**Provisional programme, September 16th ,2017
FIMM/NVAMG 2017 International Scientific Spine Conference**

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| Saturday, September 16th, 2017 Scientific Spine Conference - Diagnostics, interventions and epidemiology – the latest insights  |
| Location: | **NH Hotel Utrecht Center** |  |
| Time | **Title** | **Speaker**  |
| 08.00-08.45 | Registration and Reception |   |
| 08.45-09.00 | Welcome - Henk Bultman and Nando Liem (NVAMG) | Henk Bultman and Nando Liem |
| 09.00-09.30 | [Teaching the Diagnosis of Pain- a workshop for teachers](#_Teaching_the_Diagnosis) | Simon Vulfsons |
| 09.30-10.00 | [Modic Changes - MRI](#_Modic_Changes_on) | Hugues Brat |
| 10.00-10.30 | Modic Changes - Clinical aspects | Marc-Henri Gauchat  |
| 10.30-11.00 | Coffee/tea Break |  |
| 11.00-11.30 | [Subgrouping low back pain: Modic changes](#_Subgrouping_low_back) | Michiel Schepers  |
| 11.30 - 12.00 | [An update on intradiscal procedures](#_An_update_on) | Michiel Schepers  |
| 12.00-13.15 | Lunch |   |
| 13.15-13.45 | Nociception | Wolfgang van Heymann |
| 13.45-14.15 | [The principles of Counter strain](#_The_Principles_of) | Mike Kuchera |
| 14.15-14.45 | Buffer session |  |
| 14.45-15.15 | Coffee/tea Break |  |
| 15.15-15.45 | Cervicogenic dizziness/ upper cervical spine | Mengemann Heinz  |
| 15.45-16.15 | [Clinical Trial and Evidence on the effectiveness and safety of Chuna manual medicine](#_Clinical_Trial_and) | Byung-Cheul Shin and Me-riong Kim                  |
| 16.15-16.45 | Epidemiological Outcomes Netherlands | Wouter Schuller |
| 16.45-17.15 | Epidemiological Outcomes Israel | Yacov Fogelman |
| 17.15-17.45 | Expert opinion-case presentation | Nando Liem and guest |
| 17.45-18.15 | Closing speaker - Vision for FIMM 2018 | Henk Bultman-French delegates |
| 18.15 | Drinks - informal reception / Freshen up for Social Program |  |
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| 19.00 | Social Program | Walking Dinner (3 course) optional |

## Teaching the Diagnosis of Pain–a workshop for teachers – Simon Vulfsons

The accurate diagnosis of pain is the cornerstone of successful treatment of any patient suffering from musculoskeletal pain.

As experienced therapists, we all find ways to practice diagnosis in the clinic but is there a good system for teaching this to our students  simply, quickly and enjoyably?

In this workshop, those who are interested in teaching students and junior physicians the art of musculoskeletal diagnosis will learn:

1. a simple heuristic for history taking and physical examination
2. A congruent pathophysiologic approach to the diagnosis of musculoskeletal pain
3. a simple tool for evaluating the bio-psych-social-spiritual involvement on the patient
4. and how to define a therapeutic plan

This workshop will be run by Dr. Simon Vulfsons, director of the Institute for Pain Medicine, Rambam Health Care Service, Haifa Israel

Past chairman ISMM (Israel Society for Musculoskeletal Medicine)
President FIMM (International Federation of Musculoskeletal Medicine)

## Modic Changes on MRI

Hugues Brat, MD – Groupe3R, Sion, Switzerland

Disc aging:

* Early disc degeneration (also known as chemical disc) is related to a proteolytic activity and generates pro-inflammatory cytokines (If, IL, Tnfa) stimulating nociceptors at the vertebral endplate and outer annular fibers.
* Late disc degeneration (known as mechanical disc) allows diffusion of chemical nociceptor stimulators through annular and endplate fissures, reaching centripetal neo-innervation with new pain receptors.

Modic on MRI:

MRI depicts Modic 1 changes such as inflammatory processes, erosions and reactive bone marrow edema as **hypo**intense endplate changes on T1 weighted images (T1w), **hyper**intense on T2w and **hyper**intense on T1w after fat suppression and i.v. Gadolinium injection.

Modic 2 changes are chronic erosions and bone marrow fatty transformation, seen as **hyper**intense endplate changes on T1w, **hyper**intense on T2w and **hypo**intense on T1w after fat suppression and i.v. Gadolinium injection.

Modic 3 changes are represented by osteosclerosis and **hypo**intense on all sequences.

There is **no continuum** from Modic 1 to 2 and from 2 to 3.

Clinical correlation:

Modic 1 changes are predictive of a symptomatic disc and extension is proportional to functional impairment (not to pain severity).

Mixed Modic 1 and 2 might be symptomatic as well, but less than pure Modic 1.

Imaging Treatment:

A target population with chronic spine pain, Modic 1 changes, epidural steroid infiltration with partial effect and a positive discography for a chemical disc, might benefit from an intradiscal steroid injection in order to sedate pain at 1 month.

Modic 1 Differential Diagnosis:

Acute annular tear with peridiscal inflammation might mimic Modic 1 changes.

Early ankylosing spondylitis is a common pitfall.

Severe facet joint inflammatory rash can extend to posterior aspect of endplates.

Conclusion: Modic 1 changes are

* related to degeneration,
* indicate endplate edema and inflammatory changes,
* are painful,
* limit function in proportion to their extend,
* should not be mistaken with AS,
* can sometimes be sedated with intradiscal steroids.

## Subgrouping low back pain: Modic changes

In low back pain more and more subgroups can be identified.
One of these is painfull endplate edema called modic type 1 and Modic type 2 changes as dseen on the MRI .
Recent studies have shown that there are treatment options which seem promising for this specific population.
These treatments will be discussed.

Michiel Schepers MD
Musculoskeletal medicine and pain management
Senior instructor S.I.S (Spine intervention society)

## An update on intradiscal procedures

And overview will be given on the current minimal invasive treatments for discogenic low back pain.
Rationale and evidence or lack of evidence will be discussed.

Michiel Schepers MD
Musculoskeletal medicine and pain management
Senior instructor S.I.S (Spine intervention society)

## The Principles of Counterstrain

(Prof. Dr. Michael L. Kuchera)

Counterstrain is an indirect form of manual medicine technique that involves:

* Diagnosis of somatic dysfunction including identification of the most tender point within the tissue texture abnormality to be treated.
* A therapeutic component described as “spontaneous release by positioning” in which the body is positioned in such a fashion that pressure tenderness over the point is reduced by at least 70%.

The diagnosed somatic dysfunction is considered to be the result of a continuing, inappropriate strain neuroreflex that can be inhibited or reversed by maintaining a position of mild strain in the direction exactly opposite that of the false strain reflex for an appropriate period of time (usually 90 seconds) … thus “counterstrain.” Counterstrain technique is easily taught and its principles are readily transferable to most regions and tissues of the musculoskeletal system.

New research has documented that in addition to reducing pain and reestablishing normal joint and muscle function, counterstrain may also be useful in reducing edema and producing a mechanotransduction mediated reduction in pre-inflammatory interleukins and apoptosis.

Because of the underlying mechanisms-of-action proposed or documented for counterstrain and because by definition it is administered by positioning the passive patient away from pain and tissue tension, this manual technique is particularly useful in treating acutely painful conditions, reducing or eliminating myofascial trigger points, and/or for use in the emergency department or hospitalized settings.

Exemplars will be provided in this presentation and a practical workshop demonstrating the ease and efficacy of counterstrain will be presented on Sunday, September 17.

## Clinical Trial and Evidence on the effectiveness and safety of Chuna manual medicine

Byung-Cheul Shin 1,2, Kyeong-Tae Lim 1,3, , Eui-Hyoung Hwang 1,2, In Heo 1,3, Byung-Jun Kim 1,3

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***Purpose:***

A research project (June, 2015 ~ May, 2018) is ongoing for creating clinical evidence of Chuna manual medicine (CMM) in Korea. The aim of this presentation is to introduce results from the research project which contains evidence from systematic reviews and clinical trials from a randomized controlled trial (RCT) on CMM for non-acute low back pain (LBP).

***Methods/Design:***

We conducted systematic reviews on the effectiveness and safety of CMM for various kinds of conditions. Especially, musculoskeletal disorders are analyzed more in depth. Additionally, a pilot, three-armed, multicenter, pragmatic randomized controlled pilot trial (n=60; Chuna group (n=20), usual care group (n=20), or Chuna plus usual care group (n=20) for 6 weeks of treatment) was conducted in 2016 for exploring the feasibility of main clinical trial. From the pilot randomized trial, then we analyzed powered sample size (n=194) and are conducting a confirmative, pragmatic, multi-centered, randomized controlled clinical trial from 2017 to 2018. The primary outcome was numerical rating scale (NRS) of LBP, Secondary outcomes include NRS of leg pain, Oswestry disability index (ODI) and lumbar range of motion (ROM).

***Results:***

Based on the systematic review on the effectiveness of CMM, 28 systematic reviews and 778 RCTs of CMM for various conditions were found. Of them, about 120 RCTs were related on musculoskeletal conditions. For safety, 47 adverse reactions from 78 literatures were detected. For pilot trial, total 60 patients were included in the intention-to-treat analysis and there were significant differences (P<0.01) in NRS scores of each group (Chuna ; -3.28±1.73, UC ; -1.95±1.85, Chuna+UC -1.75±2.02). Based on pilot trial, we estimated 194 were powered sample size for main trial. A confirmative, pragmatic RCT is ongoing (118/194 were randomly allocated, 60.8%) and will be finished May, 2018.

***Conclusions:***

Based on our systematic reviews, musculoskeletal conditions are most clinically researched area and CMM showed an improvement on pain and function, but quality of evidence is low. Though safety is generally good, however several serious adverse events were reported. Through our clinical trial results, we suggest that CMM might have a comparative effectiveness on non-acute LBP. A large, well-designed main study based on pilot results will be finalized in 2018.

***Key words;*** Clinical Trial, Evidence, Chuna manual medicine, systematic review, effectiveness, safety, randomized controlled trial