



BIostatISTICS MCQ

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Principles & Applications

# Mcq On Meiotic Error

**Yi-Tong Ma**



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Bibliography of Agriculture ,1974

**Mechanisms that Prevent and Correct Trisomy** Daniel Benjamin Cortes,2015 Female meiosis in most metazoans facilitates the creating of a single haploid oocyte from an original diploid precursor with replicated chromosomes In order to promote accurate



retention of only a single set of each homologous chromosome female meiosis orchestrates a complex series of events that result in the placement of a small acentrosomal meiotic spindle in a perpendicular orientation adjacent to the cortex in both meiosis I and meiosis II By placing the spindle perpendicular to the cortex along its major axis prior to anaphase oocytes ensure that each division retains a single copy of each homolog in meiosis I or sister chromatid in meiosis II Defects in the processes of meiosis can result in chromosome missegregation which induces aneuploidy One specific form of aneuploidy wherein three copies of each chromosome are present trisomy is of special importance due to its prevalence in human infertility and due to the presence of viable trisomic humans with developmental disorders Down Syndrome Klinefelter etc In humans trisomy is believed to arise mostly through errors in female meiosis highlighting the error prone nature of female meiosis Several mechanisms known to promote trisomy occur during female meiosis and in the mitotic divisions thereafter usually as a result of meiotic errors Herein we highlighted some of these known mechanisms and their consequences in human mouse and *C elegans* Despite the breadth of knowledge about defects of female meiosis and early embryonic development however little is known about any mechanisms that might rescue trisomy in vivo Case studies of triploX and Down Syndrome reproductive humans show fewer than expected trisomic progeny based on traditional Mendelian segregation Using triploX strains of the nematode *Caenorhabditis elegans* as well as some mutants with unpaired chromosomes we investigated whether female meiosis and the sequential mitotic divisions bear mechanisms that reduce the incidence of trisomy or more generally the incidence of unpaired chromosome inheritance In this work we show evidence that three distinct mechanisms act through meiosis and early mitosis in *C elegans* embryos to selectively expel unpaired chromosomes First we observed a measurable reduction in inheritance of unpaired chromosomes occurring during anaphase of meiosis I that is dependent on the active placement of the meiotic contractile ring Next we provide evidence for a mechanisms that relied on the rotation of meiotic spindles from their metaphase orientation parallel to the cortex into their anaphase orientation perpendicular to the cortex In that work we ASPM a marker of spindle poles as being consistently asymmetric on spindle poles in a way that predicted spindle orientation following rotation Unpaired chromosomes in a triploX strain frequently associated with an ASPM enriched pole which ended up at the cortex after rotation Finally we provide preliminary evidence for a mechanisms that provides further correction after meiosis has completed The incidence of trisomy was further reduced by mitotic correction in the first few divisions of embryonic development producing a population of mosaic embryos which were partially returned to disomy Together these data support a model where trisomy is detected by meiotic and mitotic embryos and a series of mechanisms activate in order to reduce the chances of excess chromosome inheritance

**Recombination and Meiosis** Richard Egel, Dirk-Henner Lankenau, 2008-07-25 Once per life cycle mitotic nuclear divisions are replaced by meiosis I and II reducing chromosome number from the diploid level to a haploid genome and recombining chromosome arms by crossing over In animals all this happens during formation of eggs and sperm in

yeasts before spore formation The mechanisms of reciprocal exchange at crossover chiasma sites are central to mainstream meiosis To initiate the meiotic exchange of DNA surgical cuts are made as a form of calculated damage that subsequently is repaired by homologous recombination These key events are accompanied by ancillary provisions at the level of chromatin organization sister chromatid cohesion and differential centromere connectivity Great progress has been made in recent years in our understanding of these mechanisms Questions still open primarily concern the placement of and mutual coordination between neighboring crossover events Of overlapping significance this book features two comprehensive treatises of enzymes involved in meiotic recombination as well as the historical conceptualization of meiotic phenomena from genetical experiments More specifically these mechanisms are addressed in yeasts as unicellular model eukaryotes Furthermore evolutionary subjects related to meiosis are treated

## Reviewing **Mcq On Meiotic Error**: Unlocking the Spellbinding Force of Linguistics

In a fast-paced world fueled by information and interconnectivity, the spellbinding force of linguistics has acquired newfound prominence. Its capacity to evoke emotions, stimulate contemplation, and stimulate metamorphosis is really astonishing. Within the pages of "**Mcq On Meiotic Error**," an enthralling opus penned by a highly acclaimed wordsmith, readers set about an immersive expedition to unravel the intricate significance of language and its indelible imprint on our lives. Throughout this assessment, we shall delve to the book is central motifs, appraise its distinctive narrative style, and gauge its overarching influence on the minds of its readers.

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