

# High Content Imaging: Dissecting Complexity of Biology

" High content screening (HCS), which combines automated fluorescence microscopy with quantitative image analysis, allows the acquisition of unbiased multi-parametric data at the single cell level."

## **Capabilities and Highlights**

- · >10 years of experience, with ~100 accumulated projects, >30 types of applications
- · MOA study with correlation of morphological changes to compound activity
- · Simultaneous multiple-parameter measurement within single cell level
- · High content screen/analysis at automated and high-throughput
- · Flexibility and adaptability during development process at QC control

#### General Application

- · 2D/3D Cytotoxicity
- · Protein degradation
- · Receptor cluster formation
- · Receptor internalization
- Membrane-cytosol-nucleus
  translocation
- Mitochondria classification (membrane potential)
- · Autophagy
- · Cell cycle
- · Apoptosis
- · In-cell Western

#### CNS

- · Neuroinflammation
- · Alzheimer's disease (AD)
- · Parkinson's disease (PD)
- · Amyotrophic lateral sclerosis (ALS)
- Other Neurological disorders & Pathway related assay

#### Oncology

- · Angiogenesis and tumor formation
- · Phagocytosis
- · Foci formation

#### **Kidney Disease**

- · MDCK 3D cyst model
- · Renal fibrosis



#### **Metabolic Disease**

- · Adipogenesis
- · Cholesterol lipolysis

#### **Safety Assessment**

- Drug induced liver toxicity evaluation
- · Drug induced cholesterolosis
- $\cdot$  Drug induced steatosis
- $\cdot$  Drug induced phospholipidosis
- · Micronucleus
- · DNA damage



## **Contact Us**

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## **WuXi Biology**

### **Screening for Mitofusin Activators**

An imbalance in mitochondrial dynamics is thought to contribute to the progression of neurodegeneration, cancer, and cardiac diseases.



In Mfn2-null MEFs there is a low mitochondrial aspect ratio (AR), which indicates poorly fused mitochondria. This assay was used to screen mitofusin activators targeting Mfn2 (outer mitochondrial membrane GTPase, critical for mitochondrial fusion).

Leflunomide: immunosuppressive disease-modifying antirheumatic drug (DMARD).



Mitofusin Mfn2 activators restore mitochondrial aspect ratio (AR) in mouse embryonic fibroblasts (MEFs).

#### Case study: Neurite Outgrowth & Neuronal Differentiation

The cells neuronal differentiation could be profiled and quantified using HCS, including the neurite outgrowth evaluation, neuron-specific markers characterization, etc.

#### **Neurite Outgrowth**





**Cells were induced to neuron-like under NGF incubation.** The neurite outgrowth process could be measured by neurite profile and quantification using HCS.

#### **Neuronal Differentiation**



Cells were differentiated towards a **Dopaminergic** phenotype that express **Tyrosine Hydroxylase** (TH) with robust neurite outgrowth (TUJ1).The differentiated cells showed increased sensitivity to **6-OHDA** 



Cells were differentiated towards a **Cholinergic** phenotype that express Choline Acetyltransferase (ChAT) with robust neurite outgrowth( $\top UJ1$ ). The differentiated cells showed increased sensitivity to **Acrylamide**.