



## A Clinico Etiological Study of Cutaneous Drug Eruptions

Authors

**Dr M. Nithya Kumari M.D., Dr K.Sridevi, M.D., Associate Professor,  
Dr G.Ramesh Kumar, Professor**

Corresponding Author

**Dr K.Sridevi**

Department of D.V.L., S.V. Medical College, Tirupati

Email: [kotasridevidr@gmail.com](mailto:kotasridevidr@gmail.com)

### ABSTRACT

**Background:** *To study the clinical spectrum of Cutaneous adverse drug reactions (ACDRS) in hospitalized patients and to establish a causal link between the drug and the reaction by using WHO causality definitions.*

#### Objectives:

1. *To calculate prevalence of ACDRS*
2. *To evaluate various etiological factors*
3. *To study clinical patterns*

**Methods:** *All patients attending to the OPD and IPD of D.V.L. and patients referred from other departments with suspected ACDRs were included in the study.*

**Results:** *In the present study most common offending drug group is antimicrobials (48.6%) followed by 20% of anticonvulsants, 12.8% cases of NSAIDs and 18.6% others. Common morphological types of ACDRs were acneiform eruptions, F.D.E, maculopapular rash, TEN-SJS, Dapsone syndrome and Pruritis.*

**Keywords:** *Adverse Cutaneous drug reactions- Etiology- Drug eruptions*

### INTRODUCTION

The drug eruptions are unwanted and unintended mucocutaneous reactions which occur on administration of diagnostic or therapeutic agent.

The prevalence of drug reactions depends on many factors including genetic and racial factors. Some drugs which are banned in developed countries are still in use in India. In addition, India is known for its indigenous medicines which can also be a source of drug eruptions.

Incidence of drug eruptions in our country varies between 6 to 30 % and about 8% hospital

admission are due to drug eruptions. Internationally drug eruption occurs in 2-3% of Patients.

Drug eruptions range from pruritus to severe life-threatening Stevens-Johnson Syndrome (SJS) or toxic epidermal necrolysis.

The diagnosis of cutaneous eruption is based on detailed history and correlation between drug intake and the onset of rash. Cutaneous drug eruptions are visible and hence reporting is earlier and better when compared to reactions of internal organs and other systems.

## MATERIALS AND METHODS

### Source of Data

The present prospective study was done over a period of one year from January 2013 to 2014 in the department of D.V.L, S.V. Medical College, Tirupati.

The study cases are all patients of either sex attending the D.V.L Department, patients referred from other departments with suspected ACDRs.

The protocol was approved by Human Research Ethics Committee (HREC) of S.V.Medical College, Tirupati.

All cases of ACDRs willing to participate and having causality assessment scale are included in the study.

Patients not willing to participate in the study and patients dropping out the study were excluded. Investigations like complete hemogram, L.F.T., R.F.T., Skin Biopsy, VDRL and HIV testing were carried out

### The Criteria for the diagnosis of ACDRs

1. The time interval between the introduction of the drug and the onset of reaction should be within specific time described in literature for each reaction.  
For example Maculopapular rash : <7 days, Urticaria : 7-21 days, SJS-TEN : 1-3 weeks etc.
2. Improvement in the condition of patient after withdrawal of drug .
3. Based on the W.H.O causality definition, ACDRs were categorised as Certain, Probable, Possible and Unlikely

## RESULTS AND DISCUSSION

A total 22,073 patients attended department of D.V.L. during study period of which 70 cases had ACDRs.

The prevalence ratio is 0.0032% in the study, mean age of patients is 33.47%. In this study most of them i.e. 32/70 were in the age group of 21-40 years followed by 14/70 in 41-50 years of age, then 10/70 in 11-20 years of age, 8/70 in the age

group of >50 years and 6/70 cases were 0-10 years of age.

The male to female ratio 2.181(48 male and 22 female).

### Morphological Types of ACDRs

| Type of Drug Reaction                                   | No. of Cases | Percentage (%) |
|---|--------------|----------------|
| Acne form eruptions                                     | 17           | 24.28          |
| Fixed drug eruptions                                    | 13           | 18.52          |
| Morbilliform rash                                       | 10           | 14.28          |
| Toxic Epidemolytic Necrolysis                           | 6            | 8.75           |
| Dapsone syndrome  | 3            | 4.28           |
| Erythema multiforme                                     | 1            | 1.43           |
| Purpura   | 1            | 1.4            |
| Exfoliative dermatitis                                  | 2            | 2.85           |
| Pruritis  | 3            | 4.28           |
| Stevens Johnson Syndrome                                | 4            | 5.71           |
| Urticaria   | 2            | 2.85           |
| Lichenoid eruptions                                     | 2            | 2.85           |
| Psoriasisiform eruptions                                | 2            | 2.85           |
| Peeling of palmar skin                                  | 1            | 1.43           |
| Acute generalized exanthematouspustulosis               | 1            | 1.43           |
| Eczematous eruption                                     | 1            | 1.43           |
| Drug rash with eosinophilia & systemic symptoms (DRESS) | 1            | 1.43           |
| TOTAL   | 70           | 100            |

ACDRs vary in the pattern of morphology and distribution. Acneiform eruptions were the most common drug eruptions (24.28%) folloed by fixed drug eruptions (18.52%).



**Acneiform Eruptions**



**Exfoliates dermatitis**

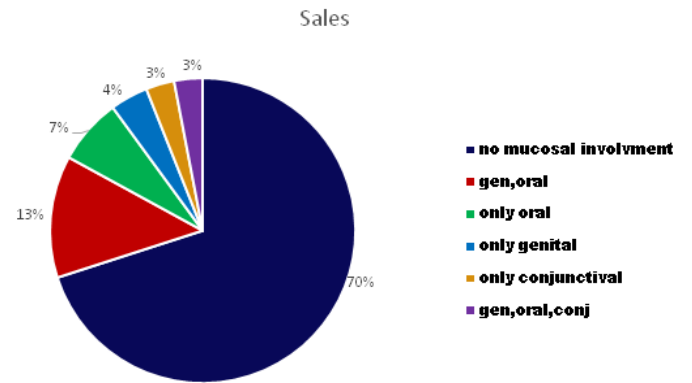


**Fixed Drug eruption on Lip**



**Carbamazepine Induced Maculopapular Rash**

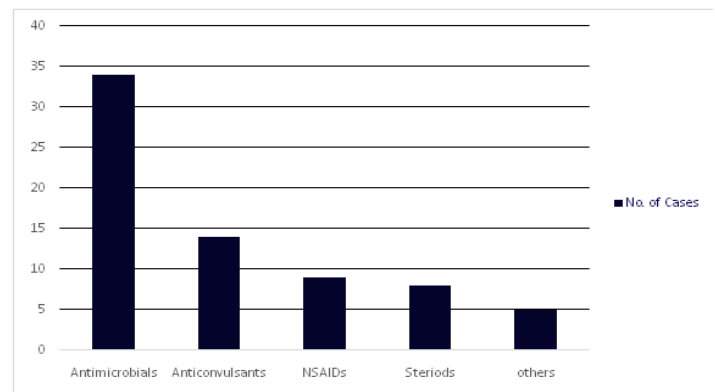
**Mucosal Involvement in ACDRs**



In our study, visceral involvement may predict poor prognosis seen in patients with S.J.S.-TEN and Dapsone syndrome.

Commonly incriminated drugs were Phenytoin (14.28%) followed by diclofenac (10%) followed by cotrimoxazole (7.21%).

When looked at drug groups macrolides 48% (13/70 cases) followed by antiepileptics 20% (14/70 cases), NSAIDs (7/70)cases, steroids 11.4% (8/70) cases. This is in concordance with an earlier report from North India.



Antimicrobials and NSAIDs are commonly prescribed drugs by physicians and general practitioners, even quacks for trivial illness so there are more chances of developing reactions to these groups.

**CONCLUSION**

After a cutaneous drug eruption has been diagnosed and treated, clear information must be provided to the patient regarding his/her drug rash.

Advised the patient to carry a card, informing about their reaction and the offending drug is necessary.

Finally cutaneous drug reaction are to be reported to the manufacturer and regulatory agency especially if skin eruption is rare, serious or unexpected.

Drug reactions are common reason for litigation, warning a patient about potential adverse effects and before prescribing a medicine previous history of drug eruption should be taken.

## REFERENCES

1. Litt JZ. Litt's D.E.R.M. Drug Eruption and Reactions Manual, 14 thEdi.London: Informa Healthcare ; 2008.VI.
2. Raksha MP, Marfatia YS. Clinical Study of cutaneous drug eruptions in 200 patients. Indian J Dermatol Venereol Leprol. 2008 Jan-Feb; 74 (1):80.
3. Hotchandani SC, Bhatt JD, Shah MK. A prospective analysis of drug-induced acute cutaneous reactions reported in patients at a tertiary care hospital. Indian J Pharmacol. 2010 Apr;42(2): 118-9.
4. BigbyM.Rates of Cutaneous reactions to drugs. Arch Dermatol.2001 Jun; 137(6): 765-70.
5. Craig KS, Edward WC, Anthony AG. Cutaneous drug reactions. Pharmacol Rev 2001; 53: 57-79.
6. Noel MV, Sushma M, Guido S. Cutaneous adverse drug reactions in hospitalized patients in a tertiary care centre. Indian J Pharmacol 2004; 36:292-5.
7. Pudukadan D, Thappa DM. Adverse cutaneous drug reactions : Clinical pattern and causative agents in a tertiary care centre in South India. India J DermatolVenereolLeprol. 2004 Jan -Feb; 70 (1):20-4.
8. Lazarou J, Pomeranz BH, Corey PN. Incidence of adverse drug reactions in hospitalized patients: a meta-analysis of prospective studies. JAMA. 1998 Apr 15; 279(15): 1200-5.
9. Chatterjee S, Gosh AP, Barbhuiya J, Dey SK. Adverse cutaneous drug reactions: A one year survey at a Dermatology outpatient clinic of a tertiary care hospital. Indian J Pharmacol 2006; 38: 429-31.
10. Ajayi FO, Sun H, Perry J . Adverse drug reactions : a review of relevant factors. J ClinPharmacol. 2000 Oct; 40(109):1093-101.
11. Sullivan JR, Shear NH. Drug eruptions and other adverse drug effects in aged skin. ClinGeriatr Med. 2002 Feb; 18(1): 21-42.
12. Solensky R, Mendelson LM. Systemic reactions to antibiotics. Immunol Allergy Clin N Am. 2001; 21: 679-97.
13. Anjaneyan G, Gupta R, Vora RV. Clinical study of adverse cutaneous drug reactions at a rural based tertiary care centre in Gujarat. Nati J Physiol Pharm Pharmacol 2013; 3:129-136.