



Haptic simulators accelerate laparoscopic simulator training, but skills are not transferable to a non-haptic simulator: a randomized trial

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Received: 10 March 2022 / Accepted: 24 June 2022 / Published online: 2 August 2022
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Abstract

Background Laparoscopy requires specific psychomotor skills and can be challenging to learn. Most proficiency-based laparoscopic training programs have used non-haptic virtual reality simulators; however, haptic simulators can provide the tactile sensations that the surgeon would experience in the operating room. The objective was to investigate the effect of adding haptic simulators to a proficiency-based laparoscopy training program.

Methods A randomized controlled trial was designed where residents ($n = 36$) were randomized to proficiency-based laparoscopic simulator training using haptic or non-haptic simulators. Subsequently, participants from the haptic group completed a follow-up test, where they had to reach proficiency again using the non-haptic simulator. Participants from the non-haptic group returned to train until reaching proficiency again using the non-haptic simulator.

Results Mean completion times during the intervention were 120 min (SD 38.7 min) and 183 min (SD 66.3 min) for the haptic group and the non-haptic group, respectively ($p = 0.001$). The mean times to proficiency during the follow-up test were 107 min (SD 41.0 min) and 58 min (SD 23.7 min) for the haptic and the non-haptic group, respectively ($p < 0.001$). The haptic group was not faster to reach proficiency in the follow-up test than during the intervention ($p = 0.22$). In contrast, the non-haptic group reached the required proficiency level significantly faster in the follow-up test ($p < 0.001$).

Conclusion Haptic virtual reality simulators reduce the time to reach proficiency compared to non-haptic simulators. However, the acquired skills are not transferable to the conventional non-haptic setting.

Keywords Laparoscopy · Simulation · Haptics · Training

Laparoscopic surgery requires specific psychomotor skills of the surgeon who has to work in a three-dimensional space guided by two-dimensional images and has limited tactile feedback compared to open surgery. Performing laparoscopic procedures safely requires depth perception and excellent hand–eye coordination. These skills can be practiced safely outside the operating room, increasing the

demand for efficient simulation-based training, e.g., using virtual reality simulators [1–7]. Previous studies have shown that virtual reality simulators can shorten the time for novice surgeons to reach proficiency, and the skills are transferable to the operating room [8, 9]. Therefore, virtual reality simulators are widely used for laparoscopy training and become a standard part of modern surgical education [10–12]. The virtual reality simulators can also provide an automated assessment of trainees' performance, enabling them to use a mastery learning approach [13].

However, most laparoscopic training programs with demonstrated clinical transfer were created on virtual reality simulators without haptic feedback; this has been a concern from a clinical point of view as this differs from real surgeries. Although the tactile sensations in laparoscopic surgery are less evident than open surgery, haptic sensations can still be perceived through the instruments by the operating surgeon [14, 15]. These tactile sensations, especially pulling and grasping, are important as understanding them provides

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valuable information; excessive grasping can damage tissue, whereas inadequate tension can lead to tissue slippage. [16–18].

Therefore, virtual reality simulators with haptics for laparoscopy have been developed. These are more realistic but costly and more complex [19, 20], and there could be a risk for a higher frequency of technical malfunctions, disturbing training. [21, 22]. Previous studies have investigated the effect of haptic devices on virtual reality simulators and found that the haptic devices accelerate the initial learning curve of trainees [19, 23–28], but these studies only included very few repetitions or a limited amount of training time. The impact of haptic devices in a proficiency-based training program for laparoscopy has not been explored.

Instructor-based feedback has proven to be a valuable tool in reducing time to reach proficiency in laparoscopic simulator training programs without negatively impacting the retention of skills [29]. However, different training setups, in this case, using a haptic simulator versus the non-haptic simulator, might require different amounts of instructor assistance.

The objectives of this trial were to investigate if training with virtual reality with a haptic device compared with a non-haptic simulator resulted in reduced time to reach proficiency for surgical novices, how it affected their retention of skills and the need for instructor feedback and frequency of simulator malfunctions.

Materials and methods

A single-center randomized superiority trial was planned according to the CONSORT statement [30]. The trial was exempt from ethical approval by the Regional Committee on Biomedical Research Ethics (H-20063093). The trial was registered at Clinicaltrials.gov (NCT05191589).

Setting

Data collection was done at the simulation center at Copenhagen Academy for Medical Education and Simulation (CAMES) in Copenhagen, Denmark [31].

Participants

Participants were surgeons without any previous laparoscopic experience. Participants were recruited from the surgical departments in the eastern part of Denmark and were invited by e-mail and received written and verbal information before filling out the informed consent forms. The inclusion and exclusion criteria used for the trial were as follows:

Inclusion criteria for both intervention and follow-up: (1) Residents working in the Eastern part of Denmark; (2) Provided informed consent before inclusion.

Exclusion criteria for intervention: (1) Having previously participated in studies involving laparoscopic training; (2) Having participated in laparoscopic training programs at any simulation center; (3) Having prior experience with laparoscopic surgery (having performed any laparoscopic procedures as primary surgeon, including supervised procedures); (4) Having performed any supervised laparoscopy procedures as primary surgeons during intervention; (5) Did not give consent; (6) Did not speak Danish on a conversational level.

All participants were given a unique trial identification number before randomization.

Intervention and control

Participants were randomized to proficiency-based training with either a haptic (haptic group) or non-haptic laparoscopy simulator (non-haptic group) during the intervention. All participants were invited back after a 3–6 weeks break for a follow-up test where both groups had to practice until they reached proficiency again using the non-haptic setup. No simulator training was allowed, nor any operations as primary surgeon in any laparoscopic procedures during this break.

The simulator training program consisted of four basic skills tasks (Instrument Navigation; Grasping; Lifting and grasping, and Fine dissection) and a procedural task (salpingectomy due to a bleeding ectopic pregnancy) on a virtual reality simulator [8, 13]. All participants had to reach proficiency for all modules, which entails passing twice within five consecutive attempts. The predefined proficiency level was established in previous trials and includes the parameters such as instrument time, instrument path length, instrument angular path length, tissue damage, bleeding, and energy damage [8, 13]. The metrics of these metrics are summarized in the [Appendix](#). Participants booked training sessions by e-mail, and a maximum of one 2-h training session per day was allowed to minimize cognitive overload and fatigue.

Randomization

A 1:1 randomization was performed centrally using a web-based system from Sealed Envelope (London, United Kingdom). The allocation sequence was computer-generated and used varying block sizes of four and six, a sequence that was kept concealed from the principal investigator throughout the trial.

Materials and equipment

We used six LapSim® virtual reality simulators (Software version 2019.1) from Surgical Science (Gothenburg, Sweden) – three simulators with haptics and three simulators without. The only difference between the two setups was the tactile sensation provided by the haptic simulators. To prevent overworking the simulators leading to a potential risk of further technical malfunctions, participants were randomized to a different simulator for each training session within their allocation to minimize bias from the same person training on a simulator which could experience technical problems.

All simulators were connected to a 27" screen and were height adjustable to ensure the best viewing condition and the most ergonomically correct working position. Each simulator was separated by dividing walls, and noise-canceling Bose Quiet Comfort III headsets were worn by all participants during the trial when training [31].

Outcome measures

The primary outcome was the total time (min) spent to reach the predefined proficiency level for all five tasks. The secondary outcome was the time (min) to reach proficiency on the conventional non-haptic setting after 3–6 weeks of no laparoscopic training.

Exploratory outcomes were instructor time (s) spent on feedback during training and number of malfunctions and time spent solving them during the intervention and follow-up test.

We also calculated the cumulative time (min) training to proficiency, meaning the sum of the time to proficiency during the intervention and the follow-up test.

Sample size calculation

The sample size was calculated based on the primary outcome, time to reach proficiency, using data from a previous trial conducted by the same research group [32]. For the non-haptic group, it was assumed that a mean time of 320 min was needed to reach proficiency, while for the haptic group, a mean time of 240 min was expected. Standard deviations of 70 were assumed for both groups. Using a

two-sided significance level of 0.05 and a power of 0.90, the minimum sample size required was 34 participants, 17 in each group.

Statistical analysis

The data was analyzed using SPSS® version 27.0 (IBM, Armonk, NY, USA) and RStudio 2021© (RStudio, Boston, MA, USA). Independent samples *t*-tests were used for inter-group comparisons for primary, secondary, and exploratory outcomes during the intervention and the follow-up test.

To analyze the training effect, over time, for both the haptic and the non-haptic group, the mixed model with repeated measurements and unstructured covariance matrix model was used for time to reach proficiency and instructor-based feedback. The basic model was $Y = a + bI + ct + dtI$, where *I* is the indication of the intervention, *t* is time (time1 corresponding to the intervention phase and time2 corresponding to the follow-up phase), and *a* through *d* are coefficients of the regression equation.

Finally, Fisher's exact test was used to compare the number of technical malfunctions between the haptic and non-haptic simulator.

Results

The patient demographics for the trial are presented in Table 1 and the trial flowchart in Fig. 1. We found that the haptic group reached proficiency significantly faster than the non-haptic group during the intervention ($p = 0.001$). In contrast, the non-haptic group reached proficiency significantly faster than the haptic group ($p < 0.001$) during the follow-up test. Furthermore, we found that the haptic group had no significant improvement, regarding time spent, from the trial's intervention to the follow-up test ($p = 0.22$). However, the non-haptic group improved their time to reach proficiency significantly faster from the intervention to the follow-up test ($p < 0.001$). There was no significant difference between the two groups for the cumulated time spent reaching proficiency ($p = 0.42$) (Table 2) (Fig. 2).

During the intervention, we found that the haptic group required significantly less instructor assistance than the non-haptic group ($p < 0.001$). For the follow-up test, we

Table 1 Baseline characteristics for participants who completed the intervention

Groups	Haptic group (<i>n</i> = 19)	Non-Haptic group (<i>n</i> = 17)
Sex (number of men/women)	4/15	5/12
Age (median/range)	29 (28–39)	29 (27–33)
Dexterity (number of right-/left-handed)	15/4	14/3

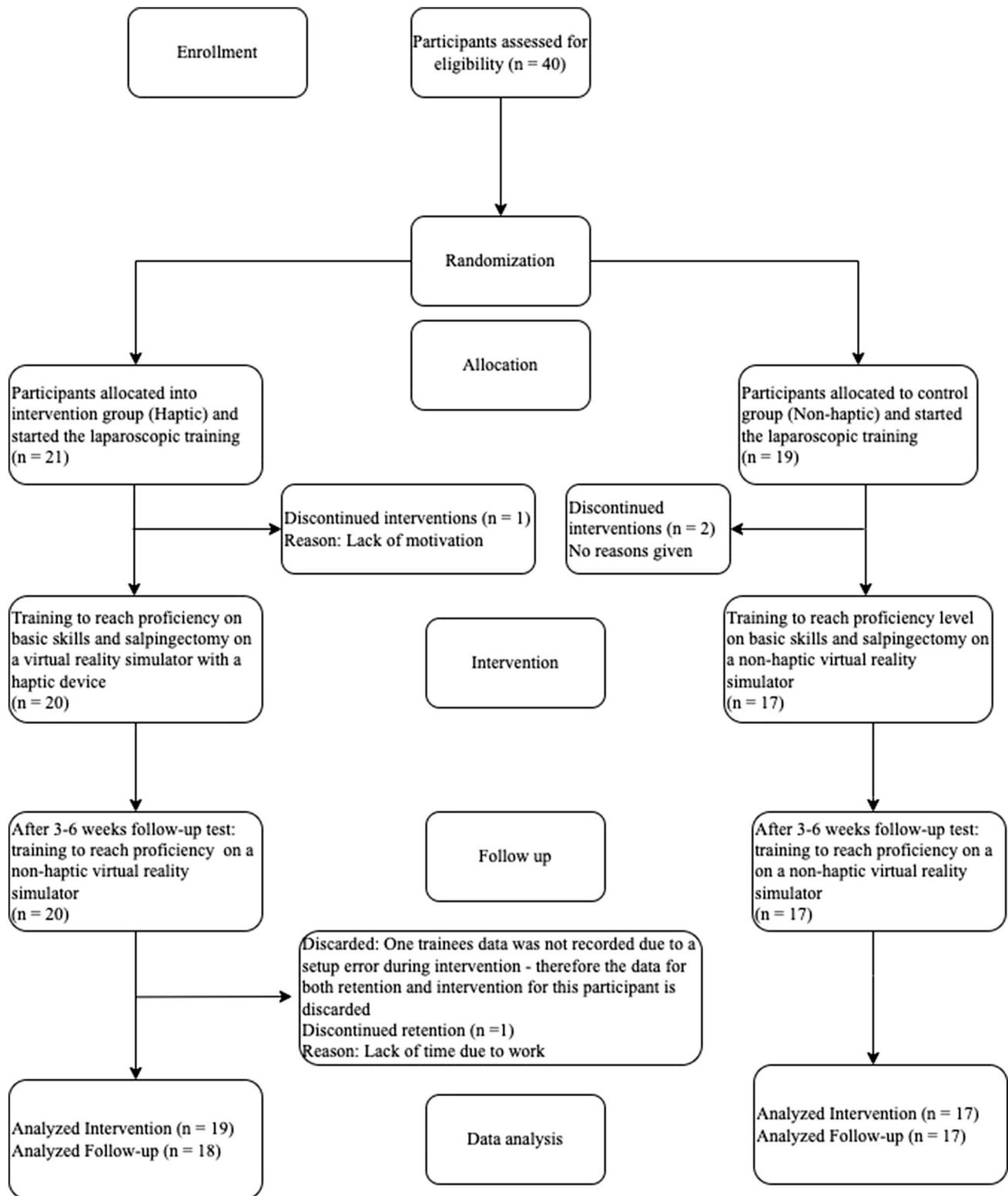
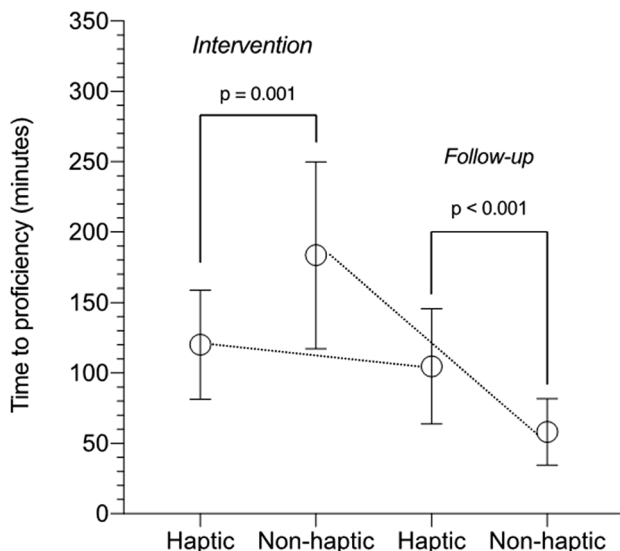


Fig. 1 Flowchart according to consort statement

Table 2 Mean training time to reach proficiency for intervention and follow-up

	Haptic group	Non-haptic group	<i>p</i> values
Intervention (min)	120 (95% CI [83; 149]) (<i>n</i> = 19)	183 (95% CI [149; 217]) (<i>n</i> = 17)	0.001
Follow up (min)	107 (95% CI [87; 128]) (<i>n</i> = 18)	58 (95% CI [46; 70]) (<i>n</i> = 17)	< 0.0001
Cumulated time (min)	219 (95% CI [185; 253]) (<i>n</i> = 18)	244 (95% CI [204; 284]) (<i>n</i> = 17)	0.42

**Fig. 2** Error plot (mean with 95% CI) for time to reach proficiency for the haptic group and the non-haptic group during the intervention and follow-up test

found that the time needed for instructor assistance was significantly higher for the haptic group compared to the non-haptic group ($p < 0.001$).

The haptic group required significantly more feedback from the instructor for the follow-up test than the assistance needed during the intervention phase ($p < 0.001$). In contrast, the non-haptic group used significantly less help from the instructor during the follow-up phase ($p < 0.001$). We found that nine out of 17 participants in the non-haptic group needed no assistance from the instructor during the follow-up test. There was no significant difference in the cumulated time spent getting instructor-based feedback ($p = 0.38$) (Table 3) (Fig. 3).

Lastly, technical malfunctions were reported and registered during both the intervention and follow-up—five incidents were documented out of 4497 attempts resulting in a technical malfunction rate of 0.1% for the entire trial. Four of

the five technical malfunctions were during the intervention; three occurred for the haptic group and one in the non-haptic group. Finally, only one technical malfunction occurred during the follow-up test using non-haptic simulators. There was no significant difference in the number of technical malfunctions between the haptic and non-haptic setup ($p = 0.10$). All technical malfunctions were solved on-site with a simple restart of the exercise.

Discussion

We found that training with a haptic compared with a non-haptic simulator reduced the time to reach proficiency in a simulation-based laparoscopy training program. Previous studies have examined the impact of haptic devices on virtual reality simulators but with contradictory results [21]. The effect of haptic devices on laparoscopic proficiency-based training programs has not been thoroughly examined, since most previous trials only tested the haptic device system over a few repetitions or a limited amount of training time before a post-test or transfer test, or did not look at the isolated influence of the haptic devices [19, 22, 24, 26, 27, 33]. We also minimized potential confounders using the same simulator software [19, 24–27, 33].

A trial by Hagelsteen et al. examined the combined effect of haptics and 3D-vision over non-haptics with 2D-vision on the performance curve of trainees and found this to accelerate skill acquisition [33]. However, 3D-vision alone has also been shown to reduce time to proficiency [32] and Hagelsteen's study could not independently estimate haptics' effect. To our knowledge, this is the first trial to examine the isolated effect of using haptic simulators on proficiency-based training. Using a proficiency-based design, compared with time- or repetition-based training, which only focuses on the initial part of the learning curve, is a strength of our trial.

During the follow-up test where both groups trained using the non-haptic setup, we found that the haptic group performed worse than the non-haptic groups, meaning that the accelerated acquisition of skills using the haptic

Table 3 Meantime required for instructor-based feedback during the intervention and follow-up test

	Haptic group	Non-haptic group	<i>p</i> values
Intervention (s)	124 (95% CI [83; 149]) (<i>n</i> = 19)	376 (95% CI [276; 468]) (<i>n</i> = 17)	0.001
Follow up (s)	343 (95% CI [243; 471]) (<i>n</i> = 18)	24 (95% CI [8; 40]) (<i>n</i> = 17)	0.001
Cumulated time (s)	466 (95% CI [320; 511]) (<i>n</i> = 18)	405 (95% CI [308; 501]) (<i>n</i> = 17)	0.38

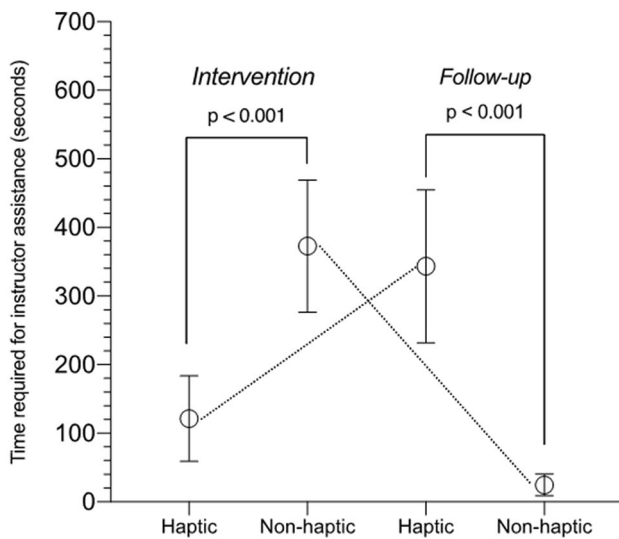


Fig. 3 Error plot (mean with 95% CI) for time required for instructor assistance for the haptic group and the non-haptic group during the intervention and follow-up test

devices could not be transferred to the non-haptic setup, as they needed significantly longer time to reach proficiency than the non-haptic group.

This can be explained using the Guidance Hypothesis [34] which argues that providing trainees with guidance or help—in our case, the haptic device—when learning psychomotor skills such as those acquired during laparoscopy simulator training can be beneficial for faster skill acquisition [35, 36]. However, participants can become dependent upon the help, which can be detrimental to long-term motor learning [34].

We chose a follow-up test where both groups trained on the non-haptic simulators till proficiency because studies on clinical transfer from laparoscopy simulator training have used non-haptic simulators [9, 13, 37]. Our results show that the trainees who practice on the simulator with a haptic device may not necessarily achieve the same skill level and have reduced transfer of skills to the conventional non-haptic system, demonstrating that the trainees may not receive the full effect of the training program when training is accelerated using haptic simulators and proficiency levels from non-haptic simulators are used. Our findings illustrate that there might be a risk using proficiency levels defined for non-haptic simulators on haptic simulators, as skill acquisition might be reduced because proficiency levels are specific to the context in which they were determined [38].

Our findings are supported by the increased need for instructor assistance in the haptic group during in the follow-up test compared with the intervention phase, even though they are solving identical tasks as during the intervention,

just on a non-haptic system. We observed that the participants from the haptic group became frustrated when transferred to the non-haptic simulator, as they perceived the tasks as more difficult, as seen by their increased need for instructor guidance. In contrast, the non-haptic group needed less instructor assistance during the follow-up test, and they trained more independently than during the intervention. During the follow-up test, we found that the participants from the haptic group spent almost the same amount of time training as during the intervention, demonstrating that to a large extent, they had to re-learn the skills necessary to reach proficiency again. This indicates the haptic device might have accelerated the participants' skills acquisition too much by making it easier for them to reach proficiency during the intervention. This could result in lower consolidation of skills learned [39].

Because training without haptics is more difficult, the non-haptic group trained longer before reaching proficiency during the intervention, which could explain the better retention and lower need for instructor feedback during the follow-up test. Making skills acquisition faster does not necessarily make it better—simulation-based training should not focus on acquiring skills the fastest way possible if this compromises the quality of the targeted skills. A potentially unrecognized benefit of a non-haptic laparoscopy simulator might be the increased difficulty when training without haptics, leading to a higher level of skills acquisition. A study conducted by Ali et al. demonstrated that increasing the difficulty of laparoscopy tasks on a virtual reality simulator increased training quality, resulting in better skills acquisition [40].

Previous studies have examined how different interventions, 3D-vision, and instructor-based feedback, affected time to proficiency in a laparoscopy training program. Both 3D-vision and instructor-based feedback reduced time to proficiency, just as the haptic device did in this study. However, these studies demonstrated no negative impact on the retention of skills after using 3D-vision or instructor-based feedback when all the participants were invited back to do a retention test [32, 36, 41]. Using haptics compared with non-haptics might have accelerated training too much, resulting in a less effective learning process.

Dismissing the potential benefits of creating a more realistic training opportunity using haptic devices should be done with caution, as haptics has been described as a paradigm shift within laparoscopic virtual reality training [42]. However, direct implementation on already-established proficiency-based training programs should be done with caution also, as they create a new task. Instead, we would advise that implementation should be done by conducting further validation studies so that the proficiency setting would be redefined.

A limitation of our study is that we did not examine transfer to a clinical setting. Hence, we cannot conclude anything of the impact of haptic simulator training on actual clinical performance.

More research into the effect of haptic simulators, whether they result in better clinical transfer, and credible newly established pass/fail-standards are needed before they can be recommended as the standard for future simulation-based training for laparoscopy.

Haptic devices are a costly add-on [43], and the use of more advanced simulators can potentially lead to a high frequency of technical malfunctions and thereby a need for technical support [44]; however, this was not the case, as technical malfunctions were very rare in our study.

Our initial assumptions regarding the time needed to reach proficiency for both groups during the intervention were not as expected. However, the effect is unaffected. There is a significant difference in the time to proficiency required in the two groups during both intervention and follow-up. For sample size calculation, assumptions were based on data from previous trials conducted by the same research group. There has been an update of the software since this trial [32] and this could potentially be the reason for the difference in training time (former version 2014 to the current version 2019), as some tasks may have become easier in newer versions. To our knowledge, no other studies have compared training on different versions of the same simulator software, and this merits further investigation in the future. Proficiency levels defined for one software version might not be applicable in a newer version of the software if it has undergone substantial changes, thereby changing the context for which it was defined [38].

Conclusion

The use of a haptic simulator reduces the time needed to reach proficiency in a proficiency-based laparoscopic training program. However, the acquired skills cannot be transferred to the conventional non-haptic simulator setup. Accelerating training too much using haptics may limit the effect of skills training if non-haptic proficiency levels are used on a haptic simulator.

Appendix

Basic skill 1: Instrument navigation.

Parameter	Requirement for proficiency level
Left instrument time (s)	<25
Left instrument misses	<2
Left instrument path length (m)	<1.4
Left instrument angular path (degrees)	<250
Right instrument time (s)	<25
Right instrument misses	<2
Right instrument path length (m)	<1.4
Right instrument angular path (degrees)	<250
Tissue Damage	<5
Maximum Damage (mm)	<10

Basic skill 2: Grasping.

Parameter	Requirements for proficiency level
Left instrument time (s)	<45
Left instrument path length (m)	<2
Left instrument angular path (degrees)	<300
Right instrument time (s)	<45
Right instrument path length (m)	<2
Right instrument angular path (degrees)	<300
Tissue damage (frequency)	<3

Basic skill 3: Lifting and grasping.

Parameter requirements for proficiency level	Requirements for proficiency level
Total time (s)	<120
Left instrument misses (%)	<60
Left instrument path length (m)	<3.2
Left instrument angular path (degrees)	<600
Right instrument misses (%)	<60
Right instrument path length (m)	<3.2
Right instrument angular path (degrees)	<600
Tissue damage (frequency)	<5
Maximum damage (mm)	<15
Grasper collided with left box (frequency)	<10
Left box lifted (frequency)	<15
Grasper collided with right box (frequency)	<10

Basic skills 4: Fine dissection.

Parameters	Requirements for proficiency level
Total time (s)	<150
Ripped or burned blood vessels	<0
Energy damaged on blood vessels (%)	<20
Ripped small vessels (%)	<25
Burned small vessels with or without stretch (%)	<25

Parameters	Requirements for proficiency level
Grasper path length (m)	<0.5
Grasper angular path (degrees)	<120
Grasper outside view (frequency)	<2
Grasper outside view (s)	<4
Cutter path length (m)	<0.9
Cutter path length (m)	<0.8
Cutter angular path (degrees)	<200
Cutter outside view (frequency)	<2
Cutter outside view (s)	<4

Procedural modul: Ectopic Pregnancy.

Parameters for the procedure: salpingectomy on the LapSim® virtual reality simulator. To reach the proficiency level all the requirements for the proficiency level must be fulfilled using correct operation technique.

Parameters	Requirements for proficiency level
Total time (s)	<280
Left instrument path length (m)	<2
Left instrument angular path (degrees)	<350
Right instrument path length (m)	<3
Right instrument angular path (degrees)	<450
Blood loss (ml)	<180
Pool of blood (ml)	<10
Ovary Diathermy damage (s)	<3
Tube Cut: Uterus distance (mm)	<10
Removed dissected tissue (Yes/No)	Yes
Bleeding vessel cut (Yes/No)	No

Declarations

Disclosures Anishan Vamadevan, Morten Stadeager, Lars Konge, Flemming Bjerrum have no conflicts of interest or financial ties to disclose.

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