

2020, Vol. 25, No. 2, 129-142 http://dx.doi.org/10.1037/met0000238

Evaluating a Theory-Based Hypothesis Against Its Complement Using an AIC-Type Information Criterion With an Application to Facial Burn Injury

Leonard Vanbrabant Ghent University and Municipal Health Service (GGD) West-Brabant, Noord-Brabant, Breda, the Netherlands Nancy Van Loey Association of Dutch Burn Centers, Noord-Holland, Beverwijk, the Netherlands, and Utrecht University

Rebecca M. Kuiper Utrecht University

Abstract

An information criterion (IC) like the Akaike IC (AIC), can be used to select the best hypothesis from a set of competing theory-based hypotheses. An IC developed to evaluate theory-based order-restricted hypotheses is the Generalized Order-Restricted Information Criterion (GORIC). Like for any IC, the values themselves are not interpretable but only comparable. To improve the interpretation regarding the strength, GORIC weights and related evidence ratios can be computed. However, if the unconstrained hypothesis (the default) is used as competing hypothesis, the evidence ratio is not affected by sample-size nor effect-size in case the hypothesis of interest is (also) in agreement with the data. In practice, this means that in such a case strong support for the order-restricted hypothesis against its complement using the GORIC (weights). We show how to compute the GORIC value for the complement, which cannot be achieved by current methods. In a small simulation study, we show that the evidence ratio for the order-restricted hypothesis versus the complement increases for larger samples and/or effect-sizes, while the evidence ratio for the order-restricted hypothesis versus the unconstrained hypothesis remains bounded. An empirical example about facial burn injury illustrates our method and shows that using the complement as competing hypothesis results in much more support for the hypothesis of interest than using the unconstrained hypothesis as competing hypothesis.

Translational Abstract

In an informative hypothesis, academic expertise (i.e., theory) about the population of interest is included in the hypothesis in terms of order-restrictions on the model parameters. As an example, in an ANOVA setting, we might expect that the group means follow a certain order (e.g., $H_1: \mu_1 < \mu_2 < \mu_3 < \mu_4$). As another example, in a linear regression model, we might expect that the (standardized) regression coefficients are subject to multiple one-sided restrictions (e.g., $H_2: \beta_1 > 0$; $\beta_2 > 0$; $\beta_3 > 0$). In the absence of a competing informative hypothesis, an order-restricted hypothesis is typically evaluated against the unconstrained hypothesis H_u (all orderings are allowed). However, a problem arises if the order-restricted hypothesis H_1 is in agreement with the data: Then, the estimated parameters for H_1 are identical to the estimated parameters of H_u . Therefore, we introduce an AIC-type/information-theoretic method for evaluating an informative hypothesis H_m against its complement, where the complement H_c is defined as H_c : not H_m (all orderings are allowed except the ordering in H_1). The method is illustrated using an empirical example about facial burn injury. In addition, the method is implemented in the user-friendly R package **restriktor**.

Keywords: complement, evidence ratio, level probabilities, model selection, order restrictions

The evaluation of a researcher's theory is often key to research. For example, consider a study about facial burn injury. A burn event can have an avers impact on a person's quality of life. The scars can affect physical appearance and may constitute a source of

The data and the ideas in this article are currently not posted or shared elsewhere. Rebecca M. Kuiper is supported by a grant from the Netherlands organization for scientific research: NWO-VENI-451-16-019.

Correspondence concerning this article should be addressed to Leonard Vanbrabant, Municipal Health Service (GGD) West-Brabant, Doornboslaan 225-227, 4816 CZ Breda, the Netherlands. E-mail: l.vanbrabant@ggdwestbrabant.nl

This article was published Online First October 31, 2019.

Leonard Vanbrabant, Department of Data-analysis, Ghent University, and Municipal Health Service (GGD) West-Brabant, Noord-Brabant, Breda, the Netherlands; Nancy Van Loey, Association of Dutch Burn Centers, Noord-Holland, Beverwijk, the Netherlands, and Department of Clinical Psychology, Utrecht University; Rebecca M. Kuiper, Department of Methods and Statistics, Utrecht University.

Leonard Vanbrabant is now at the Department of Methods and Statistics, Utrecht University.

rumination acting as a reminder to the event (Nolen-Hoeksema, Wisco, & Lyubomirsky, 2008). Based on previous research (Van Loey et al., 2014), it is expected that injury characteristics that may be perceived as distressing such as facial burn injury and larger burns may be triggers for the activation and prolongation of rumination. In addition, a gender effect is also expected because disfiguring scars resulting from burns may be of greater importance to woman as compared with men (Ghriwati et al., 2017). We therefore hypothesized that the means of rumination for men with and without facial burn injury and the mean of rumination for women without facial burn injury would be lower than the mean of rumination for women with facial burn injury. In symbols, this hypothesis can be stated as: H_1 : { $\mu_{men; no facial burns}$, $\mu_{men; facial burns}$, $\mu_{\text{women; no facial burns}} \} \le \mu_{\text{women; facial burns}}$, where μ are the population means for rumination for the four groups determined by gender and facial burns, adjusted for the population effects of some covariates. Note that no particular order is assumed among the first three means. This form of theory-based hypothesis is known as an order-restricted (OR) hypothesis or informative hypothesis (Hoijtink, 2012) because the order of the means is restricted based on theory and/or academic expertise.

To evaluate such OR hypothesis, three methods can be distinguished, that is, OR hypothesis testing (e.g., Kudô, 1963), model selection using OR information criteria (e.g., Anraku, 1999; Mulder, & Raftery, in press), and model selection using the Bayes factor (e.g., Mulder, Hoijtink, & Klugkist, 2010). In this current article, we focus on model selection using information criteria. Akaike's IC (AIC; see, e.g., Akaike, 1973, 1998) is probably the most familiar and widely used information criterion employed in the social and behavioral sciences. Nevertheless, the AIC is not suitable when the model parameters (e.g., means or regression coefficients) are subject to order restrictions. A modification of the AIC that can deal with simple order restrictions¹ (i.e., Order-Restricted Information Criterion [ORIC]) in the exponential family was proposed by Anraku (1999). Kuiper, Hoijtink, and Silvapulle (2011) generalized the ORIC (GORIC) to accommodate any linear inequality restrictions in multivariate normal linear models (except for range restrictions, which bounds a parameter to a specific interval, e.g., $-1 \le \mu \le 1$). Information criteria like the AIC, ORIC, and GORIC are calculated as minus two times the maximum log-likelihood (under the hypothesized model) plus twice a penalty term value. The main difference between the methods is in calculating the penalty term value, which is less straightforward to compute in case of order restrictions.

The evaluation of an OR hypothesis (e.g., H_1) requires at least one competing hypothesis. Sometimes researchers have another hypothesis of interest and want to know which hypothesis is best, for example H_1 versus H_2 : $\mu_{\text{men; no facial burns}} \leq \mu_{\text{men; facial burns}} \leq \mu_{\text{women; no facial burns}} \leq \mu_{\text{women; facial burns}}$. The difference between the two is that in H_2 the means are fully ordered. In practice, researchers often do not have such a specific competing hypothesis and only the unconstrained hypothesis H_u , where no restrictions are imposed on the model parameters, remains included as competing hypothesis in the set. Therefore, in this article, we focus solely on the set of hypotheses with one OR hypothesis H_m and the unconstrained hypothesis H_u .

The hypothesis with the lowest GORIC value is the preferred one. The GORIC values themselves are not interpretable and only the differences between the values can be inspected. To improve the interpretation, so-called GORIC weights (w_m) can be computed, which are derived from the Akaike weights (Akaike, 1978; Burnham & Anderson, 2002) and are comparable with posterior model probabilities (see Burnham & Anderson, 2002, pp. 302– 305). An IC weight w_m represents the relative likelihood of hypothesis *m* given the data and the set of *M* hypotheses (Burnham & Anderson, 2002; Kuiper, 2011; Wagenmakers & Farrell, 2004). For example, if we compare hypothesis H_1 against hypothesis H_u , we can examine the ratio of the two corresponding weights, that is w_1/w_u . This evidence ratio is considered as the strength of evidence in favor of model *m* (in this example, m = 1) of being the best model (Burnham & Anderson, 2002; Wagenmakers & Farrell, 2004).

The evidence ratio should increase for larger samples and/or effect-sizes. However, if the OR hypothesis of interest H_m is in agreement with the data, increasing the sample-size and/or effectsize does not affect the evidence ratio if the unconstrained hypothesis is used as competing hypothesis (assuming that H_m is true and remains in agreement with the data). In that case, both hypotheses H_m and H_u are in line with the data, since H_u is always in line with the data, and consequently both hypotheses have the same maximized log-likelihood value. Then, the difference in GORIC values equals the difference in penalty term values, which are independent of sample-size and effect-size. The latter case is illustrated in Figure 1, where we generated 500 data sets according to an ANOVA model with four uncorrelated ordered means which are in agreement with H_1 , with a sample-size of n = 50 per group and various effect-sizes f (Cohen, 1988, pp. 274-275). The results (cf. see triangles in Figure 1) show that at first the mean evidence ratio (on a log scale) of w_1/w_μ increases slightly for increasing effectsizes and that afterward it stabilizes at an upper-bound value of approximately $exp(1.09) \approx 2.97$ on the original scale. It is at this point that the data are for each simulation run in agreement with H_1 and thus the maximized log-likelihood values of H_1 and H_{μ} are the same. The boundary value equates the exponential difference of the penalty term values between H_{μ} and H_{1} , that is, $\exp(5.00-3.91) = \exp(1.09) \approx 2.97$; as will become clear later on. Consequently, strong support for the OR parameters is not expressed in a high evidence ratio if compared to the unconstrained hypothesis and many research questions may be erroneously dismissed as irrelevant. It is important to note that the boundary issue is not specific to the order-restricted IC literature but can also be found in the Bayes factor literature (e.g., Mulder et al., 2009, 2010). They solve the boundary issue by comparing the orderrestricted hypothesis against its complement.

The objective of this study is to show that this upper bound issue can also be solved by replacing the unconstrained hypothesis by the complement of the hypothesis of interest (cf. see circles in Figure 1). The complement is defined as $H_c = \neg H_m$, where \neg denotes "not." For example, for the OR hypothesis H_1 : { $\mu_{men; no facial burns}$, $\mu_{men; facial burns}$, $\mu_{women; no facial burns}$ } $\leq \mu_{women; facial burns}$ with four means there are four ways in which the four means can be ordered in such an ordering. Hypothesis H_1 consists of one of these four combinations, therefore the complement represents the 4 – 1 = 3 remaining ways in which the four means can be ordered. In

¹ Simple order restrictions are of the form $\mu_1 \leq \mu_2 \leq \ldots \leq \mu_J$, where *J* is the total number of groups and where \leq can be replaced by an =.



Figure 1. Mean evidence ratio (on a log scale and based on 500 simulations) for hypothesis H_1 : { μ_1 , μ_2 , μ_3 } $\leq \mu_4$ compared with the unconstrained hypothesis (mean w_1/w_u) and compared with its complement (mean w_1/w_c) when H_1 is true, for n = 50, and various effect-sizes (*f*).

this "simple" case, it is easy to write out the complement but this is often not the case. In many cases, it is a cumbersome or even impossible task to write up all possible combinations that belong to the complement, because the number of combinations increases excessively with the number of parameters. For example, for the OR hypothesis H_2 with four means there are 24 ways (i.e., 4! = $4 \times 3 \times 2 \times 1$) in which the four means can be ordered in a simple order ordering. Moreover, the GORIC is often not defined when the complement comprises multiple combinations. This is because, the GORIC is only defined for restrictions that form a closed convex cone² (ccc, e.g., H_1 and H_2).

The novelty of this article is that we introduce the GORIC for the situation that the restrictions in the complement are not a ccc. We show (a) how to determine the log-likelihood for the complement and (b) how to determine the penalty term value for the complement.

The remainder of this article is organized as follows. First, we provide some technical background about the computation of the GORIC and the corresponding penalty term value for the unconstrained hypothesis and an OR hypothesis H_m . Second, we introduce how the GORIC is computed for the complement of H_m . Third, we illustrate our method with the empirical example introduced at the beginning of this section. Finally, we give some concluding remarks and recommendations.

Technical Background

Before we introduce the GORIC for the complement, some technical background is inevitable. The results given in this part are for the linear regression model, where the regression coefficients are subject to linear inequality and/or linear equality restrictions. The method can readily be adapted to multivariate normal linear models. This is briefly discussed in the last section.

Linear Model and Order-Restricted Hypotheses

Consider the standard linear regression model,

$$y_i = \boldsymbol{x}_i^T \boldsymbol{\theta} + \boldsymbol{\epsilon}_i, \ i = 1, \ \dots, \ n, \tag{1}$$

where $\mathbf{\theta} = (\theta_1, \ldots, \theta_p)^T$ is the parameter vector of interest, $\mathbf{x}_i = (\mathbf{x}_{i1}, \ldots, \mathbf{x}_{ip})^T$ is a vector of predictor variables³ for person *i*, and $\mathbf{\epsilon}_i = (\mathbf{\epsilon}_1, \ldots, \mathbf{\epsilon}_n)^T$ is a vector of normally distributed random errors: $\mathbf{\epsilon}_i \sim \mathcal{N}(0, \sigma^2)$. Let the (unconstrained) maximum likelihood estimates be denoted by $\hat{\mathbf{\theta}}$ and the order-restricted mle's denoted by $\tilde{\mathbf{\theta}}_m$. The latter is the solution of maximizing the likelihood under the restrictions in H_m , a well-studied restricted optimization problem in the statistical literature (Nocedal & Wright, 2006).

We consider three types of hypotheses, namely $H_u: \mathbf{0} \in \mathbb{R}^p$, where \mathbb{R}^p is the p-dimensional Euclidean space, $H_m: \mathbf{0} \in C$, where C is also a space in \mathbb{R}^p and is a (reallocated) closed convex cone (Kuiper, Hoijtink, & Silvapulle, 2012), and $H_c: \neg H_m$, which is not necessarily a (reallocated) closed convex cone. Because, most applications only involve linear restrictions, we only consider linear hypotheses.

The GORIC

The GORIC for the unconstrained hypothesis H_u is defined as

$$GORIC_u = -2 \times LL_u + 2 \times PT_u, \tag{2}$$

where LL_u is the maximized log-likelihood value and the penalty term value is defined as $PT_u = 1 + p$. Note that $GORIC_u$ equals the AIC for H_u .

The GORIC for the OR hypothesis H_m is defined as

$$GORIC_m = -2 \times LL_m + 2 \times PT_m, \tag{3}$$

where LL_m is the maximized log-likelihood value for the OR hypothesis H_m and PT_m is the penalty term value for H_m . The penalty term value equals

$$\mathrm{PT}_m = 1 + \sum_{j=0}^p j \times LP_j(p, \mathbf{\Sigma}, H_m).$$

In an univariate regression model $\Sigma = (\mathbf{X}^T \mathbf{X})^{-1}$ is the unscaled⁴ covariance matrix of the parameters with $\mathbf{X} = (\mathbf{x}_1^T, \dots, \mathbf{x}_n^T)^T$ of order $n \times p$ and $LP_j(p, \Sigma, H_m)$ are the level probabilities (chi-bar-square weights). A level probability LP_j , is the probability that the OR mle $\tilde{\mathbf{\theta}}_m$ has *j* levels (under the null-hypothesis), where j = p - "the number of active restrictions"; and the *LP*'s sum to 1. We will clarify the computation of PT_m using two examples. Consider Figure 2a, where

² If we only consider linear hypotheses, then a closed convex cone (ccc) can be written in the form $R\theta \ge \mathbf{r}$, where R is a matrix with known constants, \mathbf{r} a vector with known constants and $\boldsymbol{\theta}$ a vector with the model parameters. If the restrictions in H_m or H_c cannot be written in this form, then they do not form a ccc. For example, the complement of $H : \mu_1 \ge 0$; $\mu_2 \ge 0$, that is $H_c : \theta_1 \le 0 \& \theta_2 \ge 0$; $\theta_1 \le 0 \& \theta_2 \le 0$ cannot be written in the form $R\theta \ge \mathbf{r}$. For more details regarding ccc's, we refer the interested reader to Berman (1973).

³ In case of an intercept, $x_{i1} = 1$ for all *i*'s and θ_1 is interpreted as the intercept.

⁴ The calculation of the level probabilities is invariant for positive constants like σ^2 (known or unknown; Silvapulle & Sen, 2005, p. 32) or even for $\tilde{\sigma}^2$, the OR mle of σ^2 .



Figure 2. Illustration to illuminate the computation of the penalty term value of H_m . The gray shaded area is the permissible area under H_m . (a) $H_3: \theta_1 \ge 0, \theta_2 \ge 0$. (b) $H_4: \theta_1 \ge \theta_2$.

the unconstrained parameter space is determined by the two parameters θ_1 and θ_2 and is divided into four quadrants (Q₁ to Q₄). If we assume that θ_1 and θ_2 are independent of each other (i.e., $\Sigma = I$, where **I** is an identity matrix), then each quadrant is assigned a level probability of 0.25 under H_0 : $\theta_1 = \theta_2 = 0$. The permissible gray shaded area is defined by the order restrictions H_3 : $\theta_1 \ge 0$, $\theta_2 \ge 0$. Then, the probability that j = 2, that is, none of the restrictions are active (i.e., j = p - 0 = 2 - 0 = 2), is 0.25 (Q₁). The probability that j = 1, that is, that one restriction is active (i.e., j = 2 - 1 = 1), is 0.25 + 0.25 = 0.50 (Q₂ and Q₄). The probability that j = 0, that is, that both restrictions are active (i.e., j = 2 - 2 = 0), is 0.25 (Q₃). Consequently, the penalty term value for the OR hypothesis H_3 can be computed as $PT_3 = 1 + 0 \times 0.25 + 1 \times 0.50 + 2 \times 0.25 = 2$. In addition, consider Figure 2b, where the parameter space is restricted by the order restrictions $H_4: \theta_1 \ge \theta_2$. Because the order restriction divides the unconstrained parameter space into two spaces, Q_1 and Q_2 are now two half-spaces. Again, assume that $\Sigma = I$ and again we have two parameters (i.e., p = 2), but now we only have one order restriction (i.e., $q_1 = 1$). Consequently, there can be zero or at maximum one active restriction and thus the probability that j = 0, that is, that we have two active restrictions, is 0. This is because, if we impose one order restriction on two parameters, one parameter is allowed to vary freely, while the other parameter is restricted by the value of this free parameter. The probability that j = 1, that is, that the order restriction is active, is 0.5 (Q_2). The probability that j = 2, that is, that the order restriction is not active, is 0.5 (Q_1) . Hence, the penalty term value for the OR hypothesis H_4 is computed by $PT_4 =$ $1 + 0 \times 0 + 1 \times 0.5 + 2 \times 0.5 = 2.5.$

Equipped with this knowledge, we rewrite the penalty term PT_m . This becomes helpful for determining the penalty for the complement of H_m , which is discussed later on. Let $q_1 > 0$ be the number of order restrictions and $q_2 > 0$ the number of equality restrictions. Then, $p = q_1 + q_2 + F$, with F the number of remaining, free parameters. Then, the penalty for H_m can be rewritten as:

$$PT_{m} = 1 + \sum_{j=0}^{p} j \times LP_{j}(p, \mathbf{\Sigma}, H_{m})$$

= 1 + 0 × LP₀ + 1 × LP₁ + ... + p × LP_p
= 1 + (F + 0) × LP_{F+0} + ... + (p - q_{2}) × LP_{p-q_{2}}
= 1 + (F + 0) × LP_{F+0} + ... + (F + q_{1}) × LP_{F+q_{1}}
= 1 + F + 0 × LP_{F+0} + ... + q_{1} × LP_{F+q_{1}}, (4)

using that LP_0 to LP_{F-1} are 0 (because F free parameters in H_m denote that there are always at least F "inactive" restrictions); and that LP_{p-q_2+1} to LP_p are 0 (because the q_2 equality restrictions in H_m are always active); and that the LP's sum to 1.

From the penalty term value PT_m , it follows that for q_1 order restrictions and q_2 equality restrictions, and thus $F = p - q_2$ free parameters, the penalty term value for a hypothesis with solely equality restrictions equals $1 + F = 1 + p - q_2$, which equals the penalty term value of the AIC.

In case of order restrictions, the exact computation of the level probabilities when $\Sigma \neq I$ and for q > 4 is a difficult task in general because the probabilities can no longer be expressed in closed form. Fortunately, the probabilities can be approximated by using the multivariate normal probability distribution function with additional Monte Carlo steps (Grömping, 2010) or they can be computed easily and sufficiently precise by Monte Carlo simulation (Silvapulle & Sen, 2005; Wolak, 1987).

Introduction of the GORIC for H_c

Here, we introduce the GORIC for the complement of H_m , which is defined as

$$GORIC_c = -2 \times LL_c + 2 \times PT_c, \tag{5}$$

where LL_c is the maximized log-likelihood value for the complement of H_m and PT_c is the penalty term value. Recall that for the computation of the GORIC value for H_m the order-restricted hypothesis is required to be a ccc. However, the complement H_c is in many cases not a ccc. For example, the complement of $H_3: \theta_1 \ge 0$, $\theta_2 \ge 0$, that is, H_c , is constructed by the quadrants Q_2 , Q_3 , and Q_4 (see Figure 3a), and can be written as

$$\theta_1 \leq 0 & \& & \theta_2 \geq 0 \quad (Q_2)$$

and
$$H_c: \quad \theta_1 \leq 0 & \& & \theta_2 \leq 0 \quad (Q_3)$$

and
$$\theta_1 \geq 0 & \& & \theta_2 \leq 0 \quad (Q_4).$$

Note that in this case, the complement can be written out easily but, for many hypotheses, it is a difficult or even impossible task to write up the complement. Because, H_c is not a ccc, the LL_c and the PT_c values cannot be computed directly like the LL_m and the PT_m values.

Computing the Log-Likelihood for H_c

To compute the LL_c value, we first need to ascertain whether the restrictions in H_m are in line with the data or not. If at least one inequality restriction is violated, then the data are automatically in line with the complement and the LL_c value equals the LL_u value. This is illustrated for H_3 : $\theta_1 \ge 0$, $\theta_2 \ge 0$ in Figure 3a, where the permissible area is Q_1 and the quadrants Q_2 , Q_3 , and Q_4 form the complement. Because the unconstrained mle's $\hat{\theta}$ lie in Q₃ (here, both restrictions are violated), the data are in line with the complement and the LL_c is equal to LL_u . Note that the same applies if the mle's lie in Q₂ or Q₄. On the other hand, if the data are in line with the restrictions in H_3 , then we have to find the mle's of θ that are closest to $\theta \in H_c$, given Σ , which is denoted by θ_c . Let us inspect a bivariate case with $\Sigma = I$ (solid circles), as depicted in Figure 3b. The solid circles of the contour plot indicate that the two parameters θ_1 and θ_2 are uncorrelated. As a reminder, the lines of the contour plot correspond to parameter values which have equal

log-likelihood values and lines closer to $\hat{\theta}$ result in a higher log-likelihood value, because $\hat{\boldsymbol{\theta}}$ is the value for which the loglikelihood is maximized (without imposing restrictions on the parameters). Clearly, the solution $\hat{\theta}_c$ is on the boundary of the restricted parameter space H_m . Because there are many boundary solutions (see thick black lines), we have to search for a solution that has the shortest distance between $\hat{\theta}$ and the two boundaries, given Σ . Fortunately, we do not have to investigate each point on the thick black lines but only the points $\hat{\boldsymbol{\theta}}_{c1}$ and $\hat{\boldsymbol{\theta}}_{c2}$. The point $\hat{\boldsymbol{\theta}}_{c1}$ is computed by treating the inequality restriction for θ_1 as equality restriction (i.e., $\theta_1 = 0$, $\theta_2 \ge 0$). Analogously, for the point $\hat{\theta}_{c2}$, where θ_2 is treated as equality restriction (i.e., $\theta_1 \ge 0$, $\theta_2 = 0$). Thus, in general, there are in total q_1 possibilities to be investigated. Notably, in case of equality restrictions, all q_2 equalities are "freed." The LL corresponding to that point that results in the highest log-likelihood value, given Σ , equals the LL_c value (here, $\tilde{\boldsymbol{\theta}}_{c1}$).

As mentioned above, the solution of $\boldsymbol{\theta}_c$ is dependent on the covariance matrix $\boldsymbol{\Sigma}$. To clarify this, again consider Figure 3b. The solid contour lines show the solution of $\boldsymbol{\tilde{\theta}}_c$ if $\boldsymbol{\Sigma}$ is an identity matrix (i.e., $\boldsymbol{\tilde{\theta}}_{c1}$). It can easily been seen that, if the covariance matrix is not an identity matrix (e.g., see dot-dashed line) that the solution of $\boldsymbol{\tilde{\theta}}_c$ alters (here, it is $\boldsymbol{\tilde{\theta}}_{c2}$).

Computing the Penalty Term Value for H_c

The penalty term value for H_m can be seen as the expected number of "inactive" restrictions (i.e., j = p – "the number of active restrictions" which implies that free parameters denote "inactive" restrictions) plus one for the variance term. The expected number of "inactive" restrictions is the sum of the expected number of "inactive" restrictions for $1 + p - q_2$ subspaces (where 0 to $p - q_2$ restrictions are "inactive"). If we apply this

Figure 3. The gray shaded area (i.e., Q_1) is the permissible area under H_3 and the other quadrants (white area) are the permissible areas under its complement H_c . (a) $H_3 : \theta_1 \ge 0, \theta_2 \ge 0$. The mle's $\hat{\theta}$ lie in Q_3 and is thus in agreement with H_c . (b) $H_3 : \theta_1 \ge 0, \theta_2 \ge 0$, for $\Sigma = \mathbf{I}$ (solid lines) and for $\Sigma \neq \mathbf{I}$ (dashed lines). The mle's $\hat{\theta}$ lie in Q_1 and is thus *not* in agreement with H_c .



134

principle on the complement, then only two distinct subspaces can be distinguished (and not $1 + p - q_2$). The first subspace is the one not in agreement with the complement (cf. Q_1 ; i.e., the space fully in agreement with H_m). For this subspace, the number of "inactive" restrictions equals the number of free parameters in H_c , denoted by F^{c} . Note that the free parameters in $H_{m}(F)$ remain free in H_{c} and the q_2 equality restrictions in H_m are "freed" in H_c . Thus, $F^c =$ $F + q_2 = p - q_1$. The probability of F^c levels in H_c ($LP_{F^c}^c$) equals the probability of having $p - q_2$ levels in H_m (LP_{p-q_2} ; cf. probability of ending up in Q_1 is $LP_{p-q_2} = LP_2 = 0.25$), that is, the probability finding the mle in H_m (under the null). The second subspace is the one fully in agreement with the complement (cf. not Q_1 , that is, Q_2 , Q_3 , and Q_4) and has $p = F^c + q_1$ "inactive" restrictions (note that there are no equality restrictions in H_c). The corresponding level probability is denoted by $LP_{F^c+q_1}^c$ and equals the probability of not finding the *mle* in H_m but in H_c which equals $(1 - LP_{p-q_2})$. Mimicking Equation 4, the penalty term value for H_c is given by

$$PT_{c} = 1 + (F^{c} + 0) \times LP_{F^{c}+0}^{c} + (F^{c} + q_{1}) \times LP_{F^{c}+q_{1}}^{c}$$

= 1 + (p - q_{1}) + q_{1} × (1 - LP_{p-q_{2}})
= 1 + p - q_{1} \times LP_{p-q_{2}}. (7)

Another way of establishing PT_c follows from the fact that the complement is the whole space minus the space fully in agreement with H_m and, therefore, its penalty equals the penalty of the whole space (i.e., 1 + p) minus the expected number of "inactive" restrictions in the space H_m . The expression for the latter equals the last part in Equation 4, that is, $q_1 \times LP_{F+q_1}$. Then, it follows that the expected number of "inactive" restrictions in H_c equals

$$\mathbf{PT}_c = 1 + p - q_1 \times LP_{F+q_1}.$$
(8)

This expression is equal to Equation 7, because $F + q_1 = p - q_2$.

To illustrate, the penalty term value for the complement of H_3 : $\theta_1 \ge 0, \theta_2 \ge 0$ is computed by $PT_c = 1 + 2 - 2 \times 0.25 = 2.5$ and the penalty term value for the complement of $H_4: \theta_1 \ge \theta_2$ is computed by $PT_c = 1 + 2 - 1 \times 0.5 = 2.5$. Because the complement of H_4 (i.e., $\theta_1 \le \theta_2$) is a ccc, we can use the PT_m formula as well, which also renders a penalty of 2.5. In Appendix A, we illustrate the computation of the PT_m and the PT_c values in case of three parameters.

In the next section, we show by means of a brief illustration and the results from a simulation study that the evidence ratio for the order-restricted hypothesis compared with its complement is boundless.

Unbounded GORIC Weights in Case of the Complement

Once the GORIC values for H_m and its complement are known, the GORIC weights can be easily obtained as follows

$$w_{s} = \frac{\exp\{-0.5(\text{GORIC}_{s})\}}{\exp\{-0.5(\text{GORIC}_{m})\} + \exp\{-0.5(\text{GORIC}_{c})\}},$$
 (9)

where the subscript *s* equals *m* or *c* for hypothesis H_m and hypothesis H_c , respectively. From these weights, we can determine the evidence ratio for H_m against its complement w_m/w_c . This ratio is interpreted as the strength of evidence for H_m given

the data and H_c (Kuiper, 2011; Wagenmakers & Farrell, 2004). For example, for Figure 3b with $\Sigma \neq I$ (e.g., dot-dashed lines), n = 50, and f = 0.20, the evidence ratio for $H_3: \theta_1 \ge 0, \theta_2 \ge 0$ 0 compared with H_c equals $w_3/w_c = 0.92/0.08 = 11.50$. This ratio tells us that hypothesis H_3 is the best out of the two (because it is larger than 1) and that H_3 is 11.50 times more supported than H_c . Notably, because we sampled from H_3 , H_3 is true. To contrast, if we want to determine the evidence ratio for H_3 against the unconstrained hypothesis H_{μ} , that is, w_3/w_{μ} , we have to replace the GORIC_c by the GORIC_u in Equation 9 and s equals m or u for hypothesis H_m and hypothesis H_u , respectively. Note that w_3 now not equates 0.92 from above, because the weights depend on the set of hypotheses. Therefore, if H_c is replaced by H_{u} , the weights must be recomputed for the two hypotheses in the set. Then, the evidence ratio equals $w_3/w_{\mu} =$ $0.62/0.38 \approx 1.63$. This clearly shows the advantage of using the complement as competing hypothesis. Namely for the same data, we could obtain support for H_3 of 1.63 (the maximum support) or a support of 11.50 when comparing to H_c (using n = 50).

In Appendix B, we present a simulation study in which we investigated the performance of the evidence ratio weights. The simulation results show the benefits of evaluating an OR hypothesis against its complement. While, for small effect-sizes and/or sample-sizes, the difference between the evidence ratio for the true H_m when using the complement or unconstrained as competing hypothesis is minimal, the difference increases rapidly and profoundly for larger effect-sizes and/or sample-sizes. More importantly, the evidence ratio for the true H_m against its complement is boundless for increasing effect-sizes and/or sample-size and might therefore be more compelling, whereas the evidence ratio for the unconstrained hypothesis as competing hypothesis has an upper bound. Therefore, we recommend to replace the unconstrained hypothesis by the complement of the hypothesis of interest as competing hypothesis.

Burns Example

To illustrate the method, we analyze the empirical example introduced in the introduction in which we sought to determine possible risk factors for ruminating thoughts after a burn injury. The data are based on a cohort study consisting of 245 individuals with burns, aged 18- to 74-years-old. The response variable is rumination. Moreover, for the current illustration, we included gender (0 = men, 1 = women) and facial burns (0 = no, 1 = yes) together with its interaction as predictor variables and Hospital Anxiety and Depression Scale (HADS; M = 3.85, SD = 3.66), age (M = 41.06, SD = 13.94), and the number of surgical operations, which is a measure of severity of the burns (SO; M = 1.14, SD = 1.76) as covariates.

Reconsider, the hypothesis of interest

 H_1 : { $\mu_{\text{men; no facial burns}}$, $\mu_{\text{men; facial burns}}$, $\mu_{\text{women; no facial burns}}$ } $\leq \mu_{\text{women; facial burns}}$. A natural choice to evaluate the OR hypothesis H_1 would be an order-restricted 2 × 2 ANCOVA model. Because an ANCOVA is just a special case of the linear regression model, the model can be written as a linear function. To obtain adjusted means for a person with an average score on the covariates, the covariates HADS, age and SO are centered at their average and are denoted by Z_HADS, Z_age and Z_SO, respectively. Then, the model can be written as follows:

Rumination_{*i*} = $\theta_1 + \theta_2$ facialBurns_{*i*} + θ_3 gender_{*i*}

+
$$\theta_4$$
 gender_i × facialBurns_i + θ_5 Z_HADS_i
+ θ_6 Z_age_i + θ_7 Z_SO_i + ϵ_i ,
for $i = 1, \dots, 245$.

On the left-hand side of the = operator, we have the response variable rumination and on the right-hand side we have the factors facial burns and gender and its interaction, and the centered covariates Z_HADS, Z_age and Z_SO. The interaction between gender and facial burns is included using the \times operator. Then, the four adjusted means with average scores on the covariates are computed as:

$$\mu_{\text{men}; \text{ no facial burns}} = \theta_1$$

$$\mu_{\text{men}; \text{ facial burns}} = \theta_1 + \theta_2$$

$$\mu_{\text{women; no facial burns}} = \theta_1 + \theta_3$$

$$\mu_{\text{women; facial burns}} = \theta_1 + \theta_2 + \theta_3 + \theta_4.$$

The R (R Core Team, 2019) code and the output from the analyses can be found in Appendix C.

The results show that the OR hypothesis H_1 is 0.891/0.109 \approx 8.198 times more supported by the data than its complement. For comparison, the results for the unconstrained hypothesis show that hypothesis H_1 is 0.734/0.266 \approx 2.754 times more supported by the data, which is its maximum support. Assuming H_1 is true, using the complement of H_m instead of H_u , we have now have more compelling evidence.

Summary and Discussion

In this article, we introduced the evaluation of an order-restricted (OR) hypothesis against its complement using the GORIC (weights). The GORIC is an information criterion that can be used to evaluate competing hypotheses in univariate and multivariate normal linear models, where the regression parameters θ are subject to linear (in)equality restrictions. The interpretation can be improved by computing GORIC weights and related evidence ratios reflecting the strength of evidence for one hypothesis versus another.

We advise that one should evaluate their theory against its complement H_c instead of the unconstrained hypothesis H_{μ} . The advantage of our method is that the evidence ratio for an OR hypothesis H_m compared with its complement is boundless. The evidence ratio for H_m compared to H_u is neither increased by a larger sample-size nor by a larger effect-size, if the data are in agreement with the hypothesis of interest (i.e., theory). Consequently, using the complement as competing hypothesis leads to much more support for the hypothesis of interest assuming it is true, compared with using the unconstrained hypothesis as competing hypothesis. In case the complement is not true, then the results are comparable to evaluating the order-restricted hypothesis against the unconstrained hypothesis. Furthermore, in case that H_m is almost true, then there is less support for H_m when compared with its complement than when compared with the unconstrained hypothesis. This is because the log-likelihood values are almost identical and the difference between the penalty term values is larger between H_m and H_u than between H_m and H_c . Consequently, in such cases, it is also better to evaluate H_m against its complement as the support for H_m against H_u might be overstated because H_m is not true. Besides that the complement has practical benefits,

it is also more substantive: When looking at the interpretation of the complement and the unconstrained hypothesis, the latter can be seen as all possible theories including the target hypothesis H_m , whereas the former denotes the possible theories without the target hypothesis H_m . We believe that this makes more sense if you would like to know whether your theory is better than all other theories (and how much). Comparing your theory against all possible theories including the targeted one makes less sense to us.

The method was illustrated using an empirical example about facial burn injury. In six easy steps (shown in Appendix C), we showed how to compute the evidence ratio of the researchers theory against its complement using the R package **restriktor** (see http://www.restriktor.org).

We assumed that researchers often do not have specific competing hypotheses. While, this is probably often the case, it is conceivable that the set of hypotheses contains more than one competing hypothesis. In these cases, the problem that the evidence ratio for H_m against H_u is not affected by increasing samplesize and/or effect-size after a specific value can still occur. For example, consider the set with three hypotheses: H_2 , $H_5 : \mu_1 \leq$ $\mu_2 \leq \mu_3 = \mu_4$ (which is a subset of H_2) and the unconstrained hypothesis H_u . If H_5 is true, then all three hypotheses are true and all evidence ratios are bounded (Kuiper et al., 2011, p. 107). However, the evaluation of a set of multiple OR hypotheses against its complement is less straightforward because determining the complement for multiple hypotheses might not always be trivial (especially for software).

The results presented in this article are for the univariate linear regression model but fortunately they can easily be adapted for the multivariate normal linear model. One should keep in mind that, unlike in the univariate setting, where $\tilde{\boldsymbol{\theta}}$ does *not* depend on the order-restricted covariance matrix, denoted by $\tilde{\boldsymbol{\Sigma}}$, in the multivariate normal linear model $\tilde{\boldsymbol{\theta}}$ does depend on $\tilde{\boldsymbol{\Sigma}}$ and $\tilde{\boldsymbol{\Sigma}}$ on $\tilde{\boldsymbol{\theta}}$ (Kuiper et al., 2012). Hence, an iterative procedure is needed to calculate them. The procedure is implemented in restriktor.

References

- Akaike, H. (1973). Information theory and an extension of the maximum likelihood principle. In B. Petrov & F. Caski (Eds.), *Proceedings of the* second international symposium on information theory (pp. 267–281). Budapest, Hungary: Akademiai Kiado.
- Akaike, H. (1978). On the likelihood of a time series model. *The Statistician*, 27, 217–235. http://dx.doi.org/10.2307/2988185
- Akaike, H. (1998). Information theory and an extension of the maximum likelihood principle. In E. Parzen, K. Tanabe, & G. Kitagawa (Eds.), Selected papers of Hirotugu Akaike. Springer series in statistics (Perspectives in Statistics). New York, NY: Springer. http://dx.doi.org/10 .1007/978-1-4612-1694-0_15
- Anraku, K. (1999). An information criterion for parameters under a simple order restriction. *Biometrika*, 86, 141–152. http://dx.doi.org/10.1093/ biomet/86.1.141
- Berman, A. (1973). Convex cones and linear inequalities. Cones, matrices and mathematical programming. Lecture notes in economics and Mathematical Systems (Operations Research, Computer Science, Social Science). Berlin, Heidelberg: Springer. http://dx.doi.org/10.1007/978-3-642-80730-5_1
- Burnham, K., & Anderson, D. (2002). Model selection and multimodel inference: A practical information-theoretic approach (2nd ed.). New York, NY: Spring-Verlag.

- Cohen, J. (1988). *Statistical power analysis for the behavioral sciences* (2nd ed.). Hillsdale, NJ: Erlbaum.
- Ghriwati, A., Sutter, M., Pierce, B., Perrin, P., Wiechman, S., & Schneider, J. (2017). Two-year gender differences in satisfaction with appearance after burn injury and prediction of five-year depression: A latent growth curve approach. *Archives of Physical Medicine and Rehabilitation*, 98, 2274–2279. http://dx.doi.org/10.1016/j.apmr.2017.04.011
- Grömping, U. (2010). Inference with linear equality and inequality constraints using R: The package ic.infer. *Journal of statistical software*, 33, 1–31. http://dx.doi.org/10.18637/jss.v033.i10
- Hoijtink, H. (2012). Informative hypotheses: Theory and practice for behavioral and social scientists. Boca Raton, FL: Taylor & Francis.
- Kudô, A. (1963). A multivariate analogue of the one-sided test. *Biometrika*, 50, 403–418. http://dx.doi.org/10.2307/2333909
- Kuiper, R. (2011). Model selection criteria: How to evaluate order restrictions (Doctoral dissertation). Utrecht University, Utrecht, the Netherlands. Retrieved from https://dspace.library.uu.nl/handle/1874/224499
- Kuiper, R., Hoijtink, H., & Silvapulle, M. (2011). An Akaike-type information criterion for model selection under inequality constraints. *Bi-ometrika*, 98, 495–501. http://dx.doi.org/10.1093/biomet/asr002
- Kuiper, R., Hoijtink, H., & Silvapulle, M. (2012). Generalization of the order-restricted information criterion for multivariate normal linear models. *Journal of Statistical Planning and Inference*, 142, 2454–2463. http://dx.doi.org/10.1016/j.jspi.2012.03.007
- Mulder, J., Hoijtink, H., & Klugkist, I. (2010). Equality and inequality constrained multivariate linear models: Objective model selection using constrained posterior priors. *Journal of Statistical Planning and Inference*, 140, 887–906. http://dx.doi.org/10.1016/j.jspi.2009.09.022

- Mulder, J., Klugkist, I., van de Schoot, R., Meeus, W., Selfhout, M., & Hoijtink, H. (2009). Bayesian model selection of informative hypotheses for repeated measurements. *Journal of Mathematical Psychology*, *53*, 530–546. http://dx.doi.org/10.1016/j.jmp.2009.09.003
- Mulder, J., & Raftery, A. (in press). BIC extensions for order-constrained model selection. *Sociological Methods & Research*.
- Nocedal, J., & Wright, S. (2006). *Numerical optimization*. New York, NY: Spring-Verlag.
- Nolen-Hoeksema, S., Wisco, B. E., & Lyubomirsky, S. (2008). Rethinking Rumination. *Perspectives on Psychological Science*, *3*, 400–424. http:// dx.doi.org/10.1111/j.1745-6924.2008.00088.x
- R Core Team. (2019). R: A language and environment for statistical computing [Computer software manual]. Retrieved from https://www .R-project.org/
- Silvapulle, M., & Sen, P. (2005). Constrained statistical inference: Order, inequality, and shape restrictions. Hoboken, NJ: Wiley.
- Van Loey, N. E., Oggel, A., Goemanne, A.-S., Braem, L., Vanbrabant, L., & Geenen, R. (2014). Cognitive emotion regulation strategies and neuroticism in relation to depressive symptoms following burn injury: A longitudinal study with a 2-year follow-up. *Journal of Behavioral Medicine*, 37, 839–848. http://dx.doi.org/10.1007/s10865-013-9545-2
- Wagenmakers, E., & Farrell, S. (2004). AIC model selection using Akaike weights. *Psychonomic Bulletin & Review*, 11, 192–196. http://dx.doi .org/10.3758/bf03206482
- Wolak, F. (1987). An exact test for multiple inequality and equality constraints in the linear regression model. *Journal of the American Statistical Association*, 82, 782–793. http://dx.doi.org/10.1080/01621459.1987.10478499

Appendix A

Example of Computing the PT_c in Case of Three Parameters

Consider Figure A1, the unconstrained parameter space is determined by the three parameters θ_1 , θ_2 , and θ_3 (and is of course the whole space and not just the one depicted in Figure A1). The gray shaded area is a closed convex cone and is defined by the order restrictions H_{A1} : $\theta_1 \leq \theta_2$, $\theta_1 \leq \theta_3$. The level probabilities corresponding to H_{A1} (assuming $\Sigma = I$)) equal $LP_0 = 0, LP_1 =$ $\frac{1}{6}$, $LP_2 = \frac{1}{2}$ and $LP_3 = \frac{1}{3}$. Next, we will elaborate on the values for LP_0 and LP_3 . The level probability $LP_3 = \frac{1}{3}$ is the probability that the vector with OR estimates $\tilde{\boldsymbol{\theta}}$ is identical to the unconstrained estimates (i.e., $\tilde{\theta} = \hat{\theta}$ and i = 3). This probability equals the proportion of the gray shaded area compared with the whole cube and of course also of H_{A1} versus the whole space. A level probability of $LP_0 = 0$, that is, the probability that there are j = 0levels, which means that it is impossible that the vector with OR estimates $\tilde{\boldsymbol{\theta}}$ has three active restrictions, which makes sense with two order restrictions. Hence, the penalty term value for H_{A1} equals $PT_{A1} = 1 + 0 \times 0 + \frac{1}{6} \times 1 + \frac{1}{2} \times 2 + \frac{1}{3} \times 3 = 3\frac{1}{6}$. Using the expression in Equation 8, the penalty term value for H_c equals $PT_c = 1 + p - q_1 \times LP_{p-0} = 1 + 3 - 2 \times \frac{1}{3} = 3\frac{1}{3}$. For comparison, $PT_{\mu} = 1 + 3 = 4$.



Figure A1. The permissible, gray area is defined by $H_{A1}: \theta_1 \le \theta_2, \theta_1 \le \theta_3$, depicted for θ_1, θ_2 and θ_3 between -1 and 1.

This article is intended solely for the personal use of the individual user and is not to be disseminated broadly This document is copyrighted by the American Psychological Association or one of its allied publishers.

Appendix B

Simulation Study

Design

We generated 500 samples according to the ANOVA model⁵ $y_i = \mu_1 x_{i1} + \ldots + \mu_4 x_{i4} + \epsilon_i$, $i = 1, \ldots, n$, where we assume that the residuals are standard normally distributed. We considered the OR hypothesis $H_1: \mu_1 \le \mu_2 \le \mu_3 \le \mu_4$, its complement $H_c: \neg H_1$ and the unconstrained hypothesis H_{μ} : μ_1 , μ_2 , μ_3 , μ_4 . Note that H_c does not equal $\mu_1 \ge \mu_2 \ge \mu_3 \ge \mu_4$; it does contain this but also the other (22) orderings of simple ordering combinations of μ_1 to μ_4 (excluding the one ordering in H_m). Data were generated under hypothesis H_1 with four uncorrelated independent means of size n = 30, 50, 100, 200, 500 per group and for a variety of differences among the population means, using effect-size f = 0, 0.10, $0.20, \ldots, 1$ (Cohen, 1988, pp. 274–275). Notably, f = 0 corresponds to sampling from the boundary of both H_m and H_c . If we sample values from a H_1 population with increasing effect-size, this will evidently lead to more and more support for H_1 . Let the differences between the means, d, be equally spaced, where d is defined as $d = \frac{2f\sqrt{p}}{\sqrt{\sum_{i=1}^{p}(2i-1-p)^2}}$ under the restriction that $\sum_{i=1}^{p} \mu_i = 0$ and $\sigma = 1$. Then, the *p* ordered means can be computed as $\mu_i = \frac{-(p-1)d}{2} + (i-1)d$. Appendix D shows the computed population means for the various effect-sizes (f).

Simulation Results

All results are obtained using the R package **restriktor** (see http://www.restriktor.org) employing the GORIC function. The results of the simulation study are presented in Figure B1, B2, and B3, and are obtained by computing the mean value of the relative evidences in each of the 500 simulation runs. Furthermore, to improve visibility, we took the natural logarithm values of the means and we used a varying range of sample-sizes and effect-sizes.

The results clearly illustrate the benefits of evaluating H_m versus its complement: The mean evidence ratio for H_1 versus H_c (mean w_1/w_c) increases rapidly for larger effect-sizes (see Figure B1a) and sample-sizes (see Figure B1b), while the mean evidence ratio using the unconstrained hypothesis as competing hypothesis (mean w_1/w_u) is clearly bounded after a certain value (see Figures B1c and B1d, respectively). To illustrate, consider for example Figure B1a, where the mean evidence ratio for H_1 versus H_c (mean w_1/w_c) for a medium effect-size (f = 0.30) and n = 100 is $\exp(2.63) \approx$ 13.87 (on the original scale), while the mean evidence ratio for H_1 versus H_u (mean w_1/w_u) is bounded at $\exp(1.92) \approx 6.82$. Note that the value 1.92 equals the difference in penalty term values; with $PT_u - PT_1 = (1.00 + 4.00) - (1.00 + 2.08) \approx 1.92$, which equals the difference in GORIC values, because the log-likelihood values are here the same (i.e., $LL_{\mu} = LL_{1}$).

For small effect-sizes and small samples, the mean evidence ratio for H_1 using H_c is slightly lower than when using H_{μ} . For example, for f = 0.10 and n = 30 the mean evidence ratio for w_1/w_c is exp(1.50) ≈ 4.48 and for w_1/w_u the mean evidence ratio is $exp(1.61) \approx 5.00$. In this case, using the complement is a bit more conservative; although the conclusion is not different of course. Furthermore, the evidence ratio for small effect-sizes ($f \leq$ 0.20) does not increase very rapidly (see Figure B1b), independent of sample-size. This is because, when examining small effects, the complement is often true (even though the data were generated under H_1). This is illustrated in Figure B2. For example, if f = 0, the mle's are (except from some sampling variation) in 23/24 (approximately 95.8%) of the time not in agreement with H_1 (and thus in agreement with H_c). Thus, both hypotheses H_c and H_{μ} have the same maximized log-likelihood value with a probability of $\text{prob}_{cu} = 23/24$. When f increases, the data/the mle's will be more and more in agreement with H_1 , and thus not with its complement H_c and hence the proportion of equal maximized log-likelihood values of H_u and H_c (and thus prob_{cu}) decreases. Logically, the proportion of equal maximized log-likelihood values of H_1 and H_{μ} , that is $1 - \text{prob}_{cu}$, then increases.

The presented results so far are for the scenario that H_1 is true, but we are also interested in the performance if H_1 is not true (i.e., H_c is true). Figure B3 shows the results for the situation that the complement is true. Data were generated under the complement of H_1 , for which we choose $H_c: \mu_1 \ge \mu_2 \ge \mu_3 \ge \mu_4$. The means are given in Appendix D and are now in reversed order compared with the previous simulation. Again, we considered the OR hypothesis H_1 , its complement H_c , and the unconstrained hypothesis H_u . The results in Figure B3a show that the mean evidence ratio for H_1 versus H_c (mean w_1/w_c) and for H_1 versus H_u (mean w_1/w_u) decreases rapidly for larger f. This is because, when the effect-size and/or the sample-size increases, the data/mle's will be more and more in agreement with the complement H_c and of course also with the unconstrained hypothesis H_{μ} . The results shown in Figure B3b are based on the same numerical results shown in Figure B3a but now for H_c versus H_1 (mean w_c/w_1) and for H_u versus H_1 (mean w_u/w_1). They clearly show the nice property that if the complement (and also H_{μ}) is true, both evidence ratios w_c/w_1 and w_{μ}/w_{1} show more support for larger effect-sizes and samples sizes. Stated otherwise, both are boundless.

⁵ Note that the ANOVA model is a special case of the multiple regression model discussed in the main text.



Figure B1. Mean of the evidence ratio (on a log scale) for the situation that the OR hypothesis H_1 is true (based on 500 simulations). (a) Hypothesis H_1 is compared with its complement H_c (mean w_1/w_c), for various effect-sizes (f) and for n = 30, 50, 100 and 200. (b) Hypothesis H_1 is compared with its complement H_c (mean w_1/w_c), for various sample-sizes (n) and for f = 0.10, 0.20, 0.30 and 0.40. (c, d) same as (a, b) but now for hypothesis H_1 versus the unconstrained hypothesis H_u (mean w_1/w_u).



Figure B2. Proportion of data sets (based on 500 simulations) that result in equal log-likelihood values for the complement of $H_1: \mu_1 \le \mu_2 \le \mu_3 \le \mu_4$, that is, H_c , and the unconstrained hypothesis H_u (i.e., $LL_c = LL_u$), for various effect-sizes (*f*) and n = 30, 50, 100, 200.



Figure B3. Mean of the evidence ratio (based on 500 simulations) for the situation that the complement H_c of the OR hypothesis H_1 is true, for various effect-sizes (f) and for n = 30 and 200. (a) The closed circles denote H_1 versus H_c (mean w_1/w_c), for n = 30 (solid line) and n = 200 (dashed line). The closed triangles denote H_1 versus the unconstrained hypothesis H_u (mean w_1/w_u). (b) Same as (a) but now for H_c versus H_1 (mean w_c/w_1) and H_u versus H_1 (mean w_u/w_1), for n = 30 and n = 200. Note that Figure B3b is on a log scale.

Appendix C

R Code to Run the Burns Example

In what follows, we describe all steps to compute the evidence ratio for hypothesis H_1 compared with its complement and H_1 compared with the unconstrained hypothesis using the R package **restriktor**.

Step 1: Load your data set into R.

burns <- read.csv("burns.csv", header = TRUE, sep = " ")</pre>

More information about how to load your data into R, can be found online at http://restriktor.org/tutorial/importdata.html.

Step 2: Center the covariates HADS, age, and SO at their average. This can be done in R as follows:

```
burns$Z_HADS <- burns$HADS - mean(burns$HADS, na.rm = TRUE)
burns$Z_age <- burns$age - mean(burns$age, na.rm = TRUE)
burns$Z_S0 <- burns$S0 - mean(burns$S0, na.rm = TRUE)</pre>
```

Step 3: Fit the unconstrained linear regression model using the lm() function.

For clarity reasons, we explicitly added an intercept term by specifying the value 1. The interaction between gender and facial burns is included using the : operator.

Step 4: Create the restriction syntax for restriktor. Now that the model is defined in R, we are left with specifying the order restrictions. This is done in restriktor by specifying a so-called restriction syntax. Order restrictions are defined by means of inequality restrictions (< or >) or by equality restrictions (= =). In addition, a convenient feature of the restriktor restriction syntax is the option to define new parameters that are linear in the original model parameters. This can be done using the := operator. In this way, we can compute the four adjusted means and impose order restrictions among these means. The restriction syntax is enclosed within single quotes. Then, for hypothesis H_3 the restriction syntax might looks as follows:

```
myRestrictions <- '
ml := .Intercept.
m2 := .Intercept. + facialBurns
m3 := .Intercept. + gender
m4 := .Intercept. + facialBurns + gender + gender.facialBurns
m1 < m4
m2 < m4
m3 < m4 '</pre>
```

It is important to note that variable/factor names of the interaction effects in objects of class lm contain a semicolon (:) between the variable names (e.g., gender:facialBurns). To use these parameters in the restriction syntax, the semicolon must be replaced by a dot (.; e.g., gender.facialBurns). In addition, the intercept of a fitted objects of class lm is denoted in the output as (Intercept) and not as 1 anymore. To use the intercept in the restriction syntax, the parentheses must also be replaced by a dot (i.e., .Intercept.). More information about the restriction syntax can be found online at http://restriktor.org/tutorial/syntax.html.

EVALUATING A HYPOTHESIS AGAINST ITS COMPLEMENT

Step 5: Compute the GORIC weights and the evidence ratio using the goric() function from the restriktor package.

The first argument to the goric() function is the unconstrained fitted object of class lm. The second argument is the restriction syntax specified in the previous step. To compare H_3 with its complement H_c , the argument comparison has to be set to "complement" (by default it is set to "unconstrained").

Step 6: Interpret the results. A brief overview can be requested using the print() function. A more detailed overview can requested by the summary() function as follows:

```
summary(out.c, brief = TRUE)
Restriktor: generalized order-restricted information criterion (GORIC):
Results:
```

	model	loglik	penalty	goric	goric.weights
1	myRestrictions	-660.024	6.987	1334.022	0.891
2	Complement	-661.942	7.173	1338.230	0.109

Relative GORIC-weights:

	vs myRestrictions	vs complement
myRestrictions	1.000	8.198
complement	0.122	1.000

```
.
```

The order-restricted hypothesis myRestrictions has 8.198 times more support than its complement.

Below are the results for H_1 compared with the unconstrained hypothesis. out.u $\leq -$ goric(fit.lm,

```
constraints = myRestrictions,
comparison = "unconstrained")
summary(out.u, brief = TRUE)
```

Restriktor: Generalized order-restricted information criterion (GORIC): Results:

	model	loglik	penalty	goric	goric.weights
1	myRestrictions	-660.024	6.987	1334.022	0.734
2	unconstrained	-660.024	8.000	1336.048	0.266

Relative GORIC-weights:

	vs myRestrictions	vs unconstrained
myRestrictions	1.000	2.754
unconstrained	0.363	1.000

Note: In case of equal log-likelihood (loglik) values, the relative weights are solely based on the difference in penalty values.

Appendix D

Population Means for First Simulation Study

Effect-size		Populatio	on means	
f	μ_1	μ_2	μ_3	μ_4
0	0	0	0	0
.1	134	044	.044	.134
.2	268	089	.089	.268
.3	402	134	.134	.402
	:	:	:	:
1	-1.341	447	.447	1.341

Note. In the second simulation, we used the reverse ordering of these means.

Received December 14, 2018 Revision received May 24, 2019 Accepted July 16, 2019 ■

$ \widehat{ P } A = \mathbf{P} S = \mathbf{P} S + \mathbf{P} S = \mathbf{P} S + \mathbf{P} S = \mathbf{P} S $
APA JOURNALS
ORDER INFORMATION
Start my 2020 subscription to Psychological Methods ® ISSN: 1082-989X
PRICINGAPA Member/Affiliate\$101Individual Nonmember\$222Institution\$910
Call 800-374-2721 or 202-336-5600 Fax 202-336-5568 TDD/TTY 202-336-6123
Subscription orders must be prepaid. Subscriptions are on a calendar year basis. Please allow 4-6 weeks for delivery of the first issue.
Learn more and order online at: www.apa.org/pubs/journals/met
Visit at.apa.org/circ2020 to browse APA's full journal collection. All APA journal subscriptions include online first journal articles and access to archives. Individuals can receive online access to all of APA's scholarly journals through a subscription to APA PsycNET®, or through an institutional subscription to the PsycARTICLES® database.