

Hydroscav H₂S & RSH REMOVAL

Product Description

Hydroscav is a water-soluble blend of amine resin solution in a complex alcohol system. This product is extremely effective in removing H₂S from gas, water, crude and fuel oils.

Despite what the name infers, Hydroscav is **not** a scavenger in chemical terms. Rather than 'scavenging' H_2S , it converts H_2S & RSH into a stable non hazardous Sulphate salt (SO_4^{-2}). This eliminates several issues in the process system, often associated with triazine / formaldehyde based scavengers.

Hydroscav is also effective in removing mercaptans from liquids in the same manner, however, the reaction time required to remove the mercaptans is slightly longer.

Product Characteristics

Hydroscav converts H₂S & RSH into a non toxic sulphate salt and water:

- 1. **Hydroscav** will chelate the H₂S & RSH molecules
- It then converts the molecules into a liquid salt SO₄-2
- 3. This solution then attaches to the Amine molecule
- 4. The water based molecule then mixes with the produced water
- 5. The result is a water salt with a very low COD
- 6. The water salt is stable and the reaction is irreversible

$$R (Hydroscav) + H_2S & Mercaptans$$

$$R' SO_4^{-2} + H_2O$$

Dosage Rates

Gas: 1ppm of H₂S requires 0.2ppm of *Hydroscav* (measured in gas phase)

Water: 1ppm of H₂S requires 3 ppm of *Hydroscav* (measured in liquid phase)

Crude: 1ppm of H₂S requires 3–6 ppm of *Hydroscav* (measured in liquid

Fuel Oils: phase) 1ppm of H₂S requires 3-4 ppm of *Hydroscav* (measured in

liquid phase)

Removal of mercaptans: for each ppm of mercaptans you will require between 3–5ppm of *Hydroscav* depending on the sulphur speciation of the mercaptans.



Application

Like any chemical, application is the key. Applied with a good mixing mechanism, the performance of the chemical will be enhanced. For oil and gas production, *Hydroscav* can be injected down hole, and it is not affected by high temperatures. For treatment of crude, condensates or fuels oils, ideally Hydroscav should be injected at or near the inlet valve of the transfer pump, as theasset is being transferred from one tank to another. Alternatively, Hydroscav can be added to a storage tank and circulated, however, this may require longer time at possibly higher dosage rates depending on the capacity of the tank and the effectiveness of the circulation pump.

Gas

Hydroscav needs to be injected under pressure to ensure good migration. The reaction is instantaneous in the gaseous phase and the separation of the newly formed Suphate salts is very fast. The only drawback is that **Hydroscav** cannot be used where there are high levels of CO2 as the CO2 will reduce the chemicals effectiveness.

Water

Hydroscav is extremely effective in removing H_2S and mercaptans from water. The effect of the chemical (both used and unused) on water is very interesting. The chart below shows changes to the water after different rates of injection.

Sample Description	рН	Conductivity uS/cm	Turbidity (NTU)	Silica as SiO2 (ppm)	Total Iron (ppm)	H2S in Liquid Phase (ppm)
Raw Water Feed	7.67	29, 400	211	7.5	3.0	5.5
Hydroscav @ 15 ppm	8.35	29600	203	7.6	0.8	Nil
Hydroscav @ 25 ppm	8.45	29300	198	7.3	0.8	Nil
Hydroscav @ 30 ppm	8.51	29200	199	8.7	0.5	Nil
Hydroscav @ 40 ppm	8.59	29000	199	10.7	0.5	Nil
Hydroscav @ 50 ppm	8.52	30000	199	8.5	0.5	Nil
Hydroscav @ 60 ppm	8.52	30100	180	6.0	0.8	Nil
Hydroscav @ 80 ppm	8.67	30000	166	3.8	0.8	Nil
Hydroscav @ 120 ppm	8.65	30000	133	10.0	0.8	Nil
Hydroscav @ 160 ppm	8.72	30000	75	9.0	0.5	Nil
Hydroscav @ 200 ppm	8.84	30000	2.93	9.9	0.3	Nil
Hydroscav @ 1000 ppm	10.02	29800	3.78	Nil	0.0	Nil

Total Iron is dramatically reduced even at low dosage whereas H_2S is reduced to NIL as both undissociated H_2S and dissociated HS ions are neutralized by **OURScavanger**. This is important as it represents a drastic reduction in Total Suspended Solids (TSS).



The pH remains within 8 - 9, which indicates that the chemical does not bestow higher alkalinity on the produced water. This is an important finding where there are high levels of TSS.

Crude & Fuels

Hydroscav converts the H₂S and Mercaptans into sulphate salts. The chemical then separates from the hydrocarbons due to the higher specific gravity. For extremely heavy crudes, adjustments to the chemical can be made to increase the specific gravity and speed up the separation process. **Hydroscav** will not affect the hydrocarbons in any way.

Naphtha Sample: H2S = 112ppm; RSH = 191 (ASTM D3227)

Hydroscav @	100 ppm		400 ppm		800 ppm	
	H2S	RSH	H2S	RSH	H2S	RSH
SAMPLE	112	191	112	191	112	191
IMMEDIATE	63	131	59	96	42	80
AFTER 4HR	49	97	14	63	NIL	39
AFTER 8HR	40	73	NIL	34	NIL	22
AFTER 12HR	33	62	NIL	24	NIL	3.4

The amount of Hydroscav required to remove H_2S from any system is typically around 30% less than that of a triazine based Scavenger. The reaction is stable and the result is a non toxic sulphate salt that can be easily handled.

Frequently Asked Questions

How fast is Hydroscav able to removal of H_2S ? The reaction with Hydroscav and H_2S is instant once contact is made. RSH reaction time can take longer depending on the RSH species. Typically, the treatment time required is more a function of the efficiency of the application and nature of the asset that is being treated.

Are light thiols or other sulphur compounds effected or only H_2S ? All sulphur compounds react with the chemical.

What is the level of Effectiveness of Hydroscav to remove H_2S / RSH in the liquid phase? 650 mg/L (combined H_2S & RSH) in liquid phase was tested and was successful. No upper limit has been established.

How do you remove the used and unused *Hydroscav* **from Fuel Oil?** The product is 100% miscible with water and this is a SINGLE PHASE system only. In liquid hydrocarbon applications, the sulphates which are created by *Hydroscav* stay in the water phase and are separated from hydrocarbons during normal separation.



What is the effect of the scaling potential of the system? *Hydroscav* will sightly increase the pH of the water, therefore, analysis has to be carried out to determine the effect of this increase in pH, particularly where there are high levels or suspended solids or calcium carbonates in the produced water.

What is the potential of scaling with the reacted or un-reacted *Hydroscav*? NONE - No potential scaling in ordinary environments. The increase of SO₄⁻² in the water phase will depend on the mole-mole equilibrium between quantity of H2S / RSH and *Hydroscav*. Cat ions such as Barium, phosphorous and others associated SO4-2 Cat ions may cause fouling or deposits once it has reached maximum solubility. Excess THIScavenger will not cause any deposits.

Do you use an air oxidiser? If so, how can you ensure that you have enough O₂ under anaerobic conditions? - No, enclosed system; hence no Oxygen ingress.

What is formed at higher temperatures? It does not decompose thermally within the operating temperature range up to 300 Deg. C

Did you find increased sulphate levels in the water after treatment? Yes - the amount of increase is directly related to the amount of H_2S / RSH treated by the chemical.

How effective does the mixing have to be? No special mixing tooling required with water. In gas lines, inject with sprayer nozzle under positive pressure. For heavy crudes and HSFO, it is recommended to inject as early as possible or at pump inlet to ensure efficient mixing.

How fast or complete is the phase separation afterwards? There is complete separation from the hydrocarbon phase, and the speed of the separation can be adjusted by adjusting the specific gravity of the chemical.

Disclaimer

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