β-Blocker Therapy and Risk of Chronic Obstructive Pulmonary Disease – A Danish Nationwide Study of 1·3 Million Individuals

Anne Orholm Nielsen a, Lars Pedersen b, Birgitte Fischer Sode c, Morten Dahl a,⁎

a Department of Clinical Biochemistry, Zealand University Hospital, Lykkebakvej 1, 4600 Køge, Denmark
b Department of Clinical Epidemiology, Aarhus University Hospital, Olef Palms Allé 43-45, 8200 Aarhus N, Denmark
c Department of Thoracic Anaesthesiology, the Heart Centre, Rigshospitalet, Blegdamsvej 9, 2100 Copenhagen, Denmark

A R T I C L E   I N F O

Article history:
Received 5 September 2018
Received in revised form 20 December 2018
Accepted 15 January 2019
Available online 29 January 2019

Keywords:
β-Blockers
Obstructive pulmonary disease
Pharmacology
Epidemiology

A B S T R A C T

Background: The possible association between β-adrenoceptor antagonists (β-blockers) and risk of COPD is controversial. The objective of the present study was to test whether β-blocker use is associated with susceptibility to the disease.

Methods: A total of 301,542 new users of β-blockers and 1,000,633 new users of any other antihypertensive drugs aged 30–90 years without any history of COPD hospitalizations were included in the present study and followed in the Danish National Patient Registry for incident admissions for COPD and COPD death between 1995 and 2015. Multiple adjusted cox regression models were used to examine the association between use of β-blockers and COPD hospitalization. Additionally, subgroup analyses based on underlying diseases at baseline or duration of treatment were performed.

Findings: People treated with β-blockers continuously for more than 6 months had a lower risk of COPD hospitalization during follow-up compared to people treated with any other antihypertensive drugs (adjusted hazard ratio [HRadjusted] 0·80, 95% CI 0·79–0·82). Risk of COPD hospitalization was lower in the groups treated with β-blockers among patients with ischemic heart disease (0·72, 0·69–0·75), cardiac arrhythmias (0·76, 0·72–0·80), asthma (0·69, 0·61–0·79), hypertension (0·91, 0·86–0·96), and diseases of the pulmonary circulation (pulmonary embolism and cor pulmonale) (0·72, 0·59–0·87). All-cause mortality as well as risk of COPD death during follow-up was lower in the group treated with β-blockers compared to the group treated with any other antihypertensive drugs (0·56, 0·53–0·59).

Interpretation: Treatment with β-blockers seems to reduce risk of COPD hospitalization and mortality compared to treatment with any other antihypertensive drugs.

Funding: The Danish Council for Independent Research in Denmark (grant no. 4183-005698), The Research Foundation of Health Science in Region Zealand (grant no. RSSF2017000661 and no. 15-000342), The Research Foundation of Medical Science (A.P. Møller Foundation, grant no. 16-68), The Research Foundation in memory of King Christian 10th (grant no. 142/2017), Aase & Ejnar Danielsen’s Research Foundation (grant no. 10-001946), and Lundbeck Foundation (grant no. R252-2017-1690).

© 2019 Published by Elsevier Ltd. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/).

1. Introduction

Chronic obstructive pulmonary disease (COPD) is a common respiratory health problem characterized by persistent respiratory symptoms, such as dyspnea, cough, and airflow limitations [1]. The airflow limitation is caused by parenchymal destruction and narrowing of the small airways leading to an increase in small-airway resistance [2,3]. The respiratory symptoms can be ameliorated with β2-agonists; drugs that stimulate the β2-adrenergic receptors causing relaxation of smooth muscles in the airways and thereby bronchodilation [1].

β-adrenergic receptors are situated on smooth muscle cells in many organs and are divided into 3 subtypes, β1, β2, and β3-adrenoceptors, according to their specific actions [4]. β1-adrenoceptors are found mainly in the heart where activation increases heart rate and force of contraction [4]. β2-adrenoceptors are primarily found in the airways where activation leads to muscle relaxation and bronchodilation [5]. β-adrenoceptor antagonists (β-blockers) are drugs that block the β-adrenergic receptors and are mainly used in treatment of hypertension, cardiac arrhythmia, myocardial infarction, and ischemic heart disease.
β-adrenoceptors, the di-
2.3. Statistical Analyses

Hazard ratios (HR) for risk of COPD hospitalization and death from COPD in users of β-blockers were calculated using Cox regression models with adjustment for known confounders: age, sex, calendar year, comorbidities, income level, employment status, and marital status. In the analyses of risk of COPD hospitalization, death is censored.

Taking into account different indications for choice of treatment, we performed subgroup analyses based on different comorbidities at baseline. Subgroups that were specified in the statistical analysis plan (details provided in Supplementary Material) included diabetes mellitus, hypertension, ischemic heart disease, diseases of the pulmonary circulation (pulmonary embolism or cor pulmonale), cardiac arrhythmias (atrial fibrillation, tachycardia, or ventricular disorders), and asthma. In addition, we performed subgroup analyses stratified for age and sex as specified in the statistical analysis plan.

All analyses were performed using SAS statistical software package v. 9.4 for Windows. The study was approved by Statistics Denmark and The Danish Health Data Board (Reg. no.: 704586).

3. Results

3.1. Patient Characteristics

Novel users of either β-blockers or any other antihypertensive drugs without COPD at baseline aged 30–90 years between 1995 and 2015 were included in the current study. Baseline demographic statistics of the cohort are listed in Table 1. The sample population consisted of 301,542 users of β-blockers with a mean age of 58.1 years at baseline, and 1,000,633 users of any other antihypertensive drugs with a mean age of 61.6 years at baseline. Among users of β-blockers, 151,420 (50.2%) were females, whereas 535,471 (53.5%) were females in the group of users of any other antihypertensive drugs. Among users of β-blockers, 34,058 (11.3%) were widows/widowers and 193,930 (64.3%) were married. 164,558 (16.4%) in the group of users of any other antihypertensive drugs (54.7% vs. 48.6%) which was in line with the distribution of employed people in the two groups (47.1% vs. 39.7%). No differences in the relative numbers of unemployed or people receiving social support were observed between the two groups. Users of β-blockers were more likely to have had a history of ischemic heart disease (22.3% vs. 6.3%) and cardiac arrhythmia (15.2% vs. 3.8%). However, users of any other antihypertensive drugs were more likely to have diabetes mellitus type 1 or type 2 (5.5% vs. 2.6%).

3.2. Risk of COPD Hospitalization

Among the 301,542 users of β-blockers, 17,813 incident COPD hospitalizations were registered between 1995 and 2015, corresponding to 2,743,220 person-years, and an incident rate of 649 cases per 100,000 person-years. In the same period, 71,129 incident cases of COPD hospitalizations were observed among the 1,000,633 users of any other antihypertensive drugs, corresponding to 7,737,523 person-years, and an incident rate of 919 cases per 100,000 person-years.

After adjustment for all covariates, users of β-blockers had an overall 19.7% lower risk of COPD hospitalization compared to users of any other antihypertensive drug during follow-up (adjusted hazard ratio [HRadjust] = 0.80, 95% CI 0.79–0.82). HR for COPD hospitalization in the first 5 years after treatment start was 0.75 (0.73–0.77), while exclusion of the first 5 years of follow-up resulted in a HR for COPD hospitalization of 0.85 (0.83–0.87) (Table 2). Stratifying for age and gender did not change the overall association between β-blockers and reduced risk of COPD hospitalization (Table 2). Furthermore, the association of β-blockers remained in those participants taking β-blockers continuously for 5 years (0.84, 0.82–0.87).

Cumulative risk of COPD hospitalizations in the 2 study groups are shown in Fig. 1.

When analyzing the association between risk of COPD hospitalization and β-blockers with respect of their selectivity (or not) on bronchial tone, we did not find any significant difference between β1-selective and non-selective β-blockers. Users of β1-selective blockers had an overall 21% lower risk of COPD hospitalization (adjusted hazard ratio 0.79, 95% CI 0.78–0.81), whereas users of non-selective β-blockers had an overall 16% lower risk of COPD hospitalization (adjusted hazard ratio 0.84, 95% CI 0.82–0.87) compared to users of any other antihypertensive drug (Table 2).

3.3. Subgroup Analyses

As data on indication for the current treatment with either β-blockers or any other antihypertensive drug is not available in the registries, we performed subgroup analyses based on underlying diseases at baseline to avoid confounding by indication (Table 2). No major differences in risk of COPD hospitalization in users of β-blockers from the overall hazard ratio were seen between the groups of patients with ischemic heart disease (0.72, 0.69–0.75), cardiac arrhythmias (0.76, 0.72–0.80), hypertension, ischemic heart disease, diseases of the pulmonary circulation (pulmonary embolism or cor pulmonale), cardiac arrhythmias (atrial fibrillation, tachycardia, or ventricular disorders), and asthma.

شاركاء 1

| Baseline characteristics of the cohort. | Users of β-blockers | Users of any other AH drugs |
|------------------------------------------|---------------------|-----------------------------|
| Number of participants                   | 301,542             | 1,000,633                   |
| Mean age - years                         | 58.1                | 61.6                        |
| Age – n. (%)                             |                     |                             |
| 30–39                                    | 28,255              | 54,896 (5.5)                |
| 40–49                                    | 56,645              | 153,093 (15.3)              |
| 50–59                                    | 77,840              | 237,779 (23.8)              |
| 60–69                                    | 72,240              | 245,404 (24.5)              |
| 70–79                                    | 48,974              | 199,144 (19.9)              |
| 80–89                                    | 17,588              | 110,317 (11.0)              |
| Sex – n. (%)                             |                     |                             |
| Female                                   | 151,420             | 535,471 (53.5)              |
| Male                                     | 150,122             | 465,162 (46.5)              |
| Charlson comorbidity index – n. (%)      |                     |                             |
| None                                     | 221,332             | 723,912 (72.3)              |
| 1–2                                      | 70,020              | 229,269 (22.9)              |
| 3+                                       | 10,190              | 47,452 (4.7)                |
| Comorbidities – n. (%)                   |                     |                             |
| Diabetes type 1/type 2                   | 7852 (2.6)          | 55,010 (5.5)                |
| Hypertension                             | 29,393              | 90,501 (9.0)                |
| Ischemic heart disease                   | 67,180              | 63,237 (6.3)                |
| Diseases of the pulmonary circulation    | 1452 (0.5)          | 5267 (0.5)                  |
| Cardiac arrhythmia                       | 45,758              | 38,068 (3.8)                |
| Asthma                                   | 1801 (0.6)          | 9108 (0.9)                  |
| Income level – n. (%)                    |                     |                             |
| Lowest percentile                        | 67,664              | 257,797 (25.8)              |
| Low/mid percentile                       | 68,908              | 256,527 (25.6)              |
| Mid/high percentile                      | 79,776              | 245,733 (24.6)              |
| Highest percentile                       | 85,134              | 240,383 (24.0)              |
| Employment status – n. (%)               |                     |                             |
| Employed                                 | 142,128             | 396,886 (39.7)              |
| Unemployed                               | 18,111              | 4503 (4.6)                  |
| On the social                            | 49,158              | 155,232 (15.5)              |
| Retired                                  | 92,085              | 401,819 (40.2)              |
| Civil status – n. (%)                    |                     |                             |
| Widow/widower                            | 34,058              | 164,558 (16.4)              |
| Divorced                                 | 37,398              | 127,632 (12.8)              |
| Married                                  | 193,930             | 591,312 (59.1)              |
| Single                                   | 36,096              | 116,938 (11.7)              |

AH = antihypertensive. n. = number.
diabetes mellitus type 1/type 2 (0·95, 0·87–1·05) no effect on COPD hospitalization was seen for β-blockers (Table 2).

Furthermore, propensity score adjustment by inverse probability weighting did not change the overall association between use of β-blockers and COPD. Similar results were found when dividing β-blockers into β₁-selective blockers and non-selective blockers (Supplementary Table S4).

3.4. Mortality

All-cause mortality was substantially lower in users of β-blockers during follow-up. Among the 301,542 users of β-blockers, 52,070 deaths were registered between 1995 and 2015, corresponding to 2,824,553 person-years, and an incident rate of 1843 deaths per 100,000 person-years. In the same period, 251,720 deaths were observed among the 1,000,633 users of any other antihypertensive drugs, corresponding to 8,043,900 person-years, and an incident rate of 3129 deaths per 100,000 person-years.

During follow-up, use of β-blockers was associated with a lower risk of death from COPD compared to use of any other antihypertensive drugs (0·56, 0·53–0·59). The association between β-blockers and reduced risk of COPD death seemed to be persistent when excluding the first 5 years of follow-up (0·72, 0·67–0·78) (Table 3).

Cumulative risk of all-cause mortality in the 2 study groups are shown in Fig. 2.

4. Discussion

In this large Danish population-based study, we have evaluated the risk of COPD hospitalization in patients treated with β-blockers in the period 1995–2015. The study revealed that treatment with β-blockers, regardless of their selectivity on bronchial tone, was associated with reduced risk of COPD hospitalization, all-cause mortality, and death from COPD compared with other antihypertensive drugs.

This is in line with former studies showing that β-blockers reduce risk of exacerbations in patients with COPD. Du et al. performed in 2014 a meta-analysis including 15 observational cohort studies of β-blocker treatment in patients with COPD and revealed that treatment with β-blockers significantly decreased the risk of exacerbation of COPD [10]. Bhatt et al. performed in 2015 a prospective multicenter observational study including subjects with COPD GOLD stage 2 to 4 and found that β-blockers were associated with a significant reduction in

![Fig. 1. Cumulative risk of COPD hospitalizations.](image-url)
COPD exacerbations regardless of severity of airflow obstruction [11]. The mechanisms behind this beneficial effect of β-blockers on risk of COPD exacerbations have not been clarified. However, it has been shown in a murine model that long-term pharmaceutically blocking of pulmonary β-adrenoceptors led to overexpression of β-receptors [17], and that this in turn could decrease bronchoconstriction [18]. Moreover, long-term administration of β-blockers has been shown to reduce airway inflammation and decrease mucus production in another murine asthma model [19].

Even though several studies have indicated a beneficial effect of β-blockers in patients with COPD, there is still a certain reluctance to prescribe β-blockers to patients with airway symptoms such as cough and shortness of breath, or if the patient has a history of cigarette smoking. The explanation behind the 19–7% overall lowered risk of COPD hospitalization found in this study could be that only the healthiest patients are prescribed β-blockers. To avoid for this “healthy user phenomenon” we have looked at the effect of β-blockers after excluding the first 5 years of follow-up. By this, we assume that we are only following patients free of airway symptoms at baseline, as patients already presenting airway symptoms at baseline are at higher risk of COPD hospitalization in the first 5 years. The adjusted HR after exclusion of the first 5 years of follow-up still showed the beneficial association between β-blockers and risk of COPD hospitalization, suggesting that the reduction seen with β-blockers is a class effect.

Another way of adjusting for the healthy user phenomenon is by stratifying for underlying diseases at baseline. In the groups of patients with ischemic heart disease and cardiac arrhythmia, risk of COPD hospitalization was decreased in users of β-blockers compared to users of any other antihypertensive drugs. This could be explained by the cardioprotective effect of β-blockers, as studies have shown that many COPD exacerbations may have cardiovascular causes [11]. The decreased risk for COPD hospitalization in patients with underlying asthma is surprising as many studies have shown significant reduction in lung function in asthmatics treated with β-blockers [20]. An explanation for this finding could be that some patients with known asthma in treatment with β-blockers when hospitalized with airway symptoms are misclassified as asthma hospitalization when they have COPD exacerbation.

Adjusted hazard ratio for death from COPD during follow-up in users of β-blockers was 0.56 (95% CI 0.53–0.59), suggesting that β-blockers are associated with reduced mortality regardless of underlying diseases. Exclusion of the first 5 years of follow-up did not change this association. This is in line with other studies evaluating risk of mortality in COPD patients treated with β-blockers following myocardial infarction, concluding that β-blockers’ apparent benefits on exacerbations are not solely due to a “healthy” user phenomenon [11,12]. Supporting this, the above mentioned meta-analysis by Du et al. found a 28% reduction in overall mortality in COPD patients treated with β-blockers [10]. However, the present study is to our knowledge the first study to compare risk of mortality in patients treated with either β-blockers versus any other antihypertensive drug.

Major strengths of the current study include 1) a very large unselected sample size, even in the subgroups, and 2) data are collected from completed databases with an almost 100% coverage of the Danish population. However, the study has some limitations and biases as well. A review of 1581 patients with COPD from the Danish Patient Registry showed a positive predictive value for COPD of 92% [21], however, studies have shown that a diagnosis based on clinical assessment or based on spirometry may be incorrect, as external obstruction of the small airways by pulmonary congestion, such as in unrecognized heart failure, also causes obstruction [22–24]. Importantly, the results from this study remained when stratifying the analysis for ischemic heart disease and cor pulmonale. As the study is based on register-based data, we cannot know for sure if the enrolled participants are actually taking the drugs. However, the Danish National Prescription Registry includes data on drugs handed out to the recipient in drugstores, with additional information on pack size and price. The findings in this study are therefore based on data on drugs sold to the patients and not only prescriptions made from the general practitioner. We therefore assume that individuals are actually in treatment with the drug if it is bought on a regular basis consistently with pack size, as suggested by Kildemoes and colleagues in 2011 [13]. Another potential limitation is the lack of diagnoses reported by general practitioners. In the present study, only patients hospitalized with COPD were included, which can tend to
underestimate our risk estimate. For this reason, we supplemented the primary endpoint with all-cause mortality and COPD mortality specifically. Furthermore, we do not have data regarding the participants’ smoking habits. Trying to take this into account, we have adjusted for socio-economic data, such as income and employment levels as well as marital status. Several studies have revealed that these socio-economic factors influence smoking status. Pennanen et al. showed that lower education, income, and living without a spouse are factors associated with an increased heaviness of smoking index [25], Zhang et al. found that current smoking is associated with low socio-economic status and being never married, while former smoking is associated with secondary education level, middle-high income, and being widowed [26]. Thus, we do not think that smoking substantially influenced our study results. Finally, included patients are mainly whites of Danish descent, and the results may not automatically apply to other ethnic groups or to other countries with a significant different lifestyle.

To our knowledge, this is the first study to examine the association between β-blocker use and risk of COPD compared specifically with other antihypertensive drugs. This is important because other cardiovascular drugs (ACE inhibitors and angiotensin II receptor antagonists) also seem to have beneficial effects in COPD [27, 28].

In conclusion, the current study has revealed that long-term treatment with β-blockers was associated with reduced risk of COPD hospitalization, all-cause mortality, and death from COPD compared with treatment with other antihypertensive drugs, and thus β-blockers can be considered as safe, even in patients with COPD.

5. Outstanding questions

To further explore whether patients receiving β-blockers have reduced COPD hospitalizations and COPD deaths, a future randomized clinical trial is warranted.

Supplementary data to this article can be found online at https://doi.org/10.1016/j.eclinm.2019.01.004.

Author Contributions

All authors conceived and designed the study. AON, BS, and MD collected the data. AON and LP did the statistical analyses. All authors contributed to data interpretation. AON wrote the original draft of the paper and all authors reviewed and edited drafts and approved the final version for submission. The corresponding author had full access to all the data in the study and had the final responsibility for the decision to submit for publication.

Conflict of Interests

All authors declare no competing interests.

Acknowledgements/Funding

The project was financially supported by The Danish Council for Independent Research in Denmark (grant no. 4183-005698), The Research Foundation of Health Science in Region Zealand (grant no. RSSF2017000661 and no. 15-000342), The Research Foundation of Medical Science (A.P. Müller Foundation) (grant no. 16-68), The Research Foundation in memory of King Christian 10th (grant no. 142/2017), Aase & Ejnar Danielsen’s Research Foundation (grant no. 10-001946), and Lundbeck Foundation (grant no. R252-2017-1690). The funding sources are public or nonprofit organizations and support science in general. They had no influence in paper design, data collection, data analysis, interpretation, or writing of the paper.

References

[1] From the “Global Strategy for the Diagnosis, Management and Prevention of COPD,” Global Initiative for Chronic Obstructive Lung Disease (GOLD). [cited 2018 Jun 24]. Available from http://goldcopd.org/; 2018.
[2] McDonough JE, Yuan R, Suzuki M, et al. Small-airway obstruction and embryogenesis in chronic obstructive pulmonary disease. N Engl J Med 2011;365(17):1567–75.
[3] Hogg JC, A pathologist’s view of airway obstruction in chronic obstructive pulmonary disease. Am J Respir Crit Care Med 2012;186(5):5–7.
[4] Rang H, Dale M, Ritter J, Moore P. Pharmacology. 5th ed.Elsevier Churchill Livingstone; 2003; 797s Bd.
[5] Nielsen AO, Jensen CS, Arredouani MS, Dahl R, Dahl M. Variants of the ADRB2 Gene in COPD: Systematic Review and Meta-Analyses of Disease Risk and Treatment Response. COPD 2017;14:451–60.
[6] Baker JC, Wilcox RG. β-blockers, heart disease and COPD: current controversies and uncertainties. Thorax 2017;72(1):271–6.
[7] Lewis RV, Löffthouse C. Adverse reactions with beta-adrenoceptor blocking drugs. An update. Drug Saf 1993;9(4):272–9.
[8] Clague HW, Ahmad D, Carruthers SG. Influence of cardioselectivity and respiratory disease on pulmonary responsiveness to beta-blockade. Eur J Clin Pharmacol 1984;27(5):517–23.
[9] van der Woude HJ, Zagaarsma J, Postma DS, Winter TH, van Huls I, Aalbers R. Detrimental effects of β-blockers in COPD*: a concern for nonselective β-blockers. Chest 2005;127(3):818–24.
[10] Du Q, Sun Y, Ding N, Lu L, Chen Y. Beta-Blockers Reduced the Risk of Mortality and Exacerbation in Patients with COPD: A Meta-Analysis of Observational Studies. PLoS ONE 2014;9(11):e130484. https://doi.org/10.1371/journal.pone.0130484.
[11] Bhatt SP, Wells JM, Kinney GL, et al. β-Blockers are associated with a reduction in COPD exacerbations. Thorax 2016;71(1):8–14.
[12] Coper S, Görend R, Rossignon P, et al. Association of beta-blocker treatment with mortality following myocardial infarction in patients with chronic obstructive pulmonary disease and heart failure or left ventricular dysfunction: a propensity matched-cohort analysis from the High-Risk Myocardial Infarction Database Initiative. Eur J Heart Fail 2017;19(2):271–9.
[13] Kildehorns HW, Sørensen HT, Hallas J. The Danish National Prescription Registry. Scand J Public Health 2011;39(7 Suppl):38–41.
[14] Pedersen CB. The Danish Civil Registration System. Scand J Public Health 2011;39(7 Suppl):22–5.
[15] Schmidt M, Schmidt SA, Sanagedaal JL, Ehrenstein V, Pedersen L, Sørensen HT. The Danish National Patient Registry: a review of content, data quality, and research potential. Clin Epidemiol 2015;7:449–90.
[16] Helweg-Larsen K. The Danish Register of Causes of Death. Scand J Public Health 2011;39(7 Suppl):22–5.
[17] Callaerts-Vegh Z, Evans KJJ, Dudekula N, et al. Effect of acute and chronic administration of β-adrenoceptor ligands on airway function in a murine model of asthma. Proc Natl Acad Sci U S A 2004;101(14):4948–53.
[18] McGraw DW, Forbes SL, Mak JCW, et al. Transgenic overexpression of β2-adrenergic receptors in airway epithelial cells decreases bronchoconstriction. Am J Physiol-Lung Cell Mol Physiol 2000;279(2):L1379–89.
[19] Nguyen LP, Omokubi O, Paria S, et al. Chronic exposure to beta-blockers attenuates inflammation and mucus content in a murine asthma model. Am J Respir Cell Mol Biol 2008;38(3):256–62.
[20] Morales DR, Jackson C, Lipworth BJ, Donnan PT, Guthrie B. Adverse respiratory effect of acute β-blocker exposure in asthma: a systematic review and meta-analysis of randomized controlled trials. Chest 2014;145(4):779–88.
[21] Thomesen RW, Lange P, Helquist B, et al. Validity and under-recording of diagnosis of COPD in the Danish National Patient Registry. Respir Med 2011;105(7):1063–8.
[22] Güder G, Brenner S, Stöhr S, Hans U, Ruten FH. Chronic obstructive pulmonary disease in heart failure: accurate diagnosis and treatment. Eur J Heart Fail 2014;16(12):1273–82.
[23] Ruten FH, Moons KGM, M-JM Cramer, et al. Recognising heart failure in elderly patients with stable chronic obstructive pulmonary disease in primary care: cross-sectional diagnostic study. BMJ 2005;331(7529):1379.
[24] Brenner S, Güder G, Berliner D, et al. Airway obstruction in systolic heart failure-COPD or congestion? Int J Cardiol 2013;168(3):1910–6.
[25] Pennanen M, Bons Us, Korhonen T, et al. Smoking, nicotine dependence and nicotine intake by socio-economic status and marital status. Addict Behav 2014;39(7):1145–51.
[26] Zhang DM, Hu Z, Orton S, et al. Socio-economic and psychosocial determinants of smoking and passive smoking in older adults. Biomed Environ Sci BES 2013;26(6):453–67.
[27] Vasiileiadis IE, Goudis CA, Giannakopoulou PT, Liu T. Angiotensin converting enzyme inhibitors and angiotensin receptor blockers: a promising medication for chronic obstructive pulmonary disease? COPD 2018;15(2):148–56.
[28] Mancini GB, Enmian M, Zhang N, Levesque LE, FitzGerald JM, Brophy JM. Reduction of morbidity and mortality by statins, angiotensin-converting enzyme inhibitors, and angiotensin receptor blockers in patients with chronic obstructive pulmonary disease. J Am Coll Cardiol 2006;47(12):2554–60.