

Materials List for:

Assaying Blood Cell Populations of the *Drosophila melanogaster* Larva

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Materials

Name	Company	Catalog Number	Comments
6 cm/9 cm Petri dishes			One for each genotype to be evaluated
Water squirt bottle			
Metal spoon/spatula			
Thin paintbrush			e.g., a "liner"
Glass cavity dish			
PAP pen: Super PAP PEN IM3580	Beckman Coulter		
Glass slides			Each slide will have 5 or more PAP PEN squares drawn on them. Size of squares depends on the imaging objective and magnification of the microscope camera; e.g., 2 mm squares.
Moist chamber			This will be used to prevent slides and wells from drying out: sealed container with wet paper towels lining the sides/bottom
Schneider's <i>Drosophila</i> cell culture media	Invitrogen		
Cold block			This is a metal block (a.k.a. heating block) chilled in bucket containing ice; preferably black-colored or other dark, non-reflective color
2 x 1 ml syringes with needles (27 G ½")	Becton Dickinson		For dissections.
Optional: Surgical spring scissors (cutting edge 2 mm)	Fine Science Tools		
Glass beads, 212 - 600 µm	Sigma		
2 ml Eppendorf tubes	Eppendorf		One per genotype evaluating
Vortex mixer	Fisher Scientific		
Transgenic <i>Drosophila</i> larvae with fluorescently marked hemocytes.			Suitable transgenes include: HmlΔ-DsRed (Makhijani <i>et al.</i> , 2011), MSNF9mo-mCherry (Tokusumi <i>et al.</i> , 2009), BcF6-CFP and -GFP (Gajewski <i>et al.</i> , 2007), or HmlΔ-GAL4 (Sinenko and Mathey-Prevot, 2004), Pxn-GAL4 (Stramer <i>et al.</i> , 2005), He-GAL4 (Zettervall <i>et al.</i> , 2004), Crq-GAL4 (by H. Agaisse (Stramer <i>et al.</i> , 2005)), or eater-GAL4 (Tokusumi <i>et al.</i> , 2009) combined with UAS-GFP or other fluorescent protein transgenes.

Fluorescence dissecting microscope	Leica		Here: Leica M205, optional with camera, imaging software and measuring module
Inverted fluorescence microscope with camera attachment	Leica or Keyence		With or without tile scanning function (e.g., Leica DMI series, Keyence BIOREVO BZ-9000 series)