

## High-Throughput Quantitative Methylation Profiling with EpiTYPER<sup>®</sup> and the MassARRAY<sup>®</sup> System

- > A complete system for discriminating methylated vs. non-methylated DNA; includes software, reagents, and MALDI-TOF mass spectrometry for detection and quantification.
- Interrogate 10s 100s of samples and CpG sites in amplicons from 200 600 bp and detect down to 5% differences in methylation.



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### **Technology Overview**

Agena Bioscience's DNA methylation analysis technology (EpiTYPER) is one of the most reliable quantitative methods available today for analyzing DNA methylation changes<sup>1,2</sup>. The technology, which has been referenced in more than 1,000 peer-reviewed journal articles, includes the following components:

- EPIDESIGNER Software for genomic target selection and PCR primer design.
- EPITYPER REAGENT SETS Includes all reagents and consumables needed after bisulfite treatment of DNA.
- MASSARRAY SYSTEM MALDI-TOF mass spectrometer for robust and precise signal detection and quantification.
- EPITYPER REPORTING SOFTWARE For data analysis and graphical presentation of the level of methylation at each CpG site in each sample.

#### Benefits of the EpiTYPER DNA Methylation Analysis Technology

## Efficient

- Go from bisulfite-treated DNA to data in 8 hours.
- Covers multiple CpGs in amplicons of up to 600 bp.
- Compatible with many sample types, including formalin-fixed paraffin-embedded tissue.



High precision (5% CV).

## M Sensitive

 Detects down to 5% change in methylation levels.

## Cost Effective

- 96- and 384-well formats available
- Multiple CpGs analyzed in one simple reaction, from amplicons as long as 200-600 bp.

## 😧 Simple Workflow

- No need to design CpG-specific primers.
- No PCR product purification needed.
- Ideal for investigating a few or several hundred target regions.
- Convenient software solutions for comparison between samples.



#### ASSAY WORKFLOW

#### **STEP 1** ASSAY DESIGN

Enter your target sequences and the software determines primer designs for the most complete DNA coverage. In addition to optimized primer sequences, EpiDesigner delivers an easy-to-read graphical interpretation of the amplicons designed over your target regions, as well as annotating distinct CpG sites covered by the assays.

#### **STEP 2** BISULFITE TREATMENT

Bisulfite treatment converts any non-methylated cytosine residues into uracil, while methylated cytosine residues are unaffected. This step results in the generation of methylation-dependent sequence changes in the DNA template.



#### STEP 3 PCR, IN VITRO TRANSCRIPTION, AND RNA CLEAVAGE DESIGN



RNA transcription and uracil-specific cleavage using RNAse A.

The EpiTYPER assay starts with PCR using T7promoter-tagged reverse primers to amplify the target regions while preserving the bisulfite-induced sequence changes. After SAP treatment, in vitro transcription is performed and the resulting RNA transcripts are specifically cleaved at uracil residues. The resulting fragments differ in size and mass, depending on the sequence changes generated through bisulfite treatment. This difference allows the data analysis software to generate quantitative information for each analyzed target fragment.



#### **STEP 4** DATA ACQUISITION AND ANALYSIS

The EpiTYPER reaction products are dispensed onto a SpectroCHIP® Array (chip). The chip is then placed in the MALDI-TOF mass spectrometer for data acquisition. The results are automatically loaded into a database for data analysis with EpiTYPER software.



Easily determine percent methylation

## **EpiTYPER Software**

The EpiTYPER software provides an advanced and convenient solution for the quantitative analysis of CpG methylation. Numerical and graphic tools are available and the data are automatically matched to a reference sequence. Basic statistical analysis and confidence ratings are available for built-in quality control.

#### DATA ANALYZER MODE

Customized visual presentation of amplicon data in the most useful format for your needs. Sequence view shows the entire forward and reverse nucleotide sequence for the selected amplicon.



# Amplican QCC03\_JM32053\_255: Sample 100\_meth\_2: Sum of 7 of 7 Spectra Minimum Concernation of the second se

#### CUSTOM VIEWS OF AMPLICONS

Graphical representations of the CpG sites as epigrams within the selected amplicon enable easy comparison of samples and CpG sites.

Catalog No.	Item	# Wells/Pads	Chip Format
MassCLEAVE Reagent Kits			
13165F	MassCLEAVE, PCR Reagent And SpectroCHIP Kit	960	CPM 96
11377D	MassCLEAVE, PCR Reagent And SpectroCHIP Kit	3840	CPM 384
13165	MassCLEAVE, PCR Reagent And SpectroCHIP Kit	960	96
11377	MassCLEAVE, PCR Reagent And SpectroCHIP Kit	3840	384
Bisulfite Conversion Kits			
10131	EZ 96 DNA Bisulfite Treatment Kit (96-well Plate Format)	192	N/A
10132	EZ DNA Bisulfite Treatment Kit (Tube Format)	50	N/A

#### **ORDERING INFORMATION**

Reagent sets are available in 96- and 384-well formats, and are designed for use with the MassARRAY System.

#### References

- 1. A systematic comparison of quantitative high-resolution DNA methylation analysis and methylation-specific PCR. Claus R, et al. Epigenetics 7(2012) 772-780.
- Quantitative high-throughput analysis of DNA methylation patterns by base-specific cleavage and mass spectrometry. Ehrich M, et al. PNAS 102(2005) 15785-15790.

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