

Standard Specification for Wrought Cobalt-28Chromium-6Molybdenum Alloys for Surgical Implants (UNS R31537, UNS R31538, and UNS R31539)¹

This standard is issued under the fixed designation F1537; the number immediately following the designation indicates the year of original adoption or, in the case of revision, the year of last revision. A number in parentheses indicates the year of last reapproval. A superscript epsilon (ε) indicates an editorial change since the last revision or reapproval.

1. Scope*

- 1.1 This specification covers the chemical, mechanical, and metallurgical requirements for three wrought cobalt-28chromium-6molybdenum alloys used for surgical implants. The properties specified apply specifically to wrought bar, rod, and wire.
- 1.2 The values stated in inch-pound units are to be regarded as standard. The values given in parentheses are mathematical conversions to SI units that are provided for information only and are not considered standard.

2. Referenced Documents

2.1 ASTM Standards:²

E8/E8M Test Methods for Tension Testing 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

E112 Test Methods for Determining Average Grain Size

E354 Test Methods for Chemical Analysis of High-Temperature, Electrical, Magnetic, and Other Similar Iron, Nickel, and Cobalt Alloys

F75 Specification for Cobalt-28 Chromium-6 Molybdenum Alloy Castings and Casting Alloy for Surgical Implants (UNS R30075)

F799 Specification for Cobalt-28Chromium-6Molybdenum Alloy Forgings for Surgical Implants (UNS R31537, R31538, R31539) 2.2 Aerospace Material Specifications:³

AMS 2248 Chemical Check Analysis Limits, Corrosion and Heat Resistant Steels and Alloys, Maraging and Other Highly-Alloyed Steels, and Iron Alloys

AMS 2269 Chemical Check Analysis Limits, Nickel, Nickel Alloys and Cobalt Alloys

AMS 2630 Inspection, Ultrasonic Product over 0.5 Inch (12.7 mm) Thick

2.3 ISO Standards:⁴

ISO 5832–12 Implants for Surgery—Metallic Materials—Part 12: Wrought Cobalt-Chromium-Molybdenum AlloyISO 6892 Metallic Materials Tensile Testing at Ambient Temperature

ISO 9001 Quality Management Systems—Requirements

3. Terminology

- 3.1 Definitions of Terms Specific to This Standard:
- 3.1.1 *lot*—the total number of mill products produced from the same melt heat under the same conditions at essentially the same time.

4. Ordering Information

- 4.1 Inquiries and orders for material under this specification shall include the following information:
 - 4.1.1 Quantity,
 - 4.1.2 ASTM designation, alloy number, and date of issue,
 - 4.1.3 Mechanical properties (See Section 7),
 - 4.1.4 Form (bar, rod or wire),
- 4.1.5 Applicable dimensions including size, thickness, width, and length (exact, random, or multiples) or drawing number,
 - 4.1.6 Condition (See Section 5),
 - 4.1.7 Special tests (if any), and
 - 4.1.8 Other requirements.

5. Materials and Manufacture

5.1 Product shall be furnished as specified below:

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

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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 Society of Automotive Engineers (SAE), 400 Commonwealth Dr., Warrendale, PA 15096-0001, http://www.sae.org.

⁴ Available from American National Standards Institute (ANSI), 25 W. 43rd St., 4th Floor, New York, NY 10036, http://www.ansi.org.

- 5.1.1 The annealed condition is typically supplied as a hot rolled and annealed product.
- 5.1.2 The hot worked condition is typically supplied as a hot rolled and unannealed product.
- 5.1.3 The warm worked condition is typically supplied as a thermomechanically processed product to achieve a strain-hardened structure.

6. Chemical Requirements

- 6.1 The cobalt-28chromium-6molybdenum alloys shall conform to the chemical requirements prescribed in Table 1. The supplier shall not ship material that is outside the limits specified in Table 1 for the applicable alloy.
- 6.1.1 Requirements for the major and minor elemental constituents are listed in Table 1. Also listed are important residual elements. Analysis for elements not listed in Table 1 is not required to verify compliance with this specification.
- 6.2 *Product Analysis*—The product analysis is either for the purpose of verifying the composition of a heat or lot or to determine variations in the composition within the heat.
- 6.2.1 Acceptance or rejection of a heat or lot of material may be made by the purchaser on the basis of this product analysis.
- 6.2.2 Product analysis tolerances do not broaden the specified heat analysis requirements but cover variations between laboratories in the measurement of chemical content. Product analysis limits shall be as specified in Table 2.

7. Mechanical Requirements

7.1 Tensile Properties:

- 7.1.1 Perform at least two tension tests from each lot. Should any of these 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.
- 7.1.2 Tensile test results for which any specimen fractures outside the gauge length shall be considered acceptable, if both the elongation and reduction of area meet the minimum

TABLE 1 Chemical Composition

				-						
Composition % (mass/mass)										
Element	Alloy 1 UNS R31537 (Low Carbon)		Alloy 2 UNS R31538 (High Carbon)		Alloy 3 UNS R31539 (Dispersion Strengthened)					
	min	max	min	max	min	max				
Carbon		0.14	0.15	0.35		0.14				
Aluminum					0.30	1.00				
Lanthanum					0.03	0.20				
Chromium	26.0	30.0	26.0	30.0	26.0	30.0				
Molybdenum	5.0	7.0	5.0	7.0	5.0	7.0				
Nickel		1.0		1.0		1.0				
Iron		0.75		0.75		0.75				
Silicon		1.0		1.0		1.0				
Manganese		1.0		1.0		1.0				
Nitrogen		0.25		0.25		0.25				
CahaltA	Dolones		Polonoo		Dalamas					

^A Approximately equal to the difference of 100 % and the sum percentage of the other specified elements. The percentage of cobalt difference is not required to be reported.

TABLE 2 Product Analysis Tolerances^{A,B}

Permissible Variation Under the Minimum		
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Limit or Over the Maximum Limit, % (mass/		
mass) ^C		
0.02		
0.05		
0.10		
0.01		
0.30		
0.15		
0.05		
0.03		
0.05		
0.03		
0.02		

^A See Test Methods E354.

requirements specified. Refer to subsections 7.11.4 and 7.11.5 of Test Methods E8/E8M.

7.1.3 The mechanical properties of test specimens shall conform to the requirements specified in Table 3.

7.2 Hardness:

- 7.2.1 Hardness values shall be determined in accordance with Test Methods E18.
- 7.2.2 Hardness values are for information only and shall not be used as a basis for rejection.

8. Microstructure Requirements

8.1 Bar, rod, and wire conforming to this specification shall have a homogeneous microstructure with an average grain size of ASTM No. 5 or finer when measured in accordance with Test Methods E112.

9. Ultrasonic Inspection

9.1 For finished thicknesses 0.250 in. (6.35 mm) and greater, inspection shall be per AMS 2630 Class A1. Equivalent test methods may be substituted when agreed to by purchaser and supplier. AMS 2630 indicates a minimum size of 0.500 in. (12.7 mm). The use of this specification to a

TABLE 3 Mechanical Requirements

			-		
Condition	Ultimate Tensile Strength min, psi (MPa)	Yield Strength (0.2 % offset), min, psi (MPa)	Elongation ^A min, %	Reduction in Area min, %	Hardness HRC, Typical
Annealed	130 000	75 000	20	20	25
	(897)	(517)			
Hot	145 000	101 000	12	12	28
Worked	(1000)	(700)			
Warm	170 000	120 000	12	12	35
Worked	(1172)	(827)			

^A Elongation of material 0.062 in. (1.575 mm) or greater in diameter or thickness shall be measured using a gauge length of 2 in. or 4D or 4W. The gauge length must be reported with the test results. The method for determining elongation of material under 0.062 in. (1.57 mm) in diameter or thickness may be negotiated. Alternately, a gauge length corresponding to 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 Refer to AMS 2269 for chemical check analysis limits (except nitrogen).

 $^{^{\}it C}$ For elements in which only a maximum percentage is indicated, the "under minimum limit" is not applicable.

D Refer to AMS 2248.

minimum size of 0.250 in. (6.35 mm) is done with agreement of producer and user members of the committee.

9.2 For finished thicknesses less than 0.250 in. (6.35 mm) and for shape bar that cannot be inspected at finish, intermediate size bars or billets shall be ultrasonically inspected per AMS 2630 Class A1 or as agreed upon by purchaser and supplier.

10. Significance of Numerical Limits

10.1 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.

11. Certification

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

12. Quality Program Requirements

12.1 The alloy supplier shall maintain a quality program or quality management system, such as ISO 9001.

13. Keywords

13.1 cobalt alloys (for surgical implants); cobalt-28chromium-6molybdenum; metals (for surgical implants)

APPENDIXES

(Nonmandatory Information)

X1. RATIONALE

- X1.1 The purpose of this specification is to characterize composition and properties to ensure consistency in wrought cobalt-28chromium-6molybdenum bar, rod, and wire used in the manufacturing of medical devices.
- X1.2 The carbon content of commercially available F1537 barstock varies to meet specific customer needs. The chemical composition was expanded into low and high carbon alloys (Alloys 1 and 2 in Table 1). Alloy 2 contains a greater volume fraction of carbides compared to Alloy 1.
- X1.3 Alloy 3 was added in Table 1 since bar, rod, and wire produced with this composition meet the current ASTM F1537 requirements. Alloy 3 is a dispersion-strengthened alloy containing fine aluminum-lanthanum oxides/nitrides. 5,6

- X1.4 The minimum mechanical properties of wrought bar, rod, and wire conforming to this specification are superior to the minimum mechanical properties of Specification F75 material in the as-cast condition.
- X1.5 Some complex metallic phases, such as carbides, oxides, or carbonitrides, or combinations thereof, may be present in the microstructure of this alloy.
- X1.6 ISO standards are listed for reference only. Although the ISO 5832–12 standard listed in Section 2 is similar to the corresponding ASTM standard, they are not identical. Use of the ISO standard instead of the preferred ASTM standard may be agreed upon between purchaser and supplier. The mechanical properties of warm-worked condition material, subsection 5.1.3, of this specification are equivalent to the ISO 5832–12 cold-worked condition.
- X1.7 Wrought material, typically hot-worked and unannealed with a surface suitable for forging, shall be fabricated and supplied in accordance with this specification when used as forging stock in the manufacture of forgings in accordance with Specification F799.

⁵ Wang, K. K., Gustavson, L. J., and Dumbleton, J. H., U.S. Patent #4,668,290, "Dispersion Strengthened Cobalt-Chromium-Molybdenum Alloy Produced By Gas Atomization", Filed August 13, 1985, Issued May 26, 1987.

⁶ Wang, K. K., Gustavson, L. J., and Dumbleton, J. H., "The Development of a New Dispersion Strengthened Vitallium Alloy for Medical Implants", *Modern Developments in Powder Metallurgy*, Vol 20, 1988, pp. 361–375, Metal Powder Industries Federation, 105 College Road East, Princeton, NJ 08540-6692.



X2. BIOCOMPATIBILITY

X2.1 The alloy compositions covered by this specification have been employed successfully in human implant applications in contact with soft tissue and bone for over a decade (reference 510k #K852964 for Alloy 3, which has been used since 1985).

X2.2 No known surgical implant material has ever been shown to be completely free of adverse reactions in the human

body. Long term clinical experience of the use of the material referred to in this specification, however, has shown that 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 (F1537 - 08) that may impact the use of this standard. (Approved Nov. 15, 2011.)

(1) Revised section on ultrasonic inspection to make it explicit with regards to specification, inspection type, and sizes. This section clearly defines how ultrasonic inspection will be performed on small sizes and special shapes.

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