

Cell Line Authentication Service

STR Profile Report

Sample Submitted By: NIH/NCI

Shouhui Yang

Email Address: shouhui.yang@nih.gov

ATCC Sales Order: SO0895664

FTA Barcode: STRB8048

Cell Line Designation: CFPAC-1

Date Sample Received: Monday, October 04, 2021

Report Date: Friday, October 15, 2021

Methodology: Seventeen short tandem repeat (STR) loci plus the gender determining locus, Amelogenin, were amplified

using the commercially available PowerPlex® 18D Kit from Promega. The cell line sample was processed using the ABI Prism® 3500xl Genetic Analyzer. Data were analyzed using GeneMapper® ID-X v1.2 software (Applied Biosystems). Appropriate positive and negative controls were run and confirmed for each

sample submitted.

Data Interpretation: Cell lines were authenticated using Short Tandem Repeat (STR) analysis as described in 2012 in ANSI

Standard (ASN-0002) Authentication of Human Cell Lines: Standardization of STR Profiling by the ATCC Standards Development Organization (SDO) and in Capes-Davis et al., Match criteria for human cell line

authentication: Where do we draw the line? Int. J. Cancer. 2012 Nov 8. doi: 10.1002/ijc.27931

ATCC performs STR Profiling following ISO 9001:2008 and ISO/IEC 17025:2005 quality standards.

There are no warranties with respect to the services or results supplied, express or implied, including, without limitation, any implied warranty of merchantability or fitness for a particular purpose. Neither ATCC nor Promega is liable for any damages or injuries resulting from receipt and/or improper, inappropriate, negligent or other wrongful use of the test results supplied, and/or from misidentification, misrepresentation, or lack of accuracy of those results. Your exclusive remedy against ATCC, Promega and those supplying materials used in the services for any losses or damage of any kind whatsoever, whether in contract, tort, or otherwise, shall be, at Promega's option, refund of the fee paid for such service or repeat of the service.

The ATCC trademark and trade name, any and all ATCC catalog numbers are trademarks of the American Type Culture Collection. PowerPlex is a registered trademark of Promega Corporation. Applied Biosystems, ABI Prism and GeneMapper are registered trademarks of Life Technologies Corporation.

Technical questions?

Ordering questions?

800-638-6597 or 703-365-2700 Fax 703-365-2750 Email: sales@atcc.org



ATCC Sales Order: SO0895664

	Test Results	s for Submitted	d Sample		A ⁻	TCC Referenc	e Database Pro	ofile
Locus		Query Pro	file: CFPAC-1		Database F		C-1; Pancreatic omo sapiens)	Carcinoma;
D3S1358	16							
TH01	8				8			
D21S11	30	31.2						
D18S51	12							
Penta_E	10	12						
D5S818	10	11			10	11		
D13S317	12				12			
D7S820	8	10			8	10		
D16S539	9	11			9	11		
CSF1PO	10				10			
Penta_D	11	13						
Amelogeni	n X	Y			Х	Y		
vWA	17				17			
D8S1179	11	15						
TPOX	8				8			
FGA	21	22						
D19S433	13	15						
D2S1338	18	23						
Number of sha	ared alleles between	query sample and	database profile	э:				13
Total number	of alleles in the datab	ase profile:						13
Percent match	h between the submitt	ed sample and th	e database prof	le:				100
The allele ma	tch algorithm compare	es the 8 core loci	plus amelogenin	only, even though	alleles from all loc	i will be reported	when available.	
please do no	ighlighted in grey (8 c I t publish the allele ca grams showing raw da	alls from all the S	Amelogenin) ca TR loci tested.	n be made public i	to verify cell identity	y. In order to prot	ect the identity of	the donor,
Cell lines with	of Test Results 80% match are consuthentication of related		ed; i.e., derived f	rom a common an	cestry. Cell lines wi	ith between a 55	% to 80% match r	equire further
The s	submitted sample pr	ofile is human,	but not a matc	h for any profile	in the ATCC STF	R database.		
	submitted profile is a ogenin): CRL-1918	an exact match	for the followin	g ATCC human	cell line(s) in the	ATCC STR da	tabase (8 core l	oci plus
The s	submitted profile is s	similar to the fol	lowing ATCC h	numan cell line(s):			
An S	TR profile could not	be generated.						
Additional (Comments:							

e-Signature, Technician:	gsykes 10/15/2021
e-Signature, Reviewer:	Bchase 10/15/2021

n/a





ATCC Sales Order: SO0895664

Addendum: Comparative Output from the ATCC STR Profile Database

% Match	ATCC® Cat. No.	Designation	D5S818	D13S317	D7S820	D16S539	vWA	TH01	AMEL	TPOX	CSF1P0
100	STRB8048	CFPAC-1	10,11	12	8,10	9,11	17	8	X,Y	8	10
100	CRL-1918	CFPAC-1; Pancreatic Carcinoma; Human (Homo sapiens)	10,11	12	8,10	9,11	17	8	X,Y	8	10

Definitions of terms used in this report:

Peak Area Difference (PAD):

Refers to a heterozygous peak imbalance.

Two alleles at a single locus should amplify in a similar manner; and therefore produce peaks of similar height and area. Peaks which are above threshold (50 rfu) but are not of similar area, within 50% of each other, are referred to as a PAD. Due to their nature cell lines do not amplify in the same manner as a sample taken from a fresh buccal swab. PAD is far more common in cell line samples.

Stutter:

A stutter peak is a small peak which occurs immediately before the true peak. It is defined as being a single repeat unit smaller than the true peak. The stutter peak should be less than 15% of the true peak. The stutter is caused by the polymerase.

+4 Peak:

A +4 is similar to a stutter but occurs immediately after the true peak. A stutter peak should be less than 5% for a homozygous and 10% for a heterozygous.

Below Threshold Peak(s):

Cell lines can produce unusual profiles and occasionally a peak will amplify poorly and be below threshold. Where we find a below threshold peak which we believe is valid we indicate it as a below threshold peak. Our cell line analysis criteria, Homozygous and Heterozygous peaks must be equal to or above the set height threshold for it to be considered a true peak.

Ladder/ Off Ladder Peak(s):

The allelic ladder consists of most or all known alleles in the population and allows for precise assignment of alleles. Those which do not align are termed 'off ladder.

Artifact:

A non-allelic product of the amplification process, an anomaly of the detection process, or a by-product of primer synthesis

Pull-up

A term used to describe when signal from one dye color channel produces artificial peaks in another, usually adjacent, color.

Spike:

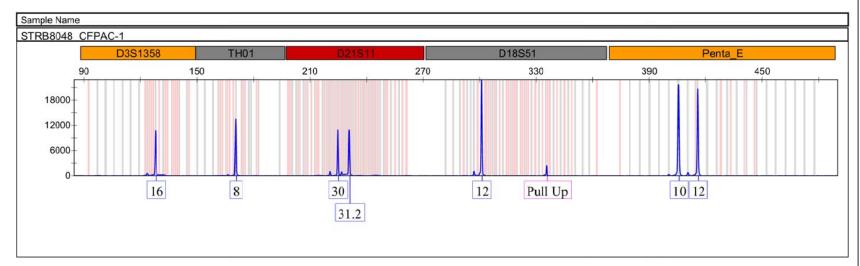
An extraneous peak resulting from dust, dried polymer, an air bubble, or an electrical surge.

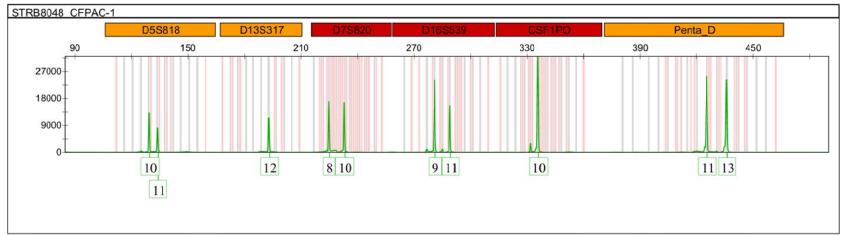
Dye blob:

Free dye not coupled to primer that can be injected into the capillary (A known and documented dye blob is often found at the D3S1358 locus.)



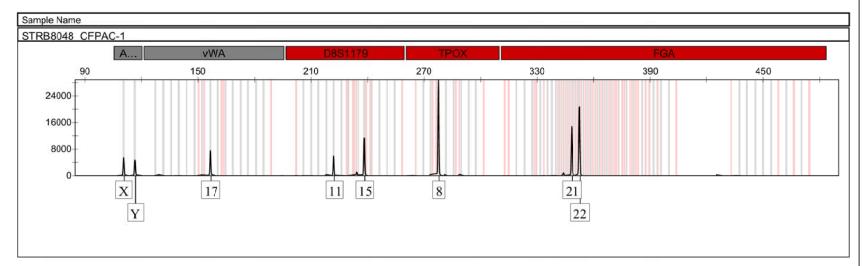


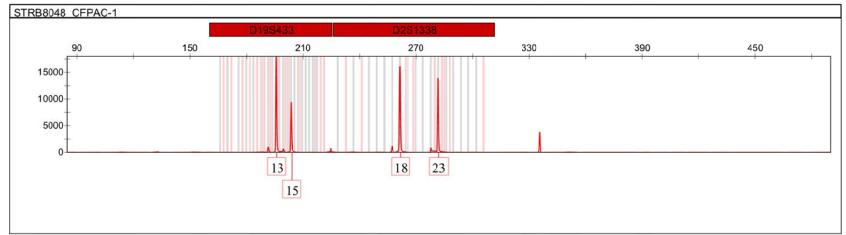














Cell Line Authentication Service

STR Profile Report

Sample Submitted By: National Institutes of Health

Shouhui Yang

Email Address: shouhui.yang@nih.gov

ATCC Sales Order: SO0895664

FTA Barcode: STRB8050

Cell Line Designation: Capan-2 P9

Date Sample Received: Monday, March 28, 2022

Report Date: Wednesday, March 30, 2022

Methodology: Seventeen short tandem repeat (STR) loci plus the gender determining locus, Amelogenin, were amplified

using the commercially available PowerPlex® 18D Kit from Promega. The cell line sample was processed using the ABI Prism® 3500xl Genetic Analyzer. Data were analyzed using GeneMapper® ID-X v1.2 software (Applied Biosystems). Appropriate positive and negative controls were run and confirmed for each

sample submitted.

Data Interpretation: Cell lines were authenticated using Short Tandem Repeat (STR) analysis as described in 2012 in ANSI

Standard (ASN-0002) Authentication of Human Cell Lines: Standardization of STR Profiling by the ATCC Standards Development Organization (SDO) and in Capes-Davis et al., Match criteria for human cell line

authentication: Where do we draw the line? Int. J. Cancer. 2012 Nov 8. doi: 10.1002/ijc.27931

ATCC performs STR Profiling following ISO 9001:2008 and ISO/IEC 17025:2005 quality standards.

There are no warranties with respect to the services or results supplied, express or implied, including, without limitation, any implied warranty of merchantability or fitness for a particular purpose. Neither ATCC nor Promega is liable for any damages or injuries resulting from receipt and/or improper, inappropriate, negligent or other wrongful use of the test results supplied, and/or from misidentification, misrepresentation, or lack of accuracy of those results. Your exclusive remedy against ATCC, Promega and those supplying materials used in the services for any losses or damage of any kind whatsoever, whether in contract, tort, or otherwise, shall be, at Promega's option, refund of the fee paid for such service or repeat of the service.

The ATCC trademark and trade name, any and all ATCC catalog numbers are trademarks of the American Type Culture Collection. PowerPlex is a registered trademark of Promega Corporation. Applied Biosystems, ABI Prism and GeneMapper are registered trademarks of Life Technologies Corporation.

Technical questions?

Ordering questions?

800-638-6597 or 703-365-2700 Fax 703-365-2750 Email: sales@atcc.org



ATCC Sales Order: SO0895664

	Test Result	s for Submitted	Sample		Α.	TCC Reference	ce Database Pro	ofile
Locus		Query Profile	e: Capan-2 P9		Database I		n-2; Pancreatic omo sapiens)	Carcinoma;
D3S1358	17	18						
TH01	9.3				9.3			
D21S11	31							
D18S51	13							
Penta_E	11							
D5S818	11	12			11	12		
D13S317	11	12			11	12		
D7S820	9	11			9	11		
D16S539	9	13			9	13		
CSF1PO	11	12			11	12		
Penta_D	13	15						
Amelogenin	Х				Х			
vWA	17				17			
D8S1179	12	13						
TPOX	8				8			
FGA	21	24						
D19S433	13	15						
D2S1338	19	25						
Number of shared	l alleles between	query sample and	database profile:	:	•		•	14
Total number of a	lleles in the datab	ase profile:						14
		ted sample and the						100
The allele match a	algorithm compar	es the 8 core loci p	lus amelogenin d	only, even though	alleles from all loc	i will be reported	d when available.	
	iblish the allele c	core STR loci plus A alls from all the ST ata are attached.		n be made public t	o verify cell identity	/. In order to pro	tect the identity of	the donor,
Explanation of To Cell lines with 80% profiling for auther	% match are cons	idered to be related dness.	d; i.e., derived fro	om a common and	cestry. Cell lines w	ith between a 55	5% to 80% match r	require further
The subr	nitted sample p	rofile is human, b	ut not a match	for any profile i	n the ATCC STF	R database.		
	nitted profile is a nin): HTB-80	an exact match fo	or the following	ATCC human	cell line(s) in the	ATCC STR da	atabase (8 core l	loci plus
The subr	nitted profile is	similar to the follo	owing ATCC hu	uman cell line(s)):			
An STR	orofile could not	be generated.						
Additional Con	nments:							

e-Signature, Technician:	Ichamp 3/30/2022
e-Signature, Reviewer:	Bchase 3/30/2022

n/a





ATCC Sales Order: SO0895664

Addendum: Comparative Output from the ATCC STR Profile Database

% Match	ATCC® Cat. No.	Designation	D5S818	D13S317	D7S820	D16S539	vWA	TH01	AMEL	TPOX	CSF1PO
100	STRB8050	Capan-2 P9	11,12	11,12	9,11	9,13	17	9.3	Х	8	11,12
100	HTB-80	Capan-2; Pancreatic Carcinoma; Human (Homo sapiens)	11,12	11,12	9,11	9,13	17	9.3	Х	8	11,12

Definitions of terms used in this report:

Peak Area Difference (PAD):

Refers to a heterozygous peak imbalance.

Two alleles at a single locus should amplify in a similar manner; and therefore produce peaks of similar height and area. Peaks which are above threshold (50 rfu) but are not of similar area, within 50% of each other, are referred to as a PAD. Due to their nature cell lines do not amplify in the same manner as a sample taken from a fresh buccal swab. PAD is far more common in cell line samples.

Stutter

A stutter peak is a small peak which occurs immediately before the true peak. It is defined as being a single repeat unit smaller than the true peak. The stutter peak should be less than 15% of the true peak. The stutter is caused by the polymerase.

+4 Peak:

A +4 is similar to a stutter but occurs immediately after the true peak. A stutter peak should be less than 5% for a homozygous and 10% for a heterozygous.

Below Threshold Peak(s):

Cell lines can produce unusual profiles and occasionally a peak will amplify poorly and be below threshold. Where we find a below threshold peak which we believe is valid we indicate it as a below threshold peak. Our cell line analysis criteria, Homozygous and Heterozygous peaks must be equal to or above the set height threshold for it to be considered a true peak.

Ladder/ Off Ladder Peak(s):

The allelic ladder consists of most or all known alleles in the population and allows for precise assignment of alleles. Those which do not align are termed 'off ladder.

Artifact:

A non-allelic product of the amplification process, an anomaly of the detection process, or a by-product of primer synthesis

Pull-up:

A term used to describe when signal from one dye color channel produces artificial peaks in another, usually adjacent, color.

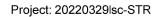
Spike

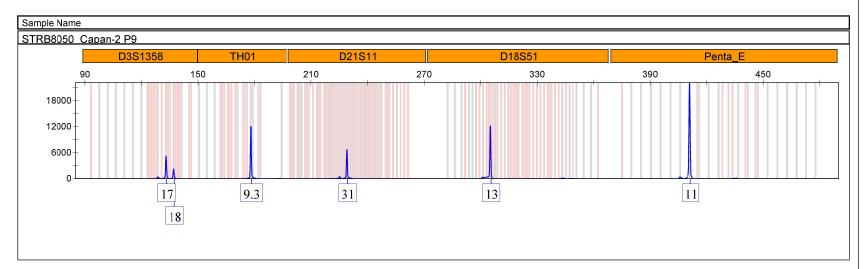
An extraneous peak resulting from dust, dried polymer, an air bubble, or an electrical surge.

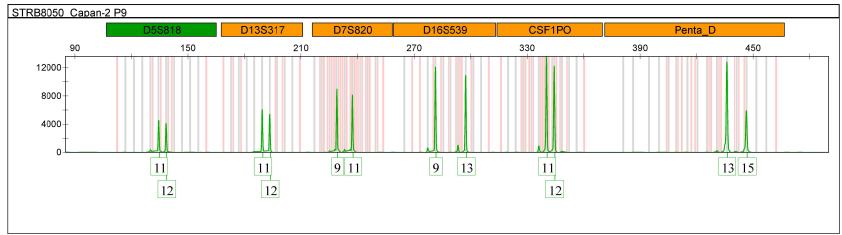
Dve blob:

Free dye not coupled to primer that can be injected into the capillary (A known and documented dye blob is often found at the D3S1358 locus.)

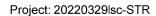


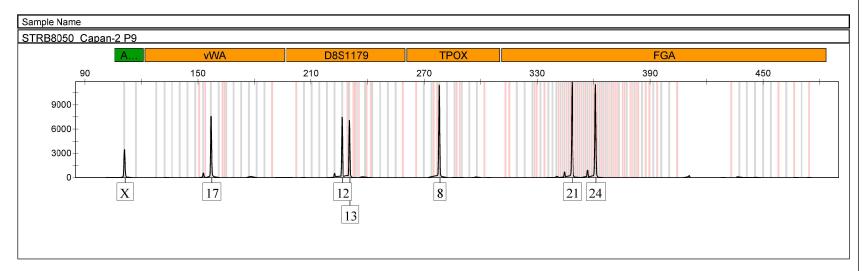


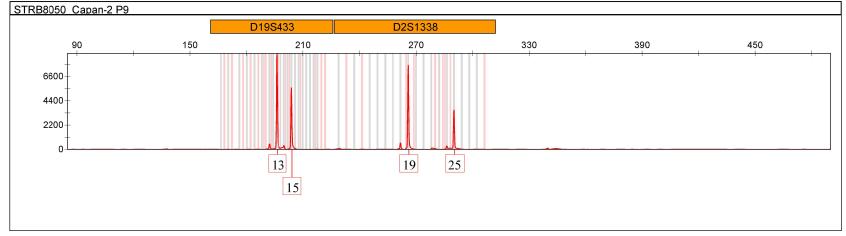














Cell Line Authentication Service

STR Profile Report

Sample Submitted By: National Institutes of Health

Shouhui Yang

Email Address: shouhui.yang@nih.gov

ATCC Sales Order: SO0895664

FTA Barcode: STRB8042

Cell Line Designation: ASPC-1

Date Sample Received: Monday, March 28, 2022

Report Date: Wednesday, March 30, 2022

Methodology: Seventeen short tandem repeat (STR) loci plus the gender determining locus, Amelogenin, were amplified

using the commercially available PowerPlex® 18D Kit from Promega. The cell line sample was processed using the ABI Prism® 3500xl Genetic Analyzer. Data were analyzed using GeneMapper® ID-X v1.2 software (Applied Biosystems). Appropriate positive and negative controls were run and confirmed for each

sample submitted.

Data Interpretation: Cell lines were authenticated using Short Tandem Repeat (STR) analysis as described in 2012 in ANSI

Standard (ASN-0002) Authentication of Human Cell Lines: Standardization of STR Profiling by the ATCC Standards Development Organization (SDO) and in Capes-Davis et al., Match criteria for human cell line

authentication: Where do we draw the line? Int. J. Cancer. 2012 Nov 8. doi: 10.1002/ijc.27931

ATCC performs STR Profiling following ISO 9001:2008 and ISO/IEC 17025:2005 quality standards.

There are no warranties with respect to the services or results supplied, express or implied, including, without limitation, any implied warranty of merchantability or fitness for a particular purpose. Neither ATCC nor Promega is liable for any damages or injuries resulting from receipt and/or improper, inappropriate, negligent or other wrongful use of the test results supplied, and/or from misidentification, misrepresentation, or lack of accuracy of those results. Your exclusive remedy against ATCC, Promega and those supplying materials used in the services for any losses or damage of any kind whatsoever, whether in contract, tort, or otherwise, shall be, at Promega's option, refund of the fee paid for such service or repeat of the service.

The ATCC trademark and trade name, any and all ATCC catalog numbers are trademarks of the American Type Culture Collection. PowerPlex is a registered trademark of Promega Corporation. Applied Biosystems, ABI Prism and GeneMapper are registered trademarks of Life Technologies Corporation.

Technical questions?

Ordering questions?

800-638-6597 or 703-365-2700 Fax 703-365-2750 Email: sales@atcc.org



FTA Barcode: STRB8042
ATCC Sales Order: SO0895664

	Test Results for Submitted Sample						ATCC Reference Database Profile				
Locus		Query Pro	ofile: ASPC-1		Database Pr	Database Profile: AsPC-1; Pancreatic Cancer; Human (Homo sapiens)					
D3S1358	16										
TH01	7	9.3			7	9.3					
D21S11	28	30									
D18S51	18										
Penta_E	5	12									
D5S818	12				12						
D13S317	9	12			9	12					
D7S820	12	13			12	13					
D16S539	11				11						
CSF1PO	10	13			10	13					
Penta_D	9	12									
Amelogenin	Х				Х						
vWA	17				17						
D8S1179	13	15									
TPOX	8	10			8	10					
FGA	24										
D19S433	14										
D2S1338	22	23									
Number of shared	alleles between	query sample an	d database profi	le:				14			
Total number of al	leles in the datab	ase profile:						14			
Percent match bet								100			
The allele match a	algorithm compare	es the 8 core loci	plus amelogenii	n only, even though a	alleles from all loc	i will be reported	l when available.				
NOTE: Loci highli please do not pu Electropherogram	blish the allele ca	alls from all the S		an be made public to	verify cell identity	v. In order to pro	tect the identity o	f the donor,			
profiling for auther	6 match are consintication of related	dness.		from a common anco	•		% to 80% match	require further			

	The submitted sample profile is human, but not a match for any profile in the ATCC STR database.
X	The submitted profile is an exact match for the following ATCC human cell line(s) in the ATCC STR database (8 core loci plus Amelogenin): CRL-1682
	The submitted profile is similar to the following ATCC human cell line(s):
	An STR profile could not be generated.

Additional Comments:

n/a

e-Signature, Technician:	Ichamp 3/30/2022
e-Signature, Reviewer:	Bchase 3/30/2022





ATCC Sales Order: SO0895664

Addendum: Comparative Output from the ATCC STR Profile Database

% Match	ATCC® Cat. No.	Designation	D5S818	D13S317	D7S820	D16S539	vWA	TH01	AMEL	TPOX	CSF1PO
100	STRB8042	ASPC-1	12	9,12	12,13	11	17	7,9.3	Х	8,10	10,13
100	CRL-1682	AsPC-1; Pancreatic Cancer; Human (Homo sapiens)	12	9,12	12,13	11	17	7,9.3	Х	8,10	10,13

Definitions of terms used in this report:

Peak Area Difference (PAD):

Refers to a heterozygous peak imbalance.

Two alleles at a single locus should amplify in a similar manner; and therefore produce peaks of similar height and area. Peaks which are above threshold (50 rfu) but are not of similar area, within 50% of each other, are referred to as a PAD. Due to their nature cell lines do not amplify in the same manner as a sample taken from a fresh buccal swab. PAD is far more common in cell line samples.

Stutter

A stutter peak is a small peak which occurs immediately before the true peak. It is defined as being a single repeat unit smaller than the true peak. The stutter peak should be less than 15% of the true peak. The stutter is caused by the polymerase.

+4 Peak

A +4 is similar to a stutter but occurs immediately after the true peak. A stutter peak should be less than 5% for a homozygous and 10% for a heterozygous.

Below Threshold Peak(s):

Cell lines can produce unusual profiles and occasionally a peak will amplify poorly and be below threshold. Where we find a below threshold peak which we believe is valid we indicate it as a below threshold peak. Our cell line analysis criteria, Homozygous and Heterozygous peaks must be equal to or above the set height threshold for it to be considered a true peak.

Ladder/ Off Ladder Peak(s):

The allelic ladder consists of most or all known alleles in the population and allows for precise assignment of alleles. Those which do not align are termed 'off ladder.

Artifact:

A non-allelic product of the amplification process, an anomaly of the detection process, or a by-product of primer synthesis

Pull-up:

A term used to describe when signal from one dye color channel produces artificial peaks in another, usually adjacent, color.

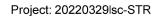
Spike

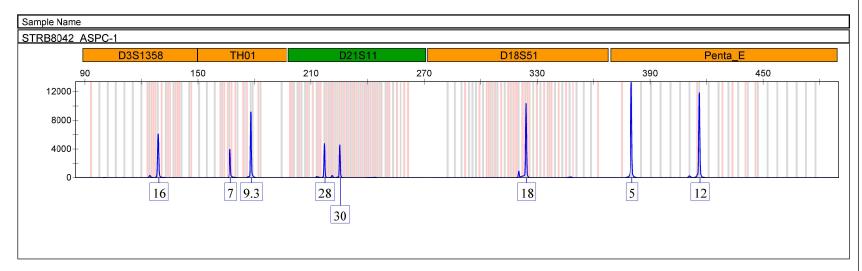
An extraneous peak resulting from dust, dried polymer, an air bubble, or an electrical surge.

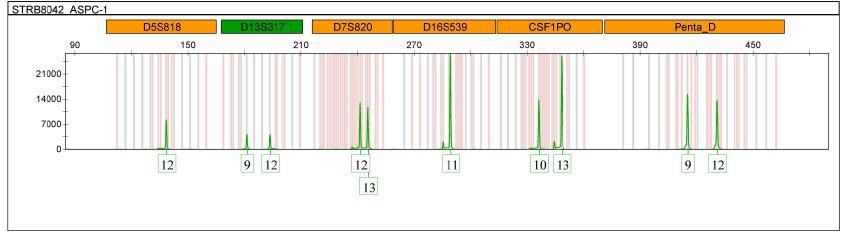
Dye blob:

Free dye not coupled to primer that can be injected into the capillary (A known and documented dye blob is often found at the D3S1358 locus.)

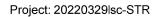


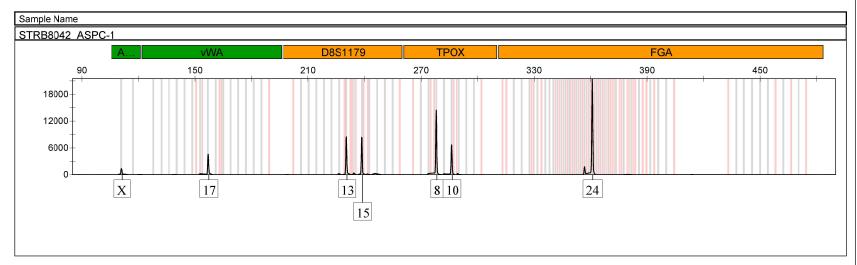


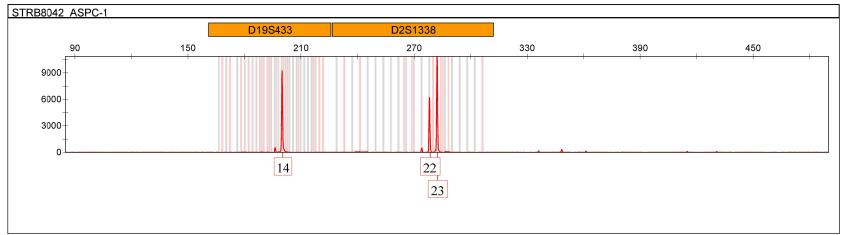














Frederick National Laboratory for Cancer Research

DATE: July 10th, 2015

TO: Perwez Hussain, Ph.D.

THROUGH: Carissa Grose

Group Leader, Cloning and Nucleic Acids

Protein Expression Laboratory

FROM: Allison Meade

Research Associate I

Molecular Detection Group Protein Expression Laboratory

SUBJECT: Short Tandem Repeat Analysis (STR) CSAS 17674

SAMPLES: 7 cell lines

MDG received the above DNA samples from your laboratory for STR analysis. DNA was extracted from each sample and was amplified by PCR using the AmpF ℓ STR® Identifiler PCR amplification kit. This is a STR multiplex assay that amplifies 15 tetranucleotide repeat loci and the Amelogenin gender determination marker in a single PCR amplification. Sample PCR amplicon, as well as positive control amplicon, were then denatured at 95°C and the fragments analyzed on an Applied Biosystems 3130xl genetic analyzer in Hi-Di formamide with a size standard. Fragments were labeled and identified using GeneMapper 4.0.

The resulting allelic distributions of the samples are listed in the attached table. The numerical values represent the number of tetramer repeats for each allele of the 16 target loci. The allelic distribution of the positive control is also provided as a reference.

RESULTS:

	Panc-1	MiaPaCa-2	Capan-1	Capan-2	CFPAC-1	Pacn10.05	ASPC-1	Test Positive Control	Expected Positive Control
D8S1179	14, 15	16, 16	14,16	12, 13	11, 15	13, 14	13, 15	13, 13	13, 13
D21S11	28, 28	29, 31.2	28, 30	31, 31	30, 31.2	30, 30	28, 30	30, 30	30, 30
D7S820	8, 10	12, 13	10, 11	9, 11	8, 10	8, 9	12, 13	10, 11	10, 11
CSF1P0	10, 12	10, 10	11, 11	11, 12	10, 10	12, 12	10, 13	10, 12	10, 12
D3S1358	17, 17	16, 16	15, 15	17, 18	16, 16	14, 14	16, 16	14, 15	14, 15
TH01	7, 8	9, 10	6, 6	9.3, 9.3	8, 8	6, 9.3	7, 9.3	8, 9.3	8, 9.3
D13S317	11, 11	12, 13	9, 9	11, 12	12, 12	12, 12	9, 12	11, 11	11, 11
D16S539	11, 11	10, 13	13, 14	9, 13	9, 11	9, 12	11, 11	11, 12	11, 12
D2S1338	23, 24	25, 25	20, 24	19, 25	18, 23	17, 18	22, 23	19, 23	19, 23
D19S433	11, 16	15, 15	14, 15	13, 15	13, 15	13, 14	14, 14	14, 15	14, 15
vWA	15, 15	15, 15	16, 16	17, 17	17, 17	16, 16	17, 17	17, 18	17, 18
TPOX	8, 11	9, 9	8, 11	8, 8	8, 8	11, 11	8, 10	8, 8	8, 8
D18S51	12, 12	12, 12	12, 12	13, 13	12, 12	15, 15	18, 18	15, 19	15, 19
D5S818	11, 13	12, 13	11, 11	11, 12	10, 11	13, 13	12, 12	11,11	11,11
FGA	21, 21	22, 22	24, 24	21, 24	21, 22	20, 20	24, 24	23, 24	23, 24
Amelogenin	X, X	X, X	X, X	X, X	X, Y	X, X	X, X	X, X	X, X

If you have any questions, please feel free to contact me.

Allison Meade Carissa Grose
ATRF, C2015 ATRF, C2026
8560 Progress Dr. 8560 Progress Dr.
Frederick, MD 21701 Frederick, MD 21701
Phone: 301-846-1888 Phone: 301-360-3427
Fax: 301-846-6289 Fax: 301-846-6289

DISCLAIMER: Characterization of a cell line by examining differences in the number of short tandem repeats (STR) is a technology originally developed for use as a forensic tool. While this technology has been applied for the identification and characterization of human tumor cell lines, there are some technical issues to be considered in the interpretation of the data obtained. Issues that may be encountered are polyploidy and increased mutation rates (loss/gain of number of STR) due to continuous cell culture¹.

References:

¹ Parson W., Kirchebner R., Mühlmann R., Renner K., Kofler A., Schmidt S., Kofler R. *(2005) FASEB J.* 19, 434-436.