# **TSA-100**

# A TOTAL SULFUR ANALYZER

# For Continuous Sulfur Monitoring in Liquid and Gas

Applied Analytics **TSA-100**, designed for accuracy, versatility, fast-response, and ruggedness in any range of total sulfur measurements. This innovative multi-component analyzer has critical advantages over older standards such as the lead acetate tape method, a high-maintenance process with slow measurements, cumbersome adjustments, and toxic consumable. Most significantly, the **TSA-100** is engineered for insusceptibility to component cross-interference, meaning that the analyzer can provide accurate total sulfur measurements when interfering compounds like H<sub>2</sub>S, SO<sub>2</sub>, and mercaptans are all present in the sample.

The system consists of four major subunits: 1) a xenon light source, which sends light through fiber optic cables into the 2) flow cell, in which the unique characteristics of the sample define its interaction with the light; the light then enters a 3) holographic grating, which separates white light into its components such that the 4) diode array detector can measure the intensity of radiation at each wavelength. These measurements report on the absorption of light by the various components in the sample, and thus reveal the concentrations of the components. The analyzer is solid state (no moving parts), making TSA-100 suitable for even the harshest industrial environments. Using the full high-resolution spectrum from the diode array, our analyzer can easily switch between extreme wavelength ranges to provide instantaneous and continuous measurements at 0-10 ppm and 0-100% all in the same instrument.

AAI offers effectively two different methods of determining total sulfur. In the *DIRECT METHOD* (H2S-220), the analyzer mathematically derives the concentration of each sulfur compound from the full, high-resolution absorbance spectra it obtains from the original process stream. There is no need to mechanically consolidate all sulfur into one specific component, as by combustion in the standard method. The concentrations of sulfur based components like H2S, COS, and mercaptans are computed individually from the spectrum and added together, such that the computer outputs a total sulfur reading. The direct method effectively measures total sulfur in the original process stream as multiple components.

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The second is the conventional *STANDARD METHOD* (**TSA-100**), which involves placing the hydrocarbon sample in a hot combustion tube in an oxygen-rich environment (ASTM D 5453 – 06) The sulfur is oxidized to  $SO_2$ . The analyzer uses the absorption of the single component ( $SO_2$ ) to determine its concentration, and the total sulfur. The **TSA-100** provides the ideal  $SO_2$ - monitoring solution in case the standard method is your route. The **TSA-100** is recommended for streams containing high-level aromatics.

Comparison to paper tape: The paper tape method has several important shortcomings which help elucidate why it may be an outdated system. In this method,  $H_2S$  and lead acetate react to form a color spot of lead sulfide precipitate on the tape. The rate that the color changes is related to  $H_2S$  concentration.

The most obvious flaw with the paper tape is an economic one: as a consumable, the tape must be continually purchased and replaced. Furthermore, due to the hazardous nature of lead acetate (causes fetal mortality, possibly carcinogenic), the tape must be returned after use to the manufacturer for disposal. The purchase and disposal of paper tape become constant expenditures which, over the long run, are of great financial concern.

In addition, a moving parts sulfur-detection system relying on paper tape requires substantial maintenance, frequently incapacitated by sample saturation or tape replacement. The regular response time can be as long as several minutes, depending on the sulfur concentration. Lastly, this method is quite vulnerable to inaccuracies stemming from crossinterference among the sample components.

The advantages of the TSA-100 over paper tape are economically, environmentally, and technologically significant. The spectrophotometric principles of the analyzer eliminate the need for costly, polluting consumable like lead acetate; the only part that would ever need to be replaced is the inexpensive and long-lasting xenon lamp. The analyzer is solid state (no moving parts), and since it connects to the sample through fiber optic cables, it can be installed far from any explosive or poisonous gases.

While lead acetate tape gives slow, discrete measurements, the TSA-100 provides instantaneous and continuous measurements. Using the full high-resolution spectrum obtained by the diode array, the analyzer can easily switch between extreme wavelength ranges, allowing measurements of 0-10 ppm and 0-100% in the same instrument. Comparable range flexibility using paper tape might require dilutions, producing an additional sampling concern and possibly inviting inaccuracies.

The cross-interference of  $H_2S$ , mercaptans (the functional S-H group), SO<sub>2</sub>, and other sulfur compounds is a concern which makes the paper tape method incompatible with the current technological demands of industry. AAI's innovative analysis corrects for this interference, allowing simultaneous measurement of all sulfur components and the *total sulfur*.



Figure 1: Absorbance spectra of SO<sub>2</sub>, different concentrations



Figure 2: Absorbance spectra of  $H_2S$ , different concentrations



Figure 3: Absorbance spectra of H<sub>2</sub>S and methyl mercaptan



## Total sulfur by conversion to SO<sub>2</sub> (combustion sampling system)

The sulfur based compounds in the sample stream are pyrolyzed to give  $SO_2$ . The flow of the sample gas and  $O_2$  are regulated to keep the correct mixture for optimal conversation. These gases are mixed in a chamber before they are injected into the furnace where all sulfur compounds are converted into  $SO_2$ . The  $SO_2$  is then transferred into the analyzer's flow cell where it is measured by a UV process diode array spectrophotometer. The  $SO_2$  stream is returned to a waste line that can be vented or sent to the flare. The sampling system allows for automatic zero (blank) with Nitrogen at preset time intervals.





#### **TSA-100 Specifications**

Measurement principle

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Tatal sulfan ha	
Total sultur by	ine sullur
conversion to SO2	stream are
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system)	to keep the
	These gase
	are injected
	compounds
	The SO2 is
	flow cell w
	process dic
	The SO2 st
	can be vent
	he sampling
	zero (blank
Detector	1024 eleme
Light source	Long life Xe
Components	H2S COS r
Flow cell	Fiber optics
	(Fiber lengt
	(flow cell le
Repeatability	±0.5% of so
Measurement range	0-10PPM; (
Accuracy	±1% of sca
Zero drift	2% for 24 h
Response time	20 sec
Ambient Temperature	0-55C(32 to
	-20C to 55
Flow cell	
Temperature	-50 to 3000

Pressure I/O Analog outputs

**Digital communication** PLC Fault relay

UV absorbance The process fluid and a stream of O<sub>2</sub> flow juartz tube that is heated to 1100 °C. nat is generated is then passed to a nere it is measured by UV absorbance. based compounds in the sample . pyrolyzed to give SO2. the sample gas and O2 are regulated correct mixture for optimal conversation. es are mixed in a chamber before they d into the furnace where all sulfur s are converted into SO2. then transferred into the analyzer's here it is measured by a UV de array spectrophotometer. tream is returned to a waste line that ted or sent to the flare. T g system allows for automatic ) with Nitrogen at preset time intervals ents diode array e lamp mercaptans SO2 s transmit light to and from cell th depending on installation) ngth depending on H2S concentration) cale D-100PPM; 0-1000PPM; 0-1%; 0-100% le (or ±1PPM) nour (5% for low PPM range) o130F) standard C (-4 to 131 F) optional

C (-60 to 570F) 200 bar (3000 psig)

two galvanically isolated 4-20mA (additional channels are optional) RS232 Modbus, TCP/IP Modbus OPC server included One SPDT alarm relay

### Spectrometer Wavelength range

Slit Width **EP** resolution

Wavelength Accuracy Wavelength reproducibility Photometric accuracy Photometric noise

Photometric stability

Power

Analyzer Dimensions

Sampling system

#### Area Classifications

General purpose NEMA-4X IP 65 US Class I Div. 2 Groups C&D Zone 2(22) Category G D3 US Class I Div. 1 Groups B,C&D EEx d II C T6 Category G D2 NEMA 6 Class 1 div 1 B,C,D

EMC and LVD Supplies

190-800nm 1nm >1.6 toluene in hexane, ratio absorbance at 269nm / 266nm <±0.5nm (NIST 2034) <±0.04nm (NIST 2034) ±0.005 (NIST '930e) <0.002AU 32 scans at 0 AU at 250nm <0.002AU/h at 0AU at 340nm +-10C 80 to 240 Volts AC 40 to 60 Hz, 20 Watts 16"(H)x14"(W)x8"(D) (41x36x20cm) 24"(H)x24"(W)x12"(D)

Standard Optional (Z purged)

Optional (X purged)

Optional (Enclosure) Complies with all relevant European directives SO2 in N2 gas for calibration N2 for Zero O2 for combustion Air for purging



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