Supplementary Material for - A hybrid-membrane migration method to isolate high-purity adipose-derived stem cells from fat tissues

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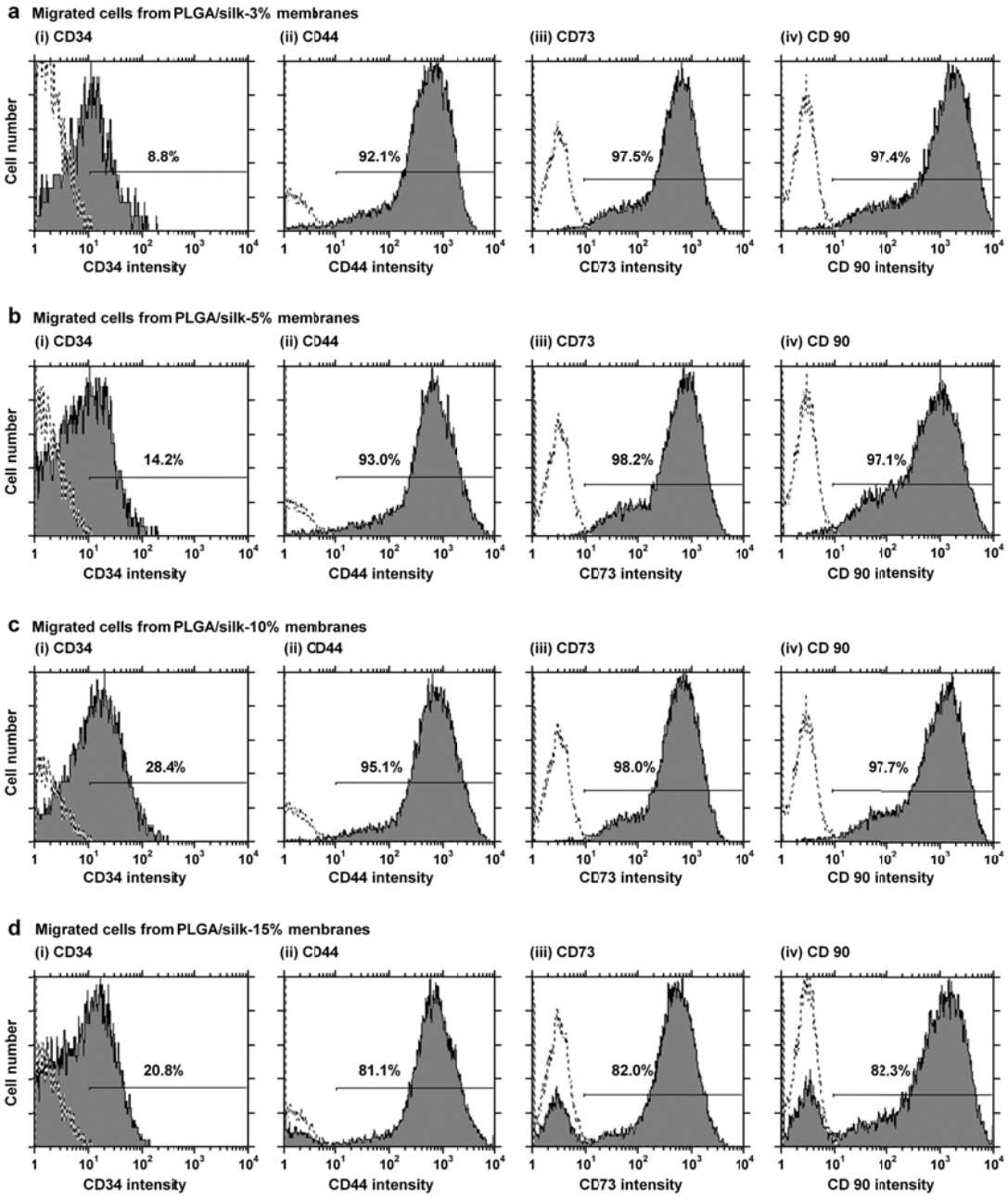
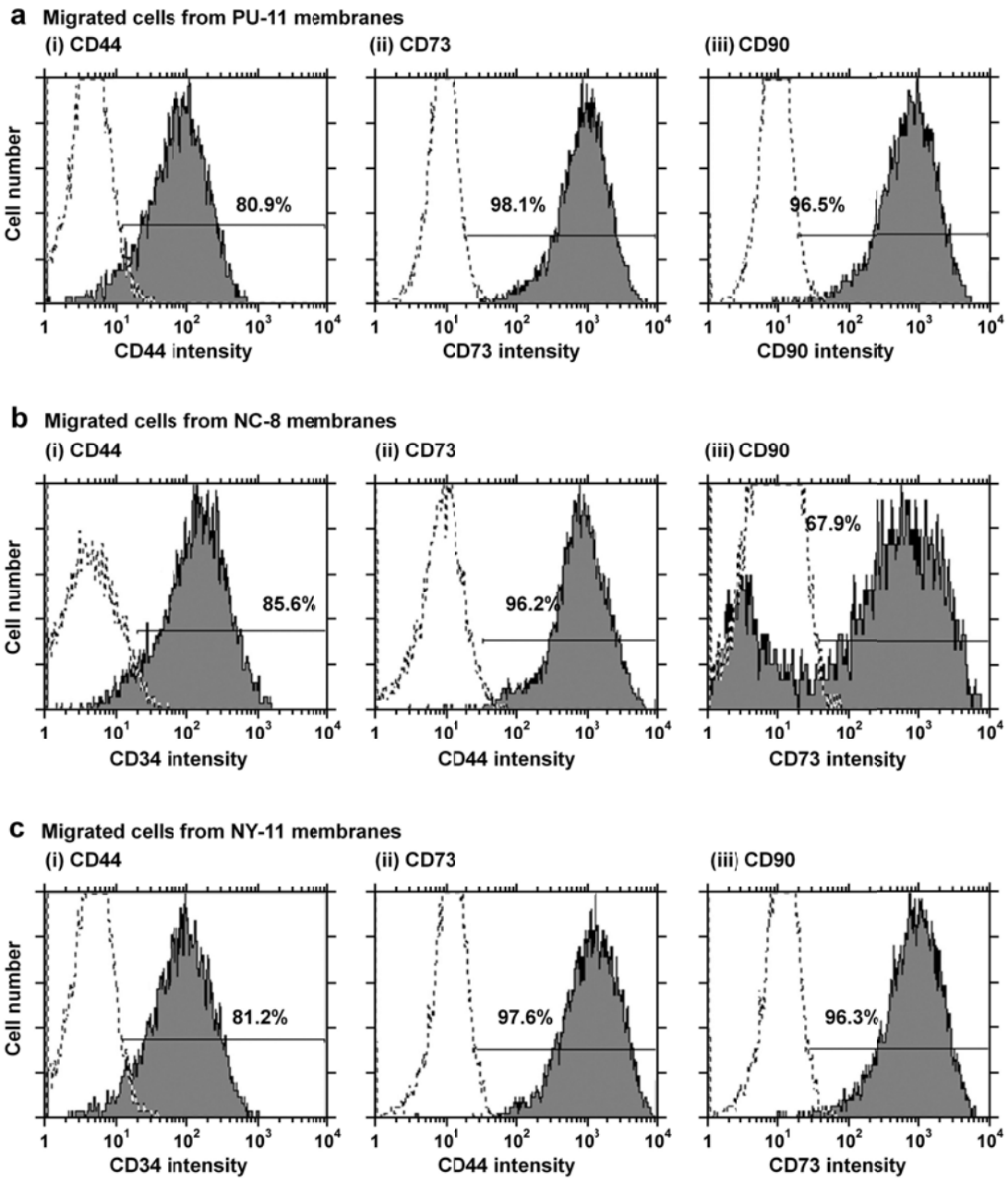
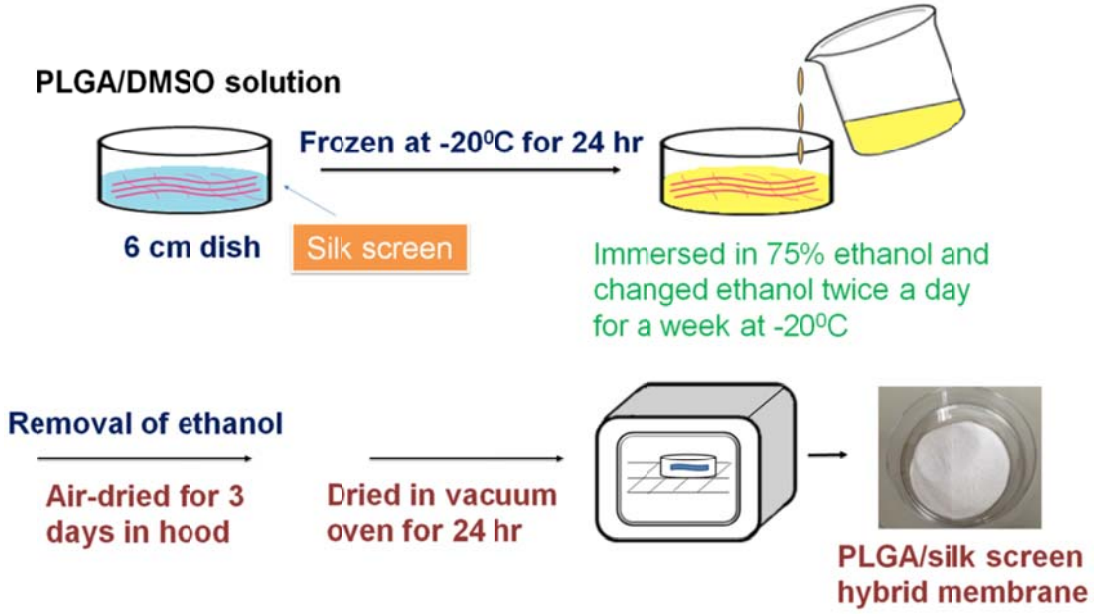
SUPPLEMENTARY FIGURES



Supplementary Figure 1. CD34 (i) and MSC markers (CD44 (ii), CD73 (iii), and CD90 (iv)) expression analyzed by flow cytometry. Flow cytometry scattergrams of the migrated cells from (**a**) PLGA/silk-3%, (**b**) PLGA/silk-5%, (**c**) PLGA/silk-10% and (**d**) PLGA/silk-15% membranes and subsequently cultured for 15 days after SVF was permeated through the membranes. The dotted lines indicate cells labeled with the isotype controls.



Supplementary Figure 2. MSC markers (CD44 [i], CD73 [ii], and CD90 [iii]) expression analyzed by flow cytometry. Flow cytometry scattergrams of the migrated cells from (**a**) PU-11, (**b**) NC-8 and (**c**) NY-11membranes and subsequently cultured for 15 days after SVF was permeated through the membranes. The dotted lines indicate cells labeled with the isotype controls.



Supplementary Figure 3. Manufacturing procedures of the PLGA/silk membranes prepared by the freeze-extraction method.

SUPPLEMENTARY TABLE

Supplementary Table 1 Characteristics of membranes used in this study.

Membranes	Materials	Pore size (μm)	Porosity (%)	Thickness (μm)
Silk-PLGA-3%	Silk & PLGA	24.4		
Silk-PLGA-5%	Silk & PLGA	22.4		
Silk-PLGA-10%	Silk & PLGA	20.8		
Silk-PLGA-15%	Silk screen & PLGA	18.2		
PU-11	Polyurethane	11.0	86	1200
NC-8	Nitrocellulose	8.0	84	135
NY11	Nylon	11.0	6	65