

- Hierarchical models
 - Rats example
 - Factory example
- 8 schools example
- Exchangeability
- Computational aspects

Bayesian Statistics and Data Analysis Lecture 7

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Section 1

Hierarchical models



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Hierarchical model

- Example: Treatment effectiveness
 - in hospital j the survival probability is θ_j
 - observations y_{ij} tell whether patient i survived in hospital



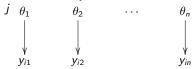


Hierarchical models

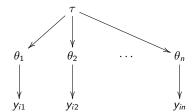
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Hierarchical model

- Example: Treatment effectiveness
 - in hospital j the survival probability is θ_j
 - observations y_{ij} tell whether patient i survived in hospital



• sensible to assume that θ_i are similar



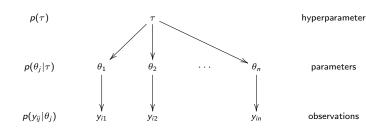
- natural to think that θ_j have common population distribution
- θ_j is not directly observed and the population distribution is unknown



Hierarchical model: terms

Lvl 1: observations given parameters $p(y_{ii}|\theta_i)$

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Joint posterior

$$p(\theta, \tau|y) \propto p(y|\theta, \tau)p(\theta, \tau)$$

 $\propto p(y|\theta)p(\theta|\tau)p(\tau)$



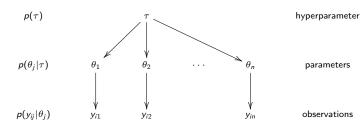
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Hierarchical model: terms

Lvl 1: observations given parameters $p(y_{ij}|\theta_j)$

Lvl 2: parameters given hyperparameters $p(\theta_j|\tau)$



Joint posterior

$$p(\theta, \tau|y) \propto p(y|\theta, \tau)p(\theta, \tau)$$

 $\propto p(y|\theta)p(\theta|\tau)p(\tau)$



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Comparisons

• "Separate model" (model with separate/independent effects) a a

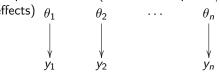




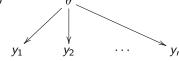
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Comparisons

• "Separate model" (model with separate/independent effects) θ_1 θ_2 \dots θ_n



• "Joint/pooled model" (model with a common effect / pooled model) θ

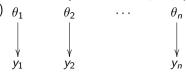




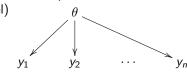
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Comparisons

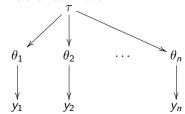
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Hierarchical model



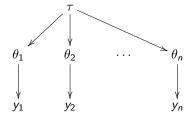


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Predictive distributions for hiearchical models

- Two types of predictive distributions
 - 1. A new observation in an existing group
 - 2. A new observation in a new group





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- Medicine testing
- Type F344 female rats in control group given placebo
 - count how many get endometrial stromal polyps
 - familiar binomial model example



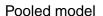
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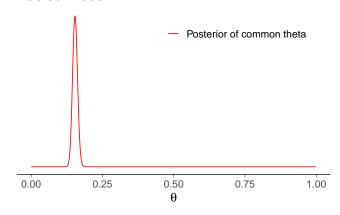
- Medicine testing
- Type F344 female rats in control group given placebo
 - count how many get endometrial stromal polyps
 - familiar binomial model example
- Experiment has been repeated 71 times

	0/20	0/20	0/20	0/20	0/20	0/20	0/20	0/19	0/19	0/19
	0/19	0/18	0/18	0/17	1/20	1/20	1/20	1/20	1/19	1/19
	1/18	1/18	2/25	2/24	2/23	2/20	2/20	2/20	2/20	2/20
	2/20	1/10	5/49	2/19	5/46	3/27	2/17	7/49	7/47	3/20
	3/20	2/13	9/48	10/50	4/20	4/20	4/20	4/20	4/20	4/20
	4/20	10/48	4/19	4/19	4/19	5/22	11/46	12/49	5/20	5/20
	6/23	5/19	6/22	6/20	6/20	6/20	16/52	15/46	15/47	9/24
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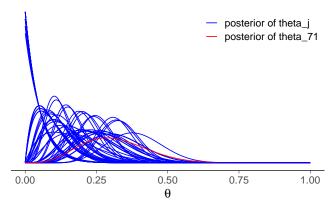






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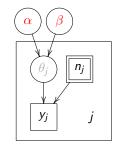


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• Hierarchical binomial model for rats prior parameters α and β are unknown

$$\theta_j | \boldsymbol{\alpha}, \boldsymbol{\beta} \sim \mathsf{Beta}(\theta_j | \boldsymbol{\alpha}, \boldsymbol{\beta})$$

$$y_j|n_j,\theta_j\sim \text{Bin}(y_j|n_j,\theta_j)$$



- Joint posterior $p(\theta_1, \dots, \theta_J, \alpha, \beta|y)$
 - multiple parameters

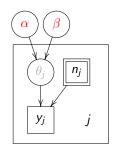


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• Hierarchical binomial model for rats prior parameters α and β are unknown

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- Joint posterior $p(\theta_1, \dots, \theta_J, \alpha, \beta|y)$
 - multiple parameters
 - factorize $\prod_{i=1}^{J} p(\theta_i | \alpha, \beta, y) p(\alpha, \beta | y)$



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- Population prior Beta $(\theta_i | \alpha, \beta)$
- Hyperprior $p(\alpha, \beta)$?
 - α, β both affect the location and scale
 - BDA3 (p. 110) has (vague) $p(\alpha, \beta) \propto (\alpha + \beta)^{-5/2}$



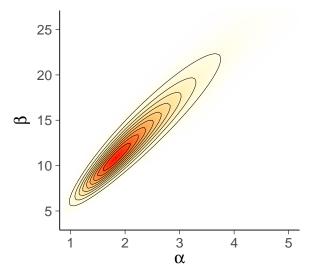
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 - BDA3 (p. 110) has (vague) $p(\alpha, \beta) \propto (\alpha + \beta)^{-5/2}$
- What type of predicitive distributions can we have?



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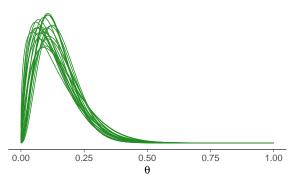
The marginal of α and β





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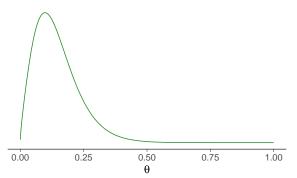






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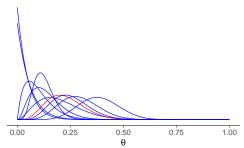






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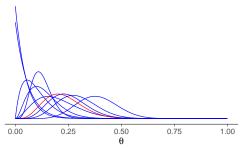




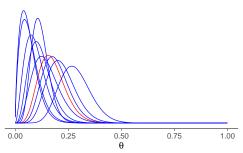


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Separate model



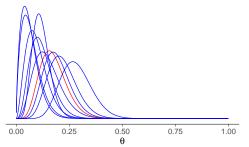
Hierarchical model



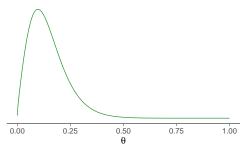


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Hierarchical model



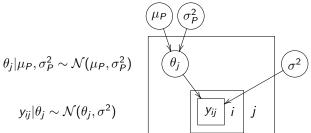
Population distribution (prior) for $\boldsymbol{\theta}_j$





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- Factory has 6 machines which quality is evaluated
- Assume hierarchical model
 - each machine has its own (average) quality θ_j and common variance σ^2

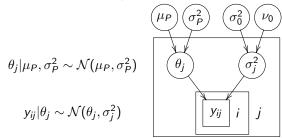


• Can be used to predict the future quality produced by each machine and quality produced by a new similar machine



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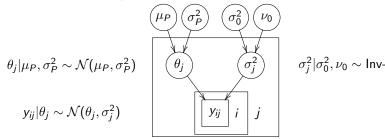


 $\sigma_j^2 | \sigma_0^2,
u_0 \sim {\sf Inv}$



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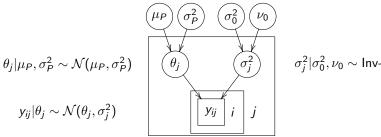


• What type of predicitive distributions can we have?



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- Factory has 6 machines which quality is evaluated
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- What type of predicitive distributions can we have?
- Can be used to predict the future quality produced by each machine and quality produced by a new similar machine



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- Example: SAT coaching effectiveness
 - in USA commonly used Scholastic Aptitude Test (SAT) is designed so that short term practice should not improve the results significantly
 - schools have anyway coaching courses
 - test the effectiveness of the coaching courses



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- Example: SAT coaching effectiveness
 - in USA commonly used Scholastic Aptitude Test (SAT) is designed so that short term practice should not improve the results significantly
 - schools have anyway coaching courses
 - test the effectiveness of the coaching courses
- SAT
 - standardized multiple choice test
 - mean about 500 and standard deviation about 100
 - most scores between 200 and 800
 - different topics, e.g., V=Verbal, M=Mathematics
 - pre-test PSAT



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- Effectiveness of the SAT coaching
 - students had made pre-tests PSAT-M and PSAT-V
 - part of students were coached
 - linear regression was used to estimate the coaching effect y_j for the school j (could be denoted with $\bar{y}_{,j}$, too) and variances σ_i^2
 - y_i approximately normally distributed, with variances assumed to be known based on about 30 students per school
 - data is group means and variances (not personal results)



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• Data: School
$$\begin{vmatrix} A & B & C & D & E & F & G & H \\ y_j & 28 & 8 & -3 & 7 & -1 & 1 & 18 & 12 \\ \sigma_j & 15 & 10 & 16 & 11 & 9 & 22 & 20 & 28 \end{vmatrix}$$



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Hierarchical normal model for group means

• J experiments, unknown θ_i and known σ^2

$$y_{ij}|\theta_j \sim \mathcal{N}(\theta_j, \sigma^2), \quad i = 1, \dots, n_j; \quad j = 1, \dots, J$$

• Group *j* sample mean and sample variance

$$\bar{y}_{.j} = \frac{1}{n_j} \sum_{i=1}^{n_j} y_{ij}$$

$$\sigma_j^2 = \frac{\sigma^2}{n_j}$$



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$$\sigma_j^2 = \frac{\sigma^2}{n_i}$$

• Use model

$$\bar{y}_{.j}|\theta_j \sim \mathcal{N}(\theta_j, \sigma_i^2)$$

this model can be generalized so that, σ_j^2 can be different from each other for other reasons than n_i

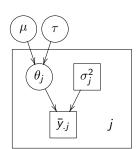


Hierarchical normal model for group means

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$$\theta_j | \mu, au \sim \mathcal{N}(\mu, au)$$

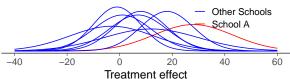
$$ar{y}_{.j}| heta_j \sim \mathcal{N}(heta_j, \sigma_j^2)$$





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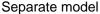


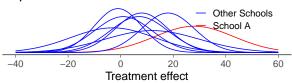




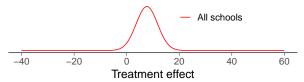
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Hierarchical normal model: 8 schools





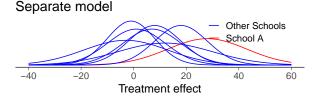
Pooled model



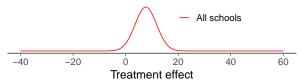


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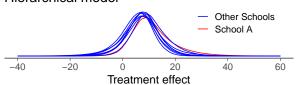
Hierarchical normal model: 8 schools



Pooled model



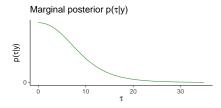
Hierarchical model





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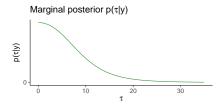
Hierarchical normal model: 8 schools

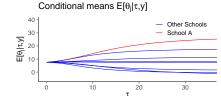




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Hierarchical normal model: 8 schools

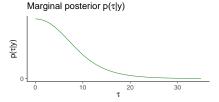


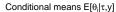


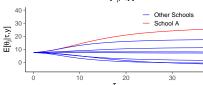


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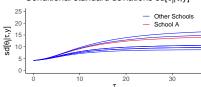
Hierarchical normal model: 8 schools







Conditional standard deviations $sd[\theta_i|\tau,y]$





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Section 2

Exchangeability



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Exchangeability

- Justifies why we can use
 - a joint model for data
 - a joint prior for a set of parameters
- Less strict than independence (IID)
- IID implies exchangeability



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Exchangeability

• Exchangeability
Random variables $\theta_1, \ldots, \theta_J$ (or y_1, \ldots, y_J) are
exchangeable if the joint distribution p is invariant to the permutation of indices
i.e

$$p(\theta_1, \theta_2, \theta_3) = p(\theta_{\pi(1)}, \theta_{\pi(2)}, \theta_{\pi(3)})$$

for any permutation π of the indicies. E.g.

$$p(\theta_1, \theta_2, \theta_3) = p(\theta_2, \theta_3, \theta_1)$$



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$$p(\theta_1, \theta_2, \theta_3) = p(\theta_2, \theta_3, \theta_1)$$

Can we come up with a situation where this doesn't hold?



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Can we come up with a situation where this doesn't hold? E.g. when we have a trend over indicies (like in time series data)



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Can we come up with a situation where this doesn't hold? E.g. when we have a trend over indicies (like in time series data)

• Exchangeability implies symmetry: If there is no information which can be used a priori to separate θ_j form each other, we can assume exchangeability. ("Ignorance implies exchangeability")



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Exchangeability: DeFinettis theorem

Let $(x_n)_{n=1}^{\infty}$ to be an infinite sequence of exchangeable random variables. De Finetti's theorem then says that there is some random variable θ so that x_j are conditionally independent given θ , and joint density for x_1, \ldots, x_J can be written in the *iid mixture* form

$$p(x_1,\ldots,x_J) = \int \left[\prod_{j=1}^J p(x_j|\theta)\right] p(\theta)d\theta$$

• Exchangeability: the data behave as if they were i.i.d., once we account for some unknown parameter θ



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- Suppose we're observing coin flips, but we're not sure about the bias θ
 - We think the flips are 'the same kind of thing' (no reason to treat flip 1 differently from flip 2): Exchangeability
 - But we might not believe they are strictly independent because were uncertain about the bias: they share information through the unknown θ



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Example

- Suppose we're observing coin flips, but we're not sure about the bias θ
 - We think the flips are 'the same kind of thing' (no reason to treat flip 1 differently from flip 2): Exchangeability
 - But we might not believe they are strictly independent because were uncertain about the bias: they share information through the unknown θ
- Given θ the flips are i.i.d.
- Without conditioning on θ , the flips are only exchangeable, not iid.



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Exchangeability

- Exchangeability does not mean that the results of the experiments could not be different
 - e.g. if we know that the experiments have been in two different laboratories, and we know that the other laboratory has better conditions for the rats, but we do not know which experiments have been made in which laboratory
 - a priori experiments are exchangeable
 - model could have unknown parameter for the laboratory with a conditional prior for rats assumed to come form the same place (clustering model)



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Exchangeability and additional information

- Example: bioassay
 - y_i number of dead animals are not exchangeable alone



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Exchangeability and additional information

- Example: bioassay
 - y_i number of dead animals are not exchangeable alone
 - x_i dose is additional information



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Exchangeability and additional information

- Example: bioassay
 - yi number of dead animals are not exchangeable alone
 - x_i dose is additional information
 - (x_i, y_i) exchangeable and logistic regression was used

$$p(\alpha, \beta|y, n, x) \propto \prod_{i=1}^{n} p(y_i|\alpha, \beta, n_i, x_i) p(\alpha, \beta)$$



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- Example: hierarchical rats example
 - all rats not exchangeable



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- Example: hierarchical rats example
 - all rats not exchangeable
 - in a single laboratory rats exchangeable



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- Example: hierarchical rats example
 - all rats not exchangeable
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- Example: hierarchical rats example
 - all rats not exchangeable
 - in a single laboratory rats exchangeable
 - laboratories exchangeable
 - \rightarrow hierarchical model can be used



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Partial or conditional exchangeability

- Conditional exchangeability
 - if y_i is connected to an additional information x_i, so that y_i are not exchangeable, but (y_i, x_i) exchangeable use joint model or conditional model (y_i|x_i).



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Partial or conditional exchangeability

- Conditional exchangeability
 - if y_i is connected to an additional information x_i , so that y_i are not exchangeable, but (y_i, x_i) exchangeable use joint model or conditional model $(y_i|x_i)$.
- Partial exchangeability
 - if the observations can be grouped (a priori), then we can use a hierarchical model



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Section 3

Computational aspects



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The funnel posterior

- Hiearchical models
 - Group-level or global parameters, e.g.

$$au \sim p(au)$$

• Local or individual-level parameters

$$\theta_i \sim \mathcal{N}(0, \tau)$$

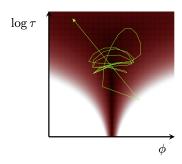
- Creates a "funnel-like" posterior geometry:
- Comes from the variance in the different layers:
 - When τ is small, the θ_i 's are concentrated around 0
 - When τ is large, the θ_i 's are widely dispersed



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Why problematic?

- 1. Pathological geometry: difficult to explore efficiently
- 2. Divergences: HMC will risk divergencies (also a good diagnostic)



Betancourt (2020)

demo



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Handling the funnel

- 1. Reduce step size (adapt_delta closer to 1)
- 2. Reparametrize using non-centered parametrization
 - 2.1 Centered parametrization

$$\theta_i \sim N(\mu, \tau)$$



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Handling the funnel

- 1. Reduce step size (adapt_delta closer to 1)
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 - 2.1 Centered parametrization

$$\theta_i \sim N(\mu, \tau)$$

2.2 Non-centered parametrization

$$\eta_i \sim N(0,1)$$

$$\theta_{\it i} = \mu + \tau \eta_{\it i}$$

