



A mixed autoregressive probit model for ordinal longitudinal data

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SUMMARY

Longitudinal data with binary and ordinal outcomes routinely appear in medical applications. Existing methods are typically designed to deal with short measurement series. In contrast, modern longitudinal data can result in large numbers of subject-specific serial observations. In this framework, we consider multivariate probit models with random effects to capture heterogeneity and autoregressive terms for describing the serial dependence. Since likelihood inference for the proposed class of models is computationally burdensome because of high-dimensional intractable integrals, a pseudolikelihood approach is followed. The methodology is motivated by the analysis of a large longitudinal study on the determinants of migraine severity.

Keywords: Autoregressive errors; Composite likelihood; Longitudinal data; Migraine severity; Mixed models; Ordinal probit; Pairwise likelihood.

1. INTRODUCTION

Pain severity is often measured on rating scales that involve 4–11 categories ranging from the absence of symptoms to the most severe pain (e.g. Von Korff *and others*, 2000). For chronic and recurrent pain conditions, such as migraine and back pain, studying the symptom severity over a time period is crucial to detect common- and person-specific pain trigger conditions. To this aim, patients record the pain severity in a diary over some time period. See Bolger *and others* (2003) for general design, technology, and analysis questions. With the availability of electronic data collection methods such as palmtop computers, the frequency of such assessments can be very high. Thus, it is important to develop statistical methods that are able to deal adequately with large longitudinal ordinal response data in cross-sectional setups.

There exist several methods to deal with short longitudinal setups involving ordinal responses measured typically over 4–7 time points. Many of them require the inclusion of random effects to deal with

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the dependence between subject-specific measurements (e.g. Hedeker and Gibbons, 1994; Gibbons and Hedeker, 1997; Liu and Hedeker, 2006; Todem *and others*, 2007). Estimation in such models are often based on Gauss–Hermite quadrature for the integration of random effects. Another proposal involves the global odds ratio suggested by Dale (1986) (see Molenberghs and Lesaffre, 1994; Williamson *and others*, 1995). Still another approach is based on Markov transition models. Lee and Daniels (2007) extend this method from binary (Heagerty, 2002) to ordinal longitudinal data involving 6 time points. Böckenholt (1999) uses a first-order Markov process on the category indicators to capture the time dependence. His model is able to fit longer ordinal time series but requires that all time points are equidistant and common to all units.

For studying binary time series, Piorecky *and others* (1996) use generalized estimating equations (Liang and Zeger, 1986) to adjust for the dependency between measurements. Generalized estimating equations could also be used for ordinal-valued time series if one is only interested in inference for regression parameters (see, e.g. Liang *and others*, 1992; Lipsitz and Kim, 1994; Heagerty and Zeger, 1996; Fahrmeir and Pritscher, 1996; Delfino *and others*, 2001).

All the above approaches are limited by the number of person-specific measurements or by other restrictions such as common equidistant time points. Motivated by a longitudinal study on migraine severity determinants, we propose a class of mixed ordered probit models with an autocorrelated component to capture subject-specific time series variability. In Section 2, we describe the model class. In Section 3, we develop a computationally convenient composite likelihood approach for inference and model selection. Section 4 illustrates the application to the migraine pain severity data. The paper closes with some final remarks.

2. MIXED AUTOREGRESSIVE ORDINAL PROBIT MODELS

Let Y_{ij} represent a categorical response with h possible ordered categories and let \mathbf{x}_{ij} be a vector of p exploratory variables observed at time t_{ij} for observation $j = 1, \dots, n_i$ on subject $i = 1, \dots, m$. As usual in longitudinal studies, the m subjects are assumed to be independent. The ordinal response Y_{ij} may be viewed as a censored observation from a hidden continuous variable Y_{ij}^* ,

$$Y_{ij} = y_{ij} \Leftrightarrow \alpha_{y_{ij}-1} < Y_{ij}^* \leq \alpha_{y_{ij}}, \quad y_{ij} \in \{1, \dots, h\},$$

where $-\infty \equiv \alpha_0 < \alpha_1 < \dots < \alpha_{h-1} < \alpha_h \equiv \infty$ are suitable threshold parameters. The important case of binary response corresponds to $h = 2$ and a single threshold parameter α_1 . Among several possible specifications for the relationship between the unobserved Y_{ij}^* and the vector of regressors \mathbf{x}_{ij} , a common choice is a linear mixed model of type

$$Y_{ij}^* = \mathbf{x}_{ij}^T \boldsymbol{\beta} + U_i + \epsilon_{ij}, \quad (2.1)$$

where $\boldsymbol{\beta}$ is a vector of p unknown coefficients, also termed fixed effects, while the U_i are m mutually independent random effects describing the heterogeneity among different subjects and the ϵ_{ij} are underlying errors. Popular assumptions for the marginal distribution of ϵ_{ij} are logistic and normal distributions, leading to the cumulative logit and cumulative probit models for the observed Y_{ij} , respectively. Additionally, we assume independence between ϵ_{ij} and U_i . For more details, see Agresti (2002, section 7). Here, we choose a probit model and assume that the random effects are normally distributed, $U_i \stackrel{\text{i.i.d.}}{\sim} \mathcal{N}(0, \sigma^2)$. We consider these distributional assumptions for ease of interpretation and mathematical manageability, although the methodology discussed in this paper holds more generally.

Model identifiability for the resulting multivariate probit model requires restrictions on both the location and the scale of the unobserved process Y_{ij}^* . These requirements are met when the errors ϵ_{ij} have

unit variance, and the first cut point α_1 or, alternatively, the intercept β_1 is fixed to zero (see, e.g. Chib and Greenberg, 1998).

Probit models with random effects have a particularly convenient interpretation. In fact, it is straightforward to move from a subject-specific interpretation to a population-level interpretation. For example, consider the probability that subject i experiences a certain level y_{ij} at time t_{ij}

$$\text{pr}(Y_{ij} = y_{ij}; \boldsymbol{\theta}) = \text{pr}(Y_{ij}^* \in (\alpha_{y_{ij}-1}, \alpha_{y_{ij}}]; \boldsymbol{\theta}) = \Phi\left(\frac{\alpha_{y_{ij}} - \mathbf{x}_{ij}^T \boldsymbol{\beta}}{\sqrt{\sigma^2 + 1}}\right) - \Phi\left(\frac{\alpha_{y_{ij}-1} - \mathbf{x}_{ij}^T \boldsymbol{\beta}}{\sqrt{\sigma^2 + 1}}\right), \quad (2.2)$$

where $\Phi(z)$ denotes the cumulative probability function of a standard normal variable and $\boldsymbol{\theta}$ is the parameter vector, including the cut points $\boldsymbol{\alpha} = (\alpha_2, \dots, \alpha_h)^T$, the regressor coefficients $\boldsymbol{\beta}$, and the variance component σ^2 . While the subject-specific effect of the covariates on the response is described by $\boldsymbol{\beta}$, from expression (2.2), it follows that the average population effect is governed by the rescaled coefficient $\boldsymbol{\beta}^{\text{pop}} = \boldsymbol{\beta}/\sqrt{\sigma^2 + 1}$.

Commonly, probit models with random effects are constructed by assuming that the underlying errors ϵ_{ij} are mutually independent, $\epsilon_{ij} \stackrel{\text{i.i.d.}}{\sim} \mathcal{N}(0, 1)$. It follows that the joint distribution of the hidden variables for the i th subject $(Y_{i1}^*, \dots, Y_{in_i}^*)^T$ is multivariate normal with standardized mean vector

$$\left(\frac{\mathbf{x}_{i1}^T \boldsymbol{\beta}}{\sqrt{\sigma^2 + 1}}, \dots, \frac{\mathbf{x}_{in_i}^T \boldsymbol{\beta}}{\sqrt{\sigma^2 + 1}}\right)^T \quad (2.3)$$

and correlation matrix with constant nondiagonal entries given by $\sigma^2/(\sigma^2 + 1)$.

Model fitting is typically performed by maximum likelihood. Denote by $\mathbf{y} = (\mathbf{y}_1^T, \dots, \mathbf{y}_m^T)^T$ the vector of all observations, with $\mathbf{y}_i = (y_{i1}, \dots, y_{in_i})^T$ being the subvector of observations pertaining to the i th patient. Similarly denote the vectors of the corresponding hidden variables \mathbf{Y}^* and \mathbf{Y}_i^* , respectively. The likelihood function for the usual probit model with underlying independent errors involves m intractable integrals

$$\begin{aligned} \mathcal{L}(\boldsymbol{\theta}; \mathbf{y}) &= \prod_{i=1}^m \int_{-\infty}^{\infty} \prod_{j=1}^{n_i} \text{pr}(Y_{ij} = y_{ij} | \mathbf{x}_{ij}, u_i; \boldsymbol{\theta}) f(u_i; \boldsymbol{\theta}) du_i \\ &= \prod_{i=1}^m \int_{-\infty}^{\infty} \prod_{j=1}^{n_i} \left(\int_{\alpha_{y_{ij}-1}}^{\alpha_{y_{ij}}} f(y_{ij}^* | \mathbf{x}_{ij}, u_i; \boldsymbol{\theta}) dy_{ij}^* \right) f(u_i; \boldsymbol{\theta}) du_i \\ &= \prod_{i=1}^m \int_{-\infty}^{\infty} \prod_{j=1}^{n_i} (\Phi(\alpha_{y_{ij}} - \mathbf{x}_{ij}^T \boldsymbol{\beta} - u_i) - \Phi(\alpha_{y_{ij}-1} - \mathbf{x}_{ij}^T \boldsymbol{\beta} - u_i)) \phi\left(\frac{u_i}{\sigma}\right) du_i, \quad (2.4) \end{aligned}$$

where $\phi(z)$ denotes the probability density function of a standard normal variable. This likelihood may be approximated by Gauss–Hermite quadrature or, more accurately, by adaptive Gauss–Hermite quadrature (see, e.g. Pinheiro and Bates, 2000).

Although the above described probit mixed model is widely used, its underlying equal correlation assumption seems unsatisfactory for many longitudinal studies, especially for those characterized by moderate to long subject-specific series. Better models should take into account the serial correlation within each subject-specific time series. In this paper, we propose to model the within-subject serial correlation by a smooth temporal decaying correlation function as, for example the exponential correlation model (Diggle and others, 2002), $\text{corr}(\epsilon_{ij}, \epsilon_{ik}) = \exp(-\delta|t_{ij} - t_{ik}|)$, where t_{ij} denotes the measurement time

of observation y_{ij} . This correlation function reduces to the autoregressive model of order 1, $\gamma^{|t_{ij}-t_{ik}|}$ with $\gamma = e^{-\delta}$, for equispaced observations times. Correspondingly, the correlation between 2 hidden continuous variables is formed by a constant subject-specific level plus a smooth serial component

$$\text{corr}(Y_{ij}^*, Y_{ik}^*) = \frac{\sigma^2}{\sigma^2 + 1} + \frac{e^{-\delta|t_{ij}-t_{ik}|}}{\sigma^2 + 1}. \quad (2.5)$$

Thus, by assuming serial correlation among the errors, we obtain a multivariate probit model with the same marginal interpretation as in (2.2) but with a more realistic longitudinal structure. Thereafter, the proposed class of models will be termed mixed autoregressive ordinal probit (MAOP) models.

Further, model flexibility may be obtained by allowing the parameter δ to depend on a subject-specific factor S_i with q different levels. Thus, the model may describe different memory effects in different groups of subjects. For example, in the migraine data discussed in Section 4, different pain memory effects can be postulated in patients taking medications or not or in patients with different headaches types.

The cost for the versatility of the MAOP model is paid in terms of computational difficulties. The likelihood function still involves m intractable integrals but with dimensions corresponding to the cluster sizes n_1, \dots, n_m . Denote always by θ the parameter vector that now contains also the autocorrelation parameters δ . The likelihood function for the model with serially correlated errors is

$$\mathcal{L}(\theta; \mathbf{y}) = \prod_{i=1}^m \int_{-\infty}^{\infty} \left(\int_{a_{y_{i1}-1}}^{a_{y_{i1}}} \cdots \int_{a_{y_{in_i}-1}}^{a_{y_{in_i}}} f(y_{i1}^*, \dots, y_{in_i}^* | \mathbf{x}_{i1}, \dots, \mathbf{x}_{in_i}, u_i; \theta) dy_{i1}^*, \dots, dy_{in_i}^* \right) f(u_i; \theta) du_i.$$

By using the assumptions of normality for both the random effects U_i and the hidden errors ϵ_{ij} , the likelihood may be rewritten as the product of m integrals of multivariate normal densities of dimensions n_1, \dots, n_m

$$\mathcal{L}(\theta; \mathbf{y}) = \prod_{i=1}^m \int_{\tilde{a}_{y_{i1}-1}}^{\tilde{a}_{y_{i1}}} \cdots \int_{\tilde{a}_{y_{in_i}-1}}^{\tilde{a}_{y_{in_i}}} \phi_{n_i}(z_{i1}, \dots, z_{in_i}; R_i) dz_{i1}, \dots, dz_{in_i}, \quad (2.6)$$

where $\tilde{a}_{y_{ij}}$ indicates the standardized cut point, $\tilde{a}_{y_{ij}} = (a_{y_{ij}} - \mathbf{x}_{ij}^T \boldsymbol{\beta}) / \sqrt{\sigma^2 + 1}$. The integrands $\phi_{n_i}(z_{i1}, \dots, z_{in_i}; R_i)$ are n_i -dimensional multivariate normal densities with zero means and correlation matrix R_i whose entries are given by expression (2.5). Except for longitudinal data with small numbers of observations per subject, the direct computation of likelihood (2.6) is time consuming and possibly numerically unstable.

MAOP models for discrete-time observations are categorized Gaussian linear state space models. The celebrated Kalman filter (Kalman, 1960) allows efficient iterative computation of the exact likelihood function in Gaussian linear state space models but cannot be applied to censored observations. Reliable approaches use several kinds of Monte Carlo approximations based typically on Kalman filter-type iterations (see, e.g. Durbin and Koopman, 2001). A Bayesian analysis of binary time series allowing for covariates using Markov chain Monte Carlo methods and the simulation smoother of De Jong and Shephard (1995) for block updates of the hidden process variables were developed in Czado and Song (2008). It would be feasible to extend their approach to ordinal-valued time series models using ideas of Müller and Czado (2005, 2009) to update the threshold parameters.

Unfortunately, these computer-intensive approaches may be difficult to apply in large longitudinal data sets, such as the migraine data analyzed in Section 4. Moreover, even if the computational cost would be tolerable, a full likelihood approach might be impractical due to the difficulty of assessing the adequacy of the multivariate assumptions underlying the model. These considerations lead us to consider a composite likelihood approach (Lindsay, 1988).

3. COMPOSITE LIKELIHOOD INFERENCE

The term composite likelihood denotes a rich class of pseudolikelihoods constructed by compounding valid likelihoods based on data subsets. Recent applications include genetics, spatial statistics, time series, and longitudinal data analysis (see Varin, 2008, for a recent review).

Here, we focus on the composite likelihood constructed combining likelihoods for pairs of observations, also called pairwise likelihood (Le Cessie and Van Houwelingen, 1994). Since pairs formed from closest observations are likely to be more informative, it is convenient to restrict to the pseudolikelihood constructed from the marginal probabilities of observed pairs of outcomes less distant than q units,

$$p\ell^{(q)}(\boldsymbol{\theta}; \mathbf{y}) = \sum_{i=1}^m \sum_{j < k}^{n_i} \log \text{pr}(Y_{ij} = y_{ij}, Y_{ik} = y_{ik}; \boldsymbol{\theta}) \mathbb{1}_{[-q, q]}(t_{ij} - t_{ik}),$$

where $\mathbb{1}_{\mathcal{A}}(x)$ is the indicator of the event $\{x: x \in \mathcal{A}\}$. Note that $p\ell^{(q)}(\cdot; \mathbf{y})$ is a weighted log-pairwise likelihood with dummy weights used to exclude pairs too far apart. A recent detailed discussion of weighted versions of pairwise likelihood can be found in Joe and Lee (2009).

In contrast to a full likelihood approach, the pairwise likelihood for MAOP models involves only 2D Gaussian integrals,

$$\text{pr}(Y_{ij} = y_{ij}, Y_{ik} = y_{ik}; \boldsymbol{\theta}) = \int_{\tilde{\alpha}_{y_{ij}-1}}^{\tilde{\alpha}_{y_{ij}}} \int_{\tilde{\alpha}_{y_{ik}-1}}^{\tilde{\alpha}_{y_{ik}}} \phi_2 \left(z_{ij}, z_{ik}; \frac{\sigma^2}{\sigma^2 + 1} + \frac{e^{-\delta_{w_i}|t_{ij}-t_{ik}|}}{\sigma^2 + 1} \right) dz_{ij} dz_{ik}.$$

The maximum composite likelihood estimator for $\boldsymbol{\theta}$ solves the composite likelihood score equation,

$$u^{(q)}(\boldsymbol{\theta}; \mathbf{y}) = \sum_{i=1}^m u_i^{(q)}(\boldsymbol{\theta}; \mathbf{y}_i) = \sum_{i=1}^m \sum_{j < k}^{n_i} u_{i,jk}(\boldsymbol{\theta}; \mathbf{y}_i) \mathbb{1}_{[-q, q]}(t_{ij} - t_{ik}),$$

where $u_{i,jk}(\boldsymbol{\theta}; \mathbf{y}_i) = \nabla \log \text{pr}(Y_{ij} = y_{ij}, Y_{ik} = y_{ik}; \boldsymbol{\theta})$. Since $u^{(q)}(\boldsymbol{\theta}; \mathbf{y})$ is a linear combination of proper score functions associated with each pairwise term forming the pseudolikelihood, it follows that, under standard assumptions (Molenberghs and Verbeke, 2005, section 9), the maximum pairwise likelihood estimator $\hat{\boldsymbol{\theta}}^{(q)}$ is consistent and asymptotically normally distributed. See also Cox and Reid (2004) for a discussion on situations in which consistency of maximum pairwise likelihood estimators may not hold, such as in long-memory temporal processes.

The asymptotic variance of $\hat{\boldsymbol{\theta}}^{(q)}$ assumes the typical ‘‘sandwich’’ form,

$$\Sigma^{(m)}(\boldsymbol{\theta}) = \mathbf{H}^{(q)}(\boldsymbol{\theta})^{-1} \mathbf{J}^{(q)}(\boldsymbol{\theta}) \mathbf{H}^{(q)}(\boldsymbol{\theta})^{-1},$$

where $\mathbf{H}^{(q)}(\boldsymbol{\theta}) = -\mathbb{E}\{\nabla u^{(q)}(\boldsymbol{\theta}; \mathbf{Y})\}$ and $\mathbf{J}^{(q)}(\boldsymbol{\theta}) = \text{cov}\{u^{(q)}(\boldsymbol{\theta}; \mathbf{Y})\}$. The inverse of $\Sigma^{(q)}(\boldsymbol{\theta})$ is also termed as Godambe information (Song, 2007, section 3). An empirical estimate of $\mathbf{H}^{(q)}(\boldsymbol{\theta})$ is $-\nabla u^{(q)}(\hat{\boldsymbol{\theta}}^{(q)}; \mathbf{y})$. Alternatively, exploiting the information identity for each pairwise term forming the pseudolikelihood, $\mathbf{H}^{(q)}(\boldsymbol{\theta})$ may be conveniently estimated by

$$\hat{\mathbf{H}}^{(q)}(\mathbf{y}) = \sum_{i=1}^m \sum_{j < k}^{n_i} u_{i,jk}(\hat{\boldsymbol{\theta}}^{(q)}; \mathbf{y}_i) u_{i,jk}(\hat{\boldsymbol{\theta}}^{(q)}; \mathbf{y}_i)^T \mathbb{1}_{[-q, q]}(t_{ij} - t_{ik}), \quad (3.1)$$

thus avoiding the need to derive Hessian matrices. The natural empirical estimate of $\mathbf{J}^{(q)}(\boldsymbol{\theta})$ is

$$\hat{\mathbf{J}}^{(q)}(\mathbf{y}) = \sum_{i=1}^m u_i^{(q)}(\hat{\boldsymbol{\theta}}^{(q)}; \mathbf{y}_i) u_i^{(q)}(\hat{\boldsymbol{\theta}}^{(q)}; \mathbf{y}_i)^T. \quad (3.2)$$

Matrices $\hat{H}^{(q)}(\mathbf{y})$ and $\hat{J}^{(q)}(\mathbf{y})$ are key ingredients for high-level inferential tasks such as hypothesis testing and model selection. The composite likelihood information criterion (CLIC) by Varin and Vidoni (2005) is a direct generalization of the Akaike (1973) criterion for model selection with composite likelihoods. The CLIC suggests to prefer models with smaller values of the quantity

$$\text{CLIC}^{(q)} = -2(p\ell^{(q)}(\hat{\boldsymbol{\theta}}; \mathbf{y}) - d^{(q)}(\mathbf{y})),$$

where $d^{(q)}(\mathbf{y})$ is an estimate of the effective number of parameters of the model. A consistent estimate of $d^{(q)}(\mathbf{y})$ is given by the trace of the matrix $\hat{\Sigma}^{(q)}(\mathbf{y})\hat{H}^{(q)}(\mathbf{y})$. This information criterion may be seen as a form of the Takeuchi (1976) information criterion for model selection with misspecified likelihoods, being the pairwise likelihood a misspecified likelihood under the working assumption of independent pairs.

Regarding the choice of the maximal admissible distance q between pairs used in the pairwise likelihood, previous work on pairwise likelihood for temporal and spatial processes suggests that the inclusion of too-distant pairs is not only computationally inefficient but may also not improve statistical efficiency (see Varin, 2008). Here, we propose to choose the tuning parameter q as the value minimizing a global fitting criterion, for example the generalized variance defined as the determinant of $\hat{\Sigma}^{(q)}(\mathbf{y})$.

The supplementary appendix (available at *Biostatistics* online) contains details on a simulation study carried out to evaluate the finite-sample performance of the proposed inferential methods. The results suggest that maximum pairwise likelihood estimators behave well for all the parameters even in case of strong serial correlation among the hidden variables. Computer code written in the R language (R Development Core Team, 2008) is also included in the supplementary material (available at *Biostatistics* online).

4. MIGRAINE SEVERITY DATA

Prince *and others* (2004) report that 45 million Americans seek medical attention for headaches yearly, at an estimated labor cost of \$13 billion. They show that only half of the migraine patients are affected by weather conditions. In contrast, some studies show little or no effect of weather conditions on migraine severity (see Cooke *and others*, 2000; Prince *and others*, 2004, for specific references). However, in these studies, only the frequency of headache occurrences, and the daily maximum or total score of an ordinal severity levels have been studied.

Current strategies for the analysis of pain severity data measured on an ordinal scale require aggregating over periods to achieve continuous average or total pain scores (e.g. Cooke *and others*, 2000; Prince *and others*, 2004; Goldstein *and others*, 2005; Raskin *and others*, 2005). Such an approach ignores effects occurring during the aggregation periods.

Here, we directly model the observed severity categories collected using a headache diary. In particular, we investigate the 4 daily ratings—recorded at morning, noon, afternoon, and bedtime—of the headache intensity of 133 Canadian (Toronto) patients in a study conducted by psychologist T. Kostecki-Dillon during the years 1993–1996. Records of the migraine severity were made on an ordinal scale with 6 categories described in Table 1.

In addition to a subject-specific questionnaire with personal and clinical information, weather conditions were recorded. They were collected from the meteorological station closest to the place where patients spends most of their time. The weather covariates include measurements related to sunshine, humidity, wind direction and speed, windchill, pressure, air quality, and many others.

Patients with a very large number of missing observations in subsequent measurements, or with less than 1 day of measurements, were omitted. The final data set comprises 119 patients with a total of 16 366 measurements, 1157 of which are missing. We assume an ignorable missing mechanism, and thus, we base inference on the pairwise likelihood formed by pairs of observed outcomes. The numbers of observations

Table 1. *Migraine data. Description of response categories with observed frequencies*

Intensity	Frequency	Condition	Description
0	9210	No headache	
1	2455	Mild headache	Aware of it only when attending to it
2	1685	Moderate headache	Could be ignored at times
3	1156	Painful headache	Continuously aware of it but able to start or continue daily activities as usual
4	526	Severe headache	Continuously aware of it, difficult to concentrate, and able to perform only undemanding tasks
5	177	Intense headache	Continuously aware of it, incapacitating unable to start or continue activity

Table 2. *Migraine data. Observed 2-step transition proportions*

	0	1	2	3	4	5
0	0.83	0.10	0.04	0.02	0.01	0.00
1	0.35	0.37	0.17	0.08	0.03	0.01
2	0.25	0.22	0.33	0.14	0.05	0.01
3	0.20	0.15	0.22	0.30	0.10	0.03
4	0.15	0.10	0.14	0.27	0.27	0.07
5	0.10	0.05	0.10	0.16	0.24	0.35

per patient vary from 16 (4 days) to 1352 (338 days). Observations are not necessarily consecutive. Often, the subject-specific observations are organized in separated measurement periods, each of them formed by consecutive observations. The minimal measurement period is 1 day (4 measurements), while the maximal one is 213 days (852 measurements).

Table 2 reports the observed proportions of the transitions between the ordinal categories in 2 consecutive measurements. Serial correlation in the data is suggested by the patterns of symptom persistence and of transitions between adjacent categories.

For illustration, we study the relationship between headache severity using university degree status and the usage of analgesics as base variables. Three weather covariates are additionally included. The first is the change in atmospheric pressure from the previous day, categorized in 3 levels, namely from high (>1013 hPa) to low pressure (≤ 1013 hPa), from low to high pressure, and unchanged level of pressure (from low to low or from high to high). The second weather covariate is the relative humidity index with 3 levels, that is less than 60% of humidity, between 60% and 80% of humidity, and more than 80% of humidity. The last weather covariate is windchill categorized into 4 classes, between -50°C and -10°C , between -10°C and 0°C , between 0°C and 10°C , and between 10°C and 30°C .

We consider the 2 binary covariates university degree and usage of analgesics as covariates of primary interest, thus they are included in all considered models. The base model is

$$\text{headache} \sim \text{university} + \text{analgesics}.$$

Furthermore, we consider 2 different autocorrelation parameters γ for subjects with analgesics intake and those without.

For model comparison, it is necessary to fit all models of interest with a pairwise likelihood constructed from the same pairs of observations, that is with the same distance q . We choose q as the value minimizing the generalized variance for the larger model

headache \sim university + analgesics + change + humidity + windchill.

According to this criterion, the overall best performance is obtained with $q = 12$. Thus, we fit all the other (nested) models with this value for q .

Table 3 shows the $2^3 = 8$ models obtained by adding all the possible combinations of the 3 weather covariates to the base model. The relative performance of the k th model with respect to the alternative models can be summarized by CLIC weights defined as $w_k = e^{-\Delta_k} / \sum_{k=1}^8 e^{-\Delta_k}$, where $\Delta_k = (\text{CLIC}_k - \min_i \text{CLIC}_i) / 2$.

Qualitative conclusions should take into account the fitted models with their relative importance expressed through the CLIC weights. In Table 4, for illustration, we show parameter estimates and standard errors only for the 2 best models, namely the model including the change in pressure and the base model.

Table 3. *Migraine data. Maximized log pairwise likelihoods with $q = 12$, CLIC statistics, and CLIC weights for various models fitted to the migraine data*

Change	Humidity	Windchill	Log-pair	CLIC	Weights
—	—	—	-2935.16	5917.15	0.27
*	—	—	-2933.58	5916.30	0.41
—	*	—	-2934.36	5918.84	0.12
—	—	*	-2933.36	5922.53	0.02
*	*	—	-2932.90	5918.33	0.15
*	—	*	-2931.75	5921.93	0.02
—	*	*	-2932.62	5924.59	0.01
*	*	*	-2931.03	5924.20	0.01

The variables included in the models are indicated by symbol *

Table 4. *Migraine data. Estimates and standard errors (SE) from the pairwise likelihood with $q = 12$ for the base model (first 2 columns) and the best model accordingly to CLIC with different autocorrelation parameters for analgesic users and nonanalgesic users (third and fourth column) and with a single common autocorrelation parameter (fifth and sixth column). The levels of the variable change are as follows: 1, change from low to high atmospheric pressure; 2, substantially unchanged atmospheric pressure; and 3, change from high to low atmospheric pressure. The baseline is “no university degree, no intake of analgesics, change from low to high pressure”*

	Estimates	SE	Estimates	SE	Estimates	SE
α_2	0.588	0.046	0.588	0.046	0.589	0.046
α_3	1.136	0.069	1.136	0.069	1.137	0.069
α_4	1.786	0.079	1.787	0.080	1.788	0.080
α_5	2.505	0.109	2.506	0.111	2.508	0.112
Intercept	-0.474	0.226	-0.522	0.223	-0.517	0.223
University	-0.523	0.172	-0.523	0.174	-0.525	0.173
Analgesics	0.558	0.202	0.561	0.205	0.557	0.205
Change 2	—	—	0.031	0.051	0.031	0.051
Change 3	—	—	0.164	0.053	0.164	0.053
γ_F	0.415	0.094	0.424	0.094	0.540	0.031
γ_T	0.556	0.030	0.557	0.030	0.540	0.031
$\gamma_T - \gamma_F$	0.142	0.098	0.133	0.098	0.000	0.000
σ^2	0.566	0.110	0.564	0.111	0.566	0.112

When considering also the other 6 fitted models, we obtain the following overall conclusions. Subjects with university degree tend to suffer from lower levels of headache, while those taking analgesics have stronger symptoms. These variables have more predictive impact on the headache symptoms than the considered weather effects. Among the latter, only the change in the atmospheric pressure is significant in that its decrease is associated with raised headache severity. The categorized relative humidity appears weakly significant and the windchill covariate even less.

Finally, in all fitted models, there is no appreciable difference between the symptom persistence for patients who took analgesics and those who did not. Indeed, the difference between the autocorrelation parameters for the analgesic users (γ_T) and the nonanalgesic users (γ_F) for all models is estimated between 0.133 and 0.148 with standard errors ranging between 0.096 and 0.099. This is confirmed by refitting the models with a common autocorrelation parameter γ for all patients: Table 4 reports the best model with and without separate autocorrelation parameters. The models with common and separate autocorrelations give CLIC = 5915.8 and CLIC = 5916.3, respectively.

5. CONCLUDING REMARKS

We have developed a pseudolikelihood approach for analyzing a large longitudinal study on migraine severity symptoms. The proposed methodology is general and may be useful for other studies with ordinal, as well as binary, outcomes.

The main advantage from pairwise likelihood inference is its computational simplicity. Moreover, since only the specification of bivariate margins is required, our approach relies on model assumptions to a lesser degree than any approach based on a full likelihood approximation. Some loss of efficiency may be experienced for the composite likelihood method compared to a full likelihood, but full likelihood is intractable for large numbers of observations per subject. The study of the efficiency of maximum pairwise likelihood estimators is possible only with a small number of observations per patients as in Joe and Lee (2009) whose results encourage the use of this pseudolikelihood.

The underlying normal assumptions leading to the multivariate probit model were considered mainly for ease of interpretation. However, there are no theoretical restrictions against considering other distributional assumptions. An alternative of possible interest is a cumulative logit model (Agresti, 2002, chapter 7) for the conditional distribution of the response given the random effects.

Other useful variants of the proposed class of models may involve robustification of the random effect distribution, for example by using a Student t -distribution instead of the traditional Gaussian distribution.

Often, in longitudinal studies, the missing data mechanism may not be assumed ignorable and thus likelihood-type analysis based on complete observations are not valid. Modifications of the pairwise likelihood for nonignorable missing data mechanisms are described in Parzen *and others* (2007).

As with standard likelihood inference, maximum pairwise likelihood estimators for variance components fail to correct for the degrees of freedom lost for estimating fixed effects and thus are prone to severe downward bias. When the number of covariates is not small compared to the number of subjects, bias in the estimate of σ^2 can be worthy of attention. Among several bias correction procedures, resampling methods such as jackknife and bootstrap are viable approaches given the low computational cost of pairwise likelihood evaluations. Furthermore, computational saving may be obtained by using first-order approximations instead of complete maximization of the pseudolikelihood for each resampled data set.

Standard errors estimated from the empirical quantities (3.1) and (3.2) may be numerically imprecise for longitudinal studies with few subjects, typically leading to overoptimistic standard errors. More robust variance estimates for small numbers of subjects may be obtained with resampling techniques such as the jackknife or the bootstrap.

SUPPLEMENTARY MATERIAL

Supplementary material is available at <http://biostatistics.oxfordjournals.org>.

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