

Variant Calling and Annotation

Genomics Lecture #8/#9

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Variant Calling

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Variant calling is an important procedure for whole-exome and whole-genome sequencing, and for some experiments also for RNA-seq.

Two major classes of variant

- Single-nucleotide variant (SNV)
- Structural variant

In this lecture, we will discuss issues and algorithms of SNV calling

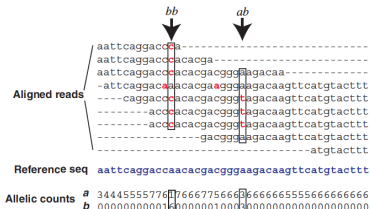
In the second half of the lecture, we will explain some of the issues and algorithms surrounding variant annotation.

Germline variants

From read mapping, we get a series of aligned columns of nucleotides and have information about

- 1 mapping quality for each read
- 2 base call quality for each position
- 3 A stack of nucleotides

- k wildtype nucleotides a
- $n - k$ nucleotides b
- $a, b \in \{a, c, g, t\}$ and $a \neq b$



Germline variants

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Ignoring for the moment everything but SNVs, our goal is to characterize each column of the alignment as homozygous wild-type (aa), heterozygous (ab), or homozygous variant (bb).

Germline variants

Thus, if the true genotype is homozygous reference (a, a), and we observe k reference bases at such a position, then the remaining $n - k$ bases must represent sequencing errors, and analogously for homozygous variant (b, b), positions

True Genotype	Number of errors
a, a	$n - k$
b, b	k

If the true genotype is heterozygous, then we can approximate the probability of the genotype as

$$\text{dbinom}(n, k, p = 0.5) = \binom{n}{k} p^k (1 - p)^{n-k} = \binom{n}{k} \frac{1}{2^n}$$

A naive algorithm

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Early NGS studies basically filtered base calls according to quality and then used a frequency filter.

Typically, a quality filter of PHRED Q20 was used (i.e., probability of error 1%). Then, the following frequency thresholds were used according to the frequency of the non-ref base, $f(b)$:

$f(b)$	genotype call
$[0, 0.2)$	homozygous reference
$[0.2, 0.8]$	heterozygous
$(0.8, 1]$	homozygous variant

A naive algorithm

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The frequency heuristic works well if the sequencing depth is high, so that the probability of a heterozygous nucleotide falling outside of the 20% – 80% region is low.

Problems with frequency heuristic:

- For low sequencing depth, leads to undercalling of heterozygous genotypes
- Use of quality threshold leads to loss of information on individual read/base qualities
- Does not provide a measure of confidence in the call

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For these reasons, a number of probabilistic methods have been developed.

We will discuss two of them and provide some algorithmic background.

- MAQ: An early algorithm.
- SNVMix: A more flexible Bayesian algorithm

The MAQ SNV calling algorithm makes use of the MAP formalism, which will be explained in the following.

MAP (Short review)

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Bayes' theorem follows from the definition of the conditional probability and relates the conditional probability $P(A|B)$ to $P(B|A)$ for two events A and B such that $P(B) \neq 0$:

$$P(A|B) = \frac{P(B|A)P(A)}{P(B)}. \quad (1)$$

Bayes Theorem

Bayes' theorem follows from the definition of the conditional probability and relates the conditional probability $P(A|B)$ to $P(B|A)$ for two events A and B such that $P(B) \neq 0$:

$$P(A|B) = \frac{P(B|A) \cdot P(A)}{P(B)}$$

The diagram shows the equation $P(A|B) = \frac{P(B|A) \cdot P(A)}{P(B)}$ with colored boxes around each term: $P(A|B)$ in a blue box, $P(B|A)$ in a red box, $P(A)$ in a green box, and $P(B)$ in a yellow box. Arrows point from the following list to these terms: 'posterior' to $P(A|B)$, 'likelihood' to $P(B|A)$, 'prior' to $P(A)$, and 'normalization constant' to $P(B)$.

- posterior
- likelihood
- prior
- normalization constant

Bayes Theorem

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Bayes' theorem is often used for a set of n mutually exclusive events E_1, E_2, \dots, E_n such that $\sum_i P(E_i) = 1$. Then, we have

$$P(E_i|B) = \frac{P(B|E_i)P(E_i)}{\sum_i P(B|E_i)P(E_i)}. \quad (2)$$

- This form of Bayes' theorem makes it clear why $B = \sum_i P(B|E_i)P(E_i)$ is called the normalization constant, because it forces the sum of all $P(E_i|B)$ to be equal to one, thus making $P(\cdot|B)$ a real probability measure

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Bayes Theorem

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In the context of bioinformatics, Bayesian inference is often used to identify the most likely model: For instance, we observe a DNA sequence and would like to know if it is a gene (M_1) or not (M_2).

Often, the model is symbolized by M and the observed data by D . Then, Bayes' theorem can be given as:

$$P(M_1|D) = \frac{P(D|M_1)P(M_1)}{P(D|M_1)P(M_1) + P(D|M_2)P(M_2)} \quad (3)$$

maximum a posteriori (MAP)

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In Bayesian statistics, maximum a posteriori (MAP) estimation is often used to generate an estimate of the maximum value of a probability distribution.

That is, if x is used to refer to the data (x can be an arbitrary expression), and θ is used to refer to the parameters of a model, then Bayes' law states that:

$$P(\theta|x) = \frac{P(x|\theta)P(\theta)}{P(x)} \quad (4)$$

maximum a posteriori (MAP)

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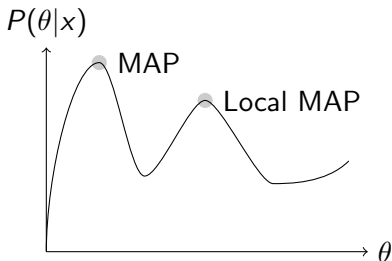
The term $P(\theta|x)$ is referred to as the posterior probability, and specifies the probability of the parameters θ given the observed data x . The denominator on the right-hand side can be regarded as a normalizing constant that does not depend on θ , and so it can be disregarded for the maximization of θ .

The MAP estimate of θ is defined as:

$$\hat{\theta} = \underset{\theta}{\operatorname{argmax}} P(\theta|x) = \underset{\theta}{\operatorname{argmax}} P(x|\theta)P(\theta) \quad (5)$$

maximum a posteriori (MAP)

One important issue about MAP estimation procedures (that we will not discuss further here), is that they tend to have the disadvantage that they “get stuck” in local maxima without being able to offer a guarantee of finding the global maximum.



MAQ

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One of the first widely used read mappers and variant callers

Li H, Ruan J, Durbin R (2008) Mapping short DNA sequencing reads and calling variants using mapping quality scores
Genome Research 18:1851–1858

- MAQ uses a number of interesting heuristics for read mapping and variant calling
- MAQ calls the genotype that maximizes the posterior probability

MAQ

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We will not review the entire MAQ mapping algorithm, but just those parts that are relevant to the variant calling process

The Mapping Quality for the assigned alignment of a read s is denoted as Q_s , the PHRED-scaled probability that the read alignment is wrong.

$$Q_s = -10 \log_{10} \Pr[\text{read is wrongly mapped}]$$

For example $Q_s = 30$ implies there is a 1:1000 probability that the read s has been wrongly mapped, $Q_s = 20$ implies a 1:100 probability, and so on.

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FASTQ and PHRED-like Quality Scores

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Recall from lecture #1:

- Illumina sequences are reported in FASTQ format.

```
@My-Illu:6:73:941:1973#0/1
```

```
GATTTGGGGTTCAAAGCAGTATCGATCAAATAGTAAATCCATTGTTCAACTCACAGTTT
```

```
+
```

```
!' '*(((***+))%%%+) (%%%) .1***-+*'')**55CCF>>>>>CCCCCCC65
```

- 1 Read identifier
- 2 sequence reported by the machine
- 3 '+' (can optionally include a sequence description)
- 4 ASCII encoded base quality scores

PHRED Quality Scores

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- The PHRED quality score is defined as

$$Q_{PHRED} = -10 \log_{10} p$$

where p is the probability that the corresponding base call is wrong.

- The PHRED quality score is nothing more than a simple transformation.

Q_{PHRED}	p	Accuracy
10	10^{-1}	90%
20	10^{-2}	99%
30	10^{-3}	99.9%
40	10^{-4}	99.99%
50	10^{-5}	99.999%

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MAQ: Mapping quality

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- MAQ uses a system of templates and hashing to index the first 28 bp of each read
- Each 28 bp sequence of the reference is similarly hashed/indexed. If a hit is found to a read MAQ calculates the sum of qualities of mismatched bases q
- Because of the templates/hash procedure, all reads with not more than two mismatches in the first 28 bp are found (pigeon hole principle)
- For simplicity, consider here quality values for an ungapped alignment of the entire read including the part of the read after the first 28 bp

MAQ: Mapping quality

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We consider the probability that a read z comes from position u of a reference sequence \mathcal{R}

$$p(z|\mathcal{R}, u) = \prod_{i \in \text{mismatched bases}} 10^{-\frac{q_i}{10}} = 10^{-\frac{\sum_i q_i}{10}}$$

That is, the probability that a read z comes from position u of reference sequence \mathcal{R} is modeled as the product of the PHRED quality scores for each of the bases that are mismatched in the alignment.

For instance, if the alignment at position u has one mismatch with PHRED base quality 20 and one with PHRED quality 10, then

$$p(z|\mathcal{R}, u) = 10^{-\frac{20+10}{10}} = 10^{-3} = 0.001$$

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MAQ: Mapping quality

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We now calculate the posterior probability of the mapping at position u , $p_s(u|\mathcal{R}, z)$ using Bayes law

$$p_s(u|\mathcal{R}, z) = \frac{p(z|\mathcal{R}, u)p(u|\mathcal{R})}{\sum_v p(z|\mathcal{R}, v)p(v|\mathcal{R})}$$

If we assume a uniform prior distribution $p(u|\mathcal{R})$, then the read is equally likely to begin at any position of the reference. The sum then goes over all positions from 1 to $L - |z| + 1$, where L is the length of \mathcal{R} and $|z|$ is the length of a read.

$$p_s(u|\mathcal{R}, z) = \frac{p(z|\mathcal{R}, u)}{\sum_{v=1}^{L-|z|+1} p(z|\mathcal{R}, v)}$$

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MAQ: Mapping quality

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Finally, we can get the PHRED scaled mapping quality, reflecting the probability that the read is wrongly mapped

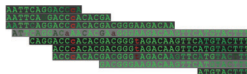
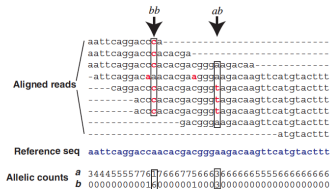
$$Q_s(u|\mathcal{R}, z) = -10 \log_{10}[1 - p_s(u|\mathcal{R}, z)]$$

Note that the explicit calculation of Q_s would require summing over the entire reference sequence, which would be impractical. MAQ therefore uses a **heuristic** to estimate Q_s , which will not be discussed further here.

MAQ: Consensus Genotype Calling

If a read is incorrectly mapped, then genotypes inferred from that read may not be reliable. Therefore, MAQ does not allow base quality q_i to exceed the mapping quality of its read z .

$$q_i = \min(q_i, Q_s(u|\mathcal{R}, z))$$



Darker background \Leftrightarrow higher mapping quality. Higher contrast \Leftrightarrow higher base quality

MAQ: Consensus Genotype Calling

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MAQ uses the quality values from mapping and of individual bases to call the most likely genotype. We assume we have a column of an alignment with k reference bases a and $n - k$ variant bases b . Any other bases are ignored as being probably sequencing errors

True Genotype	# errors	Cond. Prob. of Genotype
a, a	$n - k$	$\alpha_{n, n-k}$
b, b	k	$\alpha_{n, k}$
a, b	?	$\binom{n}{k} \frac{1}{2^n}$

MAQ: Consensus Genotype Calling

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The goal is now to decide which of the three possible genotypes has the highest posterior probability given the data (the mapping and alignment): $p(g|D)$.

MAQ now assumes the prior for the genotypes is

- $P(\langle a, a \rangle) = (1 - r)/2$
- $P(\langle b, b \rangle) = (1 - r)/2$
- $P(\langle a, b \rangle) = r$

Here, r is the probability of observing a heterozygous genotype. MAQ uses $r = 0.001$ for new SNPs, and 0.2 for known SNPs, but site-specific values for r could also be used.

MAQ: Consensus Genotype Calling

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MAQ thus calls the genotypes as

$$\hat{g} = \operatorname{argmax}_{g \in (\langle a, a \rangle, \langle a, b \rangle, \langle b, b \rangle)} p(g|D)$$

The quality of this genotype call can then be calculated as

$$Q_g = -10 \log_{10}[1 - P(\hat{g}|D)]$$

MAQ: Consensus Genotype Calling

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We now need a way of calculating $\alpha_{n,k}$, the probability of observing k errors in n nucleotides in the alignment. If we assume that error rates arise independently, and error rates are identical for all bases, then we can use a binomial distribution

$$\text{dbinom}(n, k, p = \epsilon) = \binom{n}{k} \epsilon^k (1 - \epsilon)^{n-k}$$

For instance, the probability of observing 2 erroneous nucleotides in 20, if the per read error rate is $\epsilon = 0.01$ can be calculated in R as

```
> dbinom(2,20,0.01)
[1] 0.01585576
```

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In practice, MAQ errors are correlated and are not identical for each base in the alignment. Therefore, MAQ does not use a binomial distribution, but a heuristic that reflects the probabilities of observing an alignment with the given pattern of per base error probabilities.

$$\alpha_{n,k} = c'_{n,k} \prod_{i=0}^{k-1} \epsilon_{i+1}^{\theta^i}$$

Here, ϵ_i is the i^{th} smallest base error probability for the k observed errors, $c'_{n,k}$ is a constant and θ is a parameter that controls the dependency of errors.

This equation reflects the base errors. We will not go into further detail, but if desired see the Supplemental material of the MAQ paper.

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With all of this, we can now call the posterior probabilities of the three genotypes given the data D , that is a column with n aligned nucleotides and quality scores of which k correspond to the reference a and $n - k$ to a variant nucleotide b .

$$\begin{aligned} p(G = \langle a, a \rangle | D) &\propto p(D | G = \langle a, a \rangle) p(G = \langle a, a \rangle) \\ &\propto \alpha_{n,k} \cdot (1 - r) / 2 \end{aligned}$$

$$\begin{aligned} p(G = \langle b, b \rangle | D) &\propto p(D | G = \langle b, b \rangle) p(G = \langle b, b \rangle) \\ &\propto \alpha_{n,n-k} \cdot (1 - r) / 2 \end{aligned}$$

$$\begin{aligned} p(G = \langle a, b \rangle | D) &\propto p(D | G = \langle a, b \rangle) p(G = \langle a, b \rangle) \\ &\propto \binom{n}{k} \frac{1}{2^n} \cdot r \end{aligned}$$

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Finally, the genotype with the highest posterior probability is chosen

$$\hat{g} = \operatorname{argmax}_{g \in (\langle a, a \rangle, \langle a, b \rangle, \langle b, b \rangle)} p(g|D)$$

The probability of this genotype is used as a measure of confidence in the call.

What have we learned?

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The MAQ algorithm is typical for many in genomics in that a well known statistical or algorithmic framework is used with a number of heuristics that deliver reasonable values for the parameters needed for the framework to work.

Major aspects of MAQ SNV calling algorithm

- Integrates mapping and per base quality scores
- Bayesian (MAP) framework to integrate observations and priors on genotypes
- Provides estimation of reliability of genotype call.

Expectation Maximization (EM)

We will discuss how EM is used for mixture distributions. For ease of presentation, we will discuss in detail a mixture of Gaussians, but the principles are the same for other probability distributions

The basic framework is that we assume that a data point y_j is produced as follows

- First, choose one of $i \in \{1, \dots, I\}$ components that produces the measurement
- Then, according to the parameters of component $C = i$, the actual measurement is generated

This is known as a mixture distribution, and the corresponding probability density function (pdf) is defined as

$$p(y_j|\theta) = \sum_{i=1}^I \alpha_i p(y_j|C = i, \beta_i) \quad (6)$$

Expectation Maximization (EM)

Note that in this notation, θ comprises both the weight parameters for the probability of one of the l components generating the data, as well as the various parameters for each of the components, β_i (where in general β_i can be a vector of parameters)

- Of course, $\sum_{i=1}^l \alpha_i = 1$
- The parameters β_i are associated with the PDF of component i .

We will now show how to perform maximum likelihood estimation using the Expectation Maximization (EM) framework to find values for the parameters θ that maximize the probability of the data. This involves maximization of the log-likelihood for θ .

$$\log L(\theta) = \log p(\mathbf{y}|\theta) \quad (7)$$

Expectation Maximization (EM)

We can flesh out the formula as follows

$$\begin{aligned}\log L(\theta) &= \log p(\mathbf{y}|\theta) \\ &= \log \left\{ \prod_{j=1}^J p(y_j|\theta) \right\} \\ &= \sum_{j=1}^J \log p(y_j|\theta) \\ &= \sum_{j=1}^J \log \left\{ \sum_{i=1}^I \alpha_i p(y_j|C=i, \beta_i) \right\}\end{aligned}$$

- Since the log is outside the sum in the last expression, there is no analytic (closed form) optimization.

Expectation Maximization (EM)

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The EM algorithm is essentially like a push-me-pull-you algorithm that goes back and forth between

- find an estimate for the likelihood function
- maximizing the whole term



Expectation Maximization (EM)

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If there is time in the practical session, I will explain the derivation of the EM method and show in detail how the maximization expressions are derived for a simple distribution – mixture of Gaussians. For today, I will show only a high level summary. For the practical, you will be expected to implement a simplified version of EM – known as gene counting (will explain at end of this lecture)

Expectation Maximization (EM)

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The EM algorithm will be explained using a mixture of Gaussians (SNVMix uses some slightly less familiar distributions). Recall the form of a multivariate Gaussian distribution for a k -dimensional vector $\mathbf{x} = [x_1, x_2, \dots, x_k]$:

$$\mathbf{x} \sim \mathcal{N}(\boldsymbol{\mu}, \boldsymbol{\Sigma}) \quad (8)$$

i.e. the probability density function is

$$f(x_1, x_2, \dots, x_k) = \frac{1}{(2\pi)^{k/2} |\boldsymbol{\Sigma}|^{1/2}} e^{-\frac{1}{2}(\mathbf{x}-\boldsymbol{\mu})^T \boldsymbol{\Sigma}^{-1}(\mathbf{x}-\boldsymbol{\mu})} \quad (9)$$

Expectation Maximization (EM)

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We thus wish to estimate the parameters for a mixture of Gaussians. We need to estimate both the mixture parameters $\pi_1, \pi_2, \dots, \pi_c$ with $\sum_{i=1}^c \pi_i = 1$, but also the means and variances for each of the individual Gaussian distributions, $\mu_1, \mu_2, \dots, \mu_c$, and $\Sigma_1, \Sigma_2, \dots, \Sigma_c$.

We thus want to maximize the log likelihood given by

$$L(\theta | \mathbf{x}_1, \mathbf{x}_2, \dots, \mathbf{x}_n) = \log \prod_{i=1}^n \sum_{k=1}^c \pi_k f(\mathbf{x}_i | \mu_k, \Sigma_k) \quad (10)$$

or equivalently

$$L(\theta | \mathbf{x}_1, \mathbf{x}_2, \dots, \mathbf{x}_n) = \sum_{i=1}^n \log \sum_{k=1}^c \pi_k f(\mathbf{x}_i | \mu_k, \Sigma_k) \quad (11)$$

Expectation Maximization (EM)

We can now calculate the probability that a particular data point \mathbf{x}_j belongs to a particular component k

We write the posterior probability that an observation \mathbf{x}_j belongs to component k as

$$\hat{\tau}_{jk} = \frac{f(\mathbf{x}_j | \mu_k, \Sigma_k) \hat{\pi}_k}{\sum_{i=1}^c f(\mathbf{x}_j | \mu_i, \Sigma_i) \hat{\pi}_i} \quad (12)$$

- The posterior probability $\hat{\tau}_{jk}$ is unknown but can be easily **estimated** if we use the current values of the parameters for the Gaussians.
- $\hat{\tau}_{jk}$ is thus an estimate for the probability that observation j was generated by component k given the data and current parameter estimates
- This is the **Expectation** step of the EM algorithm

Expectation Maximization (EM)

Given our current estimates of the component membership for each of the datapoints, we can maximize the values of the mixture parameters as well as of the Gaussians by setting their first derivative to zero etc (individual steps not shown here). This leads to the following

$$\hat{\pi}_k = \frac{1}{n} \sum_{i=1}^n \hat{\tau}_{ik} \quad (13)$$

$$\hat{\mu}_k = \frac{1}{n} \sum_{i=1}^n \frac{\hat{\tau}_{ik} \mathbf{x}_i}{\hat{\pi}_k} \quad (14)$$

$$\hat{\Sigma}_k = \frac{1}{n} \sum_{i=1}^n \frac{\hat{\tau}_{ik} (\mathbf{x}_i - \hat{\mu}_k) (\mathbf{x}_i - \hat{\mu}_k)^T}{\hat{\pi}_k} \quad (15)$$

This is the **Maximization** step of the EM algorithm

Expectation Maximization (EM)

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Thus, the individual steps of the EM algorithm are thus

- 1 Initial component parameters (with a reasonable guess)
- 2 For each data point, calculate posterior probability of membership to each component using the current parameter values
- 3 Then, based on these estimates, maximize the log likelihood of the parameters given the data
- 4 Repeat until convergence¹

¹and hope you have not landed in a local maximum.

SNVMix

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We will now discuss an algorithm called SNVMix, that uses the EM framework to estimate optimal parameters for calling SNPs in a Bayesian framework.

- This algorithm was first described here:
Shah SP et al. (2009) Mutational evolution in a lobular breast tumor profiled at single nucleotide resolution.
Nature **461**:809-13.
- an improved version (which we will not discuss) was later presented here:
Goya R et al. (2010) SNVMix: predicting single nucleotide variants from next-generation sequencing of tumors.
Bioinformatics **26**:730-6.

SNVMix: Model specification

Variant
Calling and
Annotation

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The core of the model is a specification of the genotypes and the conditional probabilities of the observed distribution of alleles – similar to MAQ.

- Let $G_i = k, k \in \{\langle a, a \rangle, \langle a, b \rangle, \langle b, b \rangle\}$ be a multinomial random variable representing the genotype at nucleotide position i ($a = \text{ref}, b$ is non-ref).
- Let the observed allele frequency $X_i = [a_i, b_i]^T$, i.e., a vector of counts of the reference and non-reference alleles at position i
- Then $N_i = a_i + b_i$ is the observed read depth at position i .

SNVMix: Model specification

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The central idea is that we assume the allele counts are generated by a class conditional density

Thus

$$X_i \sim \text{Binom}(a_i | \mu_k, N_i) \quad (16)$$

The probability of the observed read counts (number of a_i bases amongst all N_i bases at position i) is thus conditioned on the underlying genotype $G_i = k$, and μ_k is the corresponding parameter of a Binomial distribution for genotype k .

SNVMix: Model specification

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If we actually knew the genotype, then it is simplicissimo to calculate the probability of the allele counts using the binomial distribution.

Thus

$$P(X_i) = \text{Binom}(a_i | \mu_k, N_i) = \binom{N_i}{a_i} \mu_k^{a_i} (1 - \mu_k)^{N_i - a_i} \quad (17)$$

Intuitively, we would expect the values of μ_{aa} to be close to 1, those for μ_{ab} to be close to 0.5 and those for μ_{bb} to be near zero. However, we do not know the exact values for real data, which may depend on things such as the sequencing error rate².

² And for cancer data, on the relative mixture of normal and cancerous tissue in a biopsy. 

SNVMix: Model specification

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Annotation

The prior probability of observing a genotype k at any position of the sequenced genome is represented as a multinomial variable π .

- $0 \leq \pi_k \leq 1, \quad \forall k$
- $\sum_{k=1}^3 \pi_k = 1$
- Note that in general we will expect the values of π to be highly skewed towards observing homozygous reference bases (since most genomic positions are not variant in any one individual)
- SNVMix is thus a classic generative mixture model to explain the observed data.

SNVMix: Model specification

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Annotation

The marginal distribution of X_i (in which we have marginalized – removed – the influence of the actual genotype) can then be calculated as the convex combination of the class conditional Binomial densities, weighted by the multinomial π :

$$p(X_i) = \sum_{k=1}^3 \pi_k \binom{N_i}{a_i} \mu_k^{a_i} (1 - \mu_k)^{N_i - a_i} \quad (18)$$

Again, the sum is taken over k representing the three genotypes aa , ab , and bb .

SNVMix: Model specification

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We can then use this equation to calculate the log likelihood of our entire dataset, which comprises positions $1 \dots T$.

$$\log p(X_{1:T} | \mu_{1:K}, \pi) = \sum_{i=1}^T \log \sum_{k=1}^3 \pi_k \binom{N_i}{a_i} \mu_k^{a_i} (1 - \mu_k)^{N_i - a_i} \quad (19)$$

Our problem is that the model parameters $\theta = (\pi, \mu)$ are not known. If the true genotype were somehow known, we could simply calculate them from the training data. But, instead, we will learn (estimate) the parameters from data by using maximum a posteriori (MAP) expectation maximization (EM).

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Assuming we have solved for the parameters (we will get to that shortly), then we can easily calculate the posterior probability of any genotype using Bayes rule

$$p(G_i = k | X_{1:N}, \pi, \mu_k) = \frac{\pi_k \text{Binom}(X_i | \mu_k, N_i)}{\sum_{j=1}^3 \pi_j \text{Binom}(X_i | \mu_j, N_i)} \quad (20)$$

For notational simplicity, we will denote $p(G_i = k | X_{1:N}, \pi, \mu_k)$ as $\gamma_i(k)$, the marginal probability of the genotype for position i given all the data and the model parameters.

SNVMix: Prior Distributions

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Bayesian mixture models use hyperparameters, i.e., parameters of a prior distribution; the term is used to distinguish them from parameters of the model used for the final analysis. We will use two underlying distributions to calculate these hyperparameters for SNVMix.

- $\pi \sim \text{Dirichlet}(\pi|\delta)$
- $\mu \sim \text{Beta}(\mu_k|\alpha_k, \beta_k)$

Beta distribution

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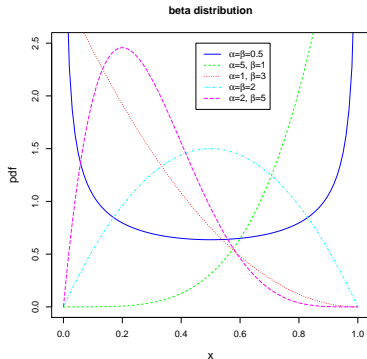
Beta distribution: A family of continuous distributions defined on $[0, 1]$ and parametrized by two positive shape parameters, α and β

$$p(x) = \frac{1}{B(\alpha, \beta)} \cdot x^{\alpha-1} (1-x)^{\beta-1}$$

here, $x \in [0, 1]$, and

$$B(\alpha, \beta) = \frac{\Gamma(\alpha + \beta)}{\Gamma(\alpha) \cdot \Gamma(\beta)}$$

where Γ is the Gamma function
(extension of factorial).



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Beta distribution

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Important in the current context is that the Beta distribution is the conjugate prior to the binomial distribution.

- That is, we can express our prior belief about the value of the μ_k parameter of the binomial distribution for read counts using a Beta distribution
- We say that μ_k is conjugately distributed according to a Beta distribution: $\mu_k \sim \text{Beta}(\mu_k | \alpha_k, \beta_k)$.
- This requires us to express our prior belief about μ_{aa} , μ_{ab} , and μ_{bb} by specifying values for α_k, β_k

Beta distribution

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For instance, let us say we are not very sure about what we think the value of $\mu_{a,b}$ should be, perhaps because we do not know if the sample being sequenced contains tumorous or non-tumorous tissues³.

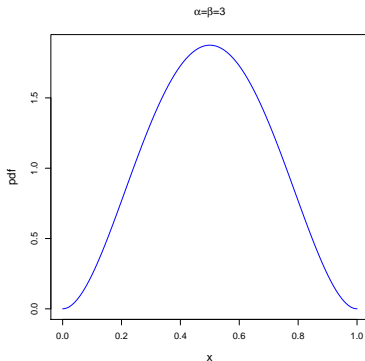
- We might then try $\alpha_{ab} = \beta_{ab} = 3$

```
x <- seq(0.0, 1.0, 0.01)
y <- dbeta(x, 3, 3)
title <- expression(paste(alpha,"=",beta,"=3"))
plot(x, y, type="l",main=title,
xlab="x",ylab="pdf",col="blue",lty=1,cex.lab=1.25)
```

³Tumor tissue may be characterized by the loss of heterozygosity (LOH) of large chromosomal regions.

Beta distribution

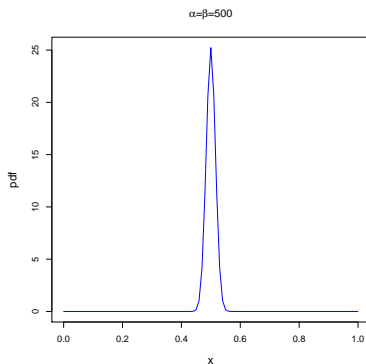
$$\alpha_{ab} = \beta_{ab} = 3$$



- x is here representing a value for μ_{ab} , and the y axis reflects our belief about the prior probability of this value
- Question: Are we very sure about μ_{ab} ?

Beta distribution

$$\alpha_{ab} = \beta_{ab} = 500.$$



- Question: How sure are we now about μ_{ab} ?

Beta distribution in SNVMix

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The expected value of a $\text{Beta}(\alpha + \beta)$ distribution is simply

$$\frac{\alpha}{\alpha + \beta}$$

In SNVMix, these values are defined as

- $\alpha_{aa} = 1000$, $\beta_{aa} = 1$, that is, our prior belief in reference reads given homozygous reference sequence is 0.999001
- $\alpha_{ab} = 500$, $\beta_{ab} = 500$, that is, our prior belief in reference reads given a het true sequence is 0.5
- $\alpha_{bb} = 1$, $\beta_{bb} = 1000$ (vice versa to $\alpha_{aa} = 1000$, $\beta_{aa} = 1$)

SNVMix: M-step updating equation for μ

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The maximization step updating equation basically adds to observed counts for a certain true genotype to our prior.

$$\mu_k^{new} = \frac{\sum_{i=1}^T a_i^{I(G_i=k)} + \alpha_k}{\sum_{i=1}^T N_i^{I(G_i=k)} + \alpha_k + \beta_k - 2} \quad (21)$$

- Note that in this notation, $I(G_i = k)$ is an indicator function so that the expression is zero unless $G_i = k$
- The update is simply the proportion of the observed reference reads with “pseudocounts” added from the Beta prior (amongst all positions called to genotype k).

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nomenclature

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Dirichlet

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Annotation

The Dirichlet distribution is the multivariate generalization of the beta distribution and represent the conjugate prior of the multinomial distribution. Thus, just as SNVMix used the Beta distribution as a prior for μ (binomial distribution of read counts), it uses the Dirichlet as a prior for π (multinomial distribution for the three possible genotypes).

In SNVMix, the values for the prior are set to

$\delta(\langle a, a \rangle)$	$\delta(\langle b, b \rangle)$	$\delta(\langle b, b \rangle)$
1000	100	100

Dirichlet

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Thus, the prior is skewed toward π_{aa} assuming that most positions will be homozygous for the reference allele. The pseudocounts are essentially equivalent to having seen $1000 + 100 + 100 = 1200$ positions with the distribution 83.3% $\langle a, a \rangle$, and 8.3% each for $\langle b, b \rangle$ and $\langle b, b \rangle$.

The weight of the prior belief is reflected in the number of pseudocounts. For instance, the following counts result in the same proportion but there is much less weight of prior belief

$\delta(\langle a, a \rangle)$	$\delta(\langle b, b \rangle)$	$\delta(\langle b, b \rangle)$
10	1	1

Dirichlet

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A Dirichlet distribution of order k (3 in our example) is a PDF that represents the belief (“probability”) that the probabilities of k distinct events (in our case, the genotypes $\langle a, a \rangle, \langle a, b \rangle, \langle b, b \rangle$) are x_i given that each event has been observed $\alpha_i - 1$ times.

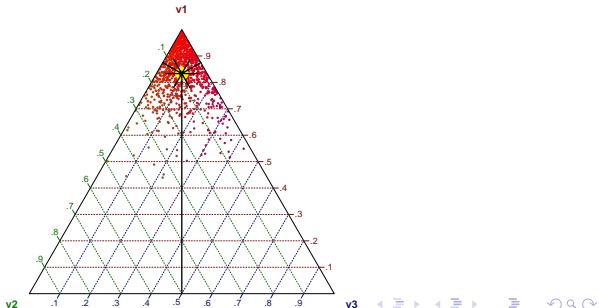
$$f(x_1, x_2, \dots, x_{k-1}; \alpha_1, \alpha_2, \dots, \alpha_k) = \frac{1}{B(\alpha)} \prod_{i=1}^k x_i^{\alpha_i - 1} \quad (22)$$

- Note that by convention f has $k - 1$ arguments. Since $\sum_{i=1}^k x_i = 1$ there is no need to show the k^{th} argument.
- $B(\alpha)$ is the Beta function

$$B(\alpha) = \frac{\prod_{i=1}^k \Gamma(\alpha_i)}{\Gamma\left(\sum_{i=1}^k \alpha_i\right)} \quad (23)$$

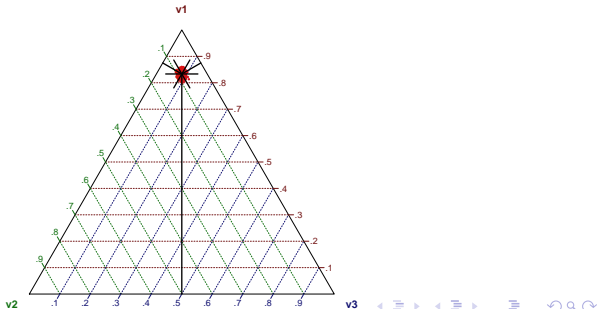
Dirichlet distribution with a low number of pseudocounts (relatively weak prior):

- $\delta(\langle a, a \rangle) = 10$
- $\delta(\langle a, b \rangle) = 1$
- $\delta(\langle b, b \rangle) = 1$



Dirichlet distribution with a higher number of pseudocounts
(relatively strong prior):

- $\delta(\langle a, a \rangle) = 1000$
- $\delta(\langle a, b \rangle) = 100$
- $\delta(\langle b, b \rangle) = 100$



SNVMix: M-step updating equation for π

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Annotation

The update equation for π is similar to that for μ

$$\pi_k^{new} = \frac{\sum_{i=1}^T I(G_i = k) + \delta(k)}{\sum_{j \in \{\langle a,a \rangle, \langle a,b \rangle, \langle b,b \rangle\}} \sum_{i=1}^T I(G_i = j) + \delta(j)} \quad (24)$$

SNVMix: Initialize EM

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We are now in a position to initialize the EM

We need:

- Mapped NGS reads comprising $i = 1, \dots, T$ genomic positions, each of which has N_i reads with a_i reference and b_i nonreference bases.
- Initialize $\pi_k = \frac{\delta(k)}{N_\delta}$ where $N_\delta = \frac{\delta(k)}{\sum_j \delta(j)}$
- Initialize $\mu_k = \frac{\alpha_k}{\alpha_k + \beta_k}$
- pick a tolerance to judge convergence

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SNVMix: Run EM

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The EM algorithm iterates between the E-step where we assign the genotypes using Equation (20) and the M-step where we re-estimate the model parameters with equations (24) for π , (21) for μ .

At each iteration we evaluate the complete data log-likelihood as given by Equation (19) and the algorithm terminates when this quantity no longer increases

SNVMix1 vs SNVMix2

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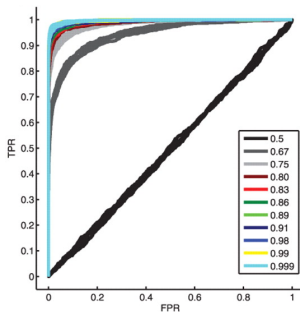
Variant
nomenclature

Annotation

The SNVMix algorithm was later extended to include mapping and base qualities into the same Bayesian framework, primarily by adapting the formulas used for the EM equations. We will not discuss this here^a

^aGoya R et al. (2010) *Bioinformatics* 26:730–736.

Performance of SNVMix2 algorithm on simulated data with increasing levels of certainty in the base call



Annotation and Variant Nomenclature

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The output of variant calling programs is a set of calls with chromosomal coordinates, e.g.

```
10 72535007 C T
```

- i.e., position 72614421 on chromosome 10, which in the reference sequence has a thymine (T) is found to have an alternate cytosine (C) base
- This is fine, but biologists or MDs generally need gene-based annotations such as

```
HGVS=C10orf27:NM_152710:exon8:c.710G>A:p.R237Q
```

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Annotation

Annotation algorithms take genomic sequences and transcripts, and metadata about genes (e.g., positions of exons on a chromosome), and infer the transcript-level annotation for chromosomal variants.

- You are expected to know the basic variant nomenclature from the HGVS
- If you are not familiar with this, study how to represent missense, nonsense, and deletion mutations in cDNA and protein at <http://www.hgvs.org/mutnomen/>
- DNA: A,C,G,T
 - c.435C>A
- Protein: 1- or 3-letter code
 - p.A212P, Ala212Pro
- Use HGNC⁴ Gene symbols, e.g., *FBN1* for Fibrillin-1
- Nice Tool: <http://www.humgen.nl/mutalyzer/1.0.1/>

⁴HUGO Gene Nomenclature Committee

Standardized Nomenclature for Mutations

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Annotation

Short review of HGVS nomenclature for mutations

DNA ...

- Single-nucleotide Substitution c.123A>G
- Deletion c.123de1A
- Duplication c.123dupA
- Insertion c.123_124insC

Deletions & Insertions

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- c.546delT
- c.546del
- c.586_591del
- c.586_591delTGGTCA oder c.586_591del6
- c.546_547insT (not c.546insT – which is ambiguous)
- c.1086_1087insGCGTGA

Alleles

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- Nomenclature for two alleles in one gene may be needed for autosomal recessive diseases:
- c . [546C>T] + [2398de1T]

Frameshift

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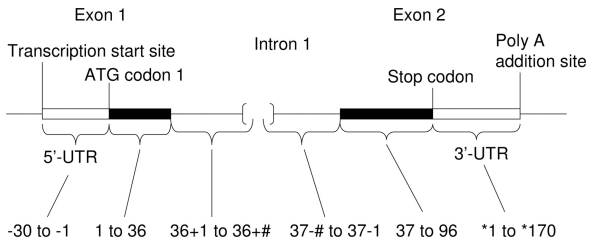
Annotation

- Short form p.Arg83fs
- Long form: p.Arg83SerfsX15
 - The frameshift has led to an amino acid substitution (Arg83Ser), which as a rule is located at the first amino acid affected by the frameshift mutation (“fs”)
 - We then see the length of the remaining protein sequence up to the next stop codon (X15, i.e., 15 amino acids)
 - Note that the amino acid sequence following the frameshift is almost always different from the wildtype sequence
 - We expect to find a premature termination codon not very far downstream from frameshift mutations (why?)

Numbering

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indicates any positive integer number

- Note: splice mutations are numbered according to the closest exonic nucleotide position: z.B. $36+1G>C$, $37-2A>G$

Annotation

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Annotation

We will now show how to take the output of variant calling programs (calls with chromosomal coordinates) and to produce gene/transcript-based annotations

There are a number of programs out there that are commonly used to process VCF files. Two of the best are

- Annovar: Wang K et al., (2010) *Nucleic Acids Res* **38**:e164.
- Variant Effect Predictor (At the EBI)

The latter can be used online (google it)

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The algorithms involve a lot of heuristics, which have to deal with the “weirdness” of our genome: overlapping genes, genes located in introns of other genes, multiple coding/non-coding transcripts of a single gene, antisense transcripts etc etc.

We will not attempt to demonstrate the complete process, but will present a simplified version of how a deletion variant gets annotated.

Deletion

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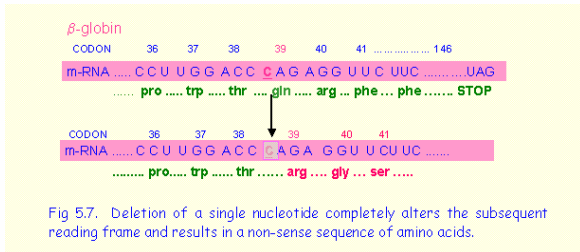


Fig 5.7. Deletion of a single nucleotide completely alters the subsequent reading frame and results in a non-sense sequence of amino acids.

- Example of a deletion of a single nucleotide causing a frameshift

Deletion

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Annotation

On the chromosome, a deletion mutation is represented as

1 1887113 1887113 C -

That is,

- Deletion of a “C” at position 1887113 of chromosome 1

Our goal is now to find out whether this deletion occurs within an exon of a gene, and if so, the determine the effect on the cDNA sequence

- KIAA1751:uc001aim.1:exon18:c.2193delG:p.Q731fs

Annotation: Where does the data come from

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Annotation

The read mapping algorithms use a reference genome sequence, usually hg18 (March 2006) or hg19 (Feb 2009). One can download annotation files from various sources including Ensembl and UCSC. These files include information about genes and transcripts, and are used to infer the effects of variants.

We will demonstrate this with the UCSC known genes data.⁵ One can download a series of files with the entire genome sequence and information about each transcript, its exon structure, and the nucleotide sequence of the splice mRNA (cDNA).

⁵see <http://genome.ucsc.edu>

UCSC KnownGenes

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Annotation

field	example	explanation
name	uc001aaa.3	Name of gene
chrom	chr1	ref. chromosome
strand	+	+ or - strand
txStart	11873	Transcription start
txEnd	14409	Transcription end
cdsStart	11873	Coding region start
cdsEnd	11873	Coding region end
exonCount	3	Number of exons
exonStarts	11873,12612,13220,...	Exon start positions
exonEnds	12227,12721,14409,...	Exon start end positions

Annotation: bird's eye view

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Annotation

- 1 Identify the gene(s) that overlaps with the variant⁶
- 2 Determine whether the variant is located in an exon⁷
- 3 Determine whether the variant is located in the coding sequence⁸
- 4 If the variant is a SNV, determine whether the encoded amino acid is changed⁹, if so annotate as **missense**
- 5 If the variant is located right before or after an exon/intron boundary, annotate as **splicing**
- 6 If the variant removes/adds nucleotides from the CDS, annotate as **deletion/insertion**¹⁰

⁶ If you do not find such a gene, the variant is **upstream**, **downstream**, or **intergenic**.

⁷ If not, the variant is **intronic**

⁸ If not, the variant is **3' or 5' UTR**

⁹ If not, the variant is a **synonymous substitution**

¹⁰ If the variant removes/adds $2n+1$ or $2n+2$ nucleotides, also annotate as **frameshift**

Annotation: Overview

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Annotation

Algorithm 1 Create annotation

```
1: get_annotation(chr, pos, ref, var)
2: gene  $\leftarrow$  get_overlapping_gene(chr, pos)
3: if gene == NULL then
4:   return ‘‘variant non genic’’
5: end if
6: if ann  $\leftarrow$  get_deletion(chr, pos, ref, var) != NULL then
7:   return ann
8: else
9:   // Other functions for other mutation classes
10: end if
```

Annotation: Get overlapping gene

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Algorithm 2 Get overlapping gene

```
1: gene  $\leftarrow$  get_overlapping_gene(chr, pos)
2: for all  $g \in \mathcal{G}$  do //loop over candidate genes
3:   txStart  $\leftarrow$  g.txStart
4:   txEnd  $\leftarrow$  g.txEnd
5:   if pos < txStart then
6:     continue //variant is 5' to gene
7:   else if pos > txEnd then
8:     continue //variant is 3' to gene
9:   end if
10:  return g // variant is located within gene model
11: end for
```

• Note that for genes on "-", the meaning of txStart and txEnd is "reversed"

Annotation: Get rvarstart

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Algorithm 3 Get rvarstart

```
1: get_rvarstart(gene g, pos)
2: cumlenintron  $\leftarrow$  0 // cumulative length of introns
3: cumlenexon  $\leftarrow$  0 // cumulative length of exons
4: for  $k \in g.getExons()$  do //loop over exons
5:     cumlenexon  $\leftarrow$  cumlenexon + g.getExonLen(k)
6:     if  $k > 0$  then //no intron before first exon!
7:         cumlenintron  $\leftarrow$  cumlenintron + g.getIntronLen(k)
8:     end if
9:     if pos located in exon  $k$  then
10:         rvarstart  $\leftarrow$  start - txStart - cumlenintron + 1;
11:         return rvarstart // start of variant within mRNA
12:     end if
13: end for
```

Annotation: Get deletion

Algorithm 4 Get rvarstart

```
1: get_deletion(gene g, pos,ref)
2: rvarstart  $\leftarrow$  get_rvarstart(g, pos)
3: cdspos  $\leftarrow$  rvarstart - g.getCDSstart() + 1
4: mut  $\leftarrow$  sprintf("c%ddel%c",cdspos,ref)
5: return mut
```

- This will return something like c.2193delG, a deletion of a "G" at position 2193 of the coding sequence
- N.B.: This function is highly simplified and will only work for the + strand.

Annotation: Final words

Variant
Calling and
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This should be enough to give you some idea of how genomic variants are annotated. We have of course left out a large number of functionalities

- Figure out effect on protein based on translating the DNA sequence
- Functions for other kinds of variants
- Figuring out how to number variants in genes on the minus strand
- Reporting either all consequences of a given variant (there may be a different consequence for each of the isoforms of a gene)
- Alternatively, prioritize the variants to report only the most relevant variant

Variant
Calling

naive

MAP

MAQ

EM

SNVMix

Variant
nomenclature

Annotation

Summary

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In these two lectures we have examined how to call variants. We have studied two classes of algorithms, MAP estimation and expectation maximization, and how they are used in two variant calling algorithms, MAQ and SNVMix. Finally, we have examined how called variants are annotated as to their consequences for genes and transcripts, which is usually necessary for biological or medical interpretation of variants.

What you should now know:

- The kinds of data used in variant calling (mapping quality, base quality, depth, ...)
- How this data is exploited to improve variant calling
- Bayes' law and how it can be used to estimate parameters
- Be able to interpret the major formulae of MAP and EM (no need to memorize)
- Variant nomenclature and basic ideas of variant annotation (“bird’s eye view”)

The End of the Lecture as We Know It

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Variant
Calling

naive

MAP

MAQ

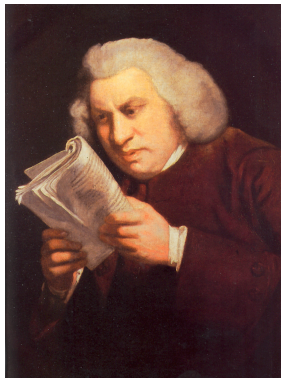
EM

SNVMix

Variant
nomenclature

Annotation

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Lectures were once useful; but now, when all can read, and books are so numerous, lectures are unnecessary. If your attention fails, and you miss a part of a lecture, it is lost; you cannot go back as you do upon a book... People have nowadays got a strange opinion that everything should be taught by lectures. Now, I cannot see that lectures can do as much good as reading the books from which the lectures are taken. I know nothing that can be best taught by lectures, except where experiments are to be shown. You may teach chymistry by lectures. You might teach making shoes by lectures!