

Uninterrupted Culture Media Complete Recommended Protocol

Continuous Single Culture-NX Complete (CSCM-NXC) and Continuous Single Culture Complete (CSCM-C)

4/5-Well Dish

CSCM-NXC and CSCM-C are optimized to be used in an uninterrupted culture system without medium renewal or dish change on Day 3 when using 500 μ L of CSCM-NXC and CSCM-C Medium for embryo culture.

MEDIUM PREPARATION

CSCM-NXC and CSCM-C are complete, ready-to-use medium containing Human Serum Albumin (HSA) as the protein component. It is not necessary to add any other protein before use.¹

DISH PREPARATION


A sufficient volume of CSCM-NXC or CSCM-C is required so that oocyte collection, insemination, and embryo culture dishes can be prepared.

Oocyte Collection and Insemination Dishes (60 x 15 mm):

On the day before oocyte retrieval (Day -1) prepare dishes with eight 100 μ L volume drops of pre-supplemented uninterrupted culture medium along the perimeter of the dish with an 11 mL volume oil overlay.

Embryo Culture Dishes (4/5-well culture dish): On the day of oocyte collection (Day 0), prepare 4/5-well dishes by aseptically dispensing 500 μ L volume of pre-supplemented uninterrupted culture media and 400 μ L volume oil overlay in each well. If using a 5-well dish, the medium in the center well is used for rinsing the embryos prior to placing them into the remaining 4-wells for extended culture.

Time-lapse Surveillance System: CSCM-NXC and CSCM-C work effectively in all time-lapse surveillance systems and should be used accordingly to the individual system's protocol.


 To ensure proper temperature and pH of medium and oil, all dishes should be pre-equilibrated overnight to 37°C in a 5–6% CO₂ incubator on the day before oocyte collection (Day -1) and on the day of oocyte collection (Day 0). Minimum incubation time should be no less than 6 hours.

CO₂ AND PH

CSCM-NXC and CSCM-C can be used in ambient oxygen incubators but we recommend using reduced oxygen when possible. Following CO₂ equilibration, the pH of the uninterrupted culture medium should be within the acceptable pH range of 7.25–7.40 with a targeted pH of ~7.26–7.35.

Tri-gas Incubator: Ideally, we recommend 6% CO₂, 6% O₂, and 88% N₂ for optimal fertilization and embryo growth.

CO₂ Incubator: Alternatively, we recommend a 5–6% CO₂ for sufficient fertilization and embryo growth.

 pH levels are affected by the altitude above sea level (elevation), as the elevation increases the gas pressure over the culture medium decreases requiring a higher CO₂ pressure to achieve the optimal pH for the uninterrupted culture medium (7.26–7.35). To adjust pH, increase CO₂ levels to lower pH and decrease CO₂ levels to raise pH to fall within the specified range. pH should always be verified with pH measurements using a pH meter.

OOCYTE RECOVERY

Immediately upon oocyte collection and identification, it is recommended that oocytes be placed into the pre-equilibrated oocyte collection dish (prepared on Day -1). Following the trimming of the cumulus coronal complex from each oocyte, place 1–3 oocytes into a fresh drop of the oocyte collection dish and return to the controlled gas incubator at 37°C for the desired recovery period. Oocytes should be incubated for a minimum of 1 hour prior to insemination or denuding if ICSI is to be performed. Oocytes should remain in the collection dish for up to 4 hours prior to conventional IVF or insemination by Intracytoplasmic Sperm Injection (ICSI).

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INSEMINATION OF OOCYTES FOR CONVENTIONAL IVF

Upon oocyte recovery, trimming of cumulus complex and incubation for up to 4 hours (minimum 1 hour), remove the oocytes from the collection dish and transfer the oocytes to the pre-supplemented uninterrupted culture medium droplets in the pre-equilibrate insemination dish (prepared on Day -1). It is recommended to aseptically dispense 50,000–100,000/mL motile sperm per micro droplet containing 1–3 oocytes. Return the insemination dish to the incubator and check for normal fertilization 16–20 hours post insemination.

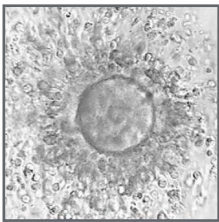
ICSI

Following >1 hour post oocyte denuding, remove denuded oocytes from incubator and inseminate with sperm as per standard ICSI protocol. Immediately following insemination, place 1–3 inseminated oocytes into a fresh drop of the pre-supplemented uninterrupted culture medium in the pre-equilibrated insemination dish (prepared on Day -1), and return dish to the incubator and check for normal fertilization 16–20 hours post insemination.

EMBRYO CULTURE POST FERTILIZATION

Following fertilization assessments with the identification of the presence of normal fertilization (two pronuclei and two polar bodies), transfer 2PN zygotes into the pre-equilibrated 4/5-well CSCM-NXC or CSCM-C culture dish previously prepared on Day 0. We recommend embryos to be group cultured (maximum of 5 embryos per well) in a static, uninterrupted culture system, without dish change or medium renewal on Day 3, until the desired stage of embryo development is achieved.

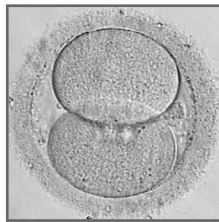
DAY 0
Oocyte



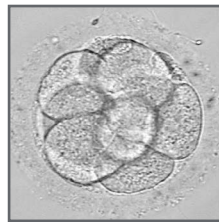
DAY 1
2PN



DAY 2
2-4 Cells



DAY 3
8 Cells



DAY 4
Compacting-Cavitating
Cells



DAY 5/6
Blastocyst Stage



1. FUJIFILM Irvine Scientific has not validated these procedures and each laboratory should consult its own laboratory procedures and protocols which have been specifically developed and optimized for your individual medical program.

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