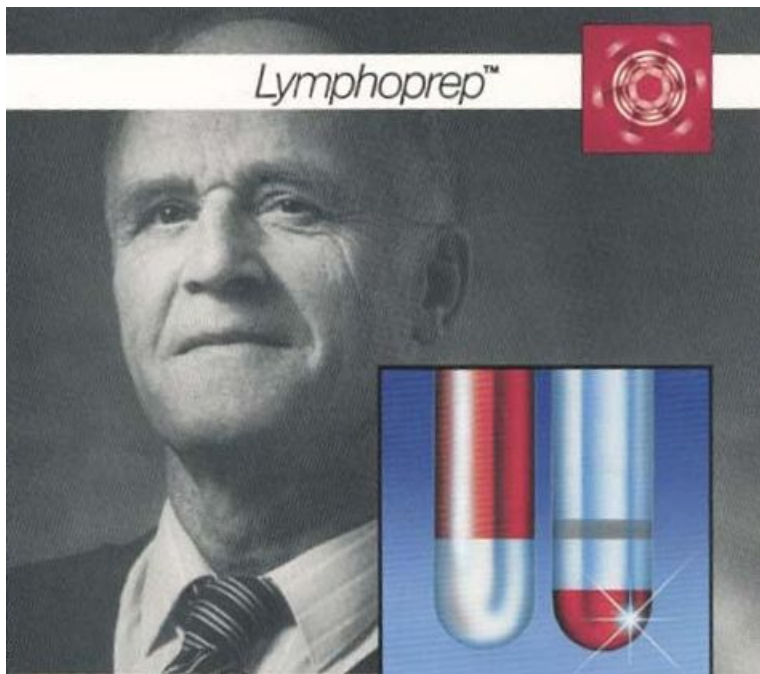
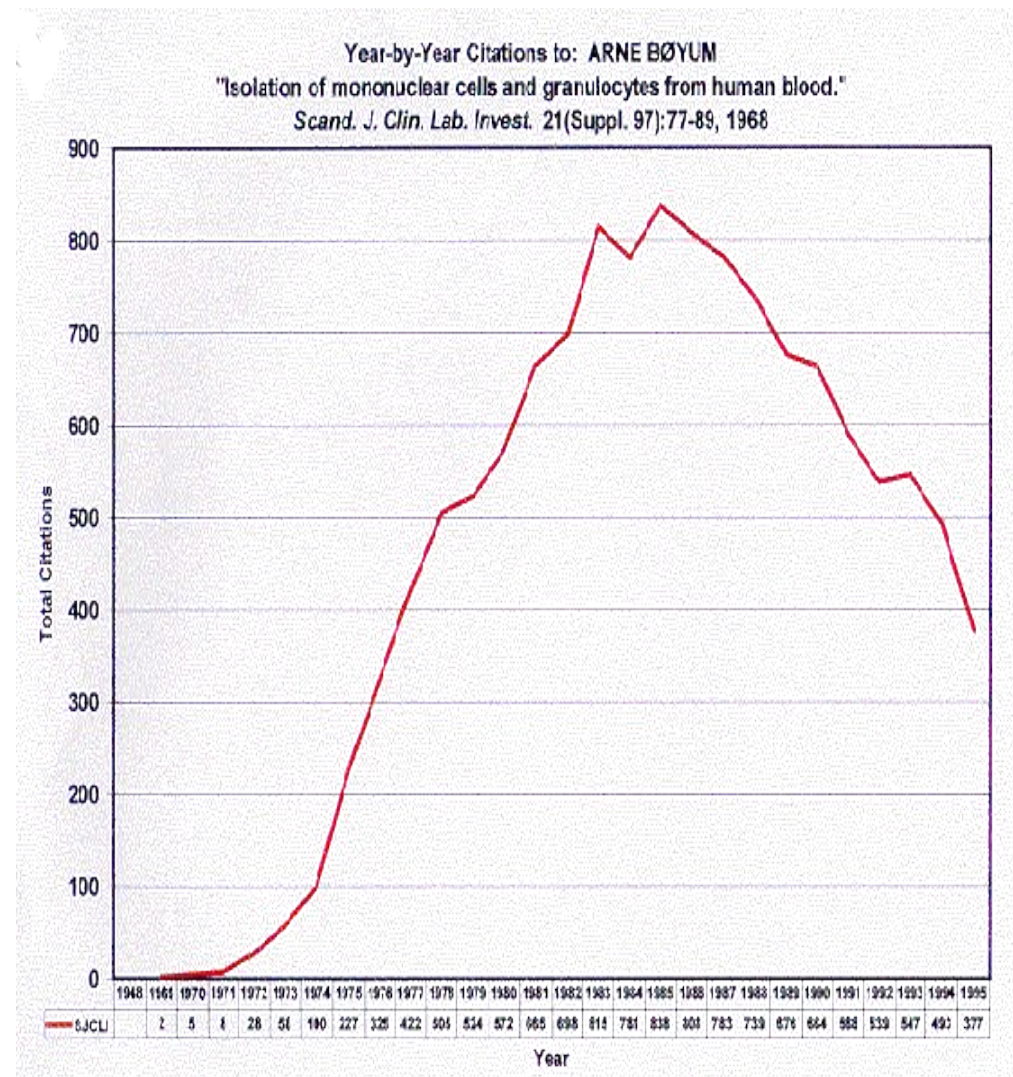




# Автор метода выделения и разделения лейкоцитов периферической крови человека в градиенте плотности



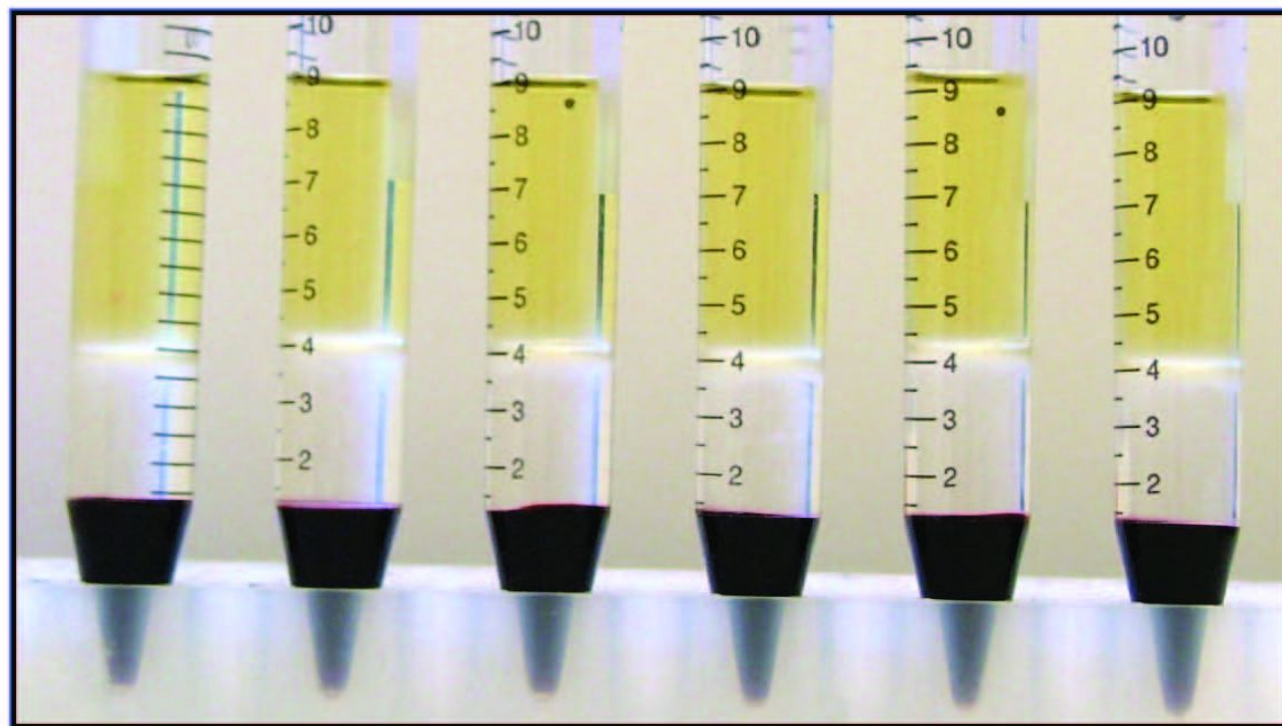
**Arne Bøyum**  
Division for Toxicology  
Norwegian Defence  
Research  
Establishment  
N-2007 Kjeller Norway



# Метод выделения и разделения лейкоцитов периферической крови в градиенте плотности фиколл-гипак (фиколл-дитриазоат натрия, $d=1,077 \text{ г/см}^3$ )



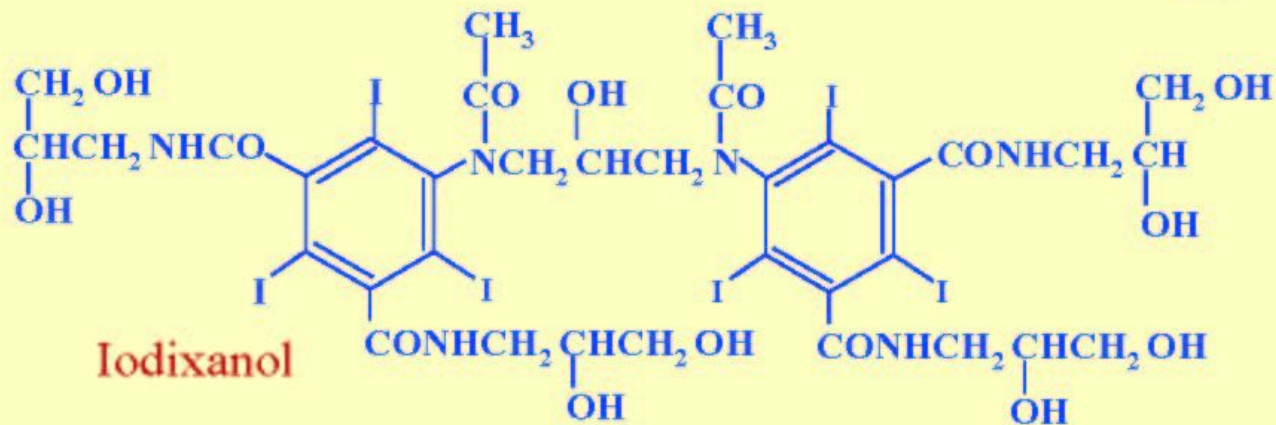
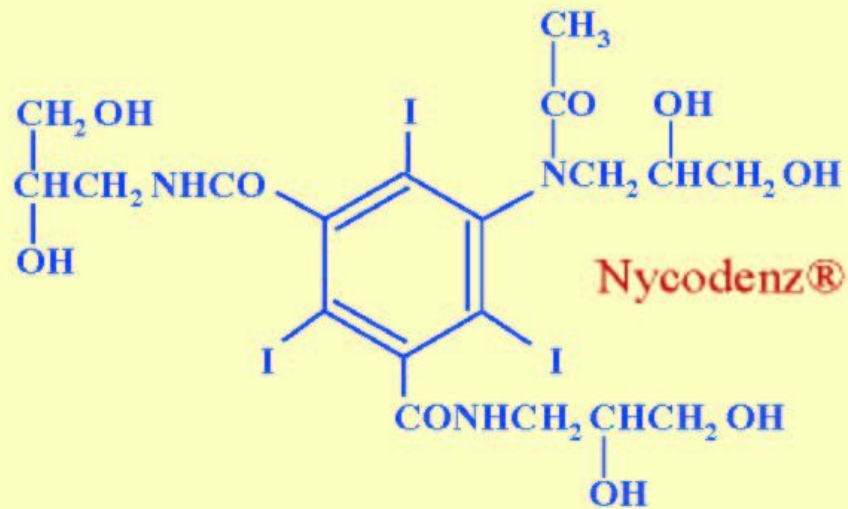
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Norwegian Defence  
Research  
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**Figure 2:** Purification of human PBMCs from six donors on *Lymphoprep*<sup>TM</sup>



**Sodium diatrizoate**

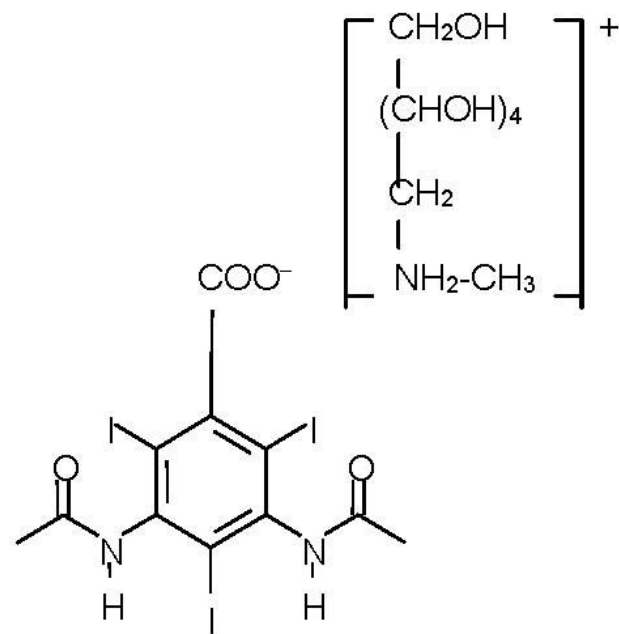
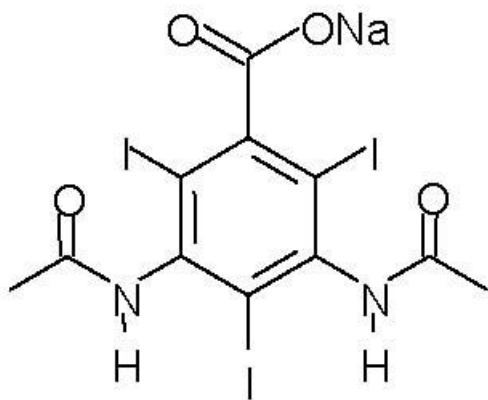


*Figure 1: Molecular structure of iodinated density gradient media*

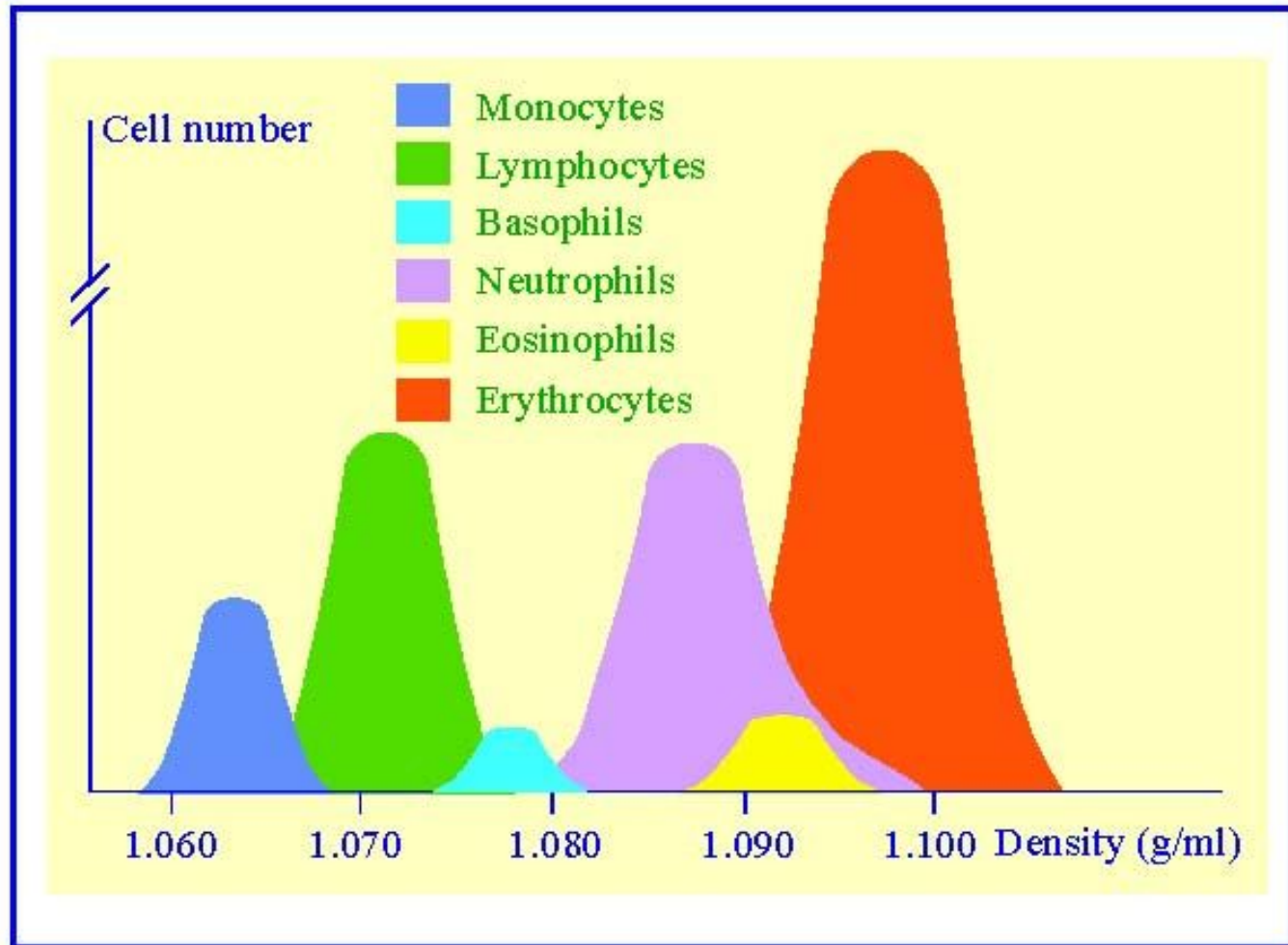
# Верографин 76% СПОФА (Прага)

Смесь N-метил-D-глюкаминовой и натриевой солей диатриазоевой кислоты (3,5-бис-(ацетиламино)-2,4,6-трийодбензойной кислоты) (86,85:13,15%) в ампулах по 20 мл для инъекций. Аналог верографина - 76% раствор урографина выпускается Байер Шеринг Фарма АГ (Германия D-13342 Берлин).

## Натрия диатриазоат



## Меглюмин диатриазоат



**FIGURE 1.** Density of human blood cells.

# Lympholyte<sup>®</sup> Cell Separation Media

CEDARLANE<sup>®</sup> has specifically designed cell separation centrifugation media for the isolation of viable lymphocytes from mouse, rat, rabbit, human and other mammalian cell populations. All products are supplied as sterile liquid with varying densities.

Applications for **Lympholyte<sup>®</sup>** include:

- isolation of viable lymphocytes from lymphoid organs (particularly spleen) by removal of red cells and dead cells.
- removal of dead cells from lymphocyte suspensions; for example, after treatment with antibody plus complement, or following cell culture.
- **Lympholyte<sup>®</sup>-H** and **Lympholyte<sup>®</sup>-Mammal** are preferred for the isolation of viable lymphocytes from peripheral blood.

Product	Density (@ 25°C, g/ml)	Cat.#	Size	Main Application
<b>Lympholyte<sup>®</sup>-M</b>	1.0875 ± 0.001	<b>CL5030</b>	5x30ml	Isolation of viable MOUSE lymphocytes from lymphoid tissue.
		<b>CL5031</b>	100ml	
		<b>CL5035</b>	500ml	<b>Not recommended for mouse blood, see Lympholyte<sup>®</sup>-Mammal</b>
<b>Lympholyte<sup>®</sup>-Rat</b>	1.0940 ± 0.001	<b>CL5040</b>	5x30ml	Isolation of viable RAT lymphocytes from lymphoid tissue
		<b>CL5041</b>	100ml	
		<b>CL5045</b>	500ml	
<b>Lympholyte<sup>®</sup>-H</b> CE	1.0770 ± 0.001	<b>CL5010</b>	5x30ml	Isolation of viable HUMAN lymphocytes from peripheral blood
		<b>CL5015</b>	100ml	
		<b>CL5016</b>	6x100ml	
		<b>CL5020</b>	500ml	
		<b>CL5026</b>	6x500ml	
<b>Lympholyte<sup>®</sup>-Rabbit</b>	1.0965 ± 0.001	<b>CL5050</b>	5x30ml	Isolation of viable RABBIT lymphocytes from lymphoid tissue
<b>Lympholyte<sup>®</sup>-Mammal</b>	1.0860 ± 0.001	<b>CL5110</b>	5x30ml	Isolation of viable lymphocytes from peripheral blood of most mammalian species
		<b>CL5115</b>	100ml	
		<b>CL5120</b>	500ml	
<b>Lympholyte<sup>®</sup>-poly</b>	1.113 ± 0.001	<b>CL5070</b>	100ml	Isolation of viable HUMAN polymorphonuclear cells
		<b>CL5071</b>	250ml	
<b>Lympholyte<sup>®</sup>-1.1</b>	1.100 ± 0.001	<b>CL5095</b>	500ml	A high density Lympholyte <sup>®</sup> solution which can be diluted with PBS (without Ca <sup>2+</sup> /Mg <sup>2+</sup> ). Isolation of pancreatic islet cells prior to expanding them <i>in vitro</i>
<b>Metrizamide AG</b> (Analytical Grade)	(powder)	<b>CL5080</b>	50g	Fractionation of nucleic acids, proteins, and polysaccharides; Separation of lysosomes from mitochondria and peroxisomes
<b>Polysucrose 400</b>	(powder)	<b>CL5401</b>	100g	To prepare density gradient media for separation of different spleen, bone marrow and other tissues
	(powder)	<b>CL5405</b>	500g	
<b>Polysucrose 70</b>	(powder)	<b>CL5701</b>	100g	For cell isolation and animal perfusion
	(powder)	<b>CL5705</b>	500g	

The CE Mark indicates that CEDARLANE<sup>®</sup> Laboratories has met the registration requirements for the *In Vitro* Diagnostic Directive 98/79/EC.

Please contact...

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LABORATORIES LIMITED

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