Do TETRA (Airwave) Base Station Signals Have a Short-Term Impact on Health and Well-Being? A Randomized Double-Blind Provocation Study

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BACKGROUND: “Airwave” is the new communication system currently being rolled out across the United Kingdom for the police and emergency services. It is based on the Terrestrial Trunked Radio Telecommunications System (TETRA). Some police officers have complained about skin rashes, nausea, headaches, and depression as a consequence of using their Airwave handsets. In addition, a small subgroup in the population self-report being sensitive to electromagnetic fields (EMFs) in general.

OBJECTIVES: We conducted a randomized double-blind provocation study to establish whether short-term exposure to a TETRA base station signal has an impact on the health and well-being of individuals with self-reported “electrosensitivity” and of participants who served as controls.

METHODS: Fifty-one individuals with self-reported electrosensitivity and 132 age- and sex-matched controls participated in an open provocation test; 48 sensitive and 132 control participants went on to complete double-blind tests in a fully screened semianechoic chamber. Heart rate, skin conductance, and blood pressure readings provided objective indices of short-term physiological response. Visual analog scales and symptom scales provided subjective indices of well-being.

RESULTS: We found no differences on any measure between TETRA and sham (no signal) under double-blind conditions for either controls or electrosensitive participants, and neither group could detect the presence of a TETRA signal at rates greater than chance (50%). When conditions were not double blind, however, the self-reported electrosensitive individuals did report feeling worse and experienced more severe symptoms during TETRA compared with sham.

CONCLUSIONS: Our findings suggest that the adverse symptoms experienced by electrosensitive individuals are due to the belief of harm from TETRA base stations rather than to the low-level EMF exposure itself.

KEY WORDS: electromagnetic fields, electromagnetic hypersensitivity, electrosensitivity, idio-pathic environmental intolerance, mobile phone. Environ Health Perspect 118:735–741 (2010). doi:10.1289/ehp.0901416 [Online 14 January 2010]

“Airwave” is the new communication system currently being rolled out across the United Kingdom for the police and emergency services. It is based on the Terrestrial Trunked Radio Telecommunications System (TETRA), which offers enhanced digital communication that improves emergency services’ performance and public safety. However, a growing number of concerns have been raised about the possible adverse health effects related to this new technology. For example, some police officers have complained about skin rashes, nausea, sleeplessness, headaches, and depression as a consequence of using their Airwave handsets (Farrel 2002; Police Federation News 2005). Others have complained about unusual symptoms, which they attribute specifically to TETRA base station signals. In addition, there is a small subgroup of persons in the general population who believe they suffer from “electrosensitivity.” As the term suggests, sufferers believe that their symptoms are caused by devices emitting electromagnetic fields (EMFs), and anecdotal evidence points to TETRA as the source of some of these symptoms. Prevalence rates for self-reported electrosensitivity vary. For example, in Iran the prevalence rate is currently available on the possible impact of TETRA on health. Randomized double-blind studies on Global System for Mobile Communication (GSM) and Universal Mobile Telecommunications System (UMTS) have reported no reproducible evidence of adverse health effects for either healthy controls or individuals with perceived electrosensitivity (Eltiti et al. 2007a; Regel et al. 2006; Rubin et al. 2006). Moreover, neither electrosensitive nor healthy individuals can detect the presence of EMFs at rates greater than chance (Eltiti et al. 2007a; Kwon et al. 2008; Regel et al. 2006; Rubin et al. 2006, 2010; Zwamborn et al. 2003). In a recent study using functional magnetic resonance imaging Landgebre et al. (2008) showed that the subjective perception of symptoms was triggered by the anticipation of being exposed to an EMF. This finding led to the implication that these symptoms are due to beliefs and anticipation of harm rather than to the EMFs themselves. Consequently, the World Health Organization has suggested that terms such as “electrosensitivity” should be replaced by IEI-EMF to acknowledge the absence of evidence for a causal connection.

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between EMFs and reported symptoms (Hansson Mild et al. 2006).

The health concerns regarding TETRA handsets and base stations are predominantly focused on the 17.6-Hz pulse frequency feature and its possible impact on biological systems [Advisory Group on Non-Ionising Radiation (AGNIR) 2001]. In 2001, the AGNIR reviewed all of the available evidence, including calcium efflux, brain waves, and epilepsy. The group concluded that any effect on biological systems would require a nonlinear biological interaction that operated on the time scale of the carrier frequency that could demodulate the amplitude-modulated component (Challis 2005; NR PB 2001; Valberg et al. 2007). Given this, the AGNIR concluded that TETRA was highly unlikely to pose a health risk but nevertheless recommended that well-designed laboratory studies be conducted (NRPB 2001). Only two studies investigating TETRA technology have been published since the AGNIR report was released. Green et al. (2005) found no evidence that TETRA handsets produced significant changes in calcium physiology in the brain as a consequence of TETRA exposure, whereas Barker et al. (2007) reported that neither GSM nor TETRA handset signals affected blood pressure or any other physiological parameter. The present study is the first report of the short-term health effects of TETRA base stations.

The aims of the present study were to determine whether symptoms reported by IEI-EMF suffers are caused by short-term exposure to radiofrequency (RF) EMFs as produced by TETRA base station signals and to determine whether exposure to these signals affects a selection of the adult population that do not report sensitivity to EMFs. We tested TETRA base station signals on both healthy controls and IEI-EMF suffers under open provocation and double-blind conditions while measuring a range of objective and subjective indicators of well-being, as well as the participants’ ability to detect the presence or absence of the signal.

Materials and Methods

Participants. All participants completed the Electromagnetic Hypersensitivity Questionnaire (EHQ) (Elititi et al. 2007b) prior to testing, which provided an assessment of their current state of health and whether participants attributed their symptoms to EMFs. Volunteers in the sensitive group were selected on the basis of a self-report of being sensitive to EMFs, particularly those produced by mobile communication handsets and base stations. Volunteers in the control group confirmed that they were not sensitive to EMFs. Volunteers were excluded if they fulfilled one or more of the following criteria: were outside the age range of 18–80 years, had a history of brain injury, had a diagnosis of epilepsy or claustrophobia, had received a diagnosis or any treatment for a mental disease, had been fitted with a pacemaker, or had any physical impairment or illness or were taking any medication that could have, in some way, affected the results of the study. Participants were recruited through local and national newspaper advertisements, letters of invitation, specialist health publications, personal recommendation, and Web site contact; they were reimbursed their travel expenses and received a small payment. All testing took place between April 2007 and January 2009.

Figure 1. Diagram of the exposure chamber.

Design. A mixed design was used in which two groups (between subjects: sensitive and controls) were exposed to two exposure conditions (within subjects: TETRA and sham). Each participant took part in an open provocation session and two double-blind sessions. Each session was spaced at least 1 week apart and was conducted at about the same time of day. Subjective well-being and physiological functioning were measured during each session throughout the exposure conditions.

The initial open provocation session consisted of two tests where both the experimenter and the participant knew when the base station was “on” and when it was “off.” This was followed by a brief double-blind test comprising four short-duration trials with two on and two off conditions. Sessions 2 and 3 comprised the double-blind component of the study, with just one exposure (TETRA or sham) being administered during each of these sessions. All participants received both exposure conditions counterbalanced across sessions 2 and 3. Exposure conditions were block-randomized within group and programmed into the exposure system control computer by an external consultant (Belloulle 2008). Double-blinding remained in place until completion of the data collection phase.

Our sample size was calculated on the assumption that RF-EMFs has a small effect on human health (d = 0.40). Hence, 66 participants per group would yield statistical power of 0.90 for within-subjects effects, which allowed us to be 90% confident about differences found between sham and TETRA exposure conditions. The same level of confidence for between-subjects effects (i.e., group by exposure condition interaction) would require 132 participants per group.

Materials and equipment: screened semi-anechoic chamber. The testing room had a shielding effectiveness between 55 and 60 dB at 420 MHz. The appropriate depth absorber (300 mm) was used to achieve a uniform field at 420 MHz and conformed to the BS EN 61100-4-3:2006 “on” + A1: 2008 specification (National Physical Laboratory, Teddington, Middlesex, UK). Participants were seated 4.95 m from the antenna of the base station. A screen was placed 2.8 m from the participant, blocking the antenna from view, and was used to back-project task instructions for the participant during testing. The projector was positioned outside the testing room at the back wall, and projection was made through a screened window behind the antenna. A screened window in the near wall allowed visual contact between experimenter and participant (Figure 1).

Exposure system. We used the TETRA signal release 1 [specification 390 392-2; European Telecommunications Standards Institute (ETSI), Sophia-Antipolis Cedex,
France), which comprised a time-division multiple-access frame structure with four time slots per frame on a single carrier. The chosen frequency was 420 MHz with a 25-kHz bandwidth. The signal was emitted at a power flux density of 10 mW/m² (uncertainty estimate, 1 dB) over the area in which the participant was seated. Calculating a value for the Specific Absorption Rate (SAR) for whole body exposure is extremely complex; the variation in both the electrical properties of individual tissues and their spatial distributions means that SAR will never be uniform across the tissues of the body. In addition, SAR calculations should ideally be based on measurements using either a physical or numerical phantom, neither of which was available to us. Nevertheless, we would like to provide an estimated SAR for the present study. We calculated an approximate corresponding SAR for the power level used with this formula: body surface area × power flux density + body weight. We used assumed values for body surface area and body weight; thus, for a person with a body surface area of 1.9 m² and a body weight of 70 kg, the approximate corresponding SAR for this power level is 271 µW/kg (1.9 × 10mW/70 kg). The maximum power produced by TETRA base station transmitters is comparable to those of mobile phone base station transmitters (NRPB 2001). In the absence of any other available evidence, we broadly based the power level for this study on one conducted by Mann et al. (2000) who measured signal strengths of mobile phone base stations around the United Kingdom. With reference to the International Commission on Non-Ionizing Radiation Protection (1998) guidelines the power level used in the present study is 0.5% of the 2.1W/m² reference level for the general public. The properties of the TETRA signal were modeled directly on the TETRA system that is currently used by the emergency services, thus replicating as closely as possible actual base-station emissions. Whether or not traffic is carried on the channel has a profound effect on the waveform. Thus, to allow for the harmonics of the time slots, a ratio of time slot occupancy of 50% traffic/no traffic was applied over the period of the tests. The sham condition comprised a no-signal condition. [A copy of the technical reference manual is available upon request (Belloul 2008).]

Regular calibrations confirmed that power density levels remained within 1 dB variability tolerance throughout the study period. The field uniformity was independently tested and verified by the National Physical Laboratory.

**Biographical information.** Participation began with a medical history interview, which provided biographical information and assessed participants’ state of health prior to testing.

**Subjective well-being.** Visual analog scales (VASs) and symptom scales were used to measure subjective well-being. The VAS comprised a 10-cm line, with 0 cm representing “not at all” and 10 cm “extremely,” for anxiety, tension, arousal, relaxation, discomfort, and fatigue. The corresponding descriptors were “anxious,” “tense,” “agitated,” “relaxed,” “discomfort,” and “tired.” Participants were asked to mark anywhere on the line corresponding to how they felt at that time. The symptom scales were derived from the EQH (Eltiti et al. 2007b) and comprised 57 symptoms on a 5-point scale (“not at all” to “a great deal”); participants were required to select how much they were suffering from each symptom.

**Physiological measures.** Physiological measures were being assessed using the means ± SDs for blood volume pulse (BVP), heart rate (HR), and skin conductance (SC) recorded throughout the open provocation and double-blind tests. We recorded physiological data using a ProComp Infiniti eight-channel encoder with Biograph Infiniti software [version 2.0.1; Thought Technology Ltd., Plattsburgh, New York, USA (2003)] that we ran on a Dell Latitude notebook (Dell Products UK, Dublin, Ireland). Signals were sampled at a rate of 2,048 samples/sec for BVP and 256 samples/sec for SC. The BVP was submitted to a fourth-order Butterworth low-pass filter with a 10-Hz cutoff frequency. The HR was calculated from the filtered BVP by calculating the time locations for the BVP peaks and valleys based on the locations on which the derivative of the BVP reached zero (dicrotic notches were ignored by the algorithm). HR was then estimated based on the time between peaks: HR = 1/(interpeak interval). All signals were resampled at 8 samples/sec to have a uniform rate. BVP signals were detrended because the important information in this signal was on the peak-to-peak values (Eltiti et al. 2007a).

“On”/“Off” judgments. There were six double-blind on/off judgments: four 5-min trials after the open provocation component and two 50-min trials. Participants were required to judge whether the base station was “on” or “off” and to indicate whether their confidence was “low,” “moderate,” or “high” directly after each trial.

**Procedure.** The open provocation test and the 5-min double-blind tests were conducted in session 1. Sessions 2 and 3 were double-blind and contained one exposure condition each. Ordering of exposure condition was randomized across participants. Table 1 shows the procedures in detail.

**Statistical analyses.** We adjusted the alpha levels using the Bonferroni correction to control for type 1 error. Analysis of variance (ANOVA) was used to measure mean differences of data with normal distributions. Where the distributions were skewed and could not be normalized, we used nonparametric methods instead; Mann-Whitney U-tests were used for between-group comparisons, and Wilcoxon signed rank tests for within-group comparisons. Nonparametric tests lack the flexibility of ANOVAs, which calculate interaction effects. To mimic this type of analysis using nonparametric methods,

| Table 1. Procedures for open provocation and double-blind tests. |
|---------------------------------------------------------------|
| **Task** | **Duration** |
| Session 1 |  |
| Background information | Medical history interview, psychologic questionnaires, WAIS-R verbal memory test |
| Open provocation (e.g., sham, TETRA) | VAS completed every 5 min, symptoms reported, physiologic measurements taken continuously |
| Cognitive tests | Participants completed Backward Digit Span Task and Letter Cancellation Task |
| Double-blind trials (e.g., sham, sham, TETRA, TETRA) | At the end of each of the four short-duration trials, participants specified whether they believed the base station had been “on” or “off” and how confident of this judgment they were |
| Session 2 and 3, double-blind (e.g., session 2, TETRA; session 3, sham) | Exposure per session (TETRA/sham) Physiologic measurements were taken continuously over the entire exposure period |
| Low load | Participants watched Planet Earth DVD, completed VAS every 5 min, and recorded any symptoms |
| High load | Participants performed an operation span task; the task was interrupted every 5 min to allow for the completion of VAS and symptom scales |
| Cognitive tests | Participants completed Backward Digit Span Task and Letter Cancellation Task |
| On/off judgment | Participants made a judgment as to whether the base station was “on” or “off” |
we performed between-subjects analyses on difference scores.

Results

Biographical information. Fifty-one “sensitive” participants and 144 control volunteers completed at least one session (Figure 2). However, during the testing, we lost data for five controls because of technical failures, and one person withdrew after session 1. Additionally, we tested controls more quickly than sensitives and found that the ratio of males to females was significantly higher for the controls compared with the sensitive group. Therefore, we tested an additional six females and randomly selected six males (Figure 2). Our final sample of 183 (51 sensitives, 132 controls) included 10 emergency service workers, of these, two were sensitive participants. An independent-samples t-test showed that the sensitive groups (mean ± SD = 42 ± 16; range, 18–73) and control groups (41 ± 19; range, 18–80) were comparable in terms of age (t[181] = 0.289, p > 0.77) and sex (sensitives, 61% female; controls, 51% female; χ²(1) = 1.487, p > 0.2).

![Flow of participants through each stage of testing.](image)

The EHQ responses indicated that 11 (21.6%) sensitive participants reported being “a little bit” sensitive, 10 (19.6%) “moderately,” 11 (21.6%) “quite a bit,” and 12 (23.5%) “a great deal.” Six (11.8%) reported that they were not sensitive to EMFs but specified EMF-emitting objects as being associated with their symptoms. We conducted telephone interviews with these participants to establish eligibility to participate as sensitive before attending session 1. One person in the sensitive sample did not complete the EHQ before participating but was invited after a telephone interview. Assuming that any differences between these subgroups would be greatest under open provocation conditions, we performed Kruskal–Wallis tests on the open provocation physiological measures, VASs, and symptom scales. The outcomes for all these comparisons suggest that these groups were broadly comparable (χ²(4) ≤ 5.659, p > 0.28).

Table 2. Additional health and social behavior information [n (%)].

| Variable | Sensitive Yes | Control Yes | Sensitive No | Control No | χ²
|---------|--------------|-------------|--------------|-------------|--------|
| Chronic illness | 5 (9.8) | 15 (11.4) | 46 (80.2) | 117 (88.6) | 0.002 |
| Headache-proneb | 24 (47.1) | 25 (18.9) | 27 (52.9) | 100 (75.6) | 11.9* |
| Taking medication | 21 (41.2) | 59 (44.7) | 30 (58.8) | 73 (55.3) | 0.07 |
| Electric shockc | 15 (29.4) | 23 (17.4) | 34 (66.7) | 109 (82.6) | 2.99 |
| Smoker | 6 (11.8) | 14 (10.6) | 45 (88.2) | 118 (89.4) | 0 |
| Consumes alcohol | 41 (80.4) | 108 (81.8) | 10 (19.6) | 24 (19.2) | 0 |

*Values reflect χ² with Yates continuity corrections; p-values with or without this correction are very similar. bMissing data for 7 (5.3%) control participants. cMissing data for 2 (3.9%) sensitive participants. *p < 0.008 (Bonferroni corrected α = 0.008).
participants, independent of group, felt worse during TETRA compared with sham. Table 3 shows that, under TETRA conditions compared with sham, controls reported significantly higher levels of anxiety, tension, arousal, and discomfort and lower levels of relaxation during TETRA compared with sham. Analyses of difference scores (TETRA – sham) revealed that sensitive participants reported a significantly greater increase in levels of anxiety, tension, arousal, and discomfort during the TETRA exposure compared with controls.

Under double-blind conditions, we found no significant overall difference between groups (p ≥ 0.1). For exposure condition, relaxation alone remained as significant (z = –2.713, p = 0.007), with higher relaxation being reported during TETRA (mean ± SE = 6.76 ± 0.12) compared with sham (6.54 ± 0.11). Table 4 shows that, when controlling for baseline, however, both within- and between-group comparisons revealed no significant differences for any of the variables.

**Symptom scales.** We also analyzed total symptom score, which measures symptom severity, and the total number of symptoms reported by means of nonparametric statistics.

During the open provocation test, we found an overall difference between groups for both the total symptom score (z = –5.985, p < 0.001) and total number of symptoms reported (z = –6.448, p < 0.001), indicating that sensitive participants experienced a greater severity (mean ± SE = 5.92 ± 1.23, n = 51) and number of symptoms (4.63 ± 0.86, n = 51) than did controls (1.04 ± 0.17 and 0.92 ± 0.13, respectively; n = 132). We also found an overall difference for exposure condition (TETRA, sham) for symptom severity only (z = –2.255, p = 0.024), suggesting that all participants experienced greater symptom severity when exposed to TETRA (2.89 ± 0.552) compared with sham (1.91 ± 0.322). The total number of symptoms reported remained comparable across exposure conditions (z = –1.874, p > 0.06). Control participants reported no differences between exposure conditions for either symptom severity or total number of symptoms reported (Table 5). In contrast, the sensitive participants reported a significantly higher total symptom score and more symptoms during TETRA than during sham. The between-group analyses of difference scores (TETRA – sham) revealed a significant difference for both total symptom score and total number of symptoms reported (Table 5).

Under double-blind conditions, we found a significant difference between groups for the total symptom score (z = –5.917, p < 0.001) and total number of symptoms reported (z = –5.282, p < 0.001), with sensitive participants reporting greater symptom severity (mean ± SE = 5.23 ± 1.06, n = 48) and number of symptoms (4.03 ± 0.71, n = 48) compared with controls (1.20 ± 0.18 and 0.92 ± 0.10, respectively; n = 132). We found no overall effect for exposure. Thus, sensitive reported a greater severity and number of symptoms.

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### Table 3. Descriptives (means ± SEs) and statistical tests (z-scores) for the open provocation VAS for sensitive and control participants.

| VAS measure  | Sham       | TETRA       | Difference score<sup>a</sup> | TETRA vs. sham | Sensitive<sup>b</sup> vs. Control<sup>c</sup> |
|--------------|------------|-------------|-----------------------------|----------------|-----------------------------------|
| Anxiety      | 1.51 ± 0.22 | 0.99 ± 0.10 | 2.24 ± 0.27                 | 0.72 ± 0.16    | 0.13 ± 0.07                        |
| Tension      | 1.51 ± 0.23 | 1.00 ± 0.10 | 2.31 ± 0.29                 | 0.80 ± 0.18    | 0.20 ± 0.06                        |
| Arousal      | 1.54 ± 0.22 | 0.96 ± 0.11 | 2.20 ± 0.22                 | 0.86 ± 0.17    | 0.16 ± 0.06                        |
| Discomfort   | 1.92 ± 0.29 | 1.25 ± 0.11 | 2.71 ± 0.31                 | 0.79 ± 0.24    | 0.16 ± 0.06                        |
| Fatigue      | 3.05 ± 0.34 | 2.04 ± 0.17 | 3.19 ± 0.34                 | 0.13 ± 0.21    | 0.09 ± 0.10                        |
| Relaxation   | 7.44 ± 0.28 | 8.16 ± 0.15 | 6.65 ± 0.29                 | –0.79 ± 0.24   | –0.22 ± 0.08                       |

All data were analyzed using corresponding t-tests with very similar results.

<sup>a</sup>Difference score mean ± SE. TETRA – sham; positive value indicates TETRA > sham. <sup>b</sup>Wilcoxon signed rank test. <sup>c</sup>Mann-Whitney U-test. <sup>d</sup>Relaxation is reversed; therefore, a high score tends toward “extremely” relaxed. *p < 0.05, **p < 0.01, ***p < 0.008 (Bonferroni corrected α = 0.008).

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### Table 4. Descriptives (means ± SEs) and statistical tests (z-scores) for the double-blind VAS for sensitive and control participants.

| VAS measure  | Sham BL | TETRA BL | Difference score<sup>a</sup> | TETRA vs. sham | Sensitive<sup>b</sup> vs. Control<sup>c</sup> |
|--------------|---------|----------|--------------------------------|----------------|-----------------------------------|
| Anxiety      | 1.46 ± 0.24 | 0.97 ± 0.19 | 1.95 ± 0.24                  | 1.13 ± 0.08    | 0.03 ± 0.37                        |
| Tension      | 1.52 ± 0.25 | 0.96 ± 0.09 | 2.06 ± 0.26                  | 1.20 ± 0.09    | 0.03 ± 0.36                        |
| Arousal      | 1.41 ± 0.24 | 1.01 ± 0.10 | 2.12 ± 0.26                  | 1.16 ± 0.21    | –0.11 ± 0.37                       |
| Discomfort   | 1.42 ± 0.26 | 0.97 ± 0.09 | 2.07 ± 0.26                  | 1.08 ± 0.09    | –0.07 ± 0.32                       |
| Fatigue      | 2.17 ± 0.29 | 1.49 ± 0.14 | 2.70 ± 0.29                  | 1.60 ± 0.13    | –0.20 ± 0.45                       |
| Relaxation   | 7.70 ± 0.26 | 8.39 ± 0.12 | 6.00 ± 0.25                  | 6.74 ± 0.12    | 0.25 ± 0.48                        |

BL, baseline. All data were analyzed using corresponding t-tests with very similar results.

<sup>a</sup>Difference score mean ± SE for double-blind tests – (TETRA – TETRA baseline) – (sham – sham baseline); positive value indicates TETRA > sham. <sup>b</sup>Double-blind within-group analyses were performed on the difference between exposure and baseline scores using Wilcoxon sign rank tests (i.e., TETRA – baseline vs. sham – baseline). <sup>c</sup>Double-blind between-group analyses were performed on the difference of the difference scores using Mann-Whitney U-tests; that is, sensitive (TETRA – baseline) – (sham – baseline) vs. control (TETRA – baseline) – (sham – baseline). *Relaxation is reversed; therefore, a high score tends toward “extremely” relaxed.

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### Table 5. Symptom scales descriptives (means ± SEs) and statistical tests (z-scores) for the open provocation and double-blind tests for sensitive and control participants.

| Test                  | Sham       | TETRA      | Difference score<sup>a</sup> | TETRA vs. sham | Sensitive<sup>b</sup> vs. Control<sup>c</sup> |
|-----------------------|------------|------------|-----------------------------|----------------|-----------------------------------|
| Open provocation      |            |            |                             |                |                                   |
| Total symptom score   | 3.94 ± 1.00 | 1.13 ± 0.19 | 7.90 ± 1.75                 | 3.98 ± 1.44    | –0.17 ± 0.18                       |
| Total number of symptoms reported | 3.55 ± 0.80 | 1.03 ± 0.16 | 5.71 ± 1.08                 | 2.16 ± 0.74    | –0.23 ± 0.14                       |
| Double-blind          |            |            |                             |                |                                   |
| Total symptom score   | 6.63 ± 1.65 | 1.25 ± 0.22 | 3.84 ± 0.88                 | 1.15 ± 0.19    | –2.78 ± 1.56                       |
| Total number of symptoms reported | 5.06 ± 1.04 | 0.91 ± 0.13 | 2.99 ± 0.67                 | 0.92 ± 0.12    | –2.07 ± 1.03                       |

<sup>a</sup>Difference score: TETRA – sham; positive value indicates TETRA > sham. *Wilcoxon signed rank tests. *Mann-Whitney U-tests. *p < 0.001.

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Table 6. Descriptive (means ± SEs for original untransformed data) and statistical tests (F-scores) for physiologic measures for sensitive and control participants by exposure during open provocation and double-blind tests.

| Test/measure | Sham | TETRA | Condition | Sensitive vs. control | Group × condition |
|--------------|------|-------|-----------|-----------------------|-------------------|
| **Open provocation** |      |       |           |                       |                   |
| BVP mean *a* | 36.02 ± 0.03 | 36.10 ± 0.06 | 36.01 ± 0.03 | 36.13 ± 0.10 | NA | NA | NA |
| BVP SD *a* | 1.15 ± 0.13 | 1.12 ± 0.09 | 1.10 ± 0.13 | 1.14 ± 0.11 | NA | NA | NA |
| SC mean | 6.99 ± 0.35 | 7.61 ± 0.44 | 6.99 ± 0.35 | 7.66 ± 0.45 | 0.24 | 0.06 | 0.45 |
| SC SD | 0.93 ± 0.11 | 0.95 ± 0.07 | 0.94 ± 0.12 | 0.99 ± 0.08 | 0.08 | 0.12 | 0.08 |
| HR mean | 71.78 ± 1.50 | 67.58 ± 0.88 | 71.66 ± 1.46 | 66.94 ± 0.88 | 2.11 | 7.16** | 1.02 |
| HR SD a | 7.46 ± 0.73 | 7.11 ± 0.59 | 7.00 ± 0.78 | 6.84 ± 0.54 | NA | NA | NA |
| **Double-blind** |      |       |           |                       |                   |
| BVP mean *a* | 36.30 ± 0.14 | 36.71 ± 0.49 | 36.23 ± 0.08 | 36.16 ± 0.06 | NA | NA | NA |
| BVP SD | 1.19 ± 0.09 | 1.58 ± 0.35 | 1.17 ± 0.12 | 1.27 ± 0.08 | 0.18 | 0.15 | 0.20 |
| SC mean | 9.50 ± 0.88 | 8.67 ± 0.47 | 8.84 ± 0.74 | 7.98 ± 0.40 | 1.03 | 0.92 | 0.42 |
| SC SD | 1.63 ± 0.13 | 1.68 ± 0.09 | 1.62 ± 0.11 | 1.49 ± 0.09 | 0.78 | 0.45 | 3.34 |
| HR mean | 77.01 ± 1.32 | 73.51 ± 1.08 | 77.07 ± 1.29 | 72.69 ± 0.91 | 0.25 | 5.59* | 0.33 |
| HR SD | 9.40 ± 0.79 | 8.83 ± 0.59 | 10.63 ± 1.16 | 8.23 ± 0.47 | 0.04 | 3.17 | 1.11 |

NA, not applicable. Nonparametric statistics were also performed on the untransformed data with virtually the same results (copies of this analysis are available upon request).

*a*These data did not lend themselves to transformation because participants’ scores were tightly grouped around the mean; therefore, ANOVAs were not conducted on these data. *p < 0.05. **p < 0.01.

Table 7. Successful and failed judgments by trial for sensitive and control participants [n (%)].

| Trial | Sensitive [n = 50] | Control [n = 132] | Sensitive vs. control |
|-------|-------------------|------------------|----------------------|
|       | Correct | Incorrect | χ² | Correct | Incorrect | χ² | Correct | Incorrect | χ² |
| **Trial 1** | 26 (52.0) | 24 (48.0) | 0 | 66 (50.0) | 66 (50.0) | 0 | 0.01 |
| **Trial 2** | 29 (56.0) | 22 (44.0) | 0.33 | 61 (46.2) | 71 (53.8) | 0.54 | 1.03 |
| **Trial 3** | 31 (62.0) | 19 (38.0) | 2.17 | 55 (41.7) | 77 (58.3) | 3.82* | 5.23* |
| **Trial 4** | 25 (50.0) | 25 (50.0) | 0 | 55 (41.7) | 77 (58.3) | 2.96 | 0.71 |
| **Trial 5** | 22 (45.8) | 26 (54.2) | 0.08 | 72 (64.5) | 60 (45.5) | 0.97 | 0.75 |
| **Trial 6** | 16 (32.7) | 32 (66.7) | 4.2* | 69 (52.3) | 63 (47.7) | 0.1 | 4.34* |

Values reflect χ² with Yates continuity corrections. *p*-Values with or without the correction are very similar.

*a*Duration of exposure, 5 min. *b*Duration of exposure, 50 min. *c*Sensitivity group: n = 48 for sessions 2 and 3. *p < 0.05.

Figure 3. The probability of getting 0–6 of 6 correct: shown as a percentage by group of expected outcome versus observed outcome. (A) Controls [n = 132]. (B) Sensitives [n = 48].

This investigation is the first to examine the short-term effects of a TETRA base station signal on human health and well-being. We found no evidence to suggest that TETRA base station signals have a negative impact on health and well-being in either the control or sensitive groups. In addition, neither group could reliably detect the presence of a TETRA signal. These findings concur with two recent studies conducted on TETRA handset signals (Barker et al. 2007; Green et al. 2005) as well as a growing body of evidence on GSM and UMTS signals, which found no differences between active and sham conditions and no ability to detect the presence of EMFs (Elitti et al. 2007; Kwon et al. 2008; Lykov et al. 2001b; Regel et al. 2006; Rubin et al. 2006; Zwamborn et al. 2003). Moreover, Rubin et al. (2010) concluded in their systematic review that the evidence linking IEM-EMF to EMFs is weaker now than in their first review (Rubin et al. 2005). Taken together, these findings suggest that RF-EMFs in the range of approximately 400–2100 MHz do not have a negative impact on human health.

This study has some limitations. Our study tested individuals who identified themselves as sensitive to EMFs. Unlike a handset, which one can know to be on or off, reporting a sensitivity specifically to the TETRA base station signal is difficult, because it is unlikely that one can know where the signal is...
coming from. Therefore, this study addresses the general question of whether people with a perceived sensitivity to EMFs are affected by TETRA. We did not ascertain the exact onset of symptoms for each sensitive participant, and our study does not address long-term consequences of EMF exposure. We were not able to recruit the target number of sensitive participants in the 2.5-year period of funding for the study. Even with these caveats, we had sufficient power to detect a reliable difference in the open provocation condition. When participants knew they were being exposed to a TETRA signal, we found changes in measures, but these results were not replicated under double-blind conditions. Sensitive participants continued to report more symptoms and experienced a greater overall severity of symptoms than did controls, but this effect was independent of exposure condition. This pattern of results replicates our previous findings using GSM and UMTS signals, which showed that only during open provocation conditions did sensitive participants report increased symptom and reduced well-being under active signal conditions (Elitti et al. 2007a). Overall, these results suggest that it is not acute exposure to a base station signal that causes symptoms, but the knowledge of that exposure.

The long-term health consequences of EMF exposure are most often associated with physical health risks such as cancer. However, the long-term consequences of prolonged stress and anxiety that accompany the expectations associated with the everyday experiences of IEI-EMF individuals may be equally serious and, to our knowledge, have not been studied. For example, evidence suggests that ongoing worry is associated with low HR variability and increased HR during waking and sleeping, which, in turn, is associated with an increased risk of cardiovascular disease (Brosschot et al. 2007). Such findings indicate the importance of cognitive factors in IEI-EMF. Therefore, we recommend that future research should include a case-control study to elucidate the long-term impact of IEI-EMF on health.

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