Nociception Level (NoL) index alteration after standardized nociceptive stimulus decreases with higher doses of remifentanil

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Background: Several indexes have been recently used to monitor nociception intensity during general anesthesia (GA). The PMD-100 monitor (Medasense Biometrics, Israel) is a novel monitor of nociception, presenting the Nociception Level (NoL) Index. The NoL index is a multiparametric index derived from heart rate (HR), HR variability, plethysmograph wave amplitude, skin conductance and its fluctuations. This index ranges from 0 to 100, with lower value meaning lower pain intensity. We tested the NoL alteration during a standardized noxious stimulus at various doses of remifentanil (RF) i.v. infusion, with the hypothesis that the higher the RF dose, the lower the NoL alteration.

Methods: 40 patients received desflurane-RF based GA with an epidural analgesia for laparotomy. A moderate noxious stimulus (electrical stimulation 70mA, 100Hz, 30sec) was applied to the forearm of the patients at 4 RF doses varying from 0.005 to 0.15 µg·kg⁻¹·min⁻¹. For each RF dosage, the pre- and the post-stimulation NoL peak values, the difference (ΔNoL) and HR were recorded and compared using a linear mixed effects model and CI of 95%. Study # NCT02602379.

Results: The median pre stimulation NoL basal values ranged for 3 to 5 with no significant difference when RF infusion increased. The median post stimulation values at RF doses of 0.005, 0.05, 0.1 and 0.15 µg·kg⁻¹·min⁻¹ were, respectively, 39, 15, 8 and 8.5, showing statistical significant difference between 0.005 and 0.05 µg·kg⁻¹·min⁻¹ (p<0.0001) and between 0.05 and 0.1 µg·kg⁻¹·min⁻¹ (p<0.01) but not at higher doses e.g. between 0.1 and 0.15 µg·kg⁻¹·min⁻¹ (p=0.49). Accordingly, the ΔNoL was greater at 0.005 than at 0.05 µg·kg⁻¹·min⁻¹ (6.5 vs 3, p=0.0001) and at 0.05 than at 0.1 µg·kg⁻¹·min⁻¹ (6.5 vs 3, p=0.01) but showed no difference between 0.1 and 0.15 µg·kg⁻¹·min⁻¹ (3 vs 3.5, p=0.70). The correlation between RF dosage and post-stimulation NoL value was higher (p=0.61) than with HR (p=0.40).

Conclusion: When a patient is exposed to a standardized noxious stimulation, NoL reaches higher peak values and shows greater alteration when the patient is receiving lower doses of RF. In this study, a plateau of minimal NoL variation was reached at 0.1 µg·kg⁻¹·min⁻¹ of RF. The NoL index showed greater correlation with RF dose than HR did. These results suggest great potential of the NoL index as a tool to monitor nociception intensity during anesthesia. However, further studies are needed to assess whether the NoL index can better guide intraoperative opioid administration and monitor the nociception level than classic clinical signs like HR or blood pressure.