# Tunable Diode Laser Technology

# **Direct Adsorbtion**

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## **TDL Basics-How does it work**

## What can be measured

## **Types of installations**

## **Products**

## **Combustion Control**

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# **Tunable Diode Lasers**

# How Do They Work

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## **Diode Laser**

- "A laser diode, or LD, is an electrically pumped semiconductor laser in which the active medium is formed by a <u>p-n junction</u> of a <u>semiconductor diode</u> similar to that found in a <u>light-emitting diode</u>.
- The laser diode is the most common type of laser produced. Laser diodes have a very wide range of uses that include, are not limited to, fiber optic communications, barcode readers, laser pointers, CD/DVD/Blu-ray reading and recording, laser printing, scanning and increasingly <u>directional lighting sources</u>.<sup>1</sup>



<sup>1</sup>Text and pictured from (http://en.wikipedia.org/wiki/Laser\_diode)



## TruePeak - General Layout TDLS200



#### Basic Requirements

- Laser
- Detector
- Optics
- Electronics
- Pressure
- Temperature



## Single Peak Spectroscopy



The Tunable Diode lasers have very narrow wavelength emission The linewidth is typically only around 0.00004nm The laser scans the bandwidth, measuring the peak & baseline TRUEPEAK 200/ 220 performs **1000 scans per second** Direct Peak Area integration: accurate regardless of shape of peak



## **Operation – TruePeak TDL**

Coarse wavelength adjustment Laser @ a fixed temperature Cooler@ a fixed temperature

Fine wavelength adjustment A current ramp is fed to the laser



Light is transmitted It passes through gas to be measured





Light is absorbed by the target gas The amount = analyte concentration

The light is then focused on the detector The amount of light at detector = gas concentration

It also provides Diagnostics on the measurement



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## **Operation – TruePeak TDLS..... Page 2**

#### Light absorbed by the target (Peak)

The peak is proportional to the analyte concentration

#### **Enlarged to show concentration detail**







#### Analyzer flips the Peak. Why? Makes it easier to comprehend

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## 2<sup>nd</sup>. Gen Tuneable Diode Lasers

#### There are 2 highly common measurement techniques. They are-Direct Absorption (left spectra) and Second Harmonic or 2f (right spectra)





Changes in background gases affect the shape of the absorption peak

Competitor's Second Harmonic (2f) peak height is AFFECTED

**TruePeak** uses Direct Absorption and Peak Area is UNAFFECTED



## Direct Absorption = Area under Peak Stays the Same





**Process Gases** 

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**O2** in changing background gas concentrations



## Second Harmonic (2f). Peak height is affected

Yes you can try to fit changes into lab determined curves to adjust for <u>changes</u> in: Background gases \* Process Temperature \* Process Pressure Clearly this is not perfect. It is however better than doing nothing It does however lead to false hopes that the compensation is working correctly Who can predict what samples are the proper ones to take?



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## **TruePeak - Length Does Matter**



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## What Can TDL Measure?



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## **TDLS200**

In Situ Analysis without sample conditioning – May be configured as Extractive Fast Response (5-20 seconds) Interference Rejection (high and variable light obstruction) **Process Pressures up to 20 Bar (Application Dependent) Process Temperature up to 1500°C+ (Application Dependent) Optical Measurement, no sensor contact with process** Aggressive Options EX: high particulate content, corrosives etc **Flexible Installation Options** Class 1, Div 2 Group B, C, D when purged **ATEX Category 3 Zone2 Safety Integrity Level:** SIL 1 Assessed Gases measured:  $O_2$ , CO,  $CH_4$ ,  $CO_2$ ,  $H_2S$ ,  $NH_3$ , HCN, HCI, HF, C<sub>2</sub>H<sub>2</sub>, H<sub>2</sub>O, CO%+CO<sub>2</sub>%,

 $NH_3ppm + H_2O\%, CO+CH_4$ 





## TDLS220

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Extractive Analysis sample conditioning may be needed Fast Response (5-20 seconds) **Interference Rejection (high and variable light obstruction)** Sample gas temperature up to 120°C (248°F) (When ambient temperature is  $\leq 40^{\circ}$ C) Sample Pressures up to 7 Bar (100psi) **Optical Measurement, no sensor contact with process Aggressive Options EX: high particulate content, corrosives etc Flexible Installation Options** Class 1, Div 2 Group B, C, D when purged Gases measured: O<sub>2</sub>

Max Range 0-100%, Min Range 0-1% More gas measurements to come





## **TDLS500**

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Extractive Analysis sample conditioning may be needed Fast Response (7-20 seconds) Interference Rejection (high and variable light obstruction) Sample gas temperature up to 100°C (212°F) Ambient temperature upper limit is 40°C – Designed for controlled environment Sample Pressures up to 7 Bar (100psi) **Optical Measurement, no sensor contact with process** Multi-pass Integrated Cavity Output Spectroscopy (ICOS) **Optical Path Lengths up to 10,000 meters for high sensitivity** Sub-ppb detection limits **Flexible Installation Options** Class 1, Div 2 Group B, C, D when purged Gases measured: Inlet Outlet (to pump) Fiber-coupled Lens diode laser  $C_2H_2$ ,  $NH_3$ ,  $CO_1H_2S$ Detector Gas sample  $C_{2}H_{2} + m - C_{2}H_{2}$  $C_{2}H_{2} + NH_{3}$ Data collection laser control Display/readout

electronics

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DKDCAMA

and analysis system

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# **Installation Options**









#### • CROSS STACK/PIPE (IN-SITU)

- Measurement across the process
- Path integrated measurement
- Validation options
  - » Off line verification/calibration with calibration cell
  - » On line verification with dynamic spiking (bump cell)

#### **Installation Options**



#### **BYPASS LEG**

- Process flow through measurement leg
   or-
- Process slipstream through measurement leg
- Allows isolation from process
- Validation options
- Large diameter pipe run
  - Isolate from process, flow gas standard
  - On line verification with dynamic spiking

#### **FLOW CELL**

- Pull or push sample through flow cell
- Validation options
  - Isolate from process, flow gas standard



#### **INSITU ALINGMENT**

- The analyzer laser beam must be able to pass from one side of the process to the other.
- The flanges and nozzles must be within +/- 2 Degrees of center line













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#### UTILITY PANEL

#### - A **Utility Panel** provides a central location for:

- Nitrogen supply for purges
- Validation gas supply
- Purge control
- Validation control
- 110 VAC line power in and 24VDC out to each analyzer
- Analog signals
- Digital signals
- Analyzer interface
- Yokogawa supplies a single interconnect cable that connects the Utility Panel to the Launch unit for power and signal requirements
- Utility Panels for 1 to 4 analyzers are available



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#### **SINGLE UTILITY PANEL**



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# Combustion Control Applications



#### Efficiency

Minimize excess  $O_2/CO$ 

#### Throughput

Minimize heat capacity

#### **Emissions**

Reduce NOx, CO, CO<sub>2</sub>

#### Safety

- Avoid fuel rich conditions

- Identify burner flame out
- Identify process tube leaks





### Placement

- Oxygen concentrations can have high distribution in large systems (vertical and horizontal)
- Vertical distribution is due to tramp air (air leaks)
- Horizontal distribution is due to burner variations and flow effects
- Placement is critical to allow control, distributions can be 50% to >100% of the average excess oxygen from the burners
- Errors for low temperature in-situ probes placed further away from the burners are dominated by tramp air effects
- Errors for high temperature CCE analyzers are dominated by burner effects. Multiple analyzers are typically installed. Decisions on which values to use are significant (low, average)





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### Speed

 Sensors have fast response (5-10s typical). Filters and diffusion elements can significantly affect response time (can be tested).

### Interferences

 Any combustible gases in the process (CO, HC's, H<sub>2</sub>, etc) will burn with oxygen at the sensor, consuming oxygen and forcing the measurement low.

Example:  $(5\% O_2 \text{ in the presence of } 1\% C_3H_8)$ 

 $C_3H_8 + 5O_2 = 3CO_2 + 4H_20$ 

### 5% O<sub>2</sub> level would read "zero"

## **The Past**

### Solid State Sensors (combined with ZrO<sub>2</sub>)

- Thick/thin film
- Catalytic bead

## Optical

- NDIR, Gas Filter Correlation







### **Combustion systems are changing:**

- Emissions limits are lower. Low NOx burners have reduced CO emissions. Measurement is more difficult.
- Furnaces are larger with more burners. Catching breakthrough from a "bad" burner requires improved sensitivity
- NOx emissions can limit system operations.
- Efficient combustion is critical in allowing maximum firing rates

## Increasing cost of fuel and feedstock puts a higher emphasis on combustion control.

**Efficiency = Lower Fuel Costs + Higher System Throughput** 



# **CO breakthrough** determines the ideal control point, prior to breakthrough:

- Highest efficiency
- Highest temperatures produced
- Combustibles are consumed

**CO trim control** can delivery optimum efficiency and flame temperature while remaining safe

**CO levels** from burners have been a moving target

- Older burners CO levels 100's of ppm
- Low NOx burners CO levels <50ppm</li>
- Ultra low-NOx burners CO levels <10ppm</li>



#### **Efficiency Measurements**

Oxygen	<ul> <li>Primary combustion efficiency measurement. Easy to use for control</li> <li>Typically also used as safety measurement</li> </ul>
СО	<ul> <li>Ideal set point measurement (for excess air)</li> <li>Pre-cursor to combustibles breakthrough</li> </ul>



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Proprietary info goes here ...



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#### What is the Correct Excess Oxygen Value ?



**CO** Measurement can determine O<sub>2</sub> setpoint

# TDL is seeing increased use in combustion oxygen measurements

- Path average measurement reduces distribution errors
- No interference from combustibles or CO
- No potential ignition source during upset conditions
- Fast response (5 seconds)
- Ability to provide Measurement in Gas Temps up to 1500 C



### **Measurement Suite using TDL**



\*Consult Yokogawa for Temperature, CH4, and  $\rm H_2O$  . These are application dependant

**Operator Test. Adjust O2 downward to cause CO breakthroughs.** 



- NOx limits can result in firing rate (capacity) limits
- NOx credits can be sold
- NOx is formed in the combustion process through reaction of nitrogen and oxygen in burner air feed
- Reducing excess air, reduces nitrogen and oxygen, resulting in reduced NOx emissions
- CO<sub>2</sub> emissions are also reduced through efficiency improvements
- CO emissions can be measured and controlled near real time

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#### Three Conditions present safety concerns ...

#### Fuel rich burner conditions

CO levels increase as a precursor to hydrocarbon breakthrough

#### - Burner flame out

- Temperature drops rapidly
- Oxygen increases rapidly

#### Process tube leaks

- Moisture may increase (steam cracking)
- Oxygen may not change significantly
- CO may not change significantly

- Hydrocarbons (methane) increase rapidly
- Moisture drops rapidly
- Hydrocarbons increase (methane)
- Temperature may not change significantly
- Measuring and understanding furnace conditions (indicated in red) can help identify safety concerns and their causes.
- This can only be accomplished by having enough measurements points to discriminate between differing safety concerns.
- $\rightarrow$  O<sub>2</sub> and CO measurements are not sufficient.

#### Solution: Measurement Suite using TDL (Patent pending)



### Large Scale Combustion (Furnaces and Heaters)

- **Path vs. Point** decision (O<sub>2</sub> and CO)
- Measurement location (**distribution**, tramp air)
- Response time needs (safety + control)
- **Control method** (single air control, multiple fuel controls)
- **Safety issues** (Ignition sources, combustibles measurement)



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- Emissions limits are lower.
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- Furnaces are larger with more burners. Catching breakthrough from a "bad" burner requires improved sensitivity
- NOx emissions can limit system operations. Efficient combustion is critical in allowing maximum firing rates

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