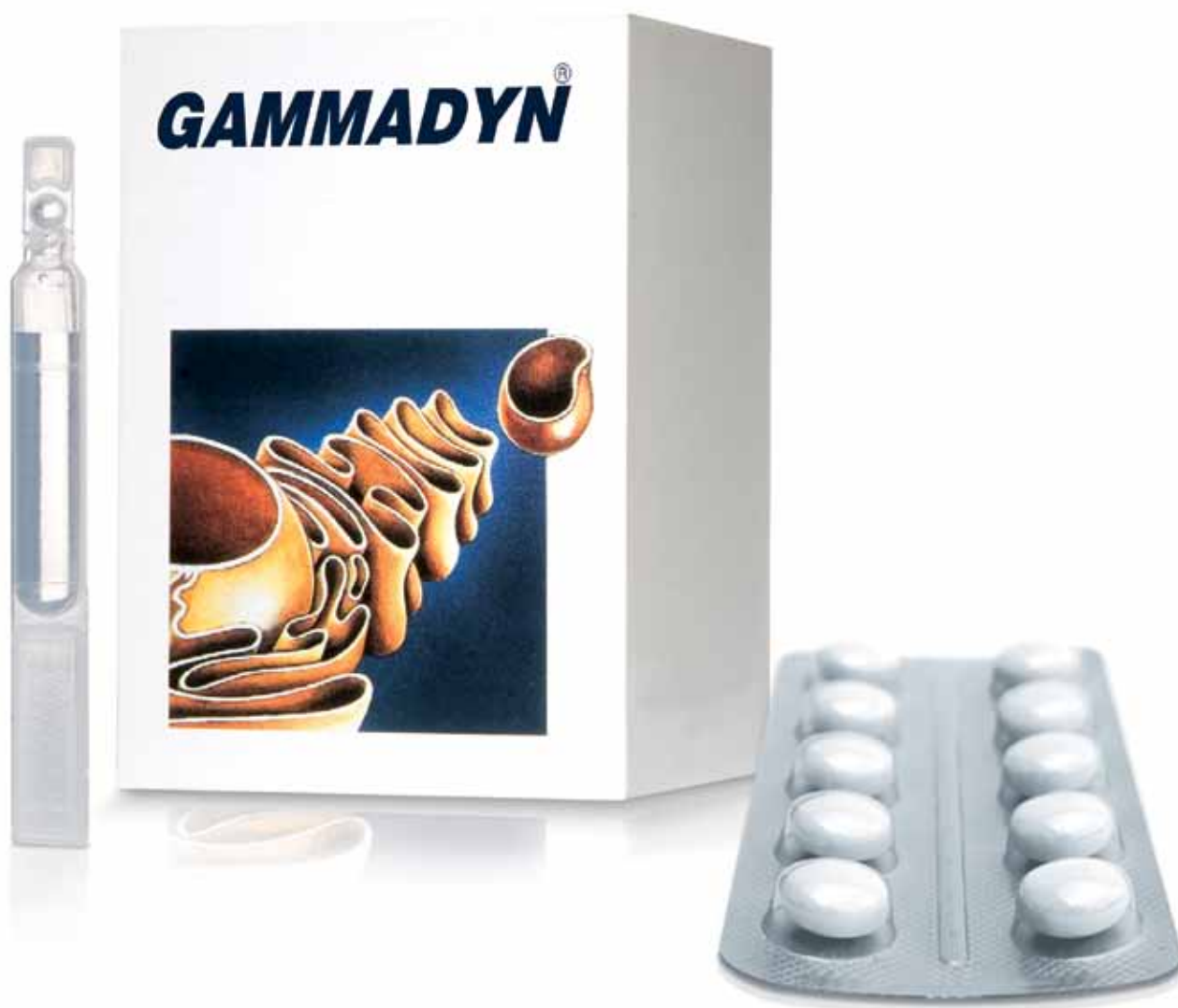


unda[®]

Gammadyn[®]

OLIGO ELEMENTS TO STIMULATE ENZYME
AND HORMONE FUNCTION



The aim of this comprehensive product reference guide is to assist in building more effective and successful patient treatment strategies. The reference guide is to be used as an adjunct to Seroyal Continuing Education and is not intended to be utilized as a diagnostic tool nor replace any other required education. The remedies indicated in this reference guide are potent medicines that can have profound therapeutic effects on patients. As with any intervention, the dosage may need to be adjusted for those with chronic conditions, very sensitive individuals and those taking multiple medications. Practitioners are solely responsible for the care and treatment provided to their own patients. The information provided by the speaker or speakers in the Seroyal Continuing Education program together with any written material provided do not necessarily represent the views of Seroyal and are not intended as medical advice or an endorsement of any products. This information is for professional use only and is not meant to diagnose, treat, cure, prevent any disease or replace traditional treatment, and has not been evaluated by the FDA or Health Canada.

Seroyal

The Function of Trace Minerals

Over a century ago, French chemist and biologist Gabriel Bertrand discovered that trace minerals (mineral salts) are naturally present within the body. These minerals play an extremely important biological role as key components in the control of cellular metabolism in homeostatic regulation. At the cellular level, trace minerals are used at minute intracellular dilutions, released in ionized form for immediate transportation and utilized throughout the body. Trace minerals function as:

Enzymatic co-factors. Enzymes are the biological catalysts required in most chemical reactions that occur in biological systems. Enzymes allow the body to perform metabolic functions at rates in excess of a million times greater than would be possible without them. Formed out of protein molecules, most enzymes depend on specific, low-concentration trace minerals to provide metal ions as either stabilizers or activators, as well as help determine the enzyme's specificity for a given substrate. Trace minerals become the co-factors to help normalize enzymatic function.

- Example:** As a stabilizer, the trace mineral metal ion combines with the protein molecule in a permanent structure. Cytochrome C, an enzyme involved in oxidation-reduction processes, contains iron as an integral part of its structure.
- Example:** As an activator, the protein forms a bond with the trace mineral-metal only during the reaction itself, so that the metal ion acts as an enzyme catalyst. Alkaline Phosphatase, an enzyme essential for bone formation, functions only in the presence of zinc ions.
- Example:** Trace minerals affect the molecular configuration of enzymes, and can also induce distortions in the substrate, rendering it more susceptible to enzyme modification. Carboxypeptidase, a proteolytic enzyme, contains a zinc stabilizer that alters peptide configuration allowing hydrolysis at the carboxy-terminal.

Hormone modulators: Trace minerals are essential in the production of hormones and vitamins in the body.

- Example:** iodine is essential in the production of T3 and T4
- Example:** cobalt is essential in the existence of Vitamin B12
- Example:** iron is essential in the composition of hemoglobin

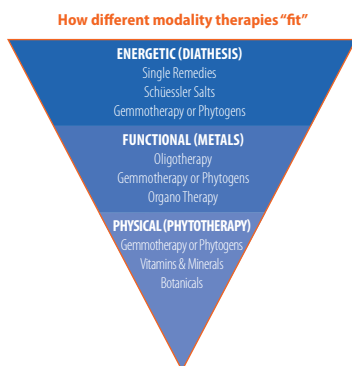
Structural components: Minerals possess either a quantitative or qualitative effect, depending on whether they work at the macro (quantitative) level of structural formation and functional processes, or at the micro (qualitative) level of cellular enzyme activation. Trace minerals work at a micro level (qualitative in action) but may have additional roles and work at a macro (or quantitative) level.

- Example:** fluoride aids in the normal growth of teeth and bone
- Example:** silica aids in the growth of the connective tissue

What are Oligo Elements?

"Oligo elements" or "oligotherapy" are trace minerals in a highly bioavailable form, administered in small doses and at low concentrations, providing ions the body cannot synthesize and which are

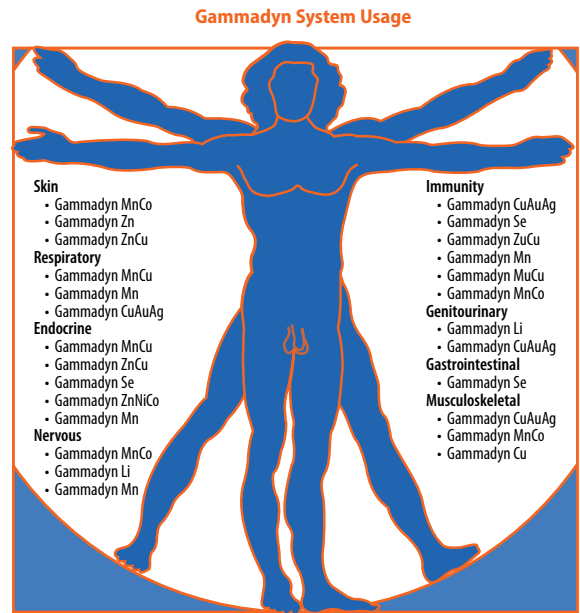
indispensable for cellular enzymatic functions. With a good safety profile, oligo elements have: no known side effects; no known drug to drug interaction; and will not cause any pharmaco-dependency. Oligo therapy can be well integrated with other modalities.



Oligo Elements and Physiological Disorders

Oligo elements are suited to treating functional pathologies and bringing the body back to homeostasis. It is a holistic approach that takes not just the physical but also the constitution of the individual into account. By regulating a particular constitution, practitioners are able to diminish or modulate reactivity in relation to a specific pathological predisposition.

Metabolic disorders involving Oligo elements can be divided into two main groups: disorders of overload and disorders of deficiency.



Disorders of Overload

Overload disorders are due primarily to accidental intoxication from adulterated foods and environmental pollutants, such as pesticides, heavy metals, or element disequilibrium.

Metal toxicity has become a major contributing factor in many metabolic disorders and functional diseases. For example, aluminum, a common contaminant in soft drink cans and some cookware, is known to induce neurofibrillary formations in the brains of higher animals and has been found in significant concentrations in the brains of individuals afflicted with Alzheimer's disease.

Toxic metals directly interfere with specific enzyme functions. Zinc-dependent enzymes are a prime example. In the presence of even low levels of lead, a close neighbor of zinc in the periodic table, these enzymes can bind to lead and thus be inactivated. Mercury and cadmium are also known toxic enzyme inhibitors. The toxic effect of these heavy metals stems from their ability to distort the substrate binding site of the enzyme rendering it useless. Targeted doses of oligo elements displace toxic inhibitors and reactivate enzyme functions.

An imbalance in the intracellular oligo element equilibrium can cause a build-up of one element at the expense of the other, thus leading to a qualitative insufficiency. This disequilibrium toxicity will respond to oligo therapy. For example, Alkaline Phosphatase is an enzyme activated by zinc, but the complex is highly changeable. Other divalent cations, such as magnesium or cobalt, if present in excessive levels, can bind to the enzyme and deactivate it. Large doses of zinc will only contribute to disequilibrium and are unable to reactivate the enzymes. Only direct administration of targeted doses of oligo elements will reactivate these enzymes.

As therapeutic enzymatic regulators, oligo elements aid in the treatment of biochemical dysfunctions without disturbing the existing equilibrium. Supplying these needed mineral co-factors at precise concentrations unblocks diseased metabolic pathways, allowing healing mechanisms to operate and prepare the patient to respond more readily and successfully to other modalities.

Disorders of Deficiency

Quantitative mineral deficiencies are the result of either a decrease of absorption or increased excretion. Absorption may be affected by alimentary supply. Deficient foods as a result of mineral depleted agricultural soils can deprive the body of essential metal cofactors. As a result, vegetarians have been determined at risk for mineral deficiencies in manganese, zinc, chromium, magnesium, copper and iron. Frequently, absorption is affected by decreased bioavailability of foods due to chelation with inassimilable compounds, such as phytates, phosphates, and antibiotics, or intestinal pathology (malabsorption syndromes), including dysbiosis of the microflora and other diseases.

Deficiencies may also be the result of increased excretion. For example, infants exposed to intrauterine alcohol have decreased plasma zinc levels and increased urinary excretion, resulting in decreased protein synthesis. There is a tight relationship between element equilibrium and metabolism. For example, zinc plays an important role in the synthesis of amino acids as a cofactor for Glutamate Dehydrogenase, magnesium is necessary for digestion as a primary cofactor for protein Peptidase, copper helps prevent free radical pathology as a part of Superoxide Dismutase, and manganese is essential for DNA synthesis and repair in DNA Polymerase. The scientific literature contains volumes of examples of minerals and their affect on the body's metabolism. The proper functioning of each of these processes is dependent on a highly integrated and balanced relationship of one mineral to another. As indicated above, a toxicity, deficiency or imbalance in any one mineral can be at the expense of another, resulting in metabolic disorder.

Oligo elements administered in the precise form and concentrations capable of rapid absorption quickly provide essential metals to reactivate cofactor-deprived or -inhibited enzymes. These reactivated enzymes are then capable of returning the cell to normal functioning capacity where they can then utilize other nutrients to promote normal metabolism and cellular homeostasis.

Used as supplementation, oligo elements are suited to treating functional pathologies stemming from trace mineral deficiencies caused by: nutritional imbalances, increased requirements; or insufficient absorption.

The Law of Arndt-Schultz

Conventional pharmacology dictates that a certain amount of an active substance is required in order for it to act upon receptors, enzymatic reactions or any other mechanism, and have an effect. This implies a linear dose-related effect relationship: the higher the dose, the stronger the action and the increased toxicity.

Oligo elements dosage is different. Its effectiveness is based more upon the Law of Arndt-Schultz: low doses stimulate, medium doses regulate and high doses depress. Concentrations higher than those required to maintain essential functions may have a secondary or pharmacological effect.

In oligotherapy, enzymatic kinetics follow specific physical laws of optimum concentrations that determine the formation of enzyme-cofactor and -substrate complexes, and the rate at which enzyme activity occurs. Enzymes are influenced by cellular environmental factors and function within a very narrow range of cofactor concentrations.

Benefits of unda® Gammadyn® in Oligotherapy

Manufacturing excellence and stringent testing make unda® Gammadyn® the market leader in oligotherapy products.

Convenient. Small travel-size ampoules for cleaner, easier, and more effective administration than sprays. Helps retain maximum potency and prevent contamination.

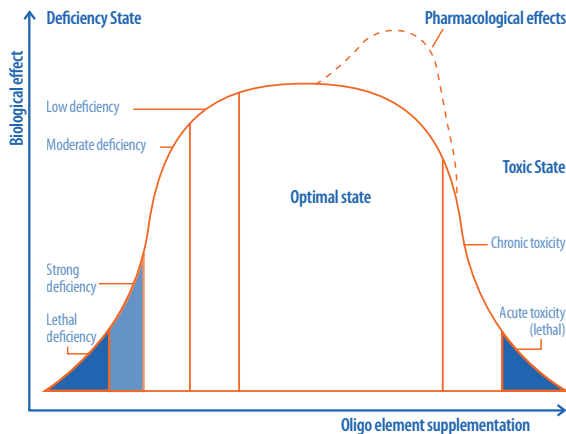
Quick Absorption. Liquid formulations keep active ingredients stable and highly bioavailable. Administered in solution for sublingual absorption, where sublingual mucosa, rich in surface capillaries, allow for the unimpaired absorption of the oligo elements into the circulation system and directly to the cell site in their intended concentrations.

Patient compliant. Superior tasting with no aftertaste.

Dose and Administration

Take one ampoule one to two times daily on an empty stomach or as recommended by your healthcare practitioner. Squeeze contents directly into mouth. Hold under tongue for about 20 seconds and swallow.

Arndt-Schultz law of pharmacology
Biological effects on the function
of oligo element supplementation



Gammadyn® Overview

COBALT (CO)



INGREDIENTS

Each 2 ml (0.07 fl.oz.) ampoule contains 0.059 mg of cobalt from cobalt gluconate.

MODE OF ACTION

Cobalt plays an essential role in the formation of vitamin B12 (cobalamin), and thus performs a number of physiological functions. Cobalt is involved in the regulation of the sympathetic and parasympathetic nervous system via the adrenergic receptors. Cobalt via B12 can help reduce spasms and increase blood flow. Cobalt stimulates erythropoietin secretion, facilitating the synthesis of erythrocytes. Cobalt via B12 can help prevent congenital anomalies and promote the healthy formation of mature sperm and ovum. Cobalt is also a cofactor for glycylglycine dipeptidase, an enzyme which hydrolyzes specific dipeptides.

INDICATIONS

GENERAL

Cardiorespiratory disorders: Peripheral circulatory spasms
Prediabetic condition due to secretal exhaustion of the Islets of Langerhans
Psychosomatic disorders: Migraine, stage fright
Spasmodic disorders

SPECIFIC

Anxious Agitation
Lumbago
Migraine
Parasite (intestinal)
Travel Sickness

KNOWN USES

Cobalt has been traditionally used with manganese in oligotherapy for the maintenance of healthy circulation; it has been used together with nickel for supporting the digestive system, and with zinc and nickel for supporting the endocrine system.

COPPER (CU)



INGREDIENTS

Each 2 ml (0.07 fl.oz.) ampoule contains 0.725 mg of copper from copper gluconate.

MODE OF ACTION

Copper forms the central element in a number of metalloenzymes which can help neutralize many toxins and high levels of certain biochemicals in the blood. Copper can stimulate the immune system and increases multiplication of B-lymphocytes and the synthesis of antibodies. Most copper is stored in the liver, heart, brain, spleen, kidneys and usually bound to proteins such as ceruloplasmin, cytochrome oxidase (liver detoxification) and superoxide dismutase (SOD). Copper is essential for decreasing acute and chronic bacterial and viral infections. Copper is also required in the production of various hormones (catecholamine).

INDICATIONS

GENERAL

Cardiorespiratory problems
Chronic and acute inflammation
Digestive disorders
Infections

SPECIFIC

Alcoholism (chronic Anemia)	Arthritis
Bronchitis	Candidiasis
Diarrhea	Hemorrhoids
Hepatic Detoxification	Influenza
Laryngitis (chronic)	Lumbago
Pancreatitis	Premenstrual syndrome (PMS)
Prostatitis (acute)	Warts

KNOWN USES

Copper has been traditionally used as an essential oligoelement and for states of infection and inflammation. In oligotherapy, copper has multiple, enzymatic functions, most notably the cytochrome C oxydase, which is responsible for the oxygenation in the respiratory chain and in the elaboration of elastin and collagen fibres. Copper is indicated for all infectious and viral states, such as flu and inflammatory rheumatism. Copper is a mineral that is a factor in the maintenance of good health, and also helps to produce red blood cells and connective tissue in humans (NHFD, 2004a).

COPPER-GOLD-SILVER (CU-AU-AG)



INGREDIENTS

Each tablet contains 0.063 mg of copper from copper gluconate, 0.0014 mg of gold from gold sodium thiomalate, and 0.021 mg silver from silver lactate. unda® Gammadyn Cu-Au-Ag is available in tablet form to ensure proper dispensing and product stability.

MODE OF ACTION

Cu-Au-Ag is indicated for arthritis and chronic rheumatism. Copper deficiencies affect the formation of collagen and its tensile strength. Copper affects the central nervous system and is associated with locomotor disorders. These oligo-elements potentiate adrenal function and can decrease fatigue associated with hypoadrenalism. Gold has been used successfully in chronic rheumatoid arthritis (American Rheumatism Assoc., 1970). This combination of oligo-elements is also important in immune function and helps activate the mechanisms of cellular and humoral immunity.

INDICATIONS

GENERAL

Antibiotic: Staphylococci, Streptococci
Cellulohumoral immunodepletion
Collagenosis, psoriasis
Cutaneous disorders: common acne
Cystitis
Evolutive tuberculosis: laryngitis - adenitis - osteomyelitis
Functional hypercorticism or adrenal exhaustion - prediabetes
General fatigability, loss of intellectual vitality
Loss of general vitality profound senescence
Osteoarticular and muscular disorders

SPECIFIC

Abscesses	Acne (juvenile)
Arteriosclerosis	Arthritis (chronic)
Boils	Burns
Constipation	Cystitis
Fatigue (infection or disease)	Gallstones
Hallucinations	Hemorrhoids
Herpes (zoster)	Hives
Insect Bites	Low Blood Pressure
Osteitis	Periodontal Disease
Psoriasis	Rheumatism (chronic)
Sleeping Disorder (waking early)	Sore Throat

KNOWN USES

CuAuAg is used in acute cases of rhinopharyngitis presenting with complications of otitis, adenitis, tonsillitis, and sinusitis. It can also act as a flu preventative in the fatigued and elderly. It rapidly modifies intelligence, imagination and reasoning power when they become signs of anergic diathesis. When concentration difficulties are accompanied by an overall reduction of vitality, CuAuAg can be given. CuAuAg increases the bodily defences especially in Anergic diathesis common to the three types of inflammatory rheumatism (chronic evolutive polyarthritis, ankylosing spondylarthritis, and acute articular rheumatism).

FLUORINE

(F)



INGREDIENTS

Each 2 ml (0.07 fl.oz.) ampoule contains 0.2 mg fluorine from sodium fluoride.

MODE OF ACTION

Fluorine is an element that regulates the metabolism of phosphorus and calcium. Fluorine acts to normalize the parathyroid gland and is necessary for the formation of strong, hard bones and teeth that resist decay. Geographical deficiencies of fluoride have been linked to increases of brittle bones in the elderly resulting in disabling hip fractures. As an oligotherapy remedy, fluorine is known to have a positive effect on calcium retention in the osseous tissues, and has been used for ligament and osseous conditions and as part of the preventive treatment for osteoporosis (Padrazzi, 1988). Fluorine It can also play an essential role in inflammatory diseases, such as rheumatoid arthritis. Fluorine makes up an integral part of potassium fluorosilicate (K₂SiF₆) believed to be the active anti-inflammatory compound found in marine sponges.

INDICATIONS

GENERAL

Arthritis and rheumatism (chronic)
Disorders of the vertebral statics: scoliosis
Hyperparathyroidism
Osteoarticular and muscular disorders
Osteoporosis
Retardation of calcification - prophylaxis of dental caries
- toothache - osteoligamentary weakness: ligamentary hyperlaxity

SPECIFIC

Arthritis (chronic)	Arthrosis (with osteoporosis)
Decalcification	Dental Cavities
Growth Disturbances	Lumbago
Menopause (osteoporosis)	Osteitis
Osteochondritis	Osteomalacia
Osteoporosis	Pregnancy
Rheumatism (chronic)	Scoliosis
Varicose Veins	

KNOWN USES

Fluorine is traditionally used in oligotherapy for the maintenance of healthy ligaments and bones. It is generally associated with oligotherapy remedy MnCo.

IODINE

(I)



INGREDIENTS

Each 2 ml (0.07 fl.oz.) ampoule contains 24 mcg of iodine from potassium iodide.

MODE OF ACTION

The primary role of iodine is as a component of thyroid hormone, and ultimately the regulation of cellular oxidation. The thyroid hormones (T3 and T4) accelerate cellular reactions, increase oxygen consumption, and increase growth and protein synthesis. Iodine is a cofactor for enzymes involved in the immune system. These include catalase and myeloperoxidase, which are both bacteriocidal in nature.

INDICATIONS

GENERAL

Cardio respiratory disorders
Hormonal homeostasis: dysmenorrhea, functional myxoedema, endocrine dysfunction, juvenile diabetes
Viral diseases - mycosis

SPECIFIC

Abscesses	Amenorrhea
Arthritis (rheumatoid)	Burns
Fatigue (psychic)	Goiter
Hypertension (essential)	Hypertension (stress)
Impotence	Libido, weak (male)
Myocardial Infarction	Obesity (hypothyroid)
Premenstrual Syndrome (PMS)	Sciatica
Seborrhea (hyper)	Seborrhea (hypo)
Thyroid (hyper)	Vertigo

KNOWN USES

Iodine has traditionally been used in oligotherapy for the maintenance of normal thyroid function. Deficiencies are linked to endocrine disturbances that could lead to increased risk of breast and ovarian cancer. One of the most damaging effects on iodine deficiency is on the development of the brain and may cause mental retardation, as well as hypothyroidism, goiter, and other growth and developmental abnormalities.

LITHIUM

(Li)



INGREDIENTS

Each 2ml (0.07 fl. oz.) ampoule contains 4.07 mg lithium gluconate in a 10% alcohol and 5% glucose aqueous solution.

MODE OF ACTION

Lithium is a trace element that intervenes in the transmission of nerve cells by affecting membrane potential. It has a sedative and non-hypnotic action. Lithium also acts upon urinary function by stimulating the elimination of urea and uric acid. Lithium has been successfully used in the prevention of both manic and depressive mood swings. Studies at Rockland State Hospital, New York (1975) suggest that lithium may play a role in controlling alcoholism. Lithium can change drinking habits by affecting or countering the action of alcohol on the brain. Lithium may boost the level of acetylcholine.

INDICATIONS

GENERAL

Behavior problems, insomnia, manic depression, anxiety, hyperirritability, gout and various types of pain. Lithium can also help in various eating disorders (anorexia nervosa).

SPECIFIC

Anorexia	Anxiety
Depression	Dysmenorrhea
Fatigue (non-specific)	Gout
Hallucinations	Hypertension (stress)
Lumbago	Lumbago (functional)
Migraine (tension)	Obesity (anxious overeating)
Sleep disorder (bedwetting)	

MAGNESIUM (MG)



INGREDIENTS

Each 2 ml (0.07 fl.oz.) ampoule contains 0.1044 mg magnesium from magnesium gluconate.

MODE OF ACTION

Approximately 60% of the body's magnesium is located within the skeletal structure of the body and is essential for normal bone formation. The remainder of the body's magnesium is found in the intracellular form. It performs several functions: controls membrane permeability, muscular contraction, nerve impulse conduction, intracellular fluid regulation, blood viscosity, activation of enzyme systems, and the regulation of protein synthesis and calcium metabolism. As an oligotherapy remedy, magnesium is involved in numerous physiological activities. It primarily affects the initiation of post-synaptic potential (Padrazzi, 1988). Supplementary magnesium has been used successfully in the treatment of various cardiovascular and related metabolic conditions, such as elevated cholesterol and triglycerides (and imbalances of the HDL/LDL ratio), arrhythmias, and for tetanic muscular conditions (such as occur during alcohol withdrawal) and for the premenstrual syndrome. Magnesium plays an essential role in the hormonal regulation via insulin, estrogen and thyroid hormone. Magnesium stimulates the secretion of the adrenal gland (corticosteroids) and of the parathyroid gland.

INDICATIONS

GENERAL

Digestive disorders	Leukocyte formation
Myocardial insufficiency	
Osteoarticular and muscular disorders	
Parasites	Psychosomatic illnesses
Water retention	

SPECIFIC

Adrenal (hypo)	Alcoholism (chronic)
Amenorrhea	Asthma
Constipation	Dysovulation
Enterocolitis	Fatigue (psychic)
Gallbladder Dysfunction	Gallstones
Goiter	Hayfever
Hepatobiliary Insufficiency	Muscle Cramps
Obesity (digestive)	Osteoporosis
Parasite	Premenstrual syndrome (PMS)
Sore Throat (chronic)	Warts
Travel Sickness	

KNOWN USES

Magnesium is a complementary remedy for dystonia diathesis, and is indicated for spasmophilic pains, colitis, and neurovegetative dystonia. It is a complementary remedy for dystonic conditions, such as with MnCo and sulfur remedies to alleviate alternation of constipation and diarrhea. Used with MnCo, it alleviates hemorrhoids related to intestinal disturbances. Used with sulfur, it treats asthma accompanied by migraine, eczema, and urticaria. When used in combination as CuMgMnZn, it can be used for alleviating acute pleurisy, tonsillitis, bronchitis, pneumonia, acute rhinopharyngitis, colds, and flu.

MANGANESE (MN)



INGREDIENTS

Each 2 ml (0.07 fl.oz.) ampoule contains 0.0728 mg of manganese from manganese gluconate.

MODE OF ACTION

Manganese plays a chief role in the synthesis of glycoproteins and glycolipids such as collagen. Thus, manganese is important in tissue healing and connective tissue disorders. Mn is a cofactor for many enzymes and activates the mitochondrial form of superoxide dismutase (SOD), a critical antioxidant that is adversely affected by a manganese deficiency. Manganese is required for the production of neurotransmitters. Mn is essential for the biosynthesis of protein via numerous enzyme systems (phosphotases, DNA polymase, peptidases, etc.).

INDICATIONS

GENERAL

Collagen and connective tissue disease, allergies, and inflammatory disorders.

General endocrine function, neurological and psychosomatic disturbances including migraines

SPECIFIC

Arthritis (chronic)	Asthma
Eczema	Gallbladder Dysfunction
Hayfever	Hypermenorrhea
Impotence	Insect Bites
Memory Decline	Menopause (cardiovascular)
Myocardial Infarction	Neuralgia
Rheumatism (chronic)	
Sleep Disorders (difficulty waking)	
Urticaria	Vertigo

KNOWN USES

Traditionally, manganese has been used in oligotherapy to alleviate symptoms associated with allergy.

MANGANESE-COBALT (MN-CO)



INGREDIENTS

Each 2 ml (0.07 fl.oz.) ampoule contains 0.0728 mg of manganese from manganese gluconate, and 0.0726 mg of cobalt from cobalt gluconate.

MODE OF ACTION

This combination of bio-activating metals is formulated for individuals in mid-life or later. Cobalt plays a control role in the formation of cobalamin (vitamin B12) and in preventing anemia. Cobalt also acts as a co-factor in antioxidant therapy for the enzyme catalase. Manganese is a co-factor for a variety of enzymes and is the central element in the mitochondrial form of superoxide dismutase (SOD).

INDICATIONS

GENERAL

Mn-Co is widely used in psychogenic illnesses, skin disorders, osteoporosis and inflammatory disease.

SPECIFIC

Amenorrhea	Anemia
Anorexia	Anxiety
Arthrosis	Constipation
Dystonia	Enterocolitis
Fatigue (mental strain)	Fatigue (non-specific)
Gallstones	Gout
Headache	Hemorrhoids
Herpes (simplex)	Libido, weak (female)
Lumbago (osteoporotic)	Memory Function
Migraine (hormonal)	Muscle Cramps and Spasms
Numbness	Obesity (water retention)
Sinusitis (chronic)	Varicose Veins
Water Retention	

KNOWN USES

Manganese-cobalt is traditionally used in oligotherapy for the maintenance of healthy circulation, and as a combination oligotherapy remedy, primarily for its action on dystonic diathesis. It is indicated for renal insufficiency, arteriosclerosis, circulation problems, and neurovegetative dystonia. It has also been used for conditions such as senile myocarditis (as complementary treatment), coronaritis (as complementary treatment), menopausal anemia (as complementary treatment), varicosis, arteriole hypertension, extra systoles, and tachycardia (as complementary treatment).

MANGANESE-COPPER (MN-CU)



INGREDIENTS

Each 2 ml (0.07 fl.oz.) ampoule contains 0.0728 mg of manganese from manganese gluconate, and 0.0726 mg of copper from copper gluconate.

MODE OF ACTION

The biocatalytic complex Mn-Cu is essential in normalizing enzyme activity associated with all forms of inflammatory disorders. Both copper and manganese play an essential role in the formation of superoxide dismutase (SOD), an enzyme that prevents free radical damage to cellular membranes and tissues. Copper can stimulate the immune system and increase B-lymphocyte production. Mn and Cu are both important in general endocrine functions, including the hypothalamopituitary complex, the adrenal gland (ascorbic acid oxidase) and the stimulation of the thyroid gland. Mn and Cu are required for glucose utilization and the synthesis of glycolipids and glycoproteins such as collagen.

INDICATIONS

GENERAL

Allergies, adenopathies, humoral immunodepletion
Colds, influenza - asthma
Chronic infectious diseases
Cutaneous: acne rosacea, dermatitis, burns, wounds
Enterocolitis, duodenal disorders, constipation and diarrhea
Hormonal homeostasis: dysregulation of the sexual cycle, frigidity
Osteoarticular disorders
Psychosomatic disorders
Urinary disorders: nephritis

SPECIFIC

Anorexia	Anxiety (chronic)
Articular Pain	Arthritis (chronic)
Bronchitis (chronic)	Cirrhosis
Cystitis	Decalcification
Dysmenorrhea	Dysovulation
Eczema	Emphysema
Enterocolitis	Gastritis
Goiter	Growth Disturbance (excess)
Hayfever	Hepatobiliary Insufficiency
Infertility (male)	Laryngitis (chronic)
Lumbago	
Menopause (fatigue and depression)	
Osteomalacia	Periodontal Disease
Pregnancy	Prostatitis
Sciatica	Scoliosis
Sinusitis	Sleeping Disorders
Thyroid (hypo)	Warts

KNOWN USES

Mn-Cu can be used to alleviate hyposthenia. It normalizes enzyme activity associated with all forms of inflammatory disorders. It is important in endocrine functions, including the hypothalamopituitary complex, the adrenal gland and the stimulation of the thyroid gland. Mn-Cu is primarily indicated for physical, intellectual, and psychological fatigue; bronchial fragility; duodenal problems; deforming arthritis; and infectious or allergic oto-rhino-laryngology.

MANGANESE-COPPER-COBALT (MN-CU-CO)



INGREDIENTS

Each 2 ml (0.07 fl.oz.) ampoule contains 0.0728 mg of manganese from manganese gluconate, 0.0726 mg of copper from copper gluconate, and 0.0726 mg of cobalt from cobalt gluconate.

MODE OF ACTION

The biocatalytic complex of Mn-Cu-Co is especially useful in a variety of inflammatory disorders. Both manganese and copper are central elements in a variety of metalloenzymes including superoxide dismutase (SOD) and thus help prevent free radical pathologies and associated inflammatory disorders. Cobalt acts as a cofactor in antioxidant therapy for the enzyme catalase. These three oligo-elements are essential for normal bone and collagen formation. Deficiencies will result in abnormal bone development and decalcification. Cobalt via vitamin B12 can help reduce spasms and increase blood flow.

INDICATIONS

GENERAL

Cardiorespiratory disorders
Colitis
Cutaneous disorders: acne
Hormonal homeostasis: thyroid insufficiency
Osteoarticular disorders
Psychosomatic disorders: asthenia

SPECIFIC

Anaemia	Decalcification
Gastritis	Low Blood Pressure
Periodontal Disease	Seborrhea (hypo)

KNOWN USES

Traditionally used in oligotherapy to normalize the state of asthenia. Essential factor for glutathione peroxidase (GSH-Px) an enzyme that catalyzes the reduction of hydrogen peroxidase and various organic peroxides, which are highly reactive compounds produced by normal metabolic and detoxification processes. GSH-Px prevents the free radical damage of these substances in the cell which prevents cellular damage. GSH-Px also reduces the production of inflammatory prostaglandins and leukotrienes. In oligotherapy, selenium is a trace mineral that intervenes with the glutathione peroxidase mechanism and that neutralises free radicals, opposing ageing and cellular degeneration. It modifies the terrain in the course of muscular and cutaneous affections.

PHOSPHORUS (P)



INGREDIENTS

Each 2 ml (0.07 fl.oz.) ampoule contains 0.14 mg of phosphorus from disodium phosphate.

MODE OF ACTION

Phosphorus exists as phosphate in the body and is found in both skeletal and intracellular fluids. Phosphorus is found in all cells of plants and animals and is essential for normal growth and development. Its mode of action is mainly by way of bonding, polymer formation, hydration, chemical transport and buffering. Intracellularly, phosphorus is involved in four metabolic functions:

1. Energy transport and formation via ATP and ADP
2. A major constituent in cellular membranes as phospholipids
3. Structural requirement for nucleoproteins (DNA, RNA)
4. Buffering, calcium transport and osmotic pressure of intracellular fluids

Phosphorus also plays key roles in circulatory function (RBC), nerve transmission (myelin sheath) and muscle contraction (ATP). Phosphorus is a central component of vitamin B6.

INDICATIONS

GENERAL

Osteoarticular and muscular disorders, cramps
Cardiorespiratory disorders and circulation problems
Hormonal homeostasis: dysregulation of thyrocalcitonin secretion

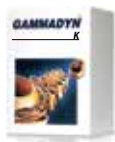
SPECIFIC

Fatigue (mental strain)	Fatigue (psychic)
Goiter	Lumbago (osteoporotic)
Menopause (fatigue and depression)	
Menopause (cardiovascular)	Muscle Cramps
Numbness	Osteochondritis
Osteomalacia	Osteoporosis
Rheumatism (chronic)	Scoliosis

KNOWN USES

Phosphorus has been traditionally used in oligotherapy to act on neurovegetative dystonia (particularly for tetanus), spasmophilia, and Dupuytren disease. Phosphorus has also been used to treat painful back in young people and muscular contractures. It is also used for diuresis disturbance and acts on parathyroid.

POTASSIUM (K)



INGREDIENTS

Each 2 ml (0.07 fl.oz.) ampoule contains 0.04 mg of potassium from potassium gluconate.

MODE OF ACTION

Potassium in the body occurs intracellularly and serves many functions, including:

1. Intracellular body fluid regulation, including osmotic pressure, buffering, viscosity, CO₂ transport (RBC), and solubilization of proteins.
2. Membrane effects, including membrane permeability, sodium pump action, muscular contraction, and nerve impulse conduction.

Potassium is often employed in the control of water retention. Potassium is also used with magnesium in the treatment of angina and arrhythmias and in the treatment of other degenerative diseases.

INDICATIONS

GENERAL

Adrenal function
Arthritis (chronic)
Electrolyte balance
Nerve transmission
Water retention

SPECIFIC

Adrenal (hyper)	Alcoholism (chronic)
Arthritis (poly)	Articular Pain
Diarrhea	Fatigue (due to mental strain)
Gallbladder Dysfunction	Gallstones
Hepatobiliary Insufficiency	Lumbago
Neuralgia	Pre-menstrual syndrome (PMS)
Rheumatism (chronic)	Sciatica

KNOWN USES

Traditionally, potassium has been used in oligotherapy to alleviate symptoms associated with rheumatoid arthritis, and for patients with an elevated sedimentation rate. It seems that potassium is used primarily for its action on chronic rheumatism. In addition, it is also used for cramps and muscular manifestations. Generally, potassium remedy is associated with Manganese and Copper-Silver.

SELENIUM (SE)



INGREDIENTS

Each 2 ml (0.07 fl.oz.) ampoule contains 40 mcg of selenium from sodium selenite.

MODE OF ACTION

Selenium is an essential factor for glutathione peroxidase (GSH-Px), an enzyme that catalyzes the reduction of hydrogen peroxide and various organic peroxides, highly reactive compounds that are products of normal metabolic and detoxification processes. GSH-Px scavenges these free radicals within the cells, in the cytosol and the mitochondrial matrix, to prevent damage to the cell, the mitochondria, the microsome and the lysosome membranes. GSH-Px functions in cooperation with superoxide dismutase (SOD). GSH-Px also reduces the production of inflammatory prostaglandins and leukotrienes. Selenium deficiency results in cardiomyopathy. Free radical damage is also known to be a major component in cataracts, acne, periodontal disease and tendinitis and bursitis. It has been theorized that selenium prevents damage to the inhibitors in DNA that control the cells' tendency to multiply.

INDICATIONS

GENERAL

Auto-immune disorders
Free radical damage
Gastrointestinal disorders
Immune deficiencies
Inflammatory disorders
Male reproductive function
Skin and mucous membrane disorders

SPECIFIC

Acne	Aging
Asthma	Bursitis
Candidiasis	Cataracts
Celiac disease	Cervical dysplasia
Crohn's disease	Dermatitis herpetiformis
Eczema	Multiple sclerosis
Myopathy	Osteoarthritis (chronic)
Periodontal disease	Psoriasis
Rheumatoid arthritis (chronic)	Seborrheic dermatitis
Tendinitis	Ulcerated skin
Vasculitis	Viral hepatitis

KNOWN USES

Selenium has been traditionally used in oligotherapy to neutralize free radicals and to protect against ageing, cellular degeneration, and muscular and cutaneous infections.

SULFUR (S)



INGREDIENTS

Each 2 ml (0.07 fl.oz.) ampoule contains 0.3 mg of sulfur from sodium thiosulfate.

MODE OF ACTION

Sulfur compounds are metabolically important because of their ability to interconvert disulfide and sulfhydryl groups in oxidation-reduction reactions. Disulfide and sulfhydryl bonds provide the stabilization and configuration for protein molecules. Mucopolysaccharides (especially chondroitin sulfate and collagen) contain sulfur. Sulpholipids are found in the liver, brain and kidneys. Sulfur is found in several amino acids including cysteine and methionine. Taurine, the precursor for bile acid production, also requires sulfur via cysteine and plays a role in neuropathologies.

INDICATIONS

GENERAL

Cardiorespiratory disorders: asthma
Cutaneous disorders: dermatosis, dandruff
Digestive disorders: hepatobiliary insufficiency
Osteoarticular and muscular: intervertebral disk prolapse, arthritis (chronic) and arthrosis
Psychosomatic: migraine

SPECIFIC

Acne	Adrenal (hyper)
Amenorrhoea	Arthritis (chronic, rheumatoid)
Arthrosis (with osteoporosis)	Articular Pain
Asthma	Bronchitis (chronic)
Cholesterol	Dysmenorrhoea
Eczema (chronic)	Emphysema
Epileptic seizures	Gallstones
Gout	Hayfever
Hepatobiliary Insufficiency	Herpes (simplex)
Herpes (zoster)	Insect Bites
Laryngitis (chronic)	Libido, weak (male)
Migraine	Neuralgia
Pregnancy	Prostatitis (acute)
Psoriasis	Rheumatism (chronic)
Sciatica	Sinusitis
Vertigo	

KNOWN USES

Traditionally used in oligotherapy in the course of repeating affections such as migraines, eczema, superficial body growths, arthritis and osteoarthritis. Sulfur is a complement to manganese in the treatment of arthritic or atro-tuberculosis states. For symptomatic treatment (oligostim), use sulfur as a terrain modifier in the course of cutaneous repetitive affections and rheumatic complaints.

ZINC (ZN)



INGREDIENTS

Each 2 ml (0.07 fl.oz.) ampoule contains 0.0674 mg of zinc from zinc gluconate.

MODE OF ACTION

Zinc plays a major role in the synthesis and metabolism of protein. Because of its importance in protein synthesis, it is essential to healing, growth and sexual maturation. Zinc activates over 70 enzymes, including nucleic acid polymerases (cell growth and repair), superoxide dismutase (antioxidant) and peptidases (protein digestion). Zinc acts in a synergistic manner with vitamins A and C, and the minerals magnesium and manganese. The zinc requirement of most adults is in the vicinity of 10-20 mg/day. Strial and white spots on the fingernails may indicate long-term zinc depletion. Zinc is now used for a variety of medical problems, including: postsurgical use; parenteral nutrition; sickle cell anemia (prevents sickling of red blood cells); rheumatoid arthritis (anti-inflammatory); acne and other skin disorders; loss of taste; impaired immunity and gonadal insufficiency. Research has demonstrated that zinc deficiency can compromise cellular immune function. Patients on total parenteral nutrition developed T-cell dysfunction in association with severe zinc deficiency. After 12 days of intravenous zinc supplementation (12 mg/day) T-cell function increased dramatically and averaged 160 percent of control.

INDICATIONS

GENERAL

Adrenal insufficiency
Cutaneous disorders: acne
Gonadic disorders - impotence, infertility
Hypothalamo - pituitary disorders
Immunity homeostasis - candidiasis

SPECIFIC

Alcoholism (chronic)	Asthma
Cholesterol	Cirrhosis
Growth Disturbances	Hayfever
Infertility	Libido (weak)
Lumbago	Prostatitis
Thyroid (hyper)	

KNOWN USES

Zinc has been traditionally used in oligotherapy to help maintain immunity. It helps in tissue formation, and also helps the body to metabolize proteins, fats, and carbohydrates. Its anti-inflammatory activity, which is the basis of therapeutic use, other than acrodermatitis enteropathica, is not well known: production of cytokines, antioxidant activity.

ZINC-COPPER (ZN-CU)



INGREDIENTS

Each 2 ml (0.07 fl.oz.) ampoule contains 0.0674 mg of zinc from zinc gluconate, and 0.0726 mg of copper from copper gluconate.

MODE OF ACTION

The catalytic complex of Zn-Cu is essentially used in problems associated with the regulation of steroid secretion. These oligo-elements also play an essential role in skin and inflammatory disorders via superoxide dismutase. Zinc and copper are essential for proper endocrine function (hypothalamo-pituitary, adrenal, etc.). Copper functions as the central element in the metallo-enzyme, dopamine-hydroxylase, which is essential for the synthesis of norepinephrine by the adrenal gland. Deficiencies of zinc and copper effect affect neurotransmitters and histamine release in the brain, thus effecting also affecting neurological behavior.

INDICATIONS

GENERAL

Adrenocortical dysregulations
Cutaneous disorders:
Dysthyroidism with a hypo-tendency
General immunodepletion (humoral and cellular)
Hormonal homeostasis: all syndromes of hypothalamo-pineal gland- glandular dysregulation, hypomenorrhea, sterility, menopause, impotence
Inflammatory syndromes
Pancreas: dysregulations of insuline secretion

SPECIFIC

Growth Disturbances	Impotence
Libido, weak (male)	Menopause (neuroendocrine)
Psoriasis	Sleep Disorder (bedwetting)

KNOWN USES

Zinc-copper has been traditionally used in oligotherapy to support the thyroid and endocrine systems. It has been used to address the functional problems of benign prostatic hypertrophy (BPH), hormonal development abnormalities in children. ZnCu can also resolve the symptoms of pituitary dysfunction as well as treat the pituitary/genital maladaptation syndrome. In summary, zinc can assist the pituitary and gonads, while copper assists the adrenals.

ZINC-NICKEL-COBALT (ZN-NI-CO)



INGREDIENTS

Each 2 ml (0.07 fl.oz.) ampoule contains 0.0674 mg of zinc from zinc gluconate, 0.0726 mg of nickel from nickel gluconate, and 0.0726 mg of copper from copper gluconate.

MODE OF ACTION

This association of three trace-elements has a first-rate catalytic role in hypophyso-pancreatic dysfunctions and prediabetic conditions: indeed, they concentrate preferably in the pancreas and, by their hyperglycemic, synergistic effect, improve the tissular fixation of glucose. Zinc forms specific metallo-enzymes such as carboxypeptidase and renal dipeptidase that digest protein. Zinc also plays an active role in glucose metabolism. Cobalt is the central element in various peptidases and stimulates glycogen and insulin release by the liver and pancreas. Nickel is important for liver function and glycogen release from tissues. Nickel has been shown to enhance the uptake of glucose into rat adipose tissue and the incorporation of glucose into glycogen. Nickel also plays an important role in several enzymes that deal with carbohydrate metabolism.

INDICATIONS

GENERAL

Digestive disorders: food allergies
Pituitary-pancreatic dysregulations

SPECIFIC

Adrenal (hyper)	Diabetes (mellitus)
Gallbladder Dysfunction	Hypoglycemia
Indigestion	Libido, weak (male)
Pancreatitis	Sleep Disorders (bedwetting)

REFERENCES

- Abraham, G. E., et al., "Serum and Red Cell Magnesium Levels in Patients with Premenstrual Tension," *Am. J. Clinical Nutr.*, 34: 2364-2366, 1981.
- Alberts, B., et al., "The Plasma Membrane," *In The Molecular Biology of the Cell*, Garland Pub. New York, 1983.
- Allen, J., et al., "Severe Zinc Deficiency in Humans; Association with T-Lymphocyte Dysfunction," *Ann. Int. Med.* 95: 154-157, 1981.
- Andersen ME, Gearhart JM, Clewell HJ. Pharmacokinetic data needs to support risk assessments for inhaled and ingested manganese. *Neurotoxicology.* 1999;20(2-3):161-71
- Anonymous, "Role of Zinc in Enzyme Regulation and Protection of Essential Thiol Groups," *Nutrition Reviews*, Vol. 44: 9, 309-311, 1986.
- Antoniou, L., "Zinc In Clinical Medicine," *Drug Therapy* pp. 92-102, 1984.
- Arora A, and Shukla Y. Induction of preneoplastic altered hepatic foci following dietary sulphur supplementation. *Hum Exp Toxicol.* 2004;23(5):229-34.
- Aschner M, Dorman DC. Manganese: Pharmacokinetics and molecular mechanisms of brain uptake. *Toxicol Rev.* 2006;25(3):147-54
- Ashmead, H., et al., "Intestinal Absorption of Metal Ions and Chelates," Charles C. Thomas Publ., Springfield, 11, 1985.
- Aston, B., "Manganese and Man," *J. of Orthomolecular Psychiatry*, 9: 4, 237-249, 1980.
- Bazhora, H., et al., "The Action of the Oligo-Elements Cu, Mn, et. Co on the Synthesis of Antibodies from Lymphoid Tissue," *Mikrobiolog Zh*, 36: 6, 771-776, 1974.
- Bert, L., et al., "Carboxypeptidase A," *In Zinc Enzymes*, T. Spiro, ed., John Wiley & Sons, New York, pp. 27-70, 1983.
- Bertrand, G., 8th Int. Congress, Appl. Chem., 28: 30, 1912.
- Burbeau, A., et al., "Zinc, Taurine and Epilepsy," *Arch. Neurol.* 30: 52, 1 974.
- Burch, R., et al., "Symposium on Trace Elements," *In: The Medical Clinics of North America* 60: 4, 1976.
- Blodgett RC, Pietrusko RG. *Scand J Rheumatol Suppl.* 1986; 63:67-78. Long-term efficacy and safety of auranofin: a review of clinical experience.
- Bonham M, o'Connor JM, Hannigan BM, and Strain JJ. Review article: The immune system as a physiological indicator of marginal copper status? *British Journal of Nutrition.* 2002; 87: 393-403.
- Boron, Chromium, Copper, Iodine, Iron, Manganese, Molybdenum, Nickel, Silicon, Vanadium, and Zinc. A Report of the Panel on Micronutrients, Subcommittees on Upper Reference Levels of nutrients and of Interpretation and Uses of Dietary Reference Intakes, and the Standing Committee on the Scientific Evaluation of Dietary Reference Intakes [Internet] [cited 22-June-2007]. National Academy Press. Washington, D.C. 2000. Available from: <http://www.nap.edu/>.
- Brown RA and Wells TM. Interaction and aggregation of sodium aurothiomalate and poly L-lysine. *J Pharm Pharmacol.* 1992; 44(6):467-71.
- Bruce, A. et al. The effect of selenium and vitamin E on glutathione peroxidase levels and subjective symptoms in patients with arthrosis and rheumatoid arthritis.
- Chircorian A and Barrios Am. Inhibition of lysosomal cysteine proteases by chrotherapeutic compounds: a possible mechanism for the antiarthritic activity of Au(I). *Bioorg Med Chem Lett.* 2004; 14(20):5113-6.
- Christensen JM, Poulsen OM, and Thomsen M. A short-term cross-over study on oral administration of soluble and insoluble cobalt compounds: sex differences in biological levels. *Int Arch Occup Environ Health.* 1993;65(4):233-40.
- Clark, LC et al. *Journal of Nutrition.* 1986;116:170.
- The Cooperating Clinics Committee of the American Rheumatism Assoc.: A Controlled Trial of Cyclophosphamide in Rheumatoid Arthritis, *New Eng. J. Med.*, 283: 883-889, 1970.
- Clin Cancer Res. 2006; 1(12):2178-84. Young, R. S., "Cobalt in Biology and Biochemistry," Academic Press, London, 1979.
- Coleman, J. E., "Cd Binding to Apoalkaline Phosphatase as Functions of Phosphate, Magnesium pH, and Cd (II) Concentration," *J Biol. Chem.*, 247: 4718, 1972.
- The Cooperating Clinics Committee of the American Rheumatism Assoc.: A Controlled Trial of Cyclophosphamide in Rheumatoid Arthritis, *New Eng. J. Med.* 283: 883-889, 1970.
- Coze, J. F., et al., "Utilization of Manganese in the Treatment of Chronic Laryngitis," *Sem Hop Paris*, 57: (21/22), 1139-1143, 1981.
- Coze, J. B., et al., "Utilization du Manganese Cuive dans le Traitement des Pharyngites Chroniques," *SEM Hop Paris*, 57: 21-24, 1981.
- Cullinan, P., "The Mechanism of Action of Homeopathic Remedies - Towards a Definitive Model," *J of Comp. Med.*, July, 1985.
- Danscher G. Light and electron microscopic localization of silver in biological tissue. *Histochemistry.* 1981; 71(2):177-82.
- Dawson and Laws, "The Oligo Elements in Rheumatology," *Journee' de Medecine Fonctions Nelle*, 27 Avril, 1980.
- Draper, H. H., et al., "Advances in Nutrition Research," *In: Nutrition and Osteoporosis*, Plenum Press, New York, 2: 79, 1980.
- Ducros V, Favier A, and Guigues M. Selenium bioavailability as selenite (74Se) and as a selenium drug (76Se) by stable isotope methodology. *J Trace Elem Electrolytes Health Dis.* 1991 Sep; 5(3):145-54.
- Eisler R. Gold concentration in abiotic materials, plants, and animals: a synoptic review. *Environ Monit Assess.* 2004; 90(1-3):73-88.
- Ernestam S, Lampa J, Rogberg S, Rönnelid J, Kalreskog L and Hafström I. Evidence for immunostimulatory effects of intramuscular gold in patients with rheumatoid arthritis: correlation with skin reactions. *Rheumatol.* 2003; 30(8):1748-55.
- Fame, J., "Les Oligoelements en Rheumatology," *Il Cuivre, selenium, manganese, lithium etc.*, *Tempo Medical Belgique* 55: June 1985.
- Ferdinand, W., "The Enzyme Molecule," John Wiley and Sons, New York, 1976.
- Fineli, D. S., et al., "Interaction of Zinc and Lead on Aminolevulinat Dehydratase," *Biochem. Biophys. Res. Commun.*, 65: 303-311, 1975.
- Fingji, E. and Woodbury, D., "Absorption of Drugs," *In: The Pharmacological Basis of Therapeutics*, 5th ed., Macmillan Publ., New York, pp. 5-9, 1975.
- Forbes, R., et al., "Bioavailability of Trace Mineral Elements," *Ann Rev Nutr.*, 3: 212-231, 1983.
- Frederickson, H., et al., "The Neurobiology of Zinc," *In: Neurology and Neurobiology*, Vol. 11, A. Liss Pub., New York, 1983.
- Gottlieb NL. Comparative pharmacokinetics of parenteral and oral gold compounds. *J Rheumatol Suppl.* 1982; 8:99-109.
- Garfinkel, D, "Is Aging Inevitable? The Intracellular Zinc Deficiency Hypothesis of Aging," *Medical Hypothesis* 19: 117-137, 1986.
- Garison, R. and Somer, E., *The Nutrition Desk Reference* Keats Publ., Co., New Canaan, Conn., 1985. Goldhaber, SB. Trace element risk assessment: essentiality vs. toxicity. *Regulatory Toxicology and Pharmacology.* 2003;38:232-242
- Greaves M. Gemmotherapy and Oligotherapy Regenerators of Dying Intoxicated Cells. 2002. Xlibris;USA
- Greaves M. Gemmotherapy and Oligotherapy: Regenerators of Dying Intoxicated Cells. Xlibris Corporation: USA; 2002.
- Guyton, A., "Textbook of Medical Physiology," W. B. Saunders Co., Philadelphia, 1976.
- Health Canada. Dietary Reference Intake. [Internet] [Updated Aug 4, 2005] [Accessed February 23, 2007] Available at www.hc-sc.gc.ca/fn-an/nutrition/reference/table/ref_elements_tbls_e.html
- Kutsky, R., "Handbook of Vitamins, Minerals and Hormones," Van Nostrand Reinhold Co., San Francisco, 1981.
- Gregson, R. P, et al., "Fluorine is a Major Constituent of the Marine Sponge Halichondria Moorei," *Science*, 206: 1108-1109, 1979.
- Griem P and Gleichmann E. Gold antirheumatic drug: desired and adverse effects of Au(I) and Au(III) [corrected] on the immune system. *Z Rheumatol.* 1996; 55(6):430.
- Guyton, A. C., *Medical Physiology*, 5th ed., Saunders, Philadelphia, 1976.
- Hurley, L., "Clinical and Experimental Aspects of Manganese in Nutrition," *In: Clinical Biochemical and Nutritional Aspects of the Trace Elements*, A. R. Liss, New York, pp. 369-378, 1982.
- Ibata, Y., et al., "Electron Microscopic Demonstration of Zinc in Hippocampus Formation Using Timm's Sulfide-
- Institute of Medicine, Food and Nutrition Board [IOM]. Dietary Reference Intakes for Vitamin A, Vitamin K, Arsenic, Boron, Chromium, Copper, Iodine, Iron, Manganese, Molybdenum, Nickel, Silicon, Vanadium, and Zinc. A Report of the Panel on Micronutrients, Subcommittees on Upper Reference
- Institute of Medicine, Food and Nutrition Board (IOM). Dietary Reference Intakes for Vitamin A, Vitamin K, Arsenic, Boron, Chromium, Copper, Iodine, Iron, Manganese, Molybdenum, Nickel,
- Irving, J. T., "Theories of Mineralization of Bone," *Clin. Orthop*, 97: 225, 1973.
- Keen, C. L., et al., "Manganese in Biochemistry of the Ultratrace Elements," Frieden, (ed.), Plenum Press, New York, pp. 89-132, 1984.
- Kelley DS, Daudu PA, Taylor PC, Mackey BE, and Turnlund JR. Effects of low-copper diets on human immune response. *American Journal of Clinical Nutrition.* 1995; 62:412-6.
- Kosman, M. E., "Management of Potassium Problems During Long Term Diuretic Therapy," *J. A. M. A.* 230:5, 1974.
- Koryem HK, Taha KM, Ibrahim IK, and Younes LK. Liver toxicity profile in gold-treated Egyptian rheumatoid arthritis patients. *Int J Clin Pharmacol Res.* 1998;18(1):31-7.
- Kutsky, R., "Handbook of Vitamins, Minerals and Hormones," Van Nostrand Reinhold Co, San Francisco, 1981.
- Lansdown AB. Critical observations on the neurotoxicity of silver. *Crit Rev Toxicol.* 2007; 37(3):237-50.
- Levels of nutrients and of Interpretation and Uses of Dietary Reference Intakes, and the Natural Health Products Directorate: Compendium of Monographs [Internet] 2004 [Ottawa, Ontario; cited 22-June-2007]. Available from: <http://hc-sc.gc.ca/a.Copper-Draft>
- Limpscomb, W N., "Carboxypeptidase A: A Protein and an Enzyme," *Advan. Protein Chemistry*, 25: 1-78, 1971.
- Martin, W G., "The Neglected Nutrient Sulfur," *The Sulfur Institute Journal* Spring, 1968.
- Marovinic, J., "Endemic Goiter and Cretinism at the Dawn of the Third Millennium," *Ann. Rev Nutr* 3: 341-412, 1983.
- McKenna JK, Hull CM, and Zone JJ. Argyria associated with colloidal silver supplementation. *Int J O Dermatol.* 2003; 42(7):549.
- McNeil DL. Oral gold therapy in steroid-dependent asthma, nasal polyposis, and aspirin hypersensitivity. *Ann Allergy.* 1990; 65(4):288-90.
- Milano, R., et al., "Copper and Inflammation: A Possible Rationale for the Pharmacological Manipulation of Inflammatory Disorders," *Agents and Actions*, 16 (6): 504-513, 1985.
- Morassut P, Goldstein R, Cyr M, Karsh J, and McKendry RJ. Gold sodium thiomalate compared to low dose methotrexate in the treatment of rheumatoid arthritis—a randomized, double blind 26-week trial. *Rheumatology.* 1989; 16(3):302-6.
- National Academy Press. Washington, D.C. 2000. Available from: <http://www.nap.edu/>.
- Natural Health Products Directorate: Compendium of Monographs: Selenium [Internet] 2007 [Ottawa, Ontario; cited 23-Nov-2007]. Available from: <http://hc-sc.gc.ca/a.Copper-Draft b. Zinc-Draft>
- Natural Health Products Directorate (NHPD). Compendial monograph Magnesium. [Internet] [Updated April 3, 2007] [Accessed July 17, 2007] Available at http://www.hc-sc.gc.ca/dhp-mps/prodnatur/applications/licen-prod/monograph/mono_list_e.html
- Natural Medicines Comprehensive Database (NMCD). Potassium. [Internet] [Accessed July 16, 2007] Available at www.naturaldatabase.com
- Neville, J. N. et al., "Nutritional Status of Alcoholics," *Am. J Clin. Nutr.*, 21: 1329-1340, 1968.
- Nève J, Hanocq M, Peretz A, Khalil FA, and Pelen F. Absorption and metabolism of oral zinc gluconate in humans in fasting state, during, and after a meal. *Biol Trace Elem Res.* 1992 Jan-Mar;32:201-12.
- O'Brien MD, Camilleri M, von der Ohe MR, Philips SF, Pemberton JH, Prather CM, Wiste JA, Hanson RB. Mobility and tone of the left colon in constipation: a role in clinical practice? *Am J Gastroenterol.* 1996;91(12):2532-8
- O'Dell, B., "Biochemistry of Copper," *Medical Clinics of North America*, 60 (4): 687-703, 1976.

- Okunuki, K., et. al., "Structure and Function of Cytochromes," University Park Press, New York, 1968.
- Padrazzi P. L'Oligotherapie Reactionnelle. Editions Similia, 1988.
- Padrazzi P. L'Oligotherapie Reactionnelle. 1988. Editions Similia.
- PDRhealth. Potassium. [Internet] [Accessed July 16, 2007] Available at www.pdrhealth.com.
- Percival SS. Copper and Immunity. *Am J Clin Nutr.* 1998; 67(suppl): 1064S-8S.
- Percival SS. Copper and Immunity. *Am J Clin Nutr.* 1998; 67(suppl): 1064S-8S.
- Picard, M., "Utilization Therapeutique des Oligoelements," Ed. Maloine, Paris, 1982. Pfeiffer, C., "Mental and Elemental Nutrients," Keats Publ., New Canaan, Conn, 1975.
- Pfeiffer CC. Mental and Elemental Nutrients. 1975. Keats Publishing; Connecticut: 296-297
- Pixit, P. K., et. al., "Effects of Metal Ions and Sulfhydryl-Inhibitors on Glucose Metabolism by Adipose Tissue," *Amer. J. Physiol.* 213: 849-856, 1967.
- Pories, W. J., et. al., "Effects of Zinc Deficiency and Zinc Supplementation on Adrenal, Plasma, Steroids and Prasad, K.ed. Vitamins, Nutrition and Cancer. New York: Karger; 1984.
- Proc NZ Workshop on Trace Elements in NZ. Dunedin: U. of Otago; 1981:92.
- Reeves, P G., et. al., "Interrelationship of Thyroxine, Citrate and Renal Calcification in the Magnesium Deficient Rat," *Proc. Soc. Exp. Biol. Med.*, 137: 1104-1109, 1971.
- Sabichi AL, Lee JJ, Taylor RJ, Thompson IM, Miles BJ, Tangen CM et al. Selenium accumulation in prostate tissue during a randomized, controlled, short-term trial of l-selenomethionine: a Southwest Oncology Group Study.
- Schroeder, H., "The Trace Elements and Man," The Devin-Adair Co., Conn., 1975.
- Scudder, P R., et. al., "Serum Copper and Related Variables in Rheumatoid Arthritis," *Annals of the Rheumatic Diseases*, 37: 67-70 1978.
- Shah P, Griffith Sm, Shadforth MF, Fisher J, Dawes PT, Poulton KV, et al. Can gold therapy be used more safely in rheumatoid arthritis? Adverse drug reactions are more likely in patients with nodular disease, independent of HLA-DR3 status. *J Rheumatol.* 2004; 31(10):1903-5.
- Shearn, Martin, "The Medical Clinics of North America: Rheumatic Diseases," W. B. Saunders Co., Philadelphia, 61: 2, 1977.
- Shirina LI, Mazo VK. Mineral substance in human nutrition. Manganese: absorption and bioavailability. *Vopr Pitan.* 2006;75(5):4-14
- Siepmann M, Spank S, Kluge A, Schappach A, and Kirch W. The pharmacokinetics of zinc from zinc gluconate: a comparison with zinc oxide in healthy men. *Int J Clin Pharmacol Ther.* 2005;43(12):562-5.
- Stadel, B., "Dietary Iodine and Risk of Breast, Endometrial and Ovarian Cancer," *The Lancet*, April, 1976.
- Steadman CJ, Philips SF, Camilleri M, Talley NJ, Haddad A, Hanson R. Control of muscle tone in the human colon. *Gut.* 1992;33(4):541-6
- Stéphan F, Revuz J. *Ann Dermatol Venerol.* Zinc salts in dermatology. *Annales de Dermatologie et de Swales, J. D.*, "Magnesium Deficiency and Diuretics," *Br Med. J.*, 285: 1377-1378, 1982.
- Symptomology and Occurence," *Acta Medica Scandinavica (Supp.)* 661: 3-4. 1982. Stryer, L., "Introduction to Enzymes," In: *Biochemistry*, Freeman and Co., San Francisco, 1975.
- Stebbing, A. R. D., "Hormonesis - The Stimulation of Growth by Low Levels of Inhibitors," *The Science of the Total Env.*, 22: 213-234, 1982.
- Shah, B. G. "Chelating Agents and Bioavailability of Minerals," *Nutrition Research*, 1: 617-622, 1981.
- Standing Committee on the Scientific Evaluation of Dietary Reference Intakes [Internet] [cited 22-June-2007]. National Academy Press. Washington, D.C. 2000. Available from: <http://www.nap.edu/>.
- Silver Technique," *J. Histochem. Cytochem.* 17: 171-175, 1969.
- Thymus in Rats," *Life Sci.* 24: 177-181, 1979.
- Turnland JR, Jacob RA, Keen CL, Strain JJ, Kelley DS, Domek JM et al. Long-term high copper intake: effects on indexes of copper status, antioxidant status, and immune function in young men. *American Journal of Clinical Nutrition.* 2004; 79:1037-44.
- Underwood, E., "Trace Elements in Human and Animal Nutrition," Academic Press, New York, pp. 7, 1977.
- Veien NK, Hattel T, Justesen O, Norholm A. Oral challenge with nickel and cobalt in patients with positive patch tests to nickel and/or cobalt. *Acta Derm Venerol.* 1987;67(4):321-5
- Van der Velden VH, Hulsman AR. Autonomic innervation of human airways: structure, function, and pathophysiology in asthma. *Neuroimmunomodulation.* 1999;6(3):145-59
- Vénérologie. 2004;131(5):455-60.
- Veien NK, Hattel T, Justesen O, Norholm A. Oral challenge with nickel and cobalt in patients with positive patch tests to nickel and/or cobalt. *Acta Derm Venerol.* 1987;67(4):321-5.
- Veien NK, Hattel T, Justesen O, and Norholm A. Oral challenge with metal salts. (I). Vesicular patch-test-negative hand eczema Contact Dermatitis. 1983; 9(5):402-6.
- Virtamo, J. et al. Serum selenium and the risk of coronary heart disease and stroke. *Am J Epidemiol.* 1985;122:276-82.
- Wester, P O., et. al., "The Importance of Magnesium Ion, Magnesium Deficiency, Widdowson, E., et. al., "Chemical Composition of the Body," in Comar, C., ed., *Mineral Metabolism Academic Press*, 2: 57, 1964.
- Young, R. S., "Cobalt in Biology and Biochemistry," Academic Press, London, 1979.
- Young, R. S., Cobalt in "Biochemistry of the Essential Ultratrace Elements," Frieden, E., ed., Plenum Press, New York, 1984.

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THE PUREST HOMEOPATHIC SOURCE IN THE WORLD

Established over half a century ago in Belgium, unda® is renown for manufacturing exceptional homeopathic products utilized in supporting immune, lymphatic and endocrine systems to properly process and eliminate toxins from the body. In the production of all homeopathic remedies unda® uses only pure materials and herbs that are biodynamically grown or wildcrafted. unda® produces a broad range of homeopathic products in various potencies including: the unique Numbered Compounds, Gemmotherapy macerates, Schüssler Tissue Salts, Gammodyn Oligo Elements, Organotherapy, creams and oils, as well as homeopathic compatible dental care.



Numbered Compounds

unda® Numbered Compounds are the only complex remedies of their type, formulated with both plants and metals, acting on organotropic and energetic sensitivity levels respectively. Each remedy has its own fingerprint and is combined synergistically based on the integrated principles and theories of anthroposophy, oligotherapy, botany, Traditional Chinese Medicine and homeopathy. unda® Numbered Compounds are key in Biotherapeutic Drainage™ as they work on a physiological and “terrain” level, making them an excellent addition to practitioners’ prevention and treatment armamentarium.



Gemmotherapy

Gemmotherapy uses the extracts of fresh buds, shoots, roots or stems from developing plants, where the plant’s life essence is at its peak in the young growths. unda® Gemmotherapies are prepared in a natural glycerine and organic ethanol medium, which is then filtered and potentized at a 1/10th dilution (1X Hahnemannian) in a pure water, natural glycerine and organic ethanol medium. These complex remedies are macerated for increased patient compliance. This methodology captures the most complete set of highly-concentrated active constituents necessary for tissue regeneration, favourable growth development and essential drainage properties. unda® Gemmotherapies facilitate Biotherapeutic Drainage™ and is an excellent adjunct to the unda® Numbered Compounds.



Schüssler Salts

Schüssler Tissue Salts act on the cellular level to balance and restore cell function. unda® Tissue Salts are prepared using methods based on Dr. W. H. Schüssler Biochemical System of Medicine (in accordance with the German Homeopathic Pharmacopia (HAB)).

Seroyal Continuing Education

Seroyal offers research and practice-based continuing education. The mandate of Seroyal’s professional education program is to provide healthcare practitioners with world-class, clinically relevant information to assist in the integration of therapeutic modalities for a more complete, holistic approach. International experts in natural medicine and science will share knowledge, clinical experiences and reference materials to help expand your therapeutic capabilities and support your practice.

To learn more visit:

www.seroyalseminars.com

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