

# Bayesian Analysis with Stata: application to neonatal mortality in the UK

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MCMC

Neonatal  
Mortality

2005 data in  
Stata

2005 data in  
Mata

2005 data in  
WinBUGS

1999-2009  
data

Conclusions

# Bayesian Analysis With Stata



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## Bayesian Analysis with Stata

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### Selecting a Dirichlet Process Prior

*Posted by John in [Bayesian Analysis with Stata](#) on August 15, 2014*

This is a continuation my previous posting on non-parametric Bayesian analysis and this time I will try to show how a Dirichlet process can be used to create a family of distributions that provide much more flexible priors than the standard options such as the normal or gamma.

Last time we saw how we can represent a distribution over a finite number,  $J$ , of possible values by a set of probabilities,  $\theta_j, j=1\dots J$ , and how we can create a prior for those probabilities by using a Dirichlet

<http://staffblogs.le.ac.uk/bayeswithstata/>

# Pros and Cons of Bayes

## Advantages

- Direct answers to research questions
- Purely model based: no ad hoc modifications
- Computation can be done by simulation

# Pros and Cons of Bayes

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## Advantages

- Direct answers to research questions
- Purely model based: no ad hoc modifications
- Computation can be done by simulation

## Disadvantages

- Bayes is unfamiliar to some people
- Users must specify their priors
- Computation by simulation can be slow
- Bayesian analysis is not yet a core part of Stata

## Outline of the talk

- MCMC: Computation by simulation in Stata
- Neonatal mortality data
- Model the data for 2005
  - Stata
  - Mata
  - WinBUGS
- Model time trends 1999-2009; predict 2010

# MCMC: Computation by Simulation

## MCMC

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Conclusions

Researcher specifies:

Model:  $p(y|\theta)$

Prior:  $p(\theta)$

Posterior:

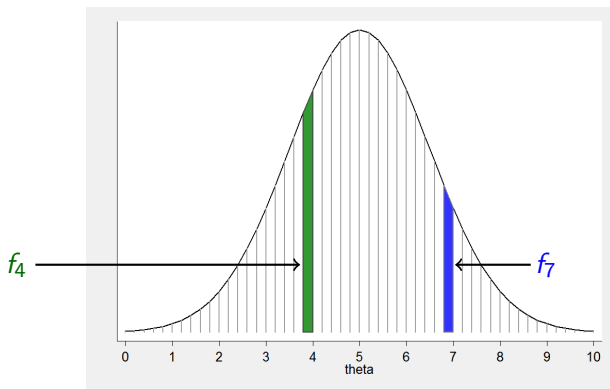
$$p(\theta|y) = \frac{p(y|\theta)p(\theta)}{\int p(y|\theta)p(\theta)d\theta} \propto p(y|\theta)p(\theta)$$

Approximate the posterior by simulating many values of  $\theta$

*For example:*

mean of simulations approximates mean of posterior

Posterior distribution  $p(\theta|y)$



$\theta = \dots, 4.0, 7.8, 5.2, \dots, 2.1, 7.0, 5.5, 4.0, \dots$

4.0 and 7.0 must occur in the ratio  $f_4:f_7$



# Markov chain Monte Carlo

## MCMC

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$\theta = \dots 4.0 \curvearrowright 7.8 \curvearrowright 5.2 \dots 2.1 \curvearrowright 7.0 \curvearrowright 7.0 \curvearrowright 4.0 \dots$

Either move to a new value or repeat the old value with transition probabilities  $t(\text{old}, \text{new})$

## Markov chain Monte Carlo

## MCMC

Neonatal  
Mortality2005 data in  
Stata2005 data in  
Mata2005 data in  
WinBUGS1999-2009  
data

Conclusions

 $\theta = \dots 4.0 \rightsquigarrow 7.8 \rightsquigarrow 5.2 \dots 2.1 \rightsquigarrow 7.0 \rightsquigarrow 7.0 \rightsquigarrow 4.0 \dots$ 

Either move to a new value or repeat the old value with transition probabilities  $t(\text{old}, \text{new})$

$$\begin{bmatrix} f_4 & f_7 \end{bmatrix} \begin{bmatrix} 1 - t(4,7) & t(4,7) \\ t(7,4) & 1 - t(7,4) \end{bmatrix} = \begin{bmatrix} f_4 & f_7 \end{bmatrix}$$

## Markov chain Monte Carlo

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$\theta = \dots 4.0 \rightsquigarrow 7.8 \rightsquigarrow 5.2 \dots 2.1 \rightsquigarrow 7.0 \rightsquigarrow 7.0 \rightsquigarrow 4.0 \dots$

Either move to a new value or repeat the old value with transition probabilities  $t(\text{old}, \text{new})$

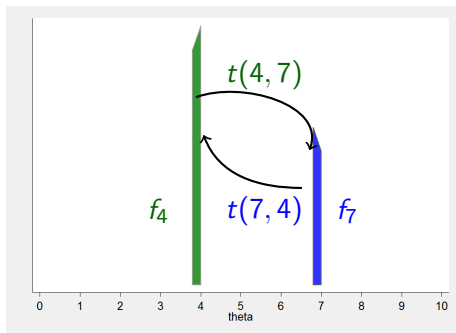
$$\begin{bmatrix} f_4 & f_7 \end{bmatrix} \begin{bmatrix} 1 - t(4, 7) & t(4, 7) \\ t(7, 4) & 1 - t(7, 4) \end{bmatrix} = \begin{bmatrix} f_4 & f_7 \end{bmatrix}$$

$$f_4[1 - t(4, 7)] + f_7 t(7, 4) = f_4$$

Detailed Balance:  $f_4 t(4, 7) = f_7 t(7, 4)$

for all pairs of values

## Metropolis-Hastings



$$P(\text{propose 4 to 7}) = \pi(4, 7)$$

$$P(\text{accept 4 to 7}) = a(4, 7)$$

Detailed Balance becomes,

$$f_4 \pi(4, 7) a(4, 7) = f_7 \pi(7, 4) a(7, 4)$$

e.g. set  $a(7,4)=1$  and make,

$$a(4, 7) = \frac{f_7 \pi(7, 4)}{f_4 \pi(4, 7)}$$

# Multi-parameter models

## Block updating

propose new values for all of the parameters and update in a single MH step

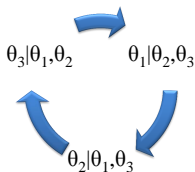
## Multi-parameter models

### Block updating

propose new values for all of the parameters and update in a single MH step

### Gibbs sampling

update each parameter in turn keeping the others fixed at their current value



## Bayesian Computing

Metropolis-Hastings: rejected moves lead to repeated values

$$\theta = \dots, 4.0, 4.0, 5.2, \dots, 2.1, 7.0, 5.5, 5.5, 4.0, \dots$$

Many repeats  $\Rightarrow$  slow convergence  $\Rightarrow$  a long chain

The trick is to choose proposals that,

- Move freely across the posterior
- Have a 'reasonable' chance of being accepted

## Bayesian Computing

Metropolis-Hastings: rejected moves lead to repeated values

$$\theta = \dots, 4.0, 4.0, 5.2, \dots, 2.1, 7.0, 5.5, 5.5, 4.0, \dots$$

Many repeats  $\Rightarrow$  slow convergence  $\Rightarrow$  a long chain

The trick is to choose proposals that,

- Move freely across the posterior
- Have a 'reasonable' chance of being accepted

Bayesian computing = Designing efficient algorithms

- Inefficient algorithms can take days to run
- *Bayesian Analysis with Stata* presents guidelines and some programs
- WinBUGS, OpenBUGS, JAGS, Stan provide black-box solutions



## A Simple Example

**Data:** twenty values from  $N(5, sd=2)$

**Researcher specifies:**

$$\text{Model: } p(y|\theta) = N(\theta, 2)$$

$$\text{Prior: } p(\theta) = N(4, 1)$$

**Posterior:**

$$p(\theta|y) \propto \left[ \prod \exp(-0.125(y_i - \theta)^2) \right] \exp(-0.5(\theta - 4)^2)$$

**log Posterior:**

$$\log [p(\theta|y)] = \text{constant} + \sum -0.125(y_i - \theta)^2 - 0.5(\theta - 4)^2$$

# A Stata Program

## MCMC

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Conclusions

```
set obs 20
gen y = rnormal(5,2)
local theta = 4
forvalues iter=1/100 {
    gen LogL = -0.125*(y-'theta')^2
    qui su LogL
    local logpost = r(sum) -0.5*('theta'-4)^2
    local newtheta = 'theta' + rnormal(0,0.25)
    qui replace LogL = -0.125*(y-'newtheta')^2
    qui su LogL
    local newlogpost = r(sum) -0.5*('newtheta'-4)^2
    if log(runiform()) < ('newlogpost' - 'logpost') local theta = 'newtheta'
    di %6.2f 'theta'
    drop LogL
}
```

# A Stata Program

## Simulate the data

```
set obs 20
gen y = rnormal(5,2)
local theta = 4
forvalues iter=1/100 {
    gen LogL = -0.125*(y-'theta')^2
    qui su LogL
    local logpost = r(sum) -0.5*('theta'-4)^2
    local newtheta = 'theta' + rnormal(0,0.25)
    qui replace LogL = -0.125*(y-'newtheta')^2
    qui su LogL
    local newlogpost = r(sum) -0.5*('newtheta'-4)^2
    if log(runiform()) < ('newlogpost' - 'logpost') local theta = 'newtheta'
    di %6.2f 'theta'
    drop LogL
}
```

# A Stata Program

Initial value

```
set obs 20
gen y = rnormal(5,2)
local theta = 4
forvalues iter=1/100 {
    gen LogL = -0.125*(y-'theta')^2
    qui su LogL
    local logpost = r(sum) -0.5*('theta'-4)^2
    local newtheta = 'theta' + rnormal(0,0.25)
    qui replace LogL = -0.125*(y-'newtheta')^2
    qui su LogL
    local newlogpost = r(sum) -0.5*('newtheta'-4)^2
    if log(runiform()) < ('newlogpost' - 'logpost') local theta = 'newtheta'
    di %6.2f 'theta'
    drop LogL
}
```

# A Stata Program

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    local newlogpost = r(sum) -0.5*('newtheta'-4)^2
    if log(runiform()) < ('newlogpost' - 'logpost') local theta = 'newtheta'
    di %6.2f 'theta'
    drop LogL
}

```

Evaluate log-posterior

# A Stata Program

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    local newtheta = 'theta' + rnormal(0,0.25)
    qui replace LogL = -0.125*(y-'newtheta')^2
    qui su LogL
    local newlogpost = r(sum) -0.5*('newtheta'-4)^2
    if log(runiform()) < ('newlogpost' - 'logpost') local theta = 'newtheta'
    di %6.2f 'theta'
    drop LogL
}

```

Make a proposal

# A Stata Program

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forvalues iter=1/100 {
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    qui su LogL
    local logpost = r(sum) -0.5*('theta'-4)^2
    local newtheta = 'theta' + rnormal(0,0.25)
    qui replace LogL = -0.125*(y-'newtheta')^2
    qui su LogL
    local newlogpost = r(sum) -0.5*('newtheta'-4)^2
    if log(runiform()) < ('newlogpost' - 'logpost') local theta = 'newtheta'
    di %6.2f 'theta'
    drop LogL
}

```

Evaluate new log-posterior

# A Stata Program

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```

set obs 20
gen y = rnormal(5,2)
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forvalues iter=1/100 {
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    qui su LogL
    local logpost = r(sum) -0.5*('theta'-4)^2
    local newtheta = 'theta' + rnormal(0,0.25)
    qui replace LogL = -0.125*(y-'newtheta')^2
    qui su LogL
    local newlogpost = r(sum) -0.5*('newtheta'-4)^2
    if log(runiform()) < ('newlogpost' - 'logpost') local theta = 'newtheta'
    di %6.2f 'theta'
    drop LogL
}

```

Accept or Reject



## MCMC

Neonatal  
Mortality

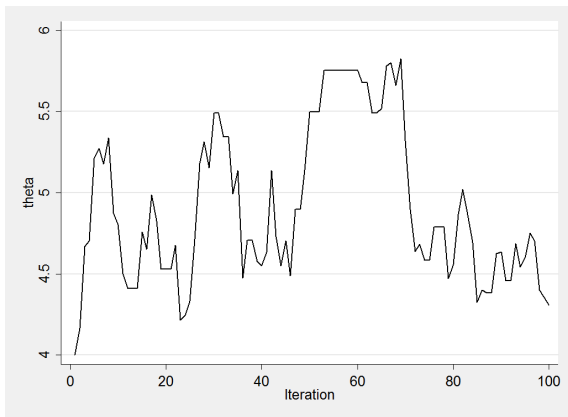
2005 data in  
Stata

2005 data in  
Mata

2005 data in  
WinBUGS

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data

Conclusions



## Simpler Code

```
set obs 20
gen y = rnormal(5,2)

program logpost
    args logp b
    local theta = 'b'[1,1]
    scalar 'logp' = 0
    logdensity normal 'logp' y 'theta' 2
    logdensity normal 'logp' 'theta' 4 1
end

matrix b = 4
mcmcrun logpost b using temp.csv, replace ///
    sampler(mhsnorm , sd(0.25)) par(theta) burn(100) update(5000)
insheet using temp.csv, comma clear
mcmctrace theta
```

## Simpler Code

```
set obs 20
gen y = rnormal(5,2)

program logpost
  args logp b
  local theta = 'b'[1,1]
  scalar 'logp' = 0
  logdensity normal 'logp' y 'theta' 2
  logdensity normal 'logp' 'theta' 4 1
end
```

Parameters in row matrix b  
Log-posterior returned in scalar logp

```
matrix b = 4
mcmcrun logpost b using temp.csv, replace ///
  sampler(mhsnorm , sd(0.25)) par(theta) burn(100) update(5000)
insheet using temp.csv, comma clear
mcmctrace theta
```

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```

set obs 20
gen y = rnormal(5,2)

program logpost
    args logp b
    local theta = 'b'[1,1]
    scalar 'logp' = 0
    logdensity normal 'logp' y 'theta' 2
    logdensity normal 'logp' 'theta' 4 1
end

matrix b = 4
mcmcrun logpost b using temp.csv, replace ///
    sampler(mhnorm , sd(0.25)) par(theta) burn(100) update(5000)
insheet using temp.csv, comma clear
mcmctrace theta
    
```

logdensity knows the formulae  
for standard distributions

## Simpler Code

### MCMC

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Conclusions

```
set obs 20
gen y = rnormal(5,2)

program logpost
    args logp b
    local theta = 'b'[1,1]
    scalar 'logp' = 0
    logdensity normal 'logp' y 'theta' 2
    logdensity normal 'logp' 'theta' 4 1
end

matrix b = 4
mcmcrun logpost b using temp.csv, replace ///
    sampler(mhsnorm , sd(0.25)) par(theta) burn(100) update(5000)
insheet using temp.csv, comma clear
mcmctrace theta
```

mcmcrun creates the chain  
and saves the values

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## Simpler Code

```
set obs 20
gen y = rnormal(5,2)

program logpost
    args logp b
    local theta = 'b'[1,1]
    scalar 'logp' = 0
    logdensity normal 'logp' y 'theta' 2
    logdensity normal 'logp' 'theta' 4 1
end

matrix b = 4
mcmcrun logpost b using temp.csv, replace ///
    sampler(mhsnorm , sd(0.25)) par(theta) burn(100) update(5000)
insheet using temp.csv, comma clear
mcmctrace theta
```

mcmctrace plots the chain

## MCMC

Neonatal  
Mortality

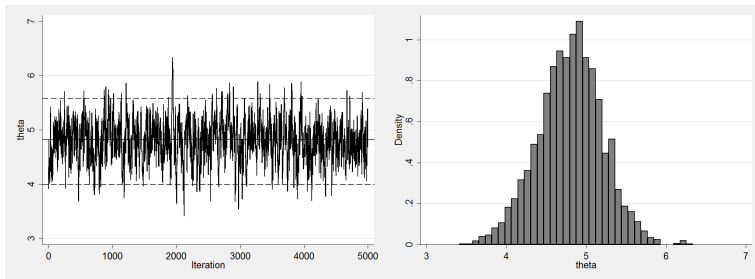
2005 data in  
Stata

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Conclusions



## Robust Model

```
set obs 20
gen y = rnormal(5,2)
qui replace y = 12 in 10
program logpost
    args logp b
    local theta = 'b'[1,1]
    scalar 'logp' = 0
    logdensity t 'logp' y 'theta' 2 4
    logdensity normal 'logp' 'theta' 4 1
end
matrix b = 4
mcmcrun logpost b using temp.csv, replace ///
sampler(mhsnorm , sd(0.25)) par(theta) burn(100) update(5000)
insheet using temp.csv, comma clear
mcmctrace theta
```



## Programs introduced in the book

### MCMC

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Conclusions

Family	Example	Purpose
	logdensity	Calculate log-likelihoods & log-priors
mhs	mhsnorm	Various Metropolis-Hastings samplers
mcmc	mcmctrace	Run or inspect an MCMC analysis
gbs	gbsslice	Gibbs samplers
wbs	wbsrun	Communication with WinBUGS

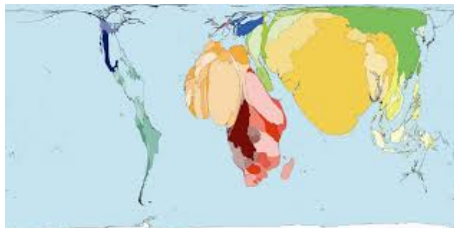
# Neonatal Mortality in the UK

Mortality during the first 28 days of life following a live birth

Often divided into

Early Neonatal (0-7 days): pregnancy

Late Neonatal (8-27days): environmental



# Why do UK & USA do poorly?

MCMC

Neonatal Mortality

2005 data in Stata

2005 data in Mata

2005 data in WinBUGS

1999-2009 data

Conclusions

BMJ

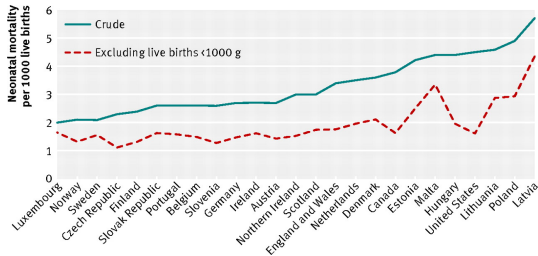
BMJ 2012;344:e746. doi: 10.1136/bmj.e746 Published 17 February 2012

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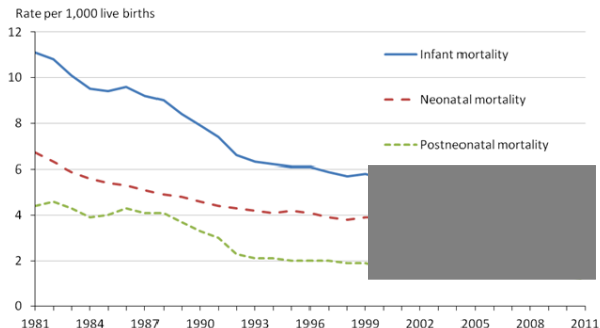
## Influence of definition based versus pragmatic birth registration on international comparisons of perinatal and infant mortality: population based retrospective study

BMJ OPEN ACCESS

K S Joseph professor<sup>1</sup>, Shiliang Liu research scientist<sup>2</sup>, Jocelyn Rouleau data manager<sup>2</sup>, Sarka Lisronkova postdoctoral fellow<sup>1</sup>, Jennifer A Hutcheon postdoctoral fellow<sup>1</sup>, Reg Saave professor<sup>2</sup>, Alexander C Allen professor<sup>3</sup>, Michael S Kramer professor<sup>3</sup>, for the Fetal and Infant Health Study Group of the Canadian Perinatal Surveillance System



# UK Neonatal Mortality 1999-2010



<http://www.ons.gov.uk/ons/rel/vsob1/child-mortality-statistics-childhood-infant-and-perinatal/2011/sty-infant-mortality.html>

# Prior knowledge

## FETAL AND NEONATAL MEDICINE Richard E. Behrman, *Editor*

---

### *Neonatal mortality risk in relation to birth weight and gestational age: Update*

**Beverly L. Koops, M.D., Linda J. Morgan, M.D., and  
Frederick C. Battaglia, M.D., *Denver, Colo., and Hanover, N.H.***

# Prior knowledge

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Frederick C. Battaglia, M.D., *Denver, Colo., and***

## The New England Journal of Medicine

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ASSOCIATION OF YOUNG MATERNAL AGE WITH ADVERSE REPRODUCTIVE OUTCOMES

ALISON M. FRASER, M.S.P.H., JOHN E. BROCKERT, M.P.H., AND R.H. WARD, Ph.D.

# Prior knowledge

## FETAL AND NEONATAL MEDICINE Richard E. Behrman, *Editor*

---

*Neonatal mortality risk in relation to birth  
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**Beverly L. Koops, M.D., Linda J. Morgan, M.D.  
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Number 15

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1995, 17

Number 17

**WITH ADVERSE REPRODUCTIVE OUTCOMES  
BROCKERT, M.P.H., AND R.H. WARD, Ph.D.**

### INCREASED MATERNAL AGE AND THE RISK OF FETAL DEATH

RUTH C. FRETTS, M.D., M.P.H., JULIE SCHMITTDIEL, M.A., FRANCES H. McLEAN, B.Sc.N.,  
ROBERT H. USHER, M.D., AND MARLENE B. GOLDMAN, Sc.D.

deaths  $\sim$  Poisson(Rate per 1000  $\times$  Births/1000)

$$\log(\text{Rate}) = \mu + \alpha[\text{age}] + \beta[\text{bwt}]$$

$$\text{Constraint: } \alpha[20-24] = \beta[3000-3499] = 0$$

$\mu$  represents the log rate in the baseline group

$\alpha$  represents the log relative rate (baseline:20-24 years)

$\beta$  represents the log relative rate (baseline: 3000-3499 grams)



deaths  $\sim$  Poisson(Rate per 1000  $\times$  Births/1000)

$$\log(\text{Rate}) = \mu + \alpha[\text{age}] + \beta[\text{bwt}]$$

$$\text{Constraint: } \alpha[20-24] = \beta[3000-3499] = 0$$

$\mu$  represents the log rate in the baseline group

$\alpha$  represents the log relative rate (baseline:20-24 years)

$\beta$  represents the log relative rate (baseline: 3000-3499 grams)

### ML analysis in Stata

```
. glm deaths ib2.age ib6.bwt , fam(poi) off(lnBirths)
```

Maternal Age	Relative Rate	$\alpha$	Birth Weight	Relative Rate	$\beta$
<20	2 (1,4)	0.69 (0.34)	<1000	100(10,1000)	4.61 (1.15)
20-24	1	0.0	1000-1499	10(2,50)	2.30 (0.80)
25-29	1(0.5,2)	0.00 (0.34)	1500-1999	3(1,9)	1.10 (0.55)
30-34	1(0.5,2)	0.00 (0.34)	2000-2499	1(0.5,2)	0.00 (0.34)
35-39	1(0.5,2)	0.00 (0.34)	2500-2999	1(0.5,2)	0.00 (0.34)
40+	2(1,4)	0.69 (0.34)	3000-3499	1	0.0
			3500-3999	1(0.5,2)	0.00 (0.34)
			4000+	2(1,4)	0.69 (0.34)

Mortality in baseline category

0.25 per 1,000 births (0.08,0.75)      $\mu$  -1.39(0.55)

## log-posterior(slow and inefficient)

```
program logpost
  args logp b
  tempvar lnMU j MU
  gen 'lnMU' = lnBirths + 'b'[1,1]
  replace 'lnMU' = 'lnMU' + 'b'[1,b] if bwt != 6
  replace 'lnMU' = 'lnMU' + 'b'[1,a] if age != 2
  gen 'MU' = exp('lnMU')
  scalar 'logp' = 0
  logdensity poisson 'logp' deaths 'MU'
  logdensity normal 'logp' 'b'[1,1] -1.39 0.55
  logdensity normal 'logp' 'b'[1,2] 4.61 1.15
  logdensity normal 'logp' 'b'[1,3] 2.30 0.80
  logdensity normal 'logp' 'b'[1,4] 1.10 0.55
  logdensity normal 'logp' 'b'[1,5] 0 0.34
  logdensity normal 'logp' 'b'[1,6] 0 0.34
  logdensity normal 'logp' 'b'[1,7] 0 0.34
  logdensity normal 'logp' 'b'[1,8] 0.69 0.34
  logdensity normal 'logp' 'b'[1,9] 0.69 0.34
  logdensity normal 'logp' 'b'[1,10] 0 0.34
  logdensity normal 'logp' 'b'[1,11] 0 0.34
  logdensity normal 'logp' 'b'[1,12] 0 0.34
  logdensity normal 'logp' 'b'[1,13] 0.69 0.34
end
```

# Convergence: Trace Plots

MCMC

Neonatal  
Mortality

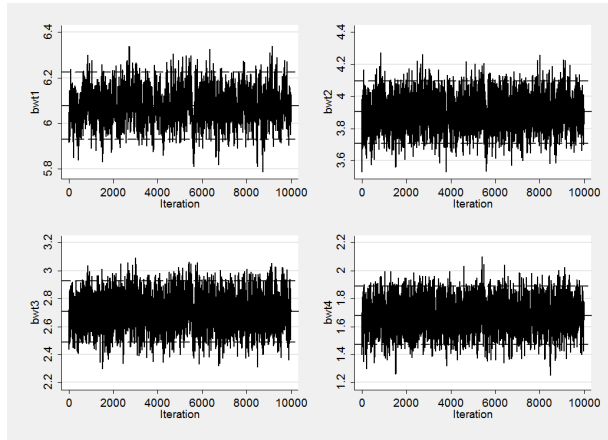
2005 data in  
Stata

2005 data in  
Mata

2005 data in  
WinBUGS

1999-2009  
data

Conclusions



# Convergence: Section Plots

MCMC

Neonatal  
Mortality

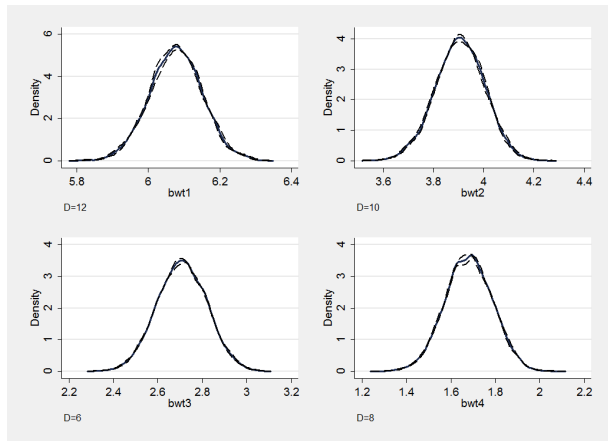
2005 data in  
Stata

2005 data in  
Mata

2005 data in  
WinBUGS

1999-2009  
data

Conclusions



# Parameter Estimates

MCMC

Neonatal Mortality

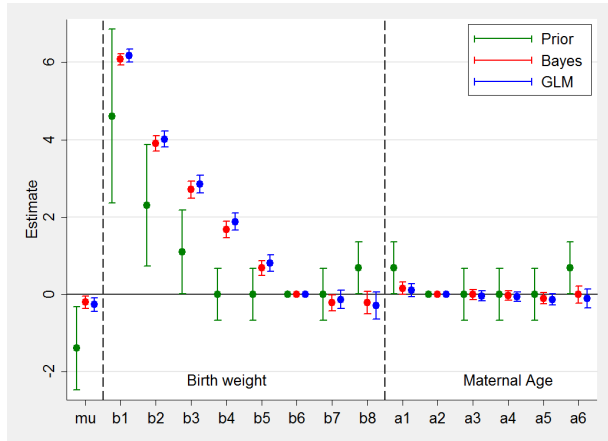
2005 data in Stata

2005 data in Mata

2005 data in WinBUGS

1999-2009 data

Conclusions



## Prediction

Predictive distribution of new values  $y^*$  given previous values  $y$

$$p(y^*|y) = \int p(y^*|\theta)p(\theta|y)d\theta$$

approximate the integral in an MCMC algorithm

simulate new data  $y^*$  from  $p(y^*|\theta)$  using the current  $\theta$

```

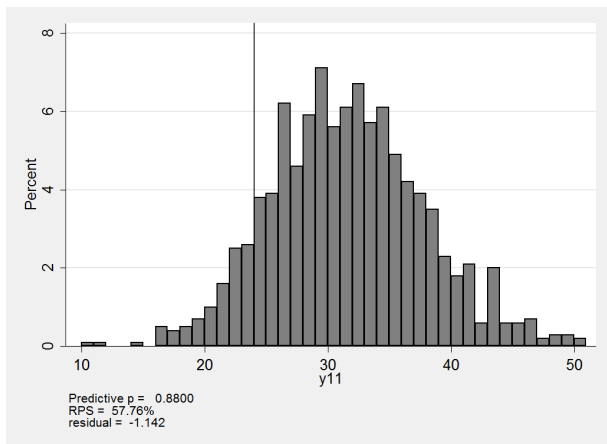
program y , rclass
    args b

    tempvar lnMU j MU y
    tempname pd
    gen 'lnMU' = lnBirths + 'b'[1,1]
    qui replace 'lnMU' = 'lnMU' + 'b'[1,b] if bwt != 6
    qui replace 'lnMU' = 'lnMU' + 'b'[1,a] if age != 2
    gen 'MU' = exp('lnMU')
    gen 'y' = rpoisson('MU')
    mkmat 'y' , matrix('pd')
    matrix 'pd' = 'pd'
    return matrix pred = 'pd'
end

mcmcrun logpost b using temp.csv, replace ///
    samplers( 13(mhnorm , sd(0.1)) ) burn(1000) updates(20000) ///
    thin(20) par(mu bwt1-bwt5 bwt7 bwt8 age1 age3-age6) pred(y)
  
```

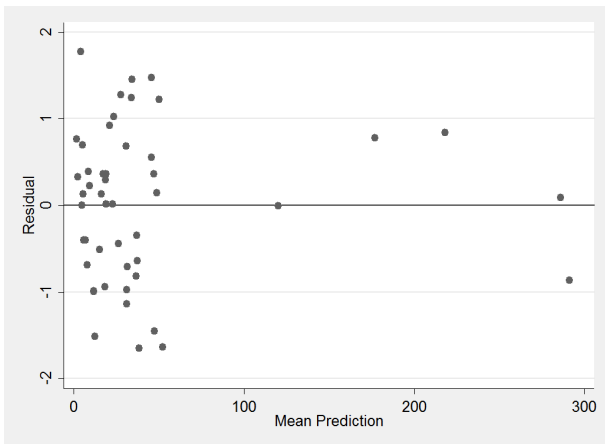


## Predictive Distribution of Observation 11



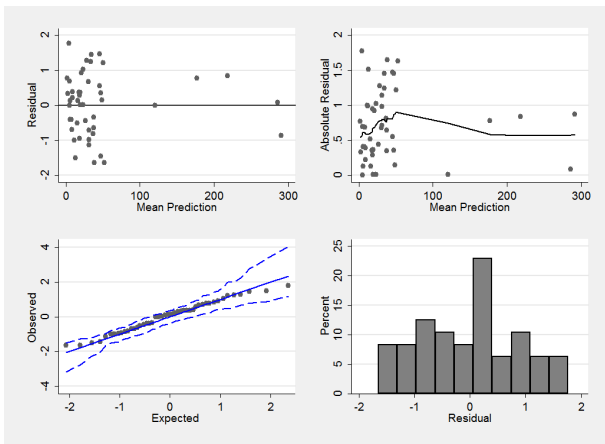
```
mcmccheck , d(deaths[11]) df(paed2005.dta) p(y11) pf(mcmc2005.dta)  
gopt(width(1))
```

# Residuals vs Fit



```
mcmccheck , d(deaths) df(paed2005.dta) p(y) pf(mcmc2005.dta)
```

# Summary Plot



```
mcmccheck , d(deaths) df(paed2005.dta) p(y) pf(mcmc2005.dta) plot(summary)
```

```

set matastrict on
mata:
mata clear
real scalar logpost(real matrix X,real rowvector b,real scalar ipar)
{
    real colvector lnM
    real scalar i

    lnM = X[,2] : + b[1]
    for(i=1;i<=48;i++) {
        if( X[i,5] != 2 ) lnM[i] = lnM[i] + b[X[i,3]]
        if( X[i,6] != 6 ) lnM[i] = lnM[i] + b[X[i,4]]
    }
    return( sum(X[,1]:*lnM :- exp(lnM)) -1.652893*(b[1]-1.39)*(b[1]-1.39)
    -0.378072*(b[2]-4.61)*(b[2]-4.61)-0.78125*(b[3]-2.3)*(b[3]-2.3)
    -1.652893*(b[4]-1.1)*(b[4]-1.1)-4.32526*(b[5]*b[5]+b[6]*b[6]+b[7]*b[7]
    +(b[8]-0.69)*(b[8]-0.69)+(b[9]-0.69)*(b[9]-0.69)+b[10]*b[10]+b[11]*b[11]
    +b[12]*b[12]+(b[13]-0.69)*(b[13]-0.69)) )
}
end

```

```

use paediatric.dta, clear
keep if year == 2005
drop if bwt == 9
gen Rate = 1000*deaths/births
gen lnBirths = log(births/1000)
gen b = bwt + (bwt<6)
gen a = 7 + age + (age<2)
matrix theta = (-1.5,4.5,2.5,1,0,0,0,1,1,0,0,0,1)
matrix s = J(1,13,0.1)
mcmcrun logpost X theta using temp1.csv, samp( 13(mhnorm , sd(s)) ) ///
    burn(1000) adapt update(50000) thin(5) jpost dots(0) ///
    par(mu bwt1-bwt5 bwt7 bwt8 age1 age3-age6) replace ///
    data(X=(deaths lnBirths a b age bwt) theta s) mata
  
```

## WinBUGS Analysis

- Write text files containing:
  - The model file (c.f. logpost)
  - The data in WinBUGS format
  - The initial values
  - The script (batch commands to control the fit)
- Run the script file
  - Results stored in a text file
- Read results into Stata
- Process the results

## WinBUGS model file

```

model {
  for( i in 1:6 ) {
    for( j in 1:8 ) {
      log(m[i,j]) <- mu + alpha[i] + beta[j] + LB[i,j]
      D[i,j] ~ dpois(m[i,j])
    }
  }
  mu ~ dnorm(-1.39,3.306)
  alpha[1] ~ dnorm(0.69,8.651)
  alpha[2] <- 0
  alpha[3] ~ dnorm(0.0,8.651)
  alpha[4] ~ dnorm(0.0,8.651)
  alpha[5] ~ dnorm(0.0,8.651)
  alpha[6] ~ dnorm(0.69,8.651)
  beta[1] ~ dnorm(4.61,0.756)
  beta[2] ~ dnorm(2.30,1.563)
  beta[3] ~ dnorm(1.10,3.306)
  beta[4] ~ dnorm(0.0,8.651)
  beta[5] ~ dnorm(0.0,8.651)
  beta[6] <- 0
  beta[7] ~ dnorm(0.0,8.651)
  beta[8] ~ dnorm(0.69,8.651)
}

```

## WinBUGS data file

MCMC

Neonatal  
Mortality

2005 data in  
Stata

2005 data in  
Mata

2005 data in  
WinBUGS

1999-2009  
data

Conclusions

Store data logBirths and deaths in Stata matrices LB and D

```
. wbslist (matrix LB D) using data.txt , replace
```

```
list( LB=structure(.Data=c(
-1.248, -0.931, -0.224, 0.901, 2.218, 2.840, 2.420, 1.152,
-0.496, -0.076, 0.646, 1.840, 3.173, 3.812, 3.468, 2.370,
-0.197, 0.153, 0.914, 2.043, 3.343, 4.081, 3.840, 2.866,
-0.200, 0.240, 0.967, 2.090, 3.366, 4.186, 4.037, 3.151,
-0.601, -0.172, 0.538, 1.530, 2.750, 3.565, 3.449, 2.610,
-2.017, -1.483, -0.863, 0.143, 1.273, 2.016, 1.839, 1.055),.Dim = c(6,8)),
D=structure(.Data=c(120,20,8,7,19,17,10,3,
234,33,29,35,27,35,26,6,
272,50,35,42,57,50,25,8,
288,60,27,44,50,39,31,13,
190,24,21,14,23,24,19,6,
35,10,7,5,6,5,8,3),.Dim = c(6,8)))
```



# WinBUGS initial values file

MCMC

Neonatal  
Mortality

2005 data in  
Stata

2005 data in  
Mata

2005 data in  
WinBUGS

1999-2009  
data

Conclusions

```
. wbslist (mu=-1.4,alpha=c(0.7,NA,0,0,0,0.7), ///  
          beta=c(4.6,2.3,1.1,0,0,NA,0,0.7)) ///  
          using init.txt , replace
```

```
list( mu=-1.4,alpha=c(0.7,NA,0,0,0,0.7),  
      beta=c(4.6,2.3,1.1,0,0,NA,0,0.7))
```

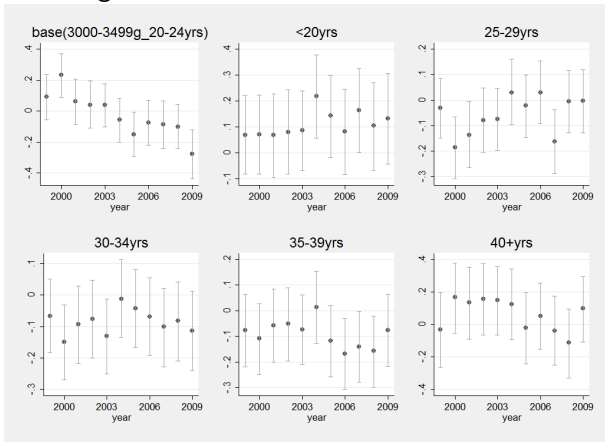
# WinBUGS script file

```
. wbsscript using script.txt, model(model.txt) ///  
  data(data.txt) init(init.txt) burn(1000) update(10000) ///  
  set(mu alpha beta) coda(NMR) log(NMR.log) replace
```

```
display('log')  
check('E:/StataUsers/Data/model.txt')  
data('E:/StataUsers/Data/data.txt')  
compile(1)  
inits(1,'E:/StataUsers/Data/init.txt')  
gen.inits()  
refresh(100000)  
update(1000)  
set('mu')  
set('alpha')  
set('beta')  
update(10000)  
coda(*,'E:/StataUsers/Data/NMR')  
save('E:/StataUsers/Data/NMR.log')
```

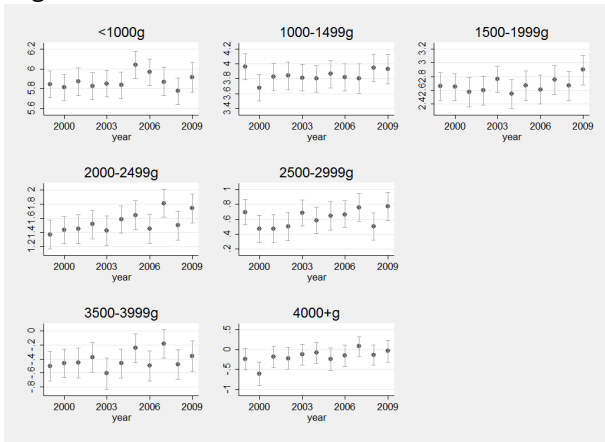
# Analyses for 1999-2009

## Baseline and age effects



# Analyses for 1999-2009

## Birth weight effects

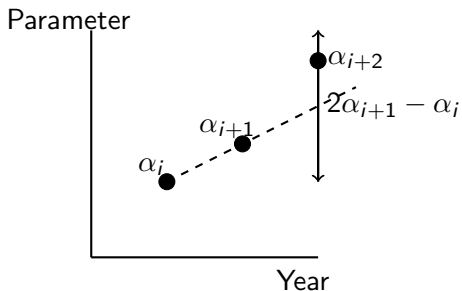


## Smoothing the parameters

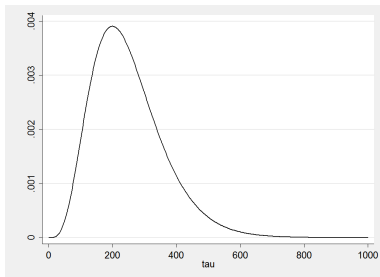
The parameter estimates show improbable jumps between years and would be more credible if the trends were smoother.

$$\alpha_{i+2} \sim N(2\alpha_{i+1} - \alpha_i, \tau)$$

$\tau$  acts as a smoothing parameter

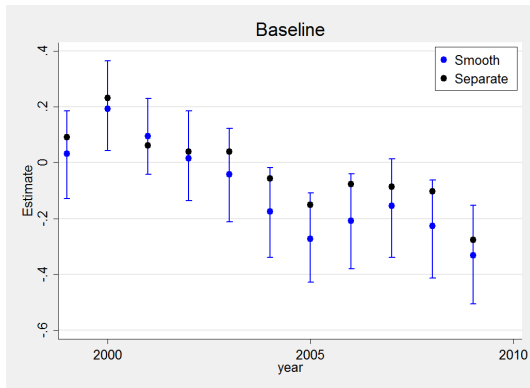


## Prior for tau



Prior  $G(5,50)$ , prior mean = 250, prior sd = 112  
 $\tau=250 \Rightarrow \text{sd}=0.063$

# Smoothed Estimates



# Stronger smoothing

MCMC

Neonatal Mortality

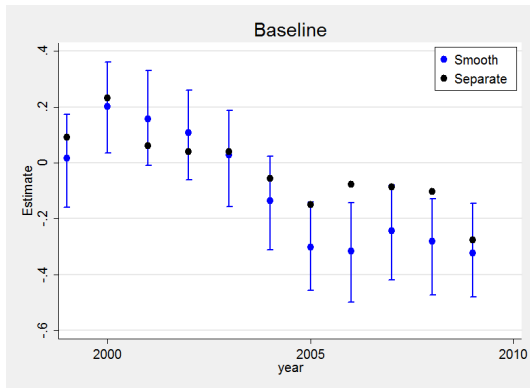
2005 data in Stata

2005 data in Mata

2005 data in WinBUGS

1999-2009 data

Conclusions



Prior  $G(50,5)$ , prior mean 250, prior  $sd=35$



## Average predictions for 2010

mean prediction (actual number)

BWT(g)	AGE(yrs)					
	< 20	20–	25–	30–	35–	40+
< 1000	90(72)	217(205)	317(257)	283(278)	210(173)	84(49)
1000–	13(17)	32(21)	35(46)	43(31)	33(26)	12(11)
1500–	11(4)	30(25)	42(28)	40(34)	29(17)	12(8)
2000–	12(11)	33(24)	45(37)	37(34)	27(22)	10(8)
2500–	17(10)	49(38)	65(51)	53(44)	34(27)	11(8)
3000–	12(10)	35(37)	50(48)	42(49)	26(17)	8(5)
3500–	6(3)	19(21)	30(13)	27(19)	17(15)	5(5)
4000+	2(0)	8(6)	15(13)	14(12)	9(10)	3(2)

Slightly better predictions if  $\tau$  is allowed to differ with the parameter

MCMC

Neonatal  
Mortality

2005 data in  
Stata

2005 data in  
Mata

2005 data in  
WinBUGS

1999-2009  
data

Conclusions

- The Bayesian approach has many advantages
- MCMC is straightforward but can be slow
- MCMC allows flexibility in modelling (cf ml)
- MCMC is practical in Stata for small problems
- Larger problems require Mata or WinBUGS
- Other issues covered in the book include:
  - Convergence checking
  - Gibbs sampling
  - Model comparison & tests
  - Validation of software
  - Writing Bayesian programs for general use

## Recommendations

- Stata should provide facilities for communication with other software, e.g.
  - Stata  $\longleftrightarrow$  WinBUGS
  - Stata  $\longleftrightarrow$  R
- Stata needs to be able to handle datasets that are too large for the Editor
- Bayesian analysis should be fully integrated into Stata
- It would be possible to re-write WinBUGS/JAGS in Mata
- This is a job for StataCorp