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## Challenges and recommendations for magnetic hyperthermia characterization measurements

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### ABSTRACT

**Purpose:** The localized heating of magnetic nanoparticles (MNPs) *via* the application of time-varying magnetic fields – a process known as magnetic field hyperthermia (MFH) – can greatly enhance existing options for cancer treatment; but for broad clinical uptake its optimization, reproducibility and safety must be comprehensively proven. As part of this effort, the quantification of MNP heating – characterized by the specific loss power (SLP), measured in W/g, or by the intrinsic loss power (ILP), in  $\text{Hm}^2/\text{kg}$  – is frequently reported. However, in SLP/ILP measurements to date, the apparatus, the analysis techniques and the field conditions used by different researchers have varied greatly, leading to questions as to the reproducibility of the measurements.

**Materials and Methods:** An interlaboratory study (across  $N = 21$  European sites) of calorimetry measurements that constitutes a snapshot of the current state-of-the-art within the MFH community has been undertaken. Identical samples of two stable nanoparticle systems were distributed to all participating laboratories. Raw measurement data as well as the results of in-house analysis techniques were collected along with details of the measurement apparatus used. Raw measurement data was further reanalyzed by universal application of the corrected-slope method to examine relative influences of apparatus and results processing.

**Results:** The data show that although there is very good intralaboratory repeatability, the overall interlaboratory measurement accuracy is poor, with the consolidated ILP data having standard deviations on the mean of ca.  $\pm 30\%$  to  $\pm 40\%$ . There is a strong systematic component to the uncertainties, and a clear rank correlation between the measuring laboratory and the ILP. Both of these are indications of a current lack of normalization in this field. A number of possible sources of systematic uncertainties are identified, and means determined to alleviate or minimize them. However, no single dominant factor was identified, and significant work remains to ascertain and remove the remaining uncertainty sources.

**Conclusion:** We conclude that the study reveals a current lack of harmonization in MFH characterization of MNPs, and highlights the growing need for standardized, quantitative characterization techniques for this emerging medical technology.

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## 1. Introduction

Cancer remains a leading public health challenge facing humanity in the twenty first century. In 2018 there were 17 million new cases worldwide, with an anticipated increase to 27.5 million by 2040 [1]. The most established methods of cancer treatment at present are surgery, radiotherapy and chemotherapy. These techniques have shown significant progress in recent decades, and are complemented today by other more recently developed techniques such as

immunotherapy [2] or hormonotherapy [3]. Despite the progress made, there remains a significant need for innovative approaches which improve patient outcomes, while minimizing the trauma and collateral damage associated with established cancer therapies.

Magnetic field hyperthermia (MFH), also referred to as magnetic fluid hyperthermia, is an emerging technique capable of complementing or replacing established cancer therapies [4,5]. MFH requires magnetic nanoparticles (MNPs)

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 Supplemental data for this article can be accessed [here](#).

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to be introduced to the tumor tissue, which are then activated by the application of a radio frequency time-varying magnetic field. The MNPs dissipate heat, elevating the temperature of the cancer cells to induce weakness or death, or to render them more sensitive to chemo- or radiotherapy [6]. The technique has already shown great promise in human [4,7] and animal trials. With appropriate refinement, MFH is anticipated to offer new capabilities in cancer therapy, while inflicting minimal strain on the patient's physiology [8].

When evaluating MNP heating efficiency for hyperthermia, researchers typically make calorimetry measurements, and report the heating power  $P$  dissipated per unit mass of MNPs,  $m_{\text{MNP}}$ . The efficiency of this process is reported as the specific loss power (SLP):

$$SLP = P/m_{\text{MNP}}, \quad (1)$$

measured in Watts per gram [9]. The properties of the externally-applied time-varying magnetic field dictate the extent of heating produced by the MNPs, and thus Kallumadil et al. [10] proposed the intrinsic loss power (ILP) as an approximation suitable for comparing the outcomes of heating efficiency measurements conducted at different magnetic field frequencies or amplitudes. It is given by:

$$ILP = SLP/H_0^2 f, \quad (2)$$

where  $f$  is the frequency and  $H_0$  the magnitude of the time-varying field,  $H(t) = H_0 \sin(2\pi f t)$ .

Equation (2) is strictly only valid for low field amplitudes and frequencies as dictated in part by the linear response theory (LRT) regime for superparamagnetic nanoparticles [11,12]. One way to describe the LRT condition is to note that the power dissipation in Equation (1) is generated by the cyclic response of the magnetization of the material,  $M(t)$ , in response to the time-varying magnetic field:

$$P = \mu_0 f \oint M(t) dH(t), \quad (3)$$

where  $\mu_0$  is the permeability of free space. The LRT condition then applies when the magnetization is linearly proportional to the magnetic field.

Carrey et al. [12] have shown that this corresponds to the condition that the parameter:

$$\xi = \mu_0 M_s V H_0 / k_B T \text{ should be } \leq 1, \quad (4)$$

where  $M_s$  is the saturation magnetization of a nanoparticle of volume  $V$  at temperature  $T$ , and  $k_B$  is the Boltzmann constant. Carrey et al. have also shown that the LRT approximation is generally suitable for materials with high magnetocrystalline anisotropies, where the anisotropy field  $H_K$  is much higher than  $H_0$ , but that it may also be applicable in less anisotropic materials if  $V$  is reduced [12]. For  $f$  in the range typically used in MFH, namely ca.  $10^5$ – $10^6$  Hz, it has been shown [11] that the LRT holds for polydisperse systems with a polydispersity index  $\geq 0.1$ . For typical iron-oxide based MNPs, the LRT region has been found experimentally to apply at the clinically relevant  $H_0$  amplitudes of a few kA/m and frequencies of several hundred kHz respectively [10].

Keen interest in MFH has resulted in a competitive research environment, with different laboratories vying to publish the latest attention-grabbing  $SLP$  or  $ILP$  values [13]. However, despite the large number of research groups and publications, there is as yet no consensus on a harmonized approach to conducting either the measurements or the data analysis used for determining the  $SLP$ . Furthermore, until now no interlaboratory comparison of MNP heating efficiency measurements has been published. Testing is required to examine whether the results produced by the different techniques vary, and to quantify the extent of the variation. Without these verification steps, the comparison of  $SLP$  measurements reported by different laboratories is of questionable significance. In the extreme, it is not possible to judge which are the most efficient particles, despite this being a vital issue for MFH development.

From both the standardization and product development perspectives then, reliable and accurate calorimetric characterization of MNPs is a key requirement for the successful technology transfer and clinical implementation of MFH [14]. Some efforts have been made in this direction, including a study of relevant factors and recommendations for best practice  $SLP$  measurements [15]; and a study and discussion of factors relevant for MFH standardization [16]. However, while these works illustrated how the results of in-house measurements on a specific calorimetry apparatus can differ with changing sample properties, measurement protocols and analysis techniques, the impact within the wider hyperthermia community has yet to be studied. It is therefore timely for an interlaboratory survey of the current technical capabilities in  $SLP$  characterization, to understand the cross-compatibility of the apparatus used, and to lay the foundation for future prenormative research and standardization. There is a similar need for validated  $SLP$  measurement methods. This is a complex topic, albeit one with established procedural guidelines to probe the robustness, precision and trueness of the measured value [17]. With MFH expanding ever further into preclinical and clinical trials, it is important to understand the current state of accuracy, and level of compatibility, between the different measurement methods in use today [18].

To this end, an interlaboratory comparison study was devised and conducted under the auspices of the RADIOMAG EU COST action TD 1402 [19]. A total of 21 laboratories contributed their measurements to the study, providing an unprecedented snapshot of the current state-of-the-art in measurement centers across the Europe. Here, we present the key findings and recommendations which resulted from this interlaboratory study.

## 2. Materials and methods

All of the participating laboratories used calorimetry-based methods and magnetic nanoparticle samples in liquid suspension. The specific apparatus, measurement and analysis techniques varied greatly between laboratories. The study was designed accordingly.

## 2.1. Study design

The goal of the study was to determine the reproducibility of *SLP* and/or *ILP* measurements acquired at different participating laboratories using a universal measurement protocol, and the same MNP preparations. Qualified reference materials for this purpose are not currently available. Guidelines state that materials other than certified reference materials may be employed as a provisional benchmark in order to test whether candidate reference materials and/or candidate test methods approach the required level of certainty [20]. A representative test material is defined as a material taken from a single batch, which offers sufficient homogeneity and stability regarding the specified properties, and which is implicitly assumed to be fit for use in development for target properties not yet validated.

To this end, RADIOMAG project members with expertise in MNP synthesis produced eight distinct batches of iron oxide nanoparticles, from which two were selected for distribution to the measurement laboratories. The nanoparticle systems selected differed in composition, magnetic properties and concentration, and exhibited distinct hyperthermia behaviors. The systems were characterized before distribution to the measurement partners, viz.: iron oxide concentration was estimated using a colorimetric assay after appropriate acidic digestion in HCl 5 mol L<sup>-1</sup> [21], and the hydrodynamic diameters of the suspensions were measured by a backscattering DLS method (at 135° angle), not necessitating any dilution [22].

The study comprised two rounds. The measurement protocol in Round 1 represented the consortium's best initial effort at establishing suitable sample preparation and data collection methods. After assessing these results, in Round 2 the sample preparation and measurement details were refined, and additional system characterization tests were included.

## 2.2. Measurement apparatus

To find the experimental conditions which best matched those available across the participating laboratories, a questionnaire was circulated to all partners. The results provided an interesting snapshot of experimental capabilities in the field. The best match of measurement parameters accessible to almost all the participants was as summarized in Table 1.

Separately, a detailed survey of the apparatus used was conducted. This revealed a large variation in equipment types, with: nearly adiabatic *versus* non-adiabatic systems; commercial *versus* custom build magnetic field generators; thermocouple *versus* fiber optic probe thermometers; copper *versus* Litz wire coils; and various coil designs with different numbers of turns and sample volumes, etc.

## 2.3. MNP samples

The choice of samples for the study was driven in part by the goal of applying correlation and repeatability analysis methods to the collected data (see Section 2.6 below). In particular,

**Table 1.** Field frequency, intensity and sample volume parameters chosen as the best match for the different capabilities of the measurement laboratories.

Frequency	Field intensity	Sample volume
300 kHz	15 kAm <sup>-1</sup> (Peak) – 10.6 kAm <sup>-1</sup> (RMS)	1 ml

**Table 2.** Properties of the liquid suspensions distributed for measurement:  $c_{\text{FeOx}}$  = iron oxide concentration;  $\sigma_s$  = saturation magnetization per unit mass of iron oxide, measured at 300 K;  $M_s$  = saturation magnetization per unit volume of iron oxide, measured at 300 K;  $d_{\text{core}}$  = average magnetic core diameter;  $d_{\text{Hydro}}$  = intensity (z) average hydrodynamic diameter;  $PDI$  = polydispersity index; the latter two parameters determined from second order cumulant fitting of the DLS correlogram.

Sample ID	$c_{\text{FeOx}}$ [mg <sub>FeOx</sub> /mL]	$\sigma_s$ [Am <sup>2</sup> /kg <sub>FeOx</sub> ]	$M_s$ [kA/m]	$d_{\text{core}}$ [nm]	$d_{\text{Hydro}}$ [nm]	$PDI$
Sample 1	40.6	69.3	356	7.0	170	0.21
Sample 2	3.9	94.4	485	8.3	90	0.24

the ASTM E691-18 guidelines on “Standard Practice for Conducting an Interlaboratory Study to Determine the Precision of a Test Method” [23] state that analysis should be undertaken on a single “test result”, this being defined as “the value of a characteristic obtained by carrying out a specified test method”. Given that our intention was to determine the precision of the magnetic hyperthermia test method, even though the participating laboratories in the study had similar, but not identical apparatus (and therefore could not be expected to employ the same field intensity and frequency), it was apparent that the “test result” parameter would need to be the *ILP* rather than the *SLP*. This therefore meant that a key requirement for the choice of samples was that they should exhibit behavior consistent with the LRT conditions described in Equations (3) and (4) above.

Given the already-mentioned observation that many iron-oxide based MNPs have been reported as exhibiting LRT behavior in response to fields and frequencies comparable to those listed in Table 1 [10], it was therefore natural to start the search for suitable samples from that material type. After some preliminary testing, two promising MNP samples (Samples 1 and 2) were selected as representative of those typically studied in hyperthermia characterization, and as likely LRT candidates. Both were aqueous suspensions of multicore magnetic nanoparticles with organic coatings, with the magnetic cores being composed of the iron oxides magnetite (Fe<sub>3</sub>O<sub>4</sub>) and/or maghemite (γ-Fe<sub>2</sub>O<sub>3</sub>). A summary of their primary characteristics is shown in Table 2. The measured saturation magnetizations at 300 K (356 ± 22 kA m<sup>-1</sup> and 485 ± 30 kA m<sup>-1</sup> respectively) were consistent with Sample 1 being primarily maghemite, and Sample 2 being primarily magnetite [24]. Both samples were measured in each round of the study, but fresh aliquots were delivered in Round 2 to avoid the possible confounding effects of interlaboratory differences in storage conditions.

## 2.4. Measurement protocol

For each round, a measurement protocol and a results table were distributed to each partner, along with the samples.

The protocols were detailed, but allowed for some flexibility in recognition of the different apparatus in each laboratory.

### 2.4.1. Round 1 protocol

**2.4.1.1. Sample preparation.** *Inspection* – Prior to measurement, samples were to be visually inspected and any signs of sedimentation or precipitation noted. Subsequent sonication or vortexing for a period of not more than 5 min was permitted, but was not mandatory, being instead a matter of local practice. (If such a step was included, an additional 30 min was to be added to the thermalization time.) *Thermalization* – Samples were to be equilibrated at room temperature for at least 2 h before measurements commenced. *Filling* – A standard empty vial (chromatographic 1.8 ml) was supplied along with the sample material, with a request to transfer 1.0 ml of the undiluted sample fluid into the vial using a pipette or micropipette. If the supplied vial did not fit the local apparatus, the measurer was advised to use their normal sample receptacle, and use a volume as close to 1 ml as possible.

**2.4.1.2. Apparatus positioning.** *Sample Positioning* – No guidance was provided for placement of the sample within the excitation coils. This allowed the measurers to use their normal procedures and provided an accurate representation of their in-house measurement technique. *Temperature Probe Positioning* – The temperature probe was to be centered as much as possible within the solution. If the sample volume was too small, then it was understood that the available positions might be limited. In any case, the measurer was asked to aim for the temperature probe to be totally immersed within the sample, while not touching the sides of the container.

**2.4.1.3. Measurements.** *Adiabatic and Non-adiabatic Apparatus* – Both were used in the study, although most laboratories used non-adiabatic systems. (The less common adiabatic calorimeters typically include an “active” vacuum insulation jacket or similar means to minimize or control heat transfer between the sample and its environment.) For the adiabatic systems, the measurement protocol was slightly changed by omitting the cooling curve part of the measurement (see below). *Time Resolution* – Sample temperatures were to be recorded once per second if the measurement equipment permitted. If the measurement equipment was not compatible with this requirement, the measurer was asked to use the nearest possible setting. *Initial Temperature Recording* – Measurers were asked to record the sample temperature for 200 s prior to switching on the applied time-varying field. *Field Exposure Time* – The field was to be switched on for 300 s. The frequency and amplitude were requested to be as close to 300 kHz and  $15 \text{ kAm}^{-1}$  as possible. In those cases where the apparatus could not match the requested values, the measurer was asked to report the actual values used. *Cooling curve* – After the field was switched off, the temperature was to continue to be recorded as the sample cooled. The measurement was to be

terminated only once the sample temperature had returned to that observed before the field was switched on, and this baseline state had been maintained for 30 s. *Repeats* – The measurement procedure was to be repeated three times for each sample. For each repetition, the same values for all measurement parameters (field intensity, frequency, sample volume, exposure time, etc.) were to be used. *Ambient Temperature* – The measurers were asked to track and report the ambient temperature in their laboratory over the measurement period.

**2.4.1.4. Analysis.** In addition to providing the raw measurement data in the measurement report sheet, each measurer was also requested to calculate the SLP from the measurement data using their own in-house technique. For this calculation, the iron concentration values for each sample were provided in the report sheet, and the measurers were asked to report the SLP in watts per gram of iron.

### 2.4.2. Round 2 protocol

The measurement protocol set was refined before Round 2, based on the lessons learnt from analyzing the results of Round 1. A summary of the changes made is as follows.

*Temperature Limits* – Measurers were requested to switch off the field early if the sample temperature reached  $60^\circ\text{C}$ . This was to avoid the possible confounding effects of water evaporation from the sample. *Probe Positioning* – Further guidance on optimal positioning of the temperature probe(s) was included. *Refinement of Instructions* – General improvements of the instructions section of the protocol to improve the accessibility and ease of understanding, including an explanation of the concepts of “adiabatic” and “non-adiabatic” systems, and a diagram illustrating the entire measurement protocol from start to finish, as well as a simplified data-reporting spreadsheet. *Water Measurement* – To help assess the environmental losses in the non-adiabatic systems, a pure water sample was circulated for measurement in the same manner as the MNP samples, to assess environmental heat transfer both into and out of the sample space during measurement.

## 2.5. SLP calculations

The raw data collected from each laboratory comprised three successive heating and (for non-adiabatic systems) cooling curves for each sample, all collected under the same conditions. To preserve anonymity during data processing, each laboratory was randomly assigned a unique identifier consisting of two letters and two numbers. These identifiers were used to label the results.

Various approaches exist for calculating the SLP from heating/cooling curves, and each of the measurers had their own method for calculating it. Therefore, in each round, the participants were asked to analyze their data using their in-house methods, and also to provide the raw data for single-operator recalculations based on the Corrected Slope Method [15] (CSM), a method that was used to analyze and

compensate for the environmental heat losses in the non-adiabatic systems.

Briefly, in the CSM the thermal loss behavior of the apparatus is determined by measuring the cooling behavior of a hot sample in the absence of a magnetic field excitation. Plotting the numerical derivative of the cooling curve reveals an apparatus-specific  $\Delta T_{LLR}$  above the baseline temperature  $T_0$  over which the environmental losses are directly proportional to the measured  $\Delta T = T(t) - T_0$ . Within this 'linear loss region' the power  $P$  in Equation (1) is replaced by  $P - L \Delta T$ , where the 'linear loss factor'  $L$  is a sample-specific constant. By restricting data analysis to the  $\Delta T_{LLR}$  region, the  $L$  parameter could be determined directly from the heating curves, alongside the  $SLP$  [25].

## 2.6. Correlation and repeatability analysis

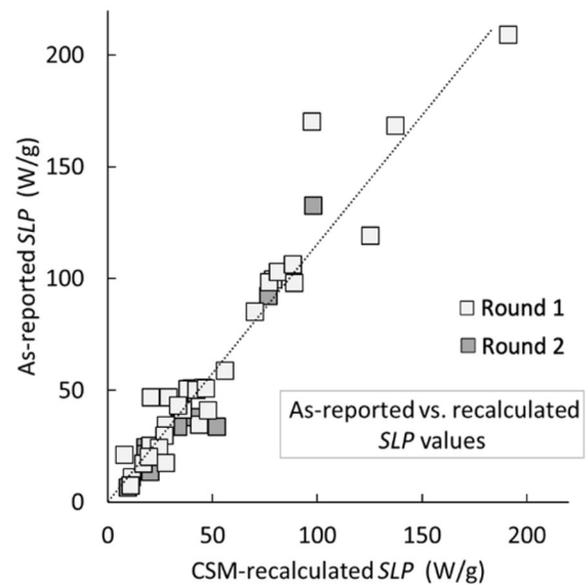
For correlation analysis of the data, both the Pearson and Spearman statistical methods were used. In both cases they were calculated using readily available spreadsheet functions. The Pearson correlation coefficient  $r_{xy}$  is a measure of the linear correlation between two variables  $x$  and  $y$ , with  $r_{xy} = +1$  or  $-1$  denoting a total positive or negative linear correlation, and  $r_{xy} = 0$  indicating no linear correlation at all. The Spearman rank correlation coefficient  $\rho$  is equal to the Pearson coefficient applied to the rank values of the two variables, rather than the variables themselves. It is the non-parametric version of the Pearson correlation and is used to assess the degree to which two variables are monotonically related.

For repeatability analysis of the data, Youden plots [26] and Mandel  $h$  statistics [23,27] were used. Youden plots are a means of comparing precision and bias between laboratories, and distinguishing between random and systematic uncertainties, by graphical means. The Mandel  $h$  statistic is given by  $h = (\alpha - \alpha_m)/s_x$ , where  $\alpha_m$  is the mean value for each sample, and  $s_x$  is the standard deviation of all the measurements of the laboratories. In this way,  $h$  is a measure of the deviation of a single laboratory's result from the overall mean, and can be used to identify outliers, and test significance. For the latter, a critical value of the statistic,  $h_{crit}$ , may be defined for any given significance level and number of independent measurements [23], which can further assist the comparative process.

## 3. Results

### 3.1. Initial review and analysis

Completed measurement reports and temperature-time data suitable for detailed analysis were received from 17 of the 21 responding laboratories during Round 1 of the interlaboratory study, and from 8 of 12 respondents during Round 2. Many of the laboratories were unable to exactly match the requested field parameters ( $H_0 = 15 \text{ kA m}^{-1}$ ,  $f = 300 \text{ kHz}$ ), but in 16/17 of the Round 1 cases  $H_0$  and  $f$  ranged from 6.2 to  $15.0 \text{ kA m}^{-1}$  and 194 to 377 kHz respectively. The exception

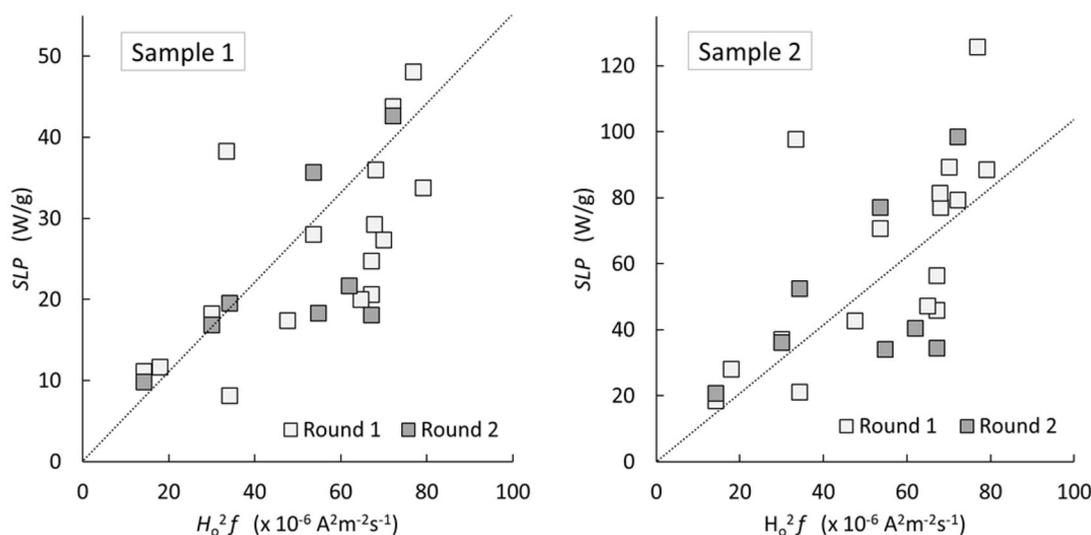


**Figure 1.** As-reported  $SLP$  values for Samples 1 and 2 plotted against CSM-recalculated values. The CSM analysis takes account of the inherent environmental losses in non-adiabatic calorimetry systems and is confined to the 'linear loss region' of the  $T(t)$  heating curves. The dotted line is a best linear fit to the data; it has a slope of ca. 1.15, indicating a tendency for the as-reported values to be higher than the recalculated ones.

was one laboratory that used  $H_0 = 15 \text{ kA m}^{-1}$  and  $f = 928 \text{ kHz}$ .

Initial inspection of the heating curves showed that in almost all cases, the three consecutive datasets recorded on the same sample were in excellent agreement with one another. Subsequent single-operator CSM reanalysis confirmed this agreement, with the intra-laboratory standard deviation in the three consecutive measurements generally being of order  $\pm 2\%$  to  $\pm 6\%$ . That said, the analysis revealed some exceptions, with run-to-run deviations of between  $\pm 10\%$  and  $\pm 25\%$  in 8/50 cases, and even higher deviations in 3/50 cases. It was further noted that in the 3/50 cases where the run-to-run deviations exceeded  $\pm 25\%$ , there was a consistent increase in the reported  $SLP$  values from run 1 to run 2 to run 3; while in the 8/50 cases with  $\pm 10\text{--}25\%$  deviations, no such trend was apparent.

As-reported  $SLP$  values were then compared to the CSM-recalculated values, as shown in Figure 1. The data show that there is a correlation between the two, with a mean ratio of  $1.15 \pm 0.36$ , indicating a tendency for the as-reported values to be higher than the recalculated ones. Similar deviations have been previously reported from the re-analysis of literature data [15]. For the CSM analysis, the cooling curves from each of the laboratories were analyzed to identify the 'linear loss region' for heat transfer to the environment. This  $\Delta T_{LLR}$  was found to be surprisingly consistent across all the laboratories, and was of order 25 K. The 'linear loss factor'  $L$  determined from the CSM fits was more variable, with the fitted values ranging from ca.  $10\text{--}50 \text{ mW K}^{-1}$  for Sample 1, with a mean of  $24 \pm 12 \text{ mW K}^{-1}$ ; and from ca. 6 to  $26 \text{ mW K}^{-1}$  for Sample 2, with a mean of  $12 \pm 6 \text{ mW K}^{-1}$  – values that are comparable with previous reports on similar apparatus of  $L$  values of order  $5\text{--}10 \text{ mW K}^{-1}$  [15].



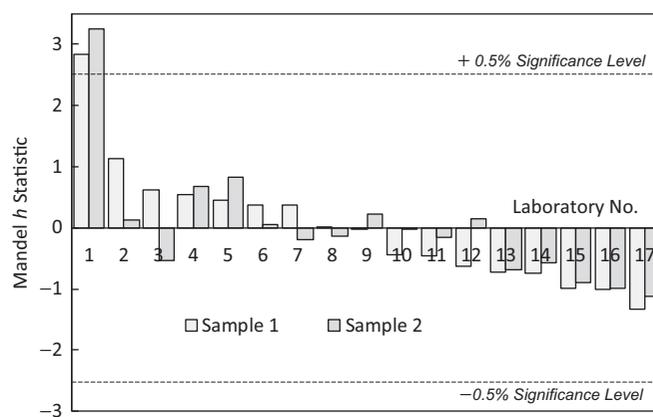
**Figure 2.** *SLP* values for Samples 1 and 2 plotted as a function of  $H_o^2 f$ , for which at least a monotonic trend is expected, according to Equations (1) and (3). The scatter evident in the data points is an indication of interlaboratory measurement variability. Dotted lines are guides to the eye only.

Further analysis proceeded using the CSM-recalculated fits. Figure 2 shows the measured *SLP* values for Samples 1 and 2, plotted as a function of  $H_o^2 f$ . According to Equations (1) and (3) it was anticipated that there should be a monotonic relationship between *SLP* and  $H_o^2 f$ . Although there is indeed a correlation between the variables – the Spearman coefficients are 0.75 and 0.80 for Samples 1 and 2 respectively – it is nonetheless clear from the data that there is significant scatter and variation in the measured data points. Furthermore, comparison in Figure 2 of the data collected in Rounds 1 and 2 of the study shows that despite significant extra control having been taken of the Round 2 measurements, there is even then no clear sign of monotonic behavior in the data. This therefore points to a possible systematic uncertainty in the measurements as undertaken in the different laboratories.

### 3.2. Repeatability analysis

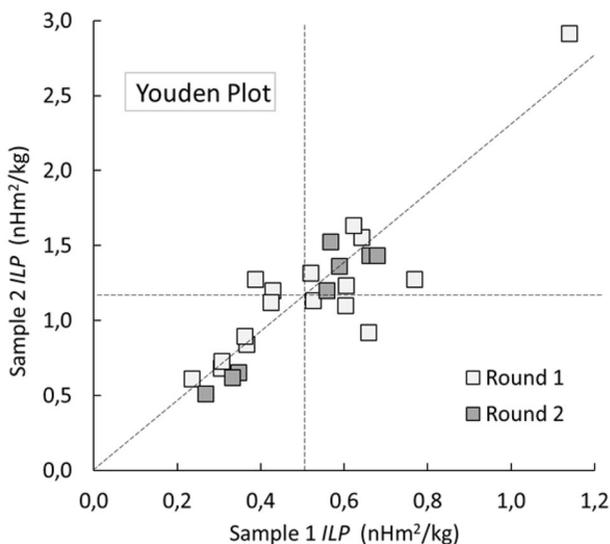
Before undertaking repeatability analysis of the data, a decision needed to be taken as to whether the *ILP* metric of Equation (2) could be used for such purposes. This step was required given that the ASTM E691-18 International Standard [23] states that repeatability analysis may be undertaken only on a single measurable metric, such as the *ILP*, rather than on a range of measurable metrics, such as the *SLP*.

First, it was noted from Table 2 that both Samples 1 and 2 were polydisperse systems with PDI values well in excess of 0.1; and that the frequencies applied were all in the range from  $10^5$  to  $10^6$  Hz. According to this criterion [11], both would therefore be expected to lie within the LRT regime. Second, the  $\zeta$  parameter of Equation (4) was calculated for both samples using the measured  $M_5$  values from Table 2. For  $H_o = 15 \text{ kA m}^{-1}$  the  $\zeta$  parameter was determined to be ca. 0.29 for Sample 1 and ca. 0.66 for Sample 2, both of which are less than one, and thus satisfy the criterion to lie within the LRT regime [12]. Third, the data from Table 2 was used to locate the samples on the LRT “validity map”



**Figure 3.** Mandel *h* statistics (relative to the mean) for Round 1 measurements, using the CSM-recalculated *ILP* values. Laboratory 1 is an outlier, with *h* values that exceed the  $\pm 0.5\%$  significance level,  $h_{crit} = 2.51$ .

introduced by Carrey et al. [12], which allowed for consideration of both particle size and anisotropy, alongside the external conditions ( $H_o$  and  $T$ ). According to this graphical analysis, the parameters for both samples lay well within the LRT validity region (see Supplementary Information, Figure S1). Fourth, despite the scatter, the data in Figure 2 are consistent with there being an underlying linear correlation between the *SLP* and  $H_o^2 f$ , as evidenced by Pearson coefficients of 0.91 and 0.85 for Samples 1 and 2 respectively. Fifth, a complementary version of Figure 2 was produced in which the *ILP* (rather than the *SLP*) was plotted against  $H_o^2 f$  (see Supplementary Information Figure S2). No correlation at all was evident between the *ILP* and  $H_o^2 f$  values, as is expected only in the LRT regime. The Pearson coefficients in this case were  $-0.12$  and  $-0.26$  for Samples 1 and 2 respectively. Taken together, these observations and considerations lead to the conclusion that at the fields and frequencies applied, the magnetizations of both samples responded linearly to the applied stimulus, and could therefore be characterized by the *ILP* parameter, as defined in Equation (2).

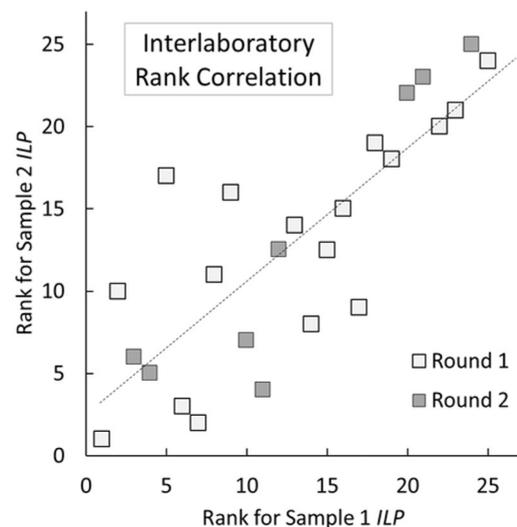


**Figure 4.** Youden plots of CSM-recalculated *ILP* values for both samples and for both rounds of measurement. The vertical and horizontal dashed lines mark the mean values for each sample,  $ILP_{1\text{-mean}}$  and  $ILP_{2\text{-mean}}$ . The diagonal line has slope  $ILP_{2\text{-mean}}/ILP_{1\text{-mean}} \simeq 2.25$ .

The measured *SLP* data was therefore converted into *ILP* data, which allowed for repeatability analyses to be undertaken using both the Mandel *h* statistic, as shown in Figure 3, and the Youden plot, as shown in Figure 4. For clarity, the Mandel *h* statistics in Figure 3 are shown for the Round 1 measurements only; the corresponding data for Round 2 are less comprehensive but very similar, and are given in the Supplementary Information, Figure S3. From inspection of both figures it is clear that in only one case did the *h* statistic exceed the  $\pm 0.5\%$  significance level ( $h_{\text{crit}} = 2.51$ , for  $p = 17$  independent measurements [23]), and that this occurred for both samples in that laboratory.

The Youden plot in Figure 4 includes the CSM-recalculated *ILP* data from both rounds of measurement. The outlier evident in the Mandel analysis is also seen here, as the data point in the top right-hand corner of the diagram, separated from the rest. However, also notable is the tendency for the data points to be spread out along the diagonal line with slope equal to the ratio of the mean *ILP* values of each sample. Such behavior is typical of Youden plots in which systematic uncertainties dominate over random errors.

Mean *ILP* values were then determined for each of the samples based on both the as-reported and CSM-recalculated data from Rounds 1 and 2 (see Table 1), but excluding the outlier data from Laboratory 1. On inspection of this data it is apparent that there is good agreement between the Round 1 and Round 2 data, and that the as-reported and CSM-recalculated values were not significantly different, despite the previously-mentioned tendency for the as-reported values to be overestimated, as seen in Figure 1. What is also apparent, however, is that the standard deviations on the means are large, at ca. 30% to 40% of the calculated mean. This is most likely a reflection of the systematic uncertainties seen in Figure 4, and is evidently a feature that was present in both rounds of measurement.



**Figure 5.** Interlaboratory rank correlation of the CSM-recalculated *ILP* values for both samples and both rounds of measurement. Each data point represents the ranking of a pair of measurements ( $ILP_1, ILP_2$ ) recorded in a single laboratory in a single round. The dotted line corresponds to the best linear fit to the data; its slope is equal to the Spearman correlation coefficient for the data,  $\rho = 0.81$ .

### 3.3. Correlation analysis

Having identified a systematic error contribution to the measured *ILP* values, a natural corollary was to explore whether there was a rank correlation in the data, wherein “some laboratories measured low, while others measured high”. To this end the CSM-recalculated *ILP* values for Samples 1 and 2, as measured in the different laboratories in both Rounds 1 and 2 ( $N = 25$  measurements for each sample) were ranked from highest to lowest *ILP*, and the results plotted in Figure 5. From this plot it appears that the strongest rank correlation is between those laboratories that recorded the smallest *ILP* values, i.e., those that were ranked ca. 18th or above. The overall Spearman correlation coefficient was determined to be  $\rho = 0.81$ , which indicates a moderately strong correlation.

### 3.4. Systematic factors analysis

In light of the large systematic variations revealed in the preceding analyses, the possible sources of such variations were considered and analyzed, as reported below.

#### 3.4.1. Sample degradation

Liquid suspensions of MNPs, as used in this study, may be liable to aggregation, sedimentation, and to a lesser extent chemical reaction. In addition, factors such as the exposure to magnetic fields, heat or sunlight during transit may influence such effects. It was therefore impossible to guarantee *a priori* that the samples all arrived at each laboratory in the same state as they were when shipped from the source. Participants were therefore asked to report any visually detectable instability in the as-received samples. In practice, no such effects were noted. (In some laboratories a vortexing step was included in the measurement protocol, but this was

**Table 3.** Means and standard deviations of the as-reported and CSM-recalculated *ILP* values for Samples 1 and 2, for both rounds of the study.

Intrinsic Loss Power, <i>ILP</i> [nHm <sup>2</sup> kg <sup>-1</sup> ]	As-reported <i>ILP</i> values				CSM-recalculated <i>ILP</i> values			
	Round 1		Round 2		Round 1		Round 2	
	Sample 1	Sample 2	Sample 1	Sample 2	Sample 1	Sample 2	Sample 1	Sample 2
Mean	0.52	1.13	0.49	1.07	0.49	1.09	0.50	1.09
Std. dev.	0.14	0.42	0.14	0.45	0.15	0.30	0.16	0.42

Data from the outlier, Laboratory 1, identified in Figure 3, was excluded from these calculations. Round 1 data was therefore from  $N = 16$  laboratories; Round 2 from  $N = 8$  laboratories.

a matter of local practice, rather than any perceived need to redisperse the samples.)

Sample aging, including environment-dependent aging [28], was another potentially confounding factor that was recognized at the outset. This was mitigated in the study protocol by asking the participants to store the samples at 5–8 °C upon receipt and to complete their measurements promptly thereafter, and by distributing fresh aliquots of centrally controlled samples in each round. In practice, all of the participants measured their samples within ca. one month of the receipt date, and there were no discernible round-to-round variations in the measured *ILP* values (Table 3).

### 3.4.2. Magnetic field amplitude

The study participants used a diverse range of (often bespoke) apparatus to generate the time-varying field,  $H(t) = H_0 \sin(2\pi f t)$ , for their calorimetry measurements, and their as-reported values for the frequency  $f$  and magnitude  $H_0$  were used in the subsequent *ILP* analysis. Although frequency can be measured to high precision, standard methods do not yet exist for the *in situ* measurement of  $H_0$ , since standard Hall probes do not function well in the  $f \approx 300$  kHz range, and specialist probes, although they do exist [29], are not yet in widespread use. Similarly, reliable magnetic heating calibration samples have not yet been developed. As such, most laboratories relied either on extrapolations from *dc* field measurements, or theoretical estimates based on the known geometry of the field generation coils in their apparatus; or, in the case of those users of commercial systems, on the manufacturer's reported  $H_0$  values. These are, however, problematic and generally unverified approaches, which may lead to unrecorded deviations between the reported  $H_0$  and the actual  $H_0$  at the sample. Furthermore, dependent on the coil geometry, field inhomogeneity over the sample volume may also be a source of unreported deviations.

To further analyze this possible source of systematic uncertainties, a *post facto* exercise was performed on the Round 1 data (excluding the outlier, Laboratory 1, so that  $N = 16$ ). It was assumed that the only source of interlaboratory deviation in the measured *ILPs* was the  $H_0$  value, and correction factors were applied – by replacing  $H_0$  with  $H_0' = (1 + x) H_0$  – to bring all the measurements in line with the overall mean. Naturally,  $x$  took both positive and negative values, hence attention was paid to its absolute value,  $|x|$ . It was found that across the 16 laboratories, the mean  $|x|$  was 14% for Sample 1, and 11% for Sample 2, with standard deviations of 8% in both cases. Perhaps more pertinently, a laboratory-specific repeatability metric  $y = |x_1 - x_2|$  was also

derived, where  $x_1$  and  $x_2$  were the  $x$  values for Samples 1 and 2 respectively. It was found that  $y$  was less than 2% in 4/16 cases, and more than 5% in 9/16 cases. The latter figure implies that it is unlikely that unrecorded deviations in  $H_0$  were a primary source of the systematic uncertainties, as if that were the case, the laboratory-specific  $y$  metrics derived here should have been close to zero in more cases.

### 3.4.3. Instrument manufacturer

Although many of the study participants used bespoke field generation apparatus, it was noted that two particular commercial systems had each been used by  $N = 5$  participants. It was found that for System 1, usable data was received from 4 partners, and that the measured mean *ILPs* for Samples 1 and 2 were ca. 13% and 16% higher, respectively, than the overall means. For System 2, usable data was received from 4 partners, but one of these was the outlier Laboratory 1; the measured mean *ILPs* for the remaining 3 laboratories, for Samples 1 and 2, were ca. 8% and 18% lower, respectively, than the overall means. Although the numbers of laboratories compared here is low, these are rather large deviations from the mean, which may indicate systematic variations between magnetic hyperthermia instrument manufacturers.

### 3.4.4. Thermometry

According to the measurement reports, 15 participants used a fiber-optic thermometer (from a wide range of manufacturers and models) in their apparatus; while 5 used a T-type copper/constantan thermocouple, all of which were supplied as standard with the commercial System 2 (see above). For the fiber-optic measurements, usable data was received from 13 partners: the measured mean *ILP* for Sample 1 was ca. 4% higher than the overall mean, while for Sample 2 it was approximately equal. For the thermocouple measurements, usable data was received from 4 partners, but one of these was the outlier Laboratory 1; the measured mean *ILPs* for the remaining 3 laboratories, for Samples 1 and 2, were ca. 8% and 18% lower, respectively, than the overall means.

Thermocouples can introduce at least two possible sources of uncertainty due to their electrical conductivity, viz.: (a) the occurrence of a spurious voltage jump upon field switching, which adds a sharp increase or decrease to the output temperature signal; and (b) continuous additional heating throughout the measurement window due to eddy currents within the thermocouple probes [30]. Both issues can be alleviated – the first through additional filtering, and the second by selecting thermocouples made from materials which

exhibit minimal conductivity – but to our knowledge the laboratories in question had not adopted such measures.

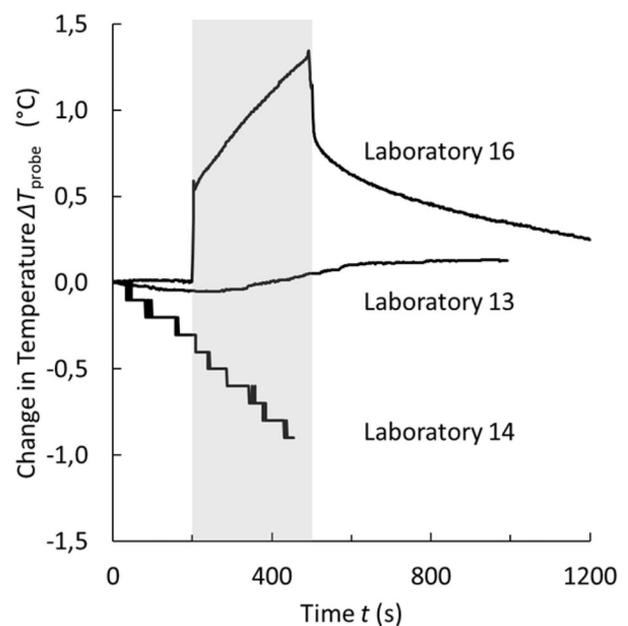
The mechanism by which such perturbative effects might lead to suppression in the *ILP* values measured *via* thermocouple probes is not yet well understood. One possibility is that the eddy-current-generated heat within the probe might inhibit its ability to respond to changes in the temperature of its environment, and that in effect, by having heat continually flowing out of the thermocouple, it might be harder to measure heat flow coming back into the thermocouple.

In addition to the probe types used, the precise manufacturer, model and associated read-out system employed may also influence the accuracy, noise-level and response time of temperature monitoring during calorimetry [31]. Furthermore, it is known that the probe position within a sample can impact on measurement [15,32]. In light of this, probe placement advice was provided in the Round 1 measurement protocol, and precise probe positioning was mandated in Round 2. However, systematic information on the extent to which these instructions were followed was not obtained, and therefore it is not clear whether probe placement factors may have contributed to the systematic uncertainties observed.

### 3.4.5. Sample environment

All but one of the laboratories used non-adiabatic calorimetry equipment for the *SLP* measurements. For the non-adiabatic systems, two major sources of heat flow have the potential to impact upon the shape of the heating curve. First, the heat loss behavior from the sample into the surrounding environment is different for every apparatus. It is decided by factors including the sample volume, and the quality of insulation employed. The heat loss behavior may have an impact on the final result, depending on the method used to calculate the *SLP*: e.g., the Corrected Slope Method is designed to analyze and compensate for the losses, while the initial slope technique does not. However, even the CSM calculations may not hold in cases where the calorimeter operates under low heat generation and/or poor thermal insulation conditions. Second, over time, waste heat from the field generation coil may penetrate the sample space, producing an additional heat flow into the sample, and an artificial enhancement of the apparent heating power.

The variability of heat loss in the non-adiabatic calorimeters was tested by inspection of the CSM-re-calculated data, and in particular the calculated 'linear loss factors' *L* (as described in Section 2.5). From the Round 1 data (excluding the outlier, Laboratory 1), the *L* parameter was found to range from 10 to 50 mW/K for Sample 1 and from 6 to 26 mW/K for Sample 2, with means of 24 and 13 mW/K respectively. These values are somewhat high compared to the 6 mW/K reported by Wildeboer et al. [15], albeit that measurement was on a sample with an *ILP* of ca. 2.8 nHm<sup>2</sup>kg<sup>-1</sup>, and some as-yet unexplored heat loss trends as a function of the *ILP* may be expected. In any case, correlation analysis of the *L* and *ILP* parameters gave Pearson



**Figure 6.** Illustration of the range of environmental heat transfer conditions under which different laboratory calorimeters operate, as measured using a centrally supplied pure water sample and the same measurement protocols as would be applied to a magnetic hyperthermia sample. The gray box highlights the period during which the magnetic field was applied. The middle curve is representative of the response recorded in most laboratories; the other two curves illustrate extreme responses.

coefficients of 0.02 and 0.00 for Samples 1 and 2 respectively, indicating that no correlation was present.

To explore possible heat-flow effects associated with the sample volume *V*, correlation analysis was performed on the Round 1 data, where the choice of sample volume had been left to user preference. Excluding the data from one laboratory which had used  $V = 40 \mu\text{L}$ , and focusing on the remaining laboratories where *V* ranged from 0.5 to 2.0 ml, the analysis gave the following Pearson coefficients for Samples 1 and 2 respectively: for *V* versus *L*,  $\rho = 0.04$  and  $-0.02$ ; and for *V* versus *ILP*,  $\rho = -0.16$  and  $0.25$ . All four  $\rho$  values were thus close to zero. This, coupled with the observation from Table 3 that the Round 2 *ILP* measurements, which were all performed on a fixed 1.0 ml volume sample, were very similar to the Round 1 measurements, leads to the conclusion that *V* was not a significant factor in the systematic uncertainties.

Lastly, with respect to possible environmental heat flows into the sample, it was noted that although most calorimeters are designed using cooling water circulation and thermal insulation to minimize or prevent this, no data was collected in Round 1 to evaluate this. A water sample was therefore supplied in Round 2 to provide a means of measuring such background effects in the users' equipment. Large variations in behavior were observed; Figure 6 illustrates this with temperature-time data as recorded in three different systems, including two extreme cases, and one typical case.

The data shown in Figure 6 from Laboratory 13 (the middle curve) was typical of most of the reported curves. This laboratory used a commercial calorimeter and a fiber-optic temperature probe. The data show that the temperature was broadly stable before the field is switched on, with a slight

fall probably caused by the cooling water circulating in the coil. When the field is applied, a slight increase in the sample temperature is recorded, which begins around halfway through the field-on period. The sample temperature does not rapidly reduce after the field is switched off, indicating that the probe has a temperature similar to the surrounding environment. This behavior is most likely due to heat produced in the coil penetrating the sample space over time. However, the overall temperature rise is minimal ( $\Delta T_{\max} \simeq 0.1$  K), and as such is unlikely to impact on the resulting *SLP*, provided that the measured samples produce a much higher temperature rise than this background effect.

Laboratory 16 (the upper curve in Figure 6) used another commercial system, but with an as-supplied T-type thermocouple for temperature measurement. When the field was switched on, an abrupt temperature increase  $\Delta T$  of ca. 0.5 K was recorded: this was most likely caused by an induced voltage in the thermocouple signal, rather than being a real change in the sample temperature [30]. Thereafter, the probe temperature increased steadily with time, most likely as the result of eddy currents flowing in the thermocouple, reaching a maximum  $\Delta T_{\max} \simeq 1.3$  K before the field was switched off. The temperature as recorded then fell abruptly by ca. 0.5 K, after which a typical cooling trend was observed, indicating that the probe was at a higher temperature than its environment. Taken together, this behavior is further evidence for extraneous perturbative effects on thermocouple measurements, as was discussed in Section 3.4.4. Solutions to these problems could be achieved by filtering electric signals and by choosing less conductive materials.

Laboratory 14 (the lower curve in Figure 6) employed a modified version of the same commercial apparatus as used in Laboratory 16, including the substitution of a fiber-optic probe for the as-supplied thermocouple, albeit the resolution of the fiber-optic was only 0.1 K, leading to 'steps' in the recorded temperature data. Before the field is switched on, a rather marked cooling effect was observed, presumably due to a refrigerating effect from the cooling water in the coil. This cooling trend continued throughout the measurement, with the field-on and field-off switching having no apparent effect. By the end of the measurement, a  $\Delta T_{\max}$  of ca.  $-0.8$  K had been recorded, which would likely be sufficient to confound *SLP* measurement on this apparatus.

However, from the received data it was clear that Laboratories 14 and 16 were exceptions rather than the rule, and as such, samples environment effects of this kind could not explain the observed systematic variations that were observed in Figure 4, nor the rank correlations seen in Figure 5.

#### 4. Discussion

Initial inspection of the completed measurement reports from the participating laboratories showed that there was generally a good level of experimental repeatability within any given laboratory, with run-to-run deviations of less than  $\pm 10\%$  in 39/50 cases, and less than  $\pm 5\%$  in 33/50 cases. In-house data analysis was also found to be generally good,

with single-operator re-analysis of the raw data using the Correlated Slope Method (CSM) showing a strong linear trend (Figure 1), albeit with a tendency toward over-estimation, consistent with previous CSM-recalculation reports [15].

Given the complexity of calorimetric measurements in general, a random uncertainty of order  $\pm 5\%$  to  $\pm 10\%$  should most probably be considered to be acceptable (so long as the values thus obtained are duly reported with the appropriate corresponding uncertainty). However, larger intra-laboratory run-to-run deviations were noted in 11/50 cases. The three largest deviations were found to correspond to incremental changes in the measurements from run to run, which was most likely a sign of instrument drift, and as such may be regarded as experimental error. In contrast, no such trend was found for the 8/50 cases of run-to-run deviations in the range from  $\pm 10\%$  to  $\pm 25\%$ , implying that these were cases where significant random experimental uncertainties were present.

In addition to these random measurement uncertainties, inspection of the CSM-recalculated Specific Loss Power (*SLP*) data plotted as a function of  $H_o^2 f$  in Figure 2 showed that there was significant scatter in the data – more so than would be expected from the intra-laboratory uncertainties alone – and that the expected linear (or at least monotonic) relationship between *SLP* and  $H_o^2 f$  was not clearly present. In light of this, careful consideration was given to whether the physical and magnetic properties of the samples studied, combined with the experimental conditions used, placed the data in the Linear Response Theory (LRT) regime as described by Equation (4). It was determined that the LRT criteria were indeed met, and that as such a linear correlation was to be expected between the *SLP* and  $H_o^2 f$  parameters, as embodied in Equation (2) as the Intrinsic Loss Power (*ILP*) parameter.

It may be noted that although the focus of the study was on the inter-laboratory comparisons, in retrospect it would have been useful to have conducted more extended studies in at least one laboratory. A good example here would have been to undertake a series of field-dependent *SLP* measurements at a single site, for which systematic uncertainties would have been minimized, allowing the anticipated linear  $SLP - H_o^2 f$  relationship to be observed directly. Unfortunately, this was not considered at the time, and the uncertain aging characteristics of the samples ruled out subsequent measurements.

Interlaboratory repeatability analysis was then performed using the *ILP* data. An initial Mandel *h* statistics analysis (Figure 3) showed that with the exception of one outlier (Laboratory 1), all of the measurements fell well inside the  $\pm 0.5\%$  significance level. The outlier was also evident in Youden analysis (Figure 4), but the Youden plot also revealed that there was a clear systematic uncertainty component to the data, with the data points spread out along the ratio-of-means diagonal. The mean *ILP* values for both samples across both rounds of measurement were then calculated (excluding Laboratory 1) in Table 3, from which it was found that the round-to-round agreement was good, but that the

measurement accuracy was poor, with standard deviations of ca.  $\pm 30\%$  to  $\pm 40\%$ .

Rank correlation analysis (Figure 5) indicated significant correlation between the *ILP*s measured, with a Spearman coefficient of ca. 0.81, consistent with the notion that “some laboratories measured low, while others measured high”. This then led to an analysis of possible sources of systematic uncertainties:

- Sample degradation was considered as a potential source, but discounted on the basis of the study design, and on the absence of any reported issues with suspension stability.
- Unreported deviations in the magnetic field amplitude were considered, and a *post facto* numerical exercise performed to establish whether deviations between the reported  $H_0$  and a postulated ‘actual’  $H_0' = (1 + x) H_0$  might account for the systematic uncertainties. This analysis was based on the supposition that if there was such a systematic deviation (for example one due to an over- or under-estimation by the user of the current flowing in the coil), it would affect both samples’ measurements, in a given laboratory, by an equal amount. It was found that in 9/16 cases the intra-laboratory deviations differed by more than 5% between Samples 1 and 2, indicating that it was unlikely to be the primary source of the systematic uncertainties. Although other sources of unreported deviations in  $H_0$  are possible – such as those due to field inhomogeneity, and/or variable placement of samples within the calorimeter – these were thought more likely to result in random, rather than systematic, uncertainties.
- Possible equipment-related issues were considered by examining the subset of data from those participants whose experiments were performed on one of two particular commercial systems ( $N=5$  of each). Interestingly, the data showed that the System 1 users reported *ILP* values ca. 13% and 16% higher than the overall mean, while the System 2 users reported *ILP* values ca. 8% and 18% lower than the overall mean, for Samples 1 and 2 respectively.
- Thermometry methods were considered, and comparisons made between users of fiber-optic thermometers and of copper/constantan thermocouples – the latter supplied as standard with the System 2 commercial systems. In this case the fiber-optic users reported *ILP* values within ca. 4% of the overall mean, while the thermocouple users again reported *ILP* values ca. 8% and 18% lower than the overall mean. Potential systematic uncertainties due to probe positioning were also considered but discounted on the basis that a uniform probe-positioning requirement was included in the Round 2 measurement protocol, with no evident impact on the systematic uncertainties.
- Sample environment factors were considered for the non-adiabatic calorimeters used by all but one participant. Analysis of the ‘linear loss factor’  $L$  determined from the CSM-recalculations showed that although there were definite interlaboratory differences, there was no correlation

between  $L$  and *ILP* (Pearson coefficients of ca. 0.02 and 0.00 for Samples 1 and 2). Similarly, there were no clear correlations between the sample volume used,  $V$ , and either the  $L$  or the *ILP* parameters. Also, uniform sample volume was included in the Round 2 measurement protocol, with no impact on the systematic uncertainties. Lastly, data acquired during Round 2 on the sample environment by having the participants measure a ‘blank’ sample of pure water showed that in most cases the heat transfer to and from the sample into its environment was small, with typical  $\Delta T_{\max}$  values of ca. 0.1 K.

Although this analysis of factors that might lead to systematic uncertainties was not exhaustive, it does represent the authors’ best efforts at understanding the correlations in Figures 4 and 5, and the large standard deviations in Table 3. None of the factors considered is an obvious candidate as the sole source of the observed systematic variation. Hence, unless there is some other factor that has escaped our attention, the source must logically be a combination of factors which, in some way that we do not yet fully understand, work together to produce the observed systematic uncertainties.

Nevertheless, it is possible as a result of this analysis to compile a list of recommended best practice for researchers in this field, as follows:

1. The magnetic field strength and its homogeneity over the sample volume should be accurately measured and the frequency verified. All properties of the magnetic field should remain stable for the complete measurement time.
2. The heat flows into and out of the sample space during normal operation should be understood and accounted for in each apparatus. The possibility of inhomogeneous heat distribution across the sample should also be considered and accounted for in an appropriate way. The measurement of ‘blank’ samples of pure water using probes placed at various positions within the sample volume is a good way to check this.
3. Thermal probes should not generate false signals or additional heat in the applied time-varying magnetic field, for which reason the use of thermocouples (unless properly shielded) is not recommended. The accuracy and response time of the chosen probe should also be verified as appropriate for the anticipated range of temperature measurements.
4. The physical and magnetic properties and stability of the material undergoing measurement should be verified in detail. Vortexing of the sample prior to measurement is recommended, to counter possible incipient aggregation effects.
5. A well-defined and repeatable measurement protocol should be employed. The protocol used in this study – comprising three consecutive heating runs plus a cooling curve measurement – is available in the [Supplementary Information](#) and is recommended. Standardization of the sample volume and placement in

the calorimeter, and of the thermal probe (or probes) positioning within the sample, is also recommended.

6. Data analysis should be undertaken using, or with reference to, current best-practice methods, such as the freely available Corrected Slope Method employed in this study. If the CSM is used, attention should be paid to the linear loss factor, and this should be reported alongside the calculated *SLP* or *ILP* for a given sample.
7. Measurement reports, especially those published in the literature, should include details on the apparatus and measurement protocols used, and should always include both the applied field amplitude  $H_o$  and frequency  $f$ , alongside the chosen *SLP* or *ILP* metric. Uncertainties should be reported as the estimated random uncertainties for the laboratory in which the measurement was made, and, in instances where comparisons are made between reported measurements in different laboratories, attention should be paid to the potential impact of ca.  $\pm 30$ -40% systematic uncertainties of the kind described in this study.

## 5. Conclusions

Of all the physical properties measured routinely in laboratories around the world, thermal properties are arguably some of the most difficult to get right. Calorimetry measurements are fraught with unexpected and undetected factors that can confound results, especially when experiments are undertaken with non-adiabatic systems. In essence, this is the problem that faces many practitioners in the emerging field of magnetic field hyperthermia today.

In the interlaboratory study reported here we have surveyed the current state-of-the-art in magnetic field hyperthermia calorimetry in Europe. We have found that although there is evidently very good repeatability within a given laboratory, the overall measurement accuracy is poor, with a significant disparity between laboratories. For the two samples that were studied, across 17 laboratories, the reported magnetic heating metrics had standard deviations on the mean of approximately  $\pm 30\%$  to  $\pm 40\%$ . These are large uncertainties. Furthermore, we have found a strong systematic component to the uncertainties, coupled with a clear rank correlation between the measuring laboratory and the reported metric. Both of these are indications of a current lack of normalization in this field.

Through analysis of the potential factors leading to these systematic uncertainties, and comparison with the data provided by the study participants, we have identified a number of possible sources of uncertainty and have considered ways in which these can be alleviated or minimized. However, no single dominant factor was identified, and significant work remains to ascertain and remove the remaining uncertainty sources. In the meantime, the results presented here clearly demonstrate the need for standardized operating procedures in the hyperthermia characterization community. In addition, the development of verified reference materials is also an increasingly pressing requirement. In this context it is interesting to note that materials other than magnetic

nanoparticles may prove to be better reference materials for use in interlaboratory or calibration measurements [33].

Although we do not yet have all the answers regarding the origins of the systematic errors identified and quantified through this study, we do nonetheless have a positive outcome in the form of an agreed set of recommended best practice guidelines for magnetic hyperthermia characterization measurements. These are:

1. Verify the homogeneity and stability of the magnetic field strength and frequency applied, ideally using a suitably calibrated probe.
2. Understand and measure the heat flows into and out of the sample space, for example by recording both heating and cooling curves, and by keeping temperature excursions to moderate levels, of order  $\pm 10$ -20 °C.
3. Use reliable thermal probes that do not generate false signals or additional heat, such as fiber optic probes. (Avoid thermocouples.)
4. Verify the physical/magnetic properties of the sample; and avoid aggregation. Choose a sample concentration appropriate for the intended measurement, even if this means diluting the sample so that the heat generated is manageable.
5. Define and use a repeatable measurement protocol, with standardized sample volumes and placement, and standardized probe positioning. (The Radiomag protocol is recommended, and is given in the [Supplementary information](#).)
6. Perform at least three measurements on any given sample and estimate the random uncertainties associated with the measurement. (Repeatability levels should typically be within  $\pm 10\%$ , and preferably within  $\pm 5\%$ ; larger values may indicate instrumental or procedural issues that should be resolved.)
7. Review and use current best-practice methods for data analysis. (The CSM "calibrated slope method" is recommended, and is freely available [25]).
8. Always report experimental details,  $H_o$  and  $f$ , alongside the chosen *SLP* or *ILP* metric; and report estimates of the local random uncertainties derived from the repeat measurements.

In conclusion, we believe that our study provides a first step in the prenormative validation of magnetic field hyperthermia research methods, and will be an aid to future standardization development, and the continued transition of this exciting new technology from the laboratory to the clinic.

## Acknowledgements

The RADIOMAG consortium members participating in this study and not referenced in the authorship are explicitly named here. (See the [Supplementary Information](#) for a complete listing, with affiliations.) Nanoparticle synthesis was undertaken by: Dermot Brougham, Daniel Horak, Sophie Laurent, Claudio Sangregorio, Nguyễn Thanh, and Ladislav Vekas. Measurements were undertaken by: Manuel Bañobre, Marko Bošković, Dermot Brougham, Julian Carrey, Nico Cassinelli, Marco

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## Disclosure statement

No potential conflict of interest was reported by the author(s).

## Supplementary material

See the [Supplementary Material](#) for details of the recommended measurement protocol used in Round 2; and for full sets of the CSM-recalculated data as used in the study.

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## Data availability statement

The data that support the findings of this study are publicly available for download. The files are uploaded in the Zenodo repository under the title “Radiomag Interlaboratory Comparison of Magnetic Hyperthermia Characterisation Measurements: Data” DOI: <https://zenodo.org/record/4281154#.YEiMBk6g82w>.

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## Supplementary Information

### Challenges and Recommendations for Magnetic Hyperthermia Characterization Measurements

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37	E. Tombácz	University of Szeged	Hungary
38	L. Duc Tung	University College London	United Kingdom
39	V. Zavisova	Slovak Academy of Sciences	Slovakia

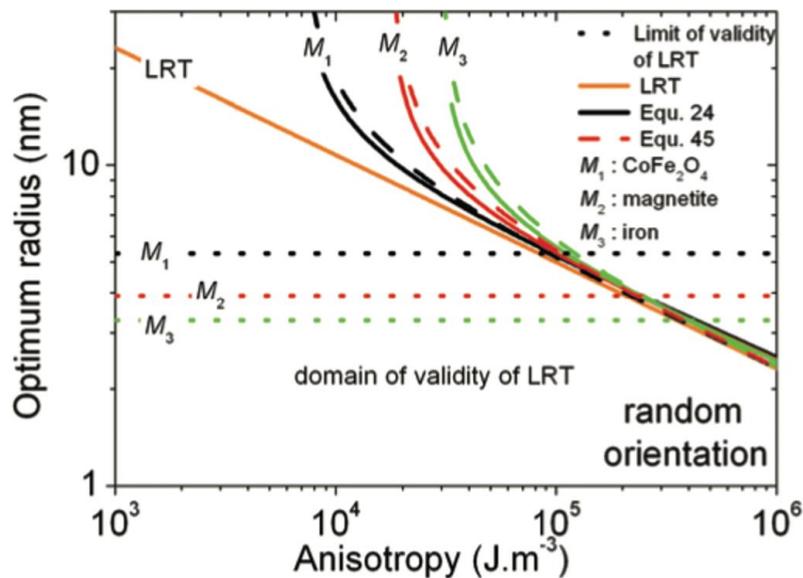
## 2. Additional Information re LRT Validity Discussion

### 2.1. Graphical Analysis

As noted in section 3.2 of the paper,

“Third, the data from **Table 2** was used to locate the samples on the LRT “validity map” introduced by Carrey *et al.*,<sup>12</sup> which allowed for consideration of both particle size and anisotropy, alongside the external conditions ( $H_o$  and  $f$ ). According to this graphical analysis, the parameters for both samples lay well within the LRT validity region (see Supplementary Information, **Figure S1**).”

The “validity map” referred to here is Figure 5(b) from Carrey 2011:\*



.. which summarises the results obtained from some of the theoretical models those authors employed to establish the reliability of the LRT approximation. The solid lines correspond to optimum radius (for maximal magnetic heating effect) predictions from both the LRT model and two other, more complex and more realistic models (Equations 24 and 45 in Carrey 2011), and the authors comment that it is only for anisotropies above ca. 100 kJ/m<sup>3</sup> (the “strongly anisotropic materials” regime), that the models agree.

However, superimposed on this diagram are horizontal dotted lines corresponding to the condition:

$$\xi = \mu_0 M_S V H_o / k_B T = 1 \quad ,$$

i.e. radii  $r_o$  for which the particle volume  $V = k_B T / \mu_0 M_S H_o$ , and the authors note that any particles that have a radius that is less than  $r_o$  lie within the “domain of validity of LRT”.

It may be noted that particles in this region are likely to be sub-optimal with respect to their magnetic heating potential, but regardless, they may be described using the linear response theory.

\* J. Carrey, B. Mehdaoui, and M. Respaud, *Simple models for dynamic hysteresis loop calculations of magnetic single-domain nanoparticles: Application to magnetic hyperthermia optimization*, *J. Appl. Phys.* **109**, 083921 (2011).

Applying this graphical representation of the “LRT validity map” to the two samples used in the Radiomag study requires estimations of the  $r_o$  values of both samples. Using the data presented in **Table 1** in the paper, viz.  $M_s = 356 \pm 22 \text{ kA m}^{-1}$  and  $485 \pm 30 \text{ kA m}^{-1}$  for Samples 1 and 2 respectively (at  $T = 300 \text{ K}$ ); and applying the experimental condition that  $H_o = 15 \text{ kA/m}$ , we obtain:

$r_o = 5.3 \pm 0.3 \text{ nm}$  for Sample 1; and

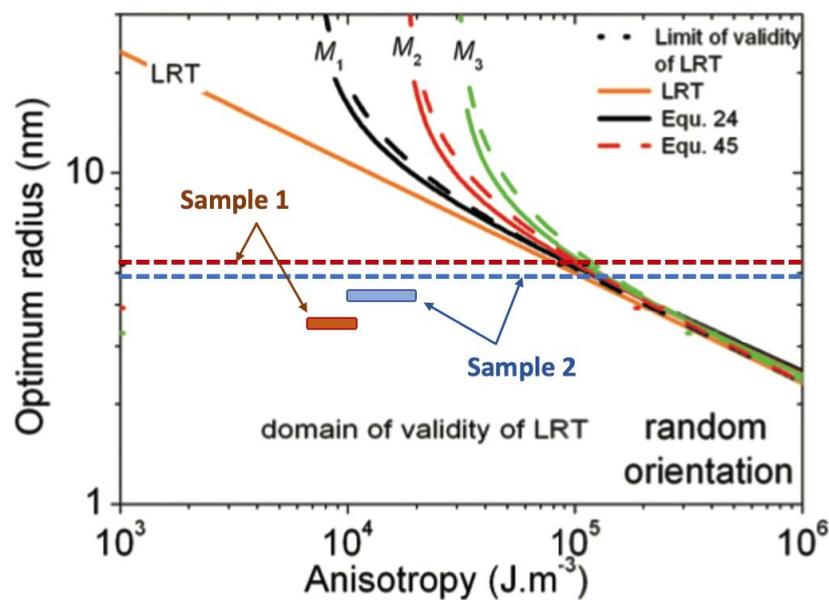
$r_o = 4.8 \pm 0.3 \text{ nm}$  for Sample 2.

Next, we take values for the effective anisotropy constants,  $K_{\text{eff}}$ , for both samples from recent experimental data published on maghemite and magnetite nanoparticles:

for maghemite nanoparticles,<sup>†</sup>  $K_{\text{eff}}$  may fall in the range from 6.5 to 10.8 kJ/m<sup>3</sup>; and

for magnetite nanoparticles,<sup>‡</sup>  $K_{\text{eff}}$  may fall in the range from 10 to 20 kJ/m<sup>3</sup>.

This then leads to the following graphical representation of the LRT status of the samples:



**Figure S1:** Graphical analysis of the position of the study samples on the “LRT validity map”.

.. from which it is apparent that both samples fall well within the domain of validity of linear response theory.

<sup>†</sup> A. I. Figueroa, J. Bartolomé, L. M. García, F. Bartolomé, A. Arauzo, A. Millán, and F. Palacio, *Magnetic Anisotropy of Maghemite Nanoparticles Probed by RF Transverse Susceptibility*, *Physics Procedia* **75**, 1050-7 (2015).

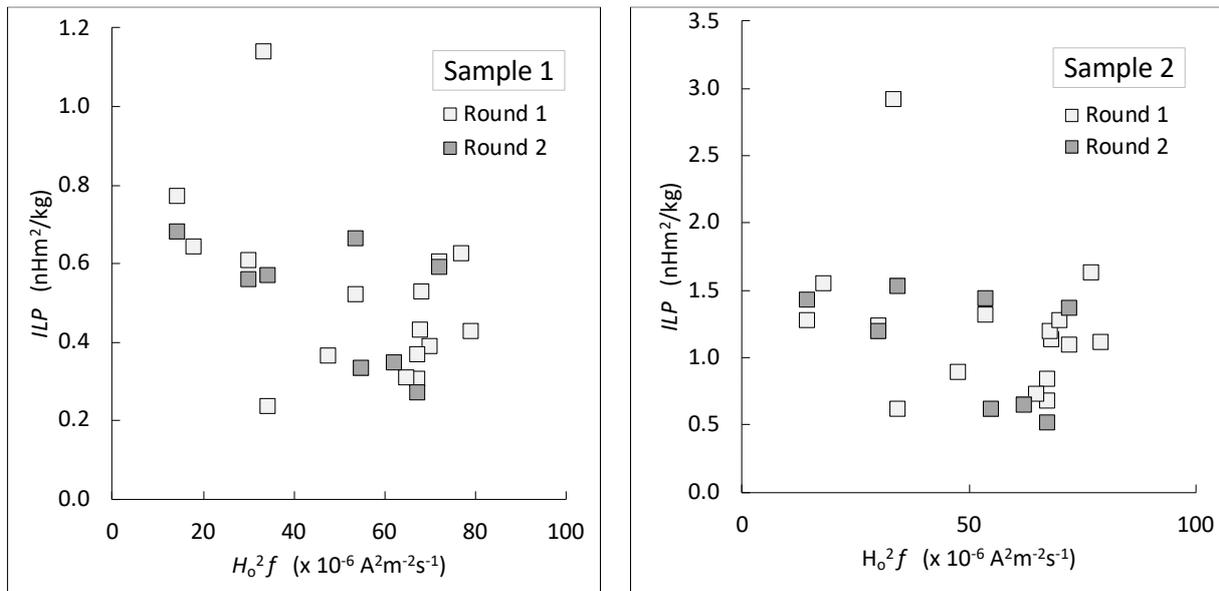
<sup>‡</sup> H. Mamiya, H. Fukumoto, J. L. Cuya Huaman, K. Suzuki, H. Miyamura, and J. Balachandran, *Estimation of Magnetic Anisotropy of Individual Magnetite Nanoparticles for Magnetic Hyperthermia*, *ACS Nano* **14**, 8421-32 (2020).

## 2.1. Graphical Analysis

As further noted in section 3.2 of the paper,

“Fifth, a complementary version of **Figure 2** was produced in which the *ILP* (rather than the *SLP*) was plotted against  $H_o^2f$  (see Supplementary Information **Figure S2**). No correlation at all was evident between the *ILP* and  $H_o^2f$  values, as is expected only in the LRT regime. The Pearson coefficients in this case were -0.12 and -0.26 for Samples 1 and 2 respectively.”

The figure referred to here is shown below:



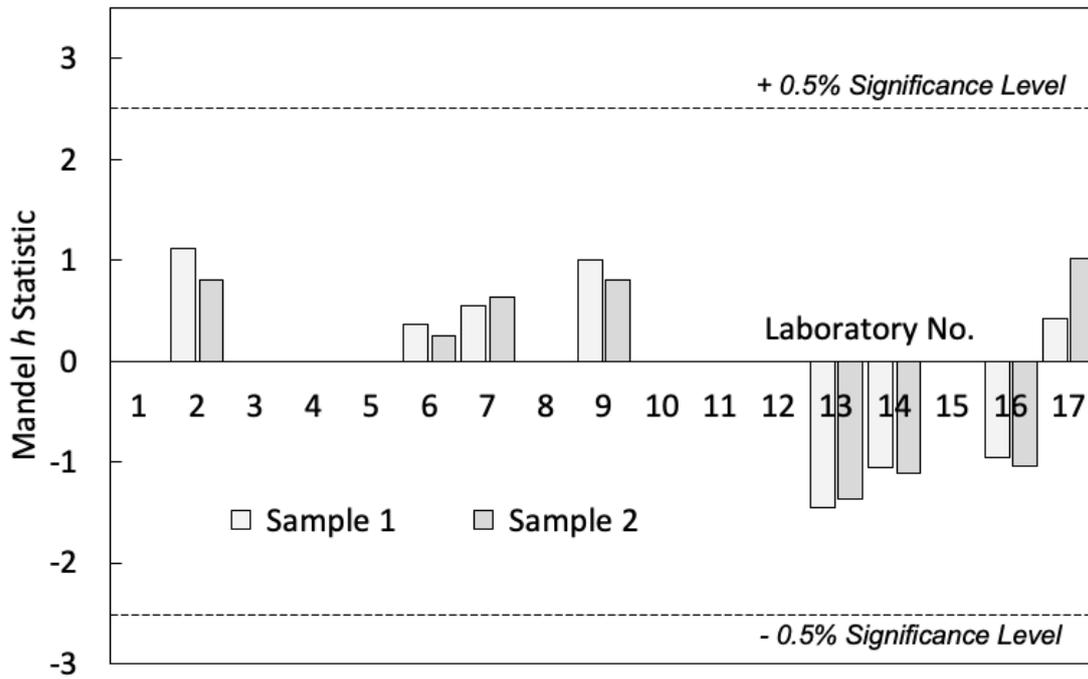
**Figure S2:** *ILP* values for Samples 1 and 2 plotted as a function of  $H_o^2f$ .

### 3. Mandel $h$ Statistics Plot for Round 2 Data

As noted in section 3.2 of the paper,

“For clarity, the Mandel  $h$  statistics in **Figure 3** are shown for the Round 1 measurements only; the corresponding data for Round 2 are less comprehensive but very similar, and are given in the Supplementary Information, **Figure S3**.”

The figure referred to here is shown below:



**Figure S3:** Mandel  $h$  statistics (relative to the mean) for Round 2 measurements, using the CSM-recalculated  $ILP$  values. Eight laboratories participated in Round 2: here they are given the same numerical label as was used in **Figure 3**.

## 4. Recommended Magnetic Hyperthermia Measurement Protocol

A transcript of the Excel file distributed to the measurement laboratories for providing measurement instructions and gathering the reported measurement data follows. This is the version distributed for the second round of the interlaboratory comparison study.

### **TAB #1: Instructions - Basic Data**

Keep in mind that it is critical to provide accurate answers, even if any errors have accidentally happened, as the participants will not be publicly identified by any means. Moreover, reporting any errors/artefacts detected and any corrective action taken is desirable, and will positively influence the study. Should you have any comment to make regarding the sample postage/delivery, storage or measurement, please encode it in the "Remarks and comments" section at the end of the form. Note that this section is not intended to make suggestions or give personal opinions; there will be a post-testing survey for the purpose.

#### *INSTRUCTIONS - HINTS*

Please follow the instructions given here to conduct the SAR measurements. If you are in doubt on how to proceed at some point or have any questions, please contact directly the experiment coordinators. Do not attempt to modify the present template unless for filling in the required data. Should you have any comments or remarks to make, please do so in the "Remarks & comments" tab.

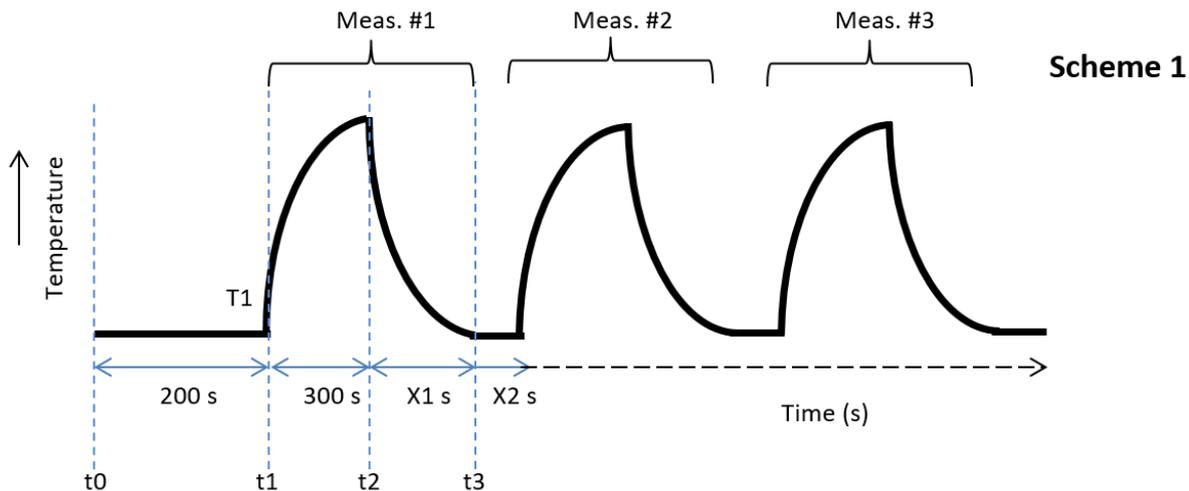
Cells to be filled by the measurers are highlighted in grey (and also in light blue in the case of the tab "Equipment") with a dotted border.

**ADIABATIC VS NON-ADIABATIC SETUPS.** It is important to understand the kind of setup you use; accordingly, we here provide the definitions of the two types considered for this particular inter-laboratory test. In simple terms, an adiabatic setup is that where there is no heat exchange with the environment (conduction, convection, radiation); this typically requires, among others, an "active" insulation jacket kept automatically to a certain temperature (that of the sample in strict adiabatic conditions, or a constant temperature in the case of semi-adiabatic or isoperibol calorimeters, where heat exchange corrections must be made). A non-adiabatic setup is any other where the former conditions (even just one) are not met.

- 1) Regardless the previous storage history, let the vial/s with the sample to equilibrate at room temperature for at least 2 hours before starting the experiment. Note that two vials of the same sample can be mixed up if required for any reason. Vials of different samples cannot be mixed up under any circumstances. Sonication or vortexing of the samples are allowed for homogenisation purposes, but only for periods of up to 5 minutes. If done, allow at least an additional 30 minutes for thermal equilibration of the sonicated/vortexed samples at room temperature.
- 2) The empty vial supplied can be placed directly into the instrument to make the measurements if the clearance of the coil permits. Transfer 1ml of the sample without diluting to the empty vial supplied using appropriate means (pipette/micropipette). The supplied vials should be used for the measurements unless your instrumental setup works only with either higher or lower sample volumes than the proposed one; in that case, take the required sample volume and report it in the relevant section "Volume used (ml)" of tabs "Sample#X" and subsequent ones. **IMPORTANT:** regardless the sample container, it is mandatory to measure and then note down the actual sample volume used in the C8 cell of the relevant tab of this file.
- 3) Always keep track of room temperature!

- 4) Positioning of the temperature probe: please centre the temperature probe as much as possible in the solution. In any case, totally immerse it into the sample (for both zero-conductivity water blank sample or the ferrofluids) and double check that it is not touching the walls of the container (whether using the supplied vials or your own containers)
- 5) Perform the following procedure with the zero-conductivity sample provided. Start the experiment by recording the temperature evolution from the initial point  $t_0$  during 200 seconds to a stable temperature  $T_1$  (see Scheme 1 below), then apply the field using a frequency of 300 kHz and the corresponding field intensity (peak, not rms) of 15 kA/m [If you cannot meet the latter conditions please use the closest possible, whichever, and report them in the corresponding cells in tabs "Sample#1" and subsequent ones] and continue recording  $T$  vs  $t$  under the applied field for 300 seconds, reaching the point  $t_2$ . After this time, remove the field and let the system cool down on its own down back to  $T_1$  for  $X_1$  s (it does not matter the time that it takes) while still acquiring the temperature evolution. Once  $T_1$  is reached again (point  $t_3$ ), keep acquiring the temperature evolution for  $X_2$  s (no less than 30 s) and stop the experiment. The whole procedure is graphically summarised in Scheme 1 at the bottom of this page. In order to ensure a better comparison between the different laboratories, we encourage you to use a time resolution of 1 second, i.e. acquire 1 data point per second. If your system does not allow you to change it, please do the corresponding averaging of the raw data prior to paste it into the relevant "EXPERIMENTAL SECTION (II)" so that the aforementioned time resolution is matched.
- 6) Repeat the procedure above three times to obtain three temperature vs time curves and three SAR values for each sample. Copy and paste the experimental temperature and time data in the section "EXPERIMENTAL DATA (II)" of tabs "Sample#4" and "Sample#6". Besides, encode the three SAR values of each sample in the corresponding cells in tabs "Sample#4" and "Sample#6" as the value of the slope of the temperature vs time curve. Use your own criteria/method to work out the slope of the curve (initial slope method, corrected slope method, Box-Lucas, etc.), but do not fit the curve starting from  $t_0$  in any case!!! Finally, use the concentration values provided in the corresponding tab for each sample.
- 7) Repeat exactly the same procedure described in point 5, this time for the ferrofluid samples to be tested.
- 8) Graphs are automatically generated from the experimental data inserted in the relevant sections, so please do not attempt to modify them. Remember that graphs in Excel can only accommodate up to 30000 data points; if exceeded, your data will be just partially displayed in the corresponding charts.
- 9) Make sure that all the sections through the tabs of this spreadsheet have been completed and email this file to XXXXX at your earliest convenience, preferably before the 23rd of April (23:59 CET).
- 10) Regarding the AC field, if your setup allows you to do so, choose "peak" instead of the "rms" signal.

Whether or not you are aware of other participants of this experiment, do not contact them regarding the results. Any enquiries about the whole test should be directed to XXXXX and XXXXX.



### TAB#2: Quick Step-by-step Instructions

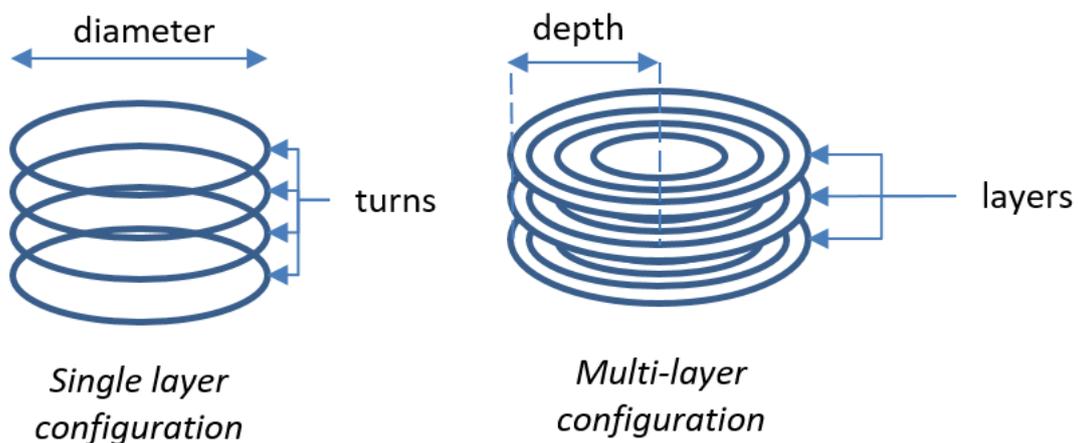
IMPORTANT: before proceeding with the following instructions you should have read carefully the previous tab "Instructions - basic data"; if not, please do so.

- 1) Sonication or vortex (5 mins. max.; not needed in the case of the water sample). Allow for thermal equilibration (30 minutes).
- 2) Transfer 1ml of the sample without diluting to the empty vial supplied using appropriate means (pipette/micropipette).
- 3) Position the temperature probe at the centre of the solution.
- 4) Using a resolution of 1 second, measure zero-conductivity water: record T vs t for 200 seconds after thermal equilibration, apply a 15 kA/m field at 300 kHz, record T vs t for 300 s, cool down back to the equilibrium temperature and keep measuring for at least 30 seconds.
- 5) Repeat the same procedure in point 4 with samples X and X.
- 6) Send data to XXXX before the 23rd of [deadline].

### TAB#3: Equipment

	<i>Click on the grey cells to expand the dropdown lists below. Choose the option and describe (light blue cells) as appropriate. Write N.A. for those options that do not apply.</i>
<b>Core method for T(t) measurements</b>	
<b>Nature of the equipment</b>	
<i>In the case of the two first options, please briefly indicate brand, model, and the customisation details:</i>	
<b>Sample holder - material</b>	
<i>If other, please specify:</i>	

<b>Sample holder - thermal insulation</b>	
<i>Please describe the means implemented in the measurement system to thermally insulate the sample (vacuum, etc.):</i>	
<b>Means of producing magnetic field</b>	
<i>Indicate the commercial system used; if a non-commercial one has been used, please give details on (alternatively, you can provide a picture):</i>	
<i>Number of turns</i>	
<i>Diameter/s</i>	
<i>Distance between turns</i>	
<i>Number of layers</i>	
<i>Distance between layers</i>	
<i>Depth</i>	
<i>Inductance (if known)</i>	
<b>Thermal insulation between the magnetic field source and the sample</b>	
<i>Please describe the means implemented in the measurement system to thermally insulate the sample from the magnetic field source:</i>	
<b>Type of temperature sensor</b>	
<i>If not part of a commercial system, please give details on the type of temperature sensor use to monitor the temperature during the measurements:</i>	



Scheme 2 depicting the different parameters required to collect the information asked about the coils used during the experiment.

#### TAB#4: Water Sample

Preliminary considerations:

(1) If the sample temperature has been controlled by any means (fridge, freezer, etc.) during storage, please indicate the temperature. If any (either controlled or uncontrolled) temperature change has happened during storage, please describe it in the "Remarks and comments" section at the end of the form.

(2) Each sample should be measured triplicate, but continuously during a single experiment as shown in Scheme 1 in the Introduction tab.

**Vial ID (barcode):**

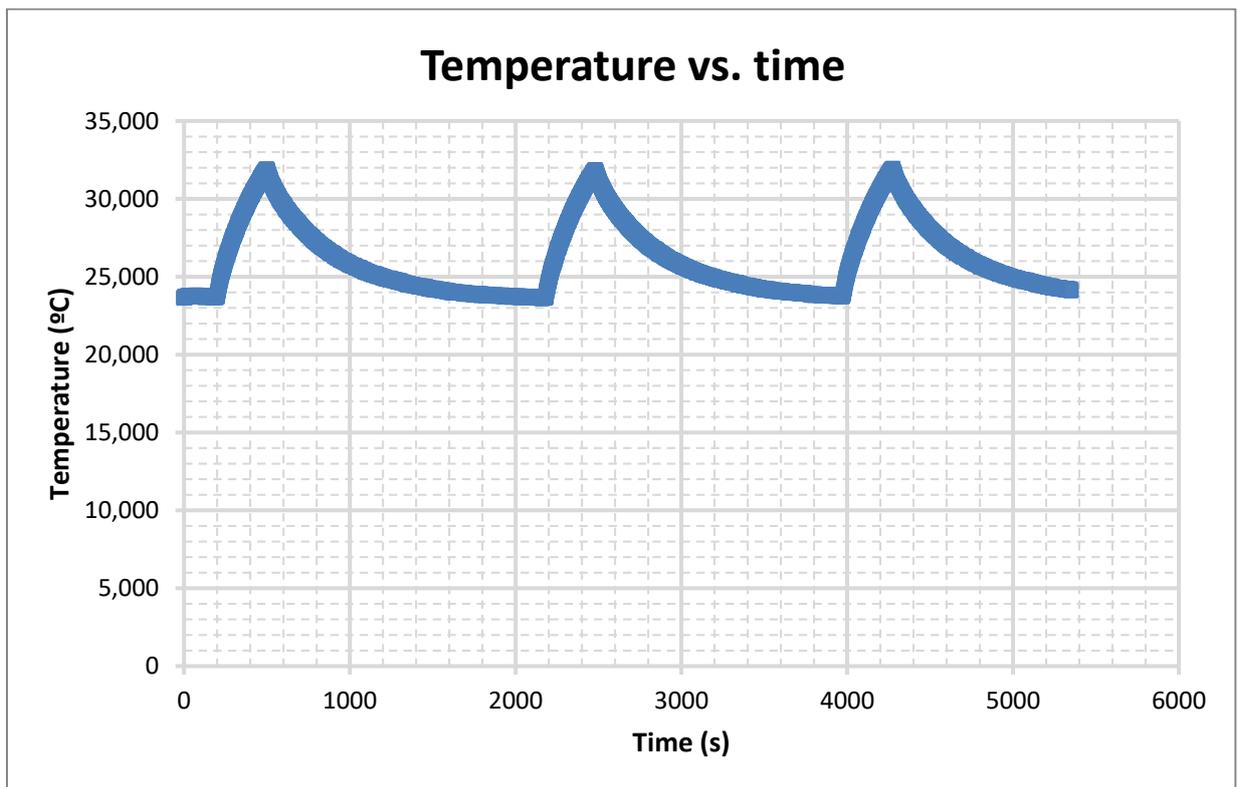
**EXPERIMENTAL DATA (II)**

*Insert the experimental T vs t data. Please use as many rows as necessary (the data will be plotted in the following tab).*

Time (s)	Temperature (°C)		
		<b>Vial ID (barcode):</b>	<input type="text"/>
		<b>Was the sample damaged upon receipt? Sedimentation/precipitation has occurred to some extent? Any other noticeable observations? (Y/N):</b>	<input type="text"/>
		Please describe if applies	<input type="text"/>
		<b>Volume used (ml):</b>	<input type="text"/>
		Have you used a regularly calibrated pipette? Y/N:	<input type="text"/>
		<i>Corrected volume</i>	0
		<b>Concentration (mg Fe/ml)</b>	N/A
		<b>Storage method</b>	
		Temperature control? (Y/N):	<input type="text"/>
		Exposed to ambient light? (Y/N):	<input type="text"/>
		Other:	<input type="text"/>

		<b>Storage temperature (°C):</b>	
		<b>Sample pre-treatment</b>	
		<b>Ultrasounds (Y/N):</b>	
		If yes, specify power, treatment time and temperature	
		If other treatment, please provide details	

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**Authors comment:** *The data shown here is just example. The graph is actually produced when the measurers insert their data points.*

**NOTES:**

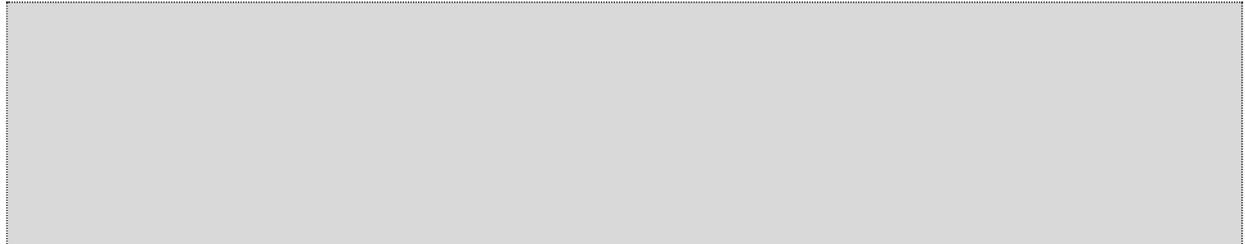
- Any comments regarding this section should be encoded in the final "Remarks&Comments" tab.
- Make sure that all the numerical values are given in the units specified at the column headings.

### **TAB#5: Other Samples**

The structure of this tab is exactly the same as the one presented in the preceding tab#4.

### **TAB#6: Remarks and Comments**

*Please make use of this tab to make any comments or clarifications regarding the experiments. Use as much space as needed.*



### **TAB#7: Corrections Log**

The following is the corrections log of the final version of the report template containing all the improvements made along all the previous rounds.

#### **March, 2016**

- 1- Cell I13's formula now points to the right range of cells containing the ILP values instead of the SAR ones in tabs "Sample#1" and "Sample#2".
- 2- The average curve in the graph in tab "Tvs.t#1" is now linked to the right data series from the corresponding data tab "Sample#1".
- 3- The content of cell B in "Instructions - basic data" tab has been removed as the date is already asked in tabs "Sample#1" and "Sample#2".

#### **April & May 2016**

- 4- Tab "Instructions - basic data": hints on the temperature probe positioning are now provided.
- 5- Tab "Instructions - basic data": differences between adiabatic and non-adiabatic setups for SAR measurements are now explained.
- 6- Tab "Instructions - basic data": experimental times (points t0, t1 and t2) have been modified, and so the Scheme 1.
- 7- Tab "Instructions - basic data": experimental procedure for the (problem) ferrofluids has been rewritten, including now the measurements of the blank sample of zero-conductivity water.
- 8- New tabs for samples #3, #4 are now available
- 9- New tabs for Tvs.t#3, Tvs.t#4 (corresponding to samples#3, #4) are now available.
- 10- Some typographic errors have been corrected.
- 11- Instructions regarding time resolution of the measurements have been added to the "Instructions - basic data" section.

### March 2017

- 12- A "QUICK STEP BY STEP INSTRUCTIONS" tab has been included after the introductory one. This quick guide is just to serve as a quick reference while doing the measurements. **IT DOES NOT SUBSTITUTE THE INTRODUCTION TAB!**
- 13- Scheme 1 in the "Instructions - basic data" tab has been modified to show the whole measurement procedure as it should happen.
- 14- The "Means of producing magnetic field" subsection in the "Equipment" section has been changed. There are more questions about the coil for producing the field. An explanatory scheme with indication of the coil parameters is now available.
- 15- The "Ambient temperature" subsection has been removed.
- 16- Data tabs are now much simpler. Average values and statistics have been removed. All the data from the three measurements (as they are now experimentally carried out continuously in a single session) may be now directly pasted to the corresponding two columns in tabs "Water\_sample", "Sample#3" and "Sample#4".
- 17- Graphs have been now integrated into the corresponding data tabs.

