

Research

Open Access

Prolactin daily rhythm in suckling male rabbits

Pilar Alvarez¹, Daniel Cardinali², Pilar Cano³, Pilar Rebollar⁴ and Ana Esquifino^{*3}

Address: ¹Departamento de Biología Celular, Facultad de Medicina, Universidad Complutense de Madrid, 28040 Madrid, Spain, ²Departamento de Fisiología, Facultad de Medicina, Universidad de Buenos Aires, 1121 Buenos Aires, Argentina, ³Departamento de Bioquímica y Biología Molecular III, Facultad de Medicina, Universidad Complutense de Madrid, 28040 Madrid, Spain and ⁴Departamento de Producción Animal, E.T.S.I. Agrónomos, Universidad Politécnica de Madrid, Spain

Email: Pilar Alvarez - pilar@med.ucm.es; Daniel Cardinali - daniel@mail.retina.ar; Pilar Cano - pelayos@med.ucm.es; Pilar Rebollar - prebollar@pan.etsia.upm.es; Ana Esquifino* - pelayos@med.ucm.es

* Corresponding author

Published: 13 January 2005

Received: 18 November 2004

Journal of Circadian Rhythms 2005, **3**:1 doi:10.1186/1740-3391-3-1

Accepted: 13 January 2005

This article is available from: <http://www.jcircadianrhythms.com/content/3/1/1>

© 2005 Alvarez et al; licensee BioMed Central Ltd.

This is an Open Access article distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/2.0>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Abstract

Background: This study describes the 24-h changes in plasma prolactin levels, and dopamine (DA), serotonin (5HT), gamma-aminobutyric acid (GABA) and taurine concentration in median eminence and adenohipophysis of newborn male rabbits.

Methods: Animals were kept under controlled light-dark cycles (LD 16:8, lights on at 08:00 h), housed in individual metal cages, and fed ad libitum with free access to tap water. On day 1 after parturition, litter size was standardized to 8–9 to assure similar lactation conditions during the experiment. Groups of 6–7 suckling male rabbits were killed by decapitation on day 11 of life at six different time points during a 24-h period.

Results: Plasma prolactin levels changed significantly throughout the day, showing a peak at the beginning of the active phase (at 01:00 h) and a second maximum during the first part of the resting phase (at 13:00 h). Median eminence DA concentration also changed significantly during the day, peaking at the same time intervals as plasma prolactin. A single maximum (at 13:00 h) was found for adenohipophysial DA concentration. Individual adenohipophysial DA concentrations correlated significantly with their respective plasma prolactin levels. A maximum in median eminence 5HT concentration occurred at 21:00 h whereas adenohipophysial 5HT peaked at 13:00 h. Median eminence 5HT concentration and circulating prolactin correlated inversely. In the median eminence, GABA concentration attained maximal values at 21:00 h, whereas it reached a maximum at 13:00 h in the pituitary gland. Median eminence GABA concentration correlated inversely with circulating prolactin. In the median eminence, taurine values varied in a bimodal way showing two maxima, at the second half of the rest span and of the activity phase, respectively. In the adenohipophysis, minimal taurine levels coincided with the major plasma prolactin peak (at 01:00 h). Circulating prolactin and adenohipophysial taurine levels correlated inversely.

Conclusion: The correlations among the changes in the neurotransmitters analyzed and circulating prolactin levels explain the circadian secretory pattern of the hormone in newborn male rabbits.

Background

The mechanisms that regulate prolactin secretion are complex [1]. Two major regulatory inhibitory inputs for prolactin secretion are dopamine [2] and gamma-aminobutyric acid (GABA) [3-6]. In addition, many other neuromodulators have been implicated in the control of prolactin secretion, among them, vasoactive intestinal peptide, thyrotropin releasing hormone and serotonin (5HT) [1]. More recently, taurine has also been implicated in the regulation of prolactin secretion [1].

It is well known that basal secretion of prolactin varies throughout the day, describing a characteristic pattern with maximal values close to the light-dark transition [7,8]. Such a circadian pattern has been described not only in rodents (rat and mouse) but also in many other species [1]. In the rat, we previously demonstrated changes of the secretory pattern of prolactin along the year [7-11], as well as a function of aging [12,13].

The rat is very immature at birth, so that newborn and suckling rats are very sensitive to manipulations that can affect adulthood [14-18]. Circadian rhythms of developing mammals seem to be entrained by the rhythmicity of their mother [19,20], and several studies have indicated that maternal melatonin is necessary to entrain the circadian rhythms in the newborn [21,22].

The rabbit is probably the best-studied laboratory animal in the wild, due to its abundance, size and importance as an agricultural pest [23,24]. Wild and laboratory rabbits are essentially nocturnal and display a clear daily pattern of activity [25]. The rabbit possesses a number of behavioral specializations that make it uniquely suited for circadian studies. Female rabbits visit their altricial young only for a few minutes once every 24 h to nurse, and survival of the young depends on the tight circadian-controlled synchronization in behavior and physiology with the mother. This unusual pattern of maternal care and the demands it places on the litter provide an excellent opportunity to analyze circadian rhythms during early development [25].

In contrast to the large amount of information available on circadian rhythms in adult mammals, studies on circadian phenomena in neonates are few [26,27]. For example, in 21 day-old male rats the daily circadian pattern of prolactin secretion seen in adults is absent [18]. Considering that no information on circadian rhythmicity of prolactin secretion in neonatal male rabbits is available, we undertook the present study to analyze whether neonatal male rabbits show defined 24-h changes in plasma prolactin levels and whether neonatal male rabbits show circadian changes in DA, 5HT, GABA and taurine concentration in median eminence and the adenohipo-

physis, all of which are well known modulators of prolactin secretion.

Methods

Animals

This study was performed using 24 multiparous, lactating Californian × New Zealand White crossbreed doe rabbits. Animals were housed in research facilities of the Animal Production Department. They were maintained under controlled light-dark cycles (LD 16:8, light on at 08:00 h), housed in individual metal cages, fed *ad libitum* using a commercial pellet diet (Lab Rabbit Chow, Purina Mills, Torrejón de Ardoz, Madrid, Spain) with free access to tap water. On day 1 after parturition, litter size was standardized to 8-9 by adding or removing kits to assure similar lactation conditions during the experiment. This study was performed according to the CEE Council Directive (86/609, 1986) for the care of experimental animals. Groups of 6-7 suckling male rabbits were killed by decapitation on day 11 of life at six different time points throughout a 24-hour cycle. The brains were quickly removed, and the median eminence and the anterior pituitary were taken out. Anterior pituitaries were weighed and homogenized in chilled (0-1 °C) 2 M acetic acid. After centrifugation (at 15000 × g for 30 min, at 5 °C), the samples were either analyzed for DA and 5HT or boiled for 10 min and further centrifuged at 14000 rpm for 20 min to measure GABA and taurine.

Hormone assay

Plasma prolactin levels were measured by a specific homologous RIA method [28] using AFP-991086 antibody supplied by the National Institutes of Health (NIH, Bethesda, MD, USA) and Dr. A. F. Parlow (Harbour-UCLA Medical Center, CA, USA). The titer of antibody used was 1:62,500. The PRL standard used was RbPRL_r-RP-1. Hormone was labeled with ¹²⁵I by the chloroamine-T method [29]. The volume of plasma for PRL determinations was 10 µl. *Staphylococcus aureus* (prepared by the Department of Plant Physiology, U.A.M., Madrid, Spain) was used to precipitate the bound fraction [28]. All samples were measured in the same assay run to avoid inter-assay variations. The sensitivity of the assay for PRL was 0.125 ng/ml and the intra-assay coefficient of variation was < 5%. The intra-assay coefficient of variation was calculated using a pool of plasma measured ten times in the same assay; mean (± S.E.M.) concentration was 106.9 ± 4.1 ng/ml.

Catecholamine and indoleamine analysis

DA and 5HT concentration was measured by high pressure liquid chromatography (HPLC) using electrochemical detection (Coulochem, 5100A, ESA; USA), as described elsewhere [12]. A C-18 reverse phase column eluted with a mobile phase (pH 4. 0.1 M sodium acetate,

0.1 M citric acid, 0.7 mM sodium octylsulphate and 0.57 mM EDTA containing 10% methanol, v/v) was employed. Flow rate was 1 ml/min, at a pressure of 2200 psi. Fixed potentials against H_2/H^+ reference electrode were: conditioning electrode: -0.4 V; preoxidation electrode: +0.10; working electrode: +0.35 V. Indoleamine and catecholamine concentration was calculated from the chromatographic peak heights by using external standards and was expressed as pg/ μ g protein. The linearity of the detector response for DA and 5HT was tested within the concentration ranges found in median eminence and adenohipophysial supernatants.

Amino acid analysis

Amino acids were isolated and analyzed by HPLC with fluorescence detection after precolumn derivatization with O-phthalaldehyde (OPA) as described elsewhere [30]. An aliquot of the tissue supernatant containing homoserine as an internal standard was neutralized with 4 M NaOH and was then incubated at room temperature with OPA reagent (4 mM OPA, 10% methanol, 2.56 mM 2-mercaptoethanol, in 1.6 M potassium borate buffer, pH 9.5) for 1 min. The reaction was stopped by adding acetic acid (0.5 % v/v). Samples were immediately loaded through a Rheodyne (Model 7125) injector system (50 μ l loop) to reach a C-18 reverse-phase column (4.6 mm ID \times 150 mm, Nucleosil 5, 100A). Elution was achieved by means of a mobile phase consisting of 0.1 M sodium acetate buffer (pH 6.5) containing 35 % methanol, at a flow rate of 1 mL/min and a pressure of 140 Bars. The column was subsequently washed with the same buffer containing 70 % methanol and re-equilibrated with the elution buffer before re-use. The filter fluorometer was set at the following wavelengths: excitation: 340 nm, emission: 455 nm. The procedure allowed a distinct separation and resolution of the amino acids measured. Amino acid content was calculated from the chromatographic peak heights by using standard curves and the internal standard. The linearity of the detector response was tested within the concentration ranges found in median eminence and adenohipophysial extracts.

Statistics

Statistical analysis of results was performed by a one-way analysis of variance (ANOVA) followed by post-hoc Tukey-Kramer's multiple comparisons tests. Curve estimation in regression analysis was made by using SPSS software, version 10.1 (SPSS Inc., Chicago, ILL). P values lower than 0.05 were considered evidence for statistical significance.

Results

Figure 1 shows the levels of prolactin throughout the day in suckling male pups. Plasma prolactin levels changed significantly throughout the day ($F = 21.1$; $p < 0.0001$),

showing two maxima, a major one at the beginning of the active phase (at 01:00 h) and a second one during the first part of the resting phase (at 13:00 h).

Figures 2,3,4,5 depict the changes in median eminence and adenohipophysial concentration of DA, 5-HT, GABA and taurine. Mean plasma prolactin concentration is plotted as a reference in every case.

Median eminence DA concentration changed in a bimodal way as a function of time of day, showing two maxima, coinciding with those of plasma prolactin at the active and resting phase of the diurnal cycle ($F = 14.1$; $p < 0.0001$, Figure 2). In the case of adenohipophysial DA concentration, a single maximum occurred during the first half of the rest phase (at 13:00 h) ($F = 29.9$; $p < 0.0001$). Only in the adenohipophysial, plasma prolactin and DA concentration correlated in a direct way. This correlation was best described by a log model with $r^2 = 0.16$, $b_0 = -123.7$ and $b_1 = 18.1$ ($F = 4.69$, $p = 0.04$).

As shown in Figure 3, a maximum in median eminence 5HT concentration occurred at the second half of the rest span ($F = 64.1$; $p < 0.0001$) whereas a maximum in adenohipophysial 5HT levels was found at the first half of rest span. Circulating prolactin and median eminence 5HT concentration correlated inversely in a linear way ($r^2 = 0.18$, $b_0 = 677.6$ and $b_1 = -4.9$, $F = 5.3$, $p < 0.03$).

Figure 4 shows the changes in median eminence and adenohipophysial GABA concentration. In the median eminence, GABA concentration attained maximal values at the rest phase, with a peak at late evening (i.e. at 21:00 h, $F = 11.1$, $p < 0.0001$). In the anterior pituitary, GABA concentration reached a maximum at 13:00 h ($F = 21.6$, $p < 0.0001$). Circulating prolactin and median eminence GABA concentration correlated inversely in a linear way ($r^2 = 0.21$, $b_0 = 25.7$ and $b_1 = -0.22$, $F = 6.6$, $p < 0.01$).

Figure 5 depicts the 24-h changes in taurine concentration. In the median eminence, taurine values varied in a bimodal way showing a peak at the second half of the rest period, a nadir at the early activity span (coinciding with the prolactin peak) and a second maximum late in the activity phase (at 05:00 h, $F = 32.9$, $p < 0.0001$). Likewise, in the adenohipophysial, taurine levels exhibited minimal values at the time of the prolactin peak (i.e., at 13:00 h, $F = 21.6$, $p < 0.0001$). Circulating prolactin and adenohipophysial taurine levels correlated inversely in a linear way ($r^2 = 0.42$, $b_0 = 11.6$ and $b_1 = -0.11$, $F = 17.4$, $p < 0.0001$).

Discussion

The present study, performed in neonatal male rabbit pups sacrificed at 6 different time intervals during a 24-h cycle, describes for the first time significant changes in

PROLACTIN

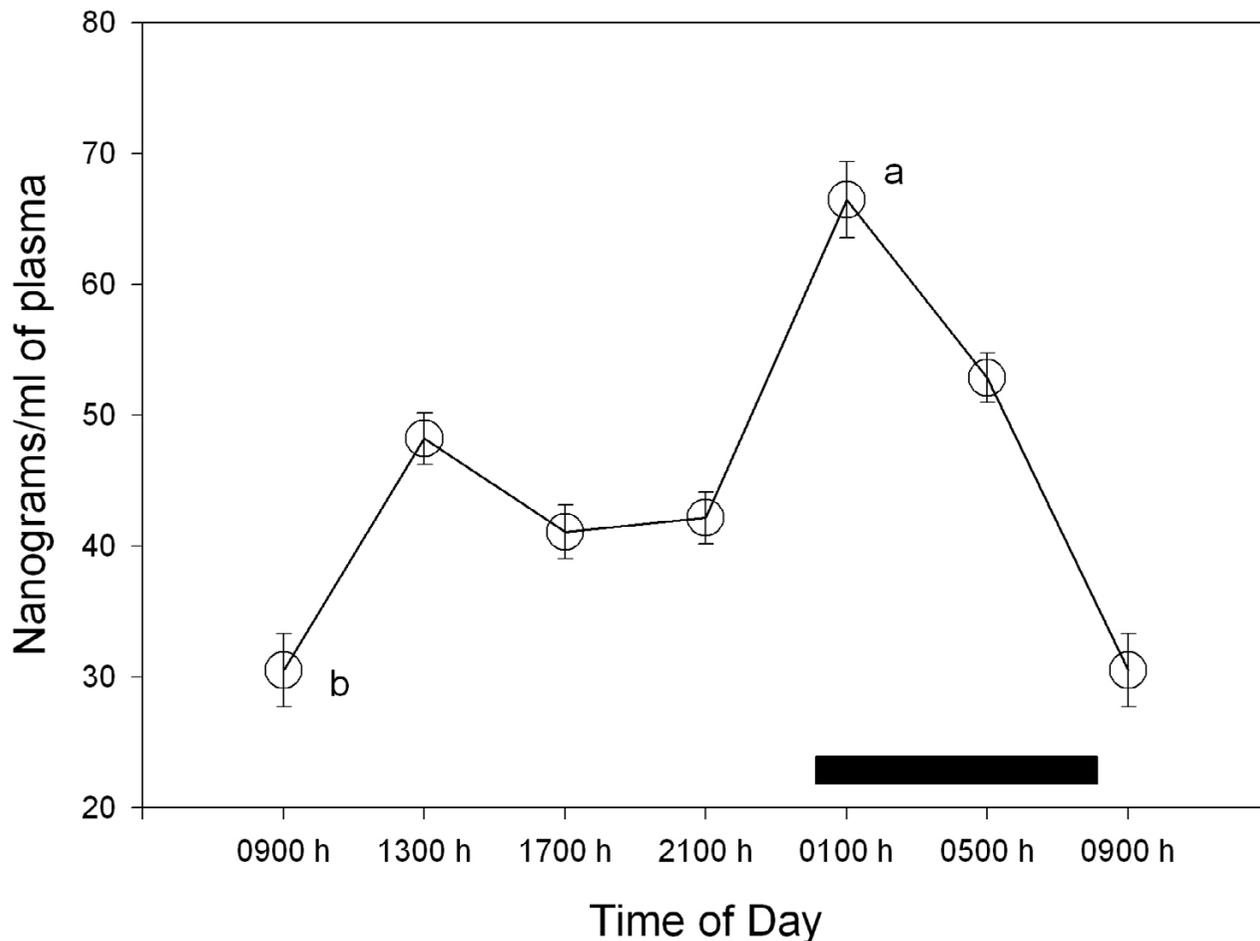


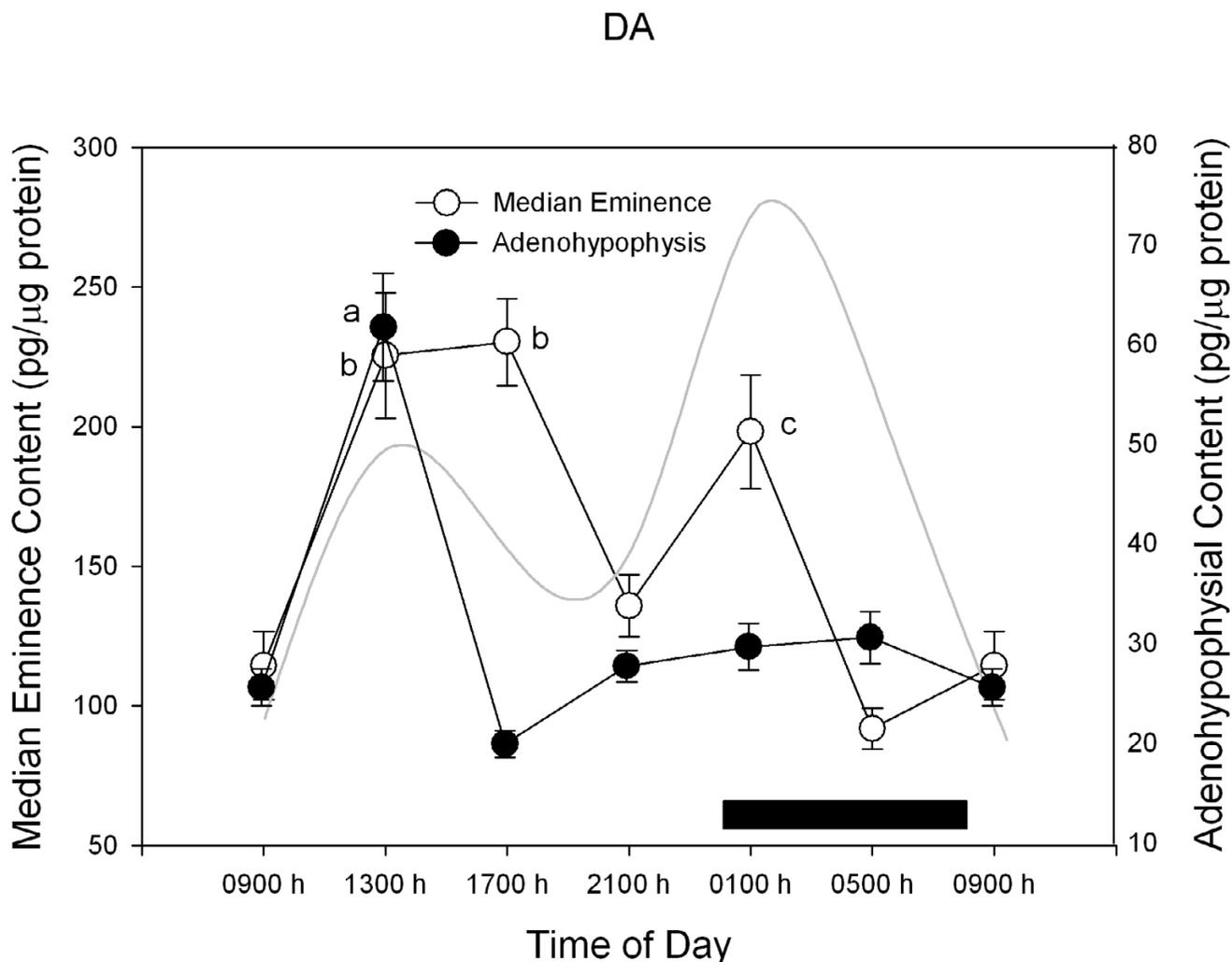
Figure 1

24-h changes in plasma prolactin levels of 11 days old male rabbit pups. Groups of 6–7 pups were killed by decapitation at 6 different time intervals throughout a 24 h cycle. Bar indicates scotophase duration. Results are the means \pm SEM. ^a $p < 0.01$ vs. all time points. ^b $p < 0.01$ vs. 01:00 h, 05:00 h and 13:00 h, Tukey-Kramer's multiple comparisons test. For further statistical analysis, see text.

plasma prolactin levels throughout the day. In concomitant measurements of median eminence and adeno-hypophysial concentration of DA, 5HT, GABA and taurine, a clear daily pattern was found in almost every case. Contrasting with neonatal rats that did not display any circadian pattern of plasma prolactin [18], a daily rhythm of plasma prolactin occurred in neonatal male rabbits, with a maximal value attained 1 h after lights-off (at 01:00 h) and a secondary peak found during the first part of the resting phase (at 13:00 h).

In adult rabbits, daily patterns of prolactin secretion depend on light/dark phases [25]. The present results indicate that, already on day 11 of life, male rabbit pups display daily changes in plasma prolactin levels, remarkably similar to those described in adult male rats (e.g., the maximum displayed 1 h after the dark onset) [7-10].

The activity of several nuclei of rabbit hypothalamus increases with age and with experience of anticipatory arousal [27]. However, no study has been published on

**Figure 2****24-h changes in median eminence and adenohipophysial DA concentration in 11 days old male rabbit pups.**

Groups of 6–7 pups were killed by decapitation at 6 different time intervals throughout a 24 h cycle. Bar indicates scotophase duration. Results are the means \pm SEM. Circulating prolactin levels are shown in shaded line. Letters indicate the existence of significant differences between time points within each tissue after a Tukey-Kramer's multiple comparisons test, as follows: ^a $p < 0.01$ vs. all time points. ^b $p < 0.01$ vs. 05:00 h, 09:00 h and 21:00 h. ^c $p < 0.01$ vs. 05:00 and 21:00 h. For further statistical analysis, see text.

the regulatory mechanism of prolactin in rabbits. Considering that DA is the major inhibitory input for prolactin secretion [1,32], the present study indicating that DA concentration in median eminence of rabbit pups is high during the rest phase of the day (when plasma prolactin levels are low), and decreases at day-night transition (coinciding with the increase in circulating prolactin), may support a cause-effect relationship. The afternoon decrease in median eminence DA concentration could be

a prerequisite for prolactin release in neonatal male rabbits [2]. However, median eminence DA concentration of male rabbit pups also presents a peak during the activity phase (01:00 h) associated with the highest prolactin levels. Therefore, the data suggest that the inhibitory regulatory influence of DA on prolactin secretion is exerted mainly during the light phase of the photoperiod, whereas during the dark phase other hypothalamic neuro-modulators could be operative, as it was previously

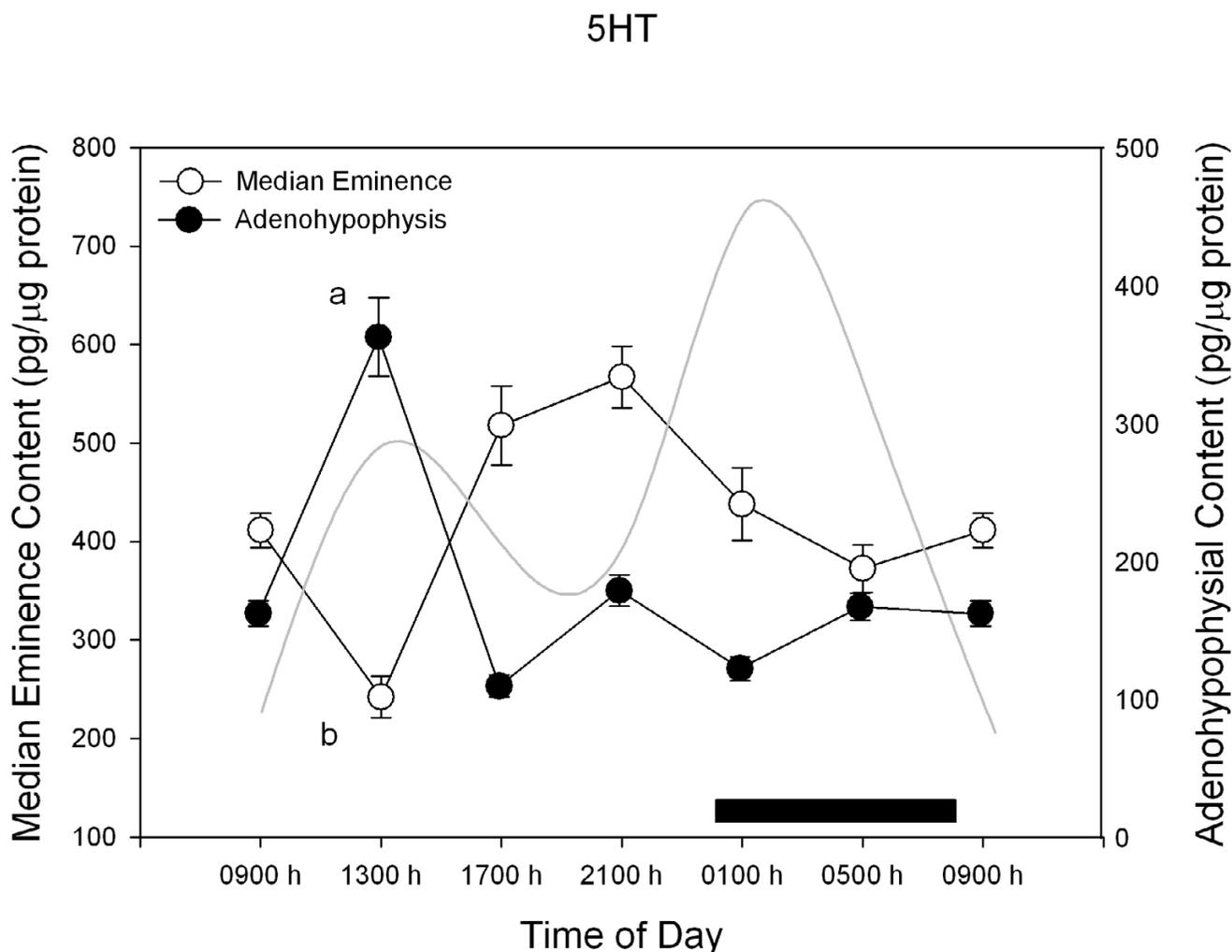


Figure 3
24-h changes in median eminence and adenohipophysial 5HT concentration in 11 days old male rabbit pups.
 Groups of 6–7 pups were killed by decapitation at 6 different time intervals throughout a 24 h cycle. Bar indicates scotophase duration. Results are the means \pm SEM. Circulating prolactin levels are shown in shaded line. Letters indicate the existence of significant differences between time points within each tissue after a Tukey-Kramer’s multiple comparisons test, as follows: ^a $p < 0.01$ vs. all time points. ^b $p < 0.01$ vs. 01:00 h, 09:00 h, 17:00 and 21:00 h. For further statistical analysis, see text.

described in rats [13]. These hypotheses must be tested rigorously (e.g., by using pharmacological blocking agents) before a definitive conclusion can be made.

Among other possible neuromodulators of prolactin secretion, the arcuate nucleus receives a dense serotonergic innervation consisting of a population of brainstem neurons arising mainly from the midbrain raphe nuclei [33] and from fibers originated in 5HT cell bodies located within the hypothalamus. There is a close proximity of

5HT fibers to dopaminergic cell bodies in the arcuate nucleus [34]. Therefore, an indirect effect of 5HT on prolactin release could be linked to the modulation of the inhibitory dopaminergic inputs to the pituitary. Our foregoing results agree with this hypothesis since 5HT concentration in median eminence changes diurnally in an opposite way to that of plasma prolactin levels, albeit without a significant correlation between them. Indeed, previous experiments in rats indicated that 5HT could probably modulate directly the secretion of prolactin [13].

GABA

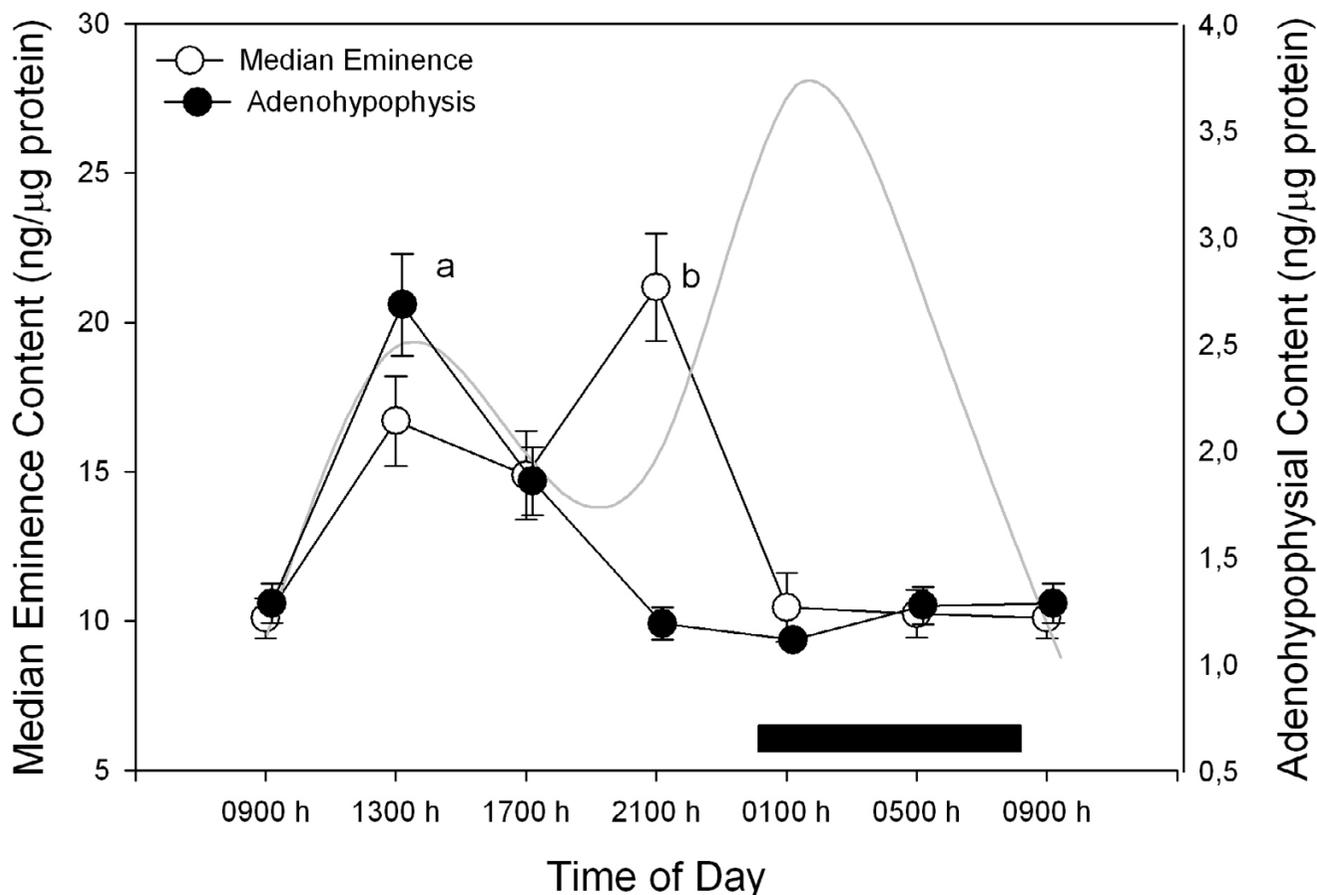


Figure 4
24-h changes in median eminence and adenohypophysial GABA concentration in 11 days old male rabbit pups.
 Groups of 6–7 pups were killed by decapitation at 6 different time intervals throughout a 24 h cycle. Bar indicates scotophase duration. Results are the means ± SEM. Circulating prolactin levels are shown in shaded line. Letters indicate the existence of significant differences between time points within each tissue after a Tukey-Kramer's multiple comparisons test, as follows: ^a p < 0.01 vs. all time points. ^b p < 0.01 vs. 01:00 h, 05:00 h and 09:00 h, p < 0.05 vs. 17:00 h. For further statistical analysis, see text.

Taurine has also been implicated in the regulation of prolactin release [5,13,35,36]. The foregoing results indicate that in median eminence and anterior pituitary of male rabbit pups taurine concentration varies inversely to plasma prolactin levels, displaying a mirror pattern. In the adenohypophysis a negative correlation between plasma prolactin and taurine levels was found, similarly to previous data obtained in rats [13]. Therefore, taurine may play a role in prolactin regulation in newborn rabbits.

A relatively dense innervation of GABA terminals exists in the external layer of the median eminence [37], and the ability of median eminence neurons to release GABA in portal blood has been demonstrated [38]. We previously demonstrated a possibly inhibitory control of GABA on prolactin secretion during the activity phase in male rats [3-6]. Results obtained in the present study in suckling male rabbits support such an inhibitory effect of GABA on plasma prolactin levels exerted mainly during the dark

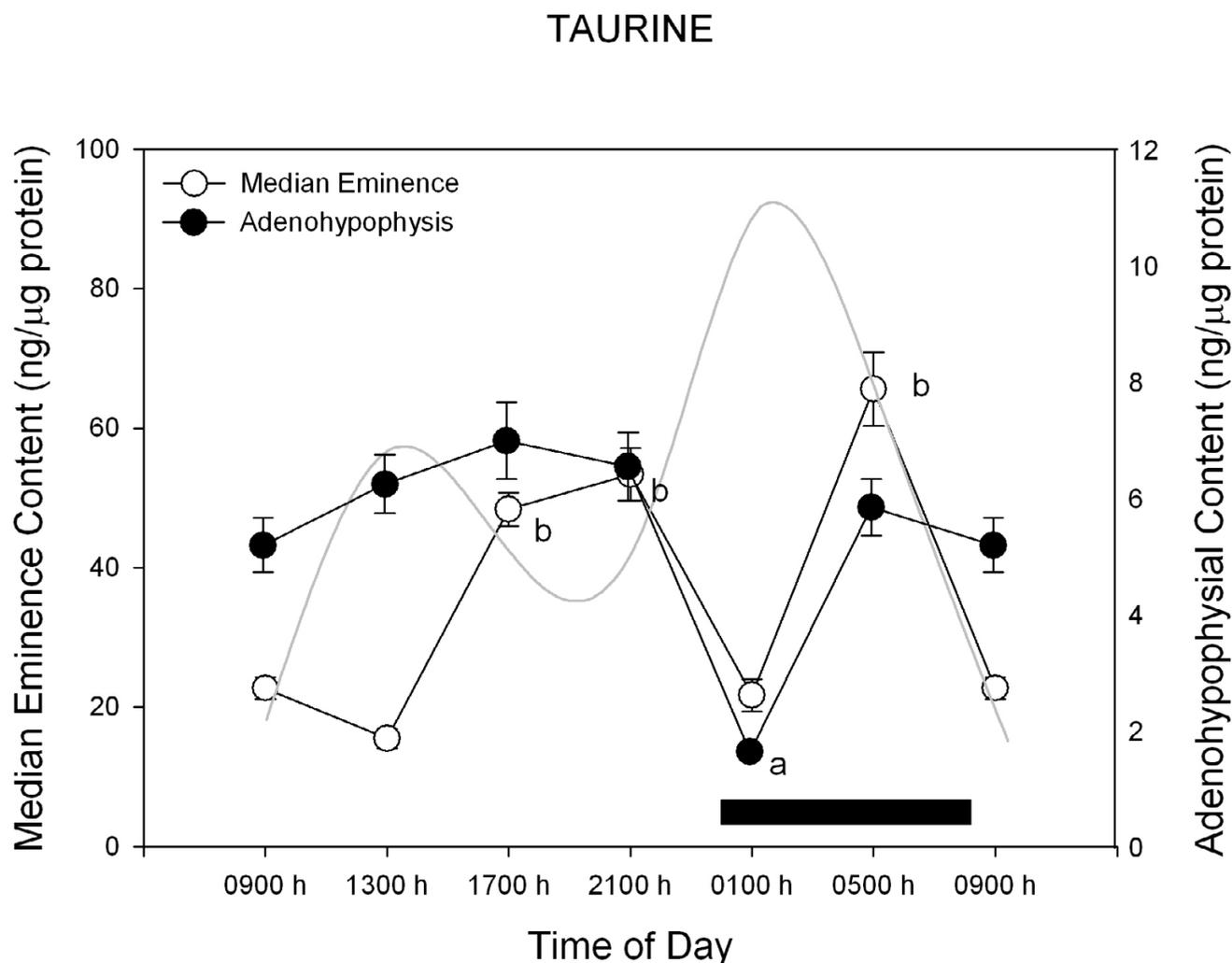


Figure 5
24-h changes in median eminence and adenohypophysial taurine concentration in 11 days old male rabbit pups. Groups of 6–7 pups were killed by decapitation at 6 different time intervals throughout a 24 h cycle. Bar indicates scotophase duration. Results are the means \pm SEM. Circulating prolactin levels are shown in shaded line. Letters indicate the existence of significant differences between time points within each tissue after a Tukey-Kramer's multiple comparisons test, as follows: ^a $p < 0.01$ vs. all time points. ^b $p < 0.01$ vs. 01:00 h, 09:00 h and 13:00 h. For further statistical analysis, see text.

phase of daily photoperiod. The data indicate that GABA concentration in median eminence decreased during the day-night transition, while plasma prolactin levels were increasing. Actually, in median eminence a negative correlation between GABA concentration and plasma prolactin was found, thus suggesting an inhibitory effect of GABA on prolactin secretion.

GABA acting on specific receptors in the anterior pituitary has been reported to suppress prolactin secretion [39,40], although whether this effect was physiological has been

questioned [40]. Data from literature suggest that the role of GABA on prolactin release is quite complex [41]. In some conditions, such as aging [13] or hyperprolactinemia [6], the inhibitory role of GABA becomes more pronounced whereas the inhibitory control exerted by DA diminishes. Our results in male rabbit pups indicated that, although no correlation between plasma prolactin and pituitary GABA concentration was found, the pattern may confirm the main role of this amino acid in the control of prolactin secretion during the dark phase of the photoperiod that was developed later. Again, all these

hypotheses must be tested. e.g. pharmacologically, before a definitive conclusion on this matter can be drawn.

Conclusions

In suckling male rabbits plasma prolactin and median eminence and anterior pituitary concentration of several neuromodulators change on a daily basis. The existence of significant correlations among several of the neurotransmitters analyzed and plasma prolactin levels may explain the circadian secretory pattern of prolactin at this age in suckling rabbits. Collectively, the present results differ from the reported absence of circadian rhythmicity of prolactin and median eminence and adeno-hypophysial neuromodulators in rats at a comparable age.

Competing Interests

The author(s) declare that they have no competing interests.

Authors' Contributions

MPA and PC carried out the experiment and the immunoassays and the analysis of catecholamines, indoleamines and amino acids. DPC and AIE designed the experiments. Also, DPC performed the statistical analysis. PR took care of the experimental animals. AIE supervised its technical implementation and drafted the manuscript. All authors read and approved the final manuscript.

Acknowledgements

This work was supported by grants from DGES, PB9-0257/97, Ministerio de Educación y Cultura, Spain.

References

- Freeman M, Kanyicska B, Lerant A, Nagy G: **Prolactin: structure, function, and regulation of secretion.** *Physiol Rev* 2000, **80**:1523-1631.
- Ben-Jonathan N, Hnasko R: **Dopamine as prolactin (PRL) inhibitor.** *Endocr Rev* 2001, **22**:724-763.
- Casanueva F, Apud AJ, Masotto C, Cocchi D, Locatelli V, Racagni G, Muller E: **Daily fluctuations in the activity of the tuberoinfundibular GABAergic system and plasma prolactin levels.** *Neuroendocrinology* 1984, **39**:367-370.
- Selgas L, Arce A, Esquifino AI, Cardinali DP: **Twenty-four hour rhythms of serum ACTH, prolactin, growth hormone and thyroid-stimulating hormone, and of median eminence norepinephrine, dopamine and serotonin, in rats injected with Freund's adjuvant.** *Chronobiol Int* 1997, **14**:253-265.
- Duvilanski BH, Selgas L, García Bonacho M, Esquifino AI: **Daily variations of amino acid concentration in mediobasal hypothalamus, in rats injected with Freund's adjuvant. Effect of cyclosporine.** *J Neuroimmunol* 1998, **87**:189-196.
- Duvilanski BH, Alvarez MP, Castrillón PO, Esquifino AI: **Daily changes of GABA and taurine concentrations in various hypothalamic areas are affected by chronic hyperprolactinemia.** *Chronobiol Int* 2003, **20**:1-14.
- García Bonacho M, Esquifino AI, Castrillón P, Reyes Toso CF, Cardinali DP: **Age-dependent effect of Freund's adjuvant on 24-h rhythms in plasma prolactin, growth hormone, thyrotropin, insulin, follicle-stimulating hormone, luteinizing hormone and testosterone in rats.** *Life Sci* 1999, **66**:1969-1977.
- Castrillón P, Cardinali DP, Pazo D, Cutrera RA, Esquifino AI: **Effect of superior cervical ganglionectomy on 24-h variations in hormone secretion from anterior hypophysis and in hypothalamic monoamine turnover, during the preclinical phase of Freund's adjuvant arthritis in rats.** *J Neuroendocrinol* 2001, **13**:288-295.
- Esquifino AI, Pazo D, Cutrera RA, Cardinali DP: **Seasonally-dependent effect of ectopic pituitary grafts on 24-h rhythms in serum prolactin and gonadotropins in rats.** *Chronobiol Int* 1999, **16**:451-460.
- Esquifino AI, Chacon F, Jimenez V, Reyes Toso CF, Cardinali DP: **24-hour changes in circulating prolactin, follicle-stimulating hormone, luteinizing hormone and testosterone in male rats subjected to social isolation.** *J Circadian Rhythms* 2004, **2**:1.
- Chacón F, Cano P, Jimenez V, Cardinali DP, Marcos A, Esquifino AI: **24-hour changes in circulating prolactin, follicle-stimulating hormone, luteinizing hormone and testosterone in young male rats subjected to calorie restriction.** *Chronobiol Int* 2004, **21**:393-404.
- Cano P, Cardinali DP, Castrillón P, Reyes Toso C, Esquifino AI: **Age dependent changes in 24-h rhythms of catecholamine concentration and turnover in hypothalamus, corpus striatum and pituitary gland of rats injected with Freund's adjuvant.** *BMC Physiology* 2001, **1**:14.
- Esquifino AI, Cano P, Jiménez , Reyes Toso CF, Cardinali DP: **Changes of prolactin regulatory mechanisms in aging: 24-h rhythms of serum prolactin and median eminence and adeno-hypophysial concentration of dopamine, serotonin, gamma aminobutyric acid, taurine and somatostatin in young and aged rats.** *Exp Gerontol* 2004, **39**:45-52.
- Dunaway JE: **Alteration in timing of PMS-induced ovulation following pinealectomy.** *Neuroendocrinology* 1969, **5**:281-289.
- Tresguerras JAF, Esquifino AI, López-Calderón A: **Effects of estradiol benzoate and castration on LH in experimental hyperprolactinemia.** *J Steroid Biochem* 1983, **19**:447-453.
- Moreno ML, Villanua MA, Esquifino AI: **Serum prolactin and luteinizing hormone levels and the activities of hypothalamic monoamine oxidase A and B and phenylethanolamine-N-methyl transferase are changed during sexual maturation in male rats treated neonatally with melatonin.** *J Pineal Res* 1992, **13**:167-173.
- Velázquez E, Esquifino AI, Zueco JA, Ruiz Albusac JM, Blázquez E: **Evidence that circadian variations of circulating melatonin levels in fetal and suckling rats are dependent on maternal melatonin transfer.** *Neuroendocrinology* 1992, **55**:321-326.
- Esquifino AI, Arce A, Villanúa MA, Cardinali DP: **Development of 24-hour rhythms in serum prolactin and luteinizing hormone levels in rats neonatally administered melatonin.** *Chronobiol Int* 1998, **15**:21-28.
- Davis FC, Gorski RA: **Development of hamster circadian rhythms. III. Role of the maternal suprachiasmatic nucleus.** *J Comp Physiol [A]* 1988, **162**:601-610.
- Davis FC, Mannion J: **Entrainment of hamster pup circadian rhythms by prenatal melatonin injections to the mother.** *Am J Physiol* 1988, **255**:R439-R448.
- Klein DC, Lines SU: **Pineal hydroxyindole-O-methyltransferase activity in growing rat.** *Endocrinology* 1969, **84**:1523-1525.
- Reppert SM, Klein DC: **Transport of maternal ³H-melatonin to suckling rats and the fate of ³H-melatonin in neonatal in the neonatal rat.** *Endocrinology* 1978, **102**:582-588.
- Manning PJ, Ringler DH, Newcomer CE: **The Biology of the Laboratory Rabbit.** New York: Academic Press; 1994.
- Thompson HV, King CM: **The European Rabbit.** Oxford University Press, Oxford; 1994.
- Jilge B, Hudson R: **Diversity and development of circadian rhythms in the European rabbit.** *Chronobiol Int* 2001, **18**:1-26.
- Takahashi K, Ohi K, Shimoda K, Tamada N, Hayashi S: **Postnatal maternal entrainment of circadian rhythms.** In *Development of circadian rhythmicity and photoperiodism in mammals. Research in perinatal medicine Volume 9.* Edited by: Reppert SM. New York : Ithaca, Perinatology Press; 1989:67-82.
- Allingham K, von Saldern C, Brennan PA, Distel H, Hudson R: **Endogenous expression of c-Fos in hypothalamic nuclei of neonatal rabbits coincides with their circadian pattern of suckling-associated arousal.** *Brain Res* 1998, **783**:210-218.
- Ubilla E, Alvarioño JMR, Esquifino AI, Agrasal C: **Effect of induction of parturition by administration of a prostaglandin F2 analogue in rabbits: possible modification of prolactin LH and FSH secretion patterns.** *Anim Reprod Sci* 1992, **27**:13-20.

29. Greenwood FC, Hunter NWM, Glover JS: **The preparation of ¹³¹I-labelled human growth hormone of high specific radioactivity.** *Biochem J* 1963, **89**:114-123.
30. García-Bonacho M, Cardinali DP, Castrillón P, Cutrera RA, Esquifino AI: **Aging-induced changes in 24-h rhythms of mitogenic responses, lymphocyte subset populations and neurotransmitter and amino acid concentration in rat submaxillary lymph nodes during Freund's adjuvant arthritis.** *Exp Gerontol* 2001, **36**:267-282.
31. Hudson R, Distel H: **The temporal pattern of suckling in rabbit pups: A model of circadian synchrony between mother and young.** In *Development of circadian rhythmicity and photoperiodism in mammals* Edited by: Reppert SM. Boston: Perinatology Press; 1989:83-109.
32. Moore KE: **Interactions between prolactin and dopaminergic neurons.** *Biol Reprod* 1987, **36**:47-58.
33. Steinbusch HWM: **Distribution of serotonin-immunoreactivity in the central nervous system of the rat-cell bodies and terminals.** *Neuroscience* 1981, **6**:557-618.
34. Kiss J, Halasz B: **Synaptic connections between serotonergic axon terminals and tyrosine hydroxylase-immunoreactive neurons in the arcuate nucleus of the rat hypothalamus. A combination of electron microscopic autoradiography and immunocytochemistry.** *Brain Res* 1986, **364**:284-294.
35. Login IS: **Direct stimulation of pituitary prolactin release by glutamate.** *Life Sci* 1990, **47**:2269-2275.
36. Arias P, Jarry H, Convertini V, Ginzburg M, Wuttke W, Moguilevsky J: **Changes in mediobasal hypothalamic dopamine and GABA release: A possible mechanism underlying taurine-induced prolactin secretion.** *Amino Acids* 1998, **15**:5-11.
37. Vincent SR, Hokfelt T, Wu JY: **GABA neuron systems in hypothalamus and the pituitary gland.** *Neuroendocrinology* 1982, **34**:117-125.
38. Gudelsky G, Apud J, Masotto C, Locatelli V, Cocchi D, Racagni G, Muller E: **Ethanolamine-O-sulfate enhances g-aminobutyric acid secretion into hypophysial portal blood and lowers serum prolactin concentrations.** *Neuroendocrinology* 1983, **37**:397-399.
39. Ondo JG, Dom R: **The arcuate nucleus: a site for gamma-aminobutyric acid regulation of prolactin secretion.** *Brain Res* 1986, **381**:43-48.
40. Lee TY, Pan JT: **Involvement of central GABAergic neurons in basal and diurnal changes of tuberoinfundibular dopaminergic neuronal activity and prolactin secretion.** *Life Sci* 2001, **68**:1965-1975.
41. Apud JA, Cocchi D, Locatelli V, Masoto C, Muller ER, Racagni G: **Biochemical and functional aspects on the control of prolactin release by the hypothalamo-pituitary GABAergic system.** *Psychoneuroendocrinology* 1989, **14**:3-17.

Publish with **BioMed Central** and every scientist can read your work free of charge

"BioMed Central will be the most significant development for disseminating the results of biomedical research in our lifetime."

Sir Paul Nurse, Cancer Research UK

Your research papers will be:

- available free of charge to the entire biomedical community
- peer reviewed and published immediately upon acceptance
- cited in PubMed and archived on PubMed Central
- yours — you keep the copyright

Submit your manuscript here:
http://www.biomedcentral.com/info/publishing_adv.asp

