

## Research Article

# Alloimmune Disorders in Recurrent Implantation Failures after IVF/ET Procedures

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## Introduction

Recurrent Implantation Failures (RIF) can be defined as the absence of implantation after 2 to 6 consecutive cycles of IVF, ICSI or frozen embryo replacement cycles where the cumulative number of transferred high-quality embryos was no less than four for cleavage-stage embryos and no less than two for blastocysts. RIF can be considered with appropriate developmental stage with determination of implantation by an increasing quantitative Human Chorionic Gonadotropin (hCG) level and usually include also biochemical pregnancies [1].

Successful pregnancy implantation is related to adequate utero-placental circulation, absence of uterine anomalies and adequate endometrial receptivity. Immunological causes and thrombophilias play important roles in implantation failure through mechanisms similar to recurrent miscarriages [2]. Pregnancy is believed to appear a period of immunomodulation with greater pro-inflammatory activity at the beginning of pregnancy and change during the second and third trimester [3].

The aim of our study was to find if a misbalance of Th1/Th2 cytokines in patients who underwent IVF/ET procedures may influence on the pregnancy outcome. Our study evaluated peripheral blood cytokines profile in patients after consecutive implantation failures following IVF/ET procedures in comparing to the patients after successful pregnancy outcome following first attempt of IVF/ET procedures.

## Material and Methods

One hundred fifty-eight patients with a history of two and more implantation failures after IVF/ET procedures were evaluated. All patients had a normal ovarian reserve as measured by the levels of Anti-Müllerian Hormone (AMH), Follicle-Stimulating Hormone (FSH) and oestradiol. The indications for IVF/ET procedures were unexplained infertility or male infertility.

The study was performed between January 2021 and January 2023. Patients were registered at the Department of Operative and Endoscopic Gynaecology at the Medical University of Lodz. The patients gave written consent for participation in the study, and the study was approved by the Ethics committee. The control group comprised seventy six women who had successfully given birth at the first attempt of ET. The indications for IVF procedures were the same as those for the study group.

The IVF/ET procedures were similar for all patients. Chromosomal abnormalities of the male and female were excluded after their karyotype analysis. Embryonic aneuploidy was evaluated using Fluorescence In Situ Hybridization (FISH) for chromosomes 13, 16, 18, 21, 22, X and Y.

**All the women were investigated to exclude the known causes of implantation failures as:**

Anatomical (congenital and acquired anatomic abnormalities of the uterus and adnexa), endometriosis, infectious fac-

tors (*Cytomegalovirus*, *Herpes simplex*, *Mycoplasma hominis*, *Ureaplasma urealyticum*, *Chlamydia trachomatis*), inherited thrombophilias (mutations MTHFR, for homocysteine, factor V Leiden mutation, mutation 20210G for prothrombin and for antithrombin III, deficiencies in protein S and protein C), endocrinological disturbances (luteal phase insufficiency, polycystic ovary syndrome, insulin resistance, diabetes mellitus, hyperprolactinaemia and hyperandrogenism).

Receptivity of the endometrium was estimated before first ET by endometrial biopsy during the cycle before the IVF cycle.

Women with medical conditions or treatment affecting their immunologic responses were excluded from the study. All studied RIF and control group women were qualified for immunological tests which consisted of PBL population profile and the PBL Th1/Th2 cytokine secretion upon mitogen stimulation. All examinations were performed in non-pregnant women in the midsecretory phase.

10ml of heparinized peripheral blood was drawn from tested women using a Vacutainer sampling system (Beckton-Dickinson, Eastern Europe Division, Heidelberg, Germany).

Immunologic assays were done in APC Medical Analyses laboratory that participates in the diagnostic immunologic division in the research institute.

**Lymphocyte Populations**

The percentage of B lymphocytes, T lymphocytes populations and NK cells was measured in the whole heparinized peripheral blood using Multitest 6-color TBNK reagent (Beckton-Dickinson and Co., BD Biosciences, San Jose, USA) with standard monoclonal antibodies: CD3FITC, CD19APC, CD16+, CD56PE, CD45PerCP-Cy5.5, CD4Pe-Cy7, CD8APC-Cy7 using Facs Canto cytometer (Beckton-Dickinson and Co., BD Biosciences, San Jose, USA) and BD FACSCanto Software. The reference value were as follows: Lymphocytes T (60-82.0%), Lymphocytes B (7-23.0%), lymphocytes helper CD3+/CD4+ (30-51%), lymphocytes supresor CD3+/CD8+ (19-39%), CD3-/CD16+/56+ (NK) (7-24%), CD5+/CD19+ (2-10%), CD 56+ (3-12.0%).

**NK cells activity:** was measured by NKTEST (Glycotope, Biotechnology, Germany), using cryopreserved K562 target cells and found by flow cytometric analysis. The reference value of the NK cytotoxic activity after incubation time of 2 hours determined as normal range was 13.7-33.5%.

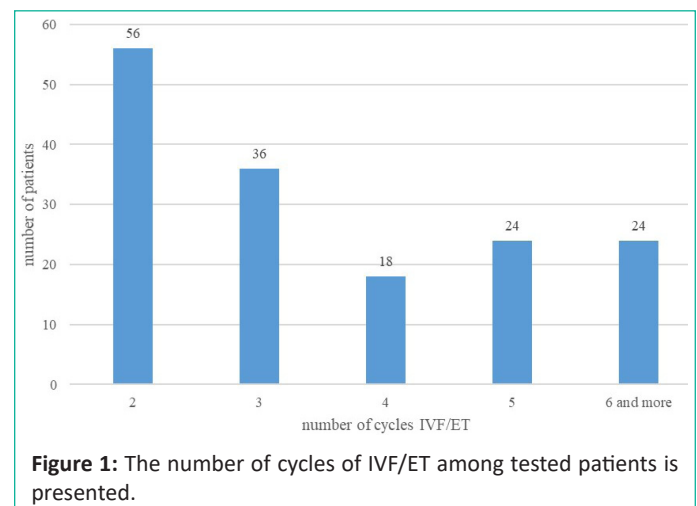
**PBL Th1/Th2 cytokine profile:** PBLs were isolated from heparinized whole blood by gradient separation. A suspension of 1x10<sup>6</sup> cells was cultured in Roswell Park Memorial Institute (RPMI) 1640 medium (enriched with 2mM l-glutamine, 10% fetal calf serum, 100U/mL penicillin, and 10µg/mL streptomycin). The lymphocytes were stimulated with phytohemagglutinin (at a mitogenic dose of 5mg/mL), and then incubated for 24h in humid air enriched with 5% CO<sub>2</sub> at 37°C. After centrifugation, the supernatant was collected for the cytokine assay. Cytokine levels were estimated using a BD™ Cytometric Bead Array Human Th1/Th2 Cytokine kit (Beckton-Dickinson and Co., BD Biosciences, San Jose, USA). The tests were performed on a FACSCanto™ cytometer (Beckton-Dickinson and Co., BD Biosciences). The concentrations of cytokines: Interferon [IFN]γ, Tumor Necrosis Factor [TNF]α, Interleukin [IL]-10, Interleukin [IL]-4, were determined and represented as pg/mL. The reference values, determined previously for fertile Polish women, were as follows: IFN-γ 209–1050pg/ml; TNFα 320–1380pg/ml; IL-10 1530–3830pg/ml; IL-4 20-120pg/ml.

**Statistical Analysis**

Median and two quartiles (first and third, Q1 and Q3, respectively) of PBL population's percentage were used as summary statistics.  $s_n$  Statistic was computed as the measure of variability:  $s_n = med\{med|x_i - x_j|, j = 1..n\}$  [1]. This is average distance between two randomly sampled observations among group. Higher  $s_n$  level reflects higher variability. If it was necessary distributions of statistics were estimated numerically. Risk of failure of IVF/ET, was modeling for cases ( $Y = 1$ ) and controls ( $Y = 0$ ) groups with model  $h[P(Y=1|X)] = \alpha + \beta^T X$ , where  $X$  is matrix of immunological predictors and  $h$  is *logit* function. Results were adjusted to age of patients. Odds Ratio (OR) was used as a measure of the effect size as well as the confidence interval, CI 95%, for this statistic at  $1 - \alpha = 0.95$  level [4].

**Results**

The mean age in the group of patients with RIF after IVF/ET was 36 years (Q1=33, Q3=39), in the control group was 34 years (Q1=31, Q3=37). The most dominant group (56 patients) consists of cases with 2 ET/IVF failures, the rest of tested patients underwent 3 to 6 and more cycles of ET/IVF procedures (from 18 to 36 patients).



**Figure 1:** The number of cycles of IVF/ET among tested patients is presented.

PBL populations percentage in RIF and control patients were estimated using median and two quartiles (first and third, Q1 and Q3, respectively) the measure of variability, showed in table 1.

**Table 1:** Characteristics of cases and controls analyzed in this study according to PBL percentage and cytotoxic NK activity.

Variable	Group	Q1	Median	Q3
Lymphocytes T	Cases	80.5	82	83.5
	Controls	73	76	78.5
Lymphocytes B	Cases	6	6	7
	Controls	10.5	13	14.5
Lymphocytes helper CD3+/CD4+	cases	38.5	39	44.5
	controls	32.0	34	36.5
Lymphocytes supresor CD3+/CD8+	cases	28.5	31	36
	controls	25	27	32
NK cells	Cases	16.7	20	24
	Controls	11	14	19.7
Lymphocytes CD5+/CD19+	cases	3.5	4	5.5
	controls	4	5	6
Lymphocytes CD 56+	cases	23	23	24.5
	controls	9.5	11	13
Activity of NKcells	Cases	19.6	28.2	37.5
	Controls	10	13.2	26.8

Q1, Q3 – first and third quartile

In the tested RIF's patients we found increased percentage of lymphocytes T, CD56+ and NK cells, comparing to the control group as well as enhanced NK cells activity. The percentage of lymphocytes B in RIF patients were decreased in significantly comparing to the control group. Compared to the reference values the percentage of NK cells was increased, however, insignificant in RIF's patients.

The mean age in the group of RIF patients was 36 years (Q1=33, Q3=39), in the control group was 34 years (Q1=31, Q3=37). The difference of 2 years is insignificant comparing the both groups. Taking into consideration the PBL percentage in patients with similar age we estimated the risk of RIF, presented in table 2.

**Table 2:** Risk of RIF after IV/ETF in dependence of the PBL percentage.

Variable	OR	CI 95%		p-value
Lymphocytes T increase at 1% point	1.077	1.010	1.149	0.01
Lymphocytes B increase at 1% point	0.472	0.298	0.748	0.01
Lymphocytes CD56 + increase at 1% point	1.119	1.027	1.219	0.01
NK cells activity increases at 1% point	1.144	1.060	1.236	0.01

OR: Odd Ratio; CI: Confidence Interval

In the tested group we observed that increase only of 1% of PBL T may influence on the risk of implantation failures almost for 8% (OR=1.077). Similarly we noted that increase of percentage of lymphocytes CD56+ in peripheral blood is associated with the rise of the risk of RIF as 12% (OD=1.119). Activity of NK cells over reference values is connected with enlarged risk of RIF as 15% (OR=1.144). Moreover, decrease of 1% of PBL B is supposed to be connected with twice decline of the risk of RIF in tested women (OR=0.472).

Mean concentrations of IFN $\gamma$ , TNF $\alpha$ , IL-10, IL-4 cytokines produced by mitogen-stimulated PBL and Th1/Th2 ratio (IFN $\gamma$ /IL10, TNF $\alpha$ /IL10) was measured in RIF and control patients.

$S_n$  – measure of variability: higher  $S_n$   $\rightarrow$  higher variability

**Table 3:** Th1/Th2 cytokine profile in patients with RIF versus successful pregnancy outcome after IVF/ET.

Parameter	RIF median (Q1-Q3)	$S_n$	Control median (Q1-Q3)	$S_n$
IFN $\gamma$	975 (770-1104.5)	230	683 (468.5-822)	290
TNF $\alpha$	1240 (1050-1821.5)	349	1020 (798.5-1185)	309
IL10	957 (600-1690)	655	967 (828.5-1595)	210
IFN $\gamma$ /IL10	0.97 (0.55-1.36)	0.54	0.71 (0.52-0.82)	0.26
TNF $\alpha$ /IL10	1.29 (0.72-2.06)	0.76	0.87 (0.52-1.36)	0.51

Q1, Q3 – first and third quartile

IFN $\gamma$ - interferon  $\gamma$ , TNF $\alpha$ -tumor necrosis  $\alpha$ , IL-10- interleukin 10,

The expression of IFN $\gamma$  and TNF $\alpha$  in PBL were found higher in patients with RIF than in the control group. The average concentration of IFN $\gamma$  among cases was 975pg/ml comparing to the controls with the average concentration 683pg/ml.

The levels of IL-10 didn't differ significantly between the two studied groups with the similar average of the concentration about 960pg/ml but 3-fold differentiation of the concentration of IL-10 was noted in the tested women with RIF. The aver-

age difference of concentration of IL-10 among patients with RIF was estimates as  $S_n$ =655pg/ml, in the controls  $S_n$ =210pg/ml. High expression of TNF $\alpha$  and IFN $\gamma$  in the tested group with the similar levels of IL-10 suggest abnormal ratios of IFN $\gamma$ /IL-10 amount 0.97 comparing to 0.71 among controls. Similar ratios of TNF $\alpha$ /IL-10 were elevated in women with RIF 1.29 comparing to the controls 0.87 for TNF $\alpha$ . This difference was not statistically significant.

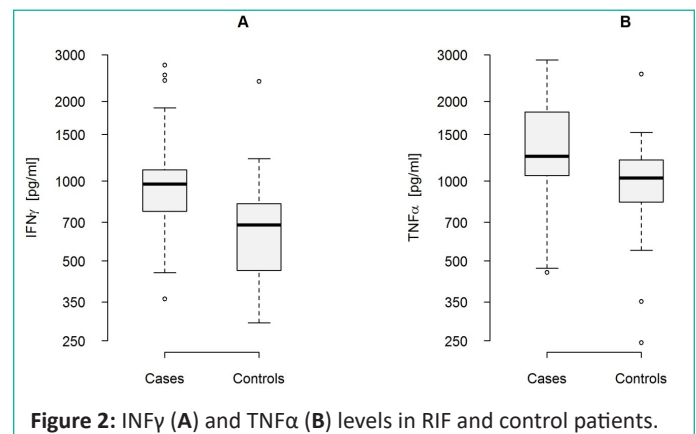
Comparing the Th1/Th2 cytokines profile in patients with same age we found increasing risk of RIF with increasing values of Th1 cytokines produced by PBL (Table 4).

**Table 4:** The risk of RIF after IVF/ERT in dependence of the cytokines levels adjusted to the same patients' age.

Variable	OR	CI 95		p-value
IFN $\gamma$ increases at 10%	1.13	0.998	1.28	0.03
TNF $\alpha$ increases at 10%	1.11	0.968	1.29	0.06

IFN $\gamma$ : Interferon  $\gamma$ ; TNF $\alpha$ : Tumor Necrosis  $\alpha$

In patients with same age we found that the risk of implantation failure increased as 11% (OR=1.11) with rising of expression of TNF $\alpha$  for 10%. Similar high expression of IFN $\gamma$  with 10% extension implicated over 2 fold increased risk of RIF (OR=1.28).



**Figure 2:** IFN $\gamma$  (A) and TNF $\alpha$  (B) levels in RIF and control patients.

We didn't noted that the change in levels of cytokine IL-10 and IL-4 is correlated with the successful implantation after IVF/ET ( $p=0.03$ ).

### Discussion

Several studies have showed modelled inflammation across pregnancy with immunomodulation of Th1/Th2 activity [3,5]. Maternal systemic inflammatory response could stimulate maternal-fetal interface unit to adverse implantation. Abnormal immune responses are significantly increased in women suffering from recurrent miscarriages [6,7]. Implantation failure after IVF procedures is still controversial with different definition of rising in the quantitative hCG level after embryo transfer [8]. Despite increasing number of gene mutations in the gene coding for the proteins responsible for thrombophilias, endometrial remodeling events indispensable to endometrial decidualization and the good quality of transferred embryos, peripheral blood immunological balance is independent required factor for the successful pregnancy outcome [1,9,10].

Our study revealed increased levels of Th1 cytokine TNF $\alpha$  and IFN $\gamma$  produced by PBL in women with RIF after IVF/ET procedures comparing to the controls. Strong correlation was observed in the elevated level of TNF $\alpha$ , similar as Zhang et al.[11] found in RM patients. Increased level of 10% level of TNF $\alpha$  with

normal levels of IFN $\gamma$  can in the same patients rise the risk of implantation failure as 11% (OR=1.11). The elevated ratio of TNF $\alpha$ /IL-10 and IFN $\gamma$ /1IL-10 also significantly influence on the risk of implantation failure ( $p<0.05$ ).

The average concentration of IL-10 didn't differ significantly between tested and controls but very wide range of concentration of IL-10 among sera of women with RIF indicates heterogeneity of peripheral blood immunological profile.

In women undergoing IVF/ET treatment, an increased count of activated not only uNK but PB NK cells (CD56 dim CD16+ CD69+) was reported to be associated with reduced rate of embryo implantation in IVF treatment [12]. The result of our study show the incidence of higher percentage of CD56+ and elevated activity of NK cells in RIF patients comparing to the control group. The increase of CD56+ levels elevates the risk of the unsuccessful implantation as 12% (OR=1.119).

Increased activity of NK cells found in RIF patients was observed as a factor provoking the risk of implantation failure estimated as 15% (OR=1.144).

Different immunological models found in patients suffering from RIF after IVF/ET procedures allow to use the treatment methods appropriate for identified disorders which gives the opportunity to increase the percentage of success [13].

### Conclusions

Our study revealed different alloimmune disorders found in the sera of patients with RIF after IVF/ET, in particular those concerning cytokine balance, lymphocyte profile and NK cells activity in peripheral blood.

The rate of immunological disorders found among patients undergoing the IVF/ET procedures reveals the necessity of performing certain immunological tests particularly among patients with at least 2 failures.

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