Microinjection via the Suprachoroidal Space: A Review of a Novel Mode of Administration

HIGHLIGHTS

› Administration into the Suprachoroidal Space (SCS): An Innovative Ocular Drug Delivery Technique

› Preclinical Studies of SCS Administration

› Clinical Trials Supporting SCS Administration of Triamcinolone Acetonide for Macular Edema Associated with Non-Infectious Uveitis
Microinjection via the Suprachoroidal Space: 
A Review of a Novel Mode of Administration

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OVERVIEW

This supplement to The American Journal of Managed Care® describes standard techniques for drug delivery to the eye with a focus on delivery of corticosteroids for non-infectious uveitis (NIU) or macular edema associated with uveitis (UME), introduces the suprachoroidal space (SCS) as an innovative target for therapies intended to treat the back of the eye, discusses SCS injection via microneedle, and summarizes clinical trial data supporting the efficacy and safety of SCS injection with a triamcinolone acetonide suspension in patients with UME.

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Microinjection via the Suprachoroidal Space: A Review of a Novel Mode of Administration S243

Thomas Ciulla, MD, MBA; and Steven Yeh, MD
FACULTY DISCLOSURES

These faculty have disclosed the following relevant commercial financial relationships or affiliations in the past 12 months.

Thomas Ciulla, MD, MBA

STOCK OWNERSHIP
Clearside Biomedical

EMPLOYMENT
Clearside Biomedical

Steven Yeh, MD

CONSULTANCIES OR PAID ADVISORY BOARDS
Bausch + Lomb, Adverum Biotechnologies, REGENXBIO
Introduction

A novel target for administration of ocular therapy—the suprachoroidal space (SCS)—and the advantages and utility of microneedle drug injection to access this area have attracted increased interest from ophthalmologists. Standard ocular drug delivery methods may involve systemic treatment, topical application, and delivery adjacent to the eye (periocular) or into the vitreous humor. Systemic therapy may be associated with adverse effects (AEs) due to systemic drug exposure, whereas local therapy can be limited by drug exposure to unaffected ocular tissues and by variable or low concentrations of the therapy delivered to target tissues due to the eye’s anatomy. For diseases affecting the posterior tissues of the eye, SCS administration offers compartmentalization of therapies away from anterior tissues and the potential for long-acting delivery to target tissues.

One disease that can affect posterior tissues and that is associated with considerable visual impairment is uveitis, which is characterized by inflammation of the uveal tract (ie, iris, ciliary body, and choroid) and adjacent structures. Uveitis can result from either infectious or noninfectious (eg, autoimmune) processes. The results of a claims-based analysis of data from the US OptumHealth Reporting and Insights database found that among patients with at least 2 medical claims related to a diagnosis of uveitis in 2012, 91% (4827) of 5299 uveitis cases were classified as noninfectious uveitis (NIU). The investigators also estimated that in 2015, the prevalence of NIU in the United States was approximately 121 patients/100,000 adults. However, these results may not be generalizable to the entire US population, as data analyzed were commercial insurance claims only.

The most common complication of NIU and the primary condition associated with vision loss in uveitis is macular edema (ME), a central accumulation of extracellular fluid in the retina. Although prevalence varies according to the underlying cause of inflammation, ME generally affects 8% to 30% of patients with uveitis. Uveitis associated with ME (UME) is the first ocular condition for which an SCS injection-based therapy has been approved.

This article summarizes standard techniques for ocular delivery with a focus on the delivery of corticosteroids for NIU and UME;
it then introduces the SCS as a delivery option for therapies intended to treat the back of the eye. Also included is a brief presentation of clinical data on SCS-TA, a triamcinolone acetonide (TA) injectable suspension (Xipere®, Bausch + Lomb) delivered via the SCS Microinjector® (Clearside Biomedical, Inc). SCS-TA is the first FDA-approved treatment to utilize the SCS and is indicated for the treatment of UME. 

**Techniques for Ocular Drug Delivery**

Numerous methods of drug delivery have been developed for treating diseases of the eye, and each has benefits and disadvantages. Standard methods include systemic delivery (eg, oral, intravenous, and subcutaneous routes) as well as local drug delivery methods (eg, topical eye drops, periocular or intravitreal [IVT] injections, and IVT implants) (Figure 1). Systemic delivery of drugs for ocular conditions can result in systemic AEs, such as gastrointestinal (GI) issues. Furthermore, local delivery methods may be limited by the occurrence of AEs associated with drug exposure to unaffected tissue and, in some cases, by the inability of drugs to penetrate the cornea or sclera or into the posterior segment when delivered via a topical or periocular route. More recently, SCS injection was introduced as an alternative method of ocular drug administration.

**Systemic and Topical Therapies in NIU and UME**

Systemic therapies—including steroidal and nonsteroidal anti-inflammatory drugs and biologic and nonbiologic immunomodulatory agents—are effective for treating UME but are often recommended in cases of bilateral disease or cases refractory to local therapy due to the potential for systemic complications. Even at low doses, but particularly when administered over the long term, systemic corticosteroid treatment for NIU and UME may be associated with AEs such as infection and GI disturbances; similarly, systemic immunomodulatory agents and nonsteroidal anti-inflammatory drugs confer the risk of GI issues when used alone or in combination with steroids. Topical eye drops are a mainstay of local ocular corticosteroid therapy in NIU; however, their efficacy can be limited by an inability to penetrate to posterior eye structures, which results in poor bioavailability for posterior conditions. Because eye drops may require frequent reaplication due to a high clearance rate and drug dilution at the ocular surface, patients may show poor adherence to treatment. Further, corticosteroid eye drops carry the risk of elevated intraocular pressure (IOP) and cataract development due to anterior segment drug exposure.

**Injection and Implantation Techniques in NIU and UME**

Another method for local drug delivery, periocular sub-Tenon injection of corticosteroids is administered by cannula or needle into the superior temporal or inferior temporal quadrant. However, sub-Tenon injection of corticosteroid medications requires permeation through the sclera; it can be associated with variable levels of corticosteroid delivered to the posterior segment, exacerbation of cataracts, and elevation of IOP. The results of the multicenter, randomized clinical trial (RCT) in UME (POINT; NCT02374060) indicated that sub-Tenon injection of TA—a corticosteroid commonly used to treat UME—was associated with inferior central subfield thickness (CST) reduction and visual acuity (VA) improvement when compared with IVT injection of TA and IVT implant of dexamethasone. Thus, use of alternative injection techniques may be preferred, particularly in treating UME.
direct method for administering therapeutic doses of a drug into
the posterior segment of the eye, and both can be accomplished in
an office setting.1,15 IVT delivery of corticosteroids is effective for
treating UME in the posterior segment, characteristically in patients
with moderate to severe unilateral UME or for whom immunomodu-
ulating treatments fail or are not tolerated.5,11,12 It is also used as an
interim treatment for UME in patients who later receive long-term
immunomodulating therapy.1 IVT treatment via implantation of
sustained-release technologies allows for delivery of a controlled
drug dose to a desired location over time; this reduces the need for
repeated treatments and potentially reduces the risks of infection
and retinal detachment.1 Possible complications inherent to IVT
delivery include increased IOP, endophthalmitis, retinal detachment,
and cataract development that partially arise from corticosteroid
exposure in the anterior chamber.3,5,15

The MUST RCT (NCT00132691) enrolled 255 participants (479 eyes)
with NIU between December 2005 and December 2008 and compared
efficacy and safety results in patients receiving either the 0.59 mg
fluocinolone acetonide (FA) intraocular implant (Retisert®, Bausch
+ Lomb) or systemic corticosteroid therapy supplemented with
further immunsuppression, when indicated.18,19 The FA intraocular
implant releases the drug into the intravitreal space over approxi-
mately 30 months.19 The outcomes of this trial illustrated the risks
and benefits of an IVT sustained-release corticosteroid implant in
treating NIU vs systemic therapy.18 After 24 months, significantly
more eyes treated with the IVT FA implant experienced control of
uveitic inflammation vs eyes from participants treated with systemic
steroids (P = .001). Both treatments were associated with a decreased
incidence of ME; use of the FA implant conferred a significantly
greater likelihood of reducing ME after 6 months than did systemic
corticosteroid therapy (P = .002), although the difference was no
longer significant by 24 months (P = .071). However, patients who
received the FA implant were over 4 times more likely to experience
an IOP elevation of at least 10 mm Hg or an absolute IOP of at least
30 mm Hg or to require medical or surgical intervention for elevated
IOP (all P < .0001).18 In a 7-year extended follow-up of participants
in the MUST study, those randomly assigned to receive systemic
therapy had better visual acuity than did those randomly assigned
to receive IVT FA implants; however, this interpretation is limited
by loss of patients to follow-up and to inconsistent reimplantation
in the FA implant group following depletion of implants.20

**SCS Administration: An Innovative Drug Delivery Technique**

Administration of medication to the SCS, which is located between
the choroid and the sclera (Figure 2), offers several advantages over
other methods of local ocular drug delivery by avoiding physical barriers (eg, the sclera). Under normal conditions, the SCS is in a largely collapsed state due to IOP and to fibers attaching the sclera to the choroid; however, it can expand and collapse with fluid entrance and egress. Because of its location adjacent to the choroid and, more generally, the posterior segment, the SCS offers a novel target for drug delivery, with the potential for broad distribution and high bioavailability of therapeutic agents and relatively easy access to posterior target tissues. Drugs injected into the SCS spread circumferentially; when injected into the pars plana region, they expand the SCS and quickly flow posteriorly due to the differential between the IOP and the posterior SCS pressure. In animal models, drugs had negligible dissemination into the vitreous as compared with those given via IVT injection. Because of this compartmentalization of administered drugs away from the anterior segment of the eye, SCS delivery may decrease the risk of some AEs associated with drug exposure to the anterior chamber, such as elevations in IOP and development or progression of cataracts associated with corticosteroid use.

The sclera adheres tightly to the ciliary body at the scleral spur, limiting anterior distribution of injectate. In animal models, drugs injected into the SCS were undetectable in the aqueous humor and had negligible dissemination into the vitreous as compared with those given via IVT injection. Because of this compartmentalization of administered drugs away from the anterior segment of the eye, SCS delivery may decrease the risk of some AEs associated with drug exposure to the anterior chamber, such as elevations in IOP and development or progression of cataracts associated with corticosteroid use.

Until recently, therapeutic access to the SCS generally has required surgical dissection through the sclera in the operating room with the target eye anesthetized and immobilized. This changed with the development of a proprietary SCS microneedle delivery system, the SCS Microinjector®, which consists of a piston syringe and 2 microneedles (900 μm and 1100 μm; different lengths are provided to account for patient variability in scleral thickness) (Figures 2 and 3). Administration via the SCS Microinjector® is a 3-step process, beginning with insertion of a 900-μm microneedle into the sclera at an angle perpendicular to the eye and 4 mm to 4.5 mm posterior to the limbus. The next step is application of gentle force to allow the needle hub to depress the ocular surface and form a seal between the needle hub and conjunctiva; this ensures that the needle opening can reach the SCS and limits reflux during the injection. The third step is slow injection into the SCS over 5 to 10 seconds, maintaining a perpendicular position and compression on the conjunctiva and then keeping the needle hub in place for 3 to 5 seconds after injection. If resistance occurs, injection at a new site is recommended; if resistance continues, a switch to the provided 1100-μm microneedle may allow for penetration into the SCS. The SCS Microinjector® provides more reliable access to the SCS than does a hypodermic needle, as hypodermic needle use involes a steep tactile learning curve to feel when the sclera has been penetrated and to avoid accidental IVT injection. Use of the SCS Microinjector® also is less invasive than is cannulation, as it does not require an incision with scleral dissection. In addition, SCS injection can be performed in the outpatient setting using local anesthesia.

**Sceral Thickness and Needle Length**

A loss of resistance is felt when the microneedle reaches the SCS due to the density of the collagen matrix that composes the sclera. Drug injection into the dense sclera is not possible; if the microneedle used is not long enough, the injection cannot proceed. Although thickness of the sclera can vary from anterior to posterior, thickness of the sclera at the pars plana site of injection varies only minimally between individuals and between quadrants of the eye. The microneedle in the SCS Microinjector® is designed to penetrate both the conjunctiva and sclera; the microneedle lengths were selected to make piercing of the choroid or retina unlikely while addressing the known minimal variability in sceral thickness.

Two post hoc analyses examined patient and clinical characteristics associated with SCS administration of SCS-TA via the 900-μm microneedle (Table 1). In a post hoc study of data from 6 clinical trials involving 3 different ocular disorders affecting the posterior segment (NIU, diabetic ME [DME], and retinal vein occlusion [RVO]), a 900-μm microneedle was used to administer SCS-TA in 70.9% of the procedures, and the remaining procedures were carried out with the 1100-μm microneedle. Statistically significant correlations were found between needle length and patient gender (P = .006) and needle length and quadrant of administration (P = .0005) (Figure 4), although further study is needed to determine whether these correlations may be used to guide preselection of needle length. There was no significant difference in needle length used among participants with NIU vs other disorders (P = .504).

A separate post hoc analysis of SCS Microinjector® needle length used in 2 clinical trials of patients with NIU was presented at the
2021 Association for Research in Vision and Ophthalmology Annual Meeting (ARVO 2021). Treatment was completed with the 900-μm microneedle in 74% of injections, and the 1100-μm microneedle was used for the remaining injections. No correlation was found between needle length and gender, although significant relationships with needle length were observed for injection quadrant and disease duration ($P = .012$ and $P = .037$, respectively). A modest inverse correlation between age and needle length also was observed. The 2 post hoc analyses on needle length usage demonstrated that the 900-μm microneedle can be used for the majority of SCS-TA injections. However, it is not yet clear whether statistical differences observed in either of the post hoc analyses are sufficiently meaningful to guide preselection of needle length.

### Preclinical Studies of SCS Administration

For small-molecule suspensions, drug particle size and the relative insolubility of molecules are important factors driving drug durability following SCS administration. The results from 1 preclinical study found that drug particles ranging from the size of gene therapy viral vectors to small-molecule suspensions (eg, 10 nm to 10 μm) remained primarily in the SCS and choroid for a period of months. The physiologic upper limit of pore size in the fenestrated choriocapillaris is estimated to be between 6 nm and 12 nm, which likely limits transcapillary passage of particles.

### Animal Model Studies of SCS Injection

In porcine and canine ex vivo models, fluid volume of up to 1 mL injected via cannulation into the SCS resulted in rapid local SCS expansion before flowing posteriorly. The potential for a high level of bioavailability after SCS injection extends beyond the choroid to the retinal pigment epithelium (RPE) and, ultimately, to the retina. In a rabbit model, SCS injection of a solution containing micronized axitinib compounded with a wetting agent, a tonicity agent, and a viscosity-enhancing agent resulted in much greater drug exposure in the retina, choroid, and posterior sclera and substantially decreased drug exposure in the vitreous humor vs IVT injection. After SCS injection, axitinib levels in the plasma and aqueous humor were undetectable. The utility of the SCS as a location for drug administration was further confirmed by the observation that a single SCS injection produced sustained and high levels of axitinib in the RPE-choroid-sclera (RCS) and retina. After 6 months, in a separate rabbit toxicology study, mean axitinib levels in the RCS and in the retina were 3 to 5 log orders higher than the in-vitro concentration of an agent needed to inhibit VEGF receptor 2 by 50%, further supporting the potential for long-acting drug delivery to posterior segment tissues through SCS injection.

Additionally, a series of studies in rabbits compared drug distribution following IVT and SCS injection of the TA injectable formulation for injection into the SCS.

### TABLE 1. Key Results from Post Hoc Analyses on Completion of SCS-TA Injection via SCS Microinjector®

<table>
<thead>
<tr>
<th></th>
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</tr>
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<tbody>
<tr>
<td>Studies included</td>
<td>2 studies in NIU:</td>
<td>2 studies in NIU:</td>
</tr>
<tr>
<td></td>
<td>• Phase 3 AZALEA (NCT03097315)</td>
<td>• Phase 3 AZALEA (NCT03097315)</td>
</tr>
<tr>
<td></td>
<td>and PEACHTREE (NCT02595398)</td>
<td>and PEACHTREE (NCT02595398)</td>
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<tr>
<td></td>
<td>1 study in DME:</td>
<td></td>
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<td></td>
<td>• Phase 2 TYBEE (NCT03126786)</td>
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<td></td>
<td>3 studies in RVO:</td>
<td></td>
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<tr>
<td></td>
<td>• Phase 2 TANZANITE (NCT02303184)</td>
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<td></td>
<td>• Phase 3 SAPPHIRE (NCT02980874)</td>
<td></td>
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<tr>
<td></td>
<td>and TOPAZ (NCT03203447)</td>
<td></td>
</tr>
<tr>
<td>Number of baseline injections</td>
<td>581</td>
<td>133</td>
</tr>
<tr>
<td>Overall procedures completed with the 900-μm microneedle, %</td>
<td>70.9%</td>
<td>74%</td>
</tr>
<tr>
<td>Characteristics significantly associated with completion of SCS-TA injection using a 900-μm microneedle</td>
<td>Gender ($P = .006$)</td>
<td>Administration quadrant of eye ($P = .012$)</td>
</tr>
<tr>
<td></td>
<td>• Female, 76%</td>
<td>• Into the superior temporal quadrant, 82%</td>
</tr>
<tr>
<td></td>
<td>• Male, 66%</td>
<td>• Into the superior nasal quadrant, 45%</td>
</tr>
<tr>
<td></td>
<td>Administration quadrant of eye ($P = .0005$)</td>
<td>Disease duration ($P = .037$)</td>
</tr>
<tr>
<td></td>
<td>• Into the superior temporal quadrant of the eye, 78%</td>
<td>• Limited (≤3 months) disease duration, 91%</td>
</tr>
<tr>
<td></td>
<td>• Into the inferior temporal quadrant, 65%</td>
<td>• Persistent (&gt;3 months) disease duration, 70%</td>
</tr>
</tbody>
</table>

DME, diabetic macular edema; NIU, noninfectious uveitis; RVO, retinal vein occlusion; SCS, suprachoroidal space; SCS-TA, triamcinolone acetonide suspension formulation for injection into the SCS.
suspension formulation Triesence (TRI; FDA approved for uveitis and several other ocular diseases) and following SCS injection of 2 different TA formulations, including TRI and an investigational prototype of a TA injectable suspension for use in the SCS (X-TA).8,30

In the first of these 2 studies, 50 rabbits received TRI as bilateral IVT injections or as bilateral SCS injections to compare drug concentrations in ocular tissues and plasma after drug delivery. At day 91 (end of study), the levels of TA in the RCS were 12-fold higher with SCS injection than with IVT injection. Further, SCS injection resulted in lower levels of TA in the vitreous and aqueous humors, iris/ciliary body, and lens compared with IVT injection over the duration of the study.8

In the second study, 20 rabbits were randomly assigned to receive bilateral SCS injections using either TRI or X-TA with the SCS Microinjector®. No TA was detected in the aqueous humor throughout the 91-day study; on day 15, minimal levels of TA were observed in the vitreous humor of eyes receiving SCS injection of either formulation, whereas comparable levels of TA were detected in the RCS for X-TA vs TRI throughout the study.8

Results from these 2 studies showed that SCS injection resulted in durable TA levels in posterior tissues over the course of 3 months. Furthermore, injection into the SCS resulted in TA sequestration away from anterior tissues. Thus, these results support a potentially lower risk of IOP elevation and cataract development associated with anterior tissue exposure to TA.8

Simulated SCS Injection
The previously mentioned X-TA suspension prototype is a proprietary formulation of TA designed for SCS microinjection; it contains surfactants chosen to improve wetting, reduce the coefficient of friction, and minimize foaming and the formation of microbubbles.8

To examine injectability of this TA formulation specifically designed for SCS delivery, investigators assessed the glide force—the required pressure applied to the SCS Microinjector® plunger to expel the formulation—of TRI vs X-TA using the SCS Microinjector® in a series of 7-second simulated microinjection procedures.8 Injections of air and water were included as controls. Glide force was significantly higher with the TRI formulation vs X-TA (P < .001), and TRI was associated with greater variability in glide force than was X-TA (Figure 5).8 The glide force profile for X-TA was notably closer to the glide force for injections of air and of water than was that of TRI (0.73 N vs 0.19 N, 0.23 N, and 1.31 N, respectively). Less force was needed to overcome the syringe-plunger’s resistance to flow when injecting X-TA vs TRI; for the physician, this may provide a more distinct tactile signal that the microneedle tip has entered the SCS or that a change to the 1100-μm microneedle

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**FIGURE 4.** Correlation of Needle Length Used in SCS Injections With Injection Location and Patient Gender in Post Hoc Study86

![Graph showing correlation of needle length used in SCS injections with injection location and patient gender in post hoc study.](image-url)

- **Percent of participants receiving injection**
  - Superior temporal: 78%
  - Inferior temporal: 65%
  - Superior nasal: 50%
  - Inferior nasal: 42%
  - Female: 76%
  - Male: 66%

- **Injected quadrant of eye**
  - Superior temporal: 22%
  - Inferior temporal: 35%
  - Superior nasal: 50%
  - Inferior nasal: 38%

- **Gender of participant**
  - Female: 24%
  - Male: 34%

**SCS**, suprachoroidal space.

is needed. Compared with that of TRI, the particle size of X-TA is smaller; this difference could contribute to reduced variability and lower necessary glide force when injected through the 30-gauge needle of the SCS Microinjector®. Consequently, the use of a TA formulation specifically designed for SCS injection may help to achieve controlled and consistent delivery due to lower necessary injection force and less variability in applied pressure as a result of expedient choice and tightly controlled particle size distribution of the drug.8

Clinical Trials Supporting SCS Administration of SCS-TA in NIU and UME

The anti-inflammatory activity of TA is well known, and the drug has been extensively studied for the treatment of NIU and UME via periocular or IVT injection.9,10 However, SCS-TA offers potential advantages in terms safety, efficacy, and ease of injection.8,14,32 FDA approval of SCS-TA to treat UME via SCS administration was based on the results of the phase 3 PEACHTREE trial (NCT02595398) (Table 2).14,32-34

PEACHTREE was a 24-week, international, multicenter study conducted between November 2015 and January 2018 in 160 participants with UME who were randomly assigned 3:2 to receive a single injection of SCS-TA via SCS Microinjector® (n = 96) or a sham SCS injection procedure (control; n = 64) at day 0 and again at week 12; follow-up visits occurred every 4 weeks for the duration of the trial, and efficacy end points were assessed at 24 weeks.14 Compared with participants in the control arm, significantly more participants in the SCS-TA arm gained at least 15 Early Treatment Diabetic Retinopathy Study (ETDRS) letters (≥3 lines) in best-corrected VA (BCVA) (P < .001). In addition, participants in the SCS-TA arm achieved a significantly greater mean reduction in CST compared with those in the control arm (P < .001). Although more participants in the SCS-TA arm than in the control arm experienced elevated IOP at the time of procedure, fewer experienced elevated IOP related to corticosteroid use (ie, events occurring after the day of the procedure). Finally, cataract AE rates were similar between the SCS-TA group and the control group.12

A post hoc analysis of outcomes from the PEACHTREE trial compared the 83 participants in the SCS-TA arm (86.5%) who completed the study without requiring rescue therapy with the 46 participants in the control arm (71.9%) who required rescue therapy (of whom 37 [80.4%] received IVT injection or periocular corticosteroids).36 Unrescued SCS-TA participants had a lower incidence of elevated IOP (≥30 mm Hg) than did rescued control participants (4.9% vs 10.9%, respectively); further, the incidence of IOP-lowering medication use was lower in unrescued SCS-TA participants than in rescued control participants (7.2% vs 13.0%, respectively).36

The MAGNOLIA study (NCT02952001) was an international, multicenter, observational extension study of 33 participants who successfully completed the PEACHTREE study without the use of rescue medications (Table 2).14,32,34 Relative to the first dose of study drug in PEACHTREE, patients in the SCS-TA arm had a median time to rescue therapy of 344.0 days.33 In addition, patients in the SCS-TA arm not receiving rescue therapy showed a significant gain in mean BCVA letters and a significant reduction in mean CST from the PEACHTREE baseline to week 24 of MAGNOLIA (48 weeks from PEACHTREE baseline; P = .004 and P = .001, respectively).35

An additional post hoc analysis used pooled data from the 33 participants in MAGNOLIA and the 127 participants from PEACHTREE who were not enrolled in MAGNOLIA to determine the probability of rescue medication use over the entire 48-week period.35 The median time to rescue relative to the last dose of study drug was significantly longer among the participants

![Glide Force Over Time for X-TA and TRI](chart)

**Figure 5.** Average Glide Force Over Time for X-TA and TRI During Syringe Compression Using an SCS Microinjector®

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**Table 2.**

<table>
<thead>
<tr>
<th>End Point</th>
<th>Control Arm (%)</th>
<th>83 Participants (%)</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>AE Incidence</td>
<td>17.2</td>
<td>68.9</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>Rescue Therapy Use</td>
<td>43.3</td>
<td>8.7</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>Efficacy</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mean Reduction in CST (lines)</td>
<td>3.7</td>
<td>6.4</td>
<td>&lt;.001</td>
</tr>
<tr>
<td>Mean BCVA Gain (letters)</td>
<td>10.9</td>
<td>13.5</td>
<td>&lt;.001</td>
</tr>
</tbody>
</table>

**S249**

**MICROINJECTION VIA THE SUPRACHOROIDAL SPACE**

**SCS, suprachoroidal space; TRI, Trience [triamcinolone acetonide injectable suspension]; X-TA, triamcinolone acetonide suspension formulation prototype for injection into the SCS.**

*Twelve replicate injections per group.

*Error bars (mean ± standard error) are present in each of the groups assessed and are standard for these types of measurements.

treated with SCS-TA than among the participants in the control arm (257.0 vs 55.5 days, respectively), and the time to rescue distributions were significantly different ($P < .001$).33

Finally, the open-label, prospective, multicenter AZALEA safety trial (NCT03097315) evaluated rates of serious AEs (SAEs) and treatment-emergent AEs (TEAEs) in 38 participants with NIU (20 participants with ME; 18 participants without ME) who were treated with SCS-TA given via SCS Microinjector® (Table 2).14,32-34

Similar to dosing in PEACHTREE, AZALEA participants received a dose of SCS-TA at baseline and at 12 weeks, with follow-up every 4 weeks to 24 weeks.44 Mean IOP was 13.3 mm Hg at baseline and 15.2 mm Hg at week 24 in the study eye. Six participants (15.8%) had an IOP elevation of more than 10 mm Hg compared with baseline in the study eye, and 2 participants (5.3%) had an IOP of more than 30 mm Hg. Cataract formation AEs were reported in 4 study eyes (10.5%); 1 of which was deemed related to treatment. Seven participants experienced a TEAE considered to be related to the study drug by an investigator, and no SAEs involving the study eye were reported.14

**Post Hoc Analysis on the Safety of SCS Injection of SCS-TA in Different Ocular Diseases**

A post hoc analysis presented at ARVO 2021 used pooled data from 8 clinical trials of SCS-TA in NIU, DME, or RVO and compared the safety of SCS-TA monotherapy via the SCS Microinjector® ($n = 166$) vs a combination of SCS-TA injection plus IVT anti-VEGF injection ($n = 460$) vs sham SCS injection plus IVT anti-VEGF injection (control; $n = 449$).37

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**TABLE 2. SCS-TA: Key Clinical Trial Results**

<table>
<thead>
<tr>
<th>Study name</th>
<th>PEACHTREE (N = 160)</th>
<th>MAGNOLIA (N = 33; 28 active participants)</th>
<th>AZALEA (N = 38)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Study design</strong></td>
<td>Phase 3 international, multicenter, controlled trial in participants with UME randomly assigned to SCS-TA or sham procedure</td>
<td>Phase 3 international, multicenter, observational extension to PEACHTREE; participants received no additional treatment (except rescue therapy, when necessary)</td>
<td>Open-label, multicenter, prospective safety trial of SCS-TA in participants with NIU with and without ME</td>
</tr>
<tr>
<td><strong>Time frame</strong></td>
<td>Baseline to 24 weeks</td>
<td>24-week extension to PEACHTREE</td>
<td>Baseline to 24 weeks</td>
</tr>
</tbody>
</table>
| **Key Results**    | Gain of $\geq 15$ ETDRS letters in BCVA  
  - Active: 46.9%  
  - Control: 15.6% ($P < .001$)  
  Mean change in ETDRS letters  
  - Active: +13.8  
  - Control: +3.0 ($P < .001$)  
  Mean change in CST  
  - Active: $-152.6 \ \mu m$  
  - Control: $-17.9 \ \mu m$ ($P < .001$)  
  Elevated IOP  
  At time of procedure  
  - Active: 8.3%  
  - Control: 0%  
  Post-procedure*  
  - Active: 11.5%  
  - Control: 15.6%  
  Cataract  
  - Active: 7.3%  
  - Control: 6.3%  
  Median time to rescue therapy relative to first treatment in PEACHTREE:  
  - 344.0 days  
  Mean change in ETDRS letters in BCVA  
  - Crossover: +16.8 ($P < .001$)  
  - Week 48: +12.1 ($P = .004$)  
  Mean change in CST  
  - Crossover: $-178.1 \ (P < .001)$  
  - Week 48: $-174.5 \ (P = .001)$  
  Number of participants with TEAEs or serious ocular AEs  
  - TEAEs: 7 (18.4%)  
  - Serious ocular AEs: 0 |

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AE, adverse effect; BCVA, best-corrected visual acuity; CST, central subfield thickness; ETDRS, Early Treatment Diabetic Retinopathy Study; IOP, intraocular pressure; ME, macular edema; NIU, noninfectious uveitis; SCS-TA, triamcinolone acetonide suspension formulation for injection into the suprachoroidal space; TEAE, treatment-emergent AE; UME, macular edema associated with uveitis.

*Rates of ocular AEs reported in this table differ slightly from data in the prescribing information due to differences in AE categories used in the FDA analysis.

Five participants who received a sham procedure were also followed through the extension study, as enrollment occurred prior to unmasking. Efficacy results are presented for active participants only.

Includes all events of elevated IOP that did not occur on the day of the procedure.

Crossover occurred at week 24 of PEACHTREE.
Of the 626 participants receiving at least 1 SCS-TA injection, 3 participants (all of whom received multiple injections) experienced ocular SAEs of interest (1 retinal detachment in a quadrant different from and 8 weeks after SC injection, and 2 cases of reduced vision ≥ 1 week after SC injection), although none of the events were deemed related to treatment. There were no SAEs involving retinal tear without detachment, lens injury, supachoroidal hemorrhage, endophthalmitis, or central retinal artery occlusion in patients receiving SCS-TA monotherapy or combination therapy. Additionally, the AE rates of eye pain experienced by patients with NIU, RVO, or DME who received SCS injections of SCS-TA were similar to those reported in registration trials of IVT anti-VEGF agents approved for ocular diseases. These results support SCS injection of SCS-TA as offering acceptable safety and incidence of eye pain compared with IVT injection.

**Discussion**

The development of a microinjector device optimized for drug delivery to the SCS addresses an unmet need in the treatment of ophthalmic diseases affecting the posterior segment of the eye. Other delivery methods have demonstrated efficacy in treating ocular diseases such as NIU and UME, but each of these approaches has been associated with considerable risk of AEs, and efficacy may be affected by limitations inherent to the eye’s anatomy. Such limitations are largely a consequence of the challenges associated with the eye’s anatomy and the difficulty both of delivering the appropriate levels of drug to the correct target areas and of avoiding distribution to locations where it might induce undesirable effects. Moreover, the structure of the SCS enables a degree of drug sequestration away from the iris/ciliary body, lens, and aqueous humor, thereby limiting drug exposure to anterior tissues. Previous methods of accessing the SCS often required an operating room setting with immobilization of patient’s eye under anesthesia and surgical dissection through the sclera. The SCS Microinjector® was designed specifically to access the SCS safely and easily using only local anesthesia.

Results from the PEACHTREE study, along with those from the MAGNOLIA and AZALEA studies, offer compelling evidence for the safety of SCS injection with the SCS Microinjector® and the effectiveness of SCS-TA delivered via the SCS Microinjector® to treat UME. The broader implication of improvements in BCVA among participants with UME in PEACHTREE are illustrated by results of an analysis conducted by the MUST Trial Research Group on the relationship between improvements in VA and visual function outcomes. A 1-line improved BCVA score (equivalent to the difference between 20/20 and 20/25 or 20/160 and 20/200 scores on the Snellen scale) was associated with significantly higher scores for general vision, driving, color and peripheral vision, and vision-specific social functioning, mental health, and dependency on the National Eye Institute Visual Function Questionnaire (all p < .0001). In the PEACHTREE study, 46.9% of participants treated with SCS-TA experienced a BCVA improvement of at least 3 lines.

SCS delivery of various drug formulations, including small-molecule suspensions and viral-vector therapies, has been evaluated in clinical trials to treat several disease states. Therapeutic agents currently under investigation for SCS delivery include the tyrosine kinase inhibitor axitinib to treat neovascular age-related macular degeneration (nAMD; NCT04626128), a viral-vector gene therapy to treat nAMD (NCT04514653) and diabetic retinopathy (NCT04567550), and a virus-like drug conjugate to treat primary indeterminate lesions and small choroidal melanoma (NCT04471530).

**Conclusions**

SCS-TA is the first drug to be FDA-approved for SCS administration. The combination of SCS-TA formulation and SCS Microinjector® represents a meaningful addition to the therapeutic landscape for treatment of UME. Additionally, the SCS Microinjector® provides new opportunities for treating ocular diseases with an SCS drug delivery procedure in the office-based setting, offering the potential for broad use across several ocular diseases with delivery of different drugs.

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**Authorship information:** Concept and design (TC and SY); acquisition of data (SY); analysis and interpretation of data (SY); critical revision of the manuscript for important intellectual content (TC and SY); supervision (TC).

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