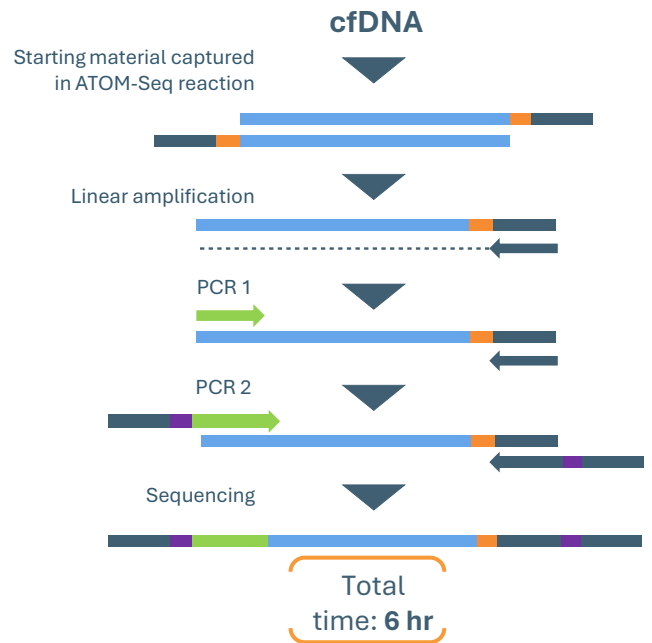


# Cell-free DNA Enrichment Workflow

NGS library preparation workflows for capturing and enriching cell-free DNA from plasma and urine

## Uniquely designed for challenging material

- Simple, ligation-free approach with no DNA end-repair
- Captures all single- and double-strand DNA
- Captures short and degraded material
- Efficient with low input quantities
- Single primer enrichment to maximise capture regardless of DNA breakpoint
- Unique molecular identifies for error suppression
- Minimal bead purification steps



## Minimal residual disease and cancer screening



ATOM-Seq's efficient capture of cell-free DNA makes it ideal for detecting any low frequency genetic signature from plasma or urine samples

Identifying rare low level tumour mutations

Track disease response

Monitoring residual disease



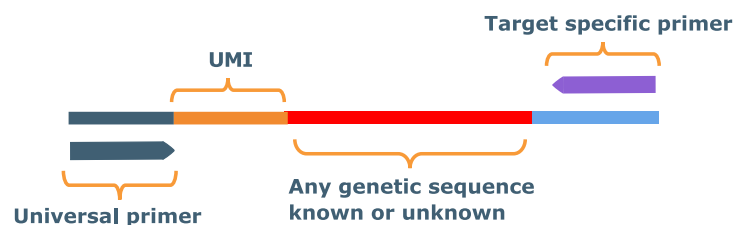
**Detect even the rarest clinical signatures** with confidence, using both UMIs and unique error-reducing workflow optimisations



**Maximum retention, minimal time,** due to minimal purification steps



**Get more from your patient sample** by capturing all cfDNA and enriching targets using a single primer

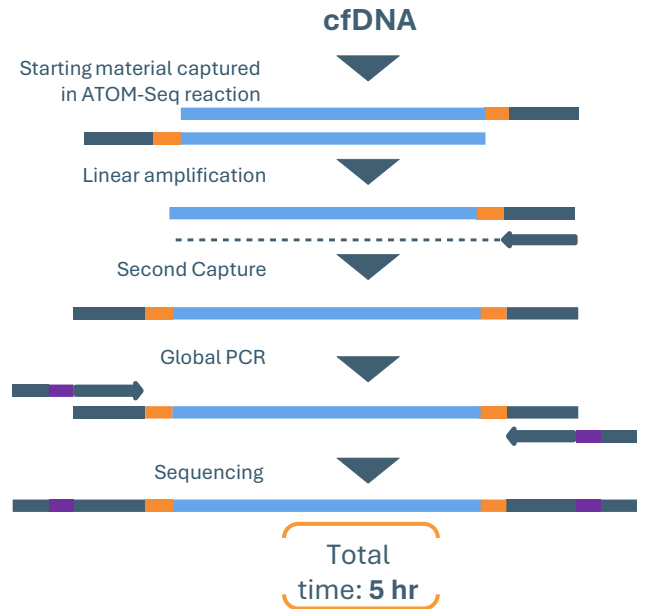


# Total Cell-Free DNA Capture Workflow

NGS library preparation workflows for capturing  
all cell-free DNA for genetic and epigenetic analysis

## Uniquely designed for challenging material

- Simple, ligation-free approach with no DNA end-repair
- Captures all single- and double-strand DNA
- Captures short and degraded material
- Efficient with low input quantities
- Single primer enrichment to maximise capture regardless of DNA breakpoint
- Unique molecular identifies for error suppression
- Minimal bead purification steps



## Workflow benefits



**Higher complexity libraries** as all starting material can be captured

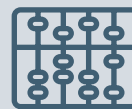


**Process poor quality samples** with resilient workflows able to process even the lowest quality sample



### Flexible workflows for

- Whole genome mutation
- Whole genome methylation
- High-GC whole genome
- Whole-sample sequencing (e.g. sample purity testing)



**Get more from methylation analysis** with linear amplifications, making the workflows resistant to DNA damage

Product Code	Product Name
SEQ001	XCeloSeq cfDNA Library Preparation Kit
SEQ004	XCeloSeq Methyl-cfDNA Library Preparation Kit

Cell-free DNA  
Enrichment Workflow



Total cfDNA Capture  
Workflow



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Custom Assay  
Development

