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TITLE: A COMPARISON OF TWO EPIDURAL ALPHA-2-AGONISTS, GUANFACINE AND CLONIDINE, IN REGARD TO DURATION OF ANTINOCICEPTION, AND

VENTILATORY HEMODYNAMIC AND

EFFECTS

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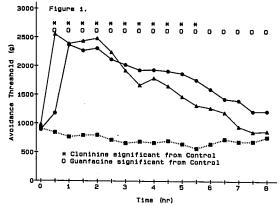
Epidural clonidine(C) produces analgesia in humans with acute and chronic pain. 1.2 However, it's use appears to be limited because of short lasting analgesia, hemodynamic depression, sedation and respiratory depression. A comparison of intrathecal guanfacine(G) and C has shown G to have longer duration of action. This study compares these two agents epidurally.

Approval was obtained from the Animal Care Committee. 5 Nubian goats, 45-55 kg, were instrumented with pulmonary and carotid artery and epidural catheters. Each goat received epidural injections of G 5 mg in 10 ml normal saline(NS), C 750 mcg in 10 ml NS, and a 10 ml NS control on 3 separate days, in an open label fashion. Antinociception [avoidance threshold, (AT)] was tested by using a point [avoidance threshold, (AT)] was tested by using a point pressure device⁵ attached to the goats hind limb in which force was gradually applied until limb withdrawal. AT, blood pressure (BP), heart rate (HR), cardiac output (CO), wedge pressure (PCWP), central venous pressure (CVP), and arterial and venous blood gases were measured at baseline and then every 30 minutes after epidural injection for 8 hrs. Statistical significance (p<0.05) was determined with an analysis of variance with repeated measure design.

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G and C produced similar increases in the AT, however, G lasted significantly longer than C (G-8 hrs. vs. C-5.5 hrs)(Fig.1). Increases in pCO, were greater and much more frequent in the C group. There were no marked differences between G and C in regard to decreases in HR, BP, CO, PCWP, CVP, or venous admixture during periods of increased AT. Epidural pretreatment with the alpha-2-antagonist idazoxan 30 min prior to epidural guanfacine prevented the

increase in AT.



To our knowledge this is the first study to document the antinociceptive effects of epidural Because of longer duration antinociception and less respiratory depression, epidural guanfacine may be a superior drug for postoperative analgesia and chronic pain syndromes. References:

- Eisenach, JC, et al., Anesthesiol 71:640-646, 1989. Eisenach, JC, et al., Anesthesiol 71:647-652, 1989. 2. 3.
- Penon, C, et al., Anesthesiol 71:A649, 1989. 4. Post, C, et al., Anesth Analg 66:317-24, 1987.
- Nolan, A, et al., J of Pharm Meth 17:39-49, 1987.

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TITLE: EFFECTS OF CLONIDINE ON C-FIBER ACTION

POTENTIAL, INTERACTION WITH LIDOCAINE AUTHORS: D. Gaumann, M.D.; P. Brunet, Ph.D.*; P. Jirounek,

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Administration of clonidine together with lidocaine for peripheral nerve block significantly prolongs the local anesthetic (LA) effect, possibly by a direct action on the nerve (1, 2). The present study examined the effect of clonidine and its interaction with lidocaine on C-fiber action potential in the vagal nerve preparation of the rabbit. Rabbit vagus nerves were desheathed and mounted in a sucrose-gap apparatus. The stimulation chamber was either perfused with Locke solution or with 5x10-4 M lidocaine. Various concentrations of clonidine (5x10-8 M to 5x10-3 M) were either added to the Locke or lidocaine solution. The preparation was stimulated by single supramaximal square wave pulses of 0.4 msec. The compound action potential (AP) of the C-fiber component was analyzed with regard to area, peak amplitude and time interval from stimulation to peak.

Clonidine added to Locke solution, at concentrations from 5x10-8 M to 5x10-5 M, had no effect on AP, while concentrations of 5x10-4 M and 5x10-3 M significantly decreased AP area (Table 1). When clonidine was added to a lidocaine solution (5x10-4 M), a low concentration of 5x10-7 M significantly decreased AP area. Clonidine at concentrations of 5x10-4 M and 5x10-3 M, in the presence of lidocaine, led to a significant decrease in AP area as compared to clonidine treatment alone (Table 1). Local anesthetic effects of clonidine (5x10-4 M) were compared to lidocaine (5x10-4 M, Table 2). There was no difference between the two drugs with regard to their effects on AP area. Clonidine evoked a less pronounced decrease in AP amplitude and delay of AP peak than lidocaine.

Clonidine alone exhibits LA effects above concentrations of 5x10-5 M. LA effects of clonidine (5x10-4 M) do not differ from those of lidocaine (5x10-4 M) with regard to the probability of fiber blockade. as indicated by AP area. Clonidine, however, slows fiber conduction less than lidocaine, resulting in a higher peak AP. In the presence of lidocaine, clonidine has synergistic local anesthetic effects at concentrations of 5x10-7 M and above 5x10-5 M. The synergistic effect of the low concentration of clonidine (5x10-7 M) with lidocaine (5x10-4 M), might explain the clinical observation that clondine prolongs the action of lidocaine in peripheral nerve block. where it is administered at approximately 1000-fold lower concentrations than lidocaine (1, 2).

References: 1) Anesthesiology 71: A644, 1989

2) Anesthesiology 73: A821, 1990

Effects of clonidine on area of action potential (AP) in the presence of Locké or lidocaine solution Table 1:

Pretreatment	Concentration of Clonidine					
	5x10 ⁻⁸ M	5x10-7 M	5x10 ⁻⁸ M	5x10 ⁻⁵ M	5x10-4 M	5x10-3 M
Locke	96.9±3.6 (n=5)	103±3.0 (n=5)	97.0±2.3 (n=6)	88.5±5.9 (n=7)	83.1±3.4† (n=11)	71.9±3.2† (n=5)
Lidocaine	98,1±3,5	86.1±5.6*†	95,5±4.9	89.0±9.7	63.2±8.8*1	17.5±4,8°
(5×10-4M)	(n=4)	(n=5)	(n=5)	(n=5)	(n=4)	(n=4)

Effects of clonidine expressed as % of pretreatment value (mean \pm SE). $^{\circ}p<0.05$ between group comparison, $\uparrow p<0.05$ within group comparison to pretreatment value.

Table 2: Effects of clonidine or ildocaine at equimolar concentration (5x10-4 M) on action potential (AP)

Drug	AP Peak Amplitude (%)	AP Area (%)	AP Peak Delay (msec)
Lidocaine (n=8)	41.3 ± 3.1†	75.8 ± 9.4	8.4 ± 0,9†
Clonidine	58.5 ± 2.4*†	82.2 ± 5.9	2.6 ± 0.4*†

Drug effects as % or difference (mean ± SE) of control value with Locke solution. *p< 0.05 between group comparison. †p<0.05 within group comparison to control value.