

**Table 2. Secondary outcomes: Dosing, Opioid Use and Dose associated adverse events in the DEX and Non-DEX Groups**

<b>Outcome measure</b>	<b>DEX Group (n=39)</b>	<b>No DEX Group (n=39)</b>	<b>p value</b>
<b>DOSING</b>			
<b>1. Dexmedetomidine, mcg/kg/hr, mean ±SD</b>	<b>0.36 ± 0.12</b>	<b>---</b>	<b>---</b>
<b>2. Morphine, mcg/kg/hr, mean ±SD</b>	<b>33 ± 18.2</b>	<b>26.7 ± 12.3</b>	<b>0.13</b>
<b>3. Fentanyl, mcg/kg/hr, mean (range)</b>	<b>1.5</b>	<b>1(1-2)</b>	<b>---</b>
<b>4. Duration of infusion, hr, median (IQR)</b>	<b>91.5 (64.25-135.25)</b>	<b>67.25 (44.5-170.5)</b>	<b>0.9</b>
<b>Supplemental opioid dosing, n (%)</b>	<b>35 (90)</b>	<b>35 (90)</b>	<b>1</b>
<b>Morphine</b>	<b>34 (97)</b>	<b>28 (80)</b>	<b>0.13</b>
<b>Fentanyl</b>	<b>1(3)</b>	<b>11 (31.4)</b>	<b>0.28</b>
<b>Duration of supplemental dosing, days, median (IQR)</b>	<b>6 (4-8)</b>	<b>7 (4-10)</b>	<b>0.32</b>
<b>Supplemental doses per patient, n (%)</b>	<b>11.6 (13)</b>	<b>15.8 (18.6)</b>	<b>0.35</b>
<b>Patients requiring withdrawal scoring, n (%)</b>	<b>8 (20)</b>	<b>13 (33)</b>	<b>1</b>
<b>Total opioid dose, mcg/kg, median (IQR)</b>	<b>1155 (450-4665)</b>	<b>1841.25 (1291.5-7442.5)</b>	<b>0.01</b>