**Multiphoton Flow Cytometer for High Throughput Analysis of Multicellular Aggregates Like Pancreatic Islets**

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The Wisconsin Alumni Research Foundation (WARF) is seeking commercial partners interested in developing a multiphoton flow cytometer that combines microfluidics with multiphoton microscopy to provide high throughput analysis of the cellular components of multicellular entities (i.e., pancreatic islets, embryoid bodies or model organisms). The instrument is capable of detecting fluorescence signals, including very weak autofluorescence associated with certain metabolites and extracellular matrix proteins.

**OVERVIEW**

Regenerative medicine requires a practical, high throughput method of assessing the cellular components of multicellular entities before transplantation. Such a method would have many clinical and research applications. For example, no means currently is available to efficiently assess the functional potential of pancreatic islet cells prior to transplantation without disrupting the islet. Additionally, the characteristics of embryoid bodies used for stem cell research cannot be screened easily prior to dissociation of the bodies.

Flow cytometers are well-established research and clinical instruments that can provide automated, quantifiable and verifiable data on cellular phenotype, differentiation and metabolic state. However, multicellular aggregates such as islets can be disrupted under the forces generated in a typical flow cytometer. Also, conventional flow cytometry monitoring systems are not well-suited for the analysis of three-dimensional cellular structures.

Multiphoton laser-scanning microscopy (MPLSM) is a fluorescence imaging technique that allows imaging of living tissue up to a depth of 600 µm. This technique is useful for live cell imaging studies of large, dense structures such as an islet. However, MPLSM has not been used for high throughput analysis of cells.

**THE INVENTION**

UW-Madison researchers have developed a system that combines the high throughput characteristics of flow cytometry with the capabilities of MPLSM. This multiphoton flow cytometry system (MPFC) enables deep, high resolution images of large diameter cells.
and aggregates.

The multiphoton laser can excite intrinsic cellular fluorophores such as NADH, FAD and collagen, allowing both spectral and lifetime data to be acquired. This information then can be used to reveal information on cellular processes like metabolism, viability and the functional potential of cells, pancreatic islets, embryoid bodies and other entities.

APPLICATIONS

• High throughput characterization of the inner cells of multicellular entities
• Noninvasive labeling, characterization and high throughput sorting of cells prior to transplantation

KEY BENEFITS

• Combines imaging and fluorescence measurements to provide more useful analysis metrics
• Enables sophisticated automated cell sorting
• Controls flow rates and minimizes shear forces, preventing damage to multicellular aggregates

STAGE OF DEVELOPMENT

The researchers have made a functional device that is being actively used.

ADDITIONAL INFORMATION

Related Technologies

Publications


Tech Fields
Research Tools - Detection
Analytical Instrumentation - Microscopy

CONTACT INFORMATION

For current licensing status, please contact Jeanine Burmania at jeanine@warf.org or 608-960-9846.