- 38) 高橋裕樹、KYSS Study Group. 全身性エリテマトーデスの症例対照研究. 第50回日本リウマチ学会総会学術総会、2006年4月(長崎)
- 39) 山口将平、吉田大輔、橋爪誠. 本邦における 特発性門脈圧亢進症の食道静脈瘤に対する治療 成績の検討-全国調査結果報告. 第13回日本門 脈圧亢進症学会総会、2006年9月(東京)
- 40) 仁科基子、柴崎智美、太田晶子、石島英樹、 泉田美知子、永井正規. 潰瘍性大腸炎医療受給 者の年次変化. 第 65 回日本公衆衛生学会、2006 年 10 月(富山)
- 41) 伊津野孝、杉田稔、玉腰暁子、永井正規、稲葉裕. 血栓性血小板減少性紫斑病 (TTP) /溶血性尿毒症症候群 (HUS) の全国患者数調査. 第 65 回日本公衆衛生学会、2006年10月(富山) 42) 土井由利子、横山徹爾. 過去 33 年間 (1972-2004年) における難病による死亡の推移に関する検討. 第 65 回日本公衆衛生学会、2006.10.25; 富山. 同抄録集 p.638.
- 43) 黒沢美智子、稲葉裕、金子史男、永井正規. ベーチェット病の臨床調査個人票データの分析. 第 65 回日本公衆衛生学会総会、2006 年 10 月(富山)
- 44)太田晶子、柴崎智美、仁科基子、石島英樹、 泉田美知子、永井正規. 呼吸器系難病患者の日 常生活への支障の程度の検討-臨床調査個人票 の解析-. 第 65 回日本公衆衛生学会、2006 年 10 月(富山)
- 45) 柴崎智美、太田晶子、石島英樹、泉田美知子、 仁科基子、永井正規. 免疫系難病患者の日常生 活への支障の程度の検討-臨床調査個人票の解 析-. 第 65 回日本公衆衛生学会、2006 年 10 月 (富山)
- 46) 黒沢美智子、稲葉裕、松葉剛. ベーチェット 病患者の QOL 調査. 第 71 回日本民族衛生学会 総会、2006 年 11 月 (那覇)
- 47)縣 俊彦、稲葉裕、黒沢美智子. 神経線維腫症Ⅱ型公費負担対象者の特性. 第71回日本民族

- 衛生学会、2006年(那覇)
- 48) 黒沢美智子、稲葉裕、永井正規. 天疱瘡(稀少難治性皮膚疾患)の臨床調査個人票電子化データの分析. 第 17 回日本疫学会総会、2007 年 1月(広島)
- 49) 稲葉裕、黒沢美智子、小林茂人. 日欧の抗好中球細胞質抗体(ANCA)関連疾患の Incidence と Prevalence の比較. 第 17 回日本疫学会学術総会、講演集、p193,2007 (広島)
- 50)縣俊彦、稲葉裕、黒沢美智子. 2 つの全国規模調査から見た神経線維腫症Ⅱ型患者の特性. 第 17 回日本疫学会学術総会、2007 年1月(広
- 島) 51)大浦麻絵、大西浩文、坂内文男、森満、玉腰 暁子、永井正規、井廻道夫、坪内博仁、大西三 朗、全国疫学調査による難治性肝疾患の全国患
- 者数の推定. 第 17 回日本疫学会、2007 年 1 月 (広島)
- 52) 坪井一哉、鈴木貞夫. ライソゾーム病患者における公費負担医療に対する調査. 第17回日本疫学会、2007年1月(広島)
- 53) 岡本和士、柴崎智美、永井正規. パーキンソン病患者数増加に関する記述疫学的検討. 第 17 回日本疫学会、2007(広島)
- 54) 鷲尾昌一、KYSS Study Group. 全身性エリテマトーデスの発症の関連要因: Kyushu Sapporo SLE (KYSS)study. 第77回日本衛生学会総会、2007年3月(大阪)
- 55) Michiko Kurosawa, Yutaka Inaba, Tsuyoshi Matsuba, Akiko Tamakoshi, Fumio Kaneko, Akiko Nishibu, Takashi Kawamura. QOL study of Behcet's disease patients in Japan. Rheumatology Supple42, S15, 2006.
- 56) Yutaka Inaba, Michiko Kurosawa, Fumio Kaneko, Tomohiko Makino, Masaki Nagai. Analysis of the electronic clinical database (2001-2004) of patients with Behchet disease receiving financial aid for treatment in Japan. Rheumatology Supple42,

S16, 2006.

57) M Kurosawa, Y Inaba, T Matsuba, A Tamakoshi, F Kaneko, A Nishibu, T Kawamura.

QOL study of Behchet's disease patients in Japan.

12th International Conference on Behchet's Diseases, 20-23, Sep.2006, (Lisbon, Portugal) 58) Y Inaba, M Kurosawa, F Kaneko, T Makino, M Nagai. Analysis of the electronic clinical database of patients(2001-2004) with Behchet's disease receiving financial aid for treatment in Japan. 12th International Conference on Behchet's Diseases, 20-23, Sep.,2006(Lisbon, Portugal)

59) Toshihiko Agata, Hidesuke Shimizu,

Hirofumi Takagi, Yutaka Inaba, Akiko Tamakoshi, Michihito Niimura, A STUDY OF LISCH NODULES (LN) AND NEUROFIBROMATOSIS 1 IN JAPAN. 17th International Congress of eye research. 2006.10 (Buenos Aires Alzentin)

- 60) Okamoto K, Kihira T, Kondo T. Lifestyle factors and risk of amyotrophic lateral sclerosis; A case-control study in Japan. 17th International Symposium on ALS/MND (Yokohama). 2006.
- 61) Tameko Kihira, Kanno S, Miwa H, Kondo T, Okamoto K. Evaluation of the role of exogenous risk factors in patients with amyotrophic lateral sclerosis. 17th International Symposium on ALS/MND. Dec. 2006, Yokohama, Japan.
- 62) Kobashi G, Ohta K, Hata A, Washio M, Okamoto K, Japan OPLL Epidemiological Study Group. Genetic and acquired factors for ossification of the posterior longitudinal ligament of the spines in Japan; a case-control study. American Society of Human Genetics, October, 2006 (New Orleans, USA)

(平成 19 年度)

63) 黒沢美智子、稲葉裕、金子史男、永井正規. 臨床調査個人票データベースを用いたベーチ エット病の予後の検討.第77回日本衛生学会、 2007(総会講演集,p699)

- 64) 坪井一哉、片岡智史. 遅発型 Pompe 病に対する酵素補充療法: 症例報告. 第 76 回日本交通医学会東海北陸地方会、2007 (名古屋) 65) 坪井一哉. ゴーシェ病における NK 細胞活性および LAK 細胞活性の定量的検討. 第 49 回日本先天代謝異常学会、2007 (山形)
- 66) 片岡智史、坪井一哉. 糖原病 II 型 (Pompe 病) における酵素補充療法の導入. 第 86 回東海臨床遺伝代謝懇話会、2007 (名古屋)
- 67) 坪井一哉. ゴーシェ病における NK 細胞活性および LAK 細胞活性の検討. 第 69 回日本血液学会、第 49 回日本臨床血液学会合同総会 2007 (横浜)
- 68) 坪井一哉. ファブリー病 11 症例に対する 酵素補充療法の検討. 第 104 回日本内科学会 講演会、 2007 (大阪)
- 69) 黒沢美智子、稲葉裕、玉腰暁子、金子史男、 西部明子、川上佳男、川村孝. ベーチェット 病の QOL フォローアップ調査. 第 66 回日本 公衆衛生学会、2007(総会抄録集 p398). (松 山)
- 70) 稲葉裕、黒沢美智子、小林茂人、尾崎承一、 永井正規. 臨床調査個人票データベースを用 いた難治性血管炎 (悪性関節リウマチ、大動 脈炎症候群) の特徴. 第66回日本公衆衛生学 会、2007(総会抄録集 p398). (松山)
- 71) 土井由利子、横山徹爾、酒井美良、中村好一. クロイツフェルト・ヤコブ病による死亡の時系列推移. 第 66 回日本公衆衛生学会2007.10 同抄録集 p.241 (松山)
- 72) DUKI M. I. Zulkarnain, DOI Yuriko, SAKAI Miyoshi, YOKOYAMA Tetsuji. Geographical Analyses of Mortality Caused by Severe Acute Pancreatitis, 1995-2004, in Japan. 第 66 回日本公衆衛生学会、2007.10 同抄録集. p.241-2(松

山)

73) 石島英樹、仁科基子、柴崎智美、太田晶子、 泉田美知子、永井正規. パーキンソン病患者の 重症度別の日常生活の状況、社会保障制度の利 用状況、治療状況. 第 66 回日本公衆衛生学会総 会、2007.10 (松山)

74) 仁科基子、太田晶子、柴﨑智美、石島英樹、 泉田美知子、永井正規. 潰瘍性大腸炎医療受給 者の臨床像の年次変化. 第 66 回日本公衆衛生学 会総会、2007.10 (松山)

75) 眞崎直子、松田智大、坂田清美、他. 神経難病患者(脊髄小脳変性症)の QOL と保健福祉サービスのニーズに関する研究. 第66回日本公衆衛生学会、2007.10. (松山)

76) 西川浩昭、縣俊彦、稲葉裕、黒沢美智子. 神経線維腫症 1 公費患者の 10 年の変遷. 第 71 回日本民族衛生学会、2007.11 第 73 巻付録 p60-1 (高岡)

77) 縣俊彦、西川浩昭、稲葉裕、黒沢美智子. 神経線維腫症 2 公費患者の 10 年の変遷. 第 71 回日本民族衛生学会、2007.11 第 73 巻付録 p62-3 (高岡)

78)鷲尾昌一、KYSS Study Group. 全身性エリテマトーデス発症の関連要因:食事因子を中心に:KYSS study. 第 18 回日本疫学会、2008.01 (東京). KYSS Study Group (Kyushu Sapporo SLE study group)の構成員は以下のとおりである。鷲尾昌一、清原千香子、堀内孝彦、塚本浩、原田実根、浅見豊子、佛淵孝夫、牛山理、多田芳史、長澤浩平、児玉寛子、井手三郎、小橋元、岡本和士、阪本尚正、佐々木敏、三宅吉博、横山徹爾、大浦麻絵、鈴木拓、森満、高橋裕樹、山本元久、阿部敬、稲

葉裕、永井正規.

79) 土井由利子、横山徹爾、酒井美良、中村好一、丹後俊郎. クロイツフェルト・ヤコブ病による死亡の地域集積性に関する検討. 第 18 回日本疫学会学術総会、2008.1 (東京)

80)島田直樹、中尾眞二、小澤敬也、永井正規、 杉田稔. 臨床調査個人票からみた再生不良性 貧血の特徴. 第 78 回日本衛生学会総会、2008.3 (熊本)

81) T Agata , H Nishikawa, Y Inaba, M Kurosawa, M Nagai. A Nation-Wide Epidemiological Study of COPD and Tobacco Smoking Rate in Japan. 6th Conference of International Society for the Prevention of Tobacco Induced Disease.2007.11 Little Rock AS USA

82) Miura K, Matsumori A, Morikawa Y, Inaba Y, Nagai M, Nakagawa H. Predictive score to predict the prognosis of cardiomyopathies: from a nationwide study in Japan. The 1st International Congress of Cardiomyopathy and Heart Failure, 2007 (Kyoto)

83) Kihira T, Hama K, Nakanishi I, Hiwatani Y, Kazimoto T, Okawa M, Miwa M, Okamoto K, Kondo T. Longitudinal changes in age at onset and survival in patients with amyotrophic lateral sclerosis in w prefecture. 18th International Symposium on ALS/MND 2007. (Tront)

- G. 知的所有権の取得状況(予定を含む)
- 1. 特許取得 なし
- 2. 実用新案登録 なし
- 3. その他 なし

Ⅱ. 研究成果の刊行に関する一覧表

研究成果の刊行に関する一覧表(平成17~19年度)

書籍

書籍							•
著者氏名	論文タイトル名	書籍全体の編 集者名	書籍名	出版社名	出版地	出版年	ページ
松森昭 三浦克之 中川秀昭	心筋症の疫学	松森昭	新・目でみる 循環器病シ リーズ15 心筋症	メジカルビュー 社	東京	2007	20-24
雑誌				•			
発表	長者氏名	論文タイトル名		発表誌名	巻号	ページ	出版年
Miyake Y, Sasaki S, Yokoyama T, Chida K, Azuma A, Suda T, Kudoh S, Sakamoto N, Okamoto K, Kobashi G, Washio M, Inaba Y, Tanaka H		environmental factors and		Ann. occup. Hyg.	49(3)	259-265	2005
Miyake Y, Sasaki S, YokoyamaT, Chida K, Azuma A, Suda T, Kudoh S, Sakamoto N, Okamoto K, Kobashi G, Washio M, Inaba Y, Tanaka H, Japan Indiopathic Pulmonary Fibrosis Study Group		Case-control study of medical history and idiopathic pulmonary fibrosis in Japan		Respirology	10	504-509	2005
黒沢美智子 稲葉裕		Behçets 病の 像の動向)最近の疫学	医学のあゆみ	215(1)	5-8	2005
Kihira T, Yoshida S, Hironishi M, Miwa H, Okamoto K, Kondo T		C hanges in the amyotrophic lat Wakayama,Japa	eral sclerosis in	Amyotrophic Lateral Sclerosis	6	155-163	2005
柴崎智美 永井正規 渕上博司 仁科基子 太田晶子 川村孝 大野良之		特定疾患治療の受給者の経年3 調査リンケージ た解析	变化一受給者	日本公衆衛生雑 誌	第52巻 第12号	1009-1020	2005
岡本和士 紀平為子 近藤智義 阪本尚正 小橋元 鷲尾昌一 三宅吉博 横山徹爾 佐々木敏 稲葉裕 要因に関する検討		化とその関連	厚生の指標	52(5)	29-33	2005	
Sakauchi F, M Toda G	Iori M, Zeniya M,	Antimitochondr negative primar cirrhosis in Japa clinical data wh applied to receiv financial aid	y biliary an: utilization of en patients	Journal of Epidemiology	16(1)	30–34	2006
柴崎智美 永	井正規	男性の全身性 デス;臨床症 差		じほう	Vol.3 No.6	659-663	2006

Washio M, Horiuchi T, Kiyohara C, Kodama H, Tada Y, Asami T, Takahashi H, Kobashi G, Abe T, Tanaka H, Nogami N, Harada M, Tsukamoto H, Ide S, Nagasawa K, Ushiyama O, Hotokebuchi T, Okamoto K, Sakamoto N, Sasaki S, Miyake Y, Yokoyama T, Mori M, Oura A, Sinomura Y, Suzuki H, Yamamoto M, Inaba Y, Nagai M	Smoking, drinking, sleeping habits, and other lifestyle factors and the risk of systemic lupus erythematosus in Japanesefemales: findings from the KYSS study	Rheumatology	16	143–150	2006
石島英樹 永井正規	パーキンソン病関連疾患の 性差	じほう	Vol.3 No.7	763—768	2006
太田晶子 永井正規	サルコイドーシスの臨床症 状における性差	じほう	Vol.3 No.8	873—877	2006
Wakai K, Kawamura T, Endoh M, Kojima M, Tomino Y, Tamakoshi A, Ohno Y, Inaba Y, Sakai H	A scoring system to predict renal outcome in IgA nephropathy: from a nationwide prospective study	Nephrol Dial Transplant	21	2800-2808	2006
土井由利子 横山徹爾 川南勝彦 石川雅彦	特定疾患治療研究対象疾患 と国際疾病分類(ICD- 10,9,8)に基づく死因コード の対応	日本公衆衛生雑誌	第53巻 第10号	777-786	2006
太田晶子 永井正規 仁科基子 柴﨑智美 石島英樹 泉田美知子	臨床調査個人票から得られ る難病の発病年齢	日本公衆衛生雑誌	第54巻 第1号	3-14	2007
太田晶子 永井正規 仁科基子 柴﨑智美 石島英樹 泉田美知子	特定疾患医療受給者の実態- 疾患別・性・年齢別受給者 数とその時間的変化	日本公衆衛生雑誌	第54巻 第1号	32-42	2007
泉田美知子 永井正規 仁科基子 柴﨑智美 太田晶子 石島英樹	特発性血小板減少性紫斑病 患者の初期の臨床症状とそ の時間的変化	日本血栓止血学会誌	第18巻 第1号	53-60	2007
大浦麻絵	全国疫学調査による難治性 の肝疾患の日本の患者数推 定	肝胆膵	第54巻 第3号	425-430	2007
Nasermoaddeli A, Miura K, Matsumori A, Soyama Y, Morikawa Y, kitabatake A, Inaba Y, Nakagawa H	Prognosis and prognostic factors in patients with hypertrophic cardiomyopathy in Japan: results from a nationwide study	Heart	93	711-715	2007
福島若葉 大藤さとこ 廣田良夫	特発性大腿骨頭壊死症の疫 学	Clinical Calcium	17(6)	11-16	2007

Doi Y, Yokoyama T, Sakai M , Nakamura Y	Creutzfeldt-Jakob disease mortality in Japan, 1979-2004: Analysis of national death certificate data	Journal of Epidemiology	17(4)	133-139	2007
土井由利子 横山徹爾 酒井美良	日本における難病による死 亡の時系列推移(1972~2004 年)	日本公衆衛生雑 誌	第54巻 第10号	684-694	2007
Okamoto K, Kihira T, Kondo T, Kobashi G, Washio M, Sasaki S, Yokoyama T, Miyake Y, Sakamoto N, Inaba Y, Nagai M	Nutritional status and risk of amyotrophic lateral sclerosis in Japan	Amyotrophic Lateral Sclerosis	8(5)	300-304	2007
Sakauchi F, Oura A, Ohnishi H, Mori M	Comparison of the clinical features of Japanese patients with primary billiary cirrhosis in 1999 and 2004: utilization of clinical data when patients applied to receive public financial aid	Journal of Epidemiology	17(6)	210-214	2007
三徳和子 松田智大 新城正紀 眞崎直子 平良セツ子 丹野高三 簑輪眞澄 坂田清美	難病疾患患者における包括 的QOLの特徴と類似点	川崎医療福祉学 会誌	17(2) 掲載予定		2007
Miura K, Matsumori A, Nasermoaddeli A, Soyama Y, Morikawa Y, Sakurai M, Kitabatake A, Nagai M, Inaba Y, Nakagawa H	Prognosis and prognostic factors in patients with idiopathic dilated cardiomyopathy in Japan: results from a nationwide study	Circ J	(in press)		
Okamoto K, Kihira T, Kondo T, Kobashi G, Washio M, Sasaki S, Yokoyama T, Miyake Y, Sakamoto N, Inaba Y, Nagai M	Lifestyle factors and risk of amyotrophic lateral sclerosis in Japan	Amyotroph Lateral Scler	(in press)		

Ⅲ. 研究成果の刊行物・別刷

Ⅲ 疫学

心筋症の疫学

松森 昭 京都大学大学院医学研究科循環器內科学 三浦克之,中川秀昭 金沢医科大学公衆衛生学

1998年まで心筋症に関する系統的な疫学調査は行われていなかったが、その年わが国における特発性心筋症の疫学像を明らかにする目的で、厚生省特定疾患特発性心筋症調査研究班と特定疾患の疫学に関する研究班の共同研究が行われた。その後も共同研究が継続され、心筋症の5年生存率が明らかとなった。本項ではこれらの結果を中心に述べる。

拡張型心筋症

わが国の厚生省特定疾患特発性心筋症調査研究班の調査(1974年)では、心筋症の発生頻度(有病率)は人口10万人あたり0.56人であるが1)、心エコー図が普及する以前のデータであり実際の頻度よりかなり低いと考えられている。拡張型心筋症については新潟県における1985年の特定疾患申請書を対象に調べた成績では2)、有病率は人口10万人あたり1.06であるが、1992年では、9.03と増加している3)。心エコー図を用いた住民検診に基づく黒田らの成績4)では、有病率は15となっている。また、鬼平と三浦5)による秋田市での調査では、拡張型の有病率は人口10万人あたり12.5人である。欧米でも8.3~36.5人とほぼ同様な頻度が報告されている6.7)。

筆者らは1998年,わが国における特発性心筋症の疫学像を明らかにする目的で,全国疫学調査を実施した^{8,9)}。調査対象の病院は,全国の一般病院から規模別に無作為抽出し,大学病院と大規模病院はすべて調査した。調査対象診療科は,内科,循環器科,小児科とし,一次調査票を郵送した診療科は2,414であった。一次調査では,1998年1月1日から12月31日の患者数を調査した。一次調査で「患者あり」との返事のあった診療科に対して拡張型,肥大型,拘束型については二次調査を依頼した。一次調査で報告された症例数は拡張型が6,341例,肥大型が7,262例,拘束型

拡張型

肥大型

拘束型

表1 ▶ 報告患者数と全国 推計患者数

1.6 多病,型 8-10 kg 6-1	4.1. 一次調查外門。東京全国推計 19 第1年第18日 - 東京	### /(95%信頼区間)
	和 告 数	
拡張型心筋症	6,341	(16,500~18,800)
肥大型心筋症	## 7,262 ### 21,900 ###	(20,600~23,200)
拘束型心筋症		(250~350)
ミドロッドリア病	280 640 6640	(500~780)
心Fabry病。	150	(95~205)
QT延長症候群	1,000	(690~1,320)
不整脈原性右室心筋症	245 520	(360~680)

が112例であった。また、ミトコンドリア病280例、心Fabry病70例、QT延長症候群389例、不整脈原性右室心筋症245例であった。1998年の全国推計患者数は、拡張型は17,700人、肥大型は21,900人、拘束型は300人、ミトコンドリア病640人、心Fabry病150人、QT延長症候群1,000人、不整脈原性右室心筋症520人と推計された(表1)。今回の調査では拡張型心筋症全国推計患者数は、人口10万人あたりは14.0人であったが、この調査は病院受診者を対象としており、初期の拡張型心筋症は無症状の者も多いため、実際の有病率はさらに高いものと考えられる。

◇年齢・性別分布

ミネアポリス,セントポール市での結果では1979年では人口10万人あたり男性 63人,女性35人であったが、その後急速に増加し1984年では男性125人、女性58人とかなり高い頻度が報告されている 10)。古賀らは19歳以下には少なく、 $40\sim50$ 歳代に多く、40歳未満では明らかな性差はないが、40歳以降は男性が多いと報告した 11 0。拡張型心筋症が中・高齢の男性に多い傾向は、鬼平 6^{5} やLilienfeld 6^{10} の報告でも認められている。

筆者らの二次調査の結果,拡張型心筋症は男性が1,400例,女性が532例で男女比は2.6であり,年齢分布は1歳から93歳にわたっていた。60歳台にピークがみられた。図1に性年齢分布を示す。家族内に発症者があるものの割合は5.2%であった(表2)。拡張型心筋症と関連が示唆されているC型肝炎ウイルスの感染は6.7%にみられ,同年齢の正常者の1.7%に比し有意に高かった。死因については,心不全による死亡が49.2%と最も多く,次いで不整脈死が20.2%であった(図2)。

ミトコンドリア病

心Fabry病

QT延長症候群

不整脈原性右室心筋症

C型肝炎ウイルス

不整脈死

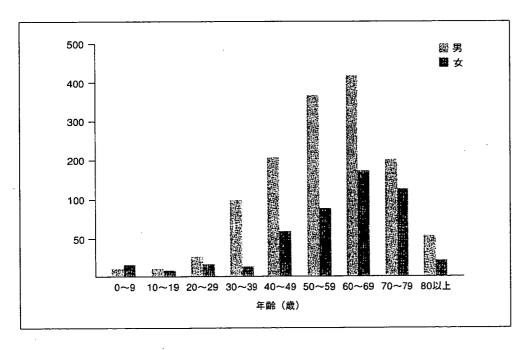


図1 ▶ 拡張型心筋症の性, 年齢別分布

表2 ▶ 家系内発症

等為此一次。並張型心的症候的學家。	症家系内発症。
なし 1,520(78.7) 1,404(-65.8) 15(57.7) 23.1)
の 7 不明 312(16.1) 3430(20:1) 5(19.2)
計: 1,932(100.0)	100.0) 福福

◇予後および予後規定因子

心筋症の予後に関しては昭和57年の厚生省特定疾患特発性心筋症調査研究班の報告¹²⁾では、5年生存率は拡張型54.3%、10年生存率はそれぞれ36.0%とされている。

筆者らは、1999年に実施した全国疫学調査二次調査対象症例の5年後の予後調査を実施した¹³⁾。全国疫学調査二次調査症例1,932例のうち医療機関からの協力が得られた1,554例について5年後の予後を追跡した。原則として医療機関からの予後情報の提供を受けたが、追跡中断例では市町村への住民票請求により生死の確認を行った。追跡を行った1,554例の5年生存率は75.7%であった。また、1998年中に初めて診断された390例での5年生存率は78.6%であった。予後関連要因に関する単変量解析においては、加齢、低い肥満度、NYHA分類、左脚ブロック、心胸比、左室拡張末期径、左室径指数、左室駆出率、診断後経過年数が、予後不良の要因であった。

重要な要因をモデルに含むCox比例ハザードモデルによる多変量解析では、男性、加齢、家族歴、左室径指数、左室駆出率が、互いに独立して有意に高いハザード比を示し、予後不良要因と考えられた。左室径係数の1標準偏差上昇あたりの多変量調整ハザード比は1.33、左室駆出率の1標準偏差低下あたりのハザード比は1.26であった。

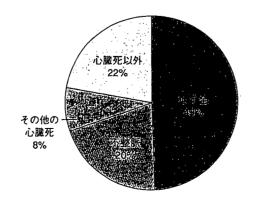
以前のわが国からの報告と比べると、本症の予後はかなり改善してきていると考えられた。また、医療機関による偏りのないわが国での本症患者の予後が明らかになった。予後要因の中では、左室径と左室駆出率がそれぞれ独立して有用な予後予測要因であると考えられた。

肥大型心筋症

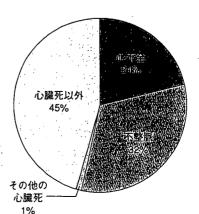
わが国ではHadaら¹⁴⁾が人口10万人に対し170人,黒田ら⁴⁾が374人の頻度を報告している。欧米では^{7, 15, 16)},10万人に対し19.7~1,100人と報告により大きな差がある。このような差の原因として調査方法が異なることが考えられ,一次検診に心電図を用い,異常者のみに心エコー図を施行した場合には,心電図異常のない肥大型心筋症が見落とされる¹¹⁾。

1999年の調査では推計患者数は21,900人で10万人あたり17.3人とこれまでの報告に比べ頻度が低かったが、今回は病院の診療録から調査した報告であり、心症状

図2 ▶ 心筋症の死因



拡張型心筋症



肥大型心筋症

5年生存率

予後関連要因

予後不良要因

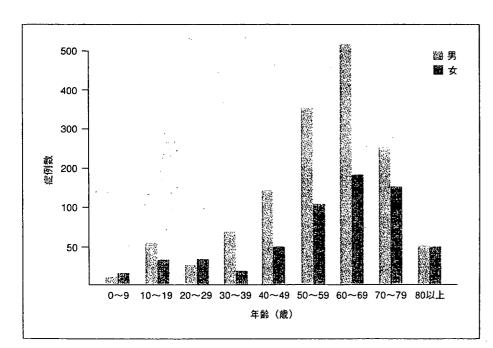
が少ない肥大型心筋症の多くは病院を受診していないと考えられ、過小評価されている可能性が高い。家族内に発症者があるものの割合は14.1%であった(表2)。肥大型では拡張型心筋症とは異なり不整脈死31.9%が心不全死21.3%より多かった(図2)。C型肝炎ウイルス抗体は9.5%に認め、同年齢の正常者の抗体陽性率に比し有意に高く、肥大型では拡張型より高くこれまでの報告と一致している^{17,18)}。古賀の報告では¹¹⁾、好発年齢は、男性では40歳代、50歳代であるが、女性では年齢分布の明らかな差はなく、男女比は10歳代、20歳代ではほぼ1:1で常染色体性優性遺伝様式に一致するが、40歳代以降は3:1以上で、単一遺伝子の異常では説明しえないとしている。

病型別にみると,非閉塞性の非対称性心室中隔肥厚(ASH)や心尖部肥大は40歳以降に好発している。したがって中年以降の非閉塞性のASH例や心尖部肥大例には単一遺伝子の異常ではなく,複数の遺伝子異常が関与する例(多因子病)や後天性因子が関与して発症する例が含まれていると推測している。前述の1998年の調査では,肥大型心筋症は男が1,490例,女が644例で男女比は2.3であり,年齢分布は1歳から100歳にわたり,60歳台にピークがみられた(図3)。

◇予後および予後規定因子

心筋症の予後に関しては昭和57年の厚生省特定疾患特発性心筋症調査研究班の報告¹²⁾では、5年生存率は91.5%、10年生存率は81.8%とされている。古賀¹¹⁾は、10年以上観察しえた59例では初診時にみられた異常Q波は半数で消失し、非対称性肥厚からびまん性肥厚へ進行することを反映した所見と考えた。中隔肥厚の進行や僧帽弁の収縮期前方運動(systolic anterior motion of mitral valve:SAM)の出現・増強を示す例は少数で、一般的な変化は新たなQ波、ブロック、心室内伝導障害の出現、SAMの減弱・消失、心室中隔肥厚の減少で、これらの所見は、HCMの心筋病変は進行性でやがて収縮力低下をきたすことを示すことが報告されている。

1998年の調査1,605例の5年生存率は86.4%で、1998年中に初めて診断された342



心不全死

常染色体性優性遺伝

図3 ▶ 肥大型心筋症の性, 年齢 分布

例での5年生存率は85.7%であった¹³⁾。予後関連要因に関する単変量解析においては、加齢、低肥満度、糖尿病の既往、NYHA、心房細動、左脚ブロック、心胸比、左室駆出率、病床数が少ない病院が、予後不良の要因であった。心尖部肥大の存在は良好な予後を示した。

重要な要因をモデルに含むCox比例ハザードモデルによる多変量解析では、加齢、NYHA、左脚ブロック、心尖部肥大の欠如、心胸比、左脚駆出率が、互いに独立して有意に高いハザード比を示し、予後不良要因と考えられた。特に左室駆出率の低下と心胸比の増大との関連が最も強く、左室駆出率の1標準偏差低下あたりの多変量調整ハザード比は1.42で、心胸比の1標準偏差上昇あたりのハザード比は1.61であった。

【文献】

- 1)服部 譲,中野為夫,服部 諄,ほか:特発性心筋症の疫学調査(第4報).厚生省 特定疾患特発性心筋症調査研究昭和51年度研究報告集,1977,p10-15.
- 2)三井田孝, 横山明裕, 和泉 徹, ほか:新潟県における拡張型心筋症の実体調査. 新潟県医師会報 441:1,1986.
- 3)和泉 徹,加藤公則:心筋症の疫学.本邦での有病率,発症率,自然歴.『循環器 NOW 6.心筋症・心筋炎』(和泉 徹編),南江堂,東京,1994,p2-9.
- 4)黒田敏男, ほか:循環器系住民検診における断層心エコー図法の意義. J Cardiol 19:933-943. 1989.
- 5)鬼平 聡, 三浦 傅:心筋症の疫学. 日本臨床 58:141-146, 2000.
- 6) Williams DG, Olsen EG: Prevalence of overt dilated cardiomyopathy in two regions of England. Br Heart J 54: 153-155, 1985.
- 7) Codd MB, Sugrue DD, Gersh BJ, Melton LJ 3rd: Epidemiology of idiopathic dilated and hypertrophic cardiomyopathy. A population-based study in Olmsted County, Minnesota, 1975-1984. Circulation 80: 564-572, 1989.
- 8) Miura K, Nakagawa H, Morikawa Y, et al: Epidemiology of idiopathic cardiomyopathy in Japan: results from a nationwide survey. Heart 87: 126-130, 2002.
- 9) Matsumori A, Furukawa Y, Hasegawa K, et al: Epidemiologic and clinical characteristics of cardiomyopathies in Japan: results from nationwide surveys. Circ J 66: 323-336, 2002.
- 10) Lilienfeld DE, Sprafka JM, Pham DL, Baxter J: Morbidity from congestive and hypertrophic cardiomyopathy in the Minneapolis St. Paul metropolitan area: 1979-1984. Cardiology 80: 71-76, 1992.
- 11) 古賀義則:心筋症の疫学. 【目でみる循環器病シリーズ14:心筋症】(松森 昭編集), メジカルビュー社, 東京, 2000, p12-17.
- 12)河合忠一, 桜井恒太郎, 岸本千晴, 富岡宣良:特発性心筋症の予後調査. 厚生省特定疾患特発性心筋症調査研究班昭和57年度研究報告集, 1983, p63-66.
- 13) 中川秀昭, 三浦克之, アリ・ナセルモアッデリ, ほか:わが国の拡張型心筋症の予後と予後要因―全国疫学調査5年後の予後調査より―. 厚生労働科学研究費補助金 (難治性疾患克服研究事業)分担研究報告書. (印刷中)
- 14) Hada Y, Sakamoto T, Amano K, et al: Prevalence of hypertrophic cardiomyopathy in a population of adult Japanese workers as detected by echocardiographic screening. Am J Cardiol 59: 183-184, 1987.
- 15) Agnarsson UT, Hardarson T, Hallgrimsson J, Sigfusson N: The prevalence of hypertrophic cardiomyopathy in men: an echocardiographic study with a review of death records. J Intern Med 232: 499-506, 1992.
- 16) Maron BJ, Gardin JM, Flack JM, et al: Prevalance of hypertrophic cardiomyopathy in a general population of young adults. Echocardiographic analysis of 4111 subjects in the CARDIA study. Coronary Artery Risk Development in (Young) Adults. Circulation 92: 785-789, 1995.
- 17) Matsumori A, Ohashi N, Hasegawa K, et al: Hepatitis C virus infection and heart diseases. A multicenter study in Japan. Jpn Circ J 62: 389-391, 1998.
- 18) Matsumori A: Hepatitis C virus infection and cardiomyopathies. Circ Res 96: 144-147, 2005.

Occupational and Environmental Factors and Idiopathic Pulmonary Fibrosis in Japan

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Idiopathic pulmonary fibrosis (IPF) is a progressive fibrosing interstitial lung disease of unknown etiology. Environmental factors, especially occupational agents, may be of great importance in the manifestation of IPF. We examined the relationship between occupational and environmental factors and IPF in Japan. A multicenter hospital-based case-control study was performed in 2001. Included were 102 cases aged 40 years or over who were within 2 years of having been diagnosed in accordance with the most recent criteria. Controls, aged 40 years or over, were 55 hospitalized patients diagnosed as having acute bacterial pneumonia and four outpatients with common colds. Data on occupational and environmental factors were obtained from a questionnaire. Multiple logistic regression analysis was used to estimate the adjusted odds ratios (ORs) and 95% confidence intervals (CIs) of IPF for single factors with adjustment for age, sex and region. Compared with controls, cases were more likely to have been managers, officials or production workers and less likely to have been protective service or materials handling workers. Clerical and related work was significantly related to a decreased risk of IPF after further adjustment for pack-years of smoking (OR = 0.42; 95% CI = 0.18-0.95). Exposure to metal dust was significantly associated with an increased risk of IPF (OR = 9.55; 95% CI = 1.68-181.12). From 20.0 to 39.9 pack-years of smoking was significantly associated with an increased risk of IPF (OR = 3.23; 95% CI = 1.01-10.84). Our results appear to confirm data from previous epidemiologic studies. Metal dust exposure may be a particularly important risk factor for IPF.

Keywords: case-control studies; metal dust; occupations; pulmonary fibrosis; smoking

INTRODUCTION

Idiopathic pulmonary fibrosis (IPF) is a progressive fibrosing interstitial lung disease of unknown etiology (Selman *et al.*, 2001). The mortality rate appears

to be increasing in Western populations (Johnston et al., 1990; Hubbard et al., 1996a; Mannino et al., 1996). Men are more likely than women to develop or die from IPF (Johnston et al., 1990; Coultas et al., 1994; Iwai et al., 1994; Hubbard et al., 1996a; Mannino et al., 1996). A study in the UK found increased deaths due to IPF in traditionally industrialized areas (Johnston et al., 1990). Thus,

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environmental factors, especially occupational agents, may be of great importance in the manifestation of IPF.

Four case-control studies demonstrated that exposure to metal dust was associated with an increased risk of IPF (Scott et al., 1990; Iwai et al., 1994; Hubbard et al., 1996b; Baumgartner et al., 2000). A historical cohort study in the workforce of a major UK engineering company found a doseresponse relationship between years of working with metal and risk of IPF and a 21-fold increase in the odds ratio (OR) among sheet-metal workers (Hubbard et al., 2000). Other occupational agents and job activities associated with IPF have been identified, including wood dust (Hubbard et al., 1996b), textile dust (Hubbard et al., 1996b), sand or stone (Hubbard et al., 1996b; Baumgartner et al., 2000), silica (Mullen et al., 1998), mould in the workplace (Mullen et al., 1998), agricultural chemicals (Iwai et al., 1994), cattle or livestock (Scott et al., 1990; Baumgartner et al., 2000), vegetable/animal dust (Baumgartner et al., 2000), farming (Baumgartner et al., 2000), raising birds (Baumgartner et al., 2000) and hairdressing (Baumgartner et al., 2000). A study of death certificates in England and Wales showed that standardized mortality ratios were elevated among members of the armed forces, miners and quarrymen, service, sports and recreation workers, and electrical and electronic workers, but found no evidence of an increased risk among persons in occupations that potentially exposed them to wood and metal dust (Harris et al., 2001). Cigarette smoking was related to an increased risk of IPF, although there was no clear exposure-response pattern with cumulative consumption of cigarettes in two casecontrol studies (Hubbard et al., 1996b; Baumgartner et al., 1997). Because epidemiologic information regarding the etiologic factors associated with IPF is sparse in Japan, the present study examined the relationship between occupational and environmental factors and the development of IPF, based on a multicenter hospital-based case-control study.

MATERIALS AND METHODS

Subjects

Eligible cases aged 40 years or over who were within 2 years of having been diagnosed with IPF were identified among 21 collaborating hospitals and their 29 affiliated hospitals during the period from 1 June to 30 November 2001. The diagnosis of IPF by the collaborating respiratory disease specialists was based on clinical history, clinical examination and high-resolution computerized tomography (HRCT) of the chest. Results of video-assisted thoracoscopic lung biopsy transbronchial lung biopsy and/or bronchoalveolar lavage, corresponding to the

international consensus statement on IPF of the American Thoracic Society and the European Respiratory Society (American Thoracic Society, 2002), were also used when available, either alone or in combination, to assist diagnosis. All cases had basal fine crackles through auscultation and predominantly peripheral, subpleural, bibasal fine reticular shadows and/or honeycombing, occasionally with traction bronchiectasis and bronchiolectasis on HRCT. There was no evidence of either coexisting collagen-vascular disease or history of known occupational exposure to agents that might produce a clinical picture similar to that of IPF in any of the cases. The physicians in charge asked eligible patients to participate in this study, and 104 patients were cooperative in answering the questionnaires while three patients refused.

Control subjects, aged 40 years or over and without prior respiratory diseases, were prospectively selected from individuals who received treatment at the respiratory ward of the 21 collaborating hospitals and their 29 affiliated hospitals during the same time period as the cases. Potential control subjects consisted of 56 hospitalized patients diagnosed as having acute bacterial pneumonia and four outpatients with common colds. Only one eligible control subject who was asked to take part in this study by a physician refused to answer the questionnaire. Controls were not, individually or in larger groups, matched to cases. Few patients with acute infectious or common diseases receive treatment at a specialized medical institution. Of the 21 collaborating hospitals, 14 were university hospitals with doctors who exclusively treated patients with serious illnesses. Thus, 95 of the 104 cases were recruited from the 21 collaborating hospitals and 34 of the 60 controls were selected from 29 hospitals that were affiliated to the collaborating hospitals. All study subjects gave their fully informed consent in writing.

The study subjects were originally restricted to males, but included in the analysis were 10 female cases and five female controls whose treatment was provided at six of the collaborating hospitals and one affiliated hospital. Incomplete data in relation to cigarette smoking caused the exclusion of two male cases and one male control. There were 102 cases and 59 control subjects left for analysis.

Questionnaires

Sets of two self-administered questionnaires were handed to cases and controls by physicians. The subjects filled out the questionnaires and mailed them to the data management center. A telephone interview was conducted by a trained research technician to complete missing or illogical data.

One of the self-administered questionnaires elicited information on age, sex, type of job held for the longest period of time, exposure to 13 specific

occupational agents, smoking habits, moulds in the house, indoor domestic pets and residential municipality. Employment data focused on type of job held for the longest period of time during the subject's work life and years of exposure were requested regarding the job and occupational agents, respectively. Occupational agents were defined as present if the subject reported ≥10 h of exposure per week. Neither the questionnaire nor a telephone interview requested a full occupational history or gave any information to help responders recall possible exposures to occupational agents that they may otherwise have overlooked in relation to their particular occupation.

The other self-administered questionnaire was a validated self-administered dietary history questionnaire that was used to assess dietary habits over a period of 1 month (Sasaki et al., 1998, 2000). In the present study, data obtained from the dietary history questionnaire were not used.

Statistical analysis

Jobs held for the longest period of time were coded using the Japanese Standard Occupational Classification and stratified into 11 major groups (professional and technical; managers and officials; clerical and related fields; sales; service; protective service; farming, fishing and forestry; transport and communication; production; materials handling; and construction and extraction). Included in this analysis were eight specific occupational agents to which three or more subjects had been exposed for more than a year. Age was classified into four categories (<50, 50–59, 60-69 and 70+ years); region into five (Kanto-Koshinetsu, Tokai, Kinki, Chugoku-Shikoku and Kyushu); cigarette smoking into three (never smoked, former smoker and current smoker); pack-years of smoking into five (none, 0–19.9, 20.0–39.9, 40.0-59.9, and 60.0+); and residential municipality into two (city and town or village). Multiple logistic regression analysis was used to estimate the adjusted ORs and 95% confidence intervals (CIs) of IPF for single factors with adjustment for age, sex and region. The reference category for all occupational factors, moulds in the house and indoor domestic pets was based on the comparison of those exposed to a single agent with all those unexposed, including potential subjects who were exposed to other etiologic factors. All computations were performed using version 8.2 of the SAS software package (SAS Institute, Inc., Cary, NC, USA).

RESULTS

Dyspnea on exertion was present at enrollment in 83 of the 104 cases (81.4%). The median (90% central range) of arterial O_2 pressure was 80.2 mmHg

Table 1. Characteristics of the study population

Variable	n (%)				
	Cases $(n = 102)$	Controls $(n = 59)$			
Sex (male)	92 (90.2)	54 (91.5)			
Age (years)					
<50	3 (2.9)	2 (3.4)			
50–59	15 (14.7)	19 (32.2)			
60–69	56 (54.9)	24 (40.7)			
70+	28 (27.5)	14 (23.7)			
Region					
Kanto-Koshinetsu	56 (54.9)	27 (45.8)			
Tokai	11 (10.8)	10 (17.0)			
Kinki	14 (13.7)	5 (8.5)			
Chugoku-Shikoku	4 (3.9)	6 (10.2)			
Kyushu	17 (16.7)	11 (18.6)			

(57.2–97.0) and that of vital capacity expressed as percentage predicted values was 77.8% (41.0–116.3) in cases. The proportions of male subjects among cases and controls were 90.2% and 91.5%, respectively (Table 1). Compared with control subjects, cases were older and had a lower prevalence of residence in Chugoku-Shikoku.

Table 2 presents adjusted ORs and 95% CIs for IPF in relation to occupational factors after controlling for age, sex and region. Because five female cases had never held a job, they were regarded as 'unexposed' in the occupational analyses. The median duration of the job held for the longest period of time was 35 years in cases and 31 years in controls. No marked difference was found in the risk of IPF among occupational groups (P = 0.50, Wald $\chi^2 = 9.38$ with 10 degrees of freedom for homogeneity of OR for each occupational group). None of the occupational groups was related to the risk of IPF with statistical significance, although at least a 2-fold increase in OR was observed among managers and officials and production workers, and there was a <0.5-fold decrease in OR among those in clerical and related fields, protective service and materials handling. Further adjustment for packyears of smoking slightly strengthened associations with two major occupational groups: managers and officials, and clerical and related occupations (adjusted ORs were 6.06, 95% CI: 0.97-118.6 and 0.42, 95% CI: 0.18-0.95, respectively). Overall, 25 cases and three controls were exposed to only one of the occupational agents being studied, and four cases and two controls were exposed to two agents. Only three and one cases were exposed to three and six occupational agents, respectively. Exposure to any of the eight kinds of dust being studied was significantly associated with an increased risk of IPF. In particular, exposure to metal dust was related to an ~10-fold increased risk of IPF. No association 262

Table 2. Adjusted odds ratios for idiopathic pulmonary fibrosis in relation to occupational factors

Factor	n (%)		Adjusted odds ratio ^a	95% confidence	
	Cases $(n = 102)$	Controls $(n = 59)$	ouds fatio	ilici vai	
Job held for the longest period of time	me				
Professional or technical	9 (8.8)	7 (11.9)	0.71	0.23-2.25	
Manager or official	9 (8.8)	1 (1.7)	4.26	0.74-80.88	
Clerical or related occupation	18 (17.7)	18 (30.5)	0.49	0.22-1.08	
Sales	11 (10.8)	6 (10.2)	1.29	0.44-4.18	
Service	6 (5.9)	3 (5.1)	1.02	0.23-5.46	
Protective service	2 (2.0)	3 (5.1)	0.33	0.04-2.19	
Farming, fishing or forestry	7 (6.9)	7 (11.9)	0.55	0.16-1.89	
Transport or communication	4 (3.9)	2 (3.4)	1.10	0.19-8.73	
Production	18 (17.7)	5 (8.5)	2.56	0.91-8.54	
Materials handling	2 (2.0)	2 (3.4)	0.46	0.05-4.34	
Construction or extraction	11 (10.8)	5 (8.5)	1.37	0.42-4.44	
Occupational agents					
Any dust ^b	33 (32.4)	5 (8.5)	5.61	2.12-17.89	
Metal	12 (11.8)	1 (1.7)	9.55	1.68-181.12	
Wood	5 (4.9)	0 (0.0)			
Asbestos	3 (2.9)	0 (0.0)			
Coal	3 (2.9)	0 (0.0)			
Stone or sand	11 (10.8)	4 (6.8)	1.75	0.52-7.01	
Solvents	4 (3.9)	0 (0.0)			
Pesticides	6 (5.9)	2 (3.4)	1.46	0.30-10.61	
Chalk	4 (3.9)	0 (0.0)			

^aAdjusted for age (<50, 50-59, 60-69 or 70+ years), sex and region (Kanto-Koshinetsu, Tokai, Kinki, Chugoku-Shikoku and Kyushu).

of exposure to stone, sand or pesticides with the risk of IPF was found. None of the control subjects reported exposure to wood, asbestos, coal, solvents or chalk. Additional adjustment for pack-years of smoking did not change the association with IPF of exposure to any of the dusts being studied and metal dust. When exposures to metal dust, stone, sand and pesticides were included in the same model with age, sex and region, a positive association between metal dust exposure and IPF was slightly attenuated but remained statistically significant (adjusted OR 9.25, 95% CI: 1.59–176.7).

Results for environmental factors are shown in Table 3. More cases than control subjects were former smokers, whereas current smoking was more prevalent in controls than in cases, although differences between groups were not statistically significant. Adjusted OR for the comparison of having smoked with never having smoked was 1.91 (95% CI: 0.71–5.15). A significantly increased risk of IPF was observed for smokers with 20.0–39.9 pack-years, but there was no dose–response association with cumulative consumption of cigarettes. Although not statistically significant, moulds in the living room and the presence of indoor hamsters were

associated with a >50% decreased risk of IPF. Moulds in the bathroom, kitchen or closets and the presence of indoor birds, cats or dogs were not measurably related to the risk of IPF. There was no clear difference between cases and controls in terms of residential municipality.

DISCUSSION

The present study demonstrated that, compared with control subjects, cases were more likely to have been managers and officials or production workers and less likely to have been protective service workers or materials handling workers, although none of the effects reached significance. Workers in clerical and related fields had a significantly decreased risk of IPF independent of age, sex, region and smoking status. Exposure to metal dust was significantly associated with an increased risk of IPF, but exposure to stone, sand or pesticides was not materially related to IPF. There was no statistically significant relationship between the environmental factors under study and IPF, although 20.0-39.9 pack-years of smoking was significantly associated with an increased risk of IPF.

Eight cases and two controls were exposed to two or more occupational agents.

Table 3. Adjusted odds ratios for idiopathic pulmonary fibrosis in relation to environmental factors

Factor	n (%)		Adjusted odds ratio ^a	95% confidence	
	Cases $(n = 102)$	Controls $(n = 59)$		111101 7 111	
Smoking status					
Never smoked	18 (17.6)	14 (23.7)	1.00		
Former smoker	80 (78.4)	34 (57.6)	2.21	0.82-6.04	
Current smoker	4 (3.9)	11 (18.6)	0.50	0.10-2.24	
Pack-years of smoking					
None	18 (17.7)	14 (23.7)	1.00		
0.6-19.9	10 (9.8)	11 (18.6)	0.87	0.25-3.10	
20.0-39.9	30 (29.4)	10 (17.0)	3.23	1.01-10.84	
40.0-59.9	29 (28.4)	15 (25.4)	2.22	0.70-7.23	
60.0+	15 (14.7)	9 (15.3)	1.59	0.46-5.64	
Moulds					
Any place ^b	56 (54.9)	36 (61.0)	0.98	0.48-2.01	
Living room	5 (4.9)	8 (13.6)	0.36	0.10-1.20	
Bathroom	51 (50.0)	28 (47.5)	1.38	0.69-2.82	
Kitchen	12 (11.8)	11 (18.6)	0.61	0.24-1.57	
Closets	17 (16.7)	9 (15.3)	1.25	0.50-3.30	
Indoor domestic pets					
Any pets ^c	40 (39.2)	25 (42.4)	0.94	0.47-1.86	
Birds	17 (16.7)	9 (15.3)	1.16	0.47-3.03	
Cats	14 (13.7)	8 (13.6)	1.24	0.45-3.58	
Dogs	15 (14.7)	10 (17.0)	0.85	0.33-2.26	
Hamsters	2 (2.0)	3 (5.1)	0.27	0.03-1.80	
Residential municipality	•		•		
Village or town	15 (14.7)	12 (20.3)	1.00		
City	87 (85.3)	47 (79.7)	1.35	0.56-3.28	

^aAdjusted for age (<50, 50-59, 60-69 or 70+ years), sex and region (Kanto-Koshinetsu, Tokai, Kinki, Chugoku-Shikoku and Kyushu).

These findings are in agreement with previous observations showing a positive relationship between exposure to metal dust and the risk of IPF (Scott et al., 1990; Iwai et al., 1994; Hubbard et al., 1996b; Baumgartner et al., 2000), but they are at variance with a case-control study that reported positive associations between farming and stone/sand dust exposure and IPF (Baumgartner et al., 2000). The mechanisms underlying the positive association between metal dust exposure and the risk of IPF are still obscure. Recent experimental research has demonstrated that particulate nickel promotes pulmonary fibrosis by inhibiting the fibrinolytic cascade (Andrew and Barchowsky, 2000). Potolicchio et al. reported that susceptibility to hard metal lung disease is associated with binding of cobalt by HLA-DP molecules (Potolicchio et al., 1997, 1999). Managers, officials and those working in clerical and related fields are not likely to be potentially exposed to metal dust. The first Whitehall study showed that mortality rates from lung cancer, chronic bronchitis and respiratory diseases were markedly increased with a decrease in employment grade (van Rossum et al., 2000). The present findings partially contradict this observation. In our current study, nine cases were managers and officials: six presidents of a company, two department managers of a company and one director of a union. One of the controls was an executive director of a company. None had been exposed to any of the occupational agents under investigation. A non-significant increased risk of IPF among managers and officials may be ascribed to unrecognized factors that are related to job grade. A case-control study in Lithuania found that the main risk factors of myocardial infarction for managers were hypertension and stress (Malinauskiene et al., 2002). Their stressful work life may have contributed to the manifestation of IPF. The excess IPF risk among managers and officials was not likely to be explained by overdiagnosis in the higher employment grades. All Japanese are covered by universal medical care insurance and all are provided with completely free access to the same medical care.

^bOverall, 22 cases and 13 controls were exposed to moulds in two or more places.

^cOverall, 8 cases and 3 controls had two or more types of indoor domestic pets.

A case-control study in the USA reported that being a former smoker and 21-40 pack-years of smoking were significantly related to an increased risk of IPF, whereas there was no association of current smoking and more than 40 pack-years of smoking with IPF (Baumgartner et al., 1997). Our results are generally in agreement with these findings, although a positive relationship between having previously smoked and IPF was not statistically significant in this study. A history of having ever smoked was associated with a 1.6-fold increased risk of IPF in the above-cited US study (Baumgartner et al., 1997) and a study in the UK (Hubbard et al., 1996b). The present findings, although not statistically significant, are similar to these observations. Overall, 34 cases and 14 controls quit smoking within 3 years of data collection. It is possible that cases were more likely to quit smoking because of diagnosis or the progression of their disease. When these subjects who stopped smoking within 3 years were considered to be current smokers, adjusted ORs (95% CIs) were 2.11 (0.72-6.19) and 1.73 (0.60-5.03) for former and current smoking, respectively. Additional adjustment for packyears of smoking slightly affected the association with IPF in managers and officials and workers in clerical and related fields, although the interaction with pack-years of smoking was not statistically significant for those occupational groups. On the other hand, adjustment for smoking did not measurably influence the effects of metal dust exposure. Thus, smoking and metal dust exposure were likely to be independent factors. A case-control study of 17 cases and 94 controls in the USA reported that patients with interstitial lung disease were 16.0 times more likely to be exposed to mould than were controls in their workplace (Mullen et al., 1998). The present results are not consistent with this finding. To our knowledge, no study has assessed the relationship between domestic pets and the risk of IPF. Our findings contradict a previous epidemiologic study in Japan showing a positive association between residence in an agricultural area and the risk of IPF (Iwai et al., 1994).

Selection and information bias are methodological issues that need careful consideration. We attempted to identify and recruit all eligible cases seen at each participating hospital during the specified study period according to the most recent diagnostic criteria. Only three eligible patients did not take part in this study. Thus, it was unlikely that selection bias for the cases occurred. It is difficult to ensure that control subjects are drawn from the same study population as the cases. This disadvantage is likely to be diluted by controlling for region. Almost all controls were hospitalized patients with acute bacterial pneumonia. Therefore, control subjects may not have been representative of the general population that generated the

cases. The prevalence values of having ever smoked in the present controls were not likely to differ from those reported elsewhere, although the prevalence values of current smoking were relatively low in this study. In a population-based case-control study of acute myocardial infarction in Fukuoka, Japan, the proportions of people who had never smoked, had formerly smoked and currently smoked were 25%, 23% and 52%, respectively, among 260 male controls below 65 years old, and 24%, 41% and 35%, respectively, among 212 male controls aged 65 years or over (Miyake and Fukuoka Heart Study Group, 2000). The corresponding figures in this study were 14%, 55% and 31%, respectively, among 29 male controls below 65 years old, and 24%, 68% and 8%, respectively, among 25 male controls aged 65 years or over. If acute bacterial pneumonia shared risk factors with IPF, the reported OR would have been underestimated. The ratio of controls to cases was below 1:1. Eligible control subjects with acute bacterial pneumonia were not likely to arise during the summer months because of seasonal variation in this disease. Moreover, eligible control patients who received treatment at the non-respiratory ward of 50 hospitals were not recruited. The statistical power of this study was extremely low, although a statistically significant association was observed. Cases may have been more likely than controls to remember specific exposures under study. However, subjects would not have been aware of the possible ill effects of occupational and environmental factors under investigation because the etiology of IPF is unknown. Thus, a difference in recall between cases and controls was not likely to have occurred. We did not collect data for a detailed occupational history. However, the impact of job activities, other than the job held for the longest period of time, on IPF was likely to be negligible and unlikely to differ between cases and controls because the median duration of the job held for the longest period of time was 30 years or more in both cases and controls. The consequence could be a minor underestimation of values in our results.

Despite these potential limitations, the present results appear to confirm data from previous epidemiologic studies. Exposure to metal dust is a particularly important risk factor for IPF in Japan, as well as in the UK and the USA. Larger studies with more precise and detailed exposure measurements are needed to assess the impact of occupational and environmental factors on the development of IPF. Investigations regarding biological mechanisms are also required.

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REFERENCES

- American Thoracic Society. (2002) American Thoracic Society/European Respiratory Society international multidisciplinary consensus classification of the idiopathic interstitial pneumonias. Am J Respir Crit Care Med; 165: 277–304.
- Andrew A, Barchowsky A. (2000) Nickel-induced plasminogen activator inhibitor-1 expression inhibits the fibrinolytic activity of human airway epithelial cells. Toxicol Appl Pharmacol; 168: 50-7.
- Baumgartner KB, Samet JM, Stidley CA, Colby TV, Waldron JA, Collaborating centers. (1997) Cigarette smoking: a risk factor for idiopathic pulmonary fibrosis. Am J Respir Crit Care Med; 155: 242–8.
- Baumgartner KB, Samet JM, Coultas DB et al. (2000) Occupational and environmental risk factors for idiopathic pulmonary fibrosis: a multicenter case—control study. Am J Epidemiol; 152: 307–15.
- Coultas DB, Zumwalt RE, Black WC, Sobonya RE. (1994) The epidemiology of interstitial lung diseases. Am J Respir Crit Care Med; 150: 967–72.
- Harris JM, Cullinan P, McDonald JC. (2001) Occupational distribution and geographic clustering of deaths certified to be cryptogenic fibrosing alveolitis in England and Wales. Chest; 119: 428–33.
- Hubbard R, Cooper M, Antoniak M et al. (2000) Risk of cryptogenic fibrosing alveolitis in metal workers. Lancet; 355: 466-7.
- Hubbard R, Johnston I, Coultas DB, Britton J. (1996a) Mortality rates from cryptogenic fibrosing alveolitis in seven countries. Thorax; 51: 711–16.
- Hubbard R, Lewis S, Richards K, Johnston I, Britton J. (1996b) Occupational exposure to metal or wood dust and aetiology of cryptogenic fibrosing alveolitis. Lancet; 347: 284–9.
- Iwai K, Mori T, Yamada N, Yamaguchi M, Hosoda Y. (1994) Idiopathic pulmonary fibrosis: epidemiologic approaches to occupational exposure. Am J Respir Crit Care Med; 150: 670-5.
- Johnston I, Britton J, Kinnear W, Logan R. (1990) Rising mortality from cryptogenic fibrosing alveolitis. Brit Med J; 301: 1017-21.

- Malinauskiene V, Grazuleviciene R, Nieuwenhuijsen MJ, Azaraviciene A. (2002) Myocardial infarction risk and occupational categories in Kaunas 25–64 year old men. Occup Environ Med; 59: 745–50.
- Mannino DM, Etzel RA, Parrish RG. (1996) Pulmonary fibrosis deaths in the United States, 1979–1991. An analysis of multiple-cause mortality data. Am J Respir Crit Care Med; 153: 1548–52.
- Miyake Y, Fukuoka Heart Study Group. (2000) Risk factors for non-fatal acute myocardial infarction in middle-aged and older Japanese. Jpn Circ J; 64: 103–9.
- Mullen J, Hodgson MJ, DeGraff CA, Godar T. (1998) Casecontrol study of idiopathic pulmonary fibrosis and environmental exposures. J Occup Environ Med; 40: 363-7.
- Potolicchio I, Festucci A, Hausler P, Sorrentino R. (1999) HLA-DP molecules bind cobalt: a possible explanation for the genetic association with hard metal disease. Euro J Immunol; 29: 2140–47.
- Potolicchio I, Mosconi G, Forni A, Nemery B, Seghizzi P, Sorrentino R. (1997) Susceptibility to hard metal lung disease is strongly associated with the presence of glutamate 69 in HLA-DPβ chain. Euro J Immunol; 27: 2741–43.
- Sasaki S, Ushio F, Amano K et al. (2000) Serum biomarkerbased validation of a self-administered diet history questionnaire for Japanese subjects. J Nutr Sci Vitaminol; 46: 285–96.
- Sasaki S, Yanagibori R, Amano K. (1998) Self-administered diet history questionnaire developed for health education: a relative validation of the test-version by comparison with 3-day diet record in women. J Epidemiol; 8: 203–15.
- Scott J, Johnston I, Britton J. (1990) What causes cryptogenic fibrosing alveolitis? A case-control study of environmental exposure to dust. Brit Med J; 301: 1015-17.
- Selman M, King TE Jr, Pardo A. (2001) Idiopathic pulmonary fibrosis: prevailing and evolving hypotheses about its pathogenesis and implications for therapy. Ann Intern Med; 134: 136-51.
- van Rossum CTM, Shipley MJ, van de Mheen H, Grobbee DE, Marmot MG. (2000) Employment grade differences in cause specific mortality. A 25 year follow up of civil servants from the first Whitehall study. J Epidemiol Community Health; 54: 178–84.