# Quantabio

| Product Information           |                |  |
|-------------------------------|----------------|--|
| AccuStart Long Range SuperMix |                |  |
| Part Number                   | 95199-025      |  |
| Number of Reactions           | 25 reactions   |  |
| Reaction Size                 | 50 uL          |  |
| Storage Temperature           | -25ºC to -15ºC |  |
| Lot Number                    | 66229981       |  |
| Reference Number              | 060823         |  |
| Expiration Date               | 06/30/2026     |  |
|                               |                |  |

## Product Description:

The AccuStart Long Range SuperMix is a 4x, ready-to-use solution that contains all the components for long-range target amplifications, including a blend of two hot-start thermostable DNA polymerases (one with strong proof-reading activity) and an optimized buffer to ensure high efficiency, sensitivity and specificity for amplification of long-range targets. This mix enables routine and easy amplification of up to 24kb target from human genomic DNA with high accuracy (> 10-fold better fidelity than Taq) and accommodates targets with broad GC-content (no separate GC buffer needed). Fast cycling conditions (≥2 kb/min) can be implemented for extremely long targets. This product is also capable of amplifying multiple targets simultaneously – up to 6 targets up to 6 kb each.

#### **Component Part Numbers:**

#### 84638 AccuStart Long Range SuperMix 0.313 mL

| Product Specifications |  |
|------------------------|--|
| 95199                  |  |
| Assay                  | Long Range PCR Master Mix Functional Assay |
| Result                 | Pass                                       |

### **Quality Control Analysis and Specifications:**

**The functionality of the AccuStart Long Range SuperMix** is evaluated in a PCR reaction to produce a major band of amplification for targets at 24 kb, 17.5 kb and 8.5 kb from 5 ng of Human Genomic DNA.

#### Notes:

Enzyme components were tested prior to formulation of the master mix and found free of contaminating endonucleases and exonucleases. Enzyme purity was >99% as determined by SDS-PAGE and negligible *E.coli* genomic DNA contamination was confirmed by qPCR. Specific activity was verified for each enzyme pre-formulation.

#### Limitations of Use

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