2. Cholesterol Sulfate
Stephanie Seneff
Wise Traditions Workshop
November 8, 2013
"If we all worked on the assumption that what is accepted as true is really true, there would be little hope of advance."

-- Orville Wright
Outline

• Introduction
• Cholesterol sulfate
• Heparan sulfate proteoglycans (HSPGs)
• Atherosclerosis is protective
• How statins really work explains why they don’t really work
• Maintaining blood stability
• Cancer to the rescue?
• Summary
Introduction
A Provocative Proposal

• Cholesterol sulfate supplies oxygen, sulfur, cholesterol, energy and negative charge to all the tissues

• Sulfate is synthesized from sulfide in skin and blood stream utilizing energy in sunlight
  – Protects from UV damage and keeps microbes out

• Endothelial Nitric Oxide Synthase (eNOS) performs the magic

The skin is a solar powered battery!
Cholesterol and Cholesterol Sulfate

Sulfation makes cholesterol water-soluble and therefore much easier to transport.
When cholesterol and sulfate “hold hands” they can navigate the waters together
Think about Sulfate!

• Cells in the skin produce vitamin D3 sulfate upon exposure to the sun
  – The precursor to vitamin D3 is 7-dehydrocholesterol

• Cells also produce an abundance of cholesterol sulfate
  – I believe this is the more important molecule!

• Many of the alleged benefits of vitamin D3 are actually benefits of cholesterol sulfate
  – Protection against cancer, diabetes and cardiovascular disease; improved immune function
Heart Disease Mortality and Sunlight*

*Grimes et al., Q. J. Med. 1996; 89:579-589
Lack of Sunlight May Raise Stroke Risk*

• Strokes are caused by blood instability: clots and hemorrhages
  – Is this due to insufficient sulfate in blood?
• Study involved 16,500 people
• Tracked the history of where they had lived
  – Looked at weather statistics for those places
• Found 60% increased risk to stroke for lowest sun exposure (relationship with both latitude and weather patterns)

*A. Mozes, HealthDay, RSS Feed, Feb 2, 2012.*
Ultraviolet Exposure and Mortality among Women in Sweden*

- 38,472 women selected in 1991-1992, aged 30-49
  - monitored for 15 years
- Questionnaire asked about frequency of sunbathing vacations and sunburn
  - *Increased* sunburn frequency associated with *reduced* all-cause mortality
  - Sunbathing vacations more than once a year *reduced* risk to cardiovascular disease and mortality

* Yang et al., Cancer Epidemiol Biomarkers Prev. 20(4):683-690, 2011
Skin Melanoma Increasing 2%/Yr since 1974*

* Andrew Schneider, Aol News, May 24, 2010

aolnews.com/2010/05/24/study-many-sunscreens-may-be-accelerating-cancer
How to Stay Healthy

• Plenty of dietary sulfur
• Plenty of dietary cholesterol
• Plenty of sun exposure
Cholesterol Sulfate
Cholesterol is a Miracle Worker

• In the brain:
  – Synapse: promotes cell-cell communication
  – Myelin sheath: insulates channel from signal loss

• In the membranes of all cells
  – Prevents ion leaks
  – Protects from pathogens (microbes)

• In the plasma lipoprotein (LDL, HDL)
  – Essential for protecting contents from oxidation and glycation damage during transport to cells

• Precursor to vital hormones
  – Vitamin D
  – All the sex hormones (testosterone, estrogen, etc.)
  – Cortisone: the stress hormone

• Aids in digestion of fats
Cholesterol is Essential for Insulin Release *

- Experiment with pancreatic beta cells
- Expose them to toxin that impairs cholesterol synthesis
- Cells secrete significantly less insulin in response to glucose challenge

* Xia et al., Endocrinology 149:10, 5136-5145, 2008.
Cholesterol is Essential for Insulin Release *

- Experiment with pancreatic beta cells
- This is one reason why statin drugs increase risk to diabetes
- Significantly less insulin in response to glucose challenge

* Xia et al., Endocrinology 149:10, 5136-5145, 2008.
Sulfate is Vastly Underappreciated!

• Sulfate is the 4th most abundant anion in the blood and protects it from coagulating.
• Detoxifies drugs, food additives, and environmental toxins like aluminum and mercury.
• Essential component of extracellular matrix proteins throughout the tissues.
• Cerebroside sulfate is a major constituent of myelin sheaths surrounding axons in neurons.
Cholesterol Sulfate Deficiency

• Cholesterol sulfate deficiency leads to filaggrin deficiency which is associated with:
  – Atopic Dermatitis
  – Eosinophilic Esophagitis
  – Asthma

• These three conditions are on the rise and are associated with autism and celiac disease
Cholesterol Sulfate Synthesis in the Skin

• Keratinocytes synthesize abundant cholesterol sulfate as they mature
  – This synthesis takes place in outer layer of the epidermis

• Cholesterol sulfate catalyzes synthesis of profilaggrin, one of the two key proteins forming the cross-linked mesh that protects from bacterial invasion and prevents water loss
Impaired wound healing in atopic dermatitis

- Water loss
- Microbe penetration

*J.A. Segre, J Clin Invest. 2006;116:1150-58*
Filaggrin Plays a Crucial Role

- Loss of filaggrin function associated with
  - Increased skin permeability
  - Susceptibility to eczema and asthma
  - Eosinophilic Esophagitis: (EoE)*

EOE: A Modern Disease*

- Allergic inflammatory condition of the esophagus
- Eosinophils invade esophagus
  - Release toxins that kill viruses and induce tissue damage
- Person has difficulty swallowing
  - Eats very slowly
- Food gets stuck in the throat

A Very Bad Idea
Cholesterol Sulfate in the Lungs*

Bronchial tubes produce cholesterol sulfate in the epithelial cells as they mature and form the outer layer of the bronchial epithelium

Cholesterol Sulfate in Horses’ Hoofs! *

- 37-43% cholesterol
- 15-20% cholesterol sulfate

Laminitis (Founder)

• Lameness in horses due to inflammation in the hoof
• Can be caused by exposure to herbicides and nitrate fertilizers

Is This due to Impaired Cholesterol Sulfate Supply to the Hoof?
Cholesterol Sulfate and Trypsin*

• Trypsin is an enzyme released by the pancreas, which digests proteins

• Too much trypsin leads to breakdown of collagen glue that maintains structural integrity of intestinal mucosa (GAGs)

• Cholesterol sulfate suppresses trypsin synthesis (protects mucosa from breakdown)

Cholesterol Sulfate Inhibits Trypsin *

* Sato et al., J. Investigative Dermatology, 11:2, 189-193, Aug 1998
Recapitulation

• Cholesterol sulfate plays many important roles in the body: skin, blood, esophagus, lungs, and pancreas

• Cholesterol sulfate in the skin and lungs keeps bacteria out and water in

• Horses’ hooves are loaded with cholesterol sulfate

• Cholesterol sulfate protects collagen from breakdown by trypsin
Heparan Sulfate Proteoglycans (HSPGs)
Sulfated Biomolecules*

• Sulfated mucopolysaccharides (glycosaminoglycans)
• Steroid sulfates
• Phenol sulfates
• Tyrosine sulfation
• Bilirubin sulfate
• Arylamine sulfates
• Choline sulfate in lichens, fungi and marine algae
• Sulfolipids

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Sulfated Glycosaminoglycans (GAGs)

- Prominent in extracellular matrix of all cells
- Amount of sulfate depends on availability
- Crucial for maintaining negative charge and protecting from infection

[Diagram showing structures of sulfated GAGs]

http://www.science-autism.org/sulphate.htm
Sulfated Glycosaminoglycans (GAGs)

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These are also known as “mucopolysaccharides”

http://www.science-autism.org/sulphate.htm
Ten Positive Effects of Mucopolysaccharides *

- A lipid-clearing effect in the blood.
- Stimulation of cellular metabolism.
- Efficient metabolism of fatty acids.
- Increase in RNA and DNA synthesis of cells.
- Increase in growth, size and quantity of normal cells.
- Anti-atherosclerosis, anti-atherogenic activities.
- Anti-inflammatory effect.
- Anti-thrombogenic and anti-coagulant activity.
- Increases the number of coronary artery branches and collateral circulation in experimental atherosclerosis.
- Accelerates healing, regeneration and repair of cardiovascular tissue

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Many of these effects can be explained by the fact that they supply sulfate.
Heparan Sulfate: Wonder Worker

Polymers of sugars with attached nitrogen and sulfates: safe glucose storage
Heparan Sulfate in Pancreas *

- Beta cells in pancreas produce insulin
- They also produce an abundance of heparan sulfate
- In type I diabetes, inflammatory cells break down basement membrane and penetrate pancreatic islets
- These inflammatory cells produce heparanase, which destroys heparan sulfate
- This leads to cell death of beta cells and diabetes

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Various Factors that Increase Sulfate*

An increase of the 35S-sulfate deposit is effected regularly by infections, by injections of toxins and proteins, by hypoxemia, dietetic influence, muscular over-exertion, weather influence and increase of the blood pressure.

SMPS = Sulfated mucopolysaccharides = glycosaminoglycans

*Report by Bernhard Muschlien

First published in the German language in the SANUM-Post magazine (17/ 1991)
A Provocative Hypothesis

Cells store glucose temporarily in HSPGs, protected by sulfate

When cholesterol and sulfate are in short supply, storage of sugars in HSPGs becomes impaired

This is the source of insulin resistance and diabetes!
Disrupted Signaling when HSPGs are Depleted*

HPGs = Heparan Sulfate Proteoglycans

*Figure 5 in P Muller and AF Schier, Developmental Cell 21, July 19, 2011, 145-158.
Sulfotransferases
(Transfer sulfate from one molecule to another)

- Carbohydrate sulfotransferase
- Galactose-3-O-sulfotransferase
- Heparan sulfate sulfotransferases (2-O, 3-O, 6-O)
- N-deacetylase/N-sulfotransferase
- Tyrosylprotein sulfotransferase
- Uronyl-2-sulfotransferase
- Estrone sulfotransferase
- Chondroitin 4-sulfotransferase
Recapitulation

• Heparan sulfate proteoglycans (HSPGs) are everywhere in the body
  – They play a crucial role in ion transport, nutrient uptake, and cell signalling
  – They protect pancreas during insulin production
• I propose that they also provide a temporary storage depot for glucose
  – Deficient sulfate leads to diabetes
• Multiple sulfotransferases attest to the importance of sulfate to the body
Atherosclerosis is Protective!
End Stage Kidney Disease and Cardiovascular Disease*

- High serum cholesterol
- High blood pressure
- High serum homocysteine
- Obesity
- Inflammation

Dialysis patients have extreme risk to cardiovascular disease

At low cholesterol, low blood pressure, low homocysteine, emaciated

My Conclusion from This

Inflammation is the ONLY “Risk Factor” that counts: All the other “Risk Factors” are protective measures!
Sulfates in the Kidneys*

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Statins and Kidney Failure*

• Kidney failure is alarmingly on the rise in the US and elsewhere
  – Associated with a 23-fold increase in risk to heart disease!

• Statins are being overprescribed for patients with kidney failure
  – Severe muscle pain & rhabdomyolysis
  – Increased risk to dementia and diabetes
  – No improvement in kidney function

* M.A. Lanaspa et al., Nature Communications, 2013; Aug 24. [Epub ahead of print]
Heart Disease: A Theory!

Cardiovascular plaque develops as an alternative mechanism to produce cholesterol sulfate from damaged LDL and homocysteine.
A Recent Paper


“Is Endothelial Nitric Oxide Synthase a Moonlighting Protein Whose Day Job is Cholesterol Sulfate Synthesis? Implications for Cholesterol Transport, Diabetes and Cardiovascular Disease.”

Entropy 2012, 14, 2492-2530.
What Happens when Cholesterol Sulfate Synthesis is Impaired??

Cardiovascular disease!!

Activities in Plaque Produce Cholesterol Sulfate to Supply the Heart!
They Knew a Long Time Ago*

- Article published in 1960
- Fed cholesterol to monkeys – induced atherosclerosis
- If sulfur-containing nutrients are added, atherosclerosis is prevented
- These nutrients provide source of sulfate to enable cholesterol transport

They Knew a Long Time Ago*

"A form of vascular disease resembling human atherosclerosis has been produced in the New World primate Cebus fatuella. ... In order to produce these phenomena, the diets had to be rich in cholesterol, choline and neutral fat but relatively low in organic sulfur compounds. Without this deprivation of organic sulfur the response of the serum lipids to cholesterol feeding was small."

“Heart Stents Still Overused, Experts Say”*

- Two misconceptions
  - Heart attack caused by blocked artery
  - Eating fatty foods induces fat build-up
- Heart attack is actually caused by thrombus formation (blood clot) that often occurs in a region of the artery where plaque is not obstructive
- Stent insertion is expensive and is not living up to the promise of protection

*Anahad O’Connor, NY Times, Aug. 16, 2013
Steps in Atherosclerosis*

1. Inflamed endothelium provides adhesion molecules to trap and hold macrophages
2. Macrophages through scavenger process take up oxidized LDL and become foam cells
3. Interleukins and growth factors promote proliferation of smooth muscle cells (artery thickening)
4. Extracellular matrix proteins are degraded
5. Vulnerable plaque eruption: thrombosis

* Libby et al., Circulation 105:1135-1143, 2002
Cardiovascular Plaque*

*Figure 3 in L. Badimón et al., Rev Esp Cardiol. 2009;62(10):1161-78 (p. 1163)
Many Good Reasons for ROS

• ROS (reactive oxygen species) are a key component of inflammation in the artery
• ROS are needed to produce sulfate*
• Oxidation of glycated LDL makes it accessible to macrophages for breakdown**
• Peroxynitrite (product of reaction between superoxide and nitric oxide) is toxic to pathogens***

Macrophages and Cholesterol*

- Macrophages in artery wall take up oxidized LDL and export extracted cholesterol to HDL-A1
- Macrophages eventually become damaged by exposure to oxidizing and glycating agents → necrotic core

Macrophage Proliferation in Plaque*

*Figure 4, C.S. Robbins et al., Nature Medicine, Aug. 2013 [ePub ahead of print]
Declaring War on Macrophages*

• Macrophages replicate inside plaque
  – Suppressing their entry into plaque is not enough!
• Proposal: develop new therapy that suppresses proliferation of macrophages in plaque

"It's a bit like killing off the army to be sure there's no collateral damage when invaders attack your country”

--David Diamond, Member of THINCs

*C.S. Robbins et al., Nature Medicine, Aug. 2013 [ePub ahead of print]
Elevated Homocysteine and Heart Disease*

- 587 patients with coronary artery disease followed over median period of 4.6 years
- Homocysteine > 15 micromol/Liter → 6.5-fold increase in death rate compared to homocysteine < 10 micromol/L

Pathway from Homocysteine to Sulfate*

When folate and B12 are deficient, nearly all homocysteine is converted to homocysteine thiolactone

Platelets and Cholesterol Sulfate

• Platelets and RBCs both synthesize cholesterol sulfate (Ch-S)
  – Ch-S is present in the atherosclerotic lesions in the aorta
  – Platelets will accept cholesterol only from HDL-A1
  – Platelet synthesis rate increases 300-fold when PAPS is available.
  – PAPS is formed from ATP and sulfate

• Platelet aggregation leads to thrombosis
  – HDL suppresses aggregation; LDL promotes it

* Yanai et al, Circulation 109, 92-96, 2004
Steps in Atherosclerosis: Reinterpreted

1. Endothelial cells lining artery walls feeding the heart release inflammatory agents
2. Macrophages infiltrate artery wall
3. Macrophages extract cholesterol from oxidized LDL and deliver it to HDL-A1.
4. Platelets extract cholesterol from HLD-A1 and convert it to cholesterol sulfate, with help from PAPS
5. Macrophages die and build up necrotic core
Putting it All Together

- homocysteine
- sulfate
- ATP
- PAPS
- Platelet
- Cholesterol Sulfate
- HDL-A1
- oxLDL
- LDL
- ApoE
- Ch
- Ch
- Lipid Raft
- Heart muscle cell
- Artery Wall
- Endothelial Cell
- Inflammatory Agents
- ROS
- insulin
- glucose
- Synthesize sulfate
- (3)
- Red Blood Cell
- Platelet
- Macrophage
- RBC
Putting it All Together

Cholesterol protects the heart from ion leaks and sulfate allows it to safely metabolize glucose
Statins Increase Plaque Calcification*

- 6673 people studied (2413 on statins 4260 not on statins)
- Mean age 59, 55% male
- Evaluated using coronary CT angiography - noninvasive method to visualize atherosclerotic features
- Statin use was associated with increased prevalence of calcified plaque and increasing numbers of coronary segments possessing calcified plaque

I hypothesize that treatments aimed at reducing the supply of cholesterol to the plaque will eventually lead to severe deficiencies in cholesterol and sulfate supply to the heart, resulting in heart failure.
Figure 5
Prevalence of CHF, by Age, 1976-80 and 1988-91

Recapitulation

• Kidney disease is associated with inversion of factors for heart disease risk
  – Kidneys depend critically on sulfates for function
  – Inflammation is required to synthesize sulfate
• Cardiovascular disease can be best characterized as a factory to supply cholesterol and sulfate to the heart and kidney
• Statin drugs, through their ability to deplete the supply of cholesterol to the plaque, can lead to heart failure and kidney failure down the road
How Statins Really Work Explains Why They Don’t Really Work
Sugar: The Bitter Truth*

• Robert Lustig, UCSF Professor of Pediatrics, has done more than anyone to promote the idea that fructose is damaging to your health
• In his 90-minute lecture, he discusses the critical role that the liver plays in detoxifying fructose
• Nearly 4 million views from YouTube

Liver Detoxifies Fructose

- Dietary fructose is toxic -- a severe glycating agent
- Liver takes up fructose and turns it into fat
- Fat is exported to tissues packaged in LDL particles
- Cholesterol is required to “wrap” synthesized fats
A Proposal

• Statins prevent liver from detoxifying fructose
• Muscles (type-II fibers) aggressively convert dietary sugars to lactate instead *
• Muscles experience severe glycation damage
• Cells die and release their contents as debris
• Insufficient clean-up by macrophages leads to collateral damage
• Fructose and glucose build up and cause widespread glycation damage
• Cascade effect

How Statins Change Fructose Metabolism

• Liver can’t come up with enough cholesterol to “wrap” synthesized fats with LDL
• Muscle cells must dispose of sugars via anaerobic metabolism (convert to lactate)
• Heart and liver benefit from a healthy fuel source
But, ...The Muscle Cells Get Wrecked*

- Patients on statins routinely get creatine phosphokinase levels monitored for possible muscle damage
- However, even when creatine phosphokinase levels are not elevated, myopathy (muscle weakness) can occur, and is often associated with muscle damage
- Patients on statins are often advised to continue statin therapy in this circumstance
  - This is bad advice

* Mohaupt et al., CMAJ 181(1-2), E11-E18, Jul 2009
Mitochondrial Disruption by Statins

Cell is Forced to Switch to Anaerobic Metabolism

Slide adapted from Statins and Miofibrillar Response, R. Laforge
Widespread glycation and ROS damage in muscle cells kills them

BUT

They don’t die properly!
Cytotoxic T-cells initiate apoptosis cascade and clean up cell debris following programmed cell death mediated by Fas-FasL signaling*

Cell Apoptosis

- Both Fas and FasL must migrate from the cytoplasm to the membrane to be activated
- This requires isoprenylation by geranylgeranyl pyrophosphate
- Statins suppress these mechanisms
  - Cell dies through necrosis instead

Statins make you grow old faster

Heart failure

Kidney failure

Diabetes

Liver failure

Glycation Damage

AGES

Fructose

Cataracts

Memory problems
Statins, Diabetes, and Cataracts*

- Large study: 6397 patients
- Divided into four groups:  
  ± diabetes, ± statins
- Looked at 50% point for age of diagnosis of cataracts
- Taking a statin gives you cataracts three years earlier

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<td>With diabetes</td>
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Statin's Promote Muscle Fiber Atrophy

- Statins induce synthesis of atrogin-1 in muscles, inducing skeletal muscle atrophy.
- In controlled experiments, Lovastatin promoted muscle fiber damage in zebrafish embryos.

Myopathy, Weakness and Chondroitin Sulfate*

- Chondroitin sulfate deficiency in muscles leads to severe limb and respiratory weakness
- Prednisone (a synthetic corticosteroid) produced a marked increase in strength

Hypothesis: prednisone sulfate supplies sulfate to synthesize chondroitin sulfate

Heparin prevents Oxidative Damage from Fenton Reaction*

• Superoxide ($O_2^-$) is a key ROS in the mitochondria
  – Iron oxidation (Fenton reaction) induces superoxide
  – $4Fe(II) + O_2 + 4H^+ \rightarrow 4Fe(III)(OH)_2^+ + 2H_2O$

• Heparin (many sulfates) induces acidic environment that changes the reaction as follows:
  – $4Fe(II) + O_2 + 4H^+ \rightarrow 4Fe(III) + 2H_2O$
  – This produces water (harmless) rather than superoxide (destructive)

“Chronic and age-related increases in ROS production will cause mitochondrial damage and dysfunction that perpetuate a catastrophic cycle of cellular injury, further ROS generation, mitochondrial decline, and cell death.”*  

* S.W. Ballinger et al., Circ Res. 2000;86. P. 965
Lactate and Heart Failure*

• Plasma lactate levels at rest are a marker for oxidative capacity
• Study involving 10,000 participants
  – Resting lactate levels measured
  – Monitored for ten years subsequently
• Lactate levels grouped into quartiles
  – Highest quartile had significant increased incidence of heart failure and death

Recapitulation

• Liver normally detoxifies fructose by converting it to fat → LDL
  – Statins impair this process
• Muscle cells respond by converting fructose to lactate
  – Supplies alternative fuel to liver, heart, and brain
  – But muscles suffer glycation damage as a consequence
• Statins interfere with distressed muscle cell's ability to signal distress
  – Cell dies an unnatural rather than a programmed cell death
  – Fructose accumulates and causes widespread glycation
• This has widespread consequences, including heart failure
• Cataracts, a sign of AGEing, happen sooner on statins
Maintaining Blood Stability
The Glycocalyx

• Negatively charged gel-like mesh lining the walls of all arteries, veins and capillaries
• Depends crucially on sulfated polysaccharides
  – Particularly heparan sulfate proteoglycans (HSPGs)
• Sulfate creates “exclusion zones”
  – Helps protect cells in wall from ion leaks and contact with enzymes suspended in blood
  – Greatly reduces amount of flowing blood (decreases resistance, lowers blood pressure)
  – Stores energy in “battery” constructed from structured water
"The Fourth Phase of Water" = the Exclusion Zone (EZ)
  – Profoundly excludes things, even small molecules
  – \( H_3O_2 \) instead of \( H_2O \) -- negatively charged!
  – Fills most of your cells and the extracellular tissues
  – "polywater" - water as a polymer

Many experiments in Pollack’s laboratory confirm the existence of EZ water
  – A hydrophilic surface (sulfate!) will promote growth of EZ water upon exposure to infrared light
  – EZ water absorbs UV light (270 nm) and this builds charge!

Water flows in small diameter tubes and the flow rate increases upon exposure to sunlight
  – Could the same thing be happening in your capillaries?

Pressure, oxygen, light, and infrared energy all build EZ water

*http://articles.mercola.com/sites/articles/archive/2013/08/18/exclusion-zone-water.aspx*
“And so it is with the exclusion zone. Order cannot persist without a continual supply of energy. The separated charges will slowly recombine, and order will give way to disorder. The exclusion zone’s outer reaches will wear thin like an eroding beach.”

– page 95, The Fourth Phase of Water

Sunlight renews the exclusion zones throughout your body!
Therapeutic Value of Sauna Therapy (Infrared light)*

• Impaired vascular function associated with hypertension, hyperlipidemia, diabetes, obesity
• Sauna therapy ameliorates endothelial dysfunction and improves health
  – Increased eNOS synthesis in aorta in animal experiments
  – Decreased body weight and body fat in obese subjects
  – Improved cardiac function in patients with heart failure
• Infrared expands exclusion zones 4-fold**

** B. Chai et al., J Phys Chem B. 2009 October 22; 113(42): 13953–13958.
Effect of Radiant Energy on Near-Surface Water *

“The main question, however, is where the energy for building these charged, low-entropy zones might come from. It is shown that radiant energy profoundly expands these zones in a reversible, wavelength-dependent manner. It appears that incident radiant energy may be stored in the water as entropy loss and charge separation.”

*B. Chai et al., J Phys Chem B. 2009 October 22; 113(42): 13953–13958.
Key Role for Sulfates

• Sulfates decorate exterior of cell, e.g., as heparan sulfate proteoglycans (HSPGs) or as cholesterol sulfate (Ch-S)
• Sulfates carry negative charge and keep suspended cells from sticking together
• Sulfates, as kosmotropes, create exclusion zone – gel-like environment to protect cell
Zeta Potential

• Zeta potential indicates degree of repulsion between similarly charged particles in a dispersion (e.g., the blood)

• High zeta potential confers stability
  – RBCs and platelets resist aggregation

• Low zeta potential causes flocculation and coagulation (blood clot)

• Proposal: a steady drop in zeta potential in the blood as we age is the source of many modern diseases
Grounding the Human Body Reduces Blood Viscosity*

• Blood viscosity is strongly influenced by the surface charge on the suspended particles (like RBCs)
• A higher repulsive surface charge increases spacing between RBCs, reduces clumping, lowers viscosity, and lowers peripheral resistance to flow (blood pressure)
• Grounding transfers electrons from soil to the body
  – Increases zeta potential, lowers blood viscosity

*G. Chevalier et al., J Alternative and Complementary Medicine, 1–9, 2012
Hypothesis

• When blood becomes deficient in negative charge (sulfates), glycocalyx becomes unhealthy
• Endothelial cells develop gaps that allow blood to seep into tissues
• Blood clots are needed to plug the holes
• Mistakes can lead to dangerous blockage
  – Deep vein thrombosis
  – Pulmonary thrombosis
Strokes are Happening at Younger Age*

- Incidence of stroke is increasing among people under 55 years old
- 959 patients followed through 2012 (aged 18 to 50 years at time of stroke)
  - Suffered from stroke between 1980 and 2010.
  - Compared with age-matched young adults without stroke
- Stroke victims have higher risk of dying prematurely
  - 3.4-fold higher risk in the 40- to 50-year-old age group
  - 6.4 fold higher risk in the 18-29 year olds
- Underlying vascular disease remains active throughout life?

"High Cholesterol Levels Are Associated with Improved Long-term Survival after Acute Ischemic Stroke"*

• 190 stroke patients studied
• Survival rates (P= 0.0001):
  
<table>
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<tr>
<th></th>
<th>3-mo</th>
<th>1-yr</th>
<th>5-yr</th>
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<td></td>
<td>100%</td>
<td>98%</td>
<td>84%</td>
<td>high-cholesterol group</td>
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<tr>
<td></td>
<td>92%</td>
<td>87%</td>
<td>57%</td>
<td>low-cholesterol group</td>
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</tr>
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DHEA Provides Sulfate?*

• In aging rats:
  – eNOS expression is lowered
  – Response of vascular smooth muscle to nitric oxide is decreased
  – Weakened resistance to oxidative damage

• DHEA supplement ameliorates these effects and delays aging process

Recapitulation

• Sulfates in the glycocalyx maintain proper function of the artery wall and smooth flow of the blood
• Gerald Pollack has revolutionized our understanding of water as a miracle molecule
• Infrared light induces growth of exclusion zone
• Grounding can improve blood stability by providing negative charge through currents from the ground
• High cholesterol leads to longer survival following stroke
• DHEA sulfate can improve aging vessels
Cancer to the Rescue?
“Cancer as a Metabolic Disease”*

• Turns cancer theory upside down!
• Impaired mitochondrial function leads to switch to aerobic glycolysis
  – Cancer cells convert glucose to lactate, producing a tiny amount of energy
  – Consume lots of glucose
• Metabolic dysfunction *precedes* genetic mutations

*T.N. Seyfried and L.M. Shelton, Nutrition and and Metabolism 7:7, 2010
“Cancer as a Metabolic Disease”*

- Turns cancer theory upside down!
- Impaired mitochondrial function leads to switching to aerobic glycolysis.
  
  Tumor is a factory where damaging (glycating) sugars are converted into healthy fuel to supply to heart and brain.

- Metabolic dysfunction precedes genetic mutations

*T.N. Seyfried and L.M. Shelton, Nutrition and Metabolism 7:7, 2010
Breast Cancer Overtreatment* 

- 200,000 Norwegian women followed consecutive 6-year periods 
- Half received regular breast exams and mammograms, while the other half did not 
- Those receiving monitoring developed 22% more incidence of breast cancer 
- Researchers concluded that the control group had probably had as many cancers as the “treatment” group, but these cancers had resolved on their own! 
- Another possibility is that the radiation in the mammogram induced breast cancer

“Heparan Sulfate and Heparanase as Modulators of Breast Cancer Progression”*

• Heparanase is a potent tumor modulator due to its protumorigenic, proangiogenic, and prometastatic activities
  – Plays a role in bone metastasis
• Heparanase breaks down heparin sulfate into fragments of 4 to 7 kDa size
• Heparinase is upregulated in all human sarcomas and carcinomas
• Syndecan-1 shedding is an important step in tumor invasion and metastasis

*A.M. Gomes et al., BioMed Research International, 2013, article ID: 852093
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Is the tumor providing heparan sulfate to the blood?

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Syndecan Shedding in Breast Cancer

Figure 1 from Weinbaum et al., Annu. Rev. Biomed. Eng. 2007. 9:121–67
Hydrolysis of Estrone Sulfate and Dehydroepiandrosterone (DHEA) Sulfate by MCF-7 Human Breast Cancer Cells*

Breast cancer tumor cells break down estrone sulfate and DHEA sulfate in order to supply sulfate to the blood stream??

*J.H. MacIndoe et al., Endocrinology Sep 1, 1988, 123(3), 1281-1287.
Curing Cancer Brings on Health Problems*

“The rise in cancer survival in Britain is a ‘double-edged sword’ a charity has warned, because the NHS is ‘woefully unprepared’ to cope with the number of patients who go on to suffer long-term health problems.”

“... one in four survivors is left suffering long-term debilitating health conditions as a result of the disease, or the treatment for it.”

Curing Cancer Brings on Health Problems, Cont’d*

• Many cancer survivors suffer extreme pain up to five years after diagnosis
• Women diagnosed with breast cancer are almost twice as likely to get heart failure
• Men diagnosed with prostate cancer are more than twice as likely to get osteoporosis
• Many treated for prostate cancer or bowel cancer suffer from incontinence

Chemotherapy for Breast Cancer*

- Chemotherapy induces anemia
- EPO (erythropoietin) sometimes given to induce RBC regeneration from bone marrow
  - This can lead to thrombosis (blood clots)
- Alternative is blood transfusions to restore hemoglobin count

These problems suggest that breast cancer is associated with unstable blood

*http://www.medpagetoday.com/HematologyOncology/Anemia/40536
Cognitive Decline and Breast Cancer*

"Patients treated with *hormonal* therapy were five times more likely to exhibit cognitive decline as compared with a control group. Chemotherapy was not a significant predictor of cognitive decline."

*http://www.medpagetoday.com/MeetingCoverage/MBCS/41486
Statins and Breast Cancer*

- Population-based case-control study of breast cancer
- 1,068 cases compared to 902 controls
- Current users of statins for 10 years or longer
  - 1.83-fold increased risk of invasive ductal carcinoma
    (accounts for 7 out of 10 cases in UK)
  - 1.97-fold increased risk of invasive lobular carcinoma
    (10 to 15% of cases)

*I hypothesize that this is because statins deplete cholesterol sulfate supply and disrupt sugar metabolism

*J.A. McDougall et al., Cancer Epidemiol Biomarkers Prev. 2013 Jul 5. (Epub ahead of print)
Cholesterol Sulfate in Prostate Cancer*

“Remarkably, cholesterol sulfate ... was observed exclusively in the cancerous tissue, enabling its clear identification”

"Hormone Use in Prostate Cancer May Harm Kidneys"*

• Analysis included 10,250 men, who were followed for an average of 4.1 years

• Androgen deprivation therapy (ADT) led to significantly increased risk of acute kidney injury in men being treated for prostate cancer
  – Hyperglycemia is a well established adverse effect of ADT, and this stresses the kidney

*Is this due to insufficient sulfate supply to the body??

Statins and Prostate Cancer*

• 388 prostate cancer cases compared to 1,552 controls
• Ever-use of any statin was associated with a significant 55% increased risk to prostate cancer
  – Highest cumulative dose group had 86% increased risk

*CC Chang et al., Prostate. 2011 Dec;71(16):1818-24
Iclusig: A Drug that’s Failing*

• Chemotherapy for leukemia
• “Fatal and life-threatening heart attack, stroke, loss of blood flow to the extremities resulting in tissue death, and severe narrowing of blood vessels in the extremities, heart, and brain requiring urgent surgical procedures to restore blood flow”

*http://www.medpagetoday.com/HematologyOncology/Leukemia/42637
Cancer and Deep Vein Thrombosis*

• Deep vein thrombosis is the second-leading cause of death among cancer patients (after infection)

• Chemotherapy increases risk of thrombosis 6.5x

Hypothesis: Removing the tumor removes the protective effect

Colon Cancer Tumor Cells Synthesize Hydrogen Sulfide*

• Hydrogen sulfide induces cell growth in tumor cells by promoting glycolysis and oxidative phosphorylation (breaking down glucose to produce ATP).
• Hydrogen sulfide is produced by the enzyme Cystathionine-β-synthase (CBS) in these cells – It is upregulated in these cells and promotes their growth
• Suppression of hydrogen sulfide synthesis is being proposed as a technique to kill the tumor


The tumor is able to synthesize sulfate from the hydrogen sulfide gas it produces??
William Li: Can we eat to starve cancer?

• You can starve a tumor (prevent blood vessel growth) by eating certain foods that inhibit angiogenesis (blood vessel growth)

• Natural inhibitors of angiogenesis:
  – Resveratrol (red grapes, wine); ellagic acid (strawberries)
  – Turmeric, garlic, glusamine (sulfate)
  – Olive oil, tea, parsley, etc. (flavonoids)

• Also protect from obesity

All of these foods either contain sulfate transporters or contain sulfur or both

*TED Talk: https://www.youtube.com/watch?v=B9bDZ5-zPtY
Recapitulation

• Cancer is a metabolic disease
  – Cancers cells consume massive amounts of glucose and produce massive amounts of lactate
• Treating cancer can lead to deteriorating health
  – Heart failure, extreme pain, osteoporosis, mental decline, anemia, deep vein thrombosis
• Breast cancer and prostate cancer supply sulfated sterols to the body
  – These are essential to maintain healthy blood
• Foods that promote sulfate synthesis and sulfate transport can cause tumor to shrink
Summary

• Sunlight-catalyzed cholesterol sulfate synthesis in the skin is essential for long-term health
  – Heart disease is a compensatory mechanism to synthesize cholesterol sulfate
• Heparan sulfate proteoglycans (HSPGs) are pervasive in the body, and sulfate depletion leads to diabetes, heart disease, and kidney disease
• Statin drugs work by destroying skeletal muscles to provide lactate for the heart
• Hemostasis depends on adequate sulfate in the blood
• Cancer may develop as a strategy to both reduce blood sugar levels and produce sulfated sterols
What's Next?

Early afternoon (1:30 p.m. – 3 p.m.)
1. Gut Microbes: How They Help Us Out
2. Autism, Depression, and Alzheimer's Disease

Late afternoon (3:30 p.m. – 5 p.m.)
1. Glyphosate: The Elephant in the Room