

## Original Research Article

# Microbiologic Profile of Diabetic Foot Ulcers: A Trial of Honey versus Povidone Iodine Dressings

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### ABSTRACT

Diabetic foot ulcers (DFU), a debilitating complication of diabetes mellitus, are often infected. Infections impede wound healing and can lead to ulcer progression and possible mortality if poorly managed. Wound dressing is vital to DFU management, and supports the prevention and treatment of wound infections.

**Aims:** To compare the effect of honey and povidone iodine dressings on wound microbial colonization and infection for Wagner Grade 2 DFU.

**Study design:** This was a randomized controlled trial on the effects of honey and povidone iodine dressings on the microbiologic profile of Wagner grade 2 DFU at the University of Port Harcourt Teaching Hospital (UPTH), Port Harcourt, spanning a year interval.

**Methodology:** Thirty patients (13 females) aged 47 to 65 years with Wagner grade 2 diabetic foot ulcers were enrolled. Data on socio-demographics, and contaminating/infecting organisms were obtained from serial swab sample collection and microbiological investigation and analysed using Statistical Package for the Social Sciences (SPSS) 20.0. A p-value < 0.05 was considered significant.

**Results:** The mean HbA1c levels were  $7.52 \pm 1.023\%$  and  $7.40 \pm 0.944\%$  for the honey and povidone iodine groups respectively ( $P = 0.73$ ). Staphylococcus aureus, Escherichia coli, Pseudomonas, Klebsiella, Proteus spp. were all cultured at baseline studies. S. aureus was the most prevalent throughout the study duration. By week 4, none of the patients in the honey group had organisms isolated. By week 5 and 6, no microbial organism was isolated from the patients in both groups.

**Conclusion:** Polymicrobial ulcer contamination occurs commonly in DFU. Optimal wound care controls microbial activity, thereby promoting wound healing.

*Keywords:* Diabetic foot ulcer, wound dressing, povidone iodine, honey, microbiologic profile

## 1. INTRODUCTION

Foot ulcers are debilitating complications of diabetes mellitus, a disorder of abnormal glucose metabolism.[1] The prevalence of DM foot lesions is 0.9% to 8.3% in Nigeria.[2][3][4] About a sixth of diabetics develop diabetic foot ulcers (DFU) in their lifetime, with associated physical, psychological, and financial disability.[2][5][6] Over 80% of non-traumatic lower limb amputations are also attributable to DFU. [2][7] According to a 2004 analysis, chronic wounds cost about \$9.7 billion in the US annually, making them the most expensive human skin condition in terms of direct medical costs.[8]

Neuropathy, angiopathy, trauma and impaired immunity act synergistically to produce ulcers and further impede the healing of these wounds [1][4][5] Excessive mechanical stress on the wound, tissue ischemia, or interstitial oedema usually contribute to impair the healing of DFU [9] Autonomic dysfunction causes anhidrosis, dry and easily-cracked skin that permits bacterial invasion. [5] Advanced glycation end-products bind to tissues, initiate florid atherosclerosis, delay collateral vessel formation, cause thrombosis, and indirectly hinder the local antimicrobial mechanism, while impaired bacterial phagocytosis and cell-mediated immunity directly produce immunosuppression, infection. [2][4][5] Long-standing bacterial colonisation and infection by diverse pathogens are common complications of healing wounds. [9]

The Wagner Classification is a commonly used DFU grading system. [1][10] It considers the ulcer depth, occurrence of osteomyelitis, and the extent of tissue gangrene. [10] Wagner grade 2 ulcers are deep, with tendon, bone, ligament, or joint involvement. [1][10] Clinically, the presence of two or more of the cardinal symptoms of inflammation—induration, erythema, elevated temperature, increased pain, and purulent discharge—indicates the presence of DFI. [6]

DFU infections are often polymicrobial. [2][5][11][12][13] *Streptococcus sp.*, *Bacteroides*, *Pseudomonas*, *Proteus*, *Enterococcus*, *Escherichia coli*, and *Acinobacter* have been isolated severally, with *S. aureus* being the most common. [14][15] Culture samples are best taken from deep tissues beneath the ulcers. [16] Culture-based approaches to microbial isolation is being replaced by highly sensitive molecular sequencing which targets the species-specific small subunit ribosomal RNA (16S rRNA) gene, and is able to characterize the wound microbiota based on microbial load, microbial diversity, and presence of pathogens. [12][17] However, this is not commonly available in developing countries. Studies have found quinolones, cephalosporins, gentamicin combined with metronidazole to be the most potent antibiotics against both gram-positive and gram-negative organisms in DFU and thus recommended them for empirical use. [11][18][19]

Wound healing is not necessarily impaired by bacterial colonization. [12][20] The administration of antibiotic based only on the isolation of microbe from the base of the wound is inappropriate and risks the development of multi-drug resistant microbial strains. [9][12] The clinical picture should be considered also.

The cornerstone of DFU treatment is multidisciplinary care with a focus on patient education, regular foot assessment, glycaemic control, and aggressive intervention (debridement, antibiotic therapy, and regular dressing). [2][8][21] Wound dressings should relieve symptoms, provide mechanical and antimicrobial protection, absorb exudates, control odour, be cost-effective and ultimately promote healing. [5][9][22][23] The choice of dressing is based on the wound features i.e. appearance and exudate. [22] There is however no model dressing for DFU. [8][22]

Honey is a supersaturated sugar solution prepared by bees, from nectar or other plant secretions. [24][25][26] The earliest medical literature gives evidence of honey's therapeutic benefits. [9] It contains carbohydrates - usually glucose and fructose (80–85%), water (15–17%), protein (0.1–0.4%), ash (0.2%), and small amounts of enzymes, amino acids, vitamins, and phenolic antioxidants. [25][26][27] It also contains hydrogen peroxide, and inhibin, and has high acidity (pH 3.2 - 4.2) which all impede bacterial growth. [14][23][28] Honey prevents biofilm development, reduces the ulcer's bioburden by debriding necrotic tissues, reduces inflammation and pain, and promotes granulation and epithelisation while minimizing scarring. [9][14][23][29] It deodorizes infected wounds by providing local bacteria with glucose and fructose as substitutes for amino acids from dead cells and serum, leading to the production of lactic acid instead of malodorous substances such as amines, ammonia, and sulphur. [30] Studies show that honey produced faster healing and wound size reduction than other conventional dressings. [9] Honey is potent against *Staphylococcus aureus* (*S. aureus*), *Pseudomonas aeruginosa*, *Klebsiella pneumoniae*, *Escherichia coli*, methicillin-resistant *S. aureus*, etc. [14][31] It is cheap, thus reducing patients' financial burden. [4][14][19]

Povidone-iodine contains a loose blend of iodine and a non-ionic surfactant. [32][33][34] It easily penetrates cell membranes, and interrupts microbial protein synthesis, respiratory chain enzymes, lipid membrane, and nucleic acid activity. [33][34] Its potency is greatest at 0.1%–1% dilution due to weakened bonds between the carrier polymer and iodine molecules. This increases the quantity of free elemental iodine in solution which is lethal to bacteria, viruses, protozoa, fungi, amoebic cysts, and protozoa. [3][14] It eradicates all common nosocomial infection-causing microbes in under 20–30 seconds. [34] Povidone-iodine is also effective against bacterial biofilm, has a good safety profile and is widely tolerated. [34][35] Despite its extensive use, resistance or antibiotic cross-resistance to it is yet unreported since it acts on multiple bacterial targets. [34][35]

In an established infection, bacteria impede healing by depriving host cells of nutrients and oxygen, produce cytotoxic substances, and inhibit immune responses, leading to local hypoxia, thrombosis, tissue death, and ulcer deterioration. [12][20] Therefore, an understanding of practical, affordable, and accessible dressing materials that control infection would significantly improve DFU management. [9][14] The aim of the present study was to compare the effect of honey and povidone iodine dressings on wound microbial colonization and infection for Wagner grade 2 DFU.

## 2. MATERIAL AND METHODS

### 2.1 Study Design

This randomized controlled study compared the microbiologic profiles of organisms contaminating healing Wagner Grade 2 DFU in patients receiving povidone iodine dressings versus honey dressings in UPTH between April 1st, 2017, and April 30th, 2018.

### 2.2 Sample Size Determination

The sample size for the study was calculated using the formula for comparison of groups [36];

$$n = \frac{2 (Z\alpha + Z\beta)^2 S^2}{d^2}$$

where  $n$  = minimum sample size;  $Z\alpha$  = significance level of 95%; corresponds to a value of 1.96;  $Z\beta$  = power of 80%; corresponds to a value of 0.84;  $S$  = standard deviation; standard deviation of the rate of healing among patients with diabetic foot ulcers using honey dressing from a similar study was 0.94[17];  $d$  = level of precision of 0.5.

Allowing for an attrition of about 10%, the sample size was rounded up to 60.

Adjustment for population <10,000 using finite population correction[36]

Adjusted sample size =  $\frac{n_0 N}{n_0 + (N-1)}$

$n_0$  = minimum sample size = 60;  $N$  = Total population of DFU from the review of records (UPTH, 2016) = 47

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Therefore, the adjusted sample size was approximated to 30

Hence, a total sample size of 30 comprising 15 patients per group was involved in the study.

### 2.3 Eligibility Criteria

The study participants were diabetics with Wagner Grade 2 foot ulcers, aged 30-65 years (lesser risk of co-morbidities such as cardiac and vascular diseases compared to older patients). But it is important to note that presence of Vasculopathy and Neuropathy. These co morbid conditions are detrimental of the outcome of the treatment.

#### 2.3.1 Inclusion Criteria

They also fulfilled the following criteria:

- Ankle brachial pressure index (ABPI) >0.9,
- Serum albumin concentration >35g/dl.
- Oxygen saturation of  $\geq 92\%$  by pulse oximetry

#### 2.3.2 Exclusion Criteria

Patients with multiple co-morbidities, severe immunosuppression, malignancy or chemotherapy, haemoglobinopathies, steroid therapy, and neutrophil count under  $2000/\text{mm}^3$ .

### 2.4 Study Procedure

#### 2.4.1 Randomization

The subjects were allocated into groups by simple randomization using an opaque envelope containing papers labelled "A" or "B". A paper was randomly selected from the opaque envelope for each of the eligible subjects, who were then assigned to the group labelled on the paper. Group A received honey dressing while Group B received povidone iodine dressing.

## 2.4.2 Blinding

The investigator was blinded (single blinding) to the dressing options to avoid bias and ensure the validity of the outcome measures. The investigator was also absent at the removal of old dressings, returned to assess the wound/foot parameters, and left before the application of new dressings by trained nurses. But each group should be marked to avoid confusion.

## 2.4.3 Details of the Study

Honey procured from a single commercial local source to guarantee uniformity and 10% povidone iodine solution were used. Honey should be natural because artificial honey also available. And the efficacy will vary. All subjects received suitable antibiotics and their ulcers were surgically debrided by the investigator or senior postgraduate orthopaedic trainees. Tissue specimens excised from the ulcer bed were instantly sent for microscopy, culture and sensitivity analysis. Optimum glycaemic control was maintained under the supervision of a physician.

Dressing was performed daily. The wound was first cleansed with 0.9% saline, covered with honey or povidone iodine-soaked gauze supported by dry sterile gauze, and then bandaged.

A weekly wound assessment was performed by the investigator who was blinded to the material (honey or povidone iodine) of the dressing. A wound swab was also taken from the ulcer bed and sent for analysis by a physician medical microbiologist. The assessment ended 6 weeks after the initial debridement or when the wound had healed, whichever occurred first.

The consumables used were honey, 10% povidone iodine (Betadine®), 0.9% saline, sterile cotton swabs, sterile gauze, crepe bandages, sterile swab sticks, sterile gloves and culture media (sheep blood agar, MacConkey agar, and Robertson cooked meat medium).

## STUDY PROTOCOL

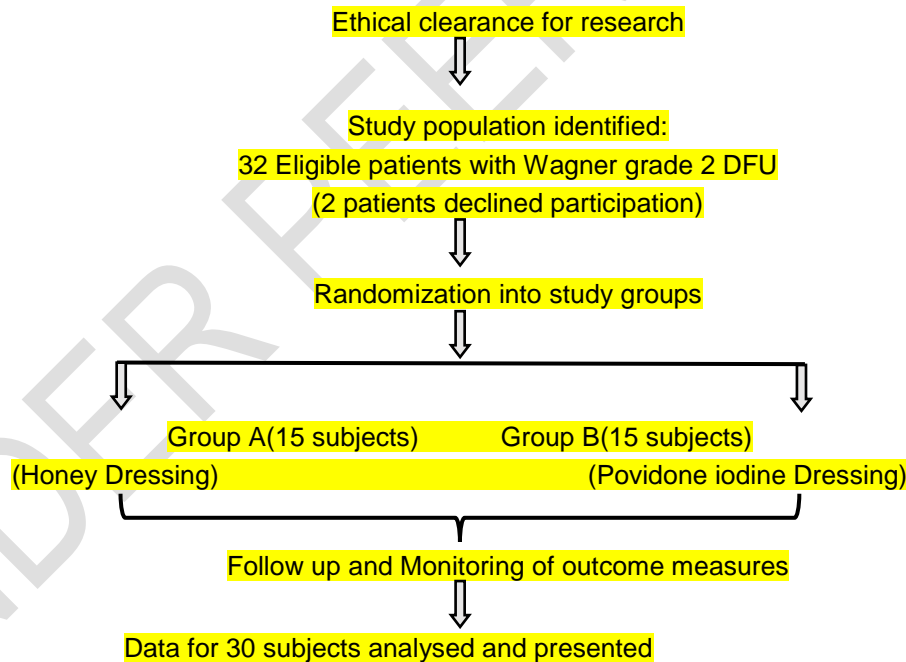


Figure 1: Diagrammatic Presentation of Study Protocol

## 2.5 Data Analysis

Data analysis utilized the IBM® Statistical Package for the Social Sciences (SPSS) version 20. Data were presented as tables and charts. Qualitative variables such as age categories were expressed as frequencies and proportions while quantitative variables such as HbA1c were summarized as means  $\pm$  standard deviation. Student's t test was used to

compare the differences in means between the groups for data with normal distribution (such as HbA1c). Chi square test or Fisher's exact test was used to compare the differences in proportions between the groups. A  $P$  value = 0.05 was considered statistically significant.

### 3. RESULTS

The study had 17 males and 13 females, aged 47-65 years with a mean  $55.53 \pm 5.041$  years and  $54.93 \pm 5.298$  years in the honey and povidone iodine groups respectively. There was no significant difference in age ( $P = 0.26$ ) and sex ( $P = 0.71$ ) between both study groups as depicted in Table 1.

**Table 1: Demography of study groups**

| Variables           | Groups in the study                      |        |       |        | Total |        |
|---------------------|--|--------|-------|--------|-------|--------|
|                     | Povidone Iodine                          |        | Honey |        | N=30  | n (%)  |
|                     | N=15                                     | n (%)  | N=15  | n (%)  |       |        |
| <b>Age category</b> |  |        |       |        |       |        |
| 45-49 years         | 3  | (20.0) | 4     | (26.7) | 7     | (23.3) |
| 50-54 years         | 4  | (26.7) | 0     | (0.0)  | 4     | (13.3) |
| 55-59 years         | 5  | (33.3) | 7     | (46.7) | 12    | (40.0) |
| >60 years           | 3  | (20.0) | 4     | (26.7) | 7     | (23.3) |
|                     | <b>Fisher's exact test=4.432; P=0.26</b> |        |       |        |       |        |
| <b>Sex</b>          |  |        |       |        |       |        |
| Female              | 7  | (46.7) | 6     | (40.0) | 13    | (43.3) |
| Male                | 8  | (53.3) | 9     | (60.0) | 17    | (56.7) |
|                     | <b>Chi-square=0.136; P=0.71</b>          |        |       |        |       |        |

The HbA1c range was 6.3% - 10.2% in the povidone iodine group, and 6.7%-9.9% for the honey group. The mean HbA1c was  $7.52 \pm 1.023\%$  and  $7.40 \pm 0.944\%$  for the honey and povidone iodine groups respectively. This difference was not significant ( $P = 0.73$ ).

*Staphylococcus aureus* was the most prevalent organism isolated (63.3%) in the initial wound swab. Diverse organisms were cultured from three ulcers (10%) at the initial swab. In all the weeks of follow-up, *S. aureus* remained the most prevalent organism as shown in Table 2.

**Table 2: Distribution of microbiological organisms identified across the study period**

| Variables          | Groups in the study    |                                  | Total<br>N=30<br>n (%) |
|--------------------|------------------------|----------------------------------|------------------------|
|                    | Honey<br>N=15<br>n (%) | Povidone-Iodine<br>N=15<br>n (%) |                        |
| <b>Baseline*</b>   |                        |                                  |                        |
| <i>S.aureus</i>    | 8 (53.3)               | 11 (73.3)                        | 19 (63.3)              |
| <i>E.coli</i>      | 3 (20.0)               | 2 (13.3)                         | 5 (16.6)               |
| <i>Pseudomonas</i> | 2 (13.3)               | 1 (6.7)                          | 3 (10)                 |
| <i>Klebsiella</i>  | 2 (13.3)               | 0 (0.0)                          | 2 (6.7)                |
| <i>Proteus</i>     | 2 (13.3)               | 0 (0.0)                          | 2 (6.7)                |
| <b>Week 1*</b>     |                        |                                  |                        |
| <i>S.aureus</i>    | 11 (73.3)              | 11 (73.3)                        | 22 (73.3)              |
| <i>E.coli</i>      | 2 (13.3)               | 3 (20.0)                         | 5 (16.7)               |
| <i>Klebsiella</i>  | 1 (6.7)                | 0 (0.0)                          | 1 (3.3)                |
| <i>Proteus</i>     | 1 (6.7)                | 0 (0.0)                          | 1 (3.3)                |
| <b>Week 2</b>      |                        |                                  |                        |
| <i>S.aureus</i>    | 10 (66.7)              | 9 (60.0)                         | 19 (63.3)              |
| <i>Pseudomonas</i> | 1 (6.7)                | 1 (6.7)                          | 2 (6.7)                |

**Week 3**

*S.aureus* 7 (46.7) 7 (46.7) 14 (46.7)

**Week 4**

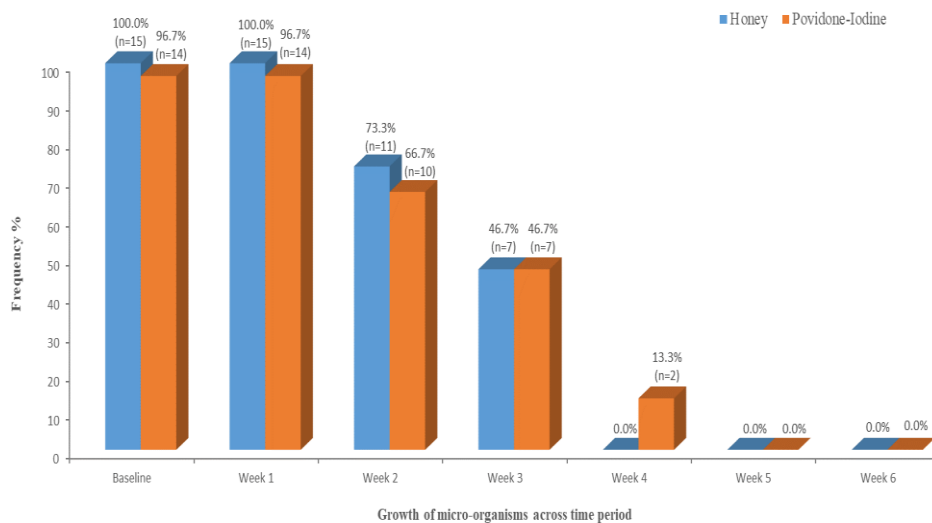
*S.aureus* 0 (0.0) 2 (13.3) 2 (6.7)

\*Some of the DFUs in the groups had more than one micro-organism isolated.

Weeks 5 and 6 reported no growth of micro-organisms

At baseline, microorganisms were isolated from all patients in group A and all **except one** (96.7%) patient in group B. By week 4, none of the patients in the honey group (0.0%) had organisms isolated while two of the patients in the povidone iodine group (13.3%) had organisms isolated. By weeks 5 and 6, no microbial organism was isolated from the patients in both groups as **shown** in Figure 2.

**Figure 2: Presence of growth of micro-organisms across time**



Before hospital presentation, the wound care product most commonly used was antibiotic powder (23.3%). **Multiple products were used by 13.3% of the patients** while 10% used none. The difference between both study groups was not significant ( $P = 0.66$ ). See Table 3.

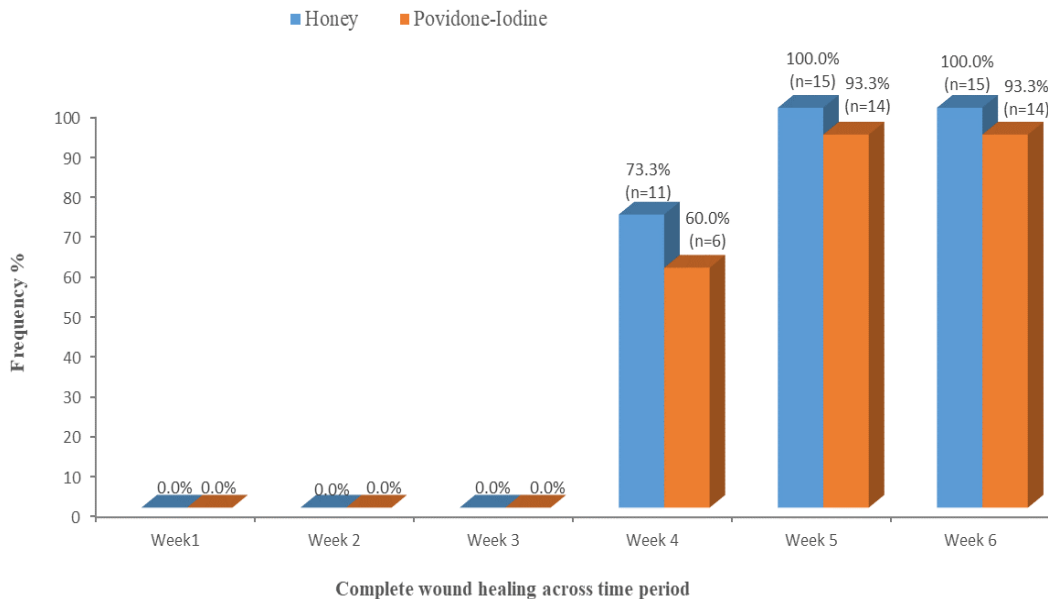
**Table 3: Distribution of wound care products used by patients before hospital presentation**

| Products used     | Groups in the study |                       | Total n (%) |
|-------------------|---------------------|-----------------------|-------------|
|                   | Honey n (%)         | Povidone-Iodine n (%) |             |
| Iodine            | 1 (6.7)             | 3 (20.0)              | 4 (13.3)    |
| Cetrimide         | 1 (6.7)             | 3 (20.0)              | 4 (13.3)    |
| Gentian violet    | 2 (13.3)            | 1 (6.7)               | 3 (10.0)    |
| Eusol             | 2 (13.3)            | 0 (0.0)               | 2 (6.7)     |
| Hydrogen peroxide | 2 (13.3)            | 1 (6.7)               | 3 (10.0)    |
| Antibiotic powder | 3 (20.0)            | 4 (26.7)              | 7 (23.3)    |
| Combined products | 3 (20.0)            | 1 (6.7)               | 4 (13.3)    |
| Nil               | 1 (6.7)             | 2 (13.3)              | 3 (10.0)    |

|              |                   |                   |                   |
|--------------|-------------------|-------------------|-------------------|
| <b>Total</b> | <b>15 (100.0)</b> | <b>15 (100.0)</b> | <b>30 (100.0)</b> |
|--------------|-------------------|-------------------|-------------------|

Fisher's exact test=5.846; **P=0.66**

**Figure 3: Distribution of subjects with complete wound healing across the study period**



#### 4. DISCUSSION

The difficulty in finding patients who met the inclusion criteria - those whose wounds were supposed to heal as close to normal as possible, highlights the burden of DM complications, especially DFU. This indicates, as some writers have suggested, that most diabetics may already be prone to wound sepsis and other complications at their first presentation with ulcers. [1][3][21]

**Duration of Diabetes, Diabetes status and response to management of diabetes are key factors to be considered while comparing the results of the study.**

Both the honey and povidone iodine groups were comparable in terms of demographic characteristics, and HbA1c level, thus ruling them out as possible confounders. DFU chronicity and risk of infection correlate positively with poor glycaemic control as observed in other studies. [37][38][39][40] Ignorance and socio-cultural influences such as belief in the supernatural causes and therapies for DFU contribute to late presentation as noted by Ogbera et al., in whose study 78% of respondents held such views. [41][42] These erroneous attitude and behaviour towards DFU are still fairly common observed among patients in resource-limited countries such as the authors' practice environment, and possibly contribute significantly to poor patient outcome.

*Staphylococcus aureus* was the most prevalent organism isolated (63.3%) in the initial wound swab, possibly from self-contamination from patients' peri-ulcer normal flora, while one ulcer (3.3%) yielded no growth probably because the patient had self-administered antibiotics prior the sample collection. Several workers predominantly cultured *S.aureus* from DFUs in their studies. [4][6][14][15][18][43] In line with the findings of other workers, [4][14][15][18] 3 ulcers (10%) contained polymicrobial organisms at the first swab. In contrast, Shukrimiet al [14] in a similar prospective comparative study in Malaysia, isolated more *Streptococcus sp.* (30%) than *Staphylococcus sp.* (16%), with 20% of the ulcers harboring polymicrobial infections, Mehta et al [13] isolated *Pseudomonas aeruginosa* most commonly (27%) in their study while Mohammed et al reported a predominance of monomicrobial infections (77.3%) and *Candida albicans* in Egypt for unclear reasons [44]. This could be due to the differences in organisms that are prevalent in the different environments. Similarly, the majority of the ulcers yielded no growth towards the point of complete wound healing. Holubova et al

surmised that antibiotic therapy was unnecessary because honey eradicated all, including the virulent microorganisms in ulcers with local signs of infection.[9]

Foot ulcers that were previously treated with cetrimide, iodine, hydrogen peroxide, or those that had no prior treatment before debridement in the honey group had an average healing time of  $4.00 \pm 0.00$  weeks. Accordingly, ulcers in the povidone iodine group that had previously received hydrogen peroxide, iodine, and gentian violet treatment healed in about  $5.00 \pm 0.00$  weeks. There was no significant difference in wound healing between both study groups based on the individual wound care products used prior to this study. These products also did not appear to have influenced the nature of the colonizing microbiota in the course of the study. This is probably because before debridement, the ulcer beds contained varying amounts of slough, scab, and necrotic tissue that were excised giving rise to fresh ulcer beds that were not exposed to the applied substances. Furthermore, healing becomes progressive after debridement. [2][3][31]

This study's strength is in its design as a randomized controlled trial, which aims to offer significant information regarding the impact of honey versus povidone iodine on the microbiologic profiles of DFU. Nevertheless, because this was a single-center study, its generalizability might be limited; as a result, multi-center investigations are advised.

## 5. CONCLUSION

Microorganisms, most commonly *Staphylococcus aureus* readily colonize DFU at different stages of their healing, with multiple species often co-existing within the ulcer. Both honey and povidone iodine were comparable in controlling the wound microbiota significantly enough to prevent wound infection and allow for healing to progress. The mere presence of organisms in an ulcer does not necessarily impede its healing.

**Aim of the study has not been explained in conclusion. There is no statement regarding the method of dressing and its superiority over other one. What is the effect of different dressings on the Microorganisms flora?**

## CONSENT AND ETHICAL APPROVAL

Ethical approval was obtained from the research and ethics committee of the UPTH before the commencement of the study. Written informed consent was obtained from all study participants after being given adequate information on the nature, scope, and reason for the research. Anonymity and confidentiality were upheld in the study. Participation in the study was voluntary, and patients' withdrawal from the study did not affect their medical care.

## REFERENCES

1. Akkus G, Sert M. Diabetic foot ulcers: A devastating complication of diabetes mellitus continues non-stop in spite of new medical treatment modalities. *World J Diabetes*. 2022;13(12):1106–21.
2. Ibrahim AM. Diabetic foot ulcer: synopsis of the epidemiology and pathophysiology. *Int J Diabetes Endocrinol*. 2018;3(2):23-28.
3. Danmusa UM, Terhile I, Nasir IA, Ahmad AA, Muhammad HY. Prevalence and healthcare costs associated with the management of diabetic foot ulcer in patients attending Ahmadu Bello University Teaching Hospital, Nigeria. *Int J Health Sci (Qassim)*. 2016;10(2):219-28.
4. Akaa PD, Ahachi NC, Kortor NJ, Mue DD, Elachi CI, Ogiator M, et al. Diabetic foot ulcers: epidemiology, management modalities and outcome at Benue State University Teaching Hospital Makurdi. *J Adv Med Med Res*. 2017;22(10):1–12.
5. Raja JM, Maturana MA, Kayali S, Khouzam A, Efeovbokhan N. Diabetic foot ulcer: A comprehensive review of pathophysiology and management modalities. *World J Clin cases*. 2023;11(8):1684–93.
6. Alhubail A, Sewify M, Messenger G, Masoetsa R, Hussain I, Nair S, et al. Microbiological profile of diabetic foot ulcers in Kuwait. *PLoS One*. 2020;15(12):e0244306.
7. Edo AE, Edo GO, Ezeani IU. Risk factors, ulcer grade and management outcome of diabetic foot ulcers in a Tropical Tertiary Care Hospital. *Niger Med J*. 2013;54(1):59–63.
8. Powers JG, Higham C, Broussard K, Phillips TJ. Wound healing and treating wounds: Chronic wound care and management. *J Am Acad Dermatol*. 2016;74(4):607–25.
9. Holubová A, Chlupáčová L, Krocová J, Cetlová L, Peters LJF, Cremers NAJ, et al. The Use of Medical Grade Honey on Infected Chronic Diabetic Foot Ulcers—A Prospective Case-Control Study. *Antibiotics (Basel)*. 2023;12(9):1364.

10. Jeon BJ, Choi HJ, Kang JS, Tak MS, Park ES. Comparison of five systems of classification of diabetic foot ulcers and predictive factors for amputation. *Int Wound J.* 2017;14(3):537–45.
11. Otu AA, Umoh VA, Essien OE, Enang OE, Okpa HO, Mbu PN. Profile, Bacteriology, and Risk Factors for Foot Ulcers among Diabetics in a Tertiary Hospital in Calabar , Nigeria. *Ulcers.* 2013;2013:Article ID 820468.
12. Tomic-Canic M, Burgess JL, O'Neill KE, Strbo N, Pastar I. Skin Microbiota and its Interplay with Wound Healing. *Am J Clin Dermatol.* 2020;21(Suppl 1):36–43.
13. Mehta VJ, Kikani KM, Mehta SJ. Microbiological profile of diabetic foot ulcers and its antibiotic susceptibility pattern in a teaching hospital, Gujarat. *Int J Basic Clin Pharmacol.* 2014;3(1):92–5.
14. Shukrimi A, Sulaiman AR, Halim AY, Azril A. A comparative study between honey and povidone iodine as dressing solution for Wagner type II diabetic foot ulcers. *Med J Malaysia.* 2008;63(1):44–6.
15. Kaimkhani GM, Siddiqui AA, Rasheed N, Rajput MI, Kumar J, Khan MH, et al. Pattern of infecting microorganisms and their susceptibility to antimicrobial drugs in patients with diabetic foot infections in a tertiary care hospital in Karachi, Pakistan. *Cureus.* 2018;10(6): e2872. doi:10.7759/cureus.2872.
16. Ramsay S, Cowan L, Davidson JM, Nanney L, Schultz G. Wound samples: moving towards a standardised method of collection and analysis. *Int Wound J.* 2016;13(5):880–91.
17. Moghazy AM, Shams ME, Adly OA, Abbas AH, El-Badawy MA, Elsakka DM, et al. The clinical and cost effectiveness of bee honey dressing in the treatment of diabetic foot ulcers. *Diabetes Res Clin Pract [Internet].* 2010;89(3):276–81.
18. Oyan B, Abere S, Gomba VE, Lawson S, Okoli N, Tonye-Abere O. Diversity of microbiological pathogens in diabetic foot ulcers in an African population. *Int J Res Med Sci.* 2023;11(9):3212-7.
19. Brenyah RC, Ephraim RKD, Jnr BAE, Asamoah J. Bacterial profile of diabetic foot ulcers of patients visiting a specialist diabetic clinic at Komfo Anokye Teaching Hospital, Kumasi, Ghana. *Br J Med Med Res.* 2014;4(27):4501–10.
20. Versey Z, da Cruz Nizer WS, Russell E, Zigic S, DeZeeuw KG, Marek JE, et al. Biofilm-Innate Immune Interface: Contribution to Chronic Wound Formation. *Front Immunol.* 2021;12:648554. doi:10.3389/fimmu.2021.648554
21. Kolarić V, Svirčević V, Bijuk R, Zupančić V. CHRONIC COMPLICATIONS OF DIABETES AND QUALITY OF LIFE. *Acta Clin Croat.* 2022 Nov;61(3):520–7.
22. Shi C, Wang C, Liu H, Li Q, Li R, Zhang Y, et al. Selection of Appropriate Wound Dressing for Various Wounds. *Front Bioeng Biotechnol.* 2020;8:182. doi:10.3389/fbioe.2020.00182
23. Wang C, Guo M, Zhang N, Wang G. Effectiveness of honey dressing in the treatment of diabetic foot ulcers: A systematic review and meta-analysis. *Complement Ther Clin Pract [Internet].* 2019;34:123–31.
24. Codex Alimentarius Commission. Codex Alimentarius Commission Standards. Codex Stan 12-1981. 2001;1–8.
25. Buba F, Gidado A, Shugaba A. Analysis of biochemical composition of honey samples from North-East Nigeria. *Biochem Anal Biochem.* 2013;2(3):139.
26. Ukom AN, Okereke IO, Ugwuona FU. PHYSICOCHEMICAL PROPERTIES AND ANTIOXIDANT ACTIVITIES OF HONEY FROM SOUTHERN NIGERIA. *J Chem Soc Niger [Internet].* 2019;44(6):1044-55
27. Hoxha F, Lamçe F, Beqo M, Kongoli R, Malollari I, Kyçyk O. Quality evaluation of commercialised honey in Tirana using physicochemical analysis. *Albanian J Agric Sci.* 2019;18(1):26–31.
28. Mijanur Rahman M, Gan SH, Khalil MI. Neurological effects of honey: Current and future prospects. Evidence-based Complement Altern Med. 2014;2014:958721.
29. Agarwal S, Bhardwaj V, Singh A, Khan MH, Goel S, Bharat M, et al. A control clinical trial of honey-impregnated and povidone iodine dressings in the treatment of diabetic foot ulcers among Northern Indian subjects. *Indian J Sci Res.* 2015;6(2):7–10.
30. Asif M. Physical and Chemical Characterizations, Storing conditions, Health Benefits and Medicinal Uses of Honey.
31. Mohamed H, Salma MA, Al Lenjawi B, Abdi S, Gouda Z, Barakat N, et al. The efficacy and safety of natural honey on the healing of foot ulcers: a case series. *Wounds a Compend Clin Res Pract.* 2015;27(4):103–14.
32. Gupta Jr S, Shinde S, Shinde RK, Gupta S. Topical management of wound: a narrative review of cadexomer iodine ointment versus povidone iodine ointment. *Cureus.* 2022;14(4):e24598.
33. Lepelletier D, Maillard JY, Pozzetto B, Simon A. Povidone Iodine: Properties, Mechanisms of Action, and Role in Infection Control and *Staphylococcus aureus* Decolonization. *Antimicrob Agents Chemother.* 2020;64(9): e00682-20
34. Bigliardi PL, Alsagoff SAL, El-Kafrawi HY, Pyon JK, Wa CTC, Villa MA. Povidone iodine in wound healing: A review of current concepts and practices. *Int J Surg [Internet].* 2017;44:260–8. doi:10.1016/j.ijssu.2017.06.073
35. Barreto R, Barrois B, Lambert J, Malhotra-Kumar S, Santos-Fernandes V, Monstrey S. Addressing the challenges in antisepsis: focus on povidone iodine. *Int J Antimicrob Agents.* 2020;56(3):106064.
36. Katz DL, Wild DM, Elmore JG, Lucan SC. Jekel's epidemiology, biostatistics, preventive medicine and public health. 4th Edition. Philadelphia, PA: Saunders/Elsevier.; 2014. 155 p.
37. Anumah FO, Mshelia-Reng R, Abubakar A, Sough T, Asudo F, Jamda MA, et al. Management outcome of diabetic foot ulcers in a teaching hospital in Abuja, Nigeria. *Age.* 2017;9(1):15-20.

38. Christman AL, Selvin E, Margolis DJ, Lazarus GS, Garza LA. Hemoglobin A1c predicts healing rate in diabetic wounds. *J Invest Dermatol.* 2011;131(10):2121–7.
39. Gulati S, Qureshi A, Srivastava A, Kataria K, Kumar P JA. A prospective randomized study to compare the effectiveness of honey dressing vs. povidone iodine dressing in chronic wound healing. *Indian J Surg.* 2014;76(3):193–8.
40. Haghightapanah M, Nejad ASM, Haghightapanah M, Thunga G, Mallayasamy S. Factors that Correlate with Poor Glycemic Control in Type 2 Diabetes Mellitus Patients with Complications. *Osong public Heal Res Perspect.* 2018 Aug;9(4):167–74.
41. Ogbera A, Ekpebegh C. Diabetes Mellitus in Nigeria; the past, present and future. *World J Diabetes.* 2014;5(6):905–11.
42. Chalya PL, Mabula JB, Dass RM, Kabangila R, Jaka H, McHembe MD, et al. Surgical management of Diabetic foot ulcers: A Tanzanian university teaching hospital experience. *BMC Res Notes.* 2011;4:365.
43. Edo AE, Eregie A. Bacteriology of diabetic foot ulcers in Benin City, Nigeria. *Diabetes Int.* 2007;21–3.
44. Hassan MA, Tamer TM, Rageh AA, Abou-Zeid AM, Abd El-Zaher EHF, Kenawy ER. Insight into multidrug-resistant microorganisms from microbial infected diabetic foot ulcers. *Diabetes Metab Syndr Clin Res Rev.* 2019;13(2):1261–70.

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