

Cell Line Authentication Report

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1. Sample information:

Sample No.: “HUP-T4”

2. Methods:

- Genomic DNA was extracted from the cell pellets provided by the customer.
- Samples, together with positive and negative control were amplified using GenePrint System (Promega).
- Amplified products were processed using the ABI3730xl Genetic Analyzer.
- Data was analyzed by GeneMapper4.0 software and then compared with the ATCC, DSMZ or JCRB databases for reference matching.

3. Results:

STR profile

Marker	Sample				Database		
	Allele1	Allele2	Allele3	Allele4	Allele1	Allele2	Allele3
D5S818	10	11			10	11	
D13S317	8	9			8	8	
D7S820	8	11			8	11	
D16S539	10	11			10	11	
VWA	17	18			17	18	
TH01	9	9			9	9	
AMEL	X	X			X	X	
TPOX	8	8			8	8	
CSF1PO	10	12			10	12	
D12S391	15	19					
FGA	23	24	25				
D2S1338	18	19					
D21S11	29	32.2					
D18S51	18	18					
D8S1179	11	16					
D3S1358	15	15					
D6S1043	13	13					
PENTAE	18	19					
D19S433	13	13					
PENTAD	9	9					
D1S1656	15	16					

4. Conclusion

A. The STR results show that there are no four alleles on the main nine locus, there is certainly no cross contamination of human cells in this cell line.

B . The match percent between the sample and the STR database profile is **94.4%/100%**, the cell name is **HUP-T4**.

Similarity	Cell line	Source	Shared	D5S818	D7S820	D13S317	D16S539	vWA	TH01	TPOX	CSF1PO	Amelogenin
		Your query		10, 11	8, 11	8, 9	10, 11	17, 18	9, 9	8, 8	10, 12	X, X
94.4 %	HUP-T4	DSMZ: ACC-223	9	10, 11	8, 11	8, 8	10, 11	17, 18	9, 9	8, 8	10, 12	X, X

C. The TH01 Loci with 8,9 was supported by the paper(Cosmic-CLP=907286; ECACC=93121056; PubMed=25877200).

Note:

1. The STR profile data was compared with the ATCC, DSMZ or JCRB databases, if the cell line is not included in the three institutions, the results are not correct. More information you provided would be useful for Cell Line Authentication.
2. Based on the ANSI Standard, cell lines with 100% match are considered to be “identical”; cell lines with $\geq 80\%$ but less than 100% match are considered to be “related”.

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