



QuantSeq 3' mRNA-Seq Library Prep

The most cited reason for doing RNA sequencing is to investigate gene expression. Whilst differential expression may be addressed by conventional mRNA sequencing, this method requires high read depth sequencing and complex data analysis.

QuantSeq 3' mRNA-Seq offers a much faster and more cost-efficient way to get expression analysis done with the added benefits of high-multiplexing capability and the robustness to deal with the most difficult of samples!



Introduction

QuantSeq 3' mRNA-Seq generates only one fragment per transcript at the 3' end instead of covering the full length of the transcript with read as conventional mRNA-Seq (Fig. 1).

The most cost-efficient way to study differential gene expression

Since QuantSeq generates only one fragment per transcript, gene expression studies with QuantSeq require a vastly lower number of reads per sample (we recommend 3-5 M reads for standard applications) than with conventional mRNA-Seq (usually 20-50 M reads recommended). This means that more samples can be pooled and sequenced on one single lane, reducing the cost per sample drastically (Fig. 2).

Fast, easy, and robust library prep in under 4.5 hours

In contrast to conventional mRNA-Seq, QuantSeq only requires 5 major steps and can be completed in about 4.5 hours! Quant-Seq does not require any pre-processing of the RNA, such as poly(A) RNA selection or ribosomal RNA depletion since the protocol intrinsically selects only poly(A) RNA. This saves time and makes the protocol very robust (Fig. 3).



Figure 1 | In contrast to common mRNA-Seq methods, QuantSeq generates only one fragment per transcript at the very 3' end.



Figure 2 | Comparison of cost per sample of conventional mRNA-Seq and QuantSeq 3' mRNA-Seq for a complete gene expression profiling experiment.



Figure 3 | Workflow comparison of conventional mRNA-Seq and QuantSeq 3' mRNA-Seq for gene expression profiling.

Vast species, quantity, and quality compatibility, even ideal for FFPE samples

QuantSeq's 3' capture approach successfully generates fragments even from degraded RNA, such as RNA derived from FFPE samples. The technology is less sensitive to variations in RNA quality than conventional mRNA-Seq which requires high-quality RNA for mRNA enrichment.

Ordering Information Catalog Numbers: 015 (QuantSeq FWD 3' mRNA-Seq Library Prep Kit) 016 (QuantSeq REV 3' mRNA-Seq Library Prep Kit) 191 - 196 (QuantSeq FWD 3' mRNA-Seq V2 Library Prep Kits – with Lexogen's patented 12 nt Unique Dual Indices)

Free and simple data analysis

With QuantSeq, the number of reads mapped to a given gene is directly proportional to its expression. 3' mRNA-Seq does not rely on correct isoform annotation and identification for determining unambiguous gene expression values. Thus, the data analysis of QuantSeq reads is simple and requires only a short time. Researchers lacking bioinformatics expertise can access Lexogen's user-friendly data analysis pipeline - free with the purchase of a QuantSeq kit, directly from our website: www.lexogen.com/data-analysis-solutions



For more information and additional resources on QuantSeq visit our website.

Find more about QuantSeq at <u>www.lexogen.com</u>. Contact us at <u>support@lexogen.com</u>, +43 1 345 1212-41, or find your local contact at <u>www.lexogen.com/distributors</u>