

Trypan UltraBlue™, Trypan Purple™, and Trypan Red Plus™

Biological Properties

Trypan UltraBlue™, Trypan Purple™, and Trypan Red Plus™ are similar to Trypan Blue in cell permeability. It is not permeable to live cells. Compared to Trypan Blue, these new trypan compounds are less toxic to cells. In particular, they have minimal effect on cell surface receptors such as G-protein coupled receptors (GPCRs). Another advantage is that the cells can be clearly observed under microscope when Trypan Red Plus™ is used while Trypan Blue makes it quite difficult to see cells under microscope.

Our Trypan UltraBlue™, Trypan Purple™, Trypan Red Plus™ and Trypan UltraRed™ can also be used to prevent fluorescent dyes (such as FDA, rhodamine 123, JC-1, TMRA, TMRM, Indo-1 AM, Fura-2 AM, calcein AM, Fluo-3 AM, Fluo-4 AM and Fluo-8 AM) from leaking out of cells. They might inhibit the activities of drug-efflux pumps since they contain a probenecid-like moiety as shown below. Compared to probenecid, they are neutral, highly soluble in water, and convenient to use. Their cellular mechanisms are still under investigation.



Figure 1. The structure of Trypan Red Plus™
(WSH = water-soluble head; PLM = probenecid-like moiety)

Our Trypan Purple™, and Trypan Red Plus™ are highly purified, and can be used up to 1 mM with minimal cell cytotoxicity. A certain volume of our concentrated Trypan UltraBlue™, Trypan Purple™, and Trypan Red Plus™ solutions can be added into the assay system to have the final concentrations of Trypan UltraBlue™, Trypan Purple™, and Trypan Red Plus™ ranging from 0.1 to 1.0 mM depending on the cell lines used. The recommended concentrations are from 0.25 to 0.75 mM.

Ordering Information

Cat. #	Product Name	Unit Size
2450	Trypan Blue, sodium salt *Cell culture tested*	100 g
2452	Trypan Blue, sodium salt *UltraPure grade* *Purified to eliminate fluorescent impurities*	10 g
2455	Trypan UltraBlue™, sodium salt *0.1 M aqueous solution*	1 mL
2456	Trypan Red Plus™ *0.1 M aqueous solution*	10 mL
2457	Trypan Red Plus™ *0.1 M aqueous solution*	100 mL
2465	Trypan Purple™ *0.1 M aqueous solution*	10 mL
2466	Trypan Purple™ *0.1 M aqueous solution*	100 mL

Storage Conditions

Store at room temperature. Expiration date is 6 months from the date of receipt.

References

1. Hirasawa H, Yanagi Y, Tamaki Y, Inoue Y, Kadonosono K. (2007) Indocyanine green and trypan blue: intracellular uptake and extracellular binding by human retinal pigment epithelial cells. *Retina*, 27, 375.
2. Nanavaty MA, Johar K, Sivasankaran MA, Vasavada AR, Praveen MR, Zetterstrom C. (2006) Effect of trypan blue staining on the density and viability of lens epithelial cells in white cataract. *J Cataract Refract Surg*, 32, 1483.
3. Li Q, Kato Y, Sai Y, Imai T, Tsuji A. (2005) Multidrug resistance-associated protein 1 functions as an efflux pump of xenobiotics in the skin. *Pharm Res*, 22, 842.
4. Orlicky J, Sulova Z, Dovinova I, Fiala R, Zahradnikova A, Jr., Breier A. (2004) Functional fluo-3/AM assay on P-glycoprotein transport activity in L1210/VCR cells by confocal microscopy. *Gen Physiol Biophys*, 23, 357.
5. Heinemann A, Ofner M, Amann R, Peskar BA. (2003) A novel assay to measure the calcium flux in human basophils: effects of chemokines and nerve growth factor. *Pharmacology*, 67, 49.
6. Abrahamse SL, Rechkemmer G. (2001) Identification of an organic anion transport system in the human colon carcinoma cell line HT29 clone 19A. *Pflugers Arch*, 441, 529.
7. Fast VG, Ideker RE. (2000) Simultaneous optical mapping of transmembrane potential and intracellular calcium in myocyte cultures. *J Cardiovasc Electrophysiol*, 11, 547.
8. Packham MA, Rand ML, Perry DW, Ruben DH, Kinlough-Rathbone RL. (1996) Probenecid inhibits platelet responses to aggregating agents in vitro and has a synergistic inhibitory effect with penicillin G. *Thromb Haemost*, 76, 239.
9. Merritt JE, McCarthy SA, Davies MP, Moores KE. (1990) Use of fluo-3 to measure cytosolic Ca²⁺ in platelets and neutrophils. Loading cells with the dye, calibration of traces, measurements in the presence of plasma, and buffering of cytosolic Ca²⁺. *Biochem J*, 269, 513.
10. Kermod JC, Zheng Q, Cook EP. (1996) Fluorescent indicators give biased estimates of intracellular free calcium change in aggregating platelets: implication for studies with human von Willebrand factor. *Blood Cells Mol Dis*, 22, 238.
11. Chaka W, Scharringa J, Verheul AF, Verhoef J, Van Strijp AG, Hoepelman IM. (1995) Quantitative analysis of phagocytosis and killing of *Cryptococcus neoformans* by human peripheral blood mononuclear cells by flow cytometry. *Clin Diagn Lab Immunol*, 2, 753.
12. Wan CP, Park CS, Lau BH. (1993) A rapid and simple microfluorometric phagocytosis assay. *J Immunol Methods*, 162, 1.
13. Wong K, Kwan-Yeung L. (1993) Sphingosine mobilizes intracellular calcium in human neutrophils. *Cell Calcium*, 14, 493.
14. Joling P, Bakker LJ, Van Strijp JA, Meerloo T, de Graaf L, Dekker ME, Goudsmit J, Verhoef J, Schuurman HJ. (1993) Binding of human immunodeficiency virus type-1 to follicular dendritic cells in vitro is complement dependent. *J Immunol*, 150, 1065.
15. Noma T, Yoshizawa I, Nakamura Y, Kawano Y, Nakajima T, Itoh M, Koku K, Maeda K, Ikezawa Z, Baba M, et al. (1992) A rapid measuring technique for allergen-induced IL2 responsiveness of lymphocytes by the propidium iodide-staining method. Detection of the etiological antigen in patients with allergic diseases. *Arerugi*, 41, 1354.
16. Martin RJ, Kusel JR. (1992) On the distribution of a fluorescent ivermectin probe (4" 5,7 dimethyl-bodipy propionylivermectin) in *Ascaris* membranes. *Parasitology*, 104 (Pt 3), 549.

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