NSI水质微生物质控样



提供:

英文证书 中文使用说明书 项目分析确认值 质控接受区间

项目	浓度	货号	价格
埃氏&耐热&总大肠菌群	1000-2000CFU/100mL	NSI_9001H	¥ 450
	< 200CFU/100mL	NSI_9001L	¥ 450
大肠埃希氏菌	耐热大肠菌群	总大肠菌群	
脱水片水解于100mL无菌水中使用			
菌落总数	<i>5-500CFU</i>	NSI_HPCQC	
脱水片水解于100mL无菌水中使用			¥ 450

以下项目 整盒出售,10支/盒 - 2580元	货号	_
P.aeruginosa(NCTC 12951) 绿脓杆菌	10662-10	_
E.aerogenes(NCTC 10006) 产气肠杆菌	10006-10	
Klebsiell spp(NCTC 8167) 克氏杆菌	8167-10	
E.faecalis(NCTC 775)粪肠球菌	775-10	

电话: 021-38029051 微信: anacrm











Certificate of Analysis

E. coli, Total Coliform, and Fecal Coliform CRM

Catalog Number: 9001L Lot Number: 200327 Manufacture Date: 03/27/ Certified Date: 04/06/2020 03/27/2020 Expiration: 05/31/2021 trix: Flash Pellet Hazards: Infectious (See MSDS)

Analyte	Strain #	ATCC#	Certified <u>Activity</u> (MPN or CFU/100 mL)	Acceptance <u>Limits</u> (MPN or CFU/100 mL)
Escherichia coli (MPN) 1	NCTC 9001	11775	88 ± 12	19 - 425
Total Coliform (MPN) 1	NCTC 9001	11775	88 ± 12	18 - 436
Fecal Coliform (MPN) 2	NCTC 9001	11775	57 ± 7	10 - 361
Escherichia coli (MF) 3	NCTC 9001	11775	77 ± 11	19 - 389
Total Coliform (MF) 8	NCTC 9001	11775	77 ± 11	26 - 187
Fecal Coliform (MF) 4	NCTC 9001	11775	21 ± 4	4 - 136

Must store at -10°C to -20°C. Activity, uncertainty, and stability is based upon this storage temperature.

This certified reference material (CRM) is a lyophilized pellet packaged under vacuum in a 5 mL glass vial. This CRM must be rehydrated and analyzed in its entirety for certified activity and associated uncertainty to be applicable.

Precautions: Microbiological Reference Materials are designed for use for quality control in appropriately equipped microbiology laboratories by trained personnel. These reference materials contain viable microorganisms and should be handled according to appropriate biosafety level guidelines and disposed of according to applicable biohazard disposal regulations.

Principle, Explanation, & Reagents

This E. Coli, Total Coliform, and Fecal Coliform certified reference material (CRM) is a dehydrated pure culture of the organism produced by a proprietary process that yields stable, reliable, and cost effective samples that are homogenous and quantifiable. It is a hyphilized sample that must be rehydrated prior to use. Microorganism suspensions preserved by our process relatin their viability, biochemical profits and susceptibility patterns. The ingredients used to prepare the suspension preserve the microorganisms for use when needed. When rehydrated, the samples are ready for immediate use. Inactive ingredients may include: Albumin, Gelatin, Dextrose, Glycerol, Na-L-Ascorbate, Mannitol, Skim Milk, Trehalose, TSB, Na-L Glutamate, Sucrose, Page's Saline.

Instructions for Use:

Do not open the sample vial until the entire COA has been reviewed. Allow no more than 30 minutes to elapse from the completion of hydration and introduction of sample to media.

- 1. Retrieve a sample from the freezer and allow the capped sample vial to equilibrate to room temperature (15-30°C). Do not open vial until equilibration is complete. This should take approximately 15 minutes.
- Retrieve a 100 mL vial of sterile water or phosphate buffer. Once the sample is at room temperature, open the sample vial and aseptically transfer the pellet to the hydration fluid.
- 3. Once the transfer is complete, shake gently to dissolve. Full dissolution will take no more than 10 minutes.
- This is your sample for analysis. Analyze according to your usual laboratory procedures



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Certificate of Analysis

HPC Control CRM

Catalog Num Lot Number: 200511 Manufacture Date: 5/11/2020 Certified Date: 5/28/2020

Hazards: Infectious

<u>Analyte</u>	Strain#	ATCC#	Certification <u>Method</u>	A	ctiv	ied <u>ity</u> PN/mL	Limits(cfu	otance or MPN/mL) MPN/mL
Heterotrophic Organisms ¹ Heterotrophic Organisms ²	NCTC 775	19443	IDEXX Simplate® (MPN/mL) Plate Count (CFU/mL)	280 344	_	0.000	2*SD 207 - 354 298 - 389	3*SD 170 - 390 276 - 412

Packaging and Storage

Sample vial must be stored unopened at -10°C to -20°C. Activity, uncertainty, and stability are based upon this storage

The Heterotrophic Plate Count CRM is a lyophilized quick dissolve sample packaged in a 5 mL glass vial under vacuum that must be rehydrated prior to use. This CRM must be analyzed in its entirety for certified activity and associated uncertainty to be applicable.

Precautions

Microbiological Reference Materials are designed for use for quality control in appropriately equipped microbiology laboratories by trained personnel. These reference materials contain viable microorganisms and should be handled according to appropriate biosafety level guidelines and disposed of according to applicable biohazard disposal regulations.

Instructions For Use

Do not open the sample vial until the entire COA has been reviewed. Allow no more than 30 minutes to elapse from the completion of hydration and introduction of sample to media.

- Retrieve a sample from the freezer and allow the capped sample vial to equilibrate to room temperature (15-30 °C). Do not open vial until equilibration is complete. This should take about 15 minutes.
- 2) Retrieve a 100 mL vial of sterile water or phosphate buffer. Once the sample is at room temperature, open the sample vial and transfer the pellet to the hydration fluid. This transfer may be performed with sterile tweezers or simply by inverting the open sample
- Once the transfer is complete, cap the hydration fluid vial. Swirt the sample to dissolve. Full dissolution should take no longer than
- 4) Mix by inversion several times. This is your sample for analysis.
- Review certified activity to determine if a dilution of the hydrated sample is required for your method. Analyze according to your usual laboratory procedures.

Principle, Explanation, & Reagents

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This HPC standard is a dehiydrated pure culture of the organism produced by a proprietary process that yields stable, reliable, and cost effective samples that are homogenous and quantifiable. It is a dried sample that must be rehydrated prior to use. Microorganism suspensions preserved our process relatin their viability, blookhenical profile, and susceptibility patterns. The ingredients used to prepare the suspension preserve the microorganisms for use when needed. When rehydrated, the samples are ready for immediate use. Inactive ingredients may include: Albumin, Gelatin, Deutrose, Oliperol, Nat-Ascorbate, Mannicol, Skim Mills, Trehalose, TSB, Nat-Glutamate, Sucrosse, Page's Sallor.

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分析证明书 III 医菌群 质控制 埃氏、总大肠菌群、耐热(粪)

产品编号: 9001 产品批号。 200327



项目	<u>施认值</u> (MPN or CFU/100mL)	接受区间 (MPN or CFU/100mL)
大肠埃希氏菌 (MPN) - 爱德士法	88 ± 12	19 - 425
总大肠菌群(MPN)- 爱德士法	88 ± 12	18 - 436
耐热大肠菌群 (MPN) - 爱德士法	57 ± 7	10 - 361
大肠块希氏菌 (MF) - SM92228法	77 ± 11	19 - 389
总大肠菌群 (MF) - SM9222B法	77 ± 11	28 - 187
耐热大肠溶剂 (MF) - SM9222D法	21 + 4	4 - 136

包装及保存

样品必须-10℃~-20℃密封保存。活性、不确定度、稳定性均基于此保存条件设置

该标准物质密封在玻璃瓶中, 水解后使用, 样品一次整片使用, 不可分割使用

该标准物质用于微生物实验室《具有专业人员及设备》的质量控制。 该标物含微生物:操作及弃置应当依据适当的生物安全等接勤作特面及微生物私害弃置管制措施。

由于微生物的敏感性及水化后只能短时间保存(不超过30分钟),诸务必阅读全常的配制说明后再开始操作。诸严格按照配制说明操作,以确保微生物可用

成50%、排作下、5000 读标物为原水产品,趁过适当程序制备、稳定、可靠、可量化、均匀且价廉 使用前必须水解,依原提供的程序每年,可保持可用性、生化轮廓及镜源性。 需要时谓用到非活性物质。水解后立即使用。非活性物质可能包括,白蛋白、明胶...

溯源、用涂、不确定度、均匀性、稳定性、保质期

产品编号: HPCQC

产品批号。 200511



菌落总数 质控样

0 2021-11-30 危险性: 越染

QQ: 2447084087. 微信: anacrm Website: www.anacrm.com

项目 接受区间 确认信 (MPN or CFU/mL) (MPN or CFU/mL) 207 - 354 异养物 - IDEXX simplate 208 - 380 异养物 - Plate count

样品必须-10℃~-20℃密封保存。活性、不确定度、稳定性均基于此保存条件设置

该有证标准物质为5mL玻璃瓶真空封装的快速漆干拌,需水解使用。该标准物质必须整片分析,不可分割使用

在整 流転權物原用于微生物实验室(具有专业人员及设备>的质量控制。 流标物含微生物,操作及存置应当依把适当的生物安全等效操作指南及微生物危害弃置管制措施。

使用说明

在认真阅读本COA前切勿打开样品。稀释后仅有30分钟的分析时间

- 冰箱取出,回温15分钟至室温。 其他准备就给前勿打开样品,端气可能破坏样品。
- 取100ml无蔑水崇稀释派。样品一旦回温至55温。为开件品转售至6种液中。该过程可以用无测镜子夹取或直接领入一旦转移元税,温紧报查,回啶服荡以使拌品5分钟内完全外解
- 上下翻转10次混匀,所得即为要分析的溶液 浏览证书浓度,根据分析方法确定是否需要稀释

原则、解释、试剂

该赛高企数标准物度是一个有机物配水产品。该产品具有稳定的,均匀,可信赖,经济实惠等特点。该标准物质是股水干燥产品。 多须未鲜使用,我们的程序过程使数生物参浮液排转可显性。生化性,及要变性,为按该样品水解并保存特性效立即使用,用到的非古性崇拜成分可能包括。白蛋白,则股,葡萄糖,甘油,Na-L-冷凝度盐,进移,保力盐等
、Na-L-冷凝度盐,进移,保力盐等

溯源、用途、不确定度、均匀性、稳定性、保质期