

The Archer® FusionPlex® Lung Kit is a targeted next-generation sequencing (NGS) assay to detect EGFR vIII and MET exon 14 skipping events along with prominent ALK, BRAF, FGFR, NRG1, NTRK, RET, and ROS1 fusions and select point mutations in 14 key gene targets associated with lung cancer. The kit is powered by Archer's proprietary Anchored Multiplex PCR (AMP™) target enrichment chemistry to detect fusions in a single sequencing assay, even without prior knowledge of fusion partners or breakpoints.

Highlights



Focused fusion detection Small panel to interrogate discrete oncogenic fusions



Splice variants
Identify expressed EGFRvIII and
MET exon 14 skipping events



Sample versatility
Purpose-built for low-input FFPE
samples



Lyophilized workflow Reduce turn-around time and eliminate master mixes

Specifications

14

Gene targets

≥10ng

Input nucleic acid required*

500K

Recommended # of reads

2.5h

Hands-on time

9h

Total time

Illumina[®]

Platform

*Input recommendations for FFPE samples vary depending on Archer Preseq® RNA QC score; 50ng input recommended in absence of PreSeq screening

Gene targets



Legend

◆ SNV or indel

● Fusion, splicing or exon skipping

Product information

Illumina

AB0121 — 8 reactions

AB0128 — 16-reaction starter

AB0129 — 48 reactions



One size fits all.

Want to customize? Combine any of hundreds of wet lab-validated designs or modify an existing Archer panel with **Archer Assay Designer**, the premiere online design tool to build an assay that fits your exact requirements.

Learn more at archerdx.com/solid-tumor

