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An incorrect graphic appears in the legend for Fig. 1. The corrected sentence appears below in bold.

Figure 1. A two-step subcellular fractionation procedure for separation of late endosomes and DCG. (A) Distribution of intracellular compartments from PC12 cells on 1–16% Ficoll gradients. PC12 cells expressing ssHRP$^{P\text{-selectin}}$ were loaded with ^{3}H-Dopamine, or labeled with ^{125}I-Trn or ^{125}I-EGF as described in Materials and Methods. Cells were homogenized in HB and the PNS was centrifuged on 1–16% Ficoll gradients and fractionated. The early/recycling endosomes are shown by the distribution of ^{125}I-Trn endocytosed for 60 min at 37°C (△). Late endosomes are shown by the distribution of ^{125}I-EGF internalized for 20 min at 37°C (■). **The distribution of NAGA activity ($OD_{420\ nm}$; ○), HRP activity ($OD_{450\ nm}$; ●), and ^{3}H-Dopamine radioactivity (filled plus signs) along 1–16% Ficoll gradients are shown.** (B) Separation of DCG and late endosomes on a secondary 0.9–1.85 M sucrose gradient. Fractions 14–20 from the 1–16% Ficoll gradients were pooled, diluted with HB, and recentrifuged on 0.9–1.85 M sucrose gradients to equilibrium. The distributions of NAGA activity (○), HRP activity (●), ^{3}H-Dopamine radioactivity (filled plus signs), ^{125}I-EGF internalized for 20 min at 37°C (■) and ^{125}I-Trn (△) after centrifugation on this gradient are shown.