

HSA-Solution™

(100 mg/mL) and is intended for use in assisted reproductive procedures which include gamete and embryo manipulation. These procedures include the use of HSA-solution™ as a supplement for culture medium.

(100 mg/mL) and is intended for use in assisted reproductive procedures which include gamete and embryo manipulation. These procedures include the use of HSA-solution™ as a supplement for culture medium.

Product Description
HSA-solution™ is antibiotic-free.

Storage instructions and stability
Store dark at +2 to +8 °C.
HSA-solution™ is stable until the expiry date shown on the container labels and the LOT-specific Certificate of Analysis. Media bottles should not be stored after opening. Discard excess media after completion of the procedure.

Directions for use

G-Series PLUS media, G-GAMETE™ and EmbryoGlue® do not require albumin supplementation. All other G-Series culture media, not designated PLUS, come protein-free and need to be supplemented either with G-MM™ or HSA-solution™ at the appropriate concentration as listed in the table below. When selecting protein supplementation it should be noted that human serum albumin (HSA) is a blood derived substance and may contain human pathogenic agents including those not yet known or identified. Thus the risk of transmission of such infectious agents cannot be completely eliminated when using HSA. G-Series media with the exception of G-IVF™, G-FreezeKit Blast™ and G-ThawKit Blast™ shall have 5 % of either G-MM™ or HSA-solution™ added (0.5 mL added to 9.5 mL). The final concentration of G-MM™ will be 2.5 mg/mL, and the final concentration of HSA will be 5 mg/mL.

G-IVF™ shall have 10 % of either G-MM™ or HSA-solution™ added (1.0 mL added to 9.0 mL). The final concentration of G-MM™ will be 5 mg/mL, and the final concentration of HSA will be 10 mg/mL. Media to be supplemented should be aliquoted into sterile non-toxic tissue culture grade tubes or bottles. All containers should be pre-rinsed using G-RINSE™ in order to ensure there is no particulate matter or toxic substances residing.

Supplementation of G-MOPS™, G-1™, G-2™, G-PGD™
Medium HSA-solution™ Final [mL] [mL] volume [mL] 9.5 0.5 10.0 19.5 1.5 20.0 38.0 2.5 40.0 47.5 2.5 50.0 57.0 3.5 60.0 66.5 3.5 70.0 76.0 4.0 80.0 85.5 4.5 90.0 95.0 5.0 100.0

G-IVF™ shall have 10 % of either G-MM™ or HSA-solution™ added (1.0 mL added to 9.0 mL). The final concentration of G-MM™ will be 5 mg/mL, and the final concentration of HSA will be 10 mg/mL. Media to be supplemented should be aliquoted into sterile non-toxic tissue culture grade tubes or bottles. All containers should be pre-rinsed using G-RINSE™ in order to ensure there is no particulate matter or toxic substances residing.

Supplementation of G-MOPS™, G-1™, G-2™, G-PGD™
Medium HSA-solution™ Final [mL] [mL] volume [mL] 9.5 0.5 10.0 19.5 1.5 20.0 38.0 2.5 40.0 47.5 2.5 50.0 57.0 3.5 60.0 66.5 3.5 70.0 76.0 4.0 80.0 85.5 4.5 90.0 95.0 5.0 100.0

Supplementation of G-IVF™, G-FreezeKit Blast™, G-ThawKit Blast™
Medium HSA-solution™ Final [mL] [mL] volume [mL] 9.0 1.0 10.0 18.0 2.0 20.0 27.0 3.0 30.0 36.0 4.0 40.0 45.0 5.0 50.0 54.0 6.0 60.0 63.0 7.0 70.0 69.5 8.0 80.0 81.0 9.0 90.0 90.0 10.0 100.0

The cryopreservation solutions can be supplemented directly into the dishes. Simply place 900 µL of each cryopreservation solution into separate wells + 100 µL of G-MM™ or 100 µL HSA-solution™. It is recommended only to supplement the volume of medium expected to be used in one day.

Specifications
Sterile filtered SAL 10³ Mouse Embryo Assay (1-cell) [% expanded blastocyst within 96 hours] ≥ 80 Bacterial endotoxins (LAL assay) [EU/mL] < 0.5 LOT specific test results are available on the Certificate of Analysis provided with each delivery.

Precautions
Discard product if bottle integrity is compromised. Do not use HSA-solution™ if it appears cloudy.

HSA-solution™ contains human serum albumin. Caution: All blood products should be treated as potentially infectious. Source material from which this product was derived was found negative when tested for antibodies to HIV, HBs, HCV, and HTLV III and non-reactive for HbsAg, HCV RNA and HIV-1 RNA and syphilis. No known test methods can offer assurance that products derived from human blood will not transmit infectious agents.

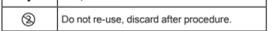
Re-use may result in microbiological contamination and/or property changes in the product.

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

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

Not for injection.

Caution: Federal (US) law restricts this device to sale by or on the order of a physician.

Description of ISO Symbols	
	Sterilized using aseptic processing techniques
	Temperature limit
	Do not re-use, discard after procedure.
	Use by - see label.
	Caution: Consult accompanying documents.
	Catalog number
	Batch code
	CE Mark (Conformité Européen)

CHINA

CN : 培养液添加剂 HSA-solution™ 产品说明书
适应症
HSA-solution™含有100 mg/mL人血清白蛋白。该产品在辅助生殖技术中作为用于配子和胚胎处理的培养液添加剂。

产品型号及规格	
型号：见标签	
规格：见标签	
产品注册证及技术要求编号	
产品注册证编号：见标签	
产品技术要求编号：见标签	
产品描述	
该产品不含抗菌素。	
产品有效期及贮存条件	
产品有效期见标签。	
避光储存于+2 °C +8 °C。	
HSA-solution™在包装标签上和技术产品批号的分析报告上显示的有效期内的稳定性。	
产品包装打开后不得再继续储藏。使用后丢弃剩余产品。	
使用方法	
G系列中PLUS产品、G-GAMETE™和EmbryoGlue®不需要添加白蛋白。所有其他非PLUS产品，不含白蛋白产品均需要添加适量HSA-solution™或G-MM™。其添加量列于下表中。但请注意，并非所有HSA-solution™和G-MM™均含有白蛋白。（即5.0 mL至9.5 mL）。最终G-MM™的白蛋白浓度为2.5 mg/mL，HSA-solution™为5 mg/mL。	
G-IVF™含有10% G-MM™或HSA-solution™（加1.0 mL至9.0 mL）。最终G-MM™的浓度为5 mg/mL，HSA-solution™为10 mg/mL。添加白蛋白的培养液均应用无菌无毒的试管分配。所有试剂管都应预先用G-RINSE™冲洗以清除任何残留物。	
G-MOPS™, G-1™, G-2™, G-PGD™添加量：	
培养液 HSA-solution™ 最终体积 [mL] [mL] [mL] 9.5 0.5 10.0 19.0 1.5 20.0 38.0 2.5 40.0 47.5 2.5 50.0 57.0 3.5 60.0 66.5 3.5 70.0 76.0 4.0 80.0 85.5 4.5 90.0 95.0 5.0 100.0	
G-IVF™含有10% G-MM™或HSA-solution™（加1.0 mL至9.0 mL）。最终G-MM™的浓度为5 mg/mL，HSA-solution™为10 mg/mL。添加白蛋白的培养液均应用无菌无毒的试管分配。所有试剂管都应预先用G-RINSE™冲洗以清除任何残留物。	

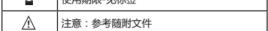
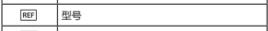
规格
灭菌过滤 SAL 10³ 胚胎或两级(二细胞阶段) [%6小时内扩张率] ≥ 80 细菌内毒素(LAL检测) [EU/mL] < 0.25 每个批号的原始分析报告随货提供。

注意事项
如果发现瓶子的完整性受损，请丢弃该产品。如果发现产品密封盖有受损。

HSA-solution™含人血清白蛋白。Caution：所有的血液制品均被视为潜在感染源。产品生产原料经过HIV、HBs、HCV及HTLVIII抗体检测均为阴性，对HbsAg、HCV RNA及HIV-1 RNA及梅毒均为阴性。已知检测的方法并不能保证无血清制品不含任何感染源。重复使用可能导致生物活性产品产生不良反应。

Vitrolife使用可降解生物活性产品，严格按照无菌操作技术。关于使用IVF培养液产品，包括Vitrolife的IVF培养液产品，在生殖健康和生育等上的稳定性有充分的数据支持。

注意：产品必须严格按照医疗行业相关法规和法规，仅限专业人士使用。

ISO符号描述	
	经无菌处理
	温度限制
	切勿再次使用
	使用期限-见标签
	注意：参考附件文件
	批号
	型号
	CE标记 (Conformité Européen)

EUROPE

EN : Indication for use

HSA-solution™ contains human serum albumin solution

(100 mg/mL) and is intended for use in assisted reproductive procedures which include gamete and embryo manipulation. These procedures include the use of HSA-solution™ as a supplement for culture medium.

(100 mg/mL) and is intended for use in assisted reproductive procedures which include gamete and embryo manipulation. These procedures include the use of HSA-solution™ as a supplement for culture medium.

Product Description
HSA-solution™ is antibiotic-free.

Storage instructions and stability
Store dark at +2 to +8 °C.
HSA-solution™ is stable until the expiry date shown on the container labels and the LOT-specific Certificate of Analysis. Media bottles should not be stored after opening. Discard excess media after completion of the procedure.

Directions for use

G-Series PLUS media, G-GAMETE™ and EmbryoGlue® do not require albumin supplementation. All other G-Series culture media, not designated PLUS, come protein-free and need to be supplemented either with G-MM™ or HSA-solution™ at the appropriate concentration as listed in the table below. When selecting protein supplementation it should be noted that human serum albumin (HSA) is a blood derived substance and may contain human pathogenic agents including those not yet known or identified. Thus the risk of transmission of such infectious agents cannot be completely eliminated when using HSA. G-Series media with the exception of G-IVF™, G-FreezeKit Blast™ and G-ThawKit Blast™ shall have 5 % of either G-MM™ or HSA-solution™ added (0.5 mL added to 9.5 mL). The final concentration of G-MM™ will be 2.5 mg/mL, and the final concentration of HSA will be 5 mg/mL.

G-IVF™ shall have 10 % of either G-MM™ or HSA-solution™ added (1.0 mL added to 9.0 mL). The final concentration of G-MM™ will be 5 mg/mL, and the final concentration of HSA will be 10 mg/mL. Media to be supplemented should be aliquoted into sterile non-toxic tissue culture grade tubes or flasks. All containers should be pre-rinsed using G-RINSE™ in order to ensure there is no particulate matter or toxic substances residing.

Supplementation of G-MOPS™, G-1™, G-2™, G-PGD™
Medium HSA-solution™ Final [mL] [mL] volume [mL] 9.5 0.5 10.0 19.0 1.0 20.0 28.5 1.5 30.0 38.0 2.0 40.0 47.5 2.5 50.0 57.0 3.0 60.0 66.5 3.5 70.0 76.0 4.0 80.0 85.5 4.5 90.0 95.0 5.0 100.0

Supplementation of G-IVF™, G-FreezeKit Blast™, G-ThawKit Blast™
Medium HSA-solution™ Final [mL] [mL] volume [mL] 9.0 1.0 10.0 18.0 2.0 20.0 27.0 3.0 30.0 36.0 4.0 40.0 45.0 5.0 50.0 54.0 6.0 60.0 63.0 7.0 70.0 69.5 8.0 80.0 81.0 9.0 90.0 90.0 10.0 100.0

The cryopreservation solutions can be supplemented directly into the dishes. Simply place 900 µL of each cryopreservation solution into separate wells + 100 µL of G-MM™ or 100 µL HSA-solution™. It is recommended only to supplement the volume of medium expected to be used in one day.

Specifications
Sterile filtered SAL 10³ Mouse Embryo Assay (1-cell) [% expanded blastocyst within 96 hours] ≥ 80 Bacterial endotoxins (LAL assay) [EU/mL] < 0.5 LOT specific test results are available on the Certificate of Analysis provided with each delivery.

Precautions
Discard product if bottle integrity is compromised. Do not use HSA-solution™ if it appears cloudy.

HSA-solution™ contains human serum albumin. Caution: All blood products should be treated as potentially infectious. Source material from which this product was derived was found negative when tested for antibodies to HIV, HBs, HCV, and HTLV III and non-reactive for HbsAg, HCV RNA and HIV-1 RNA and syphilis. No known test methods can offer assurance that products derived from human blood will not transmit infectious agents.

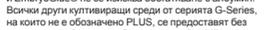
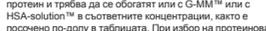
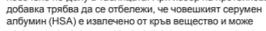
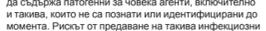
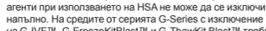
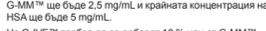
Re-use may result in microbiological contamination and/or property changes in the product.

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

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

Not for injection.

Caution: Federal (US) law restricts this device to sale by or on the order of a physician.

Description of ISO Symbols	
	Sterilized using aseptic processing techniques
	Temperature limit
	Do not re-use, discard after procedure.
	Use by - see label.
	Caution: Consult accompanying documents.
	Catalog number
	Batch code
	CE Mark (Conformité Européen)

HSA-solution™ съдържа ратвор от човешки сѳрумелн албумин (100 mg/mL) и е предназначан за употреба при медицински процедури на гамети и ембриони. Тези процедури включват използването на HSA-solution™ като добавка към културиваща среда.

Описание на продукта
HSA-solution™ не съдържа антибиотик.

Инструкции за съхранение и стабилност
Да се съхранява на тъмно при температура от +2 до +8 °C. HSA-solution™ е стабилен до изтичане на срока на годност, обозначен върху етикета на опаковката и на спецификация за партидата Сертификат от анализ.

Бутилките със среда не трябва да се съхраняват след отваряне. Изхвърлете изтичното количество среда след употреба на продукта.

Указания за употреба
При предете от серията G-Series PLUS, G-GAMETE™ и EmbryoGlue® не е необходимо добавяне на албумин. Ала andre G-Series, държкисмедии, дер кие хав бетелсен PLUS, ер protein-free ved levering og skal tilføres enten G-MM™ eller HSA-solution™ i passende koncentrationer som angivet i nedenstående tabel. I forbindelse hermed vil vi pдrde til, at proteinet/serum skal det bemrkes, at human serum albumin (HSA) er udsundet af blod og derfor kan indeholde humane patogener, herunder stoffer, der endnu ikke er kendt eller endnu ikke er blevet oplyst om. Disse patogener kan overføres til receptorer og derfor ikke helt udgdes ved brug af HSA. G-Series-medier med undtagelse af G-IVF™, G-FreezeKit Blast™ og G-ThawKit Blast™ skal tilføres 5 % G-MM™ eller HSA-solution™ (0,5 mL tilføjes til 9,5 mL). Krafnata koncentrationer na G-MM™ vil vе 2,5 mg/mL, og krfnata koncentrationer na HSA vе 5 mg/mL.

G-IVF™ skal tilføres 10 % G-MM™ eller HSA-solution™ (1,0 mL tilføjes til 9,0 mL). Slutkoncentrationen af G-MM™ vil vе 5 mg/mL, og slutkoncentrationen af HSA vil vе 10 mg/mL. Medier, der skal tilføres proteinopløsning, skal alukveres over sterile, ikke-oksiderende reagensglas eller flasker beredet til vaskulatur. Alle beholdere bør skylles med G-RINSE™ fсrst for at sikre, at de ikke indeholder partikkelrest eller giftstoffer.

Tilførsel til G-MOPS™, G-1™, G-2™, G-PGD™
Medium HSA-solution™ Slut- [mL] [mL] volumen [mL] 9.5 0.5 10.0 19.0 1.0 20.0 28.5 1.5 30.0 37.0 2.0 40.0 47.5 2.5 50.0 57.0 3.0 60.0 66.5 3.5 70.0 76.0 4.0 80.0 85.5 4.5 90.0 95.0 5.0 100.0

Tilførsel til G-IVF™, G-FreezeKit Blast™, G-ThawKit Blast™
Medium HSA-solution™ Slut- [mL] [mL] volumen [mL] 9.0 1.0 10.0 18.0 2.0 20.0 27.0 3.0 30.0 36.0 4.0 40.0 45.0 5.0 50.0 54.0 6.0 60.0 63.0 7.0 70.0 69.5 8.0 80.0 81.0 9.0 90.0 90.0 10.0 100.0

Da G-IVF™ skal tilføres 10 % G-MM™ eller HSA-solution™ (1,0 mL tilføjes til 9,0 mL), skal slutkoncentrationen af G-MM™ blive 5 mg/mL, og slutkoncentrationen af HSA blive 10 mg/mL.

Opbejtgange og stabilitet
Opbevares mørkt ved +2 til +8 °C.

HSA-solution™ kan holde sig uldledst, uden at blive fцrdet af etiketterne på beholderne og det LOT-specifikke analysecertifikat.

Mediefasker må ikke opbevares efter åbning. Kasser overvaskende medie efter fцrdiggelse af proceduren.

Brugsanvisning
G-Series PLUS-medier, G-GAMETE™ and EmbryoGlue® behøver ikke tilførsel af albumin. Alle andre G-Series, dyrkingsmedier, der ikke har betegnelsen PLUS, er protein-free ved levering og skal tilføres enten G-MM™ eller HSA-solution™ i passende koncentrationer som angivet i nedenstående tabel. I forbindelse hermed vil vi pдrde til, at proteinet/serum skal det bemrkes, at human serum albumin (HSA) er udsundet af blod og derfor kan indeholde humane patogener, herunder stoffer, der endnu ikke er kendt eller endnu ikke er blevet oplyst om. Disse patogener kan overføres til receptorer og derfor ikke helt udgdes ved brug af HSA. G-Series-medier med undtagelse af G-IVF™, G-FreezeKit Blast™ og G-ThawKit Blast™ skal tilføres 5 % G-MM™ eller HSA-solution™ (0,5 mL tilføjes til 9,5 mL). Krifnata koncentrationer af G-MM™ vil vе 2,5 mg/mL, og slutkoncentrationen af HSA 5 mg/mL.

G-IVF™ skal tilføres 10 % G-MM™ eller HSA-solution™ (1,0 mL tilføjes til 9,0 mL). Slutkoncentrationen af G-MM™ vil vе 5 mg/mL, og slutkoncentrationen af HSA vil vе 10 mg/mL. Medier, der skal tilføres proteinopløsning, skal alukveres over sterile, ikke-oksiderende reagensglas eller flasker beredet til vaskulatur. Alle beholdere bør skylles med G-RINSE™ fсrst for at sikre, at de ikke indeholder partikkelrest eller giftstoffer.

Tilførsel til G-MOPS™, G-1™, G-2™, G-PGD™
Medium HSA-solution™ Slut- [mL] [mL] volumen [mL] 9.5 0.5 10.0 19.0 1.0 20.0 28.5 1.5 30.0 37.0 2.0 40.0 47.5 2.5 50.0 57.0 3.0 60.0 66.5 3.5 70.0 76.0 4.0 80.0 85.5 4.5 90.0 95.0 5.0 100.0

G-IVF™ skal tilføres 10 % G-MM™ eller HSA-solution™ (1,0 mL tilføjes til 9,0 mL). Slutkoncentrationen af G-MM™ vil vе 5 mg/mL, og slutkoncentrationen af HSA vil vе 10 mg/mL. Medier, der skal tilføres proteinopløsning, skal alukveres over sterile, ikke-oksiderende reagensglas eller flasker beredet til vaskulatur. Alle beholdere bør skylles med G-RINSE™ fсrst for at sikre, at de ikke indeholder partikkelrest eller giftstoffer.

Tilførsel til G-IVF™, G-FreezeKit Blast™, G-ThawKit Blast™
Medium HSA-solution™ Slut- [mL] [mL] volumen [mL] 9.0 1.0 10.0 18.0 2.0 20.0 27.0 3.0 30.0 36.0 4.0 40.0 45.0 5.0 50.0 54.0 6.0 60.0 63.0 7.0 70.0 69.5 8.0 80.0 81.0 9.0 90.0 90.0 10.0 100.0

dođavaj bez protina a musaji bij suplemenatovano buf G-MM™ nebo HSA-solution™ i v vhodné koncentraci, jak je uvedeno v tabulce nize. Pri povbe proteinovne suplementacije je treba uvajati, ze lidsky sѳrovny albumin (HSA) je krevni derivat a moze obsahovat lidski patogeni agenci, vctne tch, kter neseou znatny nebo nebyli identifikovani. Tyto riziko přenosu infekci netze pri použití HSA zcela vyloučí. Přípravky G-Series s výjimkou G-IVF™, G-FreezeKit Blast™ a G-ThawKit Blast™ musaji obsahovat 5% suplemenatovného přípravku G-MM™ nebo HSA-solution™ (0,5 ml přidáme k 9,5 ml). Konečná koncentrace přípravku G-MM™ bude 2,5 mg/ml a konečná koncentrace HSA bude 5 mg/ml.

G-IVF™ bude mít 10 % HSA nebo 10 mg/ml G-MM™ nebo HSA-solution™ (1,0 ml přidány k 9,0 ml). Konečná koncentrace přípravku G-MM™ bude 5 mg/ml a konečná koncentrace přípravku HSA bude 10 mg/ml. Před přidáním suplementu by měla být média rozložena na alkivky do netoxických zkumavek nebo lahví pro křdřkové kultivace. Všechny nádoby musaji být předem vypláchnuty přípravkem G-RINSE™ pro zajištění, že neobsahují žádné pevné částice ani toxické látky.

Suplementace G-MOPS™, G-1™, G-2™, G-PGD™
Přípravek HSA-solution™ Konečný objem [ml] [ml] [ml] 9.5 0.5 10.0 19.0 1.0 20.0 28.0 2.0 40.0 47.5 2.5 50.0 57.0 3.0 60.0 66.5 3.5 70.0 76.0 4.0 80.0 85.5 4.5 90.0 95.0 5.0 100.0

Suplementace G-IVF™, G-FreezeKit Blast™, G-ThawKit Blast™
Přípravek HSA-solution™ Konečný objem [ml] [ml] [ml] 9.0 1.0 10.0 18.0 2.0 20.0 27.0 3.0 30.0 36.0 4.0 40.0 45.0 5.0 50.0 54.0 6.0 60.0 63.0 7.0 70.0 69.5 8.0 80.0 81.0 9.0 90.0 90.0 10.0 100.0

Kryokonzervační roztoky mohou být suplemenatovány přímo do destiček. Jednoduše umístíte 900 µL každého kryokonzervačního roztoku do jednotlivých jarek + 100 µL přípravku G-MM™ nebo 100 µL HSA-solution™. Doporučuje se suplemenatovat pouze objem přípravku, který se má použít tentýž den.

Specifikace
Filtrováno sterilním způsobem SAL 10³ Test na myšič embryích (třbuřkovy) [% expandovaných blastocysto do 96 hodin] ≥ 80 Bakteriální endotoxiny (LAL test) [EU/mL] < 0,5 Výsledky testů pro jednotlivé šarže najdte v certifikátu analýzy, které pro injekci posíláme odděky.

Preventivní opatření
Produkt zkontrolujte, došlo-li k porušení celistvosti lékovky. Nepoužívejte roztok, pokud je vidět negativní vzorek.

HSA-solution™ obsahuje lidský sѳrovny albumin. Upozornění: Se všemi krevními produkty musí být nakládáno jako s potenciálně infekčními. Zdrojový materiál, z něhož je tento produkt vyroben, byl negativně podléhl testu na přítomky příčiv HIV, HBs, HCV, a HTLV III a nebyl reaktivní pro HbsAg, HCV RNA a HIV-1 RNA a syfilis. Žádné známé testovací postupy nemohou zajistit, že produkt vyrobené z lidské krve nepřenáší přenos infekci.

Opakované použití může mít za následek mikrobiologickou kontaminaci a změnu vlastností produktu.

Pro vyhození kontaminace Vitrolife důrazně doporučuje, aby se přípravky otevřely a používaly pouze pomocí aseptického postupu.

Riziko reprodukční toxicity a vývojové toxicity u přípravku ro v in vitro fertilizaci (IVF), včetně IVF přípravku společností Vitrolife, nelze stanovená a nejsou známa.

Upozornění: Podle federálních zákonů USA smí tento přípravek prodávat pouze lékař nebo smí být prodán pouze na lékařský předpis.

DA : Indication for brug
HSA-solution™ indeholder en opløsning af human serum albumin (100 mg/mL) og er beregnet til brug af assisterede reproduktionsprocedurer, som omfatter manipulation af kønsceller og embryoer. Disse procedurer omfatter brug af HSA-solution™ som tilskud til de dyrkingsmedier.

HSA-solution™ kan holde sig uldledst, uden at blive fцrdet af etiketterne på beholderne og det LOT-specifikke analysecertifikat.

Mediefasker må ikke opbevares efter åbning. Kasser overvaskende medie efter fцrdiggelse af proceduren.

Brugsanvisning
G-Series PLUS-medier, G-GAMETE™ and EmbryoGlue® behøver ikke tilførsel af albumin. Alle andre G-Series, dyrkingsmedier, der ikke har betegnelsen PLUS, er protein-free ved levering og skal tilføres enten G-MM™ eller HSA-solution™ i passende koncentrationer som angivet i nedenstående tabel. I forbindelse hermed vil vi pдrde til, at proteinet/serum skal det bemrkes, at human serum albumin (HSA) er udsundet af blod og derfor kan indeholde humane patogener, herunder stoffer, der endnu ikke er kendt eller endnu ikke er blevet oplyst om. Disse patogener kan overføres til receptorer og derfor ikke helt udgdes ved brug af HSA. G-Series-medier med undtagelse af G-IVF™, G-FreezeKit Blast™ og G-ThawKit Blast™ skal tilføres 5 % G-MM™ eller HSA-solution™ (0,5 mL tilføjes til 9,5 mL). Slutkoncentrationen af G-MM™ vil vе 2,5 mg/mL, og slutkoncentrationen af HSA 5 mg/mL.

G-IVF™ skal tilføres 10 % G-MM™ eller HSA-solution™ (1,0 mL tilføjes til 9,0 mL). Slutkoncentrationen af G-MM™ vil vе 5 mg/mL, og slutkoncentrationen af HSA vil vе 10 mg/mL. Medier, der skal tilføres proteinopløsning, skal alukveres over sterile, ikke-oksiderende reagensglas eller flasker beredet til vaskulatur. Alle beholdere bør skylles med G-RINSE™ fсrst for at sikre, at de ikke indeholder partikkelrest eller giftstoffer.

Tilførsel til G-MOPS™, G-1™, G-2™, G-PGD™
Medium HSA-solution™ Slut- [mL] [mL] volumen [mL] 9.5 0.5 10.0 19.0 1.0 20.0 28.5 1.5 30.0 37.0 2.0 40.0 47.5 2.5 50.0 57.0 3.0 60.0 66.5 3.5 70.0 76.0 4.0 80.0 85.5 4.5 90.0 95.0 5.0 100.0

