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— in addition to overheads from lectures 8–L and 11–L

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| TRANSITION MODELS AS MARKOV MODELS |
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### Markov models:

- Markov property: “the future depends on the past only through the present”,
- conditional — by successive conditioning on the past, e.g.,  
$$P(Y_1, Y_2, Y_3, Y_4) = P(Y_1) \cdot P(Y_2|Y_1) \cdot P(Y_3|Y_1, Y_2) \cdot P(Y_4|Y_1, Y_2, Y_3),$$
- components of model:
  - \* initial distribution(s) — unconditional,
  - \* conditional distribution given past,
- order of model: number of time lags involved; simplest is first order where only previous observation involved,
- in general, no simple link between conditional and marginal models  $\Rightarrow$  parameters have different interpretations.

## TRANSITIONAL LOGISTIC REGRESSION

### Analysis of transitional logistic regression models:

- use previous state(s) (depending on order) as predictors along with other predictors,
- omit time point(s), where no previous state(s) available, or include them by *(i)* setting previous state(s)  $\equiv 0$ , and *(ii)* including indicator variable for obs. with no previous state(s) and its interactions,
- statistical analysis:
  - \* 2-level structure: ordinary logistic regression with robust variance estimation (clustered on subjects)<sup>1</sup>  $\sim$  GEE with independence working correlation,
  - \* other hierarchical structures: marginal or random effects logistic regression.
- recommended to check sensitivity of results to order of model.

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<sup>1</sup> Robust variance estimates gives consistent inference when (only) the conditional mean is correctly specified, and often also when the Markov property is violated.

## INDONESIAN CHILDREN STUDY

Notes on models fitted in Diggle et al. (2002):

- first and second order transitional models,
- observations without previous states omitted,
- results for first order models:
  - \* previous state: a weak predictor,
  - \* estimates for Xerophthalmia only little affected by previous state,
  - \* no evidence of interaction with previous state, for Xerophthalmia or other predictors,
  - \* robust and ordinary SE's similar  $\Rightarrow$  Markov assumption seems approximately ok,
- results for second order models:  
(not shown in tables, and I could not reproduce them from the dataset)
  - \* substantial increase in coefficient for Xerophthalmia,
  - \* reduced influence of season,— shows importance of assessing the robustness of results to model order.

## TRANSITIONAL POISSON REGRESSION

3 different models/approaches (for Poisson regression with log link):

- use  $Y_{t-1}$  as a predictor:

$$\log(\mu_{ti}) = (X'\beta)_{ti} + \gamma Y_{t-1,i},$$

— does not work well for  $\gamma > 0$  (exponential growth over time, rarely a realistic model), so primarily a model for negative correlation,

- use mean-corrected  $Y_{t-1}$  as a predictor:

$$\log(\mu_{ti}) = (X'\beta)_{ti} + \gamma (\log(Y_{t-1,i}^*) - (X'\beta)_{t-1,i}),$$

where  $Y_{t-1,i}^* = \max(d, Y_{t-1,i})$  for some  $0 < d < 1$  (to avoid problems with zero counts);

- \*  $\gamma > 0 \sim$  positive correlation is possible and does not lead to non-stationarity,
- \* model cannot be fit with standard GLM software,
- autocorrelated errors  $\sim$  transitional model but usually considered as a GLM with autocorrelation/time series errors (Davis et al., Biometrika **87**:491–505),
- transitional regression models (Brumback et al., 2000) incorporate all above models (autocorrelated errors and transitional GLMs), and are fitted by generalized least squares (S code available from Statlib).