



AMERICAN ACADEMY
OF OPHTHALMOLOGY®



September 11, 2023

The Honorable Chiquita Brooks-LaSure, Administrator
Centers for Medicare and Medicaid Services
Department of Health and Human Services
Attention: CMS-1772-P
Mail Stop C4-26-05
7500 Security Boulevard
Baltimore, MD 21244-1850

Via online submission at www.regulations.gov

Re: Medicare Program: Hospital Outpatient Prospective Payment and Ambulatory Surgical Center Payment Systems and Quality Reporting Programs

Dear Administrator Brooks-LaSure:

We appreciate this opportunity to submit comments on behalf of five leading ophthalmology organizations with regard to CMS-1772-P, *Medicare Program: Hospital Outpatient Prospective Payment and Ambulatory Surgical Center Payment Systems and Quality Reporting Programs; Organ Acquisition; Rural Emergency Hospitals: Payment Policies, Conditions of Participation, Provider Enrollment, Physician Self-Referral; New Service Category for Hospital Outpatient Department Prior Authorization Process; Overall Hospital Quality Star Rating*. Collectively, the members of our societies are responsible for performing the vast majority of all ophthalmic surgical procedures performed in the US, most of which, in the ophthalmic ASC setting.

The American Academy of Ophthalmology (AAO) is the largest association of eye physicians and surgeons in the United States. A nationwide community of nearly 20,000 medical doctors, we protect sight and empower lives by setting the standards for ophthalmic education and advocating for our patients and the public. We innovate to advance our profession and to ensure the delivery of the highest-quality eye care.

The American Society of Cataract and Refractive Surgery (ASCRS) is a medical specialty society representing 6,500 ophthalmologists in the United States and abroad who share a particular interest in and commitment to advancing the art and science of ophthalmic surgery.

The American Society of Retina Specialists (ASRS) is the largest retinal organization in the world, representing over 3,000 members. Retina specialists are board certified ophthalmologists who have completed fellowship training in the medical and surgical treatment of retinal diseases. The mission of the ASRS is to provide a collegial open forum for education, to advance the understanding and treatment of vitreoretinal diseases, and enhance the ability of its members to provide the highest quality of patient care.

The Outpatient Ophthalmic Surgery Society (OOSS) is a professional medical society that represents over 4,000 ophthalmologists, nurses, and administrators who specialize in providing high-quality ophthalmic surgical services in cost-effective ASC environments. The programs and services of OOSS are designed to ensure top-quality and sustainable patient care and safety in surgical environments that support ever-changing technology and regulation. OOSS is a member of the ASC Quality Collaboration (ASCQC), a cooperative effort of organizations and companies interested in ensuring that ambulatory surgical center (ASC) quality data is appropriately developed and reported. (ASCQC developed the claims-based quality measures incorporated within the recent rulemakings governing ASC quality reporting.)

The Society for Excellence in Eyecare (SEE) is a professional organization of ophthalmologists dedicated to educating its members about the most effective and advanced developments in ophthalmology, developing and implementing standards of practice for the effective and ethical provision of ophthalmologic services to patients, and serving as an advocate for patients in the promotion of high quality, cost-effective eye care services.

On behalf of AAO, ASCRS, ASRS, OOSS, and SEE, we are taking this opportunity to comment on this important regulation governing CY 2024 Medicare ambulatory surgical center (ASC) payment rates and the Quality Reporting Program for ASCs. Importantly, we strongly support the agency's decision in 2019 to change the ASC update factor from the Consumer Price Index – Urban (CPI-U) to the Hospital Market Basket (HMB) and are pleased that this trial has been extended from 2023 to 2025. We strongly believe that it should be adopted permanently. We will discuss below other payment policy changes that should ameliorate some of the distortions in relative payments to ASCs and HOPDs.

We draw your attention to a study by the KNG Group, *Medicare Cost Savings Tied to Ambulatory Surgery Centers*, which concluded that annual Medicare cost savings attributable to ASCs increased from \$3.1 billion in 2011 to \$4.1 billion in 2018. (We note that ophthalmology accounted for more than one-third of these savings.) Importantly, if volume migration continues at the same rate as 2011-2018, surgery centers are projected to save Medicare \$74.2 billion from 2019-2028. Policies that encourage migration embody the potential to generate even greater savings than those projected. Our comments reflect our organizations' commitment to providing the highest quality and most accessible care at lower cost to the Medicare program and its beneficiaries.

The nation's ophthalmic ASCs are committed to providing Medicare beneficiaries with access to the highest quality surgical care while lowering their cost-sharing obligations and assisting the Medicare program in the containment of health expenditures. Since 1982, ASCs have expanded their role in meeting the surgical needs of the Medicare population and have done so saving billions of dollars annually. Simply stated, at a time when public policymakers are searching for meaningful health care reform -- improving quality and access, while reducing costs -- ASCs embody the potential to be a significant part of the solution. Despite CMS' decision in 2019 to change the ASC update factor from the CPI-U to the HMB, elements of the proposed regulation, particularly the use of the rescaler to achieve budget neutrality, will continue to thwart, rather than enhance the ability of our facilities to continue to serve the nation's Medicare beneficiaries.

Under the proposed rule, facility payment for cataract removal Category I Current Procedural Terminology 66984 (CPT®) in 2024 would be \$1,165 in the ASC setting, while reimbursement for the same procedure in the HOPD would be \$2,256. The beneficiary's financial obligation in the form of copayments is \$233 in the ASC and at least \$451 in the HOPD; patient cost-sharing is always lower in the ASC. Therefore, for each cataract operation performed in an ASC instead of an HOPD, the program and beneficiary save \$1,165, and the beneficiary saves at least \$218, for a total savings of \$1,383 or more. With more than 1.8 million cataract surgery cases performed per year for Medicare fee for service beneficiaries, the impact of savings to the program and the beneficiary by performance of cataract surgery in the ASC, as confirmed now by a multitude of studies and reports, is well into the billions of dollars annually. While ASCs perform about 75 percent of cataract surgeries, there is still significant opportunity for volume migration as virtually every cataract operation can be safely and effectively performed in ASCs.

SUMMARY OF RECOMMENDATIONS

- ❖ **CMS should maintain use of the hospital market basket as the annual update mechanism for ASC payments.**
- ❖ **CMS should apply the OPPS relative weights to ASC services and discontinue the rescaling of ASC relative weights. Rescaling has had the effect of arbitrarily and inappropriately reducing ASC payment rates and causing a substantial divergence in payment rates between HOPDs and ASCs that is unrelated to the costs of delivering services in those settings**
- ❖ **CMS should develop a policy that covers all drugs that are administered at the time of cataract surgery, but are not integral or necessary to the cataract procedure, and have an FDA-approved indication to treat or prevent post-operative concerns, such as pain and inflammation (or, alternatively, indications reflected in specialty society guidelines or medical compendia) separately under Part B. Payment should also be extended to the HOPD setting.**
- ❖ **In determining payment rates for MIGS, CMS should finalize its proposed restructuring of the Intraocular Procedures APC family, assign CPT codes 66989, 66991, 0671T to APC 5493, and finalize device-intensive designation for all three procedures.**
- ❖ **With respect to payment for Dextenza and its insertion procedure, CPT code 68841, CMS should provide for separate payment in the ASC setting for the insertion procedure reported with CPT code 68841 by changing the HOPD status indicator to "J1" and continue separate payment for the drug.**
- ❖ **With respect to insertable devices, CMS should use alternative measures to calculate the device-offset percentage when necessary to ensure fair payment to ASCs including hospital invoices submitted by the manufacturer, consistent with CMS policy.**
- ❖ **CMS should adjust the New Technology IOL (NTIOL) payment rate from \$50 to \$91.06.**
- ❖ **CMS should continue assigning CPT 92229 (*imaging of retina for detection of monitoring of disease; with point-of-care automated analysis with diagnostic report;***

unilateral or bilateral) to Ambulatory Payment Classification (APC) 5733 and not finalize the proposal to reassign CPT 92229 to APC 5732.

- ❖ **CMS should permanently withdraw proposed Measure ASC-11 from the ASC Quality Reporting Program and establish a measure reporting on TASS.**

The Medicare Prescription Drug, Improvement, and Modernization Act of 2003 (MMA) mandated that CMS implement by January 1, 2008, a new ASC payment system. Our organizations and the entire ambulatory surgery community reached consensus on the appropriate contours of an equitable and rational program. In the final ASC payment regulation that became effective in 2008, CMS adhered to the spirit, if not the letter, of many of these principles, most importantly, that the new payment system should be modeled on the methodology and payment rates applicable to surgical services furnished in HOPDs.

For well over a decade, we have expressed grave concerns that the continued use of the Consumer Price Index – Urban (CPI-U) rather than the HMB as the ASC update factor as well as maintenance of the rescaler to achieve budget neutrality will continue to exacerbate the gap between the ASC and HOPD payment rates in ways that were unrelated to comparable cost differences in the provision of care in the two settings, with respect to technology and staffing costs which are identical. **We appreciate that CMS has responded to some of our concerns, particularly taking the important step of replacing the CPI-U with the HMB as the annual update factor for ASCs (at least through 2025), a key step in encouraging additional procedures to be performed in the more cost-effective ASC. To ensure that this proposal has its intended effect, however, we recommend CMS also eliminate the secondary rescaler.**

In 2003, aggregate ASC payments as a percent of HOPD rates were 85 percent. When the new system was established in 2008, the percentage had dropped to 65 percent; under the proposed 2024 rates, Medicare will reimburse ASCs, on average, 50 percent less than HOPDs performing the same procedures. We note further that whereas ASCs accounted for 6.63 percent of total expenditures in 2016, the ASC percentage of that spend is declining, representing only 5.9 percent in 2022. This situation is the result of the application of the rescaler and is entirely unrelated to the cost of providing services to Medicare patients within the respective outpatient surgical environments. At a time when ASCs offer the very real potential of augmenting access to high quality services at substantially lower cost, policymakers and the public should be concerned about the growing risk of surgery migrating back to the higher-cost HOPD. Since the advent of the new payment system, hospital market share is growing for many high-volume procedures.

In formulating ASC policy and establishing payment rates, it is imperative that the agency recognizes that most ASCs are small businesses that must run efficiently to remain in operation. There are over 6,000 Medicare-certified ASCs – about 1,200 of which specialize in ophthalmology – and over half have only one or two operating rooms. Our facilities purchase the same equipment, devices, implants, and supplies as HOPDs and must compete with hospitals for the same nurses and other personnel, while complying with the same federal and state patient health and safety requirements and the ever-growing demands of the

Medicare ASC quality reporting program. Importantly, anesthesia costs are dramatically increasing due to many factors, including declining physician reimbursement and the inordinate impact of the No Surprise Act on anesthesia providers. Our centers operate efficiently; however, receiving reimbursement that is about half that of competing hospitals compromises the ability of our facilities to continue to provide the care and technology that Medicare beneficiaries deserve.

As discussed in detail below, the agency's continued utilization of rescaling to achieve budget neutrality in the 2024 proposal, as well as the recent reclassification of procedures into new APCs and packaging policies, has exacerbated distortions in payment rates to ASCs and hospitals. In a very real sense, these policies compromise the integrity of the ASC payment system, reduce realizable program savings, increase beneficiary out-of-pocket costs, and inhibit transparency regarding price and quality among Medicare providers, thereby jeopardizing beneficiary access to affordable, high quality surgical care.

Since CMS decided almost a decade ago to overhaul the ASC payment system, our organizations have been engaged in discussions of ideas and review of data with the agency regarding the issues presented in this and recent rulemakings. We have appreciated the agency's willingness to work with the ASC industry, the ophthalmology community, and others and believe that there are many positive components to the proposed rule. With this spirit of cooperation and commitment to formulating a rational and equitable ASC payment system, we join the ASC industry and other surgical specialty organizations in offering our specific comments:

ANNUAL PAYMENT UPDATE AND REQUEST FOR COST DATA

As we have emphasized in our comments in recent years, our organizations strongly support the agency's decision to change the ASC update factor from the CPI-U to the Hospital Market Basket (HMB). The CPI-U does not reflect ASC cost growth; the HMB is a better proxy for ASC cost increases. ASCs and HOPDs treat the same patients for the same conditions and consume commensurate resources and incur similar costs. **Application of different inflation factors has unjustly expanded the gap in payments to HOPDs and ASCs. We believe that applying the same update factor to both types of facilities can potentially promote appropriate migration of services from the HOPD to ASC,** generating significant cost savings to the Medicare Program. ASC growth has been compromised by lack of parity in payment to HOPDs and ASCs. Aligning conversion factors – in addition to eliminating the rescaler, as discussed below – will equitably level the playing field between hospital and freestanding surgical facilities. We applaud the agency's decision to maintain the HMB as the ASC update factor through 2025, and we look forward to collaborating with CMS over the next two years.

CMS has, in the proposed rule, again expressed a desire to assess the feasibility of collaborating with stakeholders to collect ASC cost data in a minimally burdensome manner. For the reasons stated above, we believe that the HMB is an appropriate update factor for ASCs. **If, however, CMS elects to collect data to establish a new market basket, the agency should expand its analysis to create an index that will be applied to both the HOPD and ASC to**

ensure that payments using the same relative weights are aligned over time. In developing a data collection modality, CMS should keep in mind that ASCs already incur excessive administrative burdens in complying with current regulations; requiring formal cost reports would diminish the agency's commitment to promulgate rules and policies that allow facilities to maintain efficiency in the delivery of services to our patients. We look forward to collaborating with CMS on this endeavor.

RESCALING ADJUSTMENT APPLIED TO ASC RELATIVE VALUE WEIGHTS

AAO, ASCRS, ASRS, OOSS, and SEE strongly believe that CMS should eliminate the rescaling of the ASC relative weights, as this practice has exacerbated the gap between ASC and HOPD payments and inappropriately reduced payments to ASCs without evidence of growing differences in capital and operating costs in the two settings. As we have noted in our comments to past ASC payment rulemakings, our organizations support the utilization of the same APCs and relative weights in creating a rational and coherent payment system encompassing the services offered by both HOPDs and ASCs:

“. . . the rescaling of ASC relative weights . . . will result in further divergences in weights and payments, exacerbating exactly the types of distortions that the new system was presumably intended to correct. The only legitimate basis for change in relative payments to HOPDs and ASCs should be changes in the relative costs of providing specific outpatient services. There is little basis for believing that these variations will occur, and to the extent that they do, they should be accounted for directly through adjustments to the conversion factor.”

It is important to note that APC relative weights are already adjusted once for budget neutrality. Contrary to CMS' assertion in 2007 that rescaling would protect ASCs from decreases in payments for procedures due to changes in OPSS relative weights, experience since that time reflects otherwise. *The rescaling adjustment has had the opposite effect, decreasing the relative weights on ASC surgical procedures each year. Since 2010, our relative weights have decreased by an average of 7 percent each year. There is no evidence to suggest that there are growing differences in capital and operating costs in the two settings to support such an accelerating differential. This historical trend suggests that the application of the rescaler in the ASC environment will continue to erode the relationship between ASC and HOPD payments.* The agency is needlessly increasing Medicare program costs by making it financially impracticable to furnish these services that are clinically appropriate for the ASC, and hence pushing physicians to return these procedures to the more expensive HOPD setting. **We strongly recommend that the agency discontinue the use of the rescaler.**

We note that CMS is not required to maintain rescaling. Congress imposed a budget neutrality requirement on the new ASC payment system *only* during the inaugural implementation year of 2008; CMS is under no legal obligation to continue to apply rescaling and should not do so when it creates significant disparities in relative payments to ASCs and hospitals that are not related to the costs incurred in providing such services. Therefore, we implore the agency to encourage savings and greater access to ASCs for Medicare beneficiaries by eliminating the ASC weight scaler.

Our organizations realize that the elimination of the ASC weight rescaler would increase Medicare program costs, at least initially, until cost savings are achieved by volume shifting to the ASC setting. As an alternative, **we propose that CMS refrain from the applying the secondary rescaler to the ASC payment system and, instead, combine the OPPS and ASC utilization and mixes in services to establish a single weight scaler.** By incorporating the ASC volume into OPPS weight scaler calculations, CMS would improve the alignment of the payment systems and more accurately scale for outpatient volume across both sites of service.

Now, sixteen years after the implementation of the ASC payment system, it is past time for CMS to analyze the drivers of continuing increases in the OPPS rescaler and whether such factors should or should not be offset in the ASC setting.

COVERAGE OF NON-OPIOID DRUGS

We were pleased that CMS proposed to codify last year a modest expansion of its Non-Opioid Pain Management policy and solicit comments on several related implementation issues. To date, CMS is paying separately in the ASC for only a very limited number of specific non-opioid pain management drugs. Specifically, AAO, ASCRS, ASRS, OOSS, and SEE are concerned with the bundling of FDA-approved drugs that are administered at the time of ophthalmic surgery—either before, during or at the end of the procedure— and which can be substituted for postoperative administration of drugs which have an indication for the treatment of post-operative pain and/or inflammation and/or other sequela of the surgery. Our organizations believe that CMS should consider as part of future rulemakings to expand its proposed criteria for Non-Opioid Pain Management Drugs, in addition to the FDA label indication, to also recognize an intraoperatively-administered drug which can be substituted for postoperative drugs, when demonstrated based on the recommendation of recognized specialty societies, or guidelines contained in medical compendia.

AAO, ASCRS, OOSS, ASRS and SEE maintain that since these medications have a post-operative indication and can reduce or eliminate the need for postoperative medications, they should not be considered surgical supplies bundled into the facility payment, but instead be covered and paid for under Medicare Part B, just as topical steroids and non-steroidals are paid under Part D. ASCs face the same challenges in affording to provide these drugs as they do to provide the specific non-opioid pain management drugs -- Omidria and Dextenza are the only ophthalmic drug encompassed by existing policy for payment of non-opioid drugs -- that CMS removed from the facility fee for that reason. The problem is compounded by the reality that ASCs are receiving, as discussed in detail above, only half of the reimbursement provided to HOPDs.

Chang, et.al. recently published an article in JAMA Ophthalmology which concluded that despite offering the potential of improved compliance and greater patient convenience, findings of this study suggest that sustained-release corticosteroid use during cataract surgery was low and associated with cost increases to the health care system vs conventional postoperative eye drops. As these new products must be priced high enough to qualify for the Medicare pass-through program, unreasonable cost may have been a deterrent to their use, suggesting that the current Medicare reimbursement rules may not be appropriate for sustained-release postoperative

corticosteroids in cataract surgery.”

Without the assurance that ASCs will be able to afford to provide these treatments to patients, manufacturers may discontinue their innovation in this area.

Dexamethasone is a drug commonly used in ophthalmology because of longstanding evidence that it reduces inflammation, and therefore, pain following cataract surgery. Some drugs that have the active ingredient dexamethasone, such as Dextenza, are already indicated for both inflammation and pain. Others, such as Dexycu, are indicated for post-operative inflammation alone. *The agency should not treat label differences as determinative as to whether a drug like Dexycu can be designated as a non-opioid pain management drug. Inflammation following ocular surgery is a major risk factor for pain.* Ocular inflammation induces activation of neurons within the sensory trigeminal complex; altered activity in intracellular signaling caused by ocular inflammation also plays a role in the sensitization of ocular-related brainstem circuits, leading to the development of ocular pain. By reducing inflammation, ophthalmic dexamethasone reduces a key risk factor associated with post-operative pain.

We appreciate that CMS solicited comments in 2022 regarding whether the agency should rely on factors other than FDA-approved indications labeling in its process for determining whether a drug qualifies as a non-opioid pain management drug. *Our organizations believe that, in future rulemakings, CMS should consider as an alternative to the criterion based on FDA label to allow the recommendations of specialty societies or other national organizations such as medical compendia developers.* Specialty societies and other clinical experts already play a key role in identifying clinically recognized uses for drugs that extend beyond FDA-approved labeling. It is important to consider an alternative criterion as the guidelines contained in medical compendia are continually evolving as clinicians identify new uses for existing drugs and these uses are empirically tested by scientists and clinicians.

We and our patients are fortunate to have multiple options to meet the post-operative challenges our patients face, including excellent self-administered medications as well as innovative and effective surgeon-administered drugs. Our members and facilities believe that patients should be afforded the option of using self-administered eye drop medications post-procedure (Part D products) or to have FDA-approved drug products administered by the surgeon at the time of the ophthalmic surgery (Part B products). **Therefore, we urge CMS to develop a policy that covers and pays for drugs that are administered at the time of ophthalmic surgery, are substitutes for postoperative medications, and have an FDA-approved indication to treat/prevent post-operative issues, such as pain, inflammation, or infection, separately under Medicare Part B.**

With respect to other related and relevant implementation issues:

- We urge CMS to adopt refinements to the non-opioid authorities to help ensure patient access to drugs, like Dexycu, that were marketed prior to 2022 but that now must seek an FDA label expansion for pain indications before they can be eligible for such status. This can be accomplished by temporarily “grandfathering” for two or three years these products so that payment doesn’t lapse pending eligibility for non-opioid status.
- We likewise support the extension of separate payment for qualifying products in the

hospital outpatient setting. Recent data demonstrate that the lack of separate payment for these drugs in the HOPD setting following the expiration of its pass-through status resulted in a significant drop in utilization of products in the HOPD. This was in contrast to the stable utilization in the ASC setting where Omidria continued to be paid separately as a non-opioid pain management drug. In response to CMS' request for comments on expanding the policy to the HOPD setting, our organizations support the expansion because it would harmonize beneficiary access to non-opioid pain management products across settings of care.

- We recommend that CMS continue to use the standard ASP + 6% payment methodology to mitigate payment differentials across settings.

FACILITY PAYMENT FOR MIGS AND OTHER INTRAOCULAR PROCEDURES

Micro-invasive glaucoma surgery (MIGS) may be performed with a cataract procedure or without a cataract procedure. CPT codes 66989 and 66991 are used to report a MIGS procedure with a complex and routine cataract procedure, respectively, while CPT 0671T is used to report a MIGS procedure that is performed without a cataract procedure. CMS proposes to assign all of these procedures to an APC in the Intraocular Procedures APC family for 2024.

Before addressing the specific APC proposals, we first turn to CMS proposed restructuring of the Intraocular Procedures APC family. CMS proposes to add another APC to this family by splitting APC 5492 into two APCs. **We agree with CMS's proposal to restructure the Intraocular Procedures APC family, and we recommend CMS finalize them.**

With respect to the MIGS procedure codes, CMS proposes to assign CPT codes 66989 and 66991 to APC 5493, and proposes to assign CPT code 0671T to APC 5492. Our organizations agree with the proposed APC assignments for CPT codes 66989 and 66991 and recommend that CMS finalize the proposal to assign both codes to APC 5493. For CPT code 0671T, CMS also should assign the procedure to APC 5493 rather than APC 5492 in light of the CY 2022 hospital outpatient claims data for the procedure. Those data illustrate a geometric mean cost that is more consistent with the procedures assigned to APC 5493 than APC 5492. Given the clinical and resource similarity of all three procedures (CPT codes 66989, 66991, and 0671T), CMS should assign them to APC 5493.

We are encouraged by these proposals for improved payment for MIGS procedures and we thank CMS for their structured analysis. Since the majority of these procedures are performed in the ASC setting, it is critical to ensure that the payment rates for these procedures are adequate. There is a great deal on innovation in the MIGS device space and these new technologies can have a significant impact on lowering intraocular pressure with lower risk of complications. We encourage CMS to review applications for new HCPCS Level II C-Codes and assign device intensive status when necessary to promote patient access in the ASC setting. In addition, because adequate claims data may not be available for these new C-codes and CPT codes, we urge CMS to use hospital outpatient invoices that are submitted by the manufacturer.

ASC PAYMENT FOR DEXTENZA AND THE INSERTION PROCEDURE

DEXTENZA (Dextenza) is approved for the treatment of ocular inflammation and pain following ophthalmic surgery and is reported using J1096 (Dexamethasone, lacrimal ophthalmic insert, 0.1 mg). Dextenza is a physician-administered corticosteroid intracanalicular insert that is inserted following an ocular procedure and is currently reported using CPT code 68841 (Insertion of drug-eluting implant, including puntal dilation when performed, into lacrimal canaliculus, each). Depending on the use, the procedure may be the only one an ASC furnishes to a beneficiary, or another ophthalmic procedure may also be furnished on the same day. While ASCs incur costs for the inserted drug and for performing the procedure, under CMS's proposal, ASCs would only receive reimbursement for the drug. Because CMS proposes to assign a Q1 status indicator for CPT code 68841 for hospital outpatient payment purposes, that status indicator means no payment is made to ASCs for the procedure, which is untenable, especially since the procedure is predominantly furnished in ASCs.

In the proposed rule, CMS does not explain why it believes it is appropriate not to compensate ASCs for the costs they incur in furnishing this service. In prior rulemakings, CMS has indicated that it assigned a Q1 status indicator to CPT code 68841 because the procedure is mostly performed during other ophthalmic procedures. That is not entirely clinically accurate. The insertion procedure reported by CPT code 68841 is not performed *during another procedure*, but at the conclusion of another procedure during the same operative session. This is not a rational reason for denial of coverage. There are numerous examples of procedures being performed together more than 75% of the time, that is addressed by development of a combination CPT code. In addition, when the procedure is performed to treat a patient that has ocular itching associated with allergic conjunctivitis, the beneficiary does not receive another ophthalmic procedure that day. As a result, CMS has wrongly assigned a Q1 status indicator to the procedure – and uniquely so, as the other 70 or so procedures in the same APC have a J1 status indicator.

The impact of this incorrect status indicator assignment disproportionately falls on ASCs. While hospitals can be paid for CPT code 68841, such as when there is no other payable service on the claim, an ASC is never paid for the service, despite the fact that it incurs significant costs to furnish the service (the hospital geometric mean cost for the procedure is over \$2,000). There is no basis for this differential treatment between hospitals and ASCs and this differential adversely impacts Medicare beneficiaries. With no payment for the service, there is a financial disincentive for ASCs to furnish the service – no compensation for the facility type costs incurred by the ASCs. CPT code 68841 should be treated like other similar procedures for introducing a product into the eye (e.g., CPT codes 66030 and 0699T) and be assigned an OPPS status indicator of J1 so that payment for the procedure would be made in the ASC setting.

As indicated in our comments above, we reiterate our support for the continued separate payment for Dextenza (J1096) to ASCs as a drug with a post-operative indication that is not a surgical supply, and therefore, should be paid separately under Medicare Part B. We believe that separate payment should flow from the product being a separately payable drug under the hospital outpatient payment system (K status indicator). Dextenza should not be treated as a packaged drug because it is not integral, ancillary, or supportive of another surgical procedure. As noted above, a facility may furnish just the drug and the insertion procedure, just like a facility may furnish a drug and an administration procedure. Those drugs are not packaged and Dextenza likewise should not be packaged. As such, separate payment to ASCs for Dextenza should not

come as a result of the non-opioid pain management policy. While we support separate payment for Dextenza under the non-opioid pain management policy, we believe that policy should not be needed because Dextenza should be a K status indicator drug and thus separately payable to ASCs.

DEVICE INTENSIVE PROCEDURES

Our organizations are appreciative of the agency's efforts to address the device-intensive threshold and its impact on ASC volume. The agency's recognition of the important role that devices play in our ability to perform surgical procedures – by reducing the devices threshold over the past several years from 50 percent to 40 percent and then to 30 percent and by establishing that the threshold determinations should be based on the percentage the device accounts for in the surgery center as opposed to the percentage in the HOPD -- has enabled our members to offer more high-quality services to our patients at great savings to the Medicare program and its beneficiaries.

Recent changes to the device-intensive threshold have greatly increased the number of device-intensive codes on the ASC-CPL, but it has also shone a spotlight on how the lack of complete alignment in the HOPD and ASC payment systems serves as a barrier to access for Medicare beneficiaries. While there is a statutory cap on the patient responsibility when a procedure is done in a hospital, including an HOPD, that policy is not in place for the ASC setting. Even though the Medicare beneficiary's patient responsibility is capped, the hospital is made whole by the Medicare program. Beneficiaries who would otherwise have access to the high-quality, convenient ASC setting are disadvantaged by this lack of alignment in policy. **We recommend that CMS join the ASC community in encouraging Congress to implement an ASC copayment cap with respect to devices implanted during surgery in an ASC.**

Because of the copayment cap and its impact on many device-intensive codes, there are many areas, particularly in rural communities, where the wage index is so low that it makes it financially unsustainable for rural facilities to provide certain device-intensive procedures to Medicare beneficiaries. **To address this concern, CMS should refrain from adjusting the device portion of the payment by the local wage index.** This is consistent with the Agency's policy for separately payable drugs and biologics, and it is highly unlikely that a facility in a rural community is getting a better deal on devices than ASCs in large cities.

AQUEOUS SHUNT CODES

We appreciate that CMS has revised APC 5493 to better serve Medicare beneficiaries undergoing intensive ophthalmic procedures. For patients with severe glaucoma that have previously failed medical therapy, laser treatments, and often incisional procedures, this represents the permanent implantation of aqueous shunts. These are among the most resource-intensive procedures performed in our facilities, which is appropriately reflected in the placement of CPT code 66179 (*Aqueous shunt to extraocular equatorial plate reservoir, external approach; without graft*) in APC 5493 in the 2024 OPSS/ASC proposed rule. However, we are troubled that CPT code 66180 (*Aqueous shunt to extraocular equatorial plate reservoir, external approach; with graft*) remains in APC 5492. CPT codes 66179 and 66180 describe nearly *identical* procedures with an important discrepancy being that CPT code 66180 involves fixating a patch graft as well,

substantially increasing a facility's expenditures for that procedure as it must procure the patch graft in addition to the other surgical devices and materials involved.

Fortunately, these procedures are not as common as less invasive glaucoma interventions, but when they are required, the resources consumed are substantially more costly. **We request that CMS coherently reflect this by placing both CPT code 66179 and CPT code 66180 in APC 5493 in the 2024 OPPTS/ASC final rule.**

NEW TECHNOLOGY IOL (NTIOL) PAYMENT RATE

Embedded in our comments is the premise that CMS should promote innovation and ensure that Medicare beneficiaries have access to critical, life-saving new cures and technologies that improve beneficiary health outcomes. This should apply to all emerging technologies, including New Technology Intraocular Lenses (NTIOLs), which replace a beneficiary's natural lens that has been removed in a cataract surgery. Since 1999, when CMS established the payment rate for a new NTIOL at \$50 per lens, there has not been a payment adjustment. While there have been major technological advances in IOLs over the last two decades, there continues to be addressable unmet needs for post-cataract patients, including enhancing lens quality, reducing secondary interventions, and addressing negative dysphotopsia. In the final rule, our organizations request that the agency update the NTIOL payment rate of \$50 to \$91.04, which is reflective of the increase in the Consumer Price Index during the period 1999 to 2022.

In real dollar terms, the flat rate payment for NTIOLs has significantly lagged behind the overall economic inflation rate and is not reflective of the increased costs associated with research and development to bring new technologies to market. When the NTIOL provisions were established **more than two decades ago**, CMS stated it would revisit the appropriateness of the fee established for cataract surgery with IOL insertion when a lens determined to be an NTIOL is furnished. Now is an appropriate time to make such an adjustment.

The IOL industry continues to advance ocular innovation with several new monofocal IOL technologies. These future IOLs have the potential to provide meaningful outcomes for cataract patients.

In order to meet CMS' original objectives to enable Medicare beneficiaries' expedited access to new technologies, we strongly encourage CMS to consider increasing the per lens payment from \$50 to \$91.04 in CY 2024. Additionally, we request that the NTIOL payment be updated annually to reflect CMS' decision to increase the NTIOL payment will help ensure Medicare beneficiaries have expanded access to future innovation.

In conversation with CMS representatives, we have learned that there are currently no active classes of NTIOLs. As such, since 2011, stakeholders have submitted information to be approved as astigmatism-correcting and/or presbyopia-correcting *instead of* pursuing NTIOL status. In order to foster innovation and support patient access to modern, rapidly-evolving technologies, we propose that CMS publish annually a list of active NTIOL classes. The publication of active NTIOL classes will allow new applications to be accepted and catalyze a reemergence of participation within this program across multiple companies within industry.

RETINA IMAGING CODES

As ophthalmologists, our members have a duty to protect the vision of Medicare beneficiaries. That responsibility extends to making sure the right patients get the right treatment at the right time through regular vision screening. For diabetic patients, early detection is paramount.

Thirty million people in the U.S. have diabetes, and diabetic retinopathy affects nearly 30 percent of diabetic patients.¹ Diabetic retinopathy and diabetic macular edema are complications of diabetes and the most frequent causes of blindness in adults. Since these conditions do not have symptoms at first, early detection is critical to preserving vision. For this reason, both the American Diabetes Association (ADA) and the American Academy of Ophthalmology (AAO) recommend annual eye exams and prompt referral to a specialist if signs of diabetic retinopathy are detected.^{2,3}

The new retinal imaging procedure reportable with CPT 92229 allows early detection of sight-threatening diabetic retinopathy with immediate point-of-care feedback to the patient and primary care provider.⁴ It can be performed in primary care offices and will expand the number of beneficiaries whose retinopathy can be identified early enough to prevent irreversible vision loss. It promises a significant advance in public health and substantial program savings by reducing the need for costly treatment of advanced diabetic eye disease. For CY 2024, Addendum B released with the proposed rule indicates that CMS is proposing to reassign CPT 92229 to APC 5732. As a result, reimbursement will shrink by about 36 percent to \$37.05. Reimbursement adequate to cover practice costs is necessary to ensure wider adoption and we feel such a drastic reduction will hinder access to care for Medicare beneficiaries with diabetes who have been shown to benefit from the CPT 92229 service^{5,6}.

¹ Lundeen E.A., Wittenborn J., Benoit S.R., & Saaddine J. Disparities in Receipt of Eye Exams Among Medicare Part B Fee-for-Service Beneficiaries with Diabetes — United States, 2017. *MMWR Morb Mortal Wkly Rep* 2019;68:1020–1023. [https://dx.doi.org/10.15585/mmwr.mm6845a3external icon](https://dx.doi.org/10.15585/mmwr.mm6845a3external%20icon)

² The American Diabetes Association. (2019). Microvascular Complications and Foot Care: Standards of Medical Care in Diabetes-2020. *Diabetes Care*, 43(Suppl. 1), S135–S151. <https://doi.org/10.2337/dc20-s011>

³ American Academy of Ophthalmology. (2019, September 25). *Diabetic Retinopathy Preferred Practice Pattern*. <https://dx.doi.org/10.1016/j.ophtha.2019.09.025>

⁴ Aiello L.P., Odia I., Glassman A.R., et al. (2019). *Comparison of Early Treatment Diabetic Retinopathy Study Standard 7-Field Imaging With Ultrawide-Field Imaging for Determining Severity of Diabetic Retinopathy*. *JAMA Ophthalmology*, 2019;137(1):65-73.

⁵ Huang, J., Wang J., Channa, R., Wolf, R., Abramoff, M.D., & Alvin Liu, T.Y. (2023). Autonomous artificial intelligence exams are associated with higher adherence to diabetic retinopathy testing in an integrated healthcare system [Abstract]. *Investigative Ophthalmology & Visual Science*, 64(8), 212. Retrieved September 1, 2023, from <https://iovs.arvojournals.org/article.aspx?articleid=2785915>

⁶ Leong, A., Wang J., Wolf, R., Channa, R., Abramoff, M.D., Lehmann H., & Alvin Liu, T.Y. (2023). Autonomous artificial intelligence (AI) increases health equity for patients who are more at risk for poor visual outcomes due to diabetic

Our organizations strongly urge CMS to continue assigning CPT 92229 (*imaging of retina for detection of monitoring of disease; with point-of-care automated analysis with diagnostic report; unilateral or bilateral*) to Ambulatory Payment Classification (APC) 5733 and not finalize the proposal to reassign CPT 92229 to APC 5732. We believe keeping CPT 92229 in its current APC assignment is appropriate because the service described by CPT 92229 is comparable to other services assigned to APC 5733 both clinically and in terms of resource use.

Our organizations are concerned that lowering the APC assignment for CPT 92229 to 5732 will have deleterious effects on patient access to diabetic retinopathy screening in the clinic setting. Per the 2024 Medicare Physician Fee Schedule (MPFS) Proposed Rule, CPT 92229 is included in the list of imaging services subject to payment cap as defined by Section 5102(B) of the Deficit Reduction Act of 2005 (the DRA). The law requires the physician fee schedule payment for the technical component of these imaging services, including the technical component of the global payment, to be capped at the OPPS payment rate. If CMS finalizes the APC reassignment CPT 92229 to 5732, the 2024 MPFS rate for the service would be limited to the outpatient rate of \$37.05.

Reimbursement that adequately recognizes that practitioners incur resource costs for ongoing use of the AI software is needed. The analysis fee is a direct practice cost analogous to a supply because it is attributable to a specific imaging service provided to a specific individual patient each time it is performed. It is not an indirect cost like computer hardware or software that is purchased once or licensed annually or monthly and then utilized repeatedly to provide varying services to multiple patients.

Greater access to this technology is an important health equity issue, as significant health disparities exist in diabetes care and access to the diabetic eye exam. Without sufficient reimbursement, medical and ophthalmology practices are unlikely to adopt this sight-saving advancement and patients across the country will be unjustly disadvantaged. To remove barriers in underserved communities and individuals, devices that promise a significant advance in public health should be accurately priced to ensure its widespread adoption.

We strongly urge CMS to ensure continued patient access to the CPT 92229 service by continuing to assign CPT 92229 to APC 5733 for CY 2024.

UNLISTED CODES

An important anomaly in CMS' effort to align the ASC and HOPD payment systems is the treatment of procedures for which there is not an appropriate CPT code. In some ASCs, surgeons utilize innovative techniques or new technologies to perform a procedure; this can mean that the service is not described by a specific CPT code. These services are reimbursed in the HOPD (and, indeed, in physician offices) but are not eligible for payment in the ASC. In the proposed 2008 ASC payment rule, CMS stated that, without knowledge of the procedure's code, it cannot determine whether the procedure performed would have been excluded from

eye disease (DED) [Abstract]. *Investigative Ophthalmology & Visual Science*, 64(8), 243. Retrieved September 1, 2023, from <https://iovs.arvojournals.org/article.aspx?articleid=2785901>

the ASC payment under the rule's safety criteria.

Although an unlisted code does not allow the reporting of specific procedures, the code does include the narrowly defined anatomic region of the service that could provide the basis for a determination about the safety of the procedure in the ASC. There is no clear safety rationale for this policy and commercial insurers typically afford ASCs the flexibility to use unlisted CPT codes to make claims for payment. We note that the agency does permit HOPDs and even physician offices to use unlisted codes; allowing this practice for ASCs will enable CMS to derive savings for both the program and beneficiaries. If physicians are permitted to choose to perform a procedure with an unlisted code in HOPDs, facilities that are managed, staffed, and equipped like Medicare-certified ASCs, surgeons should be allowed to utilize unlisted codes in the ASC. **We urge CMS to revise the Federal Code of Regulations to eliminate this restriction on billing with unlisted codes.**

QUALITY REPORTING PROGRAM (QRP)

AAO, ASCRS, ASRS, OOSS and SEE very much appreciate the efforts undertaken by CMS to implement the ASC Quality Reporting Program over the past several years and the agency's acceptance of many of the suggestions proffered by our organizations. Accommodating the perspectives and concerns of the ASC and surgical communities is undoubtedly a major factor in the exceptional 98-plus percent reporting rate by facilities with respect to measures implemented to date. We believe that the following are prerequisites to the adoption of a quality measure for the ASC. A measure should:

- Relate specifically to the episode of care in the ASC;
- Evaluate the practices and quality of the care facility;
- Involve reporting by the facility of data available in the ASC chart;
- Produce outcomes data that is actionable by the ASC, embodying the potential to improve the quality of care provided in the facility; and,
- Have been tested in the ASC environment.

ASC-11 Should Be Permanently Withdrawn as an ASC Quality Measure.

Since the publication of the 2014 ASC payment rule, we have strenuously objected to the application to ASCs of Measure ASC-11 (NQF 1536): *Cataracts – Improvement in Patient's Visual Function Within 90 Days Following Cataract Surgery*. We applauded the agency's decision several years ago to withdraw the measure for purposes of mandatory reporting by facilities and were disappointed that this measure, misguided in its application to the ASC, was included in the final 2022 rule. **While we support the agency's recent decision to defer implementation of ASC-11 until after the Covid-19 public health emergency and at least through 2024, our organizations strongly believe that ASC-11 should be withdrawn altogether.**

NQF 1536, from which it is derived, is a patient-reported outcome measure taken singularly from a measure group designed for registry-only reporting by physicians and was never intended to serve as a measure of facility-level quality and has never been tested for facility reporting. The bedrock foundation of quality reporting in the ASC is that facility-level measures should relate to *an episode of care that occurs within the confines of the ASC, encompass data that*

is available within the ASC chart, be collectable by ASC staff, generate conclusions that are actionable by the facility, and have been tested in the ASC environment. ASC-11 meets none of these criteria. For the reasons stated herein, we are not surprised that only a few dozen facilities opted to report on ASC-11 on a voluntary basis, and, therefore, sparsely reported results could not possibly result in a sufficiently meaningful sampling in terms of measuring ASC quality, either on a facility-by-facility or industry-wide basis.

- **The data required for reporting by facilities is not available within the records of the ambulatory surgical center; facilities should not be required to access and report data that is only available to other providers, such as the physician’s office.**

Even if NQF 1536 was to be tested and resubmitted for endorsement by NQF at the facility level, there are insurmountable barriers to collecting data for these measures within the ASC. *All the currently implemented ASC quality measures utilize data that is housed within and collected by the ASC itself. This new measure requires reporting on data that is located in the surgeon’s office and is wholly inaccessible by the ASC.*

Data collection for ASC measures is complicated by the fact that ASCs are administratively and financially separate from physician offices. Under the Medicare program, an ASC operates exclusively for the purpose of furnishing ambulatory surgical services to patients requiring such a setting. Although the governing regulations permit the surgical facility to exist adjacent to a physician’s office under certain circumstances, Medicare ASC Conditions for Coverage state very clearly that the two entities must be physically, administratively, and financially separate from one another. *Importantly, medical record keeping must always be maintained separately and exclusively from other operations.* While CMS has referred to NQF 1536 as a “chart-based” measure, none of the data necessary to appropriately comply with the measures are located in the ASC patient records. In other words, even though a physician in the clinic may perform surgery in the ASC next door, the medical records of one entity are *never* readily accessible by the other. As a practical matter, the ASC is staffed by registered nurses, operating room technicians, and clerical staff who are neither qualified to evaluate surgical outcomes nor located in the physician’s office where pre-operative and post-operative care might be efficiently and accurately evaluated.

- **It is not appropriate to migrate a measure across settings of care that was not considered in the design, testing, and implementation of the measure and with respect to which the measure is not actionable by the facility.**

ASC-11 is inappropriate for reporting (mandatory or voluntary) by the ASC because facilities are not equipped to evaluate potential cataract outcomes. The facility is not involved in the baseline events preceding the surgery against which outcomes are measured in the post-surgical events that encompass the healing process. It is inconceivable that the ASC, which is neither licensed nor qualified to evaluate the cataract patient and make these assessments, would be involved in the professional decision-making contemplated by the measure. *Any improvement in visual function would be attributable to the individual surgeon, not to the facility where the surgery occurred.* Physician-level measures such as those incorporated within the Quality Payment Program (QPP) are formulated to assess quality *within the physician’s office.*

Importantly, *NQF 1536 has not been tested nor endorsed as a facility-level measure for the ASC setting.* Indeed, the Measures Application Partnership (MAP), in reviewing the measure,

admonished that it should be tested and NQF-endorsed for the facility level of analysis; this has not occurred.

The goal of the ASC QRP should be to reflect those aspects of patient care that are within the control of the facility (e.g., patient safety, staffing, equipment) and for which it is reasonable to hold the ASC accountable. The surgeon's decision to recommend cataract surgery is based upon factors other than the potential for functional benefit – visual acuity and visual impairment are also considered. *It is inconceivable that the ASC, which is neither licensed nor qualified to evaluate the cataract patient and make these assessments would be involved in the professional decision-making contemplated by the measure.* As discussed above, a measure should be applied to the ASC only where actions that might be undertaken to improve such quality are within the purview of the facility. *We do not believe that it is possible for ASCs to influence outcomes with respect to this measure; therefore, it is not appropriate to include the measure in the ASC Quality Reporting Program.*

- **Implementation of this measure in the facility would be extremely burdensome and resource-intensive for the reporting ASC.**

The burden on ASCs to disseminate, collect, and report on this measure would be significant – even in a survey with a modest number of patient respondents-- given that the facility's exposure to the patient is essentially limited to the care provided at the ASC on the day of surgery. The measure requires that patients complete a visual function questionnaire both before and after their scheduled surgery. The results of the pre- and post-surgery questionnaires are then compared to assess patient-perceived improvement. The patient's visual acuity is measured before and after surgery in the physician's office, not the ASC. Single-specialty facilities like ophthalmic ASCs tend to be smaller, have fewer operating rooms and employees, and, typically, will have available diminished resources to meet the administrative burdens posed by regulatory initiatives like quality reporting.

These burdens are not trivial. The measure contemplates the administration of two visual function patient questionnaires, one completed by the patient prior to surgery and the other completed during the 90-day period following surgery. These survey instruments would be delivered to the patient during his pre-op and post-op visits *with the physician*. Facilities would be required to contract with an outside entity to administer pre- and post-surgical surveys and tabulate the results, a burdensome and expensive exercise considering the number of cataract surgeries performed in surgery centers, or somehow mandate that surgeons on the medical staff of the ASC conduct visual function surveys as a condition of maintaining ASC staff privileges. The process contemplated with respect to reporting on the measure, whether performed on a mandatory or voluntary basis, places the ASC in the arbitrary and unworkable role as “middleman” between the physician and the government. The numerous modifications to the manual specifications for ASC-11 -- which have continued to be confusing, ambiguous, and unrealistic – reflect the monumental challenge that even the agency has encountered in clarifying just what would be expected of facilities.

- **Adoption of ASC-11 would not produce data that is actionable by the ASC, and, as such, does not embody the potential to improve the quality of care provided by the ASC.**

Finally, the surgeon's decision to recommend cataract surgery is based upon factors other than the potential for functional benefit – visual acuity and visual impairment are also considered. It is inconceivable that the ASC, which is neither licensed nor qualified to evaluate the cataract patient and make these assessments, would be involved in the professional decision-making contemplated by the measure. As discussed above, a measure should be applied only where actions that might be undertaken to improve such quality are within the purview of the facility. *We do not believe that it is possible for ASCs to influence outcomes with respect to this measure; therefore, it is not appropriate to include the measure in the ASC Quality Reporting Program.*

We strongly recommend that ASC-11 be permanently withdrawn as an ASC quality measure.

CMS Should Adopt a Quality Measure for TASS

In 2018, CMS invited public comment regarding the adoption of a measure, developed by the ASC Quality Collaboration to assess the number of patients diagnosed with TASS (Toxic Anterior Segment Syndrome) within two days of undergoing anterior segment surgery in the ASC. The measure was reviewed by the Measures Applications Partnership (MAP) four years ago and received conditional support pending endorsement by the National Quality Forum (NQF). CMS did not finalize adoption of this measure in the 2018 rulemaking.

TASS, an acute and serious inflammation of the anterior chamber, or segment, of the eye following cataract surgery, is directly related to extraocular substances that inadvertently enter the eye during surgery. The incidence of TASS is measurable, attributable to the ASC, and is actionable by the facility. There are published guidelines regarding cleaning and sterilization of intraocular surgical instruments to help improve quality and prevent TASS. This measure would promote collaboration between the surgeon and the facility, as the surgeon, under current practice, would report back to the facility any incidence of TASS. Further, measuring the incidence may aid in better tracking and understanding of the prevalence of TASS, as the Food and Drug Administration contends that TASS is significantly under-reported, and surveillance is underway. Specific prevention guidelines have been developed and this measure would help ensure that they are being followed appropriately. **AAO, ASCRS, ASRS, OOSS, and SEE strongly support inclusion of the TASS measure in the ASCQR program.**

CMS Should be Cognizant of the Costs of Facility Participation in Quality Reporting Surveys

Our organizations are concerned that CMS fails to adequately consider the cost burden on facilities when establishing new quality measures. For example, ASCs will be required to contract with a third-party vendor to administer the Consumer Assessment of Healthcare Providers and Systems Outpatient and Ambulatory Surgery (OAS CAHPS) beginning in 2025. CMS maintains a list of approved vendors, but it is not clear that CMS knows how much their approved vendors charge. Vendor prices vary significantly, including at least one as high as \$20,000 per year. At some point, an ASC may make the business decision to decline to participate in the ASCQR Program and accept the two percent penalty. This does not mean the facility will be any less safe – or that it will not be collecting quality metrics. But if the burden outweighs the benefit to patients and the facility, we may see ASCQR Program participation decline. **We urge CMS to add measures – or modify the measures already in the ASCQR Program – to reduce costs and encourage ASC participation.**

Thank you for providing our organizations with the opportunity to present our views on the proposed regulation regarding 2024 Medicare ASC payment rates and the ASC Quality Reporting Program. Should you have any questions or require further information, please feel free to contact us at: Brandy Keys, Director of Health Policy, AAO, bkeys@aao.org, 202.737.6662; Mark Cribben, Director of Government Affairs, ASCRS, mcribbs@ASCRS.org, 703.591.2220; Allison Madson, Vice President of Health Policy, ASRS, allison.madson@asrs.org, 312.578.8760; Michael Romansky, JD, Washington Counsel, OOSS, mromansky@OOSS.org, 301.332.6474; and, Allison Shuren, JD, Washington Counsel, SEE, allison.shuren@aporter.com, 202.942.6525.

We much appreciate your consideration of our views.

American Academy of Ophthalmology
American Society of Cataract and Refractive Surgery
American Society of Retina Specialists
Outpatient Ophthalmic Surgery Society
Society for Excellence in Eyecare