

In vitro maturation media, cysteamine concentration and glutathione level affect blastocysts development in mouse

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Received: 29 August 2007; accepted: 27 December 2007

Abstract

Background: Preparation of oocytes is one of the critical factors that determine the developmental competence of embryos produced by in vitro fertilization (IVF). Objective: In this study, the effect of cysteamine, type of media and glutathione (GSH) level on blastocysts development after in vitro maturation of mouse oocytes were investigated.

Materials and Methods: Premature female mice were primed with pregnant mare stimulating gonadotrophin (PMSG), and germinal vesicle (GV) stage oocytes were obtained 45 hr later. GV oocytes were cultured in presence of 0, 50, 100, 200 and 500 μ m cysteamine in TCM199 and MEME media. After IVM, MII oocytes were in vitro fertilized (IVF) and in vitro cultured (IVC) in order to observe embryo development. A group of In Vivo Ovulated (IVO) oocytes after priming with PMSG and HCG also were included in this study. 5,5-Dithio-bis (2nitrobenzoic acid) DTNB-recycling protocol was used for GSH assay.

Results: Rate of IVM and IVF were improved in all oocytes treated with cysteamine in the two medium except 500 μ m (81% MII rate in TCM and 64% MII in MEME). Rate of blastocyst in 100 μ m cysteamine in TCM199 and 200 μ m in MEME was higher compared to control groups (In TCM 45% and in MEME 35%). In vivo MII and GV oocytes represented the highest and lowest GSH level respectively.

Conclusion: Our results revealed that the media and concentration of cysteamine can affects on IVM, IVF and rate of blastocysts development on dose dependant manner.

Key words: *Cysteamine, IVM, Blastocysts, GSH, Oocyte, Mouse.*

Introduction

Preparation of oocytes is one of the critical factors that determine the developmental competence of embryos produced by in vitro fertilization (1). In vitro-matured (IVM) oocytes are known to have a lower developmental competence after IVF to in vivo-derived oocytes

(2). IVM oocytes have been used in some laboratories because their use makes it feasible to obtain a large number of oocytes from ovaries at relatively low cost. Many factors influence the karyoplasmic and cytoplasmic maturation of oocytes in vitro, including co-culturing with follicular cells such as cumulus cells or granulosa cells, duration of maturation, and type of media (3-8). However, the developmental ability of oocytes matured in defined media still tends to be lower than that of oocytes matured in media supplemented with chemicals such as amino acids or antioxidants (9, 10).

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TCM199 and MEME are two commercially available media used for human oocyte culture (11). For a great understanding of the effects of the different media and its supplements on IVM, subsequent embryo and fetal development, more studies are necessary. Cysteamine is a thiol compound that is known to be a scavenger of hydroxyl radical, and may contribute to maintaining the redox status in oocytes (12, 13). Addition of Cysteamine to a maturation medium increased GSH synthesis in bovine oocytes (14) and enhanced in vitro development of porcine embryos derived from intracytoplasmic sperm injection. It was shown that basic culture medium alone could influence rates of oocyte maturation (15) but addition of thiols compounds, like cysteamine to culture medium could have different effects depending on the concentration, the species and the type of oocyte (16). In this study, the effects of two culture media with different concentration of cysteamine on germinal vesicle breakdown (GVBD) and MII development, 2 cell embryo and blastocysts formation were investigated. Synthesis of GSH after administration of cysteamine to culture medium was measured and effects of its level on IVM, IVF and IVC rate was assessed.

Materials and methods

Superovulation

Oocytes were obtained from immature female ICR mice with age 4 weeks that were kept under controlled light and temperature conditions with free access to water and food. They had 12 hour light and 12 hour dark condition. Animal experiments were approved by the ethical committee of Tehran Medical University and performed with accordance with the university guidelines. Prepubertal ICR mice were primed with 5 IU of PMSG. The ovaries were removed 45 hr later and placed in TCM 199 or MEME according to subsequent culture.

IVM

Cumulus oocytes complexes (COCs) were retrieved directly from the follicles under a stereomicroscope by two 27 gauge needles. IVM medium consisted of MEME and TCM199 supplemented with 10% fetal Bovine serum, 75mIU FSH, 10ng/ml EGF (Epidermal growth factor, sigma) and different concentration of Cysteamine (Sigma, USA) 0, 50, 100, 200 and 500 μ M. the in total 15-20 oocytes were transferred to 50 μ l droplet covered with Mineral

oil and cultured in 37°C and 5 % CO₂. For IVM in each group 150 oocytes were cultured.

In vivo MII oocyte

In vivo ovulated (IVO) (17) oocytes were obtained from mice previously injected with 5 IU of PMSG to stimulate follicular development, followed 46 h later by 5IU HCG. MII oocytes were obtained from ovarian tube 16 h after HCG.

IVF

Epididymal sperm suspensions were prepared from adult mice and preincubated for 2 hr in IVF medium to ensure capacitation. IVF medium consisted of HTF medium was supplemented with 3% BSA fraction V. After IVM, MII oocytes were washed in IVF medium, and placed in 100 μ l micro-droplets under mineral oil, then 2×10^6 spermatozoa/ ml were added. Sperm and oocytes were incubated for 4 hours after which oocytes were removed, washed, and placed in 50 μ l micro-droplets of in vitro fertilization medium(HTF with 4mg/ml BSA). After 24 hours, numbers of two cells embryos were counted.

In vitro culture

Two cells embryos were washed in culture medium and cultured for subsequent development. In total 10-15 embryos were cultured in 50 μ l KSOM medium.

GSH assay

For GSH assay, Oocytes (germinal vesicle and MII stage) were washed three times in the stock buffer (0.2 M sodium phosphate buffer containing 10 mM EDTA, PH=7.2), and groups of 10-15 oocytes in 5 μ l of stock buffer were transferred to 1.5-ml microfuge tubes. Samples were stored at -80°C until assay. The intracellular concentration of GSH in oocytes was determined using 5, 5'-dithio - bis (2 - nitrobenzoic acid) - glutathione disulfide (DTNB-GSSG) reductase recycling assay (9). Briefly, 700 μ l of 0.33 mg/ml NADPH and 100 μ l of 6 mM DTNB in the stock buffer, and 190 μ l of distilled water were mixed in a microfuge tube. Then 10 microliters of 250 units/ml glutathione reductase (Sigma, USA) was added to initiate the reaction and the absorbance was monitored continuously at 412 nm using a spectrophotometer (Ziess, Japan) for 2 min and the quantity was determined from a standard curve which was constructed. The experiments were repeated 3 times with 10-15 oocytes for each group.

Statistical analysis

Statistical analyses of data from five replicate trials were carried out by INSTAT view.

Comparisons between groups of oocytes were performed using Chi-square and ANOVA with Tukey-Kramer Multiple Comparison Test as a post test. All values are expressed as Mean ± SD. A probability of $p < 0.05$ was considered to be statistically significant.

Results

IVM rate

Rates of GVBD and MII development are shown in Table 1. In all Cysteamine groups except 500 μM , rate of GVBD improved in two media. Rate of GVBD and MII in TCM199 and 100 μM cysteamine was higher than control group significantly ($p < 0.05$) (Table I). In MEME the development of GVBD and MII was higher in 200 μM and rate of MII was significantly higher than

control group ($p < 0.05$).

IVF and IVC rate

Two cells embryos and blastocysts development are shown in Table 1. Rate of two cell embryos was increased in the presence of cysteamine in a dose dependent manner. In the presence of 500 μM of cysteamine, rate of IVF was decreased more probably because of cytotoxicity effect. Rate of IVF was increased in TCM199 with 100 μM and in MEME with 200 μM cysteamine significantly ($P < 0.05$). Blastocysts development was increased in the presence of cysteamine in TCM199 significantly ($P < 0.05$). Rate of blastocysts in TCM199 with 100 μM was 45% and in MEME with 200 μM was 39%. Results showed that IVC rate in TCM199 was higher than MEME (Table I).

Table I. Percentage (%) of IVM, IVF and IVC in different groups in two media of TCM199 and MEME.

Rates Groups	GVBD rate		MII oocyte rate		Two cell embryos rate		Blastocysts	
	MEME	TCM199	MEME	TCM199	MEME	TCM199	MEME	TCM199
Control	62	79	54	60	40	49	30	32
50 μM Cys*	65	81	58	61	38	51	28	33
100 μM Cys	78	86	64	81	42	72	35	45
200 μM Cys	86	84	66	68	51	64	37	39
500 μM Cys	40	42	24	28	19	27	0	0

*- Cys: cysteamine

GSH analysis

GSH level was compared between oocytes cultured in the presence of cysteamine in two media and those matured in vivo. GSH level has been shown in different groups in two media in Figure 1, A and B. Intracytoplasmic GSH level in samples of oocytes obtained from mouse at oocyte collection time (0 h) and after IVM (24 hours cultured in TCM199) were analyzed. Also the level of glutathione in in-vivo MII oocytes was measured. Level of glutathione in In vivo oocytes was significantly higher than other groups (Figure

1A). The results indicate that the level of glutathione in MII oocyte was significantly higher in GV stage oocytes ($p < 0.001$). Also the level of Glutathione in 100 and 200 μM cysteamine were significantly higher than the control group in TCM199 and MEME respectively ($p < 0.05$).

In this experiment, for comparison of GSH synthesis in different IVM media, its level at oocyte collection time (0 h) and after IVM (24 hours cultured in MEME) were analyzed. Level of glutathione in in-vivo oocytes was significantly higher than other groups.

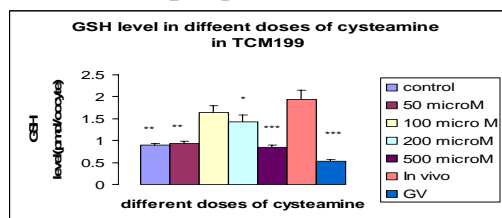


Figure 1(A): Level of GSH in different doses of cysteamine in TCM199.

*shows significance with $p < 0.05$

**shows significance with $p < 0.001$

***shows significance with $p < 0.0001$

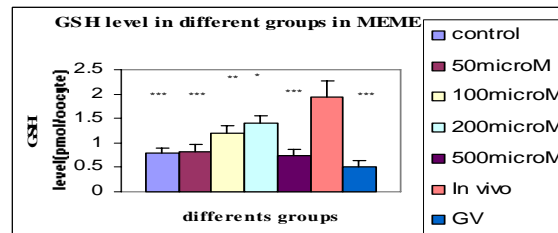


Figure 1(B): Level of GSH in different doses of cysteamine in MEME.

Discussion

The results of the present study revealed that cysteamine improved the rate of IVM, in vitro fertilization and blastocysts development in dose dependant manner. In this study, 100µm cysteamine in TCM199 and 200 µm in MEME had the highest rate of IVM and IVF and blastocysts development. Gasparinnini et al (2006) showed that 50 µm cysteamine increased rate of IVM and blastocysts in buffalo oocytes. In the study, administration of cysteamine increased the GSH synthesis (18). Difference species could be the possible reason between this result and our result. De Matos et al (2003) showed that 50, 100, 200 µm of cysteamine had no effects on IVM development but the rate of 2 cell embryo and blastocysts were increased (16). In our study, addition of cysteamine had dose dependant effects on IVM and IVC. It is noted, that they used different medium in their study (16).

Our results showed that GSH synthesis increased in the presence of cysteamine during IVM of mouse oocytes. Moreover, our study demonstrate that culture conditions stimulate a significant increase in oocyte GSH content during meiotic progression, from GV stage through MII in oocytes. The augmentation has beneficial effect on the rate of GVBD and MII developmental competencies of mouse oocytes.

In the present study, the addition of cysteamine not only improved the GSH content in MII oocytes but also had positive effect on the maturation rate of mouse Germinal vesicle oocytes. GSH is the major non-protein sulfhydryl compound in mammalian cells; it serves as a reservoir for cysteine and plays an important role in protecting the cell from oxidative damage. GSH content increases during development and oocyte maturation in the ovary, as the oocyte approaches the time of ovulation, and protects it in later stages of Fertilization (19). After fertilization, GSH participates in sperm decondensation in parallel to oocyte activation, and in the transformation of the fertilizing sperm head into the male pronucleus (20). Mercaptoethanol and cysteamine reduce cystine to cysteine and promote the uptake of cysteine enhancing glutathione (GSH) synthesis. Supplementation of mercaptoethanol, cysteine and cystine to IVM medium increased the intracellular GSH content of bovine oocytes and improved embryo development and quality, producing more embryos reaching the blastocyst stage on day 6 than embryos matured in unsupplemented medium (21). In vivo level of glutathione in MII oocytes

was significantly higher than that of in vitro condition in our study. in consistence with our finding, Zheng et al (1988) using hamster oocytes showed that the significant increase in GSH occurred during the transition from a nuclear or fibrillar GV to the time that chromosomes was condensed and spindle began to form (13). Luciano et al (2006) showed that in the presence of 100µm cysteamine the level of GSH was not increased in equine compared to control. Difference between our results and luciano might be because of species, different medium and contents of medium (22).

TCM199 improved GVBD and MII development better than MEME. Presence of 100µm cysteamine in TCM199 and 200µm in MEME had positive effects on IVM and IVF rate. In general terms, MEME is a less complex medium than TCM199. Both contain a similar range of inorganic salt, but MEME has a higher glucose concentration, fewer non-essential amino acids and a relatively small number of vitamins. TCM199 has a lower glucose concentration, both essential and non- essential amino-acids, a large range of vitamins and several other components such as cholesterol and ribose. Amino acids are known to be beneficial for embryo development, and oocyte maturation (11). Ruth et al (et al) showed that MEME had better effects on human MII development compared to TCM199. This result was not in consistent with the present study. Difference in type of oocyte (human oocyte) and absence of cysteamine in culture medium changed the results (11).

Conclusion

Overall, our result indicated that cysteamine plays a positive role in improvement of oocyte and embryo development. However, further studies are required to clarify its mechanisms.

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