



### **What was the finding?**

Overall, only **30.1%** of participants were able to identify the **most appropriate conclusion (correct direction of effect and strength of evidence)**. More students (48.2%) than practitioners (22.2%) chose the most appropriate conclusions ( $P < 0.001$ ). Looking at the four SRs separately, higher proportions of participants identified the most appropriate conclusions in SR1 and SR3, that is, the **positive SRs** (39.2% and 36.4%, respectively), compared to SR2 and SR4, that is, the negative SRs (25.0% and 20.2%, respectively). Fewer than one-half (47%) correctly identified the direction of effect against their prior beliefs. "Positive" SRs were more likely than "negative" SRs to change the participants' beliefs about the effect of the intervention (RR 1.8, 95% CI 1.3 to 2.6) and "convert" those who were previously unsure by making them choose the appropriate direction of effect (RR 1.9, 95% CI 1.3 to 2.8).

### **How much can we take out from this research/paper?**

Quite a good sample size of participants as a whole and fit the purpose of the hypothesis testing. A proper sample size estimation might be difficult without earlier studies in the similar setting but it could help in gauging a most economical sample size. Except the result on ability to interpret the most appropriate conclusions was stratified according to the different categories of participants, other results will have to be taken as indicating the whole group. On this note, the different approaches of the trimmed SRs in HCPs and medical students could exert differential effect on their performance. This added to the other uncertainties during the application such as discussion among the participants, referencing and Googling for references, etc.

The use of the 4 trimmed abstracts was unreal and its applicability to infer the same skill when reading other SRs is questionable when the full-texts are available. This might have caused the poorer performance on judging the strength of evidence. The same goes to the limit of time in assessing the abstracts. However unreal was the assessment methods, it might be the most feasible and sufficient method in judging the participants ability in interpreting the directions of effect, strengths of evidence and in reading forest plots for the treatment effectiveness. Additionally, the proficiency of participants on the (technical) language of the measures (answer sheet), relevancy of the study or the review topic to the clinical practice of some of the participants might be putting off and reduced their performance due to different backgrounds of practice.

Some of the constructs were not of sufficient clarity in their meaning and timing of measure such as the prior belief of the treatment/intervention effect, 'had seen or heard of the SRs' and 'perceptions of the value of the authors' conclusions'. These might cause them to be subjected to much personal experience of the participants and a poor measure across different participants.

The study was conducted among those who attended EBM workshops and by a mere logical deduction, the proportions of appropriate interpretation among other staffs could be lower as previous exposures and skills of EBM were not being assessed adequately. Another important question this study raises would be how the EBM workshops should be conducted to improve this skill, or what different workshop is needed. These questions will require an experimental study with a control group of people who never attended the EBM workshop or a qualitative study (approach?) to understand those who were skillful and those less skillful in interpreting SRs in order to understand the real reasons and possible causes from where interventions could improve on this. The skills in interpreting other types of clinical evidence are not included in the study [1].

### **References**

1. Munn Z, Stern C, Aromataris E, et al. What kind of systematic review should I conduct? A proposed typology and guidance for systematic reviewers in the medical and health sciences. *BMC Med Res Methodol*. 2018 Jan 10;18(1):5. doi: 10.1186/s12874-017-046.

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