Neuropeptides in the posterodorsal medial amygdala modulate central cardiovascular reflex responses in awake male rats

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Abstract

The rat posterodorsal medial amygdala (MePD) links emotionally charged sensory stimuli to social behavior, and is part of the supramedullary control of the cardiovascular system. We studied the effects of microinjections of neuroactive peptides markedly found in the MePD, namely oxytocin (OT, 10 ng and 25 pg; n=6/group), somatostatin (SST, 1 and 0.05 µM; n=8 and 5, respectively), and angiotensin II (Ang II, 50 pmol and 50 fmol; n=7/group), on basal cardiovascular activity and on baroreflex- and chemoreflex-mediated responses in awake adult male rats. Power spectral and symbolic analyses were applied to pulse interval and systolic arterial pressure series to identify centrally mediated sympathetic/parasympathetic components in the heart rate variability (HRV) and arterial pressure variability (APV). No microinjected substance affected basal parameters. On the other hand, compared with the control data (saline, 0.3 mL; n=7), OT (10 ng) decreased mean AP (MAP₅₀) after baroreflex stimulation and increased both the mean AP response after chemoreflex activation and the high-frequency component of the HRV. OT (25 pg) increased overall HRV but did not affect any parameter of the symbolic analysis. SST (1 µM) decreased MAP₅₀, and SST (0.05 µM) enhanced the sympathovagal cardiac index. Both doses of SST increased HRV and its low-frequency component. Ang II (50 pmol) increased HRV and reduced the two unlike variations pattern of the symbolic analysis (P<0.05 in all cases). These results demonstrate neuropeptidergic actions in the MePD for both the increase in the range of the cardiovascular reflex responses and the involvement of the central sympathetic and parasympathetic systems on HRV and APV.

Key words: Baroreceptor and chemoreceptor reflexes; Power spectral analysis; Symbolic analysis; Oxytocin; Somatostatin; Angiotensin II

Introduction

The posterodorsal medial amygdala (MePD) is a forebrain subcortical component of the “extended amygdala” (1) with a well-described role in the response to stressful stimuli (2), hypothalamic neuroendocrine secretion (3), and the display of social behaviors, such as aggressive and reproductive behaviors in rats (4,5). The MePD is important for higher-level, supramedullary control of cardiovascular functions (6-8), and, in this sense, sends projections to the central amygdaloid nucleus (CeA) and then to the A1 noradrenergic cells of the caudal ventrolateral medulla and the nucleus of the solitary tract (NTS; 9,10). The MePD is also interconnected with other medial amygdala subnuclei that innervate cardiovascular-related areas of the hypothalamus (9,10) and the sympathetic/parasympathetic-related parts of the periventricular hypothalamic zone (11). It is highly likely that this integrated network modulates cardiovascular responses to ensure appropriate hemodynamic responses concomitant with the expression of social behaviors (7,8,12).

The involvement of the MePD in the regulation of heart rate (HR) and arterial blood pressure (AP) were reported under different methodological conditions. For example, electrical stimulation of the MePD increased AP in mice (13). Activation of the whole mouse medial amygdala (MeA) after fear or aversive stress led to sympathetic activation...
and neurogenic hypertension (14,15), whereas bilateral neurotoxic lesions of the posterior part of the MeA attenuated the development of high AP levels in spontaneously hypertensive rats (16). Moreover, central control of short-term reflexes of the cardiovascular system integrates inputs from peripheral baroreceptors and chemoreceptors and the sympathetic/parasympathetic output to the heart and vascular beds (17). Microinjection of glutamate (2 μg/0.3 μL) into the MePD of awake rats elicits a selective activation of the central sympathetic adjustment to modulate the baroreflex response, whereas microinjection of gamma-aminobutyric acid (GABA; 61 ng/0.3 μL) activates the central parasympathetic counterpart (8). Histamine (30 μg/0.3 μL and 300 μg/0.3 μL) in the MePD shows more complex and dose-dependent responses, with a lower dose modulating part of the baroreflex response and a higher dose leading to a decrease in the parasympathetic component of the HR variability (HRV), an increase in the sympathetic/parasympathetic balance at basal conditions, and impairment of the chemoreflex bradycardic response (7).

It is notable that various neuroactive peptides can be found in the MePD. Oxytocin (OT)-binding sites occur in different MePD cellular columns (18), and local OT modulates the neural processing of social odorants and approach behavior (19). Somatostatin (SST) is found in the MePD as well (20.21) and causes a dose-dependent inhibition of the rat MeA, either directly or by activating GABAergic synaptic transmission (21). There is also an evident expression of the angiotensin II (Ang II) 1A receptor in the MePD (22). These findings are worth noting because OT reduces cardiovascular responses to acute stress in rats (23) and modulates baroreceptor reflex responses (24). SST modulates sympathetic and parasympathetic cardiovascular actions and induces hypotension and bradycardia (25). Ang II can increase AP due to centrally mediated effects (26), acting in circumventricular organs or in the NTS, either by activating the central sympathetic nervous system or by elevating plasma vasopressin, noradrenaline, and adrenaline levels (6,27). In rats, central Ang II also blunts parasympathetic activity, reduces baroreflex sensitivity (28), and stimulates chemoreflex responses (29). However, it is not currently known whether there are cardiovascular effects induced by these neuropeptides when acting in the rat MePD.

Here, we studied the basal hemodynamic state and reflex-mediated cardiovascular responses after microinjections of OT, SST, and Ang II into the MePD of awake male rats. Baroreflex and chemoreflex responses were tested. Power spectral and symbolic analyses were performed to evaluate the central involvement of sympathetic and parasympathetic components in the variability of the HR and AP. We used the same methodological approach previously described for the study of classical neurotransmitter actions in the MePD (7,8, and references therein) and found additional changes in the central control of cardiovascular reflexes elicited by these neuropeptides in this brain area, providing additional detail for better understanding of the mechanisms already described in the literature (6).

### Material and Methods

#### Animals

Adult male Wistar rats (3-4 months old, body weight 250-300 g) were kept in groups under standard laboratory conditions, with food and water *ad libitum* and room temperature around 22°C in a 12:12-h light-dark cycle (lights off at 7:00 pm). All efforts were made to minimize the number of animals and their suffering. Rats were manipulated according to international laws for ethical care and use (European Communities Council Directive of 24 November 1986, 86/609/EEC), conforming to national guidelines, and experiments were approved by the Ethics Committee of Universidade Federal de Ciências da Saúde de Porto Alegre (protocol #05/029).

#### Implantation of the MePD cannula and vascular catheters for hemodynamic evaluation

The experimental design is shown in Figure 1. Animals were anesthetized with a combination of ketamine and xylazine (90 and 10 mg/kg, *ip*, respectively) and were stereotaxically implanted with a cannula (0.6 mm outer diameter, OD) 2 mm above the right MePD, according to the following coordinates: 3.3 mm posterior to the bregma.

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**Figure 1.** Experimental design (in days) with surgical, microinjection, cardiovascular recordings and histological procedure schedules.

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**Table: Experimental Design**

| DAYS | STEREOTAXIC SURGERY | IMPLANTATION OF VASCULAR CATHETERS | CARdioVASCULAR RECORDINGS: - BASAL DATA - MICROINJECTIONS IN THE MePD - BAROREFLEX AND CHEMOREFLEX STIMULATION | HISTOLOGICAL ANALYSIS |
|------|---------------------|-----------------------------------|------------------------------------------------|---------------------|
| 1    |                     |                                   |                                                |                     |
| 3    |                     |                                   |                                                |                     |
| 6    |                     |                                   |                                                |                     |
| 7    |                     |                                   |                                                |                     |
3.6 mm lateral to the sagittal suture, and 5.7 mm below the dura mater (adapted from Ref. 30; Figure 2). After that, each rat received ketoprofen (3 mg/kg in 0.3 mL, ip), one injection per day for 2 days.

After 5 days, animals were anesthetized as cited earlier and had a polyethylene catheter [PE-10 soldered to a PE-50, 0.28 mm inner diameter (ID), Biocorp, Australia, filled with sterile 0.9% NaCl (pH 7.4) and heparin] placed into the abdominal aorta and another into the inferior vena cava through the left femoral artery and vein, respectively. Each catheter was tunneled subcutaneously and exteriorized at the back of the rat neck. Gentamicin (2 mg/rat, im) was

Figure 2. A, Photomicrograph of a coronal section of the ventral forebrain to show the microinjection (MI) site where substances were microinjected, diffused out and reached the posterodorsal medial amygdala (MePD), without lesioning it directly. In this case, the MePD was found approximately 3.30 mm below the bregma, lateral to the optic tract (opt) and ventral to the stria terminalis (st). Hematoxylin and eosin stain. B, Schematic drawing of coronal sections of the ventral part of the rat brain showing the MePD and the approximate sites for the microinjections of saline, oxytocin, somatostatin and angiotensin II (filled circles). Non-target sites are indicated by filled squares. All microinjections were administered to the right MePD and the points represent the approximate locale for drug spread. There are fewer points than the number of animals studied due to overlapping in the microinjection sites in different rats. No evident signs of parenchymal lesions in the MePD were found in the experimental groups. Coordinates are in millimeters posterior to the bregma (from 2.8 to 3.8 mm). MePV, poster-oventral part of the medial amygdaloid nucleus. Adapted from Ref. 30.
injected at the end of this procedure. The next day, the arterial catheter was attached to a 40-cm polyethylene tube (PE-50, 0.5 mm ID, Biocorp), and a strain-gauge pressure transducer (P23 Db, Gould Statham, USA) was used for direct hemodynamic measurements. Signals passed through a preamplifier (Hewlett-Packard 8805, Puerto Rico) and were delivered to a microcomputer equipped with an analog-to-digital converter board (CODAS, 1 kHz, Dataq Instruments, USA). The recorded data were analyzed on a beat-to-beat basis.

All rats of the control and experimental groups were subjected to the same recording protocol. That is, after the catheter was connected to the transducer, rats were allowed to acclimate to a Plexiglas recording box (25 × 15 × 10 cm) over 20 to 30 min while cardiovascular parameters were being continuously monitored. Values of HR and AP were obtained during the last 15 min of this period to provide basal within group and between groups data to be compared with the effects of microinjections into the MePD. All recordings occurred during the morning (from 8:00 am to 12:00 noon) to avoid unpredictable circadian variations in the results.

Immediately after these initial recordings, a microinjection needle (0.3 mm OD) was introduced into and 2 mm below the intracerebral guide cannula. Animals were then randomly assigned to 1 of 4 experimental groups that were manually microinjected with 1) vehicle (saline, 0.3 μL, n = 7) as the control group; 2) OT (10 ng/0.3 μL and 25 pg/0.3 μL, n = 6 in both groups); 3) SST (1 μM/0.3 μL, n = 8; and 0.05 μM/0.3 μL, n = 5); 4) Ang II (50 pmol/0.3 μL and 50 fmol/0.3 μL, n = 7 in both groups). All drugs were purchased from Sigma Chemical Co. (USA). These higher and lower values of OT, SST, and Ang II proved to be effective in inducing central effects when microinjected into different brain areas (described in Ref. 6 and references therein). The microinjection procedure was monitored by the displacement of liquid and an air bubble inside the catheter connected to a 2-μL Hamilton microsyringe (USA), lasted for 1 min, and the needle remained inside the cannula for an additional minute to avoid reflux. After 4 min, new cardiovascular recordings were obtained and lasted for the next 15 min. Additional MePD microinjections were completed before testing baroreceptor- and chemoreceptor-mediated reflexes. Then, every rat received 3 microinjections of each substance assigned to its experimental group as follows: one to test the MePD microinjection effects on the HR and AP values, one before testing the baroreflex responses, and one before testing the chemoreflex responses. The order for testing baroreflex or chemoreflex activities was randomly chosen for each rat in all experimental groups. Obvious drug carryover effects were avoided, and new recordings were only started when hemodynamic values returned to baseline levels and remained stable during the recording time (7,8).

**Baroreflex and chemoreflex stimulation**

Reflex responses were tested 5 min following each MePD microinjection. Baroreflex-mediated changes were measured during peak increases or decreases in mean arterial pressure (MAP) after vena cava injection of a single dose of phenylephrine (8 μg/0.1 mL, Sigma Chemical Co.), or sodium nitroprusside (100 μg/0.1 mL, Sigma Chemical Co.) dissolved in saline (8). These injections were made manually and, in all experimental groups, at similar and appropriate infusion rates, to lead to changes in MAP in the range 10–30 mmHg with no evident artifactual activations. Various data points served to compose sigmoidal curves. All changes in MAP and HR were measured, and baroreflex sensitivity was determined by fitting the MAP and HR changes to a sigmoidal logistic equation, as follows: $HR = P_1 + P_2/[1 + \exp (P_3 \times (MAP–P_4))]$ where $P_1$ is the lower HR plateau [in beats per minute (bpm)], $P_2$ is HR range between upper and lower curve plateaus (delta HR, in bpm), $P_3$ is a curvature coefficient that is independent of range, and $P_4$ is MAP at one-half of the HR range (MAP_{50}, in mmHg), which is also the point for the calculation of the maximum gain. The upper plateau was calculated by the sum of $P_1$ and $P_2$ (7, 8, and references therein).

Chemoreflex sensitivity was tested under the same methodological conditions, but after the administration of increasing intravenous doses of potassium cyanide (KCN; 60, 100, 140, and 180 μg/kg; Merck, Germany). Injected volumes ranged from 0.06 to 0.18 mL. Mean HR and MAP were measured continuously 10 s before and 15 s after each injection of these KCN doses. The inter-injection interval for each dose of KCN was around 4 min, and a new dose was administered only when appropriate HR and MAP baselines had been reestablished (7,8).

**Power spectral analysis and symbolic analysis**

These analyses demonstrate the different components in the variability of the HR and AP recordings and were applied to the results of the experimental groups as long as technically possible. For this reason, some results from the Ang II groups were not sufficient for further analyses and are shown only when they could be reliably done.

These methodological descriptions were reported previously (7,8 and references therein). Briefly, power spectral analysis was applied to pulse interval (PI) and systolic arterial pressure (SAP) series. Tachograms and systograms were created from AP signals through the beat-to-beat PI and SAP, respectively. Frequency domain analysis of HRV (PI variability) and AP variability (APV) were obtained with an autoregressive algorithm on stationary sequences of 200 beats that were randomly chosen, but without artifacts, using the stationary test based on very low quantification of oscillations. Spectral estimation by the autoregressive model, using the residual theorem, provides central frequency and associated power with the additional advantage that, even with short segments of data, they can provide a reliable spectral resolution. HRV and APV correspond to the total power spectrum of PI and SAP.
The low-frequency (LF; 0.2-0.75 Hz) and high-frequency (HF; 0.75-3.0 Hz) spectral components of PI and SAP were expressed in both absolute (ms\(^2\) and mmHg\(^2\), respectively) and normalized units (NU). NU were obtained by calculating the power of LF and HF and correlating them to the total power without the very low frequency component (frequencies that were <0.2 Hz). This method estimates the center frequency and power of each relevant oscillatory component and indicates the involvement of the central control of the sympathetic/parasympathetic systems in the cardiovascular responses. The ratio between LF and HF components (LF/HF) was considered a synthetic expression of the sympathovagal balance. The coherence between the PI and the SAP variability was assessed by means of a cross-spectral analysis. The alpha index was obtained from the square root of the ratio between the R-R interval and the SAP variability within the LF ranges and was calculated only when the magnitude of the squared coherence exceeded 0.5 (range=0 to 1) in the LF band. Using this alpha index for the LF component, beat-to-beat values of the PI and the SAP were used to estimate spontaneous baroreflex sensitivity.

Symbolic analysis was applied on the same sequences of PI and SAP data. This method consisted of the scattering of the time series over 6 bins, each identified by a number (symbol) from 0 to 5. Symbolic series is a sequence of 3 symbols. All 3 possible beat patterns are divided into 4 groups as follows: 1) no variations (0V, i.e., three identical symbols), 2) one variation (1V, two identical symbols and 1 different), 3) two like variations (2LV), and 4) two unlike variations (2UV). The 0V is a marker of sympathetic modulation, a 1V pattern reflects sympathetic and parasympathetic modulations, a 2LV pattern indicates sympathetic and parasympathetic modulations with vagal predominance, and 2UV is an exclusive marker of vagal modulation (see Ref. 31 and references therein).

Histological procedure
At the end of the experiments, rats were deeply anesthetized as described earlier and transcardially perfused with saline and 10% formalin solution. Brains were cut into 60-μm thick sections using a vibratome (Leica, Germany) and stained with hematoxylin and eosin. The microinjection site was histologically determined and compared to the images of an atlas (30). This microinjection procedure was validated for accurate drug administration in small brain regions (32)—further comments about the effects of microinjected substances in the MePD can be found elsewhere (7,8). Only those rats that had the microinjections directed to the MePD were considered for further analysis. Data from rats that had intraparenchimatous hemorrhage, mechanical lesions in the MePD, or microinjections that damaged the stria terminals were not included.

Approximate sites for drug spread in the targeted area are shown in Figure 2A and B. As described previously (8), an inherent technical limitation of the present methodological approach is that drugs may have affected all parts of local neurons and glial cells, including both cell bodies and their prolongments. The specific diffusion rate for each of the substances microinjected into the MePD was not directly evaluated, and the average radius of drug diffusion was not estimated by virtue of comparing a dye with distinct physicochemical properties with the microinjected drugs. However, it is assumed that substances can diffuse an average radius of 0.5 mm from the point of microinjection (32). Here, it is also likely that the position of the guide cannula for the microinjection let the substances spread from the medial border of the MePD through its neuropil, and the coarse axons of the optic tract, situated in the back of the microinjection site, might have limited drug diffusion outside the MePD.

Nevertheless, non-target groups were also included to control for drug effects outside the intended area (n=7 rats for each microinjected substance). Cannulae and microinjections reached the vicinity of the MePD in these non-target groups, and their data served to evaluate possible effects of the tested drugs in other medial amygdala subnuclei, other adjacent amygdaloid nuclei, in the optic tract, or in the ventral basal ganglia, all structures close to the MePD (Figure 2).

Statistical analysis
The basic requirements for the use of parametric analysis were assessed by the Kolmogorov-Smirnov and Levene tests. After that, the values of HR, SAP, diastolic AP (DAP), and MAP at basal conditions and the results of HR and AP evoked by the chemoreceptor reflex response were submitted to two-way analysis of variance (ANOVA) for repeated measures. Data obtained after the stimulation of the baroreflex were analyzed using one-way ANOVA. The Newman-Keuls post hoc test was used for multiple comparisons.

The normal distribution of data from spectral and symbolic analyses was determined by the Bonferroni test. After that, the results were analyzed using one-way ANOVA followed by the Tukey test. When Gaussian normality failed, a Kruskal-Wallis test on ranks and the Dunn’s test were performed. In all cases, the statistically significant level was set a priori as P<0.05.

Results
Present results focus on the MePD results. To be concise, the non-target control data did not reproduce in any way the effects obtained in the experimental groups microinjected with the different neuropeptides in the MePD (ANOVA, P>0.05 in all cases; data not shown).

Hemodynamic status
Basal hemodynamic values were within normal ranges for adult male rats recorded under laboratory conditions.
No relevant statistical difference was found in the HR, SAP, DAP, and MAP following microinjections of saline or OT, SST, and Ang II (ANOVA, P > 0.05 in all cases). There were no lasting effects of the microinjections in any experimental group, which provided stable baseline recordings prior to the assessment of the baroreflex and the chemoreflex responses (Table 1).

### Effects of baroreflex activation

Reflex control of the HR was analyzed by fitting sigmoidal curves to different data points of each animal. Mean values were calculated for each experimental group (Figure 3). The MAP_{50}, which describes the MAP value at one-half of the HR range in the baroreflex curve, following the injections of phenylephrine and sodium nitroprusside showed an overall statistical difference between groups [F(6, 47) = 3.37, P < 0.01]. The post hoc comparisons revealed that the MePD microinjection of OT (10 ng) or SST (1 μM) promoted a significant decrease in the MAP_{50}, shifting the barocurve to the left and to lower HR values when compared with the results from the saline microinjected group (P < 0.05 in both cases).

An overall statistical difference was also found in the maximum gain of the reflex responses when the AP was increased by phenylephrine or decreased by sodium nitroprusside when compared between the experimental groups [F(6, 47) = 2.56, P = 0.03], but post hoc comparisons did not show differences between any tested substances compared with saline. For the HR range [F(6, 47) = 1.29, P = 0.27], the lower plateau [F(6, 47) = 1.53, P = 0.19] and the upper plateau [F(6, 47) = 1.19, P = 0.09] did not differ between the experimental groups.

### Effects of chemoreflex activation

Data for chemoreflex activation are shown in Figure 4. There was a statistically significant difference in the HR after injection of the increasing doses of KCN [F(3, 123) = 69.77, P = 0.01] and between experimental groups microinjected into the MePD [F(6, 123) = 2.82, P = 0.02], but no statistical interaction was found between these two factors [F(18, 123) = 1.51, P = 0.09].

On the other hand, the values of MAP obtained after the injection of the different doses of KCN reached statistically significant levels [F(3, 123) = 17.26, P = 0.01] and, although data did not differ between the experimental groups [F(6, 123) = 1.08, P = 0.38], the interaction between these two factors was significantly different [F(18, 123) = 2.22, P < 0.01]. The post hoc test showed that, compared with saline, OT (10 ng) microinjected into the MePD led to a marked increase in the MAP following KCN (180 μg/kg; P < 0.05). No other comparison showed a significant

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**Table 1.** Basal hemodynamic data obtained before (control data) and after (test data) microinjections into the rat posterodorsal medial amygdala of saline (0.3 μL, n = 7), oxytocin (OT; 10 ng/0.3 μL and 25 pg/0.3 μL, n = 6 in both groups), somatostatin (SST; 1 μM/0.3 μL, n = 8 and 0.05 μM/0.3 μL, n = 5), or angiotensin II (Ang II; 50 pmol/0.3 μL and 50 fmol/0.3 μL; n = 7 in both groups).

| Groups       | Control (mmHg) | After (mmHg) | Control (mmHg) | After (mmHg) | Control (mmHg) | After (mmHg) | Control (mmHg) | After (mmHg) | Control (mmHg) | After (mmHg) |
|--------------|----------------|-------------|----------------|-------------|----------------|-------------|----------------|-------------|----------------|-------------|
| Saline       | 136 ± 9.0      | 131.0 ± 12.7| 90.1 ± 4.6     | 89.5 ± 9.3  | 111.6 ± 5.6    | 109.0 ± 10.1| 360.2 ± 29.3   | 350.7 ± 31.9| 132.6 ± 11.2   | 135.7 ± 7.8  |
| OT (10 ng)   | 131.7 ± 5.2    | 128.7 ± 6.6 | 93.7 ± 7.9     | 91.9 ± 9.4  | 111.9 ± 6.5    | 109.4 ± 8.8 | 370.7 ± 23.4   | 347.7 ± 12.5| 131.9 ± 4.0    | 135.4 ± 7.0  |
| SST (1 μM)   | 131.5 ± 7.7    | 124.0 ± 6.8 | 88.5 ± 3.8     | 91.6 ± 6.3  | 109.7 ± 3.3    | 112.7 ± 6.9 | 352.5 ± 39.6   | 365.9 ± 38.1| 131.5 ± 7.7    | 124.0 ± 6.8  |
| Ang II (50 pmol) | 134.0 ± 6.0  | 131.1 ± 8.5 | 94.8 ± 6.1     | 95.4 ± 10.4 | 113.5 ± 5.5    | 112.5 ± 9.5 | 354.7 ± 37.3   | 353.6 ± 53.3| 134.4 ± 11.8   | 134.7 ± 9.2  |

Data are reported as means ± SD. SAP: systolic arterial pressure; DAP: diastolic arterial pressure; MAP: mean arterial blood pressure; HR: heart rate. No statistically significant differences were found comparing data between experimental groups (one-way ANOVA).
post hoc ANOVA test for repeated measures and the Newman-Keuls test indicated that SST (1 μM) promoted an increase in this parameter compared with saline (P<0.05 in all cases). In addition, the LF component of the HRV, in NU, presented a statistically significant difference between the experimental groups [F(5,33)=2.61; P<0.05] and the post hoc test indicated that SST (1 μM and 0.05 μM) promoted an increase in this parameter compared with saline (P<0.05 in both cases). There was also an overall difference in the absolute power of the HF component of the HRV [F(5,33)=2.65; P<0.05], and the post hoc comparison showed that OT (10 ng) promoted an increase in this parameter compared with saline (P<0.05).

The sympathetic/parasympathetic balance obtained from the alpha index also showed a statistically significant difference between experimental groups [F(5,33)=2.71; P<0.05] and, specifically, SST (0.05 μM) promoted higher values compared with saline (Newman-Keuls test, P<0.05).

Symbolic analysis

Results are presented in Table 3 and Figure 5. Compared with control data, the microinjections of OT (10 ng and 25 pg) and SST (1 and 0.05 μM) did not cause evident effects (Newman-Keuls test, P>0.05), but there was a significant decrease in the 2UV pattern after Ang II (50 fmol) microinjection into the MePD when compared with saline [F(5,34)=2.282, P<0.05; Newman-Keuls test, P<0.05].

Discussion

Our present results reinforce findings that the rat MePD is a component of the central network that modulates baroreceptor- and chemoreceptor-mediated cardiovascular adjustments. These data match with the evident presence of neuropeptides and their receptors in the MePD (18-22) and indicate functional implications that deserve further evaluation, as follows.

First, it is interesting that the tested neuropeptides promoted long-lasting effects (in the order of minutes) when microinjected into the MePD, as previously reported for classical neurotransmitters [glutamate, GABA, and histamine], which appears to be a particularity of this brain area. In addition, the MePD contributes more to changes in cardiovascular reflexes than in basal HR and AP levels. As occurs for other amygdaloid nuclei (33), it is possible that transmitter-mediated excitatory, inhibitory, and/or disinhibitory mechanisms in the MePD can promote a flexible, dose- and context-dependent modulation of the central control of sympathetic/parasympathetic output (6). We also propose that the rat MePD is responsive to different

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**Figure 4. A.** Heart rate (HR) and B, mean arterial blood pressure (MAP) of the chemoreceptor reflex response induced by increasing doses of potassium cyanide (KCN, from 60 to 180 μg/kg) in rats that received microinjections into the posterodorsal medial amygdala of saline (0.3 μL, n=7), oxytocin (OT; 10 ng/0.3 μL and 25 pg/0.3 μL, n=6 in both groups), somatostatin (SST; 1 μM/0.3 μL, n=8 and 0.05 μM/0.3 μL, n=5), or angiotensin II (Ang II; 50 pmol/0.3 μL and 50 fmol/0.3 μL; n=7 in both groups). Data are reported as means±SD. Data were submitted to the two-way ANOVA test for repeated measures and the Newman-Keuls post hoc test. In A, for HR: *P<0.01 compared with 60 μg/kg KCN in the group microinjected with saline. **P<0.01 compared with 60 μg/kg KCN in the group microinjected with OT 10 ng. *P<0.01 compared with 60 μg/kg KCN in the group microinjected with OT 25 pg. 1P<0.01 compared with 60 μg/kg KCN in the group microinjected with OT 10 ng. *P<0.05 compared with 140 μg/kg KCN in the group microinjected with Ang II 50 pmol. In B, for MAP: *P<0.01 compared with 60 μg/kg KCN in the group microinjected with OT 10 ng. *P<0.05 compared with 180 μg/kg KCN in the group microinjected with Ang II 50 pmol. +P<0.05 compared with 180 μg/kg KCN in the group microinjected with saline. 1P<0.05 compared with 180 μg/kg KCN in the group microinjected with OT 10 ng. Expanded and adapted from Ref. 6 with permission from Nova Science Publishers (USA).

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statistical difference between the experimental groups and the saline group (P>0.05).

**Power spectral analysis**

Power spectral analysis results are shown in Table 2 and Figure 5. There were no statistically significant differences in the SAP [F(5,33)=1.63; P=0.17], in the mean values of HR [F(5,33)=2.01; P=0.10], in the absolute LF component [F(5,33)=1.87; P=0.125], or in the absolute HF component of the SAP between the experimental groups [F(5,33)=1.41; P=0.24].
neurotransmitters and neuropeptides that code distinct synaptic inputs for specific fine-tunings of centrally mediated cardiovascular reflexes and the dynamic display of behaviors in awake animals (7,8). It is now desirable to determine how neurotransmitters and neuropeptides in the MePD can modulate the activity of specific output pathways that reach other brain areas related to central cardiovascular control.

Second, all the neuropeptides tested here evoked a higher HRV as revealed by the power spectral analysis. The predominant central sympathetic or parasympathetic modulation of the cardiovascular system involves the balance between the spectral components in favor of the LF or the HF band, respectively (34). However, it is noteworthy that OT microinjections into the MePD did not cause specific effects on the parameters evaluated by the symbolic analysis. Then, it is possible that higher absolute LF and HF power values (after OT 25 pg and 10 ng, respectively) can be explained by the increase in HRV itself and the change of the HF peak value, which is also associated with ventilatory components and might be correlated with an increased respiratory rate following OT (10 ng) microinjection into the MePD. The participation of rat MePD in the central control of breathing is currently unknown. This is a working hypothesis that needs to be confirmed with other methodological approaches and opens a new and interesting line of research.

Indeed, OT can have site-specific roles on the central modulation of cardiovascular responses instead of similar and widespread actions. For example, although intracebroventricular injection of OT did not evoke significant cardiovascular effects (35), unilateral microinjections of an OT receptor antagonist in the NTS reduced the long-lasting vasopressor and tachycardic responses elicited by local microinjection of this neuropeptide (24). OT also counteracted the vasodepressor and bradycardic responses induced by glutamate in this nucleus (24). Data from the CeA, a component of the extended amygdala, provided another interpretation for the effects of OT (36). That is, OT microinjected into the CeA promoted an elevation in plasma corticosterone, but not epinephrine and norepinephrine. At the same time, the induction of a tachycardic response was attributable to a somatic-sympathetic/parasympathetic coupling rather than to a genuine “autonomic” activation (36). Here, the microinjection of OT into the MePD also appeared to affect central cardiovascular control only indirectly.

Otherwise, microinjection of SST into the MePD caused an increase in the HRV suggestive of a central sympathetic and parasympathetic coactivation. Because this finding was also accompanied by an increase in the LF (in NU), they rather point to changes in the central sympathetic regulation. This increase in the HRV was also accompanied by a higher alpha index, which indicates the relationship between the LF components of the HRV and the APV along with an increase in baroreflex sensitivity. The precise mechanism by which SST exerts its effects in the MePD still remains to be established. SST inhibits the presynaptic release of glutamate, activates potassium channels, and inhibits voltage-gated calcium channels via sst2 or sst5 receptors (37 and references therein). SST can also promote an inhibitory action by acting directly on its receptors or indirectly via GABAergic synapses in the MeA (21). From our present results, SST induced a central sympathetic effect, whereas, comparatively, GABA in the MePD decreased the upper

Table 2. Cardiovascular parameters of power spectral analysis calculated from sampled data of rats that received microinjections into the posterodorsal medial amygdala of saline (0.3 μL), oxytocin (OT; 10 ng/0.3 μL and 25 pg/0.3 μL), somatostatin (SST; 1 μM/0.3 μL and 0.05 μM/0.3 μL), or angiotensin II (Ang II; 50 pmol/0.3 μL).

| Measurement | Saline (n=7) | OT 10 ng (n=6) | OT 25 pg (n=6) | SST 1 μM (n=8) | SST 0.05 μM (n=5) | Ang II 50 pmol (n=7) |
|-------------|-------------|---------------|---------------|---------------|------------------|-------------------|
| HRV (ms²)   | 13.35 ± 4.18 | 29.34 ± 9.57  | 46.75 ± 18.7* | 49.80 ± 16.90* | 58.81 ± 39.59*  | 50.78 ± 22.23*    |
| fLF (Hz)    | 0.54 ± 0.10  | 0.50 ± 0.10   | 0.51 ± 0.12   | 0.45 ± 0.10   | 0.42 ± 0.07      | 0.56 ± 0.10       |
| HF (ms²)    | 0.49 ± 0.26  | 1.51 ± 0.89   | 6.48 ± 6.21*  | 3.04 ± 2.62   | 5.70 ± 4.88      | 3.80 ± 3.59       |
| LF (ms²)    | 9.24 ± 3.55  | 13.45 ± 6.41  | 15.53 ± 6.63  | 19.66 ± 8.17* | 22.38 ± 11.86*   | 14.46 ± 7.66      |
| HF (Hz)     | 1.84 ± 1.37  | 1.87 ± 10.95* | 1.76 ± 151    | 1.70 ± 1.68   | 1.67 ± 1.21      | 1.65 ± 1.29       |
| LF/HF       | 0.53 ± 0.11  | 0.53 ± 0.14   | 1.16 ± 0.22   | 1.09 ± 0.79   | 1.56 ± 0.65*     | 1.29 ± 0.93       |
| APV (mmHg²) | 8.84 ± 3.48  | 20.16 ± 10.70 | 10.54 ± 5.89  | 15.80 ± 8.82  | 12.32 ± 5.98     | 17.46 ± 12.55     |
| VLF (mmHg²) | 5.78 ± 4.09  | 11.07 ± 8.76  | 3.62 ± 2.84   | 8.90 ± 9.22   | 8.71 ± 7.13      | 10.13 ± 12.49     |
| HF (mmHg²)  | 1.72 ± 0.77  | 5.78 ± 3.31   | 4.04 ± 3.18   | 4.20 ± 3.16   | 2.73 ± 1.79      | 3.60 ± 1.93       |
| HF (ms)     | 1.33 ± 0.43  | 3.24 ± 2.38   | 2.86 ± 3.26   | 2.70 ± 1.02   | 0.88 ± 0.36      | 2.29 ± 2.06       |
| Alpha index (ms/mmHg) | 0.53 ± 0.11 | 0.53 ± 0.14 | 1.16 ± 0.52 | 1.09 ± 0.79 | 1.56 ± 0.65* | 1.29 ± 0.93 |

Data are reported as means ± SD. HRV: heart rate variability; fLF: central low frequency; fHF: central high frequency; LF: low frequency power; HF: high frequency power; APV: arterial pressure variability; VLF: very low frequency power. *P<0.05, compared to saline (one-way ANOVA and the Tukey post hoc test or the Kruskal-Wallis test and the Dunn’s post hoc test).
plateau of the baroreflex curve and induced higher values of HRV linked with the central parasympathetic control of the cardiovascular system (8). Therefore, these findings demonstrate non-overlapping effects of SST and GABA on cardiovascular responses when acting in the rat MePD.

Finally, microinjection of Ang II into the MePD increased the HRV, did not affect power spectral components, but reduced the 2UV pattern of the symbolic analysis, which

Figure 5. Power spectrum and symbolic analysis applied to a time series of cardiovascular parameters of rats. The columns show the pulse interval series, power spectrum and the symbolic pattern distribution for the data obtained from the experimental groups that received microinjections into the posterodorsal medial amygdala with A, saline (0.3 μL, n = 7), B, oxytocin 10 ng and C, 25 pg (n = 6 in both groups), D, somatostatin 1 μM (n = 8) and E, 0.05 μM (n = 5), or F, angiotensin II 50 pmol (for the power spectral analysis) and 50 fmol (for the symbolic analysis; n = 7 in both groups). Values are from representative samples of each experimental group. Mean values for all studied animals are shown in Tables 2 and 3.
was linked to cardiac vagal activity in pharmacological tests (38). In the group microinjected with 50 pmol ANG II, it is also possible that the lower bradycardic response produced by pressure increase was related to the blunting of the parasympathetic component, and the higher baroreflex gain observed could be associated with the higher tachycardic response after AP reduction. Microinjection, electrophysiological, and lesion studies demonstrate complex interactions and multiple sites for the action of Ang II on cardiovascular control and in different brain areas (26-29). It is interesting that AT1A receptors are involved in neuronal activation and cardiovascular change after an olfactory-mediated psychosocial stress in mice (39). Acting in the MeA, Ang II might inhibit stress-induced aversive responses (39). However, no strain difference (between spontaneously hypertensive and normotensive Wistar male rats) was identified in c-Fos immunoreactivity in the MeA after aversive air jet or air noise stress stimulation (40). Then, it is possible that different species and/or types and intensity of sensory stimulation can trigger the Ang II-evoked responses. Marked central Ang II actions on the cardiovascular system, partly stimulating sympathetic responses (27), were not found after its microinjection into the MePD. Rather, we found a central parasympathetic effect and, in agreement with NTS data (29), Ang II in the MePD potentiated the chemoreflex-mediated decrease in HR frequency. These new findings add and expand the idea of site-specific effects of Ang II on central cardiovascular control and reinforce the multiple modulatory actions of the MePD.

In summary, our work provides additional data for MePD modulation of the baroreflex and chemoreflex cardiovascular responses in awake rats. Local microinjections promoted neuropeptidergic-selective responses and involved the central activation of the sympathetic and parasympathetic effector systems. This MePD plastic capacity could provide the animal with a dynamic mechanism of cardiovascular control suitable for the ongoing demands of social behaviors, for which the MePD is a relevant component in integrated brain circuits (2-6,9-11). Besides the adjustment of homeostatic responses, these neuropeptidergic actions in the MePD can also be involved in the pathophysiology of cardiovascular diseases, such as hypertension (14-16), a possibility that deserves further studies as well.

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