



American Orthotic & Prosthetic Association

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AOPA In Advance SmartBrief  
*Breaking News*  
August 23, 2018

**AOPA Headlines:**

[Spinal Bracing RFP Extended to September 20th](#)

[Jurisdiction D Releases Quarterly TPE Results: Diabetic Shoes](#)

[Two New Issues Proposed for RAC Audits](#)

[Thank you to our Supplier Plus Members](#)

[Jurisdiction D Releases Quarterly TPE Results](#)

[VA Roundtable Airs Broad Challenges to VA Proposed Rule](#)

[CMS Releases Proposed Rule on DMEPOS Competitive Bidding](#)

[DME MACs Retire Draft LCD and Policy Article for Lower Limb Prostheses](#)

[CMS' Lower Limb Prostheses Interagency Workgroup Releases a Consensus Statement](#)

[CMS Issues Instructions for DME MACs: Require O&P Documentation as Part of Medical Record](#)

[DME MACs Announce the Retirement of the Dear Physician Letter for Artificial Limbs](#)

[Exhibit at the AOPA National Assembly – Sept. 26-29 in Vancouver, Canada](#)

[RSVP for the 2018 Party With a Purpose](#)

[Upcoming Events](#)

<b>Spinal Bracing RFP Extended to September 20th</b>
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AOPA, under the auspices of its Orthotics 2020 program, circulated a request for proposals early in 2018 relating to 5 subject areas for original orthotic papers, with the original deadline for receipt of applications by April 30, 2018. Proposals have been received in all five of those categories. This notification is to announce a **re-opening of the opportunity to submit grant applications/extension of the deadline for applications as to the two revised RFPs on spinal bracing (available below). Extensions do not apply to any of the other research categories/RFPs which have already advanced into the decision stage.**

AOPA will now be accepting applications for grants as to the two revised spinal bracing RFPs, available below, provided that they are received no later than **September 20, 2018 at 11:59 pm.**

In all other respects, except for this extended deadline date, all terms stated in the original AOPA announcements remain intact and in effect as to these two revised spinal bracing RFPs.

Please review closely the terms of the RFPs. One problem we have noticed with responses is that there were multiple scoliosis applications/research protocols submitted under the RFP for back bracing. The category of back bracing is neither written nor intended to solicit submissions related to scoliosis. There may come a time when we will be looking for scoliosis papers, but this is NOT that time, so please do not commit your valuable time and energies to submitting scoliosis proposals in response to this back bracing RFP.

Back bracing is a very important and primary category in the orthotics profession. We have intentionally expanded the publication/notification /outreach to a broader audience of potential investigators toward submission of proposals as to these two revised spinal bracing RFPs. We will encourage and will welcome all high-quality submissions which are in accordance with the terms of the two RFPs.

[2018 Clinical or Comparative Effectiveness RFP on Back Bracing and Factors on Favorable Patient Outcomes \(FINAL 75\)](#)

[2018 Clinical or Comparative Effectiveness RFP on Back Bracing and Factors on Favorable Patient Outcomes \(S S-150\)](#)

Please contact AOPA staff at [ymazur@AOPAnet.org](mailto:ymazur@AOPAnet.org) with any questions.

### **Jurisdiction D Releases Quarterly TPE Results: Diabetic Shoes**

Noridian, the Durable Medical Equipment Administrative Contractor (DME MAC) for Jurisdiction D, recently published the quarterly results of their Target, Probe & Educate (TPE) audits for Therapeutic Shoes for Persons with Diabetes. The audit results are based on claims for code A5500 reviewed during April 2018-June 2018 and they show an overall claim potential improper payment rate is **33%**.

The top denial reasons for the TPE results are:

- Documentation does not support basic coverage criteria
- Detailed Written Order (DWO) was not received
- Documentation was not received in response to the Additional Documentation Request (ADR) letter
- Medical record documentation was not received

View the complete results and a full list of denial reasons [here](#).

### **Two New Issues Proposed for RAC Audits**

On August 7, 2018, the Centers for Medicare and Medicaid Services published two new proposed issues for review by Performant Recovery, the Recovery Audit Contractor (RAC) for DMEPOS, Home Health, and Hospice claims nationwide. The two issues that have been proposed for RAC review are custom fabricated knee orthoses described by L1844 and L1846 and DMEPOS services delivered to a Medicare beneficiary during a Hospice benefit period.

The proposed RAC review for custom fabricated knee orthoses is a complex review meaning that claims will be reviewed to ensure that the knee orthoses delivered were medically necessary and

meet the coverage guidelines outlined in the Medicare local coverage decision (LCD) and Policy Article for knee orthoses.

The second issue that has been proposed for RAC review is an automated review for DMEPOS services that were provided during a Medicare covered hospice benefit period. As hospice is considered a Medicare Part A benefit, claims submitted to Part B contractors, including DME MACs, are typically not covered as Part B services unless they are unrelated to the patient's terminal disease. When this is the case, providers are required to submit the claim with a "GW" modifier.

AOPA will continue to monitor both the CMS and RAC websites to confirm if and when these two issues are approved for RAC review.

Questions regarding this issue may be directed to Joe McTernan at [jmcternan@AOPAnet.org](mailto:jmcternan@AOPAnet.org) or Devon Bernard at [dbernard@AOPAnet.org](mailto:dbernard@AOPAnet.org).

### Thank you to our Supplier Plus Members



### Jurisdiction D Releases Quarterly TPE Results

Noridian, the Durable Medical Equipment Administrative Contractor (DME MAC) for Jurisdiction D, recently published the quarterly results of their Target, Probe & Educate (TPE) audits. The audits are based on claims audited/reviewed during April 2018-June 2018 and the results are as follows:

- Ankle Foot Orthoses/Knee Ankle Foot Orthoses (L4360, L4361, L4386 and L4387) had an overall claim potential improper payment rate is 34%. This is an increase over last quarter's overall claim potential improper payment rate of 19%.
- Knee Orthoses (L1810, L1812, L1830, L1832, L1833, L1843, L1845 and L1852) had an overall claim potential improper payment rate is 57%. This is a decrease from last quarter's overall claim potential improper payment rate of 77%.

- Spinal Orthoses (L0625, L0626, L0627, L0630, L0631, L0637, L0641, L0642, L0643, L0648 and L0650) had an overall claim potential improper payment rate is 51%. This is an increase over last quarter's overall claim potential improper payment rate of 34%.

The top and common denial reasons for all TPE results (in no order) are as follows:

- Documentation does not support basic coverage criteria.
- Detailed Written Order (DWO) is incomplete or missing elements.
- Documentation was not received in response to the Additional Documentation Request (ADR) letter.
- Claim is the same or similar to another claim on file.
- Documentation does not include verification that the equipment was lost, stolen or irreparably damaged in a specific incident.

View the complete results and a full list of denial reasons [here](#).

Questions? Contact Joe McTernan at [jmcternan@AOPAnet.org](mailto:jmcternan@AOPAnet.org) or Devon Bernard at [dbernard@AOPAnet.org](mailto:dbernard@AOPAnet.org)

<p><b>VA Roundtable Airs Broad Challenges to VA Proposed Rule That Would Limit Amputee Veterans' Choice between Private Sector and VA Care, Medical Necessity and Other Issues</b></p>
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On July 25, AOPA was among several entities represented at a Congressional Roundtable discussion conducted by the House Veteran's Affairs Health Subcommittee. The Roundtable was presided over by the Subcommittee Chair Rep. Neal Dunn, M.D. (R-FL) and Representative Julia Brownley (D-CA), with Rep. Annie Custer (D-NH) also attending a portion of the session. The list of participants is attached. Such roundtables are informal discussions with no sworn testimony and no recording of statements, to encourage a free exchange. The primary topic for the session was VA prosthetic care, with a distinct focus of the VA's proposed rule of October 16, 2017, including controversial provisions that would make VA the sole determinant of where a Veteran could receive O&P care (private sector vs. VA clinic)—thus it would limit the long-standing choice of provider afforded to many, if not all Veterans—and a tightening of the definition of medical necessity, along with brief discussion of other matters.

The session commenced with statements by Chairman Dunn, and Ranking Minority Member Brownley. Both opposed the two key changes, noted above, which the proposed rule would change, posing the question why now? And what is broken that needs fixing? The Chairman also specifically mentioned the need to assure training funds to aid in the education of new prosthetists for future generations of Veterans, specifically citing the NCOPE/AOPA supported work force study which targeted the need for a 60% increase in number of prosthetists to keep pace with retirements and expanding population (AOPA's lead advocate on that issue, Catriona Macdonald, was also present for the session).

AOPA emphasized that VA care needs to be focused in achieving both choice and quality of providers and their care for Veterans. Other topics discussed included brief discussions of the valuable clinical guidelines developed and being implemented by VA and possible expansion of patient outcomes efforts, as well as commentary by the VA OIG reps of purported excessive referral to private sector contractors for diabetic shoes and compression footwear in Southern Nevada, and a brief mention of an upcoming OIG report of VA coding practices which intersects well with the final paragraph on the AOPA one-pager, namely ways in which current coding

operate as an impediment to innovative technologies. There was also discussion supporting H.R. 2322, the Amputee Veterans Bill of Rights.

In addition to the VA Roundtable, AOPA reps—Executive Director Tom Fise, and Legislative Counsel Colin Roskey (Lincoln Policy Group) along with NAAOP’s first Intern Advocate, Nicole ver Kuilen met with Senator Tammy Duckworth (D-IL). The AOPA reps provided a briefing, update and request for support with topics discussed including: implementation of the prosthetist/orthotist notes provision; latest status of the CMS July 2015 draft LCD and the DME MAC Dear Physician Letter (both now withdrawn though deliberations on LCD-related issues remain active and volatile); S.1191—remaining sections including the section on reinforcing the ‘minimal self-adjustment’ definition that distinguishes OTS orthotics from custom-fitted and custom-fabricated orthotics; VA matters; and the MEDPAC report on orthotics.



Picture, Seated: Senator Tammy Duckworth (D-IL) and 2018 NAAOP Advocacy Intern Nicole ver Kuilen  
Standing AOPA Executive Director, Thomas Fise and AOPA Legislative Counsel Colin Roskey.

[Click here to view the one-pager which AOPA distributed to all participants which underscores O&P perspectives on several VA issues.](#)

### **CMS Releases Proposed Rule on DMEPOS Competitive Bidding**

On July 11, 2018, the Centers for Medicare and Medicaid Services (CMS) released a proposed rule that proposes several changes to the Medicare Durable Medical Equipment, Prosthetics, Orthotics, and Supplies (DMEPOS) competitive bidding program. In addition to the proposed changes to the competitive bidding program, the proposed rule also solicits comments regarding ways to improve the gap filling methodology that is used to establish Medicare fee schedule amounts for new items and technologies.

Two items of note in the proposed rule include the fact that off the shelf (OTS) orthoses are not mentioned as a potential product category for any envisioned revisions to the competitive bidding program, and that the current DMEPOS competitive bidding program will be effectively suspended when existing contracts expire at the end of 2018 and will remain suspended until such time as any new contracts are awarded under the rules that are being proposed by CMS. While OTS orthoses are eligible for inclusion in competitive bidding under the law that created the program, CMS has, to date, not made, or announced any decision to include them as a product category in the competitive bidding program. The announcement that the competitive bidding program will effectively be suspended when current contracts expire in 2018 is a significant development in that it acknowledges the negative impact that competitive bidding has had on patient access to

medically necessary DMEPOS services, especially in rural areas. While the competitive bidding program has not directly impacted the delivery of O&P care to date, we are hopeful that CMS' commitment to improving the program will help ensure that access to O&P care for Medicare beneficiaries remains available if the time comes when OTS orthoses might be included in competitive bidding in the future.

AOPA is currently reviewing the 368 page document and will prepare a full analysis for its members in the near future.

The CMS Fact Sheet regarding the proposed rule may be viewed by [clicking here](#).

The complete proposed rule may be viewed by [clicking here](#).

Questions regarding the proposed rule may be directed to Joe McTernan at [jmcternan@AOPAnet.org](mailto:jmcternan@AOPAnet.org) or Devon Bernard at [dbernard@AOPAnet.org](mailto:dbernard@AOPAnet.org).

### **DME MACs Retire Draft LCD and Policy Article for Lower Limb Prostheses**

Earlier this month, AOPA reported that the report of the Inter Agency Workgroup that was formed to provide a consensus statement to inform Medicare coverage of lower limb prostheses had been published. As part of the consensus statement, the Inter Agency Workgroup recommended, and CMS concurred, that the draft LCD and Policy Article for Lower Limb Prostheses, that was initially released on July 16, 2015, should be removed from the DME MAC websites and that the current LCD and Policy Article should remain in force for the immediate future.

On June 21, 2018, the DME MACs released a joint publication announcing the retirement of both the draft LCD and Policy Article. AOPA is continuing to evaluate the full consensus statement of the Inter Agency Workgroup and will provide comments in the near future.

The announcement of the draft LCD and Policy Article may be viewed by [clicking here](#).

Questions regarding this issue may be directed to Joe McTernan at [jmcternan@AOPAnet.org](mailto:jmcternan@AOPAnet.org) or Devon Bernard at [dbernard@AOPAnet.org](mailto:dbernard@AOPAnet.org).

### **CMS' Lower Limb Prostheses Interagency Workgroup Releases a Consensus Statement**

In 2016 the Center for Medicare & Medicaid Services (CMS) convened the Lower Limb Prostheses Interagency Workgroup in response to the comments received in regard to the 2015 Draft Local Coverage Determination (LCD) for Lower Limb Prostheses. The Workgroup's purpose was to "develop a consensus statement that informs Medicare policy by reviewing the available clinical evidence that defines best practices in the care of beneficiaries who require lower limb prostheses." The Workgroup has completed their review and released a consensus statement outlining their findings and suggestions.

Based on the findings and recommendations of the Workgroup CMS is taking the following immediate actions:

- Instructing the Durable Medical Equipment Medicare Administrative Contractors (DME MACs) to remove the Draft LCD
- Instructing the DME MACs that coverage for lower limb prostheses will remain under the current LCD, with no changes



- Future LCD changes must follow procedures set forth in the 21<sup>st</sup> Century Cures Act
- Considering creating a National Coverage Determination (NCD) to evaluate the use of microprocessor knees (MPK) in those individuals utilizing their prostheses as a limited community ambulator (K2 functional level)

The removal of the Draft LCD has been an aim of AOPA and its lobbying efforts, since it was introduced in 2015, and the official directions from CMS to remove it is a major victory. AOPA is currently reviewing the full findings of the Workgroup and the consensus document, and there are some areas/conclusions where AOPA will likely disagree and wish to provide additional comments. For example, it appears that the consensus document may have been written before the enactment in February, 2018 of Section 50402 of the Bipartisan Budget Act of 2018 which recognizes the legitimacy of orthotist's and prosthetist's notes in the medical record for the justification of medical necessity. Even in light of the [recent letter from the CMS Deputy Administrator & Director of Program Integrity](#) instructing the implementation of Section 50402, CMS has yet to revise the Program Integrity Manual to reflect the directions from the Deputy Director, the head of CMS' own Program Integrity Center, and the current status of the prosthetist's notes continues to be misstated.

In addition the recommendation that the potential for MPK devices for K2- limited community ambulators- be done by a NCD is an important step forward as it represent CMS taking back this authority from the DME MACs. LCDs are the province of the DME MACs, while a NCD is a strict ***Federal Register*** CMS-driven rulemaking process. CMS has consistently said that the only way they could take this matter out of the authority of the DME MACs would be to invoke a NCD. AOPA will also be submitting comments on the proposed NCD and potential for microprocessor knees to be used by K2 -limited community- ambulators in accordance with established guidelines.

[Click here to access the document.](#) Then click to go to the Lower Limb Prosthetic Workgroup Consensus Document link.

AOPA will keep you posted about any additional actions taken as a result of the Workgroup's final findings. Questions? Contact Joe McTernan at [jmcternan@AOPAnet.org](mailto:jmcternan@AOPAnet.org) or Devon Bernard at [dbernard@AOPAnet.org](mailto:dbernard@AOPAnet.org).

<p><b>CMS Issues Instructions for DME MACs to Immediately Implement Provisions that Require the Recognition of Orthotist and Prosthetist Clinical Documentation as Part of the Medical Record</b></p>
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AOPA and its lobbying team have been pressing CMS from all levels, most recently, including consultation with Trump Administration officials at the Office of Management and Budget, to formally implement the provisions of Section 50402 of the Bipartisan Budget Act of 2018 (Public Law 115-123) as it related to the prosthetist's and orthotist's clinical notes. AOPA was encouraged by the DME MAC notification 2-3 weeks ago where the DME MAC Medical Directors announced that they were 'retiring' the August 2011 "Dear Physician" letter on Lower Limb Prosthetics.

AOPA would like to share a recent letter from Alec Alexander, CMS' Director of Program Integrity which indicates that CMS "has issued instruction to the Durable Medical Equipment (DME) Medicare Administrative Contractors (MAC) to implement Section 50402 immediately." Section 50402 states:

*“(5) DOCUMENTATION CREATED BY*

*17 ORTHOTISTS AND PROSTHETISTS.—For purposes of*

*18 determining the reasonableness and medical neces-*

*19 sity of orthotics and prosthetics, documentation cre-*

*20 ated by an orthotist or prosthetist shall be consid-*

*21 ered part of the individual’s medical record to sup-*

*22 port documentation created by eligible professionals*

*23 described in section 1848(k)(3)(B).”.*

Mr. Alexander’s letter is a clear assertion of CMS’ commitment to acknowledge immediate implementation of the new statutory provisions in Section 50402, accepting the orthotist and prosthetist clinical notes as part of the individual’s medical record as to “determining the reasonableness and medical necessity of orthotics and prosthetics” e.g., functional levels, identification of broken, damaged parts and their repair, and identifying components in a category included in a physician approved detailed written order. We also suggest that AOPA members consider including a copy of the letter with all claims they file.

AOPA will continue to keep you informed of any developments in this important area.

[Click here to view the letter from Mr. Alexander.](#)

To view the DME MAC announcement of the retirement of the Dear Physician letter for lower limb prostheses, [click here.](#)

### **DME MACs Announce the Retirement of the Dear Physician Letter for Artificial Limbs**

Recently, each of the four DME MAC contractors has published a revised version of the Dear Physician letter regarding artificial limbs. The revised version of the letter states that the Dear Physician letter is being retired due to pending guidance from the Centers for Medicare and Medicaid Services (CMS) on potential program changes that may be necessary to implement the recently passed legislation that requires recognition of O&P practitioner notes as part of the patient’s medical record.

The Dear Physician letter for artificial limbs, originally published in August, 2011, fundamentally changed how Medicare claims for artificial limbs were processed. Prior to the 2011 Prosthetic Dear Physician letter, practitioner notes were generally accepted as being valuable when making claim payment decisions. The Dear Physician letter made it clear that for Medicare purposes, “it is the treating physician’s records, not those of the prosthetist’s, which are used to justify payment.” This statement, and the overall tone of the Dear Physician letter lead to years of frustration where the clinical notes of qualified, educated, certified, and often licensed prosthetists, were simply ignored during the claim review process. This exclusion of valuable clinical information lead to higher claim denial rates and unacceptable delays in the appeal process.

With the February, 2018 passage of legislation that now requires documentation created by orthotists and prosthetists to be considered part of the patient’s medical record for medical



review purposes, the statement quoted above and other parts of the Dear Physician letter are no longer consistent with the law. The DME MACs have acknowledged this and have decided to retire the 2011 Dear Physician letter for artificial limbs until they receive further guidance from CMS.

While the retirement of the Dear Physician letter does not mean that the DME MACs will no longer require physician documentation to support claims for artificial limbs, it is a clear indication that they acknowledge and understand that the provisions of the original Dear Physician letter are no longer consistent with the law and therefore can no longer be used as the sole justification for denying a Medicare claim. It also indicates that the diligent AOPA efforts to convince CMS to provide guidance on this issue to their DME MAC contractors. While AOPA does not know what that eventual guidance will be, it is clear that CMS is aware of the legislation and intends to provide guidance to the DME MACs regarding the role of O&P practitioner notes in the medical review process.

[View a sample of one of the Dear Physician letters indicating its retirement.](#)

**Exhibit at the AOPA National Assembly – Sept. 26-29 in Vancouver, Canada**



AOPA is currently accepting exhibit applications for the 2018 AOPA National Assembly which is to be held September 26-29, 2018 in beautiful Vancouver, Canada. Now is your chance to sign up and showcase your products at the largest O&P tradeshow in the Western Hemisphere. This world-wide convention opportunity features 4 days of high-level networking, exhibits, the latest techniques in O&P treatment, and the finest industry specific business and clinical training programs. We hope you make plans to join us.

**Exhibiting at this important event will give you the opportunity to:**

- Build your customer base and increase sales by meeting with Facility Owners and decision-making practitioners.
- Experience face-to-face time with existing customers to answer questions and build new relationships.
- Enjoy sponsored networking opportunities, including an opening welcome reception in the exhibit hall.
- Take advantage of fun traffic-building opportunities.
- Take advantage of education sessions to learn what's happening with U.S. health care reform, Medicare, and other regulatory agencies that affect the success of your products.
- Increase visibility for your company/organization in a targeted market.
- Host a Manufacturer's Workshop and/or Product Preview Theater presentation.
- Speak to AOPA coding experts to learn what's happening with U.S. government sponsored healthcare programs.
- Increase your exposure through a wide variety of advertising and sponsorship opportunities.
- Hear from top-researchers and clinicians to learn what products and support is needed from manufacturers.
- Participate in key education programs and plenary sessions.

- Benefit from global exposure.
- Much more!

[Click here to sign up](#) today to exhibit at the 2018 AOPA National Assembly. Questions? Contact Kelly O'Neill at [kelly.oneill@AOPAnet.org](mailto:kelly.oneill@AOPAnet.org) or call (571) 431-0852.



Join your colleagues at the Vancouver West Convention Center, Vancouver, BC, Canada for AOPA's 101st Assembly! Enjoy the best in business education and advanced clinical programming, the largest O&P exhibit hall, networking with the most influential people in the profession, must attend events, and 30+ CE credits. [Register and learn more here.](#)

### RSVP for the 2018 Party With a Purpose



Mark your calendars and attend the “Party With a Purpose” on Thursday, September 27 during the 2018 AOPA National Assembly in Vancouver. Be there or be square and wear your grooviest threads, as we celebrate and re-live the fabulous 1970’s! The night includes dinner, a costume contest, Dr. Disco, a dance off, and much more! This is a special event and certain rules and restrictions may apply. For additional information about the Party with a Purpose or to register, contact Devon Bernard at [dbernard@AOPAnet.org](mailto:dbernard@AOPAnet.org).

### Upcoming AOPA Events

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| September 12, 2018    | <p><i>Medicare as Secondary Payer: Knowing the Rules</i><br/>AOPA Webinar<br/><a href="#">Learn more and register here</a></p>                   |
| September 26-29, 2018 | <p>AOPA National Assembly<br/>Vancouver, Canada<br/><a href="#">Learn more and register here</a></p>   |
| October 10, 2018      | <p><i>Year End Review: How to Wrap-Up &amp; Get Ready for the New Year</i><br/>AOPA Webinar<br/><a href="#">Learn more and register here</a></p> |