Umbilical Cord Blood

D'Agati Alfio, Leanza Vito, Leanza Gianluca, Carbonaro Antonio and Pafumi Carlo^{*}

Department of Obstetrics and Gynaecology, University of Catania, Italy

*Corresponding author

Carlo Pafumi, University of Catania, Department of Obstetrics and Gynaecology, Via Torre del Vescovo, 2 95122, Catania, Italy, Fax: 0039095312001; E-mail: pafumi@unict.it

Submitted: 01 July 2016; Accepted: 15 July 2016; Published: 22 July 2016

Abstract

Umbilical cord blood (UCB) is widely considered as a potential alternative to bone marrow for haematopoietic stem cells. Efforts on UCB research have now taken a step further with the establishment of numerous cord blood banks throughout the world.

In 15 months (from December 1999 to February 2001) we obtained 863 cord blood units by withdrawing residual blood from the placenta. The method of blood collection consisted of puncturing, with an 18-gauge needle, the umbilical cord vein and withdrawing the blood into a sterile bag immediately after clamping and newborn assistance. The blood was collected when the placenta was still in utero. Analysis of the blood was also performed to exclude any bacterial contamination. Among them 429 were collected from newborns delivered vaginally (Group 1) and 434 were collected from caesarean sections (Group 2).

In this study we evaluated the volume of blood collected and the number of CD34+ cells contained in the foetal cord blood and we demonstrated that these two parameters were similar between the two groups; The higher median volume of blood collected from infant delivered through a caesarean section seems mainly due to the different clamping time rather than to the kind of delivery.

Keywords: Haematopoietic stem cells, Umbilical cord, Cryopreservation, Cord blood bank, Nucleated cell counts.

Introduction

A lot of research has been performed all over the world on umbilical cord blood (UCB), and many UCB banks have been instituted in the U.S.A. and Europe, since Gabutti, for the first time, collected and isolated several foetal haematopoietic progenitor cells from the residual placental blood, after delivery and ligation of the umbilical cord. In Italy, also, UCB banks belonging to the GRACE group of Milan connected to the European Netcord have been established. Recently, a new UCB bank has been instituted in Sicily, particularly in Sciacca. It is currently waiting for the official affiliation to the GRACE group (ISO 9002 certification).

Considered as an alternative source of haematopoietic stem cells to bone marrow, UCB is becoming more common in the treatment of malignant and non-malignant haematologic and immunologic diseases. Rubinstein argues that UCB confers some additional advantages when compared to hematopoietic stem cells taken from bone marrow. For example, there is a lower risk of graft-versus-host disease despite HLA mismatch in transplantation with UCB, and there is no risk or pain for the donor. However, some disadvantages have been noted, like a relatively lower amount of stem cells acquired.

Materials and Methods

Since the beginning of its activity, the Obstetrics and Gynaecology Department of the University of Catania has been involved in the collection of UCB samples. From December 1999 to February 2001, 863 UCB units had been collected and sent to Sciacca's bank. Among them, 429 were collected from newborns delivered vaginally, while the remaining were collected from caesarean sections.

All vaginal deliveries were done under epidural or local infiltration anaesthesia and uterotonic drugs were not given before cord clamping. Significant obstetric complications were not recorded. All infants were in the 38th-41st week of gestation, with an Apgar score of 7 or more at the 1st and 5th minute, and remained healthy in the newborn nursery. We measured the time elapsed between birth of the buttocks and the moment when the umbilical cord was clamped. Indications for a caesarean section were previous caesarean sections, or in some cases, cephalopelvic disproportion. Four mothers were in early labour at the time of the section. All infants were at term and none suffered any cardiopulmonary distress during the neonatal period. All sections were performed under epidural anesthesia. At delivery, infants were held above of the level of the surgical table and below the height of maternal abdomen. This level was similar to the level at which vaginally delevered infants were held. The cord was handled carefully so as

to stretch it as little as possible, but it was often necessary to unwind it from the infant's neck or legs. We measured the time elapsed from the birth of the buttocks in a vertex, cephalic presentation when the breech was first delivered through the uterine incision to the first cry and the time when the umbilical cord was clamped. We evaluated the volume of blood collected and the number of CD34+ cells contained in the foetal cord blood according to the birth route: caesarean section or vaginal delivery.

The method of blood collection consisted of puncturing, with an 18-gauge needle, the umbilical cord vein and withdrawing the blood into a sterile bag immediately after clamping and newborn assistance (Figure 1). The blood was collected when the placenta was still in utero. In fact, as shown by Dunn, after vaginal delivery of an infant, the compression of the placenta due to uterine contractions forces blood from the placenta to the infant and hastens placenta transfusion. The blood contained in placenta flowes by gravity from the umbilical cord to the sterile collection bag. Collection was approved by the local ethical committee and an informed consent was obtained from the mother before each collection. The indications for a caesarean delivery were independent from the study. UCB clamping times, independently from the research, were obviously different in the two groups, owing to the technical reasons based on the type of delivery. Clamping times were particularly shorter in caesarean sections with respect to vaginal deliveries. This is particularly important because the time allowed between delivery of an infant and clamping of his/her umbilical cord is the principal determinant of the distribution of blood between the infant and the placenta.

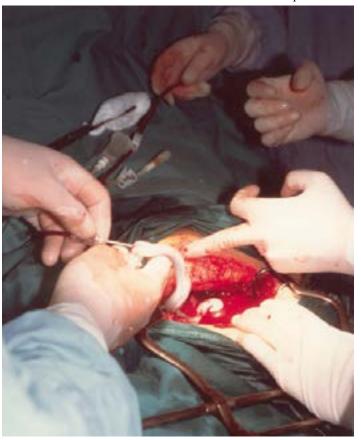


Figure 1: Umbilical cord blood collection in cesarea section.

Results

The results showed that cord blood volume and the number of CD34+ cells collected were similar between the 2 groups (Table 1). The higher median volume of blood collected from infant delivered through a caesarean section seems mainly due to the different clamping time rather than to the kind of delivery.

	Caesarean Section (n=434)	Vaginal Delivery (n=429)	p-value
Cord Blood Volume (ml)	59 ± 27	55 ± 23	0.11€
Cord Blood CD34+ (x10 ⁵)	30 ± 6	17.4 ± 2.4	0.08ŧ

Table 1: Umbilical cord blood data; *: Mann-Whitney U Test; t: Not statistically significant.

Conclusion

In conclusion, according to what was previously reported, there was no statistically significant difference between the quantity and the quality of blood collected during caesarean sections or vaginal deliveries. This has an important implication in the selection of the kind of delivery for the crucial collections performed when a pregnant woman has a child affected by a disease treatable with stem cells transplantation. These cells can be obtained from either the newborn foetal cord blood or his bone marrow, if he is HLA compatible. In such cases it is not necessary to perform a caesarean delivery to attempt to collect a higher blood volume and consequently a higher number of transplantable foetal haematopoeitic stem cells [1-11].

References

- 1. da Silva-Anoma S, Bertin KD, Ossenou O, Gaudens DA, Yao D, et al. (2001) [Prolapse of the urethral mucosa in young girls from the Ivory Coast]. Ann Urol (Paris) 35: 60-63.
- 2. Fiogbe MA, Hounnou GM, Koura A, Agossou-Voyeme KA (2011) Urethral mucosal prolapse in young girls: a report of nine cases in Cotonou. Afr J Paediatr Surg 8: 12-14.
- 3. Lang ME, Darwish A, Long AM (2005) Vaginal bleeding in the prepubertal child. CMAJ 172: 1289-1290.
- 4. Aprile A, Ranzato C, Rizzotto MR, Arseni A, Da Dalt L, et al. (2011) "Vaginal" bleeding in prepubertal age: a rare scaring riddle, a case of the urethral prolapse and review of the literature. Forensic Sci Int 210: e16-20.
- 5. Park BJ, Kim YW, Kim TE, Lee DH (2009) Urethral prolapse in a premenarchal Asian girl. Obstet Gynecol 113: 506-507.
- Falandry L (1994) [Prolapse of the urethra in black girls. Personal experience in 11 cases]. Med Trop (Mars) 54: 152-156.
- da Silva-Anoma S, Aguehounde C, Ouattara O, Dieth A, Keita A, et al. (1996) The urethral prolapse in girls: a rare condition in pediatric surgery. Our experience of 22 cases observed at the University Hospital of Cocody and Yopougon. Prog Urol 6: 392-397.

- 8. Rudin JE, Geldt VG, Alecseev EB (1997) Prolapse of urethral mucosa in white female children: experience with 58 cases. J Pediatr Surg 32: 423-425.
- 9. Sherry Boschert (2001) Topical Estrogen Soaks Heal Urethral Prolapse. OB/GYN News.
- 10. Agarwal S, Lall A, Bianchi A, Dickson A (2008) Uro-genital bleeding in pre-menarcheal girls: dilemmas of child abuse. Pediatr Surg Int 24: 745-746.
- 11. Essiet A, Ikpi E, Essiet GA, Nkposong EO (2007) Uretralprolapse: A case report and commentary on management. African Journal of Urology 13: 1.

Copyright: ©2016 Carlo Pafumi. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.