

# OptiPrep™ Application Sheet C38

## (I) Maintenance of cells in suspension for chemical/physical measurements (II) Microfluidic cell encapsulation/cell sorting

- ◆ OptiPrep™ is a 60% (w/v) solution of iodixanol in water, density = 1.32 g/ml

### PART I: Maintenance of cells in suspension for chemical/physical measurements

#### 1. Background

This is a non-centrifugal application, in which OptiPrep™ is mixed with a suspension of cells in order to prevent their sedimentation during an extended physical measurement or separation. The first published paper to report this novel application was in 2001, when Farinas, Chow and Wada [1] commented in the abstract to their paper: “Recent developments in microfluidics have enabled the design of a lab-on-a-chip system capable of measuring cellular membrane potential. The chip accesses liquid samples sequentially by sipping from a micro-plate through a capillary, mixes the samples with cells flowing through a microchannel, contacts the cells with potential-sensitive dyes, and reads out cellular responses using fluorescence detection”. Later Culbertson [2] noted that the use of OptiPrep™ enabled the rate of cell entry into the main chip channel of a microfluidic device to remain constant over the course of an entire experiment. The technique permits the identification of single cells with unique compositional characteristics that may presage the later development of a disease state. In its simplest form, a medium with the same density as the cell, which in no way interacts with the cell, maintains the cell in a state of continuous suspension.

**Section 2** briefly describes solution preparation and the physical parameters of some the solutions that have been used in these studies. **Section 3** summarizes the cell types that have been studied and types of physico-chemical measurements that were carried out. **Section 4** lists some more recent publications. **Section 6** is the reference list.

#### 2. Solution preparation

Mix OptiPrep™ (shake the bottle gently before use) with saline or a balanced salt solution to give a solution of suitable density, according to the figures in Table 1. Table 2 summarizes a few of the conditions reported in the literature. For more information on making up solutions see [Application Sheet C01](#).

- ◆ The actual density required will depend on the cell type; it can be chosen by suspending the cells in a series of density solutions and observing the disposition of the cells after centrifugation. Because of the heterogeneity of any cell population however it is impossible to select a density that will allow all the cells to remain in suspension, some will float and some will sediment. If the cell suspension contains a variety of cell types the problem will be exaggerated. For example human peripheral blood mononuclear cells have a density range of 1.058-1.078 g/ml.

**Table 1:** Density of solutions produced by dilution of OptiPrep™ with an isotonic buffered saline solution

% (w/v) iodixanol	Refractive index	Density (g/ml)	% (w/v) iodixanol	Refractive index	Density (g/ml)
5.00	1.3429	1.032	11.00	1.3523	1.063
6.00	1.3444	1.037	12.00	1.3538	1.068
7.00	1.3459	1.043	13.00	1.3554	1.074
8.00	1.3475	1.048	14.00	1.3570	1.079
9.00	1.3491	1.053	15.00	1.3585	1.085
10.00	1.3507	1.058	16.00	1.3601	1.090

**Table 2:** Selected solution properties reported in the literature

Cell Type	Solution composition	Iodixanol (% w/v)	Density (g/ml)	Ref #
HeLa	PBS + 1% albumin	11.4	1.065	3
HeLa	Hank's Balanced Salt Solution	9.6	1.054	4
Human blood lymphocytes and THP-1 <sup>1</sup> cells	Hank's Balanced Salt Solution	10.8	1.062	1
Human blood leukocytes	PBS	3-9	1.02-1.05 <sup>2</sup>	5
Human erythrocytes	PBS	15-18	1.084-1.100	6
Human erythrocytes	PBS	17	1.095	7
Human erythrocytes	PBS/dextran <sup>3</sup>	2.6		8
Human erythrocytes	PBS/5.5mM glucose/4% BSA	19	1.105	9
Jurkat/lymphocytes	PBS	10.8	1.062	2
Macrophages	PBS	8.4	1.052	10
Yeast	Water	19	1.105	11,12

<sup>1</sup> THP-1 cells are a human acute leukaemic monocytic line

<sup>2</sup> The densities were chosen to allow the cells to sediment slowly to the bottom of the chamber

<sup>3</sup> The density is difficult to estimate because of the dextran content; the latter was added to raise the viscosity

### 3. Cell types and study topic

#### Breast carcinoma cells

Iso-acoustic focusing for phenotyping [29]

Radiometric assays [50]

Viability of cells in alginate particles [13]

#### Bronchial epithelial cells

Confocal light absorption and scattering spectroscopic (CLASS) microscopy [14]

#### Cancer cell lines, distinction of

SERS-microfluidic droplet platform [58]

#### CHO cells

Microfluidic cytometry [15, 48]

#### Embryonic spinal cord cells

Ribosome reduction [57]

#### Embryonic stem cells

Gene expression in single cells [16]

#### Erythrocytes

Acoustic cell manipulation [38]

Blood viscosity changes in response to exercise [17]

Channel flow profiling [9]

Cholesterol [53]

Density matching [39]

Highly mono-dispersed microdroplet production [35]

Lysis kinetics [7]

Nitric oxide scavenging by haemoglobin [6, 8]

Phase separation in micro-channel networks [31]

Plasma membrane cholesterol [52]

Shape and mechanical properties of erythrocytes [18]

#### *Escherichia coli*

Genetically encoded synthesis of nanomaterials [32]

#### Fibroblasts

Hydrogel encapsulation [12]

### *Francisella tularensis*

Cell sorting [10]

### HeLa cells

Cell heterogeneity in metal ion responses [4]

Fluorescence-activated microfluidic devices for cell identification and sorting [3, 19, 20, 21]

Microfluidic cytometry [45, 49]

### Hepatocarcinoma cells

Microfluidic cytometry [15]

### Hepatocytes

Microfluidic cytometry [42]

### HL60 cells

Differential detection photothermal spectroscopy: label-free detection [37]

### Human stem cells

Pancreatic  $\beta$ -cell transformation [52]

### Industrial effluent analysis

Membrane bioreactors [22]

### Jurkat cells

Microfluidic cytometry [15, 43, 46]

Microfluidic devices for cell identification and sorting [2]

### Leukocytes (human)

Acoustic cell manipulation [38]

Continuous concentration of cells [23]

Density matching [39]

Microfluidic devices for cell identification and sorting; orientation-dependent elastic light scattering [5]

### Lung adenocarcinoma epithelial cells

Microfluidic cytometry [41]

### Lymphocytic cells (incl. lymphoblasts and other similar cells)

Cell membrane potential measurements using potential-sensitive dyes [1]

Cell size and density changes; response to growth factor deprivation [24]

Hydrogel encapsulation [12]

Iso-acoustic focusing for phenotyping [29]

Microfluidic cytometry [15, 43]

PCR-based sorting [30]

Viscoelastic carrier fluids [51]

### Mesenchymal stem cells

Microfluidic cytometry [47]

### Mouse bone marrow stromal cells

Microfluidic cytometry [44]

### Mouse hybridoma cells

Analysis and sorting of single cells [25]

### Mouse pro-B lymphoid cells

Cell size, density and deformability [26]

### Mouse insulinoma cells

Enzymatic crosslinking of tyramine-functionalized polymer droplets [34]

### Neural cells

Density matching [40]

Single cell barcoding – droplet microfluidics [33]

### Prostate cancer cells

Printed droplet microfluidics: dispensing of picoliter droplets and cells [36]

### Rat liver lysosomes (stained with red tracker dye)

Confocal light absorption and scattering spectroscopic (CLASS) microscopy [14]

### Review articles

Extracellular matrix [56]

Recent advances in research involving chemical and biological reactions in microdroplets [27]

### *Staphylococcus aureus*

Microchannels [54]

### Tumor-infiltrating myeloid cells

Tumour growth regulation [55]

### Yeast cells

Acoustic cell manipulation [38]

Biogenic magnetization [28]

Hydrogel encapsulation [12]

Polyglycerol microgel particles for the micro-encapsulation of yeast cells [11]

## 4. Physical measurements (recent publications)

Density matching: see refs 59 and 60

Droplet sorting/merging: see ref 61

Microfluidic cytometry: see refs 62-66

Single cell analysis: see ref 67

## 5. References

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## (II) Microfluidic cell encapsulation/cell sorting

This new application of OptiPrep™ has only very recently featured regularly in the literature; publications are listed alphabetically according to **cell type**, or “**methodology**” or “**reviews**”.

### Adipose tissue stem cells

Morandi, E.M., Verstappen, R., Zwierzina, M.E., Geley, S., Pierer, G. and Ploner, C. (2016) *ITGAV and ITGA5 diversely regulate proliferation and adipogenic differentiation of human adipose derived stem cells* Sci. Rep., **6**: 28889

### Bacteria

Jusková, P., Schmid, Y.R.F., Stucki, A., Schmitt, S., Held, M. and Dittrich, P.S. (2019) “*Basicles*”: *microbial growth and production monitoring in giant lipid vesicles* ACS Appl. Mater. Interfaces, **11**, 34698–34706

### Carcinoma cells

Eastburn, D.J., Sciambi, A. and Abate, A.R. (2013) *Ultrahigh-throughput mammalian single-cell reverse-transcriptase polymerase chain reaction in microfluidic drops* Anal. Chem., **85**, 8016-8021

Jiang, Z., Xia, B., McBride, R. and Oakey, J. (2017) *A microfluidic-based cell encapsulation platform to achieve high long-term cell viability in photopolymerized PEGNB hydrogel microspheres* J. Mater. Chem. B, **5**, 173-180

Pei, H., Li, L., Wang, Y., Sheng, R., Wang, Y., Xie, S., Shui, L., Si, H. and Tang, B. (2019) *Single-cell phenotypic profiling of CTCs in whole blood using an integrated microfluidic device* Anal. Chem., **91**, 11078–11084

Sun, D., Cao, F., Cong, L., Xu, W., Chen, Q., Shic, W. and Xu, S. (2019) *Cellular heterogeneity identified by single-cell alkaline phosphatase (ALP) via a SERRS microfluidic droplet platform* Lab Chip, **19**, 335-342

### B-lymphocytes

Eastburn, D.J., Sciambi, A. and Abate, A.R. (2013) *Ultrahigh-throughput mammalian single-cell reverse-transcriptase polymerase chain reaction in microfluidic drops* Anal. Chem., **85**, 8016-8021

Holzner, G., Du, Y., Cao, X., Choo, J., deMello, A.J. and Stavrakis, S. (2018) *An optofluidic system with integrated microlens arrays for parallel imaging flow cytometry* Lab Chip, **18**, 3631–3637

### Bone marrow cells

Asensio, M.A., Lim, Y.W., Wayham, N., Stadtmiller, K., Edgar, R.C., Leong, J., Leong, R., Mizrahi, R.A. Adams, M.S. et al (2019) *Antibody repertoire analysis of mouse immunization protocols using microfluidics and molecular genomics* mAbs **11**, 870–883

Petukhov, V., Guo, J., Baryawno, N., Severe, N., Scadden, D.T., Samsonova, M.G. and Kharchenko, P.V. (2018) *dropEst: pipeline for accurate estimation of molecular counts in droplet-based single-cell RNA-seq experiments* Genome Biol. **19**: 78

**Breast cancer cells**

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