The effectiveness of chitosan-based bioactive microfibre gelling (BMG[™]) dressing in postoperative wounds: a case series

Introduction: Maxiocel® (CD Medical) is a bioactive microfiber gelling (BMG) dressing which is made up of 100% chitosan, a polysaccharide obtained from natural resources. Chitosan is known for its inherent haemostatic, antimicrobial, biocompatible, moisture-regulating, biodegradable and anti-inflammatory characteristics. This case series investigated the effectiveness of BMG dressing in promoting the healing of postoperative wounds. Methods: Four different outpatient cases were treated using BMG dressing. Prior to dressing application, wounds were cleansed using an antimicrobial solution and debridement was carried out when necessary. Periodically, wound dimensions and photographs were recorded to ensure comprehensive documentation. Results: All four patients showed a large reduction in wound size, especially patient 2 who is 56 years old with pre-existing diabetes, hypertension and dyslipidaemia, diagnosed as having a postoperative wound secondary to infected left BKA stump. His wound showed 92.42% reduction in size in 41 days. Conclusion: BMG dressing offers an effective solution for accelerating the wound healing process, making it a promising option in modern wound care.

ioactive microfiber gelling (BMG™) dressing is a type of wound dressing material that is made of up 100% chitosan, a natural polysaccharide derived from natural resources including shellfish, mushrooms and insects. It is derived from deacetylated chitin with a minimum of 50% of its amine groups in the free form (Zhang et al, 2020). Chitosan has several characteristics that make it suitable for use as a modern dressing, such as antimicrobial, anti-inflammatory, low immunogenicity and antioxidant properties (Patrulea et al, 2015; Dragostin et al, 2016).

Chitosan exerts a beneficial impact on each of the four phases of wound healing. During the haemostasis phase, chitosan exhibits the capacity to absorb plasma, induce coagulation of erythrocytes, promote platelet aggregation and activate platelets. Red blood cells adhere to chitosan macromolecule chains, hence facilitating the haemostatic process (Whang et al, 2005). During the inflammatory phase, chitosan exerts an anti-inflammatory action via modulating the activity of tumor necrosis factor-alpha (TNF- α), which subsequently decreases the production of nitric oxide (Yoon et al, 2007). During the proliferative phase, lysozymes break down chitosan to create N-acetyl-D-glucosamine, which promotes the growth of fibroblasts and the formation and

remodelling of collagen (Axio Biosolutions). Chitosan dressing also inhibits the metallomatrix proteases (MMP 2 and MMP 9) that help to regenerate the extracellular matrix quickly. It also speeds up the granulation process in the proliferative phase (Archana et al, 2013). In the maturation phase, chitosan contains a component of N-acetyl glucosamine that helps to reduce scar formation (Nair et al, 2020).

Objectives

The aim of this case series is to discuss the efficacy of BMG dressings in facilitating wound healing in post-surgical wounds.

Method

This case series involved the application of a BMG dressing (MaxioCel®, CD Medical) on four patients. It was conducted in an outpatient setting at Wound Care Clinic (WCC) Hospital Teluk Intan. Table 1 illustrates the baseline demographic summary of enrolled patients. For all patients, the TIME concept was used to assess the wounds and Harikrishna Periwound Skin Classification (HPSC) [Table 1] was used to describe the periwound area. The wound was cleansed using a surfactant containing antimicrobial solution (Octenisept®, Schülke) and, as per the wound requirements, debridement was performed, prior to a BMG

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Key words

- Bioactive microfiber aellina
- Case series
- Chitosan
- Postoperative wounds

Declarations

The authors have no conflicts of interest to declare.

Table 1. Baseline demographics summary of enrolled patients.											
Patient ID	Age	Comorbidities	Wound location	Diagnosis	Duration of wound prior to BMG dressing	Wound characteristics	Duration of treatment (days)	Previous dressing used			
1	54	Type 2 diabetes	Left foot	Post surgical wound secondary to infected left diabetic foot ulcer	7 days	Moderate exudate HPSC class 1	85	Silver hydro fiber and hydrogel			
2	56	Type 2 diabetes, dyslipidaemia, hypertension	Left knee	Postoperative wound secondary to infected left BKA stump.	30 days	Moderate exudate HPSC class 1	41	Silver hydrofiber and hydrogel			
3	73	Lower eyelid basaloid squamous cell carcinoma, Hypertension	Left eye	Postoperative wound secondary to left eye fungating squamous cell carcinoma.	14 days	Moderate exudate HPSC class 1	63	Silver hydrofiber			
4	69	Type 2 diabetes	Left hand	Postoperative wound secondary to left hand NF.	30 days	Moderate exudate HPSC class 1	61	Silver hydrofiber and hydrogel			

Table 2. Summary of clinical observations and results.											
Patient ID	Wound size (cm²)		Wound size	Exudate level							
	Before treatment	After treatment	reduction (%)	Before treatment	After treatment						
1	156.45	14.3	90.86	Moderate	Low						
2	66	5	92.42	Moderate	Low						
3	17.86	7.5	58	Moderate	Low						
4	170	17.5	89.71	Moderate	Low						

dressing being used. Once the tendon was almost closed, we initiated the BMG dressing usage. Gamgee roll was used in the outer layer as a secondary dressing and changed on a daily basis.

Ethical approval

Written consent from the patients for the use of patient details and photographs was obtained.

Results

Table 2 summarises the clinical outcomes, including the major observations and the key results of these clinical evaluations.

Patient 1: Postoperative wound secondary to infected left diabetic foot ulcer

This case involved a 52-year-old woman with pre-existing type 2 diabetes (T2D). Initially, she presented with an infected diabetic foot ulcer

that required wound debridement. She was referred to the wound care clinic (WCC) following her discharge for wound management. Figure 1 depicts the progression of wound healing during the treatment period. The wound had a 90.86% reduction in size after 85 days of treatment.

Patient 2: Postoperative wound secondary to infected left below-knee amputation stump

This case centred on a 56-year-old man with pre-existing T2D, hypertension and dyslipidaemia, who underwent left below-knee amputation due to wet gangrene in sepsis. The patient's blood glucose levels were challenging to manage in the hospital ward as a result of infections. He had to undergo repeated wound debridement procedures for his stump. The results of the tissue culture revealed the presence of *Pseudomonas aeruginosa*, which was then treated with appropriate antibiotics

Figure 1. Patient 1: The progression of wound healing during the treatment period.



Figure 2. Patient 2: The progression of wound healing during the treatment period.



Figure 3. Patient 3: The progression of wound healing during the treatment period.



Day 1: L 3.8cm × w 4.7 cm (17.86 cm²)



Day 14: L 3.8cm × w 4.5 cm (17.1 cm²)



Day 45: L 3.5 cm × w 4cm (14cm²)



Day 63: L 3cm × w 2.5 cm (7.5 cm²)

Figure 4. Patient 4: The progression of wound healing during the treatment period.



Day 1: L 17cm × w 10cm (170cm²)



Day 18: L 15 cm × w 9 cm (135 cm²)







Day 45: L 11cm × w 4cm (44cm²)

based on the wound's sensitivity. Subsequently, he was referred to the WCC to continue his wound treatment. Figure 2 shows wound size reduction by 92.42% from initial wound size after 41 days of treatment using BMG dressing.

Patient 3: Postoperative wound secondary to left eye fungating squamous cell carcinoma. This case is a 73-year-old man with preexisting hypertension who was diagnosed

with left lower eyelid basaloid squamous cell carcinoma in 2019. He defaulted on his treatment and follow-up for the previous 2 years. He returned in November 2023 for a fungating mass with maggot infestation over his left eye. The left eye exenteration was performed at Serdang Hospital, Malaysia, within the same week. He was then referred to WCC for wound management for his left eye. Figure 3 illustrates the wound progression throughout the treatment in which wound size was reduced by 58% in 63 days.

Patient 4: Postoperative wound secondary to left hand necrotising fasciitis

A 69-year-old man with newly diagnosed T2D was admitted for sepsis secondary to left hand necrotising fasciitis. He had extensive wound debridement and ray amputation of his left thumb. Subsequently, he was referred to the WCC for his post-surgical wound. Figure 4 demonstrates the wound healing progress where his wound reduced by 89.71% from initial wound size in 73 days.

Discussion

In this case series, all four patients were treated using a BMG dressing, which is commonly used for a variety of wounds at different anatomical sites.

Although BMG dressing can be used in combination with other wound care products, this study focused on the use of BMG dressing alone to assess its individual efficacy. The dressing also conforms to the wound bed, is easy to apply and can be removed without the patient experiencing any pain, thus making it much more suitable for all types of wounds.

BMG dressing's microfibers absorb fluid from a wound and form a gel-like substance upon contact with wound exudate, thus helping to prevent periwound maceration and making it suitable for low to moderate exudating wounds. As it is in gel form, it helps in maintaining a moist environment, which aids in promoting good granulation tissue formation as shown in all four cases. Nevertheless, other elements play a role in the process of wound healing (Archana et al, 2013). Patient 3 has the lowest percentage of wound size reduction when compared to the other three patients. The slower healing could be attributed to factors such as advanced age and his immunocompromised status. BMG dressing aids in autolytic debridement by moistureretentive properties, which helps soften nonviable tissue and make it easier to remove during mechanical debridement (Archana et al, 2013).

The BMG dressing provides an optimal moist wound environment, a critical factor for expediting wound healing. It also forms a barrier that protects the wound from external contaminants while permitting passage of oxygen. The gel formation additionally serves to entrap the pathogenic bacteria from the wound site and prevent the dressing from adhering to the wound bed, hence reducing the likelihood of pain and trauma to the wound bed (Archana et al, 2013).

Another observable feature in this case series is the degree of exudate. As per the World Union of Wound Healing Societies Consensus Document, 'Wound Exudate: Effective Assessment and Management' (World Union of Wound Healing Societies, 2019), the factors that influence the level of wound exudate are wound aetiology, the stage of wound healing, the size of the wound and the presence of other medical conditions and complications in the patient. **Table 2** displays all four cases that demonstrate a decrease in exudate level, as the size of the wound decreases over time.

Limitations

The number of patients was small and, thus, it may require another study involving larger patient numbers and multiple centres in future.

Conclusion

This case series demonstrated that the chitosan-based BMG dressing is an effective adjunct in managing postoperative wounds. All four patients saw a significant reduction in wound size and exudate levels, supporting the BMG dressing's role in accelerating wound healing, promoting granulation tissue formation and maintaining a moist wound environment. The dressing's ease of use, biocompatibility and pain-free application further enhance patient comfort and treatment compliance.

References

- Archana D, Singh BK, Dutta J, Dutta PK (2013) In vivo evaluation of chitosan-pvp-titanium dioxide nanocomposite as wound dressing material. *Carbohydr Polym* 95(1): 530-9
- Axio Biosolutions (2025) Wound Healing Properties. Available at: https://axiobio.com/characterization-of-maxiocel/(accessed 19.09.2025)
- Dragostin OM, Samal SK, Dash M et al (2016) New antimicrobial chitosan derivatives for wound dressing applications. *Carbohydr Polym* 141: 28-40
- Nair HKR, Chong SS, Othman AM (2020) Validation of harikrishna periwound skin classification for wound assessment. J Wound Care 29(Sup4): S44_S48
- Patrulea V, Ostafe V, Borchard G, Jordan O (2015) Chitosan as a starting material for wound healing applications. *Eur J Pharm Biopharm* 97(Pt B): 417-26
- Whang HS, Kirsch W, Zhu YH et al (2005) Hemostatic agents derived from chitin and chitosan. *J Macromolec Sci C* 45(4): 309–23
- World Union of Wound Healing Societies Consensus
 Document (2019) Wound Exudate: Effective
 Assessment and Management. Available at:
 https://woundsinternational.com/wp-content/
 uploads/2023/02/836aed9753c3d8e3d8694bcaee336395.
 pdf (accessed 19.09.2025)
- Yoon HJ, Moon ME, Park HS et al (2007) Chitosan oligosaccharide (cos) inhibits lps-induced inflammatory effects in raw 264.7 macrophage cells. *Biochem Biophys* Res Commun 358(3): 954-9
- Zhang S, Li Jianyong, Li Jianfeng et al (2020) Application status and technical analysis of chitosan-based medical dressings: a review. RSC Adv 10(56): 34308-34322