

Correlation of Medical Anamnesis with Current Disease in Chronic Kidney Disease Patients Receiving Hemodialysis: A Retrospective Study of Causative and Exacerbating Factors from Homeopathic Point of View

ORIGINAL

Abstract

Background: Chronic diseases are illnesses or health conditions that are persistent and have long-term effects on the physical status of the human body. The manifestation of a chronic disease is the result of coexistence of multiple chronic diseases. Especially, the chronic kidney disease (CKD) is a representative chronic syndrome of multifactorial affair, characterized by a gradual progression.

Methods: This study considered 81 patients with CKD and investigated the correlation between the disease and the patients' demographics, anamnesis, and clinical presentation, as well as the potential relationship between the disease and other illnesses.

Results: A significant number of diseases can lead to the manifestation of CKD, and notably, many of them are idiopathic. However, despite the rapid development in medical science and pharmaceutical technology and a plethora of research trials, there is a lack of ability to identify and analyze the causes and pathophysiologic mechanisms that are responsible for the manifestation of CKD.

Conclusions: This retrospective study investigates the factors that may play a role in the manifestation of CKD through the knowledge and research of both conventional medicine and homeopathy.

Background

The causes of chronic diseases seemingly have unsatisfactory explanations for modern medicine. Thus, medical science defines a set of illnesses as diseases of unknown etiology (idiopathic diseases),

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in which the disease's cause is not readily apparent. Examples include idiopathic hypertension, idiopathic glomerulonephritis, fever of unknown origin, urticaria of unknown origin, pulmonary fibrosis, and, by extension, chronic autoimmune diseases, malignancies, and focal segmental glomerulosclerosis [1-2]. In addition, the "comprehensive" causative relationships between the cause and the progression of a disease do not seem to explain this complexity [3].

Chronic diseases are illnesses or other health conditions that are persistent and have long-term effects on the physical status of the human body. They disrupt and undermine the patient's health to such a degree that the energy of the vital force that automatically drives life can defy these diseases only in part and without effect both at their onset and during their progression [4]. The manifestation of a chronic disease is a multifactorial affair, especially in chronic kidney disease (CKD) which is a priori the result of the coexistence of multiple chronic diseases. This also constitutes a first approach, according to conventional medicine, as to why not all humans with the same diseases also present the same progression or complications [5]. CKD is a representative chronic syndrome, since it is characterized by a gradual progression and multifactorial nature [6-7].

Significant number of diseases can lead to the manifestation of CKD, and, notably, many of them are idiopathic. As evidenced by literature and daily clinical practice, there is an inability to identify and analyze the causes and pathophysiologic mechanisms that are responsible for the manifestation of CKD. This is despite the rapid development in medical science and pharmaceutical technology as well as a plethora of research trials. This study therefore investigates factors that may play a role in the manifestation of CKD through the knowledge and research of both conventional medicine and homeopathy.

Homeopathy approaches the human body holistically, considering it as a whole—a set of interacting

mental, emotional, and physiological processes. The fundamental principle of homeopathy is that each symptom the body develops is the expression of its defense and its effort to combat the disorder [8]. According to Professor Vithoukas, the key differences between conventional medicine and homeopathy lie in the principles encompassing them. Specifically, conventional medicine looks for the causes of a disease and targets it directly (e.g., a bacterium or a virus), whereas homeopathy tries to stimulate and strengthen the defense mechanism so that the body can address the underlying cause through its immune system. When adverse factors are more potent than the body's defenses, symptoms manifest and constitute the expression of the disorder on a physical and/or mental level [9-10]. A miasm is the body's predisposition for certain categories of diseases and something that hinders the action of the homeopathic remedy. As a result, the body's vital force is unable to fight them. Homeopathic medicine defines a miasm as a "peculiar morbid derangement of the vital force" of the human body [10-14]. According to Hahnemann, the initial exposure of the body to miasms causes local symptoms, such as skin or venereal diseases, but if these symptoms are suppressed by chemical medications, the cause—the miasm—goes deeper and begins to manifest itself in the form of diseases of internal organs. Thus, homeopathy maintains that the attempt to treat diseases by directly addressing their symptoms (as usually applied in conventional medicine) is ineffective [15-18]. Notably, according to the homeopathic analysis of miasms, the predisposed weakness of the human body's defense mechanism can be affected by three major factors: 1) the hereditary effect, 2) potent infectious diseases, and 3) previous therapies and vaccinations. In addition, for a deeper disorder, the internal miasm will manifest itself as a serious disease following exposure to stress, poor living conditions, and improper and exhaustive allopathic therapy [15]. Consequently, according to the law of disease sup-

pression, when a disease is not treated in depth and causally, then it is simply suppressed. A disease manifesting in the body must be cured and, according to the homeopathic view, by treating the symptoms only, one does not truly address the disease because one is applying a mere superficial treatment of deeper health disorders. Suppressing the symptoms of a disease often causes adverse effects on the body. The suppressed disease tends to affect deeper structures and settles in healthier tissues and organs [20-21]. The consolidated theory on disease constitutes a valuable source of information and understanding of the natural history of a disease, since it also allows for the analysis of symptoms as well as the pathophysiological behavior of the human body. Thus, it is interesting that through the life of a person, from birth to death, there is a 'continuum' in the pathological conditions a person may experience. The body, as a whole, suffers deeply any time there is an acute or a chronic condition that is either maltreated or neglected. Chronic and acute diseases in the medical history of a person constitute a rigidly related chain of immune responses in the form of a real 'continuum' that at every point in time indicates the end result of this continuum, [4]. Suppression of symptoms and diseases, which is often achieved through excess chemical agents, often overwhelms the body's natural defenses and forces the immune system to compromise. This starts a deeper line of defense, which then constitutes the beginning of a new chronic condition [19].

Study Objective

The objectives of this study are: 1) to investigate the correlation between the disease and the patients' demographics, 2) to comparatively evaluate the relationship between the disease and the existence of other infections, and 3) factors regarding the anamnesis and the clinical presentation of the patients.

Methods

This study included 81 patients with CKD and investigated the correlation between the disease and the patients' demographics, anamnesis, and clinical presentation, as well as the potential relationship between the disease and other illnesses. Women accounted for 51.8% of study participants; the remaining 48.2% were men. The median age was 60.7 years with a standard deviation of 11.39 years (range: 38 - 85 years). At the start of the study, we recorded the sex and the age of the participants, the age of onset of CKD, disease treatments, clinical presentation, and anamnesis. Specifically, clinical presentation included whether the subject developed fever (and how high) or had had high fever in recent years, infections (and the age of onset and treatment), and relapses. Reporting anamnesis included frequent use of antibiotics or chronic use of chemical medicines, as well as a family history of kidney disease or sexually transmitted diseases.

Questionnaires were completed following interview of the subjects. We studied and analyzed descriptors of the variables. Standard measures of position and dispersion were used for the description of demographics and the recording of the frequency and the relative frequency of the questions included in the main questionnaire. For the potential correlation between personal details, as well as the various factors reported regarding the health status of patients with CKD, we used Pearson X² (chi-square) heterogeneity test. Where the conditions were not met, we used Fisher's exact test. In addition, Pearson's correlation coefficient was used to check the potential relationships between quantitative variables. Reported p-values were based on two-sided tests, and $p < 0.05$ was considered statistically significant. The software SPSS (SPSS Inc., 2003, Chicago, USA) was used for the statistical analysis.

Results

Basic Descriptive Data

With regards to the underlying cause of CKD, 45.7% of subjects had chronic glomerulonephritis, 23.2% tubular necrosis, 4.9% obstructive uropathy, and the remaining 26.2% had other diseases. For history of infection, 17.3% of the subjects had urinary tract infection, and 76.5% developed some other type of infection. Only 6.2% did not develop any infection (**Table 1**). Next, we analyzed etiological factors that were possibly connected to CKD. We found that 92.6% of subjects had concomitant physical illnesses, and 6.2% presented anxiety disorders, while only one subject did not manifest other diseases (**Table 2**). Regarding chemical medicines, 93.3% of subjects used antibiotics, while 66.7% frequently used them (**Table 3**). Moreover, 82.7% made chronic use of chemical medicines (**Table 4**). In addition, 43.1% of subjects developed infection relapse (**Table 5**). Notably, 23.5% of the subjects with CKD had a family history of kidney disease (**Table 6**) and 32.1% had a history of sexually transmitted diseases (**Table 7**).

Statistically Significant Data

There was a strong correlation (p -value < 0.001) between the disease and recurrent infections. We found that 79.17% of the aggregate sample presented recurrent infections. The correlation between the disease and the development of recurrent urinary tract infections was also statistically significant (Fisher's exact p -value < 0.001) (**Table 8**). In addition, we found a strong correlation ($X^2=79.98$ and p -value < 0.001) between the disease and the chronic use of chemical medicines. We found that 95.7% of subjects with CKD were using chemical medicines (Table 9). Of the subjects asked about their history of sexually transmitted diseases, 26 of 81 patients with CKD reported a history of sexually transmitted diseases (STD). Of them, 53% reported urethritis (either gono-

coccal or non-gonococcal), 32% reported Human Papillomavirus (HPV) infection, and 15% could not report the type of the sexually transmitted disease. Furthermore, there was a correlation between CKD and the history of STD ($X^2 = 19.33$ and p -value < 0.001). Finally, the relationship between the disease and the existence of causative factors (physical and mental stress) showed a strong correlation ($X^2 = 93.29$ and a p -value < 0.001). Specifically, 92.6% of subjects with CKD presented physical stress, but only 6.2% presented mental stress (**Table 10**).

Table 1. Distribution (absolute and relative frequencies) of the health characteristics of patients enrolled in the study.

Disease	N	%
Chronic Glomerulonephritis	35	43,2
Chronic Tubular Necrosis	37	45,7
Obstructive Uropathy	4	4,9
Other	5	6,2
Total	81	100
Infection		
None	5	6,2
Urinary Tract Infection	14	17,3
Other Infections	62	76,5
Total	81	100

Table 2. Table of correlation between causative factors and CKD.

Other Causative Factors - Possible Correlation with the Development of Chronic Renal Failure	N	%
Physical Shock	75	92,6
Psychic Shock	5	6,2
None	1	1,2
Total	81	100

Table 3. Table of correlation between antibiotic use and CKD.

Treatment : Antibiotic/Other	N	%
No	5	6,7
Yes	70	93,3
Total	75	100
Frequent Use of Antibiotics		
No	27	33,3
Yes	54	66,7
Total	81	100

Table 4. Table of correlation between Chronic use of chemical medicines and CKD

Chronic Use of Chemical Medicines	N	%
No	14	17,3
Yes	67	82,7
Total	81	100

Table 5. Table of correlation between recurrent urinary infection and CKD.

Chronic Use of Chemical Medicines	N	%
No	41	56,9
Yes	31	43,1
Total	72	100

Table 6. Table of correlation between CKD and family history.

Family History of Kidney Disease	N	%
No	62	76,5
Yes	19	23,5
Total	81	100

Table 7. Table of correlation between CKD and sexually transmitted diseases.

History of Sexually Transmitted Diseases	N	%
No	55	67,9
Yes	26	32,1

Table 8. Table of correlation between the disease (Chronic Kidney Disease) and recurrent acute infections, for patients of the total sample.

Disease	Recurrent Acute Infections n (%)		Total	
	No	Yes		
CKD	5 (12,2)	76 (79,2)	81	
				$\chi^2= 53,4$
				p-value<0.001

Table 9. Table of correlation between the disease (CKD) and the use of chemical drugs for patients of the total sample.

Disease	Use of Chemical Drugs n (%)		Total	
	No	Yes		
CKD	14 (20,9)	67 (95,7)	81	
				$\chi^2= 79,98$
				p-value<0.001

Table 10. Table of correlation between the disease and the availability of the causative factors, for patients of the total sample.

Disease	Causative Factors n (%)			Total	
	No	Psychic Shock	Physical Shock		
CKD	1 (1,23)	5 (6,2)	75 (92,6)		
				81 (100)	Fisher's exact
					p-value <0.001

Discussion

Chronic, non-communicable diseases are by far a major cause of mortality worldwide [5-7]. CKD is representative of such chronic conditions. It is characterized by its gradual progression and mainly by its numerous etiological factors. The progression of CKD does not consistently depend on the extent and the severity of the original cause/disease. Patients with CKD, even when the cause of kidney damage is not active, present with a deterioration of the renal function, and it is only in rare cases that they recover to a normal kidney function. These observations indicate that it is not only the original cause that contributes to the progression of CKD but also various other factors, such as the diet, environmental conditions, physical status (defense mechanism), exposure to toxic agents, and age. A characteristic example in kidney failure is the structural and functional impairment of the nephrons, which are unable to maintain the body's homeostasis [22-26]. The homeopathic approach considers that, in patients with CKD, miasms can potentially affect the body's structure. This way, in some CKD cases, the behavior of the "sick" system is predefined by its composition. End stage CKD is a chronic syndrome, resulting from one or more chronic diseases and leading to patients having to take long-term medicines for the treatment of the underlying diseases.

The effectiveness of homeopathy has been established in numerous studies for a broad range of acute and chronic diseases [27]. A prospective study including 3,981 elderly subjects showed that the course of chronic diseases improved in a marked and sustained way in patients who received homeopathic treatment [28]. Its results strengthened the view that homeopathic treatment can play an important role in patients with chronic diseases. According to another meta-analysis, homeopathy is also effective in treating chronic allergies [29]. Of particular interest is a pooled study reviewing the 83 most important clinical trials conducted in

the last thirty years regarding treatment of acute and chronic diseases. The authors prove that homeopathy is a safe and effective treatment [30]. Furthermore, a comparative cohort study comparing homeopathy and conventional medicine for treating acute respiratory tract and ear infections concluded that homeopathy is not inferior [31]. In addition, according to a nationwide study conducted in Switzerland, patients receiving homeopathic treatment in primary care were significantly satisfied compared to those receiving conventional medicine. The study also concluded that the risk of side effects in homeopathic treatment is two or three times lower than in conventional medicine [32]. Furthermore, a study conducted in Germany observed no medical toxicity or poisoning in children inadvertently taking big quantities of homeopathic remedies [33].

Not only do homeopathic treatments present a low risk of adverse reactions, the low cost of homeopathic preparations is low and they distinguish from the concept of placebo [29, 34]. In contrast, chronic use of chemical medicines can be interpreted from a homeopathic view as a phenomenon of "symptom suppression," given that patients may develop terminal kidney failure [20]. In addition, it is well documented that a vast number of antimicrobial medicines administered for the treatment of urinary tract infections also cause nephrotoxicity [35-61]. For example, the use of nitrofurantoin administered as a long term chemical prophylaxis in recurrent urinary tract infections led to acute granulomatous interstitial nephritis [52]. The use of antimicrobial medicines in patients with kidney disorders, as well as their repeated use in recurrent infections, clearly constitutes an aggravating factor, leading to renal impairment.

Often, however, long-term administration of different chemical medicines is necessary for the treatment of CKD, despite adverse reactions. The epidemiology of renal impairment-inducing medicines is complex, and the exact mechanism of

drug nephrotoxicity is unknown [51, 62]. The administration of chemical preparations and vaccines seemingly causes a complex internal process with the participation of various organs, resulting in manifestation of toxicity. Chronic use of chemical substances/medicines can result in a status of health where not only there is no cure, but side effects and other pathologies may also be caused [35, 62-64]. This may also be the case for the regular vaccination of patients with chronic diseases who develop chronic renal impairment. As proven in daily medical practice, a large number of patients do not know they are in early stages of renal impairment and take nephrotoxic antibiotics for forthcoming infections, thus further deteriorating their renal function. As earlier mentioned, homeopathic remedies are known to have fewer side effects and lower nephrotoxicity. More specifically, in our study, 92.6% of subjects with CKD had other concomitant diseases, given that CKD is a chronic syndrome. Notably, 1% to 5% of patients receiving non-steroidal anti-inflammatory drugs develop nephrotoxic syndrome, and analgesic nephropathy is one of the chronic renal diseases leading to end stage CKD [65].

Thus, the concern arises whether analgesics and other chemical medicines have a substantial therapeutic effect, as well as whether symptoms of chronic diseases are potentially suppressed as a result of their abuse. Many patients with CKD receive immunosuppressive therapy with corticoids making them susceptible to recurrent infections, which in turn are treated with antibiotics. Thus there is a vicious circle of the disease's manifestation and treatment. Since the nephrotoxicity mechanisms of various medicines have not yet been fully elucidated, impairment of renal function is observed in a number of patients receiving chemical medicines for a certain period of time. Further studies should be conducted to elucidate which are the causative factors for renal impairment in certain patients following use of chemical medi-

nes. Despite the advancements in pharmacology, there is a potential inability to justify the factors causing nephrotoxicity in a particular group of patients. According to homeopathic theory, the disease outcome pathways and the exact causative factors should be considered during the progression of a disease for which suppressive therapy is administered. Urinary tract infections and especially recurrent febrile urinary tract infections (such as pyelonephritis) can lead to impairment of renal function. For example, recurrent urinary tract infections due to the bacterium *P. mirabilis* cause permanent renal parenchyma lesions, given that they form infection stones [66]. Studies have also shown that recurrent urinary tract infections, infection stones, high blood pressure, and diabetes mellitus appear to be high risk factors for developing chronic renal impairment. The presence of infections and recurrent infections in patients with impaired renal function can lead to CKD. Furthermore, patients with chronic renal impairment have an imbalanced immune system, which is a direct result of renal impairment as well as the primary disease that led to its manifestation [67].

Patients with recurrent infections and CKD often receive chemical medicines and/or immunosuppressants, such as corticosteroids and antibiotics, to treat their underlying conditions. In an effort to interpret the frequent relapses of infections from a homeopathic perspective, a potential miasmatic dimension of the disease arises. According to homeopathy, chronic predispositions are the reason why patients develop relapses despite receiving the correct treatment. Notably, in the phenomenon of symptom/disease "suppression," patients with CKD address these infection relapses by taking chemical medicines, such as antibiotics, antipyretics, and anti-inflammatory agents. Hahnemann states that by treating acute infections with "allopathic" therapies and with high doses of medicines, local symptoms of the diseases are suppressed, and the risk of chronic disease

is increased [15-16]. Furthermore, by aiming at balancing the body's vital reaction, homeopathic treatment seeks a holistic physiological response of the body. As a result, this approach is indicated for the treatment of both acute infections and chronic diseases [30, 68-71].

Sexually transmitted diseases are conditions of the urogenital system, often first manifesting themselves as skin conditions, such as in HPV infection, genital herpes, and syphilis [22-66]. They often manifest with relapses and, as a result with very few exceptions (e.g., genital warts), the use of antibiotics is needed to treat them. Once again, antibiotics have potential nephrotoxic properties. Approaching sexually transmitted diseases with the homeopathic concept of "disease suppression" ("when a disease is not treated in depth and causally, then it is simply suppressed"), any disease presenting itself in the human body must be thoroughly cured [19]. By treating the symptoms, one does not treat the disease itself. It is simply a symptomatic treatment of the health disorder, not a complete cure of the disease. By analyzing the health course of patients with CKD and a history of sexually transmitted diseases, it is not a course leading to cure, as stated in Hering's Law, "from a more important organ to a less important one." [72]

On the contrary, by holistically analyzing patients with CKD and STD, one observes that they lead to the manifestation of deeper pathologies. Often, the suppressive treatment of a disease has damaging effects on the body. The suppressed disease tends to affect deeper structures and to settle in more important tissues and organs. According to Hahnemann, initial exposure to miasms causes local symptoms, such as skin or venereal diseases; if however these symptoms are suppressed by chemical medication, the cause—the miasm—goes deeper and begins to manifest itself as disease of the internal organs. Therefore, one could argue that there is the potential of manifestation of a miasmatic condition in patients with CKD and a history of

sexually transmitted diseases [15, 16]. Furthermore, the deoxyribonucleic acid (DNA) of viruses, bacteria, and microorganisms leave an "imprint," a traceable frequency in the body fluids, which opens the way for the development of a highly sensitive detection system for chronic bacterial infections in human and animal diseases [73].

Our study shows that a large number of patients with CKD have a history of sexually transmitted diseases. More specifically, by focusing on gonorrhea with regards to the potential development of pre-disposition/miasmatic disposition in some patients, it is known that the disease may present recurrent manifestations. Gonorrhea tends to relapse with no apparent development of protective immunity arising from previous episodes of infection. Thus, a new hypothesis emerges regarding gonococcal infections [74]. *Neisseria gonorrhoeae* has been associated with humans for several millennia. It has adapted remarkably well to the human immune system and, accordingly, has evolved to evade destruction by the host's immune defenses. Regarding treatment of sexually transmitted diseases, the use of homeopathic remedies is recommended. Each year, the number of CKD patients as well as those with sexually transmitted diseases is increased. A characteristic example in kidney failure is the structural and functional impairment of the nephrons, which are unable to maintain the body's homeostasis. This could be true for patients with CKD, in which miasmatic conditions potentially affect the body's structure. Accordingly, in some CKD cases, the behavior of the "sick" system is predefined by its composition.

Conclusions

In conclusion, the multifactorial nature of CKD and the complexity of the human body should be approached both through the documented research of conventional medicine as well as by homeopathy. Further studies in patients with CKD and

other chronic diseases should be conducted using and comparing the abovementioned approaches and treatments. Analysis through the homeopathic theory of disease suppression, as well as that of miasms, provide a new approach, while offering a future prospect for conducting studies aimed at finding new causative factors in patients with renal impairment who later develop end stage CKD. New data and answers may therefore arise regarding the manifestation and course of renal impairment. Homeopathy provides an alternative treatment solution, with lower nephrotoxicity rates and proven effectiveness. Taking into account the polypharmacy of patients with CKD, as well as their susceptibility to recurrent infections due to immunosuppression, there is a real need for further investigation for the comparative study of chemical medicines used in conventional medicine and homeopathic preparations.

Abbreviations

CKD: Chronic kidney disease; HPV: Human Papillomavirus; STD: Sexually transmitted disease; DNA: Deoxyribonucleic acid.

Competing interests

The authors declare that they have no competing interests.

References

1. Daskalakis N, Winn M: Focal and segmental glomerulosclerosis. *Cell Mol Life Sci* 2006, 63: 2506-2511.
2. Medical Encyclopedia: Idiopathic pulmonary fibrosis. MedlinePlus. Retrieved 2007-02-13.
3. Asimov I: *The Human Body: Its Structure and Operation*. Boston: Houghton Mifflin; 1963.
4. Vithoulkas G, Carlino S: The "continuum" of a unified theory of diseases. *Med Sci Monit* 2010, 16: SR7-15.
5. National Center for Chronic Disease Prevention and Health Promotion: *Chronic Disease Overview*. Atlanta; 2012.
6. Register C: *Living with chronic illness: days of patience and passion*. New York: Bantam Books; 1989.
7. World Health Organization: *Chronic diseases*. 2011.
8. Vithoulkas G, Tiwari SK: *The Science of Homoeopathy*. New Delhi: B. Jain Publishers; 2002.
9. Vithoulkas G: *Homeopathy - Medicine for the New Millennium*. Athens: International Academy of Classical homeopathy; 2000.
10. Vithoulkas G: *The Science of Homeopathy*. USA: Grove Press; 1980.
11. King S: Miasms in homeopathy. *Classical homeopathy*, archived from the original on 2009-03-07, retrieved 2009-03-25.
12. Miasma in Webster Dictionary [<http://www.merriam-webster.com/dictionary/miasma>].
13. Little D: Miasm in *Classical homeopathy* [<http://www.simillimum.com/education/little-library/constitution-temperaments-and-miasms/mch/article.php>].
14. Speight P: *Comparisons of the Chronic Miasms*. London: Double & Brendon Ltd; 1977.
15. Hahnemann S: *Die chronischen Krankheiten, ihre eigenthümliche Natur und homöopathische Heilung [The chronic diseases, their specific nature and homoeopathic treatment]*. Dresden and Leipzig: Arnoldische Buchhandlung; 1828.
16. Hahnemann S: *The Organon of the Healing Art*. Dublin: W.F.Wakeman; 1833.
17. Hahnemann S: Essay on a new principle for ascertaining the curative powers of drugs, together with some views of the past. [*Hufeland's*] *J pract d Arzkd* 1796, 2: 391-439.
18. Hahnemann S: Fingerzeige auf den homöopathischen Gebrauch der Arzneien in der bisherigen Praxis. [*Hufeland's*] *N J d pract Arzkd* 1807, 26: 5-43.
19. Vithoulkas G: *The Science of Homeopathy*. New York: Grove Press; 1980.
20. Teixeira MZ: Is there scientific evidence that suppression of acute diseases in childhood induce *chronic diseases* in the future? *Homeopathy* 2002, 91: 207-216.
21. Hahnemann S: *The Organon of Medicine*. 6th edition Dudgeon's 5th with Boericke's 6th added.
22. Longo D, Fauci A, Kasper D, Hauser S, Jameson J, Loscalzo J: *Harrison's Principles of Internal Medicine (18th Ed)*. New York: McGraw-Hill; 2011.

23. Jacobson HR, Klahr S, Striker GE: *The Principles and Practice of Nephrology (2nd Ed)*. St. Louis: Mosby-Year Book; 1995.
24. Quirós PL, Ceballos M, Remón C, Hernández Romero MC, Benavides B, Pérez Pérez-Ruilópez MA, Lozano A, Aznar E, Rivero M, Fernández Ruiz E: Systemic arterial hypertension in primary chronic glomerulonephritis: prevalence and its influence on the renal prognosis. *Nefrologia* 2005, 25: 250-257.
25. McPhee SJ, Hammer GD: *Pathophysiology of Disease - An introduction to clinical Medicine (6th Ed)*. New York: McGraw-Hill Medical; 2009.
26. Luke RG, Strom TB: Chronic renal failure. In *Internal Medicine*. 4th edition. Edited by Stein JH. St Louis: Mosby; 1994: 2622-2645.
27. Reilly DT, Taylor MA, McSharry C, Aitchison T: Is Homeopathy a placebo response? *Lancet* 1986, 2: 1272.
28. Teut M, Lüdtke R, Schnabel K, Willich SN, Witt CM: Homeopathic treatment of elderly patients-a prospective observational study with follow-up over a two year period. *BMC Geriatr* 2010, 10: 10.
29. Ullman D, Frass M: A review of homeopathic research in the treatment of respiratory allergies. *Altern Med Rev* 2010, 15: 48-58.
30. Bellavite P, Marzotto M, Chirumbolo S, Conforti A: Advances in Homeopathy and immunology: a review of clinical research. *Front Biosci (Schol Ed)* 2011, 3: 1363-1389.
31. Haidvoogl M, Riley DS, Heger M, Brien S, Jong M, Fischer M, Lewith GT, Jansen G, Thurneysen AE: Homeopathic and conventional treatment for acute respiratory and ear complaints: a comparative study on outcome in the primary care setting. *BMC Complement Altern Med* 2007, 7: 7.
32. Marian F, Joost K, Saini KD, von Ammon K, Thurneysen A, Busato A: Patient satisfaction and side effects in primary care: an observational study comparing Homeopathy and conventional medicine. *BMC Complement Altern Med* 2008, 8: 52.
33. Zuzak TJ, Rauber-Lüthy C, Simões-Wüst AP: Accidental intakes of remedies from complementary and alternative medicine in children-analysis of data from the Swiss Toxicological Information Centre. *Eur J Pediatr* 2010, 169: 681-688.
34. Rossi E, Crudeli L, Endrizzi C, Garibaldi D: Cost-benefit evaluation of homeopathic versus conventional therapy in respiratory diseases. *Homeopathy* 2009, 98: 2-10.
35. Choudhury D, Ahmed Z: Drug-induced nephrotoxicity. *Med Clin North Am* 1997, 81: 705-717.
36. Kampmeier RH: Introduction of sulfonamide therapy for gonorrhoea. *Sex Transm Dis* 1983, 10: 81-84.
37. Rule AD, Krambeck AE, Lieske JC: Chronic kidney disease in kidney stone formers. *Clin J Am Soc Nephrol* 2011, 6: 2069-2075.
38. Bowie WR: Approach to men with urethritis and urologic complications of sexually transmitted diseases. *Med Clin North Am* 1990, 74: 1543-1557.
39. Bowie WR, Floyd JF, Miller Y, Alexander ER, Holmes J, Holmes KK: Differential response of chlamydial and ureaplasma-associated urethritis to sulphafurazole (sulfisoxazole) and aminocyclitols. *Lancet* 1976, 2: 1276-1278.
40. Bowie WR, Alexander ER, Stimson JB, Floyd JF, Holmes KK: Therapy for nongonococcal urethritis: double-blind randomized comparison of two doses and two durations of minocycline. *Ann Intern Med* 1981, 95: 306-311.
41. Hook IB, Goldstein RS (Eds): Overview of clinical nephrotoxicity. In *Toxicology of the Kidney*. 2nd edition. New York: Raven Press; 1993.
42. Namagondlu G, Low SE, Seneviratne R, Banerjee A: Acute renal failure from nitrofurantoin-induced acute granulomatous interstitial nephritis. *QJM* 2010, 103: 49-52.
43. De Ruyter A, Thin RN: Genital herpes - A guide to pharmacological therapy. *Drugs* 1994, 47: 297-304.
44. Diaz-Mitoma F, Sibbald RG, Shafran SD, Boon R, Saltzman RL: Oral famciclovir for suppression of recurrent genital herpes: a randomized controlled trial. Collaborative Famciclovir Genital Herpes Research Group. *JAMA* 1998, 280: 887-892.
45. Deck AJ, Shapiro RH, Berger RE: Epididymo-orchitis. In *The current treatment of infectious diseases*. 1st edition. Edited by Schlossberg D. St Louis: Mosby; 2000.
46. Harrison WO, Hooper RR, Kilpatrick ME, et al: Penicillin-resistant gonorrhoea: Alternative Therapy. In Seigenthaler W
47. Luke RG, Strom TB: Chronic renal failure. In *Internal Medicine*. 4th edition. Edited by Stein JH. St Louis, Mosby; 1993.
48. Kampmeier RH: Introduction to sulfonamide therapy for gonorrhoea. *Sex Transm Dis* 1983, 10: 81-84.
49. Whelton A: Nephrotoxicity of nonsteroidal anti-inflammatory drugs: physiologic foundations and clinical implications. *Am J Med* 1999; 106: 135-245.
50. Werner M, Costa MJ, Mitchell LG, Nayar R: Nephrotoxicity of xenobiotics. *Clin Chim Acta* 1995; 237: 107-154.
51. Skinner R: Nephrotoxicity-what do we know and what don't we know? *J Pediatr Hematol Oncol* 2011; 33: 128-134.
52. Namagondlu G, Low SE, Seneviratne R, Banerjee A: Acute renal failure from nitrofurantoin-induced acute granulomatous interstitial nephritis. *QJM* 2010; 103: 49-52.
53. Naughton CA: Drug-induced nephrotoxicity. *Am Fam Physician* 2008; 78: 743-750.
54. Dahanukar SA, Thatte UM, Deshmukh UD, Kulkarni MK, Bapat RD: The influence of surgical stress on the psychoneuro-endocrine-immune axis. *J Postgrad Med* 1996; 42: 12-14.
55. Luthy R (eds): *Current chemotherapy*. Washington, DC, American Society for Microbiology, (1978)

56. Kleinknecht D, Landais P, Goldfurb B: Drug associated acute renal failure. A prospective multicenter report. *Proc EDTA-ERA* 1988; 22: 1002.
57. Judson FN: Treatment of uncomplicated gonorrhea with ceftriaxone: A review. *Sex Transm Dis* 1986; 13(Suppl 3): 199-202.
58. Urol SJ: Infection-induced urinary calculi and renal failure. *Nephrol* 1987; 21: 219-223.
59. Coufalik ED, Taylor-Robinson D, Csonka GW: Treatment of nongonococcal urethritis with rifampicin as a means of determining the role of urealyticum Ureaplasma. *Br J Vener Dis* 1979; 55: 36-43.
60. Hoosen AA, O'Farrell N, van den Ende J: Microbiology of acute epididymitis in a developing community. *Genitourin Med* 1993; 69: 361-363.
61. Morin JP, Fillastre JP, Olier B: Antibiotic nephrotoxicity. *Chemioterapia* 1984; 3: 33-40.
62. Taber SS, Pasko DA: The epidemiology of drug-induced disorders: the kidney. *Expert Opin Drug Saf* 2008; 7: 679-690.
63. Babl FE, Lewena S, Brown L: Vaccination-related adverse events. *Pediatr Emerg Care* 2006; 22: 514-519.
64. Taber SS, Mueller BA: Drug-associated renal dysfunction. *Crit Care Clin* 2006; 22: 357-374.
65. Harirforoosh S, Jamali F: Effect of inflammation on kidney function and pharmacokinetics of COX-2 selective nonsteroidal anti-inflammatory drugs rofecoxib and meloxicam. *J Appl Toxicol* 2008; 28: 829-838.
66. McDougal WS, Wein AJ, Kavoussi LR, Novick AC, Partin AW, Peters CA, Ramchandani P: *Campbell-Walsh Urology (10th Ed)*. Philadelphia: Saunders; 2011.
67. Pesanti EL: Immunologic defects and vaccination in patients with chronic renal failure. *Infect Dis Clin North Am* 2001; 15: 813-832.
68. Witt CM, Lüdtke R, Baur R, Willich SN: Homeopathic treatment of patients with chronic low back pain: A prospective observational study with 2 years' follow-up. *Clin J Pain* 2009; 25: 334-339.
69. Pomposelli R, Piasere V, Andreoni C, Costini G, Tonini E, Spalluzzi A, Rossi D, Quarenghi C, Zanolin ME, Bellavite P: Observational study of homeopathic and conventional therapies in patients with diabetic polyneuropathy. *Homeopathy* 2009; 98: 17-25.
70. Brien SB, Leydon GM, Lewith G: *Homeopathy* enables rheumatoid arthritis patients to cope with their chronic ill health: a qualitative study of patient's perceptions of the homeopathic consultation. *Patient Educ Couns* 2012; 89: 507-516.
71. Witt CM, Lüdtke R, Mengler N, Willich SN: How healthy are chronically ill patients after eight years of homeopathic treatment?-Results from a long term observational study. *BMC Public Health* 2008; 8: 413.
72. Reappearance of old symptoms during Homeopathic Treatment's intervention. Papamethodiou D Master Thesis (2011).
73. Montagnier L, Aïssa J, Ferris S, Montagnier JL, Lavallée C: Electromagnetic signals are produced by aqueous nanostructures derived from bacterial DNA sequences. *Interdiscip Sci* 2009; 1: 81-90.
74. Liu Y, Feinen B, Russell MW: New concepts in immunity to *Neisseria gonorrhoeae*: innate responses and suppression of adaptive immunity favor the pathogen, not the host. *Front Microbiol* 2011; 2: 52.

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