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DEVELOPMENT OF AN IMPROVED ULTRA-HIGH STRENGTH STEEL FOR FORGED AIRCRAFT COMPONENTS

R.T. Ault 9.M. Wold R.B. Bortolo

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RESEARCH GENTER

TECHNICAL REPORT AFML-TR-71-27 FEBRUARY 1971



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Development of An Improved Ultra-High Strength

Steel for Forged Aircraft Components

R. T. Ault G. M. Waid R. B. Bertolo

Republic Steel Corporation Research Center

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The distribution of this report is limited because the report contains technology identifiable with items on the strategic subargo lists excluded from export under the U. S. Export Control Act, as implemented by AFR 310-2 and AFSCR 80-20.

FOREWARD

This final report was prepared by the Alloy Development Section, Metallurgical Division, Research Center, Republic Steel Corporation, Cleveland, Ohio, under USAF Contract F33615-69-C-1638. The contract was initiated under Project No. 7351, Metallic Materials, and Task 735105, High Strength Metallic Materials.

The work was administered under the direction of the Air Force Materials Laboratory, Metals and Ceramic Division, LLP, with H. J. Middendorp as project engineer.

This report covers work accomplished independently by Republic Steel Corporation, Research Center, from July 20, 1960 through April 30, 1969 and work accomplished under the subject contract from May 1, 1969 to December 31, 1970. The principal participants in the research were R. T. Ault, G. N. Waid, and R. B. Bertolo.

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The manuscript was released by the authors in January 1971 for publication as an AFML technical report.

This technical report has been reviewed and is approved.

I Perhantte

I. Perlmutter Chief, Metals Branch Metals and Ceramics Division Air Force Materials Laboratory

ABSTRACT

The objective of this program was to develop an ultra-high strength steel in the 300 to 320 ksi ultimate tensile strength range, with improved fatigue strength, fracture toughness, and stress corrosion resistance for greater reliability in forged landing gear components. Alloy development studies were conducted on two bainitic alloy systems and two martensific alloy systems in order to develop the best combination of mechanical properties at tensile strength levels in excess of 300,000 psi. Of the four alloy systems investigated, steels from the low alloy medium carbon Mi-Cz-Mo-Si-V martensific system developed the best combination of fracture toughness, fatigue strength and stress corrosion cracking resistance. A martensific alloy was developed with a nominal composition of

č	Mn	P	<u>s</u>	<u>S1</u>	<u>Ni</u>	Cr	Mo	ų.
0,40	0.35	<,010	<.010	2.25	1,3	0.80	0.25	0.22

which schieves the following average longitudinal properties based on lsboratory sized heats: Y.S. = 268 ksi, U.T.S. = 311 ksi, El. = 12%. R.A. = 44%, CVN = 20 ft-1bs, K_{IC} = 60 ksi $\sqrt{10.05}$, K_{ISCC} = 17 $\sqrt{10.05}$ $\sqrt{10.05}$, unnotch fatigue strength at 107 cycles of 170 ksi, and a notch (K_t = 3.0) fatigue strength of 80 ksi.

The stress corrosion studies demonstrated that variations in phosphorous, sulfur, silicon, chromium, and molybdenum, significantly influenced plane strain fracture toughness properties, but had essentially no effect on the KISCC stress corrosion cracking resistance parameter. The low alloy Ni-Cr-Mo-Si-V martensitic steels had greater SCC resistance than the best bsinitic steel.

Processing studies conducted on two bainitic alloys and one marteneztic alloy revealed that the vacuum are remelted (VAR) steels had the highest levels of fracture toughness, and the electroslag remelted (\vec{xsR}) steels had the lowest levels of fracture toughness. Investigating the influence of melting practice on fatigue properties demonstrated that for the two bainitic steels the ESR material had the highest fatigue strengths, and the VIM material the lowest fatigue strengths. For the martensitic steel the VAR material had the highest fatigue strengths followed by the ESR material and the VIM material. The experimental steels demonstrated tension-tension unnotch fatigue strengths, at 10⁷ cycles, in the range of 170,000 to 210,000 psi. Notch (K_t = 3.0) fatigue strengths at 10⁷ cycles of 80,000 psi were achieved. Thermal-mechanical working treatments demonstrated that the strength and toughness properties of ultra-high strength low allop martensitic and bainitic steels are influenced only slightly by refinement of the prior austenite grain size.

Comparison of the mechanical properties of the newly developed, low siley mertensitic steels with similar properties of currently used commercial ultrahigh strength steels nevealed that the strength-toughness and the strength-SCC resistance characteristics of the new low alloy maxtensitic steels are superior to those of the commercially produced steels. Both the unnotch and notch fatigue strengths of the new Ni-Cr-Mo-Si-V martensitic steels were superior to the similarly measured fatigue strengths of the commercial steels.

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I. INTRODUCTION

The increasingly stringent requirements, in terms of load, range, maneuverability, and performance of present and future military and commercial aircraft, places increasing demands upon the strength and reliability of the steels and other high strength materials used in these aircraft. The great need to save weight in these larger and higher performance sircraft, justifiably causes designers to look for higher and higher strength steels. The increased strength, however, must not be accompanied by decreases in fracture toughness, fatigue strength, and stress corrosion cracking resistance in order to ensure reliability in the performance of the critical load bearing components. For it is well known that very small flaws, excessive inclusion contents, or hostile environments can cause catastrophic failures at stresses well below the design level, if the material lacks sufficient fatigue strength, toughness, or stress corrosion resistance.

The objective of this program was to develop an ultra-high strength steel in the 300 to 320 ksi ultimate tensile strength range, with improved fatigue strength, fracture toughness, and stress cerrosion resistance for greater reliability in forged landing gear components and related structural airframe applications. The guaranteed minimum ultimate tensile strengths of the two currently widely used low alloy steels for landing gears are 260 ksi for 4340 steel and 270 ksi for 300 M steel. An increase in the guaranteed minimum ultimate tensile strength to 300 ksi would, therefore, represent a 10 to 15% increase in ultimate tensile strength level. The approach to achieving these goals was to conduct concurrent alloy development and processing studies, The alloy development efforts included three separate approaches as follows: A. Low Alloy Bainites, B. Medium Alloy Bainites, and C. Low Alloy Martensites. The processing studies included the influence of impurities and melting practice on mechanical properties as evidenced by the effects of vacuum induction melting, electroslag remelting, and vacuum are remelting; and the effects of thermal-mechanical working on an annealed ferrite-carbide matrix and its influence on heat treated properties of low alloy steels.

II. MATERIALS AND PROCEDURES

A. Materials

I

The great majority of the experimental alloys were melted in a 50 ib vacuum induction melting (VIM) furnace. The alloys were melted either as single 50 lb ingots or three-way split heats resulting in three 16 lb ingots. Fifteen heats (Heats El through 15) were melted in a 300 lb VIM furnace at Battelle Memorial Institute, Columbus, Ohio, as two-way split heats resulting in 50 lb ingots. Some of the earlier bainitic steels were melted in an air induction furnace and poured into 70 lb ingot molds. A prefix of the letters A, B, or C before a heat number indicate an air induction melted heat, while the letters V or Z indicate a VIM heat. All ingots were forged at 2150 P, conditioned, and rolled to 1/2-inch thick plate at 1950 F.

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Three experimental alloys (C229, C230, C231) were initially air induction melted as 85 1b ingots, forged to 2-5/8-inch diameter rounds, and electroslag remaited (ESR) by Mellon Institute, Pittsburgh, Pennsylvania. The electrodes were remelted in a water cooled copper mole using AC power at a current of 2000 to 1400 amps and a voltage of 39 to 43 volts. The melting rate was 1 to 1.2 lbs/min. An automatic electrode guide mechanism kept the electrode 20 to 30 mm into the flux, using a 3-inch flux gap. The flux used contained 60-70% CaF₂, 10-15% lime, and the balance alumina. One ingot (Heat C229) had a rough surface because the furnace controls kept cutting off and the high lime flux boiled up during remelting.

The vacuum are remelted (VAR) heats were melted at our Central Alloy District Canton, Ohio, as 350 lb electric furnace heats using a standard double slag practice and poured into 9-inch diameter electrode ingot molds. The electrode ingots were then conditioned and consumable vacuum are remelted. The VAR ingots were then forged at 2150 F to 4-inch square bars, by length, cut into 14-inch lengths, conditioned, forged again to 1-1/2-inch thick slabs, conditioned, and rolled to 1/2-inch thick plate.

B. Test Procedures

The following general procedure was used in the processing and heat treatment of all experiments1 slloys. The 1/2-inch thick place material was sectioned into rough cut 1/2-inch square tensile and Charpy coupons, gradient bars, and quenching dilatometer samples. The gradient bars were heated in a gradient furnace, with a 2200 F backwall temperature, and water quenched in order to determine the $A_{1,2}$ A_3 and carbide solution temperatures for subsequent heat treatment of the alloy. This ensured that each alloy was heat treated correctly, that is the sustanitizing temperature was selected as the minimum temperature which ensured that all carbides were in solution (in one hour) in the sustenite. All mechanical property specimens were initially normalized for 1 hour st a temperature of 100 to 150 F above the austenitizing temperature for the alloy. All heat treatments were performed in neutral sait baths and the martensitic alloys were quenched into 120 F agitated oil, and couble tempered at various temperatures for 2 + 2 hours. When refrigeration treatments were used the material was refrigerated before tempering at -110 F for 1 hour. The quenching dilatometer apeciments were used to determine the martensite start temperatures and if the alloy was heat treated baialtically, the TTT curve was determined metallographically. The mechanical property specimen coupons were then heat treated and subsequently finish ground. The Charpy V-notch specimens were of standard dimensions (0,394-imples square and 2.165-inches long). The tensile specimens were standard 0.252-inch diameter round specimens with a 1-inch gage length as shown in Figure 1(a). All tensile tests were performed at room temperature.

The plane-strain fracture toughness cests were performed at room tempersture, in ordinary air, and at -65 F in a bath of dry ice and trichloreethyleas using the notch bend specimen shown in Figure 1(b). The preparation of the test specimens consisted of cutting a 1/16-insh slot 0.230-inches deep by a grinding wheel and extending its base approximately 0.000-inches deep by electric discharge machining, using 0.003-inch thick brass shim stock which produced about a 0.004-inch wide slot. The length of the initial notch is denoted as, a_0 , in Figure 1(b). The notch was further extended about

0.120-inches by fatigue precracking the specimen in a three point bending configuration and icom losd measurements the maximum stress intensity (K_f max) at the crack tip during final stages of fatigue crack growth was calculated to be typically about 0.4% of K_Q for most alloys and never exceeding 60% of K_Q. The final crack length, s, was within the limits of 0.45 to 0.55 W where W is the specimen depth, nominally 0.800-inches. The plane-strain fracture toughness tests at both room temperature and -65 F were in accord with and met all of the current ASTM (E399-70T) recommendations and requirements for plane-strain fracture toughness testing (1).

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The stress corrosion tests utilized the same plane-strain fracture toughness specimen, fatigue precreacked in the same manner. After precracking all specimens were stored in desiccators prior to testing. All specimens were cantilever loaded "wet" (salt solution added prior to the application of load) as shown in Figure 2. The initial stress-intensity levels (K_{Ii}) were calculated according to the relationship given by Kies, et al. (2). The 3-1/27 NaCl solution was changed daily except on weekends and specimens were exposed until either the specimen failed or a minimum time of 200 hours had elapsed.

The Charpy V-motch specimens were tested at various temperatures on a 240 ft-1b capacity SATEC Impact Tester Model S1-1C, which was certified by the Army Materials and Mechanics Research Center, Watertown, Massachusetts.

Smooth and notched ($K_t = 3.0$) fatigue specimens were tested in tension-tension loading at an R value of +0.1 (R is the ratio of minimum stress to maximum stress). The unnotched fatigue specimens, shown in Figure 3(a), were lapped to produce a surface finish of 2 to 6 rms. The tests were performed on a Sonntag Hodel SF-4 fatigue testing machine cycling at 3600 cpm. The fatigue specimen and grips were surrounded with a Tygon tubing container, as shown in Figure 4, through which a flow of prepurified nitrogen gas (dewpoint of -S0 F) is passed for the duration of the test in order to maintain a constant, low humidity, testing atmosphere. The alignment of the machine, loading train, and test specimen were checked thoroughly both statically and dynamically in order to minimize bending stresses.

III, RESULTS AND DISCUSSION

The results of the alloy development studies will be presented initially, and include extensive investigations on low alloy bainitic steels, medium alloy bainitic steels, and low alloy marteneitic steels. The results of the stress corrosion investigations will be presented next, followed by the results of the processing studies. The processing studies include an extensive investigation of the effects of impurities and melting practice on tension-tension fatigue properties and the effects of thermal-mechanical working treatments on strength and toughness properties.

A. Alloy Development Studies

1. Low Alloy Bainitic Steele

The two decomposition products of austenite, in alloy carbon steels, which can be utilized to achieve ultra-high strength levels are lower bainite and martensite. It is now well established that for the same composition and equivalent strength levels lower bainite has toughness properties which are superior to tempered martensite (3,4,5). It was felt, therefore, that if a beinitic steel could be developed with strengths in excess of 300,000 psi ultimate tensile strength it should have toughness properties superior to those of a martensitic steel at the same strength level. One approach used to achieve this aim was to suitably alloy a steel to produce an aging response or secondary hardening upon aging subsequent to the isothermal bainitic treatment. This concept and these steels are hereafter referred to as baimaging steels. The basic alloy system chosen was a medium carbon Ni-Cr-Mo-V system. The compositions of the Ni-Cr-Mo-V bainaging alloys are presented in Table I, and the tensile and Charpy V-notch impact properties are presented in Table II. As mentioned earlier in the section on Test Procedures, for each bainitic alloy the A_1 , A_3 , carbide solution and M_s temperatures were determined along with the complete lover bainite portion of the TTT curve for each alloy. Therefore, the mechanical properties reported are for the optimum heat treatment for each alloy. For these low alloy bainitic steels the austenitizing temperatures ranged from 1550 F to 1750 F and the transformation times ranged from 6 hours to 48 hours to ensure that a completely bainitic structure was formed. Each alloy was isothermally transformed at a temperature of 25 F to 35 F above its Mg temperature in order to obtain maximum strength properties. The age hardening curves for these alloys indicated that if age hardoning did occur, maximum hardening was observed at a temperature of approximately 950 F for an aging time of 4 hours.

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The mechanical properties in Table II reveal that if the alloy overages or softens upon bainaging at 950 F the toughness generally increases, however, if the strength level remains constant or increases upon bainaging the toughness generally decreases. Thus the strength-toughness response to secondary hardening in these bainitic steels is very similat to the behavior observed in secondary hardening martensitic steels. The data in Tables I and II also reveal that for a given strength level an increase in Ni is beneficial to toughness and that increasing Cr and B are detrimental to toughness. Three of the alloys (B332, B333, and B355) had ultimate tensile strengths in excess of 300,000 psi in the as-transformed condition, while the highest strength obtained in the aged condition was 294,000 psi for alloy B346. These results indicate that in order to obtain an ultimate tensile strength of 300 ksi in these bainitic steels a minimum carbon content of 0.50% will be required. While a strength level of 301 ksi war obtained for alloy B355 with 0.45% carbon and 2.00% silicon this is not a desirable approach because of the very detrimental influence of the high silicon content on Charpy

impact toughuens. Based on these results six additional compositions were vacuum induction melted, processed, and mechanical properties determined. The compositions of these alloys (2389 through 2394) are presented in Table I, and the mechanical properties are presented in Table III. About 4% cobait was added to all of these alloys in order to move the bainite finish curve to the left or shorter times. All of these six alloys were isothermally transformed for 6 hours.

The date in Table III reveal that all of the alloys meet the strength requirements in the as transformed condition but that significant softening occurs upon aging all alloys. As saticipated the alloy with the highest molybdenum content (2391) demonstrated the highest retentivity of strength upon aging. As was shown by the data in Table II the elements which produce age hardening (chromium and molybdenum) cannot be raised to too high a level because a severe loss in toughness accompanies a pronounced age hardening response. The strength and toughness response of these bainitic steels us a function of chromium and molybdenum contents are shown in Figures 5 and 6. Increasing chromium content is seen to have a slight strengthening effect for sll three heat treatment conditions and an inconsistent and unpronounced effect on toughness properties. Increasing molybdenum content increases strength and decreases Charpy impact toughness as shown in Figure 6. The lower strength of the sz transformed bainite at the highest molybdenum level is not a real compositional effect but is probably attributed to a higher isothermal transformation temperature for alloy 2391. Alloys 2389, 2390, and 2391 were isothermally transformed to bainite for 6 hours at 495 F, 475 F. and 525 F respectively.

The strength-toughness properties of these as transformed bainitic steels are shown in Figure 7. The mechanical properties shown in Figure 7 and similar data on the other alloy systems illustrated throughout this reputione plotted in this manner in order to readily compare the strength-tor mess relationships of the experimental alloys with those of the currently, widely used 300 M start. As 300 M steel is the most widely used commercial alloy for forged landing gear components it seems appropriate to compare the strengthtoughness properties of the experimental alloys with the strengthtoughness properties of similarly processed 50 ib VIM laboratory produced 300 M steel. When 300 M steel is vacuum induction melted and processed in the same manner as the experimental alloys, it typically achieves an ultimate tensile strength of 286 ksi with Charpy V-notch impact energy values of 19 ft-lbs at +70 F and 16 ft-lbs at -65 F when tested in the longitudinal direction. Therefore, the coordinates drawn in Figure 7 and in other figures throughout the report are drawn at a value of 286 kai U.T.S. and 19 ft-lbs Charpy V-notch impact energy for ready comparison with the strength-toughness properties of the experimental siloys. From the data in Figure 7 it can be seen that while the desired strength levels were achieved, the Charpy impact energy values did not reach the desired levels. As the strength-toughness relationships of these low alloy bainitic steels did not look encouraging work in this alloy system was terminated.

2. Medium Allcy Bainitic Steels

At the 260 to 280 ksi ultimate tensile strength level HP 9-4-45 steel heat treated bainitically has the best combination of strength and toughness of any available commercial alloy. It's only shortcoming is that it is limited to a minimum tensile strength of about 260 ksi. Therefore, efforts were initiated to modify HP 9-4-45 steel in order to increase the strength level range of the bainitic microstructure. Initially twelve 70 lb air induction heats were made to determine if the strength level could be increased readily. After determining the TTT diagrams of the twelve alloys, four of the alloys had unsuitable TTT curves for bainitic heat treatment, therefore only eight of the alloys were heat treated for mechanical properties. The compositions of these alloys are presented in Table I and the properties are listed in Table IV. Of these eight alloys only alloy B297 showed promise toward achieving higher strength levels. It's toughness level is also very high considering that these were air melted ingots. Based on these results eleven additional medium alloy bainitic steels, with varying levels of carbon, nickel, chromium, and molybdenum were vacuum induction melted, processed and tested. The compositions of these steels (Hests Z211 through 2225) are presented in Table I and the mechanical properties are presented in Table V. Alloys 2211 through 2214 are high carbon modifications of the previous most promising medium alloy bainitic steel, (Heat B297). The dats in Table V reveal that increasing the carbon content has lowered the Mg temperature sufficiently to enable the desired strength levels to be achieved in a bsinitic microstructure. Each of these four alloys were isothermally transformed at both 25 and 75 F above their respective M_s temperatures. The two highest carbon heats, Z213 and Z214 achieved tensile strengths greater than 330,000 psi. The increase in strength level was, however, not without cost as the impact properties were decreased considerably. The strength-toughness relationships of these alloys are illustrated in Figure 8.

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Seven heats of the medium alloy bainitic steels, Z219 through Z225, were investigated to determine the effects of varying Ni, Co, Cr, Mo, and Si levels on strength and toughness properties. Heats Z219, Z220 and Z221 (see Tables I and V) are a 0.46% C, 3.5% Ni, 2.0% Co composition with varying silicon content. As shown in Figure 9 it can be seen that increasing Si produced a modest increase in strength with a commensurate decrease in Charpy impact toughness. These alloys, however, are short of the desired strength level range. Alloys Z222, Z223, and Z224 are a modified Krupp steel with higher carbon contents for strength, and cobalt added to decrease the bainite finish times. The strength level of these bainitic steels are also marginal. The tensile strength-Charpy impact toughness relationships for these heats are also shown in Figure 8. As the strength-toughness relationships of these medium alloy bainitic steels did not look promising, work in this alloy system was stopped.

3. Low Alloy Martensitic Steels

a. Medium Carbon Ni-Cr-Mo-W-V System

The low alloy martensitic steel approaches involved two alloy systems, and the first alloy system investigated was the medium carbon Ni-Cr-Mo-W-V system. As evidence existed indicating that tungsten enhances the toughness of medium cerbon martensitic steels (6) as well as the toughness of tool steels (7) it was decided to thoroughly explore the influence of tungsten on toughness in low alloy ultra-high strength steels. The composition and properties of the initial heats in the Ni-Cr-Mo-W-V system (Heats V707 through 279) are presented in Tables I and VI respectively. Alloys 275, 277, 278, and 279 are remake heats of V710, V748, V750 and V751 respectively, because of missed compositions. These alloys were designed statistically using a 1/3 replicate of a 3³ factorial experiment, confounding using the V(ABC) aliases. The mechanical property data in Table VI were analyzed statistically and the results of regression analyses for the 400 F temper data are shown below:

U.T.S., ksi = +65.5 + 468.5 (%C) + 13.1 (%Cr) + 5.9 (%Ho) + 6.9 (%W)

 $R^2 = 0.85$ Standard Error = 6.8 ksi

CVN, ft-lbs = + 41.i - 8.6 (%Cr) - 22.3 (%Mo) - 8.7 (%W) + 2.9 (%Cr²) + 7.0 (%Mo²) + 2.0 (%W²) R² = 0.74 Standard Error = 3.5 ft-lbs

 R^2 is the multiple correlation coefficient which measures the fraction of total variation about the average of the dependent variable, which has been explained by the regression equation. \mathbb{R}^2 can range between 0 and 1, values close to 1 meaning that most of the variation in the dependent variable has been explained by the regression equation. Values near 0 indicate that the regression equation has explained little of the variation in the dependent variable. The standard error or standard error of estimate is a measure of the goodness of fit of the regression equation. The smaller the standard error of estimate the better the regression line fits the data. The strength-toughness relationships of these alloys are illustrated in Figure 10. Based on these results a second set of fifteen heats were statistically designed and processed. The design of these alloys (295 through 2111) employed a 2³ factorial experiment, augmented by stor and center points. The compositions of these heats are presented in Table I and the properties are given in Table VII. It can be seen from Table I that the composition range of Cr, Mo and W was broadened considerably in heats 295 through 2111, compared to the earlier V707 series of heats. This was done with the intention of achieving ultimate tensile strengths in excess

of 290 ksi at tempering temperatures in the neighborhood of 1000 F. This is a very difficult goal to achieve and still maintain adequate toughness. The idea was to add sufficient alloy carbide formers to produce a moderate degree of secondary hardening to achieve the desired strength levels at high tempering temperatures. A pronounced secondary hardening response was to be avoided, because of its well known embrittling effect. It can be seen from Table VII that alloy 2106 did achieve an U.T.S. of 270 ksi when tempered at 1000 F, however the toughness level was low.

The strength-toughness relationships for these alloys are shown in Figure 11, and the results of the regression analyses for the 400 F temper condition are given below:

U.T.S., ksi = + 311.3 + 4.5 (%Cr) - 6.6 (%Mo) - 1.0 (%W) + 2.3 (%Mo²) - 2.5 (%W x %Mo)

 $R^2 = 0.94$

Standaró Error = 1.8 ksi

CVN, ft-lbs = + 29.2 - 6.05 (ZCr) - 5.9 (ZMo) - 9.09 (ZW) + 2.7 (ZCr x ZW) + 1.50 (ZW x ZHo)

 $r^2 = 0.51$

Standard Error = 1.8 ft-1bs

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In addition to these statistically designed Ni-Cr-Mo-W-V alloys a series of classically designed alloys was VIN and processed in order to determine the optimum levels of manyanese, silicon and vanadium in this alloy system. The compositions of these alloys (heats 1112 through Z120) are given in Table I and the mechanical properties are listed in Table VII. The data on the manganese series of heats (2112, 2113, 2114) reveal that manganese does not have a significant effect on either strength or toughness. The data for the vanadium series (bests 2118, 219, and 2120) indicates that increasing vanadium from 0.10% to 0.19% has a beneficial effect on Charpy impact toughness with no further increase in toughness when vanadium is increased to 0.31%. The effect of silicon, as shown by heats 2115, 2136 and 2117, is to markedly increase both yield and ultimate tensile strength, with a sacrifice in toughness at the highest silicon level. Based on these results three additional compositions were melted, processed, and mechanical properties determined. The compositions of these alloys are listed on Table I (2386, 2387, 2388) and the mechanical properties are listed in Table VIII. The strength and toughness properties for the SON P tempering temperature are shown as a function of sugging content in Figure 12. The data reveal that tongsten has a negligible "frect on both strength and toughness properties, sad therefore it appears that the addition of the element tungsten to these low alloy artensitic steels is not beneficial. Shile this combination of strength and toughness (312 ksi and 17 ft-16s) does not need to be apologized far, it is not as prodising as some of the alloys in the Ni-Cr-No-Si-V. low alloy mortantic system; therefore work was stopped at this point on the Ni-Cr-Mo-W-V alloy system.

b. Medium Carbon Ni-Cr-Mo-Si-V System

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The so and approach for the low alloy martensitic steels was the medium carbon Ni-Cr-Mo-Si-V alloy system. Initially fifteen VIM heats (243 through 260) were processed and evaluated and their compositions and properties are given in Tables I and IX respectively. Unless noted otherwise all of the alloys in this alloy system were refrigerated for 1 hour at -100 F prior to double tempering for 2 + 2 hours at the indicated tempering temperature. The effect of silicon content on strength and toughness properties is shown in Figures 13 and 14. In both series of alloys increasing silicon markedly increased both yield and ultimate tensile strength. In the Z43 alloy series increasing silicon also increased the room temperature Charpy impact toughness slightly, while in the Z58 alloy series increasing silicon decreased toughness slightly. The effect of silicon content on the Charpy impact energy-transition temperature curves for these six alloys is illustrated in Figures 15, 16, 17 and 18. The date in these figures illustrate that generally as silicon is increased and hence strength level is increased that the Charpy impact energy is decreased for a given test temperature. The data in Figure 16 however demonstrate the opposite effect and it is seen that as silicon content is increased toughness increases at all test temperatures. This is an important observation and indicates that further exploration of the effect of silicon content on toughness is necessary and also its possible interaction with chromium and molybdenum as the only difference in the 243 series and the 258 series of heats is the chromium and molybdenum levels. The data presented in Figures 17 and 18 slso reveal the importance of tempering temperature in evaluating toughness properties when silicon content is a variable. These data reveal that at the 600 F tempering temperature the low silicon (0.1.1) Si) alloy does not have superior toughness properties even though the alloy is much less strong than the two higher silicon alloys. This is because at a tempering temperature of 600 F the low silicon steel (258) is right in the middle of the tempered martensite embrittlement trough as illustrated in Figure 19. Because of the well known effect of silicon in retarding the kinetics of cementite precipitation the tempered martensite embrittlement range has been shifted to the right about 200 F to 700 F and 800 F for the higher silicon levels compared to the usual 500 F and 600 F range for the lower silicon levels The influence of increasing silicon upon retarding the degredation of strength properties with increasing tempering temperature is also illustrated in Figure 19. Similar observations are made for the Z43 series of alloys in Figure 20.

The effect of chromium on strength and toughness properties is illustrated in Figure 21 for both the 246 and 249 three-way-split series of heats. The Charpy impact-transition temperature curves for these six alloys are shown in Figures 22 and 23. For the 249

series zero chromium appeared to be the best level, while for the Z46 series about 0.9% chromium appeared to give the best combinstion of properties. The influence of cobalt and vanadium on toughness properties is illustrated in Figures 24 and 25. The data in Figure 24 reveal that 2% cobalt is detrimental to toughness properties and has no effect on strength properties in these low alloy martensitic steels. Increasing vanadium content from 0.10% to 0.20% is seen to be beneficial to toughness properties as shown in Figure 25. The strength-toughness relationships for alloys Z43 through Z60 are compared to similar properties for laboratory produced 300 M steel in Figures 26 and 27. It is noted from these figures that for test temperatures of both +70 F and -65 F that many of these alloys look promising from a strength-toughness viewpoint. The mechanical properties on this first series of heats, in the Ni-Cr-Mo-Si-V alloy system, have demonstrated, therefore, that increasing vanadium content from C.10% to 0.20% is beneficial to impact toughness, that 1% cobalt is detrimental to impact toughness, and that the optimum levels of Si, Cr, and Mo warrant further detailed investigation.

In order to determine the optimum levels of Si, Cr, and Mo in this alloy system, as well as any possible interaction effects between these elements, a set of fifteen statistically designed heats were vacuum induction melted and processed. In addition to these statistically designed alloys a series of classical design single compositional variable alloys were melted and processed in order to determine the optimum levels of Ni, Mn, and V in these Ni-Cr-Mo-Si-V low alloy martensitic steels. The results of the single variable experiments will be discussed first.

OPTIMUM LEVELS OF NICKEL, MANGANESE, AND VANADIUM

In addition to determining the optimum levels of Ni, Mn, and V in these low alloy martensitic steels, the effect of columbium ou strength and toughness was also investigated. The use of Cb as a strengthener and grain refiner in hot rolled, high strength low alloy (HSLA) steels is well known, however its effect on structure and properties in medium carbon martensitic steels has not previously been reported. The element vanadium is usually used for this purpose in quenched and tempered ultra-high strength steels. The compositions and mechanical properties of the (b series of heats (2332-2334) are listed in Tables I and I respectively. These data illustrated in Figure 28 reveal that strength properties are independent of columbium content, but that Charpy impact toughness increases with Cb content. The effect of Cb is therefore, similar to the effect of vanadium in enhancing toughness, however, as will be shown later vanadium has a more pronounced effect than columbium.

The compositions of the nickel series of heats are listed in Table I and the machanical properties are listed in Table X and illustrated in Figure 29. The influence of nickel was investigated in two sets of three-way-split heats (Heats 2273, 2274,

2275 and 2329, 2330, and 2331), and therefore the data points from the two separate sets of heats are not connected in Figure 29. The strength is generally higher and the toughness generally lower in the higher nickel series of heats because these heats had about a 0.40% carbon content compared to a 0.38% carbon content for the lower nickel level series of heats. The effect of nickel level on strength properties is not straight forward, but should probably be interpreted as having essentially no effect on strength properties. In the 2 to 3% Ni range the boughness properties seem to be independent of nickel level. In the lower nickel range the Charpy impact energies significantly increase when nickel increases from 0.52 to 1.66%. In addition the data in Table X reveal that the reduction in area values increase considerably (from 20% to 40%) over this same range of nickel contents. These data indicate that for 0.40% carbon martensitic steels there is no improvement in toughness when Ni is increased from 2 to 3% and that the optimum nickel content should probably be in the range from 1.5 to 2.0%.

The mechanical properties for the vanadium series of heats (2276, 2277, and 2278) shown in Figure 30 demonstrate that as the vanadium content is increased from 0.10% to 0.20% there is a slight increase in yield strength, no change in ultimate tensile strength, but a substantial increase in notch impact toughness. When the vonsdium level increases from 0.20 to 0.29% there is essentially no change in strength, but a further slight increase in toughness. Specimena from these three hests were examined metallographically in order to explain the improvement in toughness with increasing vanadium content. Figure 31 shows the microstructures of the three vanadium alloys at a magnification of 100%. Measurements of the prior sustenite grain size revealed that Heat Z276 with 0.10% V had an ASTM grain size of 5.4, and Heats 2277 (0.20% V) and 2278 (0.29% V) had ASTM grain sizes of 8.4 and 8.7 respectively. The information in Figure 30 also indicates that the percent reduction in area (% RA) increases in a parallel manner to toughness with increasing vonadium coutent. It is well known that % RA is grain size dependent and therefore, it is clear that the refinement of grain size by the addition of varadium up to 0.20% is responsible for the large improvement in toughness and RA but that a further increase in vanadium content causes only slight further grain refinement and thus only a slight further improvement in toughness and reduction in area. These three alloys were also examined by transmission electron microscopy in order to determine if other factors besides grain refinement were responsible for the improvement in mechanical properties. This investigation indicated that all three alloys had similar structures of tempered martensite and plates of Fe₃C, and no significant differences were found; therefore, it is believed that the major effect of vanadium on the improvement of mechanical properties is due to grain refinement.

The results of the manganese series of heats (2270, 2271, and 2272) presented in Table X and Figure 32 demonstrate that as the Mn content is increased, the Y.S. and U.T.S. remain essentially constant, but the toughness decreases substantially. As these

low alloy martensitic steels are not lacking in hardenability, and the finding that increasing manganese is detrimental to toughness it would appear prudent to add only sufficient manganese to the up about 0.010% maximum sulfur; about 0.30% Mn would therefore appear to be about optimum. In order to understand the reason for decreasing Charpy impact energy with increasing manganese, these alloys were examined by transmission electron microscopy. The results of this examination indicate that the precipitation of carbides during tempering was similar in all three alloys, the carbide in this case (600 F temper) being essentially FeaC. The only difference that could be discerned was that alloy Z272 with the highest Mn level had a greater number of microtwinned martensite plates than did the lower Mn alloy Z270. Figures 33 and 34 show typical regions of alloys 2270 and 2272 respectively. Although, Heat 2272 was not heavily microtwinned it did contain substantially more twins than Heat Z270.

It has been well established that microtwinned martensite has inherently less toughness than dislocated martensite (8,9) and that the lowering of the M_s temperature increases the tendency towards twinned martensite (10). Manganese lowers the M_s temperature in steels and therefore by increasing the Mn content there is a greater tendency towards the production of lower toughness, twinned martensite. It is a little surprising however, that an increase of approximately 0.50% Mn would be enough to produce a substantial difference in the toughness of the martensite if twinning were the only explanation. Therefore, it is felt that twinning does play a role in lowering the toughness of the higher manganese alloy but in addition, some subtle effect of Mn may be operative which as yet cannot be detected. Structure-property correlations in medium carbon martensitic steels are at best elusive.

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OPTIMUM LEVELS OF SILICON, CHROMIUM, AND MOLYBDENUM

In order to determine the proper levels of Si, Cr, and Mo, and possible interaction effects between these elements, a set of fifteen statistically designed heats were vacuum induction melted as two-way split heats resulting in 50 lb ingots. The composition of these heats (Heats R1 through 15) are listed in Table I and the mechanical properties are shown in Table XI. The prefix R before heats 1 through 8 stands for remake as the compositions were missed the first time on the first eight heats and hence were remade. These 50 lb heats provided sufficient material to determine fracture toughness and stress corrosion properties in addition to tensile and Charpy impact properties. In order to determine mechanical property-composition prediction equations and possible interaction effect; between the elements Si, Cr, and Mo, Heats R1 through 15 were designed statistically utilizing a 23 factorial experiment, sugmented by star and center points. The prediction equations for ultimate tensile strength, plane strain fracture toughness at both room temperature and -65 F, and

Charpy impact energy resulting from the regression analyses for the composition and mechanical properties shown in Tables I and XI are presented below:

U.T.S., ksi = 322.5 - 90.2 (% Si) +76.6 (% Cr) +24.6 (% Si²) -23.1 (% Cr²) +12.8 (% Mo)

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 $R^2 = 0.92$ Standard Error = 2.9 ksi

 $\begin{array}{rl} {\rm K_{IC},\ ksi}\,\sqrt{\rm in.} &=& -35.7\ +119.9\ (\%\ Si)\ -39.1\ (\%\ Cr)\ +31.5\ (\%\ Mo)\\ {\rm at}\ +70\ F & -29.6\ (\%\ Si^2)\ -19.8\ (\%\ Mo^2)\ -19.8\ (\%\ Cr\ x\ \%\ Mo)\\ & +16.1\ (\%\ Cr^2) \end{array}$

 $R^2 = 0.97$ Standard Error = 1.5 ksi \sqrt{in} .

 K_{IC} , ksi \sqrt{in} . = 20.46 +38.0 (7 Si) -36.7 (7 Cr) -11.9 (7 Mo) at -65 F -14.8 (7 Si²) +10.7 (7 Cr²)

 $R^2 = 0.94$ Standard Error = 1.4 ksi \sqrt{in} .

CVN, ft-lbs = -4.5 + 29.4 (7 S1) -7.0 (7 S1²) -5.5 (7 Ho) at +70 F -3.3 (7 Cr)

 $R^2 = 0.68$ Standard Error = 1.3 ft-lbs

From the values of the multiple correlation coefficient (R^2) and the values of the standard error of estimate, it can be seen that both the U.T.S. and KIC equations should have a high segree of predictability. The multiple correlation coefficients measure the fraction of total variation about the average of the dependent variable which has been explained by the regression equation (values close to 1 meaning that most of the variation in the dependent variable has been explained by the regression equation). The standard error of estimate is a measure of the goodness of fit of the regression equation. However, for the GVN equation, only 68% of the variation in average measured toughness is explained by the variations in Si, Cr, and Mo. This observation is not too surprising as it is well known that fracture initiation tests such as the Charpy V-notch impact test, the reduction in area, and fatigue properties are far more sensitive to inclusion content than are crack propagation tests such as fracture toughness. Hence, the CVN values will be reflecting the content and distribution of other microstructural festures in addition to the intended compositional variables Si, Cr, and Ho far more than the fracture toughness values. At these strength levels, it is clear that a crack propagation test such as ETC is a better toughness parameter for screening compositional variables than is the Charpy V-notch impact test. Galy the +70 F Ric - composition equation indicates an interaction effect between elements as shown by the negative % Cr x % Mo term. It is noted that the -65 F K_{IC} - composition equation differs from the +70 F K_{IC} - composition equation in that it does not contain either a Mo^2 or a Cr x Mo term. It is not unexpected that a toughness-composition response curve is different both qualitatively and quantitatively at different test temperatures.

The mechanics, property-composition prediction equations for U.T.S. and $E_{\rm YL}$ are expressed graphically in Figures 35, 36, 37. The dependence of ultimate tensile strength upon silicon content illustrated in Figure 3. receals that strength is essentially constant matil the Si level is approximately 2.1% at which point the ultimate tensile strength (U.T.S.) increases rapidly. At the same sittion level both $X_{\rm IC}$ expressions reach a maximum and the fracture soughnees decreases above about 2.1% Si as strength inclease 1 From these dats it would appear that the optimum silicor evel is about 2.1%. The influence of chromium level upon K and U.T.S. illustrated in Figure 36 discloses that U.T.S. increases sharply with increasing Cr, until about 1.7% Cr when a maximum is reached and U.T.S. begins to decrease with a further increase in chromium. The +70 F ${\tt K}_{\rm IC}$ equation indicates that ${\tt K}_{\rm IC}$ decreases until about 1.5% Cr at which point K_{IC} reaches a minimum and with a further increase in Cr, KIC increases again. The -65 F KIC equation indicates that KIC decreases until about 1.7% Cr and then a slight upturn in K_{TC} is observed with increasing Cr. It is believed that the -65 F K_{TC} versus chromium response curve is the more valid one and although fracture toughness does increase again at high Cr levels it probably never reaches the level observed at the lowest Cr contents (about 0.75%). From a fracture toughness standpoint, therefore, it would seem that a lover Cr level (about 0.75%) would be most desirable. The mechanical property versus molybdenum expressions shown in Figure 37 disclose that as Mo is increased over the range from 0.25% to 1.0% strength increases linearly and fracture toughness decreases. It would appear advisable therefore, to keep the Mo level low and only add sufficient Mo (about 0.25%) to meet hardenability requirements. The -65 F KIC - composition prediction equation is illustrated in three dimensions in Figure 38. This figure discloses that to achieve maximum fracture toughness silicon should be maintained at about 2.1%, and that molybdenum and chromium should both be limited to the lovest levels investigated which is about 0.25% and 0.75% respectively. It should be noted that the use of prediction equations to extrapolate beyond the compositional limits of the alloy design is an unwise practice.

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2. a further attempt to understand the influence of, and proper level eS Cr and Mo from a fracture toughness viewpoint several correlations of fracture toughness with Cr and Mo parameters are presented in Figures 39, 40, and 41. Data for these figures were taken from heats which had a silicon level in the range from 1.78% to 2.15% which produced an ultimate tensile strength range of 297 ksi to 315 ksi. The data in Figure 39 reveal that the ratio of the carbide forming elements (Cr + Mo) to carbon should be kept as low as possible for maximum fracture toughness. As shown in Table I all of these low alloy martensitic steels had a carbon content of about 0.40%. The variation of fracture toughness with combined chronium and molybdenum content shown in Figure 40 demonstrates again that Cr plus Mo should be held to the lowest possible levels for maximum fracture toughness. The data presented in Figure 41 indicate that the Cr to Mo ratio does not significantly affect fracture toughness at either +70 F or -65 F.

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Based on this analysis of the statistically designed alloys and the results of the single variable experiments to determine the optimum levels of Ni, Mn, and V, two additional heats were vacuum induction melted and processed. As discussed previously the optimum Ni, Mn, and V levels were determined to be about 1.8%, 0.30%, and 0.25% respectively. The results from the statistically designed heats indicated that the optimum silicon and melybdenum levels should be about 2.1% and 0.25% respectively and that chromium should be low, probably at about 0.75%. In order to further clarify the differences in slope of the +70 F KIC, and the -65 F KIC response curve as a function of chromium content (Figure 36) it was decided advisable to melt a high Cr level hest in addition to the lower Cr level heat. The higher Cr level heat would also present an opportunity to validate the predictability of the ultimate tensile strength equation as well. The compositions and mechanical properties of these two heats (2525 and 2551) along with the composition and properties of laboratory produced 300 M steel (Heat 2:49) are presented in Tables I and XII. It can be seen from the data in Table XII that the lower chromium heat Z551 (0.82% Cr) achieved an excellent combination of strength and toughness properties with an ultimate tensile strength of 311 ksi, 19 ft-lbs Charpy impact energy, and a K_{TC} of 60 ksi/in. at +70 F and 45 ksi/in. at -65 F. As expected the higher chromium heat Z525 (1.90% Cr) produced a higher strength level (318 ksi) and a lower toughness level (14 ft-1bs, and 49 ksi \sqrt{in} .). These experimental results are compared to the predicted results in Table XIII, utilizing the mechanical propertycomposition prediction equations presented earlier. This comparison reveals that the prediction equations should not be used to provide accurate quantitative estimates of mechanical properties, but that they are of value in indicating mechanical property-compositional trends qualitatively.

The Charpy impact properties of Reats 2525, 2551, and R1 through 15 are compared to the impact energies of laboratory produced 300 M steel (Heat Z449) in Figures 42 and 43. Considering the generally observed relationship of rapidly decreasing toughness with increasing strength at these ultra-high strength levels, the strength-toughness relationships of these medium carbon (about 0.40% carbon) Ni-Cr-Mo-Si-V martensitic steels are considered quite attractive. The plane strain fracture toughness properties of these alloys are compared to those of 300 M steel in Figures 44 and 45. These comparisons reveal that at both room temperature and -65 F the ultimate tensile strengthfracture toughness relationships for the new experimental martensitic steels are considerably improved compared to 300 M steel. Considering the fracture toughness properties of the slloy with the optimum composition (2551) it is seen that the tensile strength level has been increased by about 25,000 pai

while maintaining the same level of fractive toughness as in 300 M steel. As shown by the laboratory heat of 300 M steel in Figures 44 and 45 and as demonstrated by all of our experience with the HP 9Ni-4Co steels the fracture toughness properties obtained from billet stock from large diameter vacuum arc remelted ingots are almost always greater than those obtained on 4-1/2-inch diameter VIM laboratory ingots. It is, therefore, anticipated that the fracture toughness levels of the experimental martensitic rteels will be improved by about 10 to 20% when they are made commercially by electric furnace-vacuum arc remelt practice. A more detailed comparison of the mechanical properties of these new low alloy martensitic steels with similar properties of other commercially available ultra-high strength steels will be made in the final section of this report.

FRACTURE TOUGHNESS - CHARPY IMPACT CORRELATIONS

The availability of a relatively large amount of well documented fracture toughness and Charpy impact data generated in this program provided the opportunity to attempt correlations between fracture toughness and Charpy impact energy values. Even though previous attempts to correlate KIC with Charpy impact energy values have proven unsuccessful (11), the large economic and time saving convenience of the Charpy impact test compared to the plane strain fracture toughness test was sufficient motive for another attempt to determine a statistically valid correlation between K_{IC} and CVN toughness parameters. It is clear from the average CVN and K_{TC} values for Heats R1 through 15, shown in Figure 46, that there is not a simple linear relationship between K_{IC} and CVN. A total of 47 separate K_{IC} , CVN, and Y.S. observations were programed in a variety of forms in an attempt to determine a valid correlation between K_{TC} and CVN parameters. The most valid correlation found is given below as equation (a):

Eqn. (s)

$$\left(\frac{K_{IC}}{YS}\right)^{2} = -0.18 + \frac{46.3}{YS} + \frac{0.75 \text{ eVN}}{YS}$$

$$R^{2} = 0.91 \qquad \text{Standard Error} = 0.0049$$
or
$$K_{IC} = YS \left[-0.18 + \frac{\sqrt{6.3}}{YS} + \frac{0.75 \text{ eVN}}{YS}\right]^{1/2}$$

$$R^{2} = 0.66 \qquad \text{Standard Error} = 0.487 \text{ km} \text{ fm}^{1/2}$$

A second correlation not as statistically valid, but one that can be plotted in two dimensions is presented below as equation (b):

Eqn. (b)

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$$\begin{pmatrix} \frac{K_{IC}}{YS} \end{pmatrix}^{2} = -0.045 + 1.37 \ (\frac{CVN}{YS})$$

$$R^{2} = 0.82 \qquad \text{Standard Error} = 0.0058$$
or
$$K_{IC} = YS \left[-0.045 + 1.37 \ (\frac{CVN}{YS}) \right]^{1/2}$$

$$R^{2} = 0.66 \qquad \text{Standard Error} = 4.87 \text{ ksi } \sqrt{\text{in.}}$$

This equation (b) is expressed graphically in Figure 47, and the large amound scatter observed would indicate that this is not a reliable relationship to use. One of the problems in determining a statistically valid correlation between \mathtt{X}_{IC} and CVN is that it is not possible to obtain a good independent estimate of CVN and YS because YS and LVN are highly correlated. The expression given in equation (s) is significantly better than equation (b) as can be seen by the values of \mathbb{R}^2 and the standard error of estimate. Even equation (a), however, will not be a good predictive equation because there is too much scatter about the line as indicated by the standard error of estimate. In summary equation (a) is certainly the best equation to use to estimate K_{IC} values from CBN values, however, even equation (a) is only of value for a "ball park" estimate of KIC and unfortunately the experimentalist will still have to perform the fracture toughness test in order to determine a meaningful and valid KIC number. Both of these expressions were obtained from data on low alloy martensitic steels which had yield strengths in the range of 234 to 287 ksi, ultimate tensile strengths in the range of 281 to 332 ksi, K_{IC} values in the range of 34 to 70 ksi /in., and CVN values in the range of 11 to 21 ft-1bs.

B. Stress Corrosion Studies

In addition to enhancing the strength-loughness relationships in ultrahigh strength steels one of the aims of this program was to improve the stress corrosion cracking (SCC) resistance of ultra-high strength steels. Toward this aim two approaches were taken; (1) to lower the impurity elements phosphorous and sulfur to the lowest possible levels, and (2) investigate systematically the influence of the elements silicon, chromium and molybdenum on SCC resistance.

INFLUENCE OF PHOSPHOROUS AND SULFUR

The influence of the impurities phosphorcus and sulfur on the toughness of high strength steels has been investigated widely, however, little work has been done concerning the effects of these impurities on the SCC resistance of high strength steels. To investigate this effect a series of six 35 pound heats of 9-4-45 steel were vacuum induction melted (VIM) with varying phosphorous and sulfur levels and subsequently forged and rolled to 1/2" thick plate material. Tensile, Charpy impact, fracture toughness, and stress corrosicn specimens from each heat were subsequently heat treated to both bainitic and martensitic microstructures. The beinitic specimens were normalized, austenitized, and isothermally transformed at 465 F for six hours. The martensitic specimens were normalized, austenitized, oil quenched, refrigerated at -110 F and tempered at 500 F for 2 + 2 hours. The fatigue cracked stress corrosion specimens were cantilever loaded in a 3-1/2% NaCl solution which was changed daily, except on weekends, and the KISCC values were determined after specimens had run out for 500 bours. KISCC stands for the plane strain stress intensity, under SCC conditions, above which cracking is observed. The compositions and properties of the six heats (V723-V746) are shown in Table I and Table XIV. The stress corrosion curves for two of the alloys are shown in Figures 48 and 49 and curves summarizing the influence of phosphorous and sulfur levels on CVN, KIC and KISCC properties are illustrated in Figures 50 through 55. The complete delayed failure curves for all six alloys were presented in the Second Quarterly Progress Report on this contract. The stress corrosion curves in Figures 48 and 49 illustrate the severe degradation in load carrying capacity of ultra-high strength steels when fatigue cracked samples are stressed in an aqueous environment. For all phosphorous and sulfur levels the bainitic microstructure exhibited greater stress corrosion clacking resistance than the martensitic microstructure.

The effect of phosphorous content on $K_{\rm IC}$ and $K_{\rm ISCC}$ for sulfur levels of .009% and .010% is shown in Figure 50. For this sulfur level it is seen that both fracture toughness and SCC resistance are essentially independent of phosphorous level over the range of .004 to .020%. The effect of sulfur content on $K_{\rm IC}$ and $K_{\rm ISCC}$ is illustrated in Figure 51 for phosphorous contents less than .004%. It can be seen that increasing sulfur level significantly decreases fracture toughness of both the bainitic and martensitic microstructures, while there is apparently no effect of sulfur on the $K_{\rm ISCC}$ levels. The effect of phosphorous plus sulfur content a fracture to parameters is shown in Figure 52 and reveals a pronounced detrimental effect on $K_{\rm IC}$ and a slightly detrimental effect

on K_{ISCC}. With respect to toughness it can be seen from Table XIV that for the same total P + S content (0.029% for alloys V726 and V727) that sulf r is the more detrimental of the two elements. The effect of phosphorous, sulfur, and phosphorous plus sulfur on Charpy impact properties is shown in Figures 53, 54, and 55 respectively. It is evident from these figures that sulfur is far more detrimental to toughness than phosphorous. We have seen, therefore, that while increasing phosphorous and sulfur levels are detrimental to toughness properties of both bainitic and martensitic 9-4-45 steel that these impurity elements had essentially no effect on SCC resistance as characterized by $K_{\rm ISCC}$. It should be noted that only the trends of this impurity element study should be considered, and not the absolute magnitude of the toughness numbers, as these properties were obtained on small laboratory heats of 9-4-45 steel and not commercially produced vacuum arc remelted waterial.

INFLUENCE OF SILICON, CHROMIUM, AND MOLYBDENUM

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The influence of silicon, chromium and molybdenum on SCC resistance in low alloy mactensitic steels was investigated by means of the fifteen sustistica. 'y designed heats (R1 through 15) in order to determine the compositional dependence of the stress corrosion resistance parameter The K_{ISCC} values were determined from specimens which had not KISCC. failed after 200 hours in a 3-1/2% NaCl solution which was changed daily except on weekends. The stress corrosion curves for all fifteen alloys were reported in the Sixth Quarterly Progress Report on this contract and three of these stress corrosion curves are shown in Figures 56, 57, and 58. The general shape of the stress intensity-time to failure curves was the same for all fifteen heats. As shown in Figures 56, 57, and 58, and by the KISCC data in Table XI the variations in Si, Cr, and Mo produced significant changes in plane strain fracture toughness, however, the SCC resistance parameter KISCC was essentially uneffected by these compositional variations. The $K_{\rm ISCC}$ values for all fifteen alloys were in the range of 16 to 19 ksi $\sqrt{10}$. and hence the stress corrosion resistance of these 0.40% carbon martensizic steels is independent of the intended variations in Si, Cr, and Mo; 1.78 to 2,75% Si, 0.80 to 1.75% Cr, and 0.26 to 1.02% Mo (compositious listed in Table I). These same variations in Si, Cr, and Mo produced a change in fracture toughness values ranging from 40 to 64 ksi /in. (Table XI), indicating that KISCC and KIC are in no way simply related. The often made generalization that if a material's fracture toughness is increased its stress corrosion resistance will also be increased is shown here not to be either a good or valid generalization.

While the $K_{\rm ISCC}$ threshold stress intensity values were independent of composition it can be seen from the stress intensity - time to failure curves in Figures 56 through 58, from the portion of the curve where the stress intensity decreases rapidly at essentially constant time, that this essentially constant time to failure over a varying stress range shifts significantly from alloy to alloy. A regression analysis was run to determine if there was a significant composition dependence of the time to failure at an applied stress intensity equal to 60% of the alloy's K_{IC} value. The results of this regression analysis are given below: Time to Failure, linutes = 488.8 +60.3 (% Si) -735 (% Cr)at $K_{II} = 0.60 K_{IC}$ $R^2 = 0.73$ Standard Error = 24.0 Minutes

The equation indicates that at this applied stress lovel the time to failure or stress corrosion resistance is independent of molybdenum content, linearly dependent upon silicon content, and quadratically dependent upon chromium content. For certain compositions this equation is expressed graphically in Figures 59 and 40. Neither the significance nor the reason for the complex dependence of time to failure upon Cr content in known. The increased time to failure at this stress level, with increasing silicon content, is somewhat more understandable as Carter (12) has shown that increasing Si content in 4340 type steels decreases the crack growth rate in KISCC tests. In agreement with the current results, he also found that increasing silicon did not influence the K_{TSCC} threshild stress intensity level even though it did decrease the crack growth rate. This indicates that in 0.40 carbon, low alloy martensitic steels, composition can and does influence the rate of subcritical flaw growth under stress corresion cracking conditions even though it does not influence the stress intensity level at which rapid mechanical crack propagation begins. This stress intensity level at which the slowly moving crack reaches a critical length under SCC conditions and rapid mechanical crack propagation begins is known to be higher than the KIC value for the alloy in these types of steels (12.13), therefore, indicating that some type of crack blunting mechanism is operative. In steels at these ultra-high strength levels, the extent of subcritical slow crack growth is relatively short in terms of both time and crack length; therefore, not much reliance can be placed on finding a subcritical flaw during periodic inspections and, hence, the decreased crack growth rate with increasing silicon content is of no practical import. For comparison purposes, the KISCC value of 300 M steel (VIM Heat Z449) was determined to be 13 ksi /in. (Figure 61). The increase in strength level from about 286 ksi for 300 M steel to about 315 ksi for the experimental Ni-Cr-No-Si-V martensitic steels, therefore, has not produced a degradation in stress corresion cracking resistance. A more detailed comparison of the KISCC values of these experimental low alloy martensitic high strength steels with KISCE values of several commercial high strength steels will be made in the final section of this report.

As the fatigue testing portion of this program included the fatigue testing of bainitic steels as well as martensitic steels it was decided to determine the KISCC values of an experimental bainitic steel in both the VIM and VAR conditions. The stress corresion behavior of medium alloy bainitic steel Z411 and the same composition in the VAR condition (Heat 3838800) (compositions listed in Table I) are shown in Figures 62 and 63. The KISCC value of this medium alloy bainitic steel is seen to be independent of melting practice. While the KISCC value of

13 ksi $\sqrt{10}$ in seems quite low for a bainitic microstructure, the carbon content of this steel is relatively high (about 0.50% carbon) and it is felt that if this some composition whe heat treated to a maxtensitic microstructure the KISCC value would be lower than 13 ksi $\sqrt{10}$.

From an alloy development standpoint it was disappointing to find that the basic stress corresion resistance of low alloy martensitic steels was independent of the variations in Si, Ur, and No contents, thereby closing the door on one more possible avenue of improving the stress corrosion resistance of low alloy martensitic steels from a compositional point of view. It was also disappointing to learn that lowering the levels of phosphorous and sulfur down to .003% P and .002% S did not provide any enhancement in KISCC for either bainitic or martensitic ultrahigh strength steels. Paxton has investigated the effect of several impurities at different impuricy levels on the stress corrosion cracking resistance of 300 grady maraging steel and demonstrated that impurity levels had little effect on KISCC values with the total range of KISCC values varying from 7 to 15 ksi \sqrt{in} , (14). This work on the influence of impurities on the stress corrosion resistance of 197. Ni maraging steel has recently been verified on a commercial scale by the evaluation of a high purity 18% Ni (360) (5-ton heat, .003% C, .001% N, .004% S, .002% F) maraging steel forging where the KISCC level was determined to be 7 ksi /in. (15). Thus, from an impurity level standpoint, it appears that the stress corrosion cracking resistance cannot be improved for either the 18% Ni maraging steels or the low slloy martensitic or balaitic steels. From a microctructural point of view, bainitic structures have greater stress corrosion cracking resistance than martensitic structures; however, at strength levels above 300 ksi, the strength-toughness relationships of bainitic steels are no longer attractive as was demonstrated earlier in this report. The influence of grain refinement has been studied for 4340 steel and it was found that KISCC values ranged from 14-16 kci vin. independent of prior sustenite grain rize variations from ASIM 7 to 12 (16). It was observed, however, that the crack growth rates decreased with decreasing prior-sustenite grain size. From a compositional standpoint, the only work to oste on low alloy martensitic steels which has shown that stress corrosion cracking resistance can be improved by elloying is the work of Sandoz which demonstrated that RISCC increases for decreasing C and Mn levels in 4340 type stuels (17). These steels, however, were heat treated to a relatively low strength level (about 170 to 195 ksi yield strength) and it is not clear that the same compositional dependence would be observed at atrength levels near 300 ksi,

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It appears, therefore, that at strength levels in the neighborhood of 300,000 psi, low alloy steels have greater stress corrosion crecking resistance than 18% Ni maraging steels; however, these levels of resistance are not inherently high and appear to be largely independent of compositional, microstructural, and impurity variables. Thus, metallurgical means for improving the inherent stress corrosion crecking resistance of ultra-high strength steels is not immediately apparent and, therefore, the present means of successful utilization of steels at these strength levels such as shot pseuing, cadmium plating, and psinting will continue to be necessary for the foreseesble future.
C. Processing Studies

The processing studies include the investigation of the effects of three different melting practices; electroslag remelting, vacuum induction melting, and vacuum arc remelting on strength and toughness, and tension-tension fatigue properties. In addition the effect of thermal-mechanical treatments on the mechanical properties of both bainitic and martensitic steels was investigated.

1. Influence of Melting Practice on Strength and Toughness Properties

The influence of melting practice on mechanical properties was investigated for a herein bainitic steel, a medium alloy bainitic steel, and a low alloy martensitic steel. The three experimental alloys were selected from the preliminary alloy development results end vacuum induction melted (VIM), electroslag remelted (ESR), and vacuum arc reselted (VAR) to the lowest possible levels of impurities. The compositions of these three alloys do not necessarily represent the optimum composition in each alloy system, as the compositions had to be selected early in the program in order to be processed by the different melting practices. The compositions of the three experimental steels with the three different melting practices are presented in Table I. The low alloy bainitic steel was not vacuum arc remelied. The VIM heats (Z409, Z411, 2412) were melted as 50 1b ingots. The ESR beats (C229, C230, C231) were initially eir induction melted as 85 lb ingots, forged to 2-5/8-inch diameter rounds, and electrosisg remeited by Mellon Institute. The VAR heats (3838800, 3880808, 3888811) were melted initially at our Central Alley District (Canton, Ohio) as 350 lb electric furnace heats using a standard double slag practice and poured into 9-inch dismeter electrode ingot molds. These electrode ingots were then conditioned and consumable vacuum arc remelted.

The mechanical properties of these alloy melted by VIN, ESR, and VAR techniques are compared in Table XV. The VIM and VAR low alloy martensitic alloys have essentially the same strength and toughness properties; however, the ESR material (Heat C229) has high side. strength properties and low side toughness properties due to a high side carbon content of 0.44 percent. Metallographic examination of specimens from Heats Z409, C229, nd 3888808 revealed that there were no significant differences in the degree of microcleanliness of these three steels, and that all three were very clean. The low KTC and percent reduction in eres values for ESR Heat C229 cannot be totally explained by the higher carbon content, and as the alloy appeared to have a high degree of microcleanliness, the law &-c and % R.A. values are most probably due to the localized segregation of non-metallic inclusions. Slectron metallographic examination of extraction replicas from the fracture surfaces of the tensils samples of the ESR C229 alloy revealed that an unusually large aumber of both globular and angular particles were present of the freeture surface. Electron diffraction analysic of these particles tentativaly identified the globular particles as SiO2 and the angular particles as CaAl204. This localized sepregation condition is most prehabily

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due to the problems encountered during the electroslag remelting of this heat as previously described in the section on Materials and Procedures. The three medium alloy bainitic steels (2411, C230, 3898800) have essentially the same strength properties, with the VAR heat having the highest level of fracture toughness and the ESR heat the lowest level of fracture toughness. The two low alloy bainitic heats both have high strength levels for a bainitic microstructure; however, the fracture toughness level of both the VIM and ESR materials is quite low.

2. Influence of Melting Practice on Fatigue Properties

As fatigue strength is one of the most important design parameters for the successful utilization of ultra-high strength steels for landing gears, an extensive fatigue study was conducted. The influence of composition, microstructure, and melting practice on the tension-tension fatigue properties of various experimental steels was investigated. Both notched and unnotched fatigue tests were conducted at an R value (ratio of minimum stress to maximum stress) of +0.10.

UNNOTCH FATIGUE TESTS

The maximum stress versus cycles to failure (S-N) curves for the three VIM steels are shown in Figures 54, 65, and 66 and the comparison S-N curves for the ESR alloys are shown in Figures 67, 68, and 69. With the exception of alloy Z412, the degree of scatter in the fatigue data is typical and not excessively large for specialty ultra-high strength steels. The reason for the unusually high number of thread failures in alloy Z412, Figure 66 is not known. The fatigue behavior of the low alloy martensitic steel produced by vacuum arc realting (Heat 3888811) is shown in Figure 70. The fatigue behavior of the medium alloy bainitic steel produced by vacuum arc remelting (Heat 3888800) is shown in Figure 71.

The influence of melting practice on the fatigue strengths of these three alloys is illustrated in Figures 72, 73, and 74. The fatigue behavior of the low alloy martensitic steels illustrated in Figure 72 demonstrates that the VAP material had the highest fatigue strengths, and that the VIM material had the lowest fatigue strengths. The comparison S-N curves for the medium alloy bainitic steels (Figure 73) reveal that the ESR material had the highest fatigue strengths and that the VIM had the lowest fatigue strengths. The comparison S-N curves for the low alloy bainitic steels again demonstrate that the ESR material had superior fotigue life compared to the VIM material. The low alloy bainitic steel composition was not vacuum arc remelted. As melting practice has been shown to have a pronounced influence on fatigue properties for a given composition and strength level, a quantitative analysis of the inclusion contents was performed. It is well known that if environmental effects are eliminated the two primary factors controlling the fstigue behavior of high Strength steels are strength level and inclusion content. It will be mentioned later that microstructure

due to the problems encountered during the electroslag remelting of this heat as previously described in the section on Materials and Procedures. The three medium alloy bainitic steels (2411, C230, 3898800) have essentially the same strength properties, with the VAR heat having the highest level of fracture toughness and the ESR heat the lowest level of fracture toughness. The two low slloy bainitic heats both have high strength levels for a bainitic microstructure; however, the fracture toughness level of both the VIM and ESR materials is quite low.

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UNNOTCH FATIGUE TESTS

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The maximum stress versus cycles to failure (S-N) curves for the three VIM steels are shown in Figures 54, 65, and 66 and the comparison S-N curves for the ESR alloys are shown in Figures 67, 68, and 69. With the exception of alloy 2412, the degree of scatter in the fatigue data is typical and not excessively large for specialty ultra-high strength steels. The reason for the unusually high number of thread failures in alloy 2412, Figure 66 is not known. The fatigue behavior of the low alloy martensitic steel produced by vacuum arc remelting (Heat 3888811) is shown in Figure 70. The fatigue behavior of the medium alloy bainitic steel produced by vacuum arc remelting (Heat 3888800) is shown in Figure 71.

The influence of melting practice on the fatigue strengths of these three alloys is illustrated in Figures 72, 73, and 74. The fatigue behavior of the low alloy martensitic steels illustrated in Pigure 72 demonstrates that the VAP material had the highest fatigue strengths, and that the VIM material had the lowest fatigue strengths. The comparison S-N curves for the medium alloy bainitic steels (Figure 73) reveal that the ESR material had the highest fatigue strengths and that the VIM had the lowest fatigue strengths. The comparison S-N curves for the low alloy bainitic steels again demonstrate that the ESR material had superior fotigue life compared to the VIM material. The low alloy bainitic steel composition was not vacuum arc remelted. As melting practice has been shown to have a pronounced influence on fatigue properties for a given composition and strength level, a quantitative analysis of the inclusion contents was performed. It is well known that if environmental effects are eliminated the two primary factors controlling the fatigue behavior of high strength steels are strength level and inclusion content. It will be mentioned later that microstructure

also plays a role in affecting fatigue strength, but it is a secondary role compared to inclusion content. It now has been well demonstrated that the size, number, shape, and location of inclusions has a pronounced influence on the fatigue properties of steel (18, 19, 20, 21, 22, 23, 24, 25, 26, 2/, 28). Oxides and silicates in particular have been found to be most deleterious (20, 22, 24-27), while sulfides have generally been found to have little harmful effect on fatigue properties (18, 20, 24, 26, 27). For this study, as composition and strength level were held essentially constant for a given alloy system, it seemed most likely that the fatigue behavior as influenced by melting practice could most likely be explained by variations in inclusion contents, size, shape or distribution. Standard metallographic examination for microcleanliness for all six alloys (shown in Figures 72, 73, and 74) was performed and no apparent differences were observed. All six heats were very clean. Alloys 2411, C230, and 3888800 of the medium alloy bainitic steel series were selected for detailed quantitative analysis. The quantitative analysis was performed on an AMEDA instrument which is an Automatic Microscopic Electronic Data Accumulator manufactured by Fenco Corporation of Irwin, Pennsylvania. The AMEDA was used to determine total volume percentage of inclusions for all inclusion types. The AMEDA also was used to determine the size distribution of sulfide inclusions, but these data were not correlatable to fatigue strength data as sulfides are not responsible for the initiation of fatigue cracks. Unfortunately from an analysis standpoint the oxide inclusions were too small in these vacuum melted and ESR steels for size distribution data to be determined on the AMEDA. Point counting was used in a few selected instances to verify the volume percent numbers being determined on the AMEDA. Point counting was performed at a magnification of 320X, using 125 fields per specimen, with an eye piece containing a grid with 81 intersections,

The volume percent inclusion data obtained on both the AMEDA instrument and by point counting for the VIM, ESR, and VAR medium alloy bainitic steels are shown in Table XVI. Initially the volume percent measurements were determined on longitudinal sections of the threaded grip end of the failed fatigue specimen. These data reveal that the VAR alloy 3888800 has by far the lowest level of inclusions, and that the VIM melted alloy 2411 has the next highest volume percentage and the ESR alloy C230 has the highest level of inclusion content. The volume percent inclusions determined by point counting reveal the same order of rating the three alloys in terms of microcleanliness. Unfortunately these volume percent inclusion numbers do not correlate with the rating of the fatigue strengths as shown in Figure 73. The ESR alloy C230 had the highest fatigue strength but it also had the highest volume percent of inclusions. As these data were not correlateable it was thought that counting the inclusion contents adjacent to the fracture surface would be more meaningful than counting them in the threaded grip region of the specimen. The volume percent inclusion data obtained near the fracture surface region of the specimen reveal that the VIM and RSR materials have about the same percentage of inclusions and that again the VAR material has a significantly lower percentage of inclusions.

These data, however, also do not explain the relative fatigue strengths of the three alloys. The reason for the lack of correlation between the volume percent inclusion measurements and the fatigue strengths of the three alloys is thought to be due to the relatively small volume percentage of inclusions for all three alloys. Because sil three heats were really quite clean it is felt that the inclusion size and distribution at the region of initiation of the first fatigue crack is a very localized situation which is not accurately represented by an average volume percentage of inclusions. It is felt that the factor which most probably controls the fatigue life of these very clean steels is the size or orientation of the oxide or silicate inclusion located in the maximum stress region of the fatigue specimen. Unfortunately, experimentally verifying this belief was not possible. Fatigue studies by Johnson and Sewell (20) and Murray and Johnson (27) support this argument by demonstrating that for the same total number of Enclusions (in clean steels) the fatigue life varied over a considerable range.

A limited number of transverse fatigue tests were performed. A detailed study of transverse fatigue properties was not conducted because it was believed that the degree of anisotropy exhibited by 1/2-inch thick plate would have little relevance to the degree of anisotropy usually found in forgings or forging billet stock. The transverse fatigue properties were determined on the low alloy martensitic steel composition in both the VIM and ESR conditions. The longitudinal and transverse fatigue properties of VIM alloy 2351 are presented in Figures 75 and 76, and compared in Figure 77. Alloy 2351 is an additional heat of the same nominal composition as Heat 2409 and the composition is listed in Table I. As shown in Figure 77 a normal amount of anisotropy was observed with the 10' cycle fatigue strength decreasing from 170 ksi for the longitudinal direction to 140 ksi for the transverse direction. Similar data are presented an Figures 78 and 79 for the ESR low alloy martenaitic composition. In this case the observed anisotropy is minimal with the transverse fatigue strengths nearly equaling the longitudinal strengths. Metallographic examination for inclusion contents, size and distribution did not provide an explanation for the different degree of anisotropy between the VIM and ESR materiale.

The unnotched fatigue properties of laboratory produced 300 M steel (VIM heat 2449) are presented in Figure 80. A 10⁷ cycle fatigue strength of about 175,000 psi was obtained. The fatigue properties of the two VAR steels, medium alloy bainitic steel 3888800, and low alloy martensitic steel 3888811, are compared in Figure 81. It can be seen that the fatigue atrengths of the bainitic steel are slightly greater than those of the martensitic steel. Further comparison of the fatigue properties of bainitically heat treated steels as made by the data presented in Table XVII which compares the 10⁷ cycle fatigue strength to ultimate tensile strength ratio (Fatigue Ratio) of the experimental steels in this program to the Fatigue Ratio of several commercial ultra-high strength steels. As fatigue strengths increase significantly with increasing R values these comparisons are only made for tension-tension fatigue tests with an R value of either +0.06 or +0.10. Concerding the experimental alloys of this investigation it can be seen that the bainitic steels generally hav a much higher Fatigue Ratio than the martensitic steels. Although the reason for the lower fatigue strength and Fatigue Ratio of VIM bainitic alloy 2411 is not known, it is felt that the fatigue properties of this heat are not representative of the alloy. This can be seen by comparing the Farigue Ratios of the same composition in the VAR condition (Heat 3888800) and the ESR condition (Heat C230). The 107 cycle fatigue strengths of about 200,000 psi for bainitic steels VAR 3888800, and ESR C230 and C231 are noted to be considerably superior to these obtained on the commercially available martensitic steels such as 4340, 300 M and 18 Ni maraging. Other investigators have also demonstrated that the fatigue properties of lower bainite structures are superior to those of tempered martensite (33). It can also be seen from the information in Table XVII that the Fatigue Ratios of the experimental martensitic steels are quite commendable.

NOTCH FATIGUE TESTS

The notch fatigue strengths of two experimental steels are shown in Figures 82, 83, and 84. Notch fatigue tests were only performed on the VAR steels as there was insufficient plate material to obtain both unnotched and notched fatigue specimens from the VIM and ESR heats. The comparison S-N curves shown in Figure 84 reveal that the notch fatigue strengths of these two alloys come tegether to a common value of about 80,000 psi at 10^7 cycles. This is a commonly observed behavior, that while there may be differences in fatigue properties between alloys at $K_t = 1$, the notch fatigue properties of different high strength steels tend to be very similar. At similar R values the 10^7 cycle notch fatigue strengths of 50,000 psi for alloys 3888800 and 3888811 compare to a value of 60,000 psi for 300 M steel (30) and 40,000 psi for high-purity 18 Ni maraging steel (15).

3. Thermal-Mechanical Working Treatments

An investigation was conducted to determine the effects of deformation of an annealed ferrite-carbide matrix, upon the mechanical properties of a subsequently conventionally heat treated low alloy martensitic and bainitic steel. The genesis of this work was the work of Webster (34) which demonstrated that the deformation of an annealed or tempered martensitic structure for both AFC-77, a martensitic stainless steel, and 300 M steel, produced microscopic voids at the carbide-matrix interfaces; and that these voids were metastable in sustenite at high temperatures and resulted in considerable refinement of austenite grain size by acting as barriers to grain growth. A series of thermalmechanical treatments were given to commercial 300 M steel, an experimental low alloy martensitic steel, and an experimental low alloy bainitic steel in order to effect refinement of the prior sustenite grain size and subsequent mechanical properties.

The first material investigated was commercial 300 M steel, which was processed by the following six thermal-mechanical treatments:

- a. Spheroidize anneal and deform 50% at 1200 F at the completion of the annealing cycle.
- b. Spheroidize anneal, cool to room temperature and deform 50%.
- c. Spheroidize anneal, cool to room temperature, heat to 1200 F and deform 50%.
- d. Temper quenched material at 1200 F and deform 50% at 1200 F.
- e. Temper quenched material at 1200 F, cool to room temperature and deform 50%.
- f. Temper quenched material at 1200 F, cool to room temperature, heat to 800 F and deform 50%.

The 300 M steel processed as shown above was austenitized at various temperatures and examined metallographically to determine prior sustenite grain size. These results are shown in Table XVIII. The data indicate that austenitization must take place at least 50 F below the conventional austenitizing temperature of 1375 to 1600 F in order for significant grain refinement to take place. The hardness data on the oil quenched samples indicate that apparently all carbides are dissolved at temperatures as low as 1475 F. Material processed by each of the six processes was then sustenitized for 1/2 hour at 1475 and 1525 F, oil quenched and tempered for 2 + 2 hrs at 600 F. The resulting tensile and Charpy impact properties are shown in Table XIX. For both the spheroidized annealed and tempered martensitic structures rolling at 1200 F versus rolling at room temperature did not significantly effect either strength or toughness properties. For a given deformation treatment the spheroidized microstructures produced a finer grain size than the tempered martensite microstructures.

Comparing the mechanical properties of the conventionally austenitized, quenched and tempered 300 M steel to these of the grain refined thermal-mechanically processed 300 M steel it can be seen that the yield and tensile strengths were increased generally by 3 to 6%; that the reduction in area values decreased by 9 to 30%; that the elongation values decreased by 8 to 25%, and that the Charpy impact energy values varied irregularly from an increase of 5% to a decrease of 30%. These results indicate that the reduction in prior austenite grain size from about ASTM number 9 to ASTM number 13, for 300 M steel, as a result of these thermalmechanical treatments is not warranted from a mechanical property standpoint.

The experimental low alloy martensitic steel (Alloy 2350 with the same nominal composition as Alloys 2409 and 245) was processed with the following thermal-mechanical treatment cycles:

a. Spheroidize anneal and deform 50% by rolling at 1100 F at the completion of the annealing cycle and air cool effer rolling.

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- b. Spheroidize anneal, air cool to room temperature, heat to 1100 F and deform 50%; air cool.
- c. Austenitize, oil quench, temper at 1200 F, deform 50% at 1200 F; air cool.
- d. Austenitize, oil queach, temper at 1200 F, sir cool to room temperature, heat to 800 F and deform 50%; sir cool.

The 2350 alloy (composition shown in Table I), processed as shown above, was sustenitized at various temperatures and examined metallographically to determine prior sustanite grain sind. These results sre shown in Table XX. The data indicate that sustenitization must take place it least 50 P below the conventionel sustanizing temperature of 1675 to 1760 P for this alloy in order for significant grain refinement to teke place. The horaness data on the oil queached samples indicate that apparently all carbides are dissolved at temperatures as low as 1975 F. Material processed by each of the four processes was then sustenitized for 1 bour at 1600 F, oil quenched and tempered for 2 + 2 hours at 600 F. The resulting tensile and Charpy impact properties are presented in Table XXI. There does not seen to be a significant difference between the four thermal-mechanical treatments and the degree of grain refinement. Compared to the conventionally quenched and tempered saterial, the thermal-mochanically processed material achieved a small and varying increase in yield strength: the ultimate tensile strength increased about 3.5%, the % El. values were sesentially unaffected, and the % R.A. values were increased by about 20%, while the OWN values were increased by about 15%.

In order to determine if a bainitic structure would respond in a different fashion to grain refinement transmiss, low alloy bainitic steel Z412 received the following processing cycles:

- a. Spheroidize anneal and deform S0% by reling at 1200 g at the completion of the annealing cycle and air cool after relieg.
- b. Spheroidize anneal, sir cool to room temperature, heat to 1290 F and deform 50%; air cool.

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c. Austenitize, oil quench, temper at 1200 F, deform 50% at 1200 F, and air cool.

After processing as shown above, the material was sustenitized at two different temperatures (1600 F and 1650 F) and then isothermally transformed at 475 F for six hours to a fully bainitic structure. The resulting tensile and Charpy impact properties are presented in Table XXII. For all three processing treatments the lower austenitizing temperature produced a finer grain size which resulted in an increase in yield strength. It did not significantly affect any of the other tensile properties. There seemed to be some trend, but perhaps not a significant one, of decreasing Charpy impact toughness with increased yield strength and finer grain size produced

by the lower sustenitizing temperature. For a given sustenitizing temperature there did not seem to be any significant difference in mechanical properties as a function of the thermal-mechanical process, nor was there any significant difference when compared to the conventionally heat treated bainitic material. A Hall-Perch plot of the yield strength and grain size data is shown in Figure 85 along with the data for the martensitic steels. Least-square fit lines have been drawn through the data points for each alloy and the dotted lines indicate the 90 percent confidence limits. It can be seen that for the limited range of grain sizes obtained, that grain refinement had a greater effect in increasing yield strength for the bainitic steel than for the martensitic steels. Grain refinement appears to be an ineffective means of strengthening medium carbon martensitic steels. In addition toughness properties are not significantly changed and, as was mentioned previously, it has been shown that grain refinement does not improve the KTSCC values for 4340 steel (16).

IV. COMPARISON OF EXPERIMENTAL MARTENSITIC STEELS WITH COMMERCIAL HIGH STRENGTH STEELS

The alloy development studies discussed previously in Section III demonstrated that of the two bainitic alloy systems and two martenaitic alloy systems investigated, the best strength and toughness properties were obtained on alloys in the medium carbon Ni-Cr-Mo-Si-V low alloy martenaitic system. In addition the stress corrosion studies demonstrated that the low alloy Ni-Cr-He-Si-V martenaitic steels had higher SCC resistant, KISCC, values than the best bainitic steel. It is the purpose of this section, therefore, to compare the properties of the laboratory produced experimental Ni-Cr-Mo-Si-V martenaitic steels with similar properties of currently used commercially produced ultra-high strength steels.

PLANE-STRAIL FRACTURE TOUGHNESS

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Thisk section isnding gear components are sufficiently large such that the state of stress around most flaws, if present, would be plane strain. The conditions, therefore, that would lead to brittle fracture will be determined by the plane-strain fracture toughness parameter, $K_{\rm IC}$. A summary of the plane-strain fracture toughness data for H-11, '340, 300 M, HP 9-4-45, and 18 Ki marsging steels at room temperature and -65 F are illustrated in Figures 86, and 87 respectively. To the extent that was possible all of the $K_{\rm IC}$ data for 4340 steel were obtained from references (28, 35, and 36). The data for H-11 were obtained from references (37 and 38), for 300 M steel references (28, 37, 39, and 15); for 18 Ni maraging steel references (15, 37, 38, 40, and 41).

The room temperature fracture toughness data shown in Figure 86, reveal the usual trend of a creasing toughness with increasing strength level for all steels. At the 260 to 280 kri strength level HP 9-4-45 steel heat treated

in the bainitic condition has the highest fracture toughness values. The drawback of the KP 9-4-45 bainitic alloy is that in thick sections the highest strength that can be guaranteed is 260 ksi. The strength-toughness relationships for the 18 Ni maraging steels are seen to be very good. however, the 18 Ni meraging steels have not been used for sircraft landing gear components because of problems with thermal embrittlement in thick sections, low notch fatigue properties, and a low strain hardening exponent. The other three steels, H-11, 4340, and 300 M have all been used in production for aircraft landing gear forgings and, of course, the 300 M alloy is used extensively today in current sircraft. It can be seen from the fracture toughness date at both temperatures (Figures 86 and 87) that the strength-fracture toughness relationships for the new low alloy Ni-Cr-Mo-Si-V martensitic steels are considerably superior to the strength-toughness relationships of 4340, H-11, and 300 M steels. It should be noted that the fracture toughness data on the experimental-laboratory martensitic steels are thought to be conservative or low side franture toughness values. Our experience with both 300 M steel and the HP 9Ni-4Co steels has indicated that for the same composition the fracture toughness properties obtained on production material, from large ingots using electric furnace air melt-vacuum arc remelt practice, are superior to those obtained on small laboratory VIM ingots. The fracture toughness properties of the experimental steels are, therefore, expected to increase by 10 to 15 percent when determined from production VAR material. The strength-toughness relationships in Figures 86 and 87 reveal that the strength level of the new low siloy martensitic steels has been increased by about 25,000 pai (from about 250 ksi to 310 ksi) while maintaining the same level of fracture toughness.

The crack propagation resistance of these steels can be compared in terms of critical crack size rather than the absolute magnitude of K_{IC} . Fracture mechanics analyses demonstrate that fracture will occur when

$$K_{IC} = 1.1 \sigma \sqrt{\pi} C \frac{1}{\sqrt{Q}}$$
(1)

where:

σ = gross area applied stress C = critical crack depth Q = geometric flaw shape parameter

In airframe design, the applied stress is often limited to a fraction of the material ultimate tensile strength (σ_{II}) ; i.e.,

$$\sigma = f \sigma_{\mu}$$
(2)

where f is a design parameter. Therefore, equation (1) can be written as:

 $K_{IC} = 1.1 f \sigma_{\mu} \sqrt{\pi} C \frac{1}{\sqrt{Q}}$ (3)

or:

$$C = \frac{Q}{1.21 \pi f^2} \left(\frac{K_{IC}}{\sigma_{\mu}}\right)^2$$
(4)

The critical crack size is therefore proportial to $(K_{IC}/\sigma_{\mu})^2$. Using the room temperature ultimate tensile strengths and K_{IC} values from Figure 86, the critical crack size parameter $(K_{IC}/\sigma_{\mu})^2$ was calculated and are compared in Figure 88. This comparison of critical flaw size parameters reveals that the new Ni-Cr-Mo-Si-V martensitic steels at a tensile strength level of about 310 ksi have the same critical crack size as 300 M steel at 280 ksi, 4340 at 260 ksi and H-11 at 240 ksi. In other words the new experimental steels have the same flaw size tolerance as the commercial H-11, 4340, and 300 M steels at considerably higher strength levels.

CHARPY IMPACT DATA

The room temperature Charpy impact data for the new experimental martensitic steels are compared to similar dats for commercial high strength steels in Figure 89. There were insufficient CVN data on commercial steels at -65 F to make a comparison at this test temperature. All of the CVN data are from longitudinally oriented specimens and the data for the commercial steels were taken from references (15, 28, 38, and 42). The data in Figure 89 reveal that a rather large degree of scatter exists for 300 M steel, however, this is not unexpected as the Charpy impact test is known to be more sensitive to inclusion contents than the plane-strain fracture toughness test. The data also reveal that for a given strength level the Charpy impact energy values for the 18 Ni maraging steels are rather low compared to 4340, HP 9-4-45, and 300 M steels. This is now a commonly observed behavior that while the 18 Ni maraging steels have high fracture toughness, they have relatively low CVN energy values. This characteristic of 18 Ni maraging ateels is probably a result of the ease of plastic instability in this material. It can be seen from Figure 89 that the strength-Charpy impact toughness relationships of the new martensitic steels are significantly superior to all of the commercial steels.

STRESS CORROSION RESISTANCE

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It is well known that ultra-high strength steels are very susceptible to stress corrosion cracking. To avoid stress corrosion cracking in ultra-high strength landing gear components ective treatments such as cadmium plating and painting have been ap id, ond shot peening has been used to induce residual surface compressive stresses in order to suppress crack initiation. In recent years the SCC resistance of sigh strength materials has been determined by the use of fatigue cracked fracture toughness specimens, which have the two-fold advantage of reducing the inherent appreciable scatter incurred with the use of unnotched specimens, and providing a SCC resistance parameter which is quantitative and provides the possibility of use in design.

The threshold stress intensity values, KISCC, for the Ni-Cr-Ho-Si-V martensitic steels are compared to the KISCC values for the commercial high strength steels in Figure 90. The KISCC values for the commercial steels were obtained from references (12, 15, 16, 43, 44, and 45). The data reveal that in the strength level range of 260 to 280 kai, 4340, 300 M, and H-11 steels have $K_{\rm ISCC}$ values in the range of 10 to 20 ksi $\sqrt{10}$. At strength levels in the vicinity of 300 ksi, however, the maraging steels SCC resistance decreases to $K_{\rm ISCC}$ values of 7 to 12 ksi $\sqrt{10}$. The new low alloy martensitic steels at much higher strength levels (298 to 332 ksi) have $K_{\rm ISCC}$ values in the range of 16 to 19 ksi \sqrt{in} . The strength - SCC resistance relationships of the newly developed martensitic steels have, therefore, been improved compared to both 18 Ni maraging and 4340, 300 M, and H-11 steels. Even with this improvement, however, these levels of SCC resistance are not inherently high and therefore these steels will not be able to be applied from an engineering reliability standpoint by means of the fracture mechanics approach utilizing the knowledge of the stress state, defect size and periodic nondestructive inspection techniques. These new ultra-high strength low alloy steels will, the effore, have to be utilized in the same ranner that the present high strength steels have been so successfully utilized from a SCC resistance standpoint, by means of plating, painting, and shot reening.

FATIGUE PROPERTIES

The complete notch and unnotch fatigue properties of both experimental bainitic and martensitic steels were presented and discussed previously in section C-2. In Figure 91 the unmotch fatigue properties of the Ni-Cr-Mo-Si-V martensitic stells are compared to similar fatigue properties for several commercial steels. All of the data shown in Figure 91 are for tensiontension fatigue tests at R values of either +0.06 or +0.10. The rather large body of rotating-beam (R = -1.0) fatigua data on high strength steels could not be utilized for such a comparison. The fatigue data on the commercial steels were obtained from references (15, 30, 31, 32, and 38). The comparison reveals that the experimental martensitic steels have considerably higher fatigue strengths than the commercial steels. The experimental martensitic steels demonstrated 10/ cycle fatigue strengths in the range of 170 to 190 ksi while the commercial steels exhibited 10' cycle fatigue strengths in the range of 90 to 130 ksi. Both groups of steels seem to have a trend of decreasing fatigue strength with increasing tensile strength level. This effect has been observed previously (46), however, neither the reason or the significance of this trend is understood at the present time. It is believed that the reason the experimental martensitic steels have greater fatigue strengths than the commercial steels is a higher degree of microcleanliness in the experimental steels. Except as it effects tensile strength level, composition is known to have little effect on fatigue properties of ultrahigh strength steels. This basic difference between the laboratory produced steels and the commercially produced steels can be seen by comparing the fatigue strengths of laboratory produced 300 M steel versus conmercially produced 300 M steel. It is anticipated, therefore, that the new low alloy martensitic steel when melted by commercial steelmaking practice, in large ingot sizes, will have fatigue strengths in the range of 130 to 150 ksi, when tested under similar conditions.

V. SUMMARY AND CONCLUSIONS

Exhaustive and detailed alloy development and processing investigations were conducted in order to develop an ultra-high strength steel in the 300 to 320 ksl ultimate tensile strength range, with improved fatigue strength, fracture toughness, and stress corrosion resistance for greater reliability in forged lending gear components. Two bainitic alloy systems and two martensitic alloy systems were thoroughly investigated in order to develop the best combination of mechanical properties at trasile strength levels in excess of 300,000 psi. Of the four alloy systems investigated, steels from the low alloy medium carbon Ni-Cr-Mo-Si-V martensitic system developed the best combination of fracture toughness, fatigue strength and stress corresion creaking resistance.

The stress corresion strict lemonstrated that while lowering phosphorous and sulfur levels is beneficial to toughness properties it has essentially no affect on SCC resistance, as indicated by the $K_{\rm TSCC}$ parameter, for either high strength bainitic or martensitic steels. Similar studies in low alloy martensitic steels demonstrated that variations in silicon, chromium, and molybdenum significantly effected plane strain fracture toughness properties, while having no effect on $K_{\rm SSCC}$ values. The low alloy Ni-Cr-Mo-Si-V steels had higher SCC resistance than the best medium alloy bainitic steel.

The processing studies conducted on two bainitic alloys and one martensitic slloy revealed that the vacuum are remelted steels had the highest level of fracture toughness, while the electrosiag remaited materials had the lowest level of frecture toughness with the vacuum induction melted material being intermediate in toughnees properties. Considering the influence of melting practics on fatigue properties, for the two beinitic steels the ESR material had the highest fatigue strongths, and the VIH material the lowest fatigue strengths. For the martensitic steel the VAR material had the highest fatigue strengths followed by the ESR material and then the VIM material. The experimental steels demonstrated unnotch 107 cycle fatigue strengths in the range of 170,000 to 210,000 ysi. The motch ($K_{t} = 3.0$) fatigue strengths of the VAR baiaitic steel and the VAR martensitic steel were essentially the sume (80,000 psi) at 107 cycles. Thermal-mechanical working treatments demonstrated that the strength and toughness properties of ultra-kigh strength low alloy marteneitic and bainitic & sais are little influenced by refinement of the prior austenite grain size.

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Comparison with similar properties of currently used and commercially melted ultra-high strength steels, revealed that the strength- ighness properties of the new low alloy martensitic steels were superior to the strength thoughness properties of the commercially produced steels. Comparison of the threshold stress intensity (K_{ISCC}) SCC resistance parameter indicated that the new martensitic steels had higher K_{ISCC} values than the current commercial steels at a cc siderably higher strength level. The tensiontension (k = -0.10) unbotch fatigue strengths at 10⁷ cycles were in the range of 170 to 190 ksi for the newly developed maxtensitic steels compared to 90 to 130 bsi for the commercial steels.

From the alloy development and processing studies a new improved ultra-high strength martensitic steel with a nominal composition of

<u>c</u>	Mn	P	<u>s</u>	<u>Si</u>	Ni	Gr	<u>Ho</u>	Ā
0.40	0.35	<.010	<.010	2.25	1.8	0.80	0.25	0.22

has been developed. When heat treated by normalizing, austenitizing, oil quenching, refrigerating, and double tempering at 600 F the alloy develops the following average longitudinal, room temperature properties based on laboratory sized heats:

U.T.S., ksi	311
Y.S., ksi	268
Elongation, %	12
Reduction of area, %	44
CVN, ft-lbs	20
K _{TCs} ksi√in.	60
KISCC, ksi /ia.	17
Axial fatigue strength at 10^7 cycles, $K_t = 1$, ksi	170
Axial fatigue strength at 10^7 cycles, $K_t = 3.0$, ksi	80

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	Remarks	N1-Cr-Mo-V	Bainaging	Steels																						.					A
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WINDER OF	>1	0.10	0.10	0.10	0.10	0.10	0.10	0.10	0.10	0.10	0.10	0.10	0.10	0.10	0.10	0.10	0.41	0.10	0.10	0.10	50°0	0.09	0.10	0.10	0.12	0.10	0.11	0.19	0.19	0,19	0,22
TUTU	31	ł	ł	ł	\$	1	1	\$	\$	1	¥	z	ł	ŧ	ł	ł		ł	ł	1	1	ŧ	1	ŧ	t	1	1	ŧ	1	\$	1
TUTUT	Wo	0.54	0.55	0.54	0.55	1.02	1.05	0.52	0.54	1.06	1.08	1.10	2.05	1.85	2.05	1.17	1.17	1.13	1.13	1.15	1.10	1.10	1, 10	1.10	1.10	1.00	1.90	1.00	1.40	2.90	1.40
TO ONOT 1	5	0.95	0.94	1.07	1.06	1,05	2.05	1.00	1.00	1.03	2.05	3.20	3.10	2.15	2.10	2.10	2.10	2.10	2.10	2.10	2.10	2.10	2,10	1.90	2.00	1.90	3,00	1.0	1.0	1.0	•
TEOJILOO	IN	1.07	2.90	1.13	3.00	1.10	1.10	1.05	2,95	1.20	1.18	0.86	0.85	3.00	3.05	1.20	1.20	1.21	1.16	3.05	3.08	3.20	3, 20	3.20	1.30	3.07	3.08	2.85	2.85	2.85	3,07
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	SI	0.27	0.25	0.27	0.24	0.30	0,30	0.22	0.17	0.26	0.20	0.26	0,24	0.26	0.23	1.40	1.35	1.45	1.35	0.21	0.20	0.21	G. 16	Ũ. 26	2.00	0.27	0.25	0.31	0.31	0.31	0.28
	Wu	0.81	0.79	0.81)	0.77	0,90	0.90	0.85	0.76	61.0	0.70	0.80	0.74	0.77	0.72	0.91	0.88	0.81	1.30	0.80	0.73	0.70	05.0	05 0	1.50	0.91	0.82	0.73	0.74	0.75	0.78
	ပျ	0.55	0.54	0.48	0.48	0.48	0.48	0.45	0.43	0.44	0,44	0.45	0.43	0.45	0.44	0.44	0.44	<u>C, 34</u>	46.0	0.33	0.31	0.33	0.41	0.34	0.45	0.43	0.32	0.50	0.48	0.49	0.49
	Heat No.	8332	8333	8334	8335	8336	8337	8338	8339	8340	8341	8342	B343	8344	B345	B346	B347	8348	B349	8350	B351	B 352	0353	B354	5355	3356	3357	2389	2390	1963	1392

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ChEMICAL COMPOSITIONS OF EXPERIMENTAL ALLOYS

Remarks	N1-Cr-Mo-V Bainaging Steela	Medium Alloy Bainitic Steels	Medium Alloy BainitLc Steels	Ni-Cr-Mo-W-V Martensítíc Stcels
8	3,85 3,85 5	<i>、、、、、、、、、、、、、、、、、、、、、、、、、、、、、、、、、、、、</i>	22121222222222222222222222222222222222	* * * * * * * * * * * * * * * * * * *
<u>Aï</u>	1.1	• • • <u>-</u> • • • •		4 8 4 2 5 6 2 T 8 7
শ	0.21 0.21	0.084 0.087 0.081 0.080 0.086 0.086 0.086	0.10 0.10 0.10 0.10 0.11 0.11 0.11 0.11	0.09 0.09 0.00 0.09 0.01 0.00 0.09 0.01 0.01
A	8 2			0,54 2,10 2,10 0,55 0,53 0,53 0,53 0,53
W	1.43 1.48	0.15 0.15 0.33 0.31 0.31 0.31 0.35	0.28 0.25 0.25 0.25 0.25 0.25 0.25 0.25	0,48 0,990 1,955 1
15	<u>0.35</u> 0.75	0.116 0.15 0.17 0.33 0.17 0.33 0.33 0.33 0.33 0.33 0.33 0.33 0.3	0.27 0.27 0.25 0.25 1.50 1.50 1.50	0.48 2.00 0.98 0.48 0.48 2.10 2.10
NT	2.90 2.90	10.21 9.75 9.90 10.11 10.11 9.55 9.55	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~	1.95 2.05 2.05 1.97 1.95 1.95
S	.006	<.010	× • • • • • • • • • • • • • • • • • • •	
H4	. 002. 002		 003 003	
<u>S1</u>	0.28	0.20 0.20 0.15 0.15 0.15 0.15 0.15 0.15 0.15 0.1	0.08 0.08 0.11 0.28 0.28 0.28 0.28 0.28	0.33 0.33 0.33 0.33 0.33 0.33 0.33 0.33
W	0.74 0.74	00,320 22,222 22	0.17 0.15 0.15 0.24 0.24 0.29 0.29 0.29 0.29	0,71 0,75 0,75 0,75 0,75 0,75 0,91 0,91
ငး	, 49 , 50	0,42 0,44 0,44 0,45 0,45 0,45 0,45	45 45 45 45 45 47 47 47 47	33 37 37 37 37 37 37 37 37 37 37 37 37 3
4.4	04 00	00000000000000000000000000000000000000	<u> </u>	
Rea No	239 2,39	826 825 825 825 825 825 822 822 822 822 822	2222 2222 2222 2222 2222 2222 2222 2222 2222	0770 0770 0770 0770 1770 1770 1770 1770

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CHEVICIL DOMPOSITIONS OF EXPERIMENTAL ALLOYS

Remarks	Ni-Cr-Mo-W-V Martensitic Steels	N1-Cz-Mo-W-V Martensitic Steels		>
8		1 5 2 1		
A1	11111	8 8 8 8		
>1	0.10 0.10 0.09 0.09 0.10	0.11 0.11 0.11 0.11	0.00 0.00 0.00 0.00 0.00 0.00 0.00 0.0	0.10 0.10 0.10 0.10 0.10 0.10
31	1.15 1.15 1.15 1.15 1.15	2.05 1.03 2.05 1.05	1.26 1.26 2.15 2.15 2.15 2.15 2.15 2.15 2.15 2.15	1.02 1.02 1.02 1.02 1.02
윘	0.97 1.96 1.97 1.00	0.95 1.00 1.95 0.95	1. 15 2. 15	• = = = = = =
빙	0.97 0.97 0.97 1.00 -	0.49 2.05 0.01	1.24 1.22 2.63 2.65 2.65 2.65 1.23 1.23 2.65 2.65 1.23 1.23 1.23 1.25 1.23 1.23 1.25 1.23 1.25 1.23 1.23 1.25 1.25 1.25 1.25 1.25 1.25 1.25 1.25	0.98 0.97 0.98 0.98 0.96
FN	1.92 1.97 1.90 1.90 1.85	1.82 1.85 1.85 1.81	200 200 200 200 200 200 200 200 200 200	1.90 1.90 1.90 1.90 1.80
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ર્ભા	×.010	.002 .002 .002 .002		.005 .006 .003 .004
St	0.30 0.30 0.26 0.27 0.27	0.27 0.31 0.31 0.27	0.26 0.27 0.25 0.25 0.27 27 27 27 27 27 27 27 27 27 27 27 27 2	0.27 0.26 0.27 <u>1.62</u> 2.51
W	0.80 0.80 0.82 0.82 0.91 0.91	0.77 0.76 0.78 0.78	0.72 0.73 0.73 0.73 0.73 0.75 0.75 0.75 0.75 0.75 0.75 0.75 0.75	0.29 0.75 0.75 0.72 0.72
ol	0.37 0.37 0.46 0.43 0.43 0.43	0.43 0.43 0.43 0.43	0.42 0.43 0.44 0.44 0.45 0.45 0.45 0.45 0.45 0.45	0.41 0.42 0.42 0.42 0.42
Heat No.	V716 V717 V748 V749 V750 V751	275 277 278 279	295 295 295 295 295 2010 2010 2010 2010 2010 2010 2010 201	2112 2113 2114 2115 2115 2117 2117
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CHEMICAL COMPOSITIONS OF EXPERIMENTAL ALLOYS

Remarks	N1-Cr-Mo-W-V Martensitic	Steels	N1-Cr-Mo-S1-V Martensitic stor	2 		>	Cb-Series Martensiric Steels	Ni-Series Marte stitic Steels
ଥ	11	1111	8 E 2	0,97 1,01	1.06		Cb 039	
<u>A1</u>	Ŧı		.010	110	.0118 .018 .018	.026 .032 .012 .024 .033		
거	0,10	0,19	6.21 0.21 0.21	0.20	0.21 0.21 0.20	0.10 0.20 0.20 0.20	¥ 2 5	0.21 0.21 0.19 0.19 0.20
31	1.02				\$ ÷ \$		111	
N	1 2 1	0°25 0°25 0°25	0.25 0.26 0.26	0.29 0.29 0.29	0.53 0.54 0.39	0.41 0.42 0.47 0.47 0.47 0.47 0.47 0.47 0.47 0.47	0.27 0.26 0.25	0.26 0.26 0.30 0.25 0.25
빙	0.98 1.02	1.00	0.60 0.60	0,62 0,89 1,60	0,01 0,67 0,92	0,97 2,00 2,97 0,98	0,72 0.72 0.70	0.84 0.04 0.77 0.77 0.77
IN	1.89 1.90	1.72	2.00 1.80 1.80	2.15 2.10 2.05	20000 20000 20000	2,05 2,05 2,05 2,00	2,00 1,98 1,98	24.25 20.55 20.55 2.45 2.45 2.45 2.45 2.45 2.45 2.45 2
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<u>S1</u>	0.26 0.26 0.26	2,18 2,11 2,19	1,05	1.65 1.65	1,65 1,65 0,31 1,60	2.65 2.65 2.65 2.65	2,55 2,50 2,40	22,55 2,55 2,55 2,55 2,55 2,55 2,55 2,5
WU	0.74 0.74 0.75	0.53 0.53 0.53	0.34 0.34 0.35	0,36	0,36 0,36 0,36 0,36	0,35 0,35 0,36	0.34 0.33 0.35	0,36 0,36 0,36 0,34 0,34
0]	0.42 0.43 0.44	0,39 0,40 0,40	0.38 0.38 0.38	8°°0 6°°0 6°°0	0,38 0,38 0,39 0,37	0,37 0,38 0,38 0,37	0.41 0.40 0.40	0,41 0,39 0,38 0,38 0,38
Heat No.	2118 2119 2120	2386 2387 2388	243 244 245	247 248 248	255 251 255 255 255	257 258 259 260	2332 2333 2334	2273 2274 2275 2329 2330 2331 2331

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Chine L COMPOSITIONS OF EXPERIMENTAL ALLOYS

Remarks	V-Series Martensitic Steels	Mn-Serles Martensitic Steels	N1-Cr-Mo-S1-V Martens1t1c Steels	300 M
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<u>A1</u>	\$ á \$	4 I I		
>!	0, 10 0, 20 0, 29	0.20 0.21 0.21	0.22 0.22 0.22 0.21 0.21 0.21 0.21 0.21	0.24 0.20 0.06
3;		111		
0 W	0.28 0.28 0.28	0.26 0.26 0.26	0.43 0.45 0.68 0.68 0.68 0.68 0.68 0.68 0.68 0.68	0.33 0.25 0.40
빙	0.78 0.81 0.83	0,75 0,80 0,80	$\begin{array}{c} 1.00\\ 1.02\\ 1.35\\$	1.90 0.82 0.73
II	1.90 1.95 1.90	1.98 2.05 2.05	8 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9	2.00 2.00 1.72
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: W	0.26 0.26 0.25	0.26	000000000000000000000000000000000000000	0.32
ပျ	0.38 0.38 0.38	0.42 0.39 0.39	000000000000000000000000000000000000000	0.41 0.39 0.39
Heat No.	2276 2277 2278	2270 2271 2272	我就我我我我我 业众当分岁窗了每分心乱激励场与	2525 2551 2449

TARLE I (Continued)

CHEMICAL CORPOSITIONS OF EXPREMENTAL ALLOYS

<u>Remerks</u>	Low Alloy Merterafte Medium Alloy Bainite Low Alloy Bainite	Low Alloy Martensite Medium Alloy Bainite Low Alloy Bainite	Low Alloy Martensite Medium Alloy Bainite Low Alloy Bainite	Low Alloy Martenzite Medium Alloy Bainite Low Alloy Martenzite	Low Alloy Martensite Low Alloy Martensite
찌		008 004 110	. C34 . 003	• 004 • 004	
ol		.007 .005	· 002 • 009	.002 .004 .003	
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A1	04 04 04	20] 20]	8 7 X 6 1	.02 .02	3 3
>;	<u>1vete</u> 0.19 0.17 0.17	Analya 0.21 0.10 0.30	<u>uelt An</u> 0.21 0.09 0.20	Lyn1a 0.22 0.24 0.24	0.20 0.20
З Т С	IM Ane 0.28 0.24 1.36	Malt 0.27 1.50	Las. Rat 0.26 0.24 1.52	LR. Ane. 0.27 0.36 0.34	0.26 0.27
성	0,76 0.36 0.99	A1X 0.78 0.38 1.05	0.86 0.86 1.05	0.80 0.39 1.80	0.76 0.77
NT	1.75 6.75 2.75	1,83 6,80 2,90	. Elle 1.90 6.90 2.90	1.70 7.10 1.67	2.00
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13.1	0.00 0.00 0.00	2.67 0.14 0.34	2,47 0,03 0,28	2,57 0,01 2,13	2,60 2,63
W.	0.39	0,48 0,37 0,79	0,50 0,25 0,80	0.53 0.25 38	0.37 0.39
S	0.39 0.52 0.49	000 885 885 885	0,44 0,50 0,51	0.38 0.35 0.35	0.40 0.42
He#1 	2410 2411 2412	6231 6230 6231	6229 6230 6231	28888008 38388008 38388008 38888211	2350 2350

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CHEMICAL COMPOSITIONS OF EXPERIMENTAL ALLOYS

Remarks	6-4-45 (F + S)	Steels				>
ဒို	4.36	4.30	4.30	4.22	4.17	4.20
AI	3	£	8	ı	1	£
>1	60.0	0.09	0.08	0.09	0.09	0.07
윘	0.33	0.40	0.34	0.33	0.40	0.32
빙	0.28	0.36	0.30	0.30	0.28	0.31
ŦN	8.55	8.41	8.34	8.30	8.20	7.97
വ	.002	.020	,025	.025	600.	.010
ei	.003	.013	,022	.004	.020	.004
S	0, 10	0°0	0.05	0°0,0	0.08	0.06
W	0.27	0.23	0.27	0.27	0.25	0.27
01	0.41	0.42	0.44	0.42	0.42	0.44
Heat No.	V7 23	V724	V725	V726	V727	V746

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TABLE	Ι	I
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MECHANICAL PROPERTIES OF LOW ALLOY BAINAGING STEELS

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		Isotherm	elly T	rensfor:	ned					
		Low	er Bsi	nite			Bainaged	950 F,	4 Hrs	
Heat	Y.S.	U.T.S,	<u>E1</u> .	R.A.	CVN	Y.S.	U.T.S.	£1,	R.A.	CVN
No,	ksi	ksi	<u>%</u>		ft-1b	ksi	ksi	7,	7	ft-lb
B332	264	312	9	21	9	216	229	12	31	13
B333	246	304	10	29	15	214	227	12	33	14
B334	246	293	10	35	14	213	22.3	12	35	16
B335	226	272	12	44	17	205	218	14	43	17
B336	244	293	11	37	11	229	242	14	39	14
337	214	286	12	41	9	214	233	12	33	12
B338	223	262	14	49	15	201	212	15	31	<u>1</u> 8
8339	205	252	13	45	18	197	209	15	46	19
B340	220	265	12	42	14	218	230	14	45	17
B341	206	274	12	42	8	209	244	14	43	12
B342	193	286	14	37	16	200	270	11	28	10
B343	225	285	13	33	15	213	276	10	23	10
B344	151	285	13	33	20	199	284	11	29	9
8345	175	273	15	37	10	202	275	12	29	9
B346	214	294	14	28	17	178	294	7	6	7
B347	219	281	15	37	20	197	293	8	12	9
B348	182	243	17	43	24	194	254	12	23	13
B349	157	244	17	41	Ž3	165	272	12	24	10
B350	148	232	14	41	25	179	232	15	44	17
B351	175	236	14	42	20	178	228	15	44	19
B353	201	266	14	44	18	2:7	257	13	38	8
8354	158	222	18	46	30	134	282	14	35	12
B355	157	301	7	6	7	139	295	3	0	3
B355	180	245	15	41	21	157	279	9	12	7
2257	1 2 2	253	15	27	19	261	270	14	33	12

Average longitudinal, room temperature properties of air melted heats.

TABLE ILT

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MECHANICAL PROPERTIES OF LOW ALLOY BAINNOING STEELS

	CUN	ft-1bs	12.5	14	C.I	N	20.5	9.5	22	o	, 2 7	21	8.5 2	-20		
×4 0	R.A.	2	4 ľ	39	60	33	17	17	с. С	34	38	36	5	Ĩ		
31 1100	EL.	and	۲. د	12	77		3	2Ì	27	51	12	12	2.2	2.7		
A ped	0. Y. S.	ket	227	229	233	262	251	251	223	226	224	LCL	9EC	232		
	Y.S.	kg 1.	217	216	219	213	227	326	223	2.2.6	126	328	126	226		
	CVM	ft. lbs	15	15	13	12.5	11	2.2	22.5	12.5	57	24	E.	27		
A D	R. A.	4 miles	50	47	WE	42	34	38	80 0	21	39	474) 474)	£.43	th da		
d at 95	EI.	X.	16	14,	12	12	11	12	12	6	12	12	с С	14		
480 V80	U.T.S.	kei.	235	235	241	243	261	261	233	231	239	242	243	244		
	¥. S.	ku1	221	221	226	226	233	233	233	23%	236	241	236	236		
	CVN	£t. lbs	24,5	13.5	14.5	13.5	12.5	13,5	2.4.5	11.5	ტ	12.5	12.5	12.5		
ned	R.A.	No.	96 39	41	35	38	36	35	43	36	47	38	40	30		
Kana Eur	E 1.	7	11	12	:	11	11	12	11	12	11	٦ï	1.2	10		
7 8V	U, T. S.	All Land	312	60E	313	eie	305	304	299	297	306	305	315	313		
	¥. S.	kui	22.2	250	247	251	2,20	226	258	258	242	262	261	262		
	Ncat	No.	2389		Z.390		1953		2352		2393		962Z 4	7		

LongLtudinal, room temperature properties Aging time, 4 hours Second.

N. 68

	<u>MEGNANIC</u>	AL PROPERTIES O	P MEDIUM ALL	UI BAINIILU S	IELLS
Hest	Y.S. ksi	U.T.S. ksi	E1. Z	R.5. Z	CVN ft-lbs
B266	204	264	12	32	8
	197	264	12	32	8
B267	181	268	15	46	14
	182	267	15	44	14
B268	221	266	10	36	12
	222	265	10	40	13
B270	228	279	11	45	18
	229	271	11	43	18
B272	234	277	12	43	16
	234	2?7	12	4 . '	16
B273	237	277	11	45	18
	235	277	12	50	18
B297	217	285	12	40	22
	218	283	12	41	20
B299	216	256	13	52	19
	213	256	13	54	24

TABLE IV

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Room temperature properties of air melted heats tested in longitudinal direction

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TABLE V

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MECHANICAL PROPERTIES OF MEDIUM ALLOY BAINITIC STEELS

Heat <u>No.</u>	Isothermel-Transformation Temperature, ^o F	Y.S. ksi	J.T.S. ksi	E1. 7	R.A. <u>7</u>	CVN ft-1bs
Z211	445	260	301	15	66	11
	495	238	272	11	45	16
Z212	415	266	316	11	39	21
	465	251	290	11	38	15
Z213	435	288	332	7	16	15
	485	268	303	5	11	7
2214	415	288	336	ÿ	23	7
	465	280	318	8	24	9
2219	540	232	261	14	51	20
	590	208	232	15	52	20
2220	540	247	284	13	50	42
	590	218	254	14	50	2.5 TO
Z22 1	520	239	286	13	42	14
	570	221	263	13	45	17
Z222	445	218	285	12	41	16
	495	206	265	12	45	17
Z223	470	223	283	12	45	12
	520	207	254	13	46	15
Z224	490	223	281	12	40	15
	540	205	262	11	47	20
Z225	490	231	284	11	40	15
	540	212	264	12	45	15

Average longitudinal, room temperature properties

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TABLE VI

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New York

MECHANICAL PROPERTIES OF MEDIUM CARBON Ni-Cr-Mo-W-V MARTENSITIC STEELS

			400	F Te	rsem			600 F	Temp	é'T	
Heat	Test	Y.S.	U.T.S.	E1.	R.A.	CVN	Y.S.	U.T.S.	E1,	R.A.	CVN
No.	Dir.	kei	ksi	7.	7.	ft-1bs	ksi	ksi	- 7	7.	ft-1bs
V707	1	203	257	14	52	27	204	235	14	57	24
	Т	20/	254	13	48	24	204	2.35	13	48	20
V708	I.	215	290	11	36	10	215	262	11	41	10
	ፓ	220	235	11	31	15	211	259	11	41	10
V709	l	225	285	11	28	13	219	270	12	40	13
	T	218	296	11	26	11	213	267	11	30	10
V710	L	226	279	10	34	14	242	263	11	44	12
	T	222	285	8	~ `	11	240	250	8	31	10
V711	L	213	274	12		15	211	258	12	44	11
	Т	213	273	9	27	12	207	254	9	27	10
V712	L	221	263	13	45	21	216	246	14	54	19
	T	222	268	11	42	18	216	246	12	47	17
V713	L	213	271	12	45	22	215	246	13	53	20
	T	216	261	12	40	18	214	245	12	48	18
V714	L	230	284	8	21	8	234	271	10	35	9
	Т	228	293	6	15	7	229	27G	8	26	8
V715	L	213	286	11	32	13	212	266	11	38	11
	T	225	283	10	32	11	210	265	11	40	12
V716	L	195	267	13	40	21	19ó	242	12	43	16
	Т	205	270	15	41	19	198	243	13	41	15
V717	L	201	265	13	45	23	198	241	13	53	17
	Т	202	265	12	44	21	197	237	12	51	16
V748	L	217	310	8	18	12	218	282	11	37	12
	Т	226	307	8	15	12	220	279	8	17	9
v749	I.	231	283	10	26	12	237	267	11	40	11
	T	229	294	11	25	3	238	269	11	35	10
V750	L	235	300	8	18	9	255	282	9	31	10
	Т	247	304	8	16	6	250	278	7	18	8
V751	L	236	290	10	34	12	244	264	11	47	12
	Т	238	287	10	37	11	245	265	10	40	10
275	L	242	300	11	30	13	242	267	10	38	12
277	L	211	307	12	30	16	207	274	9	27	14
278	L	230	301	9	22	14	236	277	9	34	9
279	L	228	294	10	35	13	231	261	10	44	9

Average (2 specimens per condition) room temperature properties Heats 275 through 279 are re-make heats of V710, V748, V750 and V751 respectively

TABLE VII

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MECHANICAL PROPERTIES OF MEDIUM CARBON NI-Cr-MO-W-V MARTENSITIC STEELS

	t-1bs	-65 F	ç	2 1	~	. vc) o	, v	: r	4	· •	y ve	ر د	- eri	- 4	· œ	4	10	12	. ~	10	2	01	, r		12
	CVN. F	+70 F	61		21	; 0	. 00	10	9	• œ	ο σ	9	10	2	51	10	01	รา	15	13	14	5	12	12	71	Ĭ
ember	R. A.	7	1 .2	: -3 - C	5	29	29	16	25	26	23	31	39	28	36	02	35	44	50	43	43	68	36	30	47	46
DO F T	El.	2	0	γσ	01	ဆ	10	10	3		8	10	11	σ	11	12	G	11	13	11	11	12	10		12	11
ē	U.T.S.	ksi	275	281	275	275	284	285	287	263	290	293	284	281	281	276	270	257	254	252	257	303	323	259	264	262
	Y.S.	ksı	234	253	247	243	237	245	241	240	244	244	245	239	238	241	249	227	218	218	220	258	278	222	236	238
	- 168	-65 F	1	1	1	÷	1	1	1	ŧ	1	t	1	1	t	1	ı	1	1	;	1	1	4	I	1	1
	CVN, Ft	+70 F	11	æ	4	0	9	4	7	9	ľ0	4	12	8	70	12	10	12	12	14	4	12	10	8	14	14
enper	R.A.	2	29	24	20	18	29	17	1	21	24	18	36	ł	33	25	1	41	37	37	41	32	24	37	40	36
DO F T	El.	2	6	2	7	2	6	2	1	8	6	ထ	10	ŧ	10	æ	ł	. 11	2	20	00	10	σ	11	11	11
500 F	U.T.S.	ksi	294	297	292	284	297	297	294	289	287	292	301	265	289	286	283	270	271	267	270	308	319	273	274	275
	Υ. Υ.	kei	249	271	. 260	254	235	249	242	241	230	232	246	245	235	25 î	263	234	228	226	228	255	270	234	237	239
	- 103	-65 F	12	10	ര	8	7	6	ო	ŝ	9	9	10	4	8	ŝ	33	10	20	18	15	14	10	10	16	16
	CVN, FI	+70 F	13	ĸIJ	8	9	8	8	ŝ	7	6	۲.	11	89	11	10	10	14	20	20	18	20	12	ม	16	18
Temper	R.A.	z	32	25	27	19	23	22	1	15	23	19	33	20	 67	25	26	44	45	¢4 ∹ł	46	37	32	40	40	9C
400 2	El.	2	10	ፍ	9	7	6	æ	1	2	σ	6	17	2	12	<u>о</u> ,	c,	13	12	13	12	12	11	12	12	11
	U.T.S.	ksi	308	303	306	294	313	309	310	303	304	308	313	299	010	298	296	288	287	283	287	311	323	293	294	290
	Υ.S.	k#t	243	259	257	251	245	260	250	245	243	243	248	252	254	256	254	234	231	222	232	252	261	239	241	243
	Heat	No.	295	296	762	86Z	662	2100	2301	2102	2105	2106	L01 2	Z108	Z109	Z110	2111	2112	2113	2114	2115	2116	112	2116	2119	2120

Average longitudinal properties

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TABLE VII (Continued)

MECHANICAL PROPERTIES OF MEDIUM CARBON N1-Cr-Mo-W-V MARTENSITIC STEELS

		CVN, ft-lbs +70 F ~65 F		13 -		6	: 00	•	۲ ۲0	10		2	י ת	؛ ح	ā	t • •	l4 -	7			10	61				
	er	K.A.		36	34	27	25] ~	4 C	32	53		ר ד	26	20		ŝ	22	27			31	÷			
1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1		7		11	11	თ	6	~ ~	7 0	ۍ	80	0 0	n i	10	10	5	77	σ	12	1 0	207	10	2			
C f	2	k81		240	240	246	246	255) c } c	りつざ	260	262	4 •	192	270	21.7	51	256	259	110	544	239	;			
	0 1	ks1	010	017	777	221	222	213		017	217	218		077	221	212		077	219	217	4 4 1	227				
	ft-1hs	-65 ₽	1	I	6	1	1	4	1	ſ	t	t		8	1	ŧ		1	ſ	4		1				
	CVN	+70 F	14	 	ų c	0	ω	10		4 0	ת	ω	01	2 4	ע	11	0		12	11	t c	77				
'enper	R.A.	7	35	26	0 (4 (3	25	28	15) L (+	7	19	22		10	30	10	4 8	62	24		47				
800 F J	ਸ਼1.	2	10	đ	. 0	n (ַת	20	σ	. a	0	ω	6	· c	n '	10	07	ç d	Ø	2	a	c				
	IT.S.	1:81	260	257	255		107	272	272	276		270	267	6		704	262	090	202	262	245				ł	
	Y.S.	kai	227	239	232	020	077	525	234	234		230	230	233		107	227	000	6-0C	235	232	1				
	Heat	No.	295	296	297	202		662	Z100	Z101		2012	2105	2,106		1017	Z108	2100		0172	2111					

Average longitudinal properties

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TABLE VIII

MECHANICAL PROPERTIES OF NI-Cr-Mo-H-V MARTENSITIC STEELS

	Tempering					CV	N
Heat	Temperature	Y.S.	U.T.S.	E1.	R.A.	ft-	1bs
No.	o _F	ksi	ksi	<u>%</u>	7	<u>+70 F</u>	<u>-65 F</u>
2386	500	270	317	11	38	18	-
	550	269	313	10	38	18	-
	600	268	308	12	42	16	15
	650	269	304	11	44	16	
	700	268	299	11	41	16	.2
Z387	500	272	317	11	41	18	-
	550	270	312	10	40	17	-
	600	269	312	13	43	17	13
	650	271	309	11	41	17	-
	700	269	303	12	38	16	13
Z388	500	268	319	11	37	16	-
	550	268	315	10	31	15	-
	600	271	313	10	39	16	12
	650	272	310	11	35	15	-
	700	271	304	11	35	14	12

Average, longitudinal, properties

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TABLE IX

MECHANICAL PROPERTIES OF MEDIUM CARBON Ni-Cr-Mc-Si-V MARTENSITIC STEELS

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	Tempering								
Heat	Temperature	Y.S.	U.T.S.	E1.	R.A.	Charpy	Impact	Energy,	ft-lbs
No.	F	<u>ksi</u>	<u>ksí</u>	7.	7.	<u>+70 F</u>	<u>0 F</u>	<u>-65 F</u>	-200 F
							_		
Z43	400	243	298	11	46	20	20	20	14
	500	248	291	12	50	19	-	-	-
	600	243	284	9	45	17	16	1.2	7
	700	228	263	12	50	14	-	-	**
	800	207	232	11	44	22	-	-	-
Z44	400	24^	305	11	44	22	21	18	16
	500	254	299	12	47	21	-	-	-
	600	258	298	10	44	20	19	16	10
	700	247	278	13	53	16	-	-	-
	800	216	245	12	50	15	-	-	-
245	400	260	317	11	<u></u> 44	22	20	17	10
	500	263	31 0	1.1	47	19	-	-	-
	600	263	307	11	45	21	19	17	9
	700	264	298	11	49	17	-	-	-
	800	232	268	13	50	13	-	-	-
Z46	400	258	311	11	42	20	20	19	11
	500	263	306	11	45	19	-	-	-
	600	261	299	10	9 8	16	16	13	9
	700	257	285	12	51	15	œ	-	-
	800	225	250	12	47	14	-		-
247	400	256	307	10	40	21	20	16	10
	509	261	306	10	42	19	-	-	-
	600	266	306	11	41	19	16	13	10
	700	254	278	11	51	15	-	-	-
	800	221	242	11	45	14	-	-	-
Z48	400	246	313	10	40	20	20	16	7
	500	255	307	11	42	18	-	-	-
	600	256	305	б	16	16	13	12	7
	700	251	291	11	45	12	-	-	-
	800	220	266	12	40	10	-	-	-
Z49	400	256	311	13	42	21	20	14	13
	500	263	301	11	41	20	-	-	-
	500	266	301	11	46	18	16	16	12
	700	251	285	11	48	16	-	-	-
	800	219	251	10	37	14	-	-	-
250	400	235	293	11	45	20	18	18	13
	500	242	288	11	43	19	-	-	-
	600	256	297	11	41	14	15	13	11
	700	-	289	12	47	15	-	-	-
	800	222	261	12	40	12	-	-	-

TABLE IX (Continued)

MECHANICAL PROPERTIES OF MEDIUM CARBON NI-Cr-Mo-Si-V MARTENSITIC STEELS

	Tempering								
Heat	Temperature	Y.S.	U.T.S.	E1.	R.A.	Charpy	Impact	Energy,	ft-lbs
No.	° _F	ksi	ksi	7.	74	<u>+70 Z</u>	OF	<u>-63 P</u>	<u>-200 F</u>
751	400	250	25.1	12	60	12		_	
47 î	400	250	201	10	20	12	37	16	0
	500	231	205	10	20	16	17	10	2 9
	200	270	300	11	20	13	249	7.4	9
	700	233	200	11	22	11	-	-	**
	800 (00	220	213	12	54	20	15	26	10
255	400	230	203	13	20 70	45	10	10	10
	500	222	204	12	40	10	16	12	
	500	218	233	12	40	10	10	10	11
	700	209	233	12	53	13	-	-	-
	800	199	218	13	21	23	*0		10
250	400	240	305	10	37	18	10	15	10
	500	245	298	11	39	1/	-	-	-
	600	246	295	10	36	16	14	12	ວ
	700	243	186	11	39	11	•	~	124
	800	210	251	11	39	12	-	-	
257	400	256	316	10	35	18	17	B	4
	500	255	310	11	39	17	**		
	500	261	307	10	41	ló	15	13	ð
	700	259	299	11	41	14	-	**	**
	800	226	272	11	38	8		-	-
258	400	235	286	13	50	22	21	21	16
	500	224	264	13	53	18	-	-	
	600	220	253	12	51	1\$	16	16	12
	700	209	233	13	56	19	-	-	(S
	800	203	221	13	53	23	-	-	*
259	400	248	301	11	43	21	20	15	11
	500	244	296	11	39	20	-	**	4%
	600	256	294	11	43	19	15	16	10
	700	245	<u>281</u>	11	45	17	-	-	**
	800	216	252	11	41	14	**		*
Z60	400	256	312	11	40	20	17	15	. B
	500	255	308	10	39	16	34	مقد	-
	600	263	305	10	42	18	17	13	9
	700	261	300	11	40	17	*		
	800	234	275	13	46	11	**	n .	-

Average, longitudinal properties

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TABLE X

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MECHANICAL PROPERTIES OF Ni-Cr-Mo-Si-V MARTENSITIC STEELS (Ni, Mn, V, Cb SERIES)

					CV	'N	
Heat	Y.S.	U.T.S.	E1.	R.A.	ft-	lbs	
No.	ksi	<u>ksi</u>	- 74		<u>+70 F</u>	-65 F	Remarks
Z332	265	310	10	31	11.5	9,5	Columbium Series
	268	312	11	33	13	9,5	
2333	267	311	11	38	15.5	11	
	268	310	10	38	16	12	
Z334	264	306	12	35	17	14.5	
	265	310	11	38	17	13.5	¥
Z273	273	317	11	-0	16	12	Nickel Series
	271	319	11	37	14	13	l e
2274	282	324	11	40	14	13	
	281	325	11	40	14	13	
2275	279	324	11	40	16	14	
	279	324	12	43	15	13	
7329	264	311	8	19	13	10.5	
	265	310	8	20	12.5	10.5	
2330	363	306	10	31	15	12.5	
	261	306	11	3.	14.5	11	
23.71	262	303	12	3¢	18.5	15	
	263	305	11	÷;0	18.5	14.5	¥
2276	264	312	8	2)	13	11	Vanadium Series
	264	312	9	23	14	11	ł
2277	268	310	11	43	16	15	
	~ ′	4.8	-		20	17	
2278	26 4	306	11	44	20	16	
	267	307	12	45	20	17	¥
2270	271	313	10	39	19	15	Manganese Series
	271	314	12	44	19	14	ł
Z271	269	315	10	36	16	13	
	267	315	10	34	15	12	, Yanta ku
2272	266	315	11	33	15	12	Į.
	268	315	10	33	16	10	÷.

Longitudiosi properties, 600 7 tempering temperature
TAELE XI

MECHANICAL PROPERTIES OF STATISTICALLY DEGIGNED NI-CT-MO-SI-V MARTENSITIC STEELS

Heat	Y.S.	J.T.S.	E1.	R.A.	CVN.	ft-lba	TIC.	kei√ia.	^X ISCC
No.	ksi	ksi	%	7	+70 F	-65 F	+70 F	-65 F	ksi /in.
R 1	261	299	11	47	12.5	18	59.8	45.4	18
	256	296	11	4 4	22	21	62.7	46,2	
					20.5	19	64,3		
R 2	266	307	11	39	16,5	15.5	55.5	42.2	17
	261	303	11	40	15.5	16	60.C	39.0	
					14.5	17	58,9		
R 3	263	309	11	41	18	20.5	61,4	40 3	16
	261	306	11	39	19	19.5	60,0	40.7	
					20	17.5	62,4		
R 🌲	269	316	9	31	15.5	17	49.6	36.3	16
	268	312	8	25	17	14.5	48.2	37.8	
					17	15.5	49,9		
R 5	269	311	10	36	17	17.5	58.8	42.6	18
	261	301	11	38	20.5	15	59.3	40,7	
					19.5	17.5	58.7	-	
R 6	271	315	11	36	15	16	49.8	35.5	18
	267	311	11	35	18	15	50.7	35.3	
					16	15	50.5		
R 7	266	315	8	28	16	16.5	58.9	36.0	18
	264	313	10	40	17	13	57.6	38.1	
					19	14.5	5	3	
R 8	272	321	12	37	17	17		36.0	17
	274	321	9	26	18	15,5	49.7	32.2	
					17	14	52.0		
9	273	314	10	29	15	13,5	48.0	35.0	16
	270	312	8	23	16.5	15.5	47.5	35_O	
					14.5	12	48.0		_
10	284	333	8	34	16	12	40.5	31,4	18
	286	332	9	31	1:5	11	68.4	29.6	
					b.5	9	40.1		
11	265	311	11	40	16	16.5	54.7	37.5	18
	257	369	9	25	14	17.5	54.2	40.2	
					16.5	15	57.8		
12	263	300	10	42	22	21	55.0	46.3	13
	261	299	10	37	22.5	20.5	64.4	48.2	
					19.5	19	65.0		
13	264	307	11	40	21	17	\$2.0	\$4.8	18
	252	307	11	42	19	18	61.4	45.1	
		-		• ·	20	15	60.9		
14	273	316	11	35	17	14.5	47.5	34.1	19
- •	268	312	10	35	18	Ģ	46.7	36.8	
	-				18.5	10	46.1		
15	269	310	10	36	19.5	16.5	56.0	40.7	17
	261	306	10	34	19	16.5	56.6	39.2	
				÷-1		18 4	52 8		

Longitudinal properties, 600 F temperature

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TABLE XII

MECHANICAL PROPERTIES OF NI-Cr-Mo-SI-V MARTENSITIC STEELS AND 300 M STEEL R_{IC},ksi√in. Heat Y.S. U.T.S. CVN, ft-1bs El. R.A. <u>+70 F</u> ¥9. 7 <u>k61</u> kai 7. +70 F -6. 7 -65 F 2525 269 9.5 315 40 14.3 13.8 49.5 34.4 $\frac{42}{41}$ $\frac{371}{270}$ <u>316</u> 319 9.5 $\frac{14.2}{14.3}$ <u>9.5</u> 19.7 $\frac{49.1}{49.4}$ $\frac{35.6}{35.2}$ àvg. 2551 11 43 19 50,5 267 310 16 45.6 $\frac{312}{311}$ $\frac{12}{11.5}$ <u>44</u> 43.5 20 19.5 <u>59.6</u> 60.0 269 17 14.5 16.5

42

<u>43</u> <u>42.5</u>

18

<u>20</u> 19

15

 $\frac{17}{16}$

45.0

48.2

 $\frac{43.0}{45.6}$

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Longitudinal properties, 600 P tempering temperature

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<u>287</u> 286

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Avg.

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(300H)

268

240

<u>245</u> 243

TABLE XIII

COMPARISON OF EXPERIMENTAL AND PREDICTED MECHANICAL PROPERTIES FOR HEATS Z525 AND Z551

	E	xperiment	tal Resul	its	P	redicted	Results	
Heat No.	U.T.S. ksi	K 1C +70 F	KIC -65 F	CVN ft-lbs	U.T.S. <u>ksi</u>	^K IC +70 F	K -65 F	GTN ft-1bs
Z525	318	49	35	<u>1</u> 4	308	65	42	18
Z551	311	69	45	19	292	67	51	22

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TABLE XIV

TENGILE, CHARFY, FRACTURE TOUCHNESS, AND STRASS COLROSION PROPERTIES OF 9-4-45 STEEL WITH PHOSPHORUS AND SULFUE ADDITIONS

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N 14	003	•			000	. 040				czn.				.025					600.				.010			
A 21	.003	•			613					. 024				.004				000	. 120				.004			
K _{T.SCC} ks1 /1n.	14	3	16		11	t <u>1</u>		, i	-	मच	1 C	51	5	12	t	18) t		ŗ	t v 7	Q T	1	11	ŧ	17	1
k _{IC} ksi /in.	61.1	2	91.1	1	42.3	3	64.8	ŧ	0 UV		2 Y Y		2 4 -	43.0	ŧ	53.0	8	0 75		11 1 1 1	C.1/	1	55.0	ł	77.0	3
CVN ft-1bs	15	14	25	22	12	11	17	12	8	v	14	t	 	7 T	51	13	11	14	1	1.0		07	74	12	22	20
R.A. 7	41	40	5 G	43	34	33	43	36	ო	~	36	200	1 C	4 4	17	38	27	44	41	5	 	, ; ;	1	3/	50	17
121. - 22	10	No.	ន	12	æ	9	11	11	ر ي	2	5	. 63) r	~ ~		10	σ'n	10	0	12	11	• •	2	ת	13	11
U. T. S. ks1	276	1/2	267	266	280	280	269	268	283	285	269	266	286	100		707	270	277	277	269	268	776	282	202	267	2.68
Y.S.Y.	239	2010	270	97 2'	140 140	245	210	210	243	243	215	216	249	240		01 4 01 4	Z 15	241	2430	215	214	246	263		212	213
Teat Direction	⊷i E	⊣ ,	-1 E	₩,	-1 8	; 4 ,	-7 1	Et ,	~	H	ក	بع	1	è	ہ ا	3 E	H .	н а.	64	ר '	ы	1	1 5-	* ,	-3 6	H
Miszos tructure	Mar tens i.te v	Reinito		Martoneitz			174111CC			c	Buinite	36	Martemette	47 H	Baintee	14		sant terret. Co	:	Baintte	13	Mertensice	2	Baintre		
Heat No.	V723			4774				1796	(TA 1 A			1	V7.26				1-1-1-1-13	A 1 410				17 445				

Average roum temperature properties

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TABLE XV

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MECHANICAL PROPERTY COMPARISON OF VIM, ESR, AND VAR EXPERIMENTAL BAINITIC AND MARTENSITIC STEELS

Heat No.	Y.S. kei	U.T.S. ksi	B1.	R.A. 7	CUN.	ft-1b <i>u</i> -65 F	K _{IC} , ks +70 F	1 /in. -65 F	Kemarku
2409	270	312	10	41	15	14	55.4	39.0	VIM - Low Alloy Martensite
G229	287	332	7.5	25	11	8	34.3	27.7	ESR - Low Alloy Martensite
3088808	263	309	S	34	17	15	56,2	42.0	VAR - Low Alloy Martensite
1142	246	299	10	39	13	11	62.0	47.0	VIM - Medium Alloy Bainite
C230	243	290	12	45	16	15	58.8	41.5	ESR - Medium Alloy Balnice
388800	235	290	13	48	19	12	67.2	49.8	VAR - Medium Ailoy Jainite
Z422	260	318	10	33	14	8	34.5	31,9	VIM - Low Alloy Leinite
C231	264	321	10	33	12	7	29.8	23.8	ESR - Low Alloy Buintte

Average, longitudinal properties

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TABLE XVI

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VOLUME PERCENT INCLUSION DATA ON VIM, ESR, AND VAR MEDIUM ALLOY BAINITIC STEELS

Volume % Inclusions

Cyle Threaded Area Threaded Area Fracture Area [1] 1 0.34 0.040 20.411 20.1						
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	শ	Sample	Thre AMEDA	aded Area Point Counting	Fractur AMEDA	e Area Point Counting
3 0.47 - - 0.28 - 0.04 - - 0.03 - - <td></td> <td></td> <td>.034</td> <td>.040</td> <td>ŧ</td> <td>£</td>			.034	.040	ŧ	£
6 0056 - 0056 - 0038 -		ო	.047	a	i	1
8 .038 .038 .034 13 .034 .03 .034 Average .042 .044 .072 14 . .072 .072 15 .070 . .072 14 . .070 . 14 . .070 . 14 . . .070 14 . . .070 14 . . .070 15 . . . 64 . .070 . 14 . . .070 15 . . .063 11 .063 . . 11 .063 . . 11 .063 . . 12 . . . 13 . . . 10 . . . 11 . . . 12 <t< td=""><td></td><td>¢,</td><td>, 056</td><td>t</td><td>. 028</td><td>3</td></t<>		¢,	, 056	t	. 028	3
13 .034 - <td></td> <td>8</td> <td>.038</td> <td>t</td> <td>.054</td> <td>3</td>		8	.038	t	.054	3
Average		13	.034	1	3	J
10 -		Average	.042			
14 -		10	ł	·	.072	.058
4A .070 . .070 . .070 . </td <td></td> <td>14</td> <td>ł</td> <td>1</td> <td>. 044</td> <td>1 3 9</td>		14	ł	1	. 044	1 3 9
44 48 58 58 58 58 58 58 58 58 58 5					Average .050	
4B .047 - 5A .043 - 5A .063 - 5B .063 - 11 .063 - 11 .063 - 5B .064 - 11 .063 - 11 .063 - 11 .063 - 12 - .055 10 - - 10 - - 10 - - 10 - - 10 .056 - 10 .056 - 10 .056 - .050 .005 - .050 .005 - .050 .050 -		4A	.070	£	ł	1
5A .048 - <td></td> <td>4B</td> <td>.047</td> <td>1</td> <td>t</td> <td>t</td>		4B	.047	1	t	t
5B .063 . 11 .063 . 11 .063 . 11 .052 . 11 .055 . 11 .055 . 11 .055 . 11 .055 . 12 . .056 10 . . 10 . . 10 . . 10 . . 10 . . 10 . . 10 . . 10 . . 10 . . 10 . . 10 . . 10 		5.6	.048	ł	\$	ı
11		58	.063	ı	•	1
Average .055 Average .055 12009 10009 10009 Average .048 001 		and Fri	.047	Ł	.052	ı
12 13 14 10 10 10 10 10 10 10 10 10 10		Averege	.055			
10 10 10 10 10 10 10 10 10 10		12.	£	660.	3	,
10		٢Ĵ	ŧ	9	.061	. 107
800 4 Average .054 Average .054 .054 .054 .054 .054 .054 .013 .013013015015050		10	ŧ	ł	. 048	
800 4 2 - 013 - 013 - 015 - 015 - 015 - 015 - 015 - 015 - 00					Average .054	
7	800	4	ŧ	1	.013	1
% .018 - .050 .050 10 .004 .020 .020 .050 .050		*	ł	Ł	.015	8
10 .00% .020 .020 .011		¢,	010	ı	. 005	, ņ50
Average .011			. 004	. 620	a 	\$
	•				Average ,011	

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TABLE XVII

COMPARISON OF FATIGUE PROPERTIES OF EXPRETIZENTIAL STEELS WITH COMMERCIAL HIGH STRENGTH STREELS

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Reference	15 29	30 25 25	31	<u>ដ</u> រ រ	31 31 31	31 32	t SIHI,		2 2 2
Mícro- structure	Martensíte "	Martensite 11 11	Martenstte	Martensite " "	2 2 2	Bainite Martenofie H	Martousles	: 3	Balaten Bolaten Mortensste
Fatigue Ratio (10 ⁷ Strength/ U.T.S.)	. 39 . 41	.31 .39 .41	67 "	.38 .41 .12	. 37 . 40 . 35	. 50 . 50 . 49	£2 .	, 45 , 44	74. 14.
10 ⁷ Cycle Fatigue Strength kai	105 110	~90 118 120	130	118 118 118	58 58 68	140 134 130	170	140 138	-125 160 175
U, T, S. kei	~270 266	294 ~285 290	265	307 306 288 288 288 288 288	283 283	270 268 268	321	312 312	299 317 286
αi	+0.06 "	+0.10	+0,10	90°09 •••••	0,10	+0.10 +0.10	+0.10	= = :	7 2 E
Test Dir.	: !	۽ ۽ آ <i>س</i>	н	너 더 너 더	4 4 4	54 x Fr	ન 1	臣曰:	24 55 24 55
Produc t Form	Billet "	Porging 	Billet	Vorging " "	2 2	Plate - Forging	1/2" Plate	: = :	= = =
Melting Practice	VAR 19	VAR 11 11	лак	VIM- VAR VIM- VAR VAR VAR	VAR	VAR Var Var	WIN	: = :	: = =
Alloy	4340 "	300 24	8-11	9 18 Ni Maraging N (300 Grade) "	2 2	HP 9-4-45 11	1351	2409	Z412 Z412 Z469 (300 M)

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TABLE XVII (Continued)

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COMPARISON OF FATIGUE PROPERTIES OF EXPERIMENTAL STEELS WITH COMMERCIAL HIGE STRENGCH STREAS

<u> </u>	Melting. Practice	Product Form	Test Dir.	শ্বা	U.T.S. kei	10 ⁷ Cycle Fatigue Surwngth koi	Fațigue Rutio (10' Strength/ U.T.S.)	Micro- structure	Refarence
00%28%86	VAR	1/21	L	+0.10	290	195	. 67	Bainte	THIS Investigation
3688931	11	51 59 F	*	•	307	061	, 62	Martensite	Ξ
C229	RSR	1/2 ^w 18445	ц	+0.10	332	170	12,	Martensite	THIS Investigation
0823		; ; ; ; ; ; ; ; ; ; ; ; ; ; ; ; ; ; ;	E ci :	= = =	332 290 321	1.70 207 210	.51 .71 .66	יי Bainitw Bainite	= = =
	41 14 17 17 14 14 14 14 14 14 14 14 14 14 14 14 14								

8 is the ratio of minimum stress to maximum stress

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TABLE XVIII

GRAIN SIZE AND HARDNESS OF THERMAL-MECHANICALLY PROCESSED 300M STEEL

		;	i	1		Austeni	tizing	Tempera	ture*				
PT.	00.086	6.S.	세	<u>6.S.</u>	^{ਸ਼}	1525 G.S.	<u>т</u>	1550 G.S.	~ ~	1575 G.S.	^н _М	1600 G.S.	ч Ж
a	Spheroidsl Anneal Rolled at 1200 ?	13.5	55.0	12.2	53.5	1.1.8	58.5	11.2	37.0	10.7	58.0	10.2	58.5
þ.	Spheroidal Anneal Roiled at Room Temp.	14.5	56,5	13.7	55.0	12.6	58.0	11.5	56.0	11.5	58.0	11.0	56,5
ຍ່ .	Spheroidal Aineal Cowled to Rown Temp. Rollad at 1200 F	14.0	57.0	13.2	57.5	12.4	58.7	11.7	56.0	11.1	58.0	10.4	57.5
g	Quenched and Tempered Rolled at 1200 F	12.8	55.5	12.2	57.3	11.8	57.5	11.0	57.5	10.7	55.5	10.1	57.0
	Quenched and Tempered Rolled at Room Temp.	12.9	57.3	12.1	58,5	11.5	58.5	11.2	54.5	11.0	58.ù	10,4	56.0
પં	Quenched and Tempered Cooled to Room Temp. Rolled at 800 F	12.3	57.0	11.9	58.3	10.7	58.0	10.6	57.0	10.5	57.3	10.4	56.0

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G.S. = ASTM-Number *Austenitized for 1/2 hour and oil quenched

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Sample No.	Process	Aust.Temp.	ASTM <u>G.S.</u>	Y.S. <u>(ksi)</u>	U.T.S. (ksi)	R.A. (7)	E1. (1")	CVN (ft-1bs)
1-1 1-2 Avg. 1-3 1-4 Avz.	Sph. Anneal Roll at 1200 F	1475 f 1525 f	13.5 11.8	261.0 259.0 260.0 253.0 253.0 253.0	293.2 289.2 291.2 288.2 288.2 288.2 288.2	29.3 30.5 29.9 29.9 36.3 33.1	9.0 9.0 9.0 10.0 16.0 10.0	14 14 17 17 17
2-1 2-2 Avg. 2-3 2-4 Avg.	Sph. Anneal Roll at R.T.	1475 F 1525 F	14.5 12.6	258.0 260.5 259.2 253.0 251.5 252.2	289.2 290.2 289.7 288.2 288.2 288.2 288.2	36.9 32.5 34.7 39.6 38.4 39.0	9.0 <u>10.0</u> 9.5 10.0 <u>10.0</u> 10.0	15 <u>17</u> 16 18 <u>16</u> 17
3-1 3-2 Avg. 3-3 3-4 Avg.	Sph. Anneal Cool to R.T. Roll at 1200 F	1475 F 1525 F	14.0 12.4	267.1 267.1 267.1 257.0 256.5 256.7	293.2 293.2 293.2 291.2 291.2 291.2 291.2	32.5 31.1 31.8 31.1 31.9 31.5	9.0 9.0 9.0 10.0 10.0 10.0	15 <u>16</u> 15.5 19 <u>19</u> <u>19</u>
4-1 4-2 Avg. 4-9 4-4 Avg.	Tempered Mart. Roll at 1200 F	1475 F 1525 F	12.8 11.8	253.5 252.0 252.7 248.0 247.0 247.5	289.2 290.2 269.7 289.2 289.2 288.2 288.2	35.7 33.7 34.7 36.9 <u>39.6</u> 38.2	$ \begin{array}{r} 12.0 \\ \underline{11.0} \\ \overline{11.5} \\ 11.0 \\ \underline{12.0} \\ \overline{11.5} \end{array} $	19 19 19 20 22 21
5-1 5-2 Avg. 5-3 5-4 Avg.	Tempered Mart. Roll at R.T.	1475 F 1525 F	12.9 11.6	249.0 251.0 250.0 248.0 249.5 248.7	291.0 291.2 291.1 290.2 290.2 290.2 290.2	39.0 40.2 39.6 41.4 40.2 40.6	11.0 <u>11.0</u> 11.0 12.0 12.0 12.0	20 20 20 20 20 20 20
6-1 6-2 Avg. 6-3 6-4 Avg.	Tempered Mart. Roll at 800 F	1475 F 1525 F	12.3 10.7	$\begin{array}{r} 253.0\\ \underline{252.0}\\ 252.5\\ 251.0\\ \underline{251.0}\\ 251.0\\ \underline{251.0}\\ 251.0\end{array}$	293.2 293.2 293.2 291.2 291.0 291.1	40.8 35.7 38.7 38.4 40.2 39.3	11.9 <u>12.0</u> <u>11.5</u> 11.0 <u>11.9</u> 11.0	19 20 19.5 20 22 21
Avg.	Conventional Processing and Heat Treatment	1600 F	9.4	241.6	282.3	à 4 ,7	12.0	20

TABLE XIX

MECHANICAL PROPERTIES OF THERMAL-MECHANICALLY PROCESSED 300M STEEL

All material tempered at 660 F for 2 + 2 hours

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TABLE XX

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54. TH

GIAIN SIZE AND HARDNESS OF THERMAL-MECHANICALLY FROCESSED LOW ALLOY MARTENSITIC STEEL (2350)

ſ	Cabarodden Annual	G.S.	S E E R C S C	<u> </u>	C R C	1625 G.S.			10.50 G.S.	1650 F G.S. Rc	<u>1650 F</u> <u>G.S.</u> Rc <u>G.S.</u> 10 f f f 10 f 10 f	<u>1650 F</u> <u>G.S. R</u> c <u>G.S. R</u> c 10 <u>6</u> <u>6</u> <u>7</u> <u>6</u> <u>7</u> <u>6</u> <u>7</u> <u>6</u> <u>7</u> <u>6</u> <u>7</u>	<u>1650 F</u> <u>1675 F</u> <u>17</u> <u>G.S. R</u> <u>G.S. R</u> <u>G.S.</u> 10 <u>5</u> <u>5</u> <u>10</u> <u>5</u> <u>6</u> <u>5</u> <u>7</u>
6	spherolulize Anneal, Rolled at 1100 F	0.21	1.10	C.11	1.00	11.0		1.00	9.01 1.00	0.66 0.01 /.66	0.01 0.66 0.01 /.66	0.02 0.01 0.62 0.01 1.62	9.6 0.95 0.01 0.CC 9.01 1.cc
.	Spheroidize Anneal, Cooled to Room Temp, Rolled at 1100 F	11.8	55.7	10.6	56.0	6'6	-,	55.0	55.0 9.7	55.0 9.7 55.7	55.0 9.7 55.7 9.3	55.0 9.7 55.7 9.3 53.0	55.0 9.7 55.7 9.3 53.0 9.0
ບ້	Quenched and Tempered, Covled to Room Temp, Rolled at 1200 F	11.0	57.0	10.3	55.7	10, 1	56	0.0	6.0 9.6	i.0 9.6 54.7	i.0 9.6 54.7 9.9	i.0 9.6 54.7 9.9 55.5	i.0 9.6 54.7 9.9 55.5 9.7
d.	Quenched and Tempered, Cooled to Room Temp, Rolled at 800 F	12.0	57.0	11.5	55.0	10.8	55	0	.0 10.1	.0 10.1 55.7	.0 10.1 55.7 10.0	.0 10.1 55.7 10.0 56.0	.0 10.1 55.7 10.0 56.0 9.1

G.S. m ASTM number *Austenitized for 1 hour and oil quenched

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TABLE XXI

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MECHANICAL PROPERTIES OF THERMAL-MECHANICALLY PROCEEDED LOW ALLOY MAKTENSITIC STEEL (2350)

Sanple Not	From	Aust. Temp	AS TH G. S.	Y.S. kai	U.T.S. kei	B.A. 7	E1. (1") Z	CVN (ft-1bs)
1-1 1-2 Åvs.	Sph. Anneel Roll st 1100 F	1600 F	11.5	288.2 291.2 289.7	316.2 321.3 318.7	35, 1 36, 9 36, 0	9.0 10.1	17.0 19.0 18.0
3* 7 3 - 7 & V 8.	Sph. Annewl. Coul to R.T. Roll at 1100 F	1600 F	10.6	284,1 282.1 283.1	216.2 317.2 316.7	33.7 24.5 34.1	0 0 0 0 0 0	16.5 17.0 16.7
4- 1 4- 2 A vg.	Tempered Mart. Cool to R.T. Roll at 1200 F	1600 P	10.3	276.1 278.9 277.5	315.2 316.7 316.9	39.0 35.7 37.3	11.0 9.0 10.6	16.0 17.5 16.7
6 l 6 2 Åv8.	Tempered Mart. Cool to R.T. Roll at 8.00 F	1600 F	11.5	280.1 281.4 280.7	316.2 318.7 317.7	40.2 41.4 40.8	11.0 9.0 10.0	16.5 17.0 16.7
350-1 350-2 Åvg.	Conventional Processing and Neat Treatment	1700 <i>F</i>	۶°,	278.1 279.1 278.6	305.6 305.8 305.7	29.3 26.5 27.9	0°6	14.0 13.0

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All material tempered at 600 F for 2 + 2 hours, longitudinch direction

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TABLE XXII

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MECHANICAL ENDERTIES OF A THERMAL-MECHANICALLY PROCESSED IOW ALLOY BAINITIC STEEL (2412)

Semple No.	Frocass	éust, Temp#	Asim <u>G.S.</u>	Y.S. <u>kzi</u>	U.T.S. <u>Àsi</u>	R.A. <u>7</u>	E1.(1") 7	CVN ft-1bs
ì-1 1-2 Avg.	Zph. Annael Rolled at 1200 P	1600 P	11.5	$\frac{242.0}{241.0}$ 241.5	304.2 <u>303.2</u> 303.7	33.1 <u>28.5</u> 30.8	10.0 <u>10.0</u> 13.0	11.8 <u>12.0</u> 11.9
1-3 1-4 Avg.		1630 P	11.0	225.9 236.9 231.4	300.2 <u>304.2</u> 302.2	30.5 <u>31.1</u> 30.8	11.0 11.0 11.0	$ \begin{array}{r} 14.2 \\ \underline{11.5} \\ 12.8 \end{array} $
2-1 2-2 Av8.	Sph. Annes1 Cocl to R.T. Rolled st 1200 F	1800 P	12.4	251,0 <u>252.0</u> 251,5	306.2 <u>306.2</u> 306.2	35.1 <u>35.1</u>	11.0 <u>0</u> 10.0	$\frac{12.5}{14.0}$ 13.2
Z-3 Z-4 А¥8.		1650 F	11.7	230,9 <u>228,9</u> 229,9	301.2 301.2 301.2	36.9 <u>31.1</u> 34.0	9.0 <u>11.0</u> 19.0	15.2 <u>14.5</u> 14.8
3-1 3-2 Avg.	Quenched and Tempered Rolled at 1200 Y	16ò0 F	11.9	248.5 245.0 245.7	306.2 306.2 308.2	32.9 36.3 34.4	11.0 <u>11.0</u> 11.0	14.2 10.5 12.4
3-3 3-4 Avg.		1650 F	11,1	232.5 <u>229.9</u> 231.2	303.2 303.2 303.2	39.0 39.6 39.3	12.9 <u>12.0</u> 12.0	15.0 15.0 15.0
412-1 412-2 -	Conventionsl Processing and Hest Treatment	1650 F	10 6	228,9 235.9	305.2 305.2	38.4 37.6	13.0 12.0	15.0 14.2 13.2
Avs.				232.4	305,2	38.0	12.9	14.1

*Austenitized at temperature for one hour and isothermally transformed at 475 F for alx hours



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Figure 4. Fetigue Testing Apparatus



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320 \Box Strength, kei 008 U.T.S. 280 Y.S. 260 .20 .40 .60 .80 0 25 20 ≁70 F CVW, Et-lbs 15, റ 10 -65 P 5 0 .20 .80 .40 .69 0 Tuogsten, X

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С 30 0 Effect of Silicon on Charpy Impact-Transition Curves for Heats 243, 244, and 245 (400 F Temper) 400 F Temper ▲ Z44 (1.70% S1), 305 ksi
 ■ Z45 (2.65% S1), 317 ksi O Z43 (1.05% S1), 298 kai Test Temperature, ^{OF} - 65 1.70% St 2.657 31 1.05% St Figure 15. - 200 20 25 12 2 ŝ 0 Charpy Impact Energy, it-1bs

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350 0 Z58 (0.33% Si) ۵ Z59 (1.65% Si) Z60 (2.65% Si) Δ U.T.S. Δ 300 Strength, ksi ſ 250 O Y.S. 200 25 Tested at +70F Charpy Impact Energy, ft-lbs C 20 15 258 (0.33% Si) 259 (1.65% Si) 0 ۵ 260 (2.65% Si) Δ 10 7.00 400 500 600 800 Tempering Temperature, OF

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320 310 300 U.T.S. 290 Strength, ksi. 60 280 ¥.,S. 270 50 5 E 260 40 å 7 R.A. R 250 30 240 20 .20 .30 9 . 10 .40 30 25 +70 F 20 CVN, ft-lbs 15 X ~55 F 0 10 5 0 ÷ .10 .20 .30 0 .40 Vanadium, %

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Figure 31. Microstructures of Vanadium Series of Ni-Ct-Mo-Si-V Martensitic Steels

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330 320 310 V.T.S. 300 Strength, ksi 290 280 Y.S. 270 260 250 .1 .2 .4 .6 0 .8 30 25 20 CVN, ft-1bs +70 F 15 10 . -65 F 5 0 .2 .6 0 .4 .8 1.0 Manganese, %

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Figure 33. Transmission Electron Micrograph of Heat Z270, Containing 0.26% Mn, Showing Short Microtwins in Some of the Martensite Flates

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Figure 34. Transmission Electron Micrograph of Heat 2272, Containing 0.78% Mn, Showing Long Microtwins in the Martensite Plates

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Tensile Strength-Fracture Toughness Relationships for Experimental Low Alloy Mertensitic Steels Compared to 300 M Steel Figure 45.

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30 25 Charpy V-Notch Impact Energy, ft-1bs 20 ST, 15 10 5 0 50 60 30 70 40 Fracture Toughness, K_{IC}, ksi√in. Comparison of Fracture Toughness and Charpy Impact Energy Values for Heats R 1 through 15 Figure 46.

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S-N Curve for Bainitic Alloy 3588800 (VAR) Figure 71.

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Comparison of Critical Crack Sizes of the New Low Alloy Martensitic Steels with Commercial High Strength Steels

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The objective of this program was	s to develop an	ultra-high st	rength steel in the		
The objective of this program was 300 to 320 ksi ultimate tensile a	s to develop an strength range,	ultra-high st with improved	rength steel in the fatigue strength,		
The objective of this program was 300 to 320 ksi ultimate tensile a fracture toughness, and stress ca	s to develop an strength range, orrosion resist	ultra-high st with improved ance for great	rength steel in the fatigue strength, ar reliability in		
The objective of this program was 300 to 320 ksi ultimate tensile a fracture toughness, and stress ca forged landing gear components.	s to develop an strength range, orrosion resist Alloy developm	ultra-high st with improved ance for great ent studies we	rength steel in the fatigue strength, er reliability in ere conducted on two order to develop the		
The objective of this program was 300 to 320 ksi ultimate tensile a fracture toughness, and stress co forged landing gear components. bainitic alloy systems and two man bast combination of mechanical man	s to develop an strength range, orrosion resist Alloy developm artensitic allo romerties at the	ultra-high st with improved ance for great ent studies we by systems in o asile strength	rength steel in the i fatigue strength, ar reliability in ere conducted on two order to develop the plevels in excess of		
The objective of this program was 300 to 320 ksi ultimate tensile a fracture toughness, and stress ca forged landing gear components. bainitic alloy systems and two ma best combination of mechanical pa 300,000 psi. Of the four allow	s to develop an strength range, orrosion resist Alloy developm artensitic allo roperties at te systems investi	ultra-high st with improved ance for great ent studies we y systems in o asile strength gated, steels	rength steel in the fatigue strength, ar reliability in ere conducted on two order to develop the levels in excess of from the low alloy.		
The objective of this program way 300 to 320 ksi ultimate tensile a fracture toughness, and stress ca forged landing gear components. bainitic alloy systems and two ma best combination of mechanical pa 300,000 psi. Of the four alloy a medium carbon Ni-Cr-Mo-Si-V marte	s to develop an strength range, orrosion resist Alloy developm artensitic allo roperties at te systems investi ensitic evstem	ultra-high st with improved ance for great ent studies we by systems in o asile strength gated, steels developed the	rength steel in the fatigue strength, ar reliability in are conducted on two order to develop the h levels in excess of from the low alloy, best combination of		
The objective of this program was 300 to 320 ksi ultimate tensile a fracture toughness, and stress ca forged landing gear components. bainitic alloy systems and two ma best combination of mechanical pa 300,000 psi. Of the four alloy a medium carbon Ni-Cr-Mo-Si-V marter fracture toughness, fatigue stream	s to develop an strength range, orrosion resist Alloy developm artensitic allo roperties at te systems investi ensitic system ngth and stress	ultra-high st with improved ance for great ent studies we by systems in o masile strength gated, steels developed the corrosion creat	rength steel in the fatigue strength, er reliability in ere conducted on two order to develop the levels in excess of from the low alloy, best combination of acking resistance.		
The objective of this program was 300 to 320 ksi ultimate tensile a fracture toughness, and stress ca forged landing gear components. bainitic alloy systems and two ma best combination of mechanical pa 300,000 psi. Of the four alloy a medium carbon Ni-Cr-Mo-Si-V marter fracture toughness, fatigue stream martensitic alloy was developed to	s to develop an strength range, orrosion resist Alloy developm artensitic ello roparties at te systems investi ensitic system ngth and stress which achieves	ultra-high st with improved ance for great ent studies we y systems in o asile strength gated, steels developed the corrosion cra the following	rength steel in the i fatigue strength, er reliability in ere conducted on two order to develop the i levels in excess of from the low alloy, best combination of ocking resistance. average longitudina		
The objective of this program was 300 to 320 ksi ultimate tensile a fracture toughness, and stress ca forged landing gear components. bainitic alloy systems and two m best combination of mechanical pa 300,000 psi. Of the four alloy a medium carbon Ni-Cr-Mo-Si-V marte fracture toughness, fatigue stren martensitic alloy was developed a room temperature properties based	s to develop an strength range, orrosion resist Alloy developm artensitic allo roperties at te systems investi ensitic eystem ngth and stress which achieves d on laboratory	ultra-high st with improved ance for great ent studies we by systems in o asile strength gated, steels developed the corrosion cra the following sized heats:	rength steel in the fatigue strength, ar reliability in ere conducted on two order to develop the h levels in excess of from the low alloy, best combination of acking resistance. average longitudina Y.S. = 268 ksi, U.		
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