Indian Journal of

Dermatology, Venereology & Leprology

Vol 74 | Issue 1 | Jan-Feb 2008

The Indian Journal of Dermatology, Venereology and Leprology (IJDVL) is a bimonthly publication of the Indian Association of Dermatologists, Venereologists and Leprologists (IADVL) and is published for IADVL by Medknow Publications.

The Journal is **indexed/listed** with Science Citation Index Expanded, PUBMED, EMBASE, Bioline International, CAB Abstracts, Global Health, DOAJ, Health and Wellness Research Center, SCOPUS, Health Reference Center Academic, InfoTrac One File, Expanded Academic ASAP, NIWI, INIST, Uncover, JADE (Journal Article Database), IndMed, Indian Science Abstract's and PubList.

All the rights are reserved. Apart from any fair dealing for the purposes of research or private study, or criticism or review, no part of the publication can be reproduced, stored, or transmitted, in any form or by any means, without the prior permission of the Editor, IJDVL.

The information and opinions presented in the Journal reflect the views of the authors and not of the IJDVL or its Editorial Board or the IADVL. Publication does not constitute endorsement by the journal.

The IJDVL and/or its publisher cannot be held responsible for errors or for any consequences arising from the use of the information contained in this journal. The appearance of advertising or product information in the various sections in the journal does not constitute an endorsement or approval by the journal and/or its publisher of the quality or value of the said product or of claims made for it by its manufacturer.

The journal is published and distributed by Medknow Publications. Copies are sent to subscribers directly from the publisher's address. It is illegal to acquire copies from any other source. If a copy is received for personal use as a member of the association/society, one can not resale or give-away the copy for commercial or library use.

The Journal is printed on acid free paper.

EDITOR

Uday Khopkar

ASSOCIATE EDITORS

Ameet Valia Sangeeta Amladi

ASSISTANT EDITORS

K. C. Nischal

Sushil Pande Vishalakshi Viswanath

EDITORIAL BOARD

Chetan Oberai (Ex-officio) Arun Inamdar Binod Khaitan D. A. Satish D. M. Thappa H. R. Jerajani Koushik Lahiri (Ex-officio) Joseph Sundharam Kanthraj GR M. Ramam Manas Chatterjee Rajeev Sharma Sandipan Dhar Sanjeev Handa S. L. Wadhwa Sharad Mutalik Shruthakirti Shenoi Susmit Haldar Venkatram Mysore

EDITORIAL ADVISORY BOARD

Aditya Gupta, Canada C. R. Srinivas, India Celia Moss, UK Giam Yoke Chin, Singapore Gurmohan Singh, India Howard Libman, USA J. S. Pasricha, India Jag Bhawan, USA John McGrath, UK K. Pavithran, India R. G. Valia, India Robert A. Schwartz, USA Robin Graham-Brown, UK V. N. Sehgal, India

Rodney Sinclair, Australia

STATISTICAL EDITOR

OMBUDSMAN

S. R. Suryawanshi

A. K. Bajaj

IADVL NATIONAL EXECUTIVE 2006 - 2007

President Chetan M. Oberai

Immediate Past President Suresh Joshipura

EDITORIAL OFFICE

Dr. Uday Khopkar

Editor, IJDVL, Department of Dermatology,

117, 1st Floor, Old OPD Building, K.E.M.

Hospital, Parel, Mumbai - 400012, India.

E-mail: editor@ijdvl.com

President (Elect) S. Sacchidanand

Vice-Presidents Amrinder Jit Kanwar Dilip Shah Secretary Treasurer

Secretary Koushik Lahiri

Jt. Secretaries Rakesh Bansal Manas C

Manas Chatterjee

Arijit Coondoo

Published for IADVL by

MEDKNOW PUBLICATIONS

A-109, Kanara Business Centre, Off Link Road, Ghatkopar (E), Mumbai - 400075, India. Tel: 91-22-6649 1818 / 1816 Website: www.medknow.com

> www.ijdvl.com www.journalonweb.com/ijdvl www.bioline.org.br/dv

ISSN 0378-6323 E-ISSN 0973-3930

Indian Journal of

Dermatology, Venereology & Leprology

Journal indexed with SCI-E, PubMed, and EMBASE

Issue 1 Jan-Feb 2008	C	0	N	Т	E	N	
DITORIAL REPORT - 2007							
IIDVL gets into the Science Citation Index Expanded! Uday Khopkar							1
DITORIAL							
Registration and reporting of clinical trials Uday Khopkar, Sushil Pande							2
PECIALTY INTERFACE							
Preventing steroid induced osteoporosis Jyotsna Oak							5
EVIEW ARTICLE							
Molecular diagnostics in genodermatoses - simplified Ravi N. Hiremagalore, Nagendrachary Nizamabad, Vijayaraghavan Kamasam	udram						8
RIGINAL ARTICLES							
A clinicoepidemiological study of polymorphic light eruption Lata Sharma, A. Basnet							15
A clinico-epidemiological study of PLE was done for a period of one year to includ between IV and VI. The manifestation of PLE was most common in house wives or patients of PLE presented with mild symptoms and rash around neck, lower forea aggravated on exposure to sunlight. PLE was more prevalent in the months of Man disease was recurrent in 31.36% of cases.	sun exposers	ed ar ns w	eas. N hich v	Aost o was	of the	9	
Comparative study of efficacy and safety of hydroxychloroquine ar light eruption: A randomized, double-blind, multicentric study Anil Pareek, Uday Khopkar, S. Sacchidanand, Nitin Chandurkar, Geeta S. Naik		_				_	c 18
In a double-blind randomized, comparative multicentric study evaluating efficacy light eruption, a total of 117 patients of PLE were randomized to receive hydroxyo tablets for a period of 2 months (initial twice daily dose was reduced to once daily reduction in severity scores for burning, itching, and erythema was observed in p hydroxychloroquine as compared to chloroquine. Hydroxychloroquine was found studied with lesser risk of ocular toxicity.	hloroquine after 1 mo atients trea	and nth). ted v	chlor A sig vith	oquin	ne ant		ıg

Many faces of cutaneous leishmaniasis Arfan Ul Bari, Simeen Ber Rahman

Symptomatic cutaneous leishmaniasis is diverse in its presentation and outcome in a tropical country like Pakistan where the disease is endemic. The study describes the clinical profile and atypical presentations in 41 cases among 718

patients of cutaneous leishmaniasis. Extremity was the most common site of involvement and lupoid cutaneous leishmaniasis was the most common atypical form observed. Authors suggest that clustering of atypical cases in a geographically restricted region could possibly be due to emergence of a new parasite strain.

Forehead plaque: A cutaneous marker of CNS involvement in tuberous sclerosis G. Raghu Rama Rao, P. V. Krishna Rao, K. V. T. Gopal, Y. Hari Kishan Kumar, B. V. Ramachandra

In a retrospective study of 15 patients of tuberous sclerosis, eight patients had central nervous system involvement. Among these 8 cases, 7 cases had forehead plaque. This small study suggests that presence of forehead plaque is significantly associated with CNS involvement.

BRIEF REPORTS

Ligand-binding prediction for ErbB2, a key molecule in the pathogenesis of leprosy Viroj Wiwanitkit......

SCORTEN: Does it need modification? Col. S. S. Vaishampayan, Col. A. L. Das, Col. R. Verma

CASE REPORTS

Universal acquired melanosis (Carbon baby) P. K. Kaviarasan, P. V. S. Prasad, J. M. Joe, N. Nandana, P. Viswanathan

Adult onset, hypopigmented solitary mastocytoma: Report of two cases D. Pandhi, A. Singal, S. Aggarwal.....





32

28

35

38





41

23

59

CONTENTS (Contd.)

Incidental finding of skin deposits of corticosteroids without associated granulomatous inflammation: Report of three cases Rajiv Joshi

Erythromelanosis follicularis faciei *et* **colli: Relationship with keratosis pilaris** M. Augustine, E. Jayaseelan.....

Naxos disease: A rare occurrence of cardiomyopathy with woolly hair and palmoplantar keratoderma R. Rai, B. Ramachandran, V. S. Sundaram, G. Rajendren, C. R. Srinivas.....

Granular parakeratosis presenting with facial keratotic papules	
R. Joshi, A. Taneja	

Adult cutaneous myofibroma V. Patel, V. Kharkar, U. Khopkar

LETTERS TO THE EDITOR

Extragenital lichen sclerosus of childhood presenting as erythematous patches N. G. Stavrianeas, A. C. Katoulis, A. I. Kanelleas, E. Bozi, E. Toumbis-Ioannou...

Leukocytoclastic vasculitis during pegylated interferon and ribavirin treatment of hepatitis C virus infection Esra Adisen, Murat Dizbay, Kenan Hize, Nilsel İlter.....











56





44

47

CONTENTS (Contd.)

Poland's syndrome Saurabh Agarwal, Ajay Arya	62
Hereditary leiomyomatosis with renal cell carcinoma Sachin S. Soni, Swarnalata Gowrishankar, Gopal Kishan Adikey, Anuradha S. Raman	63
Infantile onset of Cockayne syndrome in two siblings Prerna Batra, Abhijeet Saha, Ashok Kumar	65
Multiple xanthogranulomas in an adult Surajit Nayak, Basanti Acharjya, Basanti Devi, Manoj Kumar Patra	67
Bullous pyoderma gangrenosum associated with ulcerative colitis Naik Chandra Lal, Singh Gurcharan, Kumar Lekshman, Lokanatha K	68
Sporotrichoid pattern of malignant melanoma Ranjan C. Rawal, Kanu Mangla	70
Acitretin for Papillon-Lefèvre syndrome in a five-year-old girl Didem Didar Balci, Gamze Serarslan, Ozlem Sangun, Seydo Homan	71
Bilateral Becker's nevi Ramesh Bansal, Rajeev Sen	73

Madarosis: A dermatological marker Silonie Sachdeva, Pawan Prasher

74

CONTENTS (Contd.)

FOCUS

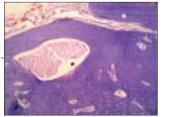
Botulinum toxin	
Preeti Savardekar	77

E-UDVL

Net Studies A study of oxidative stress in paucibacillary and multibacillary leprosy	
P. Jyothi, Najeeba Riyaz, G. Nandakumar, M. P. Binitha	80
Clinical study of cutaneous drug eruptions in 200 patients M. Patel Raksha, Y. S. Marfatia	80
Net case	
Porokeratosis confined to the genital area: A report of three cases	
Sujata Sengupta, Jayanta Kumar Das, Asok Gangopadhyay	80
Net Letters	
Camisa disease: A rare variant of Vohwinkel's syndrome	
T. S. Rajashekar, Gurcharan Singh, Chandra Naik, L. Rajendra Okade	81
Cross reaction between two azoles used for different indications	
Arika Bansal, Rashmi Kumari, M. Ramam	81
Net Quiz	
Asymptomatic erythematous plaque on eyelid	
Neeraj Srivastava, Lakhan Singh Solanki, Sanjay Singh	82

QUIZ

A bluish nodule on the arm Ragunatha S., Arun C. Inamadar, Vamseedhar Annam, B. R. Yelikar.....



83

REFEREE INDEX-2007

INSTRUCTIONS FOR AUTHORS

The copies of the journal to members of the association are sent by ordinary post. The editorial board, association or publisher will not be responsible for non-receipt of copies. If any of the members wish to receive the copies by registered post or courier, kindly contact the journal's / publisher's office. If a copy returns due to incomplete, incorrect or changed address of a member on two consecutive occasions, the names of such members will be deleted from the mailing list of the journal. Providing complete, correct and up-to-date address is the responsibility of the members. Copies are sent to subscribers and members directly from the publisher's address; it is illegal to acquire copies from any other source. If a copy is received for personal use as a member of the association/society, one cannot resale or give-away the copy for commercial or library use.

A clinicoepidemiological study of polymorphic light eruption

Lata Sharma, A. Basnet

Department of Dermatology and Venereology, Institute of Medical Sciences, Banaras Hindu University, Varanasi, India

Address for correspondence: Dr. Lata Sharma, Department of Dermatology and Venereology, Institute of Medical Sciences, Banaras Hindu University, Varanasi -221 005, India. E-mail: lataims@rediffmail.com

ABSTRACT

Background and Aims: The prevalence of polymorphic light eruption (PLE) varies between 10-20% in different countries but no such data is available from India, where exposure to sunlight is high. **Methods:** A clinico-epidemiological study of PLE was done in the skin outpatient department (OPD) of Institute of Medical Sciences Hospital from January to December. **Results:** The ages of the patients varied from 5-70 years. Out of a total of 39,112 OPD cases, 220 cases of PLE (138 females and 82 males) were recorded, giving a prevalence of 0.56% in this study population. The skin type varied between IV and VI in 96% of the cases. Housewives were 81, students 67, office persons 39, farmers 22, businessmen 6 and unemployed 5. **Discussion:** The manifestation of PLE was most common in housewives in areas exposed to the sun. Most of the PLE patients presented with mild symptoms and rash around the neck, forearms and arms which was aggravated on exposure to sunlight. PLE was more prevalence of PLE was 0.56%. It was mild in nature and only areas exposed to the sun were involved.

Key Words: Polymorphic light eruption, Clinico-epidemiological study, Photodermatosis

INTRODUCTION

Polymorphic Light Eruption is an abnormal cutaneous response which occurs on exposure to sunlight. The reported prevalence in various cohort studies from England, Sweden and Singapore varies from 10-20% and 26% in a study by Fotiades *et al.*^[1-5] No such data however, is available from our country even though the exposure to sunlight is high.

METHODS

A clinico-epidemiological study was done in patients who had been visiting our Dermatology OPD for a period of one year. Subjects taking systemic corticosteroids or photosensitizing drugs were excluded. Thus, 220 cases of PLE were registered from January to December. Patient details recorded included month and age of onset of symptoms of PLE, its severity, nature - transient, persistent or recurrent, aggravating factors, constitutional and other symptoms, results of healing of the rash and any change in the severity of symptoms. History of the disease in the family, the patient's profession, duration of exposure to sunlight during outdoor activities including travel, preference for the type of clothing; materials used during daytime - cosmetics and sunscreens, as well as types of previous treatments were noted.

Findings of the clinical examination were recorded including the skin type of cases as classified by the Fitzpatricks skin phototype scale.^[6] Details of skin lesions and the site, size, shape, color, type and secondary changes were noted. In doubtful cases, patients were asked to protect the affected part from sun exposure for 7-10 days during which the lesions of PLE should have healed without scarring. Immunological and urine examinations were done to exclude systemic lupus erythematosus and porphyria. Data thus obtained was compiled, tabulated and statistically summarized.

How to cite this article: Sharma L, Basnet A. A clinicoepidemiological study of polymorphic light eruption. Indian J Dermatol Venereol Leprol 2008;74:15-7.

Received: December, 2006. Accepted: August, 2007. Source of Support: Nil. Conflict of Interest: None D.

RESULTS

Out of the total 39,112 patients who attended the Dermatology OPD, 220 patients had PLE. The prevalence of PLE was thus calculated to be 0.56%, which included 138 females and 82 males. The age of the patients varied from 5 to 70 years with mean \pm standard deviation of 29.90 \pm 12.22 years. The ages of 131 (59.55%) cases were \leq 30 years and the ages of 76 (34.54%) cases were between 31 and 50 years.

Eighty one cases were housewives, 67 were students 39 were office persons, 22 were farmers, 6 businessmen and 5 were unemployed. Skin rash was present in all cases along with itching in 151, burning sensation in 20, both itching and burning in 13 while 36 were asymptomatic. Constitutional symptoms like fever and malaise were present in 11 cases, headache in 3 and swelling of the face in 1. The rash appeared on exposure to sunlight within 30 minutes in 65 cases and after > 30 minutes in 20 cases, on termination of exposure in 20 but the time interval was not known in 115 patients. The aggravating factor was sunlight in 103 cases, the heat of an open fire while cooking food in 6, unidentifiable in 69 cases and both sunlight and heat in 42 cases. The month of onset of rash varied from January to December. The months of presentation were March and September for 46 and 33 cases, respectively [Table 1]. There was history of recurrence in 99 cases, severity of rash increased in 69, decreased in 23 and no change was noted in 7 cases. History of atopy was present in 5 cases and in family in 23 cases. History of PLE was present in family in 22 cases including 2 each in the grandfather, father, brother and son, in the mother in 6 and in the sister in 8 cases. The material of clothing used was of a mixed type in 172 cases, cotton in 35 and synthetic in 13. In the history of treatment, 40 patients used a topical steroid, 24 antifungal, 13 sunscreens, 4 oral steroids, 19 antihistaminic, and eight used homeopathic drugs. The skin type of patients varied from III to VI with a maximum of 107 (48.64%) cases in type IV, 72 (32.73%) in V, 33 (15%) in VI and only 1 (0.45%) in type I. Exposed parts were involved in all cases. The rash was papular in most of the cases [Table 2]. The majority of the lesions were erythematous [Table 3]. Acne was associated in 42 patients, melasma in 11 and tinea corporis in 7 patients.

DISCUSSION

PLE is considered to be a disease of fair-skinned individuals with skin types I to IV.^[2] It is less common in very dark-skinned individuals in America, India and Pakistan.^[7] In

our study, 96% of the patients were of skin types IV to VI which explains its low prevalence (0.56%). The majority of the cases were in the age group of 21-30 years consistent with the existing idea that PLE is a disease of the first three decades of life.^[8] Of the cases suffering from PLE, 62.73% were females. Most of them were housewives in whom exposure to sunlight was intermittent and for a short period. The clothing used was light which gave full exposure to the neck, arms and forearms. Partial ultraviolet radiation-induced immunosuppression is said to result in a delayed type hypersensitivity response to photoinduced antigens.^[9] PLE lesions fade off near or sometimes sharply at the borders of garments but not all exposed areas are involved. It is thought that exposure of those areas throughout the year makes them more tolerant.^[10]

The onset of PLE was in the months of February and August in 13.18 and 11.82% of the cases, respectively. In March and September, 20.91 and 15% of the cases were recorded which was high when compared to the other

Table 1: Month wise distribution of PLE					
Month	Presentation		Onset		
	n	%	n	%	
January	0	0	16	7.27	
February	23	10.45	29	13.18	
March	46	20.91	23	10.45	
April	20	9.09	14	6.36	
May	03	1.36	16	7.27	
June	15	6.82	19	8.64	
July	21	9.55	17	7.73	
August	15	6.82	26	11.82	
September	33	15.00	22	10.00	
October	22	10.00	17	7.73	
November	17	7.73	13	5.91	
December	5	2.27	8	3.64	
Total	220	100.00	220	100.00	

Table 2: Site and type of lesion and number of PLE patients						
Site	n	%	Туре	n	%	
Neck	136	61.82	Papule	119	54.09	
Arm	121	55.00	Macule	43	19.55	
Forearm	105	47.73	Plaque	9	4.09	
Face	74	33.64	Vesicle	1	0.45	
Back	21	9.55	Macule + papule	10	4.55	
Neck 'V'	19	8.64	Papule + Plaque	33	15.00	
Hand	13	5.91	Macule + papule + plaque	5	2.27	
Leg	4	1.82				
Abdomen	1	0.45				
Ear	1	0.45				
	n=220		Total	220	100.00	
	11-220		10(2)	220	100.00	

Table 3: Color of the lesion and number of PLE patients					
n	Percentage				
68	30.91				
35	15.91				
32	14.55				
18	8.18				
37	16.82				
13	5.91				
12	5.45				
5	2.27				
220	100.00				
	n 68 35 32 18 37 13 12 5				

months. During these months, patients used light clothing with short sleeves during outdoor activity. This is the time when the sun shines on the equator and the days and nights are of almost equal length. Hawk has described PLE as a disease of the spring when moderate intensity of sunlight causes maximum hazard.^[11] The action spectrum of PLE is said to include both long and short wavelength ultraviolet radiation but in a study by Pryzbilla and coworkers, all cases tested were negative for a ultraviolet B photopatch test.^[8]

The majority of the cases were from the city of Varanasi where the study was conducted. Its latitude is 25° north and longitude 83° east. The prevalence of PLE is said to be higher in regions away from the equator because of the variation in the proportion of ultraviolet A and B radiations at different latitudes.^[5] Sunlight was the precipitating factor in 46.82% of the cases, the heat of an open fire in 2.73%, both of these in 19.09% although the precipitating factor was not known in 31.36% of the cases. Consistent with a report by Jansen, we found that a 30 min exposure to sunlight was required to produce the rash, the interval being slightly less than $\frac{1}{2}$ hour in 29.55% of cases, more than $\frac{1}{2}$ hour in 9.09%, but 52.27% were not aware of this.^[12]

The external aspect of the arms and forearms were involved in most of the cases possibly because these parts are placed horizontally while sitting or traveling and receive the maximum exposure. On the other hand, the position of the face is vertical while walking or working or it may not even be exposed to the sun if the person is bending forward. The exposure of covered areas in the summer months makes them vulnerable to this photodermatosis.^[113] The decline in the severity of eruption or rash on repeated sun exposure or as summer progresses, causes the observed hardening.^[12] The rash was recurrent in 99 cases but an increase in the severity of lesions was noted by 23 cases. It appears that the disease did not manifest fully in the beginning. The type of lesion was papular or macular in this study as compared to the vesicular and plaque types and extensive involvement as reported by Hawk.^[11] Fewer cases had itching, burning and constitutional symptoms than reported by Frain-Bell *et al.*^[8] It may be because the disease was milder in this part of the world. Covered areas were not affected irrespective of the type of clothing or weave tightness which suggests that it is probably preventable by all types of clothing.^[12]

Inheritance of PLE is probably polygenic or through a dominant single gene.^[13] Family history of PLE was found in 10% of the cases in the present study but varied from 6.25-12% in the studies conducted by Ross^[2] and Millard.^[13] As it is a mild photodermatosis in India, many patients were not aware of its occurrence in other members of the family.

REFERENCES

- Morison WL, Stern RS. Polymorphous light eruption: A common reaction uncommonly recognized. Acta Derm Venereol 1982;62:237-40.
- 2. Ros AM, Wennersten G. Current aspects of Polymorphous light eruptions in Sweden. Photodermatol 1986;3:298-302.
- 3. Khoo SW, Tay YK, Tham SN. Photodermatoses in a Singapore skin referral centre. Clin Exp Dermatol 1996;21:263-8.
- Pao C, Norris PG, Corbett M, Hawk JL. Polymorphic light eruption: Prevalence in Australia and England. Br J Dermatol 1994;130:62-4.
- 5. Fotiades J. Soter NA, Lum HW. Result of evaluation of 203 patients for photosensitivity in 3 years period. J Am Acad Dermatol 1995;33:597-602.
- 6. Fitzpatrick TB. The validity and practicality of sun reactive skin type I through VI. Arch Dermatol 1988;124:869.
- Magnus, IA. Dermatological photobiology: Clinical and experimental aspects. Blackwell Scientific Publications: Oxford; 1976. p. 117-63.
- Frain-Bell W, Dickson A, Herd J, Sturrock I. The action spectrum in polymorphic light eruption. Br J Dermatol 1973;89:243.
- 9. Palmer RA, Fredman PS. Ultraviolet radiation causes less immunosuppression in patients with polymorphic light eruption than in controls. J Invest Dermatol 2004;122:291-4.
- Ting WW, Vest CD, Sontheimer R. Practical and experimental consideration of sun protection in dermatology. Int J Dermatol 2003;42:505-13.
- 11. Hawk JLM, Norris PG. Abnormal responses to ultraviolet radiation. In: Freedberg IM, Eisen AZ, Wolff K, Austen KF, Goldsmith LA, Katz SI, editors, Fitzpatrick's Dermatology in General Medicine. 5th ed. McGraw-Hill: New York; 1999. p. 1:1573-89.
- 12. Jansen CT. The Natural history of Polymorphous light eruptions. Arch Dermatol 1979;115:165-9.
- 13. Millard TP, Bataille V, Snieder H, Spector TD, McGregor JM. The heritability of polymorphic light eruption. J Invest Dermatol 2000;115:467-70.