

HEMOSTATIC FUNCTION OF APHERESIS PLATELETS STORED AT 4°C AND 22°C

Kristin M. Reddoch,* Heather F. Pidcoke,[†] Robbie K. Montgomery,[†]
Chriselda G. Fedyk,[†] James K. Aden,[†] Anand K. Ramasubramanian,*
and Andrew P. Cap[†]

*Department of Biomedical Engineering, The University of Texas at San Antonio, San Antonio, Texas;
and [†]US Army Institute of Surgical Research, Fort Sam Houston, Texas

Received 17 Sep 2013; first review completed 03 Oct 2013; accepted in final form 23 Oct 2013

ABSTRACT Introduction: Platelet refrigeration decreases the risk of bacterial contamination and may preserve function better than standard-of-care room temperature (RT) storage. Benefits could include lower transfusion-related complications, decreased costs, improved hemostasis in acutely bleeding patients, and extended shelf life. In this study, we compared the effects of 22°C and 4°C storage on the functional and activation status of apheresis platelets. **Methods:** Apheresis platelets (n = 5 per group) were stored for 5 days at 22°C with agitation (RT) versus at 4°C with agitation (4°C + AG) and without (4°C). Measurements included platelet counts, mean platelet volume, blood gas analytes, aggregation response, thromboelastography, thromboxane B₂ and soluble CD40 ligand release, activation markers, and microparticle formation. **Results:** Sample pH levels were within acceptable limits for storage products (pH 6.2–7.4). Platelet glucose metabolism ($P < 0.05$), aggregation response (adenosine diphosphate: RT 0; 4°C + AG 5.0 ± 0.8 ; 4°C 5.6 ± 0.9 ; $P < 0.05$), and clot strength (maximum amplitude: RT 58 ± 2 ; 4°C + AG 63 ± 2 ; 4°C 67 ± 2 ; $P < 0.05$) were better preserved at 4°C compared with RT storage. Refrigerated samples were more activated compared with RT ($P < 0.05$), although thromboxane B₂ ($P < 0.05$) and soluble CD40 ligand release ($P < 0.05$) were higher at RT. Agitation did not improve the quality of 4°C-stored samples. **Conclusions:** Apheresis platelets stored at 4°C maintain more viable metabolic characteristics, are hemostatically more effective, and release fewer proinflammatory mediators than apheresis platelets stored at RT over 5 days. Given the superior bacteriologic safety of refrigerated products, these data suggest that cold-stored platelets may improve outcomes for acutely bleeding patients.

KEYWORDS Hemostasis, coagulation, aggregation, clot strength, cold storage, activation

ABBREVIATIONS RT room temperature; 4°C storage at 4°C; 4°C + AG storage at 4°C with gentle agitation; AP apheresis platelets; ADP adenosine diphosphate; TRAP thrombin receptor activating peptide; CD62P P-selectin; CD40L CD40 ligand; sCD40L soluble CD40 ligand; CD154 CD40L surface expression molecule; CD42b glycoprotein Ib receptor; TxB₂ thromboxane B₂; RBC red blood cell; MPV mean platelet volume; MPC mean platelet component; TEG Thromboelastography; AUC area under the curve; *R* reaction time (time to initial fibrin formation); *K* time to clot formation; α angle rate of clot formation; MA maximum amplitude (clot strength)

INTRODUCTION

Current blood-banking recommendations are that platelets be stored in incubators at 22°C, with gentle agitation for no longer than 5 days (1). This limited shelf life is necessary because of the risk of bacterial contamination that leads to life-threatening transfusion-related infections (2–4). These storage practices also result in a platelet storage lesion that is associated with a decline in platelet hemostatic function (5).

Cold storage at 4°C could prolong shelf life by diminishing the risk of bacterial sepsis, decreasing platelet metabolism, and

maintaining functionality. Transportation of 4°C platelets would be convenient because the infrastructure for other refrigerated blood products, such as red blood cells (RBCs), is already in place (6). Cold platelets may be a better hemostatic product. Several *in vitro* studies from the 1970s showed that refrigeration of platelets results in better metabolic and functional responses such as minimal lactate accumulation, better aggregation response, and adhesion to subendothelium (7–9). *In vivo* human studies have shown that 4°C platelets function better than room temperature (RT) platelets in reducing the bleeding times in thrombocytopenic patients, aspirin-treated volunteers, and in aplastic thrombocytopenic patients shortly after transfusion (10–12).

Despite promising *in vitro* and *in vivo* studies in controlling acute bleeding, the practice of cold storage for transfusion was abandoned during the 1970s because of the belief that clinically effective platelets should be both hemostatically functional and survive in circulation for several days as indicated for prophylactic transfusion. It was shown by Murphy and Gardner (13) that platelets stored for up to 18 h in cold (2°C–4°C) show decreased recovery and survival upon transfusion compared with their RT (22°C–24°C) counterparts; i.e., the life span ($t_{1/2}$) of cold- and RT-stored platelets is 1.3 and 3.9 days, respectively. Consequently, when transfused to thrombocytopenic patients, cold-stored platelets are as effective as

Address reprint requests to Anand K. Ramasubramanian, PhD, Department of Biomedical Engineering, University of Texas at San Antonio, San Antonio, TX 78256. E-mail: anand.ramasubramanian@utsa.edu; or Andrew P. Cap, MD, PhD, Coagulation and Blood Research, US Army Institute of Surgical Research, 3650 Chambers Pass, JBSA Fort Sam Houston, Texas 78234. E-mail: andrew.p.cap.mil@mail.mil.

This study was conducted under a protocol reviewed and approved by the US Army Medical Research and Materiel Command Institutional Review Board and in accordance with the approved protocol.

Support was received from the US Army Medical Research and Materiel Command and National Institutes of Health (NIGMS MBRS-RISE GM060655)

The authors declare that they have no conflicts of interest relevant to this article.

The opinions or assertions contained herein are the private views of the author and are not to be construed as official or as reflecting the views of the Department of the Army or the Department of Defense.

DOI: 10.1097/SHK.0000000000000082

Copyright © 2014 by the Shock Society

Report Documentation Page

Form Approved
OMB No. 0704-0188

Public reporting burden for the collection of information is estimated to average 1 hour per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, to Washington Headquarters Services, Directorate for Information Operations and Reports, 1215 Jefferson Davis Highway, Suite 1204, Arlington VA 22202-4302. Respondents should be aware that notwithstanding any other provision of law, no person shall be subject to a penalty for failing to comply with a collection of information if it does not display a currently valid OMB control number.

1. REPORT DATE 01 MAY 2014		2. REPORT TYPE N/A		3. DATES COVERED -	
4. TITLE AND SUBTITLE Hemostatic function of apheresis platelets stored at 4 degrees c and 22 degrees c				5a. CONTRACT NUMBER	
				5b. GRANT NUMBER	
				5c. PROGRAM ELEMENT NUMBER	
6. AUTHOR(S) Reddoch K. M., Pidcoke H. F., Montgomery R. K., Fedyk C. G., Aden J. K., Ramasubramanian A. K., Cap A. P.,				5d. PROJECT NUMBER	
				5e. TASK NUMBER	
				5f. WORK UNIT NUMBER	
7. PERFORMING ORGANIZATION NAME(S) AND ADDRESS(ES) United States Army Institute of Surgical Research, JBSA Fort Sam Houston, TX				8. PERFORMING ORGANIZATION REPORT NUMBER	
9. SPONSORING/MONITORING AGENCY NAME(S) AND ADDRESS(ES)				10. SPONSOR/MONITOR'S ACRONYM(S)	
				11. SPONSOR/MONITOR'S REPORT NUMBER(S)	
12. DISTRIBUTION/AVAILABILITY STATEMENT Approved for public release, distribution unlimited					
13. SUPPLEMENTARY NOTES					
14. ABSTRACT					
15. SUBJECT TERMS					
16. SECURITY CLASSIFICATION OF:			17. LIMITATION OF ABSTRACT UU	18. NUMBER OF PAGES 8	19a. NAME OF RESPONSIBLE PERSON
a. REPORT unclassified	b. ABSTRACT unclassified	c. THIS PAGE unclassified			

RT-stored platelets at 24 h after transfusion, although not at 72 h, suggesting cold storage may be acceptable for therapeutic transfusion (i.e., therapy for active hemorrhage) (14). In recent years, Hoffmeister and colleagues (15–17) have shown that platelet clearance and function may be entirely separate attributes and that appropriate modification (such as glycosylation) may prolong platelet survival in circulation. Concurrent reports from several groups have confirmed that platelets stored at 4°C are “healthier” than platelets stored at RT (18). Together, these studies have prompted the reconsideration of cold-stored platelets as a transfusion product to treat bleeding patients (18–20). We have recently shown that cold storage of platelet-rich plasma (PRP) enhances platelet activation and aggregation under shear, and cold storage of platelets in whole blood improves their performance in a panel of functional assays compared with RT-stored platelets (21, 22).

In this study, we characterized the effects of storage at both 22°C and 4°C for up to 5 days with apheresis platelets (APs) using functional and activation assays. It is particularly important to evaluate the effect of storage on APs because apheresis is the most common method of platelet isolation with 80% use in the United States and 50% in Europe, and it has been shown that the method of preparation of platelet concentrate (i.e., buffy coat, apheresis, or PRP) is a determinant of platelet storage lesion (23–27). We hypothesized that apheresis platelets stored at 4°C would demonstrate more viable metabolic characteristics, perform better in functional tests, form stronger clots, and release fewer inflammatory mediators compared with platelets stored at 22°C.

MATERIALS AND METHODS

Reagents and suppliers

Calcium and kaolin used in thromboelastography (TEG) were obtained from Haemonetics Corp (Braintree, MA). Multiplate agonists were obtained from Diapharma (West Chester, OH). Fluorescein isothiocyanate conjugated bovine lactadherin was purchased from Haematologic Technologies, Inc (Essex Junction, VT). Gibco 1× Hanks balanced salt solution without calcium and magnesium was obtained from Invitrogen Life Technologies (Carlsbad, CA). The following anti human monoclonal mouse antibodies were purchased from BD Biosciences (San Jose, CA): allophycocyanin conjugated anti CD62P (P selectin), clone AK 4; Phycoerythrin conjugated anti CD 154 (CD40 ligand or CD40L), clone TRAP1 (thrombin receptor activating peptide 1). Peridinin Chlorophyll Protein conjugated mouse monoclonal anti CD42b (glycoprotein Ib receptor, clone HIP 1) was purchased from BioLegend (San Diego, CA). For absolute particle counts, TruCount tubes from BD Biosciences were utilized. Thromboxane B₂ (TxB₂) enzyme immunoassay kits were purchased from Cayman Chemicals (Ann Arbor, MI), and human soluble CD40L (sCD40L) extra sensitive platinum enzyme linked immunosorbent assay kits were purchased from eBioscience (Vienna, Austria). CG4⁺ and CHEM8⁺ cartridges were purchased from Abbott Laboratories (Abbott Park, IL).

Platelet storage and handling

Single AP units were collected in acid citrate dextrose plasma from healthy donors (n = 5) using a Trima Accel Automated Blood Collection System (Terumo BCT, Lakewood, CO) under a protocol reviewed and approved by the US Army Medical Research and Materiel Command Institutional Review Board and in accordance with the approved protocol. Ten milliliter aliquots were obtained from the donor bag and sterilely transferred into 15 mL BCSI mini bags (Seattle, WA) for storage, with one minibag used for each time point and condition. The minibags were stored for 5 days in one of the following conditions: RT in a Food and Drug Administration approved platelet incubator with agitation (RT); in a walk in refrigerator at 4°C with agitation (4°C + AG), or without agitation (4°C). Platelet function, characteristics, and release markers were analyzed for all stored samples on the day of collection (baseline or day 1), day 3, and day 5. Samples stored at 4°C were allowed to come to RT for 30 min and gently massaged before testing commenced. Apheresis platelets were

centrifuged for 10 min at 3,000g to obtain platelet poor plasma where applicable. For RBC reconstitution, whole blood was collected in acid citrate dextrose vacutainer tubes, spun at 3,500g for 10 min, and the plasma and buffy coat layers were carefully aspirated from the RBCs.

Laboratory equipment and measurements All tests were performed at the US Army Institute of Surgical Research at Joint Base San Antonio/Fort Sam Houston according to routine protocols. Platelet count and mean platelet volume (MPV) were determined in duplicate using a standard cell counter (Coulter Ac T diff2 Hematology System; Beckman Coulter, Brea, CA). Mean platelet component (MPC) was determined using an Advia 120 Haematology System (Bayer Diagnostics, Tarrytown, NY). Blood pH, sodium (in mM), potassium (in mM), chloride (in mM), glucose (in mg/dL), Pco₂ (in mmHg), Po₂ (in mmHg), bicarbonate (in mM), and lactate (in mM) were determined using an i STAT point of care device (Abbott Laboratories). Thromboelastography was performed using a TEG5000 device (Haemoscope Corp, Niles, IL) with kaolin activation. Aggregation was conducted using a multiplate impedance aggregometer (Verum Diagnostica, Munich, Germany) with adenosine diphosphate (ADP), collagen, and TRAP as agonists. Platelet markers were assessed by flow cytometry with BD FACSanto I, and microparticle counts were determined using a BD FACSanto II, RUO flow cytometer equipped with a FSC PMT capable of detecting particles down to 200 nm. Particles were considered microparticles if their sizes fell within the range of 200 to 1,000 nm.

Blood gas analysis Blood gas and chemistry analyses in fresh and stored APs were performed using CG4⁺ and CG8⁺ cartridges, respectively. Platelet poor plasma samples (100 µL/cartridge) were obtained to test for metabolic parameters.

Thromboelastography All samples were tested as previously described (28). Briefly, samples were recalcified by the addition of CaCl₂ (0.2 M, 30 µL) to a TEG sample cup followed by kaolin treated APs (330 µL). All tests were performed at 37°C in duplicate.

Impedance aggregometry Impedance aggregometry was performed according to established methods (28). Briefly, 300 µL of APs diluted with platelet poor plasma to a concentration of 250 × 10³ platelets/µL was incubated with 300 µL of 3 mM NaCl₂/CaCl₂ solution (Verum Diagnostica). After incubation, 20 µL of agonist (6.5 µM ADP, 3.2 µg/mL collagen, or 32 µM TRAP) was added. The tests were carried out for 6 min in duplicate, and area under the curve (AUC) was reported. Aggregation was also performed in the presence of RBCs with a platelet count of 300 × 10³/µL and 40% hematocrit for 12 min.

Surface receptor expression Geometric mean fluorescence intensity and abundance (% positive) were determined for the following platelet markers: CD42b (glycoprotein Ib receptor), CD62P (P selectin), lactadherin (phosphatidylserine [PS] surface expression), and CD154 (CD40L surface expression molecule). Ten thousand events per sample were recorded for each marker, and forward and side scatter characteristics were used to identify the platelet population. For micro particle analysis, absolute particle counts were determined using TruCount tubes (BD Biosciences).

Enzyme linked immunosorbent assay Commercially available kits were used to assess sCD40L and TxB₂ levels released into plasma during storage. Platelet poor plasma was stored at 80°C until analysis by enzyme linked immunosorbent assay. All plasma samples were diluted before testing to ensure that the concentrations fell within the standard range prescribed by the manufacturer.

Statistical analysis The data collected were analyzed by one way analysis of variance (ANOVA) for repeated measures with a post hoc Bonferroni adjustment for pairwise comparisons. Significance from baseline (day 1) and between groups was determined when *P* < 0.05. Data in tables are mean ± SD, and data in graphs are mean ± SEM, unless otherwise stated. Microsoft Excel (Microsoft Corp, Redmond, WA) was used to manage data, and analysis was performed using the statistical software JMP (version 10; SAS Institute Inc, Cary, NC).

RESULTS

Platelet Count, MPV, and MPC

We obtained platelet counts for all AP samples to assess loss of single platelets during storage in minibags as described in Materials and Methods (Table 1A). We observed a significant drop in the platelet count in both 4°C + AG and 4°C samples by day 5 (*P* < 0.05), along with macroscopic platelet clumping, but the platelet number was unchanged in RT samples. The decrease in platelet number in 4°C samples on day 5 compared with fresh platelets (day 1 or baseline) was ~20%; however, platelet counts were stable, and the clumping effect was

TABLE 1. Platelet counts $\times 10^3$

A. Platelet count $\times 10^3$, MPV, and MPC during storage in minibags (research use only)	Baseline		RT		4°C		4°C + AG	
	Day 1	Day 3	Day 5	Day 3	Day 5	Day 3	Day 5	
	Count, $\times 10^3$ platelets/ μ L	1,084 \pm 47	1,019 \pm 66	924 \pm 149	815 \pm 152	727 \pm 153*	854 \pm 103	687 \pm 147*
MPV, fl	7.0 \pm 0.2	6.9 \pm 0.1	7.0 \pm 0.2	6.9 \pm 0.4	6.9 \pm 0.3	7.0 \pm 0.4	6.8 \pm 0.3	
MPC, g/dL	25 \pm 1.8		20 \pm 1.7*		26 \pm 2.3			

*Values (mean \pm SD) significant from baseline ($P < 0.05$).

B. Platelet count $\times 10^3$ of apheresis platelets stored in full-size Food and Drug Administration approved platelet storage bags (n = 3)	Baseline		RT		4°C	
	Day 1	Day 3	Day 5	Day 5	Day 5	
	Count, $\times 10^3$ platelets/ μ L	1,192 \pm 445		1,187 \pm 51		1,123 \pm 40

*Values (mean \pm SD) were not significantly different ($P \geq 0.05$).

abolished when refrigerated platelets were stored in full-size bags (Table 1B). Mean platelet volume remained unchanged in both RT and 4°C samples suggesting no significant change in the size of single platelets. The MPC decreased significantly after 5-day RT storage ($P < 0.05$), suggesting platelet degranulation due to RT storage. Platelets stored at 4°C maintained MPC. Stored platelet samples were visually inspected for swirling according to the guidelines set by Bertolini and Murphy (29). Room temperature platelets maintained their ability to swirl at day 5 of storage, whereas swirling in 4°C-stored samples was nonexistent, most likely due to the shape change that occurs during storage at 4°C (30).

Platelet metabolism We measured the pH and blood gases in stored platelet samples using i-STAT, and the results are tabulated in Table 2. All samples maintained acceptable levels of pH, i.e., between 6.2 and 7.4 (31). We observed an increase in lactate production and glucose consumption in both RT- and 4°C-stored platelets. The lactate levels in RT-stored platelets increased by $\sim 300\%$ and $\sim 600\%$, and glucose levels dropped by 8% and 25% by days 3 and 5, respectively ($P < 0.05$). In comparison, the lactate levels in 4°C-stored platelets increased only by 150% and 225% by days 3 and 5, respectively ($P < 0.05$), and the glucose levels did not change significantly. Correspondingly, more bicarbonate was consumed and carbon

dioxide released in RT- than 4°C-stored platelets ($P < 0.05$). The dissolved oxygen levels were comparable and remained high, suggesting adequate gas exchange during the storage period. The glucose consumption and lactate production rates were 3- and 2-fold higher, respectively, during RT storage than during cold storage over 5 days (Table 3). There was no difference between 4°C platelets stored under nonagitated or agitated conditions. Thus, these parameters suggest that 4°C-stored platelets show a significant decrease (more than 2- to 3-fold) in metabolic rate compared with RT-stored platelets over a 5-day storage. A small but significant difference in sodium and chloride levels was observed at day 5 in RT samples and in potassium levels in 4°C and 4°C + AG samples.

Platelet aggregation Next, we tested the aggregation response of fresh and stored platelets to physiologically relevant agonists including ADP, collagen, and TRAP (Fig. 1). Cold-stored platelets aggregate significantly better than both RT-stored and fresh platelets when stimulated with ADP (Fig. 1A) and collagen (Fig. 1B). However, in the case of TRAP (Fig. 1C), all storage products decreased compared with fresh platelets, and no differences were seen between groups. There was no difference between 4°C platelets stored under nonagitated or agitated conditions. These data show that 4°C-stored platelets are more responsive to activating stimuli than RT-stored platelets.

TABLE 2. Metabolic and electrolyte levels in plasma during storage

	Baseline		RT		4°C		4°C + AG	
	Day 1	Day 3	Day 5	Day 3	Day 5	Day 3	Day 5	
pH	7.24 \pm .07	7.45 \pm .16	7.28 \pm .08	7.45 \pm .21	7.44 \pm .13	7.48 \pm .25	7.44 \pm .15	
Lactate, mg/dL	1.77 \pm 0.6	6.85 \pm 0.8*	12.87 \pm 2.1*	4.51 \pm 0.6*	5.67 \pm 0.3*†	4.83 \pm 0.7*	6.05 \pm 0.4*†	
Glucose, mg/dL	321.5 \pm 8.1	294.0 \pm 22.1*	238.3 \pm 42.5*	310.2 \pm 14.2*	307.5 \pm 25.0*†	312.4 \pm 28.1*	303.5 \pm 24.6*†	
Bicarbonate, mM	18.04 \pm 1.6	10.06 \pm 2.9*	6.27 \pm 0.9*	14.00 \pm 1.7*‡	12.78 \pm 0.8*†	14.01 \pm 2.7*‡	12.38 \pm 1.0*†	
pCO ₂ , mmHg	38.5 \pm 4.7	13.8 \pm 1.5*	13.1 \pm .25*	24.1 \pm 4.2*‡	19.5 \pm 4.4*†	22.6 \pm 5.6*‡	18.7 \pm 4.7*†	
pO ₂ , mmHg	91.8 \pm 9.7	113 \pm 17.8	132 \pm 41.2	120.4 \pm 32.7	139.8 \pm 26.8*†	116.6 \pm 31.0	128.0 \pm 29.3†	
Sodium, mM	137.5 \pm 1	139.5 \pm 3.7	140.25 \pm 1.3*	137.4 \pm 1.7	136.8 \pm .96†	137.8 \pm 1.6	136.5 \pm 1.3†	
Potassium, mM	3.24 \pm .18	3.34 \pm .23	3.45 \pm .31	3.52 \pm .19*	3.60 \pm .25*	3.52 \pm .25*	3.55 \pm .27*†	
Chloride, mM	97.6 \pm .89	101.3 \pm 4.9	104.8 \pm 1.3*	99.2 \pm 2.8	99.7 \pm 1.2†	99.0 \pm 2.2	98.8 \pm 1.7†	

*Values (mean \pm SD) significant from baseline represented by $P < 0.05$.

†Values significant from day 5 RT represented by $P < 0.05$.

‡Values significant from day 3 RT represented by $P < 0.05$.

TABLE 3. Glucose consumption and lactate production rates calculated from the measurements in Table 1

	RT		4°C		4°C + AG	
	Day 3	Day 5	Day 3	Day 5	Day 3	Day 5
Glucose consumption from baseline, $\mu\text{mol}/10^{12}$ platelets/h	30.6 \pm 7.9	75.6 \pm 25.0	17.1 \pm 11.3	25.3 \pm 3.6*	14.8 \pm 7.1*	29.4 \pm 9.8*
Lactate production from baseline, $\mu\text{mol}/10^{12}$ platelets/h	71.8 \pm 7.2	97.6 \pm 23.0	36.1 \pm 5.1*	40.0 \pm 8.6*	40.3 \pm 4.9*	44.6 \pm 8.5*

*Values (mean \pm SD) significant from corresponding RT samples represented by $P < 0.05$.

We also evaluated the effect of RBCs on aggregation response by adding 40% RBCs to platelets stored for 5 days (Fig. 1, D and E). The baseline AUC is higher than in Figure 1, A–C, because aggregation was monitored for 12 min instead of 6. We observed that the addition of RBCs significantly increased the response of fresh platelets to ADP. The response of RT platelets to ADP and collagen stimulation again declined significantly both with and without RBCs compared with baseline, whereas the response of cold-stored platelets was not enhanced by the addition of RBCs. Responses of day 5 4°C samples to ADP and collagen stimulation over 12 min were similar to baseline. Unlike the 6-min data, the 4°C collagen response was not enhanced at day 5 compared with fresh. Decreases in TRAP aggregation over time were not significant in 12-min aggregation tests for any group, and responses were not changed by the addition of RBCs (Fig. 1F).

Clot formation and strength We tested the kinetics of formation, strength, and stability of clots formed from fresh or stored platelets using TEG. Thromboelastography measures time to initial fibrin formation (reaction time [R]), time to clot formation (clot time [K]), rate of clot formation (α angle), clot strength (maximum amplitude [MA]), and clot lysis at 30 min. As can be seen in Figure 2A, the time to initial fibrin formation,

R time, was significantly faster with storage at 4°C without agitation, resulting in faster clot formation at day 3 compared with both baseline and RT storage. The clot generation time, K, for the RT samples was statistically different from baseline at day 3, but differences were clinically unimportant (Fig. 2B). The α angle was not significantly changed by storage method or duration (Fig. 2C). The clot strength, as measured by MA, decreased significantly on day 5 for RT-stored platelets but remained unaffected in 4°C-stored platelets (Fig. 2D). The clot lysis, as measured by clot lysis at 30 min, increased significantly after 5 days for RT-stored platelets, but remained constant for 4°C-stored platelets (Fig. 2E). With the exception of the day 3 R values, agitation did not alter any of the TEG parameters during storage at 4°C. Taken together, the TEG assay clearly demonstrates that 4°C-stored platelets retain the ability to support normal clotting, comparable to fresh platelets even after 5 days of storage, whereas RT-stored platelets form clots that are weak and susceptible to lysis. Conversely, cold storage enhanced initiation of fibrin formation.

Platelet activation marker expression We measured the levels of receptors expressed on activated platelets including P-selectin, lactadherin binding to exposed PS and CD40L (Fig. 3, A–C). We observed a significant increase in the

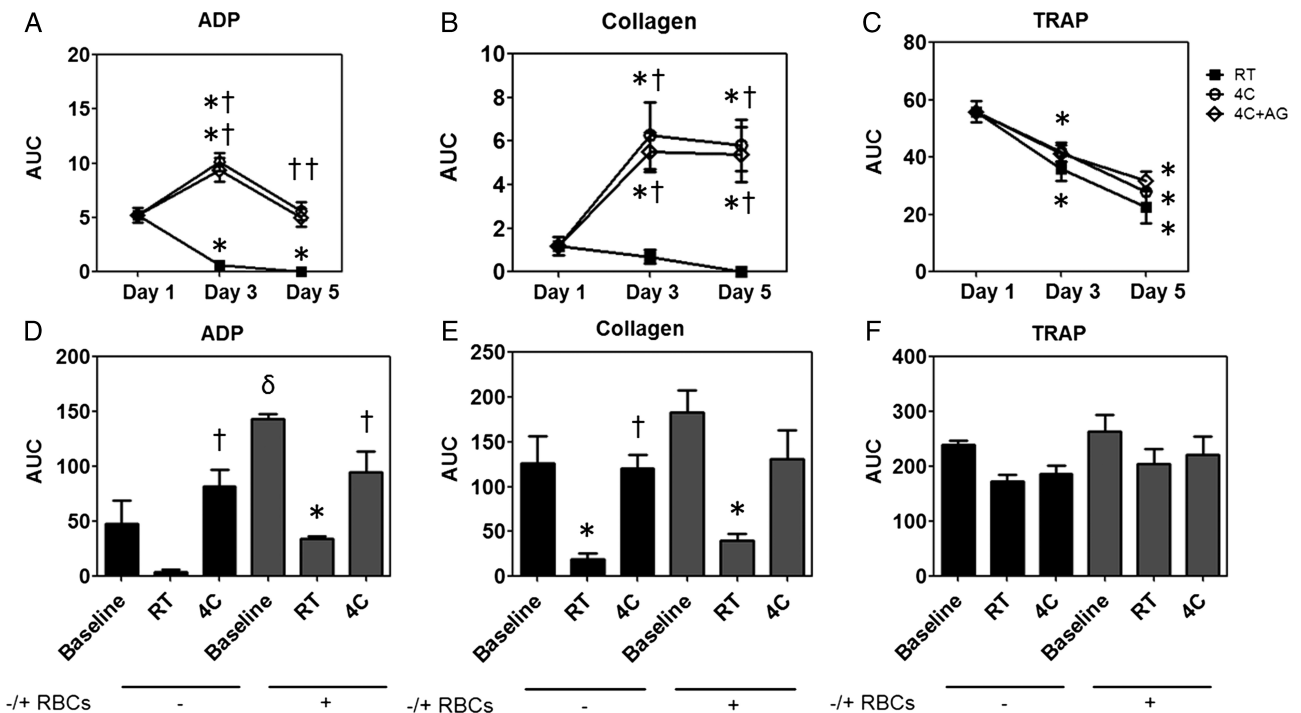


FIG. 1. Aggregation of platelets without (A, B, C) and with RBCs (D, E, F) when stimulated with (A, D) ADP, (B, E) collagen, and (C, F) TRAP. Bar graphs D, E, and F are comparisons between baseline versus day 5 RT and 4°C APs. Treatment conditions are represented as follows: RT = ■; 4°C = ⊖; 4°C + AG = ⊕. Area under the curves are represented as mean \pm SEM. Differences compared to Baseline (*), RT (†), and between samples without RBCs versus with (δ) are shown if both the one way ANOVA for repeated measures and the Bonferroni test comparisons are significant ($P < 0.05$). Note: Red blood cell AUC values are higher in graphs D–F compared with A–C because aggregation was measured over 12 min instead of 6.

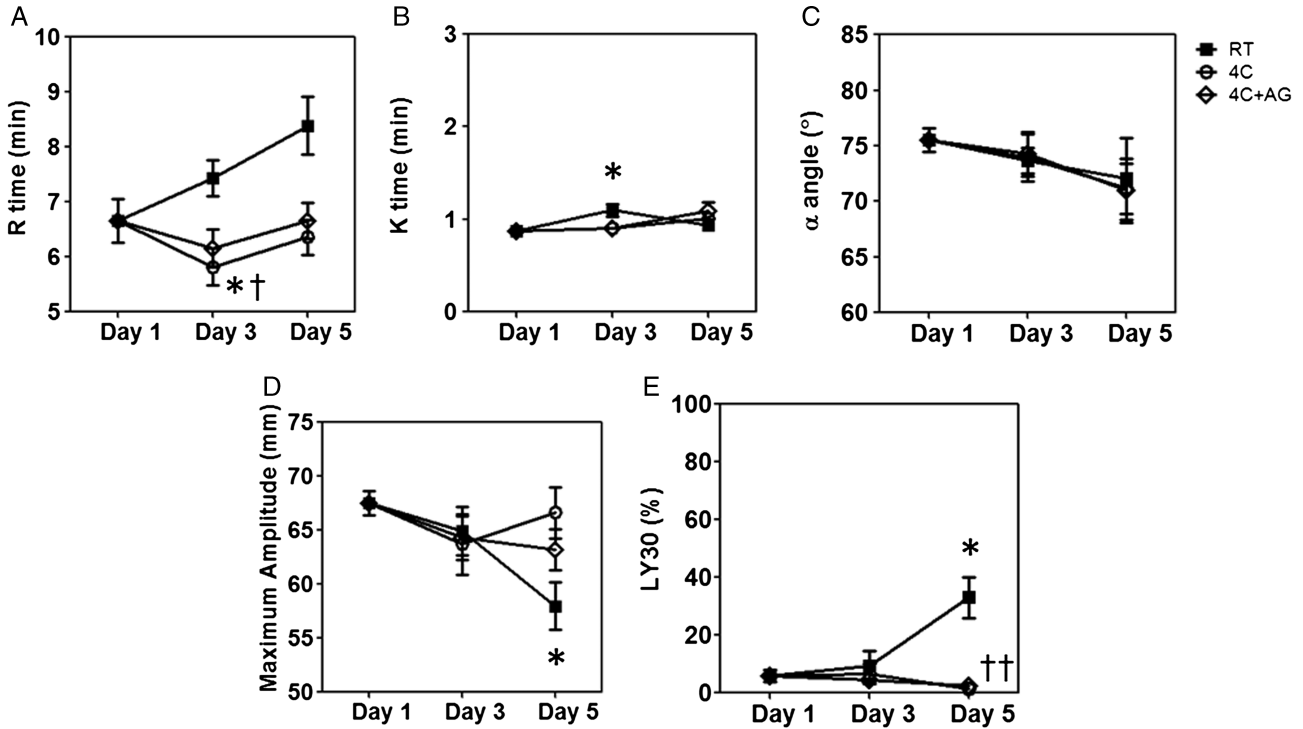


FIG. 2. Measurement of clot properties by TEG. A, R time. B, K time. C, α Angle. D, Maximum amplitude. E, Percent lysis after 30 minutes. Treatment conditions are represented as follows: RT = ■; 4°C = ○; 4°C + AG = ◇. Data are represented as mean ± SEM. Differences from baseline (*) and between treatment groups (†) are shown if results from both the one way ANOVA for repeated measures and the post hoc Bonferroni test comparisons are significant ($P < 0.05$).

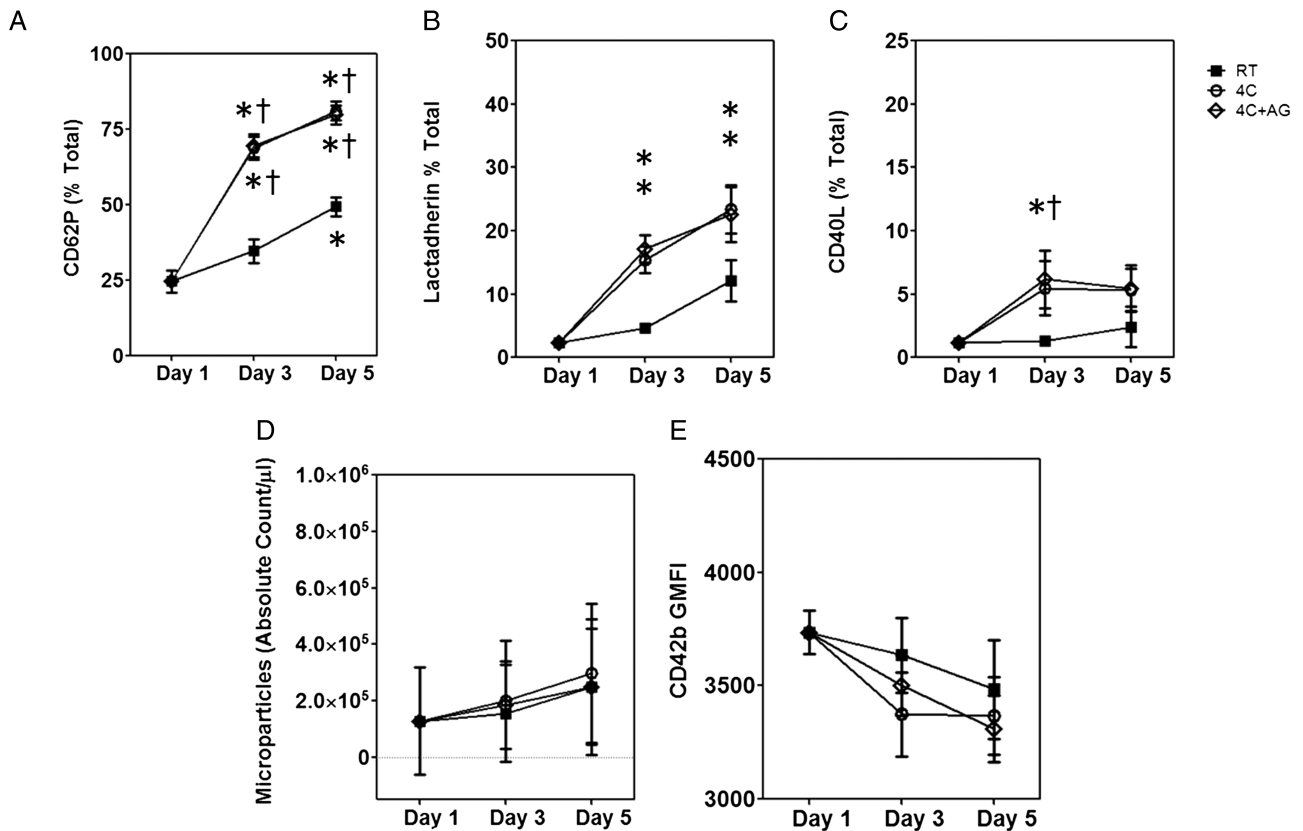


FIG. 3. Estimation of surface receptor levels by flow cytometry. A, P selectin. B, Lactadherin binding (PS exposure). C, CD40L expression. D, Micro particle release. E, Glycoprotein Iba. Treatment conditions are represented as follows: RT = ■; 4°C = ○; 4°C + AG = ◇. The expression levels are represented as mean ± SEM. Differences from baseline (*) and between treatment groups (†) are shown if results from both the one way ANOVA for repeated measures and the post hoc Bonferroni test comparisons are significant ($P < 0.05$).

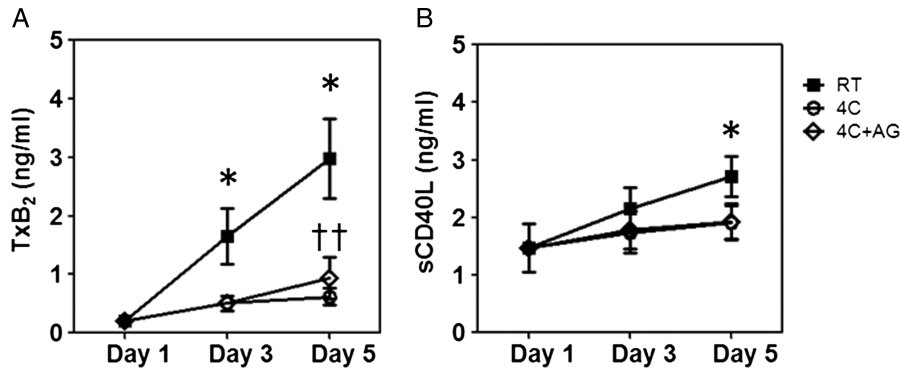


FIG. 4. Enzyme-linked immunosorbent assay quantification of soluble factors released by platelets. A, Thromboxane B₂. B, Soluble CD40L. Treatment conditions are represented as follows: RT = ■; 4°C = ○; 4°C + AG = ◇. The concentrations are represented as mean ± SEM. Differences from baseline (*) and between treatment groups (†) are shown if results from both the one way ANOVA for repeated measures and the post hoc Bonferroni test comparisons are significant ($P < 0.05$).

percentage of platelets expressing P-selectin (CD62P) over storage duration in both RT- and 4°C-stored platelets. The 4°C-stored platelets showed a greater increase in CD62P expression (Fig. 3A) compared with RT-stored platelets. Phosphatidylserine exposure, as measured by lactadherin binding (Fig. 3B), also increased with storage duration at 4°C. We used lactadherin instead of annexin V because lactadherin binding does not require Ca⁺² and has higher sensitivity (32). The CD40L expression on platelet surfaces was higher with cold storage and agitation on day 3 compared with baseline and RT conditions (Fig. 3C). Changes in microparticle count and CD42b surface expression were not significant either over time or between cold- and RT-stored samples (Fig. 3, D and E).

Release of proinflammatory mediators When platelets become activated, they release their granular contents and other metabolites into plasma. We tested the levels of sCD40L and TxB₂ released by platelets into plasma during storage (Fig. 4). Thromboxane B₂ is the stable end product of a metabolite of the arachidonic acid pathway, TxA₂, which intracellularly amplifies signals from other agonists such as ADP or thrombin. Room temperature-stored platelets released 4-fold more TxB₂ than 4°C-stored platelets after 5 days of storage, whereas the latter platelets did not release any significant quantities of TxB₂ (Fig. 4A). Soluble CD40L is a protein expressed in platelets that is known to play a role in inflammation, thrombosis, and restenosis (33). CD40L translocates from inside to the surface of agonist-activated platelets and is shed from the platelet surface as sCD40L. Room temperature-stored samples released slightly more sCD40L into plasma during storage compared with baseline on day 5 (Fig. 4B).

DISCUSSION

In this work, we demonstrated that apheresis platelets stored at 4°C are superior in metabolic and functional assays to platelets stored under standard blood banking conditions at RT. Furthermore, we find that agitation does not significantly influence the function of the 4°C-stored product. Our results suggest that 4°C-stored apheresis platelets would perform better to stop acute hemorrhage. Additional benefits of refrigeration would include decreased risk of bacterial contamination and simplified transport and storage because other

blood products must be maintained at 4°C (2–4). Cold platelets may be a viable alternative to RT-stored platelets for therapeutic, as opposed to prophylactic transfusion when immediate hemostasis is required.

As platelets stored in plasma metabolize glucose via the glycolytic pathway, they produce lactate and free hydrogen ions, which are buffered by bicarbonate in plasma to yield carbon dioxide and water (34). Glucose levels in 4°C-stored platelets did not change over time, consistent with decreased metabolic rates in the refrigerated samples. Similarly, changes in lactate and bicarbonate levels were also small. A similar effect was described in platelet concentrates from PRP stored at 4°C for up to 3 days (8). Stable pH levels correlate with preserved platelet viability; thus, the nearly constant pH measurements in the stored 4°C samples indicate that platelet quality is better maintained during refrigeration. In contrast, the RT-stored samples demonstrated a precipitous drop in glucose levels, increased lactate levels, and a corresponding decrease in bicarbonate levels, consistent with the increased metabolic rates previously described during RT storage (31, 35). The glucose consumption and lactate production rates during RT storage were comparable to those reported by Dumont and van den Broeke (31). Surprisingly, despite better metabolic indices, the platelet counts during 4°C storage were significantly lower compared with RT storage, suggesting adhesion to the bag, clumping, or cell death. This loss was abolished when platelets were stored in larger, standard-size bags, suggesting that higher surface-to-volume ratio in the smaller bags may contribute to the reduction in platelet count.

The various markers of platelet activation including CD62P expression, PS exposure, and expression of CD40L increased during 4°C storage, but only CD62P expression was greater at RT. A similar difference in CD62P expression and PS exposure between 4°C and RT storage has been reported for PRP (36). Phosphatidylserine exposure and the release of α -granule contents during RT storage are correlated with the loss in mitochondrial membrane potential and apoptosis due to depletion of glucose and lactate accumulation that occurs from continuous metabolic activity (37, 38). These markers represent different molecular events of activation: CD62P is released from platelet α -granules; PS is externalized from the inner leaflet of the platelet membrane; and CD40L is a

proinflammatory mediator translocated to the platelet membrane. All were higher in 4°C-stored platelets compared with RT despite significantly reduced metabolic rates. In addition, RT storage, but not cold storage, results in platelet degranulation, which has been correlated with activation (39). These data suggest that distinct mechanisms of activation are triggered at each storage temperature tested. While platelet activation at RT is related to aging or senescence, we speculate that cold-induced biophysical changes such as membrane rafting, glycoprotein Ib receptor clustering, and cytoskeletal rearrangement may trigger a set of downstream biochemical pathways distinct from aging that result in platelet activation at 4°C (15, 40–43).

In contrast, we found that RT-stored platelets release more proinflammatory mediators such as sCD40L and TxA₂ (measured as the more stable downstream product, TxB₂), possibly due to platelet degranulation. Elevated release of sCD40L at RT has been previously described and may play a potential role in adverse transfusion reactions (44, 45). Thromboxane A₂ is produced by the oxidation of arachidonic acid during platelet activation and is in itself a potent activator of platelets, thus may result in inappropriate coagulation upon transfusion (46). In essence, a decrease in soluble inflammatory mediators suggests that cold-stored platelets may be safer for transfusion than RT-stored platelets and may result in fewer adverse events such as transfusion-related acute lung injury. Low TxB₂ release from 4°C-stored platelets indicates that arachidonic acid is relatively preserved, suggesting that the dynamic range of platelet response to physiologic agonists after transfusion is similarly preserved and may explain our aggregometry results.

Circulating platelets are in a state of low activation but high responsiveness. The stimulation of fresh platelets with suboptimal concentrations of natural agonists not only activates the platelets but also decreases their response to subsequent stimulation with the same or different agonists (47–49). Both RT storage and 4°C storage activate platelets, but this activation leads to variable responses to chemical agonists. The markedly reduced ability of RT-stored platelets to aggregate upon stimulation with ADP, collagen, and TRAP is concordant with previous studies of platelets obtained via both the buffy coat and apheresis methods (50, 51). This phenomenon has been attributed to senescence due to progressive loss in energy-generating machinery (52). In contrast, 4°C-stored platelets aggregate as well as fresh platelets to ADP and collagen, possibly due to cold-induced platelet activation. This trend was maintained even after the addition of RBCs to mimic the physiological environment. Together, our data suggest that activation due to storage at 4°C “primes” the platelet to a state of heightened responsiveness without compromising the innate response to stimuli. Thus, it appears that the activation state attained by platelets due to 4°C storage may be different from activation states associated with either RT storage or prior activation with chemical stimuli. The similar decline in aggregation response to TRAP activation for both storage temperatures may be due to a loss of thrombin receptors (53).

The superior ability of 4°C-stored platelets to aggregate when stimulated with chemical agonists is accompanied by improved clot strength and stability. While RT storage negatively impacts clot strength and enhances fibrinolysis compared with

fresh platelets, significant loss of function was not seen in 4°C-stored samples. Specifically, TEG analysis indicates that despite preactivation, RT-stored platelets result in the formation of weaker clots that are less resistant to fibrinolysis, probably due to low thrombin generation levels (54). The maintenance of hemostatic function in 4°C-stored platelets, as estimated by TEG, is in line with previous reports (20). Although TEG measures global coagulation, the contribution of platelets is reflected predominantly in MA. In this study, the MA values confirm that platelet integrity and function are well preserved during cold storage without significantly changing AP clot strength.

We stored platelets at 4°C both with and without the method of agitation currently used by blood banks during standard RT storage. Our assays did not indicate that agitation during cold storage has a significant role in maintaining platelet health and function. Conversely, RT-stored platelets must be agitated either continuously or intermittently every 24 h to prevent poor gas exchange, acidosis (pH < 6), loss of normal morphology, and low platelet recovery (55–57). Thus, cold storage may reduce storage costs and minimize space by eliminating the need for platelet agitators during storage and transportation to remote locations. Furthermore, the cost of 4°C-stored platelets could be further lowered if reduced need for bacterial testing is proven.

In summary, our data demonstrate that cold-stored platelets have better functional competence and fewer inflammatory mediators than RT-stored platelets. Despite a few promising human trials in the 1970s, platelet refrigeration has been abandoned because of shorter circulation time after transfusion. Platelet utilization is shifting because of clinical practice guidelines that recommend 1:1:1 transfusion ratios for massive transfusion trauma protocols. The medical environment in which prophylactic transfusions predominated is giving way to one in which active bleeding is frequently the target of therapy. When treating severe hemorrhage, the superior hemostatic effectiveness of cold platelets may actually make them the better choice. Refrigerated platelets were the standard of care for more than a decade. Our results, along with a growing body of evidence, suggest the need to seriously consider the reinstatement of cold-stored platelets as a widely available therapeutic product.

ACKNOWLEDGMENTS

The authors thank Ron Bryant for collecting apheresis platelets.

REFERENCES

- Stronck DF, Rebull P: Platelet transfusions. *Lancet* 370(9585):427–438, 2007.
- Brecher ME, Hay SN, Rothenberg SJ: Evaluation of a new generation of plastic culture bottles with an automated microbial detection system for nine common contaminating organisms found in PLT components. *Transfusion* 44(3):359–363, 2004.
- Kuehnert MJ, Roth VR, Haley NR, Gregory KR, Elder KV, Schreiber GB, Arduino MJ, Holt SC, Carson LA, Banerjee SN, et al.: Transfusion-transmitted bacterial infection in the United States, 1998 through 2000. *Transfusion* 41(12):1493–1499, 2001.
- Currie LM, Harper JR, Allan H, Connor J: Inhibition of cytokine accumulation and bacterial growth during storage of platelet concentrates at 4 degrees C with retention of *in vitro* functional activity. *Transfusion* 37(1):18–24, 1997.
- Seghatchian J, Krailadsiri P: Platelet storage lesion and apoptosis: are they related? *Transfus Apher Sci* 24(1):103–105, 2001.

6. Klein HG, Spahn DR, Carson JL: Red blood cell transfusion in clinical practice. *Lancet* 370(9585):415 426, 2007.
7. McGill M, Brindley DC: Effects of storage on platelet reactivity to arterial subendothelium during blood flow. *J Lab Clin Med* 94(2):370 380, 1979.
8. Rock G, Figueredo A: Metabolic changes during platelet storage. *Transfusion* 16(6):571 579, 1976.
9. Shively JA, Gott CL, de Jongh DS: The effect of storage on adhesion and aggregation of platelets. *Vox Sang* 18(3):204 215, 1970.
10. Becker GA, Tuccelli M, Kunicki T, Chalos MK, Aster RH: Studies of platelet concentrates stored at 22 C and 4 C. *Transfusion* 13(2):61 68, 1973.
11. Slichter SJ, Harker LA: Preparation and storage of platelet concentrates. *Transfusion* 16(1):8 12, 1976.
12. Valeri CR: Circulation and hemostatic effectiveness of platelets stored at 4 C or 22 C: studies in aspirin-treated normal volunteers. *Transfusion* 16(1):20 23, 1976.
13. Murphy S, Gardner FH: Effect of storage temperature on maintenance of platelet viability deleterious effect of refrigerated storage. *N Engl J Med* 280(20):1094 1098, 1969.
14. Filip DJ, Aster RH: Relative hemostatic effectiveness of human platelets stored at 4 degrees and 22 degrees C. *J Lab Clin Med* 91(4):618 624, 1978.
15. Hoffmeister KM, Felbinger TW, Falet H, Denis CV, Bergmeier W, Mayadas TN, von Andrian UH, Wagner DD, Stossel TP, Hartwig JH: The clearance mechanism of chilled blood platelets. *Cell* 112(1):87 97, 2003.
16. Hoffmeister KM, Josefsson EC, Isaac NA, Clausen H, Hartwig JH, Stossel TP: Glycosylation restores survival of chilled blood platelets. *Science* 301(5639):1531 1534, 2003.
17. Josefsson EC, Gebhard HH, Stossel TP, Hartwig JH, Hoffmeister KM: The macrophage alphaMbeta2 integrin alphaM lectin domain mediates the phagocytosis of chilled platelets. *J Biol Chem* 280(18):18025 18032, 2005.
18. Kaufman RM: Uncommon cold: could 4 degrees C storage improve platelet function? *Transfusion* 45(9):1407 1412, 2005.
19. Egidi MG, D'Alessandro A, Mandarello G, Zolla L: Troubleshooting in platelet storage temperature and new perspectives through proteomics. *Blood Transfus* 8(Suppl 3):s73 s81, 2010.
20. Stiegler G, Fischer G, Ramanathan G, Bencur P, Weigel G, Mannhalter C: P-selectin mRNA is maintained in platelet concentrates stored at 4 degrees C. *Transfusion* 49(5):921 927, 2009.
21. Montgomery RK, Reddoch KM, Evani SJ, Cap AP, Ramasubramanian AK: Enhanced shear-induced platelet aggregation due to low-temperature storage. *Transfusion* 53(7):1520 1530, 2013.
22. Pidcoke HF, McFaul SJ, Ramasubramanian AK, Parida BK, Mora AG, Fedyk CG, Valdez-Delgado KK, Montgomery RK, Reddoch KM, Rodriguez AC, et al.: Primary hemostatic capacity of whole blood: a comprehensive analysis of pathogen reduction and refrigeration effects over time. *Transfusion* 53(Suppl 1):137S 149S, 2013.
23. Bock M, Rahrigh S, Kunz D, Lutze G, Heim MU: Platelet concentrates derived from buffy coat and apheresis: biochemical and functional differences. *Transfus Med* 12(5):317 324, 2002.
24. Krailadsiri P, Seghatchian J: Are all leucodepleted platelet concentrates equivalent? Comparison of Cobe LRS Turbo, Haemonetics MCS + LD, and filtered pooled buffy-coat-derived platelets. *Vox Sang* 78(3):171 175, 2000.
25. Metcalfe P, Williamson LM, Reutlingsperger CP, Swann I, Ouweland WH, Goodall AH: Activation during preparation of therapeutic platelets affects deterioration during storage: a comparative flow cytometric study of different production methods. *Br J Haematol* 98(1):86 95, 1997.
26. Sloand EM, Yu M, Klein HG: Comparison of random-donor platelet concentrates prepared from whole blood units and platelets prepared from single-donor apheresis collections. *Transfusion* 36(11 12):955 959, 1996.
27. Vassallo RR, Murphy S: A critical comparison of platelet preparation methods. *Curr Opin Hematol* 13(5):323 330, 2006.
28. Borchsen L, Johansson PI, Kristensen AT, Daugaard G, Ostrowski SR: The influence of platelets, plasma and red blood cells on functional hemostatic assays. *Blood Coagul Fibrinolysis* 22(3):167 175, 2011.
29. Bertolini F, Murphy S: A multicenter evaluation of reproducibility of swirling in platelet concentrates. *Transfusion* 34(9):796 801, 1994.
30. Mathai J, Resmi KR, Sulochana PV, Sathyabhama S, Baby Saritha G, Krishnan LK: Suitability of measurement of swirling as a marker of platelet shape change in concentrates stored for transfusion. *Platelets* 17(6):393 396, 2006.
31. Dumont LJ, van den Broeke T: Seven-day storage of apheresis platelets: report of an *in vitro* study. *Transfusion* 43(2):143 150, 2003.
32. Albanyan AM, Murphy MF, Rasmussen JT, Heegaard CW, Harrison P: Measurement of phosphatidylserine exposure during storage of platelet concentrates using the novel probe lactadherin: a comparison study with annexin V. *Transfusion* 49(1):99 107, 2009.
33. Andre P, Nannizzi-Alaimo L, Prasad SK, Phillips DR: Platelet-derived CD40L: the switch-hitting player of cardiovascular disease. *Circulation* 106(8):896 899, 2002.
34. Baker JM, Candy DJ, Hawker RJ: Influences of pH on human platelet metabolism. *Platelets* 12(6):333 342, 2001.
35. Kilkson H, Holme S, Murphy S: Platelet metabolism during storage of platelet concentrates at 22 degrees C. *Blood* 64(2):406 414, 1984.
36. Babic AM, Josefsson EC, Bergmeier W, Wagner DD, Kaufman RM, Silberstein LE, Stossel TP, Hartwig JH, Hoffmeister KM: *In vitro* function and phagocytosis of galactosylated platelet concentrates after long-term refrigeration. *Transfusion* 47(3):442 451, 2007.
37. Albanyan AM, Harrison P, Murphy MF: Markers of platelet activation and apoptosis during storage of apheresis- and buffy coat-derived platelet concentrates for 7 days. *Transfusion* 49(1):108 117, 2009.
38. Bertino AM, Qi XQ, Li J, Xia Y, Kuter DJ: Apoptotic markers are increased in platelets stored at 37 degrees C. *Transfusion* 43(7):857 866, 2003.
39. Macey MG, Carty E, Webb L, Chapman ES, Zelmanovic D, Okrongly D, Rampton DS, Newland AC: Use of mean platelet component to measure platelet activation on the ADVIA 120 haematology system. *Cytometry* 38(5):250 255, 1999.
40. Goussset K, Tsvetkova NM, Crowe JH, Tablin F: Important role of raft aggregation in the signaling events of cold-induced platelet activation. *Biochim Biophys Acta* 1660(1 2):7 15, 2004.
41. Rumjantseva V, Grewal PK, Wandall HH, Josefsson EC, Sorensen AL, Larson G, Marth JD, Hartwig JH, Hoffmeister KM: Dual roles for hepatic lectin receptors in the clearance of chilled platelets. *Nat Med* 15(11):1273 1280, 2009.
42. White JG, Rao GH: Microtubule coils versus the surface membrane cytoskeleton in maintenance and restoration of platelet discoid shape. *Am J Pathol* 152(2):597 609, 1998.
43. Gitz E, Koekman CA, van den Heuvel DJ, Deckmyn H, Akkerman JW, Gerritsen HC, Urbanus RT: Improved platelet survival after cold storage by prevention of glycoprotein Iba clustering in lipid rafts. *Haematologica* 97(12):e70 e72, 2012.
44. Kaufman J, Spinelli SL, Schultz E, Blumberg N, Phipps RP: Release of biologically active CD154 during collection and storage of platelet concentrates prepared for transfusion. *J Thromb Haemost* 5(4):788 796, 2007.
45. Khan SY, Kelher MR, Heal JM, Blumberg N, Boshkov LK, Phipps R, Gettings KF, McLaughlin NJ, Silliman CC: Soluble CD40 ligand accumulates in stored blood components, primes neutrophils through CD40, and is a potential cofactor in the development of transfusion-related acute lung injury. *Blood* 108(7):2455 2462, 2006.
46. Paul BZ, Daniel JL, Kunapuli SP: Platelet shape change is mediated by both calcium-dependent and -independent signaling pathways. Role of p160 Rho-associated coiled-coil containing protein kinase in platelet shape change. *J Biol Chem* 274(40):28293 28300, 1999.
47. Baurand A, Eckly A, Hechler B, Kauffenstein G, Galzi JL, Cazenave JP, Leon C, Gachet C: Differential regulation and relocalization of the platelet P2Y receptors after activation: a way to avoid loss of hemostatic properties? *Mol Pharmacol* 67(3):721 733, 2005.
48. Hardy AR, Conley PB, Luo J, Benovic JL, Poole AW, Mundell SJ: P2Y1 and P2Y12 receptors for ADP desensitize by distinct kinase-dependent mechanisms. *Blood* 105(9):3552 3560, 2005.
49. Peerschke EI: Ca²⁺ mobilization and fibrinogen binding of platelets refractory to adenosine diphosphate stimulation. *J Lab Clin Med* 106(2):111 122, 1985.
50. Curvers J, van Pampus EC, Feijje MA, Rombout-Sestrienkova E, Giesen PL, Heemskerk JW: Decreased responsiveness and development of activation markers of PLTs stored in plasma. *Transfusion* 44(1):49 58, 2004.
51. Rinder HM, Murphy M, Mitchell JG, Stocks J, Ault KA, Hillman RS: Progressive platelet activation with storage: evidence for shortened survival of activated platelets after transfusion. *Transfusion* 31(5):409 414, 1991.
52. Holme S, Heaton A: *In-vitro* platelet aging at 22-degrees-C is reduced compared to *in-vivo* aging at 37-degrees-C. *Br J Haematol* 91(1):212 218, 1995.
53. Schlagenhaut A, Kozma N, Leschnik B, Wagner T, Muntean W: Thrombin receptor levels in platelet concentrates during storage and their impact on platelet functionality. *Transfusion* 52(6):1253 1259, 2012.
54. Svendsen MS, Rojkaer R, Kristensen AT, Salado-Jimena JA, Kjalke M, Johansson PI: Impairment of the hemostatic potential of platelets during storage as evaluated by flow cytometry, thrombin generation, and thrombelastography under conditions promoting formation of coated platelets. *Transfusion* 47(11):2057 2065, 2007.
55. Mitchell SG, Hawker RJ, Turner VS, Hesslewood SR, Harding LK: Effect of agitation on the quality of platelet concentrates. *Vox Sang* 67(2):160 165, 1994.
56. Skripchenko A, Myrup A, Thompson-Montgomery D, Awatefe H, Moroff G, Wagner SJ: Periods without agitation diminish platelet mitochondrial function during storage. *Transfusion* 50(2):390 399, 2010.
57. Snyder EL, Bookbinder M, Kakaiya R, Ferri P, Kiraly T: 5-Day storage of platelet concentrates in Clx containers effect of type of agitation. *Vox Sang* 45(6):432 437, 1983.