

reference	participants' characteristics	intervention group/ control group	outcomes	critical appraisal/ conclusion																																																																																																										
<p><a href="#">Cui et al. 2010</a> Systemic administration of lidocaine reduces morphine requirements and postoperative pain of patients undergoing thoracic surgery after propofol-remifentanyl-based anaesthesia. Eur J Anaesthesiol. 2010 Jan;27(1):41-6.</p>	<p><b>inclusion criteria</b></p> <ul style="list-style-type: none"> <li>- ASA physical status I-II</li> <li>- age 18–65 yrs</li> <li>- undergoing thoracic surgery lasting 3–6 h between 1 Jan to 31 Jul 2008</li> </ul> <p><b>exclusion criteria</b></p> <ul style="list-style-type: none"> <li>- chronic pain</li> <li>- taking analgesics or opioids within 7 days of surgery</li> <li>- history of drug or alcohol abuse</li> <li>- psychiatric disorder</li> <li>- obesity</li> <li>- acute CV disorder</li> <li>- CNS disease</li> <li>- inability to communicate with investigator</li> <li>- contraindication(s) to propofol, opioids, lidocaine or to self administration of morphine via PCA</li> <li>- surgery time &gt;6 h- immediate extubation was not planned after surgery</li> </ul> <p><b>demographic data:</b></p> <table> <thead> <tr> <th></th> <th>group L</th> <th>group C</th> </tr> </thead> <tbody> <tr> <td>age (yrs)</td> <td>54(9)</td> <td>51(10)</td> </tr> <tr> <td>weight (kg)</td> <td>65(9)</td> <td>60(10)</td> </tr> <tr> <td>height (cm)</td> <td>168(8)</td> <td>170(8)</td> </tr> <tr> <td>sex (m/f)</td> <td>13/7</td> <td>13/7</td> </tr> <tr> <td>ASA status (I/II)</td> <td>6/14</td> <td>5/15</td> </tr> </tbody> </table> <p><b>patient flow and follow up:</b></p> <p><u>total patient number included:</u> 45</p> <p><u>randomised in:</u> group L: 20 group C: 20</p> <p><u>excluded:</u> 5</p> <p><u>analysed:</u> 40</p> <p><u>follow-up:</u> 2,6,12,24,36,48 h postop</p>		group L	group C	age (yrs)	54(9)	51(10)	weight (kg)	65(9)	60(10)	height (cm)	168(8)	170(8)	sex (m/f)	13/7	13/7	ASA status (I/II)	6/14	5/15	<p><b>intervention prior to anaesthesia</b></p> <ul style="list-style-type: none"> <li>- group L: IV lidocaine (33.0 µg/kg/min) at induction, stopped at skin closure</li> <li>- group C (control): IV physiological saline</li> </ul> <p><b>mode of anaesthesia</b></p> <ul style="list-style-type: none"> <li>- propofol-remifentanyl (targeted cible)</li> </ul> <p><b>surgical approach</b></p> <p>Procedure (n)</p> <p><i>Pulmonary lobectomy:</i> Group L 7 Group C 6</p> <p><i>Oesophagectomy:</i> Group L 9 Group C 9</p> <p><i>Cardiectomy:</i> Group L 4 Group C 5</p> <p><b>postoperative analgesia</b></p> <ul style="list-style-type: none"> <li>- in PACU: Titration of morphine (1.0–2.0 mg/2 min) were given to keep the VRS-4 &lt;2.</li> <li>- after 2h postop: PCA morphine, 1 mg with 5 min lo, for 48 h post op</li> </ul>	<p><b>postoperative pain [VRS-4 in the first 2 h 0=no pain, 1= slight pain, 2= moderate pain, 3= intense or severe pain]</b></p> <table> <thead> <tr> <th></th> <th>group L</th> <th>group C</th> <th></th> </tr> </thead> <tbody> <tr> <td>at 30 min</td> <td></td> <td></td> <td></td> </tr> <tr> <td>≤ 1</td> <td>16</td> <td>10</td> <td></td> </tr> <tr> <td>≥ 2</td> <td>4</td> <td>10</td> <td>S</td> </tr> <tr> <td>at 90 &amp; 120 min</td> <td></td> <td></td> <td></td> </tr> <tr> <td></td> <td></td> <td>(No data)</td> <td>NS</td> </tr> </tbody> </table> <p><b>postoperative pain [VAS (mm): mean±SD]</b></p> <table> <thead> <tr> <th></th> <th>group L</th> <th>group C</th> <th></th> </tr> </thead> <tbody> <tr> <td>at 6 h</td> <td></td> <td></td> <td></td> </tr> <tr> <td></td> <td>23.1±15.6</td> <td>34.2±15.4</td> <td>S</td> </tr> <tr> <td>from 6-48 h</td> <td></td> <td></td> <td></td> </tr> <tr> <td></td> <td>(no data)</td> <td>(no data)</td> <td>NS</td> </tr> </tbody> </table> <p><b>morphine consumption in the first 2 h (mg): mean±SD</b></p> <table> <thead> <tr> <th></th> <th>group L</th> <th>group C</th> <th></th> </tr> </thead> <tbody> <tr> <td>at 30 min</td> <td>0 (0–2.8)</td> <td>6 (0.3–11.3)</td> <td>S</td> </tr> <tr> <td>30–60 min</td> <td>0 (0–0.75)</td> <td>1.5 (0–4)</td> <td>S</td> </tr> <tr> <td>total amount of morphine given 0–120 min</td> <td>0 (0–4)</td> <td>8 (1.25–15.)</td> <td>S</td> </tr> </tbody> </table> <p><b>PCA morphine consumption (mg): mean±SD</b></p> <table> <thead> <tr> <th></th> <th>group L</th> <th>group C</th> <th></th> </tr> </thead> <tbody> <tr> <td>during 2–6 h (mg)</td> <td></td> <td></td> <td></td> </tr> <tr> <td></td> <td>2 (0–3.)</td> <td>4 (2–5.8)</td> <td>S</td> </tr> <tr> <td>during 6 to 48 h (mg)</td> <td></td> <td></td> <td></td> </tr> <tr> <td></td> <td>11 (8.3–26.5)</td> <td>20.5 (10–29)</td> <td>NS</td> </tr> <tr> <td>total doses of morphine given in 48 h</td> <td></td> <td></td> <td></td> </tr> <tr> <td></td> <td>12 (9.3–28)</td> <td>30.5 (11–47.8)</td> <td>NS</td> </tr> </tbody> </table>		group L	group C		at 30 min				≤ 1	16	10		≥ 2	4	10	S	at 90 & 120 min						(No data)	NS		group L	group C		at 6 h					23.1±15.6	34.2±15.4	S	from 6-48 h					(no data)	(no data)	NS		group L	group C		at 30 min	0 (0–2.8)	6 (0.3–11.3)	S	30–60 min	0 (0–0.75)	1.5 (0–4)	S	total amount of morphine given 0–120 min	0 (0–4)	8 (1.25–15.)	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