Abstract
Microbial biopolymers have recently been introduced as a new material for soil treatment and improvement. Biopolymers provide significant strengthening to soil, even in small quantities (i.e., at 1/10th or less of the required amount of conventional binders, such as cement). In particular, thermo-gelating biopolymers, including agar gum, gellan gum, and xanthan gum, are known to strengthen soils noticeably, even under water-saturated conditions. However, an explicitly detailed examination of the microscopic interactions and strengthening characteristics between gellan gum and soil particles has not yet been performed. In this study, a series of laboratory experiments were performed to evaluate the effect of soil–gellan gum interactions on the strengthening behavior of gellan gum-treated soil mixtures (from sand to clay). The experimental results showed that the strengths of sand–clay mixtures were effectively increased by gellan gum treatment over those of pure sand or clay. The strengthening behavior is attributed to the conglomeration of fine particles as well as to the interconnection of fine and coarse particles, by gellan gum. Gellan gum treatment significantly improved not only inter-particle cohesion but also the friction angle of clay-containing soils.

Keywords Biopolymer · Clay · Cohesion · Direct shear test · Friction angle · Gellan gum · Sand · Shear strength · Vane shear test

1 Introduction
Several studies introducing biologically based geotechnical engineering methods for soil enhancement and treatment have recently been reported. The most common approach involves the intra-soil cultivation of microbes to produce precipitates, such as the microbially induced calcite precipitation (MICP) method [71]. The resulting precipitates then form inter-particle cementation between coarse soil
particles [29, 30, 57, 85]. The MICP method has been widely explored in efforts to enhance the inter-particle cementation, which results in significant cohesion enhancement [23, 27, 53, 59] and hydraulic conductivity control [24, 28] of sandy or silty soils.

Meanwhile, different approaches using microbial excretions (e.g., biopolymers) as a binder material for various soil treatment purposes are now being actively investigated [13, 16–18, 45, 66]. Biopolymers are polymeric biomolecules produced by living organisms, where the monomeric units (i.e., monomers) are covalently bonded [84]. They are commonly used as thickeners or emulsifiers in food and medical products, due to their gel-phase rheological characteristics [51]. Since the introduction of biopolymers in construction engineering [68], their use for geotechnical engineering purposes has also been explored [13, 20, 35, 46, 66]. Recent studies on microbial biopolymers externally produced in culture tanks, such as polysaccharides, have demonstrated remarkable strengthening efficiency in such applications. These results suggest the potential utility of these biopolymers as a new construction material for environmentally friendly geotechnical engineering [5, 13, 14, 18, 19]. Among biopolymers suitable for soil treatment, gel-type biopolymers, such as gellan gum, agar gum, and xanthan gum, have several advantages, including quick (rapid) setting (gelation) [17], the ability to reduce hydraulic conductivity via bio-clogging [21], improving the shear resistance of soils [38, 52] and a unique gel-structure formation process that reveals high gel strength even under fully saturated conditions [22].

Soil type and soil particle composition (i.e., coarse and fine composition) are basic properties that affect soil index parameters, such as Atterberg limits [67, 80], and geotechnical engineering behaviors, such as undrained shear strength [73, 82], soil stiffness and compressibility [12, 15] and hydraulic conductivity [56]. In terms of soil improvement, the presence of clays in soils inhibits the cementation of cement- or lime-mixed soils, due to the expansion of a double layer of clay particles that reduce adhesion between clay particles and cement hydrates [79]. Moreover, high clay content is known to reduce the calcite precipitation efficiency due to the low void ratio and permeability of fine soils that restrict microbe or nutrient solutions transport in soil, thus limiting the application of MICP for real, in situ practices [62, 63].

Among other types of biopolymers, the gel-type biopolymers show better inter-particle interactions with clay particles than with coarse particles, such as sand [16, 17]. The interactions between gel-type biopolymers and clay particles result in greater strengthening due to direct ionic or hydrogen bonding and matrix formation, while neutral sand particles undergo no direct bonding with biopolymers. For sand, gellan gum biopolymer provides strengthening to gellan gum–sand mixtures due to the condensation and aggregation effects of the high tensile gellan gum hydrogels among the sand particles [21]. However, a detailed understanding of the interactions between gellan gum and soils and their strengthening behavior, especially for sand–clay mixtures with different clay contents, has yet to be achieved. Thus, the present study was conducted to investigate the inter-particle interactions and strengthening behavior of gellan gum-treated soils, ranging from sand to clays, through a series of laboratory tests.

2 Materials and methods

2.1 Gellan gum-treated soil mixtures

2.1.1 Gellan gum biopolymer

Gellan gum is a high molecular weight polysaccharide group biopolymer produced by the bacterium *Sphingomonas elodea*. The most notable characteristic of gellan gum is its thermo-gelation property: Its solubility and viscosity are temperature dependent. Gellan gum exhibits poor solubility at low temperature, while sufficient gum dissolution with random coil conformation forms uniform gellan gum solutions at temperatures higher than 90 °C [39]. Upon temperature decrease, randomly dispersed gellan gum coils adopt a double-helical structure through ionotropic sol–gel transition, which results in high viscous and firm gellan gum gel formation [26, 61]. Details of the gellan gum rheology and a case study of experimental implementations can be found in Chang et al. [17, 22]. Low acyl gellan gum (CAS No: 71010-52-1) was used in this study to verify the interaction between soils and gellan gum, without the presence of additional cations, which alter the gelation and strength of gellan gum gels.

2.1.2 Sand–clay mixtures

*Jumunjin* sand, a standard sand in Korea, was used for the experimental tests in this study. *Jumunjin* sand is classified as poorly graded sand (SP) according to the USCS classification, with soil properties of $D_{50} = 0.52$ mm, $C_v = 1.35$, $C_c = 1.14$, $G_s = 2.65$, $e_{\text{min}} = 0.644$, and $e_{\text{max}} = 0.892$.

For clay, *Bintang* kaolin (Belitung Island, Indonesia), a commercially available white kaolin clay material, was used. *Bintang* kaolin is classified as *CH* according to the USCS classification, having soil properties of $PL = 24$, $LL = 62$, $G_s = 2.65$, and $D_{50} = 44$ μm.

Clean sand and clay were dried in an oven before specimen preparation. Different sand–clay mixtures were
obtained by mixing dry sand and clay at the mass ratios (i.e., \( m_c / m_s \); clay-to-soil ratio in mass, where \( m_c / m_s = 1.0 \) indicates pure clay) listed in Table 1. The liquid limits of the prepared sand–clay mixtures (except pure sand) were evaluated via a fall cone test using a 30°C, 80 g British cone [7, 9]. The liquid limit of the sand–clay mixtures was \( LL = 17 \) at \( m_c / m_s = 0.2 \) and was gradually increased to 40 and 62 for \( m_c / m_s = 0.5 \) and 1.0 soils, respectively (Table 1).

Initial water content (\( w \)) values for gellan gum–soil mixtures were determined by considering both clay (\( m_c / m_s \)) and gellan gum content (\( m_b / m_s \)) to the total weight of soil (i.e., higher \( w \) values with higher clay and gellan gum contents). For example, the initial water content of the gellan gum–soil mixtures should be higher than the LL of untreated (natural) soils to provide thorough mixing due to the hydrophilic water holding and retention capacity of biopolymers [10], which increases the LL of biopolymer–soil mixtures [14]. Thus, the \( w \) values have been fixed as 60, 50, 40, and 30% for \( m_c / m_s = 1.0, 0.5, 0.2, \) and 0 soils, respectively (Table 2).

### 2.1.3 Preparation of gellan gum-treated soils

To prepare thermo-gelated gellan gum-treated soil mixtures, dry gellan gum was first dissolved and hydrated in pure water heated to 120°C, to obtain a \( m_b / m_w = 5\% \) gellan gum solution (\( m_b / m_w \) = gellan gum content to the mass of water). Thereafter, a dry sand–clay mixture and the hot \( m_b / m_w = 5\% \) gellan gum solution were uniformly mixed at \( m_w / m_s \) (gellan gum solution to the mass of soil ratio) = 20%, resulting in thermo-gelated \( m_b / m_s = 1\% \) gellan gum-treated soil with initial \( w = 20\% \). Additional heated distilled water was added and mixed thoroughly to achieve the desired initial water content, as listed in Table 2 (e.g., \( w = 30, 40, 50, \) and 60% for \( m_c / m_s = 0.5 \) soil).

For laboratory vane shear tests, the gellan gum-treated soil mixtures (\( m_b / m_s = 1\% \)) were poured into a rectangular cup made of acryl having inner dimensions of 50 mm width, 50 mm length, and 50 mm depth. The thermo-gelated specimens were tightly sealed with laboratory vacuum packing to prevent surface evaporation. Temperature change was measured via a radiation thermometer to verify complete cooling (at room temperature) of the thermo-gelated gellan gum-treated soils.

| Test | Dimensions (mm) | Soil (%) | Gellan gum (%) | Initial water content (\( w \))a | Specimensb | Condition |
|------|----------------|----------|---------------|-------------------------------|-----------|-----------|
|      |                | Sand     | Clay          | \( m_b / m_s \) | Contenta | Treatment | specimensb |                |
| Laboratory vane shear (I) | 50 \( \times \) 50 \( \times \) 50 (cup) | 0 100 1.0 | 1 | Untreateda and thermal | 40, 55, 65 | 3 EA | Immediate (without drying) |
|      |                | 50        | 50           | 0.5               | 0, 0.5, 1, 2, 3, 4, 5 | Thermal | 40 | 20, 30, 40, 50, 60 |
|      |                | 20        | 0.2          | 2, 2.5            | 0, 0.5, 1, 1.5, 2.25 | 40 | 20, 30, 40, 50 |
| Laboratory vane shear (II) | 50 \( \times \) 50 \( \times \) 50 (cup) | 0 100 1.0 | 0.5           | 2, 2.5            | 1, 2, 3, 4, 5 | 60 | 50 |
|      |                | 50        | 0.2          | 0, 0.5, 1, 1.5, 2, 2.5 | 0.5, 1, 1.5, 2, 2.5 | 40 | 50 |
|      |                | 20        | 0.8          | 1                 | 0.2, 0.4, 0.6, 0.8, 1 | 40 | 30 |
| Direct shear | Diameter 60; Height 20 | 0 100 1.0 | 0.2 | 1, 2, 3, 4, 5 | 1, 2, 3, 4, 5 | 30 | 40 |

| Test | Dimensions (mm) | Soil (%) | Gellan gum (%) | Initial water content (\( w \))a | Specimensb | Condition |
|------|----------------|----------|---------------|-------------------------------|-----------|-----------|
|      |                | Sand     | Clay          | \( m_b / m_s \) | Contenta | Treatment | specimensb |                |
|      |                | 0 100 1.0 | 1 | Untreateda and thermal | 40, 55, 65 | 3 EA | Immediate (without drying) |
|      |                | 50        | 50           | 0.5               | 0, 0.5, 1, 2, 3, 4, 5 | Thermal | 40 | 20, 30, 40, 50, 60 |
|      |                | 20        | 0.2          | 2, 2.5            | 0, 0.5, 1, 1.5, 2.25 | 40 | 20, 30, 40, 50, 60 |
|      |                | 0 100 1.0 | 0.5           | 2, 2.5            | 1, 2, 3, 4, 5 | 60 | 50 |
|      |                | 50        | 0.2          | 0, 0.5, 1, 1.5, 2, 2.5 | 0.5, 1, 1.5, 2, 2.5 | 40 | 50 |
|      |                | 20        | 0.8          | 1                 | 0.2, 0.4, 0.6, 0.8, 1 | 40 | 30 |

| Test | Dimensions (mm) | Soil (%) | Gellan gum (%) | Initial water content (\( w \))a | Specimensb | Condition |
|------|----------------|----------|---------------|-------------------------------|-----------|-----------|
|      |                | Sand     | Clay          | \( m_b / m_s \) | Contenta | Treatment | specimensb |                |
|      |                | 0 100 1.0 | 1 | Untreateda and thermal | 40, 55, 65 | 3 EA | Immediate (without drying) |
|      |                | 50        | 50           | 0.5               | 0, 0.5, 1, 2, 3, 4, 5 | Thermal | 40 | 20, 30, 40, 50, 60 |
|      |                | 20        | 0.2          | 2, 2.5            | 0, 0.5, 1, 1.5, 2.25 | 40 | 20, 30, 40, 50, 60 |
|      |                | 0 100 1.0 | 0.5           | 2, 2.5            | 1, 2, 3, 4, 5 | 60 | 50 |
|      |                | 50        | 0.2          | 0, 0.5, 1, 1.5, 2, 2.5 | 0.5, 1, 1.5, 2, 2.5 | 40 | 50 |
|      |                | 20        | 0.8          | 1                 | 0.2, 0.4, 0.6, 0.8, 1 | 40 | 30 |

| Test | Dimensions (mm) | Soil (%) | Gellan gum (%) | Initial water content (\( w \))a | Specimensb | Condition |
|------|----------------|----------|---------------|-------------------------------|-----------|-----------|
|      |                | Sand     | Clay          | \( m_b / m_s \) | Contenta | Treatment | specimensb |                |
|      |                | 0 100 1.0 | 1 | Untreateda and thermal | 40, 55, 65 | 3 EA | Immediate (without drying) |
|      |                | 50        | 50           | 0.5               | 0, 0.5, 1, 2, 3, 4, 5 | Thermal | 40 | 20, 30, 40, 50, 60 |
|      |                | 20        | 0.2          | 2, 2.5            | 0, 0.5, 1, 1.5, 2.25 | 40 | 20, 30, 40, 50, 60 |
|      |                | 0 100 1.0 | 0.5           | 2, 2.5            | 1, 2, 3, 4, 5 | 60 | 50 |
|      |                | 50        | 0.2          | 0, 0.5, 1, 1.5, 2, 2.5 | 0.5, 1, 1.5, 2, 2.5 | 40 | 50 |
|      |                | 20        | 0.8          | 1                 | 0.2, 0.4, 0.6, 0.8, 1 | 40 | 30 |

a Water content (percent ratio to the mass of soil) at the moment of specimen preparation. The actual water content at vane shear measurements (Fig. 1) can differ slightly from the initial water content due to time delays

b Number of measurements for a single condition (on average)

c Natural soil without any biopolymer treatment
For direct shear tests, the thermo-gelated gellan gum-treated soil mixtures were molded into disk shapes 60 mm in diameter and 20 mm in height. The specimens were submerged in water and cooled to maintain the initial conditions of each gellan gum-treated soil specimen.

2.2 Laboratory vane shear tests

In general, vane shear tests are used to determine soil surface shear strengths for near-zero effective confinements [8], where vane shear tests are widely performed to measure the undrained shear strength of soils in the laboratory and in situ [11].

In the present study, primary laboratory vane shear tests (I) were performed using a 12.7-mm rectangular vane (vane thickness \( t = 0.5 \text{ mm} \); perimeter ratio \( x = 5\% \)) [6] on both untreated and gellan gum-treated (with and without thermo-gelation, respectively) soil specimens, which had different soil compositions and initial water contents. Although the laboratory vane shear test presents problems with organic soils due to uncertain failure conditions around the vane circumference [50], three specimens were tested and averaged to represent a single soil condition with minimum uncertainty. The vane was pushed 30 mm into the soil from the top surface to place the blade directly in the middle of the specimens and then rotated constantly with a 10°/min angular velocity [36].

Secondary laboratory vane shear tests (II) were conducted to evaluate the effect of the gellan gum–clay interaction (i.e., \( m_b/m_c \); gellan gum-to-clay ratio in mass) on the vane shear strength behavior of the thermo-gelated gellan gum-treated soils, where \( m_b/m_c = 0.5 \) and 1.0 soils at a water content of \( w = 40\% \) were considered. The ratio \( m_b/m_c \) was maintained at 1, 2, 3, 4, or 5% for pure clays (\( m_c/m_s = 1.0 \)), while it was reduced to half those amounts for \( m_b/m_c = 0.5 \) soils, respectively, to obtain identical \( m_b/m_c \) conditions of 1–5% for both soil compositions. The aforementioned experimental procedures of soil mixing, molding, cooling, and strength measurement used for the primary laboratory vane shear tests were also adopted.

Because a 12.7×12.7 mm vane was applied, the obtained laboratory vane shear strength (\( \tau_v \)) values were \( \tau_v = M/4.29 \) (kPa), where \( M \) (N mm) is the measured torque at failure. Rod friction [76] and shear rate [3] effects were neglected due to the small penetration depth of 30 mm and the low peripheral velocity of 1.1 mm/min.

2.3 Direct shear test

The direct shear test is appropriate for evaluating qualitative stress–strain behaviors along a thin zone of shear failure, especially for granular materials where drainage is not a consideration [37, 49]. Moreover, a relatively small-scale circular-type direct shear apparatus of 60 mm in diameter has substantial advantages, such as the ability to provide consistent and reliable evaluation of strength parameters, including \( \phi_d \) and \( c_d \), as demonstrated by numerous measurements [65].

Soils with \( m_b/m_s = 1.0, 0.5, 0.2, \) or 0 (i.e., pure sand) were mixed with thermo-gelated gellan gum, as summarized in Table 2. Gellan gum-treated soils were prepared with gellan gum-to-clay ratios (\( m_b/m_c \)) of 1, 2, 3, 4, or 5% at different initial water contents (e.g., 60% for pure clay, 30% for pure sand).

Gellan gum-treated soil specimens were placed in a circular shear box having a 60-mm inner diameter and 25 mm height (HM-2701.60D). Porous stones were placed above and beneath the sample. The shear box was filled with water to saturate the gellan gum-treated soil and prevent any moisture loss from the specimen during testing. The initial dry densities before loading were 0.96, 1.23, 1.62, and 1.42 g/cm\(^3\) on average for \( m_b/m_s = 1.0, 0.5, 0.2, \) and 0 soils, respectively. The target overburden stress of \( \sigma_v = 50, 100, 200, \) and 400 kPa was applied via a pneumatic actuator for 24 h to ensure that there was no

![Fig. 1 Laboratory vane shear test results of \( m_b/m_s = 1\% \) gellan gum-treated soils with various water contents](image-url)
further vertical displacement before the horizontal shear displacement test.

A direct shear testing device (Humboldt HM-2560A) was used to perform direct shear tests. Horizontal shear displacement was applied at a rate of 0.01 mm/min for 10-mm total horizontal displacement [2]. For biopolymer-treated soils, the elapsed time of horizontal shearing must be at least 49250 s due to the high water holding capacity and low coefficient of consolidation ($C_v$) values of biopolymer-treated clays (i.e., $C_v = 0.2 \times 10^{-7}$ m$^2$/s, at void ratio = 0.9 for beta-glucan-treated clay) [14, 58]. However, although horizontal shearing has been applied slowly, it is hard to present a perfect drained condition during a direct shear test on gel-type biopolymer-treated soils due to the high water holding capacity of gellan gum [17, 21]. Thus, the direct shear test results from this study have been interpreted in terms of direct shear strength, direct shear dilation angle, direct shear cohesion and direct shear friction angle where notations become $\tau_v$, $\psi_d$, $c_d$, and $\phi_d$, respectively.

Shear load and vertical and shear displacements were measured automatically via a load cell (HM-2300.020) and an LVDT (HM-14368, HM-14180). Specimen preparation and direct shear testing were performed at the same room temperature (20 °C) to avoid viscosity variation effects of gellan gum gels with temperature change. At least three specimens were tested to obtain a representative value for each condition depending on $m_s/m_c$, $m_b/m_c$, and $\sigma_v$.

### 2.4 Scanning electron microscope (SEM) images

SEM images were taken to observe the microscale interactions between gellan gum and soil particles for 1% gellan gum-treated soils ($m_s/m_c = 1.0$ and 0.5). The undisturbed condition was investigated by sampling a 0.5-cm$^3$ piece of a cube (10 mm wide, 10 mm long, 5 mm high) of dried gellan gum-treated soil that had not been subjected to any testing. The disturbed condition was investigated by collecting fragments of 1% gellan gum-treated soil that had been crushed during unconfined compression tests. Both undisturbed and disturbed sample pieces were attached to a 25-mm-diameter SEM mount by carbon conductive tabs (PELCO Tabs$^\text{TM}$). Carbon paint (DAG-T-502) was applied to the sample edges and bottoms to provide sufficient grounding. Specimens were coated via an osmium plasma coater (OPC-60A) using osmium tetroxide (OsO4) as the source of osmium, for 20 s under vacuum. An extreme high-resolution SEM (FEI Magellan 400L XHR) was used to observe the microscopic structure and particle alignment of the gellan gum-treated soil samples.

### 3 Results and analysis

#### 3.1 Laboratory vane shear strength ($\tau_v$)

#### 3.1.1 1% gellan gum-treated soils for various initial water contents

The results for dry density and laboratory vane shear strength of the $m_s/m_c = 1$% gellan gum-treated soils at various water contents are shown in Fig. 1. The dry densities of both untreated and gellan gum-treated soils showed a single trend with water content regardless of the gellan gum treatment (Fig. 1a). The vane shear strength ($\tau_v$) increased with a reduction in the initial water content and an increase in the clay content (Fig. 1b). Gellan gum treatment increased the shear strength significantly for all soil mixtures, even with high water content.

In general, gellan gum forms uniform hydrocolloids and transforms into firm hydrogels via thermo-gelation upon cooling, and the resulting hydrogels have extremely high water holding capacity (i.e., absorbability) [61]. As a result, the overall shear strengths of the gellan gum-treated soils remain much higher than those of untreated soils, even above the LL values (Table 1) of the untreated condition, and also decrease gradually with increasing water content. Moreover, soils with lower $m_c/m_s$ show higher strengthening efficiency (i.e., the $\tau_v$ of 1% gellan gum-treated soil/ $\tau_v$ of untreated soil), which implies that the gellan gum-to-clay ratio in mass ($m_b/m_c$) is a possible dominant parameter influencing the strengthening behavior of gellan gum-treated soils.

#### 3.1.2 Different percent gellan gum-treated soils with the same initial water content

The secondary laboratory vane shear test results for different percent gellan gum-treated soils ($m_s/m_c = 0.5$ and 1.0) at a similar dry density = 1.15 g/cm$^3$ ($w = 40\%$) are shown in Fig. 2. The vane shear strength ($\tau_v$) values of both soils increase with higher gellan gum content ($m_b/m_c$) and converge after certain $m_s/m_c$ conditions are met (Fig. 2a). From the viewpoint of clay content, both soils show an upper limit of shear strength at $m_b/m_c = 4\%$ (Fig. 2b), which implies that the strengthening behavior is dominated by the gellan gum–kaolinite clay matrix formed via thermo-gelation.

Hydrated biopolymers are known to be effective coagulants for clay particles [74, 75], while electrically neutral sand particles (e.g., those with ionic- or hydrogen bonding) have no direct interaction with biopolymers [5, 16]. For this reason, the interaction between gellan gum and clay particles can be facilitated by lowering the $m_c/m_s$ values,
which results in higher $m_c/m_s$. The vane shear strength of soils thus can be effectively improved by gellan gum treatment, changing the soil composition and optimizing the gellan gum-to-clay ratio.

3.2 Shear stress and vertical strain behaviors with shear displacement

Typical direct shear test results of gellan gum-treated soils are shown in Fig. 3. The vertical strain ($\varepsilon_v$) values were calculated from the changes in the height of the sample relative to the initial height of the sample (20 mm). Negative values indicate volumetric contraction. The maximum direct shear dilation angle ($\psi_d$) values were estimated from the steepest slope ($\Delta$vertical displacement/$\Delta$horizontal displacement) of shear displacement—vertical strain curves [42, 54], as shown in Fig. 3b, d, f.

Figure 3a, b presents the shear behavior of untreated ($m_b/m_s = 0\%$) natural soils at an overburden stress of $\sigma_\text{v} = 50$ kPa. Differences between the peak shear strength and the residual shear strength increased with lower clay content, whereas the volumetric change behavior became more dilative with the same clay content. In detail, untreated soils start to show peak $\tau_t$ and $\psi_d$ values for sand contents of 80% and 100%. When the weight of the sand in the sand–clay mixtures is higher than 75%, the shear behavior of the sand–clay mixtures is governed mainly by the frictional resistance between sand grains [83]. Thus, soils containing 80% sand appear to follow the shear behavior of sand grains (with peaks), while soils of $m_c/m_s = 0.5$ and 1.0 show strain-hardening behaviors.

Figure 3c, d shows the shear behavior of gellan gum-treated soils (clay = 20% and $m_b/m_s = 1\%$) under different overburden stress conditions. Gellan gum treatment instantly increased the $\tau_t$ and $\psi_d$ of the soils (e.g., comparison of 50-kPa overburden stress results). As the overburden stress increased, $\tau_t$ of the gellan gum-treated soils increased (Fig. 3c), while the soils become less dilative (Fig. 3d).

Figure 3e, f reveals the shear behavior of different percent gellan gum-treated soils ($m_c/m_s = 0.5$ and an overburden stress of 50 kPa). As the $m_b/m_s$ ratio was increased (i.e., higher gellan gum content), the $\tau_t$ and $\psi_d$ increased, and thus, the soil showed more brittle and dilative behavior. It was found that the gellan gum treatment induced a structural agglomeration effect (i.e., forming gellan gum–clay matrices), which is similar to cementation. This finding is consistent with the general $\tau_t$ and $\psi_d$ increments observed with cemented [4, 43] and polymer-treated [40] soils.

3.3 Shear strength and overburden stress relationships

The direct shear strengths of thermo-gelated gellan gum-treated soils are shown in Fig. 4. Overall, the direct shear strength increases linearly with the increase in overburden stress, regardless of the soil composition and gellan gum treatment. The soil composition affects the strength of untreated (i.e., natural) soils: the $\tau_t$ of untreated soils decreases with increasing $m_c/m_s$, i.e., $\tau_t = 9.3^\circ$ for pure sand $\rightarrow$ $\phi_d = 8.7^\circ$ for pure clay. Meanwhile, gellan gum treatment increases the strength of the soils significantly, with $\phi_d$ and $c_d$ increases with increasing $m_b/m_s$ or $m_b/m_c$.

As shown in Fig. 4a, the $\phi_d$ of pure sand is negligibly affected by gellan gum treatment, whereas the $c_d$ is substantially increased with increasing gellan gum content. Cement treatment on sand is known to accompany increases of $c_d$, while there are opposing views on the $\phi_d$ behavior of cement–sand mixtures. Some studies indicate an increase in $\phi_d$ with cementation [25, 47, 77], contrasting with results showing a constant $\phi_d$ regardless of cement-treated sands with a high degree of compaction ($D_c$) [32, 44, 72]. However, the $\phi_d$ of gellan gum-treated sand
remains constant, regardless of the gellan gum content and $D_r$ of the specimens in this study. Cement-treated sand consists of a solid phase (sand grains and C-S-H hydrates) only after cement hydration, while gellan gum-treated sand contains solid grains and hydrogels in voids. The presence of gellan gum hydrogels in sand pores only affects $c_d$ depending on the gel concentration, without any effect on the mechanical behavior ($\phi_d, \phi_u$) of sand [21]. Thus, the shear behavior of gellan gum-treated sand should be interpreted differently than that of cement-treated sands.

Meanwhile, clay-containing soils show a remarkable increase in $\phi_d$ as well as $c_d$ with a sequential increase in $m_b/m_c$ (Fig. 4b, c, d), which is consistent with the general behavior of cement-treated clays [55, 70, 78]. The presence of a small amount of gellan gum ($m_b/m_c = 1\%$) in clay-containing soils immediately induced a remarkable increase in $\phi_d$, whereas $\phi_d$ gradually increases with higher $m_b/m_c$ ($>1\%$) conditions which implies that the gellan gum treatment not only contributes to the increase in $c_d$ but also contributes to the conglomeration of fine particles (i.e., clay) to form partially conglomerated aggregates, resulting in an increase in dilatancy and $\phi_u$, respectively.

Meanwhile, the $c_d$ of gellan gum-treated soils increased consistently with increasing $m_b/m_c$ or $m_b/m_s$ ratios. Gellan gum treatment plays an important role in $c_d$ enhancement, regardless of soil type. The $c_d$ value of gellan gum-treated soils is highly correlated with the gellan gum–clay matrix ($m_b/m_c$), which shows a similar increment in $c_d$ to the subsequent increase in $m_b/m_c$.

In general, an increase in $\phi_d$ induces higher shear strength and bearing capacity at greater depths, while the increase in $c_d$ becomes meaningful for subsurface soil strengthening. Thus, it can be concluded that gellan gum treatment can be effective for improving the shear strength of shallow subsurface sand layers, such as seashore or arid region dunes, by increasing $c_d$ (Fig. 4a). It also may be effective in improving the shear strength of deep clayey soil deposits via $\phi_d$ as well as $c_d$ increment (Fig. 4d), which results in a remarkable increase in the bearing capacity of soils. Moreover, the significant increase in shear strength for gellan gum-treated sandy soils (Fig. 4a) implies the potential value of applying gellan gum in geotechnical aseismic practices, such as increasing resistance to liquefaction. Specifically, this shear strength increase results from improved residual shear strength and shear stiffness, as shown in Fig. 3a, c, e, which enhances the post-liquefaction parameters of sands at shallow depths [1, 31, 86].

4 Discussion

4.1 Shear strength parameters of gellan gum-treated soils

Figure 5 presents the direct shear strength ($\tau_d$) of gellan gum-treated soils with various $m_b/m_c$ ratios at different overburden stress ($\sigma_v$) levels. The direct shear strength of soils with less clay content increases more rapidly with an
increase in the \( \frac{m_b}{m_s} \) ratio. That is, the direct shear strength of soils with large amounts of clay or pure sand increases gradually with an increasing \( \frac{m_b}{m_s} \) ratio. All the gellan gum-treated soil mixtures were prepared with \( \frac{m_b}{m_c} \) ranging from 0 to 5%, and the highest \( s_f \) value was exhibited at lower \( \frac{m_b}{m_c} \) conditions, for soils with lower clay content.

Regarding the gellan gum to clay \( (\frac{m_b}{m_c} = \frac{m_b/\text{m_c}}{m_c/\text{m_s}}) \) composition, gellan gum-treated soils reached a maximum \( s_f \) at \( m_b/m_c = 4% \) regardless of the \( m_c/m_s \) and \( r_v \) (Fig. 5a, b), consistent with the laboratory vane shear strength \( (s_v) \) behavior shown in Fig. 2b. The \( \tau_c \) values of gellan gum-treated soils increased with lower clay content under the same \( m_b/m_c \) conditions, as shown in Fig. 5c, d which implies the importance of the physical interlocking between granular grains on the shear strength behavior when the intergranular gellan gum–clay matrices are identical (i.e., with the same \( m_b/m_c \)). Meanwhile, as expected, the \( \tau_c \) of pure sand was generally lower than that of clay-containing soils, due to the lack of a gellan gum-clay matrix.

Figure 6 shows the shear strength parameters \( (c_d \) and \( \phi_d) \) of gellan gum-treated soils with varying \( m_b/m_c \) contents. As shown in Fig. 6a, the \( c_d \) values of gellan gum-treated soils increase sequentially with increasing \( m_b/m_c \) and reach their maximum value at around \( m_b/m_c = 4\% \). The decrease in \( c_d \) at \( m_b/m_c > 4\% \) appears to be affected by surplus gellan monomers, which can cause ionic repulsion or hydrologic swelling, resulting in a reduction of gellan gum–clay interaction. Nevertheless, it can be cautiously predicted that clay materials with larger specific surfaces (e.g., montmorillonite) should show maximum strengths at \( m_b/m_c \) values higher than 4%.

Meanwhile, as shown in Fig. 6b, the \( \phi_d \) values of the gellan gum-treated sand (clay 0%) have less dependency on changes in \( m_b/m_c \), while the \( \phi_d \) values of the gellan gum-treated clayey soils increase abruptly with the presence of a small amount of gellan gum (i.e., \( m_b/m_c = 1\%) \) and subsequently converge or slightly increase with increasing \( m_b/m_c \). Therefore, the addition of even a small amount of gellan gum can effectively improve the \( \phi_d \) of clayey soils. The larger \( \phi_d \) improvement effect observed.

![Fig. 4 Direct shear test results of gellan gum-treated (thermo-gelated) soils on \( \sigma_v-\tau_c \) planes](image-url)
with \( m_b/m_s = 0.2 \) and 0.5 soils indicates that not only the inter-granular friction of sand grains, but also the conglomerated gellan gum–clay matrix inside the inter-granular pores, or between sand particles, significantly governs the overall friction behavior (including significant \( \psi_d \) increase; Fig. 3f) of gellan gum-treated soils.

With regard to soil composition, gellan gum treatment of single-grained soils (pure sand and pure clay) appears to be more appropriate for increasing \( c_d \) (Fig. 6a), while gellan gum treatment of multi-grained soils (\( m_b/m_s = 0.2 \) and 0.5) shows significant increases in \( \phi_d \) (Fig. 6b). However, it should be noted that the \( c_d \) and \( \phi_d \) values shown in Fig. 6 are presented with variations in \( m_b/m_c \) instead of \( m_b/m_s \). This means that the absolute quantity of gellan gum becomes less at lower \( m_b/m_c \) ratios in the order 1.0 → 0.5 → 0.2. For instance, \( m_b/m_c = 5, 2, \) and 1% exhibit \( m_b/m_s = 1\% \) conditions identical to \( m_b/m_s = 0.2, 0.5, \) and 1.0 soils, respectively.

The \( c_d \) and \( \phi_d \) of gellan gum-treated soils are presented with \( m_b/m_s \) variations in Fig. 7. \( c_d \) (Fig. 7a) shows a strong correlation with \( m_b/m_c \), regardless of soil composition, while the \( \phi_d \) (Fig. 7b) becomes relatively higher for multi-grained soils, regardless of \( m_b/m_s \) conditions. The \( \phi_d \) of pure sand remains almost constant with \( m_b/m_s \) variation, while the \( \phi_d \) of pure clay becomes almost identical to the \( \phi_d \) of pure sand at high \( m_b/m_s \) conditions. \( \phi_d \) values of sand–clay mixtures become higher than those of single-grained soils, which can be ascribed to the effect of the conglomeration between gellan gum–clay matrices and sand particles. Thus, it can be concluded that the \( c_d \) characteristic of gellan gum-treated soils is strongly affected by the ratio of the absolute quantity of gellan gum to the amount of soil (\( m_b/m_s \)), rather than to the soil composition and \( m_b/m_c \) matrices; meanwhile, the \( \phi_d \) characteristic of gellan gum-treated soils is governed by the multi-grain composition and \( m_b/m_c \) variation rather than the \( m_b/m_s \) ratio.
4.2 Microscopic interaction and shear model of gellan gum-treated soils

Figure 8 shows SEM images of 1% gellan gum-treated soils with \( m_b/m_s = 1.0 \) and 0.5. Clay particles and gellan gum monomers form a firm and well-bonded gellan gum–clay matrix via hydrogen bonding [17, 48, 64], as shown in Fig. 8a–d. Gellan gum and gellan gum–clay matrices show no direct interaction (e.g., hydrogen or ionic bonds) with sand particles for both undisturbed (Fig. 8e) and disturbed conditions (Fig. 8f). Thus, it appears that the gellan gum–clay matrix dominates the strengthening behaviors of gellan gum-treated soils.

For pure clay, gellan gum chains, with average lengths of 100 nm, can form connections between kaolinite particle edges, inducing an accumulation of stacks of kaolinite particles (Fig. 8a). Because gellan gum is an anionic biopolymer [61], gellan gum monomers are expected to attach to positively charged kaolinite particle edges and enhance plate particle stacking. Inter-particle bonds can be broken or detached when gellan gum-treated soils are subjected to macro-strain disturbances, such as crushing by unconfined compression tests (Fig. 8b). Therefore, the strengthening behavior of gellan gum-treated soils can be explained as a combination of the following: (1) the formation of a gellan gum gel or gellan gum–clay matrix, which immediately enhances \( c_d \); (2) a conglomeration effect induced by gellan gum–clay matrices, which instantly increases dilatancy and \( \phi_d \); and (3) higher strength with higher water content due to the LL increase [14] induced by hydrophilic biopolymer treatment.

Figure 9 shows schematic models of the microstructural evolution of gellan gum-treated soils during direct shear tests. For pure sand (Fig. 9a), gellan gum gels are condensed among sand grains and thus do not affect \( \phi_d \) but do provide a significant contribution to \( c_d \) during shear, as already shown in Fig. 4(a). However, the \( \phi_d \) will increase when the gellan gum gels are fully dried, by forming a surface coating and enlarging inter-particle contact areas [21].

For pure clay (Fig. 9b), gellan gum gels bond with themselves and clay particles. They thereby facilitate bonding among clay particles and conglomeration of clay particles during shear, increasing both the \( \phi_d \) and \( c_d \). The \( \phi_d \) of gellan gum-treated pure clay increases due to the restricted slippage, which is governed by the high tensile gellan gum chains connecting clay particles. Gellan gum...
also provides a significant improvement in $c_d$ (Fig. 6a). Although natural clay generally contracts with shearing, gellan gum-treated pure clay shows dilative behavior during direct shearing due to the conglomeration induced by the formation of gellan gum–clay matrices (Fig. 3b).

For sand–clay mixtures (Fig. 9c), gellan gum gels enhance interactions between clay particles and between sand grains, producing an instant increase in $c_d$. Moreover, the conglomeration of the gellan gum–clay and gellan gum–clay–sand matrices induces a considerably higher $\phi_d$, causing accumulated gellan gum–clay matrices to behave as secondary grains between sand particles. Thus, the $\phi_d$ of gellan gum-treated sand–clay mixtures becomes remarkably higher than that of single grain soils (pure sand or pure clay) (Fig. 6b) and dominates the shear behavior of sand–clay mixtures.

4.3 Potential of gellan gum treatment in geotechnical engineering practices

The results of this study show that $m_c/m_s$ becomes the dominant parameter governing the strengthening behavior of gellan gum-treated soils. Field engineers thus can determine the optimum quantity of gellan gum needed to provide the most effective strengthening based on the soil
type (USCS classification) and clay content in the soil \( (m_d / m_s) \).

For low plasticity clay (CL), 5% cement treatment improves the \( c \) from 75.5 to 152.1 kPa, while the \( \phi \) increases from 27.5° to 34.2° after 28 days of curing [81]. However, for gellan gum treatment, 4% gellan gum-treated pure clay enhances the \( c_d \) from 18.5 kPa to 127.3 kPa and the \( \phi_d \) from 18.7° to 30.7° only 12 h after treatment (Fig. 6). Thus, gellan gum treatment shows potential to be applied for quick and immediate soil stabilization or improvement purposes in geotechnical engineering practices. Moreover, gellan gum can become a promising binder for the currently increasing demands on eco-friendly raw earth buildings and constructions [33, 60].

Meanwhile, despite the positive impact on the shear behavior of gellan gum biopolymer-treated soils, practical realization becomes an important concern for gellan gum biopolymer usage in real in situ practices. High-temperature control and thorough mixing or injection into soils with high fine contents become challenges for the future commercialization of gellan gum biopolymer in the field of geotechnical engineering. Thus, further research is required to develop detailed construction methods and systems, as well as relevant machinery, which have high compatibility with existing construction technologies.

Moreover, economic concerns regarding gellan gum utilization for soil treatment remain, based on its relatively high cost compared to conventional soil treatment materials such as ordinary cement. Chang et al. [21] showed that the total cost to treat one ton of soil with gellan gum via thermal treatment is approximately US$25-30, comparable to the current cost (US$10) of 10% cement treatment. However, recent analyses of economic feasibility and prospects show that the use of biopolymers for construction purposes is becoming more feasible as a result of the current rapid growth of the global biopolymer market [34, 41, 69].

5 Conclusions

Through a series of experimental programs, this study explored the strengthening behavior of gellan gum-treated soils (from sand to clay). The results of laboratory vane shear strength (\( \tau_v \)) and direct shear strength parameters \( (c_d, \phi_d, \tau_f) \) for different sand–clay mixtures treated by gellan gum indicate that gellan gum treatment enhances the strength of soils significantly and that the strengthening behavior of gellan gum-treated soils involves a combination of phenomena. These include (1) the formation of a gellan gum gel or gellan gum–clay matrix, which enhances \( c_d \); (2) conglomeration induced by gellan gum–clay matrices, which increases the \( \phi_d \); and (3) higher internal strength of gellan gum-treated soils due to the water
holding characteristic of hydrophilic gellan gum. Indeed, further strengthening due to gellan gum gels or gellan gum–clay matrix condensation with drying is also expected. For soils with clay, the strengthening has a unique characteristic that invariably depends on the gellan gum-to-clay ratio in mass \((m_g/m_c)\). In detail, the gellan gum–clay matrix of kaolinite clay is optimized at around \(m_g/m_c = 4\%\) for gellan gum–kaolinite mixtures, regardless of the soil composition (i.e., \(m_g/m_c\)) and soil water content. However, \(m_g/m_c\) conditions higher than 4% are expected to yield convergence or even a reduction in strength due to ionic repulsion or hydrophilic swelling of surplus gellan gum monomers, which remain unbonded with clay particles.

The microstructure of gellan gum-treated soils indicates the formation of direct hydrogen bonding between gellan gum monomers and clay particles, especially at positively charged edges. The resulting gellan gum–clay matrices increase the shear strength of soils by enhancing \(c_u\) and conglomeration, resulting in an increase in the \(\phi_s\) of soils. When the goal is the strengthening of soils, sand–clay mixtures have advantages that result in higher strength due to the conglomeration effect between the gellan gum–clay matrix and granular particles.

The findings obtained from this study provide a guideline for optimal gellan gum treatment in geotechnical engineering practices. The results of this study show that \(m_g/m_c\) becomes the dominant parameter governing the strengthening behavior of the gellan gum-treated soils. Thus, field engineers can determine the optimum quantity of gellan gum needed to provide the most effective strengthening based on the soil type (USCS classification) and clay content in the soil \((m_g/m_c)\). If the clay mineral is mainly kaolinite, \(m_g/m_c = 4\%\) can be recommended for maximum strength capacity, while other clay types should be evaluated in further studies.

The significant shear strength and stiffness enhancement of gellan gum-treated sandy soils for low overburden stress suggest the significant potential of gellan gum applications for geotechnical earthquake-related soil management practices where an increase in both cyclic shear resistance and post-liquefaction shear strength is desired. Considering the aspect of environmental friendliness, gellan gum treatment can be applied for various geotechnical engineering purposes such as soil stabilization, deep mixing, surface erosion reduction, temporary soil structures, and so on. Moreover, the rapidly growing global biopolymer market and carbon emission trading market are improving the economic feasibility of gellan gum–soil technologies as a promising in situ implementation approach in the near future.

Acknowledgements The research described in this paper was financially supported by a National Research Foundation of Korea (NRF) grant funded by the Korean government (MSIP) (No. 2017R1A2B4008635). A Grant (18AWMP-B114119-03) from the Water Management Research Program and a Grant (18SCIP-B105148-04) from the Construction Technology Research Program both funded by the Ministry of Land, Infrastructure and Transport (MOLIT) of the Korean government also supported this research.

Open Access This article is distributed under the terms of the Creative Commons Attribution 4.0 International License (http://creativecommons.org/licenses/by/4.0/), which permits unrestricted use, distribution, and reproduction in any medium, provided you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made.

References

1. Andrus RD, Stokoe KH II (2000) Liquefaction resistance of soils from shear-wave velocity. J Geotech Geoenvir Eng 126(11):1015–1025. https://doi.org/10.1061/(ASCE)1090-0241(2000)126:11(1015)
2. ASTM D3080/D3080 M-11 (2011) Standard test method for direct shear test of soils under consolidated drained conditions. ASTM International, West Conshohocken. https://doi.org/10.1520/D3080_D3080M-11
3. Biscontin G, Pestana JM (2001) Influence of peripheral velocity on vane shear strength of an artificial clay. Geotech Test J 24(4):423–429. https://doi.org/10.1520/GTJ1140J
4. Bolton MD (1986) The strength and dilatancy of sands. Géotechnique 36(1):65–78. https://doi.org/10.1680/geot.1986.36.1.65
5. Bouazza A, Gates WP, Ranjith PG (2009) Hydraulic conductivity of biopolymer-treated silty sand. Géotechnique 59(1):71–72. https://doi.org/10.1680/geot.2009.00137
6. British Standard Institute (1990) BS 1377-7: Methods of test for soils for civil engineering purposes. Part 7–Shear strength tests (total stress). British Standard Institute, London
7. British Standard Institute (2014) BS EN ISO 17892-1: Geotechnical investigation and testing. Laboratory testing of soil. Part 1–Determination of water content. British Standard Institute, London
8. Brunori F, Penzo MC, Torri D (2013) Soil shear strength: its measurement and soil detachability. CATENA 16(1):59–71. https://doi.org/10.1016/j.catena.2013.03.009
9. Budhu M (1985) The effect of clay content on liquid limit from a fall cone and the British Cup Device. Geotech Test J 8(2):91–95. https://doi.org/10.1520/GTJ10515J
10. Cao J, Jung J, Song X, Bate B (2018) On the soil water characteristic curves of poorly graded granular materials in aqueous polymer solutions. Acta Geotech 13(1):103–116. https://doi.org/10.1007/s11440-017-0568-7
11. Chandler RJ (1988) The in-situ measurement of the undrained shear strength of clays using the field vane. In: Vane shear strength testing in soils: field and laboratory studies vol 1014, p 13
12. Chang I, Cho G-C (2010) A new alternative for estimation of geotechnical engineering parameters in reclaimed clays by using shear wave velocity. Geotech Test J 33(3):171–182. https://doi.org/10.1520/GTJ102360
13. Chang I, Cho G-C (2012) Strengthening of Korean residual soil with \(\beta\)-1,3/1,6-glucan biopolymer. Constr Build Mater 30:30–35. https://doi.org/10.1016/j.conbuildmat.2011.11.030

© Springer
