

1 **Evaluation of Tissue Perfusion Status in Moderate to Late Preterm**

2 Running Title: Tissue Perfusion Status in Preterm Infants

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23 **Summary**

24 The aim of this study was to investigate the tissue perfusion status and circadian rhythm
25 in moderately premature infants. As a prospective study, from July 2019 to October
26 2019, the haemodynamic stability of moderate to late preterm, including such indicators
27 as perfusion index (PI), blood pressure (systolic/diastolic) (BP), heart rate (HR),
28 respiratory rate (RR), oxygen saturation (SpO₂) and body temperature were monitored
29 in the morning and at night within eight days after birth. There was no difference of
30 statistical significance between PI values in the morning and at night ($P > 0.05$). The
31 HR from days six to eight after birth was higher than days one to three ($P < 0.05$). The
32 HR increased significantly on days seven and eight compared with days four and five
33 ($P < 0.05$). The BP from days three to eight was significantly higher than on day one (P
34 < 0.05), and the BP from days four to eight was higher than on day two. There was a
35 weak positive correlation between the PI values and gestational age (GA) ($r = 0.097$),
36 HR($r = 0.067$) and time ($r = 0.284$), and a negative correlation with SpO₂ ($r = -0.113$).
37 The PI and HR of moderate to late preterm increased within eight days after birth. BP
38 was relatively lower after birth and gradually increased to a stable level on days three
39 to four. The PI and BP circadian rhythms associated with tissue perfusion were not
40 established on day eight after birth.

41 **Keywords:** Infant, Premature infant, Perfusion index, Pulse rate, Blood pressure

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45 **Introduction**

46 Premature infants refer to infants born at less than 37 weeks of gestation. [1]
47 Premature infants are deficient in body function due to inadequate maternal incubation
48 time. At the same time, the change of living environment (from inside to outside the
49 uterus) changes the hemodynamic state of premature infants. Hemodynamic instability
50 may lead to perfusion changes, leading to various perinatal complications and even
51 death.[2-5] Therefore, monitoring the hemodynamic status of preterm infants can
52 master the perfusion level of preterm infants and provide a theoretical basis for clinical
53 treatment.

54 In recent years, the wide application of pulse oximeter and noninvasive
55 sphygmomanometer in neonatal intensive care unit has promoted the preliminary
56 evaluation of neonatal blood circulation stability to a certain extent. [6] Blood pressure
57 may not be a good indicator of peripheral perfusion due to the influence of receptor
58 fluid and sympathetic nervous system.[7] Perfusion index (PI) is the ratio of pulsatile
59 blood flow to non pulsatile blood flow in the monitored tissue. It has been proved to
60 be a simple and noninvasive method to reflect the changes of peripheral perfusion.
61 [8,9] Studies have shown that perfusion index plays an important role in the evaluation
62 of disease severity, the screening of congenital heart disease, the early identification
63 of neonatal shock, and so on.[10] In addition, circadian rhythm is the basic feature of
64 life phenomena, and its changes can lead to pathological changes in human tissues.
65 [11] Some studies suggested that the introduction of a robust light dark cycle in the
66 neonatal intensive care unit can be used to guide the circadian rhythm system of

67 preterm infants, which may be conducive to the growth and development of preterm
68 infants. [12] However, there is no research on the correlation between PI and circadian
69 rhythm in preterm infants.

70 Therefore, in this paper, correlations between the PI value and blood pressure
71 (BP) (systolic/diastolic), heart rate (HR), respiratory rate (RR), oxygen saturation
72 (SpO₂) and circadian rhythm were evaluated by continuously monitoring the data of
73 preterm infants under stable conditions within eight days after birth.

74

75 **Methods**

76 The prospective study was conducted consecutively at the Neonatology
77 Department of Children's Hospital of Shanxi from July 2019 to October 2019. The
78 criteria for the subjects included: (1) moderate to late preterm at 32 to 36 weeks
79 gestational age (GA) [13]; (2) admitted within six hours after birth; (3) an Apgar score
80 of eight at 1 min and 10 at 5 min; (4) haemodynamically stable preterm newborns,
81 characterised by smooth breathing, normal colour and cry, normal position and activity,
82 normal muscle strength and muscle tone; (5) no need for oxygen or respiratory support.
83 Exclusion criteria included: newborns with infections, congenital heart diseases
84 (CHD), NEC, continuous apnoea episodes, hyperpyrexia ($\geq 37.5^{\circ}\text{C}$) and
85 pneumothorax during hospitalisation. This study was conducted in accordance with
86 the Declaration of Helsinki and approved by the Ethics Committee of Children's
87 Hospital of Shanxi.

88 All enrolled preterm infants were assessed by physical examinations and

89 laboratory tests to clarify their physical conditions (smooth breathing, normal colour
90 and cry, normal position and activity, normal muscle strength and muscle tone) after
91 admission, and every morning their conditions were checked. The pulse oximeter was
92 placed on the foot of preterm newborns in a calm state for half an hour after feeding
93 between 8:00–10:00 in the morning and 22:00–0:00 at night. The following data were
94 then recorded: PI, HR, SpO₂, BP, RR and body temperature (T). The PI and HR were
95 detected by a Masimo Radical-7 (USA) monitor, and SpO₂ was read three consecutive
96 times in six seconds to obtain the average value.

97 Statistical Analyses

98 The Kolmogorov–Smirnov test was used to quantify the normal distribution data,
99 which were expressed as mean \pm standard deviation ($m \pm SD$). Two groups of
100 parametric variables were compared by a t-test, and variance analysis was adopted for
101 multi-group comparison. $P < 0.05$ was considered statistically significant. The Pearson
102 correlation coefficient was used to analyse the correlation of variables. Multiple linear
103 regression analysis was applied to establish multiple linear regression equations. All
104 statistical calculations were processed by *SPSS 22.0* in Windows (IBM *SPSS* Statistic).

105

106 **Results**

107 A total of 95 preterm infants (32 to 36 weeks GA) were admitted to the
108 Neonatology Department of Children’s Hospital of Shanxi. One patient suffered from
109 NEC on day eight after birth, one patient was suspected of sepsis due to persistent
110 fever during hospitalisation, and six patients were excluded due to incomplete

111 information. Eighty-seven preterm infants, all infants were inborn, were involved in
112 the study. There were 2 cases of placenta previa, 2 cases of mild anemia during
113 pregnancy, 2 cases of mild pregnancy induced hypertension, 1 case of breech position,
114 2 cases of premature rupture of membranes > 6h, and 3 cases of test tube infants.
115 Intrauterine growth restriction occurred in 3 cases. There were 16 cases of natural
116 delivery and the rest were cesarean section. The average umbilical cord ligation time
117 was (1.4 ± 0.3) min. There were 48 M and 43F, with a mean GA of 34.4 ± 1.1 weeks
118 (W), BW (birth weight) of $2,142.6 \pm 384.8$ g and a mean Apgar score of 9.8 ± 0.4 at 1
119 min and 9.9 ± 0.3 at 5 min after admission.

120 The mean PI, HR, BP, SpO₂ and T values of preterm infants eight days after birth
121 are shown in Table 1. The PI values of the preterm infants were in a growing trend
122 after birth, which increased significantly from days five to eight compared to days one
123 and two ($P < 0.05$), and the PI values on days seven and eight were much higher than
124 those from days one to four ($P < 0.05$). However, there were no significant differences
125 among the PI values on the other days ($P > 0.05$). The HR increased gradually after
126 birth, with the values from days six to eight after birth much higher than those from
127 days one to three ($P < 0.05$). The HR increased significantly on days seven and eight
128 compared with those on days four and five ($P < 0.05$), and the HR increased
129 significantly on day eight compared with day six ($P < 0.05$). There were no significant
130 differences between the HR values from days one to five ($P > 0.05$). BP from days
131 three to eight was significantly higher than on day one ($P < 0.05$), days four to eight
132 were higher than on day two, while the value stayed stable on the other days ($P > 0.05$)

133 (Figure 1). The T values on days two and three after birth were higher than on day one.
134 On day six, the value was lower than on day two ($P < 0.05$), and for the remaining
135 days, the values were similar ($P > 0.05$). There was no significant difference between
136 the RR and SpO₂ during the period.

137 BP (systolic/diastolic) in the morning and night were similar ($t = 1.691, P = 0.194$;
138 $t = 0.370, P = 0.543$) from day one to day eight. The PI in the morning and night from
139 days one to eight were similar ($P > 0.05$) (Table 2). There was a difference of statistical
140 significance for PI values among different GAs ($F = 6.233, P < 0.001$), and the PI
141 value increased along with GA (Table 3).

142

143 The results of the Pearson correlation analysis showed a weak positive correlation
144 between the PI value with GA ($r = 0.097$), HR ($r = 0.067$) and time ($r = 0.284$) and a
145 negative correlation with SpO₂ ($r = -0.113, P < 0.01$) (Table 4). Multiple linear
146 regression analysis was adopted with the PI as the dependent variable, while GA, HR,
147 time and SpO₂ were independent variables. The calculation of the PI value under the
148 non-standardised regression model was based on the following equation: $PI = 2.253 +$
149 $0.057 \times GA (w) + 0.062 \times \text{time (day)} - 0.03 \times \text{SpO}_2 (\%)$ (Table 5).

150

151 Discussion

152 As an optical plethysmography parameter related to systemic perfusion, PI can
153 serve as a sensitive reflection of the perfusion level of peripheral tissues, with a
154 correlation to ventricular output.[8,14] Sivaprasath et al. proposed that the PI value was

155 positively correlated to pulse pressure and systolic and diastolic BPs to various degrees
156 among children aged 1~12 years. The decline of the PI value may predict impending
157 shock but was not reliable for detecting hypotension [15] It was found that the PI and
158 HR of moderate to late preterm in the first eight days after birth were growing slowly
159 at the points of time in our study, similar to findings from previous researches at home
160 and abroad.[16-18] A positive correlation between the PI and HR was also found, but
161 there was no significant correlation between the PI and BP.
162 A physiological theory holds that BP is decided by cardiac output (heart stroke volume
163 and HR) and peripheral vascular resistance (arterial compliance, ratio of systemic blood
164 flow to systemic vascular volume) while neonatal BP is affected by multiple factors,
165 including GA, age in days, BW, postnatal age, antenatal hormones, patent ductus
166 arteriosus and temperature. So far, there is no unified definition of hypotension,
167 weakening its credibility as an indicator for evaluation. Therefore, blood flow may be
168 a better indicator of perfusion than BP. Neonatal myocardial contractile elements were
169 significantly fewer compared with older children and adults. The immature myocardial
170 cells tended to exhibit a higher basal contractile state and were more sensitive to cardiac
171 afterload, [19] hence the mobilisation of the cardiac reserve may first be characterised
172 by an increase in HR rather than BP despite the instability of systemic blood perfusion.
173 All infants, especially premature infants, experience a series of haemodynamic changes
174 during the transitional period after birth, including intrauterine to extrauterine changes,
175 decreased pulmonary arterial pressure, shunting of blood flow from the systemic
176 circulation to the pulmonary circulation, closure of the ductus arteriosus and increased

177 volume of the systemic circulation. A HR value between 120 and 160 bpm coupled with
178 weak myocardial contraction means that the cardiac reserve could be achieved by
179 increasing HR to maintain tissue perfusion. Even in the absence of adequate tissue
180 perfusion during the compensatory period of shock, peripheral blood vessels are
181 responsive to ischemic stimuli via the sympathetic nervous system and humoral
182 regulation. This is also one of the reasons that many newborns have tachycardia with
183 or without increased BP and no hypotension during the compensatory period of shock.
184 However, the inflammatory reaction during shock can seriously affect the
185 microcirculation of adjacent tissues and skin, as indicated by the significant decrease
186 of the PI value of peripheral blood flow in the first 45 seconds after ischemic
187 stimulus.[20] Our study showed that the BP of moderate to late preterm was lower after
188 birth and tended to stabilise on days three to four after birth. The PI and HR were
189 recorded until day 10 after delivery in the preliminary study, which was found to be
190 increasing, while there was not much change in BP. However, part of the data was
191 eliminated because the patients were discharged. Therefore, arterial BP is not an
192 accurate indicator to evaluate neonatal peripheral tissue perfusion, while PI, which
193 reflects the ratio of arterial blood flow against non-arterial blood flow, is considered
194 more reliable in this regard. Theoretically, the cardiac reserve capacity would increase
195 with age. As the HR of infants and young children is lower than newborn infants, the
196 HR value of preterm infants should gradually decrease and stabilise at a particular stage.
197 However, at least on day eight after birth, we have not seen a drop in RP. Whether there
198 is a similar trend of HR for term infants will be the focus of future studies.

199 Previous studies focusing on the PI values at different ages claimed the median PI
200 of preterm infants with a GA <32W was 0.9 on the first day after birth, 1.8 at 24 hours
201 after birth,[21,22] and 3.0 for children 1~3 years old.[14] Our research showed that the
202 PI values for preterm infants with a GA 34~36W were significantly higher than those
203 with a GA 32~33W, which would increase with age, suggesting that PI was related to
204 the maturity of preterm infants. Meanwhile, the correlation analysis in our study
205 showed that PI is related to GA (W), time (day) and SpO₂ (%), which were incorporated
206 into the equation for the calculation of PI. In this equation, SpO₂ is easy to be measured
207 as both GA and time are objective indicators. The results of the equation were similar
208 to those of previous studies, making it a useful tool to predict the normal value of PI in
209 moderate to late preterm within eight days after birth. It may also be applied to term
210 infants and earlier preterm infants. Previous studies reported that a low PI value below
211 0.7 indicated left heart obstructive disease. [23] In our study, it was found that the PI
212 value of a preterm infant suffering from severe NEC on day eight eventually led to
213 surgery, before which the PI value dropped from 1.5 to a much lower level of 0.76,
214 suggesting that PI could be a signal of low systemic perfusion level. The significant
215 decline of the PI value during neonatal shock should be taken as a reference based on
216 the normal value of individuals for liquid recovery treatment. The PI equation obtained
217 in this study was helpful to evaluate the node of fluid resuscitation. We tried to
218 resuscitate a 2-day-old preterm infant with a GA of 34W with a PI value of 0.28 from
219 severe shock. When the PI value gradually increased to 1.9, saline dilatation was
220 terminated. Unfortunately, the infant developed a pulmonary haemorrhage. Therefore,

221 it is necessary to explore the node of PI value for different infants during fluid
222 resuscitation to improve the prognosis of infants in shock.

223 PI is the pulse index of blood flow, which is influenced by multiple factors, such
224 as muscle contraction, temperature, blood shunting, invasive procedures, neonatal
225 posture, circadian rhythm, etc. The data was measured when the infants were in a calm
226 state with minimum invasive procedures to ensure the accuracy of data. Due to the
227 circadian rhythm of the sympathetic-parasympathetic nervous system and various
228 activities, BP dipping of adults and older children within 24 hours is usually higher than
229 10%,[24,25] a phenomenon gradually formed as a result of the development and
230 maturity of children and the effect of various factors. This paper found no difference in
231 BP and PI values between morning and night within eight days after birth, probably due
232 to the instability of the sympathetic-parasympathetic nervous system in the early
233 postnatal period of preterm infants, which prevented the establishment of the circadian
234 rhythms associated with BP and PI at the end of day eight after birth. Yet in the primary
235 stage of the study, namely on day 10 after delivery, no difference was found in BP values
236 between morning and night while there were differences for PI. However, the data were
237 eliminated, given that the absence of some data might lead to a deviation. Therefore, it
238 remains unknown whether the circadian rhythm related to PI of moderate to late preterm
239 could be established 10 days after birth.

240 The present study had several limitations. First, the sample size of this study was
241 very small. Thus, our results should be confirmed in a further study with more infants.

242 Second, This study was only monitored for 8 days, and only two monitoring periods of

243 PI per day were set in this study. Finally, the hormone levels of infants were not
244 measured in this study. So the relationship between PI changes and hormone levels was
245 not detected.

246

247 **Conclusion**

248 This study established an equation for calculating PI value by recording and
249 analyzing GA, PI, HR, BP, SpO₂ and day values of preterm infants within 8 days after
250 birth. The equation can be used to preliminarily evaluate the peripheral perfusion of
251 middle and late preterm infants within 8 days after birth.

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254 Neonatology , Children's Hospital of Shanxi and Women Health Center of Shanxi,
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257 **Authors' contributions**

258 Liu K, Zhao Y and Yang G had full access to all of the data in the study and take
259 responsibility for the integrity of the data and the accuracy of the data analysis. Liu K
260 and Zhao Y designed the study and draft the manuscript, Yang G assisted with study
261 design, collected data and conducted data analysis. Niu S, Zhang M and Gao F
262 coordinated the data collection. All authors read and approved the final manuscript.

263 **Disclosure Statement**

264 The authors hereby declare no competing conflict of interest regarding any aspect of
265 this article.

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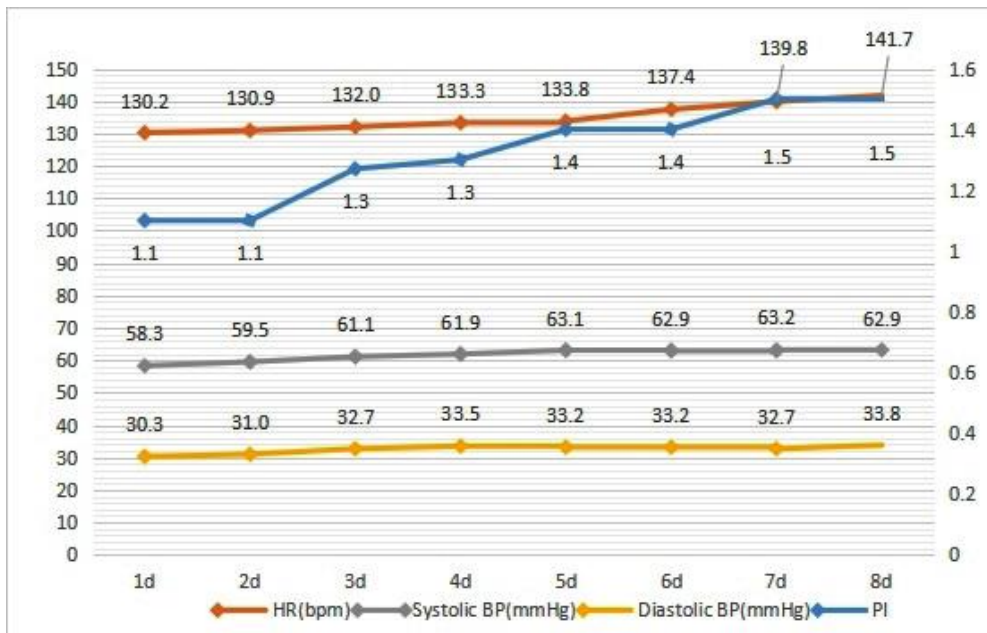
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376 Figure 1 Mean of PI, HR, BP values of preterm infants during 8 days after birth

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Table 1 Mean PI, HR, BP, SpO₂, T values of preterm infants during 8 days after birth

Time (day)	PI	HR	Systolic BP	Diastolic BP	SpO ₂ (%)	RR (bpm)	T (°C)
		(bpm)	(mmHg)	(mmHg)			
1	1.1±0.5	130.2±10.7	58.3±7.9	30.3±6.1	97.6±2.1	44.8±8.2	36.8±0.4
2	1.1±0.5	130.9±10.1	59.5±6.2	31.0±5.5	97.3±2.4	43.4±7.0	37.0±0.2
3	1.3±0.5	132.0±12.1	61.1±7.2	32.7±5.9	97.5±2.2	42.9±7.3	37.0±0.3
4	1.3±0.5	133.4±14.6	61.9±8.6	33.5±6.6	97.1±2.2	43.2±6.7	36.9±0.3
5	1.4±0.6	133.8±15.2	63.1±7.6	33.2±4.7	97.5±2.1	44.6±7.2	37.0±0.3
6	1.4±0.5	137.4±12.3	62.9±6.0	33.2±5.6	97.2±2.7	44.0±6.6	36.9±0.8
7	1.5±0.6	139.8±12.4	63.2±6.0	32.7±4.5	97.7±1.9	44.5±7.3	37.0±0.3
8	1.5±0.5	141.7±12.2	62.9±7.1	33.8±6.7	97.5±2.3	43.5±7.4	36.9±0.3
<i>F</i>	12.872	18.589	10.411	7.185	1.262	1.609	3.442
<i>P</i>	<0.001	<0.001	<0.001	<0.001	0.266	0.129	0.001

391 Note: Tukey test for post-test

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Table 2 PI values in the Morning and Night

Time (day)	morning	night	<i>t</i>	<i>P</i>
1	1.1±0.5	1.1±0.4	0.417	0.678
2	1.2±0.5	1.1±0.4	0.553	0.582
3	1.3±0.6	1.3±0.5	-0.188	0.851
4	1.3±0.6	1.3±0.5	-0.522	0.603
5	1.4±0.6	1.5±0.5	-0.82	0.415
6	1.4±0.5	1.4±0.4	-0.329	0.743
7	1.4±0.6	1.5±0.6	-1.397	0.166
8	1.5±0.6	1.5±0.5	-0.484	0.630

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Table 3 PI values at Different Gestational Ages

GA(W)	PI	<i>F</i>	<i>P</i>
32	1.2±0.4		
33	1.3±0.5		
34	1.4±0.6	6.233	<0.001
35	1.5±0.6		
36	1.4±0.6		

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Table 4 Results of Correlation Analysis

Variables	HR (bpm)	SpO ₂ (%)	PI	Systolic BP (mmHg)	Distolic BP (mmHg)	RR (bpm)	T (°C)	GA (W)	BW (g)	Time (day)
HR(bpm)	1.000									
SpO ₂ (%)	-0.014	1.000								
PI	0.067*	-0.113*	1.000							
Systolic BP (mmHg)	0.102*	0.026	0.025	1.000						
Distolic BP (mmHg)	0.114*	0.017	-0.018	0.296*	1.000					
RR(bpm)	0.110*	-0.036	0.019	0.069*	-0.055	1.000				
T(°C)	0.107*	-0.085*	0.002	0.023	0.026	0.081*	1.000			
GA(W)	-0.116*	-0.014	0.097*	0.027	-0.004	0.058	0.073*	1.000		
BW(g)	-0.102*	-0.072*	-0.011	0.084*	0.028	0.124*	0.030	0.287*	1.000	
Time (day)	0.361*	0.028	0.284*	0.197*	0.147*	0.017	0.004	-0.058	-0.011	1.000

427 Pearson test, *p<0.01

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Table 5 PI Value under Multiple Linear Regression Model

	<i>B</i>	Std. Error	<i>Beta</i>	<i>t</i>	<i>P</i>	<i>VIF</i>
Constant	2.253	0.760		2.965	0.003	
GA(W)	0.057	0.013	0.108	4.501	0.000	1.014
HR(bpm)	0.001	0.001	-0.032	-1.235	0.217	1.165
SpO ₂ (%)	-0.030	0.006	-0.120	-5.048	0.000	1.002
Time(day)	0.062	0.005	0.305	11.938	0.000	1.153