Accepted Manuscript

Evaluation of sheep ovarian tissue cryopreserved by slow freezing or vitrificaton after chick embryo chorioallantoic membrane transplantation

Mahboubeh Vatanparast, Mohammad Ali Khalili, Nahid Yari, Marjan Omidi, Mehdi Mohsenzadeh

PII: S0011-2240(17)30255-9

DOI: 10.1016/j.cryobiol.2018.01.002

Reference: YCRYO 3920

To appear in: *Cryobiology*

Received Date: 20 July 2017

Revised Date: 14 October 2017

Accepted Date: 9 January 2018

Please cite this article as: M. Vatanparast, M.A. Khalili, N. Yari, M. Omidi, M. Mohsenzadeh, Evaluation of sheep ovarian tissue cryopreserved by slow freezing or vitrificaton after chick embryo chorioallantoic membrane transplantation, *Cryobiology* (2018), doi: 10.1016/j.cryobiol.2018.01.002.

This is a PDF file of an unedited manuscript that has been accepted for publication. As a service to our customers we are providing this early version of the manuscript. The manuscript will undergo copyediting, typesetting, and review of the resulting proof before it is published in its final form. Please note that during the production process errors may be discovered which could affect the content, and all legal disclaimers that apply to the journal pertain.



- 1 Evaluation of sheep ovarian tissue cryopreserved by slow
- 2 freezing or vitrificaton after chick embryo chorioallantoic
- **3 membrane transplantation**
- 4
- 5 Mahboubeh Vatanparast^{1,2}, Mohammad Ali Khalili¹, Nahid Yari¹, Marjan
- 6 Omidi¹, Mehdi Mohsenzadeh¹

⁷ ¹Research and Clinical Center for Infertility, Yazd Reproductive Institute, Shahid

8 Sadoughi University of Medical Sciences, Yazd, Iran

9 ²Rafsanjan University of Medical Sciences, Rafsanjan, Iran.

10

- 11 Corresponding author:
- 12 Mohammad Ali Khalili, PhD
- 13 Email: <u>khalili59@hotmail.com</u>
- 14 Tel: 09133570876
- 15
- 16
- 17
- 18

19

20 Abstract

The aim of our investigations was to compare the effectiveness of two methods for cryopreservation of sheep ovarian tissue, slow freezing and vitrification. The quality of cryopreserved tissues was evaluated after 5 days of thawing and chorioallantoic membrane (CAM) transplantation. Follicular structure, stromal integrity and neovascularization were assessed. The areas of fibrosis and necrosis were measured using MICROVISIBLE software, and proliferation was assessed with Ki-67 immunostaning. Download English Version:

https://daneshyari.com/en/article/8464156

Download Persian Version:

https://daneshyari.com/article/8464156

Daneshyari.com