

Sound Choice Pharmaceutical Institute

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June 2011 Newsletter

Editorial Contents:

1) Misleading media labels moral adult stem cell therapies 'controversial'!

Embryonic stem cell proponents attempt to hijack the stem cell issue by labeling moral adult stem cell therapies as controversial. Don't be fooled by their outrageous strategies.

2) Multiple independent scientists publish studies supporting Sound Choice's hypothesis that vaccine contaminants may be linked to autism.

Autism is associated with HUNDREDS of new gene mutations not found in the parents. How did those mutations get there? Sound Choice is the organization leading this effort, and we do our science in morally compatible ways.

3) We need your help to purchase equipment!

We have \$5,000 towards this purchase. Please help us reach our \$35,000 target.

PLEASE DONATE TODAY! WE RELY ON YOUR SUPPORT!

Dr. Theresa Deisher, President, Sound Choice Pharmaceutical Institute

Misleading media labels moral adult stem cell therapies 'controversial'!

USA Today (June 29, 2011 issue <http://yourlife.usatoday.com/health/story/2011/06/Doctors-offer-unapproved-stem-cell-therapies/48933666/1>) carried a touching story

recently about New York Yankees pitcher Bartolo Colon whose fast ball is back this season after being treated with his own stem cells. Colon sat out 2010 because of injuries, and had not completed a full season since 2005. Now he is 39 and pitched a shut out May 30. How was he able to come back? He was treated with his own stem cells! So why would USA Today label the treatments 'untested', 'unapproved' and controversial when adult stem cells take a sidelined pitcher like Colon and put him back on the mound?

When faced with the evidence of the benefits of adult stem cell therapies, proponents of embryo destroying stem cells can only grab at half-baked accusations and mislabels to try to maintain their position that destroying embryos is the only way to go. What should be controversial is the question of why Colon had to go to the Dominican Republic for his adult stem cell treatments! Why can't human beings get these treatments in the United States? What about the people who cannot afford to fly to the Dominican Republic to be treated? Why can't US tax payers be treated in the US where they can be assured of the quality of their care? We covered this in our last newsletter. US tax payers cannot get moral adult stem cell treatments in the US because embryonic stem cell proponents do everything they can to keep these moral stem cells out.

Dogs, horses and cats are treated with safe, effective and affordable adult stem cells in the US. Only humans are kept from these therapies. If you asked a veterinarian why they don't use animal embryonic stem cells for their

therapies they would tell you "Embryonic stem cell therapies are expensive and the stem cells trigger immune responses and form tumors. Why would we use expensive and unsafe embryonic stem cells when we can treat the dogs and horses with safe, effective and affordable adult stem cells?" We should all be demanding that our elected officials, the NIH, the FDA, the scientists and physicians answer this question for humans: "Why the focus on embryonic stem cells that are expensive and trigger immune rejection and tumor formation? Why can't US tax payers be treated in the US with safe, effective and affordable adult stem cell therapies?"

The USA Today article mentions the 'anger' that some US scientists expressed that patients are being hood-winked by charlatans outside of the US charging high prices for adult stem cell therapies. We should be angry. However, let's direct that anger where it belongs! We should be angry at the scientists who fraudulently and viciously attacked adult stem cell research, we should be angry at the politicians who chose 'anti-life' stances over pro-patient stances, and we should be angry that the NIH and FDA have not led the way in making sure that US tax payers get safe, proven, effective and affordable adult stem cell therapies.

Multiple independent scientists publish studies supporting Sound Choice's hypothesis that vaccine contaminants may be linked to autism.

Even my local newspaper, The Seattle Times, picked up this story on June 09, 2011 (<http://community.seattletimes.nwsources.com/mobile/?type=story&id=2015268511&>).

Three publications in the journal Neuron (1,2,3) have now established that autism is caused by hundreds of gene mutations, many of these are 'spontaneous'

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new mutations in the children only. Spontaneous or 'de novo' mutations are gene mutations that the parents do not have. This means that some environmental trigger caused gene damage to hundreds of genes in hundreds of thousands of children. The scientists who published these studies, scientists from Yale, UCLA, Emory, Carnegie Mellon, Vanderbilt, Baylor, Cornell Medical College, Stanford, Princeton, Harvard, the University of WA, Brown University, Columbia University, Albert Einstein College of Medicine and others, lament the fact that their results, however, will have no immediate benefit for the children with autism or for their families. The scientists, however, expect their reports to benefit them, since they now have hundreds of new genes that they can write grants about. It seems that Sound Choice Pharmaceutical Institute is the only organization working for the patients' benefit.

These academic scientists are focused on the downstream effects of all the new mutations they have discovered. Sound Choice Pharmaceutical Institute is focused on the upstream causes, the environmental trigger, of all these new mutations. It is only by identifying the causes of all these hundreds of mutations that we can help prevent new cases of autism and find treatments for those who already suffer from this disease. What is the cause? Manufacturing vaccines using electively aborted human fetal cell lines!

Autism is not an epidemic unique to the US. Autism is now a worldwide epidemic. Since 2001 or so, third world countries are experiencing autism at the same levels as the US ⁽⁴⁾. What do children in Africa and Asia have in common with children in the US? They eat different food, drink different water, they don't watch TV or play computer games excessively, they don't sit on fire retardant couches. What they have in common since 2001 is that they are vaccinated with vaccines containing contaminating aborted human fetal DNA and human retroviruses. Humanitarian vaccination campaigns undertaken by the World Health Organization ⁽⁵⁾ and other agencies to eliminate measles have inadvertently introduced a new disease to these countries : autism. The vaccines used in those campaigns do not contain mercury! The vaccines are, however, manufactured by using aborted fetal cell lines. Vaccines that are manufactured using aborted fetal cell lines contain unacceptably high levels of contaminating human fetal DNA and human retroviruses. Those contaminants are known triggers for gene mutations!

We need YOUR help to accelerate our research to prove the dangers of using aborted fetal cells for vaccine production. We have a \$5,000 donation to purchase a refurbished piece of equipment - we need \$30,000 more. Please give generously!

- 1) *Rare de novo and transmitted copy-number variation in autistic spectrum disorders.* Levy D, et. al., 2011, Neuron, 70:886-897.
- 2) *Rare de novo variants associated with autism implicate a large functional network of genes involved in formation and function of synapses.* Gilman SR, et' al., 2011, Neuron 70:898-907.
- 3) *Multiple Recurrent De Novo CNVs, Including Duplications of the 7q11.23 Williams Syndrome Region, Are Strongly Associated with Autism.* Sanders SJ, et. al., 2011, Neuron, 70 :863-885.
- 4) *Pakistan Newswire January 6, 2004. "Autism is on the rise in India, Pakistan, Bangladesh, Sri Lanka, Nepal, Bhutan and the Maldives - the countries within the SAARC region."* .s.l. : Pakistan Newswire, 2004.
- 5) *ARC, CDC, WHO, UNICEF, UNF. Measles Initiative. The Measles Initiative.* [Online] [Cited: June 5, 2011.] <http://www.measlesinitiative.org/mi-files/Reports/Measles%20Initiative/Measles%20Initiative%20Fact%20Sheet%20Jan%202011.pdf>.

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Sound Choice Pharmaceutical Institute
1102 Columbia Street Suite 316-322
Eklind Hall

Seattle, WA 98104

OR to

P.O. Box 2247

Seattle, WA 98111