Heroic Medicine

Nova Cells Institute's Extraordinary Healing Program

We are revolutionizing medicine one turnaround at a time

Nova Cells Institute

562-916-3410

www.novacellsinstitute.com
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US Information Line: 1-562-916-3410
The Nova Cells Story

Nova Cells Institute of Mexico (NCIM) was founded by researchers and administrators fed up with the underhanded tactics and unbridled greed rampant among many private clinical stem cell programs abroad. This is one reason you will read this statement throughout NCIM’s literature and postings: “Patients & their caregivers deal directly with the scientists, technicians, doctors & other professionals who are part of the NCIM program. There are no brokers, marketers or other middlemen involved in NCIM's program.”

Not surprisingly, NCIM is a small organization run by those who do the day-to-day “grunt work.” It is in-a-sense both a collective and a family. Patients and others who have been through the program quickly pick up on and love the fact they do not have to wade through marketing pitches from professional salespeople, slick brokers or three piece suits (executives.) In addition, they do not have to deal with people whose desire to help others is largely predicated on making money. Those who handle and care for them at NCIM are without exception people who have a passion for going the extra mile and a long history of doing this, often for little or no pay or other tangible rewards.

And most importantly, NCIM is results-committed and driven. Its doctors and researchers are tireless in their quest of things that reliably turn-the-tables on human suffering. In addition, the body of science that informs NCIM’s work and offerings is very leading edge. Biochemist Dr. Abel Pena, who runs the laboratory end-of-things for Nova Cells, was trained by a leading US stem cell biologist and maintains the highest levels of safety and hygiene in his lab in Mexico. In addition, he has pioneered numerous methods for priming stem cells, i.e., getting them to respond to chemical signals in target tissues by becoming cells that tend to effect or support healing or relief. Stem cells primed using Dr. Pena’s methods have produced salutary, sometimes remarkable clinical responses in people with a wide variety of neurologic & other diseases and conditions. He also created NCIM’s proprietary "Beacon Factor"

Click to read more about this

Read more: The Nova Cells Difference

Phone: 1-562-916-3410  Email: NCInfodesk@gmail.com
Videos: Get acquainted with Nova Cells

https://www.youtube.com/watch?v=-BH7WRv5l0o

https://youtu.be/GAS7IXWLe0s?list=PLySdDdYLElg520yVeEwp
u7BmwZ1yu2QyO
“Stem cells have the remarkable potential to develop into many different cell types in the body.

“When a stem cell divides, each new cell has the potential to either remain a stem cell or become another type of cell with a more specialized function, such as a muscle cell, a red blood cell, or a brain cell.”

From the National Institutes of Health (NIH) Stem Cell Information website: http://stemcells.nih.gov/index.asp

PRINCIPLE SOURCES OF STEM CELLS

There are a number of sources of stem cells:

1) Embryonic stem cell (Nova Cells does not use these, never has & never will)

2) Stem cells taken from aborted human fetuses [Fetal cells] (Nova Cells does not use these, never has and never will)

3) Umbilical cord blood.

4) Umbilical cord Wharton’s Jelly (Stem call-laden gelatinous layer that surrounds umbilical cord blood vessels).

5) Placental blood and tissues.

6) Bone marrow & other tissues such as bone marrow, skin, fat (adipose), liver and so forth.
Stem Cell Potency

**Totipotent** – All somatic cell types, Germ Cells, Placenta
- Fertilized egg, Zygote, Morula cells
- Adult Totipotent Stem Cells

**Pluripotent** – All somatic cell types
- Inner Cell Mass (ICM) of Blastocysts
- Embryonic Stem Cells
- Adult Pluripotent Stem cells

**Lineage-committed** – Germ Layer Lineage Stem Cells
- Ectoderm associated stem cells
- Mesoderm associated stem cells
- Endoderm associated stem cells

**Multipotent** – Cell specific
- Neural stem cells
- Cord blood stem cells
- Hematopoietic stem cells
- Most bone marrow stem cells

**Unipotent** – Committed to becoming a single cell type
- Dopamine expressing neurons
- Muscle cells
- Cartilage
- Insulin-producing cells

SEE ALSO: [https://ncimx.wordpress.com/stem-cells-we-use/](https://ncimx.wordpress.com/stem-cells-we-use/)
Stem Cells Used in the Nova Cells Program

The Nova Cells program only uses adult (nonembryonic) stem cells derived from the human umbilical cord including blood and Wharton’s Jelly, the placenta, as well as those harvested from a patient’s own tissues such as bone marrow.

See also: [https://ncimx.wordpress.com/stem-cells-we-use/](https://ncimx.wordpress.com/stem-cells-we-use/)

The umbilical cord blood, **Wharton’s jelly** and placenta-derived stem cells are extracted from donors with Type O blood. The stem cells used include but are not limited to CD34+/AC133, neurogenic progenitors, mesenchymals (MSC), Primitive Neurogenic Progenitors, i.e., embryonic-Like stem cells from the cord -- Embryonic-like stem cells from cord - pluripotent neurogenic progenitors.

All medical procedures and laboratory processing is done in Mexico.

**DONOR CELLS ARE HEAVILY SCREENED FOR DISEASE, CONTAMINATION, ETC.**

All human umbilical cord and placenta-derived stem cells and cord blood serum used by Nova Cells contracted/affiliated/consulting physicians are processed in state-of-the-art Mexican hematology laboratories that conform to the highest Quality Control/Quality Assurance standards and employs state-of-the-art technology & lab methods. The blood and various tissues are acquired from healthy mothers who give birth to full term, normal, healthy babies (in Mexico). Harvested blood and tissues are then screened for all major communicable diseases using highly sophisticated lab methods, after which stem cells are separated out. These cells are then typically (but not always) expanded, i.e., induced to reproduce, in a growth medium that is free from any animal products, i.e., no mouse feeder cells, fetal calf serum or other animal serums, etc. The cells used are almost always “fresh;” that is, no freezing and storage is involved. The cells given to patients come from either patient matched blood type or Type O (Universal type) blood are gender-matched, i.e., cells from males are infused into male patients and cells from females are used to treat female patients.

**NOTE:** Many people ask about having Nova Cells use cord blood they have stored in the US or Canada. In the US, cord blood is treated by the FDA as a drug. As such, it cannot be used for any condition not approved by the FDA stateside nor can it be sent to Mexico for such purposes.
IMPORTANT POINTS

Nova Cells Institute has never worked with embryonic or fetal stem cells, and never will. Embryonic and fetal stem cells can produce tumors (These are classified as pluripotent" which means they can form teratomas and other tumors in certain instances such as being injected into the central nervous system). The adult stem cells Nova Cells uses (umbilical cord, Wharton's Jelly, placenta-derived, and a patient's own bone marrow) are "multipotent" which limits the body cell types they can become (These stem cells do not form teratomas or such. Some private stem cell clinics in Europe and the US have injected adult stem cells into organs such as kidneys and eyes which caused problems and complications. Nova Cells Institute has never injected stem cells into an organ).

Let's be very clear about these points and a few others:

(1) Nova Cells was established in 2009. There are other companies that came along later on which use "Nova" in their company title, but these are NOT affiliated in any way with Nova Cells Institute of Mexico (NCIM).

(2) Nova Cells Institute has never used embryonic or fetal stem cells and never will (They can form tumors under certain circumstances).

(3) Nova Cells Institute only uses multipotent adult stem cells derived from umbilical cord blood, Wharton's Jelly, placental tissue, and a patient's own bone marrow.

(4) Nova Cells Institute administers stem cells by intravenous drip (On rare occasions Nova Cells affiliated doctors recommend an intrathecal or spinal tap infusion of cells. Everything related to this is spelled out, a process called "informed consent". Patients who prefer not to do this are given cells by intravenous or IV drip).
More about **Wharton’s Jelly Stem Cells**

NOVA CELLS INSTITUTE HARVESTS & MAKES CLINICAL USE OF STEM CELLS ISOLATED FROM (UMBILICAL CORD) WHARTON’S JELLY. BELOW ARE LINKS TO A SAMPLING OF PAPERS PUBLISHED ON WJ STEM CELLS:

**Wharton’s Jelly-Derived Mesenchymal Stem Cells: Phenotypic Characterization and Optimizing Their Therapeutic Potential for Clinical Applications**

**BONE REGENERATION**

Bone Regeneration Using Wharton’s Jelly Mesenchymal Stem Cells

**CANCER**

Human Wharton’s jelly mesenchymal stem cell secretome display antiproliferative effect on leukemia cell line and produce additive cytotoxic effect in combination with doxorubicin

Umbilical cord tissue-derived mesenchymal stem cells induce apoptosis in PC-3 prostate cancer cells through activation of JNK and downregulation of PI3K/AKT signaling

**CYSTIC FIBROSIS**

The Potential of Wharton’s Jelly Derived Mesenchymal Stem Cells in Treating Patients with Cystic Fibrosis

**DIABETES**

Wharton’s Jelly Mesenchymal Stem Cells for the Treatment of Type 1 Diabetes
MULTIPLE SCLEROSIS

Therapeutic Effect of Transplanted Human Wharton’s Jelly Stem Cell-Derived Oligodendrocyte Progenitor Cells (hWJ-MSC-derived OPCs) in an Animal Model of Multiple Sclerosis

NEURONAL REPAIR & NEW NERVE CELL FORMATION

The Secretome of Bone Marrow and Wharton Jelly Derived Mesenchymal Stem Cells Induces Differentiation and Neurite Outgrowth in SH-SY5Y Cells

Matrix Cells from Wharton’s Jelly Form Neurons and Glia

Neurotrophic factors from human Wharton’s jelly mesenchymal stem cells induce gliogenesis

Wharton’s jelly derived mesenchymal stromal cells: Biological properties, induction of neuronal phenotype and current applications in neurodegeneration research

Conversion of Human Umbilical Cord Mesenchymal Stem Cells in Wharton’s Jelly to Dopaminergic Neurons In Vitro: Potential Therapeutic Application for Parkinsonism

TISSUE REGENERATION

Wharton’s jelly mesenchymal stem cells promote wound healing and tissue regeneration

TRAUMATIC BRAIN INJURY

Wharton’s Jelly Transplantation Improves Neurologic Function in a Rat Model of Traumatic Brain Injury
Unique to Nova Cells:
The Beacon Factor™

About the Beacon Factor (invented by NCIM’s own Dr. Abel Pena)

https://biotheorist.files.wordpress.com/2015/02/about-the-beacon-factor.pdf
Conditions Successfully Treated

Learn more about each of the disorders below by either following the link, or emailing NCInfodesk@gmail.com or by calling 1-562-916-3410. Nova Cells has successfully treated many other conditions so if you do not see your particular challenge here, just ask us about our experience with it!

https://ncimx.wordpress.com/conditions-nova-cells-has-successfully-treated/

FREE, NO OBLIGATION CASE EVALUATION - HOW TO GET YOURS: https://ncimx.wordpress.com/free-case-evaluation-how-to-get-yours/

Autism – Email NCInfodesk@gmail.com for specifics or call 1-562-916-3410

Cancer (End-stage & advanced too): http://www.novacellsinstitute.com/cancer-stem-cell-therapy.html

Cardiovascular: https://ncimx.wordpress.com/category/reversing-blockage-in-blood-vessels/

Cerebral Palsy – https://ncimx.wordpress.com/?s=Cerebral+Palsy

Chronic Fatigue Syndrome (CFS) – Email NCInfodesk@gmail.com for specifics or call 1-562-916-3410

Dementia/Lewy Body Dementia – https://ncimx.wordpress.com/?s=Dementia

Epilepsy/Other Seizure Disorders – https://ncimx.wordpress.com/?s=Seizure

Fibrosis – Email NCInfodesk@gmail.com for specifics or call 1-562-916-3410

Multiple sclerosis – https://ncimx.wordpress.com/?s=multiple+sclerosis
Parkinson’s Disease – See alzheimer’s-dementia

Spina Bifida – https://ncimx.wordpress.com/?s=Spina+Bifida

Spinal Cord Injury/Paralysis –
https://ncimx.wordpress.com/?s=spinal+cord+injury

Stroke – https://ncimx.wordpress.com/category/stroke/

Traumatic Brain Injury – Email NCInfodesk@gmail.com for specifics or call 1-562-916-3410
Patient Responses to Treatment
Ryan Weber (Anoxic Brain Injury): Improvements began surfacing right after his treatment!

(August 2014) To Grace & All The Nova Cells Family:

Sorry for not getting back to you sooner but it has been crazy busy. Mexico was great, nothing we imagined. The hospital where Ryan received his Stem Cell Treatment was so clean and I have been in many different hospitals here in the United States that was not as clean. We enjoyed the Mexican food as it was so flavorful. It made me sad to leave.

Ryan’s Stem Cell treatment was a great success! We noticed a huge difference in his muscle tone when they brought him back from the O.R. I cried when I first picked Ryan up because it was like holding a different kid. Ryan’s muscle tone is completely relaxed. Ryan has started back up with his outpatient physical therapy, warm water therapy and his massage appointments. His therapist were shocked at the difference in his muscle tone. Ryan has more control over his muscles.

Some of the things Ryan has done since his Stem Cell Treatment that he was not able to do before hand include:

HEAD CONTROL - He holds his head in a neutral position now and freely moves it to the left and right to look around. Before Stem Cells, Ryan would either have it tilted to one side or the other or front or back.
Picture taken the day after Ryan’s treatment at the San Diego Zoo. Look at how his arms are more bend, his legs are bend, head in right position. Ryan is actually sitting on the tusk.

HANDS – Ryan is able to open and close his hands freely. He regained the infant grasp reflex. With this he is able to reach out and grab items. Last night (8/26/14), he reached up and grabbed his Dad’s glasses and the later in the night he reached up and grabbed a handful of my hair. Before Stem Cells, Ryan kept his hands closed in a tight fist.

Picture of Ryan’s hands after his Stem Cell treatment
**ARMS** – Ryan actually is able to control both his arms equally. He is able to bend both elbows at his choice. Ryan is able to control his arms to reach to get items he wants. He still favors the left arm over the right but he was left handed dominant. Before Stem Cells, Ryan did have minimal control over the left arm and no control over the right arm.

![Photo of Rick and Ryan at the beach before Ryan’s Stem cell treatment.](image)

*Rick and Ryan at the beach before Ryan’s Stem cell treatment. Look at the stiff straight arms and tight closed fist hands*

**TRUNK (TORSO)** – Ryan is able to sit in upright position on our lap with little help. When Ryan sits his chest and belly are straighter. Before Stem Cells, Ryan sat on our lap and needed a lot of help to stay in upright position and usually was tilted to one side or the other.

**LEGS** – Ryan legs are able to bend at the knees. He is able to control them better alternating them to walk in his walker and today walked in the pool with very little help by himself without a walker. When Ryan sits on our laps Ryan will bend his knees and sit like any other “scientifically Normal” child. When he lays on the floor he will pick up his legs to reposition them. Before Stem Cells, Ryan legs were mainly in a locked straight position. It was a battle to have him sit on our lap, put him in his car seat or in anything
that required him to be in a bend position. He would walk in his walker but his body looked like an “S”.

**MENTAL** – Ryan is happier; he smiles and laughs more. I credit the Stem Cells as he is not always having to deal with very tight muscle which were probably really sore. When Ryan looks at you, he just smiles like he can actually see us clearly. Before he recognized the important people like his siblings, Grandparents and us as his parents but since his Stem Cell treatment-couple of our friends have talked to Ryan and this time Ryan interacted with them.

Most of Ryan’s improvements have been “small” but when you put all the small pieces together it makes for a huge improvement. It’s the little things that matter. I have to tell you a story about our day before we were picked up to go to Mexico.

We were all around and waiting for the phone call to tell us you were on your way. We were sitting outside at the patio set when I opened my pictures in my Phone to find the Creative Miracle Prayer I wanted to say before Ryan’s Stem Cell treatment. My phone usually opens to the last picture taken but that day when I opened my pictures, I started to cry because it automatically opened to a picture of Rick holding Ryan the day he was born back on 12/17/09 (See below). Rick asked what was wrong and I turned my phone to show him the picture that popped up and his jaw dropped. This is the picture that popped up.

![Image of Rick holding Ryan]
I took that as a sign from above that Ryan would be born again. With all of Ryan’s improvements that is exactly what happened! We cannot thank you all enough for making this miracle happen for Ryan. I will send pictures in another email with little explanations of them. Just wanted to get this email of Ryan’s improvements to you. You may share this with everyone and if they want to follow Ryan his Facebook page is www.facebook.com/littlewarriorryan.

Thank you for giving us our little boy back! Lots of love.

Gina & Rick Weber
Bladder Problems

**Julia Simpson’s eye, bladder and Parkinson’s symptoms disappeared quickly following Nova Cells treatment**

Emailed to Nova Cells:

This is Julia Simpson. I heard about Nova [Cells Institute](#) from other patients so I looked them up and checked out their website and testimonials. I am getting on in years being 66 and just feel the weight of the years on me. I have been pretty much healthy all of my life but as the years passed by I acquired more and more little problems which latter turn chronic. I get check-ups, take supplements, but my symptoms persisted.

I have a prolapsed bladder which makes me get up at night at least 2 times sometimes three times. This has been going on for about 6 years.

About 8 years ago a doctor prescribed [Calcitonin](#) as a nasal spray to avoid bone loss. I used it for about a month and a half. About a month into it I started to have problems with my right eye. It began tearing during the night and I had pain in it and it was red and irritated. Then headache started on that side. I did not connect this to the nasal spray. For days I had the tearing and redness even during the day and became very sensitive to sunlight. I realized that I was putting the spray in my right nostril, and the right eye was the one giving me a problem. So I then stopped using it and told my doctor. The problem is that although it stopped bothering me during the day it has never stopped giving me problems at night if I sleep on the right side, and I still had lots of pain.

I have more or less a lot of energy but I take one or two small naps a day to get reenergized.

One major thing that has been bothering me for the last 6 years is that my right hand is trembling slightly, some times more so when I get stressed. I got very worried about having Parkinson’s. I decided to get a stem cell treatment for rejuvenating my organs and hopefully helping get rid myself of the Parkinson’s like symptoms because although it is not progressing it does not go away and left me very worried about this.

On June 19, 2015 I had my stem cell treatment at NCIM . I was picked up by Abel and Grace who were so nice and professional. They answered all of my questions so I felt at ease and relaxed because I went to Mexico alone.
The city of Tijuana was just around the corner sort of speak. Just 15 minutes from the city of San Diego. The city itself was totally different from what I had imagined. It is a real nice city of 3 million people. They have roundabouts often and lots of statues of heroes, even one of Abraham Lincoln (planted in the avenue of the heroes). I understand the Lincoln statue serves as a “thumbs up” to American democracy and is a “welcome here” for US tourists. Lots of beautiful tall buildings, many made of all glass, dot the landscape everywhere. I saw McDonald’s, Costco, Office Max, Office Depot, Walmart, and a Subway Sandwich shop among others. People were milling about and walking all over the place, kind of like New York. I felt really good about Tijuana and totally safe.

Once I got to the hospital I found the staff incredibly friendly and courteous. Once I got checked in I was assessed by three doctors an internal medicine as well as a neurologist and an anesthesiologist.

The doctors did many tests and then I was infused with the Beacon Factor and got my stem cells by IV and also a percentage by spinal tap. I was not put to sleep for my spinal tap. The MDs just gave me a local anesthesia where the little puncture was made. It didn’t hurt at all. Then I got more Beacon Factor which helps the stem cells seek out damaged tissue. I was there for observation for a few hours then discharged. Abel and Grace then took me to eat at a delicious cafeteria with all kinds of Mexican dishes to choose from. Then was brought back across the border in the US.

About 6 days after my treatment all of a sudden I noticed that my eye no longer was tearing up or hurt any more. I could not believe it and decided to wait a few more days just in case it came back. It has already been 17 days and the eye is not tearing up.

My energy has shot up incredibly too. To the point where I’ve only taken naps three times in all these days, and short ones at that.

About a week into my treatment I noticed that I was not waking up at night to go to the bathroom. I was instead sleeping through the night like a baby and waking up at 8:00 or 8:30 AM each morning. I wake up so rested. I am going into deep sleep every night and am even dreaming now which was not true in the past.

My bladder, of course, is still prolapsed, but I do not have the sensation of needing to visit the bathroom over and over again each night.

But the most wonderful thing is that my hand shaking is all gone and I am ecstatic about this! About 10 days post-treatment, I was using my computer when I realized that I was not shaking any more. Of all the symptoms impacted by my treatment I am so happy and amazed that the shaking is gone! It was amazing that I felt so good I did not even notice the change at first. Thank you Nova Cells Institute! Yes, I was hoping to get rid of
my Parkinson’s symptoms, but I never thought that they would be gone in a matter of days. Maybe it was so quick because my shaking was rather mild, but even if it had taken months I would have been just as happy. I am not afraid or concerned any more.

Julia Simpson
Cancer

Killing cancer, not the cancer sufferer (Targeting & defeating cancer while leaving healthy cells unaffected)

On Sunday December 7 (2014) CBS’s famed 60 Minutes devoted a thirteen minute segment titled “Disrupting Cancer” to physician and billionaire businessman Patrick Soon-Shiong, MD’s advocacy and use of the rapid gene sequencing of tumor cells to help zero in specific drugs or other agents that will eradicate them. Later the same day, Forbes’s writer Matthew Herper published an article in which he stated something virtually everyone doing cancer work knows, namely that “Everyone is looking to use DNA sequencing to better pick cancer drugs. And in some ways, Soon-Shiong is an odd person to pick as a spokesperson for this, because he’s just getting started.” (Here Is What ’60 Minutes’ Didn’t Tell You About The Billionaire Who Is Trying To Disrupt Cancer Care).

This is not to say that with Dr. Soon-Shiong’s deep pockets and army of computer experts and researchers he will not wind up making major inroads in this area of biomedicine. And with cancer striking so many, progress is welcome whether it comes out of a one person lab or a research enterprise that fills buildings or is stretched across many continents.

Nova Cells is no stranger to finding ways to “disrupt cancer”. It has, in fact, focused on a treatment approach that combines attacking tumors using donor immune cells (from healthy young people) and specific compounds that gum up the inner machinery in cancer cells while leaving healthy cells. It has been a spectacularly successful quest that started with the treatment of a 71 year old gentleman from the Pacific Northwest who was in the last throes of end-stage prostate cancer that had spread into his liver and bones. He was, in fact, so bad off that NCIM doctors thought he might not live more than a handful of days, at best. But, like so many people struggling with cancer, he was a spirited fighter who simply would not entertain abandoning hope until, as he put it, “I take my last breathe”.

Page 24 of 79
This determined chap knew the score going into treatment, namely that he was volunteering to undergo something very experimental in nature. But he was resolutely committed to “fighting the good fight”, even if this pitched battle proved his last. Seeing his determined spirit, the MDs set about stabilizing him to help insure he could handle the biological blitzkrieg (lightening war) that NCIM’s experimental tumor-killing method was expected to unleash in his body.

Three days later the doctors began daily treatments using cancer-fighting cells (called granulocytes) that had been previously isolated from the blood of many young, healthy volunteers and then combined and infused by intravenous (IV) means. In addition, he was to be given intentionally HLA mismatched umbilical cord stem cells.

If the treatment worked as anticipated, the donor granulocytes would attack the tumors and also send a “wake up call” to the immune system, i.e., alert the fellow’s own immune cells that something alien was present which they would then attack (One problem in cancer is that tumors send out biochemical signals that suppress the sufferer’s own innate tumor-fighting cells). In addition, the HLA mismatched umbilical cord blood stem cells would be attracted by growth factors secreted by the tumors, attach to them, and then be attacked by the patient’s own immune cells which would react to them as a foreign presence.

The outcome? He did have a mild immune reaction (rash) that was easily suppressed using an IV antihistamine. This is how NCIM’s own patient care coordinator, Grace Pena, Ph.D. cand., summed up what transpired:

“Within three days his face was pinkish and his hands were getting pink. He was eating so well that the staff could not believe it. After the 7th primed donor granulocyte treatment he was walking 1/2 mile around the hospital, and feeling great. He was feeling so good in fact that he decided to return to his home state of Washington to finish up a million dollar real estate deal. NCIM doctors and staff protested wanting him to finish the entire course of therapy but he was adamant about getting home, but promised to return and complete his care once his real estate deal was completed. This he did. By October his US oncologist reported that his PSA is 1.1, his prostate is smaller than a normal 20 year olds, and he could find no cancer at all. Also, his Alkaline Phosphatase level which was about 2,000 during his treatment had dropped to something around 89.”

This fellow remains in full remission to this day (September 2015).
Following this exciting clinical success, other end-stage cancer patients were treated (prostate, breast, lung, others) using the same method. Fifty percent (50%) experienced tumor shrinkage (partial remission) while the rest (50%) went into full remission.

In addition, other cancer-fighting measures were tested both alone and in combination with the donor granulocytes & cord blood stem cells (Gr-UCSC). Those that resulted in tumor shrinkage or obliteration when used alone, as well as those that bolstered the anti-cancer impact of the Gr-UCSCs were retained.

Among these were two chemical compounds that throw a monkey wrench into the tiny power-generating factories (mitochondria) in tumor cells but do not adversely affect normal cells. They are, in technical parlance Patrick Soon-Shiong, MD, selective for malignant cells.

NCIM also selectively utilizes a unique injectable protein that amplifies the cancer-fighting activity of a patient’s own macrophages.

Returning now to that 60 Minutes segment: Dr. Soon-Shiong beamed about tackling cancer at an immunological level. Nova Cells doctors and staff wish him well in this quest, but can’t help adding that in many ways we left the starting gate years ago and are “already there” (That is, have made demonstrable progress and enjoyed great success).

If you’d like to know more about NCIM’s cancer treatment method, just click these links:

Click to read more about NCIM’s Gr-UCSC method of treating cancer

Click to access cancer patient write-ups
Linda’s little boy has cerebral palsy and has vision, cognitive and walking issues. Within a week of getting primed stem cells & the nontoxic Beacon Factor by Nova Cells Institute doctors (August 2015), Linda shared this with Nova Cells by email (gracepatients@gmail.com):

As you know, my boy has really been struggling for a long time with walking and now after his treatment he just gets up and does it! Today he was even trying to run about. Also, I see signs his vision has improved too. On top of all this, his temperament is so much better and he is no longer easily frustrated and his ability to focus & take on challenging tasks has improved. For instance, today at therapy he put objects like circles, triangles and stars into the right sorting container which left the staff totally blown away! My husband took in all these leaps forward which left him almost in a state of disbelief! I wish all the parents of children with CP would get them into treatment with your
program. Warmest regards to both you and Dr. Abel, and do watch your email for more progress updates and video clips too of our boy. Love, Linda.

Grace added:

At the hospital, even before we left to take this family back to their hotel in San Diego, the boy was sitting on the floor, then just folded one of his legs and got right up without help from his mom or anyone present. This so amazed the lady that it left her totally wide mouth open. Then at the hotel the boy did it again but this time took about 4 steps and sat down. The mother was simply awed by what was happening.

Emily Johnson: Ten year old with cerebral palsy turned a corner following her NCIM stem cell treatment

Ten year old Emily Johnson was born 16 weeks premature (weighing in at only 1 lb 6 oz.) and sustained nervous system damage later was manifest as an inability to communicate verbally, problems getting about, and seizures. Her doctors diagnosed her as having cerebral palsy. Like many children with CP Emily benefited from use of a walker and various kinds of therapy. In time her seizures abated and she learned to communicate via a speech device. Her parents, Patrick and Justin, wanted more for Emily, of course. Like most other parents of children with CP they went on a quest for therapies (including novel ones) that might reduce or remediate Emily’s disability and improve or enhance her motor and cognitive functioning. This quest brought them to Nova Cells Institute of Mexico (NCIM) in December 2011 and culminated in Emily’s being treated with pure umbilical cord and Wharton’s Jelly derived stem cells plus “Beacon Factor”, i.e., a proprietary hypoallergenic combination of peptides and cytokines that helps stem cells rapidly home in on and anchor to target tissues (NCIM’s Beacon Factor has been used by NCIM on all types of neurological diseases such as cerebral palsy, multiple sclerosis, spina bifida, and Parkinson’s disease with great success).

Emily’s response to her NCIM treatment began 2 months following her treatment and is ably captured in this update from her parents that came in this month (May 2012):

Our daughter Emily had a stem cell treatment through NCIM when she was 10 years old. We set up everything through (NCIM patient educator and care facilitator) Grace
and she was wonderful to us. She was so attentive and loving and I really felt a bond with her over the phone and felt like I knew her and could really feel her excitement for helping kids!

When we got to San Diego, Grace and (her biochemist husband) Abel picked us up and personally escorted us to Mexico! Traveling to another country can be a bit overwhelming but they made us feel at ease and we really felt like they cared about our little Emily and they were very thorough in explaining how they get the stem cells and treat them and the desired outcome.

Since the treatment we have seen some wonderful changes in our daughter. Emily is non-verbal and cannot walk without assistance. Prior to her stem cell treatment she could not ask for help or communicate her needs effectively even with a speech output device. Now, just 5 months after treatment, she clearly articulates her needs using the device and she has also learned how to use an iPad for communication as well.

She has also shown a new level of maturity. She has demonstrated through testing that she can read well over 200 words. She is also able to build sentences and follow along with academic activities using her speech device. Her temperament has also changed. She used to break down easily and get very frustrated and she was unable to control her emotions. Now she seems to be more patient and confident in herself.

We feel that these changes are just the beginning and we are hopeful and excited to see even more good things coming from her treatment!

TO ACCESS A NEWS RELEASE THAT INCLUDES MENTION OF EMILY CLICK THIS LINK OR GO TO http://bit.ly/J9AKQw
Dementias
(Alzheimer's, Lewy Body)

Alzheimer’s: How Nova Cells is helping people turn the tables on this disease

https://biotheorist.files.wordpress.com/2015/02/ncim-dementia1.pdf
Messed up cellular “garbage disposal system” found to play a role in Alzheimer’s disease (And Nova Cell’s BEACON FACTOR coaxes lysosomes to do their job!)

Many neurodegenerative diseases involve a failure of lysosomes (the “garbage disposal” system within cells including neurons) to do their job properly. Now comes evidence that defective or compromised lysosomes may contribute to Alzheimer’s disease, something ably laid out in a 6-30-15 article on the Medical Express website at http://medicalxpress.com/news/2015-06-failure-cells-garbage-disposal-contribute.html. Here is a salient quote from this article:

"Lysosomes, the “garbage disposal” systems of cells, are found in great abundance near the amyloid plaques in the brain that are a hallmark of Alzheimer’s disease. Scientists have long assumed that their presence was helpful—that they were degrading the toxic proteins that trigger amyloid plaque formation.

However, in Alzheimer’s patients, these lysosomes lack the ability to do their jobs properly, and instead of helping, the accumulation of lysosomes may even contribute to the disease, Yale University researchers report the week of June 29 in the Proceedings of the National Academy of Sciences.

The new findings raise the possibility that coaxing lysosomes to do their jobs could help to prevent the toxic processes that eventually destroy the minds of Alzheimer’s patients.

The big question is how to get lysosomes to do their job as they should. While medicine has little that can pull this off, Nova Cells possesses an effective, nontoxic way to do so: Namely its proprietary Beacon Factor. Not surprisingly, many patients with neurologic diseases and conditions that involve fouled up lysosomes have responded quickly and beautifully to intravenous infusions of the Beacon Factor. Click to read some of the case histories."
Canadian Lady with Progressive Supranuclear Palsy, Corticobasal degeneration (CBD) and prefrontal dementia has an amazing response to her NCIM treatment

Progressive Supranuclear Palsy (PSP) is a devastating neurodegenerative disease that oftentimes robs its sufferers of their memories, balance and much more. As bad as this insidious disease is, things get even worse when the patient has other neurologic challenges such as Corticobasal degeneration (CBD) (which causes cell die off in various parts of the brain including in many instances those critical to thinking and executive decision).

Recently Nova Cells was approached by a Canadian gentleman whose wife has PSP, CBD and prefrontal dementia. NCIM doctors reviewed scans of this lady’s brain which revealed clear evidence that her frontal lobes had been impacted by her disease. Her
husband was advised that if his wife was treated with primed stem cells and the Beacon Factor, she would probably see very mild improvements that would most likely not be manifest until about 2 months after her treatment.

As this was a “heroic measures” case and the husband did not have an exaggerated hopes concerning outcome (“If she could just relax her right arm and hand, which has caused her so much pain, and be understood a little more, this would be a Godsend”), the patient was booked for treatment on Thursday, April 30.

NCIM patient educator and care coordinator, Grace Odgers, Ph.D. candidate was present during the lady’s treatment and had this to say about her condition and response coming out of the gate:

The patient is really in a bad condition. But as soon as she came out of the surgery she was acknowledging things and people around her. She was locking eyes with her husband and looking around which is something she had not done in a quite a while.

And, while her left arm and hand worked normally, her right arm was totally stiff and bent at the elbow and drawn up against her chest. Her right hand formed a stiff inflexible fist which she could not open or relax at all.

About an hour after her treatment the attending physician came in to check on her and noticed that she was more alert and also that her right arm and hand were now relaxed! He took her hand and opened it more and more until it was fully extended. Then her husband asked her to shake hands with him and she did which amazed everyone! We were all absolutely flabbergasted and emotional.

Soon the lady was smiling a lot, especially compared to when she first arrived. Prior to her treatment her face was somewhat stiff and expressionless and though she tried to smile, all she could manage was a forced half smile.

The doctors mentioned that when she went into the surgery room and they commenced treating her, she extended her totally stiff right leg about 2 minutes into her infusion.

The couple returned to Canada on Monday the 4th. Grace called the lady’s husband shortly after they got home and was told by him that his wife’s balance had improved significantly (He added that this is “great” because she has a history of falling).
LEWY BODY DEMENTIA: How Arthur Rechlo regained ground lost to Alzheimer’s cousin

Lewy Body Dementia is sometimes called “Alzheimer’s cousin”. Like Alzheimer’s it is an incurable brain disease that steadily whittles away most sufferers’ cognitive functions including their ability to focus, sleep well, or walk normally. Those with advanced LBD may have trouble swallowing liquids and solids.

Currently LBD affects about 1.4 million Americans. Both radio and TV personality Casey Kasem and Oscar® winning actor Robin Williams had LBD, and legendary Chicago Blackhawks coach Stan Matika was recently diagnosed with it.

LBD has no cure and the best medicine can presently do is relieve symptoms and alleviate patient suffering. The one thing doctors never see is a person with advanced LBD who can hardly walk or talk begin doing so. This all changed after 76 year old Canadian Arthur Rechlo had an experimental treatment in Mexico.

Click to read all about Arthur’s amazing turnaround.

http://www.stalbertgazette.com/article/20150228/SAG9604/302289988/-1/sag
Epilepsy turnaround for an angel (Angel Sosa)

Angel Sosa is 4 years old and has not exactly had an easy ride in life. Born to loving parents and a large family he entered this world saddled with lots of neurological and physical challenges. Among them: Infantile spasms, a difficult-to-treat form of childhood-onset epilepsy called Lennox–Gastaut syndrome (LGS) or simply Lennox syndrome, and developmental delays. According to his loving Mum, Monique, he was “having 10 clusters of 10-11 seizures lasting six (6) minutes each day and was having ten tonic seizures a day sometimes lasting up to 3 minutes without breathing."

As you might expect little Angel’s early life was spent seeing doctors and therapists and taking drugs to control his epilepsy (Three currently: Felbamate, Valproic acid, and Zonisemide). Naturally, his parents kept an eye on the news and the Web for any medical advances that might offer any hope of remediating their son’s intractable epilepsy and other challenges. With the advent and growth of adult (nonembryonic) stem cell treatment programs outside the US they saw evidence that perhaps some form of stem cell therapy could help their little boy. As they sorted through the various private stem cell medicine operations abroad they came across accounts of the many turnarounds in children effected by Nova Cell’s primed stem cell treatments and intravenous use of its proprietary Beacon Factor. This encouraged Monique to reach out to NCIM patient educator & care facilitator, Grace (Ph.D. candidate), by phone. Grace then walked her through what NCIM doctors and scientists had accomplished and documented in treated patients and then ably tackled all her questions and concerns. Satisfied with what she had learned, the Sosa’s, enrolled their son in the NCIM treatment program and read the patient handbook (This link is to the abridged version of NCIM’s patient handbook. Those who wish to access the full version must register). Later, on the heels of Angel’s approval for treatment by NCIM physicians including a pediatric neurologist, she and her husband booked him for care on 6 February (2014).

When the “big day” arrived for Angel, Grace noted this about the little boy:

While waiting for his treatment Angel was very stiff and somewhat arching backward. His eyes were moving around the room randomly and, though he would look at you if you called his name after a couple of times, he did so only for a split second and it seemed that he saw right through you. His mom had to
give him his dose of medications because he was getting a little more stiff and she knew they were wearing off.

Angel was then given primed umbilical cord stem cells and the Beacon Factor. Grace noted this at the time:

About an hour after his stem cell treatment Angel’s mom mentioned to us that both she and his father were impressed with how relaxed Angel is including his normally stiff legs. As the attending physician was checking him in recovery Angel did something amazing: He fixed his gaze on him for about 5 seconds. The doctor was both amazed and incredulous. Then Angel’s father said ‘Oh yea, he already has focused his gaze on his mom and I.’ Within an hour Angel was already aware of his surroundings and the people around him.

In addition, Angel’s legs which were normally ice cold were now pink and warm and remained so.

After Angel got home and settled in Monique shared this with Grace by phone and later by email:

The Thursday morning of his stem cell treatment he had 3 tonic seizures and then after stem cells no more seizures till late Sunday night he had 2 small quick seizures. All this week he’s only had 3 seizures! And no big seizures at all! No more 6 minute clusters :)

In therapy his therapists have noticed improvements already! He feels so much stronger! He is tolerating standing a little more and one of his therapists noticed he was using the muscles on his arm and shoulders. Everyone has
noticed that he’s more alert and happy and has more energy! On Tuesday he looked at me and smiled Mind you, he has never smiled for no reason.

Monique added that Angel’s three (3) year old sister was super excited by the fact her brother was following her all over the place with his eyes, something she instinctively regarded as a form of playing with her.

She concluded with this:

Thank you NOVA!!! I’m so excited about seeing more gains with my angel!
Thank you Grace and Abel!

The Sosa’s have set up a Facebook page for Angel which is at https://www.facebook.com/angel.sosa.52?fref=ts. This page contains photos, videos, posts and more.

CHECK OUT THIS EXCITING UPDATE POSTED ON ANGEL SOSA’S FACEBOOK PAGE (3-3-2014):

Since having done stem cells February 6, angel went 24 days without a Tonic seizure!! Before stem cells he was having them everyday 2-4 a day. It only lasted a couple seconds and recovered great afterwards. And with the cluster of infantile spasms some days it seems like there’s none and other days he will have 1-2 maybe lasting only a couple seconds and before he was having 2-4 clusters a day lasting 6-10 mins everyday!! I have also completely weaned him off zonegran his last dose was February 28. So now he’s only on 2 meds depakote and felbamate.
Roger Easterling of California Gets A New Lease on Life (Rapidly progressing multiple sclerosis)

Roger Easterling, 48, began having tinkling sensations and numbness in his hands and legs during the summer of 2014. At first he tried to ignore this but soon he found this impossible to do as one leg and foot were now working very badly, and one of his hands had become stiff and difficult to move. Soon both hands were drawn up in a fist. He was also getting tired very easily and edgy.

After seeing a neurologist and going through all kinds of tests and scans the verdict came down: Multiple sclerosis.

Roger began taking the drugs his doctor prescribed but all he did was deteriorate. He was now afraid, not just for himself but the family that depended on him including his wife and teenage children.

One of the things that really bothered Roger was watching his once considerable physical and mental energies drain away, combined with the onset of depression. And though he was sleeping more and had begun napping for twenty minutes three and four times daily, he was constantly tired and almost never felt refreshed or ready to tackle routine tasks and chores.

Like many people with a devastating disease that wasn't responding to conventional care, Roger and his wife began scouring the Web for promising new treatments including stem cells. This led him to Nova Cells during February (2015).

During the first week of March Roger went to Mexico where he received primed umbilical cord stem cells and the Beacon Factor. Within hours he was moving his stiff hand freely and within two weeks both hands were relaxed and working normally.

In the days since then Roger has enjoyed additional gains:

Prior to his stem cell & Beacon Factor treatment Roger would go golfing with his children every weekend but found that it took four hours to do nine holes and he left the country club exhausted. Now he is golfing the same course in 2 hours time and finishes up full of pep.
In the months prior to his stem cell treatment Roger and his family would go to church every Sunday, but because of his MS he did not have the strength to stand up to sing. Now he is able to do this.

In addition, whereas once he got tuckered out just carrying a gallon of water from his car into his house, now he was taking on physically demanding tasks without getting weary. For example, recently he pulled out a defective toilet in one of his bathrooms and carried it out of his house for hauling away and disposal.

Roger’s depression too is now pretty much a thing of the past.

Naturally, Roger and his family and the entire Nova Cells staff is excited by his turnaround and anticipate seeing more gains and improvements in the weeks and months ahead.

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**Multiple Sclerosis: Three recent cases**

**Mr. J. R.** is a 49 year old Mexican national who has been struggling with Progressive Relapsing Multiple Sclerosis for six years, which was rapidly progressing prior to his coming to NCIM (January 2012).

He arrived at NCIM’s treatment facility unable to walk and with his speech so slurred only his family was able to understand him. He also had difficulty swallowing as well as being unable to move his tongue very well. According to NCIM co-administrator & patient care coordinator, Grace, Mr. J. R.’s arms functioned normally but everything from the waist down plus his tongue and throat did not. He was treated with stem cells derived from three umbilical cords, Wharton’s Jelly and his own bone marrow (primed using Nova Cell’s unique laboratory method) along with NCIM’s proprietary Beacon factor.

Six weeks after his treatment Mr. J. R. was able to be understood even on the phone. He was so amazed and ecstatic with the improvement in his ability to communicate that he could not resist calling people to chat plus family and friends whom he often invited to him home to visit with him.

About three months following his treatment he was walking again, specifically taking a few steps at a time and climbing a few stairs — with assistance.
As of August (2012) Mr. J.R. can walk (slowly) *without assistance* and can also touch his palate with his tongue. He is hoping to be able to return to NCIM in January 2013 for another treatment.

**Ms. Elisa** is a 38 year old lady diagnosed with Secondary-Progressive Multiple Sclerosis four years ago. Among the things she has been wrestling with:

- Poor balance
- Lots of pain in her legs
- Rigidity in her hands and one leg
- Migraines
- Unable to walk well (Alternates using a walker and wheelchair to get about)
- Severe depression
- Poor appetite with corresponding weight loss

Ms. Elisa was treated during December 2011 with primed stem cells from Wharton’s Jelly, cord blood and her own bone marrow, along with NCIM’s proprietary Beacon Factor.

About two months after her treatment Ms. Elisa noticed her migraine headaches were becoming less frequent. In addition, she was able to balance herself to the point of not being afraid to fall any more though she continues to use her walker to get about. The rigidity in the hand and leg on one side of body has diminished and the other leg is no longer rigid though there is some light twitching which she is managing with tonic water and quinine. Also, Ms. Elisa is eating better, has gained some weight, and though she feels sad she no longer feels controlled by depression (The severe bouts of depression having completely disappeared). She reports seeing no further disease progression.
Mr. M. S., the brother of a prominent Mexican politician, is not a Nova Cell’s patient but instead was treated by his primary care doctor in Mexico using stem cells processed and primed by NCIM. He has been struggling with multiple sclerosis for eight (8) years and complained of severe fatigue, blurred and double vision in one eye, problems with his bladder, constipation, lots of muscle spasticity plus loss of muscle strength.

He was treated in February 2012 and responded so amazingly well his family brought him back to Tijuana in the middle of June to get a booster treatment.

Among the benefits Mr. M.S. has reported: His spasticity is much better especially following his second treatment, his vision is improving and he is getting more control of his bladder. His family is taking him to physical therapy which has resulted in his gaining some physical strength. Perhaps most impressive of all to Mr. M. S. is the fact that his condition has stopped progressing.
Parkinson's disease

Julia Simpson’s eye, bladder and Parkinson’s symptoms disappeared quickly following Nova Cells treatment

Emailed to Nova Cells:

This is Julia Simpson. I heard about Nova Cells Institute from other patients so I looked them up and checked out their website and testimonials. I am getting on in years being 66 and just feel the weight of the years on me. I have been pretty much healthy all of my life but as the years passed by I acquired more and more little problems which latter turn chronic. I get check-ups, take supplements, but my symptoms persisted.

I have a prolapsed bladder which makes me get up at night at least 2 times sometimes three times. This has been going on for about 6 years.

About 8 years ago a doctor prescribed Calcitonin as a nasal spray to avoid bone loss. I used it for about a month and a half. About a month into it I started to have problems with my right eye. It began tearing during the night and I had pain in it and it was red and irritated. Then headache started on that side. I did not connect this to the nasal spray. For days I had the tearing and redness even during the day and became very sensitive to sunlight. I realized that I was putting the spray in my right nostril, and the right eye was the one giving me a problem. So I then stopped using it and told my doctor. The problem is that although it stopped bothering me during the day it has never stopped giving me problems at night if I sleep on the right side, and I still had lots of pain.

I have more or less a lot of energy but I take one or two small naps a day to get reenergized.

One major thing that has been bothering me for the last 6 years is that my right hand is trembling slightly, some times more so when I get stressed. I got very worried about having Parkinson’s. I decided to get a stem cell treatment for rejuvenating my organs and hopefully helping get rid myself of the Parkinson’s like symptoms because although it is not progressing it does not go away and left me very worried about this.

On June 19, 2015 I had my stem cell treatment at NCIM. I was picked up by Abel and Grace who were so nice and professional. They answered all of my questions so I felt at ease and relaxed because I went to Mexico alone.
The city of Tijuana was just around the corner sort of speak. Just 15 minutes from the city of San Diego. The city itself was totally different from what I had imagined. It is a real nice city of 3 million people. They have roundabouts often and lots of statues of heroes, even one of Abraham Lincoln (planted in the avenue of the heroes). I understand the Lincoln statue serves as a “thumbs up” to American democracy and is a “welcome here” for US tourists. Lots of beautiful tall buildings, many made of all glass, dot the landscape everywhere. I saw McDonald’s, Costco, Office Max, Office Depot, Walmart, and a Subway Sandwich shop among others. People were milling about and walking all over the place, kind of like New York. I felt really good about Tijuana and totally safe.

Once I got to the hospital I found the staff incredibly friendly and courteous. Once I got checked in I was assessed by three doctors an internal medicine as well as a neurologist and an anesthesiologist.

The doctors did many tests and then I was infused with the Beacon Factor and got my stem cells by IV and also a percentage by spinal tap. I was not put to sleep for my spinal tap. The MDs just gave me a local anesthesia where the little puncture was made. It didn’t hurt at all. Then I got more Beacon Factor which helps the stem cells seek out damaged tissue. I was there for observation for a few hours then discharged. Abel and Grace then took me to eat at a delicious cafeteria with all kinds of Mexican dishes to choose from. Then was brought back across the border in the US.

About 6 days after my treatment all of a sudden I noticed that my eye no longer was tearing up or hurt any more. I could not believe it and decided to wait a few more days just in case it came back. It has already been 17 days and the eye is not tearing up.

My energy has shot up incredibly too. To the point where I’ve only taken naps three times in all these days, and short ones at that.

About a week into my treatment I noticed that I was not waking up at night to go to the bathroom. I was instead sleeping through the night like a baby and waking up at 8:00 or 8:30 AM each morning. I wake up so rested. I am going into deep sleep every night and am even dreaming now which was not true in the past.

My bladder, of course, is still prolapsed, but I do not have the sensation of needing to visit the bathroom over and over again each night.

But the most wonderful thing is that my hand shaking is all gone and I am ecstatic about this! About 10 days post-treatment, I was using my computer when I realized that I was not shaking any more. Of all the symptoms impacted by my treatment I am so happy and amazed that the shaking is gone! It was amazing that I felt so good I did not even notice the change at first. Thank you Nova Cells Institute! Yes, I was hoping to get rid of
my Parkinson’s symptoms, but I never thought that they would be gone in a matter of
days. Maybe it was so quick because my shaking was rather mild, but even if it had
taken months I would have been just as happy. I am not afraid or concerned any more.

Julia Simpson
Sleep Apnea

Central sleep apnea episodes fall from 66 per night to 7 in Ryan Neighbors, reports mom (Shelly)

In a Feb. 10 2014 email to Grace at Nova Cells Institute, Shelly Neighbors shared this exciting news about her daughter Ryan:

I have so much to tell you and show you!

Ryan is moving her legs on her own controlled!! We’re not walking—but this is a step in the right direction.

Not only that—but she recently had a sleep study to check on her central sleep apnea. Before stem cells, she was having 66 episodes per night. After we only had 7! They are thinking about taking her off her oxygen!

We hope to see you all again before this fall. I have videos to send but I’m at the gym. I send them when I get home. We love you and hope all is well.

Xoxo
Shelly Neighbors

Check out these prior blog entries on Ryan: https://ncimx.wordpress.com/?s=Ryan
John’s medical troubles began in 2010 with “feeling downright lousy”, the cause of which proved mystifying and hard-to-pinpoint for his regular doctors. After two months of feeling crappy he became so ill that his wife insisted on taking him to a local hospital. After various tests the hospital doctors came up with a diagnosis: John had an infection in various parts of his body including his neck bones. They proceeded to treat him with powerful antibiotics and also scraped the infected material from the vertebrae in his neck and various other places. This process helped eliminate John’s infection but, unfortunately, during one of the procedures doctors damaged or cut a nerve in his neck. This left him a quadriplegic who could not move his legs or arms or fingers. In addition, his left leg was permanently bent at the knee which meant he could only sleep comfortable on his side.

During late July 2012 John went to Mexico where he was treated with Nova Cell Institute of Mexico’s (NCIM) proprietary Beacon Factor and then primed adult (nonembryonic) stem cells.

Within 13 days following his treatment John was able to straighten out his perpetually bent left leg. He also began feeling tingling in his fingers.
By day 20 post-treatment John could flex his left leg and a few weeks later found he could lift one of his arms and touch his nose.

On October 7 (2012) while sitting in the passenger seat of his car he grabbed a tissue and wiped something from the top of his right arm with his left hand!

His wife reports that “His gains are holding up” and they are seeing more of them. Physical therapy sessions are reinforcing John’s restored bodily functioning and helping building up his strength.

Lately John has found that he can stand up more and more by himself while holding on to something and is even taking what his wife calls “baby steps”.
Spina Bifida

**Bryson Chailer: Rapid & impressive improvements in child with spina bifida with 2013 blog updates!**

I brought my son, Bryson Chailer, to Mexico during April (2012) for his first stem cell treatment and I just wanted to give you an update. First of all, let me say, Grace and Abel Pena are two of the sweetest, most caring, most hospitable people I have ever met. They are genuine and truly care about their patients and made our experience top notch. The Mexican doctor and his team and the nursing staff were all fantastic as well. We were treated professionally and with love and everything was explained to us so we could understand every single step of the entire process. We couldn’t be more impressed and will send people your way because of our wonderful experience with Nova Cells Institute.

Within 48 hours of his treatment I saw changes in Bryson and want to share these with everyone: I noticed right away that he had more energy, which is almost unimaginable, because he was already a little fire ball anyway, but he was wound for sound after the stem cell implantation! I noticed right away that he slept more soundly that night and was not tossing and turning and having interrupted sleep. He slept this way last night as well. Bryson has a vesicostomy due to bladder issues, reflux into his kidneys, and the catheter not working properly. This is temporary and to give his bladder a rest and time to repair and will be reversed. Since he urinates out of the stoma, his diaper is wet only at the top and I noticed the morning after the stem cells, his diaper was wet from top to bottom, which I think means he had been urinating, not only from the stoma, but from his penis as well. His diaper was just saturated all in the front. I noticed this same thing this morning. He pooped in the middle of the night and had not been given any kind of stool softener or enema, which is usually what we have to do to get him to poop. He pooped this morning without any enema also. His appetite has increased tremendously. He is eating more food and more often.

Also, more recently Bryson has begun having movement in his right foot and toes! He has always had movement in his left foot and toes, but never on his right foot and toes (Bryson’s level of spina bifida is at L1-L5, so he is paralyzed from the waist down). Having movement in his feet and toes is astounding! I don’t think he has any sensory or sensation yet, but he definitely has something going on to have movement. He wiggles
his toes when I run my nails down the bottom of his feet. He is actually responding to
the Babinski’s reflex test, which makes your toes flex outward when stimulated.

These are just the few things I’ve seen in my son since his treatment by NCIM. I will let
you know of any other improvements as they happen.

**Kara Underwood**

You can read more on a blog site set up to showcase Bryson’s story and especially his
response to his treatment with adult (nonembryonic) stem cells: http://www.stemcellsforbryson.blogspot.com/

**FACEBOOK ENTRIES (2012)**

**Kara Underwood**

5-11-12

Just wanted to share some progress I saw in Bryson last night! I noticed when he was on
his tummy, he lifted his right leg off the ground and held it there for quite some time! I
was just sitting there watching in amazement! Both of his legs are moving like crazy at
night too!

**Kara Underwood**

5-12-12

My mom and I were at Alco today letting Bryson pick out a toy and when he rides in the
shopping cart, his legs usually just hang down and dangle, but he actually straightened
his left leg from the dangling, bended position! Mom and I both just looked at each
other and I bent it back down and he lifted it again, and again! One happy momma and
one happy Nana! Praise God for taking the stem cells right where they need to be! Bryson continues to make progress every single day!

**Kara Underwood**

5-14-12

Such great news to share!!!! Bryson just went to therapy and the therapist says his left leg is definitely moving and extending by using his quads and hamstrings that are firing! Also, his right hip flexor is firing! God is making the stem cells work quicker than we could ever even imagine! To God be the glory!

**Kara Underwood**

5-14-12

Take That Spina Bifida!!!!!

http://www.youtube.com/watch?v=MzVCY3LHO5M&feature=share

Bryson doing leg lifts. This video was uploaded from an Android phone

**Kara Underwood**

6-4-12

Knew I would get a good report from the therapist! It just keeps on getting better! Bryson now has slight firing in his left hip! So, he has active hamstring and quad function in the left leg, now slight firing of the left hip, active firing in the right hip, and movement in both feet and toes! God is so good and thank you Jesus!

**Kara Underwood**

6-15-12

Update on Bryson since stem cell treatment! WARNING-may contain too many details and too much information! Lo!

Because Bryson has spina bifida, he has what’s called neurogenic bladder and bowel. This means the nerves don’t get signals to the brain to go potty like everyone else, so he has to take stool softeners and laxatives on a daily basis in order to go potty. He is currently refusing to take his meds and I have tried everything and I was at my wits end, so I’ve just left it in God’s hands for the past several days, and without any medication, he has pooped totally on his own, with no stool softener or laxatives! He also felt
stomach pains for the first time ever from having diarrhea, which should not happen to someone who is paralyzed from the waist down! This may not sound exciting to most people, but to a mom of a child with spina bifida, this is huge! Go stem cells go! I’ve never thanked Jesus for poop and a stomach ache before, but you bet I am now!!!!!

Kara Underwood

7-1-12

I’m one excited momma right now! Bryson just crawled on his hands and knees for the first time! He crawled 4 big steps before his legs gave way! He has never done anything but army crawl, using his arms, with his legs dragging behind him! He actually crawled on on his hands and knees by sitting on his knees and pulling them forward by using his legs! Yay! My God is so good all the time! Thank you Jesus for your continued healing! Go stem cells go!

Kara Underwood

7-1-12

My little man just crawled on his hands and knees about 6 feet without stopping! Praise God! Can I get a woot woot!

Kara Underwood

7-2-12

Thanks to God and Nova Cells Institute, Bryson is doing better than ever after his stem cell treatment and he is walking so great in his RGO braces now!

VIDEO: Go Stem Cells Go!!!!

Bryson walking in RGO braces after stem cell treatment. He is getting better and better!

“Bryson’s Story” an 8 minute, 40 second video compilation “showing progress from birth until stem cell treatment in 2012”:
http://www.youtube.com/watch?v=JqySzJ7smKA&feature=share

Kara Underwood commented on TLC (July 23, 2012)

Praying for Mama! God is still the Great Physician and Healer and the bible says that if you are all believing in total healing, and you are all standing in faith together, your Mama will be healed! Something to think about that will make a huge difference in her quality of like is stem cell treatment. Stem cell transplants are doing phenomenal things for so many diseases today and my son, age 3, who has spina bifida and is paralyzed from the waist down, got stem cell treatment 3 months ago and he is already seeing major results! He is moving his toes, crawling on his hands and knees and is walking in braces with a walker now! He is also off all medications as well! God led us down the path of stem cells for our son and He is making sure those stem cells go right where they need to be in his body and my son is being healed more and more every day! Praise God! God bless your entire family and you are all in my prayers, especially Mama :)

Stand By Mama in Her Fight Against ALS: Cake Boss: TLC

Kara Underwood

8-20-2012

Bryson is getting so good at walking in braces with his walker! Miss Elsie is just there for added support and barely has to touch his walker now! He’s doing all the work totally on his own! I am so proud of my little man! We were told Bryson would absolutely NEVER do this and look at him go! Praise God! Nova Cells Institute and their stem cells rock!

LATEST VIDEO: http://www.youtube.com/watch?v=3yggQiXU_wA&feature=share

Monday, August 20, 2012 – Rural TV Feature on Bryson and Donation Information!

http://stemcellsforbryson.blogspot.com/2012/08/well-night-we-have-all-been-waiting-for.html?spref=fb

JULY 2014: VIDEO UPDATE: Bryson Chailer walking without braces!
Kara Underwood posted this video of her son, Bryson, walking without braces to YouTube on Thursday, 7-24-2014:
https://www.youtube.com/watch?v=yIr5DJyq6k&feature=youtu.be

Bryson Chailer has some a long way thanks to faith, therapy and, yes, stem cell medicine!

If you are not acquainted with Bryson’s story here is a link to an earlier Nova Cells Institute blog entry that contains information, links, photos and more: https://ncimx.wordpress.com/2012/05/06/bryson-chailer-rapid-impressive-improvements-in-child-with-spina-bifida/

Photo of Bryson with his Mum, Kara Underwood, taken during December 2013

SEPTEMBER 2014: Facebook video & written update by Kara Underwood on her son, Bryson

Kara Underwood

Walking on the treadmill for the first time in his life! He’s in a harnessing system and only has braces from the knees down! We were told he would NEVER in his lifetime do this! All I have to say to every doctor that said NO and he CAN’T and he NEVER WILL is this…….we serve an awesome mighty God who says my son WILL WALK! He promises healing to all who ask for it as we receive it for our son in the name of Jesus!

RIKKY FORESMAN

SUN GAZETTE NEWSPAPER ARTICLE - ‘Blazin’ a trail’ - Dad: Son develops feeling, increased motion in his legs after stem cell therapy (January 16, 2014)

The Rikky Foresman saga continues: “Rikky is standing!”

Blazin for Rikky – March 3, 2014

https://www.facebook.com/BlazinforRikky

Rikky is standing!!!
READ MORE ON RIKKY FORESMAN AT
https://ncimx.wordpress.com/?s=Foresman

MARCH 2015 VIDEO OF RIKKY

http://www.novacellsinstitute.com/gallery-ncim.html

Tatyana Kushniryuk shares wonderful news about her son Rowan’s response to treatment (Spina Bifida)
My son Roman was born with Spina Bifida in 2010. He is a L4-L5 level which means he cannot void on his own and does not have bowel control. His legs were very weak and he could not walk independently until he was 2.5 years old. He was able to hold on to a walker and push it slowly but the big change came after stem cells. He received his first stem cell treatment in China, and in Mexico twice since then. The latest change was from Mexico.

We arrived to San Diego a few days before treatment and checked into our hotel downtown. The next few days we spent exploring the city and taking a trip to the San Diego SeaWorld courtesy of Warrior Families Beating Spina Bifida Foundation. Roman really enjoyed that trip because he got to see the sea lions and pet the little sharks. He was not as excited about Shamu as I was.

The next day we were escorted to Mexico across the border with Grace and Abel Pena. They were really great and picked us up in their car and drove us to the hospital. The staff greeted us as we registered in as patients, signed some papers and answered
questions about his blood type etc. I can understand how some people may be cautious signing things and giving consent to doctors in a different country but since I knew Grace and Abel from our previous trip to Mexico, I was at ease.

The doctors and nurses were very accommodating and made sure we were comfortable with our room and the procedure. It was much faster than I remember it the first time. The anesthesiologist and doctors talked about how they would only put him to sleep for a very short time and not keep him under sedation when not necessary. They had performed the spinal taps many times and knew their jobs very well. I felt confident in their care and did not have second thoughts about the procedure. I only wish that Roman cooperated as well as I thought he would. He did not like the taste of versed (the relaxing serum before procedures) so we opted to take him without it to the procedure room. The hospital has the coolest stair case that has a slide like side so they can transport people in a wheelchair up and down the twisty ramps.

During the procedure, Grace and I went to eat at a local place so that I would not be nervous and we hurried back because Roman was not going to be in the procedure very long. After we came back, Roman was not awake yet but was brought in soon after. He did great; there was no pain in his back or the IV. He was a bit freaked out thinking that the procedure was just beginning after looking at his IV. It took a while to convince him that we are all done.

After about half an hour, he began eating everything, and I mean everything he could get his hands on. The Jell-O, the rice, the juice, the fruit cup. We had asked for refills. He was feeling really great; He said “Is this Mexico? I like Mexico”…while he’s eating and watching Netflix on our laptop. (by the way, wifi works there and I would take a laptop with Netflix if you have it.)

We had a great experience and soon after we left back across the border to our hotel room. We stopped on the way while Grace and Able treated us to some local cuisine of ice cream, then sandwiches while walking around the outside mall.

Since the trip we have noticed Roman waking up in the middle of the night saying he feels his butt more and that he has to go pee. One day after about 1.5 months he had a wet diaper. He has never had a wet diaper in 6 years! Since then his recurrent infections suddenly stopped and he has been having more strength in his legs to run and walk without his AFO shoes. I am excited seeing these results they are truly worth it! The peeing is becoming more consistent and we are seeing good results in his muscle strength as well.

I encourage anyone that wants to give their kiddo a chance to improve their life to take a chance and make this trip happen.
Tatyana Kushniryuk

A Stem Cell Breakthrough for Little Ryan Neighbors by her Mum, Michelle (Click link to access all 3 pages)
**Ryan Neighbors News Release on “The Shine” (Europe’s largest spina bifida charity)**

On Sunday an NCIM press release on little Ryan Neighbors appeared on 3 PR services (24-7 Press Releases, Fire PR in the UK, and Free Press Release) and in the handful of hours since has created a bit of a positive stir. For one thing, Europe’s largest spina bifida charity picked up on tweets of the press releases and ran them on their main website at [http://paper.li/SHINEUKCharity/1319719408](http://paper.li/SHINEUKCharity/1319719408).

New video of Ryan standing ALMOST all on her own:


We are starting fundraising again this weekend. and our goal is to be back (to Nova Cells) by this summer/fall

Ryan’s core strength is MIND BLOWING. She’s breathing so much better and eating all sorts of different foods. She is also writing her name and is able to stabilize her trunk and balance when in a crawling position and when standing. Her legs aren’t doing as much as we hoped, BUT she has gained so much strength in her core and in other places I never even thought of it helping her legs. I know for a fact we wouldn’t be where we are today without your help and the stem cells. Hopefully a 2nd round will do much more.

We love you guys and can’t wait to see you again.

GOD IS GOOD. Love you guys MUCHO!!!!

Shelly Neighbors

**AND IN A FEBRURY 10, 2014 EMAIL SHELLY HAD THIS TO SHARE WITH NCIM’S GRACE:**

I have so much to tell you and show you!

Ryan is moving her legs on her own CONTROLLED!! We’re not walking – but this is a step in the right direction.

Not only that – but she recently had a sleep study to check on her central sleep apnea. Before stem cells, she was having 66 episodes per night. After we Only had 7! They are thinking about taking her off her oxygen!

We hope to see you all again before this fall. I have videos to send but I’m at the gym. I send them when I get home. We love you and hope all is well.

Page 60 of 79
Xoxo

Shelly Neighbors

**Central sleep apnea episodes fall from 66 per night to 7 in Ryan Neighbors, reports mom (Shelly)**

IN A FEB. 10 2014 EMAIL TO GRACE AT NOVA CELLS INSTITUTE, SHELLY NEIGHBORS SHARED THIS EXCITING NEWS ABOUT HER DAUGHTER RYAN:

I have so much to tell you and show you!

Ryan is moving her legs on her own CONTROLLED!! We’re not walking – but this is a step in the right direction.

Not only that – but she recently had a sleep study to check on her central sleep apnea. Before stem cells, she was having 66 episodes per night. After we Only had 7! They are thinking about taking her off her oxygen!

We hope to see you all again before this fall. I have videos to send but I’m at the gym. I send them when I get home. We love you and hope all is well.

Xoxo
Shelly Neighbors

**CHECK OUT THESE PRIOR BLOG ENTRIES ON RYAN:**
https://ncimx.wordpress.com/?s=Ryan

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**DANI SMITH**

Danielle “Dani” Nicole Smith entered September the world in 2006 with spina bifida (Myelomeningocele). Her Mum had this to share about her condition and the treatments that followed:

“With Myelomeningocele her spinal cord did not completely form and resulted in partial paralysis from the waist down. In her short life, Danielle has had five major operations. She had neurosurgery to place her spinal cord back in her body, three brain neurosurgeries, and double eye surgery. Followed by three stem cell treatments and a whole lot of physical therapy. She wears AFOs (leg braces) and uses crutches to walk. Despite everything that she has been through, Danielle
has a pleasant demeanor and is a typical six-year old. She is an absolute joy to us and we can’t imagine our life without her.”

During June 2013 little Dani was treated in Mexico by Nova Cells Institute (NCIM) using primed stem cells and its proprietary Beacon Factor. Here is an abridged version of what Dani’s Mum shared about all this on a blog set up to showcase her progress:

**July 15, 2013**

**DANIELLE HAD HER STEM CELL TREATMENT!!!**

On Tuesday, June 25th, we were picked up by Grace and Abel Pena who drove us to the Women's and Children Hospital in Mexico. Grace and Abel are truly wonderful and caring people!

First, Dr. Trujillo, an Internist came to see Dani and check her vitals. The nurses drew blood for the lab. Shortly after this the nurses gave Dani a dose of "the Beacon Factor" developed by Abel. This is to help "home" the cells to the target area. We waited for the Beacon Factor to work it's way into Danielle's system for 3-4 hours. During this time, we were visited by Dr. Montero the Anesthesiologist and the Neurologist. The Neurologist had Dani perform tasks, such as lifting her legs and arms to test her abilities.

At 6pm. the doctors took Dani in for the spinal procedure. A small amount of fluid was removed from her spine and the donated stem cells injected. A portion of the stem cells were also administered through IV. The procedure went well. Within an hour Dani was waking up and hungry. We departed to cross the border back into the US at around 9pm. It took an hour to cross the border. Attempting to cross earlier in the day can take 3-4 hours.

Dani had a headache and backache the next day. We stayed in the hotel room and relaxed while Jeff and Samantha played in the pool. I am glad that we had a few days after the procedure for Dani to begin feeling better before we flew home.

Since we've been home Dani has been great! I have slowly added back her daily exercises. She is back on the treadmill and I am working my way up to a four week at-home intensive physical therapy program. Danielle knows that we hope this surgery will allow her to walk independently and she is so excited!! She is trying to walk everywhere herself and doesn't want help. I told her that it will take a little time to see the results. When she had her own stem cells with the same procedure it was 2 months post treatment that she took her first step and 4 months post treatment for multiple steps. We expect the same timeframe of 2-4 months to begin to see really positive results.
Thank you all for helping to make this happen!!! And..... please keep the prayers coming!!

**August 20, 2013**

It's been almost two months since Danielle's stem cell treatment. She is such a determined little girl that really wants to walk independently. Dani has worked her way back to a full load of physical therapy every day, weekends off. She is using her top weights on the Universal Exercise Unit. School started today and she was up at 5:30am to exercise. We practiced walking outside for two weeks to build up her endurance for distances before returning to school.

Danielle can feel hot and cold on both of her feet. She can feel all of her toes and has more sensation on her legs!! Dani can also feel around her spinal closure on her back (basically the whole bottom half of her back). Dani is walking and attempting to walk all over the house. She is so excited and doing a great job.

We are in the process of planning for Dani to take an intensive physical therapy program. It is so important to follow up the stem cell treatment with intensive PT.
VIDEO OF DANI SMITH BELOW: Published on Nov 20, 2014

Danielle "Dani" Smith has Spina Bifida and told she would never walk. With a stem cell treatment from Nova Cells Institute Mexico, Dani is walking today. Give kids a chance to grow

https://www.youtube.com/watch?v=RUBvyFtoVm4
My name is Kristin Smith and my 2 ½ year old son, Gabe, has myelomeningocele type spina bifida affecting the L3-L4 region of his spine. He also has a VP shunt in place. When he was 1 ½ years old we took him to China for his first stem cell treatment where he received eight (8) intravenous IV treatments with umbilical cord stem cells along with acupuncture, physical therapy, fire cupping and massage. The cost was $30,000 USD. Prior to this treatment Gabe was unable to use his legs and had low hip movement as well as low sensation and poor blood flow in his legs and feet. Following his treatment he was able to move his hips enough to use an RGO brace (which basically supports his legs and moves them once he initiates movement with his hips) that allows him to walk with a walker. In addition, he had greater sensation to his knees and was able to support himself on his knees while holding onto something. He also experienced more sensation in his penis.

In May 2012 we went to Tijuana, Mexico for his second stem cell treatment with Nova Cells Institute (NCIM). We had heard many wonderful things about this group especially with regard to the rapid results that families were seeing in their spina bifida-afflicted children following treatment with Nova Cell’s primed stem cells. At the same time we also heard many negative things about the north Baja area, as I’m sure most people from the States have, e.g., violence, kidnapping, dirty, dangerous, etc. We went despite the rumors and are so thankful that we did! To our surprise Tijuana is very clean, full of nice people and a lot of uniformed young children walking around and playing after school. We ate in a local restaurant around 8 pm and never for a second felt insecure or in any danger whatsoever. It was the opposite of what we were told by well meaning friends and other who were basically just speculated wildly. We had great
food and while waiting to cross the border (back into the US) were able to buy some great souvenirs from the local people who were working hard to support their families.

As for the treatment itself, NCIM’s Grace and Abel picked us up at our hotel in San Diego and were simply the best hosts you could ever hope to find. Once at the clinic, our son did not cry when we handed him to the doctor because Abel was in the OR with him throughout the entire procedure which made him feel totally at ease. The procedure went perfectly and his legs almost immediately warmed up (better blood flow) which made both he and us so very happy. Grace stayed with us during Gabe’s treatment and recovery, comforting us and answering all our questions, and then went with my husband to get some food. Amazingly, after they returned with the food our big boy ate an entire plate of chicken and rice thirty. Hardly (30) minutes had elapsed since his treatment!

Within the first week after Gabe’s treatment we began seeing improvements. Among them:

He was able to consistently crawl while pulling his legs underneath himself.

He was able to climb up 10 stairs without help, pulling his legs up all by himself.

Also, his therapist realized that he was no longer relying on his RGO to move his legs, but was using his own leg muscles to walk!

All of these gains represented firsts for Gabe, mind you.

Now it has been almost two (2) months since our treatment and he is moving his legs from the knee, and has a lot more strength. He has more sensation further down his legs and is a lot faster and more mobile. Also, he is able to move around during play in ways that he never could before.

I would be remiss if I did not point out the vast difference between Gabe’s treatments in China and then Mexico (Nova Cells): The Chinese stem cell & other treatments took place over many weeks, cost $30K USD, and produced some modest physical improvements. The Nova Cell’s primed stem cell treatment took a single day, cost far less than the Chinese treatment, and produced very impressive improvements in Gabe literally coming out of the gate and continuing to this very day.

To say we are all more than delighted with Nova Cells and the wonderful people who make up the NCIM family is an understatement.
Readers should feel free to contact me at kristinsmith32@gmail.com but please put something in the subject line pertaining to stem cells so I do not think it is spam.

**JULY 21, 2012 VIDEO OF GABE:**

http://www.youtube.com/watch?v=u4xsGl926yo&sns=fb (28 seconds)

**ADDITIONAL VIDEOS OF GABE CAN BE FOUND BY CLICKING THIS LINK:** https://www.youtube.com/results?search_query=mospina85

**REGULAR UPDATES ON GABE’S PROGRESS ARE POSTED TO “GABE WILL WALK” ON FACEBOOK:**

https://www.facebook.com/#!/GabeWillWalk?fref=ts (NOTE: You will need a Facebook account to access this community page. If you do not have a Facebook account you can easily set one up by clicking this link https://www.facebook.com/r.php?locale=en_US)

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**Samantha Stiles benefits greatly from NCIM stem cell treatment**

Samantha “Sammie” Stiles, age 3 at the time of her treatment for spina bifida on 7-4-2012

Little Samantha came to NCIM wearing braces and using crutches to get about which made it possible for her to walk albeit with great difficulty.

In the months following her treatment by NCIM doctors using primed stem cells Samantha’s mother Jennifer reported she is walking more independently and appears to have gained strength in her legs. She added that Samantha “...walks often holding onto my hand or her little toy cart which she was not strong enough to do, she has a lot more confidence. I feel like she has gained sensation further down her legs. She says she can feel her great toes and she tells me she has to go to the bathroom more often like she has gained some sort of feeling. We have had to move her crutches up 3 notches because she has grown taller. I haven’t noticed any more ankle strength but more muscle tone and strength in her legs and now is standing taller & straighter with more confidence without her braces”.

More recently, Jennifer Stiles wrote NCIM that Samantha is has been getting ever stronger and as-a-result does not use her crutches at all anymore and is now walking and dancing all over the place using only her little braces. The Stile family are hoping that Samantha will be able get rid of her braces in the near future.
View this video of Samantha walking independently (note the huge grin)

Samantha's video

If you need a viewer, here is a free download that we at NCIM used to watch this video:
Vidolan.org/index.html
Canadian man 10 years out from a stroke recovers lost body functions in 7 days time!

Photo – Truro Daily News (2013)

Dan Huntley is an outgoing chap who had a stroke almost 10 years ago that left him paralyzed on his left side and which in-a-sense turned his world upside down. While his doctors cautioned him to accept his disability and make the best of things, Dan kept looking for ways to restore his former independence. A December 2013 Truro Daily News article on Dan titled “Truro stroke survivor has renewed hope with new car” reflects Dan’s quest for this (You can access at http://www.trurodaily.com/News/Local/2013-12-19/article-3549224/Truro-stroke-survivor-has-renewed-hope-with-new-car/1).

After reading about Arthur Rechlo’s incredible turnaround in the St. Albert Gazette (Click to access), Dan felt that any treatment that could make such a difference in an advanced dementia patient might do something good for him. He then contacted Nova Cells Institute and began a dialog with patient educator & care coordinator, Grace Odgers, Ph.D.

Long story short: Dan was treated over two consecutive days in early May (5-7 and 5-8). Within a very short period of time he reported that his energy level had shot up to what it had been when he was in his 40s and 50s, and on top of this he was feeling “tingling” in his paralyzed left side.
Dan is back home in Canada now and welcomes calls from well wishers and those interested in learning more about his Nova Cells treatment and his response to-date. Those interested in connecting with Mr. Dan should email Grace at gracepatients@gmail.com or call her at 1-562-916-3410. Those with Skype can connect with Mr. Dan via his Skype account ID which is dan.huntleyalerxgillis (If you do not have Skype, it is free at http://www.skype.com. Skype members can call or instant text other members for free).

**From an email sent Grace on 5-11-2015 by Mr. Dan:**

*I can feel bottom of left foot normal and itchy. I can relax and sleep like before stroke. Now I know I have a left side on my body. Foot working little by little. WOW!*

**And in a 5-12-2015 phone call to Grace, Mr. Dan shared these new developments:**

(1) The depression and anxiety that has dogged his steps during most of the years following his stroke has **utterly dissipated**.

(2) His memory has come roaring back (He is remembering things his stroke robbed him of going back decades).

(3) He can now lean on his left side and stand.

(4) He senses that his paralyzed left hand is about to come to life and begin working.

**And this from an email sent Grace on 5-14-15 by Mr. Dan:**

*3 pm Thursday May 14. Praise the Lord, today I walked some without assistance. No cane. I will be going 22 May for bladder tests and hope my bladder is good by then. Wow, I am coming back to life. This is truly a miracle in the making. This is the next generation in medicine [I have felt this way since I first investigated the Nova Cells treatment. Our medical system is money hungry. They must give us this great chance to be healthy as the good Lord intended us to be]. God bless you all! -Dan*

**And there is this from an email sent Grace by Mr. Dan on Saturday, 5-16-15**

(9:06 pm Nova Scotia):

*I can feel the toes on my left foot.*

Mr. Dan contacted the *Truro Daily News* about his turnaround and was interviewed by a reporter on 5-20-2015. He tells Grace that “CTV news may follow my treatment with an interview some time later”. 
In addition, he plans on seeing his neurologist in the very near future. Once this MD sees the rapid improvements he has made, he will ask for approval to enter a physical therapy program to reinforce his gains and build on them.

Dan Huntley now standing on his own! (This photo sent to NCIM on 5-17-2015)

TRURO DAILY NEWS article on Dan Huntley

Truro stroke survivor has renewed hope with new car

Published on December 19, 2013

TRURO – Dan Huntley didn't take his doctors' advice when told to accept a disability following a stroke.

Huntley, an Antigonish native who moved to Truro this past summer, suffered a stroke eight years ago this coming March and was told he would never walk again. He proved his doctors wrong and has since gotten his driver's license back.
Dan Huntley’s amazing stroke turnaround the focus of a 5-21-15
Truro Daily News (Canada) article


“I believe in it and I believe it’s the next generation in medicine, just like penicillin was,” said the Truro resident. “Regardless of what Canada and the U.S. says about it. They don’t know about it. I didn’t know about it, but I know about it now.”

Truro Daily News > News > Local

Truro man gaining strength after stem cell treatment

Raisa Tetsunish
Published on May 20, 2015

TRURO – Dan Huntley answered his door without his cane.

If it were three weeks ago, it would have been a different story. He would’ve used either his cane or a wheelchair. But things have changed for the 66 year-old after he travelled to Mexico for stem cell treatment.

“I call it fantastic,” said Huntley, who survived a stroke almost 10 years ago. “I met a lot of Godly people. It was a Godly experience.”

Huntley left for Tijuana on May 7 for a 24-hour treatment process with laboratory primed
6-27-15 update from NCIM’s chronic stroke turnaround hero, Dan Huntley (Canada)

On June 27th Nova Cells Institute (NCIM) chronic stroke patient, Dan Huntley (Photo – Truro Daily News (2013), sent this email update to NCIM’s patient educator and care coordinator, Grace Odgers, Ph.D. candidate:

It sounds like you are very busy helping the needy, Grace. Update: I now have movement in my left elbow which was previously locked at a 90 degree to 45 degree angle. Also, my walking continues to improve and I can now actually feel the leg muscles working. Amen, praise God, and God bless you, Grace, and Abel, and all the angels at the hospital in Mexico. Say “Hi “to Dr. R. for me and to all the nurses also. A big thank you and love to all, Dan ❤️👍😊😉

You can read more about “Mr. Dan’s” remarkable turnaround following his NCIM treatment at https://ncimx.wordpress.com/category/stroke/
Nova Cells - FAQs (Frequently Asked Questions)

Q. Where is Nova Cells located?

A. The corporate headquarters and all lab and patient treatments are in Mexico and all treatments are done in clinics and hospitals conveniently located near the San Diego-Tijuana border.

Q. Do Nova Cells affiliated clinics and its stem cell program adheres to the highest standards?

A. NCIM affiliated and contracted clinics are all in good standing in terms of meeting regulatory standards set forth for medical clinics by the Department of Health, State of Baja, CP, Mexico and so forth.

Q. What kind of stem cells does Nova Cells utilize to treat people?

(1) Bone marrow stem and precursor or progenitor cells taken from a small bone marrow sample (~100-150 mL) harvested from the patient.

(2) Umbilical cord blood stem cells. These come from disease-screened cord blood from Type O blood that is gender matched to the patient-recipient.

(3) Umbilical cord Wharton’s jelly-derived stem cells.

(4) Placental stem cells. These come from disease-screened placentas donated by healthy young women in Mexico who gave birth to healthy babies.

Q. How can Nova Cells doctors and scientists be sure the cells they use are safe?

A. The use of adult stem cells from cord blood and autologous bone marrow has been going on with respect to the treatment of certain blood borne conditions including leukemia for many years now in the United States and elsewhere. These typically involve the use of chemotherapy or radiation to eradicate the patient’s diseased bone marrow followed by an infusion of non-diseased bone marrow from a donor or the patient or else stem cell–rich cord blood as a way to repopulate the bone marrow with healthy tissue. Chemotherapy and radiation carries with it short-term side effects and long-term risks. But the use of adult stem cells such as those from healthy HLA matched donor
bone marrow and umbilical cord has generally proved safe over both the short- and long-term.

For its part, Nova Cells takes great, even extraordinary care to insure that the cells produced in its affiliated laboratories are carefully processed using equipment, technologies, reagents, growth factors and such that are suitable for producing cells for use in humans, and that these cells are free of microbial contamination and healthy and robust. It also contracts only physicians and other medical personnel who are highly qualified and experienced; men and women who work out of clinics and hospitals that are licensed and approved to handle not just the medical diagnostic and treatment aspects of patient care, but emergencies as well.

Q. Who chooses the specific stem or progenitor (precursor) cells or blend of stem or progenitor cells that are used to treat a specific disease or condition?

A. Typically, NCIM’s team determines which specific stem or progenitor cell or cells or combination thereof is most likely to have clinical benefits in a given instance. In some cases, a physician’s experience points to way to the best treatment choice. In others such as when there is little or no reliable clinical experience to tap nor anything of a clinical experience nature published in the mainstream scientific literature, a decision is made based on whatever can be determined from published scientific and medical literature in tandem with insights provided by theory, logic, and reason.

Q. What about Nova Cells contracted laboratories? What exactly do they do?

The hematology laboratories Nova Cells utilizes in Mexico are equipped with ultra-sophisticated equipment and adhere to the highest standards for laboratories that process cells for use in people. In addition, NCIM’s Director of Laboratory Services, Dr. Abel Pena, both utilizes and has created a number of highly innovative technologies and methods which help insure that the cells used – whether from the patient’s own body or from a donor – are both robust and apt to confer maximum clinical benefit.

Also: In some instances, adult stem cells are laboratory differentiated into whatever cell type is needed and then infused or transplanted into the target organ or tissue. For example, let’s say a patient has severe brain damage due to a stroke. In this instance bone marrow mesenchymal cells would likely be utilized and then be transformed in one of Nova Cells’ laboratories into neuronal cells. These would then turned over to the medical team for intrathecal (lumbar puncture) infusion into the spinal canal or catheter infusion into the patient’s brain or. NOTA BENE: No media containing animal derived compounds or proteins is ever used during the culturing or differentiation of cells employed by the Nova Cells program.
In other instances the stem cells are primed to become the desired bodily cell. To do stem cell priming, NCIM’s director of laboratory services, Dr. Abel Pena, takes the stem cells and utilizes specific proteins to “prime” them to respond to biochemical signals in the target tissue by becoming bodily cells that support healing and restoration.

In addition to NCIM’s proprietary priming method, stem cells are also exposed to a Beacon or homing factor (made up of specific naturally occurring and nontoxic molecules that occur in all people) that help the primed stem zero in on the target tissue(s) or organ(s).

**Q. Is there a phone number I can call to get more information on the Nova Cells program?**

A. Yes, there is. For those who prefer to speak in English or Spanish, call **1-562-916-3410** (10 am to 7 pm Pacific Time, M-F).
1-562-916-3410
NCInfodesk@gmail.com

Nova Cells Director of Laboratory Services, biochemist Abel Pena
The Nova Cells process at-a-glance

**Documentation**

Enrollment in the Nova Cell Program. Provide copies of medical records.

**Evaluation**

Evaluation of medical history, diet and lifestyles by Nova Cell doctors and staff. Nova Cell MDs and its scientific team may request additional testing or information.

**Acceptance in the program or not**

Acceptance or denial of acceptance into the Nova Cell program.

**Treatment proposal**

For those accepted into the Nova Cell Program: Nova Cells medical team craft a treatment plan proposal that is submitted to the patient.

**Customized diet & lifestyle plan**

Following acceptance of the treatment plan, Nova Cells doctors and other professionals create a customized dietary & lifestyle guide and submit this to the patient.

**Treatment & follow-up**

Patient is then treated and tracked for responses.
How to get more information or register

YOU CAN GET MORE INFORMATION BY PHONE OR EMAIL:

**Phone:** 1-562-916-3410.

**E-mail:** NCinfodesk@gmail.com

TO GAIN ACCESS TO NOVA CELL'S DETAILED HANDBOOK: