Research Article

Safety of Autologous Cord Blood Cells for Preterms: A Descriptive Study

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Received 23 March 2018; Revised 26 June 2018; Accepted 19 July 2018; Published 15 August 2018

Academic Editor: Leonora Buzanska

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Background. Preterm birth complications are one of the leading causes of death among children under 5 years of age. Despite advances in medical care, many survivors face a lifetime of disability, including mental and physical retardation, and chronic lung disease. More recently, both allogenic and autogenic cord blood cells have been applied in the treatment of neonatal conditions such as hypoxic-ischemic encephalopathy (HIE) and bronchopulmonary dysplasia (BPD).

Objective. To assess the safety of autologous, volume- and red blood cell- (RBC-) reduced, noncryopreserved umbilical cord blood (UCB) cell infusion for preterm infants.

Method. This study was a phase I, open-label, single-arm, single-center trial to evaluate the safety of autologous, volume- and RBC-reduced, noncryopreserved UCB cell (5 × 10^7 cells/kg) infusion for preterm infants <37 weeks gestational age. UCB cell characteristics, pre- and postinfusion vital signs, and laboratory investigations were recorded. Clinical data including mortality rates and preterm complications were recorded. Results. After processing, (22.67 ± 4.05) ml UCB cells in volume, (2.67 ± 2.00) × 10^8 cells in number, with (22.67 ± 4.05) × 10^6 CD34+, (3.72 ± 3.25) × 10^5 colony forming cells (CFU-GM), and (99.7 ± 0.17%) vitality were infused to 15 preterm infants within 8 hours after birth. No adverse effects were noticed during treatment. All fifteen patients who received UCB infusion survived. The duration of hospitalization ranged from 4 to 65 (30 ± 23.6) days. Regarding preterm complications, no BPD, necrotizing enterocolitis (NEC), retinopathy of prematurity (ROP) was observed. There were 1/15 (7%) infant with intraventricular hemorrhage (IVH), 5/15 (33.3%) infants with ventilation-associated pneumonia, and 10/15 (66.67%) with anemia, respectively.

Conclusions. Collection, preparation, and infusion of fresh autologous UCB cells to preterm infants is feasible and safe. Adequately powered randomized controlled studies are needed.

1. Introduction

Preterm delivery is a global health problem. The rate of preterm birth ranges from 5% to 18% of babies born across 184 countries. An estimated 15 million babies are born preterm every year [1]. Preterm birth complications are the leading cause of death among children under 5 years of age, which are responsible for nearly 1 million deaths in 2015. The morbidity associated with preterm birth often extends to later life, resulting in enormous physical, psychological, and economic costs [2]. Inflammation, ischemia, and free radical toxicity lead to multiorgan damage in preterm infants, characterized by reduced numbers of tissue cells, blood vessels, and progenitor cell [3–6]. Current management has been shown to reduce preterm complications and overall morbidity. However, many survivors still face a lifetime of disability, including mental and physical retardation, and chronic lung disease [1]. It has been reported that among infants born with gestational ages of 22 to 28 weeks, 16% are complicated with severe intraventricular hemorrhage (IVH), 36% with late-onset sepsis (LOS), and 68% with bronchopulmonary dysplasia (BPD) [7]. The current treatments such as pulmonary
surfactant administration, noninvasive respiratory support, and antibiotic administration are single-organ or symptom-targeted. Neonatologists are in urgent need for new systemic multiorgan-targeted treatments.

Human umbilical cord blood cells (UCBC) are abundant in stem cells. These primitive cells can home into damaged tissues, produce anti-inflammatory and immune-modulatory factors by paracrine effects, and differentiate into tissue cells [8]. Potential effects on respiratory distress syndrome (RDS), sepsis, and hypoxic-ischemic brain damage have been suggested in animal models [9–11]. Recently, these potential effects have been proved to be safe and feasible in clinical applications. Allogenic umbilical cord blood-derived mesenchymal stem cells (MSCs) have been applied in adults with acute RDS [12] and preterm infants with BPD [13], and autologous UCBC has been applied to neonates with HIE [14].

Recently, delayed cord clamping in premature neonates have been reported to improve neonatal mortality and morbidity. The American College of Obstetricians and Gynecologists now recommends a delay in umbilical cord clamping in preterm infants for at least 30–60 seconds after birth [15]. The potential mechanism was that delayed cord clamping was accompanied by an increased supply of RBCs and valuable progenitor cells [16].

Based on these evidence, we hypothesized that autologous cord blood infusion was safe for preterm infants. We report the outcomes of the infusion of autologous, volume- and RBC-reduced, noncryopreserved cord blood cell to 15 premature neonates.

2. Methods

This study was a phase I, open-label, single-arm, single-center trial to evaluate the safety of autologous, volume- and red blood cell- (RBC-) reduced, noncryopreserved umbilical cord blood cells (UCBC) (5 × 10^7 cells/kg) infusion for preterm infants <37 weeks gestational age.

2.1. Patients. We initiated this pilot study in December 2009. Inborn infants admitted to the Neonatal Intensive Care Unit (NICU) of Guangdong Women and Children’s Hospital were eligible if they were (1) preterm: <37 weeks gestation, (2) without congenital abnormalities, (3) without maternal chorioamnionitis, (4) had available UCB, and (5) the mother was negative for hepatitis B (HBsAg and/or HBeAg) and C virus (anti-HCV), syphilis, HIV (anti-HIV-1 and -2) and IgM against Cytomegalovirus, rubella, toxoplasma, and herpes simplex virus. The study protocol was approved by the ethics committee of Guangdong Women and Children’s Hospital. All patients in the study were given an intensive care therapy in accordance with the departmental guidelines which included therapies including positive pressure mechanical ventilation, noninvasive respiratory support, oxygen therapy, and exogenous surfactant (Curosurf, Chiesi, Parma, Italy) replacement. Chest radiographs were performed at admission and 8 hours after CBT on the first day of life in all surviving patients. Blood gas was monitored every 24 hours until weaning from ventilation. All clinical diagnoses were defined according to a standard reference [17]. Soon after the preterm infant was delivered, written consent was signed by the parents, and autologous cord blood infusion was applied to the baby in addition to routine pulmonary surfactant replacement and mechanical ventilation support as indicated.

2.2. Cord Blood Process. Guangdong Cord Blood and Stem Cell Bank is a public provincial blood bank affiliated to the Guangdong Women and Children’s Hospital, which collects cord blood of every delivery in this hospital. Therefore, the cord blood of all the subjects had been routinely collected during the delivery. The procedure of cord blood collection and transfusion was performed in accordance with the cord blood bank guidance [18]. The umbilical cord was clamped for the collection using a blood-collection bag (WEGO, China) containing 28 ml of citrate-phosphate-dextrose-anticoagulant right after the baby was born and before the placenta was delivered. The umbilical vein was sterilized and punctured with a 17-gauge needle. UCB collections were made by trained obstetricians or cord blood bank collection staff who were present at the hospital during weekdays for 8–12 hours per day. When collection was completed, the blood bag tubing was closed and sealed. Cord blood labeled with the full name of the donor, group type, and volume of the blood product was stored in 4 degree and sent to the Cord Blood and Stem Cell Bank for processing immediately. Before processing, 2 ml samples were taken from all collected CB units to test for the presence of virus (human immunodeficiency virus, hepatitis B virus, hepatitis C virus, and Cytomegalovirus) and bacterial infections (including Treponema Pallidum). A sample of peripheral blood was collected from the mother and tested for the presence of maternal transmissible diseases. And the results were obtained soon before the transfusion started. After the sample was taken, it was volume- and RBC-reduced after 30 minute incubation with 6% Hespan (Bethlehem, USA) following established CBB procedures using the SEPAX S-100 automated processing system (Biosafe, Geneva, Switzerland) if the unit contained >30 ml of UCB or manually if the unit was <30 ml. The mononuclear layer was isolated by density gradient centrifugation (1000g, 30 min, RT, Beckman, American), then was transferred to cryobags. Excessive nucleated cell-poor plasma was expelled. Meanwhile, MNC count, CD34 cell, CFU-GM, and steril- ity detection (Sheldon Manufacturing Inc., Cornelius, OR, USA) were performed. Cell viability was measured via 7-aminoactinomycin D (7-AAD) detection kit through flow cytometry analysis (BD Bioscience, USA). All infusions were administered in Guangdong Women and Children Hospital. Infusate and subject identities were double-checked by the research and clinical nursing staff. Infusions were also monitored by the research and clinical staff. Cells were infused over 15 minutes, followed by a 2 ml saline flush to clear the intravascular line.

2.2.1. Assessment of Safety. Shortly, before, during, and until 24 hours after transfusion, heart rate, systolic, diastolic, and mean arterial blood pressure and arterial blood oxygen saturation level were monitored in peripheral blood continually.
and documented. Moreover, laboratory investigations in peripheral blood were monitored and kept stable during the whole treatment period, detailed in Table 1. Infusion reactions and signs of circulatory overload were checked.

2.2.2. Results. From January 1, 2009, till June 5, 2016, fifteen infants were enrolled for the treatment, gestational age ranged from 28 2/7 to 34 1/7 (31.2 ± 1.62) weeks and birth weight ranged from 1200 to 2220 (1582.7 ± 252.8) grams; 12/15 (80%) were delivered by cesarean section. All 15 patients who received the cord blood infusion survived. The duration of hospitalization ranged from 4 to 65 (30 ± 23.6) days. Details were shown in Table 2.

2.3. Characteristics of Cord Blood Processing. Cord blood volume collected ranged from 27 to 76 ml, mean (47.13 ± 19.10) ml; volume postprocessing ranged from 16 to 30 ml, mean (22.67 ± 4.05) ml; cells collected ranged from 0.97 to 8.11 (×10⁸), mean (3.10 ± 2.17 × 10⁸); cells postprocessing ranged from 0.86 to 7.83 (×10⁸), mean (2.67 ± 2.00 × 10⁸); cells concentration postprocessing ranged from 5.85 to 40.8 × 10⁹/ml, mean (13.10 ± 10.35 × 10⁹/ml); CFU-GM ranged from 0.72 to 11.27 (×10⁵), mean (3.72 ± 3.25 × 10⁵); amount of CD34+ cells in units varied widely, ranged from 0.1 to 16.22 × 10⁶, mean (22.67 ± 4.05) ml; and viability of postprocessing units was high, ranged from 99.5 to 100%, mean (99.7 ± 0.17%). Details are shown in Table 3.

2.4. Infusion. Infused NC ranged from 4.48 to 5.0 × 10⁷/kg, mean (4.97 ± 0.13 × 10⁷/kg); time between collection (birth) and initiation of infusion ranged from 4.5 to 9 hours after birth, mean (6.77 ± 1.52) h; infused volume ranged from 2 to 28 ml, mean (10.27 ± 6.18 ml); pathogen detection (including bacteria culture, fungus culture, human immunodeficiency virus, hepatitis B virus, hepatitis C virus, Cytomegalovirus, and Treponema Pallidum) results were all negative.

2.5. Cord Blood Safety. The patient’s vital signs and laboratory investigations were monitored during the whole treatment period, details were shown in Table 1. No significant infusion reactions were noted. No signs of circulatory overload and graft-versus-host disease (GVHD) were detected. Heart rate, mean arterial pressure, and oxygen saturation did not vary significantly before and after infusions.

2.6. Clinical Presentation and Complications

2.6.1. Mortality. The fifteen patients who received infusions all survived.

2.6.2. Nervous System. Three patients had birth asphyxia, among them one suffered from IVH. None of the patients developed abnormal clinical features of central nervous system disorders such as convulsions, apnea, or dysphagia.

2.6.3. Respiratory System. 12/15 (80%) presented with tachypnea and grunting soon after birth. The infants were diagnosed with RDS, 2/15 (13.3%) cases were grade I, 5/15 (33.3%) cases were grade II, 6/15 (40%) cases were grade III, and 2/15 (13.3%) cases were grade IV; and 12/15 (80%) received one dose PS replacement and 8/15 (53.3%) received intubation-surfactant replacement extubation–nasal continuous positive airway pressure (INSURE) therapy; however, one patient needed reintubation. 4/15 (26.7%) received mechanical ventilation; the median duration was 3.2 ± 1.8 days. The duration of oxygen therapy was (5.3 ± 3.0) days. No patient suffered from BPD, and chest radiographs showed improvement.

2.6.4. Infection. 5/15 suffered from ventilation-associated pneumonia (VAP), of which two were cases of Klebsiella pneumonia, one was a case of Pseudomonas aeruginosa pneumonia, one was a case of Acinetobacter baumannii pneumonia, and 1 suffered from late onset sepsis, infected with Klebsiella pneumonia proved by blood culture.

2.6.5. ROP. No patients suffered from ROP.

2.6.6. NEC. No patients suffered from NEC.

2.6.7. Anemia. 10/15 (66.67%) suffered from anemia (≤140 g/l); 2/15 (13.33%) needed RBC transfusion.

3. Discussion

In our study, we treated 15 preterm infants with autologous, volume and RBC-reduced cord blood cells. The treatment was started within 8 hours after birth. No adverse effect of cell therapy was noticed. No patient died during treatment. No preterm complications such as BPD, NEC, or ROP were observed. Our study presents preliminary data on the safety of autologous cord blood cell therapy in preterm infants. We postulated that several factors contributed to the safety issue, among them, the most important one was the autologous cell source. Based on the autologous cell source, no GVHD-related complication was observed. Moreover, autologous cell source avoided ethical issues. A second factor that contributed to the safety issue was cord blood minimal-processing procedure. In our study, only density gradient centrifugation was employed to separate nucleated cells. Since our cell infusions were started within 8 hours after birth, no cryopreservation was needed; thus, no chemicals were added into the cord blood cells for cryopreservation. This minimal-processing procedure and immediate transfusion after processing helped to avoid contamination and possible chemical toxicity. It also alleviated decreased of viability which may happen during storage.

In our study, mean (47.13 ± 19.10) ml cord blood with a total TNC of (3.10 ± 2.17) × 10⁸ mononuclear cells was collected before processing. After processing, cord blood volume and TNC were reduced to (22.67 ± 4.05) ml and (2.67 ± 2.00) × 10⁸, respectively, including (22.67 ± 4.05) × 10⁶ CD34+ and (3.72 ± 3.25) × 10⁴ colony forming cells (CFU-GM) in a vitality (99.7 ± 0.17%).

Recently, delayed umbilical cord clamping 30–60 seconds after birth in preterms had been recommended by the American College of Obstetricians and Gynecologists and had been reported to reduce preterm-related complications. It is possible that delayed cord clamping increases supply of RBC and valuable stem and progenitor cells (SPC), thus may improve mortality and morbidity in premature neonates.
Table 1: Clinical findings previous and post infusion.

| Serial number | HB Pre | HCT Pre | WBC Pre | PLT Pre | pH Pre | Post-12 h | Post-24 h | PO2 Pre | Post-12 h | Post-24 h | PCO2 Pre | Post-12 h | Post-24 h | Fio2 Pre | Post-12 h | Post-24 h | Post-12 h | Post-24 h |
|---------------|-------|--------|--------|--------|-------|----------|----------|-------|----------|----------|-------|----------|----------|--------|----------|----------|----------|----------|
| 1             | 177   | 144    | 37.3   | 10.8   | 9.2   | 304      | 335      | 7.34  | 7.27     | 7.45     | 9.3   | 8.9      | 8.9      | 4.8    | 7        | 3.2      | 0.3     | 0.21     |
| 2             | 167   | 137    | 37.4   | 10.9   | 10.2  | 284      | 243      | 7.26  | 7.4      | 7.37     | 6.7   | 12.7     | 6.4      | 6.8    | 3.7      | 4.6      | 0.4     | 0.25     |
| 3             | 135   | 132    | 37.3   | 14.9   | 13.2  | 252      | 248      | 7.25  | 7.2      | 7.33     | 4.64  | 5.07     | 9.9      | 7.25   | 8.02     | 4.26     | 0.35    | 0.25     |
| 4             | 145   | 144    | 39.9   | 8.8    | 10.1  | 405      | 336      | 7.25  | 7.28     | 7.39     | 6.5   | 9.1      | 6.1      | 7.6    | 6.6      | 4.4      | 0.35    | 0.21     |
| 5             | 171   | 180    | 50.6   | 9.6    | 17.9  | 265      | 161      | 7.42  | 7.39     | 7.34     | 3.9   | 4.5      | 6.4      | 3.7    | 3.9      | 5.1      | 0.55    | 0.3      |
| 6             | 120   | 116    | 35     | 14     | 13.5  | 320      | 256      | 7.34  | 7.36     | 7.49     | 8.3   | 6.6      | 5.6      | 5.9    | 4.8      | 3.2      | 0.45    | 0.21     |
| 7             | 148   | 139    | 39.1   | 35     | 14.2  | 274      | 446      | 7.3   | 7.34     | 7.37     | 11.6  | 11.3     | 6.95     | 5.1    | 4.28     | 4.34     | 0.35    | 0.21     |
| 8             | 212   | 203    | 60.5   | 54.3   | 13.7  | 14      | 413      | 420   | 7.42     | 7.28     | 7.45  | 10.48    | 10.25    | 8.31   | 4.78     | 4.82     | 3.62    | 0.25     |
| 9             | 149   | 136    | 43.7   | 42.1   | 9.62  | 10.3     | 338      | 340   | 7.28     | 7.42     | 7.43  | 9.77     | 8.23     | 7.14   | 7.82     | 4.5      | 4.23    | 0.25     |
| 10            | 166   | 158    | 47.6   | 46.7   | 12.3  | 16.3     | 271      | 189   | 7.32     | 7.3     | 7.3   | 9.84     | 10.2     | 8.13   | 6.66     | 5.45     | 6.03    | 0.5      |
| 11            | 164   | 109    | 31.8   | 14.4   | 9.4   | 229      | 154      | 7.34  | 7.35     | 7.36     | 6.4   | 5.3      | 5.2      | 4.9    | 4.5      | 0.5      | 0.3     | 0.21     |
| 12            | 169   | 136    | 49.9   | 7.1    | 17    | 296      | 318      | 7.32  | 7.5      | 7.37     | 9     | 14       | 7.6      | 4.7    | 2.5      | 3.4      | 0.25    | 0.21     |
| 13            | 159   | 105    | 31.2   | 12.9   | 10.6  | 329      | 465      | 7.28  | 7.5      | 7.48     | 8.3   | 6.2      | 8        | 7.6    | 3        | 4.1      | 0.3     | 0.21     |
| 14            | 142   | 136    | 46     | 42     | 11.8  | 9.6      | 235      | 245   | 7.39     | 7.4      | 7.58  | 10.06    | 9.82     | 6.17   | 5.3      | 4.82     | 2.92    | 0.25     |
| 15            | 136   | 122    | 48     | 46.3   | 11.3  | 9.3      | 236      | 240   | 7.32     | 7.36     | 7.4   | 11.1     | 10.95    | 10.32  | 6.47     | 5.37     | 4.83    | 0.45     |
| Mean          | 157   | 139    | 46.3   | 39.3   | 13.1  | 12.3     | 296.7    | 293.1 | 7.32     | 7.36     | 7.4   | 8.39     | 8.87     | 7.39   | 5.98     | 4.91     | 4.18    | 0.37     |
| SD            | 22.04 | 25.5   | 6.18   | 9.30   | 6.4   | 3.03     | 56.70    | 97.88 | 0.06     | 0.08     | 2.31  | 2.86     | 1.54      | 1.28   | 1.5      | 0.82      | 0.10    | 0.05     |
|   | Sex | GA (weeks) | BW (g) | Delivery mode | APGAR | 1 minute | 5 minute | RDS grade | PS | Duration of mechanical ventilation (D) | Duration of oxygen therapy (D) | Reintubation | Time return to BW (D) | 1 | Time return to BW (D) | BPD | VAP | IVH | ROP | LOS | NEC | Duration of hospitalization (D) | Prognosis |
|---|-----|-----------|--------|---------------|-------|-----------|----------|-----------|----|--------------------------------------|-------------------------------|--------------|----------------------------|---|-----------------------|-----|-----|-----|-----|-----|-----|--------------------------------|----------|
| 1 | M   | 30 + 5    | 1630   | CS            | 9     | 9         | 3        | Y         | 1  | 5                                    | 5                             | N            | 20                         | 5  | Nil                   | Nil | Nil | Nil | Nil | Nil | Nil | 18                            | Cured    |
| 2 | M   | 30 + 5    | 1510   | CS            | 5     | 9         | 4        | Y         | 1  | 3                                    | 9                             | N            | 21                         | 5  | Nil                   | Nil | Nil | Nil | Nil | Nil | Nil | 53                            | Cured    |
| 3 | M   | 33       | 1630   | VD            | 8     | 6         | 3        | Y         | 1  | 4                                    | 11                            | N            | 36                         | 5  | Nil                   | Nil | Nil | Nil | Nil | Nil | Nil | 46                            | Cured    |
| 4 | M   | 31 + 2   | 1510   | VD            | 9     | 10        | 2        | Y         | 1  | 2.5                                  | 3                             | N            | 20                         | 5  | Nil                   | Nil | Y   | Nil | Y   | Nil | Nil | 65                            | Cured    |
| 5 | F   | 28 + 2   | 1400   | CS            | 9     | 10        | 3        | Y         | 1  | 4                                    | 9                             | Y            | 32                         | 5  | Nil                   | Y   | Nil | Nil | Nil | Nil | Nil | 30                            | Cured    |
| 6 | M   | 30 + 1   | 1600   | CS            | 8     | 10        | 4        | Y         | 1  | 2.5                                  | 4                             | N            | 26                         | 5  | Nil                   | Y   | Nil | Nil | Y   | Nil | Nil | 4                             | Cured    |
| 7 | M   | 29 + 1   | 1450   | CS            | 9     | 10        | 2        | Y         | 1  | 7                                    | 8                             | N            | 28                         | 5  | Nil                   | Nil | Nil | Nil | Nil | Nil | Nil | 10                            | Cured    |
| 8 | M   | 31 + 5   | 1700   | CS            | 8     | 9         | 1        | Y         | 1  | 2                                    | 2                             | N            | 14                         | 5  | Nil                   | Nil | Nil | Nil | Nil | Nil | Nil | 47                            | Cured    |
| 9 | M   | 32 + 5   | 1940   | CS            | 9     | 10        | 2        | Y         | 1  | 3                                    | 3                             | N            | 13                         | 5  | Nil                   | Nil | Nil | Nil | Nil | Nil | Nil | 24                            | Cured    |
| 10| M   | 29 + 6   | 1400   | CS            | 10    | 10        | 1        | N         | 0  | 0                                    | 3                             | N            | 14                         | 5  | Nil                   | Y   | Nil | Nil | Nil | Nil | Nil | 82                            | Cured    |
| 11| M   | 32 + 5   | 1660   | CS            | 9     | 10        | 3        | Y         | 1  | 5.5                                   | 9                             | N            | 12                         | 5  | Nil                   | Nil | Nil | Nil | Nil | Nil | Nil | 23                            | Cured    |
| 12| M   | 31       | 1240   | CS            | 9     | 9         | 2        | Y         | 1  | 4                                    | 5                             | N            | 14                         | 5  | Nil                   | Y   | Y   | Nil | Y   | Nil | Nil | 20                            | Cured    |
| 13| M   | 30       | 1600   | VD            | 9     | 9         | 3        | Y         | 1  | 3                                    | 4.5                            | N            | 12                         | 5  | Nil                   | Nil | Nil | Nil | Nil | Nil | Nil | 19                            | Cured    |
| 14| F   | 32 + 2   | 1450   | CS            | 9     | 10        | 2        | N         | 0  | 1.5                                   | 2.5                            | N            | 16                         | 5  | Nil                   | Nil | Nil | Nil | Nil | Nil | Nil | 5                             | Cured    |
| 15| F   | 34 + 1   | 2220   | VD            | 9     | 10        | 3        | N         | 0  | 1                                    | 2                             | N            | 14                         | 5  | Nil                   | Nil | Nil | Nil | Nil | Nil | Nil | 4                             | Cured    |
|   | **MEAN ± SD** |          |        |               |       |           |          |           |    | **3.2 ± 1.8**                        | **5.3 ± 3.0**                |              | **30 ± 23.6**          |    |                      |    |    |    |    |    |    |                          |          |
| No | Volume collected | Volume postprocessing | Cells collected (108) | Cells postprocessing (108) | Cell concentration postprocessing × 10^6/ml | CFU-GM/10^5 | CD34+/10^6 | Viability | Infused NC × 10^7/kg | Infused age (H) | Infused volume | Pathogen detection |
|----|------------------|-----------------------|----------------------|---------------------------|-------------------------------------------|-------------|-----------|-----------|-------------------|----------------|---------------|-------------------|
| 1  | 34               | 22                    | 1.82                 | 1.46                      | 6.65                                      | 1.84        | 1.1       | 100       | 4.48              | 5              | 11            | Negative          |
| 2  | 35               | 20                    | 1.6                  | 1.32                      | 5.82                                      | 1.98        | 0.7       | 99.5      | 5                 | 6              | 13            | Negative          |
| 3  | 61               | 24                    | 2.44                 | 1.67                      | 6.95                                      | 2.3         | 1.14      | 99.8      | 5                 | 7              | 12            | Negative          |
| 4  | 33               | 16                    | 0.97                 | 0.86                      | 5.4                                       | 0.72        | 0.31      | 99.8      | 5                 | 4              | 14            | Negative          |
| 5  | 36               | 22                    | 1.43                 | 1.21                      | 5.5                                       | 1.57        | 0.59      | 99.8      | 5                 | 9              | 13            | Negative          |
| 6  | 28               | 16                    | 1.43                 | 1.23                      | 7.7                                       | 1.53        | 0.9       | 99.5      | 5                 | 4.5            | 11            | Negative          |
| 7  | 55               | 28                    | 5.81                 | 5.43                      | 26.3                                      | 11.27       | 5.69      | 99.8      | 5                 | 6              | 3             | Negative          |
| 8  | 65               | 20                    | 2.99                 | 2.37                      | 11.85                                     | 1.66        | 1.66      | 99.9      | 5                 | 7              | 8             | Negative          |
| 9  | 81               | 20                    | 6.14                 | 4.33                      | 21.65                                     | 9.96        | 3.9       | 99.9      | 5                 | 8              | 5             | Negative          |
| 10 | 27               | 28                    | 2.21                 | 1.97                      | 7.05                                      | 3.35        | 0.55      | 99.8      | 5                 | 8              | 28            | Negative          |
| 11 | 46               | 24                    | 2.96                 | 2.65                      | 13.6                                      | 3.32        | 1.01      | 99.5      | 5                 | 7              | 6             | Negative          |
| 12 | 37               | 24                    | 1.51                 | 1.4                       | 5.85                                      | 1.04        | 0.08      | 99.9      | 5                 | 9              | 12            | Negative          |
| 13 | 24               | 22                    | 1.72                 | 1.58                      | 8.53                                      | 2.82        | 1.1       | 99.8      | 5                 | 6              | 9             | Negative          |
| 14 | 69               | 24                    | 8.11                 | 7.83                      | 40.8                                      | 6.08        | 16.22     | 99.8      | 5                 | 7              | 2             | Negative          |
| 15 | 76               | 30                    | 5.3                  | 4.7                       | 22.8                                      | 6.39        | 1         | 99.5      | 5                 | 8              | 7             | Negative          |
| Mean | 47.13            | 22.67                 | 3.10                 | 2.67                      | 13.10                                     | 3.72        | 2.40      | 99.7      | 4.97              | 6.77           | 10.27         | Negative          |
| SD  | 19.10            | 4.05                  | 2.17                 | 2.00                      | 10.35                                     | 3.25        | 4.10      | 0.17      | 0.13              | 1.52           | 6.18         |                   |
[15, 16]. However, delayed umbilical cord clamping mainly supply RBC instead of MNC to the infants. Therefore, we used noncryopreserved autologous cord blood cells infusion soon after preterm birth which contains mainly MNC with a lower volume. As is known that SPC mainly exits in MNC layer. It is considered as the most effective component in cord blood and strongly associated with a lower risk of developing preterm complications [19, 20]. Further, in view of the vulnerable heart function of preterm infants, they could only accept limited volume of infusion. Therefore, to achieve more MNC in a lower volume, we used volume- and RBC-reduced cord blood cells in our study to decrease the burden to the heart.

BPD is the main complication that contributes to morbidity and mortality in extremely premature newborns. It has been reported that among infants born with gestational ages of 22 to 28 weeks, 68% suffered from BPD [7]. The pathophysiologic features of BPD include abnormal lung growth characterized by reduced numbers of alveoli, blood vessels, prominent fibrosis, and secondary pulmonary hypertension [21]. Cord blood angiogenic progenitor cells and endothelial progenitor cells were reported to be decreased in preterm infants with BPD [5, 6]. Current therapy for BPD included noninvasive ventilation strategies, inhaled nitric oxide, antioxidants, vitamin A, caffeine, and corticosteroids. However, so far there are no specific therapies that have been widely adopted. It has been reported that MSCs may release anti-inflammatory paracrine factors. These factors have effect on both lung injury and sepsis [22, 23]. Perventricular leukomalacia (PVL) is another severe preterm complication, affecting 16% infants born with gestational age of 22 to 28 weeks [7]. PVL reflected perinatal damage from inflammation and oxidation to the developing brain, which was one of main reasons responsible for cerebral palsy [24]. Current therapy for hypoxic-ischemic damage was hypothermia. However, hypothermia therapy is contraindicated in preterm infants because of their immature thermoregulation [25, 26]. HUCBC has been shown to be effective in newborn models of hypoxic injury [4]; furthermore, autologous intravenous UCB infusion is safe and feasible in neonates with HIE and young children with acquired neurological disorders [7, 27]. In our study, there was one case complicated with IVH; however, no clinical presentation related to the central nervous system was observed. The underlying mechanism might be that the UCBC migrate to damaged sites, form anti-inflammatory or immunomodulatory factors, then proliferate into neurons [28, 29]. Sepsis is a common and major cause of death in preterm infants [30, 31]. Among very low birth weight infants (VLBW; < 1500 g), rates of sepsis range between 11% and 46% [32]. Neonatal sepsis and systemic inflammatory response syndrome (SIRS) are associated with brain damage [31, 33]. Current therapy for neonatal sepsis is the antibiotic administration. However, antibiotic resistance is a therapeutic problem in preterms. Substantial evidence from models of both lung injury and sepsis suggested that MSCs have an anti-inflammatory effect on host tissue, partly through the release of paracrine factors, reprogramming host macrophages [22, 23, 34]. In addition, MSCs reduced alveolar bacterial counts and improved alveolar macrophage phagocytosis after direct bacterial injury mediated by FGF7, LL-37 [35, 36]. All these investigations have laid down the foundation for cord blood cell therapy for preterms with pulmonary disorders complicated with sepsis and brain damage. NEC and ROP are two main complications of preterm infants; however, there were no cases observed in our study; the underlying reason might be the limited enrolled number and relatively large gestational age of infants enrolled in our study. To achieve more evidence regarding NEC and ROP, a large cohort study will be needed in our future study.

Studies on safety and feasibility of whole autologous cord blood transplant in preterm were also reported [19, 20]. Rudnicki and colleagues compared whole autologous cord blood infusion with allogeneic red blood cells in the treatment of preterm with anemia and showed autologous CB infusion was as effective and safe as allogeneic RBC transfusion [19]. In our study, 10/15 (66.67%) infants suffered from anemia (≤140 g/l); 2/15 (13.33%) needed RBC transfusion. Allogeneic RBCs transfusion is the main therapy for severe anemia [17]. In this study, we explore the safety of volume- and RBC-reduced cord blood cells to treat preterm infants. Both autologous cord blood infusion possess its advantage. However, further multiple center randomized controlled studies are needed regarding short-term and long-term outcomes.

Regarding the administration route, there were reports on the advantage of damaged site administration when compared to intravenous infusion. However, on the one hand, the potential preterm complications were due to multiorgan damage. To achieve the multiorgan-targeted effect, we used intravenous infusion as the administration route, which may result in cells being trapped in organs such as lung and brain. On the other hand, in the report supported site administration, allogeneic-MSC was used in intratracheal administration to treat hyperoxia-induced lung damage; it seemed to attenuate the side effects of rejection.

In our study, we chose the infusion timing to be very soon after birth which is within the first 8 postnatal hours. Although some infants were delivered at midnight, we tried to process as early as within 8 hours after their birth. As it had been reported that it would take more than 1 week for progenitor cells to differentiate into damaged tissue cells, we administered CBC during the first hours after birth, so that it might provide enough time for these cells for differentiation.

In conclusion, we demonstrated autologous, volume- and RBC-reduced, noncryopreserved cord blood cells transfusion soon after birth was safe and feasible in preterm infants. Autologous cord blood infusion avoid GVHD; meanwhile, the reduced volume would protect the fragile cardio-function of the preterm infants. This autologous, volume- and RBC-reduced, method guaranteed the safety of application. In addition, this was an autologous transfusion instead of "cell transplantation therapy", and therefore it was not regulated by the FDA regulation licensing public cord blood banks distributing unrelated banked cord blood units for allogeneic transplantation in 2012. However, our study was the single-center descriptive study with limited number of preterm infants, further multicenter randomized controlled trials are needed to prove the effectiveness.
Data Availability

The data used to support the findings of this study are available from the corresponding author upon request.

Conflicts of Interest

The authors indicated no potential conflicts of interest.

Authors’ Contributions

Jie Yang and Zhuxiao Ren have equivalent contribution to the study.

Acknowledgments

This work was supported by Guangzhou Technology Program (grant numbers 201707010398, 201804010380).

References

[1] H. Blencowe, S. Cousens, M. Z. Oestergaard et al., “National, regional, and worldwide estimates of preterm birth rates in the year 2010 with time trends since 1990 for selected countries: a systematic analysis and implications,” The Lancet, vol. 379, no. 9832, pp. 2162–2172, 2012.

[2] S. Petrou, Z. Mehta, C. Hockley, P. Cook-Mozaffari, J. Henderson, and M. Goldacre, “The impact of preterm birth on hospital inpatient admissions and costs during the first 5 years of life,” Pediatrics, vol. 112, no. 6, pp. 1290–1297, 2003.

[3] O. Khwaja and J. J. Volpe, “Pathogenesis of cerebral white matter injury of prematurity,” Archives of Disease in Childhood - Fetal and Neonatal Edition, vol. 93, no. 2, pp. F153–F161, 2008.

[4] S. A. Mitsialis and S. Kourembanas, “Stem cell-based therapies for the newborn lung and brain: possibilities and challenges,” Seminars in Perinatology, vol. 40, no. 3, pp. 138–151, 2016.

[5] C. D. Baker, V. Balasubramaniam, P. M. Mourani et al., "Cord blood angiogenic progenitor cells are decreased in bronchopulmonary dysplasia," The European Respiratory Journal, vol. 40, no. 6, pp. 1516–1522, 2012.

[6] A. Borghesi, M. Massa, R. Campanelli et al., “Cirulating endothelial progenitor cells in preterm infants with bronchopulmonary dysplasia,” American Journal of Respiratory and Critical Care Medicine, vol. 180, no. 6, pp. 540–546, 2009.

[7] B. J. Stoll, N. I. Hansen, E. F. Bell et al., “Neonatal outcomes of extremely preterm infants from the NICHD neonatal research network,” Pediatrics, vol. 126, no. 3, pp. 443–456, 2010.

[8] T. C. Lund, A. E. Boitano, C. S. Delaney, E. J. Shpall, and J. E. Wagner, "Advances in umbilical cord blood manipulation—from niche to bedside," Nature Reviews. Clinical Oncology, vol. 12, no. 3, pp. 163–174, 2015.

[9] J. Walter, L. B. Ware, and M. A. Matthey, “Mesenchymal stem cells: mechanisms of potential therapeutic benefit in ARDS and sepsis,” The Lancet Respiratory Medicine, vol. 2, no. 12, pp. 1016–1026, 2014.

[10] R. S. Alphonsie, S. Rajabali, and B. Thebault, "Lung injury in preterm neonates: the role and therapeutic potential of stem cells," Antioxidants & Redox Signaling, vol. 17, no. 7, pp. 1013–1040, 2012.

[11] A. Drobyshyevsky, C. M. Cotten, Z. Shi et al., “Human umbilical cord blood cells ameliorate motor deficits in rabbits in a cerebral palsy model,” Developmental Neuroscience, vol. 37, no. 4-5, pp. 349–362, 2015.

[12] Y. Chang, S. H. Park, J. W. Huh, C. M. Lim, Y. Koh, and S. B. Hong, "Intratracheal administration of umbilical cord blood-derived mesenchymal stem cells in a patient with acute respiratory distress syndrome," Journal of Korean Medical Science, vol. 29, no. 3, pp. 438–440, 2014.

[13] Y. S. Chang, S. Y. Ahn, H. S. Yoo et al., "Mesenchymal stem cells for bronchopulmonary dysplasia: phase 1 dose-escalation clinical trial," The Journal of Pediatrics, vol. 164, no. 5, pp. 966–972.e6, 2014.

[14] C. M. Cotten, A. P. Murtha, R. N. Goldberg et al., “Feasibility of autologous cord blood cells for infants with hypoxic-ischemic encephalopathy,” The Journal of Pediatrics, vol. 164, no. 5, pp. 973–979.e1, 2014.

[15] "Committee opinion no. 684 summary: delayed umbilical cord clamping after birth," Obstetrics & Gynecology, vol. 129, no. 1, pp. 232–233, 2017.

[16] C. Lawton, S. Acosta, N. Watson et al., “Enhancing endogenous stem cells in the newborn via delayed umbilical cord clamping,” Neural Regeneration Research, vol. 10, no. 9, pp. 1359–1362, 2015.

[17] T. Gomella, M. Cummingham, and E. Fabien, Neonatology, Lange, New York, 6th edition, 2009.

[18] National Health and Family Planning Commission of the People’s Republic of China, “Umbilical cord blood and stem cell bank management regulation,” http://www.Moh.Gov.cn/zhuzhan/wsbmgz/201308/9da745d55fe749a2b82b95ee65294b55.shtml (Chinese).

[19] M. Kotowski, Z. Litwinska, P. Klos et al., “Autologous cord blood transfusion in preterm infants—could its humoral effect be the key to control prematurity-related complications? A preliminary study,” Journal of Physiology and Pharmacology, vol. 68, no. 6, pp. 921–927, 2017.

[20] J. Rudnicki, M. P. Kawa, M. Kotowski et al., “Clinical evaluation of the safety and feasibility of whole autologous cord blood transplant as a source of stem and progenitor cells for extremely premature neonates: preliminary report,” Experimental and Clinical Transplantation, vol. 13, no. 6, pp. 563–572, 2015.

[21] W. H. Northway Jr, R. C. Rosan, and D. Y. Porter, “Pulmonary disease following respirator therapy of hyaline-membrane disease — bronchopulmonary dysplasia,” The New England Journal of Medicine, vol. 276, no. 7, pp. 357–368, 1967.

[22] L. A. Ortiz, M. DuTreil, C. Fattman et al., “Interleukin 1 receptor antagonist mediates the antiinflammatory and anti-fibrotic effect of mesenchymal stem cells during lung injury,” Proceedings of the National Academy of Sciences of the United States of America, vol. 104, no. 26, pp. 11002–11007, 2007.

[23] S. Danchuk, J. H. Ylostalo, F. Hossain et al., “Human multipotent stromal cells attenuate lipopolysaccharide-induced acute lung injury in mice via secretion of tumor necrosis factor-a-induced protein 6,” Stem Cell Research & Therapy, vol. 2, no. 3, p. 27, 2011.

[24] A. H. MacLennan, S. C. Thompson, and J. Gecz, “Cerebral palsy: causes, pathways, and the role of genetic variants,” American Journal of Obstetrics and Gynecology, vol. 213, no. 6, pp. 779–788, 2015.

[25] S. Shankaran, A. R. Laptcoop, R. A. Ehrenkranz et al., “Whole-body hypothermia for neonates with hypoxic-ischemic
encephalopathy,” *The New England Journal of Medicine*, vol. 353, no. 15, pp. 1574–1584, 2005.

[26] R. P. Sutsko, K. C. Young, A. Ribeiro et al., “Long-term reparative effects of mesenchymal stem cell therapy following neonatal hyperoxia-induced lung injury,” *Pediatric Research*, vol. 73, no. 1, pp. 46–53, 2013.

[27] W. Sun, L. Buzanska, K. Domanska-Janik, R. J. Salvi, and M. K. Stachowiak, “Voltage-sensitive and ligand-gated channels in differentiating neural stem-like cells derived from the nonhematopoietic fraction of human umbilical cord blood,” *Stem Cells*, vol. 23, no. 7, pp. 931–945, 2005.

[28] A. Saha, S. Buntz, P. Scotland et al., “A cord blood monocyte-derived cell therapy product accelerates brain remyelination,” *JCI Insight*, vol. 1, no. 13, article e86667, 2016.

[29] Z. A. Englander, J. Sun, Laura Case, M. A. Mikati, J. Kurtzberg, and A. W. Song, “Brain structural connectivity increases concurrent with functional improvement: evidence from diffusion tensor MRI in children with cerebral palsy during therapy,” *Neuroimage: Clinical*, vol. 7, pp. 315–324, 2015.

[30] J. E. Lawn, S. Cousens, J. Zupan, and Lancet Neonatal Survival Steering Team, “4 million neonatal deaths: when? Where? Why?,” *The Lancet*, vol. 365, no. 9462, pp. 891–900, 2005.

[31] S. Vergnano, E. Menson, N. Kennea et al., “Neonatal infections in England: the NeonIN surveillance network,” *Archives of Disease in Childhood - Fetal and Neonatal Edition*, vol. 96, no. 1, pp. F9–14, 2011.

[32] B. Alshaikh, K. Yusuf, and R. Sauve, “Neurodevelopmental outcomes of very low birth weight infants with neonatal sepsis: systematic review and meta-analysis,” *Journal of Perinatology*, vol. 33, no. 7, pp. 558–564, 2013.

[33] B. J. Stoll, N. I. Hansen, I. Adams-Chapman et al., “Neurodevelopmental and growth impairment among extremely low-birth-weight infants with neonatal infection,” *Journal of the American Medical Association*, vol. 292, no. 19, pp. 2357–2365, 2004.

[34] K. Németh, A. Leelahavanichkul, P. S. T. Yuen et al., “Bone marrow stromal cells attenuate sepsis via prostaglandin E2-dependent reprogramming of host macrophages to increase their interleukin-10 production,” *Nature Medicine*, vol. 15, no. 1, pp. 42–49, 2009.

[35] J. W. Lee, A. Krasnodembskaya, D. H. McKenna, Y. Song, J. Abbott, and M. A. Matthay, “Therapeutic effects of human mesenchymal stem cells in ex vivo human lungs injured with live bacteria,” *American Journal of Respiratory and Critical Care Medicine*, vol. 187, no. 7, pp. 751–760, 2013.

[36] A. Krasnodembskaya, Y. Song, X. Fang et al., “Antibacterial effect of human mesenchymal stem cells is mediated in part from secretion of the antimicrobial peptide LL-37,” *Stem Cells*, vol. 28, no. 12, pp. 2229–2238, 2010.