Maggot debridement therapy in the treatment of diabetic foot ulcers











Authors (from left to right): Harikrishna K.R.Nair, Nazni Wasi Ahmad, AA Ismail, Alabed Ali A Alabed, Ahmad Soffian

The incidence of diabetic foot ulcers (DFU) in Malaysia is about 15 % of diabetic cases and is increasing in trend as the population ages. DFUs are one of the major complications of long-standing diabetes, which may lead to amputation if left untreated or undertreated. Management of the diabetic foot requires a thorough knowledge of the major risk factors for amputation, frequent routine evaluation and meticulous preventive maintenance. The most common risk factors for ulcer formation include diabetic neuropathy, structural foot deformity and peripheral arterial occlusive disease. Maggot debridement therapy (MDT) is a type of biological therapy involving the introduction of live, sterile maggots (fly larvae) into non-healing skin and soft tissue wounds of a human to clean out the necrotic (dead) tissue or slough within a wound, and at the same time promote granulation tissue and epithelisation. We recruited nine patients with DFUs and used MDT to debride the wounds. The wound bed was assessed and the percentage of wound healing was assessed. Pain score was assessed and any side effects or complications such as myasis and bleeding were documented. All wounds showed reduction in slough and necrotic tissue with wound healing as assessed by area measurements. Pain reduction was noted and there were no untoward effects.

Harikrishna KR Nair, Wound Care Unit, Department of Internal Medicine, Hospital Kuala Lumpur and Reseach Scholar, Faculty of Medicine Lincoln University College Malaysia;

Nazni Wasi Ahmad,

Entomology Unit, Insistute of Medical Research, Ministry of Health Malaysia;

AA Ismail Community Medicine Department, Faculty of Medicine Lincoln University College Malaysia;

Ali A Alabed, Community Medicine Department, Faculty of Medicine Lincoln University College Malaysia;

Ahmad Soffian, Wound Care Unit, Department of Internal Medicine, Hospital Kuala Lumpur iabetes can lead to slow wounds healing in many patients. This happens primarily in older patients, the number of which is increasing, resulting in rising cost for the delivery of healthcare. It has been said that, the annual cost to manage these wounds exceeds 20 billion dollars (Harding et al, 2002) with a loss of over two million work days. Diabetic foot ulcers (DFU) are more difficult to treat and may lead to minor or major amputation of affected part if not treated or undertreated (International Diabetes Federation, 2010).

There are a lot of treatments available in association with wound healing in patients with diabeties, such as biotherapies, photobiomodulation, topical oxygen therapy and various advanced dressings. The primary focus of this study is biological therapy using maggots. Maggot debridement therapy (MDT) is the application of live fly larvae to

wounds to facilitate in wound debridement (cleaning), antimicrobial effect and/or healing. An infestation of maggot on a living host is called myiasis. When that infestation is limited to a wound, it is called wound myiasis. MDT is basically a therapeutic wound myiasis, controlled in ways that optimise efficacy and safety. The myiasis is carefully controlled by selecting the species and strain of fly (the species that being in this study is Lucilia cuprina), sterilising the larvae, using special dressings to maintain the larvae on the wound, and integrating quality control measures throughout the process (Nair et al, 2020). The Institute of Medical Research Malaysia found the species Lucilia cuprina to be the ideal maggot to be used in the therapy as it is the endemic species of Lucilia in Malaysia.

Historically, military surgeons were the earliest to find the beneficial effects of maggot-infested wounds. They noted that injured soldiers abandoned on the battlefield become better

and that their wound recovery was obviously improved when those wounds were infested with maggots (Pechter and Sherman, 1983). William Baer having observed maggot-infested wounds during his service, as an orthopedic surgeon in World War I, was the first to apply maggots systematically to non-healing wounds while he was a professor at Johns Hopkins and Children's Hospital in Baltimore, Maryland (Baer et al, 1931).

This research is to determine the clinical efficacy of MDT in terms of wound healing in DFUs carried out in Hospital Kuala Lumpur's Wound Care Unit. The time period of this report is six months from the initial approval of the research proposal. The aim of this study was to observe wound-related pain associated with MDT, the possible complications that may arise such as myiasis, bleeding from the time of wound debridement to wound bed preparation.

Methodology

Wound dressing and treatment

Patients with DFUs were evaluated for MDT and written consent for treatment was obtained from those selected. The maggots (*Lucilia cuprina*)

were obtained from the Medical Entomology Unit, IMR, Kuala Lumpur. Each vial contained 100-500 sterile maggots that were viable and used minimally for two to three MDT treatments. Distilled water was added to the vial containing maggots to loosen them. The maggots were pipetted out of the vial onto a piece of gauze. Assessment of the wound site along with pain score assessment were done on every visit. The wound perimeter was surrounded with a cage dressing, which is made up of gauze and micropore, used to adhere the gauze around the wound to prevent the maggots from escaping out of the wound bed. The gauze containing maggots was applied directly onto the wound bed. We used 10 maggots for every 1cm² of wound surface. The wound was then covered with light gauze to absorb the fluid, after this the entire foot was loosely bandaged with a crepe bandage. The bandage and gauze were changed as necessary before the washout. A washout of the wound was performed after 72 hours of maggot application. This was performed using distilled water. Maggots were then reapplied as needed. If no change seen after three consecutive applications then MDT was abandoned and other

Case 1

- An 81-year-old Malay gentleman with underlying diabetes on treatment
- Had a diabetic foot ulcer for more than 6 weeks, which was getting worse
- Maggot debridement therapy was performed six times, no side effects were observed (e.g. myiasis or bleeding)
- The wound improved by 31%

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Pre-maggot debridement therapy	Post-maggot debridement therapy
Area (length x width): 102cm	Area (length x width):7 0cm
Wound bed: Tendon exposed Sloughy tissue (100%)	Wound bed: Tendon exposed Granulation tissue (80%)
Pain score: 3	Pain score: 1





Case 2

- A 71-year-old Chinese gentleman with diabetes and mycosis fungoides stage II (mycosis fungoides is a cutaneous T-cell lymphoma) being treated by chemotherapy
- Patient claimed the wound was originated from a radiotherapy session
- Treated with one round of maggot debridement therapy, no side effects were observed (e.g. myiasis or bleeding)
- The wound improved by 12%

Pre-maggot debridement therapy	Post-maggot debridement therapy
Area (length x width): 31.35cm	Area (length x width): 27.5cm
Wound bed: Sloughy tissue (70%) Granulation tissue (30%)	Wound bed: Sloughy tissue (30%) Granulation tissue (70%)







Case 3

- A 65-year-old Malay gentleman, being treated for diabetes and hypertension since 2014
- Unresolved right cellulitis and goitre, referred to Wound Care Clinic for further management
- Treated with five rounds of maggot debridement therapy, no side effects were observed (e.g. myiasis or bleeding)
- The wound improved by 36%

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Pre-maggot debridement therapy	Post-maggot debridement therapy
Area (length x width): 55cm	Area (length x width): 35cm
Wound bed: Sloughy tissue (50%) Granulation tissue (30%) Epithelialisation (20%)	Wound bed: Granulation (70% Epithelialisation (30%)

Pain score: 3





forms of debridement carried out as needed.

Along the evaluation process, the patient will have their DFUs graded according to the University of Texas Medical Branch (UTMB) grading (Armstong et al, 1996). Treatment will be considered complete once the wound is healed, defined as the wound/ulcer being suitable enough for split-thickness skin grafting (STSG), flap coverage or self-healing as judged clinically. DFUs that are found suitable for desloughing and STSG or flap coverage at the same setting will also be considered as healed

Pain was measured using the Visual Analog Scale (VAS).

Results

Assessment of all nine patients who were enrolled in this study to date, positive outcome were achieved in 66% of all patients. There are six patients who have completed the study, their data showed:

- Successful debridement of wound surface area in which less than 5% of wound bed was covered by slough or necrotic tissue after MDT
- Improved healing rate, most wound suitable for wound closure by STSG or flap coverage

within one week

- Time to heal has reached optimal result with good wound bed preparation with minimum of two application, where there was a reduction in slough and necrotic tissue
- Reduction in wound-related pain from 3–2 to 2–0.

The other three patients still show significant improvement by at least 40% of slough debrided within the time frame. Furthermore, MDT was not harmful with no side effects reported in this study. The details of the DFU aetiologyi and treatments can be seen in Cases 1–9.

Discussion

Wound healing is basically a physiological process of repairing, regenerating, remodelling of the injured tissue which consist of four components: homeostasis; inflammation; proliferation; and remodelling. (Stechmiller and Schultz, 2007). In each phase, new cells are produced and transported into the injured tissue to aid healing, or cells already present alter their activity to secrete new cytokines or perform different tasks, in response to changing conditions in the wound, such as bleeding, hypoxia, alterations in cytokine concentrations. The cells then will undergo apoptosis, programmed cell death, and are

Case 4

Pain score: 1

- A 59-year-old, Malay gentleman. Known case of diabetes, with hypertension on treatment.
- Treated with three rounds of maggot debridement therapy, no side effects were observed (e.g. myiasis or bleeding)
- The wound improved by 67%

	Before maggot debridement therapy	Post maggot debridement therapy
	Area (length x width): 45cm	Area (length x width): 15cm
	Wound bed: Bone exposes Sloughy tissue (60%) Granulation tissue (40%)	Wound bed: Bone exposed Granulation tissue (100%)

Pain score: 0





Case 5

- A 51-year-old Indian lady with underlying diabetes on treatment
- Wound debridement done under the Orthopaedic department, Hospital Kuala Lumpur
- Treated with four rounds of maggot debridement therapy, no side effects were observed (e.g. myiasis or bleeding)
- The wound improved by 36%

Pre-maggot debridement therapy	Post-maggot debridement therapy
Area (length x width): 36cm	Area (length x width): 22.5cm
Wound bed: Sloughy tissue (40%) Granulation tissue (60%)	Wound bed: Sloughy tissue (80%) Granulation tissue (20%)
Pain score: 2	Pain score: 0





Case 6

- A 57-year-old Malay gentleman with underlying diabetes for 2 years, on treatment
- Develop an ulcer on the right big toe on 15.6.2020 because of blister due to pressure
- Treated with five rounds of maggot debridement therapy, no side effects were observed (e.g. myiasis or bleeding)
- The wound improved by 50%

Before maggot debridement therapy	Post maggot debridement therapy
Area cm (length x width): 24cm	Area cm (length x width): 12cm
Wound bed: Sloughy tissue (40%) Granulation tissue (60%)	Wound bed: Sloughy tissue (80%) Granulation tissue (20%)
Pain score: 2	Pain score: 0





Case 7

- A 65-year-old, Malay gentleman with underlying diabetes and hypertension, which have been undertreatment since 2014
- Unresolved left cellulitis and referred to Wound Care Clinic for futher management.
- Treated with four rounds of maggot debridement therapy, no side effects were observed (e.g. myiasis or bleeding)
- The wound improved by 66%

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Pre-maggot debridement therapy	Post-maggot debridement therapy
Area (length x width): 35cm	Area (length x width): 12cm
Wound bed: Necrotic tissue (80%) Sloughy tissue (20%)	Wound bed: Granulation tissue (100%)
Pain score: 2	Pain score: 0





Case 8

- A 76-year-old, Indian gentleman with underlying siabetes on treatment
- Wound debridement and ray's amputation done in hospital
- Treated with three rounds of maggot debridement therapy, no side effects were observed (e.g. myiasis or bleeding)
- The wound improved by 34%

Pre-maggot debridement therapy	Post-maggot debridement therapy
Area (length x width): 24cm	Area (length x width): 12cm
Wound bed: Sloughy tissue (80%) Granulation tissue (20%)	Wound bed: Sloughy tissue (20%) Granulation tissue (80%)
Pain score: 3	Pain score: 2





References

Armstrong D, Lavery LA, Harkless LB (1996) Treatment-based classification system for assessment and care of diabetic feet J Am Podiatr Med Assoc 86(7): 311-6. https://doi. org/10.7547/87507315-86-7-311

Harding KG, Morris HL, Patel GK (2002) Science, medicine and the future: healing chronic wounds. BMJ 324(7330):160-3. https://doi. org/10.1136/bmj.324.7330.160

Nari HKR, Ahmad NW, Teh CH (2020) Maggot Debridement Therapy in Malaysia, Int 11 ow Extrem Wounds 11;1534734620932397. https://doi. org/10.1177/1534734620932397

Pechter EA, Sherman RA (1983) Maggot therapy: the surgical metamorphosis. Plast Reconstr Surg 172(4):567-70 https:// doi.org/10.1097/00006534-198310000-00032

Baer WS (1931) The treatment of chronic osteomyelitis with the maggot (larva of the blow fly). J Bone Joint Surg Am 13:438-75

Cowan LJ, Stechmiller JK, Philips P et al (2013) Chronic wounds, biofilms and use of medicinal larvae. Ulcers 487024. https://doi. org/10.1155/2013/487024

Horobin AJ, Shakesheff KM, Pritchard DI (2006) Promotion of human dermal fibroblast migration, matrix remodelling and modification of fibroblast morphology within a novel 3D model by Lucilia sericata larval secretions. J Invest Dermatol 126(6):1410-8, https://doi. org/10.1038/sj.jid.5700256

Prete PE (1997) Growth effects of Phaenicia sericata larval extracts on fibroblasts: mechanism for wound healing by maggot therapy. Life Sci 60(8):505-10. https://doi.org/10.1016/s0024-3205(96)00688-1

Sherman RA, Sherman J, Gilead L (2001) Maggot débridement therapy in outpatients. Arch Phys Med Rehabil 2001 82(9):1226-9. https://doi.org/10.1053/ apmr.2001.24300

Sherman RA (2002) Maggot versus conservative debridement therapy for the treatment of pressure ulcers. Wound Repair Regen 10(4):208-14. https:// doi.org/10.1046/j.1524-475x.2002.10403.x

Sherman RA (2003) Maggot therapy for treating diabetic foot ulcers unresponsive to conventional therapy. Diabetes Care 26(2):446-51. https://doi.org/10.2337/ diacare.26.2.446

engulfed by macrophages to be removed once no longer needed. Normally, these four stages in the healing process progressed quickly and smoothly based on its natural sequences. However, in certain cases such as presence of infection, biofilm or ischaemia, the healing may cease, leading to non-healing wound. Wound healing may cease at any phase (or even while undergoing a combination of phases), but commonly it is within the inflammatory phase: dead, infected debris may not be adequately removed from the wound bed, and/or it might not be possible for the body to eradicate the local infection, and/ or the proteases and other destructive products of inflammation by clearing the newly formed cellular and extracellular matrix as fast as it is being laid down (Stechmiller and Schultz, 2007). It is in this context that debridement, disinfection, or cellular proliferation and migration are so important, as they can push the non-healing wound into the next phase of healing.

There have been several studies conducted to $determine \ how \ the \ maggots \ increase \ granulation$ in the wound bed. A study conducted in 2006 demonstrated an increased migration (but not proliferation) of the fibroblasts which was attributed to the action of serine and metalloproteinases. (Horobin et al, 2006). Another study found high levels of gamma interferon and interleuken-10 in the excretions of maggots that were thought to increase granulation tissue formation (Prete, 1997).

About 5-30% of wounds treated with MDT associated with pain or discomfort is reported in some studies. (Sherman et al, 2001; Sherman, 2002;2003.) At the same time, these patients also have chronic wound pain even before treatment with MDT. Therefore, they are likely to experience pain or discomfort during and after treatment, and also can be readily identified, warned, and treated appropriately with analgesics if needed. Commonly, pain generally occurs after the first 24 hour, as the maggots increase in size. Analgesic is a good choice to control the pain, but if not, taking out the dressing and removing of the maggots will reduce and halt the pain or discomfort in no time.

Finally, MDT is a great modality to be used in managing DFUs especially for wound debridement. However, a more robust trial should be conducted to show the significance of MDT.

- A 53-year-old, Indian lady with underlying diabetes being treated for more than three years
- Rays amputation done under Orthopaedic Surgeon in Klang on 22 June 2020, at presentation her tendon was slighlty necroctic and sloughy
- Treated with four rounds of maggot debridement therapy, no side effects were observed (e.g. myiasis or bleeding)

■ The wound improved by 58%	
Pre-maggot debridement therapy	Post-maggot debridement therapy
Area (length x width): 108cm	Area (length x width): 45cm
Wound bed: Tendon exposed Sloughy tissue (60%) Granulation tissue (40%)	Wound bed: Granulation tissue (80%) Epithelialisation(20%)
Pain score: 3	Pain score: 2



