

Seminar for Statistics

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# Comparison of EM-algorithm and MLE using Cholesky decomposition

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#### Abstract

The intent of this work is to compare The EM algorithm to a MLE approach in the case of multivariate normal mixture models using the Cholesky decomposition. The EM algorithm is widely used in statistics and is proven to converge, however in pathological cases convergence slows down considerably. MLE doesn't have this particular error, but is computationally costly. The Cholesky decomposition cuts down the necessary parameters almost in half....

methods(not done) results(not done)

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## Chapter 1

# Introduction to normal mixture models

here intro to normal mixtures

A good and thorough introductory book is the work of McLachlan and Peel 2000 and the reader is encouraged to study that to learn in depth about normal mixtures. We will here give a short overwiev of normal mixtures to fix notation and nomenclature.

Let  $\mu \in \mathbb{R}^p$ ,  $\Sigma \in \mathbb{R}^{p \times p}$  and  $\phi(\mu, \Sigma)$  be the normal distribution with mean  $\mu$  and covariance matrix  $\Sigma$ .

Normal mixture model are designed for situations where we assume that a given dataset originates from more than one population of explaining variables.

$$Y_1, \ldots, Y_{\infty}$$

**Definition 1.0.0.1.** Suppose we have a random sample  $Y_1, \ldots, Y_n$  with probability density function  $Y_j \sim f(y_j)$  on  $\mathbb{R}^p$  We assume that the density  $f(y_j)$  of  $Y_j$  can be written in the form

$$f(y_j) = \sum_{i=1}^{K} \pi_i \phi_i(y_i)$$

The  $\pi_i$  are called the component densities of the mixture.

explain in scetch EM algo

explain idea to use parameter optimizer instead, EM has pathological insufficiencies, like 'getting stuck' for many iterations. we hope we need less iterations, and as concequence less time. 'special' idea: using cholesky decomp.

#### 1.1 choice of notation

describe difference in notation between ceuleux & govaert and our covariance matrix decomposition.

The classification of models in this paper relies heavily on the work of Celeux and Grovaert, however, out of necessity for clarity, we break with their notation. So as to not confuse the reader we describe here in depth the differences in notation between Celeux and Govaert and ours.

explanation for the volume, shape and orientation descriptors

The basis of classification in CnG is the decomposition of a symmetric matrix into an orthogonal and a diagonal component. A symmetric positive definite matrix  $\Sigma$  can be decomposed as follows

$$\Sigma = \lambda \boldsymbol{D} \boldsymbol{A} \boldsymbol{D}^{\mathsf{T}}$$

with D an orthogonal matrix and A a diagonal matrix and  $\lambda = \sqrt[p]{det(\Sigma)}$  the p-th root of the determinant of  $\Sigma$ .

This decomposition has an appealing geometric interpretation, with D as the *orientation* of the distribution, A the *shape*, and  $\lambda$  the *volume*. The problem of notation comes from standard conventions in linear algebra, where the letters A and D are usually occupied by arbytrary and diagonal matrices respectively. Furthermore, we intend to apply a variant of the Cholesky decomposition to  $\Sigma$ , the  $LDL^{\top}$  decomposition. This obviously raises some conflicts in notation.

Therefore we, from here on, when reffering to the decomposition as described by cng, will use the following modification of notation:

$$\begin{aligned} \boldsymbol{D} &\longmapsto \boldsymbol{Q} \\ \boldsymbol{A} &\longmapsto \boldsymbol{\Lambda} \\ \boldsymbol{\lambda} &\longmapsto \boldsymbol{\alpha} \\ \boldsymbol{\Sigma} &= \boldsymbol{\lambda} \boldsymbol{D} \boldsymbol{A} \boldsymbol{D}^\top = \boldsymbol{\alpha} \boldsymbol{Q} \boldsymbol{\Lambda} \boldsymbol{Q}^\top \end{aligned}$$

These were chosen according to general conventions of linear algebra. Q is usually chosen for orthonormal matrices;  $\Lambda$  is often a choice for eigen vectors and  $\alpha$  was somewhat arbitrarily chosen.

make clear that the models can not be translated one to one to ldlt model make nice table(maybe sideways to account for parameter list)

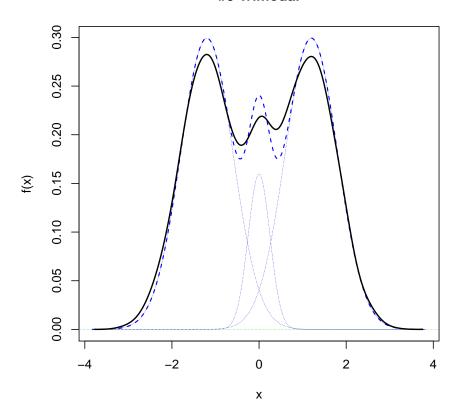
count													$1 + pK + K \frac{p(p-1)}{2}$	$K + pK + K\frac{p(p-1)}{2}$
parameters													$\lambda, d_{i,k}, l_{i,j,k} \ j > i - 1 + pK + K \frac{p(p-1)}{2}$	$\lambda_k, d_{i,k}, l_{i,j,k} \ j > i  K + pK + K \frac{p(p-1)}{2}$
$_{\perp}TQT$	same as $C\&G$						don't exist						$lpha oldsymbol{L}_k oldsymbol{D}_k oldsymbol{L}_k^ op$	$lpha_koldsymbol{L}_koldsymbol{D}_koldsymbol{L}_k^ op$
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parameters	α	$\alpha_k$	$lpha, \lambda_i$	$lpha_k, \lambda_i$	$lpha, \lambda_{i,k}$	$lpha_k, \lambda_{i,k}$	$\alpha, \lambda_i, q_{i,j}$	$lpha, \lambda_{i,k}, q_{i,j}$	$lpha_k, \lambda_i, q_{i,j}$	$\alpha_k, \lambda_{i,k}, q_{i,j}$	$lpha, \lambda_i, q_{i,j,k}$	$lpha_k, \lambda_i, q_{i,j,k}$	$lpha, \lambda_i, q_{i,j,k}$	$lpha_k, \lambda_i, q_{i,j,k}$
orientation	ı	ı	coordinate axes	coordinate axes	coordinate axes	coordinate axes	ednal	ednal	ednal	ednal	variable	variable	variable	variable
$_{ m shape}$	ednal	ednal	ednal	ednal	variable	variable	ednal	variable	ednal	variable	equal	ednal	variable	variable
volume	ednal	variable	ednal	variable	ednal	variable	ednal	ednal	variable	variable	ednal	variable	equal	variable
$oldsymbol{\Sigma}_k$ C&G	$\alpha m{I}$	$lpha_k m{I}$	$\Delta \Delta$	$lpha_k \mathbf{A}$	$lpha oldsymbol{\Lambda}_k$	$lpha_k \mathbf{\Lambda}_k$	$lpha oldsymbol{Q} oldsymbol{V} oldsymbol{Q}$	$lpha oldsymbol{Q} oldsymbol{\Lambda}_k oldsymbol{Q}^{ op}$	$lpha_k oldsymbol{Q} oldsymbol{\Lambda} oldsymbol{Q}^{ op}$	$lpha_k oldsymbol{Q} oldsymbol{\Lambda}_k oldsymbol{Q}^ op$	$lpha oldsymbol{Q}_k oldsymbol{\Lambda} oldsymbol{Q}_k^ op$	$lpha_k oldsymbol{Q}_k oldsymbol{\Lambda} oldsymbol{Q}_k^{ op}$	$lpha oldsymbol{Q}_k oldsymbol{\Lambda}_k oldsymbol{Q}_k^{ op}$	$lpha_k oldsymbol{Q}_k oldsymbol{\Lambda}_k oldsymbol{Q}_k^ op$
Model	EII	VII	EEI	VEI	EVI	VVI	EEE	EVE	VEE	VVE	EEV	VEV	EVV	VVV

### 1.2 problems of EM

the EM algo has stalling problems especially close to a local optimum show an example using nor1mix

> library("nor1mix")
> set.seed(2019)
> plot(MW.nm9, lty=2, col = "blue", p.norm=FALSE, p.comp=TRUE)
> x9 <- rnorMix(5000, MW.nm9)
> lines(density(x9), lwd=1.8)# "clearly" 3 components

#### #9 Trimodal



then an illustration of MW examples of pathological cases

# Chapter 2

# placeholder

placeholder

6 placeholder

## **Bibliography**

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8 BIBLIOGRAPHY

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