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Hyperbaric Oxygenation Therapy (HBOT) A good possibility in Neurorehabilitation but not accepted enough and generally used

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Danube Lecture

11th Danube Teaching Course

Kazimierz Dolny, May 13 - 15, 2015

Neurorehabilitation, basic program Restoration of the disturbed brain functions

- Rebuilding of the damaged neural network (post acute phase)
- Activation of the rebuilded neural network (HBOT)
- Reactivation of the neural network, rehabilitation program, (stimulation of proprioception)
- Special reactivation of neural network, (individual neurorehabilitation progam)

Activation of cortical network pressurized oxygen

- Reactivating of idling neurons
- Improving of oxygen cell respiration
- Restorating of blood brain barrier and cell membranes
- Promotion f neovascularization
- · Acting as scavenger of free radicals
- · Stimulation of the adaptive immune system

Neuro-Rehabilitation Methods stimulation of the proprioception

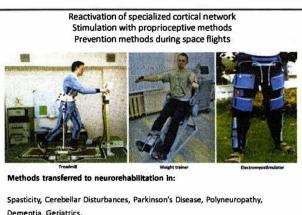


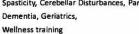
ADELI SYSTEM

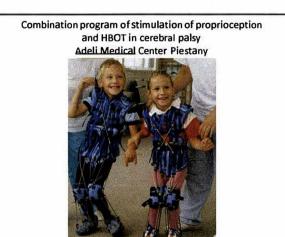
Cerebral paisy Traumatic Brain Deficits Stroke Apallic Syndrome Vertebral Spine Decompensation Syndrome Parkinson's Syndrome



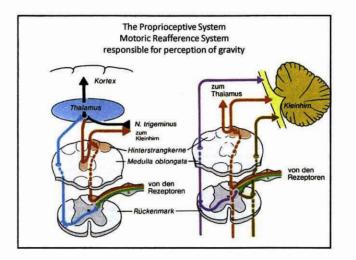
Souce: ADELI Fiver Dementia

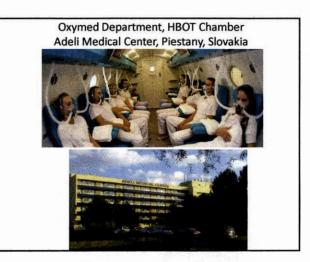






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The pressurized environment is not new

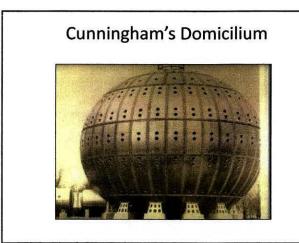
The first recorded use in history was a diving bell Alexander the Great used in the siege of Tyrus in 332 BC. Oxygen was discovered by Priestly 1774

- He warned that increased pressure may be toxic
- This held the field back for many years

1920s - 30s

Remarkable clinical results were obtained by Orville Cunningham, Professor of Anesthesia, University of Kansas 1928

The six stories stainless steel Domicilium was erected for Cunningham by Timken Ball Bearing Company in Cleveland



Pressurized O₂ adheres to all gas laws of physics

Henry's Law states there is a direct relationship between pressure and the amount of gas dissolved in solutes

Under hyperbaric condition oxygen is increased in:

- bone
- urine
- plasma
- lymph
- and most importantly in the cerebrospinal fluid

Under pressure free molecular oxygen is delivered directly to the cell for immediate metabolic use without energy exchange.

Edward Teller, Ph.D.

Dose equals

- Strength of pressure
- Time (length) of exposure
- Frequency
- Total number of treatments

How is it administered -Pressure Vessel

- Multiplace chamber
- Monoplace chamber
- Low pressure portable chamber

Types of pressure chamber

- Multiplace Chamber
- Monoplace Chamber
- Low Pressure Portable Chamber

History of Hyperbaric Oxygenation Chamber **Oxymed Department Adeli Center Piestany**









2014 Oxymed Center, Pies

Multiplace Chamber





Monoplace Chamber



Hyperbaric Oxygenation Treatment Administration

- Hyperbaric Oxygen Therapy is using 100% Oxygen at a greater pressure than in atmosphere
- 20-60 sessions are recommended (for insurance)
- Some patients are needing 100 sessions and more depending from the diagnosis
- Duration of one session 60 to 90 minutes

Effects of Pressurized Oxygen in Acute Brain Insult - 1

- · Reduces adhesion of WBCs (white blood cells) to endothelium
- · Perfuses all tissue spaces
- Life sustaining O2 available via retrograde perfusion in absence of a trickle phenomena
- Delivers metabolically available O₂ without chemical energy transfer - enough to sustain life without blood

Effects of Pressurized Oxygen in Acute Brain Insult - 2

- Under pressure, O₂ adheres to all the gas laws of physics
- Displaces all other gases in the body: – N₂, CO
- Follows the law of mass action
- Completely saturates hemoglobin
- Increases plasma O₂ by 2000%
- Dissolves in cerebrospinal fluid, lymph, bone and urine

Effects of Pressurized Oxygen in Acute Brain Insult - 3

- Reduces cerebral edema & ICP
- Limits the ischemic cascade
- Reduces CNS lactate peak in hypoxia
- Neutralizes toxic amines
- Disaggregation of platelets
- Increases Phagocytic activity of PMN cells (white blood cells)

Effects of Pressurized Oxygen in Chronic Brain Insult - 1

- Reactivates idling neurons
- Enhances plasticity
- Efficiently elevates diffusional driving force for O₂, thereby increasing tissue oxygen availability
- Promotes phagocytosis (internal debridement)
- Ameliorates multiple biochemical changes

Effects of Pressurized Oxygen in Chronic Brain Insult - 2

- Restores the integrity of the blood brain barrier and cell membranes
- Improves cell respiration, reduces cell byproducts – cytokines
- Promotes neovascularization
- Promotes epithelization

Effects of Pressurized Oxygen in Chronic Brain Insult - 3

- · Acts as scavenger of free radicals
- Bacteriostatic effects, synergizes with certain antibiotics
- Neutralizes certain toxins: clostridium, anaerobes
- Stimulates the adaptive immune system, especially in elderly (mice)

Influence on certain drugs

HBOT may enhance the effectiveness of certain drugs and extend the longevity of the product

Application in Neurology (1)

- Acute neurological conditions: Stroke, traumatic brain injury (TBI), encephalitis, spinal cord injury
- Progredient neurological deseases: Alzheimer dementia, ALS, MS, vascular dementia

Chronic neurological conditions: CP, apallic syndrome, traumatic brain deficit, spinal cord lesion

Application in Neurology (2)

- Acute Intoxication: **CO-intoxication**
- Decompression: diving accidents, air embolism
- Radiation accident

Cochrane Recommandations for HBOT

Traumatic Brain Injury - Artru (1976), Holbach (1974), Rockswold (1992), Ren (2001)

- Neonatal hypoxic encephalopathy Hutchinson (1966), Liu (2006)
- · Cerebral Palsy Machado (1989), Montgomery (1999), Packard (2000), Collet (2001), Mathai (2005)

Own experiences in Apallic Syndrome

Common project with HBOT Center

Fort Lauderdale (USA)

Dr. Richard Neubauer

Case M. J.

MJ, 31ª, male, AS, CO-intoxication, attempted suicide, remission state II-III during 12 years

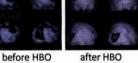
HBOT: 60 sessions

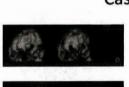
HBO home treatment: 350 sessions 1,5 ATA

Additional: physiotherapy logopedics

Significant improvement minimal cognitive defects, spastic signs









SPECT: Significant improvement of perfusion

Case J. N.

JN, 21^a, male, AS, traumatic remission stage II-III HBOT: 64 sessions 1.5 - 1.75 ATA

Additional treatment: physiotherapy

Significant improvement Light defect symptoms: cerebellar, spastic symptoms, speech disturbances (pseudobulbar), cognitive deficits

Case M. A. B.

MAB, 89a, female, AS, traumatic, full stage

1.1 ATA (1 hour each session)

HBOT: 38 sessions, Treatment stopped because disallowed One week after start improvement: spasticity and encephalopathy reduced, mobilization possible, walking with help, speech understandable

SPECT Significant improvement of perfusion

care. Mild encephalopathy as rest state.

Improvement to significant amelioration of quality of life with good contact to surrounding and less demand of nursing

HBOT in Apallic Syndrome

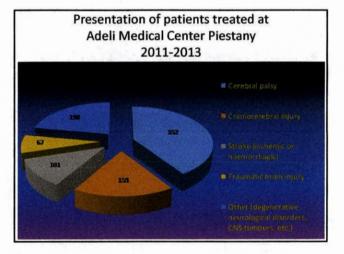
in full stage

- → as early as possible
- in remission stage with temporary interruption state II-III → after 3 months
- · in later remission stages and in defect stage without sufficient result of running rehabilitation → in any case

Continuation of classic treatment and consequent neurorehabilitation program

Clinical effects Apallic Syndrome and HBO-treatment:

reorganization of cortical connections to brain stem centers supported by the activation of reticular system
Revitalization of the cortical network (idling neurons)
Revitalization of damaged axons
Reactivation of dendrite system



Future aspects of HBOT in Neurology

- Acute conditions:
 - traumatic brain injury, stroke, encephalitis, spinal cord injury, neonatal hypoxic encephalopathy
- **Progredient conditions:**
- MS, progredient dementia of different origin, ALS
- Chronic states:
- Cerebral palsy
- Apallic syndrome
- Hypoxic encephalopathy
- Vascular dementia, mixed dementia
- Alzheimer dementia
- Traumatic spinal cord lesions

3. Summary of the 11th International Danube Teaching Course in Kazimierz (by Professor Konrad Rejdak, MD, PhD)

11th International Danube Teaching Course took place on 13-15 May 2015 in Kazimierz Dolny – historical centre of medieval culture of Poland.

The meeting was organised under the auspices of the International Danube Symposium for Continuing Education in Neurological Sciences, Department of Neurology Medical University of Lublin (Head: Prof dr K. Rejdak) and Lublin Branch of Polish Neurological Society (Prof dr Z. Stelmasiak).

The scientific programme focused on the newest aspects of multiple sclerosis, epilepsy, headache and child neurology but also included the special session on stem cell therapy. The highest educative and scientific level of this conference was guaranteed by the attendance of many internationally recognised experts and scientists from the country and abroad including prof. F. Gerstenbrand, prof. A. Korczyn, prof. G. Bauer, prof. P. Sorensen, prof. I. Blatt, prof M. Brazdil and others.

There were 7 Danube Lectures: by

Prof. F. Gerstenbrandt – Hyperbaric Oxygenation Therapy - a good possibility in neurorehabilitation, but not enough accepted and generally used

Prof. A. Korczyn – Medically unexplained symptoms

Prof. A. Słowik – Asymptomatic Carotid Artery Disease Update: Surgery, Stent or Watchful Waiting?

Prof. K. Selmaj – Future trend in MS therapy – the rule of generic drugs

Prof. P. S. Sørensen – Neutralizing antibodies to biological MS therapies

Prof. G. Bauer - Locked-in syndrome and related conditions

Prof. I. Blatt – Status Epilepticus

Prof. M. Brázdil - Epileptic seizures arising from the anterior cingulate cortex

Moreover the program included 6 scientific sessions: neuroprotection (3 lectures), multiple sclerosis (6 lectures), stroke and ultrasound exmination (5 lectures), child neurology (3 lectures) and epilepsy (9 lectures).

The meeting was very successful with around 450 participants including students and residents who were given a free admission to the scientific part of the meeting.



Photo: (from left) dr Szczepanska-Szerej, prof. K. Rejdak, prof. F. Gerstenbrandt, prof A. Korczyn, prof. Z. Stelmasiak, prof. P. Nowacki, prof. K. Mitosek-Szewczyk



International Danube Symposium for Neurological Sciences and Continuing Education

Lublin Branch of Polish Neurological Society

Department of Neurology Medical University of Lublin

XI Międzynarodowe Warsztaty Szkoleniowo-Naukowe

PROGRAM FINAL PROGRAMME



11th Danube Teaching Course

13 - 15 czerwca 2015, Kazimierz Dolny

www.danube2015.skolamed.pl