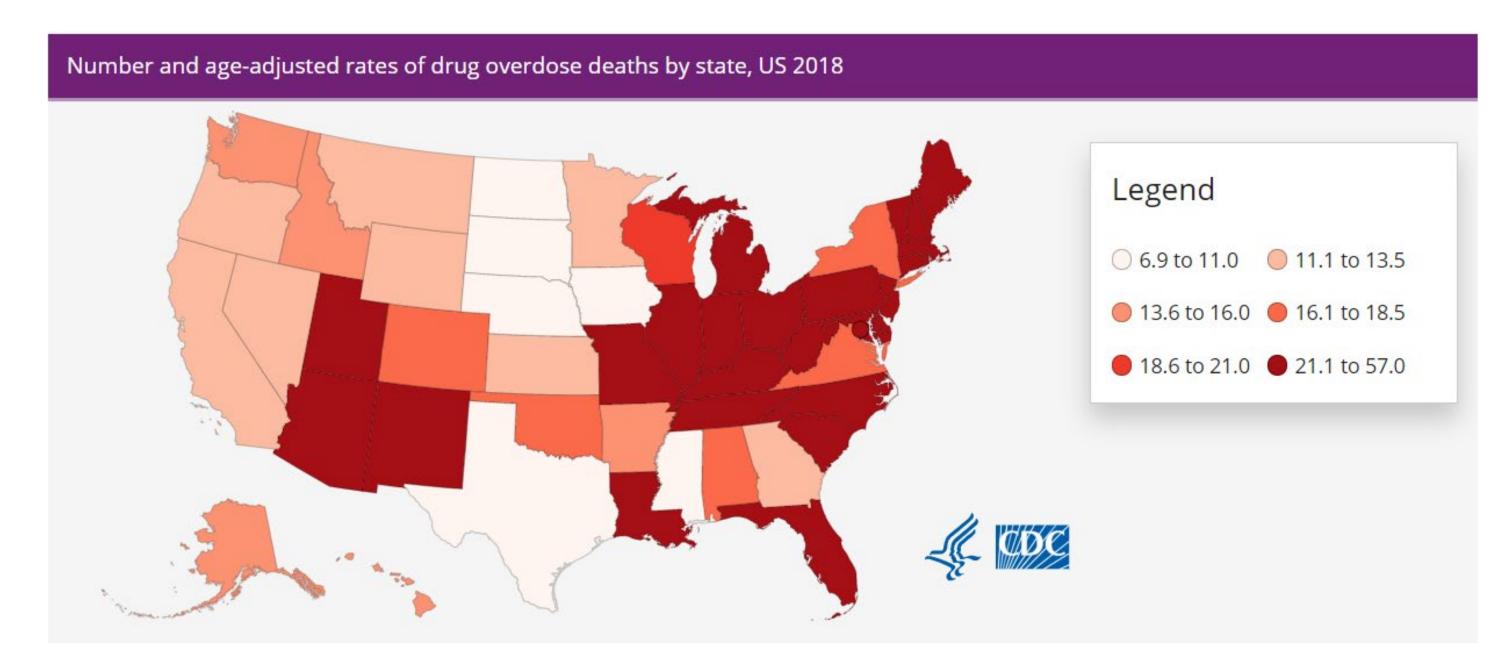
Local wound infiltration with liposomal bupivacaine decreases post-cesarean pain scores



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ABSTRACT & INTRODUCTION

Surgical patients are routinely prescribed opioid analgesics that are strongly associated with drug overdose deaths.



- Liposomal bupivacaine is a long-acting local anesthetic that received FDA approval in 2011 for • local surgical site infiltration to promote postoperative analgesia
- This anesthetic has demonstrated an improvement in post-operative pain control in various surgical realms, indicating a decreased need for opiates and subsequently decreased risks. However, few studies have investigated its use at the time of cesarean delivery.

METHODS

- Retrospective study of patients who underwent cesarean delivery from May-December 2019 at Miami Valley Hospital in Dayton, OH
- Inclusion criteria: Age 18-45, BMI <60
- Liposomal bupivacaine group (n=57) received intraoperative liposomal bupivacaine in addition to standard of care analgesics, which includes intrathecal or epidural morphine
- Control group (n= 59) underwent C section within 1 week of LBG counterpart and within 3 weeks of gestational age of LBG counterpart
- Statistical analysis was completed using IBM SPSS Statistics

PRIMARY OUTCOMES

- Patient opiate use in morphine-equivalent doses (MED) from 24-48 hours postoperatively
- Patient pain ratings via 11-point Likert Scale averaged every 12 hours

Patients who received liposomal bupivacaine during C-section had significantly lower pain scores 0-24 hours after surgery.

RESULTS

Table 1. Morphine equivalent dosing (MED) for post-operative timeframes

| Outcome | Control (N=57) | Liposomal Bupivacaine (n=59) | \boldsymbol{P} |
|---------|----------------|------------------------------|------------------|
| 0-6 h | 15.1 (21.1) | 10.3 (13.0) | 0.15 |
| 6-12 h | 5.6 (71.) | 4.8 (6.6) | 0.54 |
| 12-24 h | 18.9 (28.2) | 13.6 (14.6) | 0.21 |
| 24-48 h | 44.4 (82.2) | 31.3 (29.7) | 0.26 |

Data are mean (SD) MED averaged over the 12 hour period

Table 2. Pain scores for post-operative timeframes

| Outcome | Control (N=57) | Liposomal Bupivacaine (n=59) | P |
|----------------------|----------------|------------------------------|--------|
| 0-12 h | 4.7 (1.9) | 1.5 (1.7) | < 0.01 |
| 12-24 h | 4.4 (2.3) | 2.1 (1.9) | < 0.01 |
| 24-36 h | 3.7 (2.0) | 3.7 (1.7) | 0.91 |
| 36-48 h | 4.4(2.1) | 4.4 (1.6) | 0.92 |
| 48-60 h ^a | 3.8 (2.0) | 5.3 (1.7) | < 0.01 |
| 60-72 h ^b | 4.0 (2.2) | 5.6 (1.6) | < 0.01 |
| 72-84 h ^c | 3.5 (2.1) | 5.9 (1.4) | < 0.01 |
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Data are mean (SD) pain scores averaged over the 12 hour period

CONCLUSIONS & FUTURE DIRECTIONS

- Confounding factors include differences between post-operative opiate pain medication administration by nursing care which could lead to lack of significant differences in opiate use.
- Future directions include a prospective randomized control trial emphasizing patient and healthcare worker education on post-operative analgesic usage. Additional studies should assess differences between liposomal and systemic bupivacaine

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^a Pain scores available for 49 and 52 patients in the CG and LBG, respectively.

b Pain scores available for 38 patients in each group.

^c Pain scores available for 27 and 29 patients in the CG and LBG, respectively.