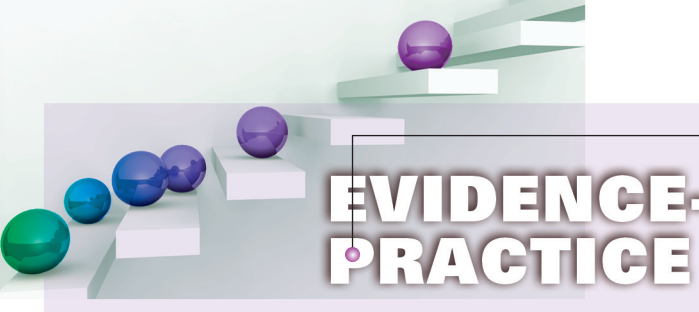




EVIDENCE-BASED PRACTICE

Step by Step

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Critical Appraisal of the Evidence: Part III

The process of synthesis: seeing similarities and differences across the body of evidence.

This is the seventh article in a series from the Arizona State University College of Nursing and Health Innovation's Center for the Advancement of Evidence-Based Practice. Evidence-based practice (EBP) is a problem-solving approach to the delivery of health care that integrates the best evidence from studies and patient care data with clinician expertise and patient preferences and values. When delivered in a context of caring and in a supportive organizational culture, the highest quality of care and best patient outcomes can be achieved.

The purpose of this series is to give nurses the knowledge and skills they need to implement EBP consistently, one step at a time. Articles will appear every two months to allow you time to incorporate information as you work toward implementing EBP at your institution. Also, we've scheduled "Chat with the Authors" calls every few months to provide a direct line to the experts to help you resolve questions. See details below.

In September's evidence-based practice (EBP) article, Rebecca R., our hypothetical staff nurse, Carlos A., her hospital's expert EBP mentor, and Chen M., Rebecca's nurse colleague, rapidly critically appraised the 15 articles they found to answer their clinical question—"In *hospitalized adults* (P), how does a *rapid response team* (I) compared with *no rapid response team* (C) affect the *number of cardiac arrests* (O) and *unplanned admissions to the ICU* (O) during a *three-month period* (T):"—and determined that they were all "keepers." The team now begins the process of evaluation and synthesis of the articles to see what the evidence says about initiating a rapid response team (RRT) in their hospital. Carlos reminds them that evaluation and synthesis are synergistic processes and don't necessarily happen one after the other. Nevertheless, to help them learn, he will guide them through the EBP process one step at a time.

STARTING THE EVALUATION

Rebecca, Carlos, and Chen begin to work with the evaluation table

they created earlier in this process when they found and filled in the essential elements of the 15 studies and projects (see "Critical Appraisal of the Evidence: Part I," July). Now each takes a stack of the "keeper" studies and systematically begins adding to the table any remaining data that best reflect the study elements pertaining to the group's clinical question (see Table 1; for the entire table with all 15 articles, go to <http://links.lww.com/AJN/A17>). They had agreed that a "Notes" section within the "Appraisal: Worth to Practice" column would be a good place to record the nuances

of an article, their impressions of it, as well as any tips—such as what worked in calling an RRT—that could be used later when they write up their ideas for initiating an RRT at their hospital, if the evidence points in that direction. Chen remarks that although she thought their initial table contained a lot of information, this final version is more thorough by far. She appreciates the opportunity to go back and confirm her original understanding of the study essentials.

The team members discuss the evolving patterns as they complete the table. The three systematic

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Go to www.ajnonline.com and click on "Podcasts" and then on "Conversations" to listen to our interview with Ellen Fineout-Overholt and Bernadette Mazurek Melnyk.

Table 1. Final Evaluation Table

First Author (Year)	Conceptual Framework	Design/Method	Sample/Setting	Major Variables Studied (and Their Definitions)	Measurement	Data Analysis	Findings	Appraisal: Worth to Practice
Chan PS, et al. <i>Arch Intern Med</i> 2010;170(1):18-26	None	SR Purpose: effect of RRT on HMR and CR - Searched 5 databases from 1950–2008 and “grey literature” from MD conferences - Included only 1) RCTs and prospective studies with 2) a control group or control period and 3) hospital mortality well described as outcome - Excluded 5 studies that met criteria due to no response to e-mail by primary authors	N = 18 out of 143 potential studies Setting: acute care hospitals; 13 adult, 5 peds Average no. beds: NR Attrition: NR	IV: RRT DV1: HMR (including DNR, excluding DNR, not treated in ICU, no HMR definition) DV2: CR	RRT: was the MD involved? HMR: overall hospital deaths (see definition) CR: cardio and/or pulmonary arrest; cardiac arrest calls	- Frequency - Relative risk	13/16 studies reporting team structure 7/11 adult and 4/5 peds studies had significant reduction in CR CR: - In adults, 21%–48% reduction in CR; RR 0.66 [95% CI, 0.54–0.80] - In peds, 38% reduction in CR; RR 0.62 [95% CI, 0.46–0.84] HMR: - In adults, HMR RR 0.96 [95% CI, 0.84–1.09] - In peds, HMR RR 0.79 [95% CI, 0.63–0.98]	Weaknesses: - Potential missed evidence with exclusion of all studies except those with control groups - Grey literature search limited to medical meetings - Only included HMR and CR outcomes - No cost data Strengths: - Identified no. of activations of RRT/1,000 admissions - Identified variance in outcome definition and measurement (for example, 10 of 15 studies included deaths from DNRs in their mortality measurement) Conclusion: - RRT reduces CR in adults, and CR and HMR in peds Feasibility: - RRT is reasonable to implement; evaluating cost will help in making decisions about using RRT - Risk/Benefit (harm): benefits outweigh risks

McGaughy J, et al. <i>Cochrane Database Syst Rev</i> 2007;3: CD005529	None	SR (Cochrane review) Purpose: effect of RRT on HMR • Searched 6 databases from 1990–2006 • Excluded all but 2 RCTs	N = 2 studies Acute care settings in Australia and the UK Attrition: NR	IV: RRT DV1: HMR	HMR: Australia: overall hospital mortality without DNR UK: Simplified Acute Physiology Score (SAPS) II death probability estimate	OR	OR of Australian study, 0.98 (95% CI, 0.83–1.16) OR of UK study, 0.52 (95% CI, 0.32–0.85)	Weaknesses: • Didn't include full body of evidence • Conflicting results of retained studies, but no discussion of the impact of lower-level evidence • Recommendation "need more research" Conclusion: • Inconclusive
Winters BD, et al. <i>Crit Care Med</i> 2007;35(5): 1238–43	None	SR Purpose: effect of RRT on HMR and CR • Searched 3 databases from 1990–2005 • Included only studies with a control group	N = 8 studies Average no. beds: 500 Attrition: NR	IV: RRT DV1: HMR DV2: CR	HMR: overall death rate CR: no. of in-hospital arrests	Risk ratio	HMR: • Observational studies, risk ratio for RRT on HMR, 0.87 (95% CI, 0.73–1.04) • Cluster RCTs, risk ratio for RRT on HMR, 0.76 (95% CI, 0.39–1.48) CR: • Observational studies, risk ratio for RRT on CR, 0.70 (95% CI, 0.56–0.92) • Cluster RCTs, risk ratio for RRT on CR, 0.94 (95% CI, 0.79–1.13)	Strengths: • Provides comparison across studies for ○ Study lengths (range, 4–82 months) ○ Sample size (range, 2,183–199,024) ○ Criteria for RRT initiation (common: respiratory rate, heart rate, blood pressure, mental status change; not all studies, but noteworthy: oxygen saturation, "worry") • Includes ideas about future evidence generation (conducting research)—finding out what we don't know Conclusion: • Some support for RRT, but not reliable enough to recommend as standard of care

CI = confidence interval; CR = cardiopulmonary arrest or code rates; DNR = do not resuscitate; DV = dependent variable; HMR = hospital-wide mortality rates; ICU = intensive care unit; IV = independent variable; MD = medical doctor; NR = not reported; OR = odds ratio; Peds = pediatrics; RCT = randomized controlled trial; RR = relative risk; RRT = rapid response team; SR = systematic review; UK = United Kingdom

reviews, which are higher-level evidence, seem to have an inherent bias in that they included only studies with control groups. In general, these studies weren't in favor of initiating an RRT. Carlos asks Rebecca and Chen whether,

Chen in their efforts to appraise the MERIT study and comments on how well they're putting the pieces of the evidence puzzle together. The nurses are excited that they're able to use their new knowledge to shed light on the

as well as a good number of journals have encouraged their use. When they review the actual guidelines, the team notices that they seem to be focused on research; for example, they require a research question and refer to

It's not the number of studies or projects that determines the reliability of their findings, but the uniformity and quality of their methods.

now that they've appraised all the evidence about RRTs, they're confident in their decision to include all the studies and projects (including the lower-level evidence) among the "keepers." The nurses reply with an emphatic affirmative! They tell Carlos that the projects and descriptive studies were what brought the issue to life for them. They realize that the higher-level evidence is somewhat in conflict with the lower-level evidence, but they're most interested in the conclusions that can be drawn from considering the entire body of evidence.

Rebecca and Chen admit they have issues with the systematic reviews, all of which include the MERIT study.¹⁴ In particular, they discuss how the authors of the systematic reviews made sure to report the MERIT study's finding that the RRT had no effect, but didn't emphasize the MERIT study authors' discussion about how their study methods may have influenced the reliability of the findings (for more, see "Critical Appraisal of the Evidence: Part II," September). Carlos says that this is an excellent observation. He also reminds the team that clinicians may read a systematic review for the conclusion and never consider the original studies. He encourages Rebecca and

study. They discuss with Carlos how the interpretation of the MERIT study has perhaps contributed to a misunderstanding of the impact of RRTs.

Comparing the evidence. As the team enters the lower-level evidence into the evaluation table, they note that it's challenging to compare the project reports with studies that have clearly described methodology, measurement, analysis, and findings. Chen remarks that she wishes researchers and clinicians would write study and project reports similarly. Although each of the studies has a process or method determining how it was conducted, as well as how outcomes were measured, data were analyzed, and results interpreted, comparing the studies as they're currently written adds another layer of complexity to the evaluation. Carlos says that while it would be great to have studies and projects written in a similar format so they're easier to compare, that's unlikely to happen. But he tells the team not to lose all hope, as a format has been developed for reporting quality improvement initiatives called the SQUIRE Guidelines; however, they aren't ideal. The team looks up the guidelines online (www.squire-statement.org) and finds that the Institute for Healthcare Improvement (IHI)

the study of an intervention, whereas EBP projects have PICOT questions and apply evidence to practice. The team discusses that these guidelines can be confusing to the clinicians authoring the reports on their projects. In addition, they note that there's no mention of the synthesis of the body of evidence that should drive an evidence-based project. While the SQUIRE Guidelines are a step in the right direction for the future, Carlos, Rebecca, and Chen conclude that, for now, they'll need to learn to read these studies as they find them—looking carefully for the details that inform their clinical question.

Once the data have been entered into the table, Carlos suggests that they take each column, one by one, and note the similarities and differences across the studies and projects. After they've briefly looked over the columns, he asks the team which ones they think they should focus on to answer their question. Rebecca and Chen choose "Design/Method," "Sample/Setting," "Findings," and "Appraisal: Worth to Practice" (see Table 1) as the initial ones to consider. Carlos agrees that these are the columns in which they're most likely to find the most pertinent information for their synthesis.

SYNTHESIZING: MAKING DECISIONS BASED ON THE EVIDENCE

Design/Method. The team starts with the “Design/Method” column because Carlos reminds them that it’s important to note each study’s level of evidence. He suggests that they take this information and create a synthesis table (one in which data is extracted from the evaluation table to better see the similarities and differences between studies) (see Table 2¹⁻¹⁵). The synthesis table makes it clear that there is less higher-level and more lower-level evidence, which will impact the reliability of the overall findings. As the team noted, the higher-level evidence is not without methodological issues, which will increase the challenge of coming to a conclusion about

the impact of an RRT on the outcomes.

Sample/Setting. In reviewing the “Sample/Setting” column, the group notes that the number of hospital beds ranged from 218 to 662 across the studies. There were several types of hospitals represented (4 teaching, 4 community, 4 no mention, 2 acute care hospitals, and 1 public hospital). The evidence they’ve collected seems applicable, since their hospital is a community hospital.

Findings. To help the team better discuss the evidence, Carlos suggests that they refer to all projects or studies as “the body of evidence.” They don’t want to get confused by calling them all studies, as they aren’t, but at the

same time continually referring to “studies and projects” is cumbersome. He goes on to say that, as part of the synthesis process, it’s important for the group to determine the overall impact of the intervention across the body of evidence. He helps them create a second synthesis table containing the findings of each study or project (see Table 3¹⁻¹⁵). As they look over the results, Rebecca and Chen note that RRTs reduce code rates, particularly outside the ICU, whereas unplanned ICU admissions (UICUA) don’t seem to be as affected by them. However, 10 of the 15 studies and projects reviewed didn’t evaluate this outcome, so it may not be fair to write it off just yet.

Table 2: The 15 Studies: Levels and Types of Evidence

	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15
Level I: Systematic review or meta-analysis	X	X	X												
Level II: Randomized controlled trial				X											
Level III: Controlled trial without randomization															
Level IV: Case-control or cohort study					X	X									
Level V: Systematic review of qualitative or descriptive studies															
Level VI: Qualitative or descriptive study (includes evidence implementation projects)							X	X	X	X	X	X	X	X	X
Level VII: Expert opinion or consensus															

Adapted with permission from Melnyk BM, Fineout-Overholt E, editors. Evidence-based practice in nursing and healthcare: a guide to best practice. 2nd ed. Philadelphia: Wolters Kluwer Health / Lippincott Williams and Wilkins; 2010.

1 = Chan PS, et al. (2010); 2 = McGaughey J, et al.; 3 = Winters BD, et al.; 4 = Hillman K, et al.; 5 = Sharek PJ, et al.; 6 = Chan PS, et al. (2009); 7 = DeVita MA, et al.; 8 = Mailey J, et al.; 9 = Dacey MJ, et al.; 10 = McFarlan SJ, Hensley S.; 11 = Offner PJ, et al.; 12 = Bertaut Y, et al.; 13 = Benson L, et al.; 14 = Hatler C, et al.; 15 = Bader MK, et al.

Table 3: Effect of the Rapid Response Team on Outcomes

	1 ^a	2 ^a	3 ^a	4 ^a	5 ^a	6 ^a	7	8	9	10	11	12	13	14	15
HMR	— adult ↓ ^b peds	—	—	—	↓ ^b	—	NE	↓ ^c	↓ ^b	NR	NE	—	↓ ^c	NE	↓ ^{b, d}
CRO	NE	NE	NE	NE	↓ ^c	↓ ^b	NE	NE	↓ ^b	↓ ^c	↓ ^b	↓ ^c	NE	↓ ^c	↓ ^c
CR	↓ ^b peds and adult	NE	↓ ^b	—	NE	—	↓ ^b	↓ ^c	NE	NE	NE	NE	↓ ^b	NE	NE
UICUA	NE	NE	NE	—	NE	NE	NE	NE	↓ ^b	↑ ^c	NE	NE	NE	—	↓ ^b

1 = Chan PS, et al. (2010); 2 = McGaughey J, et al.; 3 = Winters BD, et al.; 4 = Hillman K, et al.; 5 = Sharek PJ, et al.; 6 = Chan PS, et al. (2009); 7 = DeVita MA, et al.; 8 = Mailey J, et al.; 9 = Dacey MJ, et al.; 10 = McFarlan SJ, Hensley S.; 11 = Offner PJ, et al.; 12 = Bertaut Y, et al.; 13 = Benson L, et al.; 14 = Hatler C, et al.; 15 = Bader MK, et al.

CR = cardiopulmonary arrest or code rates; CRO = code rates outside the ICU; HMR = hospital-wide mortality rates; NE = not evaluated; NR = not reported; UICUA = unplanned ICU admissions

^a higher-level evidence; ^b statistically significant findings; ^c statistical significance not reported; ^d non-ICU mortality was reduced

The EBP team can tell from reading the evidence that researchers consider the impact of an RRT on hospital-wide mortality rates (HMR) as the more important outcome; however, the group remains unconvinced that this outcome is the best for evaluating the purpose of an RRT, which, according to the IHI, is early intervention in patients who are unstable or at risk for cardiac or respiratory arrest.¹⁶ That said, of the 11 studies and projects that evaluated mortality, more than half found that an RRT reduced it. Carlos reminds the group that four of those six articles are level-VI evidence and that some weren't research. The findings produced at this level of evidence are typically less reliable than those at higher levels of evidence; however, Carlos notes that two articles

having level-VI evidence, a study and a project, had statistically significant (less likely to occur by chance, $P < 0.05$) reductions in HMR, which increases the reliability of the results.

Chen asks, since four level-VI reports documented that an RRT reduces HMR, should they put more confidence in findings that occur more than once? Carlos replies that it's not the number of studies or projects that determines the reliability of their findings, but the uniformity and quality of their methods. He recites something he heard in his Expert EBP Mentor program that helped to clarify the concept of making decisions based on the evidence: the level of the evidence (the design) plus the quality of the evidence (the validity of the methods) equals the strength of the evidence, which is

what leads clinicians to act in confidence and apply the evidence (or not) to their practice and expect similar findings (outcomes). In terms of making a decision about whether or not to initiate an RRT, Carlos says that their evidence stacks up: first, the MERIT study's results are questionable because of problems with the study methods, and this affects the reliability of the three systematic reviews as well as the MERIT study itself; second, the reasonably conducted lower-level studies/projects, with their statistically significant findings, are persuasive. Therefore, the team begins to consider the possibility that initiating an RRT may reduce code rates outside the ICU (CRO) and may impact non-ICU mortality; both are outcomes they would like to address. The evidence doesn't provide equally

promising results for UICUA, but the team agrees to include it in the outcomes for their RRT project because it wasn't evaluated in most of the articles they appraised.

As the EBP team continues to discuss probable outcomes, Rebecca points to one study's

data in the "Findings" column that shows a financial return on investment for an RRT.⁹ Carlos remarks to the group that this is only one study, and that they'll need to make sure to collect data on the costs of their RRT as well as the cost implications of the outcomes. They determine that

the important outcomes to measure are: CRO, non-ICU mortality (excluding patients with do not resuscitate [DNR] orders), UICUA, and cost.

Appraisal: Worth to Practice.

As the team discusses their synthesis and the decision they'll make based on the evidence,

Table 4. Defined Criteria for Initiating an RRT Consult

	4	8	9	13	15
Respiratory distress (breaths/min)	Airway threatened Respiratory arrest RR < 5 or > 36	RR < 10 or > 30	RR < 8 or > 30 Unexplained dyspnea	RR < 8 or > 28 New-onset difficulty breathing	RR < 10 or > 30 Shortness of breath
Change in mental status	Change in LOC Decrease in Glasgow Coma Scale of > 2 points	ND	Unexplained change	Sudden decrease in LOC with normal blood glucose	Decreased LOC
Tachycardia (beats/min)	> 140	> 130	Unexplained > 130 for 15 min	> 120	> 130
Bradycardia (beats/min)	< 40	< 60	Unexplained < 50 for 15 min	< 40	< 40
Blood pressure (mmHg)	SBP < 90	SBP < 90 or > 180	Hypotension (unexplained)	SBP > 200 or < 90	SBP < 90
Chest pain	Cardiac arrest	ND	ND	Complaint of nontraumatic chest pain	Complaint of nontraumatic chest pain
Seizures	Sudden or extended	ND	ND	Repeated or prolonged	ND
Concern/worry about patient	Serious concern about a patient who doesn't fit the above criteria	NE	Nurse concern about overall deterioration in patients' condition without any of the above criteria (p. 2077)	Nurse concern	• Uncontrolled pain • Failure to respond to treatment • Unable to obtain prompt assistance for unstable patient
Pulse oximetry (SpO₂)	NE	NE	NE	< 92%	< 92%
Other				• Color change of patient • Unexplained agitation for > 10 min • CIWA > 15 points	• UOP < 50 cc/4 hr • Color change of patient (pale, dusky, gray, or blue) • New-onset limb weakness or smile droop • Sepsis: ≥ 2 SIRS criteria

4 = Hillman K, et al.; 8 = Mailey J, et al.; 9 = Dacey MJ, et al.; 13 = Benson L, et al.; 15 = Bader MK, et al.

cc = cubic centimeters; CIWA = Clinical Institute Withdrawal Assessment; hr = hour; LOC = level of consciousness; min = minute; mmHg = millimeters of mercury; ND = not defined; NE = not evaluated; RR = respiratory rate; SBP = systolic blood pressure; SIRS = systemic inflammatory response syndrome; SpO₂ = arterial oxygen saturation; UOP = urine output

Rebecca raises a question that's been on her mind. She reminds them that in the "Appraisal: Worth to Practice" column, teaching was identified as an important factor in initiating an RRT and expresses concern that their hospital is not an academic medical center. Chen reminds her that even though theirs is not a designated teaching hospital with residents on staff 24 hours a day, it has a culture of teaching that should enhance the success of an RRT. She adds that she's already hearing a buzz

of excitement about their project, that their colleagues across all disciplines have been eager to hear the results of their review of the evidence. In addition, Carlos says that many resources in their hospital will be available to help them get started with their project and reminds them of their hospital administrators' commitment to support the team.

ACTING ON THE EVIDENCE

As they consider the synthesis of the evidence, the team agrees

that an RRT is a valuable intervention to initiate. They decide to take the criteria for activating an RRT from several successful studies/projects and put them into a synthesis table to better see their major similarities (see Table 4^{4, 8, 9, 13, 15}). From this combined list, they choose the criteria for initiating an RRT consult that they'll use in their project (see Table 5). The team also begins discussing the ideal make up for their RRT. Again, they go back to the evaluation table and look

Table 5. Defined Criteria for Initiating an RRT Consult at Our Hospital

Pulmonary	
Ventilation	Color change of patient (pale, dusky, gray, or blue)
Respiratory distress	RR < 10 or > 30 breaths/min or unexplained dyspnea or new-onset difficulty breathing or shortness of breath
Cardiovascular	
Tachycardia	Unexplained > 130 beats/min for 15 min
Bradycardia	Unexplained < 50 beats/min for 15 min
Blood pressure	Unexplained SBP < 90 or > 200 mmHg
Chest pain	Complaint of nontraumatic chest pain
Pulse oximetry	< 92% SpO ₂
Perfusion	UOP < 50 cc/4 hr
Neurologic	
Seizures	Initial, repeated, or prolonged
Change in mental status	• Sudden decrease in LOC with normal blood glucose • Unexplained agitation for > 10 min • New-onset limb weakness or smile droop
Concern/worry about patient	Nurse concern about overall deterioration in patients' condition without any of the above criteria
Sepsis	
	• Temp, > 38°C • HR, > 90 beats/min • RR, > 20 breaths/min • WBC, > 12,000, < 4,000, or > 10% bands

cc = cubic centimeters; hr = hours; HR = heart rate; LOC = level of consciousness; min = minute; mmHg = millimeters of mercury; RR = respiratory rate; SBP = systolic blood pressure; SpO₂ = arterial oxygen saturation; Temp = temperature; UOP = urine output; WBC = white blood count

over the “Major Variables Studied” column, noting that the composition of the RRT varied among the studies/projects. Some

evidence that led to the project, how to call an RRT, and outcome measures that will indicate whether or not the implementation

As they consider the synthesis of the evidence, the team agrees that an RRT is a valuable intervention to initiate.

RRTs had active physician participation (n = 6), some had designated physician consultation on an as-needed basis (n = 2), and some were nurse-led teams (n = 4). Most RRTs also had a respiratory therapist (RT). All RRT members had expertise in intensive care and many were certified in advanced cardiac life support (ACLS). They agree that their team will be comprised of ACLS-certified members. It will be led by an acute care nurse practitioner (ACNP) credentialed for advanced procedures, such as central line insertion. Members will include an ICU RN and an RT who can intubate. They also discuss having physicians willing to be called when needed. Although no studies or projects had a chaplain on their RRT, Chen says that it would make sense in their hospital. Carlos, who's been on staff the longest of the three, says that interdisciplinary collaboration has been a mainstay of their organization. A physician, ACNP, ICU RN, RT, and chaplain are logical choices for their RRT.

As the team ponders the evidence, they begin to discuss the next step, which is to develop ideas for writing their project implementation plan (also called a protocol). Included in this protocol will be an educational plan to let those involved in the project know information such as the

of the evidence was successful. They'll also need an evaluation plan. From reviewing the studies and projects, they also realize that it's important to focus their plan on evidence implementation, including carefully evaluating both the process of implementation and project outcomes.

Be sure to join the EBP team in the next installment of this series as they develop their implementation plan for initiating an RRT in their hospital, including the submission of their project proposal to the ethics review board. ▼

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