



VONESTEP CULTURE MEDIUM

VITROMEDia

VITROMED GmbH

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VONESTEP

CULTURE MEDIUM



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USE

^vONESTEP Culture Medium was developed for in vitro procedures involving the fertilization and culture of human embryos from the 1cell stage to the blastocyst stage.

COMPONENTS

Sodium chloride Potassium chloride Magnesium sulphate Potassium phosphate Sodium bicarbonate L-Glutamic acid Calcium lactate Sodium pyruvate Calcium-D-pantothenate Alanyl-glutamine L-Phenylalanine L-Aspartic acid Glycine L-Asparagine L-Tyrosine L-Proline L-Serine L-Arginine L-Cystine L-Histidine L-Isoleucine L-Leucine L-Lysine L-Methionine L-Threonine L-Trytophan L-Valine EDTA Glucose Sodium Citrate Hyaluronate Phenol red Gentamicin HSA

QC Tests

Sterility sterile (SAL10-3)

Endotoxins <0.25 EU/ml

Human Sperm Survival Assay

Mouse Embryo Assay (blastocysts after 120h) ≥ 80%

Time Lapse Embryo Imaging with morphokinetics and cell count

Schedule of dish preparation and procedures for use of ^vONESTEP Culture Medium

9	Name of Product REF VONESTEP V-O	SM-20	Unit Size 20 ml GLASS Bottle
	 VONESTEP Medium can I development from Day 0 to For Embryo Culture we rea culture (up to 3 - 4 Embr Embryo Culture 	pe used to culture o day 6 commend 30 – 50 μ yos per drop) or 10	e Embryos at all stages of I drops of medium for group) – 15 µL per drop for single
рН	After equilibration with 6% CO2 Medium should be in a pH rang	2 for a minimum of 4 ge of 7.20 - 7.40 with	hours or overnight, ^v ONESTEP a desired pH of 7.26 - 7.35
DAY -1	 Prepare ^vONESTEP Medium them Prepare V-HEPES plus me 	n dishes at the day edium in a 37°C wa	before use and preincubate armer for Oocyte / Cumulus
DAY O	 For conventional IVF, insem incubate overnight For ICSI remove the cumulu VDENUPET After ICSI transfer injected of For conventional IVF prepa 	inate oocytes in VON Is mass from the ooc Docytes directly to V(Ire VONESTEP Mediur	NESTEP Medium and sytes in ^v HYLASE using DNESTEP Medium dishes n dishes for use on Day 1
DAY 1	 Assess fertilization of oocyte For IVF remove oocytes fro them in fresh equilibrated ^v To minimize out-gassing of return the culture dish to the 	es treated by IVF or 10 m the cumulus cells ONESTEP Medium dia CO ₂ and a drift in the e incubator within 2	CSI using VDENUPET and transfer shes prepared on Day 0 he pH of the medium quickly - 3 minutes
DAY 2	 Embryo assessment optional excessively stressed by the Embryo Transfer and cryo individual lab protocol 	al but not recommer removal from incubo opreservation are o	nded so that Embryos are not ator for assessment optional depending on the
DAY 3	Assess Embryos for cleavage a within 2-3 minutes for continued Embryo Transfer and cryoprese lab protocol	nd quickly return the d culture AND / OR ervation are optional	e culture dish to the incubator depending on the individual
DAY 4	Embryo Assessment AND / OR Embryo Transfer and cryoprese	rvation optional dep	pending on the individual lab
DAY 5	Assess Embryo morphology and Cryopreserve Blastocysts	d do Embryo Transfei	r AND / OR
	VITR	ON	AED ia



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