SAMPLE INTEGRITY



# Sample Integrity Solutions

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#### 1 IN EVERY 200 SAMPLES IS MIXED UP1

It is estimated that 1 in every 200 samples is mixed up. Many of these mistakes originate outside the lab and traditional quality checks are not enough to detect them. Sample mix-ups lead to incorrect results and can damage a lab's reputation.

#### DOWNSTREAM MOLECULAR ANALYSIS IS UNSUCCESSFUL IN 22.5% OF CASES<sup>2</sup>

Many samples fail downstream analysis due to low quantity or poor-quality DNA; meaning wasted time and no results. DNA quality measurements based on absorbance detect nucleic acid rather than amplifiable DNA fragments and often provide inaccurate results.



- FOR MOLECULAR DIAGNOSTIC LABS Prevent sample mix-ups and ensure DNA is of sufficient quality before expensive downstream analysis.
- **FOR BIOBANKS** Verify sample origin to prevent erroneous study findings.
- FOR TOXICOLOGY LABS Detect fraudulent samples by matching submissions to a reference genetic fingerprint.

# Three sample quality measurements from a single assay:

#### SAMPLE IDENTIFICATION



Verify sample identity by comparing genetic fingerprints. Prevent mix-ups and mismatches even if they originated outside of the lab.

#### DNA QUANTITY

# Is there enough DNA?

Understand how much DNA is present within a sample. Ensure there is enough for all applications.

#### **DNA QUALITY**



Decide if the DNA in a sample is suitable for downstream analysis by measuring the number of amplifiable DNA fragments rather than free nucleotides and other contaminants.



## **Panel Selection**

Each of the iPLEX<sup>®</sup> Pro Sample Integrity panels enable rapid and highly accurate sample identification and DNA quality measurements. SNPs and gender markers are used to generate a sample's unique genetic fingerprint. The automated reporter compares this to a reference profile and generates an easy to interpret Match or Mismatch result. In the same reaction, amplification is compared to built-in controls to determine how many amplifiable DNA copies are present in the sample.

These pre-designed panels are available for on-demand ordering to get you testing samples quickly. Each leverages the proven iPLEX Pro chemistry and can be processed in the same run with other applications.

#### **IPLEX PRO SAMPLE ID PANEL**

This panel determines a sample's identity, DNA quality and DNA quantity. It targets 44 SNPs with high minor allele frequency across major HapMap populations and 3 gender markers.

#### **IPLEX PRO EXOME QC PANEL**

This panel determines a sample's identity, DNA quality and DNA quantity. To screen for sample degradation, it also provides the number of intact, amplifiable DNA copies across a size range of 100, 200, 300, 400 and 500 nucleotides. It targets 21 SNPs in exonic regions of the genome and 3 gender markers.

#### Comparing DNA eliminates misidentification:

#### COMPARISON: Non-Tumor vs. Non-Tumor

Automated algorithm assigns a match score based on the SNP results of two samples. The software reporter will clearly identify matched and mismatched samples

#### COMPARISON: Tumor vs. Non-Tumor

Algorithm is adjusted to account for potential loss of heterozygosity (LOH) in tumor specimens. The software reporter will clearly identify matched and mismatched samples

# SAMPLE INTEGRITY

Sample Integrity Panel Comparison					
	iPLEX Pro Sample ID Panel	iPLEX Pro Exome QC Panel			
Sample Identification	44 SNPs, 3 gender markers	21 SNPs, 3 gender markers			
DNA Quantity	5 controls	25 controls			
DNA Quality	Assay measures amplifiable copies of DNA	Assay measures amplifiable copies of DNA at several fragment lengths			
DNA Input	5 ng+	5 ng+			
Genomic Region Targeted	Intronic & Exonic Regions	Exonic Regions			
Automated Reporting	Included	Included			

iPLEX Pro Sample ID Panel - SNPs & Genes					
albumin_1	rs11781516	rs1994997	rs3819854		
albumin_10	rs13050660	rs2010253	rs717302		
albumin_5	rs1335873	rs2040411	rs727811		
albumin_8	rs1357617	rs2046361	rs729172		
albumin_9	rs1360288	rs2056277	rs740910		
AMEL_XY	rs136337	rs2076848	rs8037429		
ARSD_XY	rs1382387	rs214054	rs826472		
TGIF2L_XY	rs1413212	rs2247221	rs876724		
rs1005533	rs1454361	rs2518968	rs891700		
rs1024116	rs1463729	rs251934	rs901398		
rs1028528	rs1468118	rs2714854	rs914165		
rs10495407	rs1493232	rs2831700	rs9583190		
rs10771010	rs1982986	rs354439	rs964681		

iPLEX PI	ro Exome QC I	Panel - SNPs	& Genes
POLR2A	TPT1	rs10495563	rs4478844
POLR2A	VIM	rs1065457	rs586421
TBP	POLR2A	rs1127379	rs6420424
ALB	RPL27A	rs11998387	rs6977125
IPO8	HBS1L	rs1200349	rs7653897
POLR2A	TPT1	rs12594531	rs773901
POLR2A	IPO8	rs1344	rs9131
TBP	PDCD2	rs17548783	AHSP
IPO8	TBP	rs2246209	AMEL
HBG2	PPIA	rs2273171	ARSD
GUSB	HBS1L	rs2301771	
AHSP	PDCD2	rs3743165	
RPL27A	rs1045728	rs3884596	

# iPLEX Pro Sample Integrity Reporting Software

The iPLEX Pro Sample Integrity panels are accompanied by reporting software that makes sample identity and DNA quality determination quick and easy. Automated reports clearly show sample information, gender, the number of amplifiable DNA copies and a match/mismatch analysis.

#### SUMMARY REPORT

The summary report provides a quick overview, highlighting any unexpected sample matches or mismatches. It also lists samples that may have failed quality control due to poor quality or low quantity DNA.

Type of R Panel Ve Date of R Sar Compar Elapsed	eport: Local irsion: 4 eport: May 29 <sup>th</sup> , 2018 at 10: nples: 102 isons: 3570 Time: 5 seconds	42:07			
Plates Used	to Generate Report				
Plate ID	Exp	eriment ID	Experiment #	Samples	
Unexpected	Results				
Mismatches	Non-Tumor vs. Non-Tumor	Q			
Matches	Non-Tumor vs. Non-Tumor	0			
Matches	Non-Tumor vs. Tumor	Ď.			
					_

- RUN INFORMATION Displays details of the run including name, date and number of samples analyzed.
- QC STATUS Lists samples that may have failed QC due to poor quality or low quantity DNA
- MATCH / MISMATCH INFORMATION Clearly identifies samples did not agree with the prescribed matching criteria

#### MATCH REPORT

The Match Report lists identifying information, SNP call and matching score for two samples involved in a comparison.

- MATCH / MISMATCH INFORMATION Clearly identifies samples did not agree with the prescribed matching criteria
- HISTORICAL DATABASE Performs a global search across the entire database to identify sample relationships.
- LOCAL COMPARISON Allows for a quick analysis of sample relationships within an individual run.



### SAMPLE INTEGRITY



#### **SAMPLE REPORT**

Each Sample Report contains identifying information, DNA copy number, gender, SNP calls and Unexpected Match/Mismatches.

#### **QC REPORT**

Both the iPLEX Sample ID and Exome QC panels use in-assay control to determine the number of amplifiable DNA copies present in a sample. The Exome QC panel provides this quality analysis at several DNA fragment lengths and generates a sample degradation curve.





#### DNA Quality Data

 Amplifiable Copy Number:
 4334 ± 549
 4188
 3737
 3964



4706 5076

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#### ASK ABOUT OUR OTHER APPLICATIONS

The MassARRAY® is a versatile genetic analysis tool and not limited to a single application. Ask for information regarding our other applications including Oncology, Pharmacogenetics and Hereditary Genetic Testing.

#### **ORDERING INFORMATION**

Catalog No.	Item	Chemistry	Chip format	# of Samples
13117F	Complete iPLEX Sample ID Panel Set CPM	iPLEX Master Mix	96	960
13117	Complete iPLEX Sample ID Panel Set	iPLEX Master Mix	96	960
13118	Complete iPLEX Sample ID Panel Set	iPLEX Master Mix	384	3,840
25093F	Complete iPLEX Pro Sample ID Panel Set CPM	iPLEX Pro	96	960
25093	Complete iPLEX Pro Sample ID Panel Set	iPLEX Pro	96	960
25094	Complete iPLEX Pro Sample ID Panel Set	iPLEX Pro	384	768
25095	Complete iPLEX Pro Sample ID Panel Set	iPLEX Pro	96	3,840
13175F	Complete Exome QC Panel Set CPM	iPLEX Pro	96	960
13175	Complete Exome QC Panel Set	iPLEX Pro	96	960
13176	Complete Exome QC Panel Set	iPLEX Pro	384	3,840

#### References

Marberger M, McConnell JD, Fowler I, et al. Biopsy Misidentification Identified by DNA Profiling in a Large Multicenter Trial. Journal of Clinical Oncology. 2011;29(13):1744-1749. doi:10.1200/JCO.2010.32.1646.

2. Hussam Al-Kateb et al. Identification of major factors associated with failed clinical molecular oncology testing performed by next generation sequencing (NGS). Molecular Oncology, 9 (2015) 1737-1743.

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