

## The Detection and Reduction of Mercury in Humans Using Oral DMPS (2,3-dimercaptopropane-1-sulfonate) with Support Nutritional

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### Abstract

**Background:** *Micromercurialism* is a term designating chronic low-level mercury toxicity, the clinical testing, diagnosis and treatment of which has been essentially non-existent in our health care system<sup>1</sup>. This is likely due to the vague and non-specific nature of the presenting signs and symptoms [Table 1]. Dental amalgam is the greatest source of mercury in the general population<sup>2</sup>, and is released in vapour form, which is detectable, and quantifiable<sup>3, 4a, 4b</sup>. Tissue-bound mercury levels are not adequately measured by standard blood and urine analysis and there are many inconsistencies and false negatives associated with hair analysis<sup>5</sup>. The determination of an elevated body burden of mercury can best be made by measuring urinary mercury, after provocation with the chelating agent **2,3-dimercaptopropane sulfonate (DMPS)**, a simple, inexpensive procedure that provides objective confirmatory evidence<sup>6</sup>. Attempts both to detect and detoxify individuals of mercury have been hampered by the use of intravenous DMPS chelation and lack of clinician's awareness of proper testing techniques.

**Objective:** To determine the ability of an oral DMPS preparation to provide both accurate detection and removal of tissue-bound mercury by measuring pre- and post-treatment urine mercury by DMPS provocation analysis.

**Methods:** From January 1, 2005 through April 30, 2006 a cohort of 147 people distributed between medical clinics in New Zealand and the United States subsequently performed acceptable follow-up testing and was included in this study. The participants were selected based on the documentation of tissue burden of mercury obtained by an oral DMPS provocation 2-hour urine collection and analysis. Collection protocols required no seafood for 3 days preceding the collection. This baseline quantitative analysis was then compared against a repeat provoked urine following either one or two months of treatment using oral DMPS and a support formula (**MercOut**·www.MercOut.com), each taken once daily. Dosing of DMPS was initially 100 mg given at bedtime and progressed to a maximum 200 mg DMPS over the 30-day detoxification period.

**Results:** 63% of the cohort (93 of 147) showed a reduction of measured urine mercury levels greater than 50% within 30 to 60 days of using the oral DMPS formulation. The average mercury reduction in this group was a highly significant 69% [Chart 1].

**Conclusion:** 93 of the 147 cohort (63%) fell into either **Groups C or D, showing a very significant reduction from between 51% to 100% in their repeat urine Hg levels as compared to initial baseline.** The average mercury urine reduction in these two groups was 69%. The results show that the use of a formulation containing oral DMPS significantly reduced the tissue concentrations of mercury in humans within 60 days.

### References

- <sup>1</sup> Ely JT, A, Fundenberg HH, Muirhead RJ, LaMArche MG, Krone CA, Buscher D; Urine Mercury in micromercurialism: bimodal distribution and diagnostic implications. *Bull. Environ. Toxicol.* **63** (1999) 553-9
- <sup>2</sup> World Health Organisation: Elemental Mercury and Inorganic Mercury: Human Health Perspectives (1993) 11 (Contains 10 pages of references)
- <sup>3</sup> Ely J.T., Risk Factor for Parenteral Intoxication by Mercury from Dental Amalgam. *Bull. Environ. Toxicol.* **67** (2001), 309-316
- <sup>4a</sup> Weiner JA, Nylander M (1995) An estimation of the uptake of mercury from amalgam fillings based on urinary excretion of mercury in Swedish subjects. *The Science of the Total Environment*, 168:255–265.
- <sup>4b</sup> Barregard L, Sallsten G, Jarvholm B (1995) People with high mercury uptake from their own dental amalgam fillings. *Occupational and Environmental Medicine*, 52:124–128.

- <sup>4c</sup> Aposhian, H.V., D.C. Bruce, W. Alter, R.C. Dart, K.M. Hurlbut, M.M. Aposhian, Urinary Mercury after Administration of DMPS: Correlation with Dental Amalgam Score *FASEB J.* **6**: (19922472-2476;.)
- <sup>5</sup> G. Drasch, E. Wanghofer and G. Roeder, Are blood, urine, hair and muscle valid biomarkers for the internal burden of men with the heavy metals mercury, lead and cadmium? *Trace Elements Electrolytes* **14** (1997), 116–123
- <sup>6</sup> H.V. Aposhian, R.M. Maiorino, D. Gonzales-Ramirez, M. Zuniga-Charles, Z. Xu, et al., Mobilization of heavy metals by newer, therapeutically useful chelating agents, *Toxicology* **97** (1995), 23–38.

## Chart 1 -- Patient Outcomes

The following data shows baseline mercury levels taken prior to 30- to 60-day DMPS/nutritional supplement program ("Pre-Hg") and after the program was completed and a repeat urine challenge was performed ("Post-Hg"), with the % Improvement indicated.

**Results Analysis:** 93 of the 147 total included cohort (63%) fell into either Groups C or D, showing a very significant reduction of between **51% to 100%** in their repeat urine mercury (Hg) levels as compared to baseline. The average mercury urine reduction in these two groups was **69%**.

### Group A: 0-25% Improvement (25 people)

Name	Pre-Hg	Post-Hg	% Improve	Name	Pre-Hg	Post-Hg	% Improve
Alberto	5.9	5.9	0%	Jayne	54	43	20%
Dennis	16	16	0%	Murray	54	43	20%
Melissa	23	23	0%	Maria	15	12	20%
Carol	28	27	4%	John	64	50	22%
Diana	31	29	6%	Robert	8.1	6.3	22%
Randy	92.5	85	8%	Shushila	22	17	23%
Donald	13	12	8%	Glenys	56	43	23%
Eric	23	20	13%	Jo	49	38	23%
Dawn	73	69	15%	Suzanne	40	31	23%
Alyse	9.8	8.3	15%	Becky	60	46	23%
Bob	13	11	15%	Tom	25	19	24%
Pat	29	24	17%	Janet	16	12	25%
Angelika	10	8.1	19%				

### Group B: 26 to 50% Improvement (29 people)

Name	Pre-Hg	Post-Hg	% Improve	Name	Pre-Hg	Post-Hg	% Improve
Marion	29	21	28%	George	8.4	4.9	42%
Rosemarie	45	31	31%	Stuart	28	16	43%
Steven	29	20	31%	Adolf	14	8	43%
Ladene	7.4	5	32%	Ted	32	18	44%
Chris	46	31	33%	Hectorine	25	14	44%
Justine	33	22	33%	Laurie	25	14	44%
Janis	15	9.8	35%	Vivian	110	60	45%
Rick	6.2	4	35%	Michael	20	11	45%
Michael	42	27	36%	Bill	7.4	4.1	45%
Macy	19	12	37%	T.	26	14	46%
Tom	24	15	38%	Robert	18	9.5	47%
Christine	49	30	39%	Louise	10	5.1	49%
Mary-Lyn	28	11	39%	Steve	66	33	50%
Sandra	110	66	40%	Bob	34	17	50%
Ada	7.4	4.4	41%				

**Chart 1 -- Patient Outcomes (cont'd)**

**Group C: 51 to 75% Improvement (65 people)**

Name	Pre-Hg	Post Hg	% Improve	Name	Pre-Hg	Post-Hg	% Improve
Liz	5.7	2.8	51%	Marcus	30	11	63%
Diane	70	34	51%	MaryLou	65	24	63%
Shirley	27	13	52%	Fraser	47	17	64%
Janet	42	20	52%	Karen	15	5.4	64%
Alister	25	12	52%	Larry	76	27	64%
Erica	4.5	2.1	53%	Leas	69	25	64%
Peggy	70	33	53%	Adela	14	5.1	64%
David	5.5	2.6	53%	Graeme	34	12	65%
Geoffrey	35	16	54%	Maria	28	9.8	65%
Larry	6.8	3.1	54%	Lee	32	11	66%
Verna	65	30	54%	Gaylene	18	6.1	66%
Bronwynne	60	27	55%	Ann	140	47	66%
Ursula	20	9	55%	Dianne	17	5.7	66%
Derry	42	19	55%	James	27	9.3	66%
Walter	33	15	55%	Reowtie	170	57	66%
Jack	210	95	55%	Sude	51	17	67%
Charise	85	37	56%	Michelle	72	24	67%
Louisa	25	11	56%	Susan	69	23	67%
Phyllis	11	4.7	57%	Karen	20	6.4	68%
Harry	22	9.5	57%	Linton	15	4.8	68%
Mark	140	58	59%	Dianne	80	25	69%
Mary	32	13	59%	Ann	83	24	71%
John	57	23	60%	Maufred	28	8.2	71%
Vikki	36	14	61%	Bronwyn	72	20	72%
Vince	44	17	61%	Bill	7.4	2.1	72%
Tom	11	4.3	61%	Ron	18	4.8	73%
Gary	83	32	61%	Jose	13	3.5	73%
Natalie	37	14	62%	Mezare	92	25	73%
Scott	17	6.4	62%	Antoine	8.9	2.4	73%
David	45	17	62%	Marc	92	24	74%
Randy	20	7.7	62%	Betty	25	6.3	75%
Gary	24	8.9	63%	Wes	91	23	75%
Elly	63	23	63%				

**Group D: 76% to 100% Improvement (28 people)**

Name	Pre-Hg	Post-Hg	% Improve	Name	Pre-Hg	Post-Hg	% Improve
Harold	27	6.5	76%	Francoise	52	8.2	84%
Jay	49	11	78%	Mary	37	5.2	86%
Howard	35	7.7	78%	Rosalie	41	4.5	86%
Otto	58	12	79%	George	27	3.8	86%
Beth	23	4.9	79%	James	44	5.8	87%
Tony	27	5.4	80%	Diane	7.3	0.9	88%
Philip	12	2.4	80%	Kathy	150	18	88%
John	20	3.9	81%	Carol	52	4.9	91%
Marty	92	17	82%	N.	15	1.4	91%
Joe	340	55	83%	Anjal	23	1.8	92%
Nancy	66	11	83%	Steven	90	2.4	97%
Paul	21	3.3	84%	Gary	38	1.2	97%
Landon	22	3.6	84%	Carol	49	6.5	99%
Jim	26	4.1	84%	A.J.	21	0	100%

**Table 1: Symptoms and Pathology of Mercury in the Human Body**

**Physical**

Autism  
 Amyotrophic lateral sclerosis  
 Ankylosing spondylitis  
 Myasthenia gravis  
 Paresthesias and neuralgias  
 Vision, taste, smell & hearing disturbances  
 Vertigo & tinnitus  
 Multiple Sclerosis  
 Parkinson's disease  
 Alzheimer's Dementia  
 Other dementias  
 Hypothyroidism/Cold Extremities  
 Infertility  
 Poor libido  
 Impotency

**Physical** (cont'd)

Other endocrine problems  
 Rheumatoid arthritis  
 Juvenile arthritis  
 Arrhythmias  
 Myocardial infarction  
 Lupus erythromatosus  
 Other autoimmune diseases  
 Multiple chemical sensitivities  
 Chronic fatigue  
 Fibromyalgia  
 Sciatica  
 Gastritis & Colitis  
 Irritable bowel syndrome  
 Crohn's disease  
 Sleep disorders  
 Yeast syndrome

**Psychological/Mental:**

"Brain fog" or poor focus/concentration  
 Rage or being quick to anger  
 Mood swings  
 Indecisiveness  
 Panic attacks  
 Attention Deficit Disorder (ADHD)  
 Hyperactivity  
 Learning disabilities  
 Depression  
 Unexplainable sadness  
 Joylessness  
 Fearfulness  
 Obsessive-compulsive disorder  
 Manic-depressive disorder  
 Sleep disturbances  
 Anorexia nervosa  
 Bulimia

**Oral Formula Used in Research**

<b>Dosage: 3 Morning capsules and 1 Nighttime capsule</b>	
<b>Amount per 3 Morning capsules</b>	
Vitamin C (ascorbic acid)	360 mg
Calcium (from citrate)	50 mg
Magnesium (from aspartate)	12 mg
Zinc (from picolinate)	20 mg
Selenium (from l-selenomethionine)	200 mcg
Copper (from copper glycinate)	0.60 mg
Manganese (from manganese aspartate)	0.38 mg
Chromium (from polynicotinate)	25 mcg
Molybdenum (from molybdenum aspartate)	99 mcg
<b>MercOut Support Complex</b>	1426 mg
MSM (methylsulfonylmethane), NAC (N-Acetyl-Cysteine), Milk Thistle Extract (80% silymarin), Boron, Turmeric Root Extract (min. 95% curcuminoids), Alpha Lipoic Acid, L-Methionine, L-Glutamine, Glycine, L-Cysteine.	
<b>Amount per 1 Nighttime capsule</b>	
DMPS (dimercaptopropanesulfonic acid)	125 mg
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