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1: Read Mapping

Read Mapping: Mapping billions of DNA fragments (reads) against a reference genome to identify genomic variants

- Requires approximate string matching
- Computationally expensive alignment using **quadratic-time** dynamic programming algorithm
- Bottlenecked by memory bandwidth**

Three general types of read mappers:

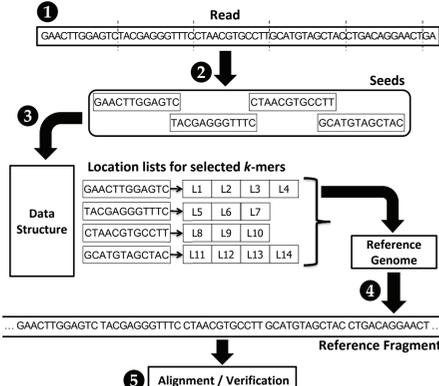
- Suffix-array based mappers
- Hash table based mappers
- Hybrid

2: Hash Table Based Mappers

Seed-and-extend procedure to map reads against a reference genome allowing e indels. They have:

- High **sensitivity** (can tolerate many errors)
- High **comprehensiveness** (can find more mappings)
- BUT**
- Low **speed**

The most recent fastest hash table based read mapper, **mrFAST with FastHASH** [Xin+, BMC Genomics 2013]



3: Problem

For lower runtimes, **location filters** can efficiently determine whether a candidate mapping location will result in an **incorrect mapping before** performing the computationally **expensive incorrect verification** by alignment. They should be **fast**.

4: Our Goal

Design and implement a new filter that **rejects incorrect mappings** before the alignment step

- Minimize the occurrences of unnecessary alignment
- Maintain **high sensitivity and comprehensiveness**
- Obtain **low runtime** and **low false positive rate**

Accelerate read mapping by overcoming memory bottleneck with **3D-stacked memory** and its **PIM** for data-intensive computation

- Very fast parallel operations on big data sets **near memory**

6: GRIM-Filter Mechanism

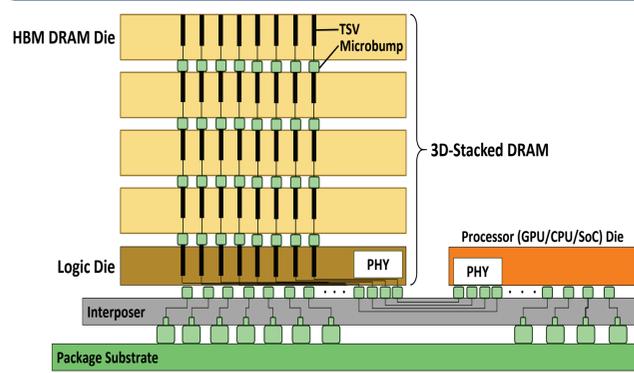
GRIM-Filter is based on two key ideas:

- Introduce **parallelism** to **q-gram string matching**
- Utilize a 3D-stacked DRAM to **alleviate** the memory bandwidth issue of our algorithm and **parallelizes** most of the filter.

GRIM-Filter has two main steps:

- Precomputation:** Divide the reference genome into consecutive bins and generate *existence bitvectors* for each bin.
- Filtering Algorithm:** **Filter** locations by quickly determining whether a read can map to a specific segment of the genome.

5: 3D-Stacked Logic-in-Memory DRAM



Recent technology that tightly couples memory and logic vertically with very high bandwidth connectors.

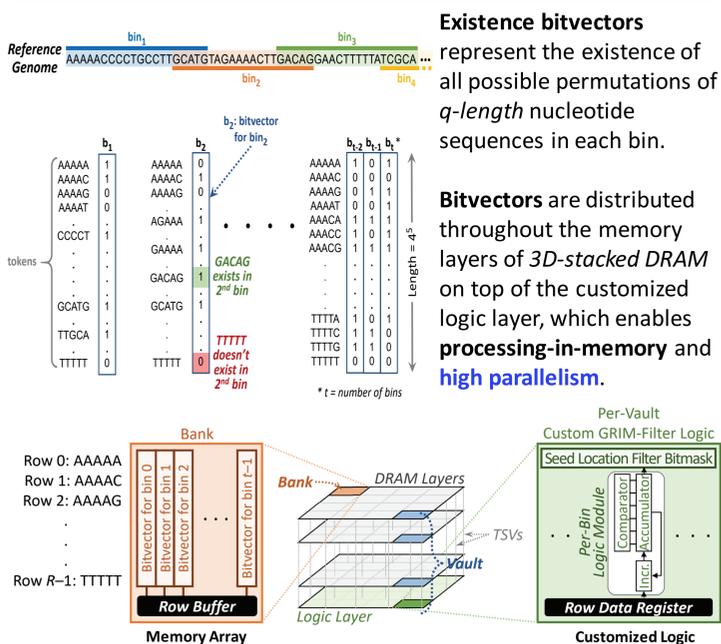
Numerous Through Silicon Vias (TSVs) connecting layers, enable **higher bandwidth** and **lower latency** and **energy consumption**.

Customizable logic layer for application-specific accelerators.

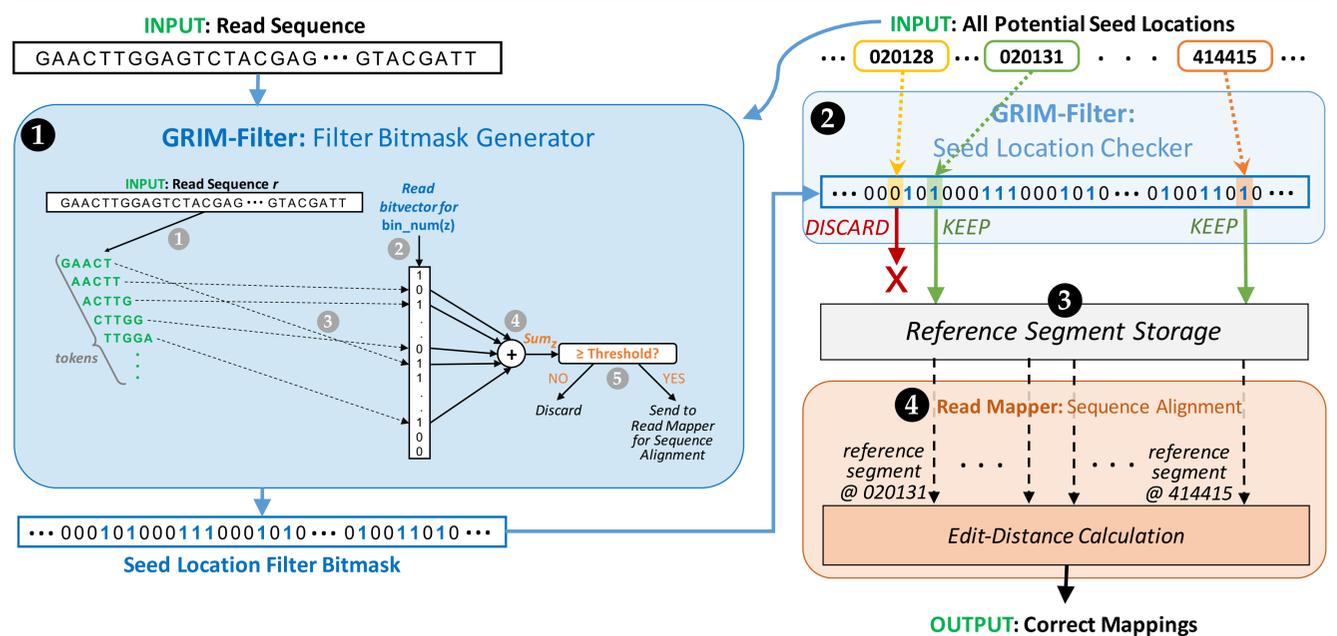
Logic layer enables **fast, massively parallel operations** on large sets of data, and provides the ability to run these operations **near memory** to alleviate the memory bottleneck.

Processing in 3D-stacked memory is extremely good at accelerating embarrassingly parallel simple bit operations.

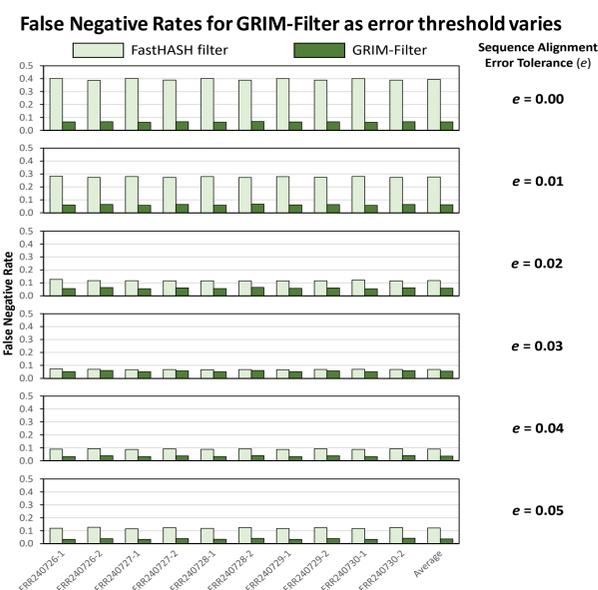
7: Bins & Bitvectors



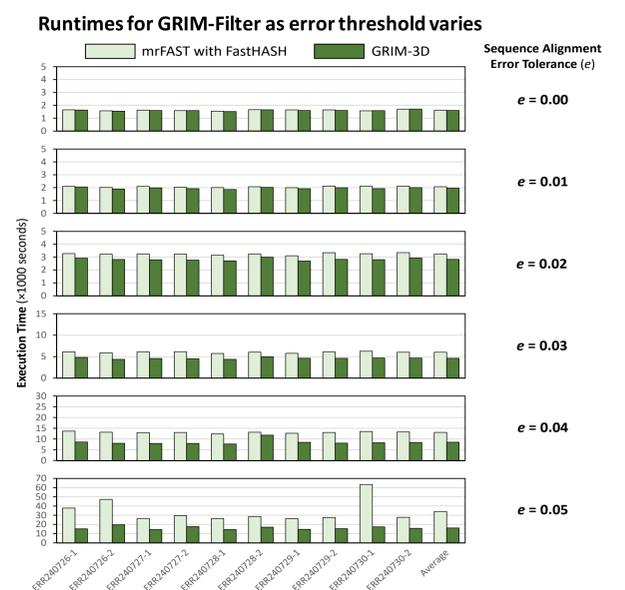
8: GRIM-Filter Walkthrough



9: Results & Conclusion



- Baseline: **mrFAST with FastHASH** mapper code [Xin+, BMC Genomics 2013]. However, GRIM-Filter is **fully complementary** to other mappers, too.
- Key Results of GRIM-Filter:**
 - 5.59x-6.41x less false negative locations**, and
 - 1.81x-3.65x end-to-end speedup** over the state-of-the-art read mapper mrFAST with FastHASH.
- We show the inherent **parallelism** of our filter and **ease of implementation** for **3D-stacked memory**. There is great promise in adapting DNA read mapping algorithms to state-of-the-art and emerging memory and processing technologies.
- Other Results:**
 - Examined sensitivity to **number of bins**: 450x65536
 - Examined sensitivity to **q-gram size**: 5
 - Found to be the best tradeoff between **memory consumption**, **filtering efficiency**, and **runtime**.



*In the figures shown, smaller bars indicate better performance