

# Product Catalog

iFlow™ Droplet & Single-Cell

Microfluidic Systems & Solutions



# iFlow™ Droplet System -Touchscreen Version



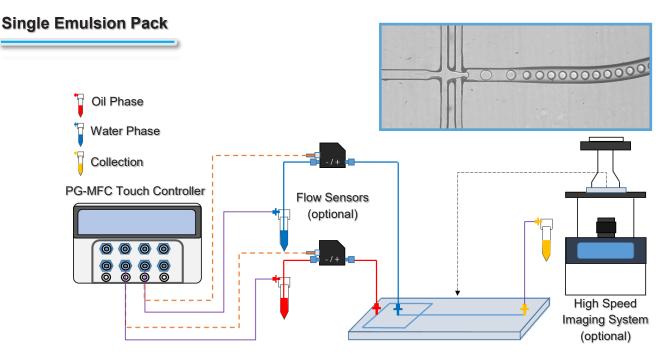


Microfluidic droplet generation allows greater control over size and uniformity than conventional batch production.

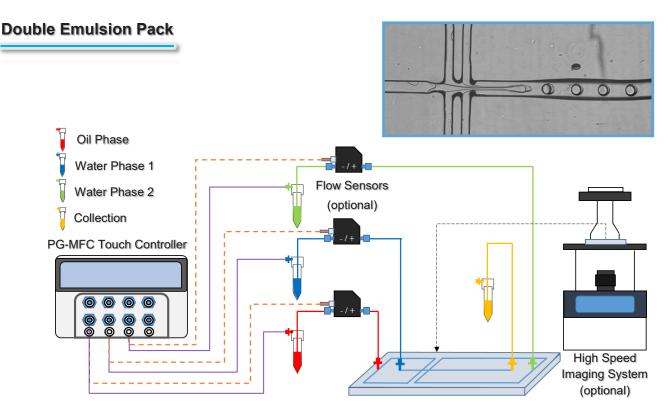
PreciGenome's droplet generation package provides a convenient open platform compatible with a wide range of chip designs. It may also be configured for either small volume production (<0.5 ml) or large volume (1.5 ml - 1 L). Single and double emulsion configurations are both available, and the iFlow Touch pressure controller provides a convenient AIO control and display unit.

# **Key Benefits High Performance** Scalable **Open Platform** & Efficiency **Throughput** • Direct plug reservoirs: <0.5ml • Tunable size (chip dependent) Reagents • Microfluidic chips • Reservoir kits: 1.5 ml - 1 L Uniform production • Poisson encapsulation **Simple Operation Cost Effective** Easy setup • Affordable configuration Compact size Low cost consumables Intuitive UI w/ touchscreen





In single emulsion, one dispersed phase forms uniform droplets within the continuous phase. This setup is for water-in-oil emulsion. Oil-in-water emulsion would switch the two phases' positions.



In double emulsion, an additional continuous phase encapsulates the initial droplets formed in the first emulsion event. This setup is for water-in-oil-in-water emulsion. Switching water phase 2 and the oil phase would result in a three-inlet single emulsion.

# iFlow™ Droplet System

# - Light Version

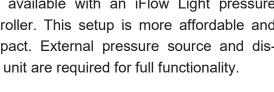






PreciGenome's droplet generation package is also available with an iFlow Light pressure controller. This setup is more affordable and compact. External pressure source and display unit are required for full functionality.

## Diagram for Single Emulsion



## **System Contents**

PG-MFC-LT-2CH light pressure controller

2x reservoir kits (1.5, 15, 50, 100, or 1000 ml)

2x reservoir tube racks (15 or 50 ml)

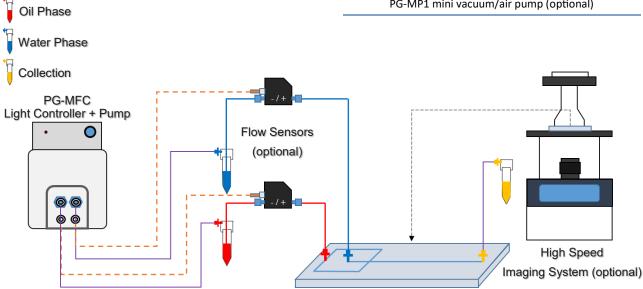
2x droplet generation chips (assorted models)

PG-LFS flow sensor (assorted full scale, optional)

**Tubings & fittings** 

PG-HSV high speed imaging system (optional)

PG-MP1 mini vacuum/air pump (optional)

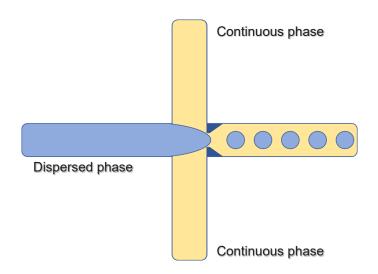


Though the pressure source is different for this package, the principles of droplet generation are the same regardless of pressure controller used. With the right chip and setup, the iFlow Light may be configured for single emulsion shown above, or double emulsion (not shown.)



# Working Principle

The system requires two or more unmixable liquid phases, referred to as the dispersed phase, sometimes called the droplet phase, and the continuous phase. The schematic below illustrates the device with junction in focused-flow geometry designed for droplet generation.



Schematic illustration of enlarged junction, controlled break-up droplet at the orifice. The flow through the orifice enables a controlled droplets break-up, which is required for yielding monodisperse micro-emulsions.

# Microfluidic Droplet Generator Chips

PreciGenome offers a variety of droplet generator chips in different materials to meet most of our customers' application requirements.

Two types of materials, including polymers (PC, COC, COP etc.) and glass, are commonly used to fabricate microfluidic chips. Material of the chip is selected depending on the application requirements, including chip design, types of solvent or reagent used for experiment, needs of the application, budget, and fabrication time etc. Usually, for research purposes the materials for the chip fabrication generally prioritize performance of the device. For mass production of a products, the factors of production cost, reliability and ease of use are considered first.



# **Applications**



- Droplet-based PCR
- DNA/RNA sequencing
- Drug delivery
- Protein crystallization

- Cell culture
- Chemical synthesis
- Microparticle synthesis
- Gel particle synthesis

#### **Droplet-based PCR**

Digital PCR system as a new generations of Polymerase Chain Reaction (PCR) system has been an important tool in genomics and biological fields. Droplet PCR operates by assembling ingredients, forming droplets, combining droplets, thermocycling, and then processing results by using water-in-oil systems.

#### **DNA/RNA** sequencing

Droplet-based microfluidic systems have been used for DNA/RNA sequencing

#### **Drug Delivery**

Droplet microfluidics enables useful platforms for drug delivery vehicles and drug molecules as novel functional materials. Because of uniform size, monodisperse size distribution, and desired properties, droplet microfluidics demonstrates promising potential for production of complex drug systems.

#### **Protein crystallization**

Droplet microfluidics technology has been used for investigating the conditions necessary for protein crystallization.

#### Cell culture

Droplets are able to be used as incubators for single cells. Due to the high throughput, incubation in up to millions of droplets offers powerful capacity of characterizing cell population based on cells' kinetic behavior such as protein secretion, enzyme activity, etc.

#### **Chemical synthesis**

Droplet-based microfluidics has become an important method for chemical synthesis. Droplets are able to act as individual reactions, which are free from contamination from outside.

#### Microparticle synthesis

Advanced particles and particle-based materials, such as polymer particles, microcapsules, nanocrystals, and photonic crystal beads can be synthesized by the droplet generation system. The system is also able to synthesize microparticles/nanoparticles, like PLGA microparticles, colloidal CdS and CdS/CdSe core-shell nanoparticles.

### **Gel Particle Synthesis**

In the last decade, the gel particles (hydrogels, microgels, and nanogels) has been an area of interest for many researchers and industries. Because of high throughput, mono-dispersity of particles, and low cost, droplet-based microfluidic systems have been widely used.

# **Droplet Size & Frequency Tuning**



76.1 µm 858 hz

64.1 µm 1169 hz

55.5 μm 2200 hz

With its user-friendly interface and tunable pneumatic pumps, the iFlow pressure controller allows fine control over pressure ratios to achieve a wide range of droplet sizes and production rates with high consistency between different scales of production.

Selected data is for microfluidic chips with 38 µm nozzle width in a single emulsion configuration (three-inlet is capable of double emulsion, not shown.)

Water 2 psi Oil 2 psi	0	75.3 µm 950 hz	Water 2 psi Oil 2 psi	0
Water 3 psi Oil 4 psi	0	65.7 µm 1488 hz	Water 3 psi Oil 4 psi	0
Water 4 psi Oil 6 psi	0	58.8 µm 2078 hz	Water 4 psi Oil 6 psi	0

### Two-Inlet Chip (15 ml Reservoir Kits)

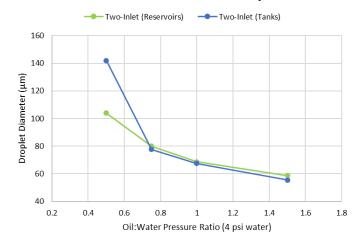
Water 2 psi Oil 3 psi	0	86.5 µm 465 hz	Water 2 psi Oil 3 psi	0	87.4 µm 651 hz
Water 3 psi Oil 6 psi	0	73.5 µm 1488 hz	Water 3 psi Oil 6 psi	0	73.5 µm 2425 hz
Water 4 psi Oil 9 psi	0	63.9 µm 2976 hz	Water 4 psi Oil 9 psi	0	51.3 µm 9954 hz

Three-Inlet Chip (15 ml Reservoir Kits)

## Three-Inlet Chip (500 µl Chip Tanks)

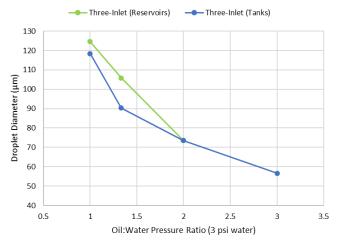
Two-Inlet Chip (500 µl Chip Tanks)





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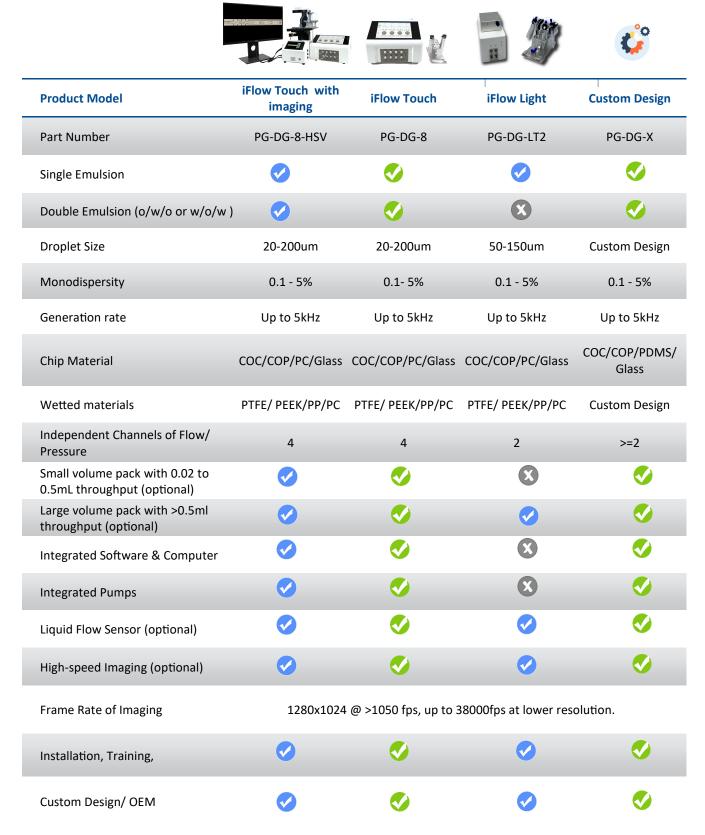
## Oil:Water Pressure Ratio vs. Droplet Size



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# iFlow™ Droplet System Comparison Chart

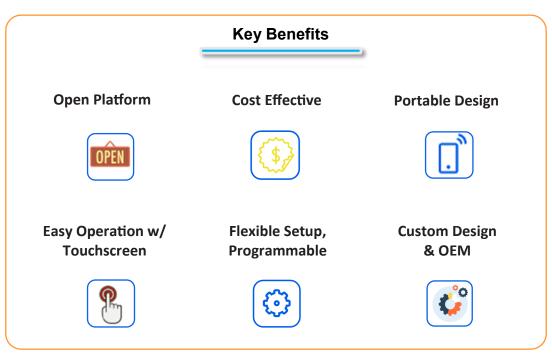


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# iFlow™ Single Cell Analysis R&D System





## **Applications**

- Single cell isolation & PCR amplification
- Single cell antibody discovery
- Single cell analysis
- Cell line development
- DNA/RNA sequencing library

- Biopharmaceutical discovery
- Drug-resistance studies
- Double emulsion
- Enzyme evolution
- Synthetic biology

Catalog #	Name		
PG-SC-8-HSV	System w. iFlow Touch™ controller and highspeed imaging		
PG-SC-8	System w. iFlow Touch™ controller		

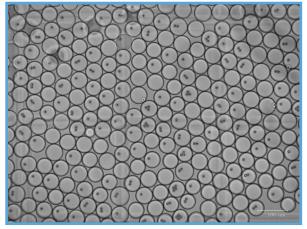


# Single Cell Encapsulation Efficiency

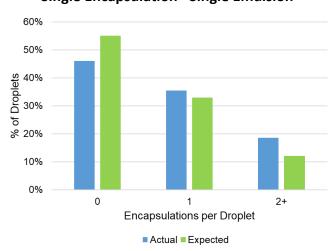
## **Encapsulation vs. Poisson Distribution**

Using 10 µm polystyrene beads as a model for mammal cells, droplet encapsulation follows the Poisson distribution. *Data shown is for 55-60 µm droplets; expected value is with one-third of initial bead concentration (chip design dependent.)* 

Single Encapsulation - 10  $\mu m$  beads



Single Encapsulation - Single Emulsion



## Beads Concentration vs. Encapsulation Rates

The Poisson distribution also holds for other bead concentrations. As bead concentration decreases, a greater percentage of droplets have no or one encapsulation events. *Data shown is for 55-60 µm droplets; expected value is one-third of initial bead concentration (chip design dependent.)* 

Initial Bead Concentration	% 0 Encapsulations		% 1 Encapsulations		% 2+ Encapsulations	
(beads/ml)	Actual	Expected	Actual	Expected	Actual	Expected
7,470,000	80.3	75.5	18.1	21.2	1.6	3.3
12,900,000	69.4	68.8	24.5	25.7	6.1	5.5
20,580,000	46.1	55.0	35.4	32.9	18.5	12.1

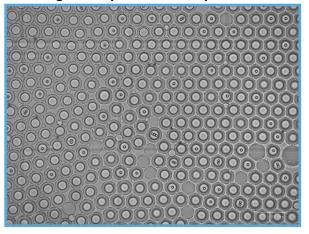


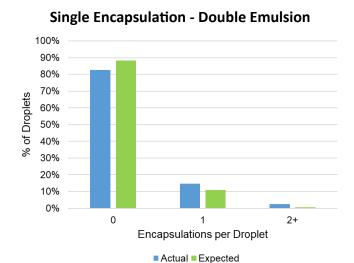
# Single Cell Encapsulation Efficiency

## **Double Emulsion Encapsulation**

Droplet encapsulation also follows the Poisson distribution in double emulsion systems. Data shown is for 30 µm inner shell; expected value is with one-half of initial bead concentration (chip design dependent.)

Single Encapsulation - 10 µm beads





**Initial Bead Concentration** % 0 Encapsulations % 1 Encapsulations % 2+ Encapsulations (beads/ml) Actual **Expected** Actual **Expected** Actual **Expected** 17,610,000 82.6 88.3 14.8 11.0 2.6 0.7



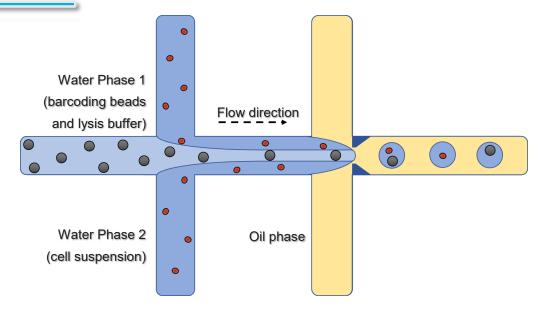
Surfactants and/or stabilizing agents are needed to maintain stable double emulsions, and pressure tuning can adjust droplet size. Factors such as presence of cells or inorganic salts may affect surfactant choice. The following surfactant mixes were used for each phase:

Phase	Solvent	Surfactant		
I	DI H <sub>2</sub> O OR	0.5-1% TWEEN-20 OR		
Inner Water	1x PBS	0.5-2% BSA		
Oil	HFE 7500	2.2% Krytox 157		
Outer Water	DI H <sub>2</sub> O OR	1-3% Triton X-100		
	1x PBS	1-370 1111011 A-100		



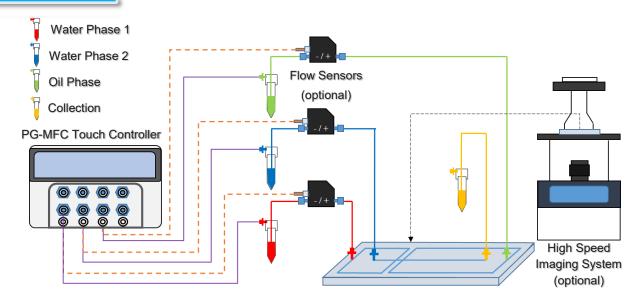
# Application: Drop-Seq

## **Working Principle**



In Drop-Seq, the microfluidic chip allows two water phases to mix before being dispersed in the oil phase. This allows barcoding beads and cells to be captured together in droplets.

## System Diagram



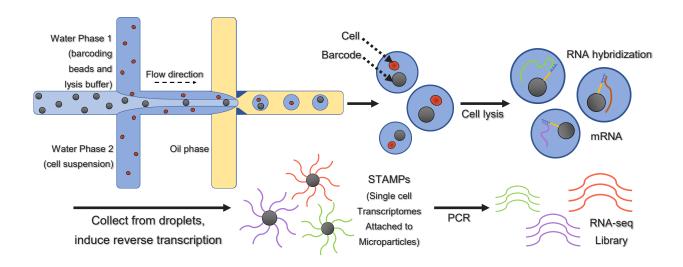
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# **Drop-Seq Encapsulation Efficiency**

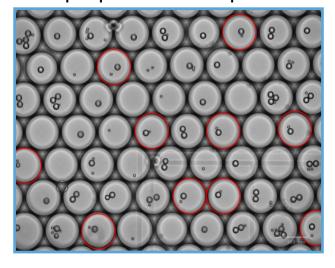
## **Drop-Seq Introduction**

An expansion of single cell encapsulation, Drop-Seq uses droplets to pair single cells with single barcoding microparticles. The resulting droplets serve as miniature independent reaction chambers for cell lysis and RNA hybridization, after which the products may be collected and sequenced to generate a library of thousands of single-cell transcriptomes.



## **Drop-Seq Encapsulation**

Drop-Seq Model - 10 and 20 µm beads



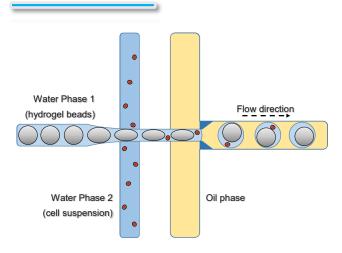
Droplet encapsulation similarly follows the Poisson distribution for a Drop-Seq model. Encapsulation was done with 10 and 20  $\mu$ m polystyrene beads in 100  $\mu$ m droplets, where 10  $\mu$ m beads modeled cells and 20  $\mu$ m beads modeled barcoding microparticles. Approx. 14.4% of droplets contained one of each bead (vs. 13.5% modeled by the Poisson distribution.)

# PreciGenome

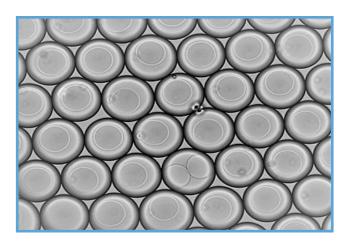
# Application:

# Hydrogel Beads Encapsulation

## Working Principle

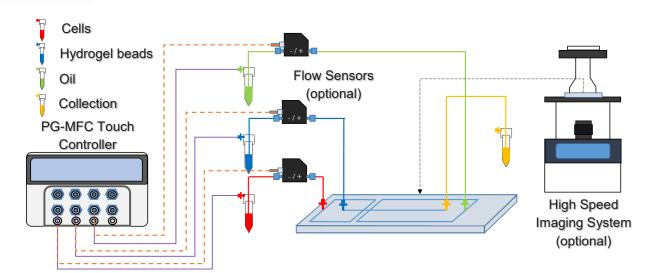


With specially designed microfluidic channels in the same schematic microfluidic chip as in the previous Drop-Seq example, hydrogel beads can reach 90% hydrogel encapsulation rate on our single cell analysis R&D platform.



By attaching functional groups, hydrogel beads have diverse applications, such as cell culture, drug delivery study, etc. Combining mono-dispersed hydrogel beads with different attachments and microfluidic droplet technology allows a high encapsulation rate of single cell and single bead. Suitable droplets can thus be miniature independent reaction chambers. Researchers can study individual cell behaviors with different environments. By adding barcodes on hydrogel beads, single cell RNA seq can be achieved. Researchers are able to study rare cancer cell mutation, response to new drugs, and CRISPR screening, etc. Using this platform, cell isolation and sorting can be achieved as well.

## **System Diagram**



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# Notes



## **Some of Our Customers**





















PreciGenome is located in the heart of Silicon Valley, San Jose, California, USA. We have been focusing on developing nanoparticle synthesis systems and solutions for our customers since we started our business. Our technology enables rapid prototyping with high quality and reliable performance for lipid nanoparticles, liposomes, PLGA, etc.

#### **HEADQUARTER**

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