Time to spontaneous ductus arteriosus closure in full-term neonates

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ABSTRACT

Objective: The mean closure time of the ductus arteriosus (DA) in full-term neonates is presumed to be 1–2 days after birth; however, whether this rate is accurate throughout the neonatal period is still unclear. In addition, the clinical determinants that influence DA closure remain unknown.

Methods: Echocardiography was performed 1826 times in 1442 participants (732 boys, 710 girls). An iE33 colour Doppler echocardiograph supplied by Philips Electronics was employed to examine DA flow. Data regarding sex, birth date, examination date, method of delivery, mother’s age, past deliveries, neonatal body weight and body height were also collected. The Statistical Analysis System makes statistical clarification of these queries possible. We examined the persistence of DA in full-term neonates and appropriate for gestational age (AGA) neonates in the early neonatal period using colour Doppler echocardiography, and a subsequent analysis with SAS.

Results: After performing multivariable analyses, the median DA persistency times were 27.42 and 45.10 h after birth in boys and girls, respectively. A statistically significant sex difference was observed (\( p<0.0001 \)). Additionally, significant time differences were observed between vaginal and scheduled caesarean deliveries, at 26.97 and 28.93 h, respectively (\( p<0.0245 \)). No significant differences were observed in the other variables.

Conclusions: Spontaneous DA closure time curves were clarified for the first time throughout the early neonatal period in full-term and AGA neonates. It was revealed that both sex and delivery method play important roles in time to DA closure.

One of the most important fetal circulation components, the ductus arteriosus (DA) is destined to close immediately after birth. It is known that the DA in full-term neonates usually closes 1–2 days after birth. Gentile et al. reported that the DA was undetectable in 42% of neonates by 24 h of age, and in 78% by 40 h. The actual persistency rate of the DA through the neonatal period is not clear since it is quite difficult to detect the specific DA closure time in individual neonates. In order to detect the accurate closure time, echocardiography must be continuously performed in the neonate for 1 or 2 days, which is nearly impossible in clinical practice. Another method is to gather many participants and examine each at fixed, predetermined time points. Although certainly reliable and accurate, these data are not continuous.

Survival analysis is used to analyse lifetime data, measured between a study’s start and either when an event of interest occurs or if the event does not occur within the study period, at the latest examination. The latter data are called right-censored data and are usually analysed by the Kaplan-Meier method. The DA closure data, however, are a mixture of right-censored, left-censored and interval-censored data and seldom contain accurate closure times. The latter two cannot be analysed using the Kaplan-Meier method. The main reason we cannot have an accurate
and continuous DA persistency rate is that there have been no statistical methods to deal with these three types of censored data. The Statistical Analysis System (SAS) LIFEREG (parametric maximum likelihood estimator for interval censoring data) and ICLIFETEST (non-parametric maximum likelihood estimator for interval censoring data) procedures allow statistical analysis of all censored data (right, left and interval) and can, therefore, analyse the DA data.

The SAS has programs which allow for detecting aspects influencing a targeted event. We chose some clinical factors that might influence the closure of DA. A multivariable analysis was performed to determine items that influence DA closure.

Colour Doppler images from conventional two-dimensional (2D) echocardiography have been widely used to detect stenotic, regurgitating, and shunt flows in children. Colour Doppler has also been used to evaluate DA conditions in neonates. In this study, we examined DA in full-term and appropriate for gestational age neonate participants using colour Doppler echocardiography, and analysed the data according to the SAS procedure to obtain a DA closure time curve.

The objectives of our study were: (1) to acquire the persistency rate curve as per the hours after birth and (2) to identify significant clinical determinants that influence the closure of the DA.

**MATERIALS AND METHODS**

**Study design**

In order to determine the closure time curve of DA in normal neonates, a statistically regulated number of neonatal participants were enrolled in this study to ensure sufficient power. Each of the participants underwent 2D colour Doppler echocardiography at an irregular age (hours) after birth (table 1D). Some participants were followed until DA closure.

Data of DA closure for each participant only have the DA was persistent or not at the examination and the time the examination was performed. We divided the DA closure data into three types. In the first type, called the ‘left’, the DA has already closed at the first examination, and we usually draw the time course from the left to the right. The time of the event is located to the left of the time of the first examination. In the second type, called the ‘interval-censored’, the closure of DA is presumed to be between the two examinations. In the third one, called the ‘right’, the DA does not close during the examination period. It is presumed that the closure time of DA is after the examination period, and the closure time is to the right of the end of the examination period.

The SAS program is the only one program that can deal with these three kinds of censored data. Both a non-parametric procedure (ICLIFETEST) and an accelerated-failure time model assuming log-logistic

| Table 1 Characteristics of participants |
|-----------------------------------------|
| (A) Clinical variables                    |
|                                        | Boys | Girls | Total | p Value |
| Number of participants                  | 732  | 710   | 1442  |
| Number of examinations                  | 897  | 929   | 1826  |
| Gestational age (week)                  | 38.7 (2.3) | 38.7 (2.3) | 0.34 |
| Range                                   | 37.0–41.9 | 37.0–41.7 |
| Birth weight (g)                        | 2990 (479) | 2890 (448) |
| Range                                   | 2280–3870 | 2225–3815 |
| Birth height (cm)                       | 48.3 (2.6) | 47.5 (2.6) |
| Range                                   | 42.8–53.2 | 43.1–52.7 |
| Parturient number                       | 343 (50) | 341 (50) | 684 |
| Multipara                               | 389 (51) | 369 (49) | 758 0.66 |
| Delivery style                          | 424 (50) | 419 (50) | 843  |
| Caesarean                               | 308 (51) | 291 (49) | 599 0.67 |

| (B) Examination profile                 |
|                                        | Boys | Girls | Total |
| Times of Examinations                  |     |      |       |
| 1                                      | 600 | 559   | 1159  |
| 2                                      | 103 | 104   | 207   |
| 3                                      | 25  | 33    | 58    |
| 4                                      | 4   | 9     | 13    |
| 5                                      | 0   | 3     | 3     |
| 6                                      | 0   | 2     | 2     |

| (C) Event occurrences and duration of observation |
|                                                |
| N                  | Minimum | Median | Maximum |
| With event         | 1204    |        |        |
| Boys               | 633*    | 7.67   | 62.60  | 167.62 |
| Girls              | 571*    | 16.78  | 70.01  | 161.40 |
| Without event      | 238     |        |        |
| Boys               | 99      | 2.12   | 28.85  | 148.57 |
| Girls              | 139     | 4.30   | 38.88  | 137.95 |

| (D) Distribution of age at examination     |
| Age (hour)                                 |
| Number of examinations                     |
| 0 to <8                                   | 41     |
| 8 to <16                                  | 60     |
| 16 to <24                                 | 128    |
| 24 to <32                                 | 244    |
| 32 to <40                                 | 133    |
| 40 to <48                                 | 132    |
| 48 to <56                                 | 232    |
| 56 to <64                                 | 121    |
| 64 to <72                                 | 118    |
| 72 to <80                                 | 192    |
| 80 to <88                                 | 91     |
| 88 to <96                                 | 61     |
| 96 to <104                                | 103    |
| 104 to <112                               | 61     |
| 112 to <120                               | 26     |
A single observer (HN) took all the measurements. The echocardiography examinations were performed 1826 times (897 in boys, 929 in girls) in 1442 participants. The distribution of the number of examinations for each participant is shown in table 1B. The observation durations are shown in table 1C. The distribution of age (hour) at examination is shown in table 1D. The event means to be detected are DA closure during the observation period. Although there were no significant differences between the sexes in the observation period, there was a significant difference in the number of events between sexes.

The participants’ clinical information was collected from their outpatient or inpatient records. Body height (measured in the prone position), and body weight were measured at the time of echocardiography.

**Statistical analyses**

The number of neonates needed for this study was determined according to Cohen’s report.³ The effect size (the difference of means divided by SD) was presumed to be 0.20. Statistical power was set at 0.85 and type I error in the two-sided test was allowed to be 0.05. Accordingly, at least 450 examinations were needed in each group in order to accurately compare the mean values.

The F-test and Student t test, respectively, were employed to determine whether the sample variance and means of the two categories were statistically equivalent. The \( \chi^2 \) test was employed to determine whether the proportions of the samples were statistically equivalent. The LIFEREG procedure in the SAS (V9.3; SAS Institute Inc, Cary, North Carolina, USA) software was employed to analyse the data on the accelerated-failure time model. The ICLIFETEST procedure was used to determine the presence of statistical differences, while Cox regression was used to compare clinical differences between the two categories. A multivariable analysis was performed in order to determine which clinical characteristics influenced DA closure. A p value of less than 0.05 was considered statistically significant.

**RESULTS**

The characteristics of the participants are shown in table 1. Between the sexes, there were no significant differences in group size, number of examinations, gestational age, parturient number or delivery methods. Additionally, there were no significant sex differences in the distribution of the number of examinations (table 1B). However, a significant sex difference (p<0.0001) was identified in the proportion of DA closures.

The persistency rate curve had a sigmoid-like curve shape (figures 1–3).
The multivariable analyses of DA closure using the ICLIFETIME and LIFEREG procedures are shown in table 2. Significant differences between sexes (p<0.0001) and delivery methods (p=0.0245) were noted, while there were no significant differences in the other clinical determinants.

**The sex difference**

The persistency rate curve in terms of sex was shown in figure 1. Figure 1A showed the non-parametric life survival time curve obtained with ICLIFETEST. The life survival time curves, hypothesising logarithmic logistic distribution and confirmed by the curves shown as

**Figure 1** Ductus arteriosus closure time curve by sex (p<0.0001). The life survival time curves were based on the hypothesised logarithmic logistic distribution (B using LIFEREG) and confirmed by the curves using ICLIFETEST (A). DA, ductus arteriosus.

**Figure 2** Ductus arteriosus closure time by delivery method (p=0.0245). The life survival time curves, which were based on the hypothesised logarithmic logistic distribution (B using LIFEREG), and confirmed by the curves using ICLIFETEST (A). DA, ductus arteriosus.
The persistency rates in girls were higher than that of boys throughout the examination period \((p<0.0001)\). The median DA survival times (MST) were 27.42 and 45.10 h after birth in boys and girls, respectively. The female:male ratio of MST was 1.645. The DA closure time in girls was significantly slower than that in boys.

**Comparison of DA closing time between vaginal and caesarean delivery**

The persistency rate curve in terms of delivery method is shown in figure 2. Figure 2A, B show the survival curves using ICLIFETEST and LIFEREG, respectively. Throughout the examination period, the persistency rates in neonates delivered via caesarean section were significantly higher than those delivered vaginally.

### Table 2  The multiple variance of coefficients in ductus arteriosus closing by ICLIFETEST (non-parametric) and LIFEREG (parametric)

| Number | MST ICLIFETEST | MST LIFEREG | p Value (ICLIFETEST) |
|---|---|---|---|
| **Sex** | | | |
| Female | 710 | 45.10 | 36.77 |
| Male | 732 | 27.42 | 26.89 | <0.0001 |
| **PN** | | | |
| Primipara | 684 | 27.40 | 30.97 |
| Multipara | 758 | 30.30 | 31.61 | 0.6254 |
| **Method** | | | |
| Vaginal | 843 | 26.97 | 30.25 |
| Caesarean | 599 | 28.93 | 32.75 | 0.0245 |
| **Height (cm)** | | | |
| ≥48 | 761 | 29.28 | 31.62 |
| <48 | 681 | 27.40 | 30.96 | 0.5612 |
| **Weight (g)** | | | |
| ≥2500 | 127 | 24.40 | 28.24 |
| <2500 | 1315 | 28.93 | 31.60 | 0.1620 |
| **GD (day)** | | | |
| ≥272 d | 760 | 28.93 | 31.31 |
| <273 | 682 | 26.47 | 31.31 | 0.3626 |

GD, gestational days; Height, body height; MST, median survival time; PN, parturient number; Weight, body weight.
higher compared to the neonates who were delivered vaginally \((p=0.0245)\). The MST was 30.25 and 32.75 h after birth among the participants delivered vaginally and via caesarean section, respectively. The vaginal: caesarean delivery MST ratio was 0.932. The DA closure time in the participants delivered vaginally was significantly faster than in those delivered via caesarean section.

The persistency curves of boys and girls delivered vaginally and abdominally are summarised in figure 3. Figure 3A, B show the survival curves using ICLIFETEST and LIFEREG, respectively. The differences in DA closure between delivery methods were more clearly defined among the girls than the boys.

**DISCUSSION**

A precise mean time of the DA in full-term neonates has not yet been established. Moreover, no data are available regarding the accurate persistency rate throughout the neonatal period.

There are no programs which are able to deal with three kinds of censored data; ‘left’, ‘interval’ and ‘right’ censored, except the SAS program. These programs are based on the statistical method of the maximum likelihood estimation.

We applied multivariable regression analysis using the SAS programs to the data on DA closure time. We discovered successive persistency rate curves of DA throughout the early neonatal period and identified factors which influence the closure time for the first time.

The DA survival curves had sigmoid shapes. These results seemed reliable, since sigmoid curves frequently appear in many biological field situations, such as the increase in the population, as well as the relationship between death rate distribution and quantity of toxic agents, haemoglobin oxygen saturation and oxygen tension, and so on.

**Factors that influenced DA closure**

There were significant differences in sex and delivery method (table 2), and the \(p\) value for sex \((p<0.0001)\) was lower than that for the delivery method \((p=0.0245)\).

**Sex**

It has been reported that female patients have failures of DA closures about twice as often as male patients.\(^4\)\(^5\)

Flinsenberg et al.\(^6\) reported that oestrogen 17 \(\beta\)-estradiol induced an endothelium-independent relaxation of chicken DA. It is presumed that oestrogen may play an important role in the persistence of DA. No study has been reported on the theme in humans and therefore, the true reason is not known.

We reported a significant sex difference in the left ventricular end-diastolic volume index (body surface area calculated using the index formula reported by Haycock et al.\(^7\)) in the early neonatal period using three-dimensional echocardiography.\(^8\) These results revealed that the sex differences seemed to already appear both morphologically and functionally.

**Delivery method**

Differences have been reported in the neonatal laboratory data and clinical conditions between the vaginal and caesarean delivery. However, there have been no reports on the relationship between the DA closure time and delivery method. This is the first report that examines these relationships.

Smith reported that prostaglandin E2 (PGE2) induces relaxation of the DA through the Gs protein adenyl cyclase. Increased cyclic adenosine monophosphate (cAMP) and PGE2 is one of main mechanisms of patency of DA.\(^9\) Interleukin-6 (IL-6) stimulates synthesis of PGE2. Mojaveri et al.\(^10\) reported that mean serum IL-6 levels in neonates delivered vaginally (10.9 pg/mL) were higher than those in neonates delivered by elective caesarean section (6.6 pg/mL), but the difference was not significant \((p=0.86)\). The relationship between serum IL-6 level and DA closure is still unclear.

Using full-spectrum near-infrared spectroscopy, Isobe et al.\(^11\) reported that the cerebral oxygenation in neonates delivered abdominally significantly decreased compared to that in neonates delivered vaginally at 8.5 min post-delivery. Urelesberger et al.\(^12\) reported that arterial oxygen saturation and heart rate were significantly lower in infants delivered abdominally than in those delivered vaginally. Noh et al.\(^13\) examined oxidative markers in the umbilical cord blood and reported that the markers in neonates who were delivered through planned caesarean section were significantly higher compared to those in infants delivered vaginally. These reports indicate that neonates delivered abdominally are at greater risk for problems related to their respiratory or circulatory systems.

**Other covariates**

No significant relationships were seen between DA closure and the parturient number, body weight, body height or gestational days (table 2).

**Study limitations**

The DA closure time curves are based on all participants’ data and not on individual participant data. It is impossible to expect an exact DA closure time for each neonate.

**Application of the SAS programs to other fields**

The life data analysis used in this paper could potentially be applied to many diseases. Lately, interest has grown in progression-free survival for slowly progressive diseases that are difficult to cure (Y So, G Johnston, SH Kim. Analyzing interval-censored survival data with STSTM software. SAS global forum 2010).\(^14\)\(^15\) Progression-free survival is often used as a clinical end point in randomised controlled trials for some slowly progressive diseases.
cancer therapies; however, there has been no way to study the morbidity rate and age of onset in patients with these diseases. The procedure used in this paper could be usefully applied to the analysis of these diseases.

Another advantage to the SAS program is the ability to freely choose examination intervals. Examining many patients at particular fixed intervals or ages can be a burden to examiners. It is preferable that examination intervals be conducted irregularly and still produce an accurate survival curve.

CONCLUSION
Using DA closure data acquired by echocardiography and analysed through SAS programs, successive persistency rate curves of DA throughout the early neonatal period were revealed. Factors that influence the closure time were identified for the first time. These results could lead to new progressions in the endocrinological, neurological or other fields of the early neonatal period.

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Collaborators HN examined all the participants and drafted the work. YK supervised the writing of the paper and gave the final approval. CH, MW, YN and SN analysed the data using SAS programs.

Competing interests None declared.

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