

Une histoire de mots inattendus et de génomes Sophie Schbath

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Une histoire de mots inattendus et de génomes

Sophie Schbath





ALEA 2017, Marseille, 22 mars 2017

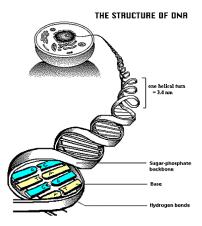
Sophie Schbath (INRA - MaIAGE)

Histoire de mots

Introduction

DNA and motifs

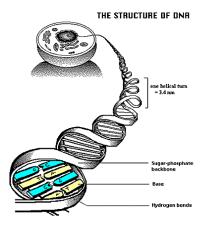
- DNA : Long molecule, sequence of nucleotides
- Nucleotides : A(denine), C(ytosine), G(uanine), T(hymine).



...GTTCAATCGTAGGTAGGTACTGAATGGTAGGTATGTTGA...

DNA and motifs

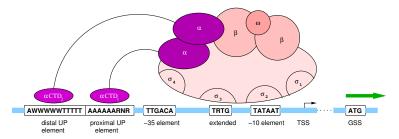
- DNA : Long molecule, sequence of nucleotides
- Nucleotides : A(denine), C(ytosine), G(uanine), T(hymine).
- Motif (= oligonucleotides) : short sequence of nucleotides, e.g. AGGTA



...GTTCAATCGT<u>AGGTAGGTA</u>CTGAATGGT<u>AGGTA</u>TGTTGA...

DNA and binding sites

 Functional motif : recognized by proteins or enzymes to initiate a biological process



Some functional motifs

- Restriction sites : recognized by specific bacterial restriction enzymes ⇒ double-strand DNA break.
 E.g. GAATTC recognized by *Eco*RI
- Chi motif : recognized by an enzyme which processes along DNA sequence and degrades it ⇒ enzyme degradation activity stopped and DNA repair is stimulated by recombination.
 E.g. GCTGGTGG recognized by *RecBCD* (*E. coli*)
- *parS*: recognized by the *Spo0J* protein ⇒ organization of *B. subtilis* genome into macro-domains.
 TGTTAACACGTGAAACA
- **promoter** : structured motif recognized by the RNA polymerase to initiate gene transcription.

E.g. TTGAC - - TATAAT (*E. coli*).

Some functional motifs

● **Restriction sites** : recognized by specific bacterial restriction enzymes ⇒ double-strand DNA break.

E.g. GAATTC recognized by EcoRI

very rare along bacterial genomes

- Chi motif : recognized by an enzyme which processes along DNA sequence and degrades it ⇒ enzyme degradation activity stopped and DNA repair is stimulated by recombination. E.g. GCTGGTGG recognized by *RecBCD* (*E. coli*) very frequent along *E. coli* genome
- parS : recognized by the Spo0J protein ⇒ organization of B. subtilis genome into macro-domains.

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very frequent into the ORI domain, rare elsewhere

 promoter : structured motif recognized by the RNA polymerase to initiate gene transcription.

E.g. TTGAC — — — TATAAT (*E. coli*). particularly located in front of genes

Prediction of functional motifs

Most of the functional motifs are unknown in the different species.

For instance,

- which would be the Chi motif of S. aureus? [Halpern et al. (07)]
- Is there an equivalent of parS in E. coli? [Mercier et al. (08)]

Statistical approach : to identify candidate motifs based on their statistical properties.

The most over-represented 8-letter words under M1

The most over-represented families anbcdefg under M1

E. coli $(\ell = 4.6 \ 10^6)$

H. influenzae ($\ell = 1.8 \ 10^6$)

word	obs	exp	score	motif	obs	exp	score
gctggtgg	762	84.9	73.5	gntggtgg	223	55.3	22.33
ggcgctgg	828	125.9	62.6	anttcatc	469	180.3	21.59
cgctggcg	870	150.8	58.6	anatcgcc	288	87.8	21.38
gctggcgg	723	125.9	53.3	tnatcgcc	279	84.5	21.18
cgctggtg	619	101.7	51.3	gnagaaga	270	83.6	20.10

Statistical questions on word occurrences

Here are some quantities of interest.

- Number of occurrences (overlapping or not) :
 - Is N^{obs}(**w**) significantly high?
 - Is $N^{obs}(\mathbf{w})$ significantly higher than $N^{obs}(\mathbf{w}')$?
 - Is $N_1^{\text{obs}}(\mathbf{w})$ significantly more unexpected than $N_2^{\text{obs}}(\mathbf{w})$?
- Distance between motif occurrences :
 - Are there significantly rich regions with motif w
 - Are two motifs significantly correlated?
- Waiting time till the first occurrence :
 - Is the presence of a motif w significant?

A model to define what to expect

Assessing the significance of an observed value (count, distance, occurrence, etc.) requires to define a **null model** to set what to expect.

A model for random sequences :

- Markov chain models : a Markov chain of order m (Mm) fits the h-mers frequencies for h = 1, ..., (m + 1).
- Hidden Markov models allow to integrate heterogeneity.

A model to define what to expect

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A model for the occurrence processes :

- (compound) Poisson processes allow to fit the number of occurrences and then to study the significance of inter-arrival times ([Robin (02)], or to compare the exceptionality of a word in two sequences ([Robin et al. (07)]).
- Hawkes processes allow to estimate the dependence between occurrence processes ([Gusto and S. (05)], [Reynaud and S. (10)])

Markov chains of order m : model Mm

Let $X_1 X_2 X_3 \cdots X_{\ell} \cdots$ be a stationary Markov chain of order *m* on $\mathcal{A} = \{a, c, g, t\}$, i.e.

$$\mathbb{P}(X_i = b \mid X_1, X_2, ..., X_{i-1}) = \mathbb{P}(X_i = b \mid X_{i-m}, ..., X_{i-1}).$$

Transition probabilities are denoted by

$$\pi(a_1\cdots a_m,b)=\mathbb{P}(X_i=b\mid X_{i-m}\cdots X_{i-1}=a_1\cdots a_m),$$

whereas the stationary distribution is given by

$$\mu(a_1a_2\cdots a_m):=\mathbb{P}(X_i=a_1,\ldots,X_{i+m-1}=a_m),\quad\forall i.$$

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The MLE are

$$\widehat{\pi}(a_1\cdots a_m, a_{m+1}) = \frac{N^{\text{obs}}(a_1\cdots a_m a_{m+1})}{N^{\text{obs}}(a_1a_2\cdots a_m+1)}, \quad \widehat{\mu}(a_1\cdots a_m) = \frac{N^{\text{obs}}(a_1\cdots a_m)}{\ell - m + 1}$$

$$\rightarrow \widehat{\mathbb{E}}N(a_1\cdots a_m a_{m+1}) \simeq N^{\mathrm{obs}}(a_1\cdots a_m a_{m+1})$$

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Occurrences of words may overlap in DNA sequences (no space between words).

 \Rightarrow occurrences are not independent.

 Occurrences of overlapping words will tend to occur in clumps. For instance, they are 3 overlapping occurrences of CAGCAG below :

TAGACAGATAGACGAT CAGCAGCAGCAG ACAGTAGGCATGA...

• On the contrary, occurrences of non-overlapping words will never overlap.

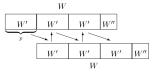
Overlapping occurrences (2)

All results on word occurrences will depend on the overlapping structure of the words.

Classically, this structure is described thanks to the periods of a word :

p is a period of $\mathbf{w} := w_1 w_2 \cdots w_h$ iff $w_i = w_{i+p}$, $\forall i$

meaning that 2 occurrences of **w** can overlap on h - p letters.



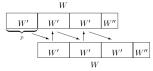
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We also define the overlapping indicator :

 $\varepsilon_{h-p}(\mathbf{w}) = 1$ if p is a period of **w**, and 0 otherwise

Detecting words with significanly unexpected counts

Problem

Let $N(\mathbf{w})$ be the number of occurrences of the word $\mathbf{w} := w_1 w_2 \cdots w_h$ in the sequence $X_1 X_2 X_3 \cdots X_\ell$ (model M1) :

$$N(\mathbf{w}) = \sum_{i=1}^{\ell-h+1} Y_i$$

where

$$Y_i = \mathbf{1}\{\mathbf{w} \text{ starts at position } i\} \sim \mathcal{B}(\mu(\mathbf{w}))$$

and

$$\mu(\mathbf{w}) = \mu(\mathbf{w}_1) \prod_{j=1}^{h-1} \pi(\mathbf{w}_j, \mathbf{w}_{j+1}).$$

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Question : how to decide if $N^{obs}(\mathbf{w})$ is significantly unexpected (under model M1)?

Ideally : one should compute the *p*-value $\mathbb{P}(N(\mathbf{w}) \ge N^{\text{obs}}(\mathbf{w}))$ or at least compare $N^{\text{obs}}(\mathbf{w})$ with the expected count $\mathbb{E}N(\mathbf{w})$ Sophie Schbath (INBA - MalAGE) Histoire de mots ALEA 2017

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Hint

If the Y_i's were independent, we would have

$$\sum_{i=1}^{\ell} Y_i \sim \mathcal{B}(\ell, \mu(\mathbf{w})) \text{ approx by } \begin{cases} \mathcal{P}(\ell\mu(\mathbf{w})) & \text{if } \ell\mu(\mathbf{w}) \text{ small}, \\ \mathcal{N}(\ell\mu(\mathbf{w}), \ell\mu(\mathbf{w})(1-\mu(\mathbf{w}))) & \text{if } \ell\mu(\mathbf{w}) \sim \infty. \end{cases}$$

But the Y_i's are not independent (overlaps) :

- For non-overlapping words, such as ATGAC, $Y_i = 1 \Rightarrow Y_{i+1} = 0$.
- For overlapping words, such as ATGAT,

$$\mathbb{P}(Y_{i+3} = 1 \mid Y_i = 1) > \mathbb{P}(Y_{i+3} = 1).$$

Scores of exceptionality

• In the 80's, the ratio $\frac{N^{obs}(\mathbf{w})}{\mathbb{E}N(\mathbf{w})}$ was used with

$$\mathbb{E}N(\mathbf{w}) = (\ell - h + 1)\mu(\mathbf{w}) = (\ell - h + 1)\mu(w_1)\prod_{j=1}^{h-1}\pi(w_j, w_{j+1})$$

 \rightarrow problem with the variability around 1 : Var(N)?

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- \rightarrow problem with the variability around 1 : Var(N)?
- Normalization by EN(w) like for a Poisson variable [Brendel et al. (86)]

$$rac{\mathsf{V}^{\mathsf{obs}}(\mathbf{w}) - \mathbb{E} \mathcal{N}(\mathbf{w})}{\sqrt{\mathbb{E} \mathcal{N}(\mathbf{w})}}$$
 .

 \rightarrow problem with the variability around 0 : Var(N) $\neq \mathbb{E}(N)$.

Scores (2)

• The variance formula was published in 1992 by Kleffe & Borodowsky. The overlapping structure of the word clearly appears.

$$\begin{aligned} \text{Var}[N(\mathbf{w})] &= (\ell - h + 1)\mu(\mathbf{w})[1 - \mu(\mathbf{w})] \\ &+ 2\mu(\mathbf{w})\sum_{d=1}^{h-1} (\ell - h - d + 1) \left[\varepsilon_{h-d}(\mathbf{w})\prod_{j=h-d+1}^{h} \pi(w_{j-1}, w_j) - \mu(\mathbf{w})\right] \\ &+ 2\mu^2(\mathbf{w})\sum_{t=1}^{\ell-2h+1} (\ell - 2h - t + 2) \left[\frac{1}{\mu(w_1)}\pi^t(w_h, w_1) - 1\right] \end{aligned}$$

One then uses the *z*-score and the Central Limit Theorem :

$$\frac{N(\mathbf{w}) - \mathbb{E}N(\mathbf{w})}{\sqrt{\text{Var}(N(\mathbf{w}))}} \longrightarrow \mathcal{N}(0, 1) \text{ as } \ell \to \infty.$$



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 \rightarrow problem when parameters (π,μ) are unknown and have to be estimated by their MLE $(\widehat{\pi},\widehat{\mu})$

Indeed,

$$\operatorname{Var}(N - \widehat{\mathbb{E}(N)}) \neq \widehat{\operatorname{Var}(N)}$$

Scores (3)

• Prum, Rodolphe, de Turckheim (95) proposed an appropriate normalizing factor $\hat{\sigma}$ for $N - \widehat{\mathbb{E}N}$ which depends on the overlapping structure of the word.

It leads to the following score

$$\frac{N(\mathbf{w}) - \widehat{\mathbb{E}N(\mathbf{w})}}{\widehat{\sigma(\mathbf{w})}} \longrightarrow \mathcal{N}(0, 1) \text{ as } \ell \to \infty.$$

and an approximation of the *p*-value :

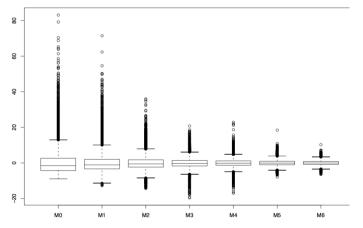
$$\mathbb{P}(\textit{\textit{N}} \geq \textit{\textit{N}}^{obs}) \simeq \mathbb{P}\left(\mathcal{N}(0,1) \geq \frac{\textit{\textit{N}}^{obs} - \widehat{\mathbb{EN}}}{\widehat{\sigma}}\right)$$

A similar score has been derived under model Mm.

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Histoire de mots

Which model to use?



Scores of exceptionality for the 65,536 8-letter words in the *E.coli* backbone.

Example : gctggtgg occurs 762 times in the *E. coli*'s genome (leading strands, $\ell = 4.610^6$).

model	fit	expected	score	<i>p</i> -value	rank
M00	length	70.783			
M0	bases	85.944	72.9	$< 10^{-323}$	3
M1	dinucl.	84.943	73.5	$< 10^{-323}$	1
M2	trinucl.	206.791	38.8	$< 10^{-323}$	1
МЗ	tetranucl.	355.508	22.0	1.4 10 ⁻¹⁰⁷	5
M4	pentanucl.	355.312	22.9	2.310 ⁻¹¹⁶	2
M5	hexanucl.	420.867	19.7	1.0 10 ⁻⁸⁶	1
M6	heptanucl.	610.114	10.6	1.5 10 ⁻²⁶	3

Influence of the model (2)

	gctggtgg 762 occ.			cgctgg 28 occ.	ccggccta 71 occ.	
M0	85.944	< 10 ⁻³²³ (3)	85.524	< 10 ⁻³²³ (2)	70.445	0.47 (25608)
M1	84.943	< 10 ⁻³²³ (1)	125.919	< 10 ⁻³²³ (2)	48.173	10 ⁻³ (13081)
M2	206.791	< 10 ⁻³²³ (1)	255.638	10^{-283} (3)	35.830	10 ⁻⁸ (4436)
M3	355.508	1.4 10 ⁻¹⁰⁷ (5)	441.226	10 ⁻⁷⁸ (15)	14.697	10 ⁻⁴⁹ (47)
M4	355.312	2.310 ⁻¹¹⁶ (2)	392.252	10 ⁻¹²⁰ (1)	15.341	10 ⁻⁴⁶ (21)
M5	420.867	1.0 10 ⁻⁸⁶ (1)	633.453	10 ⁻²² (24)	27.761	10 ⁻¹⁸ (36)
M6	610.114	1.5 10 ⁻²⁶ (3)	812.339	0.16 (14686)	25.777	10 ⁻²⁶ (4)

Expected counts and *p*-values (rank) under models Mm,

$$m=0,1,\ldots,6,$$

estimated from the E. coli's genome (4638858 bps, leading strands).

The Gaussian approximation appeared to be not accurate for expectedly rare words $(\mathbb{E}(N(\mathbf{w})) = O(1) \text{ as } \ell \to +\infty)$.

Here "w" is rare along the sequence.

- If w is not self-overlapping : N(w) ~ Pois(E[N(w)]) (Chen-Stein method).
- In the general case, N(w) is approximated by a Geometric-Poisson distribution with parameter ((1 a(w))E[N(w)]; a(w)) (S. (95)).

Both (compound) Poisson approximations are still valid when plugging the estimated parameters of the Markov model. In the general case : clump decomposition



$$N(\mathbf{w}) = \sum_{c=1}^{\widetilde{N}(\mathbf{w})} K_c$$

where

• $\widetilde{N}(\mathbf{w})$ is the number of clumps

• *K_c* is the size of the *c*-th clump

In the general case : clump decomposition



$$N(\mathbf{w}) = \sum_{c=1}^{N(\mathbf{w})} K_c$$



is the overlapping proba.

where

- *N*(w) is the number of clumps
 can be approximated by *Pois*((1 − *a*(w))E[*N*(w)]) (Chen-Stein)
- K_c is the size of the c-th clump

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where

- *N*(w) is the number of clumps
 can be approximated by *Pois*((1 − *a*(w))E[*N*(w)]) (Chen-Stein)
- K_c is the size of the c-th clump follows a geometric distribution G(a(w))

Chen(75), Stein(71), Arratia et al. (89)

$$egin{aligned} Y_i &\sim \mathcal{B}(p_i) & N = \sum_{i \in I} Y_i \ Z_i &\sim \mathcal{P} \mathrm{o}(p_i) ext{ indep.} & Z = \sum_{i \in I} Z_i \ d_{\mathsf{TV}}(\mathcal{L}(N) - \mathcal{L}(Z)) &\leq 2(b_1 + b_2 + b_3) \end{aligned}$$

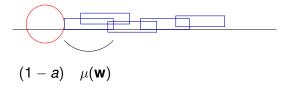
where $b_1 = \sum_{i \in I} \sum_{j \in B_i} \mathbb{E} Y_j \mathbb{E} Y_j$, B_i is any neighborhood of *i* in *I*

$$b_2 = \sum_{i \in I} \sum_{j \in B_i \setminus \{i\}} \mathbb{E}(Y_i Y_j)$$

$$b_3 = \sum_{i \in I} \mathbb{E} | \mathbb{E}(Y_i - p_i | \sigma(Y_j, j \notin B_i)) |.$$

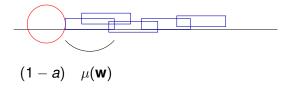
Geometric distribution for the clump size

Probability for a clump of w to start at a given position

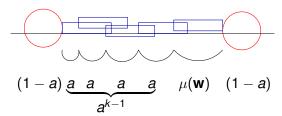


Geometric distribution for the clump size

Probability for a clump of w to start at a given position



Probability for a k-clump of w to start at a given position



Under model M1 (with known parameters)

- the exact distribution of the count $N(\mathbf{w})$ can be computed
 - via its generating function [Régnier (00)],
 - via the duality equation P(N(w) ≥ x) = P(T_x ≤ ℓ) where T_x is the position of the *x*-th occurrence; The distribution of T_x can be obtained by recursion or via its generating function (Robin & Daudin (99), Stefanov (03)).
- large deviation technique can be used to directly approximate the p-value [Nuel (04)] :

$$\mathbb{P}(N \ge N^{\text{obs}}) \simeq \exp(-\ell I(N^{\text{obs}})).$$

It is a very accurate (but numerically costly) method for very exceptional words.

Prediction and identification of functional DNA motifs

Chi motifs in bacterial genomes

- Motif involved in the repair of double-strand DNA breaks. Chi needs to be frequent along bacterial genomes.
- Chi motifs have been identified for few bacterial species. They are not conserved through species.
- Known Chi motifs are 5 to 8 nucleotides long and can be degenerated.
- Moreover, Chi activity is strongly orientation-dependent (direction of DNA replication).

It is present preferentially on the leading strands (high skew).

The skew of a motif \mathbf{w} is defined by $N^{obs}(\mathbf{w})/N^{obs}(\overline{\mathbf{w}})$ where $\overline{\mathbf{w}}$ is the reverse complementary of \mathbf{w} .

E. coli as a learning case

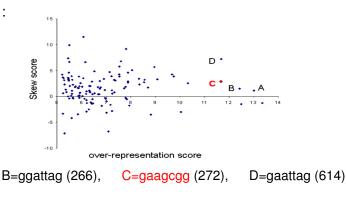
- 8-letter word GCTGGTGG
- 762 occurrences on the leading strands ($\ell = 4.6 \, 10^6$)
- Among the most over-represented 8-letter words (whatever the model Mm)
 - \Rightarrow its frequency cannot be explained by the genome composition.
- Its rank is improved if one analyzes only the backbone genome (genome conserved in several strains of the species).
- Its skew equals 3.20 (*p*-value of 3.310⁻¹¹).

The skew significance can be evaluated thanks to the Gaussian approximation of word counts.

Identification of Chi motif in S. aureus

Halpern et al. (07)

- Analysis of the *S. aureus* backbone ($\ell = 2.44 \, 10^6$).
- 8-letter words : none of the most over-represented and skewed motifs were frequent enough.
- 7-letter words :



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A=gaaaatg (1067),

Toward more complex motifs

Signature Motif of the Ter Macrodomain of E. coli

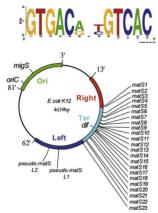
The MatP/matS Site-Specific System Organizes the Terminus Region of the *E. coli* Chromosome into a Macrodomain

Romain Mercier,¹ Marie-Agnès Petit,² Sophie Schbath,³ Stéphane Robin,⁴ Meriem El Karoul,² Frédéric Boccard,^{1,*} and Olivier Espéli^{1,*}

Cell (2008)

Use of R'MES software :

- exceptional frequency
- exceptional contrast



Definition : a word whose one or more positions may tolerate different nucleotides. The IUPAC alphabet maybe used (R=A or G, Y=C or T, N=A or C or G or T, etc.)

Examples :

- the Chi motif of *H. influenzae* is gNtggtgg
- the matS motif of E. coli is gtgacRNYgtcac
- \rightarrow one will consider them like a family ${\cal W}$ of words :

$$N(\mathcal{W}) = \sum_{\mathbf{w} \in \mathcal{W}} N(\mathbf{w})$$

Gaussian approximation [S. (95)] :

- $\mathbb{E}(N(\mathcal{W})) = \sum_{\mathbf{w} \in \mathcal{W}} \mathbb{E}(N(\mathbf{w}))$
- Var(N(W)) : need for $Cov(N(\mathbf{w}), N(\mathbf{w}'))$
 - \rightarrow one needs to know all possible overlaps between \boldsymbol{w} and \boldsymbol{w}'

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- Var(N(W)) : need for Cov(N(w), N(w'))
 - \rightarrow one needs to know all possible overlaps between \boldsymbol{w} and \boldsymbol{w}'

Compound Poisson approximation [Roquain and S. (07)] :

- mixed clumps need to be considered (again it requires all possible overlaps between any w and w' in the W family)
- the clump size is no more geometric, the overlap probability a(w) is replaced by a matrix A = (a(w, w'))
- we still get a compound Poisson distribution for N(W)

Here is an example of a PWM of length h = 5:

$$\mathbf{m} = \begin{pmatrix} 1 & 0.25 & 0 & 0.25 & 0.3 \\ 0 & 0 & 0 & 0.25 & 0.1 \\ 0 & 0.75 & 1 & 0.25 & 0 \\ 0 & 0 & 0 & 0.25 & 0.6 \end{pmatrix} \begin{array}{c} A \\ C \\ G \\ T \end{array}$$

 $m_{a,j}$ = probability of letter *a* at motif position *j*

Such representation induces a set of "compatible" words having different probabilities (or "weights")

How to count occurrences of a PWM?

...GTTCGT<u>AGGTA</u>CGG<u>TACTG</u>ATGGT<u>AAGTA</u>TG<u>AGGCT</u>... weights 0.05 0 0.02 0.1

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Classical approach : to count the number of "hits" i.e. $\sum_{i} 1\{\nu_i \ge \alpha\}$

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Othewise, there exists dedicated results [Touzet & Varré (07)], [Pape et al. (08)], [Turatsinze et al. (08)].

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Another approach : weighted count

Drawbacks of the classical approach :

- choice of the threshold α
- the hits are not weighted anymore

 \rightarrow New approach : to directly study the distribution of the weighted count defined by

$$T(\mathbf{m}) = \sum_{i=1}^{\ell-h+1} \nu_i.$$

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Note : if \mathbf{m} is a word, there is a unique compatible word and both counts are equal to the total word count

Expectation and variance of $T(\mathbf{m})$ can be analytically derived

A Gaussian approximation can be performed

A compound Poisson approximation has been derived for $T = \sum_{c=1}^{C} K_c$:

- the number *C* of clumps of compatible words can be approximated by a Poisson variable with explicit parameter (Chen-Stein method)
- the distribution of K_c , the total weight of the *c*th clump can be simulated

A compound Poisson approximation is better than a Gaussian approximation as soon as occurrences of compatible words are rare (h large enough and card(compatible words)<< 4^{h}).

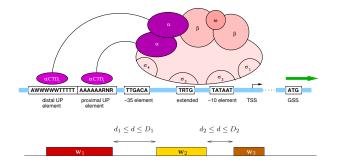
NBS motif

	1	2	3	4	5	6	7	8	9	10	11	12	13	14
A	.88	0	0	0	0	0	0	0	0	0	0	0	0	.88
С	0	0	0	.11	1	.88	.73	.73	.88	1	.11	0	0	0
G	0	0	0	.11	0	0	.12	.12	0	0	.11	0	0	0
Т	.12	1	1	.78	0	.12	.15	.15	.12	0	.78	1	1	.12

To be done :

- check the Gaussian approximation is good when few 0's in the PWM
- study the influence of the parameter estimation
- generalize to Markovian sequences

Structured motifs



What is the probability for a structured motif to occur in a given sequence?

Difficulty : even for 2 boxes, previous results on word counts cannot be used because the overlapping structure is too complicated. The structure of the motif need to be considered.

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- The 2 next approaches rely on the exact distribution of the following intersite distances in Markovian sequences (recursive formula or probability generating function) [Robin and Daudin (01)], [Stefanov (03)] :
 - $T_{\alpha,\mathbf{w}}$, the waiting time to reach pattern **w** from state α
 - $T_{\mathbf{w},\mathbf{w}'}$, the waiting time to reach pattern \mathbf{w}' from pattern \mathbf{w}

Structured motifs (3)

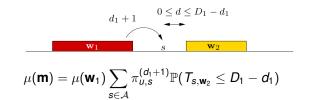
A first order approximation ([Robin et al. (02)])

• The probability $\mathbb{P}(N(\mathbf{m}) = 0) = 1 - \mathbb{P}(N(\mathbf{m}) \ge 1)$ is approximated by

$$(1 - \mu(\mathbf{m})) \left(1 - \gamma(\mathbf{m})\right)^{\ell - |\mathbf{m}|}$$

where $\mu(\mathbf{m}) = \mathbb{P}(\mathbf{m} \text{ occurs at position } i)$
 $\gamma(\mathbf{m}) = \mathbb{P}(\mathbf{m} \text{ at } i | \mathbf{m} \text{ not at } i - 1)$

The occurrence probability of m is calculated like



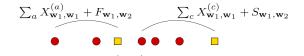
where T_{s,w_2} is the waiting time to reach pattern w_2 from state s

Structured motifs (4)

An exact approach via random sums ([Stefanov et al. (06)])

- Assumption : w₂ should not occur in between the 2 boxes.
- The explicit formula for the pgf of τ_m is given thanks to the following decomposition : /

$$\tau_{\mathbf{m}} \stackrel{\mathcal{D}}{=} T_{\alpha,\mathbf{w}_{1}} + \sum_{b=1}^{L'} \left(\underbrace{\sum_{a=1}^{L_{1}} X_{\mathbf{w}_{1},\mathbf{w}_{1}}^{(ab)} + F_{\mathbf{w}_{1},\mathbf{w}_{2}}^{(b)} + T_{\mathbf{w}_{2},\mathbf{w}_{1}}^{(b)}}_{\stackrel{\mathbb{D}}{=} T_{\mathbf{w}_{1},\mathbf{w}_{2}} \mid \text{failure}} \right) + \underbrace{\sum_{c=1}^{L_{2}} X_{\mathbf{w}_{1},\mathbf{w}_{1}}^{(c)} + S_{\mathbf{w}_{1},\mathbf{w}_{2}}}_{\stackrel{\mathbb{D}}{=} T_{\mathbf{w}_{1},\mathbf{w}_{2}} \mid \text{success}}$$



 L_1 , L_2 and L' are independent geometric variables.

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