JCRB C	ell Ba	ınk		Notes]									
Distributi	on			Responsible	e person : Kurematsu, M]							
Source Information					Product Information									
Registered c	ell No:	JCRB00	98		Passage Number: P16*		Viable cell number (a): 2.08x10^6 cells/ml							
Cell strain name: KURAMOCHI number: 122386					Type of ampoules: glass		Total cell number (b): 2.23x10^6 cells/ml Results of quality con						trols.	
Passage Number: P13* [ype of Source: from					No of ampoules:34		Viability(a/b x 100):	96.06	%	Karyotype	Myco.	Bac./ Fung	Isoenzyme	
Location of Amp: N1-N-5					Lot number:09029	98	No of colonies/100 cells:	NT	%					
Date Pr	epared:	?			Location of am	p: N2-E-2	Growth curve estimations:	NT						
Human asc	ites ova	rian cance	er	Culture Conditions:	Passage History: From P13* toP16*, 16-days.									
undifferent	ated ca	rcinoma			Subculture Method: Harvested with 0.25% trypsin and 0.02% EDTA (at R.T., 5min)									
Date	Date Total Number Passage				(Anchorage dependent culture)									
mm/dd/yy	culture	of plates	number	ledium Information:	Medium:	RPMI1640	0)71698 prej	pared ; r	nixed with	serum on	073098	Inoculume	e Size
	days				Serum:	10% FCS	10	ot# MITSU	BISHI	PVF01			Start	End
08/17/98	0	100m/m,1	P14*	Retrieve culture: The	cells in 1-amp. were	washed 2-times and su	uspended 2ml. The number of the	cells was co	ounted w	ith Tatai 4-f	olds dill.(4	412:24, 1/	2block).	
				Viability; 94.50%. V	Whole viable cell num	bers: 2.64x10^6 cells.	. They were inoculated to 100m./r	n. dish.						
				Approximately cell de	ensity: 2.64x10^5 cel	ls/ml, 4.47x10^4 cells/	/cm^2.						4.5x10^4/cm	5.0x10^4/cm2
08/18/98	1	100m/m,1	P14*	Took the photographs	s and medium ware cl	hanged.								
08/21/98	4	100m/m,3	P15*	Subculture:Cells were	e harvested with 0.259	% trypsin (5min at R.7	Γ.), suspended to 10ml and counte	ed with Tata	i at 4-fol	lds dill.(538:	4 1-block)	. Viabilit	3	
				Whole viable cell nun	nbers 8.61x10^6 cells	s. Inoculated to new cu	alture dish with MC210. An appro-	oximate. cel	l density	:1.20x10^5	cells/ml, 2	.03x10^4	2.0x10^4/cm	1.4x10^5/cm2
08/25/98	8	100m/m,3	P15*	Added fresh medium(5ml) into each dishes and mixed well(final 15ml).										
08/28/98	11	100m/m,24	P16*	Subculture: Cells were harvested with 0.25% trypsin, suspended to 10ml and counted with Tatai at 10-folds dill. (319:41 1/2-block). Viability:88.6							:88.61 %			
	Whole viable cell numbers 2.55x10 ⁷ cells. Inoculated to new 24xculture dishes . An approximate. cell density:1.06x10 [^]							0^5 cells/m	, 1.80x10 [/]	⁴ cells/ci	1.8x10^4/cm	9.4x10^4/cm2		
08/31/98	14	100m/m,24	P16*	Took the photographs and medium ware changed.										
09/02/98	16			Cell freezing: Before	harvesting, took phot	ographs. Cells from 1	7 dishes were harvested, suspende	ed to 45ml n	nedium c	containing 59	%DMSO.			
	and counted cell number with Tatai at 10-folds dill. (521:12, 1-block). Whole viable cell numbers: 9.38×10^{47} cells.													
				Cell density: 2.08x10	⁶ cells/ml. Viability	: 96.06%. The suspens	sion was dispensed and frozen to 4	40-glass am	poules a	s distributioi	1.			2.1x10^6/amp
				The residual cells wer	re inoculated to 2-test	tubes and 1-blood aga	ar dish for sterility test.							
	100m/m,2 P1/* Furthermore, the residual cells were washed once, and inoculated to 100m/m dishes for sterilize test.								1.7x10^4/cm	2				
				Isoenzyme analysis sa	ample was made from	6xculture dishes.								
				A	wint Desses Number									
				Ampoules nad miss-p	print. Passege Numbe	er were lost asterisk								
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-													1	