Factors associated with pregnancy-related concerns in women with inflammatory rheumatic diseases – an analysis of a nationwide pregnancy cohort

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Conclusions

At least one in five women with IRD is "very concerned" about pregnancy and infant health due to the underlying rheumatic disease. A higher RAID score, nulligravidae, treatment changes and younger age were associated with higher levels of concern. Therefore, patient concerns should be an essential part of individual counselling for women with IRD who want to become pregnant or are already pregnant.

Background

Pregnancies in women with chronic diseases are often accompanied by concerns about possible complications during pregnancy and about the health of the child^[1]. In women with inflammatory rheumatic diseases (IRD) this can lead to a reduced number of offspring[2]. To date, surveys and interviews, but no data quantifying these worries and factors influencing them, have been published. (1) Lange et al. Z Geburtshilfe Neonatol 2015:219(4):161-9. (2) Clowse et al. Arthritis Care Res 2012:64(5):668-74.

Objectives

To investigate and quantify pregnancy concerns in women with various underlying IRD, and to identify factors that have an impact on these sorrows.

Patients and Methods

- Data source: German pregnancy registry Rhekiss, a nationwide, multicentre, web-based longitudinal cohort study, initiated in 2015 to investigate pregnancies and related issues in women with various IRD
- Enrolment: Women with IRD can either be enrolled if they are planning a pregnancy (cohort 1) or if they are already pregnant prior to gestational week 20 (cohort 2)
- Data reporting: At regular, pre-defined follow-up visits by rheumatologist and patient
- Selection criteria: Women who had answered the question at baseline visit regarding their concerns about pregnancy and child's health due to the IRD between 09/2015 and 10/2022
- Data analysis:
 - Descriptive analysis of categorical answers and patient characteristics
 - Impact of different factors associated with patients' anxieties was estimated by multivariable ordinal logistic regression using the proportional odds model and combining the two cohorts. Missing values were replaced by single imputation.

Results



Not really

32.7

7.7

1.9

58.0%

11.1%

54 1%

68 5%

0.4

36

32.0

79

1.4

50.0%

22.2%

31.4%

50.0%

455 womer A little

219

32.0

8.2

2.0

61.3%

10.5%

53 9%

69 3%

0.5

A lot

140

31.5

61.1%

15.8%

61.4%

70.3%

0.6

3.2

8

Not at all

55

34 3

8.6

1.0

41.9%

35.3%



155

32.9

8.7

1.7

4.7%

40.3%

51.2%

Not really

Cohort 2: 784 women A little

398

32.6

7.9

2.3

52.4%

11.1%

44.5%

A lot

176

32.1

7.7

3.2

61.9%

12.3%

45.8%

Table 1:
Characteristics of
patients stratified
by cohort and by
level of concern.
*Either due to the plan to
become pregnant or due
to the ongoing pregnancy.
RAID (Rheumatoid Arthritis

composite index to as

disease hurden in rhe

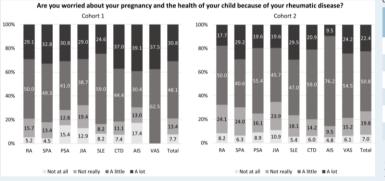
diseases, range 0-10 w

numbers indicatina wo

6	
of fied	Number of patients
by	Age in years (mean)
rn.	Disease duration in year
	Moderate to severe dise
n to	RAID score (mean)
lue incv.	Current flare
thritis even-item	Treatment change*
sess the ımatic	Nulligravidae
vith higher orse status.	# Comorbidities (mean)

Number of patients				
Age in years (mean)				
Disease duration in years (mean)				
Moderate to severe disease				
RAID score (mean)				
Current flare				
Freatment change*				
Nulligravidae				

Figure: Level of concerns stratified by cohort and by IRD diagnosis.



45.2% 54.7% 53.8% 59.9% 0.8 0.6 0.5 0.8

Table 2: Factors associated with higher level of concerns estimated by logistic regression.

	Odds ratio [95% confidence interval]
RAID score increase	1.35 [1.27; 1.43]
Nulligravidae	1.29 [1.01; 1.65]
Treatment change	1.26 [1.00; 1.60]
Age	0.96 [0.94; 0.99]
# Comorbidities	1.13 [0.98; 1.29]
Cohort 2 vs. cohort 1	0.74 [0.60; 0.93]

*Abbreviations: AIS. autoinflammatory syndrome: CTD. connective tissue disease (w/o SLE): JIA, juvenile idiopathic arthritis: PSA, psoriatic arthritis; RA, rheumatoid arthritis; SLE, systemic lupus erythematosus; SPA snandulaarthritis: VAS vasculitis

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