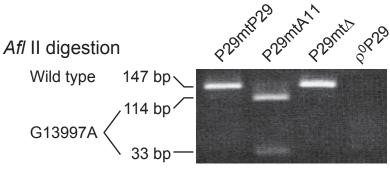
Table 1 Genetic characteristics of parent cells and their trans-mitochondrial cybrids

Cell lines <sup>a</sup>	Nuclear genotypes (genetic marker) <sup>b</sup>	mtDNA - genotypes	Fusion combination			
			nuclear	×	mtDNA	Selection
			donors		donors	
Nuclear donors						
ρ <sup>0</sup> P29	P29 (HAT <sup>r</sup> , BrdU <sup>s</sup> )	mtDNA less				
mtDNA donors						
B82mtP29	B82 (HAT <sup>s</sup> , BrdU <sup>r</sup> )	Wild type	$\rho^0$ B82	×	en <sup>c</sup> P29	BrdU + UP
B82mtA11	B82 (HAT <sup>s</sup> , BrdU <sup>r</sup> )	G13997A	$\rho^0$ B82	×	enA11	BrdU + UP⁻
B82mt∆	B82 (HAT <sup>s</sup> , BrdU <sup>r</sup> )	∆mtDNA4696	ρ <sup>0</sup> B82	×	platelets	UP <sup>-</sup>
Trans-mitochondrial	cybrids					
P29mtP29	P29 (HAT <sup>r</sup> , BrdU <sup>s</sup> )	Wild type	$\rho^0$ P29	×	enB82mtP29	HAT + UP
P29mtA11	P29 (HAT <sup>r</sup> , BrdU <sup>s</sup> )	G13997A	ρ <sup>0</sup> P29	×	enB82mtA11	HAT + UP
P29mt∆	P29 (HAT <sup>r</sup> , BrdU <sup>s</sup> )	∆mtDNA4696	ρ <sup>0</sup> P29	×	enB82mt∆	HAT + UP

- As mtDNA donors, we used B82mtP29, B82mtA11, and B82mt $\Delta$  cybrids shearing the same nuclear background of B82 cells for excluding variations of nuclear-coded cytoplasmic factors in mtDNA donors. B82mtP29 cybrids carrying nuclear DNA from B82 cells and mtDNA from P29 cells were obtained by fusion of  $\rho^0$ B82 cells with enucleated P29 cells and subsequent cultivation in the selection medium with BrdU and UP $^-$ .  $\rho^0$ B82 cells can survive in the selection medium with BrdU due to their lacking thimidine kinase activity, and cannot survive in the selection medium without uridine and pyruvate (UP $^-$  medium) due to their lacking mtDNA. Thus, BrdU and UP $^-$  eliminate unenucleated P29 cells and unfused  $\rho^0$ B82 cells, respectively, and allow exclusive growth of the B82mtP29 cybrids. B82mt $\Delta$  cybrids carrying nuclear DNA from B82 cells and  $\Delta$ mtDNA4696 were obtained by fusion of  $\rho^0$ B82 cells with platelets from mito-mice carrying  $\Delta$ mtDNA4696 [23] in the UP $^-$  selection medium. As G13997A mtDNA donors, we used B82mtA11 cybrids obtained in our previous work [16].
- All the mtDNA donors sharing the B82 nuclear background lacking thymidine kinase activity cannot survive in the presence of a hypoxanthine/aminopterin/thymidine (HAT). On the contrary, nuclear donors ρ<sup>0</sup>P29 cells can grow in the HAT selection medium due to their prossessing thimidine kinase activity, but not in UP<sup>-</sup> selection medium due to their complete respiration defects by mtDNA depletion. Thus, HAT and UP<sup>-</sup> allow exclusive growth of the P29mtP29, P29mtA11, and P29mtΔ cybrids.
- c en represents enucleated.

Figure 1





## Xho I digestion





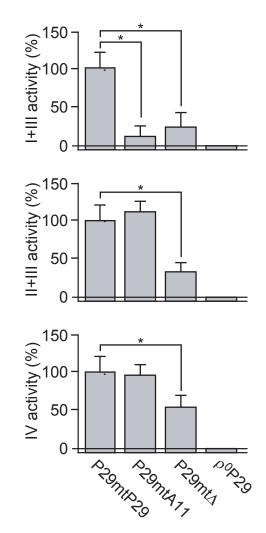
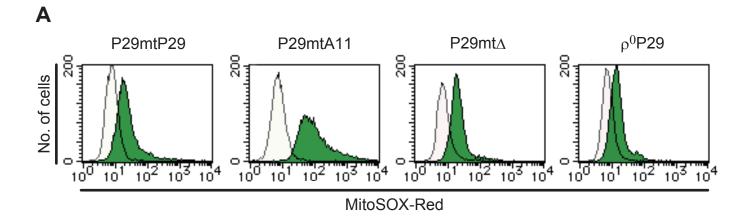


Figure 2



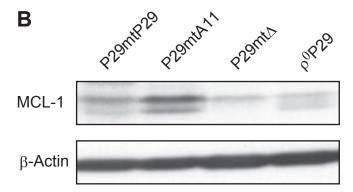


Figure 3

