Evaluation of Autoregressive Covariance Structures for Test-Day Records of Holstein Cows: Estimates of Parameters

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ABSTRACT

The objective of this study was to compare test-day (TD) models with autoregressive covariance structures for the estimation of genetic and environmental components of variance for milk, fat and protein yields, and somatic cell score (SCS) in Holstein cows. Four models were compared: model I (CS model) was a simple TD repeatability animal model with compound symmetry covariance structure for environmental effects, model II (AR_{pe} model) and model III (AR_e model) had first-order autoregressive covariance structures for TD permanent or residual environmental effects, respectively, and model IV (305-d model) was a simple animal model using 305-d records. Data were 106,472 first-lactation TD records of 12,071 Holstein cows calving from 1996 through 2001. Likelihood ratio tests indicated that AR_{pe} and AR_e models fit the data significantly better than the CS model. The AR_e model resulted in slightly smaller estimates of genetic variance and heritability than did the CS model. Estimates of residual variance were always smaller with the CS model than with the AR_e model with the autoregressive covariance structure among TD residual effects. Estimates of heritability with different TD models were in the range of 0.06 to 0.11. The 305-d model resulted in estimates of heritability in the range of 0.11 to 0.36. The autoregressive covariance structure among TD residual effects may help to prevent bias in heritability estimates for milk, fat and protein yields, and SCS.

(**Key words:** milk yield, test day, autoregressive covariance)

Abbreviation key: AIC = Akaike information criterion, AR(1) = first-order autoregressive, CS = compound symmetry, TD = test day.

INTRODUCTION

Direct use of test-day (TD) records in genetic evaluations has several advantages over the use of 305-d lactation records. Effects specific for each individual record may be accounted for with TD models. Most importantly, TD records may be adjusted for the curvilinear shape of the lactation curve (Swalve, 1995b). With TD models, variable amounts of information from incomplete lactations and from different recording systems may also be included. This information may allow for the use of records as they are measured to obtain earlier predictions of genetic potential and thereby help in reducing generation interval (Swalve, 2000).

Test-day records may be viewed as repeated measures of a single trait within a lactation. A main issue with such a model is to account for the covariance structure of the repeated records. The extreme assumption with the simple repeatability animal model of constant environmental and genetic correlations among different TD records may not be a realistic assumption (Henderson, 1984). Simple repeatability model assumes independent residual effects and constant environmental correlation. Correlated records are expected to be less informative than independent ones. Ignoring such correlations when actually present may result in biased estimates of other variance components parameters. Alternatively, monthly TD records can be viewed as different but correlated traits. No predetermined structure or pattern is assumed with this approach for variances or covariances of different TD records.

The structure to account for (co)variation among random correlated effects of repeated TD records should have fewer parameters than the multiple trait approach to avoid over-parameterization. Most importantly, the proposed structure should effectively model the relationships among the effects in the model. The first-order autoregressive process, AR(1), may satisfy these conditions. The AR(1) covariance structure has only one more parameter than the simple repeatability model, which has the compound symme-

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try (CS) covariance structure, and allows for nonconstant covariances (Wade and Quaas, 1993).

Harville (1979) proposed the use of an autoregressive process to model covariance structures for random effects of repeated measures in animal breeding. Similarly, Quaas (1984) suggested an AR(1) process to model the residual covariance structure when animals have repeated measures. Kachman and Everett (1989) used the AR(1) structure to model environmental covariances among TD records within a lactation. A correlation coefficient of 0.6 was used to model nonconstant covariances among TD residual effects. Similarly, Vasconcelos et al. (2004) used the AR(1) structure to predict TD records of uncompleted lactations.

Carvalho et al. (1998) compared expected genetic gain and estimates of genetic parameters with models assuming or not assuming an AR(1) covariance structure among permanent environmental effects within a lactation. They reported that the AR(1) covariance structure for permanent environmental effects within lactation was effective for partitioning total variance and removing noise that would otherwise be confounded with genetic effects. In a subsequent study, Carvalho et al. (2002a) used a similar model and estimated variance components for milk yield of Holstein, Bruna, and Modicana breeds. The estimates of variance components were different for the different breeds.

Recently, random regression models have been widely studied and evaluated for genetic evaluation at national level in many countries. Random regression models have the advantage of flexibility to account for the environmental and genetic components of the shape of lactation curve. However, random regression models require the estimation of large number of parameters and may not be adequate at early or late stages of the lactation. Kettunen et al. (1998) reported negative values for the genetic correlation between early and late TD records within lactation. Meuwissen and Pool (2001) reported similar accuracy of predicting missing records with autoregression and random regression models. However, random regression models required 4 times more dispersion parameters to be estimated compared with the autoregression models.

An AR(1) covariance structure for the residual effects of TD records has not been evaluated before in animal models. Additionally, estimates of genetic parameters with an AR(1) model need to be compared with those from the currently used 305-d model and from simple repeatability models for TD records.

The objective of this study was to compare 2 differently defined autoregressive covariance structures among TD environmental effects. Estimates of vari-

ance components for milk, fat and protein yields, and SCS from models with the AR(1) covariance structures were compared with those from the simple repeatability model and with those from a standard 305-d lactation model.

MATERIALS AND METHODS

Data and Editing

Test-day and 305-d lactation records were from Dairy Records Management Systems in Raleigh, NC. Traits included milk, fat and protein yields, and SCS for first lactations of Holstein cows. The original data set was edited to eliminate records of nonHolsteins, of cows with improper identification, of lactations initiated by abortion, of cows with unknown sires, of cows with missing herd codes, and of cows with unknown birth or calving dates.

Test-day intervals were set to 30 d from 6 to 305 d in milk. Records were assigned to TD based on the DIM when they were recorded rather than their ordinal sequence. Only lactations with twice-daily milking were included in the final data set. Lactation records were required to have at least 5 TD records. To eliminate outliers, milk, fat and protein yields, and SCS greater than 3 standard deviations from their unadjusted means were deleted.

Each sire was required to have at least 10 daughters with a first-lactation record. All herds were required to have at least 15 cows. Lactation records were categorized as treated or not with bST. A minimum of 4 TD records treated with bST was required for any lactation record to be classified as a bST-influenced record. Records with partial bST treatments or those that did not agree with these guidelines were deleted.

The final data set included 12,071 first-lactation records of Holstein cows calving from 1996 through 2001. About 9% of the cows were classified as bST treated. More details about the data with numbers of available TD and 305-d records for each trait are in Table 1.

Models and Methodology

Two approaches for modeling records for estimation of variance component parameters were compared. The first approach used TD records with a single-trait animal model with repeated measures. Repeated measures were accounted for by making 3 different assumptions about the covariance structures for permanent environmental and residual effects. Model I (CS model) assumed a compound symmetry covariance structure for overall environmental effects. For model II (AR_{pe} model), a first-order autoregressive covariance structure was fitted for permanent environmental ef-

Table 1. Numbers of available test-day and 305-d records for milk, fat and protein yields and SCS.

Records	Milk	Fat	Protein	SCS
All TD records	106,472	89,254	89,259	84,952
TD 1	10,459	8404	8408	8342
TD 2	10,923	9088	9088	8714
TD 3	10,944	9216	9216	8738
TD 4	10,960	9274	9274	8781
TD 5	10,968	9259	9259	8799
TD 6	10,931	9234	9234	8717
TD 7	10,864	9170	9170	8647
TD 8	10,898	9189	9189	8691
TD 9	10,528	8875	8875	8404
TD 10	8997	7545	7546	7119
305-d records	12,071	10,130	10,086	10,161
Completed 305-d records	4340	3209	3210	2842

fects of TD records. Model III (AR_e model) was the same as model II but with the first-order autoregressive covariance structure fitted for residual effects of TD records instead of for permanent environmental effects. The second approach with model IV (305-d model) was to use a single-trait animal model without repeated measures with 305-d lactation records instead of TD records.

The equation for the linear mixed model in matrix notation for the first 3 models was:

$$\mathbf{y} = \mathbf{X}\beta + \mathbf{Z}_1\mathbf{a} + \mathbf{Z}_2\mathbf{pe} + \mathbf{e},$$

where $\mathbf{y}$ is a vector of TD records of observations of a trait, β is the vector of fixed effects, $\mathbf{a}$ is the vector of random animal additive genetic effects, $\mathbf{pe}$ is the vector of random cow permanent environmental effects, $\mathbf{e}$ is the vector of random TD residual effects, and $\mathbf{X}$, $\mathbf{Z}_1$, and $\mathbf{Z}_2$ are incidence matrices relating TD observations to fixed, random animal additive genetic, and random cow permanent environmental effects, respectively.

The first moment for all 3 TD models was assumed to be $\mathbf{E}[\mathbf{y}] = \mathbf{X}\beta$. The second moments about the means were assumed to be:

$$\text{Var} = \begin{bmatrix} \mathbf{a} \\ \mathbf{pe} \\ \mathbf{e} \end{bmatrix} = \begin{bmatrix} \mathbf{A}_N\sigma_a^2 & \mathbf{0} & \mathbf{0} \\ \mathbf{0} & \mathbf{I}_n\sigma_{pe}^2 & \mathbf{0} \\ \mathbf{0} & \mathbf{0} & \mathbf{I}_n \otimes \mathbf{I}_t\sigma_e^2 \end{bmatrix} \text{ for CS model,}$$

$$\text{Var} = \begin{bmatrix} \mathbf{a} \\ \mathbf{pe} \\ \mathbf{e} \end{bmatrix} =$$

$$\begin{bmatrix} \mathbf{A}_N\sigma_a^2 & \mathbf{0} & \mathbf{0} \\ \mathbf{0} & \mathbf{I}_n \otimes \mathbf{AR}_{pe_t}\sigma_{pe}^2 & \mathbf{0} \\ \mathbf{0} & \mathbf{0} & \mathbf{I}_n \otimes \mathbf{I}_t\sigma_e^2 \end{bmatrix} \text{ for AR}_{pe} \text{ model,}$$

$$\text{Var} = \begin{bmatrix} \mathbf{a} \\ \mathbf{pe} \\ \mathbf{e} \end{bmatrix} =$$

$$\begin{bmatrix} \mathbf{A}_N\sigma_a^2 & \mathbf{0} & \mathbf{0} \\ \mathbf{0} & \mathbf{I}_n\sigma_{pe}^2 & \mathbf{0} \\ \mathbf{0} & \mathbf{0} & \mathbf{I}_n \otimes \mathbf{AR}_e\sigma_e^2 \end{bmatrix} \text{ for AR}_e \text{ model,}$$

where $\mathbf{A}_N$ is the numerator relationship matrix of order N (the number of animals), $\mathbf{I}_N$ is an identity matrix of order n (the number of cows with records), $\mathbf{I}_t$ is an identity matrix with variable order t (with t the number of TD records of a cow, possibly as many as 10), $\mathbf{AR}_{pe_t}$ and $\mathbf{AR}_e$ are first-order autoregressive correlation matrices of order t among TD permanent and residual environmental effects, respectively, and σ_a^2 , σ_{pe}^2 and σ_e^2 are variances of additive genetic, permanent, and residual environmental effects, respectively, and $\otimes$ is the direct product operator.

The 3 TD models also included common fixed effects of herd test date (HTD), bST treatment, and age at calving in 2-mo intervals from 22 to 38 mo. Effect of DIM as (DIM/30.5) within TD interval was included as a quadratic polynomial to adjust for the shape of the lactation curve. Fixed effects for the 305-d model were: herd-year-season (HYS), bST treatment, and age at calving.

Another TD model (US model) with an unstructured covariance was applied to the data to evaluate the environmental covariances and variances among different monthly TD records with variable numbers of intervals. Similar fixed effects and first-moment model assumptions of the other TD models were also assumed for the unstructured covariance model. However, the unstructured covariance model assumes no particular pattern of covariances or variances for overall environmental effects of TD records. The US model can lead to computational difficulties because it includes too many parameters to be estimated.

Data were analyzed with a single-trait animal model using the ASREML program, release 1.0 (Gilmour et al., 2002). This statistical package uses an average information algorithm and sparse matrix methods. Convergence was assumed when both the log-likeli-

hood and estimated parameters did not change for at least 3 consecutive restarts.

The CS model can be considered nested within, or is a reduced form of the AR_{pe} and AR_e models. The CS, AR_{pe} , and AR_e models share the same fixed effects. The AR_{pe} and AR_e models had the same variance components as the CS model except for one additional variance component. Therefore, the log-likelihood of the CS model may be quantitatively compared with those of the AR_{pe} and AR_e models with the likelihood ratio test with 1 degree of freedom. The log-likelihood ratio statistic is calculated as twice the difference of the log-likelihood values of the complete and reduced models, which can be compared with the critical χ^2 values for the desired probability level and appropriate degrees of freedom based on the difference in number of estimated variance component parameters with each model under consideration (McCulloch and Searle, 2000).

Estimates of 305-d genetic variances were also obtained using TD estimates with CS, AR_{pe} , and AR_e models as: $[305^2 \times \sigma_a^2]$ for yield traits, where σ_a^2 is the TD additive genetic variance. For yield traits, estimates of 305-d phenotypic variance were derived using estimates from the simple repeatability model with CS covariance structure as: $[305^2 \times \sigma_a^2 + 305^2 \times \sigma_{pe}^2 + 30.5^2 \times 10 \times \sigma_e^2]$, where σ_{pe}^2 and σ_e^2 are the permanent environmental and residual variances, respectively. Using estimates from AR_{pe} and AR_e models, estimates of 305-d phenotypic variance for yield traits were derived using the formulas: $[305^2 \times \sigma_a^2 + 30.5^2 \times \sigma_{pe}^2 \times \sum_{i=1}^{10} \sum_{j=1}^{10} \rho^{|i-j|} + 30.5^2 \times 10 \times \sigma_e^2]$ and $[305^2 \times \sigma_a^2 + 305^2 \times \sigma_{pe}^2 + 30.5^2 \times \sigma_e^2 \times \sum_{i=1}^{10} \sum_{j=1}^{10} \rho^{|i-j|}]$, respectively, where ρ is

the AR(1) correlation coefficient. For SCS, similar approaches were used to obtain 305-d estimates of variance components using estimates with TD models. However, the procedure was different by considering that 305-d SCS records are derived by averaging the available TD records rather than accumulating them as with yield traits. For example, the estimates of the phenotypic variance of the 305-d SCS as calculated from TD estimates of variance components with the

AR_e model is: $[\sigma_a^2 + \sigma_{pe}^2 + (100)^{-1} \times \sigma_e^2 \times \sum_{i=1}^{10} \sum_{j=1}^{10} \rho^{|i-j|}]$.

RESULTS AND DISCUSSION

Covariance Structures

The data for all 4 traits of milk, fat and protein yields, and SCS were first analyzed with the US model

Figure 1. Plot of estimates of environmental variances and covariances (kg^2) by intervals between monthly test-day (TD) records for milk yield with unstructured covariance structure model (A) and with CS, AR_{pe} , and AR_e models (B). The CS model is a TD model with compound symmetry covariance structure for TD environmental effects; AR_{pe} model assumes autoregressive covariance structure for TD permanent environmental effects; and AR_e model assumes autoregressive covariance structure for TD residual effects.

with an unstructured covariance matrix for the overall environmental effects of TD records. Estimates of covariances with US model can be plotted over the number of intervals between monthly TD records. Estimates of environmental variances and covariances at different TD intervals with US, CS, AR_{pe} , and AR_e models are plotted in Figures 1 and 2 for milk yield and SCS, respectively. Similar plots were also created for fat and protein yields and will be referred to in the text. The purpose of the plots was to observe the pattern of change of the estimates throughout the lactation rather than to quantify the magnitude of estimates with the different models.

Figure 2. Plot of estimates of environmental variances and covariances by intervals between monthly test-day (TD) records for SCS with unstructured covariance structure model (A) and with CS, AR_{pe} and AR_c models (B). The CS model is a TD model with compound symmetry covariance structure for TD environmental effects; AR_{pe} model assumes autoregressive covariance structure for TD permanent environmental effects and AR_c model assumes autoregressive covariance structure for TD residual effects.

The primary analyses using the US model revealed larger environmental covariances between adjacent TD records than those further apart for milk, fat and protein yields, and SCS. Estimates of overall environmental correlations using the unstructured covariance models were in the range of 0.20 to 0.67 for milk, 0.13 to 0.50 for fat, 0.15 to 0.63 for protein, and 0.20 to 0.61 for SCS TD records. On average, TD records at the beginning and at the end of the lactation had the smallest estimates of correlations with the other TD records.

Estimates of covariances did not decline to zero regardless of the number of intervals between all combinations of TD records. Estimates of covariances seem to reach a plateau mostly after the third or fourth interval between TD records, which may suggest the presence of a common environmental variance for all TD records. Any covariance structure, to be useful,

should be able to account for this effect. The patterns of change of estimates of unstructured covariances were the same for all traits: milk, fat and protein yields, and SCS.

The pattern of change of estimates of covariances with the US model agrees well with the defined characteristics of the first-order autoregressive covariance structure. The AR(1) covariance structure allows for a decrease in covariance with increasing interval and assumes the correlation to be strictly a function of number of intervals (Littell et al., 2002). Depending on the value of the correlation coefficient and the number of intervals between measures, the AR(1) covariance structure may allow for correlations to reach an approximate plateau.

Gadini (1997) used 2 trait analyses of TD records for milk, fat and protein yields, and SCS to estimate genetic and phenotypic correlations and variances among TD records but did not report estimates of environmental covariances. The environmental covariances derived from genetic and phenotypic estimates showed a pattern similar to the one in this study except for a few unusual estimates that did not follow the general pattern. The calculated estimates of environmental correlations were also very close to the estimates in this study and were in the range of 0.28 to 0.67 for milk yield, 0.24 to 0.53 for fat yield, 0.23 to 0.62 for protein yield, and 0.21 to 0.64 for SCS. Meyer et al. (1989) reported a similar pattern for environmental correlations among TD records for milk, fat and protein yields. Norman et al. (1999) reported comparable findings with smaller correlations with increased number of intervals between TD records. They also reported the smallest correlations were for pairs of TD records at early and late stages of lactation.

Except for SCS, the largest environmental variances were generally for first or last TD records. However, the largest estimates of variances were not more than 30, 40, 47, and 11% larger than the smallest estimates of variances for milk, fat, protein, and SCS records, respectively. Likewise, Ali and Schaeffer (1987) and Meyer et al. (1989), using fixed regression models, and White et al. (1999) using a random regression model reported the largest variances for milk yields at the beginning and at the end of lactation. Swalve (1995c) found similar patterns of estimates of environmental variances over the course of lactation for milk, fat and protein yields.

The estimates of covariances with the CS model did not change, regardless of the number of intervals between TD records that resulted in smaller estimates of covariances between adjacent records and larger estimates of covariances between more separated TD for milk, fat, and protein records (Figures 1B and 2B).

Estimates of overall environmental correlations (r_e) among TD records with the CS model were obtained as the ratio of estimates of permanent to the total environmental variances ($r_e = \sigma_{pe}^2 / (\sigma_{pe}^2 + \sigma_e^2)$, where σ_{pe}^2 and σ_e^2 are the permanent environmental and residual variances, respectively). Estimates of overall environmental correlations were 0.47, 0.30, 0.40, and 0.45 for milk, fat, protein, and SCS, respectively. These estimates are assumed the same among all TD records, regardless of the number of intervals among the records. Estimates of environmental variances with the CS model were generally similar to the mean of environmental variance estimates obtained with the US model. The CS model obviously does not seem to be a close approximation for the covariances among most TD records.

The estimates of environmental covariances with the AR_{pe} model decreased as the number of intervals between records increased but at a decreasing rate (Figures 1B and 2B). The environmental covariances between TD records did not reach a plateau, regardless of the number of intervals between any of the 10 TD records. This result may be explained by the large estimates of the autoregressive correlation coefficient (ρ) with the AR_{pe} model (0.82 to 0.90).

Estimates of overall environmental correlations (r_e) for different TD intervals with the AR_{pe} model can be obtained as function of residual and permanent environmental variances and the AR(1) rho correlation coefficients as $r_e = (\sigma_{pe}^2 \times \rho_{pe}^n) / (\sigma_{pe}^2 + \sigma_e^2)$, where σ_{pe}^2 is the permanent environmental variance, ρ_{pe}^n is the autoregressive correlation coefficient for number of intervals (n) between the TD records, and σ_e^2 is the residual variance. Estimates of overall environmental correlations were in the ranges of 0.23 to 0.63 for milk, 0.10 to 0.41 for fat, 0.17 to 0.57 for protein, and 0.23 to 0.55 for SCS. Estimates of overall environmental variances with the AR_{pe} model were in very close agreement with estimates obtained with the US model. Several studies have reported the use of a similar definition of the AR(1) covariance structure as the AR_{pe} model to account for the nonconstant environmental covariances among TD records (Carvalho et al., 1998; 2002a,b; Vasconcelos et al., 2004).

The AR_e model with the AR(1) covariance structure among the residuals resulted in covariance matrices most analogous to the general covariance structures as estimated with the US model for all traits. As shown in Figures 1B and 2B, estimates of environmental covariances with the AR_e model were very close to the median points for unstructured covariances for most TD intervals. Estimates of covariances among adjacent records decreased markedly at the beginning and

continued to decrease, but at a much slower rate, as the number of intervals between the TD records increased, which resulted in a plateau after the third or fourth TD intervals. The estimate of the total environmental covariance between any 2 records cannot drop below the estimate of permanent environmental variance. Norman et al. (1999) found that correlations among TD records for milk yield traits and SCS were well modeled with the autoregressive correlation structure. They compared the AR(1) with other structures such as identity (I), intercept (J), and heterogeneous variances at the beginning, middle, or late stages of lactation among other structures.

Estimates of overall environmental correlations (r_e) with the AR_e model can be obtained as $r_e = (\sigma_{pe}^2 + \sigma_e^2 \times \rho_{pe}^n) / (\sigma_{pe}^2 + \sigma_e^2)$, where σ_{pe}^2 is the permanent environmental variance, σ_e^2 is the residual variance and ρ_{pe}^n is the autoregressive correlation coefficient for number of intervals (n) between TD records. Estimates of overall environmental correlations with the AR_e model were in the range of 0.39 to 0.62 for milk, 0.24 to 0.42 for fat, 0.32 to 0.56 for protein, and 0.39 to 0.56 for SCS. The ranges of the estimates of overall environmental correlation with the AR_e model were similar to the range of estimates obtained with the unstructured covariance model.

Based on the above comparison of the environmental covariance structures with the different TD models, a preliminary conclusion may be drawn that the AR(1) covariance structure for the TD permanent environmental and residual effects may be a suitable approximation for modeling of environmental covariance matrices for TD milk, fat and protein yields, and SCS. More quantitative tests will be carried out to compare the performance of the suggested models.

Likelihood Ratio Tests

A more objective way to compare the models is to calculate the log-likelihoods of the models and conduct tests of significance among them whenever possible. The log-likelihood values are shown in Tables 2, 3, 4, and 5 for milk, fat and protein yields, and SCS, respectively. Based on the likelihood ratio tests, AR_{pe} and AR_e models fit the data significantly better than the simple repeatability model with the CS covariance structure (CS model) for all 4 traits. All of the calculated likelihood ratio statistics were much larger than the critical χ^2 values. The smallest calculated statistic was 2158 between the AR_e model and the CS model for fat yield. This value is about 199 times larger than the critical χ^2 value at $P = 0.001$ with 1 degree of freedom, which is 10.83. The Akaike information crite-

Table 2. Estimates of variance components (kg^2), heritability, repeatability, and standard errors for test-day and 305-d milk yields.¹

Component ²	CS model		AR _{pe} model		AR _e model		305-d model	
	Estimate	SE	Estimate	SE	Estimate	SE	Estimate	SE
σ_a^2	2.59	0.39	2.06	0.33	2.22	0.35	281,454	54,470
σ_{pe}^2	11.31	0.37	17.16	0.33	9.26	0.34		
σ_e^2	12.62	0.06	7.03	0.07	14.63	0.10	1,799,260	53,530
σ_p^2	26.51	0.23	26.24	0.22	26.11	0.22	2,080,714	31,950
ρ			0.88	<0.01	0.38	<0.01		
h^2	0.10	0.01	0.08	0.01	0.09	0.01		
h_{305}^2 ³	0.17		0.10		0.15		0.14	0.03
r	0.52	<0.01	0.29–0.66		0.44–0.65			
Log-L (LRT) ⁴	-1865		-1822 (8586)		-1831 (6704)		-717	
AIC ⁵	3730		3644		3663			

¹CS model is a simple repeatability model with compound symmetry covariance structure for TD environmental effects; AR_{pe} model assumes autoregressive covariance structure for TD permanent environmental effects; AR_e model assumes autoregressive covariance structure for TD residual effects; and 305-d model is simple animal model using 305-d records.

²Additive genetic variance (σ_a^2), permanent environmental variance (σ_{pe}^2), residual variance (σ_e^2), phenotypic variance (σ_p^2), AR(1) correlation coefficient (ρ), heritability (h^2), heritability of 305-d lactations (h_{305}^2), and repeatability of TD records (r).

³Using estimates with TD models except for 305-d model.

⁴Log-L = Logarithm of likelihood values divided by 100 (LRT: likelihood ratio test statistic for comparing CS model with AR_{pe} and AR_e models).

⁵AIC = Akaike information criteria: $-2 (\log\text{-likelihood}) + 2 (\text{number of variance parameters})$, values are divisions of 100.

Table 3. Estimates of variance components (kg^2), heritability, repeatability, and standard errors for test-day and 305-d fat yield.^{1,2}

Component ³	CS model		AR _{pe} model		AR _e model		305-d model	
	Estimate	SE	Estimate	SE	Estimate	SE	Estimate	SE
σ_a^2	0.44	0.06	0.49	0.06	0.42	0.06	1064	136
σ_{pe}^2	1.22	0.06	2.01	0.05	1.00	0.05		
σ_e^2	2.90	0.02	2.05	0.02	3.11	0.02	1903	113
σ_p^2	4.55	0.04	4.55	0.04	4.53	0.03	2967	56
ρ			0.83	0.01	0.23	0.01		
h^2	0.10	0.01	0.11	0.01	0.09	0.01		
h_{305}^2 ⁴	0.23		0.13		0.22		0.36	0.04
r	0.36	0.01	0.19–0.48		0.31–0.47			
Log-L (LRT) ⁵	-959		-945 (2828)		-948 (2158)		-339	
AIC ⁶	1918		1890		1897			

¹CS model is a simple repeatability model with compound symmetry covariance structure for TD environmental effects; AR_{pe} model assumes autoregressive covariance structure for TD permanent environmental effects; AR_e model assumes autoregressive covariance structure for TD residual effects; and 305-d model is simple animal model using 305-d records.

²Estimates of variance components are multiples of 100 for CS, AR_{pe}, and AR_e models.

³Additive genetic variance (σ_a^2), permanent environmental variance (σ_{pe}^2), residual variance (σ_e^2), phenotypic variance (σ_p^2), AR(1) correlation coefficient (ρ), heritability (h^2), heritability of 305-d lactations (h_{305}^2), and repeatability of TD records (r).

⁴Using estimates with TD models except for 305-d model.

⁵Log-L = Logarithm of likelihood values divided by 100 (LRT: likelihood ratio test statistic for comparing CS model with AR_{pe} and AR_e models).

⁶AIC = Akaike information criteria: $-2 (\log\text{-likelihood}) + 2 (\text{number of variance parameters})$, values are divisions of 100.

Table 4. Estimates of variance components (kg^2), heritability, repeatability, and standard errors for test day and 305-d protein yield.^{1,2}

Component ³	CS model		AR _{pe} model		AR _e model		305-d model	
	Estimate	SE	Estimate	SE	Estimate	SE	Estimate	SE
σ_a^2	0.24	0.04	0.18	0.03	0.21	0.03	288.23	52.30
σ_{pe}^2	0.82	0.03	1.36	0.03	0.65	0.03		
σ_e^2	1.21	0.01	0.71	0.01	1.39	0.01	1289	48.80
σ_p^2	2.27	0.02	2.25	0.02	2.24	0.02	1578	27.44
ρ			0.86	0.01	0.35	0.01		
h^2	0.11	0.02	0.08	0.01	0.09	0.01		
h_{305}^2 ⁴	0.20		0.10		0.19		0.18	0.03
r	0.467	0.005	0.235–0.601		0.381–0.600			
Log-L (LRT) ⁵	–645		–616 (5948)		–622 (4638)		–317	
AIC ⁶	1291		1232		1245			

¹CS model is a simple repeatability model with compound symmetry covariance structure for TD environmental effects; AR_{pe} model assumes autoregressive covariance structure for TD permanent environmental effects; AR_e model assumes autoregressive covariance structure for TD residual effects; and 305-d model is simple animal model using 305-d records.

²Estimates of variance components are multiples of 100 for CS, AR_{pe} and AR_e models.

³Additive genetic variance (σ_a^2), permanent environmental variance (σ_{pe}^2), residual variance (σ_e^2), phenotypic variance (σ_p^2), AR(1) correlation coefficient (ρ), heritability (h^2), heritability of 305-d lactations (h_{305}^2), and repeatability of TD records (r).

⁴Using estimates with TD models except for 305-d model.

⁵Log-L = Logarithm of likelihood values divided by 100 (LRT: likelihood ratio test statistic for comparing CS model with AR_{pe} and AR_e models).

⁶AIC = Akaike information criteria: $-2 (\log\text{-likelihood}) + 2 (\text{number of variance parameters})$, values are divisions of 100.

Table 5. Estimates of variance components, heritability, repeatability, and standard errors for test day and 305-d SCS.¹

Component ³	CS model		AR _{pe} model		AR _e model		305-d model	
	Estimate	SE	Estimate	SE	Estimate	SE	Estimate	SE
σ_a^2	0.18	0.04	0.19	0.04	0.18	0.04	0.17	0.04
σ_{pe}^2	1.27	0.04	1.75	0.04	1.11	0.04		
σ_e^2	1.59	0.01	1.08	0.01	1.74	0.01	1.42	0.04
σ_p^2	3.04	0.03	3.02	0.03	3.02	0.03	1.59	0.03
ρ			0.90	<0.01	0.28	0.01		
h^2	0.06	0.01	0.06	0.01	0.06	0.01		
h_{305}^2 ³	0.11		0.07		0.11		0.11	0.03
r	0.48	0.01	0.28–0.58		0.43–0.59			
Log-L (LRT) ⁴	–716		–698 (3686)		–703 (2700)		–67.0	
AIC ⁵	1433		1396		1406			

¹CS model is a simple repeatability model with compound symmetry covariance structure for TD environmental effects; AR_{pe} model assumes autoregressive covariance structure for TD permanent environmental effects; AR_e model assumes autoregressive covariance structure for TD residual effects; and 305-d model is simple animal model using 305-d records.

²Additive genetic variance (σ_a^2), permanent environmental variance (σ_{pe}^2), residual variance (σ_e^2), phenotypic variance (σ_p^2), AR(1) correlation coefficient (ρ), heritability (h^2), heritability of 305-d lactations (h_{305}^2), and repeatability of TD records (r).

³Using estimates with TD models except for 305-d model.

⁴Log-L = Logarithm of likelihood values divided by 100 (LRT: likelihood ratio test statistic for comparing CS model with AR_{pe} and AR_e models).

⁵AIC = Akaike information criteria: $-2 (\log\text{-likelihood}) + 2 (\text{number of variance parameters})$, values are divisions of 100.

rion (AIC) gave similar results for the AR_{pe} and AR_e models compared with the CS model for all 4 traits. The better fit of the AR_{pe} and AR_e models to the data than the CS model may indicate that the estimates of the variance components with AR_{pe} and AR_e models are more accurate, given the data.

Generally, differences in the log-likelihood and AIC values between AR_{pe} and AR_e models were not as great as differences between AR_{pe} and CS models or AR_e and CS models. Based on the log-likelihood and AIC values, the AR_{pe} model seems to have better fit to the data than the AR_e model. However, no quantitative test is available to verify the difference in fit of AR_{pe} and AR_e models, because both models have the same number of estimated variance components parameters, and the χ^2 value thus has zero degrees of freedom.

Estimation of Variance Components

Estimates of variance components with all 4 models are presented in Table 2 for milk yield, in Table 3 for fat yield, in Table 4 for protein yield, and in Table 5 for SCS. Estimates of heritability were in the range of 0.08 to 0.11 for yield traits, and heritability was 0.06 for SCS with different TD models. Estimates of heritability with the 305-d model were in the range of 0.14 to 0.36 for yield traits, and heritability was 0.11 for SCS. The standard errors of all estimates of variance components and heritability were small, which indicates that the size of the data set was adequate. Furthermore, none of the estimates of parameters was fixed at boundaries or out of the parameter space.

Estimates of heritability in the literature are variable with both TD and 305-d lactation models. Estimates vary for different populations or datasets, countries, regions, periods or year, methods, and models of analysis and other factors. Estimates with a cubic spline random regression model were as small as 0.09, 0.06, 0.09, and 0.02 for TD milk, fat, protein, and SCS, respectively (DeGroot, 2003). Similarly, Tsuruta (1998) reported small estimates of heritability in the range of 0.10 to 0.26, 0.05 to 0.12, 0.09 to 0.21, and 0.03 to 0.09 for TD milk, fat, protein, and SCS, respectively. Other researchers have reported greater estimates of heritability for TD yield traits (Auran, 1976; Danell, 1982; Pander et al., 1992). Reents et al. (1994) reported estimates of heritability for SCS of 0.06 to 0.10 for TD records in the first lactation. Heritability estimates for milk yield using 305-d lactation records have been reported to be as large as 0.49 (Pander et al., 1992) and 0.39 (Swalve, 1995a) and as small as 0.24 (Visscher and Goddard, 1995).

Estimates of autoregressive correlation coefficients (ρ) for the residuals with the AR_e model were small

and in the range of 0.23 to 0.38. The small estimates of the correlation coefficient for residual effects is because AR_e model also includes permanent environmental effects that are assumed to be common for all TD records for each cow. Estimates of the autoregressive correlation coefficient with the AR_{pe} model for the permanent environmental effects, which are expected to be highly repeatable among different TD records, were large (0.83 to 0.90). Carvalho et al. (1998) reported estimates of the AR(1) correlation coefficients among permanent environmental effects in the range of 0.57 to 0.83 for milk yield in Holstein and Lucerna cattle. Their moderate estimates may be because their model included an additional permanent environmental effect along with the autocorrelated ones and independent residual effects.

Estimates of repeatability (r) with the different TD models may be obtained by the general formula of $r = (\sigma_a^2 + \rho_{pe}^n \sigma_{pe}^2 + \rho_e^n \sigma_e^2) / (\sigma_a^2 + \sigma_{pe}^2 + \sigma_e^2)$, where ρ_{pe}^n is the autoregressive correlation coefficient among TD permanent environmental effects and is 1 for all models except for AR_{pe} model, ρ_e^n is the autoregressive correlation coefficient among TD residual effects and is 0 for all models except for the AR_e model, n is the number of intervals between TD records, σ_a^2 is TD additive genetic variance, σ_{pe}^2 and σ_e^2 are TD permanent environmental and residual variances, respectively. Estimates of repeatability with the CS model were in the range of 0.36 to 0.52 for different traits. Estimates of repeatability with the AR_{pe} model were 0.29 to 0.66 for milk yield, 0.19 to 0.48 for fat yield, 0.23 to 0.60 for protein yield and 0.28 to 0.58 for SCS. Estimates of repeatability with the AR_e model had larger lower limits and similar upper limits as estimates with the AR_{pe} model. The smaller lower limits of repeatability with the AR_{pe} model are due to the large estimates of the correlation coefficient with the AR_{pe} model. The large correlation coefficient allows the permanent environmental covariance between TD records with the AR_{pe} model to considerably decrease with the increase in number of intervals between records without reaching plateau.

The AR_e model with the AR(1) covariance structure among the residual effects resulted in smaller estimates of heritability for all yield traits than with the CS model. Estimates of additive genetic and permanent environmental variances for yield traits were always smaller with the AR_e model than with the CS model. Similarly, estimates of total environmental and phenotypic variances were slightly smaller with the AR_e model than with the CS model for all traits.

Estimates of residual variances were always less with the CS model than with the AR_e model. The in-

Table 6. Relative amount of time (%) required for genetic evaluation with different models compared with AR_e model.¹

	CS model	AR _{pe} model	305-d model
Milk	9.2	27.1	-93.2
Fat	-10.7	12.0	-94.6
Protein	-1.4	9.7	-94.4
SSC	-12.9	22.4	-105.8

¹AR_e model assumes autoregressive covariance structure for TD residual effects; CS model is a simple repeatability model with compound symmetry covariance structure for TD environmental effects; AR_{pe} model assumes autoregressive covariance structure for TD permanent environmental effects; and 305-d model is simple animal model using 305-d records.

creases in estimates of residual variance with the AR_e model compared with estimates with the CS model were about 16% for milk yield, 7% for fat yield, 14% for protein yield, and 10% for SCS. The residual variance can become smaller if a nonzero correlation among repeated measures on a given subject is ignored.

Estimates of heritability for 305-d lactations were also calculated using linear functions of estimates of variance components obtained with the TD models. The estimates of 305-d heritability were larger when the estimates of TD variance components were obtained with either the CS model or AR_e than when they were obtained with the AR_{pe} model (Tables 2 through 5). This result may be due to the larger estimate of permanent environmental variance and the associated 305-d estimate of phenotypic variance with the AR_{pe} model compared with the CS and AR_e models. The 305-d heritability estimates from the CS model were slightly larger than those estimated by the AR_e model. Except for fat yield and SCS, the estimates of 305-d heritability with the CS model or AR_e model were larger than those obtained directly with the 305-d model.

The amount of computational time required to obtain genetic evaluations with different models were compared relative to the AR_e model (Table 6). The use of the AR_e model will require a relatively similar amount of computational time as the CS model (from 9.2% less to 12.9% more time). In all cases, the AR_{pe} model required more computational time than did the AR_e model (9.7 to 27.1% more time). All TD models required a considerable increase in the amount of computational time required compared with the 305-d model.

CONCLUSIONS

The autoregressive covariance structure for TD permanent environmental or residual effects is more suitable to model the covariance among TD environmental

effects than the simple repeatability model for milk, fat and protein yields, and SCS data. The autoregressive covariance structure could account for the nonconstant environmental correlations between TD records with variable interval lengths. The CS model did not effectively account for decreasing environmental correlations between pairs of TD records separated by more intervals. Estimates of residual variance, genetic variance, and heritability with the CS model may thus be biased. Models with the AR(1) covariance structures can help to obtain more accurate estimates of heritability for TD milk, fat and protein yields, and SCS of Holstein cows. The AR(1) covariance structures for the permanent environmental or residual effects may be a reasonable approximation for modeling of environmental covariance matrices among TD milk, fat and protein yields, and SCS of Holstein cows.

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