

Impact of Missing Data on Mixtures and Clustering

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Impact of Missing Data on Mixtures and Clustering

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Take home message

Missing data may change preconceptions

- Mixtures: EM has unexpected behaviour concerning degeneracy dynamic
- Clustering: the missing data pattern may convey some information on partition

These topics are in the research agenda of Statisticians since:

The larger the datasets, the more missing data may appear...

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Outline

1 Introduction

- 2 Impact of MAR data on the EM algorithm
- 3 Impact of MNAR data on clustering
- 4 Concluding remarks

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Outline

1 Introduction

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Missing data: notations

- $y = \{y_1 | \dots | y_n\}^T$: full dataset with *n* individuals
- $y_i = (y_{i1}, \dots, y_{id}) \in \mathcal{Y} = \mathbb{R}^d$ with individual $i \in \{1, \dots, n\}$
- $c = \{c_1 | \dots | c_n\}^T \in \{0, 1\}^{n \times d}$: pattern of missing data for the full dataset
- $c_i = (c_{i1}, \ldots, c_{id}) \in \{0, 1\}^d$: pattern of missing data for individual $i \in \{1, \ldots, n\}$

$$c_{ij} = 1 \Leftrightarrow y_{ij}$$
 is missing

• y_i^{obs} : the observed variables values for indiv. i (and $y^{\text{obs}} = \{y_1^{\text{obs}}| \dots |y_n^{\text{obs}}\}^T$) • y_i^{mis} : the missing variables values for individual i (and $y^{\text{mis}} = \{y_1^{\text{mis}}| \dots |y_n^{\text{mis}}\}^T$)

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Typology of the missingness mechanisms

Missing completely at random (MCAR):

$$\mathsf{p}(c|y;\psi) = \mathsf{p}(c;\psi) \quad \forall y$$

Missing at random (MAR):

$$\mathsf{p}(c|y;\psi) = \mathsf{p}(c|y^{\mathrm{obs}};\psi) \quad \forall y^{\mathrm{mis}}$$

Missing not at random (MNAR): the mechanism is not MCAR nor MAR

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Ignorable vs. non ignorable model

A missing mechanism is ignorable if likelihoods can be decomposed as

$$L(\theta, \psi; \underbrace{y^{\text{obs}}, c}_{\text{observed data}}) = L(\psi; c | y^{\text{obs}}) \times L(\theta; y^{\text{obs}})$$

Some simple algebra show that this occurs when missing mechanism is not MNAR

Inference of θ

"If the missing mechanism is ignorable then likelihood-based inferences for θ from $L(\theta; y^{\text{obs}})$ will be the same as likelihood based inference for θ from $L(\theta, \psi; y^{\text{obs}}, c)$." [Little and Rubin, 2002 Section 6.2]

- M(C)AR is ignorable
- MNAR is not ignorable

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Mixture model and clustering

- Partition (K clusters): $z = (z_1 | \dots | z_n)^T \in \{0, 1\}^{n \times K}$ where
 - $z_i = (z_{i1}, \ldots, z_{iK}) \in \{0, 1\}^K$
 - **z**_{ik} = 1 if y_i belongs to cluster k, $z_{ik} = 0$ otherwise

• Mixture model: y_1, \ldots, y_n are i.i.d. from the *d*-variate Gaussian mixture

$$f(y_i; \pi, \theta) = \sum_{k=1}^{K} \pi_k f_k(y_i; \theta_k)$$

- $\pi_k = p(z_{ik} = 1), \ \pi = (\pi_1, \ldots, \pi_K)$
- $f_k(.; \theta_k) = \phi(.; \mu_k, \Sigma_k)$ is the d-variate Gaussian distribution with mean vector μ_k and covariance matrix Σ_k
- $\theta = (\theta_1, \ldots, \theta_K)$ is the whole mixture parameter
- Clustering: MAP principle from the mixture output to estimate the partition

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Outline

2 Impact of MAR data on the EM algorithm

- Gaussian mixture degeneracy without missing data
- Gaussian mixture degeneracy with missing data

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Degeneracy genesis: unbounded likelihood

d-variate Gaussian mixture

$$f(y_i; \boldsymbol{\theta}) = \sum_{k=1}^{K} \pi_k \underbrace{\frac{1}{(2\pi)^{d/2} |\boldsymbol{\Sigma}_k|^{1/2}} \exp\left(-\frac{1}{2} (y_i - \mu_k)^T \boldsymbol{\Sigma}_k^{-1} (y_i - \mu_k)\right)}_{\phi(y_i; \mu_k, \boldsymbol{\Sigma}_k)}$$

Sampling:
$$y_1, \ldots, y_n \stackrel{i.i.d.}{\sim} p(.; \theta)$$
 without any missing data $(y^{obs} = y)$
Likelihood: $\ell(\theta; y) = \ln L(\theta; y) = \sum_{i=1}^n \ln f(y_i; \theta)$

particular center
$$\mu_2 = y_i \implies \lim_{|\Sigma_2| \to 0} \ell(\theta; y) = +\infty$$

[Kiefer and Wolfowitz, 1956] [Day, 1969]

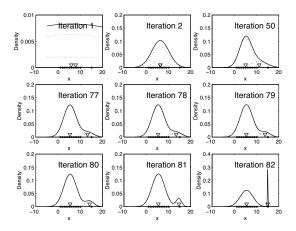
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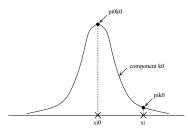
EM behaviour: illustration



- degeneracy may occur even when starting from large variances
- convergence can be slow when far from the degenerate limit
- convergence extremely fast near degeneracy

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EM behaviour: results



$$u_0 = \left[\frac{1}{p_{i_0k_0}}, \{p_{ik_0}\}_{i \neq i_0}\right]$$

degeneracy of component k_0 at y_{i_0}

 $\begin{array}{c} \Leftrightarrow \\ \| u_0 \| \to 0 \\ \text{[Biernacki and Chrétien, 2003]} \\ \text{[Ingrassia and Rocci, 2009]} \end{array}$

Proposition 1: Existence of a bassin of attraction

 $\exists \epsilon > 0 \text{ s.t. if } \|u_0\| \leq \epsilon \text{ then } \|u_0^+\| = o\|u_0\| \text{ with probability 1.}$

Proposition 2: Speed towards degeneracy is exponential

 $\begin{array}{l} \exists \epsilon > 0, \, \alpha > 0 \text{ and } \beta > 0 \text{ s.t. if } \|u_0\| \leq \epsilon \text{ then, with probability 1,} \\ |\Sigma_{k_0}^+| \leq \alpha / |\Sigma_{k_0}| \cdot \exp\Big(-\beta / |\Sigma_{k_0}|\Big). \end{array}$

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2 Impact of MAR data on the EM algorithm

Gaussian mixture degeneracy without missing data

Gaussian mixture degeneracy with missing data

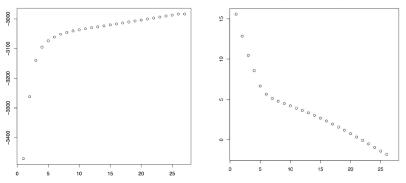
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EM behaviour illustration

- Breast cancer tissue of the UCI database repository: 106 units, 9 variables.
- 10% of missing data randomly generated
- K = 4 clusters

Log-likelihood according to the number of iterations



Decrease of the log-determinant of the degenerated component

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Detail from the illustration

	1	2	3	4	5	6	7	8	9
1	211.00		0.09	30.75	151.98	4.94	14.27	27.24	217.13
2	196.86	0.02	0.09	28.59	82.06	2.87	7.97	27.66	200.75
3	144.00	0.12	0.05	19.65	70.43	3.58		7.57	160.37
4	172.52	0.13	0.04		192.22	5.12	19.32	32.19	174.93
5	121.00	0.17	0.09	24.44	144.47	5.91	22.02	10.59	141.77
6	223.00	0.12	0.08	33.10	197.01	5.95	30.45	12.96	252.48
7		0.17	0.23	34.22	94.35	2.76	31.28	13.88	180.61
8	303.00	0.06	0.04	22.57		4.54	21.83	5.72	321.65
9	250.00	0.09	0.09	29.64	180.76	6.10	26.14	13.96	280.12
10	391.00	0.06	0.01	35.78		7.41	22.13	28.11	400.99
11	176.00	0.09	0.08	20.59	79.71		18.23	9.58	191.99
12	145.00		0.11	21.22	82.46	3.89	20.30	6.17	162.51
13	124.13	0.13	0.11	20.59			18.46	9.12	134.89
14	103.00	0.16	0.29	23.75	78.26	3.29	22.32	8.12	124.98

Table : Data belonging to the degenerated component.

- Cvg. towards a degenerated component (no plateau of the log-likelihood)
- Degeneracy relatively slow: log-likelihood linear according to the nb of it.
- Number of points of the degenerated solution greater than the space dimension
 d (but the number of complete points lower than d)

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Intermediate conclusion on missing data

<u>Like</u> the complete data y case

- Likelihood is unbounded
- EM can be attracted by degenerate solutions

<u>Unlike</u> the complete data y case

- Risk to consider a degenerated solution as valid
- Risk of losing a lot of time in useless iterations

Statisticians should be aware of such dangerous EM behaviour...

... since missing data are more and more frequent

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Understanding degeneracy speed on a toy example

- Univariate framework, no mixture, only one observed data: y
- Maximum likelihood estimator (Unbounded likelihood!): $\mu = y$, $\sigma^2 = 0$
- Suppose equivalently that n-1 data are unobserved (unchanged likelihood)
- Here is one iteration of a (useless) EM algorithm (it. q)

$$\mu^{(q+1)} = \frac{(n-1)\mu^{(q)} + y}{n} \text{ and } \sigma^{2(q+1)} = \frac{(n-1)\sigma^{2(q)} + (y-\mu^{(q+1)})^2}{n}$$

Linear grow of the log-likelihood (have a look also when *n* increases!) $\ell(\theta^{(q)}; x) \sim -0.5q \log \frac{n-1}{n}$

Geometrical convergence rate towards 0 for the variance

$$\sigma^{2(q)} \sim \sigma^{2(0)} \left(\frac{n-1}{n}\right)^q$$

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Influence of the missing data rate

% missing data	0	5	10	15	20	25	30
% deg.	16	4	12	11	46	51	100
Average nb of iterations before deg.	2	13	13	82	304	138	215

Table : Frequency and speed of degeneracy (deg.) according to the rate of missing data on the breast cancer data set.

When the rate of missing data increases:

- The rate of degeneracy increases
- The number of iterations before degeneracy seems to (globally) increase

Again, statisticians should be aware of such dangerous EM behaviour...

... since missing data are more and more frequent

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A model-based MNAR clustering approach

Medical study illustration

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Proposed MNAR models in clustering

Question we address now

Since MNAR is not ignorable, which distribution $p(c|y, z; \psi)$ to propose?

Hypothesis 1: conditional independence

$$p(c_i|y_i, z_{ik} = 1; \psi) = \prod_{j=1}^d p(c_{ij}|y_i, z_{ik} = 1; \psi)$$

Hypothesis 2: linear function within canonical link functions ρ

$$\mathsf{p}(c_{ij} = 1 \mid y_i, z_{ik} = 1; \psi) = \rho(\alpha_{kj} + \beta_{kj} y_{ij})$$

• $\psi = (\alpha, \beta)$ where $\alpha = (\alpha_{11}, \dots, \alpha_{1d}, \dots, \alpha_{K1}, \dots, \alpha_{Kd})^T \in \mathbb{R}^{Kd}$ and $\beta = (\beta_{11}, \dots, \beta_{1d}, \dots, \beta_{K1}, \dots, \beta_{Kd})^T \in \mathbb{R}^{Kd}$

• ρ is the cdf of any continuous distribution (logit, probit)

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A by-product zoology of MNAR models

	Effect variabl		Effect on members	the class hip <i>k</i>	Nb parameters	
	Depends Depends on <i>i</i> on <i>k</i>		Depends on <i>j</i>	Depends on <i>k</i>	Continuous	
$MNARz^{j}y^{k}$	 ✓ 	√	✓	✓	2Kd	
MNARyz ^j	\checkmark	×	\checkmark	\checkmark	(K+1)d	
MNARy ^k z	\checkmark	\checkmark	×	\checkmark	K(d+1)	
MNAR <i>yz</i>	\checkmark	×	×	\checkmark	(K+d)	
MNAR <i>y</i>	\checkmark	×	×	×	d	
MNARy ^k	\checkmark	\checkmark	×	×	Kd	
MNAR <i>z</i>	×	×	×	\checkmark	K	
MNAR <i>zⁱ</i>	X	×	\checkmark	\checkmark	Kd	

Remarks:

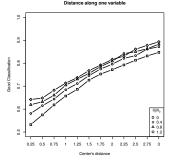
- $MNARz^j y^k$ is the most complex model
- MNARz, MNARz^{*j*}: the only effect of missingness is on the class membership, $\psi = (\alpha_{11}, ..., \alpha_{1d}, ..., \alpha_{K1}, ..., \alpha_{Kd})^T$, $p(c_{ij} = 1 | y_i, z_{ik} = 1; \psi) = \rho(\alpha_{kj})$
- MCAR is a specific and simple case

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MNARz analysis: pattern c gives information on partition z!

Draw Bayes error of a MNARz model with two components and 20% of missing data

 $\pi_k = 0.5, \; \|\mu_2 - \mu_1\|$ varies, $\Sigma_1 = \Sigma_2 = \mathsf{I}, |lpha_2 - lpha_1|$ varies



Both μ_k and α_k act on the Bayes error

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Reinterpretation of the MNARz and MNARz^j models as MAR

Commonly used in Machine Learning [Jones, 1996], [Little and Rubin, 2002], [Josse et al., 2019]

Mixture model for y^{obs} and Bernoulli distribution for C \Leftrightarrow MAR mixture model for $\tilde{y}^{obs} = (y^{obs}|c)$

For example,

$$y^{\rm obs} = \left(\begin{array}{ccc} ? & 2.6 & 5\\ {\rm blue} & 1.9 & 4\\ {\rm red} & 2.3 & ? \end{array}\right), \quad c = \left(\begin{array}{ccc} 1 & 0 & 0\\ 0 & 0 & 0\\ 0 & 0 & 1 \end{array}\right)$$

then \tilde{y}^{obs} is expressed as

$$\tilde{y}^{\rm obs} = \left(\begin{array}{ccccccc} ? & 2.6 & 5 & 1 & 0 & 0 \\ {\rm blue} & 1.9 & 4 & 0 & 0 & 0 \\ {\rm red} & 2.3 & ? & 0 & 0 & 1 \end{array} \right)$$

Proposition 3: in terms of maximum likelihood

The maximum likelihood estimate associated to the dataset $\tilde{y}^{\rm obs}$ under MAR model is the one associated to the dataset $y^{\rm obs}$ under MNARz or MNARz^j models.

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Concluding remarks

Identifiability

Previous works: [Teicher, 1963] (without NA), [Miao et al., 2016] (for MNAR data)

Proposition 4

Assume that

- **I** The marginal mixture $\sum_{k=1}^{K} \pi_k f_k(y_i; \theta_k)$ is identifiable
- **2** There exists a total ordering \leq of $\mathcal{F}_j \times \mathcal{R}$, for $j \in \{1, \ldots, d\}$ fixed, where $\mathcal{F}_j = \{f_{1j}, \ldots, f_{Kj}\}$ and $\mathcal{R} = \{\rho_1, \ldots, \rho_K\} = \{\rho(.; \psi_1), \ldots, \rho(.; \psi_K)\}$. The total ordering is s.t. $\forall k < \ell$, $F_k = \rho_k f_{kj} \leq F_\ell = \rho_\ell f_{\ell j}$ implies

$$\lim_{u \to +\infty} \frac{\rho_{\ell}(u) f_{\ell j}(u)}{\rho_k(u) f_{k j}(u)} = 0$$

Then the mixture model with one of the MNAR \ast mechanisms is identifiable up to label swapping

All MNAR models are identifiable (or at least generically identifiable) for probit/logit

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Estimation procedure overview

- Use EM or Stochastic EM (SEM) algorithms
- MNARz and MNARz^j: EM and SEM are very simple
- MNARy*: the SE step requires a within Gibbs loop, sometimes involving itself a Sampling Importance Resampling (SIR)

	E	M	SEM		
MNARz MNARz ^j		(\checkmark		
	Probit	Logit	Probit	Logit	
MNARy*	no closed form	no closed form, optim. pb	~	require algorithms as SIR (costly)	

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Model selection

Can select between MCAR and MNAR* with any information criterion (BIC, ICL)

Even if the missing mechanism is ignorable for MCAR...

 \dots need to model c to compare a MCAR and a MNAR model

CAUTION

- It is just a selection between several proposed MNAR models
- It is not deciding if missingness procedure is "genererically" MNAR or not

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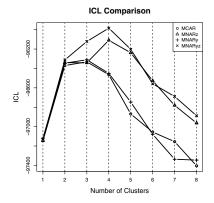
Hospital data description

- Number of patients: $n = 5 \ 146$
- Number of features: d = 7
 - Age
 - Size
 - Weight
 - Cardiac frequency
 - Hemoglobin concentration
 - Temperature
 - Minimum Diastolic and Systolic Blood Pressure
- Percentage of missing data: 6.4%

Doctors are convinced that their missing data are MNAR

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ICL comparison



• MCAR, MNARy and MNARz are equivalent until K = 3

• MNARz and MNARyz clearly indicate presence of an additional cluster (K = 4)

It seems to be an illustation of the effect of c through MNARz and MNARyz

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To conclude

Summary

- Statisticians should properly consider the potential missing data impact both from an algorithmic and from a modeling point of view
- EM: be careful about degeneracy which seems to exacerbated/masqued
- MNAR: interest of the simple but meaningful model MNARz, link with usual methods

Ongoing works

- EM: propose mechanism to identify/discard degenerate runs
- MNAR: extend to categorical, count and mixed data

Thanks!