# Quantabio

Product Information			
PerfeCTa <sup>®</sup> SYBR <sup>®</sup> Green SuperMix			
Part Number	95054-100		
Number of Reactions	100 reactions		
Reaction Size	50 µL		
Storage Temperature	-25ºC to -15ºC		
Lot Number	028634		
Reference Number	121318		
Expiration Date	12/31/2021		

## Product Description:

PerfeCTa SYBR Green SuperMix is a user-friendly, 2X concentrated reaction mix that simplifies setup and reduces errors with optimized reference dye and pre-blended AccuVue plate loading dye for visual confirmation of reagent addition and mixing. This proprietary buffer technology stabilizes a high concentration of SYBR Green I dye to ensure maximum optical signal with low abundance or small targets (such as microRNA). Successful detection with a nonspecific, dsDNA intercalating dye requires precise target amplification as off-target primer elongation will contribute to overall fluorescent signal and lead to over-reported relative abundance values. This reagent is powered by a highly-processive, ultra-pure Taq DNA polymerase mutant with stringent, ultra-pure AccuStart™II antibody hot start technology that allows ambient room-temperature setup and maximal enzyme kinetics after rapid, irreversible denaturation at 95°C.

## Component Part Numbers:

#### 84016 PerfeCTa SYBR Green SuperMix, 1.25mL

Product Specifications				
95054				
Assay	qPCR ß actin Plasmid DNA Functional Assay for SYBR Green SuperMix	DNase	RNase	
Result	Pass	Pass	Pass	

# **Quality Control Analysis and Specifications:**

# Nuclease Assay:

**DNase:** DNase activity must be below the detectable limits of 100 pg DNase I equivalent as assayed using a fluorogenic substrate following a 1 hour incubation at 37°C with each kit component at 1X concentration.

**RNase:** RNase activity must be below the detectable limits of 1 pg RNase A equivalent as assayed using a fluorogenic substrate following a 1 hour incubation at 37°C with each kit component at 1X concentration.

**qPCR ß actin Plasmid DNA Functional Assay for SYBR Green SuperMix:** Real-time PCR detection of log-fold serial dilutions of a control DNA from 10 copies to  $1 \times 10^7$  copies. Cq standard curve analysis must have coefficient of determination ( $r^2$ )  $\geq$ 0.990 with a slope between -3.20 and -3.65.

#### Limitations of Use

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