Royal Hospital for Women (RHW) GUIDELINE COVER SHEET



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SUMMARY	Patients undergoing In Vitro Fertilisation (IVF) treatment, have an embryo transferred into their uterus, with the goal that the embryo implants into the lining of the uterus to achieve a pregnancy. The embryo(s) transferred to the patient can either originate from the cycle in which they were created (fresh cycle) or be frozen (cryopreserved) and thawed before transfer (thaw cycle). It is an Australia and New Zealand Assisted Reproduction Database (ANZARD) reporting requirement for an Assisted Reproductive Technology (ART) organisation including the Fertility Research Centre (FRC) to follow-up the outcome of all IVF treatments resulting in pregnancy and birth. This is a guideline regarding the follow-up of IVF pregnancy outcomes.		
Key Words	IVF, Pregnancy		



In Vitro Fertilisation (IVF) Pregnancy Outcomes

RHW GUID008

Contents

1	BAC	ACKGROUND3				
2	DEF	DEFINITIONS				
3	RESPONSIBILITIES					
	3.1	Medical, Midwifery, Nursing, Allied health	4			
4 PRO		OCEDURE	5			
	4.1	Clinical Practice	5			
	4.1.1	Determining Pregnancy Outcome:	5			
4.2 4.3 4.4		Documentation	5			
		Education Notes				
		Implementation, Communication, and Education Plan:	6			
	4.5	Related Policies/procedures	7			
	4.6	References	7			
5	ABC	RIGINAL HEALTH IMPACT STATEMENT DOCUMENTATION	7			
6	CUL	CULTURAL SUPPORT7				
7	NAT	NATIONAL STANDARDS				
8	REV	EVISION AND APPROVAL HISTORY8				
Α	ppendix	x A	8			
	8.1	Birth Outcome questions	۶			



In Vitro Fertilisation (IVF) Pregnancy Outcomes

RHW GUID008

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Within this document we will use the term woman, this is not to exclude those who give birth and do not identify as female. It is crucial to use the preferred language and terminology as described and guided by each individual person when providing care.

1 BACKGROUND

Patients undergoing In Vitro Fertilisation (IVF) treatment, have an embryo transferred into their uterus, the embryo implants into the lining of the uterus to achieve a pregnancy. The embryo(s) transferred to the patient can either originate from the cycle in which they were created (fresh cycle) or be frozen (cryopreserved) and thawed before transfer (thaw cycle). It is an Australia and New Zealand Assisted Reproduction Database (ANZARD) reporting requirement for an Assisted Reproductive Technology (ART) organisation including the FRC to follow-up the outcome of all IVF treatments resulting in pregnancy and birth. This is a guideline regarding the follow-up of IVF pregnancy outcomes.

2 DEFINITIONS

ANZARD	Australia and New Zealand Assisted Reproduction Database (ANZARD) is a				
	Clinical Quality Registry comprising information on all assisted reproductive				
	technology (ART) treatment cycles undertaken in Australian and New Zealand				
	fertility clinics.				
ARTEMIS	Computer program which stores data for all patients at FRC undergoing ART				
	procedures.				
ART	Assisted Reproductive Technology				
ART	an entity accountable for the delivery of services at one or more ART units				
organisation					
Biochemical	the absence of an identifiable pregnancy on ultrasound examination despite a				
Pregnancy	positive urine or blood β-hCG pregnancy test.				
Birth outcome	Including all livebirths, still births and neonatal deaths.				
Blighted Ovum	A condition that occurs when a gestational sac develops without an embryo (also				
	called anembryonic gestation).				
Clinical	A clinical pregnancy must fulfil at least one of the following criteria:				
Pregnancy	1. Pregnancy known to be ongoing at 20 weeks				
	2. Evidence by ultrasound of an intrauterine sac and/or foetal heart.				
	3. Examination of products of conception reveal chorionic villi				



In Vitro Fertilisation (IVF) Pregnancy Outcomes

RHW GUID008

	4. A definite ectopic pregnancy that has been diagnosed laparoscopically or by				
	ultrasound				
Ectopic	A complication of pregnancy in which the fertilised egg implants outside the uterus				
Pregnancy					
EDD	Estimated Due Date				
Embryo	Refers to the procedure where an embryo is transferred into the woman's uterus				
Transfer	using a thin catheter				
EPAS	Early Pregnancy Assessment Service				
FRC	Fertility and Research Centre				
IVF Treatment	Assisted reproduction technology involving the in vitro (outside of body) handling of				
	human oocytes (eggs) and sperm or embryos for the purposes of establishing a				
	pregnancy.				
Livebirth	A birth of an infant that meets the WHO definition and is 20 weeks or more				
	gestation or 400 grams or more in birthweight. WHO definition is: The complete				
	expulsion or extraction from its mother of a product of conception, irrespective of				
	the duration of the pregnancy, which, after such separation, breathes or shows any				
	other evidence of life, such as beating of the heart, pulsation of the umbilical co				
	definite movement of voluntary muscles, whether or not the umbilical cord ha				
	cut or the placenta is attached; each product of such a birth is considered livebor				
Missed	A missed (or silent) miscarriage is one where the baby has died or not developed				
miscarriage					
Neonatal death	h Is the death of a liveborn infant within 28 days of birth.				
Non-viable	· · · · · · · · · · · · · · · · · · ·				
pregnancy	detectable foetal heartbeat and/or abnormalities are present in the gestational sac.				
Patient	Woman undergoing IVF treatment				
Positive	Pregnancy tests check your urine or blood for a hormone called human chorionic				
Pregnancy	gonadotropin (β-hCG). This hormone is produced in the placenta of pregnant				
Test	women made during pregnancy. β-hCG is more than 100 mlU/ml				
Stillbirth	A stillbirth is the birth of an infant after 20 weeks or more gestation or 400 grams or				
	more birthweight that shows no signs of life.				
Viable	Refers to a pregnancy in which the foetus is developing intrauterine and has a				
pregnancy	detectable heartbeat				

3 RESPONSIBILITIES

3.1 Medical, Midwifery, Nursing, Allied health

3.2 Medical Director

Oversee all policy development and final approval of all clinical documentation in accordance with Reproductive Technology Accreditation Committee (RTAC) Guidelines



In Vitro Fertilisation (IVF) Pregnancy Outcomes

RHW GUID008

3.3 Medical staff

Development and management of individualised and comprehensive IVF treatment inclusive of monitoring of results and counselling

3.4 Registered Nurses/Midwives

Recognition, education and support to patients undertaking IVF treatment via superovulation inclusive of direct patient contact, triaging and escalation where appropriate

3.5 Allied Health

Counselling and emotional support particularly if there are poor outcomes

4 PROCEDURE

4.1 Clinical Practice

- Book the patient for a β-hCG test 10 days post embryo transfer to determine if a positive pregnancy test has been achieved. If positive (>100mlU/ml), then repeat in 1 week to check for appropriate rise in β-hCG (should double every 48-72hours). If <100mlU/ml, then repeat in 48 hours.
- Schedule patient for an ultrasound at 7 weeks of pregnancy to determine viability.
- Obtain a copy of the ultrasound report and place it in the patient's medical record.
- Review pregnancy status in ARTEMIS and update the record accordingly:
 - Pregnancy non-viable refer patient to Early Pregnancy Assessment Service (EPAS)
 - o Pregnancy viable advise the patient to visit her GP for ongoing care
- Document the estimated due date (EDD) to generate the patient's pregnancy status in ARTEMIS.

4.1.1 Determining Pregnancy Outcome:

- Access the pregnancy log monthly on ARTEMIS to check if the patient is due to give birth.
- Check the eMR for Birth Summary documentation if patient gave birth within SESLHD.
- Contact patient via phone or email (once they have given birth) to inquire about birth outcome (See Attachment 1)
- Record birth outcomes into ARTEMIS.

4.2 Documentation

- Antenatal yellow card
- Artemis Database



In Vitro Fertilisation (IVF) Pregnancy Outcomes

RHW GUID008

Artemis Documentation Instructions:				
To see which patients are due:				
Treatments Tab Select Pregnancy Review				
	Click "Refresh" Button			
Birth Outcomes				
Search	Select Patient			
Latest Cycle Tab	Click Pregnancy			
Enter	Delivery Date Delivery Method Gender Tick either Live birth, Still birth of NND (neo-natal death) Enter Baby Name Fill out congenital abnormalities if they exist. If they do not exist keep blank. Enter Hospital Details: If birth was at RHW tick RHW If birth was at another hospital tick Other			
Multiple Pregnancy?	On Left hand side Click "Add Baby" if multiple pregnancy to enter additional baby.			
Once Report Completed	Tick "Report/ Pregnancy" complete, ensure outcome is "Pregnant- Live Birth"			

• Appendix A: Birth Outcome questions

4.3 Education Notes

- The Australian and New Zealand Assisted Reproduction Database (ANZARD)
 collects information on assisted reproductive technology (ART) and donor sperm
 insemination (DI) treatments undertaken in Australian and New Zealand fertility
 clinics and the resulting pregnancy and birth outcomes.
- The data collected in ANZARD is used for a variety of purposes, including in the
 production of the Assisted Reproductive Technology in Australia and New Zealand
 annual report series, to monitor ART treatment practices, success rates and perinatal
 outcomes, to inform standards for accreditation of fertility clinics, and to provide
 feedback to clinics on their data compared to national standards.

4.4 Implementation, Communication, and Education Plan:

The guideline will be distributed to all medical, nursing and midwifery staff via @health email. The guideline will be discussed at ward meetings, education and patient quality and safety meetings. Education will occur through in-services, open forum and local ward implementation strategies to address changes to practice. The staff are asked to respond to an email or sign an audit sheet in their clinical area to acknowledge they have read and



In Vitro Fertilisation (IVF) Pregnancy Outcomes

RHW GUID008

understood the guideline. The guideline will be uploaded on to the intranet and staff are informed how to access it

4.5 Related Policies/procedures

- In Vitro Fertilisation (IVF) Management and Treatment of a Superovulation Cycle Clinical Business Rule
- In Vitro Fertilisation (IVF) -Embryo Transfer Clinical Business Rule

4.6 References

- Fertility Society of Australia and New Zealand. (2021). RTAC Scheme. Retrieved March 30, 2023, from https://www.fertilitysociety.com.au/wp-content/uploads/RTAC-Scheme-20-December-2021.pdf
- Zegers-Hochschild, F., Adamson, G. D., Dyer, S., Racowsky, C., De Mouzon, J., Sokol, R. Z., Rienzi, L., Sunde, A., Schmidt, L., Cooke, I. D., Simpson, J. L., & Van Der Poel, S. (2017). The International Glossary on Infertility and Fertility Care, 2017. Human Reproduction, 32(9), 1786–1801.
 https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5850297/
 https://doi.org/10.1093/humrep/dex234

5 ABORIGINAL HEALTH IMPACT STATEMENT DOCUMENTATION

- Considerations for culturally safe and appropriate care provision have been made in the development of this guideline and will be accounted for in its implementation.
- When clinical risks are identified for an Aboriginal and/or Torres Strait Islander woman or family, they may require additional supports. This may include Aboriginal health professionals such as Aboriginal Liaison Officers, health workers or other culturally specific services

6 CULTURAL SUPPORT

- For a Culturally and Linguistically Diverse CALD woman, notify the nominated crosscultural health worker during Monday to Friday business hours
- If the woman is from a non-English speaking background, call the interpreter service:

 NSW Ministry of Health Policy Directive PD2017 044-Interpreters Standard

 Procedures for Working with Health Care Interpreters.

7 NATIONAL STANDARDS

- Standard 1 Governance
- Standard 5 Comprehensive Care
- Standard 6 Communicating for Safety



In Vitro Fertilisation (IVF) Pregnancy Outcomes

RHW GUID008

8 REVISION AND APPROVAL HISTORY

Date	Revision No.	Author and Approval
August 2024	1	NUM Fertility and Research Centre
23.9.24	1	RHW BRGC

Appendix A

8.1 Birth Outcome questions

Baby Name:	
DOB:	
Gender:	Male - Female -
Delivery Method	Vaginal □ Caesarean □
Weight (grams)	
Hospital	RHW Other
Abnormalities (Record any known congenital abnormalities here)	