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Mothers Have Lower Testosterone Than Non-Mothers: Evidence from the Philippines

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Abstract

Testosterone (T) is lower among fathers and men in committed relationships, suggesting that the hormone mediates the trade-off between mating and parenting effort. The function of T in women, and responses of the hormone to relationships or motherhood, are less well understood. Here we report relationships between T and pairbonding and motherhood in a random sample of 67 women (20.9 to 22.4 years old) participating in a population-based birth cohort study in the Philippines. Testosterone was measured in saliva collected at bedtime and at waking the following morning to capture circadian dynamics. Compared to non-mothers and non-pairbonded women, mothers and pairbonded women had 32% ($p < 0.0001$) and 23% ($p < 0.004$) lower waking T, respectively, but similar evening T. The lower waking T in mothers largely reflected reduced T in mothers of young offspring (< 2 years), with mothers of older offspring (2+ years) having intermediate T. These differences were independent of measures of breastfeeding, contraceptive pill use, menstrual cycle, sleep quality, education, employment and socioeconomic status. Our findings highlight a similar relationship between parenting and committed relationships and T in women as documented in men, and suggest that caregiving of dependent young may modulate female T. Future research should clarify whether this cross-sectional association reflects a suppressive effect of motherhood on T, whether these relationships vary across cultures, and the role of T within the endocrine architecture regulating female reproductive and caregiving strategies.

Keywords

reproduction; life history; parenting; sex steroids; behavioral ecology

INTRODUCTION

The role of testosterone (T) as a mediator of reproductive strategies has been the focus of considerable attention, with most work conducted on males. In human males T is proposed to mediate reproductive strategy in part through anabolic effects on sexually dimorphic traits, including skeletal growth, lean mass and strength (Bhasin et al., 2001; Sinha-Hikim et

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al., 2003), and also through effects on behaviors related to competition (Elias, 1981; Mazur et al., 1992; Salvador et al., 2003) and libido (Anderson et al., 1992; Wang et al., 2000). In many birds and mammals, T is elevated during periods of competition and mating, and is reduced in the context of caregiving in some species with biparental care (Goymann et al., 2007; Wingfield et al., 1990; Wynne-Edwards, 2001). There is evidence for a similar responsiveness to pairbonding and caretaking status among human males. Men who are in stable pairbonds have been shown to have lower T than non-pairbonded men in some (Booth and Dabbs, 1993; Gray et al., 2004; Mazur and Michalek, 1998; van Anders and Watson, 2006) but not all populations investigated (Flinn et al., 1998; Gray, 2003; Gray et al., 2006). Fathers also tend to have lower T than single men (Berg and Wynne-Edwards, 2001; Gray et al., 2002; Muller et al., 2009), and there is evidence that these relationships are stronger than with pairbonding alone in some (Gray et al., 2006; Kuzawa et al., 2009) but not all populations (Gray, 2003; Gray et al., 2004). These findings in males have been interpreted as support for the hypothesis that T mediates the tradeoff between behavioral and energetic effort devoted to mating and parenting in humans, much as it does in other taxa (Goymann et al., 2007; Wingfield et al., 1990).

Comparably few studies have focused on the importance of motherhood or pairbonding status to T levels in females, but evidence to date highlights similar relationships to those observed in males. Females have been found to exhibit differences in T based on relationship status (van Anders et al., 2007b), sexual orientation and pairbonding (van Anders and Watson, 2006) and whether they reside in the same city as their pairbond partner (van Anders and Watson, 2007). Women with multiple, simultaneous romantic partners showed higher T compared to single women or monogamous women (van Anders et al., 2007b), not unlike the finding of higher T in Swahili men with multiple wives (Gray, 2003). There is some evidence that testosterone promotes behaviors related to mating activity in females, including libido (Braunstein et al., 2005; Buster et al., 2005; Davis et al., 2008; Guay, 2001; van Anders et al., 2007a), competition (Edwards et al., 2006; Oliveira et al., 2009), dominance (Grant and France, 2001), and aggression (Harris et al., 1996). Much as has been argued for men, a tendency to reduce expression of these behaviors might provide reproductive benefits to females when priorities have shifted from securing mates to caring for dependent young (Fite et al., 2005).

To our knowledge, no study to date has specifically evaluated whether motherhood predicts T in women. In this journal, we recently documented that fathers living in Cebu City, the Philippines, have lower T than non-fathers (Kuzawa et al., 2009) and that, unlike in other samples investigated (Gray et al., 2002; Burnham et al., 2003; Gray et al., 2004), there was no association with pairbonding independent of fatherhood status in this population. Working with a same-aged sample of women from this population, here we report T in waking and pre-bedtime saliva samples, which we relate to pairbonding and parenting status while adjusting for potentially important confounding factors.

MATERIALS AND METHODS

Study population

Data come from the Cebu Longitudinal Health and Nutrition Survey (CLHNS), a population-based birth cohort study that has tracked a large cohort of men and women who were born in 1983-84 (Adair et al., 2001; Adair et al., 1993; Kuzawa and Adair, 2003). Women were 20-22 years of age at the time of interview and sample collection. Body weight (kg) and height (cm) were measured using standard anthropometric techniques (Lohman et al., 1988). The body mass index (BMI) was calculated as the ratio of weight (kg)/height (m²). Dietary intake was measured using two 24-hour recalls on consecutive days and the mean was used in analyses. Energy intake was calculated using Philippines

Food Composition Tables produced by the Food and Nutrition Research Institute of the Philippines (FNRI, 1997). Information on current pregnancy status, self-reported contraceptive use, breastfeeding, and a range of socioeconomic, educational and employment questions were obtained in interviews. Participants in this study have traditionally been compensated for their time with a modest “bother fee” (\$2 USD), which was also provided in this study. This research was conducted under conditions of informed consent with human subjects clearance from the Institutional Review Boards of Northwestern University and the University of North Carolina, Chapel Hill.

Motherhood and pairbonding—Women were classified as pairbonded if they reported currently living with a partner and/or were married. Using the household roster, mothers were defined as women who were living with one or more biological offspring, with age of offspring also noted. Although we did not have information on offspring living outside of the household, this was not likely an important issue. First, in Cebu mothers are the primary caregivers for dependent offspring, and offspring raised away from their mothers are likely rare (Liwag et al., 1998). Second, past research has demonstrated the importance of direct interaction and caregiving as an influence on maternal endocrine profile, which is of greatest interest to our research questions; thus, the rare mother who was not living with and raising her own offspring, and thus would be misclassified as a non-mother in our study, might be expected to share more hormonal similarities with non-mothers (e.g. Feldman et al., 2007; Fleming et al., 1997).

Menstrual cycle changes—Circulating T in females originates from adrenal and ovarian sources (Abraham 1974). Although some have concluded that menarcheal changes in T are minor enough to be ignored in behavioral research (Dabbs 1990), the ovarian contribution to circulating T changes across the menstrual cycle, and has generally been found to rise late in the follicular phase (Abraham 1974). In a study that used a sensitive equilibrium dialysis method capable of measuring free T at the low concentrations found in women, free T in 34 healthy women (the closest correlate of the salivary T measured here) was found to reach a pre-ovulatory peak late in the follicular phase, at 3 days prior to the LH peak, then gradually declined (Sinha-Hikim et al., 1998). To account for this potential influence on T, we estimated which women were in the late follicular phase at the time of saliva collection. During in-home interviews, women who did not report being pregnant were asked the timing of their last menstrual period. From this information we classified women as being in the late follicular phase if they reported that their last menstrual period began 7-14 days prior to saliva collection. In the event that saliva collection was more than 28 days after the interview, we assumed a 28-day cycle.

Lactation suppresses gonadotropin secretion (McNeilly, 1980) and has been reported to reduce testosterone in some but not all studies (e.g. Alder et al., 1986; Dada and Laditan, 1982). In addition, lactational suppression of ACTH could reduce production of adrenal androgens, with secondary effects on the adrenal contribution to circulating T. To account for possible lactational influences on T, we tested whether women who were currently breastfeeding had lower T, and if so, the extent to which this contributed to any differences seen in T among mothers, and especially, mothers of infants and young children of breastfeeding age.

Contraceptive pill use can reduce free T via effects on ovarian T production and also by increasing sex hormone binding globulin concentrations (Greco et al., 2007). Although their use is relatively uncommon in this population, we controlled for oral contraceptive use, self-reported during in-home interviews, as a predictor and potential confounding influence on the associations that we investigated.

Saliva collection

Each participant was provided with instructions and two tubes for saliva collection. The first sample was collected immediately prior to bed. After collection, the participant sealed the tube and kept it at room temperature. Participants were instructed to place the second tube next to their bed and to collect the second sample immediately upon waking the following morning. Respondents reported time of saliva collection, and at waking, wake time, usual wake time, and a 5-level self-rated scale reflecting sleep quality the prior night. Both tubes were collected later that day by an interviewer, who placed the tubes on ice packs in a cooler. They were then transported to a freezer where they were stored at -35 C until shipment on dry ice to the Laboratory for Human Biology Research at Northwestern University, where they were stored at -80 C. They were thawed, supernatant separated, and aliquoted into smaller tubes for analysis of individual analytes.

Testosterone measurement

Salivary T concentrations were determined in the Laboratory for Human Biology Research at Northwestern University using an enzyme immunoassay protocol specifically developed for use with saliva samples (Salimetrics No. 1-2402, State College, PA). All samples were run in duplicate, and low and high controls were included with each assay. The intra-assay coefficients of variation were 2.5% and 6.7% for high and low controls, respectively. The inter-assay coefficients of variation were 5.6% and 6.7%, respectively.

Sample selection

During the 2005 survey, 893 females of the original cohort of 1477 liveborn females, ages 21.5 years (range 20.8-22.6 years), were located and interviewed. Of these, 835 individuals agreed to participate in saliva collection. From these samples, we randomly selected saliva samples from 67 non-pregnant women for testosterone measurement. Baseline characteristics of the final sub-sample of 67 women who were included in the testosterone analyses were compared with those who were in the sample at baseline but excluded from the analysis. There were no significant or borderline significant differences (all $p > 0.1$) in baseline characteristics between the testosterone analysis sub-sample and the remaining original baseline cohort, respectively, in household income (311 vs. 312 pesos), birth weight (3005 vs. 2993 g), mother's height (150.4 vs. 150.5 cm), mother's weight at parturition (50.1 vs. 50.1 kg), mother's age (27.1 vs. 26.0 years), mother's highest completed grade (6.5 vs. 7.1) or father's highest completed grade (6.9 vs. 7.3), suggesting that sample inclusion was not biased.

Statistical analysis

All analyses were performed with version 10 of Stata (Stata Corporation, College Station, TX). Anthropometric, evening and waking T and daily energy intake (kcal) were all analyzed as continuous variables. We first used linear regression to test for significant differences in waking and evening T in relation to pairbonding and motherhood status and also to identify important potential confounding influences on T, adjusting only for time of sample collection, wake time that day and usual wake time. Next, we used multiple linear regression to evaluate relationships between waking and evening T and both motherhood and pairbonding while adjusting for covariates and potential confounders. Final models considered the age of the youngest offspring as a predictor, with mothers categorized as having young offspring (<2 years) and older offspring (2+ years) adjusting for time of sample collection, wake time that day and usual wake time, contraceptive pill use and menstrual cycle phase.

RESULTS

The sample included 67 women with an average age of 21.5 years (range 20.9 to 22.4 years), of whom 36% were mothers. Only 7% of non-mothers lived in households headed by themselves (and their partner), while more than half of mothers lived in their own households. Consistent with this, mothers were more likely to be employed, were less likely to be currently enrolled in school, and lived in lower income households. They were also more likely to be pairbonded or to report currently using oral contraceptives (Table 1). When compared to non-mothers, mothers had lower unadjusted waking T but not lower evening T.

Table 2 reports mean waking and evening T stratified on motherhood and pairbonding status in addition to three potential confounding factors known to influence T levels in females: using contraceptive pills, breastfeeding and being in the late follicular phase of the menstrual cycle. Measures of socioeconomic status (per capita household income, assets scale, currently employed), education (highest grade completed), sleep quality the night prior to waking sample collection (5-level self-rated scale) and daily caloric intake were not significant or borderline significant as predictors of waking or evening T (all $p > 0.3$) and are therefore not shown. Mothers had lower waking T but not evening T compared to non-mothers. When mothers were stratified according to the age of their youngest offspring, mothers with young offspring (< 2 years of age) had significantly lower waking T and evening T compared to non-mothers, while mothers of older offspring (2+ years) only had borderline significantly lower T and only in the morning. Pairbonded women had lower waking T but their evening T was similar to that of non-pairbonded women. As expected, women late in the follicular phase of the menstrual cycle had higher waking T compared to women in other phases of the menstrual cycle, while contraceptive pill users had lower T at both times of day compared to non pill users.

Although women who were currently breastfeeding had lower waking T and borderline lower evening T compared to women not currently breastfeeding (Table 2), breastfeeding did not appear to be an important influence on the lower T found in mothers. Although mothers as a group had lower T than non-mothers, of the 14 mothers with young offspring, the 7 who were breastfeeding had modestly higher waking T (61.5 pg/ml) compared to the 7 who were not breastfeeding (47.9 pg/ml). Consistent with this, breastfeeding was no longer associated with lower T in the morning (-0.7 pg/ml, $p < 0.954$ compared to non-breastfeeders) or in the evening (-2.3 pg/ml, $p < 0.775$ compared to non-breastfeeders) after adjusting for being a mother of a young offspring (<2 years).

We next ran a series of regression models designed to disentangle the independent associations of motherhood and pairbonding status with T. Given the low frequency in our sample of mothers who were not pairbonded ($n=4$) and of pairbonded non-mothers ($n=5$), we caution that these results should be interpreted as preliminary. Adjusting for usual wake time, wake time that day and time of saliva collection, waking T was similar in the 5 non-mothers who were pairbonded and the 38 non-mothers who were not pairbonded (86.5 pg/ml vs. 94.0 pg/ml, respectively, $p < 0.4$). Of the 24 mothers, the 4 who were not pair-bonded had similar mean waking T to the 20 mothers who were pairbonded (53.9 pg/ml vs. 65.3 pg/ml, $p < 0.7$). When motherhood and pairbonding were considered simultaneously in a regression model that adjusted for circadian time variables, mothers had a significant 28.1 pg/ml (SE 10.9 pg/ml) reduction in waking T compared to non-mothers ($t = -2.58$, $p < 0.012$), while pairbonding status was no longer a significant predictor of waking T after adjusting for motherhood (-3.6 pg/ml, SE 10.8 pg/ml, $p < 0.737$).

Final regression models evaluated relationships between motherhood status and waking and evening T adjusting for potential confounding influences identified above. In final models (Table 3) mothers of young offspring had lower T in the morning (significant) and evening (not significant) compared to non-mothers, adjusting for time of saliva collection, usual wake time, and wake time that day, contraceptive pill use and being in the late follicular phase of the menstrual cycle (Figure 1). Mothers of older offspring had non-significantly lower waking T than non-mothers, and a statistically significant but modest (~6 pg/ml) increase in evening T compared to non-mothers.

DISCUSSION

In this sample of Filipino women, mothers had lower waking T than non-mothers, with differences greatest among mothers with younger offspring. Although the number of breastfeeding women was relatively small, available data suggested minimal contributions of lactation as an influence on these differences. Although pairbonded women had lower T than non-pairbonded women, these differences were more modest than those seen in mothers, and, although small sample size limited strong inferences, were largely accounted for by the association between motherhood and T. The present findings are to our knowledge the first demonstration that T is lower in the context of parenting among women.

Our findings among the females in this sample share similarities to patterns that we recently reported in males from the same cohort (Kuzawa et al., 2009), and that have been documented in males in other populations (Gray et al., 2006). Notably, in both men and women at Cebu, parenting is a strong predictor of lower T levels, while pairbonding status was less important as an independent predictor of T. Similar studies of T in females in other populations have focused on relationship status as a predictor, without accounting for parenting status, and have found mixed evidence for lower T in pairbonded women. Similar to our results, two studies have documented no relationship between pairbonding and T in heterosexual women (van Anders et al., 2007b; van Anders and Watson, 2006). However, another study, which controlled for sexual orientation, showed that women living in the same city as their partner had lower T levels compared to single women and those in long distance relationships (van Anders and Watson, 2007).

Why we find evidence for lower T specific only to motherhood—with no difference by pairbonding status alone—is uncertain, but could trace to recent cultural shifts in the nature of pairbonding in Philippine society. Long-term pairings in which men and women are not legally married are increasingly common in Cebu (NSO, 2004). The growing incidence of couples avoiding legal marriage may reflect an effort of men and women to circumvent the lack of legalized divorce in the Philippines (Gloria, 2007). While it remains speculative, such non-legalized pairings may involve different endocrine responses than are elicited by formalized marital relations that have typically been practiced in the Philippines in the past.

While traditional patterns of marriage and pairbonding may have shifted in the Philippines, levels of maternal involvement with their offspring have remained relatively stable (Liwag et al., 1998; Medina, 2001). Although fathers have traditionally played a relatively minor role in offspring care, this is gradually changing, and sibling care is also quite prevalent (ECD, unpublished data; Harper, 2010; Kuzawa et al., 2009). However, in general, girls are socialized into domestic, parenting roles from a young age and mothers remain the primary source of nurturing and caregiving for their children across socioeconomic strata in the Philippines (Liwag et al., 1998). This remains the case even as millions of women enter the workforce to help support the family financially (Medina, 2001). Our findings suggest cross-cultural differences in the relationship between pairbonding and T, highlighting the need for

future comparative work to be couched in a biocultural framework that addresses the similarities and differences in the roles related to pairbonding, marriage and childcare.

The similarities in relationships between T and pairbonding and parenting among males and females in Cebu are striking. Recent comparative studies also document similarities in T response and function in the males and females of other taxa, such as fish and birds (Moller et al., 2005). For instance, in certain fish taxa, the degree of male sexually-selected ornamentation is positively related to peak female androgen levels, suggesting that sexual selection for higher T in males has correlated consequences on T levels in females of that species (Mank, 2007). It is presently unclear how sex differences in T might co-vary across species in mammals or primates, or the degree to which androgen levels, or related behavioral and somatic effects in women, might be a correlated consequence of selection on hormone regulation and target tissue effects in human males (Ketterson et al., 2005). The broad similarities in the relationship between parenting status and T in males and females in our sample underscore the need to investigate this possibility.

It is also possible that T has effects that are adaptive to females, and thus have been shaped by natural selection for a female-specific function. There is evidence from other species that selection has influenced the level of T and its behavioral effects in females independent of effects in males, as suggested for instance by the finding that peak female T is highest in colonial bird species in which females play a prominent role in territorial defense (Moller et al., 2005). Although less research into T has been conducted in female primates, the work to date supports the hormone's importance as a modulator of female reproductive strategies. For instance, in Wied's black tufted-ear marmosets (a species of New World monkey), mothers have been found to exhibit reduced caregiving effort in association with elevated T (Fite et al., 2005). In the great apes, T is higher for more of the cycle in chimpanzees than in gorilla or orangutan, which has been related to the longer period of mating during each cycle in chimpanzees (Nadler et al., 1985).

Although conflicting evidence exists (e.g. Davis et al., 2005; Kivlighan et al., 2005; Mazur et al. 1997), it has been speculated that T is associated with a wide range of behaviors in women that may influence mate choice, sexual behavior, and parenting, which could improve mating success among non-mothers (Braunstein et al., 2005; Buster et al., 2005; Davis et al., 2008; Grant and France, 2001; Guay, 2001; Harris et al., 1996; van Anders et al., 2007a). To the extent that the lower T observed among mothers in our sample reflects a hormonal response to motherhood, rather than a trait-level predictor of motherhood status, reduced T in the context of motherhood could help shift female priorities away from mating and in favor of caretaking and parenting, much as has been proposed for men (Gray and Campbell, 2009).

It is important to note, however, that the sources of T are distinct in men and women. In women, the concentration of free T, which is indexed by the salivary measures used in this study, is a function of steroid production in the adrenal and ovaries and binding by sex hormone binding globulin. Lower T in mothers likely traces to a combination of factors that shift across the menstrual cycle and with changing patterns and intensity of offspring investment. For instance, breastfeeding has been associated with suppression of ovarian and adrenal hormone production, which could influence circulating T. To the extent that T motivates mating-related behaviors in women (Buster et al., 2005), such a mechanism could help shift metabolic and behavioral priorities away from initiating a new pregnancy during the most energetically intensive period of investment in the current offspring (Valeggia and Ellison 2009). Although we did find that breastfeeding women had lower T, this association was weaker than the association between T and motherhood, and mothers of young offspring who were not breastfeeding actually had modestly lower T than women who were

breastfeeding. In light of our small sample sizes, we emphasize the need to investigate these questions with a larger sample of mothers and with more variation in lactational status.

In addition to any direct effects of reproductive or energetic status on hormone production, past work has demonstrated that exposure to infant cues, such as odors or recorded cries, development of maternal attachment and mother-infant bonding, and rates of affectionate maternal behavior, can directly elicit changes in neuroendocrine regulation and hormonal levels in mothers (Feldman et al., 2007; Fleming et al., 1997a; Fleming et al., 1997b; Storey et al., 2000). These studies have focused primarily on cortisol, oxytocin, and female sex steroids, such as estradiol and progesterone, and comparably little is known about female T responses to offspring cues and caregiving. Most studies of T in mothers have focused on the immediate postpartum period, and results have been inconsistent. Still, in the early postpartum period higher T in mothers has been associated with reduced affectionate contact and positive feelings about the infant (Fleming et al., 1997b), and also with maternal depression and higher anger scores (Hohlagschwandtner, 2001).

Although our data are cross-sectional, the patterns that we document are consistent with an effect of caretaking on female T levels. Mothers play a central role as caretakers in the Philippines, and often have primary responsibility for meeting a child's needs. In our sample, less than 50% of the fathers listed themselves as individuals within the household who were responsible for childcare (Kuzawa et al., 2009). Fathers who reported being involved in childcare had particularly low evening T levels, which we speculated could indicate a direct role of interacting with the child across the day on T suppression. Similarly, our present finding that T was lowest among mothers with young offspring, whereas mothers of older children had intermediate T levels, could indicate that the more frequent and demanding activities of caring for infants and young children, compared to older offspring, contribute to the lower T in these women. Our findings provide a strong rationale to include T measurement in future endocrine studies of maternal caretaking.

Several limitations of this study warrant mention. First, we only have a single waking and evening saliva sample for each participant. Past work on T has shown that collecting samples across several days and using the average in analyses enhances reliability (Dabbs, 1990). Similarly, a validation study that compared serum free and total T to salivary T measured with an enzyme immunoassay protocol similar to that used here found that relationships between salivary measures and total T were similar in males and females ($r = 0.54$ and 0.56 , respectively), but that relationships with serum free T--reflecting the unbound fraction--were weaker in females ($r = 0.42$) than in males ($r = 0.64$) (Shirtcliff et al., 2002). Another potential source of error traces to menstrual changes in T, which we adjusted for using each woman's recalled timing of last menstrual period and assuming a 28-day cycle for each woman. Repeat measurement of blood samples would have allowed more precise characterization of menstrual changes in T, which are secondary to the contribution of ovarian T to the pool of circulating T. Unless these factors influence mothers and non-mothers differently in our sample, these various sources of error are not likely to introduce bias but will merely limit our ability to detect statistically-significant relationships between T and social context (Shirtcliff et al., 2002). It is possible that some of the borderline relationships in our study, such as the lower evening T among mothers, would have reached the level of statistical significance with better characterization of typical T levels and with better statistical control of each woman's position in the menstrual cycle.

As is the case with most prior studies of T and social context in males or females, our data are cross-sectional and thus do not permit us to distinguish whether women with high T are less likely to become pregnant from the alternative that mothering has a direct suppressive effect on T. Both trait and state effects are possible, and both could occur simultaneously.

Although only circumstantial evidence, the finding that T is lowest in mothers of young infants and children, and intermediate in mothers of older children, makes a trait-level relationship less likely, as it suggests that mothers may have relatively high or low levels of T depending on the age of their currently youngest offspring. A longitudinal design that follows women prospectively as they transition from non-mothers into motherhood, and that assesses acute T responses to interacting with offspring, will be necessary to more definitively establish the causal direction of effects underlying the relationships that we document.

In summary, in a population of young Filipino women, T levels were roughly one third lower in mothers than in non-mothers in the morning, but there were no differences in T measured in the evening prior to bed. Differences were larger and more significant among mothers with young infants, perhaps indicating a role of demanding caregiving as an influence on T. Although small sample sizes precluded definitive tests of their independent associations, T was more strongly related to motherhood status than to pairbonding status in these women. These findings are similar to our recent findings among the males in this cohort, in which parenting but not pairbonding predicted lower T (Kuzawa et al., 2009). Future research is needed to clarify cultural variation in these relationships, establish whether motherhood lowers T in a longitudinal design, and assess the possible role of reduced T in female caretaking and reproductive strategies.

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Fig. 1.

Mean ($\pm$ SD) waking and evening salivary free testosterone in non-mothers, mothers whose youngest offspring is less than 2 years of age, and mothers of older children (2+ years of age). Sample sizes in parentheses. Adjusted for time of saliva measurement, usual wake time, wake time that day, oral contraceptive use and being in the late follicular stage of the menstrual cycle. Significant differences: Waking T, mothers of young offspring vs. older offspring $p < 0.051$, mothers of young offspring vs. non-mothers $p < 0.002$; Evening T, mothers of older offspring vs. non-mothers ($p < 0.044$) and mothers of young offspring ($p < 0.004$).

Table 1

Characteristics of sample.

	All (n=67)	Non-mothers (n=43)	Mothers (n=24)	p ^a
Biological variables				
Waking testosterone (pg/ml)	82.5 (33.0)	93.1 (32.3)	63.4 (25.1)	0.0002
Evening testosterone (pg/ml)	45.0 (20.1)	47.6 (20.3)	40.3 (19.3)	0.156
Collection time (waking)	6.8 (2.3)	6.8 (2.3)	6.5 (1.1)	0.576
Collection time (evening)	20.6 (4.8)	20.2 (5.9)	21.5 (1.4)	0.261
Height (cm)	150.3 (5.6)	150.7 (5.8)	149.6 (5.3)	0.434
BMI (kg/m ²)	20.5 (3.1)	20.2 (3.0)	21.0 (3.2)	0.281
Energy intake (kcal/day)	1609 (860)	1528 (680)	1755 (1115)	0.502
Triceps skinfold (mm)	20.2 (5.9)	20.4 (6.2)	20.0 (5.5)	0.921
Social context				
Age (years)	21.5 (0.3)	21.5 (0.3)	21.6 (0.3)	0.319
Education (highest grade)	11.1 (3.1)	11.9 (3.1)	9.9 (2.7)	0.011
Household size (persons)	6.5 (2.9)	6.6 (2.7)	6.2 (3.3)	0.470
Per capita weekly				
household income (pesos)	93 (104)	109 (122)	66 (52)	0.030
Lives in own household (%)	23.1%	7.0%	54.6%	0.0001
Reproductive status				
Uses oral contraceptives (%)	13.4%	2.3%	33.3%	0.0001
Pairbonded (%)	37.3%	11.6%	83.3%	0.0001
Currently breastfeeding (%)	13.4%	-	37.5%	-
Mother (%)	35.8%			

Mean (SD) unless otherwise noted

^a p-value from 2-sided t-test or chi-square comparing mothers and non-mothers

Table 2

Mean waking and evening testosterone (pg/ml) by motherhood, pairbonding, pill use, breastfeeding and menstrual cycle phase.

Variable (n)	Waking T	p ^a	Evening T	p ^a
Non-mother (43)	93.1 (4.5)		47.6 (3.0)	
Mother - all (24)	63.4 (6.0)	0.0001 ^b	40.3 (4.0)	0.200 ^b
Young offspring (14)	54.7 (7.6)	0.0001 ^b	30.6 (4.9)	0.004 ^b
Older offspring (10)	75.6 (9.0)	0.096 ^b	53.8 (5.8)	0.236 ^b
Non-pairbonded (42)	90.2 (4.7)		47.7 (3.0)	
Pairbonded (25)	69.6 (6.2)	0.004	40.5 (3.9)	0.180
Non pill use (58)	87.0 (4.0)		47.8 (2.4)	
Pill user (9)	53.7 (10.2)	0.003	26.9 (6.2)	0.003
Not breastfeeding (58)	85.5 (4.2)		46.8 (2.5)	
Breastfeeding (9)	63.2 (10.6)	0.043	33.6 (6.4)	0.057
Other phase (45)	76.5 (4.7)		42.3 (2.9)	
Late follicular (22)	94.7 (6.7)	0.018	50.4 (4.2)	0.173

Mean (SEM) testosterone adjusted for time of saliva collection, usual wake time, and wake time that day.

^a p-value for significant difference in T measured at same time of day (all linear regression).

^b Compared to non-mothers

Table 3

Multiple regression models predicting waking and evening testosterone.

	Waking T	p	Evening T	p
Mother (offspring < 2 years) ^b	-33.6 (10.2)	0.002	-9.1 (6.2)	0.149
Mother (offspring 2+ years) ^b	-8.5 (10.6)	0.428	13.7 (6.6)	0.044
Contraceptive use	-14.0 (11.4)	0.226	-15.5 (7.2)	0.035
Late follicular	13.6 (7.9)	0.092	7.4 (5.0)	0.142
Model R ²		0.360		0.321

β (SE) predicting waking testosterone (pg/ml). Also adjusted for time of saliva collection, wake time and usual wake time (not shown).

^b Compared to non-mothers