The Risk of Prostate Cancer in Pilots: A Meta alysi.

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BACKGROUND: Aviation exposes pilots to various occupationally related hazards, including sizing radiate and sinical combustion. The possible increased risk of prostate cancer among pilots in comparise significant and subject of debate. This systematic review and meta-analysis aimed to determine the quality supporting evidence and magnitude of this association.

METHODS: All studies pertaining to prostate cancer in pilots were retrieved study that assessed the incidence of prostate cancer relative regardless of language or size. A random effect model was was assessed using the Q statistic and I².

tom multiple database and from a manual search. Any the incidence in the general population was included to pool release risks (RR) across studies. Heterogeneity

- **RESULTS:** Eight studies with a low risk of bias were include the meta-a. In a had an increased risk of developing prostate cancer compared to the general population of 95% controlence interval (Cl), 1.5–2.7]. The analysis was associated with substantial heterogeneity ($l^2 = 7$). Several arroups had significantly increased risk, such as African American pilots (RR 10.00; 95% Cl, 5.04–19.86) and the pilots (RR 3.30; 95% Cl, 2.03–5.39).
- **CONCLUSION:** Pilots are at least twice as likely to the prostate cer compared to the general population. The implications of these findings are important concering thigh provide the prostate cancer and the large number of pilots in the workforce.

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KEYWORDS: aviation, pilots, prostate cer i

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viati lave been examined in regards arious aspecu to pilot health, s analyz' the specific risk facay ¹ tors to the development of ors the le utility of electrocardioheart dise erstano. or u grams, t X-ra and MRIs for screening purposes in this also been made to try to underpopulation. stand if pilot. at an increased risk for the development of cancers. It has be roposed that aviators are exposed to potential carcinogens such as ionizing radiation during flight²² and jet fuel combustion products.¹⁹ Nonionizing electromagnetic fields¹⁰ and disruption of the circadian rhythm²⁸ are also potential contributing factors. It is important to understand whether pilots are at an increased risk for certain diseases based on occupational exposures so their health status can be properly evaluated, maintained, and when necessary, treated. Prostate cancer is one of the malignancies that has been investigated in the literature.

Prostate cancer is the second most common type of male cancer worldwide. The most recent data from 2012 estimated

that there were 1.1 million cases and over 307,000 deaths worldwide.³² In the United States, the risk of developing prostate cancer is estimated to be one in six.²⁶ This cancer is particularly relevant to the field of aviation since about 95% of pilots in the United States are male.⁷ Moreover, prostate cancer is also strongly associated with age.^{14,21} As populations continue to age and the public use of aviation-based transport continues to rise, the average age of pilots will continue to increase. Over the last 20 yr in the United States, the average age of pilots has increased from 40.5 to 44.7 according to the Federal Aviation Administration.^{8,9} If this trend continues, the incidence of prostate cancer will continue to increase. It is

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imperative to understand if a pilot's occupational exposures further increase the risk of prostate cancer.

The increased risk of prostate cancer among aviation pilots is a subject of debate because there is ambiguity in the literature. Some studies suggest that they are indeed at an increased risk while other studies seem to suggest that they are not.^{1,11} Determining the incidence of prostate cancer in pilots compared to the general population is important to advance our understanding of the potential risks as well as to help inform policies and screening protocols specific to aviators. Therefore, the aim of this study was to perform a systematic review and meta-analysis to determine whether pilots are at an increased risk of developing prostate cancer compared to the general population.

METHODS

This study was conducted according to guidance from the Cochrane Handbook of Systematic Reviews and is reported according to preferred reporting items for systematic reviews and meta-analyses recommendations (PRISMA).^{15,20}

Literature Search

A comprehensive literature search of several databases was r formed from each database's inception to November 2013 any language. The databases included Ovid Medline in-proces and other nonindexed citations, Ovid Medline, and PubMed. An experienced librarian from Mayo Clinic de d conducted the search strategy with input from gators. *y* inve This search was duplicated by an experience. rari Civil Aviation Medical Institute at the .eral A on Adm. istration to ensure the complete of the sea. protocol. Both librarians used controlled v supplem d with d the incidence of keywords to search for studies that as. also manual prostate cancer in pilots rched PubMed, Ovid Medline, and th efense Sechnical Internation Center, pert int articles to ensure the comand crossed-referen. pleteness of the search lol.

Study Sei .ion

All st. s wer onsidered regardless of publication lanaies were eligible if they compared .0015. guage or prostate cancer in pilots to the general popthe inciden. ulation. Abstra and titles that resulted from executing the search strategy were independently evaluated by two reviewers for potential eligibility, and the full text versions of all potentially eligible studies were obtained. Two reviewers working independently considered the full text reports for eligibility. Disagreements were harmonized by consensus and, if not possible by consensus, through arbitration by a third reviewer.

Data Extraction

Information on the studies' characteristics and demographics was recorded, such as authors, publication year, country, number of years in the evaluation, type of pilot population studied, and outcome. The incidence of prostate cancer was reported as either a standardized incidence ratio (SIR) or as an incidence rate ratio (IRR) in all studies.

Assessment of Methodological Quality (Risk of Bias)

The methodological quality of the included studies was assessed by using the Newcastle-Ottawa scale.³¹ This scale consists of three domains (cohort selection, comparability of outcome) and evaluates each study's overall risk operator. Two revers independently assessed the quality of constudy.

Outcome Definition

The primary outcome, incident of presente can use defined as new onset prostate can use urine the study period as determined by public restines. Incident the standardized to the spective point tion determine the SIR or IRR.

Statistic ' ' ' ysis and Su up Analysis

or IKR was retrieved om each study as well as the 90% The S or 9 confidence interval (CI) from each study. The I² statistic d to estim the percentage of total variation across was to het seneity rather than chance (ranging from studie alues of \leq 25%, 50%, and \geq 75% represent 0 to 100 × moderate, and high inconsistency, respectively. The ranmodel was used to pool results, thereby accounting de variance between studies.⁶ This model was chosen because f the anticipated significant heterogeneity between the studies. omprehensive Meta-Analysis, version 2 (Englewood, NJ) was used for statistical analysis. All P-values are two tailed and the threshold for significance was set at P < 0.05.

The a priori hypothesis is to conduct subgroup analysis based on race (white or African American), the type of pilot (military or civilian), and estimated exposure to radiation (low, medium, or high). Although the SIR and IRR are both relative effects measures (risk ratios) and may approximate each other, their estimation methods differ. Therefore, using subgroup analysis, we explored whether the pooled effect size differed between studies reporting IRR and SIR. The relative estimates from subgroups were compared using the ANOVA test to determine if a statistically significant difference was present among the estimates derived from each subgroup.

RESULTS

The initial search resulted in 44 publications and, after abstract and full text reviews, 8 studies met the inclusion criteria (**Fig. 1**). More than 128,000 pilots were evaluated. The year of publication ranged from 1996 to 2011, and earliest data included in the studies were from 1946. Three studies took place in North America, while the remaining five took place in Europe.

Table I shows details of the baseline characteristics of the included studies. Risk of bias of the included studies was found to be low according to the Newcastle-Ottawa quality



Fig. 1. Flowchart showing the literature search yield and selected studies.

assessment scale. Pilots were twice as likely to develop protate cancer compared to the general population (RR 2.0; 95) CI, 1.5–2.7). The analysis was associated with high heterogeneity ($I^2 = 79\%$) that was explained by subgroup analysis (**Fig. 2**).

Table I. Study Characteristics.

Studies that reported an SIR had an RR of 1.36 (95% CI, 1.18-1.56) compared to the study that reported IRR which had an RR of 3.84 (95% CI, 2.40-6.13). The RR of 2.56 (95% CI, 2.01-3.27) in whites was lower than that in African Americans who had an RR of 10.00 (95% CI, 5.04-19.86). Civilian pilots had an RR of 1.36 (95% CI, 1.01-1.83) while those with military backgrounds had an RR of 3.30 (95% CI, 2.03-5.39). Lastly, the estimated radiation exposure risk was analyzed in terms of low, moderate, and highed in the original studies. There was no statistical¹ 'iffer-.gnifica analysis 1 ence among these subgroups. All subs ımmarized in Table II.

DISCUSSION

This systematic revie nd meta-a. is r' vs that pilots have twice the risk for opment of h .te cancer as the genubgroups analyzed, military pilots eral population. ...mong and Africe erican pile nd an even higher risk. It was ed that there was angher risk in the study that also n IRR compared to the studies that reported a SIR. The repor reasoi r this is the ht to be that the study which reported IRR wa only st that included men of African ancestry. d the highest risk of all the subgroups that This subgr palyzed, and African ancestry is a known strong risk factor ' ppment of prostate cancer.¹⁸

athough some of the risk factors for prostate cancer are own, the etiology of this disease process is still poorly

STUDY	PILOT POPULATION	TY LENGTH	IZE	SERVICE	TYPE OF PILOTS	AGE RANGE	RISK FACTORS
Band 1996 ¹	Canada	1992	2680	Civilian	Professional and General	Not Specified	Radiation exposure
del Junco 2011 ⁵	U.S. A	1991-2	337	Military	Professional	35-64	Race, age
Gundestrup 1999 ¹¹	D ark	1921–1995	3790	Civilian	Professional and General	Not Specified	Type of aircraft, flight hours, radiation exposure
Haldorsen 2°	prway	1946–1994	3815	Civilian	Professional and General	Not Specified	Radiation exposure, smoking status
Hammar 200₂	Sweae.	1957–1994	105,025	Military and Civilian	Professional and General	20-80+	Service branch, flight hours, altitude, distance
Pukkala 2002 ²³	Denmark, Finland, Iceland, Norway, and Sweden	1946–1997	10,032	Civilian	Professional	Not Specified	Flight hours, radiation exposure, circadian rhythm disturbance, smoking status
Rafnsson 2000 ²⁴	Iceland	1955–1997	458	Civilian	Professional and General	Not Specified	Flight hours, radiation exposure, circadian rhythm disturbance
Yamane 2006 ³³	U.S. Air Force	1989–2002	1959	Military	Professional	17–60	Age

Note: For all studies the history of cancer and race (% white) was not specified.

Model	Study name					
		Risk ratio	Lower limit	Upper limit		
	Band et al.	1.9	1.3	2.7	- 1	
	del Junco et al. a	2.0	1.3	3.2		
	del Junco et al. b	7.1	2.0	24.5		
	del Junco et al. c	2.6	1.7	4.0		
	del Junco et al. d	3.0	2.0	4.6		
	del Junco et al. e	12.1	3.9	37.2		
	del Junco et al. f	11.2	3.3	37.3		
	Gundestrup et al.	0.8	0.2	3.2		
	Haldorsen et al. a	0.7	0.2	2.5		
	Hammar et al.	1.2	0.9	1.6		
	Pukkala et al.	1.2	0.9	1.6		
	Rafnsson et al.	1.3	0.4	4.0		
	Yamane	1.4	1.2	1.7		
Fixed		1.6	1.4	1.8		
Random		2.0	1.5	2.7		
I^2	=79%				0	

Risk ratio and 95% CI



Meta Analysis

Fig. 2. Forest plot and overall study analysis.

understood. There is little in the literature about what migh increase prostate cancer risk in pilots. Some hype suggest exposure to ionizing radiation during flight,² fuer mbustion products,¹⁹ electromagnetic fields,¹⁰ a of cir-'isrupt' cadian rhythm²⁸ are plausible causes, nsi risk factors, del Junco et al.⁵ ascertaj a higher of prostate cancer among a subgroup of American ts. The socioeconomic status of pilots right ibly be another risk factor, but this is not well. erstood.25

This study was not signed to answer the destion of causality and is therefore inable in hed light on potential etiologies. Future studies the readed to try to determine the reason that pilots have a preased is of developing prostate er. , ne meantime, it may be prudent to consider whether nore aggressive screening practices might be necessary for viator populations.

It is important to note that studies which assessed the mortality of pilots did not find an increase in mortality due to prostate cancer.^{2,3} This may suggest that the increased incidence in pilots is because they are more frequently examined than the general population. However, since screening for prostate cancer during a flight physical is not required, this hypothesis is unlikely to account for the entire increase in incidence seen in this study. Another possible explanation could be that pilots live longer since they are healthier than the general population and prostate cancer is a disease of old age. However, the incidence

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COVARIATL		NO. COHORTS	EFFECT SIZE	LOWER LIMIT	UPPER LIMIT	l ^{2%}	<i>P</i>-VALUE FOR DIFFERENCE	
Effect size type	e							
IRR		6	3.84	2.40	6.13	67.93	0.01	
SIR		7	1.36	1.18	1.56	13.78		
Race								
Black		3	10.00	5.04	19.86	0.00	0.01	
White		3	2.56	2.01	3.27	0.00		
Pilot type								
Civilian		5	1.36	1.01	1.83	28.94	0.01	
Military		7	3.30	2.03	5.39	85.31		
Estimated rad	iation exposure							
Low		6	0.92	0.64	1.33	0.00	0.28	
Medium		3	1.08	0.63	1.86	0.00		
High		6	1.32	1.03	1.69	0.00		

Table II.

IRR = incidence rate ratio; SIR = standardized incidence ratio.

was standardized by age, which reduces the impact of confounding by age. Lastly, errors in the ascertainment of cause of death in observational studies are common. Future research may better clarify whether prostate cancer mortality in pilots is different from that of the general population.

The results derived from observational studies are subject to confounding. Additionally, there was high heterogeneity between studies. Our a priori analysis explains this heterogeneity. It is most likely due to the diversity of the populations included in the individual studies as well as the variance in when the data were collected. Some studies included data from 1946 while others included data only from 1991. Another limitation is that in one study, there is a potential for overlap of patients among the different cohorts.²³ Since the majority of the studies included only pilots from within their own countries, this limitation is not a concern in other studies.

The strengths of this review include the exhaustive and reproducible search strategy, inclusion of non-English studies, and a large sample size of over 128,000 pilots from 8 studies. Most previous articles that addressed the question of whether pilots are at an increased risk of developing prostate cancer did not focus specifically on prostate cancer but rather on cancers in general. Therefore, they would include at most two or three articles on prostate cancer and conclude that the data were mixed. To our knowledge, this is the largest systematic revier that has been performed to date for answering the question of whether pilots are at an increased risk for developing prostate cancer.

Consideration must be given to screening for cancer in pilots. The U.S. Preventive Service .sk Fo has recently recommended against routine scree o fo cancer using prostate-specific antigen only atory tes that can be used as a screening to Many wo assume e applied i that this recommendation sho ilots. This might be reasonable if ilous had a rage risk. However, it appears that they twice as like. develop this malignancy.

ed for more studies on this This review high. ∖th∕ subject. We need to be. derstap hy aviators are at an eff increased ris¹ ler to 1. vely preserve the health status of p or more investigative work s. An her incen is that • study av actually underestimate the increased r incidence has been rising over risk of pile the years part. but not completely, due to increased screening efforts. The suby del Junco et al.⁵ seems to suggest that, over time, aviators are developing prostate cancer at an even faster rate than the general population. If this is true, then studies which rely upon data from the mid-1900s might not truly represent the increased risk that is now present in the early 2000s. Given the prevalence of prostate cancer in the general population and the elevated at-risk status of pilots, it is imperative that we gain a more robust understanding of the true risk and the mechanisms underlying that risk. Lastly, shared decision-making tools are needed to communicate the risk of prostate cancer to pilots and aid them in the decision regarding screening.

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