THE MULTIFLOW® FAMILY OF KITS

Multiple endpoints. One step.

The Method

In just one step:

reagents are added to cells...

outer membranes are lysed, nuclei are stained...

and fluorescent antibodies bind to nuclear epitopes.

MultiFlow add-and-read kits were developed from the ground up to be simple and efficient.

When microtiter plates are used to collect these multiplexed data, the amount of test compound necessary for analysis is dramatically reduced.

At this point nuclei are ready for flow cytometric analysis.

Benefits

- High information content
- Multiplexed assay format

- Simple and efficient
- Fast, flow cytometric analysis
- Scalable compatible with microtiter plates and robotic liquid handlers

Available Kits

Kit Name Applications

MultiFlow – Cleaved PARP	Apoptosis	
MultiFlow – γH2AX	Double strand DNA breaks	Each kit also provides: Cell density Cell proliferation Cytotoxicity
MultiFlow – Phospho-Histone H3	Mitotic cells	
MultiFlow – p53	Genotoxic stress	
MultiFlow DNA Damage Kit – p53, γH2AX, Phospho-Histone H3	DNA damage responseGenotoxic Mode of Action	
MultiFlow DNA Damage Kit – Cleaved PARP, γH2AX, Phospho- Histone H3	DNA damage responseGenotoxic Mode of Action	
MultiFlow DNA Damage Kit – p53, γH2AX, Phospho-Histone H3, Cleaved PARP	DNA damage responseGenotoxic Mode of ActionApoptosis	



MultiFlow® - Phospho-Histone H3 Kit

Multiple endpoints. One step.

The Method

In just one step:

reagents are added to cells...

outer membranes are lysed, nuclei are stained...

and fluorescent antibodies bind to nuclear epitopes.

MultiFlow add-and-read kits were developed from the ground up to be simple and efficient.

When microtiter plates are used to collect these multiplexed data, the amount of test compound necessary for analysis is dramatically reduced.

At this point nuclei are ready for flow cytometric analysis.

Endpoints

- Mitotic cells via Phospho-Histone H3
- Cell density
- Cell proliferation
- Cytotoxicity

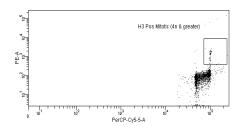
Benefits

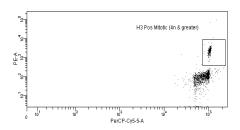
- High Information Content
- Multiplexed Assay format

- Simple and efficient
- Fast, flow cytometric analysis
- Scalable compatible with microtiter plates and robotic liquid handlers

Sample Plots

Compared to the negative control (left), four hours of exposure to vinblastine induced a large number of phospho-histone H3-positive events (right).







MULTIFLOW® - CLEAVED PARP KIT

Multiple endpoints. One step.

The Method

In just one step:

reagents are added to cells...

outer membranes are lysed, nuclei are stained...

and fluorescent antibodies bind to nuclear epitopes.

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When microtiter plates are used to collect these multiplexed data, the amount of test compound necessary for analysis is dramatically reduced.

At this point nuclei are ready for flow cytometric analysis.

Endpoints

- Apoptosis via cleaved PARP
- Cell density
- Cell proliferation
- Cytotoxicity

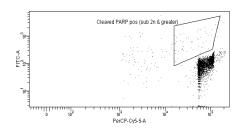
Benefits

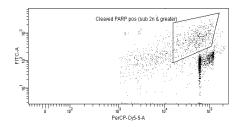
- High Information Content
- Multiplexed Assay format

- Simple and efficient
- Fast, flow cytometric analysis
- Scalable compatible with microtiter plates and robotic liquid handlers

Sample Plots

Compared to the negative control (left), 24 hours of continuous exposure to CCCP induces a large fluorescence shift (right).







MultiFlow® - γH2AX Kit

Multiple endpoints. One step.

The Method

In just one step:

reagents are added to cells...

outer membranes are lysed, nuclei are stained...

and fluorescent antibodies bind to nuclear epitopes.

MultiFlow add-and-read kits were developed from the ground up to be simple and efficient.

When microtiter plates are used to collect these multiplexed data, the amount of test compound necessary for analysis is dramatically reduced.

At this point nuclei are ready for flow cytometric analysis.

Endpoints

- Double strand DNA breaks via yH2AX
- Cell density
- Cell proliferation
- Cytotoxicity

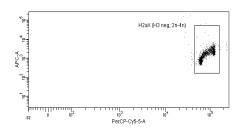
Benefits

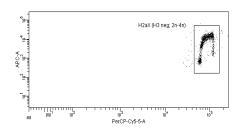
- High Information Content
- Multiplexed Assay format

- Simple and efficient
- Fast, flow cytometric analysis
- Scalable compatible with microtiter plates and robotic liquid handlers

Sample Plots

Compared to the negative control (left) 4 hours of exposure to camptothecin induced a large fluorescence shift (right).







MULTIFLOW® - P53 KIT

Multiple endpoints. One step.

The Method

In just one step:

reagents are added to cells...

outer membranes are lysed, nuclei are stained...

and fluorescent antibodies bind to nuclear epitopes.

MultiFlow add-and-read kits were developed from the ground up to be simple and efficient.

When microtiter plates are used to collect these multiplexed data, the amount of test compound necessary for analysis is dramatically reduced.

At this point nuclei are ready for flow cytometric analysis.

Endpoints

- Genotoxic stress via p53
- Cell density
- Cell proliferation
- Cytotoxicity

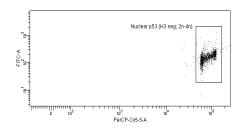
Benefits

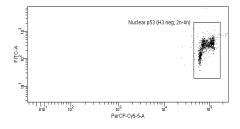
- High Information Content
- Multiplexed Assay format

- Simple and efficient
- Fast, flow cytometric analysis
- Scalable compatible with microtiter plates and robotic liquid handlers

Sample Plots

Compared to the negative control (left) 4 hours of exposure to camptothecin induced a large fluorescence shift (right).







MULTIFLOW® DNA DAMAGE KIT

P53, γH2AX, PHOSPHO-HISTONE H3

Multiple endpoints. One step.

The Method

In just one step:

reagents are added to cells...

outer membranes are lysed, nuclei are stained...

and fluorescent antibodies bind to nuclear epitopes.

MultiFlow add-and-read kits were developed from the ground up to be simple and efficient.

When microtiter plates are used to collect these multiplexed data, the amount of test compound necessary for analysis is dramatically reduced.

At this point nuclei are ready for flow cytometric analysis.

Endpoints

- Genotoxic stress via p53
- Double strand DNA breaks via yH2AX
- Mitotic cells via phospho-histone H3
- Cell density
- Cell Cytotoxicity proliferation

Benefits

- High Information Content
- Multiplexed Assay format

- Simple and efficient
- Fast, flow cytometric analysis
- Scalable compatible with microtiter plates and robotic liquid handlers



MULTIFLOW® DNA DAMAGE KIT

CLEAVED PARP, YH2AX, PHOSPHO-HISTONE H3

Multiple endpoints. One step.

The Method

In just one step:

reagents are added to cells...

outer membranes are lysed, nuclei are stained...

and fluorescent antibodies bind to nuclear epitopes.

MultiFlow add-and-read kits were developed from the ground up to be simple and efficient.

When microtiter plates are used to collect these multiplexed data, the amount of test compound necessary for analysis is dramatically reduced.

At this point nuclei are ready for flow cytometric analysis.

Endpoints

- Apoptosis via cleaved PARP
- Double strand DNA breaks via yH2AX
- Mitotic cells via phospho-histone H3
- Cell density
- Cell Cytotoxicity proliferation

Benefits

- High Information Content
- Multiplexed Assay format

- Simple and efficient
- Fast, flow cytometric analysis
- Scalable compatible with microtiter plates and robotic liquid handlers



MULTIFLOW® DNA DAMAGE KIT

P53, γH2AX, PHOSPHO-HISTONE H3, CLEAVED PARP

Multiple endpoints. One step.

The Method

In just one step:

reagents are added to cells...

outer membranes are lysed, nuclei are stained...

and fluorescent antibodies bind to nuclear epitopes.

MultiFlow add-and-read kits were developed from the ground up to be simple and efficient.

When microtiter plates are used to collect these multiplexed data, the amount of test compound necessary for analysis is dramatically reduced.

At this point nuclei are ready for flow cytometric analysis.

Endpoints

- Genotoxic stress via p53
- Double strand DNA breaks via γH2AX
- Mitotic cells via phospho-histone H3
- Apoptosis via cleaved PARP
- Cell Cytotoxicity density and cell proliferation

Benefits

- High Information Content
- Multiplexed Assay format

- Simple and efficient
- Fast, flow cytometric analysis
- Scalable compatible with microtiter plates and robotic liquid handlers

