abcam

Product datasheet

CYP2D6 Activity Assay Kit (Fluorometric) ab211078

3 Images

Overview

Product name CYP2D6 Activity Assay Kit (Fluorometric)

Detection method Fluorescent

Sample type Cell Lysate, Microsomes, Tissue Lysate

Assay type Enzyme activity (quantitative)

Species reactivity Reacts with: Mammals, Other species

Product overview CYP2D6 Activity Assay Kit (Fluorometric) (ab211078) allows rapid measurement of native or

recombinant cytochrome P450 2D6 (CYP2D6) activity in biological samples such as liver microsomes. The assay utilizes a non-fluorescent CYP2D6-selective substrate that is converted into a highly fluorescent metabolite detected in the visible range (Ex/Em = 390/468 nm), ensuring a high signal-to-background ratio with little interference by autofluorescence. CYP2D6 specific activity is calculated by running parallel reactions in the presence and absence of the potent CYP2D6-selective inhibitor quinidine and subtracting any residual activity detected with the

inhibitor present.

The kit contains enough reagents to perform 100 sets of paired reactuibs (in presence/absence

of inhibitor).

This product is manufactured by BioVision, an Abcam company and was previously called K703 Cytochrome P450 2D6 (CYP2D6) Activity Assay Kit (Fluorometric). K703-200 is the same size

as the 200 test size of ab211078.

Cytochrome P450 2D6 (CYP2D6, EC 1.14.14.1) is a member of the cytochrome P450 monooxidase (CYP) family of microsomal xenobiotic metabolism enzymes. CYPs are membrane-bound hemeproteins responsible for Phase I biotransformation reactions, in which lipophilic drugs and other xenobiotic compounds are converted to more hydrophilic products to facilitate excretion from the body. CYP2D6 catalyzes oxidation of lipophilic bases with an aromatic ring and a nitrogen atom and is highly expressed in liver and brain tissue. The enzyme is responsible for metabolism of nearly 25% of all small molecule drugs commonly used by humans, particularly psychiatric drugs such as antidepressants, antipsychotics and stimulants. The CYP2D6 gene is highly polymorphic in the human population, with CYP2D6 activity ranging from complete metabolic deficiency to ultra-rapid metabolism. Due to this wide phenotypic variability, CYP2D6 is

frequently implicated in drug toxicity and clinical drug/drug interactions.

Platform Microplate reader

Notes

1

Properties

Storage instructions

Store at -20°C. Please refer to protocols.

Components	200 tests
β-NADP ⁺ Stock 100X (2 mg)	1 vial
AHMC Standard	1 x 25µl
CYP2D6 Assay Buffer	1 x 100ml
CYP2D6 Inhibitor (Quinidine) (150 μg)	1 vial
CYP2D6 Substrate (200 μg)	1 vial
NADPH Generating System 100X (25 mg)	1 vial
Recombinant Human CYP2D6 (15 mg)	1 vial

Function

Responsible for the metabolism of many drugs and environmental chemicals that it oxidizes. It is involved in the metabolism of drugs such as antiarrhythmics, adrenoceptor antagonists, and tricyclic antidepressants.

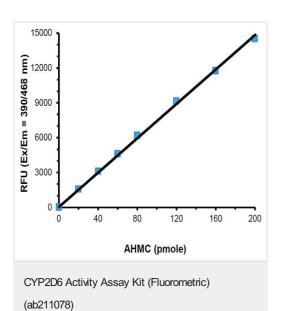
Sequence similarities

Belongs to the cytochrome P450 family.

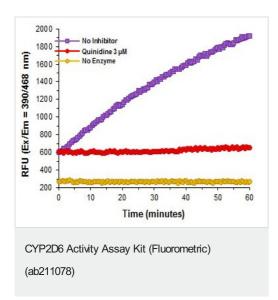
Cellular localization

Endoplasmic reticulum membrane. Microsome membrane.

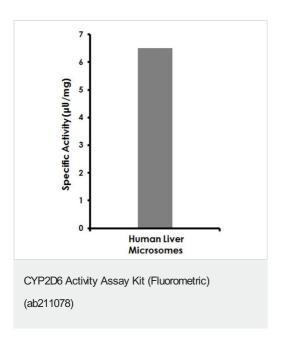
Images



Typical AHMC standard calibration curve. One mol of AHMC corresponds to the metabolism of one mol of CYP2D6 substrate.



Reaction kinetics of fluorogenic substrate metabolism in human liver microsomes (0.5 mg/mL) at 37°C in the presence and absence of the selective CYP2D6 inhibitor quinidine (the no inhibitor reaction contained a final concentration of 0.5% acetonitrile).



Specific activity of CYP2D6 in pooled human liver microsome sample (0.5 mg/mL).

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