

**CORROSION RESISTANT MATERIAL FOR REDUCED FOULING, HEAT
TRANSFER COMPONENT WITH IMPROVED CORROSION AND FOULING
RESISTANCE, AND METHOD FOR REDUCING FOULING**

Cross Reference to Related Application

[0001] This application relates to and claims priority to US Provisional Patent Application No. 60/751,985, filed December 21, 2005, entitled "Corrosion Resistant Material For Reduced Fouling, A Heat Exchanger Having Reduced Fouling And A Method For Reducing Heat Exchanger Fouling in a Refinery" and US Provisional Patent Application No. 60/815,844, filed June 23, 2006 entitled "A Method of Reducing Heat Exchanger Fouling in a Refinery" and US Provisional Patent Application No. 60/872,493, filed December 4, 2006 entitled "An Insert and Method For Reducing Fouling In A Process Stream," the disclosures of which are hereby incorporated herein specifically by reference.

Field of the Invention

[0002] This invention relates to the reduction of sulfidation or sulfidic corrosion and the reduction of depositional fouling in general and in particular the reduction of sulfidation/sulfidic corrosion and the reduction of depositional fouling in heat transfer components, which include but are not limited to heat exchangers, furnaces and furnace tubes located in refining facilities and petrochemical processing facilities and other components used for transporting or conveying process streams, which may be prone to fouling. In particular, the present invention relates to the reduction of corrosion and fouling associated with process streams. The present invention is directed to a method of reducing fouling in a heat transfer component, which combines the use of a corrosion resistant material having the desired surface roughness with the application of vibration, pulsation and internal turbulence promoters.

Background of the Invention

[0003] Heat transfer components are used in refinery and petrochemical processing applications at various locations within the facilities to adjust the temperature (i.e., heat or cool) of the processed fluid (e.g., crude oil or derivatives thereof). The heat transfer components (e.g., a heat exchanger, furnaces, and furnace tubes) may be near the furnace to pre-heat the temperature of the oil prior to entry into the furnace (i.e., late-train). A typical tube-in-shell heat exchanger includes a plurality of tubes through which the oil may flow through and

around. A hot fluid and a cold fluid enter separate chambers or tubes of the heat exchanger unit. The hot fluid transfers its heat to the cold fluid. The heat exchanger is designed to efficiently transfer heat from one fluid to another. The hot and cold fluids are never combined. Heat transfer occurs through the tube wall that separates the hot and cold liquids. By employing the correct flow rate and maximizing the area of the partition, heat exchanger performance can be optimally controlled. A variety of other heat exchanger designs, such as spiral heat exchangers, tube-in-tube heat exchangers and plate-and-frame heat exchangers operate essentially on the same principles.

[0004] During normal use with contact between the oil and the heat exchanger, corrosion and the build-up of deposits occurs. This build-up of deposits is often called fouling. Fouling adversely impacts the optimal control of the heat exchanger. Fouling in this context is the unwanted deposition of solids on the surfaces of the tubes of the heat exchanger, which leads to a loss in efficiency of the heat exchanger. Fouling is not limited to heat exchangers. Fouling may occur in other heat transfer components and transfer components for transferring process fluids. The loss in heat transfer efficiency results in higher fuel consumption at the furnace and reduced throughput. The buildup of foulants in fluid transfer components results in reduced throughput, higher loads on pumping devices and plugging of downstream equipment as large pieces of foulant periodically dislodge and flow downstream. As a result of fouling, the transfer components and heat transfer components must be periodically removed from service to be cleaned. This decreases overall facility reliability due to shutdowns for maintenance. This also leads to increased manpower requirements due to the number of cleaning crews required to service fouled heat exchanger and process fluid transfer tubes. Another detriment is an increase in volatile organic emission resulting from the cleaning process.

[0005] During normal use, the surfaces of the tubes of the heat exchanger are subject to corrosion as a result of the prolonged exposure to the stream of crude and other petroleum fractions. Corrosion on the surfaces of the tubes creates an uneven surface that can enhance fouling because the various particles found in the petroleum stream may attach themselves to the roughened surface. Fouling is not limited solely to the crude oils being processed. The vacuum residual streams are often used to heat the crude within the tubes. These streams often contain solids and are high fouling. Fouling can be associated with other process streams including but not limited to air. Fouling can be associated with other process streams including but not limited to process gases (e.g., air).

[0006] While the problems of fouling extend beyond petroleum refining and petrochemical processing, the presence of crude oil presents numerous obstacles in preventing fouling that are unique to petroleum refining and petrochemical processing not present in other industries. Crude oil, in the context of fouling, is in reality more than simply a petroleum product produced from an underground reservoir. Crude oil is a complex mixture of organic and inorganic components which may result in a variety of foulant deposits on the surfaces of the heat exchanger including but not limited to both surfaces of the heat exchanger tubes, the baffles and the tube sheets. For example, crude oil as it is received at the refinery often contains corrosion byproducts such as iron sulfide, which are formed by the corrosion of drilling tubulars, pipelines, tanker holds and crude storage tanks. This material, under the right conditions, will deposit within heat exchangers resulting in depositional fouling. Crude oils often contain aqueous contaminants, some of which arrive at the refinery. Desalting is used to remove most of this material, but some of these contaminants pass through the desalter into the crude preheat train. These dissolved salts can also contribute to depositional fouling. Sodium chloride and various carbonate salts are typical of this type of foulant deposit. As more and more chemicals are used to enhance production of crude from old reservoirs, additional inorganic materials are coming to the refineries in the crude oil and potentially contributing to fouling.

[0007] Crude oils are typically blended at the refinery, and the mixing of certain types of crudes can lead to another type of foulant material. The asphaltenic material that is precipitated by blending of incompatible crudes will often lead to a predominantly organic type of fouling, which with prolonged heating, will form a carbonaceous or coke-like foulant deposit. Crude oils often also contain acidic components that directly corrode the heat exchanger materials as well. Naphthenic acids will remove metal from the surface and sulfidic components will cause sulfidic corrosion which forms iron sulfide. This sulfidic scale that is formed is often referred to as sulfide induced fouling.

[0008] Synthetic crudes are derived from processing of bitumens, shale, tar sands or extra heavy oils and are also processed in refinery operations. These synthetic crudes present additional fouling problems, as these materials are too heavy and contaminant laden for the typical refinery to process. The materials are often pre-treated at the production site and then shipped to refineries as synthetic crudes. These crudes may contain fine particulate siliceous inorganic matter, such as in the case of tar sands. Some may also contain reactive olefinic materials that are prone to forming polymeric foulant deposits within heat exchangers. As can

be understood from this discussion, crude oils are complex mixtures capable of forming a wide-range of foulant deposit types.

[0009] Currently, there are various techniques available for reducing fouling in refinery operations. One technique is avoiding the purchase of high-fouling crudes or corrosive crudes. This, however, reduces the pool of feedstock that is potentially available to the refinery. Additionally, the crude oil can be tested to determine whether or not the crude oil is compatible with the refinery. Again, this can reduce the feedstock potentially available to the refinery. Anti-foulant agents may also be added to the refinery stream. While these techniques are useful in reducing the rate of fouling within the heat transfer components, fouling may still occur under certain circumstances. The heat exchangers must still be routinely removed from service for cleaning to remove the build-up of contaminants. Furnace tubes must be taken off-line and steam-air decoked or pigged because of foulant deposition. Other alternative cleaning methods include the use of mechanical devices (e.g., "SPIRELF" and "brush and basket" devices). These devices, however, have low reliability and high maintenance needs.

[0010] There is a need to significantly reduce fouling in heat transfer components in refinery and petrochemical processing operations that does not encounter the drawbacks associated with the current techniques.

Summary of the Invention

[0011] It is an aspect of the present invention to provide a heat transfer component that is resistant to fouling. The heat transfer component is used to either raise or lower the temperature of a process fluid or stream. The process fluid or stream is preferably crude oil based and is processed in a refinery or petrochemical facility. The present invention, however, is not intended to be limited solely to the use of crude oils, other process streams are considered to be well within the scope of the present invention. The heat transfer component may be a heat exchanger, a furnace, furnace tubes or any other component within a refinery or petrochemical facility that is capable of transferring heat from one medium to another which is also susceptible to fouling including but not limited to Crude Preheat, Coker preheat, FCC slurry bottoms, debutanizer exchanger/tower, other feed/effluent exchangers and furnace air preheaters in refinery facilities and flare compressor components and steam cracker/reformer tubes in petrochemical facilities. The heat transfer component contains at least one heat transfer element. It is contemplated that the heat transfer component is a heat exchanger for heating crude oil in a refinery stream prior to the crude entering a furnace, whereby the heat exchanger is resistant to fouling. The heat exchanger may be a tube-in-shell type heat exchanger having a tube bundle located within a housing. The present invention is not intended

to be limited to tube-in-shell exchangers; rather, the present invention has application within other exchangers which are prone to fouling when subject to petroleum and/or vacuum residual streams. The tube-in-shell exchanger includes a housing having a wall forming a hollow interior. The wall has an inner surface that is adjacent the hollow interior. The heat transfer element may be a tube bundle located within hollow interior of the housing. The crude oil is heated within the hollow interior of the heat exchanger housing as the crude oil flows over the tube bundle. The tube bundle preferably includes a plurality of heat exchanger tubes.

[0012] It is an aspect of the present invention to provide a heat transfer component for heating a process stream that is resistant to sulfidation corrosion and corrosion fouling when exposed to the process stream. The heat transfer component includes a housing having a wall forming a hollow interior, wherein the wall having an inner surface, at least one heat transfer element located within the housing for heating the process stream within the hollow interior of the housing. In accordance with the present invention at least one of the inner surface and the at least one heat transfer element having a surface roughness of less than 40 micro inches (1.1 μm). The surface roughness is preferably less than 20 micro inches (0.5 μm). More preferably, the surface roughness is less than 10 micro inches (0.25 μm). The at least one heat transfer element being formed from a composition that is resistant to sulfidation corrosion and corrosion induced fouling.

[0013] The heat transfer component may be a heat exchanger and the at least one heat transfer element is a tube bundle having a plurality of heat transfer tubes, wherein each of the heat transfer tubes having an inner diameter surface and an outer diameter surface, wherein at least one of the inner diameter surface and the outer diameter surface having a surface roughness of less than 40 micro inches (1.1 μm). It is preferable that at least one of the interior surface of the wall of the heat transfer component and the inner and/or outer surfaces of the plurality of heat exchanger tubes is formed in accordance with this invention to have a surface roughness of less than 40 micro inches (1.1 μm). Preferably, the surface roughness is less than 20 micro inches (0.5 μm). More preferably, the surface roughness is less than 10 micro inches (0.25 μm). It is contemplated that both the inner and outer surfaces of the plurality of heat exchanger tubes may have the above-mentioned surface roughness. Such a surface roughness significantly reduces fouling. The smooth surface within the inner diameter of the tubes reduces fouling of the petroleum stream flowing through the tubes. The smooth surfaces on the outer diameter of the tubes and on the inner surface of the housing will reduce fouling of the vacuum residual stream within the housing. It is also contemplated that the surfaces of the baffles located within the heat exchanger and the surfaces of the tube sheets, which secure the

tubes in place may also have the above-mentioned surface roughness. Such a surface roughness would significantly reduce fouling on these components.

[0014] It is an aspect of the present invention to provide a sleeve that is resistant to sulfidation corrosion and corrosion fouling for use in a heat transfer component. The heat transfer component includes at least one heat transfer element having an inner diameter surface and an outer diameter surface, wherein the heat transfer element being formed from a first material. The sleeve in accordance with the present invention has a first surface adapted to contact one of the inner diameter surface and the outer diameter surface of the heat transfer element and a second surface adapted to contact the process stream, wherein the second surface having the surface roughness of less than 40 micro inches (1.1 μm). The sleeve being formed from a composition that is resistant to sulfidation corrosion and corrosion induced fouling.

[0015] The use of composition that is resistant to sulfidation or sulfidic corrosion and fouling for forming at least a portion of the heat transfer component and/or the sleeve significantly reduces fouling and corrosion, which produces numerous benefits including an increase in heating efficiency, a reduction in the overall amount of energy needed to heat the crude oil, an increase in refinery throughput and a significant reduction in refinery downtime.

[0016] In accordance with one aspect of the present invention, the composition for the sleeve or the heat transfer component is a steel composition containing X, Y, and Z, wherein X is a metal selected from the group consisting of Fe, Ni, Co and mixtures thereof, wherein Y is Cr, and wherein Z is at least one alloying element selected from the group consisting of Si, Al, Mn, Ti, Zr, Hf, V, Nb, Ta, Mo, W, Sc, Y, La, Ce, Pt, Cu, Ag, Au, Ru, Rh, Ir, Ga, In, Ge, Sn, Pb, B, C, N, O, P, and S. The sleeve or the heat transfer element contains Cr-enriched layer formed thereon, wherein the Cr-enriched layer also being formed from the steel composition X, Y and Z. The ratio of Y to X in the Cr-enriched layer being greater than the ratio of Y to X in the remaining portion of the heat transfer element or sleeve. The Cr-enriched layer may be formed by one of electro-polishing at least on surface of heat transfer element, electroplating, thermal spray coating, laser deposition, sputtering, physical vapor deposition, chemical vapor deposition, plasma powder welding overlay, cladding, and diffusion bonding. A protective layer may be formed on an outer surface of the Cr-enriched layer. The protective layer comprises an oxide selected from the group consisting of a magnetite, an iron-chromium spinel, a chromium oxide, and mixtures thereof. The protective layer is formed when the Cr-enriched layers is subjected to a process stream at high temperatures up to 1100°C.

[0017] In accordance with another aspect of the present invention, the composition for the sleeve or the heat transfer component is a carbon steel having an aluminum or aluminum alloy layer located thereon.

[0018] In accordance with another aspect of the present invention, the composition for the sleeve or the heat transfer component is one aluminum, titanium, chromium and alloys of aluminum, titanium, chromium.

[0019] In accordance with another aspect of the present invention, the composition for the sleeve or the heat transfer component is a chromium-enriched oxide containing material formed from the composition δ , ϵ , and ζ . ζ is a steel containing at least about 5 to about 40 wt. % chromium. ϵ is a chromium enriched oxide (M_3O_4 or M_2O_3 or mixtures thereof) formed on the surface of the steel ζ . M is a metal containing at least 5 wt. % Cr based on the total weight of the metal M. δ is a top layer formed on the surface of the chromium-enriched oxide ϵ , comprising sulfide, oxide, oxysulfide, and mixtures thereof. The top layer δ comprises iron sulfide ($Fe_{1-x}S$), iron oxide (Fe_3O_4), iron oxysulfide, iron-chromium sulfide, iron-chromium oxide, iron-chromium oxysulfide, and mixtures thereof. The metal M of the chromium enriched oxide (M_3O_4 or M_2O_3 or mixtures thereof) may comprise Fe, Cr, and constituting elements of the steel ζ . The steel ζ may be selected from low chromium steels, ferritic stainless steels, martensitic stainless steels, austenitic stainless steels, duplex stainless steels and precipitation-hardenable alloys.

[0020] In accordance with another aspect of the present invention, the composition of a heat exchange surface for the sleeve or the heat transfer component is a silicon containing steel composition including an alloy and a non-metallic film formed on the surface of the alloy. The alloy is formed from the composition η , θ , and ι . η is a metal selected from the group consisting of Fe, Ni, Co, and mixtures thereof. θ is Si. ι is at least one alloying element selected from the group consisting of Cr, Al, Mn, Ti, Zr, Hf, V, Nb, Ta, Mo, W, Sc, La, Y, Ce, Ru, Rh, Ir, Pd, Pt, Cu, Ag, Au, Ga, Ge, As, In, Sn, Sb, Pb, B, C, N, P, O, S and mixtures thereof. The non-metallic film comprises sulfide, oxide, carbide, nitride, oxysulfide, oxycarbide, oxynitride and mixtures thereof. The metal η may comprise at least about 60 wt% to about 99.98 wt. % of the alloy. The alloying metal θ may comprise at least about 0.01 wt% to about 5.0 wt. % of the alloy. Preferably, the alloying metal θ comprises at least about 0.01 wt% to about 3.0 wt. % of the alloy. The alloying element ι comprise at least about 0.01 wt% to about 40.0 wt% of the alloy. Preferably, the alloying element ι comprises at least about 0.01 wt% to about 3.0 wt% Al of the alloy. The alloying element ι may comprise at least about 0.01 wt% to about 30.0 wt% Cr of the alloy. The alloying element ι may comprise at least about

0.01 wt% to about 3.0 wt% Al and at least about 0.01 wt% to about 30.0 wt% Cr of the alloy. The non-metallic film on the surface of the alloy may comprise at least 1 nm thick Si-partitioned non-metallic film. The non-metallic film is a multi-layer non-metallic film and consists of at least 10 atomic percent Si based on the concentration of the non-metallic film.

[0021] To further reduce and/or limit fouling, the heat transfer component may include or be operatively coupled to a vibration producing device to impart a vibrational force to the heat transfer element which results in the development of a shear motion in the liquid flowing within the heat exchanger. This shear motion or turbulent flow within the heat transfer component limits the formation of any foulant on the surfaces of the component by reducing the viscous boundary layer adjacent the walls of the heat transfer element. The heat transfer component may include or be operatively coupled to a pulsation producing device to apply pressure pulsations to the process stream.

[0022] It is another aspect of the present invention to provide a method of reducing fouling in a heat exchanger for a process stream. The heat exchanger has a plurality of existing heat exchanger tubes. The method includes removing at least a portion of the plurality of existing heat exchanger tubes from the heat exchanger; and installing a plurality of replacement heat exchanger tubes. Each of the plurality of replacement heat exchanger tubes being formed from a composition that is resistant to sulfidation corrosion and fouling, described above, and a surface roughness of less than 40 micro inches (1.1 μm). Alternatively, each replacement tube may contain a sleeve located therein. The sleeves being formed from a composition that is resistant to sulfidation corrosion and corrosion induced fouling.

[0023] It is another aspect of the present invention to provide a method of providing sulfidation corrosion resistance and corrosion induced fouling resistance to a metal surface that is subject to a process stream at high temperatures. The method includes providing a metal layer having a surface roughness of less than 40 micro inches (1.1 μm) formed from a composition comprising X, Y, and Z, wherein X is a metal selected from the group consisting of Fe, Ni, Co and mixtures thereof, wherein Y is Cr, and wherein Z is at least one alloying element selected from the group consisting of Si, Al, Mn, Ti, Zr, Hf, V, Nb, Ta, Mo, W, Sc, Y, La, Ce, Pt, Cu, Ag, Au, Ru, Rh, Ir, Ga, In, Ge, Sn, Pb, B, C, N, O, P, and S, wherein a Cr-enriched layer is located on the metal layer, wherein the Cr-enriched layer also being formed from the steel composition X, Y, and Z, wherein the ratio of Y to X in the Cr-enriched layer being greater than the ratio of Y to X in the metal layer. The method further includes forming an protective layer on a surface of the Cr-enriched layer.

[0024] It is yet another aspect of the present invention to provide a method of providing sulfidation corrosion resistance and corrosion induced fouling resistance to a heat transfer component surface. The method includes providing a silicon containing steel composition including an alloy, wherein the alloy is formed from the composition η , θ , and ι , in which η is a metal selected from the group consisting of Fe, Ni, Co, and mixtures thereof, θ is Si, and ι is at least one alloying element selected from the group consisting of Cr, Al, Mn, Ti, Zr, Hf, V, Nb, Ta, Mo, W, Sc, La, Y, Ce, Ru, Rh, Ir, Pd, Pt, Cu, Ag, Au, Ga, Ge, As, In, Sn, Sb, Pb, B, C, N, P, O, S and mixtures thereof. The method further includes forming a non-metallic film on the surface of the alloy having a surface roughness of less than 40 micro inches (1.1 μm), wherein the non-metallic film comprises sulfide, oxide, carbide, nitride, oxysulfide, oxycarbide, oxynitride and mixtures thereof.

[0025] It is yet another aspect of the present invention to provide a method of providing sulfidation corrosion resistance and corrosion induced fouling resistance to a heat transfer component surface. The method includes providing a steel surface containing at least about 5 to about 40 wt. % chromium. The method further includes forming a chromium-enriched oxide (M_3O_4 or M_2O_3 or mixtures thereof) on the surface of the steel, wherein M is a metal containing at least 5 wt. % Cr based on the total weight of the metal M. The method also includes forming a top layer formed on the surface of the chromium-enriched oxide having a surface roughness of less than 40 micro inches (1.1 μm), comprising sulfide, oxide, oxysulfide, and mixtures thereof.

Brief Description of the Drawings

[0026] The invention will now be described in connection with the following drawings in which like reference numerals designate like elements and wherein:

[0027] Fig. 1 is an example of heat exchanger having a plurality of heat exchanger tubes for use in a refinery operation;

[0028] Fig. 2 is a schematic view illustrating the various layers forming the steel composition utilized in forming the heat transfer components in accordance with an embodiment of the present invention;

[0029] Fig. 3 is a schematic view illustrating the various layers forming the aluminum clad carbon steel utilized in forming the heat transfer components in accordance with another embodiment of the present invention;

[0030] Fig. 4 is a partial sectional view of a heat exchanger tube in accordance with the present invention;

[0031] Figs. 5 and 6 are images illustrating the fouling on a conventional heat exchanger tube after a field trial;

[0032] Figs. 7 and 8 are images illustrating the significant reduction in fouling on a heat exchanger tube in accordance with the present invention after a field trial;

[0033] Fig. 9 is an insert sleeve in accordance with the present invention

[0034] Figure 10 is a partial side view in section of a heat transfer component having its inner diameter formed in accordance with the surface disclosed herein;

[0035] Figure 11 is a partial side view in section of a heat transfer component having its outer diameter formed in accordance with the surface disclosed herein;

[0036] Figure 12 is a partial side view in section of a heat transfer component having both its inner diameter and its outer diameter formed in accordance with the surface disclosed herein;

[0037] Figure 13 shows a cross section of a surface treated as explained with respect to the first example herein;

[0038] Figure 14 is a graph illustrating atomic concentration versus sputter depth for a test specimen;

[0039] Figure 15 is a graph illustrating dimensionless temperature change over time for a test specimen;

[0040] Figure 16 is a graph illustrating dimensionless temperature change over time for another test specimen;

[0041] Fig. 17 is a partial side view in section of a heat transfer component having its inner diameter formed in accordance with the surface disclosed herein;

[0042] Fig. 18 is a partial side view in section of a heat transfer component having its outer diameter formed in accordance with the surface disclosed herein;

[0043] Fig. 19 is a partial side view in section of a heat transfer component having both its inner diameter and its outer diameter formed in accordance with the surface disclosed herein;

[0044] Fig. 20 shows a cross section of a surface treated as explained with respect to the first example herein; and,

[0045] Fig. 21 shows a cross section of a surface treated as explained with respect to the second example herein.

Detailed Description of Preferred Embodiments

[0046] The present invention will now be described in greater detail in connection with the attached figures. Fig. 1 is a tube-in-shell heat exchanger 10, which is located upstream from a furnace (not shown) and employs the principles of the present invention. The tube-in-shell heat exchanger 10 disclosed herein illustrates one application of the present invention to reduce sulfidation or sulfidic corrosion and depositional fouling in refinery and petrochemical applications. The tube-in-shell exchanger 10 is just one heat transfer component falling under the scope of the corrosion reduction and fouling mitigation measures in accordance with the present invention. The principles of the present invention are intended to be used in other heat exchangers including but not limited to spiral heat exchangers, tube-in-tube heat exchangers and plate-and-frame heat exchangers having at least one heat transfer element. The principles of the present invention are intended to be employed in other heat transfer components including furnaces, furnace tubes and other heat transfer components which may be prone to petroleum and/or vacuum residual fouling. The heat exchanger 10 is used to pre-heat crude oil in a refinery operation prior to entry into the furnace. The heat exchanger 10 includes a housing or shell 11, which surrounds and forms a hollow interior 12. A bundle 13 of heat exchanger tubes 14 is located within the hollow interior 12, as shown in Fig. 1. The bundle 13 includes a plurality of tubes 14. The tubes 14 may be arranged in a triangular configuration or a rectangular configuration. Other tube arrangements are contemplated and considered to be well within the scope of the present invention. Each tube 14 has a generally hollow interior 15 such that the crude oil to be heated flows there-through. The heating or warming fluid (e.g., vacuum residual stream) flows through the hollow interior 12 to pre-heat the crude oil stream as the stream flows through the hollow interior 15 towards the furnace. Alternatively, it is contemplated that the crude oil may flow through the hollow interior 12 of the housing 11. The housing 11 and the tubes 14 are preferably formed from a steel composition. It is contemplated that the housing 11 and the tubes 14 may be formed from the same material. It is also contemplated that the housing 11 and the tubes 14 may be formed from different materials. Typically, the tubes and the housing are formed from a carbon or low chromium content steel.

[0047] As described above, heat exchangers are typically subject to fouling after prolonged exposure to crude oil. The presence of fouling reduces the performance of the heat exchanger. Figs. 5 and 6 illustrate the effects of fouling on the surface of a heat exchanger tube. The presence of fouling reduces throughput and increases fuel consumption. Figs. 5 and

6 illustrate the amount of fouling present within the heat exchanger tube after five months of operation. This fouling represents an approximately 31% reduction in the heat exchanger efficiency in the refinery. The foulant contains sodium chloride, iron sulfide and carbonaceous materials. As shown in Figs. 5 and 6, significant amounts of pitting are present. Pitting can further exacerbate the fouling problem.

[0048] By contrast, Figs. 7 and 8 illustrate the reduction in fouling utilizing heat exchanger tubes 14 which embody the principles of the present invention. The surface cross-sections illustrated in Figs. 7 and 8 illustrate a marked reduction in fouling. These tubes were located in the same heat exchanger and subject to the same operating conditions over the same five month period. While the foulant present in the exchanger tube 14 also contained sodium chloride, iron sulfide and carbonaceous material, the amount of foulant was significantly reduced. The thickness of the foulant was reduced to less than 10 microns. The tubes having the reduced surface roughness also exhibited less pitting. The conventional tubes illustrated in Figs. 5 and 6 exhibited a mean foulant deposit weight density of 46 mg/cm². By contrast, the tubes 14 constructed using principles in accordance with the present invention illustrated at least 50% reduction in the mean foulant deposit weight density. The sample tubes exhibited a mean foulant deposit weight density of 22 mg/cm². Deposit weight density was determined by the National Association of Corrosion Engineers (NACE) method TM0199-99. The reduction in fouling shown in Figs. 7 and 8 illustrate the benefits of the present invention.

[0049] The reduction in fouling may be obtained as a result of controlling the surface roughness of the inner diameter surface and the outer diameter surface of the tubes 14 and/or the interior surface of the shell 11. Controlling the surface roughness of the inner diameter surface of the tubes mitigates the fouling of process fluid or crude oil within the tubes 14. Controlling the surface roughness of the outer diameter surface of the tubes 14 and the inner surface of the shell 11 mitigates fouling associated with the heating fluid (e.g., vacuum residual) flowing within the hollow interior 12. In accordance with the present invention, at least one of the interior surface of the hollow interior 12 and the surfaces of the tubes 14 has a surface roughness of less than 40 micro inches (1.1 μ m). Surface roughness can be measured in many ways. Industry prefers to use a skidded contact profilometer. Roughness is routinely expressed as the arithmetic average roughness (Ra). The arithmetic average height of roughness component of irregularities from the mean line is measured within the sample length L. The standard cut-off is 0.8 mm with a measuring length of 4.8 mm. This measurement conforms to ANSI/ASME B46.1 "Surface Texture- Surface Roughness, Waviness and Lay," which was employed in determining the surface roughness in accordance with the present

invention. A uniform surface roughness of less than 40 micro inches (1.1 μm) produces a significant reduction in fouling.

[0050] Further reductions in surface roughness are desirable. It is preferable that the surface roughness be below 20 micro inches (0.5 μm). It is more preferable that the surface roughness be below 10 micro inches (0.25 μm). It is preferable that both the inner diameter surface and the outer diameter surface have the described surface roughness. The desired surface roughness may be obtained through various techniques including but not limited to mechanical polishing and electro-polishing. In the samples illustrated in Figs. 5 and 6, the surface roughness of the tubes was variable between 38 and 70 micro inches. The tubes in Figs. 5 and 6 were not polished. The tubes illustrated in Figs. 7 and 8, which form the basis for the present invention were polished to a more uniform 20 micro inches (0.5 μm). This was accomplished using conventional mechanical polishing techniques. The tubes were then electro-polished in an acidic electrolyte to produce a reflective surface having a surface roughness below 10 micro inches (0.25 μm). The treated tubes exhibited a marked reduction in fouling.

[0051] In accordance with the present invention, it is preferable that the tubes 14 be formed from a steel composition that is resistant to sulfidation or sulfidic corrosion and depositional fouling. The use of such a steel composition significantly reduces fouling, which produces numerous benefits including an increase in heating efficiency, a reduction in the amount of energy needed to pre-heat the crude oil, and a significant reduction in refinery downtime and throughput. It is preferable that the tubes 14 and/or the housing 11 of the pre-heat exchanger have several layers, as illustrated in Figs. 2 and 4. The primary layer 21 is a steel composition containing three primary components or constituents X, Y and Z. X denotes a metal that is selected from the group preferably consisting of Fe, Ni, and Co. X may also contain mixtures of Fe, Ni and Co. Y denotes Cr. In accordance with the present invention, a steel composition contains Cr at least greater than 1 wt.% based on the combined weight of the three primary constituents X, Y and Z. Higher Cr contents are desirable for improved sulfidation or sulfidic corrosion resistance. It is preferable that the Cr content be higher than 5 wt.% based on the combined weight of three primary constituents X, Y and Z. It is more preferable that the Cr content be higher than 10 wt.% based on the combined weight of three primary constituents X, Y and Z. Z is preferably an alloying element.

[0052] In accordance with the present invention, Z preferably includes at least one alloying element selected from the group consisting of Si, Al, Mn, Ti, Zr, Hf, V, Nb, Ta, Mo, W, Sc, Y, La, Ce, Pt, Cu, Ag, Au, Ru, Rh, Ir, Ga, In, Ge, Sn, Pb, B, C, N, O, P, and S. Z may

also contain mixtures of Si, Al, Mn, Ti, Zr, Hf, V, Nb, Ta, Mo, W, Sc, Y, La, Ce, Pt, Cu, Ag, Au, Ru, Rh, Ir, Ga, In, Ge, Sn, Pb, B, C, N, O, P, and S. The weight percent of an alloying element is preferably higher than 0.01 wt.%, and more preferably higher than 0.05 wt.%, and most preferably higher than 0.1 wt.%, based on the combined weight of three primary constituents X, Y and Z. The combined weight percent of all alloying elements in a steel composition is preferably less than 10 wt.%, and more preferably less than 5 wt.%, based on the combined weight of three primary constituents X, Y and Z. While other compositions are considered to be within the scope of the present invention, the above-described composition has been found to reduce fouling.

[0053] Table 1 illustrates non-limiting examples of a steel composition that is resistant to sulfidation or sulfidic corrosion and corrosion induced fouling for use in both refining and petrochemical applications. Other materials exhibiting similar properties are considered to be well within the scope of the present invention provided such materials fall within the scope of the prescribed ranges.

TABLE 1

Name (Grade)	UNS Number	Constituent X in wt.%	Constituent Y in wt.%	Constituent Z in wt.%
T11	K11562	Balanced Fe	1.25Cr	0.5Mo, 0.5Si, 0.3Mn, 0.15C, 0.045P, 0.045S
T22	K21590	Balanced Fe	2.25Cr	1.0Mo, 0.5Si, 0.3Mn, 0.15C, 0.035P, 0.035S
T5	S50100	Balanced Fe	5Cr	0.5Mo, 0.5Si, 0.3Mn, 0.15C, 0.04P, 0.03S
T9	J82090	Balanced Fe	9Cr	1.0Si, 0.35Mn, 0.02C, 0.04P, 0.045S
409	S40900	Balanced Fe	10.5Cr	1.0Si, 1.0Mn, 0.5Ni, 0.5Ti, 0.08C, 0.045P, 0.045S
410	S41000	Balanced Fe	11.5Cr	0.15C, 0.045P, 0.03S
430	S43000	Balanced Fe	16Cr	1.0Si, 1.0Mn, 0.12C, 0.045P, 0.03S
XM-27 E-Brite	S44627	Balanced Fe	25Cr	0.5Ni, 0.75Mo, 0.4Si, 0.4Mn, 0.05Nb, 0.2Cu, 0.01C, 0.02P, 0.02S, 0.015N

SeaCure	S44660	Balanced Fe	25Cr	1.5Ni, 2.5Mo, 1.0Si, 1.0Mn, 0.05Nb, 0.2Cu, 0.025C, 0.04P, 0.03S, 0.035N
304	S30400	Bal.Fe, 8Ni	18Cr	2.0Mn, 0.75Si, 0.08C, 0.04P, 0.03S
304L	S30403	Bal.Fe, 8Ni	18Cr	2.0Mn, 0.75Si, 0.035C, 0.04P, 0.03S
309S	S30908	Bal.Fe, 12Ni	22Cr	2.0Mn, 0.75Si, 0.75Mo, 0.08C, 0.045P, 0.03S
310	S31000	Bal.Fe, 19Ni	24Cr	2.0Mn, 1.5Si, 0.75Mo, 0.25C, 0.045P, 0.03S
316	S31600	Bal.Fe, 11Ni	16Cr	2.0Mn, 0.75Si, 2.0Mo, 0.08C, 0.04P, 0.03S
316L	S31603	Bal.Fe, 11Ni	16Cr	2.0Mn, 0.75Si, 2.0Mo, 0.035C, 0.04P, 0.03S
321	S32100	Bal.Fe, 9Ni	17Cr	2.0Mn, 0.75Si, 0.4Ti, 0.08C, 0.045P, 0.03S
2205	S32205	Bal.Fe:4.5Ni	22Cr	2.0Mn, 1.0Si, 3.0Mo, 0.03C, 0.14N, 0.03P, 0.02S
2507	S32507	Bal.Fe:6Ni	24Cr	1.2Mn, 0.8Si, 3.0Mo, 0.5Cu, 0.03C, 0.2N, 0.035P, 0.02S
AL-6XN	N08367	Bal.Fe:24Ni	20Cr	6.2Mo, 0.4Si, 0.4 Mn, 0.22N, 0.2Cu, 0.02C, 0.02P, 0.03S, 0.035N
Alloy 800	N08800	Bal.Fe:30Ni	19Cr	0.15Ti, 0.15Al

[0054] The chromium enrichment at the surface of the non-fouling surface is advantageous. Therefore, the steel composition preferably includes a chromium enriched layer 22. The Cr-enriched layer 22 is formed on the primary layer 21. The layer 22 may be formed on both the inner surface and the exterior surface of the tubes. The thickness of the Cr-enriched layer 22 is greater than 10 angstroms. The Cr-enriched layer 22 contains the same three primary components or constituents X, Y and Z. X denotes a metal that is selected from the group preferably consisting of Fe, Ni, and Co. X may also contain mixtures of Fe, Ni, Co and Ti. Y denotes Cr. It is contemplated that Y may also comprise Ni, O, Al, Si and mixtures thereof.

The percentage of Cr is higher in layer 22 when compared to the primary layer 21. In accordance with the present invention, Cr content in layer 22 is at least greater than 2 wt.% based on the combined weight of three primary constituents X, Y and Z. It is preferable that the Cr content be higher than 10 wt.% based on the combined weight of three primary constituents X, Y and Z. It is more preferable that the Cr content be higher than 30 wt.% based on the combined weight of three primary constituents X, Y and Z. In the layer 22, the ratio of Y to X is greater than the ratio of Y to X in the layer 21. The ratio should be greater by a factor of at least 2. The ratio should preferably be greater by a factor of at least four. More preferably, the ratio should be a greater by a factor of eight. Z is preferably an alloying element.

[0055] For example, 5-chrome steel (T5) nominally contains about 5 wt.% chromium per about 95 wt.% iron to give an untreated surface ratio of 0.05 in the primary layer 21. In the Cr-enriched layer 22, the ratio increased to at least 0.1, preferably to 0.2 and most preferably to 0.4 chromium atoms per iron atom in the surface layer of the heat exchanger tube. For 316L stainless steel, which has nominally 16 wt.% Cr, 11 wt.% Ni, 2 wt.% Mn, 2 wt.% Mo, the bulk ratio of chromium to iron would be $16/69 = 0.23$. After treatment to enrich the surface chromium, the ratio may rise to at least 0.46, preferably 0.92 and most preferably 1.84.

[0056] In the Cr-enriched layer 22, Z preferably includes at least one alloying element selected from the group consisting of Si, Al, Mn, Ti, Zr, Hf, V, Nb, Ta, Mo, W, Sc, Y, La, Ce, Pt, Cu, Ag, Au, Ru, Rh, Ir, Ga, In, Ge, Sn, Pb, B, C, N, O, P, and S. The weight percent of an alloying element is preferably higher than 0.01 wt.%, and more preferably higher than 0.05 wt.%, and most preferably higher than 0.1 wt.%, based on the combined weight of three primary constituents X, Y and Z.

[0057] It is contemplated that the Cr-enriched layer 22 may be formed on two sides of the primary layer 21 such that both the interior surface and the exterior surface contain a Cr-enriched layer. The Cr-enriched layer 22 may be formed on the primary layer 21 using one of several techniques. The Cr-enriched layer may be formed by electro-polishing the tube in a solution containing chromic acid. This is effective when the Cr content in the steel composition is less than about 15 wt. %. It is also contemplated that the Cr-enriched layer 22 may be formed using various other formation techniques including but not limited to electroplating chromium onto another alloy such as a carbon steel, bright annealing, passivation, thermal spray coating, laser deposition, sputtering, physical vapor deposition, chemical vapor deposition, plasma powder welding overlay, cladding, and diffusion bonding. It is also possible to choose a high chromium alloy including but not limited to 304L stainless

steel, 316 stainless steel and AL6XN alloy. In accordance with the present invention, the secondary layer 22 may be mechanically polished and/or electro-polished as described above in order to obtain a uniform surface roughness of less than 40 micro inches (1.1 μm), preferably less than 20 micro inches (0.5 μm) and more preferably less than 10 micro inches (0.25 μm). The desired surface roughness can also be achieved using fine abrasive polishing or metal peening.

[0058] The Cr-enriched layer 22 may be formed on the primary layer 21 by bright annealing the tube. Bright annealing is an annealing process that is carried out in a controlled atmosphere furnace or vacuum in order that oxidation is reduced to a minimum and the surface remains relatively bright. The process conditions such as atmosphere, temperature, time and heating/cooling rate utilized during the bright annealing process will be dependent on the metallurgy of the alloy being acted upon. The skilled artisan can easily determine the conditions based on the alloy's metallurgy. As a non-limiting example, the austenitic stainless steel such as 304L can be bright annealed in either pure hydrogen or dissociated ammonia, provided that the dew point of the atmosphere is less than -50°C and the tubes, upon entering the furnace, are dry and scrupulously clean. Bright annealing temperatures usually are above 1040°C . Time at temperature is often kept short to hold surface scaling to a minimum or to control grain growth.

[0059] In accordance with the present invention, a protective layer 23 is preferably formed on the Cr-enriched layer 22. The Cr-enriched layer 22 is necessary for the formation of the protective layer 23. The protective layer may be an oxide layer, a sulfide layer, an oxysulfide layer or any combination thereof. The protective layer 23 preferably includes a material such as a magnetite, an iron-chromium spinel, a chromium oxide, oxides of the same and mixtures thereof. The layer 23 may also contain a mixed oxide sulfide thiospinel. While it is possible to form the protective layer 23 on the Cr-enriched layer 22 prior to installation of the tubes 14 within the housing 11 of the pre-heat exchanger 10, the protective layer 23 is preferably formed on the Cr-enriched layer 22 after the tubes 14 are located within the exchanger 10 and the pre-heat exchanger is operational. The protective layer 23 forms when the Cr-enriched layer is exposed to the process stream at high temperatures. In a late-train heat exchanger application, the protective layer forms at temperatures up to 400°C . In applications in a furnace or outside the late-train heat exchanger, the protective layer forms at temperatures up to 600°C . In petrochemical applications including use in steam cracker and reformer tubes, the protective layer forms at temperatures up to 1100°C . The thickness of the protective layer 23 is preferably greater than 100 nm, more preferably greater than 500 nm, and most preferably

greater than 1 micron. As illustrated in figures 6 and 7, a field trial of 5-chrome steel revealed about 1 micron thick Cr-enriched magnetite layer formed during about 4 months of period. Since the stream oil flowing within a heat exchanger tube is a highly reducing and sulfidizing environment, the protective layer 23 can further convert to a mixed oxide-sulfide layer or a thiospinel-type sulfide layer after prolonged exposure. Applicants note that the formation of the protective layer 23 is a result of the electro-polishing of the Cr-enriched layer 21.

[0060] The formation of the protective layer further reduces fouling. The foulants, which form on the protective layer 23 exhibit significantly less adhesion characteristics when compared to foulants, which form on surfaces that do not have the protective layer. One benefit of this reduced adhesion lies in the cleaning of the heat exchange surface. Less time is required to remove any foulants from the tubes. This results in a decrease in downtime such that the pre-heat exchanger can be serviced in a more efficient manner and placed back online sooner. Also, with a less adherent deposit, on-line cleaning methods may become more effective or at least more rapid, which will further reduce downtime and throughput loss.

[0061] There are numerous additional benefits of reducing the surface roughness of the tubes 14. One of the benefits is the shifting from a linear growth rate of the foulant, which results in the continuous thickening of the foulant deposit; to an asymptotic growth rate which reaches a finite thickness and then stops thickening.

[0062] The tubes 14 disclosed above may be used to form new heat exchangers. The tubes 14 can also be used in existing exchangers as replacement tubes. The use of the tubes 14 should produce significant benefits in the refinery operations. In addition to reducing fouling, there is a reduction in the number of scheduled downtimes the heat exchangers operate more efficiently because the harmful effects of fouling are reduced. In addition, as demonstrated in the field test, the use of the tubes 14 will also prolong tube life due to reduced pitting corrosion.

[0063] The tubes 14 in accordance with the present invention may be used to retrofit an existing heat exchanger during a scheduled downtime. The existing tubes can be removed from the heat exchanger. The tubes 14 having the above described surface roughness and/or material composition are installed in the interior 12 of the housing 11. While it is preferable to replace all of the existing heat exchanger tubes with replacement tubes having the above-described construction in order to maximize the reduction in fouling, the present invention is not intended to be so limited. It is contemplated that only a portion of the existing heat exchanger tubes be replaced with replacement tubes. While such a construction may not result in the same reduction in fouling, a degree of fouling mitigation will be obtained. The determination of the number and location of existing tubes to be replaced by the replacement

tubes can be determined by a physical inspection of the tubes within the bundle within the heat exchanger. The tubes located closest to the furnace may be more prone to fouling. As such, it is also contemplated that tubes located most closely to the furnace may be replaced with tubes 14.

[0064] It may not be cost effective to replace all or a portion of the tubes of the heat transfer component with tubes 14 constructed in accordance with the principles of the present invention. In accordance with another aspect of the present invention, an insert 50 is provided for use in retrofitting existing heat transfer components to mitigate fouling. The insert 50 will be described in connection with Fig. 9. The insert 50 is sized such that the outer diameter of the insert is sized to abut the inner diameter surface of the tube 60. The tube 60 is secured to a tube sheet 70. Since it is intended that the insert 50 is to be retrofit in existing operation heat transfer components, it is contemplated that a layer of corrosion 61 (e.g., an oxidized layer) may exist between the tube 60 and the insert 50. It is also possible to use the insert for fitting over a tube. In this case, the insert is sized to closely surround the tube, which is susceptible to fouling, build-up or corrosion on its outer diameter surface. The insert 50 in accordance with the present invention may be constructed from any of the materials described herein having the above described surface roughnesses. It is important that the insert or sleeve 50 contact the tube 60 such that the heat transfer properties are not adversely or significantly diminished.

[0065] One advantage of the use of inserts 50 is that it allows one to retrofit an existing heat exchanger and convert it thereby to a non-fouling heat exchanger. This avoids the cost and time associated with the construction of a new bundle. For example, an existing heat exchanger subject to repeated fouling and requiring significant downtime due to required cleaning could be retrofitted with stainless steel inserts and electro-polished in situ to achieve a smooth, corrosion-resistant, non-fouling tube inner diameter surface. This would be much cheaper than replacing the entire heat exchanger bundle with a stainless steel equivalent (including tube sheets and baffles) and using electro-polished stainless steel tubes.

[0066] An additional advantage is that in some applications, it is not possible to use solid tubes made of a given alloy, even if this could prevent fouling on the inner diameter. As such, it is possible to use an insert with new or replacement tubes. For example, there are real world examples where fouling on the tube inner diameter could be prevented by use of an electro-polished stainless steel tube instead of the currently used 5-chrome tube. However, in this case, the shell side, or the tube outer diameter, is exposed to an aqueous environment at elevated temperatures where stainless steel cannot be used. The potential for stress corrosion cracking is the issue. The advantage of the present invention is that an electro-polished

stainless steel insert can be used within the 5-chrome tube. The electro-polished stainless steel inserts reduces fouling on the tube inner diameter and the 5-chrome tube itself is adequate with respect to the aqueous environment on the tube outer diameter. Similarly, highly corrosion resistant and reduced surface roughness titanium alloy inserts could be inserted within a less corrosion resistant outer tube for use in applications where a highly corrosive process fluid is passing through the tubes and a less corrosive fluid is in contact with the outer diameter. These are just two examples of the generic situation where the use of electro-polished inserts may be the only feasible route to reducing tube inner diameter fouling.

[0067] The insert 50 is relevant to fouling beyond the petroleum and petrochemical industries. Other potential applications include for example, black liquor fouling and paper slurry pipe systems in the paper and pulp industry, microbiological contamination (bio-fouling) in the water treatment and distillation industry, product contamination in the pharmaceutical and semiconductor industries, reduction of contamination pick-up and increasing the effectiveness of conventional decontamination techniques for recirculation piping in the nuclear power industry and piping and exchangers used in the food, dairy and beverage industries where product build-up is a problem. Polymer sheeting in chemical reactor piping and heat exchangers used to produce polymers could also benefit as well as heat exchangers used to remove water during crystallization processes. There are many other examples of heat exchanger and piping fouling that can be ameliorated by this invention.

[0068] Many potential metal types can be used to form the insert and the metal chosen will depend on the nature of the fluid stream and the type of fouling that is being prevented. Though austenitic stainless steels such as types 304 and 316 can be used to fabricate the tube inserts, other stainless steel alloys could also be used, such as martensitic stainless steels such as type 410, ferritic stainless steels such as type 430. Other high performance alloys could also be considered such as superaustenitic stainless steels such as AL-6XN, duplex steels such as alloy 2205 and superferritic grades. Nickel based alloys will also be useful in certain environments and include the Ni-Cu, Ni-Cr-Fe, and Ni-Cr-Mo-Fe families of alloys. Cobalt and titanium alloys are also possible as well as pure metal inserts. Any metal or alloy that can be electropolished or otherwise smoothed is a possible material for use as an insert. The metals described herein are not limited to use with the insert; rather, it is contemplated that the materials may be used to form at least a portion of the heat transfer component.

[0069] Electropolishing or other means of reducing the surface roughness, as described above, may be used either before or after the tube with the insert is installed in the heat exchanger bundle to obtain the desired surface roughness. In some instances, it might be

most advantageous to install the inserts into tubes that are already themselves installed within a heat exchanger and then electropolish the inserts. In other cases, it might be preferred to produce a stockpile of heat exchanger tubes with installed inserts and electropolish these before they are rolled into the tubesheets. The former scenario might be appropriate to retrofit a relatively new heat exchanger bundle in which the tubes are still in good shape or in construction of a new heat exchanger bundle, whereas the latter might be appropriate for retrofitting an older bundle for which re-tubing might be in order.

[0070] As is currently the case with installing inserts into heat exchanger tubes, good metal-to-metal contacting must be insured in order to minimize heat transfer losses that could occur due to air gaps between the outer tube and the tube liner. In all cases, the tube inner diameter must be as clean as possible and free of solids or liquids before the insert is expanded. Even with cleaning, a layer of corrosion product may exist. Ensuring a clean surface is relatively straightforward for new tubing, but can be more problematic for used tubing. Hydroblasting of the used tube, drying and light mechanical honing may be required. Following hydrostatic expansion of the insert, mechanical rolling of the ends of the inserts is also required to produce a good mechanical seal between the insert and the outer tube. A tube would be similarly prepared for use with an "outer" insert or sleeve.

[0071] A variation of the present invention will now be described in greater detail in connection with Fig. 3. Fig. 3 illustrates an aluminum or aluminum alloy coated carbon steel that may be effective in reducing corrosion and mitigating fouling. A carbon steel layer 31 is coated or clad with an aluminum layer 32. The aluminum layer or aluminum alloy may be applied by immersion of the steel in molten aluminum or aluminum alloy or by thermal spraying of aluminum powder or wire that is atomized. When used in a tube 14, the aluminum layer 32 is located on both the inner diameter surface and the outer diameter surface of the tube 14 similar to the Cr-enriched layer 22.

[0072] In accordance with another aspect of the invention, the heat transfer component, such as exchange tubes, can be constructed from a silicon-containing steel composition containing an alloy and a non-metallic film formed on the surface of the alloy. The alloy is a composition formed of the materials η , θ , and ι . In this case, η is a metal selected from the group consisting of Fe, Ni, Co, and mixtures thereof. θ is Si. Component ι is at least one alloying element selected from the group consisting of Cr, Al, Mn, Ti, Zr, Hf, V, Nb, Ta, Mo, W, Sc, La, Y, Ce, Ru, Rh, Ir, Pd, Pt, Cu, Ag, Au, Ga, Ge, As, In, Sn, Sb, Pb, B, C, N, P, O, S and mixtures thereof. The non-metallic film comprises sulfide, oxide, carbide, nitride, oxysulfide, oxycarbide, oxynitride and mixtures thereof, and is formed on top of the alloy.

[0073] The non-metallic film formed on the surface of an alloy contains at least one of a Si-partitioned oxide, sulfide, carbide, nitride, oxysulfide, oxycarbide, oxynitride and mixtures thereof. The temperature at which the non-metallic film forms varies. In a late-train heat exchanger applications, the non-metallic film forms at temperatures up to 400°C. In applications in a furnace or outside the late-train heat exchanger, the non-metallic film forms at temperatures up to 600°C. In petrochemical applications including use in steam cracker and reformer tubes, the non-metallic film forms at temperatures up to 1100°C. The temperatures utilized during the formation of the non-metallic film will be dependent on the metallurgy of the steel being acted upon. The skilled artisan can easily determine the upper temperature constraints based on the steel's metallurgy. . The Si-partitioned oxide or oxysulfide film effectively retards iron transport in the non-metallic film (e.g iron sulfide corrosion scale), thus sulfidation corrosion is substantially mitigated. Optionally, further surface smoothing of the alloy surface of a silicon-containing steel composition can provide the heat exchange surface, such as heat exchanger tubes in refinery applications, with superior fouling resistance.

[0074] In particular, for the alloy, the metal η , selected from the group consisting of Fe, Ni, Co, and mixtures thereof, can have a concentration that ranges from at least about 60 wt. % to about 99.98 wt. %, preferably at least about 70 wt. % to about 99.98 wt. %, and more preferably at least about 75 wt. % to about 99.98 wt. %. It is preferred that the metal η is Fe. It is well known that pure Fe has much better sulfidation resistance than pure Ni and pure Co.

[0075] The metal θ is Si. The alloy contains Si at least about 0.01 wt. % to about 5.0 wt. %, and preferably at least about 0.01 wt. % to about 3.0 wt. %. The metal Si in the alloy promotes formation of Si-partitioned oxide of oxysulfide in the non-metallic film formed on top of the alloy when it is exposed to a high temperature stream, such as a crude oil stream at temperatures up to 400°C or higher.

[0076] The metal ι is at least one alloying element selected from the group consisting of Cr, Al, Mn, Ti, Zr, Hf, V, Nb, Ta, Mo, W, Sc, La, Y, Ce, Ru, Rh, Ir, Pd, Pt, Cu, Ag, Au, Ga, Ge, As, In, Sn, Sb, Pb, B, C, N, P, O, S and mixtures thereof. The concentration of metal ι of the alloy ranges from at least about 0.01 wt. % to about 30.0 wt. %, preferably at least about 0.01 wt. % to about 30.0 wt. %, and more preferably at least about 0.01 wt. % to about 25.0 wt. %.

[0077] It is preferred that the metal ι is Al. The alloy contains Al at least about 0.01 wt. % to about 5.0 wt. %, and preferably at least about 0.01 wt. % to about 3.0 wt. %. Al provides the alloy with synergistic sulfidation corrosion resistance. Thus, the alloy in this case contains Si in an amount of at least about 0.01 wt. % to about 5.0 wt. %, and preferably at least

about 0.01 wt. % to about 3.0 wt. %, and Al in an amount of at least about 0.01 wt. % to about 5.0 wt. %, and preferably at least about 0.01 wt. % to about 3.0 wt. %.

[0078] The metal τ can also include Cr. In that case, the alloy contains Cr at least about 0.01 wt. % to about 40.0 wt. %. Cr provides the alloy with synergistic sulfidation corrosion resistance. Thus, the alloy according to this composition will contain Si in an amount of at least about 0.01 wt. % to about 5.0 wt. %, and preferably at least about 0.01 wt. % to about 3.0 wt. %, and Cr in an amount of at least about 0.01 wt. % to about 40.0 wt. % and preferably at least about 0.01 wt% to about 30.0 wt%.

[0079] The metal τ can also be a combination of elements. For example, the alloy can contains Si (component θ) in an amount of at least about 0.01 wt. % to about 5.0 wt. %, and preferably at least about 0.01 wt. % to about 3.0 wt. %, Al in an amount of at least about 0.01 wt. % to about 5.0 wt. %, and preferably at least about 0.01 wt. % to about 3.0 wt. %, and Cr in an amount of at least about 0.01 wt. % to about 40.0 wt. % and preferably at least about 0.01 wt% to about 30.0 wt%.

[0080] A non-limiting example of the alloy having the composition η , θ , and τ is listed in Table 2, below.

Alloy Name	Wt.% of Elements
EM-1001	Balanced Fe:3.0Si:0.1C
EM-1002	Balanced Fe:2.5Si:0.5Mn:0.15C
EM-1003	Balanced Fe:1.5Si:5.0Cr:0.5Mo:0.3Mn:0.15C:0.04P:0.03S
EM-1004	Balanced Fe:1.5Si:5.0Cr:0.5Mo:0.5Mn:0.1C
EM-1005	Balanced Fe:2.5Si:5.0Cr:0.5Mo:0.5Mn:0.1C
EM-1006	Balanced Fe:2.5Si:5.0Cr:0.5Al:0.1C
EM-1007	Balanced Fe:2.5Si:5.0Cr:0.5Al:0.5Mo:0.5Mn:0.1C
EM-1008	Balanced Fe:2.5Si:5.0Cr:0.5Al:0.5Mo:0.5Mn:0.5Ti:0.1C

[0081] The non-metallic film formed on the surface of the alloy comprises sulfide, oxide, carbide, nitride, oxysulfide, oxycarbide, oxynitride and mixtures thereof. The non-metallic film can comprise at least a 1 nm thick Si-partitioned non-metallic film and can consist of at least 10 atomic percent Si based on the concentration of the non-metallic film. Preferably, the Si-partitioned non-metallic film is an oxide or oxysulfide. The Si-partitioned oxide or oxysulfide film effectively retards iron transport, thus sulfidation corrosion is substantially mitigated. The Si-partitioned oxide or oxysulfide film is preferably formed on the exposed surface of the heat transfer component (e.g., heat exchanger or insert), for example on one or both of the exposed exterior surface and the interior surface of a late-train crude preheat exchanger tube.

[0082] The non-metallic film can be formed in-situ within the heat transfer component. The initial non-metallic film is preferably formed by exposing the alloy to a crude oil stream at high temperatures up to 1100°C.

[0083] Thus, the non-metallic film is formed on the surface of the alloy to construct a surface that reduces sulfidation corrosion and reduces depositional fouling, in heat exchangers for example, especially in late-train crude preheat exchangers. The material forms a surface having a surface roughness of less than 40 micro inches, preferably less than 20 micro inches and more preferably less than 10 micro inches.

[0084] The non-metallic film can be formed on the inner diameter (ID), the outer diameter (OD) or both the ID and the OD of the alloy, depending on the need for mitigating corrosion and fouling. The non-metallic film is formed on the surface of the alloy by exposing the ID, OD, or both the ID and OD of the alloy to a high temperature, as described above.

[0085] The non-metallic film also can be formed before exposing the alloy of a heat transfer component in a crude oil stream at high temperatures up to 400°C. The non-metallic film can be formed on the surface of the alloy by exposing the alloy to a low oxygen partial pressure environment at a temperature of from about 300°C to 1100°C for a time sufficient to effect the formation of the non-metallic film. Preferably the non-metallic film comprises at least a 1 nm thick Si-partitioned non-metallic film and consisting of at least 10 atomic percent Si based on the concentration of the non-metallic film on the surface of the alloy.

[0086] A low oxygen partial pressure environment can be generated from gases selected from the group consisting of CO₂, CO, CH₄, NH₃, H₂O, H₂, N₂, Ar, He and mixtures thereof. As a non-limiting example, CO₂/CO and H₂O/H₂ gas mixtures can be used. A time sufficient to effect the formation of a non-metallic film comprising at least a 1 nm thick Si-partitioned non-metallic film and consisting of at least 10 atomic percent Si based on the concentration of the non-metallic film on the surface of the alloy ranges from 1 min to 100 hrs. The thickness of the non-metallic film ranges from at least about 1 nm to about 100 μm, preferably from at least about 10 nm to about 50 μm, more preferably from at least about 100 nm to about 10 μm. The non-metallic film provides superior corrosion and fouling resistance beneficial in heat exchanger tubes in refinery applications.

[0087] The non-metallic film may be formed on the surface of the alloy by the bright annealing method. Bright annealing is an annealing process that is carried out in a controlled atmosphere furnace or a vacuum furnace to provide low oxygen partial pressure environments in order that oxidation is reduced to a minimum and the surface remains relatively bright. The

process conditions such as atmosphere, temperature, time and heating/cooling rate utilized during the bright annealing process are dependent on the metallurgy of the alloy being acted upon. The skilled artisan can easily determine the conditions based on the alloy's metallurgy. As a non-limiting example, the bright annealing can be done in either pure hydrogen or argon or dissociated ammonia, provided that the dew point of the atmosphere is less than -40°C . Bright annealing temperatures usually are above about 1038°C . Time at temperature is often kept short to hold surface scaling to a minimum or to control grain growth. Vacuum furnaces can generally achieve the best atmospheric quality for bright annealing purposes. Vacuum levels in the furnace must be better than 1×10^{-3} Torr. Fast cooling in vacuum furnaces is generally achieved by back filling the chamber with argon or nitrogen and then re-circulating this gas at high velocity through a heat exchanger to remove heat.

[0088] Fig. 10 shows a heat exchange component, in this case a pipe 200, having a base 202 formed of the alloy η , θ , and ι and the non-metallic film 204 formed on the surface of the alloy base on the ID.

[0089] Fig. 11 shows a heat exchange component, in this case a pipe 200, having a base 202 formed of the alloy η , θ , and ι and the non-metallic film 204 formed on the surface of the alloy base on the OD.

[0090] Fig. 12 shows a heat exchange component, in this case a pipe 200, having a base 202 formed of the alloy η , θ , and ι and the non-metallic film 204 formed on the surface of the alloy base on both the ID and the OD.

[0091] Examples of preparing a sample in accordance with the composition disclosed herein follow.

Example 1

[0092] Silicon-containing steels listed in above table are prepared by arc melting. The arc melted steels are hot rolled into thick sheets of about 1/2 inch thickness. The sheets are annealed at 1100°C overnight in inert argon atmosphere and furnace-cooled to room temperature. Rectangular samples of 0.5 inch x 0.25 inch are cut from the sheets. The sample faces are polished to either 600 grit finish or Linde B ($0.05 \mu\text{m}$ alumina powder) finish and cleaned in acetone. The sample is exposed to 60:40 vol.% of a crude mix (e.g. 60 vol.% Maya and 40 vol.% Olmeca crude mix) at 400°C for 4 hours in a tubing bomb test apparatus. After testing, the specimen is cleaned in toluene and acetone sequentially and characterized by selected analytical instruments.

[0093] Both surface and cross sectional images of the tested specimen are examined using a Scanning Electron Microscopy (SEM). The atomic percent of elements in the silicon-containing steel composition is determined by standard Auger Electron Spectroscopy (AES) analyses. A focused electron beam irradiates a specimen surface and produces Auger electrons, whose energies are characteristic of the element from which they are generated. Compositional depth profiling of elements is accomplished by using an independent ion beam to sputter the sample surface while using AES to analyze each successive depth.

Example 2

[0094] The commercially available ALCOR Hot Liquid Process Simulator (HLPS) is used to evaluate the relative fouling potentials of the crude oils or blends described in the examples below. The test unit procedure designed for this testing is as follows.

[0095] In accordance with standard ALCOR HLPS test procedure, ALCOR runs are carried out by charging the one-liter reservoir with a crude oil or blend, heating the liquid (up to 150°C) and pumping it across a vertically positioned, carbon-steel rod with a flow rate of 3.0 mL/minute. The spent oil is collected in the top section of the ALCOR reservoir, which is separated from the untreated oil by a sealed piston, thereby allowing for once-through operation. The system is pressurized with nitrogen (400-500 psig) prior to each test run to ensure gases remain dissolved in the oil during the test. The rod is electrically heated to a preset temperatures and held constant throughout the run. The rod surface temperature used for these tests is 275°C. Thermocouple readings are recorded for the bulk fluid inlet temperature, outlet temperature (T_{outlet}), and the temperature for the surface of the rod (T_{rod}). The heated surface thermocouple is positioned inside the rod.

[0096] During the fouling tests, asphaltenes deposit on the heated surface and are thermally degraded to coke, which build up on the surface of the test rod. The coke deposit causes an insulating effect that reduces the efficiency and/or ability of the heated surface to heat the oil passing over it. The resulting reduction in outlet bulk fluid temperature continues over time as more foulant builds up on the surface. This reduction in temperature is referred to as the outlet liquid Delta T and is dependent on the type of crude oil/blend, testing conditions and other effects. Thus, Delta T is expressed as:

$$[0097] \Delta T = T_{\text{outlet}} - T_{\text{outlet max}}$$

[0098] Delta T measures heat transfer of the foulant layer. The Dimensionless Delta T is expressed as:

$$[0099] \text{Dimensionless } \Delta T = (T_{\text{outlet}} - T_{\text{outlet max}}) / (T_{\text{rod}} - T_{\text{outlet max}}).$$

[00100] Dimensionless ΔT corrects for heat transfer characteristics of the oil tested.

[00101] The test time for these runs is 180 minutes. Noteworthy is that the flow regime for the ALCOR system is laminar and therefore direct correlation with field experience is difficult. However, the unit has been proven effective in evaluating differences in relative fouling potentials between crude oils and blends.

[00102] The ALCOR unit standard fouling test parameters and operating test conditions used for whole crudes/blends are summarized below.

Flow Rate/Type: 3.0 mL/minute / once through operation
Metallurgy: carbon-steel (1018), alloy 1, alloy 4, heater rods
System Pressure: 400-500 psi
Rod Surface Temperature(s): 275°C
System Temperature Setting (reservoir, pump and lines): 150°C
Actual Bulk Fluid Inlet Temperature: 105-120°C
Time: 30 minutes of stirring and pre-heat within reservoir are allowed prior to the start of run.

[00103] The ALCOR test rod specimen includes silicon-containing steels, alloy 1 and alloy 4, that are prepared by arc melting. The arc melted alloys are hot rolled into thick sheets of about 1/2 inch thickness. The sheets are annealed at 1100°C overnight in inert argon atmosphere and furnace-cooled to room temperature. The ALCOR rod specimen is machined from the sheets. The sample faces are polished to either 600 grit finish or Linde B (0.05 μm alumina powder) finish and cleaned in acetone.

[00104] For the first specimen, alloy 1 is exposed to 60:40 vol.% of a crude mix at 400°C for 4 hours in a tubing bomb test apparatus. After testing, the specimen is characterized by SEM. Figure 13 depicts a surface and cross sectional images of the tested specimen. A non-metallic film comprising outer iron sulfide (Fe_{1-x}S) and inner Si-partitioned oxysulfide is observed. The non-metallic film is formed on the silicon-containing steel surface. The same tested specimen is characterized by AES.

[00105] Figure 14 depicts AES depth profile from the top surface of the tested specimen. The concentration of various elements (O, S, C, Fe, and Si) in atomic % is plotted as a function of sputter depth in nm. The non-metallic film in this case appears to be a mixture of outer iron sulfide (Fe_{1-x}S) and inner Si-partitioned oxysulfide. The thickness of outer iron sulfide film is about 800 nm, and the thickness of the inner Si-partitioned oxysulfide film is about 1000 nm. The Si concentration of the inner Si-partitioned oxysulfide film varies through thickness, but is as high as 14 atomic %. The thickness of the Si-partitioned oxysulfide film

which contains at least 10 atomic percent Si based on the concentration of the non-metallic film is about 500 nm.

[00106] In the second specimen, following the ALCOR HLPS test method described above, the ALCOR rod made out of alloy 1 is tested in the ALCOR unit. Figure 15 depicts the test results of alloy 1 (shown as Fe-3.0Steel in the figure) along with standard rod (shown as 1018CS in the figure). Dimensionless ΔT of alloy 1 remains almost flat for 180 minutes of testing time. This result suggests depositional fouling is substantially reduced by use of the silicon-containing steel composition of this instant invention. In comparison, Dimensionless ΔT of 1018CS decreases with time for 180 minutes of testing time. This result suggests that asphaltenes of the crude mix deposit on the heated ALCOR rod surface and are thermally degraded to coke, which build up on the surface of the test rod.

[00107] In the third specimen, following the ALCOR HLPS test method described above, the ALCOR rod made out of alloy 4 is tested in the ALCOR unit. Figure 16 depicts the test results of alloy 4 (shown as Bal.Fe:5Cr:1.5Si in the figure) along with standard rod (shown as 1018CS in the figure). Dimensionless ΔT of alloy 4 remains almost flat for 180 minutes of testing time. This result suggests depositional fouling is substantially reduced by use of the silicon-containing steel composition of this invention. In comparison, Dimensionless ΔT of 1018CS decreases with time for 180 minutes of testing time. This result suggests that asphaltenes of the crude mix deposit on the heated ALCOR rod surface and are thermally degraded to coke, which build up on the surface of the test rod.

[00108] The surface treatment can be used in any heat exchange component or with any surface susceptible to fouling or corrosion. It is also possible to use the treatment with other mechanisms for reducing fouling and corrosion. For example, the surface treatment can be used with heat exchange components that are also subject to vibration or pulsed fluid flow, which are mechanisms used to disrupt fouling and corrosion deposition.

[00109] In one exemplary embodiment, the surface treatment can be applied to a heat exchange component in the form of a heat exchanger tube, especially those formed as tube bundles retained in a heat exchanger housing. It is also contemplated that the surface treatment can be used in replacement tubes or in tube sheaths used to cover an inner or outer diameter of a corroded or fouled tube or a new tube as well, depending upon the situation.

[00110] In accordance with another aspect of the invention, the heat transfer component, such as heat exchange tubes or inserts, can be constructed from a chromium-enriched oxide containing the material having a composition containing δ , ϵ , and ζ . In this case, ζ is a steel containing at least about 5 to about 40 wt. % chromium. ϵ is a layer formed on

the surface of the steel ζ and is a chromium-enriched oxide (M_3O_4 or M_2O_3 or mixtures thereof) wherein the metal M contains at least 5 wt. % Cr based on the total weight of the metal M. δ is a top layer formed on the surface of the chromium-enriched oxide ε comprising sulfide, oxide, oxysulfide, and mixtures thereof.

[00111] The steel ζ contains at least about 5 to about 40 wt. % chromium and is selected from low chromium steels, ferritic stainless steels, martensitic stainless steels, austenetic stainless steels, duplex stainless steels and precipitation-hardenable alloys. A non-limiting example of the base metal ζ is listed in Table 3, below.

Base Metal, ζ	Alloy	UNS No.	Alloy Compositions (Weight %)
Low chromium steels	T5	S50100	Bal.Fe:5Cr:0.5Mo, 0.5Si, 0.3Mn, 0.15C, 0.04P, 0.03S
	T9	J82090	Bal.Fe:9Cr:1.0Si, 0.35Mn, 0.02C, 0.04P, 0.045S
Ferritic stainless steels	409	S40900	Bal.Fe:10.5Cr:1.0Si, 1.0Mn, 0.5Ni, 0.5Ti, 0.08C, 0.045P, 0.045S
	410	S41000	Bal.Fe:11.5Cr:0.15C, 0.045P, 0.03S
	430	S43000	Bal.Fe:16.0Cr:1.0Si, 1.0Mn, 0.12C, 0.045P, 0.03S
Austenetic stainless steels	304	S30400	Bal.Fe: 8Ni:18Cr:2.0Mn, 0.75Si, 0.08C, 0.04P, 0.03S
	310	S31000	Bal.Fe:19Ni:24Cr:2.0Mn, 1.5Si, 0.75Mo, 0.25C, 0.045P, 0.03S
	253 MA	S30815	Bal.Fe:11Ni:21Cr:1.7Si, 0.04Ce, 0.17N, 0.08C
	RA8 5H	S30615	Bal.Fe:14.5Ni:18.5Cr:3.5Si:1.0Al, 0.2C
Duplex stainless steels	2205	S32205	Bal.Fe:4.5Ni:22Cr:2.0Mn, 1.0Si, 3.0Mo, 0.03C, 0.14N, 0.03P, 0.02S
	2507	S32507	Bal.Fe:6Ni:24Cr:1.2Mn, 0.8Si, 3.0Mo, 0.5Cu, 0.03C, 0.2N, 0.035P, 0.02S

[00112] The chromium-enriched oxide ε (M_3O_4 or M_2O_3 or mixtures thereof) can be formed on the surface of the steel ζ before use of the chromium-enriched oxide containing material, for example in a late-train crude preheat exchanger. The metal M of the chromium enriched oxide ε (M_3O_4 or M_2O_3 or mixtures thereof) preferably contains at least 5 wt. % Cr, preferably at least 10 wt. % Cr, based on the total weight of the metal M. The metal M of the chromium enriched oxide ε (M_3O_4 or M_2O_3 or mixtures thereof) preferably comprises Fe, Cr, and constituting elements of the steel ζ . Constituting elements of the steel ζ contain Ni, Co, Mn, Al, Si, B, C, N, P, Y, Ce, Ti, Zr, Hf, V, Nb, Ta, Mo, W, and mixtures thereof. The metal M of the chromium enriched oxide ε (M_3O_4 or M_2O_3 or mixtures thereof) is preferred to

be Cr-rich. Excessive partitioning of Cr in the chromium enriched oxide ϵ (M_3O_4 or M_2O_3 or mixtures thereof) attributes to superior corrosion resistance of the steel ζ . Chromium preferably enters to vacancy sites of the oxide lattice and effectively suppresses iron transport from the steel, a prerequisite for sustaining corrosion.

[00113] The chromium enriched oxide ϵ (M_3O_4 or M_2O_3 or mixtures thereof) is formed on the surface of the steel ζ by exposing the steel ζ to a low oxygen partial pressure environment at a temperature of from about 300°C to 1100°C for a time sufficient to effect the formation of a chromium enriched oxide (M_3O_4 or M_2O_3 or mixtures thereof) on the surface of the steel. Thermodynamically, an oxide forms on a metal surface when the oxygen potential in the environment is greater than the oxygen partial pressure in equilibrium with the oxide. For example, the partial pressure of oxygen in equilibrium with Cr_2O_3 at 1000°C is about 10^{-21} atm. This implies that the formation of Cr_2O_3 is favored thermodynamically at 1000°C in environments with oxygen partial potentials higher than 10^{-21} atm.

[00114] A low oxygen partial pressure environment can be generated from gases selected from the group consisting of CO_2 , CO, CH_4 , NH_3 , H_2O , H_2 , N_2 , Ar, He and mixtures thereof. As a non-limiting example, CO_2/CO and H_2O/H_2 gas mixtures can be used. A time sufficient to effect the formation of the chromium enriched oxide ϵ (M_3O_4 or M_2O_3 or mixtures thereof) on the surface of the steel ranges from 1 min to 100 hrs. The thickness of the chromium-enriched oxide ϵ (M_3O_4 or M_2O_3 or mixtures thereof) ranges from at least about 1 nm to about 100 μm , preferably from at least about 10 nm to about 50 μm , more preferably from at least about 100 nm to about 10 μm . The chromium-enriched oxide ϵ provides superior corrosion and fouling resistance beneficial in heat exchanger tubes in refinery applications.

[00115] The chromium enriched oxide ϵ (M_3O_4 or M_2O_3 or mixtures thereof) may be formed on the surface of the steel ζ by bright annealing method. Bright annealing is an annealing process that is carried out in a controlled atmosphere furnace or vacuum to provide low oxygen partial pressure environments in order that oxidation is reduced to a minimum and the surface remains relatively bright. The process conditions such as atmosphere, temperature, time and heating/cooling rate utilized during the bright annealing process are dependent on the metallurgy of the alloy being acted upon. The skilled artisan can easily determine the conditions based on the alloy's metallurgy. As a non-limiting example, the austenitic stainless steel such as 304L can be bright annealed in either pure hydrogen or argon or dissociated ammonia, provided that the dew point of the atmosphere is less than -40°C and the tubes, upon entering the furnace, are dry and scrupulously clean. Bright annealing temperatures usually are above about 1038°C. Time at temperature is often kept short to hold surface scaling to a

minimum or to control grain growth. Vacuum furnaces can generally achieve the best atmospheric quality for bright annealing purposes. Vacuum levels in the furnace must be better than 1×10^{-3} Torr. Fast cooling in vacuum furnaces is generally achieved by back filling the chamber with argon or nitrogen and then re-circulating this gas at high velocity through a heat exchanger to remove heat.

[00116] For the top layer δ , sulfide and oxysulfide form on the surface of the chromium-enriched oxide ϵ in-situ during use of the chromium-enriched oxide containing material, in a late-train crude preheat exchanger for instance. The temperature at which the protective layer forms varies. In a late-train heat exchanger applications, the protective layer forms at temperatures up to 400°C . In applications in a furnace or outside the late-train heat exchanger, the protective layer forms at temperatures up to 600°C . In petrochemical applications including use in steam cracker and reformer tubes, the protective layer forms at temperatures up to 1050°C . The temperatures utilized during the formation of the protective layer will be dependent on the metallurgy of the steel being acted upon. The skilled artisan can easily determine the upper temperature constraints based on the steel's metallurgy. The top layer δ may contain some impurities formed from the elements constituting the chromium enriched oxide ϵ and the steel ζ . Preferably, the top layer δ comprises iron sulfide (Fe_{1-x}S), iron oxide (Fe_3O_4), iron oxysulfide, iron-chromium sulfide, iron-chromium oxide, iron-chromium oxysulfide, and mixtures thereof. The thickness of the top layer δ ranges from at least about $0.1\ \mu\text{m}$ to about $50\ \mu\text{m}$, preferably from at least about $0.1\ \mu\text{m}$ to about $30\ \mu\text{m}$, more preferably from at least about $0.1\ \mu\text{m}$ to about $10\ \mu\text{m}$.

[00117] Thus, the chromium-enriched oxide containing material so formed is utilized to construct a surface that reduces sulfidation corrosion and reduces depositional fouling, in heat exchangers for example, especially in late-train crude preheat exchangers. The surface has a surface roughness of less than 40 micro inches ($1.1\ \mu\text{m}$), preferably less than 20 micro inches ($0.5\ \mu\text{m}$), and more preferably less than 10 micro inches ($0.25\ \mu\text{m}$). The chromium-enriched oxide ϵ (M_3O_4 or M_2O_3 or mixtures thereof) can be formed on the inner diameter (ID), the outer diameter (OD) or both the ID and the OD of the steel ζ , depending on the need for mitigating corrosion and fouling. The chromium enriched oxide ϵ (M_3O_4 or M_2O_3 or mixtures thereof) is formed on the surface of the steel ζ by exposing the ID, OD or both the ID and OD of the steel ζ , as described above.

[00118] Fig. 17 shows a heat exchange component, in this case a pipe 100, having a base 102 formed of steel ζ with a layer 104 of chromium-enriched oxide ϵ and top layer 106 formed as a coating δ on the chromium-enriched oxide layer on the ID.

[00119] Fig. 18 shows a heat exchange component, in this case a pipe 100, having a base 102 formed of steel ζ with a layer 104 of chromium-enriched oxide ϵ and top layer 106 formed as a coating δ on the chromium-enriched oxide layer on the OD.

[00120] Fig. 19 shows a heat exchange component, in this case a pipe 100, having a base 102 formed of steel ζ with a layer 104 of chromium-enriched oxide ϵ and top layer 106 formed as a coating δ on the chromium-enriched oxide layer on both the ID and the OD.

Example 1

[00121] A first example of forming the surface is as follows. A low chromium steel, T5, specimen is exposed to a gas mixture containing 50% H_2O and 50% He at 566°C with a flow rate of about 500 cc/min. The specimen is heated to 566°C at a rate of 20 °C/min, held for 24 hours, and cooled down to room temperature. After heat treatment of the specimen in a low oxygen partial pressure environment, the specimen is characterized by Scanning Electron Microscopy (SEM).

[00122] Fig. 20 depicts a surface and cross sectional images of the heat treated specimen. A top layer comprising magnetite (Fe_3O_4) and a chromium-enriched oxide (M_3O_4) can be observed. The concentration of the metal M of a chromium enriched oxide (M_3O_4) is Balanced Fe:12.0 Cr:1.5Si in wt.%. The thickness of the top Fe_3O_4 layer is about 5 μm , and the thickness of a chromium-enriched oxide (M_3O_4) is about 4 μm .

Example 2

[00123] In a second example, the specimen as prepared in Example 1 is used for corrosion and fouling testing. The specimen is exposed to 60:40 vol.% of a crude mix at 400°C for 4 hours in a tubing bomb test apparatus. After testing, the specimen is cleaned in toluene and acetone sequentially and characterized by SEM.

[00124] Fig. 21 depicts a surface and cross sectional images of the tested specimen. A top layer is comprised of iron sulfide (Fe_{1-x}S) and magnetite (Fe_3O_4). The thickness of the top layer is about 5 μm . A chromium-enriched oxide (M_3O_4) is retained. The concentration of the metal M of a chromium enriched oxide (M_3O_4) is Balanced Fe:12.0 Cr:1.5Si in wt.%. The thickness of a chromium-enriched oxide (M_3O_4) is about 4 μm .

[00125] The surface treatment can be used in any heat exchange component or with any surface susceptible to fouling or corrosion. It is also possible to use the treatment with other mechanisms for reducing fouling and corrosion. For example, the surface treatment can

be used with heat exchange components that are also subject to vibration or pulsed fluid flow, which are mechanisms used to disrupt fouling and corrosion deposition.

[00126] In one exemplary embodiment, the surface treatment can be applied to a heat exchange component in the form of a heat exchanger tube, especially those formed as tube bundles retained in a heat exchanger housing. It is also contemplated that the surface treatment can be used in replacement tubes or in tube sheaths used to cover an inner or outer diameter of a corroded or fouled tube.

[00127] It is another aspect of the present invention to combine heat transfer components having the above described corrosion resistant materials with either a pulsation generating device or a vibrational generating device to further reduce and mitigate fouling. The devices are generically designated in Fig. 1 with reference numeral 3.

[00128] It is contemplated that the pulsation device will comprise any means for applying liquid pressure pulsations to the tube side liquid. In the simplest concept, the device may comprise a reciprocating pump type mechanism with a cylinder connected to the inlet/outlet conduits of the exchanger and a reciprocating piston in the cylinder to vary the internal volume of the cylinder. As the piston moves within the cylinder, the liquid will alternately be drawn into the cylinder and then expelled from it, creating pulsations in the conduit to which the device is connected. The use of a double-acting pump of this kind with one side connected to the inlet conduit and the other connected to the outlet conduit is particularly desirable since it will create the desired pressure pulsations in the tubes regardless of the pressure drop occurring in the exchanger tube bundle. Variation in the frequency of the pulsations may be afforded by variations in the reciprocation speed of the piston and any desired variations in pulsation amplitude may be provided by the use of a variable displacement pump, e.g. a variable displacement piston pump, swashplate (stationary plate) pump and its variations such as the wobble plate (rotary plate) pump or bent axis pump.

[00129] The present invention is not intended to be limited to the above-described pump; rather, it is contemplated that other types of pumps may also be used as the pulsation device including diaphragm pumps and these may be practically attractive since they offer the potential for activation of the diaphragm by electrical, pneumatic or direct mechanical means with the movement of the diaphragm controlled to provide the desired frequency and amplitude (by control of the extent of diaphragm movement). Other types may also be used but gear pumps and related types such as the helical rotor and multi screw pumps which give a relatively smooth (non pulsating) fluid flow are less preferred in view of the objective of introducing pulsations which disrupt the formation of the troublesome boundary layer. Other

types which do produce flow pulsations such as the lobe pump, the vane pump and the similar radial piston pump, are normally less preferred although they may be able to produce sufficient pulsation for the desired purpose. Given the objective is to induce pulsations, other types of pulsator may be used, for example, a flow interrupter which periodically interrupts the liquid flow on the tube side. Pulsators of this type may include, for example, siren type, rotary vane pulsators in which the flow interruption is caused by the repeated opening and closing of liquid flow passages in a stator/rotor pair, each of which has radial flow openings which coincide with rotation of the moving rotor member. The rotor may suitably be given impetus by the use of vanes at an angle to the direction of liquid flow, e.g. by making radial cuts in the rotor disc and bending tabs away from the plane of the disc to form the vanes. Another type is the reed valve type with spring metal vanes which cover apertures in a disc and which are opened temporarily by the pressure of the fluid in the tube, followed by a period when the vane snaps closed until fluid pressure once more forces the vane open.

[00130] In order to optimize the impact within the heat transfer component, it is preferable to locate the pulsation device close to the exchanger in order to ensure that the pulsations are efficiently transferred to the liquid flow in the tube bundle, that is, the pulsations are not degraded by passage through intervening devices such as valves. Normally, the frequency of the liquid pulsations will be in the range of 0.1 Hz to 20 kHz. The amplitude of the pulsation as measured by the incremental flow rate through the heat exchange tubes could range from about the order of the normal heat exchanger flow rate at the lower end of the range of pulsation frequencies to less than 10^{-6} of the normal flow rate at the higher frequencies; because of pressure drop limitations in the heat transfer component operation and/or dissipation of higher frequencies in the fluid, the upper limit of the pulse amplitude will decrease with increased frequency. Thus, for example, in the lower half of this frequency range, the amplitude of the pulsations could be from about 10^{-2} to about the normal flow rate and with frequencies in the upper half of the range, from about 10^{-6} to 0.1 of the normal flow rate through the exchanger.

[00131] It is contemplated that the vibration producing device may be any means that is capable of imparting a vibration force on the heat exchanger unit. The vibration producing device may be of the kind disclosed in co-pending US Patent Serial No. 11/436,802. The vibration producing device may be externally connected to the heat exchange unit to impart controlled vibrational energy to the tubes of the bundle. The vibration producing device can take the form of any type of mechanical device that induces tube vibration while maintaining structural integrity of the heat exchanger. Any device capable of generating

sufficient dynamic force at selected frequencies would be suitable. The vibration producing device can be single device, such as an impact hammer or electromagnetic shaker, or an array of devices, such as hammers, shakers or piezoelectric stacks. An array can be spatially distributed to generate the desired dynamic signal to achieve an optimal vibrational frequency. The vibration producing device may be placed at various locations on or near the heat exchange unit as long as there is a mechanical link to the tubes. Sufficient vibration energy can be transferred to the tubes of the heat exchanger at different vibration modes. There are low and high frequency vibration modes of tubes. For low frequency modes (typically below 1000Hz), axial excitation is more efficient at transmitting vibration energy, while at high frequency modes, transverse excitation is more efficient. The density of the vibration modes is higher at a high frequency range than at a low frequency range (typically below 1000 Hz), and vibration energy transfer efficiency is also higher in the high frequency range. Further, displacement of tube vibration is very small at high frequency (>1000 Hz) and insignificant for potential damage to the tubes.

[00132] It will be apparent to those skilled in the art that various modifications and/or variations may be made without departing from the scope of the present invention. While the present invention has been described in the context of the heat exchanger in a refinery operation. The present invention is not intended to be so limited; rather, it is contemplated that the desired surface roughness and materials disclosed herein may be used in other portions of a refinery operation where fouling may be of a concern. Reducing the surface roughness of other corrosion resistant materials such as aluminized carbon steel, titanium, electroless nickel-coated carbon steel and other corrosion resistant surfaces are extensions of this concept as delineated below. It is contemplated that the method of reducing fouling disclosed herein can be combined with other reduction strategies to reduce fouling. This includes combining the low surface roughness and/or material compositions disclosed herein with vibrational, pulsation, helical shell-side baffles and internal turbulence promoters. Thus, it is intended that the present invention covers the modifications and variations of the method herein, provided they come within the scope of the appended claims and their equivalents.

What Is Claimed Is:

1. A heat transfer component for heating a process stream, comprising:
a housing having a wall forming a hollow interior, wherein the wall having an inner surface;
at least one heat transfer element located within the housing for heating the process stream within the hollow interior of the housing;
wherein at least one of the inner surface and the at least one heat transfer element having a surface roughness of less than 40 micro inches (1.1 μm).
2. The heat transfer component for heating a process stream, comprising:
wherein the at least one heat transfer element having an inner diameter surface and an outer diameter surface, wherein the heat transfer element being formed from a first material;
a sleeve, wherein the sleeve having a first surface adapted to contact one of the inner diameter surface and the outer diameter surface of the heat transfer element and a second surface adapted to contact the process stream, wherein the second surface having the surface roughness of less than 40 micro inches (1.1 μm).
3. The heat transfer component according to claim 2, wherein the sleeve being formed from a composition that is resistant to sulfidation corrosion and corrosion induced fouling.
4. The heat transfer component according to claim 1, wherein the at least one heat transfer element being formed from a composition that is resistant to sulfidation corrosion and corrosion induced fouling.
5. The heat transfer component according to claim 3 or 4, wherein the composition is a steel composition comprising:
X, Y, and Z,
wherein X is a metal selected from the group consisting of Fe, Ni, Co and mixtures thereof,
wherein Y is Cr, and
wherein Z is at least one alloying element selected from the group consisting of Si, Al, Mn, Ti, Zr, Hf, V, Nb, Ta, Mo, W, Sc, Y, La, Ce, Pt, Cu, Ag, Au, Ru, Rh, Ir, Ga, In, Ge, Sn, Pb, B, C, N, O, P, and S.

6. The heat transfer component according to claim 5, wherein each of the at least one heat transfer element having Cr-enriched layer formed thereon, wherein the Cr-enriched layer also being formed from the steel composition X, Y and Z, wherein the ratio of Y to X in the Cr-enriched layer being greater than the ratio of Y to X in the remaining portion of the heat transfer element.

7. The heat transfer component according to claim 6, wherein the Cr-enriched layer is formed by one of electro-polishing at least on surface of heat transfer element, electroplating, thermal spray coating, laser deposition, sputtering, physical vapor deposition, chemical vapor deposition, plasma powder welding overlay, cladding, and diffusion bonding.

8. The heat transfer component according to claim 7, further comprising:
an protective layer formed on an outer surface of the Cr-enriched layer.

9. The heat transfer component according to claim 8, wherein the protective layer comprises an oxide selected from the group consisting of a magnetite, an iron-chromium spinel, a chromium oxide, and mixtures thereof.

10. The heat transfer component according to claim 9, wherein the protective layer being formed within the at least one heat transfer element when subjected to a process stream at high temperatures up to 400°C.

11. The heat transfer component according to claim 9, wherein the protective layer being formed within the at least one heat transfer element when subjected to a process stream at high temperatures up to 600°C.

12. The heat transfer component according to claim 9, wherein the protective layer being formed within the at least one heat transfer element when subjected to a process stream at high temperatures up to 1100°C.

13. The heat transfer component according to claim 3 or 4, wherein the composition is a carbon steel having an aluminum or aluminum alloy layer located thereon.

14. The heat transfer component according to claim 3 or 4, wherein the composition is one aluminum, titanium, chromium and alloys of aluminum, titanium, chromium.

15. The heat transfer component according to claim 3 or 4, wherein one of the at least one heat transfer and the sleeve having a heat exchange surface formed from the composition, wherein the composition comprising:

a chromium-enriched oxide containing material formed from the composition δ , ϵ , and ζ , wherein,

ζ is a steel containing at least about 5 to about 40 wt. % chromium;

ϵ is a chromium enriched oxide (M_3O_4 or M_2O_3 or mixtures thereof) formed on the surface of the steel ζ , wherein M is a metal containing at least 5 wt. % Cr based on the total weight of the metal M; and,

δ is a top layer formed on the surface of the chromium-enriched oxide ϵ , comprising sulfide, oxide, oxysulfide, and mixtures thereof.

16. The heat transfer component of claim 15, wherein the top layer δ comprises iron sulfide ($Fe_{1-x}S$), iron oxide (Fe_3O_4), iron oxysulfide, iron-chromium sulfide, iron-chromium oxide, iron-chromium oxysulfide, and mixtures thereof.

17. The heat transfer component according to any one of claims 15 or 16, wherein the metal M of the chromium enriched oxide (M_3O_4 or M_2O_3 or mixtures thereof) comprises Fe, Cr, and constituting elements of the steel ζ .

18. The heat transfer component according to any one of claims 15 to 17, wherein the steel ζ is selected from low chromium steels, ferritic stainless steels, martensitic stainless steels, austenitic stainless steels, duplex stainless steels and precipitation-hardenable alloys

19. The heat transfer component according to claim 3 or 4, wherein one of the at least one heat transfer and the sleeve having a heat exchange surface formed from the composition, wherein the composition comprising:

a silicon containing steel composition including an alloy and a non-metallic film

formed on the surface of the alloy,

wherein the alloy is formed from the composition η , θ , and ι , in which η is a metal selected from the group consisting of Fe, Ni, Co, and mixtures thereof, θ is Si, and ι is at least one alloying element selected from the group consisting of Cr, Al, Mn, Ti, Zr, Hf, V, Nb, Ta, Mo, W, Sc, La, Y, Ce, Ru, Rh, Ir, Pd, Pt, Cu, Ag, Au, Ga, Ge, As, In, Sn, Sb, Pb, B, C, N, P, O, S and mixtures thereof, and

wherein the non-metallic film comprises sulfide, oxide, carbide, nitride, oxysulfide, oxycarbide, oxynitride and mixtures thereof.

20. The heat transfer component of claim 19, wherein the metal η comprises at least about 60 wt% to about 99.98 wt. % of the alloy.

21. The heat transfer component according to claim 19 or 20, wherein the alloying metal θ comprises at least about 0.01 wt% to about 5.0 wt. % of the alloy.

22. The heat transfer component according to any one of claims 19 to 21, wherein the alloying element ι comprises at least about 0.01 wt% to about 40.0 wt% of the alloy.

23. The heat transfer component according to any one of claims 19 to 22, wherein the alloying metal θ comprises at least about 0.01 wt% to about 3.0 wt. % of the alloy.

24. The heat transfer component according to any one of claims 19 to 23, wherein the alloying element ι comprises at least about 0.01 wt% to about 3.0 wt% Al of the alloy.

25. The heat transfer component according to any one of claims 19 to 24, wherein the alloying element ι comprises at least about 0.01 wt% to about 30.0 wt% Cr of the alloy.

26. The heat transfer component according to any one of claims 19 to 25, wherein the alloying element ϵ comprises at least about 0.01 wt% to about 3.0 wt% Al and at least about 0.01 wt% to about 30.0 wt% Cr of the alloy.

27. The heat transfer component according to any one of claims 19 to 26, wherein the non-metallic film on the surface of the alloy comprises at least 1 nm thick Si-partitioned non-metallic film.

28. The heat transfer component of claim 27, wherein the non-metallic film is a multi-layer non-metallic film and consists of at least 10 atomic percent Si based on the concentration of the non-metallic film.

29. The heat transfer component according to any one of the preceding claims, wherein the heat transfer component is a heat exchanger and the at least one heat transfer element is a tube bundle having a plurality of heat transfer tubes, wherein each of the heat transfer tubes having an inner diameter surface and an outer diameter surface, wherein at least one of the inner diameter surface and the outer diameter surface having a surface roughness of less than 40 micro inches (1.1 μm).

30. The heat transfer component according to any one of the preceding claims, wherein the surface roughness is less than 20 micro inches (0.5 μm).

31. The heat transfer component according to any one of the preceding claims, wherein the surface roughness is less than 10 micro inches (0.25 μm).

32. The heat transfer component according to any one of the preceding claims, further comprising:

one of a vibration producing device to impart a vibrational force to the heat transfer element and a pulsation producing device for apply pressure pulsations to the process stream.

33. A method of reducing fouling in a heat transfer component according to anyone of the preceding claims, the method comprising:

applying one of fluid pressure pulsations to the process stream flowing through the at least one heat transfer element and vibration to the heat transfer component to effect a reduction

of the viscous boundary layer adjacent the at least one heat transfer element to reduce the incidence of fouling and promote heat transfer from the heat transfer element to the process stream.

34. The method of reducing fouling according to claim 33, wherein the method comprising:

applying fluid pressure pulsations to the process stream.

35. The method of reducing fouling according to claim 33, wherein the method comprising:

applying vibration to the at least one heat transfer element.

36. A method of reducing fouling in a heat exchanger for a process stream, wherein the heat exchanger having a plurality of existing heat exchanger tubes, the method comprising:

removing at least a portion of the plurality of existing heat exchanger tubes from the heat exchanger; and

installing a plurality of replacement heat exchanger tubes, wherein each of the plurality of replacement heat exchanger tubes having a surface roughness of less than 40 micro inches (1.1 μm).

37. The method according to claim 36, wherein each of the replacement heat exchanger tubes being formed from a composition that is resistant to sulfidation corrosion and corrosion induced fouling.

38. A method of reducing fouling in a heat transfer component for a process stream, wherein the heat transfer component having at least one heat transfer element for heating the process stream, wherein the heat transfer element having an inner diameter surface and an outer diameter surface, the method comprising:

installing a sleeve on at least one of the heat transfer elements, wherein the sleeve being formed from a composition that is resistant to sulfidation corrosion and corrosion induced fouling,

wherein the sleeve having a first surface adapted to contact one of the inner diameter surface and the outer diameter surface of the heat transfer element and a second surface adapted to contact the process stream, wherein the second surface having a surface roughness of less than 40 micro inches (1.1 μm).

39. A method of providing sulfidation corrosion resistance and corrosion induced fouling resistance to a metal surface that is subject to a process stream at high temperatures, the method comprising:

providing a metal layer having a surface roughness of less than 40 micro inches (1.1. μm) formed from a composition comprising X, Y, and Z, wherein X is a metal selected from the group consisting of Fe, Ni, Co and mixtures thereof, wherein Y is Cr, and wherein Z is at least one alloying element selected from the group consisting of Si, Al, Mn, Ti, Zr, Hf, V, Nb, Ta, Mo, W, Sc, Y, La, Ce, Pt, Cu, Ag, Au, Ru, Rh, Ir, Ga, In, Ge, Sn, Pb, B, C, N, O, P, and S, wherein a Cr-enriched layer is located on the metal layer, wherein the Cr-enriched layer also being formed from the steel composition X, Y, and Z, wherein the ratio of Y to X in the Cr-enriched layer being greater than the ratio of Y to X in the metal layer; and

forming a protective layer on a surface of the Cr-enriched layer.

40. A method of providing sulfidation corrosion resistance and corrosion induced fouling resistance to a heat transfer component surface, the method comprising:

providing a silicon containing steel composition including an alloy, wherein the alloy is formed from the composition η , θ , and ι , in which η is a metal selected from the group consisting of Fe, Ni, Co, and mixtures thereof, θ is Si, and ι is at least one alloying element selected from the group consisting of Cr, Al, Mn, Ti, Zr, Hf, V, Nb, Ta, Mo, W, Sc, La, Y, Ce, Ru, Rh, Ir, Pd, Pt, Cu, Ag, Au, Ga, Ge, As, In, Sn, Sb, Pb, B, C, N, P, O, S and mixtures thereof; and,

forming a non-metallic film on the surface of the alloy having a surface roughness of less than 40 micro inches (1.1. μm), wherein the non-metallic film comprises sulfide, oxide, carbide, nitride, oxysulfide, oxycarbide, oxynitride and mixtures thereof.

41. A method of providing sulfidation corrosion resistance and corrosion induced fouling resistance to a heat transfer component surface, the method comprising:

providing a steel surface containing at least about 5 to about 40 wt. % chromium;

forming a chromium-enriched oxide (M_3O_4 or M_2O_3 or mixtures thereof) on the surface of the steel, wherein M is a metal containing at least 5 wt. % Cr based on the total weight of the

metal M; and,

forming a top layer formed on the surface of the chromium-enriched oxide having a surface roughness of less than 40 micro inches (1.1. μm), comprising sulfide, oxide, oxysulfide, and mixtures thereof.

42. The method according to claims 37 or 38, wherein the composition is a steel composition comprising:

X, Y, and Z,

wherein X is a metal selected from the group consisting of Fe, Ni, Co and mixtures thereof,

wherein Y is Cr, and

wherein Z is at least one alloying element selected from the group consisting of Si, Al, Mn, Ti, Zr, Hf, V, Nb, Ta, Mo, W, Sc, Y, La, Ce, Pt, Cu, Ag, Au, Ru, Rh, Ir, Ga, In, Ge, Sn, Pb, B, C, N, O, P, and S.

43. The method according to claim 39 or 42, wherein the composition further comprising a Cr-enriched layer formed thereon, wherein the Cr-enriched layer also being formed from the steel composition X, Y and Z, wherein the ratio of Y to X in the Cr-enriched layer being greater than the ratio of Y to X in the remaining portion of composition.

44. The method according to claim 43, wherein the Cr-enriched layer is formed by one of electro-polishing the composition, electroplating, thermal spray coating, laser deposition, sputtering, physical vapor deposition, chemical vapor deposition, plasma powder welding overlay, cladding, and diffusion bonding.

45. The method according to claim 44, further comprising:

an protective layer formed on an outer surface of the Cr-enriched layer.

46. The method according to claim 45, wherein the protective layer comprises an oxide selected from the group consisting of a magnetite, an iron-chromium spinel, a chromium oxide, and mixtures thereof.

47. The method according to claim 46, wherein the protective layer being formed when the composition is subjected to a process stream at high temperatures up to 400°C.

48. The method according to claim 47, wherein the protective layer being formed when the composition is subjected to a process stream at high temperatures up to 600°C.

49. The method according to claim 48, wherein the protective layer being formed when the composition is subjected to a process stream at high temperatures up to 1100°C.

50. The method according to claims 37 or 38, wherein the composition is a carbon steel having an aluminum or aluminum alloy layer located thereon.

51. The method according to claims 37 or 38, wherein the composition is one of aluminum, titanium, chromium and alloys of aluminum, titanium, chromium.

52. The method according to claims 37 or 38, wherein the composition comprising: a chromium-enriched oxide containing material formed from the composition δ , ϵ , and ζ , wherein,

ζ is a steel containing at least about 5 to about 40 wt. % chromium;

ϵ is a chromium enriched oxide (M_3O_4 or M_2O_3 or mixtures thereof) formed on the surface of the steel ζ , wherein M is a metal containing at least 5 wt. % Cr based on the total weight of the metal M; and,

δ is a top layer formed on the surface of the chromium-enriched oxide ϵ , comprising sulfide, oxide, oxysulfide, and mixtures thereof.

53. The method of claim 52, wherein the chromium enriched oxide (M_3O_4 or M_2O_3 or mixtures thereof) is formed on the surface of the steel by exposing the steel to a low oxygen partial pressure environment at a temperature of from about 300°C to 1100°C for a time sufficient to effect the formation of a chromium enriched oxide (M_3O_4 or M_2O_3 or mixtures thereof) on the surface of said a steel.

54. The method of claim 53, wherein the time sufficient to effect the formation of the chromium enriched oxide (M_3O_4 or M_2O_3 or mixtures thereof) on the surface of the steel ranges

from 1 min to 100 hrs.

55. The method of claim 53, wherein the low oxygen partial pressure environment is formed from gases selected from the group consisting CO₂, CO, CH₄, NH₃, H₂O, H₂, N₂, Ar, He and mixtures thereof.

56. The method of claim 53, wherein the low oxygen partial pressure environment is a gas mixture of CO₂ and CO.

57. The method of claim 53, wherein the low oxygen partial pressure environment is a gas mixture of H₂O and H₂.

58. The method of claim 53, wherein the low oxygen partial pressure environment is pure hydrogen or argon having the dew point of the atmosphere is less than -40°C.

59. The method of claim 53, wherein the low oxygen partial pressure environment is vacuum having a vacuum level better than 1×10^{-3} Torr.

60. The method according to anyone of claims 53 to 59, wherein the top layer δ comprises iron sulfide (Fe_{1-x}S), iron oxide (Fe₃O₄), iron oxysulfide, iron-chromium sulfide, iron-chromium oxide, iron-chromium oxysulfide, and mixtures thereof.

61. The method according to any one of claims 53 to 60, wherein the metal M of the chromium enriched oxide (M₃O₄ or M₂O₃ or mixtures thereof) comprises Fe, Cr, and constituting elements of the steel ζ .

62. The method according to any one of claims 53 to 61, wherein the steel ζ is selected from low chromium steels, ferritic stainless steels, martensitic stainless steels, austenitic stainless steels, duplex stainless steels and precipitation-hardenable alloys.

63. The method according to claim 37 or 38, wherein the composition comprising:
a silicon containing steel composition including an alloy and a non-metallic film formed on the surface of the alloy,

wherein the alloy is formed from the composition η , θ , and ι , in which η is a metal selected from the group consisting of Fe, Ni, Co, and mixtures thereof, θ is Si, and ι is at least one alloying element selected from the group consisting of Cr, Al, Mn, Ti, Zr, Hf, V, Nb, Ta, Mo, W, Sc, La, Y, Ce, Ru, Rh, Ir, Pd, Pt, Cu, Ag, Au, Ga, Ge, As, In, Sn, Sb, Pb, B, C, N, P, O, S and mixtures thereof, and

wherein the non-metallic film comprises sulfide, oxide, carbide, nitride, oxysulfide, oxycarbide, oxynitride and mixtures thereof.

64. The method of claim 40 or 63, wherein the metal η comprises at least about 60 wt% to about 99.98 wt. % of the alloy.

65. The method according to claim 63 or 64, wherein the alloying metal θ comprises at least about 0.01 wt% to about 5.0 wt. % of the alloy.

66. The method according to any one of claims 63 to 65, wherein the alloying element ι comprises at least about 0.01 wt% to about 40.0 wt% of the alloy.

67. The method according to any one of claims 63 to 66, wherein the alloying metal θ comprises at least about 0.01 wt% to about 3.0 wt. % of the alloy.

68. The method according to any one of claims 63 to 67, wherein the alloying element ι comprises at least about 0.01 wt% to about 3.0 wt% Al of the alloy.

69. The method according to any one of claims 63 to 68, wherein the alloying element comprises at least about 0.01 wt% to about 30.0 wt% Cr of the alloy.

70. The method according to any one of claims 63 to 69, wherein the alloying element comprises at least about 0.01 wt% to about 3.0 wt% Al and at least about 0.01 wt% to about 30.0 wt% Cr of the alloy.

71. The method according to any one of claims 63 to 70, wherein the non-metallic film on the surface of the alloy comprises at least 1 nm thick Si-partitioned non-metallic film.

72. The method of claim 71, wherein the non-metallic film is a multi-layer non-metallic film and consists of at least 10 atomic percent Si based on the concentration of the non-metallic film.

73. The method according to any one of claims 63 to 72, wherein the non-metallic film is formed by exposing the alloy to a high temperature up to 1100°C.

74. The method according to any one of claims 63 to 72, wherein the non-metallic film is formed by exposing the alloy to a process stream at high temperatures up to 1100°C.

75. The method according to any one of claims 63 to 72, wherein the non-metallic film is formed on the surface of the alloy by exposing the alloy to a low oxygen partial pressure environment at a temperature of from about 300°C to 1100°C for a time sufficient to effect the formation of a non-metallic film on the surface of said an alloy.

76. The method of claim 75, wherein the time sufficient to effect the formation of the non-metallic film on the surface of the alloy ranges from 1 min to 100 hrs.

77. The method of claim 75 or 76, wherein the low oxygen partial pressure environment is formed from gases selected from the group consisting CO₂, CO, CH₄, NH₃, H₂O, H₂, N₂, Ar, He and mixtures thereof.

78. The method of claim 75 or 76, wherein the low oxygen partial pressure environment is a gas mixture of CO₂ and CO.

79. The method of claim 75 or 76, wherein the low oxygen partial pressure environment is a gas mixture of H₂O and H₂.

80. The method of claim 75 or 76, wherein the low oxygen partial pressure environment is pure hydrogen or argon having the dew point of the atmosphere is less than -40°C.

81. The method of claim 75 or 76, wherein the low oxygen partial pressure environment is vacuum having a vacuum level better than 1×10^{-3} Torr.

82. The method according to any one of claims 36 to 81, wherein the surface roughness is less than 20 micro inches (0.5 μm).

83. The method according to any one of claims 36 to 82, wherein the surface roughness is less than 10 micro inches (0.25 μm).

84. The method according to any one of claims 36 to 83, further comprising:
imparting one of a vibrational force to the heat transfer element and pressure pulsations to the process stream.