

Contract service

Proteomics specialists offer protein quantitative analysis of metabolic pathway-wide proteome profiling using the latest MS instrument & MRM Standard Mix (Page 2).

Available panel: Human major metabolic enzyme (338 proteins)

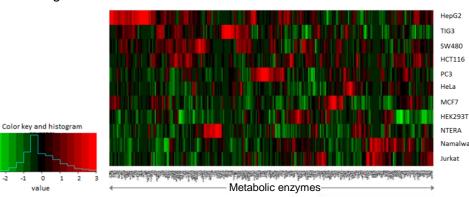
Sample preps: Frozen pellets of human cultured cells or frozen human tissue

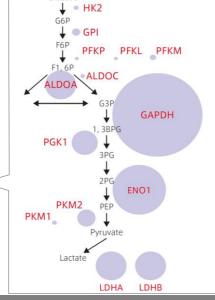
*Sample must be sent with serum free condition.

Example #1: Quantitative Analysis of Metabolic Enzymes in 11 Human Cancer Cell Lines

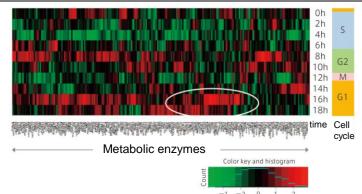
Bottom: Quantitative analysis was performed against 266 x human major metabolic enzymes in 11 human cancer cells. The expression levels were shown in the heat map below. Distinguished expression patterns among cell lines, which are clustered by the profile similarities. iMPAQT enables to show important factors on each pathway by calculating enzyme expression as quantitative value.

Right: Bubble charts of expression levels for glycolysis enzymes in HeLa cells. Expression levels of upstream enzymes were low, while those of downstream ones were high.





Example #2: Profiles of Human Metabolic Enzymes among Cell Cycles



After synchronizing cell division with thymidine block, HeLa cells were collected at every 2 hours and 236 x metabolic enzyme expressions were measured.

In S phase, expressions of many metabolic enzyme are suppressed (Green). In G1 phase, the expression level related to glycolysis and TCA cycle were greatly increased

※Please ask your local distributor about quote.

This service uses synthetic peptides as standard, so obtained quantitative value is standardized by synthetic peptides (It does not reflect protein purification at pre-treatment and efficiency of enzyme digestion). Obtained quantitative value is calculated from concentration of inner standard peptide.

X All products here are research use only, not for diagnostic use * Specs might be changed for improvement without notice

- Company name and product name are trademark or registered mark
- * Please contact your local distributors for orders, quote request and inquir

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Bio-REV Pte. Ltd.

Block 211 Woodlands Avenue 9 #08-78/79 Woodlands Spectrum II

Singapore 738 960 TEL: +65-6555-9001 FAX: +65-6555-9030

Email: sales@bio-rev.com

Technical Support: techserv@bio-rev.com

Funakoshi Co., Ltd.

Address: 9-7 Hongo 2-Chome, Bunkyo-ku,

Tokyo 113-0033 JAPAN Phone: +81-3-5684-6296 : +81-3-5684-6297 Email: export@funakoshi.co.jp

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A new proteomics technology by LC-MS/MS/MRM

Quantitative proteomics tools & analysis service by iMPAQT method



More details: Web page No

Quantitative tools 81385



We offer product & service to quantify multiple proteins at one time by mass spectrometer (LC-MS/MS)

iMPAQT*1, New technology for Quantitative Proteomics

*1 in vitro proteome-assisted MRM for Protein Absolute QuanTification Matsumoto, M., et al., Nat. Methods, 14(3), 251~258(2017).[PMID: 28267743]

- Developed by Drs. Nakayama & Matsumoto of Kyushu University.
- Highly ionized peptides by actual measurement of genome-wide human recombinant proteins
- Quantitative profiling of ~400 proteins/hour at once
- A powerful tool for pathway-wide proteomics, subtype clarification and no antibodies available targets

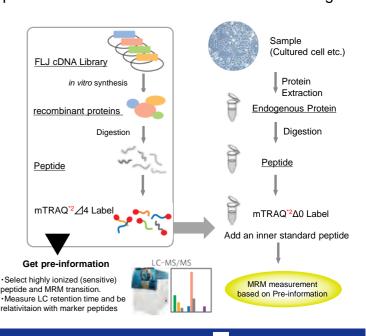
Technology Outline

- 1 In vitro synthesis of 18,000 recombinant human proteins, followed by digestion to peptides & mTRAQ*2 (heavy) labeling.
- 2 Acquire pre-information (MRM transition parameters, ionization values, relative retention time) of every peptides by LC-MS/MS.
- 3 MRM-measurement based on the pre-information for the samples & internal standards labeled with mTRAQ*2 (light for samples, heavy for standards)

√ Can perform LC-MS/MS analysis

Product available!

*2 mTRAQ reagent is product of AB Sciex.





Funakoshi offer



✓ No compatible equipments... ✓ No technitians/skills...

Service available!



Quantitative Proteomics Service by iMPAQT



Mixture of standard peptides for human metabolic enzyme.

MRMplus Standard Mix

Quantitative Proteomics tool for iMPAQT

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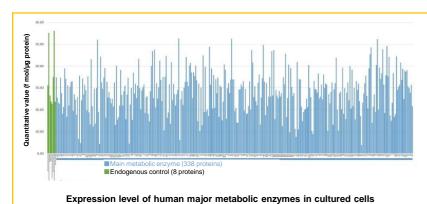


Standard peptides mixture for LC-MS/MS MRM - 338x Human major metabolic enzymes can be quantified-**MRMplus Standard Mix**

Workflow: Preparation of Scheduled MRM by MRMplus®



A standard peptides mixture of human major metabolic enzymes selected by iMPAQT.



Features

Measurable peptides in the Mix!

Total 338 of major human metabolic enzymes from carbohydrate metabolism, lipid metabolism, amino acids metabolism, nucleic acid metabolism,

Only 2 days!

(Sample prep.: 1 day + MS analysis: 1 day) Expression level of 338x metabolic enzymes in one profile.

Other Features

- Simple: Well-validated 338 human major metabolic enzymes & 8 endogenous controls are pre-mixed, just mTRAQ-label & Go!
- Easy: Pre-set parameters (MRM transition, relative retention time) will be provided.

Application

- Visualization of up/down-regulations on whole metabolic pathway.
- Drug discovery/Pharmacology
- Basic research for proteomics

Discription		
Maker	Product code	Size
MRMplus	Standard Mix, Metaboli	c Enzyme Deck
FMI	FMI-METMIX	200μg
Content:	peptide cocktail (lyoph	ilized), method file, datasheet
For 25 ind	ependent projects.	
XTo orde	r, "letter of assurance" is r	needed.

Related Products

■ Proteomics Sample Preparation Kit for Cultured Cell 2G

Cell lysis & peptide preparation for mass spectrometry by cultured cell sample, optimized for mTRAQ "conjugation for LC-MS/MS

*1: mTRAQ is a product from AB Sciex

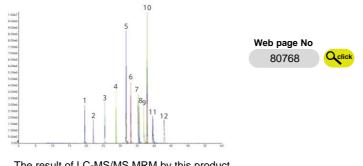
■Protocol overview

- Cell lysis
- BCA assay*2
- Protease digestion & alkylation
- mTRAQ *2 Labeling.
- Freeze-drying (-65oC).
- *2: BCA assay kit and mTRAQ reagents are not provided but required.

Discription		
Maker	Product code	Size
Proteomi	cs Sample Preparation K	it for Cultured Cell 2G
FMR	FMR-003	1 kit
Content: T	ris HCl, SDS, urea, soluble b	uffer, protease, reagent A/B/C, store tube
Enable to	analyze 10 times .	
:XTo orde	er "letter of assurance" is no	eded

■ MRMplus Retention Time Marker

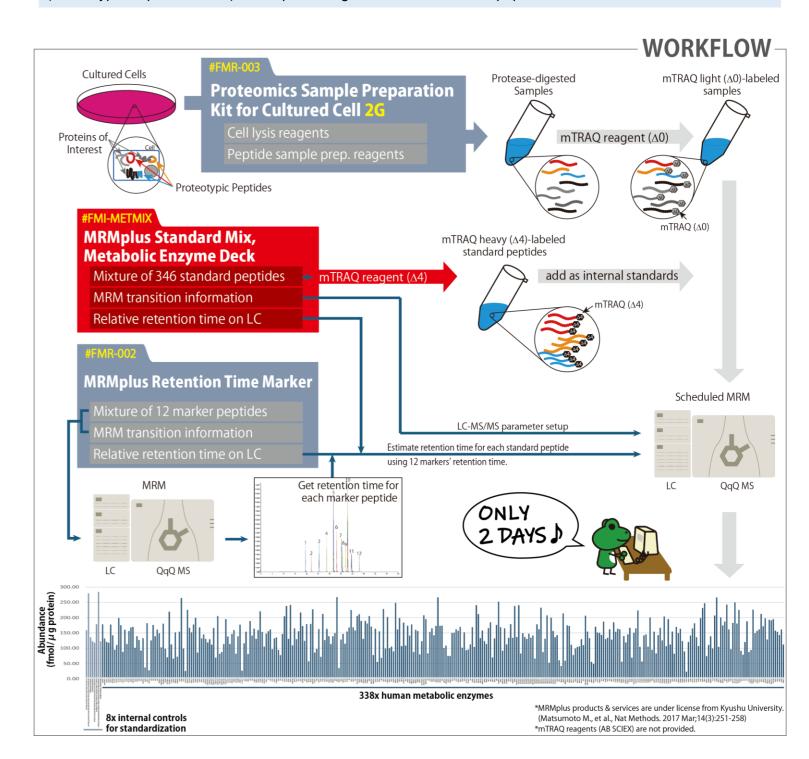
A mixture of 12 synthetic peptides for calibration of retention times for each peptides in LC-MS/MS system. Compatible with MRMplus Standard Mix.



The result of LC-MS/MS MRM by this product

iscription Maker	Product code		Size
MRMplus	Retention Time Ma	arker	
FMR	FMR-002	-80°C	12µl

MRM (Multiple Reaction Monitoring) and is one of the quantitative proteomics methods by mass spectrometry. MRM uses triple-quadrupole LC-MS/MS and enable us to distinguish peptides by their amino acid sequences specifically by detecting product/fragment ions of selected MS parameters (MRM transitions). It is possible to quantify proteins of interest (POIs) by measuring MRM transitions of specific sequences (Proteotypic Peptides: PTPs) in samples along with internal standard peptides.



^{*}This product is licensed from Kyushu university

^{*}This product requires triple-quadrupole (QqQ) mass spectrometry (AB Sciex 4000QTRAP or later).

^{*}mTRAQ reagent (AB Sciex) is required but not provided.

^{*}Compatible with conventional LC: Column inner diameter should be less than 2.1mm.