# SOPHiA DDM™ GlInger Genomic Integrity Solution

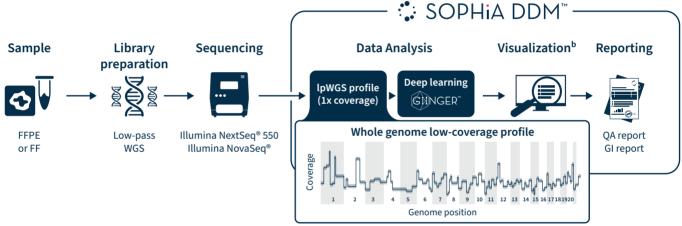
Expand identification of HRD+ cancer samples with a universal application

**SOPHIA DDM™ Glinger Genomic Integrity Solution** is a bioinformatic pipeline that defines the genomic instability (GI) status in ovarian cancer samples.

Powered by the **Glinger™** analytical algorithm, exclusively available on the **SOPHiA DDM™ Platform**, this deep learning-based approach enables you to analyze low-pass Whole Genome Sequencing (WGS) data (~1x coverage) in a decentralized workflow.

The solution offers a **versatile**, **universal** approach to genomic instability measurement that can complement capture-based BRCA assays for a complete homologous recombination deficiency (HRD) assessment without extra sample processing<sup>a</sup>.

### Streamlined bioinformatic workflow



The SOPHiA GENETICS team enabled a **seamless integration** of GlInger™ algorithm into our already validated in-house capture-based library preparation. We are now able to test the Genomic Integrity Index for all our ovarian cancer samples quickly, accurately, and seamlessly.



## Benefits of SOPHiA DDM™ GIInger Genomic Integrity Solution



#### **Maximize insights**

from your low-pass WGS data (1x coverage) with the high analytical performance of the GIInger™ algorithm



### Flexible, adaptable

WGS approach that complements your existing hybrid capture-based BRCA analysis workflow for HRD assessment



#### **Easily interpret**

GI status with a clear PDF report, downloadable from the SOPHiA DDM™ Platform



## Universal solution, robust across multiple wet lab technologies

GII variations across wet lab technologies<sup>a</sup> (n=91)<sup>1</sup> GII variations Sequencer | Library Prep | Workflow -15 -10 10 Illumina - NovaSeq® | Library Prep A | Combined (targeted + lpWGS) Illumina - NovaSeq® | SOPHiA GENETICS - LP3 | Combined (targeted + lpWGS) Illumina - NextSeq® | SOPHiA GENETICS - LP3 | Combined (targeted + lpWGS) Illumina - NextSeg® | Library Prep B | lpWGS Illumina - NovaSeq® | Library Prep C | lpWGS Illumina - NextSeq® | Library Prep A | lpWGS Illumina - NextSeq® | SOPHiA GENETICS - LP2 | lpWGS Illumina - NextSeq® | Library Prep D | lpWGS Illumina - NovaSeq® | Library Prep B | IpWGS Seq. A | SOPHiA GENETICS - LP3 | Combined (targeted + lpWGS) Seq. A | Library Prep E | Combined (targeted + lpWGS) Seq. B | SOPHIA GENETICS LP2 | lpWGS Illumina - NovaSeq® | SOPHiA GENETICS LP3 | lpWGS

GI analysis by GIInger™ is robust across multiple sequencer types, library preps, and workflows.

GIInger™ is also compatible with SOPHiA GENETICS

Universal Library Prep
(ULP), streamlining NGS
workflows with one single protocol across all RUO
SOPHiA DDM™ Oncology applications.

Based on independent analyses of set of candidate samples processed in replicates across different wet lab workflows (n=91). All analyses of SOPHiA DDM™ GIInger Genomic Integrity Solution with other library preparation kits and sequencers other than those defined in the product Instructions for Use (IFU) are for information use only.

# Highly concordant with the SOPHiA DDM™ HRD Solution

Analytical concordance to SOPHiA DDM™ HRD Solution was evaluated using<sup>2</sup>:

- 129 DNA samples from ovarian cancer tissue
- 50 ng DNA input with DQN>3 and tumor content >30%
- Libraries sequenced in combined workflow compared to WGS only<sup>c</sup>

		SOPHiA DDM™ GIInger Genomic Integrity Solution GI Status				
		Positive	Negative	Negative*	Rejected	Inconclusive
SOPHIA DDM™ HRD Solution GI Status	Positive	63	_	_	-	_
	Negative	-	58	-	-	-
	Negative*	_	2	4	-	_
	Rejected	-	1	-	1	-
	Inconclusive	_	_	_	_	_

## **Specifications**

Product type	Dry lab solution - bioinformatic pipeline (reagents not included)			
Sample type	FFPE or FF ovarian cancer tissue			
WGS coverage depth	~1x recommended (0.4x minimum)			
Limit of detection	30% tumor content			
Sequencer compatibility <sup>a</sup>	Illumina NextSeq® 550 and NovaSeq®			
Max. upload size	100 GB <sup>d</sup>			
Product code	DL0121ILLRSM			

Want to know more?

Contact us at:

info@sophiagenetics.com

aSOPHIA DDM™ Glinger Genomic Integrity Solution was verified for use with SOPHIA GENETICS LP3 and Universal Library Prep and Illumina NextSeq® 550 and NovaSeq® only. Please contact local sales representative or info@sophiagenetics.com for further information on use with other wet lab technologies.

Pall results are exclusively provided via downloadable files; 'Libraries prepared using LP3 reagents and sequenced on Illumina® NovaSeq 6000. dData from sequencing runs exceeding 100 GB should be split and uploaded in multiple batches. FFPE, formalin-fixed, paraffin-embedded; FF, fresh-frozen; HRD, homologous recombination deficiency; lpWGS, low-pass whole genome sequencing; QA, quality. '. Data on File. '2. SOPHiA DDM™ Glinger Genomic Integrity Solution. Instructions for Use. v1.1.. SOPHiA DDM™ Glinger Genomic Integrity Solution is for Research Use Only and is not intended for purposes other than research. SOPHiA DDM™ Glinger Genomic Integrity Solution is not for diagnostic, therapeutic, or treatment purposes. All product and company names are trademarks" or registered® trademarks of their respective holders. Use of them does not imply any affiliation with endorsement by them.





