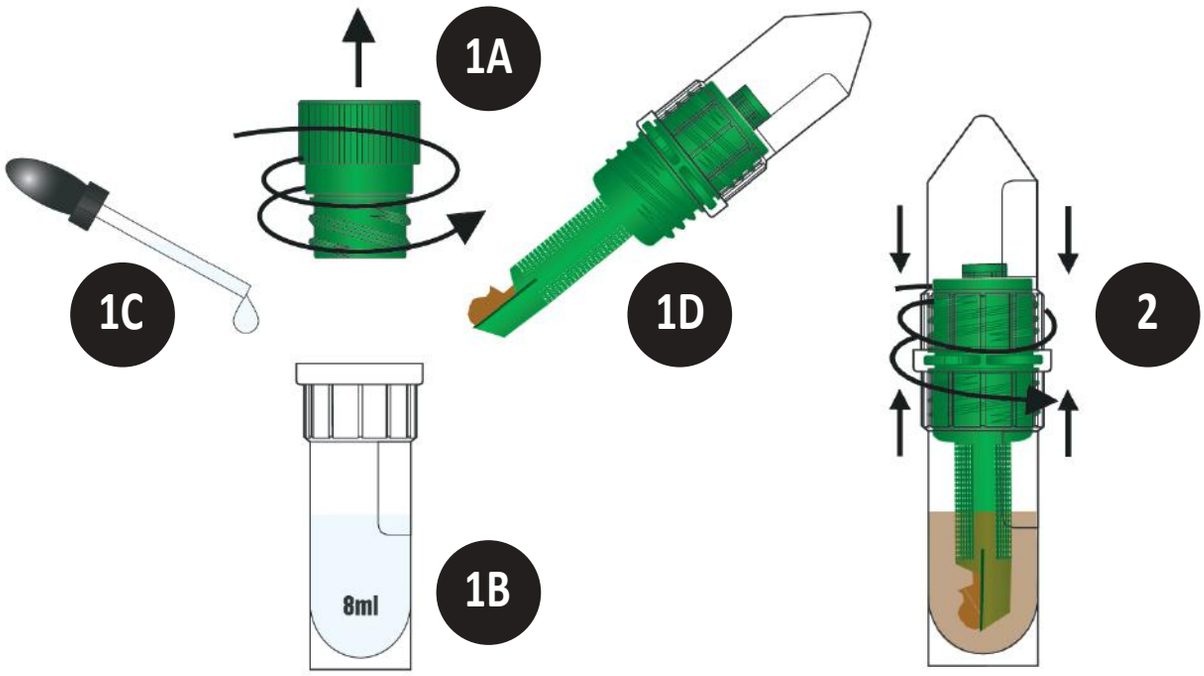
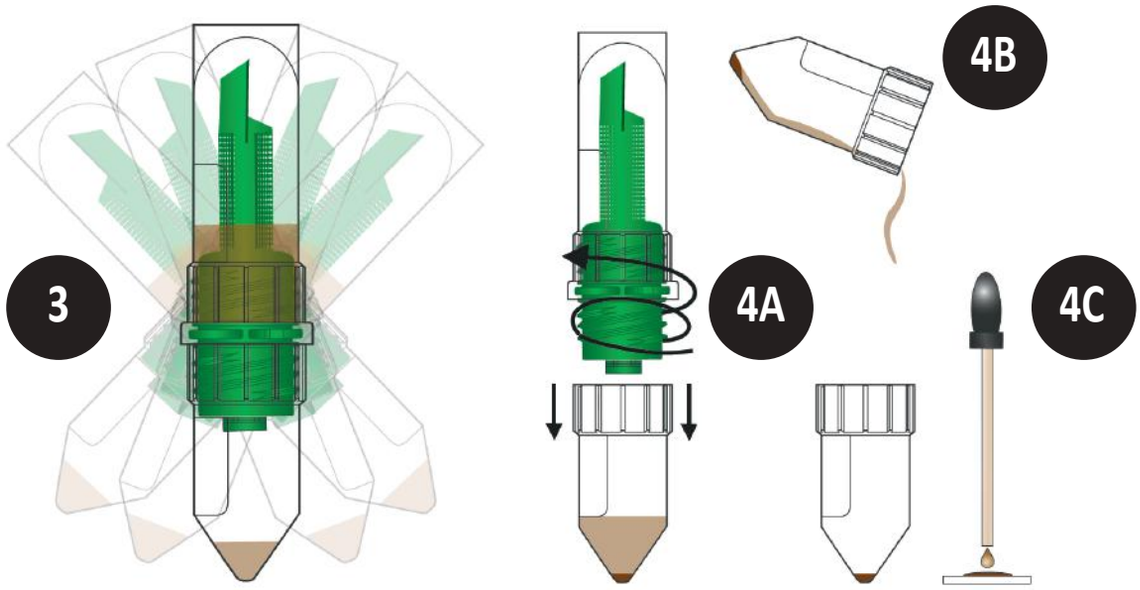
APACOR

Midi Parasep® SF Faecal Parasite Concentrator





CS	**Midi Parasep® SF** Koncentrátor parazitů ve stolici
DE	**Midi Parasep® SF** Konzentrator für Stuhlparasiten
ES	**Midi Parasep® SF** Concentrador de parásitos fecales
FR	**Midi Parasep® SF** Concentrateurs de Parasites Fécaux
HR	**Midi Parasep® SF** Koncentratori crijevnih parazita bez otapala
IT	**Midi Parasep® SF** Concentratore di parassiti fecali
NL	**Midi Parasep® SF** Fecale Parasieten Concentrator
PL	**Midi Parasep® SF** System do zagęszczenia kału przy analizie parazytów
PT	**Midi Parasep® SF** Concentrador de parasitas fecais
SI	**Midi Parasep® SF** Koncentrator parazitov v blatu

APA103
V8 09/2018

CE marking (European directive 98/79/CE on in vitro diagnostic medical devices)
Označení CE (Evropská směrnice 98/79 / ES o diagnostických zdravotnických prostředcích in vitro)
CE-Kennzeichnung (EG-Richtlinie 98/79 / EG über In-vitro-Diagnostika)
Marcado CE (directiva europea 98/79 / CE sobre productos sanitarios para diagnóstico in vitro)
Marquage CE (directive européenne 98/79 / CE relative aux dispositifs médicaux de diagnostic in vitro)
CE označavanje (Europska direktiva 98/79 / EZ o in vitro dijagnostičkim medicinskim uređajima)
Marcatura CE (Direttiva Europea 98/79 / CE relativa ai dispositivi medico-diagnostici in vitro)
CE-markering (Europese richtlijn 98/79 / EG betreffende de in vitro diagnostische medische hulpmiddelen)
Oznakowanie CE (dyrektywa europejska 98/79 / WE w sprawie wyrobów medycznych do diagnostyki in vitro)
Marcação CE (directiva europeia 98/79 / CE relativa aos dispositivos médicos de diagnóstico)
Oznaka CE (Evropska direktiva 98/79 / ES o in vitro diagnostičnih medicinskih pripomočkih)

IVD

For in vitro diagnostic use
K diagnostickému použití in vitro
Für in-vitro-Diagnostik
Para uso diagnóstico in vitro
Pour diagnostic in vitro
Za in vitro dijagnostičke svrhe
Per uso diagnostico in vitro
Voor in vitro diagnostisch gebruik
Do diagnostyki in vitro
Para uso diagnóstico in vitro
Za in vitro diagnostično uporabo

REF

Catalogue number
Katalogové číslo
Katalognummer
Número de catálogo
Numéro de catalogue
Kataloški broj
Numero di catalogo
Catalogus nummer
Numer katalogowy
Número de catálogo
Kataloška številka

LOT

Batch code
Kód šarže
Loskennzeichen
Código de lote
Code de lot
Serija broj
Codice del lotto
Batchcode
Kod partii
Código do lote
Kodo serije

Expiry date MM/YYYY
Datum ukončení platnosti MM / YYYY
Gültig bis MM / JJJJ
Fecha de caducidad MM / AAAA
Date d'expiration MM / AAAA
Datum isteka MM / GGGG
Data di scadenza MM / AAAA
Vervaldatum MM / YYYY
Termin ważności MM / YYYY
Data de validade MM / AAAA
Datum prenehanja veljavnosti MM / LLLL

Storage temperature limitation
Omezení skladovací teplota
Lagertemperaturbegrenzung
Límite de temperatura
Limitation de la température de stockage
Ograničenje temperature skladištenja
Limitazione della temperatura di stoccaggio
Begrenzing bewaartemperatuur
Ograniczenie temperatury bagażu
Limitação de temperatura de armazenamento
Omejitev temperature za shranjevanje

Manufacturer
Výrobce
Hersteller
Fabricante
Fabricant
Proizvođač
Fabbricante
Fabrikant
Producent
Fabricante
Proizvajalec

Consult instruction for use
Konzultujte návod k použití
Consult Gebrauchsanweisung
Consulte las instrucciones de uso
Consultez Mode d'emploi
Posavjetujte se Naputak za primjenu
Consultare istruzioni per l'uso
Raadpleeg Gebruiksaanwijzing
Skonsultuj Instrukcja użycia
Consulte Instruções de uso
Consult Navodila za uporabo

EN

See label for storage conditions and expiry date. Please adhere to the following guidelines when handling Midi Parasep® SF. To avoid cross contamination the Midi Parasep® SF device should remain closed at all times except when introducing the sample or when retrieving the final concentrated sample for examination.

Sample Preparation

If using prefilled Midi Parasep® SF, start at 1D

1A Unscrew lid.

1B Add 8.0ml of fixative.

1C Add one drop of surfactant (eg: Apacor Triton X solution) to the chamber.

1D Introduce a pea sized faecal sample to the fixative. Mix in thoroughly with the Midi Parasep® SF spoon. If the sample is hard, break it up with the end of the spoon.

Emulsification

2 Seal Midi Parasep® SF by screwing in the filter/sedimentation cone unit. Vortex or shake to emulsify with the sedimentation cone pointing upwards.

Centrifugation

3 Invert Midi Parasep® SF and centrifuge at 400g for 2 minutes. (J. Clin. Microbiol. doi:10.1128/JCM.00838-15) Midi Parasep® SF fits all 50ml centrifuge buckets.

NOTE: To calculate the required RPM for any centrifuge

$$RPM = \sqrt{\frac{g}{1.12r}} \times 1000$$

RPM Rotor Speed in revs/min

g centrifugal force (max. 1000g)

r radius, horizontal distance between sedimentation cone tip and spindle centre measured in mm

Examination

4A Unscrew and discard the filter and mixing tube.

4B Pour off all the liquid above the sediment.

4C Pipette one drop of sediment onto a slide and cover with coverslip. Alternatively, follow laboratory SOP for slide preparation.

CS

Podmínky skladování a exspirace jsou uvedeny na nálepce. Při práci s Midi Parasep® SF dodržujte prosím následující návod. Abychom zabránili kontaminaci, musí Midi Parasep® SF koncentrátor zůstat po celou dobu uzavřený s výjimkou zavádění vzorku nebo když je koncentrovaný vzorek předán ke zkoumání.

Příprava vzorku

Při použití předplněné Midi Parasep® SF, začněte u 1D.

1A Odšroubujte víčko.

1B Přidejte 8,0 ml fixačního.

1C Jednu kapku surfaktantu (např. Apacor Triton X roztok) do míchacího prostoru.

1D Zavést kopeček fekálně vzorku s fixačním pomocí lžíce na konci filtru Midi Parasep® SF. Vmícháme důkladně lžící Midi Parasep® SF. V případě, že vzorek je těžké, rozdělit to s koncem lžíce.

Emulgace

2 Filtrační díl pevně sešroubujte dohromady se zásobníkem roztoku a krátce promíchejte.

Centrifugace

3 Vložte do centrifugy a centrifugujte při 400 g po dobu 2 minut. (J. Clin. Microbiol. doi:10.1128/JCM.00838-15) Pro Midi Parasep® SF můžete použít všechny 50 ml adaptéry.

Poznámka: Pro výpočet RPM použijte následující vzorec

$$RPM = \sqrt{\frac{g}{1.12r}} \times 1000$$

RPM Otáčky rotoru

g Odstředivá síla centrifugy (max.1000g)

r Rádius, horizontální vzdálenost mezi koncem sedimentační zkumavky a středem osy, měřeno v mm

Zkoumání vzorku

4A Vyjměte Midi Parasep® SF, odšroubujte filtrační díl a zlikvidujte (tento díl zůstává uzavřen).

4B Vylijte veškerou kapalinu nad usazeninou.

4C Napipetujte jednu kapku sedimentu na sklíčko mikroskopu a zakryjte krycím sklíčkem. Alternativně postupujte podle laboratorních standardních postupů pro přípravu mikroskopických sklíček.

DE

Haltbarkeit und Aufbewahrung : Siehe Packungsaufdruck
Bitte beim Verwenden von Midi Parasep® SF die nachfolgenden Anweisungen beachten. Um Kreuz-kontamination zu vermeiden, sollte das Midi Parasep® SF Röhrchen, außer bei Probenzugabe und Entnahme des Sediments zur mikroskopischen Untersuchung, immer verschlossen bleiben.

Probenvorbereitung
Bei der Verwendung von vorgefüllten Midi Parasep® SF bei 1D beginnen.

1A Deckel abschrauben.

1B 8,0ml Fixierlösung zugeben.

1C Apacor Triton X Lösung (1:20 Verdünnung) zugeben.

1D Eine erbsengroße Kotprobe zum Fixierlösung zugeben. Mit dem Midi Parasep® SF Löffel gut mischen. Bei harten Stuhlproben mit dem Löffel zerdrücken.

Emulgieren
2 Den Filterteil mit dem Sedimentationsröhrchen des Midi Parasep® SF mit dem Probenröhrchen fest zusammenschrauben. Die Probe gut mischen mittels Votexmischer bzw. kräftig schütteln, bis eine homogene Emulsion entsteht. Es ist wichtig, daß der Konusboden des Sedimentationsröhrchen nach oben zeigt.

Zentrifugation
3 Midi Parasep® SF mit dem konusförmigen, spitz zulaufenden Teil nach unten in die Zentrifuge stellen. 2 Minuten bei 400xg zentrifugieren. (J. Clin. Microbiol. doi:10.1128/JCM.00838-15)
Midi Parasep® SF passt in all gängigen 50ml Zentrifugenröhrchen-Aufsätze.

RPM-Berechnung für all gängigen Zentrifugen

$$RPM = \sqrt{\frac{g}{1.12r}} \times 1000$$

RPM Rotordrehzahl in Umdrehungen/min
g Zentrifugalkraft (max 1000g)
r Radius, Abstand zw. dem unteren Ende des konischen Röhrchen und der Zentrifugenspindel, in mm

Probenuntersuchung

4A Midi Parasep® SF aufdrehen und den Filterteil entsorgen (dieser Teil sollte beim Aufdrehen verschlossen bleiben).

4B Den Überstand vorsichtig abgießen.

4C Einen Tropfen Sediment auf einen Objektträger pipettieren und mit Deckglas bedecken. Alternativ folgen Sie den Laborstandard für die Präparation des Objektträgers.

ES

Mirar la etiqueta para ver condiciones de almacenaje y fecha caducidad. Cuando se manipule Midi Parasep® SF se ruega seguir las instrucciones. Para evitar contaminaciones cruzadas el Midi Parasep® SF ha de permanecer siempre cerrado, excepto cuando se introduce la muestra o cuando se extrae la preparación final con objeto de ser examinada.

Preparación de la muestra
Si utiliza Midi Parasep® SF prellenado, comience en 1D.

1A Desenroscar el tapón.

1B Añadir 8.0 ml de fijador.

1C Si se requiere una gota de surfactante (solución Apacor Triton X) para emulsionar.

1D Introducir una muestra de heces del tamaño de un guisante en el fijador. Agitar vigorosamente con la cuchara del dispositivo Midi Parasep® SF. Si la muestra es de consistencia dura, trocearla con la punta de la cuchara

Emulsionado
2 Enroscar la cámara de mezcla con la unidad de filtro/cono de sedimentación. Vortear o agitar para emulsionar con el cono de sedimentación hacia arriba.

Centrifugación
3 Invertir el Midi Parasep® SF y centrifugar a 400g durante 2 minutos. (J. Clin. Microbiol. doi:10.1128/JCM.00838-15)
El Midi Parasep® SF se adecúa a todas las cestas de centrifugatión de 50 ml.

Nota: Para calcular la RPM requeridas para cualquier centrifuga

$$RPM = \sqrt{\frac{g}{1.12r}} \times 1000$$

RPM Velocidad del rotor
g Fuerza centrífuga (max 1000g)
r Radio, distancia entre la punta del cono y el centro del rotor medida en mm.

Examen

4A Desenrosque y elimine la cámara de mezcla junto cone el filtro.

4B Decante el liquido sobrenadante del sedimento.

4C Pipetee una gota de sedimento en un portaobjetos y cúbralo con un cubreobjetos. Alternativamente, siga el Procedimiento de funcionamiento estándar del laboratorio para la preparación de portaobjetos.

FR

Voir étiquette pour stockage et date d'expiration.
Respectez les consignes suivantes lorsque vous manipulez le Midi Parasep®. Pour éviter la contamination croisée, le Midi Parasep® SF devrait rester fermé, sauf lors de la saisie de l'échantillon ou quand vous prenez l'échantillon concentré final pour l'examination.

Préparation de l'échantillon
Si vous utilisez Midi Parasep® SF pré-remplie, commencer à 1D.

1A Dévissez le bouchon.

1B Ajoutez 8,0ml de fixateur.

1C Et ajoutez une goutte de surfactant (par ex: solution Apacor Triton X) pour émulsifier.

1D Introduire un échantillon fécal de la taille d'un pois au fixateur. Mélanger soigneusement avec la cuillère Midi Parasep® SF. Si l'échantillon est trop dur, rompre avec la fin de la cuillère.

Émulsification
2 Scellez le Midi Parasep® SF en le vissant dans le compartiment de cône de filtrage. Tourbillonez ou secouez pour émulsionner avec le cône de sédimentation pointé vers le haut.

Centrifugation
3 Retournez le Midi Parasep® SF et centrifugez le à 400g pendant 2 minutes. (J. Clin. Microbiol. doi:10.1128/JCM.00838-15)
Midi Parasep® SF s'adapte à tous les seaux de centrifugeuses 50ml.

RAPPEL: Calcul du nombre de tours par minute en fonction du rayon de la centrifugeuse.

$$RPM = \sqrt{\frac{g}{1.12r}} \times 1000$$

RPM tours par minute.
g accélération (max.1000g)
r rayon de la centrifugeuse en mm (depuis l'axe central jusqu'à la pointe du cône)

Examination

4A Dévissez et jetez le filtre et le tube de mélange.

4B Décantez tout le liquide au-dessus du sédiment.

4C Pipeter une goutte de sédiment sur une lame de microscope et couvrir avec une lamelle couvre-objet. Vous pouvez également suivre la procédure opératoire standard du laboratoire pour la préparation des lames de microscope.

HR

Pogledajte naljepnicu za uvjete čuvanja i rok valjanosti.
Molimo pridržavajte se sljedećih smjernica prilikom rukovanja Midi Parasep®-om. Kako biste izbjegli kros-kontaminaciju Midi Parasep® SF bi trebao biti zatvoren cijelo vrijeme osim kada stavljate uzorak ili prilikom uzimanja krajnjeg koncentriranog uzorka za mikroskopiranje.

Pripema Uzorka
Ako koristite napunjcnc Midi Parasep® SF, započeti u 1D.

1A Otvorite poklopac.

1B Dodajte 8,0 ml fiksatora.

1C Dodajte Apacor Triton X otopina u komoru za miješanje.

1D Uzmite žličicu uzorka na fixativu koristeći žličicu na kraju Midi Parasep® SF filtera. Dobro promiješajte s Midi Parasep® SF žličicom. Ako je uzorak tvrd, razbijte ga s krajem žličice.

Emulgiranje
2 Zatvorite Midi Parasep® SF tako da umetnete filter / sedimentacijski konus. Vorteksirajte ili protresite kako bi emulgirali sa sedimentacijskim konusom prema gore.

Centrifugiranje
3 Okrenite Midi Parasep® SF i centrifugirajte na 400g 2 minute. (J. Clin. Microbiol. doi:10.1128/JCM.00838-15)
Midi Parasep® SF odgovara svim 50 ml adapterima za centrifuge.

NAPOMENA: Preračunavanje RPM Za Svaku Centrifugu

$$RPM = \sqrt{\frac{g}{1.12r}} \times 1000$$

RPM brzina rotora u okr./min.
g centrifugalna sila (max. 1000g)
r radijus, horizontalna udaljenost izmedu sedimentacijskog konusa i centra vrtnje mjerena u mm

Pregled

4A Otvorite i bacite filter i komoru za miješanje.

4B Odlijte svu tekućinu iznad sedimenta.

4C Pipetirajte jednu kapljicu sedimenta na mikroskopski klizač i pokrijte pokrivačem. Alternativno, slijedite laboratorijske standardne postupke za pripremu mikroskopa.

IT

Leggere le indicazioni dell etichetta su conservazione e data di scaden. Si prega di seguire le seguenti avvertenze quando si utilizza il kit Midi Parasep® SF. Per evitare cross-contaminazioni il concentratore Midi Parasep® SF dovrebbe rimanere sempre chiuso tranne quando si debba introdurre il campione o quando debba essere recuperato il campione dopo la concentrazione (sedimento) per la successiva analisi.

Preparazione del campione

Se si utilizza Midi Parasep® SF preriempito, iniziare dal punto 1D.

1A Svitare il tappo.

1B Aggiungere 8.0 ml di fissativo

1C Se richiesto, aggiungere una goccia di surfattante (es. soluzione Apacor Triton X) per emulsionare.

1D Introdurre un campione fecale della dimensione di un pisello al fissatore. Mescolare bene con il cucchiaio del Midi Parasep® SF. Se il campione è duro, romperlo con l'estremità del cucchiaio.

Omogenizzazione

2 Chiudere ermeticamente il Midi Parasep® SF avvitando sul flacone di raccolta il cono di sedimentazione connesso con il filtro. Agitare a mano o con il vortex con il cono di sedimentazione rivolto verso líalto.

Centrifugazione

3 Invertire il Midi Parasep® SF e centrifugare a 400g per 2 minuti. (J. Clin. Microbiol. doi:10.1128/JCM.00838-15)
Il Midi Parasep® SF si adatta a tutte le centrifughe con rotori per provette da 50 ml.

Nota: per tutti i tipi di centrifuga la conversione da g a RPM avviene tramite questa formula:

$$RPM = \sqrt{\frac{g}{1.12r}} \times 1000$$

RPM Velocità del rotore in giri/ minuto
g Forza centrifuga (massimo 1000g)
r Raggio, distanza orizzontale tra la punta del cono di sedimentazione e il centro del rotore misurato in mm

Esame del campione

4A Svitare la camera di miscelazione annessa al filtro ed eliminarla.

4B Eliminare il sovranatante.

4C Pipetta una goccia di sedimento su di un vetrino e coprire con coprioggetto. In alternativa, seguire la procedura operativa standard di laboratorio per la preparazione vetrino da microscopio.

NL

Zie etiket voor bewaring en vervaldatum. Houdt u aan de volgende richtlijnen bij het omgaan met Midi Parasep® SF. Om kruisbesmetting te voorkomen, moet de Midi Parasep® SF altijd gesloten blijven, behalve bij het invoeren van het staal of bij het ophalen van het definitieve geconcentreerde staal voor onderzoek.

Staalvoorbereiding

Bij gebruik van voorgevulde Midi Parasep® SF, beginnen bij 1D.

1A Schroeft u het deksel los.

1B Voeg 8,0 ml fixatief toe.

1C Voegt een druppel surfactant (bv: Apacor Triton X oplossing) om te emulgeren.

1D Breng een kleine hoeveelheid feces monster ann het fixatief in de monsterbuis. Meng grondig met de Midi Parasep® SF lepel. Als het monster hard is, breek het dan open met de lepel.

Emulsificatie

2 Sluit de Midi Parasep® SF af door de filter unit in de sedimentatie kegel te schroeven. Schud met de sedimentatie kegel naar boven gericht om te emulgeren.

Centrifugatie

3 Keer de Midi Parasep® SF om en centrifugeer aan 400g gedurende twee minuten.
(J. Clin. Microbiol. doi:10.1128/JCM.00838-15)
Midi Parasep® SF past op alle 50 ml centrifuges.

Voor het berekenen van de benodigde RPM voor een centrifuge

$$RPM = \sqrt{\frac{g}{1.12r}} \times 1000$$

RPM Rotor snelheid in toeren per minuut
g Centrifugale kracht (maximaal 1000g)
r Radius, horizontale afstand tussen centrum van de centrifuge en de tip van de buis, gemeten in mm.

Onderzoek

4A Schroef los en gooi de mengkamer en filter weg.

4B Giet alle vloeistof weg die zich boven het sediment bevindt.

4C Pipet een druppel sediment op een microscoopglaasje en bedek met dekglaasje. Alternatief volg laboratorium standaard werkprocedure voor microscoopglaasje bereiden.

PL

Warunki przechowywania oraz data ważności zestawu.
na etykiecie. Proszę uważnie przeczytać instrukcję wykonania oznaczenia (Midi Parasep® SF) a następnie postępować z jej zaleceniami. Aby uniknąć przypadkowego zanieczyszczenia fiolka powinna być zamknięta przez cały czas przechowywania. Fiolkę Midi Parasep® SF otwieramy podczas pobierania próbki oraz podczas analizy zatężonego materiału biologicznego.

Przygotowanie próbki
Jeśli używasz 'prefilled' Midi Parasep® SF, zaczynaij od 1D.

1A Otworzyć probówkę a następnie.

1B Dodaj 8.0ml utrwalacza.

1C Oraz 1 kroplę surfaktantu (np. Apacor Triton X rozwiązanie).

1D Pobierz próbkę kału wielkości ziarna grochu do utrwalacza. Wymieszaj dokładnie próbkę w probówce Midi Parasep® SF. Jeśli próbka jest zbyt twarda, rozdrobnij ją końcem łyżeczki.

Przygotowanie emulsji
2 Połączyć ze sobą dwie części probówki Midi Parasep® SF (1: część wirówkowa probówki zaopatrzona w łopatkę oraz filtr; 2 część probówki z roztworem oraz materiałem biologicznym). Dokładnie wymieszać zawartość probówki (część stożkowa powinna być skierowana ku górze).

Wirowanie
3 Probówkę Midi Parasep® SF wirować przy 400 g przez 2 minut. (J. Clin. Microbiol. doi:10.1128/JCM.00838-15)
Midi Parasep® SF pasuje do wszystkich 50ml wiadra wirówki.

UWAGA: Dla każdej wirówki należy obliczyć prędkość wirowania.

$$RPM = \sqrt{\frac{g}{1.12r}} \times 1000$$

RPM Prędkość wirowania (obroty/min)
g siła odśrodkowa (maksimum 1000g)
r Promień ramienia rotora

Pobranie przygotowanej próbki do badań

4A Część stożkowa zawiera przygotowany do badań materiał. Drugą część probówki (część filtrującą) zawierającą zanieczyszczenia należy zutylizować.

4B Następnie ostrożnie zlać nadsącz (materiał nie związany w osadzie oraz płyn pozostający nad osadem).

4C Pipetuj jedną kropelkę osadu na szynę mikroskopową i przykryj szkiełko nakrywkowe. Alternatywnie, postępuj zgodnie z laboratoryjnymi standardowymi procedurami działania preparatu slajdów mikroskopowych.

PT

Veja as condições de armazenamento e a data de validade na etiqueta. Quando manusear o Midi Parasep® SF deve seguir as instruções de utilização. Para evitar contaminações cruzadas o Midi Parasep® SF deve permanecer sempre fechado, excepto quando introduz a amostra ou quando extrai a preparação final para ser examinada.

Preparação da amostra
Se estiver a utilizar o Midi Parasep® SF pré-cheio, comece em 1D.

1A Desenroscar a tampar.

1B Adicione 8,0ml de fixador.

1C Se necessário uma gota de surfactante (solução Apacor Triton X) para emulsionar.

1D Introduza uma amostra de fezes do tamanho de uma ervilha no fixador. Misture cuidadosamente com a colher Midi Parasep® SF. Se a amostra for dura, parta-a com a ponta da colher.

Emulsão
2 Enroscar a câmara de mistura com a unidade de filtro/cone de sedimentação. Agitar no vortex para emulsionar com o cone de sedimentação apontando para cima.

Centrifugação
3 Inverter o Midi Parasep® SF e centrifugar a 400g por 2 minutos. (J. Clin. Microbiol. doi:10.1128/JCM.00838-15)
Midi Parasep® SF é adequado a todos os copos de centrifuga de 50ml.

Nota: Para calcular as RPM para qualquer centrifuga

$$RPM = \sqrt{\frac{g}{1.12r}} \times 1000$$

RPM Velocidade do rotor
g Força centrifuga (máximo 1000g)
r Raio, distância entre a ponta do cone e o centro do rotor medida em mm

Visualização

4A Desenrosque e elimine a câmara de mistura juntamente com o filtro.

4B Decante o liquido sobrenadante do sedimento.

4C Pipetar uma gota de sedimento em uma lâmina de microscópio e cobrir com lamínula. Alternativamente, siga o procedimento operacional padrão do laboratório para preparação de lâminas de microscópio.

SI

Shranjevanje in rok uporabe: glej nalepko!
Prosimo, da pri uporabi koncentratorja Midi Parasep® SF upoštevate naslednja priporočila. Koncentrator Midi Parasep® SF naj bo vedno zaprt. Odprite ga samo med dodajanjem vzorca blata in odvzemom koncentriranega vzorca za mikroskopsko analizo. S tem preprečite navzkrižno kontaminacijo.

Priprava vzorca
(V primeru, da uporabljate pripravljene koncentratorje, napolnjene z reagenti, začnete pri točki 1D)

1A Odvijte zamašek koncentratorja.

1B Dodajte 8,0 ml fiksativa.

1C Ter 1 kapljico surfaktanta (npr. Apacor Triton X raztopina).

1D Uvesti fekalni vzorec z grahami v fiksacijo. Temeljito premešajte z Midi Parasep® SF žličko. Če je vzorec pretrd, ga s konico žličke razdrobite.

Emulzifikacija
2 Koncentrator tesno zaprite in ga premešajte ročno ali z vortexom. Filtrirni del z vzorcem naj bo pri tem obrnjen navzdol.

Centrifugiranje
3 Koncentrator obrnite in ga centrifugirajte 2 minuti pri 400g. (J. Clin. Microbiol. doi:10.1128/JCM.00838-15)
Koncentrator ustreza vsem 50 ml nastavkom v centrifugah.

Opomba: Za izračun potrebne hitrosti (obrati na minuto), lahko za katerokoli centrifugo, uporabite naslednjo formulo

$$RPM = \sqrt{\frac{g}{1.12r}} \times 1000$$

RPM Hitrost rotorja v obratih na minuto
g Centrifugalna sila (maks. 1000g)
r polmer, razdalja med konico koncentratorja in osjo rotorja, merjena v mm

Pregled vzorca

4A Filtrirni del koncentratorja odvijte in zavrzite.

4B Vso tekočino nad sedimentom odlijte.

4C Potopite eno kapljico usedline na drsnik za mikroskop in pokrijte s pokrovom. Druga možnost je, da sledite laboratorijskemu standardnemu obratovalnemu postopku za pripravo mikroskopskega drsnika.

ALCORFIX™ SAFETY DATA SHEET

This Safety Datasheet complies with the requirements of Regulation (EC) No 1907/2006

SECTION 1 IDENTIFICATION OF THE SUBSTANCE/ MIXTURE AND THE COMPANY/UNDERTAKING

1.1 Product Identifier: AlcorFix™
108810, 108885, 108886, 108887, 108889, 149995, 248200, 249200, 249300, 249420

1.2 Relevant identified uses of the substance or mixture and uses advised against: Solution for fixation/conservation of biological samples.

1.3 Details of the supplier of the Safety Data Sheet:
Apacor Limited, Unit 5 Sapphire Centre, Fishponds Road, Wokingham, Berkshire, RG41 2QL, England
+44 (0) 118 979 5566
technical@apacor.com

1.4 Emergency telephone number:
+44 (0)118 979 5566
(Monday-Friday 0900-1700 excluding UK Public Holidays)

SECTION 2 HAZARDS IDENTIFICATION

2.1 Classification of the substance or mixture

2.1.1 Classification according to Regulation (EC) No 1272/2008 [CLP]:
Acute toxicity, Oral (Category 4), H302
Acute toxicity, Inhalation (gas) (Category 4), H332
Serious eye damage (Category 1), H318
Hazardous to the aquatic environment (Category 2), H411
Flammable liquids (Category 2), H225

See Section 16 for the full text of H-Statements mentioned in this Section.

2.2 Label elements
Labelling according to Regulation (EC) No 1272/2008 [CLP]

Pictogram 
Signal word Danger

Hazard statement(s)
H225 – Highly flammable liquid and vapour
H302 - Harmful if swallowed
H318 - Causes serious eye damage
H332 - Harmful if inhaled
H411 - Toxic to aquatic life with long lasting effects

Precautionary statements:
P210 - Keep away from heat, hot surfaces, sparks, open flames and other ignition sources. No smoking.
P280 - Wear protective gloves/protective clothing/eye protection/face protection.
P305 + P351 + P338 - IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.
P310 - Immediately call a POISON CENTER or doctor/ physician.
P370 + P378 - In case of fire: Use dry sand, carbon dioxide (CO_2), water spray, dry chemical or alcohol resistant foam to extinguish.

2.3 Other hazards
None.

SECTION 3 COMPOSITION/INFORMATION ON INGREDIENTS

3.1 Substances

3.2 Mixtures

Hazardous ingredients according to Regulation (EC) No 1272/2008
Component: **Ethanol**
CAS No: 64-17-5
EC No: 200-578-6
Index No: 603-002-00-5
Classification: Flam. Liq. 2 (H225)
Concentration: 25%

Component: **Zinc sulphate**
CAS No: 7733-02-0
EC No: 231-793-3
Index No: 030-006-00-9
Classification: Acute Tox. 4 (H302), Eye Dam. 1 (H318), Aquatic Acute 1 (H400), Aquatic Chronic 1 (H410)
Concentration: 7.9%

Component: **Acetic Acid**
CAS No: 64-19-7
EC No: 200-580-7
Index No: 607-002-00-6
Classification: Skin Corr. 1A (H314), Flam. Liq. 3 (H226)
Concentration: 4.8%

Component: **Isopropanol**
CAS No: 67-63-0
EC No: 200-661-7
Index No: 603-117-00-0
Classification: Eye Irrit. 2 (H319), STOT SE 3 (H336), Flam. Liq. 2 (H225)
Concentration: 1%

Component: **Methyl Alcohol**
CAS No: 67-56-1
EC No: 200-659-6
Index No: 603-001-00-X
Classification: Acute Tox. 3 (H301), Acute Tox. 3 (H311), Acute Tox 3. (H331), STOT SE 1 (H370), Flam. Liq. 2 (H225)
Concentration: 1%

3.3 Other Information
Additional non-hazardous ingredients:
Polyvinyl alcohol (minimum 1g/l)
DI water

SECTION 4 FIRST AID MEASURES

4.1 Description of first aid measures
Consult a physician. Show this safety data sheet to the doctor in attendance.
In case of eye contact: Rinse thoroughly with plenty of water for at least 15 minutes and consult a physician.

In case of skin contact: Wash off immediately with soap and plenty of water while removing all contaminated clothes and shoes.
If swallowed: Clean mouth with water and drink afterwards plenty of water.
If inhaled: Move to fresh air.

4.2 Most important symptoms and effects, both acute and delayed

No information available.

4.3 Indication of any immediate medical attention and special treatment needed

Notes to physician: Treat symptomatically.

SECTION 5 FIRE FIGHTING MEASURES

5.1 Extinguishing media

Suitable extinguishing media (use media appropriate to the circumstances and environment): dry sand, carbon dioxide (CO2), water spray, alcohol-resistant foam, dry chemical.

5.2 Special hazards arising from the substance or mixture

No information available

5.3 Advice for firefighters

As in any fire, wear self-contained breathing apparatus, MSHA/NIOSH (approved or equivalent) and full protective gear.

SECTION 6 ACCIDENTAL RELEASE MEASURES

6.1 Personal precautions, protective equipment and emergency procedures

Ensure adequate ventilation, especially in confined areas.

6.2 Environmental precautions

Should not be released into the environment. Prevent product from entering drains. Prevent further leakage or spillage if safe to do so.

6.3 Methods and material for containment and cleaning up

Absorb spill with inert material (eg dry sand or earth), then place in a chemical waste container. After cleaning, flush away traces with water.

6.4 Reference to other sections

For disposal, see Section 13.

SECTION 7 HANDLING AND STORAGE

7.1 Precautions for safe handling

Do not breathe vapours or spray mist. Ensure that ventilation is adequate before using this product. Avoid contact with skin and eyes. Take necessary personal protective precautions before using this product. Keep away from heat and flame. Take precautionary measures against static discharges.

7.2 Conditions for safe storage, including any incompatibilities

Technical measures/precautions: Store in a tightly closed container. Store in a cool, dry, well-ventilated area away from incompatible substances.
Incompatible products: Avoid strong bases. Oxidizing agent.

7.3 Specific end use(s)

No other specific end uses(s) are specified apart from those listed in Section 1.2.

SECTION 8 EXPOSURE CONTROLS/PERSONAL PROTECTION

8.1 Control parameters

Component	Ethanol 64-17-5	Zinc sulphate 7733-02-0	Acetic Acid 64-19-7	Isopropanol 67-63-0	Methyl Alcohol 67-56-1
UK	STEL: 3000 ppm STEL: 5760 mg/m³ TWA: 1000 ppm TWA: 1920 mg/m³			STEL: 500 ppm STEL: 1250 mg/m³ TWA: 400 ppm TWA: 999 mg/m³	STEL: 250 ppm STEL: 333 mg/m³ TWA: 200 ppm TWA: 266 mg/m³ Skin
France	TWA: 1000 ppm TWA: 1900 mg/m³ STEL: 5000 ppm STEL: 9500 mg/m³		STEL: 10 ppm STEL: 25 mg/m³	STEL: 400 ppm STEL: 980 mg/m³	TWA: 200 ppm TWA: 260 mg/m³ STEL: 1000 ppm STEL: 1300 mg/m³
Spain	STEL: 1000 ppm STEL: 1910 mg/m³		STEL: 15 ppm STEL: 37 mg/m³ TWA: 10 ppm TWA: 25 mg/m³	STEL: 400 ppm STEL: 1000 mg/m³ TWA: 200 ppm TWA: 500 mg/m³	S* TWA: 200 ppm TWA: 266 mg/m³
Germany	TWA: 500 ppm TWA: 960 mg/m³ Ceiling / Peak: 1000 ppm Ceiling / Peak: 1920 mg/m³Skin	TWA: 0.1 mg/m³ TWA: 2 mg/m³ Ceiling / Peak: 0.4 mg/m³ Ceiling / Peak: 4 mg/m³	TWA: 10 ppm TWA: 25 mg/m³ Ceiling / Peak: 20 ppm Ceiling / Peak: 50 mg/m³	TWA: 200 ppm TWA: 500 mg/m³ Ceiling / Peak: 400 ppm Ceiling / Peak: 1000 mg/m³	TWA: 200 ppm TWA: 270 mg/m³ Ceiling / Peak: 800 ppm Ceiling / Peak: 1080 mg/m³ Skin
Italy					TWA: 200 ppm TWA: 260 mg/m³ Skin
Portugal	TWA: 1000 ppm		STEL: 15 ppm TWA: 10 ppm TWA: 25 mg/m³	STEL: 400 ppm TWA: 200 ppm	STEL: 250 ppm TWA: 200 ppm TWA: 260 mg/m³
The Netherlands	Skin STEL: 1900 mg/m³ TWA: 260 mg/m³				Skin TWA: 133 mg/m³ TWA: 100 ppm
Finland	TWA: 1000 ppm TWA: 1900 mg/m³ STEL: 1300 ppm STEL: 2500 mg/m³		TWA: 5 ppm TWA: 13 mg/m³ STEL: 10 ppm STEL: 25 mg/m³	TWA: 200 ppm TWA: 500 mg/m³ STEL: 250 ppm STEL: 620 mg/m³	TWA: 200 ppm TWA: 270 mg/m³ STEL: 250 ppm STEL: 330 mg/m³ Skin
Denmark	TWA: 1000 ppm TWA: 1900 mg/m³		TWA: 10 ppm TWA: 25 mg/m³	TWA: 200 ppm TWA: 490 mg/m³	TWA: 200 ppm TWA: 260 mg/m³ Skin
Austria	STEL 2000 ppm STEL 3800 mg/m³ TWA: 1000 ppm TWA: 1900 mg/m³		STEL 20 ppm STEL 50 mg/m³ TWA: 10 ppm TWA: 25 mg/m³	STEL 800 ppm STEL 2000 mg/m³ TWA: 200 ppm TWA: 500 mg/m³	Skin STEL 800 ppm STEL 1040 mg/m³ TWA: 200 ppm TWA: 260 mg/m³
Switzerland	STEL: 1000 ppm STEL: 1920 mg/m³ TWA: 500 ppm TWA: 960 mg/m³	STEL: 4 mg/m³ TWA: 0.1 mg/m³ TWA: 2 mg/m³	STEL: 20 ppm STEL: 50 mg/m³ TWA: 10 ppm TWA: 25 mg/m³	STEL: 400 ppm STEL: 1000 mg/m³ TWA: 200 ppm TWA: 500 mg/m³	Skin STEL: 800 ppm STEL: 1040 mg/m³ TWA: 200 ppm TWA: 260 mg/m³
Poland	TWA: 1900 mg/m3		STEL: 30 mg/m3 TWA: 15 mg/m3	STEL: 1200 mg/m3 TWA: 900 mg/m3	STEL: 300 mg/m3 TWA: 100 mg/m3
Norway	TWA: 500 ppm TWA: 950 mg/m3 STEL: 500 ppm STEL: 950 mg/m3		TWA: 10 ppm TWA: 25 mg/m3 STEL: 20 ppm STEL: 37.5 mg/m3	TWA: 100 ppm TWA: 245 mg/m3 STEL: 150 ppm STEL: 306.25 mg/m3	TWA: 100 ppm TWA: 130 mg/m3 Skin STEL: 150 ppm STEL: 162.5 mg/m3
Ireland	STEL: 1000 ppm		TWA: 10 ppm TWA: 25 mg/m3 STEL: 15 ppm STEL: 37 mg/m3	TWA: 200 ppm STEL: 400 ppm Skin	TWA: 200 ppm TWA: 260 mg/m3 STEL: 600 ppm STEL: 780 mg/m3 Skin
European Union			TWA 10 ppm TWA 25 mg/m3		TWA: 200 ppm TWA: 260 mg/m3 Skin

Derived No Effect Level (DNEL) No information available

Predicted No Effect Concentration (PNEC) No information available

8.2 Exposure controls

Engineering measures: Ensure adequate ventilation, especially in confined areas.
Personal protective equipment
Respiratory protection: No special protective equipment required.
Hand protection: Wear appropriate protective gloves.
Eye protection: Wear tightly fitting safety goggles or safety glasses with side-shields.
Skin and body protection: Protective clothing to protect exposed skin.
Hygiene measures: Handle in accordance with good industrial hygiene and safety practice.
Environmental exposure controls: No information available.

SECTION 9 PHYSICAL AND CHEMICAL PROPERTIES

9.1 Information on basic physical and chemical properties

a) Appearance: clear liquid

b) Odour: pungent

c) Odour threshold: no information available

d) pH: no information available

e) Melting point / freezing point: no information available

f) Initial boiling point / boiling range: 84°C

g) Flash point: 16°C

h) Evaporation rate: no information available

i) Flammability (solid, gas): no information available

j) Upper/lower flammability or explosive limits: no information available

k) Vapour pressure: no information available

l) Vapour density: no information available

m) Relative density: no information available

n) Solubility (ies) : soluble in water

o) Partition coefficient: n-octanol/water: no information available

p) Auto-ignition temperature: no information available

q) Decomposition temperature: no information available

r) Viscosity: no information available

s) Explosive properties: no information available

t) Oxidising properties: no information available

9.2 Other information: no information available

SECTION 10 STABILITY AND REACTIVITY

10.1 Reactivity

10.2 Chemical stability

Stable under normal conditions.

10.3 Possibility of hazardous reactions

No information available.

10.4 Conditions to avoid

Heat, flames and sparks.

10.5 Incompatible materials

No particular materials.

10.6 Hazardous decomposition products

Under normal use – none.

SECTION 11 TOXICOLOGICAL INFORMATION

11.1 Information of toxicological effects

Acute toxicity:

Product: based on known/supplied information, does not present an acute toxicity hazard.

Inhalation: no data available.

Eye contact: no data available.

Skin contact: no data available.

Ingestion: no data available.

≤ 60.3% of the mixture consists of ingredients of unknown toxicity.

The following values are calculated based on GHS document chapter 3.1.

Oral	1,363.00mg/kg	
Dermal	5,158.00mg/kg	
Inhalation:	Gas	4,263.00mg/l
	Mist	20.90mg/l
	Vapour	829.22mg/l

Chemical Name	LD50 Oral	LD50 Dermal	LC50 Inhalation
Ethanol	7060mg/kg (Rat)		124.7mg/L (Rat) 4 h
Zinc sulphate	500mg/kg (Rat)		
Acetic acid	3310mg/kg (Rat)	1060mg/kg (Rabbit)	11.4mg/L (Rat) 4 h
Methyl alcohol	6200mg/kg (Rat)	15800mg/kg (Rabbit)	22500 ppm (Rat) 8 h 64000 ppm (Rat) 4 h
Isopropanol	1870mg/kg (Rat)	4059mg/kg (Rabbit)	72600mg/m3 (Rat) 4 h

Skin corrosion/irritation: no data available

Serious eye damage/eye irritation: no data available

Respiratory or skin sensitisation: no data available

Germ cell mutagenicity: no data available

Carcinogenicity: no data available

Reproductive toxicity: no data available

Specific target organ toxicity - single exposure: no data available

Specific target organ toxicity - repeated exposure: no data available

Aspiration hazard: no data available

SECTION 12 ECOLOGICAL INFORMATION

12.1 Toxicity

Toxic to aquatic life with long lasting effects.

Chemical Name	Toxicity to Algae	Toxicity to Fish	Toxicity to Daphnia and other aquatic invertebrates
Ethanol		12.0 - 16.0: 96 h Oncorhynchus mykiss mL/L LC50 static 100: 96 h Pimephales promelas mg/L LC50 static 13400 - 15100: 96 h Pimephales promelas mg/L LC50 flow-through	9268 - 14221: 48 h Daphnia magna mg/L LC50 2: 48 h Daphnia magna mg/L EC50 Static 10800: 24 h Daphnia magna mg/L EC50
Zinc sulphate	0.056: 72 h Pseudokirchneriella subcapitata mg/L EC50 static 64.8: 72 h Chlorella vulgaris mg/L EC50 2.4: 96 h Chlorella vulgaris mg/L EC50	0.162: 96 h Oncorhynchus mykiss mg/L LC50 flow-through 0.03 - 0.05: 96 h Oncorhynchus mykiss mg/L LC50 semi-static 0.34 - 0.93: 96 h Oncorhynchus mykiss mg/L LC50 static 0.218 - 0.42: 96 h Pimephales promelas mg/L LC50 flow-through 0.06: 96 h Pimephales promelas mg/L LC50 static 0.23 - 0.48: 96 h Pimephales promelas mg/L LC50 0.168 - 0.25: 96 h Pimephales promelas mg/L LC50 semi-static 0.15: 96 h Cyprinus carpio mg/L LC50 semi-static 16.85 - 27.18: 96 h Cyprinus carpio mg/L LC50 static 3 - 4.6: 96 h Lepomis macrochirus mg/L LC50 flow-through 3.55 - 6.32: 96 h Lepomis macrochirus mg/L LC50 static 0.63: 96 h Poecilia reticulata mg/L LC50 49.23 - 64.16: 96 h Poecilia reticulata mg/L LC50 semi-static 0.48 - 1.72: 96 h Poecilia reticulata mg/L LC50 static	0.75: 48 h Daphnia magna mg/L EC50 0.538 - 0.908: 48 h Daphnia magna mg/L EC50 Static
Acetic acid		79: 96 h Pimephales promelas mg/L LC50 static 75: 96 h Lepomis macrochirus mg/L LC50 static	65: 48 h Daphnia magna mg/L EC50 Static 47: 24 h Daphnia magna mg/L EC50
Isopropanol	1000: 96 h Desmodesmus subspicatus mg/L EC50 1000: 72 h Desmodesmus subspicatus mg/L EC50	9640: 96 h Pimephales promelas mg/L LC50 flow-through 11130: 96h Pimephales promelas mg/L LC50 static 1400000: 96 h Lepomis macrochirus µg/L LC50	13299: 48 h Daphnia magna mg/L EC50
Methyl alcohol		28200: 96 h Pimephales promelas mg/L LC50 flow-through 100: 96 h Pimephales promelas mg/L LC50 static 19500 - 20700: 96 h Oncorhynchus mykiss mg/L LC50 flow-through 18 - 20: 96 h Oncorhynchus mykiss mL/L LC50 static 13500 - 17600: 96 h Lepomis macrochirus mg/L LC50 flow-through	

12.2 Persistence and degradability
No information available.

12.3 Bioaccumulative potential
No information available.

Chemical Name	log Pow
Ethanol	-0.32
Acetic acid	-0.31
Isopropanol	0.05
Methyl alcohol	-0.77

12.4 Mobility in soil
No information available.

12.5 Results of PBT and vPvB assessment
No information available

12.6 Other adverse effects
No information available.

12.7 Additional information
No information available.

SECTION 13 DISPOSAL CONSIDERATIONS
13.1 Waste treatment methods
Waste from residues / unused products: In accordance with local and national regulations. Should not be released into the environment.
Contaminated packaging: Empty containers should be disposed of at an approved waste handling site for recycling or disposal.

SECTION 14 TRANSPORT INFORMATION
14.1 UN number: UN2924

14.2 UN proper shipping name: Flammable Liquid, Corrosive, n.o.s. (Ethanol, Acetic Acid)

14.3 Transport hazard class(es): 3, Subsidiary Class: 8

14.4 Packing group: II

14.5 Environmental hazards

14.6 Special precautions for user

14.7 Transport in bulk according to Annex II of MARPOL73/78 and the IBC Code
Not intended to be transported in bulk.

Note: Per 49 CFR – when shipping 30ml or less per inner packaging and the gross weight does not exceed 64lbs, use the 173.4 small quantity exception.

SECTION 15 REGULATORY INFORMATION
15.1 Safety, health and environmental regulations/ legislation specific for the substance or mixture

Chemical Name	French RG number
Ethanol	RG 84
Isopropanol	RG 84
Methyl alcohol	RG 84

TSCA	Complies
EINECS/ELINCS	-
DSL/NDSL	-
PICCS	-
ENCS	-
IECSC	-
AICS	-
KECL	-

Legend
TSCA - United States Toxic Substances Control Act Section 8(b) Inventory
EINECS/ELINCS - European Inventory of Existing Commercial Chemical Substances/EU List of Notified Chemical Substances
DSL/NDSL - Canadian Domestic Substances List/Non-Domestic Substances List
PICCS - Philippines Inventory of Chemicals and Chemical Substances
ENCS - Japan Existing and New Chemical Substances
IECSC - China Inventory of Existing Chemical Substances
AICS - Australian Inventory of Chemical Substances
KECL - Korean Existing and Evaluated Chemical Substances

15.2 Chemical Safety Assessment
No information available.

SECTION 16 OTHER INFORMATION
Full text of H-Statements referred to under sections 2 and 3
H225 - Highly flammable liquid and vapour
H226 - Flammable liquid and vapour
H301 - Toxic if swallowed
H302 - Harmful if swallowed
H311 - Toxic in contact with skin
H314 - Causes severe skin burns and eye damage
H318 - Causes serious eye damage
H319 - Causes serious eye irritation
H331 - Toxic if inhaled
H332 – Harmful if inhaled
H336 - May cause drowsiness or dizziness
H370 - Causes damage to organs.
H400 - Very toxic to aquatic life
H410 - Very toxic to aquatic life with long lasting effects
H411 - Toxic to aquatic life with long lasting effects

Amended sections are indicated by a line in the border.

The information supplied in this SDS is correct to the best of our knowledge. We do not accept any liability for loss, injury or damage, which may result from its use.

10% FORMALIN SAFETY DATA SHEET

This Safety Datasheet complies with the requirements of Regulation (EC) No 1907/2006

SECTION 1 IDENTIFICATION OF THE SUBSTANCE/ MIXTURE AND THE COMPANY/UNDERTAKING

1.1 **Product Identifier: 10% Formalin**

145200, 145300, 145400, 145420, 145700, 145800, 145900, 1460, 146200, 146300, 146400, 108900, 108910, 148980, 148998, 149910, 151000

1.2 **Relevant identified uses of the substance or mixture and uses advised against:** laboratory chemical (in vitro diagnostic)

1.3 **Details of the supplier of the Safety Data Sheet:**

Apacor Limited, Unit 5 Sapphire Centre, Fishponds Road, Wokingham, Berkshire, RG41 2QL, England
+44 (0) 118 979 5566
technical@apacor.com

1.4 **Emergency telephone number**:

+44 (0)118 979 5566
(Monday-Friday 0900-1700 excluding UK Public Holidays)

SECTION 2 HAZARDS IDENTIFICATION

2.1 Classification of the substance or mixture

Classification according to Regulation (EC) No 1272/2008 [CLP]:
Acute toxicity, Oral (Category 4), H302
Skin sensitisation (Category 1), H317
Acute toxicity, Inhalation (Category 4), H332
Germ cell mutagenicity (Category 2), H341
Carcinogenicity (Category 1B), H350

See Section 16 for the full text of H-Statements mentioned in this Section.

2.2 Label elements

Labelling according to Regulation (EC) No 1272/2008 [CLP]

Pictogram

Signal word **Danger**

Hazard statement(s)

H302 Harmful if swallowed
H317 May cause an allergic skin reaction.
H332 Harmful if inhaled.
H341 Suspected of causing genetic defects
H350 May cause cancer
Contains Formaldehyde.

Precautionary statements:

P280 Wear protective gloves/protective clothing/eye protection/face protection.
P301 + P312 + P330 IF SWALLOWED: Call a POISON CENTER or doctor/physician if you feel unwell. Rinse mouth.
P308 + P313 IF exposed or concerned: Get medical advice/ attention.
P333 + P313 If skin irritation or rash occurs: Get medical advice/attention.

See Section 16 for the full text of H-Statements mentioned in this Section.

2.3 Other hazards

This substance/mixture contains no components considered to be either persistent, bioaccumulative and toxic (PBT), or very persistent and very bioaccumulative (vPvB) at levels of 0.1% or higher.

SECTION 3 COMPOSITION/INFORMATION ON INGREDIENTS

3.2 Mixtures

Hazardous ingredients according to Regulation (EC) No 1272/2008

Component: **Formaldehyde**
CAS No: 50-00-0
EC No: 200-001-8
Index No: 605-001-00-5
Classification: Acute Tox. 3 (H301 + H311 + H331), Skin Corr. 1B (H314), Skin Sens. 1 (H317), Muta. 2 (H341), Carc. 1B (H350)
Concentration: < 5%

Component: **Methanol**
CAS No: 67-56-1
EC No: 200-659-6
Index No: 603-001-00-x
Registration No: 01-2119433307-44-xxxx
Classification: Flam. Liq. 2 (H225); Acute Tox. 3 (H301 + H311 + H331); STOT SE 1 H370
Concentration: < 1%

See Section 16 for the full text of H-Statements mentioned in this Section.

SECTION 4 FIRST AID MEASURES

4.1 Description of first aid measures

Consult a physician. Show this safety data sheet to the doctor in attendance.

If inhaled: If breathed in, move person into fresh air. If not breathing, give artificial respiration. Consult a physician.

In case of skin contact: Wash off with soap and plenty of water. Take victim immediately to hospital. Consult a physician.

In case of eye contact: Rinse thoroughly with plenty of water for at least 15 minutes and consult a physician.

If swallowed: Do NOT induce vomiting. Never give anything by mouth to an unconscious person. Rinse mouth with water. Consult a physician.

4.2 Most important symptoms and effects, both acute and delayed

The most important known symptoms and effects are described in the labelling (Section 2.2) and/or Section 11.

4.3 Indication of any immediate medical attention and special treatment needed

No data available.

SECTION 5 FIRE FIGHTING MEASURES

5.1 Extinguishing media

Suitable extinguishing media: Use water spray, alcohol-resistant foam, dry chemical or carbon dioxide.

5.2 Special hazards arising from the substance or mixture

Carbon oxides

5.3 Advice for firefighters

Wear self-contained breathing apparatus and full protective gear.

SECTION 6 ACCIDENTAL RELEASE MEASURES

6.1 Personal precautions, protective equipment and emergency procedures

Use personal protective equipment. Avoid breathing vapours, mist or gas. Ensure adequate ventilation. Remove all sources of ignition. Evacuate personnel to safe areas. Beware of vapours accumulating to form explosive concentrations. Vapours can accumulate in low areas. For personal protection see Section 8.

6.2 Environmental precautions

Prevent further leakage or spillage if safe to do so. Do not let product enter drains.

6.3 Methods and material for containment and cleaning up

Contain spillage, and then collect and place in container for disposal according to local regulations (see Section 13). Keep in suitable, closed containers for disposal.

6.4 Reference to other sections

For disposal, see Section 13.

SECTION 7 HANDLING AND STORAGE

7.1 Precautions for safe handling

Avoid contact with skin and eyes. Avoid inhalation of vapour or mist. Keep away from sources of ignition—no smoking. Take measures to prevent the build-up of electrostatic charge. For precautions see Section 2.2.

7.2 Conditions for safe storage, including any incompatibilities

Store in a cool place. Keep container tightly closed in a dry and well-ventilated place. Containers which are opened must be carefully resealed and kept upright to prevent leakage.

7.3 Specific end use(s)

No other specific uses are specified apart from those listed in Section 1.2.

SECTION 8 EXPOSURE CONTROLS/PERSONAL PROTECTION

8.1 Control parameters

Exposure limits: this product, as supplied, does not contain any hazardous materials with occupational exposure limits established by the region specific regulatory bodies.

	Formaldehyde 50-00-0	Methanol 67-56-1
Austria	STEL: 0.5 ppm STEL: 0.6 mg/m³ TWA: 0.5 ppm TWA: 0.6 mg/m³	STEL: 800 ppm STEL: 1040 mg/m³ TWA: 200 ppm TWA: 260 mg/m³
Belgium	STEL: 0.3 ppm STEL: 0.38 mg/m³	STEL: 250 ppm STEL: 333 mg/m³ TWA: 200 ppm TWA: 266 mg/m³
Denmark	STEL: 0.3 ppm STEL: 0.4 mg/m³ TWA: 0.3 ppm TWA: 0.4 mg/m³	STEL: 400 ppm STEL: 520 mg/m³ TWA: 200 ppm TWA: 260 mg/m³
France	TWA: 0.5 ppm STEL: 1 ppm	STEL: 1000 ppm STEL: 1300 mg/m³ TWA: 200 ppm TWA: 260 mg/m³
Germany	STEL: 0.6 ppm STEL: 0.74 mg/m³ TWA: 0.3 ppm TWA: 0.37 mg/m³	STEL: 800 ppm STEL: 1080 mg/m³ TWA: 200 ppm TWA: 270 mg/m³
Ireland	STEL: 2 ppm STEL: 2.5 mg/m³ TWA: 2 ppm TWA: 2.5 mg/m³	TWA: 200 ppm TWA: 260 mg/m³
Italy		TWA: 200 ppm TWA: 260 mg/m³
Poland	STEL: 1 mg/m³ TWA: 0.5 mg/m³	STEL: 300 mg/m³ TWA: 100 mg/m³
Portugal	STEL: 0.3 ppm	STEL : 250 ppm TWA : 200 ppm TWA : 260 mg/m³
Spain	STEL: 0.3 ppm STEL: 0.37 mg/m³	STEL: 250 ppm STEL: 333 mg/m³ TWA: 200 ppm TWA: 266 mg/m³
Sweden	STEL: 0.6 ppm STEL: 0.74 mg/m³ TWA: 0.3 ppm TWA: 0.37 mg/m³	STEL: 250 ppm STEL: 350 mg/m³ TWA: 200 ppm TWA: 250 mg/m³
The Netherlands	STEL: 0.5 mg/m³ TWA: 0.15 mg/m³	TWA: 133 mg/m³
UK	STEL: 2 ppm STEL: 2.5 mg/m³ TWA: 2 ppm TWA: 2.5 mg/m³	STEL: 250 ppm STEL: 333 mg/m³ TWA: 200 ppm TWA: 266 mg/m³

8.2 Exposure controls

8.2.1 Appropriate engineering controls

Handle in accordance with good industrial hygiene and safety practice. Wash hands before breaks and at the end of workday.

8.2.2 Personal protective equipment

(a) Eye/face protection: Tightly fitting safety goggles. Faceshield (8-inch minimum). Use equipment for eye protection tested and approved under appropriate government standards such as NIOSH (US) or EN 166(EU).

(b) Skin protection: Handle with gloves. Gloves must be inspected prior to use. Use proper glove removal technique (without touching glove's outer surface) to avoid skin contact with this product. Dispose of contaminated gloves after use in accordance with applicable laws and good laboratory practices. Wash and dry hands. The selected protective gloves should satisfy the specifications of EU Directive 89/686/EEC and the standard EN 374 derived from it.

(c) Body Protection: Complete suit protecting against chemicals. The type of protective equipment must be selected according to the concentration and amount of the dangerous substance at the specific workplace.

(d) Respiratory protection: Where risk assessment shows air-purifying respirators are appropriate use a full-face respirator with multi-purpose combination (US) or type ABEK (EN 14387) respirator cartridges as a backup to engineering controls. If the respirator is the sole means of protection, use a full-face supplied air respirator. Use respirators and components tested and approved under appropriate government standards such as NIOSH (US) or CEN (EU).

8.2.3 Environmental exposure controls
Prevent further leakage or spillage if safe to do so. Do not let product enter drains.

SECTION 9 PHYSICAL AND CHEMICAL PROPERTIES

9.1 Information on basic physical and chemical properties

a) Appearance Form: liquid

b) Odour no data available

c) Odour threshold no data available

d) pH no data available

e) Melting point / freezing point no data available

f) Initial boiling point and boiling range 100°C at 1.013 hPa

g) Flash point 85°C

h) Evaporation rate no data available

i) Flammability (solid, gas) no data available

j) Upper/lower flammability or explosive limits

Upper 70% (V), Lower 7% (V)

k) Vapour pressure 53hPa at 39°C

l) Vapour density no data available

m) Relative density 1.080g/cm^3

n) Solubility (ies) completely miscible

o) Partition coefficient: n-octanol/water no data available

p) Auto-ignition temperature no data available

q) Decomposition temperature no data available

r) Viscosity no data available

s) Explosive properties no data available

t) Oxidising properties no data available

9.2 Other information

No data available.

SECTION 10 STABILITY AND REACTIVITY

10.1 Reactivity

No data available.

10.2 Chemical stability

Stable under recommended storage conditions.

10.3 Possibility of hazardous reactions

No data available.

10.4 Conditions to avoid

Heat, flames and sparks.

10.5 Incompatible materials

No materials to be mentioned in particular.

10.6 Hazardous decomposition products

Carbon oxides.

SECTION 11 TOXICOLOGICAL INFORMATION

11.1 Information of toxicological effects

Acute toxicity: no data available
Skin corrosion/irritation: no data available
Serious eye damage/eye irritation: no data available
Respiratory or skin sensitisation: no data available
Germ cell mutagenicity: no data available
Carcinogenicity: IARC: 1 - Group 1: Carcinogenic to humans (Formaldehyde)
Reproductive toxicity: no data available
Specific target organ toxicity - single exposure: no data available
Specific target organ toxicity - repeated exposure: no data available
Aspiration hazard: no data available

Additional Information

Chemical Name	
Formaldehyde	LD50 oral 600mg/kg (Rat) LD50 dermal 270mg/kg (Rabbit) LC50 inhalation 0.578mg/L (Rat) 4 h
Methanol	LD50 oral - rat - 5628mg/kg LC50 inhalation - rat - 4h – 83.2mg/l/4h

SECTION 12 ECOLOGICAL INFORMATION

12.1 Toxicity

Ecotoxicity effects: contains no substances known to be hazardous to the environment or not degradable in waste water treatment plants.

Toxicity to Fish	
Formaldehyde	0.032 - 0.226: 96 h Oncorhynchus mykiss mL/L LC50 flow-through 100- 136: 96 h Oncorhynchus mykiss mg/L LC50 static 1510: 96 h Lepomis macrochirus µg/L LC50 static 22.6 - 25.7: 96 h Pimephales promelas mg/L LC50 flow-through 23.2 - 29.7: 96 h Pimephales promelas mg/L LC50 static 41: 96 h Brachydanio rerio mg/L LC50 static
Methanol	LC50 - Pimephales promelas – 28200mg / L 96h

Toxicity to Daphnia and other Aquatic Invertebrates	
Formaldehyde	11.3 - 18: 48 h Daphnia magna mg/L EC50 Static 2: 48 h Daphnia magna mg/L LC50
Methanol	EC50 - Daphnia magna - >10000mg/l

12.2 Persistence and degradability

No data available.

12.3 Bioaccumulative potential

No data available.

Chemical Name	log Pow
Formaldehyde	0.35
Methanol	-0.77

12.4 Mobility in soil

No data available.

12.5 Results of PBT and vPvB assessment

This substance/mixture contains no components considered to be either persistent, bioaccumulative and toxic (PBT), or very persistent and very bioaccumulative (vPvB) at levels of 0.1% or higher.

12.6 Other adverse effects

No data available.

12.7 Additional information

None.

SECTION 13 DISPOSAL CONSIDERATIONS

13.1 Waste treatment methods

Product: Dispose of in accordance with all federal, state, and local regulations. This combustible material may be burned in a chemical incinerator equipped with an afterburner and scrubber. Offer surplus and non-recyclable solutions to a licensed disposal company.

Contaminated packaging: Dispose of as unused product.

SECTION 14 TRANSPORT INFORMATION

IATA/DOT/IMDG/TDG: Not regulated.

14.1 UN number: -

14.2 UN proper shipping name: -

14.3 Transport hazard class(es): -

14.4 Packing group: -

14.5 Environmental hazards: -

14.6 Special precautions for user: -

SECTION 15 REGULATORY INFORMATION

15.1 Safety, health and environmental regulations/ legislation specific for the substance or mixture

No data available.

15.2 Chemical Safety Assessment

No chemical safety assessment has been carried out for this product.

SECTION 16 OTHER INFORMATION

Full text of H-Statements referred to in Sections 2 and 3

H225 Highly flammable liquid and vapour.

H301 + H311 + H331 Toxic if swallowed, in contact with skin or if inhaled

H302 Harmful if swallowed.

H314 Causes severe skin burns and eye damage.

H317 May cause an allergic skin reaction.

H332 Harmful if inhaled.

H341 Suspected of causing genetic defects.

H350 May cause cancer.

H370 Causes damage to organs.

Acute Tox. Acute toxicity

Carc. Carcinogenicity

Flam. Liq. Flammable liquids

Muta. Germ cell mutagenicity.

Skin Corr. Skin corrosion

Skin Sens. Skin sensitisation

STOT SE Specific target organ toxicity - single exposure

Amended sections are indicated by a line in the border.

The information supplied in this SDS is correct to the best of our knowledge. We do not accept any liability for loss, injury or damage, which may result from its use.

SAF SAFETY DATA SHEET

This Safety Datasheet complies with the requirements of Regulation (EC) No 1907/2006

SECTION 1 IDENTIFICATION OF THE SUBSTANCE/ MIXTURE AND THE COMPANY/UNDERTAKING

1.1 Product Identifier: SAF
(Sodium Acetate-Acetic Acid-Formalin Solution)
145500, 1461, 1461A, 146500, 146501, 108920, 148960, 148965, 149920, 149960, 153000, 249400

1.2 Relevant identified uses of the substance or mixture and uses advised against: laboratory chemical (in vitro diagnostic)

1.3 Details of the supplier of the Safety Data Sheet:
Apacor Limited, Unit 5 Sapphire Centre, Fishponds Road, Wokingham, Berkshire, RG41 2QL, England
+44 (0) 118 979 5566
technical@apacor.com

1.4 Emergency telephone number:
+44 (0)118 979 5566
(Monday-Friday 0900-1700 excluding UK Public Holidays)

SECTION 2 HAZARDS IDENTIFICATION

2.1 Classification of the substance or mixture

Classification according to Regulation (EC) No 1272/2008 [CLP]:
Acute toxicity, Oral (Category 4), H302
Skin sensitisation (Category 1), H317
Acute toxicity, Inhalation (Category 4), H332
Germ Cell Mutagenicity (Category 2), H341
Carcinogenicity (Category 1B), H350

See Section 16 for the full text of H-Statements mentioned in this Section.

2.2 Label elements
Labelling according to Regulation (EC) No 1272/2008 [CLP]

Pictogram

Signal word **Danger**

Hazard statement(s)
H302 Harmful if swallowed.
H317 May cause an allergic skin reaction.
H332 Harmful if inhaled.
H341 Suspected of causing genetic defects.
H350 May cause cancer.
Contains Formaldehyde

Precautionary statements:
P280 Wear protective gloves/protective clothing/eye protection/face protection.
P301 + P310 - IF SWALLOWED: Immediately call a POISON CENTER or doctor/ physician
P308 + P313 - IF exposed or concerned: Get medical advice/ attention

2.3 Other hazards
No data available.

SECTION 3 COMPOSITION/INFORMATION ON INGREDIENTS

3.2 Mixtures

Hazardous ingredients according to Regulation (EC) No 1272/2008
Component: **Formaldehyde**
CAS No: 50-00-0
EC No: 200-001-8
Index No: 605-001-00-5
Classification: Acute Tox. 3 (H301 + H311 + H331); Skin Corr. 1B (H314); Skin Sens. 1 (H317); Muta. 2 (H341); Carc. 1B (H350);
Concentration: < 5%

Component: **Methanol**
CAS No: 67-56-1
EC No: 200-659-6
Index No: 603-001-00-x
Registration No: 01-2119433307-44-xxxx
Classification: Flam. Liq. 2 (H225); Acute Tox 3 (H301 + H311 + H331); STOT SE 1 (H370)
Concentration: < 1%

Component: **Acetic Acid**
CAS No: 64-19-7
EC No: 200-580-7
Index No: -
Registration No: -
Classification: Skin Corr. 1A (H314) ; Flam. Liq 3 (H226)
Concentration: ≤ 2%

SECTION 4 FIRST AID MEASURES

4.1 Description of first aid measures
Consult a physician. Show this safety data sheet to the doctor in attendance.

If inhaled: If breathed in, move person into fresh air. If not breathing, give artificial respiration.

In case of skin contact: Wash off immediately with soap and plenty of water for at least 15 minutes while removing all contaminated clothes and shoes.

In case of eye contact: Rinse thoroughly with plenty of water for at least 15 minutes and consult a physician.

If swallowed: Do NOT induce vomiting. Never give anything by mouth to an unconscious person. Rinse mouth with water. Consult a physician.

4.2 Most important symptoms and effects, both acute and delayed
The most important known symptoms and effects are described in the labelling (Section 2.2) and/or in Section 11.

4.3 Indication of any immediate medical attention and special treatment needed
Notes to physician: treat symptomatically.

SECTION 5 FIRE FIGHTING MEASURES

5.1 Extinguishing media
Suitable extinguishing media: Use water spray, alcohol-resistant foam, dry chemical or carbon dioxide.

5.2 Special hazards arising from the substance or mixture
Carbon oxides.

5.3 Advice for firefighters
Wear self-contained breathing apparatus and full protective gear.

SECTION 6 ACCIDENTAL RELEASE MEASURES
6.1 Personal precautions, protective equipment and emergency procedures
Use personal protective equipment. Avoid breathing vapours, mist or gas. Ensure adequate ventilation. Remove all sources of ignition. Evacuate personnel to safe areas. Beware of vapours accumulating to form explosive concentrations. Vapours can accumulate in low areas. For personal protection see Section 8.

6.2 Environmental precautions
Prevent further leakage or spillage if safe to do so. Do not let product enter drains.

6.3 Methods and material for containment and cleaning up
Contain spillage and place in container for disposal according to local regulations (see Section 13). Keep in suitable, closed containers for disposal.

6.4 Reference to other sections
For disposal, see Section 13.

SECTION 7 HANDLING AND STORAGE
7.1 Precautions for safe handling
Avoid contact with skin and eyes. Avoid inhalation of vapour or mist. Keep away from sources of ignition—no smoking. Take measures to prevent the build-up of electrostatic charge. For precautions, see Section 2.2.

7.2 Conditions for safe storage, including any incompatibilities
Store in a cool place. Keep container tightly closed in a dry and well-ventilated place. Containers which are opened must be carefully resealed and kept upright to prevent leakage.

7.3 Specific end use(s)
No other specific uses are specified apart from those listed in Section 1.2.

SECTION 8 EXPOSURE CONTROLS/PERSONAL PROTECTION
8.1 Control parameters
This product, as supplied, does not contain any hazardous materials with occupational exposure limits established by the region specific regulatory bodies.

	Formaldehyde 50-00-0	Methanol 67-56-1	Acetic Acid 64-19-7
Austria	STEL: 0.5 ppm STEL: 0.6 mg/m³ TWA: 0.5 ppm TWA: 0.6 mg/m³	STEL: 800 ppm STEL: 1040 mg/m³ TWA: 200 ppm TWA: 260 mg/m³	STEL: 20 ppm STEL: 50 mg/m³ TWA: 10 ppm TWA: 25 mg/m³
Belgium	STEL: 0.3 ppm STEL: 0.38 mg/m³	STEL: 250 ppm STEL: 333 mg/m³ TWA: 200 ppm TWA: 266 mg/m³	STEL: 15 ppm STEL: 38 mg/m³ TWA: 10 ppm TWA: 25 mg/m³
Denmark	STEL: 0.3 ppm STEL: 0.4 mg/m³ TWA: 0.3 ppm TWA: 0.4 mg/m³	STEL: 400 ppm STEL: 520 mg/m³ TWA: 200 ppm TWA: 260 mg/m³	STEL: 20 ppm STEL: 50 mg/m³ TWA: 10 ppm TWA: 25 mg/m³
France	TWA: 0.5 ppm STEL: 1 ppm	STEL: 1000 ppm STEL: 1300 mg/m³ TWA: 200 ppm TWA: 260 mg/m³	STEL: 10 ppm STEL: 25 mg/m³
Germany	STEL: 0.6 ppm STEL: 0.74 mg/m³ TWA: 0.3 ppm TWA: 0.37 mg/m³	STEL: 800 ppm STEL: 1080 mg/m³ TWA: 200 ppm TWA: 270 mg/m³	STEL: 20 ppm STEL: 50 mg/m³ TWA: 10 ppm TWA: 25 mg/m³
Ireland	STEL: 2 ppm STEL: 2.5 mg/m³ TWA: 2 ppm TWA: 2.5 mg/m³	TWA: 200 ppm TWA: 260 mg/m³	STEL: 15 ppm STEL: 37 mg/m³ TWA: 10 ppm TWA: 25 mg/m³
Italy		TWA: 200 ppm TWA: 260 mg/m³	TWA: 10 ppm TWA: 25 mg/m³
Poland	STEL: 1 mg/m³ TWA: 0.5 mg/m³	STEL: 300 mg/m³ TWA: 100 mg/m³	STEL: 30 mg/m³ TWA: 15 mg/m³
Portugal	STEL: 0.3 ppm	STEL : 250 ppm TWA : 200 ppm TWA : 260 mg/m³	STEL: 15 ppm TWA: 10 ppm TWA: 25 mg/m³
Spain	STEL: 0.3 ppm STEL: 0.37 mg/m³	STEL: 250 ppm STEL: 333 mg/m³ TWA: 200 ppm TWA: 266 mg/m³	STEL: 15 ppm STEL: 37 mg/m³ TWA: 10 ppm TWA: 25 mg/m³
Sweden	STEL: 0.6 ppm STEL: 0.74 mg/m³ TWA: 0.3 ppm TWA: 0.37 mg/m³	STEL: 250 ppm STEL: 350 mg/m³ TWA: 200 ppm TWA: 250 mg/m³	STEL: 10 ppm STEL: 25 mg/m³ TWA: 5 ppm TWA: 13 mg/m³
The Netherlands	STEL: 0.5 mg/m³ TWA: 0.15 mg/m³	TWA: 133 mg/m³	
UK	STEL: 2 ppm STEL: 2.5 mg/m³ TWA: 2 ppm TWA: 2.5 mg/m³	STEL: 250 ppm STEL: 333 mg/m³ TWA: 200 ppm TWA: 266 mg/m³	

8.2 Exposure controls
8.2.1 Appropriate engineering controls
Handle in accordance with good industrial hygiene and safety practice. Wash hands before breaks and at the end of workday.

8.2.2 Personal protective equipment
(a) Eye/face protection: Tightly fitting safety goggles. Faceshield (8-inch minimum). Use equipment for eye protection tested and approved under appropriate government standards such as NIOSH (US) or EN 166(EU).

(b) Skin protection: Handle with gloves. Gloves must be inspected prior to use. Use proper glove removal technique (without touching glove's outer surface) to avoid skin contact with this product. Dispose of contaminated gloves after use in accordance with applicable laws and good laboratory practices. Wash and dry hands. The selected protective gloves should satisfy the specifications of EU Directive 89/686/EEC and the standard EN 374 derived from it.

(c) Body Protection: Complete suit protecting against chemicals. The type of protective equipment must be selected according to the concentration and amount of the dangerous substance at the specific workplace.

(d) Respiratory protection: Where risk assessment shows air-purifying respirators are appropriate use a full-face respirator with multi-purpose combination (US) or type ABEK (EN 14387) respirator cartridges as a backup to engineering controls. If the respirator is the sole means of protection, use a full-face supplied air respirator. Use respirators and components tested and approved under appropriate government standards such as NIOSH (US) or CEN (EU).

8.2.3 Environmental exposure controls
Prevent further leakage or spillage if safe to do so. Do not let product enter drains.

SECTION 9 PHYSICAL AND CHEMICAL PROPERTIES

9.1 Information on basic physical and chemical properties

a) Appearance aqueous solution Form: colourless liquid

b) Odour characteristic

c) Odour threshold no data available

d) pH no data available

e) Melting point / freezing point no data available

f) Initial boiling point and boiling range 102°C

g) Flash point >105°C

h) Evaporation rate no data available

i) Flammability (solid, gas) no data available

j) Upper/lower flammability or explosive limits no data available

k) Vapour pressure no data available

l) Vapour density >1

m) Relative density 1.071

n) Solubility (ies) Soluble in water

o) Partition coefficient: n-octanol/water no data available

p) Auto-ignition temperature no data available

q) Decomposition temperature no data available

r) Viscosity no data available

s) Explosive properties no data available

t) Oxidising properties no data available

9.2 Other information

No data available.

SECTION 10 STABILITY AND REACTIVITY

10.1 Reactivity

No data available.

10.2 Chemical stability

Stable under recommended storage conditions.

10.3 Possibility of hazardous reactions

No data available.

10.4 Conditions to avoid

Heat, flames and sparks.

10.5 Incompatible materials

No materials to be mentioned in particular.

10.6 Hazardous decomposition products

Carbon oxides.

SECTION 11 TOXICOLOGICAL INFORMATION

11.1 Information of toxicological effects

Acute toxicity: no data available
Skin corrosion/irritation: no data available
Serious eye damage/eye irritation: no data available
Respiratory or skin sensitisation: no data available
Germ cell mutagenicity: no data available
Carcinogenicity: IARC: 1 - Group 1: Carcinogenic to humans (Formaldehyde)
Reproductive toxicity: no data available
Specific target organ toxicity - single exposure: no data available
Specific target organ toxicity - repeated exposure: no data available
Aspiration hazard: no data available

Additional Information

Chemical Name	
Formaldehyde	LD50 oral 600 mg/kg (Rat) LD50 dermal 270 mg/kg (Rabbit) LC50 inhalation 0.578 mg/L (Rat) 4 h
Methanol	LD50 oral - rat - 5628 mg / kg LC50 inhalation - rat - 4h – 83.2 mg/l/4h
Acetic Acid	LD50 oral 3310 mg/kg (Rat) LD50 dermal 1060 mg/kg (Rabbit) LC50 inhalation 11.4 mg/L (Rat) 4 h

SECTION 12 ECOLOGICAL INFORMATION

12.1 Toxicity

Ecotoxicity effects: contains no substances known to be hazardous to the environment or not degradable in waste water treatment plants.

Toxicity to Fish	
Formaldehyde	0.032 - 0.226: 96 h Oncorhynchus mykiss mL/L LC50 flow-through 100- 136: 96 h Oncorhynchus mykiss mg/L LC50 static 1510: 96 h Lepomis macrochirus µg/L LC50 static 22.6 - 25.7: 96 h Pimephales promelas mg/L LC50 flow-through 23.2 - 29.7: 96 h Pimephales promelas mg/L LC50 static 41: 96 h Brachydanio rerio mg/L LC50 static
Methanol	LC50 - Pimephales promelas – 28200 mg / L 96h
Acetic Acid	75: 96 h Lepomis macrochirus mg/L LC50 static 79: 96 h Pimephales promelas mg/L LC50 static

Toxicity to Daphnia and other Aquatic Invertebrates	
Formaldehyde	11.3 - 18: 48 h Daphnia magna mg/L EC50 Static 2: 48 h Daphnia magna mg/L LC50
Methanol	EC50 - Daphnia magna - >10000 mg/l
Acetic Acid	47: 24 h Daphnia magna mg/L EC50 65: 48 h Daphnia magna mg/L EC50 Static

12.2 Persistence and degradability

No data available.

12.3 Bioaccumulative potential

No data available.

Chemical Name	log Pow
Formaldehyde	0.35
Methanol	-0.77
Acetic Acid	0

12.4 Mobility in soil

No data available.

12.5 Results of PBT and vPvB assessment

No data available.

12.6 Other adverse effects

No data available.

12.7 Additional information

None.

SECTION 13 DISPOSAL CONSIDERATIONS

13.1 Waste treatment methods

Product: Dispose of waste in accordance with all federal, state, and local regulations.

Contaminated packaging: Dispose of as unused product.

SECTION 14 TRANSPORT INFORMATION

IATA/DOT/IMDG/TDG: Not regulated.

14.1 UN number: -

14.2 UN proper shipping name: -

14.3 Transport hazard class(es): -

14.4 Packing group: -

14.5 Environmental hazards: -

14.6 Special precautions for user: -

SECTION 15 REGULATORY INFORMATION

15.1 Safety, health and environmental regulations/ legislation specific for the substance or mixture

No data available.

15.2 Chemical Safety Assessment

No chemical safety assessment has been carried out for this product.

SECTION 16 OTHER INFORMATION

Full text of H-Statements referred to in Sections 2 and 3

H225 Highly flammable liquid and vapour.

H226 Flammable liquid and vapour.

H301 + H311 + H331 Toxic if swallowed, in contact with skin or if inhaled.

H302 Harmful if swallowed.

H314 Causes severe skin burns and eye damage.

H317 May cause an allergic skin reaction.

H332 Harmful if inhaled.

H341 Suspected of causing genetic defects.

H350 May cause cancer.

H370 Causes damage to organs.

Acute Tox. Acute toxicity

Carc. Carcinogenicity

Flam. Liq. Flammable liquids

Muta. Germ Cell Mutagenicity

Skin Corr. Skin corrosion

Skin Sens. Skin sensitisation

STOT SE Specific target organ toxicity - single exposure

Amended sections are indicated by a line in the border.

The information supplied in this SDS is correct to the best of our knowledge. We do not accept any liability for loss, injury or damage, which may result from its use.

TRITON X SOLUTION SAFETY DATA SHEET

This Safety Datasheet complies with the requirements of Regulation (EC) No 1907/2006

SECTION 1 IDENTIFICATION OF THE SUBSTANCE/ MIXTURE AND THE COMPANY/UNDERTAKING

1.1 Product Identifier: 1472, 172018

TRITON X Solution

Used at concentration of ≤0.1% in: 145300, 145400, 145420, 145500, 145800, 145900, 146300, 146400, 146500, 146501, 108900, 108910, 108920, 108935, 148960, 148965, 148980, 148998, 149910, 149920, 149960, 151000, 153000, 249400

1.2 Relevant identified uses of the substance or mixture and uses advised against: for laboratory use (in vitro diagnostic).

1.3 Details of the supplier of the Safety Data Sheet:

Apacor Limited, Unit 5 Sapphire Centre, Fishponds Road, Wokingham, Berkshire, RG41 2QL, England
+44 (0) 118 979 5566
technical@apacor.com

1.4 Emergency telephone number:

+44 (0)118 979 5566
(Monday-Friday 0900-1700 excluding UK Public Holidays)

SECTION 2 HAZARDS IDENTIFICATION

2.1 Classification of the substance or mixture

Classification according to Regulation (EC) No 1272/2008 [CLP]:
Serious eye damage (Category1), H318

See Section 16 for the full text of H-Statements mentioned in this Section.

2.2 Label elements

Labelling according to Regulation (EC) No 1272/2008 [CLP]

Pictogram 

Signal word **Danger**

Hazard statement(s)

H318 Causes serious eye damage

Precautionary statements:

P280 Wear protective gloves/protective clothing/eye protection/face protection.
P305+P351+P338 IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.
P313 Get medical advice/attention.

See Section 16 for the full text of H-Statements mentioned in this Section.

2.3 Other hazards

None known.

SECTION 3 COMPOSITION/INFORMATION ON INGREDIENTS

3.2 Mixtures

Hazardous ingredients according to Regulation (EC) No 1272/2008

Component: **Triton X-100** (concentration 10–20%) (included in the Candidate List of Substances of Very High Concern (SVHC) according to Regulation (EC) No 1907/2006 (REACH))
CAS No: 9002-93-1
EC No: -
A registration number is not available for this substance as the substance or its use are exempted from registration according to Article 2 REACH Regulation (EC) No 1907/2006, the annual tonnage does not require a registration or the registration is envisaged for a later registration deadline.
Classification: Acute Tox. 4 (H302); Serious Eye Dam. 1 (H318)
Concentration: 5–10%

See Section 16 for the full text of H-Statements mentioned in this Section.

SECTION 4 FIRST AID MEASURES

4.1 Description of first aid measures

Consult a physician. Show this safety data sheet to the doctor in attendance.

If inhaled: If breathed in, move person into fresh air. If not breathing, give artificial respiration. Consult a physician.

In case of skin contact: Take off immediately all contaminated clothing. Rinse skin with water/shower.

In case of eye contact: rinse out with plenty of water. Immediately consult an ophthalmologist.

If swallowed: immediately make victim drink water (2 glasses at most). Consult a physician.

4.2 Most important symptoms and effects, both acute and delayed

Irritation and corrosion. Risk of serious damage to eyes.

4.3 Indication of any immediate medical attention and special treatment needed

No data available.

SECTION 5 FIRE FIGHTING MEASURES

5.1 Extinguishing media

Suitable extinguishing media: Use water spray, foam, dry chemical or carbon dioxide. (Use extinguishing measures that are appropriate to local circumstances and the surrounding environment.)
Unsuitable extinguishing media: For this substance/mixture no limitations of extinguishing agents are given.

5.2 Special hazards arising from the substance or mixture

Not combustible. Ambient fire may liberate hazardous vapours.

5.3 Advice for firefighters

Special protective equipment for firefighters: In the event of fire, wear self-contained breathing apparatus.
Further information: Prevent fire extinguishing water from contaminating surface water or the ground water system.

SECTION 6 ACCIDENTAL RELEASE MEASURES

6.1 Personal precautions, protective equipment and emergency procedures

Advice for non-emergency personnel: Do not breathe vapours, aerosols. Avoid substance contact. Ensure adequate ventilation. Evacuate the danger area, observe emergency procedures, consult an expert.

Advice for emergency responders: Protective equipment see Section 8.

6.2 Environmental precautions

Prevent further leakage or spillage if safe to do so. Do not let product enter drains. Discharge into the environment must be avoided.

6.3 Methods and material for containment and cleaning up

Soak up with inert absorbent material and dispose of as hazardous waste. Keep in suitable, closed containers for disposal.

6.4 Reference to other sections

For disposal, see Section 13.

SECTION 7 HANDLING AND STORAGE

7.1 Precautions for safe handling

Avoid inhalation of vapour or mist. For precautions see Section 2.2.

7.2 Conditions for safe storage, including any incompatibilities

Tightly closed. Recommended storage temperature see product label.

7.3 Specific end use(s)

No other specific uses are specified apart from those listed in Section 1.2.

SECTION 8 EXPOSURE CONTROLS/PERSONAL PROTECTION

8.1 Control parameters

Contains no substances with occupational exposure limit values.

8.2 Exposure controls

8.2.1 Appropriate engineering controls

Handle in accordance with good industrial hygiene and safety practice. Wash hands before breaks and at the end of workday.

8.2.2 Personal protective equipment

(a) Eye/face protection: Tightly fitting safety goggles. Faceshield (8-inch minimum). Use equipment for eye protection tested and approved under appropriate government standards such as NIOSH (US) or EN 166(EU).

(b) Skin protection: Handle with gloves. Gloves must be inspected prior to use. Use proper glove removal technique (without touching glove's outer surface) to avoid skin contact with this product. Dispose of contaminated gloves after use in accordance with applicable laws and good laboratory practices. Wash and dry hands. The selected protective gloves should satisfy the specifications of EU Directive 89/686/EEC and the standard EN 374 derived from it.

(c) Body Protection: Complete suit protecting against chemicals. The type of protective equipment must be selected according to the concentration and amount of the dangerous substance at the specific workplace.

(d) Respiratory protection: Where risk assessment shows air-purifying respirators are appropriate use a full-face respirator with multi-purpose combination (US) or type ABEK (EN 14387) respirator cartridges as a backup to engineering controls. If the respirator is the sole means of protection, use a full-face supplied air respirator. Use respirators and components tested and approved under appropriate government standards such as NIOSH (US) or CEN (EU).

8.2.3 Environmental exposure controls

Prevent further leakage or spillage if safe to do so. Do not let product enter drains. Discharge into the environment must be avoided.

SECTION 9 PHYSICAL AND CHEMICAL PROPERTIES

9.1 Information on basic physical and chemical properties

a) Appearance Form: clear, liquid; Colour: light yellow

b) Odour no data available

c) Odour threshold no data available

d) pH 9.7

e) Melting point / freezing point approx. 6°C

f) Initial boiling point and boiling range 200°C

g) Flash point 251°C

h) Evaporation rate no data available

i) Flammability (solid, gas) no data available

j) Upper/lower flammability or explosive limits no data available

k) Vapour pressure <1 hPa at 25°C

l) Vapour density no data available

m) Relative density 1.070 g/cm^3

n) Solubility (ies) Soluble in water

o) Partition coefficient: n-octanol/water no data available

p) Auto-ignition temperature no data available

q) Decomposition temperature no data available

r) Viscosity no data available

s) Explosive properties no data available

t) Oxidising properties no data available

9.2 Other information no data available

SECTION 10 STABILITY AND REACTIVITY

10.1 Reactivity

No data available.

10.2 Chemical stability

Stable under recommended storage conditions.

10.3 Possibility of hazardous reactions

No data available.

10.4 Conditions to avoid

No data available.

10.5 Incompatible materials

Strong acids. Strong bases. Strong oxidizing agents.

10.6 Hazardous decomposition products

Other decomposition products—no data available. In the event of fire: see Section 5.

SECTION 11 TOXICOLOGICAL INFORMATION

11.1 Information of toxicological effects

Acute toxicity: no data available

Skin corrosion/irritation: no data available

Serious eye damage/eye irritation: no data available

Respiratory or skin sensitisation: no data available

Germ cell mutagenicity: no data available

Carcinogenicity: IARC: no component of this product present at levels greater than 0.1% is identified as probable, possible or confirmed human carcinogen by IARC.

Reproductive toxicity: no data available

Specific target organ toxicity - single exposure: no data available

Specific target organ toxicity - repeated exposure: no data available

Aspiration hazard: no data available

Additional information: RTECS: not available. To the best of our knowledge, the chemical, physical, and toxicological properties have not been thoroughly investigated.

11.2 Further information

Triton X-100
Acute oral toxicity: LD50 Rat: 1,800 mg/kg (RTECS)
Germ cell mutagenicity: Genotoxicity in vitro Mutagenicity (mammal cell test): Mouse lymphoma test Result: negative

SECTION 12 ECOLOGICAL INFORMATION

12.1 Toxicity

No data available.

12.2 Persistence and degradability

No data available.

12.3 Bioaccumulative potential

No data available.

12.4 Mobility in soil

No data available.

12.5 Results of PBT and vPvB assessment

PBT/vPvB assessment not available as chemical safety assessment not required/not conducted.

12.6 Other adverse effects

Discharge into the environment must be avoided.
Components: Triton X-100
Toxicity to fish
LC50 Lepomis macrochirus: 2,800 - 3,200 µg/l; 96 h
Toxicity to daphnia and other aquatic invertebrates
LC50 Daphnia magna: 11.2 mg/l; 48 h

12.7 Additional information

No data available.

SECTION 13 DISPOSAL CONSIDERATIONS

13.1 Waste treatment methods

Product: Offer surplus and non-recyclable solutions to a licensed disposal company.

Contaminated packaging: Dispose of as unused product.

SECTION 14 TRANSPORT INFORMATION

IATA/DOT/IMDG/TDG: Not regulated.

14.1 UN number: -

14.2 UN proper shipping name: -

14.3 Transport hazard class(es): -

14.4 Packing group: -

14.5 Environmental hazards: -

14.6 Special precautions for user: -

SECTION 15 REGULATORY INFORMATION

15.1 Safety, health and environmental regulations/ legislation specific for the substance or mixture

Substances of very high concern (SVHC): This product does contain substances of very high concern above the respective regulatory limit (>0.1% w/w), Regulation (EC) No 1907/2006 (REACH), Article 57).
Contains: Triton X-100.

15.2 Chemical Safety Assessment

No chemical safety assessment has been carried out for this product.

SECTION 16 OTHER INFORMATION

Full text of H-Statements referred to in Sections 2 and 3

H302 Harmful if swallowed

H318 Causes serious eye damage

Acute Tox. Acute Toxicity

Serious Eye Dam. Serious Eye Damage

Amended sections are indicated by a line in the border.

The information supplied in this SDS is correct to the best of our knowledge. We do not accept any liability for loss, injury or damage, which may result from its use.

Midi Parasep® SF Faecal Parasite Concentrator

Code	Product
149900	Midi Parasep® SF
149910	Midi Parasep® SF Formalin & Triton X
149920	Midi Parasep® SF SAF & Triton X
149931	Midi Parasep® SF SafEFix™ Ecological Fixative
249300	Midi Parasep® SF AlcorFix™

Discard in accordance with your standard and local operating procedures for clinical waste.

Products can be ordered direct from Apacor or from an appointed distributor
Visit our website for all the latest information **www.apacor.com** or e-mail: **sales@apacor.com**

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