

Data Sheet

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LUXOL FAST BLUE

Klüver Barrera



CODE DESCRIPTION TESTS NUMBER

Luxol fast blue Kluver Barrera



04-200812

In Vitro Diagnostic – medical device IVD in **Class A**, Reg. UE 2017/746 Basic UDI: 080339762W01030799Y5 UDI-DI: 08033976231460



100 test



Product for the preparation of cyto-histological samples for optical microscopy. To show myelin and phospholipids in histologic sections.

PRINCIPLE

Luxol fast blue dye is a derivative of tetrabenzotetrazo-porphyrin. Kluver has demonstrated porphyrins have a selective affinity for myelin (see references). Luxol fast blue's affinity for central nervous system is usually ascribed to the bonds it forms with phospholipidic structures such as lecithin and sphingomyelin.

METHOD

- 1) Deparaffinise and bring section to ethanol 95°.
- 2) Prepare the incubation box by adding some drops of distilled water on filter paper in Petri dish and lay down the slide; put on the slide 10 drops of reagent A, close the incubation box and incubate at 56°C overnight in oven.
- 3) Extract the slide from oven and wash it with ethanol 95° (crystalline residues of reagent A should melt).
- 4) Wash in distilled water.
- 5) Put on the section 10 drops of reagent B: leave to act 30 seconds.
- 6) Differentiate in ethanol 70° until myelinic fibres become blue on colourless background (Sometimes differentiation can be difficult; repeat the step 5 for 30 seconds and put the slide again in ethanol 70°)
- 7) Wash well in distilled water (at least 2 times).
- 8) Prepare the incubation box again and introduce the slide; put on the section 10 drops of reagent C and 5 drops of reagent D: close the incubate box and incubate for 20 minutes at 56°C in oven.
- 9) Differentiate in ethanol 95° until Nissl substance results pale pink.
- 10) Dehydrate in absolute ethanol, clear in xylene and mount.



The picture is for illustrative purposes only



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Technical details

	Procedure time	20 minutes + overnight	
	Complementary equipment	Not requested	
Method specifications	Results	Myelin:	Turquoise blue
		Neurons and glial nuclei:	Pink - violet
		Nissl substance:	Pale pink
Components	A) Luxol fast blue alcoholic solution	30 ml	
	B) Basic differentiating buffer	30 ml	
	C) Cresyl violet aqueous solution	30 ml	
	D) Acid activation buffer	30 ml	
Storage	Storage	Store the preparation at 15 - 25°C. Keep the containers tightly closed.	
	Storage temperature	15 - 25°C	
	Stability	After the first opening, the product is reusable until the expiry date, if correctly stored.	
	Validity	2 years	
Warning	Product classification	The product is intended for professional laboratory use for healthcare professionals. Carefully read the information on the label (danger symbols, risk and safety phrases) and always consult the safety data sheet. Do not use if the primary container is damaged. In the event of a serious accident, we recommended that you immediately inform Bio-Optica Milano S.p.A and the competent authorities.	
	Disposal	Hazardous preparation: observe all state and local environmental regulations regarding waste disposal.	

REVISION n°	REASON	REVISION DATE
001	Regulation adjustment UE 2017/746 - IVDR	16/05/2022