Review

Maggot Therapy: The Science and Implication for CAM
Part I—History and Bacterial Resistance

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It is now a universally acknowledged fact that maggot therapy can be used successfully to treat chronic, long-standing, infected wounds, which have previously failed to respond to conventional treatment. Such wounds are typically characterized by the presence of necrotic tissue, underlying infection and poor healing. Maggot therapy employs the use of freshly emerged, sterile larvae of the common green-bottle fly, *Phaenicia (Lucilia) sericata*, and is a form of artificially induced myiasis in a controlled clinical situation. In this review article, we will discuss the role of maggots and their preparation for clinical use. Maggot therapy has the following three core beneficial effects on a wound: debridement, disinfection and enhanced healing. In part I we explore our current understanding of the mechanisms underlying these effects.

Keywords: Maggot debridement therapy – MRSA – antimicrobial – *Lucilia sericata* – wounds

Introduction I—The Rise and Fall of Maggot Therapy

Numerous clinical reports have been published that describe the outstanding effects of maggot therapy, most notably on debridement, cleansing, disinfection and healing of indolent wounds, many of which have previously failed to respond to conventional treatment (1–11). Current day maggot therapy, with its multi-action approach to wound cleansing and healing, is highly successful.

Records of maggots in wounds, however, and the recognition of improvement in the wound state as a consequence of infestation, date back to the 16th century (12). In 1829, Baron Dominic Larrey, Napoleon’s battlefield surgeon, described how men had arrived at his field hospital with healing maggot-infested wounds (13). The wounds were sustained in battle, but, owing to the presence of maggots, were not infected and showed accelerated healing. Such positive accounts were made by many surgeons who followed, but it was William Baer, Professor of Orthopaedic Surgery at the John Hopkins School of Medicine in Maryland, USA, who is believed to be the founder of modern maggot therapy (14).

It was Baer who pioneered the use of sterile maggots as a reputable method of wound therapy, following observations he made about the value of maggots in traumatic wounds on the battlefield in France during World War 1. Such was the success of Baer’s work that by the mid-1930s almost 1000 North American surgeons employed maggot therapy (15) and by the end of the decade it was in use in over 300 hospitals in the US and Canada. However, by 1940, a new era was dawning. This era which saw the introduction and widespread use of antibiotics following the mass production of penicillin (16). So despite the obvious success of maggot therapy, by the mid-1940s it had practically disappeared from use. In Part 1 of this review, we introduce the stages involved in the wound healing process, the advantages in the use of maggots for the cleaning (debridement) of infected wounds, and the possible mechanisms underlying the debridement of wounds by maggots. The antimicrobial activity of maggots to treat methicillin-resistant *Staphylococcus aureus* (MRSA)-infected wounds

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Maggot Therapy: Selection of the Flies

Many dipteran species are capable of infesting living vertebrate hosts (a condition termed myiasis). Maggot therapy is essentially artificially induced myiasis, performed in a controlled environment by experienced medical practitioners. Myiasis-causing flies may be grouped into two categories as follows: obligate and facultative parasites. Obligate parasites require the ingestion of living tissue in order to complete their lifecycles (22). Larvae of obligate parasites can cause severe damage to healthy tissue and are therefore unsuitable for use in maggot therapy. Facultative parasites are able to parasitize living hosts if conditions are favorable, but more commonly develop on carrion and therefore have greater potential for therapeutic use.

Selection of a suitable fly species for use in maggot therapy is of paramount importance, determining both the safety and success of the treatment. It is imperative to select a species that feeds almost exclusively on necrotic tissue. William Baer chose the larvae of Phaenicia sericata, the common green-bottle, as the most appropriate species for this application and this is the species still used by practitioners today. Phaenicia larvae are facultative parasites, unable to ingest or significantly damage healthy human tissue (2). Infestations of living hosts by Phaenicia do, however, occur, most commonly in sheep to induce an often fatal condition known as sheep strike. Exactly why Phaenicia attack the healthy tissue of sheep and appear unable to do the same to human tissue is as yet unknown.

Female Flies, Eggs, Larvae and Preparation for Clinical Use

In the wild, adult female Phaenicia lay a large number of eggs (2000–3000) over the course of a few weeks, a necessity as relatively few will survive to adults. The eggs are laid in clusters directly onto the chosen food source, upon which the emerging larvae will feed. Larval development requires a moist environment to prevent desiccation, so larvae are generally found in nutritious, damp places such as decaying animal corpses or moist, necrotic wounds (22). Eggs hatch within 18–24 h, depending on optimal conditions, into first instar larvae (maggots), ~1–2 mm in length, which immediately and actively begin to feed. It is this vigorous feeding activity, which is beneficial to an infected or necrotic wound. Maggots feed by the extracorporeal secretion of a wide spectrum of proteolytic enzymes that liquefy the host tissue (23–26). This semi-digested liquid material is then ingested as a source of nutrients. The maturing first instar larvae continue to feed by the extracorporeal secretion of a wide spectrum of proteolytic enzymes that liquefy the host tissue (23–26). This semi-digested liquid material is then ingested as a source of nutrients. The maturing first instar larvae continue to feed by the extracorporeal secretion of a wide spectrum of proteolytic enzymes that liquefy the host tissue (23–26). This semi-digested liquid material is then ingested as a source of nutrients. The maturing first instar larvae continue to feed by the extracorporeal secretion of a wide spectrum of proteolytic enzymes that liquefy the host tissue (23–26). This semi-digested liquid material is then ingested as a source of nutrients. The maturing first instar larvae continue to feed by the extracorporeal secretion of a wide spectrum of proteolytic enzymes that liquefy the host tissue (23–26). This semi-digested liquid material is then ingested as a source of nutrients. The maturing first instar larvae continue to feed by the extracorporeal secretion of a wide spectrum of proteolytic enzymes that liquefy the host tissue (23–26). This semi-digested liquid material is then ingested as a source of nutrients. The maturing first instar larvae continue to feed by the extracorporeal secretion of a wide spectrum of proteolytic enzymes that liquefy the host tissue (23–26). This semi-digested liquid material is then ingested as a source of nutrients. The maturing first instar larvae continue to feed by the extracorporeal secretion of a wide spectrum of proteolytic enzymes that liquefy the host tissue (23–26). This semi-digested liquid material is then ingested as a source of nutrients. The maturing first instar larvae continue to feed by the extracorporeal secretion of a wide spectrum of proteolytic enzymes that liquefy the host tissue (23–26). This semi-digested liquid material is then ingested as a source of nutrients. The maturing first instar larvae continue to feed by the extracorporeal secretion of a wide spectrum of proteolytic enzymes that liquefy the host tissue (23–26).
upon emergence from the egg and undergo rigorous testing to ensure their microbiological status (3). Larvae are then maintained under aseptic conditions prior to wound application.

**Debridement (Wound Clearing)**

Maggot therapy has the following three core beneficial effects on a wound: debridement, disinfection and enhanced healing. Debridement is the removal of cellular debris and non-viable necrotic tissue from the wound bed. This is a first, essential step before healing can commence. Removal of necrotic tissue abolishes many of the associated bacteria and also reduces wound odor. The removal of necrotic tissue, which acts as a microbial substrate, may also reduce the risk of infection. During the inflammatory stage of wound healing host leukocytes play an important role in debridement of wound sites, degrading damaged extracellular matrix (ECM) components through the release of proteases. The injury is initially filled with a provisional wound matrix consisting predominantly of fibrin and fibronectin. Key proteases are involved in ECM degradation (see below). These are released from neutrophils, macrophages, fibroblasts, epithelial and endothelial cells. As healing proceeds, and new ECM constituents such as collagen, elastin and proteoglycans are synthesized, damaged ECM is removed by these proteases (27).

**Chronic Wounds**

Chronic wounds do not proceed through the normal healing process and are typically characterized by prolonged inflammation, inhibition of cell proliferation (28,29), incomplete ECM remodeling and a failure to epithelialize (30). Over expression and inefficient debridement of temporary ECM components, e.g. fibronectin and fibrin, contribute to the failure of chronic wounds to heal. The entire environment of a chronic wound must be rebalanced for wound repair to proceed to completion, an undertaking which is unlikely to occur without extraneous intervention and one of the explanations as to why chronic wounds may persist for many years.

There are a number of existing methods for the debridement of chronic wounds as described by Schultz et al. (27). These include surgical and sharp debridement (using scalpel or scissors to remove debris and necrotic tissue), mechanical debridement using methods such as wet-to-dry dressings, wound irrigation and whirlpool techniques, enzymatic debridement using the application of exogenous enzymes, and autolytic debridement using hydrogels and hydrocolloids. Each of these techniques has associated disadvantages such as extended treatment times, pain and mechanical damage to underlying healthy tissue.

**Maggot Debridement Therapy**

The alternative is maggot therapy. Maggots debride wounds quickly and effectively, without damage to viable tissue. Maggots are photophobic and will naturally move into the deep crevices that may be beyond the reach of a surgeon’s scalpel. Reports have been published marveling at the benefits of maggot debridement therapy (MDT) in all sorts of wounds, including abscesses, burns, gangrenous wounds, arterial and venous ulcers, osteomyelitis, diabetic foot ulcers and pressure sores (7,9,31–33). One such study compared MDT with conservative debridement therapy for the treatment of pressure sores (34). Here, 80% of maggot-treated wounds (n = 43) were completely debrided, while only 48% of conventionally treated wounds (n = 49) were completely debrided. Also, by using maggots, total wound surface area decreased, whereas during conventional debridement therapy, the total wound area had increased (p = 0.001) (34). The report concluded that maggot therapy was a more effective and efficient way of debriding chronic pressure sores than the conventional treatments prescribed.

**Mechanisms of MDT**

How exactly maggots remove devitalized, necrotic tissue from the wound is currently actively being investigated. Research into the debridement mechanisms underlying maggot therapy has revealed that maggots secrete a rich soup of digestive enzymes while feeding, including carboxypeptidases A and B (35), leucine aminopeptidase (35), collagenase (23,36) and serine proteases (trypsin-like and chymotrypsin-like enzymes) (35,37). Recently, workers in Nottingham, UK, demonstrated in vitro a range of enzymes secreted by P. sericata larvae (26). Four proteolytic enzymes, comprising two serine proteases, a metalloproteinase and an aspartyl proteinase, were detected, with molecular weights ranging from 20 to 40 kDa, with activity across a wide pH range. A chymotrypsin-like serine proteinase exhibited excellent degradation of ECM components laminin, fibronectin, and collagen types I and III (26), and may therefore play a significant role in the digestion of wound matrix and effective debridement.

The mechanical action of numerous wriggling maggots in a necrotic debris-filled wound has also been suggested in aiding wound debridement. Maggots possess a pair of mandibles (hooks) which assist with locomotion and attachment to tissue. This probing and maceration of wound tissue with maggot mouthhooks may enhance debridement (38), but these hooks are used during feeding to disrupt membranes and thus facilitate the penetration of proteolytic enzymes (3). Together, this mechanical action and the secretion of powerful, proteolytic enzymes may be the secret of efficient tissue debridement.

**Disinfection (Introduction to Antibiotic Activity)**

For wounds to heal, and progress through stages of destruction and proliferation onto maturation, infection needs to be eliminated. The majority of wounds are polymicrobial, hosting a range of both anaerobic and aerobic bacteria (18,39). Antimicrobial treatment of clinically infected and non-healing wounds, should, therefore, encompass broad-spectrum anti-
microbials in order to cleanse the wound effectively. The application of maggots to an infected wound results in the rapid elimination of such infecting microorganisms (2,6,40,41). The most frequently isolated pathogen from acute and chronic wounds is Staphylococcus aureus. S. aureus is carried innocuously by ~30% of the general population (42) [40–70% of hospital staff (43,44)], usually on the moist skin in the nose, axillae (armpits) and perineum (groin), but can become pathogenic when able to enter damaged skin. S. aureus has caused great concern owing to its ability to acquire resistance to a range of antimicrobials.

Penicillin Methicillin Resistance and MRSA

In 1948, 4 years after the widespread introduction of penicillin, over 50% of nosocomial S. aureus were penicillin-resistant (45) owing to the production of penicillinase (β-lactamase), an enzyme which inactivates β-lactam antibiotics (46). Currently, the majority (80–90%) of S. aureus are penicillin-resistant. In 1960, a structural modification of penicillin saw the synthetic production of methicillin, which was active against penicillin-resistant strains of S. aureus. The launch of methicillin, however, failed to control the proliferation of resistant strains of bacteria and the first clinical isolate of MRSA was reported in 1961 (47). Since then, MRSA has continued to disseminate rapidly, causing serious hospital and community infections all over the world, with global increases in both the numbers of infected patients and mortality. The recent isolation of vancomycin-resistant strains of S. aureus (VRSA) in Japan (48) severely reduces the repertoire of drugs available to treat infections caused by resistant strains of S. aureus.

In the literature, there is an ever increasing trend supporting the clinical use of maggots for treating wounds infected with MRSA (6,40,49,50). This support, initially anecdotal, was strengthened by case studies and most recently, strong laboratory evidence indicates that maggots do possess the ability to kill clinical isolates of MRSA (51,52). As an example, Fig. 1 shows a wound 5 cm in diameter and totally covered with a thick layer of viscous slough. One pot of larvae was applied and left for 48 h, after which there was an immediate and marked improvement to the wound. Two further applications of larvae were made. At this point, only 6 days after maggot therapy had commenced, the wound, which had not responded to conventional treatment over 18 months, was now completely free from slough. It was filling rapidly with healthy granulation tissue and a swab failed to detect any presence of MRSA (Fig. 2). Larval therapy was now discontinued and the wound continued to progress normally and healed uneventfully (6).

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