Accurate Assembly of Circular RNAs with TERRACE

Tasfia Zahin¹

Qian Shi^{1,*}

Xiaofei Carl Zang^{2,*}

Mingfu Shao^{1,2}

¹Department of Computer Science and Engineering, School of Electrical Engineering and Computer Science, The Pennsylvania State University

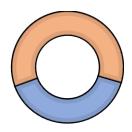
²Huck Institutes of the Life Sciences, The Pennsylvania State University

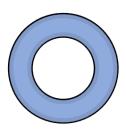


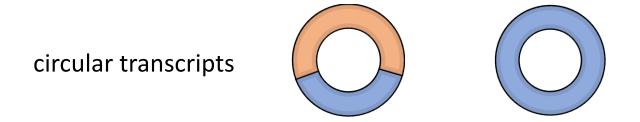


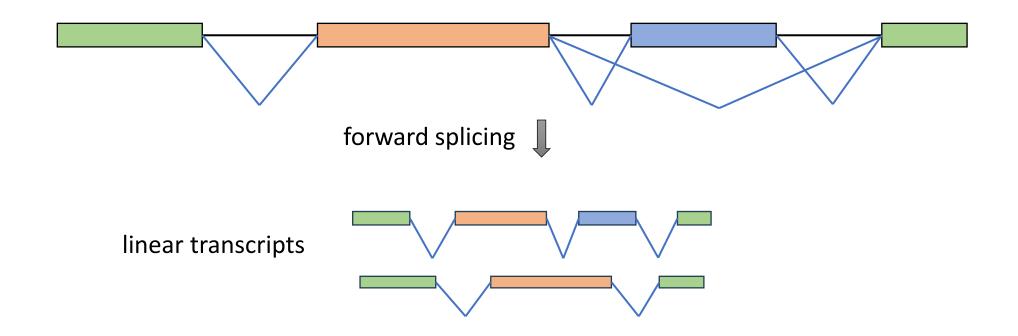


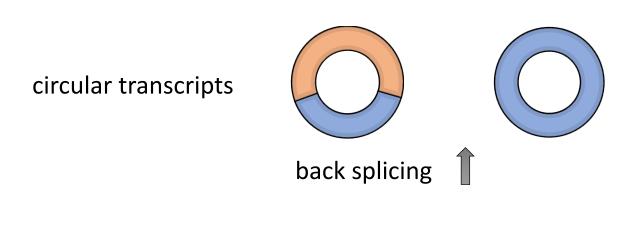
circular transcripts

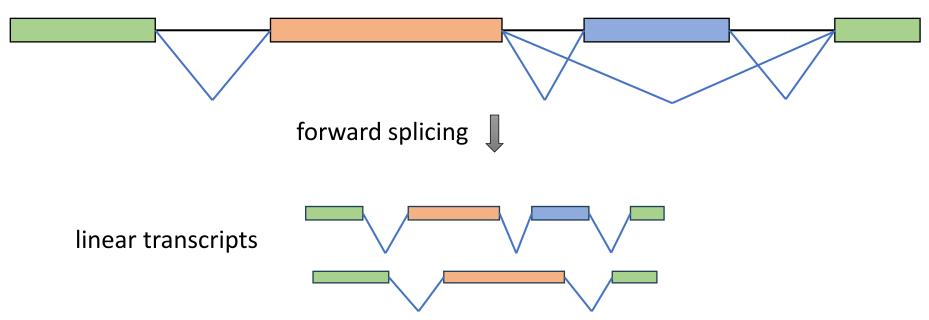


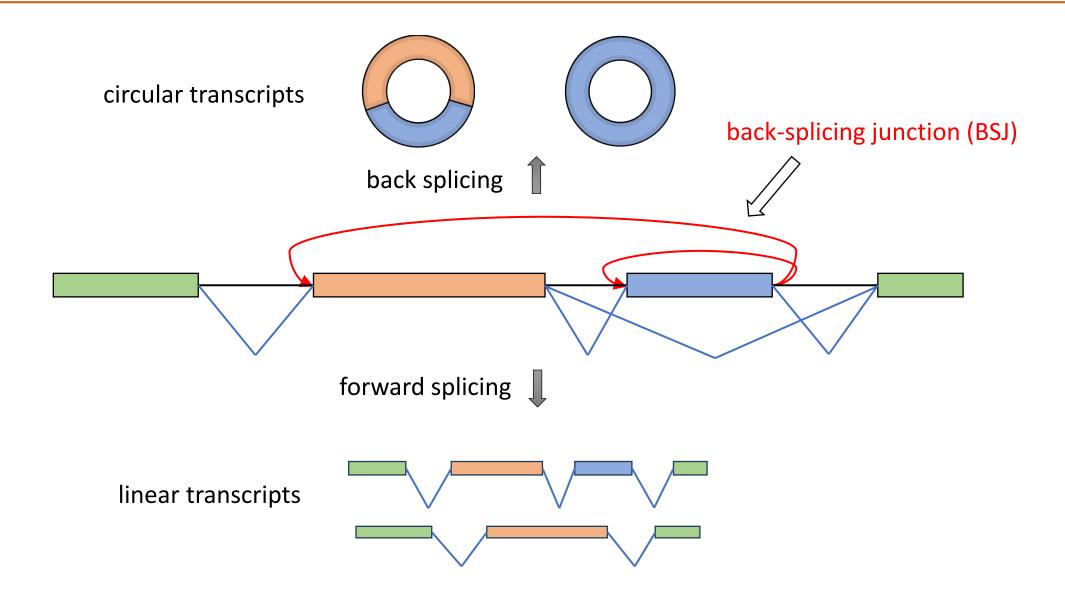












widely expressed in human tissues



More than 60% of human genes express at least one circular RNA

widely expressed in human tissues



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very stable in nature

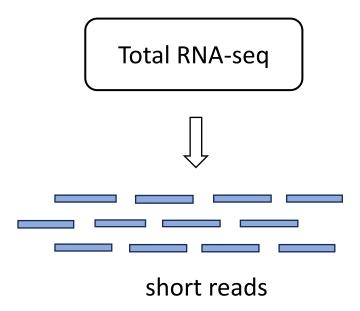


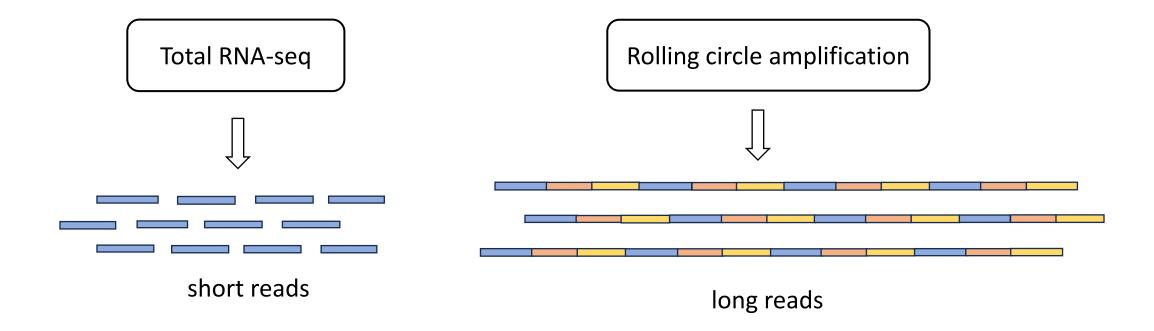


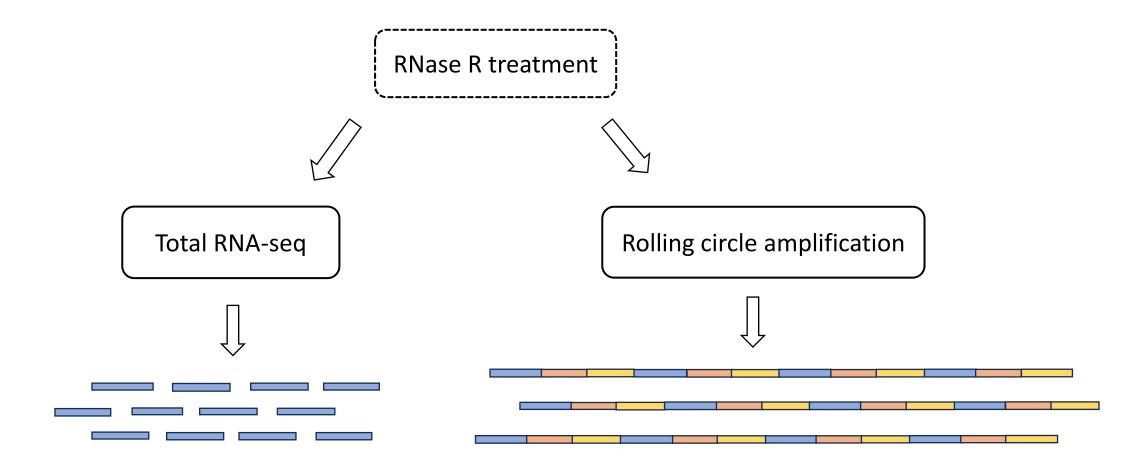


Circular RNAs have at least 2.5 times longer half-life than linear RNAs in mammary cells

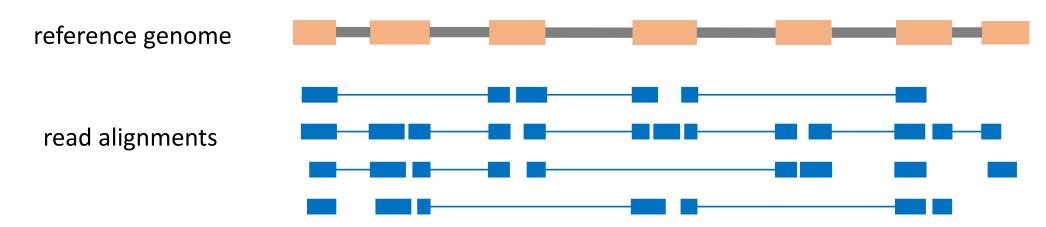
More than 60% of human genes express widely expressed in human tissues at least one circular RNA very stable in nature Circular RNAs have at least 2.5 times longer half-life than linear RNAs in mammary cells circ-Foxo3 interacts with certain proteins reliable disease biomarkers to form a complex and inhibits cell cycle diagnostic and therapeutic targets progression in cancer



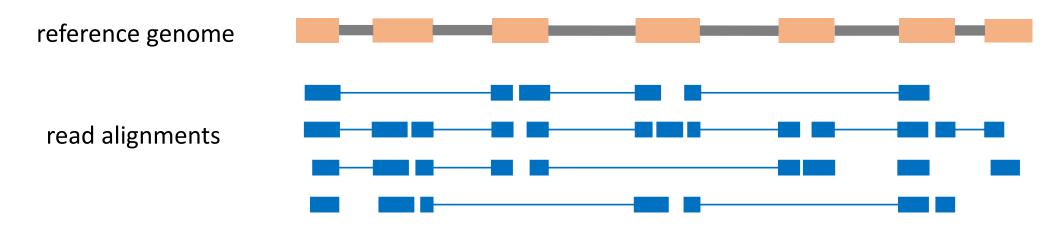




Input: a set of paired-end total RNA-seq reads aligned to a reference genome

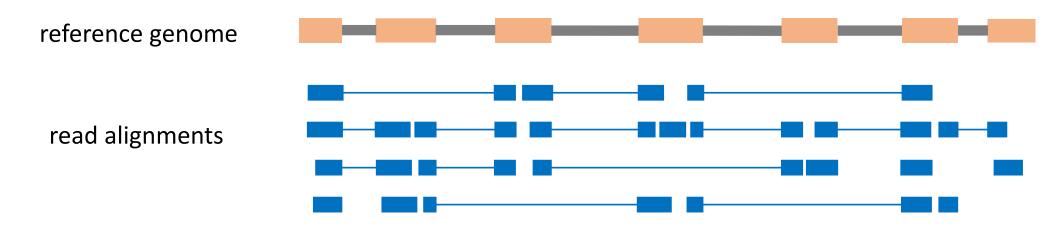


Input: a set of paired-end total RNA-seq reads aligned to a reference genome



Optional: reference gene annotation

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Optional: reference gene annotation

Goal: reconstruct a set of full-length circular transcripts



Several computational methods exist for circular RNA detection, however:

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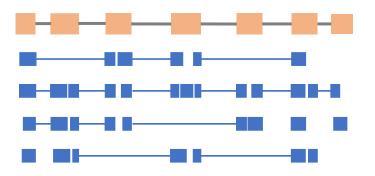
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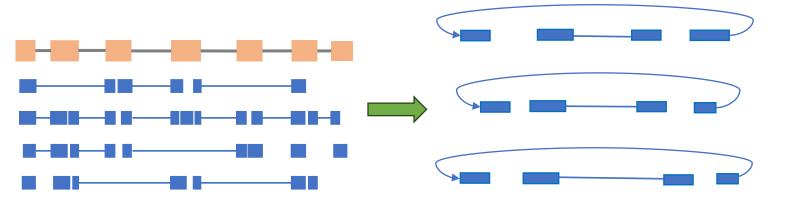


Develop a new circular RNA assembler

- does full length detection
- does not rely on reference annotation
- achieves high assembly accuracy

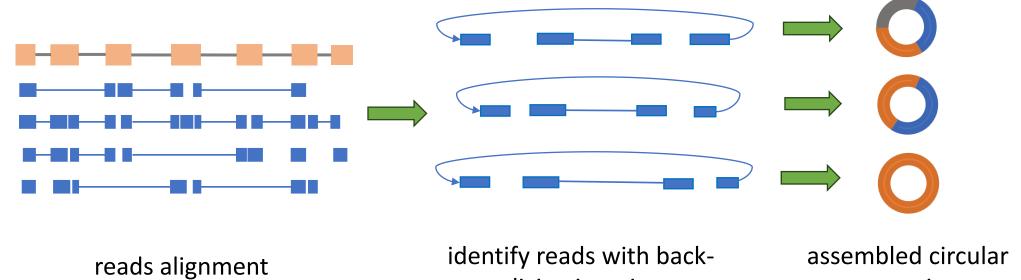


reads alignment



reads alignment

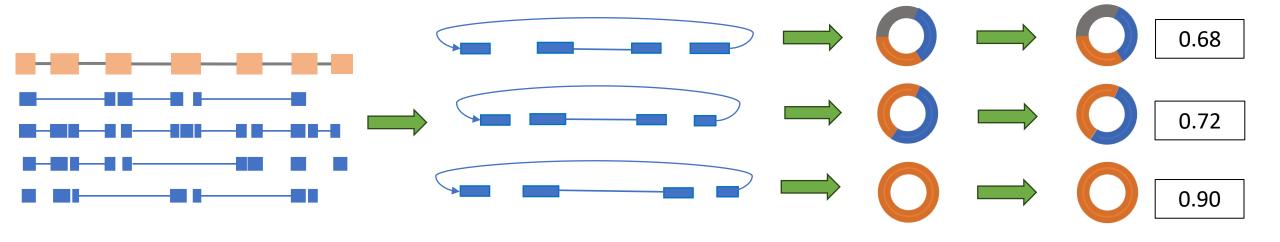
identify reads with backsplicing junctions



splicing junctions

transcripts

reads alignment



assembled circular

transcripts

identify reads with back-

splicing junctions

27

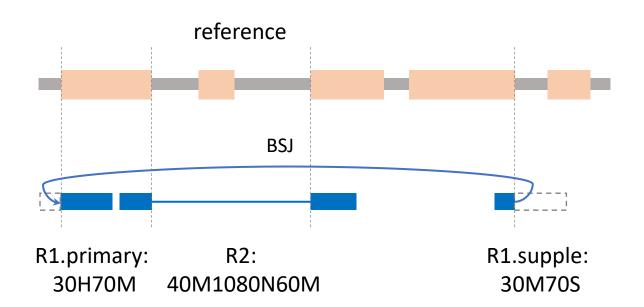
scored circular

transcripts

Identifying Back-spliced Reads

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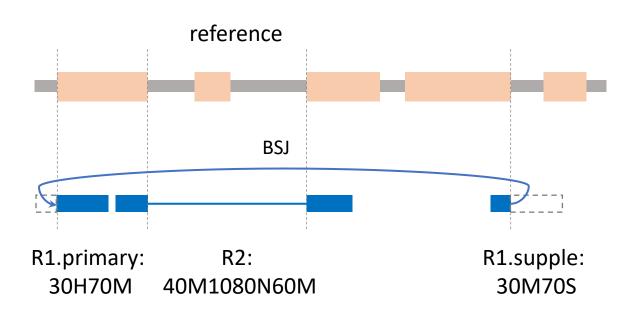
From chimeric alignments

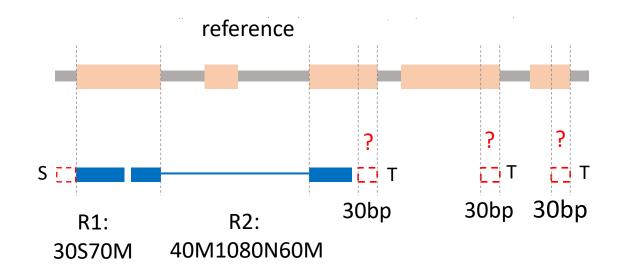


Identifying Back-spliced Reads

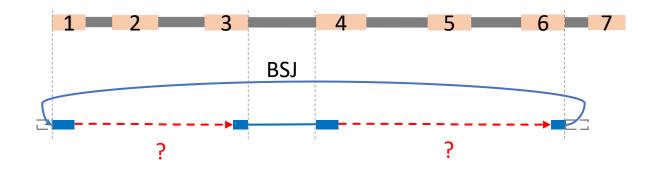
From chimeric alignments

From new junction mapping algorithm

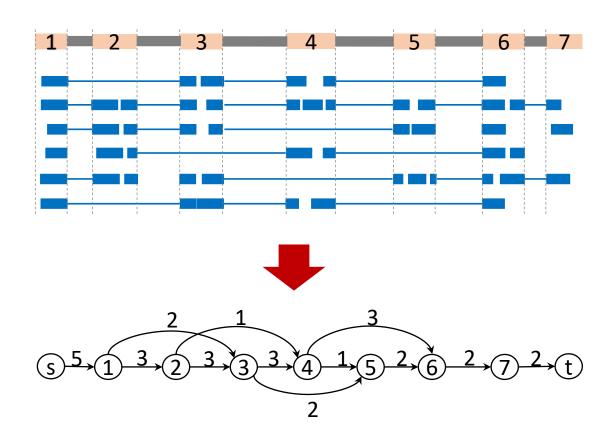


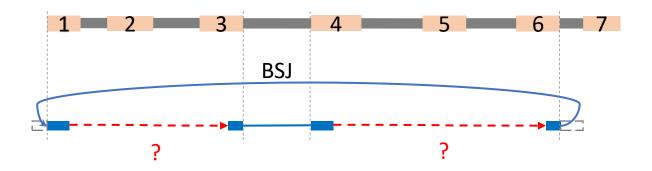


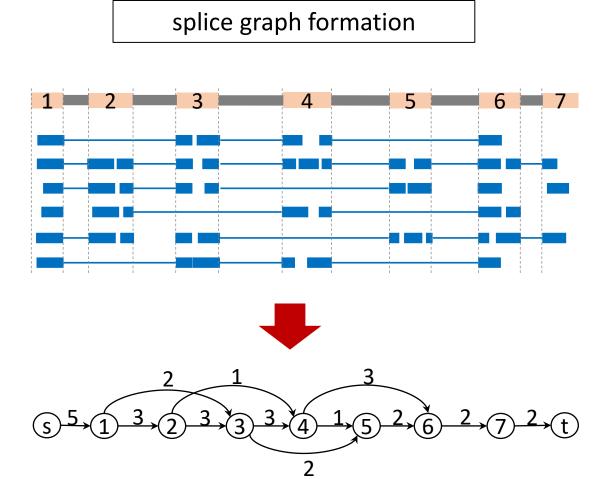
Jaccard index (S, T) > 0.9



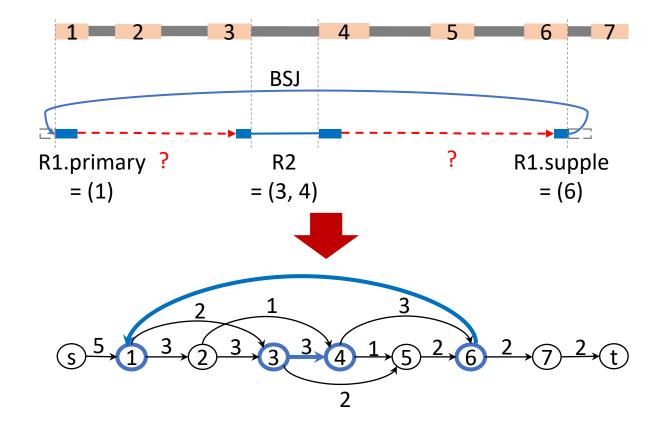
splice graph formation





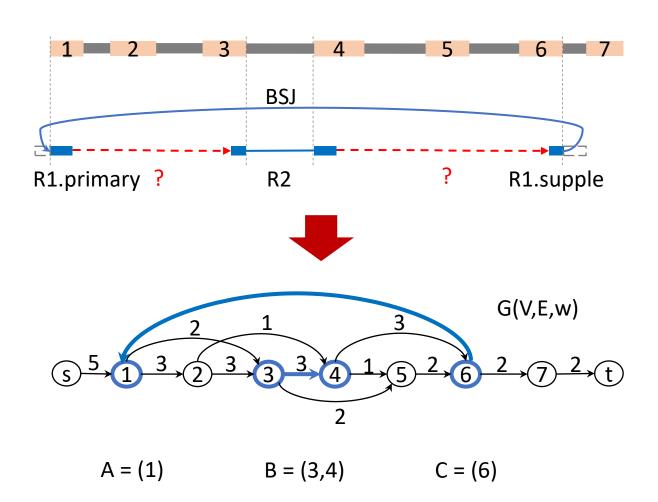


back-spliced read represented by 3 paths in splice graph



Formulation and Algorithm for Bridging

Given: G(V,E,w), $A = (a_1, a_2, ..., a_i)$, $B = (b_1, b_2, ..., b_j)$, and $C = (c_1, c_2, ..., c_k)$

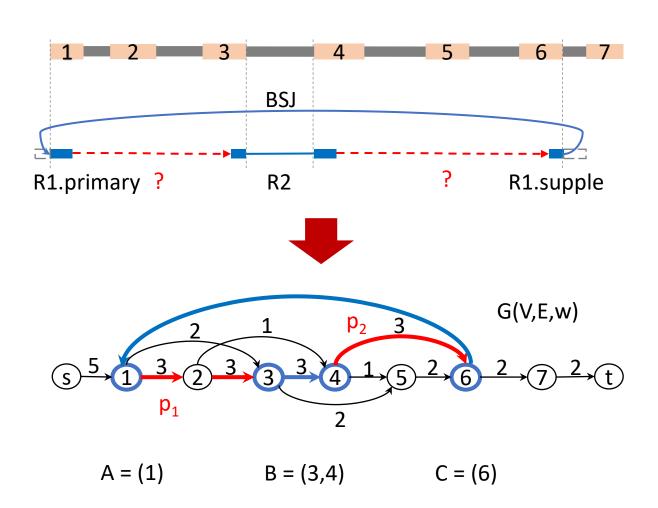


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score(p): smallest weight over all the edges in p.



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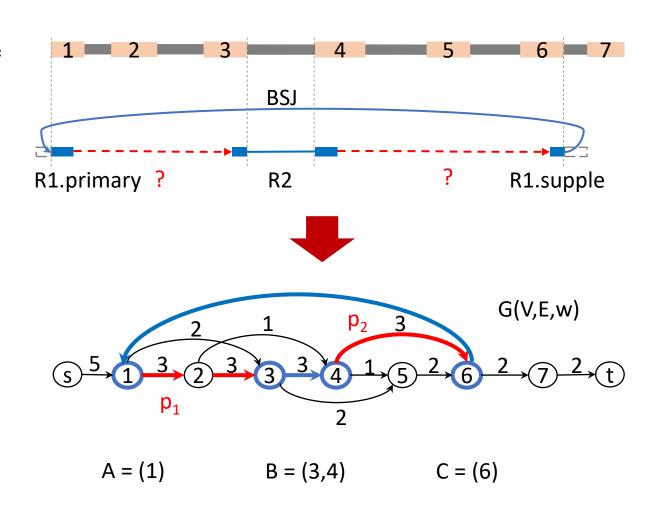
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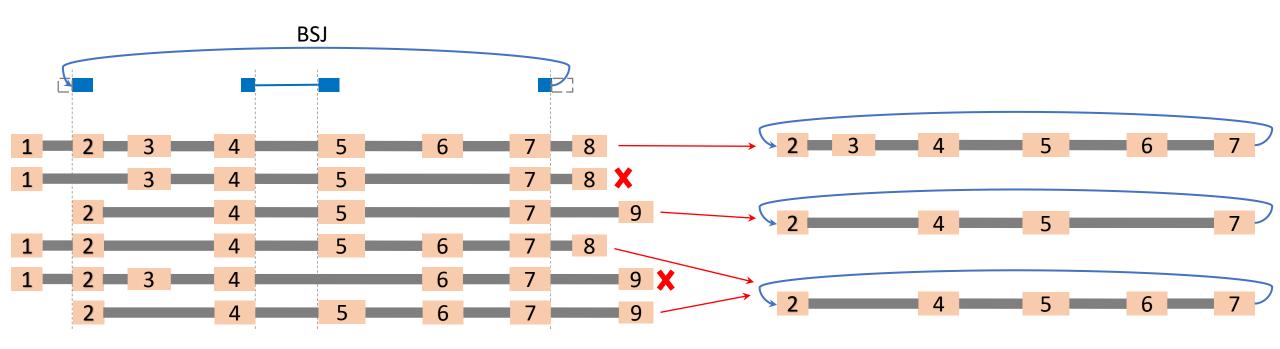
Adopt an efficient dynamic programming algorithm $(O(|V|^2 \cdot |E|))$ to find optimal p_1 and p_2 , previously proved to effective for linear transcript assembly.

Collect the top 10 optimal bridging paths (P).



Use of Reference Annotation

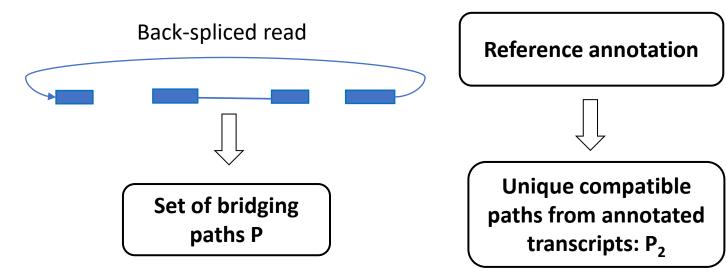
• Set of circular paths from reference annotation, if provided (P_2) .



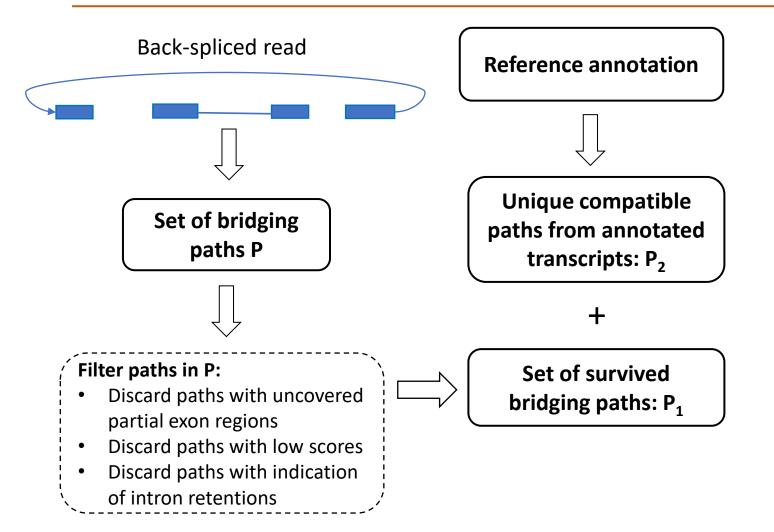
annotated linear transcripts

compatible full-length circular paths

Selection of Candidate Paths



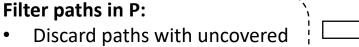
Selection of Candidate Paths



Selection of Candidate Paths

Back-spliced read

Set of bridging paths P



- partial exon regions
 Discard paths with low scores
- Discard paths with indication of intron retentions

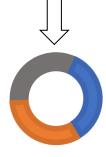
Reference annotation

Unique compatible paths from annotated transcripts: P₂

Set of survived bridging paths: P₁

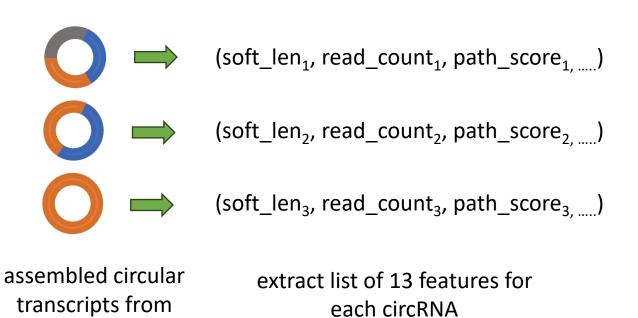
Selection Criteria:

- If $P_1 \cap P_2 \neq \emptyset$, pick the path in $P_1 \cap P_2$ with maximum score.
- If $P_1 \cap P_2 = \emptyset$ and $P_1 \neq \emptyset$, pick the path in P_1 with maximum score.
- If P₁ = Ø and P₂ contains a single path, pick that path.
- If P₁ = Ø and P₂ contains multiple path, discard the back-spliced read.
- If P₁ U P₂ = Ø, discard the back-spliced read.



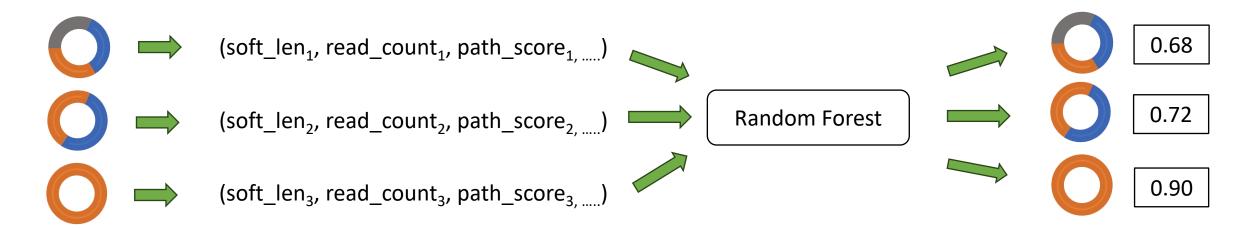
Scoring Assembled Transcripts

Scoring Assembled Transcripts



one sample

Scoring Assembled Transcripts



assembled circular transcripts from one sample

extract list of 13 features for each circRNA

train a machine learning model and test all samples

assign confidence scores to circRNAs

Experimental Setup

Datasets: Short-read total RNA-seq datasets of 8 human tissues (accession number PRJCA000751 from BIGD)

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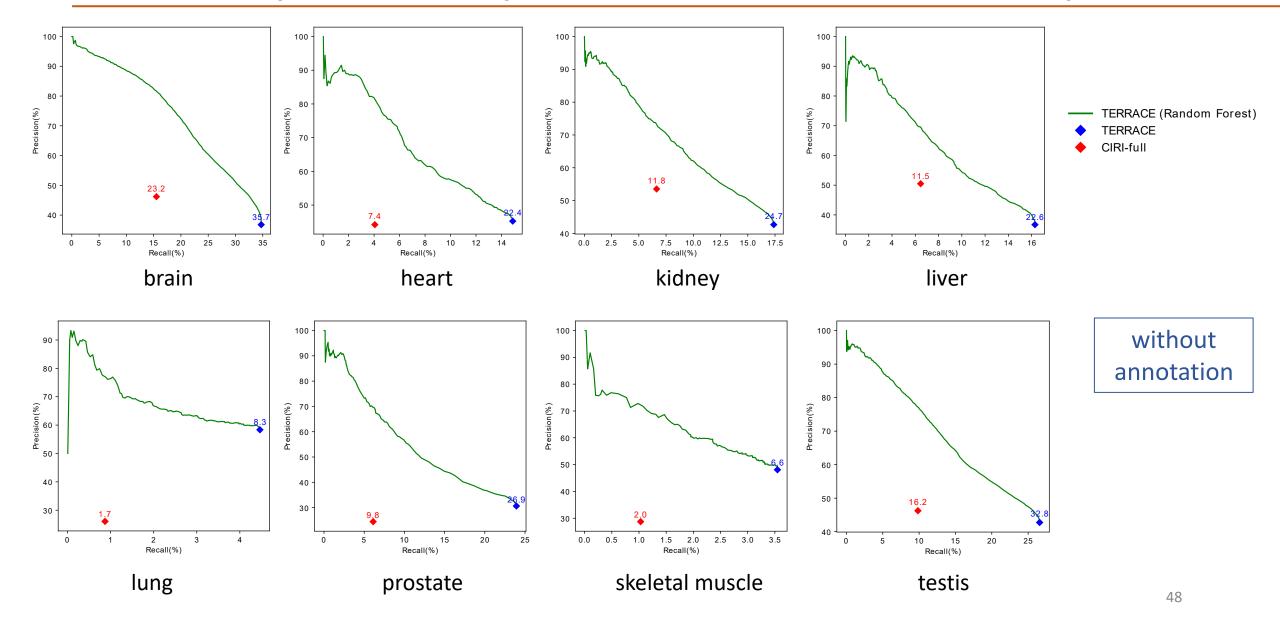
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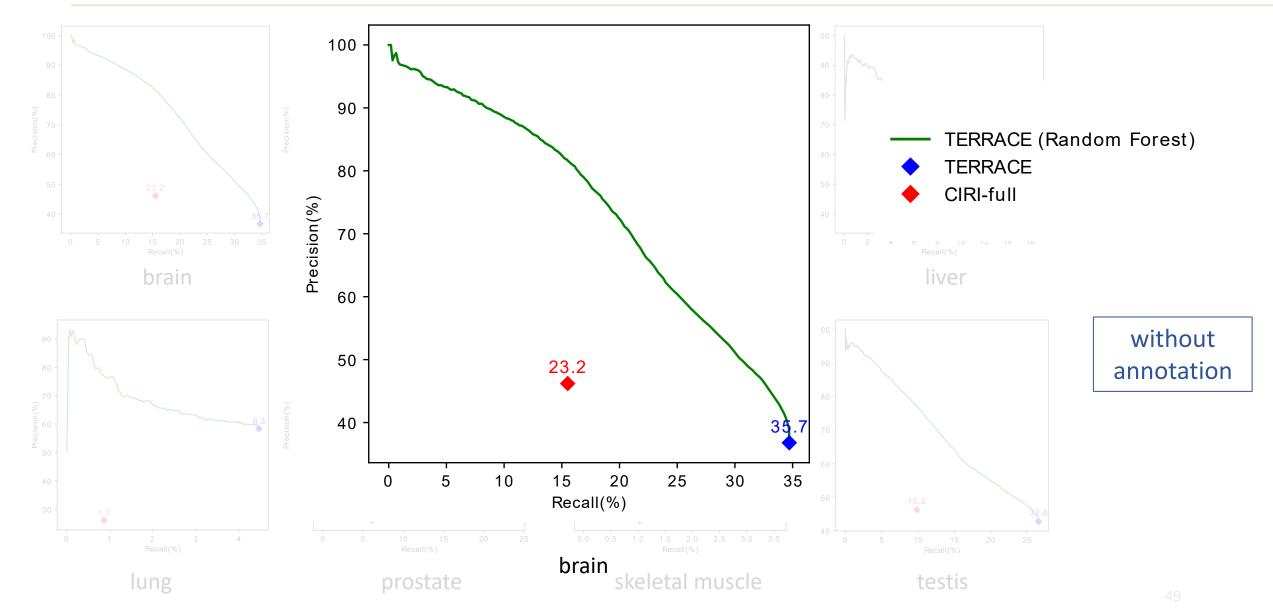
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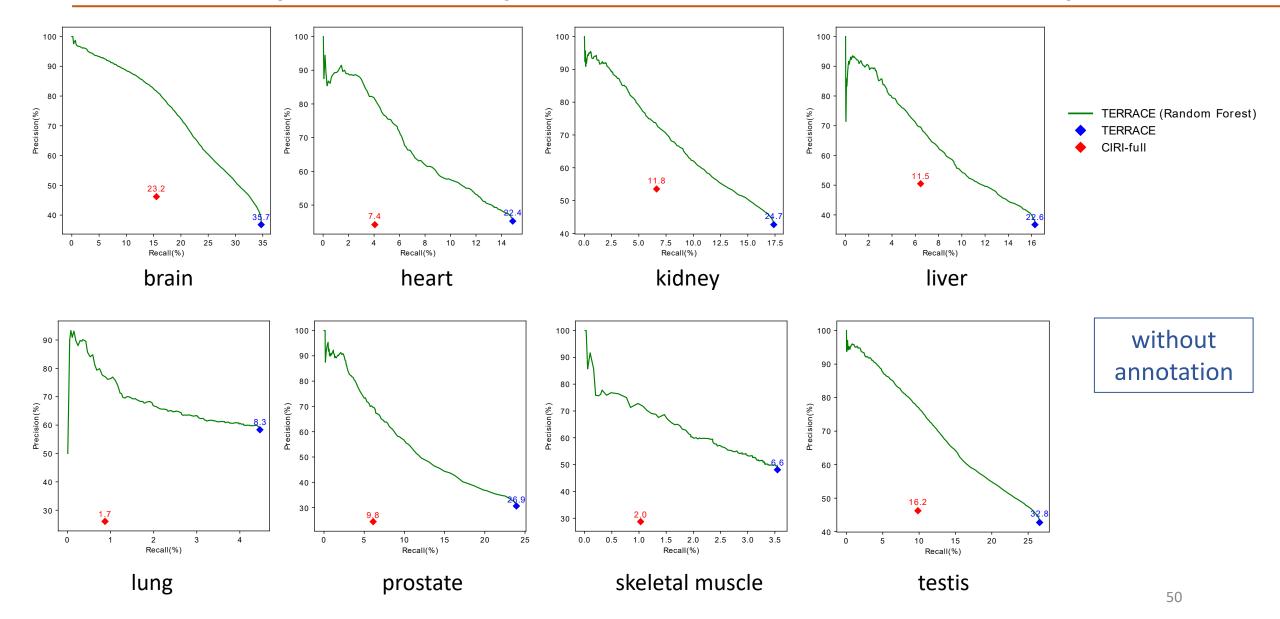
Methods Compared: CIRI-full, CIRCexplorer2, CircAST

Evaluation: number of matching transcripts with ground truth

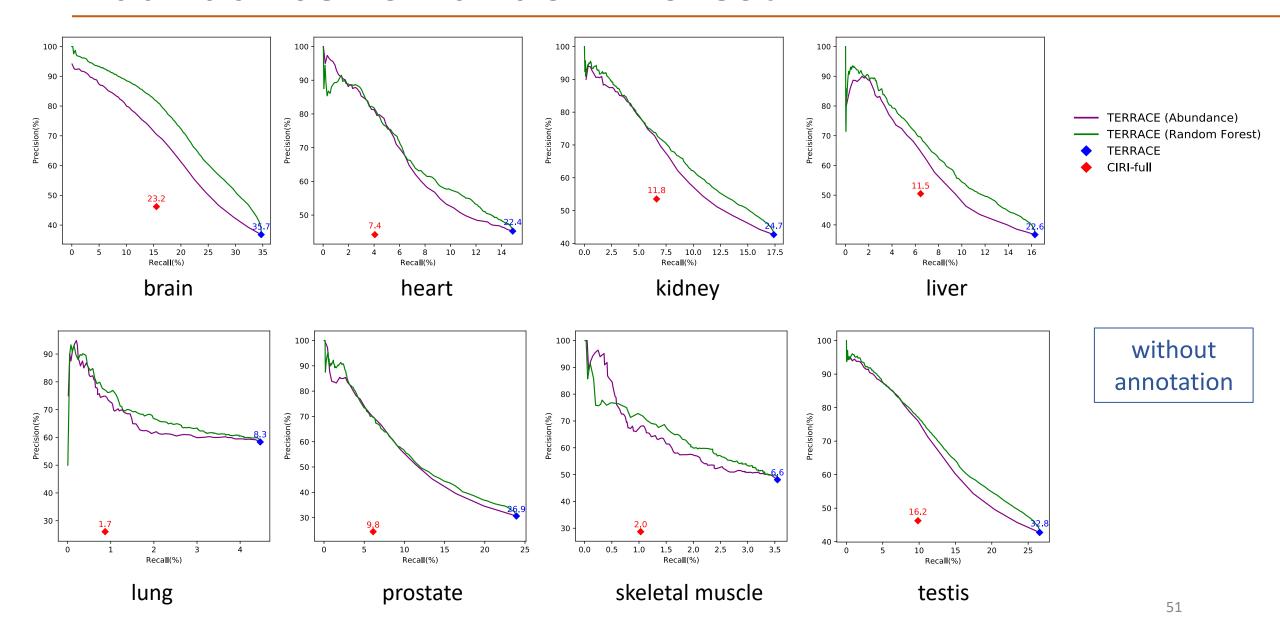
Metric: F-score, precision-recall curve



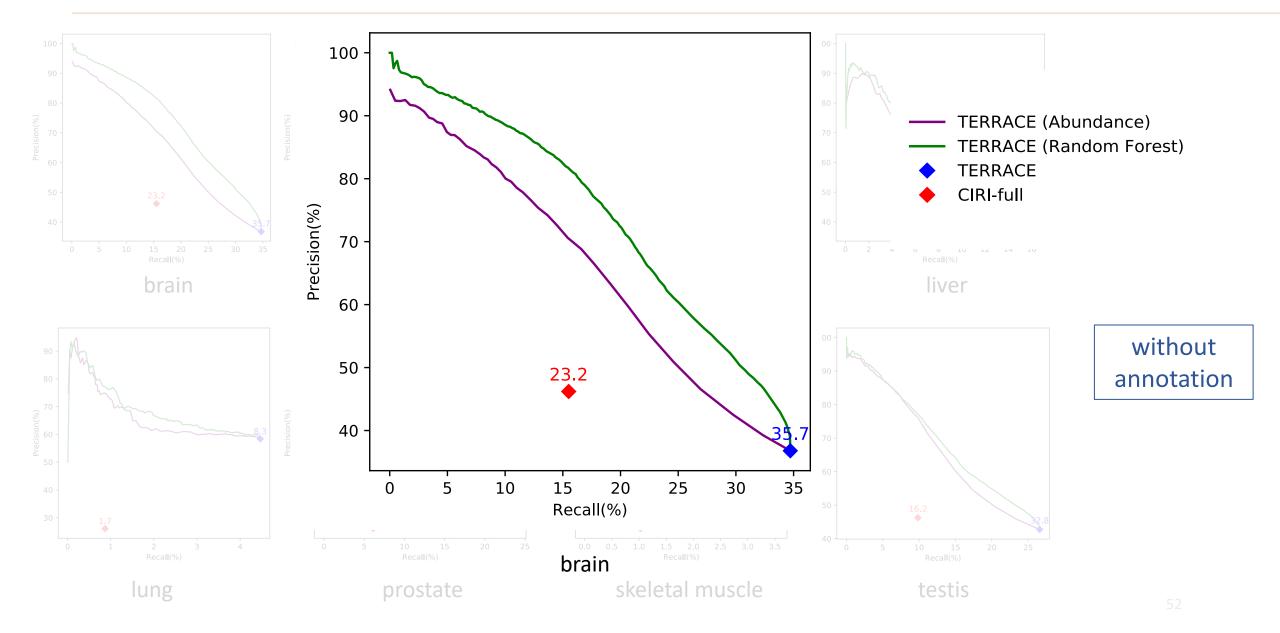




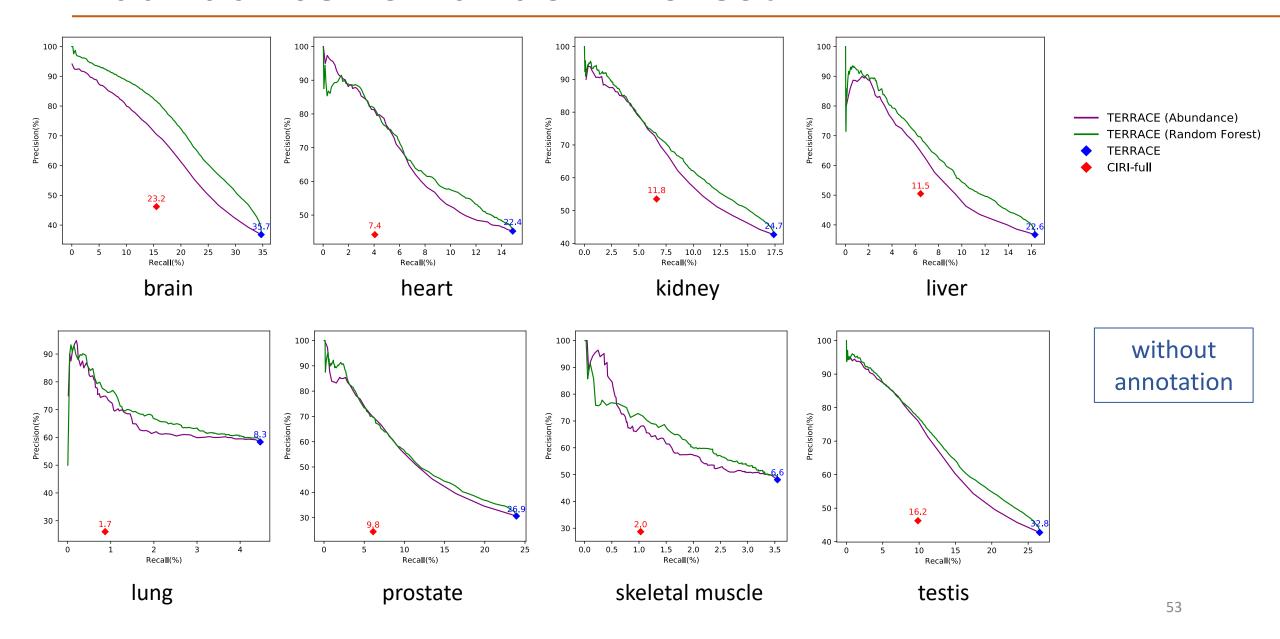
Abundance vs Random Forest

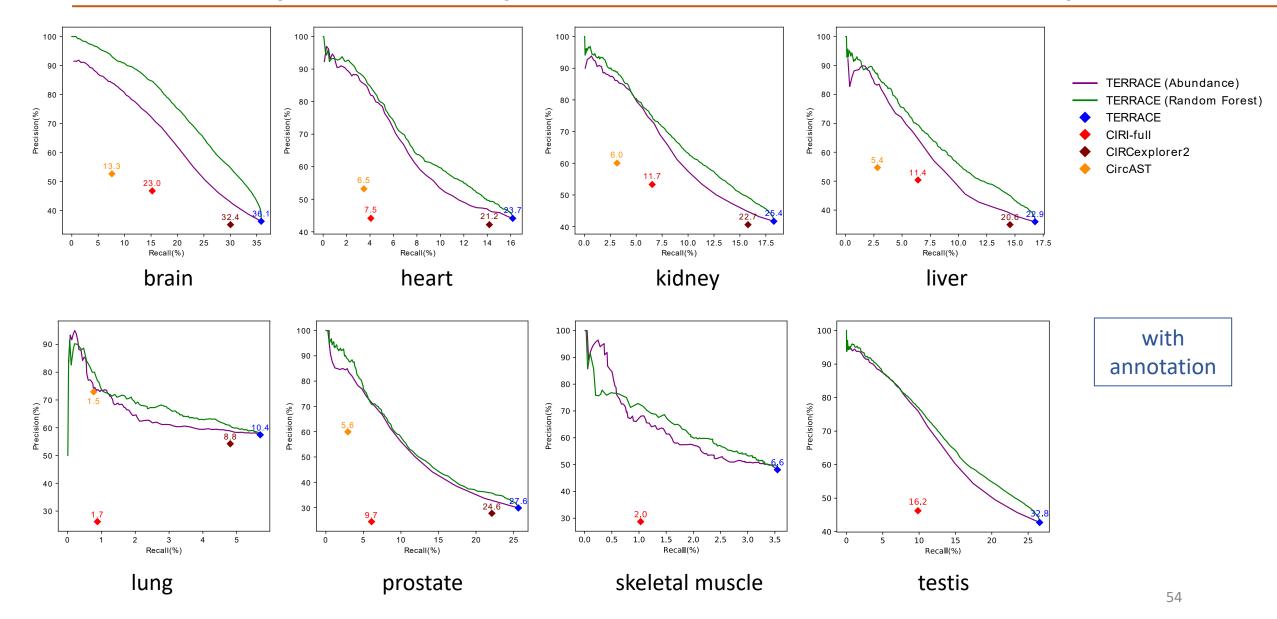


Abundance vs Random Forest

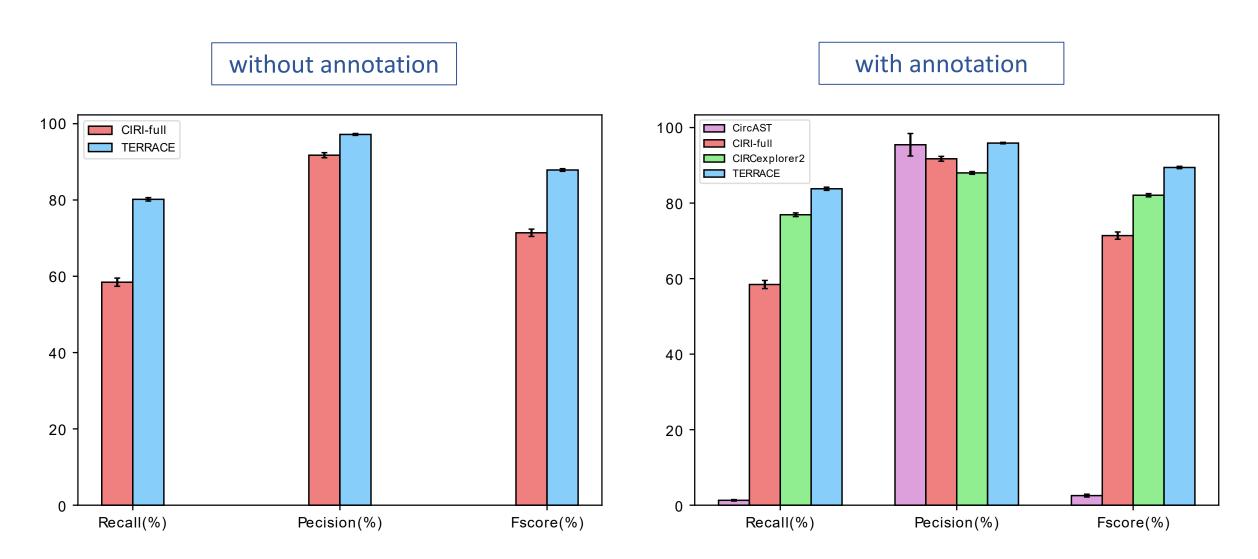


Abundance vs Random Forest





Assembly Accuracy on Simulated Data



Summary

- We present TERRACE, a full-length circular RNA assembler.
- Four algorithmic innovations: identifying accurate back-spliced reads, formulating assembly into bridging, designing new heuristics for path selection, assigning confidence scores to circRNAs.
- TERRACE allows accurate detection of circular RNAs without requiring reference gene annotation, particularly useful for species lacking well-annotated transcriptomes.
- Tool availability: https://github.com/Shao-Group/TERRACE.

Acknowledgements

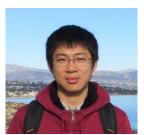
Co-authors

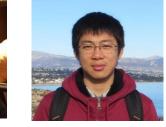


Qian Shi



Carl Zang





Mingfu Shao

Funding support



DBI-2019797

and 2145171



R01HG011065

Members of Shao Group







Xin Yuan



Qimin Zhang



Xiang Li



Zhezheng Song



Abhishek Talesara



Saadya Rao

Thank You!

Run Time and Memory Usage

CPU time (minutes) for different tools on human tissues

sample	methods w/o annotation		methods with annotation					
	CIRI-full	TERRACE	CIRCexplorer2	CircAST	CIRI-full	TERRACE		
Lung	265	29	0.14	249	266	26		
Brain	471	42	0.21	1234	489	45		
Skeletal	334	40	0.1	260	337	50		
Heart	380	32	0.1	488	389	35		
Testis	427	39	0.14	922	472	40		
Liver	330	31	0.08	423	345	31		
Kidney	352	31	0.1	554	370	32		
Prostate	253	52	0.24	229	307	57		
Average	352	37	0.14	545	372	40		

Peak memory usage (GB) for different tools on human tissues

sample	methods w/	o annotation	methods with annotation					
	CIRI-full	TERRACE	CIRCexplorer2	CircAST	CIRI-full	TERRACE		
Lung	10.1	16.9	0.14	0.07	11	16.7		
Brain	80.4	11.2	0.16	.16 0.11		11.4		
Skeletal	22.2	19.8	0.14	0.1	22.8	20.2		
Heart	30.8	10.9	0.14	0.61	31.4	11.1		
Testis	74.8	5.5	0.15	0.09	76.2	5.6		
Liver	271.9	17.7	0.14	0.08	31.7	17.9		
Kidney	32	10.3	0.14	0.08	33.1	10.1		
Prostate	21.7	23.4	0.15	0.08	22.4	23.6		
Average	68	14.5	0.15	0.15	38.7	14.6		

pAUC (without annotation)

	TERI	RACE vs CIRI-fi	ull	TERRACE vs. CIRI-full				
sample	cons	trained by rec	all	constrained by precision				
	TERRACE	CIRI-full	Δ%	TERRACE	CIRI-full	Δ%		
Lung	65.2	52.3	24.6	66.8	14.3	365.9		
Brain	1380.1	1047.8	31.7	969.7	339	185.9		
Skeletal	75.7	62.5	21	59.9	17.3	246		
Heart	347.6	294.4	18	334.2	113	195.5		
Testis	851.9	645.4	31.9	618.4	196.4	214.9		
Liver	527.1	450.4	17	255.9	136.8	87		
Kidney	553.9	483.3	14.6	267.9	133.8	100.1		
Prostate	495.7	364.1	36.1	598	161.4	270.3		

Statistics

sample	# reads	#circRNAs	w/o annotation				With annotation			
			TERRACE		CIRI-full		TERRACE		CIRCexplorer2	
			# detected	# correct	# detected	# correct	# detected	# correct	# detected	# correct
Lung	87M	18136	1388	810	606	158	1798	1033	1608	872
Brain	82M	35801	33785	12428	12024	5553	35365	12835	30611	10754
Skeletal	93M	10908	805	387	390	112	1053	494	983	434
Heart	79M	11223	3692	1670	1032	456	4113	1815	3770	1591
Testis	90M	42633	26509	11333	9070	4195	27329	11603	21740	9188
Liver	87M	11978	5314	1951	1533	774	5588	2010	4989	1744
Kidney	93M	22521	9176	3915	2791	1494	9869	4115	8747	3554
Prostate	83M	8114	6342	1942	2029	496	6973	2081	6469	1794