Prostaglandin-modulated umbilical cord blood hematopoietic stem cell transplantation

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Umbilical cord blood (UCB) is a valuable source of hematopoietic stem cells (HSCs) for use in allogeneic transplantation. Key advantages of UCB are rapid availability and less stringent requirements for HLA matching. However, UCB contains an inherently limited HSC count, which is associated with delayed time to engraftment, high graft failure rates, and early mortality. 16,16-Dimethyl prostaglandin E2 (dmPGE2) was previously identified to be a critical regulator of HSC homeostasis, and we hypothesized that brief ex vivo modulation of dmPGE2 could improve patient outcomes by increasing the “effective dose” of HSCs. Molecular profiling approaches were used to determine the optimal ex vivo modulation conditions (temperature, time, concentration, and media) for use in the clinical setting. A phase 1 trial was performed to evaluate the safety and therapeutic potential of ex vivo modulation of a single UCB unit using dmPGE2 before reduced-intensity, double UCB transplantation. Results from this study demonstrated clear safety with durable, multilineage engraftment of dmPGE2-treated UCB units. We observed encouraging trends in efficacy, with accelerated neutrophil recovery (17.5 vs 21 days, \( P = .045 \)), coupled with preferential, long-term engraftment of the dmPGE2-treated UCB unit in 10 of 12 treated participants. This study was registered at www.clinicaltrials.gov as #NCT00890500. (Blood. 2013;122(17):3074-3081)

Introduction

Umbilical cord blood (UCB) is a valuable source of hematopoietic stem cells (HSCs) for use in allogeneic transplantation when a suitable adult donor is unavailable.1,2 However, many of the publically accessible UCB units are currently unfit for use in adult transplantation because of a low HSC content, which is associated with delayed time to engraftment, high graft failure rates, and early mortality.3,4 Several approaches are being evaluated to address this limiting stem cell dose problem, including cotransplantation of multiple UCB units,5,6 cotransplantation with progenitor cells from an alternative source,7,8 direct intramarrow injection of UCB units,9 or expanding HSCs in culture.10-12 Although each strategy has its benefits, significant limitations are associated with these approaches, including induction of differentiation of long-term HSCs during extended culture, high manufacturing cost, and the risk of introducing infectious agents.

An alternative strategy to improve outcomes for patients undergoing UCB transplantation is to use a brief pulse treatment with a small molecule modulator to enhance the homing and engraftment potential of HSCs. One potential ex vivo-enhancing agent, the stable prostaglandin E2 (PGE2) derivative 16,16-dimethyl PGE2 (dmPGE2), was previously identified in a chemical screen using zebra fish embryos to be a potent regulator of vertebrate HSC homeostasis.13 dmPGE2 modulates HSCs primarily through the G-protein-coupled prostaglandin receptors PTGER2 (EP2) and PTGER4 (EP4), which use cyclic adenosine monophosphate (cAMP) as a secondary messenger to upregulate the expression of genes involved in homing (eg, CXCR4), proliferation (eg, cyclinD1), and cell survival (eg, survivin).14,15 dmPGE2 also increases Wnt signaling in HSCs and enhances self-renewal, which is important for the long-term maintenance of HSCs.16 An extensive set of characterization studies in zebra fish, murine, and nonhuman primate models demonstrated the ability of dmPGE2 to enhance hematopoietic engraftment of HSCs.13-15,17

On the basis of promising preclinical evidence and prior human safety data,18,19 a phase 1 clinical trial was initiated to evaluate the safety and efficacy of using an ex vivo treatment with dmPGE2.
to improve HSC engraftment following reduced-intensity double UCB transplantation. Double UCB transplantation provides a unique platform to examine the potential of enhanced hematopoietic engraftment of dmPGE2-modulated UCB compared with unmanipulated UCB by virtue of the competitive engraftment between UCB units. In addition, the second unmanipulated UCB unit provides a safety mechanism in the event of damage to the treated UCB unit during the ex vivo manipulation. Herein, we describe a point-of-care, ex vivo pulse treatment strategy designed to accelerate engraftment in UCB transplantation, describe the molecular studies that lead to the optimization of this ex vivo modulation process, and report outcomes treated on this phase 1 clinical trial.

### Materials and methods

#### Patients and treatment program

The primary objective of this phase 1 trial was to evaluate the safety of dmPGE2-treated UCB (dmPGE2-UCB) cotransplantation with an unmanipulated UCB unit in patients with hematologic malignancies. Secondary objectives were to determine the kinetics of engraftment and the fractional chimerism of dmPGE2-UCB after transplantation. Participants with hematologic malignancies for whom no HLA-matched donor was available received conditioning with fludarabine (180 mg/m²), melphalan (100 mg/m²), and antithymocyte globulin (4 mg/kg) and received graft-versus-host disease (GVHD) prophylaxis with sirolimus (target trough concentration, 3–12 ng/mL) and tacrolimus (target trough concentration 5–10 ng/mL), as described previously.25 UCB units were required to be ≥4/6 HLA-allele matched with the recipient and each other. Each UCB unit was required to be ≥1.5 × 10⁷ total nucleated cells (TNCs)/kg before cryopreservation, and the combined cell dose was required to be ≥3.7 × 10⁷ TNC/kg. UCB units were hierarchically selected from international cord blood banks based on TNC count, HLA match, and unit age. Units against which participants had preformed anti-HLA antibodies were excluded.

On the day of transplantation, 2 cryopreserved UCB units were thawed and resuspended in a saline solution (0.9% NaCl) containing 5% human serum albumin (Baxter or Talecris) and 8% Dextran 40 (Hospira) (LMD/HSA). A total of 2 cohorts of patients were enrolled. In cohort 1, one of the 2 UCB units was incubated with 10 µM of dmPGE2 (Fate Therapeutics) for 60 minutes at 4°C in LMD/HSA (9 patients). In the initial 6 patients, the smaller unit (by TNC before cryopreservation) was treated with dmPGE2, and in subsequent 3 patients, the larger unit was treated. Cohort 2 consisted of 12 patients in whom the larger of the 2 UCB units was incubated with 10 µM of dmPGE2 for 120 minutes at 37°C in LMD/HSA. After incubation, excess dmPGE2 was removed using a second centrifugation procedure and was resuspended in LMD/HSA for infusion. All patients received both UCB units within 4 hours of each other, with the larger UCB (whether dmPGE2-treated or not) always administered first. Standard posttransplantation care was delivered to all participants. In accordance with the Declaration of Helsinki, all participants provided informed consent to participate, and this trial was prospectively registered at www.clinicaltrials.gov (#NCT00890500). The study was approved by the Office for Protection of Research Subjects at the Dana-Farber/Harvard Cancer Center.

Patient baseline characteristics were reported descriptively. Neutrophil engraftment was defined as the first of 3 consecutive days with neutrophil recovery to at least 0.5 × 10⁹ cells/L. Platelet engraftment was defined as the first day of a platelet count of at least 20 × 10⁹ cells/L, without supporting transfusion in the prior 3 days. Donor chimerism was determined from peripheral blood mononuclear cells by analyses of informative short tandem repeat loci using the ABI Profiler-Plus Kit (Applied Biosystems) and the ABI 310 Genetic Analyzer. Overall survival (OS) was defined as the time from transplant to death from any cause, whereas progression-free survival (PFS) was defined as the time from transplant to malignant disease progression or death from any cause. Surviving patients were censored at their date of last known follow-up. OS and PFS estimates were calculated using the method of Kaplan and Meier.22 A comparative control cohort of 53 participants treated at our institution using the same UCB unit selection criteria, conditioning, and GVHD prophylaxis regimen was used for comparison (supplemental Table 1, available on the Blood Web site). All P values are based on 2-sided tests and were computed using SAS v9.2 (SAS Institute, Cary, NC).

#### Primary cells

Cryopreserved human UCB CD34⁺ primary cells were purchased from AliCells or Stem Cell Technologies (Vancouver, BC, Canada).

#### RNA preparation

RNA was extracted using the PicoPure RNA Isolation kit (Life Technologies) using the manufacturer’s recommended protocol. Total RNA was quantified using the Nanodrop 2000 Spectrophotometer (Thermo Scientific). RNA integrity was confirmed using an Agilent 2100 Bioanalyzer (Agilent Technologies).

#### Genome-wide microarray analysis

Biotinylated antisense RNA was prepared from 10 to 100 ng of total RNA involving the Message Amp II kit (Life Technologies) following the standard 2-round amplification protocols and hybridized to Affymetrix U133-plus-2.0 GeneChips according to the manufacturer’s instructions. Arrays were processed in the GeneChip Fluidics Station 450 and were scanned on the 3000 7G Scanner (Affymetrix). Probe intensities were normalized according to a log-scale robust multiarray analysis (RMA) method (Affymetrix) and were visualized in Spotfire for Genomics 4.5 (Tibco). Raw expression data files are available on Gene Expression Omnibus (GSE46569).

Parameterized paired t tests (Benjamini-Hochberg false discovery rate < .05, adjusted P value<q-value<.05, and fold change > or < threefold) detected probes with significant changes due specifically to the dmPGE2 treatment conditions. Biological pathway enrichment analysis of the upregulated probes was performed against the Gene Ontology (GO) database ( Singular Enrichment to GO Biological Process and false discovery rate < .01).

#### Microfluidic quantitative PCR using the Fluidigm platform

The messenger RNA sequences of the dmPGE2 signature genes were taken from the National Center for Biotechnology Information Gene database, and amplification primers were designed using Primer3. (supplemental Table 2) For real-time polymerase chain reaction (PCR) transcript quantitation, we used the BioMark Dynamic Array microfluidics system and GE Dynamic Array 96.96 chips (Fluidigm) using the manufacturer’s protocol. Amplification results were analyzed using BioMark Real-Time PCR Analysis software. Samples with Cycle Thresholds above 28 or amplified products with inappropriate melting curve properties were excluded from the calculations. Log₂ fold change results are displayed in Spotfire for Genomics 4.0 in a heat-map format.

#### Human CD34⁺ HSC murine homing

Human CD34⁺ cells isolated from UCB were treated with 10 µM of dmPGE2 or vehicle (dimethylsulfoxide) for 2 hours at 37°C in StemSpan-SFEM (Stem Cell Technologies) or a minimal media containing LMD/HSA. For the last 15 minutes of the incubation, Vybrant Dye Dil (Life Technologies) was added to label the cells. After incubation, 1 × 10⁶ cells from each condition were washed and injected in sublethally irradiated NOD.Cg-Prkdcscid Il2rgtm1Wjl/SzJ (NSG) mice in groups of 5 per condition. At 20 hours after the injections, hind limb long bones were harvested and the bone marrow (BM) extracted. Flow cytometry was performed on the BM using LSRII (Becton Dickinson) and gating criteria based on remaining DiI-stained CD34⁺ cells and control mouse BM. This homing experiment was performed in duplicate using CD34⁺ cells from 2 separate UCB units. Data were plotted as the increase in homed events relative to the vehicle-treated cells. Results were averaged for both donor groups in each of the conditions tested. All protocols using animals in this study were approved by the Institutional Animal Care and Use Committee at Indiana University School of Medicine (supplemental Methods).
Table 1. Patient and UCB characteristics

<table>
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<tr>
<th>Patient characteristics</th>
<th>Cohort 1</th>
<th>Cohort 2</th>
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<tbody>
<tr>
<td>Sample size (n)</td>
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<td>12</td>
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<tr>
<td>Median age, y (range)</td>
<td>43.0 (29-64)</td>
<td>57.5 (19-66)</td>
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<td>4 (33.3)</td>
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<td>UCBB unit characteristics</td>
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<td>dmPGE2 processing</td>
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<td>−7.87%</td>
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<td>CD34+ viability (7-AAD)</td>
<td>−1.40%</td>
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ALL, acute lymphoblastic leukemia; AML, acute myeloblastic leukemia; CMV, cytomegalovirus; GM, granulocyte-macrophages (in CFUs); MDS, myelodysplastic syndrome; NHL/CLL, non-Hodgkin lymphoma/chronic lymphocytic leukemia.

Results

Cohort 1

Nine participants with high risk hematological malignancies were treated using ex vivo modulation conditions based on preclinical studies (10 μM of dmPGE2, exposure for 60 minutes at 4°C in LMD/HSA media) between May 2009 and June 2010. Patient and UCB unit characteristics are described in Table 1. No safety concerns were identified in this cohort; however, neither an improvement in rates of neutrophil recovery nor preferential engraftment of the dmPGE2-UCB over the unmanipulated UCB units was observed in this cohort; however, neither an improvement in rates of neutrophil recovery nor preferential engraftment of the dmPGE2-UCB over the unmanipulated UCB units was observed (Figure 1A). The median times to neutrophil and platelet engraftment were 24 and 72.5 days, respectively. Two of 7 patients undergoing engraftment demonstrated prolonged hematopoiesis from the dmPGE2-UCB units, and 2 patients had primary or late graft failure (Figure 1B-C).

Optimization of the ex vivo modulation

Given the lack of accelerated engraftment in the initial cohort of patients, we sought to determine if the ex vivo incubation conditions were optimal in the clinical setting. To evaluate the effects of temperature, CD34+ cells were incubated at 4°C, 25°C, or 37°C for 2 hours in the presence of dmPGE2 or vehicle. After the incubation, genome-wide expression arrays were used to identify genes upregulated by dmPGE2. Only 2 probe sets across the entire human transcriptome were significantly upregulated (>fourfold) when the CD34+ cells were incubated at 4°C (Figure 2A). Performing the ex vivo incubation at 25°C also resulted in a modest biological response, with 19 probe sets being significantly upregulated. In contrast, increasing the incubation temperature to 37°C resulted in a robust biological response with 192 probe sets being significantly upregulated, suggesting that physiological temperatures are required for cryopreserved human CD34+ cells to effectively activate the prostaglandin signaling pathway. The gene ontology pathway analysis for the dmPGE2 signature showing genes involved in proliferation, migration, and receptor signaling is shown in Figure 2B.

To determine the optimal incubation time, CD34+ cells were treated with 10 μM of dmPGE2 or vehicle (dimethylsulfoxide) for increasing amounts of time at 37°C. After the incubation, dmPGE2-induced changes in gene expression were measured using a high-throughput microfluidic quantitative PCR-based system. Amplification primers were designed for 6 reference genes and 90 of the top dmPGE2 upregulated genes, selected based on the genome-wide expression analysis and biological significance for homing, proliferation, and survival (supplemental Table 1). Figure 3A shows that 2-hour incubation at 37°C is required to induce the maximal activation of the prostaglandin pathway. We also demonstrated that the prostaglandin must be present during the entire 2-hour incubation to achieve a full biological response (supplemental Figure 1). This eliminates the possibility of using a brief pulse treatment (<60 minutes) to initiate a signaling event that could mature once the enhanced HSCs are infused into the patient.

To determine the optimal concentration of dmPGE2 for ex vivo modulation, cryopreserved human CD34+ cells were treated with increasing concentrations of dmPGE2 (0.1 μM, 1 μM, 10 μM, 50 μM, and 100 μM) for 2 hours at 37°C. Supraphysiological concentrations of dmPGE2 are permissible because the molecule is removed with the supernatant during washing prior to infusion. Expression changes in the prostaglandin signature genes were...
monitored using a microfluidic quantitative reverse transcription (RT)-PCR platform. The amount able to induce a maximal pathway response was $10^{-6}$ M (Figure 3B). Higher concentrations of dmPGE2 were not able to increase pathway activation at 4°C (supplemental Figure 2).

The majority of dmPGE2 preclinical homing, engraftment, and optimization studies were performed in “complete” media such as StemSpan-SFEM (Stem Cell Technologies), which contains glucose, amino acids, and other nutrients designed to support long-term culturing of HSCs. Unfortunately, this type of complete media cannot be used in a clinical setting because differentiated cell types (eg, granulocytes) that do not survive the cryopreservation process lyse upon thawing and incubation in this media, which causes a significant reduction in the TNC counts (supplemental Figure 3).

Typically, the media used in a clinical setting for thawing and washing cryopreserved UCB contain LMD/HSA to minimize cell loss. We sought to determine whether the ex vivo modulation could be performed in these clinically compatible, nutrient-free media. To characterize the effects of media on the level of pathway activation, CD34+ cells were treated with $10^{-6}$ M of dmPGE2 for

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**Figure 1. Cohort 1 clinical results.** (A) Donor chimerism of the dmPGE2-treated UCB unit during 360 days posttransplantation (percentage of total) as determined from peripheral blood mononuclear cells by analyses of informative short tandem repeat loci. (B) Cumulative incidence of neutrophil engraftment of $>500$ cells/mm$^3$, in cohort 1 ($n = 9$, blue) compared with historical institutional control participants ($n = 53$, black). (C) Cumulative incidence of platelet engraftment in cohort 1 ($n = 9$, blue) compared with historical institutional control participants ($n = 53$, black).

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**Figure 2. Evaluation of ex vivo modulation temperature with cryopreserved human UCB CD34+ cells with dmPGE2.** (A) Genome-wide expression analysis on Affymetrix U133 plus 2.0 GeneChips of human UCB CD34+ cells treated with $10^{-6}$ M of dmPGE2 for 2 hours at 4°C, 25°C, and 37°C. RMA log$_2$ normalized expression levels for cells treated with dmPGE2 (y-axis) in comparison with vehicle-treated cells (x-axis). There were 2, 19, and 192 probe sets (red) with expression levels changing greater (or less) than fourfold due to dmPGE2 treatment at 4°C (left), 25°C (center), and 37°C (right), respectively. (B) GO enrichment analysis showing biological processes enriched in the upregulated probes induced by treatment with dmPGE2 at 37°C. Genes upregulated by dmPGE2 treatment at 37°C in CD34+ are listed for each GO category.
2 hours at 37°C in either StemSpan-SFEM or LMD/HSA. After the incubations, prostaglandin-induced expression changes were analyzed using genome-wide expression arrays. Figure 3C demonstrates that both media formulations were able to support activation of the prostaglandin pathway by dmPGE2; however, threefold more probe sets were upregulated in StemSpan-SFEM (297 vs 99 probe sets). We used a mouse homing model to determine whether the incubation media affect the functional properties of the enhanced HSCs. The homing properties of human CD34^+ cells were significantly improved in both types of media with a 2.2-fold increase in the number of human cells that had migrated to the BM of mice when the CD34^+ cells were pulse treated in StemSpan-SFEM (P < .001) compared with a 1.6-fold increase with LMD/HSA (P = .002) (Figure 3D). The difference in homing properties between the 2 types of media (1.6-fold vs 2.2-fold) is also statistically significant (P = .03). A subtle decrease in viable cell recovery (7-aminoactinomycin D [7-AAD] and colony-forming units [CFU] in culture) was observed in the LMD/HSA relative to nutrient-rich media. On the basis of these results, we decided to move forward with the clinically proven LMD/HSA media for the initial clinical studies.

To characterize the engraftment properties of HSCs treated using the optimized conditions, lethally irradiated mice were injected with decreasing numbers of mouse BM cells (200K, 100K, 50K, or 10K) treated with 10 μM of dmPGE2 or vehicle for 2 hours at 37°C. Results from this experiment demonstrate that ex vivo treatment with dmPGE2 significantly increased the survival rates of mice transplanted with limiting numbers of HSCs (supplemental Figure 4). We also observed accelerated recovery of neutrophils and platelets using this murine transplantation model (supplemental Figure 5A-B).

Cohort 2
A total of 12 additional participants were accrued between August 2010 and August 2011 and were treated according to the optimized ex vivo dmPGE2 modulation protocol (10 μM of dmPGE2 and 2-hour exposure at 37°C in LMD/HSA media). Clinical and UCB...
Discussion

The development of HSCs during embryogenesis is regulated by pathways that regulate HSC homing and engraftment after transplantation. In adulthood, dmPGE2 was originally identified as a small molecule that could increase HSC formation in the developing aorta of the zebra fish embryo, and enhanced HSC engraftment in murine HSC transplantation after brief ex vivo exposure. Further studies established that dmPGE2 improved human UCB engraftment in immunodeficient mice, and demonstrated safety in long-term primate transplantation studies. Given that the limiting number of HSCs in UCB units results in delayed hematopoietic engraftment, graft failure, and incomplete immunologic reconstitution, we sought to determine if modulation of UCB with dmPGE2 before transplantation could improve these outcomes. Of note, our study is the first in which the function of a small molecule discovered in the zebra fish system has reached a human clinical trial. UCB units from the initial cohort of 9 participants were treated under ex vivo modulation conditions designed to maximize viability of the HSCs. After demonstrating safety, but lack of clear efficacy, we enrolled a second cohort of patients using ex vivo modulation conditions designed to increase the prostaglandin pathway stimulation. Results from this second group of patients demonstrated a correlation between the enhanced biologic activity, accelerated engraftment, and preferential hematopoiesis from the modulated unit in a double UCB transplantation model.

In translating this academic discovery into the clinic, one of the challenges that we faced was determining the optimal incubation conditions to use in the clinical setting. The incubation conditions used for the preclinical studies varied significantly with respect to the temperature, duration, dose, and media tested. Molecular profiling approaches were used to identify a set of ex vivo modulation conditions, which enable rapid and robust activation of the prostaglandin pathway in cryopreserved human cord blood HSCs. The key changes to the incubation protocol were increasing the temperature from 4°C to 37°C and extending the incubation from 1 hour to 2 hours. In making these changes, we had to consider the potential impacts of the optimized conditions on cell viability, especially in the clinically established nutrient-free media. Before human use, we confirmed that these optimized conditions were not only safe but resulted in enhanced engraftment and hematopoiesis in murine models.

There were limitations to the conclusions that can be made from this phase 1 clinical trial. In the second cohort of patients, it was always the larger unit before cryopreservation that was modulated with dmPGE2, and it was this unit that dominated hematopoiesis in 10 of 12 treated patients. Although firm conclusions cannot be drawn based on the small sample sizes tested, 2 important factors should be noted. First, in the latter patients enrolled in cohort 1, the
same protocol of treating and infusing the larger unit first was used as in cohort 2, and no signal hinting at early engraftment or chimerism dominance was noted; this factor led to the early closure of the trial and the redesign of the ex vivo expansion process. In addition, in cohort 2, although the larger unit (based on TNC before cryopreservation) was to be infused first, the UCB units, in fact, were extraordinarily comparable with respect to TNC (1.8 vs 1.7 × 10⁷ TNCs/kg, P = .43), CD34⁺ cells (0.74 vs 0.56 × 10⁷ CD34⁺/kg, P = .71), and CFU growth (4.5 vs 6.9, P = .77) (Table 1). In dual UCB transplantation, 1 unit dominates in the vast majority of patients on the basis of incompletely understood biological factors. Factors such as TNC dose, CD34⁺ cell dose, CD³⁺ T-cell dose, HLA match, CXCR4 expression, and CFU assay have been suggested as relevant contributors to this phenomenon. To determine if order of infusion or cell size was relevant in the outcomes we reported in this manuscript, we compared the UCB dominance patterns with those in the historical control cohort, and noted that UCB dominance was nearly random in the control group, with ~50% of participants undergoing engraftment with each of the transplanted units, using varying thresholds for the determination of UCB dominance. The dominance seen with the dmPGE₂-treated patients in the active cohort suggests that enhanced homing may be relevant to this process. We have previously demonstrated that CXCR4 expression is enhanced on human UCB cells treated with dmPGE₂. As a chemokine receptor to SDF1, increased CXCR4 expression may enhance engraftment. We also demonstrated that dmPGE₂ functions as a modulator pathway for Wnt signaling via cAMP, which could lead to enhanced self-renewal of the HSCs. Rather than increasing the number of HSCs, this pharmacologic ex vivo modulation increases the “effective stem cell dose” for use in transplantation. Many additional cell types (eg, regulatory T cells, CD4⁺ T cells, CD8⁺ T cells, monocytes, dendritic cells, and natural killer cells) in UCB units express the EP2/4 receptors and have been shown to be affected by dmPGE₂. It is likely that effects on differentiated cell types in cord blood are involved in stimulating engraftment and driving hematopoiesis. We are currently in the process of characterizing the molecular responses across these cell types and the potential mechanism(s) by which a pulse treatment with dmPGE₂ may affect UCB unit dominance, GVHD, rates of infections, and rates of viral reactivation.

Other groups have attempted to expand HSCs to improve engraftment and clinical outcomes after UCB transplantation. A recent trial established that the notch ligand, Delta1, could be used to expand UCB in culture for 14 days, and these cells could be infused with a second UCB unit. Although the expanded UCB unit did engraft early, these cells did not contribute to long-term hematopoiesis and likely acquired a committed myeloid progenitor phenotype during ex vivo culture. Similarly, a mesenchymal stem cell coculture UCB expansion trial demonstrated a 40-fold expansion of CD34⁺ cells; however, transplantation of these cells with a second unexpanded UCB unit similarly resulted in the loss of the expanded cells with time. We have demonstrated sustained multilineage dmPGE₂-UCB–derived hematopoiesis for 27 months after transplantation, but further optimization of these and other ex vivo protocols may prove that each approach has usefulness in the transplantation setting.

Despite the limited conclusions that can be drawn from the clinical data, the potential implications of this prospective competitive engraftment clinical trial are broad. First, by generating more effective HSCs, the minimal nucleated cell dose required for adult UCB transplantation might be lowered, increasing the number of acceptable UCB units currently available in public UCB banks. Currently, fewer than 5% of UCB units in the National Marrow Donor Program inventory are of adequate size for use in single UCB transplantation for average-weight American adults. By increasing the effective stem cell dose by as little as fourfold, 98% of these units would be accessible for this use (Michael Boo, J.D., National Marrow Donor Program, email communication with D. Shoemaker). In addition, by increasing the number of accessible units, the likelihood of identifying better HLA-matched units would be increased, potentially leading to improved UCB transplantation outcomes. From a resource point of view, reducing the need for costly second UCB units and reducing the length and complexity of hospital stays represent a substantial potential cost savings for health care payers. In addition, unlike the significant time requirements associated with HSC expansion approaches, this simple ex vivo manipulation procedure with dmPGE₂ is inherently exportable to all stem cell–processing facilities.

In summary, in this preliminary human experience, we have demonstrated that ex vivo modulation of UCB stem cells by using dmPGE₂ results in enhanced and more rapid engraftment in human UCB transplantation. On the basis of these positive results, we are now expanding the use of dmPGE₂ in randomized phase 2 trials, in addition to initiating new phase 1 studies in single UCB transplantation and autologous peripheral blood stem cell transplantation.

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Prostaglandin-modulated umbilical cord blood hematopoietic stem cell transplantation