Critical Care Guideline and Consensus Statement



57

58

59

60

61

62

63

64

65 66

67

68

69

70

71

72

78

79

98

100

101

102

103

5 6 7 8

2

3

4

9

10

11

13

14

15

16

17

18

19

20

21

22

23

24

25

26 27

28

29

30

31

32

33

34

35

36

37 38

39

40

41

42

43 44

Red Blood Cell Transfusion in Critically Ill Q1 Q37 Adults

An American College of Chest Physicians Clinical Practice Guideline

Angel O. Coz Yataco, MD; Israa Soghier, MD; Paul C. Hébert, MD; Emilie Belley-Cote, MD; Margaret Disselkamp, MD; David Flynn, MS; Karin Halvorson, MD; Jonathan M. Iaccarino, MD; Wendy Lim, MD; Christina C. Lindenmeyer, MD; Peter J. Miller, MD; Kevin O'Neil, MD; Kathryn M. Pendleton, MD; Lisa Vande Vusse, MD; and Daniel R. Ouellette, MD

> BACKGROUND: Blood products frequently are administered to critically ill patients. Consid-73 ering recent trials and practice variability, a comprehensive review of current evidence was ⁷⁴ deemed essential to offer pertinent guidance to critical care practitioners. This American College of Chest Physicians (CHEST) guidelines panel examined the literature on RBC transfusions among critically ill patients overall and specific subgroups, including patients with gastrointestinal bleeding, acute coronary syndrome (ACS), cardiac surgery, isolated troponin elevation, and septic shock, to provide evidence-based recommendations.

> STUDY DESIGN AND METHODS: A panel of experts developed 6 Population, Intervention, 81 Comparator, and Outcome questions addressing RBC transfusions in critically ill patients 82 and performed a comprehensive evidence review. The panel applied the Grading of Rec- 83 ommendations, Assessment, Development, and Evaluations approach to assess the certainty 84 of evidence and to formulate and grade recommendations. A modified Delphi technique was 85 used to reach consensus on the recommendations.

> RESULTS: The initial search identified a total of 3,082 studies, and after the initial screening, 88 38 articles were reviewed. Among them, 23 studies met inclusion criteria, comprising 22 89 randomized controlled trials and 1 cohort study. Based on the analysis of these studies, the 90 panel formulated 2 strong and 4 conditional recommendations. The overall quality of evi- 91 dence for recommendations ranged from very low to moderate.

> 93 CONCLUSIONS: In most critically ill patients, a restrictive strategy was preferable to a permissive approach because it does not increase the risk of death or complications, but does decrease RBC use significantly. Data from critically ill subpopulations also supported a restrictive approach, except in patients with ACS, for whom favoring a restrictive approach could increase adverse outcomes. CHEST 2024; **■**(**■**):**■**-**■**

KEY WORDS: acute coronary syndrome; cardiac surgery; critically ill; GI bleed; red blood cell; sepsis; septic shock; transfusion

45 46 47

48

53

54

55

blood cell; RCT = randomized controlled trial; RR = risk ratio AFFILIATIONS: From the Critical Care Medicine Division and Pulmonary Medicine Division (A. O. C. Y.), Integrated Hospital-Care Institute, the Department of Gastroenterology, Hepatology and

Nutrition (C. C. L.), Cleveland Clinic; the Cleveland Clinic Lerner

ABBREVIATIONS: ACS = acute coronary syndrome; CHEST = American College of Chest Physicians; ESICM = European Society of

Intensive Care Medicine; GIB = gastrointestinal bleeding; LOS = length

of stay; MD = mean difference; MI = myocardial infarction; RBC = red

College of Medicine of Case Western Reserve University (A. O. C. Y.), 104 Cleveland, OH; the Division of Pulmonary and Critical Care Medicine 105 (I. S.), Department of Medicine, Salem Hospital/Massachusetts General Brigham, Salem; the Boston University Chobanian & Avedisian School of Medicine (D. F.), Boston, MA; the American College of Chest 107 Physicians (I. S. and J. M. I.), Glenview, IL; the Department of Critical 108 Care and Pulmonary Medicine (M. D.), Lexington Veterans Affairs Healthcare System, Lexington, KY; the Department of Medicine (K. H.), John A. Burns School of Medicine, University of Hawaii, 110

Summary of Recommendations and Suggestions

These recommendations do not apply to critically ill adults with hemodynamic instability due to acute hemorrhage, or those with neurological injuries or trauma.

- 1. In critically ill patients, we recommend a restrictive RBC transfusion strategy over a permissive RBC transfusion strategy (Strong Recommendation, Moderate Certainty of Evidence).
- 2. In critically ill patients with acute gastrointestinal bleeding, we recommend a restrictive RBC transfusion strategy over a permissive RBC transfusion strategy (Strong Recommendation, Moderate Certainty of Evidence).
- 3. In critically ill patients with acute coronary syndrome, we suggest against a restrictive RBC transfusion strategy (Conditional Recommendation, Low Certainty of Evidence).
- 4. In critically ill patients undergoing cardiac surgery, we suggest a restrictive RBC transfusion strategy over a permissive RBC transfusion strategy in the perioperative period (Conditional Recommendation, Moderate Certainty of Evidence).
- 5. In critically ill patients with isolated elevation of serum troponin without other evidence of cardiac

Honolulu, HI; the Section of Pulmonary, Critical Care, Allergy and Immunologic Disease (P. J. M.), Section on Hematology and Oncology, Department of Medicine, Section on Critical Care Medicine, Department of Anesthesiology, Wake Forest School of Medicine, Winston-Salem; the Wilmington Health and MICU (K. O.), Novant New Hanover Regional Medical Center, Wilmington, NC; the Division of Pulmonary, Allergy, Critical Care and Sleep Medicine (K. M. P.), Department of Medicine, University of Minnesota, Minneapolis, MN; the Division of Pulmonary, Critical Care and Sleep Medicine (L. V. V.), Department of Medicine, University of Washington, Seattle, WA; the Division of Pulmonary and Critical Care Medicine (D. R. O.), Henry Ford Hospital, Detroit, MI; the Bruyere Research Institute (P. C. H.), University of Ottawa, Ottawa; the Population Health Research Institute, (E. B.-C.), and the Department of Medicine (W. L.), McMaster University, Hamilton, ON, Canada.

DISCLAIMER: American College of Chest Physician guidelines are intended for general information only, are not medical advice, and do not replace professional medical care and physician advice, which always should be sought for any medical condition. The complete disclaimer for this guideline can be accessed at https://www.chestnet.org/Guidelines-and-Resources.

CORRESPONDENCE TO: Angel O. Coz Yataco, MD; email: cozyata@ccf.org

Copyright © 2024 The Author(s). Published by Elsevier Inc under license from the American College of Chest Physicians. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/).

DOI: https://doi.org/10.1016/j.chest.2024.09.016

ischemia, we suggest a restrictive RBC transfusion strategy over a permissive RBC transfusion strategy (Conditional Recommendation, Very Low Certainty of Evidence). 6. In patients with septic shock and end-organ hypoperfusion, we suggest against adding permissive RBC transfusion thresholds to usual care (Conditional Recommendation, Low Certainty of Evidence).

Remarks: Studies evaluating protocol driven approaches to goal-directed therapy in septic shock were not considered in the evidence review.

Background

In the United States, approximately 25% of critically ill patients receive RBC transfusions, totaling approximately 1.8 million units annually. The primary indication is low hemoglobin (80%), whereas less frequently encountered indications include active bleeding (27%) and hemodynamic instability (23%). Since the publication of the TRICC trial, 5 > 30 trials have examined RBC transfusion strategies in a variety of clinical settings. Recently, new studies and updated meta-analyses and guidelines have been published. Despite this evidence, significant variability exists in clinical practice regarding the indications for RBC transfusions, with most occurring in patients with hemoglobin levels of > 7 g/dL.

Although RBC transfusions can be life-saving, they carry significant risks of adverse effects, including transfusionrelated acute lung injury, transfusion-associated circulatory overload, and immunomodulating effects that may increase the risk of nosocomial infections. 1,6 These side-effects may be severe and even lifethreatening. The entire process, from distribution to administration of RBCs, incurs substantial costs, which vary globally.^{7,8} Optimal health care delivery minimizes unnecessary RBC transfusions, preserving them for patient groups with proven benefit. By optimizing the management of limited resources like RBCs, both individual patients and the broader at-risk critically ill population stand to benefit.9 Given new evidence and ongoing variability in practice, an expert panel identified, synthesized, and weighted the evidence to provide clinical recommendations for RBC transfusion in critically ill patients.

Methods

Standardized methodology for clinical practice guidelines as per American College of Chest Physicians (CHEST) 221 policy was followed. At this juncture, with several studies 222 underway in critically ill patients with neurologic injuries 223 and the trauma population, we decided to forego recom-224 mendations in these critical care subgroups. 225 **Expert Panel Composition** 226 227 228 229 230 231 232 233 Committee. 234 235 Conflicts of Interest 236 237 238 239 240 241 Question Development 242 243 244 245 246 247 248

The cochairs nominated a diverse and multidisciplinary panel based on their expertise encompassing critical care medicine, cardiology, hematology, and gastroenterology and hepatology. The final panel consisted of the 2 guideline cochairs, 9 panelists, 2 methodologists, 1 medical librarian, and 1 liaison to CHEST Guidelines Oversight

Financial relationships for each chair and the panelists were reviewed by the CHEST Professional Standards Committee for potential conflicts of interest according to the CHEST Conflict of Interest Policy. 10

The panel developed 6 clinical questions using the Population, Intervention, Comparator, and Outcome format regarding the transfusion of RBC in different clinical scenarios. The panel ranked outcomes for each question to determine critical and important outcomes a priori (e-Table 1).

Literature Search

249

250

251

252

253

254

255

256

257

258

259

260

261

262

263

264

265

266

267

268

269

270

271

272

273

274

275

A comprehensive search of MEDLINE, Embase, and the Cochrane Central Register of Controlled Trials using relevant key words was performed in May 2021 (e-Table 2), with an updated search performed in MED-LINE in January 2024. Searches were limited to English language randomized controlled trials (RCT), cohort studies, and case-control studies with at least 30 participants. Systematic reviews and prior guidelines were reviewed for context and completeness.

Results

The hemoglobin thresholds prompting RBC transfusion varied across studies, with a restrictive threshold generally defined as a hemoglobin level of 7 to 8 g/dL and a permissive threshold typically ranging from a hemoglobin level of 8.5 to 10 g/dL (Table 1). Hemodynamic instability typically has been defined as hypotension (mean arterial pressure, < 65 mm Hg, or systolic BP, < 100 mm Hg), tissue hypoperfusion caused by acute bleeding, or both.

Question 1: Should critically ill patients be treated with a restrictive or permissive RBC transfusion strategy?

Study Selection and Data Extraction

Relevant citations identified during the literature search were reviewed in duplicate using predefined criteria over 2 rounds of study selection: reviewing titles and abstracts in the first round and reviewing full texts in the second round (e-Figs 1-6). Data were 282 extracted, analyzed, and summarized for each individ- 283 ual Population, Intervention, Comparator, Outcome question.

276

277

278

280

284

285

286

287

298

299

300

301

315

316

317

319

320

321

322

323

324

325

326

327

Assessing the Quality of Evidence

Risk of bias was assessed using the Cochrane Risk of Bias 288 Tool for RCTs and the Risk of Bias in Nonrandomized 289 Studies of Intervention based on study design. 11,12 A 290 meta-analysis was performed when possible using a 291 random effects model. Results are reported as risk ratios 292 (RRs) for dichotomous outcomes and mean differences (MDs) for continuous outcomes with accompanying 95% CIs. The overall certainty of the evidence was 296 assessed for each outcome of interest using the Grading of Recommendations, Assessment, Development, and Evaluations approach (e-Table 3).¹³

Development of Recommendations

The panel reviewed and discussed the evidence. Recom- 302 mendations were drafted using the Grading of Recom- 303 mendations, Assessment, Development, Evaluations approach, with strong recommendations us- 305 ing the wording "we recommend" and conditional recommendations using the wording "we suggest." 14 307 Panel members voted individually via SurveyMonkey on the direction and strength of the recommendation.¹ Per CHEST policy, consensus was achieved with 80% agreement in directionality with at least 75% of the panel participating. The guidelines were reviewed 313 and approved by the Guidelines Oversight Committee 314 and CHEST presidential leadership.

1. In critically ill patients, we recommend a restrictive $_{318}$ RBC transfusion strategy over a permissive RBC transfusion strategy (Strong Recommendation, Moderate Certainty of Evidence).

Justification

This recommendation, applicable to most critically ill patients, is supported by evidence from several highquality RCTs involving approximately 16,000 patients. Comparisons between a restrictive and a permissive RBC 328 transfusion strategy yielded no significant differences in ICU mortality (RR, 1.00; 95% CI, 0.8-1.25), 5,16-19 30-day

TABLE 1 Hemoglobin Thresholds in Studies Included per Recommendation

	Hemoglobin Threshold, g/dL	
Population	Restrictive	Permissive
Overall critically ill	7.0-8.0	9.0-10.0
Gastrointestinal bleeding	7.0-8.0	8.0-10.0
Acute coronary syndrome	7.0-8.0	10.0
Underwent cardiac surgery	7.5-8.0	8.5-10.0
Isolated troponin elevation	N/A	N/A
Septic shock	7.0	9.0

Data are presented as ranges. N/A = no studies available.

mortality (RR, 0.99; 95% CI, 0.87-1.13), 5,17-30 or 1-year mortality (RR, 0.99; 95% CI, 0.87-1.13).31,32 Although the restrictive group exhibited a slightly longer ICU length of stay (LOS) than the permissive group (MD, +0.12 days; 95% CI, +0.01 to +0.23 days), this finding is not clinically significant. Additionally, no difference in hospital LOS was found (MD, -0.2 days; 95% CI, -0.51 to +0.12 days).^{5,17-21,23-29,33-36}

The restrictive approach proved superior to the permissive strategy in reducing adverse event rates (RR, 0.45; 95% CI, 0.22-0.94), but not in reducing secondary infections (RR, 1.03; 95% CI, 0.94-1.12). Similarly, no differences were found in organ-specific or systemspecific adverse events, including cardiac, renal, pulmonary, and thromboembolic complications (Table 2).5,18-30,33-36

Given the absence of a discernible impact on mortality and the potential reduction in overall adverse events, a restrictive approach was determined to be the preferred strategy. This approach minimized RBC use without any clinical consequences in most critically ill patients. Since

the publication of the TRICC trial,⁵ all studies have favored a more restrictive approach with the possible exception of the recently published MINT trial.²⁰ In the ^{Q11} trials evaluating ICU mortality, the clinical teams were encouraged strongly to transfuse 1 RBC unit at a time and repeat hemoglobin measurements after each unit. Using this approach, the number of RBC units transfused was decreased by 50%.5,16-19

Given the complexity of ICU care, specific subpopulations were addressed in subsequent recommendations.

What Others Are Saying

This recommendation aligned with those of other professional societies, including the Critical Care Societies Collaborative, the European Society of Intensive Care Medicine (ESICM), and the Association for the Advancement of Blood & Biotherapies. 37-39

Research Priorities

Although the use of restrictive RBC transfusion strategies alone may not need much further inquiry, using individual patient data meta-analysis may help to identify more specific subgroup effects, particularly in patients with various forms of cardiovascular diseases, ranging from acute ischemia to chronic heart diseases. Research on measures of oxygen use or biomarkers of oxygen delivery in tissue beds and overall may allow more targeted approaches to RBC transfusions.

Question 2: Should critically ill patients who have acute gastrointestinal bleeding be treated with a restrictive or permissive RBC transfusion strategy?

2. In critically ill patients with acute gastrointestinal bleeding, we recommend a restrictive RBC transfusion

TABLE 2 Pooled Analysis Comparing Restrictive vs Permissive Strategies in Overall Critically Ill Patients

Outcome	Relative Risk (95% CI)	Absolute Risk (95% CI)
ICU mortality	1.00 (0.80-1.25)	0 fewer per 1,000 (46 fewer-57 more)
1-y mortality	0.99 (0.87-1.13)	4 fewer per 1,000 (54 fewer-54 more)
30-d mortality	0.99 (0.87-1.13)	1 fewer per 1,000 (14 fewer-14 more)
ICU length of stay		0.12 d higher (0.01 higher-0.23 higher)
Hospital length of stay		0.2 d lower (0.51 lower-0.12 higher)
Adverse events	0.45 (0.22-0.94)	8 fewer per 1,000 (11 fewer-1 fewer)
Secondary infections	1.03 (0.94-1.12)	3 more per 1,000 (6 fewer-13 more)
Cardiac adverse events	0.94 (0.77-1.16)	4 fewer per 1,000 (16 fewer-11 more)
Renal adverse events	0.99 (0.89-1.10)	1 fewer per 1,000 (9 fewer-8 more)
Pulmonary adverse events	0.98 (0.88-1.08)	2 fewer per 1,000 (15 fewer-10 more)
Thromboembolism	0.83 (0.60-1.15)	4 fewer per 1,000 (9 fewer-3 more)

strategy over a permissive RBC transfusion strategy (Strong Recommendation, Moderate Certainty of Evidence).

Justification

441

442

443

444

445

446

447

448

449

450

451

452

453

454

455

456

457

458

459

460

461

462

463

464

465

466

467

468

469

470

471

472

473

474

475

476

477

478

479_{Q12}

480

481

482

483

484

485

486

487

488

489

490

491 492

493

494

495

Acute gastrointestinal bleeding (GIB) is a lifethreatening condition that often necessitates ICU care, including the critical decision of when to transfuse RBCs. We identified 3 relevant RCTs that included patients with acute upper GIB and an observational study of patients with acute lower GIB. 24-26,40

Villanueva et al²⁴ randomized 921 patients with acute upper GIB and early access to endoscopy to either a restrictive or permissive RBC transfusion strategy. The restrictive approach led to lower rates of rebleeding, fewer adverse events, and lower 6-week all-cause mortality. An open-label, cluster RCT with 936 patients reported no significant mortality difference between both RBC transfusion strategies, but noted lower transfusion rates and decreased health care costs with the restrictive approach.²⁶ A smaller single-center openlabel RCT showed that a restrictive approach was noninferior to a permissive strategy.²⁵ In aggregate, the restrictive transfusion strategy reduced short-term mortality (RR, 0.68; 95% CI, 0.48-0.97) without affecting hospital LOS (MD, -0.69 days; 95% CI, -1.98 to +0.60 days).

The restrictive approach proved superior to the permissive strategy with lower risk of acute transfusion reactions and serious adverse transfusion effects. However, no significant differences were found in the risk of infections, need for surgery in upper GIB (Table 3), or organ-specific or system-specific adverse events, including cardiac, renal, pulmonary, and thromboembolic (EP-2). The data on lower GI bleeding was limited to a retrospective study, which did not show differences in mortality or need for surgery between a restrictive and a permissive approach. 40

Given the lower risk of mortality and adverse reactions, 496 the adoption of a restrictive transfusion strategy emerged as the preferred recommendation. The restrictive strategy was superior in patients with and without portal hypertension-related GIB. Notably, portal pressure gradients increased in patients treated with a permissive strategy within the first 5 days of bleeding onset.²⁴

497

498

499

500

501

502

503

504

505

506

507

508

509

510

511

512

513

515

516

517

518

519

520

521

522

523

524

525

526

527

528

529

530

531

532

533

537

538

539

540

550

What Others Are Saying

This recommendation aligned with those of the American Association for the Study of Liver Diseases, the American College of Gastroenterology, and the American Gastroenterological Association. 41-44

Research Priorities

Studying the feasibility, implementation, and adherence 514 to a restrictive transfusion strategy in acute GIB is a priority. Additionally, examining interprofessional knowledge transmission through a systems-based approach that adheres to the principles of quality improvement is essential.

Question 3: Should critically ill patients with acute coronary syndrome be treated with a restrictive or permissive RBC transfusion strategy?

3. In critically ill patients with acute coronary syndrome, we suggest against a restrictive RBC transfusion strategy (Conditional Recommendation, Low Certainty of Evidence).

Justification

The optimal transfusion strategy for patients with acute coronary syndrome (ACS) has been controversial, balancing improved oxygen delivery to the myocardium 534 against the potential expansion of vascular volume and 535 increase in blood viscosity from overtransfusion.⁴⁵ This 536 recommendation is supported by 4 RCTs involving a total of 4,324 patients.²⁰⁻²³

TABLE 3 Pooled Analysis Comparing Restrictive vs Permissive Strategies in Gastrointestinal Bleeding

TABLE 5 1 001ed / that/515 Comparing Restrictive vs	r crimosive strategi		_ 54
Outcome	Relative Risk (95% CI)	Absolute Risk (95% CI)	_ 54:
30-d mortality	0.68 (0.48-0.97)	27 fewer per 1,000 (43 fewer-2 fewer)	54
Hospital length of stay		0.69 d lower (1.98 lower-0.6 higher)	0 54
Need for surgery in upper gastrointestinal bleeding	0.67 (0.16-2.91)	13 fewer per 1,000 (33 fewer-74 more)	54
Acute transfusion reactions	0.35 (0.20-0.61)	37 fewer per 1,000 (45 fewer-22 fewer)	54 54
Adverse transfusion effects ^a	0.73 (0.58-0.91)	54 fewer per 1,000 (83 fewer-18 fewer)	54 54
Infections in upper gastrointestinal bleeding	0.96 (0.79-1.17)	11 fewer per 1,000 (58 fewer-47 more)	_ 54

^aSerious adverse events defined as an event that endangers the health or safety of the patient.

551 The pooled analysis showed a trend toward higher 552 mortality with the restrictive approach (RR, 1.13; 553 95% CI, 0.67-1.91), but this difference was not 554 statistically significant. The MINT trial substantially 555 influenced this analysis because of its large sample size 556 because it recruited 3 times more patients than the other 557 studies combined. The 30-day mortality in the restrictive 558 group was 9.9%, compared with 8.3% in the permissive 559 group, a difference not reaching statistical significance 560 (RR, 1.19; 95% CI, 0.96-1.47). However, cardiac death 561 rates were 5.5% in the restrictive group and 3.2% in the 562 563 permissive group (RR, 1.74; 95% CI, 1.26-2.40). 564 Moreover, the point estimates for myocardial infarction 565 (MI) or death and for recurrent MI consistently favored 566 the permissive strategy.²⁰ Although a statistically 567 significant increase in adverse outcomes such as death 568 and recurrent MI might not have been found, concern 569 exists that a restrictive approach allowing hemoglobin 570 levels of 7 or 8 g/dL might increase the risk of adverse 571 outcomes in patients with acute MI. From these data, it 572 was unclear whether a gradient effect was present in 573 which risk progressively increased to < 10 g/dL or a 574 threshold effect at 10 g/dL. That is, these data do not 575 indicate whether 9 g/dL is as safe as 10 g/dL. Patient 576 577 symptoms and physiologic variables should be 578 considered when choosing a transfusion threshold for 579 patients with ACS. 580 581 582 583 584 585 586 587

588

589

590

591

592

593

594 595

596 597

598

599 600

601 602

603

604

605

Long-term follow-up of patients described a 1-year mortality of 23.1% in the restrictive and 20.4% in the permissive group (RR, 1.87; 95% CI, 0.74-4.69).³¹ Moreover, no significant differences were identified in hospital LOS, ICU LOS, or the risk of adverse events (Table 4). Overall, the increased point estimates in the risk of 30-day mortality, need for revascularization, and cardiac and renal adverse effects associated with a

restrictive transfusion strategy outweighed the potential benefits on lower rates of infection and thromboembolism.

606

607

608

609

610

611

612

613

614

615

616

617

618

619

620

621

622

623

624

625

626

627

628

629

630

631

632

633

634

636

637

638

639

640

641

642

643

644

645

646

647

648 649

650 651

653 654

655 656

657

658

659

660

Q35 652

What Others Are Saying

The ESICM guidelines, released before the publication of the MINT trial, suggested an RBC transfusion threshold of 9 to 10 g/dL in patients with ACS.^{20,37} However, given that it is unclear if a difference in outcomes occurs at higher hemoglobin levels, even in the range of a permissive approach, the panel opted to suggest against a restrictive approach.

Research Priorities

Studies exploring the impact on patient outcomes of a restrictive hemoglobin threshold of 8 g/dL or 9 g/dL are necessary. Further exploration of effect modifiers within the MINT trial may help to guide clinicians. Subgroup analysis, especially focused on type 1 MI vs 2 MI, as well as patients with heart failure and chronic kidney disease, are welcomed.

Question 4: Should critically ill patients undergoing cardiac surgery be treated with a restrictive or permissive RBC transfusion strategy?

4. In critically ill patients undergoing cardiac surgery, we suggest a restrictive RBC transfusion strategy over a permissive transfusion strategy in the perioperative 013 635 period (Conditional Recommendation, Moderate Certainty of Evidence).

Justification

This recommendation is substantiated by 7 high-quality clinical trials mostly of patients undergoing coronary artery bypass graft, valvular surgery, or both. 27-29,33-36 Despite the moderate certainty of evidence, a

TABLE 4 Pooled Analysis Comparing Restrictive vs Permissive Strategies in Patients With Acute Coronary Syndrome

Outcome	Relative Risk (95% CI)	Absolute Risk (95% CI)	
30-d mortality	1.13 (0.67-1.91)	10 more per 1,000 (26 fewer-73 more)	
Need for revascularization	1.09 (0.73-1.63)	2 more per 1,000 (6 fewer-13 more)	
Hospital length of stay		MD 0.02 d more (0.37 fewer-0.41 more)	
Adverse transfusion effects	1.87 (0.31-11.06)	1 more per 1,000 (1 fewer-10 more)	
Infections	0.87 (0.17-4.40)	10 fewer per 1,000 (61 fewer-252 more)	
Cardiac adverse events	1.16 (0.94-1.45)	10 more per 1,000 (4 fewer-29 more)	
Renal adverse events	1.06 (0.84-1.32)	7 more per 1,000 (20 fewer-39 more)	
Thromboembolism	0.75 (0.46-1.24)	5 fewer per 1,000 (10 fewer-5 more)	
Pulmonary adverse events	0.96 (0.77-1.19)	3 fewer per 1,000 (17 fewer-14 more)	

MD = mean difference.

conditional recommendation was issued because the studies showed serious imprecision in the point estimates for several outcomes.

661

662

663

664

665

666

667

668

669

670

671

672

673

674

675

676

677

678

679

680

681

682

683

684

685

686

687

688

689

690

691

692

693

694

695

696

697

698

699

700

701

702

703

704

705

706 707

708

709

710 711

712

713

714

715

Three studies examined 30-day mortality comparing both transfusion strategies in patients undergoing cardiac surgery. 27-29 Mazer et al 27 enrolled 5,243 patients, showing that a restrictive strategy was noninferior to a permissive approach and led to 20% less patients receiving RBC transfusions. Two smaller studies, although using slightly different transfusion thresholds, yielded similar outcomes. 28,29 In aggregate. no significant difference was found between the restrictive and permissive RBC transfusion threshold for 30-day mortality (RR, 1.12; 95% CI, 0.95-1.32). The hospital LOS did not differ between strategies (MD, -0.02 days; 95% CI, -0.19 to +0.15 days), whereas the ICU LOS favored the permissive strategy (MD, 0.12 days; 95% CI, 0.03-0.21 days), a finding without clinical significance. 27-29,33-36

One small study evaluated adverse transfusion reactions, but reported no events in either group. 36 No differences were found between the restrictive and permissive strategies in terms of infections, thromboembolism, or cardiac, renal, or pulmonary complications (Table 5).27-29,33-36

Overall, no important differences in outcomes or adverse events were noted between strategies among the 8,208 patients enrolled in the 7 trials. However, considering the 40% lower number of RBC units transfused to patients in the restrictive group, a restrictive strategy is the preferred approach.²⁸

What Others Are Saying

This recommendation aligned with those of other professional societies, including the ESICM and the Society of Thoracic Surgeons, which recommend a restrictive approach in this patient population.^{37,46}

716

717

718

719

720

721

722

723

724

727

728

729

730

731

732

733

734

735

736

737

738

739

740

741

750

756

757

758

770

Research Priorities

Further research may clarify the optimal RBC transfusion strategy for patients undergoing cardiac surgery. The TRICS-IV trial, currently underway, compares the two transfusion strategies in moderate to high-risk patients 65 years of age and younger to ensure 725 that the benefits of a restrictive strategy also apply to this 726 younger high-risk patient population.⁴⁷

Question 5: Should critically ill patients with an isolated elevation of serum troponin levels without other evidence of cardiac ischemia be treated with a restrictive or a permissive RBC transfusion strategy?

5. In critically ill patients with isolated elevation of serum troponin without other evidence of cardiac ischemia, we suggest a restrictive RBC transfusion strategy over a permissive RBC transfusion strategy (Conditional Recommendation, Very Low Certainty of Evidence).

Justification

No data are available regarding RBC transfusion 742 thresholds for critically ill patients with isolated elevated 743 troponin levels without evidence of cardiac ischemia, 744 defined as clinical symptoms, ECG changes, or both 745 746 consistent with ischemia.⁴⁸ Despite the absence of 747 evidence, the panel identified this as an important 748 clinical question and formulated a recommendation 749 using the collective experience approach recommended by the Grading of Recommendations, Assessment, 751 Development, and Evaluations criteria, considering 752 several factors. First, no universally agreed-on definition 753 for elevated troponin exists. This is attributable to the 754 availability of various troponin assays for clinical use, 755

TABLE 5 Pooled Analysis Comparing Restrictive vs Permissive Strategies in Patients Who Have Undergone Cardiac Surgery

Outcome	Relative Risk (95% CI)	Absolute Risk (95% CI)	— 75 — 76
30-d mortality	1.12 (0.95-1.32)	8 more per 1,000 (3 fewer-21 more)	70
ICU length of stay		MD 0.12 d more (0.03 more-0.21 more)	94
Hospital length of stay		MD 0.02 d lower (0.19 lower-0.15 higher)	76
Infections	1.07 (0.94-1.22)	6 more per 1,000 (5 fewer-19 more)	7
Cardiac adverse events	1.00 (0.75-1.32)	0 fewer per 1,000 (14 fewer-18 more)	7
Renal adverse events	1.03 (0.86-1.23)	2 more per 1,000 (7 fewer-12 more)	70
Thromboembolism	0.82 (0.36-1.88)	2 fewer per 1,000 (8 fewer-10 more)	7
Pulmonary adverse events	1.05 (0.89-1.24)	6 more per 1,000 (14 fewer-30 more)	7

MD = mean difference.

resulting in variability in both reported units and accepted normal troponin values. 49 Moreover, troponin levels represent a test value, rather than a specific clinical diagnosis, particularly in critically ill patients, in whom elevated troponin levels may indicate diverse conditions unrelated to ACS. 50-53 Considering the heterogeneity of conditions leading to elevated troponin levels in the absence of acute cardiac ischemia, the risk-benefit assessment regarding RBC transfusion strategy will depend on individual clinical circumstances. In general, we suggest adopting a restrictive RBC transfusion strategy as the first-line approach. However, the decision to transfuse should consider various clinical factors, including vasculopathy, intravascular volume status, troponin level and rate of rise, biventricular cardiac function, myocardial strain or trauma, renal dysfunction, sepsis, and surrogates of end-organ perfusion. In certain circumstances, clinicians may choose to transfuse RBCs to increase oxygen-carrying capacity.

What Others Are Saying

No recommendations from other professional societies have been published.

Question 6: Should critically ill patients with septic shock with end-organ hypoperfusion be treated with RBC transfusion in addition to usual care or usual care alone?

6. In patients with septic shock and end-organ hypoperfusion, we suggest against adding permissive RBC transfusion thresholds to usual care (Conditional Recommendation, Low Certainty of Evidence).

Remarks: Studies evaluating protocol driven approaches to goal-directed therapy in septic shock were not considered in the evidence review.

Justification

Sepsis care has evolved over the last 2 decades, with usual care including prompt antibiotic and fluid

administration and hemodynamic stabilization. Moreover, RBC transfusion practices in septic shock have shifted ^{54,55} toward a restrictive approach. ^{39,56,57} Consequently, the addition of RBC transfusions to usual care, aiming for higher hemoglobin levels, implies a permissive transfusion strategy, whereas usual care adheres to a restrictive approach. Studies evaluating RBC transfusions as part of a resuscitation bundle were not included, because the effect of transfusions could not be isolated from the effect of other bundle elements.

This recommendation is informed by 3 RCTs and long-term follow-up of an RCT cohort. 16,19,30,32 Two of the RCTs focused on general critically ill patients, whereas 1 RCT included patients with cancer, all of whom had septic shock, with higher RBC transfusion rates in the permissive groups compared with the restrictive groups.

TRISS, the largest study including > 1,000 patients, reported similar 30-day mortality rates in the permissive (35%) and restrictive (33%) approaches.³⁰ In patients with cancer, lower, albeit not statistically significant, 30day mortality (hazard ratio, 0.74; 95% CI, 0.53-1.04; P = .08) and ICU mortality rates (33.6% vs 43.7%; P = .071) were reported in the permissive compared with the restrictive group, 19 findings consistent with a smaller study in general critically ill patients. 16 The pooled analysis showed no significant difference between permissive and restrictive RBC transfusion strategies in terms of ICU mortality (RR, 0.84; 95% CI, 0.62-1.13) or 30-day mortality (RR, 0.93; 95% CI, 0.72-1.21). Longer-term follow-up at 1 year reported mortality rates of 55.8% in the permissive group and 53.5% in the restrictive cohort (RR, 0.97; 95% CI, 0.85-1.09; P = .62). No difference was found in terms of ICU and hospital LOS between both strategies.

Given that the studies evaluating adverse transfusion reactions were not powered a priori to detect such events, the number of events was very low. No strategy was superior regarding the need for renal replacement, cardiac, or pulmonary complications (Table 6). ^{19,30}

TABLE 6] Pooled Analysis Comparing Permissive vs Restrictive Strategies in Patients With Septic Shock

Outcome	Relative Risk (95% CI)	Absolute Risk (95% CI)	
ICU mortality	0.84 (0.62-1.12)	71 fewer per 1,000 (169 fewer-58 more)	
30-d mortality	0.93 (0.72-1.21)	27 fewer per 1,000 (108 fewer-81 more)	
Adverse transfusion reactions	0.33 (0.01-8.15)	1 fewer per 1,000 (2 fewer-11 more)	
Need for renal replacement therapy	0.98 (0.61-1.57)	2 fewer per 1,000 (34 fewer-50 more)	
Cardiac complications	0.60 (0.27-1.31)	11 fewer per 1,000 (19 fewer-8 more)	
Pulmonary complications	1.05 (0.60-1.82)	10 more per 1,000 (77 fewer-159 more)	

ARTICLE IN PRESS

Overall, although the data show that the permissive strategy might result in benefit, it does not exclude possible harm because the RR crosses the null threshold. In the absence of clear benefit and with similar rates of adverse effects, neither strategy is deemed clinically favorable. However, a restrictive approach results in fewer RBC units transfused, optimizing resource use and decreasing costs.30

What Others Are Saying

881

882

883

884

885

886

887

888

889

890

891

892

893

894

895

896

897

898

899

900

901

902

903

904

905

906

907

908

909

910

911

912

913

914

915

916

917

918

919

920

921

922

923

924

925

926

927

928

929

930

931

932

933

934

935

The recommendation to not add a permissive RBC transfusion strategy to usual care aligns with the most recent ESICM and the Surviving Sepsis Campaign guidelines.37,58

Research Priorities

Future studies including populations typically underrepresented in sepsis trials, such as patients with malignancies (solid and hematologic), liver disease, and chronic coronary disease, are welcome. Additionally, further evidence is needed regarding the benefits of RBC transfusions in severe hypoxemia or tissue hypoperfusion.

Additional Considerations

The statements on cost, equity, acceptability, feasibility, and implementation were consolidated, given the significant overlap across various Population, Intervention, Comparator, and Outcome questions.

Cost

Economic considerations are important when deciding on RBC transfusion strategies for critically ill patients. In the United States, the hospital cost of an RBC unit is approximately \$207,59 with administrative, logistic, and labor costs totaling up to \$1,183.60,61 A restrictive strategy spares 36% of patients from RBC transfusions, reduces the number of RBC units transfused by 50%, and reduces costs by 33% compared with a permissive approach.^{28,30}

Cost-effectiveness analysis of RBC transfusions in patients undergoing cardiac surgery revealed that expenses from surgery to the third postoperative month were slightly higher in the permissive group than in the restrictive group, driven by RBC costs. However, the differences in quality-adjusted life-years were negligible. Therefore, the restrictive strategy in these patients was considered cost-effective.⁶²

Although a definitive evaluation of cost-effectiveness is challenging because of limited studies in the

critically ill population, a restrictive approach is deemed cost-effective overall and among most subpopulations because no significant impact on overall mortality in any major subgroups was observed, except for ACS. Therefore, in the general critically ill population, in the absence of benefit resulting from a permissive strategy, a restrictive strategy would be considered a preferred option under most circumstances.

936

937

938

939

940

941

942

943

944

945

946

947

949

951

952

953

954

955

956

957

959

960

961

964

965

966

967

968

969

970

971

972

973

974

975

977

978

979

980

981

982

983

984

985

986

987

988

989

Equity

Access to RBC transfusions is influenced by geographic 948 location and resource allocation. A restrictive RBC transfusion strategy can reduce inequity by ensuring that 950 individuals most in need of RBCs receive them. This may be even more relevant during major blood shortages or in rural areas with limited resources. Moreover, in countries with higher rates of blood-borne infections, a restrictive approach will reduce exposure. However, the impact of provider biases on transfusion practices in settings of limited RBC availability remains 958 unclear.

Acceptability and Feasibility

The evidence suggested that most practitioners and 962 centers have adopted a restrictive RBC transfusion strategy, indicating stakeholder acceptability.⁶³ However, some patients may reject transfusions based on personal values or religious beliefs.⁶⁴ The panel believed that implementing restrictive transfusion strategies is feasible through behavior modification interventions, including education, institutional guidelines, and audit and feedback. Furthermore, additional blood conservation strategies, such as reduced laboratory testing, optimization of perioperative antiplatelet and anticoagulation regimens, intraoperative blood conservation, and small-volume blood sampling, are 976 crucial. These interventions have demonstrated effectiveness in reducing the odds of transfusion, inappropriate transfusion rates, and the number of RBC units transfused per patient. 65-67

Implementation

For patients reluctant to accept transfusions based on personal values or religious beliefs, a thorough discussion with the patient or surrogate should occur before deciding to transfuse. As soon as an RBC transfusion is decided on, the optimal implementation strategy encompasses a restrictive approach and transfusing 1 RBC unit at a time. This

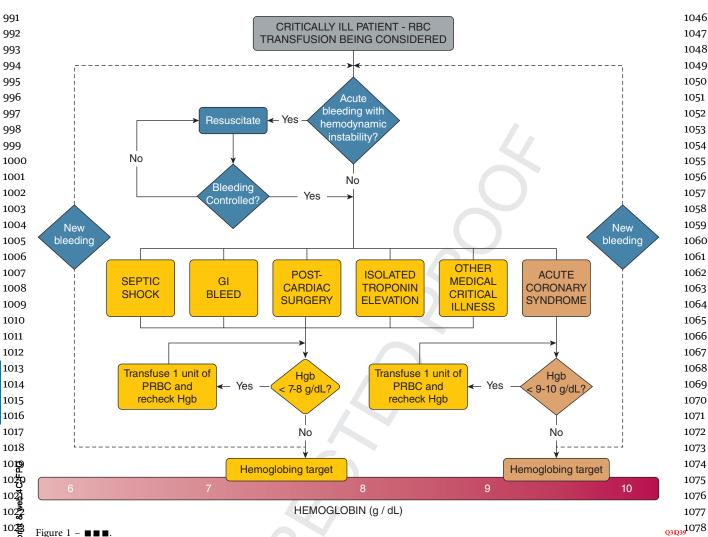


Figure 1 - ■ ■ ■

threshold and single-unit recommendation does not apply to patients actively bleeding at a rate exceeding the ability to transfuse single units or await hemoglobin test results safely. The recommendation can be applied to patients whose acute bleeding has subsided. If acute bleeding occurs during a patient's stay, this approach should be suspended and then reapplied after bleeding is controlled (Fig 1).

The panel suggested conducting audits or observational studies using hospital databases to understand current practices. Proven implementation strategies to overcome barriers to changing transfusion include academic detailing, audit-feedback approaches, standard order sets, computerized order entry decision support, reminders, and alerts. These approaches require resources, leadership, and clinical oversight.⁶⁸⁻⁷²

Summary

In the United States, approximately 5 million patients are admitted to the ICU annually, and one-quarter of them receive RBC transfusions during their stay. 1-3,73 Most clinical trials support a restrictive transfusion strategy, showing no significant differences in mortality or adverse outcomes overall and in all patient subgroups, except for critically ill patients with ACS. 20,22,23 Adopting a restrictive strategy could decrease the number of patients receiving RBC transfusions by approximately 40%.38 On a large scale, this could represent sparing 0.5 million patients from RBC transfusions.

A limitation of this guideline is the quality of the evidence, which ranged from moderate to very low. For conditions like ACS and septic shock, the number of studies was small, and the inferences

were not strong. No studies specifically addressed critically ill patients with elevated troponin levels. Moreover, the studies did not assess additional aspects of the transfusion process (storage age, donor characteristics, processing, storage solutions). These guidelines provide an opportunity for institutions to develop local policies, monitor their impact on transfusion practices, and to create a framework to longitudinally optimize RBC use.

Funding/Support

1101

1102

1103

1104

1105

1106

1107

1108

1109

1110

1111

1113

1114

1115

1116

1118

1119

1120

1121

1122

1123

1125

1126

1127^{Q20}

1128

1129

1130

1131

1133

1134

1135

1136

1137

1138

1139

1140

1141

1143

1144

1145

1146

1147

1148

1149

1150

1151

1152

1153

1154

1155

1142222

1132^{Q21}

1124219

1117_{Q18}

The authors have reported to CHEST that no funding was received for this study.

Financial/Nonfinancial Disclosures

None declared.

Acknowledgments

Author contributions: A. C. Y., I. S., and D. R. O had full access to all the data in the study and take full responsibility for the integrity of the data and the accuracy of the data analysis. P. C. H., E. B.-C., M. D., P.F., K. H., J. M. I., W. L., C. C. L., P. J. M., K. O., K. M. P., and L. V. V. contributed substantially to the study design, data interpretation, and writing of the manuscript.

Additional information: The e-Appendix, e-Figures, and e-Tables are available online under "Supplementary Data."

References

- 1. Vincent JL, Jaschinski U, Wittebole X, et al. Worldwide audit of blood transfusion practice in critically ill patients. Crit Care. 2018;22(1):102.
- 2. Free RJ, Sapiano MRP, Chavez Ortiz JL, Stewart P, Berger J, Basavaraju SV. Continued stabilization of blood collections and transfusions in the United States: Findings from the 2021 National Blood Collection and Utilization Survey. Transfusion. 2023;63(suppl 4):S8-S18.
- 3. Raasveld SJ, de Bruin S, Reuland MC, et al. Red blood cell transfusion in the intensive care unit. JAMA. 2023;330(19): 1852-1861.
- 4. United Kingdom Blood Services. Effective transfusion in surgery and critical care. In: Handbook of Transfusion Medicine. 5th ed. United Kingdom Blood Services website. https://www.transfusionguidelines. org/transfusion-handbook/7-effective-transfusion-in-surgery-and-
- 5. Hebert PC, Wells G, Blajchman MA, et al. A multicenter, randomized, controlled clinical trial of transfusion requirements in critical care. Transfusion Requirements in Critical Care Investigators, Canadian Critical Care Trials Group. N Engl J Med. 1999;340(6):409-417.
- 6. Rohde JM, Dimcheff DE, Blumberg N, et al. Health care-associated infection after red blood cell transfusion: a systematic review and meta-analysis. JAMA. 2014;311(13):1317-1326.
- 7. Jones JM, Sapiano MRP, Mowla S, Bota D, Berger JJ, Basavaraju SV. Has the trend of declining blood transfusions in the United States ended? Findings of the 2019 National Blood Collection and Utilization Survey. Transfusion. 2021;61(suppl 2):
- 8. American Red Cross. Blood shortage continues, please give blood now. American Red Cross website. Accessed October 27, 2023. https://www.redcross.org/about-us/news-and-events/press-release/2023/red-cross-announces-national-blood-shortage.html#:

~:text=When%20supplies%20are%20low%20it,meet%20hospital%
20and%20patient%20needs

1156

1157

1158

1159

9760

1161

1162

42163

1166

9467

1168

d2k70

1172

1173

1174

1175

1176

1177

1178

1183

1184

1185

1186

1187

1189

1190

1191

1192

1193

1194

1195

1196

1197

1198

1199

1200

1201

1202

1203

1204

1207

1210

- 9. Lotterman S, Sharma S. Blood transfusion. StatPearls. StatPearls Publishing; 2023. Updated June 20, 2023. National Library of Medicine website. https://www.ncbi.nlm.nih.gov/books/NBK4
- 10. American College of Chest Physicians (CHEST). Chest guideline development and submissions. American College of Chest Physicians website. https://www.chestnet.org/guidelines-and-topiccollections/guidelines/topic-submissions
- 11. Boutron, et al. Considering bias and conflicts of interest among the 1164 included studies. In: Cochrane Handbook for Systematic Reviews of 1165 Interventions. Version 6.3. 2022. The Cochrane Collaboration website. Accessed November, 2022, https://training.cochrane.org/ handbook/current/chapter-07
- 12. Higgins, et al. Assessing risk of bias in a randomized trial. In: Cochrane Handbook for Systematic Reviews of Interventions. Version 1169 6.3. 2022. The Cochrane Collaboration website. Accessed November, 2022, https://training.cochrane.org/handbook/current/chapter-08
- 13. Balshem H, Helfand M, Schunemann HJ, et al. GRADE guidelines: 1171 3. Rating the quality of evidence. J Clin Epidemiol. 2011;64(4): 401-406.
- 14. Andrews JC, Schunemann HJ, Oxman AD, et al. GRADE guidelines: 15. Going from evidence to recommendation-determinants of a recommendation's direction and strength. J Clin Epidemiol. 2013;66(7):726-735.
- 15. Jaeschke R, Guyatt GH, Dellinger P, et al. Use of GRADE grid to reach decisions on clinical practice guidelines when consensus is elusive. BMJ. 2008;337:a744.
- 1179 16. Mazza BF, Freitas FG, Barros MM, Azevedo LC, Machado FR. Blood transfusions in septic shock: is 7.0 g/dL really the 1180 appropriate threshold? Rev Bras Ter Intensiva. 2015;27(1):36-43. 1181
- 17. Hebert PC, Wells G, Marshall J, et al. Transfusion requirements in 1182 critical care. A pilot study. Canadian Critical Care Trials Group. JAMA. 1995;273(18):1439-1444.
- 18. Walsh TS, Boyd JA, Watson D, et al. Restrictive versus liberal transfusion strategies for older mechanically ventilated critically ill patients: a randomized pilot trial. Crit Care Med. 2013;41(10): 2354-2363.
- 19. Bergamin FS, Almeida JP, Landoni G, et al. Liberal versus restrictive transfusion strategy in critically ill oncologic patients: the transfusion $\,$ 1188 requirements in critically ill oncologic patients randomized controlled trial. Crit Care Med. 2017;45(5):766-773.
- 20. Carson JL, Brooks MM, Hebert PC, et al. Restrictive or liberal transfusion strategy in myocardial infarction and anemia. N Engl J Med. 2023;389(26):2446-2456.
- 21. Cooper HA, Rao SV, Greenberg MD, et al. Conservative versus liberal red cell transfusion in acute myocardial infarction (the CRIT Randomized Pilot Study). Am J Cardiol. 2011;108(8): 1108-1111.
- 22. Carson JL, Brooks MM, Abbott JD, et al. Liberal versus restrictive transfusion thresholds for patients with symptomatic coronary artery disease. Am Heart J. 2013;165(6):964-971.e1.
- 23. Ducrocq G, Gonzalez-Juanatey JR, Puymirat E, et al. Effect of a restrictive vs liberal blood transfusion strategy on major cardiovascular events among patients with acute myocardial infarction and anemia: the REALITY randomized clinical trial. JAMA. 2021;325(6):552-560.
- 24. Villanueva C, Colomo A, Bosch A, et al. Transfusion strategies for acute upper gastrointestinal bleeding. N Engl J Med. 2013;368(1):
- 25. Kola G, Sureshkumar S, Mohsina S, Sreenath GS, Kate V. Restrictive 1205 versus liberal transfusion strategy in upper gastrointestinal bleeding: 1206 a randomized controlled trial. Saudi J Gastroenterol. 2021;27(1):
- 26. Jairath V, Kahan BC, Gray A, et al. Restrictive versus liberal blood 1208 transfusion for acute upper gastrointestinal bleeding (TRIGGER): a 1209 pragmatic, open-label, cluster randomised feasibility trial. Lancet. 2015;386(9989):137-144.

ARTICLE IN PRESS

- 1211 27. Mazer CD, Whitlock RP, Fergusson DA, et al. Restrictive or liberal
 1212 red-cell transfusion for cardiac surgery. N Engl J Med. 2017;377(22):
 2133-2144
- 1213
 28. Murphy GJ, Pike K, Rogers CA, et al. Liberal or restrictive transfusion after cardiac surgery. N Engl J Med. 2015;372(11): 997-1008.
- 1216 29. Hajjar LA, Vincent JL, Galas FR, et al. Transfusion requirements after cardiac surgery: the TRACS randomized controlled trial. *JAMA*. 2010;304(14):1559-1567.
- 30. Holst LB, Haase N, Wetterslev J, et al. Lower versus higher hemoglobin threshold for transfusion in septic shock. N Engl J Med. 2014;371(15):1381-1391.
- 31. Gonzalez-Juanatey JR, Lemesle G, Puymirat E, et al. One-year major cardiovascular events after restrictive versus liberal blood transfusion strategy in patients with acute myocardial infarction and anemia: the REALITY randomized trial. *Circulation*. 2022;145(6):486-488.
- 32. Rygard SL, Holst LB, Wetterslev J, et al. Long-term outcomes in patients with septic shock transfused at a lower versus a higher haemoglobin threshold: the TRISS randomised, multicentre clinical trial. *Intensive Care Med.* 2016;42(11):1685-1694.
- 33. Koch CG, Sessler DI, Mascha EJ, et al. A randomized clinical trial of red blood cell transfusion triggers in cardiac surgery. *Ann Thorac Surg.* 2017;104(4):1243-1250.
- 1230 34. Laine A, Niemi T, Schramko A. Transfusion threshold of
 1231 hemoglobin 80 g/L is comparable to 100 g/L in terms of bleeding in cardiac surgery: a prospective randomized study. J Cardiothorac Vasc Anesth. 2018;32(1):131-139.
- 35. Bracey AW, Radovancevic R, Riggs SA, et al. Lowering the hemoglobin threshold for transfusion in coronary artery bypass procedures: effect on patient outcome. *Transfusion*. 1999;39(10): 1070-1077.
- 36. Shehata N, Burns LA, Nathan H, et al. A randomized controlled pilot study of adherence to transfusion strategies in cardiac surgery.
 Transfusion. 2012;52(1):91-99.
- 1239 37. Vlaar AP, Oczkowski S, de Bruin S, et al. Transfusion strategies in non-bleeding critically ill adults: a clinical practice guideline from the European Society of Intensive Care Medicine. *Intensive Care Med.* 2020;46(4):673-696.
- 1242 38. Carson JL, Stanworth SJ, Guyatt G, et al. Red blood cell transfusion:
 1243 2023 AABB international guidelines. *JAMA*. 2023;330(19):
 1244 1892-1902.
- 1245
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 1246
 <li
- 1247 40. Kherad O, Restellini S, Martel M, et al. Outcomes following restrictive or liberal red blood cell transfusion in patients with lower gastrointestinal bleeding. *Aliment Pharmacol Ther*. 2019;49(7): 919-925.
- 41. Garcia-Tsao G, Abraldes JG, Berzigotti A, Bosch J. Portal
 hypertensive bleeding in cirrhosis: risk stratification, diagnosis, and management: 2016 practice guidance by the American Association for the study of liver diseases. *Hepatology*. 2017;65(1):310-335.
- 1254 42. Laine L, Barkun AN, Saltzman JR, Martel M, Leontiadis GI. ACG clinical guideline: upper gastrointestinal and ulcer bleeding. Am J Gastroenterol. 2021;116(5):899-917.
- 43. Kaplan DE, Ripoll C, Thiele M, et al. AASLD practice guidance on risk stratification and management of portal hypertension and varices in cirrhosis. *Hepatology*. 2023.
- 1259 44. Henry Z, Patel K, Patton H, Saad W. AGA clinical practice update on management of bleeding gastric varices: expert review. Clin Gastroenterol Hepatol. 2021;19(6):1098-1107.e1.
- 45. Roubinian NH, Carson JL. Acute myocardial infarction and blood transfusion: lessons learned from animal models and clinical studies.
 Blood Transfus. 2023;21(3):185-188.
- 1264 46. Tibi P, McClure RS, Huang J, et al. STS/SCA/AmSECT/SABM update to the clinical practice guidelines on patient blood management. *J Extra Corpor Technol.* 2021;53(2):97-124.

 National Institutes of Health Clinical Center. Transfusion requirements in younger patients undergoing cardiac surgery (TRICS-IV). NCT04754022. ClinicalTrials.gov. National Institutes of Health. https://clinicaltrials.gov/study/NCT04754022. 1266

1267

1268

1269

1270

1271

1272

1273

1274

1275

1276

1277

1278

1279

1280

1281

1282

1283

1284

1285

1286

1287

1288

1289

1290

1291

1292

1293

1294

1295

1296

1297

1298

1299

1300

1301

1302

1303

1304

1305

1306

1307

1308

1309

1310

1311

1312

1313

1314

1315

1316

1317

1318

1319

1320

- Thygesen K, Alpert JS, Jaffe AS, et al. Fourth universal definition of myocardial infarction (2018). Circulation. 2018;138(20):e618-e651.
- Lan NSR, Bell DA. Revisiting the biological variability of cardiac troponin: implications for clinical practice. *Clin Biochem Rev*. 2019;40(4):201-216.
- Roongsritong C, Warraich I, Bradley C. Common causes of troponin elevations in the absence of acute myocardial infarction: incidence and clinical significance. *Chest.* 2004;125(5):1877-1884.
- Klein Gunnewiek JMT, van de Leur JJJPM. Elevated troponin T concentrations in critically ill patients. *Intensive Care Med*. 2003;29(12):2317-2322.
- Kelley WE, Januzzi JL, Christenson RH. Increases of cardiac troponin in conditions other than acute coronary syndrome and heart failure. Clin Chem. 2009;55(12):2098-2112.
- 53. Belley-Cote EP, Whitlock RP, Ulic DV, et al. The PROTROPIC feasibility study: prognostic value of elevated troponins in critical illness. *Can J Anaesth.* 2019;66(6):648-657.
- Dellinger RP, Carlet JM, Masur H, et al. Surviving Sepsis Campaign guidelines for management of severe sepsis and septic shock. Crit Care Med. 2004;32(3):858-873.
- Rhodes A, Evans LE, Alhazzani W, et al. Surviving Sepsis Campaign: international guidelines for management of sepsis and septic shock: 2016. Crit Care Med. 2017;45(3).
- Dixit S, Khatib KI. Blood transfusion practices in sepsis. In: Prabhakar H, Tandon MS, Kapoor I, Mahajan C, eds. *Transfusion Practice in Clinical Neurosciences*. Singapore: Springer Nature; 2022: 541-546.
- Muady GF, Bitterman H, Laor A, Vardi M, Urin V, Ghanem-Zoubi N. Hemoglobin levels and blood transfusion in patients with sepsis in internal medicine departments. *BMC Infect Dis.* 2016;16(1): 569.
- Evans L, Rhodes A, Alhazzani W, et al. Surviving Sepsis Campaign: international guidelines for management of sepsis and septic shock 2021. Crit Care Med. 2021;49(11).
- Mowla SJ, Sapiano MRP, Jones JM, Berger JJ, Basavaraju SV. Supplemental findings of the 2019 National Blood Collection and Utilization Survey. *Transfusion*. 2021;61(suppl 2):S11-S35.
- 60. Spahn DR, Goodnough LT. Alternatives to blood transfusion. *Lancet*. 2013;381(9880):1855-1865.
- Shander A, Hofmann A, Ozawa S, Theusinger OM, Gombotz H, Spahn DR. Activity-based costs of blood transfusions in surgical patients at four hospitals. *Transfusion*. 2010;50(4): 753-765.
- 62. Reeves BC, Pike K, Rogers CA, et al. A multicentre randomised controlled trial of Transfusion Indication Threshold Reduction on transfusion rates, morbidity and health-care resource use following cardiac surgery (TITRe2). *Health Technol Assess*. 2016;20(60):1-260.
- Soril LJJ, Noseworthy TW, Stelfox HT, Zygun DA, Clement FM. Facilitators of and barriers to adopting a restrictive red blood cell transfusion practice: a population-based cross-sectional survey. CMAJ Open. 2019;7(2):E252-E257.
- 64. Berend K, Levi M. Management of adult Jehovah's Witness patients with acute bleeding. *Am J Med.* 2009;122(12):1071-1076.
- 65. Soril LJJ, Noseworthy TW, Dowsett LE, et al. Behaviour modification interventions to optimise red blood cell transfusion practices: a systematic review and meta-analysis. *BMJ Open*. 2018;8(5):e019912.
- 66. Grau JB, Fortier JH, Kuschner C, et al. Implementing a protocol to optimize blood use in a cardiac surgery service: results of a pre-post analysis and the impact of high-volume blood users. *Transfusion*. 2017;57(10):2483-2489.
- Siegal DM, Belley-Cote EP, Lee SF, et al. Small-volume blood collection tubes to reduce transfusions in intensive care: the STRATUS randomized clinical trial. *JAMA*. 2023;330(19):1872-1881.

68. Gupta M, Ranapurwala MF. Clinical audits—a quality improvement tool in transfusion medicine. Global Journal of Transfusion Medicine. 2022;7(2):115-122.

- 69. Jagathkar G, Samavedam S. How to audit transfusion practices in the intensive care unit? Indian J Crit Care Med. 2019;23(suppl 3):
- 70. Adams ES, Longhurst CA, Pageler N, Widen E, Franzon D, Cornfield DN. Computerized physician order entry with decision support decreases blood transfusions in children. Pediatrics. 2011;127(5):e1112-e1119.
- 71. Shiekh Sroujieh L, Monroy D, Warren E. Using electronic health 1376 record (EHR) best practice alert (BPA) to improve RBC transfusion practices and adherence to the guidelines. Chest. 2016;150(4):599A.
- 72. Crispin P, Akers C, Brown K, et al. A review of electronic medical 1379 records and safe transfusion practice for guideline development. Vox 1380 Sang. 2022;117(6):761-768.
- 73. Viglianti EM, Iwashyna TJ. Toward the ideal ratio of patients to intensivists: finding a reasonable balance. JAMA Intern Med. 2017;177(3):396-398.