



QIAcuity[®] Digital PCR System

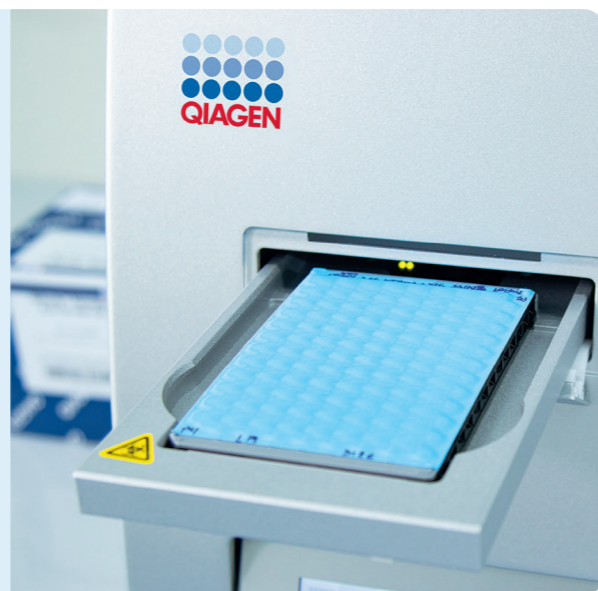
Fast. Scalable. Reliable.



The Magic is Inside

It's in the seamless integration of all digital PCR (dPCR) workflow components into an all-in-one walkaway instrument, delivering the speed and throughput every laboratory needs.

It's in the microfluidic nanoplate technology that puts every run ahead of the curve with its precision and sensitivity.



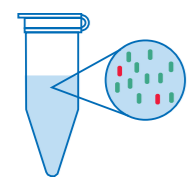
Got a needle in a haystack problem?

Digital PCR holds the answer.

You know the struggles of performing sensitive research applications to identify faint genetic event against a strong background, especially when the positives are lost in a dense pool of negatives. Finding that rare allele or mutant sequence is a typical needle in a haystack problem. This is where digital measurement comes in handy.

Digital PCR (dPCR) is a nucleic acid quantification technique that works by dividing a bulk qPCR-like

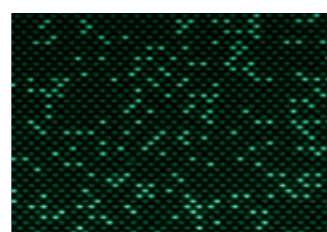
reaction mixture into numerous individual reactions called partitions and then measuring the endpoint fluorescence of each partition to determine the presence (1) or absence (0) of the target. This makes digital PCR less reliant on the kinetics of the PCR reaction and eliminates the need for standard curves as in qPCR. Statistical methods (Poisson law) are then used to calculate the absolute concentration of the target based on the number of positive and negative partitions.



Red - Target and Green - Background (gDNA, cDNA; primers/probes; master mix)



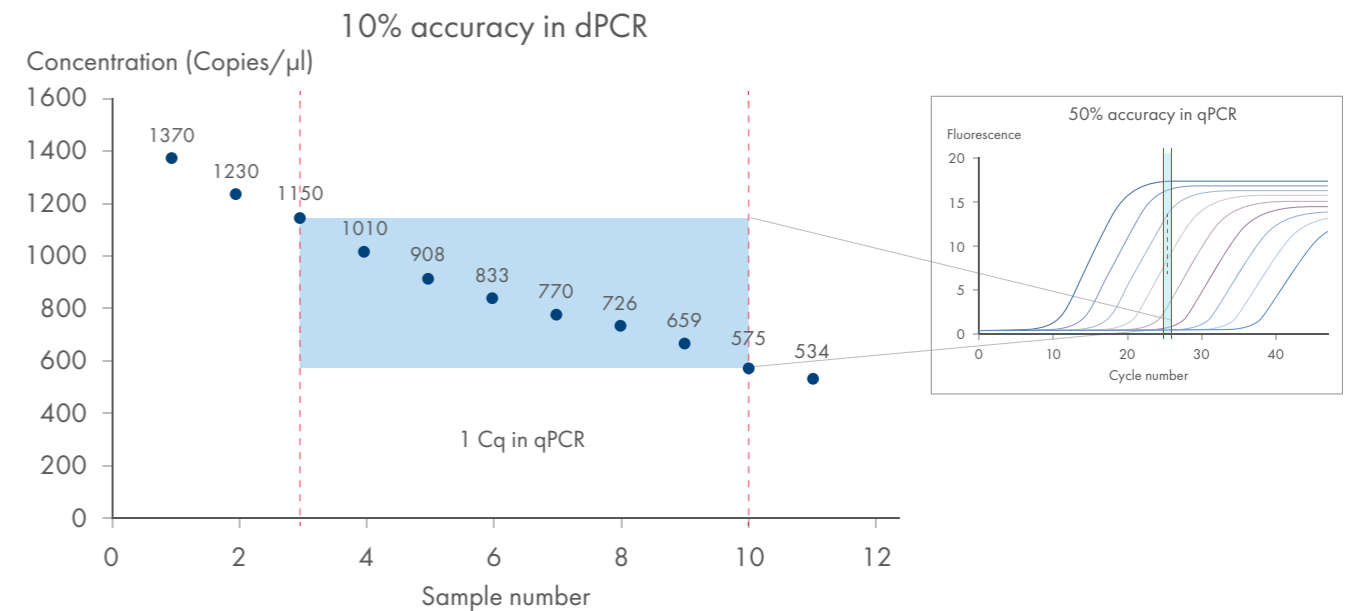
Random distribution of molecules into partitions



Absolute quantification: Copies/ μ l

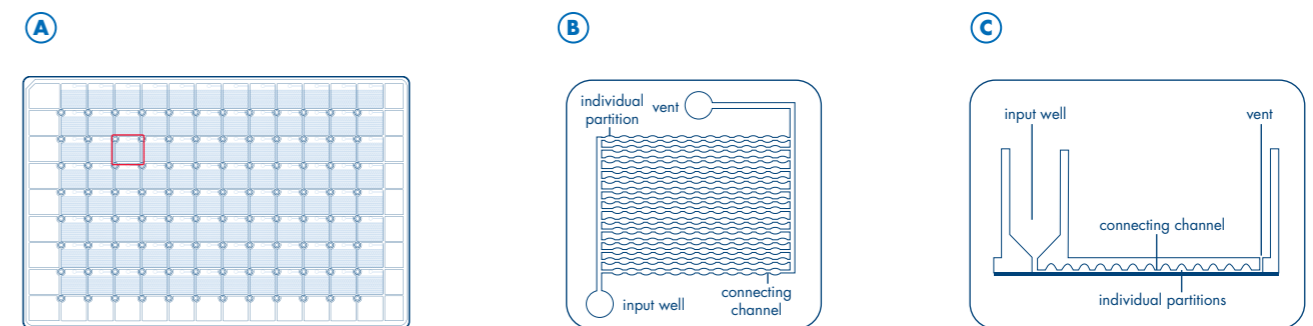
Absolute quantification provides reproducible data that can be more easily compared between laboratories. Further, due to the linear response

of the technology, dPCR offers a more precise measurement than qPCR and makes it surprisingly easy to detect the positives.



No droplets. No chips. No crystals. Digital PCR in nanoplates.

The QIAcuity Digital PCR System uses a microfluidic nanoplate technology to overcome challenges with inconsistent droplet generation, complex workflow, slow droplet readout, and limitations concerning the uncertainty of assays.



A Nanoplate with 96 well B Single well detail C Cross section view of the partitions

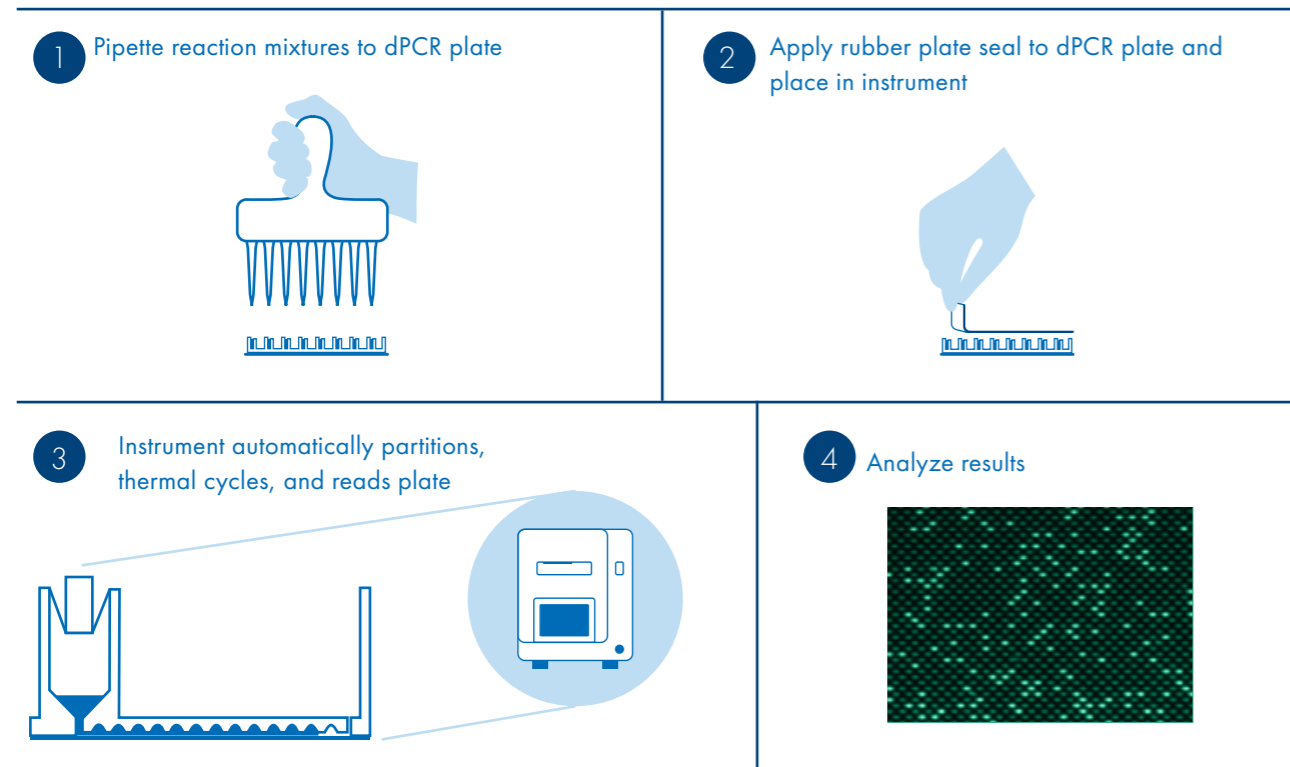
5 reasons why

- Fixed partitions prevent variation in size and coalescence, maximizing consistency
- Sealed nanoplates eliminate the risk of contamination
- Simultaneous reading of all partitions/well allows quicker time-to-result
- qPCR-like plates provide a more familiar workflow, improving ease of use
- Plates are amenable to front-end automation (e.g., on the QIAgility), minimizing hands-on steps

A simple and rapid workflow

The nanoplate-based QIAcuity Digital PCR System provides a qPCR-like workflow, in which sample preparation includes the transfer of diluted samples and the addition of master mix, probes and primers to an 8-, 24- or 96-well nanoplate. The system then

automates a fully integrated dPCR workflow – partitioning, thermocycling and imaging – enabling walk-away operation and delivering results in about two hours.



➔ Explore a virtual workflow demo: www.qiagen.com/qiacuity-demo

QIAcuity Digital PCR System

Features and benefits

With a fully integrated design, walk-away automation, ease of use, advanced multiplexing, scalable instrument and flexible plate configuration for high throughput and highly sensitive detection, the QIAcuity system can displace qPCR, ddPCR and existing dPCR systems as the method of choice for quantification of nucleic acid targets.

• Scalable design

The QIAcuity system comes in scalable instrument configurations with a single thermal cycler and capacity to run up to 4 plates or a dual thermal cycler and capacity to run up to 8 plates.

• The highest throughput digital PCR system ever

An 8-plate capacity allows up to 1248 samples to be analyzed in a single workday using a 96-well nanoplate.

• Ultra-high multiplexing

Up to 6 channels (including one reference channel) can be configured for multiplex quantification of up to 5 target DNA or RNA molecules in a given assay, saving time and reagents.

• Fully automated digital PCR

The QIAcuity system integrates reaction partitioning, thermal cycling and imaging into a single fully automated instrument that takes users from sample to result of up to 96 samples in 2 hours and up to 768 samples in 5 hours.

• Simplified transition from qPCR

The QIAcuity system is compatible with qPCR detection chemistries such as hydrolysis probes and EvaGreen dye, simplifying the transition from qPCR assays.



QIAcuity instruments

	QIAcuity One	QIAcuity Four	QIAcuity Eight
Plates processed	1	4	8
Detection channels (multiplexing)	2 or 5	5	5
Thermocycler(s)	1	1	2
Time to result	Approx. 2 h	First plate approx. 2 h Every ~60 min a following plate	First plate approx. 2 h Every ~30 min a following plate
Throughput (samples processed in a work day)	Up to 384 (96-well) Up to 96 (24-well)	Up to 672 (96-well) Up to 168 (24-well)	Up to 1248 (96-well) Up to 312 (24-well)

Detection channels and fluorophores	
Detection channels	Recommended dyes
Green	FAM
Yellow	VIC, HEX
Orange	TAMRA
Red	ROX
Crimson	Cy5

You see the most with the QIAcuity Nanoplate 26K

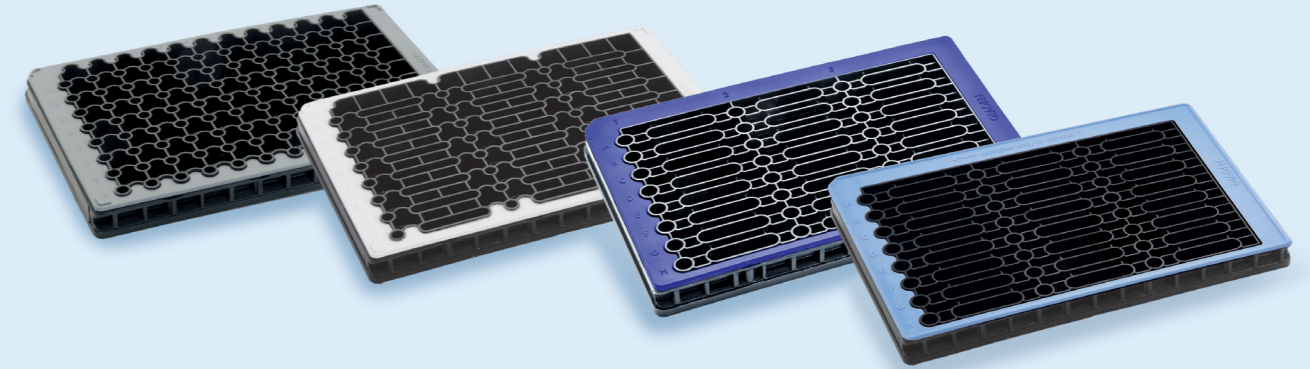
Finding a rare event (<10–20 copies per reaction) means confirming the presence or absence of a signal and not just precise quantification. Loading volume is critical when applied to samples with rare targets.

Imagine analyzing a DNA eluate containing a very rare event of

0.1 cp/μl, i.e., 3 cp in a 30 μl eluate. How many copies can you detect by digital PCR?

The QIAcuity Nanoplate 26K allows more loading, allowing you to see the most compared to that offered by any other dPCR method, including comparable plate methods.

QIAcuity Nanoplates



Features and benefits

The QIAcuity system offers distinct nanoplate configurations with flexible sample formats that accommodate a wide range of throughput and sensitivity requirements.

Plate type	Samples/plate	Partitions/well	Input volume	Key applications
Nanoplate 26K 8-well	8	approx. 26,000	40 μl	Rare mutation detection, liquid biopsy, pathogen detection
Nanoplate 26K 24-well	24	approx. 26,000	40 μl	Rare mutation detection, liquid biopsy, pathogen detection
Nanoplate 8.5K 24-well	24	approx. 8500	12 μl	CNV detection, NGS library quantification
Nanoplate 8.5K 96-well	96	approx. 8500	12 μl	CNV detection, NGS library quantification

	QIAGEN Nanoplate 26K	MAP16 dPCR plate	ddPCR	
	40 μl	9 μl	20 μl	Reaction volume
	26 μl	6 μl	13 μl	Possible sample volume from the 30 μl eluate (assuming 4x master mix and 10x assay)
	20 μl	8.5 μl	16 μl (assuming 16K droplets)	Volume analyzed
3 cp in the eluate	2.6 cp	0.6 cp	1.3 cp	Copies transferred
	1.3 cp	0.57 cp	1.04 cp	Copies analyzed
10 cp in the eluate	8.7 cp	2 cp	4.3 cp	Copies transferred
	4.3 cp	1.9 cp	3.46 cp	Copies analyzed

QIAcuity Software

QIAcuity Control Software is an integral part of the QIAcuity system. It offers a GUI (graphical user interface) for basic functionalities such as plate setup, changing the order of plates to be processed, and monitoring run status in real-time.

After a run, the data are stored in the instrument's memory and sent to the QIAcuity Software Suite for analysis.

The QIAcuity Software Suite, provided with the instrument and installed on a separate computer, controls one or multiple

QIAcuity instruments, either connected directly to one instrument or using an existing local area network (LAN).

When integrated into a local area network, the computer hosts the QIAcuity Software Suite as a server that is accessible via LAN to other computers serving as clients.

It allows multiple users to access the software from other rooms or offices and analyze data via a standard browser without installing the software on multiple computers or accessing and exchanging data via the internet.

Example run and analysis views in the software

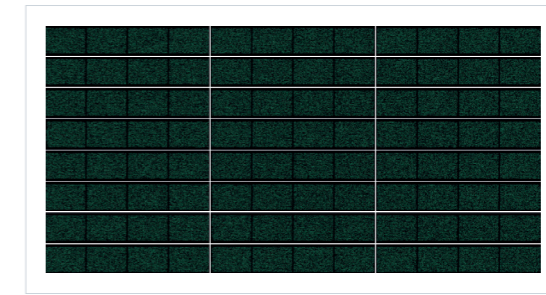
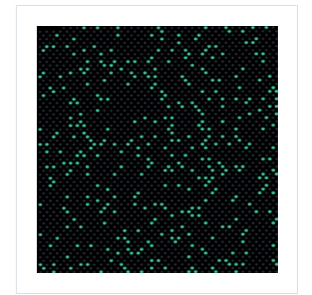
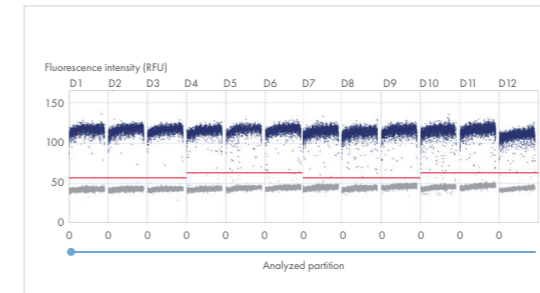


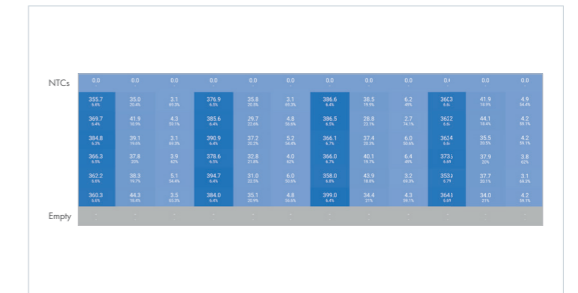
Plate image



Signal map



1D Scatterplot

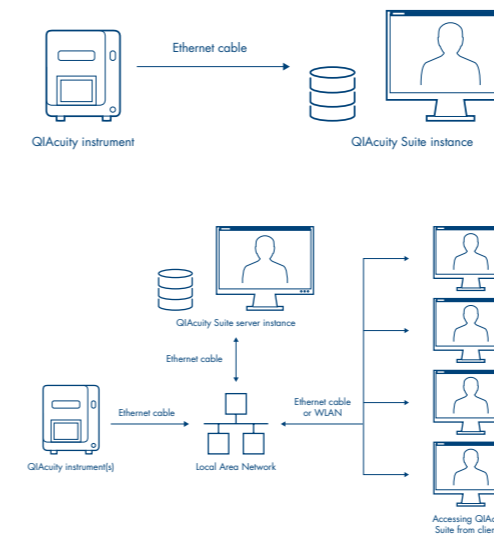


Heat map

Features and benefits

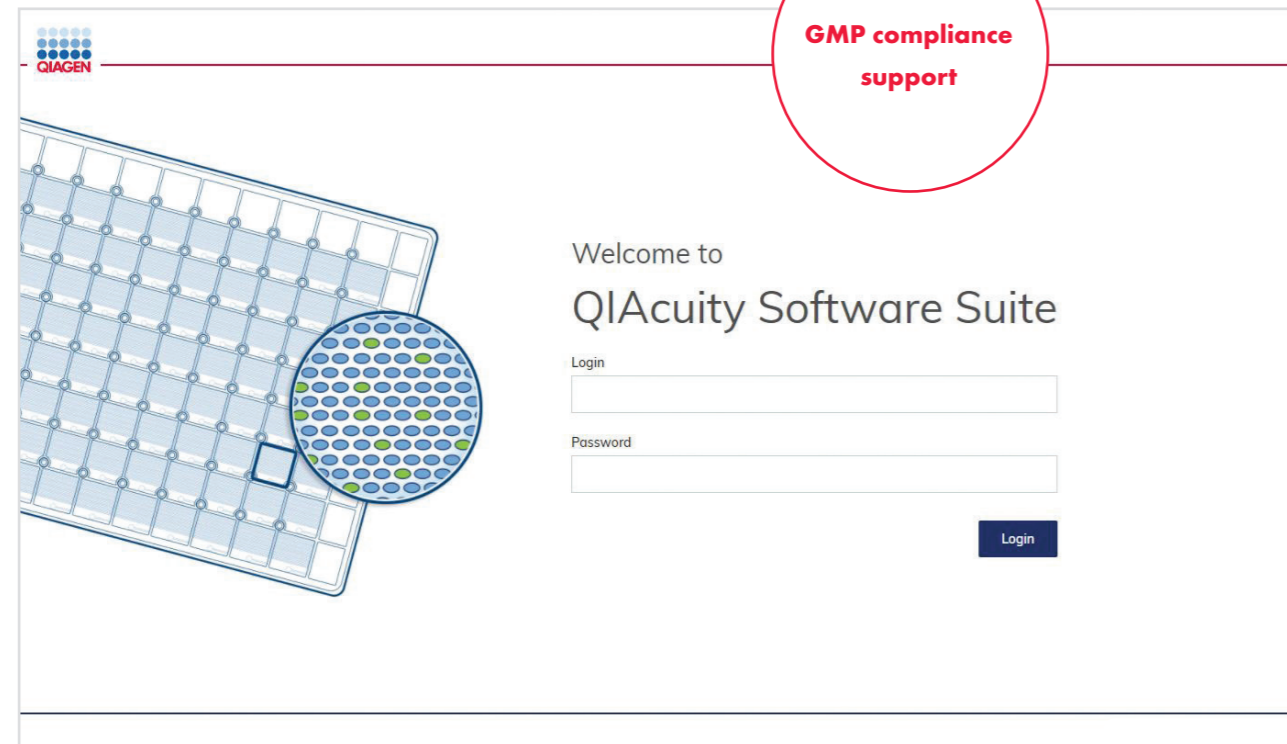
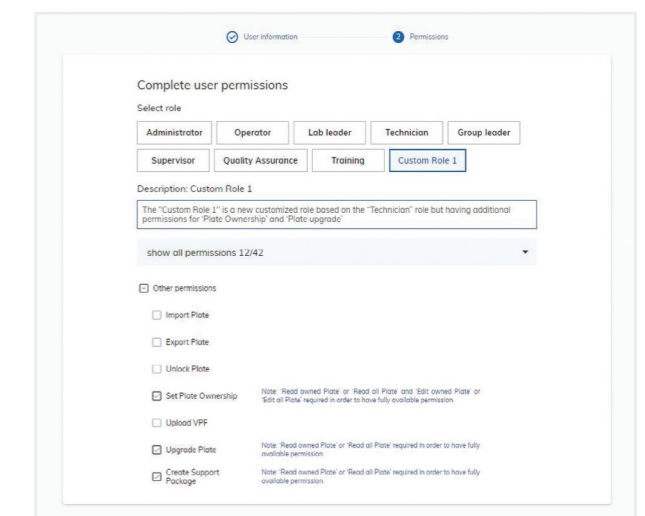
Easy to use – access, design and analyze from anywhere

- Direct connection via cable
- Network connection for multiple instruments and client access



Enables 21 CFR Part 11 compliance support in a GMP setting

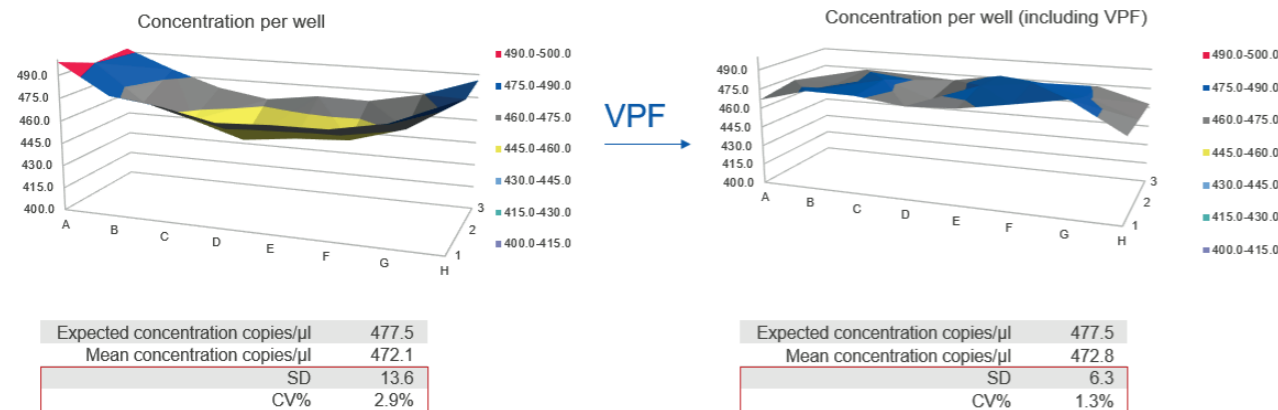
- Audit trail and traceability
- Electronic signatures
- Advanced user management with customized roles and permissions



Improving precision of concentration results by using a volume precision factor (VPF)

A precise determination of the cycled sample volume is needed to calculate target concentrations in dPCR. In general, nanoplates provide partitions of fixed sizes that enable an exact and reproducible sample concentration calculation. To compensate for even the slightest volume variations between different wells of a plate or between different plate batches, a volume precision factor (VPF) is available for each plate. The VPF is a set of factors

for each well included in the software. It consists of 96 individual factors that can address the well-to-well variability and reduce variations between different molding forms resulting in batch-to-batch variability. This increases the precision of concentration measurements in dPCR, particularly for sensitive applications such as rare mutation analysis.



Applying VPF to address well-to-well variability

Hyperwell option added to certain analysis for higher accuracy

Multiple wells can be grouped and analyzed as a single well to achieve higher accuracy. For the analysis, hyperwells are treated as a single well but with more partitions. This may be

helpful for rare event detection if the sample volume to be analyzed exceeds the volume that can be loaded into a single well.

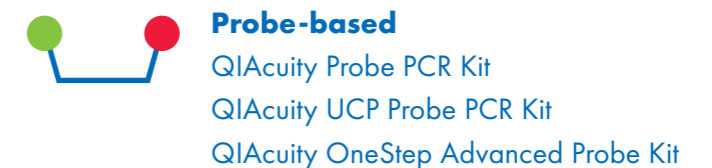
➔ For more information about QIAcuity Digital PCR System, visit: www.qiagen.com/qiacuity

QIAcuity reagents

The QIAcuity system is optimized for hydrolysis probes and EvaGreen dye, allowing you to expand applications using flexible dPCR chemistry.

A reagent for every need

- Optimized for best performance in nanoplate microfluidic
- Includes special reference dye needed for dPCR analysis and counting analyzable partitions
- Highly concentrated master mixes enabling larger sample volumes
- All mixes for single-plex and multiplex use
- QIAcuity OneStep Advanced Probe Kit with thermostable RT (HotStart) enabling multi-plate runs in high throughput
- QIAcuity UCP Probe PCR Kit with nucleic acid-depleted reagents minimizing contaminating DNA background



QIAcuity assays

Thanks to the high sensitivity and superior precision and accuracy, a wide range of samples and applications can benefit from digital PCR.

Wet-lab verified dPCR assays on GeneGlobe



- For DNA targets; for detection of copy number variation or mutations related to cancer and oncogenesis
- For quantification of microRNA targets
- For quantification of RNA/lncRNA targets and gene expression studies
- For detection of bacterial 16S rRNA and fungal ribosomal rRNA sequences; for species identification, detection of virulence genes and antibiotic resistance genes



Rare mutation detection
dPCR LNA Mutation Assays



Pathogen detection
dPCR Microbial DNA
Detection Assays



Copy number variation
dPCR Copy Number Assays
dPCR CNV Probe Assays



Gene expression
QuantiNova LNA PCR Assays



miRNA detection
miRCURY LNA
miRNA PCR Assays



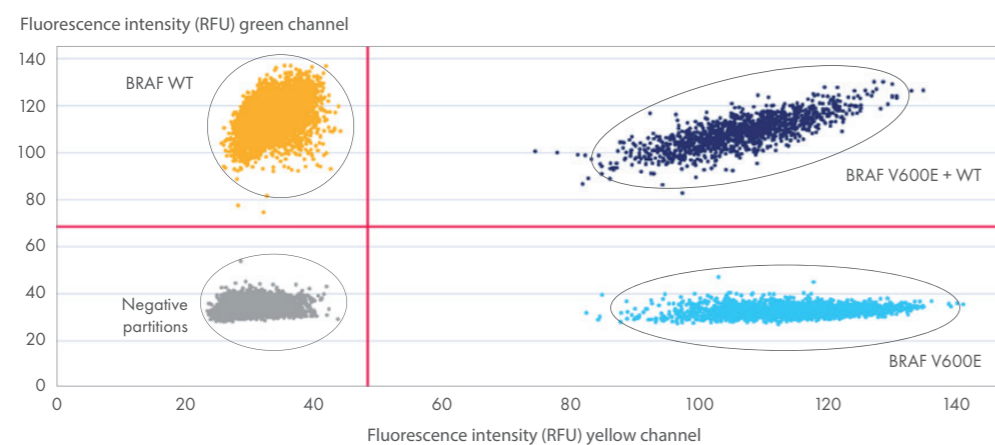
Cell and gene therapy
QIAcuity Cell and Gene Therapy
(CGT) dPCR Assays
CGT Viral Vector Lysis Kit
QIAcuity Residual DNA Quantification Kits



Find and configure your dPCR assay with ease at
www.geneglobe.qiagen.com/products/analysis-type/analysis-type-dpcr

dPCR LNA Mutation Assays

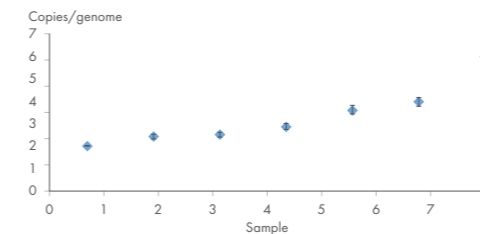
- Locked nucleic acid (LNA) technology increases assay specificity and sensitivity
- Duplex assay design detects mutated and wild-type sequences
- Two dye combinations allow detection of two targets in the same reaction



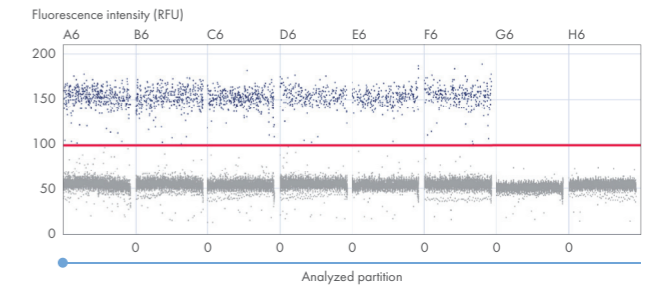
Detection of BRAF V600E mutation. dPCR LNA mutation detection/BRAF V600E assay showing four populations (single positives, single negatives, dual positives and dual negatives) in a 2D scatter plot and the ability to identify the threshold between the four populations.

dPCR Copy Number Assays

- Predesigned assays for all genes in the human genome deliver reliable results
- Three design locations per gene – 5', middle, 3' – to amplify your region of interest
- Simple and straightforward EvaGreen-based dPCR format enhances usability



MYC copy number determination in MCF-7 cell line. Copy number plot of MYC normalized with TERT as reference. The samples S1–S7 are WT/MCF-7 mixtures containing increasing amounts of MCF-7 DNA: S1=0%, S2=11%, S3=20%, S4=33%, S5=43%, S6=50% and S7=100%. MYC copy number determined using the QIAcuity System matched the expected numbers: S1=2, S2=2.4, S3=2.8, S4=3.3, S5=3.7, S6=4 and S7=6. The WT, MCF-7 and mixture samples were analyzed with 4 ng/reaction.

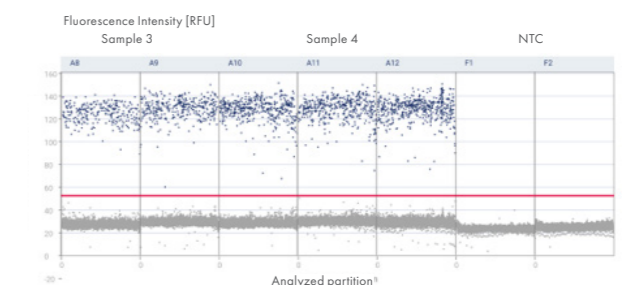
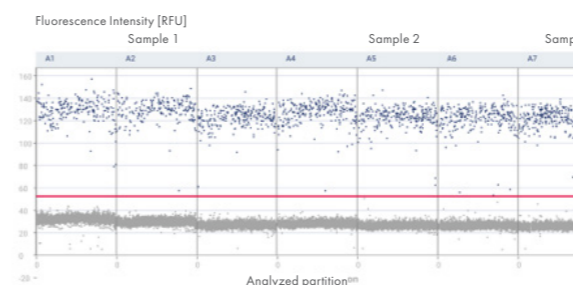
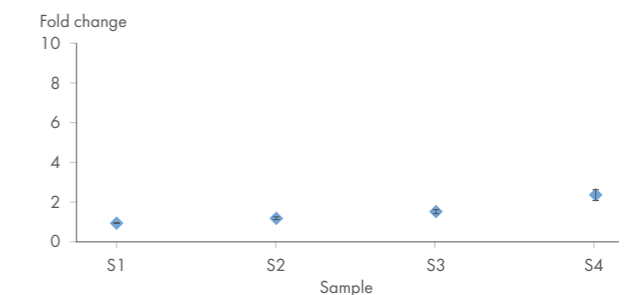


dPCR Copy Number Assay validation. 1D scatter plot showing single-well assay validation data of dPCR Copy Number Assay MYC with different human gDNA input amounts (**A6–C6**: 6 ng/reaction, **D6–F6**: 4 ng/reaction, **G6**: NTCs; Green channel for EvaGreen detection).

QuantiNova LNA PCR Assays

- Over 1.3 million assays detect any human, mouse or rat mRNA or lncRNA
- Short LNA-enhanced primers provide exceptional sensitivity and specificity
- EvaGreen-based dPCR allow accurate and convenient transcript analysis

IL-4 gene expression analysis – detecting small expression changes with the highest precision. Synthetic IL4 RNA was spiked into non-IL4 expressing Universal Human Reference RNA (Thermo Fisher Scientific). IL4 fold-expression changes in samples S2, S3 and S4 were calculated using S1 as reference sample and HPRT as reference target. The mean fold change (from 3 technical replicates/sample) in IL4 expression: S1=0 (reference), S2=1.3, S3=1.5 and S4=2.3.

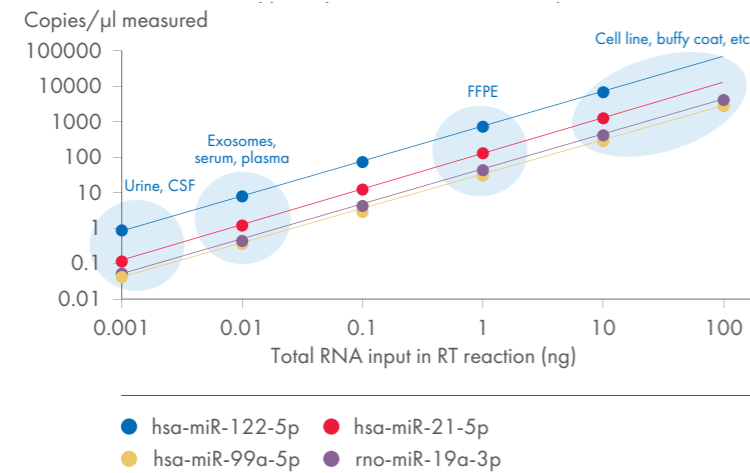


1D Scatter Plot of IL4 QuantiNova LNA PCR Assay showing the resolution

miRCURY LNA miRNA PCR Assays

- One RT reaction for all miRNA and two LNA-enhanced miRNA-specific primers for highest specificity
- EvaGreen-based dPCR allow absolute quantification of miRNA expression changes
- Full miRBase coverage enables miRNA profiling from any organism

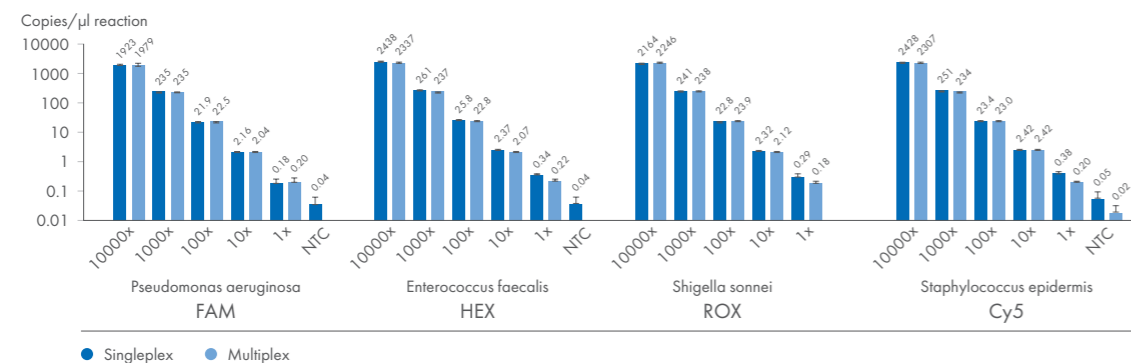
Typical yields from various samples



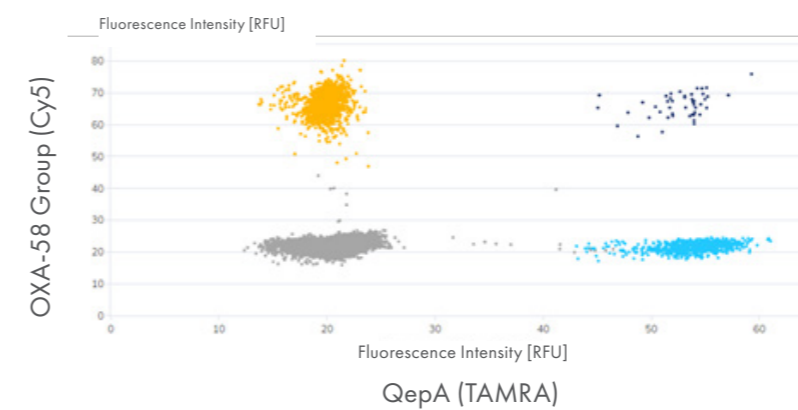
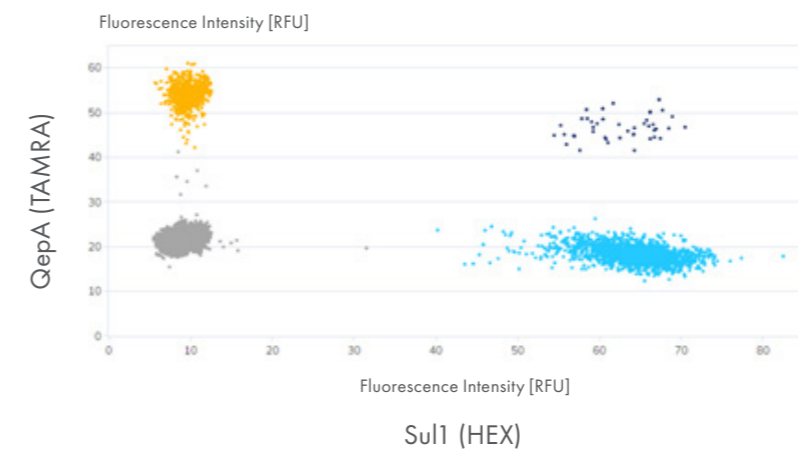
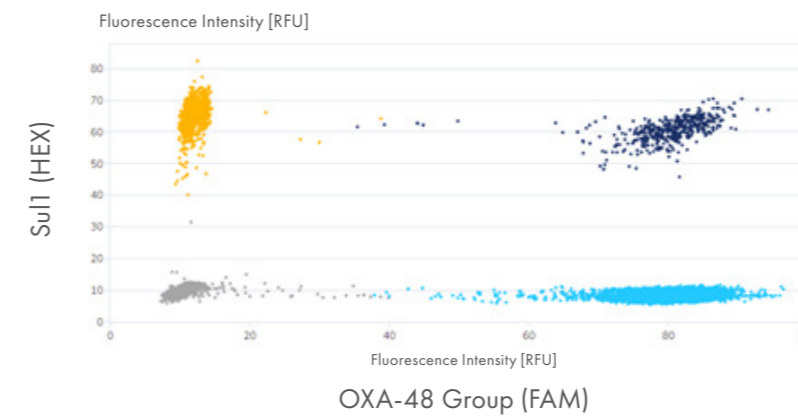
Reliable miRNA detection from different samples at 1 pg RNA input without pre-amplification

dPCR Microbial DNA Detection Assays

- Assays for more than 680 targets detecting microbial species, virulence genes, viruses or antibiotic resistance genes
- Dye selection enables multiplexing of up to 5 targets per reaction
- Combine microbial DNA and viral RNA targets in one reaction using the QIAcuity OneStep Advanced Probe Kit



Microbial detection in multiplex on the QIAcuity. Single-plex versus multiplex setup quantifying four different bacterial targets. The data shows a very similar and precise quantification of all targets for concentrations between 0.2–2500 cp/µl.



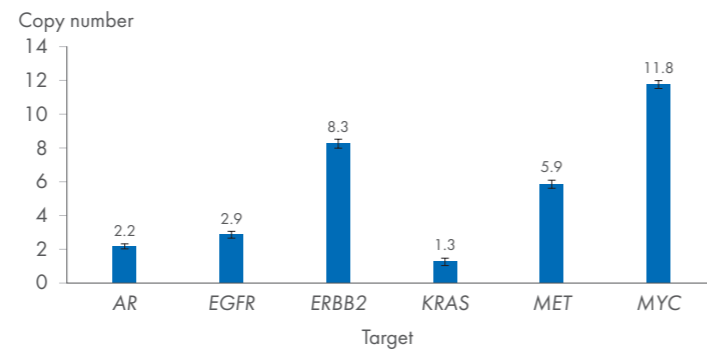
Signal separation between channels in multiplex on the QIAcuity. Four assays targeting four bacterial resistance genes were run in multiplex reactions. 2D scatter plots of various dye combinations from the 4-plex runs.



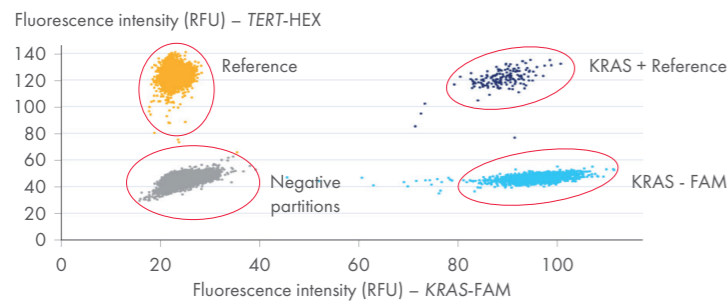
Download the **QIAcuity Application Guide** for detailed information about setting up experiments and analyzing results for applications, including copy number variation analysis, rare mutation detection and gene expression.

dPCR CNV Probe Assays

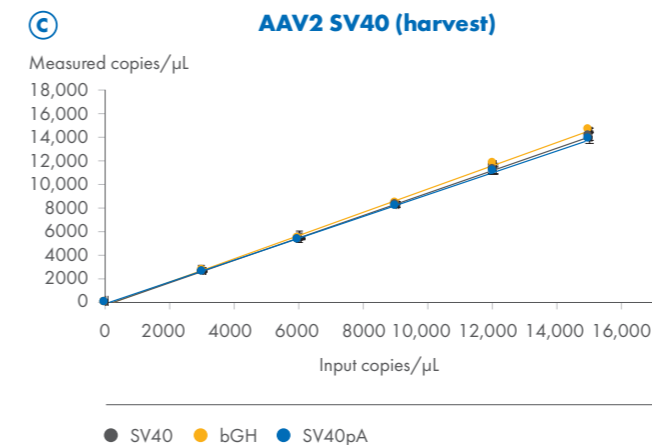
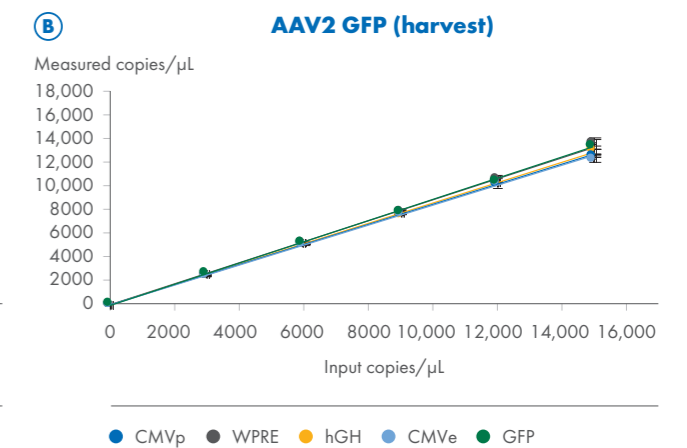
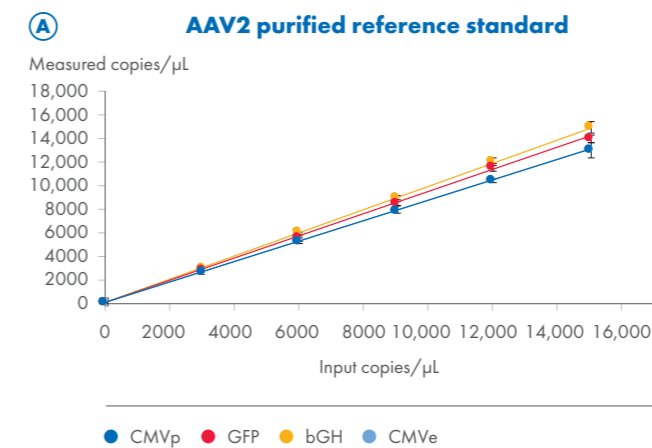
- Highly sensitive discrimination between small copy number changes, such as one copy difference between samples with minimal template input
- Multiplexing of up to five CNV targets in a single reaction without replicates saves precious samples and increases sample throughput cost-effectively



dPCR detects CNVs in different target genes without the need for replicates. CNV analysis was performed in an SK-BR3 cell line using a Nanoplate 8.5k 96-well and dPCR CNV Probe Assay on the QIAcuity dPCR system.



2D scatterplots of duplex reactions for KRAS (target) and TERT (reference). The KRAS target was labeled with a FAM probe and the reference TERT with a HEX fluorophore for multiplexing. CNV analysis was done in an SK-BR3 cell line using a Nanoplate 8.5k 96-well and dPCR CNV Probe Assay on the QIAcuity dPCR system.

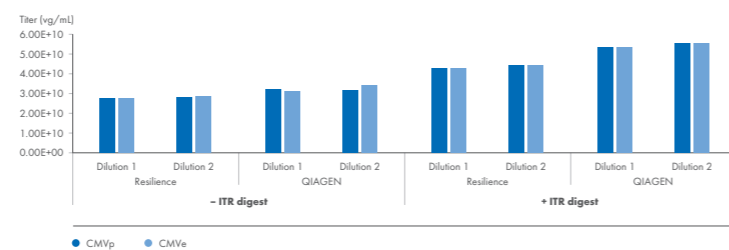


AAV2	CMVe	CMVp	WPRe	hGH pA	GFP	ITR	SV40p	bGH	SV40 pA
Coefficient of variation (8.5k) [%]	2.4	2.7	2.7	3.6	3.4	4.7	4.8	3.4	3.2
Coefficient of variation (26k) [%]	2.9	2.4	2.1	1.7	3.7	5.2	2.2	1.4	1.7
R²	1.00	1.00	1.00	1.00	1.00	1.00	1.00	1.00	1.00
Deviation (8.5k vs 26k) [%]	1.07	2.49	5.35	4.35	2.6	1.73	0.59	0.07	0.17

High linearity from 2.5–15,000 copies/μL independent of the targets and purity of the AAV sample quantified with CGT dPCR Assays. Coefficients of variation (CV) and R² values for all assays on 8.5k and 26k Nanoplates are shown. Deviations between the quantification on 8.5k and 26k Nanoplates were based on the calculated titers of at least three dilution steps.

CGT Viral Vector Lysis Kit and QIAcuity Cell and Gene Therapy (CGT) dPCR Assays

- Offer a standardized workflow for the lysis of AAV2, AAV5, AAV6, AAV8 and AAV9 and adenoviruses, enabling robust, accurate and precise viral titer determination
- Lysate is optimized for QIAcuity CGT dPCR Assays in combination with the QIAcuity Probe PCR Kit
- Assays enable singleplexed and multiplexed cell and gene therapy applications, including viral titer and vector copy number measurements



Increase of quantification with QIAGEN lysis over Resilience lysis				Increase of quantification when performing an ITR digest			
	ITR	CMVp	CMVe		ITR	CMVp	CMVe
- ITR digest	20%	16%	17%	Resilience	71%	57%	55%
+ ITR digest	18%	25%	25%	QIAGEN lysis	69%	69%	65%

Effect of ITR digest on differently processed AAV samples compared to heat lysis (Resilience protocol vs CGT Viral Vector Lysis Kit protocol). Purified AAV2 samples were processed using the CGT Viral Vector Lysis Kit (QIAGEN) or following the Resilience protocol (Resilience). QIAGEN lysis increases quantification by at least 16%. Digestion of secondary structure within ITR regions increases quantification by at least 55% depending on the target and processing workflow followed.

Genomic Services

To realize the full potential of dPCR, you need high-quality multiplex assays to answer your specific biological questions in the most accurate way. However, assay design takes time and slows you down. Partnering with our Genomic Services is a straightforward and reliable way to extend your in-house resources and ensure robust assay performance.

What can you expect?

Highly efficient assay design

process: Optimized design tools, highly skilled design experts, and decades of experience

Versatile assay design offering:

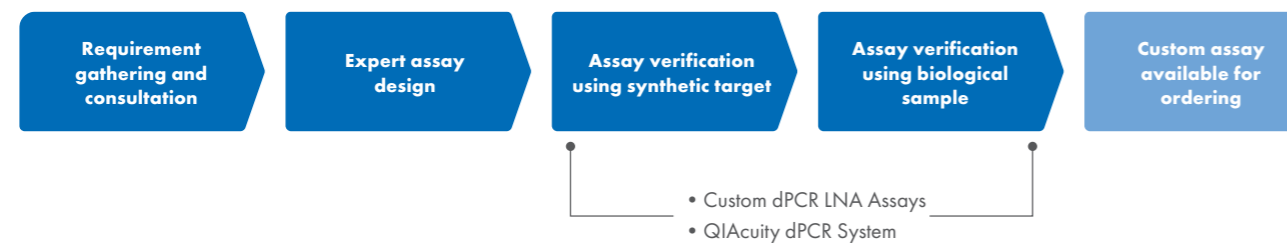
High-level multiplexing across any species and application, such as mutation detection, CNV, gene expression, and species detection

Enhanced specificity and

sensitivity: Assay design based on proprietary LNA technology

Highly flexible service offering:

Get a custom-designed assay or add an in-depth wet-lab verification service



Learn more about the service and consult an expert at www.qiagen.com/custom-dPCR-assay-design

QIAcuity services

Our instrument service plans are offered at various levels, so you can choose the one that best fits your needs and budget. Let our highly-skilled certified service team keep your instrument maintained so you can focus on results

- Maximize uptime and productivity
- Receive priority support and service
- Reduce the risk of non-compliance
- Control costs
- Minimize disruption of laboratory performance

All parts, labor and travel costs are included for standard repairs, and the annual preventive maintenance gives you peace of mind.



Get a full overview of the QIAcuity service solutions at www.qiagen.com/qiacuity-services

Digital MIQE guidelines

As part of the PCR community, you're well aware of the reproducibility crisis in research and the daily challenges in a molecular biology laboratory using qPCR, dPCR, or any comparable techniques.

To support the community, a group of experts published the Minimum Information for Publication of Quantitative Real-Time PCR Experiments (MIQE) guidelines in 2009, targeting the reliability of results for credible publications, promoting reproducibility between laboratories, and increasing experimental transparency.

Fast forward to 2013, owing to a growing interest in dPCR because of its accessibility and affordability, the Minimum Information for Publication of Quantitative Digital PCR Experiments (dMIQE) guidelines were published to ensure global standardization. The guidelines were further updated in 2020.

- Why do we need such guidelines?
- What are these guidelines?
- How are these guidelines revolutionizing dPCR experiments?



Follow our three-part webinar series on dMIQE guidelines to find out: www.qiagen.com/dMIQE-webinars

Voice of our customers



"We tested QIAGEN's QIAcuity dPCR for quantification of viral titer, vector copy number and residual host cell DNA – all critical to in-process quality control in gene therapy. It is easy to use, fast, scalable and complies with requirements for GMP. The system is a great addition to our analytical development and testing services, process development and R&D platforms which is available to our clients now."

Dana Cipriano, Senior Vice President, Testing and Analytical Services, Center for Breakthrough Medicines in King of Prussia, PA



"Digital PCR has been somewhat of a revolution in the field of copy number analysis because of the resolution. What I like about the QIAcuity system is the simple and fast workflow and that you use less plastic tips and plates in the process which is good for the environment."

Dr. Johanna Andersson-Assarsson, Department of Molecular and Clinical Medicine, University of Gothenburg



"With the new, high-throughput QIAcuity Eight, we were able to detect new variants of SARS-CoV-2 in wastewater samples successfully. This fast and scalable technology can provide a valuable addition to our environmental, biological testing services, which we will offer to our clients in the near future."

Dr. Franz Durandet, President of I.A.G.E. in Montpellier, France



"When working with low microbial biomass concentrations, qPCR is a good tool for quantification, but dPCR is the best approach. The QIAcuity dPCR instrument allows us to consistently detect and quantify microorganisms in soil, rock and water."

Prof. John R. Spear, Department of Civil and Environmental Engineering, Colorado



"Our lab loves using the QIAcuity digital PCR system for absolute quantification of targets from a wide range of samples. The workflow is straightforward, easy to learn, and generates incredibly consistent and sensitive results."

Drew Capone and colleagues, University of North Carolina at Chapel Hill, NC



"Digital PCR has higher detection rates at low DNA concentrations and can handle high concentrations of PCR inhibitors present in marine coastal ecosystems. We found the QIAcuity especially straightforward and fast. It can quantify eDNA from invasive species with more accuracy and sensitivity, independent of the amplification efficiency."

Per Sundberg, CEO, SeAnalytics AB, Gothenburg, Sweden

Ordering Information

Product	Contents	Cat. no.
QIAcuity One, 2plex Platform System FUL-1	One-plate digital PCR instrument for detecting up to 2 fluorescent dyes, notebook computer, Nanoplate Roller, USB flash memory, and QIAcuity Software Suite: includes installation, training, full agreement for 1 year with a 2-business day response time, and 1 preventive maintenance visit	911015
QIAcuity One, 5plex Platform System FUL-1 *	One-plate digital PCR instrument for detecting up to 5 fluorescent dyes, notebook computer, Nanoplate Roller, USB flash memory, and QIAcuity Software Suite: includes installation, training, full agreement for 1 year with a 2-business day response time, and 1 preventive maintenance visit	911035
QIAcuity Four Platform System FUL-1 *	Four-plate digital PCR instrument for detecting up to 5 fluorescent dyes, notebook computer, barcode scanner, Nanoplate Roller, USB flash memory, and QIAcuity Software Suite; Includes installation, training, full agreement for 1 year with a 2-business day response time, and 1 preventive maintenance visit	911045
QIAcuity Eight Platform System FUL-1 * †	Eight-plate digital PCR instrument for detecting up to 5 fluorescent dyes, notebook computer, barcode scanner, Nanoplate Roller, USB flash memory, and QIAcuity Software Suite: includes installation, training, full agreement for 1 year with a 2-business day response time, and 1 preventive maintenance visit	911055
QIAcuity Nanoplate 26k 8-well (10)	8-well dPCR Nanoplate with 26K partitions and 40 µl reaction volume per well, including Nanoplate seals	250031
QIAcuity Nanoplate 26k 24-well (10)	24-well dPCR Nanoplate with 26K partitions and 40 µl reaction volume per well, including Nanoplate seals	250001
QIAcuity Nanoplate 8.5k 24-well (10)	24-well dPCR Nanoplate with 8.5K partitions and 12 µl reaction volume per well, including Nanoplate seals	250011
QIAcuity Nanoplate 8.5k 96-well (10)	96-well dPCR Nanoplate with 8.5K partitions and 12 µl reaction volume per well, including Nanoplate seals	250021
Nanoplate Seals (11)	Nanoplate seal for sealing QIAcuity Nanoplates	250099
Nanoplate Tray (2)	Nanoplate Tray improving plate-handling during pipetting or carrying	250098
QIAcuity Probe PCR Kit (1 ml, 5 ml and 25 ml)	4x concentrated QIAcuity Probe Mastermix and Water	250101, 250102, 250103

Ordering Information

Product	Contents	Cat. no.
QIAcuity EG PCR Kit (1 ml, 5 ml and 25 ml)	3x concentrated QIAcuity EvaGreen Mastermix and Water	250111, 250112, 250113
QIAcuity OneStep Advanced Probe Kit (1 ml and 5 ml)	OneStep Advanced Probe Master Mix (4x), OneStep RT Mix (100x), Enhancer GC, QN Internal Control RNA, RNase-Free Water	250131, 250132
QIAcuity UCP Probe PCR Kit (1 ml and 5 ml)	4x concentrated QIAcuity UCP Probe Mastermix and UCP Water	250121, 250122
dPCR LNA Mutation Assays (200 rxn and 1000 rxn)	Single tube containing ready-to-use 30x-concentrated assay with choice of FAM + HEX or Atto 550 + ROX detection dyes	250200, 250201 (in GeneGlobe)
dPCR Copy Number Assays (200 rxn and 1000 rxn)	Single tube containing ready-to-use 25x-concentrated assay	250205, 250206 (in GeneGlobe)
dPCR CNV Probe Assays (reference, gene-specific and centromeric reference)	Single tube containing ready-to-use, 20x-concentrated assays	250210, 250212, 250213 (in GeneGlobe)
CGT Viral Vector Lysis Kits (100 rxn and 1000 rxn)	CGT Sample Stabilizer, DNase I, Nuclease-Free Water, CGT Lysis buffer, CGT DNase I buffer and CGT Dilution Buffer	250272, 250273
QIAcuity Cell and Gene Therapy (CGT) dPCR Assays (ten assays)	20x ready-to-use primer-probe mix, available in multiple fluorophore choices, FAM, HEX and Cy5	250230–250256
QIAcuity Residual DNA Quantification Kits (CHO, HEK293 and E. coli)	resDNA Quant Mastermix (2x), 1x Positive Control, 1x Internal Control, 3x RNase-Free Water; resDNA Quant Standard (1x) and Rehydration Buffer	250220–250225
dPCR Microbial DNA Detection Assays	One tube with the lyophilized assay, dye (fluorophore) configurable, 200 reactions (40 µl reaction in Nanoplate 26k)	250207 (in GeneGlobe)
QuantiNova LNA PCR Assays for Digital PCR (200 rxn and 750 rxn)	Pre-designed mRNA/lncRNA-specific primer mixture in a single tube	249990, 249992 (in GeneGlobe)
miRCURY LNA miRNA PCR Assays for Digital PCR	Contains forward and reverse primers for SYBR® Green-based, real-time qPCR reactions and EvaGreen-based digital PCR reactions	339306 (in GeneGlobe)

* Additional instrument and Service bundles are available.

† For all systems, Installation and Training is included but are additionally available as separate service offerings. For specific catalog numbers and additional information, visit www.qiagen.com or contact your local sales representative.

 To learn more about QIAGEN's dPCR technology visit www.qiagen.com/dPCR

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