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李宁 销售工程师 <u>培</u>训经理



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	Home Healthcare Nurse AN: 00004045-201110000-00004 October 2011
	 HERTER, REBECCA; KAZER, MEREDITH; WALLACE PHD, APRN Home Healthcare Nurse AN: 00004045-201006000-00005 June 2010
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以查找2010至2013年出版的有关warfarin treatment for heart failure in men的文章为例



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₩⊠案100: dabigatran heart failure	Qualls LG; Greiner MA; Eapen ZJ; Fonarow GC; Mills RM; Klaskala W; Hernandez AF; Curtis LH. Clinical Cardiology. 36(12):757-65, 2013 Dec. [Journal Article. Research Support, Non-U.S. Gov't]	▲查询相似文献
warfarin 搜 索返回: 46 个文本结果 排序依据: ▼	UI: 24114926 Authors Full Name Qualls, Laura G; Greiner, Melissa A; Eapen, Zubin J; Fonarow, Gregg C; Mills, Roger M; Klaskala, Winslow; Hernandez, Adrian F; Curtis, Lesley H.	Full lext
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· 过滤方式 ∷ ◎ 新增到检索历史	 Warfarin-induced leukocytoclastic vasculitis: a case report and review of literature. [Review] Hsu CY; Chen WS; Sung SH. 	摘要数据 完整数据
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例: warfarin treatment for heart failure in men of 2010-2013

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Abstract

Background—: Aspirin use in heart failure (HF) is controversial. The drug has proven benefit in comorbidities associated with HF; however, retrospective analysis of angiotensin-converting enzyme inhibitor trials and prospective comparisons with warfarin have shown increased risk of morbidity with aspirin use. This study aims to evaluate the association of low-dose aspirin with mortality and morbidity risk in a large community-based cohort.

Methods and Results—: This was a retrospective cohort study of patients attending an HF disease management program. Aspirin use at baseline and its association with mortality and HF hospitalization in the population was examined. Of 1476 patients (mean age, 70.4±12.4 years; 63% men), 892 (60.4%) were prescribed aspirin. Low-dose aspirin (75 mg/d) was prescribed to 828 (92.8%) patients. Median follow-up time was 2.6 (0.8–4.5) years. During the follow-up period, 464 (31.4%) patients died. In adjusted analysis, low-dose aspirin use was associated with reduced mortality risk compared with nonaspirin use (hazard ratio=0.58; 95% confidence interval, 0.46–0.74), and this was confirmed by a propensity-matched subgroup analysis. Low-dose aspirin use was associated with reduced mith reduced risk of HF hospitalization compared with nonaspirin use in the total population (adjusted hazard ratio=0.70; 95% confidence interval, 0.54–0.90). In adjusted analysis, there was no difference in mortality or HF hospitalization between high-dose aspirin users (>75 mg/d) and nonaspirin users.

Conclusions—: In this study, low-dose aspirin therapy was associated with a significant reduction in mortality and morbidity risk during long-term follow-up. These results suggest that low-dose aspirin may have a continuing role in secondary prevention in HF and underline the need for more trials of low-dose aspirin use in HF.

Aspirin use in heart failure (HF) is controversial. The drug has proven benefit in patients with established ischemic heart disease (IHD), a common comorbidity of HF. Aspirin is also recommended in diabetics at high risk of cardiovascular events and as second-line treatment of vascular disorders and atrial fibrillation —all frequently occurring comorbidities in an HF population. However, it has been reported that aspirin use may blunt the beneficial effect of renin–angiotensin–aldosterone system (RAAS) modifying therapy in patients with HF, and several trials have shown increased risk of HF hospitalization when using aspirin. Furthermore, older patients with HF may be at risk of adverse events related to aspirin use especially gastrointestinal hemorrhage.

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Editorial see p 237

Clinical Perspective on p 250



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Results

Patient Demographics

Data were available for 1476 patients with median follow-up time of 2.15 (0.81–4.54) years. Minimum follow-up time was 1 day, and maximum follow-up time was 11.93 years. The mean age of the population

was 70.4±12.4 years, and 930 (63.0%) patie TT1 baseline to 892 (60.4%) patients.

Of those patients prescribed aspirin, 828 (was used by 15 (1.7%) patients and a dose agent was prescribed concomitantly to 16 aspirin to 248 (27.8%) patients, and triple t

Descriptions of the total population, lownonaspirin users are given in Table 1.

Mortality

A total of 464 (31.4%) patients died during 30 (46.9%) high-dose aspirin users, and 19

In unadjusted analysis, there was a lower i than nonaspirin users (hazard ratio [HR]=C In adjusted analysis, low-dose aspirin user compared with nonaspirin users (HR=0.58 adjusted model including age, sex, BNP, cr heart rate, smoking status, comorbidities, dose aspirin use remained significant (HR=

Characteristic	Total Population (N=1476)	Low-Dose Aspirin Users (n=828)	High-Dose Aspirin Users (n=64)	Aspirin Nonusers (n=584)	■ 显示图
Demographics					↓下载Ⅰ
Age, y *t	70.4±12.4	71.9±11.3	71.9±9.8	68.1±13.8	the date
Male	930 (63.0)	533 (64.4)	40 (62.5)	357 (61.1)	國中國稅
Clinical characteristics					◎ 输出至
Systolic blood pressure, mm Hg	127.2±39.0	125.9±22.7	130.4±24.3	125.6±21.2	
Diastolic blood pressure, mmHgtt	73.2±36.7	71.2±13.4	77.0±14.1	72.5±13.3	同新 增到
BNP, pg/mL*	317 (139-668)	352 (157-709)	262 (131-556)	274 (115598)	
Creatinine, µmol/L	103 (87-129)	105 (88-130)	103 (84-141)	101 (85128)	
Ejection fraction, %	40.2±14.5	39.9±14.0	40.2±15.3	40.5±15.1	
HFrEF	797 (64.8)	464 (66.9)	37 (63.8)	296 (61.9)	
Comorbidities					
Ischemic heart disease*	664 (45.0)	482 (58.2)	27 (42.2)	155 (26.5)	
Atrial fibrillation	552 (37.4)	278 (33.6)	21 (32.8)	253 (43.3)	
Chronic obstructive pulmonary disease	164 (11.1)	97 (11.7)	6 (9.4)	61 (10.4)	
Dyslipidemia*	424 (28.7)	275 (33.2)	23 (35.9)	126 (21.6)	
Diabetes meilitus	318 (21.5)	199 (24.0)	15 (23.4)	104 (17.8)	
Hypertension	624 (42.3)	376 (45.4)	26 (40.6)	222 (38.0)	
Peripheral vascular disease	18 (1.2)	12 (1.4)	0	6 (1.0)	
Stroke	47 (3.2)	26 (3.1)	0	21 (3.6)	
Medications					
Loop diuretic	1378 (93.3)	783 (94.6)	58 (90.6)	537 (92.0)	
ACE inhibitor	1242 (84.1)	699 (84.4)	60 (93.8)	483 (82.7)	
Angistensin receptor blocker	403 (27.3)	217 (26.2)	24 (37.5)	162 (27.7)	

In unadjusted analysis, there was no statistically significant difference in mortality between patients with high-dose aspirin use and those with no aspirin use (HR=1.40; 95% CI, 0.95–2.05). In multivariable analysis, there remained no statistically significant difference in mortality between these groups (HR=0.98; 95% CI, 0.59–1.63).

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编辑修改定题通告

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我的课题 我的检索与定题通告 我的期刊目录订阅服务 安捷工具栏					
请注意:执行任何自动定题情报通告中的检索内容会清除当前所有检索历史。若确认执行,请点击"执行检索"。					
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[Use LINK to view the full text] Accession Number 00126334-90000000-98664. Author Luczkowiak, Joanna MSc 1; Martinez-Prats, Lorena MSc 1; Sierra, Olalla 1; Fiorante, Silvana MD 2; Rubio, Rafael MD 2; Pulido, Federico MD 2; Otero, Joaquin R. MD 1; Delgado, Rafael MD 1 Institution [1]Laboratory of Molecular Microbiology, Hospital Universitario 12 de Octubre, Madrid, Spain (2)HIV Unit, Hospital Universitario 12 de Octubre, Madrid, Spain Title Lack of the detection of XMRV or Polytropic MLV-related sequences in blood cells from HIV-1 infected patients in Spain.[Article] Source JAIDS Journal of Acquired Immune Deficiency Syndromes. Status Publish Ahead of Print, POST ACCEPTANCE, 29 September 2011 Abstract Background: Xenotropic murine leukemia virus-related virus (XMRV) and polytropic MLV-related virus are recently described human gammaretroviruses that have been associated with prostate cancer and chronic fatigue syndrome (CFS). These studies have been controversial since a number of laboratories have been unable to find evidence of XMRV in similar groups of patients or controls. Since the existence of XMRV raises many questions, we decided to study its presence in a group of patients infected with HIV-1 with a high proportion of intravenous drug use (IDU) and co-infection by HCV. (C) 2011 Lippincott Williams & Wilkins, Inc. DOI Number
10.1097/QAI.0b013e318238b596 This link leads to available full-text or the complete reference. http://wiidca.ouid.com/ouidurb.coi/T_IC_MODE=prid@DACE=fulleart@DECE=7004E5848/7c1
(2> [Use LINK to view the full text] Accession Number 00126334-90000000-98652. Author Mayer, Kenneth H. MD 1; Ducharme, Robert BA 2; Zaller, Nickolas PhD 3; Chan, Philip A. MD 4; Case, Patricia ScD 5; Abbott, David 6; Irma, Rodriguez MS 7; Cavanaugh, Timothy MD 8 Institution (1)The Fenway Institute, The Miriam Hospital, Beth Israel Deaconess Medical Center, Harvard Medical School (2)The Miriam Hospital



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