

3: EVALUATION OF DEVELOPMENTAL COMPETENCE OF ZONA PELLUCIDA-FREE MOUSE TWO-CELL EMBRYOS BY TIME-LAPSE MONITORING SYSTEM

LEE, JAEWANG¹; KIM, JIHYUN²; CHOI, YUN JUNG³; JUN, JIN HYUN⁴

¹Department of Biomedical Laboratory Science, Eulji University, Seongnam City, South Korea

²Department of Obstetrics and Gynecology, Bundang CHA Fertility Center, Seoul, South Korea

³Seoul Rachel Fertility Center, Seoul, South Korea ⁴Department of Biomedical Laboratory Sciences, Eulji University, Seongnam City, South Korea

Objective

The aim of this study was to clarify the role of the zona pellucida (ZP) in embryonic development of mice embryos, to investigate the effect of blastomeres alignment of ZP-free embryos on developmental competence, and to explore whether the time of compaction appearance correlates with preimplantation embryonic development and trophoblastic outgrowth potential.

Design

Animal experiment

Material and Methods

Mouse 2-cell embryos were collected on 1.5 days post coitum (dpc), and their ZP were removed using acid Tyrode's solution. Embryonic development from 2-cell to the outgrowth stage (on 1.5 to 7.5 dpc) was monitored using a time-lapse monitoring system.

Results

There was no difference in blastocyst formation between ZP-free and ZP-intact embryos. Pattern of blastomere alignment in ZP-free 4-cell embryos did not significantly affect blastocyst development and outgrowth potential. Interestingly, ZP-free 4-cell embryos with 5-point contacts in blastomeres showed significantly shorter mean cumulative time of development from 2-cell to blastocyst stage compared with those which had 4-point contacts. In addition, early compacted embryos before the 8-cell stage but with intact ZPs showed low developmental competence to blastocyst stage. This study has shown that the removal of the ZP does not affect blastocyst development and outgrowth potential in vitro in mice 2-cell embryos.

Conclusions

Collectively, we suggest that modification of ZP could apply various purposes without decreasing developmental competence in mice 2-cell embryos.

Support

This project was financially supported by a Korea Healthcare Technology R&D Project grant from the Ministry of Health and Welfare (A120043).