

# Effects of Early Embryo Cleavage on Embryo Quality and Pregnancy Outcome

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### **Abstract**

**Objective:** The aim of this study was to determine the embryo development rate, embryo quality and cleavage status of the Early Cleaved (EC) and NEC (No Early Cleavage) embryos and to analyze early embryo cleavage as a parameter for embryo selection process after Intracytoplasmic Sperm Injection (ICSI).

**Materials and Methods:** A total of 184 patients were included in the study. All embryos were checked for early cleavage at 25 hr to 27 hrs post ICSI. Embryo quality and embryo cleavage rates of the EC and NEC embryos were compared. For pregnancy assessment the patients were divided into two groups according to the presence of early cleaved embryos that were transferred. In the first group, all the embryos transferred were EC embryos and in the second group at least one of the embryos was NEC embryos.

**Results:** Normal embryo development rate (normal blastomere number at the time of embryo transfer) was found significantly higher in the EC embryos than in the NEC embryos (82.9% and 65.6% respectively) (p: 0.001). Embryo qualities were higher in the EC group. Pregnancy rates were significantly higher in the EC group (46.5% and 37.5% respectively).

**Conclusion:** EC embryos develop more normally, have better quality and have higher pregnancy rates when transferred hence should be used as an important parameter for embryo selection after ICSI

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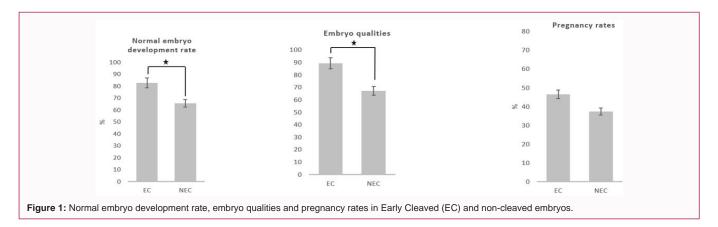
# Introduction

A useful and easy method to increase pregnancy and implantation rates by selecting the transferred embryo which has higher potential is crucial in assisted reproductive techniques [1-3]. The identification of most viable and most compotent embryos is important for Intracytoplasmic Sperm Injection technique (ICSI). Many scoring and grading criteria have been developed for selecting the best embryos for transfer. Oocyte morphology, size, pronucleus morphology, blastomere number and size, fragmentation rate, zona pellucida morphology and blastocyst morphology are the most used selection criteria for transfer [4-6]. Authors have shown that Early Cleavage (EC) of an embryo, defined as the first mitotic division happening 25 hr to 27 hrs after microinjection, is a strong indicator of competence of the developing embryo and embryo viability [5,7,8-20]. Many studies have reported EC is strongly correlated with embryo morphology [14-17], development rate to the blastocyst stage [18], chromosome abnormalities [19], embryo viability [8,12,20], implantation rate [15,21], and abortion rate [10]. Within the fertilized oocytes 5% to 38% of them were shown to cleave early [6,9].

The aim of this study was to determine the embryo development rate, embryo quality and cleavage status of the Early Cleaved (EC) and NEC (No Early Cleavage) embryos and to analyze early embryo cleavage as a parameter for embryo selection process after Intracytoplasmic Sperm Injection (ICSI).

## **Material and Methods**

A total of 84 patients were included in the study. All embryos were checked for early cleavage



at 25 hr to 27 hrs post ICSI. Embryo quality and embryo cleavage rates of the EC and NEC embryos were compared. For pregnancy assessment the patients were divided into two groups according to the presence of early cleaved embryos that were transferred. In the first group, all the embryos transferred was EC embryos and in the second group none of the embryos transferred were EC or a mixture of EC and NEC embryos.

#### Ovarian stimulation and ICSI procedure

Ovulation induction was performed using a short or long Gonadotrophin-Releasing Hormone (GnRH) analogue suppression protocol or a GnRH antagonist protocol using human menopausal gonadotrophins or recombinant Follicle-Stimulating Hormone (FSH). Oocyte-Cumulus Complexes (OCC) was recovered 36 hrs after the administration 10000 IU of human chorionic gonadotrophin. Denuding procedure were performed using enzymatic (Hyase 10X Vitrolife, Sweden) and mechanical techniques at the same time. The nuclear maturation of the oocytes was assessed with an inverted microscope (Nikon, Japan). Metaphase II oocytes with a polar body and metaphase I oocytes that extrude their polar body within 4 hrs after oocyte pick-up, were injected with a motile sperm. Microinjection was performed as described previously by Van Steirteghem et al. [25].

# Assessment of fertilization and embryo development

Further culture of injected oocytes was performed in 25 µl microdrops of culture medium (G-IVF, Vitrolife) under light paraffin oil (OVOIL, Vitrolife). Presence of fertilizations was checked 16 to 18 hours after OPU and described as fertilized if Two Pronuclei (2PN) and two polar bodies were observed in the ooplasm. Fertilization rates were calculated by dividing the number of fertilized oocytes with the total number of mature oocytes obtained. Fertilized oocytes were checked for embryonic development on day 2 and 3. Embryo development rates were calculated by dividing the number of embryos developed with the total number of mature oocytes. The cleavage status and the quality of the embryos were assessed by an inverted microscope (X40) according to the criteria of Staessen et al. [26]. A and B grade embryos were grouped as top quality embryos whereas grade C and D embryos were grouped as poor quality. Top quality embryo rates were calculated by dividing the number of top quality embryos with the total number of embryos. Mean rates for all the parameters were calculated by averaging rates calculated for each couple.

# Embryo transfer and pregnancy assessment

Embryo transfers were performed by the same clinician by the

guidance of ultrasonography on day 3. Wallace (Smiths, Kent, UK) catheter were used for embryo transfer. A  $\beta\text{-hCG}$  value of  $\geq 15$  mIU/mL twelve days after the embryo transfer indicated a positive pregnancy.

#### **Statistics**

The data were analysed using SPSS (Statistical Package for Social Sciences) software (SPSS Inc., Chicago, IL, USA) for Windows 10.0. We used Shapiro-Wilk's test to evaluate the distribution of the data. Mann-Whitney U-test was used for abnormally distributed variables and Student's-t-test for normally distributed variables. Chi-square test was used to compare the pregnancy rates between the groups. For all analyses, statistical significance was assessed at p<0.05.

#### Results

Normal embryo development rate (normal blastomere number at the time of embryo transfer) were found significantly higher in the EC embryos than in the NEC embryos (82.9% and 65.6% respectively) (p=0.001). Embryo qualities were better (top quality embryo rate) in the EC embryos (89.3% and 67.2% resp) (p=0.0042). Pregnancy rates were found significantly higher in the first group than the second group (46.5% and 37.5% respectively) (p=0.04) (Figure 1).

## **Discussion**

Many parameters may influence the outcome of ICSI but embryo selection is one of the most important factors that affect the outcome. Many parameters have been used to make this decision, including pronuclear morphology, blastomere morphology, blastocyst grading and choosing good most compotent and most viable embryos to increase implantation, pregnancy, and live birth rates is of great importance.

The use of early cleaved embryos to select the best embryos was first reported by Edwards et al. [27]. Until that time several studies have confirmed that EC is a strong indicator of embryo competency for selecting viable embryos [7-11,13,17,28,29]. Early cleavage time interval used in this study was 25 hr to 27 hrs after microinjection, which is the most frequently used time interval in most of the studies [23,30]. Many studies report the importance of EC to be used as embryo selection criteria before transfer and hence to help to avoid multiple pregnancies which is an important problem of assisted reproduction techniques. Many groups have reported an association with the quality of embryos and with rates of pregnancy and ongoing pregnancy when EC embryos were selected for embryo transfer [12,13,16,17,20,29]. Our findings confirm this study with an increased embryo quality in EC embryos than non EC embryos and

higher pregnancy rates when all the embryos that were transferred were EC embryos. We also confirm the study of Lee et al. [31] who reported that embryo quality was higher in early cleavage embryos than in non-early cleavage embryos. Also, Lee et al. [31] showed that early-cleavage embryos have a higher rate of normal development which is in accordance with our data. We have observed a higher rate of normal embryo development according to the NEC group in the EC group. In this regard, our results are similar to those of Lee et al. [31] results.

To increase pregnancy rates, implantation rates and to decrease multiple pregnancy rates, we conclude that early cleavage parameter should also be used to select best embryos for transfer. Further studies are needed to evaluate the behavior of most compotent embryos to decrease the number of embryos transferred and to reduce the risk of multiple pregnancies.

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