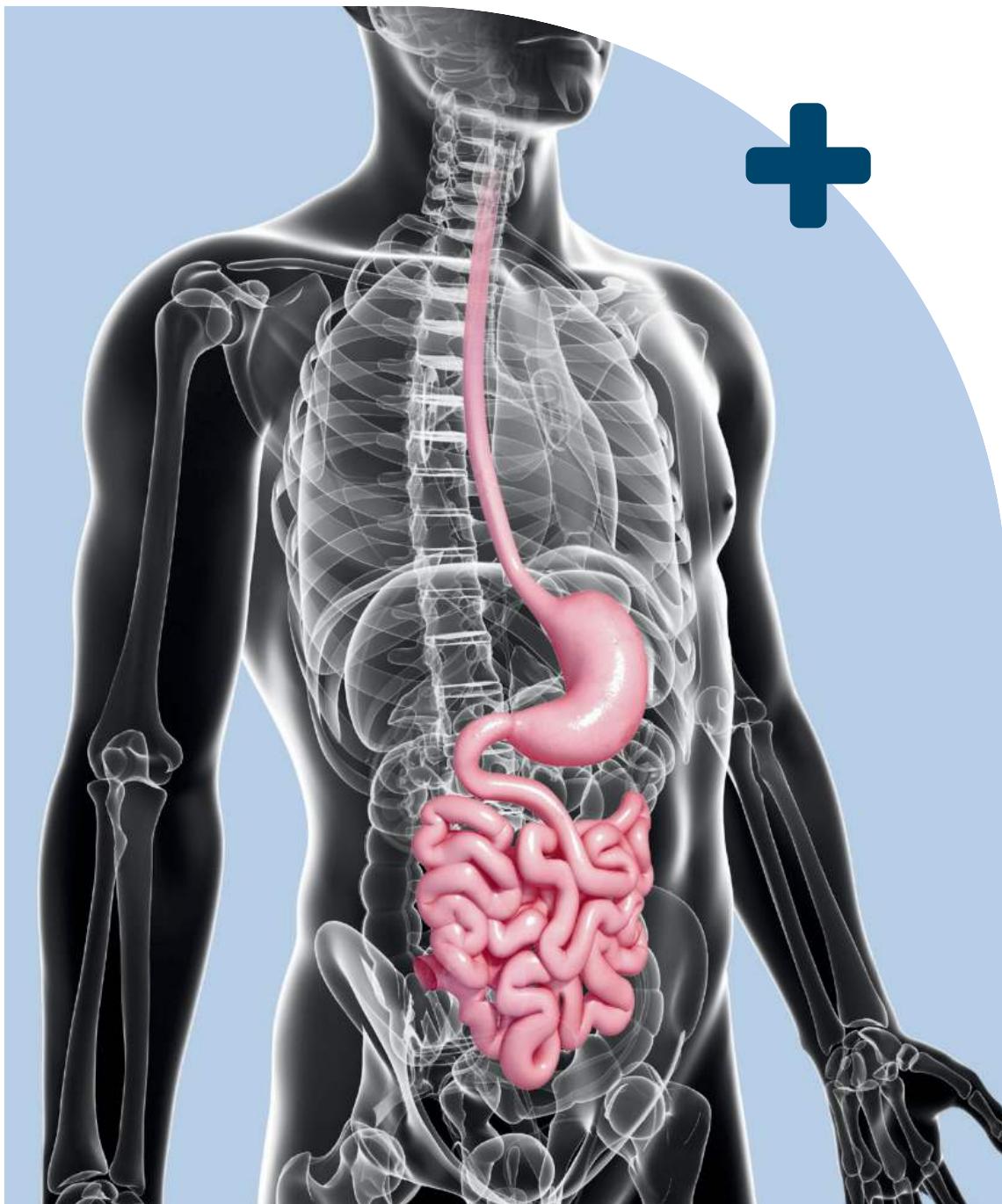


# Gastrointestinal Mucosal Healing Gel



Esophageal



Gastric



Intestinal



PT. Wickland Medika Imantara  
Medical, Aesthetics & Laboratory Supplier

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## PRODUCT FEATURE

### Acid-Sensitive Smart Biomaterials

(Available Size: 5 ml, 10 ml/Storage Condition: 0°C-40 °C)



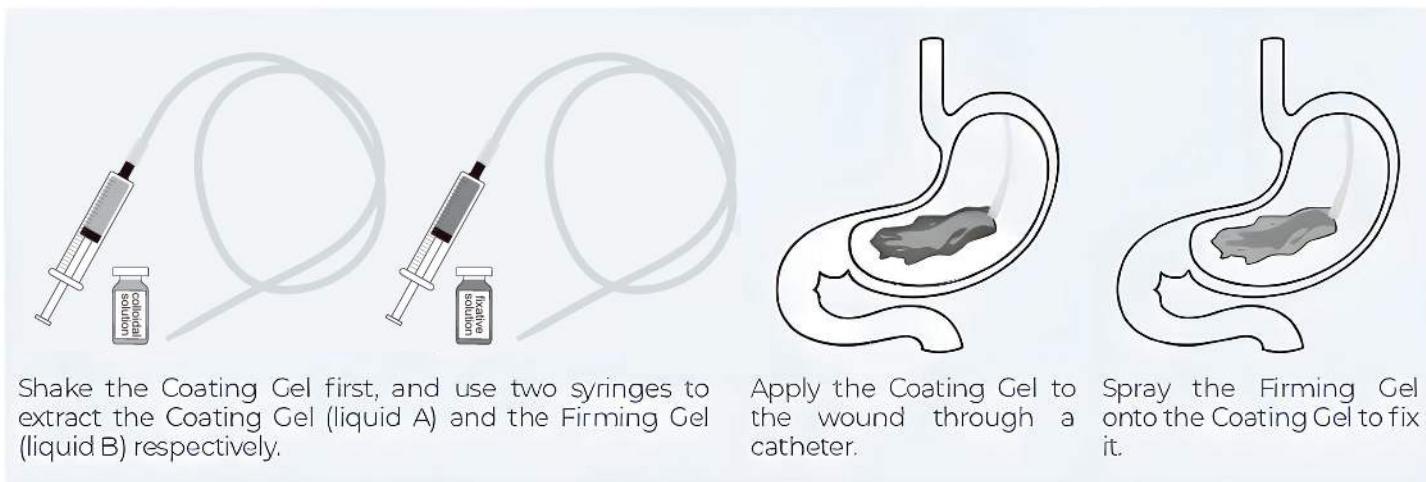
1/ Innovative materials

3/ World's first

2/ Exclusive patent

4/ Filling the gap in the field

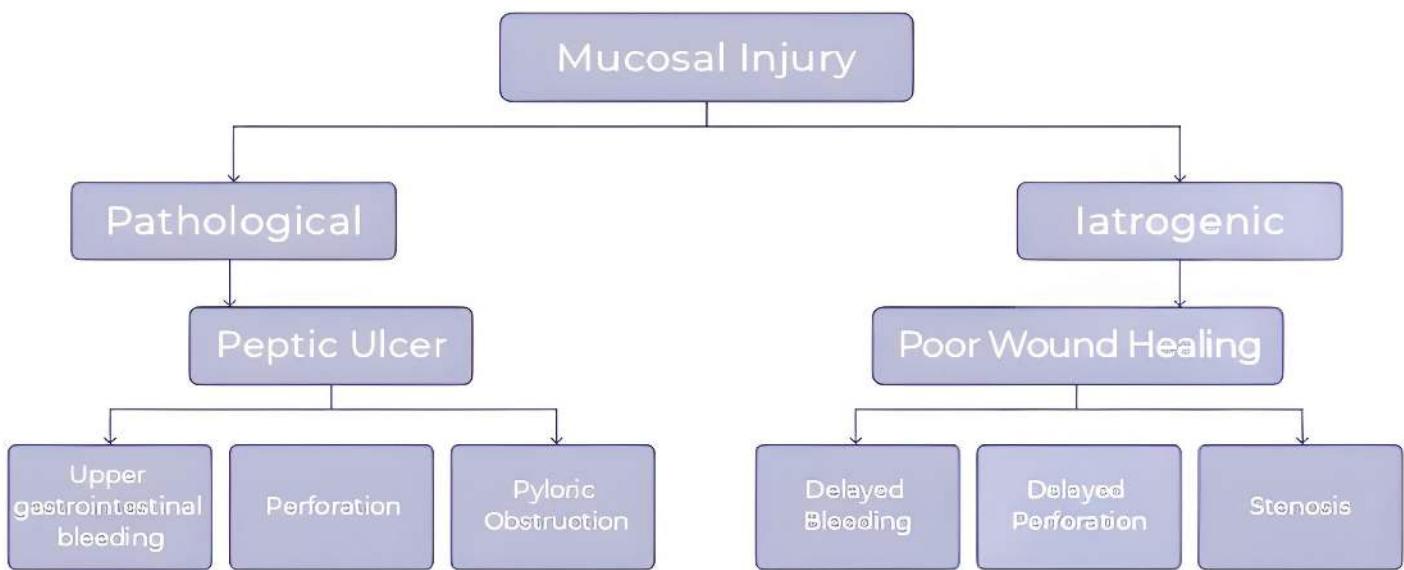
## EASY AND SAFE OPERATION



## COMPETITIVE ADVANTAGES

- The Mucosal Healing Gel is a highly innovative medical product that fills a significant gap in the gastrointestinal market.
- It effectively reduces the occurrence of postoperative complications such as bleeding and perforation after endoscopic surgery, leading to lower treatment costs and improved patient outcomes.
- By promoting wound healing, our product also helps to shortening hospitalization time and reduce hospital costs.
- Improved healing quality and reduced possibility of recurrence or reoperation result in reduced economic burden on patients and healthcare systems.
- By expanding the scope of gastroscope/colonoscopy/esophagoscope surgery, our product helps to improve the level of treatment and popularity of the hospital.
- Reduced physical and mental burden on patients leads to an improved quality of life and increased patient satisfaction.

## WHAT CAN HEALING GEL SOLVE?



## PHYSICAL, CHEMICAL AND SAFETY CHARACTERISTICS OF MUCOSAL HEALING GEL

### Four major physical and chemical characteristics

- Acid-sensitive, the material solidifies when exposed to acid
- Anti-enzymatic, the material is not digested
- Achieve coupling with the wound surface, not easy to fall off
- Convenient for injection and adapting to the wound surface

### Three major safety characteristics

- Non-toxic
- Non-irritating
- Non-allergic

## WOUND PROTECTION MECHANISM      PRODUCT INGREDIENTS

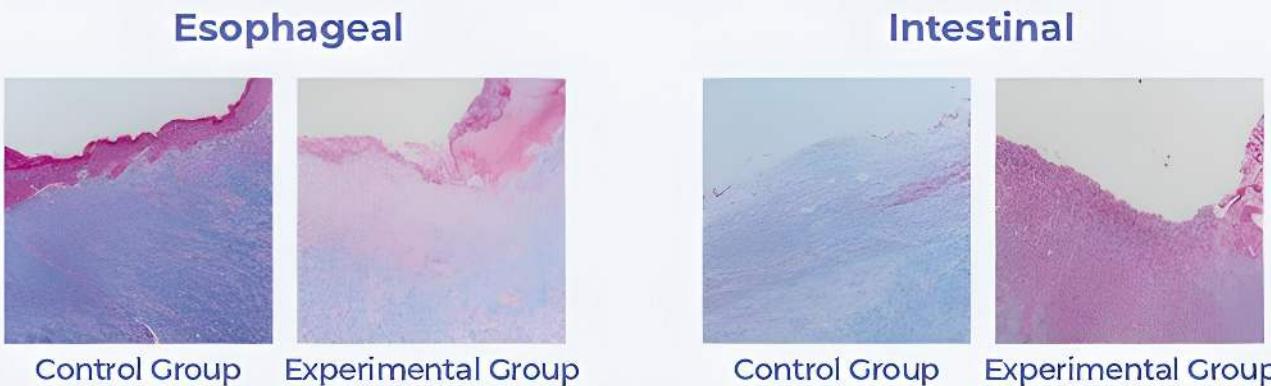
### Quadruple Recovery

-  Protect the wound
-  Promote tissue and cell regeneration
-  Accelerate healing
-  Promote gastrointestinal function recovery

The **Coating Gel** is a viscous, sterile, stain free, odorless liquid containing **sodium alginate**.

The **Firming Gel** is a clear, sterile liquid containing **polylysine** and **purified water**.

# INHIBIT FIBROSIS AND REDUCE SCAR HYPERPLASIA

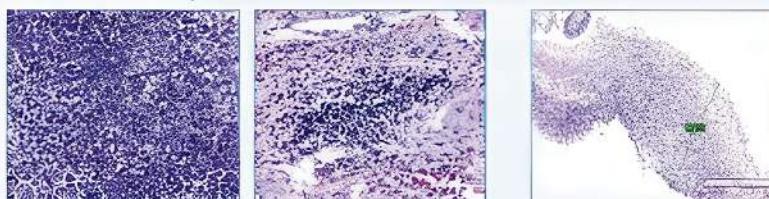


Blue staining represents collagen fibers

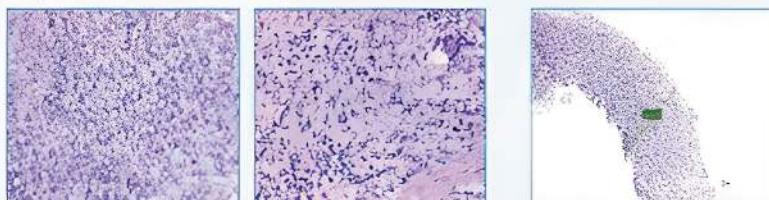
# REDUCE THE INFLAMMATORY RESPONSE AND PROMOTES REGENERATION OF NEW TISSUE

## Inflammatory response of ulcer surface

### Control Group



### Experimental Group



The experimental group showed a mild inflammatory response, while the control group showed necrosis, basal hemorrhage, fibrosis, and aggregation of marginal basal lymphocytes, and the depth of inflammatory infiltration was more in the control group than in the experimental group.

## Promotion of epithelial cell and capillary formation

### Control Group



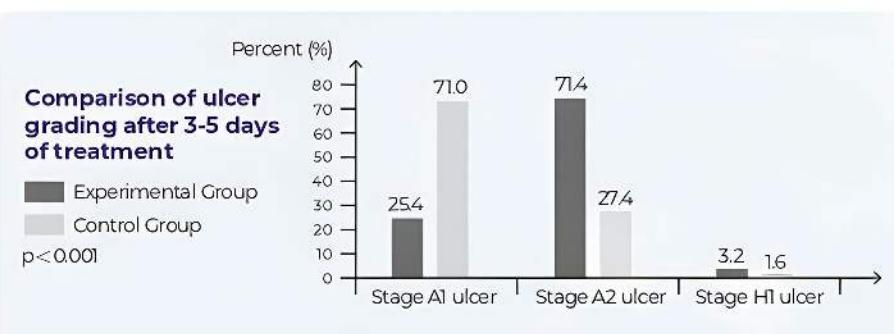
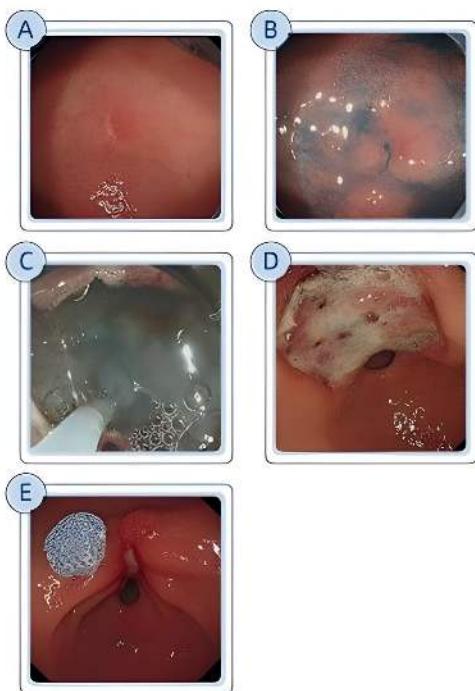
### Experimental Group



There was no marginal mucosal hyperplasia, lymphocyte aggregation, necrosis or hemorrhage in the experimental group compared to the control group.

The experimental group had neatly arranged glandular cells and more CD34(+) perfused vessels.

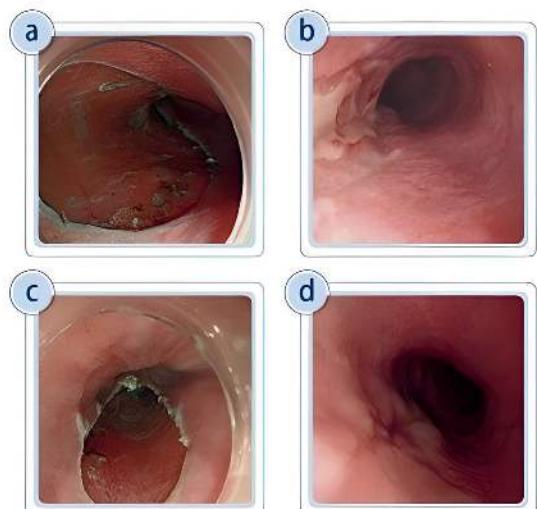
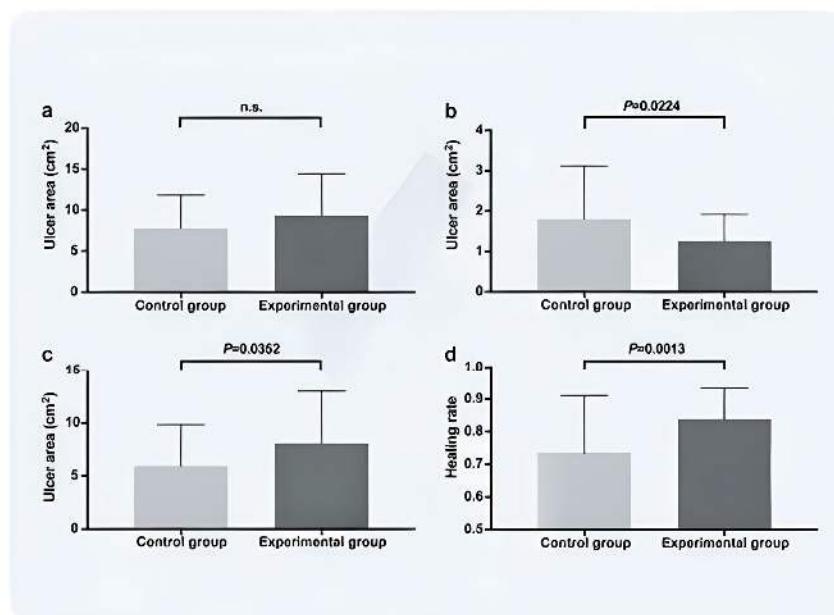
## THE ULCER STAGE AT 3-5 DAYS AFTER ESD SURGERY



### Changes in a gastric antral ulcer induced by ESD with Mucosal Healing Gel.

(A) An early-stage cancer on the lesser curvature of the antrum.  
 (B) Endoscopic image after spraying with indigo-carmine dye.  
 (C) Application of gel to the ulcer immediately after ESD.  
 (D) The ulcer was in the active stage according to the Sakita and Fukutomi classification at 3 days after ESD.  
 (E) The healing stage was observed at 28 days after ESD.

## EFFECTIVENESS OF HEALING GEL IN PROMOTING WOUND HEALING

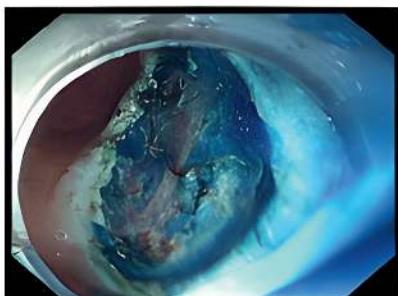


(a) Artificial ulcers left by ESD in the control group.  
 (b) Ulcer conditions in the control group 2 weeks later.  
 (c) Artificial ulcers left by ESD in the experimental group.  
 (d) Ulcer conditions in the experimental group 2 weeks later.

The healing rates were significantly higher in the experimental group than in the control group ( $83.89 \pm 9.51\%$  vs.  $73.28 \pm 17.81\%$ , respectively,  $P=0.0013$ ).

## GASTRIC CLINICAL TRIALS

### Experimental Group



During Surgery  
(4cm\*3.5cm)



3 Days after Surgery  
(2.5cm\*2cm)



28 Days after Surgery  
(Basic Healing)

### Control Group



During Surgery  
(4cm\*3.5cm)



3 Days after Surgery  
(4cm\*3.5cm)



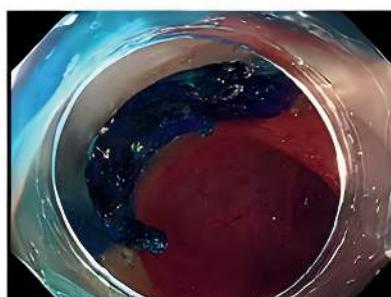
28 Days after Surgery  
(3cm\*2cm)

## INTESTINAL CLINICAL TRIALS

### Experimental Group



Before Surgery

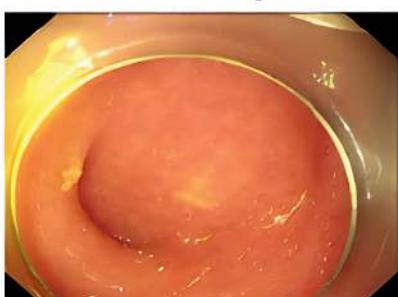


During Surgery  
(4.0cm \* 2.2cm)

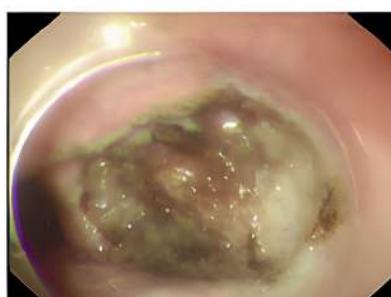


14 days after surgery  
(0.8cm \* 0.2cm)

### Control Group



Before Surgery



During Surgery  
(2.8cm \* 2.5cm)



14 days after surgery  
(2.2cm \* 2.2cm)