

#PFS052-10-EX For 2 mL x5 Reaction

PUREfrex® is NOT included.

in vitro research use only Store at -80°C before opening

Feb 2023



GeneFrontier Corporation www.genefrontier.com

SHARP Kashiwa Building, 4F 273-1 Kashiwa, Kashiwa-shi, Chiba 277-0005 Japan

Introduction

1. About PURE frex®

PUREfrex® is a reconstituted cell-free protein synthesis kit which GeneFrontier has developed based on the PURE system technology. The target protein can be synthesized by adding the template DNA (or mRNA) to the reaction mixture. The PURE system is a unique cell-free protein synthesis system, which has originally developed by Professor Takuya Ueda at the University of Tokyo, and consists of only purified factors necessary for transcription, translation and energy regeneration (Ref. 1). Therefore it enables to adjust the composition of the reaction mixture.

PURE frex® has been raised in purity by improving the methods for preparing ribosomes, tRNAs and all proteins in the reaction mixture compared with the original PURE system (Ref. 2). As the result, the contaminating lipopolysaccharide from E. coli is reduced to around 0.1 EU per 1 μ L of reaction and other contaminants, such as RNase and β -galactosidase, are also reduced.

All of proteins in PURE frex® have no tags, the synthesized protein can be purified and detected by any tags.

References) 1. Shimizu *et al.* (2001) *Nat. Biotecnol.*, vol. 19, p. 751 2. Shimizu *et al.* (2005) *Methods*, vol. 36, p. 299

Introduction

2. About EF-P

EF-P (elongation factor P) is one of the translation factors in *Escherichia coli* (*E. coli*) and a homolog of eukaryotic initiation factor 5A (eIF5A). EF-P improves the synthesis of protein containing consecutive proline residues such as Pro-Pro-Pro and Pro-Pro-Gly by promoting the formation of peptide bonds. Lysine at 34th of EF-P is post-translationally modified to β-lysillysine and the modification is important for the activity.

Because PURE $frex \otimes 1.0$, PURE $frex \otimes 2.0$ and PURE $frex \otimes 2.1$ don't contain EF-P, the synthesis efficiency of some proteins containing consecutive proline residues are low. Addition of EF-P to the PURE $frex \otimes 1.0$ reaction mixture increase the productivity of such proteins.

EF-P (#PFS052-0.5) is a supplement for PURE frex® that contains recombinant post-translationally modified EF-P isolated from E. coli.

Note

EF-P and PURE frex® is developed for $in\ vitro$ research use only. EF-P and PURE frex® should not be used for the therapy, diagnostic or administration to animals including human and should not be used as food or cosmetics etc.

To avoid the contamination of nuclease, nuclease-free-treated water, reagents and materials should be used. We also recommend wearing gloves and mask

For information concerning commercial use of EF-P, please contact GeneFrontier.



GeneFrontier Corporation www.genefrontier.com

Distributor



COSMO BIO USA [Outside Japan] 2792 Loker Ave West, Suite 101 Carlsbad, CA 92010, USA email: info@cosmbiousa.com Phone/FAX: (+1) 760-431-4600 URL: www.cosmobiousa.com



COSMO BIO CO., LTD.

TOYO EKIMAE BLDG. 2-20, TOYO 2-CHOME, KOTO-KU. TOKYO 135-0016, JAPAN Phone: +81-3-5632-9610 FAX: +81-3-5632-9619 URL: https://www.cosmobio.co.ip/

Kit components

EF-P *1

50 μL x5

40 µM EF-P (in 30 % glycerol buffer)

Store at -80°C*2

Dilution Buffer

500 μL x1

Store at lower than -20°C

Kit components

Store at -80 °C before opening

*1\

Standard final concentration of EF-P is $0.1-2~\mu M$. We recommend to check the optimal concentration of EF-P because it depends on a protein of interest.

*2)

For storage at -80°C, the rest of solution should be frozen rapidly in liquid nitrogen or dry ice/ethanol. Please divide into aliquots, if necessary, and avoid refreeze and thaw as much as possible.

Protocol

Here is a standard protocol for synthesizing proteins using EF-P and PURE frex® 2.1 (#PF213). For example, please assemble 20 µL of reaction mixture as below, in which the final concentration of each reagent is 0.5 mM Cysteine, 4 mM GSH and 1 µM EF-P.

- Thaw Solution I, Cysteine and GSH by incubation at room temperature or 37 °C for 1 minute completely, and then cool on ice.
- 2. Thaw Solution II, III and EF-P on ice.

. . . .

- 3. Mix each solution by vortex and centrifuge briefly to collect each solution at the bottom.
- 4. Assemble the reaction mixture in a tube as follows. (Add the template DNA to 0.5-3 $ng/\mu L$ per 1 kbp)

Water	6.5-X μL
Solution I *3	8 µL
10 mM Cysteine	1 μL
80 mM GSH	1 μL
Solution II	1 μL
Solution III	2 μL
40 μM EF-P	0.5 μL
Template DNA	XμL
Total	20 μL

Protocol

- 5. Incubate the tube at 37°C for 2-6 hours with heat block or water bath.
- 6. Analyze the synthesized product.

*3)

Please note that the volume of Solution I of PURE frex $^{\circ}$ 2.1 (#PF213) is different from PURE frex $^{\circ}$ 2.0 (#PF201).