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## ORIGINAL ARTICLE

# Psoriasis affects individuals of African descent and white Brazilians similarly

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### KEYWORDS

Psoriasis;  
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### Abstract

**Introduction:** Cultural, socio-demographic and environmental factors such as tropical climate and exposure to sun could have an impact on the incidence or clinical course of psoriasis. Here we describe the main clinical aspects of psoriasis in Brazilian patients and also investigate whether any particular feature can distinguish the disease occurring in Brazil from that occurring in other countries.

**Material and methods:** We recorded the clinical features of 151 psoriasis patients seen in a Brazilian public dermatological care unit between 2006 and 2008.

**Results:** Males and females were similarly affected. The reported races were as follows: whites, 47 cases (41.6%), interracial individuals (mixed race), 42 cases (37.2%) and blacks, 24 cases (21.2%). Chronic plaque-type psoriasis was the most prevalent clinical form (110 cases, 72.8%) followed by palm and sole involvement (21 cases, 13.9%).

**Conclusions:** We demonstrated that psoriasis in these Brazilian subjects was similar to that observed in subjects from other countries, but interracial and black populations were affected as much as whites. Considering the high rate of interracial populations among Brazilians we cannot exclude the possibility that Afro-descendants may have inherited Caucasian genes associated with psoriasis. Poor socio-economic conditions of Afro-descendants can limit their possibilities of receiving adequate treatments, impairing their health-related quality of life.

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**PALABRAS CLAVE**

Psoriasis;  
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Población brasileña;  
Negros

## La psoriasis afecta de forma similar a los pacientes brasileños de ascendencia africana y caucásica

**Resumen**

**Introducción:** Los factores culturales, sociodemográficos y ambientales tales como el clima tropical o la exposición solar pueden tener un impacto en la incidencia o el curso clínico de la psoriasis. En este artículo describimos los principales aspectos clínicos de la psoriasis en pacientes brasileños e investigamos si existe alguna característica que permita distinguir la enfermedad que ocurre en Brasil de la que se encuentra en otros países.

**Material y métodos:** Se recogieron las características clínicas de 151 pacientes con psoriasis evaluados en un centro dermatológico público de Brasil entre 2006 y 2008.

**Resultados:** Los hombres y las mujeres estaban afectados de forma similar. La frecuencia de afectación según la raza era la siguiente: blancos 47 casos (41,6%), mestizos 42 casos (37,2%) y negros 24 casos (21,2%). Las formas clínicas más prevalentes fueron la psoriasis crónica en placas (110 casos, 72,8%) seguida de la psoriasis palmoplantar (21 casos, 13,9%).

**Conclusiones:** Demostramos que la psoriasis en estos sujetos brasileños es similar a la que se observa en sujetos de otros países, pero los mestizos y los negros están afectados tanto como los blancos. Teniendo en cuenta la elevada proporción de población mestiza entre los brasileños, no podemos descartar la posibilidad de que los descendientes africanos hayan podido heredar los genes caucásicos asociados a la psoriasis. Las pobres condiciones socioeconómicas de los descendientes africanos pueden limitar sus posibilidades para recibir tratamientos adecuados, lo que altera su calidad de vida.

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**Introduction**

Psoriasis is a chronic inflammatory disease that affects skin and joints<sup>1</sup>. Estimates of prevalence are relatively high, varying from 0.5 to 4.6%, depending on the geographic location, but especially affecting Caucasians<sup>1,2</sup>. The etiology of the disease is unknown, but genetic predisposition and environmental factors can influence the occurrence and severity of the disease<sup>1,3,4</sup>. Epidermal proliferation is a key feature of psoriasis although immune mediated mechanisms are crucial for maintaining the inflammatory lesions<sup>5,6</sup>. Plaque psoriasis is the most frequent clinical form of the disease, affecting up to 80% of patients<sup>3,4</sup>. The onset of symptoms can be divided into two periods, teenagers (type 1) or individuals in the fifth decade of life (type 2), without distinct sex prevalence<sup>7</sup>. Most studies concerning epidemiology and clinical aspects of psoriasis focus on European and North American populations. However, very few reports address the epidemiological and clinical features of psoriasis in developing regions such as South America<sup>8,9</sup>. In Brazil, miscegenation, besides the typical tropical climate and sun exposure, can have a beneficial impact such as delayed onset and progression of psoriatic lesions. Here we describe the main clinical aspects of psoriasis in Brazilian patients that were evaluated in a public dermatological outpatient care unit in Rio de Janeiro state. We also investigate whether any particular feature can distinguish the disease occurring among Brazilian patients from that occurring among patients from other countries.

**Methodology**

We carried out a retrospective analysis of selected records of 151 patients that have been seen at the public

dermatological outpatient care unit of the Hospital Gafreé-Guinle/UNIRIO from April 2006 to September 2008. Demographic data such as age, sex, race, and geographic origin were considered. Patients were submitted to dermatological examination, and clinical aspects such as duration of illness before diagnosis and lesion features (clinical form, distribution and shape of lesions) were also investigated. All patients fulfilled the inclusion criteria for the diagnosis of psoriasis, which were based on established clinical parameters<sup>1</sup> and histopathological criteria<sup>6</sup>. The severity of psoriasis was assessed. To measure the activity and severity of psoriasis we used the Physician Global Assessment (PGA). The PGA is one of the most commonly used tools for assessing psoriasis activity and for following clinical response to treatment. There are two main forms of assessment completed by a physician: a static and a dynamic one, in which the physician assesses the global improvement compared to baseline; the latter is hardly reproducible and is based on the observer's memory, so the static assessment has been generally made to assess overall psoriasis, using a score between 0 and 6 (0=clear [no signs of psoriasis], 1=almost clear [minimal], 2=mild [slight plaque elevation, scaling and/or erythema], 3=mild to moderate [intermediate between mild and moderate], 4=moderate [moderate plaque elevation, scaling and/or erythema], 5=moderate to severe [marked plaque elevation, scaling and/or erythema] and 6=severe [very marked plaque elevation, scaling and/or erythema])<sup>10-18</sup>. In this period, the new dermatological outpatients were evaluated to estimate the prevalence of psoriasis in the Department of Dermatology. Prevalence is usually defined as the proportion of individuals in a given population with a disease in a specified time period.

This study was approved by the Ethical Committee of the Escola de Medicina e Cirurgia (UNIRIO, MEC, Brazil).

## Results

The new dermatological outpatients in the general clinic were analyzed for skin color and were reported as follows: interracial individuals (mixed race): 61.2%, whites: 37%, blacks: 1.9%, and 78.2% of the patients were primarily from Rio de Janeiro state. Psoriasis was responsible for 5.2% (95% confidence interval: 3.1–7.6%) of consultations at the Department of Dermatology in this period of 17 months. Among the 151 psoriatic patients selected for the study, the disease was equally distributed between males (71 cases, 46.3%) and females (80 cases, 53.6%). The mean age  $\pm$  standard deviation was  $42.1 \pm 20.9$  years (median 44 years, ranging from 3 to 83 years) for males and  $35.8 \pm 20.8$  years (median 36 years; ranging from 3 to 80 years) for females. Patients aged between 51–60 years (19.9%), 41–50 years (17.6%) and 31–40% (14%) accounted for the majority of cases, followed by patients aged between 0–10 years (16.2%). The period in which women reached medical care for the first consultation after the onset of symptoms was more precocious ( $3.6 \pm 5.4$  years; median 1 year, ranging from one month to 25 years) than for men ( $4.7 \pm 6.9$  years; median 2 years, ranging from two months to 33 years). One hundred thirteen cases were analyzed for skin color and were reported as follows: whites, 47 cases (41.6%), interracial individuals (mixed race), 42 cases (37.2%) and blacks, 24 (21.2%). No significant difference in terms of race was observed among males and females (Table 1). The most frequent clinical forms of psoriasis are recorded (Table 1), but chronic plaque psoriasis was the most prevalent (110 cases, 72.8%). Indeed, palm and sole involvement accounted for 13.9% of cases. The mean age of chronic plaque patients at the onset of symptoms was  $39.8 \pm 20.6$  years, with no sex differences. The majority of patients were also interracial individuals or blacks (66 cases). Patients usually exhibited single lesions (46 in 133 cases), but two (30 in 133 cases) or three lesions (20 in 133 cases) were equally very common. The affected body regions were mainly the scalp (28 cases, 21%), elbows (25 cases, 18.8%), and knees (21 cases, 15.8%). Other locations such as nails, face, back, abdomen, and chest were referred in less than 1% of cases. Only two cases of psoriatic arthritis, and no cases of inverse psoriasis or generalized pustular psoriasis were found. One hundred twenty two cases were analyzed for the severity of psoriasis

(PGA score) and are reported as follows: 0 (none of the patients), 1 (9 in 122 cases), 2 (32 in 122 cases), 3 (33 in 122 cases), 4 (21 in 122 cases), 5 (13 in 122 cases) and 6 (14 in 122 cases). The reported associated diseases were hypertension (9), atopy (7), HIV/AIDS (2), hepatitis (2), diabetes mellitus (3), and hyperthyroidism (1). Only five patients referred a family history of psoriasis.

## Discussion

Psoriasis affects people worldwide<sup>1–3</sup>; however, cultural, socio-demographic, and environmental factors could have an impact on the incidence or clinical course of the disease in some geographic regions. Here we analyzed psoriasis cases from an outpatient care unit located in Rio de Janeiro, Brazil. We demonstrated that the clinical form of the disease was similar to that observed in other countries, but interracial and black populations were affected as much as whites. No difference regarding gender was observed in the present study, even after taking into account different races. Considering the mean age of the onset of symptoms, our patients were equally distributed into type I and type II<sup>7</sup>. As expected, chronic plaque psoriasis was the most common clinical presentation with no difference for gender or skin color. Conversely, psoriatic arthritis was rarely reported, probably because these patients were mainly seen by Rheumatology services. Interracial and black subjects did not present a more severe clinical form of the disease, as has been suggested by previous authors<sup>19</sup>. The low intensity and severity of psoriasis in this sample (PGA score 5 [13 in 122 cases] and 6 [14 in 122 cases]) could be attributed to the tropical climate in Brazil, a country located close to the equator. The country is exposed to high levels of ultraviolet radiation from intense sunlight throughout the year, which leads to a much better prognosis of the disease<sup>20,21</sup>. In a Colombian study, 18.6% of 86 patients presented severe psoriasis<sup>22</sup>.

In our study, blacks comprised 18% of patients seen in our care unit, and our results indicate that psoriasis also affects an Afro-descendant population. Such percentage is high considering the official data showing that the Brazilian population consists of 38.2% of interracial individuals and 5.9% of blacks<sup>23</sup>, which include Afro-descendants. These

**Table 1** Main patients' demographic characteristics related to clinical forms of psoriasis observed in a Brazilian population

Clinical forms of psoriasis	Frequency of cases	Age (years) <sup>a</sup>	Sex (M/F)	Period of illness (years) <sup>a</sup>	Race <sup>b</sup>		
					White	Interracial	Black
Chronic plaque	110 (72.8%)	$39.8 \pm 20.6$	55/55	$4.2 \pm 6.0$	34	30	14
Palm and sole	21 (13.9%)	$24.7 \pm 20.4$	6/15	$2.1 \pm 3.0$	7	7	4
Erythrodermic	8 (5.3%)	$61.0 \pm 13.1$	5/3	$4.7 \pm 8.6$	2	2	2
Pustular	7 (4.6%)	$42.3 \pm 9.1$	2/5	$4.8 \pm 7.1$	1	2	4
Psoriatic arthritis	3 (2.0%)	$56.3 \pm 10.2$	2/1	$12.3 \pm 11.2$	1	1	0
Guttata	2 (1.3%)	$28.5 \pm 3.5$	1/1	$2.2 \pm 2.6$	2	0	0

F: female. M: male.

<sup>a</sup>Results are expressed as mean  $\pm$  standard deviation.

<sup>b</sup>Data were not available for 38 cases.

data are surprising as psoriasis is classically considered a Caucasian disease that affects less than 0.1% of Asians and is considered rare among Africans<sup>1-3,24</sup>. Note that two Latin Americans studies led to different findings. Gonzalez et al analyzed 86 patients and reported that race distribution was as follows: interracial individuals (85%), whites (14%) and blacks (1%)<sup>22</sup>. Trujillo et al analyzed 200 patients and reported the following distribution: interracial individuals (10.5%), whites (85.5%), blacks (3.5%) and unknown (0.5%)<sup>20</sup>. Yet, population-based studies showed that psoriasis affects a significant fraction of African-Americans, although it was observed a reduction in the prevalence of approximately 52% compared to Caucasians<sup>19</sup>.

In contrast, whites predominated in a previous Brazilian study, though it also included patients seen in a private dermatology practice<sup>9</sup>. In this study, we hypothesize that blacks may have inherited Caucasian genes associated with psoriasis<sup>25</sup> because of the high number of interracial populations in Brazil. In fact, this hypothesis is consistent with the literature. Green reported that psoriasis among indigenous Australians of “full-blood” descent may be rare or non-existent; minimal cases have been reported among Aborigines, and psoriasis diagnoses were associated with Aboriginal people of mixed descent<sup>26</sup>. Further research indicated that the condition appeared to be relatively uncommon among Nigerians and Mongolians, and more common among Kenyans and people from the Faroe Islands<sup>27-29</sup>. It is worth noting that the different methodology employed in research design, such as population-based research compared with hospital-based research, makes it difficult to conclude whether these differences were indeed a result of racial variation<sup>30</sup>.

Indeed, other factors such as behavior and environmental conditions can also influence the development of the disease<sup>24,31</sup>. Perhaps a major component of regional variation in the frequency, severity and morbidity of psoriasis is climate variation. A population-based study reported a seasonal variation in psoriasis diagnoses, in which over 65% of the cases were diagnosed in winter and spring, as opposed to approximately 30% of cases diagnosed in summer and autumn<sup>32</sup>. Farber and Nail reported that almost 90% of the respondents indicated that cold weather made their psoriasis worse, approximately 80% claimed that hot weather made their psoriasis better, and 80% stated that sunlight made their psoriasis better<sup>33</sup>. With respect to geographical or climatic features, an author recorded the highest prevalence level of the disease in the central area of Spain, whose weather is drier and colder than in the northern and southern/Mediterranean regions of Spain<sup>34</sup>.

In country-specific studies, the estimated prevalence of psoriasis ranges from 0% in Australian Aborigines and Andean Indians to 11.8% in the inhabitants of Kazakhstan (an Arctic region of the Soviet Union)<sup>35</sup>. More comprehensive studies reported that the prevalence of psoriasis was 1.4% in Spain<sup>34</sup>, 1.5% in the United Kingdom<sup>36</sup> and 2.9% in South Africa<sup>37</sup> and Italy<sup>38</sup>. The present study analyzed the epidemiology of psoriasis, but the prevalence of psoriasis (5.2%) was estimated for new dermatological outpatients in a limited clinical setting, with a small population. Therefore, it does not reflect the true prevalence of this specific disease in the Brazilian population. We believe that larger population-based studies should provide a broader picture of

the incidence and/or prevalence of psoriasis in Brazil. Trujillo et al reported that psoriasis represents 6% of the dermatologic consultations in Cuba<sup>20</sup>. This large number is justified because the health system in Cuba<sup>20</sup>, as well as in Brazil<sup>39</sup>, is free, allowing easy access to data concerning the entire population.

Although this study was not designed to estimate the race prevalence of psoriasis, our results suggest that this disease is common in the African-descendant Brazilian population. Different ethnic backgrounds can mirror the health-related quality of life, and African-Brazilians are more likely to have a low social status compared to that of Caucasians<sup>40</sup>. Indeed, poor socio-economic factors can limit African-Brazilian patients' possibilities of receiving adequate treatments, and such factors can have an effect on these patients' health-related quality of life. Accordingly, these results should be considered in health care policy making in Brazil, particularly when developing policies in public health programs for psoriasis patients.

## Conflict of interest

Authors have no conflict of interest to declare.

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## References

1. Christophers E. Psoriasis-epidemiology and clinical spectrum. *Clin Exp Dermatol.* 2001;26:314-20.
2. Gudjonsson JE, Elder JT. Psoriasis: epidemiology. *Clin Dermatol.* 2007;25:535-46.
3. Lebwohl M. Psoriasis. *Lancet.* 2003;361:1197-204.
4. Griffiths CEM, Barker JNWN. Pathogenesis and clinical features of psoriasis. *Lancet.* 2007;370:263-71.
5. Sabat R, Philip S, Höflich C, Kreutzer S, Wallace E, Asadullah K, et al. Immunopathogenesis of psoriasis. *Exp Dermatol.* 2007;16:779-98.
6. Lowes MA, Bowcock AM, Krueger JG. Pathogenesis and therapy of psoriasis. *Nature.* 2007;445:866-73.
7. Sampogna F, Gisondi P, Melchi CF, Amerio P, Girolomoni G, Abeni D, et al. Prevalence of symptoms experienced by patients with different clinical types of psoriasis. *Br J Dermatol.* 2004;151:594-9.
8. Aslanian FM, Lisboa FF, Iwamoto A, Carneiro SC. Clinical and epidemiological evaluation of psoriasis: clinical variants and articular manifestations. *J Eur Acad Dermatol Venereol.* 2005;19:141-2.
9. Azulay RD, Salvatti C, Rodrigues PC. Estudo comparativo da psoríase em duas instituições dermatológicas no período de 11 anos. *An Bras Dermatol.* 1975;50:33-47.
10. Feldman SR, Krueger GG. Psoriasis assessment tools in clinical trials. *Ann Rheum Dis.* 2005;64(Suppl. 2):ii65-8.
11. Gottlieb AB, Chaudhari U, Baker DG, Perate M, Dooley LT. The National Psoriasis Foundation Psoriasis Score (NPF-PS) System versus the Psoriasis Area Severity Index (PASI) and Physician's

- Global Assessment (PGA): a comparison. *J Drugs Dermatol.* 2003;2:260–6.
12. García-Diez A, Foraster CF, Sebastián FV, Tudela LL, Llach XB, Fernández GS. What characterizes the severity of psoriasis? Results from an epidemiological study of over 3,300 patients in the Iberian region *Dermatology.* 2008;216:137–51.
  13. Puiga L, Bordas X, Carrascosa JM, Daudén E, Ferrándiz C, Hernanz JM, et al. Documento de consenso sobre la evaluación y el tratamiento de la psoriasis moderada/grave del Grupo Español de Psoriasis de la Academia Española de Dermatología y Venereología. *Actas Dermosifiliogr.* 2009;100:277–86.
  14. Langley RG, Ellis CN. Evaluating psoriasis with Psoriasis Area and Severity index, Psoriasis Global Assessment, and Lattice System Physician's Global Assessment. *J Am Acad Dermatol.* 2004;51:563–9.
  15. Berth-Jones J, Grotzinger K, Rainville C, Pham B, Huang J, Daly S, et al. A study examining inter- and intrarater reliability of three scales for measuring severity of psoriasis: Psoriasis Area and Severity Index, Physician's Global Assessment and Lattice System Physician's Global Assessment. *Br J Dermatol.* 2006;155:707–13.
  16. Ashcroft D, Wan Po AL, Williams HC, Griffiths CE. Clinical measures of disease severity and outcome in psoriasis: a critical appraisal of their quality. *Br J Dermatol.* 1999;141:185–91.
  17. Van der Kerkhof PC. The psoriasis area and severity index and alternative approaches for the assessment of severity: persisting areas of confusion. *Br J Dermatol.* 1997;137:661–2.
  18. Puig L. ¿Es realmente útil el PASI como parámetro de medida de severidad en la psoriasis? [Accessed on July 14, 2009]. Available from: [http://www.aedv.es/grupo\\_psoriasis/archivo/utilidad%20PASI.pdf](http://www.aedv.es/grupo_psoriasis/archivo/utilidad%20PASI.pdf).
  19. Gelfand JM, Stern RS, Nijsten T, Feldman SR, Thomas J, Kist J, et al. The prevalence of psoriasis in African Americans: results from a population-based. *J Am Acad Dermatol.* 2005;52:23–6.
  20. Trujillo IA, García MAD, Gemeil OT, Barbosa FT, Lincheta LF, Pérez Hernández MP. Psoriasis vulgar. Estudio descriptivo de 200 pacientes. *Rev Cubana Med.* 2002;41:12–5.
  21. Ozawa M, Ferenczi K, Kikuchi T, Cardinale I, Austin LM, Coven TR, et al. 312-nanometer ultraviolet B light (narrow-band UVB) induces apoptosis of T cells within psoriatic lesions. *J Exp Med.* 1999;189:711–8.
  22. González C, Castro LA, de la Cruz G, Arenas CM, Beltrán A, Santos AM. Caracterización epidemiológica de la psoriasis en el Hospital Militar Central. *Rev Asoc Col Dermatol.* 2009;17:11–7.
  23. Instituto Brasileiro de Geografia e Estatística, IBGE, Ministério do Planejamento Orçamento e Gestão, Brazil [Accessed on December 21, 2008]. Available from: [http://www.ibge.gov.br/home/estatistica/populacao/censo2000/populacao/tabela\\_brasil.shtm](http://www.ibge.gov.br/home/estatistica/populacao/censo2000/populacao/tabela_brasil.shtm).
  24. Namazi MR. Why is psoriasis uncommon in Africans? The influence of dietary factors on the expression of psoriasis *Int J Dermatol.* 2004;43:391–2.
  25. Bonfiglioli R, Conde RA, Sampaio-Barros PD, Louzada-Junior P, Donadi EA, Bertolo MB. Frequency of HLA-B27 alleles in Brazilian patients with psoriatic arthritis. *Clin Rheumatol.* 2008;27:709–12.
  26. Green AC. Australian aborigines and psoriasis. *Australas J Dermatol.* 1984;25:18–24.
  27. Lomholt G. Prevalence of skin diseases in a population: A census study from the Faroe Islands. *Dan Med Bull.* 1964;11:1–7.
  28. Verhagen AR, Kolen JW. Psoriasis in Kenya. *Arch Dermatol.* 1967;96:39–41.
  29. Yip SY. The prevalence of psoriasis in the Mongoloid race. *J Am Acad Dermatol.* 1984;10:965–8.
  30. Plunkett A, Marks R. A review of the epidemiology of psoriasis vulgaris in the community. *Australas J Dermatol.* 1998;39:225–32.
  31. Bowcock AM. The genetics of psoriasis and autoimmunity. *Annu Rev Genomics Hum Genet.* 2005;6:93–122.
  32. Bell LM, Sedlack R, Beard CM, Perry HO, Michet CJ, Kurland LT. Incidence of psoriasis in Rochester, Minn, 1980–1983. *Arch Dermatol.* 1991;127:1184–7.
  33. Farber E, Nail M. The natural history of psoriasis in 5,600 patients. *Dermatologica.* 1974;148:1–18.
  34. Ferrándiz C, Bordas X, García-Patos V, Puig S, Pujol R, Smandiá A, et al. Prevalence of psoriasis in Spain (Epiderma Project: phase I). *J Eur Acad Dermatol Venereol.* 2001;15:20–3.
  35. Raychaudhuri SP, Farber EM. The prevalence of psoriasis in the world. *J Eur Acad Dermatol Venereol.* 2001;15:16–7.
  36. Gelfand JM, Weinstein R, Porter SB, Neimann AL, Berlin JA, Margolis DJ. Prevalence and treatment of psoriasis in the United Kingdom. A population-based study. *Arch Dermatol.* 2005;141:1537–41.
  37. Hartshorne ST. Dermatological disorders in Johannesburg, South Africa. *Clin Exp Dermatol.* 2003;28:661–5.
  38. Saraceno R, Mannheimer R, Chimenti S. Regional distribution of psoriasis in Italy. *J Eur Acad Dermatol Venereol.* 2008;22:324–9.
  39. Conselho Nacional de Secretários de Saúde. Legislação do SUS. Brazil [Accessed on July 28, 2009]. Available from: [http://bvsms.saude.gov.br/bvs/publicacoes/progestores/leg\\_sus.pdf](http://bvsms.saude.gov.br/bvs/publicacoes/progestores/leg_sus.pdf).
  40. Arruda LH, de Moraes AP. The impact of psoriasis on quality of life. *Br J Dermatol.* 2001;144(Suppl.):58:33–6.