

RapidWarm™ Blast



US

en: Indication for use

Media for warming of vitrified human blastocyst stage embryos.

Contraindications

RapidWarm™ Blast contains gentamicin. Do not use in patients with known hypersensitivity/allergy to the component.

Product Description

SUPPLEMENTED WITH HSA

RapidWarm™ Blast contains media for the warming of vitrified human blastocyst stage embryos. The media consists of a MOPS buffered solution containing gentamicin as an antibiotic agent and human serum albumin.

Warm 1™ Blast contains no cryoprotectant.

Warm 2™ Blast contains no cryoprotectant.

For use after warming to +37 °C.

The summary of safety and clinical performance can be found at www.vitrolife.com

Storage instructions and stability

Store dark at +2 to +8 °C.

RapidWarm Blast is stable until the expiry date shown on the bottle labels and the LOT specific Certificate of Analysis.

Media bottles should not be stored after opening. Discard excess media after completion of procedure.

Use of storage device

Use a legally marketed storage device indicated for use in human blastocyst vitrification procedures, in order to assure sufficient cooling and warming rates.

Use a closed storage system to prevent the potential risk of viral or other contamination of samples.

Perform the actual vitrification and warming according to the Instructions for Use for the storage device.

Directions for use

The product shall be used by an IVF professional.

The patient/parent group is an adult or reproductive-age population that undergoes IVF treatment or fertility preservation.

The following is the general procedure for using RapidWarm Blast.

Note that the timing has to be accurate.

Warning:

Place 0.5–1 ml of each of the following media into separate wells of a 4-well plate and warm to 37 °C:

• Warm 1 Blast

• Warm 2 Blast

• Warm 3 Blast

All manipulations of the blastocysts are carried out at 37 °C (on the heat stage).

Remove the cryovial containing the vitrified blastocysts from the cryostorage container. Follow the warming instructions for the specific cryovial used.

Immediately after warming, transfer blastocysts into Warm 1 Blast.

Allow the blastocysts to fall from the device and sink to the bottom. Leave for 2 min.

Transfer the blastocysts into Warm 2 Blast and let the blastocysts remain in the solution for 3 min.

Transfer the blastocysts into Warm 3 Blast and let the blastocysts remain in the solution for 10 min.

Rinse the culture in culture media several times and continue culture according to laboratory practice. Vitrolife recommends using EmbryoGleue® for this procedure.

See also instructions provided with EmbryoGleue® for this procedure.

Specifications

Aseptically prepared

Mouse Embryo Assay, 1-cell [% re-expanded blastocyst 24 hours post-test] ≥ 70

Bacterial endotoxins (LAL assay) [EU/ml] < 0.5

pH tested

Osmolarity tested

LOT-specific test results are available on the Certificate of Analysis provided with each delivery.

Precautions

The long-term safety of vitrification and/or blastocyst collagen on children born following this method of embryo cryopreservation procedure has not been established.

The risk of reproductive toxicity and developmental toxicity for IVF media, including Vitrolife's IVF media, have not yet been determined.

The safety and effectiveness of vitrification has not been fully evaluated in human embryos that have not yet reached the blastocyst stage of development.

Discard product if bottle integrity is compromised. Do not use RapidWarm Blast if it appears cloudy.

RapidWarm™ Blast is safe to use.

Caution: All glass products should be treated as potentially infectious. Source material from which this product was derived was found negative when tested for antibodies to HIV, HBc, HBe, and HTLV. No test results are available for antibodies to HCV, RNA-HIV-1 and syphilis. No known test methods can offer assurance that products derived from human blood will not transmit infectious agents.

Re-use is not recommended in microbiological contamination and/or product changes in the product.

To avoid contamination Vitrolife strongly recommends that media should be opened and used only with aseptic technique.

Any serious incident that has occurred in relation to the device should be reported to the manufacturer and the competent authority of the Member State in which the user and/or provider is established.

Not for injection.

Discard the medical hazard waste when the procedure is finished.

Symbols/glossary

*Medical devices – Symbols to be used with information to be supplied by the manufacturer (ISO 15223-1:2021).

- Warm 1 Blast
- Warm 2 Blast
- Warm 3 Blast

关于产品的所有操作需在+37 °C (在加温台上) 进行。

将装有玻璃珠的冷冻冻存载体从冷冻冻存装置中移出，通常使用带有特定冷冻液的复苏培养液。

取出后立即立即将载体转移至Warm 1 Blast。

将载体移至Warm 2 Blast，并使其在复苏中停留2分钟。

将载体移至Warm 3 Blast，并使其在复苏中停留5-10分钟。

将载体移至Warm 3 Blast，并使其在复苏液中停留5-10分钟。

将载体移至Warm 1 Blast。

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Eliminare il prodotto e l'integrità del flacone è compresa. Non utilizzare RapidWarm Blast se presenta un esposto terreno.

RapidWarm Blast contiene albumina sierica umana. Attenzione: tutti gli smodernati devono essere trattati come potenzialmente infettivi. Le materie prime impiegate per questi prodotti sono risultate negative ai test per gli scopi di controllo. Istruzioni per l'uso: Il non utilizzo può causare HIV, HCV e HIV-1 RNA e sifilide. Nessun metodo di prova può garantire che i prodotti derivati dal sangue umano non trasmettano agenti infettivi.

Il risultato può provocare contaminazione microbiologica e/o delle proprieità del prodotto.

Per evitare contaminazioni, Vitrolife raccomanda di prenotare e utilizzare il prodotto esclusivamente con tecniche asettiche.

Seguire le procedure e le attenzioni dell'utente del

Stato mentre si cui risiede l'utente o il paziente

eventuali incidenti gravi che si verificano in relazione

al dispositivo.

Non per iniezione.

Smaltire il prodotto secondo la pratica clinica standard per i risultati pericolosi al termine della procedura.

Attenzione: è legge europea (Degr) Stati Uniti d'America la vendita del prodotto disposto da prescrizione medica da parte di un professionista esperto del suo utilizzo.

It: Naudojimo indikacija

Toksini svaru mitinčios blastočiosčios stadijos žmogaus embriju atidėjimui.

Kontraindikacijos

RapidoWarm Blast® yra spūstėja ir yra gerintinkiu.

Nenuodaukti pacientams, kuriam nustatytas padidėjusias ar ateriosi sudėtinėjai medžiagai.

Gaminjimo aprašas

PAPILOTYTA HSA

„RapidoWarm Blast“ yra tarpis, skirtas vitrifikavui.

blastočiosčios stadijos žmogaus embrionų atidėjimui.

„RapidoWarm Blast“ yra spūstėja ir yra gerintinkiu.

„RapidoWarm Blast“ yra spūstėja ir yra gerintinkiu.

Naudojimo aprašymas: +37 °C.

Saugomos ir minėjimo veiksmingumo santrukia galima rasti www.vitrolife.com

Laikejimo nurodymai ir stabiliumas

Laikejimo laikas nuo +2 iki +8 °C.

„RapidoWarm Blast“ yra statikai iki antalyklos etekui

ir partios analizės serifielės nurodymos tinkamumo naudotinos.

Atidarytu buteliu su terpių laikymu išskirtus.

Blago procedūra, repausuotuose terpių likučiuose išskirtus.

Laikejimo išskirtos naudojimo instrukcijos.

Naudojimo nurodymai

Gaminys turintis naudotus IVF profesionalus.

Tikslinei pacientui grupė su yra saugomos arba

reprodukcinio amžiaus populiacija, kuriai taikomas IVF

arba yra išskirtos iš priešingimo gydymas.

Tolius patiekamus RapidWarm Blast®

naudotino procedūros.

Atkreipite dėmesį, ar laiko skaičiavimas yra tikslus.

Atkreipite dėmesį, ar