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PROSTHETIC & ORTHOTIC MAGAZINE

ISSUE 9 • SUMMER 201

Patient Profile

This Little
Warrior Wears
His Helmet
Proudly and Wins!



i-limb™ Continues to Improve

Battle Continues for Legislation to Benefit O&P Providers and Patients

Patellar Tendon Brace Considered to be Underutilized Orthotic Device







Dexterity and Function

ince the i-limb's inception, Touch Bionics has been building and improving on their concept of motorized components in each of the 5 digits. Historically, prosthetic hands have had motorized components in the thumb, index, and middle finger, but thanks to new developments, the i-limb has taken the functionality of prosthetic hands to a higher level. Patients are now capable of preforming tasks they never thought could be possible.

Like most myoelectric hands, the i-limb is controlled by muscle contractions and electrodes positioned properly on predetermined areas around the residual limb. The muscle contractions are detected by the electrodes, which cause the prosthetic hand to respond. As the user's ability to control the muscle contractions improves, the prosthetic hand can be programmed to maximize the dexterity and function.

Some Unique Features of the i-limb

- 1. Gesture Control This allows the user's pre-programmed grip options to change with one of four simple gestures. (Moving the i-limb in one of four directions or gestures activates the grip option.)
- 2. App Control In any given day users may require numerous different grip options. Users can select up to 24 pre-programmed grip options and customize an additional 12 grips using the my i-limb app on their mobile device, which is synched to their prosthetic hand.
- 3. Proximity Control Bluetooth-enabled devices called "grip

chips" can be placed near objects to activate a grip when the i-limb is placed close to the chip. For example, grip chip could be placed close to your computer keyboard to automatically enable a grip with the index finger pointed keyboard typing. 🦋







ressed for battle, 10-month old Paul Hassler is happy, confident, and winning his war with plagiocephaly and torticollis, thanks to the otherworldly headgear he will soon outgrow. Cheryl Hassler, Paul's mother, experienced the same perplexity and concern as any new mother when she recognized a flat spot on the right side of his head at birth (plagiocephaly). Following the doctor's advice, his parents kept an eye on it for a time, and noticed that Paul's head was also tilted to the right side.

Born a twin, Paul's cramped position in the uterus he shared with his brother may have resulted in a tightness in the muscle that connects the breastbone and collarbone to the skull, his parents discovered. This condition (torticollis) caused Paul to continue to tilt his head, and to preferentially lie with his head on that side—increasing the tendency of the plagiocephalic flat spot to develop further.

While the flat spot itself was not an issue, Cheryl also noticed what appeared to be a "lazy eye" on his right side. The physical therapist to whom they were referred, however, diagnosed the condition promptly, identifying a 10 mm rightward shift of his eye and ear.

The Hasslers eventually found their way from their

home in Coatesville, PA, to Lawall Prosthetics and Orthotics, where orthotist Harry Lawall III, CPO discussed the need for a cranial orthotic—a custom-created helmet with shape-correcting capabilities.

Left unchecked, the danger of the rightward shift of Paul's eye and ear continuing was very real. At that time, when he was nearly seven months old, his right eye and ear had shifted more than half an inch, according to Lawall.

"He started at 5/8 inch of a diagonal difference, which is a little above average for this condition. Our goal by the end of treatment is to get the diagonal difference down to about 1/4 inch," he explained.

Since experts believe that a baby's skull resists reshaping as the bone thickens, 12 months of age is generally regarded as the "deadline" date beyond which such therapies are ineffective.

After just two months of treatment, however, Paul's measurements indicated that his deviation from center had then been corrected to only 3/8 inch—already a 1/4 inch improvement from the original measurement—and was now only 1/8 inch away from their goal.

"He's been wearing the helmet for five weeks now," Cheryl confirmed, "and the first time he was re-measured, he was down 1/8 inch, and this time he was also down another 1/8 inch, so it's working. Harry (Lawall) thinks it should be off by Paul's first birthday—which is in October. It all depends on the child, but in Paul's case, he's taken to it and is doing great."

From the first time he donned the helmet she made a game, telling him he was a football star in training. "To this day, when I put it on, I'll use my nails and tap it and say, 'Is it on? Is it okay?' He even reaches for it and wants it back on when he comes out of the pool," she marveled.

Top Technology to the Rescue

Lawall Prosthetics & Orthotics has been a leader in cranial orthotics since receiving one of the first twenty scanners in the country back in 2003. The company recently added state-of-the-art SmartSoc™ 3D scanning systems to three of three of their office locations.

"Paul was one of my first scanned patients," said Lawall, "which made the fitting much easier than doing a traditional plaster mold of his head shape, and allowed less room for error in the final fitted helmet. A bad scan is better than a great cast, they claim, and although we are still seeing great results with the casts, fitting pediatric patients with a scanner is just less taxing for both the family and the orthotist."

The anxiety level of a young child can be greatly increased by being wrapped in plaster, he pointed out. "With the scan, I just take 12 pictures of the head, rotating around it, and we're done. It takes about two minutes."

The most advanced device is only half the battle, however. A lot depends on compliance, Lawall said.

"Paul is a very compliant patient, so we're seeing good results. These helmets also require a lot of follow-up visits; it's a big commitment for the families. We're constantly measuring, constantly making sure that the helmet is fitting correctly and needed adjustments are made promptly."

Instead of creating a new helmet each time to keep up with the child's growth, he explained, the original helmet is designed to accommodate anticipated growth, with interior layers that can be removed as necessary.

"We include about four layers of padding; then, as the head grows and begins to reshape, we take out layers in the flat area to promote growth in that quadrant.

The helmet, which must be worn 24 hours a day except when bathing or in the pool, is something the child adjusts to gradually.

"The first day it was only an hour, then we increased it and now he's up to 23 hours a day. He sleeps in it," Cheryl noted. "It was kind of weird, but in our case, he seemed to

just take to the helmet, and didn't mind wearing it, so he was a good patient, really."

Concerns and Discoveries

She confessed to the natural anxiety of a mother faced with fitting an alien device on her baby's head, initially blaming herself for Paul's condition before learning that twins are simply more vulnerable to this condition due to the added pressures of a crowded womb where movement is constricted; it's not predictable or preventable.

Premature singletons are also more susceptible to positional plagiocephaly, she learned—because while kept in hospital care, they are typically placed on their backs; pressure on the back of their skulls can allow a flat spot to develop.

(Paul and his twin, Cody, were born five weeks prematurely, and spent their first five days in the neonatal intensive-care unit (NICU).)

"We also had anxieties about what Cody was going to think about the helmet —because they were used to seeing each other every day," Cheryl recalled. "Believe it or not, it doesn't bother him. He gave Paul a weird look the first day, but that was it. Then he just kind of went 'Whatever,' and moved on. They both adapted to it without any issues."

The helmet weighs about a pound, and the added weight may play a role in correcting the effects of the torticollis, helping to strengthen the neck and address any muscular deficit there.

Although it's not difficult to put on and take off the helmet, it took a few days for Cheryl and her husband Bill to learn and get comfortable with the process.

"Sometimes it needs adjusting; Harry just added some more padding to the back because the helmet kept shifting forward," she said. "It also needs simple daily maintenance, in the form of cleaning and disinfecting with alcohol."

There are other small adjustments that become necessary, she reported. "When we feed him a bottle, he has to lay his head in a certain way; I'd rather have him in a comfortable position than me."

Their greatest challenge, however, has been the ex-

"Because he is so little, his body has to work harder to regulate the temperature, and he sweats badly inside the helmet. We've been trying to keep him inside, in the air conditioning, on these ungodly hot days."

Dressed for Success

Does wearing the helmet slow Paul down? -hinder de-(Continued on page 8)

Battle Continues for Legislation to Benefit O&P **Providers and Patients**

t has been more than a year since the O&P community (made up of providers and patients) united like never before to oppose the Draft Local Coverage Determination (LCD) bill for lower limb prostheses. Since then, there have been a couple more important instances that have alerted key congressional offices to the fact that the two groups are aligned on the majority of issues when it comes to healthcare policy.

Generally speaking, what benefits patients usually benefits providers. Therefore, while providers and patients continue to lobby together on most issues, health plans, businesses, and other payers typically take the opposite stance.

This common interest between providers and patients was on full display recently, when this past spring, the Amputee

held Coalition Congressional Lobby Day and the following American week the Orthotic & Prosthetic Association (AOPA) hosted a Policy Forum. These two events were designed to educate members of congress on the needs within the O&P field and also focus the field's efforts to fight for the best orthotic and prosthetic coverage.

On the Amputee

Coalition's Lobby Day, more than 25 leaders of the O&P profession met in Washington, DC to educate members of Congress on issues effecting the field and its patients. The key issues discussed were the LCD bill and the federal fairness bill (formerly known as parity). This bill is centered on the idea of requiring insurance plans to cover orthotics and prosthetics as they cover other medical benefits. For example, it would prohibit arbitrary limits or exclusions of devices. The federal fairness bill is focusing on getting the nearly 30 states that don't have parity to pass this legislation. The federal fairness bill is also looking to address the fact that large businesses that self-insure their employees are exempt from following state insurance laws.

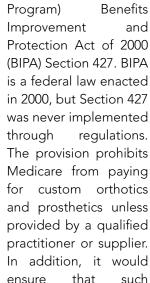
AOPA's Policy Forum was well attended by practitioners, patients and other leaders within the O&P field. Former Senator Bob Kerrey set the tone of the forum and the end result was a one-page bill that listed the key O&P policy priorities.

"Those priorities were as follows:

Separation of Orthotics & Prosthetics from Durable Medical Equipment - This would separate orthotics & prosthetics (O&P) from durable medical equipment (DME) in federal law and regulations to ensure that the O&P field does not continue to be subjected to DME-based restrictions.

Provider Qualifications and Recognition of Clinical Notes

- This would require implementation of the Medicare, Medicaid and SCHIP (State Children's Health Insurance



practitioner's clinical notes are recognized as part of the patient's medical record for purposes of determining medical necessity.

Expansion of Competitive Bidding - This item opposes efforts to further expand competitive bidding to all prosthetics and orthotics, as the Obama Administration has proposed in its most recent federal bidding proposals, and presses for clarification that competitive bidding only applies to off-theshelf orthotics that are subject to minimal self-adjustment.

Draft LCD for Lower Limb Prostheses - This would rescind, or at least place a moratorium on issuance of the Draft LCD for Lower Limb Prostheses and ensure that any future LCD is



developed in a transparent way with robust stakeholder input to ensure it does not restrict access to appropriate patient care."

Separation from the umbrella of DME care would be a great step because it would lay the groundwork for many desired policy changes. None of the four points highlighted in this bill are new ideas. They are concepts that have been proposed by various O&P organizations for years. The advantage now is the strong level of engagement and participation the whole O&P community had in creating the bill and the focus on overall reform concepts. Former Senator Kerrey took the process a step further by meeting with former colleagues in the senate to help push the bill forward. In the days following Kerrey's meetings O&P community members met with their Congressional leaders to increase the impact. At this point it seems if Congress is able to pass a Medicare bill, the O&P bill will also go through. But the reality of Congress being able to pass a Medicare bill is debatable.

Thomas, Peter W. "How Patients Benefit From O&P Policy Reforms" The O&P Edge, June 2016.

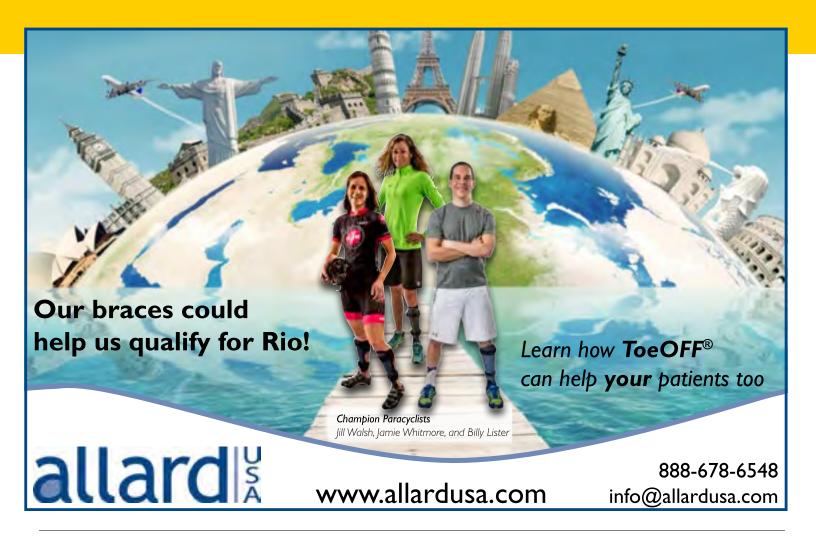
Certified Orthotist Joins Lawall's Hershey, PA Staff



Erin Aulicino, CO, LO recently joined the Lawall **Prosthetics Orthotics** team, and is working at our Hershey, PA facility located the Penn Hershev Medical Center. She received her Masters of Science in Prosthetics 8 **Orthotics** from the

University of Pittsburgh and worked as an orthotist for five years in Pittsburgh.





(Patient Profile continued from page 5)

This Little Warrior Wears His Helmet Proudly – and Wins!

velopment or activity levels?

Cheryl laughed. "Not so you'd notice! He's not only keeping up, he's surpassed his twin! We have them in walkers and he is a crazy child! He runs into everything at full steam ahead. The other day he actually crashed through a baby gate, then looked back at me and laughed, like 'Ha-ha! I got through it!'

"We wondered if it's giving him a false sense of security, since now he whacks his head and doesn't feel anything!"

Lawall acknowledges that families are sometimes concerned because their child has been wearing the helmet for six months, and has grown accustomed to having this 24-hour protective barrier in place.

"After they're done with treatment, sometimes the helmet will still fit, so we can do a weaning off process," he explained. "You leave the helmet on for two hours, then take it off for two hours—just so they get accustomed to having it off."

The bottom line, however, is that it's working, said Cheryl, and it's working much more quickly than she had expected. "People who haven't seen Paul for a while can actually see the difference, now. I was expecting him to make slow and gradual progress, but this, to me, is going fast!"

Advice to Other Parents of Plagiocephalic Children:

Cheryl and Bill Hassler stress the importance of (a) finding an experienced, progressive and trustworthy orthotic provider to guide and support you and your child through the process, (b) researching the subject and asking your doctor lots of questions—and above all, (c) not hesitating, but moving forward as quickly as possible after the diagnosis.

"The Lawall folks have absolutely been helpful—just terrific to us," Cheryl noted. "Harry explained to us that although there will be some shifting in shape as the child's head grows, once it reaches a certain point it will not correct itself. So you only have that brief window of opportunity and need to make your decision in time. Waiting to see what happens is a good way to run out of time. Don't hesitate. Get it done."

Bill agreed strongly, "You are your child's advocate right now. We didn't want Paul to have issues as he was older, if we did not have it corrected. Your child will have to live with your decision for a long time, so it ought to be the right one."

"I would highly recommend the cranial helmet to anybody whose child has this problem," added Cheryl. "Don't hesitate to talk to your doctor about it, and don't hesitate to get it. I've seen firsthand what it's doing for my son." 💓

Lawall Practitioner Attends International Symposium

att Moran, CPO, of Lawall Prosthetics & Orthotics, attended the First International Symposium on Innovations in Amputation Surgery & Prosthetic Technologies (IASPT). This was a two-day event held at the Rehabilitation Institute of Chicago for surgeons/physicians, prosthetists, physical and occupational therapists. The main objective for prosthetists at this symposium was to instruct them on advanced prosthetic issues, particularly for patients with unique surgeries and implantable devices.

In recent years, many innovative surgical techniques and technologies have been developed that have the potential for greatly enhancing the quality of life for persons with limb loss. With the field's rapid evolution, it is essential that all surgery for people with amputations is performed in light of current and coming technologies. Therefore, the symposium also focused on promoting recent innovations regarding upper and lower limb surgeries, therapies and prostheses.

Some of the topics that were discussed were implanted sensory motor systems, osseointegration/direct skeletal attach-

ment, Targeted Muscle Reinnervation (TMR) for treatment of painful neuromas and for enhanced prosthesis control, and residual limb lengthening. One of the most exciting parts of the course was the patient demonstration. Attendees had the opportunity to see and speak directly with patients who were actually using these new cutting edge technologies.

Lawall strongly encourages our practitioners to continually seek continuing education within the field. It is through participation in seminars and classes that we can better serve our patients and ensure them they are continuously being offered the latest technology available. **







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A Frequently Overlooked Orthotic Option: The Patellar Tendon Brace

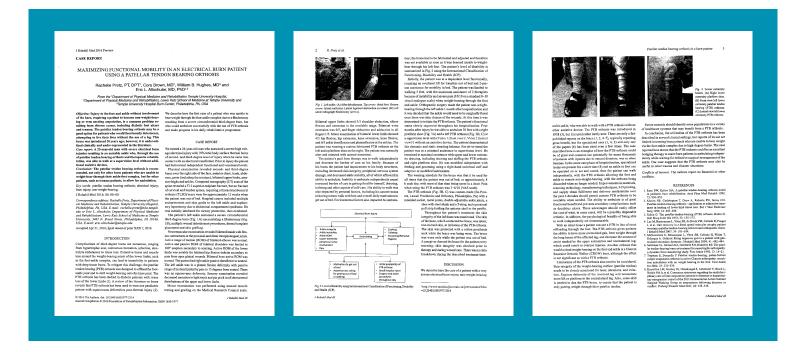
awall practitioner, Bo Powers, CPO, had the privilege of assisting Rachelle Pretz, PT, DPT, Cora Brown, MD, William Hughes, MD, and Eric Altschuler, MD, PhD, of Temple University Hospital in a case study of a patient who suffered from an electrical burn and was prescribed a Patellar Tendon Brace (PTB).

The purpose of this study was to determine if the PTB offloaded the patient's foot and ankle enough to allow healing while promoting mobility. Patellar Tendon Braces are custom braces designed to offload the foot and ankle by dispersing pressure to the patellar tendon and popliteal area of a patient's leg. The PTB was designed in 1958, but it is currently underutilized within the field of orthotics.

The candidate for this case study was a 26-year-old male, who suffered a high voltage electrical injury resulting in 70% of his body having 2nd and 3rd degree burns. "The patient's left ankle sustained a severe circumferential third degree burn necessitating a fibulectomy; multiple wound debridement procedures, dermal template placement and skin grafting." (Pretz, 1) Prior to his injury the candidate led a normal life, he was independent and able to work.

After his injury the patient had a strong desire to ambulate independently and wanted to decrease his family's burden of having to care for him. Prior to therapy he was very dependent and required assistance getting food for himself, dressing, toileting, and other aspects of self-care due to the severity of his wounds. His level of motivation would serve him well throughout his rehabilitation process.

Initially, the patient was only able to walk about 5 feet with the support of two therapists using maximum assistance. His pain scale while doing this was a 10-10. His physicians decided to fuse his ankle once the wounds healed and made him non-weight bearing for one month. "The patients functional status slowly improved throughout his hospitalization. Four months after the injury he was able to ambulate 30 feet with a right platform shoe and a left PTB orthosis at supervision level without an assistive device. The patient demonstrated fair dynamic and static standing balance. For sit-to-stand, the patient was at a minimal assistance to supervision level. He remained at maximal assistance with upper and lower extremity dressing, including donning and doffing the PTB











orthosis and right platform shoe." (Pretz, 2)

Bo Powers, CPO fabricated the PTB orthosis with a custom molded socket, double adjustable joints, steel shank in the shoe, a T-strap, and a proximal pull strap to pull the molded anterior shell into the patella tendon area and maximize load bearing. The patient's skin integrity had to be monitored closely, and he was instructed to wear the brace at all times when

weight bearing or at least 8 hours a day. In reality, most of his day was spent in a seated position, but having the brace on gave him the independence to move when necessary, it also reduced his pain level to a 3-10.

"The case described here is an exemplar of how the PTB orthosis could be of great and common use potentially for many thousands of patients with injuries due to natural disasters, war or other traumas. In the acute care phase of hospitalization, specialized teams are present for a short period of time and an ankle or foot can be operated on or set and casted, then the patient can walk independently, with the PTB orthosis allowing the foot and ankle to remain non-weight-bearing, and the orthosis being discarded when no longer needed" (Pretz, 3).

As mentioned earlier, the PTB is an underutilized orthotic device. Perhaps because it is an extensive brace and can initially be seen as costly. However, upon closer analysis, the benefit of maintaining mobility can reduce medical expenses in the long run. And the emotional benefit of maintaining a level of independence through some level of mobility and being able to progress in therapy is an invaluable benefit to any patient. 🦋

Rachelle Pretz, PT, DPT, Cora Brown, MD, William B. Hughes, MD, and Eric L. Altschuler, MD, PhD "Maximizing Functional Mobility In An Electrical Burn Patient Using A Patellar Tendon Bearing Orthosis," Journal of Rehabilitation Medicine Vol 48, 2016 pp 1-3.

Patellar Tendon Brace HIGHLIGHTS

The Patellar Tendon Brace (PTB) can be designed with either a custom foot section and a walking base or attached to a patient's shoe, depending on the patient's needs and anatomical presentation. Below are some photos highlighting the unique designs of the PTB:

- 1. Patella Tendon Bar with popliteal push. By pushing into the patella tendon and popliteal area the pressure is transferred and the foot and ankle are offloaded.
- 2. The back wall is dipped to allow room for the medial hamstring and increase comfort while sitting.
- 3. Custom shaped anterior shell increases the weight-bearing surface and locks the patient into the orthosis and prevents migration within the device.
- 4. Proximal strap helps insure the maximum squeeze between the patellar tendon and popliteal is achieved.
- 5. The walking base with a rocker bottom allows a patient with a custom foot section to ambulate on any surface. But more importantly, the rocker bottom helps ensure the patient quickly rolls over and minimizes the timespent weight bearing. But more importantly, the rocker bottom helps ensure the patient quickly rolls over and minimizes time spent weight bearing on the leg.
- 6. When the PTB is attached to a shoe, a double adjustable joint is used with metal uprights and steel shank is placed inside of the shoe.



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