

# **Technical Information and User Manual**

# NE4-H2S-500

# Electrochemical Hydrogen Sulphide (H<sub>2</sub>S) Gas Sensor

For Industrial Applications

Apart from pages 4 and 5 (Specification and Dimensions sections), the data in this document does not constitute a specification but is intended as a guide, informing the instrument designer of the performance characteristics of the sensor which were observed by Nemoto Sensor Engineering scientists.

It should be read in conjunction with the official datasheet for the device, which includes the full technical specification for the NE4-H2S-500 Gas Sensor.

Nemoto Sensor Engineering Co. Ltd. has a policy of continuous development and improvement of its products. As such the specification and data outlined in this document may be changed without notice



## **INTRODUCTION**

Nemoto & Co. Ltd was established in 1941 and continues to develop unique technologies for Safety, Security and Health markets worldwide. Using our unique experience of fine chemical preparation and printing, we were able to enter the gas sensor market in 1979 with a range of high-quality hot-wire type sensors (pellistors). Nemoto is now one of the world's leading manufacturers of chemical sensors and has so far delivered over 30-million devices to the market.

As a result of three years' development at our Tokyo R&D centre, we released our first electrochemical gas sensor in 2000.

The NE4-H2S-500 Gas Sensor is a 3-Electrode electrochemical gas sensor designed for the detection and measurement of Hydrogen Sulphide in the range 0-500ppm, in a wide range of industrial process stream analysis applications. By adhering to industry standards for size and connection orientation, the NE4-H2S-500 can be retrofitted easily to existing product designs.

By using our experience of design for manufacture and our high volume production facilities in Japan and China, we have successfully reduced the cost of the NE4-H2S-500 whilst being able to maintain the highest performance quality.

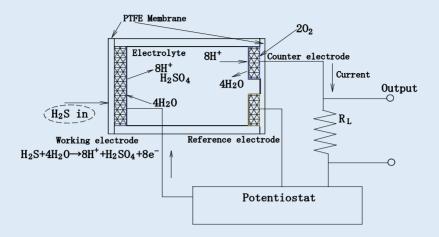
| Variant Part Number | Range     | Signal Output<br>(nA/ppm) | Comments   |
|---------------------|-----------|---------------------------|--|
| NE4-H2S             | 0-100 ppm | 500 nA/ppm                | Lower cost but incompatible with sensors from other manufacturers.   |
| NE4-H2S-100         | 0-100 ppm | 700 nA/ppm                | Standard sensor for Safety<br>applications, compatible with sensors<br>from other manufacturers.                     |
| NE4-H2S-200         | 0-200 ppm | 500 nA/ppm                | For measurements in process streams,<br>or in applications where high levels of<br>H2S are likely to be encountered. |
| NE4-H2S-500         | 0-500ppm  | 200 nA/ppm                | For measurements in process streams.   |

#### NOTE: Nemoto offers 4 variants of the NE4-H2S series sensors:

The performance data published on pages 8-11 of this manual were derived from tests on the NE4-H2S sensor. Although this sensor has a lower signal output than the NE4-H2S-500, the results are considered by Nemoto to apply equally to the NE4-H2S-500 sensor.



## PRINCIPLES OF OPERATION



The NE4-H2S-500 consists of 3 porous noble metal electrodes separated by an acidic aqueous electrolyte, housed within a plastic (PPO) enclosure. Gas enters the cell via a gas phase diffusion barrier (capillary). An electrolyte reservoir ensures an excess of electrolyte is available at all times, and the sensor is vented to ensure that the internal and external pressure of the sensor is always in equilibrium.

In operation, gas enters the cell via the capillary and filter, and comes into contact with the 'working' electrode. Any Hydrogen Sulphide present undergoes the following (oxidation) reaction:

### $H_2S + 4H_2O \quad \rightarrow \quad H_2SO_4 + 8H^+ + 8e^-$

The  $H_2SO_4$  generated dissipates into the electrolyte, whilst the hydrogen ions (H<sup>+</sup>) migrate into the electrolyte within the cell. The electrons (e<sup>-</sup>) generated at the working electrode are collected by the external circuit via a metal strip in contact with it, in the form of a small (nA) electric current.

The reaction at the working electrode is balanced by a reciprocal (reduction) reaction at the 'counter' electrode, using Oxygen from the surrounding atmosphere.

### $O_2 + 4H^{\scriptscriptstyle +} + 4e^{\scriptscriptstyle -} \rightarrow \ 2H_2O$

The electrons consumed in this reaction are supplied by the external circuit via a metal strip in contact with the counter electrode.

Thus water is consumed whilst Hydrogen ions are generated at the working electrode, whilst the water is recreated and hydrogen ions are consumed at the counter electrode. At the same time, the reaction at the working electrode generates electrons, whilst the reaction at the counter electrode consumes electrons. By connecting the working and counter electrodes together via a special circuit, the flow of electrons between the two electrodes may be measured as a nA level current signal proportional to the ppm concentration of Hydrogen Sulphide.

The 'reference' electrode maintains the healthy operation of the cell. It is surrounded by electrolyte, sees no gas and no current is allowed to be drawn from it. Its electrochemical potential hence always remains constant at a level known as the "rest air potential" and this is used to regulate the potential of the working electrode, regardless of the current it is generating during operation. The use of a reference electrode in this way (i.e. threeelectrode operation) helps to extend the working range of the sensor, improves linearity and results in a number of performance benefits compared with similar sensors working with 2-electrodes only.



# **FEATURES**

Electrochemical gas sensors have the following superiority to conventional semiconductor type and hot-wire type gas sensors.

- Linear output in proportion to gas concentration
- High reproducibility
- Highly gas specific
- Unaffected by humidity
- Stable output for long periods
- Low power consumption because no heater is used (can be battery operated)
- Small and lightweight (can be used in portable devices)
- No mechanical structure so highly resistant to shocks and vibrations.

The NE4-H2S-500 has been developed from our accumulation of technologies in production of hot-wire type gas sensors, long research experience into catalysts, fine printing, and assembling of sensors. The NE4-H2S-500 is small and less-expensive, but has high sensitivity, long life, and leak-free performance even under severe operating conditions.

#### Air vent

The electrolyte used for chemical sensors is very hygroscopic, i.e. it has affinity for water, and its volume varies depending on ambient temperature and humidity. This variation causes pressure inside the sensor to rise and fall. In the worst case the electrolyte may leak out of the sensor and damage the circuitry around it.

To prevent this, the NE4-H2S-500 utilizes an air vent capability. This maintains equilibrium between internal and external pressures and allows the sensor to be used in any orientation and under high temperature and humidity conditions.

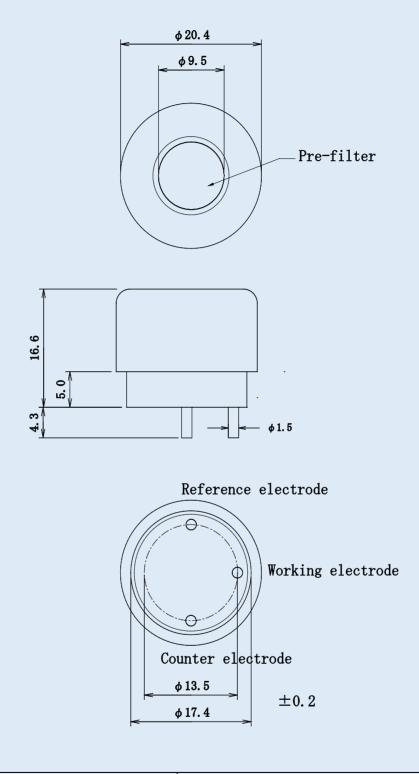


## **SPECIFICATIONS:**

| Output signal                    |   | 200 +/- 50 nA/ppm H <sub>2</sub> S<br>(Linear to 100ppm) |
|----------------------------------|---|--|
| Zero offset in clean air         |   | < +/- 2 ppm equivalent                                   |
| Response time (T90)              | (T <sub>90</sub> )                      | < 30sec.   |
| Repeatability                    | (Same day measurement)                  | < +/- 2%   |
| Long Term Stability              | Zero offset                             | < +/-2ppm equivalent / year                              |
|                                  | Sensitivity to H <sub>2</sub> S         | < +/- 10% signal / year                                  |
| Temperature<br>dependence        | (Zero offset swing)                     | < +/-3ppm (-20°C to +50°C)                               |
| Temperature range                | In Service                              | -20°C to +50°C   |
| remperature range                | Recommended in Storage                  | -0°C to +20°C  |
|                                  | In Service                              | 15 – 90%RH<br>(non-condensing)                           |
| Humidity range                   | Recommended in Storage                  | 15 – 90%RH<br>(non-condensing)                           |
| Pressure range in service        |   | 0.9 – 1.1 atm  |
| Detection ronge                  |   | 0 - 500ppm   |
| Detection range                  | Maximum Overload Range                  | 2,000ppm   |
| Recommended load resistor        |   | 10Ω  |
| Recommended maximum storage time |   | 6 months   |
| Warranty                         | Against faulty workmanship or materials | 24 months  |



## **DIMENSIONS:**



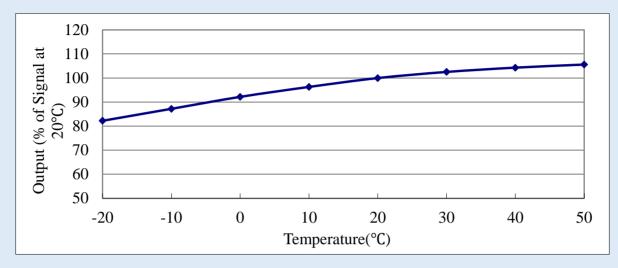
| C | Case Material | PPO           |
|---|---------------|---------------|
| C | Cap Color     | Yellow        |
| ١ | Weight        | 5 g (approx.) |



## PERFORMANCE DATA

#### 1) Uncompensated Temperature Dependence:

The following graph illustrates the typical effect of temperature on the output signal of the NE4-H2S-500, for a sensor calibrated at 20°C:



### 3) Cross Sensitivities:

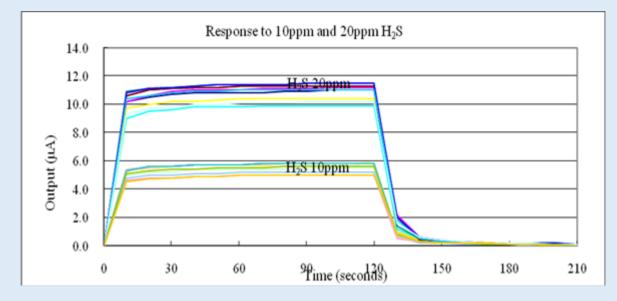
The following table gives cross sensitivity information for a variety of commonly encountered gases (H<sub>2</sub>S = 100)

| Test Gas          | % cross-sensitivity |
|-------------------|---------------------|
| Hydrogen Sulphide | 100                 |
| Carbon monoxide   | < 5                 |
| Hydrogen          | < 2                 |
| Methane           | 0                   |
| Carbon dioxide    | 0                   |
| Sulfur dioxide    | < 23                |
| Nitric oxide      | < 3                 |
| Nitrogen dioxide  | < -30               |
| Ammonia           | 0                   |
| Ethyl acetate     | 0                   |
| Chlorine          | 0                   |
| Ethylene          | 0                   |



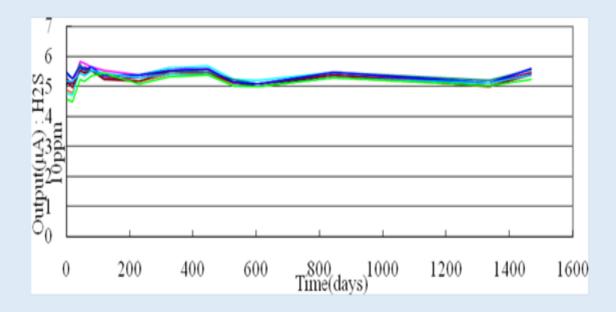
## 4) Response Characteristics

The following are plots of typical responses and recovery times for exposures to Hydrogen Sulphide at 10 ppm and 20 ppm concentrations:



### 5) Long-Term Drift Characteristics:

The following graph illustrates the typical Long term sensitivity stability of the sensor over 1500 days of operation.



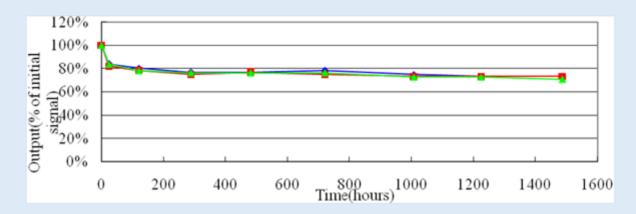
This data was generated in bench tests in a controlled clean environment. In service, in a real application, the long term stability will also be dependent on the environmental conditions of the application.



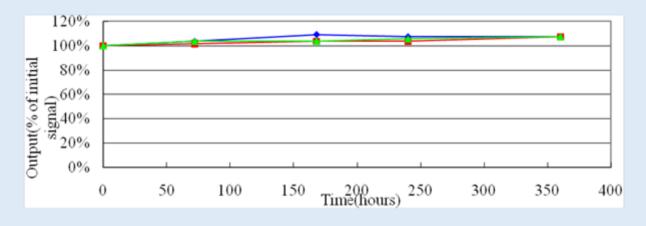
## **TOLERANCE TO ENVIRONMENTAL EXTREMES**

#### 1) Tolerance to temperature and Humidity extremes

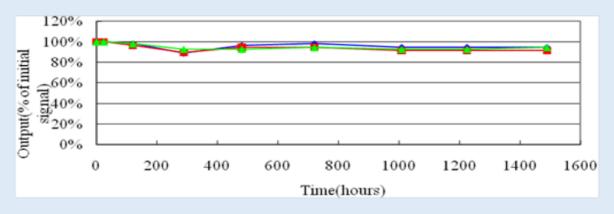
**High Temperature / Low Humidity:** The following graph illustrates the signal output stability for a sensor exposed to  $+80^{\circ}$ C at close to 0%RH over 1500 hours. The H<sub>2</sub>S concentration used was 10ppm H<sub>2</sub>S in air.



**High Temperature / High Humidity:** The following graph illustrates the signal output stability for a sensor exposed to  $+60^{\circ}$ C and 90% RH over 350 hours. The H<sub>2</sub>S concentration used was 10ppm H<sub>2</sub>S in air.



**Low Temperature:** The following graph illustrates the signal output stability for a sensor exposed to -20°C over 1500 hours. The  $H_2S$  concentration used was 10ppm  $H_2S$  in air.





## 2) <u>Tolerance to thermal shock</u>

The following table shows the zero and sensitivity change following an exposure to -20°C for 30 minutes, followed immediately with an exposure to +50°C for 30 minutes. This 60 minute cycle was repeated 10 times. The gas concentration used for testing was 100ppm.

|     | Before tempe               | Before temperature cycling After temperature cycling |                            | Sensitivity                                    |                        |
|-----|----------------------------|--|----------------------------|--|------------------------|
| No. | Zero offset in air<br>(µA) | Sensitivity to<br>100ppm H <sub>2</sub> S (µA)       | Zero offset in air<br>(µA) | Sensitivity to<br>100ppm H <sub>2</sub> S (μA) | variation ratio<br>(%) |
| 1   | 0.01                       | 56.2   | 0.02                       | 56.2   | 100.0                  |
| 2   | 0.01                       | 54.1   | 0.03                       | 54.0   | 99.8                   |
| 3   | 0.01                       | 56.9   | 0.03                       | 56.8   | 99.8                   |
| 4   | 0.01                       | 54.3   | 0.03                       | 54.1   | 99.6                   |
| 5   | 0.01                       | 55.8   | 0.02                       | 55.9   | 100.2                  |

#### 3) <u>Tolerance to mechanical shock</u>

The following table shows the zero and sensitivity change following a drop test: The sensor was dropped from a height of 1m on to a concrete floor 5 times. The gas concentration used for testing was 100ppm.

|     | Before test                | (micro A)                            | After test (micro A)       |                                   | Sensitivity            |
|-----|----------------------------|--------------------------------------|----------------------------|-----------------------------------|------------------------|
| No. | Zero offset in air<br>(µA) | Sensitivity to<br>100ppm H₂S<br>(µA) | Zero offset in air<br>(µA) | Sensitivity to<br>100ppm H₂S (µA) | variation ratio<br>(%) |
| 1   | 0.03                       | 55.4                                 | 0.03                       | 55.5                              | 100.2                  |
| 2   | 0.03                       | 56.5                                 | 0.03                       | 56.4                              | 99.8                   |
| 3   | 0.02                       | 54.7                                 | 0.04                       | 54.9                              | 100.4                  |

### 4) Tolerance to Sulphur Dioxide (SO<sub>2</sub>)

The following table shows the zero and sensitivity change following an exposure to 50ppm Sulphur Dioxide (SO<sub>2</sub>) for 30 minutes. The gas concentration used for testing was 100ppm.

|     | Before exposure            |  | After exposure             |  | Sensitivity            |
|-----|----------------------------|--|----------------------------|--|------------------------|
| No. | Zero offset in air<br>(µA) | Sensitivity to<br>100ppm H <sub>2</sub> S (µA) | Zero offset in air<br>(µA) | Sensitivity to<br>100ppm H <sub>2</sub> S (µA) | variation ratio<br>(%) |
| 1   | 0.00                       | 55.4   | 0.02                       | 55.4   | 100.0                  |
| 2   | 0.00                       | 56.1   | 0.02                       | 56.2   | 100.2                  |
| 3   | 0.00                       | 54.1   | 0.03                       | 54.6   | 100.9                  |
| 4   | 0.00                       | 58.1   | 0.03                       | 58.6   | 100.9                  |
| 5   | 0.00                       | 54.1   | 0.03                       | 54.0   | 99.8                   |



#### 5) Tolerance to Ammonia (NH3)

The following table shows the zero and sensitivity change following an exposure to 200ppm Ammonia for 120 minutes. The gas concentration used for testing was 100ppm.

|     | Before exposure            |  | After exposure             |  | Sensitivity            |
|-----|----------------------------|--|----------------------------|--|------------------------|
| No. | Zero offset in air<br>(µA) | Sensitivity to<br>100ppm H <sub>2</sub> S (µA) | Zero offset in air<br>(µA) | Sensitivity to<br>100ppm H <sub>2</sub> S (µA) | variation ratio<br>(%) |
| 1   | 0.02                       | 56.1   | 0.00                       | 55.4   | 98.8                   |
| 2   | 0.02                       | 57.0   | 0.00                       | 56.1   | 98.4                   |
| 3   | 0.02                       | 55.2   | 0.00                       | 54.1   | 98.0                   |
| 4   | 0.02                       | 59.0   | 0.00                       | 58.1   | 98.5                   |
| 5   | 0.02                       | 55.0   | 0.00                       | 54.1   | 98.4                   |

### 6) <u>Tolerance to Nitrogen Dioxide (NO<sub>2</sub>)</u>

The following table shows the zero and sensitivity change following an exposure to 50 ppm Nitrogen Dioxide (NO<sub>2</sub>) for 120 minutes. The gas concentration used for testing was 100ppm.

|     | Before exposure            |  | After exposure             |  | Sensitivity            |
|-----|----------------------------|--|----------------------------|--|------------------------|
| No. | Zero offset in air<br>(µA) | Sensitivity to<br>100ppm H <sub>2</sub> S (µA) | Zero offset in air<br>(µA) | Sensitivity to<br>100ppm H <sub>2</sub> S (µA) | variation ratio<br>(%) |
| 1   | 0.00                       | 54.8   | -0.01                      | 55.0   | 100.4                  |
| 2   | -0.01                      | 55.8   | 0.01                       | 56.5   | 101.3                  |
| 3   | 0.00                       | 53.5   | 0.00                       | 54.0   | 100.9                  |
| 4   | -0.01                      | 58.2   | 0.00                       | 58.0   | 99.6                   |
| 5   | 0.00                       | 53.7   | 0.00                       | 54.5   | 101.5                  |

### 7) Tolerance to Hydrogen

The following table shows the zero and sensitivity change following an exposure to 500ppm Hydrogen for 10 hours. The gas concentration used for testing was 100ppm.

|     | Before e                   | xposure  | After ex                   | Sensitivity                                    |                        |
|-----|----------------------------|--|----------------------------|--|------------------------|
| No. | Zero offset in air<br>(µA) | Sensitivity to<br>100ppm H <sub>2</sub> S (µA) | Zero offset in air<br>(µA) | Sensitivity to<br>100ppm H <sub>2</sub> S (µA) | variation ratio<br>(%) |
| 1   | 0.03                       | 56.3   | 0.02                       | 56.1   | 99.6                   |
| 2   | 0.03                       | 57.1   | 0.02                       | 57.5   | 100.7                  |
| 3   | 0.02                       | 55.3   | 0.02                       | 55.1   | 99.6                   |
| 4   | 0.03                       | 58.9   | 0.02                       | 59.8   | 101.5                  |
| 5   | 0.03                       | 54.8   | 0.02                       | 55.0   | 100.4                  |



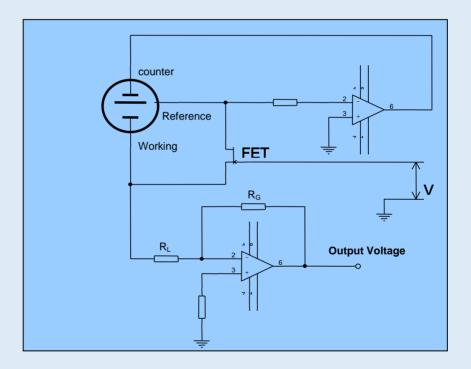
#### 8) Tolerance to Silicone vapours

|     | Before exposure            |  | After ex                   | Sensitivity                                    |                        |
|-----|----------------------------|--|----------------------------|--|------------------------|
| No. | Zero offset in air<br>(µA) | Sensitivity to<br>100ppm H <sub>2</sub> S (µA) | Zero offset in air<br>(µA) | Sensitivity to<br>100ppm H <sub>2</sub> S (µA) | variation ratio<br>(%) |
| 1   | -0.01                      | 55.0   | -0.01                      | 55.0   | 100.0                  |
| 2   | 0.01                       | 56.5   | 0.01                       | 56.1   | 99.2                   |
| 3   | 0.00                       | 54.0   | 0.00                       | 54.2   | 100.4                  |
| 4   | 0.00                       | 58.0   | 0.00                       | 58.6   | 101.0                  |
| 5   | 0.00                       | 53.4   | 0.00                       | 54.2   | 99.5                   |

The following table shows the zero and sensitivity change following an exposure to 200ppm HMDS (HexaMethylDiSiloxane) vapour for 120 minutes. The gas concentration used for testing was 100ppm.

# **NOTES FOR CIRCUIT DESIGN**

The basic measuring circuit for all 3-electrode electrochemical gas sensors is shown below:



• In this arrangement, the output voltage =

Gas Concentration (ppm) x Cell Output (A) x  $R_G(\Omega)$ 

• So if R<sub>g</sub> is 100k, cell output is 200nA/ppm and gas concentration is 10ppm then

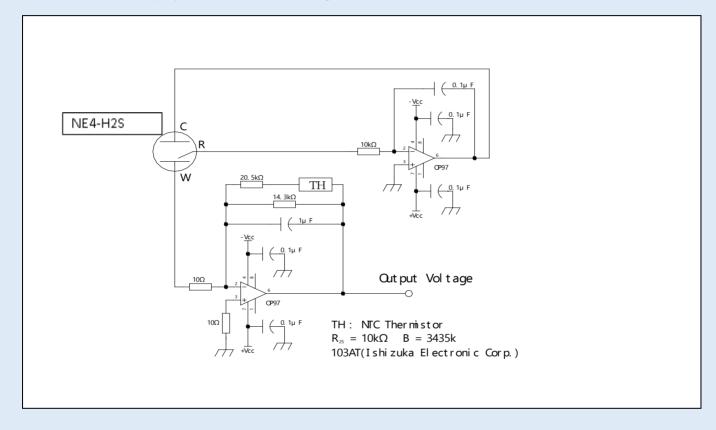
VOUT = 10 x (500x10<sup>-9</sup>) x (100x10<sup>3</sup>) = 0.20V

 R<sub>L</sub> is the cell load resistor (typically 5-50Ω). Speed of response can be increased by reducing the value of R<sub>L</sub>, but signal noise may be increased as a consequence. The recommended values are shown on sensor datasheets



- Amplifiers should be high quality precision low input offset types, e.g. OP90 or similar
- Some sensors require temperature compensation circuitry. A simple compensation network can be incorporated into this circuit by replacing RG with a thermistor. Typically this is a NTC thermistor that has 3435K of B constant adjusting the output accuracy to within ± 10% in the range of -10°C ~ 50°C. Any thermistor with a B-constant around 3500K and resistance value (R<sub>25</sub>) of 10 KΩ can be used. Alternatively, temperature compensation may be undertaken using software lookup tables.

The circuit Nemoto employs for all its internal testing of NE4-H2S-500 is shown below:



Nemoto recognises that the companies who use its gas sensing devices are themselves experts in circuitry design, often with more expertise than Nemoto in this area. The information given here is hence for initial guidance only, and Nemoto does not insist that instrument designers reproduce our circuitry guidance precisely. If the instrument designer deviates from this guidance significantly, however, Nemoto advises that we should be consulted to ensure that the proposed circuit design will function correctly.

Failure to adhere to the recommended circuitry outlined in this document without consultation with Nemoto may result in the suspension of the warranties which apply to the device.



## **GENERAL NOTES ON HANDLING, MANUFACTURE AND INSTRUMENT DESIGN**

#### 1) Long-term drift of gas sensitivity

All electrochemical gas sensors lose sensitivity over time due to small changes on the surface of the working electrode, reducing its oxidation capability. To reduce this, the NE4-H2S-500 uses a newly developed electrode catalyst that will not deteriorate by more than 5% / year. Typically, these changes are limited to less than 5% but we recommend that this deterioration should be taken into account when designing application circuits.

#### 2) Environmental effects on gas sensitivity

Due to the hygroscopic nature of the electrolyte used in electrochemical sensors, moisture is absorbed from or released to the surrounding atmosphere. In high humidity moisture is absorbed, causing an increased sensitivity. In low humidity moisture is released back to the atmosphere and the sensitivity decreases. Conventional electrochemical gas sensors show annual variation of gas sensitivity as much as  $10 \sim 20\%$ .

The NE4-H2S-500 utilises an advanced electrolyte management design and, combined with the unique electrode catalyst structure, these changes can be greatly reduced. Under normal operating conditions, gas sensitivity should change by no more than  $\pm 5\%$  of the output value. The NE4-H2S-500 is designed so that all changes due to moisture uptake/release are completely reversible.

If the gas intake area of the sensor is blocked with water drops or other liquid, gas cannot enter the sensor. The NE4-H2S-500 is fitted with an integral hydrophobic barrier to prevent this, but we recommend the use of additional membrane barriers if the sensor in highly condensing RH conditions.

H<sub>2</sub>S gas is almost the same weight as air. H<sub>2</sub>S detecting devices should be installed around least 5ft from the floor.

If the sensor is to be used in more irregular atmospheres, please contact us for assistance.

#### 3) Storage of sensors

Electrochemical sensors should be stored in a clean air under room temperature, preferably  $0^{\circ}C \sim 20^{\circ}C$  and in non-condensing RH conditions. The maximum storage period would be 6 months after delivery. For sensors stored for more than 6 months, the life in service will be shortened by the excess storage period. Unlike semiconductor type or hot-wire type gas sensors, the gas sensitivity of electrochemical gas sensors will change as time passes regardless of whether sensor has been used or not.

#### 4) Mounting of sensors

Electrode pins must be connected correctly to ensure operation.

If a thermistor is used for temperature compensation, it must be located near the sensor and away from heat sources such as transformers.

The NE4-H2S-500 can be mounted in any orientation.

NE4-H2S-500 connection pins cannot be soldered as excess heating may cause the deformation of the housing and eventually leakage of electrolyte.



#### 5) Calibration and gas testing

Calibration of detectors or densitometers should be done after the output value has been stabilized in clean air.

Evaluation of gas sensitivity should be made in clean, humidified, air. When a test gas is blown directly to the gas intake area, higher gas sensitivity may be observed. It is therefore best to test and calibrate gas detection instruments and sensors in diffusion mode. This can be achieved by using a suitable test housing where a low flow rate is used (<11/min) and where the air is agitated to ensure equal gas diffusion throughout.

Note that Nemoto's own internal testing systems place the sensor in a large chamber, with the gas introduced by injection. The chamber includes a fan which <u>gently</u> agitates the gas inside the chamber to ensure the test gas is fully mixed with the air and does not stratify in the chamber. In this way, Nemoto's own testing very closely simulates the action of the sensor in a typical application.

Other methods, including the use of flow-through hoods and pumped sampling assemblies, may of course be used by instrument manufacturers, but it should be recognised that the method used to expose the sensor to test gas will have a small effect on the accuracy and repeatability of the results obtained, and the correlation of these results with Nemoto's own routine QA test results.

#### 6) <u>Other</u>

Unless otherwise advised by Nemoto, voltage should not be supplied directly to the electrode pins.

Do not bend the pins.

Do not apply more than 5 Kg/cm<sup>2</sup> of force to the sensor.

Take care not to block the gas intake area as it may prevent gas entering the sensor.

Never put foreign material in the gas intake area as it may cause the electrolyte leakage

Do not expose the sensor to excess vibration or shock.

If the sensor housing is damaged, do not use the sensor.

After the sensor is exposed to high concentration gas for long periods, the output signal may require time to recover to normal operation.

Do not blow organic solvents, paints, chemical agents, oils, or high concentration gases directly onto sensors.

Do not disassemble the sensor as this may cause electrolyte leakage.



## **DEFINITIONS**

**Baseline / Baseline shift:** Baseline means the output level in clean air. The output current value at 20°C would be less than 700nA, but this tends to increase as the ambient temperature rises more than 30°C. The baseline shift means this variation of the output level, i.e. a maximum of 1400nA would be put out at 50°C. This baseline shift should be taken into account to optimise overall accuracy. In this manual, the output values are calculated to be equivalent to  $H_2S$  gas concentrations.

**Gas sensitivity / Output signals:** Using the NE4-H2S-500,  $200 \pm 50$ nA is generated at 1ppm of H<sub>2</sub>S gas. For instance, the generated current value will be about 2µA at 100ppm of H<sub>2</sub>S gas (100ppm x 200nA). This generated current is generally recorded as a voltage produced by a Current – Voltage converting circuit as illustrated in our recommended circuit, as the conversion is done through a resistor of 10 K $\Omega$ .

**Response time (T**<sub>90</sub>): This is the time taken to reach to 90% of the maximum output value in clean air.

**Repeatability:** This is the maximum variation of output signals when tests are repeated under the same measuring conditions (temperature, humidity, gas concentration etc.). The repeatability of NE4-H2S-500 is  $\pm$  2%, and this means that all of the test results would fall in the range of 98% ~ 102%.

**Temperature dependence:** All electrochemical sensors are affected by changes in the ambient temperature and the output increases as the ambient temperature rises. This is caused by the rate of oxidation reaction on the surface of the catalyst, the dispersibility of the gas in the capillary, and the thermal effects on the mobility of ions in the electrolyte. This temperature dependency can be compensated relatively easily by using a NTC thermistor.