

**A STUDY OF PREMENSTRUAL
DYSPHORIC DISORDER (PMDD) IN A
SAMPLE OF EGYPTIAN WOMEN**

*A.H. Khalil **, *Seif el Dawla, A.E **, *Zarif, N.M. **
*M.H. Mostafa ***, *K.M. Ismail *** and *Abou Seeda, M.R. ***

*** Department of Psychiatry, Ain Shams University**

**** Department of Obstetrics and Gynaecology, Ain Shams Univer**

A STUDY OF PREMENSTRUAL DYSPHORIC DISORDER (PMDD) IN A SAMPLE OF EGYPTIAN WOMEN

*A.H. Khalil **, *Seif el Dawla, A.E **, *Zarif, N.M. **

*M.H. Mostafa ***, *K.M. Ismail *** and *Abou Seeda, M.R. ***

* Department of Psychiatry, Ain Shams University

** Department of Obstetrics and Gynaecology, Ain Shams Univer

ABSTRACT

Up to 75% of women report some preme symptoms, but less than 10% have symptoms enough to qualify for a diagnosis of PMDD. to diagnosis is establishing a pattern of PMDD symptoms that recur during the late phase of the menstrual cycle and remit after r Underlying psychiatric and medical disorde might mimic PMDD should be ruled out dressed. The aim of this study was to study th alence of PMDD in a random sample of E; women, to explore the symptom profile of thi der in that sample and to search for any poss lationship between PMD regarding occurren or severity with different personality and demographic factors. 987 females in the chi ing period, their age between 15-45 years; them have regular menses were recruited by a ified random sample; the whole sample was sc for premenstrual symptomatology using Moo strual Distress Questionnaire (MDQ). Bc PMDD and the control groups were assessed following tools :

1) Eysenck personality questionnaire (1964) assessment of different personality var

2) Taylor Manifest Anxiety Scale (Janet

1953) ; for assessment of anxiety traits.

3) Beck Depression Inventory (BDI) (Beck, 1961) : for detection of depression states.

4) Guilford Scale (Guilford and Zimmerman) for assessment of depressive traits.

5) Fahmy and El Sherbini Socio-economic Sheet of social status (1988). PMDD represented 6.69% of the sample examined in the study, and it was found to be related to work, troublesome home atmosphere, menorrhagia, dysmenorrhoea, neuroticism, criminality, anxiety, and depressive traits and state. Depressive state and neuroticism were the primary determinants of the development of the disease.

Introduction

Menstrual syndrome (PMS) is a condition characterized by a cyclic occurrence of physical, affective and behavioural symptoms in the 7-10 days before the menstruation and ending with its onset. This is a common condition affecting up to 80% of females in the child bearing age (Facchinetti et al. 1998).

On the other hand, Premenstrual Dysphoric Disorder (PMDD), which was first established to have strict diagnostic criteria in the DSM-IV (El-Metwally, 2002 and Muzina, 1998), is a condition severe enough to interfere with the women's social, occupational and marital life. It is much less common, affecting about 2-5% of females in the child bearing age (Steinberg et al, 1999, Gehal, 1999, Young et al, 1998,

and Rubinow et al, 1998).

Despite the marked suffering faced by women who have this disorder, they used to consider this as an accepted part of their life. Many of them, even those who are aware of the size of their problem did not ever think to consult for treatment of such a condition. The cost of this illness is paid on the expense of the patient herself, her family, her work and thereby the community as a whole.

In Egypt, women are used to endure different stressors and don't complain especially if the problem is related to psychiatric symptoms for fear of being stigmatized with having mental illness and so prefer to present with physical symptoms and consult a physician rather than a psychiatrist, and this may give an

example of how culture can affect the presentation of a disease.

The aim of this study was to study the prevalence of PMDD in a random sample of Egyptian women, to explore the symptom profile of this disorder in that sample and to search for any possible relationship between PMD regarding occurrence and/or severity with different personality and socio-demographic factors.

Subjects and methods :

The work was preceded by a pilot study, to determine the size of the sample needed, and to assess the ability of the subjects to understand the used questionnaires.

The sample consisted of 987 females in the childbearing period, their age between 15-45 years; all of them have regular menses.

Any woman with a history of medical and or gynaecological diseases or a history of any hormonal therapy including contraceptive pills or steroid therapy and diuretic therapy were excluded from the study.

The samples were recruited by a stratified random sample from :

1) medical students in the 2nd, 4th, and 6th years of the faculty of medicine.

2) High nursery school students

in the 1st and 4th years.

3) Secondary nursery school dents 3rd year.

4) House officers.

5) Employees, both professi and non professional, from Shams University hospitals as as the Faculty of medicine.

6) Illiterate house wife's from atives of children attending paediatric outpatient clinic.

All the women fulfilled above mentioned inclusion cri and were accepting to participat the study after signing the stand consent.

A personal data collection s was filled at first, and included the demographic, menstrual, obriic, medical, and family history. sheet covered all the data needed assessment of the socio-econo standard according to Fahmy El-Sherbini (1988).

The whole sample was scree for premenstrual symptomatol using Moos Menstrual Dist Questionnaire (MDQ) which originally created by Moos (19 to assess symptom changes in ntion to menstruation. The Ar version of the MDQ which was veloped by Farahat (1988) thro translation/retranslation process the one used in this study.

According to the results, women scored 125 or above were considered to be a case in need for further confirmation; and those who scored less than 125 were considered controls.

To confirm this finding, DSM-IV search criteria were used. This included a prospective daily rating of symptoms for two consecutive cycles and was guided by the calendar remenstrual experiences-COPE protocol (Lewinsohn et al, 1990).

For the diagnosis of PMDD there should be at least a 30% increase in symptom scores from the follicular phase to the luteal phase (Ekholm et al. 1993). To warrant pharmacological treatment, worsening should increase by at least 50% (Steiner et al. 1995).

Those who failed to be classified as PMDD and fulfilled the criteria for PMS were separated as a separate category.

The Arabic version used in the present study was developed through a translation-retranslation process. The inter-rater reliability was established during the pilot study ($K=0.88 - 0.93$).

Both the PMDD and the control groups were assessed by the following tools:

1) Eysenck personality questionnaire (EPQ) (1964) assessment of

different personality variables.

2) Taylor Manifest Anxiety Scale (Janet Taylor, 1953) : for assessment of anxiety traits.

3) Beck Depression Inventory (BDI) (Beck, 1961) : for detection of depression states.

4) Guilford Scale (Guilford and Zimmerman) for assessment of depressive traits.

5) Fahmy and El Sherbini Socio-economic Sheet of social status (1988). The findings were analysed using the statistical package for the social sciences (SPSS).

RESULTS

The total sample included in this work involved 987 women, out of them, 514 women (52.08%) did not have any significant premenstrual symptoms and were considered to be the control group.

65 women (6.59%) fulfilled the criteria of PMDD and were considered as the patient group.

296 women (29.98%) fulfilled the criteria of PMS.

36 women (3.65%) had severe premenstrual symptoms which turned out to be a premenstrual exacerbation of a present psychiatric disorder, and were excluded from the study.

76 women (7.7%) dropped out, and also were excluded from the study.

Descriptive and demographic data :

There was no significant difference ($P = 0.882$, $U = 16520$ Mann-Whitney test) age with the mean age (26.6 ± 7.1 years) for the patient group and the mean age (26.8 ± 7.7 years) for the control group.

The mean age of onset of PMDD was 19.4 ± 5.2 years.

Also, there was no significant difference between both patient and control groups regarding the weight, height, degree of education, marital status.

According to Fahmy and El Sherbini scale (1988), women were clas-

sified into four social classes although the prevalence of the disease PMDD was higher in the lower social class, yet, there was no significant difference between both groups.

On the other hand, a statistically significant difference was found between both groups, regarding the incidence of PMDD, with a higher incidence of PMDD among the working group. ($P = 0.0001$, $U = 68.04$ Mann-Whitney test).

Family background :

18 women with PMDD (27%) were suffering from a troubled home atmosphere, in comparison with 53 women (10.31%) in the control group. The difference between both groups was found to be highly significant ($P < 0.0001$) (1) :

Home atmosphere	PMDD	Control	Chi-square	P
Calm	47 (72.31%)	461 (89.69%)		
troublesome	18 (27.69%)	53 (10.31%)	16.2	<0.0001

Mann-Whitney Test.

7.69% of the patient group had a positive past history of psychiatric illness in the patient group in comparison to 4.67% in the control group, and the difference was not statistically significant ($P = 0.2924$).

Menstrual History :

Both menorrhagia and dysmenorrhoea were significantly higher in the PMDD group. While, 19 women (29.23%) with PMDD suffered

rhagia only 69 (13.42%)
 1 in the control group suffered
 hat complaint, and the differ-
 as statistically significant (P=

women (55.38%) with PMDD
 positive history of dysmenor-
 compared to 212 (41.25%)
 a in the control group (P =

tric history :

ere was no statistically signifi-
 difference between both the
 e and the control groups re-
 g the parity, mode of deliv-

the other hand, both infertility
 abortions were found to be
 in the patient group, howev-
 difference was not statistical-

ly significant (P = 0.1099. and P =
 0.1846 respectively).

Symptom profile :

Out of the eight symptom clus-
 ters created by Moos, we found that
 negative affect was the most com-
 monly represented (29.63%) fol-
 lowed by pain score (25.93%) and
 then the behavioural changes
 (14.81%), table (2).

Within the negative affect clus-
 ter, irritability and mood swings
 (77.78% for each) were the com-
 monest symptoms, and within the
 pain cluster, fatigue was the com-
 monest symptom (72.22) followed
 by generalised aches (70.37%)
 while, within the behavioural chang-
 es cluster, preference of stay at
 home was the commonest symptom
 (46.3%).

Symptoms in Moos Questionnaire	Percentage of severity
negative affect	29.63%
score	25.93%
vioural changes	14.81
onomic reactions	7.41%
r retention	3.7%
entration score	1.85
sal	0%
rol	0%

the commonest symptoms in the
 clusters were as following :

east tenderness (48.15 %) the
 onest symptom in the water

retention cluster, Suffocation and
 palpitation (33.33%) were the com-
 monest symptoms in the control
 cluster, nausea and vomiting
 (33.33%) were the commonest

symptoms in the autonomic cluster, difficulty in concentration (38.89%) was the commonest symptom in the concentration cluster, and being affectionate (77.78%) was the commonest symptom in the arousal cluster.

Psychometric assessment :

A) Personality traits as assessed by the EPQ. out of the five subscales assessed by the EPQ only neuroticism and criminality showed a highly significant difference between the patient and control groups, being higher in the PMDD group ($P = 0.0001$ and 0.0342 respectively).

B) Anxiety symptoms as measured by Taylor Manifest Anxiety

Scale : the level of anxiety found to be higher in the PMDD group with a highly significant difference from the control group ($P = 0.0001$).

C) Depressive traits as measured by Guilford Scale : there was a highly significant difference between both groups with more depressive traits in the PMDD group.

D) Depressive state as assessed by the Beck inventory : there was a highly significant difference between both groups with more depressive symptoms in the PMDD group.

Session	PMDD	Control	Chi square	P
Normal	9 (13.84%)	193 (37.55%)		
Subnormal	55 (84.62%)	302 (58.75%)		
Very normal	1 (1.54%)	19 (3.7%)	16.33	<i>0.0003</i>

Session	PMDD	Control	Chi square	P
Normal	7 (10.77%)	201 (39.11%)		
Subnormal	18 (27.69%)	182 (35.41%)		
Intermediate	27 (41.54%)	114 (22.18%)		
Very normal	13 (20%)	17 (3.3%)	53.8	<i><0.0001</i>

Session	PMDD	Controls	Chi square	P
Normal	7 (10.77%)	127 (24.76%)		
Subnormal	16 (24.62%)	189 (36.84%)		
Intermediate	27 (41.45%)	175 (34.11%)		
Very normal	15 (23.07%)	22 (4.29%)	40.02	<i><0.0001</i>

Session	PMDD	Controls	Chi square	P
Normal	27 (41.54%)	379 (74.16%)		
Subnormal	38 (58.46%)	103 (20.16%)	41.19	<i><0.0001</i>

Session	PMDD	Controls	Chi square	P
Normal	21 (32.31%)	251 (49.12%)		
Subnormal	44 (67.69%)	259 (50.68%)	6.749	<i>0.0342</i>

Using the Logistic regression for analysis, it was found that the Beck Score (depressive), and the EPQ-N (neuroti-

cism) score were the primary determinants of the disease process as shown in the following table :

	Parameter estimate	Standard error	Pr > chi square
Beck score	0.0542	0.0165	0.001
EPQ - N	0.2006	0.0467	0.0001

Meaning that the Beck Score and the EPQ-N together are the main indicators for the development of PMDD and the probability to have the disease depends on these two variables.

Using multiple regression analysis, there was no correlation between the home atmosphere, work, or marital status and the severity of PMDD.

Also, there was no significant correlation between Beck (depressive states), Guilford (depressive traits) and Taylor (anxiety symptoms) and the severity of PMDD.

Regarding multiple correlation analysis for all the EPQ (personality traits) subscales, in relation to the severity of the disease (Moos Q), there was a statistically significant negative correlation between EPQ-L (Lie score) and the severity of PMDD ($r = -0.31663$ $P = 0.0102$).

DISCUSSION

Terms like PMS and PMDD have been used interchangeably, while they do not refer to the same finding (Vanselow 1998). Also Rapkin (1992) noticed that the term PMS and LLPDD (late luteal phase Dysphoric disorder) have been used

interchangeably in the literature. The importance of this point is most of the talks in the literature about PMS, and this in spite of the fact that most of the reviewed articles were originally about PMDD.

In Egypt, females in the child bearing period (15 - 45) consist about 13.456.404 women according to the final results of the 1996 census and according to the statistical year book (central agency for population mobilization and statistics, 1996) and since the prevalence estimate of PMDD reaches up to 10% of females in the child bearing period, so by approaching PMDD we are dealing with a condition that affects about 1.345.640 women in their period of maximal productivity and to efficiency. Those women are expected to suffer about 1400 - 2800 days through out their life (Yon 1997) providing that the estimated age of onset is around 26 years offset at the age of menopause (35 years). This constitutes a gross burden on both social and economic levels, both directly and indirectly, which represents an enormous economic suffering in a developing country like Egypt.

We did not find any correlation

teen age and severity of PMDD ($r = 0.0931$ and $P > 0.1$). Our findings coincide with Farahat (1988); Khalil, (1986) and Halbreich et al, (1992) who found no correlation between age and severity of the symptoms. On the other hand, Freeman et al (1995, 1990), Romacharam et al (1992) reported that symptom severity is inversely related to age. A possible explanation to our finding is that the young age groups of women may suffer from late age of marriage and its social consequences which might render them more liable to PMDD to a level comparable to that of the old age groups. Other possibility might be due to the lower mean age in our study, (26.6 ± 7.1 years) as compared to (33.1 ± 5.3) in Freeman study.

Although no statistically significant difference was found between the PMDD and the Control group regarding their socioeconomic status, yet, it can be noticed that there is more representation of social class I in the PMDD (35.4%), more representation of social class IV in the control group (55%), indicating a tendency towards more occurrence of the disorder in the higher social class. Reid (1985), explained this finding by the fact of greater accessibility of medical care among those individuals, furthermore, in the lower social classes, women are preoccupied mainly with their essential basic needs for life more than their psychological welfare.

Higher incidence of PMDD was reported in the working group of women, this finding was observed by other researchers like: Lee and Rittenhouse (1991), Schnurr (1988) who found that there is increased likelihood of PMS in women working outside the home, which could be explained by the presence of multiple stressors at work including multiple interpersonal relations. Andersch (1986), found that about 10% of women are absent from work on at least two occasions each year because of PMS.

The relationship and family dimensions of PMS have received inadequate attention in the literature both in terms of aetiology and management (Condon, 1996). Most of the women presented with premenstrual complaints do so because a social problem has reached a point of crises, and in many instances premenstrual complaints disguise unhappy family relationships (Vanseelow, 1998). This was matched with our findings of the more occurrence of PMDD in females with troublesome home atmosphere.

More women with PMDD were found to suffer from menorrhagia (Chi square = 11.4, $P = 0.0034$). The same finding was reported previously by Bancroft et al (1995), who reported that premenstrual depressive mood changes were noticeably worse and more prolonged in women with menorrhagia.

In a country like Egypt, this may be explained by the fact that women are brought up to believe that menstruation is something that should be hidden and not even to be talked about especially in the presence of men due to shyness, so, it would be a critical situation for a female if she could not secure her appearance and privacy in case of excessive menstrual flow as occurs in menorrhagia. This is especially important for working women and students, who are exposed to outdoor interactions. This may lead women to avoid such situations by absence from work or school, and this may represent an unwanted stressor that females may react toward it by anxiety and depression.

Although PMS and dysmenorrhoea may coexist, the variable nature of this association suggests that different pathophysiologic mechanisms are involved (Moos, 1969). In our study we have found a statistically significant difference between both patients and controls regarding the presence of dysmenorrhoea, being more in the PMDD group. Bancroft et al (1993) noticed that premenstrual depressive changes were noticeably worse and more prolonged in women with severe dysmenorrhoea. A possible explanation of such association is that the expectation of severe pain interfering with normal life might increase the negative attitude towards menstruation, i.e. act like a stressor, and women may react to it by the Dysphoric

symptoms. Women with PMS apparently achieve pregnancy readily hence, there is no association with infertility (Reid, 1985), in our study we found higher percentage of infertile women in the PMDD group, the difference was not statistically significant (Chi square = 2.555, $P = 0.1099$). This may be due to small number of infertile women in the both groups : 4 in the patient group and 14 in the control group.

Rohde et al (1992) compared frequency and type of premenstrual symptoms between a sample of fertile women and normal controls and found that 52% of the infertile women and 42% of the control group fulfilled the criteria for PMDD. Infertile women reported more often mood lability, irritability, anxiety/tension and depression of interest.

One of the aims of this study was to define the symptom profile of PMDD in the Egyptian women and to compare it with the results of other different studies. This may enable us to understand how the culture may affect the presentation of this syndrome.

Out of the 8 clusters of the questionnaire, the negative affect was the most commonly represented one in our patients, followed by pain score and the behavioral changes. The findings of presence of the negative premenstrual affect includes: anxiety, irritability,

d swings, depression, and ten-
 arahat, 1988, Parlee, 1974, and
 s, 1968 shared our study the
 of the same questionnaire
 Q). Of the three studies the
 nian one had the highest score,
 h seems to be due to different
 e of stressful stimuli affecting
 les in our society. These stress-
 timuli play a definite role in the
 women experience the period
 eding menstruation (Kinch and
 lberg, 1984). Also, the scores
 ded in our study was much
 er than other studies (mean

score of the negative affect scale
 was 35.65 ± 7.46 in comparison to
 22.1 ± 6.9 in Farahat study; 17 ± 8.1
 in Moos study and 18.3 ± 7.4 in
 Parlee study) this can be explained
 by the fact that these three studies
 were dealing with PMS, which is
 different in nature from PMDD as it
 is presenting primarily by mood
 symptoms, so eliciting high scores
 in the negative cluster of symptoms.

The frequency of each one of the
 symptoms within the cluster is vari-
 able in the different studies as fol-
 lows :

Authors	Commonest symptoms recorded
Present work	Irritability and mood swings followed by restlessness, depression, anxiety.
Winklers (1997a)	Anxiety, tension and nervousness (among women with PMS as well as PMDD).
Widom (1996)	Irritability (the single most common symptom in PMS sufferers)
Winklers et al, (1993)	Irritability followed by depressed mood (sample of women with PMS and PMDD)
Winklers et al, (1992) (A IV mood disorder work group)	Depressed mood, mood swings, anxiety / tension.
Winklers (1983)	Depressed mood, anxiety and sleep disturbances.
Winklers et al. (1982)	Severe tension, irritability, and anxiety.

Personality variables :

When with the use of the same
 for assessment of personality
 variables, there is an evident dis-
 crepancy in the results of different
 studies dealing with the effect of

personality on PMS (Mortola,
 1992). In the present work, out of
 the 5 subscales assessed by the
 EPQ, only two items showed a sta-
 tistically significant difference be-
 tween the cases and controls. First is
 neuroticism, which was found to be

significantly higher in the cases (Chi-square = 41.19; $P < 0.0001$) than in the controls. This is in agreement with most of the studies which used EPQ for assessment of personality variables such as Ekholm et al, 1998, Ussher et al, 1992, Hallman et al, 1987, and Mira et al, 1985.

Also, these results agree with other studies that used other tools for personality assessment like Freeman et al, 1995 who used tridimensional personality questionnaire (TPQ), and found a modest but significant elevation in the TPQ factors primarily harm avoidance which is broadly similar but not identical to other measures of neuroticism (Wetzel et al, 1992). Also, Parry et al, 1996, used Millon Clinical Multiaxial Inventory (MCMI) and found higher levels of certain variables that lie under the heading of neuroticism in the EPQ, like, irritability with erratic moodiness, recurring mood shifts and anxiety.

Significant difference between cases and controls regarding criminality was found in this work ($P = 0.3420$). This is in agreement with

Parry et al, 1996, who used MCMI and found higher impulsivity destructibility under the heading hypomania on the one hand self-derogatory and suicide thoughts under borderline-cyclothymia category on the other.

Freeman et al, 1995 concluded that PMS and personality are independent of each other but the personality influence the symptom expression of PMS; that the presence of PMS in a person with high harm avoidance is likely to manifest itself more in terms of depressed mood and somatic complaints whereas in a person with high novelty seeking, PMS is more likely to manifest itself in terms of mood swings and food cravings.

Co-morbidity of PMDD with mood disorders :

Yonkers (1997) summarised studies that reported the lifetime co-morbidity rate in their patients with PMDD and a history of mood disorders with rates ranging from 30% to 70% was found as shown in the following table :

Study	No. of patients	% Mood disorders
Freeman et al. 1995	26	58
Fava et al. 1992	32	34
Harrison et al. 1989	56	70
Pearlstein and Stone, 1994	78	46
Yonkers et al. 1996	243	33

is important to put into consideration that Rubino et al (1984) found that the life time prevalence of depressive illness in the general population is about 25% with a peak incidence in the mid 30s.

Our study in agreement with previous studies showed a highly significant difference between patient and control groups regarding the presence of depressive symptoms measured by Beck Scale for depressive state), being much more in the patient group, also, a similar result was found regarding the presence of depressive traits (as measured by the Guilford scale for depressive traits).

No significant difference between the patient and control groups regarding past history of psychiatric disorders was found in this study, a finding that can be explained by the fact that, the mean age in the patient group was 26 ± 7.1 years, which is well below the peak age for the risk of affective disorders.

Co-morbidity of PMDD with anxiety :

In the present work, a highly significant difference between patients and control groups regarding the level of anxiety symptoms as measured by the Taylor anxiety scale for anxiety state, being higher in the patients group. This is consistent with most of the studies that searched the relation between PMDD and anxiety (Fava et al, 1992, Mikhail, 1986, Stout et al, 1986 and Le Melleo et al, 1995).

Conclusion :

PMDD represented 6.69% of the sample examined in the study, and it was found to be related to work, troublesome home atmosphere, menorrhagia, dysmenorrhoea, neuroticism, criminality, anxiety, and depressive traits and state. Depressive state and neuroticism were the primary determinants of the development of the disease.

REFERENCES

1. Elliott, H (2002) : Premenstrual dysphoric disorder. A guide for treating clinician. *N C Med J*, 63 (2) : 72-5.
2. Facchinetti, F; Tarabusi, M and Nappi, G. (1998) : Premenstrual syndrome and Anxiety Disorders : A Psychobiological Link. *Psych Psychosom*, 67 : 57-60.
3. Muzina, K.S. (1998) : Commonly asked questions about premenstrual dysphoric disorder. *Cleveland Clinic Journal of Medicine*, 65 (3) : 9.
4. Steinberg, S.; Annable, L.; Young's, N. and Liyanage, N. (1999) : A placebo-controlled clinical trial of L-tryptophan in premenstrual dysphoric disorder. *Biol Psychiatry*, 45 : 313-320.
5. Gehlert, S.; Chang, C. and Hartlage, S. (1999) : Symptom pattern of premenstrual dysphoric disorder as defined in the Diagnostic and Statistical Manual of Mental Disorders-IV. *Journal of Women's Health*, 8 : 75-85.
6. Young, S.A.; Hurt, P.H.; Benedek, D.M. and Howard, R.S. (1999) : Treatment of premenstrual dysphoric disorder with sertraline during the luteal phase : A randomized, double-blind, placebo-controlled crossover trial. *J Clin Psychiatry*, February, 59 (2) : 76-80.
7. Rubinow, D.R. Schmidt, P.J. and Roca, C.A. (1998) : Hormone surges in reproductive endocrine-related mood disorders : Diagnostic issues. *Psychopharmacology Bulletin*, 34 (3) : 289-295.
8. Moos, R.H. (1968) : The development of a menstrual distress questionnaire. *Psychosom Med*, 30 (6) : 853-867.
9. Farahat, M.M. (1988) : Epidemiology of premenstrual syndrome in rural and urban Egyptians, Master Thesis. Ain Shams University.
10. Mortola J.F ;Girton, L. Beck, L. and Yen, S.S.C. (1990) : Diagnostic criteria for premenstrual syndrome by a simple, prospective, and reliable instrument : The Calendar of Premenstrual Experiences. *Obstet Gynecol*, 75 : 302-7.

Ekholm,U.B.; Ekholm,N.O. and Backstrom, T. (1998) : Premenstrual Syndrome : Comparison between different methods to diagnose cyclic-ity using daily symptom ratings. *Acta Obstet Gynecol Scand*, 77 : 551-557.

Steiner, M. Steinberg, S. Stewart, D. et al. (1995) : Fluoxetine in the treatment of premenstrual dysphoria. Canadian Fluoxetine Premenstrual dysphoria Collaborative Study Group. *N Engl J Med*, 332 : 1529-1534.

Vanselow, W. (1998) : A comprehensive approach to premenstrual complaints. *Australian Family Physician*, 27 (5) : 354-361.

Rapkin, A.J. (1992) : The role of serotonin in premenstrual syndrome. *Clin Obstet Gynecol*, 35 : 629-36.

Yonkers, K.A. (1997) : Anxiety symptoms and anxiety disorders : How are they related to premenstrual disorders *Clin Psychiatry*, 58 (3) : 62-67.

Mikhail, N.R. (1986) : The problem of premenstrual syndrome in general practice in El-Salam district, Ismalia. (With special consideration to psychological factors). Ain Shams University, Master thesis.

Halbreich, U., Endicott, J. and Sachacht, S. (1982) : The diversity of premenstrual changes as reflected in the premenstrual assessment form. *Acta Psychiatr Scand*, 65 : 46-65.

Freeman, E.; Rickels, K.; Sondheimer, S.J. and Polansky, M. (1990) : Ineffectiveness of progesterone suppository treatment for premenstrual syndrome. *JAMA*, 264 : 349-353.

Freeman, E.W.; Rickels, K.; Sondheimer, S.J. and Polansky M. (1995) : A double-blind trial of oral progesterone, alprazolam, and placebo in treatment of severe premenstrual syndrome. *JAMA*, 274 : 51-57.

Ramcharan, S.; Love, E.J.; Fick, G.H. et al. (1992) : The epidemiology of premenstrual symptoms in a population-based sample of 2650 urban women : attributable risk and risk factors. *J Clin Epidemiol*, 45 : 377-392.

Reid, R.L. (1985) : Premenstrual syndrome. *Curr. Probl. Obstet Gynecol Fertil*; 8 (2) 1-57.

22. Lee, K.D. and Rittenhouse, C.A. (1991) : Prevalence of perimenstrual symptoms in employed women. *Women Health* 1 (17) : 17-32.
23. LeMelledo, J.M.; Bradwejn, J.; Koszycki, D. et al. (1995) : Premenstrual dysphoric disorder and response to cholecystokinin-tetrapeptide. *Gen. Psychiatry*, 52 : 605.
24. Schnurr, P.P. (1988) : Some Correlates of Prospectively Defined menstrual Syndrome. *Am J Psychiatry*, 145 (4) : 491-494.
25. Andersch, B. (1986) : Premenstrual complaints : Prevalence of premenstrual symptoms in a Swedish Urban population. *Journal of Psychiatric Obstetrics and Gynecology*, 5 : 39-49.
26. Condon, J.T. (1996) : Premenstrual Syndrome in primary care. *Primary care psychiatry*, 2 : 15-23.
27. Barnhart, K.T. Freeman, E.W. and Sondheimer, S.J. (1995) : A clinician's guide to the premenstrual syndrome. *Medical Clinics of North America*, 79 (6).
28. Moos, R.H. (1969) : Typology of menstrual cycle symptoms. *Am. J. Obstet. Gynecol.* 103 : 390.
29. Parlee, M.B. (1974) : Stereotypic beliefs about menstruation : a methodological note on the Moos Menstrual Distress Questionnaire and new data. *Psychosom. Med*, 36 : 229-240.
30. Kinch, R.A. and Steinberg, S. (1984) : Premenstrual tension and menorrhagia, *Advances in clinical obstetrics and gynecology*. Vol. 1 Edited by Osofsky, H.J. Baltimore, the Williams & Wilkins Company, PP 173-185.
31. Yonkers, K.A. (1997) : The association between premenstrual dysphoric disorder and other mood disorders. *J Clin Psychiatry*, 58 (15) : 19-23.
32. Condon, J.T. (1996) : Premenstrual Syndrome in primary care. *Primary care psychiatry*, 2 : 15-23.
33. Merikangas, K.R.; Foeldenyi, M. and Angst, J. (1993) : The Zurich study. XIX-Patterns of menstrual disturbances in the community : results of the Zurich Cohort Study. *Eur Arch Psychiatr*

Clin Neurosci, 243 : 23-32.

Hurt, S.W.; Schnurr, P.P.; Severino, S.K. et al. (1992) : Late Luteal Phase Dysphoric Disorder in 670 women evaluated for premenstrual complaints. *Am J Psychiatry*, 149 : 525-530.

DSM, TV. (1998) : Premenstrual syndrome and anxiety disorders : A psychobiological link. *Psychother Psychosom*, 7 : 57-60.

Clare, A.W. (1983) : Psychiatric and social aspects of premenstrual complaint. *Psychological Medicine Monograph Supplement*, 4 : 1-58.

Woods, N.F.; Most, A. and Dery, G.K. (1982) : Prevalence of perimenstrual symptoms. *Am J Public Health*, 72 : 1257-1264.

Mortola, J.F. (1992) : Issues in the diagnosis and research of premenstrual syndrome. *Clin Obstet Gynecol*. 35 (3) : 587-598.

Ekholm, U.B. Ekholm, N.O. and Backstrom, T. (1998) : Premenstrual Syndrome : Comparison between different methods to diagnose cyclicality using daily symptom ratings. *Acta Obstet Gynecol Scand*, 77 : 551-557.

Ussher, J.M. and Wilding, J.M. (1992) : Interactions between stress and performance during the menstrual cycle in relation to the premenstrual syndrome. *J Reprod Infant Psychol*, 10 : 83-101.

Hallman, J. : Oreland, L Edman, G. et al. (1987) : Thrombocyte monoamine oxidase activity and personality traits in women with severe premenstrual syndrome. *Acta Psychiatr Scand*, 76 : 225-234.

Mira M., Vizzard, J. and Abraham, S. (1985) : Personality characteristics in the menstrual cycle, *J Psychosom Obstet Gynecol*, 4 : 329-334.

Freeman, E.W.; Schweizer, E. and Rickels, K. (1995) : Personality factors in women with premenstrual syndrome. *Psychosomatic Medicine*, 57 : 453-459.

Wetzel, R.D.; Knesevich, M.A.; Brown, S.L. et al . (1992) : Correlates of tridimensional personality questionnaire scales with selected Minnesota Multiphasic Personality Inventory scales. *Psychol Rep*, 71 : 1027-1038.

45. Parry, B.L., Ehlers, C.L., Mostofi, N. and Phillips, F. (1996) : Personality traits in LLPDD and normal controls during follicular and luteal menstrual cycle phases. *Psychol Med*, 26 : 197-202.
46. Freeman, E.W.; Rickels, K.; Schweizer, E. and Ting, T. (1995) : Relationships between age and symptom severity among women seeking medical treatment for premenstrual symptoms. *Psychological Med* 25 : 309-315.
47. Yonkers, K.A (1997) : Treatment of Premenstrual dysphoric disorder. *Current Review of Mood & Anxiety Disorders*, 1 : 215-237.
48. Yonkers, K.A. (1997) : The association between premenstrual dysphoric disorder and other mood disorders *Clin Psychiatry*, 58 (15) : 19-25.
49. Fava, M., Pedrazzi, F.; Guaraldi, GP. et al. (1992) : Comorbidity between anxiety and depression among patients with late luteal phase dysphoric disorder. *J Anxiety Disord*. 6 : 325-335.
50. Harrison, W.M.; Endicott, J.; Nee, J. et al. (1989) : Characteristics of women seeking treatment for premenstrual syndrome. *Psychosom* 30 (4) : 405-411.
51. Pearlstein, T.B. and Stone, A.B. (1994) : Long-term fluoxetine treatment of late luteal phase dysphoric disorder. *J Clin Psychiatry*, 55 : 332-337.
52. Yonkers, K.A.; Halbreich, U.; Freeman, E. et al. (1996) : Sertraline in the treatment of premenstrual dysphoric disorder. *Psychopharm Bull*, 1996, 32 : 71-76.
53. Yonkers, K.A.; Gullion, C.; Williams, A. et al. (1996) : Paroxetine in the treatment of premenstrual dysphoric disorder. *J Clin Psychopharm* 16 : 3-8.
54. Robins L.N.; Helzer J.E.; Weissman M.M. et al. (1984) : Lifetime prevalence of specific psychiatric disorders in three sites. *Arch Gen Psychiatry*, 41 : 949-958.
55. Mikhail, N.R. (1986) : The problem of premenstrual syndrome in general practice in El-Salam district, Ismailia (with special consideration of psychological factors). Master thesis. Ain Shams University.

Stout, A.L.; Steege, J.F.; Blazer, D.G. and George, L.K. (1986) : Comparison of lifetime psychiatric diagnoses in premenstrual syndrome clinic and community samples. *J. Nerv Ment Dis*, 174 : 517-522.

Choi, P.Y.L. and Mc Keown, S. (1997) : What are young undergraduate women's qualitative experiences of the menstrual cycle? *J. Psychosom. Obstet. Gynecol.* 18 : 259-265.

