

# ROCnReg: An R Package for Receiver Operating Characteristic Curve Inference with and without Covariates

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**Abstract** This paper introduces the package **ROCnReg** that allows estimating the pooled ROC curve, the covariate-specific ROC curve, and the covariate-adjusted ROC curve by different methods, both from (semi) parametric and nonparametric perspectives and within Bayesian and frequentist paradigms. From the estimated ROC curve (pooled, covariate-specific, or covariate-adjusted), several summary measures of discriminatory accuracy, such as the (partial) area under the ROC curve and the Youden index, can be obtained. The package also provides functions to obtain ROC-based optimal threshold values using several criteria, namely, the Youden index criterion and the criterion that sets a target value for the false positive fraction. For the Bayesian methods, we provide tools for assessing model fit via posterior predictive checks, while model choice can be carried out via several information criteria. Numerical and graphical outputs are provided for all methods. This is the only package implementing Bayesian procedures for ROC curves.

## Introduction

The receiver operating characteristic (ROC) curve (Metz, 1978) is, unarguably, the most popular tool used for evaluating the discriminatory ability of continuous-outcome diagnostic tests. The ROC curve displays the false positive fraction (FPF) against the true positive fraction (TPF) for all possible threshold values that can be used to dichotomise the test result. The ROC curve thus provides a global description of the trade-off between the FPF and the TPF of the test as the threshold changes. Plenty of parametric and semi/nonparametric methods are available for estimating the ROC curve, either from frequentist or Bayesian viewpoints and we refer the interested reader to Pepe (1998, Chapter 5), Zhou et al. (2011, Chapter 4), Inácio et al. (2020), and references therein.

It is known that in many situations, the outcome of a test and, possibly, its discriminatory capacity can be affected by covariates. Two different ROC-based measures that incorporate covariate information have been proposed: the covariate-specific or conditional ROC curve (see, e.g., Pepe, 2003, Chapter 6) and the covariate-adjusted ROC curve (Janes and Pepe, 2009). The formal definition of both curves is given in Section [Notation and definitions](#). Succinctly, a covariate-specific ROC curve is an ROC curve that conditions on a specific covariate value, thus describing the accuracy of the test in the ‘subpopulation’ defined by that covariate value. On the other hand, the covariate-adjusted ROC curve is a weighted average of covariate-specific ROC curves. Regarding estimation, since the seminal paper of Pepe (1998), a plethora of methods have been proposed in the literature for the estimation of the covariate-specific ROC curve and associated summary measures. Without being exhaustive, we mention the work of Faraggi (2003), Rodríguez-Álvarez et al. (2011a,b), Inácio de Carvalho et al. (2013), and Inácio de Carvalho et al. (2017). A detailed review can be found in Rodríguez-Álvarez et al. (2011c), Pardo-Fernández et al. (2014) and Inácio et al. (2020). With respect to the covariate-adjusted ROC curve, estimation has been discussed in Janes and Pepe (2009), Rodríguez-Álvarez et al. (2011a), Guan et al. (2012), and Inácio de Carvalho and Rodríguez-Álvarez (2018).

There are a few R packages for ROC curve analysis available on the Comprehensive R Archive Network and, as far as we are aware, all of them implementing frequentist approaches. The package **sROC** (Wang, 2012) contains functions to perform nonparametric, kernel-based, estimation of ROC curves, **pROC** (Robin et al., 2011) offers a set of tools to visualise, smooth, and compare ROC curves, and **nsROC** (Pérez Fernández et al., 2018) also allows estimating ROC curves, building confidence bands, as well as comparing several curves both for dependent and independent data (i.e., data arising from paired and unpaired study designs, respectively). However, covariate information cannot be explicitly taken into account in any of these packages. The packages **ROCRegression** (available at <https://bitbucket.org/mxrodriguez/rocregression>) and **npROCRegression** (Rodríguez-Álvarez and Roca-Pardinas, 2017) provide routines to estimate semiparametrically and nonparametrically, under a frequentist framework, the covariate-specific ROC curve. We also mention **OptimalCutpoints** (López-Ratón et al., 2014) and **ThresholdROC** (Perez Jaume et al., 2017) that provide a collection of functions for point and interval estimation of optimal thresholds for continuous diagnostic tests. To the best of our knowledge, there is no statistical software package implementing Bayesian inference for ROC curves and associated summary indices and optimal thresholds.

To close this gap, in this paper we introduce the **ROCnReg** package that allows conducting

Method	Description
<b>Pooled ROC curve</b>	
emp	(Frequentist) empirical estimator ( <a href="#">Hsieh and Turnbull, 1996</a> ).
kernel	(Frequentist) kernel-based approach ( <a href="#">Zou et al., 1997</a> ).
BB	Bayesian bootstrap method ( <a href="#">Gu et al., 2008</a> ).
dpm	Nonparametric Bayesian approach based on a Dirichlet process mixture of normal distributions ( <a href="#">Erkanli et al., 2006</a> ).
<b>Covariate-specific ROC curve</b>	
sp	(Frequentist) parametric and semiparametric induced ROC regression approach ( <a href="#">Pepe, 1998; Faraggi, 2003</a> )
kernel	Nonparametric (kernel-based) induced ROC regression approach ( <a href="#">Rodríguez-Álvarez et al., 2011a</a> ).
bnp	Nonparametric Bayesian model based on a single-weights dependent Dirichlet process mixture of normal distributions ( <a href="#">Inácio de Carvalho et al., 2013</a> ).
<b>Covariate-adjusted ROC curve</b>	
sp	(Frequentist) semiparametric method ( <a href="#">Janes and Pepe, 2009</a> ).
kernel	Nonparametric (kernel-based) induced ROC regression approach ( <a href="#">Rodríguez-Álvarez et al., 2011a</a> ).
bnp	Nonparametric Bayesian model based on a single-weights dependent Dirichlet process mixture of normal distributions and the Bayesian bootstrap ( <a href="#">Inácio de Carvalho and Rodríguez-Álvarez, 2018</a> ).

**Table 1:** Overview of ROC estimation methods included in the **ROCnReg** package.

Bayesian inference for the (pooled or marginal) ROC curve, the covariate-specific ROC curve, and the covariate-adjusted ROC curve. For the sake of generality, frequentist approaches are also implemented. Specifically, in what concerns estimation of the pooled ROC curve, **ROCnReg** implements the frequentist empirical estimator described in [Hsieh and Turnbull \(1996\)](#), the kernel-based approach proposed by [Zou et al. \(1997\)](#), the Bayesian Bootstrap method of [Gu et al. \(2008\)](#), and the Bayesian nonparametric method based on a Dirichlet process mixture of normal distributions model proposed by [Erkanli et al. \(2006\)](#). Regarding the covariate-specific ROC curve, **ROCnReg** implements the frequentist normal method of [Faraggi \(2003\)](#) and its semiparametric counterpart as described in [Pepe \(1998\)](#), the kernel-based approach of [Rodríguez-Álvarez et al. \(2011a\)](#), and the Bayesian nonparametric model, based on a single-weights dependent Dirichlet process mixture of normal distributions, proposed by [Inácio de Carvalho et al. \(2013\)](#). As for the covariate-adjusted ROC curve, the **ROCnReg** package allows estimation using the frequentist semiparametric approach of [Janes and Pepe \(2009\)](#), the frequentist nonparametric method discussed in [Rodríguez-Álvarez et al. \(2011a\)](#), and the recently proposed Bayesian nonparametric estimator of [Inácio de Carvalho and Rodríguez-Álvarez \(2018\)](#). Table 1 shows a summary of all methods implemented in the package. In addition, **ROCnReg** also provides functions to obtain ROC-based optimal thresholds to perform the classification/diagnosis of individuals as, say, diseased or nondiseased, using two different criteria, namely, the Youden index and the criterion that sets a target value for the false positive fraction. These are implemented for both the ROC curve, the covariate-specific, and the covariate-adjusted ROC curve.

The remainder of the paper is organised as follows. In Section [Notation and definitions](#) we formally introduce the (pooled or marginal) ROC curve, the covariate-specific ROC curve, and the covariate-adjusted ROC curve. The description of the Bayesian estimation methods implemented in the **ROCnReg** package is given in Section [Methods](#). In Section [Package presentation and illustration](#) the usage of the main functions and capabilities of **ROCnReg** are described and illustrated using a synthetic dataset mimicking endocrine data. The paper concludes with a discussion in Section [Summary and future plans](#).

## Notation and definitions

This section sets out the formal definition of the pooled or marginal ROC curve, the covariate-specific ROC curve, and the covariate-adjusted ROC curve. It also describes the most commonly used summary measures of discriminatory accuracy, namely, the area under the ROC curve, the partial area under the ROC curve, and the Youden Index. For conciseness, we intentionally avoid giving too many details

and refer the interested reader to Pepe (2003) (and references therein) for an extensive account of many aspects of ROC curves with and without covariates.

In what follows, we denote as  $Y$  the outcome of the diagnostic test and as  $D$  the binary variable indicating the presence ( $D = 1$ ) or absence ( $D = 0$ ) of disease. We also assume that along with  $Y$  and the true disease status  $D$ , a covariate vector  $X$  is also available, and that it may encompass both continuous and categorical covariates. For ease of notation, the covariate vector  $X$  is assumed to be the same in both the diseased ( $D = 1$ ) and nondiseased ( $D = 0$ ) populations, although this is not always necessarily the case in practice (e.g., disease stage is, obviously, a disease-specific covariate). By a slight abuse of notation, we use the subscripts  $D$  and  $\bar{D}$  to denote (random) quantities conditional on, respectively,  $D = 1$  and  $D = 0$ . For example,  $Y_D$  and  $Y_{\bar{D}}$  denote the test outcomes in the diseased and nondiseased populations, respectively.

### Pooled ROC curve

In the case of a continuous-outcome diagnostic test, the classification is usually made by comparing the test result  $Y$  against a threshold  $c$ . If the outcome is equal or above the threshold,  $Y \geq c$ , the subject will be diagnosed as diseased. On the other hand, if the test result is below the threshold,  $Y < c$ , he or she will be classified as nondiseased. The ROC curve is then defined as the set of all possible pairs of false positive fractions,  $\text{FPF}(c) = \Pr(Y \geq c | D = 0) = \Pr(Y_{\bar{D}} \geq c)$ , and true positive fractions,  $\text{TPF}(c) = \Pr(Y \geq c | D = 1) = \Pr(Y_D \geq c)$ , that can be obtained by varying the threshold value  $c$ , i.e.,

$$\{(\text{FPF}(c), \text{TPF}(c)) : c \in \mathbb{R}\}.$$

It is common to represent the ROC curve as  $\{(p, \text{ROC}(p)) : p \in [0, 1]\}$ , where

$$p = \text{FPF}(c) = 1 - F_{\bar{D}}(c), \quad \text{ROC}(p) = 1 - F_D \left\{ F_{\bar{D}}^{-1}(1 - p) \right\}, \quad (1)$$

with  $F_{\bar{D}}(y) = \Pr(Y_{\bar{D}} \leq y)$  and  $F_D(y) = \Pr(Y_D \leq y)$  denoting the cumulative distribution function (CDF) of  $Y$  in the nondiseased and diseased groups, respectively. Several indices can be used as global summary measures of the accuracy of a test. The most widely used is the area under the ROC curve (AUC), defined as

$$\text{AUC} = \int_0^1 \text{ROC}(p) dp. \quad (2)$$

In addition to its geometric definition, the AUC has also a probabilistic interpretation (see, e.g., Pepe, 2003, p. 78)

$$\text{AUC} = \Pr(Y_D \geq Y_{\bar{D}}), \quad (3)$$

that is, the AUC is the probability that a randomly selected diseased subject has a higher test outcome than that of a randomly selected nondiseased subject. The AUC takes values between 0.5, in the case of an uninformative test that classifies individuals no better than chance, and 1.0 for a perfect test. We note that an AUC below 0.5 simply means that the classification rule should be reversed. As it is clear from its definition, the AUC integrates the ROC curve over the whole range of FPFs. However, depending on the clinical circumstances, interest might lie only on a relevant interval of FPFs or TPFs, which leads to the notion of partial area under the ROC curve (pAUC). The pAUC over a range of FPFs  $(0, u_1)$ , where  $u_1$  is typically low and represents the largest acceptable FPF, is defined as

$$\text{pAUC}(u_1) = \int_0^{u_1} \text{ROC}(p) dp. \quad (4)$$

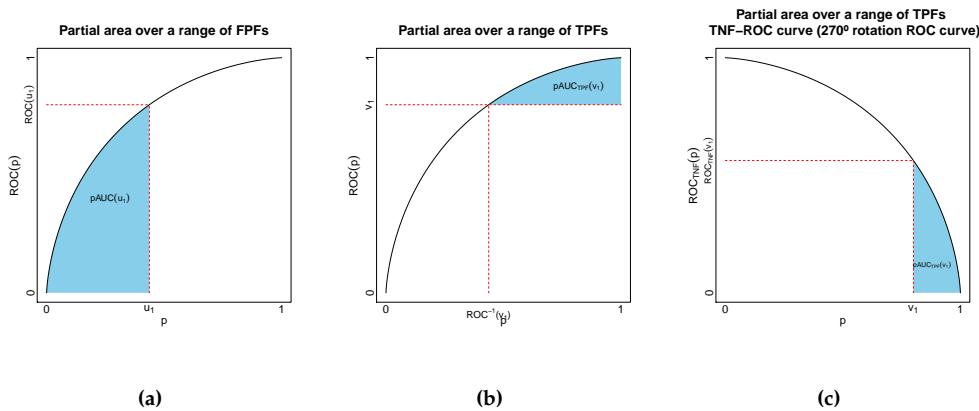
On the other hand, the pAUC over a range of TPFs  $(v_1, 1)$ , where  $v_1$  is typically large and represents the lowest acceptable TPF, is defined as

$$\text{pAUC}_{\text{TPF}}(v_1) = \int_{v_1}^1 \text{ROC}_{\text{TNF}}(p) dp, \quad (5)$$

where  $\text{ROC}_{\text{TNF}}$  is a  $270^\circ$  rotation of the ROC curve, which can be expressed as

$$\text{ROC}_{\text{TNF}}(p) = F_{\bar{D}} \{ F_D^{-1}(1 - p) \}. \quad (6)$$

The curve in (6) is referred to as the true negative fraction (TNF) ROC curve, since TNF ( $= 1 - \text{FPF}$ ) is plotted on the  $y$ -axis. We shall highlight that the argument  $p$  in the ROC curve stands for a false positive fraction, whereas in the  $\text{ROC}_{\text{TNF}}$  curve it stands for a true positive fraction. In Figure 1 we graphically illustrate the two partial areas.



**Figure 1:** (a) Shaded area in blue represents the partial area under the ROC curve over the interval  $(0, u_1)$  of FPFs. (b) Shaded area in blue represents the partial area under the ROC curve over the interval  $(v_1, 1)$  of TPFs. (c) The same as in (b) but now represented as an area under the true negative fraction ROC curve.

Another summary index of diagnostic accuracy is the Youden index ([Shapiro, 1999](#); [Youden, 1950](#))

$$YI = \max_c \{TPF(c) - FPF(c)\} \quad (7)$$

$$= \max_c \{F_{\bar{D}}(c) - F_D(c)\} \quad (8)$$

$$= \max_p \{ROC(p) - p\}. \quad (9)$$

The YI ranges from 0 to 1, taking the value of 0 in the case of an uninformative test and 1 for a perfect test. As for the AUC, a YI below 0 means that the classification rule should be reversed. The value  $c^*$  which maximises Equation (7) (or, equivalently, Equation (8)) is frequently used in practice to classify subjects as diseased or nondiseased. It should be noted that the Youden index is equivalent to the Kolmogorov-Smirnov measure of distance between the distributions of  $Y_D$  and  $Y_{\bar{D}}$  ([Pepe, 2003](#), p. 80).

### Covariate-specific ROC curve

The conditional or covariate-specific ROC curve, given a covariate value  $\mathbf{x}$ , is defined as

$$ROC(p | \mathbf{x}) = 1 - F_{\bar{D}}\{F_D^{-1}(1 - p | \mathbf{x}) | \mathbf{x}\}, \quad (10)$$

where  $F_{\bar{D}}(y | \mathbf{x}) = \Pr(Y_{\bar{D}} \leq y | \mathbf{X}_{\bar{D}} = \mathbf{x})$  and  $F_D(y | \mathbf{x}) = \Pr(Y_D \leq y | \mathbf{X}_D = \mathbf{x})$  are the conditional CDFs of the test in the nondiseased and diseased groups, respectively. In this case, a number of possibly different ROC curves (and therefore discriminatory accuracies) may be obtained for different values of  $\mathbf{x}$ . Thus, the covariate-specific ROC curve is an important tool that helps to understand and determine the optimal and suboptimal populations where to apply the tests on. That is, the covariate-specific ROC curve allows determining the populations, defined by  $\mathbf{x}$ , homogeneous with respect to  $\mathbf{x}$ , where the diagnostic test has a ‘good’ or ‘poor’ discriminatory capacity. Similarly to the unconditional case, the covariate-specific TNF-ROC curve is given by

$$ROC_{TNF}(p | \mathbf{x}) = F_{\bar{D}}\{F_D^{-1}(1 - p | \mathbf{x}) | \mathbf{x}\}, \quad (11)$$

and the covariate-specific AUC, pAUC, and Youden index are

$$\text{AUC}(\mathbf{x}) = \int_0^1 \text{ROC}(p | \mathbf{x}) dp, \quad (12)$$

$$\text{pAUC}(u_1 | \mathbf{x}) = \int_0^{u_1} \text{ROC}(p | \mathbf{x}) dp, \quad (13)$$

$$\text{pAUC}_{\text{TPF}}(v_1 | \mathbf{x}) = \int_{v_1}^1 \text{ROC}_{\text{TNF}}(p | \mathbf{x}) dp, \quad (14)$$

$$\text{YI}(\mathbf{x}) = \max_c |\text{TPF}(c | \mathbf{x}) - \text{FPF}(c | \mathbf{x})| \quad (15)$$

$$= \max_c |F_{\bar{D}}(c | \mathbf{x}) - F_D(c | \mathbf{x})| \quad (16)$$

$$= \max_p |\text{ROC}(p | \mathbf{x}) - p|. \quad (17)$$

The value  $c_x^*$  that achieves the maximum in (15) (or (16)) is called the optimal covariate-specific YI threshold and can be used to classify a subject, with covariate value  $\mathbf{x}$ , as diseased or nondiseased.

### Covariate-adjusted ROC curve

The covariate-specific ROC curve and associated AUC, pAUCs, and YI described in the previous section depict the accuracy of the test for specific covariate values. However, it would be undoubtedly useful to have a global summary measure that also takes covariate information into account. Such summary measure was developed by [Janes and Pepe \(2009\)](#), who proposed the covariate-adjusted ROC (AROC) curve, defined as

$$\text{AROC}(p) = \int \text{ROC}(p | \mathbf{x}) dH_D(\mathbf{x}), \quad (18)$$

where  $H_D(\mathbf{x}) = \Pr(\mathbf{X}_D \leq \mathbf{x})$  is the CDF of  $\mathbf{X}_D$ . That is, the AROC curve is a weighted average of covariate-specific ROC curves, weighted according to the distribution of the covariates in the diseased group. Equivalently, as shown by [Janes and Pepe \(2009\)](#), the AROC curve can also be expressed as

$$\begin{aligned} \text{AROC}(p) &= \Pr\{Y_D > F_{\bar{D}}^{-1}(1-p | \mathbf{X}_D)\} \\ &= \Pr\{1 - F_{\bar{D}}(Y_D | \mathbf{X}_D) \leq p\}. \end{aligned} \quad (19)$$

As will be seen in Section [Methods](#), Expression (19) is very convenient when it comes to estimating the AROC curve. Also, it emphasises that the AROC curve at a FPF of  $p$  is the overall TPF when the thresholds used for defining a positive test result are covariate-specific and chosen to ensure that the FPF is  $p$  in each subpopulation defined by the covariate values.

In contrast to the pooled ROC curve (see Expressions (1) and (6)) and the covariate-specific ROC curve (see Expressions (10) and (11)), the AROC curve (and its 270° rotation) cannot be expressed in terms of the (conditional) CDFs of the test in each group. This does not, however, preclude the possibility of defining AROC-based summary accuracy measures, yet more care is needed. Thus, for the AROC curve, the area under the AROC, as well as the partial areas and YI are expressed as follows

$$\text{AAUC} = \int_0^1 \text{AROC}(p) dp, \quad (20)$$

$$\text{pAAUC}(u_1) = \int_0^{u_1} \text{AROC}(p) dp, \quad (21)$$

$$\text{pAAUC}_{\text{TPF}}(v_1) = \int_{\text{AROC}^{-1}(v_1)}^1 \text{AROC}(p) dp - \{1 - \text{AROC}^{-1}(v_1)\} v_1, \quad (22)$$

$$\text{YI}_{\text{AROC}} = \max_p \{\text{AROC}(p) - p\}. \quad (23)$$

Note, in particular, that the expressions for both the partial area under the AROC curve over a range of TPFs (see also Figure 1b) and for the YI are defined in terms of the AROC curve. For the YI, once the value that achieves the maximum in (23) is obtained, say  $p^*$ , covariate-specific threshold values can be calculated as follows

$$c_x^* = F_{\bar{D}}^{-1}(1 - p^* | \mathbf{X}_D = \mathbf{x}).$$

Note that, by construction, these threshold values will ensure that the FPF is  $p^*$  in each subpopulation defined by the covariate values; however, the TPF may vary with the covariate values, i.e.,

$$\text{TPF}(c_x^*) = 1 - F_D(c_x^* | \mathbf{X}_D = \mathbf{x}).$$

To finish this part, we mention that when the accuracy of a test is not affected by covariates, this does not necessarily mean that the covariate-specific ROC curve (which in this case is the same for all covariate values) coincides with the pooled ROC curve. It does coincide, however, with the AROC curve (see [Janes and Pepe, 2009](#); [Pardo-Fernández et al., 2014](#); [Inácio de Carvalho and Rodríguez-Álvarez, 2018](#), for more details). As such, in all cases where covariates affect the test results, even though they might not affect its discriminatory capacity, inferences based on the pooled ROC curve might be misleading. In such cases, the AROC curve should be used instead. This also applies to the selection of (optimal) threshold values, which might be covariate-specific (i.e., possibly different for different covariate values).

## Methods

For space reasons, we focus ourselves here on the Bayesian methods for ROC curve inference (with and without covariates) implemented in the **ROCNReg** package. A detailed description, as well as usage examples, of the frequentist approaches are available as Supplementary Material at <https://bitbucket.org/mxrodriguez/rocnreg>.

### Pooled ROC curve

In what follows, let  $\{y_{\bar{D}i}\}_{i=1}^{n_{\bar{D}}}$  and  $\{y_{Dj}\}_{j=1}^{n_D}$  be two independent random samples of test outcomes from the nondiseased and diseased groups of size  $n_{\bar{D}}$  and  $n_D$ , respectively.

#### *Bayesian bootstrap based estimator*

The function `pooledROC.bb` implements the Bayesian bootstrap (BB) approach proposed by [Gu et al. \(2008\)](#). Their estimator relies on the notion of placement value ([Pepe, 2003](#), Chapter 5), which is simply a standardisation of the test outcomes with respect to a reference group. Specifically,  $U_D = 1 - F_{\bar{D}}(Y_D)$  is to be interpreted as a standardisation of a diseased test outcome with respect to the distribution of test results in the nondiseased population. The ROC curve can be regarded as the CDF of  $U_D$

$$\Pr(U_D \leq p) = \Pr\{1 - F_{\bar{D}}(Y_D) \leq p\} = 1 - F_{\bar{D}}\{F_{\bar{D}}^{-1}(1 - p)\} = \text{ROC}(p), \quad 0 \leq p \leq 1. \quad (24)$$

This representation in (24) of the ROC curve provided the rationale for the two-step algorithm of [Gu et al. \(2008\)](#), which can be described as follows. Let  $S$  be the number of iterations.

**Step 1: Computation of the placement value based on the BB.**

For  $s = 1, \dots, S$ , let

$$U_{Dj}^{(s)} = \sum_{i=1}^{n_{\bar{D}}} q_{1i}^{(s)} I(y_{\bar{D}i} \geq y_{Dj}), \quad j = 1, \dots, n_D,$$

where  $(q_{11}^{(s)}, \dots, q_{1n_{\bar{D}}}^{(s)}) \sim \text{Dirichlet}(n_{\bar{D}}; 1, \dots, 1)$ .

**Step 2: Generate a realisation of the ROC curve.** Based on (24), generate a realisation of  $\text{ROC}^{(s)}(p)$ , the cumulative distribution function of  $(U_{D1}^{(s)}, \dots, U_{Dn_D}^{(s)})$ , where

$$\text{ROC}^{(s)}(p) = \sum_{j=1}^{n_D} q_{2j}^{(s)} I(U_{Dj}^{(s)} \leq p), \quad (q_{21}^{(s)}, \dots, q_{2n_D}^{(s)}) \sim \text{Dirichlet}(n_D; 1, \dots, 1).$$

The BB estimate of the ROC curve is obtained by averaging over the ensemble of ROC curves  $\{\text{ROC}^{(1)}(p), \dots, \text{ROC}^{(S)}(p)\}$ , that is,

$$\widehat{\text{ROC}}^{\text{BB}}(p) = \frac{1}{S} \sum_{s=1}^S \text{ROC}^{(s)}(p),$$

and a  $(1 - \alpha) \times 100\%$  pointwise credible band can be obtained from the  $\alpha/2 \times 100\%$  and  $(1 - \alpha/2) \times 100\%$  percentiles of the same ensemble ( $\alpha \in (0, 1)$ ). Note that these pointwise credible bands for the ROC curve are to be interpreted as credible intervals for the corresponding constituents TPFs.

The Bayesian bootstrap estimator leads to closed-form expressions for the AUC and pAUC, which

are, respectively, given by

$$\begin{aligned}\text{AUC}^{(s)} &= \int_0^1 \text{ROC}^{(s)}(p) dp = 1 - \sum_{j=1}^{n_D} q_{2j}^{(s)} U_{Dj}^{(s)}, \\ \text{pAUC}^{(s)}(u_1) &= \int_0^{u_1} \text{ROC}^{(s)}(p) dp = u_1 - \sum_{j=1}^{n_D} q_{2j}^{(s)} \min \left\{ u_1, U_{Dj}^{(s)} \right\}.\end{aligned}$$

It is easy to show that

$$\text{pAUC}_{\text{TPF}}^{(s)}(v_1) = \int_{v_1}^1 \text{ROC}_{\text{TNF}}^{(s)}(p) dp = \sum_{i=1}^{n_{\bar{D}}} q_{1i}^{(s)} \max \left\{ v_1, U_{\bar{D}i}^{(s)} \right\} - v_1,$$

where

$$U_{\bar{D}i}^{(s)} = \sum_{j=1}^{n_D} q_{2j}^{(s)} I \left( y_{Dj} \geq y_{\bar{D}i} \right), \quad i = 1, \dots, n_{\bar{D}},$$

and it is also easy to demonstrate that the  $\text{ROC}_{\text{TNF}}$  curve is the survival function of the placement value  $U_{\bar{D}} = 1 - F_{\bar{D}}(Y_{\bar{D}})$ . With respect to the Youden index, it is obtained by maximising, over a grid of possible threshold values, the following expression

$$\text{YI}^{(s)} = \max_c \left\{ F_{\bar{D}}^{(s)}(c) - F_{\bar{D}}^{(s)}(c) \right\},$$

where

$$F_{\bar{D}}^{(s)}(c) = \sum_{i=1}^{n_{\bar{D}}} q_{1i}^{(s)} I(y_{\bar{D}i} \leq c) \quad \text{and} \quad F_{\bar{D}}^{(s)}(c) = \sum_{j=1}^{n_D} q_{2j}^{(s)} I(y_{Dj} \leq c).$$

As for the ROC curve, point estimates for the AUC, pAUC,  $\text{pAUC}_{\text{TPF}}$ , YI, and  $c^*$  can be obtained by averaging over the respective ensembles of  $S$  realisations, with credible bands derived from the percentiles of such ensembles.

### *Dirichlet process mixture of normal distributions based estimator*

The Bayesian nonparametric approach, based on a Dirichlet process mixture (DPM) of normal distributions, for estimating the pooled ROC curve (Erkanli et al., 2006) is implemented in the `pooledROC.dpm` function. In this case, as implicit by the name, the CDFs of the test outcomes in each group are estimated via a Dirichlet process mixture of normal distributions, that is, it is assumed that the CDF, say in the diseased group (the one in the nondiseased group,  $\bar{D}$ , follows analogously), is of the form

$$F_D(y) = \int \Phi(y | \mu, \sigma^2) dG_D(\mu, \sigma^2), \quad G_D \sim \text{DP}(\alpha_D, G_D^*(\mu, \sigma^2)), \quad (25)$$

where  $\Phi(y | \mu, \sigma^2)$  denotes the CDF of the normal distribution with mean  $\mu$  and variance  $\sigma^2$  evaluated at  $y$ . Here  $G_D \sim \text{DP}(\alpha_D, G_D^*)$  is used to denote that the mixing distribution  $G_D$  follows a Dirichlet process (DP) (Ferguson, 1973) with centring distribution  $G_D^*$ , for which  $E(G_D) = G_D^*$ , and precision parameter  $\alpha_D$ . Usually, due to conjugacy reasons,  $G_D^*(\mu, \sigma^2) \equiv N(\mu | m_{D0}, S_{D0})\Gamma(\sigma^{-2} | a_D, b_D)$  and this is the centring distribution used by the `pooledROC.dpm` function. Note that here  $S_{D0}$  denotes the variance of the normal distribution and  $a_D$  and  $b_D$  are, respectively, the shape and rate parameters of the gamma distribution. All hyperparameter values are fixed.

For ease of posterior simulation and because it provides a highly accurate approximation, we make use of the truncated stick-breaking representation of the DP (Ishwaran and James, 2001), according to which  $G_D$  can be written as

$$G_D(\cdot) = \sum_{l=1}^{L_D} \omega_{Dl} \delta_{(\mu_{Dl}, \sigma_{Dl}^2)}(\cdot),$$

where  $(\mu_{Dl}, \sigma_{Dl}^2) \stackrel{\text{iid}}{\sim} G_D^*(\mu, \sigma^2)$ , for  $l = 1, \dots, L_D$ , and the weights follow the so-called (truncated) stick-breaking construction:  $\omega_{D1} = v_{D1}$ ,  $\omega_{Dl} = v_{Dl} \prod_{r < l} (1 - v_{Dr})$ ,  $l = 2, \dots, L_D$ , and  $v_{D1}, \dots, v_{D,L_D-1} \stackrel{\text{iid}}{\sim} \text{Beta}(1, \alpha_D)$ . Further, one must set  $v_{DL_D} = 1$  in order to ensure that the weights add up to one. The CDF in (25) can therefore be written as

$$F_D(y) = \sum_{l=1}^{L_D} \omega_{Dl} \Phi(y | \mu_{Dl}, \sigma_{Dl}^2),$$

where we shall note that  $L_D$  is not the exact number of components expected to be observed, but rather

an upper bound on it, as some of the components may be unoccupied. Some comments are in order regarding the specification of the hyperparameters' values. In what concerns the centring distribution,  $m_{D0}$  represents the prior belief about the components' means and  $S_{D0}$  represents the confidence in such prior belief. Similarly, the values of  $a_D$  and  $b_D$  can be chosen to represent prior belief about the components' variance. Of course, when setting these parameters, it is crucial to consider the measurement scale of the data. By default, test outcomes are standardised (so that the resulting mean is zero and the variance is one) in the `pooledROC.dpm` function and the default values are as follows

$$m_{D0} = 0, \quad S_{D0} = 10, \quad a_D = 2, \quad b_D = 0.5.$$

Because test outcomes are standardised we expect the means of the components to be near zero and hence why  $m_{D0} = 0$ . The parameter  $S_{D0}$  then controls where the drawn  $\mu_{Dl}$  can lie and the value of 10 implies that approximately 95% of the values roughly lie within  $-6$  and  $6$ . Further, note that  $a_D = 2$  leads to a prior with infinite variance that is centred around a finite mean ( $b_D = 0.5$ ) and therefore favours variances less than one. Considering that the standardised data have a variance of one, it is reasonable to expect the within component variance to be smaller than the overall variance. The option of not standardising the test outcomes is also available in `pooledROC.dpm` and in such a case the defaults for the centring distribution hyperparameters' values are as following

$$m_{D0} = \bar{y}_D, \quad S_{D0} = 100s_D^2/n_D, \quad a_D = 2, \quad b_D = s_D^2/2,$$

with  $\bar{y}_D = \frac{1}{n_D} \sum_{j=1}^{n_D} y_{Dj}$  and  $s_D^2 = \frac{1}{n_D-1} \sum_{j=1}^{n_D} (y_{Dj} - \bar{y}_D)^2$ . Regarding the precision parameter of the DP,  $\alpha_D$ , it has a direct relationship with the number of occupied mixture components. One possible strategy for specifying  $\alpha_D$  is to fix it to a small value to favour a small number of occupied components relative to the sample size. In the `pooledROC.dpm` function we set  $\alpha_D = 1$ , a commonly used default value (Gelman et al., 2013, p. 553). Lastly, by default  $L_D = 10$ . Before proceeding we shall emphasise that these two configurations of hyperparameters values (for standardised and not standardised test outcomes) have proved to work well for a different range of test outcomes distributions but it is certainly not our goal to encourage users to use it blindly and indeed thought should be dedicated to this important task. Nevertheless, output from the function `pooledROC.dpm` may be post processed and (informal) model fit diagnostics obtained; see more in Section [Package presentation and illustration](#) and in the Supplementary Materials.

Because the full conditional distributions for all model parameters are available in closed-form, posterior simulation can be easily conducted through Gibbs sampler (see the details, for instance, in Ishwaran and James 2002). At iteration  $s$  of the Gibbs sampler procedure, the ROC curve is computed as

$$\text{ROC}^{(s)}(p) = 1 - F_D^{(s)} \left\{ F_{\bar{D}}^{-1}(s)(1-p) \right\}, \quad s = 1, \dots, S,$$

with

$$F_D^{(s)}(y) = \sum_{l=1}^{L_D} \omega_{Dl}^{(s)} \Phi \left( y \mid \mu_{Dl}^{(s)}, \sigma_{Dl}^{2(s)} \right), \quad F_{\bar{D}}^{(s)}(y) = \sum_{k=1}^{L_{\bar{D}}} \omega_{\bar{D}k}^{(s)} \Phi \left( y \mid \mu_{\bar{D}k}^{(s)}, \sigma_{\bar{D}k}^{2(s)} \right), \quad (26)$$

and where the inversion is performed numerically. There is a closed-form expression for the AUC (Erkanli et al., 2006) given by

$$\text{AUC}^{(s)} = \sum_{k=1}^{L_{\bar{D}}} \sum_{l=1}^{L_D} \omega_{\bar{D}k}^{(s)} \omega_{Dl}^{(s)} \Phi \left( \frac{b_{kl}^{(s)}}{\sqrt{1 + a_{kl}^{2(s)}}} \right), \quad b_{kl}^{(s)} = \frac{\mu_{Dl}^{(s)} - \mu_{\bar{D}k}^{(s)}}{\sigma_{Dl}^{(s)}}, \quad a_{kl}^{(s)} = \frac{\sigma_{\bar{D}k}^{(s)}}{\sigma_{Dl}^{(s)}}.$$

Also, when  $L_D = L_{\bar{D}} = 1$ , there are closed-form expressions for the pAUC and pAUC<sub>TPF</sub> which are used in the package (see Hillis and Metz, 2012). For the pAUC/pAUC<sub>TPF</sub>, when  $L_D > 1$  or  $L_{\bar{D}} > 1$ , the integrals are approximated numerically using Simpson's rule. The Youden index/optimal threshold is computed as in the Bayesian bootstrap method, with the obvious difference that here the CDFs are expressed as in (26). At the end of the sampling procedure, we have an ensemble of  $S$  ROC curves and AUCs/pAUCs/pAUC<sub>TPF</sub>s/YIs/optimal thresholds which, as before, allows obtaining point and interval estimates.

### Covariate-specific ROC curve

We now let  $\{(x_{\bar{D}i}, y_{\bar{D}i})\}_{i=1}^{n_{\bar{D}}}$  and  $\{(x_{Dj}, y_{Dj})\}_{j=1}^{n_D}$  be two independent random samples of test outcomes and covariates from the nondiseased and diseased groups of size  $n_{\bar{D}}$  and  $n_D$ , respectively. Further, for all  $i = 1, \dots, n_{\bar{D}}$  and  $j = 1, \dots, n_D$ , let  $x_{\bar{D}i} = (x_{\bar{D}i,1}, \dots, x_{\bar{D}i,q})^\top$  and  $x_{Dj} = (x_{Dj,1}, \dots, x_{Dj,q})^\top$  be  $q$ -dimensional vectors of covariates, which can be either continuous or categorical.

The function `cROC.bnp` implements the Bayesian nonparametric approach for conducting inference

about the covariate-specific ROC curve of [Inácio de Carvalho et al. \(2013\)](#), which is based on a single-weights dependent Dirichlet process mixture of normal distributions ([De Iorio et al., 2009](#)). Specifically, under this method, the conditional CDF in the diseased group is modelled as follows

$$F_D(y_{Dj} | \mathbf{x}_{Dj}) = \int \Phi(y_{Dj} | \mu_D(\mathbf{x}_{Dj}, \boldsymbol{\beta}), \sigma^2) dG_D(\boldsymbol{\beta}, \sigma^2), \quad G_D \sim DP(\alpha_D, G_D^*(\boldsymbol{\beta}, \sigma^2)),$$

with the conditional CDF in the nondiseased group  $\bar{D}$  following in an analogous manner. As in the no-covariate case, by making use of Sethuraman's truncated representation of the DP, we can write the conditional CDF as

$$\begin{aligned} F_D(y_{Dj} | \mathbf{x}_{Dj}) &= \sum_{l=1}^{L_D} \omega_{Dl} \Phi(y_{Dj} | \mu_D(\mathbf{x}_{Dj}, \boldsymbol{\beta}_{Dl}), \sigma_{Dl}^2), \\ \omega_{D1} &= v_{D1}, \quad \omega_{Dl} = v_{Dl} \prod_{r < l} (1 - v_{Dr}), \quad l = 2, \dots, L_D, \\ v_{Dl} &\stackrel{iid}{\sim} \text{Beta}(1, \alpha_D), \quad l = 1, \dots, L_D - 1, \quad v_{DL_D} = 1. \end{aligned}$$

It is worth mentioning that although the variance of each component does not depend on covariates, the overall variance of the mixture does depend on covariates as it can be written as

$$\text{var}(y_{Dj} | \mathbf{x}_{Dj}) = \sum_{l=1}^{L_D} \omega_{Dl} \sigma_{Dl}^2 + \sum_{l=1}^{L_D} \omega_{Dl} \left\{ \mu_D(\mathbf{x}_{Dj}, \boldsymbol{\beta}_{Dl}) - \left( \sum_{l=1}^{L_D} \omega_{Dl} \mu_D(\mathbf{x}_{Dj}, \boldsymbol{\beta}_{Dl}) \right)^2 \right\}.$$

Note that by assuming that the weights,  $w_{Dl}$ , do not vary with covariates, the model might have limited flexibility in practice ([MacEachern, 2000](#)). This issue can, however, be largely mitigated by using a flexible formulation for  $\mu_D(\mathbf{x}_{Dj}, \boldsymbol{\beta}_{Dl})$ , which is needed not only for the model to be able to recover nonlinear trends, but also to recover flexible shapes that might arise due to a dependence of the weights on the covariates. As such, the function `cROC.bnp` in **ROCrReg** allows to model the mean function of each component using an additive smooth structure

$$\mu_D(\mathbf{x}_{Dj}, \boldsymbol{\beta}_{Dl}) = \beta_{Dl0} + f_{Dl1}(x_{Dj,1}) + \dots + f_{Dlq}(x_{Dj,q}), \quad l = 1, \dots, L_D, \quad (27)$$

where the smooth functions,  $f_{Dlm}$  ( $m = 1, \dots, q$ ), are approximated using a linear combination of cubic B-splines basis functions. To avoid notational burden we have assumed that all  $q$  covariates are continuous and modelled in a flexible way. However, the function `cROC.bnp` can also deal with categorical covariates, linear effects of continuous covariates, as well as interactions. For the reasons mentioned before, we recommend that all continuous covariates are modelled as in (27). Nonetheless, posterior predictive checks, as illustrated in Section [Package presentation and illustration](#), can also be used to informally validate the fitted model. We write

$$\mu_D(\mathbf{x}_{Dj}, \boldsymbol{\beta}_{Dl}) = \mathbf{z}_{Dj}^\top \boldsymbol{\beta}_{Dl}, \quad l = 1, \dots, L_D, \quad j = 1, \dots, n_D, \quad (28)$$

where  $\mathbf{z}_{Dj}^\top$  is the  $j$ th row of the design matrix that contains the intercept, the continuous covariates that are modelled in a linear way (if any), the cubic B-splines basis representation for those modelled in a flexible way, the categorical covariates (if any), and their interaction(s) (if believed to exist). Also,  $\boldsymbol{\beta}_{Dl}$  collects, for the  $l$ th component, the regression coefficients associated with the aforementioned covariates. For the covariate effects modelled using cubic B-splines, an important issue is the selection of the number and location of the knots at which to anchor the basis functions, as this has the potential to impact inferences, more so for the former than the latter. The selection of the number of knots can be assisted by a model selection criterion, for example, (the adaptation to the case of mixture models of) the deviance information criterion (DIC) ([Celeux et al., 2006](#)), the log pseudo marginal likelihood (LPML) ([Geisser and Eddy, 1979](#)), and the widely applicable information criterion (WAIC) ([Gelman et al., 2014](#)). In turn, for the location of the interior knots themselves we follow [Rosenberg \(1995\)](#) and use the quantiles of the covariate values.

The regression coefficients and variances associated with each of the  $L_D$  components are sampled from the conjugate centring distribution  $(\boldsymbol{\beta}_{Dl}, \sigma_{Dl}^{-2}) \stackrel{iid}{\sim} N_{Q_D}(\mathbf{m}_D, \mathbf{S}_D) \Gamma(a_D, b_D)$ , with conjugate hyperpriors  $\mathbf{m}_D \sim N(\mathbf{m}_{D0}, \mathbf{S}_{D0})$  and  $\mathbf{S}_D^{-1} \sim \text{Wishart}(\nu_D, (\nu_D \Psi_D)^{-1})$  (a Wishart distribution with degrees of freedom  $\nu_D$  and expectation  $\Psi_D^{-1}$ ) and where  $Q_D$  is the dimension of the vector  $\mathbf{z}_{Dj}$ . Hyperparameters  $\mathbf{m}_{D0}$  and  $\Psi_D$  must be chosen to represent the prior belief about the regression coefficients associated to each mixture component and about their covariance matrix, respectively, whereas  $\mathbf{S}_{D0}$  and  $\nu_D$  are chosen to represent the confidence in the prior belief of  $\mathbf{m}_{D0}$  and  $\Psi_D$ , respectively. As in the no-covariate case, by default in `cROC.bnp` test outcomes and covariates are standardised, which not only facilitates specification of the hyperparameter values but also improves the mixing of the Markov

chain Monte Carlo (MCMC) chains. The default values are as follows:

$$\mathbf{m}_{D0} = \mathbf{0}_{Q_D}, \quad \mathbf{S}_{D0} = 10I_{Q_D}, \quad \nu_D = Q_D + 2, \quad \Psi_D = I_{Q_D}, \quad a_D = 2, \quad b_D = 0.5.$$

When test outcomes and covariates are not standardised, the defaults are the following:

$$\mathbf{m}_{D0} = \hat{\boldsymbol{\beta}}_D, \quad \mathbf{S}_{D0} = \hat{\Sigma}_D, \quad \nu_D = Q_D + 2, \quad \Psi_D = 30\hat{\Sigma}_D, \quad a_D = 2, \quad b_D = \hat{\sigma}_D^2/2,$$

where  $\hat{\boldsymbol{\beta}}_D$  and  $\hat{\sigma}_D$  are the least squares estimates from fitting the linear model  $y_{Dj} = \mathbf{z}_{Dj}\boldsymbol{\beta}_D + \sigma_{Dj}\varepsilon_{Dj}$ , where  $E(\varepsilon_{Dj}) = 0$ ,  $\text{var}(\varepsilon_{Dj}) = 1$ , and  $\hat{\Sigma}_D$  is the estimated covariance matrix of  $\hat{\boldsymbol{\beta}}_D$ . With regard to the specification of  $a_D$  and  $L_D$ , as in the DPM model (no-covariate case) we set them, respectively, to 1 and 10. The blocked Gibbs sampler is used to simulate draws from the posterior distribution and details about it can be found, for instance, in the Supplementary Materials of [Inácio de Carvalho et al. \(2017\)](#).

Similarly to the analogous model for the no covariate case, at iteration  $s$  of the Gibbs sampler procedure, the covariate-specific ROC curve is computed as

$$\text{ROC}^{(s)}(p | \mathbf{x}) = 1 - F_D^{(s)} \left\{ F_D^{-1}(s)(1 - p | \mathbf{x}) | \mathbf{x} \right\}, \quad s = 1, \dots, S,$$

with

$$F_D^{(s)}(y | \mathbf{x}) = \sum_{l=1}^{L_D} \omega_{Dl}^{(s)} \Phi \left( y | \mathbf{z}^\top \boldsymbol{\beta}_{Dl}^{(s)}, \sigma_{Dl}^{2(s)} \right), \quad F_{\bar{D}}^{(s)}(y | \mathbf{x}) = \sum_{k=1}^{L_{\bar{D}}} \omega_{\bar{D}k}^{(s)} \Phi \left( y | \mathbf{z}^\top \boldsymbol{\beta}_{\bar{D}k}^{(s)}, \sigma_{\bar{D}k}^{2(s)} \right), \quad (29)$$

and where the inversion is performed numerically. A point estimate for  $\text{ROC}(p | \mathbf{x})$  can be obtained by computing the mean of the ensemble  $\{\text{ROC}^{(1)}(p | \mathbf{x}), \dots, \text{ROC}^{(S)}(p | \mathbf{x})\}$ , with pointwise credible bands derived from the percentiles of the ensemble. Although the results presented in [Erkanli et al. \(2006\)](#) can be extended to derive a closed-form expression for the covariate-specific AUC, for computational reasons, in **ROCNReg** the integral in (12) is approximated using Simpson's rule, and the same applies for the partial areas. Conditionally on a specific covariate value, the computation of the Youden index and of the optimal threshold proceeds in a similar way as in the DPM model (see [Inácio de Carvalho et al., 2017](#), for details). As for the covariate-specific ROC curve, point and interval estimates can be obtained from the corresponding covariate-specific ensemble of each summary measure.

We finish this section by noting that a particular case of the above estimator arises when the effect of all continuous covariates is assumed to be linear and only one component is considered, i.e.,

$$F_D^{(s)}(y | \mathbf{x}) = \Phi \left( y | \tilde{\mathbf{x}}^\top \boldsymbol{\beta}_D^{(s)}, \sigma_D^{2(s)} \right), \quad \text{and} \quad F_{\bar{D}}^{(s)}(y | \mathbf{x}) = \Phi \left( y | \tilde{\mathbf{x}}^\top \boldsymbol{\beta}_{\bar{D}}^{(s)}, \sigma_{\bar{D}}^{2(s)} \right), \quad (30)$$

with  $\tilde{\mathbf{x}}^\top = (1, \mathbf{x}^\top)$ . In this case, it is easy to show that

$$\text{ROC}^{(s)}(p | \mathbf{x}) = 1 - \Phi \left\{ a^{(s)}(\mathbf{x}) + b^{(s)} \Phi^{-1}(1 - p) \right\}, \quad (31)$$

where

$$a^{(s)}(\mathbf{x}) = \tilde{\mathbf{x}}^\top \frac{(\boldsymbol{\beta}_{\bar{D}}^{(s)} - \boldsymbol{\beta}_D^{(s)})}{\sigma_D^{(s)}}, \quad \text{and} \quad b^{(s)} = \frac{\sigma_{\bar{D}}^{(s)}}{\sigma_D^{(s)}}. \quad (32)$$

With this configuration, the model for the covariate-specific ROC curve can be regarded as a Bayesian counterpart of the induced ROC approach proposed by [Faraggi \(2003\)](#) (and detailed in the Supplementary Material). We denote it as the Bayesian normal linear model (for the test outcomes).

### Covariate-adjusted ROC curve

The estimation of the AROC curve rests on the following three steps:

1. Estimation of the conditional distribution of test outcomes in the nondiseased group,  $F_{\bar{D}}(y_{\bar{D}i} | \mathbf{x}_{\bar{D}i})$ .
2. Computation of the placement value  $U_D = 1 - F_{\bar{D}}(Y_D | \mathbf{X}_D)$  where, by a slight abuse of notation, we are designating it by the same letter used for the unconditional case.
3. Estimation of the cumulative distribution function of  $U_D$ .

The approach proposed by [Inácio de Carvalho and Rodríguez-Álvarez \(2018\)](#) for estimating the AROC curve is implemented in function **AROC.bnp** and it combines a single-weights dependent Dirichlet process mixture of normal distributions in Step 1 and the Bayesian bootstrap in Step 3. Again, here, in

Step 1, we also recommend to use cubic B-splines transformations of all continuous covariates. Using the same notation as before, we model the conditional density as

$$F_{\bar{D}}(y_{\bar{D}i} \mid \mathbf{x}_{\bar{D}i}) = \sum_{l=1}^{L_{\bar{D}}} \omega_{\bar{D}l} \Phi(y_{\bar{D}i} \mid \mathbf{z}_{\bar{D}i}^\top \boldsymbol{\beta}_{\bar{D}l}, \sigma_{\bar{D}l}^2).$$

The same prior distributions and default values as in the `cROC.bnp` function are adopted for  $\boldsymbol{\beta}_{\bar{D}l}$  and  $\sigma_{\bar{D}l}^2$ . Once Step 1 has been completed and given a posterior sample from the parameters of interest, the corresponding realisation of the placement value of a diseased subject in the nondiseased population is easily computed as

$$U_{Dj}^{(s)} = 1 - F_{\bar{D}}^{(s)}(y_{Dj} \mid \mathbf{x}_{Dj}) = \sum_{l=1}^{L_{\bar{D}}} \omega_{\bar{D}l}^{(s)} \Phi\left(y_{Dj} \mid \mathbf{z}_{Dj}^\top \boldsymbol{\beta}_{\bar{D}l}^{(s)}, \sigma_{\bar{D}l}^{2(s)}\right), \quad j = 1, \dots, n_D, \quad s = 1, \dots, S.$$

Finally, in Step 3, the cumulative distribution function of  $\{U_{Dj}^{(s)}\}_{j=1}^{n_D}$  is estimated through the Bayesian bootstrap

$$\text{AROC}^{(s)}(p) = \sum_{j=1}^{n_D} q_j^{(s)} I\left(U_{Dj}^{(s)} \leq p\right), \quad (q_1^{(s)}, \dots, q_{n_D}^{(s)}) \sim \text{Dirichlet}(n_D; 1, \dots, 1).$$

As before, closed-form expressions do exist for the AAUC and pAAUC

$$\begin{aligned} \text{AAUC}^{(s)} &= \int_0^1 \text{AROC}^{(s)}(p) dp = 1 - \sum_{j=1}^{n_D} q_j^{(s)} U_{Dj}^{(s)}, \\ \text{pAAUC}^{(s)}(u_1) &= \int_0^{u_1} \text{AROC}^{(s)}(p) dp = u_1 - \sum_{j=1}^{n_D} q_j^{(s)} \min\{u_1, U_{Dj}^{(s)}\}, \end{aligned}$$

and the  $\text{pAAUC}_{\text{TNF}}$  (Equation (22)) is computed using numerical integration methods. With regard to the YI, it is obtained by directly plugging in  $\text{AROC}^{(s)}(p)$  in Expression (23).

A point estimate for  $\text{AROC}(p)$  can be obtained by computing the mean of the ensemble  $\{\text{AROC}^{(1)}(p), \dots, \text{AROC}^{(S)}(p)\}$ , that is,

$$\widehat{\text{AROC}}(p) = \frac{1}{S} \sum_{s=1}^S \text{AROC}^{(s)}(p),$$

and the percentiles of the ensemble can be used to provide pointwise credible bands/credible intervals. The same applies for the AAUC, pAAUC, and YI.

## Package presentation and illustration

This section describes the main functions in the **ROCNReg** package and illustrates their usage using, due to confidentiality reasons, a synthetic dataset mimicking endocrine data from a cross-sectional study carried out by the Galician Endocrinology and Nutrition Foundation. A detailed description of the original dataset can be found in [Tomé Martínez de Rituerto et al. \(2009\)](#). The original data have also been previously analysed in [Rodríguez-Álvarez et al. \(2011a,b\)](#) and [Inácio de Carvalho and Rodríguez-Álvarez \(2018\)](#). The synthetic data can be found in the **ROCNReg** package under the name `endosyn` and a summary of it follows.

```
R> library("ROCNReg")
R> data("endosyn")
R> summary(endosyn)

  cvd_idf        age      gender      bmi
Min.   :0.0000  Min.   :18.25  Men   :1317  Min.   :12.60
1st Qu.:0.0000  1st Qu.:29.57  Women:1523  1st Qu.:23.19
Median :0.0000  Median :39.28          Median :26.24
Mean   :0.2433  Mean   :41.43          Mean   :26.69
3rd Qu.:0.0000  3rd Qu.:50.84          3rd Qu.:29.74
Max.   :1.0000  Max.   :84.66          Max.   :46.20
```

The dataset is comprised of 2840 individuals (1317 men and 1523 women, variable `gender`), with an age range between 18 and 85 years old. Variable `bmi` contains the body mass index (BMI) values,

and `cvd_idf` is the variable that indicates the presence (1) or absence (0) of two or more cardiovascular disease (CVD) risk factors. Following previous studies, the CVD risk factors considered include raised triglycerides, reduced HD-cholesterol, raised blood pressure, and raised fasting plasma glucose. Note that from the 2840 individuals, about 24% present two or more CVD risk factors.

Using the **ROCnReg** package, in the subsequent sections we will illustrate how to ascertain, through the pooled ROC curve, the discriminatory capacity of the BMI (which acts as our diagnostic test in this example) in differentiating individuals with two or more CVD risk factors (those belonging to the diseased class  $D$ ) from those having none or just one CVD risk factor (and that therefore belong to the nondiseased group  $\bar{D}$ ). We will also show how to evaluate, through the covariate-specific ROC curve, the possible modifying effect of age and gender on the discriminatory capacity of the BMI. Finally, the last part of this section focuses on the covariate-adjusted ROC curve as a global summary measure of the BMI discriminatory ability, when taking the age and gender effects into account. In the Supplementary Material we show the usage of the package for those methods not described here in the main text.

### Pooled ROC curve

The **ROCnReg** package allows estimating the pooled ROC curve by means of the four methods listed in Table 1. Here we only present the syntax for the functions `pooledROC.BB` and `pooledROC.dpm` that correspond, respectively, to the Bayesian bootstrap estimator and the approach based on a Dirichlet process mixture (of normal distributions). The function `pooledROC.emp`, which implements an empirical estimator, and the function `pooledROC.kernel`, which is based on kernel methods, are illustrated in the Supplementary Material. The input arguments in the functions are method-specific (details can be found in the manual accompanying the package), but in all cases numerical and graphical summaries can be obtained by calling the functions `print.pooledROC`, `summary.pooledROC`, and `plot.pooledROC`, which can be abbreviated by `print`, `summary`, and `plot`. Recall that our aim is to ascertain, using the `endosyn` dataset, the discriminatory capacity of the BMI in differentiating individuals with two or more CVD risk factors from those having just one or none CVD risk factors.

```
R> set.seed(123, "L'Ecuyer-CMRG") # for reproducibility
R> pROC_dpm <- pooledROC.dpm(marker = "bmi", group = "cvd_idf", tag.h = 0,
+ data = endosyn, standardise = TRUE, p = seq(0, 1, l = 101), ci.level = 0.95,
+ compute.lpm1 = TRUE, compute.WAIC = TRUE, compute.DIC = TRUE,
+ pauc = paucontrol(compute = TRUE, focus = "FPF", value = 0.1),
+ density = densitycontrol(compute = TRUE),
+ prior.h = priorcontrol.dpm(L = 10), prior.d = priorcontrol.dpm(L = 10),
+ mcmc = mcmccontrol(nsave = 8000, nburn = 2000, nskip = 1),
+ parallel = "snow", ncpus = 2, cl = NULL)
```

Before describing in detail the previous call, we first present the control functions that are used. In particular

```
paucontrol(compute = FALSE, focus = c("FPF", "TPF"), value = 1)
```

can be used to indicate whether the pAUC should be computed (by default it is not computed), and in case it is computed (i.e., `compute = TRUE`), whether the focus should be placed on restricted FPFs (pAUC, see (4)) or on restricted TPFs (pAUC<sub>TPF</sub>, see (5)). In both cases, the upper bound  $u_1$  (if focus is the FPF) or the lower bound  $v_1$  (if focus is the TPF) should be indicated in the argument `value`. In addition to the pooled ROC curve, AUC, and pAUC (if required), the function `pooledROC.dpm` also allows computing the probability density function (PDF) of the test outcomes in both the diseased and nondiseased groups. In order to do so, we use

```
densitycontrol(compute = FALSE, grid.h = NA, grid.d = NA)
```

By default, PDFs are not returned by the function `pooledROC.dpm`, but this can be changed by setting `compute = TRUE`, and through `grid.h` and `grid.d` the user can specify a grid of test results where the PDFs are to be evaluated in, respectively, the nondiseased and diseased groups. Value `NA` signals auto initialisation, with default a vector of length 200 in the range of the test results. Regarding the hyperparameters for the Dirichlet process mixture of normals model (used for the estimation of the PDFs/CDFs of the test outcomes in each group), they can be controlled using

```
priorcontrol.dpm(m0 = NA, S0 = NA, a = 2, b = NA, alpha = 1, L = 10)
```

A detailed description of these hyperparameters is found in Section [Methods](#). Finally, to set the various parameters controlling the MCMC procedure (which in our case is simply a Gibbs sampler) we use

```
mcmccontrol(nsave = 8000, nburn = 2000, nskip = 1)
```

Here `nsave` is an integer value with the total number of scans to be saved, `nburn` is the number of burn-in scans, and `nskip` is the thinning interval. Unless due to memory usage reasons, we recommend not thinning and instead monitoring the effective sample size of the MCMC chain.

Coming back to the `pooledROC.dpm` function, through `marker` the user specifies the name of the variable containing the test results; in our case, these are the values of the BMI. The name of the variable that distinguishes diseased (two or more CVD risk factors,  $D$ ) from nondiseased individuals (none or one CVD risk factor,  $\bar{D}$ ) is represented by the argument `group`, and the value codifying nondiseased individuals in group is specified by `tag.h`. The `data` argument is a data frame containing the data and all needed variables. Setting `standardise = TRUE` (the default) will standardise (i.e., subtract the mean and divide by the standard deviation) the test outcomes. The set of FPFs at which to estimate the pooled ROC curve is specified in the argument `p`, and argument `ci.level` allows specifying the level for the credible intervals (by default: 0.95). The LPML, WAIC, and DIC are computed by setting, respectively, the arguments `compute.lpml`, `compute.WAIC`, and `compute.DIC` to `TRUE`. Argument `pauc` is an (optional) list of values to replace the default values returned by the function `paucontrol`. Here, we ask for the pAUC to be computed, with focus on restricted FPFs and upper bound  $u_1 = 0.1$ . Similarly, the argument `density` is an (optional) list of values to replace the default values returned by the function `densitycontrol`, as it is the argument `mcmc`. Through `prior.h` and `prior.d` arguments we specify the hyperparameters in the nondiseased and diseased groups, respectively. Again, both arguments are (optional) lists of values to replace the default values returned by the function `priorcontrol.dpm`. We shall remember that different hyperparameters' default values are set depending on whether test outcomes are standardised or not. Finally, arguments `parallel`, `ncpus` and `c1` allow to perform parallel computations (based on the R-package `parallel`). In particular, through `parallel` the user specifies the type of parallel operation: either "no" (default), "multicore" (not available on Microsoft Windows operating systems) or "snow". Argument `ncpus` is used to indicate the number of processes to be used in parallel operation (when `parallel = "multicore"` or `parallel = "snow"`), and `c1` is an optional parallel or snow cluster to be used when `parallel = "snow"`. If `c1` is not supplied (as in our example), a cluster on the local machine is created for the duration of the call.

A numerical summary of the fitted model can be obtained by calling the function `summary`, that provides, among other information, the estimated AUC (posterior mean) and 95% credible interval (recall that we set in the call to the function `ci.level = 0.95`) and, if required, the LPML, WAIC, and DIC, separately, in the nondiseased (denoted here as Group H) and diseased (Group D) groups.

```
R> summary(pROC_dpm)

Call:
pooledROC.dpm(marker = "bmi", group = "cvd_idf", tag.h = 0, data = endosyn,
  standardise = TRUE, p = seq(0, 1, l = 101), ci.level = 0.95,
  compute.lpml = TRUE, compute.WAIC = TRUE, compute.DIC = TRUE,
  pauc = paucontrol(compute = TRUE, focus = "FPF", value = 0.1),
  density = densitycontrol(compute = TRUE), prior.h = priorcontrol.dpm(L = 10),
  prior.d = priorcontrol.dpm(L = 10), mcmc = mcmccontrol(nsave = 8000,
  nburn = 2000, nskip = 1), parallel = "snow", ncpus = 2, cl = NULL)
```

Approach: Pooled ROC curve - Bayesian DPM

```
-----  
Area under the pooled ROC curve: 0.759 (0.74, 0.777)*  
Partial area under the pooled ROC curve (FPF = 0.1): 0.168 (0.139, 0.199)*  
* Credible level: 0.95
```

Model selection criteria:

	Group H	Group D
WAIC	12490.485	4017.063
WAIC (Penalty)	8.431	5.468
LPML	-6245.247	-2008.541
DIC	12490.276	4016.920
DIC (Penalty)	8.326	5.396

Sample sizes:

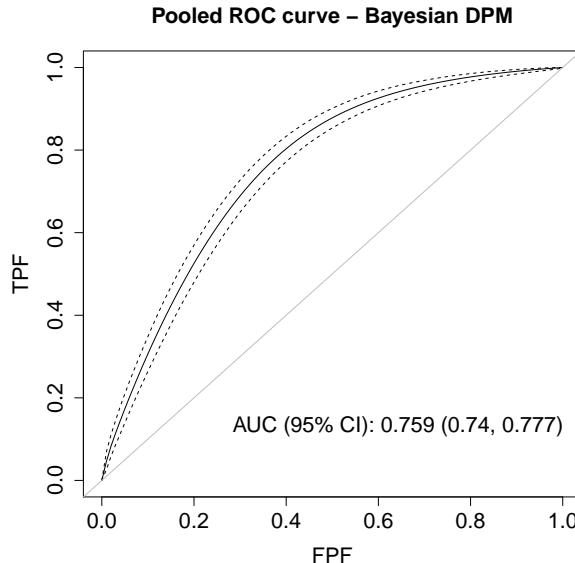
	Group H	Group D
Number of observations	2149	691
Number of missing data	0	0

To complement these numerical results, the **ROCrnReg** package also provides graphical results that can be used to further explore the fitted model. Specifically, the function `plot` depicts the estimated

pooled ROC curve and AUC (posterior means), jointly with `ci.level` $\times 100\%$  (pointwise) credible intervals (here 95%)

```
R> plot(pROC_dpm, cex.main = 1.5, cex.lab = 1.5, cex.axis = 1.5, cex = 1.5)
```

The result of the above code is shown in Figure 2.

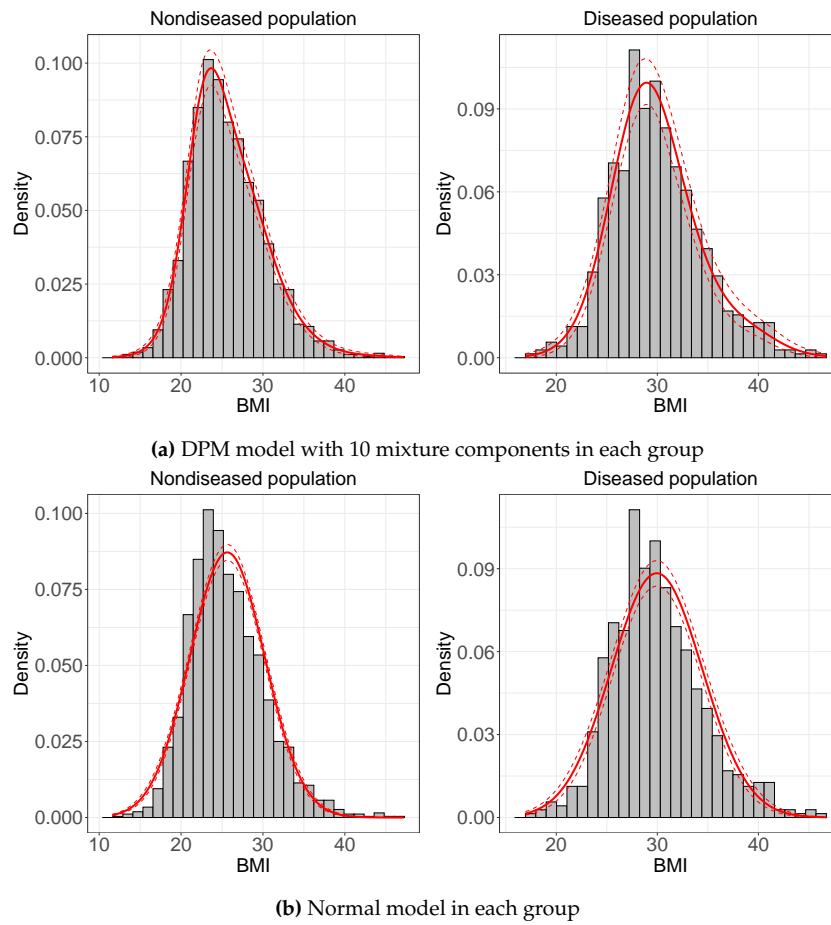


**Figure 2:** Graphical results as provided by the `plot.pooledROC` function for an object of class `pooledROC.dpm`. Posterior mean and 95% pointwise credible band for the pooled ROC curve and corresponding posterior mean and 95% credible interval for the AUC.

By means of `density = densitycontrol(compute = TRUE)` in the call to the function, the estimates of the PDFs of the BMI in both groups are to be returned. This information can be accessed through component `dens` in the object `pROC_dpm` (i.e., `pROC_dpm$dens`), which is a list with elements `h` and `d` associated with the nodiseased and diseased groups, respectively. Each of the two elements is itself another list of two components: (1) `grid`, a vector that contains the grid of test results at which the PDFs have been evaluated (estimated); and (2) `dens`, a matrix with the PDFs at each iteration of the MCMC procedure. We can use these results to plot, e.g, the posterior mean (and 95% pointwise credible bands) of the PDF of the BMI in the healthy and diseased populations (see Figure 3a obtained using the R package `ggplot2` by Wickham, 2016). As can be observed, the estimated densities obtained under the DPM method follow very closely the histograms of the data. Further, the estimated densities available in `dens` can be used, as advised by (Gelman et al., 2013, p. 553), to monitor convergence of the MCMC chains. The well-known label switching problem often leads to poor mixing of the chains of the component-specific parameters, but this may not impact convergence and mixing of the induced density/distribution of interest. For instance, Figure 4 shows the trace plots of the MCMC iterations (after burn-in) of the PDFs of the BMI in the two groups, for different (and randomly selected) values of the BMI, and Figure 5 depicts the corresponding effective sample sizes and Geweke statistics (obtained using the R package `coda` by Plummer et al., 2006). Note that all plots give evidence of a good mixing and do not suggest lack of convergence. For conciseness, the R code for reproducing Figures 3a, 4 and 5 is not provided here, but in the replication code that accompanies the paper.

It is worth noting that the function `pooledROC.dpm` also allows fitting a normal distribution in each group; this is just a particular case (for which  $L_D = L_{\bar{D}} = 1$ ) of the more general DPM model. In order to fit such model, one simply needs to set  $L = 1$  in the `prior.h` and `prior.d` arguments. The code follows.

```
R> set.seed(123, "L'Ecuyer-CMRG") # for reproducibility
R> pROC_normal <- pooledROC.dpm(marker = "bmi", group = "cvd_idf", tag.h = 0,
+ data = endosyn, standardise = TRUE, p = seq(0, 1, l = 101), ci.level = 0.95,
+ compute.lpm1 = TRUE, compute.WAIC = TRUE, compute.DIC = TRUE,
+ pauc = pauccontrol(compute = TRUE, focus = "FPF", value = 0.1),
+ density = densitycontrol(compute = TRUE),
+ prior.h = priorcontrol.dpm(L = 1), prior.d = priorcontrol.dpm(L = 1),
```



**Figure 3:** Histogram of the (observed) BMI and posterior mean jointly along with 95% pointwise credible bands (red lines) of the PDF of the BMI obtained using (a) a Dirichlet process mixture of normals model (object pROC\_dpm); and (b) a normal model (object pROC\_normal). Left: Nondiseased individuals (none or one CVD risk factor). Right: Diseased individuals (two or more CVD risk factors).

```
+ mcmc = mcmccontrol(nsave = 8000, nburn = 2000, nskip = 1),
+ parallel = "snow", ncpus = 2)
```

For the sake of space we omit from the summary the call to the function

```
R> summary(pROC_normal)
```

```
Call: [...]
```

```
Approach: Pooled ROC curve - Bayesian DPM
```

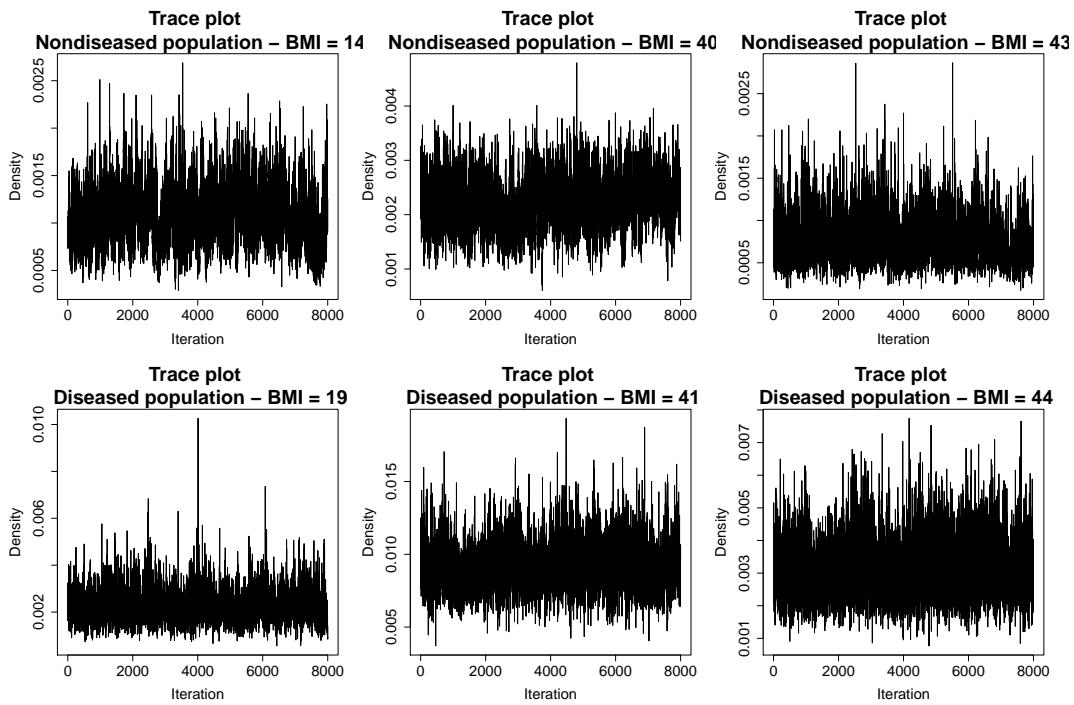
```
-----  
Area under the pooled ROC curve: 0.748 (0.728, 0.768)*  
Partial area under the pooled ROC curve (FPF = 0.1): 0.224 (0.194, 0.253)*  
* Credible level: 0.95
```

```
Model selection criteria:
```

	Group H	Group D
WAIC	12639.952	4049.004
WAIC (Penalty)	2.431	2.267
LPML	-6319.976	-2024.502
DIC	12639.505	4048.714
DIC (Penalty)	1.986	1.987

```
Sample sizes:
```

	Group H	Group D
--	---------	---------



**Figure 4:** Trace plots of the MCMC draws (after burn-in) of the PDFs of the BMI based on the model pROC\_dpm. Results are shown separately for the nondiseased and diseased populations and for different values of the BMI.

Number of observations	2149	691
Number of missing data	0	0

The fit of the DPM and normal models, in each group, can be compared on the basis of the WAIC, DIC, and/or the LPML. Remember that for the LPML, the higher its value, the better the model fit, while for the WAIC and DIC it is the other way around. By comparing these values, provided in the summary of each fitted model, we can conclude that the three criteria favour, in both the diseased and (especially in the) nondiseased groups, the more general DPM model. This is also corroborated by the plot of the fitted densities in each group shown in Figure 3b.

We now estimate the pooled ROC curve using the Bayesian bootstrap estimator (function pooledROC.BB), and comparisons with the results obtained using the DPM approach are provided.

```
R> set.seed(123, "L'Ecuyer-CMRG") # for reproducibility
R> pROC_BB <- pooledROC.BB(marker = "bmi", group = "cvd_idf", tag.h = 0, data = endosyn,
+ p = seq(0, 1, l = 101), pauc = paucontrol(compute = TRUE, focus = "TPF", value = 0.8),
+ B = 5000, ci.level = 0.95, parallel = "snow", ncpus = 2)

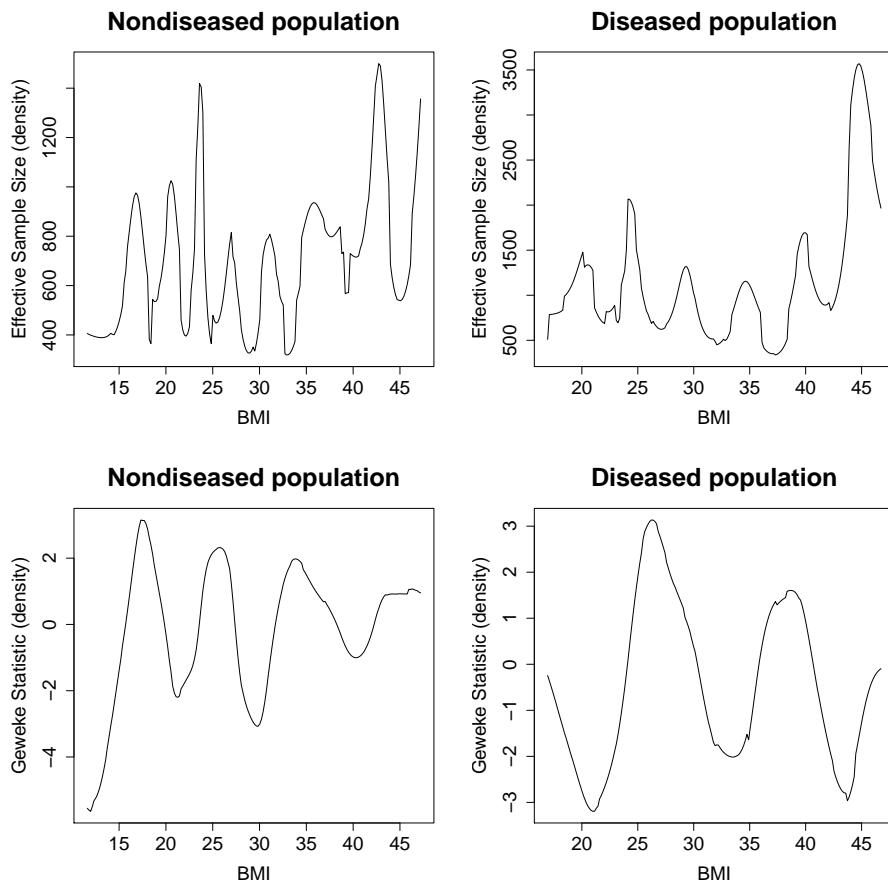
R> summary(pROC_BB)

Call: [...]

Approach: Pooled ROC curve - Bayesian bootstrap
-----
Area under the pooled ROC curve: 0.76 (0.74, 0.779)*
Partial area under the pooled ROC curve (FPP = 0.1): 0.17 (0.14, 0.201)*
* Credible level: 0.95

Sample sizes:
          Group H      Group D
Number of observations   2149       691
Number of missing data      0         0
```

Note that the posterior means for the AUC and pAUC obtained using the DPM method (0.759 and 0.168, respectively) and the Bayesian bootstrap approach (0.760 and 0.170) are almost identical. This



**Figure 5:** Effective sample size and Geweke statistic of the MCMC chains (after burn-in) and of the PDFs of the BMI based on model pROC\_dpm in the nondiseased and diseased populations. In both cases, results are shown along BMI values.

can also be observed when plotting the estimated ROC curves under the two methods (Figure 6).

We finish this section by showing how to use **ROCnReg** to obtain an (optimal) threshold value which could be further used to ‘diagnose’ an individual as diseased (two or more CVD risk factors) or healthy/nondiseased (none or only one CVD risk factor). To that aim, and for pooledROC objects (i.e., those obtained using functions pooledROC.dpm, pooledROC.BB, pooledROC.emp, and pooledROC.kernel), we use the function compute.threshold.pooledROC, which allows obtaining (optimal) threshold values using two criteria: the YI and the one that sets a target value for the FPF. For illustration, we show here the results using the YI criterion.

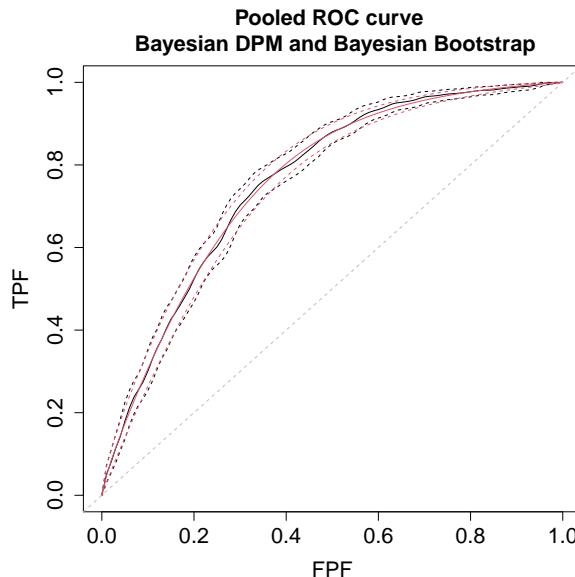
```
R> th_pROC_dmp <- compute.threshold.pooledROC(pROC_dpm, criterion = "YI",
+ ci.level = 0.95, parallel = "snow", ncpus = 2)
R> th_pROC_dmp

$call
compute.threshold.pooledROC(object = pROC_dpm, criterion = "YI",
                               ci.level = 0.95, parallel = "snow", ncpus = 2)

$thresholds
  est      ql      qh
26.46877 26.07129 26.85029

$YI
  est      ql      qh
0.4045776 0.3721684 0.4366298

$FPP
  est      ql      qh
```



**Figure 6:** Estimated ROC curve using the Bayesian bootstrap approach (in black) and the DPM method (in red). Solid lines represent the posterior means and dashed lines the 95% pointwise credible bands.

```
0.3808336 0.3469580 0.4159478
```

```
$TPF
  est      q1      qh
0.7854112 0.7528575 0.8161865
```

The function returns the posterior mean (est) and ci.level  $\times 100\%$  (here 95% since ci.level = 0.95) credible interval (lower bound: q1, upper bound: qh) for the YI and associated threshold value, as well as for the FPF and TPF associated with this cutoff value. For our example, the (posterior mean of the) YI is 0.40 and the YI-based threshold value is a BMI value of 26.5, which falls in the nutritional status defined as pre-obesity by the World Health Organization. By using this YI-based threshold value, we would have a FPF of 0.38 and a TPF of 0.79.

### Covariate-specific ROC curve

We now turn our attention to the inclusion of covariates in ROC analysis. As shown in Table 1, with **ROCNReg** the user can estimate the covariate-specific ROC curve by means of three approaches. As for the functions in **ROCNReg** for estimating the pooled ROC curve, the input arguments are method-specific, and we refer the reader to the manual for details. For all methods, numerical and graphical summaries are obtained using functions `print.cROC`, `summary.cROC` and `plot.cROC`. Here we describe how to use the function `cROC.bnp` that implements the Bayesian nonparametric approach for estimating the covariate-specific ROC curve detailed in Section [Methods](#). Also, for objects of this class, **ROCNReg** provides the function `predictive.checks`, which implements tools for assessing model fit via posterior predictive checks.

Recall that, when including covariate information in ROC analysis, interest resides in evaluating if and how the discriminatory capacity of the test varies with such covariates. In particular, in our endocrine study we aim at evaluating the possible effect of both age and gender in the discriminatory capacity of the BMI. In what follows, with this aim in mind, two different models are fitted using the function `cROC.bnp`. One which considers a normal distribution in each group and that incorporates the age effect in a linear way, and a second one which caps the maximum number of mixture components in each group at 10 (i.e.,  $L_D = L_{\bar{D}} = 10$ ) and that models the age effect using cubic B-splines (and thus allows for a nonlinear effect of age). Following Rodríguez-Álvarez et al. (2011a,b), both models consider the interaction between age and gender. For clarity, we first focus on the code that models the age effect in a linear way and use it to describe in detail the different arguments of the `cROC.bnp` function.

```
R> # Dataframe for predictions
R> agep <- seq(22, 80, l = 30)
```

```
R> endopred <- data.frame(age = rep(agep,2), gender = factor(rep(c("Women", "Men"),
+ each = length(agep))))
```

```
R> set.seed(123, "L'Ecuyer-CMRG") # for reproducibility
R> cROC_bp <- cROC.bnp(formula.h = bmi ~ gender*age, formula.d = bmi ~ gender*age,
+ group = "cvd_idf", tag.h = 0, data = endosyn, newdata = endopred,
+ standardise = TRUE, p = seq(0, 1, 1 = 101), ci.level = 0.95, compute.lpm1 = TRUE,
+ compute.WAIC = TRUE, compute.DIC = TRUE, pauc = paucontrol(compute = FALSE),
+ prior.h = priorcontrol.bnp(L = 1), prior.d = priorcontrol.bnp(L = 1),
+ density = densitycontrol(compute = TRUE),
+ mcmc = mcmccontrol(nsave = 8000, nburn = 2000, nskip = 1),
+ parallel = "snow", ncpus = 2)
```

As can be seen, many arguments coincide with those of the function `pooledROC.dpm` (described in the previous section). We thus focus here on those that are specific to `cROC.bnp`. The arguments `formula.h` and `formula.d` are `formula` objects specifying the model for the regression function (see Equation (28)) in, respectively, the nondiseased and diseased groups. They are similar to the formula used with the `glm` function, except that nonlinear functions (modelled by means of cubic B-splines) can be added using function `f` (an example will follow later in this section). Note that in both cases, the left-hand side of the formulas should include the name of the test/marker (in our case `bmi`). In our application, and for both groups, the model for the component's means includes, in addition to the linear effect of age and gender, the (linear) interaction between these two covariates (i.e.,  $\text{gender} \times \text{age} \equiv \text{gender} + \text{age} + \text{gender} : \text{age}$ ). Through the `newdata` argument the user can specify a new data frame containing the values of the covariates at which the covariate-specific ROC curve and AUC (and also pAUC and PDFs, if required) are to be computed. Finally, `prior.h` (the same holds for `prior.d`) is an (optional) list of values to replace the defaults returned by `priorcontrol.bnp` which allows setting the hyperparameters for the single-weights dependent Dirichlet process mixture of normals model (see Section [Methods](#) and the manual accompanying the package for more details)

```
priorcontrol.bnp(m0 = NA, S0 = NA, nu = NA, Psi = NA, a = 2, b = NA,
alpha = 1, L = 10)
```

In our example, we only modified the upper bound for the number of components in the mixture model, which by default is 10, and set it to 1.

In this case, the summary of the fitted model provides the following information.

```
R> summary(cROC_bp)
```

```
Call: [...]
```

```
Approach: Conditional ROC curve - Bayesian nonparametric
```

```
-----
```

```
Parametric coefficients
```

```
Group H:
```

	Post. mean	Post. quantile 2.5%	Post. quantile 97.5%
(Intercept)	26.1459	25.8765	26.4096
genderWomen	-0.9160	-1.2726	-0.5680
age	1.1949	0.9180	1.4690
genderWomen:age	1.1948	0.8455	1.5394

```
Group D:
```

	Post. mean	Post. quantile 2.5%	Post. quantile 97.5%
(Intercept)	29.1865	28.7625	29.6115
genderWomen	2.0826	1.3705	2.7665
age	0.6578	0.2162	1.0904
genderWomen:age	-0.7711	-1.4655	-0.0956

```
ROC curve:
```

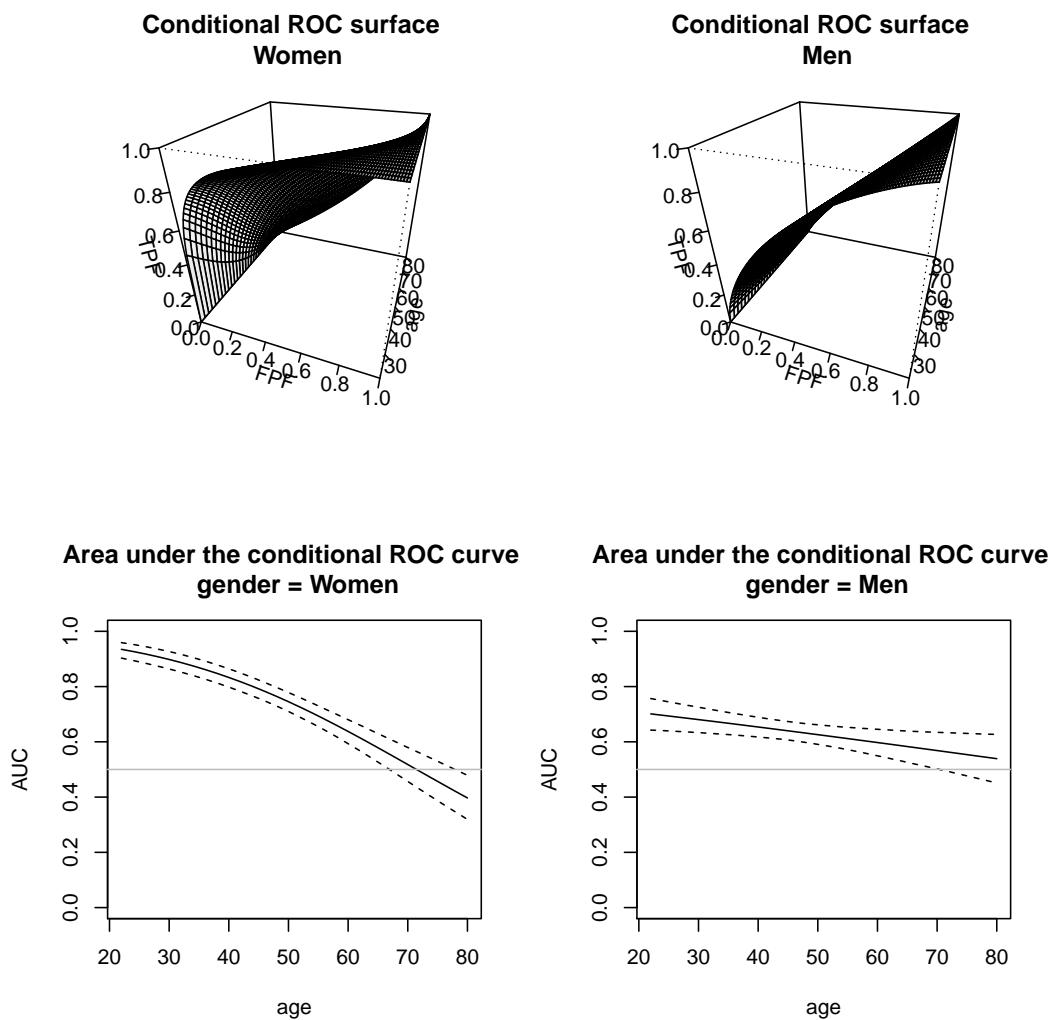
	Post. mean	Post. quantile 2.5%	Post. quantile 97.5%
(Intercept)	-0.6959	-0.8177	-0.5776
genderWomen	-0.6863	-0.8695	-0.5046
age	0.1229	0.0045	0.2415
genderWomen:age	0.4499	0.2745	0.6245

b	0.9391	0.8824	0.9975
<b>Model selection criteria:</b>			
	Group H	Group D	
WAIC	12174.986	4007.980	
WAIC (Penalty)	6.283	5.646	
LPML	-6087.492	-2003.990	
DIC	12173.664	4007.329	
DIC (Penalty)	4.994	5.053	
<b>Sample sizes:</b>			
	Group H	Group D	
Number of observations	2149	691	
Number of missing data	0	0	

The first aspect to note is that, in this case, the summary function does not provide the estimated AUC as there is one (possibly different) AUC for each combination of covariate values. Also, given that: (1) only one component has been considered for modelling the CDFs of test results in the diseased and nondiseased groups, and (2) covariate effects have been modelled in a linear way, the summary function provides the posterior mean (and quantiles) of the (parametric) coefficients associated with the regression functions (Equation (30)) and with the covariate-specific ROC curve (Equation (32)). We note that since in the call to the function we have specified `standardise = TRUE` (and consequently both the test outcomes and covariates are standardised), the regression coefficients are on the scale of the standardised covariates. If we focus on the coefficients for the covariate-specific ROC curve, it seems that the discriminatory capacity of the BMI decreases with age, with the decrease being more pronounced in women (note that the expression of the covariate-specific ROC curve in Equation (31) implies that positive coefficients correspond to a decrease in discriminatory capacity). These results are possibly better judged by plotting the estimated covariate-specific ROC curves and associated AUCs. This can be done using the `plot` function. For the covariate-specific ROC curve, the depicted graphics will depend on the number and nature of the covariates included in the analyses. In particular, for our application, we obtain, separately for men and women, the covariate-specific ROC curves (and AUCs) along age. These are shown in Figure 7, obtained using the code

```
R> op <- par(mfrow = c(2,2))
R> plot(cROC_sp, ask = FALSE)
R> par(op)
```

Although in this example we have modelled the age effect linearly and only one mixture component was considered, **ROCNReg** also allows for modelling the effect of continuous covariates in a nonlinear way, either using cubic B-spline basis expansions (through the function `cROC.bnp`) or kernel-based smoothers (via the function `cROC.kernel` which is described in the Supplementary Material). Also, as noted before, using only one mixture component for the single-weights dependent Dirichlet process mixture of normals model (function `cROC.bnp`) is equivalent to consider a (Bayesian) normal model, which might be too restrictive for most data applications. In what follows, we provide more flexibility to the model for the covariate-specific ROC curve by means of (1) increasing the number of mixture components, and (2) modelling the age effect in a nonlinear way (recall our considerations in Section [Covariate-specific ROC curve](#) about the lack of flexibility of the single-weights dependent Dirichlet process mixture of normals model when covariates effects on the components' means are modelled linearly). The former is done by modifying the value of `L` in the arguments `prior.h` and `prior.d`, with 10 being the default value. Regarding the latter, this is done by making use of the function `f` when specifying the component's mean functions through `formula.h` and `formula.d`. In particular, in our application we are interested in modelling the factor-by-curve interaction between age and gender (i.e., we model the age effect 'separately' for men and women). This is done using, e.g., `bmi ~ gender + f(age, by = gender, K = c(3,5))`. Through the argument `K` we indicate the number of internal knots that are used for constructing the cubic B-spline basis that is used to approximate the nonlinear effect of age (with the quantiles of age used to anchor the knots). Note that we can specify a different number of internal knots for men and women (`K = c(3,5)`), where the order of vector `K` should match the ordering of the levels of the factor `gender`. We also note that to assist in the selection of the number of interior knots (in **ROCNReg** the location is always based on the quantiles of the corresponding covariates), the user can make use of the WAIC, DIC, and/or LPML. For instance, for this application, we fitted different models with different number of internal knots and we have chosen the model that provided the lowest WAIC (this was done in both the nondiseased and diseased populations and we remark that the number of knots does not need to be the same in the two populations). The final model is shown below.



**Figure 7:** Graphical results as provided by the `plot.cROC` function for an object of class `cROC.bnp`. Results for the model that includes the linear interaction between age and gender and one mixture component. Top row: Posterior mean of the covariate-specific ROC curve along age, separately for men and women. Bottom row: Posterior mean and 95% pointwise credible band for the covariate-specific AUC along age, separately for men and women.

```
R> # Levels of gender, and its ordering.
R> # Needed if we want to specify different
R> # number of knots for men and women
R> levels(endosyn$gender)

[1] "Men"   "Women"

R> set.seed(123, "L'Ecuyer-CMRG") # for reproducibility
R> cROC_bnp <- cROC.bnp(
+ formula.h = bmi ~ gender + f(age, by = gender, K = c(0,0)),
+ formula.d = bmi ~ gender + f(age, by = gender, K = c(4,4)),
+ group = "cvd_idf", tag.h = 0, data = endosyn, newdata = endopred,
+ standardise = TRUE, p = seq(0, 1, 1 = 101), ci.level = 0.95, compute.lpml = TRUE,
+ compute.WAIC = TRUE, compute.DIC = TRUE, pauc = pauccontrol(compute = FALSE),
+ prior.h = priorcontrol.bnp(L = 10), prior.d = priorcontrol.bnp(L = 10),
+ density = densitycontrol(compute = TRUE),
+ mcmc = mcmccontrol(nsave = 8000, nburn = 2000, nskip = 1),
+ parallel = "snow", ncpus = 2)
```

```
R> summary(cROC_bnp)

Call: [...]

Approach: Conditional ROC curve - Bayesian nonparametric
-----

Model selection criteria:
                         Group H      Group D
WAIC           11833.000   3909.828
WAIC (Penalty)    31.236    38.583
LPML          -5916.766  -1955.449
DIC            11829.750   3904.532
DIC (Penalty)     29.611    35.934

Sample sizes:
                         Group H      Group D
Number of observations    2149       691
Number of missing data        0         0

R> op <- par(mfrow = c(2,2))
R> plot(cROC_sp, ask = FALSE)
R> par(op)
```

The graphical results are shown in Figure 8. Note that, especially for women, age displays a marked nonlinear effect. Recall that for objects of class `cROC.bnp`, and if required in the call to the function, the `summary` function provides, separately for the diseased and nondiseased/healthy groups, the WAIC, LPML, and DIC. Note that, in both cases, the three criteria support the use of the more flexible model that uses cubic B-splines and 10 mixture components for modelling the distribution of the BMI (model `cROC_bnp`) over the more restrictive Bayesian normal linear model (model `cROC_bp`). Because the WAIC, LPML, and DIC are relative criteria, posterior predictive checks are also available in **ROCNReg** through the function `predictive.checks`. Specifically, the function generates replicated datasets from the posterior predictive distribution in the two groups  $D$  and  $\bar{D}$  and compares them to the test values (BMI values in our application) using specific statistics. For the choice of such statistics we follow [Gabry et al. \(2019\)](#), who suggest choosing statistics that are ‘orthogonal’ to the model parameters. Since we are using a location-scale mixture of normals model for the test outcomes, we use here the skewness and kurtosis and check how well the posterior predictive distribution captures these two quantities.

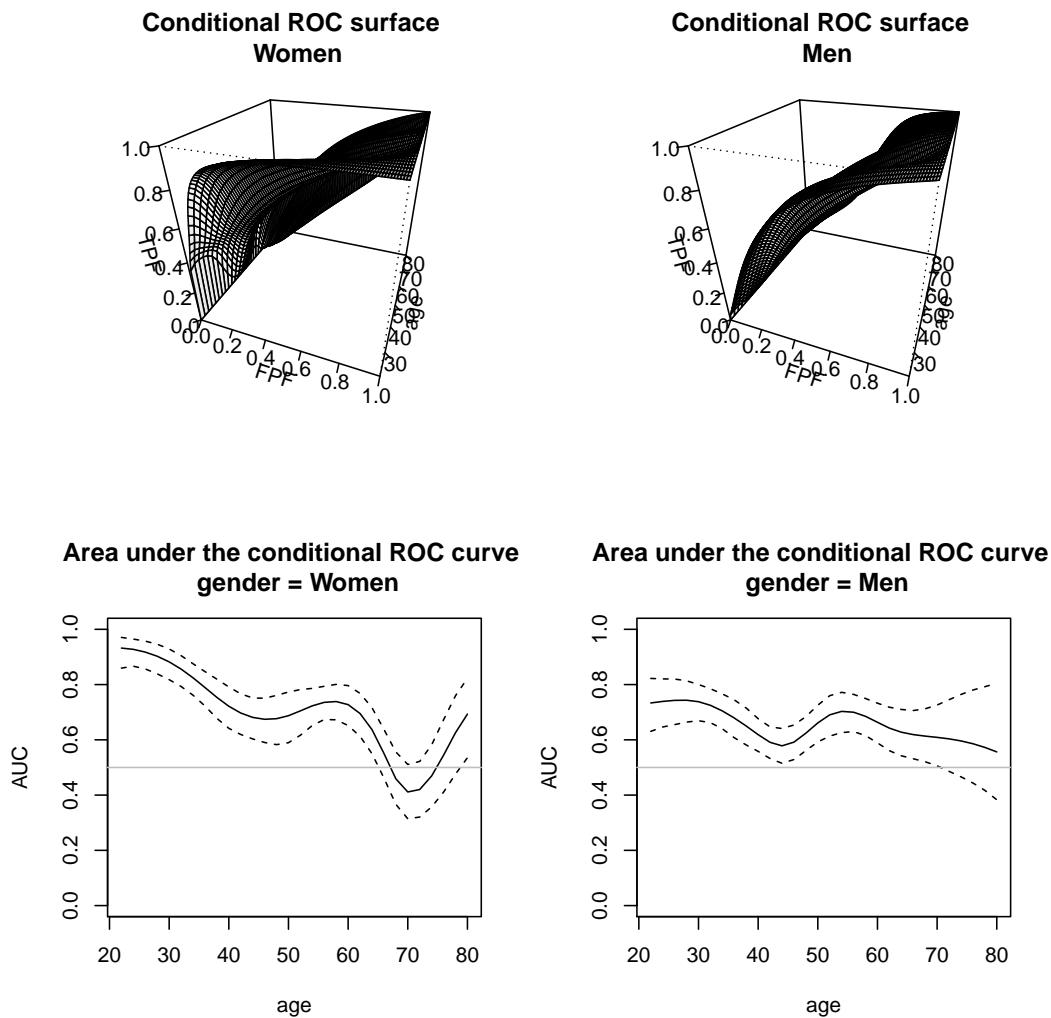
```
R> op <- par(mfrow = c(2,3))
R> pc_cROC_bp <- predictive.checks(cROC_bp,
+ statistics = c("kurtosis", "skewness"), devnew = FALSE)
R> par(op)

R> op <- par(mfrow = c(2,3))
R> pc_cROC_bnp <- predictive.checks(cROC_bnp,
+ statistics = c("kurtosis", "skewness"), devnew = FALSE)
R> par(op)
```

Results are shown in Figure 9. As can be seen, the model that includes the factor-by-curve interaction between age and gender and 10 mixture components performs quite well in capturing both quantities, while the Bayesian normal linear model fails to do so. Also shown in Figure 9 (and provided by function `predictive.checks`) are the kernel density estimates of 500 randomly selected datasets drawn from the posterior predictive distribution, in each group, compared to the kernel density estimate of the observed BMI (in each group). Again, the more flexible model, as opposed to the Bayesian normal linear model, is able to simulate data that are very much similar to the observed BMI values.

As for the pooled ROC curve, **ROCNReg** also provides a function that allows obtaining (optimal) threshold values for the covariate-specific ROC curve. For illustration, instead of the threshold values based on the Youden index, we now use the criterion that sets a target value for the FPF. The code for model `cROC_bnp`, when setting the FPF = 0.3, is as follows.

```
R> th_fpf_cROC_bnp <- compute.threshold.cROC(cROC_bnp, criterion = "FPF", FPF = 0.3,
+ newdata = endopred, ci.level = 0.95, parallel = "snow", ncpus = 2)
R> names(th_fpf_cROC_bnp)
```

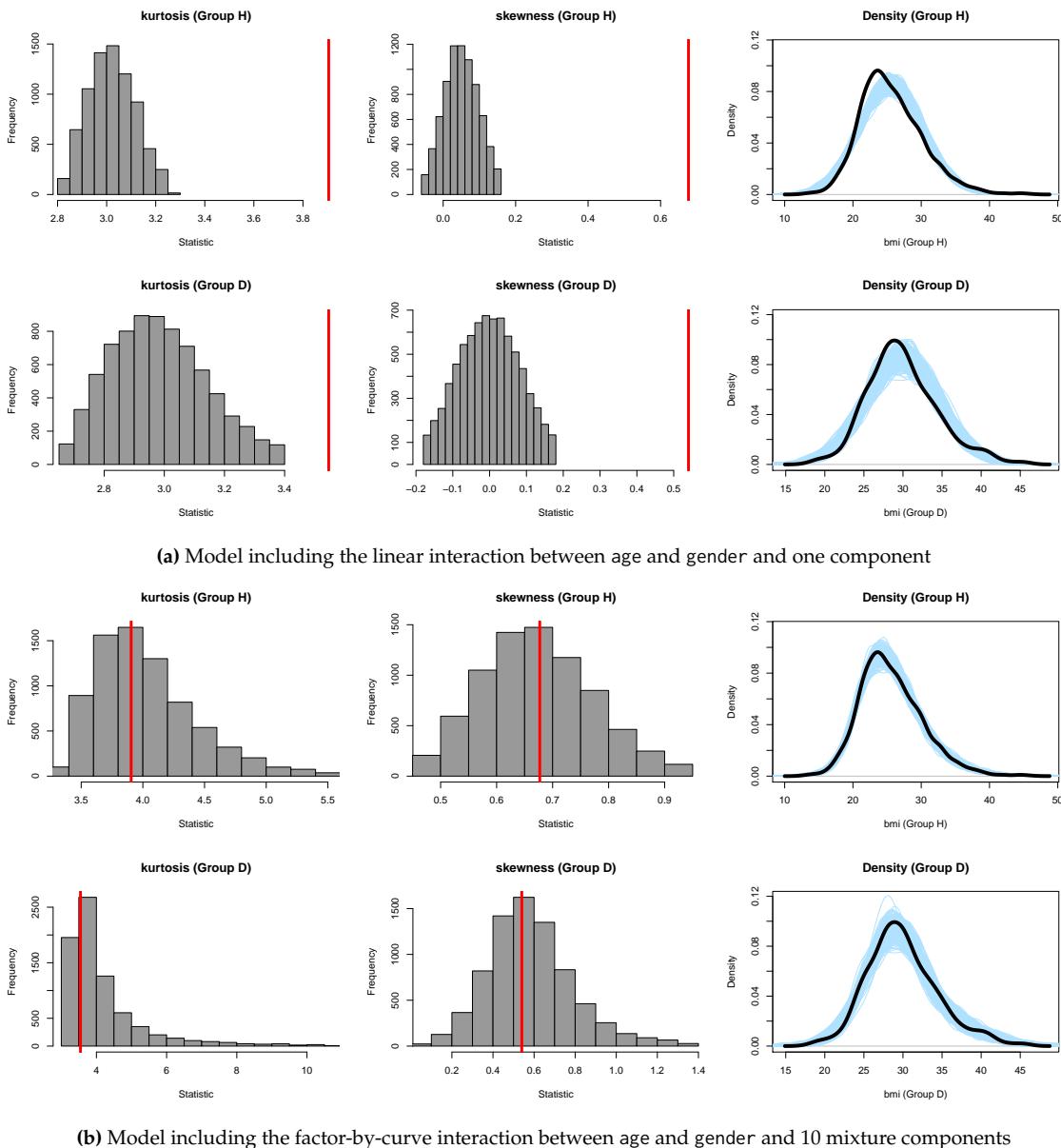


**Figure 8:** Graphical results as provided by the `plot.cROC` function for an object of class `cROC.bnp`. Results for the model that includes the factor-by-curve interaction between age and gender and 10 mixture components. Top row: Posterior mean of the covariate-specific ROC curve along age, separately for men and women. Bottom row: Posterior mean and 95% pointwise credible band for the covariate-specific AUC along age, separately for men and women.

```
[1] "newdata"    "thresholds" "TPF"      "FPF"      "call"
```

In addition to the data frame `newdata` containing the covariate values at which the thresholds are computed, the function `compute.threshold.cROC` also returns the covariate-specific thresholds corresponding to the specified FPF as well as the covariate-specific TPF attached to these thresholds. In both cases, the function returns the posterior mean and the  $ci.level \times 100\%$  (here 95%) pointwise credible intervals. Although `ROCNReg` does not provide a function for plotting the results obtained using `compute.threshold.cROC`, graphical results can be easily obtained. For simplicity, we only show here the code for the covariate-specific threshold values (`thresholds`), but a similar code can be used to plot the covariate-specific TPFs. Both plots are shown in Figure 10. As can be observed, for a FPF of 0.3, the BMI age-specific thresholds tend to increase with age both for men and women, although for the latter there is a slight decrease after an age of about 70 years old. The age-specific TPFs corresponding to the thresholds for which the FPF is 0.3 show a nonlinear behaviour and these are in general higher for women than for men (of the same age).

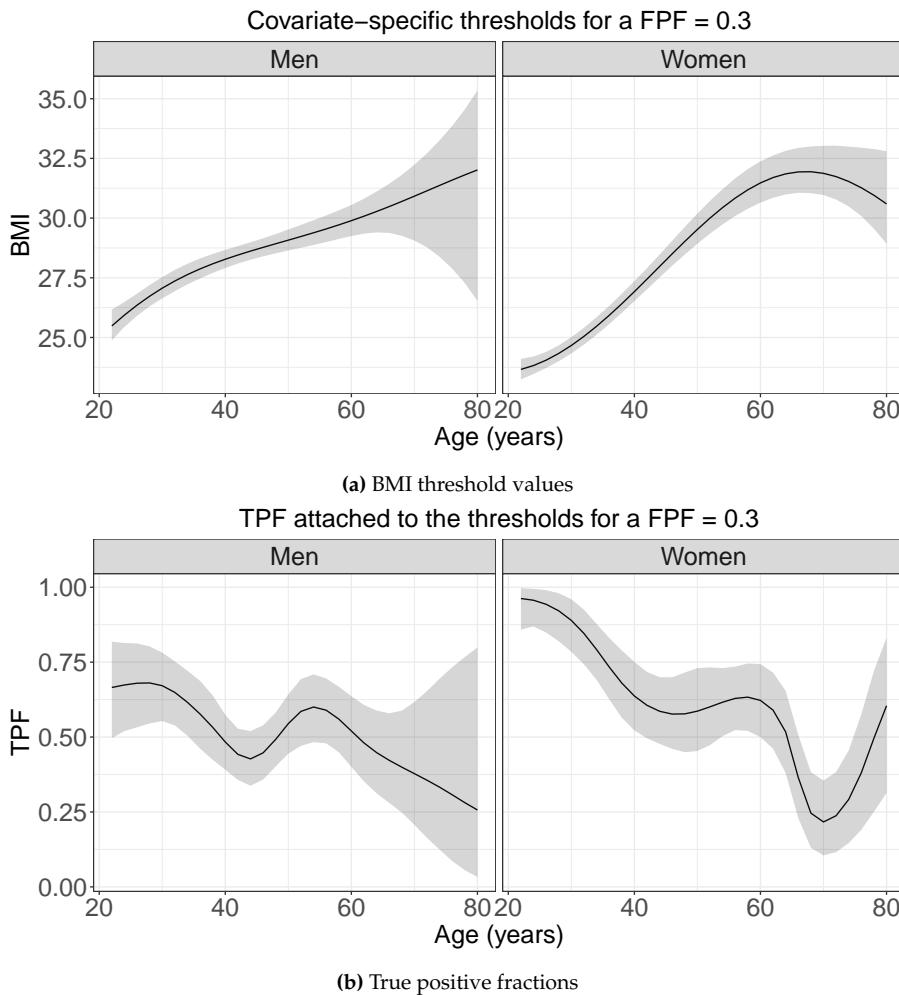
```
R> df <- data.frame(age = th_fpf_cROC_bnp$newdata$age,
+ gender = th_fpf_cROC_bnp$newdata$gender, y = th_fpf_cROC_bnp$thresholds[[1]][,"est"],
+ ql = th_fpf_cROC_bnp$thresholds[[1]][,"ql"],
+ qh = th_fpf_cROC_bnp$thresholds[[1]][,"qh"])
```



**Figure 9:** Graphical results as provided by the `predictive.checks` function for an object of class `cROC.bnp`. Histograms of the statistics *skewness* and *kurtosis* computed from 8000 draws from the posterior predictive distribution in the diseased and nondiseased group. The red line is the estimated statistic from the observed BMI values. The right-hand side plots show the kernel density estimate of the observed BMI (solid black line), jointly with the kernel density estimates for 500 simulated datasets drawn from the posterior predictive distributions.

```
R> g0 <- ggplot(df, aes(x = age, y = y, ymin = ql, ymax = qh)) + geom_line() +
+   geom_ribbon(alpha = 0.2) +
+   labs(title = "Covariate-specific thresholds for a FPF = 0.3",
+       x = "Age (years)", y = "BMI") +
+   theme(strip.text.x = element_text(size = 20),
+         plot.title = element_text(hjust = 0.5, size = 20),
+         axis.text = element_text(size = 20),
+         axis.title = element_text(size = 20)) + facet_wrap(~gender)
R> print(g0)
```

For conciseness we have not shown here how to perform convergence diagnostics of the MCMC chains for models fitted using the function `cROC.bnp`. In very much the same way as shown in the



**Figure 10:** Top row: Posterior mean (solid black line) and 95% pointwise credible band for the BMI threshold values, along age, corresponding to a FPF of 0.3. Bottom row: Posterior mean (solid black line) and 95% pointwise credible band of the TPFs, along age, corresponding to the BMI threshold values for which FPF = 0.3.

previous section for the object `pROC_dpm`, using the information contained in component `dens` in the list of returned values (if required), one can produce trace plots of the conditional densities at some sampled values, as well as obtain the corresponding effective sample sizes and Geweke statistics. Some results are provided in the Supplementary Material, and the associated code can be found in the replication code that accompanies this paper.

### Covariate-adjusted ROC curve

In this section we illustrate how to conduct inference about the covariate-adjusted ROC curve using `ROCrNReg`. Similarly to the covariate-specific ROC curve, three approaches are available for estimating the AROC curve. The function `AROC.bnp` is illustrated below, while `AROC.sp` and `AROC.kernel` are exemplified in the Supplementary Material.

Recall that the AROC curve is a global summary measure of diagnostic accuracy that takes covariate information into account. In the context of our endocrine application we seek to study the overall discriminatory capacity of the BMI for detecting the presence of CVD risk factors when adjusting for age and gender. Here we focus on how to estimate the AROC curve using the `AROC.bnp` function. The function syntax is exactly similar to the one of `cROC.bnp`, with the only difference being that we only need to specify the arguments related to the nondiseased population. The code and respective summary follow.

```
R> set.seed(123, "L'Ecuyer-CMRG") # for reproducibility
R> AROC_bnp <- AROC.bnp(
```

```

+ formula.h = bmi ~ gender + f(age, by = gender, K = c(0,0))
+ group = "cvd_idf", tag.h = 0, data = endosyn, standardise = TRUE,
+ p = seq(0, 1, l = 101), ci.level = 0.95, compute.lpml = TRUE, compute.WAIC = TRUE,
+ compute.DIC = TRUE, pauc = pauccontrol(compute = FALSE),
+ prior.h = priorcontrol.bnp(L = 10), density = densitycontrol(compute = TRUE),
+ mcmc = mcmccontrol(nsave = 8000, nburn = 2000, nskip = 1),
+ parallel = "snow", ncpus = 2)

R> summary(AROC_bnp)

Call: [...]

Approach: AROC Bayesian nonparametric
-----
Area under the covariate-adjusted ROC curve: 0.656 (0.629, 0.684)*
* Credible level: 0.95

Model selection criteria:
                    Group H
WAIC           11833.000
WAIC (Penalty)    31.236
LPML          -5916.766
DIC            11829.750
DIC (Penalty)    29.611

Sample sizes:
                    Group H     Group D
Number of observations      2149       691
Number of missing data        0         0

```

The area under the AROC curve is 0.656 (95% credible interval: (0.629, 0.684)) thus revealing a reasonable good ability of the BMI to detect the presence of CVD risk factors when teasing out the age and gender effects. As for the pooled ROC curve and the covariate-specific ROC curve, a plot function is also available (result in Figure 11a).

```
R> plot(AROC_bnp, cex.main = 1.5, cex.lab = 1.5, cex.axis = 1.5, cex = 1.3)
```

Finally, we compare the AROC curve with the pooled ROC curve that was obtained earlier by using a DPM model with 10 components in each group. In Figure 11b we show the plots of the two curves and, as can be noticed, the pooled ROC curve lies well above the AROC curve, thus evidencing the need for incorporating covariate information into the analysis.

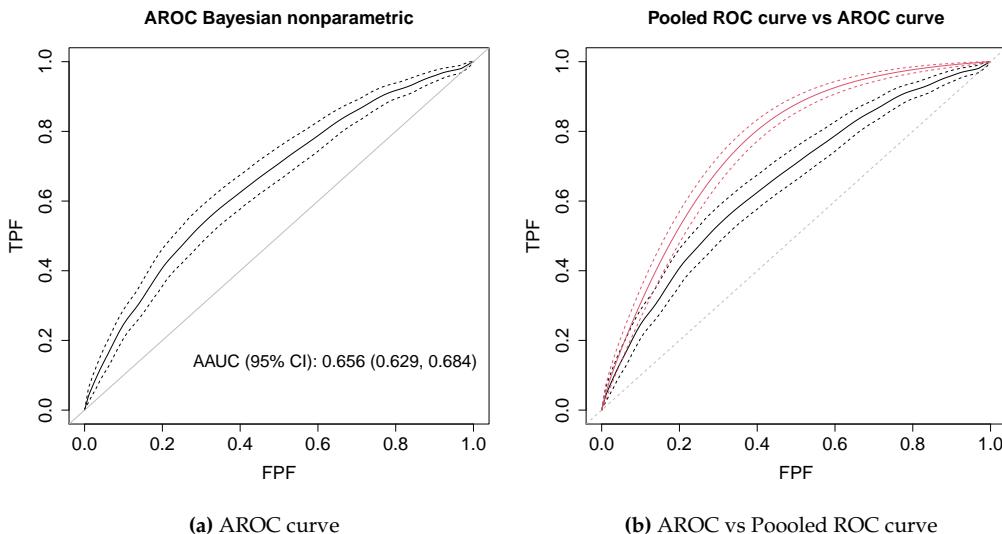
```

R> plot(AROC_bnp$p, AROC_bnp$ROC[,1], type = "l", xlim = c(0,1), ylim = c(0,1),
+ xlab = "FPF", ylab = "TPF", main = "Pooled ROC curve vs AROC curve", cex.main = 1.5,
+ cex.lab = 1.5, cex.axis = 1.5, cex = 1.5)
R> lines(AROC_bnp$p, AROC_bnp$ROC[,2], col = 1, lty = 2)
R> lines(AROC_bnp$p, AROC_bnp$ROC[,3], col = 1, lty = 2)
R> lines(pROC_dpm$p, pROC_dpm$ROC[,1], col = 2)
R> lines(pROC_dpm$p, pROC_dpm$ROC[,2], col = 2, lty = 2)
R> lines(pROC_dpm$p, pROC_dpm$ROC[,3], col = 2, lty = 2)
R> abline(0, 1, col = "grey", lty = 2)

```

## Computational aspects

We finish this section with some comments on computational aspects. In our experience, the methods with the largest computing times are those implemented in `cROC.bnp` when  $L_D > 1$  and in `cROC.kernel` when confidence bands are to be constructed. In the first case, the main reason behind the computational burden is the need to invert  $F_D(\cdot | \mathbf{x})$  in order to obtain the covariate-specific ROC curve (see equation (10)). Note that when  $L_D > 1$ , the conditional distribution function in the nondiseased group is given by a mixture of normal distributions and the corresponding quantile function needs to be computed for each covariate(s) value we might be interested in and for each iteration of the Gibbs sampler procedure. Regarding the `cROC.kernel` function, the computing time is mainly driven by the number of bootstrap samples used for constructing the confidence bands. In Table 2 we show the time, in seconds, needed for fitting the pooled, the covariate-specific, and the covariate-adjusted ROC



**Figure 11:** (a) Age and gender-adjusted ROC curve: posterior mean and 95% pointwise credible band. (b) Age and gender-adjusted ROC curve (in black) and pooled ROC curve (estimated using a DPM of normals model) (in red). Solid lines represent the posterior means and dashed lines the 95% pointwise credible bands.

curve using the Bayesian nonparametric and the kernel approaches for the synthetic endocrine data and when both parallel (with 2 and 4 processes) and no parallel options are used. We note that for the Bayesian approaches we also computed the densities/conditional densities, as well as the WAIC, LPML, and DIC, which further increase the computing time (in the case of the AROC curve these were only computed in the nondiseased population). With respect to the kernel-based approach (in this case the fit is done separately for men and women and the corresponding results are presented in the Supplementary Material) we have used 500 bootstrap samples to construct the confidence bands. As it can be appreciated, for these two intense tasks, using 4 processes drastically improves the computation time. All computations were performed in a iMac with 3.6GHz quad core Intel i7 processor and 32GB RAM running under a macOS Catalina 10.15.5 operating system.

	No parallel	Snow (2 cores)	Snow (4 cores)
pooledROC.dpm	138	118	111
pooledROC.kernel	376	196	105
cROC.bnp	2052	1117	680
cROC.kernel	Men: 1159 Women: 1885	Men: 528 Women: 916	Men: 279 Women: 466
AROC.bnp	126	115	112
AROC.kernel	Men: 847 Women: 1707	Men: 404 Women: 833	Men: 214 Women: 438

**Table 2:** Time in seconds (rounded to the nearest second) needed to fit the pooled, the covariate-specific, and the covariate-adjusted ROC curve for the Bayesian nonparametric and kernel approaches.

## Summary and future plans

In this paper we have introduced the capabilities of the R package **ROCNReg** for conducting inference about the pooled ROC curve, the covariate-specific ROC curve, and the covariate-adjusted ROC curve and their associated summary indices. As we have illustrated, the current version of the package provides several options for estimating ROC curves, both under frequentist and Bayesian paradigms, either parametrically, semiparametrically, or nonparametrically. To the best of our knowledge, this is the first software package implementing Bayesian inference for ROC curves. Several additions/extensions are planned in the future and these, among others, include:

- Implement the most time-consuming parts in C or C++.

- Incorporate methods for non-binary disease status (e.g., no disease, mild disease, severe disease). That is, implement ROC surface models.
- Implement new (optimal) threshold criteria (e.g., YI including costs).

## Computational details

The results in this paper were obtained using R 4.0.3 with the **ROCnReg** 1.0-5 package. The **ROCnReg** package has several dependencies: **graphics**, **grDevices**, **parallel**, **splines**, **stats**, **moments** (Komsta and Novomestky, 2015), **nor1mix** (Maechler, 2019), **Matrix** (Bates and Maechler, 2019), **spatstat** (Baddeley and Turner, 2005), **np** (Hayfield and Racine, 2008), **lattice** (Sarkar, 2008), **MASS** (Venables and Ripley, 2002) and **pbivnorm** (Genz and Kenkel, 2015). R itself and all packages used are available from the Comprehensive R Archive Network (CRAN) at <https://CRAN.R-project.org/>.

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