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Prevalence of pre- and postpartum depression in Jamaican women

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Abstract

Background: Maternal depression during pregnancy has been studied less than depression in postpartum period. The aims of this study were to find out the prevalence of prepartum and postpartum depression and the risk factors associated in a cohort of Afro-Jamaican pregnant women in Jamaica.

Methods: The Zung self-rating depression scale instrument was administered to 73 healthy pregnant women at 28 weeks gestation and at 6 weeks postpartum for quantitative measurement of depression. Blood samples were collected at 8, 28, 35 weeks gestation and at day 1 and 6 weeks postpartum to study the thyroid status.

Results: Study demonstrated depression prevalence rates of 56% and 34% during prepartum and postpartum period, respectively. 94% women suffering depression in both periods were single. There were significant variations in both FT₃ and TT₄ concentrations which increased from week 8 to week 28 prepartum ($p < 0.05$) and then declined at the 35th week ($p < 0.05$ compared with week 28) and 1 day post delivery study ($p < 0.05$ compared with week 35). The mean values for TSH increased significantly from week 8 through week 35. The mean values at 1 day postpartum and 6 week postpartum were not significantly different from the 35 week values. For FT₃, TT₄ and TSH there were no significant between group differences in concentrations. The major determinants of postpartum depression were moderate and severe prepartum depression and change in TT₄ hormone concentrations.

Conclusion: High prevalence of depression was found during pre- and postpartum periods. Single mothers, prepartum depression and changes in TT₄ were factors found to be significantly associated with postpartum depression.

Background

Maternal depression during pregnancy has been studied less than depression in postpartum period. A recent study, however, shows that depression seems to be as common during pregnancy as after delivery [1]. Depending upon

various scales used to measure depression, the prevalence rate has been found to be between 5% and 26% for antenatal depression [2-6] and between 7% and 30% for postpartum depression [7-10]. The etiology of both pre- and postpartum depression still remains illusive. Various psy-

cho-social and endocrine factors have been connected to both pre- and postpartum depression [2,3,11-14]. In addition, postpartum depression has been linked to a variety of endocrine root causes- especially postpartum thyroid dysfunction [15-17]. Antenatal depression has also been considered to increase the risk for postnatal depression [11,3]. This paper presents our findings on the prevalence of pre- and postpartum depression and risk factors responsible in a cohort of pregnant Afro-Jamaican women.

Methods

One hundred and forty (140) healthy Afro-Jamaican pregnant women attending antenatal clinic in the Department of Obstetrics, Gynecology and Child Health, University Hospital of the West Indies, Jamaica (WI) during the period May 2000 to February 2001 consented to participate in this study. Women with a history of thyroid disease, any medical illness, depression or substance abuse were excluded. The mean age was 27 years, parity varied from primigravidae to 4 live births and the mean length of gestation was 8 weeks at booking. A relatively large number of these mothers (67) had to be excluded (45 failed to keep their antenatal appointments, 5 had miscarriages, 4 had initial thyroid dysfunction, 7 had premature deliveries, 1 developed severe hypertension during pregnancy and 5 were delivered by cesarean section). Thus a total of 73 pregnant women completed this study.

This study was duly approved by the UWI/UHWI Ethics Committee.

Depression assessment

Each mother was given a questionnaire at booking to obtain a clinical profile and relevant demographic data. The Zung self-rating depression scale (SDS) questionnaire was administered at 28 weeks gestation and at 6 weeks postpartum. A participant was considered to have no psychopathy if Zung was <50, minimal to mild depression if Zung score was 50–59, moderate to marked depression if Zung score was 60–69, and severe to extreme depression if Zung score was 70 and over. Depending on the depres-

sion status, the cohort was subdivided into four subgroups (i) women depressed during prepartum only (ii) women depressed prepartum and who continued to be depressed postpartum (iii) women depressed postpartum only and (iv) women showing no signs of depression in either period.

Hormonal profile

Blood samples were collected at 8, 28, 35 weeks of gestation and at 1 day and six weeks after childbirth. Serum total thyroxine (TT₄), free triiodothyronine (FT₃), thyrotropin (TSH) were determined using standard radioimmunoassay kits (Diagnostic Products Corporation, Los Angeles, California, USA). The sensitivity value for each assay was 0.25 pg/dl; 0.2 pg/ml; and 0.03 :IU/ml, respectively. All samples for each test were assayed in the same batch.

Statistical analysis

Data are expressed as means ± SE or counts as appropriate. The data were analyzed by repeated measures analysis of variance (RMANOVA) with the between group factor being the four subgroups (i) women depressed during prepartum only (ii) women depressed prepartum and who continued to be depressed postpartum (iii) women depressed postpartum only and (iv) women showing no signs of depression in either period and the measurements done over time (8 wks 28 wks 35 wks 1 day postpartum and 6 wks postpartum) as the repeated measures factor. In analyses where there were significant interactions between the group factor and the repeated measures factor, we compared differences between the depression categories at each experiment and differences between the means of measured variables within each clinical category at each experiment, by Tukey method. In these comparisons the error term was the root mean square error from the RMANOVA analysis with its associated degrees of freedom.

The Zung self-rating depression scale (SDS) was used to classify participants into depression categories. For this analysis these categories were treated as an ordinal scale.

Table 1: Results of Zung self-rating depression scale (SDS) questionnaire administered to 73 women at 28 weeks of gestation and at 6 weeks postpartum.

SDS Index	Equivalent Clinical Global Impression	28 weeks Gestation Number of Cases	6 weeks Postpartum Number of cases
Below 50	Normal range, no psychopathology	32 (43.8%)	48 (65.75%)
50–59	Presence of minimal to mild depression	23 (31.5%)	18 (24.66%)
60–69	Presence of moderate to marked depression	13 (17.8%)	7 (9.59%)
70 and over	Presence of severe to extreme depression	5 (6.9%)	None

Table 2: Demographic characteristics by depression groups

Variables	Never Depressed N = 25	Prepartum Depression only N = 23	Prepartum & Postpartum Depression N = 18	Postpartum Depression only N = 7
Mean Age (Yrs)	27	27	25.9	28.4
Single	14(56%)	14(61%)	17(94%)	5 (71%)
Primigravidae	13 (52%)	11 (48%)	9 (50%)	3 (43%)
Multiparous	12 (48%)	12 (52%)	9 (50%)	4 (57%)
Employed	15 (60%)	15 (65%)	8 (44%)	6 (86%)
Unemployed	10 (40%)	8 (35%)	10 (56%)	1 (14%)

To assess effects of changing thyroid hormones on the odds of changing depression categories postpartum we used only the measurements done at 28 wk prepartum and 6 weeks postpartum. We assessed the odds of being in a particular depressed category in the postpartum period for participants using an ordinal logistic model adjusting for prepartum depression category and differences in thyroid hormones level (6 wk postpartum – wk 28 prepartum values) as covariate. Inferential tests were considered statistically significant if $p < 0.05$ (two tail).

Data analysis was performed using Stata version 8 for Windows (Statacorp, College station, TX).

Results

Pre- and postpartum depression

Self-rating depression scale administered to 73 women indicated that 41 women (56.16%) were having depression at 28 weeks prepartum. Out of these 41, 23 (31.5%) had mild, 13 (17.8%) had moderate and 5 (6.9%) had severe depression. Out of these 41, only 18 (24.66%) women were found to be depressed at 6 weeks postpartum. Remaining 23 (31.5%) women recovered anywhere between 28 weeks prepartum and 6 weeks postpartum period. The data further indicated that 4 out of the 5 women who were severely depressed at 28 weeks prepartum were only moderately depressed at 6 weeks postpartum; and the remaining had no depression. In addition, 7 (9.59%) different women were found to be depressed only at 6 weeks postpartum. Therefore, a total of 25 (34.25%) women suffered mild to marked depression during the postpartum period (Table 1).

Demography and depression

Analysis of the data on marital status demonstrated that 31 (75.6%) out of 41 women who were depressed at 28 weeks prepartum were single or legally not married. 17 (94.4%) out of 18 women who continued to have depression at 6 weeks postpartum were also single. Similarly, 5 (71.4%) out of 7 women who suffered depression in postpartum period only, were single. Age, parity, miscarriages

and employment status was not associated with depression (Table 2).

Thyroid status and depression

There were significant changes in the mean thyroid hormones concentrations over time (Fig 1). For both FT_3 and TT_4 , mean values increased from week 8 to week 28 prepartum ($p < 0.05$) and then declined at the 35th week ($p < 0.05$ compared with week 28) and 1 day post delivery study ($p < 0.05$ compared with week 35). There was a small increase in the mean FT_3 values at 6 weeks postpartum compared to the 1 day postpartum values but this was not statistically significant. In contrast, the mean values for TT_4 decreased further compared to the 1 day postpartum values ($p < 0.05$). The mean values for TSH increased significantly from week 8 through week 35. The mean values at 1 day postpartum and 6 week postpartum were not significantly different from the 35 week values. For FT_3 , TT_4 and TSH there were no significant between group differences in concentrations. However there was a significant group by study interaction for FT_3 . At weeks 8 and 28 prepartum, the women who were never depressed tended to have higher mean FT_3 values compared to women who were depressed. At week 35, the mean values for FT_3 of women who were never depressed were significantly lower ($p < 0.05$) than mean FT_3 values for women who were depressed prepartum.

There were no significant relationships between thyroid hormone concentrations and Zung scores at 28 weeks or at 6 week post partum. The major determinants of postpartum depression were moderate and severe prepartum depression and change in TT_4 hormone concentrations (Table 3). Thus the odds of being depressed postpartum increased by factors of 4.8 in the moderately depressed category and 2.1 in the severely depressed category relative to not being depressed respectively adjusting for changes in TT_4 . For every unit increase in the magnitude of the difference in mean TT_4 values between week 28 prepartum and 6 week postpartum the odds of being more depressed increases by a factor of 0.75 ($p < 0.003$).

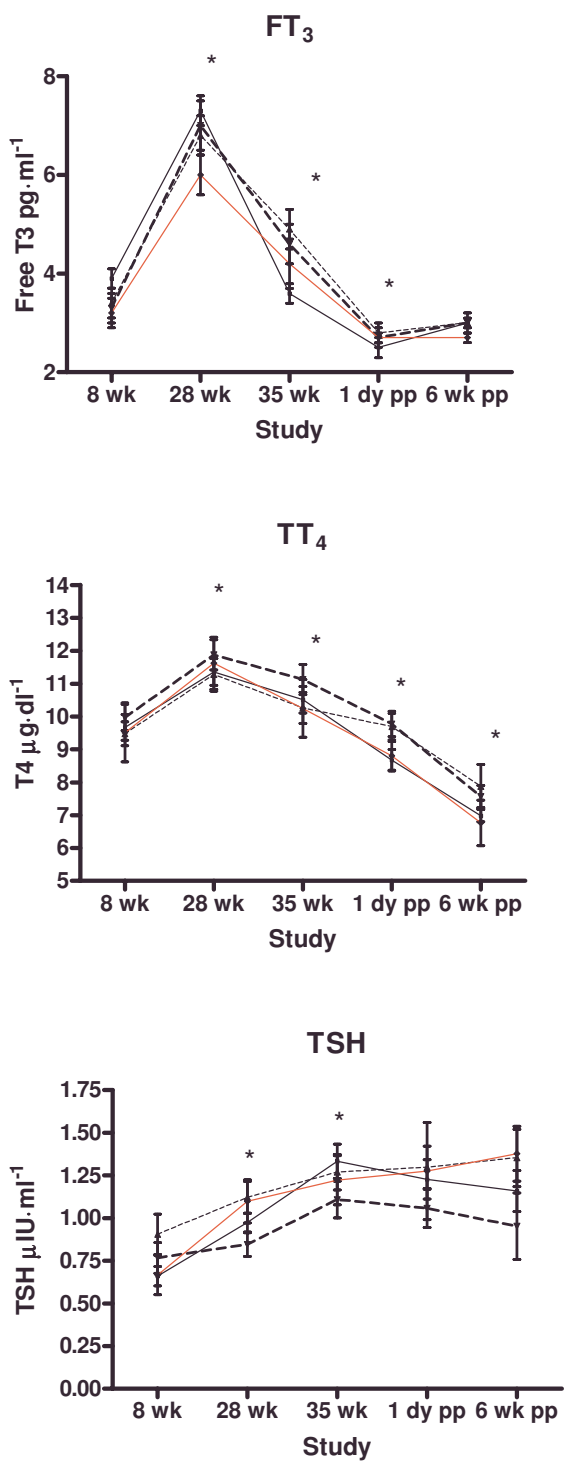


Figure 1
Thyroid hormones by depression groups. * Significantly different from previous values $p < 0.05$.

Discussion

The study demonstrated a higher prevalence rate of 56% maternal depression at 28 weeks prepartum as compared to 34 % at 6 weeks postpartum confirming that depression during pregnancy is as common as after delivery [1]. Our prevalence rates for both pre- and postpartum depression are higher than the rates reported earlier [1-3,7,8,10]. This may be attributed to the fact that greater percentages of women during both periods suffered from minimal to mild depression only, a score that might have been ignored by previous authors, who may have regarded it as clinically non-significant depression. Further analysis of the data on postpartum depression showed that prepartum depression was the major risk factor for postpartum depression [3,18].

Though the postnatal depression was observed only at 6 weeks postpartum, it is possible that the onset could have been within the first week after the delivery as reported earlier [9].

This study revealed that single mothers are more prone to depression both during pre- and postpartum periods. This can be interpreted in two ways (i) either that being single implies lack of a stable supporting union which influences depression and/or (ii) there is a predominance of single mothers in Jamaica (50 of the 73 mothers in this study were single). In both circumstances, whatever the cause of depression, the majority of subjects would be single. Employment, parity and the previous miscarriages did not seem to have any effect in producing depression.

In the sub-group of women representing 9.6% of the cohort, who developed depression in the postpartum period, relative hypothyroidism was observed during the late gestation and early postpartum periods. In addition, changes in mean TT₄ levels were significantly related to postpartum depression. This finding is supported by previous studies [15-17] that suggest that postpartum thyroid dysfunction may be responsible for postpartum depression. However, this is in contrast to the study by Lucas et al [19] that has reported no link between postpartum thyroid dysfunction and postpartum depression.

Conclusion

The results of our study limited only to corporate area, suggested almost equally high incidence of depression in single Jamaican mothers both during pre- and postpartum periods. In addition, relative hypothyroidism developed between late gestation and postpartum period could have been responsible for postpartum depression in a sub-group of mothers. In the light of the results, it is suggested that women who develop depression during pregnancy should be monitored for thyroid functions and social support be provided to single mothers to avoid the risk of

Table 3: Prepartum determinants of a more severe depression post partum category relative to less severe depression postpartum category

Prepartum Depression Category	Factor change in odds*	p value
Mild	1.7	ns
Moderate	4.8	<0.002
Severe	221	<0.001
Difference in T4 values	0.75	<0.003

* Relative to not depressed category

postpartum depression. A wider cross-sectional study in Jamaica is further needed to confirm these results.

Competing interests

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

Authors' contributions

JW M.Phil. Student who carried out this work.

OP chief supervisor, conceived and contributed to study design, reviewed the statistical analysis, interpretation of results and preparation of manuscript.

SK provided access to the subjects in the hospital, supervisory committee member, provided input on pregnancy matters.

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References

- Evans J, Heron J, Francomb H, Oke S, Golding J: Cohort study of depressed mood during pregnancy and after child birth. *BMJ* 2001, **323**(7307):257-260.
- Kitamura T, Sugawara M, Sugawara K, Toda MA, Shima S: **Psychosocial study of depression in early pregnancy.** *Br J Psychiatry* 1996, **168**(6):732-738.
- Johanson R, Chapman G, Murray D, Johnson I, Cox J: **The North Staffordshire Maternity Hospital prospective study of pregnancy associated depression.** *J Psychosom Obstet Gynaecol* 2000, **21**(2):93-97.
- Pajulo M, Savonlahti E, Sourander A, Helenius H, Piha J: **Antenatal depression, substance dependency and social support.** *J Affect Disord* 2001, **65**(1):9-17.
- Luoma I, Tamminen T, Kaukonen P, Laippala P, Purra K, Salmelin R, et al.: **Longitudinal study of maternal depressive symptoms and child well-being.** *J Am Acad Child Adolesc Psychiatry* 2001, **40**(12):1367-1374.
- Moses-Kolko EL, Roth EK: **Antepartum and postpartum depression: healthy mom, healthy baby.** *J Am Med Womens Assoc* 2004, **59**(3):181-191.
- Da Costa D, Larouche J, Drista M, Brender W: **Psychosocial correlates of prepartum and postpartum depressed mood.** *J Affect Disord* 2000, **59**(1):31-40.
- Morris-Rush JK, Freda MC, Bernstein PS: **Screening for postpartum depression in an inner-city population.** *Am J Obstet Gynecol* 2003, **188**(5):1217-1219.
- Yamashita H, Yoshida K: **Screening and intervention for depressive mothers of new-born infants.** *Seishin Shinkeigaku Zasshi* 2003, **105**(9):1129-1135.
- Bloch M, Daly RC, Rubinow DR: **Endocrine factors in the etiology of postpartum depression.** *Compr Psychiatry* 2003, **44**(3):234-246.
- Areias ME, Kumar R, Barros H, Figueiredo E: **Correlates of postnatal depression in mothers and fathers.** *Br J Psychiatry* 1996, **69**(1):36-41.
- Glasser S, Barell V, Boyko V, Ziv A, Lusky A, Shoham A, et al.: **Postpartum depression in an Israeli cohort; demographic, psychosocial and medical risk factors.** *J Psychosom Obstet Gynaecol* 2000, **21**(2):99-108.
- McCoy SJ, Beal JM, Watson GH: **Endocrine and postpartum depression. A selected review.** *J Reprod Med* 2003, **48**(6):402-408.
- Hendrick V, Altshuler LL, Suri R: **Hormonal changes in the postpartum and implications for postpartum depression.** *Psychosomatics* 1998, **39**(2):93-101.
- Amino N, Tada h, Hidaka Y: **Screening for postpartum thyroid dysfunction in the general population is beneficial.** *J clin Endocrinol Metab* 1999, **84**(6):1813-1816.
- Kent GN, Stuckey BGA, Allen JR, Lambert T, Gee V: **Postpartum thyroid dysfunction: clinical assessment and relationship to psychiatric morbidity.** *Clin Endocrinol* 1999, **51**(4):429-438.
- Barca MF, Knobel M, Tomimiri E, Cardia MS, Medeiros-Neto G: **Prevalence and characteristics of postpartum thyroid dysfunction in Sao Paulo Brazil.** *Clin Endocrinol* 2000, **53**(1):21-31.
- Heron J, O'Connor TG, Evans J, Golding J, Glover V, The ALSPAC Study Team: **The course of anxiety and depression through pregnancy and the postpartum in a community sample.** *J Affect Disord* 2004, **80**(1):65-73.
- Lucas A, Pizarro E, Granada ML, Salinas I, Sanmarti A: **Postpartum thyroid dysfunction and postpartum depression: are they two linked disorders.** *Clin Endocrinol* 2001, **55**(6):809-814.

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