larger than that in AGA individuals. Renal size did not significantly differ between AGA and SGA individuals. Plasma creatinine and creatinine clearance did not differ between groups. Renal function and size were not correlated with SBP or DBP values.

Discussion

In this study, individuals born very prematurely (<32 weeks) showed higher SBP at age 20 years than did full-term individuals of the same age. Our findings are consistent,

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**Table 3** Mean 24-h, daytime and night-time blood pressure values, blood pressure load, renal function and size, and renin concentration in SGA and AGA preterms and controls

|                         | SGA | AGA | Controls | SGA vs AGA | AGA vs Controls |
|-------------------------|-----|-----|----------|------------|-----------------|
| **N**                   | 21  | 29  | 30       | 21         | 29              |
| **Mean SD**             |     |     |          |            |                 |
| **Baseline SBP (mmHg)** | 124.1 10.9 | 125.4 9.9 | 117.7 8.2 | 0.641 <0.001* | 0.336           |
| **Baseline DBP (mmHg)** | 70.0 9.8 | 70.0 7.8 | 68.0 7.1  | 0.997        |                 |
| **24-h blood pressure** |     |     |          |            |                 |
| SBP (mmHg)              | 118.0 7.9 | 119.4 8.6 | 115.9 7.9 | 0.534        | 0.042*          |
| DBP (mmHg)              | 68.4 7.3 | 68.5 4.9 | 67.6 6.4  | 0.994        | 0.600           |
| **Daytime blood pressure** |     |     |          |            |                 |
| SBP (mmHg)              | 122.7 8.7 | 123.1 8.5 | 119.6 7.6 | 0.816        | 0.031*          |
| DBP (mmHg)              | 73.0 8.0 | 72.2 5.1 | 71.7 6.0  | 0.652        | 0.819           |
| **Night-time blood pressure** |     |     |          |            |                 |
| SBP (mmHg)              | 107.9 8.3 | 110.9 9.1 | 107.9 9.4 | 0.254        | 0.118           |
| DBP (mmHg)              | 58.1 7.3 | 59.6 5.6 | 58.8 7.1  | 0.434        | 0.556           |
| SBP load (%)            | 17.9 20.0 | 18.1 20.2 | 12.3 16.5 | 0.817        | 0.049*          |
| DBP load (%)            | 10.1 14.8 | 7.7 7.5 | 13.8 17.3 | 0.795        | 0.251           |
| Nocturnal dip SBP (%)   | 12.0 5.3 | 9.9 4.4 | 9.8 9.5   | 0.204        | 0.928           |
| Nocturnal dip DBP (%)   | 20.3 7.3 | 17.3 6.4 | 18.1 6.5  | 0.161        | 0.497           |
| Creatinine clearance (ml/min/1.73 m²) | 106.5 22.1 | 119.4 31.0 | 109.6 31.9 | 0.116        | 0.255           |
| Creatinine (μmol/l)     | 71.6 9.1 | 70.7 12.3 | 72.0 13.2 | 0.811        | 0.881           |
| Total kidney length (cm) | 20.3 1.4 | 20.7 1.6 | 21.7 1.8  | 0.238        | 0.087           |
| Total kidney volume (cm³) | 252.5 80.2 | 263.2 49.3 | 310.3 79.7 | 0.585        | 0.021*          |
| Renin (ng Al/ml/h)      | 1.8 0.8 | 2.2 0.9 | 2.3 1.1   | 0.097        | 0.852           |

Data expressed by mean [standard deviation (SD)] and regression P value adjusted for gender

SGA small for gestational age prematurely born individuals, AGA appropriate for gestational age prematurely born individuals, controls appropriate for gestational age term-born individuals, SBP systolic blood pressure, DBP diastolic blood pressure

* P<0.05
however, with no effect of IUGR on SBP in these very preterm individuals. This would seem to suggest that prematurity alone is a predisposing risk factor for higher SBP. Interestingly, compared with SGA participants, more mothers of AGA participants had suffered from hypertension during pregnancy. Yet, we found no significant associations between prenatal circumstances, such as maternal hypertension and corticosteroid use, and adult blood pressure. Prevalence of hypertension was low, however, so that the analysis lacked power.

These results compare well with our earlier work, which documented a similar mean SBP of 123 mmHg and a high prevalence of hypertension (10%) in individuals born very prematurely recruited from the same cohort [16]. Likewise, Irving et al. reported higher blood pressure in 24-year-olds born prematurely (mean gestational age 33.4 weeks) and no effect of IUGR [28]. Preterm children in a Finnish study showed cardiovascular changes, including SBP, already at the age of 5 years [29]. A Swedish study of 165,136 men also showed an inverse association between gestational age, ranging from 35–44 weeks, and SBP [14]. Johansson et al. described that blood pressure in preterm children and SGA in term children at the age of 9.6 years did not differ from that in term children with appropriate birth weight. Catecholamine levels were higher in the preterm and SGA groups, however, suggesting increased sympathoadrenal activity [30].

In contrast, Singhal et al. found 15-year-olds born prematurely (mean GA 31 weeks; N=216) not to be at higher risk of vascular disease compared with term same-age peers [31]. Also, a large cohort study in Brazil could not detect different blood pressure values for 15-year-old individuals born with a gestational age less than or more than 37 weeks [32]. In that study, adolescent blood pressure was elevated only when there was IUGR. One possible explanation for these contrasting results is the obscure effect of early determinants of increased blood pressure during adolescence that Barker et al. described [3, 33].

Bayrakci et al. described abnormal circadian blood pressure regulation in SGA preterm children aged between 5 and 17 years [34]. Nocturnal dipping, especially in those born light for birth date, was less than that of controls. Their SBP was not different from that in controls. Blood pressure data were converted into SDS scores because subjects differed in age, height, and gender, as described by Wühl et al. [35]. For our adult population, such SDS scores were not available. Recently, Bacchetta et al. described increased DBP at the mean age of 7.6 years in very preterm individuals (mean gestational age 27.3 weeks) but no difference between growth-restricted individuals (either intrauterine or extrauterine) [36].

Siewert-Delle and Ljungman suggested that in preterm individuals, gestational age (30–38 weeks) has a great impact on adult blood pressure [15]. In our study, a possible effect of IUGR may therefore have been concealed. We found only a slightly stronger regression coefficient of gestational age to adult SBP for the preterm participants. Significance disappeared, however, in this smaller separate analysis. For that matter, an earlier study in a larger sample of individuals from our cohort who were born very prematurely neither revealed a relation between gestational nor adult SBP [16]. Therefore, our data suggest weaker relation between gestational age and adult SBP in the lowest ranges of gestation.

O’Brien et al. described different mean and 95th percentile 24-h ambulatory blood pressure values (p95)

| Table 4 | Prevalence of hypertension using the criteria for hypertension measured with an ambulatory blood pressure monitor (ABPM) defined by the European Society of Hypertension [42] |
|---------|---------------------------------|---------------------------------|---------------------------------|
|         | SGA                             | AGA                             | Controls                        |
|         | N     | %    | N     | %    | N     | %    |
| Normal 24-h blood pressure | 20    | 95   | 26    | 90   | 26    | 87   |
| Borderline 24-h hypertension | 0     | 0    | 1     | 3    | 4     | 3    |
| Mild, moderate, or severe 24-h hypertension | 1     | 5    | 2     | 7    | 27    | 93   |
| Normal daytime blood pressure | 20    | 95   | 27    | 93   | 27    | 90   |
| Borderline daytime hypertension | 0     | 0    | 0     | 0    | 3     | 10   |
| Mild, moderate, or severe daytime hypertension | 1     | 5    | 2     | 7    | 0     | 0    |
| Normal night-time blood pressure | 20    | 95   | 24    | 83   | 26    | 87   |
| Borderline night-time hypertension | 0     | 0    | 3     | 10   | 2     | 1.5  |
| Mild, moderate, or severe night-time hypertension | 1     | 5    | 2     | 7    | 2     | 1.5  |
| Total    | 21    | 100  | 29    | 100  | 30    | 100  |

Normal BP 24 hr ≤130/80 mmHg, Daytime BP ≤135/85 mmHg, Nighttime BP ≤120/70 mmHg. Borderline hypertension between normal blood pressure values and hypertension, Hypertension 24 h >135/85 mmHg, Daytime >140/90 mmHg, Nighttime >125/75 mmHg
for men and women as well as for different age categories [37]. For age range 17–29 years, the p95 were 136/81 for men (N=107) and 125/78 for women (N=174).

On the basis of these reference values, the prevalence of hypertension in our cohort was one in SGA, two in AGA, and two in controls (not significant). Prehypertensive blood pressure is a risk factor for hypertension in later life [38]. Screening at a young adult age could be effective to trace individuals at risk of adult hypertension and would enable early preventive treatment.

Strengths and limitations

A major strength of this study is that perinatal parameters for prematurely born individuals were obtained prospectively. Second, the study design provided for separate evaluation of the effects of premature birth and IUGR.

The main limitation lies in the relatively small group sizes, notably that of the SGA group. For this reason, reliable prevalence rates of hypertension (stages 1 and 2) and relative risks cannot be given. Much larger samples are needed to confirm our findings. Second, selection bias could have been introduced because recruitment was among individuals who participated in the POPS 19 study, which in itself had a response rate of 62.1%. The nonresponse group in the POPS 19 study differs significantly from the response group by male gender, non-Dutch origin, low maternal education, severe handicap, and special education [21]. However, there was no difference in SGA status. Third, we had to rely on the mother’s recall of her child’s birth weight and gestational age in 40% of controls. We assumed that a mother would have remembered premature birth or low birth weight. Most mothers knew the exact term date (40 weeks), from which we calculated gestational age. Birth weight could mostly be retrieved from birth announcement cards and diaries. As indicated earlier, uncertainty about these data was reason for exclusion. The chance of recall bias would therefore seem to be low. Moreover, parental recall of birth weight has been demonstrated to be a reliable variable in a statistical analysis [39, 40]. Lastly, controls were recruited from a medical school and may be considered to be better educated than the general population, especially on health issues. Thus, they may be in better health. Their blood pressure was comparable, however, to that of same-age individuals at risk of adult hypertension and would enable statistical analysis [39, 40]. Lastly, controls were recruited from a medical school and may be considered to be better educated than the general population, especially on health issues. Thus, they may be in better health. Their blood pressure was comparable, however, to that of same-age individuals at risk of adult hypertension and would enable statistical analysis [39, 40].

In conclusion, our results suggest that very premature birth increases SBP at a young adult age. IUGR does not seem to attenuate this effect. Considering our previous findings of decreased renal size, increased microalbuminuria, and decreased renal blood flow, very preterm individuals who suffered from IUGR are likely to have reduced nephron endowment. Systematic monitoring of blood pressure in young adults born very prematurely may identify individuals at risk for developing hypertension. Such a follow-up program may reveal the clinical significance of our findings.

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No conflict of Interest

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