

TOPIC MODELING ANALYSIS OF DIFFERENTIAL GENE EXPRESSION

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1. Differential gene expression. The “log-fold change” statistic is commonly used in microarray and RNA sequencing experiments to quantify expression differences between two conditions (e.g., [2, 3]). To motivate the ideas below, I write the log-fold change for gene j and condition k as a ratio of two conditional expectations,

$$(1) \quad \text{lfc}(j, k) \equiv \log_2 \frac{E[x_j \mid \text{condition} = k]}{E[x_j \mid \text{condition} \neq k]},$$

where x_j is the measured expression level (e.g., UMI count) of gene j . In experiments where the conditions are inferred—for example, by running a machine learning algorithm to cluster the expression profiles—this quantity could represent the difference in gene expression between cells inside and outside a cluster.¹

The aim of the next sections is to define an analogue to the log-fold change statistic for topic modeling.

2. The multinomial topic model and Poisson non-negative matrix factorization. Here we briefly describe the multinomial topic model, and its connection to Poisson non-negative matrix factorization (Poisson NMF).

We begin with the “bag of words” description, which was used to describe the LDA model [1]. In this view, each document (or gene expression profile) i is represented as a vector of terms/genes, $w_i = (w_{i1}, \dots, w_{is_i})$, where s_i is the size of document i . (The order of the words or genes appearing in this vector doesn’t matter, hence the “bag of words.”) Each $w_{it} \in \{1, \dots, m\}$ is term/gene j with probability $p(w_{it} = j \mid z_{it} = k) = f_{jk}$, in which we have introduced z_{it} , a variable indicating which topic $k \in \{1, \dots, K\}$ the word/gene is drawn from. The topic indicator variables for document i are in turn generated according to $p(z_{it} = k) = l_{ik}$.

This process also defines a *multinomial* model for an $n \times m$ matrix of counts x_{ij} :

$$(2) \quad x_{i1}, \dots, x_{im} \sim \text{Multinom}(x_{i1}, \dots, x_{im}; s_i, \pi_i),$$

where $x_{ij} = \sum_{t=1}^{s_i} \delta_j(w_{it})$ is the number of times term/gene j appears in document/cell i , and the probabilities π_{ij} are weighted sums of the “factors” f_{jk} ,

$$(3) \quad \pi_{ij} = \sum_{k=1}^K l_{ik} f_{jk}.$$

The log-likelihood for the multinomial topic model, ignoring terms that do not depend on the model parameters, has a simple expression:

$$(4) \quad \log p(x) = \sum_{i=1}^n \sum_{j=1}^m x_{ij} \log(\sum_{k=1}^K l_{ik} f_{jk}).$$

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¹Supposing n_k out of a total of n gene expression profiles (cells) are from condition k , then $\text{lfc}(j, k)$ can be computed as $\text{lfc}(j, k) = \log_2 \{n_{jk} / (n_j - n_{jk}) \times (n - n_k) / n_k\}$, where n_j is the total expression of gene j among all expression profiles, and n_{jk} is the total expression of j among all cells in condition (or cluster) k .

As we have shown elsewhere, the multinomial topic model is closely related to a Poisson non-negative matrix factorization of the count data,

$$(5) \quad x_{ij} \sim \text{Poisson}(\lambda_{ij}),$$

where $\lambda_{ij} = \sum_{k=1}^K \hat{l}_{ik} \hat{f}_{jk}$. Given a Poisson NMF fit, an equivalent multinomial topic model can be easily recovered, as we have shown elsewhere.

3. Gene expression differences in topics. Returning to the question of assessing differential gene expression, there are two new twists when done in the context of topic modeling:

1. The cluster (topic) assignments are probabilistic.
2. The cluster assignments are made at the level of genes, not cells.

I propose a log-fold change statistic to address these two points. It compares the probability of gene j occurring ($w = j$) given topic k ($z = k$) versus the probability given assignment a topic other than k ($z \neq k$):

$$(6) \quad \text{lfc}(j, k) \equiv \log_2 \frac{p(w = j | z = k)}{p(w = j | z \neq k)}.$$

For a given gene j and topic k , $\text{lfc}(j, k)$ can be calculated as

$$(7) \quad \begin{aligned} \text{lfc}(j, k) &= \log_2 \left\{ \frac{p(w = j, z = k)}{p(w = j, z \neq k)} \times \frac{p(z \neq k)}{p(z = k)} \right\} \\ &= \log_2 \left\{ \frac{\sum_{i=1}^n \sum_{t=1}^{s_i} \delta_j(w_{it}) \phi_{ijkt}}{\sum_{i=1}^n \sum_{t=1}^{s_i} \delta_j(w_{it}) (1 - \phi_{ijkt})} \times \frac{\sum_{i=1}^n \sum_{j'=1}^m \sum_{t=1}^{s_i} \delta_{j'}(w_{it}) (1 - \phi_{ij'kt})}{\sum_{i=1}^n \sum_{j'=1}^m \sum_{t=1}^{s_i} \delta_{j'}(w_{it}) \phi_{ij'kt}} \right\}, \end{aligned}$$

where ϕ_{ijkt} denotes the posterior probability of $z_{it} = k$ given $w_{it} = j$,

$$(8) \quad \begin{aligned} \phi_{ijkt} &\equiv p(z_{it} = k | w_{it} = j) \\ &= \frac{p(w_{it} = j | z_{it} = k) p(z_{it} = k)}{\sum_{k'=1}^K p(w_{it} = j | z_{it} = k') p(z_{it} = k')} \\ &= \frac{l_{ik} f_{jk}}{\sum_{k'=1}^K l_{ik'} f_{jk'}}. \end{aligned}$$

Since the topic assignments z_{it} do not depend on t —that is, we can drop the “ t ” subscript from the ϕ_{ijkt} ’s—the expression for the lfc simplifies:

$$(9) \quad \text{lfc}(j, k) = \log_2 \left\{ \frac{\sum_{i=1}^n x_{ij} \phi_{ijk}}{\sum_{i=1}^n x_{ij} (1 - \phi_{ijk})} \times \frac{\sum_{i=1}^n \sum_{j'=1}^m x_{ij'} (1 - \phi_{ij'k})}{\sum_{i=1}^n \sum_{j'=1}^m x_{ij'} \phi_{ij'k}} \right\}.$$

At the maximum-likelihood solution (MLE) of the l_{ik} ’s and f_{kl} ’s, the lfc statistic simplifies further:

$$(10) \quad \text{lfc}(j, k) = \log_2 \left\{ \frac{\sum_{i=1}^n x_{ij} \phi_{ijk}}{\sum_{i=1}^n x_{ij} (1 - \phi_{ijk})} \times \frac{\sum_{i=1}^n s_i (1 - l_{ik})}{\sum_{i=1}^n s_i l_{ik}} \right\}.$$

This is because, at the MLE, the loadings s_{ik} , $k = 1, \dots, K$, for a given document/cell i should be proportional to the sums $\sum_{j=1}^m x_{ij} \phi_{ijk}$.

Finally, it is convenient that the lfc (7) will be the same if we replace the multinomial topic model parameters l_{ik} and f_{jk} with the corresponding parameters of the Poisson NMF, \hat{l}_{ik} and \hat{f}_{jk} (proof not given).

REFERENCES

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