# Nanopore automata

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#### 1 Abstract

State machine algorithms for aligning Nanopore reads. Initial goal is simple reusable code for aligning a nanopore read to a reference sequence. No attempt at optimization yet.

# 2 Specification

#### 2.1 Parameterization algorithm

Given the following inputs

- Reference genome (FASTA)
- Segment-called reads (FAST5/HDF5)

Perform the following steps

• Perform Baum-Welch to fit a rich model

Rich model incorporates segment statistics.

#### 2.2 Reference search algorithm

Given the following inputs

- Reference genome
- $\bullet$  Segment-called reads (FAST5/HDF5)
- Parameterized rich model

Perform the following steps

• Perform Viterbi alignment

### 2.3 Implementation

Libraries etc.

 $\mathrm{HDF}5...$ 

#### 2.4 Evaluation

Strategy...

Data sets...

#### 3 Methods

Model & inference algorithms.

#### 3.1 Model

- $\bullet$  Order-N transducer.
- Input: nucleotide
- Output: nucleotide, segment mean, duration
- Emissions:
  - categorical (base k-mer)
  - mixture of Normal/gamma (mean/duration)
- Transitions:
  - Match: emit single segment, absorb 1 base
  - Insert: affine gap insertion of bases: emits segments, absorbs no bases
  - Delete: affine gap deletion of bases: emits no segments, absorbs bases
  - Merge: emit single segment, absorb 2 or 3 bases

- Split: emit single segment, absorb 0 bases
- Skip: emit single segment, absorb  $2 \dots K$  bases (large K, low extension penalty)

This can be achieved by a Mealy transducer with  $3 \times 4^N$  states. The factor of  $4^N$  accounts for the order-N context. For each such context, the three states are MAT, INS and DEL.

#### Parameters:

- Gap opening & extension probabilities  $\theta_{go}$ ,  $\theta_{gx}$
- Merge probability  $\theta_{mo}$ , probability that it's a 3-merge is  $\theta_{mx}$
- Split probability  $\theta_s$
- Skip probability  $\theta_{ko}$ , skip extension probability  $\theta_{kx}$

The transition table is as follows:

Transition	From	To	Weight	Input $x$	Output $(y, m, d)$
Match	MAT	MAT	$(1 - \theta_{go})(1 - \theta_{mo})(1 - \theta_s)(1 - \theta_{ko})$	$x\in \Omega$	$y \sim \text{Categorical}(\mathbf{p}_{x,c}^m),$
					$m \sim \text{Normal}(\mu_{x,c}^m, \tau_{x,c}^m),$
					$d \sim \operatorname{Gamma}(\alpha_{x,c}^m, \beta_{x,c}^m)$
Insert	MAT	INS			
	INS	INS			
	INS	MAT			
Delete	MAT	DEL			
	DEL	DEL			
	DEL	MAT			
Merge	MAT	MAT			
Split	MAT	MAT			
Skip	MAT	MAT			

Here  $y \in \Omega$  where  $\Omega$  is the nucleotide alphabet and  $c \in \Omega^N$  is the context.

- 3.2 Baum-Welch algorithm
- 3.3 Viterbi algorithm

- 4 Results
- 5 Discussion

# 6 Acknowledgments

# 7 Figure Legends

## 8 Appendix

#### 8.1 Gamma distribution

$$x \sim \operatorname{Gamma}(\alpha, \beta)$$
  
 $\operatorname{E}[x] = \alpha/\beta$   
 $\operatorname{Var}[x] = \alpha/\beta^2$ 

Shape parameter  $\alpha$ , rate parameter  $\beta$ .

$$P(x|\alpha,\beta) = \frac{x^{\alpha-1}\beta^{\alpha}\exp(-x\beta)}{\Gamma(\alpha)}$$

where  $\Gamma$  is the gamma function

$$\Gamma(\alpha) = \int_0^\infty z^{\alpha - 1} \exp(-z) dz$$

Note  $\Gamma(n) = (n-1)!$  for positive integer n.

#### 8.2 Normal distribution

$$x \sim \text{Normal}(\mu, \tau)$$

Mean  $\mu$ , precision  $\tau$  (precision is reciprocal of variance).

$$P(x|\mu,\tau) = \sqrt{\frac{\tau}{2\pi}} \exp\left(-\frac{\tau}{2}(x-\mu)^2\right)$$