

# Particle Size Analysis for Homogenization Process Development

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HORIBA Seminar

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# Outline

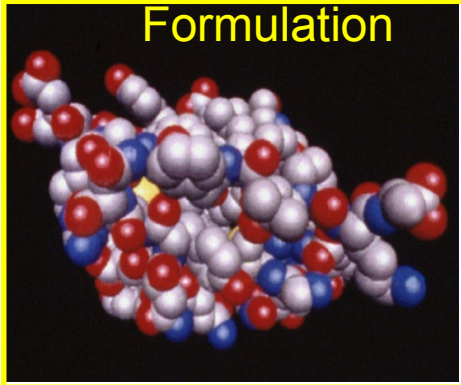
- Background
- Study Objective
- Homogenization Equipment Operating Principles
- Emulsion Preparation Process
- Instrument for Particle Size Evaluation
- Laboratory Study Results
- Summary

# Micro- and Nano- Particulate Systems

- Micro- and nano- suspensions and emulsions are commonly used in pharmaceutical, chemical, and consumer products.
- The control of the size and its distribution of these small-scale materials is important to their stability and performance during applications.
- Today, I am going to give you an example of how we developed a homogenization process for manufacturing pharmaceutical emulsions used in a pulmonary drug delivery system.

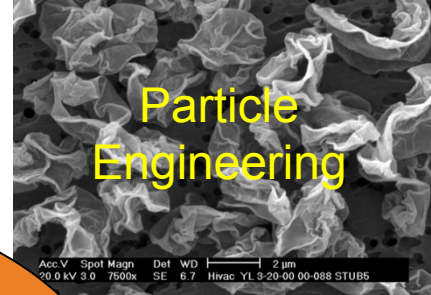
# Pulmonary Drug Delivery Product Life Cycle

## Formulation



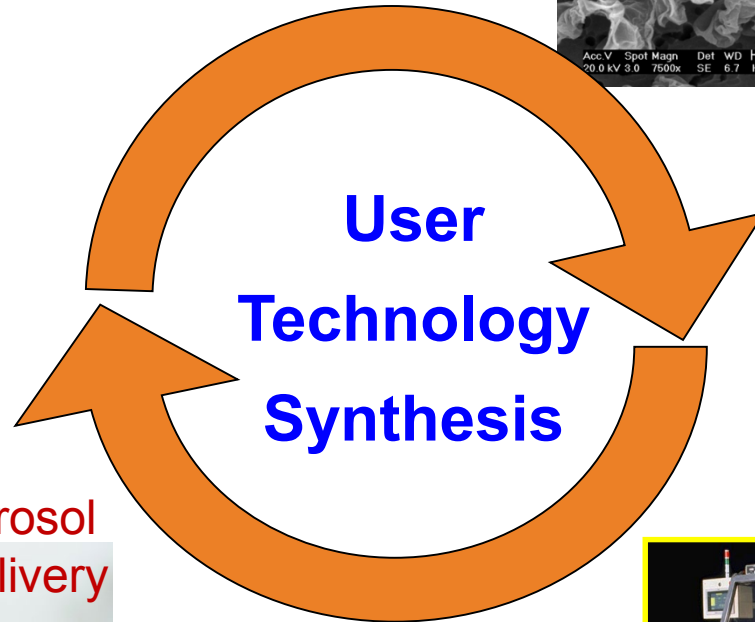
Dose  
Chemical/Physical Stability  
Solid-state properties  
Excipients

## Particle Engineering



Aerodynamics  
Dispersability  
Process control  
Scale up

Patient's needs  
Reproducible  
Easy to use & reliable



## Powder Processing



## Aerosol Delivery



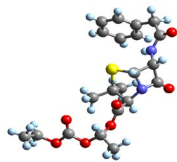
Fine powders  
Low mass, tight RSD  
Remain dispersable  
Billions/year

## Filling & Packaging



# Methods to Produce Fine Particles for Pharmaceutical Applications

**Molecule**

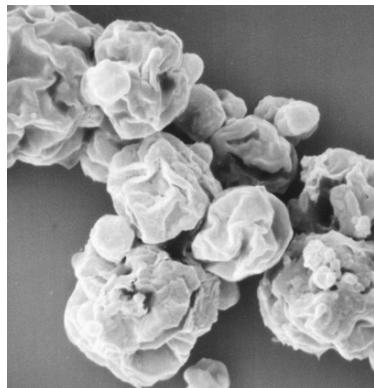


**Spray Drying**



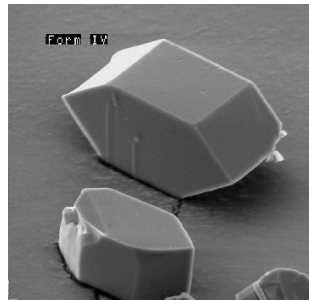
**Emulsion Based**

**SCF**



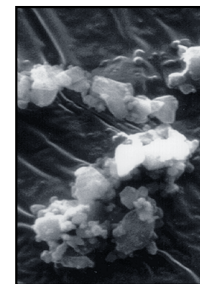
**Solution Based**

**Lyophilization**



**Crystallization**

**Attrition / Jet Milling**



**Fine Particles  
(MMAD ~ 1-5  $\mu\text{m}$ )**

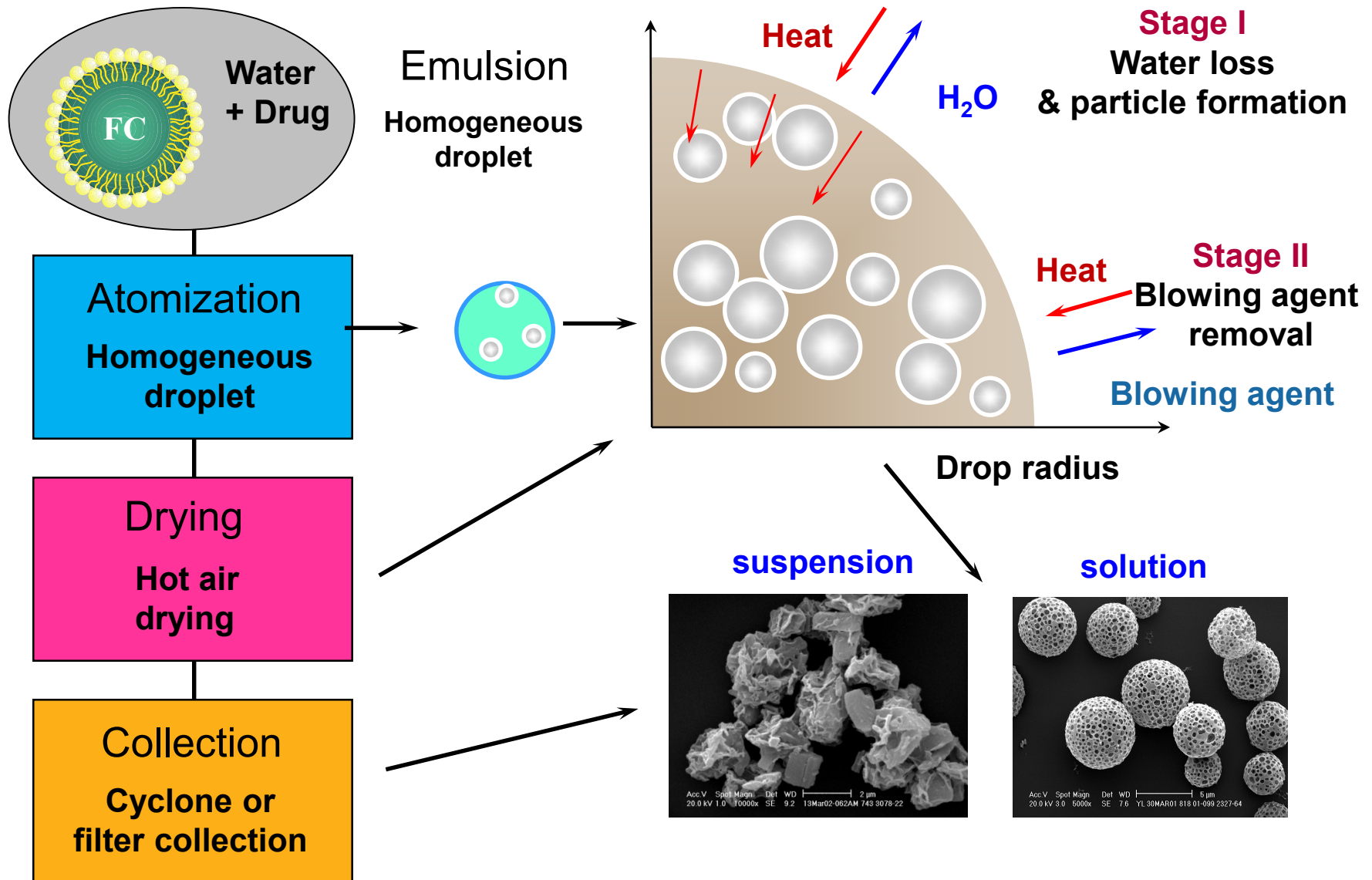
**PulmoSphere®  
Technology**

# PulmoSphere Particle Characteristics

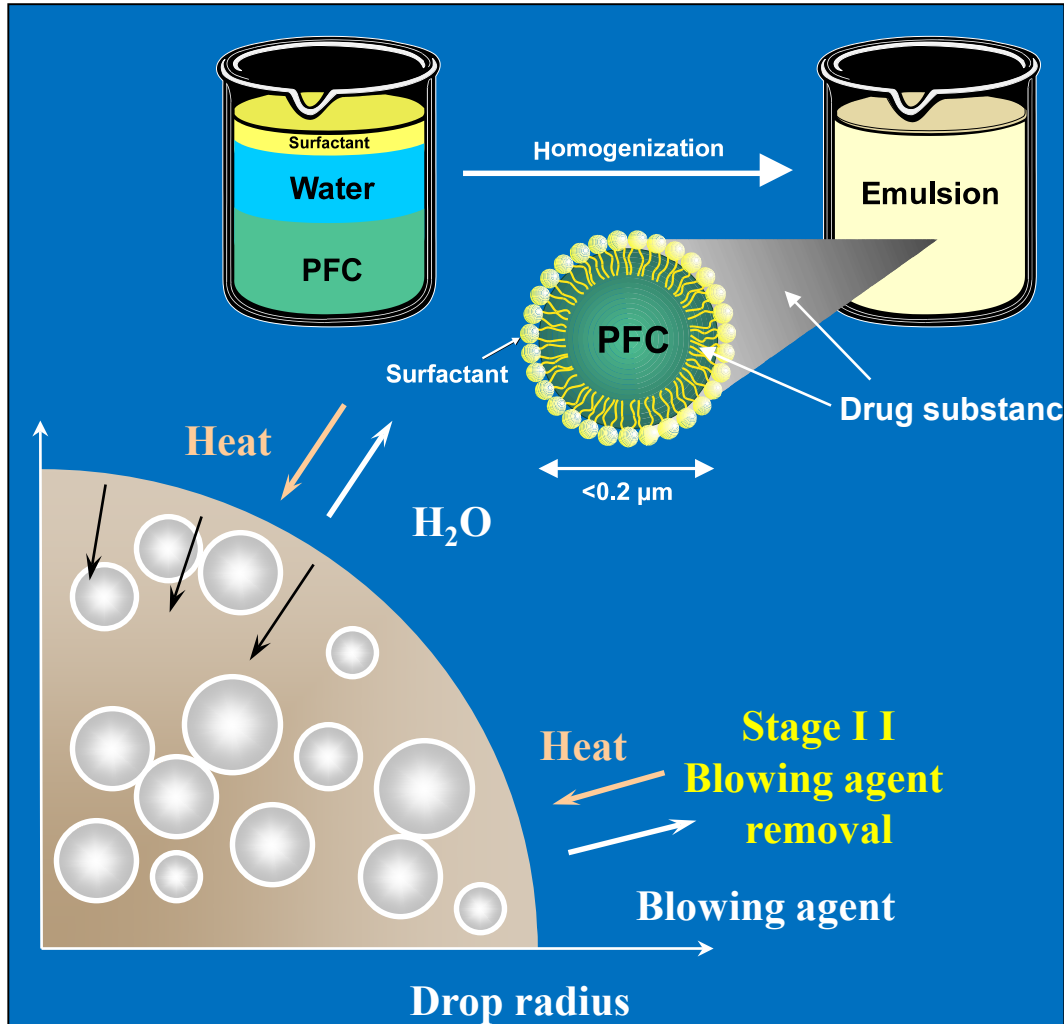
- Particle Physical Properties
  - Hollow and porous
  - Surface roughness
  - Low density
- Particle Performance Attributes
  - Flowability
  - Dispersibility
  - Aerodynamic



# Manufacture of *PulmoSphere* Particles



# Novartis PulmoSphere Technology



- Manufacture of a Fluorocarbon-in-Water Emulsion
- Excipients primarily composed of phospholipids
  - Perfluorocarbons added as a processing aid
  - Removed in the process



# Particle Size Analysis for Homogenization Process

- ◆ **The objective of this study is to develop a robust homogenization process for making pharmaceutical emulsions by evaluating droplet size distribution**

*Homogenization is a fluid mechanical process that involves the subdivision of droplets or particles into nanometer or micron sizes to create a stable emulsion or dispersion for further processing. This technology is one of most efficient means for size reduction.*

# Criteria for Evaluating High-Pressure Homogenizers

- Mean Particle Size
- Particle Size Distribution
- Emulsion Stability
- Cycle Time
- CIP/SIP Capability
- Scale Up Capability
- Routine Operation
- Maintenance

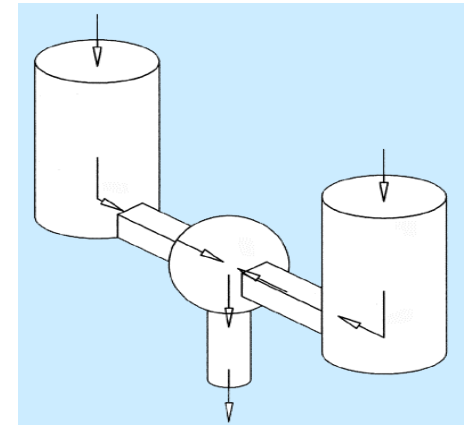
# Homogenization Principles

- High-pressure processing equipment for reducing droplet or suspension particle size primarily involves four mechanisms:
  - Shear - is caused by elongation and subsequent breakup of droplets, due to acceleration of a liquid
  - Turbulence - is caused by high velocity fluid resulting in chaotic motion to tear apart the globules
  - Impact - is caused by impinging of pressurized fluid on a hard surface breaking globules into smaller droplets
  - Cavitation - is caused by an intense pressure drop, leading to formation of vapor bubbles in the liquid, which implode causing shock waves in the fluid

*Homogenizers, available from different manufacturers operate based on combination of these mechanical forces*

# Equipment for Evaluation - Microfluidics

- Microfluidics: M-110EH
  - Microfluidics combines high flow with high-pressure, scalable fixed-geometry interaction chambers that impart high shear rates to product formulations
  - The entire product experiences identical processing conditions, producing the desired results, including: uniform particle and efficient droplet size reduction



# Process and Product Parameters

## ■ Processes

- Configuration (gap, length, shape, and size)
- Pressure drop
- Residence time
- Cooling efficiency (temperature control)

## ■ Product

- Concentration
- Viscosity (ratio and individual)
- Interfacial tension (surfactant amount and adsorption rate)
- Coalescence rate (Gibbs elasticity)
- Temperature sensitivity

# Emulsion Preparation Process

- Aqueous phase prepared with lipid surfactant
- Addition of modifier
- Pre-mix: Oil phase is slowly added to aqueous phase while mixing with a high-speed rotor/stator mixer
- High pressure homogenization

# Process Parameters Evaluated

- Pressure Drop
- Configuration
- Number of Passes
- Temperature Control

# Desirable Quality Attributes

- Mean Particle Size
  - Less than 0.8 micron (fine emulsion)
- Polydispersity
  - RSD Less than 10% (narrow distribution)
- Emulsion Stability
  - Less than 10-20% change in particle size or particle size distribution over extended period of time (long hold time)



# Instrumentation Used For Evaluation

- *Instrument for determining emulsion size*
  - Photo sedimentation – CPS
  - Dynamic light scattering – Malvern Zetasizer
  - Static light scattering – HORIBA LA-950
- *Criteria for choosing particle size analyzer*
  - Wide dynamic range
  - Broad applications
  - Accuracy and precision
  - Short cycle times (sample prep, measurement, and cleaning)
  - Ease of operation and maintenance
  - Regulatory compliance
  - At-line/on-line application

# Instrument for Particle Size Analysis

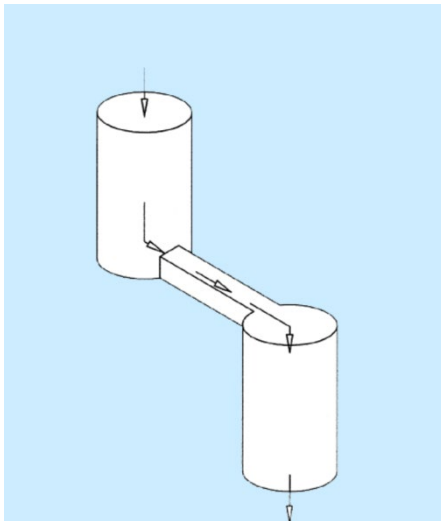
- Emulsion Sizing
  - HORIBA LA-950
  - Laser light scattering technique
  - Mie theory
  - 0.01 – 3000 micron
  - Good reproducibility
  - Fill, auto-alignment, blank, measurement, and rinse in less than 60 seconds



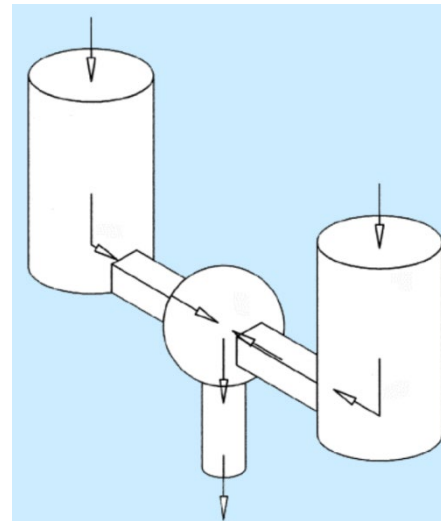
# Microfluidics Study

- Chambers                      G10Z and F20Y
- Pressures                      15 and 25 kpsig
- Passes                          1-5

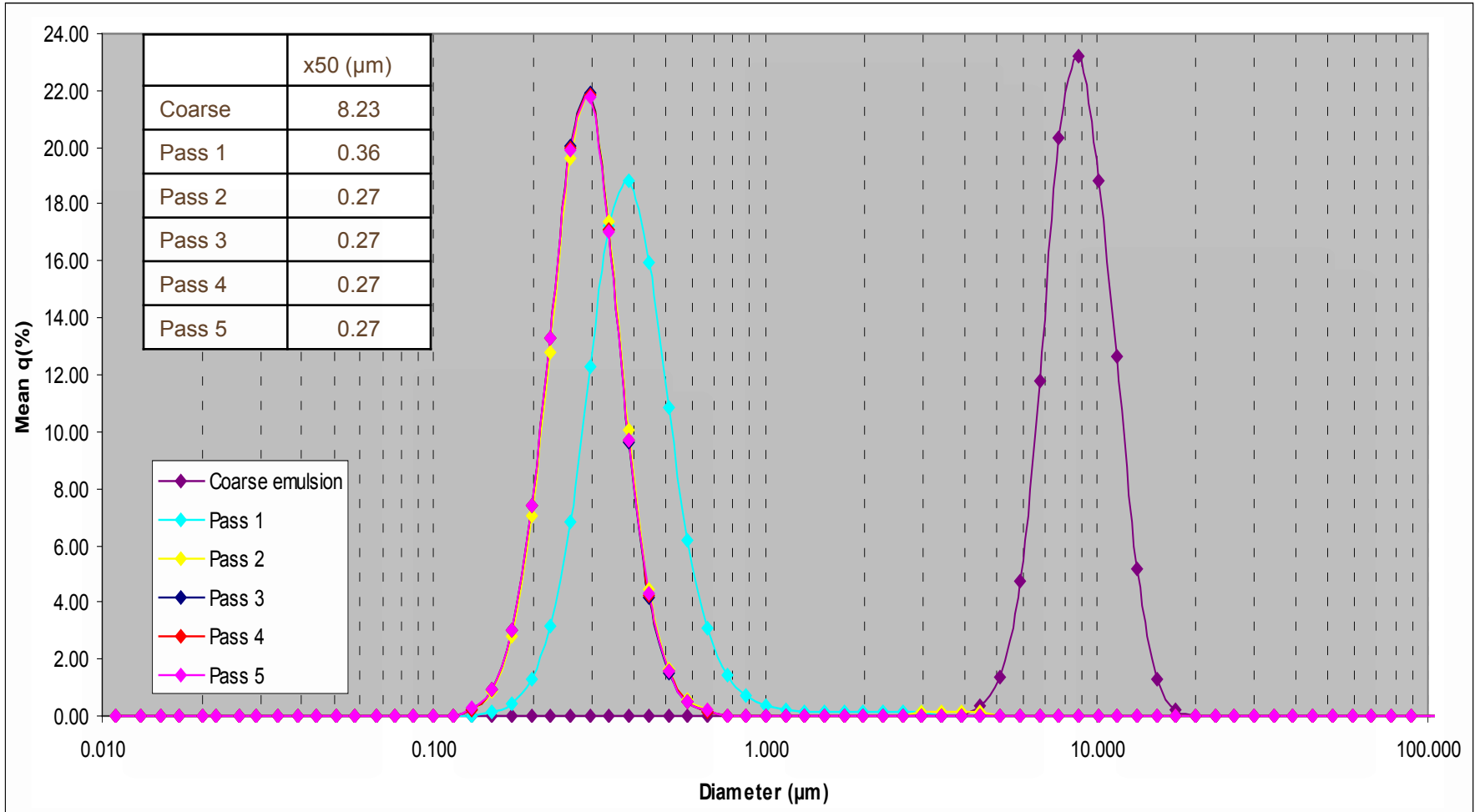
Z - Type



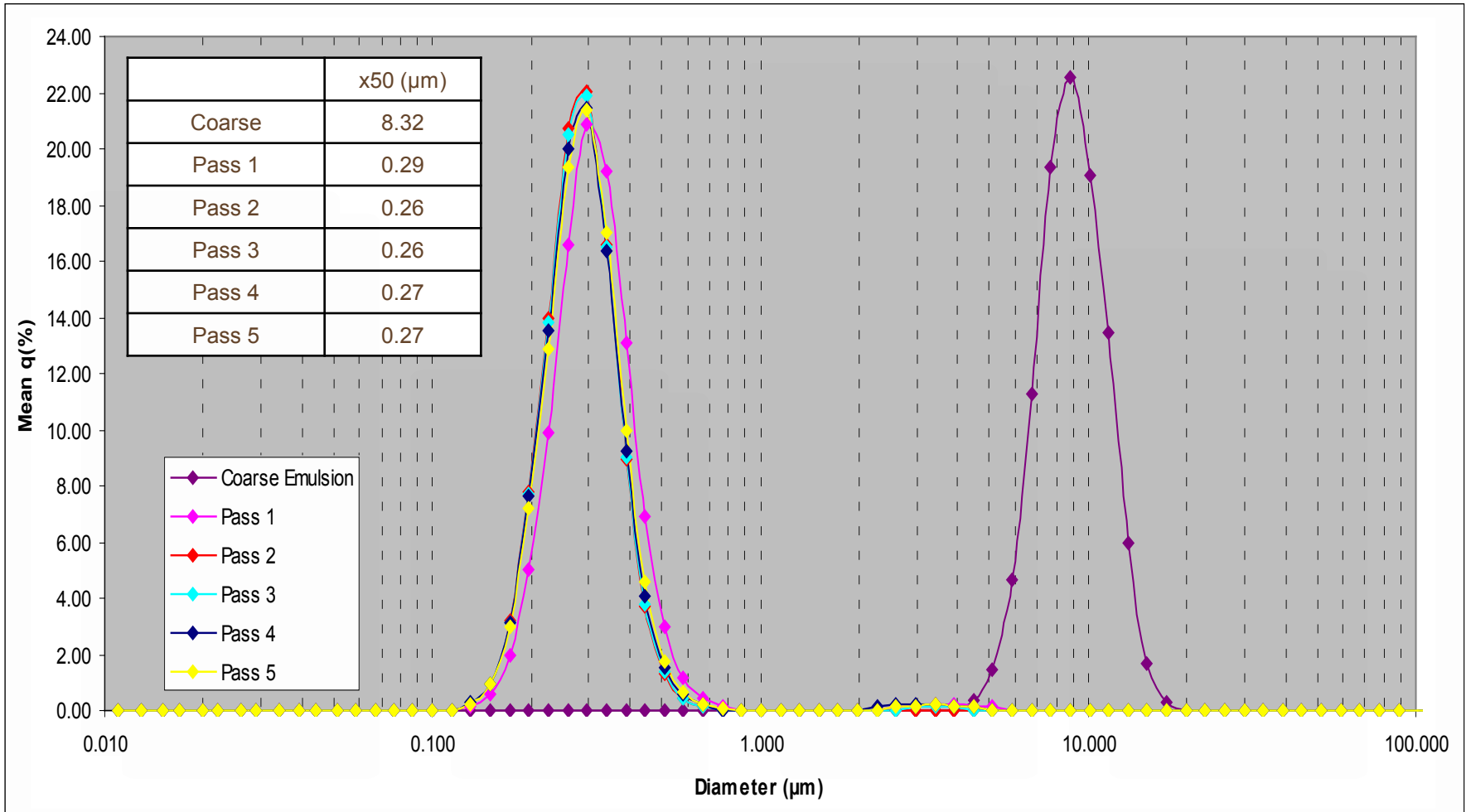
Y - Type



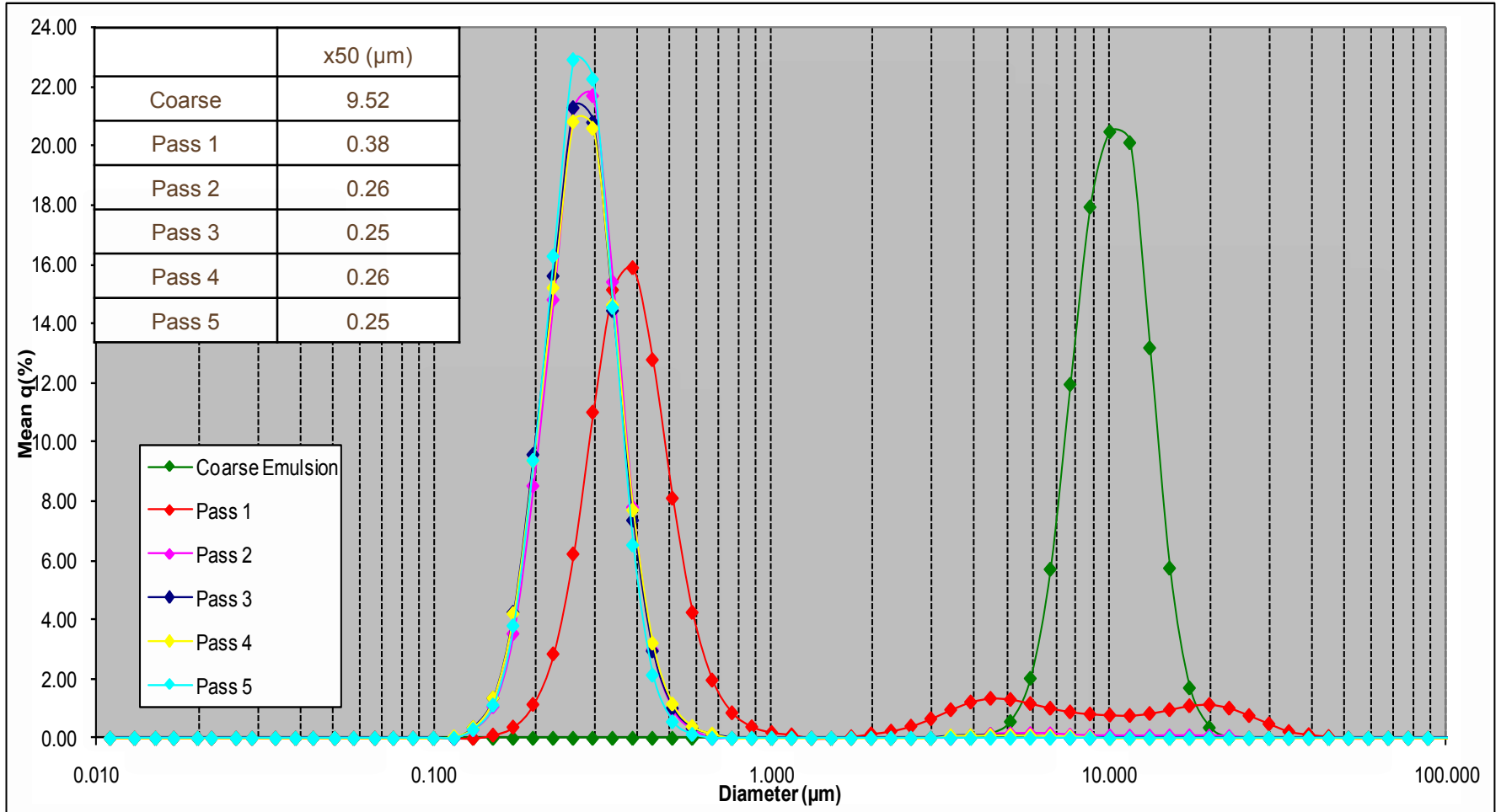
# Microfluidics - G10Z at 15 kpsig



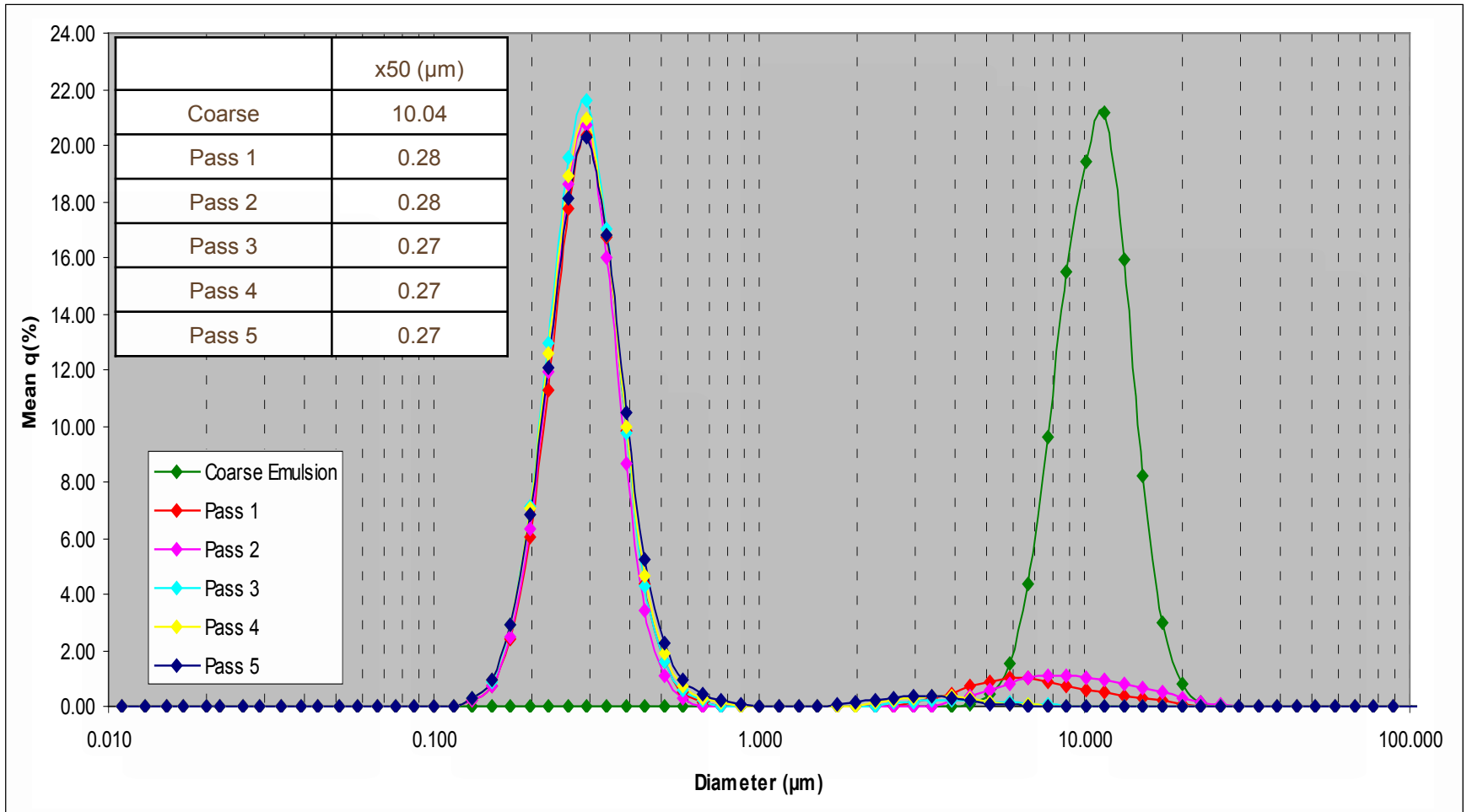
# Microfluidics - G10Z at 25 kpsig



# Microfluidics - F20Y at 15 kpsig



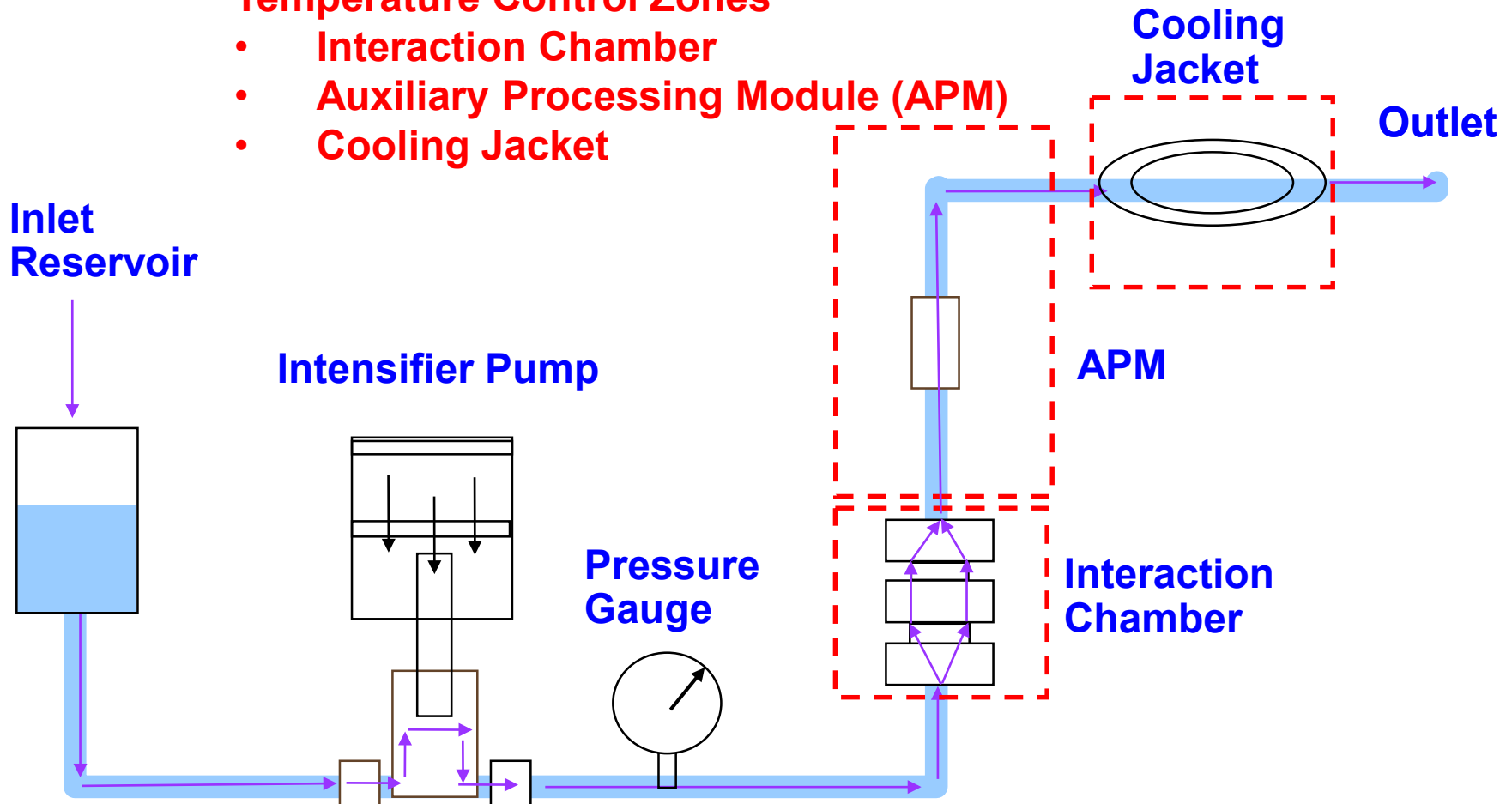
# Microfluidics - F20Y at 25 kpsig



# Microfluidics Study – Effects of Temperature Control

## Temperature Control Zones

- Interaction Chamber
- Auxiliary Processing Module (APM)
- Cooling Jacket





# Microfluidics Study – Effects of Temperature Control

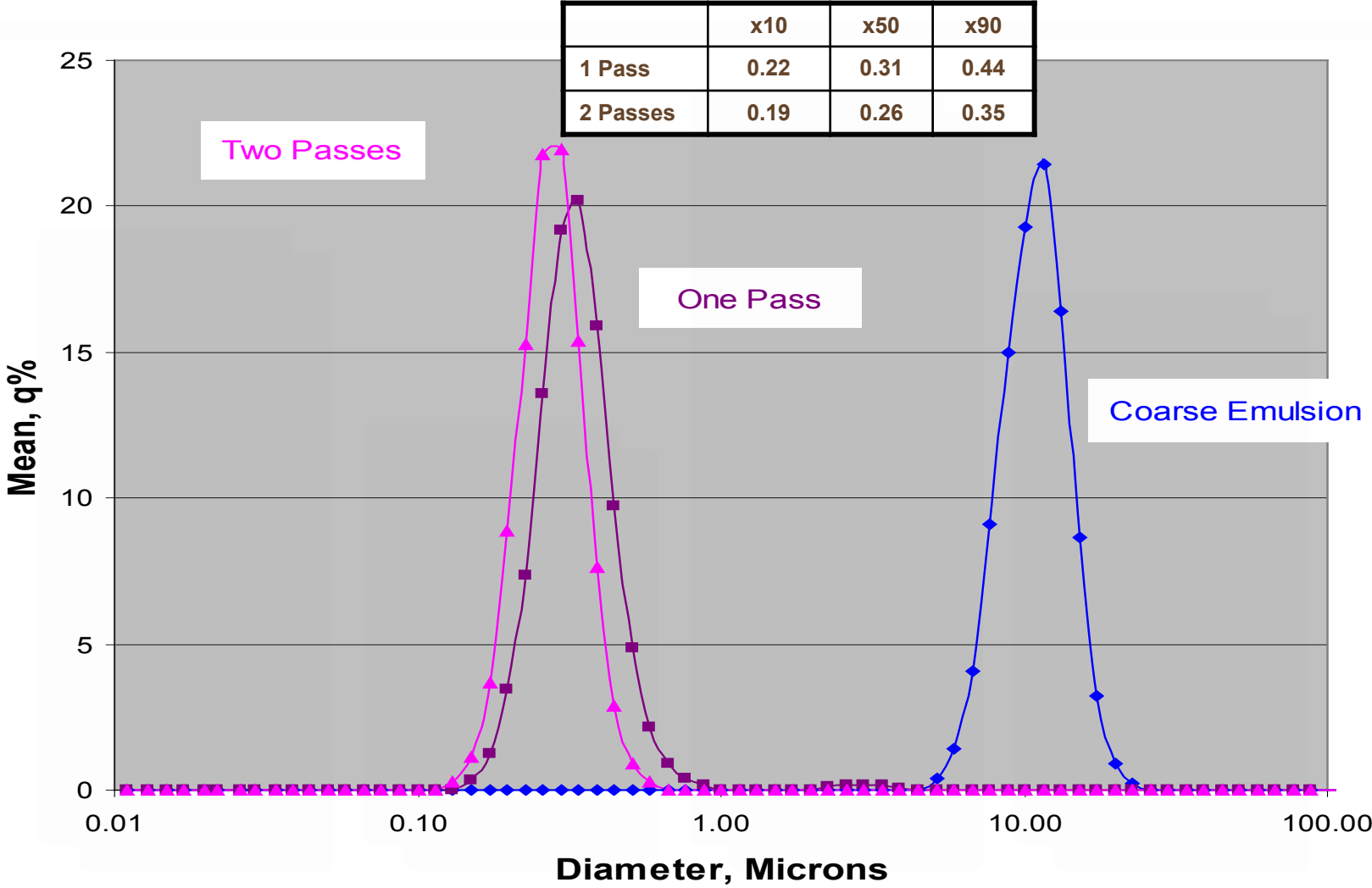
-	Interaction Chamber	APM	Cooling Jacket
Interaction Chamber	-	+	+
APM		++	+++
Cooling Jacket			+

**Interaction Chamber + Exit Line + Cooling Jacket = ++**

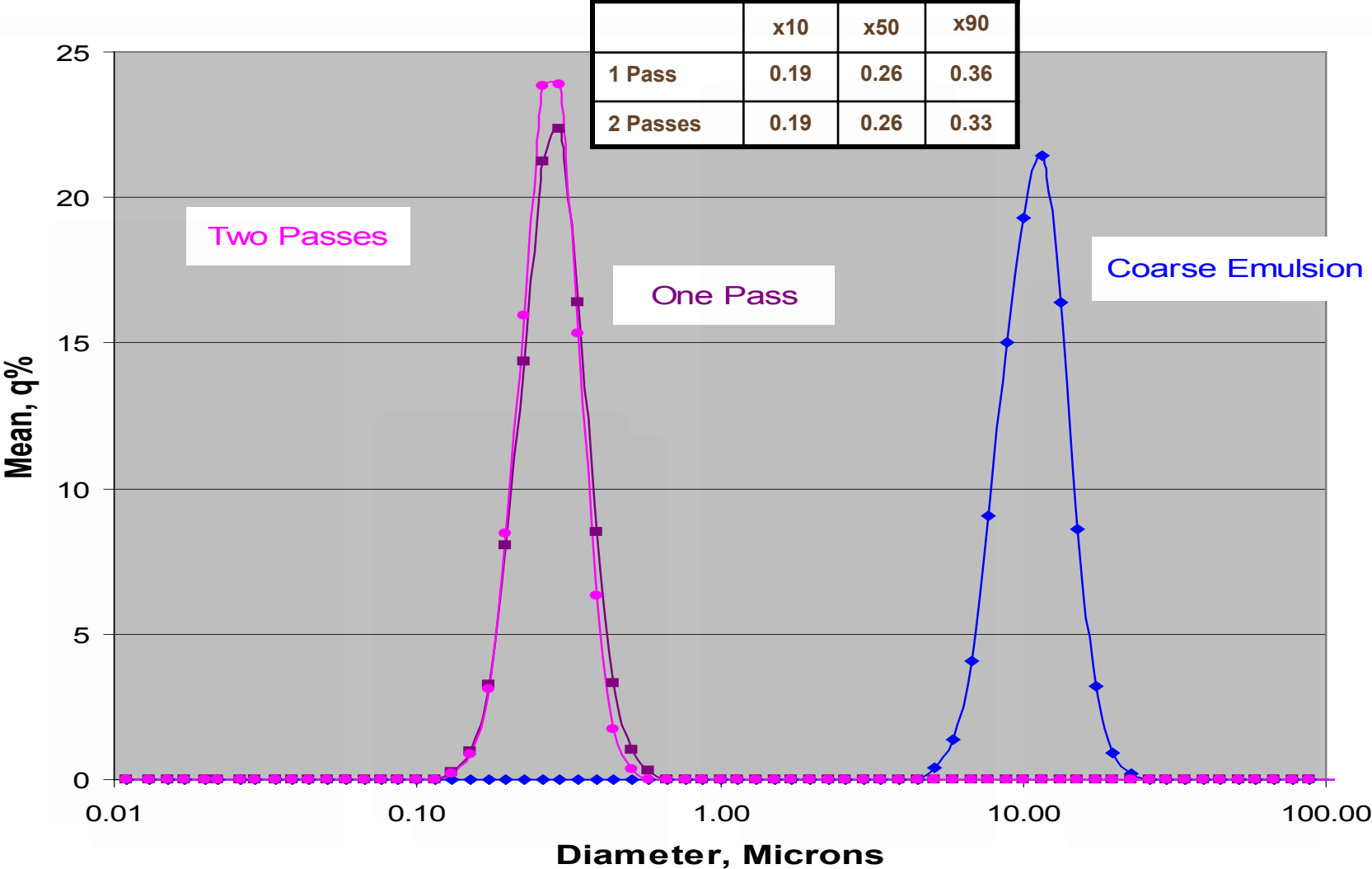
# Microfluidics Study – Temperature Control of APM and Cooling Jacket

Condition	Chamber	Pressure kpsig	Particle Size, Micron					
			Pass 1			Pass 2		
			x10	x50	x90	x10	x50	x90
1	G10Z	15	0.26	0.39	0.62	0.19	0.26	0.35
2	G10Z	20	0.21	0.29	0.40	0.19	0.25	0.33
3	G10Z	25	0.20	0.27	0.37	0.19	0.26	0.34
4	F12Y	15	0.22	0.31	0.44	0.19	0.26	0.35
5	F12Y	20	0.19	0.27	0.37	0.19	0.26	0.34
6	F12Y	25	0.19	0.26	0.36	0.19	0.26	0.33

# Microfluidics – F12Y at 15 kpsig with Optimized Cooling



# Microfluidics – F12Y at 25 kpsig with Optimized Cooling



# Emulsion Stability Studies

Homogenizer	Sample	Time Point		x10	x50	x90	Mean	Mode
Avestin C-50	Homogenized Emulsion Pass #3	0	Ave	0.18	0.26	0.38	0.35	0.25
Avestin C-50	Homogenized Emulsion Pass #3	24hrs	Ave	0.19	0.28	0.41	0.35	0.28
Avestin C-50	Homogenized Emulsion Pass #3	96hrs	Ave	0.22	0.32	0.59	0.46	0.31
Microfluidics M-110EH	Homogenized Emulsion Pass #1	0	Ave	0.19	0.27	0.39	0.33	0.27
Microfluidics M-110EH	Homogenized Emulsion Pass #1	24hrs	Ave	0.20	0.29	0.42	0.34	0.28
Microfluidics M-110EH	Homogenized Emulsion Pass #1	96hrs	Ave	0.23	0.32	0.49	0.39	0.32
Microfluidics M-110EH	Homogenized Emulsion Pass #2	0	Ave	0.19	0.26	0.35	0.27	0.27
Microfluidics M-110EH	Homogenized Emulsion Pass #2	24hrs	Ave	0.18	0.25	0.35	0.26	0.26
Microfluidics M-110EH	Homogenized Emulsion Pass #2	96hrs	Ave	0.20	0.28	0.38	0.29	0.28

# Emulsion Study Summary

- High precision and reproducible data from HORIBA LA-950 particle size analyzer provide the critical information for evaluating different equipment and processing conditions.
- Both Y and Z types interaction chambers from Microfluidics produce emulsions with fine size and fairly uniform distribution. Y type is slightly more efficient than Z type.
- Cooling study using Microfluidics demonstrates that immediate quench of the processed emulsions is a critical process parameter to control the emulsions stability.
- When employing new cooling strategy, Microfluidics F12Y interaction chamber is able to produce fine and single-mode emulsions in less than two passes.