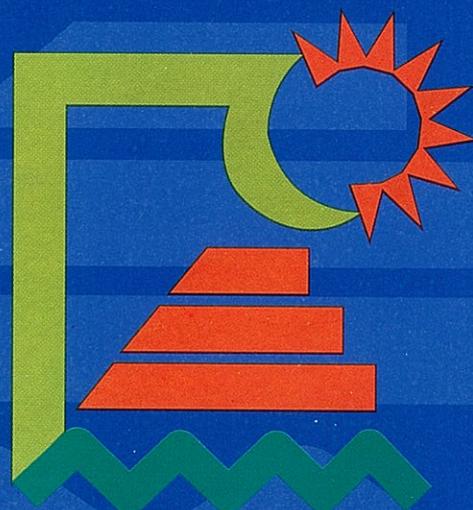


ISSN 0186-4866

Medicina Interna de México

REVISTA DE LA ASOCIACIÓN DE MEDICINA INTERNA DE MÉXICO



Volumen 16
Suplemento, 2000

Cancun



25TH World Congress of INTERNAL MEDICINE



included complete cell count, platelet count, urinalysis, chemistry panel, renal and pharyngeal culture. **Results:** All the patients were female, the age was between 16 – 36 years (mean 24.6 ± 8.05 years), The basal and final results were creatinine clearance 23.78 ± 8.9 ml/min. vs 78.6 ± 8.9 ml/min ($p < 0.001$), twenty hours proteinuria 9.46 ± 0.44 gr./lt. vs 0.90 ± 0.44 gr./lt. ($p < 0.001$), plasmatic urea 76.4 ± 11.3 mg/dl vs. 25.8 ± 11.3 gr./dl ($p < 0.05$), plasma creatinine 3.76 ± 0.49 gr./dl vs 1.12 ± 0.49 mg/dl ($p < 0.01$). The more frequently complications were infectious disease of viral (1 case) and bacterial (3 cases) etiology. We didn't see complications and decease. **Conclusions:** The intravenous cyclofosfamide pulse improve the renal function in patients with diffuse proliferative nephritis and it could be related to non toxic effect. Its application could be not require hospitalization.

EFFECTS OF KAMPO MEDICINES ON TH1/TH2 BALANCE. Kenji Watanabe, Tsutomu Nakada, Kazuki Santa and Toshihiko Hanawa. Oriental Medicine Research Center, Kitasato Institute, Tokyo, Japan.

Introduction Kampo Medicine is a Japanese herbal medicine originated from China. It has become popular and common in Japan in these days. More than 70% of internal medical doctors use Kampo medicine in their daily clinical work in Japan, because Kampo medicine is much more effective for several diseases like allergic diseases, auto-immune diseases. However the mechanism of Kampo medicine is not completely understood. For allergic diseases, Th1/Th2 balance might be one of the candidates of mechanism. **Purpose** The purpose of this study is to clarify the effect of Kampo medicines on Th1/Th2 balance in mice and human. **Methods** 1) Several Kampo medicines were administered to C57BL/6 mice and Balb/c mice for two weeks. To see the change of the subpopulations of lymphocytes, spleen cells were stained by monoclonal antibodies and analyzed by flow cytometer. Spleen cells were cultured together with anti-CD3 antibody or LPS to detect IFN-g and IL-4, or IL-12 respectively. 2) Two Kampo medicines (Juzentaiho-to and Shosaiko-to) were administered to the healthy volunteers for two weeks. Peripheral blood mononuclear cells were isolated before and after the administration of Kampo medicines, and submitted for immuno-staining to detect the change of lymphocyte subpopulations and also submitted to detect the cytokine secretion. **Results** 1) Among several Kampo medicines Gorei-san and Saireito shifted Th1/Th2 balance to Th1 side in C57BL/6 mice, while Hangekoboku-to stimulated Th1 cells in Balb/c mice. 2) In human, both Juzentaiho-to and Shosaiko-to stimulated the secretion of IFN-g and IL-4. **Conclusion** The results obtained in these studies indicated that some Kampo medicines have a potentiality to change Th1/Th2 balance in both mice and human, and this change may explain the effectiveness of Kampo medicines.

EFFECT ON LDL SUBCLASS DISTRIBUTION OF HORMONAL REPLACEMENT THERAPY IN POSTMENOPAUSAL WOMEN WITH TYPE 2 DIABETES UNDER SATISFACTORY AND NON SATISFACTORY METABOLIC CONTROL. Leobardo Sauque, Carlos A Aguilar Salinas, Onix Arita, Angelina Lòpez, Ma. Luisa Velasco, Luz E. Guillén, Francisco Gómez Pérez, Juan A Rull. Clínica de Diabetes y Metabolismo de Lípidos, Instituto Nacional de la Nutrición Salvador Zubirán, México, DF.

Objective: There is a lack of information about the effects of hormonal replacement therapy (HRT) in women with diabetes, especially during moderate chronic hyperglycemia. Our objectives were to study the effects of HRT on the lipid profile and LDL subclass distribution in women with type 2 diabetes under satisfactory and non satisfactory metabolic control. **Material and methods:** 54 postmenopausal women were included. 24 had a HbA1C $< 8\%$ during at least 3 months before the beginning of the study; the remaining 30 had HbA1C $> 8\%$. After a 6-week run in period with diet, cases were randomized to receive either placebo (HbA1C $< 8 = 13$, HbA1C $> 8 = 17$) or HRT (HbA1C $< 8 = 11$, HbA1C $> 8 = 13$) during 12 weeks. HRT consisted in conjugated estrogens 0.625mg/d PO plus medrogestrone 5mg/d during the last ten days of the cycle. At the beginning and the end of each treatment period, the LDL subclass distribution was estimated by density gradient ultracentrifugation. **Results:** At the baseline and during the study hyperglycemic cases had HbA1C level significantly higher than the near-normoglycemic controls (baseline 10.2 ± 2.9 vs $6.5 \pm 0.7\%$, $p < 0.01$). They had a trend for higher total and LDL cholesterol, triglycerides and lower HDL-cholesterol in comparison to near-normoglycemic controls. Also they had significantly higher triglycerides concentrations in VLDL, IDL, and LDL-1 particles and cholesterol content in LDL-1 and -2 particles. HRT decreased LDL-cholesterol in both groups (150 ± 36 vs 125 ± 45 and 144 ± 31 vs 124 ± 41 mg/dl respectively, $p > 0.05$). In normoglycemic cases a small increase in HbA1C was observed (6.5 ± 0.7 vs $7.4 \pm 1\%$, $p = 0.04$). HRT significantly increased plasma triglycerides and the triglycerides content of the all apoB lipoproteins containing particles, specially in the hyperglycemic cases. In all cases HRT did not modify the proportion of LDL represented by denser LDL's. **Conclusion:** HRT did not modify the LDL subclass distribution, even in the presence of moderate chronic hyperglycemia in women with type 2 diabetes. HRT resulted in small increase in HbA1C in the near normoglycemic cases, but it did not affect glycemic control in hyperglycemic subjects. The HRT-induced triglycerides elevation was significantly larger in hyperglycemic cases.

TREATMENT OF SINOVITIS RESISTANT IN RHEUMATIC DISEASES TO OTHER THERAPIES WITH METOTREXATE (MTX) INTRAARTICULAR (IAA). Leónides Cuapio M, D. Bañuelos Rmz, Jorge Rojas Rodríguez. Hospital de Especialidades CMN MAC Puebla, Pue., México.

