Morphological markers of oocyte quality

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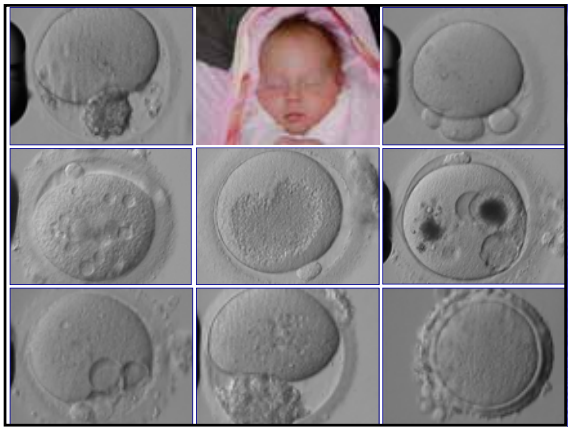
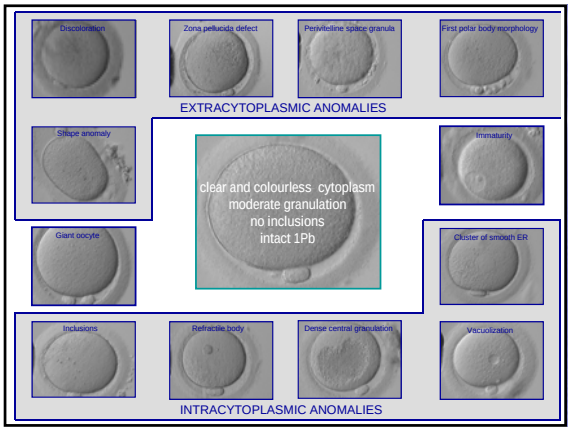
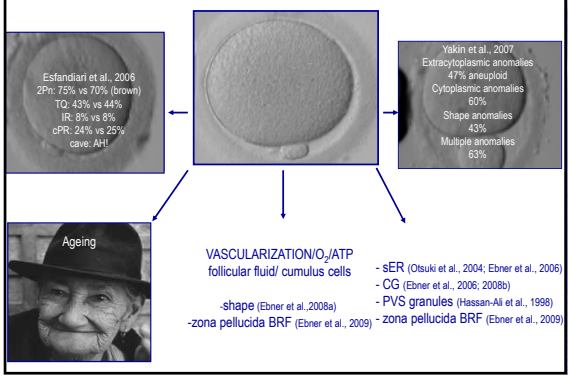
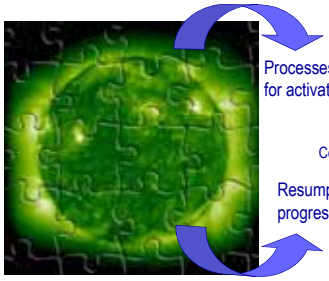




Possible influences on oocyte quality



Oocyte maturation



Processes that prepare the egg for activation and fertilization

Ca^{2+} release
Glutathione production
Competence for exocytosis

Resumption of meiosis and progression to metaphase II

Purines and cAMP
vs
MPF and Ca^{2+}/IP3

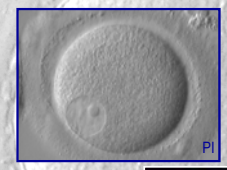
„IMMATURITY“
Desynchronization of nuclear and cytoplasmic maturation

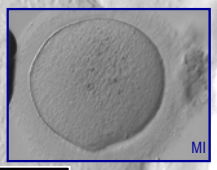
„OVERMATURITY“
Aged oocytes in vivo or in vitro

Nuclear maturity

Does nuclear maturity mean nuclear maturity?

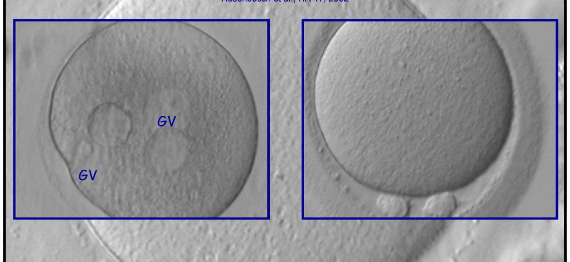
1. Nuclear maturity





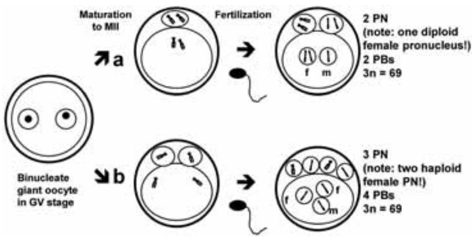
Giant oocyte

Balakier et al., HR 17, 2002
Rosenbusch et al., HR 17, 2002



Frequency approximately 0.3%
Mean diameter 200 µm (vs. 155 µm)
Contribution to digynic triploidy

Schematic representation of meiosis in giant oocytes



Rosenbusch et al., 2002

Oocyte diameter

- a critical oocyte size is necessary for resumption of meiosis (Otoi et al., 2000)
- size is determined by strong adhesion between oolemma and inner zona surface (Tartia et al., 2009)
- around ovulation GLYT1 is activated which mediates glycin accumulation which in turn acts as osmolyte and thus controls cell volume (Baltz and Tartia, 2009)

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TABLE 1

Comparison of the embryonic parameters of groups A, B, and C by the chi-square test.

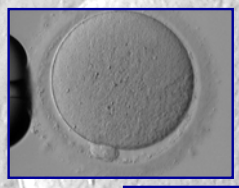
Day	Parameter	Group A	Group B	Group C	P value: A versus B versus C
1	Number of oocytes	40	60	40	
	Mean oocyte diameter (µm)	≤108.80[a]	108.80[a]–114.26[b]	≥114.26[b]	
	Fertilized	33 (82.5%)	55 (91.7%)	30 (75%)	[illegible]
2	≥4 cells	12 (36.7%)	21 (55.4%)	15 (50%)	[illegible]
	Good quality	16 (51.6%)	22 (42.3%)	10 (34.5%)	[illegible]
3	≥8 cells	12 (57.1%)	16 (53.3%)	24 (77.4%)	.117
	Good quality	10 (47.6%)	9 (21%)	11 (52.4%)	[illegible]

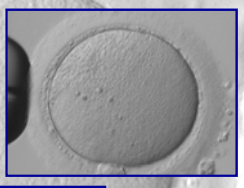
[a] 25th percentile.
[b] 75th percentile.

Romao et al., 2010

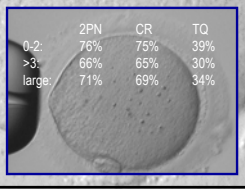
INTRACYTOPLASMIC ANOMALIES

Incorporations



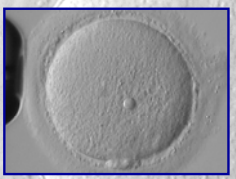


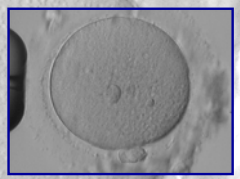
	2PN	CR	TQ
0-2:	76%	75%	39%
>3:	66%	65%	30%
large:	71%	69%	34%



Fancsovits et al., 2004

Refractile bodies



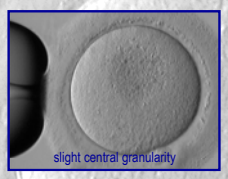


Viewed by transmitted electron microscopy, the refractile bodies showed the conventional morphology of lipofuscin inclusions and consisted of a mixture of lipids and dense granule materials

Larger lipofuscin inclusions (>5 µm) were associated with significantly reduced fertilization and unfavorable blastocyst development

Otsuki et al., 2007

Central granularity

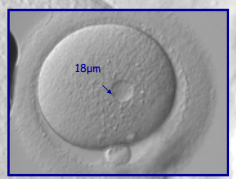


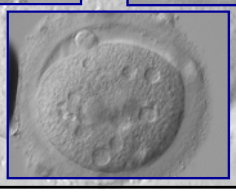
Relationship between central granularity and pregnancy outcome

- Anomaly was observed in 8% of the cycles (35% of the eggs were positive)
- Fertilization rate, embryo quality were inconspicuous
- Ongoing pregnancy rate was 12.8% (from slight form of CG), the implantaion rate 4.3%

Kahraman et al., 2000

Vacuolization





Formation of vacuoles

Vacuoles are membrane-bound cytoplasmical inclusions filled with fluid that is virtually identical with perivitelline fluid

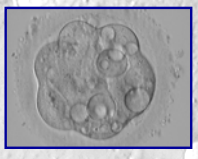
Vacuoles can arise spontaneously around extrusion of the first polar body
Van Blerkom, 1990

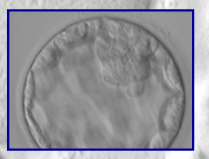
Vacuoles can form from preexisting vesicles derived from the ER or GA
El.Shafie et al., 2000

Vacuoles can be generated unintentionally by ICSI
Ebner et al., 2005

Occurence and developmental consequences of vacuoles

- 47 out of 1198 MII-oocytes showed at least one vacuole (3.9%)
- Fertilization rate was influenced negatively (48.9% vs 65.3%)
- A threshold was found above which fertilization did not occur (14 µm)
- Vacuolized oocytes had a blastocyst formation rate of only 12.5% compared to unaffected gametes (48.6%) (p<0.05)

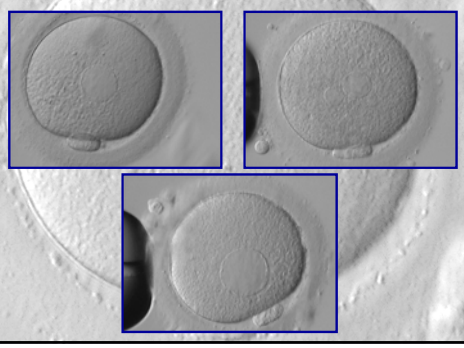




Ebner et al., 2005

EXCEPTION

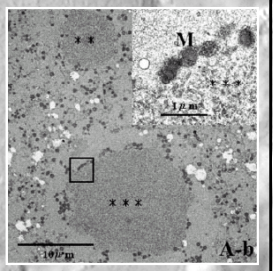
Aggregation of smooth endoplasmic reticulum



Relationship between sER clusters and outcome

Ebner et al., RBM 16, 2008; Otsuki et al., HR 19, 2004

- 6.2 to 9.4% of the cycles affected
- To our experience less than 2% of oocytes are affected (25% in pos cycles)
- Only MII oocytes
- Normal fertilization if rupture of sERC is avoided
- At lightmicroscopical level not all sERCs can be seen (2-9µm)!
- Blastocyst formation was 18%



Otsuki et al., 2004; Ebner et al., 2008

Relationship between sER clusters and outcome

- No relation to stimulation protocol, age, endometriosis but to E_2, dose of gonadothrophins, duration of COH
- sERC presence resulted in a desastrous outcome
  - IR, PR — no difference
  - Biochemical pregnancies — 58% vs 22% (P<0.01)
  - Take-home baby rate — 42% vs 78.% (p<0.001)
  - Increase in obstetric problems (33% vs. 5%) and lower birth weight (2500g vs. 3100g)
  - 2% stillbirths (not to forget one Beckwith-Wiedemann syndrome in the Otsuki paper)

Ebner et al., 2008

CASE REPORT

Smooth endoplasmic reticulum aggregations in all retrieved oocytes causing recurrent multiple anomalies: case report

At 36 weeks' gestation the patient was delivered by cesarean section. The male newborn was depressed at birth and was taken to the neonatal intensive care unit. Follow-up evaluation of the baby showed patent ductus arteriosus, dysgenesis of the cerebral sulcus, variation of septum pellucidum, and closed external meatus of the right ear. Moreover, ultrasonography showed infantile polycystic kidney. Neonatal infection developed and antibiotic supplementation was begun. Days later, respiratory depression required mechanic ventilation. On postnatal day 14 the baby died of cardiopulmonary arrest.

OOCYTE AGING

IN VIVO or IN VITRO acquired cellular, molecular, biochemical, morphological and epigenetic changes of gametes closely associated with poor developmental potential and recuced fecundity

- Decreased Fertilization
  - Polyspermy
  - Digyny
  - Parthenogenesis
- Chromosomal disorders
- Retarded development

Aged oocytes: morphological changes

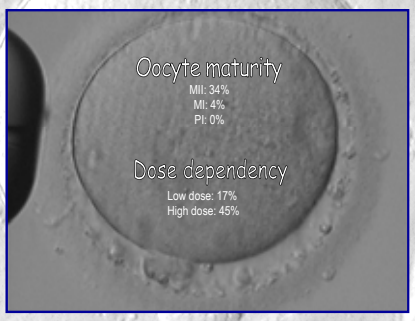
Table 1 Changes in morphology and cell biology during mammalian oocyte aging

	Fresh oocytes	Aged oocytes	Reference
PM	Microvilli extensions display intact structure	Microvilli extensions display structural alterations and are budded off into the PVS	Kim et al. (1996); Longo (1974); Pickering et al. (1988); Szollosi (1971); Webb et al. (1986)
Zona pellucida	Zona pellucida appears as a granulofibrillar, interconnected reticulum with pores	Zona pellucida displays a 'cobblestone' appearance and becomes harder	Goud et al. (2005b); Longo (1981); Miao et al. (2005); Xu et al. (1997)
PVS	Small	Large	Miao et al. (2004)
CG	CGs are densely populated in a line just beneath the oolemma, with a typical normal CG-free domain above the meiotic apparatus	CGs undergo migration and partial exocytosis	Dodson et al. (1989); Goud et al. (2005b); Gulyas (1979); Longo (1974); Szollosi (1971); Xu et al. (1997)
Microfilament	A thick microfilament domain exists in the oocyte cortex	Disrupted or lost	Kim et al. (1996)
Spindle	Spindles display vertical orientation to the oolemma and each pole is associated with a ring of centrosome proteins	Spindles become elongated and/or smaller and few microtubular foci are detectable at the cortex	Eichenlaub-Ritter et al. (1986); Eichenlaub-Ritter et al. (1988); Goud et al. (2004); Longo (1974); Meyer and Longo (1979); Segers et al. (2008); Soewarto et al. (1995); Wang et al. (2001)
Chromosomes	Chromosomes are intact and arranged symmetrically on the metaphase plate	Chromosomes display PCS and are scattered throughout the degenerating spindle and some chromosomes show centripetal migration, dispersion, decondensation and formation of a single chromatin mass	Eichenlaub-Ritter et al. (1988); Mailhes et al. (1998); Rodman (1971); Steuerwald et al. (2005); Szollosi (1971); Van Wissen et al. (1991); Zenzes and Casper (1992)
Mitochondria	Mitochondria are intact	Membrane potential decrease and mitochondrial matrix swell	Wilding et al. (2001)
PBI	PBI is intact and adjacent to the MII spindle	PBI degenerates and deviates from the MII spindle	Miao et al. (2004)

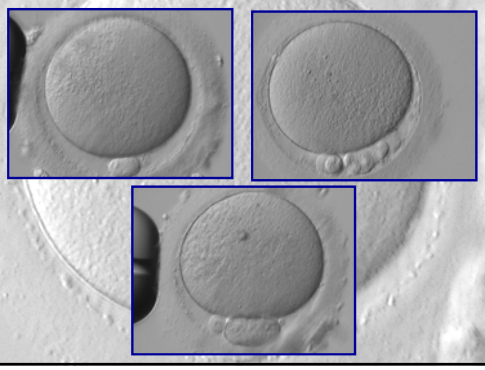
Miao et al., 2009

Perivitelline space granularity

Hassan-Ali, HR 13, 1998



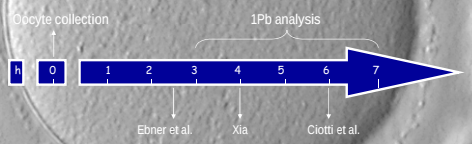
First polar body morphology



First polar body and outcome

Xia, HR 12, 1997
Ebner et al., F&St 1999, HR 2000, HR 2002
Ciotti et al., HR 19, 2004

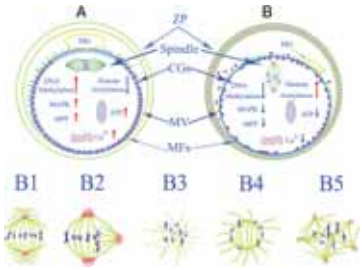
Since first polar body has a short half-life time it is thought to be an indicator of postovulatory age of the oocyte



Morphology of polar body has nothing to do with chromosomal situation of the gamete

Aged oocytes:

cellular, biochemical and epigenetic aspects



Miao et al., 2009

Factors affecting oocyte aging

Table II Effects of various environmental factors on oocyte aging

Aging environment	Effects	References
Temperature	Fertilization of room temperature aged (27°C) oocytes results in mouse full-term births. Oocytes aged in a refrigerator (4°C) or incubator (37°C) boost the developmental potential	Lin et al. (2008c); Lin et al. (2008b); Wakayama et al. (2004)
In vivo and in vitro	NOD: similar morphological alterations and cytoskeletal organization	Longo (1980); Miao et al. (2005); Webb et al. (1984)
	YFP: oocytes aged in vivo display a larger spindle and microtubule asters. Spindles in oocytes aged in vitro are close to the PM and display different orientations. In vitro culture retards oocyte aging	Ahlhuis et al. (1998); Adonis et al. (1997)
ICC	Accelerate oocyte aging by secreting a soluble APF into the medium	Miao et al. (2005); Chao et al. (2008)
ROS	Superoxide induces oocyte zona pellucida hardening, cytoplasmic microtubule dynamics increase and major CGs losses. H_2O_2 renders fresh oocytes resistant to aging but enhances the further aging in aged oocytes. Low levels of H_2O_2 induce the aging of fresh and aged oocytes, while higher concentrations of HOO compromise oocyte viability	Goud et al. (2008)

Miao et al., 2009

Conclusion

- The developmental fate of an oocyte is strongly dependant on the quality of the follicle (O_2, apoptosis)
- Controlled ovarian hyperstimulation recruits follicles of different qualities
- Either nuclear or cytoplasmic maturation may be affected both of which can influence oocyte morphology
- Oocyte aging is underestimated
- Potential negative predictors are aggregation of sER, vacuolization, dense central granulation and undetectable meiotic spindles

Conclusion II

In the context of oocyte morphology and outcome Van Blerkom and Henry (1992) suggested a interesting hypothesis

DYSMORPHISM

EARLY in maturation ⟷ LATE in maturation

aneuploidies

GIANT OOCYTES
CENTRAL GRANULATION
MEIOTIC SPINDLE

developmental failure

AGGREGATION of sER
VACUOLIZATION

Thanks for your

attention!