Also, this rule does not have tribal implications under Executive Order 13175, Consultation and Coordination with Indian Tribal Governments, because it does not have a substantial direct effect on one or more Indian tribes, on the relationship between the Federal Government and Indian tribes, or on the distribution of power and responsibilities between the Federal Government and Indian tribes.

E. Unfunded Mandates Reform Act

The Unfunded Mandates Reform Act of 1995 (2 U.S.C. 1531–1538) requires Federal agencies to assess the effects of their discretionary regulatory actions. In particular, the Act addresses actions that may result in the expenditure by a State, local, or tribal government, in the aggregate, or by the private sector of \$100,000,000 (adjusted for inflation) or more in any one year. Though this rule will not result in such an expenditure, we do discuss the effects of this rule elsewhere in this preamble.

F. Environment

We have analyzed this rule under Department of Homeland Security Directive 023-01, Rev. 1, associated implementing instructions, and **Environmental Planning COMDTINST** 5090.1 (series), which guide the Coast Guard in complying with the National Environmental Policy Act of 1969 (42) U.S.C. 4321–4370f), and have determined that this action is one of a category of actions that do not individually or cumulatively have a significant effect on the human environment. This rule involves a safety zone lasting 1.5 hours that will prohibit entry within a 1 square mile area of the Neuse River on December 5, 2020, from 4 p.m. to 5:30 p.m. It is categorically excluded from further review under paragraph L60(a) of Appendix A, Table 1 of DHS Instruction Manual 023-01-001-01, Rev. 1. A Record of **Environmental Consideration** supporting this determination is available in the docket. For instructions on locating the docket, see the ADDRESSES section of this preamble.

G. Protest Activities

The Coast Guard respects the First Amendment rights of protesters. Protesters are asked to call or email the person listed in the **FOR FURTHER INFORMATION CONTACT** section to coordinate protest activities so that your message can be received without jeopardizing the safety or security of people, places or vessels.

List of Subjects in 33 CFR Part 165

Harbors, Marine safety, Navigation (water), Reporting and recordkeeping requirements, Security measures, Waterways.

For the reasons discussed in the preamble, the Coast Guard amends 33 CFR part 165 as follows:

PART 165—REGULATED NAVIGATION AREAS AND LIMITED ACCESS AREAS

■ 1. The authority citation for part 165 continues to read as follows:

Authority: 46 U.S.C. 70034, 70051; 33 CFR 1.05–1, 6.04–1, 6.04–6, and 160.5; Department of Homeland Security Delegation No. 0170.1.

■ 2. Add § 165.T05–0645 to read as follows:

§ 165.T05-0645 Safety Zone; Neuse River, Airshow, New Bern, NC.

- (a) Location. The following area is a safety zone: All navigable waters of the Neuse River in New Bern, North Carolina, inside an area starting from approximate positions: Latitude 35°06′32″ N, longitude 077°01′54″ W, then north to latitude 35°06′55" N, longitude 077°02′04" W, then east to latitude 35°07'06" N, longitude 077°01'27" W, then southeast to latitude 35°06'49" N, longitude 077°01'12" W, then south to latitude 35°06'08" N longitude 077°01′18" W, then west to latitude 35°06′02″ N, longitude 077°01′57" W, then north to the point of origin, for a total area of approximately 1 mile square.
- (b) *Definitions*. As used in this section—

 ${\it Captain~of~the~Port~(COTP)}~{\it means~the}~{\it Commander, Sector~North~Carolina}.$

Designated representative means a Coast Guard Patrol Commander, including a Coast Guard commissioned, warrant, or petty officer designated by the Captain of the Port North Carolina (COTP) for the enforcement of the safety zone.

(c) Regulations. (1) The general regulations governing safety zones in § 165.23 apply to the area described in paragraph (a) of this section.

- (2) Entry into or remaining in this safety zone is prohibited unless authorized by the COTP North Carolina or the COTP North Carolina's designated representative. Unless permission to remain in the zone has been granted by the COTP North Carolina or the COTP North Carolina or the COTP North Carolina's designated representative, a vessel within this safety zone must immediately depart the zone when this section becomes effective.
- (3) The Captain of the Port, North Carolina can be reached through the

Coast Guard Sector North Carolina Command Duty Officer, Wilmington, North Carolina, at telephone number 910–343–3882.

- (4) The Coast Guard and designated security vessels enforcing the safety zone can be contacted on VHF–FM marine band radio channel 13 (165.65 MHz) and channel 16 (156.8 MHz).
- (d) *Enforcement*. The U.S. Coast Guard may be assisted in the patrol and enforcement of the safety zone by Federal, State, and local agencies.
- (e) Enforcement period. This regulation will be enforced from 4 p.m. through 5:30 p.m. on December 5, 2020.

Dated: November 17, 2020.

Matthew J. Baer,

Captain, U. S. Coast Guard, Captain of the Port North Carolina.

[FR Doc. 2020–25688 Filed 11–27–20; 8:45 am]

BILLING CODE 9110-04-P

DEPARTMENT OF VETERANS AFFAIRS

38 CFR Part 4

RIN 2900-AP88

Schedule for Rating Disabilities: Musculoskeletal System and Muscle Injuries

AGENCY: Department of Veterans Affairs. **ACTION:** Final rule.

SUMMARY: This document amends the Department of Veterans Affairs (VA) Schedule for Rating Disabilities ("VASRD" or "rating schedule") by revising the portion of the rating schedule that addresses the musculoskeletal system. The purpose of this revision is to ensure that this portion of the rating schedule uses current medical terminology and provides detailed and updated criteria for the evaluation of musculoskeletal disabilities.

DATES: This rule is effective February 7, 2021.

FOR FURTHER INFORMATION CONTACT: Gary Reynolds, M.D., Regulations Staff (211C), Compensation Service, Veterans Benefits Administration, Department of Veterans Affairs, 810 Vermont Avenue NW, Washington, DC 20420, (202) 461–9700. (This is not a toll-free number.)

SUPPLEMENTARY INFORMATION: The National Defense Authorization Act of 2004, secs. 1501–07, Public Law 108–136, Stat. 1392, established the Veterans' Disability Benefits Commission (the "Commission"). Section 1502 of Public Law 108–136 mandated the Commission to study

ways to improve the disability compensation system for military veterans. The Commission consulted with the Institute of Medicine (IOM) (now named the National Academy of Medicine) to review the medical aspects of current policies. In 2007, the IOM released its report titled "A 21st Century System for Evaluating Veterans for Disability Benefits." (Micahel McGeary et al. eds. 2007).

The IOM report noted that the VA Rating Schedule for Disabilities was inadequate in areas because it contained obsolete information and did not sufficiently integrate current and accepted diagnostic procedures as well as the lack of current knowledge of the relationships between conditions and comorbidities. Following the release of the IOM report, VA created a musculoskeletal system workgroup to: (1) Improve and update the process that VA uses to assign levels of disability after it grants service connection; (2) improve the fairness in adjudicating disability benefits for service-connected veterans; and (3) invite public participation.

VA began rulemaking to remove obsolete diagnostic codes, modernize the names of selected diagnostic codes, revise descriptions and criteria, and add new diagnostic codes. VA published a proposed rule to revise the regulations involving the musculoskeletal system within VASRD on August 1, 2017 (82 FR 35719). Specifically, VA proposed to rename conditions to reflect current medicine, remove obsolete conditions, clarify ambiguities, and add conditions that previously did not have diagnostic codes. Interested persons were invited to submit comments on or before October 2, 2017. VA received comments from the National Organization of Veterans' Advocates, American Association of Nurse Practitioners, Paralyzed Veterans of America, and nine individuals. VA has made limited changes based on these comments, as discussed below.

General Terminology Changes

Two separate comments recommending specific terminology changes were received.

One commenter suggested incorporating terminology used by claimants or seen in service treatment records into the VASRD regulations. The commenter stated that field medics do not always incorporate medical terminology or use treatises when entering information in a servicemember's medical record. The commenter also noted that individual claimants may not have sufficient medical training to utilize specific

technical terminology when claiming a given disability. A stated intent of the current update to the rating schedule, as stated in the preamble to the proposed rule, is to employ current medical terminology in order to clarify and standardize the disability criteria. Accordingly, VA relies on medical standards and treatises when updating terminology.

As to the effect of technical terminology in part 4 on a veteran attempting to claim disability, there is none. Claimants are not required to possess medical knowledge or expertise when describing a claimed condition; they are simply required to describe their disability and/or symptoms as they experience and observe them. Brokowski v. Shinseki, 23 Vet. App. 79, 86-87 (2009). Moreover, VA reviews medical records with the understanding that different examiners, at different times, will not describe the same disability in the same language; it is the responsibility of the rating specialist to interpret reports of examination in the light of the whole recorded history, reconciling the various reports into a consistent picture so that the current rating may accurately reflect the elements of disability present. 38 CFR 4.2. Accordingly, VA reviews the entire evidentiary record in light of the disability claimed, circumstances of military service, and all other applicable records to create a cohesive picture of the disability in question; it is not the responsibility of the claimant or a military medical provider to employ terminology that necessarily matches the VASRD. Thus, VA makes no changes related to this comment.

Another commenter suggested use of the phrases "greater than or equal to" and "less than or equal to" rather than "limited to XX degrees or more" or "limited to XX degrees or less" for criteria based on numerical range of motion measurements. While this comment was taken into consideration, VA notes the phrases "limited to XX degrees or more" or "limited to XX degrees or less" are consistent with medically-accepted language used in the VASRD for range of motion measurement and elsewhere, and are well-understood and applied by VA claims processors efficiently and accurately. Accordingly, VA makes no changes based on this comment.

Musculoskeletal Diagnostic Codes

I. Diagnostic Codes (DCs) 5002–5009

One commenter asked if there was a DC for infectious arthritis. While there is not a standalone DC for infectious arthritis, infectious arthritis may be

evaluated under DCs 5004 through 5009, depending on the infection associated with the arthritic findings. VA makes no change based on this comment.

Another commenter requested that VA use the same non-exhaustive list of conditions listed in proposed DC 5002's Note (1) for other selected DCs (5054, 5055, and 5250-5255). The list of conditions in DC 5002 is being provided to further explain the change from this DC contemplating a specific condition to contemplating a category of conditions. The other DCs suggested by the commenter are unlike proposed DC 5002 because they employ criteria based on a specific procedure (DCs 5054 & 5055) or defined range of motion measurement (DCs 5250-5255). VA makes no changes based on this comment.

Lastly, a commenter expressed concern that the directive to "assign the higher evaluation" under DC 5002 could result in situations where an active disease process results in a lower evaluation than if the residuals of the disease itself were evaluated. The directive in proposed Note (3) for DC 5002 specifically addresses this concern. As indicated in the preamble to the proposed rule, the purpose of Note (3) is to prevent ratings for both residuals and active disease process at the same time; instead, the Note requires claims processors to assign the evaluation more advantageous to the claimant: An evaluation for active disease process OR an evaluation for the residual effects of the disease (including combined and/or bilateral factors, where applicable). Accordingly, VA makes no change based on this comment.

II. DCs 5010-5024

One commenter suggested that arthritis ratings under DC 5010 resulting from separate traumas should not receive a combined evaluation under 38 CFR 4.25. VA makes no changes based on this comment, as the evaluations under the VASRD are based on the average impairment in earnings due to disabilities resulting from military service; the specific incidents or causes during military service are generally immaterial to a rating. As a practical matter, attempting to categorize functional impairment by specific traumatic instances would prove ineffective and often impossible, as specific instances of trauma are not necessarily captured in the treatment record for an individual.

One commenter asked how DC 5011 would help evaluate a case of facial fractures, hearing loss, a collapsed sinus, eye injury and so forth. VA notes

that DC 5011 does not provide specific evaluation criteria; rather, it serves as a standalone diagnostic code to track instances of decompression illness (also known as generalized barotrauma or the bends). As noted in the preamble to the proposed rule, residual manifestations of decompression illness often involve other body systems; the proposed evaluation criteria specifically directs claims processors to evaluate residuals under the appropriate body system. Accordingly, specific residual injuries will be evaluated under the most appropriate diagnostic code in the VASRD, in accordance with the findings and disability present. VA makes no changes based on this comment.

Another commenter questioned what effect the changes to DCs 5010, 5013 and 5014 would have on determinations under 38 CFR 3.309. 38 CFR 3.309 identifies diseases subject to presumptive service connection where certain circumstances of military service are otherwise met. This section pertains to establishing service connection; it does not involve the evaluation of any specified disability. The current rulemaking has no impact on the provisions of section 3.309 and therefore VA makes no changes based on this comment.

Another commenter recommended using the phrase "medically-directed therapy" as opposed to "prescribed therapeutic procedure" in the Note to DC 5012. While this comment was taken into consideration, VA's selected term has a specific meaning and indicates a prescribed course of treatment, as determined by a qualified medical professional, as evidence of the severity of the disability and disease, in the professional opinion of the provider. "Medically-directed" does not have the same meaning as "prescribed" and its use here would leave open for interpretation therapies that are either suggested at a lower level of necessity or directed by someone who is not licensed/qualified to prescribe treatment for malignancies. VA makes no changes based on this comment.

One commenter suggested adding a Note to DC 5014 indicating that, if medical evidence does not specifically indicate or state there are no residuals, there is insufficient evidence to apply the provisions of DC 5014. VA appreciates this comment but notes that 38 CFR 4.2 specifically instructs claims processors to return examinations as inadequate for evaluation purposes if the examination report does not contain sufficient detail or if a diagnosis is not supported by the findings on examination. Accordingly, the suggested

Note would be duplicative of current regulations and VA makes no change.

Also, a commenter suggested adding notes to indicate where hydrarthrosis, synovitis, and periostitis could be evaluated since VA proposed removing specific DCs for these conditions. As noted in the preamble to the proposed rule, hydrarthrosis and synovitis are signs of underlying conditions that are already captured within the evaluation criteria of other DCs. Likewise, periostitis is a non-specific inflammatory process caused by underlying conditions that can be rated in accordance with the primary diagnosis. VA sees no need to limit these signs to specific DCs; they will be evaluated with an underlying diagnosis. VA makes no changes based on this comment.

Finally, on further review, the sentence following DC 5024 is more aptly described as a Note to DCs 5013 through 5024. As such, the final rule recharacterizes it as a Note and removes as unnecessary the proposed limitation that gout only be evaluated under DC 5003.

III. DCs 5051-5056 (Introductory Notes)

One commenter requested clarification as to why joint resurfacing and total joint replacement qualify for 100 percent disability compensation during the convalescent period, but partial joint replacement does not. VA recognizes that partial joint replacement (more accurately referred to as subtotal joint replacement) may result in disability in a manner similar to joint resurfacing and/or total joint replacement. However, VA currently lacks sufficient data to determine that partial joint replacement warrants a temporary post-surgical rating in lieu of a rating based on the effects of the underlying disability. To that end, VA will consider adding criteria specific to subtotal joint replacement in a future rulemaking, once sufficient evidence is received and reviewed to provide adequate evaluation criteria.

One commenter asked if revision procedures were eligible for the same compensation as the original procedures. While this comment was asked about hip replacement, it could be applied to all of the prosthetic replacement DCs. If the original complete prosthetic component is replaced, or, in addition to replacement of the original component, additional components are installed, then the revision procedure should be evaluated in the same manner as the initial procedure. In other words, if the revision fully replaces the original total prosthetic joint replacement, VA treats

the complete revision procedure in the same manner as the initial total joint replacement. To that end, in this final rule, VA has recharacterized the proposed note at the beginning of the "Prosthetic Implants and Resurfacing" subsection as Note (1) and added a Note (2) that directs claim processors to only evaluate revision procedures in the same manner as the original procedure if the revision completely replaces the original components.

For organization and clarity, VA has also moved three other notes to the beginning of the "Prosthetic Implants and Resurfacing" subsection and added a clarifying instruction. Specifically, the note immediately following DC 5111 has been moved to the beginning of the subsection and redesignated as Note (3). DC 5053's note and DC 5056's Note (1), which were identical, have been moved and redesignated as Note (4). An instruction that clarifies when the 100 percent evaluation period begins and ends for DCs 5054 and 5055 is provided as Note (5). And Note (2) under DC 5056 has been moved and redesignated as Note (6).

IV. DCs 5054 and 5055

Multiple comments were received for DCs 5054 and 5055. Generalized objections included two commenters who shared their personal histories involving revision procedures/surgeries on their hips as the underlying basis for their objections. Two commenters also expressed reservations with the reduction in the convalescent period for these DCs because of non-sedentary or physically demanding occupations, as well as additional service-connected disabilities that potentially complicate the evaluation. In regard to using personal experiences to justify any objection to the proposed changes, VA notes that 38 U.S.C. 1155 (the statute that governs implementation of the ratings schedule) provides that ratings shall be based, as far as practicable, upon the average impairments of earning capacity resulting from such injuries in civilian occupations. Accordingly, VA formulates the VASRD based on average impairments in civil occupations, not isolated personal experiences or the demands of specific occupations. In addition, the reduction in convalescent periods is based on average recovery times, as noted in the proposed rulemaking and sources cited therein. There are provisions to address exceptional individual circumstances on a case-by-case basis that fall outside the scope of this rulemaking. No changes are made based on those comments.

Another commenter disputed the study cited in the preamble to the proposed rule. The commenter used a quotation from the authors characterizing the methodological quality as moderate to low and comparisons of rates and speeds of return to work being hampered by large variations in patient selection and measurement methods. VA disagrees that the limitations identified by the commenter should invalidate the justification to reduce the convalescent period from 12 months to 4 months for hip and knee replacements. There are multiple studies within the medical literature which demonstrate sufficient functional recovery well short of 12 months. The study cited in the proposed rule focused upon a specific outcome (return to work without restriction), rather than completion of the associated rehabilitation program. VA convalescence rates are awarded at the 100 percent level—which, in accordance with the criteria throughout 38 CFR part 4, equates to a complete inability to work. Following the convalescent period, VA assigns a nonconvalescent evaluation based on residual functional impairment, the purpose of which is to assess residual disability and compensate for average earnings loss based on said residual disability.

One commenter proposed that a reduction in benefits for these DCs occur only after mandatory examination. Post-convalescence reductions for these conditions occur without a mandatory examination, due to the common nature of these medical procedures as well as the expected outcome and residuals, as supported by medical evidence cited in the preamble to the proposed rule. As stated in 38 CFR 4.1, the percentage ratings represent as far as can practicably be determined the average impairment in earning capacity resulting from such diseases and injuries and their residual conditions in civil occupations. VA acknowledges that there may be individual circumstances which require additional consideration due to worsethan-expected residuals or the factual need for additional convalescence. In these circumstances, a claimant may submit a claim with pertinent treatment records to support an increased evaluation for residuals or additional convalescence, all without requiring a mandatory examination. VA makes no changes based on this comment.

Another commenter proposed to extend the convalescent period whenever a revision procedure is performed. While a revision procedure may require additional time in the hospital following the procedure, this time typically amounts to a few days. Additionally, while the recovery may be potentially slower following a revision, VA is currently unaware of published medical literature which quantifies this recovery in a manner sufficient to identify a unique and/or extended period of convalescence for purposes of the VASRD. Should such evidence exist at a future date, VA will review it and consider revisions to the criteria as necessary. At this time, however, VA makes no changes based on this comment.

One commenter disagreed with the proposed reduction in the convalescent period because (1) there was little to no public support for such a reduction and (2) the studies used to support the reduction were not specific to veterans. The language in 38 U.S.C. 1155 specifically contemplates a schedule of ratings based on the average impairment in earnings from civil occupations, with revisions from time to time in accordance with experience. If a particular disability's effect on earnings capacity measurably changes (usually through a combination of improved medical management and job market changes), VA complies with its statutory authority by revising the criteria contained in the VASRD to ensure evaluations are consistent with available data. VA is unaware of any study pertinent to the disabilities at issue that quantifies a different impact of a specific disability or disabilities on the general population comparative to the veteran population. Should such information become available, VA will review it along with all other available scientific, medical, and economic data available to ensure the VASRD provides the most accurate and adequate evaluations. At this time, however, VA makes no revisions based on these comments.

One commenter offered an alternative schema to VA's proposal for DC 5054. This commenter recommended a separate DC be created for hip resurfacing. The commenter provided multiple sources to justify a minimum evaluation within the criteria for this alternative schema (citing multiple sources which compared resurfacing to prosthetic replacement). The commenter also criticized VA's proposed revision for DC 5054, asserting it was contradictory to government and industry standards. The commenter asserted that the purpose and advantage of hip resurfacing is bone preservation, not improved range of motion or activity. Finally, the commenter stated that VA should evaluate resurfacing and total arthroplasty under separate DCs.

VA makes no changes based on these comments for several reasons. First, VA disagrees with the statement that a minimum evaluation for hip resurfacing post convalescence similar to total arthroplasty is required. As noted in the preamble to the proposed rule, joint resurfacing preserves more of the original anatomy of the joint, leading to greater functional potential, and ultimately less occupational disability or impairment in earnings capacity compared to a total arthroplasty. Also, the sources cited by the commenter refer to the hip resurfacing procedure itself, the unique complications associated with resurfacing, and how it compares to total arthroplasty. While relevant in individual cases, potential complications in and of themselves do not consistently predict either residual occupational disability or average impairment in earnings capacity in a manner consistent with VA's authority to maintain and revise the VASRD. Additionally, as stated previously in response to similar comments, should individual complications arise, VA has the means to address these unique situations on a case-by-case basis either through additional convalescence or increased evaluations. With regard to the comment that VA's proposed revision is contrary to government and industry standards, VA notes that the commenter did not provide resources which establish either government or industry standards for the evaluation of resurfacing or residual disability in light of occupational impairment or earnings loss, and VA is unaware of an official government or industry standard upon which to base any changes to the proposed rule.

However, to further clarify VA's intent to provide a minimum evaluation following only total joint replacement, VA has added language to the Note following final DCs 5054 and 5055 clarifying that the minimum evaluation does not apply to resurfacing. Regarding the comment that range of motion as a residual for hip resurfacing would not be addressed under other DCs, VA notes that the (proposed and now final) rule directs the rater to use DCs 5250 through 5255 to evaluate such residuals. DCs 5251, 5252, and 5253 address decreased range of motion of the hip joint as a potential residual. Additionally, VA notes that the commenter's reference to "bone preservation" is consistent with VA's explanation in the preamble of the proposed rule (noting that resurfacing "preserves more of the original anatomy"). In any event, the intent of the VASRD is to assess and evaluate

residual disability and occupational impairment. Currently, VA is unaware of medical or economic data to support an evaluation for hip resurfacing based on the quantity of bone preserved. Additionally, VA notes that a single DC for both resurfacing and prosthetic component replacement is more appropriate than having separate DCs, as the symptoms leading up to and resulting from both procedures are similar and predictable (loss of weight bearing capability, muscle strength/ endurance, and range of motion due to complications such as component loosening, infection, etc.).

V. DCs 5120-5173

One commenter stated that the rating for disarticulation of the shoulder in DC 5120 may conflict with the rules for rating the shoulder muscles and ankylosed joints. VA notes that a disarticulation at the shoulder joint removes all the joints along with their associated muscles of the upper extremity. Thus, there would be no muscles or joints remaining, and therefore no evaluation based on ankylosis of the joint could be assigned.

Another commenter asked why VA removed prompts from certain DCs directing claims processors to consider eligibility for special monthly compensation (SMC). The removal of the prompts from DCs in the proposed rule was an unintentional error. Accordingly, VA has re-inserted the prompts to consider SMC for all

applicable DCs.

One commenter questioned both the need and the basis for the proposed changes to DC 5170. The commenter disagreed with VA's proposed criteria modification to include different amputation degrees within one DC and argued that at least two different DCs was a more appropriate approach. As noted in the preamble to the proposed rule, VA is adding this terminology to incorporate a residual which causes a similar disability to the one captured by current DC 5170. Furthermore, the amputation levels captured in the (proposed and now final) DC cause similar effects on occupational disability and impairment of earnings capacity. By grouping conditions and injuries with similar functional impairment together, VA provides accurate and adequate evaluations that reflect actual functional impairment while also providing more efficient and timely delivery of benefits.

VI. DCs 5235-5243

One commenter requested that VA include more medical diagnoses synonymous with intervertebral disc

syndrome (IVDS) and arthritis because, in the commenter's view, claims processors are inconsistent with acknowledging other similar conditions/ diagnoses that are not specifically labeled as IVDS, arthritis, or degenerative joint disease (DJD). VA's original intent was to classify disability associated with IVDS under DC 5243 and all other intervertebral disc disabilities under DC 5242. To clarify that issue, VA has added such an instruction to final DC 5243.

VII. DC 5244

For newly proposed DC 5244, two commenters had questions, and one commenter offered to provide training assistance to claims processors learning how to evaluate this newly proposed DC. The issue of training is beyond the scope of this rulemaking and therefore VA does not respond. One commenter stated that using the term "paraplegia" was problematic because it lumped a number of disabilities together and because paraplegia has a legal meaning. Specifically, the commenter questioned if paraplegia under DC 5244 also applies to paraplegia caused by amyotrophic lateral sclerosis (ALS) or multiple sclerosis (MS) and whether anal and bladder sphincter control impairment is necessary for assigning paraplegia under this DC, as is required to qualify for SMC under 38 CFR 3.350(e)(2), which is titled *Paraplegia*. The other commenter asked if incomplete paralysis is compensable. First, VA intended DC 5244 to rate paralysis resulting from trauma, as indicated in the title. It is separate and distinct from paralysis caused by either ALS or MS, which are neurological diseases and are rated using the appropriate neurological DC hyphenated with DC 5110 (loss of use of both feet). Second, although paraplegia is the title of § 3.350(e)(2), that provision provides requirements for SMC; paraplegia awarded under DC 5244 does not require impairment of anal and bladder sphincter control. Third, with regard to the comment on incomplete versus complete paralysis, VA has provided a note in this final rule that, if traumatic paralysis does not cause loss of use of both hands or both feet, it is incomplete paralysis and must be rated using the appropriate diagnostic code (e.g., 38 CFR 4.124a, Diseases of the Peripheral Nerves).

VIII. DCs 5255 and 5257

One commenter concurred with the proposed changes to DC 5255. VA thanks the commenter for the input. Other commenters (1) asked if patellofemoral pain syndrome (PFPS) was included in DC 5255; (2) asked

what would happen to DCs 5258 and 5259, given the proposed changes to DC 5257; and (3) recommended that claims processors be provided additional guidance for evaluating malunion under DC 5255. First, PFPS is a symptom that may result from patellar instability, but is a less appropriate fit for DC 5255, which contains criteria requiring fractures or malunions. Second, VA intends no changes to DCs 5258 or 5259, as they involve different components of the knee; accordingly, the changes to DC 5257 have no impact on DCs 5258 and 5259. Lastly, VA will provide nonregulatory guidance and training to claims processors for evaluating malunion under DC 5255.

Four additional commenters had concerns with and suggested alternatives to the proposed criteria of DC 5257. The first commenter expressed concern that the term "physician prescribed" excludes nurse practitioners, though such prescriptions are well within their scope of practice. VA agrees, and has substituted "medical provider" in place of "physician" to indicate that such instructions are intended to include qualified medical providers such as nurse practicioners.

The second commenter argued that (1) there is subjectivity with measuring translation; and (2) operative intervention should not be the basis for distinguishing a 30 percent evaluation from a 20 percent evaluation. After review, VA agrees that using translation can add an unintended amount of subjectivity to the evaluation criteria. To that end, VA has revised the proposed criteria to remove the reference to translation, and, instead, will use the elements of ligament status, instability, and need for assistive devices/bracing. A 10 percent evaluation will be granted if a sprained, incompletely torn ligament, or completely torn ligament (whether repaired, unrepaired, or failed repair) causes persistent instability but does not require a prescription for either bracing or an assistive device for ambulation. A 20 percent evaluation will be granted under one of two circumstances: (a) In the presence of a sprained, incompletely torn ligament, or repaired completely torn ligament that causes persistent instability and a medical provider prescribes a brace and/or assistive device; or, (b) in the presence of an unrepaired completely torn ligament or completely torn ligament with failed repair that causes persistent instability and requires a prescription for either a brace or an assistive device for ambulation. A 30 percent evaluation will be granted for an unrepaired completely torn ligament or completely torn ligament with failed

repair that requires a prescription for both a brace and an assistive device for ambulation. As to the original comment, this final rule considers both operative intervention and prescriptions as a basis for distinguishing the 30 percent and 20 percent evaluations. As a result of these changes, proposed Note (1), providing measurements of joint translation, has been withdrawn.

The third commenter felt that VA gave no explanation for the new criteria, that the criteria should include assistive devices and/or bracing whether prescribed by a provider or not, and that the criteria requiring both an assistive device and bracing was too restrictive. In the preamble to the proposed rule, VA provided a full explanation for the evaluation criteria for knee instability, citing multiple peer-reviewed medical sources which further support the criteria used. Regarding the requirement for provider-prescribed bracing, braces and other assistive devices are commonly and readily available for purchase without prescription; the use of such devices, without a prescription, does not always demonstrate the presence of a knee disability impairing earning capacity. A qualified medical professional's prescription, however, provides objective evidence of the instability. Accordingly, for purposes of assessing the severity of knee instability, this (proposed and final) rule considers bracing in its evaluation criteria only when the brace or assistive device is prescribed by a provider. Moreover, to the extent the commenter believes that requiring bracing and an assistive device is too restrictive, this final rule provides a 20% rating where only one of the two has been prescribed.

The fourth commenter asserted that the proposed changes to DC 5257 (1) will result in compensation that is either completely detached from functional loss or not commensurate with the functional loss being evaluated; (2) completely ignore functional loss and misplace emphasis on physical abnormalities and recommended treatment; and (3) did not consider knee instability caused by conditions other than ligament damage.

VA appreciates the comment, but disagrees with the commenter's first assertion. Per 38 U.S.C. 1155, the schedule and its ratings shall be based, as far as practicable, upon the average impairments of earning capacity resulting from such injuries in civil occupations. VA compensates for functional loss that results in an impairment of earning capacity. The criteria for DC 5257, as indicated in the preamble to the proposed rule, incorporate both functional loss

elements (assistive devices & bracing), as well as diagnostic elements (sprain, incomplete ligament tear, complete ligament tear). These criteria, which rely upon published sources reflecting current medical standards, serve as accurate proxies for functional loss of the magnitude that negatively impacts earnings. Furthermore, the proposed (and now final) criteria are easily observed and measured. Additionally, given the progressive manner of the criteria, VA provides compensation commensurate with the severity of the disability.

As to the commenter's second assertion that the proposed criteria base evaluations on recommended treatment, that is not the case. The proposed (and now final) criteria compensate for residual disability after specific treatment interventions are prescribed, not on the prescribed treatment itself, as well as observable and measurable factors to create a more complete assessment for evaluation purposes.

Third, with regards to the causes for knee instability other than ligament damage, VA intended the evaluation for patellar instability to be limited to the patellofemoral complex only. Thus, this final rule clarifies the proposed criteria and requires a diagnosed condition involving the patellofemoral complex for a patellar instability evaluation. A history of surgical repair (or the lack thereof) and the prescriptions for the instability dictate whether that evaluation will be 10, 20, or 30 percent (consistent with the format for recurrent subluxation evaluations).

Given this revision, VA has added a note (Note (1)) explaining that the patellofemoral complex consists of the quadriceps tendon, patella (knee cap), and patellar tendon. Proposed Note (2), despite technical edits, still provides that certain surgical procedures do not qualify as surgical repair under the patellar instability provisions of this DC.

In further response to the commenter's contention, we note that knee instability resulting from muscle failure can be evaluated under DC 5313 or DC 5314. Furthermore, with regards to knee instability and specific occupations, which the commenter also raised, compensation is based on the average of impairment in earning capacity for civil occupations, not the severity of disability encountered in selected occupations. Lastly, the language alternatively proposed by the commenter, which stems from a 2003 VA proposal, does not accommodate patellar instability, a shortcoming VA is unwilling to accept. VA notes that the 2003 proposal was withdrawn specifically to address concerns and

issues with the rulemaking and to develop a new proposal at a later date. 69 FR 22757. Therefore, VA makes no revisions based on this commenter's input.

IX. DC 5262

Unrelated to any particular comment, VA has revised the language of DC 5262 in this final rule to provide clarity on the specific criteria distinguishing the 30, 20, and 10 percent ratings for shin splints. Moreover, VA has decided not to adopt a rule that would require imaging evidence for a compensable rating; as the preamble to the proposed rule noted, shin splints are typically diagnosed—and can be properly assessed—by history and physical examination. M. Winters et al., "Medial tibial stress syndrome can be diagnosed reliably using history and physical examination," 52(19) Br. J. Sports Med.1267-72 (2018).

As to the comments, one commenter asked two questions: (1) Is there ever a scenario where shin splints and fractured tibia/fibula do not have overlapping symptoms, and (2) Is a distal fracture rated as an ankle disability and shin splints as a knee disability? Whether or not symptoms from shin splints and a certain fracture may or may not overlap is a medical question for medical examiners in individual cases. Therefore, VA will not speculate on the answer to the first question here. In regard to the second question, VA's intent is that a tibia/ fibula malunion be rated as either an ankle or knee disability. Beyond malunion, however, uncomplicated tibia/fibula fractures should still be rated under DC 5262.

X. DCs 5278-5285

Three commenters provided input for the proposed changes to these codes. Besides the commenters who concurred, one commenter disagreed with the criteria for proposed DC 5285, contending that veterans who are not surgical candidates are punished by the proposed 20 and 30 percent criteria. To address those veterans who would potentially benefit from surgical intervention, but who are not surgical candidates, VA is adding a Note (2) to DC 5285 indicating that a veteran who is recommended surgical intervention for plantar fasciitis but is not a surgical candidate would be eligible for either the 20 or 30 percent evaluation levels. The Note proposed in the proposed rule is recharacterized as Note (1). VA has also revised the wording of DC 5285 for clarity.

Muscle Injuries

One commenter concurred with proposed DC 5330. VA thanks the commenter for the input.

Miscellaneous Issues

I. General Support for Rulemaking

Several commenters expressed support for particular revisions, as well as the rulemaking in general. Many of these comments, which were received from individuals as well as organizations in the veteran community, expressed appreciation for VA's action in updating the rating schedule for musculoskeletal disabilities. VA appreciates the time and effort expended by these commenters in reviewing the proposed rule and in submitting comments, as well as their support for this rulemaking.

II. Public Access

One commenter requested public access to the information developed by the musculoskeletal system workgroup. In the preamble to the proposed rule, VA explained that the workgroup, comprised of subject matter experts from VA, the Department of Defense, and medical academia, held two public forums in August 2010 and June 2012, discussing possible revisions to the musculoskeletal regulations. A transcript of this public forum and all related materials are on file and available for public inspection in the Office of Regulation Policy and Management. (Contact information for that office is noted in the ADDRESSES section of the proposed rule. 82 FR at

VA emphasizes that the workgroup did not participate in the deliberative rulemaking process; the workgroup discussed the general topic of the VASRD body system and provided feedback on the areas that were subject to advances since the last major revision of the body system. To this end, where changes to the scientific and/or medical nature of a given condition were made in the proposed rule, VA cited the published, publicly available source for these changes. Not only did this provide the public with access to the source for a given proposed change, it also confirmed that VA relied upon peerreviewed scientific and medical information to support a given change. While similar information may have been presented by a workgroup member, VA relied upon the published document(s) as the primary source for a change and included such sources in the administrative record for this rulemaking. VA did not propose scientific and/or medical changes to the

VASRD in the absence of publicly available, peer-reviewed sources.

Accordingly, references in the proposed rule to the workgroup serve as an explanatory background and introduction to the VASRD rewrite project; the changes made by this rulemaking are not a reflection of the workgroup or any workgroup member. All changes based on scientific and/or medical information are a reflection of cited, published materials which are available to the public. VA has made deliberative materials available (via citation in the rulemaking) and is providing access to materials from the public forum for public inspection at the Office of Regulation Policy and Management.

III. Technical Corrections

On review, the current rating schedule refers evaluations of inactive tuberculosis of the bones and joints (DC 5001) to 38 CFR 4.88b; however, § 4.88b was redesignated to § 4.88c in 1994. Therefore, the final rule simply corrects this reference.

In addition, the final rule revises the subheading for DCs 5051 to 5056 to "Prosthetic Implants and Resurfacing," which the proposed rule noted in its regulatory text, but not in its preamble.

Also, DCs 5054 and 5055 have been reorganized to provide clarity to the applicability of the evaluation criteria. The 100 percent evaluation applies to both resurfacing and replacements. However, the 90, 70, 50, and 30 percent evaluations apply only to replacements. Therefore, the subheading referencing "replacement" in these DCs was relocated to the most appropriate location.

Lastly, VA made non-substantive edits to the parenthetical of DC 5242 and the proposed language for recurrent subluxation or instability under DC 5257

IV. Other Comments Unrelated to or Outside the Scope of This Rulemaking

VA received comments dealing with issues not directly related to proposed amendments to the rating schedule for musculoskeletal disabilities. One commenter suggested adding specified conditions to the list of presumptive disabilities for Former Prisoners of War (FPOW). Similarly, one commenter expressed concern over the impact of this rulemaking on the provisions for presumptive service connection for FPOWs in 38 CFR 3.309. Another commenter noted that the changes would assist in providing necessary treatment for the listed disabilities.

VA does not respond to these comments because they are either

unrelated to this rulemaking or beyond its scope.

Regulatory Flexibility Act

The Secretary hereby certifies that this final rule will not have a significant economic impact on a substantial number of small entities as they are defined in the Regulatory Flexibility Act, 5 U.S.C. 601-612. This final rule will not affect any small entities. The impact of this rulemaking results in cost savings to the VA's compensation and pension appropriations. There are no small entities involved, associated have an affilitation with VA's compensation and pension appropriations. Therefore, pursuant to 5 U.S.C. 605(b), the initial and final regulatory flexibility analysis requirements of 5 U.S.C. 603 and 604 do not apply.

Executive Orders 12866, 13563, and 13771

Executive Orders 12866 and 13563 direct agencies to assess the costs and benefits of available regulatory alternatives and, when regulation is necessary, to select regulatory approaches that maximize net benefits (including potential economic, environmental, public health and safety effects, and other advantages; distributive impacts; and equity). Executive Order 13563 (Improving Regulation and Regulatory Review) emphasizes the importance of quantifying both costs and benefits, reducing costs, harmonizing rules, and promoting flexibility. The Office of Information and Regulatory Affairs has determined that this rule is an economically significant regulatory action under Executive Order 12866.

VA's impact analysis can be found as a supporting document at www.regulations.gov, usually within 48 hours after the rulemaking document is published. Additionally, a copy of this rulemaking and its impact analysis are available on VA's website at www.va.gov/orpm/, by following the link for VA Regulations Published from FY 2004 Through Fiscal Year to Date. This rule is not subject to the requirements of E.O. 13771 because this rule results in no more than de minimis costs

Unfunded Mandates

The Unfunded Mandates Reform Act of 1995 requires, at 2 U.S.C. 1532, that agencies prepare an assessment of anticipated costs and benefits before issuing any rule that may result in the expenditure by State, local, and tribal governments, in the aggregate, or by the private sector, of \$100 million or more (adjusted annually for inflation) in any

one year. This final rule will have no such effect on State, local, and tribal governments, or on the private sector.

Paperwork Reduction Act

This final rule contains no provisions constituting a collection of information under the Paperwork Reduction Act of 1995 (44 U.S.C. 3501–3521).

Catalog of Federal Domestic Assistance

The Catalog of Federal Domestic Assistance program numbers and titles for this rule are 64.013, Veterans Prosthetic Appliances; 64.104, Pension for Non-Service-Connected Disability for Veterans; 64.109, Veterans Compensation for Service-Connected Disability; and 64.110, Veterans Dependency and Indemnity Compensation for Service-Connected Death.

Congressional Review Act

This regulatory action is a major rule under the Congressional Review Act, 5 U.S.C. 801–808, because it may result in an annual effect on the economy of \$100 million or more. In accordance with 5 U.S.C. 801(a)(1), VA will submit to the Comptroller General and to Congress a copy of this regulatory action and VA's Regulatory Impact Analysis.

List of Subjects in 38 CFR Part 4

Disability benefits, Pensions, Veterans.

Signing Authority

The Secretary of Veterans Affairs, or designee, approved this document and authorized the undersigned to sign and submit the document to the Office of the Federal Register for publication electronically as an official document of the Department of Veterans Affairs. Pamela Powers, Chief of Staff, Department of Veterans Affairs, approved this document on April 1, 2020, for publication.

Dated: November 13, 2020.

Jeffrey M. Martin,

Assistant Director, Office of Regulation Policy & Management, Office of the Secretary, Department of Veterans Affairs.

For the reasons set out in the preamble, VA amends 38 CFR part 4, subpart B, as follows:

PART 4—SCHEDULE FOR RATING DISABILITIES

Subpart B—Disability Ratings

■ 1. The authority citation for part 4, subpart B continues to read as follows:

Authority: 38 U.S.C. 1155, unless otherwise noted.

- 2. Amend § 4.71a by:
- **a** a. Revising diagnostic codes 5001, 5002, 5003, 5009–5015, 5018, 5020, 5022, 5023, 5024, 5054, 5055, 5120, 5160, 5170, 5201, 5202, 5242, 5243, 5255, 5257, 5262, and 5271;
- b. Removing the notes following diagnostic codes 5053 and 5056 and the note at the end of the table entitled "Prosthetic Implants and Resurfacing";
- c. Adding notes following diagnostic code 5024;
- d. Revising the heading "Prosthetic Implants" to read "Prosthetic Implants and Resurfacing" and adding notes 1 through 6 to it; and
- e. Adding the diagnostic code 5244 to the table entitled "The Spine" and the diagnostic code 5285 to the table entitled "The Foot".

The revisions and additions read as follows:

100

§ 4.71a Schedule of ratings—musculoskeletal system.

ACUTE, SUBACUTE, OR CHRONIC DISEASES

	Rating
* * * * * * *	*
 5001 Bones and joints, tuberculosis of, active or inactive:	100 100 60 40 20
* * * * * *	*
5009 Other specified forms of arthropathy (excluding gout). Note (1): Other specified forms of arthropathy include, but are not limited to, Charcot neuropathic, hypertrophic, crystalline, and other autoimmune arthropathies. Note (2): With the types of arthritis, diagnostic codes 5004 through 5009, rate the acute phase under diagnostic codes.	

Note (2): With the types of arthritis, diagnostic codes 5004 through 5009, rate the acute phase under diagnostic code 5002; rate any chronic residuals under diagnostic code 5003.

5010 Post-traumatic arthritis: Rate as limitation of motion, dislocation, or other specified instability under the affected joint. If there are 2 or more joints affected, each rating shall be combined in accordance with § 4.25.

Note: The 100 percent rating will be continued for 1 year following the cessation of surgical, X-ray, antineoplastic chemotherapy or other prescribed therapeutic procedure. If there has been no local recurrence or metastases, rate based on residuals.

5013 Osteoporosis, residuals of.

		A	ACUTE, SUBACUTE, O	OR CHRONIC DIS	SEASES—Continued		
							Rating
	Osteomalacia, re						
5015	Bones, neoplasm	n, benign.					
5018	* [Removed]	*	*	*	*	*	*
5016	[Hellioved]						
5020	* [Removed]	*	*	*	*	*	*
5022	[Removed]	: * :					
	Tenosynovitis, te	ndinitis, tendinos through 5024:		under diagnostic d	codes 5013 through 502	4 as degenerative ar-	
	thritis, based on I	imitation of motion	on of affected parts.				
	*	*	*	*	*	*	*
			PROSTHETIC I	MPLANTS AND F	RESURFACING		
						Ratin	g
						 Major	Minor
Note	(1): When an eval	uation is assigne	ed for joint resurfacing o	r the prosthetic rer	placement of a joint unde	ar di-	
agr	ostic codes 5051-	-5056, an additio	nal rating under § 4.71a	may not also be a	assigned for that joint, ur	nless	
	erwise directed. (2): Only evaluate	a revision proc	edure in the same man	ner as the original	l procedure under diagn	ostic	
cod	lès 5051-5056 if a	II the original cor	mponents are replaced.	ŭ			
Note plac	(3): The term "pro cement of the nam	sthetic replacem	ent" in diagnostic code: er. in DC 5054, "prosthe	s 5051–5053 and (etic replacement" r	5055–5056 means a tota means a total replaceme	al re- nt of	
the	head of the femur	or of the acetab	ulum.				
Note	(4): The 100 percent of the 1-month to	cent rating for 1	year following implant ned under§4.30 followi	tation of prosthesi	s will commence after i	nitial	
Note	(5): The 100 perc	ent rating for 4	months following impla	ntation of prosthes	sis or resurfacing under	DCs	
	54 and 5055 will co pital discharge.	ommence after i	nitial grant of the 1-mor	nth total rating ass	signed under § 4.30 follo	wing	
Note				the 100 percent ra	ating period the earliest	date	
	*	*	*	*	*	*	*
5054	Hip, resurfacing	or replacement (prosthesis):	-1			40
			of prostnesis or resurtation of the femur or of the a				10
•	Following impla	antation of prost	hesis with painful motion		uch as to require the us	se of	
		re recidual week		of mation following	implementation of proofboo		19
					implantation of prosthes		7 5
	Minimum evalu	ation, total repla	cement only				3
					cing under diagnostic c	odes	
	Knee, resurfacing		um evaluation for resurf t (prosthesis):	acing.			
F	or 4 months follow	ing implantation	of prosthesis or resurfa	cing			10
F	rosthetic replacem With chronic re			ion or weakness in	the affected extremity.		6
	With intermedia	ate degrees of re	esidual weakness, pain		tion rate by analogy to		· ·
		5256, 5261, or 5					3
	At the conclusion	n of the 100 pe		, evaluate resurfa	cing under diagnostic c		J
020	*	*	*	*	*	*	*
			A				
			AMPUTAT	IONS: UPPER EX	KTREMITY		
						Ratin	g
						Major	Minor

Arm, amputation of:
5120 Complete amputation, upper extremity:
Forequarter amputation (involving complete removal of the humerus along with any portion of the scapula, clavicle, and/or ribs)

1100

AMPUTATIONS: UPPER EXTREMITY—Continued Rating Major Minor 190 190 Disarticulation (involving complete removal of the humerus only) **AMPUTATIONS: LOWER EXTREMITY** Rating Thigh, amputation of: 5160 Complete amputation, lower extremity: Trans-pelvic amputation (involving complete removal of the femur and intrinsic pelvic musculature along with any portion of the pelvic bones) ² 100 Disarticulation (involving complete removal of the femur and intrinsic pelvic musculature only) 290 Note: Separately evaluate residuals involving other body systems (e.g., bowel impairment, bladder impairment) under the appropriate diagnostic code. 5170 Toes, all, amputation of, without metatarsal loss or transmetatarsal, amputation of, with up to half of metatarsal loss 30 THE SHOULDER AND ARM Rating

	Пашт	9
	Major	Minor
* * * * *	*	*
5201 Arm, limitation of motion of:		
Flexion and/or abduction limited to 25° from side	40	30
Midway between side and shoulder level (flexion and/or abduction limited to 45°)	30	20
At shoulder level (flexion and/or abduction limited to 90°)	20	20
5202 Humerus, other impairment of:		
	80	70
Loss of head of (flail shoulder)	60	50
Fibrous union of	50	40
Recurrent dislocation of at scapulohumeral joint:		
With frequent episodes and guarding of all arm movements	30	20
With infrequent episodes and guarding of movement only at shoulder level (flexion and/or abduction		
at 90 °)	20	20
Malunion of:		
Marked deformity	30	20
Moderate deformity	20	20
•		
* * * * *	*	*

THE SPINE

5242 Degenerative arthritis, degenerative disc disease other than intervertebral disc syndrome (also, see either DC 5003 or 5010)

5243 Intervertebral disc syndrome: Assign this diagnostic code only when there is disc herniation with compression and/or irritation of the adjacent nerve root; assign diagnostic code 5242 for all other disc diagnoses.

5244 Traumatic paralysis, complete:

Paraplegia: Rate under diagnostic code 5110.

Quadriplegia: Rate separately under diagnostic codes 5109 and 5110 and combine evaluations in accordance with § 4.25. **Note:** If traumatic paralysis does not cause loss of use of both hands or both feet, it is incomplete paralysis. Evaluate residuals of incomplete traumatic paralysis under the appropriate diagnostic code (*e.g.*, § 4.124a, Diseases of the Peripheral Nerves).

					Rating
* *	*	*	*	*	*
	Т	HE HIP AND THI	GH		
					Rating
* * 55 Femur, impairment of:	*	*	*	*	*
Fracture of shaft or anatomical neck of:					
With nonunion, with loose motion (spir					
With nonunion, without loose motion, we Fracture of surgical neck of, with false					
Malunion of:	, jonit				
Evaluate under diagnostic codes 5256 in the highest evaluation.	6, 5257, 5260, c	or 5261 for the kne	e, or 5250–5254 for t	he hip, whichever results	
* *	*	*	*	*	*
	т	HE KNEE AND L	FG		
	·				Rating
					9
*	*	*	*	*	*
Knee, other impairment of:					
Recurrent subluxation or instability: Unrepaired or failed repair of complet	to ligament toar	caucina parcietan	t instability and a me	udical provider prescribes	
both an assistive device (e.g., cane)					
One of the following:	(,	•		
(a) Sprain, incomplete ligament te					
ical provider prescribes a brace	and/or assistive	e device (<i>e.g.,</i> can	e(s), crutch(es), walke	r) for ambulation.	
ical provider prescribes a brace (b) Unrepaired or failed repair of a scribes either an assistive device	and/or assistive complete ligame ce (e.g., cane(s)	e device (<i>e.g.,</i> cand ent tear causing pe , crutch(es), walke	e(s), crutch(es), walke ersistent instability, and r) or bracing for ambu	r) for ambulation. d a medical provider pre- lation	
ical provider prescribes a brace (b) Unrepaired or failed repair of of scribes either an assistive device Sprain, incomplete ligament tear, or of the scribes are prescribed as a scribe of the scribe	and/or assistive complete ligame ce (e.g., cane(s) complete ligame	e device (e.g., cand ent tear causing pe , crutch(es), walke ent tear (repaired,	e(s), crutch(es), walke ersistent instability, and r) or bracing for ambu unrepaired, or failed i	r) for ambulation. d a medical provider pre- lation repair) causing persistent	
ical provider prescribes a brace (b) Unrepaired or failed repair of of scribes either an assistive device Sprain, incomplete ligament tear, or constability, without a prescription from bracing for ambulation	and/or assistive complete ligame ce (e.g., cane(s) complete ligame m a medical pro	e device (e.g., cane ent tear causing pe , crutch(es), walke ent tear (repaired, ovider for an assist	e(s), crutch(es), walke ersistent instability, and r) or bracing for ambu unrepaired, or failed in ive device (e.g., cane	r) for ambulation. d a medical provider pre- lation repair) causing persistent (s), crutch(es), walker) or	
ical provider prescribes a brace (b) Unrepaired or failed repair of a scribes either an assistive device. Sprain, incomplete ligament tear, or a instability, without a prescription from bracing for ambulation	and/or assistive complete ligame ce (e.g., cane(s) complete ligame m a medical pro	e device (e.g., cane ent tear causing pe , crutch(es), walke ent tear (repaired, ovider for an assist	e(s), crutch(es), walke ersistent instability, and r) or bracing for ambu unrepaired, or failed in ive device (e.g., cane	r) for ambulation. d a medical provider pre- lation repair) causing persistent (s), crutch(es), walker) or	
ical provider prescribes a brace (b) Unrepaired or failed repair of of scribes either an assistive device. Sprain, incomplete ligament tear, or constability, without a prescription from bracing for ambulation	and/or assistive complete ligame ce (e.g., cane(s) complete ligame m a medical pro matellofemoral col	e device (e.g., canent tear causing pe, crutch(es), walke ent tear (repaired, ovider for an assist	e(s), crutch(es), walke ersistent instability, and r) or bracing for ambu unrepaired, or failed in ive device (e.g., cane 	r) for ambulation. d a medical provider pre- lation repair) causing persistent (s), crutch(es), walker) or	
ical provider prescribes a brace (b) Unrepaired or failed repair of of scribes either an assistive device Sprain, incomplete ligament tear, or constant instability, without a prescription from bracing for ambulation	and/or assistive complete ligame ce (e.g., cane(s) complete ligame m a medical promoter atellofemoral color a brace and e	e device (e.g., canent tear causing per, crutch(es), walke ent tear (repaired, ovider for an assist	e(s), crutch(es), walke ersistent instability, and r) or bracing for ambu unrepaired, or failed in ive device (e.g., cane	r) for ambulation. d a medical provider pre- lation epair) causing persistent (s), crutch(es), walker) or ical repair that requires a	
ical provider prescribes a brace (b) Unrepaired or failed repair of of scribes either an assistive device. Sprain, incomplete ligament tear, or continuous instability, without a prescription from the bracing for ambulation	and/or assistive complete ligame ce (e.g., cane(s) complete ligame m a medical promoter a brace and eatellofemoral color a brace and eatellofemoral color one of the follows:	e device (e.g., canent tear causing per, crutch(es), walke ent tear (repaired, ovider for an assist entire with recurre either a cane or a verifice with recurre lowing: A brace, canent tear cane or a verifice with recurre lowing: A brace, canent tear canent tear tear tear tear tear tear tear tea	e(s), crutch(es), walke ersistent instability, and or bracing for ambuunrepaired, or failed rive device (e.g., cane tinstability after surgualker tinstability after surgual, or walker tinstability after surgual.	r) for ambulation. d a medical provider pre- lation epair) causing persistent (s), crutch(es), walker) or ical repair that requires a ical repair that requires a	
ical provider prescribes a brace (b) Unrepaired or failed repair of of scribes either an assistive device. Sprain, incomplete ligament tear, or constability, without a prescription from bracing for ambulation	and/or assistive complete ligame ce (e.g., cane(s) complete ligame m a medical promatellofemoral color a brace and eatellofemoral color one of the follatellofemoral color atellofemoral color one of the follatellofemoral color atellofemoral color	e device (e.g., canent tear causing per, crutch(es), walke ent tear (repaired, ovider for an assist entered with recurre either a cane or a variety mplex with recurre lowing: A brace, camplex with recurred with r	e(s), crutch(es), walke ersistent instability, and or bracing for ambuunrepaired, or failed rive device (e.g., cane and instability after surgivalker after	r) for ambulation. d a medical provider pre- lation epair) causing persistent (s), crutch(es), walker) or ical repair that requires a ical repair that requires a without history of surgical	
ical provider prescribes a brace (b) Unrepaired or failed repair of of scribes either an assistive device. Sprain, incomplete ligament tear, or constability, without a prescription from bracing for ambulation	and/or assistive complete ligame ce (e.g., cane(s) complete ligame m a medical promiser a brace and corral brace and catellofemoral corrone of the folloatellofemoral corrigition from a m	e device (e.g., canent tear causing per, crutch(es), walke ent tear (repaired, ovider for an assist ent tear a cane or a variety and the country and the count	e(s), crutch(es), walke ersistent instability, and or bracing for ambuunrepaired, or failed rive device (e.g., cane)	r) for ambulation. d a medical provider pre- lation epair) causing persistent (s), crutch(es), walker) or ical repair that requires a ical repair that requires a without history of surgical ker	
ical provider prescribes a brace (b) Unrepaired or failed repair of of scribes either an assistive device. Sprain, incomplete ligament tear, or constability, without a prescription from bracing for ambulation	and/or assistive complete ligame ce (e.g., cane(s) complete ligame m a medical promatellofemoral color a brace and eatellofemoral color one of the foll atellofemoral coordinatellofemoral coordinatellofemoral coordinatellofemoral complete	e device (e.g., canent tear causing per, crutch(es), walke ent tear (repaired, ovider for an assist ent tear a cane or a warmplex with recurre owing: A brace, camplex with recurre nedical provider for ex consists of the center of the canend tear of the canend tear of the center of	e(s), crutch(es), walke ersistent instability, and or bracing for ambuunrepaired, or failed nive device (e.g., cane)	r) for ambulation. d a medical provider pre- lation epair) causing persistent (s), crutch(es), walker) or ical repair that requires a ical repair that requires a without history of surgical ker e patella, and the patellar	
ical provider prescribes a brace (b) Unrepaired or failed repair of of scribes either an assistive device. Sprain, incomplete ligament tear, or constability, without a prescription from bracing for ambulation	and/or assistive complete ligame ce (e.g., cane(s) complete ligame m a medical promatellofemoral cor a brace and eatellofemoral cor one of the follatellofemoral corription from a mofemoral complete to involve repair surgical repair	e device (e.g., canent tear causing per, crutch(es), walke ent tear (repaired, ovider for an assist ent tear a cane or a variety with recurre owing: A brace, camplex with recurre nedical provider for ex consists of the care of one or more p	e(s), crutch(es), walke ersistent instability, and or bracing for ambuunrepaired, or failed nive device (e.g., cane)	r) for ambulation. d a medical provider pre- lation repair) causing persistent (s), crutch(es), walker) or ical repair that requires a ical repair that requires a without history of surgical ker re patella, and the patellar	
ical provider prescribes a brace (b) Unrepaired or failed repair of of scribes either an assistive device. Sprain, incomplete ligament tear, or constability, without a prescription from bracing for ambulation	and/or assistive complete ligame ce (e.g., cane(s) complete ligame m a medical promatellofemoral cor a brace and eatellofemoral cor one of the follatellofemoral corription from a mofemoral complete to involve repair surgical repair	e device (e.g., canent tear causing per, crutch(es), walke ent tear (repaired, ovider for an assist ent tear a cane or a variety with recurre owing: A brace, camplex with recurre nedical provider for ex consists of the care of one or more p	e(s), crutch(es), walke ersistent instability, and or bracing for ambuunrepaired, or failed nive device (e.g., cane)	r) for ambulation. d a medical provider pre- lation repair) causing persistent (s), crutch(es), walker) or ical repair that requires a ical repair that requires a without history of surgical ker re patella, and the patellar	*
ical provider prescribes a brace (b) Unrepaired or failed repair of of scribes either an assistive device. Sprain, incomplete ligament tear, or consist in instability, without a prescription from bracing for ambulation	and/or assistive complete ligame ce (e.g., cane(s) complete ligame m a medical promatellofemoral cor a brace and eatellofemoral cor one of the follatellofemoral corription from a mofemoral complete to involve repair surgical repair	e device (e.g., canent tear causing per, crutch(es), walke ent tear (repaired, ovider for an assist ent). Implex with recurre either a cane or a warmlex with recurre lowing: A brace, camplex with recurre ledical provider for ex consists of the car of one or more perfor patellar instability.	e(s), crutch(es), walke ersistent instability, and or bracing for ambuunrepaired, or failed nive device (e.g., cane)	r) for ambulation. d a medical provider pre- lation repair) causing persistent (s), crutch(es), walker) or ical repair that requires a ical repair that requires a without history of surgical ker re patella, and the patellar	*
ical provider prescribes a brace (b) Unrepaired or failed repair of of scribes either an assistive device. Sprain, incomplete ligament tear, or constant instability, without a prescription from bracing for ambulation	and/or assistive complete ligame ce (e.g., cane(s) complete ligame m a medical produce and control at the control of the following at the control of the co	e device (e.g., canent tear causing per, crutch(es), walke and tear (repaired, ovider for an assist and tear (repaired), and tear (repaired), and tear (repaired), and tear (repaired) and	e(s), crutch(es), walke ersistent instability, and or bracing for ambuunrepaired, or failed rive device (e.g., cane). Int instability after surgualker	r) for ambulation. d a medical provider pre- lation epair) causing persistent (s), crutch(es), walker) or ical repair that requires a ical repair that requires a without history of surgical ker patella, and the patellar ents that contribute to the limited to, arthroscopy to	*
ical provider prescribes a brace (b) Unrepaired or failed repair of of scribes either an assistive device. Sprain, incomplete ligament tear, or consists in instability, without a prescription from bracing for ambulation	and/or assistive complete ligame ce (e.g., cane(s) complete ligame m a medical produce and control at the control of the following at the control of the co	e device (e.g., canent tear causing per, crutch(es), walke and tear (repaired, ovider for an assist and tear (repaired), and tear (repaired). Implex with recurrer and the recurrence with recurrer and the recurrence and th	e(s), crutch(es), walke ersistent instability, and or bracing for ambuunrepaired, or failed rive device (e.g., cane). Int instability after surguals after surguals are or walker	r) for ambulation. d a medical provider pre- lation epair) causing persistent (s), crutch(es), walker) or ical repair that requires a ical repair that requires a without history of surgical ker e patella, and the patellar ents that contribute to the limited to, arthroscopy to	*
ical provider prescribes a brace (b) Unrepaired or failed repair of of scribes either an assistive device. Sprain, incomplete ligament tear, or constability, without a prescription from bracing for ambulation	and/or assistive complete ligame ce (e.g., cane(s) complete ligame m a medical produce and control at the control of the following at the control of the co	e device (e.g., canent tear causing per, crutch(es), walke and tear (repaired, ovider for an assist and tear (repaired), and tear (repaired). Implex with recurrer and the recurrence with recurrer and the recurrence and th	e(s), crutch(es), walke ersistent instability, and or bracing for ambuunrepaired, or failed rive device (e.g., cane). Int instability after surguals after surguals are or walker	r) for ambulation. d a medical provider pre- lation epair) causing persistent (s), crutch(es), walker) or ical repair that requires a ical repair that requires a without history of surgical ker e patella, and the patellar ents that contribute to the limited to, arthroscopy to	*
ical provider prescribes a brace (b) Unrepaired or failed repair of of scribes either an assistive device. Sprain, incomplete ligament tear, or constability, without a prescription from bracing for ambulation	and/or assistive complete ligame ce (e.g., cane(s) complete ligame m a medical promatellofemoral color a brace and eatellofemoral color one of the follatellofemoral corption from a mofemoral complete surgical repair surgical repair surgical repair surgical surgica	e device (e.g., canent tear causing per, crutch(es), walke ent tear (repaired, ovider for an assist ent tear (repaired, ovider for an assist ent tear (repaired, ovider for an assist ent tear cane or a warmplex with recurrence owing: A brace, camplex with recurrence ent can be consisted for ex consists of the corresponding	e(s), crutch(es), walke ersistent instability, and or bracing for ambuunrepaired, or failed nive device (e.g., cane). Int instability after surgivalker	r) for ambulation. d a medical provider pre- lation	*
ical provider prescribes a brace (b) Unrepaired or failed repair of of scribes either an assistive device. Sprain, incomplete ligament tear, or constability, without a prescription from bracing for ambulation	and/or assistive complete ligame ce (e.g., cane(s) complete ligame m a medical promise at the lost of the following and the lost of the following at the lost of th	e device (e.g., canent tear causing per, crutch(es), walke ent tear (repaired, ovider for an assist ent tear a cane or a warmlex with recurrence owing: A brace, camplex with recurrence can a consist of the correct consists of the correct of the c	e(s), crutch(es), walke ersistent instability, and or bracing for ambuunrepaired, or failed nive device (e.g., cane). Int instability after surgivalker	r) for ambulation. d a medical provider pre- lation epair) causing persistent (s), crutch(es), walker) or ical repair that requires a ical repair that requires a without history of surgical ker e patella, and the patellar ents that contribute to the limited to, arthroscopy to * the ankle, whichever re- d either shoe orthotics or	*
ical provider prescribes a brace (b) Unrepaired or failed repair of of scribes either an assistive device. Sprain, incomplete ligament tear, or consists in incomplete ligament tear. Patellar instability: A diagnosed condition involving the parescription by a medical provider for the A diagnosed condition involving the parepair) that does not require a prescription by a medical provider for the tendon. Note (1): For patellar instability, the patellot tendon. Note (2): A surgical procedure that does not underlying instability shall not qualify as remove loose bodies and joint aspiration * * S2 Tibia and fibula, impairment of: Nonunion of, with loose motion, requiring by Malunion of: Evaluate under diagnostic codes 5256 sults in the highest evaluation. Medial tibial stress syndrome (MTSS), or some Requiring treatment for no less than 1 other conservative treatment, both local stress in the patellocal streament of the conservative treatment, both local streament is conservative treatment, both local streament is conservative treatment.	and/or assistive complete ligame ce (e.g., cane(s) complete ligame m a medical produce of a brace and eatellofemoral coor one of the folloatellofemoral complete ligame of the folloatellofemoral complete complet	e device (e.g., canent tear causing per, crutch(es), walke not tear (repaired, ovider for an assist male with recurre owing: A brace, camplex with recurre owing: A brace, camplex with recurre ex consists of the cor of one or more properties of the cor o	e(s), crutch(es), walke ersistent instability, and or bracing for ambuunrepaired, or failed rive device (e.g., cane). Int instability after surgualker	r) for ambulation. d a medical provider pre- lation epair) causing persistent (s), crutch(es), walker) or ical repair that requires a ical repair that requires a without history of surgical ker e patella, and the patellar ents that contribute to the limited to, arthroscopy to	*
ical provider prescribes a brace (b) Unrepaired or failed repair of of scribes either an assistive device. Sprain, incomplete ligament tear, or constability, without a prescription from bracing for ambulation	and/or assistive complete ligame ce (e.g., cane(s) complete ligame m a medical produce for a brace and eatellofemoral coor one of the follatellofemoral coordinatellofemoral coordinatellofemoral coordinatellofemoral coordinatellofemoral complete to tinvolve repair surgical s	e device (e.g., canent tear causing per, crutch(es), walke and tear (repaired, ovider for an assist and tear and tear or a variety of the country of the cou	e(s), crutch(es), walke ersistent instability, and or bracing for ambuunrepaired, or failed rive device (e.g., cane). Int instability after surgivaler	r) for ambulation. d a medical provider pre- lation epair) causing persistent (s), crutch(es), walker) or ical repair that requires a ical repair that requires a without history of surgical ker e patella, and the patellar ents that contribute to the limited to, arthroscopy to * The ankle, whichever re- d either shoe orthotics or d either shoe orthotics or	*
ical provider prescribes a brace (b) Unrepaired or failed repair of of scribes either an assistive device. Sprain, incomplete ligament tear, or constability, without a prescription from bracing for ambulation	and/or assistive complete ligame ce (e.g., cane(s) complete ligame m a medical property of a brace and eatellofemoral coor one of the follatellofemoral coordinatellofemoral coordinatellofemoral coordinatellofemoral coordinatellofemoral complete to the follatellofemoral coordinatellofemoral complete to the follatellofemoral complete to the following that involve repairs surgical repair surgical repair surgical repair to the following that is the following tha	e device (e.g., canent tear causing per, crutch(es), walke not tear (repaired, ovider for an assist male with recurre either a cane or a variety with recurre towing: A brace, camplex with recurre edical provider for ex consists of the corresponding of the corre	e(s), crutch(es), walke ersistent instability, and or bracing for ambuunrepaired, or failed rive device (e.g., cane). Int instability after surgivaler	r) for ambulation. d a medical provider pre- lation epair) causing persistent (s), crutch(es), walker) or ical repair that requires a ical repair that requires a without history of surgical ker e patella, and the patellar ents that contribute to the limited to, arthroscopy to * The ankle, whichever re- d either shoe orthotics or one orthotics or other con-	*
ical provider prescribes a brace (b) Unrepaired or failed repair of of scribes either an assistive device. Sprain, incomplete ligament tear, or constability, without a prescription from bracing for ambulation	and/or assistive complete ligame ce (e.g., cane(s) complete ligame m a medical property of a brace and eatellofemoral coor one of the follatellofemoral coordinatellofemoral coordinatellofemoral coordinatellofemoral coordinatellofemoral complete to tinvolve repairs surgical repair surgical	e device (e.g., canent tear causing per, crutch(es), walke not tear (repaired, ovider for an assist mplex with recurre either a cane or a variety and the complex with recurrence owing: A brace, camplex with recurrence of the complex of the complex consists of the comple	e(s), crutch(es), walke ersistent instability, and or bracing for ambuunrepaired, or failed rive device (e.g., cane). Int instability after surgivaler	r) for ambulation. d a medical provider pre- lation epair) causing persistent (s), crutch(es), walker) or ical repair that requires a ical repair that requires a without history of surgical ker e patella, and the patellar ents that contribute to the limited to, arthroscopy to * The ankle, whichever re- d either shoe orthotics or one orthotics or other con-	*

THE ANKLE Rating 5271 Ankle, limited motion of: Marked (less than 5 degrees dorsiflexion or less than 10 degrees plantar flexion) 20 Moderate (less than 15 degrees dorsiflexion or less than 30 degrees plantar flexion) 10 THE FOOT Rating 5285 Plantar fasciitis: No relief from both non-surgical and surgical treatment, bilateral 30 No relief from both non-surgical and surgical treatment, unilateral 20 Otherwise, unilateral or bilateral 10 Note (1): With actual loss of use of the foot, rate 40 percent. Note (2): If a veteran has been recommended for surgical intervention, but is not a surgical candidate, evaluate under the 20 percent or 30 percent criteria, whichever is applicable. THE SKULL Rating

(Authority: 38 U.S.C. 1155)

0.4. 10.5-01

- 3. Amend § 4.73 by:
- a. Designating the introductory note as Note (1) and revising it;
- b. Adding introductory note (2); and
- c. Adding add diagnostic codes 5330 and 5331 to the table entitled "Miscellaneous".

The revising and additions read as follows:

§ 4.73 Schedule of ratings—muscle injuries.

Note (1): When evaluating any claim involving muscle injuries resulting in loss of use of any extremity or loss of use of both buttocks (diagnostic code 5317, Muscle Group XVII), refer to

§ 3.350 of this chapter to determine whether the veteran may be entitled to special monthly compensation.

Note (2): Ratings of slight, moderate, moderately severe, or severe for diagnostic codes 5301 through 5323 will be determined based upon the criteria contained in § 4.56.

MISCELLANEOUS

Rating

5330 Rhabdomyolysis, residuals of:

Rate each affected muscle group separately and combine in accordance with §4.25.

Note: Separately evaluate any chronic renal complications within the appropriate body system.

5331 Compartment syndrome:

Rate each affected muscle group separately and combine in accordance with §4.25.

- 4. Amend appendix A to part 4 as follows:
- a. In § 4.71a, revise diagnostic codes 5001, 5002, 5003, 5012, 5024, 5051, 5052, 5053, 5054, 5055, 5056, 5243, 5255, and 5257;
- b. In § 4.71a, remove the diagnostic code 5235–5243;
- c. In § 4.71a, add in numerical order diagnostic codes 5009, 5010, 5011, 5013, 5014, 5015, 5018, 5020, 5022, 5023, 5120, 5160, 5170, 5201, 5202, 5235, 5236, 5237, 5238, 5239, 5240, 5241, 5242, 5244, 5262, 5271, and 5285; and
- d. In § 4.73, add an introduction note and diagnostic codes 5330 and 5331.

The revisions and additions read as follows:

Appendix A to Part 4—Table of Amendments and Effective Dates Since 1946

Sec.	Diagnostic code No.			
* *		* * *	*	*
4.71a	5001 5002 5003		1.	
* *		* * *	*	*
	5010 5011 5012 5013 5014 5015 5018 5020 5022 5023	Title, evaluation, note February 7, 2021. Title, criteria February 7, 2021. Title, criteria February 7, 2021. Criterion March 10, 1976; title, note February 7, 2021. Title February 7, 2021. Title February 7, 2021. Title February 7, 2021. Removed February 7, 2021. Removed November 30, 2020. Removed February 7, 2021. Title February 7, 2021. Title February 7, 2021. Criterion March 1, 1963; title, criteria February 7, 2021.		
* *		* * *	*	*
	5052			
* *		* *	*	*
	5120 5160	Title, criterion February 7, 2021. Title, criterion, note February 7, 2021.		
* *	5170	* * * * Title February 7, 2021.	*	*
* *		* * * Criterion February 7, 2021. Criterion February 7, 2021.	*	*
* *	5236 5237 5238 5239 5240 5241 5242	Replaces 5285–5295 September 26, 2003. Replaces 5285–5295 September 26, 2003. Replaces 5285–5295 September 26, 2003.	* 2021.	*
	5243 5244	2021. Added February 7, 2021.	iber 26, 2003; Title	rebruary 7,
* *	5255	* * Criterion July 6, 1950; criterion February 7, 2021.	*	*
* *	0200	* * *	*	*
. *	5257	Evaluation July 6, 1950; criterion and note February 7, 2021.	-	*
* *	5262	* * * * Criterion February 7, 2021.	*	*
* *	5271	* * Criterion February 7, 2021.	*	*
* *		* * * Added February 7, 2021.	*	*
	3203	* * *	*	*
4.73		Introduction Note criterion July 3, 1997; second Note added F	February 7, 2021.	^
* *		* * *	*	*
	5330 5331	Added February 7, 2021. Added February 7, 2021.		

	Sec.		Diagnostic code No.					
	*	*	*		*	*	*	*
follo		_	and		20, 5160, 5170), and 5242;	The revisions and a follows:	dditions read as
5009	Revise diagnostic , 5010, 5011, 501 , 5018, 5020, 502	12, 5013, 50	14, 533	Add diagn 0, and 533	ostic codes 52 1;	44, 5285,	Appendix B to Part 4- of Disabilities	—Numerical Inde
	Diagnostic code	No.						
			А		sculoskeletal S cute, or Chronic			
	*	*	*		*	*	*	*
					oost-traumatic an r than post-traum		ore joints, as an active pro	cess.
	*	*	*		*	*	*	*
			Other specified Post-traumatic		hropathy (exclud	ling gout).		
			Decompression					
			Osteoporosis, re		nt, primary or sec	condary.		
			Osteomalacia, r					
5015			Bones, neoplas	m, benign.				
	*	*	*		*	*	*	*
5018			[Removed]					
5020	*	*	[Removed]		*	*	*	*
3020			[Hemoved]					
	*	*	*		*	*	*	*
			[Removed]	rification				
					ndinosis or tendi	nopathy.		
5054	*	*	Hin recurfacing	or replacer	* nent (prosthesis)	*	*	*
					ement (prosthes			
	*	*	*		*	*	*	*
				Amputati	ions: Upper Ext	remity		
	amputation of:		Complete ampl	itation uppe	ar extremity			
3120		•••••	Complete ampe	itation, uppc	or extremity.			
	*	*	*		*	*	*	*
				Amputati	ions: Lower Ext	remity		
	, amputation of:		Complete ampu	ıtation, lowe	r extremity.			
	*	*	*		*	*	*	*
5170			Toes, all, ampu tarsal loss.	tation of, wit	thout metatarsal	loss or transme	etatarsal, amputation of, w	ith up to half of met
	*	*	*		*	*	*	*
					Spine			
	*	*			•	_	±	.
5242	*	*	Degenerative a	rthritis, dege	* enerative disc dis	* sease other tha	n intervertebral disc syndı	rome (also, see eith
			DC 5003 or 5				,	,
	*	*	*		*	*	*	*
5244			Traumatic naral	lucie comple	ato			

Diagnostic c	ode No.					
*	*	*	*	*	*	*
			The Foot			
*	*	*	*	*	*	*
285	Pla	ntar fasciitis.				
*	*	*	*	*	*	*
			MUSCLE INJURIES			
*	*	*	*	*	*	*
			Miscellaneous			
*	*	*	*	*	*	*
330 331	Rha	abdomyolysis, residua mpartment syndrome	als of.			
*	*	*	*	*	*	*

- 6. Amend appendix C to part 4 as
- a. Revising the entries for "Amputation" and "Arthritis";
 ■ b. Adding in alphabetical order an
- entry for "Arthropathy";
- c. Revising the entry for "Bones";
- d. Adding in alphabetical order entries for "compartment syndrome", "decompression illness", and "heterotopic ossification"
- e. Revising the entry for "Hip";

- f. Removing entries for "Hydrarthrosis, intermittent", and "Myositis ossificans"
- g. Revising entries for "Osteomalacia", "Osteoporosis, with joint manifestations", and "Paralysis";
- h. Removing entry for "Periostitis";
- i. Adding in alphabetical order an entry for "Plantar fasciitis";
- j. Revising entry for "Prosthetic implants";
- k. Adding in alphabetical order entries for "Rhabdomyolysis, residuals of" and "Spine: Degenerative arthritis, degenerative disc disease other than intervertebral disc syndrome";
- l. Removing entry for "Synovitis"; and m. Revising entry for "Tenosynovitis" The revisions and additions read as

Appendix C to Part 4—Alphabetical **Index of Disabilities**

Diagnostic code No. Amputation: Complete amputation, upper extremity 5120 Above insertion of deltoid 5121 Below insertion of deltoid 5122 Digits, five of one hand 5126 Digits, four of one hand: 5127 Thumb, index, long and ring Thumb, index, long and little 5128 Thumb, index, ring and little 5129 Thumb, long, ring and little 5130 Index, long, ring and little 5131 Digits, three of one hand:. Thumb, index and long 5132 Thumb, index and ring 5133 Thumb, index and little 5134 Thumb, long and ring 5135 Thumb, long and little 5136 Thumb, ring and little 5137 Index, long and ring 5138 Index, long and little 5139 Index, ring and little 5140 Long, ring and little 5141 Digits, two of one hand: Thumb and index 5142 Thumb and long 5143 5144 Thumb and little 5145 5146 Index and long

						Diagnostic code No.
Index and ring						514
						514
						514
						515 515
Single finger:						310
						515
Index finger						515
~. ~						515
						515 515
Forearm:						310
	pronator teres					512
Below insertion of p	pronator teres					512
Leg:						F.4.
						516 516
						516
						516
					alf of metatarsal loss	51
						51
						51° 51°
Thigh:	s, without mete	atarsar involvement				31.
	on, lower extre	emity				510
Upper third						51
Middle or lower thin	ds					51
*	*	*	*	*	*	*
thritis:						
	n post-traumat	ic				50
						50
						50
						50
						50 [.] 500
_ ` ` ` `		• ,				500
						500
						50
thropathy						50
*	*	*	*	*	*	*
ones:						
Neoplasm, benign						50
						50
Shortening of the lower	extremity					52
*	*	*	*	*	*	*
empartment syndrome						53
*	*	*	*	*	*	*
compression illness						50
*	*	*	*	*	*	*
terotopic ossification						50
):						
Flail joint						52
•		•	•	*	•	
teomalacia, residuals of		,	•	•	•	50
ncomalacia, residuais Ul						30
*	*	*	*	*	*	*
teoporosis, residuals of .						50
*	*	*	*	*	*	*
aralysis:						60
						80
						52
						32
, , , , , , , , , , , , , , , , , , ,						
*	*	*	*	*	*	*

						Diagnostic code No.
*	*	*	*	*	*	*
Prosthetic implants:						5050
Ankle replaceme	nt					505
Elbow replaceme	ent					505
Hip, resurfacing						
Knee, resurfacin	g or replacement					505
Shoulder replace	ement					505
Wrist replaceme	nt					505
*	*	*	*	*	*	*
Rhabdomyolysis, res	iduals of					5330
*	*	*	*	*	*	*
Spine:						
	hritis, degenerative	disc disease other th	nan intervertebral disc	svndrome		524
9	, angemenant			,		
*	*	*	*	*	*	*
Tenosynovitis, tendin	itis, tendinosis or te	ndinopathy				502

[FR Doc. 2020–25450 Filed 11–27–20; 8:45 am] **BILLING CODE 8320–01–P**

FEDERAL COMMUNICATIONS COMMISSION

47 CFR Parts 1 and 90

[WP Docket No. 07-100; FCC 20-137; FRS 17146]

4.9 GHz Band

AGENCY: Federal Communications

Commission.

ACTION: Final rule.

SUMMARY: In March 2018, the Federal Communications Commission (Commission) released a Sixth Further Notice of Proposed Rulemaking (Sixth FNPRM) seeking comment on ways to stimulate expanded use of and investment in the 4.9 GHz (4940-4990 MHz) band, including allowing licensees the flexibility to engage in spectrum leasing and broadening existing eligibility requirements. On September 8, 2020, the Public Safety and Homeland Security Bureau and the Wireless Telecommunications Bureau issued a Public Notice freezing the 4.9 GHz band to stabilize it while the Commission considered changes to the 4.9 GHz band rules (Freeze Public Notice). In this document, the Commission adopts rules permitting one statewide 4.9 GHz band licensee per state, the State Lessor, to lease some or all of its spectrum rights to third parties-including commercial and public safety users—in those states that the Commission has not identified as a diverter of 911 fees. The Report and Order does not limit or modify the rights of any incumbent public safety

licensees. The new rules also eliminate the requirement that leased spectrum must be used to support public safety but requires lessees to adhere to the informal coordination requirements applicable to the band.

DATES: Effective December 30, 2020, except for § 90.1217, which is delayed. We will publish a document in the **Federal Register** announcing the effective date.

ADDRESSES: Federal Communications Commission, 45 L St. NE SW, Washington, DC 20554.

FOR FURTHER INFORMATION CONTACT:

Jonathan Markman of the Wireless Telecommunications Bureau, Mobility Division, at (202) 418–7090 or Jonathan.Markman@fcc.gov. For information regarding the PRA information collection requirements contained in this PRA, contact Cathy Williams, Office of Managing Director, at (202) 418–2918 or Cathy.Williams@fcc.gov.

SUPPLEMENTARY INFORMATION: This is a summary of the Commission's Report and Order in WP Docket No. 07-100, FCC 20–137 adopted September 30, 2020 and released October 02, 2020. The full text of the Report and Order, including all Appendices, is available by downloading the text from the Commission's website at https:// www.fcc.gov/document/fcc-expandsaccess-and-investment-49-ghz-band-0. Alternative formats are available for people with disabilities (braille, large print, electronic files, audio format), by sending an email to FCC504@fcc.gov or calling the Consumer and Governmental Affairs Bureau at (202) 418-0530 (voice), (202) 418-0432 (TTY).

The Commission will send a copy of this *Report* in a report to be sent to Congress and the Government Accountability Office pursuant to the Congressional Review Act, *see* 5 U.S.C. 801(a)(1)(A).

Final Regulatory Flexibility Analysis

The Regulatory Flexibility Act (RFA) requires that an agency prepare a regulatory flexibility analysis for notice and comment rulemakings, unless the agency certifies that "the rule will not, if promulgated, have a significant economic impact on a substantial number of small entities." Accordingly, the Commission has prepared a Final Regulatory Flexibility Analysis (FRFA) concerning the possible impact of the rule changes contained in this Report and Order on small entities. As required by the Regulatory Flexibility Act of 1980, as amended (RFA), an Initial Regulatory Flexibility Analysis (IRFA) was incorporated in the Sixth Further Notice of Proposed Rulemaking (Sixth FNPRM) released in March 2018 in this proceeding (83 FR 20011, May 7, 2018). The Commission sought written public comment on the proposals in the Sixth *FNPRM*, including comments on the IRFA. No comments were filed addressing the IRFA. This present Final Regulatory Flexibility Analysis (FRFA) conforms to the RFA.

Paperwork Reduction Act

The requirements in § 90.1217 constitute new or modified collections subject to the Paperwork Reduction Act of 1995 (PRA), Public Law 104–13. They will be submitted to the Office of Management and Budget (OMB) for review under section 3507(d) of the PRA. OMB, the general public, and