PREDICTING EPIDURAL SPACE SPREAD USING ULTRASOUND COLOR DOPPLER IMAGING IN INTERLAMINAR EPIDURAL STEROID INJECTION: A PROSPECTIVE OBSERVATIONAL STUDY

YJ Kim*, H Kim, H-J Kim, WU Koh. Asan Medical Center, Seoul, Korea, Republic of

10.1136/rapm-2021-ESRA.53

Background and Aims While the use of fluoroscopy-guided transforaminal epidural steroid injection (TFESI) to help spread the injectate toward the ventral side has increased, this procedure entails a radiation risk. Recently, ultrasound has been widely applied in the medical field; among ultrasound methods, color Doppler is useful for predicting the direction of the injectate. This study describes a novel technique employing color Doppler to help predict epidural space spread in interlaminar epidural steroid injection (ILES).

Methods This observational study prospectively enrolled 35 patients scheduled for lumbar epidural steroid injection (ESI). Ultrasound-guided epidural injection was conducted and real-time images using color Doppler were recorded during injections of 5 mL of 0.1% ropivacaine containing contrast dye with dexamethasone. Fluoroscopy-guided TFESI was performed if it was difficult to perform the procedure based on ultrasound images.

Results The analysis included 30 images from 30 patients. The observed sensitivity, specificity, positive predictive value, and negative predictive values of the ultrasound color Doppler were 100%, 89.5%, 84.6%, and 100%, respectively. The agreement with ultrasound color Doppler was 93.3%.

Conclusions The main advantage of ultrasound-guided ILES is the lack of radiation exposure and contrast medium

requirement. Color Doppler may be a reliable imaging modality to predict epidural space spread during ultrasound-guided ILES. It is worth predicting the spread in the anterior epidural space (AES) by first attempting ultrasound-guided ESI. If the injectate has not spread to the AES, fluoroscopy-guided TFESI may be a good option after confirming the improvement of the patient's symptoms.

LOW-DOSE, OPIOID-FREE SUBARACHNOID BLOCK AND US-GUIDED SUPRAINGUINAL FICB AS AN ANAESTHETIC/ANALGESIC TECHNIQUE IN PATIENTS WITH ASYMPTOMATIC, MILD TO MODERATE AORTIC STENOSIS UNDERGOING HIP FRACTURE SURGERY

H Abrahams*, R Matos-Puig. GJGM Regional Hospital, Stanger, South Africa

10.1136/rapm-2021-ESRA.54

Background and Aims Aortic Stenosis (AS) is a relatively common valvular lesion in patients over 75. The significant risks associated with surgery may be mitigated by choice of anaesthetic technique and careful perioperative planning. Haemodynamic goals broadly aim to avoid hypotension and tachycardia and to preserve coronary perfusion pressure and systemic vascular resistance. Neuraxial blockade is traditionally taught to be a (relative) contraindication in this particular group of patients.

Aim: To assess perioperative pain scores and intraoperative hypotension after a technique comprising a low-dose, slow-titrated SAB and US-guided SIFI FICB in patients with mild-moderate disease as evidenced on TTE undergoing hip fracture surgery.

Methods IRB was obtained.

8 patients were enrolled prospectively over 6 months (Nov 2019 – April 2020)

All patients had a preoperative TTE to categorise disease severity.

Invasive arterial monitoring was instituted preoperatively in all patients.

US-guided SIFI FICB followed by a SAB (25G pencil point needle) was performed.

(30 ml 0.25% isobaric Bupivacaine and 7.5 mg hyperbaric Bupivacaine x 1 min, respectively).

No opioids were used.

Procedures Medical Center Institutional Review Board
101, Yongsan-ro, Yongsan-gu, Seoul 053, Republic of Korea. ☎ 02-2270-0007 FAX 02-452-0000

IRB No.	2017-08-019
TITLE	The utilization of ultrasound Doppler for the interlaminar epidural steroid injection in low back pain patients
PRINCIPAL INVESTIGATOR	Yoon Heung Kim
SPONSOR	NA
Considered Options	PMCRB has reviewed your documents and approved it on June 25, 2017
Status	APPROVED
Date of Review	26-Jun-2017
Continuing Review Report Interval	12 months

This is to certify that the information contained herein is true and correct as reflected in the records of the PMCRB Institutional Review Board. If you have any questions, contact PMCRB (PMCRB@pmc.or.kr).

pmc@pmc.or.kr

ALL PMCRB APPROVED INVESTIGATORS MUST COMPLY WITH THE FOLLOWING:

1. Conduct the research as required by the protocol.
2. Use only the Document Form bearing the PMCRB APPROVED stamp.
3. Provide your research subjects with a certified translated version of Consent Form in the subject's first language.
4. Should you request from the PMCRB of any changes to the research activity (except when necessary to protect human subjects), immediately report to the PMCRB and seek emergency changes for the protection of human subjects.
5. Report to the PMCRB the death, hospitalization, or serious illness of any study subject.
6. Promptly report to the PMCRB any new information that may adversely affect the safety of the subject or the conduct of the trial.
7. Provide reports to the PMCRB concerning the progress of the research, when requested.
8. Should you request of study discontinuation from the PMCRB before use.
9. Complete the approved consent process without coercion or undue influence, and provide the potential subject sufficient opportunity to consider whether to opt to participate.

PMCRB compliance and ethics members meet KCP & GCP-GCP requirement and all applicable laws and regulations.

Chairman: Eun, Ahnang Young 2017/07

Abstract 53 Figure 1

Abstract 54 Figure 1