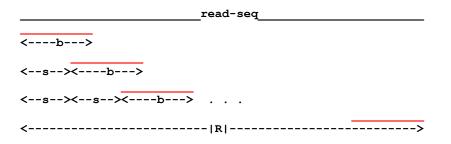
## Supplementary Text 1. Procedure for choosing b-mer length, and BF loading and querying

**Loading Bloom filter.** In this stage, all target sequences in the target set  $T = \{T_1, T_2, ...\}$  are scanned using a sliding substring of length b which is called b-mer. For each target sequence  $T_i$ , all possible  $|T_i| - b + 1$  b-mers are scanned and then inserted to the Bloom filter after specifying the corresponding bit vector positions by computing the k hash values for each b-mer.

	target-seq
<b></b>	
<b+1></b+1>	
<b+2></b+2>	
<	T >

**Querying Bloom filter.** In this step, all reads in the read set  $R = \{R_1, R_2, ...\}$  are queried using the same *b*-mer in the loading stage. If at least one hit is found for a read, the read is dispatched to the corresponding node. We can use sliding *b*-mer windows (with step size of one base, s = 1) or jumping *b*-mer windows (with step size greater than one base) to interrogate each read  $R_j$  as explained below.



Suppose that the minimum seed or exact match length to report a candidate hit for an aligner is l. We choose  $b \leq l$  and then load the Bloom filters as mentioned in the *Loading Bloom filter* stage. In the querying stage, the *b*-mers of each read sequence is interrogated against all Bloom filters from different partitions. If b = l, the reads are scanned using sliding window with step size of one base, i.e. s = 1. By choosing b < l, the interrogation step will be faster but with more extra dispatched reads, and the step size is computed as follows to cover all possible seeds or exact matches of length l between the target sequence  $T_i$  and read sequence  $R_j$ . If the *l*-mer starting at position p is covered by at least one *b*-mer, to cover the next *l*-mer starting at position p + 1 we should have  $s + b \leq l + 1$ . Therefore,  $s \leq l - b + 1$ .

