Feature	STRkit	LongTR	TRGT	Straglr	STRdust	Notes
Copy number output		-	✓	✓	-	
Allele size confidence intervals	✓	-	✓	✓	-	
De novo proximate SNV phasing	√ + output	-	✓	-	-	STRkit and TRGT can use heterozygous SNVs to cluster STR alleles; STRkit can output them to VCF.
Existing phased SNV incorporation	-	$\checkmark\checkmark$	-	-	-	
Haplotagged alignment file support	✓	✓	✓	-	✓	All but Straglr can used phased read data (from, e.g., WhatsHap) to call STR alleles.
Methylation handling	-	-	$\checkmark\checkmark$	-	-	
ONT read support	✓	✓	Explicitly forbidden	✓	-	Theoretically works on ONT data, but forbidden by software license.
Read-level data	✓	-	Partial	<b>√</b>	✓	STRkit: JSON output with read-level peak ID + sequence data; TRGT: overlapping reads in BAM; Straglr: TSV output with read-level copy numbers; STRdust: read-level sequence output in VCF.
Copyleft software license	Yes (GPLv3)	Yes (GPLv2)	-	Yes (GPLv3)	Yes (MIT)	TRGT's license restricts it to be only used with PacBio sequencing data, and the software cannot be forked and subsequently re-distributed.
Multi-threading/processing	✓	-	✓	✓	$\checkmark$	
Pre-built Docker image	<b>√</b> √	-	-	-	-	STRkit is available as a pre-built Docker image, which can be incorporated into pipelines/workflow definitions (e.g., Nextflow, WDL).