

LAB REPORT _

HOLLY WILKINSON (F) 10/10/2025

Patient Details
AGE (DOB):

ORDER NUMBER: B1B2K9K12 SAMPLE ID: 2509050006 **Specimen Details**

TYPE: PERIPHERAL BLOOD RECEIVED: 17/09/2025

CONDITION: ACCEPTABLE (GOOD)

Partner Details

NAME: LUKE WILKINSON

AGE (DOB): GENDER: MALE

FERTILYSIS REPRODUCTIVE IMMUNOLOGY			
NK cell profile/Regulatory T-Cell assay (Treg	gs) Result		Ref. Range/Normal
CD56+/CD16+ NK lymphocytes	10.53 %		<12.0
Regulatory T-Cells (Tregs)	0.30 % (L)		>0.8
NK cell cytotoxicity assay	Result	Intralipid Inhibition	Ref. Range/Normal
50:1	25.32 %	20.17%	<40 %
25:1	19.61 %	15.39 %	<30 %
12.5:1	15.10 %	8.64 %	<20 %
Th1/Th2 cytokine ratio assay	Result		Ref. Range/Normal
(CD3+CD4+) IFNy/IL-10	13.11 %		Normal <15> Increased <30> High
(CD3+C D4+) TNFα/IL-10	8.16 %		Normal <15> Increased <30> High
Leukocyte antibody detection (LAD) (crossmatch)		Result	Ref. Range/Normal
T cells (IgM)		Weak Positive (11.29%)	positive
T cells (IgG)			
T cells (IgG)		Negative (9.07%)	positive
_		Negative (9.07%) Weak Positive (14.49%)	positive positive
B cells (lgM) B cells (lgG)			· ·
B cells (IgM)	tive T or B cells will be reported.	Weak Positive (14.49%)	positive
B cells (IgM) B cells (IgG) *For results identified as positive, the percentage of posit	·	Weak Positive (14.49%)	positive
B cells (IgM) B cells (IgG) *For results identified as positive, the percentage of position (2-field) typing at	·	Weak Positive (14.49%) Negative (9.07%)	positive positive
B cells (IgM) B cells (IgG) *For results identified as positive, the percentage of positive HLA DQA-1 high resolution (2-field) typing at Allele 1	·	Weak Positive (14.49%) Negative (9.07%) Result DQA1*03:03 (No Match detected) DQA1*05:01 (No Match detected)	positive positive Ref. Range/Normal
B cells (IgM) B cells (IgG) *For results identified as positive, the percentage of positive HLA DQA-1 high resolution (2-field) typing at Allele 1 Allele 2	·	Weak Positive (14.49%) Negative (9.07%) Result DQA1*03:03 (No Match detected)	positive positive Ref. Range/Normal
B cells (IgM) B cells (IgG) *For results identified as positive, the percentage of positive. HLA DQA-1 high resolution (2-field) typing at Allele 1 Allele 2 Partner Allele 1	·	Weak Positive (14.49%) Negative (9.07%) Result DQA1*03:03 (No Match detected) DQA1*05:01 (No Match detected)	positive positive Ref. Range/Normal no matching no matching
B cells (IgM) B cells (IgG) *For results identified as positive, the percentage of position (2-field) typing at Allele 1 Allele 2 Partner Allele 1 Partner Allele 2	·	Weak Positive (14.49%) Negative (9.07%) Result DQA1*03:03 (No Match detected) DQA1*05:01 (No Match detected) DQA1*01:03 (No Match detected)	Ref. Range/Normal no matching no matching no matching
B cells (IgM) B cells (IgG) *For results identified as positive, the percentage of positive. HLA DQA-1 high resolution (2-field) typing at Allele 1 Allele 2 Partner Allele 1 Partner Allele 2 Maternal KIR typing Possible The following genotype is likely for 2DL1, 2DL	nalysis	Weak Positive (14.49%) Negative (9.07%) Result DQA1*03:03 (No Match detected) DQA1*05:01 (No Match detected) DQA1*01:03 (No Match detected) DQA1*03:01 (No Match detected) Result	Ref. Range/Normal no matching no matching no matching no matching
B cells (IgM) B cells (IgG) *For results identified as positive, the percentage of positive. HLA DQA-1 high resolution (2-field) typing at Allele 1 Allele 2 Partner Allele 1 Partner Allele 2 Maternal KIR typing Possible	nalysis genotype 3, 2DL4,2DS4, 3DL1, 3DL2, 3D	Weak Positive (14.49%) Negative (9.07%) Result DQA1*03:03 (No Match detected) DQA1*05:01 (No Match detected) DQA1*01:03 (No Match detected) DQA1*03:01 (No Match detected) Result	Ref. Range/Normal no matching no matching no matching no matching no matching
B cells (IgM) B cells (IgG) *For results identified as positive, the percentage of positive. HLA DQA-1 high resolution (2-field) typing at Allele 1 Allele 2 Partner Allele 1 Partner Allele 2 Maternal KIR typing Possible The following genotype is likely for the patient:	nalysis genotype 3, 2DL4,2DS4, 3DL1, 3DL2, 3D	Weak Positive (14.49%) Negative (9.07%) Result DQA1*03:03 (No Match detected) DQA1*05:01 (No Match detected) DQA1*01:03 (No Match detected) DQA1*03:01 (No Match detected) Result PL3, 2DP1, 3DP1 Genotype KIR AA	Ref. Range/Normal no matching no matching no matching no matching Ref. Range/Normal Ref. Range/Normal

INTERPRETATION OF RESULTS AND RECOMMENDATIONS: LOW TO MODERATE RISK

DIAGNOSIS: MILD IMMUNE TOLERANCE DEFICIENCY DETECTED

DESCRIPTION: The patient presents with **low levels of cytotoxic NK cells** and **low NK cell cytotoxicity**, both of which are **favorable**. The

Th1/Th2 cytokine ratio is also within favorable parameters, suggesting an immunologically receptive environment in this respect. Regulatory T cell (Treg) levels, however, are reduced, which may impair immune tolerance at the

maternal-fetal interface and has clinical significance.

The Leukocyte Antibody Detection (LAD) test was weakly positive, which while it can be considered as mildly favorable finding there is still room for improvement. A weakly positive LAD suggests partial alloimmune priming, which may still be enhanced further to better support maternal immune adaptation to the embryo. Furthermore, HLA-DQA1

Disclaimer: The information contained herein is intended for educational/informational purposes only. It does not constitute medical advice or counselling, it does not represent a medical examination by itself, nor it is a substitute for medical advice or medical treatment and should not be considered in any way as prescriptive. The results and the information in this test have not been approved by any official body to be used directly for diagnostic testing and the medical information contained herein is not intended as a substitute for conventional medical testing used for diagnosis and treatment. The information contained in this document is intended for use by a healthcare professional and the recipient of this report should always either consult with their treating physician, regarding medical diagnosis and treatment, or seek the advice of a qualified health care professional, concerning personal health and medical conditions. Each treatment must be prescribed by a qualified medical doctor, as medical treatments may have negative side effects. Upon receiving the results of the test, only a treating physician should decide whether treatment is required and which is the correct treatment for the patient, based on the patient's clinical history, additional medical results, known allergies and other relevant medical data. Further suggested diagnostic testing may only take place in a licensed medical facility by healthcare professionals and under circumstances approved by medical law. In any case, FERTILYSIS will not be liable for damages and claims due to improper use or misinterpretation by the party ordering the test (contracting party) both of the information contained in this report and of the company's services in general.

LAB REPORT

compatibility testing indicates no match, which is considered favorable and associated with reduced risk of alloimmune recognition.

Maternal KIR genotype is high risk (KIR AA), but this is counterbalanced by the fact that the paternal HLA-C genotype does not include HLA-C2 alleles. Therefore, while the maternal KIR profile would normally raise concern, the absence of paternal C2 moderates the overall risk.

TREATMENT:

The primary immunological concerns in this case are the **low Treg levels** and the **lack of enough protective LAD antibodies**. **Lymphocyte Immunization Therapy (LIT) is recommended** to boost maternal blocking antibodies, especially in light of the negative LAD result.

Each case should be assessed individually by an experienced reproductive immunologist or fertility specialist with expertise in immune-related infertility. These treatments aim to promote immune tolerance and improve the chances of successful implantation and pregnancy.

Note: Many of these treatments are still evolving, and research continues. It is essential for couples to consult with a qualified specialist to determine the best course of action based on their unique situation and the latest scientific evidence.

RETESTING:

No further immune testing is immediately required unless clinically indicated. However, a repeat LAD test may be recommended after LIT to confirm the development of blocking antibodies.

P.D. Venieratos Ph.D. Reproductive Biologist