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Paschal Ogadi Mokwenye

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Evaluation of Gas Hydrate in Gas Pipeline Transportation

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Bachelor of Engineering, Teesside University, 2016

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Submitted to the Graduate Faculty

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2020

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December 1, 2020

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To my mom Nelly Mokwenye and my dad Lawrence Mokwenye.

The world's greatest parents and my biggest motivators.

ABSTRACT

Gas hydrate formation and deposition is an inherent problem in the gas production and transmission industry. It causes plugging in pipelines and damages to equipment such as pumps and separators. The continuous constriction of pipelines by hydrates can cause a pressure difference between the upstream region behind the hydrate which has high pressure and the downstream region ahead of the hydrate which has low pressure, this pressure difference can cause pipelines to rupture, putting personnel and equipment in danger. The safest and cost-effective way of preventing hydrate formation is by predicting the temperature or pressure in which they form. In the past years, many correlations have been developed to predict the conditions of hydrate formation, but the models require a complex computation, or their accuracy is limited to certain gas mixtures. This study reviews the conditions of hydrate formation and numerous mathematical models developed to predict their formation temperature for natural gas. The study also involves the development of a new model to calculate the hydrate formation temperature of natural gas, the equation was developed by carrying out nonlinear regression of 460 experimental data points in MATLAB software. Hydrate formation temperatures calculated by the model were compared to experimental values as well as other reputable models. The proposed model showed superior accuracy in calculating the reference experimental hydrate formation temperature over models developed

by Hammerschmidt (1934), Motiee (1991), Towler and Mokhatab (2005), Safamirzaei M. (2015), and Chavoshi S. (2018). Also, the model produced the lowest Average Relative Deviation and Average Absolute Deviation from experimental values. Despite having three adjustable parameters the model produced an accuracy ranking of 1 to 3 across a range of specific gas gravity from 0.55 – 1 compared to other models. The model was also applied to gas compositions obtained from The Bakken and it showed a reliable prediction of the hydrate formation temperature when compared to the Katz gas gravity chart. The model offers a simple and good mathematical model for the prediction of hydrate formation temperature of the Bakken Formation.

CHAPTER 1 INTRODUCTION

A major problem in the gas industry is the effect of water vapor during the transportation and measurement of natural gas. Water liquefies, freezes, and accumulates within the system causing interruptions in pipeline transmission of gas. The solid crystals that form resemble snow and the flow of gas causes it to compress and accumulate at the lower spots of the pipeline, these snow-like crystals are called gas hydrates and they are known to cause plugging in gas pipelines. Figure 1.1 shows a typical example of a gas hydrate from a pipeline (Hammerschmidt, 1934).



Figure 1.1 Gas hydrate showing its snow-like appearance

Gas hydrates are clathrate structures in which guest molecules are enclosed in cages of a host lattice. They are crystalline forms of water in which the stabilization of the solid is facilitated by the presence of gas molecules inside the crystal matrix (cage) at temperatures significantly above the normal freezing point of water. It is composed of

gas molecules (methane predominantly) surrounded by a cavity of water molecules (Saleh, 2002). Gas hydrates usually form when 90% of the cage is occupied which gives the gas a solid volume ratio of about 160% (Riedel, et al 2010). The molecular arrangement of gas hydrates is shown in Figure 1.2.

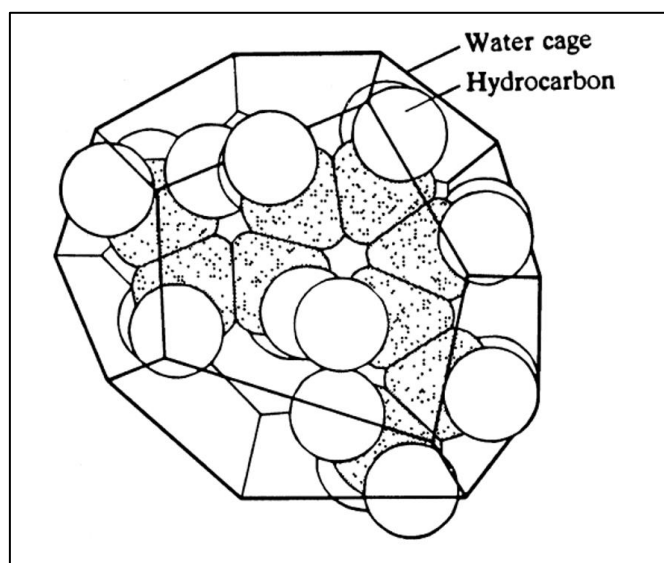


Figure 1.2 Molecular configuration of gas hydrates (Saleh, 2002)

Gas hydrate cages usually contain a single gas molecule and are composed of hydrogen-bonded water molecules. The structure of hydrates is called clathrates due to the entrapped (caged) nature of hydrogen molecules inside the crystal lattice of water molecules (Saleh, 2002), They form when water and gas combine at low temperature and high pressure in pipelines transporting natural gas, hydrates are lighter than water and as a result, they often reside at the point of interface between oil and water (Bellarby, 2009). The crystals have a honeycombed structure with small channels that allow gases to pass through, but with further accumulation, the flow in the pipe is entirely blocked. Gas hydrates also exist in subsea and ocean floors where conditions are ideal for formation, they trap large amounts of energy near subsea

seeps and around midline reservoirs (Saleh, 2002). Favorable conditions of natural gas hydrates exist in the sediment of Polar Regions and sediments covering 90% of the ocean floor. Most natural gas hydrates have more than 99% of their hydrocarbon as methane, low amounts of CO₂, and H₂S are present (Puall, et al 2001).

1.1 PROBLEM STATEMENT

Gas hydrates form inside the pipelines transporting natural gas at subsea levels and low-temperature regions. They form around the surfaces of the pipe and gradually accumulate until they constrict the flow or cause variations in flow pressure thereby causing damage to production equipment such as pumps, compressors, and separators at either end of the pipe. They can cause catastrophic rupture of the pipes and the failure of the gas transportation system. As a result, the safest and most cost-effective way of preventing gas hydrates is by predicting the temperature or pressure in which it will form for a natural gas mixture.

Many correlations have been developed to predict the conditions of hydrate formation, but the models require a complex computation, or their accuracy is limited to certain gas mixtures. Some of them can only be used in a limited range of temperature, pressure, or specific gravity and show a significant error at higher pressure. Also, some correlations are only useful when a comprehensive analysis of gas is accessible.

1.2 RESEARCH OBJECTIVES

The objective of this study is to develop a mathematical model for the prediction of gas hydrate formation temperature across a wide range of gas compositions, and pressure. Other objectives of this study, include:

- To determine the accuracy of the model by comparing the result of the model with experimental data as well as other reputable mathematical models.
- To estimate the hydrate formation temperature of natural gas compositions obtained from the Bakken Formation.

1.3 METHOD OVERVIEW

To fulfill the objectives of this study, the following steps were taken.

- The literature review presented previous research on hydrate formation and empirical correlations from reputable studies such as Hammerschmidt (1934), Motiee (1991), Towler and Mokhatab (2005), Chavoshi S. 2018, and Safamirzaei M. (2015). Also, their correlations were compiled in an Excel sheet.
- Gas composition data for The Bakken Formation were collected from various research and publications.
- A new model was developed by carrying Non-linear regression on 460 experimental data points using MATLAB.
- The new model was applied to the 460 data points in Excel and the calculated hydrate formation temperature was compared to experimental values as well as values calculated by other models.

- The model was evaluated by comparing its Average Relative Deviation and Average Absolute Deviation from reference experimental values, the deviation was compared to other models to assess its accuracy over other models.
- Gas composition data obtained from the Bakken Formation were applied to the model to calculate the hydrate formation temperature and compared with the Katz 1945 gravity chart.

1.4 THESIS OUTLINE

This thesis consists of six chapters. The outline of this thesis is as follows:

- Chapter 1 introduces gas hydrates and how they exist in the gas industry and nature. It also outlines the statement of the problem, research objectives, and methodology.
- Chapter 2 gives an in-depth overview of gas hydrates which includes how they form, mechanisms in play, causes and effects, the structures, methods of prevention, potential as an energy source in the future, and previous experiments and empirical correlations of hydrate formation temperature.
- Chapter 3 describes the methodology behind the development of the model, evaluation of the model, and data collection of gas compositions from the Bakken Formation.
- Chapter 4 presents the results of the calculated hydrate formation temperature of the proposed model and compares it to experimental data as well as values from other models. Also, the Average Relative Deviation and Average

Absolute Deviation of the model were compared to reference experimental values and that of other models.

- Chapter 5 discusses the result of the temperature-pressure curve of the new model as well as the Average Relative Deviation and Average Absolute Deviation from the reference experimental values.
- Chapter 6 presents the conclusions of the study including the recommendations for future work.

CHAPTER 2 LITERATURE

REVIEW

2.1 DISCOVERY OF GAS HYDRATES

In 1810, Humphrey Davy discovered a crystalline compound formed by chlorine and water, this later became the first known gas hydrate. An acetylene hydrate was later reported by Cailletet in 1878, he discovered that a sudden decrease in pressure helped in the formation of crystalline compounds. The first carbon disulfide and double hydrate hydrogen sulfide were reported by Schutzenberger and in 1882 Bordet discovered the double hydrate of phosphine and carbon dioxide. Double hydrates are compounds having a definite melting point and are not mixtures of a single hydrate, this is because the decomposition temperature of the single hydrate may be overall different from the decomposition temperature of a double hydrate. In 1897, Sully Thomas and De Forcrand found that carbon tetrachloride and acetylene forms double hydrates, they also discovered double hydrates of sulfur dioxide and ethylene in the following compounds: ethylene bromide, methyl Iodide, methylene chloride, methyl

bromide, methylene chloride, and methylene iodide (Hammerschmidt, 1934). The oil and gas industry began to take interest in gas hydrates in the 1930s when gas hydrate was found to cause blockages in pipelines. Hammerschmidt in 1934, discovered that natural gas hydrates were responsible for blocking gas pipelines.

2.2 GAS HYDRATE FORMATION

Gas hydrates are formed when water and gas (having lower molecular weight) combine at low temperature and generally high pressure (e.g. temperatures below 25 °C and pressures greater than 1.5MPa for natural gas hydrates) (Koh, et al 2001), precise conditions vary depending on the composition of the fluid. They exist onshore beneath the permafrost or offshore in shallow depths in the ocean (Sami, et al 2013). The gases that can form gas hydrates include methane (predominantly), propane, ethane, butane, chlorine, CO₂, nitrogen even oxygen can create hydrates. At high pressure, gas hydrates can exist at temperatures greater than the freezing point of water. The solid hydrate compounds form with the aforementioned gases at elevated pressures in the presence of water. At equilibrium conditions the hydrates cause a higher amount of water to be removed from the vapor phase than in the case of liquid water at the same pressure and temperature, this is due to the lower vapor pressure of the hydrates (Hammerschmidt, 1934).

2.2.1 Stages of Hydrate Formation

There are three main stages of hydrate formation: gas dissolution, hydrate nucleation, and lastly agglomeration as shown in Figure 2.1.

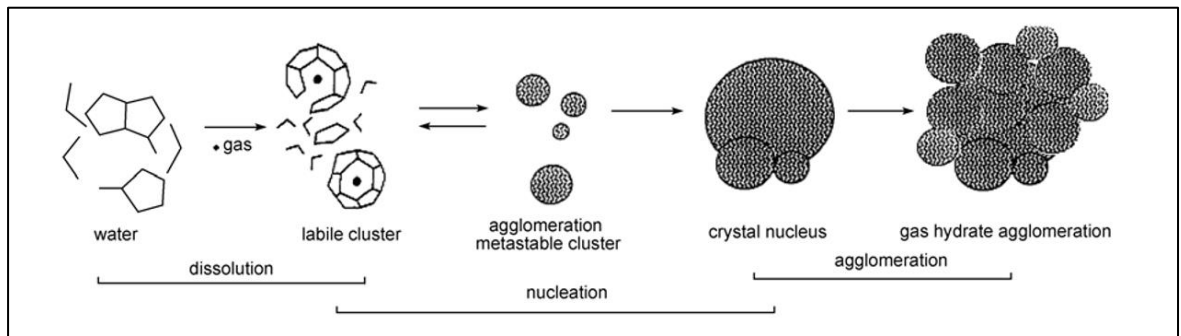


Figure 2.1 Stages of Hydrate Formation (Tang, et al 2010).

Tang, et al, 2010 found that under suitable conditions, water molecules form quasi-cavities from hydrogen bonds. Gas molecules are trapped inside these quasi-cavities when the gases dissolve, forming labile clusters. The clusters which are in quasi-equilibrium agglomerate and continue to attach until a critical radius is reached from which a stable hydrate nucleus is formed. It was found that during hydrate formation, ethane and methane occupy the smaller cavities while propane occupies large cavities. Also, when hydrates form there is a significant depletion of gas content, therefore gases in the fluid stream are consumed significantly (Tang, et al 2010).

2.2.1 Mechanisms of Hydrate Formation

The formation and deposition of hydrates in pipelines involve several processes from the formation of hydrate particles to the deposition and accumulation (growth) of the hydrate layer on the surface of the pipe. To forecast this dynamic deposition of hydrates in a pipeline, a model was developed by Zhang, et al 2019 in which the heat transfer and the hydrodynamic hydrate behavior in the pipeline were investigated. The model consisted mainly of two parts: Hydrate deposition and water condensation.

2.2.1.1 Heat Transfer

The temperature difference between the cold environment and the hot gas in the pipeline is large enough to cause heat transfer between gas pipelines in cold regions and the seafloor (Gu, et al 2019), as a result, water condenses when the temperature of the fluid falls below the dew point temperature. In cold regions and subsea levels, low temperature and high flow pressure in pipelines causes the molecules of natural gas to diffuse around the surface of water to form hydrates which adhere to the inner surface of the pipe, this leads to variations in fluid velocity, effective inner diameter and the pressure in the pipeline relative to position and time (Rao, et al 2013). The heat exchange between the external environment and fluid leads to a decrease in fluid temperature and this takes place along the direction of fluid flow as shown in Figure 2.2 (Zhang, et al 2019).

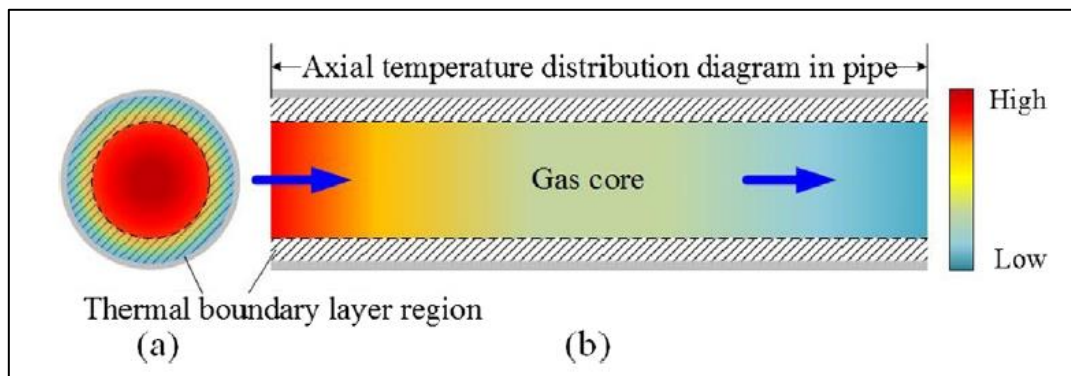


Figure 2.2 Fluid flow in a pipe showing its Thermal boundary layer region (Zhang, et al 2019)

2.2.1.2 Dissolved Water Condensate

There are two sources of liquid moisture in gas pipelines in cold regions and subsea levels: small droplets in the gas core and the liquid condensate film near the wall (Zhang, et al 2019). A heat boundary layer having a high variation in temperature

forms on the surface of the pipe as shown in Figure 2.2. This is due to a significant difference in temperature between the fluid and the environment during long-distance gas transportation in cold regions and subsea levels (Dorstewitz, et al 1994). When the fluid temperature falls below the dew point, the free water condenses and precipitate around the heat boundary layer, a liquid film forms on the pipe wall and there is a significant drop in temperature along the axis of the pipeline, as a result, water condenses on the surface of insoluble particles and small droplets form on the gas core (Gorbunov, et al 1998). Gas molecules continue to diffuse around the liquid condensate in the fluid system which consists of a thin liquid film near the pipe wall and small droplets in the gas core. A hydrate shell forms rapidly on the surface of liquid droplets when the pressure and temperature satisfy equilibrium conditions for hydrate formation, the shell grows inside the droplet to form hydrates. Also, the liquid film on the surface of the pipe wall crystallizes with gas molecules to form hydrates. These two types of hydrates deposit and accumulate on the inner wall of the pipeline to form larger hydrate layers. These processes continue until the thickness of the hydrate increases as shown in Figure 2.3.

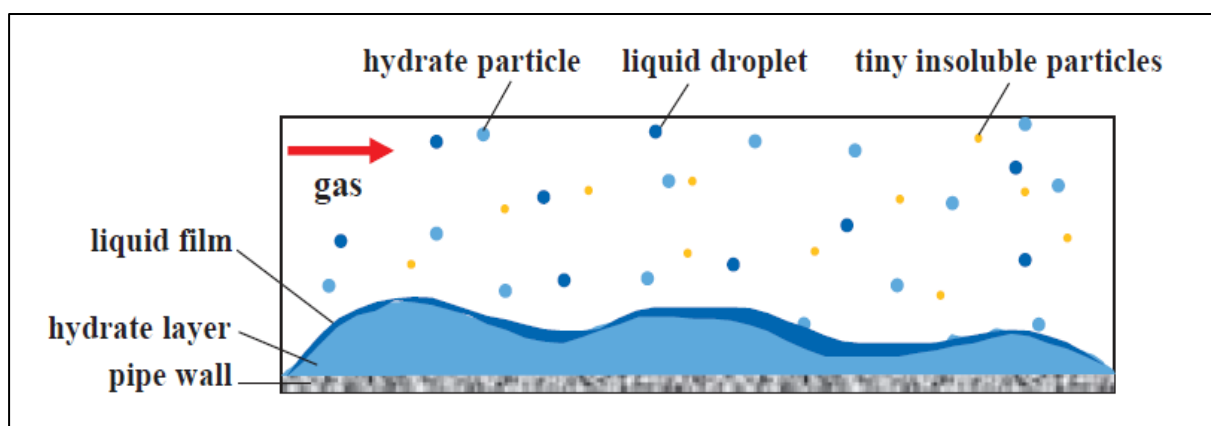


Figure 2.3 Schematic of hydrate formation and deposition mechanism in subsea and cold region long-distance gas transportation pipelines (Zhang, et al 2019).

The formed hydrate particles and the condensate droplets are transported downstream by the high-speed fluid due to their densities being in the same range.

2.2.1.3 Hydrate Layer Growth

The growth of hydrate layers is primarily due to the deposition process which combines the hydrates from scattered drops and thin condensate film (Zhang, et al 2019). The hydrate particles adhere to the pipeline wall due to strong adhesive forces between the wet pipe wall and the hydrates. New hydrate particles form from condensed water keeping the hydrate concentration in the pipe in constant dynamic equilibrium.

2.2.2 Causes of Hydrate Formation

The causes of gas hydrate are subdivided into primary and secondary causes.

2.2.2.1 Primary Causes of Hydrate Formation

The formation of gas hydrates in natural gas transportation pipelines depends primarily on the temperature, pressure, and composition of the gas-water vapor mixture (Hammerschmidt, 1934). After all primary conditions are in effect, hydrate formation is accelerated by pressure, high velocities of the gas stream, and pulsations. As shown by the melting point diagram in Figure 2.4, high pressure and low temperature are favorable to the formation of hydrates. On a practical basis, water vapor is the only component that can be controlled regarding the composition of the gas. Although removing moisture from the gas will eliminate the possibility of any hydrate forming, it is not necessary for the gas to be free of water vapor since hydrates only form when the gas reaches the dew point. Also, if the partial pressure of water vapor in the gas is less than the vapor pressure, the gas hydrate will lose water

and dissociate. The driving force behind phase change from gas to solid hydrates is thermal sub-cooling. In this case, the phase change is caused by the temperature difference between the bulk temperature and the hydrate dissociation temperature (Hammerschmidt, 1934).

The conditions of temperature and pressure are represented in disassociation curves where hydrates separate into gas and water as shown in Figure 2.4. The point of hydrate formation lies within the curve, this means that hydrates are certain to form within the curve but will not form immediately if the temperature/pressure point lies outside the disassociation curve (Saleh, 2002). The risk of hydrate agglomeration increases further inside the curve although there is a time delay with an undefined duration. The curves are generated based on experimental values or numerical predictions using the Katz gas gravity chart (Zhang, et al 2019).

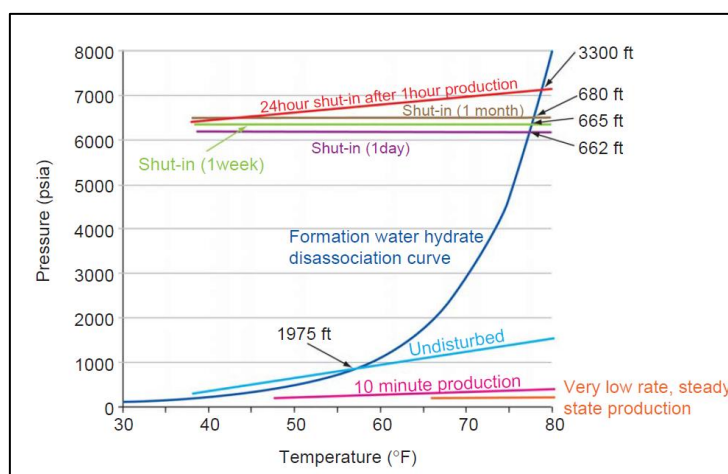


Figure 2.4 Disassociation curves showing the conditions of temperature and pressure (Zhang, et al 2019).

Figure 2.5 shows Katz’s 1945 gas gravity chart.

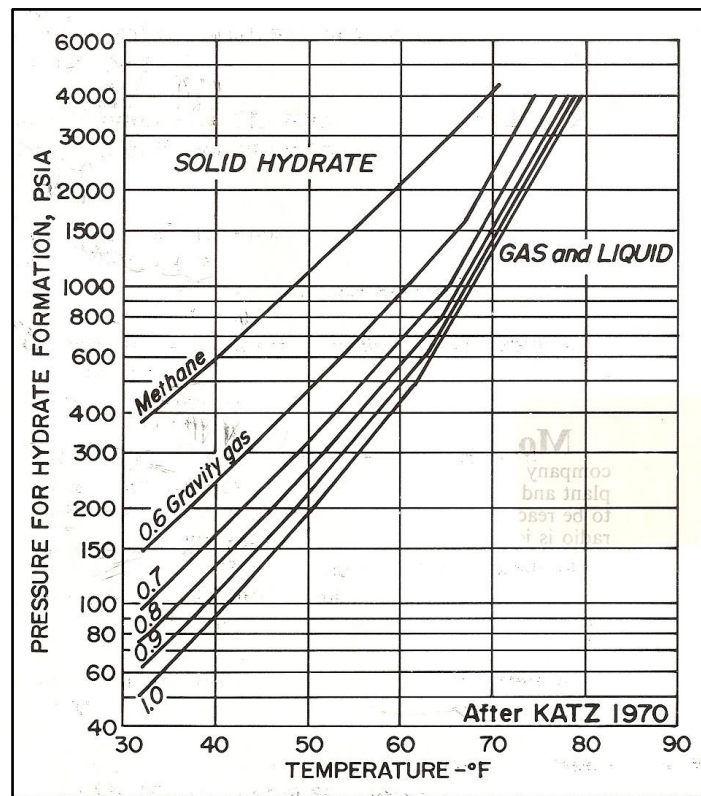


Figure 2.5 Katz gas gravity chat (Bahadori and Vuthaluru, 2009)

2.2.2.2 Secondary Causes of Hydrate Formation

As previously stated, a definite pressure, temperature, and gas-water composition are necessary for gas hydrates to form. However, it is not certain that hydrates will crystallize even after these conditions are met (Hammerschmidt, 1934). Certain secondary factors influence the formation of hydrates, these are:

1. Gas Stream velocity
2. Arrangement of the molecule of crystals
3. Water saturation (water vapor)
4. Porosity

2.2.2.2.1 Gas stream velocity

High gas velocity, pressure pulsation of the gas (due to compressors), or introducing small amounts of hydrate crystal can hasten the formation of hydrates. Under operating conditions, high gas stream velocity in the pipeline creates almost ideal conditions for the formation of hydrates provided the proper conditions of temperature, pressure, and water composition are established (Eucken. 1925).

2.2.2.2.2 Arrangement of crystal molecules

The general behavior of crystal formation described by Eucken (1925) explains the secondary causes of hydrate formation, it states that the formation of a crystal requires mostly a definite adjustment or generally a certain arrangement of molecules, this may not always be established at once and the lack of which is characterized by the liquid phase. Ideally, a certain amount of time elapses before the required number of molecules align into the correct position either by coincidence or accident. It is only after a nucleus or elementary crystal is formed that crystallization proceeds smoothly, this exerts a certain directional force on the neighboring liquid molecules and causes them to merge. As a result, hydrate formation is promoted by forces which tend to stir or mix, because the probability of the molecules aligning into the correct position is increased by those forces. Also, high-pressure pulsations and velocity impact a mixing action on the droplets of condensed moisture (Hammerschmidt, 1934).

2.2.2.2.3 Water Saturation (Water Vapor)

During the process of transporting natural gas, water is removed from the gas before transporting through pipelines, this is called separation and dehydration. However, these processes only remove the free water in the gas, therefore the pipeline system is a water-saturated gas transportation system. The phenomenon of hydrate deposition in

pipelines was observed by (Lingelem, et al 1994), they suggested that for a water-saturated gas system the mechanism is similar in a continuous gas system, having small proportions of free water.

Gorbunov, et al 1998 verified that condensation of water vapor takes place on the surface of small-size insoluble particles, it also determined that vapor heterogeneous-phase nucleation forms alone on the surface of suspended particles with soluble sites. Gas molecules constantly diffuse in the gas core around the surface of the condensate liquid drops.

2.2.2.2.4 Porosity

Natural gas produced from the reservoir stratum generally contains small-sized insoluble particles, as a result, in an environment where small-sized insoluble particles are suspended, supersaturated vapor molecules collide constantly and undergo initial heterogeneous phase nucleation on particle surfaces. Nicholas, et al. 2008 found in a study that porosity influenced the thickness of hydrate deposition in the pipelines significantly. Free water is the main constraint of hydrate formation/deposition when conditions of pressure and temperature reach hydrate phase equilibrium in a water-saturated gas system. The study proposed that the condensate of free vapor on the cold walls of the pipe formed the free water in a water-saturated system.

2.2.2.3 Other Factors Affecting Hydrate Formation

Other factors that influence hydrate formation include (Zhang, et al 2019)

1. Adhesion between hydrate particles and the pipe wall,
2. Spatial distribution characteristic of the gas-liquid phase, and

3. Fluid flow rate.

Particle surface energy, the thickness of the dielectric layer, and Hamaker number affect the nature of deposition of the particles (Chein, et al 2005). A parametric analysis was carried out by (Chaudhari, et al 2018) to investigate the effect of mixture velocity, liquid loading, and interfacial tension on hydrate formation with the use of hydrate risk correlation which uses the steady-state and transient dynamic multiphase flow simulation on a long subsea tieback. The study deduced that the flow assurance risk increases with an increase in mixture velocity and liquid loading, however it decreases with a decrease in the interfacial tension.

2.2.3 Gas Hydrates Structures

Gas hydrates form pentagonal cubic structures in their crystal lattice. The hydrocarbon molecules (in this case methane) are lodged in the center of the cubes and bounded (caged) by water molecules. There are three main structures of gas hydrates that have been found Structure I (SI), Structure II (SII), and Structure H (SH) having a cubic (isometric) lattice (Sloan, et al 2008). All the structures contain large and small cavities but only molecules with the appropriate size and geometry enter the cavities (Dorstewitz, et al 1994). Figure 2.6 shows the physical geometry and lattice structure of methane hydrates, it shows that the gas molecule is located at the center of the water cavity.

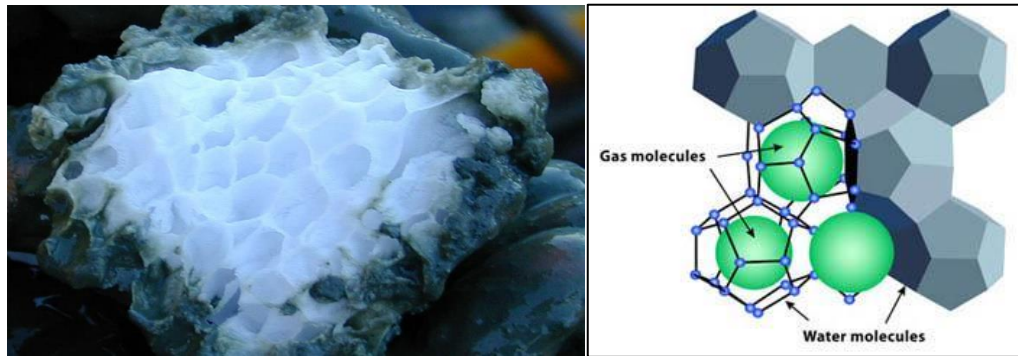


Figure 2.6 Physical geometry and Lattice structure of Gas hydrates (Studentenergy.org)

Methane and natural gas typically form hydrate Structure (I) but forms Structure II or H if higher hydrocarbons are present in the gas mixture as is the case with thermogenic gas components (Riedel, et al 2010). Figure 2.7 shows the lattice structure of the gas hydrate.

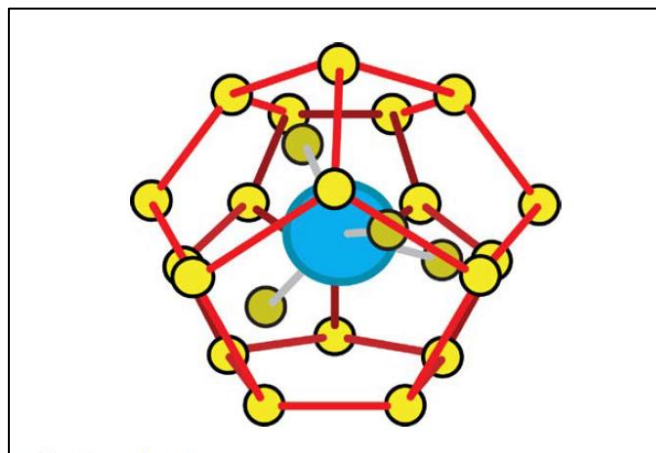


Figure 2.7 Lattice structure of gas hydrates (Bellarby, 2009)

2.2.3.1 The Structure I (SI)

Structure (I) has two main structures of cavities: a small pentagonal dodecahedral cavity which consists of 12 pentagonal rings of water (20 molecules of water) and a large tetrakaidekahedral cavity consisting of 2 hexagonal and 12 pentagonal rings of water (24 molecules of water) (Koh, et al 2002). The arrangement of the cubic

cavities is an in-body-centered packing and the cavities are large enough to contain methane, ethane, and gases having a similar range of molecular diameter such as hydrogen sulfide and carbon dioxide hydrates (Sami, et al 2013).

2.2.3.2 Structure II (SII)

Structure (II) also has two main structures: a pentagonal dodecahedral cavity and a large hexakaidecahedral cavity which consists of 12 pentagonal and 4 hexagonal rings of water (28 molecules of water). They are packed like diamonds in an octagonal shape, this leads to some cavities being large enough to contain not only ethane and methane but also larger gas molecules such as isobutane and propane.

2.2.3.3 Structure H

This structure requires both small molecules such as methane and larger molecules of typical gas condensates or oil fractions. Figure 2.8 shows the cell units and structures of hydrates (Sami, et al 2013).

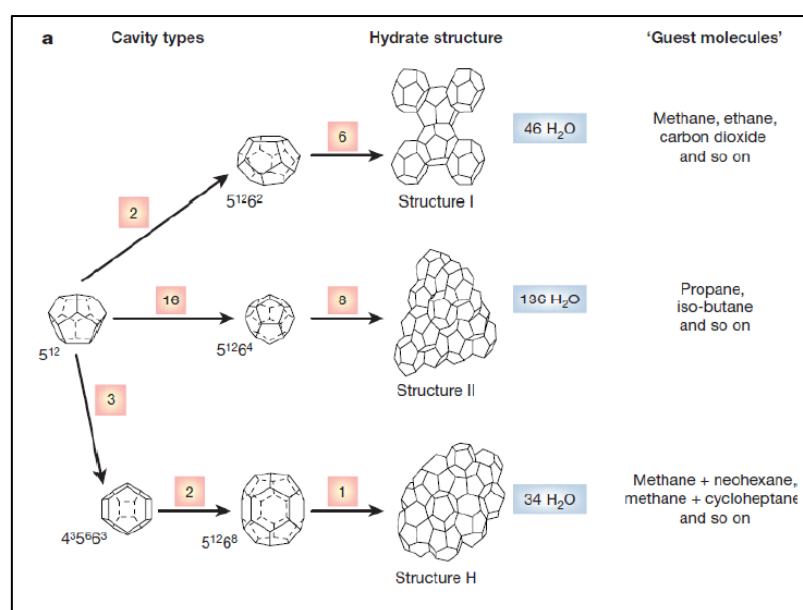


Figure 2.8 Cell unit structures of gas hydrate structure I, II, and H (Sami, et al 2013).

2.3 EFFECTS OF GAS HYDRATES

Gas hydrates have a wide range of effects on the oil and gas industry from upstream drilling and production to the midstream transportation, compression, and separation systems of natural gas. Gas hydrates are responsible for plugging in natural gas pipelines, they deposit on walls of the pipeline thereby reducing the effective flow diameter of the pipe. This causes fluctuations in the pressure of the fluid stream by causing an increase in pressure in areas where there are hydrate deposits and a decrease where there are no deposits (Sami, et al 2013).

2.3.1 Gas Hydrate Occurrence During Drilling in Offshore

Regions

The complexity and challenges of offshore drilling increases with increase in water depth. One main challenge is the formation of gas hydrates, during deepwater drilling it is likely to encounter shallow sediments containing natural gas, if encountered, the gas could enter the drilling fluid causing gas hydrate formation under high pressure and low temperature (Poberezhny, et al 2019). Also, gas hydrates could easily form when drilling mud circulation is stopped and gas enter into the fluid, in cases where the drilling mud is not treated with hydrate inhibitors, this can lead to unexpected gas kick during drilling which can block the annular clearance, pipe or other equipment such as the blowout preventer (BOP) which may result in catastrophic failure of the system in some situations. (Xiaolan, et al 2011).

Solid hydrates can plug well kill-lines and chokes causing problems in well control.

Hydrates can stop the circulation of drilling fluid and prevent the drill string from

moving if it forms in the annulus between the drilling string and casing, this can seriously affect drilling operations. Hydrates can also form in the riser, plugging the flow of drilling fluid. Figure 2.9 shows a summary of the major problems caused by gas hydrates while drilling through gas hydrate formations (Sami, et al 2013).

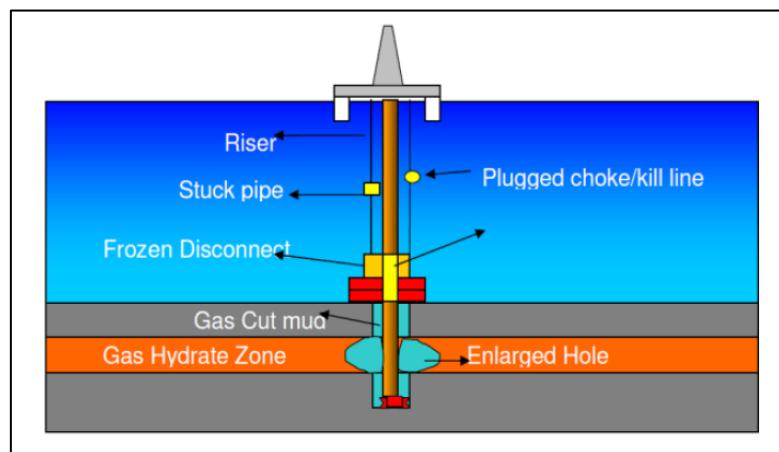


Figure 2.9 Pictorial summary of the major problems encountered during drilling operations through a hydrate formation (Sami, et al 2013).

2.3.2 Problems with Flow Assurance

Pipeline transportation is a common means of transporting oil and gas from the wellhead to production sites. Gas hydrate particles form when the pressure and temperature of the fluid in the pipeline falls within the hydrate zone in the phase diagram, these hydrate particles could eventually plug the pipeline. When gas hydrates plug transmission pipelines, operations become uneconomical, production stops for weeks or sometimes months in large extended pipelines. The propagation of hydrates tends to develop a plug that gradually separates the pipe into two pressure zones: a high-pressure zone between the high-pressure gas source and the plug, and a low-pressure zone between the plug and the gas recovery section. Figure 2.10 shows a typical hydrate plug in a gas transportation pipeline (Sami, et al 2013).



Figure 2.10 Gas hydrate plugging of gas transportation pipeline (Polartrec.com)

A pipe leak and explosion can occur due to the growth in pressure. The higher pressure can destroy the pipe when there is a large difference in pressure between the upstream and downstream sections. Both failures can put personnel in danger and damage equipment.

2.3.3 Corrosion in Gas Pipelines

Gas hydrates cause pitting corrosion of pipelines that are in acidic or neutral environments. Acidic gases such as CO_2 and H_2S , when dissolved in water accelerate the internal corrosion of pipelines as the gas components of hydrate formation (Poberezhny, et al. 2017). Specimen with gas hydrates formed on the inner surface show an increased rate of localized and general corrosion on their surfaces as shown in Figure 2.11.



Figure 2.11 General view of corrosion damage of the specimen after exposure to gas hydrates (Poberezhny, et al 2019).

(Poberezhny, et al 2019) investigated the influence of gas hydrates on the fatigue test of a St 20 steel pipe where tests were performed at pure bending in air. Specimens were tested after exposure to gas hydrate and without any pretreatment. The curve in Figure 2.12. shows a three-stage kinetics growth of fatigue crack, also a slight higher-level cyclic deformation on the fatigue growth curve was observed in the pipe after exposure to gas hydrates, it may be attributed to corrosion damage of the pipe surface.

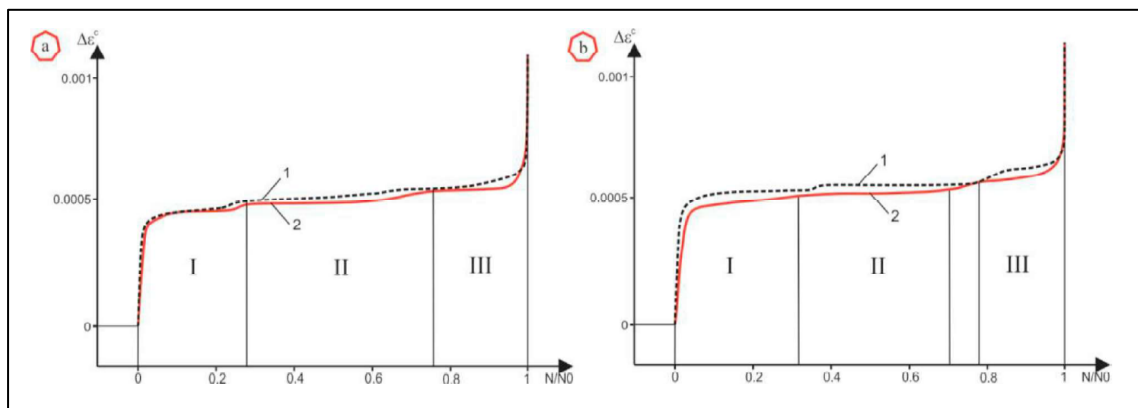


Figure 2.12 Fatigue crack growth curves for steel specimen (a) without exposure, (b) after exposure to hydrates (Poberezhny, et al 2019)

The rate of growth of fatigue crack was higher by 5-7% for the pipe specimen after exposure to hydrates compared to the specimen without pretreatment, it was attributed to the increase in surface damage due to the formation of hydrates. The fatigue test shows that the duration of the low-frequency fatigue stage was influenced by hydrates causing a shortening of the stage (III) shown in Figure 2.12, this corresponds to the serviceability of the pipeline. It also shows an increase in the deformation for steel specimen exposed to hydrates. Similar behaviors were noted for materials of sea pipelines and drill pipes (Poberezhny, et al 2019).

2.4 CURRENT HYDRATE PREVENTION

METHODS

The most effective strategy for managing hydrates in an oil production system is staying outside the temperature-pressure envelope where hydrates are stable.

Examples of this are insulating the flow path or adding a heat source which can keep the temperature of the fluid sufficiently above the predicted hydrate formation temperature during steady-state operation, this will also allow for adequate time to reduce or depressurize the flow. The purpose of reducing the pressure is to lower the temperature of hydrate formation to below the ambient temperature to allow for an extended shut-in time (Saleh, 2002). There are four main methods of combating gas hydrate plugs to ensure a continuous flow.

1. Hydraulic method
2. Thermal method
3. Mechanical method
4. Chemical method

2.4.1 Hydraulic Method

The hydraulic removal method involves depressurizing the fluid stream thereby dissociating the hydrate plug, due to the porous structure of the plugs in the gas pipelines, this method appears promising. However, it is only suitable for gaseous hydrocarbon as depressurization in liquid causes vaporization. Lowering the flow pressure does not dissolve a hydrate plug immediately, this is because the

disassociation of hydrates is highly endothermic (absorbs heat from the surroundings) which delays the break-up of the plug (Saleh, 2002).

2.4.2 Thermal Method

The thermal method involves an in-situ delivery of high temperature (heat) flow towards the hydrate plug area through the pipeline wall to raise the temperature of the system above the hydrate formation temperature. This method can be achieved for external pipelines but not suitable for subsea pipelines.

2.4.3 Mechanical Method

The mechanical method involves pigging to clean the pipeline of hydrate deposits. This method involves moving a large spherical or cylindrical disc made of a flexible material, having an outside diameter almost equal to the inside diameter of the pipeline. An example of these flexible materials includes neoprene rubber. The disadvantage of this method is that shutting down production to pig the pipeline is expensive and causes downtime (Sorheim, 2005). Figure 2.13 shows an example of a pipeline pigging operation.



Figure 2.13 Pipeline pigging (Picchemicals, 2019).

2.4.4 Chemical Method

The technology adopted by the oil and gas industry for the prevention of gas hydrates in pipeline transportation involves the introduction of a thermodynamic inhibitor such as methanol into the natural gas fluid flow. However, as oil and gas exploration moves into extreme environments (such as deep-sea, offshore exploration, and production) the temperature and pressure conditions become more severe in the field, as such the concentration of inhibitors required for the prevention of hydrate formation increases significantly, most times to excess levels. Hence other technologies have been developed in the form of low-dosage chemicals such as kinetic or anti-agglomerate inhibitors. These compounds function by retarding the hydrate formation time to longer periods than the resident time of the gas inside the hydrate-prone area of the pipeline. Compared to thermodynamic inhibitors these low dosage inhibitors offer significant environmental and economic advantages. The main aim of low dosage inhibitors is to interfere with the mechanism of hydrate formation (Koh, et al 2002).

The chemical process involves the use of two types of additives thermodynamic inhibitors (THI) and Low dosage hydrate inhibitors (LDHI), they are further subdivided into the following (Sami, et al 2013)

1. Thermodynamic Inhibitors (THI)
 - i. Alcohols (e.g. Methanol)
 - ii. Glycols
2. Low-dosage hydrate inhibitors (LDHI)

- i. Surfactants
- ii. Kinetic Inhibitors
- iii. Anti-agglomerates

2.4.4.1 Thermodynamic Inhibitors (THI)

Thermodynamic inhibitors reduce the water movement and reaction by shifting the hydrate phase boundary to a higher pressure and lower temperature, this can prevent hydrate formation effectively (Sami, et al 2013). Considering that hydrates are similar to ice, chemical deicers that remove or prevent ice from accumulating can also work for gas hydrates.

2.4.4.2 Alcohols (Methanol)

Alcohols have proven effective in combating ice formation and gas hydrate control. They contain the hydroxyl (OH) group which ensures solubility in water. The simplest form of alcohol is methanol (CH₃OH) and it is the most available/widely used in the gas industry. The addition of sufficient amounts of these compounds creates a condition whereby the temperature and/or pressure required for hydrate formation needs to be lower for it to form (Koh, et al 2002). Methanol has a relatively low molecular weight, this allows it to penetrate the hydrate cavities and layers to effectively dissolve the hydrates.

However, due to this low molecular weight, methanol is lighter than water and as a result, may prove ineffective injecting it from the wellhead downhole into the hydrate plug submerged with oil. It is effective when dealing with very light oil.

Thermodynamic inhibitors such as methanol are used to prevent hydrate formation after a shut-down where the range of time is much longer.

Plots showing the effectiveness of alcohols in hydrate inhibition are used to formulate the required dosage rates for hydrate inhibition of completion and intervention fluids, keeping in mind that less methanol or glycol should be used when brine is used in drilling fluid due to its inherent inhibition characteristics. Figure 2.14 is a curve showing the effectiveness of various alcohols in hydrate inhibition (Bellarby, 2009).

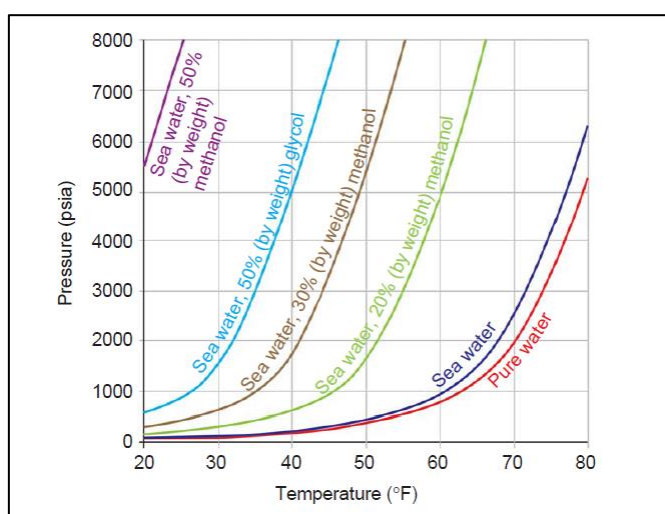


Figure 2.14 Hydrate inhibition with methanol and glycol (Bellarby, 2009)

In some fields, it is common for higher concentrations of methanol (up to 60 mass %) to be used and may also require the addition of glycol. These large volumes of methanol and glycol raise concerns about the environment, higher cost, and logistics. Low dosage hydrate inhibitors in contrast are new categories of chemical additives that can be effective at low volumes (mass %) for the same application. (Sami, et al 2013).

2.4.4.3 Glycols

Glycols are other types of hydrate inhibitors used in the industry. Two frequently used glycols are Triethylene glycol (TEG) ($\text{HO-C}_2\text{H}_4\text{-O-C}_2\text{H}_4\text{-O-C}_2\text{H}_4\text{-OH}$) and Monoethylene glycol (MEG) ($\text{HO-C}_2\text{H}_4\text{-OH}$) also diethylene glycol (DEG) is used

occasionally (Elhady, 2005). MEG is denser and more viscous than water, it is flammable (but its flash point is higher than methanol). TEG is slightly denser and more viscous than methanol, this is due to the higher molecular weight of TEG. The greater viscosity and density of the inhibitors compared to methanol provide a useful means of removing hydrates when the glycol is injected above the hydrate plug if the glycol can migrate down the tubing and stay atop the plug before it circulates into water. Compared to methanol, glycols are easier to extract in the production system, for this reason, they are frequently used in wet gas pipelines (Brustad, et al 2005).

2.4.4.4 Low-Dosage Hydrate Inhibitors (LDHI)

LDHI can prevent the formation of gas hydrates during the nucleation and agglomeration stages. Examples of commercial low-dosage hydrate inhibitors are Inhibex 100, Inhibex 501, PVP, and GHI1 (Bellarby, 2009). Ping et al 2010 found that neither GHI1 nor PVP was able to inhibit gas hydrate nucleation, but GHI1 has a greater ability to inhibit hydrates than PVP. This higher inhibition ability is caused by diethylene glycol monobutyl-ether. The categories of low-dosage hydrate inhibitors are:

1. Surfactants
2. Kinetic Inhibitors
3. Anti-agglomerates

2.4.4.4.1 Surfactants

These chemicals disperse hydrate crystals as they form, preventing the accumulation of the hydrates in a particular location in the pipeline. A typical example of such a chemical is Lecithin which is environmentally friendly and is used as a food

antioxidant. Lecithin is sometimes used as a drilling fluid additive to stop blowout preventers from freezing when gas influx occurs in deep-water. Pakulski, (2007) reported an increase in hydrate formation when natural or introduced surfactant (such as anti-agglomerates) are present.

2.4.4.4.2 Kinetic Inhibitors

Kinetic inhibitors are polymers of various types such as polyvinyl caprolactam, they work by preventing or reducing the nucleation and subsequent crystal growth of the hydrates, thereby delaying the hydrate formation (Sami, et al 2013). A one mass % concentration is sufficient to control the formation process of gas hydrates until the gas is transported to its destination where the thermodynamic conditions for hydrate formation do not exist. A major disadvantage to these inhibitors is that once nucleation occurs, these inhibitors are unable to prevent further crystallization of hydrates (Bellarby, 2009).

2.4.4.4.3 Anti-agglomerates

These inhibitors work by making the surface of the hydrate hydrophobic (avoid or repel when it makes contact with water) enabling it to be dispersed in the oil phase. Anti-agglomerates do not prevent hydrates from forming but hinders already formed gas hydrates from agglomerating into lumps that cause plugging, an example of an anti-agglomerate is quaternary ammonium salt (QUATS) which is the main active component used in corrosion inhibitors (Sami, et al 2013).

2.5 GAS HYDRATES AS A POSSIBLE SOURCE OF ENERGY

Gas hydrate is commonly known as the new clean energy (Zhang, et al 2019).

Methane gas hydrate occurs in large deposits in the form of sedimentary rocks and solid sediments within 2000m of the earth's crust in deep-water and permafrost regions (Giavarini, et al 2011). The volume of methane hydrates under the same standard temperature and pressure condition contains 164 times more methane than one volume of gaseous methane. Potential deposits of gas in hydrates that are distributed on land and offshore are more than $1.5 \times 10^{16} \text{m}^3$.

Methane gas can be extracted from the gas hydrates which can then be used as natural gas. Gas extraction is typically done by depressurization, thermal, and/or use of chemical inhibitors. During thermal stimulation, gas is released from the gas hydrate deposits by warming the formation through the direct heating of the formation or injection of heated fluid. Depressurization involves lowering the flow pressure of the gas hydrates, it is more preferred and economical than thermal stimulation, it does not require expending large energy and can be used to dissociate larger volumes of gas hydrates rapidly. Its main disadvantage is that it requires hydrate deposits with high porosity, also during the transportation phase the recovered gas and water may recrystallize into gas hydrates in the transportation pipelines leading to pipe plugging.

Hydrates occur in most arctic sedimentary regions and deep waters, this makes gas hydrate a widespread potential resource, they are naturally stable under cold temperature (deep-water or arctic) or high pressure (deep-water). The amount of gas

(mostly methane) entrapped in natural hydrates is about 50 times greater than natural gas from conventional resources (Milkov and Sassen, 2002). A region of hydrate stability exists which is determined by overlapping the geothermal temperature gradient and hydrostatic pressure with the hydrate disassociation curve for seawater, Figure 2.15 shows a typical hydrate stability zone for three varying water depths. The curve indicates that the hydrate stability zone varies with the geothermal gradient and seabed temperature. The lower portion of the stability zone indicates the point where natural gas caged in hydrates (solid) transition to natural gas (vapor). This is often identified as a bottom simulating reflector (BSR) on seismic charts though it is not a reflection of multiple seabeds.

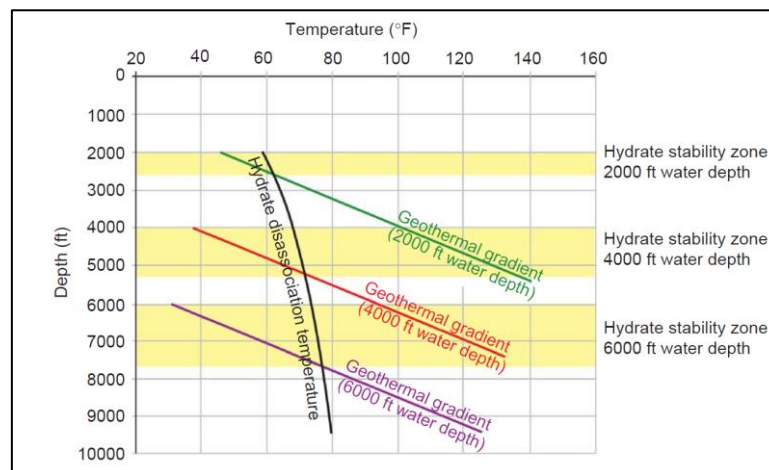


Figure 2.15 Natural hydrates stability (Bellarby, 2009).

In some circumstances, hydrates can form the cap for gas reservoirs. In a situation where the cap is penetrated by production wells due to the cap being thin, the melting of hydrates around the wellbore could breach the cap during the production of the warmer underlying gas (Bellarby, 2009).

2.6 RESEARCH AND EXPERIMENTS ON GAS HYDRATES

The Following studies are notable research and experiments on the formation and effects of gas hydrate.

2.6.1 Prediction of Hydrate Deposition in Pipelines to Improve Gas Transportation Efficiency and Safety By (Zhang, Et Al 2019)

The study simulated a 48 km long subsea pipeline transporting natural gas to predict the hydrate phase equilibrium region in the gas pipeline. The result shown in Figure 2.16 indicates that the region of hydrate phase equilibrium in the pipeline is around 220-48000m away from the pipe inlet. This suggests that hydrate deposition takes place at a distance of 220m away from the inlet.

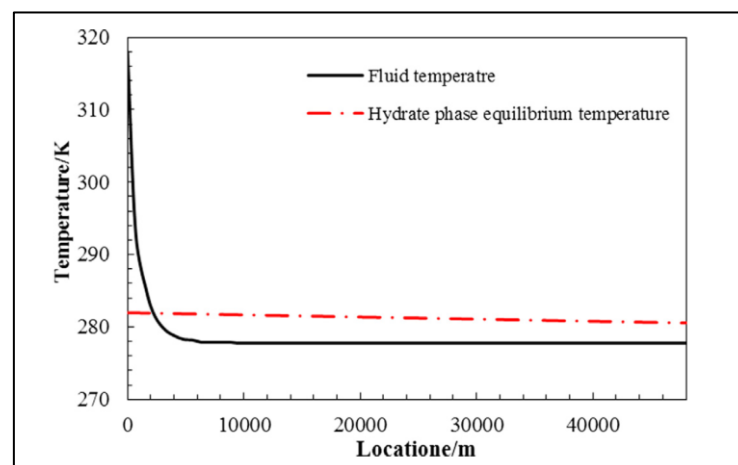


Figure 2.16 Prediction of hydrate phase equilibrium region in the subsea gas pipeline (Zhang, et al 2019).

Hence the safety and efficiency of the gas transportation system are significantly reduced by the formation and deposition of hydrates. It can be seen from the curve that the outlet pressure of the pipeline decreased significantly under the effect of hydrate deposition which decreased the effective inner diameter of the pipe and increased the flow velocity, this leads to an increase in pressure in the pipeline. Figure 2.17 shows a comparison between the effective radius of the pipeline and the proposed radius obtained from a test model, the results show that with an increase in hydrate deposition the effective radius of the pipe reduces. Also, there is a region of high-risk of hydrate deposition in the pipeline in which the two models showed similar results within the range of (5-20m) from the pipe inlet. Fluctuations in the outlet pressure of the pipeline due to hydrate deposition can damage pipeline joints and pumps, inducing severe accidents in production (Zhang, et al 2019).

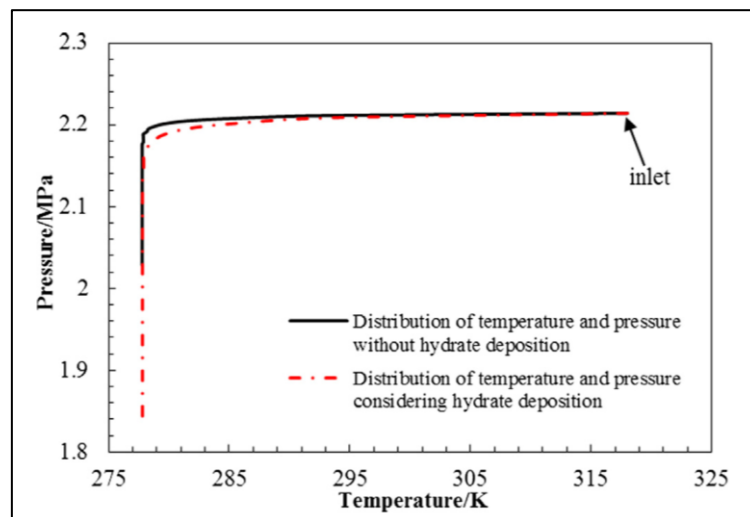


Figure 2.17 Comparison between temperature and pressure distribution in the pipeline (Zhang, et al 2019)

2.6.2 Hydrate Formation in Pipelines By (Dorstewitz And Mewes, 1995)

Hydrate formation was investigated experimentally in a horizontal pipe flow of gas and water by Dorstewitz, 1995. The pressure, temperature, and volumetric flow rate of each fluid were measured. The tubing made of glass had an inner diameter of 15mm. The formation of plugs starts on the pipe wall because the heat transfer is directed radially outwards. The onset of hydrate formation on the pipe wall is located at the interface between water and gas. Flow regime changed from the stratified flow of water and gas to intermittent flow pattern. Water was conveyed into the upper part of the pipe. The whole perimeter of the wall was covered by hydrates and a closed hydrate layer was achieved. The hydrate layer grew radially towards the center of the pipe. In certain volumetric flow rates and heat fluxes the hydrate, layer growth lead to plugging of the pipeline as shown in Figure 2.18.

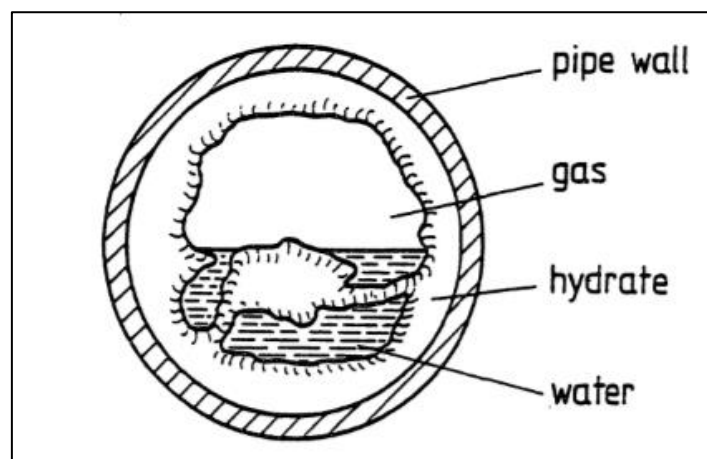


Figure 2.18 Hydrate formation pattern in a 15mm test pipe (Dorstewitz and Mewes, 1995).

Due to the void distribution, the value of the related pressure loss is low. The onset of hydrate formation occurred after 50 minutes. Only small hydrate particles were observed on the pipe wall. The particles were dragged along with the flow and the

hydrates decompose in the bulk flow. Due to further deposition of hydrates, the whole perimeter of the pipe was covered by hydrates after 59 minutes and a solid hydrate layer developed.

2.6.3 Empirical Correlations

Other methods have been developed to predict the hydrate formation conditions of gas mixtures without the need for expensive experiments. The methods of predicting hydrate formation include empirical correlations, graphical calculations, software packages, and thermodynamic models. In this study, empirical correlations have been reviewed. The empirical correlations evaluated included: Hammerschmidt (1934), Motiee (1991), Towler and Mokhatab (2005), Safamirzaei M. (2015), and Chavoshi S. 2018. All the equations calculate the hydrate formation temperature (T-explicit) for gas mixtures. Most correlations used to estimate the hydrate formation conditions calculate for T-explicit because the pressure is usually known from the process and/or flow transfer requirements, and hydrate formation temperature is the parameter required.

In 1934, Hammerschmidt proposed a correlation for gas hydrate formation, shown in Equation 1 (Hammerschmidt, 1934).

$$T = 8.9P^{0.285} \dots \dots \dots (1)$$

Where temperature T is in (°F) and pressure P is in (Psi). This equation, although simple does not consider the effect of gas specific gravity on hydrate formation.

Motiee (1991) developed Equation (2). Where temperature T is in (°F) and pressure P is in (Psi).

$$T = -238.24469 + 78.99667 \log P - 5.352544(\log P)^2 + 349.473877\gamma - 150.854675\gamma^2 - 27.604065\gamma \log P \dots \dots \dots (2)$$

The equation is widely used in the gas industry due to its accuracy in predicting the hydrate formation temperature for various natural gas mixtures but has 6 adjustable parameters which are relatively high and can lead to errors when carrying out quick hand calculations.

In 2005, Towler and Mokhatab proposed a relatively simple equation for estimating hydrate temperatures as a function of the pressure and the gas gravity shown in Equation 3.

$$T = 13.47 \ln P + 34.27 \ln \gamma - 1.675 \ln \gamma \ln P - 20.35 \dots \dots \dots (3)$$

Where temperature T is in (°F) and pressure P is in (Psi). (Towler & Mokhatab, 2005)

In 2015, Safamirzaei proposed a T-explicit correlation for specific gravity $0.55 \leq \gamma \leq 1$ shown in Equation 4. (Safamirzaei M.,2015).

$$T = A\gamma^B (\ln P)^C \dots \dots \dots (4)$$

Where:

$$A = 194.681789$$

$$B = 0.044232$$

$$C = 0.189829$$

Where P is in (kPa) and T is in (K). The study and its subsequent equation are not peer-reviewed, also the method in which the model was derived was not stated in the study.

Chavoshi S. 2018 developed a hydrate formation equation shown in Equation 5 by fitting a polynomial function to 100 experimental data points using the curve fitting tool in MATLAB software.

$$T = 242 \times \gamma^{0.02} \times P^{0.021} \dots \dots \dots (5)$$

where P is the pressure in (kPa), γ is specific gravity, and T is the temperature in (K).

Additionally, there are other equations such as Kobayashi (Kobayashi, Song, Sloan, & Bradley, 1987) and Ameripour and Barrufet (Ameripour & Barrufet, 2009). The main advantage of these correlations is their simplicity and portability. In most cases the required input data are accessible and can be calculated with a simple calculator, also the results agree with the experimental data and sometimes better than results obtained from commercial software. Despite the advantages, these correlations have limitations, most of them can only be used in a limited range of temperature, pressure, and specific gravity and they show a significant error at higher pressure. Some correlations are only accurate for certain gas mixtures such as sweet gas or with pure formers, also some correlations are only useful when a comprehensive analysis of gas is accessible. Some models use the artificial neural network (ANN) such as (Elgibaly & Elkamel, 1998), (Zahedi, Karami, & Yaghoobi, 2009), and (Khomehchi, Shamohammadi, & Yousefi, 2013). These models are often complicated and are not suitable for hand calculations (Safamirzaei M., 2015).

CHAPTER 3 METHODOLOGY

3.1 DATA COLLECTION

In this study, 460 experimental data points were collected from 15 different studies on hydrate formation, the studies are listed in Appendix A. The units of temperature for the data points were converted to Kelvin (K) and the pressure converted to Kilopascals (kPa). Gas compositions of natural gas from the Bakken Formation were collected from various wellheads of oil and gas producing wells by the Energy and Environmental Research Center (EERC) and presented in the study by (Wocken et al, 2012) and also (Eenews, 2013) as shown in Table 3.1.

Table 3.1 Gas Composition Data from Wellheads in the Bakken Formation in North Dakota (Provided to the EERC by several North Dakota operators)

Component	Gas Composition (Mol%)							
	A	B	C	D	E	F	G	H
H2O (Water)	0.02	0.29	0	0	0	0	0	0
N2 (Nitrogen)	5.21	7.10	1.72	0.86	1.435	1.715	9.9	2.09
CO2 (Carbon dioxide)	0.57	0.51	1.72	0.86	1.435	1.715	9.9	2.09
H2S (Hydrogen sulfide)	0.01	2.00	0.19	0.18	0.08	0.05	1.00	0.12

C1 (Methane)	57.67	59.30	70.23	48.07	73.93	68.05	52.9	66.17
C2 (Ethane)	19.94	17.73	13.94	18.78	13.25	14.2	11.32	13.15
C3 (Propane)	11.33	9.42	6.7	14.87	5.55	8.05	8.52	7.01
I-C4 (Isobutane)	0.97	0.70	5.5	16.38	4.32	6.22	6.46	9.37
N-C4 (N-butane)	2.83	2.03	0	0	0	0	0	0
I-C5 (Isopentane)	0.38	0.27	0	0	0	0	0	0
N-C5 (N-pentane)	0.55	0.38	0	0	0	0	0	0
C6 (Hexane)	0.22	0.16	0	0	0	0	0	0
C7	0.09	0.07	0	0	0	0	0	0
C8	0.04	0.03	0	0	0	0	0	0
C9	0.01	0.01	0	0	0	0	0	0
Specific gravity	0.87	0.84	0.79	1.04	0.76	0.82	0.93	0.85

The experimental data points and gas composition data were transferred to excel to calculate the hydrate formation temperature for all the empirical models reviewed in this study.

3.2 MODEL DEVELOPMENT

The model was developed by carrying out nonlinear regression on the 460 experimental data points in MATLAB 2020 using the curve fitting tool. Using the Custom Equation tool on MATLAB, a custom equation was obtained which achieved a 0.9742 R-square value indicating a 97.42% accuracy fitting the curve of all 460 experimental data points. The following steps were taken to carry out the regression analysis.

1. All 460 experimental data points of pressure, specific gravity, and temperature were imported into MATLAB as x, y, and z respectively with x and y being the input (independent variables) and z the output (dependent variable)
2. The curve fitting tool was selected, and the Nonlinear Least Square method was selected as shown in Figure 3.1

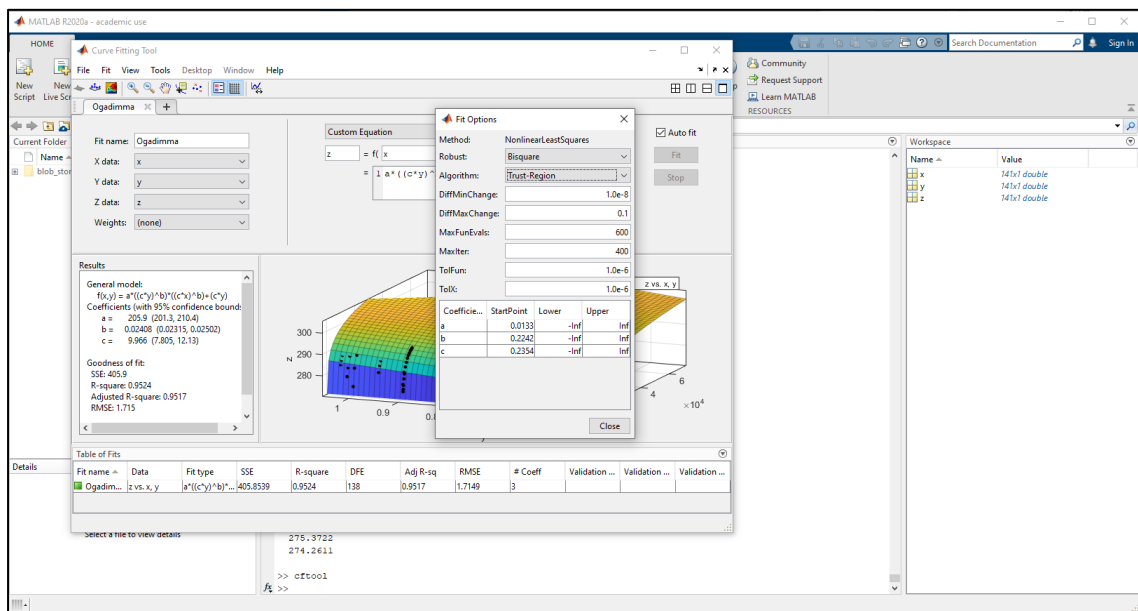


Figure 3.1 The Nonlinear Least Square method

3. By continuous trial, various equations were written inside the custom equation toolbox until an equation that fit the curve of all 460 data points more accurately was written. It produced an R-square value of 0.9742, this indicates a 97.42% coverage of the experimental data points as shown in Figure 3.2

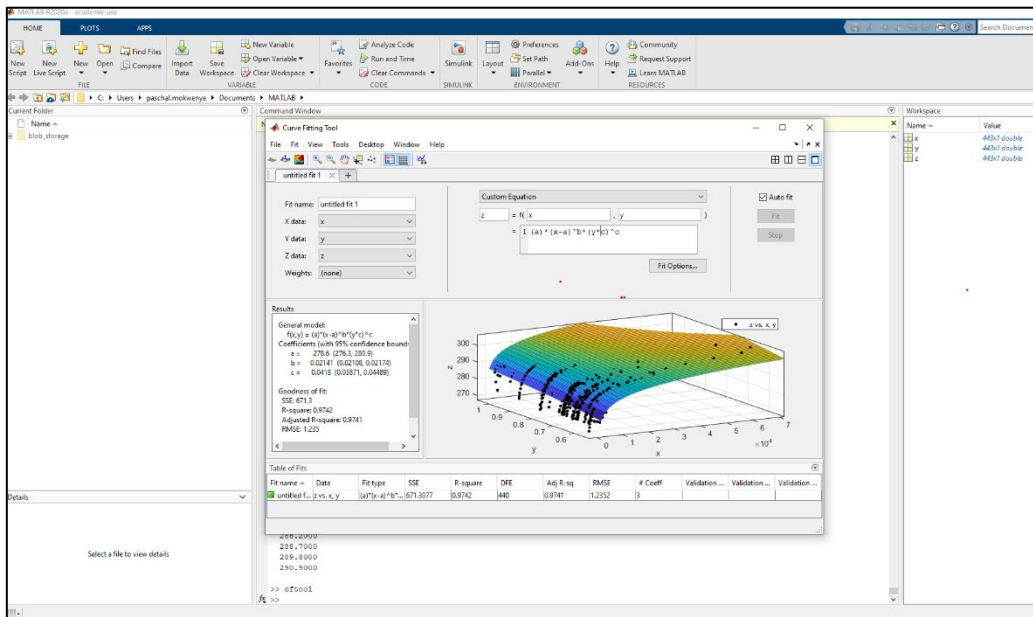


Figure 3.2 Custom equation giving 97.42% accuracy

- The final equation called New Correlation was obtained, and the coefficients of A, B, and C were defined from the regression shown in Figure 3.3

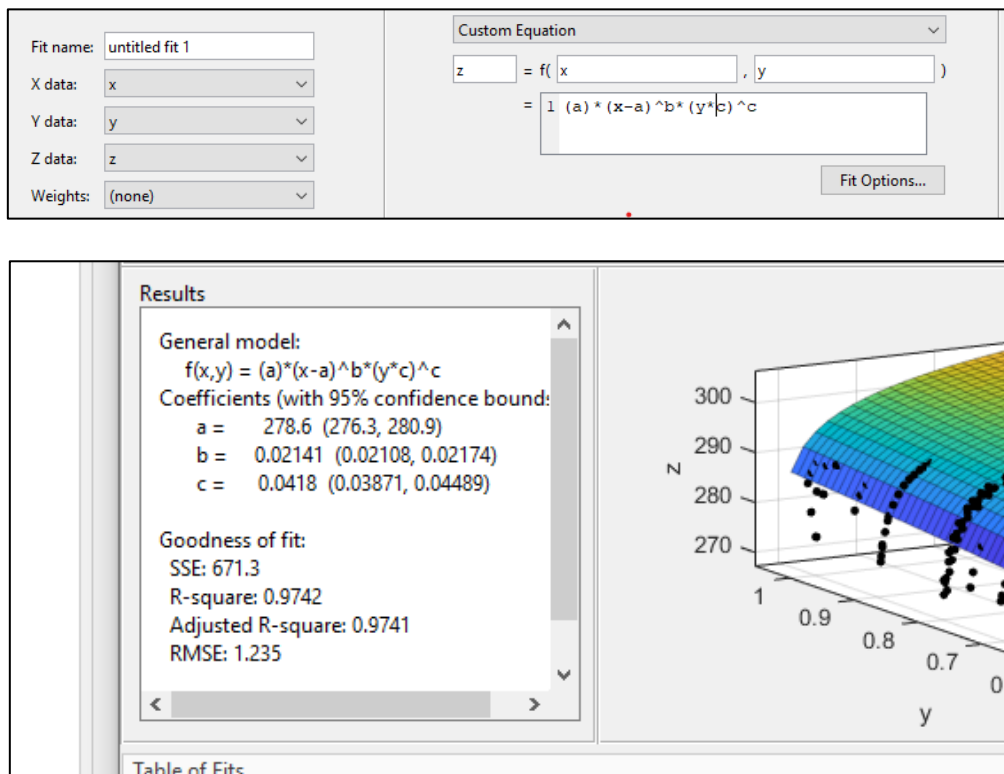


Figure 3.3 Custom equation and coefficients of A, B, and C

The equation for New Correlation is given in Equation 6, with the derived coefficients of A, B, and C given below. These coefficients were tuned optimally by MATLAB to cover the temperature range of 260 to 305 K.

$$T = A \times (P - A)^B \times (\gamma \times C)^C \dots \dots \dots (6)$$

$$A = 278.6$$

$$B = 0.02141$$

$$C = 0.0418$$

Where

$T =$ Hydrate formation temperature (in Kelvin)

$\gamma =$ Specific gravity

$P =$ Pressure (in Kilopascals)

The molecular weight and specific gravity of the gas can be calculated from the gas composition of the gas mixture as shown in Equations 6 and 7.

Molecular Weight (MW^{NG})

$$= \sum \left[\frac{\text{mol \%}}{100} \times \text{Molecular weight of gas} \right] \dots \dots \dots (7)$$

$$\text{Specific gravity } (\gamma) = \frac{MW^{NG}}{MW^{air}} \dots \dots \dots (8)$$

$MW^{NG} =$ Molecular weight of Natural gas

$MW^{air} =$ Molecular weight of Air

The correlation was carried out in Excel with the values of pressure and specific gravity used to calculate the hydrate formation temperature.

The gas composition data of the Bakken formation obtained from EERC were designated as Gas Composition A to H.

3.3 MODEL EVALUATION

The proposed model New Correlation was evaluated based on its accuracy in calculating the reference experimental hydrate formation temperature from the 460 data points, it was also compared to models developed by Hammerschmidt (1934), Motiee (1991), Towler and Mokhatab (2005), Safamirzaei M. (2015), and Chavoshi S. 2018.

The accuracy of the models was evaluated based on their deviation from reference experimental values. The Average Relative Deviation (ARD) and Average Absolute Deviation (AAD) was used to assess the deviation of the models, it shows the suitability of a correlation to calculate the hydrate formation temperature by highlighting its deviation from the reference experimental value (Safamirzaei M. 2015). The equation for Average Relative Deviation (ARD) and Average Absolute Deviation (AAD) are shown in Equations 9 and 10.

$$ARD (\%) = \frac{100}{n} \sum_{i=1}^n \left| \frac{T_{(K)}^{exp} - T_{(K)}^{cal}}{T_{(K)}^{exp}} \right| \dots \dots \dots (9)$$

$$AAD = \frac{1}{n} \left| T_{(K)}^{exp} - T_{(K)}^{cal} \right| \dots \dots \dots (10)$$

where:

n = number of data points

$T_{(K)}^{exp}$ = Experimental hydrate formation temperature

$T_{(K)}^{cal}$ = Calculated hydrate formation temperature

The models will also be evaluated based on their number of adjustable parameters, the higher the parameters the more the models will require complex calculations to

calculate the hydrate formation temperature, this will lead to errors when carrying simple hand calculations.

Using the New correlation, the hydrate formation temperature was calculated and compared to experimental values as well as other correlations. The gas compositions of natural gas from The Bakken were applied to the New Correlation and the hydrate formation temperature was calculated and also compared to values of Katz 1945 gas-gravity chart.

CHAPTER 4 RESULTS

The result of the hydrate formation temperature calculated for all the models including the New Correlation are given in Appendix B. Calculations were made using the pressure and specific gravity of 460 experimental data points obtained from the studies 15 listed in Appendix A. The specific gravity of the gas mixtures ranges from 0.55 – 1.

4.1 AVERAGE RELATIVE DEVIATION (ARD) AND AVERAGE ABSOLUTE DEVIATION (AAD)

Table 4.1 shows the Average Relative Deviation (ARD) and Average Absolute Deviation (AAD) of the models from the reference 460 experimental values of hydrate formation temperature. Table 4.1 also shows the number of adjustable parameters of each model. The models include the New Correlation (NC), Hammerschmidt (1934), Motiee (1991), Towler and Mokhatab (2005), Chavoshi S. 2018, and Safamirzaei M. (2015). A lower value of ARD indicates a lower deviation from experimental values and therefore a higher accuracy.

Table 4.1 ARD and AAD of all empirical models compared to experimental hydrate formation temperature.

Correlations	Average Relative Deviation (ARD)	Average Absolute Deviation (AAD)	Number of Adjustable Parameters
New Correlation	0.393	1.115	3
Towler and Mokhatab (2005)	0.485	1.395	4
Motiee (1991)	0.592	1.675	6
Hammerschmidt (1934)	0.998	2.892	2
Chavoshi S. (2018)	0.529	1.497	3
Safamirzaei M. (2015)	0.406	1.157	3

The model by Motiee (1991) has the highest number of adjustable parameters with 6, this is followed by Towler and Mokhatab (2005) with 4 parameters. Models with a high number of parameters require a long and complex calculation to obtain results, this is prone to errors when doing quick hand calculations. The New correlation, Safamirzaei M. (2015) and Chavoshi S. (2018) have 3 adjustable parameters while Hammerschmidt (1934) has 2 parameters.

Table 4.2 shows the ARD and AAD of the models compared to experimental values obtained from the study by Wilcox W. et al (1941) with specific gravity ranging from 0.597 to 0.668.

Table 4.2 ARD and AAD of models compared to experimental values from Wilcox W. et al (1941).

Models	Average Relative Deviation ARD	Average Absolute Deviation AAD
New Correlation	0.241	0.700
Towler and Mokhatab (2005)	0.437	1.282
Motiee (1991)	0.618	1.779
Hammerschmidt (1934)	1.275	3.747
Chavoshi S. (2018)	0.314	0.909
Safamirzaei M. (2015)	0.360	1.048

Table 4.3 shows the ARD and AAD of the models compared to experimental values obtained from Sloan D. (1990) with specific gravity ranging from 0.57 to 0.8.

Table 4.3 ARD and AAD of the models compared to experimental values obtained from Sloan D. (1990)

Models	Average Relative Deviation ARD	Average Absolute Deviation AAD
New Correlation	0.353	1.006
Mokhatab 2005)	0.441	1.268
Motiee (1991)	0.670	1.901
Hammerschmidt (1934)	0.955	2.778
Chavoshi S. 2018	0.496	1.407
Safamirzaei M. (2015)	0.436	1.250

Table 4.4 shows the ARD and AAD of all 6 models compared to experimental values obtained from the study carried out by Bahadori and Vuthaluru (2009).

Table 4.4 ARD and AAD of all 6 models compared to experimental valves obtained from (Bahadori and Vuthaluru, 2009)

Models	Average Relative Deviation ARD	Average Absolute Deviation AAD
New Correlation	0.550	1.543
Towler and Mokhatab (2005)	0.646	1.841
Motiee (1991)	0.451	1.277
Hammerschmidt (1934)	1.393	3.973
Chavoshi S. (2018)	0.662	1.853
Safamirzaei M. (2015)	0.465	1.305

Table 4.5 shows the ARD and AAD of all 6 models compared to experimental values obtained from the study carried out by Davarnejada R. (2014) on the Lavan gas field which has a specific gravity of 0.65.

Table 4.5 ARD and AAD all 6 models compared to experimental valves obtained from (Davarnejada R. 2014)

Models	Average Relative Deviation ARD	Average Absolute Deviation AAD
New Correlation	0.480	1.380
Towler and Mokhatab (2005)	0.951	2.787
Motiee (1991)	0.336	0.962
Hammerschmidt (1934)	2.080	6.163
Chavoshi S. (2018)	0.463	1.310
Safamirzaei M. (2015)	0.320	0.918

CHAPTER 5 DISCUSSION

The most accurate way of determining the hydrate formation conditions of natural gas transported in pipelines is by measuring the formation pressure and temperature directly, this process is expensive, time-consuming, and not feasible for long and subsea pipelines. As a result, correlations, and thermodynamic models have been developed to estimate the hydrate formation conditions without carrying out experiments. When using thermodynamic models, it is possible to improve the model's accuracy by adjusting parameters, if the predicted results significantly deviate from experimental values. The ARD and AAD of the New Correlation was compared to that by Hammerschmidt (1934), Motiee (1991), Towler and Mokhatab (2005), Chavoshi S. 2018, and Safamirzaei M. (2015) to assess its accuracy. The result of the pressure-temperature curve of all the models will be discussed in this chapter.

5.1 Analysis of Pressure -Temperature Curves, Average Relative Deviation (ARD) and Average Absolute Deviation (AAD) of the Models

Figure 5.1 shows the pressure-temperature curve of 460 experimental data points compared to the values calculated by the 6 models. Specific gravity range from 0.55-1. The New correlation (colored yellow) showed the highest accuracy in following the

experimental curve compared to other models. It tracked the experimental curve more closely across the range of pressure and specific gravity. There is a slight deviation at higher pressure, but overall, the New Correlation followed the experimental points very closely.

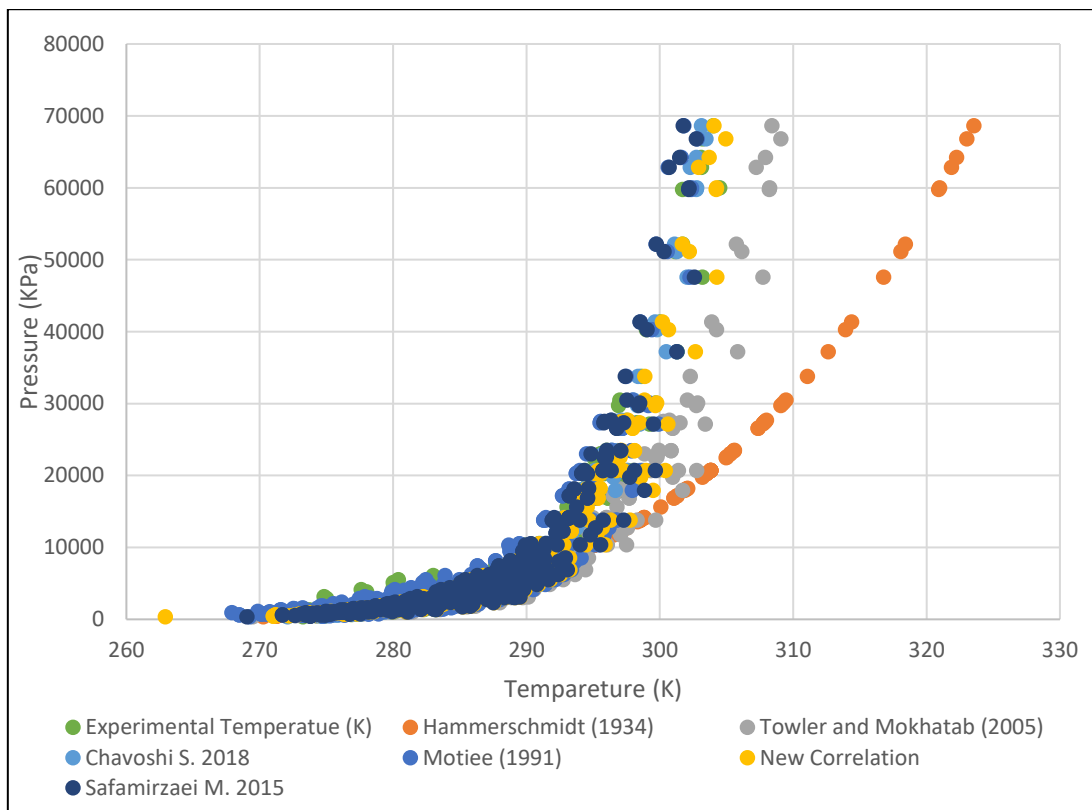


Figure 5.1 Pressure – temperature curve of experimental values from 460 data points compared to all 6 models

Figure 5.2 shows the bar chart of ARD and AAD for all 6 models, the New correlation has the lowest ARD of 0.3932 and AAD of 1.1149 compared to other models. This shows that the New correlation has superior accuracy in calculating the HTF across the range of specific gravity, due to its lower deviation from experimental values compared to the other models. Safamirzaei M. (2015) model shows the second-highest accuracy followed by the Towler and Mokhatab (2005)

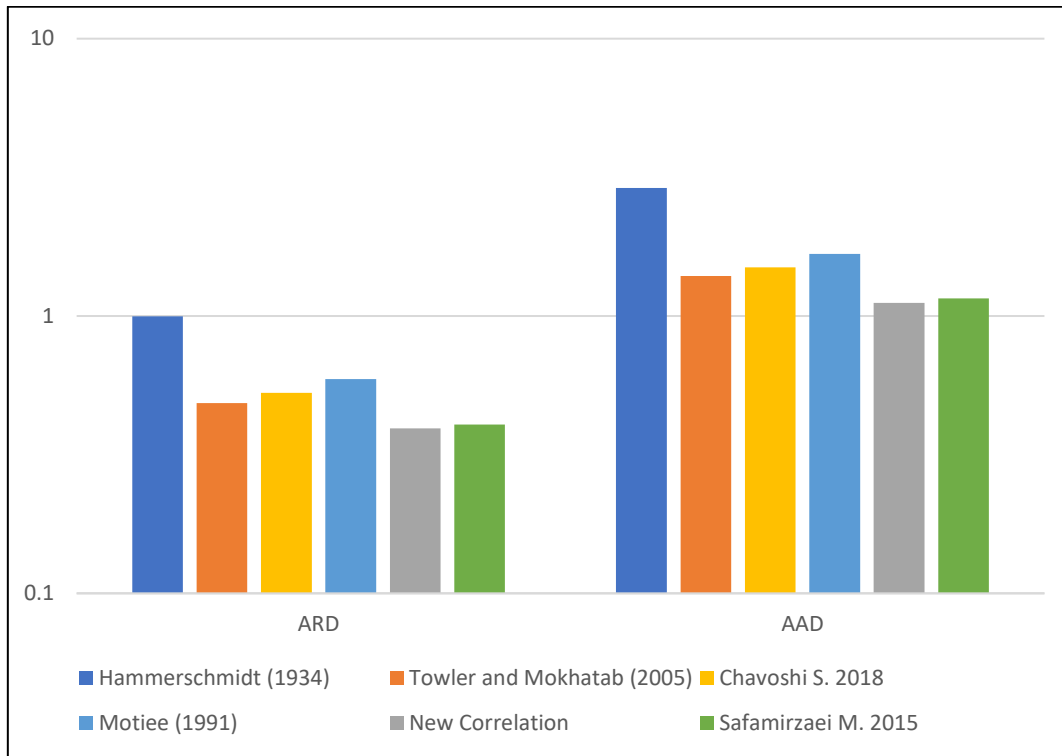


Figure 5.2 ARD and AAD of the models from 460 experimental values.

Figure 5.3 shows the pressure-temperature curve of experimental values obtained from Wilcox W. et al (1941) compared to all 6 models. The study has 32 data points with the specific gravity ranging from 0.59-0.67. The New Correlation shows excellent accuracy, tracking the experimental points more accurately than all the other models. This is confirmed by the values of ARD and AAD in Figure 5.4 which shows that New Correlation has the lowest ARD of 0.2413 and AAD of 0.4644. It is followed by Chavoshi S. 2018 which has the second-lowest ARD and AAD.

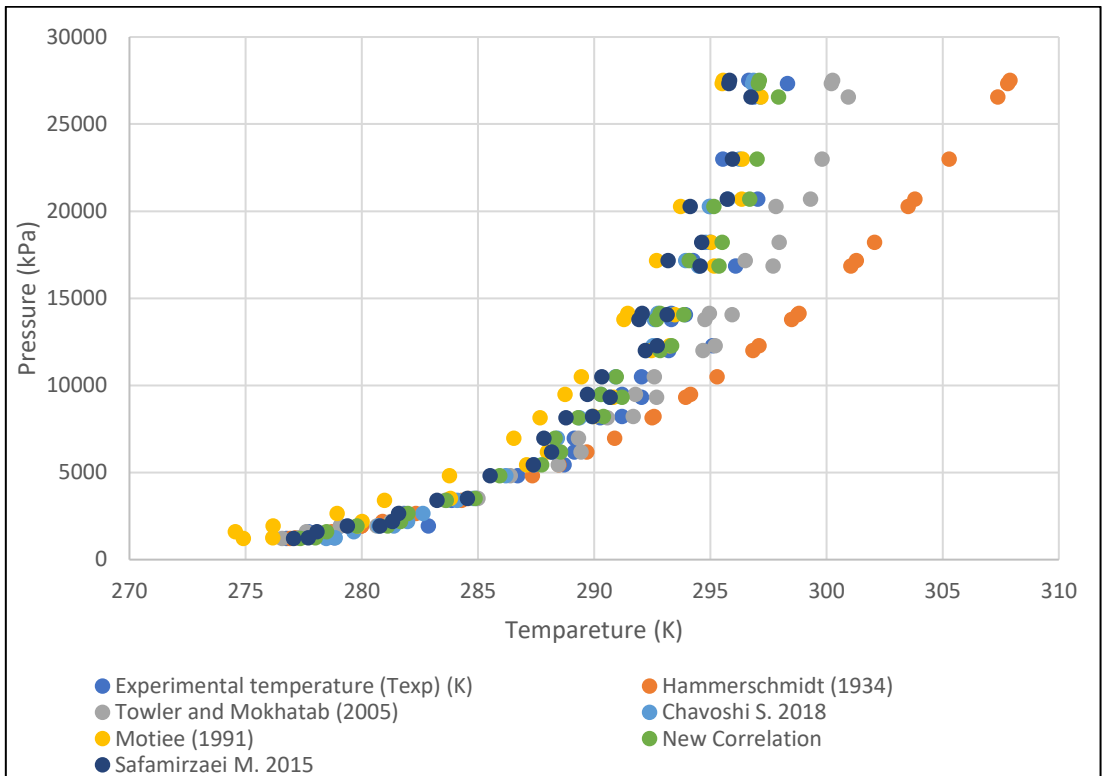


Figure 5.3 Pressure - temperature curve of experimental data obtained from Wilcox W. et al (1941) and all 6 models.

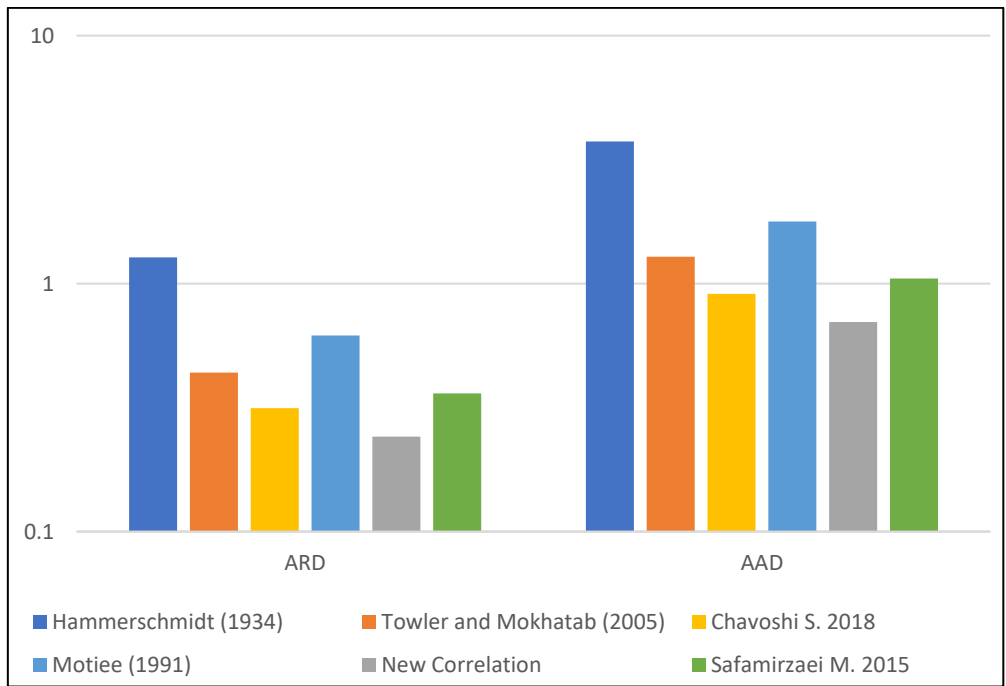


Figure 5.4 ARD and AAD of all 6 models from the experimental values of Wilcox W. et al (1941).

Figure 5.5 shows the pressure-temperature curve of experimental values obtained from Sloan D. (1990) compared to the 6 models. The study has 123 data points with the specific gravity ranging from 0.58-0.80. The New Correlation produced the highest accuracy following the experimental points more accurately than the other models. This is followed by Safamirzaei M. 2015 which produced the second-highest accuracy. The chart of ARD and AAD, showed in Figure 5.6, shows that the New Correlation has the lowest deviation from experimental values with ARD of 0.3524 and AAD of 1.0057. Safamirzaei M. 2015 has the second-lowest deviation with ARD of 0.4360 and AAD of 1.2498.

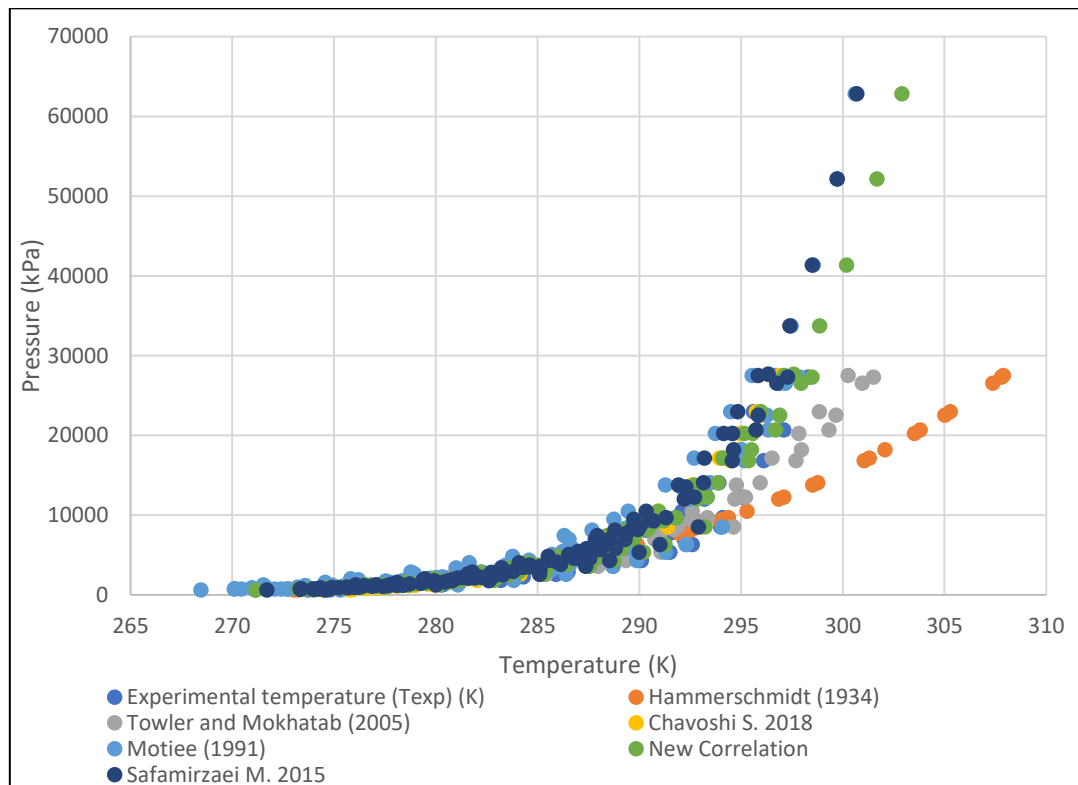


Figure 5.5 Pressure – temperature curve of experimental values obtained from Sloan D. (1990) compared to the 6 models.

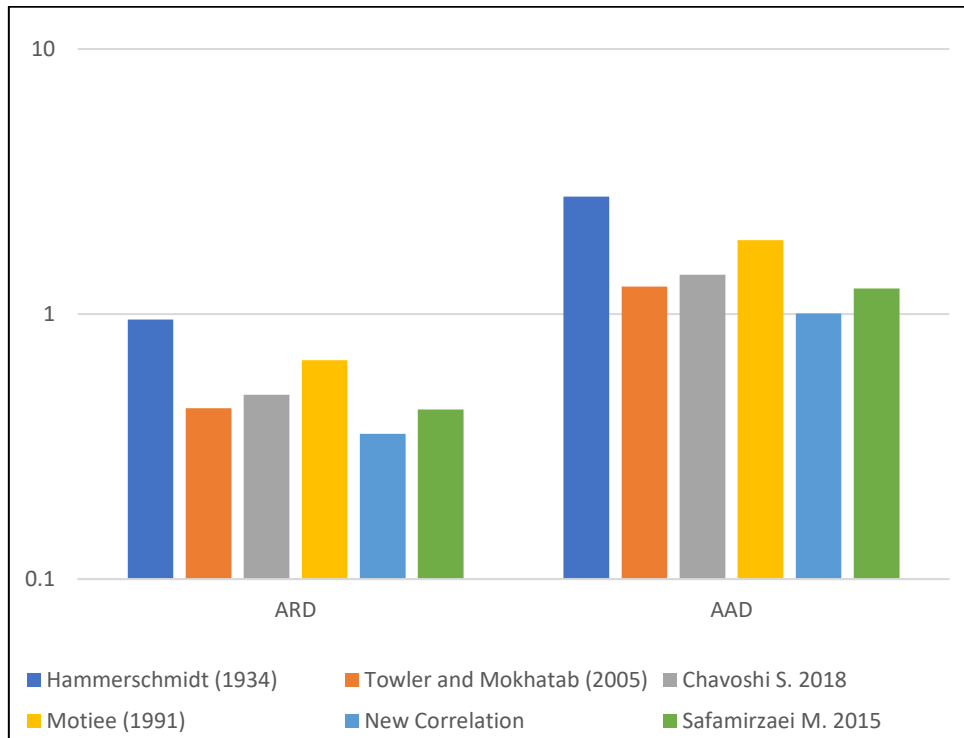


Figure 5.6 ARD and AAD of all 6 models for experimental values obtained from Sloan D. (1990)

Figure 5.7 shows the pressure-temperature curve of experimental data collected from (Bahadori and Vuthaluru, 2009) compared to values from all 6 models. The study has 22 data points with the specific gravity ranging from 0.55-1. The model by Motiee (1991) produced the highest accuracy, tracking the experimental point, this is followed by the Safamirzaei M. (2015) model. The New Correlation produced the third-highest accuracy compared to the other models, it followed the experimental curve closely except for slight deviations at high pressure. The chart of ARD and AAD in Figure 5.8 shows that the model by Motiee (1991) with the ARD of 0.4514 and AAD of 1.2765. Safamirzaei M. (2015) showed the second-lowest ARD and AAD, followed by the New Correlation produced the third-lowest ARD of 0.5503 and AAD of 1.5427.

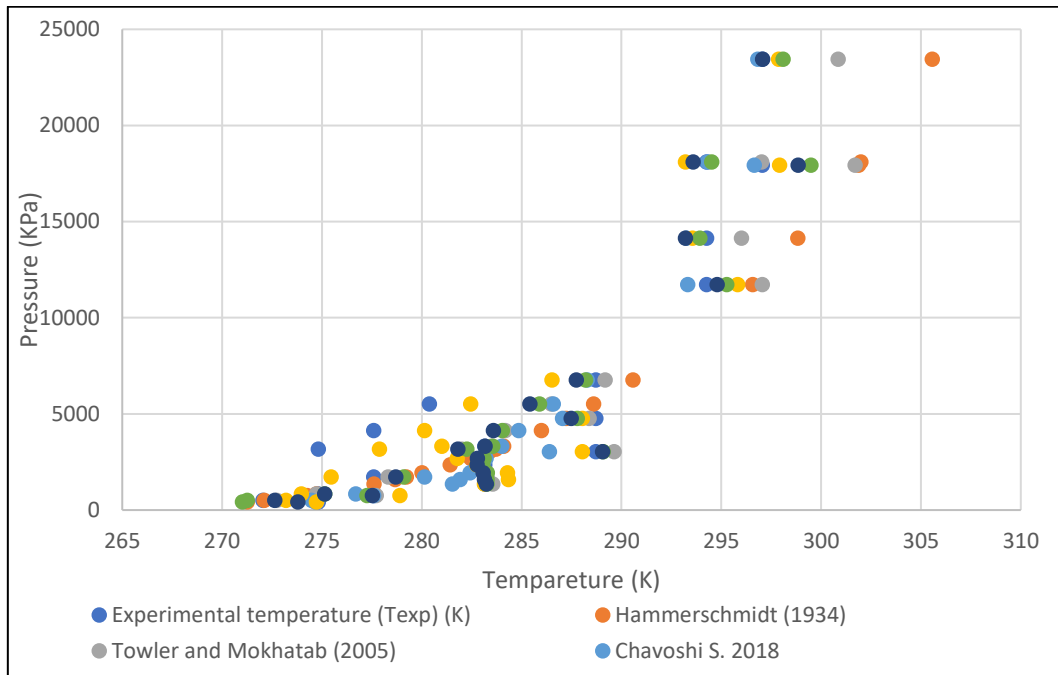


Figure 5.7 Pressure – temperature curve of experimental data from (Bahadori and Vuthaluru, 2009) compared to the 6 models

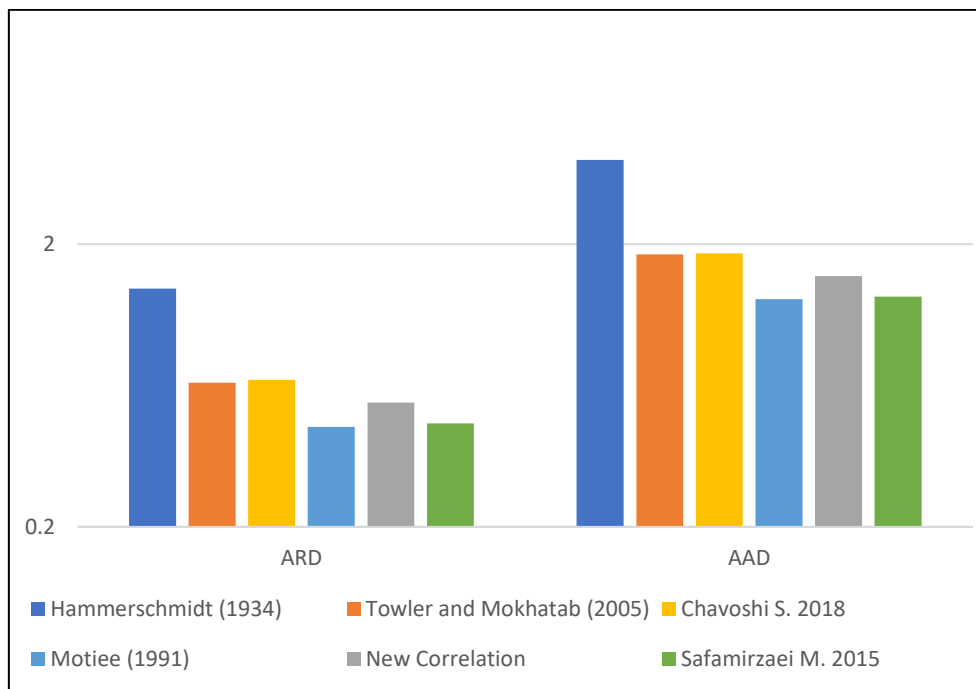


Figure 5.8 ARD and AAD of the models compared to experimental data from (Bahadori and Vuthaluru, 2009)

Figure 5.9 shows the pressure-temperature curve of experimental data collected by Davarnejad, R. (2014) from the Lavan gas field with a specific gravity of 0.65. The curve is compared to that of all 6 models. The curve shows that Safamirzaei M. (2015) produced the highest accuracy, tracking the experimental point closely, this is followed by the model by Motiee (1991). The New Correlation produced the third-highest accuracy following the experimental curve closely up to the pressure of 10000 kPa where it starts to deviate. The chart of ARD and AAD in Figure 5.10 shows that Safamirzaei M. (2015) has the lowest ARD (0.3203) and AAD (0.9184) followed by Motiee (1991). The New Correlation produced the third-lowest ARD and AAD with values of 0.4796 and 1.3803, respectively.

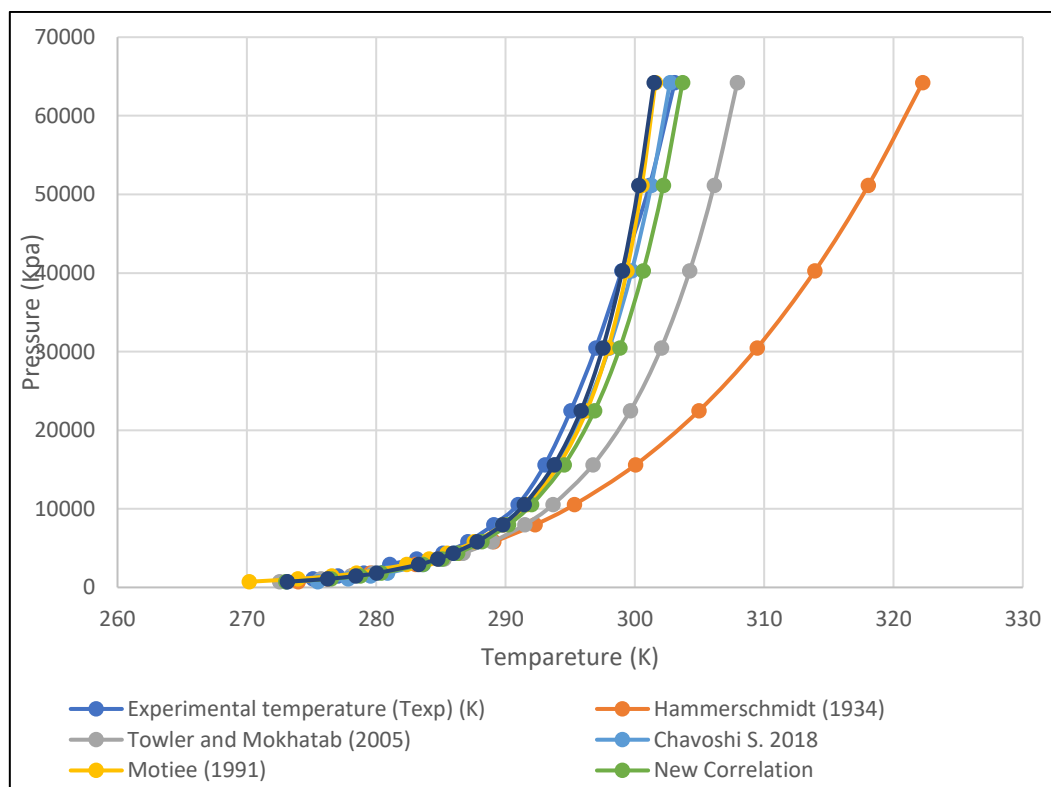


Figure 5.9 Pressure – temperature curve of experimental data collected by Davarnejad, R. (2014) compared to the 6 models.

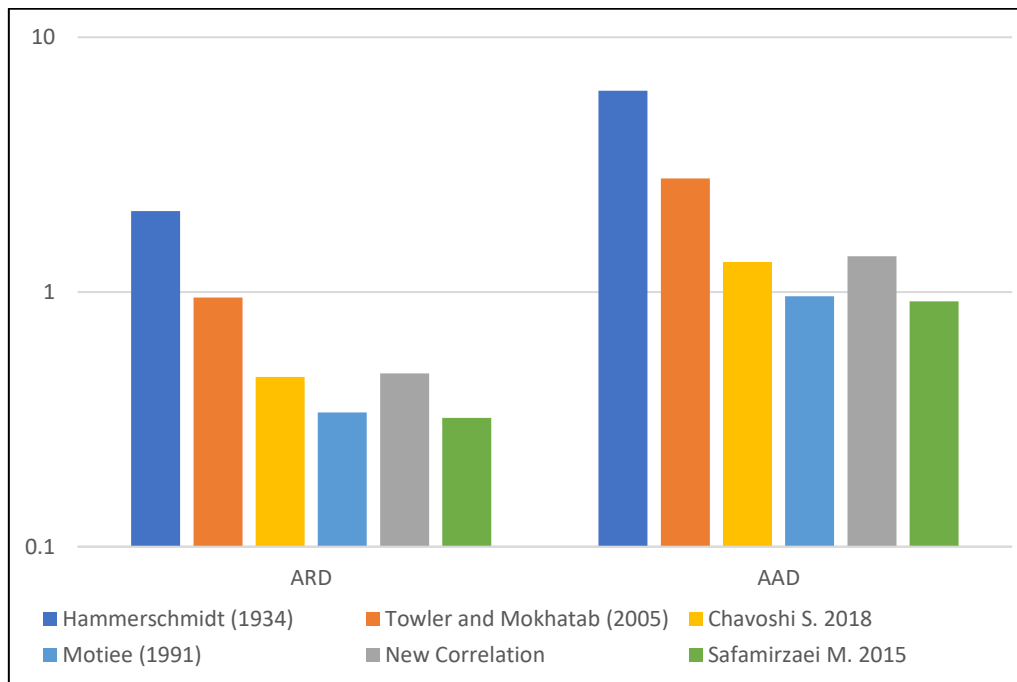


Figure 5.10 ARD and AAD of the models compared to experimental data from Davarnejad, R. (2014)

The New Correlation produced consistently high accuracy and overall superior accuracy in predicting the hydrate formation temperature of natural gas mixtures compared to other models. The experimental values of pressure and specific gravity range from 330 to 68600 kPa and 0.55 to 1.0 respectively. The New correlation having only three adjustable parameters, requires less complex calculations than those proposed by Motiee (1991), Towler and Mokhatab (2005), and Safamirzaei M. (2015) which have logarithmic functions and as such require more complex calculations. The New Correlation offers a simple mathematical equation that can accurately predict the hydrate formation temperature of various gas mixtures.

Figure 5.11 shows the scatterplot of Experimental HTF from all 460 data points $T(\text{exp})$ versus Calculated HTF from New Correlation $T(\text{cal})$. The plot shows that the points of New Correlation align accurately with experimental hydrate formation temperature indicating a strong agreement with the experimental data, except for some slight deviations at high temperature.

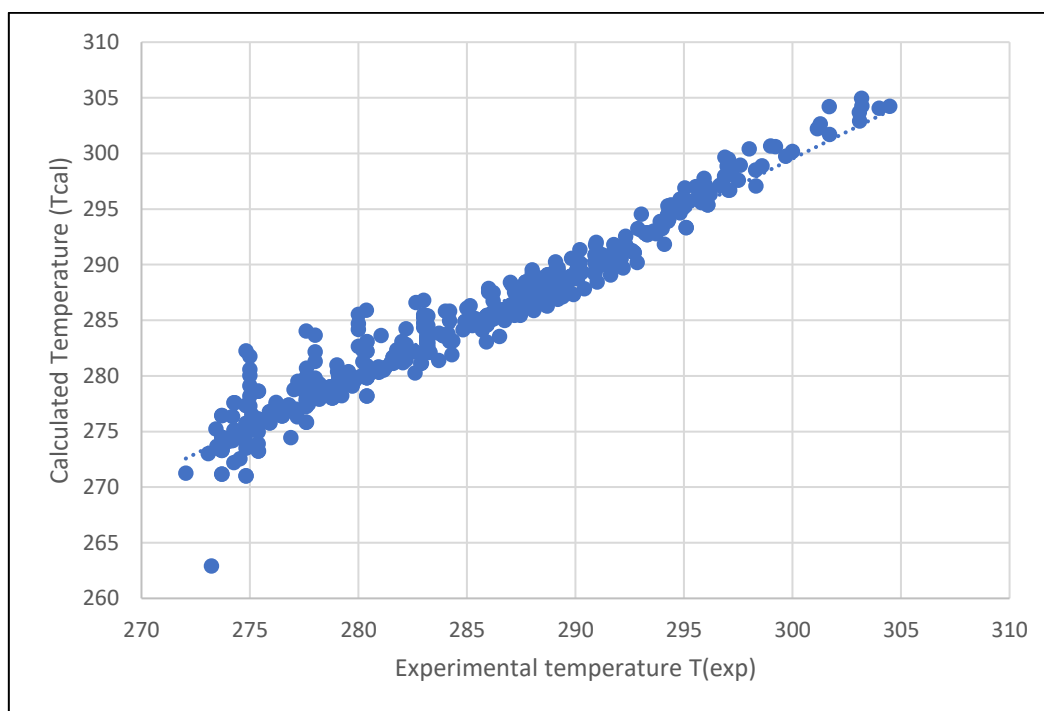


Figure 5.11 Scatterplot of Experimental temperature $T(\text{exp})$ (K) versus Calculated Temperature $T(\text{cal})$ (K) from New Correlation

5.2 Analysis of the Hydrate Formation Temperature (HTF) for The Bakken Gas Compositions

Figure 5.12 shows the pressure-temperature curve of The Bakken gas composition A with a specific gravity of 0.87. The values of HFT were calculated using the New Correlation and compared to Katz's 1945 gravity chart having a specific gravity of

0.9. The curve of New Correlation shows very good accuracy tracking the Katz's 1945 gravity chart, with a slight deviation at higher pressures from 10000 kPa and above.

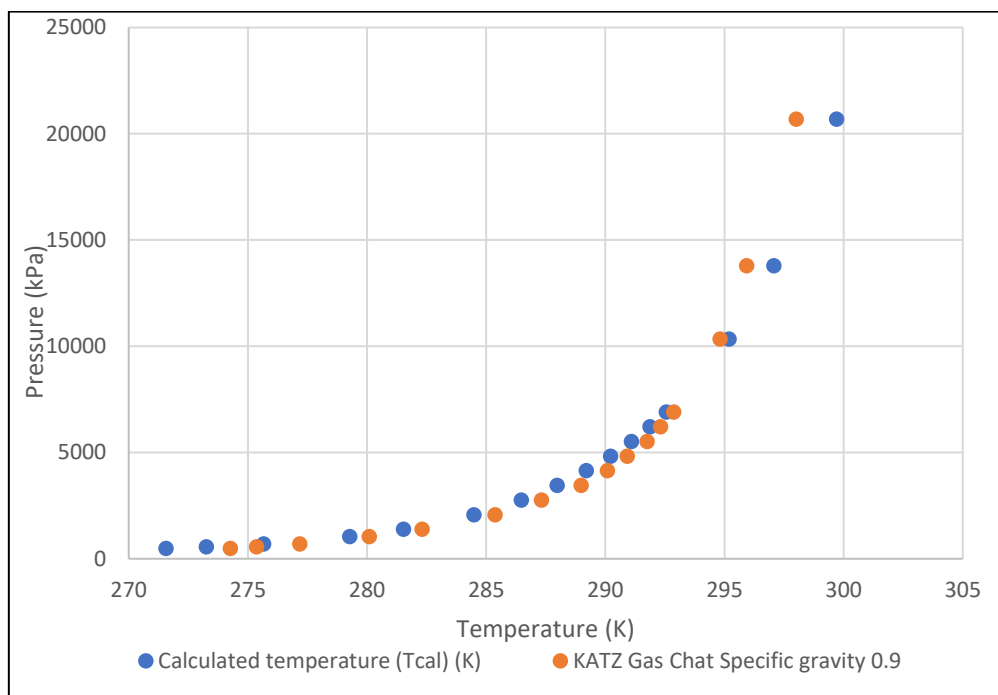


Figure 5.12 Pressure – temperature curve for The Bakken Gas Composition A

The curve also indicates that the hydrate formation temperature increases rapidly at lower pressures than at higher pressures. For example, within the pressure range of 482.63 to 6205.28, there is an increase in hydrate formation temperature of 20.31 K, this is a large increase in temperature for a small increase in pressure. However, moving to a higher pressure range of 6894.75 to 20684.27 kPa there is an increase in hydrate formation temperature of 7.14 K, this represents a small increase in temperature for a large increase in pressure.

The same increase in HFT relative to pressure can be seen in the curve of Gas Composition B shown in Figure 5.13.

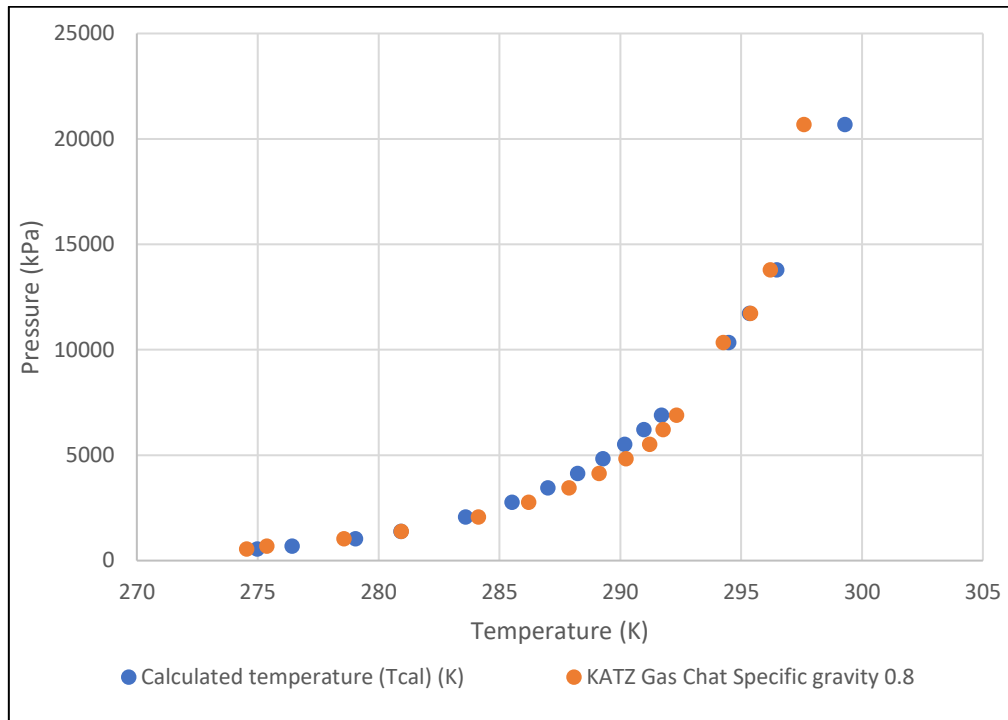


Figure 5.13 Pressure – temperature curve for The Bakken Gas Composition B

Figure 5.13 shows the pressure-temperature curve of The Bakken gas composition B with a specific gravity of 0.84 compared to Katz's 1945 gravity chart having a specific gravity of 0.8. The curve of New Correlation shows very good tracking of Katz's 1945 gravity chart with slight deviations at higher pressures of 14000 kPa and above.

Figures 5.14 shows the pressure-temperature curves for hydrate formation temperature calculated for The Bakken gas compositions C, D, E, F, G, and H. The curves indicate that an increase in the specific gravity causes an increase in the hydrate formation temperature at high pressures. For example, the gas composition E which has a specific gravity of 0.76 produced a HTF of 298.20 K at the pressure of 20684.27 kPa. However, gas composition D having a specific gravity of 1.04 produced a higher HTF of 302.22 (K) at the same pressure. This shows a 4.02 K

increase in HFT moving from the specific gravity of 0.76 to 1.04 suggesting that gases with higher specific gravity produce higher hydrate formation temperature.

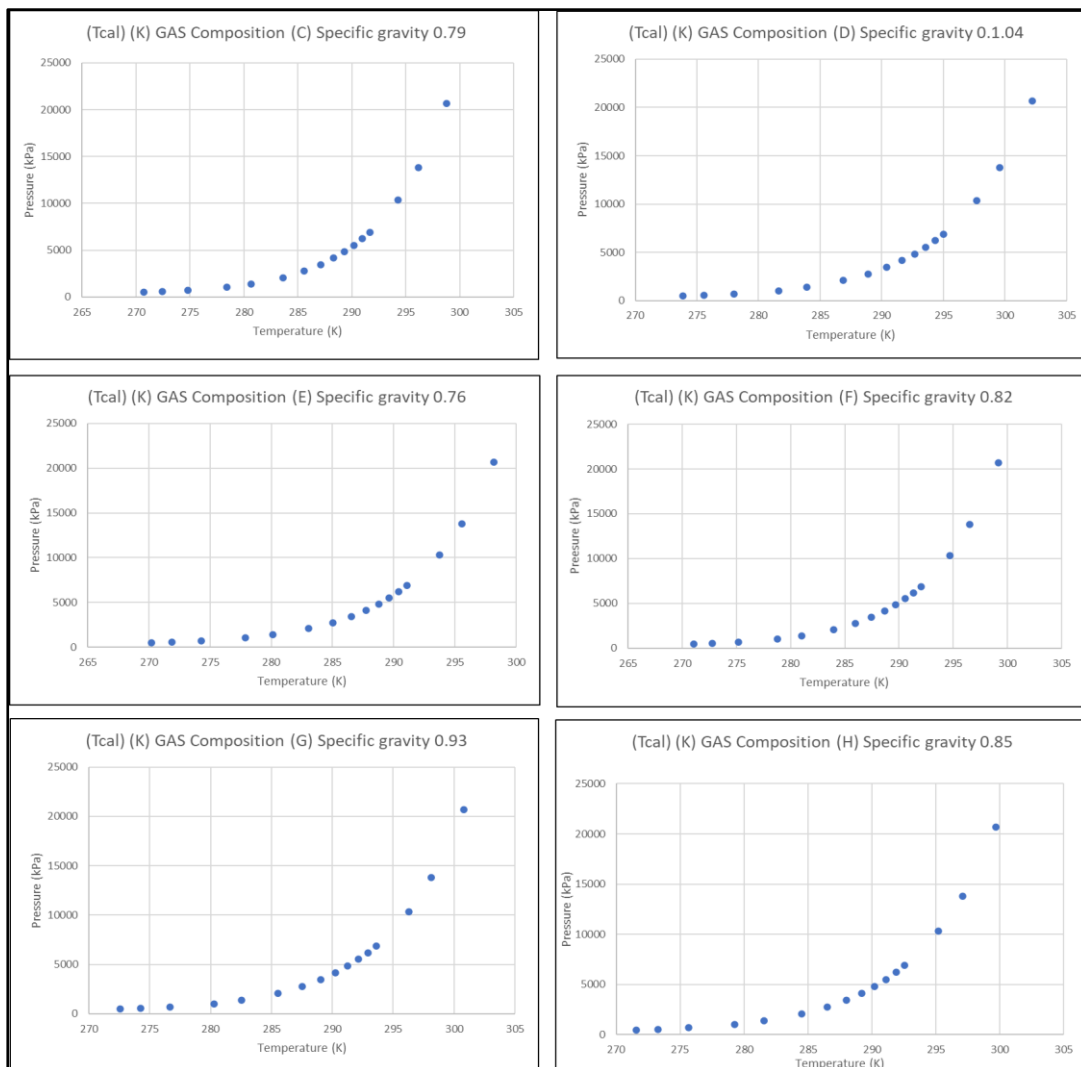


Figure 5.14 Pressure - temperature curve for hydrate formation temperature calculated for The Bakken gas compositions C-H.

Overall, the result of the calculated HTF for The Bakken gas composition A and B compared to Katz's 1945 gravity chart show that the New Correlation calculated the HTF with reliable accuracy and the pressure-temperature curve followed the Katz gravity chart sufficiently.

CHAPTER 6 CONCLUSION AND RECOMMENDATIONS

6.1 CONCLUSION

The model (New Correlation) proposed in this study shows a high potential for estimating the hydrate formation temperature for natural gas with a wide range of gas compositions.

The model's equation, which was developed by nonlinear regression of 460 experimental data points in MATLAB was compared to models by Hammerschmidt (1934), Motiee (1991), Towler and Mokhatab (2005), Chavoshi S. 2018, and Safamirzaei M. (2015). Overall, the New Correlation produced the highest accuracy in predicting the hydrate formation temperature compared to the other models when all data points were considered.

The following findings were made from the analysis of models.

1. The pressure-temperature curve of 460 experimental data points obtained from 15 studies shows that New Correlation produced the most accurate tracking of experimental points compared to the other models. The Average Relative Deviation (ARD) and Average Absolute Deviation (AAD) show that the New Correlation has the lowest deviation from experimental values.
2. The pressure-temperature curve of experimental values obtained from Wilcox

3. W. et al (1941) having 32 data points with the specific gravity ranging from 0.59-0.67 show that the New correlation produced the highest accuracy in following the experimental points across various specific gravities. The Average Relative Deviation (ARD) and Average Absolute Deviation (AAD) show that the New Correlation has the lowest deviation from experimental values.
4. The pressure-temperature curve of HFT obtained from Sloan D. (1990) having 123 experimental data points with the specific gravity ranging from 0.58-0.80 shows that the New correlation produced the most accurate tracking of HFT points than the other models. The Average Relative Deviation (ARD) and Average Absolute Deviation (AAD) show that the New Correlation has the lowest deviation from experimental values.
5. The pressure-temperature curve of HFT obtained from the study by (Bahadori and Vuthaluru, 2009) having 22 data points show that Motiee (1991) produced the most accurate results of HFT, this is followed closely by Safamirzaei M. (2015). The New correlation produced the third-highest accuracy compared to the other models. ARD and AAD also show that Motiee (1991) has the lowest deviation. The New Correlation has the third-lowest ARD and AAD.
6. The result of HTF for The Bakken Gas Composition with a specific gravity of 0.837 and 0.869 calculated with the New correlation shows that it produced reliable accuracy compared to Katz 1945 gravity chart. There were slight deviations at the higher pressure.

Compared to other reputable models the New Correlation provides consistently high accuracy in predicting the hydrate formation temperature of gas mixtures

with a specific gravity ranging between 0.55 to 1. The model produced the highest accuracy at the range of 0.55-0.8 specific gravity. Also, the New correlation offers less complicated calculations having only three adjustable parameters and without the use of complex logarithmic functions as in the case of other models. It also offers a reliable accuracy in estimating the hydrate formation temperature of natural gas from the Bakken Formation.

6.2 RECOMMENDATIONS

The following recommendation is suggested for future research:

1. Further investigation of correlations of hydrate formation temperatures and pressure.
2. Further tuning of the adjustable parameters to obtain more accurate hydrate formation temperature at higher pressures.
3. Further development of the equation on MATLAB to cover the data points more accurately.
4. Further development of models to predict hydrate formation temperature and pressure.

NOMENCLATURE

ARD	Average Relative Deviation
AAD	Average Absolute Deviation
HFT	Hydrate Formation Temperature
K	Kelvins
kPa	Kilo Pascals
NC	New Correlation
P	Pressure
T(exp)	Experimental Temperature
T(cal)	Calculated Temperature
γ	Specific gravity
<i>Mol %</i>	Mol percentage
MW^{air}	The molecular weight of air
MW^{NG}	The molecular weight of Natural gas

APPENDIX A

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APPENDIX B

	Specific gravity	Pressure (MPa)	Experimental Temperature (K)	Hammerschmidt (1934)	Towler and Mokhatab (2005)	Chavoshi S. 2018	Motiee (1991)	New Correlation	Safamirzaei M. 2015
Davarn ejad, R. (2014).	0.650283952	720	273.08	273.9707836	272.5210923	275.4745036	270.1759873	272.9982404	273.1317095
	0.650283952	1080	275.09	276.249091	275.7176958	277.8301194	273.9416086	276.5065362	276.2500843
	0.650283952	1450	277.03	278.0776387	278.0402815	279.5542858	276.5619575	278.7628952	278.4250623
	0.650283952	1810	279.04	279.5589984	279.7886176	280.8592144	278.4702113	280.3668972	280.0157119
	0.650283952	2900	281.04	283.0367347	283.5049113	283.6532635	282.3431733	283.6121195	283.2741455
	0.650283952	3620	283.12	284.8416385	285.2532475	284.9773257	284.0790016	285.0895484	284.7533444
	0.650283952	4350	285.14	286.4256131	286.7015148	286.0788169	285.4751322	286.2981896	285.9543022
	0.650283952	5800	287.08	289.0789481	288.9695411	287.8123427	287.5854531	288.1716553	287.7928535
	0.650283952	7980	289.08	292.2878634	291.4851061	289.7473599	289.8175138	290.2321001	289.7751073
	0.650283952	10520	290.96	295.3127756	293.6637122	291.4336914	291.6583076	292.0086932	291.4462687
	0.650283952	15600	293.04	300.0591092	296.7698697	293.854972	294.1347004	294.5379281	293.7604101
	0.650283952	22480	295.04	304.9630707	299.6502543	296.118233	296.2754942	296.8861864	295.8389537
	0.650283952	30460	296.98	309.4479798	302.0452589	298.0133741	297.9415271	298.8444589	297.5209881
	0.650283952	40260	298.99	313.9224537	304.2443966	299.7642075	299.3801414	300.6490025	299.0305106
	0.650283952	51140	301.14	318.0532917	306.1302676	301.2738254	300.5443084	302.2022972	300.2995872
	0.650283952	64200	303.08	322.250807	307.9233282	302.7161995	301.5916572	303.6846736	301.4853421
Davarn ejad, R. (2014).	0.794322761	330	273.22	270.2629548	269.4595679	272.0846861	269.2274629	262.9044129	269.0356737
	0.794322761	660	276.88	273.5152436	274.7951432	276.0741405	275.4946336	274.4312199	274.8641138
	0.794322761	1000	279.23	275.7961653	277.9936183	278.4936557	278.9931942	278.2016534	278.1201716
	0.794322761	1340	281.16	277.5728093	280.2464743	280.2105657	281.3411697	280.5112134	280.3195785
	0.794322761	1670	283.17	279.0103886	281.9411336	281.5090441	283.044059	282.1418196	281.9267798
	0.794322761	2340	285.24	281.3957465	284.5377469	283.5102984	285.5478104	284.5262695	284.3160059
	0.794322761	3010	287.32	283.3318279	286.4759211	285.0133498	287.3334757	286.2458276	286.0451032
	0.794322761	4350	289.18	286.4256131	289.3104557	287.2258878	289.8169161	288.7026496	288.4962212

	0.79432 2761	5690	291.32	288.895 5086	291.377 5283	288.850 1941	291.532 0502	290.466 676	290.229 2192
	0.79432 2761	8040	293.32	292.366 7566	294.038 7364	290.954 9051	293.621 0585	292.718 2313	292.397 2432
	0.79432 2761	12700	295.24	297.515 0479	297.557 8743	293.761 7078	296.177 6584	295.680 2706	295.162 8082
	0.79432 2761	19760	297.24	303.173 4299	300.960 6612	296.501 457	298.426 7817	298.543 021	297.735 6967
	0.79432 2761	27120	299.22	307.687 3333	303.397 8146	298.479 4182	299.902 9337	300.598 545	299.521 7806
	0.79432 2761	37180	301.28	312.609 3413	305.826 4062	300.463 5563	301.262 0467	302.654 0869	301.257 4887
	0.79432 2761	47560	303.19	316.770 1223	307.721 7197	302.021 1695	302.245 1498	304.264 494	302.583 0086
Bahador i, A., & Vuthalur u, Hari. B. (2009).	0.55240 1687	3157.8	274.82	283.716 4225	282.000 5391	283.235 4068	277.872 4706	282.251 0391	281.805 3215
	0.55240 1687	4136.85	277.59	285.984 1357	284.170 6253	284.846 2663	280.129 2691	284.025 385	283.574 3892
	0.55240 1687	5515.81	280.37	288.599 7662	286.482 3378	286.572 3342	282.443 3605	285.889 6745	285.408 5002
	0.60073 6835	1723.69	277.59	279.224 5326	278.301 746	280.126 8216	275.451 6788	279.092 7999	278.689 9038
	0.60073 6835	3309.48	283.15	284.097 9555	283.492 6401	283.990 6268	280.996 0843	283.553 9291	283.164 7267
	0.60073 6835	6756.86	288.71	290.578 2392	289.172 4733	288.279 4565	286.515 846	288.203 0726	287.738 3057
	0.60073 6835	18098.7 4	294.26	301.991 9109	297.012 9133	294.306 3729	293.196 4127	294.514 9124	293.582 2567
	0.65079 8238	2688.96	283.15	282.447 3883	282.919 9037	283.208 031	281.762 5015	283.112 2834	282.772 4447
	0.65079 8238	14134.2 5	294.26	298.819 9762	296.001 4173	293.251 3509	293.548 5175	293.914 2014	293.198 4033
	0.70085 9641	827.37	274.82	274.722 3584	274.709 2557	276.693 9621	273.967 4029	275.133 8379	275.127 8253
	0.70085 9641	2344.2	283.15	281.409 05	282.847 2274	282.812 0272	282.754 8276	283.053 6753	282.758 4869
	0.70085 9641	4757.38	288.71	287.228 089	288.377 6773	287.046 8013	288.032 2879	287.782 9025	287.482 6638
	0.70085 9641	23442	297.04	305.558 8564	300.839 8509	296.823 2219	297.864 7615	298.087 7137	297.055 8935
	0.80098 2446	496.42	272.04	272.100 7081	272.728 4615	274.473 6519	273.197 5569	271.254 1523	272.633 9521
	0.80098 2446	1930.53	283.15	280.007 4969	283.172 2431	282.414 5278	284.288 3866	283.279 3538	283.068 6631
	0.80098 2446	11721.0 9	294.26	296.562 5712	297.041 5994	293.316 2703	295.818 7237	295.264 1005	294.794 3891
	0.90110 5252	758.42	277.59	274.248 3921	277.714 7991	277.580 5179	278.902 8523	277.241 2095	277.515 6217
	0.90110 5252	1585.79	283.15	278.664 3739	283.305 9856	281.913 6052	284.340 4869	283.254 4291	283.127 7552
	0.90110 5252	17926.3 7	297.04	301.864 9377	301.689 4967	296.643 076	297.917 1063	299.486 6906	298.839 6097
	1.00122 8058	413.69	274.82	271.253 7388	274.724 7848	274.647 8488	274.723 8211	271.009 9295	273.784 5718
	1.00122 8058	1344.48	283.15	277.593 9376	283.543 6373	281.530 6399	283.136 7091	283.264 2687	283.229 5397
	1.00122 8058	3033.69	288.71	283.394 3677	289.632 4379	286.383 1234	288.035 8513	289.082 5439	289.042 7481

Elgibaly , A. A., & Elkamel , A. M. (1998).	1.03042	1810	284	279.558 9984	286.166 4316	283.456 8141	284.626 9859	285.813 689	285.775 422
	1.03042	2310	286	281.300 2215	287.984 9695	284.912 4998	286.044 6432	287.547 8853	287.516 6857
	1.03042	3080	288	283.515 6211	290.129 7682	286.638 9581	287.630 8659	289.533 3391	289.514 1426
	1.01540 4	990	279	275.737 7478	281.456 3459	279.805 6517	280.989 3904	280.987 8512	281.079 6713
	1.01540 4	1710	283	279.170 3879	285.538 5401	283.035 5984	284.497 6887	285.225 683	285.177 8977
	1.01540 4	2990	288	283.278 7549	289.712 1286	286.376 4086	287.738 1256	289.153 5045	289.123 1409
	0.95243 3	1420	282	277.942 7529	283.248 9986	281.572 4177	283.736 0597	283.087 8987	283.010 0369
	0.95243 3	2140	285	280.741 4657	286.336 8806	284.008 1114	286.406 2706	286.067 7026	285.978 5665
	0.95243 3	3000	287	283.305 3231	288.880 1189	286.030 0098	288.463 7971	288.403 459	288.328 3823
	0.76497 6	945	275	275.469 5195	277.013 8031	277.953 6556	277.627 3433	277.293 012	277.224 369
	0.76497 6	1290	278	277.333 5027	279.420 2976	279.776 1607	280.201 4213	279.780 9622	279.572 0321
	0.76497 6	2430	283	281.677 1661	284.316 984	283.521 5366	285.103 663	284.338 9264	284.104 5821
	0.64630 1	68600	304	323.526 3187	308.381 5967	303.100 6509	301.789 6964	304.039 7149	301.745 2188
	0.64630 1	23500	296	305.594 2139	299.929 6253	296.357 8856	296.400 5857	297.095 5365	296.006 752
	0.64630 1	7000	289	290.934 7426	290.374 7147	288.915 6859	288.757 9037	289.313 2804	288.889 6042
	0.60028 4	1520	275	278.384 7865	277.290 4586	279.383 7978	274.301 8657	278.177 6564	277.781 7221
	0.60028 4	2890	280	283.009 5136	282.403 9982	283.179 2036	279.859 2692	282.642 1012	282.250 2361
	0.60028 4	3970	283	285.627 064	284.930 8172	285.073 6668	282.434 4544	284.744 3761	284.351 3483
	0.57800 2	1840	275	279.672 5802	278.287 4136	280.294 8557	274.706 0219	279.105 2885	278.675 8665
	0.57800 2	2530	278	281.981 2459	280.832 9555	282.175 6173	277.497 2355	281.300 7814	280.879 3237
	0.57800 2	4770	283	287.252 15	285.901 8137	285.958 3871	282.716 4993	285.490 9988	285.059 2615
	0.56783	2160	275	280.808 8134	279.326 0183	281.140 402	275.516 8793	280.013 6794	279.574 9212
	0.56783	2960	278	283.198 6674	281.849 8066	283.006 7921	278.243 8029	282.145 9554	281.717 5112
	0.56783	4030	280	285.756 6827	284.321 4987	284.846 6647	280.806 5011	284.181 6864	283.750 5433
	0.56443 9	2370	275	281.490 3998	279.987 7088	281.654 9716	276.126 7478	280.578 5266	280.138 9068
	0.56443 9	4410	280	286.547 0878	284.965 2364	285.351 9973	281.344 0212	284.698 064	284.257 3325
	0.56443 9	6090	283	289.550 9206	287.552 4392	287.292 7519	283.884 9884	286.785 4562	286.301 2384
	0.55959 5	2860	275	282.927 444	281.378 332	282.719 9981	277.461 638	281.744 3585	281.305 3358
	0.55959 5	3810	278	285.274 4269	283.679 5497	284.427 9473	279.873 8315	283.640 9622	283.202 2403

	0.55959 5	5090	280	287.847 5961	286.003 5629	286.163 2872	282.216 2727	285.525 4592	285.064 4349
KATZ Gas Chat Specific gravity (γ) 0.6	0.6	20684.2 7188	294.4	303.800 2808	298.061 2715	295.125 568	293.988 3914	295.355 3008	294.321 2086
	0.6	13789.5 1459	293.705 5556	298.515 2927	294.834 3024	292.623 3107	291.429 7973	292.759 4775	292.002 8783
	0.6	10342.1 3594	292.038 8889	295.119 1449	292.544 7314	290.860 8079	289.502 6066	290.918 8709	290.309 0518
	0.6	6894.75 7293	289.122 2222	290.781 5353	289.317 7624	288.394 7098	286.628 7551	288.318 2768	287.847 7017
	0.6	6205.28 1564	288.15	289.734 0785	288.479 2312	287.757 3215	285.851 7958	287.639 7477	287.193 1694
	0.6	5515.80 5835	287.316 6667	288.599 759	287.541 8332	287.046 4495	284.968 4883	286.879 1109	286.453 8471
	0.6	4826.33 0105	286.344 4444	287.358 9963	286.479 0989	286.242 653	283.948 2553	286.013 4042	285.605 7016
	0.6	4136.85 4376	285.65	285.984 1449	285.252 2622	285.317 5367	282.745 6105	285.008 381	284.613 0301
	0.6	3447.37 8647	283.705 5556	284.434 1188	283.801 2224	284.227 215	281.288 7768	283.809 6111	283.419 546
	0.6	2757.90 2917	281.761 1111	282.643 4442	282.025 2931	282.898 4379	279.455 0113	282.322 5705	281.929 0211
	0.6	2068.42 7188	279.538 8889	280.496 7009	279.735 7221	281.194 5091	277.008 4548	280.359 7659	279.956 3649
	0.6	1378.95 1459	276.205 5556	277.754 8489	276.508 7531	278.810 3679	273.402 5982	277.454 7669	277.071 1914
	0.6	1034.21 3594	273.427 7778	275.992 9273	274.219 1821	277.131 062	270.732 3629	275.231 0335	274.943 5465
KATZ Gas Chat Specific gravity (γ) 0.7	0.7	20684.2 7188	296.761 1111	303.800 2808	299.847 6496	296.036 8482	297.176 1776	297.264 5668	296.334 8634
	0.7	13789.5 1459	294.816 6667	298.515 2927	296.678 8428	293.526 8645	294.887 6299	294.651 9633	294.000 6718
	0.7	10342.1 3594	293.288 8889	295.119 1449	294.430 5386	291.758 9194	293.152 0401	292.799 4584	292.295 2566
	0.7	6894.75 7293	291.761 1111	290.781 5353	291.261 7318	289.285 2066	290.548 2349	290.182 0533	289.817 0667
	0.7	6205.28 1564	290.927 7778	289.734 0785	290.438 3141	288.645 8502	289.841 4475	289.499 138	289.158 0563
	0.7	5515.80 5835	289.816 6667	288.599 759	289.517 8115	287.932 7832	289.036 5854	288.733 5843	288.413 6758
	0.7	4826.33 0105	288.983 3333	287.358 9963	288.474 2318	287.126 5048	288.105 2865	287.862 2813	287.559 7276
	0.7	4136.85 4376	287.872 2222	285.984 1449	287.269 5073	286.198 5319	287.005 3085	286.850 7614	286.560 2645
	0.7	3447.37 8647	286.483 3333	284.434 1188	285.844 6208	285.104 8436	285.669 9039	285.644 2422	285.358 615
	0.7	2757.90 2917	284.816 6667	282.643 4442	284.100 7005	283.771 9635	283.984 7557	284.147 5889	283.857 8924
	0.7	2068.42 7188	282.594 4444	280.496 7009	281.852 3963	282.062 7734	281.729 8001	282.172 0962	281.871 7399
	0.7	1378.95 1459	279.261 1111	277.754 8489	278.683 5895	279.671 2705	278.393 9899	279.248 3184	278.966 8269
	0.7	1034.21 3594	277.038 8889	275.992 9273	276.435 2853	277.986 7793	275.915 3556	277.010 2101	276.824 6253

	0.7	689.475 7293	273.566 6667	273.742 5752	273.266 4785	275.629 8352	272.264 288	273.420 4038	273.679 7792
KATZ Gas Chat Specific gravity (γ) 0.8	0.8	20684.2 7188	297.594 4444	303.800 2808	301.395 0807	296.828 5091	298.687 801	298.928 4215	298.090 303
	0.8	13789.5 1459	296.205 5556	298.515 2927	298.276 6563	294.311 8132	296.669 2996	296.301 1947	295.742 284
	0.8	10342.1 3594	294.261 1111	295.119 1449	296.064 099	292.539 1403	295.125 3108	294.438 321	294.026 7662
	0.8	6894.75 7293	292.316 6667	290.781 5353	292.945 6746	290.058 8123	292.791 552	291.806 2656	291.533 8959
	0.8	6205.28 1564	291.761 1111	289.734 0785	292.135 3489	289.417 7461	292.154 9363	291.119 5279	290.870 9817
	0.8	5515.80 5835	291.205 5556	288.599 759	291.229 4819	288.702 7722	291.428 5197	290.349 6892	290.122 1915
	0.8	4826.33 0105	290.233 3333	287.358 9963	290.202 4945	287.894 3377	290.586 1548	289.473 5094	289.263 1847
	0.8	4136.85 4376	289.122 2222	285.984 1449	289.016 9246	286.963 8832	289.588 8437	288.456 3278	288.257 8009
	0.8	3447.37 8647	287.872 2222	284.434 1188	287.614 693	285.867 2701	288.374 8683	287.243 0555	287.049 033
	0.8	2757.90 2917	286.205 5556	282.643 4442	285.898 5002	284.530 8257	286.838 3373	285.738 0251	285.539 4204
	0.8	2068.42 7188	284.122 2222	280.496 7009	283.685 9429	282.817 0648	284.774 9826	283.751 475	283.541 5023
	0.8	1378.95 1459	280.927 7778	277.754 8489	280.567 5186	280.419 1666	281.709 2188	280.811 3322	280.619 381
	0.8	1034.21 3594	278.566 6667	275.992 9273	278.354 9613	278.730 1707	279.422 1854	278.560 6967	278.464 4894
	0.8	689.475 7293	275.372 2222	273.742 5752	275.236 5369	276.366 9237	276.041 1642	274.950 7975	275.301 0138
	0.8	551.580 5835	274.538 8889	272.610 6696	273.520 3441	275.074 8939	274.101 7818	272.554 2905	273.491 602
KATZ Gas Chat Specific gravity (γ) 0.9	0.9	20684.2 7188	298.011 1111	303.800 2808	302.760 0116	297.528 5606	298.523 2616	300.403 7739	299.647 34
	0.9	13789.5 1459	295.927 7778	298.515 2927	299.686 0277	295.005 9292	296.774 8066	297.763 5805	297.287 0565
	0.9	10342.1 3594	294.816 6667	295.119 1449	297.505 0014	293.229 0756	295.422 4187	295.891 5126	295.562 5779
	0.9	6894.75 7293	292.872 2222	290.781 5353	294.431 0175	290.742 8979	293.358 7063	293.246 4668	293.056 6864
	0.9	6205.28 1564	292.316 6667	289.734 0785	293.632 2397	290.100 3198	292.792 2624	292.556 3397	292.390 3095
	0.9	5515.80 5835	291.761 1111	288.599 759	292.739 2821	289.383 6596	292.144 2912	291.782 7015	291.637 6082
	0.9	4826.33 0105	290.927 7778	287.358 9963	291.726 9303	288.573 3184	291.390 8604	290.902 1973	290.774 1144
	0.9	4136.85 4376	290.094 4444	285.984 1449	290.558 2558	287.640 6696	290.496 2162	289.879 9954	289.763 4792
	0.9	3447.37 8647	288.983 3333	284.434 1188	289.176 0073	286.541 4702	289.403 6699	288.660 7351	288.548 3974
	0.9	2757.90 2917	287.316 6667	282.643 4442	287.484 2719	285.201 8739	288.015 7562	287.148 2766	287.030 8995
	0.9	2068.42 7188	285.372 2222	280.496 7009	285.303 2456	283.484 0712	286.144 0024	285.151 922	285.022 5455

	0.9	1378.95 1459	282.316 6667	277.754 8489	282.229 2617	281.080 5177	283.348 285	282.197 2682	282.085 1609
	0.9	1034.21 3594	280.094 4444	275.992 9273	280.048 2354	279.387 5384	281.252 8525	279.935 5248	279.919 0135
	0.9	689.475 7293	277.177 7778	273.742 5752	276.974 2515	277.018 7178	278.141 8777	276.307 809	276.739 0138
	0.9	551.580 5835	275.372 2222	272.610 6696	275.282 5161	275.723 6408	276.351 1125	273.899 4742	274.920 1508
	0.9	482.633 0105	274.261 1111	271.966 9619	274.270 1643	274.951 5509	275.252 7879	272.197 6077	273.806 6654
Wilcox, W. I., Carson, D. B., & Katz, D. L. (1941).	0.66840 8755	1254.84 5827	278.816 6667	277.161 2497	277.290 9389	278.860 2461	276.176 0821	277.996 6125	277.704 4484
	0.66840 8755	1923.63 7285	282.872 2222	279.982 3971	280.648 0074	281.373 2426	279.820 6202	281.119 7542	280.787 0148
	0.66840 8755	12265.7 7322	295.094 4444	297.099 2688	295.206 0921	292.535 652	293.256 5023	293.331 2737	292.713 0921
	0.66840 8755	16843.8 9207	296.094 4444	301.046 9208	297.698 5238	294.490 6428	295.170 8218	295.369 8037	294.560 043
	0.66840 8755	20684.2 7188	297.038 8889	303.800 2808	299.312 4867	295.763 5522	296.350 2355	296.691 2979	295.730 1718
Gas C	0.59666 8956	27323.9 2315	298.316 6667	307.799 1444	300.213 8086	296.822 924	295.509 4684	297.072 8974	295.796 1115
	0.59666 8956	1599.58 3692	277.705 5556	278.721 9369	277.612 205	279.649 5863	274.552 0953	278.477 6549	278.073 6864
	0.59666 8956	3392.22 0588	283.872 2222	284.300 8315	283.598 972	284.099 3242	280.982 9584	283.636 9176	283.243 1669
	0.59666 8956	6963.70 4866	289.15	290.882 0931	289.326 7983	288.422 8622	286.542 3026	288.315 1778	287.838 3004
	0.59666 8956	10500.7 1536	292.038 8889	295.291 8951	292.597 8107	290.921 3749	289.456 7975	290.948 5563	290.328 2073
Gas C 2nd Series	0.59666 8956	14127.3 5769	293.316 6667	298.813 937	294.960 4491	292.739 4918	291.444 2195	292.846 1671	292.071 5631
	0.59666 8956	1923.63 7285	279.094 4444	279.982 3971	279.081 3344	280.735 0395	276.188 8963	279.788 7586	279.380 4405
	0.59666 8956	2647.58 6801	281.816 6667	282.328 0021	281.625 2382	282.624 5535	278.932 8575	281.981 9599	281.583 3241
	0.59666 8956	4819.43 5348	286.705 5556	287.345 9665	286.395 6588	286.202 191	283.769 8734	285.937 5654	285.526 2439
	0.59666 8956	8135.81 3606	290.261 1111	292.491 8723	290.565 69	289.366 6361	287.668 4271	289.314 1257	288.792 1528
	0.59666 8956	9480.29 1278	291.205 5556	294.145 612	291.783 6779	290.297 4962	288.749 0878	290.294 1908	289.716 9829
	0.59666 8956	13782.6 1983	293.316 6667	298.509 1437	294.763 7032	292.587 6577	291.282 4877	292.688 1589	291.928 0741
	0.59666 8956	20263.6 9168	294.983 3333	303.517 5761	297.833 1549	294.965 4335	293.727 7061	295.154 9405	294.133 1721
	0.59666 8956	17161.0 509	294.261 1111	301.290 3949	296.509 6531	293.937 8115	292.693 7989	294.090 7281	293.191 0368
	0.59666 8956	27503.1 8684	296.65	307.896 9429	300.265 8863	296.863 6878	295.547 3248	297.114 9193	295.832 0459
Gas D	0.64944 1297	22994.0 1557	295.538 8889	305.283 6295	299.813 5921	296.251 1707	296.376 7586	297.015 6098	295.948 5691
	0.64944 1297	1206.58 2526	276.761 1111	276.919 0502	276.573 0092	278.470 2865	274.895 9915	277.361 0518	277.060 9988
	0.64944 1297	2185.63 8062	281.594 4444	280.894 4977	281.257 6129	281.966 366	280.010 038	281.671 5233	281.322 5158

	0.64944 1297	3516.32 622	284.927 7778	284.598 6005	285.007 0014	284.796 0991	283.817 8735	284.881 7901	284.545 0676
	0.64944 1297	5446.85 8262	288.705 5556	288.480 8531	288.457 653	287.425 4577	287.098 1909	287.748 5528	287.379 1672
	0.64944 1297	8204.76 1179	291.205 5556	292.581 2554	291.687 9458	289.908 9005	289.974 3421	290.395 1677	289.928 3222
Gas D 2nd Series	0.64944 1297	11996.8 7769	293.205 5556	296.836 4935	294.683 7085	292.231 2104	292.473 4179	292.836 2402	292.209 8247
	0.64944 1297	6170.80 7777	289.15	289.679 5638	289.441 6284	288.179 6744	287.994 2309	288.557 3239	288.165 8933
	0.64944 1297	9307.92 2346	292.038 8889	293.943 3756	292.682 6489	290.677 9387	290.822 0827	291.206 4738	290.694 3636
	0.64944 1297	14065.3 0488	293.927 7778	298.759 4697	295.937 9338	293.208 9996	293.471 6113	293.857 1738	293.142 8537
	0.64944 1297	18202.1 5925	294.983 3333	302.067 6782	297.970 9166	294.800 8593	295.029 3046	295.512 7762	294.628 808
	0.64944 1297	26544.8 1558	296.872 2222	307.368 6861	300.945 8819	297.145 8971	297.174 3566	297.940 6419	296.747 2448
Sun, C.- Y., Chen, G.-J., Lin, W., & Guo, T.-M. (2003).	0.65610 6473	22545.8 5635	295.927 7778	305.004 4319	299.775 9013	296.189 224	296.471 1798	297.015 6625	295.972 0199
	0.65610 6473	1044	274.2	276.048 3512	275.578 4939	277.681 894	273.932 1215	276.337 5223	276.103 8958
	0.65610 6473	1580	277.2	278.640 1047	278.841 828	280.108 7344	277.589 5595	279.495 8364	279.155 5645
	0.65610 6473	2352	280.2	281.433 7113	281.975 026	282.458 7521	280.919 9143	282.296 8242	281.957 33
	0.65610 6473	3126	282.2	283.634 7791	284.215 5127	284.151 2827	283.192 5121	284.220 5983	283.890 2997
	0.65610 6473	3964	284.2	285.614 0253	286.085 939	285.572 0236	285.020 2147	285.794 7478	285.462 0463
	0.65610 6473	5121	286.2	287.903 8431	288.102 8238	287.111 9724	286.920 133	287.470 2827	287.116 479
	0.65610 6473	6358	288.2	289.973 0063	289.806 7994	288.419 476	288.467 9555	288.873 8738	288.483 2001
	0.65610 6473	7212	289.2	291.238 4309	290.799 3678	289.183 8415	289.345 3616	289.687 9742	289.266 7551
82.45% CH4 + 10.77% CO2 + 6.78% H2S	0.70005 9936	8220	290.2	292.600 9384	292.636 7861	290.355 715	291.701 6614	291.319 6163	290.903 7843
	0.70005 9936	1114	276.2	276.434 3319	277.017 3062	278.421 4298	276.567 1193	277.607 1795	277.385 854
	0.70005 9936	1385	278.2	277.782 7857	278.719 0024	279.697 4556	278.433 4697	279.282 0943	278.999 9371
	0.70005 9936	1815	280.2	279.578 0217	280.832 1069	281.290 1209	280.677 0117	281.252 24	280.950 3399
	0.70005 9936	2265	282.2	281.155 2568	282.563 081	282.601 5261	282.453 732	282.803 3445	282.506 0507
	0.70005 9936	3110	284.2	283.593 4759	285.040 8615	284.489 3668	284.901 2324	284.957 6229	284.671 2317
	0.70005 9936	4065	286.2	285.831 6576	287.133 691	286.093 7314	286.880 647	286.736 3563	286.446 7962
	0.70005 9936	4570	287.2	286.865 3437	288.048 842	286.798 1269	287.720 9256	287.505 9769	287.208 676

	0.70005 9936	4890	288.2	287.478 6965	288.577 7649	287.206 0318	288.199 5612	287.949 0123	287.645 111
	0.70005 9936	6110	289.2	289.582 873	290.318 4703	288.552 5628	289.738 4896	289.399 7352	289.061 8221
	0.70005 9936	6862	290.2	290.733 5077	291.225 5929	289.256 7721	290.518 4048	290.152 2567	289.788 5142
	0.70005 9936	7650	290.9	291.846 1976	292.075 1531	289.917 8535	291.235 1252	290.855 4313	290.462 1116
82.91% CH4 + 7.16% CO2 + 9.93% H2S	0.68484 0337	8024	291.2	292.345 7597	292.174 0948	290.081 0521	291.072 7759	290.896 398	290.473 2394
	0.68484 0337	1192	278.2	276.844 5103	277.233 1122	278.694 8568	276.486 7396	277.882 8004	277.621 77
	0.68484 0337	1932	282.2	280.012 8417	281.017 1114	281.535 5892	280.566 5072	281.435 8294	281.119 4034
	0.68484 0337	2460	284.2	281.769 3149	282.910 2396	282.967 6512	282.509 4287	283.110 6701	282.801 8031
	0.68484 0337	3303	286.2	284.081 9143	285.219 1582	284.724 1025	284.790 4469	285.098 151	284.797 5434
	0.68484 0337	4212	288.2	286.141 6043	287.124 0463	286.181 4028	286.598 9915	286.706 7383	286.400 2557
Deaton, W. M., & Frost, E. M.	0.80260 126	3103	282.65	283.575 358	286.848 6081	285.254 6811	287.706 9478	286.575 2658	286.382 38
(Sample A)	0.80260 126	627.422 9137	274.816 6667	273.255 3837	274.559 3745	275.838 0225	275.305 8196	274.025 8585	274.581 9006
	0.80260 126	1261.74 0585	280.316 6667	277.195 3031	279.930 3882	279.914 7179	281.078 3389	280.172 9961	280.001 7208
	0.80260 126	1806.42 6411	283.15	279.545 379	282.689 2768	282.032 1307	283.830 6257	282.829 9511	282.620 2916
	0.80260 126	2571.74 447	285.927 7778	282.105 642	285.404 925	284.131 999	286.398 6989	285.299 6314	285.100 4782
	0.80260 126	3592.16 855	288.705 5556	284.776 8884	287.974 0231	286.132 9376	288.699 3494	287.556 9777	287.364 9595
	0.80260 126	4295.43 3794	290.038 8889	286.314 0949	289.348 6032	287.209 3078	289.878 8578	288.744 3716	288.545 9527
	0.80260 126	5364.12 1174	291.483 3333	288.336 7369	291.056 7142	288.552 4944	291.294 5977	290.206 4203	289.985 3378
	0.80260 126	6301.80 8166	292.538 8889	289.885 5763	292.295 2757	289.530 3741	292.286 5267	291.259 7609	291.010 2887
(Sample B)	0.65083 6699	599.843 8845	273.705 5556	273.027 7412	271.094 3005	274.424 96	268.450 7045	271.156 9537	271.686 6181
	0.65083 6699	737.739 0304	275.372 2222	274.100 2428	272.725 4572	275.620 023	270.437 0702	273.238 3394	273.333 4998
	0.65083 6699	992.845 0502	277.594 4444	275.754 4106	275.066 5869	277.344 3466	273.204 0817	275.835 5775	275.625 613
	0.65083 6699	1337.58 2915	280.372 2222	277.561 389	277.416 0756	279.085 6725	275.881 5066	278.171 277	277.846 9188
(Sample C)	0.69338 0674	723.949 5158	273.705 5556	273.999 8028	273.508 0409	275.859 9636	272.406 3859	273.783 6992	273.951 2778
	0.69338 0674	806.686 6033	274.816 6667	274.583 2443	274.354 7127	276.487 5641	273.397 6997	274.784 3639	274.800 2782
	0.69338 0674	903.213 2054	275.927 7778	275.212 1378	275.239 0168	277.144 5843	274.419 0503	275.773 7559	275.675 2227
	0.69338 0674	972.160 7783	276.483 3333	275.632 4792	275.814 5763	277.573 052	275.076 1098	276.392 6684	276.238 3862
	0.69338 0674	1172.10 874	277.594 4444	276.741 7755	277.277 9809	278.665 45	276.719 397	277.895 7625	277.648 6092

	0.69338 0674	1241.05 6313	278.15	277.092 7396	277.725 193	279.000 1404	277.213 7521	278.338 3746	278.073 5337
	0.69338 0674	1461.68 8546	279.816 6667	278.129 6526	279.005 4428	279.960 4948	278.608 6975	279.571 0506	279.274 8922
(Sample D	0.67926 2806	2213.21 7091	283.15	280.985 8702	281.970 1303	282.293 9866	281.436 5067	282.287 3791	281.968 9268
	0.67926 2806	751.528 545	273.705 5556	274.199 3493	273.498 7181	275.963 0764	272.074 569	273.900 5059	273.996 4524
	0.67926 2806	882.528 9335	274.816 6667	275.081 5741	274.758 9858	276.895 8397	273.550 1757	275.338 1366	275.246 2842
	0.67926 2806	1089.37 1652	276.483 3333	276.300 5617	276.410 4923	278.122 9455	275.440 0275	277.079 8732	276.848 2992
	0.67926 2806	1399.63 5731	278.705 5556	277.850 0258	278.376 0898	279.590 5176	277.624 478	279.008 7444	278.704 8339
	0.67926 2806	1675.42 6022	280.372 2222	279.032 2521	279.786 7363	280.648 5179	279.148 7653	280.325 7283	280.005 3901
(Sample E	0.65278 5123	2096.00 6217	282.038 8889	280.591 7223	280.997 8899	281.747 4635	279.819 5176	281.441 7643	281.094 9103
	0.65278 5123	944.581 7492	275.372 2222	275.466 9841	274.716 9844	277.070 8273	272.848 2716	275.457 1219	275.283 0581
	0.65278 5123	1716.79 4566	280.316 6667	279.197 2994	279.425 1961	280.569 1109	278.140 845	280.035 1535	279.687 486
	0.65278 5123	3454.27 3404	285.927 7778	284.450 6722	284.934 6786	284.718 8769	283.825 695	284.824 9188	284.491 7976
(Sample F	0.63879 8761	5253.80 5057	289.261 1111	288.142 0884	287.960 564	287.112 8303	286.388 9352	287.315 6098	286.940 1576
	0.63879 8761	765.318 0595	273.705 5556	274.297 1639	272.741 1495	275.729 5604	270.125 8525	273.366 3846	273.395 37
	0.63879 8761	1241.05 6313	277.594 4444	277.092 7396	276.560 4021	278.543 0127	274.598 2557	277.386 0948	277.066 9076
	0.63879 8761	1730.58 4081	280.372 2222	279.251 6828	279.187 2464	280.494 7223	277.522 1736	279.838 9109	279.476 6206
	0.63879 8761	1730.58 4081	280.372 2222	279.251 6828	279.187 2464	280.494 7223	277.522 1736	279.838 9109	279.476 6206
	0.63879 8761	3461.16 8161	285.927 7778	284.467 2021	284.663 3701	284.607 4892	283.218 8708	284.580 3895	284.232 5882
	0.63879 8761	4909.06 7193	288.705 5556	287.514 3261	287.424 3762	286.703 9166	285.886 7239	286.874 2225	286.507 2379
	0.63879 8761	4895.27 7678	288.705 5556	287.488 5684	287.402 1529	286.686 981	285.865 7974	286.855 905	286.489 2344
	0.63879 8761	8652.92 0403	292.038 8889	293.149 5264	291.902 4118	290.136 9653	289.922 3836	290.536 5994	290.040 2671
(Sample G	0.63983 6517	9390.65 9433	292.705 5556	294.040 7802	292.568 8134	290.645 3401	290.515 5576	291.082 0039	290.556 2268
	0.63983 6517	758.423 3022	273.705 5556	274.248 4156	272.693 4567	275.686 1138	270.097 4173	273.301 4383	273.344 2253
	0.63983 6517	1234.16 1555	277.594 4444	277.058 2802	276.539 4576	278.519 4691	274.602 6319	277.362 2195	277.045 6535
	0.63983 6517	1723.68 9323	280.372 2222	279.224 53	279.178 2721	280.480 3142	277.539 9183	279.829 3806	279.468 2749
	0.63983 6517	3468.06 2918	285.927 7778	284.483 7084	284.700 6004	284.628 6237	283.281 7325	284.612 8857	284.266 1719
	0.63983 6517	5033.17 2824	288.705 5556	287.743 8484	287.642 5675	286.863 5873	286.116 7795	287.056 1945	286.687 4023
	0.63983 6517	5033.17 2824	288.705 5556	287.743 8484	287.642 5675	286.863 5873	286.116 7795	287.056 1945	286.687 4023
(Sample H	0.69136 0127	7729.02 2926	291.483 3333	291.953 1824	291.999 0973	289.907 9127	291.034 4827	290.769 8117	290.364 7951
	0.69136 0127	758.423 3022	274.205 5556	274.248 4156	273.829 2208	276.113 4723	272.739 6034	274.187 6397	274.282 2228
	0.69136 0127	944.581 7492	275.372 2222	275.466 9841	275.547 207	277.389 1585	274.728 1846	276.118 9734	275.983 0236

	0.69136 0127	1254.84 5827	277.594 4444	277.161 2497	277.770 2187	279.048 6015	277.221 1251	278.389 2006	278.119 4584
	0.69136 0127	1758.16 311	280.372 2222	279.359 527	280.409 8594	281.031 9488	280.063 7651	280.878 4887	280.568 5908
(Sample I	0.63218 6651	2130.48 0004	282.038 8889	280.709 25	280.687 0584	281.663 3233	278.960 3464	281.177 9149	280.810 1003
	0.63218 6651	792.897 0887	273.705 5556	274.489 0746	272.868 684	275.877 2103	270.091 7834	273.570 1498	273.545 543
	0.63218 6651	972.160 7783	275.316 6667	275.632 4792	274.480 9735	277.060 6013	272.010 3336	275.327 2739	275.111 761
	0.63218 6651	1310.00 3886	277.594 4444	277.430 0264	276.840 2607	278.801 4298	274.733 7839	277.676 4989	277.337 2043
(Sample J	0.62942 92	1813.32 1168	280.372 2222	279.571 6385	279.351 4991	280.686 9834	277.447 3912	279.998 1435	279.625 254
	0.62942 92	882.528 9335	273.705 5556	275.081 5741	273.652 3485	276.474 201	270.950 9644	274.462 5963	274.320 1983
	0.62942 92	951.476 5065	274.261 1111	275.508 6784	274.247 6762	276.911 2902	271.656 7831	275.098 5851	274.895 1382
	0.62942 92	1020.42 4079	274.816 6667	275.914 1922	274.801 3373	277.318 408	272.307 5055	275.673 7432	275.425 2713
	0.62942 92	1027.31 8837	274.816 6667	275.953 6542	274.854 6313	277.357 6277	272.369 8527	275.728 352	275.476 0714
	0.62942 92	1172.10 874	275.927 7778	276.741 7755	275.898 1257	278.126 6659	273.580 3602	276.773 9972	276.462 7727
	0.62942 92	1441.00 4274	277.594 4444	278.037 4033	277.532 6763	279.335 5912	275.437 3251	278.337 3949	277.978 7892
	0.62942 92	1454.79 3789	277.594 4444	278.099 0071	277.608 0495	279.391 4644	275.521 8003	278.407 6813	278.047 8512
	0.62942 92	2027.05 8644	280.372 2222	280.352 4556	280.233 3364	281.344 5432	278.400 6249	280.780 8677	280.408 8589
	0.62942 92	1992.58 4858	280.372 2222	280.230 6345	280.097 5848	281.243 2171	278.254 7894	280.661 1821	280.288 837
	0.62942 92	2006.37 4372	280.372 2222	280.279 5424	280.152 1652	281.283 9519	278.313 4638	280.709 3366	280.337 1194
	0.62942 92	2840.64 0005	283.15 2222	282.874 1545	282.903 9076	283.345 32	281.202 4614	283.087 1035	282.726 5826
	0.62942 92	3764.53 7482	285.372 2222	285.172 3005	285.132 491	285.025 8489	283.442 8163	284.959 62	284.600 4855
	0.62942 92	4033.43 3017	285.927 7778	285.764 0573	285.678 5093	285.439 108	283.978 1526	285.413 3256	285.051 6708
	0.62942 92	4026.53 8259	285.927 7778	285.749 2419	285.664 9694	285.428 8529	283.964 942	285.402 0948	285.040 5192
	0.62942 92	4054.11 7288	285.927 7778	285.808 3949	285.718 9909	285.469 7707	284.017 6296	285.446 8972	285.085 0004
	0.62942 92	4364.38 1367	286.483 3333	286.454 8379	286.302 6061	285.912 1954	284.583 5004	285.929 9583	285.563 6687
	0.62942 92	4674.64 5445	287.038 8889	287.069 2079	286.846 1235	286.324 8395	285.105 0069	286.378 3758	286.006 3908
	0.62942 92	4681.54 0202	287.038 8889	287.082 5249	286.857 7876	286.333 7016	285.116 1407	286.387 9849	286.015 8599
	0.62942 92	5067.64 661	287.594 4444	287.806 8855	287.484 9772	286.810 6254	285.711 2248	286.903 8611	286.523 0575
	0.62942 92	5433.06 8747	288.15 2222	288.456 943	288.036 0188	287.230 3007	286.228 2449	287.355 8997	286.965 5349
	0.62942 92	5812.28 0398	288.705 5556	289.099 2725	288.569 9759	287.637 5503	286.724 0477	287.792 9771	287.391 5337
	0.62942 92	6984.38 9138	289.816 6667	290.912 1216	290.023 8369	288.749 3403	288.048 1425	288.979 1789	288.537 9803
(Sample K	0.71862 1772	8384.02 4868	290.927 7778	292.811 1645	293.116 4505	290.628 2836	292.347 6373	291.766 1373	291.361 7753
	0.71862 1772	1068.68 738	274.261 1111	276.186 533	277.068 1897	278.324 3719	276.978 527	277.579 3923	277.394 3368

	0.71862 1772	1220.37 2041	275.372 2222	276.988 9465	278.102 2216	279.101 2044	278.112 2586	278.625 0604	278.388 8151
	0.71862 1772	2247.69 0878	280.372 2222	281.098 948	282.860 4391	282.703 9273	283.074 5829	283.059 82	282.779 9093
	0.71862 1772	3164.69 3598	283.15	283.734 0436	285.526 0859	284.742 5293	285.671 7121	285.386 3796	285.118 1075
(Sample L	0.58220 0927	4591.90 8357	285.983 3333	286.908 2985	285.691 6772	285.771 3466	282.641 1782	285.330 121	284.906 9284
	0.58220 0927	1261.74 0585	273.705 5556	277.195 3031	275.374 42	278.123 1858	271.513 3096	276.438 3093	276.053 697
	0.58220 0927	1427.21 476	274.816 6667	277.975 3765	276.358 6431	278.843 8688	272.655 6292	277.360 5318	276.951 8558
	0.58220 0927	1420.32 0002	274.816 6667	277.944 2024	276.319 9665	278.815 5132	272.611 0615	277.324 7811	276.916 7951
	0.58220 0927	2027.05 8644	277.594 4444	280.352 4556	279.160 8672	280.906 0005	275.814 6824	279.866 9268	279.443 1162
	0.58220 0927	2854.42 9519	280.372 2222	282.912 1374	281.894 6071	282.932 43	278.763 4635	282.198 0855	281.785 4216
	0.58220 0927	4047.22 2531	283.15	285.793 6337	284.683 2367	285.014 6099	281.636 0463	284.506 6342	284.092 1019
	0.58220 0927	7425.65 3605	287.705 5556	291.538 0972	289.530 4484	288.670 3988	286.303 7468	288.431 8314	287.920 978
	0.58220 0927	10328.3 4643	289.816 6667	295.104 0339	292.165 6745	290.677 5363	288.668 0071	290.544 3727	289.914 6725
Salufu, S., & Nwakwo, P. (2013).	1	413.685 4376	274.816 6667	271.253 6889	274.706 0118	274.641 0437	274.724 7618	270.995 8308	273.769 6144
	1	1344.47 7672	283.15	277.593 9266	283.526 28	281.523 7193	283.147 3649	283.249 724	283.214 1517
	1	3033.69 3209	288.705 5556	283.394 3761	289.616 0309	286.376 1003	288.053 1801	289.067 7212	289.027 0648
	0.7	827.370 8752	274.816 6667	274.722 3642	274.691 3651	276.687 1765	273.928 8462	275.119 7329	275.112 8983
	0.7	2344.21 748	283.15	281.409 1053	282.830 5762	282.805 1297	282.722 2826	283.039 2059	282.743 189
	0.7	4757.38 2532	288.705 5556	287.228 0938	288.361 7803	287.039 7587	288.003 7402	287.768 1427	287.467 0613
	0.7	23442.1 748	297.038 8889	305.558 963	300.825 8295	296.815 9826	297.845 3803	298.072 4699	297.039 8097
	0.65	758.423 3022	274.816 6667	274.248 4156	272.924 5695	275.773 0219	270.653 8414	273.481 536	273.534 8349
	0.65	2688.95 5344	283.15	282.447 3749	282.903 3373	283.201 0691	281.726 2483	283.097 7481	282.757 0828
	0.65	14134.2 5245	294.261 1111	298.819 9783	295.986 7611	293.244 1539	293.521 1015	293.899 1247	293.182 4881
	0.6	1723.68 9323	277.594 4444	279.224 53	278.284 6823	280.119 9433	275.409 8623	279.078 4796	278.674 7725
	0.6	3309.48 3501	283.15	284.097 9641	283.476 333	283.983 6623	280.957 4811	283.539 3898	283.149 3622
	0.6	6756.86 2147	288.705 5556	290.578 2424	289.156 9754	288.272 3823	286.480 7394	288.188 2898	287.722 6879
	0.6	18098.7 3789	294.261 1111	301.991 9093	296.998 5372	294.299 1482	293.166 1382	294.499 8031	293.566 319
Chen, G.J., and Guo, T.M.:	0.63109 0548	841.999 781	274	274.819 2621	273.318 6907	276.215 9487	270.600 5108	274.084 8923	273.990 1979
	0.63109 0548	940.999 7552	274.612	275.445 2369	274.198 1771	276.861 5075	271.645 9418	275.036 4764	274.842 8779
	0.63109 0548	989.999 7425	275.113	275.737 7463	274.599 7879	277.156 7986	272.118 72	275.457 0341	275.228 5223

	0.63109 0548	1113.99 971	276.449	276.434 3304	275.533 4245	277.844 4884	273.206 6313	276.406 2571	276.116 2332
	0.63109 0548	1435.99 9626	278.174	278.014 9411	277.542 2327	279.329 9093	275.494 4138	278.342 3509	277.985 9524
	0.63109 0548	2004.99 9478	281.234	280.274 6768	280.183 0116	281.294 7321	278.391 9137	280.735 483	280.365 009
	0.63109 0548	2846.99 9259	284.349	282.891 6873	282.957 0164	283.373 565	281.301 0025	283.133 324	282.774 6416
	0.63109 0548	3761.99 9021	286.739	285.166 5722	285.161 8805	285.036 8381	283.514 8725	284.986 5749	284.629 2434
	0.63109 0548	4034.99 895	287.461	285.767 4196	285.716 1373	285.456 4835	284.057 6906	285.447 3234	285.087 4383
	0.63109 0548	4306.99 888	288.072	286.337 815	286.232 2578	285.847 8101	284.558 2094	285.874 8823	285.511 2987
	0.63109 0548	4628.99 8796	288.627	286.980 6872	286.802 6835	286.280 9356	285.105 8384	286.345 9336	285.976 6539
	0.63109 0548	5024.99 8693	289.182	287.728 8563	287.452 11	286.774 8455	285.722 2126	286.880 5274	286.502 5537
	0.63109 0548	5370.99 8603	289.904	288.348 7768	287.978 9378	287.176 141	286.216 6752	287.313 0292	286.926 1677
	0.63109 0548	5816.99 8487	290.403	289.107 0729	288.610 0563	287.657 6164	286.802 4734	287.829 9408	287.430 1514
	0.63109 0548	7029.99 8171	291.622	290.978 1107	290.108 5435	288.804 0365	288.164 761	289.052 9675	288.611 8942
	0.63109 0548	8365.99 7824	292.839	292.788 2043	291.485 089	289.861 1909	289.380 7292	290.172 532	289.679 5653
Jossang, A. and Stange, E.: "A	0.70313 0909	4758.99 8762	289.631	287.231 1779	288.422 2536	287.067 4273	288.109 5248	287.824 0522	287.526 0014
	0.70313 0909	30075.9 9218	299.688	309.252 8046	302.823 475	298.399 9246	299.190 3323	299.739 8943	298.481 3266
	0.70313 0909	59961.9 844	304.483	320.961 7117	308.212 9958	302.755 1126	302.356 7239	304.230 9654	302.173 9016
	0.70313 0909	4758.99 8762	287.522	287.231 1779	288.422 2536	287.067 4273	288.109 5248	287.824 0522	287.526 0014
	0.70313 0909	29694.9 9228	296.891	309.057 3892	302.723 8928	298.320 046	299.126 8021	299.657 3211	298.411 2381
	0.70313 0909	59771.9 8445	301.686	320.902 4125	308.188 2057	302.734 9353	302.343 3821	304.210 1973	302.157 352
	0.70313 0909	66814.9 8262	303.176	323.016 1226	309.058 2849	303.443 9205	302.804 8923	304.939 7856	302.735 9051
Sloan, E.D. Jr.:	0.66840 8755	1254.99 9674	278.8	277.162 011	277.291 9023	278.860 9641	276.177 1573	277.997 5504	277.705 3542
	0.66840 8755	1923.99 9499	282.9	279.983 7177	280.649 487	281.374 3551	279.822 1813	281.121 0793	280.788 3419
	0.66840 8755	4122.99 8927	288.7	285.954 8895	286.638 8178	285.914 1743	285.815 7942	286.275 5243	285.953 6453
	0.66840 8755	6963.99 8188	292.2	290.882 5194	290.757 8805	289.078 798	289.559 6298	289.687 007	289.287 7182
	0.66840 8755	9679.99 7482	294.1	294.376 6612	293.345 6518	291.084 8408	291.754 0326	291.809 2748	291.301 3167
	0.66840 8755	12269.9 9681	295.1	297.103 3633	295.208 7976	292.537 767	293.258 6414	293.333 4861	292.715 124
	0.66840 8755	16839.9 9562	296.1	301.043 9093	297.696 7057	294.489 2121	295.169 4666	295.368 316	294.558 7137
	0.66840 8755	20679.9 9462	297.1	303.797 4265	299.310 8615	295.762 2677	296.349 0717	296.689 9663	295.729 0035
	0.66840 8755	27319.9 9289	298.3	307.796 9951	301.498 9964	297.496 8015	297.872 5673	298.485 1875	297.284 5378
	0.59666 8956	1599.99 9584	277.7	278.723 6669	277.614 2754	279.651 113	274.554 4288	278.479 5318	278.075 5465

	0.59666 8956	3391.99 9118	283.9	284.300 2932	283.598 4521	284.098 9347	280.982 4274	283.636 4856	283.242 735
	0.59666 8956	6963.99 8188	289.2	290.882 5194	289.327 1338	288.423 1173	286.542 6112	288.315 4487	287.838 5605
	0.59666 8956	10500.9 9727	292.1	295.292 2006	292.598 0245	290.921 5389	289.456 9818	290.948 7281	290.328 3671
	0.59666 8956	13779.9 9982	293.3	298.506 8065	294.762 1892	292.586 4895	291.281 2405	292.686 943	291.926 9687
	0.59666 8956	1923.99 9499	279.1	279.983 7177	279.082 8339	280.736 1495	276.190 5474	279.790 0774	279.381 7609
	0.59666 8956	2647.99 9311	281.8	282.329 199	281.626 4789	282.625 4781	278.934 1679	281.983 011	281.584 3807
	0.59666 8956	4819.99 8746	286.7	287.347 0317	286.396 5898	286.202 8936	283.770 778	285.938 3249	285.526 991
	0.59666 8956	8135.99 7883	290.3	292.492 1119	290.565 8703	289.366 7737	287.668 5891	289.314 2709	288.792 2907
	0.59666 8956	9480.99 7534	291.2	294.146 4352	291.784 2712	290.297 9504	288.749 6078	290.294 6679	289.717 4303
	0.59666 8956	17159.9 9554	294.3	301.289 5901	296.509 1633	293.937 4319	292.693 4106	294.090 3344	293.190 6857
	0.59666 8956	20264.9 9473	295	303.518 4585	297.833 667	294.965 8318	293.728 1001	295.155 3525	294.133 5342
	0.59666 8956	22989.9 9402	295.6	305.281 1415	298.838 4207	295.748 364	294.492 3081	295.964 1567	294.840 2599
	0.59666 8956	27499.9 9285	296.7	307.895 2044	300.264 9614	296.862 9638	295.546 6529	297.114 1729	295.831 4079
	0.64944 1297	1206.99 9686	276.8	276.921 1731	276.575 7349	278.472 308	274.899 0822	277.363 7207	277.063 561
	0.64944 1297	2185.99 9431	281.6	280.895 7002	281.258 9165	281.967 3449	280.011 406	281.672 6659	281.323 6639
	0.64944 1297	3515.99 9085	284.9	284.597 8256	285.006 2678	284.795 5426	283.817 1533	284.881 1738	284.544 4522
	0.64944 1297	5446.99 8583	288.7	288.481 0962	288.457 8562	287.425 6132	287.098 3777	287.748 72	287.379 3306
	0.64944 1297	8204.99 7866	291.2	292.581 5613	291.688 1733	289.909 0761	289.974 538	290.395 3534	289.928 4983
	0.64944 1297	11999.9 9688	293.2	296.839 5657	294.685 7584	292.232 8058	292.475 0724	292.837 9088	292.211 36
	0.64944 1297	6170.99 8395	289.2	289.679 8658	289.441 872	288.179 8614	287.994 4505	288.557 5238	288.166 0869
	0.64944 1297	9307.99 7579	292.1	293.943 4644	292.682 7126	290.677 988	290.822 1365	291.206 5257	290.694 4124
	0.64944 1297	14069.9 9634	293.9	298.763 5937	295.940 5634	293.211 053	293.473 6743	293.859 3144	293.144 7966
	0.64944 1297	18199.9 9527	295	302.066 096	297.969 9791	294.800 1233	295.028 6035	295.512 0123	294.628 1301
	0.64944 1297	26549.9 9309	296.9	307.371 5763	300.947 4197	297.147 1141	297.175 4241	297.941 8992	296.748 3233
	0.64944 1297	22549.9 9413	295.9	305.007 0278	299.659 841	296.129 8858	296.266 6708	296.890 1031	295.839 4065
	0.80260 126	626.999 8369	274.8	273.251 9461	274.554 1887	275.834 1152	275.299 9816	274.018 7385	274.576 4423
	0.80260 126	1261.99 9672	280.3	277.196 5801	279.931 9667	279.915 9248	281.079 9549	280.174 5767	280.003 2492
	0.80260 126	1805.99 953	283.2	279.543 7509	282.687 4598	282.030 731	283.828 8606	282.828 259	282.618 6008
	0.80260 126	2570.99 9331	285.9	282.103 4343	285.402 6972	284.130 27	286.396 6495	285.297 6462	285.098 481
	0.80260 126	3591.99 9066	288.7	284.776 493	287.973 6603	286.132 6541	288.699 0334	287.556 6627	287.364 6451
	0.80260 126	4295.99 8882	290.1	286.315 255	289.349 6145	287.210 1012	289.879 7124	288.745 2412	288.546 814

	0.80260 126	5363.99 8605	291.5	288.336 5223	291.056 5385	288.552 3559	291.294 4549	290.206 2705	289.985 1914
	0.80260 126	6301.99 8361	292.6	289.885 8732	292.295 5077	289.530 5576	292.286 7098	291.259 9578	291.010 4792
	0.80260 126	8535.99 7779	294	293.003 3354	294.628 2305	291.381 3176	294.075 858	293.233 772	292.900 0585
	0.65083 6699	599.999 8439	273.7	273.029 0494	271.096 3498	274.426 4582	268.453 2302	271.159 7715	271.688 7141
	0.65083 6699	737.999 808	275.4	274.102 1293	272.728 2432	275.622 0686	270.440 4218	273.241 6611	273.336 2766
	0.65083 6699	992.999 7417	277.6	275.755 3156	275.067 815	277.345 254	273.205 5073	275.836 8564	275.626 7943
	0.65083 6699	1337.99 9652	280.4	277.563 3591	277.418 5312	279.087 4982	275.884 2529	278.173 6202	277.849 2012
	0.65083 6699	1778.99 9537	282.6	279.440 2057	279.664 2055	280.762 111	278.350 1368	280.254 1174	279.903 6954
	0.65083 6699	2088.99 9457	283.7	280.567 6661	280.930 4845	281.710 8095	279.700 449	281.383 328	281.034 3821
	0.65083 6699	2247.99 9415	284.3	281.099 9544	281.508 7457	282.145 1094	280.307 4581	281.890 9274	281.544 3313
	0.65083 6699	2667.99 9306	285.9	282.387 0697	282.859 0112	283.161 8275	281.701 3485	283.060 0463	282.719 9685
	0.65083 6699	2860.99 9256	286.5	282.930 1875	283.409 5766	283.577 4411	282.260 2562	283.531 1846	283.193 4005
	0.69338 0674	723.999 8117	273.7	274.000 1717	273.508 5845	275.860 3661	272.407 0266	273.784 3612	273.951 8264
	0.69338 0674	806.999 7901	274.8	274.585 3697	274.357 7498	276.489 8179	273.401 2319	274.787 8519	274.803 3035
	0.69338 0674	902.999 7651	275.9	275.210 8015	275.237 1677	277.143 2088	274.416 9296	275.771 738	275.673 4054
	0.69338 0674	971.999 7471	276.5	275.631 5227	275.813 2802	277.572 0864	275.074 637	276.391 2943	276.237 1235
	0.69338 0674	1171.99 9695	277.6	276.741 2089	277.277 253	278.664 9056	276.718 5893	277.895 0364	277.647 9152
	0.69338 0674	1240.99 9677	278.2	277.092 4571	277.724 8359	278.999 8731	277.213 3588	278.338 024	278.073 1955
	0.69338 0674	1461.99 962	279.8	278.131 0328	279.007 1077	279.961 7458	278.610 492	279.572 6242	279.276 4402
	0.69338 0674	2212.99 9424	283.2	280.985 1522	282.250 531	282.409 569	282.009 9633	282.529 5334	282.224 9227
	0.67926 2806	751.999 8044	273.7	274.202 7132	273.503 6347	275.966 7093	272.080 3821	273.906 3466	274.001 3759
	0.67926 2806	882.999 7703	274.8	275.084 5703	274.763 1691	276.898 9412	273.555 0255	275.342 7307	275.250 3928
	0.67926 2806	1088.99 9717	276.5	276.298 525	276.407 814	278.120 951	275.437 0029	277.077 1512	276.845 733
	0.67926 2806	1399.99 9636	278.7	277.851 6913	278.378 1288	279.592 044	277.626 7074	279.010 6832	278.706 7326
	0.67926 2806	1674.99 9564	280.4	279.030 5356	279.784 7397	280.647 0176	279.146 6335	280.323 8957	280.003 5674
	0.67926 2806	2095.99 9455	282.1	280.591 6991	281.543 3284	281.971 5795	280.996 1427	281.909 8789	281.589 6754
	0.63879 8761	764.999 801	273.7	274.294 9207	272.737 8635	275.727 152	270.121 8916	273.362 5563	273.392 1191
	0.63879 8761	1240.99 9677	277.6	277.092 4571	276.560 0415	278.542 7457	274.597 8459	277.385 7453	277.066 5706
	0.63879 8761	1730.99 955	280.4	279.253 3165	279.189 1428	280.496 1362	277.524 2398	279.840 625	279.478 3285
	0.63879 8761	3460.99 91	285.9	284.466 797	284.662 9842	284.607 1973	283.218 4883	284.580 0658	284.232 2648
	0.63879 8761	8652.99 7749	292.1	293.149 6227	291.902 4825	290.137 0197	289.922 4444	290.536 6568	290.040 3214

	0.63879 8761	9390.99 7557	292.7	294.041 177	292.549 0943	290.636 1244	290.475 4018	291.062 4856	290.535 5831
	0.63983 6517	757.999 8028	273.7	274.245 411	272.689 0448	275.682 8802	270.092 0964	273.296 2715	273.339 8546
	0.63983 6517	1233.99 9679	277.6	277.057 4695	276.538 4215	278.518 7019	274.601 454	277.361 2134	277.044 6844
	0.63983 6517	1722.99 9552	280.4	279.221 8092	279.175 1106	280.477 9567	277.536 474	279.826 5202	279.465 4255
	0.63983 6517	3467.99 9098	285.9	284.483 5557	284.700 455	284.628 5137	283.281 5886	284.612 7638	284.266 05
	0.63983 6517	5032.99 8691	288.7	287.743 5292	287.642 2942	286.863 3789	286.116 5234	287.055 9694	286.687 1814
	0.63983 6517	7728.99 7989	291.5	291.953 1488	291.030 6175	289.459 184	289.188 9086	289.829 9943	289.371 7783
	0.69136 0127	757.999 8028	274.2	274.245 411	273.824 8492	276.110 2337	272.734 4742	274.182 4562	274.277 837
	0.69136 0127	944.999 7542	275.4	275.469 518	275.550 6698	277.391 7358	274.732 1383	276.122 6828	275.986 4069
	0.69136 0127	1254.99 9674	277.6	277.162 011	277.771 1782	279.049 3199	277.222 1816	278.390 1398	278.120 3655
	0.69136 0127	1757.99 9543	280.4	279.358 891	280.409 1312	281.031 3997	280.062 9985	280.877 8239	280.567 9277
	0.69136 0127	2130.99 9446	282.1	280.711 0105	281.915 1207	282.169 2622	281.627 6638	282.233 2133	281.925 3697
	0.71862 1772	1068.99 9722	274.3	276.188 2666	277.070 4663	278.326 0799	276.981 0448	277.581 7412	277.396 5432
	0.71862 1772	1219.99 9683	275.4	276.987 0665	278.099 8441	279.099 4158	278.109 6746	278.622 7013	278.386 5459
	0.71862 1772	1606.99 9582	277.6	278.752 7378	280.246 3536	280.718 933	280.400 1566	280.684 5217	280.403 8994
	0.71862 1772	2247.99 9415	280.4	281.099 9544	282.861 5084	282.704 7421	283.075 6511	283.060 7695	282.780 864
	0.71862 1772	3164.99 9177	283.2	283.734 824	285.526 8381	284.743 1066	285.672 4265	285.387 0265	285.118 7558
	0.71862 1772	4591.99 8805	286	286.908 4755	288.426 2854	286.977 2109	288.348 2057	287.852 1543	287.572 4142
	0.58220 0927	1261.99 9672	273.7	277.196 5801	275.376 0599	278.124 3849	271.515 227	276.439 8688	276.055 2039
	0.58220 0927	2026.99 9473	277.6	280.352 2477	279.160 634	280.905 8283	275.814 4253	279.866 7241	279.442 9128
	0.58220 0927	2854.99 9257	280.4	282.913 7039	281.896 201	282.933 6158	278.765 1445	282.199 4217	281.786 7633
	0.58220 0927	4046.99 8947	283.2	285.793 1547	284.682 7955	285.014 2792	281.635 6026	284.506 2729	284.091 7432
	0.58220 0927	7424.99 8068	287.7	291.537 1873	289.529 7433	288.669 8636	286.303 0979	288.431 2649	287.920 4366
	0.70312 4418	1764.99 9541	281.6	279.386 0727	280.674 4115	281.149 714	280.570 4392	281.104 4031	280.805 7421
	0.70312 4418	2516.99 9345	283.9	281.942 2052	283.446 7021	283.253 0238	283.394 7942	283.579 2106	283.289 0705
	0.70312 4418	3846.99 8999	287.7	285.356 8997	286.760 3709	285.787 7262	286.585 3113	286.424 8415	286.140 4315
	0.70312 4418	4460.99 884	288.9	286.649 4133	287.917 0326	286.677 8138	287.651 4548	287.400 1196	287.107 7808
	0.70312 4418	5832.99 8483	290.9	289.133 492	290.011 6441	288.296 7448	289.519 5437	289.151 1372	288.824 9555
	0.67368 2203	2185.99 9431	283.3	280.895 7002	281.760 2254	282.174 0804	281.099 9999	282.104 464	281.780 039
	0.67368 2203	2929.99 9238	285.7	283.117 9954	284.059 9819	283.915 2269	283.428 4002	284.100 701	283.786 8991
	0.67368 2203	3577.99 9069	287.2	284.743 785	285.628 6039	285.108 9904	284.961 3436	285.433 7957	285.121 7756

	0.67368 2203	4612.99 88	289.4	286.949 5114	287.623 3167	286.634 2728	286.846 0086	287.106 0537	286.781 6258
	0.67368 2203	5660.99 8527	291	288.846 723	289.230 5698	287.869 2136	288.311 9105	288.440 2715	288.089 8479
	0.61771 8353	13549.9 9648	293.6	298.300 3807	295.043 5642	292.685 966	292.051 9928	293.003 6083	292.276 9042
	0.61771 8353	27679.9 928	297.5	307.992 9552	300.709 2953	297.109 5842	296.365 5717	297.587 0553	296.321 3105
	0.61771 8353	41339.9 8925	300	314.365 8555	303.890 8233	299.622 8658	298.536 8685	300.175 2894	298.492 9946
	0.61771 8353	52159.9 8643	301.7	318.407 079	305.734 8049	301.089 2591	299.712 7129	301.682 2244	299.721 456
	0.61771 8353	62849.9 8365	303.1	321.846 9496	307.213 5335	302.270 3749	300.611 8438	302.894 7381	300.691 2874
	0.61771 8353	20239.9 9473	295.8	303.501 5232	298.226 3127	295.162 7815	294.545 6188	295.575 4848	294.577 9817
	0.61771 8353	33749.9 9122	298.6	311.052 0118	302.281 8965	298.349 2312	297.461 3819	298.864 6774	297.403 2143
	0.62942 92	882.999 7703	273.7	275.084 5703	273.656 5696	276.477 2978	270.955 9913	274.467 1758	274.324 293
	0.62942 92	951.999 7523	274.3	275.511 8338	274.252 0272	276.914 4873	271.661 9183	275.103 1634	274.899 3214
	0.62942 92	1171.99 9695	275.9	276.741 2089	275.897 3894	278.126 1225	273.579 5129	276.773 274	276.462 0817
	0.62942 92	2840.99 9261	283.2	282.875 1457	282.904 9085	283.346 0725	281.203 4874	283.087 9533	282.727 4361
	0.62942 92	3764.99 9021	285.4	285.173 3417	285.133 4613	285.026 5827	283.443 7723	284.960 4278	284.601 2899
	0.62942 92	4363.99 8865	286.5	286.454 0615	286.301 9124	285.911 6691	284.582 8315	285.929 3852	285.563 1018
	0.62942 92	5067.99 8682	287.6	287.807 5277	287.485 527	286.811 0438	285.711 7433	286.904 3127	286.523 5005
	0.62942 92	5432.99 8587	288.2	288.456 8212	288.035 9166	287.230 2228	286.228 1496	287.355 816	286.965 4531
	0.62942 92	5811.99 8488	288.7	289.098 8063	288.569 592	287.637 2573	286.723 6931	287.792 6632	287.391 2284
	0.62942 92	6984.99 8183	289.8	290.913 0048	290.024 527	288.749 8691	288.048 762	288.979 7408	288.538 5199
	0.62942 92	8383.99 7819	290.9	292.811 1301	291.469 331	289.858 9923	289.327 0881	290.154 3721	289.658 8749
Nasrifar, K. and Moshfeg hian, M.:	0.57475 5795	908.199 7637	273.476	275.243 2937	272.561 942	276.138 4242	267.900 6858	273.665 882	273.438 7154
	0.57475 5795	1111.79 9711	274.987	276.422 4675	274.179 8438	277.313 8695	269.851 3223	275.312 4958	274.961 9
	0.57475 5795	1336.89 9652	276.593	277.558 158	275.654 5866	278.389 6649	271.589 376	276.725 7393	276.319 6093
	0.57475 5795	1476.29 9616	277.695	278.194 2551	276.447 9398	278.970 1255	272.508 6057	277.459 8287	277.038 3843
	0.57475 5795	1594.09 9585	278.766	278.699 0936	277.062 0016	279.420 2376	273.212 5208	278.017 6815	277.589 3077
	0.57475 5795	1851.69 9518	279.71	279.716 5164	278.260 1631	280.300 5913	274.566 9699	279.084 1904	278.651 0282
	0.57475 5795	2205.99 9426	280.938	280.962 0362	279.660 5471	281.333 0448	276.118 1188	280.300 537	279.870 4531
	0.57475 5795	2635.69 9314	281.914	282.293 4531	281.084 0559	282.386 4452	277.659 643	281.510 952	281.087 2022

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