



ORIGINAL PAPER

Treatment for hyperactive children: homeopathy and methylphenidate compared in a family setting

H Frei^{1*} and A Thurneysen²

¹Spezialarzt FMH für Kinder und Jugendliche, FA Homöopathie SVHA, Laupen, Switzerland; and ²University of Berne, Switzerland

The sharp increase of the prescription of methylphenidate (MPD) in hyperactive children in recent years is a matter of increasing uneasiness among professionals, parents and politicians. There is little awareness of treatment alternatives. The purpose of this prospective trial was to assess the efficacy of homeopathy in hyperactive patients and to compare it MPD. The study was performed in a paediatric practice with conventional and homeopathic backgrounds. Children aged 3–17 y, conforming to the DSM-IV criteria for attention deficit hyperactivity disorder (ADHD) with a Conners Global Index (CGI) of 14 or higher were eligible for the study. All of them received an individual homeopathic treatment. When clinical improvement reached 50%, the parents were asked to reevaluate the symptoms. Those who did not improve sufficiently on homeopathy were changed to MPD, and again evaluated after 3 months. One hundred and fifteen children (92 boys, 23 girls) with a mean age of 8.3 y at diagnosis were included in the study. Prior to treatment the mean CGI was 20.63 (14–30), the mean index of the homeopathy group 20.52 and of the MPD-group 20.94. After an average treatment time of 3.5 months 86 children (75%) had responded to homeopathy, reaching a clinical improvement rating of 73% and an amelioration of the CGI of 55%. Twenty-five children (22%) needed MPD; the average duration of homeopathic (pre-) treatment in this group was 22 months. Clinical improvement under MPD reached 65%, the lowering of the CGI 48%. Three children did not respond to homeopathy nor to MPD, and one left the study. In cases where treatment of a hyperactive child is not urgent, homeopathy is a valuable alternative to MPD. The reported results of homeopathic treatment appear to be similar to the effects of MPD. Only children who did not reach the high level of sensory integration for school had to be changed to MPD. In preschoolers, homeopathy appears a particularly useful treatment for ADHD. *British Homeopathic Journal* (2001) 90, 183–188.

Keywords: hyperactive children; ADHD; homeopathy; methylphenidate

Introduction

The trends in the prevalence of attention deficit hyperactivity disorder (ADHD) and the prescription of methylphenidate (MPD) in children and adoles-

cents in North America have shown a marked increase during the past decade.^{1,2} Reported prescription rates range from 1.1% in Michigan (children 0–19 y),³ 3.4% in Ontario (students grades 7, 9, 11, 13)⁴ to 8–10% in south-eastern Virginia (students grades 2–5, with a maximum of 18–20% of grade 5 white boys).⁵ The increase does not seem to be limited to the US and Canada: In Switzerland, as in many other Western countries, the frequency of the diagnosis of ADHD and prescription of MPD have also risen remarkably during the past few years.^{6,7}

*Correspondence: H Frei, Spezialarzt FMH für Kinder und Jugendliche, FA Homöopathie SVHA, Kreuzplatz 6, CH-3177 Laupen, Switzerland.
E-mail: dr.heiner.frei@swissonline.ch
Received 19 June 2000; revised 8 August 2000; accepted 11 December 2000

Along with this rise comes a concern for more accurate diagnosis of ADHD,⁸ and reports of abuse of MPD which has similarities with cocaine in terms of pharmacodynamics and pharmacokinetics.^{9–12} Other problems include noncompliance with frequent dosing and wear-off or rebound effects.¹³

For parents of hyperactive children the fact that their child is receiving long-term treatment with a substance that falls under the legislation for narcotics (in Switzerland) is often a cause of major concern. Many of them refuse such a treatment unless the schools exert extreme pressure. One of the main social causes for the rise in the prescription of MPD may be found in the lowering of public education budgets in recent years, leading to larger school classes in which hyperactive behaviour is less tolerable.

It is not surprising therefore, that professionals seek options in pharmacotherapy and parents look for alternative treatments, despite the lack of controlled research on their efficacy and safety.^{13–15}

The purpose of this trial was to assess the efficacy of homeopathic treatment^{16–18} in ADHD, answering the following questions:

- What percentage of children can be sufficiently treated with homeopathy and need no other medication? How many need MPD? And how many do not respond to these treatments at all?
- What is the effect of homeopathic treatment and MPD as rated by the CGI?^{19–21}
- How do parents rate clinical improvement, including feedback from school?
- Time horizons: how long is needed to reach an adequate treatment effect in homeopathy? What was the duration of homeopathic treatment in patients who finally received MPD?

Methods

Children between 3 and 17 y conforming to the DSM-IV diagnostic criteria for ADHD were eligible for the study.^{22*} The diagnostic procedures included meticulous history taking, a general and neurological examination (as described by the author earlier²³) and an

*DSM-IV diagnostic criteria for ADHD:

(1) presence of either six symptoms of inattention or six symptoms of hyperactivity–impulsivity, which have persisted for at least 6 months to a degree that is maladaptive and inconsistent with development level; (2) presence of some symptoms that caused impairment before age 7 y; (3) presence of some impairment from symptoms in two or more settings (eg school or work and at home); (4) clear evidence of clinically significant impairment in social, academic, or occupational functioning; (5) the symptoms do not occur exclusively during the course of a pervasive developmental disorder, schizophrenia, or other psychotic disorder and are not better accounted for by another mental disorder (eg mood disorder, anxiety disorder, dissociative disorder, or a personality disorder).

assessment of the hyperactivity and attention deficit symptoms according to the Conners 10 item rating scale (Conners Global Index¹⁹). Patients with a CGI of 14 or higher were included in the study. If there was any doubt concerning the diagnosis of ADHD, patients were referred to a child and adolescent psychiatrist or psychologist or a paediatric neurologist for further testing (36 children, 31% of all patients).

Each child was first treated with homeopathy. To be effective, the homeopathic medicament has to match the *individual* symptoms of the patient, ie the symptoms that are *not* commonly present in most hyperactive children and therefore distinguish him from the others. This process of individual adaptation of the treatment may require some time, and include trials of possible medicaments, until the optimal effect is reached.

The matching of patient-symptoms and homeopathic remedies was performed following the procedures of Hahnemann,^{16–18} assisted by a computer-program (Amokoor²⁴) based on the works of Boenninghausen.^{25–27} The prescribing technique has been described by the author in earlier publications.^{28,29} In this trial homeopathic preparations of the following medicaments were used successfully (number of patients in parenthesis): *Lyc.* (12), *calc.* (7), *sulph.* (7), *bell.* (6), *caust.* (6), *phos.* (6), *ign.* (5), *nux-v.* (5), *arg-n.* (4), *sep.* (4), *lach.* (3), *merc.* (3), *puls.* (3), *sil.* (3), *ars.* (2), *staph.* (2), *agar.* (1), *bar-c.* (1), *bry.* (1), *chin.* (1), *hep.* (1), *hyos.* (1), *nat-m.* (1) and *stram.* (1).³⁰ All patients received liquid LM-potencies¹⁸ (LM-3 to LM-30) every day or every second day, depending on the severity of their symptoms. Each potency (eg LM-3) was used for 4 weeks, moving on to the next higher level (eg LM-6) after a treatment-free interval of several days to one week. If the child's reaction to the medication was insufficient (wrong choices usually do not change the hyperactivity symptoms), the next most similar remedy was prescribed. Once an adequate response had been reached, the children received the next higher potency of the same medicine.

For clinical assessment of treatment the parents had to report the changes observed in every symptom they initially reported, ie hyperactivity 'considerably improved', 'slightly improved', 'unchanged' or 'worse'. After reporting the changes of every individual symptom they were asked to summarize the overall clinical improvement as a percentage. When the overall amelioration reached 50% or more, the treatment was reassessed by the CGI rating scale. The timing of this reassessment thus was individual, depending on the time required to find the correct homeopathic medicine. Patients who did not reach sufficient clinical improvement, or whose behaviour remained unacceptable despite a certain response to homeopathy were changed to MPD after reevaluation. The point at which a patient was deemed a treatment

failure thus was individual, dependent on environmental tolerance for his behaviour. Many children had a long-term homeopathic treatment, before a crisis (usually school pressure) made MPD necessary.

Two weeks after the initiation of MPD-treatment, the CGI was determined to distinguish responders from non-responders. The final evaluation of this treatment by CGI followed 3 months after the optimal adjustment of MPD-dosage.

Patients

One hundred and fifteen children (92 boys, 23 girls) conformed to the eligibility criteria. Their mean age at diagnosis was 8.3 y. In the homeopathy group 76% of the patients were boys, 24% girls, with a mean age of 7.9 y at diagnosis. In the MPD group 92% were boys, 8% girls, with a mean age of 9.6 y. Non-responders and drop-outs were all boys with a mean age of 9.0 y.

Results

Treatment modalities in ADHD patients

Eighty-six patients (75%) responded sufficiently to homeopathy, and 25 (22%) needed MPD. Only three patients (3%) did not respond neither to homeopathy nor to MPD (Figure 1). One child left the study.

Comparison of response to homeopathy and MPD

The mean value of the CGI ratings of all patients prior to treatment was 20.63, the homeopathy group 20.52, and the MPD group 20.94. During homeopathic treat-

ment the mean CGI rating fell to 9.27 corresponding to an amelioration of 55%, and with MPD to 10.96, corresponding to an amelioration of 48% (Figure 2). The CGI prior to change to MPD was determined only in a small group of patients. It reached an average value of 13.0 which corresponds to an amelioration of 37%.

Clinical improvement ratings

Clinical improvement ratings by parents in homeopathy treated children were 73%, and with MPD 65% (Figure 3). Most of the patients who had eventually received MPD had a treatment effect from homeopathy, but the mean clinical improvement was 43%, considerably lower than in children who responded well to homeopathy.

Time horizons

The average time needed to reach an optimal homeopathic treatment effect was 3.5 months (range 1–16 months, Figure 4), the mean duration of homeopathic treatment in those patients who finally needed MPD was 22 months (range 4–62 months, Figure 5).

Discussion

In an earlier placebo-controlled study, Lamont showed, that homeopathy is an effective treatment in hyperactive children.³¹ Instead of long discussions whether or not homeopathy is placebo, its effects should be assessed by the same scales that are applied in mainstream medicine. In a situation where alternative treatments are frequently used, it is essential to know what can be expected of them.

It is surprising that 75% of the studied children reached a satisfactory amelioration with homeopathy in a *family* setting. Conversely most children who needed MPD, did so because of *school* pressure and not the situation at home.

The observed parent ratings of clinical improvement and the lowering of the CGI under homeopathy were slightly better than under MPD. This finding may be due to the short duration of action of MPD (4 h in the normal and 8 h in the retard form³²), which often leads to difficult times at noon and in the evening (observation of the authors). Therefore it is mainly the school-situation that profits from MPD. The difference between clinical amelioration and CGI ratings can be explained by the fact that every amelioration in a hyperactive child is an enormous relief for family and school. The higher clinical improvement ratings reflect this relief, while the detailed 10-item ratings with the CGI show a more realistic picture of what has really been achieved.

It may be argued that all children who received MPD also had a homeopathic pre-treatment, and that they may therefore react better to MPD than children without pre-treatment. The authors do not think that

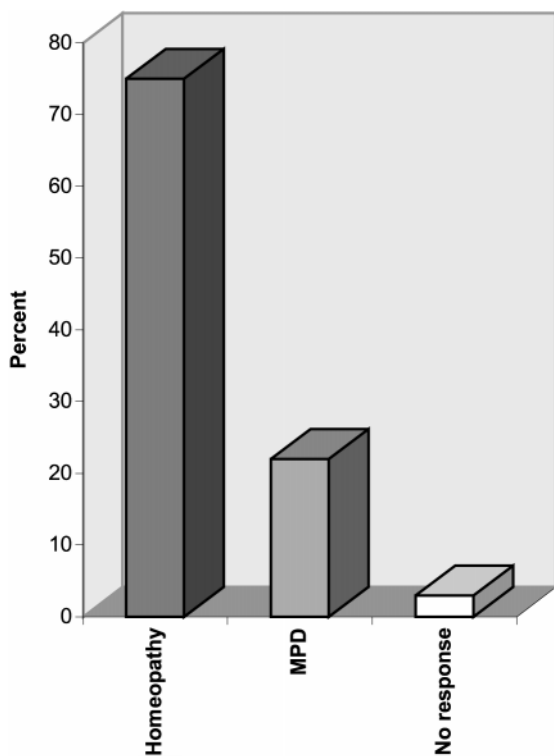


Figure 1 Treatment response in ADHD patients.

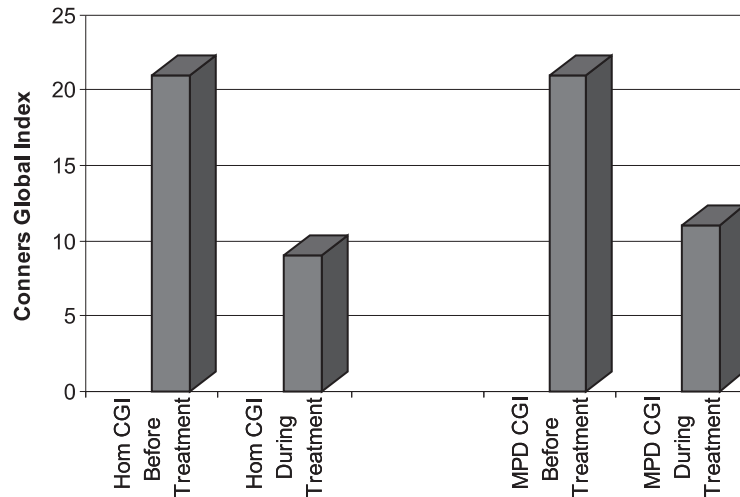


Figure 2 Changes of CGI under treatment with homeopathy and methylphenidate.

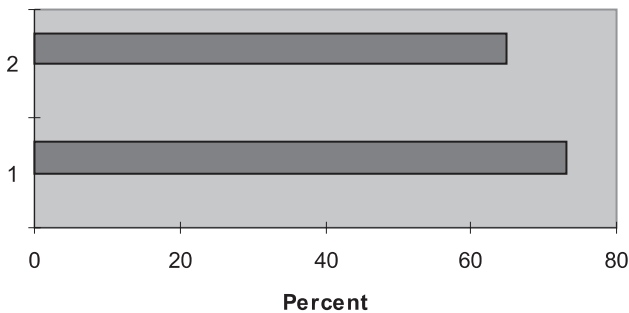


Figure 3 Clinical amelioration in ADHD-patients during treatment with homeopathy (1) and methylphenidate (2).

this is the case, because in the treatment-free intervals between the homeopathic medicines, most children show a reappearance of the hyperactivity symptoms.

This finding favours the impression, that homeopathy is, like MPD, a palliative treatment. Long-term follow-up studies over several years would be necessary to settle the question as to, whether or not a curative effect can be expected.

A problem in homeopathy is the delay until the optimal amelioration is reached. Since it is necessary to make an individualized prescription, it is difficult to treat in a situation where an improvement has to be immediate. The choice of the correct medication is dependent on the individuality of the symptoms, if a patient only has the ‘standard symptoms’ of ADHD and nothing peculiar, the homeopathic physician may have to make ‘therapeutic trials’ to find the correct medicine. The administration of a wrong remedy usually does not change anything, while giving the

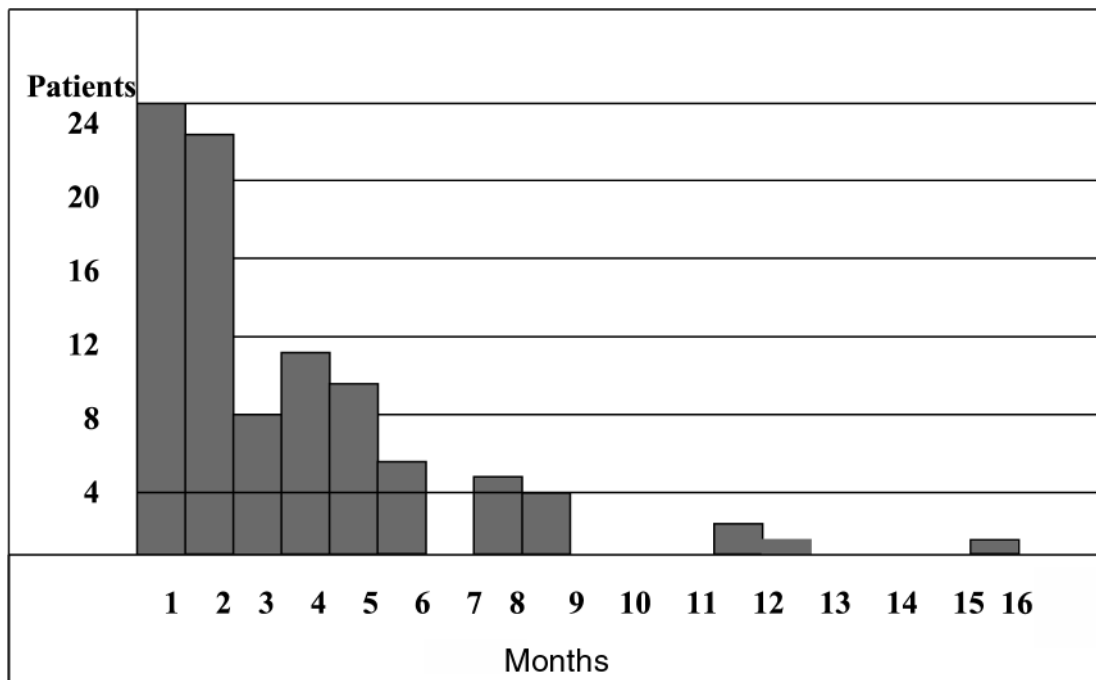


Figure 4 Time required for reaching an optimal homeopathic treatment-effect with ADHD.

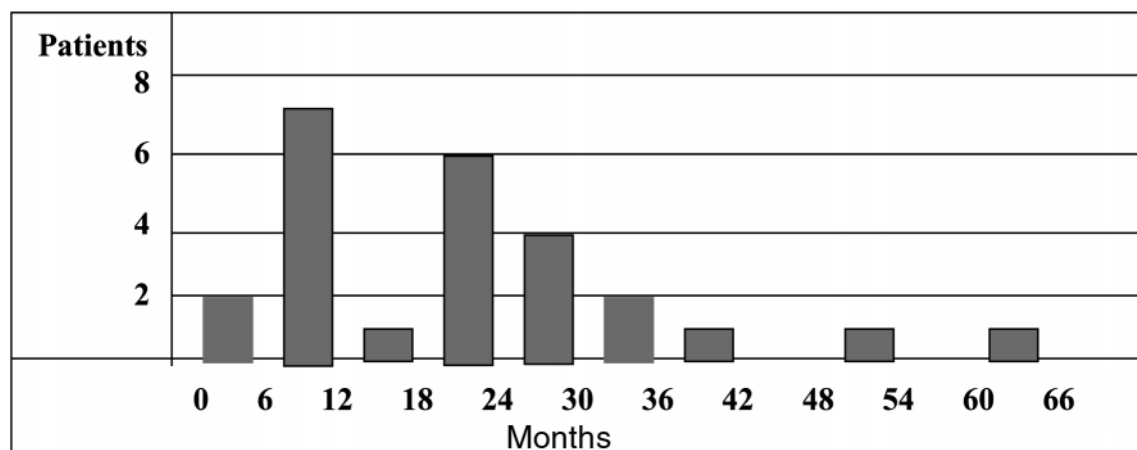


Figure 5 MPD-group: duration of homeopathic treatment prior to MPD.

right one leads to a clear, substantial improvement within 4 weeks. Figure 4 shows, that the majority of children responding to homeopathy do so within 6 months. If a child has not responded by then, it is unlikely that it will, and MPD treatment may be considered at this time.

Major advantages of homeopathy over MPD are the easy administration (once every day or once every second day), a continuous treatment effect over 24 h, no side effects except for a possible short initial aggravation, and no abuse potential. For many parents this last point is the most important concern. At preschool-age, when MPD has many side effects,³³ homeopathy may be the first choice, as well as for students, who do not need to have an immediate relief. Finally, there is an extremely low number of non-responders if both methods are available (3%).

Acknowledgements

The authors owe sincere thanks to the following people, who have contributed to this work with their advice concerning procedures and/or patients: Dr P Hämmerli, Dr M Bettler, Dr F Blaser, child and adolescent psychiatry, Dr M Stucki, Dr F Kaufmann and D Walther, MA, pediatric psychology, DL Lindstrom, MS, statistics, Dr R Hassink and Dr L Grimm, pediatric neurology, Dr M Ryffel, MPD-treatment, Dr KH Gypser and Dr U Steiner, homeopathy.

References

- 1 Robinson LM, Sclar DA, Skaer TL, *et al.* National trends in the prevalence of attention-deficit/hyperactivity disorder and the prescription of methylphenidate among school-age children: 1990–1995. *Clin Pediatr (Phila)* 1999; **38**: 209–217.
- 2 Goldman LS, Genel M, Bezman RJ, *et al.* Diagnosis and treatment of attention-deficit/hyperactivity disorder in children and adolescents. Council on Scientific Affairs, American Medical Association. *JAMA* 1998; **279**: 1100–1107.
- 3 Rappley MD, Gardiner JC, Jetton JR, *et al.* The use of methylphenidate in Michigan. *Arch Pediatr Adolesc Med* 1995; **149**: 675–679.
- 4 Ivis FJ, Adlaf EM. Prevalence of methylphenidate use among adolescents in Ontario. *Can J Public Health* 1999; **90**: 309–312.
- 5 LeFever GB, Dawson KV, Morrow AL, *et al.* The extent of drug therapy for attention deficit-hyperactivity disorder among children in public schools. *Am J Public Health* 1999; **89**: 1359–1364.
- 6 Baumgaertel A, Wolraich ML, Dietrich M, *et al.* Comparison of diagnostic criteria for attention deficit disorders in a German elementary school sample. *J Am Acad Child Adolesc Psychiatry* 1995; **34**: 629–638.
- 7 Andres Carrasco MA, Catala MA, Gomez-Beneyto M, *et al.* Study of the prevalence of attention deficit hyperactivity disorder in ten-year-old children living in the Valencia metropolitan area. *Actas Luso Esp Neurol Psiquiatr Cienc Afines* 1995; **23**: 184–188.
- 8 Schneider SC, Tan G. Attention-deficit hyperactivity disorder. In pursuit of diagnostic accuracy. *Postgrad Med* 1997; **101**: 231–232, 235–240.
- 9 Llana ME, Crismon ML. Methylphenidate: increased abuse or appropriate use? *J Am Pharm Assoc (Wash)* 1999; **39**: 526–530.
- 10 Massello W III, Carpenter DA. A fatality due to intranasal abuse of methylphenidate (Ritalin). *J Forensic Sci* 1999; **44**: 220–221.
- 11 Crutchley A, Temlett JA. Methylphenidate (Ritalin) use and abuse. *S Afr Med J* 1999; **89**: 1076–1079.
- 12 Musser CJ, Ahmann PA, Theye FW, *et al.* Stimulant use and the potential for abuse in Wisconsin as reported by school administrators and longitudinally followed children. *J Dev Behav Pediatr* 1998; **19**: 187–192.
- 13 Garland EJ. Pharmacotherapy of adolescent attention deficit hyperactivity disorder: challenges, choices and caveats. *J Psychopharmacol* 1998; **12**: 385–395.
- 14 Stupperfeld T, Parry T. Utilisation of alternative therapies in attention deficit hyperactivity disorder. *J Paediatr Child Health* 1999; **35**: 450–453.
- 15 Baumgaertel A. Alternative and controversial treatments for attention-deficit/hyperactivity disorder. *Paediatr Clin North Am* 1999; **46**: 977–992.
- 16 Hahnemann FS. *Organon der Heilkunst*, 6th edn. Heidelberg, Germany: Haug, 1989.
- 17 Hahnemann FS. *Die Chronischen Krankheiten*, 5th edn. Heidelberg, Germany: Haug, 1991.
- 18 Hahnemann FS. *Reine Arzneimittellehre*, 4th edn. Heidelberg, Germany: Haug, 1989.

- 19 Conners CK. *Conners' Rating Scales* (Revised), Technical Manual. Toronto, Canada, 1997.
- 20 Conners CK. Rating scales in attention-deficit/hyperactivity disorder: use in assessment and treatment monitoring. *J Clin Psychiatry* 1998; **59**(Suppl 7): 24–30.
- 21 Conners CK. Clinical use of rating scales in diagnosis and treatment of attention-deficit/hyperactivity disorder. *Paediatr Clin North Am* 1999; **46**: 857–870, vi.
- 22 American Psychiatric Association. *Diagnostic and Statistical Manual of Mental Disorders*, 4th edn. Washington DC: American Psychiatric Association, 1994.
- 23 Frei H. The 'clumsy' child: differential diagnosis and therapeutic indications. A review. *Schweiz Med Wochenschr* 1986; **116**: 294–299.
- 24 Steiner U. *Amokoor—Homeopathy Software* based on Boenninghausen Cv. Goldau, Switzerland, Steiner, 1992.
- 25 Boenninghausen C.v. *Therapeutisches Taschenbuch für homöopathische Aerzte* 1897. Hamburg, Germany: Von der Lieth, 1990.
- 26 Boenninghausen Cv. *Die Aphorismen des Hippokrates*. Göttingen, Germany: Burgdorf, 1979.
- 27 Boenninghausen Cv. *Kleine Schriften*. Heidelberg, Germany: Gypser, 1984.
- 28 Frei H. Die Heringsche Regel und ihre Auswirkung auf die Hierarchie der Symptome. *Z Klass Hom* 1999; **43**: 47–52.
- 29 Frei H. Die Rangordnungen der Symptome von Hahnemann, Boenninghausen, Hering und Kent, evaluiert anhand von 175 Kasuistiken. *Z Klass Hom* 1999; **43**: 143–155.
- 30 Hering C. *The Guiding Symptoms of our Materia Medica*. New Dehli, India: Jain, reprinted, 1991.
- 31 Lamont J. Homoeopathic treatment of attention deficit disorder. *Br Hom J*, 1997; **86**: 196–200.
- 32 Morant J, Ruppner H. *Arzneimittelkompendium der Schweiz*. Basel, Switzerland: Documed, 2000.
- 33 Byrne JM, Bawden HN, De Wolfe NA, et al. Clinical assessment of psychopharmacological treatment of preschoolers with ADHD. *J Clin Exp Neuropsychol* 1998; **20**: 613–627.