

See discussions, stats, and author profiles for this publication at: <https://www.researchgate.net/publication/263816135>

Complementary Treatment of Obesity and Overweight with Salacia Reticulata and Vitamin D

Article in *International Journal for Vitamin and Nutrition Research* · October 2013

DOI: 10.1024/0300-9831/a000162 · Source: PubMed

CITATIONS

8

READS

486

4 authors:



Michael Ofner

MEDYCO International

39 PUBLICATIONS 297 CITATIONS

[SEE PROFILE](#)



Andreas Tomaschitz

Medical University of Graz & Specialist Clinic for Rehabilitation Bad Gleichenberg, ...

219 PUBLICATIONS 9,358 CITATIONS

[SEE PROFILE](#)



Manfred Wonisch

Hartmannspital Vienna and Private Practice-Sports Cardiology

169 PUBLICATIONS 1,445 CITATIONS

[SEE PROFILE](#)



Gerhard Litscher

Medical University of Graz

682 PUBLICATIONS 5,410 CITATIONS

[SEE PROFILE](#)

Some of the authors of this publication are also working on these related projects:



Thermology and Complementary Medicine [View project](#)



The International Symposium on Auriculotherapy AUG 10-12, 2017 SINGAPORE [View project](#)

International Journal for Vitamin and Nutrition Research

4/13

www.verlag-hanshuber.com/IJVNR

Editor-in-Chief

R. F. Hurrell

Associate Editors

T. Bohn · M. Eggersdorfer · M. Reddy

HUBER



Contents

Original Communications	Nissensohn M., Sánchez-Villegas A., Fuentes Lugo D., Henríquez Sánchez P., Doreste Alonso J., Skinner A. L., Warthon Medina M., Lowe N. L., Hall Moran V., and Serra-Majem L.: Effect of Zinc Intake on Mental and Motor Development in Infants: A Meta-Analysis	203
	Ofner M., Tomaschitz A., Wonisch M., and Litscher G.: Complementary Treatment of Obesity and Overweight with <i>Salacia Reticulata</i> and Vitamin D	216
	Root M. M. and Dawson H. R.: DASH-Like Diets High in Protein or Monounsaturated Fats Improve Metabolic Syndrome and Calculated Vascular Risk.	224
	Ataseven, F., Aygün C., Okuyucu A., Bedir A., Küçük Y., and Küçüködük, S.: Is Vitamin D Deficiency a Risk Factor for Respiratory Distress Syndrome? . .	232
	Faizan M., Stubhaug I., Menoyo D., Esatbeyoglu T., Wagner A. E., Struksnæs G., Koppe W., and Rimbach G.: Dietary Alpha-Tocopherol Affects Tissue Vitamin E and Malondialdehyde Levels but Does not Change Antioxidant Enzymes and Fatty Acid Composition in Farmed Atlantic Salmon (<i>Salmo salar</i> L.)	238
	Roncero-Ramos I., Pastoriza S., Navarro M. P., and Delgado-Andrade, C.: Consumption of Model Maillard Reaction Products has no Significant Impact on Ca and Mg Retention or on Tissue Distribution in Rats	246
Research Note	Chevallereau G., Gleyses X., Roussel L., Hamdan S., Beauchet O., and Annweiler C.: Proposal and Validation of a Quick Question to Rate the Influence of Diet in Geriatric Epidemiological Studies on Vitamin D	254

International Journal for **Vitamin and Nutrition Research**

Your article has appeared in a journal published by Hans Huber Publishers.

This e-offprint is provided exclusively for the personal use of the authors.

It may not be posted on a personal or institutional website or to an
institutional or disciplinary repository.

If you wish to post the article to your personal or institutional website or to
archive it in an institutional or disciplinary repository, please use either a pre-
print or a post-print of your manuscript in accordance with the publication
release for your article and our “Online Rights for Journal Articles”
(<http://www.verlag-hanshuber.com/informationen>).

HUBER



Original Communication

Complementary Treatment of Obesity and Overweight with *Salacia Reticulata* and Vitamin D

Michael Ofner¹, Andreas Tomaschitz², Manfred Wonisch³,
and Gerhard Litscher⁴

¹Department of Sports Physiology, University of Vienna, Austria

²Department of Cardiology, Medical University of Graz, Austria

³Department of Cardiology, Private Hospital Hansa Graz, Austria

⁴Stronach Research Unit for Complementary and Integrative Laser Medicine, Research Unit of Biomedical Engineering in Anesthesia and Intensive Care Medicine, and TCM Research Center Graz, Medical University of Graz, Austria

Received: May 21, 2013; Accepted: October 3, 2013

Abstract: *Background:* The Indian plant root *Salacia reticulata*, which is rich in alpha-glucosidase inhibitors, is used for metabolic disorders in Ayurvedic medicine. Vitamin D₃ is also used in the treatment of some metabolic diseases. Our goal was to determine its potential effect for humans with obesity. *Material:* In a randomized open-label study, we investigated 40 healthy participants aged 30–60 years, physically active, with a body mass index (BMI) of 25–45. The participants were randomly allocated into two groups. Body weight, BMI, and body composition were measured. Both groups (A and B) received a guideline for lifestyle and fitness training for 4 weeks. Group B additionally took one capsule containing 200 mg of *Salacia reticulata* and 1.6 µg (i. e. 64 IU) Vitamin D₃ (SRD) 3 times/day with the meals. *Results:* Significant weight and body-fat reduction within 4 weeks was observed. Group A lost 1.8 kg or 2.1 %, group B lost 5.3 kg or 6.1 % (p=0.03), therefore BMI reduction was achieved. While Group A lost 1.4 % of body fat, group B reduced it by 4.5 % (p=0.01). *Conclusion:* These promising results suggest that the combination of *Salacia reticulata* and Vitamin D₃ might be highly valuable and potent to treat overweight and obesity, especially in addition to a modifying lifestyle program. Further research is needed in addition to this study to clarify pathways and effect mechanisms.

Key words: obesity, vitamin D, *Salacia reticulata*, overweight, exadipin, lifestyle, training

Introduction

Salacia reticulata is an Indian plant root and has been used for treatment of diabetes, metabolic disorders, and obesity for hundreds of years, especially in Ayurvedic medicine. The known active ingredients of the root are alpha-glucosidase inhibitors, salacinol,

kotalanol, thiocylitol, and mangiferin [1, 5]. Multiple effects in the metabolism of glucose have been shown in animal trials. For example, the incidence and the risks of diabetes decreased in these studies, as well as levels of blood glucose, due to the reduction of intestinal glucose absorption and the regulation of reactive insulin secretion [6, 13]. Further positive effects on

lipid metabolism were reported, such as reduction of fat accumulation and metabolic disease prevention [14–17]). The incidence and prevalence of metabolic syndrome were decreased, and a significant reduction in body weight could be shown in animal trials [9, 14–17]). *Salacia reticulata* suppresses the development of fat cells [1]. It was reported that the root might be effective in the treatment of certain metabolic diseases [14, 18]. One human study was published concluding that *Salacia reticulata* was able to significantly reduce glycosylated hemoglobin (HbA1c) and thus might be an option for the treatment of type II diabetes [10]. In this study, patients received verum Ayurvedic treatment with a tea containing an unspecified amount of *Salacia reticulata* for three months, randomly preceded or followed by either three months' placebo treatment (tea without *Salacia reticulata*) [10]. The use of the root seems to be safe in animal trials. No skin reactions, teratogenicity, or hepatopathology have been reported [19–22].

Vitamin D₃ (cholecalciferol) has become of more interest in the last years, resulting in over 20,000 publications available in PubMed (as of September 2013). Among other beneficial influences, it was reported that serum triglyceride concentrations as well as HbA1c levels decreased significantly following vitamin D₃ treatment, compared to placebo [23, 24]. The underlying mechanisms are still not clear; some theorize that the mechanisms are the regulation of beta-cell activity and the production of anti-inflammatory cytokines which protect beta cells from destruction. There is still some controversy as reported by Seshari *et al.* [25].

The present study investigated in an open-label trial the combined effect of *Salacia reticulata* and Vitamin D₃ on body weight and body fat during a 4-week intervention period.

These two substances seem to provide multiple positive effects for the treatment of metabolic disorders, but so far no study has evaluated the combination of *Salacia reticulata* and vitamin D₃ in the treatment of overweight, obesity, and metabolic disorders in humans.

Methods

The combination of *Salacia reticulata* and vitamin D₃ (SRD) is available as Exadipin® (AapoSpa, Austria), which was used for our study. It is licensed as a supplement for special medical treatments, and contains 200 mg *Salacia reticulata* root powder (with 60 mg saponins) as well as 1.6 µg vitamin D₃ (i. e. 64 IU) per

capsule. The recommendation is to take it with every meal, at least 3 times a day.

The participants were invited and actively asked to participate in the study in fitness centers in the federal state of Styria, Austria. All participants were informed about the study when they joined the fitness training club with the intention to lose weight. The participants received the necessary information for the study in a comprehensible written form. All the participants confirmed that they agreed with the procedure (informed consent) before examination and treatment. The study was approved by the local ethics committee (IRB).

Inclusion Criteria

- Healthy participants without ongoing drug treatment
- Overweight or obesity [body mass index (BMI) 25–45]
- Ability to do sports
- Age: 30–60
- Body composition: obese, not muscular

Exclusion Criteria

- Diabetic subjects or subjects suffering from cardiovascular diseases
- Taking any medication or additional dietary supplements

In total 40 participants (8 men, 32 women) of Caucasian ethnicity were included in 2 groups (20 participants each). The ratio of male and female participants did not differ in the two groups.

Assessment

After inclusion in the study, the participants were randomized into 2 groups via an online tool (www.randomizer.org). Each participant was examined by a professional sports scientist. The following parameters were obtained and measured at the start of the trial (in the fasting state in the morning by the same investigator):

- age and sex
- body weight (with a calibrated scale (Soehnle 7730, Leifheit AG, Nassau, Germany))
- height (biometric)
- BMI (calculated)
- body composition [impedance measurement with Soehnle 7850 (Leifheit AG, Nassau, Germany)]

Both groups (A and B) received a guideline for lifestyle, which includes the following basic recommendations (and additional explanations):

- Take care **WHAT** you buy in supermarkets
- Use your knowledge about nutrition and lifestyle – be **SMART**
- Don't drink **ALCOHOL** and sugary soft-drinks
- Don't life ascetically, but **SIN** rarely
- **REDUCE** carbohydrates and fat
- Eat food only in highest **QUALITY**
- Don't **LIE** to yourself
- Do sports and do it with **FUN**
- Control your body-**WEIGHT** continuously

Additionally, participants received a standardized fitness training plan consisting of twice-weekly muscle-strengthening and aerobic cardio training programs (each session 45 minutes), which were supervised by a professional trainer. Group B additionally took one capsule of SRD 3 times a day with meals.

The participants performed the exercise with the guidance of professional fitness trainers and nutrition specialists. After a period of 4 weeks all parameters mentioned above were measured again.

Additionally each participant completed a questionnaire addressing how strictly the guidelines (incl. SRD intake) had been adhered to by answering one question on a spectrum from 0 (= nothing adhered to) to 10 (= everything adhered to).

Statistical Methods and Analysis

Microsoft Excel 2007 was used for data collection and SPSS 16 was used for data analysis.

First, the Kolmogorov-Smirnov Test was used for evaluation if the data set was normally distributed. The two-sided unpaired *t*-test was used for unpaired groups, and the paired *t*-test for paired groups. The level of significance was set at $p \leq 0.05$. Analysis was performed assuming treat intention to treat.

Results

The mean and standard deviation (SD) of different variables were calculated for the 40 participants (Table I). The age range of the participants in both groups A and B was 30–60 years; height ranged from 154 to 182 cm in both groups A and B, as well. At the time of inclusion, the weight range of group A was 64.3–123.7 kg, and 63.4–134.4 kg in group B. The BMI of group A ranged from 25.0–43.8 kg/m², and from 25.2–43.9 kg/m² in group B. The body fat mass showed a range of 17.0–54.1 % in group A, and 30.3–46.5 % in group B.

After 4 weeks of intervention, the parameters had changed as follows. The mean body weight was 83.7 kg (63.3–122.9 kg) in group A, representing a decrease of 1.8 kg or 2.1 %. The mean body weight of group B was 81.7 kg (60.9–124.6 kg), which corresponds to a reduction of 5.3 kg or 6.1 %. The difference of means was 3.5 kg, which was significantly higher ($p = 0.03$) compared to control group (Figure 1).

The mean BMI in group A was 29.4 kg/m² (22.4–43.5 kg/m²), signifying a decrease of 0.6 kg/m² and a relative loss of 2.0 %. The mean BMI was 29.3 kg/m² (22.8–40.7 kg/m²) in group B, corresponding to a decrease of 1.9 kg/m² and a relative loss of 6.1 %.

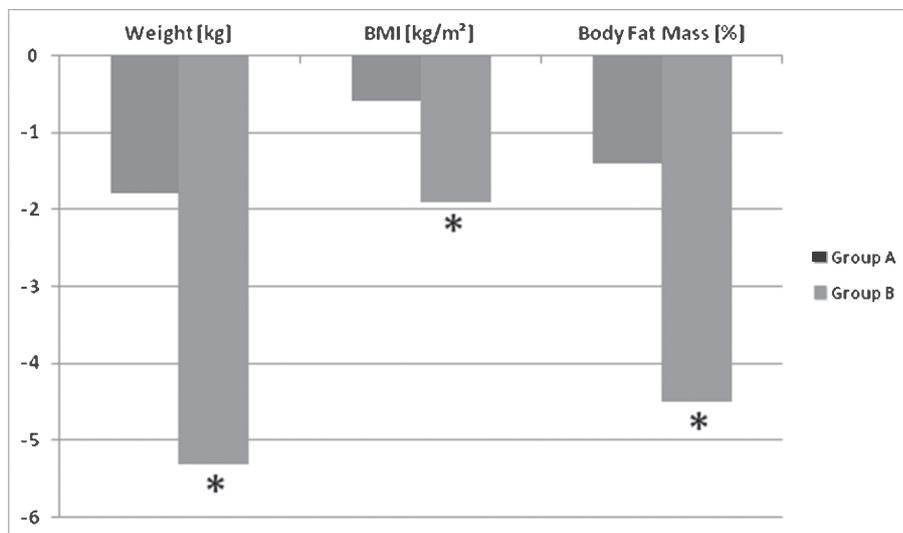


Figure 1: Weight-, BMI-, and fat loss of both groups. This chart shows the lost weight [kg], BMI [kg/m²], and body fat mass [%] of both groups within 4 weeks. Group A received lifestyle intervention, group B received in addition one capsule of a combination of Salacia reticulata and low-dose Vitamin D with every meal. * $p < 0.05$.

Table I: Mean and standard deviation of measured parameters of both groups. Group A received lifestyle intervention for 4 weeks, group B received additionally one capsule of a combination of *Salacia reticulata* (200 mg) and low-dose vitamin D (1.6 µg, i.e. 64 IU) with every meal. Date 1 was at inclusion, date 2 was 4 weeks later.

Variable	Group A		Group B	
	mean	SD	mean	SD
Examination Date 1				
Age [years]	46.8	9.4	48.1	9.6
Height [cm]	169.2	7.7	167.9	6.3
Weight [kg]	85.5	8.6	87.0	8.8
BMI [kg/m ²]	30.0	5.3	31.2	4.7
Body Fat mass [%]	34.8	9.7	36.1	5.6
Examination Date 2				
Weight [kg]	83.7	8.9	81.7	8.5
BMI [kg/m ²]	29.4	9.7	29.3	4.5
Body Fat mass [%]	33.4	5.4	31.6	5.2

Mean difference between both groups was 1.3 kg/m² ($p=0.02$), in favor of group B.

The mean percentage of body fat mass was 33.4 % (15.6–52.9) in group A, which signified a reduction of 1.4 % and a relative loss of 4.0 %. The mean body fat mass in group B was 31.6 % (20.4–41.0 %), representing a decrease of 4.5 % and a relative loss of 12.5 %. A mean difference between both groups of 3.1 % ($p=0.01$) was calculated.

The questionnaire (how strictly the guideline was adhered to) showed a mean value of 7.05 (range 4–8) in group A, compared to 7.15 (4–9; $p=0.76$) in group B.

Discussion

The scientific interest in Vitamin D₃ has grown in recent years. Cholecalciferol is a natural substance produced in the human body from 7-dehydrocholesterol by interaction with ultraviolet (UV), especially UVB, light in the skin. Vitamin D₃ can also be taken orally with nutrition or as nutrition supplements. Adults (over 70 years of age) are given the recommendation (DGE) to take 20 µg/day, which corresponds to 800 IU (International Units). No side effects were reported up to a daily intake of about 4000 IU [26]. A deficiency or under-supply of Vitamin D₃ can be a risk factor for several diseases: autoimmune diseases like multiple sclerosis [27], Crohn's disease [28], diabetes mellitus [29], infections like tuberculosis (TB) or infections of the respiratory system [30–32], hypertension [33], different kinds of cancer [34–36], osteopenia and osteo-

porosis, cardiovascular diseases [29, 37, 38], metabolic syndrome [29], muscle disorders [39], reduced brain function, and rachitis (rickets) [40]. This deficiency is suggested to occur because of UVB deficiency and/or unbalanced nutrition containing too small a Vitamin D₃ concentration. The endogenous production of Vitamin D₃ in the body also depends on skin pigmentation: the more pigmented the skin, the less efficient the transformation of the pre-vitamin to Vitamin D₃. Vitamin D₃ supplementation is often recommended in the winter months, as well as for dark-skinned people. In some studies the prevalence of Vitamin D deficiency was over 78.2 % [41, 42].

Improvement of vitamin D₃ status resulted in the reduction of inflammatory markers, such as serum highly-sensitive C-reactive protein (hsCRP), serum amyloid A (SAA), interleukin (IL)-2, IL-6, IL-10, and tumor necrosis factor (TNF)- α , and diabetic complications in type 2 diabetes patients [43]. Some data suggest a potential role for Vitamin D₃ in the treatment of diabetes-associated hepatic complications [44]. Nazarian *et al.* indicated that orally administered high-dose Vitamin D₃ supplementation improves insulin sensitivity in subjects with impaired fasting glucose and suggested that high-dose Vitamin D₃ supplementation might provide an inexpensive public health measure in preventing, or at least delaying, the progression from impaired fasting glucose to diabetes [45]. Other studies found that serum triglyceride concentrations as well as HbA1c decreased significantly under Vitamin D₃ treatment compared to placebo [23, 24].

Finally, McGill showed clearly that serum Vitamin D₃ is inversely related to weight, BMI, and markers of type II diabetes mellitus (TIIDM) (large waist,

raised HbA1c) [46]. However, a study dealing with genetic association concluded that lower 25-hydroxyvitamin D is likely to increase BMI only to a small extent [46]. Other studies support the assumption that Vitamin D supplements do not influence body weight, at least in adult obese individuals [47, 48].

During the study participants took no permanent drugs or medications. Age, height, and weight were similar in both groups, which led to homogenous and comparable initial conditions. An average weight loss of more than 5 kilograms in 4 weeks was a big success for the participating members taking SRD. The difference of weight loss in groups A and B was significant, suggesting that SRD might be a potent agent of treating overweight and obesity.

Even more interesting for us was the fact that significantly more body fat (in total more than 4.5 %) was lost in the group supplemented with SRD compared to the control group. We assume that a combination of exercise and nutrition is still the best formula for losing fat, and that this might be improved by an oral intake of SRD.

No significant difference could be observed in the means of the questionnaire scale. Probably both groups took similar nutrition and did comparable exercise.

Akase *et al.* and Shimada *et al.* found that *Salacia reticulata* seems to be a potent substance for treating metabolic diseases, especially overweight and obesity, but its combination with Vitamin D₃ leads to better results [14, 17]. In literature, the reasons for these effects are multifactorial. The ingredients (alpha-glucosidase inhibitors, salacinol, kotalanol, thiocyclitol, and mangiferin + Vitamin D₃) are thought to interact with different enzymes and to block uptake mechanisms of glucose and fat at the small intestine as well as into cells [6, 14, 15, 17]. These interesting findings might be relevant for future metabolic guidelines if further studies were to support the data. Overall, this might be a very potent combination for future metabolic treatments.

The limitations of the study should be mentioned briefly as follows. A placebo supplement would have been an option instead of just training (control group). In addition, no blood laboratory parameters (glucose, lipids, vitamin D, etc.) were measured. These might provide more support for a clear conclusion. However, our goal was to get an impression as to whether the supplement might have effects on overweight and obesity in humans, and not just in animal models, as reported previously by other authors.

As a follow-up study, a randomized double-blind, placebo-controlled clinical trial is planned in order

to obtain clear data on metabolic pathways, effects on blood sugar and fat profiles, anti-oxidation and anti-inflammation indicators, and Vitamin D serum levels. Of course, a larger sample size, a prolonged duration of intervention, and an increase of the Vitamin D dose could also be taken into account in further studies.

Conclusion

Salacia reticulata and Vitamin D₃ have been well studied individually for the treatment of metabolic diseases. Especially for Vitamin D₃, there are enough data to recommend the supplementation of this agent to treat common metabolic diseases of human beings. *Salacia reticulata* is an Indian plant root, which has been used for hundreds of years in Ayurvedic medicine for treating of those diseases as well, but up until now research has been focused mainly on animal trials.

Our data suggest that the combination of *Salacia reticulata* and Vitamin D₃ might be highly valuable and potent for the treatment of overweight and obesity, especially in addition to a lifestyle modification program focused on nutrition and exercise. The findings encourage further research to clarify the pathway and effectiveness of *Salacia reticulata* in the treatment of obesity, diabetes, and other metabolic diseases.

Acknowledgements

We would like to thank the staff of the Fitnessstudios Lifestyle in Styria for recruiting participants.

Conflicts of Interest

No potential conflict of interest relevant to this article is reported. The authors declare no competing financial interests. The authors assume responsibility for the overall content and integrity of this article.

References

1. Shimada, T., Nagai, E., Harasawa, Y., Watanabe, M., Negishi, K., Akase, T., Sai, Y., Miyamoto, K. and Aburada, M. (2011) *Salacia reticulata* inhibits differentiation of 3T3-L1 adipocytes. *J. Ethnopharmacol.* 136 (1), 67–74.

2. Eskandari, R., Kuntz, D.A., Rose, D.R. and Pinto, B.M. (2010) Potent glucosidase inhibitors: de-O-sulfonated ponkoranol and its stereoisomer. *Org. Lett.* 12 (7), 1632–1635.
3. Eskandari, R., Jones, K., Rose, D.R. and Pinto, B.M. (2010) Probing the active-site requirements of human intestinal N-terminal maltase glucoamylase: the effect of replacing the sulfate moiety by a methyl ether in ponkoranol, a naturally occurring alpha-glucosidase inhibitor. *Bioorg. Med. Chem. Lett.* 20 (19), 5686–5689.
4. Im, R., Mano, H., Matsuura, T., Nakatani, S., Shimizu, J. and Wada, M. (2009) Mechanisms of blood glucose-lowering effect of aqueous extract from stems of Kothala himbutu (*Salacia reticulata*) in the mouse. *J. Ethnopharmacol.* 121 (2), 234–240.
5. Ozaki, S., Oe, H. and Kitamura, S. (2008) Alpha-glucosidase inhibitor from Kothala-himbutu (*Salacia reticulata* WIGHT). *J. Nat. Prod.* 71 (6), 981–984.
6. Matsuura, T., Yoshikawa, Y., Masui, H. and Sano, M. (2004) [Suppression of glucose absorption by various health teas in rats]. *Yakugaku Zasshi* 124 (4), 217–223.
7. Collene, A.L., Hertzler, S.R., Williams, J.A. and Wolf, B.W. (2005) Effects of a nutritional supplement containing *Salacia oblonga* extract and insulinogenic amino acids on postprandial glycemia, insulinemia, and breath hydrogen responses in healthy adults. *Nutrition* 21 (7–8), 848–854.
8. Heacock, P.M., Hertzler, S.R., Williams, J.A. and Wolf, B.W. (2005) Effects of a medical food containing an herbal alpha-glucosidase inhibitor on postprandial glycemia and insulinemia in healthy adults. *J. Am. Diet. Assoc.* 105 (1), 65–71.
9. Huang, T.H., He, L., Qin, Q., Yang, Q., Peng, G., Harada, M., Qi, Y., Yamahara, J., Roufogalis, B.D. and Li, Y. (2008) *Salacia oblonga* root decreases cardiac hypertrophy in Zucker diabetic fatty rats: inhibition of cardiac expression of angiotensin II type 1 receptor. *Diabetes Obes. Metab.* 10 (7), 574–585.
10. Jayawardena, M.H., de Alwis, N.M., Hettigoda, V. and Fernando, D.J. (2005) A double blind randomised placebo controlled cross over study of a herbal preparation containing *Salacia reticulata* in the treatment of type 2 diabetes. *J. Ethnopharmacol.* 97 (2), 215–218.
11. Li, Y., Huang, T.H. and Yamahara, J. (2008) *Salacia* root, a unique Ayurvedic medicine, meets multiple targets in diabetes and obesity. *Life Sci.* 82 (21–22), 1045–1049.
12. Williams, J.A., Choe, Y.S., Noss, M.J., Baumgartner, C.J. and Mustad, V.A. (2007) Extract of *Salacia oblonga* lowers acute glycemia in patients with type 2 diabetes. *Am. J. Clin. Nutr.* 86 (1), 124–130.
13. Yoshino, K., Miyauchi, Y., Kanetaka, T., Takagi, Y. and Koga, K. (2009) Anti-diabetic activity of a leaf extract prepared from *Salacia reticulata* in mice. *Biosci. Biotechnol. Biochem.* 73 (5), 1096–1104.
14. Akase, T., Shimada, T., Harasawa, Y., Ikeya, Y., Nagai, E., Iizuka, S., Nakagami, G., Iizuka, S., Sanada, H., and Aburada, M. (2009) Preventive Effects of *Salacia reticulata* on Obesity and Metabolic Disorders in TSOD Mice. *Evid. Based Complement. Alternat. Med.* Jun 8.
15. Huang, T.H., Yang, Q., Harada, M., Uberai, J., Radford, J., Li, G.Q., Yamahara, J., Roufogalis, B.D. and Li, Y. (2006) *Salacia oblonga* root improves cardiac lipid metabolism in Zucker diabetic fatty rats: modulation of cardiac PPAR-alpha-mediated transcription of fatty acid metabolic genes. *Toxicol. Appl. Pharmacol.* 210 (1–2), 78–85.
16. Kishino, E., Ito, T., Fujita, K. and Kiuchi, Y. (2006) A mixture of the *Salacia reticulata* (Kotala himbutu) aqueous extract and cyclodextrin reduces the accumulation of visceral fat mass in mice and rats with high-fat diet-induced obesity. *J. Nutr.* 136 (2), 433–439.
17. Shimada, T., Nagai, E., Harasawa, Y., Akase, T., Aburada, T., Iizuka, S., Miyamoto, K. and Aburada, M. (2010) Metabolic disease prevention and suppression of fat accumulation by *Salacia reticulata*. *J. Nat. Med.* 64 (3), 266–274.
18. Kishino, E., Ito, T., Fujita, K. and Kiuchi, Y. (2009) A mixture of *Salacia reticulata* (Kotala himbutu) aqueous extract and cyclodextrin reduces body weight gain, visceral fat accumulation, and total cholesterol and insulin increases in male Wistar fatty rats. *Nutr. Res.* 29 (1), 55–63.
19. Shimoda, H., Asano, I. and Yamada, Y. (2001) [Antigenicity and phototoxicity of water-soluble extract from *Salacia reticulata* (Celastraceae)]. *Shokuhin Eiseigaku Zasshi* 42 (2), 144–147.
20. Im, R., Mano, H., Nakatani, S., Shimizu, J. and Wada, M. (2008) Safety evaluation of the aqueous extract Kothala himbutu (*Salacia reticulata*) stem in the hepatic gene expression profile of normal mice using DNA microarrays. *Biosci. Biotechnol. Biochem.* 72 (12), 3075–3083.
21. Ratnasooriya, W.D., Jayakody, J.R. and Premakumara, G.A. (2003) Adverse pregnancy outcome in rats following exposure to a *Salacia*

- reticulata (Celastraceae) root extract. *Braz. J. Med. Biol. Res.* 36 (7), 931–935.
22. Yoshikawa, M., Ninomiya, K., Shimoda, H., Nishida, N. and Matsuda, H. (2002) Hepatoprotective and antioxidative properties of *Salacia reticulata*: preventive effects of phenolic constituents on CCl₄-induced liver injury in mice. *Biol. Pharm. Bull.* 25 (1), 72–76.
 23. Aljabri, K.S., Bokhari, S.A. and Khan, M.J. (2010) Glycemic changes after Vitamin D supplementation in patients with type 1 diabetes mellitus and Vitamin D deficiency. *Ann. Saudi Med.* 30 (6), 454–458.
 24. Jorde, R. and Grimnes, G. (2011) Vitamin D and metabolic health with special reference to the effect of Vitamin D on serum lipids. *Prog. Lipid Res.* 50 (4), 303–312.
 25. Seshadri, K.G., Tamilselvan, B. and Rajendran, A. (2011) Role of Vitamin D in Diabetes. *J. Endocrinol. Metab.*; 1 (2), 47–56.26.
 26. Grant, W.B. and Holick, M.F. (2005) Benefits and requirements of Vitamin D for optimal health: a review. *Altern. Med. Rev.* 10 (2): 94–111.
 27. Ascherio, A., Munger, K.L. and Simon, K.C. (2010) Vitamin D and multiple sclerosis. *Lancet Neurol.* 9(6), 599–612.
 28. Joseph, A.J., George, B., Pulimood, A.B., Seshadri, M.S. and Chacko, A. (2009) 25 (OH) Vitamin D level in Crohn's disease: association with sun exposure & disease activity. *Indian J. Med. Res.* 130 (2), 133–137.
 29. Judd, S.E. and Tangpricha, V. (2009) Vitamin D deficiency and risk for cardiovascular disease. *Am. J. Med. Sci.* 338 (1), 40–44.
 30. Ginde, A.A., Mansbach, J.M. and Camargo, A.A. (2009) Association Between Serum 25-Hydroxyvitamin D Level and Upper Respiratory Tract Infection in the Third National Health and Nutrition Examination Survey. *Arch. Int. Med.* 169, 384–90.
 31. Cannell, J.J., Zasloff, M., Garland, C.F., Scragg, R. and Giovannucci, E. (2008) On the epidemiology of influenza. *Virology* 375, 29.
 32. Urashima, M., Segawa, T., Okazaki, M., Kurihara, M., Wada, Y. and Ida, H. (2010) Randomized trial of Vitamin D supplementation to prevent seasonal influenza A in schoolchildren. *Am. J. Clin. Nutr.* 91 (5), 1255–1260.
 33. Sowers, M.F.R. and Gadegbeku, C.A. (2009) Vitamin D deficiency in younger women is associated with increased risk of high blood pressure. *American Heart Association Abstract P253K* (Chicago).
 34. Park, S.Y., Murphy, S.P., Wilkens, L.R., Nomura, A.M., Henderson, B.E. and Kolonel, L.N. (2007) Calcium and Vitamin D intake and risk of colorectal cancer: the Multiethnic Cohort Study. *Am. J. Epidemiol.* 165 (7), 784–793.
 35. Peterlik, M., Grant, W.B. and Cross, H.S. (2009) Calcium, Vitamin D and cancer. *Anticancer Res.* 29 (9), 3687–3698.
 36. Trump, D.L., Deeb, K.K. and Johnson, C.S. (2010) Vitamin D: considerations in the continued development as an agent for cancer prevention and therapy. *Cancer J.* 16 (1), 1–9.
 37. Amital, H., Szekanecz, Z., Szucs, G., Danko, K., Nagy, E., Csepany, T., Kiss, E., Rovensky, J., Tuchynova, A., Kozakova, D., Doria, A., Corocher, N., Agmon-Levin, N., Barak, V., Orbach, H., Zandman-Goddard, G. and Shoenfeld, Y. (2010) Serum concentrations of 25-OH Vitamin D in patients with systemic lupus erythematosus (SLE) are inversely related to disease activity: is it time to routinely supplement patients with SLE with vitamin D? *Ann. Rheum. Dis.* 69 (6), 1155–1157.
 38. Artaza, J.N., Mehrotra, R. and Norris, K.C. (2009) Vitamin D and the cardiovascular system. *Clin. J. Am. Soc. Nephrol.* 4 (9), 1515–1522.
 39. Plotnikoff, G.A. and Quigley, J.M. (2003) Prevalence of severe hypovitaminosis D in patients with persistent, nonspecific musculoskeletal pain. *Mayo Clin. Proc.* 78 (12), 1463–1470.
 40. Hollis, B.W. (2005) Circulating 25-hydroxyvitamin D levels indicative of Vitamin D sufficiency: implications for establishing a new effective dietary intake recommendation for vitamin D. *J. Nutr.* 135 (2), 317–322.
 41. Hidayat, R., Setiati, S. and Soewondo, P. (2010) The association between Vitamin D deficiency and type 2 diabetes mellitus in elderly patients. *Acta Med. Indones.* 42 (3), 123–129.
 42. Grober, U. (2010) [Vitamin D – an old vitamin in a new perspective]. *Med. Monatsschr. Pharm.* 33 (10), 376–383.
 43. Shab-Bidar, S., Neyestani, T.R., Djazayeri, A., Eshraghian, M.R., Houshiarrad, A., Kalayi, A., Shariatzadeh, N., Khalaji, N. and Gharavi, A. (2012) Improvement of Vitamin D status resulted in amelioration of biomarkers of systemic inflammation in the subjects with type 2 diabetes. *Diabetes Metab. Res. Rev.* Jul; 28 (5): 424–430.
 44. George, N., Peeyush Kumar, T., Antony, S., Jayanarayanan, S. and Paulose, C.S. (2012) Effect of Vitamin D3 in reducing metabolic and oxidative

stress in the liver of streptozotocin-induced diabetic rats. *Br. J. Nutr.* Jan 6, 1–9.

of the metabolic syndrome and diabetes in overweight and obesity. *Nutr. J.* 7, 4.

45. Nazarian, S., St. Peter, J.V., Boston, R.C., Jones, S.A. and Mariash, C.N. (2011) Vitamin D3 supplementation improves insulin sensitivity in subjects with impaired fasting glucose. *Transl. Res.* 158 (5), 276–281.
46. McGill, A.T., Stewart, J.M., Lithander, F.E., Strik, C.M. and Poppitt, S.D. (2008) Relationships of low serum Vitamin D3 with anthropometry and markers

Michael Ofner, MD, PhD, MBA

Department of Sports Physiology
University of Vienna
Auf der Schmelz 6
1150 Vienna
Austria
michael.ofner@medyco.net