

PRIMETIME™ qPCR PROBES

Double- and single-quenched probes for use in 5' nuclease assays



DYES AND QUENCHERS FOR EVERY EXPERIMENT

PrimeTime qPCR Probes provide reliable sensitivity even in demanding applications such as multiplexing and digital PCR. PrimeTime qPCR Probes are available in a wide variety of dye-quencher combinations (Table 1) that are compatible with common qPCR instruments.

ACHIEVE CONSISTENT RESULTS

All PrimeTime Probes are HPLC purified, and then verified by mass spectrometry, to deliver batch-to-batch consistency and minimize the need for troubleshooting.

Table 1. Commonly used fluorophores and quenchers

Fluorophore	Emission wavelength (nm)	Quencher	* Probes with 6-FA HEX fluorophore traditional, single
6-FAM*	520	ZEN/Iowa Black™ FQ	either Iowa Black Hole Quencher- licenses required
SUN™*	554		
JOE™*	555		† Black Hole Quen be used as a que party licenses red
HEX*	555		
MAX™*	557		Double-quenche custom order.
Cy® 3	564	lowa Black RQ†‡	
ATTO™ 550§	575		§ ATTO-labeled po custom order.
ROX	608		¥ Cy 5 is also avail probe with BHQ licenses required
Texas Red® -X	617		
ATTO 647N§	662		¶ Available as rese
Cy 5 [¥]	668		
Cy 5.5	706	Black Hole Quencher®-3¶	

- AM, SUN, JOE, MAX, or res are also available as le-quenched probes with ck FQ (license free) or Black -1 (additional third-party ed for diagnostic use).
- encher-2 (BHQ2) may also uencher (additional thirdequired for diagnostic use).
- ned probes available as a
- probes available as a
- ilable as a single-quenched 22 (additional third-party d for diagnostic use).
- earch use only.

IMPROVE ASSAY SENSITIVITY WITH DOUBLE-QUENCHED PROBES

Reduce background and increase assay sensitivity with ZEN or TAO Double-Quenched Probes. Our exclusive internal quenchers are 9 bases from the 5' fluorophore and work in combination with the 3' Iowa Black quencher for maximum probe performance (Figure 1).

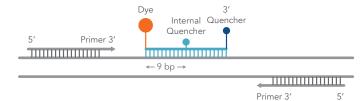


Figure 1. Schematic of a PrimeTime qPCR 5' Nuclease Assay using a double-quenched probe that includes a dye, a ZEN or TAO internal quencher, and a 3' quencher.

With nearly 4 times lower background fluorescence (Figure 2A) and approximately 30% increased signal (Figure 2B), ZEN Double-Quenched Probes simply perform better. See performance data for TAO Double-Quenched Probes at www.idtdna.com/qPCRprobes.

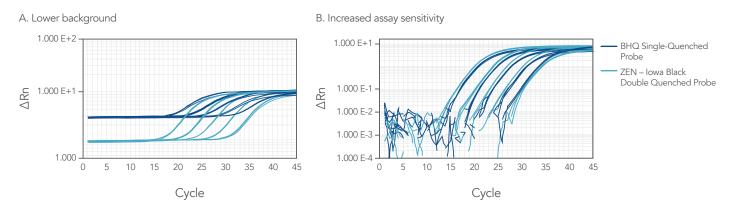


Figure 2. Increased signal detection and assay sensitivity from ZEN Double-Quenched Probes. (A) ZEN Probes (light blue) provide greater dye quenching, producing lower background and, therefore, higher signal intensities than standard single-quenched probes (BHQ Probes; dark blue). (B) ZEN Probes increase assay sensitivity, as demonstrated by the earlier Cq values observed compared to standard, BHQ single-quenched probes.

ACHIEVE MAXIMUM QUENCHING FOR LONG PROBES

Effective quenching for ZEN Double-Quenched Probes as long as 40 bases means more effective designs, even for AT-rich targets.

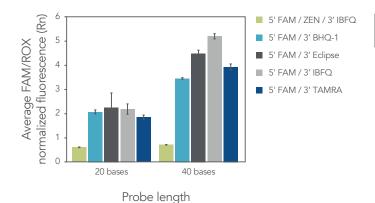


Figure 3. Only ZEN Double-Quenched Probes maintain low background signal with increasing probe length. Probes of 2 lengths (20 or 40 bases) with 5 different quenchers were compared in 10 singleplex qPCRs. Six replicate reactions with each probe type were run with 50 ng of cDNA and the TaqMan® Gene Expression Master Mix (Thermo Fisher) under standard cycling conditions on the Applied Biosystems 7900HT system. Key: IBFQ = lowa Black FQ Quencher (IDT); BHQ-1 = Black Hole Quencher-1 (Biosearch Technologies); MGB Eclipse® = Eclipse quencher (ELITech Group).

ORDERING INFORMATION

Visit www.idtdna.com/qPCRprobes to enter your sequence and choose modifications

> FOR MORE INFORMATION, VISIT WWW.IDTDNA.COM/qPCRPROBES

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