

## Fundamental Research

### A Strategy for Structural Exploration of Homoeopathic Medicines

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#### Abstract

Absence of drug material in high potency suggests that physics is the only way to understand homoeopathic medicines. Experiments carried out also demonstrate that they are different in nature from controls. But their structure is still unknown. To solve this scientific problem, a new approach of studying medicines prepared from precise variable sources like X-rays and magnetic fields is suggested to identify the factor responsible for medicinal action. Study of Causticum and medicines prepared from plants grown in different localities is also shown to be of help.

#### Keywords

X-ray, Magnetic field, Causticum, and vegetable medicines.

#### Introduction

The higher potencies of homoeopathic medicines have always been controversial since their introduction about two centuries ago. They are the major causes of scepticism against homoeopathy. Even some veteran homoeopaths, like Richard Hughes of England in the past, do not believe them. These medicines contain nothing materially and thus contradict our materialistic commonsense. Hahnemann was the first to recognize this fundamental scientific problem and the necessity of searching for a solution within the science of physics.<sup>1</sup>

#### Tools: Experimental Methods of Physics

Garth Boericke and Rudolph Smith began investigation of higher potencies to explore their structure using nuclear magnetic resonance (NMR) spectroscopy in 1963.<sup>2</sup> Their painstaking work is of great historical importance as it could demonstrate explicitly that homoeopathic medicines are different from controls (Vehicle, i.e., host or unsuccessed dilutions). And that tools of quantum physics could be employed further to explore their nature. The difference in NMR patterns is, however, small but significant and persistent.

Several other experimental methods, like infrared spectroscopy, Raman laser spectroscopy, dielectric constant, surface tension etc., followed the work of Boericke and Smith. These experimental methods have also supported objective physicochemical changes in high potencies,<sup>3</sup> but results obtained in this manner still remain obscure.

#### New Samples: Medicines from Variable Sources

One reason behind the inconclusive results seems to be that we do not choose such medicines for study that could be varied (at our will) and about which we have precise information. Such medicines do exist in homoeopathy. We should think over potentised X-ray and potentised magnetic field. One extra advantage of choosing

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them is that their lower potencies and even mother tinctures do not contain them as atoms or molecules.

Homoeopathic medicine X-ray is obtained by exposing a vial containing alcohol to X-rays. These X-rays are not of one precise wavelength but of continuous type from a Coolidge tube. The alcohol is now taken as the mother tincture and potencies are raised. X-ray is not a trivial medicine. In fact, it is a very powerful centrifugal remedy suitable to mixed miasm dominated by Sycosis.

There is a lot of difference between X-rays of wavelengths 1 Å and 100 Å. Their hazardous effects are also not the same. If their provings as homoeopathic medicines are carried out, they are unlikely to show the same Symptomatology. Taking monochromatic X-rays, we can prepare different medicines like X-ray 1Å, X-ray 10 Å and X-ray 100 Å. The precise information about these medicines is the wavelength (or frequency) alone. Their potencies can be raised from mother tinctures to say 3C, 6C, etc. in centesimal scale. Some medicines can also be prepared by taking composite wavelengths like 1Å, 50Å, and 100Å and other by taking them one by one on the same vial of alcohol. Here more samples may be possible by taking these wavelengths in different orders on different vials of alcohol. Thus we have a lot of meaningful samples of X-rays for exhaustive study. The variation in intensity of X-rays employed and duration of exposure of the vial may also prove useful in the study. Weak cosmic variations are unlikely to interfere.

The magnet is also a source of medicine in homoeopathy. The pathogenesis of it as a whole and of each pole separately are given in Hahnemann's *Materia Medica Pura*.<sup>4</sup> Medicines from it are *Magnetis poli ambo* (magnet), *Magnetis polus arcticus* (North pole), *Magnetis polus australis* (South pole). The precise information about these medicines is the exact intensity of magnetic field applied alone with the duration of exposure. Varying these parameters, several

mother tinctures can be prepared and their potencies can be raised. *Magnetis polus arcticus* and *Magnetis polus australis* are of special interest. If we take the former as a "source", the latter is a "sink." Weak geomagnetism is unlikely to interfere. A systematic study of the number of such samples in an orderly manner is likely to result in new understanding about homoeopathically prepared medicines made from energy fields.

Vegetable medicines are also of keen interest. A plant is not made up of one chemical but of many. It is well known that plants of the same species grown in different areas have considerable variations in chemical composition but have the same therapeutic effect as homoeopathic medicines. This suggests that a (vegetable) medicine is not the aggregate of its constituents. It may also be possible that either therapeutically dominating group contributes only in the preparation of potencies or the human system accepts the contribution of this group only. One more possibility is that among composite therapeutic signals from a vegetable remedy only one signal gets selected in one prover. Thus, with many provers a medicine can acquire a wide spectrum of symptoms, even of contrary ones, and during a cure only the *similimum* signal is selected for action by a living organism. So a comparative study of potencies raised from a vegetable grown in three or more areas of different soil composition and climate is likely to prove meaningful. Here mother tinctures of different origin like British, French, German, Indian, etc. may also solve the purpose.

*Causticum* is a medicine of mineral origin. It is a polychrest of high order. Hahnemann prepared it by distilling a mixture of recently slaked lime and previously burned and melted potassium bisulphate. Thus, it was the product of the crude pharmaceuticals of his time. Mother tinctures of *Causticum* so produced at different times (and by different persons) could hardly be the same chemically. Worse yet, it has rarely ever been

manufactured by Hahnemann's method.<sup>5</sup> Yet his symptoms seem to be valid for any type of Causticum produced according to different rules.<sup>5</sup> So by collecting mother tinctures of Causticum from different manufacturers, their potencies can be raised for comparative study. These high potencies may also be purchased directly from them to reduce the cost of research.

### Conclusion

According to Bernard Poitevin B., "Understanding of the mechanism of action of homoeopathic medicines in high dilution is gradually coming closer to the outstretched hand of scientific research, thanks in particular to improvements in methods of physical analysis of solvents."<sup>6</sup> A series of potencies of such medicines is needed for statistical analysis and comparative studies, whose sources, especially the precise ones, can be varied in acceptable terms. Such a study would likely be able to identify the common factor responsible for medicinal action.

### References

1. Bernal, G.G. (1993). Homeopathy and physics. *British Homoeopathic Journal*, 82: pp210.
2. Smith, R.B., and Boericke, G. W. (1968). Changes caused by succussion on NMR pattern and bioassay of Bradykinin triacetate (BKTA) succussions and dilutions. *Journal of American Institute of Homoeopathy*, 61: pp 197-212
3. Poitevin, B. (1995), Mechanism of action of homoeopathic medicines. *British Homoeopathic Journal*, 84: pp 33
4. Hahnemann, S. (2000). *Materia Medica Pura*. B. Jain Publishers Pvt. Ltd., New Delhi
5. Walach, H. (1999). Magic of signs: A nonlocal interpretation of Homoeopathy. *Journal of Scientific Exploration*, 13: pp 294
6. Poitevin, B. (1995), Mechanism of action of Homoeopathic medicines. *British Homoeopathic Journal*, 84: pp 39.