Standard Specification for
Wrought 18Chromium-14Nickel-2.5Molybdenum Stainless
Steel Bar and Wire for Surgical Implants (UNS S31673)¹

1. Scope

1.1 This specification covers the chemical, mechanical, and
microstructural requirements for wrought 18 chromium-14 nickel-
2.5 molybdenum stainless steel bar and wire used for the
manufacture of surgical implants.

1.2 The values stated in either SI units or inch-pound units
are to be regarded separately as standard. The values stated in
each system may not be exact equivalents; therefore, each
system shall be used independently of the other. Combining
values from the two systems may result in non-conformance
with the standard.

2. Referenced Documents

2.1 ASTM Standards:²
A262 Practices for Detecting Susceptibility to Intergranular
Attack in Austenitic Stainless Steels
A484/A484M Specification for General Requirements for
Stainless Steel Bars, Billets, and Forgings
A555/A555M Specification for General Requirements for
Stainless Steel Wire and Wire Rods
A751 Test Methods, Practices, and Terminology for Chemical
Analysis of Steel Products
E8/E8M Test Methods for Tension Testing of Metallic Materials
E10 Test Method for Brinell Hardness of Metallic Materials
E18 Test Methods for Rockwell Hardness of Metallic Materials
E29 Practice for Using Significant Digits in Test Data to
Determine Conformance with Specifications
E45 Test Methods for Determining the Inclusion Content of Steel

² Available from SAE International (SAE), 400 Commonwealth Dr., Warrendale,

3. Terminology

3.1 Definitions of Terms Specific to This Standard:
3.1.1 bar, n—rounds, flats, or other shapes from 0.1875 in.
[4.76 mm] to 4 in. [101.60 mm] in diameter or thickness.
(Other sizes and shapes by special order.)

3.1.2 fine wire, n—wire as described in 3.1.5, less than 0.063
in. [1.60 mm] in diameter or thickness.

3.1.3 forging bar, n—bar as described in 3.1.1, used for the
production of forgings, may be furnished in the hot worked
condition.

3.1.4 lot, n—the total number of mill products produced
from the same melt heat under the same conditions at essen-
tially the same time.

3.2.2 ISO Standards:³
ISO 5832–1 Implants for Surgery—Metallic Materials—
Part 1: Wrought Stainless Steel
ISO 6892 Metallic Materials—Tensile Testing
ISO 9001 Quality Management Systems—Requirements

3.3 ASTM Standards:
F981 Practice for Assessment of Compatibility of Biomate-
rials for Surgical Implants with Respect to Effect of
Materials on Muscle and Bone
F1350 Specification for Wrought 18 Chromium-14 Nickel-
2.5 Molybdenum Stainless Steel Surgical Fixation Wire
(UNS S31673)
IEEE/ASTM SI 10 American National Standard for Metric
Practice

3.3.1 AMS Standards:
AMS 2630 Inspection, Ultrasonic Product Over 0.5 inch
(12.7 mm) Thick
AMS 2632 Ultrasonic Inspection of Thin Materials

3.4 Other Standards:
E18 Test Methods for Rockwell Hardness of Metallic Materials

3.5 Document Summary:
For Annual Book of ASTM Standards volume information
refer to the standard’s Document Summary page on
the ASTM website.

¹ This specification is under the jurisdiction of ASTM Committee F04 on
Medical and Surgical Materials and Devices and is the direct responsibility of
Subcommittee F04.12 on Metallic Materials.

approved in 1971. Last previous edition approved in 2013 as F138 – 13. DOI:
10.1520/F0138-13A.

² For referenced ASTM standards, visit the ASTM website, www.astm.org, or
contact ASTM Customer Service at service@astm.org. For Annual Book of ASTM
Standards volume information, refer to the standard’s Document Summary page on
the ASTM website.

³ Available from American National Standards Institute (ANSI), 25 W. 43rd St.,

* A Summary of Changes section appears at the end of this standard
3.1.5 wire, \( n \)—rounds, flats or other shapes less than 0.1875 in. [4.76 mm] in diameter or thickness.

4. General Requirements for Delivery

4.1 In addition to the requirements of this specification, all requirements of the current editions of Specifications A484/A484M and A555/A555M shall apply.

4.2 In the case where a conflict exists between this specification and those listed in 2.1 and 2.3, this specification shall take precedence.

5. Ordering Information

5.1 Inquiries and orders for material under this specification shall include the following information:

5.1.1 Quantity (weight or number of pieces);
5.1.2 ASTM designation and date of issue;
5.1.3 Form (bar, wire, fine wire);
5.1.4 Condition (see 6.1);
5.1.5 Mechanical properties (if applicable, for special conditions);
5.1.6 Finish (see 6.2);
5.1.7 Applicable dimensions including size, thickness, width, and length (exact, random or multiples) or drawing number;
5.1.8 Special tests, if any; and
5.1.9 Other requirements.

6. Materials and Manufacture

6.1 Condition:

6.1.1 Bar and wire shall be furnished, as specified, in the hot worked, annealed, cold worked, or extra hard condition (see Table 1).

6.1.2 Fine wire shall be furnished, as specified, in the cold drawn condition (see Table 2).

6.2 Finish:

6.2.1 Types of finish available for bar and wire products are cold drawn, pickled, ground, ground and polished, or as specified in the purchase order.

6.2.2 Types of finish available for fine wire products are descaled or pickled, abrasive-blasted, cold drawn, ground, ground and polished, or as specified in the purchase order.

7. Chemical Requirements

7.1 The heat analysis shall conform to the requirements as to chemical composition specified in Table 3.

7.1.1 The compositional requirement shall meet the following:

\[
\% \text{Cr} + 3.3 \times \% \text{Mo} \geq 26.0
\]  

(1)

7.1.2 Requirements for the major and minor elemental constituents are listed in Table 3. Also listed are important residual elements. Analysis for elements not listed in Table 3 is not required to certify compliance with this specification.

7.1.3 Methods and practices relating to chemical analysis required by this specification shall be in accordance with Test Methods, Practices, and Terminology A751.

7.2 Product Analysis—Product analysis tolerances do not broaden the specified heat analysis requirements, but cover variations between laboratories in the measurement of chemical content. The supplier shall not ship material that is outside the limits specified in Table 3. Product analysis limits shall be as specified in Table 4.

7.2.1 The product analysis is either for the purpose of verifying the composition of a heat or manufacturing lot or to determine variations in the composition within the heat.

7.2.2 Acceptance or rejection of a heat or lot of material may be made by the purchaser on the basis of this product analysis.

7.2.3 Methods and practices relating to chemical analysis required by this specification shall be in accordance with Test Methods E354.

8. Metallurgical Requirements

8.1 The material shall exhibit no delta ferrite, chi, or sigma phases when it is examined metallographically at 100x magnification when etched in accordance with Practice E407.

8.2 The microcleanliness of the steel as determined by Method A of Test Methods E45, except using Plate I-r, on representative billet or bar samples from the heat shall not exceed the following:

<table>
<thead>
<tr>
<th>Inclusion Type</th>
<th>A (Sulfide)</th>
<th>B (Alumina)</th>
<th>C (Silicate)</th>
<th>D (Globular Oxides)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Thin</td>
<td>1.5</td>
<td>1.5</td>
<td>1.5</td>
<td>1.5</td>
</tr>
<tr>
<td>Heavy</td>
<td>1.0</td>
<td>1.0</td>
<td>1.0</td>
<td>1.0</td>
</tr>
</tbody>
</table>

9. Mechanical Properties

9.1 Tensile Properties:

9.1.1 Tensile properties shall be determined in accordance with Test Methods E8/E8M.

<table>
<thead>
<tr>
<th>Condition</th>
<th>Diameter or Thickness, in. [mm]</th>
<th>Ultimate Tensile Strength, min, psi [MPa]</th>
<th>Yield Strength (0.2% offset), min, psi [MPa]</th>
<th>Elongation(^a) in 4D or 4W, min, %</th>
<th>Brinell(^b) Hardness, max, HB</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hot worked(^c)</td>
<td>all</td>
<td>(\ldots)</td>
<td>(\ldots)</td>
<td>(\ldots)</td>
<td>250</td>
</tr>
<tr>
<td>Annealed</td>
<td>0.063 and over [1.60]</td>
<td>71 000 [490]</td>
<td>27 500 [190]</td>
<td>40</td>
<td>(\ldots)</td>
</tr>
<tr>
<td>Cold worked</td>
<td>0.063 to 1.500 [1.60 to 38.1]</td>
<td>125 000 [860]</td>
<td>100 000 [690]</td>
<td>12</td>
<td>(\ldots)</td>
</tr>
<tr>
<td>Extra-hard</td>
<td>0.063 to 0.250 [1.60 to 6.35]</td>
<td>196 000 [1350]</td>
<td>(\ldots)</td>
<td>(\ldots)</td>
<td>(\ldots)</td>
</tr>
</tbody>
</table>

\(^a\) The gage length must be reported with the test results. \(4D = 4 \times \text{diameter} ; 4W = 4 \times \text{width} .\) Alternatively, a gage length corresponding to Test Methods E8/E8M or ISO 6892 may be used when agreed upon between supplier and purchaser. (5.65 times the square root of So, where So is the original cross sectional area).

\(^b\) 29-kN [3000-kgf] load.

\(^c\) Typically supplied as hot rolled bar for forging applications.

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9.1.2 Material shall conform to the appropriate requirements as to mechanical properties specified in Table 1 and Table 2.

9.1.3 The level of mechanical properties for material in conditions other than those included in Table 1 and Table 2, shall be specified in the purchase order.

9.1.4 Bar and wire in the cold worked condition can be supplied to a higher tensile strength and corresponding lower elongation as specified on the purchase order.

9.1.5 Fine wire in the cold drawn condition can be supplied to a higher tensile strength and corresponding lower elongation as specified on the purchase order.

9.2 Hardness:

9.2.1 Hardness values shall be determined in accordance with Test Method E10 or Test Methods E18.

9.2.2 When desired, hardness limits may be specified by the purchaser. Hardness determinations shall be made on the product cross section, midway between the center and surface, if the cross section is adequate.

9.2.3 Hardness values are for information only and shall not be used as a basis for rejection.

9.3 Number of Tests:

9.3.1 Perform at least one tension test from each lot. Should any of the test pieces not meet the specified requirements, test two additional test pieces representative of the same lot, in the same manner, for each failed test piece. The lot shall be considered in compliance only if all additional test pieces meet the specified requirements.

9.3.2 Tensile test results for which any specimen fractures outside the gage length shall be considered acceptable, if the elongation meets the minimum requirement specified. Refer to section 7.11.4 of Test Methods E8/E8M. If the elongation is less than the minimum requirement, discard the test and retest. Retest one specimen for each specimen that did not meet the minimum requirements.

10. Dimensions and Permissible Variations

10.1 Units of Measure:

10.1.1 Selection—This specification requires that the purchaser selects the units (SI or inch-pound) to be used for product certification. In the absence of a stated selection of units on the purchase order, this selection may be expressed by the purchaser in several alternate forms listed in order of precedence.

10.1.1.1 If the purchaser and supplier have a history of using specific units, these units shall continue to be certified until expressly changed by the purchaser.

10.1.1.2 In the absence of historic precedence, if the units used to define the product on the purchaser’s purchase order (PO), specification, and engineering drawing are consistent, these units shall be used by the supplier for product certification.

10.1.1.3 If the purchaser’s selection of units is unclear, the units of measure shall be agreed upon between the purchaser and supplier.

10.1.2 Conversion of Units—If the supplier’s test equipment does not report in the selected units, the test equipment units may be converted to the selected units for certification purposes. Accurate arithmetic conversion and proper use of significant digits should be observed when performing this conversion. IEEE/ASTM SI 10 provides guidelines for the use of SI units. Annex A of that standard provides conversion tables and Annex B provides rules for conversion and significance.
11. Special Tests

11.1 Bar, forging bar, wire, and fine wire conforming to this specification shall be capable of passing the intergranular corrosion susceptibility test in accordance with Practice E of Practices A262.

11.1.1 Samples in the hot worked condition shall be annealed prior to Practice E of Practices A262, sensitization heat treatment.

11.2 Bar, forging bar, wire, and fine wire conforming to this specification shall have a grain size of ASTM No. 5 or finer when measured in accordance with Test Methods E112.

11.2.1 It is preferred that samples for grain size determination be selected after the hot working operation or after the final annealing operation prior to the final cold working operation.

11.2.2 If samples are selected after a final cold working operation, specimens shall be tested in accordance with Test Methods E112 or as agreed to between the supplier and purchaser.

11.3 All centerless ground or peeled and polished round bar \( \geq 0.375 \text{ in.} \) [9.5 mm] in nominal diameter shall be ultrasonically inspected at final diameter according to AMS 2630, Class A1. Equivalent test methods may be substituted when agreed upon between the purchaser and supplier.

11.4 Billet shall be ultrasonically inspected prior to being hot rolled if ultrasonic inspection is not performed at a final diameter as specified in 11.3. Acceptance criteria shall be agreed upon between the purchaser and supplier.

11.4.1 Alternately, the purchaser may request that billet be ultrasonically inspected prior to being hot rolled even if ultrasonic inspection is performed at a final diameter. Acceptance criteria shall be agreed upon between the purchaser and supplier.

11.5 Any other special requirements shall be specified by the purchaser.

12. Significance of Numerical Limits

12.1 The following applies to all specified numerical limits in this specification. To determine conformance to these limits, an observed or calculated value shall be rounded to the nearest unit in the last right hand digit used in expressing the specification limit, in accordance with the rounding method of Practice E29.

13. Certification

13.1 The supplier shall provide a certification that the material was tested in accordance with this specification and met all requirements. A report of the test results shall be furnished to the purchaser at the time of shipment.

14. Quality Program Requirements

14.1 The supplier shall maintain a quality program, such as defined in ISO 9001, or similar.

15. Keywords

15.1 metals (for surgical implants); stainless steel; surgical applications wire; surgical implants

APPENDIXES

(Nonmandatory Information)

X1. RATIONALE

X1.1 The primary reason for this specification is to characterize composition and properties to ensure consistancy in the starting material used directly, or as modified by forging, in the manufacturing of medical devices.

X1.2 This low carbon alloy is selected to provide an extra measure of assurance that the material will be free from susceptibility to intergranular corrosion.

X1.3 There is a general consensus that a homogeneous metallurgical structure will be superior with respect to corrosion and fatigue resistance. Based upon this, metallurgical requirements include fine-grained austenitic structure free of ferrite, with low micro-inclusion content, and the capability of passing an intergranular corrosion susceptibility test.

X1.4 Acceptable metal conditions include hot worked, annealed, and all cold worked conditions, the choice dependent upon the implant design and application.

X1.5 Upper composition limits for nickel and lower composition limits for molybdenum have been changed in order to meet the latest requirements specified in ISO 5832–1, Composition D.

X1.6 A maximum nitrogen limit was previously added in accordance with the specified element requirements of similar austenitic stainless steels standardized by ASTM.

X1.7 The maximum copper value is considered a practical limit based on a statistical evaluation of commercially available material. Published information has shown no adverse effect for compositions containing up to 1.0 % copper content.

X1.8 The nickel range had previously been increased to ensure that compositions melted to the upper end of the
molybdenum range would be free of delta ferrite.

X1.9 ISO standards are listed for reference only. Although ISO standards listed in Section 2 are similar to the corresponding ASTM standards, they may not be identical. Use of an ISO standard in addition to or instead of a preferred ASTM standard may be negotiated between the purchaser and supplier.

X1.10 Molybdenum-enriched chi and sigma intermetallic compounds must not be present in the microstructure because of reduced austenitic corrosion resistance and possible embrittlement effects.

X1.11 Delta ferrite is a magnetic phase that must be absent in order to provide a completely nonmagnetic microstructure that will not cause torque, displacement, or heating in a Magnetic Resonance Imaging (MRI) environment.

X1.12 Units of Measure:

X1.12.1 ASTM Policy—ASTM is promoting the use of rationalized SI (metric) units in their standards. The F12.04 Committee has modified this specification to facilitate the transition by the medical materials industry to SI between now and 2018. In the first phase of this transition, running to 2013, the specifications will be structured to allow the use of either SI or inch-pound units. The choice of primary units in each specification will be determined by the industry using the specification. The change to SI units during this period may be initiated by the purchaser through his purchase documentation. In the second phase of this transition the specifications will be written with SI as the primary units. Harmonization with corresponding ISO documents should be considered when assigning the SI values.

X2. Biocompatibility

X2.1 The material composition covered by this specification has been employed successfully in human implant applications in contact with soft tissue and bone for over a decade. Due to the well characterized level of local biological response established by this material, it has been used as a control material in Practice F981.

X2.2 No known surgical implant material has ever been shown to be completely free of adverse reactions in the human body. However, long term clinical experience has shown an acceptable level of biological response can be expected, if the material is used in appropriate applications.

SUMMARY OF CHANGES

Committee F04 has identified the location of selected changes to this standard since the last issue (F138 – 13) that may impact the use of this standard. (Approved Oct. 1, 2013)

(1) Revised 11.4 and added 11.4.1.