Approved and current. Effective starting 10/3/2022. COA-456255 (version 2.0) Human CYP2B6 + P450 Reductase +Cytochrome b5 Supersomes

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Human CYP2B6 + P450 Reductase + Cytochrome b₅ SUPERSOMES[™]

Catalog Number......456255 Lot Number......2404010 Storage Conditions..STORE AT -80°C Date Released2024 April Expiration Date......2034 April

Package Contents	0.5 nmol cytochrome P450 in 0.5 mL		
Protein Content	14 mg/mL in 100mM potassium phosphate (pH 7.4)		
Cytochrome c Reductase Activity	2300 nmol/(min x mg protein)		
Cytochrome b5 Content	150 pmol per mg		
Cytochrome P450 Content	1000 pmol per mL		
7-Ethoxy-4-Trifluoromethyl- coumarin Deethylase Activity	18 pmol product/(min x pmol P450)		

7-Ethoxy-4-trifluoromethylcoumarin deethylase activity has replaced 7-ethoxycoumarin deethylase activity for this enzyme. 7-Ethoxy-4trifluoromethylcoumarin deethylase is more sensitive and specific for CYP2B6. We have found that specific activities can vary with different lots of substrate. This activity is catalyzed by CYP2B6 which is expressed from human CYP2B6 cDNA using a baculovirus expression system. Baculovirus infected insect cells (BTI-TN-5B1-4) were used to prepare these microsomes. These microsomes also contain cDNA-expressed human P450 reductase and human cytochrome b_5 . A microsome preparation using wild type virus (Catalog No. 456201) should be used as a control for native activities.

METHOD: A 0.25 mL reaction mixture containing 10 pmol P450, 1.3 mM NADP+, 3.3 mM glucose-6-phosphate, 0.4 U/mL glucose-6-phosphate dehydrogenase, 3.3 mM magnesium chloride and 0.1 mM 7-ethoxy-4-trifluoromethylcoumarin in 50 mM potassium phosphate (pH 7.4) was incubated at 37°C for 15 minutes. After incubation, the reaction was stopped by the addition of 50 μ L 20% trichloroacetic acid and centrifuged (10,000 x g) for 1 minute. 100 μ l of the supernatant was added to 1.9 mL 100 mM Tris (pH 9) and the fluorescence was determined with excitation at 410 nm and emission at 510 nm in a spectrofluorometer. The activity was quantitated by subtracting the fluorescence of the blank and comparing to a standard curve for 7-hydroxy-4-trifluoromethylcoumarin.

Time Course of Product Formation

ADVICE

- Thaw rapidly in a 37°C water bath. Keep on ice until use.
- Aliquot to minimize freeze-thawing cycles. Less than 20% of the catalytic activity is lost after 6 freeze thaw cycles.
- Metabolite production with 7-ethoxy-4-trifluoromethylcoumarin is linear with respect to enzyme concentration up to at least 80 pmol P450 per ml.
- Metabolite production with 7-ethoxy-4-trifluoromethylcoumarin is approximately linear for 50 minutes (see graph above).
 Other substrates may not exhibit similar linearity with respect to incubation time.
- Western immunoblotting indicates the expressed CYP2B6 has the same mobility as CYP2B6 in human liver microsomes.
- Comparison of Western immunoblotting intensity and spectral P450 contents for this product and human lymphoblastexpressed CYP2B6 indicates that a substantial amount of apoprotein is found in this product.

For research use only. Not for use in diagnostic or therapeutic procedures.

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INSECT CELL MICROSOMES SAFETY INFORMATION

HAZARD WARNING:

The product was produced using baculovirus (*Autographa californica*) infected insect cells (BTI-TN-5B1-4). This virus is not known to be pathogenic to humans or other mammals.

SAFETY RECOMMENDATIONS:

When using this product, follow good laboratory safety procedures:

Do not eat, drink or smoke.

Avoid contact with skin or eyes.

Do not inhale aerosols.

Do not pipette by mouth.

Wear suitable protective clothing, gloves and eye protection.

Steam sterilize product or treat product with a 1% solution of sodium hypochlorite prior to disposal.

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23 April 2024

Quality Assurance

Date

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Revision History				
Rev.	Change No.	Description of Change	Revised By	Revised Date
1	CC-02544	Updated to Corning branding, protected variable fields, replace safety section with safety statement and clarify SDS [Safety Data Sheet] information, addition of QA signature and date	L. Brown	7/23/14