# LD16-08



# Measurement of hydrocarbons in UHP Oxygen using the MultiDetek 2 and PlasmaDetek-E



This application note shows different methods that have been developed by LDetek for measuring the hydrocarbons in a stream of Oxygen (other matrixes can be analysed as well since the PlasmaDetek-E is strictly selective to hydrocarbons). This application note is the continuity of the application note LD16-02. It is then suggested to first read the LD16-02 to be advised about the technology that was used.

# **LDETEK SOLUTION:**

The chromatograms shown in figure 1 and in figure 3 have been performed using a single column with a Plasmadetek-E. The system uses nitrogen as a unique carrier gas. The simplicity of this configuration makes this solution very efficient and simple for operation. No maintenance is required for such micro plasma based system.

## Method for chromatogram in figure 1:

The MultiDetek2 configuration uses a 10 port diaphragm valve mounted in an injection/backflush mode. The sample gas is then introduced in the packed porous polymer type column for eluting the Air/O2 followed by CH4 in one direction to the detector before to reverse the flow in the column to get the NMHC redirected to the detector. The selective mode of the PlasmaDetek-E makes the detector being strictly selective to hydrocarbons and makes oxygen/air being invisible to the detector. The detector response and LDLs are demonstrated in figure 2. Please note that different performances can be achieved depending on the maximum and minimum ranges required.

### Method for chromatogram in figure 3:

Using the same valve configuration combined with a different column type and system conditions, the analysis of separated C2's becomes possible. For some applications, where it is critical to measure C2s individually, especially acetylene at low concentrations, this method is preferred. The addition of C3+ measurement can be done in the same system to cover the whole hydrocarbon analysis solution. It offers better accuracy, sensitivity and simplicity than conventional FID or Argon plasma method. Figures 4-5 and 6 show an example of performances using this method in real life conditions.

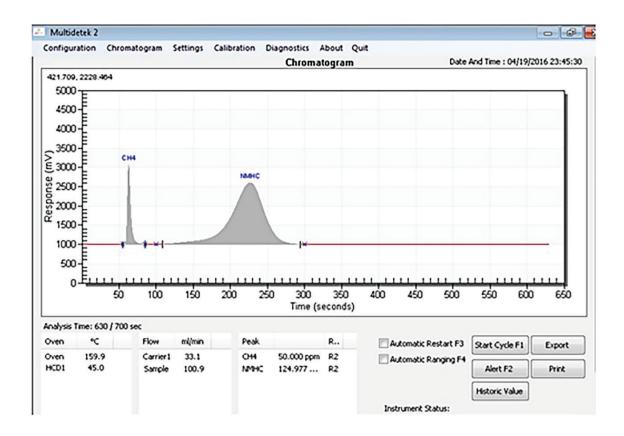


Figure 1: Chromatogram of CH4 & NMHC in oxygen matrix

COMPONENT	CONCENTRATION	PEAK HEIGHT	NOISE	LDL (3X NOISE)
CH <sub>4</sub>	50.000 ppm	2211 mV	0.8 mV	0.054 ppm
NMHC	124.977 ppm	1622 mV	0.4 mV	0.092 ppm

Note: other LDL could be obtained with different injection volume and chromatographic condition

Figure 2

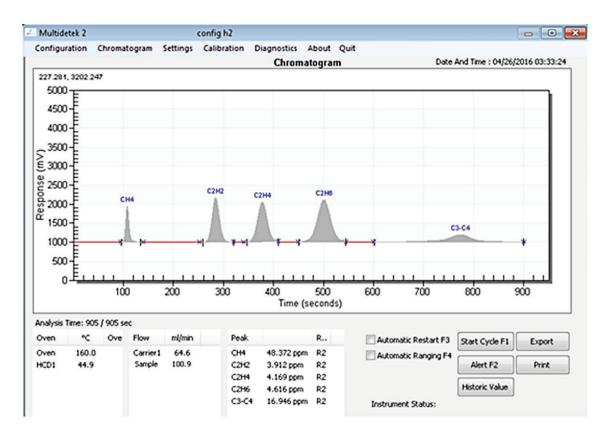


Figure 3: Chromatogram of trace CH4,C2H2, C2H4, C2H6 & C3+ in oxygen matrix

COMPONENT CONCENTRATION PEAK HEIGHT NOISE LDL (3X NOISE)	
CH <sub>4</sub> 48.372 ppm 951 mV 0.3 mV 0.046 ppm	
$\mathrm{C_2H_2}$ 3.912 ppm 1241 mV 3.3 mV 0.031 ppm	
${\rm C_2H_4}$ 4.169 ppm 1111 mV 3.3 mV 0.037 ppm	
${\rm C_2H_6}$ 4.616 ppm 1146 mV 3.3 mV 0.040 ppm	
C <sub>3</sub> + 16.946 ppm 220 mV 0.9 mV 0.207 ppm	

 $\label{thm:local_problem} \textbf{Note: other LDL could be obtained with different injection volume and chromatographic condition}$ 

Figure 4

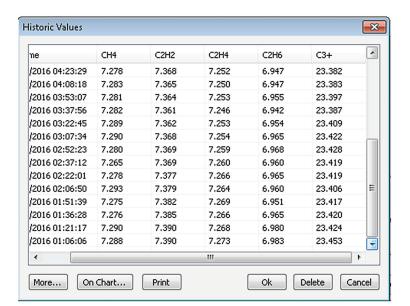


Figure 5: Example of stability results over 10 consecutive cycles

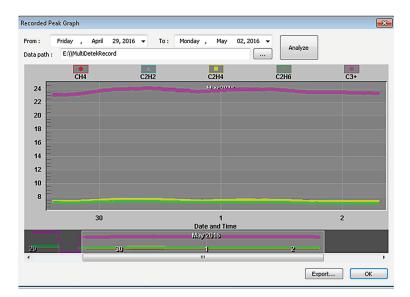


Figure 6: Example of stability results over a 3 day period

