

Petroleum Engineering Department

Faculty Research & Lab Facilities



Spring 2017



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- O PETROLEUM ENGINEERING



PE Faculty and Research Areas Hazim Abass – Hydraulic fracturing research Manika Prasad – Petrophysics of Organics, Clay, Sand, and Shale • Rosmer Brito – Midstream, Production System • Jorge Sampaio – Drilling and engineering **Design and Optimization** modeling and simulation Alfred Eustes – Drilling for petroleum & non- Azra Tutuncu – Geomechanics and petroleum unconventional gas and oil institute • Will Fleckenstein – Drilling and hydraulic Philip Winterfeld – Numerical simulation, flow fracturing and transport phenomena in porous media Ramona Graves – Reservoir Characterization Yu-Shu Wu – CO₂-EOR, CO₂ sequestration, and CEMMC geothermal, hydrology • Hossein Kazemi – IOR/EOR, reservoir studies at • Xiaolong Yin - Pore-scale physics and flow, MCERS suspension, phase behavior Jennifer Miskimins – Stimulation and FAST • Luis Zerpa – EOR, reservoir, flow assurance, gas consortium hydrate in nature Erdal Ozkan – Well testing / MCERS / Unconventional reservoir engineering PETROLEUM ENGINEERING COLORADO SCHOOL OF MINES































































Well Categories / Piceance Basin Failure Probability

Category	Barriers	Independent Failure Events	Description	Risk Level
1	1	3	Shallow Surface Casing Top of Production Casing Cement Below Over-Pressured Hydrocarbon Reservoir	High
2	1	3	Shallow Surface Casing Top of Production Casing Cement Below Under-Pressured Hydrocarbon Reservoir	
3	2	3	Shallow Surface Casing Top of Production Casing Cement Above Top of Gas	
4	2	3	Shallow Surface Casing Top of Production Casing Cement Above Surface Casing Shoe	
5	3	3	Deep Surface Casing Top of Production Casing Cement Below Under-Pressured Hydrocarbon Reservoir	
6	3	3	Deep Surface Casing Top of Production Casing Coment Above Top of Gas	
7	4	3	Deep Surface Casing Too of Production Casing Composit Above Surface Casing Shoe	

Wells in all basins in Colorado were categorized by construction process. This includes determining the number of well control barriers in place. The categories allowed for the pinpointing of potential issues with specific types of well bore construction processes.

	CRIGHINAL WELLCOLAT	CATASTINOPHIC BARMER FAILURES	CAENSTROPHIC FAILURE 95
CATEGORY 1	0	0	0.00%
CREASERY 2	-41	0	0.00%
CATEGORY 3	145	2	1.56%
COLUMN RF 4	10 10	9	9,00%
CREASED S	1,769	2	0.11%
CREBSORY 6	5,235	4	0.08%
CATESCRY 7	1,862	ĩ	0.05%
CALIBREAKA S	8	9	9.50%
CATEGORY 9	\$0	û	0.02%
CATEGORY 19	0	0	0.00%
CATEGORY 11	105	0	0.00%
CATERORY 12	1	9	0.00%
TOTAL	10,912	9	0.06%
A TOTAL WELLS	194 18,998		

The historical probability of catastrophic wellbore failure in the Piceance Basin is 8 out of 10,000 wells.

















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Wellbore Stress Modeling Project Publications

- Fleckenstein, W.W., Eustes, A.W., Rodriguez, W. J., Berger, A., Sanchez, F. J.: "Borehole Stresses in Cemented Wellbores", Exploration & Production – Volume 10 Issue 2, (October 2012), 27-31
- Fleckenstein, W.W., Eustes, A.W., Rodriguez, W. J., Berger, A., Sanchez, F. J.: "Cemented Casing: The True Stress Picture", AADE-05-NTCE-14, American Association of Drilling Engineers Annual Conference, Dallas, Texas, April 5-7, 2005
- Berger, A., Fleckenstein, W.W., Eustes, A.W., and Tronhauser, G.: "Effect of Eccentricity, Voids, Cement Channels, and Pore Pressure Decline on Collapse Resistance of Casing,", SPE 90045, Society of Petroleum Engineers Annual Technical Conference and Exhibit, Houston, Texas, September 26– 29, 2004
- Rodriguez, W. J., Fleckenstein, W. W., Eustes, A. W., "Simulation of Collapse Loads on Cemented Casing", Journal of Petroleum Technology, (August 2004): 59-60
- Rodriguez, W. J., Fleckenstein, W. W., Eustes, A. W., "Simulation of Collapse Loads on Cemented Casing Using Finite Element Analysis", SPE 84566, Society of Petroleum Engineers Annual Technical Conference and Exhibit, Denver, Colorado, October 5-8, 2003.
- Fleckenstein, W. W., Eustes, A. W., et al., "Burst-Induced Stresses in Cemented Wellbores." SPE Drilling & Completion (June 2001): 74-80.



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- Patents
 - Downhole Tools and Methods for Selectively Accessing a Tubular Annulus of a Wellbore (US 8,991,502 B2) – Basis of Startup "FracOPTIMAL" – sold development option to major service company.
- Pending Patent Applications
 - o Method and Apparatus for Accessing a Tubular Annulus of a Wellbore
 - o Method and Apparatus for Testing a Tubular Annular Seal
 - o Method and Apparatus for to Rotate Subsurface Wellbore Casing




























































































































































Focus – Fluid Dynamics and Fluid Properties

- Research Directions
 - o Computational and experimental multiphase fluid dynamics
 - Phase behavior of petroleum systems
- Current Projects
 - Multiphase flow in porous media
 - Particle transport in porous media
 - Petroleum fluids phase behavior
 - o Nano-scale flows
 - Particulate flows
- Current Students
 - o 8 PhDs, 1 MS, 2 undergraduates

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 - $_{\rm O}\,\text{Mr.}$ Joe Chen




Laboratory Facilities and Instruments

Available for Research

1st Edition, January 2017

Somayeh Karimi, PhD Candidate

Ilkay Eker, PhD Candidate

and

Petroleum Engineering Faculty

Table of Contents

Preface	76
Petroleum Engineering Department, CSM	
Rock Property Measurements	
Core Lab CMS TM -300 Core Measurement System	77
rolosity, permeability and kinkenberg permeability	
Core Lab PDPK TM -400 Core Measurement System Permeability of core slabs and whole cores	78
Gas Adsorption Gas adsorption isotherms for source rocks, and other material such as aluminum silicate	79
Micromeritics ASAP 2020TM Unconventional reservoir rock properties such as pore size distribution, specific surface area, and adsorption/desorption isotherms	80
Quadrasorb EVO/ SI Specific surface area, pore size distribution, total pore volume, and adsorption/desorption isotherms	81



	Low Frequency Velocity Measurements Rock elastic properties such as Young modulus, bulk and shear modulus, and Poisson's ratio	82
	NER AutoLab 1500 Electrical resistivity, permeability, and compressional and shear wave velocities	83
	XRadia MicroXCT-400 High resolution 3D images from rock samples	84
Flu	id Pronerty Measurements	
110	Microfluidic and Nanofluidic Porous Media Analogues Phase behavior and enhanced oil recovery in micro and nano-scale	85
	Slim Tube System Minimum Miscibility Pressure	86
	Core Lab PVT SYSTEM 400/1000 Hydrocarbon fluid Phase behavior	87
	DBR Gasometer Gas volume at ambient pressure and temperature	89
	Anton Paar DMA 4200 M Densitometer Live oil density	90
	KRÜSS Spinning Drop Tensiometer SITE100 Interfacial tension of surfactant-oil systems	91
	Fluid Measurement System (FMS) Acoustic velocities in fluids (brine, oil and gas) at reservoir pressure and temperature conditions	92
Ro	ck and Fluid Property Measurement	
	Core Lab ACES-200 Automated Centrifuge System Capillary pressures, residual fluid saturations, end-point relative permeabilities, fluid recovery, and wettability	93
	Mass-based Spontaneous Imbibition Device Spontaneous imbibition measurements	94
	Chandler Formation Response Tester (FRT) 6100 Core Flooding System Absolute and relative permeability, and oil recovery experiments	95



Linear Core Flooding System for Formation Damage Studies Fluid and gas permeabilities, and fluid recovery	96
KRüSS Drop Shape Analyzer DSA100 Contact angle between two immiscible fluids and a polished solid surface (i.e. rock), and interfacial tension between two immiscible fluids	97
The Spectral Induced Polarization (SIP) Resistivity Measuring Instrument Complex resistivity of saturated rocks in the frequency range of 1 mHz to 45 KHz	98
Impedance Analyzer – Keysight E4990A Complex resistivity of saturated rocks in the frequency range of 20 Hz to 120 MHz	99
Network Analyzer – Keysight ENA Series Complex resistivity of saturated rocks in the frequency range of 300 KHz to 20 GHz	100
Anisotropic Acoustic – Electrical Joint Measurement System P- and S-wave velocities and complex electrical conductivity as functions of angle	101
Magritek [®] Low-Field 2-MHz NMR Porosity, pore size distribution, hydrocarbon compositions and fluid viscosity	103
UNGI Coupled True Triaxial Core Measurement Assembly Compressional and shear wave velocities, deformation, resistivity, permeability in three orthogonal directions	104
UNGI Coupled Pore Pressure Penetration Apparatus (Triaxial Assembly) Lateral and axial deformations, compressional and shear wave velocities, resistivity and permeability under triaxial stress state with confining and pore pressures at high temperatures	105
Drilling, Hydraulic Fracturing and Production Simulation Laboratory Research and Education Sandvik DE130 Diamond Core Drilling Rig Drill horizontal and vertical wells	106
Hydraulic Fracturing System with Tri-axial Stress Loading Hydraulic fracture initiation and propagation and its properties such as pressure and temperature profile during injection and rate of leak-off	107
Proppant Transport in Horizontal Wellbores & Perforations Proppant transport in horizontal wells and perforations	108
Proppant Transport in Complex Fracture Network Proppant transport and concentration in complex fracture network	109



	Artificial lift Experimental Facility A new artificial lift equipment to test the process of removing stagnated liquid from low flow rate horizontal gas wells	110
	6-in. Flow Loop Experimental Facility The effects of well trajectory on flow behavior in a test section	112
<u>Ou</u> 1	tside Petroleum Engineering Department, CSM JEOL JSM-700F Field Emission Scanning Electron Microscope (FE-SEM) Micro and nano-scale images of samples, texture, fabric, and morphology of the grains, matrix, and cement	115
	JEOL IB-0910CP Cross-Section Polisher Rock surfaces are polished and prepared for SEM imaging using the polisher.	116
	FEI QUANTA 600I Environmental Scanning Emission Microscope (E-SEM) Crystallization or phase transformation, hydrate sample imaging, and particles in suspension	117
	TESCAN Integrated Mineral Analyzer (TIMA) and FEI QEMSCAN Particle and grain size and shape, element X-ray mapping, lithotyping	118
	TESCAN MIRA3 LMH Schottky Field Emission Electron Microscope (FE-SEM) Nanometer scale images, topography and compositional analysis	119
	High-Field Nuclear Magnetic Resonance (NMR) Information about structure characterization and molecular dynamics	120



Preface

The purpose of this booklet is to inform the Petroleum Engineering Department faculty and students with a list of the laboratory facilities and equipment available for research in PE department and other CSM departments. A brief description of each instrument and a photo of the instrument is presented. Furthermore, we intend to promote experimental research in the department.

Please contact Joe Chen (jychen@mines.edu) for assistance in coordinating use of the devices.



Core Lab CMS[™]-300 Core Measurement System

The CMS[™]-300 Core Measurement System is an automated unsteady state, pressure decay, permeability and porosity measurement instrument. The instrument can be used to conduct measurements under confining stress to simulate reservoir conditions. Measurements can be conducted on cylindrical cores with 1 or 1.5" diameter and 0.75 to 3.12" length. Helium and Nitrogen are used to conduct the measurements.

The following is a partial list of parameters that can be determined with the instrument:

- 1- Porosity
- 2- Permeability
- 3- Klinkenberg permeability



Figure 1: Core Lab CMS[™]-300 Core Measurement System (image adapted from catalog)



Core Lab PDPK[™]-400

The PDPK[™]-400 Pressure-Decay Profile Permeameter is a pressure decay system that is used to determine core permeability and heterogeneity. The instrument can be used to measurement permeability in x and y direction on a core as long as 3 meters. Cleaned and dried core slabs as well as whole cores can be used in measurements. The range of rock permeability for reliable measurements is 0.001 mD to greater than 30 D.



Figure 2: Core Lab PDPK[™]-400, Pressure-Decay Profile Permeameter (image adapted from catalog)



Gas Adsorption

The gas adsorption system is used to determine adsorption isotherms of different solvents on source rocks such as shales and coal bed methane. Zeolite 13x, an aluminum silicate or ceramic material which is a strong adsorbent, may be used for demonstration purposes. The system is manually operated and consists of high accuracy pressure gauges, two stainless steel open cylinder cells, stainless steel lines and high pressure needle valves all rated to 3500 psi. The two cylinders are referred to as the dosing cell and the uptake cell. The system is also designed to fit wholly into a temperature water bath allowing for temperature regulation during an experiment. Ultimately, the gas storage capacity of the samples can be determined at different pressures through material balance calculations.

The combination of the dosing cell volume, uptake cell volume, valve and the line volumes represent the system volume. For high pressure experiments, minimum sample cell cylinders volumes (available from Swagelok) are able to withstand pressures up to 3000 psi is 150 cc. In this case, a large amount of sample is required to reduce the system volume to sample ratio. Generally, subtle reduction in the amount adsorbed by the adsorbent material is experienced when system volume is lowered. Optimum ratios for the dosing cell and uptake cell have to be determined prior to experiment.



Figure 3: Gas Adsorption System



Micromeritics ASAP 2020[™]

This Micromeritics ASAP 2020TM is a gas adsorption instrument designed for core samples with very small pore sizes that range between 3 to 200 nanometers. It is mostly useful to measure the pore sizes of unconventional reservoir rocks. It has two outgassing and one analysis ports. Depending on the pore sizes, each sample measurement can take from a few hours (only surface area) to a few days (nanometer-sized pores in organic matter). The samples are measured in their powder form, and the gases that can be used for this instrument are Argon, N₂, Hexane, CO₂, and Water Vapor. The samples are first outgassed for 24 hours at 200 °C under vacuum condition while flowing inert gas through the sample to remove any contaminant gas. Then, the sample is subjected to increasing partial pressures to obtain a full isotherm. The following is a list of parameters which can be measured using ASAP 2020TM:

- 1. Specific Surface Area
- 2. Isotherm Adsorption & Desorption
- 3. Pore Size Distribution
- 4. Total Pore Volume
- 5. Average Pore Size



Figure 4: Micromeritics ASAP 2020[™]



Quadrasorb EVO/ SI

The Quadrasorb EVO/ SI is a versatile gas adsorption instrument designed for high throughput. With four independent analysis stations it is capable of analyzing four samples simultaneously. Quadrasorb EVO/ SI is available in a KR/MP configuration, which is equipped with a turbomolecular pump and 10-torr or 1-torr pressure transducer for measuring low surface area samples with krypton and microporous samples with nitrogen or argon. The following is a partial list of important parameters which can be determined from Quadrasorb Adsorption experiments:

- 1- Specific Surface Area
- 2- Isotherm Adsorption & Desorption
- 3- Pore Size Distribution
- 4- Total Pore Volume
- 5- Average Pore Size

Flovacdegasser is used in conjunction with the QUADRASORB EVO/ SI for outgassing at vacuum condition purposes.



Figure 5: QUADRASORB EVO/ SI (image adapted from user manual)



Low Frequency Velocity Measurements

The technique used to determine elastic properties at low frequencies consists of a stress/strain system that deforms the rock at a frequency range of 1 to 2000 Hz. Measurements can be conducted at different confining and pore pressure stages to simulate reservoir conditions. Additionally we can control temperature of the pressure vessel (0 – 100 °C). Besides measuring low frequency velocities we are also able to perform ultrasonic (1 MHz) velocity measurements.

The following is a partial list of parameters that can be determined using the set up:

- 1- Young's modulus
- 2- Poisson's ratio
- 3- Bulk modulus
- 4- Shear modulus
- 5- Compressional and shear wave velocities
- 6- Attenuation





Figure 6: (a) Schematic of the low frequency measurement assembly. For seismic frequencies, strains are measured on both the sample and aluminum standard. Ultrasonic transducers permit wave propagation measurement near 1 MHz. Fluid lines permit control and exchange of pore fluids independent of confining pressure, (b) Photograph of completed measurement assembly



NER AutoLab 1500

AutoLab 1500 is a servo-hydraulic operated system for biaxial measurements with software-controlled arbitrary stress paths on rock specimens up to 50.8 mm (2.0 in) in diameter at in-situ stress conditions with pore pressure and temperature controls. The high pressure biaxial system consists of a pressure vessel with an internal piston for differential stress and servo-hydraulic intensifiers for differential stress, confining and pore pressure. This instrument allows us to make measurements at reservoir pressures up to 69 MPa (10,000 psi) and temperatures up to 120°C (248°F).

The key features of the instrument are the following:

- Servo-hydraulic control of confining pressure, pore pressure, flow rate, strain rate, and force.
- Control of stresses and temperatures at reservoir conditions.
- Pore pressure intensifier compatible with water, brine, oil, and gas (including CO2).
- AutoLab software for system data acquisition and reduction.
- Integrated electronics console for servo-amplifiers and signal conditioning.

A list of the parameters measured using NER Autolab 1500 are as following:

- 1. Compressional and shear wave velocities
- 2. Electrical resistivity
- 3. Permeability



Figure 7: NER AutoLab 1500



XRadia MicroXCT-400

The micro X-ray CT (computed tomography) machine provides high resolution 3D images of samples with up to 2 " in diameter. The weight of the sample must be less than 15 kg. Image resolution of up to 1 μ m can be achieved. Four different lenses allow us to cover a varying field of view size and a variable resolution. It is a non-destructive method to explore the internal structure of samples which might be unobservable with conventional 2D techniques, such as SEM.

The instrument chamber can house a pressure cell for studying in-situ change in internal microstructure with application of pore and confining pressures up to 2500 psi as well as a temperature controls between -5 °C to 50 °C. The following is a list of parameters that can be measured using the instrument:

- 1. Rock microstructure for oil and gas exploration
- 2. In situ measurement during imaging (e. g. ultrasonic velocities)
- 3. Semiconductor packaging development and failure analysis
- 4. Life-science research
- 5. Advanced material characterization



Figure 8: XRadia MicroXCT 400 (image adapted from catalog)



Microfluidic and Nanofluidic Porous Media Analogues

Microfluidics and nanofluidics are technologies developed to control and manipulate fluids at micro- and nanoscale. Using micro- and nanofabrication, pores of controlled dimensions, surface properties, and complexities can be made on silicon or polymer substrates to facilitate direct visualization and fundamental studies of fluid flow through porous media. Microfluidic porous media analogues (μ PMA) are primarily polymer based and the pore size ranges from a few micrometers to hundreds of micrometers. Nanofluidic porous media analogues (nPMA) are built by bonding silicon to pyrex, and the channels have critical dimensions in the range of tens to hundreds of nanometers. These devices are being used to study enhanced oil recovery, phase behavior, and nano-scale single- and multiphase flows in unconventional reservoirs.



Figure 9: A – μ PMA "chips" being prepared on a master substrate; B – A fully assembled μ PMA; C – A section of pore network etched onto μ PMA; D – Water flooding pattern in the same section; E – SEM image of channels in a μ PMA



Figure 10: Pore networks on nPMA. A – 300 nm channels; B – 30 nm channels



Slim Tube System

The Slim Tube is equipped with a temperature controlling enclosure, with both 60' and 10' 1/4" slim tubes (60' for lighter oils, 10' for v. heavy oils), a constant pressure backpressure system, a constant volume positive displacement pump and the "cannon" the large SS tube that pressurizes the gas to the desired pressure and utilizes water to displace the gas into the slim tube at a constant rate, constant pressure, and constant temperature. Minimum miscibility pressure of liquid hydrocarbon and gases such as N2, Co2, and hydrocarbon gases are measured using the slim tube.



Figure 11: Slim Tube System



Core Lab PVT SYSTEM 400/1000

This PVT system can operate at reservoir pressure and temperature conditions up to a maximum of 14,500 psi and 400° F. The PVT cell is equipped with a camera that provides full visual observation of PVT experiments. We use the PVT cell to characterize reservoir fluid (oil and gas) systems, and to determine the effect of different gases (such as N2, CO2 and pure hydrocarbon components) on the reservoir oil behavior. The following is a partial list of important parameters which can be determined from PVT experiments:

- 1- Bubble point pressure
- 2- Dew point pressure
- 3- Gas oil ratio
- 4- Formation volume factor
- 5- Oil and gas compositions

We must use auxiliary instruments such as a gasometer, gas chromatograph (GC), and densitometer in conjunction with PVT cell experiments.



Figure 12: Core Lab PVT System 400/1000 (Core Lab catalog)





Figure 13: PVT cell at 45 ° from horizontal to capture condensate accumulation, and a view of the cell interior on the top left corner (modified from Core Lab catalog)



DBR Gasometer

A sample of a live oil (or gas) sample from the PVT cell is transferred to the gasometer. Then it is slowly flashed to the ambient conditions to measure the resulting gas and liquid volumes for GC analysis. The measured volume is used to determine GOR. The gasometer cylinder capacity can increase up to 10 liters at ambient pressure and temperature conditions.



Figure 14: DBR Gasometer



Anton Paar DMA 4200 M Densitometer

The densitometer is an automated instrument used to measure oil density at different pressure and temperature conditions. The range of operating temperature for our densitometer is -10° C to 200° C. The pressure of the sample can be up to 500 bar (~ 7200 psi). The sample volume required for measurements is 2 cc.

Method: DMA4200 Density Sample:	8:33.49 PM Administrator		
Density	Periodic Time 2409.051 us	~	
Density (not visccorr.) 0.00076 g/cm ²	Cell Temperature 20.00 °c	1	
Density Condition	Pressure 16.53 psi		
* API Gravity 60 *F Out of range	-	2	
		Check	
Menu	Quick Method	Start	
	and an		

Figure 15: Anton Paar DMA 4200 M Densitometer



KRüSS Spinning Drop Tensiometer SITE100

The spinning drop tensiometer is an instrument to measure very low IFTs of the oil-water systems in the range of 10 to 10^{-6} mN/m. Such systems generally contain surface active agents. The spinning speed of the capillary tube can go up to 15,000 rpm for measurements. We can decrease or increase temperature using a recirculating bath. The temperature for measurements can vary in the range of -20° C to 200° C.



Figure 16: KRÜSS Spinning Drop Tensiometer SITE100



Fluid Measurement System (FMS)

The fluid measurement system (FMS) is used to measure acoustic velocities in fluids at reservoir pressure and temperature conditions up to a maximum of 20,000 psi and 200° C. The FMS cell is equipped with a computer that provides full control and data acquisition of the FMS experiments. We use FMS to characterize reservoir fluid (brine, oil, and gas) systems, and to determine the effect of different gases (such as methane, N₂, CO₂ and pure hydrocarbon components) on reservoir oil behavior. The following is a partial list of important parameters which can be determined from FMS experiments:

- 1- Fluid velocity under pressure change
- 2- Fluid velocity under temperature change
- 3- Fluid permeability



Figure 17: Fluid Measurement System (FMS)



Core Lab ACES-200 Automated Centrifuge System

This high speed centrifuge system is capable of spinning cores at different rotational speeds (maximum practical speed is 13,000 rpm). The temperature can also be increased from the ambient temperature to the reservoir temperature. The centrifuge can use conventional (high porosity-high permeability) core plugs as well as unconventional (low porosity-low permeability) core plugs. The centrifuge is equipped with a high resolution camera which makes it possible to measure small produced fluid volumes from cores. Using rotor PIR 16.5, experiments on three core plugs can be conducted simultaneously. The core plugs dimensions should be maximum 1.5" diameter and up to 2" length.

We have used the centrifuge to determine the wettability related properties of the oil-water and gasliquid flow in cores. Gravity drainage (or, the fluid replacement concept) is the major outcome of the centrifuge experiments. Specifically, the following is a partial list of important parameters that can be determined with the centrifuge:

- 1- Drainage and imbibition capillary pressure curves for oil-water and gas-liquid systems
- 2- Relative permeability end-points
- 3- Oil recovery by gravity drainage
- 4- Pore size distribution
- 5- Wettability



Figure 18: Core Lab ACES-200 Automated Centrifuge System



Mass-Based Spontaneous Imbibition Device

The mass-based spontaneous imbibition experimental setup is used to measure fluid mass produced in spontaneous imbibition experiments. The equipment can be used to study fluid-rock interaction at room conditions. Various salinity brines, oil, dilute acid, and surfactant are some of the fluids used to study fluid-rock interactions in spontaneous imbibition experiments. Moreover, rock samples such as carbonate, sandstone, or shales with quantity as low as of 5 gm up to 200 gm are used in spontaneous imbibition experiments.



Figure 19: Mass-based spontaneous imbibition device



Chandler Formation Response Tester (FRT) 6100 Core Flooding System

There are two core flooding systems at PE research laboratory facilities. One of the core flooding systems is used in the reservoir characterization laboratory for reservoir engineering studies. The core flooding apparatus is used to determine core properties and the oil recoveries using water displacing oil or N₂ and CO_2 displacing oil-water mixture. In addition, different salinity brines and surfactant and polymer solutions can be used in core flooding experiments. The second core flooding system in the formation stimulation laboratory is used for stimulation and formation damage studies.

Our core flooding systems can operate at reservoir pressure and temperature conditions up to a maximum of 5,500 psi injection pressure, up to a maximum of 6,000 psi confining pressure, and 300° F temperature. The core plugs dimensions used in experiments are 1 and 1.5″ diameter and up to 12″ in length.

The following is a partial list of the parameters that can be determined from core flooding experiments:

- 1- Absolute permeability
- 2- Relative permeability
- 3- Oil recovery



Figure 20: Chandler Formation Response Tester (FRT) 6100 Core Flooding System



Linear Core Flooding System for Formation Damage Studies

The core holder is used to study formation damage at reservoir pressure conditions with a confining pressure of up to 6000 psi, and pore pressure up to 5500 psi. It operates at room temperature at this point; however, it can be upgraded to reservoir temperature condition. Nitrogen gas is used as pore pressure fluid (helium or argon gas can also be used). We are capable of using cores with up to 4" in diameter and 12" in length.



Figure 21: Linear core holder unit



KRüSS Drop Shape Analyzer DSA100

Our DSA100 unit is an automated instrument which gives us the capability to measure contact angle as well as liquid surface tension and calculates IFT. The contact angle is measured between two fluids (usually formation/synthetic brine and oil) and a polished solid surface such as rock. In addition, using the Minidosing MS PD-E1700, we can measure the parameters at pressure and temperature conditions up to maximum 10,000 Psi and 180° C.



Figure 22: KRüSS Drop Shape Analyzer DSA100



The Spectral Induced Polarization (SIP) Resistivity Measuring Instrument

The SIP resistivity measuring instrument measures the real and imaginary components of resistance for brine-saturated porous rocks in the frequency range of 1 mHz to 45 KHz.

The following is a partial list of the important parameters which can be determined from SIP experiments:

- 1- Resistivity
- 2- Tortuosity



Figure 23: Resistivity Measuring Instrument

Impedance Analyzer – Keysight E4990A

This Impedance Analyzer is a complex resistivity measurement used for bench top measurements in a range of frequencies (20Hz to 120MHz). The analyzing probe can be used for samples with a maximum length of 11mm and variable diameters. The following is a partial list of important parameters which can be determined from these experiments:

- 1- Conductivity
- 2- Permittivity

We use auxiliary instruments, for example Refractometer, to measure fluid conductivity if the sample analyzed is saturated.

Figure 24: Impedance Analyzer (Keysight manual)

Network Analyzer – Keysight ENA Series

This Network Analyzer is a complex resistivity measurement used for bench top measurements in a range of frequencies (300 kHz to 20 GHz). The analyzing probe can be used for any sample; the only specification is the need for a flat polish surface, to allow full contact with the sample and reliable measurements. The following is a partial list of important parameters which can be determined from these experiments:

- 1- Conductivity
- 2- Permittivity

Figure 25: Network Analyzer (Keysight manual)

Anisotropic Acoustic–Electrical Joint Measurement System

We use the system to characterize the geophysical rock properties and anisotropy, and to determine the effect of pressure and fluids on reservoir rock physics parameters. Specifically, the joint anisotropic acoustic-electrical measurement system is used to measure P- and S-wave velocities and complex electrical conductivity as functions of angle simultaneously on rock samples at various confining and pore pressure stages up to 4000 psi. The following is a list of parameters measured using the instrument:

- 1- Compressional and shear wave velocities and their anisotropies
- 2- Formation elastic properties and stiffness tensor
- 3- Acoustic attenuation and attenuation tensor
- 4- Complex electrical conductivity and conductivity tensor

Figure 26: Schematic of anisotropic acoustic – electrical joint measurement system

In the acoustic acquisition system, the pulser initiates a pulse signal to the acoustic transducers and trigger signal to oscilloscope isochronously, while the recording equipment, or oscilloscope (Tektronix TDS 3014C), receives signals from acoustic transducers and trigger signals from the pulser in different channels, and displays the wave amplitudes as a function of time.

The complex conductivity acquisition system mainly consists of the Spectral Induced Polarization (SIP) system, which comprises a four-channel acquisition array and nominal frequency range of 1 mHz to 45 kHz.

Figure 27: Electrode array and circuit simplified diagram of SIP system. 1 and 4 represent current electrodes, 2 and 3 represent potential electrodes. (Adapted from Zimmermann et al., 2008)

Magritek[®] Low-Field 2-MHz NMR

This Nuclear Magnetic Resonance (NMR) system operates a frequency of 2MHz and a magnetic field strength of 0.05T. For saturated porous media such as rocks and soils, the NMR response of this system is dependent on the size of the pore space as well as the hydrogen index of the saturating fluid. This non-destructive measurement leaves the core completely intact, while detecting hydrogen nuclei contained in the pore space through alternating magnetic fields. This allows for pore space properties to be determined without alteration of the core or pore space environment. Additional information about interstitial fluids and core mineralogy can be determined with some NMR methods. The following is a partial list of important parameters which can be determined from NMR experiments:

- 1- Pore Size Distribution
- 2- Porosity
- 3- Permeability
- 4- Fluid Viscosity
- 5- Oil and Gas Compositions

These results are commonly used in conjunction with nitrogen adsorption and resistivity studies for a complete analysis of pore-space in both laboratory and downhole logging applications.

Figure 28: 2-MHZ Magritek NMR laboratory setup for core analysis (Magritek website)

UNGI Coupled True Triaxial Core Measurement Assembly

The true triaxial measurement assembly is a unique experimental apparatus loaned to UNGI Geomechanics Research Laboratory by Dr. Ali Mese of Geomechanics Engineering and Research, PLLC. Cylindrical core samples are used in the apparatus under true triaxial stress conditions with capability of three independent principal stress magnitude application with elevated pore pressure.

The apparatus is designed to use cylindrical core samples of 2" in diameter with varying lengths with the two independent orthogonal stress magnitudes in the horizontal plane loaded with fluid pressure. The stress magnitudes applied are limited to the pressure capacities of the ISCO hydraulic syringe pumps with 0.001 psi precision. Pumps with capability of 20,000 psi, 10,000 psi and 7,500 psi are currently utilized in the true triaxial measurement assembly to implement the three principal stress magnitudes and pore pressure, respectively. The stress magnitudes can be increased using different pumps with higher pressure capacity and/or placing the standalone sample cell into the MTS load frame at the UNGI Geomechanics Laboratory if core samples needs to be tested at higher stress state. The entire assembly sits in an enclosed insulation box allowing precise temperature control with elevated temperature measurements.

The following is a list of the parameters we can determine using the instrument:

- 1- Compressional and shear wave velocities
- 2- Deformation
- 3- Resistivity
- 4- Permeability in three orthogonal directions
- 5- Differences between static and dynamic moduli
- 6- Permeability anisotropies as a function of stress

Figure 29: Coupled True Triaxial Core Measurement Assembly

UNGI Coupled Pore Pressure Penetration Apparatus (Triaxial Assembly)

The coupled pore pressure penetration assembly is a sample cell for core samples to be tested under triaxial stress state and was donated to UNGI Geomechanics Laboratory by Dr. Ali Mese of Geomechanics Engineering and Research, PLLC. The apparatus has been utilized for cylindrical core samples of 1.5" diameter with varying lengths to conduct coupled measurements of lateral and axial deformations, compressional and shear wave velocities, resistivity and permeability under triaxial stress state with confining and pore pressures up to 20,000 psi and 10,000 psi, respectively under controlled elevated temperatures up to 90°C (within 0.03°C precision) using an insulation box for the full assembly. The two hydraulic lines with a porous filter at the bottom cap allows circulation of any pore fluid at specific elevated pressure.

We can measure differential pressure at the top of the sample while different fluids are circulated at the bottom. The data used to study rock-fluid interaction and osmosis, while the data measured while circulation is happening within the sample is used for permeability and directional resistivity measurements.

Axial, confining, pore and circulation pressures as well as injection and back pressure systems are controlled by independently computerized ISCO syringe pumps with 7,500 psi to 20,000 psi capacities. The mixing of the pore fluid brines containing salts and/or chemical additives and the pressurizing hydraulic fluid (mineral oil or deionized water) is prevented in individual pumps as well as between the pressurizing fluid and the core sample for accurate measurements of true rock- pore fluid interactions. Real time monitoring of the pressure, deformation, resistivity and permeability is accomplished for the long duration of the experiments to capture the equilibrium state reached in shale/mudrock samples.

Figure 30: Coupled Pore Pressure Penetration Apparatus (Triaxial Assembly)

Research and Education Sandvik DE130 Diamond Core Drilling Rig

This rig is sponsored by Apache Corporation for research and education of the Mines Community. The Sandvik DE130 is a drilling unit for both surface and underground applications. It can drill in any orientation from horizontal and vertical and at depths of up to 1,200 m with the "N" rotary drive. It can be used to wireline core to 1,200m. The unit can push with 10,350 lbf and pull with 13,820 lbf. It is capable of a drilling torque of 137 to 619 ft-lbf. The hydraulic system has a maximum working pressure of 3,625 psi. The fluids system uses a FMC L1122 Triplex capable of 75.1 gpm at 1,000 psi.

Figure 31: Sandvik DE130 Diamond Drill Core

Figure 32: Mines' DE130 delivered on September 6, 2016

Hydraulic Fracturing System with Tri-axial Stress Loading

The hydraulic fracturing testing system, can be used to investigate fracture initiation and propagation in samples up to 8 inch \times 8 inch \times 8 inch with tri-axial stress conditions. The maximum stress loading that can be applied in x, y, and z directions are 4500, 4500, and 6000 psi, respectively. Temperature can be adjusted from the ambient condition to about 60 °C. The system is equipped with an ISCO pump to provide the injection fracturing pressure. The following is a partial list of parameters that are measured from experiments:

- 1- Pressure profile during injection
- 2- Temperature profile during injection
- 3- Rate of leak-off before and after fracturing
- 4- Acoustic wave propagation
- 5- Fracture pattern and fracture morphology

Figure 33: Hydraulic fracturing system with tri-axial stress loading


Proppant Transport in Horizontal Wellbores & Perforations

Our multiple cluster perforated horizontal wellbore is used to study proppant transport. The slurry is mixed in the tank, and run through the 30-foot long polycarbonate. Sand which exited through the perforation clusters are collected and analyzed to study proppant transport in multi-cluster perforated horizontal wellbores.



Figure 34: Schematic of proppant transport in horizontal wellbore experimental setup



Figure 35: Images of proppant transport in horizontal wellbore experimental setup



Proppant Transport in Complex Fracture Network

We can study proppant transport in complex fracture systems. The experimental setup can be used to study proppant concentration and proppant transport behavior in each fracture network as well.





Figure 36: Schematic and images of experimental setup used to study proppant transport in a complex fracture system



Artificial Lift Experimental Facility

This experimental facility has been designed to test the performance and operation of new artificial lift equipment. The purpose of this device is to remove stagnated liquids from horizontal gas where low flow pressure and low flow rates (5 to 10 BPD approx.) are expected. The device prototype consists of a mandrel with two check valves that allow the injection of gas to remove stagnated liquids through a double tubing arrangement.



Figure 37: Gallop Installation Scheme

The mandrel is installed inside a 4' ID casing which contains a water inlet valve that allows the entrance of low pressure liquids (less than 20 psi) form a continuous water supply line. As the liquid level reaches certain height within the casing, the mandrel allows the liquids to flow towards the tubing. Once the tubing is filled with water, compressed air from an external line is injected (between 30 and 60 psi of pressure) into one of the tubes, pushing the liquids along the horizontal section towards a 42' long crystal PVC vertical line of $\frac{3}{4}$ ' ID. This vertical section is followed by a recollection pipe which allows the injected gases out and measures the volume of liquids produced through a pressure transducer.

The designed instrumentation for the facility should allow us to measure the ratio of injected gas to recovered fluids. While continuous operation is done at 65 °F and under 60 psi, the facility has been designed to operate at up to 80 psi and up to 140 °F. Future modifications to the layout include the considerations for analyzing changes on the inclination of the horizontal section of the well.



Figure 38: Casing with Mandrel inserted



Figure 39: Top of the facility



The facility has been designed to recirculate the water inject/produced so to minimize the environmental footprint.



Figure 40: Schematic of the experimental test facility

Table 1: Fluids, operating conditions, and instrument			
Fluids	Gas: Air Liquid: Tap water		
Operating Conditions	 Pressure: 60 psi (Max 100 psi) Temperature: Ambient (Max 140 F) Gas flow rate (in): 0 to 0.03 MMSCFD (Superficial vel.: 0 to 95 ft/s) Water flow rate (in): 0 to 11,314 BBPD (Superficial vel.: 0 to 150 ft/s) 		
Test Section	 Casing material: PVC Casing dimensions: 4 in ID, 10.6 ft long Production and injection lines: ¾ in ID. 		



6-in. Flow Loop Experimental Facility

The facility operates with gas (Air), water (Tap Water) and oil (Low Viscosity Mineral Oil), and has been designed to study the effects of well trajectory on flow behavior in a test section, which is made of transparent acrylic pipe to observe the flow pattern along the flow loop. The inclination angle can be changed with a 3-ton capacity electric hoist. Based on the inclination angle, several well configurations can be simulated in this facility. Operating maximum pressure is 35 psi. The tests are conducted at ambient temperature. However, the operating conditions change depending on the project.

The experimental facility is designed to handle the following liquid and gas flow rates:

Fluid	Maximum Flow Rate	Minimum Flow Rate	Source		
Tap Water	35 gpm	2.9 gpm	Water Tank + Electric Submergible Pump		
Mineral Oil	35 gpm	2.9 gpm	Oil Tank + Progressive Cavity Pump		
Air	1150 CFM	3.9 CFM	Air Blower		

Table 2: Expected Flow Conditions

Test Section

Pipe Material: Acrylic Diameter of Pipe: 6-in Test Section: 32 ft Inclination Angle: 0 – 90 degree Maximum number of undulations: NA

Measured Parameters	Instrumentation		
Liquid Holdup	Quick Closing Valves		
	Conductivity Probes		
Elow Pattorn	Surveillance Cameras		
	High Speed Camera		
Pressure Gradient	Differential Pressure Transducer		
Slug Flow Characterization (translational velocity, slug length and frequency)	Conductivity Probes		
Severe Slugging Characterization (Cycle duration, slug frequency and maximum expected pressure)	Pressure Transducers		

Table 3: Instrumentation and Flow Characteristics





Figure 41: Schematic of 6 in ID flow loop

Laboratory Facilities and Instrument Available for Research Outside Petroleum Engineering Department

O PETROLEUM ENGINEERING

JEOL JSM-700F Field Emission Scanning Electron Microscope (FE-SEM)

The FE-SEM in the Electron Microscopy Laboratory in the Department of Metallurgical and Materials Engineering is a high resolution scanning electron microscope for nano analysis of structures and surface details. The instrument has a maximum resolution of 1.2 nm at 30kV.

The following parameters can be identified and analyzed using the FE-SEM on ion-milled samples:

- 1- Texture
- 2- Fabric
- 3- Morphology of the grains, matrix, and cement



Figure 42: Field Emission Scanning Electron Microspore (FE-SEM)



JEOL IB-0910CP Cross-Section Polisher

The JEOL cross-section polisher uses Argon ions to remove surface damage from a cut and grinds the surface of the specimen, creating a polished surface suitable for observation in a scanning electron microscope that is free of preparation artifacts.



Figure 43: The JEOL cross-section polisher



FEI QUANTA 600I Environmental Scanning Emission Microscope (E-SEM)

The E-SEM in the Electron Microscopy Laboratory in the Department of Metallurgical and Materials Engineering is a high performance instrument with three operating vacuum modes to accommodate a wide range of samples. The conventional mode is the High Vacuum Mode, which operates at up to 10^{-6} torr. The Low Vacuum Mode is used for imaging of non-conductive samples that cannot be coated with metal and the pressure specification for this mode ranges between 0.1 - 1.0 torr, using water vapor from a built-in water reservoir. The last mode is the Environmental Mode (E-SEM) where water vapor or auxiliary gas supplied by user can be used. The Environmental Mode operates between 1.0 - 10 torr. The E-SEM has a large chamber of 15" enabling the rotation of large specimens. The instrument also has a hot stage, with a temperature rating of 1500° C.

The following parameters can be identified and analyzed from E-SEM:

- 1- Imaging of hydrated samples
- 2- Crystallization or phase transformation
- 3- Particles in suspensions
- 4- Tensile testing, with heating or cooling



Figure 44: Environmental Scanning Emission Microspore (E-SEM)



TESCAN Integrated Mineral Analyzer (TIMA) and FEI QEMSCAN

Both TIMA and QEMSCAN instruments in Automated Mineralogy Laboratory in Geology & Geological Engineering Department is a fully automated SEM-based analysis system that provides quantitative mineralogical and textural data on the basis of automated point counting. The instrument contains a custom-built electron-beam platform equipped with four energy dispersive X-ray spectrometers (EDS) for mineral and compound identification within a wide range of sample types.

The software for each instrument allow for the automated stepping of the electron beam across samples at a user-defined pixel resolution (typically 1 - 40 micrometers). At each pixel, the system collects a backscatter electron (BSE) signal and an EDS spectrum. A mineral or phase identification is made on the basis of the BSE value and elemental intensities. Analysis was performed at an acceleration voltage of 25 kV and a specimen current of 5 nA for the QEMSCAN and a beam intensity of 14.5 for TIMA.

The analysis from these instruments provide quantitative mineralogical and textural data, false-color mineral maps, and robust statistical data which include

- 1. Highly accurate mineral (phase) abundance (i.e. modal abundance) maps
- 2. Element X-ray mapping
- 3. Particle and grain size
- 4. Particle and grain shape
- 5. Mineral associations
- 6. Lithotyping
- 7. Porosity quantification
- 8. Organic matter scans
- 9. Mineral (phase) liberation



Figure 45: TIMA



TESCAN MIRA3 LMH Schottky Field Emission Electron Microscope (FE-SEM)

The FE-SEM in the Department of Geology and Geological Engineering is a state-of-the-art high resolution scanning electron microscope for nano analysis of structures and surface details. The SE detector has a maximum resolution of 1.2 nm at 30kV and 2.5 nm at 3 kV, and the In-Beam SE detector has a maximum resolution of 1 nm at 30kV.

We use secondary electron imaging for topography contrast study, backscatter electron imaging for phase contrast, and energy-dispersive X-ray spectroscopy for compositional analysis.



Figure 46: Field Emission Scanning Electron Microspore (FE-SEM image adapted from CSM website)



High-Field Nuclear Magnetic Resonance (NMR)

The state-of-the-art NMR instruments (JEOL 500MHz liquid state [Pulse Field Gradient – PFG] NMR and Bruker 400MHz solid state NMR) in the Chemistry Department can provide important information for structure characterization as well as molecular dynamics study. We have one 5 mm multinuclear double tuned liquid state NMR probe, two 4 mm multinuclear triple/double tuned Magic Angle Spinning Solid State NMR probes, and one diffusion probe (single axial DIFF 60) equipped with exchangeable 1H, 7Li and 13C coils.

The Chemistry Department has used both NMR instruments to study molecular structures, molecular interactions, and ion/molecule diffusivity. In Petroleum Engineering, we have used the instrument to measured self-diffusion coefficients for a couple of components.



Figure 47: 500MHz liquid-state (left image-adapted from CSM website) and 400 MHz solid-state NMR (right image-adapted from CSM website)