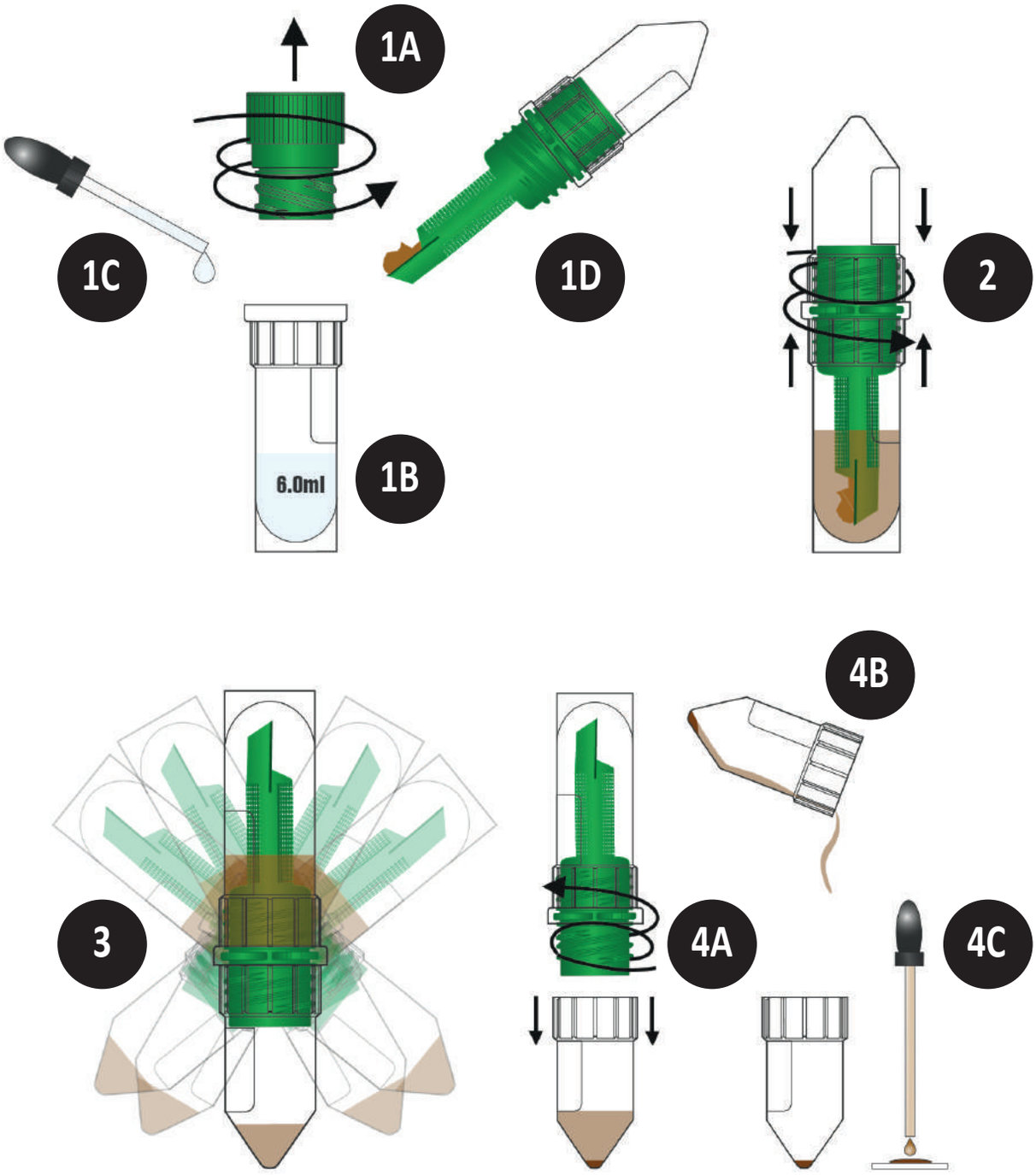
APACOR

Midi Parasep® Faecal Parasite Concentrator



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

APA101
V8.6
06/2022

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®. To avoid cross contamination the Midi Parasep® device should remain closed at all times except when introducing the sample or when retrieving the final concentrated sample for examination.

Sample Preparation
For empty Midi Parasep®.

1A Unscrew lid.

1B Add 6.0ml of fixative.

1C Add one drop of surfactant (eg:Triton X-100) to the chamber.

Alternatively, use reagent ready Midi Parasep®

1D Introduce a level scoop of formed faecal sample (equivalent 1g) to the fixative using the spoon at the end of the Midi Parasep® filter. For liquid samples, add 2 level scoops (equivalent 2g) to the fixative. Add 2.0ml of Ethyl Acetate to the mixing chamber. Mix in thoroughly with the Midi Parasep® spoon. If the sample is hard, break it up with the end of the spoon.

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

Centrifugation
3 Invert Midi Parasep® and centrifuge at 1200g for 3 minutes. Midi Parasep® 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
Open very slowly to avoid aerosol release.

4A Unscrew and discard the filter and mixing tube.

4B Pour off all the liquid above the sediment.

4C Pipette one drop of saline or Lugol's Iodine solution onto a slide, add one drop of deposit to the saline or Lugol's Iodine, mix sample and cover with cover-slip.

CS

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

Příprava vzorku
Pro prázdný Midi Parasep®.

1A Odšroubujte víčko.

1B Přidejte 6,0 ml fixačního.

1C Jednu kapku surfaktantu (např. Triton X-100) do míchacího prostoru.

Případně použijte Midi Parasep připravený k reagencie.

1D Přidejte zarovnanou odměrku vytvořeného vzorku stolice (ekvivalent 1 g) do fixačního prostředku pomocí lžíce na konci filtru Midi Parasep®. U tekutých vzorků přidejte k fixativu 2 zarovnané odměrky (ekvivalent 1 g). Přidejte 2,0 ml ethyl acetátu do míchací komory. Důkladně promíchejte lžičkou Midi Parasep®. Pokud je vzorek tvrdý, rozlomte ho koncem lžíce.

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

Centrifugace
3 Vložte do centrifugy a centrifugujte při 1200 g po dobu 3 minut. Pro Midi Parasep® 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
Otevřete jej velmi pomalu, aby se zabránilo uvolňování aerosolu.

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

4B Vylijte veškerou kapalinu nad usazeninou.

4C Napipetujte jednu kapku fyziologického roztoku nebo Lugolův roztok jodu na snímku, přidejte jednu kapku vkladu do fyziologického roztoku nebo Lugolův jódu, smíchejte vzorek a zakryjte krycí sklíčko.

DE

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

Probenvorbereitung
Für leere Midi Parasep®.

1A Deckel abschrauben.

1B 6,0ml Fixierlösung zugeben.

1C Triton X-100 (1:20 Verdünnung) zugeben.

Gebruik als alternatief reagensklare Midi Parasep®

1D Geben Sie mit dem Löffel am Ende des Midi Parasep®-Filters einen gestrichenen Messlöffel der gebildeten Kotprobe (entspricht 1 g) in das Fixiermittel. Geben Sie bei flüssigen Proben 2 gestrichene Messlöffel (entsprechend 1 g) zum Fixiermittel hinzu. Geben Sie 2,0 ml Ethylacetat in die Mischkammer. Mit dem Midi Parasep® Löffel gründlich einmischen. Wenn die Probe hart ist, brechen Sie sie mit dem Ende des Löffels auf.

Emulgieren
2 Den Filterteil mit dem Sedimentationsröhrchen des Midi Parasep® 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® mit dem konusförmigen, spitz zulaufenden Teil nach unten in die Zentrifuge stellen. 3 Minuten bei 1200xg zentrifugieren. Midi Parasep® 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
Midi Parasep® sehr langsam öffnen, um eine Aerosol-Freisetzung zu vermeiden

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

4B Den Überstand vorsichtig abgießen.

4C 1 Tropfen NaCl bzw. Lugol's Jod-Lösung auf einen Objektträger träufeln. 1 Tropfen des Sediments dazugeben, mischen und mit einem Deckglas zudecken.

ES

Mirar la etiqueta para ver condiciones de almacenaje y fecha caducidad. Cuando se manipule Midi Parasep® se ruega seguir las instrucciones. Para evitar contaminaciones cruzadas el Midi Parasep® 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
Para Midi Parasep® vacío.

1A Desenroscar el tapón.

1B Añadir 6.0 ml de fijador.

1C Si se requiere una gota de surfactante (Triton X-100) para emulsionar.

Como alternativa, utilice Midi Parasep® preparado para reactivos.

1D Introduzca una cucharada al ras de la muestra fecal formada (equivalente a 1 g) en el fijador utilizando la cuchara al final del filtro Midi Parasep®. Para muestras líquidas, agregue 2 cucharadas rasas (equivalente a 1 g) al fijador. Agregue 2,0 ml de acetato de etilo a la cámara de mezcla. Mezclar bien con la cuchara Midi Parasep®. Si la muestra es dura, rómpala con el extremo 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® y centrifugar a 1200g durante 3 minutos. El Midi Parasep® 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
Abrir lentamente para evitar la salida de aerosoles

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

4B Decante el liquido sobrenadante del sedimento.

4C Dispensar una gota de solución salina o solución de yodo-lugol, mezclar con la muestra y cubrir con un cubre-objetos.

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® 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
Pour Midi Parasep® vide.

1A Dévissez le bouchon.

1B Ajoutez 6,0ml de fixateur.

1C Et ajoutez une goutte de surfactant (par ex: Triton X-100) pour émulsifier.

Alternativement, utilisez le Midi Parasep® prêt à réagir.

1D Introduire une cuillère rase d'échantillon fécal formé (équivalent 1g) dans le fixateur à l'aide de la cuillère à l'extrémité du filtre Midi Parasep®. Pour les échantillons liquides, ajouter 2 cuillères rases (équivalent 1g) au fixateur. Ajouter 2,0 ml d'acétate d'éthyle dans la chambre de mélange. Bien mélanger avec la cuillère Midi Parasep®. Si l'échantillon est dur, cassez-le avec le bout de la cuillère.

Émulsification
2 Scellez le Midi Parasep® 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® et centrifugez le à 1200g pendant 3 minutes. Midi Parasep® 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
Débloquer délicatement pour dégazer (éther!!) puis ouvrir le tube.

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

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

4C Déposez une goutte de sérum physiologique ou de solution d'iode de Ludol sur une lame, ajoutez une goutte de sédiment à la solution saline ou iodisé, mélangez l'échantillon et couvrez avec couvre-lamelle.

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® bi trebao biti zatvoren cijelo vrijeme osim kada stavljate uzorak ili prilikom uzimanja krajnjeg koncentriranog uzorka za mikroskopiranje.

Pripema Uzorka
Za prazan Midi Parasep®.

1A Otvorite poklopac.

1B Dodajte 6,0 ml fiksator.

1C Dodajte Triton X u komoru za miješanje.

Alternativno, koristite Midi Parasep® spreman za reagens.

1D Unesite ravnu mjericu formiranog uzorka izmeta (ekvivalentno 1 g) u fiksativ koristeći žlicu na kraju Midi Parasep® filtera. Za tekuće uzorke dodajte 2 mjerene mjerice (ekvivalentno 1 g) fiksatoru. Dodajte 2,0 ml etil acetata u komoru za miješanje. Temeljito promiješajte Midi Parasep® žlicom. Ako je uzorak tvrd, razbijte ga krajem žlice.

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

Centrifugiranje
3 Okrenite Midi Parasep® i centrifugirajte na 1200 g 3 minute. Midi Parasep® 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
Otvorite vrlo polagano da biste izbjegli oslobadanje aerosola.

4A Otvorite i bacite filter i komoru za miješanje.

4B Odlijte svu tekućinu iznad sedimenta.

4C Otpipetirajte jednu kap fiziološke otopine ili lugolove otopine na stakalce, dodajte jednu kap depozita u fiziološku ili lugolovu otopinu na stakalcu, promiješajte uzorak i pokrijte pokrovnim stakalcem.

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®. Per evitare cross-contaminazioni il concentratore Midi Parasep® 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
Per Midi Parasep® vuoto.

1A Svitare il tappo.

1B Aggiungere 6,0 ml di fissativo.

1C Se richiesto, aggiungere una goccia di surfattante (es.Triton X-100) per emulsionare.

In alternativa, utilizzare Midi Parasep® pronto per il reagente

1D Introdurre un misurino raso di campione fecale formato (equivalente a 1 g) nel fissativo utilizzando il cucchiaio all'estremità del filtro Midi Parasep®. Per i campioni liquidi, aggiungere 2 misurini rasi (equivalente a 1 g) al fissativo. Aggiungere 2,0 ml di acetato di etile nella camera di miscelazione. Mescolare accuratamente con il cucchiaio Midi Parasep®. Se il campione è duro, romperlo con l'estremità del cucchiaio.

Omogenizzazione

2 Chiudere ermeticamente il Midi Parasep® 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® e centrifugare a 1200g per 3 minuti. Il Midi Parasep® 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
Aprire molto lentamente per evitare aerosol

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

4B Eliminare il sovranatante.

4C Pipettare una goccia di soluzione salina o iodina di Lugol sul vetrino, aggiungere una goccia di deposito sulla soluzione, mescolare il campione e coprirlo.

NL

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

Staalvoorbereiding
Voor lege Midi Parasep®.

1A Schroeft u het deksel los.

1B Voeg 6,0 ml fixatief toe.

1C Voegt een druppel surfactant (bv:Triton X-100) om te emulgeren.

Gebruik als alternatief reagensklare Midi Parasep®.

1D Breng een afgestreken schep gevormd fecaal monster

(equivalent van 1 g) aan op het fixeermiddel met behulp van de lepel aan het uiteinde van het Midi Parasep®-filter. Voeg voor vloeibare monsters 2 afgestreken maatscheppen (equivalent van 1 g) toe aan het fixeermiddel. Voeg 2,0 ml ethylacetaat toe aan de mengkamer. Goed mengen met de Midi Parasep® lepel. Als het monster hard is, verbreek het dan met het uiteinde van de lepel.

Emulsificatie

2 Sluit de Midi Parasep® 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® om en centrifugeer aan 1200g gedurende drie minuten. Midi Parasep® 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
Open de buis langzaam om aerosol vorming te voorkomen.

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

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

4C Pipeteer één druppel saline of Lugol's Iodine oplossing op een objectglaasje, voeg één druppel sediment toe aan de saline of Lugol's Iodine, meng het monster en dek af een dekglaasje.

PL

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

Przygotowanie próbki
Do pustego Midi Parasep®.

1A Otworzyć probówkę a następnie.

1B Dodaj 6,0ml utrwalacza.

1C Oraz 1 kroplę surfaktantu (np.Triton X-100).
Alternatywnie można użyć gotowego odczynnika Midi Parasep®.

1D Za pomocą łyżki znajdującej się na końcu filtra Midi Parasep® wprowadzić do utrwalacza płaską miarkę uformowanej próbki kału (odpowiednik 1 g). W przypadku próbek płynnych dodaj 2 płaskie miarki (odpowiednik 1 g) do utrwalacza. Dodaj 2,0 ml octanu etylu do komory mieszania. Dokładnie wymieszać łyżką Midi Parasep®. Jeśli próbka jest twarda, rozbij ją końcem łyżki.

Przygotowanie emulsji
2 Połączyć ze sobą dwie części probówki Midi Parasep® (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® wirować przy 1200 g przez 3 minut. Midi Parasep® 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
Po odwirowaniu bardzo ostrożnie otworzyć probówkę.

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 Umieść jedną kroplę soli fizjologicznej lub płynu Lugola na szkiełku podstawowym. Dodaj jedną kroplę zagęszczonego materiału do soli fizjologicznej lub płynu Lugola na szkiełku, wymieszaj próbkę i przykryj szkiełkiem nakrywkowym.

PT

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

Preparação da amostra
Para Midi Parasep® vazio.

1A Desenroscar a tampar.

1B Adicione 6,0ml de fixador.

1C Se necessário uma gota de surfactante (Triton X-100) para emulsionar.

Alternativamente, use o Midi Parasep® pronto para reagente

1D Introduza uma colher rasa de amostra fecal formada (equivalente a 1g) no fixador usando a colher na extremidade do filtro Midi Parasep®. Para amostras líquidas, adicione 2 colheres rasas (equivalente a 1g) ao fixador. Adicione 2,0 ml de acetato de etila à câmara de mistura. Misture bem com a colher Midi Parasep®. Se a amostra estiver dura, quebre-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® e centrifugar a 1200g por 3 minutos. Midi Parasep® 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
Abrir lentamente para evitar a saida de aerossóis

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

4B Decante o liquido sobrenadante do sedimento.

4C Pipete uma gota de solução salina ou de solução Iodada de Lugol para uma lâmina, adicione uma gota do deposito na solução salina ou solução Iodada de Lugol, misture a amostra e cubra com uma lamela.

SI

Shranjevanje in rok uporabe: glej nalepko!
Prosimo, da pri uporabi koncentratorja Midi Parasep® upoštevate naslednja priporočila. Koncentrator Midi Parasep® 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
Za prazan Midi Parasep®.

1A Odvijte zamašek koncentratorja.

1B Dodajte 6.0 ml fiksativa.

1C Ter 1 kapljico surfaktanta (npr. Triton-X-100).

Alternativno, koristite Midi Parasep® spreman za reagens

1D Uneti ravnu mericu formiranog uzorka fekalija (ekvivalentno 1 g) u fiksativ koristeći kašiku na kraju Midi Parasep® filtera. Za tečne uzorke, dodajte 2 merice (ekvivalentno 1 g) u fiksativ. Dodajte 2,0 ml etil acetata u komoru za mešanje. Dobro promešajte sa Midi Parasep® kašikom. Ako je uzorak tvrd, razbijte ga krajem kašike.

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 3 minuti pri 1200g. 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
Koncentrator odprite zelo počasi, da preprečite uhajanje aerosol.

4A Filtrirni del koncentratorja odvijte in zavrzite.

4B Vso tekočino nad sedimentom odlijte.

4C Na objektno stekelce kapnite eno kapljico fiziološke ali lugol jodove raztopine, dodajte eno kapljico sedimenta, vzorec premešajte in pokrijte s krovnim stekelcem.

APAFIX™ 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: Apafix™

108801, 108887 145002, 145003, 145005 146003, 146004, 146005, 149901, 160001, 902500, 907500

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, United Kingdom
+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]:
The mixture does not present a physical or chemical hazard.
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]

Hazard statement(s)

NC Not Classified

Precautionary statements:

P261 – Avoid breathing vapour/spray
P280 - Wear protective gloves/protective clothing/eye protection/face protection.
Contains Acetic Acid

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: **Acetic Acid**
CAS No: 64-19-7
EC No: 200-580-7
Index No: 607-002-00-6
Registration No: -
Classification: Skin Corr. 1A (H314), Flam. Liq. 3 (H226), Eye Dam. 1 (H318)
Concentration: < 10%

Component: **Menthol**
CAS No: 89-78-1
EC No: 201-939-0
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: **Thymol**
CAS No: 89-83-8
EC No: 201-944-8
Index No: 604-032-00-1
Registration No: -
Classification: Acute Tox. 4 (H302), Skin Corr. 1B (H314), Aquatic Chronic 2 (H411)
Concentration: < 1%

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. Consult a physician if discomfort continues.

If swallowed: Clean mouth with water and drink afterwards plenty of water. Consult a physician.

If inhaled: Move to fresh air. Consult a physician if discomfort continues.

4.2 Most important symptoms and effects, both acute and delayed

The most important 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

Notes to physician: Treat symptomatically. Have facilities in place to wash skin and eyes in case of exposure. Severe cases of exposure should receive prompt medical attention.

Eye contact – May cause eye irritation. May cause redness.

Skin Contact – May irritate the skin.

Ingestion – May irritate the mouth and throat. Small amounts will leave taste in mouth, larger amounts may cause nausea and vomiting.

Inhalation – Acute: May irritate the respiratory system and cause coughing. Delayed: Prolonged exposure to vapours or mists can cause damage to the mucous membranes of the respiratory system.

SECTION 5 FIRE FIGHTING MEASURES

5.1 Extinguishing media

This product is non-combustible. Water spray, dry powder, carbon dioxide or alcohol resistant foam.

5.2 Special hazards arising from the substance or mixture

In case of fire, toxic or irritating fumes or vapours may be formed. Contact with metals may form hydrogen gas which is flammable and can result in explosion.

5.3 Advice for firefighters

Wear self-contained breathing apparatus for firefighting.

SECTION 6: ACCIDENTAL RELEASE MEASURES

6.1 Personal precautions, protective equipment and emergency procedures

Use personal protective equipment, gloves and protective eye glasses. 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 (e.g. 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.

7.2 Conditions for safe storage, including any incompatibilities

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

7.3 Specific end use(s)

The identified uses for this product are detailed 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.

	Acetic Acid 64-19-7
Austria	STEL: 20ppm STEL: 50mg/m^3 TWA: 10ppm TWA: 25 mg/m^3
Belgium	STEL: 15ppm STEL: 38 mg/m^3 TWA: 10ppm TWA: 25mg/m^3
Denmark	STEL: 20ppm STEL: 50 mg/m^3 TWA: 10ppm TWA: 25 mg/m^3
France	STEL: 10ppm STEL: 25 mg/m^3
Germany	STEL: 20ppm STEL: 50 mg/m^3 TWA: 10ppm TWA: 25 mg/m^3
Ireland	STEL: 15ppm STEL: 37 mg/m^3 TWA: 10ppm TWA: 25 mg/m^3
Italy	TWA: 10ppm TWA: 25 mg/m^3
Poland	STEL: 30 mg/m^3 TWA: 15 mg/m^3
Portugal	STEL: 10ppm TWA: 10ppm TWA: 25 mg/m^3
Spain	STEL: 15ppm STEL: 37 mg/m^3 TWA: 10ppm TWA: 25 mg/m^3
Sweden	STEL: 10ppm STEL: 25 mg/m^3 TWA: 5ppm TWA: 13 mg/m^3

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 the workday. Ensure adequate ventilation, especially in confined areas.

8.2.2 Personal protective equipment

(a) Eye/face protection: Tightly fitting safety goggles. Face shield (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

See section 6.2

SECTION 9: PHYSICAL AND CHEMICAL PROPERTIES

9.1 Information on basic physical and chemical properties

a) Appearance: Form: liquid
Colour: Clear

b) Odour: pungent/acetic acid + distinct thyme smell

c) Odour threshold: no data available

d) pH: ≤1

e) Melting point / freezing point: no data available

f) Initial boiling point / boiling range: no data available

g) Flash point: Not applicable. The mixture is non-flammable.

h) Evaporation rate: no data available

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

j) Upper/lower flammability or explosive limits: Not applicable. The mixture is non-flammable.

k) Vapour pressure: no data available

l) Vapour density: no data available

m) Relative density: no data available

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

Reactions characteristic of weak acids.

10.2 Chemical stability

Stable under recommended handling and storage conditions under Section 7.

10.3 Possibility of hazardous reactions

May react vigorously or exothermically. Pressure may build up if reaction occurs in a sealed container. Will not polymerise.

10.4 Conditions to avoid

Avoid heat, direct sunlight and moisture. Avoid storage in freezing conditions. Avoid storage with incompatible materials. Avoid storage in an unstable manner or in a situation that would result in exposure to the product. It is advisable to store the product within some form of containment to prevent spillages reaching drainage systems.

10.5 Incompatible materials

Alkalis. Oxidising agents. Metals.

10.6 Hazardous decomposition products

Under normal use – none

SECTION 11: TOXICOLOGICAL INFORMATION

11.1 Information of toxicological effects

Acute toxicity:

Product: The mixture has not been tested for toxicological properties. This information refers to acetic acid as a pure substance.

Inhalation: Industry – Dermal; Long term systemic effects 22mg/kg/day Haematological effects

Eye contact: no data available

Skin contact: no data available

Ingestion: Sodium salt of acetic acid, pH 6-7

Skin corrosion/irritation: Dose: 0.5ml, 4 hr, Rabbit Primary dermal irritation index: 1.1 OECD Guideline 404 10% solution. Slightly irritating.

Serious eye damage/eye irritation: OECD 405, rabbit, 10% solution, 4 hour, 0.1ml. Erythema = 2.67, corneal swelling = 87%.

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

The mixture has not been tested for ecotoxicological properties.

The following information contained in section 12 refers to acetic acid as a pure substance.

Acute toxicity – fish: LC50, 96 hours: > 1000 mg/l, Onchorhynchus mykiss (Rainbow trout)
OECD 203 (Fish, Acute Toxicity Test)
Freshwater, semi-static.
Mortality

Acute toxicity – aquatic invertebrates: EC_{50}, 48 hours: > 300.82 mg/l, Daphnia magna
OECD Guideline 202.
Static, freshwater.
Mobility.
Test substance potassium acetate; result based on the acetate ion.

Acute toxicity – aquatic plants: EC_{50}, 72 hours: > 300.82 mg/l, Static, saltwater, Skeletonema costatum.
Test substance potassium acetate; result based on the acetate ion.

Acute toxicity – microorganisms: EC_{50}: 850 mg/l, Industry - Dermal; Long term systemic effects 22 mg/kg/day
Pseudomonas putida, static, freshwater, 16 hour

Acute toxicity – terrestrial: Not available.

Chronic toxicity – fish early life stage: Not available.

Short term toxicity – embryo and sac fry stages: Industry - Dermal; Long term systemic effects 22 mg/kg/day
No supplied or registered information.
Chronic toxicity – aquatic invertebrates: Not available.
Toxicity to soil: Not available.
Toxicity to terrestrial plants: Not available.
12.2 Persistence and degradability
Phototransformation: Water - DT_{50}: 26.7 days
Degradation by hydroxyl radicals.
Calculated value.
Stability (hydrolysis): Scientifically unjustified.
Biodegradation: Water – Degradation (%) 96%: 20 days
Readily biodegradable – Half life: 2 days
Biological oxygen demand: No information available.
Chemical oxygen demand: No information available.
12.3 Bioaccumulative potential
BCF: 3.16, QSAR calculation. Fish, freshwater. Not bioaccumulating.
12.4 Mobility in soil
No information available.
12.5 Results of PBT and vPvB assessment
PBT or vPvB not classified according to current EC criteria.
12.6 Other adverse effects
No information available.
12.7 Additional information
No information available.

SECTION 13: DISPOSAL CONSIDERATIONS
Proper waste management of the mixture and its container must be determined in accordance with directive 2008/98/EC.
13.1 Waste treatment methods
13.1.1 Product/packaging disposal
Do not pour into drains or waterways. Recycle of dispose of waste in compliance with current, local legislation, preferably via a certified waste company.

SECTION 14: TRANSPORT INFORMATION
IATA/DOT/IMDG/TDG - Not regulated
14.1 UN number: Not applicable
14.2 UN proper shipping name: Not applicable
14.3 Transport hazard class(es): Not applicable
14.4 Packing group: Not applicable
14.5 Environmental hazards: Not applicable
14.6 Special precautions for user: Not applicable

SECTION 15: REGULATORY INFORMATION
15.1 Safety, health and environmental regulations/ legislation specific for the substance or mixture
National regulations: Industry – Dermal; Long term systemic effects 22mg/kg/day
EU legislation: Regulation (EC) No 1907/2006 of the European Parliament and of the Council of 18 December 2006 concerning the Registration, Evaluation, Authorisation and Restriction of Chemicals (REACH) (as amended).
Regulation (EC) No 1272/2008 of the European Parliament and of the Council of 16 December 2008 on classification, labelling and packaging of substances and mixtures (as amended).
Regulation (EU) 453/2010.
15.2 Chemical Safety Assessment
A chemical safety assessment was not carried out for this mixture.

SECTION 16: OTHER INFORMATION
The information in this datasheet is based on our current level of knowledge and on national and international regulations. The mixture must not be used for other purposes than those specified in Section 1. It is at all times the responsibility of the use to take all necessary measures to comply with legal requirements and local regulations. The information in this safety data sheet must be regarded as a description of the safety requirements relating to the mixture and not as a guarantee of the properties thereof.

Full text of H-Statements referred to under sections 2 and 3
H226 – Flammable liquid and vapour
H314 - Causes severe skin burns and eye damage
H318 - Causes serious eye damage
P261 – Avoid breathing vapour/spray
P280 - Wear protective gloves/protective clothing/eye protection/face protection.
The above information is believed to be correct but does not purport to be all inclusive and shall be used as a guide. Apacor shall not be held liable for any damages resulting from handling or from contact with the above product, since the user's working conditions are not known by Apacor.

EC REP
MDSS GmbH
Schiffaraben 41
30175 Hanover
Germany

ETHYL ACETATE 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: Ethyl Acetate

145003, 145400, 145420, 145501, 145900, 146004, 146400, 146501, 1473, 900000, 901000, 903000, 905000, 906000, 908000

Synonyms, Trade Names: ETHYL ACETATE 98 - 100%, ACETIC ACID ETHYL ESTER, ACETOXYETHANE

REACH Registration Number: 01-2119475103-46-XXXX

CAS-No: 141-78-6

EU Index No: 607-022-00-5

EC No: 205-500-4

1.2 Relevant identified uses of the substance or mixture and uses advised against: laboratory chemical for the removal of fat from faecal samples.

1.3 Details of the supplier of the Safety Data Sheet: Apacor Limited, Unit 5 Sapphire Centre, Fishponds Road, Wokingham, Berkshire, RG41 2QL, United Kingdom

+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]:

Flam. Liq. 2 - H225

STOT SE 3 - H336

Eye Irrit. 2 – H319

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 Statements

H225 Highly flammable liquid and vapour.

H319 Causes serious eye irritation.

H336 May cause drowsiness or dizziness.

Precautionary Statements

P210 Keep away from heat/sparks/open flames/hot surfaces. No smoking.

P261 Avoid breathing vapour/spray.

P303+361+353 IF ON SKIN (or hair): Take off immediately all contaminated clothing. Rinse skin with water/shower.

P304+340 IF INHALED: Remove person to fresh air and keep comfortable for breathing.

P305+351+338 IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing.

P501 Dispose of contents/container in accordance with national regulations.

Supplemental Label Information

EUH066 Repeated exposure may cause skin dryness or cracking.

2.3 Other hazards

Not classified as PBT/vPvB by current EU criteria.

SECTION 3 COMPOSITION/INFORMATION ON INGREDIENTS

3.1 Substances

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

Product Name: ETHYL ACETATE

REACH Registration Number: 01-2119475103-46-XXXX

CAS-No: 141-78-6

EU Index No: 607-022-00-5

EC No: 205-500-4

Composition Comments: The data shown are in accordance with the latest EC Directives.

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: Remove affected person from source of contamination. Get medical attention if any discomfort continues.

In case of skin contact: Remove contaminated clothing immediately and wash skin with soap and water. Get medical attention if any discomfort continues.

In case of eye contact: Rinse immediately with plenty of water. Remove any contact lenses and open eyelids wide apart. Continue to rinse for at least 15 minutes. Get medical attention immediately. Continue to rinse.

If swallowed: Move affected person to fresh air and keep warm and at rest in a position comfortable for breathing. Rinse mouth thoroughly with water. Give plenty of water to drink. Get medical attention if any discomfort continues.

4.2 Most important symptoms and effects, both acute and delayed

If inhaled: Central nervous system depression including narcotic effects such as drowsiness, narcosis, reduced alertness, loss of reflexes, lack of coordination and vertigo.

In case of skin contact: Prolonged contact may cause redness, irritation and dry skin.

In case of eye contact: May cause temporary eye irritation.

4.3 Indication of any immediate medical attention and special treatment needed

Notes for doctor: No specific recommendations. If in doubt, get medical attention promptly.

SECTION 5 FIRE FIGHTING MEASURES

5.1 Extinguishing Media: Suitable extinguishing media: alcohol-resistant foam, carbon dioxide, dry powder or water fog.

5.2 Special Hazards Arising from the Substance or Mixture

Specific Hazards: Carbon oxides.

5.3 Advice for Fire-fighters

Protective Equipment for Fire-fighters: Positive-pressure self-contained breathing apparatus (SCBA) and appropriate protective clothing.

SECTION 6 ACCIDENTAL RELEASE MEASURES

6.1 Personal precautions, protective equipment and emergency procedures

Follow precautions for safe handling described in this safety data sheet. Take precautionary measures against static discharges. Avoid inhalation of vapours and contact with skin and eyes.

6.2 Environmental precautions

Spillages or uncontrolled discharges into watercourses must be immediately alerted to the Environmental Agency or other appropriate regulatory body.

6.3 Methods and material for containment and cleaning up

Absorb spillage with inert, damp, non-combustible material. Flush contaminated area with plenty of water. Collect and place in suitable waste disposal containers and seal securely. For waste disposal, see Section 13.

6.4 Reference to other sections

Wear protective clothing as described in Section 8 of this safety data sheet.

SECTION 7 HANDLING AND STORAGE

7.1 Precautions for safe handling

Avoid spilling. Avoid contact with skin and eyes. Keep away from heat, sparks and open flame. Provide adequate ventilation.

7.2 Conditions for safe storage, including any incompatibilities

Storage precautions: Store in tightly-closed, original container in a dry, cool and well-ventilated place. Keep away from heat, sparks and open flame.

Storage class: Flammable liquid storage.

7.3 Specific end use(s)

The identified uses for this product are detailed in Section 1.2.

SECTION 8 EXPOSURE CONTROLS/PERSONAL PROTECTION

8.1 Control parameters

Name	STD	TWA – 8 Hrs	STEL – 15 Min
Ethyl Acetate	WEL	200 ppm	400 ppm

WEL = Workplace Exposure Limit

Ingredient Comments

DNEL Industry Inhalation. 1468 mg/m3
DNEL Consumer Inhalation. 734 mg/m3
DNEL Industry Dermal Long Term 63mg/kg/day
DNEL Industry Inhalation Long Term 734 mg/m3
DNEL Consumer Dermal Long Term 37mg/kg/day
DNEL Consumer Inhalation Long Term 367mg/m3
PNEC Freshwater 0.26
PNEC Soil 0.22mg/kg
PNEC Sediment 0.34mg/kg
PNEC STP 650mg/l

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. Butyl rubber gloves are recommended.

(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. Other Protection: Wear rubber apron. Wear rubber footwear.

(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), fitted with type A2 gas filter cartridge.

SECTION 9 PHYSICAL AND CHEMICAL PROPERTIES

9.1 Information on basic physical and chemical properties

a) Appearance Form: colourless liquid

b) Odour Fruity

c) Odour threshold no data available

d) pH no data available

e) Melting point / freezing point -83.8°C

f) Initial boiling point and boiling range 76-77°C

g) Flash point -4°C closed cup

h) Evaporation rate 4.5 (diethyl ether=1)

i) Flammability (solid, gas) no data available

j) Upper/lower flammability or explosive limits 2.2% lower, 11.5% upper

k) Vapour pressure no data available

l) Vapour density 3.04

m) Relative density 0.899 – 0.903 @ 20°C

n) Solubility (ies) soluble in water

o) Partition coefficient: n-octanol/water 0.68

p) Auto-ignition temperature 427°C

q) Decomposition temperature no data available

r) Viscosity 0.4508 mPas @ 20°C

s) Explosive properties no data available

t) Oxidising properties no data available

9.2 Other information
Mol. Weight 88.11

SECTION 10 STABILITY AND REACTIVITY

10.1 Reactivity
No known reactivity hazards associated with this product.

10.2 Chemical stability
Stable under normal ambient temperature conditions and recommended use.

10.3 Possibility of hazardous reactions
Hazardous Polymerisation: Will not polymerise.

10.4 Conditions to avoid
Avoid excessive heat for prolonged periods of time. Avoid heat, flames and other sources of ignition.

10.5 Incompatible materials
Materials To Avoid: Strong oxidising substances.

10.6 Hazardous decomposition products
Oxides of: Carbon.

SECTION 11 TOXICOLOGICAL INFORMATION

11.1 Information of toxicological effects
Acute toxicity: (Oral LD50): 4934 mg/kg Rabbit OECD 401
(Dermal LD50): > 20000 mg/kg Rabbit OECD 404
Skin corrosion/irritation: Repeated exposure may cause skin dryness or cracking.
Serious eye damage/eye irritation: Slightly irritating.
Respiratory or skin sensitisation: Irritating to respiratory system. Vapours have a narcotic effect and may cause headache, fatigue, dizziness, nausea and vomiting.
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
Additional Information
Ingestion: May cause discomfort if swallowed. Narcotic effect.

SECTION 12 ECOLOGICAL INFORMATION

12.1 Toxicity
Ecotoxicity: The product components are not classified as environmentally hazardous. However, large or frequent spills may have hazardous effects on the environment.

Toxicity to Fish	
Ethyl Acetate	LC50 96 hours 230 mg/l Pimephales promelas (Fat-head Minnow)

Toxicity to Daphnia and other Aquatic Invertebrates	
Ethyl Acetate	NOEC 72 hours > 100 mg/l Daphnia magna

12.2 Persistence and degradability
The product is readily biodegradable.

12.3 Bioaccumulative potential
No data available. Partition coefficient: 0.68

12.4 Mobility in soil
The product is soluble in water. Surface Tension: 24 mN/m 20.

12.5 Results of PBT and vPvB assessment
Not Classified as PBT/vPvB by current EU criteria.

12.6 Other adverse effects
Not determined.

SECTION 13 DISPOSAL CONSIDERATIONS

13.1 Waste treatment methods
General Information: Waste to be treated as controlled waste. Disposal to licensed waste disposal site in accordance with local Waste Disposal Authority. Do not puncture or incinerate even when empty. Dispose of waste and residues in accordance with local authority requirements.

SECTION 14 TRANSPORT INFORMATION

General: Wear protective clothing as described in Section 8 of this safety data sheet.

14.1 UN number (ADR/RID/IMDG/ICAO) 1173

14.2 UN proper shipping name Ethyl Acetate

14.3 Transport hazard class(es)
ADR/RID/IMDG/ICAO Class: 3
ADR Label No: 3
IMDG Class: 3
ICAO Class/Division: 3
Transport label:

14.4 Packing group
ADR/RID/IMDG/ICAO Packing group: II

14.5 Environmental hazards No

14.6 Special precautions for user
EMS: F-E, S-D
Emergency Action Code: •3YE
Hazard Identification No. (ADR/RID): 33
Tunnel Restriction Code: (D/E)

SECTION 15 REGULATORY INFORMATION

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

National Regulations: The Chemicals (Hazard Information and Packaging for Supply) Regulations 2009 (S.I 2009 No. 716).

EU Legislation: Regulation (EC) No 1272/2008 of the European Parliament and of the Council of 16 December 2008 on classification, labelling and packaging of substances and mixtures (as amended).

Regulation (EC) No 1907/2006 of the European Parliament and of the Council of 18 December 2006 concerning the Registration, Evaluation, Authorisation and Restriction of Chemicals (REACH) (as amended).

This product may impact SEVESO storage regulations.

Guidance: CHIP for everyone HSG228.

Workplace Exposure Limits EH40.

Safety Data Sheets for Substances and Preparations.

Approved Classification and Labelling Guide (Sixth edition) L131.

DSEAR

Water Hazard Classification: WGK 1

15.2 Chemical Safety Assessment

A chemical safety assessment has not been carried out.

Inventory Information: TSCA EINECS PICCS NZIOC KECL ISHL ENCS

AICS DSL IECS

SECTION 16 OTHER INFORMATION

Hazard Statements in Full

H225 Highly flammable liquid and vapour.

H319 Causes serious eye irritation.

H336 May cause drowsiness or dizziness.

EUH066 Repeated exposure may cause skin dryness or cracking.

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.

EC REP

MDSS GmbH
Schiffaraben 41
30175 Hanover
Germany

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, 145800, 145900, 1460, 146200, 146300, 146400, 108900, 108910, 148998, 149910, 151000, 900000, 903000, 905000, 908000

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, United Kingdom
+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

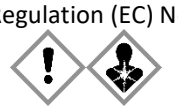
2.1 Classification of the substance or mixture

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.

EC REP

MDSS GmbH
Schiffaraben 41
30175 Hanover
Germany

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, 145501 1461, 146500, 146501, 108920, 149920, 249400, 901000, 906000

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, United Kingdom
+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.

EC REP

MDSS GmbH
Schiffaraben 41
30175 Hanover
Germany

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, 145501, 145800, 145900, 146300, 146400, 146500, 146501, 108900, 108910, 108920, 108935, 148998, 149910, 149920, 151000, 249400, 900000, 901000, 903000, 905000, 906000, 908000

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, United Kingdom
+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.

EC REP

MDSS GmbH
Schiffaraben 41
30175 Hanover
Germany

Midi Parasep® Faecal Parasite Concentrator

Code	Product
145000	Midi Parasep®
145002	Midi Parasep® Apafix™
145003	Midi Parasep® Apafix™ + 80ml Ethyl Acetate
145005	Midi Parasep® Apafix™
145200	Midi Parasep® Formalin
145300	Midi Parasep® Formalin & Triton X
145400	Midi Parasep® Formalin & Triton X + 80ml Ethyl Acetate
145500	Midi Parasep® SAF & Triton X
145501	Midi Parasep® SAF & Triton X + 80ml Ethyl Acetate

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: **orders@apacor.com**

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APA101
V8.6
06/2022