



Evaluating the Impact of Peri-Simulation Education on Medical Student Performance: A Randomized Controlled Trial

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Abstract

Introduction: Limited research exists evaluating pre-clinical medical student performance in Simulation Patient Encounters (SPEs), nor the effectiveness of teaching interventions as part of the learning. We sought to determine if a teaching intervention prior to simulation training improves medical student clinical decision-making skills compared to baseline performance.

Methods: A prospective randomized single-center crossover study comparing performance of second-year medical students in two SPEs. The groups were then randomized to receive an educational intervention either the day before the SPEs or at the conclusion of the two SPEs. The educational intervention was a 10-min video. Analysis was designed to evaluate if a difference existed between performance outcomes using a standardized checklist.

Results: The video prior cohort scored higher than the no video prior group in both case scenarios. In the anemia case scenario, video prior cohort scored 41.6% (23.69) vs. 38.9% (22.19) in the no video cohort. In the diverticular bleed case scenario, the video prior cohort scored 37.6% (19.94) vs. 37.1% (19.69) in the no video prior cohort. Additionally, for the diverticular bleed scenario there was a statistically significant higher number of prompting in the video prior cohort than the no video prior, with an average of 2.44 prompts vs. 1.56 respectively (p=0.04).

Conclusion: Our results did not find an impact of the educational video on the performance scores in the SPEs. This study had several limitations and calls for future studies with a more controlled environment to determine timing, efficacy, and impact of educational interventions in simulation training.

Keywords: SPE; Clinical reasoning; Diagnostic error; Medical education; Educational intervention

Abbreviations

SPE: Simulated Patient Encounter; HPI: History of Presenting Illness; PE: Physical Exam; MCQ: Multiple Choice Questionnaire; VI: Vertical Integration; GI: Gastrointestinal

Introduction

Mannequin-based simulation trainings (SIMs) are a cornerstone of modern medical education. It is increasingly used as it has developed into a highly effective teaching modality for a broad range of settings [1,2]. SIMs have shown to be an efficient, and cost-effective to develop such clinical skills, especially when compared to traditional teaching methods of slideshows and lectures [3-8]. Additionally, they allow students to learn in a safe and controlled environment without the risk of harm to real patients [9]. The advent of COVID-19 prohibited medical students from entering the clinical environments which further highlighted the importance of clinical development without patient contact [10,11]. In addition to the cost-effectiveness and risk avoidance, there are benefits to pre-clinical medical student learning; SIMs can be directed to a target a broad range of specific knowledge and skills targeting the development of clinical interviewing, physical

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examination, and diagnostic skills [12,13]. When designed well, this training allows for detailed data collection; assessment of score performance; and provide real-time feedback to learners [1,12,14,15]. Despite the increased utilization and employment of SIMs in medical school education, significant challenges remain. Although medical simulation has great potential, SIMs have significant variation in methodology, design, and outcomes. Research on SIMs within medical education has demonstrated insufficient technical quality to produce reliable results. This variation may be attributed to the lack of standardized guidelines for conducting SIMs for medical student training [1,3,16]. Incorporating teaching methods that is specific to the simulation learning objectives is a key component of optimizing simulation training. However, there is limited such research on such peri-SIM evaluating the impact of educational interventions on performance outcomes. Barriers to effective simulation from the institutional standpoint include the need for skilled simulation operators, financial support for the resources, and educators trained in simulation [17,18]. Several qualitative and quantitative evaluations measuring performance outcomes from SPEs have been identified. Quantitative outcomes are preferred due to a higher resolution of data [3,13,19,20]. Validated quantitative outcome measures of learner performance in simulations include time-to-action, pre-test assessments, and performance scoring using checklists of definable actions [3,19-22]. Although several studies have attempted to validate methods of simulation for medical student clinical skills acquisition, systematic reviews of the SIMs have proven limited by variations in methodology [19,20,23]. Medical schools should aim to approach simulation training in parallel to the rigorous method of evidence-based practice in modern medicine. Proper practice confers application of proper technique, and conversely improper practice can beget poor habits. Further research is needed to assess the impact of simulations on pre-clinical medical student skills competence and whether specific educational interventions can improve learning. Thus, our study aimed to add to the literature our experience with the use of simulation training as a learning modality for medical student education. We sought to determine how the timing of an educational intervention impacted the performance of second year medical students in a SPE setting as measured by a standardized skills checklist.

Materials and Methods

Study participants

All second-year medical students attending the University of Toledo College of Medicine and Life Sciences were eligible for the study. The trial was explained in detail to the group, and students were assured of the anonymity and confidentiality of personal information for all responses throughout the study period. Students that did not consent for the study did not have their coursework grades accessed and they were not asked to complete the voluntary feedback survey. A total of 174 medical students were eligible for and participated in this single center, IRB approved double blinded randomized trial.

Study design

A mixed methods study was implemented that included a randomized-controlled trial with two intervention arms designated as "gold" and "blue". To ensure that all students had a similar overall learning experience, all students received an educational video. The blue group received instructions to view the video the day prior to the SPEs. The gold group received instructions to view the video at the conclusion of the SPEs. The SPE's were a formative, non-graded,

activity designed as an opportunity for medical student learning. A two-stage randomization strategy was implemented. First, students were randomized into 32 groups of approximately equal size (4 to 6 students each). The 32 groups were then randomized into either the gold group or the blue group. The blue group was randomized to receive the video prior to the SPEs, and gold group was randomized as the control with no video prior. The randomization process was carried out by a faculty of UTCOMLS blinded from the study. The groups had a pre-randomization to the location of the simulation room and the SPE scenario. The blue and gold intervention arms each had half of the students participate in the upper GI bleed scenario first followed by the diverticular bleed second and the other half participated in the diverticular bleed scenario first followed by the upper GI bleed scenario second (Figure 1). Assignment of students and the randomization process was concealed from the students and independent from the research team. After randomization, the students received the intervention in a double blinded setting (both the subject and the investigator were blinded, such that the subjects ID and the intervention group were not linked together until after the end of the study when all results were unblinded for analyzation. The groups were assessed for completing key aspects of the history of presenting illness, physical exam, assessment/plan/intervention, and number of times the facilitator had to prompt students to keep the sim moving forward. Prompting statements were standardized to include: "are there any tests/labs/imaging or other-diagnoses that you may want to consider for this patient." The reporting guidelines for health care simulation research extensions to the CONSORT and STROBE statements were utilized to guide study design and write-up with attached checklists [24].

Simulated patient scenarios

Each simulation scenario was 15 min long followed by a 5-min debriefing session. There were four simulations running simultaneously with 8 SPE's per group session, for a total 64 SPE's. Both scenarios mirrored a total of two simulation groups running at the same time. There was a total of 4 operators running the simulated patient scenarios. The operators ran the same case, in the same room and manakin for the duration of the study. The operators were given a standardized training session for their respective cases in attempt to keep the language and overall atmosphere the same for each of the groups. The two SPE scenarios were developed by faculty, medical physicians, at UTCOMLS independent of the research investigators. The study team did not have an impact on the patient scenarios or presentation of the scenarios in any manner. The scenarios included were one adult patient with an acute lower GI bleed from diverticulitis and one adult patient with a chronic occult upper GI bleed. The information the students were expected to obtain were developed by the faculty independent of the study team as well, which were used for the performance evaluation as explained in the measurement of outcomes section below. The students were given the following information verbally from the simulation operators, "You will have 15 min to conduct a focused history and physical exam. As you develop your differential diagnosis you may order labs, tests, and imaging that you believe is necessary for the work-up of the patient. Perform any necessary interventions as you see fit."

Educational intervention

The educational intervention was designed by collaboration between UTCOMLS faculty and the study team. The intervention encompassed a 10-min recorded power-point presentation. The learning objectives were to teach, and review the evaluation,

management, diagnosis and treatment of intestinal ischemia and bleeding.

Data collection

Two blinded independent evaluators viewed video recordings of the groups during their SPEs. Both reviewers were 4th year medical students who were observed independently by a faculty for 3 videos to ensure accuracy of the assessment. The evaluators recorded whether the students performed the checklist of activities for the respective chronic upper GI bleed and acute diverticular bleed SPE's (Supplemental Table 1). These checklists were modified versions of the checklists used at UTCOMLS for their in-person patient interviews as part of the medical student educational curriculum. The modification was performed by two physicians on faculty at the medical school.

Statistical analysis

The primary objective is to compare the performance effect as measured by adapted analysis of student performance in a group setting of simulated patient encounters. The hypothesis that the Intervention = 0 was tested utilizing independent t-tests to compare mean group performance scores in the separate intervention arms. A p-value less than, or equal to 0.05 was regarded as statistically significant. This analysis approach makes two main assumptions (in addition to the normality assumption for the students paired t-test): No period effect and no intervention-period interaction. Secondary objectives will be to compare the sub-categorization of the scoring checklist. The change in accuracy of group performance in patient encounters was compared between the two intervention arms, video prior vs. no video. Comparisons of proportions were conducted via chi squared test. All statistical tests were performed using excel.

Results

170 students participated in the two SPEs. The total scores in the video prior cohort were higher than the no video prior group in both case scenarios (Table 1). In the Anemia case scenario, video prior cohort scored 41.6% (23.69) vs. 38.9% (22.19) in the no video cohort (p=0.26). In the diverticular bleed case scenario, the video prior cohort scored 37.6% (19.94) vs. 37.1% (19.69) in the no video prior cohort (p=0.85). In the diverticular bleed scenario, there was a statistically significant higher number of prompting in the video

prior cohort with an average of 2.44 prompts vs. 1.56 in the no video cohort (p=0.04). The average scores of the anemia case were higher in room 2 compared to room 1, with 42.2% vs. 38.3% respectively. The average scores of the diverticular case were higher in room 3 vs. room 4, with 39.1% vs. 35.6% respectively. However, there was not a statistically significant difference in scores by the room in which the case was presented (Table 2).

Discussion

Importance of medical student clinical skills training

SPEs were previously a feature of the United States Medical Licensing Exam (USMLE) Step 2 Clinical Skills (CS), however recently the CS portion of Step 2 was discontinued [25]. The CS portion of Step 2 was used by residency programs to screen applicants, and prior to its discontinuation, many programs indicated a desire for higher resolution data from the score report of Step 2 CS [26,27]. In response, co-sponsors of the USMLE program have stated an intention to develop novel assessments of clinical competency [28]. Although USMLE Step 2 CS was discontinued, most residency program directors previously endorsed Step 2 CS as factoring heavily into their admissions decision-making [27]. The removal of Step 2 CS has created a vacuum in the standardized assessment of medical students' clinical and communicative prowess [25]. A high granularity of data is desirable when considering the viability of potential clinical assessment methods [25,27]. Clinical skills and communication abilities are seen by many RPDs as lagging behind the book knowledge of incoming residents. The evaluation of soft skills including communication and professionalism is undoubtedly more complex than the accurate assessment of content knowledge. The present study typifies the need for continued research into the process-based evaluation of medical students' clinical competency.

Medical student learning through simulation

Cook et al. reported a moderate effect of simulation in health professionals compared to other instruction methods in a systematic review and meta-analysis [29]. Previous studies on laparoscopy and Advanced Cardiac Life Support (ACLS) have demonstrated simulation training improves clinical outcomes [1]. However it is unclear whether simulation training improves patient outcomes across other skill domains, and further studies comparing the efficacy

Table 1: Two-tailed Independent T Test analysis of performance score totals and sub-categorical scores comparing the intervention to no intervention expressed as the mean (SD), percent correct, and 95% CI. Sub-categories included the History of Presenting Illness (HPI), Physical Exam (PE), Assessment/Plan/Intervention (A/P/I). Categories are expressed with the (total number) of eligible points. The normality assumption was checked based on the Shapiro-Wilk Test (alpha =0.05).

Anemia					
Category (points)	SPE-1 (no video)	95% CI	SPE-2 (video prior)	95% CI	P-Value
Total (57)	22.19 (3.78) 38.9%	20.33-24.04	23.69 (3.63). 41.6%	21.91-25.47	0.26
HPI (20)	9.00 (2.66) 45.0%	7.69-10.30	8.69 (2.55). 43.4%	7.44-9.94	0.74
Vitals & PE (12)	3.94 (2.02) 39.4%	2.95-4.93	5.13 (1.82). 51.3%	4.24-6.02	0.09
A/P/I (25)	9.25 (1.77) 37.0%	8.38-10.12	9.88 (2.19). 39.5%	8.80-10.95	0.38
Prompting (*)	2.44 (1.26)	1.82-3.06	2.50 (1.59)	1.72-3.28	0.9
Diverticular Bleed					
Category (points)	SPE-1 (Video Prior)	95% CI	SPE-2 (No video)	95% CI	P-Value
Total (53)	19.94 (3.66) 37.6%	18.14-21.73	19.69 (4.00) 37.1%	17.73-21.65	0.85
HPI (20)	6.94 (1.65) 34.7%	6.13-7.75	6.31 (2.24). 31.6%	5.21-7.41	0.38
Vitals & PE (10)	3.94 (1.61) 32.8%	3.15-4.73	4.31 (1.54). 35.9%	3.56-5.07	0.51
A/P/I (23)	6.56 (1.75) 28.5%	5.71-7.42	6.63 (2.16). 28.8%	5.57-7.68	0.93
Prompting (*)	2.44 (1.21)	1.85-3.03	1.56 (1.09)	1.03-2.10	0.04

Supplemental Table 1: Checklist for performance scoring in Sim 1 and Sim 2 respectively. Red text indicates categories; black text indicates 1 point available for completion.

Chronic Upper GI bleed	Acute Diverticular Bleed
HPI	HPI
Patient Name	Patient Name
Chief complaint	Chief complaint
location/radiation	location/radiation
quantity/severity	quantity/severity
timing (Onset/frequency/duration)	timing (Onset/frequency/duration)
setting in which it occurs/has it happened before	setting in which it occurs/has it happened before
exacerbating factors	exacerbating factors
remitting factors	remitting factors
associated symptoms	associated symptoms
patient perspective	patient perspective
medications	medications
allergies	allergies
tobacco use	tobacco use
alcohol use	alcohol use
illicit drug	illicit drug
ROS (8 categories for a point)	ROS (8 categories for a point)
Family History	Family History
Vaccinations	Vaccinations
Past Medical Hx	Past Medical Hx
Past surgical hx	Past surgical hx
Physical Exam & vitals	Physical Exam & vitals
Auscultate abdomen x 4 quadrants	Auscultate abdomen x 4 quadrants
palpate/percuss	palpate/percuss
Look to assess symmetry/contour	Look to assess symmetry/contour
Assess for Peritoneal signs (rebound tenderness/guarding)	Assess for Peritoneal signs (rebound tenderness/guarding)
Rectal exam	Rectal exam
Auscultated Heart areas	Heart rate
Auscultated Lung Fields	RR
Heart rate	Temperature
RR	BP
Temperature	o2 sat
BP	Assessment/Plan/Interventions
o2 sat	Ask for CXR
Assessment/Plan/Interventions	Identified Normal lung fields
Ask for CXR	Ask for VBG or ABG
Identified Normal lung fields	Ask for CBC
Ask for VBG or ABG	Identified Anemia aka low hemoglobin
Identified acidosis aka Low PH	Ask for CMP
Ask for CBC	Identified acidosis
Identified Anemia aka low hemoglobin	Identified prolonged PTT/INR
Identified normocytic anemia	Ask for Fecal Occult Blood test (FOBT)
Ask for CMP	Ask for EKG
Identified elevated BUN	Identified sinus tachycardia
Ask for coagulation panel	Performed blood Type and cross
Identified prolonged PTT/INR	started 2 large bore IV access

Ask for Fecal Occult Blood test (FOBT)	Administer Fluids
Ask for EKG	Administered PRBC
Identified sinus rhythm	oxygen (nasal cannula)
Performed blood Type and cross	Vasopressor administered
Ordered Troponin	Warfarin Reversal agent PCC/FFP administered
Started 2 large bore IV access	Ordered EGD
Administer Fluids	Ordered Colonoscopy
Administer Oxygen (nasal cannula)	Ordered CT abdomen & pelvis
Ordered EGD	NG Tube
Ordered Colonoscopy	prompting
Ordered CT Abdomen	
Ordered CT angiogram	
Ordered NG tube	
Administered PRBC	
prompting	

Table 2: Two-tailed Independent T test of combined total score of SPE-1 & SPE-2 comparing intervention to no intervention, and simulation room of the cases.

	Combined SPE-1 & -2 Score	95% CI	P-Value
Video Prior	43.63 (4.88) 39.7%	41.23-46.02	0.339
No Video	41.88 (5.32) 38.1%	39.27-44.48	
Room 1 (anemia)	21.81 (3.67) 38.3%	20.01-23.61	0.09
Room 2 (anemia)	24.06 (3.53) 42.2%	22.33-25.79	
Room 3 (diverticular)	20.75 (3.51) 39.1%	19.03-22.47	0.16
Room 4 (diverticular)	18.88 (3.90) 35.6%	16.97-20.78	

of the multitude of simulation modalities, especially on pre-clinical medical students, are needed [1,23]. Issenberg et al. [16] identified 10 features of simulation that contribute to effective learning in 2005. These features listed in descending order from most effective to least effective include providing feedback, repetitive practice, curriculum integration, range of difficulty, multiple learning strategies, capture clinical variation, controlled environment, individualized learning, defined outcomes, and simulator validity. The meta-analysis by Cook in 2012 confirmed these factors, but also found that group instruction was associated with improved outcomes [20]. Our study incorporated most of these factors, apart from having a controlled simulation environment and individualized learning.

Simulation methodology

Our study assessed the impact of a 10-min video intervention on performance scores of second-year medical students in two gastrointestinal SPEs. Students either received the intervention the day prior to the SPE experience or after the conclusion of the second SPE. In this population of second-year medical students, we found no significant difference in performance scores when comparing the group that received the video-intervention the day prior to the simulations versus those that did not receive the intervention prior. Although statistical significance was not reached, the video-prior group scored higher in both simulation exercises. On further analysis there were differences in the total average score of the SPEs depending on the room, and operator, in which the activity was performed. Although these differences did not reach statistical significance, it highlights the importance of standardizing the instructions for operators and having highly trained competent operators for simulation training.

A meta-analysis comparing simulation interventions found the time of the simulation to be a significant factor affecting performance [20]. In our study, the sims were limited to 10-min sessions which may not have been enough time for them to have fully assessed the patient.

Simulation performance assessments

No significant inter-rater reliability was observed, owing to the skills checklist composed of largely objective measures. There was room for rater subjectivity on some of the physical examination items (e.g., “Look to assess symmetry/contour”) and diagnostic testing items (“Identified normal lung fields on chest X-ray”). A planned multiple-choice questionnaire pre and post-test as part of the study protocol was not administered due to time constraints. Thus, the student's baseline knowledge was unable to be assessed and may have been a confounder of the results observed. Additionally, without the MCQ testing we were unable to define if the simulation had an impact on objective knowledge testing. A post-SPE feedback survey could have elucidated if learners felt the simulation improved their application of knowledge in the clinical setting. Lower scores were observed in the acute diverticular bleed SPEs compared to the chronic GI bleed SPEs. This could have been due to slight differences in between the two grading rubrics; however, the acute nature of the former presentation could have contributed to learner stress levels and thus cognitive load. The acute scenario showed less difference in overall performance between the intervention group and the treatment group compared to the chronic scenario. Additionally, learners in the acute scenario performed lower on nearly all subsections of the checklist compared to learners in the chronic scenario, potentially reflecting a relationship between cognitive load and acuity of care. In the acute care setting, perceived case difficulty has shown to be a strong predictor of cognitive load [30]. Prompting was the only subsection in which learners in the acute setting outperformed those in the chronic setting—in the acute SPE, the intervention group required on average nearly one more cue prompt per SPE. This could simply be because acute care affords practitioners less time to pause, deliberate, and possibly get stuck during decision-making.

Educational interventions for simulation

Carter et al. [31] performed a cross over study involving third year medical students to determine if a 60-min lecture prior to a 20-min SPE involving peripheral vascular disease affected third year medical

student performance scores as measured by a standardized checklist. The students who received the pre-SPE training scored higher than did those who did not receive the lecture. Our study observed similar results with increased performance in the physical exam and vitals subcategory among learners receiving a pre-SPE educational intervention; however, the results observed in our study failed to reach statistical significance. Other studies have observed SPEs to be especially effective at improving medical student history-taking and physical examination skills [4,32]. Further research is needed to assess the impact of SPEs on pre-clinical learners who have a limited to no clinical experience and potentially a lower degree of confidence in their clinical skills compared to third-year students participating in clinical rotations.

Future considerations

For the performance scores as the day progressed, there may have been a knowledge gap. These sims were held as part of a full day of GI activities which may have influenced students' abilities to perform during the sims. All groups participated in a 'GI Olympics' education day, in which they had a team-based learning activity in the anatomy lab followed by an introduction to suturing session before participating in the SPE scenarios evaluated in this study. Additionally, the students later in the day may have had cognitive fatigue, whereas the students earlier in the day had less exposure to the GI activities which may have weakened their performance. Due to the students having interactive activities throughout the day there was ample opportunity that the students may have shared information about the simulation cases which may have impacted their performance. Cognitive load theory suggests that certain types of education are un conducive to learning new knowledge and skills due to the cognitive fatigue they impose [33]. Managing cognitive load plays an important role in the setting of medical education, in which problems are complex and students are expected to learn a high volume of material [34,35]. The design of the present study relates to cognitive load theory because the use of a pre-SPE intervention video was intended to help students schematize the complex process of reaching a diagnosis by starting with a chief complaint. We observed higher skill checklist completion percentages in the pre-SPE video intervention group compared to learners who finished the SPE before receiving intervention, albeit without statistical significance.

Conclusion

This article has presented an overview of the difficulty in conducting research relevant to medical student simulations training using a failed experimental design. It discusses technical considerations when approaching the design, assessment of, and implementation of simulation. This article provides meaningful recommendations for future simulation execution and research study considerations. The key findings from this study relate the importance of developing standardized protocols for assessing medical student performance in simulation training.

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