A Study of Efficacy of Topical Phenytoin in the Management of Diabetic Ulcer

^{*}Dr.A.M.Syed Ibrahim ¹M.S.,Dr.R. Ramakrishnan² M S, Dr.Mohamed Yasar ³m.S.,

¹Professor, Department of General Surgery, Madurai Medical College And Hospital, Madurai, Tamil Nadu, India, ²POST GRADUATE, Department of General Surgery, Madurai Medical College And Hospital, Madurai, Tamil Nadu, India ³Post Graduate, Department of General Surgery, Madurai Medical College and Hospital, Madurai, Tamil Nadu, India. Corresponding author: *Dr.A.M.Syed Ibrahim

Abstract

Background: Diabetic ulcer is the most frequent reason for hospitalization in patients with diabetes. It has increased the cost of treatment and hospitalization of these patients. Currently a lot of attention is being placed on the development of expensive topical growth factors for wound healing. Thus there remains a quest for better wound healing agents. One such agent is phenytoin which is cheap, easy to use and readily available for medical practice. Phenytoin (diphenylhydantoin) was initially introduced into therapy for the effective control of convulsive disorders. A common side effect with systemic phenytoin treatment is the development of fibrous overgrowth of gingiva. This apparent stimulatory effect of phenytoin on connective tissue suggested an encouraging possibility for its use in wound healing. To study the efficacy of topical phenytoin dressing over conventional dressing in the management of diabetic ulcers, in terms of days required for healing, rate of granulation tissue formation, quality of graft bed, graft uptake, effect on bacterial growth and side effects of topical phenytoin.

Method: All patients undergo detailed clinical examination and relevant investigations and the wounds are thoroughly debrided. The surface area is measured tracing the outline on transparent paper. This outline is transferred to a graph paper for area measurement. The patients are followed up on a daily basis for 14 days in both study and control groups. The control group and study group are subjected to twice-daily dressing. Both the groups are first cleaned with normal saline and 5% povidone iodine. The study group is then subjected to topical phenytoin dressing. Sterile gauze is soaked in preparation of i.v. phenytoin and placed over the wound Once these parameters are assessed, both the groups subjected to split thickness skin grafting. Both groups are given the same systemic antibiotics during the postoperative period. Both the groups were compared in terms of discharge, slough, wound area reduction and duration of

hospital stay.

Results: This study has shown better granulation tissue formation, graft uptake, negative bacterial growth and decreased hospital stay in patients receiving phenytoin dressing than patients receiving betadine dressing. There were no side effects in patients with phenytoin dressing.

Conclusion: Phenytoin is a cheap, readily available and easy to use alternative in the treatment of diabetic ulcers.

Keyords: Diabetic ulcers, phenytoin dressing, betadine dressing, granulation tissue, graft uptake, graft bed, bacterial growth, hospital stay, phenytoin side effects.

Date of Submission: 01 -08-2017Date of acceptance: 23-08-2017

I. Introduction

Diabetes mellitus comprise a group of common metabolic disorder that share the phenotype of hyperglycaemia. In this millennium where mankind has succeeded in deciphering the human genetic code, the issue of chronic wound management still remains an enigmatic challenge. Chronic wounds, especially non healing types, are one of the most common surgical conditions a surgeon comes across. From time immemorial Doctors have been trying many methods to treat these types of wounds. The metabolic dysregulation associated with DM causes secondary pathophysiologic changes in multiple organ systems that imposes a tremendous burden on individual.

The peculiarity of a chronic wound is that, whatever management you give, they refuse to heal, especially pressure ulcers or bed sores. The notion that wounds should be kept dry, although still held by a considerable number of clinicians, is steadily losing ground. We now know that wounds reepithelial much faster or develop granulation tissue faster when treated with dressings which allow moist wound healing. We recognize that occluding wounds does not lead to infection. Even though many modalities of wound care have come up to assist a surgeon, example the use of compression bandages to treat venous ulcers, the problem of chronic wound still remains.

A wound care revolution is currently in the making. Many techniques have been tried over the centuries to heal chronic leg ulcers. Although wound dressing have been used for at least two millennia, there exist no ideal dressing. Surgical dressing of both open and closed wound is based mainly on tradition, training and surgeons own philosophy. During the last two decades a wide variety of innovative dressings have been introduced.

Neuropathy presents in many forms including focal neuropathy and polyneuropathy and autonomous neuropathy. Patient with distal sensory neuropathy are predisposed to develop Charcot's joint which may mimic gout or degenerative joints. Treatment involves surgical debridement and antibiotic treatment. Diabetic neuropathy has been defined as peripheral nerve dysfunction after exclusion of other causes which may range from hereditary, traumatic, compressive, metabolic, toxic, nutritional infectious, immune mediated neoplastic and any other secondary systemic illness.

Classification Of Diabetic Neuropathy

A) Diffuse

1. Distal semetric sensory motor polyneuropathy

2. Autonomic neuropathy

A. submotor

B. cardiovascular C. gastro intestinal D. genitourinary

- 3. Symmetric proximal lower limb motor neuropathy. B)Focal
- 1. Cranial neuropathy
- 2. Radiculopathy
- 3. Entrapment neuropathy
- 4. Asymmetric lower limb motor neuropathy.

Non healing chronic ulcers are regularly encountered by a surgeon. The peculiarity of a these wounds are that inspite of daily dressing with expensive local applications, the wound does not heal. This problem is especially seen in diabetic ulcers, venous ulcers and pressure ulcers. Thus to treat these wound is a constant challenge for the surgeon. The notion that wounds should be kept dry, although still held by a considerable number of surgeon, is steadily losing ground. We now know that wounds develop granulation tissue when treated with dressing which allow moist wound healing. People have tried various non-conventional topical therapies in wound healing, such as aloe vera, collagen, gentian violet, benzoyl peroxide, impregnated guaze, insulin, mercurochrome, oxygen therapy, sugar and vinegar. Studies prove that topical phenytoin promotes healing of decubitus ulcer, pressure ulcer & leprosy ulcer and was found to be of superior This study was conducted to assess the percentage of graft uptake of phenytoin treated wounds and to document any local or systemic side effects of topical phenytoin applications.

II. Aim And Objectives Of The Study

To study the efficacy of topical phenytoin over conventional dressings in the management of diabetic ulcers,

- No of days required for healing
- Days required for granulation tissue formation (rate)
- Quality of graft bed & skin graft take
- Effect on bacterial load
- Side effects of topical phenytoin dressing

III. Materials And Methods

This prospective randomized comparative study included 100 patients with diabetic ulcers admitted in Madurai Medical College, Madurai from July

2016 to June 2017 satisfying all the inclusion criteria mentioned below after the clearance from the ethical committee was obtained.

The main inclusion criteria were

- grade I and II foot ulcers according to Meggitt-Wagner clinical classification
- control of diabetes mellitus with oral hypoglycemic agents or insulin

The main exclusion criteria for the study were

- o grade 111,IV,V foot ulcers according to Meggitt-Wagner clinical classification chronic ulcer of other etiology
- o other co morbid conditions like renal failure, generalized debility which adversely affect wound healing
- o Patients with allergy to phenytoin

The sample population divided into two equal and comparable groups based on willingness to undergo topical phenytoin therapy for the wound. Those not willing were subjected to conventional wound care forming the control group. Selection of patients was done by purposive sampling method. All patients underwent general physical and clinical examination for peripheral vascular status and peripheral neuropathic changes in lower extremities. Routine hematological, biochemical, urine microscopic investigations were done for each patient. The wounds were thoroughly debrided. After slough removal, the surface area was measured, tracing the outline on butter paper. This outline was transferred to a graph paper. On each occasion ulcer areas were measured twice. The average was recorded.

Sterile gauze is soaked in preparation of i.v. phenytoin and placed over the wound at 20 mg/cm2 TBSA. Conventional dressing was done with 5% w/v povidone-iodine solution. Dressings were done on twice daily basis. The patients were followed up on a daily basis for 14 days in both study and control groups. Wound culture was obtained at the start of the treatment and on the 14th day of treatment. At the end of 14 days the wounds in both the groups were inspected and compared based on the following parameters

- rate of granulation tissue formation as percentage of ulcer surface area
- quality of ulcer bed
- present dimensions and surface area of ulcer

Observed or spontaneously reported side effects (local and systemic) were documented. The patients were then subjected to split thickness skin grafting. Both the groups were given the same systemic antibiotics during the post operative period. The wounds were assessed on fifth post operative day for skin graft up take and the total number of days of hospitalization was noted. The follow up of the patients was done at one month after discharge in outpatient department to assess wound dimensions and post skin grafting complications. The results obtained were statistically evaluated and the mam parameters which were analysed were

- rate of granulation tissue formation as percentage of ulcer surface area
- graft survival and take up
- duration of hospital stay.

The variables were compared using Paired and Unpaired Student's t-test and P value of <0.05 was considered significant.

IV. Observation And Results

The 100 patients admitted for the study were divided into two equal and comparable groups. Patients subjected to topical phenytoin dressings were classified under study and those who underwent conventional moist wound dressing were classified as control.

Table 1. Age wise distribution of patients.								
Age	31-40	41-50	51-	61-	71-			
Group			60	70	80			
(yrs)								
Betadine	1	14	20	12	3			
Phenytoin	6	10	21	9	4			

 Table 1: Age wise distribution of patients.

Mean age of Betadine group is 56.12 + 8.76Mean age of Phenytoin group is 54.54 + 11.003**P** value 0.429 Not significant

Table 2: Ulcer surface area		Std.		
Group N	Mean	Deviation Median	t value	p value
ULCER AREA				
Betadine 50	37.67	7.28	2.509	0.014
Phenytoin 50	40.44	2.88		Sig

Table 3: Rate of granulation tissue formation as percentage of ulcer surface area.

 Std.

Group	Ν	Mean	Deviation	Median	t value	p value	Betadine	50	36.29
5.82	37.36	3.783	< 0.001						
Phenytoin	50	39.76	2.84	39.65	HS				

	Tuble il cluit une up us percentage of aleer sufface alea							
						Mean %of		
						graft		
	Group	N	Good	Average	Poor	uptake		
STSG	Betadine	50	35	12	3	84%		
	Phenytoin	50	42	7	1	70%		

Table 4 [.]	Graft take	un as	nercentage	of ulcer	surface	area
	Ofall take	up a	percentage	or uncer	surrace	arca

Assessment of graft take up was done at the end of the 5th post operative day as the percentage of ulcer surface area is given above.

The mean graft take up in the study group is 37.35% \pm 5.64(SD) and in the control group is 40.37% \pm 2.83(SD).

Table 5: Duration of hospital stay.

NOOF Std. Group	Ν	Mean	Deviation	Median	t value	p value	
DAYS	Betadine	50	31.80	4.63	30.00	5.706	< 0.001
Phynetoin	50	27.48	2.68	28.00		HS	

The quality of life of the patient in both the groups was assessed by the assessment of total hospital stay as number of days of admission in the hospital. The mean hospital stay in control group was 31.8 ± 4.63 (SD) days and that in the study group was 27.48 ± 2.68 (SD) days. P value is <0.001 which is highly significant

Table 6: Percentage	of negative	culture	sensitivity	at the end	of 14days.	
Creare						

Group						
	Betadine	Phynetoin	Tota			
	35	46	1			
C/S N			81			
	70.0%	92.0%	81.0%			
Р	15	4	19			
	30.0%	8.0%	18.0%			

Total	50	50	100
	100.0%	100.0%	100.0%
x2 = 4.89, p=0.027, sig			

Patients in both groups were assessed for the effect of topical phenytoin agents on the bacterial load as percentage of people who are culture sensitivity negative at the end of 14days. 90% of the study group showed negative culture sensitivity at the end of 14days whereas in control group it was 74%. In both groups, no complications occurred during the application of dressings, skin grafting or in the post operative period. The patients were followed up after one month of discharge.

The main post operative parameters noted in both the groups:

- wound size

- contracture

- pam

- infection

All these parameters were less in study as compared to control.

V. Analysis Of Data

Both groups had comparable age and sex distribution as seen in above depicted graphs.

The mean rate of granulation tissue formation in study group is 95.93cm² of total ulcer surface area and in control group is 98.09 cm["]. The results were analysed by unpaired student t-test which showed highly significant difference in rate of granulation tissue formation (p<0.0002). The mean

graft take up in the study group is 99.03 cm^2 and in the control group is 97.61 cm. The results were analysed by unpaired student t-test which showed highly significant difference in graft take up (p of 0.001). The total number of days of hospital stay for the patient was also compared. The mean number of days of hospital stay in

control group was 31.3 days and that in the study group was 27.8 days. The results were analysed by unpaired student t-test which showed highly significant difference in the number of days of hospital stay (p < 0.0002). 45% of the study group showed negative culture sensitivity at the end of 14days whereas in control group it was 37%.

VI. Discussion

This study was done as a prospective randomized controlled comparative study to compare the efficacy of topical phenytoin dressing to conventional wound dressing in the management of diabetic ulcer.

 Table 7: Comparision of present study to study by Muthu kumaraswamy et al shows following similarities.

		D t l'		D (11
	Phenytoin	Betadine	Phenytoin	Betadine
	564	58.7	54 54	56.12
	50.4	50.7	54.54	50.12
Mean age in				
-				
Years				
Graft uptake in	72.4%	58.4%	84	74
_				
Percentage				
6				
Hospital stay in	21	45	27	32
Dovs				
Days				
	<u> </u>			-
Negative bacterial	82%	54%	92%	70%
culture				
outture				

Muthukumara Swamy et al Present Study

Mean age group in Muthu kumaraswamy et al study in study group is 56.4 yrs and in the control group is 58.7yr while in the present study it is 54.54yrs in study group and 56.12yrs in control group. The percentage of graft take up in Muthu kumaraswamy et al study in the study group is 72.4% and in the control group is 58.4% while the percentage of graft take up in the study group is 84% and in the control group is 74% The duration of hospital stay in Muthu kumaraswamy et al study in control group was 45days and that in the study group was 21 days while in the present study the mean hospital stay in control group was 32days and that in the study group was 27days. The negative wound culture sensitivity at the end of 14days in Muthu kumaraswamy et al study was 82% in study group and 54% in control group. Whereas in the present study the negative culture sensitivity at the end of 14days was 92% of the study group and in control group it was 70%.

Important difference between present study and Muthu kumaraswamy et al study in that in the latter a thin layer of phenytoin powder is laid over the wound and covered with a dry gauge as the method of application.

VII. Limitations Of The Study

The most important limitation of the present study is its sample size. A randomised controlled comparative study with a much larger population may help to further substantiate the findings or reveal variations which were not observed in the present study. The cost burden on the patient is also not analysed in this study as it can be influenced by various factors other than the cost of dressings. Phenytoin dressing was found to be less expensive compared to conventional moist dressing. However, no commercial preparation of phenytoin is available in the market so far.

Summary

Increased rate of granulation tissue formation was seen in topical phenytoin group when compared to conventional group.

- Better graft take up was seen in topical phenytoin group when compared to conventional group.
- Considerable effect on bacterial load was seen in topical phenytoin group when compared to conventional group.
- Shorter duration of hospital stay was seen in topical phenytoin group when compared to conventional group.
- Topical phenytoin dressing thus is an effective, inexpensive and widely available therapeutic agent in wound healing.
- Follow up observations revealed that topical phenytoin dressing group suffered lesser post skin grafting complications like wound contracture, residual raw area and pain compared to conventional group.

VIII. Conclusion

In the present study it was concluded that topical phenytoin by decreasing bacterial load, forming healthy granulation tissue helps in better graft take up than the conventional dressing. Because of enhanced healing and overall hospital stay, the post operative complications were less in topical phenytoin dressing group. Thus topical phenytoin moist wound dressing can be considered as superior option in management of diabetic ulcers. But further studies with larger population will be needed in the future before topical phenytoin dressing can be added to the wide spectrum of treatment modalities available in the management of diabetic ulcers and ulcers of other etiologies.

References

- [1]. Muthu Kumaraswamy MG, Sivakumar G, Monoharan G. Topical phenytoin in diabetic foot ulcers. Diabetes Care. 1991; 14(10): 909-11.
- [2]. Merritt HH, Putnam TJ: Sodium diphenyl hydantoinate in the treatment of convulsive disorders. JAMA 1938; 111: 1068-73.
- [3]. Silverman AK, Fairley J, Wongs RC. Cutaneous and Immunologic reactions to phenytoin. J.Am. Acad Dermatol. 1988;18:721-41.
- [4]. ASHP Drug information 2001, American Society of Health System
- [5]. Pharmacists Bethedsa MD: 2001-81.
- [6]. Kimball OP, Horan TN. The use of Dilantin in the treatment of epilepsy. Ann
- [7]. Intern Med 1939;13:787-93.
- [8]. Shapiro M. Acceleration of gingival wound healing in non-epileptic patients receiving diphenylhydantoin sodium. Exp Med Surg 1958;16:41-53.
- [9]. Anstead GM, Hart LM, Sunahara JF. Phenytoin in wound healing Ann
- [10]. Pharmacother 1996;30:768-75.

- Talas G, Brown RA, McGrouther A Role of phenytoin in Wound healing a wound pharmacology perspective Biochem [11]. Pharmacol 1999:57(10): 1085-94.
- Genever PG, Cunliffe WJ, Wood EJ. Influence of the extracellular matrix on fibroblast responsiveness to phenytoin using [12]. in vitro wound healing model. Br J of Dermatol 1996;133:231-5.
- [13]. Dill RE, Miller EK, Weil T, Lesley S, Farmer GR, La Copino AM. Phenytoin increase gene expression for platelet derived growth factor in chain in macrophages and monocytes. J. peridontol 1993;64:169-73.
- [14]. Madagheghas, Salchian B, Tavassoli M. Use of phenytoin in healing of war and non war wounds. Int J. Dermatol 1980;28(5):347-50.
- [15]. El. Layat SG. Preliminary experience with lopical phenytoin in wound healing in a war zone Mil Med 1989;154(4):178-80.
- [16]. Lodha SC. New application of an old drug -topical phenytoin for burns, J.Burns care Rehab 1991;12(1):96.
- Yadav JK, Singhvi AM, Kumar N, Garg S. Topical phenytoin in the treatment of split-thickness skin autograft donor [17]. sites: a comparative study with polyurethane membrane drape and conventional Bums. 1993 dressing. Aug;19(4):306-10.
- [18]. Lodha SC, Lohiya MI, Vyas Mc. Role of phenytoin in healing of large abscess cavities, Br. J. surg. 1991;78:105-8.
- Pai MR, Sitaram N, Kotian MS. Topical phenytoin in diabeticulers : a double blind controlled trial. Indian J. Med Sci [19]. 2001;55(11):593-9.
- [20]. El-Nahas M, Gawish H, Tarshoby M, State O. The impact of topical phenytoin on recalcitrant neuropathic diabetic foot ulceration. J Wound Care. 2009 Jan 8(1):33-7.
- [21]. Bansal NK, Mukul. Comparison of topical phenyton with normal saline in the treatment of chronic trophic ulcers in leprosy Int J.Dermatol.1993;34(3):210-3.
- [22]. Menezes J, Rajendran A, Jacob AJ and Vaz M. The use of typical phenytoin as an adjuant in the treatment of trophic ulcers. South Cost Asia J. Trop. Med Public Health 1993;21(2):340-2.
- [23]. Bhatia A, Nanda S, Gupta U, Gupta S, Reddy BS. Topical phenytoin suspension and normal saline in the treatment of leprosy trophic ulcers. J Dermatolog Treat. 2004 Sep;15(5):321-7.
- [24].
- Pendse AK, Sharma A, Sodami A, Hada S. Topical phenytoin in wound healing. Int. Dermato 1993;32:214-7. Oluwatosin OM, Olabanji JK, Oluwatosin OA, Tijani LA, Onyechi HU. A comparison of topical honey and [25]. phenytoin in the treatment of chronic leg ulcers. Afr Med Med Sci. 2000 Mar;29(1):31-4
- [26]. Carneiro PM, Nyawawa ET. Topical phenytoin versus EUSOL in the treatment of non-malignant chronic leg ulcers. East Afr Med J. 2003 Mar;80(3): 124-9.
- Rhodes RS, Heyneman CA, Culbersten VL, Wilson SE, Phatak HM. Topical Phenytoin Treatment of stage II decubitus [27]. ulcers in the elederly. Ann Pharmaco therapy 2001;35:675-81.
- [28]. Subbanna PK, Margaret FX, George J, Tharion G, Neelakantan N, Durai Set al Topical phenytoin solution for treating pressure ulcers. Spinal Cord. 2007 Nov;45(11):739-43.
- [29]. Masgran - Peya E, Lacour M, Salman D. Topical phenytoin accelerates healing in epidermolysis bullosa simplex. Dermatology 1995; 190: 254.

*Dr.A.M.Syed Ibrahim. "A Study of Efficacy of Topical Phenytoin in the Management of Diabetic Ulcer." IOSR Journal of Dental and Medical Sciences (IOSR-JDMS) 16.8 (2017): 05-11.