Efficacy and Safety of Prophylactic Intrathecal Normal Saline for Prevention of Post Dural Puncture Headache among Women Undergoing Cesarean Section under Spinal Anesthesia: A Randomized Controlled Trial

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Keywords: PDPH, Normal Saline, Spinal Anesthesia, Intrathecal

Posted Date: June 16th, 2021

DOI: https://doi.org/10.21203/rs.3.rs-584207/v1

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Abstract

Background: Post-Dural Puncture Headache is the commonest complication of spinal anesthesia. Body of evidence revealed that Conservative management failed to show significant benefit and epidural needle and catheter techniques didn't provide conclusive evidence. On the other hand, intrathecal injection of normal saline is a simple technique and cost-effective in a resource-limited setup but it is not well examined on its effectiveness and safety profiles.

Methods and materials: After Obtaining Ethical clearance from IRB, 152 mothers scheduled for cesarean section under spinal anesthesia were allocated randomly into two groups. Data analysis was done with SPSS version 22. Descriptive statistics were run to see the overall distribution of the study subjects. Unpaired student’s T-test for continuous symmetric data and Mann-Whitney U test for non-normally distributed data were used. Categorical data were analyzed with Chi-square and fisher’s exact test where appropriate. A generalized estimating equation model was used to investigate the interaction of repeated measurements of NRS pain scores of PDPH.

Result: The overall incidence of PDPH was 29.6% while the proportion of patients who experienced PDPH was the highest among patients with control as compared to intervention (36.8% vs 22%) respectively. The GEE model revealed that the NRS pain score was 0.7, 0.4, and 0.2 unit higher at 12, 24, and 48 respectively in control as compared to the intervention.

Conclusion: prophylactic intrathecal normal saline could be an option in a resource-limited setup where the appropriate spinal needle is not accessible and management of moderate and severe PDPH is not feasible.

Registration: The protocol was registered prospectively in Clinical Trials.gov (NCT04393766).

1. Introduction

The global rate of cesarean section is increasing despite the World Health organization’s recommendation rate of cesarean section(1, 2). According to a WHO report, more than 18 million cesarean sections are performed every year(1).

Approximately, 80–90% of cesarean sections are performed with spinal anesthesia which is relatively safe as compared to general anesthesia which is accompanied by a difficult airway and risk of aspiration(3–6). However, post-dural puncture Headache is one of the commonest complications of the neuraxial block associated with penetration of dura matter and continuous flow of cerebrospinal fluid (7–14).

The International Headache Society (IHS) defines PDPH as “Headache occurring within 5 days of a lumbar puncture caused by cerebrospinal fluid (CSF) leakage through the dural puncture. It worsens 15 minutes of standing or sitting and improves after 15 minutes of lying down. It is usually accompanied by
neck stiffness and/or subjective hearing symptoms, nausea and vomiting, photophobia. It recovers spontaneously within one week, or after sealing of the leak with an autologous epidural lumbar patch within 48 hrs. There is no fever, leukocytosis, and neurologic deficit(15).”

Ninety percent of postural puncture headaches will occur within three days of dural puncture whereas sixty-six percent of the headache may happen within 48hrs of the procedure and rarely, the headache will develop between five to fourteen days after the procedure. However, some reports showed cases of post-dural puncture within 20 minutes of dural puncture and 19 months after dural puncture which were successfully managed with Epidural Blood Patch(9).

The exact mechanism of PDPH is still unknown but symptoms are generally attributed to excessive loss of CSF from the dural puncture site which results in reduced CSF pressure(12).

The lowered CSF pressure reduces the cushioning effect provided by the CSF to the brain and results in traction on intracranial pain-sensitive structures such as meningeal vessels, upper cervical and cranial nerves. The release of adenosine consequent to the sudden drop in CSF volume is also thought to cause vasodilatation of intracranial vessels and post-dural puncture headache(4).

Different factors of post-dural puncture have been mentioned in the literature among which needle size and volume of cerebrospinal fluid (CSF) lost were the independent predictors of severity of PDPH after spinal anesthesia, diagnostic lumbar puncture, or accidental dural puncture during epidural procedures(4, 7–12, 14, 16–24). The incidence of post-dural puncture headache is more likely in adults whose ages are between 30–50 years as compared to older age greater than 50 and children younger than 13 years old.

The lower weight is found to be strongly associated with the higher incidence of PDPH and cumulating evidence showed an inverse relationship between BMI and PDPH suggesting that heavier patients, in general, have higher intra-abdominal pressure, which in turn raises intra-epidural pressure and prevents cerebrospinal fluid from leaking when ADP occurs(4, 25).

Different studies showed that smokers had a considerably reduced rate of PDPH in comparison with non-smokers suggesting an inhibitory effect of tobacco smoking on PDPH that may be associated with the stimulation role of nicotine in dopamine neurotransmission(4, 15, 26).

Though pregnancy is a risk factor for PDPH due to the young age, female, sometimes sitting position, pregnancy-associated depression and anxiety, and the special popularity of regional anesthesia in this population, a meta-analysis showed that pregnancy itself does not increase the risk of PDPH. However, PDPH is associated with fatigue, sleep deprivation, and night work that lead to a higher incidence of ADP in clinical personnel when performing epidural analgesia(4).

Evidence showed that the prevalence of post-dural puncture headache in Ethiopia after spinal anesthesia was very high ranges from 38.7–42.6%(18, 23). While other studies conducted in Sub-Saharan Africa showed that the prevalence of PDPH varied from 15 to 27.5% which was strongly correlated with needle type and size (17, 27–31).
The incidence of PDPH with a randomized clinical trial conducted in Egypt on ninety obstetric patients in three groups receiving single-shot spinal anesthesia, continuous spinal anesthesia with epidural Cather and single-shot spinal with prophylactic 5ml normal saline and 10 ml normal saline before removal of the catheter after twenty-four hours was 10%, 16.6%, and 10% respectively(32).

A systemic review conducted in the United Kingdom revealed that the incidence of post-dural puncture headache after spinal anesthesia varied from 1.5%-11%. Whereas, 50% of patients developed post-dural puncture headache after accidental dural penetration with 16–18 gauges epidural needle(21).

The Cochrane database for systemic reviews identified incidence of post-dural puncture headache as 10% after spinal anesthesia, 36% after diagnostic lumbar puncture, and 81% after accidental dural puncture(9).

PDPH is getting worse with movement which leads to different undesirable consequences including delayed mother-to-child bonding, delayed breastfeeding, prolonged hospitalization, deep venous thrombosis, and thromboembolism incidents, delayed oral intake, and increase health care system cost(33).

An epidural blood patch is the known gold standard management PDPH for years when conservative management failed to be effective despite inconclusive evidence as depicted with recent systemic reviews and meta-analysis(34, 35).

Different systemic reviews and meta-analyses and randomized controlled trials revealed that conservative management for PDPH including bed rest, oral uid intake, simple oral analgesics, and caffeine failed to show significant benefit and was associated with undesirable side effects(11, 21, 34, 36–38).

A systemic review and meta-analysis by Apfel et al identified inconclusive evidence from one study with subgroup analysis. However, the quality of the included study for subgroup analysis had a small sample size, no randomization, no allocation concealment, no blinding, and proper controlling of confounders were not considered as a result both the original RCT and the systemic review recommends further RCT with high power and proper randomization(34).

A randomized clinical trial conducted to investigate the preventive benefits of intrathecal normal saline revealed a statistically significant difference in the severity and prevalence of PDPH between the groups. However, this RCT failed to assess adverse effects associated with administration of normal saline including inadequate sensory and motor block, and high spinal block and they recommend further RCT with large sample size and possible adverse effects(39).

Despite recent advancements in neuraxial techniques and different clinical and epidemiological researches on the treatment and prevention of post-dural puncture headache, the incidence of PDPH is very high and further randomized trials are in demand on its prevention approaches. Therefore, this study
aimed to investigate the efficacy and safety of prophylactic intrathecal normal saline for the prevention of PDPH after spinal anesthesia during cesarean delivery

2. Materials And Methods

2.1. Protocol and registration

This parallel double-blinded prospective randomized controlled study was conducted at Dilla University Referral Hospital from January 2020 to April 2021. Ethical clearance was obtained from Dilla University college of health sciences Institutional Review Board and it was given a Unique Identifier number(UIN:0014/19 - 11). The protocol was registered prospectively Clinical Trials.gov (NCT04393766). The objective of the study was explained and written informed consent was obtained from each participant by the data collector. Confidentiality was maintained at all levels of the study by avoiding identifiers and using codes to identify participants. The findings of the study were reported in compliance with Consolidated Standards of Reporting Trials (CONSORT) statement guidelines for randomized controlled trials(40).

2.2. Participants

Parturient who were term pregnant, ASA I and II, aged 18 to 35 years old, scheduled for cesarean section under spinal anesthesia and agreed to sign the informed consent form after comprehending the objectives of the study were included. Parturient with a history of migraine, history of PDPH, and BMI > 25 kg/m² were excluded.

2.3. Sample size determination

The required sample size was calculated using the incidence of PDPH from a previous study (16 % VS 2%) with G power software version 3.1.9.2 with the assumption of 80% power(β = 0.2) (47) that can bring a significant difference between the groups at 95% confidence interval and 5% significance level. This gave a sample size of 152 along with 10% for noncompliance.

2.4. Randomization and intervention

On the morning of surgery, every patient was re-evaluated for identification, baseline vital sign, laboratory investigation, and informed consent. Eligible patients were randomized into either treatment or control group randomly with a random sequence generated list of numbers and code which was computed with R package (Random Allocation Rule function) version 3.6.0 by the research assistant as per the schedule number given by the preoperative assessor. The list of randomly generated numbers along with the code was sealed with an opaque envelope which was later opened by the anesthetist not involved in the patient care. The data collector and the patient were blinded to the type of intervention. As per routine spinal anesthesia protocol, Group1 received 5ml intrathecal Normal saline followed by 12.5 mg of hyperbaric 0.5% Bupivacaine and Grup2 received only 12.5 mg hyperbaric 0.5% bupivacaine (Fig. 1).

2.5. Data collection tools and procedures
Data collection was carried out with pretested questionnaire which comprises Socio-demographic characteristics, preoperative patient clinical parameters, intraoperative and postoperative patient clinical variables. Data were collected with two senior Anesthetists who are not involving inpatient care. Preoperatively, the patient’s age and body mass index along with the patient’s baseline heart rate respiratory rate, and blood pressure were taken. Intraoperatively, sensory onset, peak sensory block were assessed every two minutes with pinprick until the desired level is achieved (T6-T4), and Motor block was assessed with a revised Bromage scale(41).

2.6. Outcomes and follow up

The primary outcomes of interest were incidence and severity of PDPH which were measured postoperatively at 12hrs, 24hrs, 48hrs, 72hrs, and 5th day. Most of the patients were discharged from the hospital after 48hrs but the remaining data at 72 hrs and 5th day were received with telephone contact. The secondary outcomes include sensory onset, peak sensory level, and degree of motor block, intraoperative systolic blood pressure, diastolic blood pressure, mean arterial blood pressure, heart rate, and arterial oxygen saturation every five minutes for thirty minutes. Besides, the neonatal Apgar scores, and complications related to spinal anesthesia and PDPH. Postoperatively, the incidence of post-dural puncture headache was assessed and if any, the severity was assessed with eleven points Numeric Rating Scale (NRS) at 12hrs, 24hrs, 72hrs, and 5th day. Nausea and vomiting were assessed while assessing PDPH with the Numeric Rating scale.

2.7. Data processing and Analysis

Data were entered, cleaned by Epi-info version 7, and imported to Statistical Package for Social Sciences version 22 for analysis. Descriptive statistics were run to see the overall distribution of the data. Unpaired student’s T-test for continuous symmetric data and Mann-Whitney U test for non-normally distributed data were used. Categorical data were analyzed with Chi-square and fisher’s exact test where appropriate. A generalized estimating equation model was used to investigate the interaction of repeated measurements of NRS pain scores of PDPH. The finding was reported based on the guidelines of the Consort statement. P-Value of less than 0.05 was considered as significantly associated with the outcome variable.

3. Result

3.1. Sociodemographic characteristics

As per our operational definition, the majority of participants were ASA I and most of whom had at least one cesarean section. The age of the participants varied from 18 to 38 years with a mean (± SD) of 26.3 ± 4.21 years while the weight was in the range of 50 to 102 kilogram with a mean (± SD) of 72.04 ± 9.9 kilograms (Table 1).
Table 1
Sociodemographic characteristics of participants who underwent cesarean section under spinal anesthesia in DURH, SNNPR, Ethiopia, 2021

<table>
<thead>
<tr>
<th>Variables</th>
<th>Normal saline</th>
<th>Bupivacaine alone</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>26.09 ± 3.53</td>
<td>26.52 ± 4.81</td>
<td>0.03*</td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>73.92 ± 11.28</td>
<td>70.80 ± 8.41</td>
<td>&gt;0.05</td>
</tr>
<tr>
<td>Height (cm)</td>
<td>163 ± 5.84</td>
<td>162.22 ± 5.88</td>
<td>&gt;0.05</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>27.66 ± 4.46</td>
<td>26.98 ± 3.53</td>
<td>&gt;0.05</td>
</tr>
<tr>
<td>ASA status</td>
<td></td>
<td></td>
<td>&gt;0.05</td>
</tr>
<tr>
<td>ASA I</td>
<td>11</td>
<td>65</td>
<td></td>
</tr>
<tr>
<td>ASA II</td>
<td>65</td>
<td>63</td>
<td></td>
</tr>
<tr>
<td>Parity</td>
<td></td>
<td></td>
<td>&gt;0.05</td>
</tr>
<tr>
<td>Nulliparous</td>
<td>18</td>
<td>58</td>
<td></td>
</tr>
<tr>
<td>Multiparous</td>
<td>29</td>
<td>47</td>
<td></td>
</tr>
<tr>
<td>Previous cesarean section</td>
<td></td>
<td></td>
<td>&gt;0.05</td>
</tr>
<tr>
<td>Yes</td>
<td>40</td>
<td>32</td>
<td></td>
</tr>
<tr>
<td>No</td>
<td>36</td>
<td>36</td>
<td></td>
</tr>
</tbody>
</table>

Note: ** very significant; *significant; ASA: American Society of Anesthesiology; BMI: Body Mass Index

3.2. Preoperative patient data

This study showed that the majority of indications for cesarean section was previous cesarean section scar 52(34.2%) followed by cephalopelvic disproportion 46(30.3%), Oligohydraminos 23(15.1%), and fetal distress 15(9.7%) respectively. The baseline hemodynamic parameter between the groups was comparable (Table 2).
Table 2
preoperative diagnosis and hemodynamic parameters of participants who underwent cesarean section under spinal anesthesia in DURH, SNNPR, Ethiopia, 2021

<table>
<thead>
<tr>
<th>Variables</th>
<th>Normal saline</th>
<th>Bupivacaine alone</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Indication for cesarean section</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Previous cesarean section scar</td>
<td>27(51.9)</td>
<td>25(48.1)</td>
<td>&gt; 0.05</td>
</tr>
<tr>
<td>CPD</td>
<td>26(56.5)</td>
<td>20(43.5)</td>
<td>&gt; 0.05</td>
</tr>
<tr>
<td>Oligohydramnios</td>
<td>12(52.2)</td>
<td>11(47.8)</td>
<td>&gt; 0.05</td>
</tr>
<tr>
<td>PROM</td>
<td>2(33.3)</td>
<td>4(99.7)</td>
<td>&gt; 0.05</td>
</tr>
<tr>
<td>Malpresentation</td>
<td>3(37.5)</td>
<td>5(62.5)</td>
<td>&gt; 0.05</td>
</tr>
<tr>
<td>Fetal distress</td>
<td>6(35.3)</td>
<td>11(64.7)</td>
<td>&gt; 0.05</td>
</tr>
<tr>
<td>Hematocrit (%)</td>
<td>39.59 ± 2.67</td>
<td>39.32 ± 2.79</td>
<td>&gt; 0.05</td>
</tr>
<tr>
<td>Respiratory rate</td>
<td>16.04 ± 2.29</td>
<td>17.01 ± 4.80</td>
<td>&gt; 0.05</td>
</tr>
<tr>
<td>Heart rate</td>
<td>95.78 ± 11.91</td>
<td>97.07 ± 14.80</td>
<td>&gt; 0.05</td>
</tr>
<tr>
<td>SBP</td>
<td>125.53 ± 10.28</td>
<td>124.32 ± 8.57</td>
<td>&gt; 0.05</td>
</tr>
<tr>
<td>DBP</td>
<td>77.78 ± 5.38</td>
<td>78.50 ± 4.79</td>
<td>&gt; 0.05</td>
</tr>
<tr>
<td>MAP</td>
<td>109.61 ± 7.87</td>
<td>109.04 ± 6.48</td>
<td>&gt; 0.05</td>
</tr>
<tr>
<td>SPO2</td>
<td>97.34 ± 1.11</td>
<td>97.34 ± 0.95</td>
<td>&gt; 0.05</td>
</tr>
</tbody>
</table>

Note: CPD: Cephalopelvic disproportion; PROM: Premature Rupture of Membrane; SBP: Systolic Blood Pressure; DBP: Diastolic Blood Pressure; MAP: Mean Arterial Blood Pressure; SPO2: percutaneous Oxygen saturation

3.3. Intraoperative variables

The study revealed that the time of spinal drug injection to skin incision, uterine incision, baby out time, sensory onset; surgery duration, anesthesia duration, Apgar scores, and hemodynamic parameters didn’t show a significant mean difference between the groups. Besides, the sensory onset and degree of the motor block didn’t show a significant difference between the groups (supplemental Table 1).

3.4. Incidence of post-dural puncture headache

The overall incidence of post-dural puncture was 29.6% while the proportion of patients who experienced post-dural puncture headache during five-day follow-up was the highest among patients with bupivacaine alone as compared to patients with bupivacaine and Normal saline, 36.8% vs 22% respectively. Despite a clinically significant difference, there was no statistical difference between the group on the incidence of post-dural puncture headache, p > 0.05 (Fig. 2)
The incidence of PDPH was the highest at 12 and 24 hrs between the group during the five-day follow-up period which had a significant difference at a p-value of < 0.05. However, the incidence of PDPH was almost comparable between the group at 48hrs and thereafter (Table 3).

Table 3
The proportion of patients who developed PDPH during follow-up among participants who underwent cesarean section under spinal anesthesia in Dilla University referral hospital, SNNPR, Ethiopia, 2021

<table>
<thead>
<tr>
<th>Time</th>
<th>Normal Saline</th>
<th>Bupivacaine alone</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>PDPH</td>
<td>No PDPH</td>
<td>PDPH</td>
</tr>
<tr>
<td>At 12hrs</td>
<td>9</td>
<td>67</td>
<td>19</td>
</tr>
<tr>
<td>At 24hrs</td>
<td>10</td>
<td>66</td>
<td>20</td>
</tr>
<tr>
<td>At 48hrs</td>
<td>6</td>
<td>70</td>
<td>8</td>
</tr>
<tr>
<td>At 72hrs</td>
<td>6</td>
<td>70</td>
<td>2</td>
</tr>
<tr>
<td>At 84hrs</td>
<td>3</td>
<td>73</td>
<td>4</td>
</tr>
</tbody>
</table>

NOTE: ** Very significant; *significant; PDPH: Post Dural Puncture Headache

3.4.1. Severity of PDPH

The severity of PDPH was measured with a Numeric rating scale at each follow-up period. The study revealed that the mean severity score was the highest for the first 48hrs and decreasing thereafter in both intervention and control group (Fig. 3)

3.4.2. Repeated measure interaction

The generalized Estimating Equation (GEE) model was employed to investigate the interaction of NRS pain score rating repeated measurements during the five-day follow-up. The GEE model was fitted after restructuring the NRS pain score for each measurement interval. The GEE model revealed that the NRS pain score was 0.7, 0.4, and 0.2 unit higher at 12, 24, and 48 respectively in the Bupivacaine group as compared to the Normal Saline group (Table 5).
Table 4
Repeated measure interaction estimation with generalized estimation equation for NRS pain score at different periods during follow up

<table>
<thead>
<tr>
<th>Parameter</th>
<th>β</th>
<th>Std. Error</th>
<th>95% Wald Confidence Interval</th>
<th>Hypothesis Test</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td>Lower</td>
<td>Upper</td>
</tr>
<tr>
<td>(Intercept)</td>
<td>.086</td>
<td>.0347</td>
<td>.017</td>
<td>.154</td>
</tr>
<tr>
<td>NRS 12hrs</td>
<td>.709</td>
<td>.1601</td>
<td>.395</td>
<td>1.023</td>
</tr>
<tr>
<td>NRS 24hrs</td>
<td>.480</td>
<td>.1105</td>
<td>.264</td>
<td>.697</td>
</tr>
<tr>
<td>NRS 48hrs</td>
<td>.276</td>
<td>.1189</td>
<td>.043</td>
<td>.509</td>
</tr>
<tr>
<td>NRS 72hrs</td>
<td>.046</td>
<td>.0591</td>
<td>-.070</td>
<td>.162</td>
</tr>
<tr>
<td>NRS 5TH day</td>
<td>0a</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

3.4.3. Complication

There was no apparent complication in either of the intervention or control group during the follow-up period. However, there were mild to moderate nausea and vomiting, photophobia, and neck stiffness in participants with post-dural puncture headaches where nausea and vomiting were the most frequent complication. The overall incidence of this complication was 38.9% while the incidence of nausea and vomiting was relatively higher in the bupivacaine group as compared to the normal saline group (28.9% vs 38.8%).

4. Discussion

This double Randomized controlled trial was conducted to investigate the effectiveness of prophylaxis intrathecal normal for prevention of post-dural puncture headache in parturient underwent cesarean section under spinal anesthesia.

Post Dural Puncture Headache is the most common complication of neuraxial anesthesia which has been postulated that it is resulting from intracranial hypotension due to continuous loss of CSF through the dural tear that is mainly related to needle size, shape, type, and less likely risk factor including female gender, young age, pregnancy, low BMI, previous history of PDPH, and history of migraine headache(4, 12, 20, 24, 38, 42, 43). In turn, these causes prolonged bedridden and thrombosis, decrease early breastfeeding and mother-to-child bonding, length of hospital stay, increased psychological distress, and dissatisfaction to the mother and her family(13, 16, 43–45).

This study showed that the Sociodemographic characteristics, preoperative baseline variables, sensory onset, peak sensory level, degree of motor block, intraoperative maternal hemodynamics, and Apgar scores were comparable between the groups. However, there were clinical and statistical mean differences between the groups on fifth minutes diastolic and the mean arterial blood pressure which
may be related to less sympathetic blockage among patients with normal saline due to dilution of bupivacaine.

This study revealed that the overall incidence of PDPH is very high which is comparable with other studies (16, 20, 38, 45–47). Other studies showed that the incidence of PDPH is very high among patients with control compared to an epidural and intrathecal normal saline injection unlike this study where there is no significant difference in the incidence of PDPH despite a clinically significant PDPH was observed among the control group in the first 24 hrs period (20, 46).

This study showed that the incidence of PDPH among intervention and control groups didn’t show a significant difference which is in line with other randomized control trials and systematic review and meta-analysis (16, 34, 45). However, other studies did show a significant reduction of PDPH among patients who received prophylactic intrathecal normal saline (46, 47). This discrepancy may be related to heterogeneity in the study population, variation in the type, size, and shape of spinal needle, volume of prophylactic normal saline.

In this study, patients experienced mild to moderate pain which was managed with conservative management, acetaminophen, ibuprofen, and tramadol, and no severe pain was reported. The study revealed that the mean severity score was the highest for the first 48 hrs and decreasing thereafter in both intervention and control groups which are comparable with other randomized trials (43, 47, 48).

In this study, a generalized estimating equation was used after adjusting the NRS pain score to explore the interaction in each repeated measurement. As a result, the GEE model revealed that the NRS pain score was 0.7, 0.4, and 0.2 units higher at 12, 24, and 48 respectively in the Bupivacaine group as compared to the Normal Saline group.

There were mild to moderate nausea and vomiting, photophobia, and neck stiffness in participants with post-dural puncture headache where nausea and vomiting were the most frequent complication which is comparable to other studies conducted globally on this topic.

4.1. Strength of the study

This study is a double-blind randomized trial with a relatively large sample size with a long duration of flow up which is about five days postoperatively. The study also tried to investigate the postulated adverse effects of prophylactic intrathecal normal saline including high spinal block, low motor block, and blindness related to rapid injection and obstruction of retinal smaller blood vessels.

4.2. Limitation of the study

The study was started before the COVID-19 pandemic which caused many problems where the patient flow was very low, hyperbaric bupivacaine and smaller spinal needle were not available during the lockdown. Besides, we had no PCR machine for diagnosis of COVID-19 and some of the patients might have the disease where the accurate diagnosis of PDPH was a challenge.
4.3. Implication for practice

The independent predictor of PDPH is the spinal needle size, type, and shape but it could happen even with smaller spinal needles and it is inevitable in accidental dural puncture during epidural anesthesia. PDPH is a common complication of neuraxial anesthesia which has a great impact on mother, newborn, family, and health care system particularly in patients with moderate and severe PDPH tolerant to conservative management, because the current treatment and preventive modalities are not conclusive and innovative research on preventive strategy is in demand.

4.4. Implication for further research

The study showed a clinical benefit of prophylactic intrathecal normal saline for the prevention of PDPH at least for 48hrs. However, this study used 5ml normal saline where some of the patients might have COVID-19 and/or more likely vulnerable to different psychological distress which limited our inferences to the general population. The authors recommend further multi-center randomized clinical trials placebo, 5ml, and 10ml normal saline for surgery under spinal and epidural anesthesia.

5. Conclusion

Prophylactic Intrathecal administration of normal saline for prevention of PDPH in patients undergoing cesarean delivery under spinal anesthesia is a safe and simple technique. This study failed to show the effectiveness of prophylactic intrathecal normal saline for the prevention of PDPH for a prolonged period of up to a week and further studies are required to provide a firm conclusion. However, prophylactic intrathecal normal saline could be an option in a resource-limited setup where the appropriate spinal needle is not accessible and management of moderate and severe PDPH is not feasible. We are not recommending the routine use of normal saline for the prevention of PDPH in a setting where small spinal needles are accessible.

6. Declarations

Ethics approval and consent to participate

Ethical clearance and approval were obtained from the ethical review board of the College of Health Science and Medicine.

Consent for publication

Not applicable

Availability of data and materials

Data and material can be available where appropriate.

Competing interests
The authors declare that there are no competing interests

**Funding**

This research was funded by Dilla University with a total amount of 65,480 ETB

**Acknowledgments**

The authors would like to acknowledge Dilla University for financial support and encouragement to carry out the project.

**7. References**


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Figures
Figure 1

consort flow diagram
Figure 2

Incidence of PDPH among patients who underwent cesarean section under spinal anesthesia with bupivacaine alone and bupivacaine with 5ml Normal Saline in Dilla University Referral Hospital, 2021.
Figure 3

Mean severity of PDPH at each follow-up period among participants who underwent cesarean section under spinal anesthesia in Dilla University referral hospital, Ethiopia, 2021.

Supplementary Files

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- CONSORT2010Checklist.doc
- supplementntable.docx