Disturbi Depressivi in gravidanza e nel puerperio

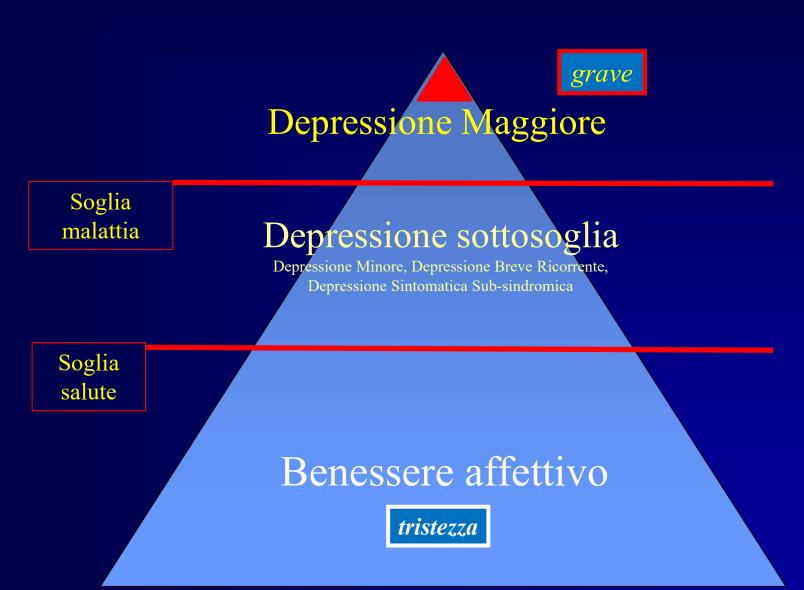
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La gravidanza non è una malattia

Depressione in gravidanza e nel puerperio



Maternity Blues

- Prevalenza: 50-85% donne.
- Esordio: entro pochi giorni dal parto (48 ore).
- Sintomi: tristezza, tendenza al pianto, sentimenti di insufficienza e di incapacità, irritabilità, ansia, difficoltà di concentrazione e di memoria, disturbi del sonno e dell'appetito, cefalea, astenia.
- Remissione: entro 2 settimane. Durata protratta: comparsa di depressione postpartum.

Burt e Stein, J Clin Psychiatry 2002; 63 (suppl 7): 9-15 Steiner et al. J Affect Disord 2003;74:67-83

Edinburgh Postnatal Depression Scale¹ (EPDS)

Name:	Address:
Your Date of Birth:	-
Baby's Date of Birth:	Phone:
As you are pregnant or have recently had a baby, we wou the answer that comes closest to how you have felt IN TH	
Here is an example, already completed.	
I have felt happy: □ Yes, all the time ⊠ Yes, most of the time □ No, not very often □ No, not at all This would mean: "I have fel Please complete the other que	It happy most of the time" during the past week, uestions in the same way.
In the past 7 days:	
1. I have been able to laugh and see the funny side of things As much as I always could Not quite so much now Definitely not so much now Not at all 2. I have looked forward with enjoyment to things As much as I ever did Rather less than I used to Definitely less than I used to Hardly at all *3. I have blamed myself unnecessarily when things went wrong Yes, most of the time Yes, some of the time Not very often No, never	*6. Things have been getting on top of me
I have been anxious or worried for no good reason No, not at all Hardly ever Yes, sometimes Yes, very often	*9 I have been so unhappy that I have been crying Yes, most of the time Yes, quite often Only occasionally No, never
*5 I have felt scared or panicky for no very good reason Yes, quite a lot Yes, sometimes No, not much No, not at all	*10 The thought of harming myself has occurred to me Yes, quite often Sometimes Hardly ever Never
Administered/Reviewed by	Date
1 Source: Cox, J.L., Holden, J.M., and Sagovsky, R. 1987. Detection of p	

necol 2010; 115;394-95

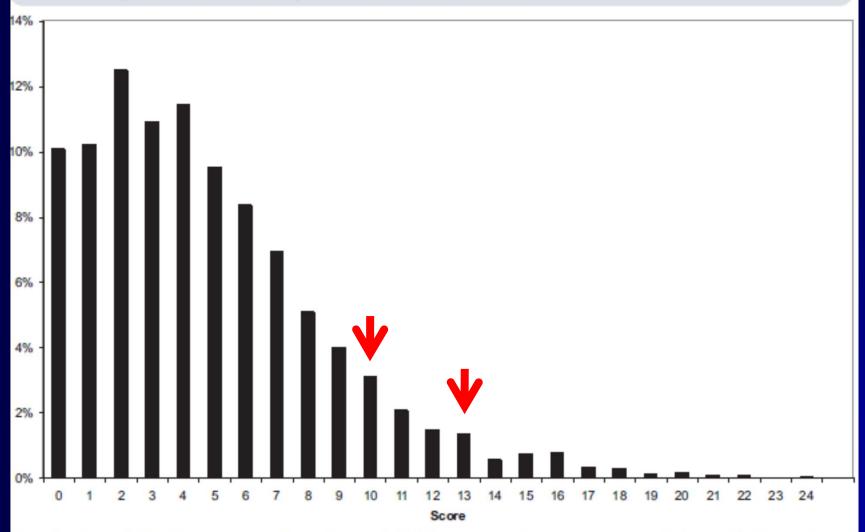
0-9: some symptoms of distress.

10-12 : symptoms of distress that may be discomforting.

13 +: the likelihood of depression is high.

Item 10: scores 1, 2 or 3 on item 10 requires further evaluation.

Distribution of antepartum and postpartum Edinburgh Postnatal Depression Scale Scores



Distribution of EPDS scores in the cohort of 1548 women who were screened from 24-28 weeks of gestation and again after delivery (total of 3168 completed screens).

Heterogeneity in perinatal depression: how far have we come? A systematic review

Hudson et al, Arch Womens Ment Health. 2017; 20: 11-23

Heterogeneous nature of this disorder

- 1. three heterogeneous patterns: low, medium or chronic-high symptom (EPDS) levels
- 2. social and psychological risk factors were the most common group of predictors related to a higher burden (high sum of score) of depressive symptoms: low education, negative life events, ethnic-minority status, unintended pregnancy, mood or anxiety symptoms during pregnancy, and prior history of psychopathology;
- 3. poorer health outcomes for children of mothers assigned to high burden symptom

Limited evidence on the specific symptoms and symptom configurations that make up PND heterogeneity

Major and Minor Depression in Pregnancy

Marchesi et al Obstet Gynecol 2009; 113: 1292-8

No state	Seneral Surv	Marine land	Manuff proof day
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0	1	2	3
add columns:			
TOTAL:			1
	Sc	omewhat difficu	
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PHQ-9

Spitzer et al, JAMA 1994; 272: 1749-56

Screening for Depression During and After Pregnancy

Obstest Gynecol 2010; 115;394-95

Screening Tool	Number of Items	Time to Complete	Sensitivity/ specificity
Edinburgh Postnatal Depression Scale (EPDS)	10	Less than 5 min	Sensitivity: 59-100% Specificity: 49-100%
Postpartum Depression Screening Scale (PDSS)	35	5–10 min	Sensitivity: 91–94% Specificity: 72–98%
Patient Health Questionnaire-9 (PHQ-9)	9	Less than 5 min	Sensitivity: 75% Specificity: 90%
Beck Depression Inventory (BDI)	21	5–10 min	Sensitivity: 47.6–82% Specificity: 85.9–89%
Beck Depression Inventory-II (BDI-II)	21	5–10 min	Sensitivity: 56–57% Specificity: 97–100%
Center for Epidemiologic Studies Depression Scale (CES-D)	20	5–10 min	Sensitivity: 60% Specificity: 92%
Zung Self-Rating Depression Scale (Zung SDS)	20	5–10 min	Sensitivity: 45–89% Specificity: 77–88%

Major and Minor Depression in Pregnancy

Marchesi et al Obstet Gynecol 2009; 113: 1292-8

154 women

Major depression was diagnosed in 19 women (12.3%) and minor depression in 28 (18.1%), whereas the remaining 107 women did not show any depressive symptoms (controls). In five women, minor depression symptoms preceded (n=3) or followed (n=2) major depression.

Table 2. Onset and Duration of Depressive Episodes in Pregnant Women

	Major Depression (n=19)	Minor Depression (n=28)
Onset of depression (trimester)		
First*	7 (36.8)	17 (60.7)
Second	4 (21.1)	7 (25.0)
Third	8 (42.1)	4 (14.3)
Duration of depression* (mo)	(3)	20 00
1	5 (26.3)	16 (57.1)
2	7 (36.8)	7 (25.0)
3-4	3 (15.8)	5 (17.9)
5 or more	4 (21.1)	_

Depressione Post-partum

- Esordio: primi tre mesi dal parto (primo mese).

- Prevalenza: 10-22%

- Sintomatologia : sovrapponibile alla Depressione Maggiore
- Effetti sullo sviluppo psichico del bambino: attaccamento insicuro e ambivalente, compromissione tono emozionale, sviluppo cognitivo e relazionale, manifestazioni psicopatologiche
- Prognosi: remissione entro 6-12 mesi

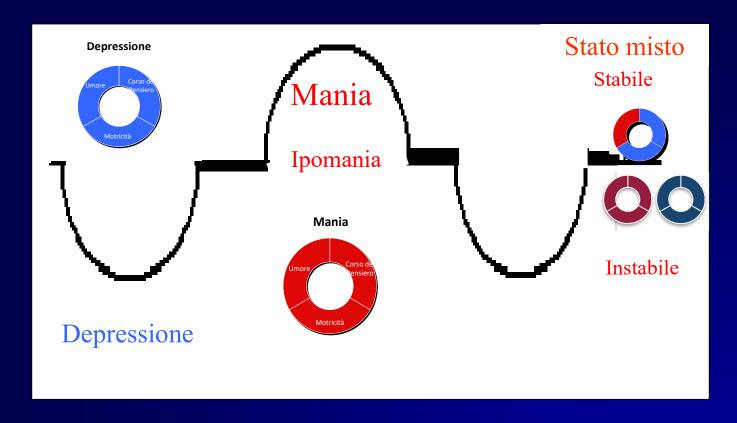
Burt e Stein, J Clin Psychiatry 2002; 63 (suppl 7): 9-15 Steiner et al. J Affect Disord 2003;74:67-83

Depressione in gravidanza e nel puerperio



Soglia malattia

Disturbi dell' Umore



Disturbo Bipolare → fasi depressive + fasi ipo/maniacali o miste Depressione Maggiore Ricorrente (Unipolare) → fasi depressive

Postpartum depression: a disorder in search of a definition

Arch Womens Ment Health (2010) 13:37-40



Diagnostic considerations

The diagnostic entities most frequently observed after birth in both the Kendell et al. (1987) and Munk-Olsen et al. (2006) studies were major mood disorders, with a particular risk for bipolar disorder. Munk-Olsen et al observed highly significant relative risks of 23.33 in the first 30 days and 6.30 in the 31–60 days after birth for bipolar disorder. Kendell et al reported an RR of 21.7 for admission with psychosis (primarily bipolar disorder) within 30 days of

Psicosi Post-partum

- Prevalenza: 0.1-0.3 % di puerpere.
- Esordio: prima settimana dopo il parto (48-72 ore).
- Sintomi:
- intense e rapide oscillazioni dello stato di coscienza;
- allucinazioni visive o uditive a contenuto triste o terrifico;
- temi deliranti di tipo persecutorio o depressivo e incentrati sulla relazione madre-bambino;
- ansia, agitazione;
- rischio di suicidio e di infanticidio.
- Prognosi:
- favorevole con remissione in 6-12 mesi nell' 80% casi;
- tendenza alla ricorrenza nel postpartum (75-90%);
- 70% Disturbo Bipolare o Depressione Maggiore (psicotici)

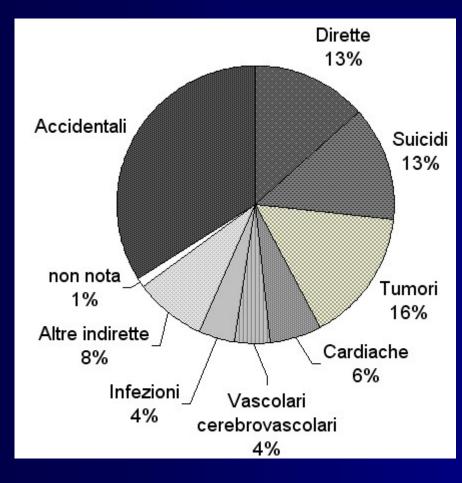






Mortalità e morbosità materna in Emilia-Romagna. Rapporto 2001-2007

Distribuzione per causa di 97 casi di morti materne



Agli iniziali 13 casi di suicidio, 5 casi sono stati aggiunti in seguito all'analisi della documentazione clinica. Suicidi → 18%

L'età media al decesso: 29,2 anni

Le morti sono avvenute:

- dopo il parto 44%;
- dopo un aborto volontario 33%
- dopo un aborto spontaneo 11%
- in gravidanza 11%

Prevalence of suicidality during pregnancy and the postpartum

Arch Womens Ment Health (2005) 8: 77-87

Summary

While suicide deaths and attempts are lower during pregnancy and the postpartum than in the general population of women, when deaths do occur, suicides account for up to 20% of postpartum deaths.

fact that cannot be accounted for solely by differences in levels of depression. Increased social support, concern for the unborn child, and more contact with health care providers may work together to reduce the risk of suicide.

- Among the pregnant women who die by suicide, teenagers were reported to be at greater risk
- as were women in cultures that stigmatize motherhood among unmarried women

Prevalence of suicidality during pregnancy and the postpartum

Arch Womens Ment Health (2005) 8: 77-87

Summary

- While suicide deaths and attempts are lower during pregnancy and the postpartum than in the general population of women, when deaths do occur, suicides account for up to 20% of postpartum deaths.
 - Consistent with lower suicide rates in pregnancy, suicide rates in the postpartum are lower than expected in the general population.
 - suicide rates are considerably elevated among groups at risk including those with psychiatric diagnoses and teenage mothers

Prevalence of suicidality during pregnancy and the postpartum

Arch Womens Ment Health (2005) 8: 77-87

Summary

- Self-harm ideation is more common than attempts or deaths, with thoughts of self-harm during pregnancy and the postpartum ranging from 5 to 14%.
- The risk for suicidality is significantly elevated among depressed women during the perinatal period, and suicide has been found to be the second or leading cause of death in this depressed population.







Mortalità e morbosità materna in Emilia-Romagna. Rapporto 2001-2007

Un contatto con i servizi psichiatrici territoriali preesistente o attivato durante il percorso assistenziale della gravidanza è stato documentato nel 28% dei casi,

una percentuale che corrisponde a metà dei casi con fattori di rischio noti.

Dal punto di vista clinico è stato possibile identificare alcuni elementi critici del percorso assistenziale:

- carente raccolta di informazioni su malattia mentale e abuso di sostanze, condizioni in alcuni casi non indagate o sottovalutate dai professionisti;
 - sono stati inoltre riscontrati difetti di comunicazione fra i servizi

Rischio di Depressione in gravidanza e nel puerperio

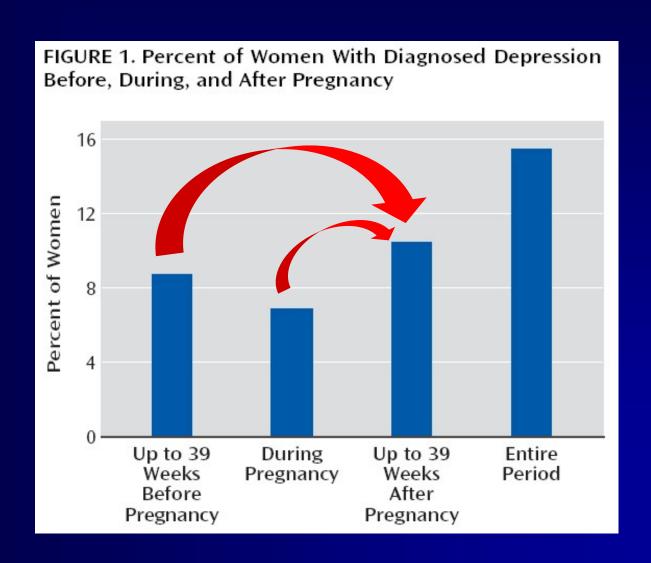
- se conosci il rischio per quella donna, puoi intervenire precocemente.

Risk factors for depressive symptoms during pregnancy: a systematic review Am J Obstet Gynecol 2010.

Factor	Total no. of studies	Total no. of subjects	Bivariate trend of association ^a	Multivariate trend of association ^a	
Anxiety ¹⁸⁻²⁸	11	4696	++++	b lack	of sample size
Life stress, composite ^{20,24,29-32,43,47-49,52,55-57,68,69,72,73}	18	9973	+++	+++	or darripro dizo
Life events, total (positive and negative)	15	9645	+++	Inconsistent	
Negative life events			++++	+++	
Daily hassles	5	1134	c	b	
Personal history of depression ^{24,32,54,62,69,74}	6	3566	+++	b	
Social support ^{20,22,24,27,28,30-35,43,48,49,52,53,55-57,60,64,68,69,73}					
Lack of social support, any source	17	5752	+++	+	
Lack of social support, partner	9	7139	++++	++++	
Domestic violence ^{24,29,30,46,54,57,67}	7	3738	+	++	
Unintended pregnancy ^{24,60,61,63,64,68}	6	11,470	+++	b	
Relationships ^{20,22,24,27,32-38,43,46,48,50,52,59,60,62,64,65,68-73}					
Cohabitation	19	12,483	++	Inconsistent	
Poor relationship quality	11	4005	+++	e N	o effect.
Demographics					
Public insurance/uninsured ^{29,30,34,50,57,58}	6	2008	+++	b	
Medicaid (US studies only)			+++	b	
Socioeconomic status ^{56,59,64,69,73}	5	2805	c	c	
Lower income ^{20,31,32,37-39,46,48,49,56,64}	11	6285	+	b	
Unemployment ^{20,27,32,35,38,39,47,49,50,64,68,71-73}	14	9417	c	Inconsistent	
Lower education 20,21,30,32-39,43,49,50,56,62,68,71-73	20	11,529	+	c	
Maternal age ^{20,22,30,32-39,43,46,48,49,59,62,64,67,68,71,73}	22	13,837	Inconsistent	c	
Maternal race/ethnicity ²⁹⁻⁴²	14	6671	Inconsistent	c	
Substance abuse			***************************************		
Smoking ^{30,32,33,36,38,41,54,56,57,59,71}	11	6641	+	c	
Alcohol use 32,33,36,44,45,54,56,57,66,71	10	10,621	Inconsistent	c	
Illicit drug use ^{30,32,33,38,46,49,57,64}	8	3010	c	Inconsistent	
Nulliparity ^{22,24,34-37,39,46-48,51,59,62,64,66,68,72,73}	18	9786	c	Inconsistent	
Poor obstetric history 19,24,32,34,39,47,49,59,64,68	10	6888	c	c	

Clinically Identified Maternal Depression Before, During, and After Pregnancies Ending in Live Births

Am J Psychiatry 2007; 164:1515–1520



Postnatal depression: A global public health perspective

Perspectives in Public Health 2009 129: 221

It is recommended that all women should have their mental and emotional health assessed postnatally so that the presence of PND can be ascertained.



prenatally





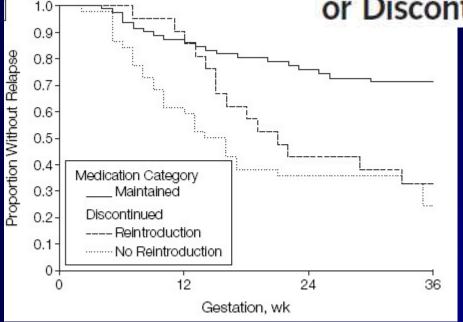


Mortalità e morbosità materna in Emilia-Romagna. Rapporto 2001-2007

Per i casi in cui è stata possibile una valutazione della qualità del percorso assistenziale, i fattori di substandard care rilevati sono stati:

- la mancata attivazione e presa in carico dei servizi territoriali psichiatrici alla dimissione di donne sintomatiche
- e/o la sospensione inappropriata della terapia con psicofarmaci durante la gravidanza.

Relapse of Major Depression During Pregnancy in Women Who Maintain or Discontinue Antidepressant Treatment



JAMA. 2006;295:499-507

Table 3. Relapse of Major Depression During Pregnancy

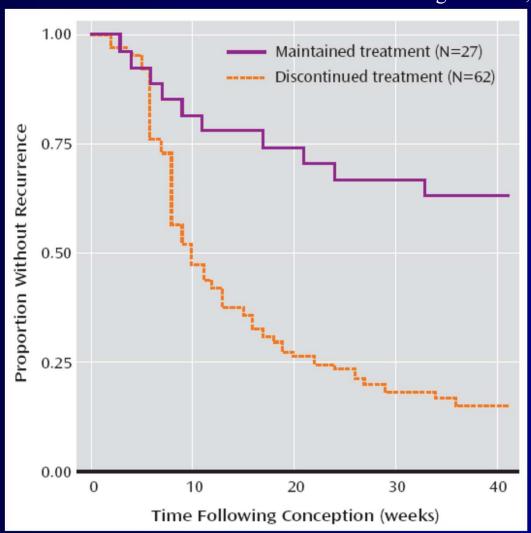
Relapse Status All Women		Medication Status				
		Maintained	Increased	Decreased	Discontinued	
No relapse)	115 (57.2)	61 (74.4)	11 (55.0)	22 (64.7)	21 (32.3)
Relapse by All	/ trimester	86 (42.8)	21 (25.6)	9 (45.0)	12 (35.3)	44 (67.7)
Firs	st	44 (51.2)	→11 (52.4)	7 (77.8)	5 (41.7)	21 (47.7)
Sec	cond	31 (36.0)	9 (42.9)	2 (22.2)	3 (25.0)	19 (43.2)
Thir	rd	11 (12.8)	1 (4.8)	0 (0.0)	4 (33.3)	4 (9.1)

Table 2. Clinical Characteristics of Pregnant V	Vomen With Histor	les of Major Depress	ion Across Differen	t Medication Treatm	ent Conditions*
Variable	All (N = 201)	Maintained (n = 82)	Increased (n = 20)	Decreased (n = 34)	Discontinued (n = 65)
Age at onset, y	44 (00 0)	10 (04 0)	0.467)	7 (04 0)	45 (04.0)
<14	44 (23.2)	19 (24.0)	3 (16.7)	7 (21.9)	15 (24.6)
14-17	45 (23.7)	19 (24.0)	5 (27.8)	4 (12.5)	17 (27.9)
18-25	50 (26.3)	19 (24.0)	7 (38.9)	11 (34.4)	13 (21.3)
>25	51 (26.8)	22 (28.0)	3 (16.7)	10 (31.2)	16 (26.2)
Duration of illness, y <5	24 (12.6)	9 (11.4)	2 (11.1)	4 (12.5)	9 (14.8)
5-14	67 (35.3)	27 (34.2)	9 (50.0)	14 (43.8)	17 (27.9)
15-20	61 (32.1)	28 (35.4)	4 (22.2)	10 (31.2)	19 (31.2)
>20	38 (20.0)	15 (19.0)	3 (16.7)	4 (12.5)	16 (26.2)
No. of prior episodes	36 (20.0)	15 (18.0)	3 (10.7)	4 (12.0)	10 (20.2)
<3	46 (23.4)	18 (22.2)	4 (20.0)	8 (23.5)	16 (25.8)
3-4	64 (32.5)	31 (38.3)	5 (25.0)	9 (26.5)	19 (30.6)
5-6	45 (22.8)	16 (19.8)	5 (25.0)	9 (26.5)	15 (24.2)
>6	42 (21.3)	16 (19.8)	6 (30.0)	8 (23.5)	12 (19.4)
No. of episodes in prior pregnancies Never pregnant	49 (25.3)	20 (25.3)	4 (21.0)	12 (36.4)	13 (20.6)
None	96 (49.5)	37 (46.8)	11 (57.9)	11 (33.3)	37 (58.7)
>1	49 (25.3)	22 (27.8)	4 (21.0)	10 (30.3)	13 (20.6)
No. of prior postpartum episodes Never pregnant	49 (26.1)	20 (26.3)	4 (22.2)	12 (36.4)	13 (21.3)
None	84 (44.7)	38 (50.0)	6 (33.3)	11 (33.3)	29 (47.5)
>1	55 (29.3)	18 (23.7)	8 (44.4)	10 (30.3)	19 (31.2)
Duration of prior episode, wk 1-6	43 (23.1)	13 (16.9)	5 (26.3)	9 (29.0)	16 (27.1)
7-12	48 (25.8)	21 (27.3)	4 (21.0)	7 (22.6)	16 (27.1)
13-36	49 (26.3)	22 (28.6)	7 (36.8)	8 (25.8)	12 (20.3)
>36	46 (24.7)	21 (27.3)	3 (15.8)	7 (22.6)	15 (25.4)
Time since onset of most recent episode, wk	28 (15.3)	10 (13.0)	4 (21.0)	5 (16.1)	9 (16.1)
25-72	54 (29.5)	16 (20.8)	9 (47.4)	12 (38.7)	17 (30.4)
73-144	47 (25.7)	19 (24.7)	4 (21.0)	7 (22.6)	17 (30.4)
>144	54 (29.5)	32 (41.6)	2 (10.5)	7 (22.6)	13 (23.2)
No. of attempts to discontinue antidepressant medication	04 (28.0)		2 (10.0)		10 (20.2)
0	49 (26.5)	21 (28.0)	3 (16.7)	13 (40.6)	12 (20.0)
1	57 (30.8)	23 (30.7)	3 (16.7)	9 (28.1)	22 (36.7)
2	35 (18.9)	17 (22.7)	5 (27.8)	5 (15.6)	8 (13.3)
>2	44 (23.8)	14 (18.7)	7 (38.9)	5 (15.6)	18 (30.0)

Article

Risk of Recurrence in Women With Bipolar Disorder During Pregnancy: Prospective Study of Mood Stabilizer Discontinuation

Viguera et al, Am J Psychiatry 2007; 164: 1817-1824



conclusions: Discontinuation of mood stabilizer, particularly abruptly during pregnancy carries a high risk for new morbidity in women with bipolar disorder, especially for early depressive and dysphoric states. However, this risk is reduced markedly by continued mood stabilizer treatment. Treatment planning for pregnant women with bipolar disorder should consider not only the relative risks of fetal exposure to mood stabilizers but also the high risk of recurrence and morbidity associated with stopping maintenance mood stabilizer treatment.

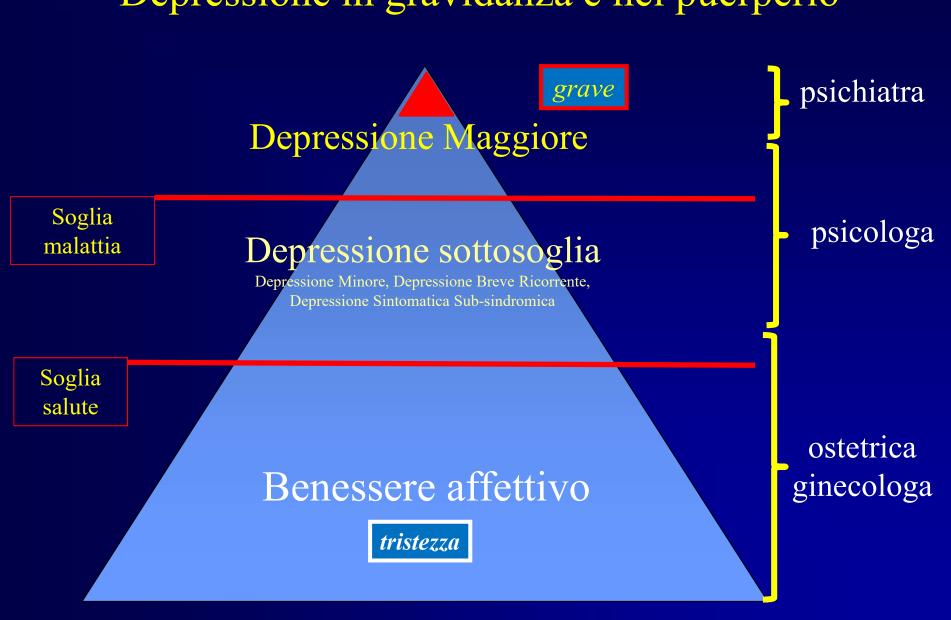
Article

Risk of Recurrence in Women With Bipolar Disorder During Pregnancy: Prospective Study of Mood Stabilizer Discontinuation

TABLE 2. Morbidity During Pregnancy Versus Treatment Status						
-			Subjects Who	Maintained	Subjects Who	Discontinued
Variable	All Subjec	ts (N=89)	Treatmer	nt (N=27)	Treatmer	it (N=62)
	N	%	N	%	N	%
Risk of at least one recurrence ^a	63/89	70.8	10/27	37.0	53/62	85.5
First recurrence risk by trimester						0.0000000000000000000000000000000000000
First	42/89	47.2	6/27	22.2	36/62	58.1
Second	15/47	31.9	3/21	14.3	12/26	46.2
Third	6/32	18.8	1/18	5.6	5/14	35.7
Recurrence polarity (all recurrences)b						
Depression	34/89	38.2	5/27	18.5	29/62	46.8
Mixed state	26/89	29.2	0/27	0.0	26/62	41.9
Hypomania	15/89	16.8	7/27	25.9	8/62	12.9
Mania	6/89	6.7	2/27	7.4	4/62	6.5
Percent of pregnancy weeks ill						
	Mean	SD	Mean	SD	Mean	SD
All cases ^c	32.8	31.5	8.8	21.3	43.3	29.6
	N	%	N	%	N	%
Stable subjects (%) ^d	26/89	29.2	17/27	63.0	9/62	14.5

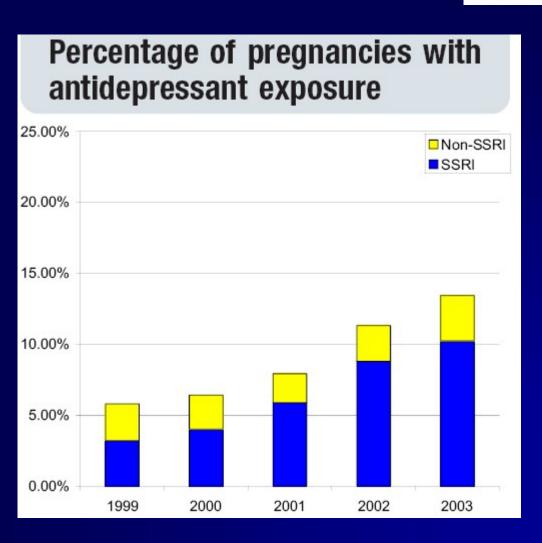
Viguera et al, Am J Psychiatry 2007; 164: 1817-1824

Depressione in gravidanza e nel puerperio



Increasing use of antidepressants in pregnancy

Am J Obstet Gynecol 2007;196;544.e1-544.e5.



Delivery outcome after maternal use of antidepressant drugs in pregnancy: an update using Swedish data

Table 1. Number of women using specific antidepressant drugs either before the first antenatal visit ('Early') or prescribed the drugs during pregnancy ('Later')

Drug name	ATC code	Early	Later
TCAs	N06AA	1662	784
Imipramine	N06AA02	10	3
Clomipramine	N06AA04	1208	592
Lofepramine	N06AA07	6	1
Amitriptyline	N06AA09	379	158
Nortriptyline	N06AA10	33	23
Protryptyline	N06AA11	1	0
Maprotiline	N06AA21	9	1
SSRIs	N06AB	10 170	4809
Fluoxetine	N06AB03	1522	892
Citalopram	N06AB04	3950	1648
Paroxetine	N06AB05	1208	405
Sertraline	N06AB06	3297	1825
Fluvoxamine	N06AB08	42	17
Escitalopram	N06AB10	153	56
Unspecified	N06AB00	86	39
MOAIs	N06AG	37	18
Moclobemide	N06AG02	37	18
SNRIs	N06AX	1351	538
Mianserin	N06AX03	85	33
Nefazodone	N06AX06	44	7
Mirtazapine	N06AX11	277	123
Bupropion	N06AX12	37	9
Venlafaxine	N06AX16	859	363
Reboxetine	N06AX18	28	6
Duloxetine	N06AX21	37	4
Unspecified	N06A	10	0
antidepressants		D _e	ois and Ka

Reis and Kallen, Psychol Med 2010; 40: 1723-33

Prevalence of Antidepressant Use during Pregnancy in Denmark, a Nation-Wide Cohort Study

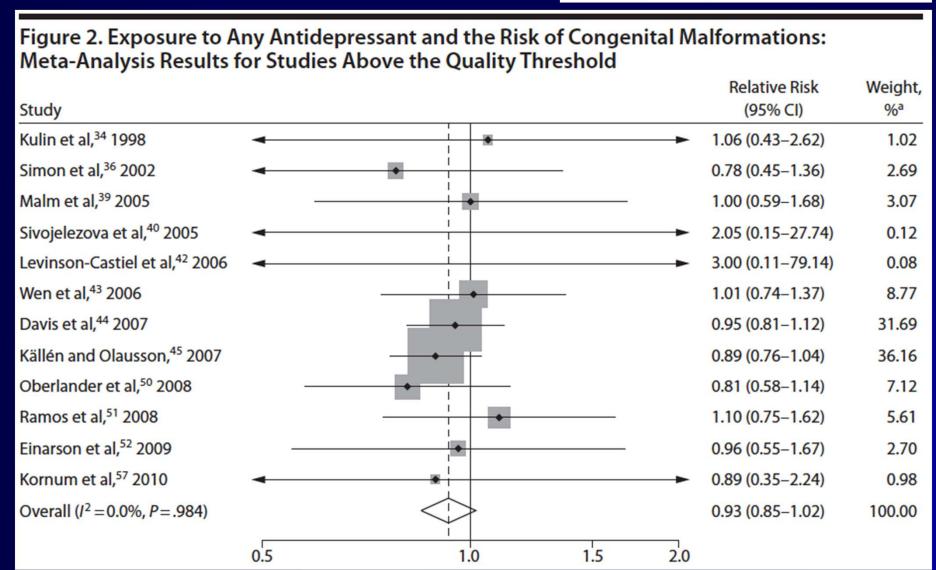
Table 3. Number of women exposed to an antidepressant during pregnancy.

	Trimester		
	First	Second	Third
Any antidepressant	18273	13039	9721
SSRI	15403 (84.29)	11370 (87.20)	8641 (88.89)
Citalopram	6657 (36.43)	4306 (33.02)	2850 (29.32)
Escitalopram	1539 (8.42)	722 (5.54)	344 (3.54)
Fluoxetine	3898 (21.33)	3618 (27.75)	2927 (30.11)
Paroxetine	1779 (9.74)	1164 (8.93)	816 (8.39)
Sertraline	3059 (16.74)	2565 (19.67)	2328 (23.95)
TCA	1101 (6.03)	748 (5.74)	479 (4.93)
Amitriptyline	578 (3.16)	292 (2.24)	120 (1,23)
Clomipramin	125 (0.68)	89 (0.68)	57 (0.59)
Dosulepin	40 (0.22)	42 (0.32)	36 (0.37)
Imipramin	61 (0.33)	32 (0.25)	15 (0.15)
Nortriptyiline	327 (1.79)	308 (2.36)	254 (2.61)
Other	3039 (16.63)	1655 (12.69)	934 (9.61)
Mianserin	270 (1.48)	113 (0.87)	57 (0.59)
Mirtazapine	876 (4.79)	348 (2.67)	142 (1.46)
Venlafaxine	1687 (9.23)	1109 (8.51)	703 (7.23)

Plos One April 2013; 8: e63034

Antidepressant Exposure During Pregnancy and Congenital Malformations: Is There an Association? A Systematic Review and Meta-Analysis of the Best Evidence

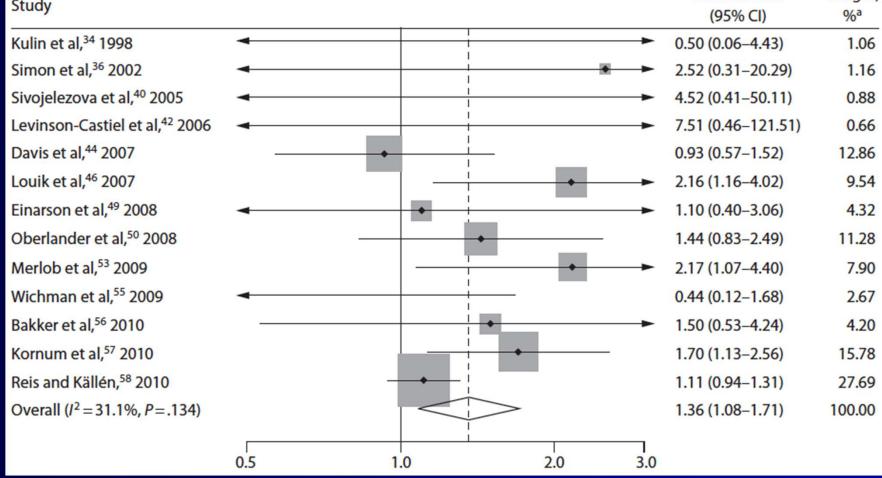
J Clin Psychiatry 2013;74(4):e293-e308



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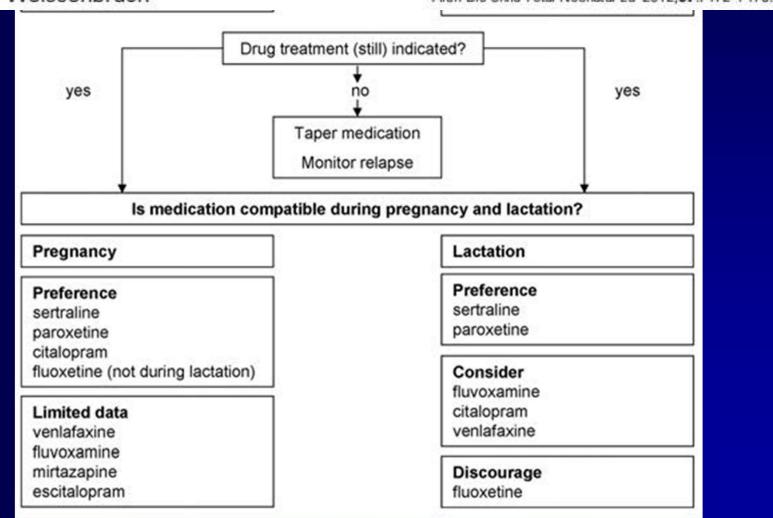
J Clin Psychiatry 2013;74(4):e293-e308

Conclusions: Overall, antidepressants do not appear to be associated with an increased risk of congenital malformations, but statistical significance was found for cardiovascular malformations. Results were robust in several sensitivity analyses. Given that the RRs are marginal, they may be the result of uncontrolled confounders. Although the RRs were statistically significant, none reached clinically significant levels.

Maternal use of SSRIs, SNRIs and NaSSAs: practical recommendations during pregnancy and lactation

S D Sie,¹ J M B Wennink,² J J van Driel,^{2,3} A G W te Winkel,⁴ K Boer,⁵ G Casteelen,⁶ M M van Weissenbruch¹

Arch Dis Child Fetal Neonatal Ed 2012;97:F472–F476.



Maternal Medication, Drug Use, and Breastfeeding



Rowe et al, Child Adolesc Psychiatric Clin N Am 2015; 24: 1-20

3 6		
Table 1 Antidepressants a	nd reported levels	in breast milk
Antidepressant	Relative Infant Dose (RID) %	Comments
Selective Serotonin	Reuptake Inhibit	tors (SSRIs)
Citalopram Escitalopram Fluvoxamine Fluoxetine Sertraline Paroxetine	3.6 ⁹⁰ 5.3 ⁹⁵ 1.6 ⁸ 5–9 ^{93,94} 0.54 ⁷ 1.4 ⁹	Compatible: SSRIs are recommended first-line agents for depression and anxiety, and are suitable when breastfeeding. There have been 2 cases of excessive somnolence, decreased feeding, and weight loss with citalopram; however, most new data suggest these side effects are rare. 91,92 Fluoxetine has been reported to cause colic, fussiness, and crying 93,94
Serotonin Norepin	ephrine Reuptake	Inhibitors
Venlafaxine Desvenlafaxine Duloxetine	8.1 ⁹⁶ 6.8 ⁹⁷ 0.1 ⁹⁸	Compatible: No adverse events reported in breastfed infants with these 3 medications
Ac. valproico Carbamazepina Litio Lamotrigina	1.4-1.7 5.9 30.1 9.2	

Depression in pregnancy: time to stop terrifying pregnant women.

J Popul Ther Clin Pharmacol. 2012;19(3):e369-70.

Recent studies have powerfully demonstrated that women with depression, whether treated with SSRIs or skip treatment, report double risk of cardiovascular malformations.

Yet, physicians and epidemiologists continue to terrify women who are at serious life threatening risks if not treated pharmacologically, with non evidence-based information.

This paper calls for immediate stop of such practice.

La comprensione,
il supporto,
l'aiuto sono spesso sufficienti,
ma in alcuni casi
il trattamento farmacologico
è necessario

Non è una colpa avere la depressione

Non avere paura Chiedi aiuto

Perinatal Psychiatry: An Emerging Specialty

- 1. Has the time come for a recognized specialty in Perinatal Psychiatry?
- 2. Or is the information available adequately disseminated and understood to support the treatment of perinatal women more broadly?

Marlene Freeman, J Clin Psychiatry 2014; 75: 1087-86

Is Perinatal Psychiatry a Needed Specialty?

1ª fase

Several health care providers are involved in recognizing and managing mental disorders in the perinatal period: social workers, nurses, gynecologists, psychologists, pediatricians, general practitioners, and psychiatrists. All of these health care providers should receive adequate training, provided by a specialized multidisciplinary team, that would include acquisition of the growing amount of evidence-based findings on perinatal psychiatry.

Is Perinatal Psychiatry a Needed Specialty?

2ª fase

Once the available information is adequately disseminated, perinatal mental health care would not require a specialized team for the majority of women, particularly those with mild-to- moderate depressive and anxiety disorders, which seldom need intensive treatment,3,4 and those with mental distress induced by negative life events, lack of social support, and domestic violence. Since these latter risk factors do not occur only in the perinatal period, they can usually be managed by nonspecialized mental health care providers.

Is Perinatal Psychiatry a Needed Specialty?

2^a fase

In contrast, a specialized multidisciplinary team should be involved in the management of severe mental disorders occurring in the perinatal period. Therefore, women with disorders other than mild-to-moderate anxiety and depressive disorders should be referred to a multidisciplinary team specialized in perinatal psychiatry. One of the components of this team should be a psychiatrist particularly versed in the pharmacologic treatment of mental disorders in the perinatal period.