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CPS Poster

A parametric quantile beta regression for modeling case fatality rates of COVID-19

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Presentation Image

A parametric quantile beta regression for modeling case fatality rates of COVID-19 Marcelo Bourguignon ¹ Diego I. Gallardo ² Helton Saulo ³ deral University of Rio Grande do Norte ²University of Atacama ³University of Bra

An important problem

tory syndrome disease (COVID-19) p ind the world. In particular, the COVII it few months. Some authors have st research in this conter hard in the past i WID-19 in Chile as the

(1) $\frac{V}{W + V}$

Chilean COVID-19 data

ID-19 data from Chile and find that the y, positivity for tests, and the percentage (ivided into 348 ses and deaths

The proposed quantile regression model ble Y follows the generalized beta distribution (Libby and Novick, ee parameters, $\alpha > 0, \beta > 0$, and $\lambda > 0$, denoted by $Y \sim$ its cumulative distribution function (CDP) is given by $F_Y(y; \Lambda, \alpha, \beta) = I_{\lambda y/(1+\lambda y-y)}(\alpha, \beta), \quad 0 < y < 1,$ (2) $y; \lambda, \alpha, \dots, B_{\mathbb{Z}}(\alpha, \beta)/B(\alpha, \beta)$ is ω $-1(1 - \omega)^{\beta-1}d\omega$ is the i

 $f_Y(y, \lambda, \alpha, \beta) = \frac{\lambda^{\alpha}y^{\alpha-1}(1-y)^{\beta-1}}{B(\alpha, \beta)[1-(1-\lambda)y]^{\alpha+\beta}}, \quad 0 < y < 1.$

ation (3) reduces to the beta distribution wh $X_2 \sim GA(\beta, \theta_2)$ are independent gamma distri $Y \stackrel{d}{=} \frac{X_1}{X_1 + X_2} \sim \text{GB3}(\lambda, \alpha, \beta)$

to a GB3 distribution, where $\lambda=\theta_1/\theta_2.$ Note that, as the CDE of the usual beta distribution, the τ -th quantile can be represented as $q(\tau;\lambda,\alpha,\beta) = \frac{z_{\alpha,\beta}(\tau)}{\lambda[1-z_{\alpha,\beta}(\tau)]+z_{\alpha,\beta}}$ $-, 0 < \tau < 1,$ (4)

th quantile of the beta distribution is quantile $z_{\alpha} g(\tau)$ as required, it is $\lambda = \frac{(1-\mu)}{\mu} \frac{z_{\alpha,\beta}(\tau)}{[1-z_{\alpha,\beta}(\tau)]}, \quad 0 < \mu < 1.$

and $g_3(\beta_i(\tau))$ $z_i^T \nu(\tau)$,

Analysis res

CFR (c n May 24 and July 23, an=0.012. for 2020 lay 24, 2021 May 24, 202

odeling the τ -th quantile of the CFR, GB3($\mu_t(\tau), \alpha_t(\tau), \beta_t(\tau)$). We conside 0.90} and used the logit link (because it is ring that the terms related to sed on the results of AIC crit

at 0. On the eters and SEs (given in pare





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Brief Description

Motivated by the case fatality rate of COVID-19, we develop a fully parametric quantile regression model based on the generalized three-parameter beta (GB3) distribution.

We first reparameterize the GB3 distribution by inserting a quantile parameter and then we develop the new proposed quantile model.

We also propose a simple interpretation of the predictor-response relationship in terms of percentage increases/decreases of the quantile.

A real COVID-19 dataset from Chile is analyzed and discussed to illustrate the proposed approach.

Abstract

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Motivated by the case fatality rate (CFR) of COVID-19, in this paper, we develop a fully parametric quantile regression model based on the generalized three-parameter beta (GB3) distribution. Beta regression models are primarily used to model rates and proportions. However, these models are usually specified in terms of a conditional mean. Therefore, they may be inadequate if the observed response variable follows an asymmetrical distribution, such as CFR data. In addition, beta

regression models do not consider the effect of the covariates across the spectrum of the dependent variable, which is possible through the conditional quantile approach. In order to introduce the proposed GB3 regression model, we first reparameterize the GB3 distribution by inserting a quantile parameter and then we develop the new proposed quantile model. We also propose a simple interpretation of the predictor-response relationship in terms of percentage increases/decreases of the quantile. A Monte Carlo study is carried out for evaluating the performance of the maximum likelihood estimates and the choice of the link functions. Finally, a real COVID-19 dataset from Chile is analyzed and discussed to illustrate the proposed approach.