

A STUDY TO ASSESS THE VALIDITY OF APPLIED KINESIOLOGY (AK) AS A DIAGNOSTIC TOOL AND AS A NONLOCAL PROXIMITY EFFECT

Stephan A. Schwartz,ⁱ Jessica Utts, PhD,ⁱⁱ S. James P. Spottiswoode,ⁱⁱⁱ Christopher W. Shade, PhD,^{iv} Lisa Tully, PhD,^v William F. Morris, DO,^{vi} and Ginette Nachman, MD, PhD^{vii}

ABSTRACT

Applied Kinesiology is a diagnostic technique widely used within the Integrative Medical community. This study asks: 1. Is there a difference in muscular strength when an individual holds a substance that is inimical to life processes (a poison solution), as compared to a substance that is essential for life (normal saline)? 2. Is this effect a transaction involving input from both the person being measured, and the kinesiologist doing the measurement, or is it only the person being measured? 3. As an extension of question 2, is the result the same when different kinesiologists take the measurement, or when no kinesiologist is involved? 4. Does belief, expectation, gender, or time cognition play a role in determining response? To answer these questions 51 participants were tested during three trials each, first by one kinesiologist, then by another and, finally, with no kinesiologist present by grip strength indicated using a hand dynamometer. For each trial a pair of randomly numbered sealed vials, each pair in a randomly numbered plastic bag, were used as the objects of the trial. In each bag one vial contained saline solution while the other was filled with a slightly smaller amount of saline solution to which had been added ionic hydroxylamine hydrochloride (NH₃OH)⁺, producing a toxic solution of 9 mg/ml. Each trial consisted of a separate muscle test for each vial. All present at the trials were blind as to which vial contained the toxin. And all who prepared the vials were blind to the trials. The force used by the kinesiologists in each of their trials was measured via a pressure pad system. The hand dynamometer trials were conducted with no kinesiologist present. Results: Of the 151 sets of trials the toxic vial was identified correctly in 80 of them (53%), resulting in a one-tailed exact binomial p-value of 0.258. Results for two of the kinesiologists were almost exactly at chance. For the third kinesiologist there was a one-tailed exact binomial p-value of 0.18 (unadjusted for multiple testing). Results for the dynamometer were also almost exactly at chance. Testing whether there was a significant difference in proportions for whom the AK test worked based on belief about whether it would work resulted in non-significant chi-square values of 0.6 ($p = 0.439$) for the trials with one kinesiologist, and 0.222 ($p = 0.136$) for the hand dynamometer trials. The final variable examined was gender. While there was no significant difference in performance for males and females for the trials of the male kinesiologist or the hand dynamometer, the combined data for the two female kinesiologists did reveal a difference. Of the 33 sessions with females, only 15 were successful (45%) while for the 18 sessions with males, 14 were successful (78%) resulting in a chi-square statistic of 4.96, $p = 0.026$. However, given all of the chi-square tests performed in this section, the results must be interpreted with caution because of multiple testing. Results indicate belief in whether or not the AK test will work was not significantly related to whether or not it actually did work. A chi-square test of the relationship between time perception and correct vial choice showed no significant relationships. The chi-square statistic for the relationship using the hand dynamometer data was 0.927, p -value = 0.629. The data in this study, particularly when seen in the larger context of a review of the literature from the AK field itself by Klinkoski and Leboeuf (1990), which considered 50 papers published between 1981 and 1987 by the International College of Applied Kinesiology, and the survey by Hall, Lewith, Brien, and Little (2008), using standard evaluation criteria (QUADAS, STARD, JADAD and CONSORT), for research methodology, as well as five prior non-clinical studies, by Radin (1984), Quintanar and Hill (1988), Braud (1989), Arnett, Friedenber, and Kendler (1999), and Kendler and Keating (2003), all together suggest: The research published by the Applied Kinesiology field itself is not to be relied upon, and in the experimental studies that do meet accepted standards of science Applied Kinesiology has not demonstrated that it is a useful or reliable diagnostic tool upon which health decisions can be based.

ⁱ Scholar-in-Residence, Atlantic University and Research Associate, Cognitive Sciences Laboratory

ⁱⁱ Professor, Department of Statistics, University of California – Irvine

ⁱⁱⁱ Managing Director and Chief Scientist, The Resource Group

^{iv} Director of International Scientific Advisory Board, Applied Biophysics Foundation

^v President and Chief Scientist, Quicksilver Scientific

^{vi} Chairman, Department of Osteopathic Manipulation Medicine, Touro College of Osteopathic Medicine

^{vii} Medical Writing Scientist, Quintiles Transnational Corporation

Correspondence: saschwartz@earthlink.net

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Project Background and Conception:

There are literally thousands of health practitioners in complementary and alternative medicine (CAM) — allopathic MDs, homeopathic MDs, osteopathic physicians, naturopaths, chiropractors, nurses, dentists, physiotherapists, and body workers — using a muscle testing protocol known as Applied Kinesiology (AK) as a diagnostic aid. This technique traces back to the 1940s and an American husband and wife physiotherapist couple, Henry and Florence Kendall, who developed a clinical procedure to ascertain the strength of a muscle, and postural procedures allowing them to isolate and test the strength of a specific muscle independent of the surrounding musculature.¹

Two decades later a Michigan chiropractor, George J. Goodheart, Jr., picked up this approach and redefined it.² Goodheart made a clinical report on a patient who came to him with an immobile shoulder that had troubled him for more than 15 years. Goodheart pressed on small nodules near the origin of the pain and the muscle strength returned to normal as did the range of motion. This led Goodheart to develop a technique for “tugging” on particular muscular trigger points to restore function.² As his clinical work continued he came to believe his muscle testing technique also had a diagnostic aspect. This aspect has become the dominant use of AK and is the focus of this study.



Trial in progress. Note participant holds vial in left hand while being tested. (1)

Typically, a patient is asked to hold his or her arm extended out from the shoulder and parallel to the floor in the coronal plane and to resist the downward pressure of the practitioner’s hand while the patient is holding or exposed to a test substance. The relative strength difference, with or without exposure to the substance, supposedly helps the kinesiologist to assess systemic imbalances, i.e., whether or not a food sensitivity exists. The practitioner might also press on key “trigger points” to find out if they cause muscle weakness. The presumption is that relative weakness is a sign that the substance, or one of the substances, is not good for that individual system. The underlying concept of AK

assumes a “proximity effect” in which holding a substance — sometimes in a bottle, or encased in a capsule, or pill — as opposed to actually ingesting it is sufficient to determine the effect of the substance on the individual being tested.

Applied Kinesiology has been refined over the past thirty years, both in Europe and in the United States, and today is used by many thousands of people as a feedback tool to assist in confirming and gathering psycho-physiological information. Professionally, certification as an Applied Kinesiologist is available from the International College of Applied Kinesiology³ and from the Touch for Health organization.⁴ To reach the highest level of certification, over 300 hours of instruction, several proficiency exams, and submission of original research papers are required. But, although there are a wealth of clinical reports⁵ and several well-known books, both academic⁶ and for the general audience,⁷ with muscle testing as their topic, the fundamental question about its diagnostic validity has rarely been directly addressed by properly controlled, randomized, and blinded experimentation.

In a review of the literature from the AK field itself, Klinkoski and Leboeuf considered 50 papers published between 1981 and 1987 by the International College of Applied Kinesiology.⁸ Twenty of these papers were classified as research papers, and Klinkoski and Leboeuf evaluated them using seven criteria they felt represented accepted standards for research methodology. These included: identification of sample size, inclusion criteria, blind and naive subjects and statistical analysis. They concluded, “Although some papers satisfied several of these criteria, none satisfied all seven of them. As none of the papers included adequate statistical analyses, no valid conclusions could be drawn concerning their report of findings.”⁸

In 2008, Hall, Lewith, Brien, and Little took an even more detailed look at the AK literature. They were explicitly interested in ascertaining whether AK offered “inter-examine reliability,” and to determine “whether there was evidence for its therapeutic effectiveness.”⁹ To do this analysis they used standard recognized evaluation tools. “Diagnostic accuracy studies were analyzed and scored for methodological quality and quality of reporting using the quality assessment tool for studies of diagnostic accuracy included in systematic reviews (QUADAS) and the Standards for Reporting of Diagnostic Studies (STARD). Clinical studies were analyzed for methodological quality using the JADAD scale and for quality of reporting using the Consolidated Standards of Reporting Trials (CONSORT).”⁹

Twenty-two papers were evaluated by Hall *et al* in this way. Their first finding was that overall the methodology of the papers was poor. “QUADAS scored 1-11 out of a possible 14, STARD scores were between 6-13 out of 25, JADAD scores were all 0 out of 5 and CONSORT 4-6 out of 22.” They felt that with these scores none of the questions that had prompted their study could be answered. They concluded by stating: “We recommend a pragmatic study of the effectiveness of kinesiology as the most appropriate initial step to determine whether kinesiology has any clinical value.”⁹

Five studies by researchers with no vested interest in AK as a clinical technique have also examined the premise of AK in a laboratory setting, and we believe should also be considered. Although none used the accepted kinesiological clinical protocol, indeed were not medical studies per se at all, we cite them because they did meet the standards of methodological rigor, i.e., blindness and randomness, etc., expected in good science, and they did address a core claim of AK.

All involved testing whether muscles were stronger or weaker in proximity to sucrose. The results were contradictory. Radin, in 1984, carried out a double-blind experiment with 58 adults testing the hypothesis that proximity to sugar reduces human grip strength. To test this hypothesis participants were asked to hold bottles identical in appearance and weight that were filled with either sugar or sand. An analysis of variance supported the hypothesis.¹⁰

Four years later, Quintanar and Hill, carried out a double-blind experiment with 90 undergraduate students in an attempt to replicate Radin’s results, using a similar protocol. When “additional controls were added to this design to reduce random sampling error, the hypothesis was not supported.”¹¹

A year later, in 1989, Braud carried out another replication of Radin’s work, employing a double-blind experiment with 50 adults. Grip strength of the dominant hand was measured by a chart-recording hand dynamometer for 12 trials. For half of the trials, the subject held an opaque, sealed bottle containing sugar; for the other half of the trials, the subject held a control bottle of equal weight

containing sand. Statistical analysis indicated no significant difference in grip strength when holding sugar versus holding sand.¹²

Ten years later, Arnett, Friedenber, and Kendler, tested the hypotheses that close physical proximity to sucrose, without its ingestion, affected muscle strength adding the additional variable of time between meals as potentially influencing the direction of this effect. Twenty eight college students performed 10 one-arm curls using the dominant arm, while holding a bottle containing either sucrose or sand in the free hand, without knowledge of the bottle's contents. Data were first collected following an overnight fast. A week later, data were collected one hour after these students had each consumed a bagel. During the fasting state, the mean maximal isometric strength while holding the bottle of sucrose was significantly greater than when holding a bottle of sand; however, no significant difference was found for the group tested after having consumed a bagel.¹³ This study did not counterbalance the order of conditions, nor did it control for the participants' amount of experience with AK.

In 2003, Kendler and Keating carried out the final precursor to the study that is the subject of this paper. Again the test was the effect of proximity to sucrose. For a sample of 76 participants a double-blind, counterbalanced assessment of grip strength was performed with and without holding a container of sucrose. No significant effect was found for proximity of glucose.¹⁴

We believe there are two major criticisms of these five studies, particularly when they are considered as an assessment of Applied Kinesiology; the current study addresses these issues as indicated:

1. These studies were all grip strength or arm curl studies carried out in non-clinical settings, and not truly tests of AK, as it is practiced in thousands of treatment rooms. With no sacrifice of methodological rigor, this study was designed so that both the set and setting of the trials were equivalent to what takes place in the treatment room of a working health professional.
2. In all of these studies cited the central variable governing the participant response was the reaction of individuals to sugar in some form. But sugar means something quite different to an athletic young individual than it does to a middle-aged, overweight, type II diabetic. None of the studies controlled for this. This study eliminates this as a confounding variable because it is structured to be as unequivocally antipodal in its conditions and the universality of reactions as possible — normal saline solution versus poison in normal saline solution.

Objectives:

The objectives of this study are to answer the following four questions:

Is there a difference in muscular strength when an individual holds a substance that is inimical to life processes, as compared to a substance that is essential for life?

If there is an effect, does the process involve input from both the person being measured and the clinician doing the measurement, or is it only the person being measured? This might help us understand the differences in results amongst the four earlier studies.

Is the result the same when different clinicians take the measurement, or when no clinician is involved?

Can AK be considered as a reliable diagnostic tool?

This last question deserves a further comment. In a study such as this one, which addresses issues in both basic research and clinical medicine, there were really three possible outcomes: 1) The null hypotheses were supported; 2) statistically significant results occurred but fell short of clinical significance; and, 3) the data supported diagnostic levels of confidence.

Methodology:

The fundamental assumption of all AK sessions involving the test of something is that substances that are inimical to life generally, or for the particular individual being tested particularly, in some degree produce weakness in the muscle tested, in comparison with control samples.

This experiment explores the proposition in a deliberately typical clinic setting to increase the clarity of the outcome. In this study one half of the vials contain a pure saline solution tainted by a notable poison, by definition life degrading, while the other half of the vials serve as controls and contain only the pure saline water solution, by definition essential to life. If the effect is ever going to occur it should reasonably be expected to occur in a binary experiment in which the two conditions being tested are as antipodal as the differences in this one.

Personnel:

Kinesiologists. There were three kinesiologists who conducted the trials in this study. By design both male and female kinesiologists participated, to see whether gender was a factor.

The male kinesiologist originally trained as a civil engineer, and worked in that capacity for a number of years, prior to becoming a chiropractic physician. He completed his chiropractic training at the National University of Health Sciences and, then, trained under programs of the International College of Applied Kinesiology, completing their standard course in 1992/93, with subsequent training at regular seminars thereafter. Notably he received much of this training from Dr. Walter Schmitt a protégé of AK's founder Dr Goodheart.

The first female kinesiologist received her chiropractic training at Palmer College of Chiropractic in 1997 and, subsequently received her certification for AK from the International College of Applied Kinesiology. She has maintained a chiropractic practice incorporating AK for a decade. This female kinesiologist served as a tester for eight trials. Subsequently, a second female kinesiologist served as the second tester.

The second female kinesiologist is a Mechanical Engineer who subsequently trained as an Applied Physiology and Touch For Health Certified Instructor. She worked as a Parts and Dimensional Engineer for about ten years in the large structural investment casting industry supplying jet engine parts to several manufacturers before obtaining instructor certification in Applied Physiology in 2004 from the Touch For Health Organization. She has maintained a private practice for five years, centering on Applied Physiology. Her involvement permitted the study to compare the results of two quite different approaches to Muscle Testing.

Participants. Fifty one (51) adult volunteers, 33 women, and 18 men took part in the 151 trials, volunteering an average of 2.5 hours, approximately what a first visit to a holistic clinician might entail. They were recruited by announcements that the study sought participants to test, and were self-

selecting. They ranged in age from their 20s to 70s. All had graduated from high school. In addition, nine (9) had some college, 10 held undergraduate degrees. Eleven (11) held Masters level degrees, two (2) were RNs, four (4) held doctorates. One declined to answer the question on education. All but two participated in all three trial conditions. These two did not take part in the hand dynamometer session because of a family emergency. Thirty three (33) had been previously tested by AK in the course of a clinical visit, 18 had no prior experience with AK.

In addition to basic demographics the study also included measures of:

1) **Time Perception.** Since neither the kinesiologist nor the participant knew which vial contained a toxin, and no actual direct contact with either the contents of the poison tainted vials or the pure saline controls occurred some form of nonlocal awareness must occur for AK to work. Schmeidler¹⁵ and Dean¹⁶ in 1974 both reported studies in which the Time Metaphor Test of Knapp and Garbutt¹⁷ positively correlated with nonlocal awareness skills. Schwartz and De Mattei¹⁸ found that individuals who self-assign themselves as time Dynamics perform less accurately than either Non-dynamics or Neutrals when they are asked to do a task involving the kind of nonlocal awareness implicit in this study. Participants in this study were asked to select a metaphor that most closely represented their understanding of time: Time is a rushing river (Dynamic). Time is an old woman spinning (Neutral). Time is a still mountain pool (Non-dynamic).

Time as Metaphor*

Dynamics	Neutrals	Non-dynamics
26	16	8
52 %	32 %	16%

*Note: One participant could not make a decision as to how he viewed time.

(Table 1)

2) **Sheep/Goat.** The now classic Sheep/Goat Effect was first reported by Schmeidler and McConnell (who coined the terms), and this belief effect is now recognized as one of the most consistently determinative variables in the intention research literature.¹⁹ Sheep — those who believe that whatever is being tested in an experiment *in which they are taking part* — is likely to work generally achieve better success than goats, who are skeptics.

Sheep /Goat

AK Can Work			AK Will Work In My Trial		
Not sure	No	Yes	Not Sure	No	Yes
18	0	33	9	1	41
35%	0%	65%	18%	2%	80%

(Table 2)

Researchers. There were seven researchers in this study each carrying out various tasks. Each was blind to the aspects of the study, in which they did not personally participate.

1) **Principal Investigator.** The Principal Investigator was present for all the trials which were held in Virginia Beach, Virginia. The PI designed the study and oversaw the conduct of the 151 sessions, including running the computer and the pressure sensor system, the hand dynamometer, and the video camera. The PI transferred the data from the signed affidavit forms signed by each participant at the end of each vial test, as well as the affidavits signed by the kinesiologists, when trials involved their participation. This data was then transferred by the PI into a single electronic database, which was sent

to Researchers 5 and 6. The PI was blind to all aspects of the vial preparation, as well as their numbering, and the numbering of the plastic bags in which the vials were placed.

2) **Researchers 2 and 3.** Researchers 2 and 3, were in Boulder, Colorado. These researchers prepared the vials, both those with the poison and the controls. They also numbered them. They were blind to which vials were allocated to which trial, and the actual trial sessions.

3) **Researcher 4.** Researcher 4 was in New York, New York, and received the vials from Researchers 2 and 3, as well as the code revealing which were controls and which contained the poison. Researcher 4 first randomized the vials and then put a numbered control vial and a numbered treated vial in small plastic bags, and numbered the bags from 1 to 150, each bag representing one trial. The vial-filled bags were then packed in a box and sent to the PI in Virginia Beach. Researcher 4 retained the codes showing which vials were controls and which contained the poison, and which from each category had been assigned to which bag. Researcher 4 was completely blind to the conduct of the actual trials.

4) **Researcher 5.** Researcher 5 was in Beverly Hills, California and carried out the analysis of the pressure pad data, which was then sent to Researcher 6. Researcher 5 was blind to all aspects of the study until all the trial data had been collected.

5) **Researcher 6.** Researcher 6 was in Davis, California and received the databases from the PI and Researcher 5. Researcher 6 analyzed these data for correlations with outcomes and demographics. Researcher 6 was blind to all aspects of the study's conduct, until all the trial data had been collected.

6) **Researcher 7.** Researcher 7 was in Durham, North Carolina and participated in the design of the study, and conducted 17 trials over two (2) sessions. Researcher 7 was blind to vial preparation and numbering. Travel distance and the complexities of scheduling made it impossible for Researcher 7 to participate in the study after the second session.

Human Experimentation:

Institutional Review Board (IRB). Pursuant to the U.S. *Code of Federal Regulations — Title 45 — Public Welfare — Department of Health and Human Services — National Institutes of Health — Office for Protection from Research Risks — Part 46 — Protection of Human Subjects*, a conforming Institutional Review Board (IRB) was established.

Informed Consent Form. Pursuant to Title 45 [Revised 18 June 1991, Effective 19 August 1991] participants in the study each signed an Informed Consent Form, which had been approved by the IRB, and which complied with all relevant guidance paragraphs of Title 45, Part 46.

Experimental Set and Setting:

Experimental Area. Because one of the purposes of this study was to approximate as closely as possible the set and setting in which Applied Kinesiology is actually carried out, and because the majority of health practitioners utilizing AK are doctors of chiropractic, the trials for this study were all conducted in a treatment room of a working chiropractic clinic in Virginia Beach, Virginia. Participants filled out their paperwork, just as they would on their first visit to the clinic. They waited in the waiting room, just as they would as patients, and were called in, one-by-one, just as patients would be in a clinical practice.

Vial Preparation:

Vial Preparation. Researchers 2 and 3 prepared the 300 vials at the laboratories of Quicksilver Scientific, LLC in Lafayette, Colorado.



A set of vials, one control, one with poison, each randomly numbered, and randomly placed in the plastic bag that was, then, randomly given a trial number.

(2)

for mice is 408 mg/kg and for rats it is 120 mg/kg. 150 test vials were filled with 6.0 ccs of isotonic (0.9% m/v) hydroxylamine hydrochloride ($\text{NH}_2\text{OH}\cdot\text{HCl}$, ACS Plus Grade, Fisher Scientific) solution and randomly and uniquely numbered.

This toxin was chosen for several reasons: 1) It is unequivocally a poison to any human system; 2) It would dissolve in water without changing the water's sensorial characteristics — no visual change, no smell, etc. — so that nothing would reveal which vial contained the toxin; 3) It could be disposed of with relative ease; and 4) Even if an accident were to occur, it would not be an unmanageable threat. Although highly toxic, in the form and potency used, as an ionic $(\text{NH}_3\text{OH})^+$ solution at a pH of ~ 3.2 , very little absorption through skin would have happened even if a vial had broken, so there was a very small margin of risk even should exposure occur. Only actually drinking the contents of the vial would have exposed participants to any significant risk, and at the level of the solution used and the quantity involved, even that would have presented a nonlethal threat. However, from the premise of AK, it clearly allowed a sharply contrasting condition when compared with the untainted saline solution.

Numbering. The vials were randomly numbered as one lot, then the vials with the toxin were boxed in one box, and those with just the untainted saline solution in another. The two boxes were sent to Researcher 4 in New York where one from each box was randomly taken and placed in a plastic bag. When this was completed the bags were randomly numbered with the trial for which they were intended. This produced 150 bags, each containing one toxin and one untainted vial.

Pressure Sensor:

Tact Array System. To continuously measure the force being exerted by the kinesiologists throughout a trial a tactile pressure sensor system manufactured by PPS of Los Angeles, California., was used. The



The pressure pad installed prior to trial.

(3)

system is stretchable and fits snugly over the palm of the hand. It is USB powered for Windows PC's and supported by proprietary PC-resident software which displays and tracks time-series, average, and peak pressures.

Hand Dynamometer:

Baseline Hand Dynamometer. The Jamar Hydraulic Hand Dynamometer gives accurate grip strength readings without the subject being able to “feel” the handle move. The results are reliably consistent with published Baseline and Jamar® studies. The internationally accepted design assures reliability, user convenience, and measurement repeat-ability. The five position adjustable handle can

accommodate any hand size. The maximum reading is recorded by a slave hand that moves with the reading hand, and remains in place until the unit is reset to zero. The strength reading can be viewed as pounds or kilograms. This permitted measurement of any grip strength differences between the vials.



Jamar Hydraulic Hand Dynamometer

(4)

Video:

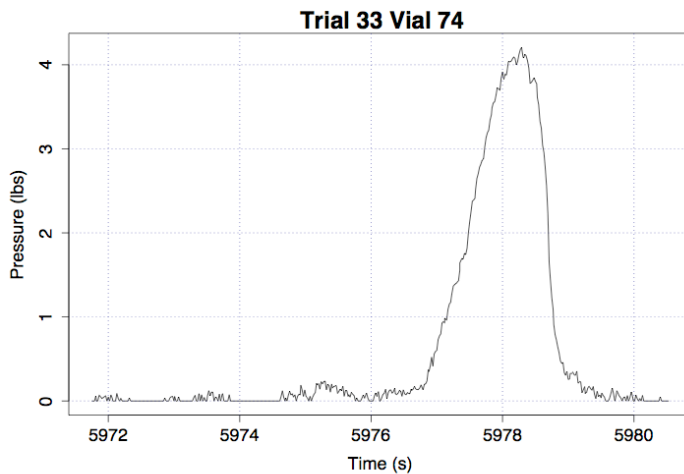
To preserve a permanent record of all trials, each was recorded in its entirety on a Sony Handycam model DCR-HC38 Digital Video Camera Recorder.

Methodology:

In order to approximate the experience of a patient in a clinic, the trials were designed for groups of 3-9 individuals per session. In that way the experience was very much like a first visit to a new physician. There were nine sessions in all, spread across parts of four months with each session running from two to three hours.

Prior to beginning participants were all briefed, their questions were answered, and they were asked to read and sign the informed consent form, and to fill out a questionnaire. The video camera was started, the pressure sensor was calibrated, and the treatment room readied to begin. By arrangement, as this was happening, the first kinesiologist arrived in the building, and was wired with the pressure sensor system by the Principal Investigator, whereupon the trials began in the treatment room, with the first participant.

The kinesiologist tested that individual's muscle strength, using their AK techniques, until satisfied that the participant was in a condition to proceed with the test. Prior to each trial the pressure sensor would be zeroed out and allowed to run for a few seconds showing a zero baseline. When this was established the PI told the kinesiologist to begin. At the end of the trial for each vial the PI, the kinesiologist, and the participant, each independently filled out, dated, and signed, a form attesting to which was used, and which produced the weaker response. At the end of each trial the vials were returned to their plastic bag, all paperwork was collected, and that trial was at an end.



A typical pressure record from a trial. Graph (1)

The first participant would leave the treatment room, the next participant was asked to enter it, and the same process proceeded until the last participant had been tested. When the entire group for the session had been tested by the first kinesiologist, that kinesiologist left the building, and the second kinesiologist, by prior arrangement, entered. There would be a break while the new kinesiologist was wired for the pressure sensor. Then the trial cycle began and the patients, in the same sequence rotation, went through a second time.

Then the second kinesiologist left the building, there was another short break, and the third trial cycle, this one with just the hand dynamometer, began and proceeded in the same sequence. On the hand dynamometer trials, the participants held the vials, asking themselves “Is this harmful to me?” pointing the dynamometer dial towards the video camera and, when they felt it was appropriate, squeezed the grip. The strength in pounds was recorded, both on video and on the trial form, the instrument was reset, and the second vial was tested in the same way, and the grip strength recorded.

Hypotheses:

Affirmative Hypothesis One: There will be a significant difference in the level of weakness associated with the presence of the toxin as compared to the presence of the normal saline solution. There will be a statistically significant correct perception of the vials with the toxin in which weakness is associated with the presence of the toxin.

Affirmative Hypothesis Two: The effect will occur under all three conditions of testing.

Affirmative Hypothesis Three: We hypothesize that if Applied Kinesiology is a viable diagnostic test that there should be no statistically significant difference in outcome regardless of which kinesiologist performs the test.

Affirmative Hypothesis Four: We hypothesize that for Applied Kinesiology to be a viable diagnostic test that that there will be no statistically significant difference arising from gender, belief, or time cognition.

Data Analysis and Results:

Data collected: For each of the 51 participants, the demographic and belief categorical variables listed in the section on “participants” were measured. During the experiment, 49 of the 51 participants contributed results for three pairs of vials, with one toxic vial and one non-toxic vial in each pair, assigned in random order. All 51 participants worked with kinesiologist PT for one pair of vials, and with either kinesiologist JA or JE for another pair. Forty-nine of the 51 participants worked with the hand dynamometer for the third pair of vials. Therefore, there were a total of 150 pairs of vials tested. For each pair of vials, it was determined whether or not the participant correctly identified the toxic vial by noting if the trial with that vial produced a weaker response than the trial with the pure saline vial.

In addition, for the two sets of trials with the kinesiologists, continuous pressure data was measured. Four measurements were derived from each of these trials: Maximum pressure, mean pressure, median pressure and integral pressure (the area under the pressure curve). These results can be used to determine if the kinesiologist, perhaps unconsciously, detected which was the toxic vial because, if so, more pressure should have been applied with the toxic vial than with the non-toxic vial. (See Graph 1 above for an example of continuous pressure data measurement.)

Results for Pressure Data:

Under the null hypothesis that it is not possible to detect which is the toxic vial, the 151 sets of trials represent a binomial experiment with $n = 151$ and $p = \frac{1}{2}$, where p is the probability of correctly selecting the toxic vial.²⁰ Under the alternative hypothesis the probability of selecting the toxic vial is higher, thus the alternative hypothesis is $p > \frac{1}{2}$. Of the 151 sets of trials the toxic vial was identified correctly in 80 of them (53%), resulting in a one-tailed exact binomial p -value of 0.258. Although the result is in the correct direction, it is not statistically significant. Results for kinesiologists PT and JE were almost exactly at chance, with 26 out of 51 correct and 4 out of 8 correct, respectively. Results for the dynamometer were also almost exactly at chance, with 25 correct out of 49 tests. For kinesiologist JA there were 25 correct guesses out of 43 tries, resulting in a one-tailed exact binomial p -value of 0.18 (unadjusted for multiple testing).

The continuous pressure data was used to test whether the kinesiologists are likely to apply more pressure when the participant is holding a toxic vial than when holding a non-toxic vial. Although a complex model could be used to combine results for all three kinesiologists, the most direct way to answer the questions of interest is to conduct separate paired t-tests for each pressure measure and each kinesiologist. For each test, the null hypothesis is that the population mean difference in the pressure measurement is 0. The alternative hypothesis is that the population mean difference is greater than 0, indicating that the pressure is greater when the participant holds a toxic vial than with a non-toxic vial.

The results of the 12 t-tests are shown in Table 3. For kinesiologist JE, all results were in the opposite direction from the alternative hypothesis, with more pressure applied with the non-toxic vials than with the toxic ones. If a one-tailed test in the opposite direction had been used, the t-test for maximum pressure alone would have been statistically significant. However the sample size was so small ($n = 6$) that no conclusions should be made about what would happen if a larger sample were available. For kinesiologists PT and JA, the mean and median pressure results are suggestive, but because so many tests were done, the p -values need to take multiple testing into account. For these two practitioners it appears that the maximum pressure applied over the time period was very similar, on average, for the toxic and non-toxic vials.

One-tailed t-Test Results for Pressure Measurements

Kinesiologist	Mean pressure	Median pressure	Max pressure	Integral pressure
Male (n = 50)	$t = 1.63$ $p = 0.055$	$t = 2.14$ $p = 0.019$	$t = 0.00$ $p = 0.50$	$t = 0.94$ $p = 0.176$
Female 1 (n = 6)	$t = -0.97$ $p = 0.812$	$t = -0.03$ $p = 0.513$	$t = -2.42$ $p = 0.970$	$t = -1.37$ $p = 0.885$
Female 2 (n = 43)	$t = 1.30$ $p = 0.100$	$t = 1.33$ $p = 0.095$	$t = -0.35$ $p = 0.635$	$t = -0.59$ $p = 0.720$

(Table 3)

Results for Belief Data:

Participants were asked whether or not they thought the experiment would work, and 33 said yes while 18 said they weren't sure. No one said no. Table 4 shows whether or not the experiment did work for that participant, for the session with the male kinesiologist and the session with the hand dynamometer. Testing whether there is a significant difference in proportions for whom it worked based on belief about whether it would work resulted in non-significant chi-square values of 0.6 ($p = 0.439$) for the trials with the male kinesiologist, and 2.222 ($p = 0.136$) for the hand dynamometer trials. There were too few trials with the other kinesiologists to do a meaningful chi-square test. These results indicate that belief in whether or not the experiment will work is not significantly related to whether or not it actually does work. Results were similar for the question about whether or not kinesiology could work in general.

Belief in Whether AK Test Would Work and Whether AK Test Did Work

Will work?	Male Kinesiologist		Hand Dynamometer	
	Did work	Did not work	Did work	Did not work
Not sure	4	6	3	7
Yes	22	19	22	17

(Table 4)

As described in the section on participants, they were asked about their time perception, categorized as dynamics, neutrals and non-dynamics. A chi-square test of the relationship between time perception and correct vial choice showed no significant relationships. For example, the chi-square statistic for the relationship using the hand dynamometer data was 0.927, p -value = 0.629.

The final variable examined was gender. While there was no significant difference in performance for males and females for the trials with the male kinesiologist or the hand dynamometer, the combined data for the two female kinesiologists did reveal a difference. Of the 33 sessions with females, only 15 were successful (45%) while for the 18 sessions with males, 14 were successful (78%) resulting in a chi-square statistic of 4.96, $p = 0.026$. However, given all of the chi-square tests performed in this section, the results must be interpreted with caution because of multiple testing.

Discussion:

The objectives of this study were to answer the following four questions:

Is there a difference in muscular strength when an individual holds a substance that is inimical to life processes, as compared to a substance that is essential for life?

If there is an effect does the process involve input from both the person being measured, and the clinician doing the measurement, or is it only the person being measured? This might help us understand the differences in results amongst the four earlier studies.

Is the result the same when different clinicians take the measurement, or when no clinician is involved?

Can AK be considered as a reliable diagnostic tool?

The task proposed by Hall, Lewith *et al* (2008): “We recommend a pragmatic study of the effectiveness of kinesiology as the most appropriate initial step to determine whether kinesiology has any clinical value,”⁹ appears to have been accomplished, and the result is that it lacks clinical value.

The data in this study, particularly when considered in the larger context of the five prior non-clinical studies, of Radin (1984), Quintanar and Hill (1988), Braud (1989), Arnett, *et al* (1999), and Kendler and Keating (2003), and the even larger horizon of the 50 papers assessed by Klinkoski and Leboeuf (1990), and the 22 papers examined by Hall, Lewith *et al* (2008), all seem to suggest one common conclusion: Applied Kinesiology does not presently provide a useful or reliable diagnostic tool upon which health decisions can be based.

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²⁰ By mistake the vials for trial 101, were used twice with different participants, several weeks apart in time. Bag 101 was the last trial in session, using the hand dynamometer in Session 5, and the first trial in Session 6 involving kinesiologist Female 2. All conditions of blindness obtained in both sessions. This duplication makes no meaningful difference to the outcome statistics, and the authors felt it was important to report exactly what happened. Thus the references elsewhere to "151 trials".