



ELEMENTAL MAPPING OF SHALE ROCKS USING LASER INDUCED BREAKDOWN SPECTROSCOPY (LIBS)

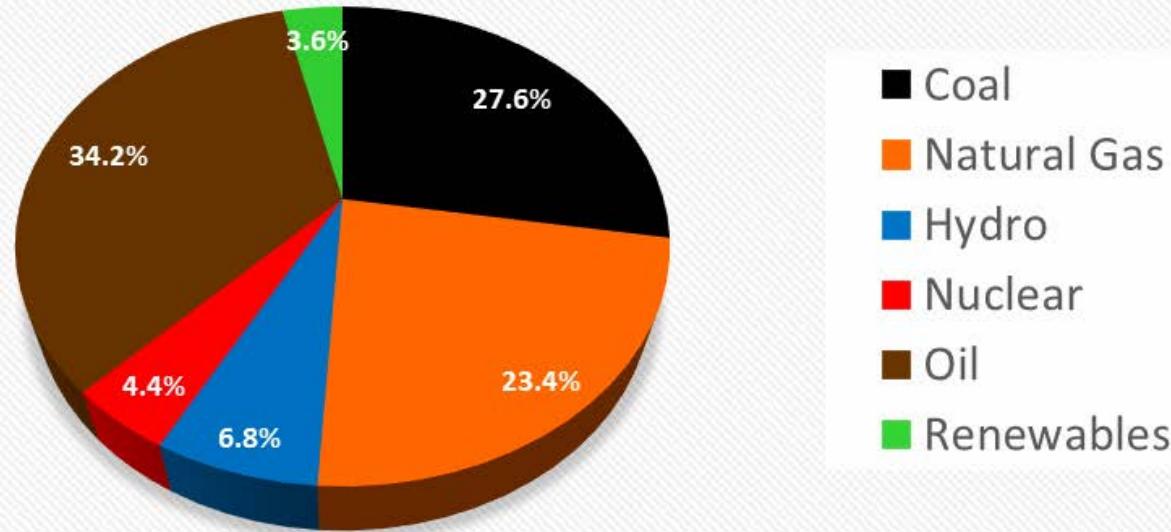
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Energy Consumption

World total primary energy consumption by fuel in 2017



Source: BP Statistical Review of World Energy, 67th Edition, June 2018

Shale Oil and Gas Resources



Resources	Oil (billion barrels)	Gas (trillion cubic feet)
Shale	345	7299
Non-shale	3012	15883
Total	3357	22882
% Shale	10	32

Source: U.S. Energy Information Administration, Independent Statistics and Analysis, June 2013

Projected Shale gas production

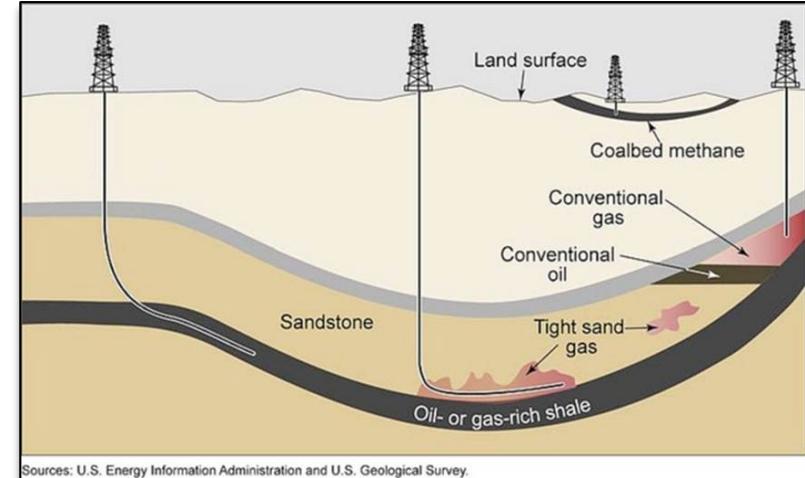


Region	2015 (bcf/day)	2040 (bcf/day)
World	42	168
USA	37	79
Other countries: Canada, China, Mexico, Argentina, Algeria		

Source: U.S. Energy Information Administration, International Energy Outlook 2016 and Annual Energy Outlook 2016

Shale Characterization

- Shale formations have oil and gas trapped within the pore spaces and are considered largest natural gas deposits
- Elemental composition can provide clues to rock properties (porosity, permeability, minerals) that could effect oil and gas accumulation
- Higher amount of carbon and hydrogen (organic material) means high gas potential
- Knowledge of mineralogy helps in selection of drilling location, resolving drilling problems, and making engineering and production decisions.
- Environmental issues associated with shale retorting require substantial monitoring and control of waste product



Sources: U.S. Energy Information Administration and U.S. Geological Survey.

Unconventional Shale



- Fine grained sedimentary rocks
 - < 2 microns
- High clay content
 - Generally > 50%
- Low permeability
 - Nano-Darcy
- High organic content
- Hydrocarbon generation
- Susceptible to hydraulically induced fractures
 - E.g. generally behaves in a brittle manner



Shale Core Logging



- XRF is a widely used technique for shale logging/mapping
- SEM-EDS for spectral mapping of shale
- Low atomic mass elements (i.e., C, H) are not measurable by XRF
- More than one technique is needed to obtain complete elemental information
- LIBS is capable of measuring these elements simultaneously.

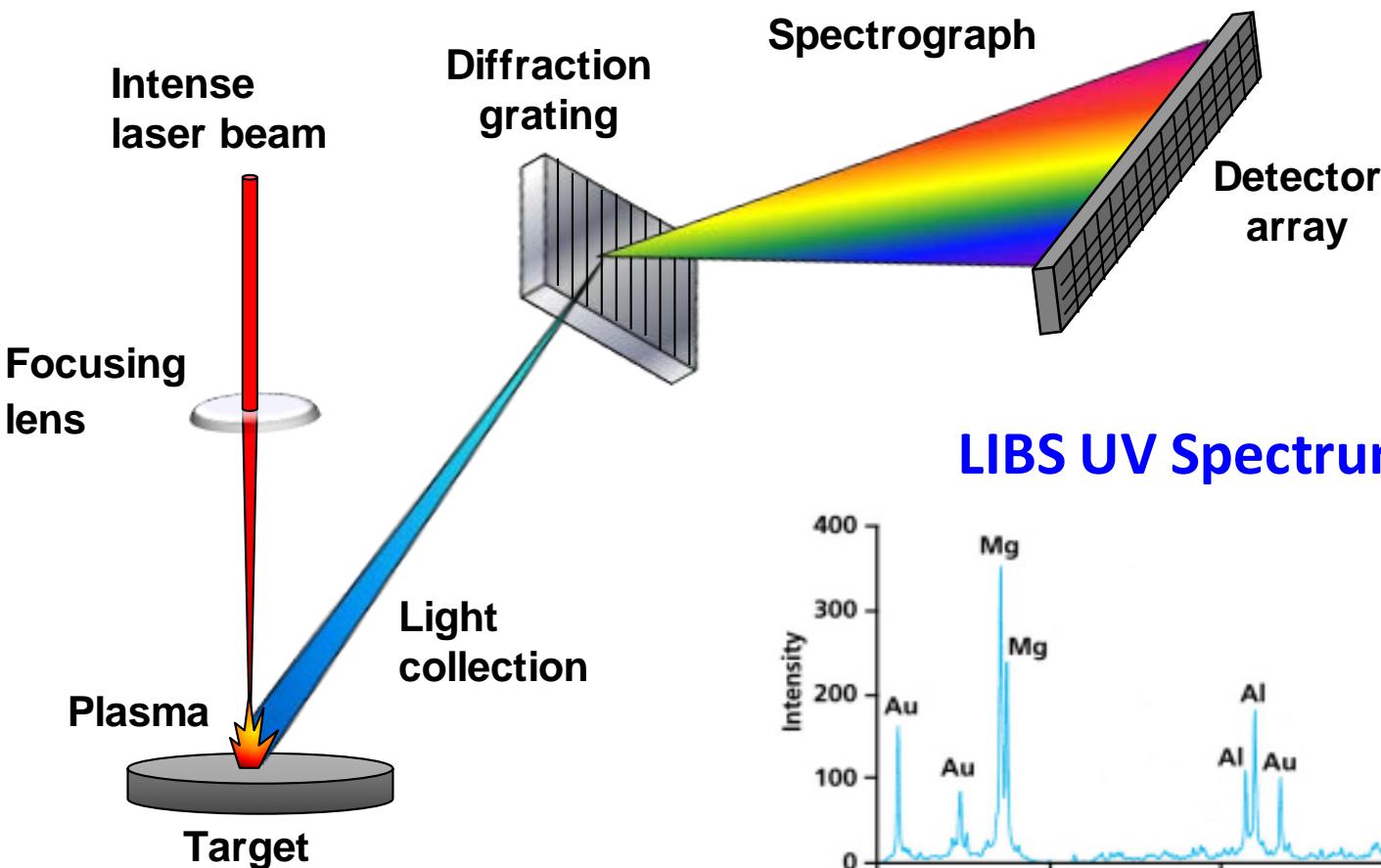


Laser Induced Breakdown Spectroscopy

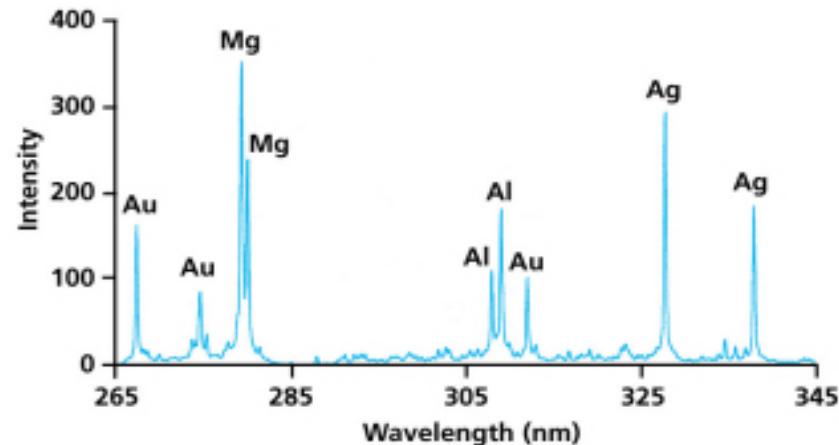


- Laser is fired upon a given sample and laser energy focused to a small spot
- A hot luminous plasma vaporizes the material, and leads to atomization and excitation of elements
- As plasma cools, emission occurs and the emitted light can be collected
- Every element in the Periodic Table gives off light at a distinct wavelength
- LIBS is capable of analyzing solid, liquid, and gaseous samples with minimum or no sample preparation
 - Matrix and/or major elements
 - Non-metals such as C, H, N, O and halogens (F)
- LIBS can perform both surface and depth analysis

Principle of LIBS



LIBS UV Spectrum



LIBS Instrument & Method Parameters



- **Experimental Conditions**

- 266 nm Nd:YAG nanosecond laser
- 50 μm laser spot size
- 81 x 81 grid pattern map
- 8 X 8 mm (64 mm²)
- 5 shots per location – Accumulated
- 6561 data points per map
- ~ 390 minutes per map
- ASI's Axiom operation software



J 200 Applied Spectra LIBS system

Shale Samples Analyzed



Sample 7498



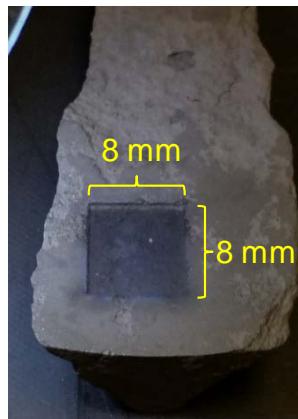
Sample 7531



Sample 9655



Sample 7504

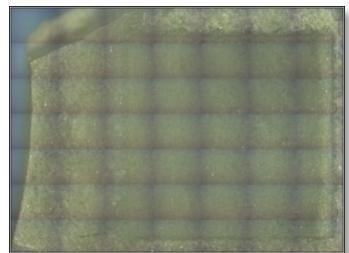
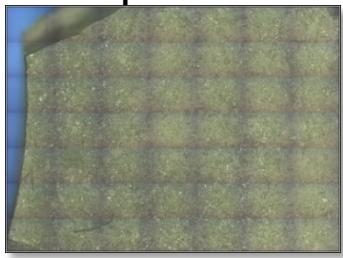


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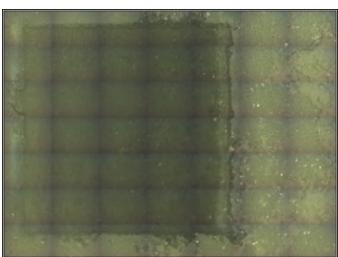
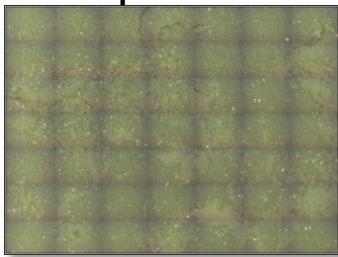


Shale Samples Analyzed

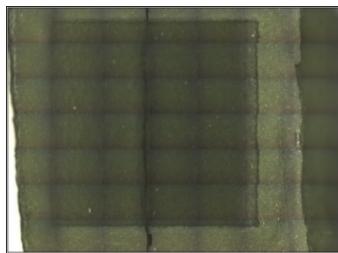
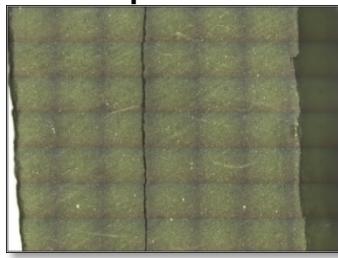
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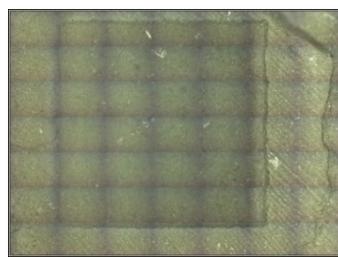
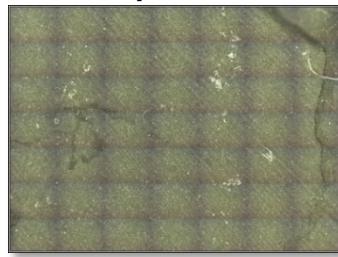
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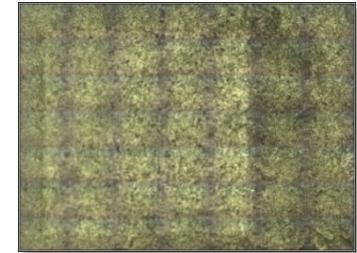
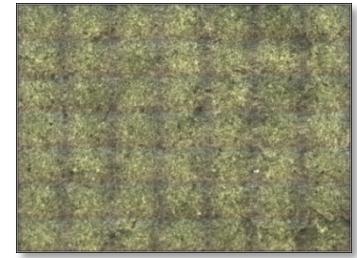
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Sample 7551



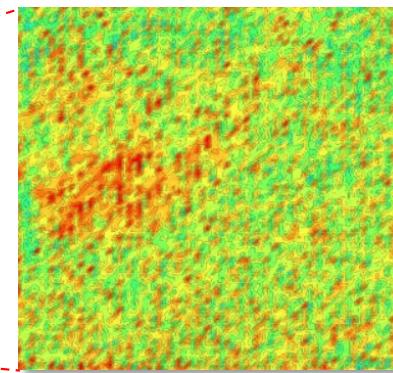
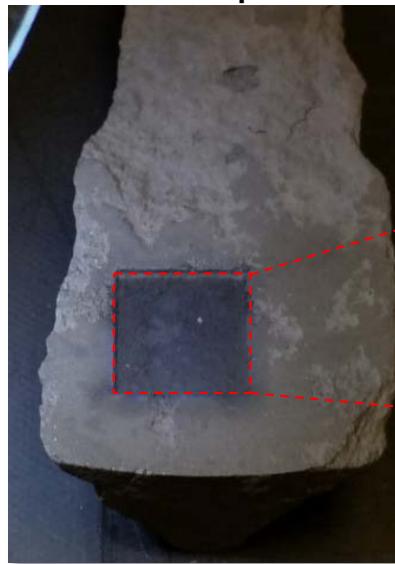
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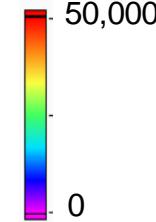
Shale Sample Mapped Area



Sample 7504



Aluminum (a.u.)

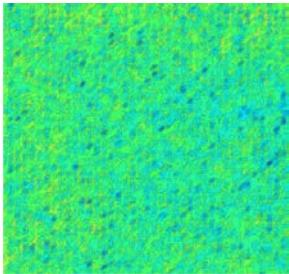


C193, H656, O777, Ca393, Mg280, Al394, Fe275, Si288, Ba493, K766, Sr407, Ti334, Na589

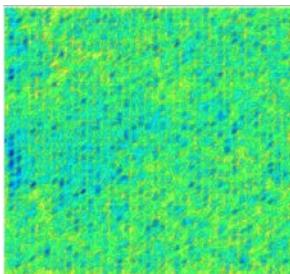
Shale Elemental Maps



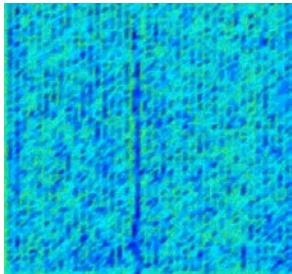
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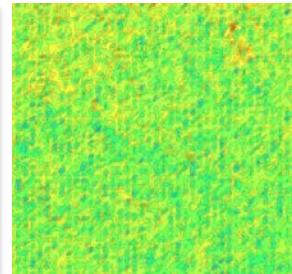
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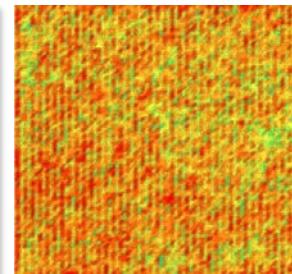
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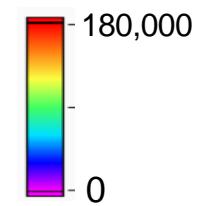
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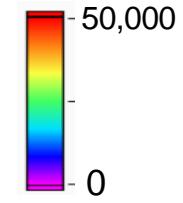
Sample 9655



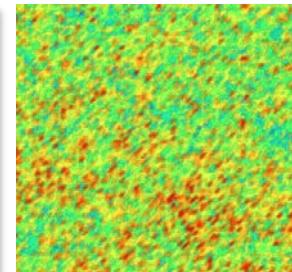
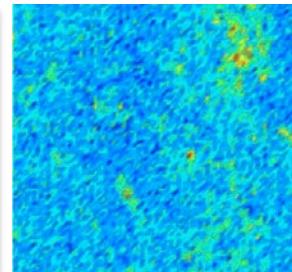
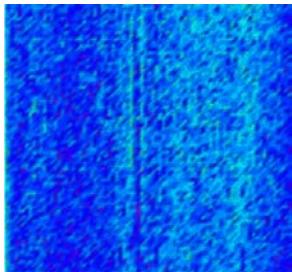
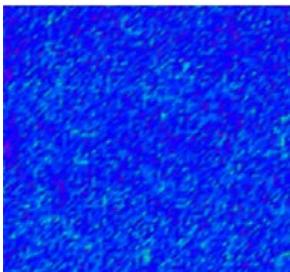
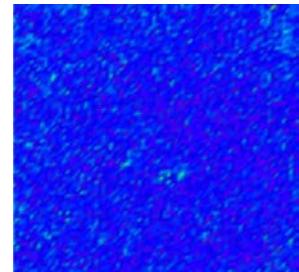
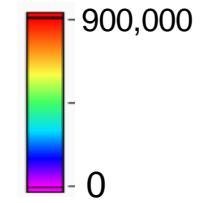
Oxygen (a.u.)



Aluminum (a.u.)



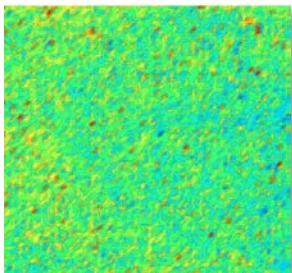
Calcium (a.u.)



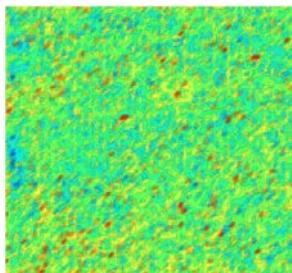
Shale Elemental Maps



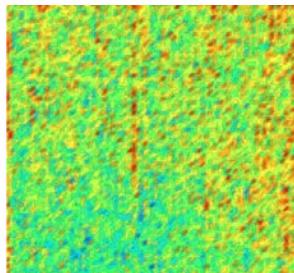
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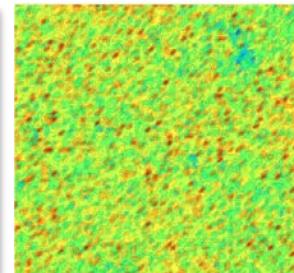
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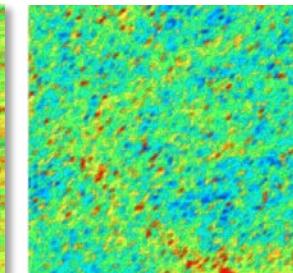
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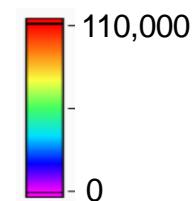
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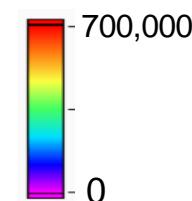
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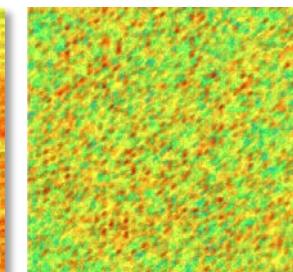
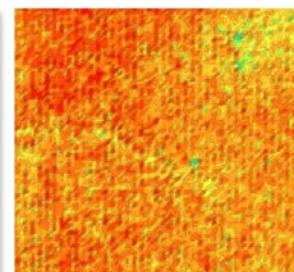
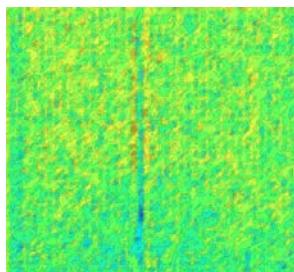
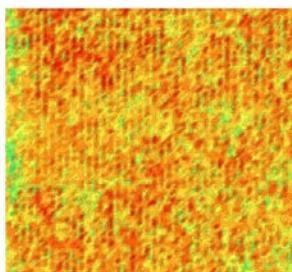
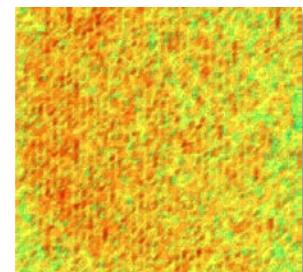
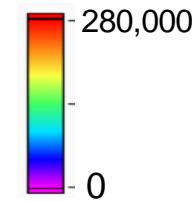
Iron (a.u.)



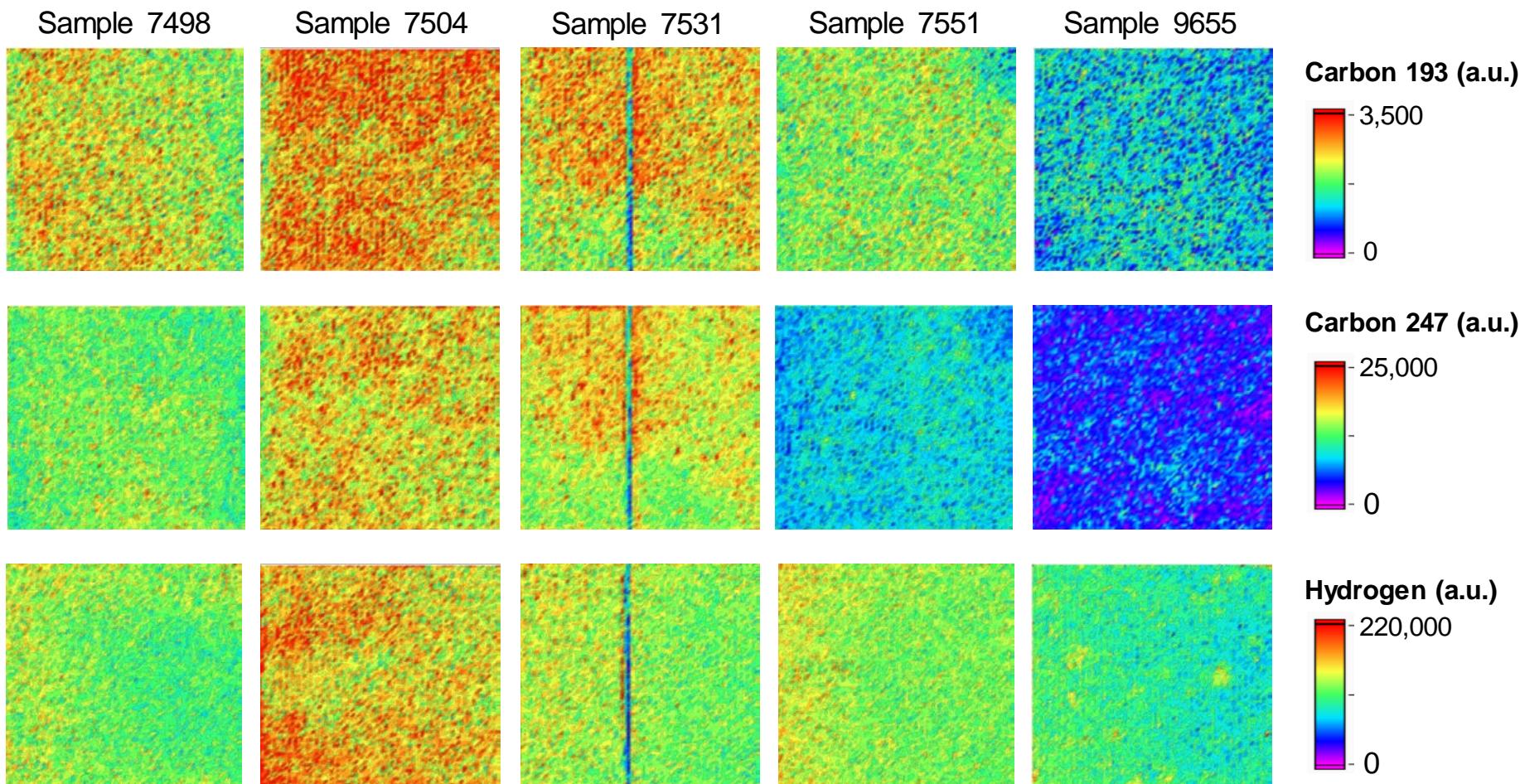
Magnesium (a.u.)



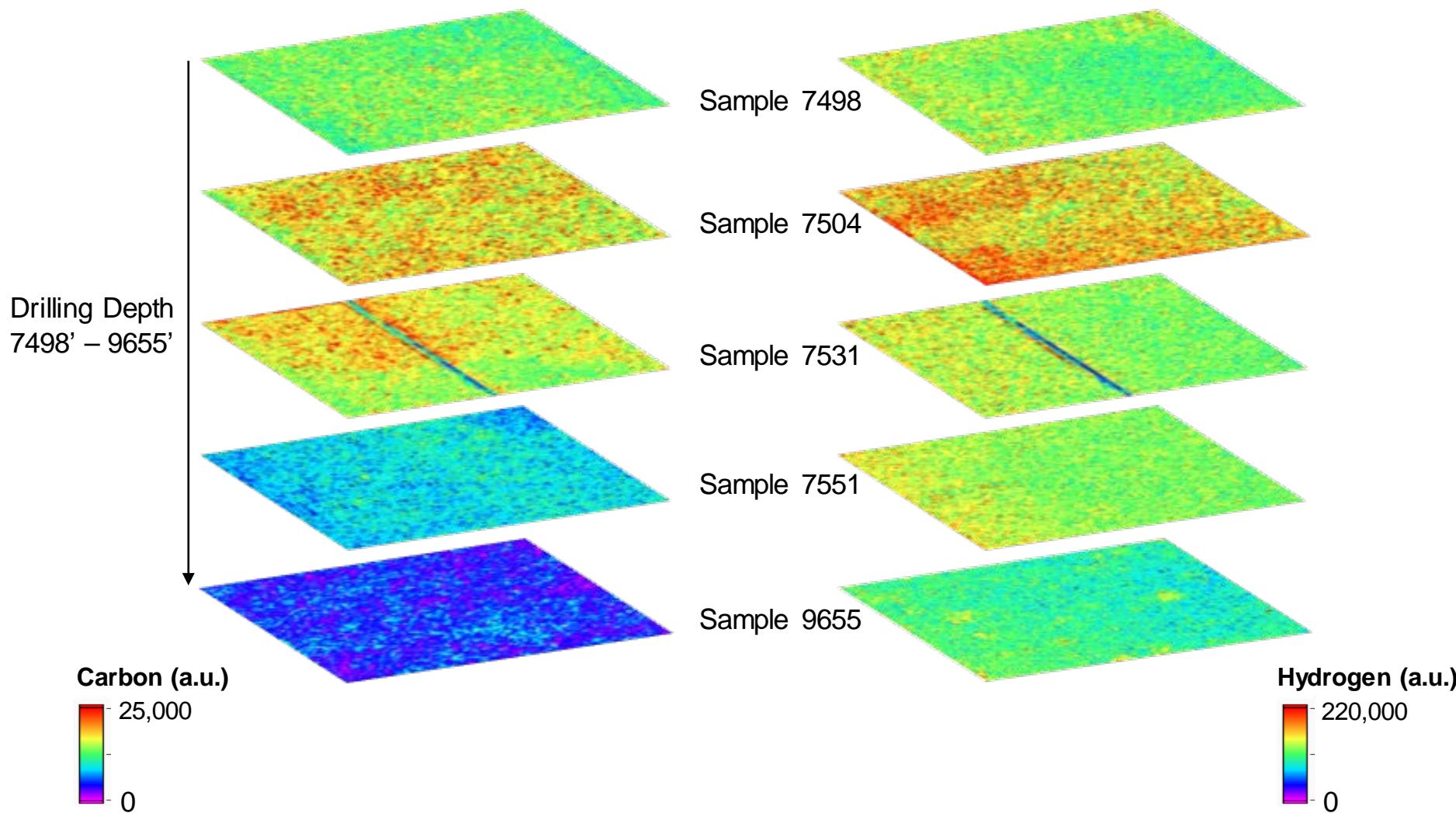
Silicon (a.u.)



Shale Elemental Maps



Shale C & H Maps Depth Profile

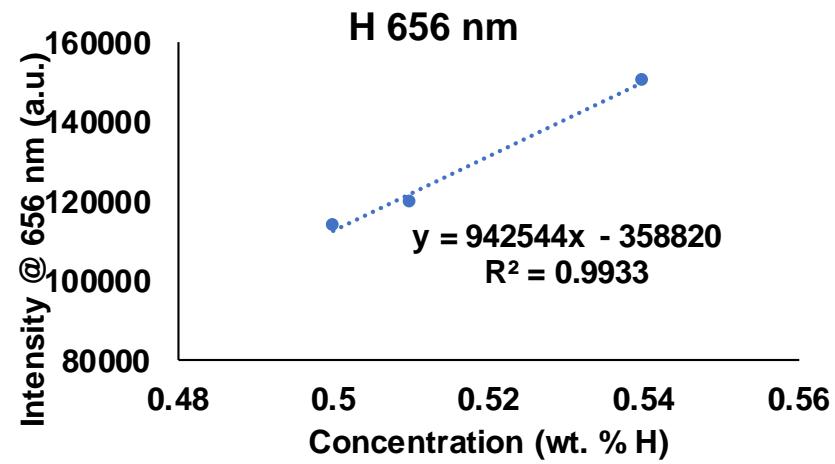
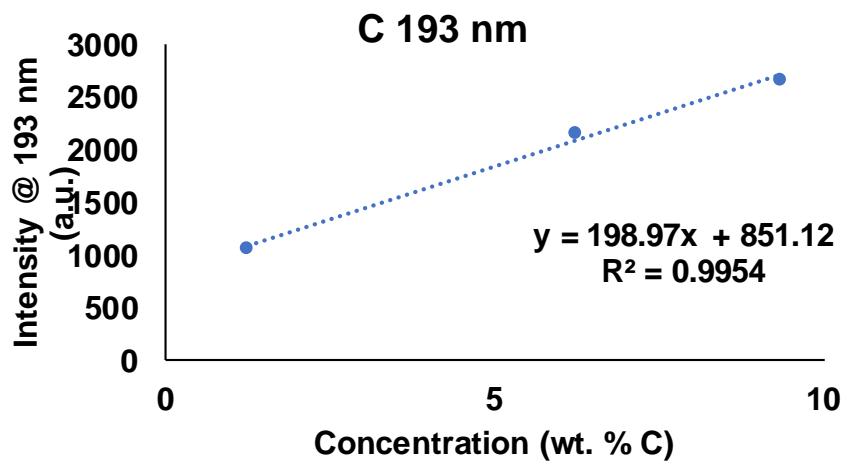


Calibration Standards

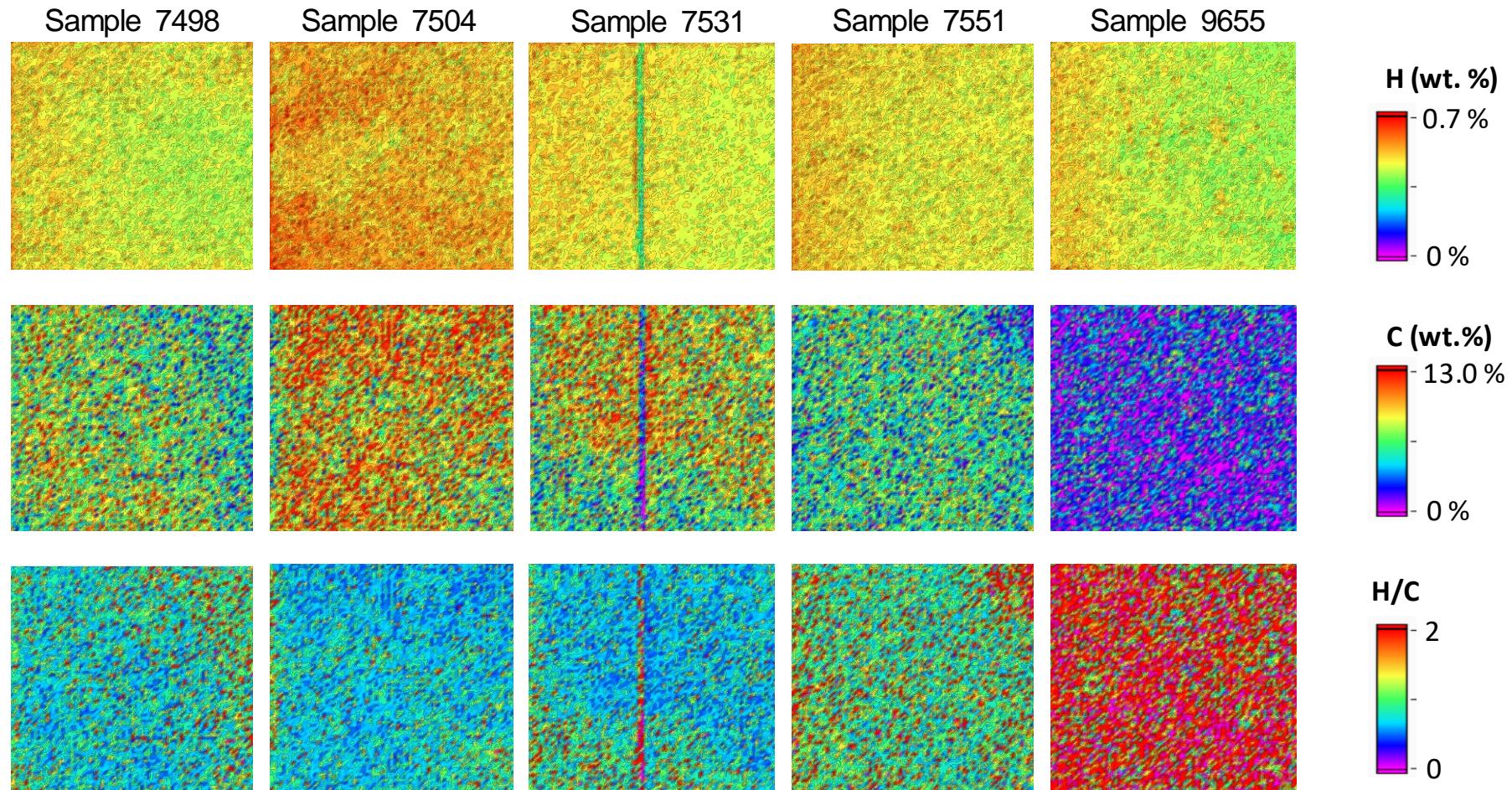


- Availability of matrix matched calibration standards is limited
- Shale samples characterized in our previous study (*Spectrochimica Acta B*, 122, 2016, 9-14) by ICP-OES and LIBS were used to calibrate the instrument
- C and H analysis was performed at the Western Kentucky University using Leco CHN TrueSpec analyzer
- Calibration standards were mapped along with samples to create calibration curves

Calibration Curves



Concentration Maps



C & H Concentrations



Total concentration of 8 mm x 8 mm area analyzed (n = 6,561)

	C 193	%RSD	H 656	%RSD
7498'	7.13 %	8.4	0.46 %	8.1
7504'	9.26 %	5.1	0.53 %	9.2
7531'	7.93 %	8.1	0.47 %	9.4
7551'	5.89 %	21.0	0.48 %	7.3
9655'	2.39 %	54.0	0.45 %	9.3

	H/C
7498'	0.85
7504'	0.76
7531'	0.84
7551'	1.05
9655'	1.70

H/C < 1 – aromatic

H/C > 1 – aliphatic

C H N Analyzer		LIBS	
	Reference Value	LIBS	% BIAS
7504'	9.33 wt. % C	9.26 % C	-0.8
7504'	0.51 wt. % H	0.53 % H	3.9

Sample	TC	TOC	TOC/TC
M7498	7.13	4.78	0.67
M7504	9.26	5.92	0.64

Elemental Concentrations



Concentration Average Value (Wt.% \pm 1 SD)						
	Depth (')	Al 394	Ca 393	Fe 275	Mg 280	Si 288
PARA2 7498	7498	0.71 \pm 0.52	1.22 \pm 0.20	1.74 \pm 0.11	0.65 \pm 0.4	21.7 \pm 0.99
PARA1 7504	7504	1.63 \pm 0.75	1.53 \pm 0.19	1.82 \pm 0.11	0.68 \pm 0.03	22.6 \pm 1.06
MSEEL 7531	7531	0.8 \pm 0.29	2.12 \pm 0.50	1.99 \pm 0.19	0.58 \pm 0.04	17.26 \pm 1.05
MSEEL 7551	7551	1.67 \pm 0.62	3.32 \pm 0.48	2.06 \pm 0.09	0.70 \pm 0.03	23.60 \pm 1.05
Bakken 9655	9655	0.73 \pm 0.59	7.67 \pm 0.66	1.72 \pm 0.14	1.06 \pm 0.07	20.09 \pm 0.90

Comparison of XRF and LIBS values (Wt.% \pm 1 SD)

Sample		Al	Ca	Fe	Si
M7498	XRF	5.18 \pm 0.13	1.60 \pm 0.01	3.85 \pm 0.03	21.27 \pm 0.12
	LIBS	0.71 \pm 0.52	1.22 \pm 0.20	1.74 \pm 0.11	21.7 \pm 0.99
M7504	XRF	3.39 \pm 0.97	1.48 \pm 0.01	2.75 \pm 0.02	22.1 \pm 0.11
	LIBS	1.63 \pm 0.75	1.53 \pm 0.19	1.82 \pm 0.11	22.6 \pm 1.06
M7531	XRF	2.16 \pm 0.11	0.84 \pm 0.01	3.21 \pm 0.03	15.44 \pm 0.11
	LIBS	0.8 \pm 0.29	2.12 \pm 0.50	1.99 \pm 0.19	17.3 \pm 1.05
M7551	XRF	2.13 \pm 0.09	3.21 \pm 0.02	3.55 \pm 0.03	21.72 \pm 0.12
	LIBS	1.67 \pm 0.62	3.32 \pm 0.48	2.06 \pm 0.90	23.6 \pm 1.05

Elements Calibration Curves

Al	$y = 3.698 x + 26528$
Ca	$y = 61.8 x + 86709$
Fe	$y = 41.28 x - 16559$
Mg	$y = 440.4 x - 86648$
Na	$y = 514.6 x + 49061$
Si	$y = 11.778 x - 52464$
Sr	$y = 579.7 x - 1871$
Ti	$y = 231.3 x - 46293$

Conclusions



- Laser induced Breakdown Spectroscopy (LIBS) can provide mineral composition and distribution in shale
- The technique provides accurate data with reasonable detection limits for most of the elements
- It can analyze light elements including C and H.
- LIBS can determine H/C ratio to predict the presence and type of gaseous hydrocarbons and impurities in scanned area
- Minimum to no sample preparation makes this technique an attractive option to avoid lengthy sample preparation procedures
- LIBS can be a robust device for in-situ shale core mapping and exploration purposes

Acknowledgment

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