



Environment, Energy and Chemistry

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Environment Energy & Chemical Solution

Dry Complex Deodorization System
Dry Combined Odor Removal System

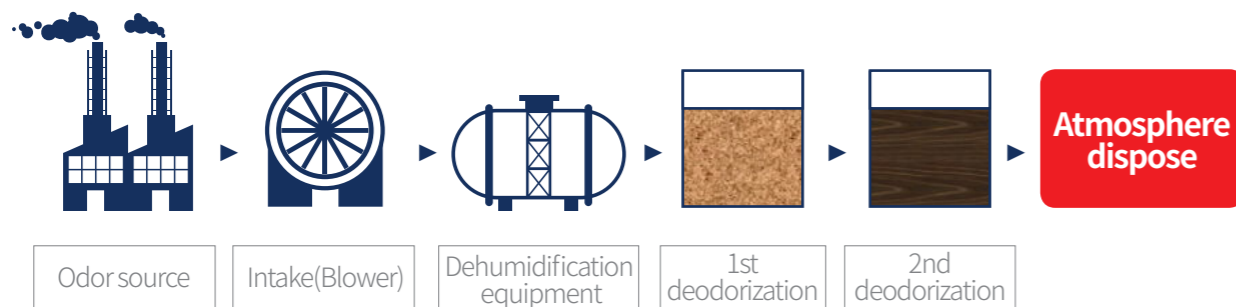
The need to remove odors



- Intensifying complaints about odor in residential areas near public treatment facilities
- Reinforcement of odor regulation according to the improvement of citizen's living standard
- It is necessary to remove harmful substances to the human body contained in odorous substances.

* A bad odor is an odor that causes an unpleasant and disgusting feeling to people.

Schematic diagram of odor removal system



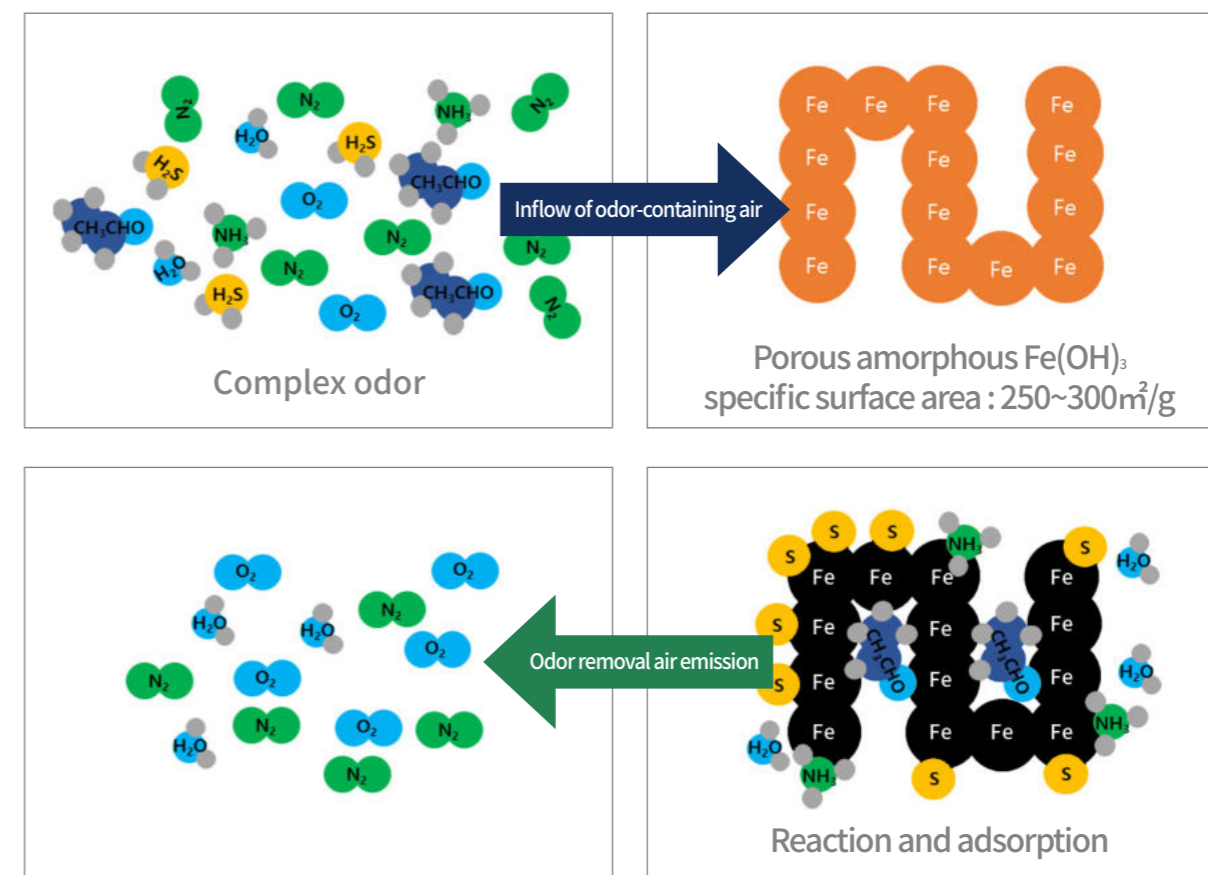
Ammonia and hydrogen sulfide are sequentially removed through a dehumidification process and iron hydroxide-based adsorbent. Then, the voc's are removed through the activated carbon tower and the air is discharged to secure clean air quality.

* VOC's : volatile organic compounds

Effect of Odor Removal

Improvement of life quality in residential areas around public facilities	By preventing the release of hydrogen sulfide and ammonia, VOC's into the atmosphere, the clean air quality is secured
Decreases of complaints due to odors and improving working conditions for field works	Securing odor source data for complex odor-causing substances and Customized odor removal process design

Primary deodorization principle (removal of hydrogen sulfide and ammonia)

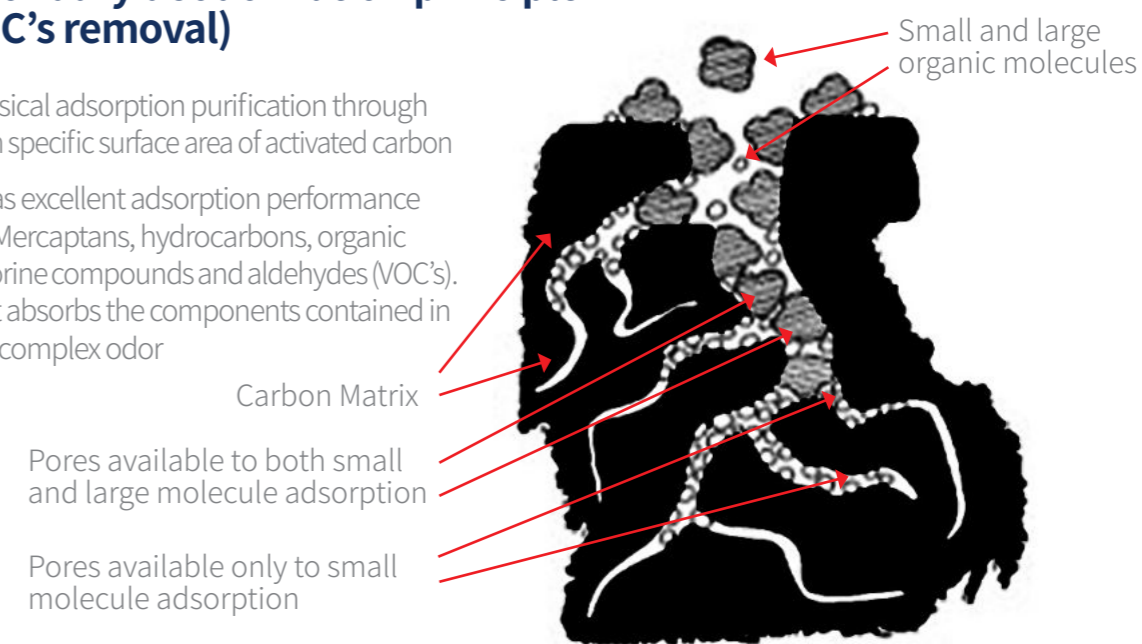


<Removal mechanism of hydrogen sulfide and ammonia>

- Hydrogen sulfide is removed by adsorption reaction through a porous iron hydroxide-based adsorbent. Then, ammonia and some voc's are adsorbed into the formed pores for primary purification.

Secondary deodorization principle (VOC's removal)

- Physical adsorption purification through high specific surface area of activated carbon
- It has excellent adsorption performance for Mercaptans, hydrocarbons, organic chlorine compounds and aldehydes (VOC's), so it absorbs the components contained in the complex odor



<Activated carbon physisorption principle>

Dry complex odor removal system



Process Features

- Removal of ammonia and moisture using dehumidification process
- Removal of hydrogen sulfide using iron hydroxide
- Removal of complex odors by removing aldehydes using activated carbon

Existing odor removal

- Short adsorbent replacement cycle due to wetting
- Unaware of the need for dehumidification
- Low removal rate due to low reactivity between hydrogen sulfide and iron oxide.
- High operating cost due to short replacement cycle
- The application of a multi-stage chemical cleaning (scrubbing) tower increases costs such as facility investment, chemicals, and electricity.
- A large amount of waste liquid and waste water generated by chemical cleaning through scrubbing
- Requires additional facilities to prevent the generation of nitrogen oxides (NOx) during combustion deodorization and high operating costs due to fuel use

Dry odor removal

- A dry process through dehumidification to prevent deterioration of adsorption capacity due to wetting
- 60% reduction in dehumidification cost through two-stage heat exchange dehumidification
- Since hydrogen sulfide is separately removed by iron hydroxide(Fe(OH)₃), it exhibits excellent compound odor removal performance compared to the existing process
- Reduction of initial installation cost due to minimum residence time
- Reduction of operating costs by using minimal power
- Unmanned operation possible and easy on-site maintenance
- No wastewater generated by the use of chemicals

Sludge Concentration Tank-derived Odor Removal Demonstration Test at Jungnang Water Regeneration Center



Conditions for removal of odors

- Demonstration test capacity : 400 L/min
- Hydrogen sulfide(H₂S) density : 1~4 ppm
- Complex odor dilution Time : 100~60,000 times
- Test bed size : 5L * 2part(top/bottom configuration)
- Odor Stream : Up-stream method



Existing odor removal equipment (4.7m³ * 3part = 14.1 m³)



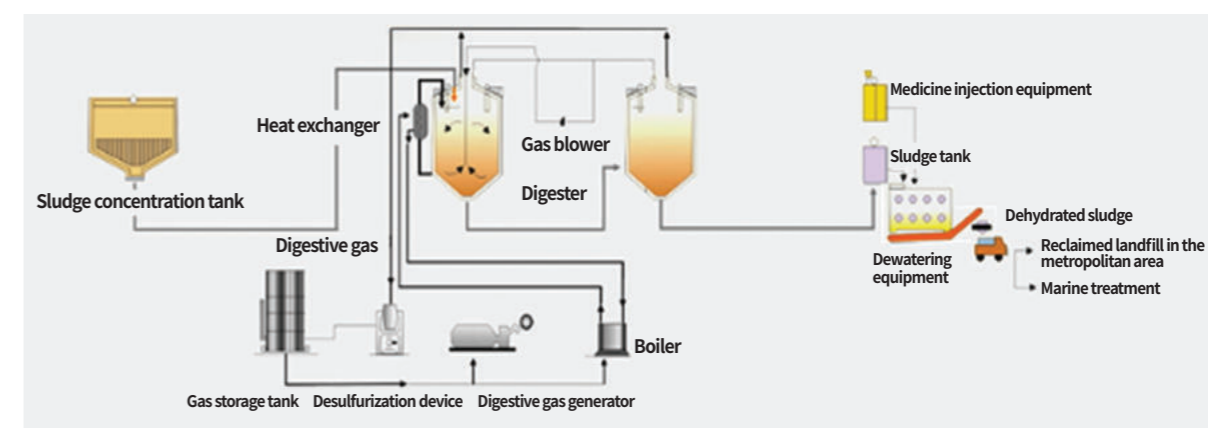
Demonstration facility (5L * 2part = 10L)



Odor stream

Composition

- 2nd deodorization
Impregnated activated carbon
- 1st deodorization
DeHyS-250



Odor generation and verification point



Inlet



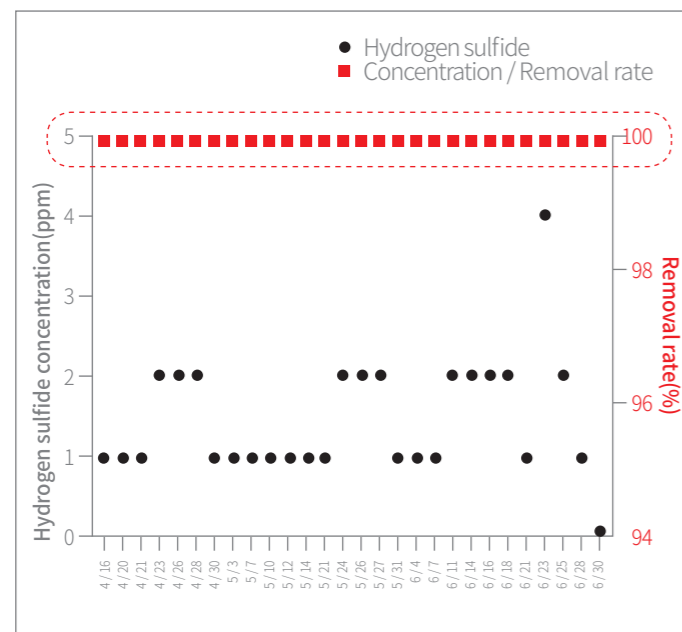
1st deodorization bed



2nd deodorization bed

Complex odor measurement and analysis : Korea EMC

Concentration Tank-derived Odor Removal Demonstration Test at Jungnang WWTP

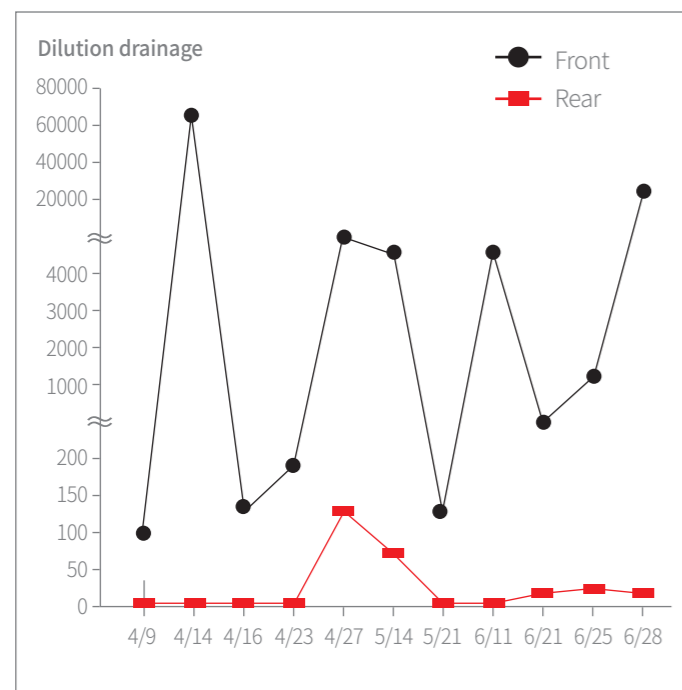


Demonstration test driving conditions

- Space velocity : 4,800hr⁻¹
- Residence time : 0.75 sec
- Throughput : 400 L/min

Hydrogen sulfide (odor-causing substance) removal rate

- **100% removal**
(Operation period: 49 days)



Diluted drainage after inflow of odor and treatment

- Dilution of inflow odor : 100 to 60,000 times
- Influence of sludge loading situation and weather
- Dilution of odor after treatment : 10~150times
- Results measured through air dilution sensory method
- Development of dry complex odor removal process with excellent odor removal performance

Division	Inflow	1 st deodorization	2 nd deodorization
Dilution factor	448	207	30

※ Complex odor measurement and analysis progress (Measuring Agency : Korea EMC)

Odor type

- Concentration tank-derived occurrence

Field test conditions

- 400 L/min, Operating hours (49 days)

Processing criteria

- Hydrogen sulfide less than 1ppm,
Dilution drainage less than 100

Sampling location

- Inflow, primary (DeHyS-250),
secondary (impregnated activated carbon)

Digested Sludge Reservoir-derived Odor Removal Demonstration Test at Tanchon Water Regeneration Center

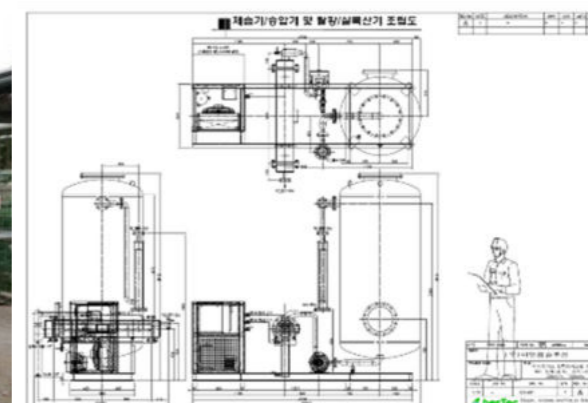


Conditions for ingress of odors

- Demonstration test capacity : 2,500 L/min
- Hydrogen sulfide (H₂S) density : 10~40 ppm
- Complex odor dilution drainage : 1,000~10,000 times
- Test bed size : 500L
- Residence time : 12sec
- Odor inflow method : Up-stream method

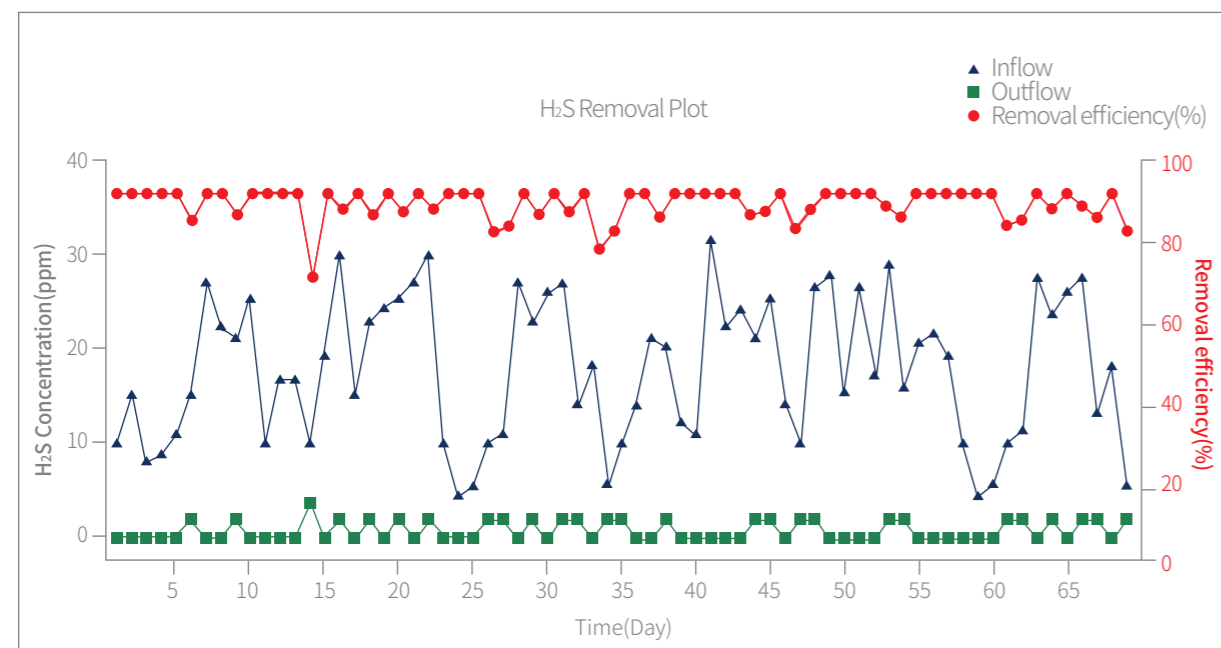
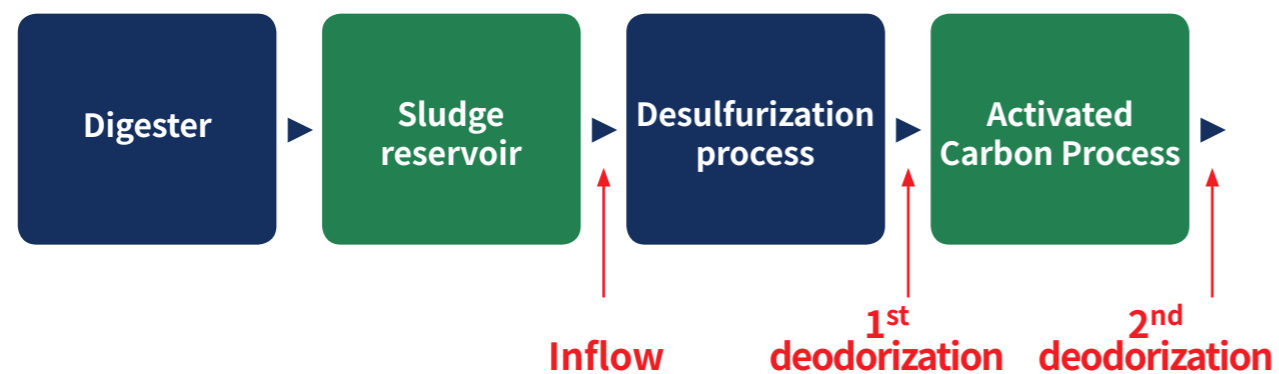


<Digestive Thalys Reservoir at Seoul Tanchon Water Regeneration Center → Pilot-plant about 70m>



<Installation site appearance & design>

Digested Sludge Reservoir-derived Odor Removal Demonstration Test at Sludge storage tank



Hydrogen sulfide inlet and outlet concentrations

- Average inlet concentration(ppm) : 18.16
- Average outlet concentration(ppm) : 0.41
- Average hydrogen sulfide removal rate(%) : 97.76

Division	Inflow	1 st deodorization	2 nd deodorization
Dilution drainage	3,000	1,000	448

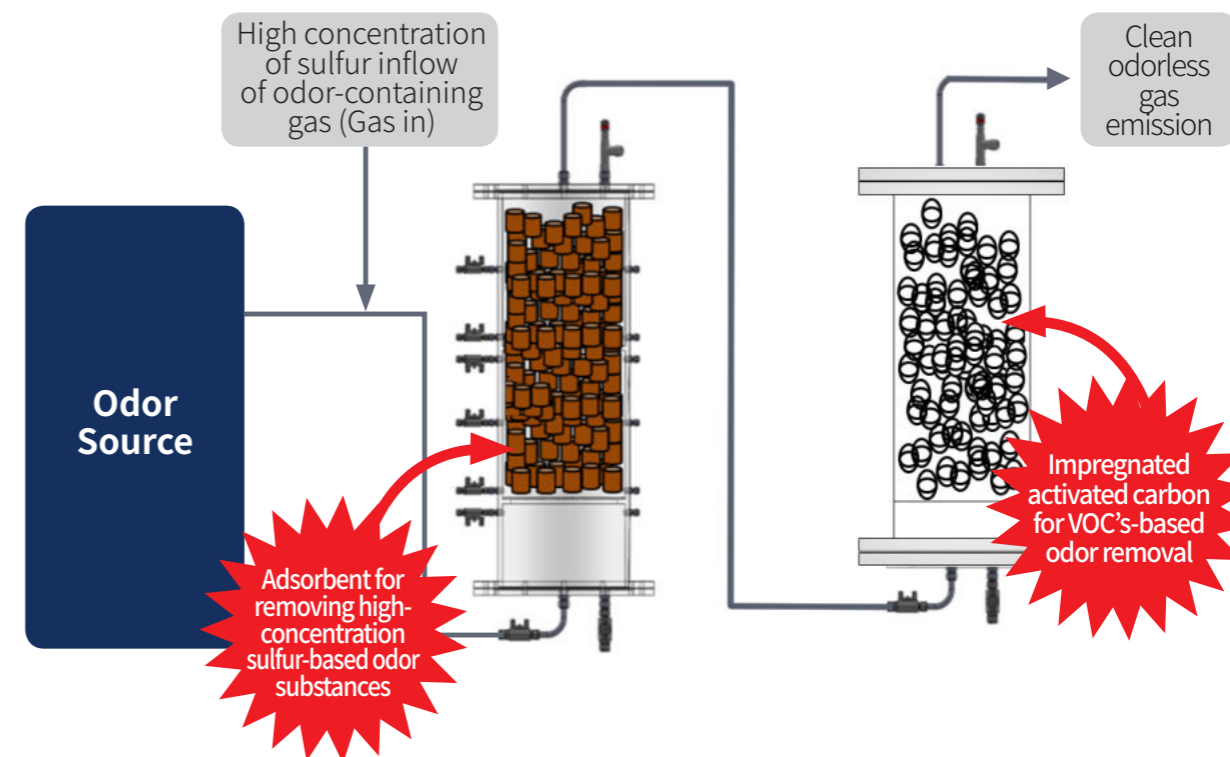
※ Complex odor measurement and analysis progress (Measuring Agency : Korea EMC)

Odor type	Field test conditions
• Digested sludge reservoir	• 2,500 L/min
Processing criteria	Sampling location
• Hydrogen sulfide less than 1ppm, Dilution drainage less than 100	• Inflow, After dry adsorption, After wet adsorption

Proposal of odor removal demonstration test in sewage treatment plant

Expected odor conditions	Demonstration test driving conditions
• Expected hydrogen sulfide(H ₂ S) concentration : 1~40 ppm	• Demonstration test capacity : 10 CMM, 8 HP
• Expected ammonia (NH ₃) concentration : 1~60 ppm	• Test bed size : 1.0 m ³
• Expected dilution factor of inflow complex odor : 1,000~60,000 times	• Odor inflow method : Up-stream method (Utilize an air blower)
	• Residence Time : 6 sec
	• Space Velocity : 600 hr ⁻¹

Schematic diagram of odor removal system



A odor removal system inhales the complex odor diluted in the air from the odor source. After the primary and secondary deodorization processes are carried out through the dehumidification process, the odor-free air is discharged. However, in the case of dry gas, dehumidification facilities are excluded.

Odor Removal Target

Division	Removal rate	1 st deodorization	2 nd deodorization
Hydrogen sulfide(ppm)	99%	1 or less	1 or less
Complex odor	80%	1,000 or less	300 or less
Moisture	-	50RH% or less	50RH% or less

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Catalytic Desulfurization System

Desulfurization catalyst system

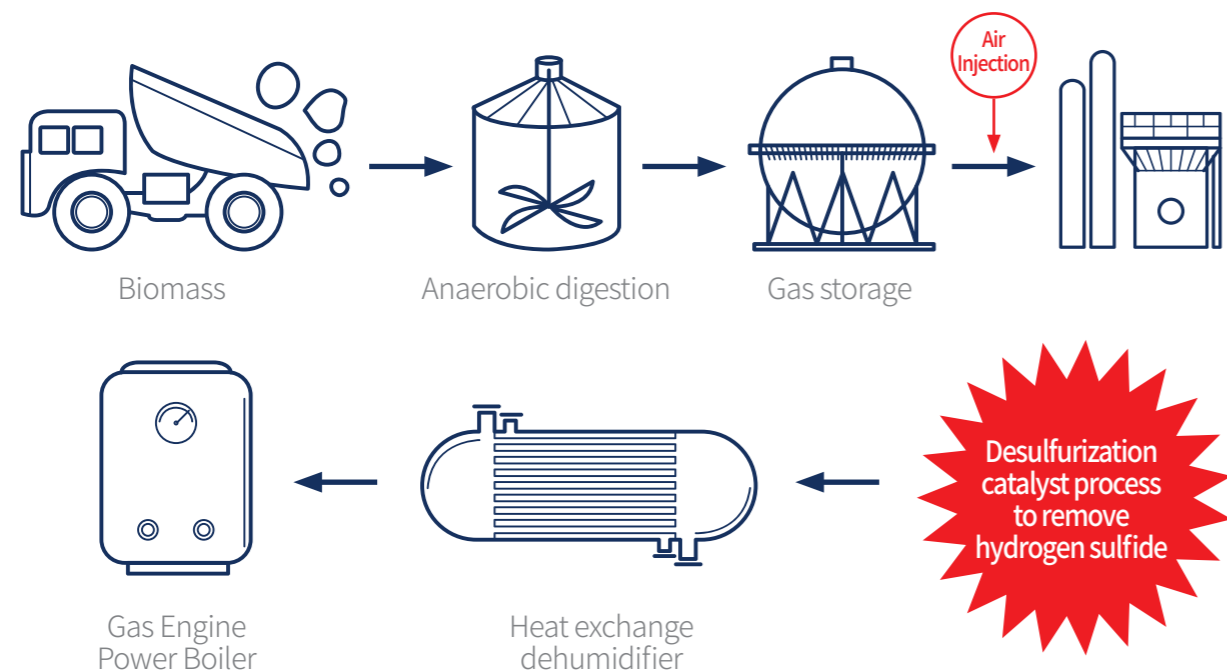
Iron oxide(Fe₂O₃) catalyst



Desulfurization Catalytic Reaction Mechanism

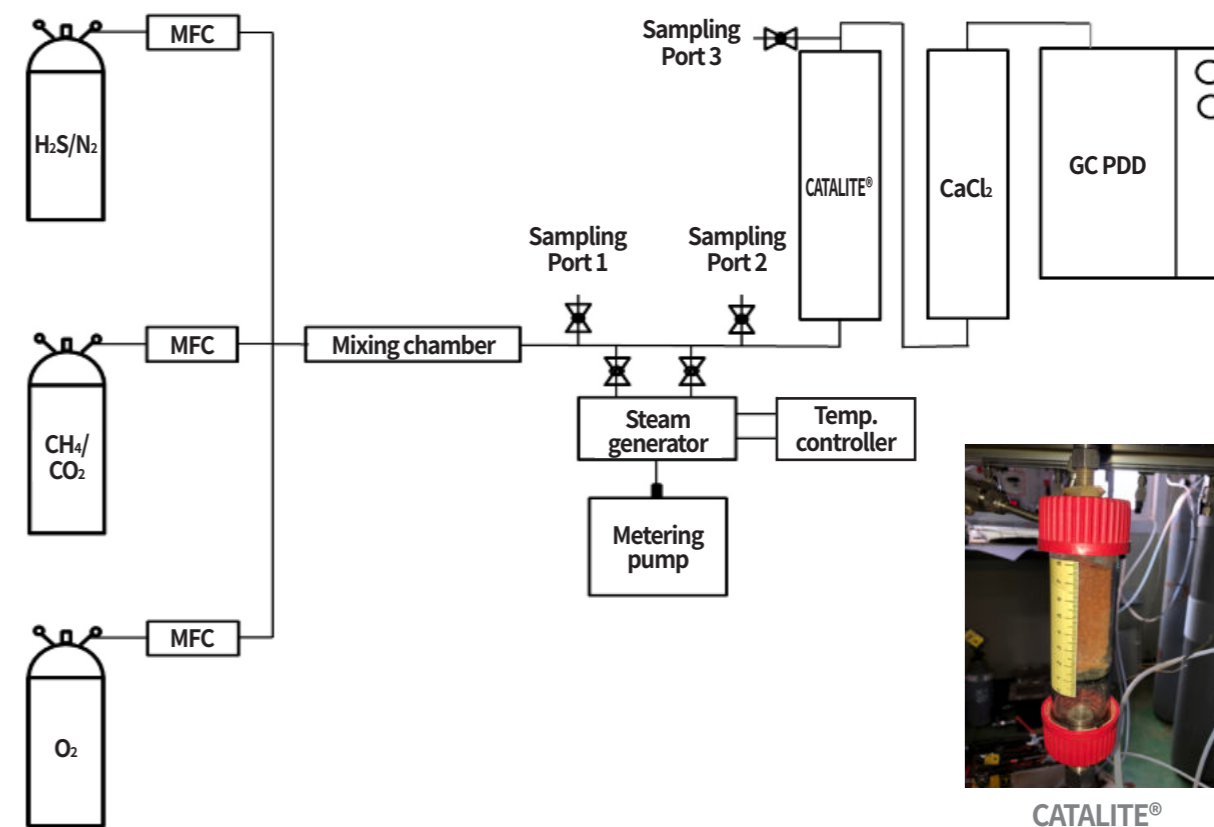


Schematic diagram of biogas production facility



Hydrogen Sulfide(H₂S) Desulfurization Catalyst Test I

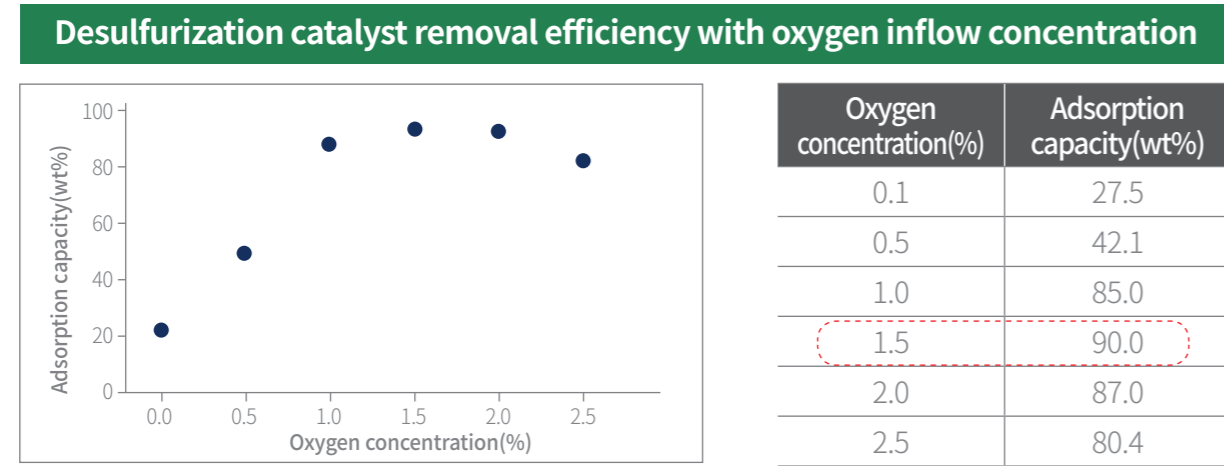
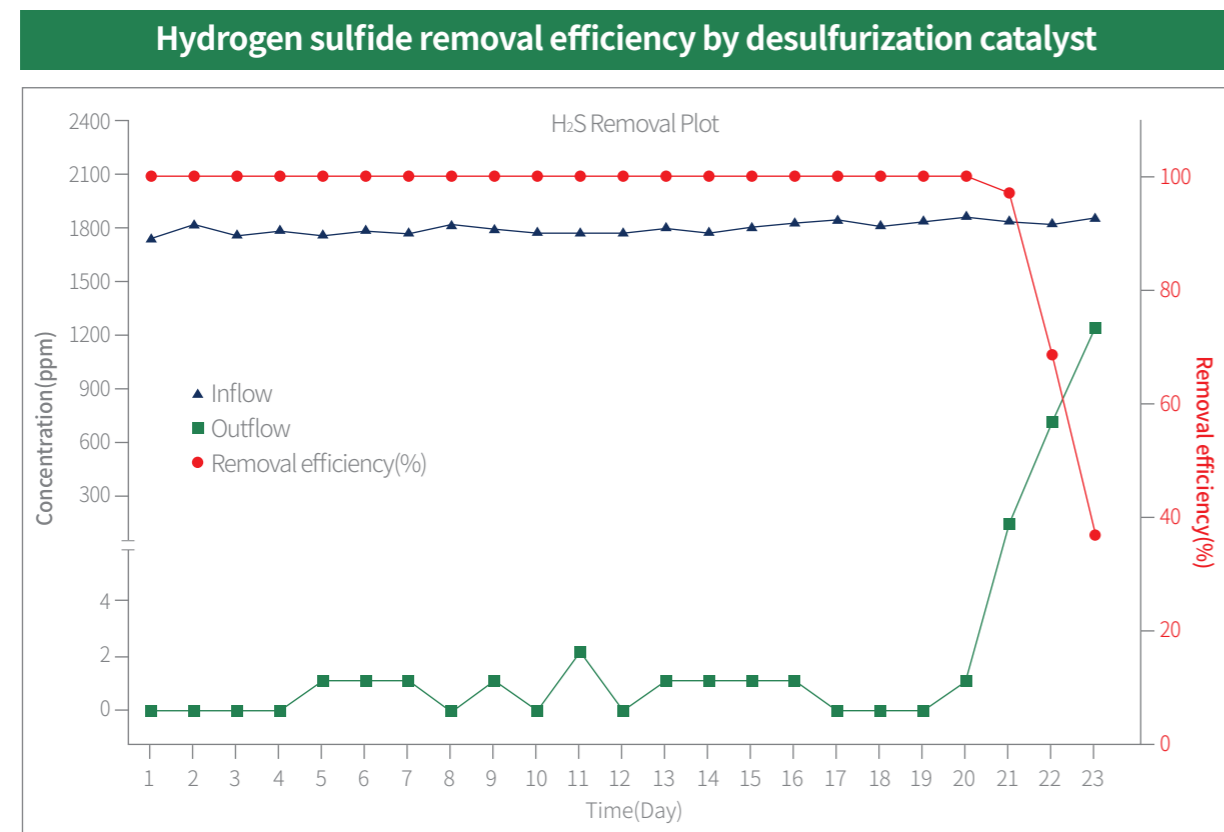
Item	Unit	CATALITE®	Remarks
Dilution drainage	3,000	wt%	Catalytic reaction experiment result criteria



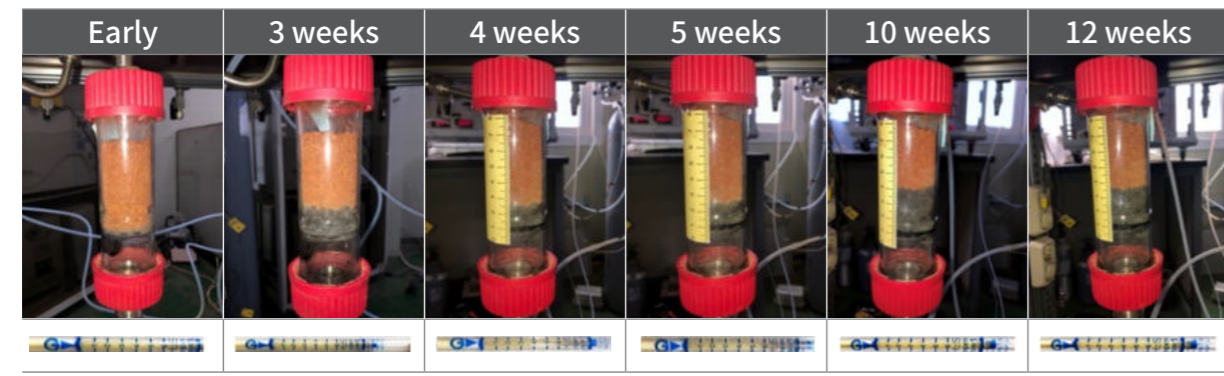
Experimental Conditions

- H₂S = 1,600 ~ 1,800 ppm
- O₂ = 1 ~ 2%, O₂
- Humidity : RH 100%
- Bed volume : 100ml
(Capa. : 36g(60ml))
- Flow rate : 50ml/min
- Space Velocity = 50 hr⁻¹
- Contact time = **72 sec**

Hydrogen sulfide(H₂S) Desulfurization Catalyst Test II



Desulfurization catalyst discoloration



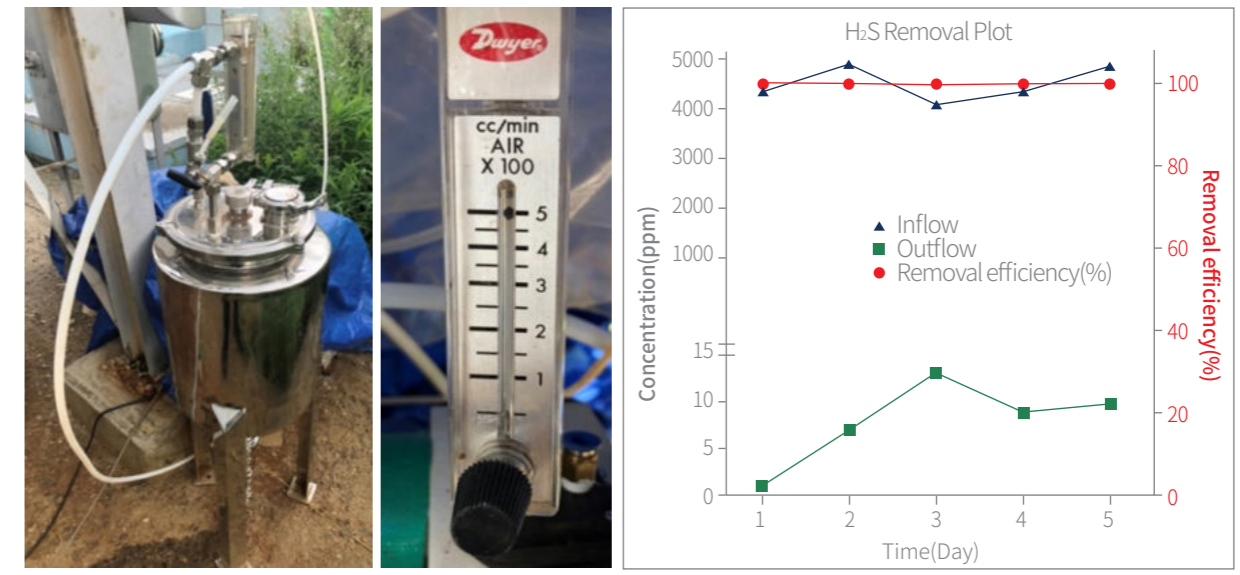
Detection of H₂S less than 1ppm as a result of checking the detector tube

Biogas pretreatment site test results I



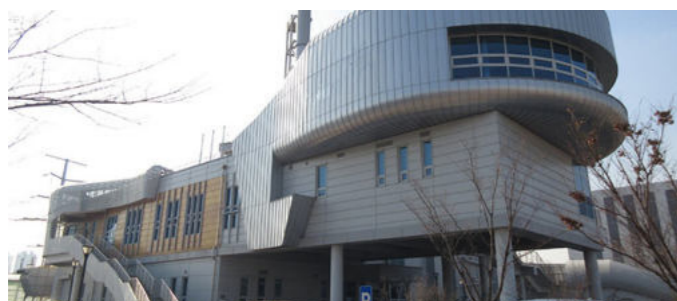
Byeolnae Food Waste Clean Center

- Sewage treatment capacity : 275,000 m³/day
- Biogas generation : 8,000m³/day
- Hydrogen sulfide(H₂S) : 5,000 ppm or more
- Siloxane : around 50 mg/m³



- Biogas Application : biogas production for use in boilers and incineration
- Process gas concentration : hydrogen sulfide 10 ppm or less
- Sampling location : 1 front and 1 rear of test bed respectively
- Conditions for field testing : 15L/min, Average hydrogen sulfide concentration (approximately 5,000 ppm), operating time (720 hr)
- Air injection : 500 ml/min, inflow composition (CH₄ 60%/CO₂ 30%/H₂S 0.5%/H₂O 5%/O₂ 1%)
- ※ Goal : Hydrogen sulfide removal rate of 99% or higher (99.5% or higher)

Biogas pretreatment site test results II

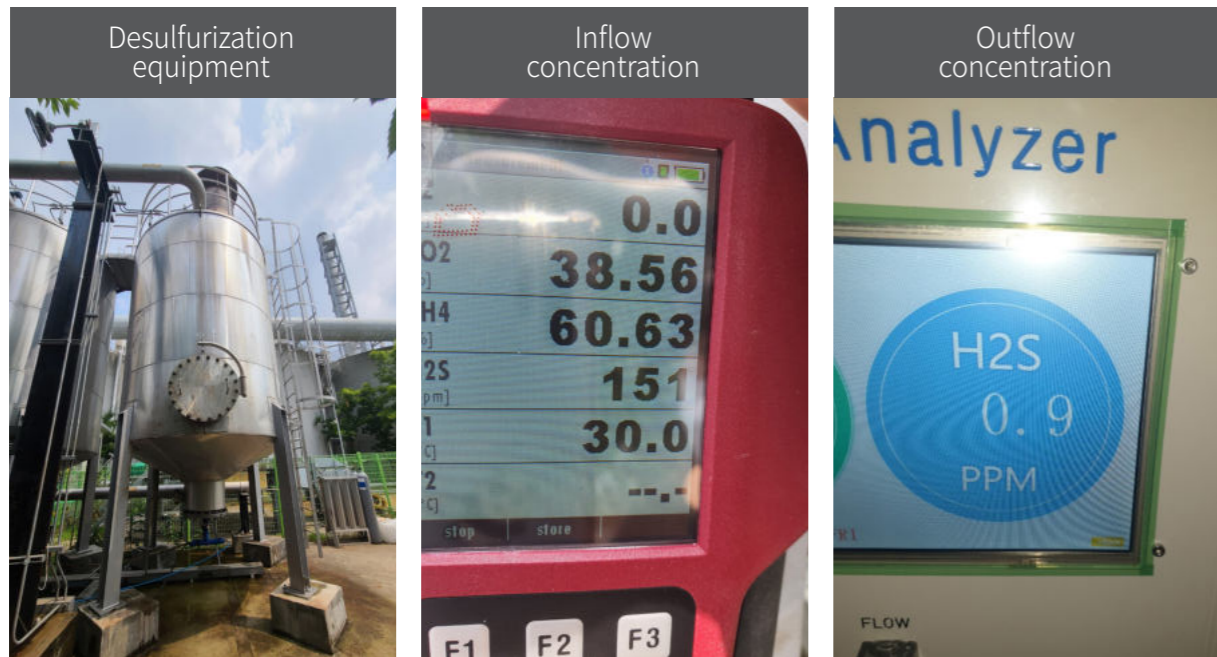


Goyang Biomass energy facility

- Food waste treatment capacity : 260 ton/day
- Biogas generation : 20,000~24,000m³/day
- Hydrogen sulfide(H₂S) Concentration : around 150 ppm



Memo



- **Biogas Usage** : Utilization of Urban Gas in Goyang City
- **Processing gas concentration** : 10 ppm or less in hydrogen sulfide (urban gas supply line 0.9 ppm)
- **Sampling Location** : Desulfurization Facility Shear and Urban Gas Supply Line
- This is the first supply facility for the desulfurization catalyst.
- Air supply blower for catalytic reaction maintains oxygen concentration according to biogas flow rate.



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Dry System for **VOC's Removal**

Dry system for removing VOC's

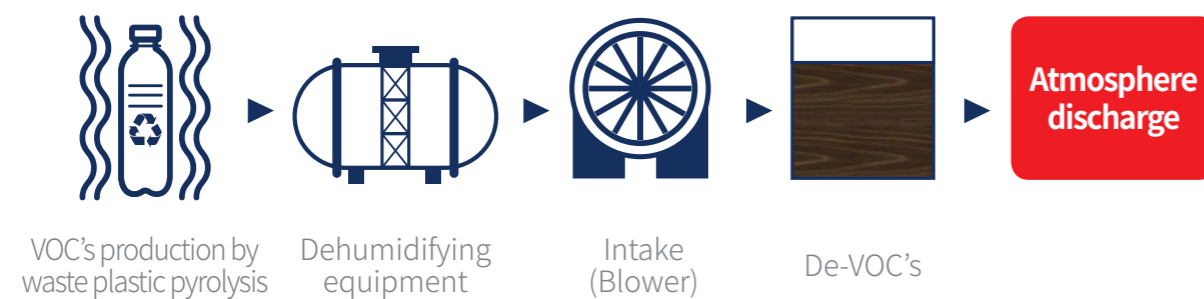
Necessity to Remove Waste Plastic VOC's

- VOC's line of odor complaints intensified near waste plastic recycling facilities
- When discharged into the atmosphere due to the occurrence of a large amount of VOC's, air quality deteriorates
- Large amounts of harmful organic compounds are discharged during melting of waste plastics.

* VOC's are volatile organic compounds including emission regulations and odorous substances.



Flow Chart of Waste Plastic Recycling VOC's Removal System



When recycling waste plastics, VOC's generated by thermal decomposition are absorbed. And the high boiling point VOC's are removed from condensate through the dehumidification facility and the low boiling point VOC's are removed from the VOC's through an activated carbon tower and discharged to the atmosphere to secure clean air quality

* VOC's: Volatile Organic Compounds

Effect of removing VOC's

Improvement of life quality in the residential areas around public facilities

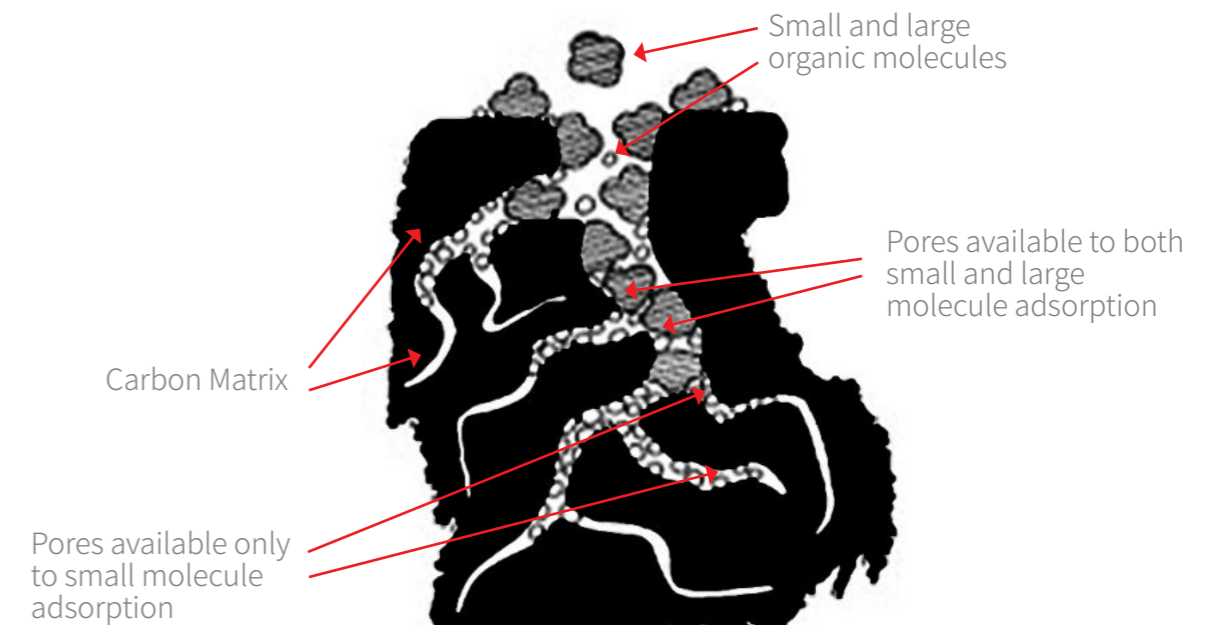
By preventing the release of VOC's into the atmosphere, the generation of fine dust is suppressed and clean air quality is secured

Decrease of complaints due to odors and improving working conditions for field works

Securing odor source data for complex odor-causing substances and Customized odor removal process design

VOC's removal principles

- Physical adsorption purification through high specific surface area of activated carbon
- It has excellent adsorption performance for mercaptans, hydrocarbons, organochlorine compounds and aldehydes. So it absorbs the components contained in complex odor



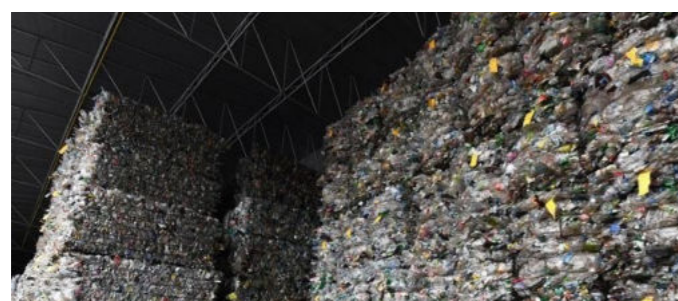
<Activated carbon physisorption principle>

Types of VOC's occurred in Waste Plastic recycling processe

<VOC's occurrence according to the type of recycled plastic by the Pocheon's company>

Type	Substance containing
PP	Propane, Polymeric carbon compounds, etc.
PE	Ethane, Polymeric carbon compounds, etc.
PS	Styrene, Toluene, Ethylbenzen, etc
ABS	Butadiene, Styrene, Toluene, Ethylbenzene, etc

Odor removal test for waste plastic recycling in Pocheon

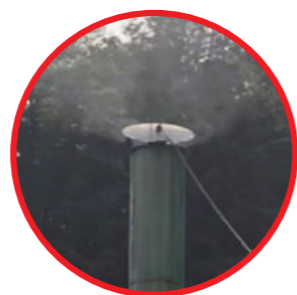


Existing inflow conditions

- Existing facility handing capacity : approximately 100 m³/min
- Types of recycled plastics : PP, PE, PS, AB, etc



Existing VOC's removal facility(30m³)



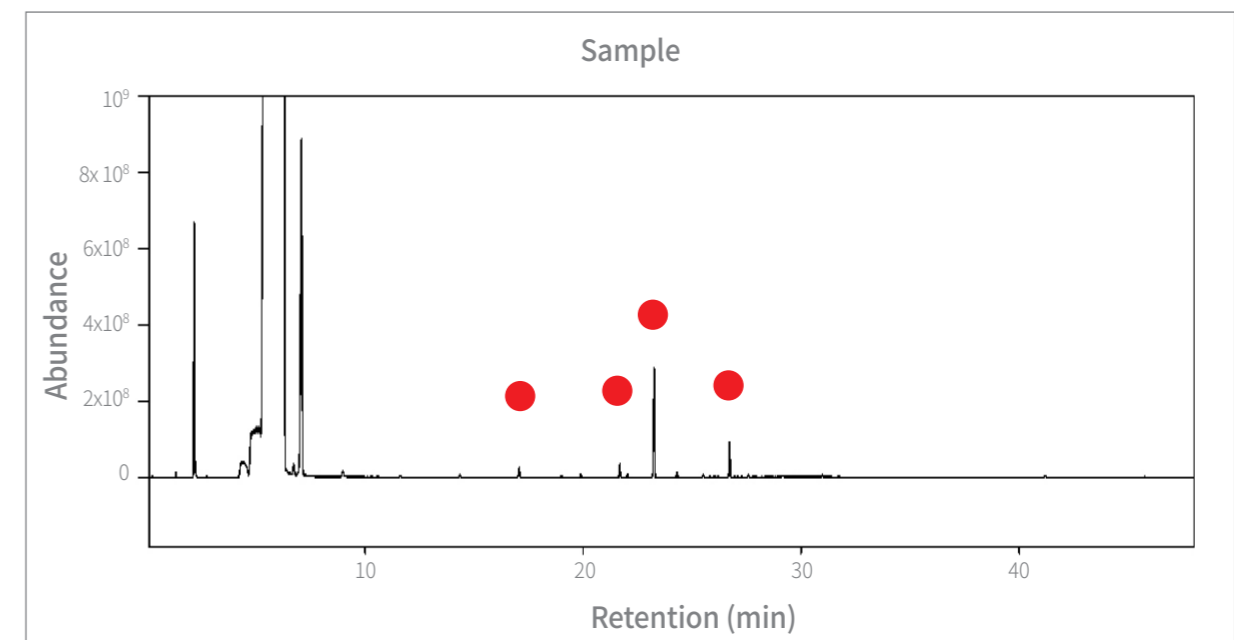
Generation of large amounts of VOC's fume



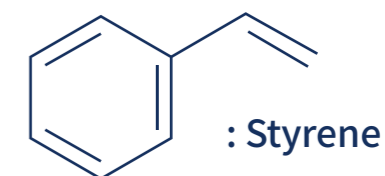
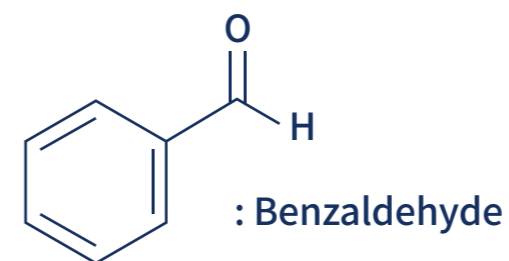
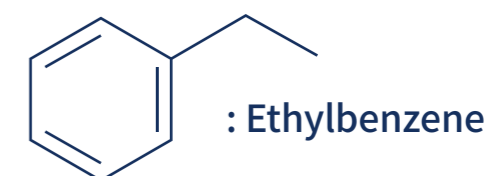
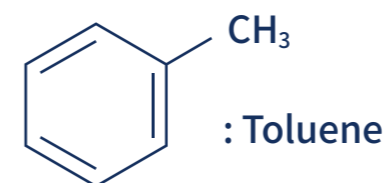
- Empirical test operating conditions
Space velocity : 300 hr⁻¹ / Residence time : 12 sec / Throughput : 500 L/min
- VOC's removal rate(Styrene, Toluene, Ethylbenzene)
100% removal (operating period: 60 days)

Qualitative analysis of plastic recycling heat process

Request for GC-MS (Seoul National University of Science and Technology) analysis

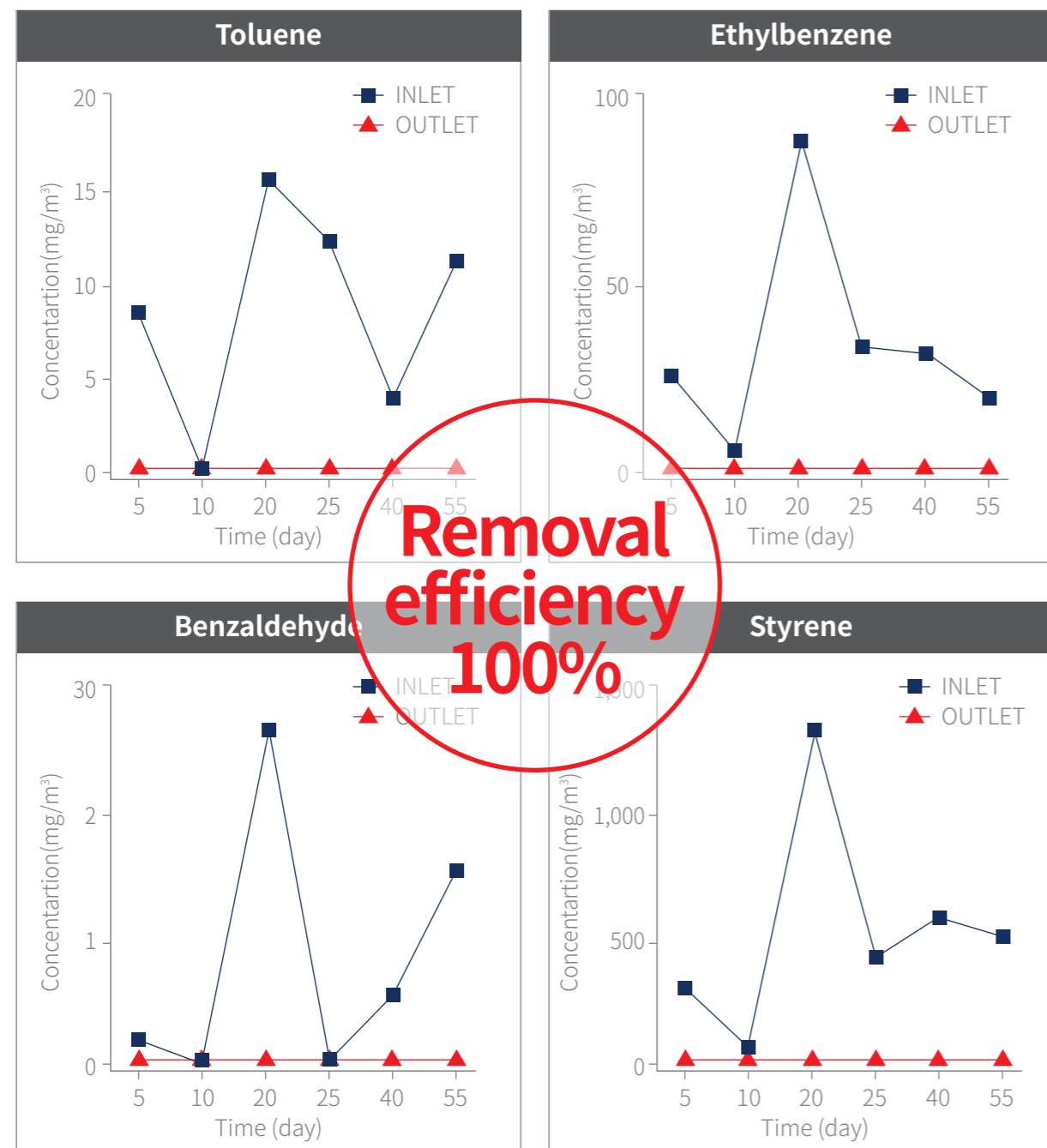


<GC-MS measurement results through hexane absorption method (Qualitative analysis)>



It mainly consists of the above components and there are many other types of aromatic organic compounds.

PILOT TEST VOC's analysis results



• Special note

- Manufacturer's plastic throughput is not constant
- unbalanced types of plastics

• Conclusion

- Wide range of influent concentration according to throughput
- Regardless of the inflow concentration, the removal rate is 100%

Emission acceptance standard

• Complex odor

Division	Emission acceptance standard (Dilution Drainage)		The range of Strict emission acceptance standard (Dilution Drainage)	
	Industrial area	Other regions	Industrial area	Other regions
Outlet	1000 or less	500 or less	500 ~ 1000	300 ~ 500
Site boundary	20 or less	15 or less	15 ~ 20	10 ~ 15

• Designated odor substance

Division	Emission acceptance standard (ppm)		The range of strict emission acceptance standard (ppm)	Application period
	Industrial area	Other regions	Industrial area	
Ammonia	2 or less	1 or less	1 ~ 2	From 2005. 2. 10
Methyl mercaptan	0.004 or less	0.002 or less	0.002 ~ 0.004	
Hydrogen sulfide	0.06 or less	0.02 or less	0.02 ~ 0.06	
Dimethylsulfide	0.05 or less	0.01 or less	0.01 ~ 0.05	
Dimethyl disulfide	0.03 or less	0.009 or less	0.009 ~ 0.03	
Trimethylamine	0.02 or less	0.005 or less	0.005 ~ 0.02	
Acetaldehyde	0.1 or less	0.05 or less	0.05 ~ 0.1	
Styrene	0.8 or less	0.4 or less	0.4 ~ 0.8	
Propionic aldehyde	0.1 or less	0.05 or less	0.05 ~ 0.1	
Butyl aldehyde	0.1 or less	0.029 or less	0.029 ~ 0.1	
n-valeraldehyde	0.02 or less	0.009 or less	0.009 ~ 0.02	
I-valeraldehyde	0.006 or less	0.003 or less	0.003 ~ 0.006	
Toluene	30 or less	10 or less	10 ~ 30	From 2008. 1. 1
Xylene	2 or less	1 or less	1 ~ 2	
Methyl ethyl ketone	35 or less	13 or less	13 ~ 35	From 2010.1. 1
Methyl isobutyl ketone	3 or less	1 or less	1 ~ 3	
Butyl acetate	4 or less	1 or less	1 ~ 4	
Propionic acid	0.07 or less	0.03 or less	0.03 ~ 0.07	
n-butylic acid	0.002 or less	0.001 or less	0.001 ~ 0.002	
n-valeric acid	0.002 or less	0.0009 or less	0.0009 ~ 0.002	
I-valeric acid	0.004 or less	0.001 or less	0.001 ~ 0.004	
I-butyl alcohol	4.0 or less	0.9 or less	0.9 ~ 4.0	

* Annex 3 of the Enforcement Rules of the Odor Prevention Act



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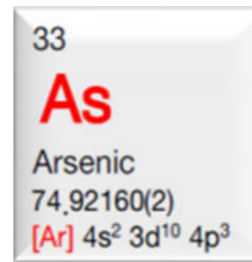
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MIRESORB®

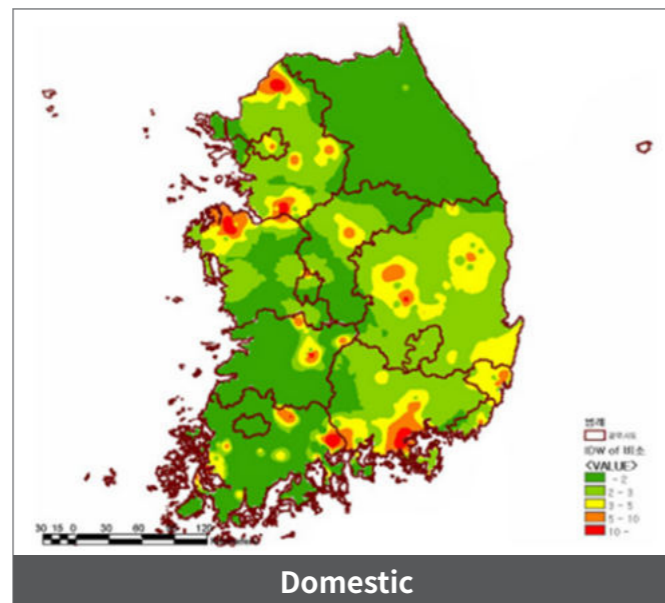
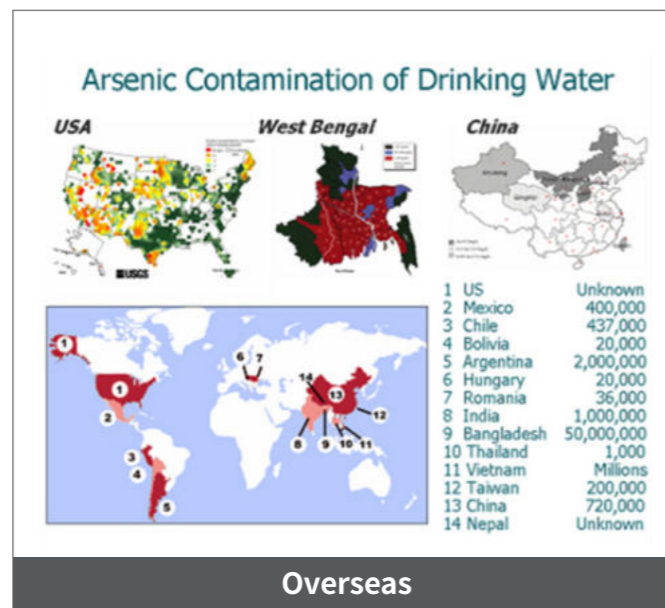
Adsorbent for arsenic removal

Arsenic (As)

- It is classified as a first-class carcinogen and mainly causes skin and liver cancer.
- It found in elemental conditions but exists mainly as sulfides
- Trivalent arsenic is about 40 to 60 times more toxic than pentavalent arsenic
- It exists in various forms such as Realgar(As₄S₄), Orpiment(As₂S₃), Arsenopyrite(FeSAs)



Arsenic contamination status of domestic and foreign groundwater



Drinking water standards in countries and organizations including WHO and USEPA: 10 µg/L

MIRESORB® Overview

MIRESORB®, an arsenic and heavy metal adsorbent in the water quality of E&C Solution Co., Ltd., is a porous adsorbent manufactured based on iron hydroxide. MIRESORB® is a product that guarantees performance through batch characteristics, column tests, on-site pilot test and actual delivery. Compared to foreign products, the performance is equal or higher and the price is 1/2 and the cost performance is very good

Granular type

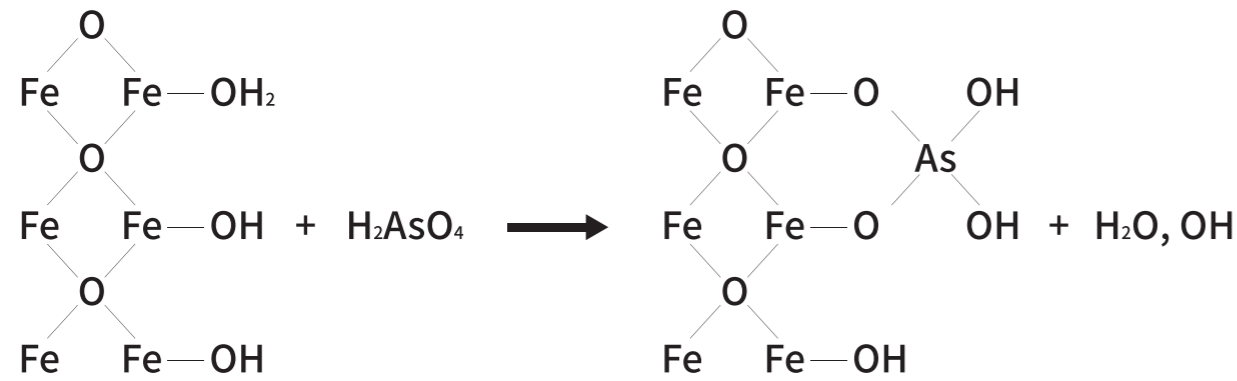


Pellet type



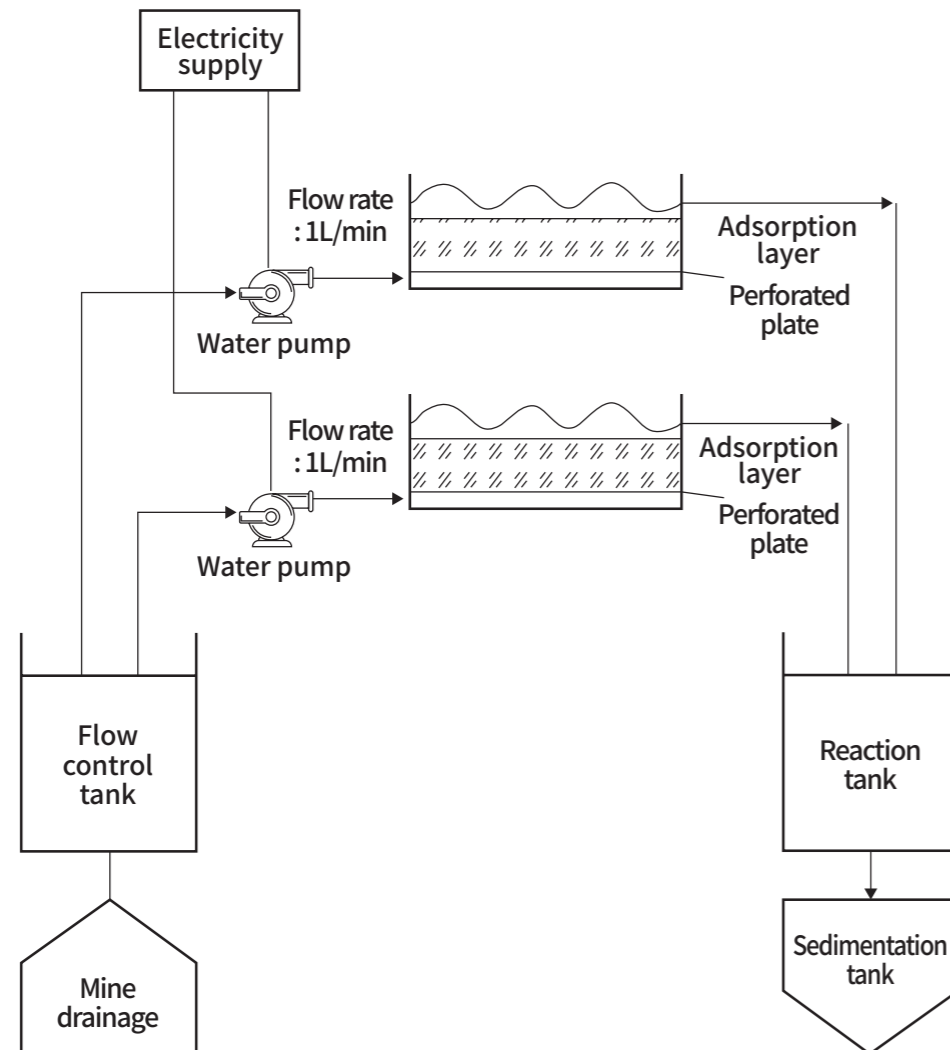
- As an insoluble inorganic compound, there is no concern about dissolution
- Performance assurance available in various EBCTs.
- Principal component : amorphous iron hydroxide(Fe(OH)₃)
- Specific surface area : 200~250m²/g
- Grain size : Customization
- Designable with packing factor
- Arsenic (V) adsorption capacity : 1 wt% or more
- Adsorption capacity of heavy metals : 1wt% or more

MIRESORB® Arsenic Adsorption Principle



<Arsenic adsorption mechanism>

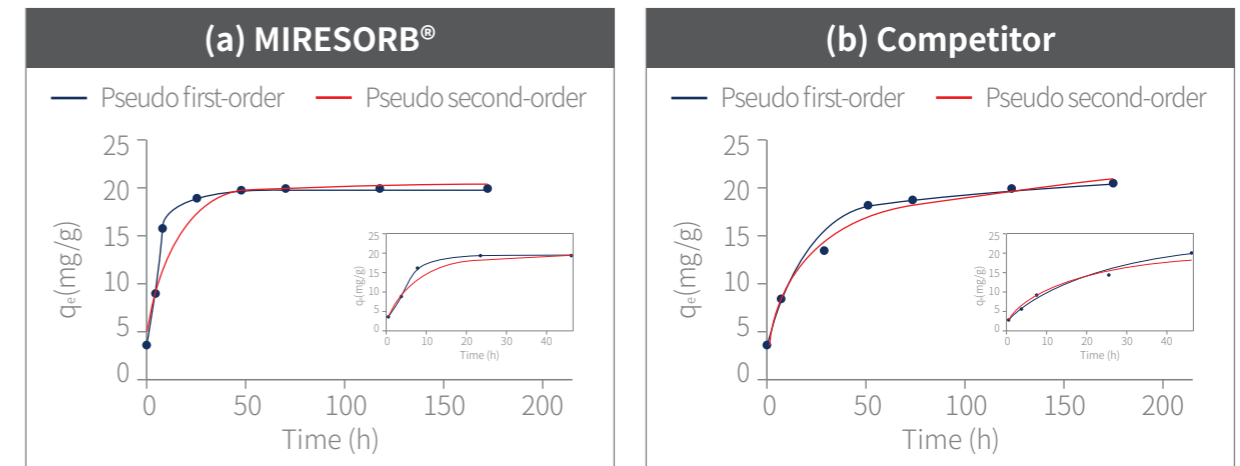
Based on adsorption principle for arsenic removal mechanism using iron hydroxide, arsenic ions are physically adsorbed on the OH-groups of the porous adsorbent.



<Pilot-plant PFD for arsenic removal>

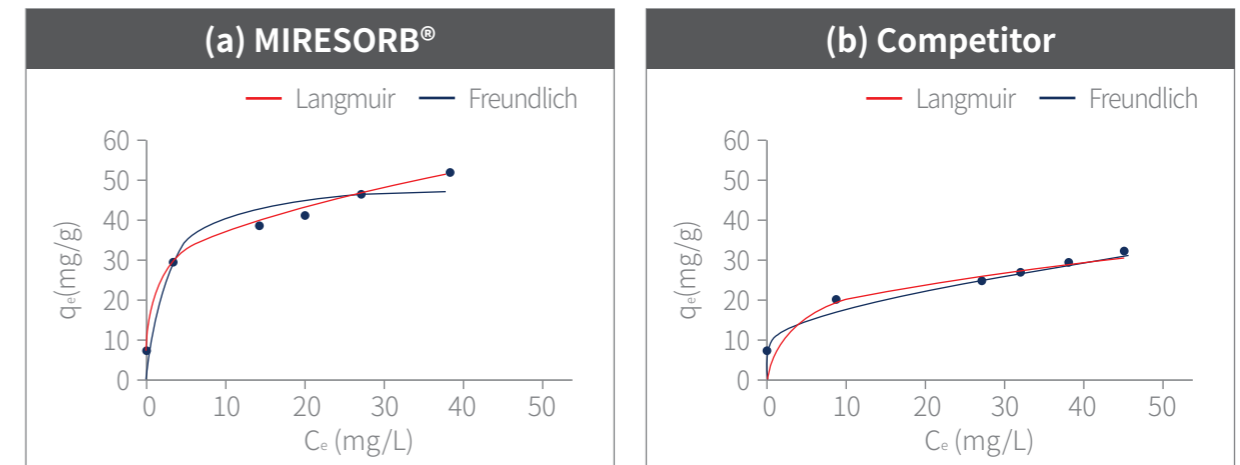
MIRESORB® arsenic removal and adsorption performance evaluation

Adsorption kinetics evaluation



Adsorbent	Pseudo First-Order			Pseudo Second-Order		
	q _e (mg/g)	K _i (1/h)	R ²	q _e (mg/g)	K _s (g/(mg·h))	R ²
MIRESORB™	19.76	0.227	0.990	20.88	0.015	0.958
Competitor	19.58	0.076	0.977	21.74	0.005	0.986

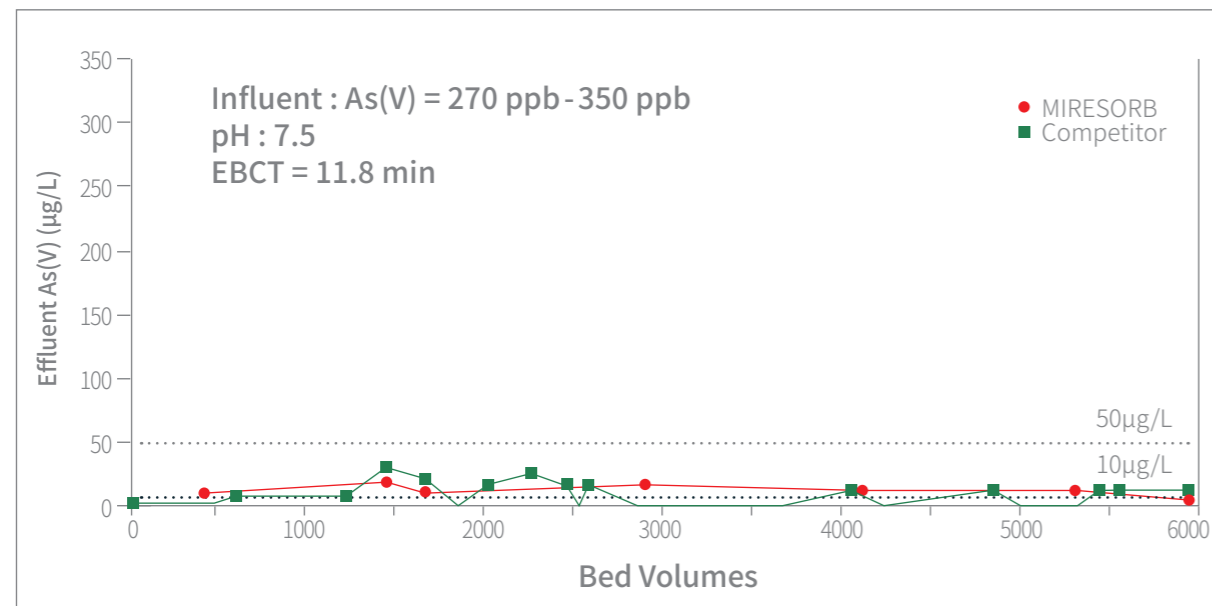
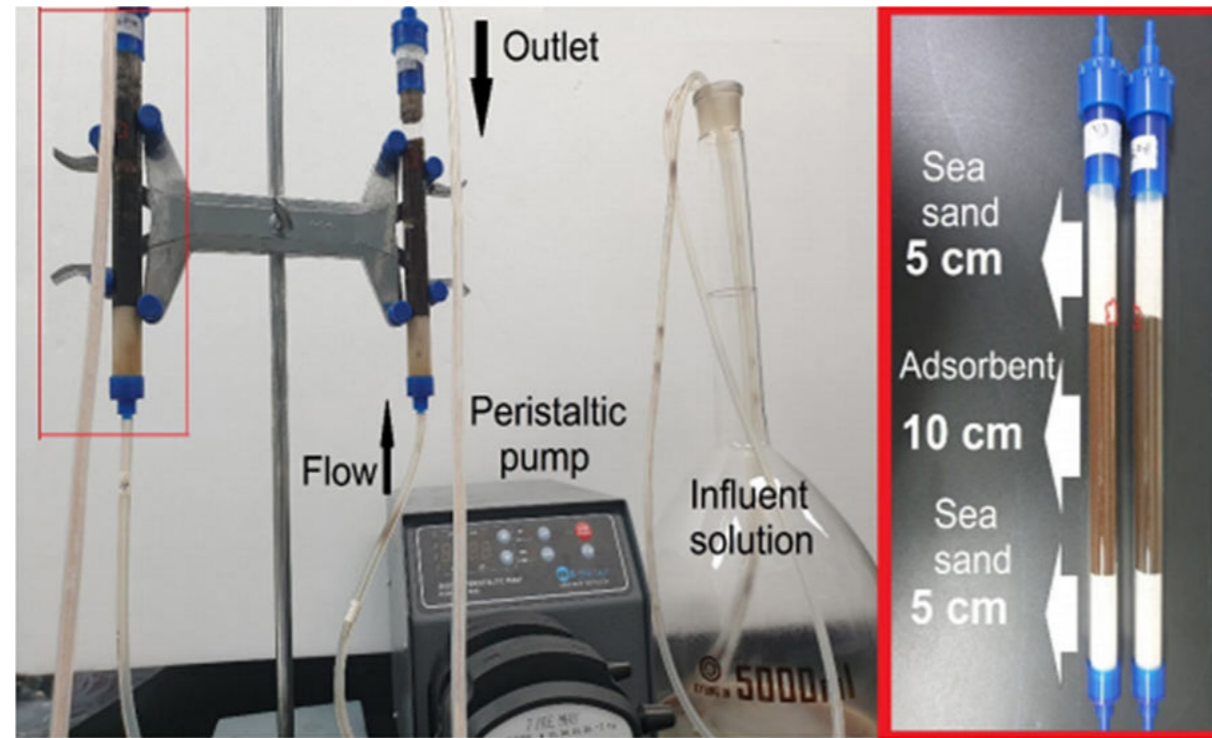
Isothermal adsorption evaluation



Adsorbent	Langmuir Isotherm			Freundlich Isotherm		
	q _m (mg/g)	K _L (L/mg)	R ²	K _F ((mg/g)·[mg/L] ^{1/n})	n(1/n)	R ²
MIRESORB™	50.387	0.38	0.91	19.04	3.68(0.27)	0.98
Competitor	29.07	0.33	0.94	12.00	4.35(0.23)	0.95

Ref. : "Arsenic (V) Removal by an Adsorbent Material Derived from Acid Mine Drainage Sludge", Applied Sciences(2021) Vol.11, No.1 47

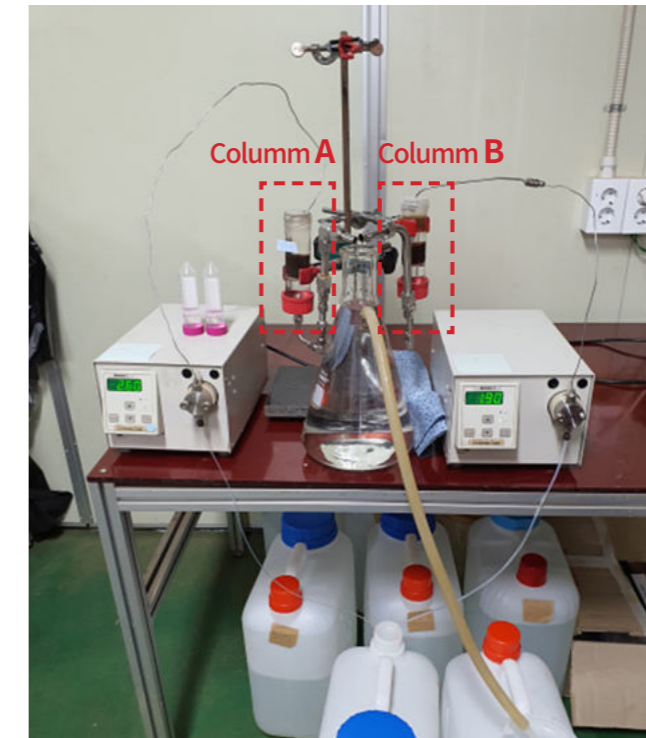
MIRESORB® column test for low concentration arsenic removal



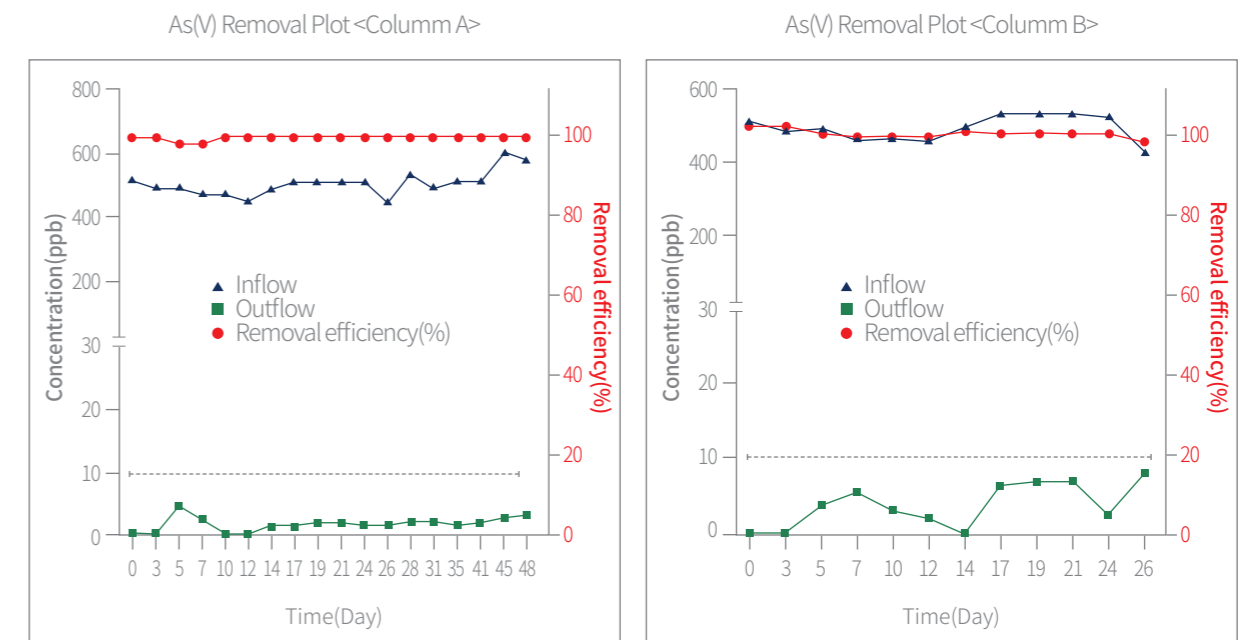
In the case of column experiments using actual mine drainage, adsorption performance is maintained even after 5,800 bed volume is removed.

Ref. : “Arsenic (V) Removal by an Adsorbent Material Derived from Acid Mine Drainage Sludge”, Applied Sciences(2021) Vol.11, No.1 47

MIRESORB® column test for high-concentration arsenic removal



<Lab test using arsenic contaminated water>



In the case of performance experiments using actual mine drainage, the AS(V) removal rate is maintained at 99% or more for 48 days.
Contaminated water used (Raw water : Mine drainage in area B)
- Arsenic (V) contained : approximately 500 ppb

MIRESORB® pilot test for low-concentration arsenic and zinc removal

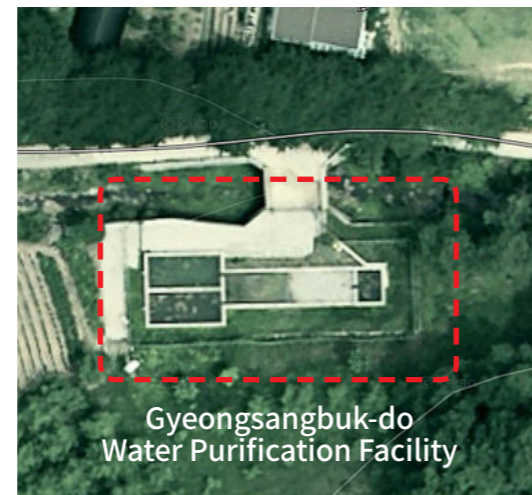


Inflow raw water condition

- Demonstration test capacity : 100ml/min
- Arsenic (V) inflow concentration : 70 ~ 400 mg/L
- Test bed size : 90L(Packed, 50L)
- Raw water inflow method : Using a metering pump



Raw water inflow site



Gyeongsangbuk-do Water Purification Facility

<Examination Point>



Installation of demonstration equipment



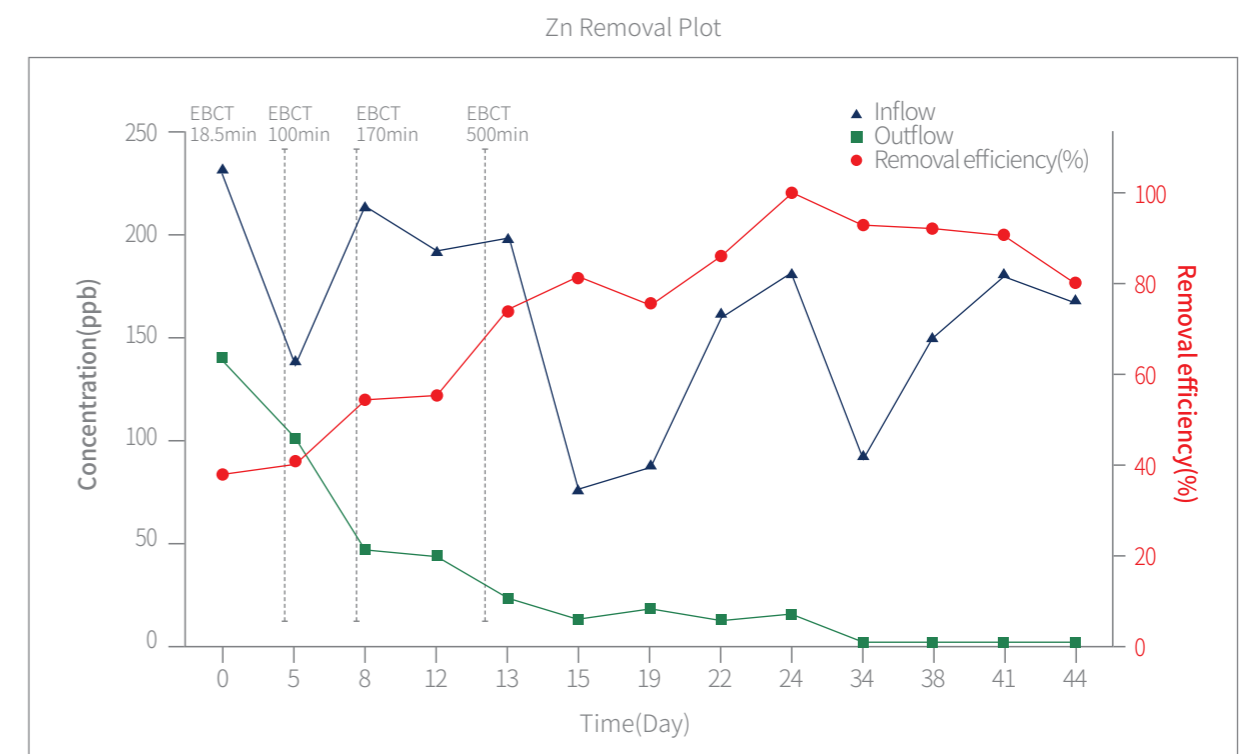
<Arsenic removal efficiency analysis>

ICP-AES, 5110 SVDV, Agilent
microscopic inorganic element analysis

MIRESORB® pilot-test results for low-concentration arsenic removal and zinc removal



< Removal efficiency of Arsenic (IV) with time(day) in pilot test >



< Removal efficiency of Zinc(II) with time(day) in pilot test >

MIRESORB® on-site application and delivery site

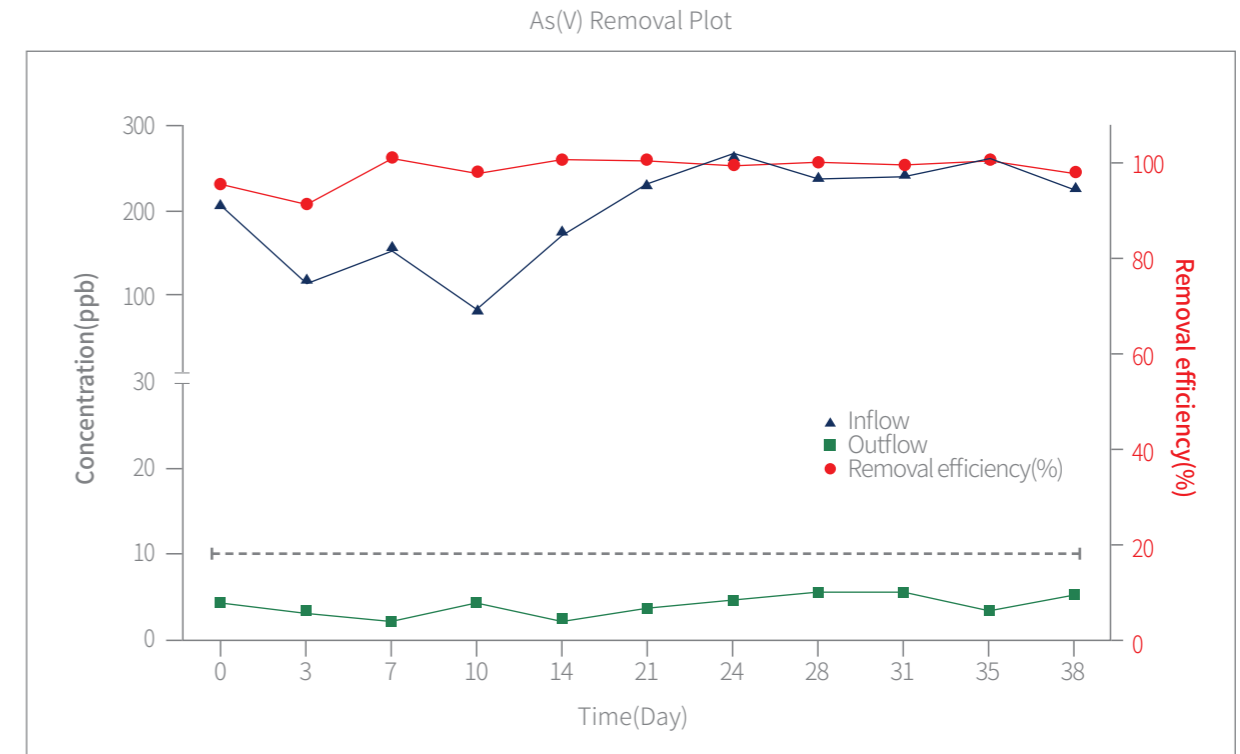
※ Arsenic and Heavy Metal Process in North Gyeongsang abandoned gold mine

- A system that coarse contaminants in raw water are removed by precipitation in storage tank, and arsenic and other heavy metals are removed by adsorption in adsorption tank, finally being discharged.
- The contaminated water is fed to the storage tank through bottom line by hydrostatic pressure and it comprises several layers such as perforated tube, limestone, arsenic-removing adsorbent and sand from the bottom.

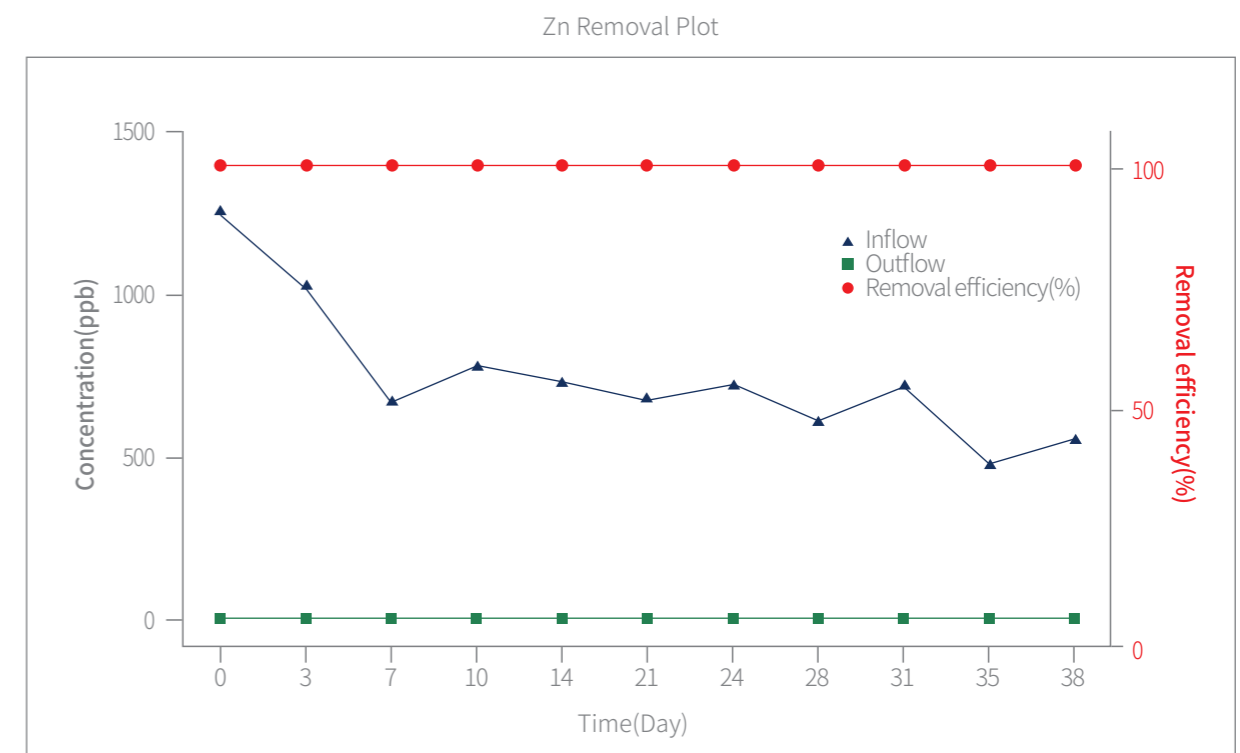


MIRESORB® results of on-site application test operation

Arsenic removal efficiency with time(day) for inflow, and resulting outflow concentrations



Zinc removal efficiency with time(day) for inflow, and resulting outflow concentrations



Analysis report

Analytical device - ICP MS

1. Pilot Test Part 7 2. Trial run result Part 7 3. Lab Test Part 5

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Inductively coupled plasma atomic emission spectrometer test report

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Inductively coupled plasma mass spectrometer test report

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Inductively coupled plasma mass spectrometer test report

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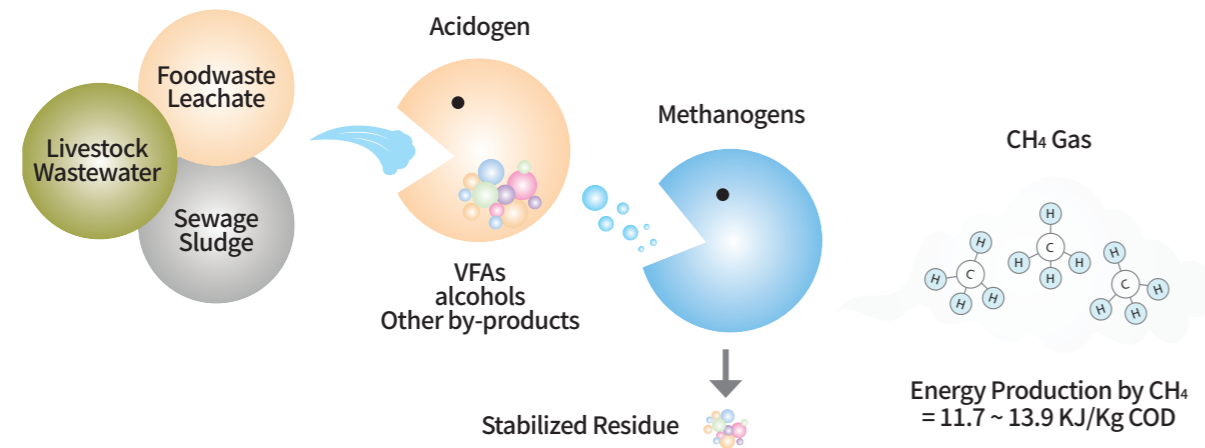
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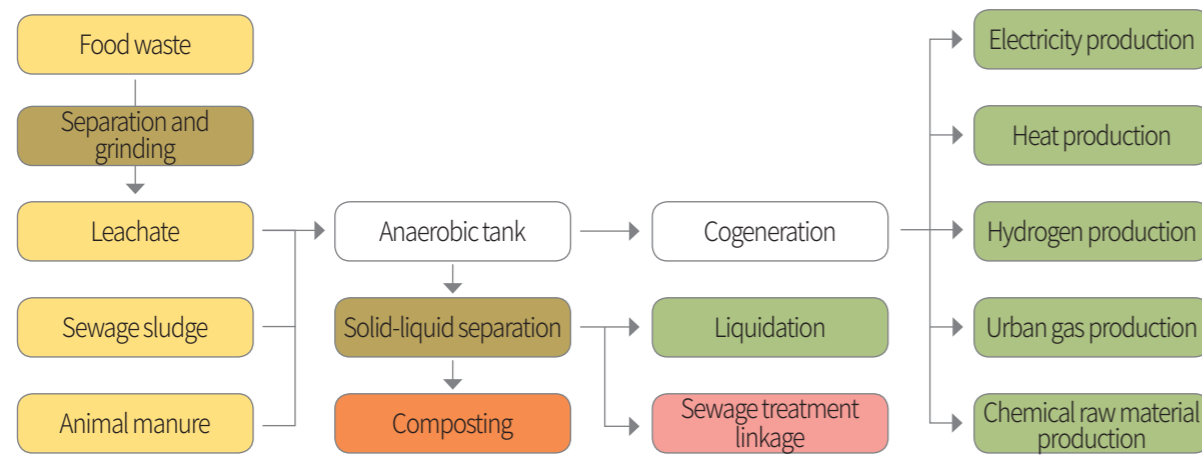
DSULFER®

Iron hydroxide stabilizer for inhibiting
hydrogen sulfide production in a digester

Biogasification



Basic biogas plant process

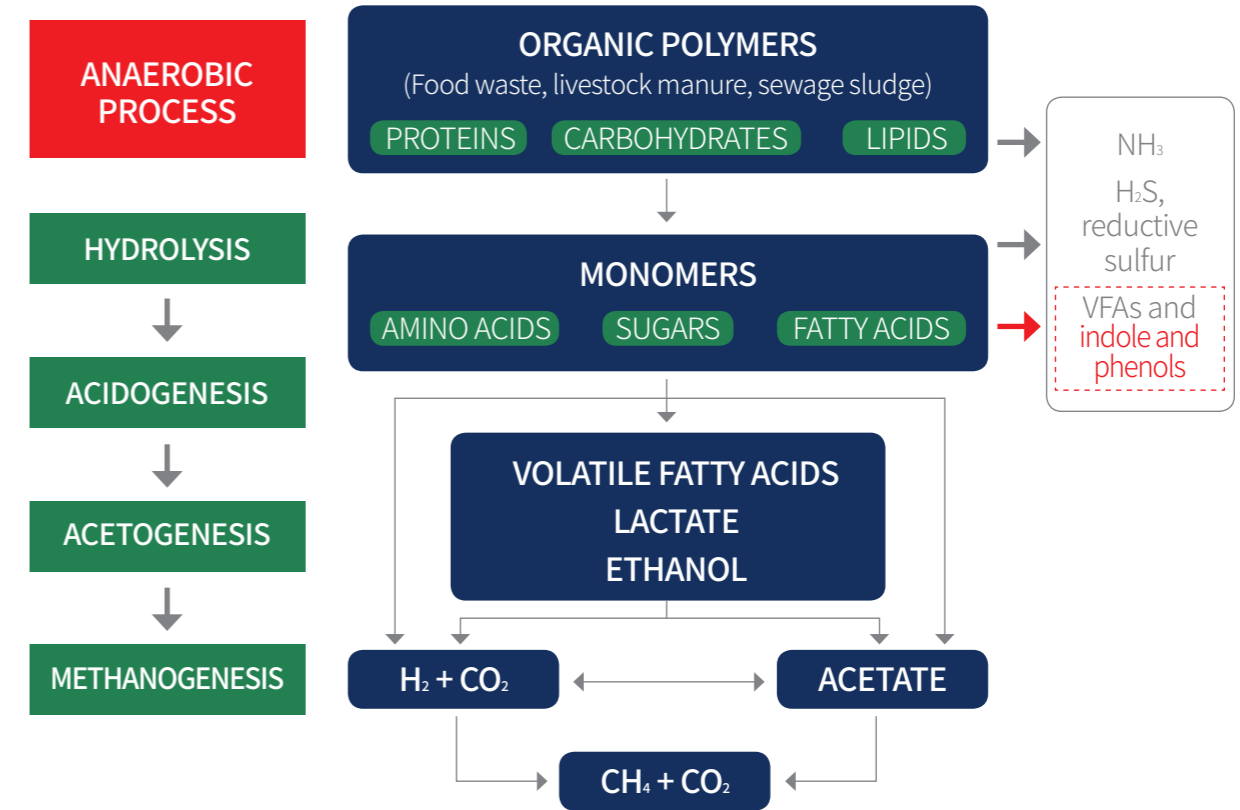


Reaction process	Reaction formula	Microorganism
Hydrolysis stage	Organic substances → Sugars, Glycerin, Fatty acids, Amino acids	Microbial in vitro enzyme
Acid production stage	Sugar → Acetic acid, Propionic acid, Butyric acid	Organic acid-producing bacteria
Methane production stage	$\text{CH}_3\text{COOH} + 2\text{H}_2\text{O} \rightarrow 2\text{CO}_2 + 8\text{H}$ (Oxidation reaction) $4\text{H}_2 + \text{CO}_2 \rightarrow \text{CH}_4 + 2\text{H}_2\text{O}$ (Reduction reaction)	Methane microorganism
Hydrogen sulfide production stage	$\text{SO}_4^{2-} + \text{organic matter} \rightarrow \text{S}^{2-} + \text{CO}_2$ $\text{S}^{2-} + 2\text{H}^+ \rightarrow \text{H}_2\text{S}$	Sulphur-reducing bacteria (SRB)

Iron hydroxide-iron chloride hydrogen sulfide reaction formula

- $2\text{Fe}(\text{OH})_3 + 3\text{HS}^- + 3\text{H}^+ \rightarrow 2\text{FeS} + \text{S} + 6\text{H}_2\text{O}$
- $2\text{Fe}(\text{OH})_3 + 3\text{HS}^- + 3\text{H}^+ \rightarrow \text{Fe}_2\text{S}_3 + 6\text{H}_2\text{O}$
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- $2\text{Fe}(\text{OH})_3 + 3\text{H}_2\text{S} \rightarrow \text{Fe}_2\text{S}_3 + 6\text{H}_2\text{O}$

By-product from anaerobic digestion process



Hazardous substances that inhibit anaerobic digestion are free ammonia, free hydrogen sulfide, heavy metals, etc., and ammonia toxicity occurs most frequently.

Hazardous substances caused by anaerobic digestion process

Ammonia	Produced by deamination reactions during protein degradation in anaerobic reactor Toxicity above NH ₃ 80 mg/L, NH ₄ ⁺ 1,500 mg/L
Sulfide	Less than 20 mg/L is required for optimal methane activity, but decreases if H ₂ S is 50 to 250 mg/L
Volatile acid	High concentration pH decreases, inhibition of reaction above 1,000 mg/L of propionic acid
Soluble sulfur compound	Maximum non-toxic concentration : 200 mg/L Heavy metals such as copper, zinc, and nickel are toxic even at low concentrations. when there is sulfide (H ₂ S gas, HS ⁻ , precipitated sulfur compounds), it is partially precipitated
Antibiotic substance	Antibiotics, cleaning agents, etc. contained in livestock manure

Digestive stabilizer (DSULFER®) features of E&Chem Solution Corp.



Prevention of corrosion damage

Corrosion can be prevented by using DSULFER® to prevent corrosion caused by the spread of sulfuric acid particles by the air supplied during the desulfurization process.



Handling stability

Iron Hydroxide is a non-toxic material and it can be easily handled with simple equipment and it doesn't deteriorate by mixing the water and doesn't require expensive weighting equipment. so it can protect the entire plant from corrosion damage and makes it to use easily without any training or special equipment



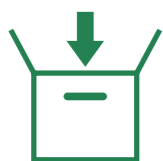
Explosion Prevention by mixing

When air is injected for desulfurization and mixed with methane, there is a risk of explosion caused by unintended air inflow but the DSULFER® is very safe.



Convenience of measurement

DSULFER® is excellent in biological stability because it is precipitated by combining with sulfur ions generated in a digester, etc., and it is fully activated a few days after injection. so hydrogen sulfide production is inhibited by the concentration of sulfur contained in the substrate.



Clean, by-product-free and easy handling

DSULFER® can be supplied in a powdery tone bag or sludge state. so it is very easy to use measure because it supplies a digester in conjunction with organic material in the most biogas plant processes



Load reduction of desulfurization equipment

DSULFER® can efficiently remove most sulfur ions from the initial digester. Therefore, expensive equipment isn't required for concentration load in the post-sulfur process. so cost can be reduced.

Comparison of E&Chem Solution Corp. digestive stabilizers

The digestive stabilizer (DSULFER®) is a principle that inhibits the generation of hydrogen sulfide by precipitating the sulfur ions generated by sulfur reducing bacteria (SRB) during the digestion process by microorganisms in the anaerobic digester easily reacting with amorphous iron hydroxide. For this purpose, it is used in an intermediate storage tank, acid fermentation tank or digester.

Unlike other additives, the digester stabilizer (DSULFER®) has an excellent effect in suppressing the generation of hydrogen sulfide because it is added together with the hydrogen sulfide contained in biogas. And the additionally included divalent trace metal has the effect of enhancing the activity of microorganisms.

In addition, unlike other companies' products, it has very good effects in harmful substances, corrosiveness, handling and storability.

Comparison of properties by additive

	E&Chem Solution Corp.	Iron chloride	Iron oxide
Harmful substance	Very good	Heavy metals, chlorine ions	Good
Harmful by-product	None	Heavy metal	None
Causticity	Very good	Very bad	Good
Sludge generation	Middle	Middle	A lot
Handleability	Very good	Very danger	Good
Methane concentration	Very good	Very good	Good
Digester biocompatibility	Very good	Faulty	Good
Removal efficiency	Very good	Very good	Good
Risk of explosion	Very good	Very good	Very good
Reaction velocity	Good	Very good	Faulty
Storability	Good	Danger	Good

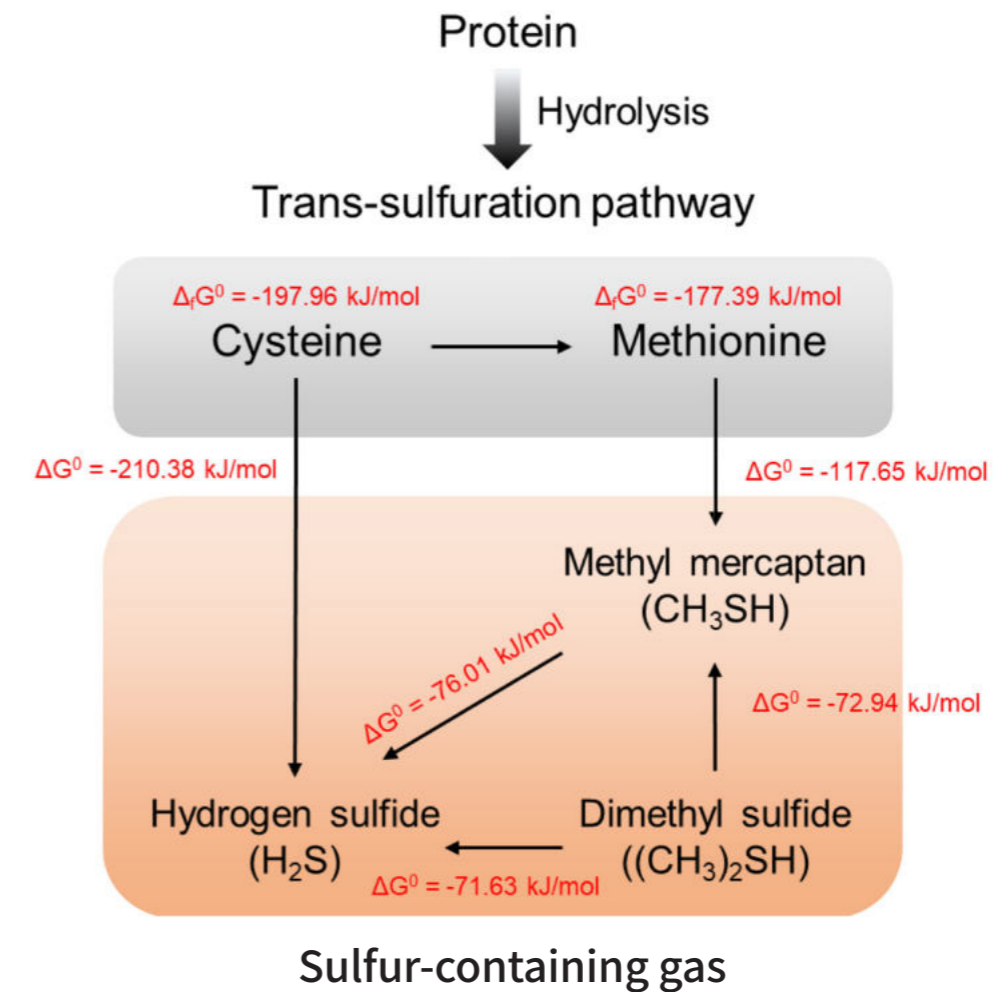
Ingredient table

Digestive stabilizer (DSULFER®) is a product with high purity through a strictly controlled production process, and it provides excellent hydrogen sulfide production inhibition performance and safety in the digester.

Particulars						
Ingredient	Iron(III) hydroxide					
Configuration	Fe(OH) ₃					
Appearance	Reddish brown powder or sludge					
Physical property	Item	Maximum value	Minimum value			
	Weight ratio of iron to dry ingredients [wt%]	60	40			
	Weight ratio of iron hydroxide to dry ingredients [wt%]	95	75			
	Bulk density [g/cm ³]	0.8	1.2			
	Water content [%]	10	5			
Trace element content	Kind	Content (mg/kg)	Elution	Verification method	Fertilizer law standard (mg/kg)	
	This product complies with the guidelines of the Enforcement Decree of the Comparative Management Act [Attachment 1] (Article 10, Paragraph 1).					
	Arsenic	0	ND*	atomic spectroscopy / process test method	5	
	Cadmium	0	ND		0.5	
	Mercury	0	ND		0.2	
	Lead	0	ND		15	
	Chromium	0	ND		30	
	Copper	-*	ND		50	
	Zinc	-	ND		130	
Nickel	0		5			

* ND: Within the criteria, -: Need to be tested for livestock manure fermentation fluid

BMP(Biochemical methane potential) Test

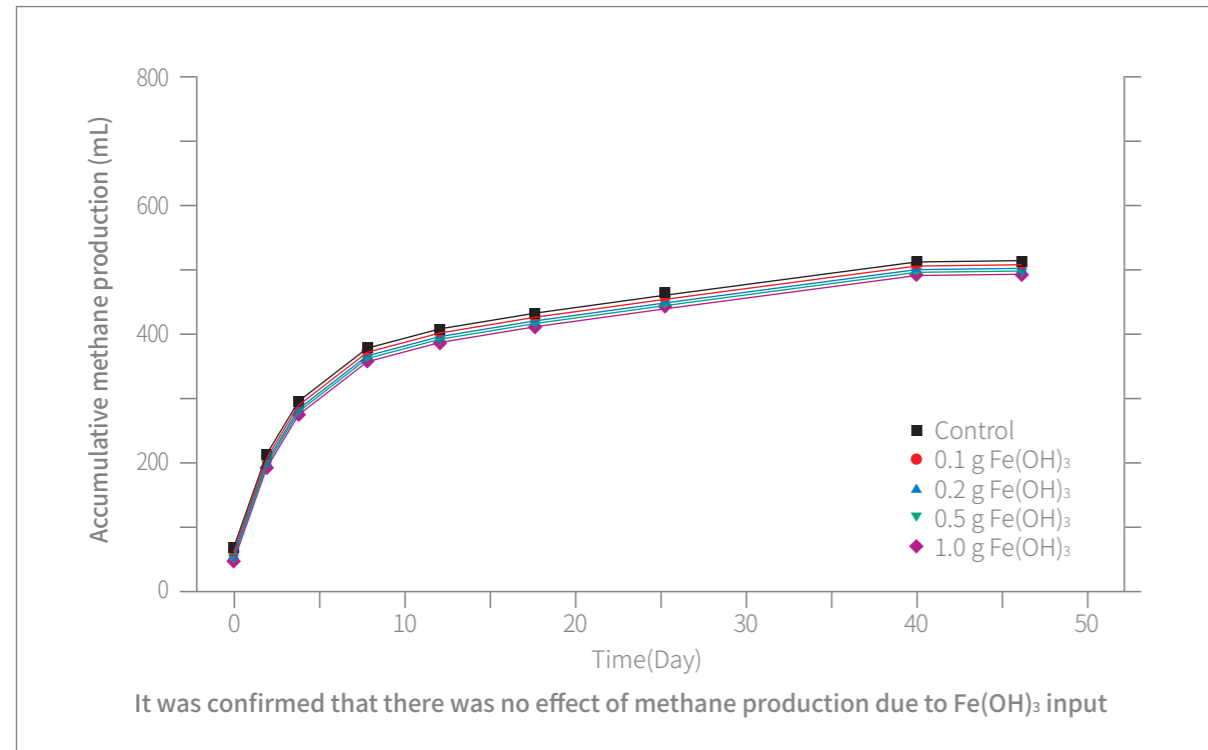


Conditions for testing

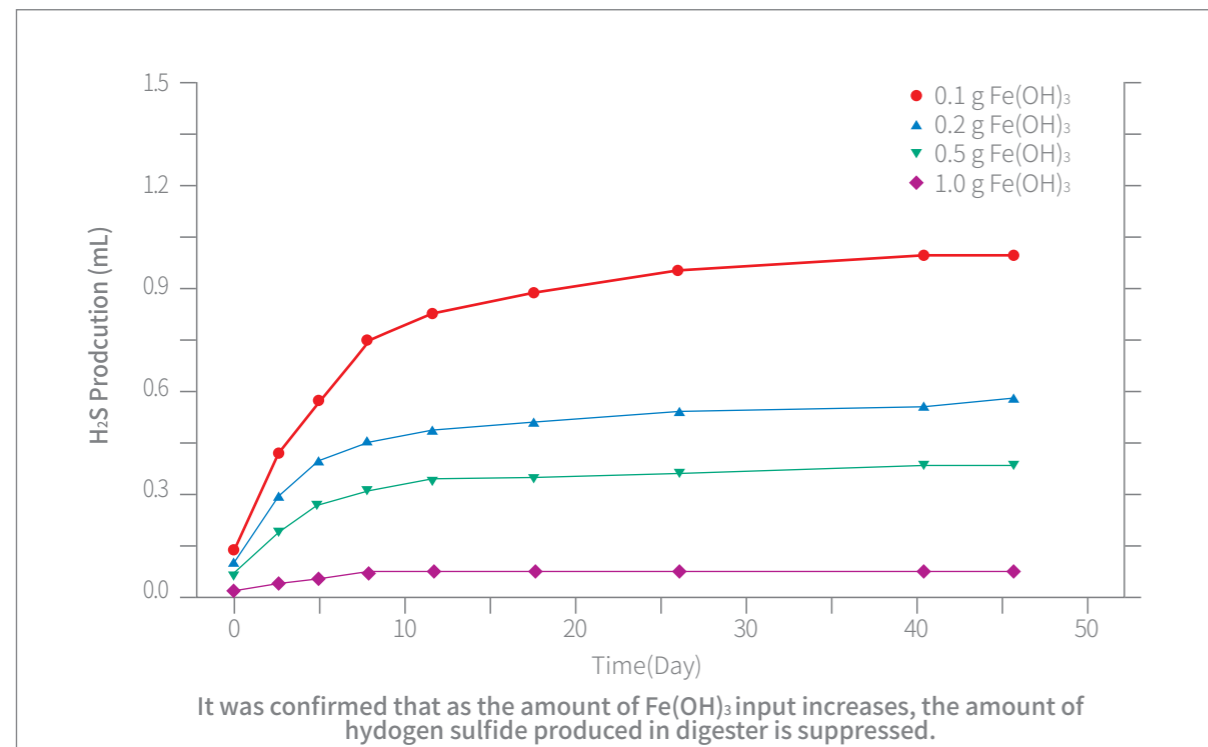
- **Operation type** : Batch
(Anaerobic digester at city D Sewage Process Plant)
- **Test capacity per batch** : 150ml
- **Concentration of injection of digestive stabilizer** : 0g, 0.1g, 0.2g, 0.5g, 1.0g
- **Microbial concentration** : 20.4 ± 0.1 g VS/L
- **Concentration of organic matter (COD)** : 5.0g TCOD/L
- **Sulfur concentration** : 20.0 mg/L
- **Digester temperature** : 37 ± 1°C
- **pH** : 7.5
(Experiments with alkalinity of 5 g/L as CaCO₃)
- **Medium** : BA medium



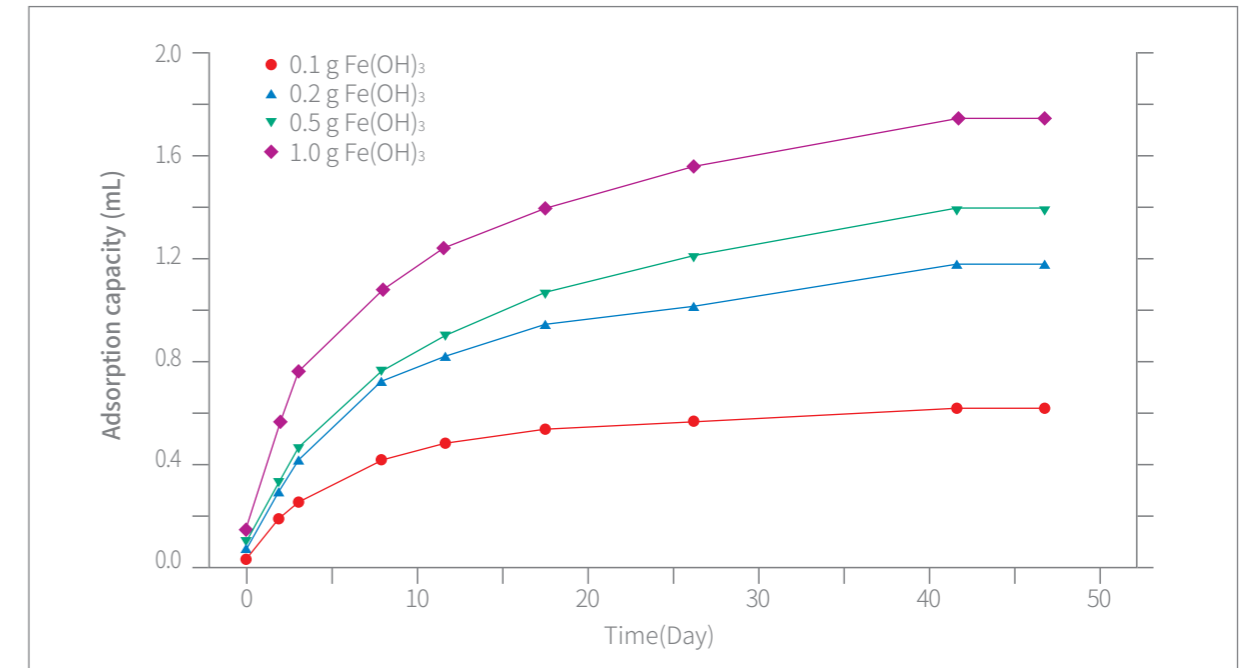
Methane production by concentration of digestive stabilizers



Hydrogen sulfide production by the concentration of digestive stabilizers

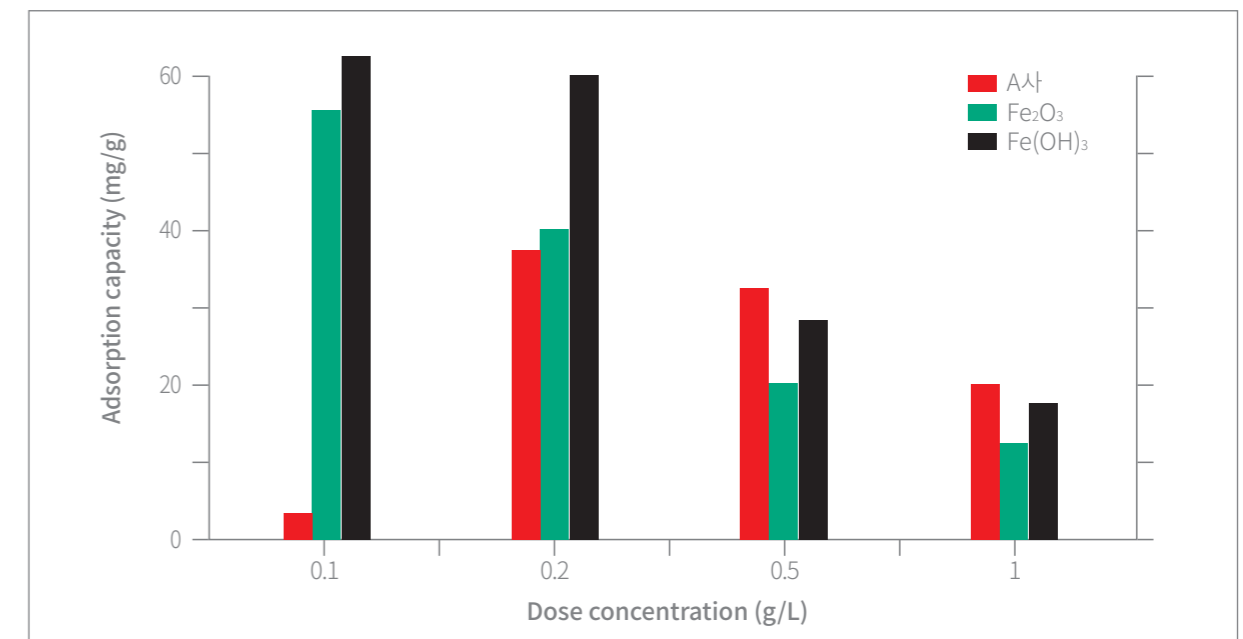


Hydrogen sulfide adsorption test of digestive stabilizer



Dosing amount	Adsorption capacity (mg/g)
0.1 g/L $\text{Fe}(\text{OH})_3$	63.4
0.2 g/L $\text{Fe}(\text{OH})_3$	60.7
0.5 g/L $\text{Fe}(\text{OH})_3$	28.4
1.0 g/L $\text{Fe}(\text{OH})_3$	17.4

Comparative testing of hydrogen sulfide adsorption capacity of digestive stabilizers



As a result of the experiment of evaluating the adsorption capacity of hydrogen sulfide through BMP tests, it was confirmed that $\text{Fe}(\text{OH})_3$ had the highest adsorption amount of hydrogen sulfide per unit mass.



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