

CALIFORNIA RIGHT TO LIFE EDUCATION FUND

P.O. Box 4343, Walnut Creek, CA 94596-4343

(925) 944-5351

E-Mail: callife@calright2life.org

Web Site: www.calright2life.org

Established 1981

April 2016

Gardasil Destroys Girl's Ovaries: Research on Ovaries Never Considered

Another article in our ongoing effort to keep you informed of negative findings regarding the Gardasil vaccine

A girl's ovaries were destroyed, with Gardasil the only potential cause. Worse, though, is that Merck either didn't bother to examine potential effects on ovaries or hid them—but did examine effects on testes.

The British Medical Journal (BMJ) has published the case report of a healthy 16-year-old Australian girl whose womanhood appears to have been stolen by Gardasil vaccinations. She has been thrust into full-fledged menopause, her ovaries irrevocably shut down, before becoming a woman. The authors, Deirdre Therese Little and Harvey Rodrick Grenville Ward (1), draw direct attention to the fact that, though the girl has been thoroughly examined and tested, there is no known explanation other than the series of three Gardasil vaccinations she had.

Making matters worse is that there may be many other such cases, but most are likely masked by the routine treatment of irregular or scanty menstruation with oral contraceptives. Indeed, it's only because this girl refused them that the truth of her situation was unmasked. Just how many other girls have lost their chance at motherhood, but don't know because their condition is masked?

The authors noted that, although the Therapeutic Goods Administration (TGA) of Australia provides data on the histology of rat testes and epididymides in the Australian Public Assessment Report for Human Papillomavirus Quadrivalent Vaccine, (Note: There is only one "quadrivalent HPV Vaccine." It's Gardasil.) no information is provided for rat ovaries. They sent a Freedom of Information request for "documented rat ovarian histology post-quadrivalent* HPV vaccination that may have been performed by the sponsor and

forwarded to the TGA." Here is their report of this highly significant missing data:

It is not known whether this event of premature ovarian failure is linked to the quadrivalent HPV vaccine. More detailed information concerning rat ovarian histology and ongoing fecundity post-HPV vaccination was sought from the Therapeutic Goods Administration (TGA). Although the TGA's Australian Public Assessment Report for Human Papillomavirus Quadrivalent Vaccine, February 2011, does report on the histology of vaccinated rat testes and epididymides, no histological report has been available for vaccinated rat ovaries.

The TGA subsequently agreed to a freedom of information application in the public interest (FOI 001-1112) requesting documented rat ovarian histology post-quadrivalent HPV vaccination that may have been performed by the sponsor and forwarded to the TGA. However, a histological report of the ovaries of vaccinated rats remained unavailable beyond a numbering of the corpora lutea present at postweaning euthanasia following the first litter.

Why did the manufacturer provide information regarding male rat testes, but not for female rat ovaries? This is more than a little shocking. It's absolutely damning! We must question the sincerity of both the manufacturer, Merck, and the TGA—not to mention questioning other regulatory agencies, such as the US's FDA and the UK's MHRA.

Potential Gardasil Risk to Ovaries

Is it conceivable that Merck didn't consider the possibility of harm to the ovaries? In point of fact, it is unreasonable to suggest that they were unaware of potential harm to ovaries. At least one Gardasil ingredient, polysorbate 80 (also called by brand names Tween 80, Alkest, and Canarcel) is a known cause of ovarian deformities, degenerative follicles, hormonal changes, and womb and vaginal changes in rats (2,3.) Worse, that ovarian damage is known to be caused by

injection of polysorbate 80—just as it is injected with Gardasil.

Another Gardasil ingredient, L-histidine, a naturally-occurring amino acid, carries significant risks, too, in the same manner that squalene does. It's a naturally-occurring substance in the human body, so injecting it could have the effect of causing an autoimmune response to that substance wherever found in the body.

A large part of one girl's life has been destroyed, and the only plausible explanation is that the cause is the Gardasil vaccination. This vaccine is sold as a cervical cancer preventative, though it has never been shown to prevent any cancer of any kind. Cancer prevention has never been more than a presumption, based on a possible connection between human papilloma virus and cervical cancer. No cause-and-effect has ever been documented.

We do not know, of course, whether Merck calculatedly avoided doing the studies on ovaries or is refusing to release data on such studies because of its damning nature. We do, though, know that the very fact that it is missing—especially in light of equivalent data on the male reproductive tract being available—must be treated as suspicious.

Certainly, the combination of one girl's loss of her ovaries, the probability of there being many others, and the utterly callous disregard for its potentially devastating effects, is more than enough reason to remove Gardasil from the market. Surely, it should be removed from governmental lists of mandated vaccines.

Sources

1 Premature ovarian failure 3 years after menarche in a 16-year-old girl following human papillomavirus vaccination, *BMJ Reports* 2012, Deirdre Therese Little, Harvey Rodrick Grenville Ward, doi:10.1136/bcr-2012-006879

2 Polysorbate 80 Causes Infertility, An Emulsifier That Can Damage Your Reproductive Health

3 Delayed effects of neonatal exposure to Tween 80 on female reproductive organs in rats. Gajdová M, Jakubovsky J, Váľky J., *Food and Chemical Toxicology*, 1993 Mar;31(3):183-90.

Original article: <http://gaia-health.com/gaia-blog/2012-10-17/gardasil-destroys-girls-ovaries-research-on-ovaries-never-considered/>

Haematopoietic stem cell transplants to treat multiple sclerosis

Several hospitals around the world (United States, Brazil, Sweden and the United Kingdom) have been taking part in an international study since 2006 in order to “evaluate the long-term effects of stem cell transplantation.” In this respect, the Royal Hallamshire Hospital in Sheffield, United Kingdom, has tested a treatment for multiple sclerosis patients.

Multiple sclerosis is an auto-immune disease that affects the central nervous system, especially the brain, optic nerves and spinal cord. The immune system of those affected destroys myelin [1] by viewing it as a foreign body. Thus, in certain places in the nervous system, the impulses are slower or completely blocked, leading to various symptoms such as numbness, muscle spasms, lack of mobility or imbalance.

The treatment tested in the United Kingdom comprises a “haematopoietic stem cell transplant [2]” (HSCT), which is performed following chemotherapy. Doctors collect a small quantity of the patient's haematopoietic stem cells in a blood sample, freeze them in order to preserve them and then readminister them intravenously after chemotherapy. The transplant allows the patient to reconstruct his/her blood and immune system following chemotherapy.

Out of the twenty patients treated, “those who were paralyzed were able to walk again.” Professor Basil Sharrack at Sheffield's Royal Hallamshire Hospital welcomed this “major success.” Doctors are optimistic but remain cautious: “the tests have shown that the marrow transplant may be able to stabilize or improve a disability for some patients with multiple sclerosis. It may, however, prove ineffective for other diseases.”

Studies are continuing to evaluate the “safety of the procedure and its long-term effects.”

[1] Myelin forms a sheath, which surrounds nerve fibres thereby protecting them and accelerating the transmission of messages or nervous impulses.

[2] Haematopoietic stem cells are multipotent adult stem cells taken from the bone marrow.

Sources: <http://genethique.org/en/haematopoietic-stem-cell-transplants-treat-multiple-sclerosis-64834.html#.VrtdWua0ejc>

Adult Stem Cells Cure Blindness in Babies Born with Congenital Cataracts

Extract of article by Wesley J. Smith

If this human breakthrough had occurred with embryonic stem cells, the front page stories would have screamed around the world. But it was adult stem cells and so the reporting was muted. Sadly, the media still – after all these years – tend to judge the newsworthiness of a story based on whether a breakthrough is embryonic. The story is sensational, nonetheless.

Adult stem cells have cured blindness and may provide a splendid treatment for cataracts.

In a recent article in the UK Telegraph comes the following:

Cataracts can be cured by using a patient's own stem cells to regrow a 'living lens' in their eye, restoring sight in just three months, scientists have shown. In research described as 'remarkable,' surgeons reversed blindness in 12 infants born with congenital cataracts by removing the damaged lens and coaxing nearby cells to repair the damage.

This is great news. And the potential is really exciting:

“An ultimate goal of stem cell research is to turn on the regenerative potential of one's own stem cells for tissue and organ repair and disease therapy,” said Dr Kang Zhang, chief of Ophthalmic Genetics and founding director of the Institute for Genomic Medicine at UC San Diego School of Medicine.

“The success of this work represents a new approach in how new human tissue or organ can be regenerated and human disease can be treated, and may have a broad impact on regenerative therapies by harnessing the regenerative power of our own body.”

Telegraph article available at
<http://www.telegraph.co.uk/news/health/news/12189238/Scientists-use-stem-cells-to-grow-living-lens-in-eye-and-cure-cataracts.html>

California Legislative Special Session Adjourned - Dangerous California Assisted Suicide Law Will Take Effect in June 2016

On March 10th following the close of California's Special Session on healthcare funding, the 90-day clock begins before California's assisted suicide law becomes operative on Day 91. The End of Life Option Act, commonly referred to as the Physician Assisted Suicide Bill will officially take effect on June 9, 2016.

The close of the special session to address Medi-Cal shortfalls, important disability services and in-home care program funding, is a reminder of the controversial legislative tactics taken by proponents in order to narrowly pass the assisted suicide bill.

Californians Against Assisted Suicide, a coalition of which California Right to Life Educational Fund is a member, remains strongly critical of both this new law and its lack of medical oversight and actual patient safeguards. We will continue to work with our coalition partners including doctors, patients and disability rights organizations to educate those impacted and vulnerable, as well as working to limit the law's harms and prevent any expansion.

For further information and ongoing developments as we continue to fight against this policy, visit the following websites: <http://stopassistedsuicide.com/> and <http://noassistedsuicideca.org/>

**Do you know someone who might
be considering abortion?**

**Make sure they get the facts first!
A LIFE depends on it...**

1-800-712-HELP (4357)

Website: <http://www.optionline.org/>

Calendar of Events

*For the latest updates of events see
www.calendarforlife.org*

#ProtestPP Saturday, April 23 - #ProtestPP is hitting the ground with the first annual nationwide protest at Planned Parenthood facilities, building on the momentum created at the coast-to-coast protests held in 2015. This nationwide protest will take place in future years on the 4th Saturday of April. For further information and to find a location near you, visit:
<http://protestpp.com/#home>

It may not be too late...

Abortion Pill Reversal

**If you know someone who has taken the
Abortion Pill (Mifeprex or RU-486)**

Call (877) 558-0333

<http://www.abortionpillreversal.com>

WHO IS CALIFORNIA RIGHT TO LIFE?

This is the newsletter of **California Right to Life Education Fund**, a 501-c-3 organization established to educate the public about pro-life issues. Donations to the EDUCATION FUND are **tax-deductible** and can be sent to P.O. Box 4343, Walnut Creek, CA 94596-0343.

California Right to Life **Committee, Inc.** is a 501-c-4 organization providing information on legislative issues affecting the right to life, and pro-life political advocacy. **CRLC, Inc. is not permitted**, under IRS regulations, to offer a tax deduction for donations. \$24.99 annually is requested for a subscription to the CRLC legislative email updates list and can be sent to 1920 Monument Blvd #309, Concord, CA 94520.

Both are affiliates of American Life League, headed by Judie Brown, and share the same "no-exceptions, no excuses" beliefs and the same dedication to promoting the Culture of Life, respecting all innocent human life from the single-cell stage to natural death.

Newsletter printed by HMR Printing

925-680-0388

Consider HMR Printing for your next printing job