Westase

Code No. 9005 Size: 1 g

Example of protoplast preparation:

(1) Saccharomyces cerevisiae, Trigonopsis variabilis

Medium YPG medium

Culture 25°C, 1 day, Shake culture (reciprocate at 120 rpm)

Conditions 0.5% Westase solution

0.5 M Na-tartrate McIlvain Buffer (pH 6.0) 30℃ Reciprocal shaking, 1 - 2 hr.

(2) Lipomyces starkeyi, Candida utilis

Medium Same as in (1)
Culture Same as in (1)

Conditions 0.5% Westase solution

0.4 M Na-tartrate McIlvain Buffer (pH 6.0) 30°C Reciprocal shaking, 1 - 2 hr.

(3) Shizosaccharomyces pombe, Hansenula mrakii, Kluyveromyces lactis,

Pichia anomala, Filobasidium floriforme, Candida colliculosa

Medium Same as in (1)

Culture Same as in (1)
Conditions 0.5% Westase solution
0.4 M Na-tartrate

McIlvain Buffer (pH 6.0) 30°C Reciprocal shaking, 3 - 4 hr.

(4) Kloeckera apiculata, Cryptococcus albidus

Medium Same as in (1)
Culture Same as in (1)

Conditions 0.5% Westase solution

0.3 M Na-tartrate
McIlvain Buffer (pH 6.0)
30°C Reciprocal shaking, 1 - 2 hr.

 Ustilago maydis, Graphiola phoenicis, Brettanomyces bruxellensis, Phaffia rhodozyma

Medium Same as in (1)

Culture 25°C, 2 days, Shake culture (reciprocate at 120 rpm)

Conditions Same as in (1)

(6) Tremella mesenterica

Medium Same as in (1)
Culture Same as in (5)

Conditions Same as in (2)

Manufactured by Ozeki corporation.

Description:

This product was prepared from liquid culture supernatant of Streptomyces rochei DB-34. This product has complex lytic activities of yeast cell mainly consisting of β -1,6 glucanase and β -1,3 glucanase activity.

Origin: Streptomyces rochei DB-34

Form: Lyophilized powder (containing celite as the excipient)

Storage: 4°C, dry condition.

Unit definition :

 β -1,6 glucanase activity : One unit is defined as the amount of enzyme

required to release 1 μ mol reducing suger from 10 mg/ml Pustulan solution in 1 min at

37°C, pH 6.0.

Lytic activity: One unit is defined as the amount of enzyme equired to cause a 1% decrease in absorbance

at 660 nm in 1 min at 30°C, pH 6.0 when using cell wall fraction of *Cryptococcus albidus* IFO

0612 as a substrare.

Quality Control Data:

Please see the Certificate of Analysis (CoA) for each lot. You can download the CoA on Takara Bio website.

Preparation of the enzyme solution:

(1) Dissolve in McIlvain Buffer.*

- * McIlvain Buffer is prepared by mixing 0.1 M Citric acid solution and 0.2 M Disodium hydrogenphosphate at the ratio of around 36.8: 63.2 (y/y), and adjust to pH 6.0.
- (2) Filter the solution with cellulose acetate filter.

Note

- (1) Na-tartrate (0.3 0.5 M) must be used as an osmotic stabilizer to form protoplast by Westase. The use of sorbitol as a stabilizer remarkably reduces the efficiency of forming protoplast.
- (2) For preparation of protoplasts by Westase, the yeast cells in exponential phase are suitable. In most yeast strains, the cells of stationary phase are not appropriate since they can result in lower efficiency.
- (3) To regenerate the cell wall of protoplasts, use 1.0 1.5 M sorbitol as anosmotic stabilizer.

Note

This product is for research use only. It is not intended for use in therapeutic or diagnostic procedures for humans or animals. Also, do not use this product as food, cosmetic, or household item, etc. Takara products may not be resold or transferred, modified for resale or transfer, or used to manufacture commercial products without written approval from Takara Bio Inc.

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Westase

本製品は、Streptomyces rochei DB-34 の液体培養上清より調製された、

 β -1,6 グルカナーゼ、 β -1,3 グルカナーゼ活性を主体とする酵母細胞壁

β-1,6 グルカナーゼ活性: 37 ℃、pH6.0 において 10 mg/ml の Pustulan

る酵素量を10とする。

きの活性を 1 U とする。

性能試験結果については、各ロットのCertificate of Analysis (CoA)をご覧

ください。CoAはタカラバイオウェブサイトからダウンロードできます。

McIlvain Buffer (0.1 M クエン酸溶液と、0.2 M リン酸水素ニナトリウム

溶液を約36.8:63.2 で混合し、pH6.0 に調製する) に適当量溶解し、セ

(1) Westase を用いるプロトプラスト化の浸透圧調整剤には、必ず酒石酸ナトリウムをご使用ください。ソルビトールなどでは十分なプロトプ

(2) プロトプラスト化には対数増殖期の酵母をご使用ください。定常期で

(3) プロトプラストの再生には、1.0~1.5 M ソルビトールをご使用くだ

は著しくプロトプラスト化の効率が落ちることがあります。

ルロースアセテートフィルターでろ過してから使用する。

凍結乾燥粉末 [賦形剤として珪藻土(セライト)を含む]

溶液から1分間に1 µmolの還元糖を遊離す

Cryptococcus albidus IFO 0612 の細胞壁画分

を基質にし、30℃、pH6.0 において、1 分間 に 660 nm における吸光度が 1% 減少すると

Streptomyces rochei DB-34

4℃、乾燥状態で保存

溶解用複合酵素剤である。

●由来

●形状

● 保存

● 活性の定義

細胞壁溶解活性:

● 品質管理

● 使用方法

● 使用上の注意

さい。

ラスト形成が起こりません。

Code No. 9005 容量: 1 g

● プロトプラスト調製条件例

(1) Saccharomyces cerevisiae, Trigonopsis variabilis

培地 YPG medium

培養条件 25℃、1 day、Shake culture (reciprocate at 120 rpm)

調製条件 0.5% Westase solution

0.5 M Na-tartrate McIlvain Buffer (pH6.0)

30°C Reciprocal shaking, $1 \sim 2 \text{ hr.}$

● 製品説明 (2) Lipomyces starkeyi, Candida utilise

培地 Same as in (1) 培養条件 Same as in (1) 調製条件 0.5% Westase solution

0.4 M Na-tartrate
McIlvain Buffer (pH6.0)

30°C Reciprocal shaking、1 \sim 2 hr.

(3) Shizosaccharomyces pombe, Hansenula mrakii, Kluyveromyces lactis, Pichia anomala, Filobasidium floriforme, Candida colliculosa

> 培地 Same as in (1) 培養条件 Same as in (1) 調製条件 0.5% Westase solution 0.4 M Na-tartrate Mcllvain Buffer (pH6.0)

30°C Reciprocal shaking $\sim 4 \text{ hr.}$

(4) Kloeckera apiculata, Cryptococcus albidus

培地 Same as in(1)

培養条件 Same as in (1) 調製条件 0.5% Westase solution 0.3 M Na-tartrate

0.3 M Na-tartrate
McIlvain Buffer (pH6.0)

30°C Reciprocal shaking $1 \sim 2 \text{ hr.}$

(5) Ustilago maydis, Graphiola phoenicis, Brettanomyces bruxellensis, Phaffia rhodozyma

培地 Same as in (1)

培養条件 25°C、2 days、Shake culture (reciprocate at 120 rpm)

調製条件 Same as in (1)

(6) Tremella mesenterica

培地 Same as in (1) 培養条件 Same as in (5) 調製条件 Same as in (2)

● 製造元 大関株式会社

●注意

本製品は研究用として販売しております。ヒト、動物への医療、臨床 診断用には使用しないようで注意ください。また、食品、化粧品、家 庭用品等として使用しないでください。

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タカラバイオ株式会社

製品についての技術的なお問い合わせ先

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