Cardiac-evoked long-range quantum entanglement in the conscious brain

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Abstract

We report observations of cardiac-evoked long-range zero quantum coherence (ceZQC) in human brain tissue which were not accompanied by an alternation of short-range quantum coherence. For every cardiac pulsation, ceZQC emerged over the entire brain tissue for a period of 270 ± 180 ms. We observed this quantum effect in 40 human volunteers, if and only if they were awake. This link to conscious awareness suggests that the information held in the long-range quantum phenomenon is used and manipulated in conscious-related computation.

INTRODUCTION

The MRI signal, which is single quantum coherence (SQC), is indirectly effected by short-range multiple quantum coherence (MQC) via T_1 and T_2 relaxation. Long-range MQC [1] may not effect the MRI signal significantly but it can be observed with the help of field modulation in so-called multiple spin echoes (MSEs) [2–4]. A basic MSE sequence consists of asymmetric magnetic field gradients, in contrast to the conventional single spin echo (SSE) methods where symmetry is essential, combined with at least two radio-frequency (RF) pulses [2–4]. For any "classical fluid", the signal-to-noise ratio in MSE, which depends on the probability of long-range multiple quantum coherence, is much lower than in SSE. For example, the signal ratio of ZQC to SQC only reaches up to 0.02 at 4 Tesla, experimentally [5, 6]. A SSE sequence could also contain MSEs if repeated fast. So-called crusher gradients (to dephase any SQC signal that remains from the previous excitation) which are added between two image acquisitions (see Fig. 1), produce an asymmetry in the subsequent acquisition [7]. In consecutive echo-planar imaging (EPI) time series, field modulations to create MSEs are accomplished in even numbered repetition. The asymmetric gradient field as shown in Fig. 1 can generate zero quantum orders [5] and higher negative orders [8]. Of course, fast imaging sequences have a high noise level (around 0.02) which means that ZQC signals are negligible in fast imaging series for any classical liquids. This is also the intention of the sequence design. However, we found, unexpectedly, ZQC signals in fast MR imaging which were evoked by the cardiac pulse.

RESULTS AND DISCUSSION

We studied 40 subjects [9] (between 18 and 46 years old) using a 3.0 Tesla whole-body MRI scanner [10]. In fast gradient-echo EPI (GE-EPI) time series [11], we found periods of predominant signal alternations re-

Figure 1. Radio frequency (RF) and Gradient scheme of two consecutive EPI acquisition. The "asymmetry" field includes all asymmetric gradients of the ZQC.

peated with the cardiac frequency (Fig. 2). In each cardiac cycle, we observed a period of around 4-6 TRs (180-270ms) (which varied between 2 TRs (90ms) and up to 10 TRs (450ms) under special circumstances) (Fig. 4B) where the averaged signal amplitude increased by up to 15 %. The evoked signal alternation confirmed a ZQC contrast mechanism caused by the sequence design. It was also verified by altering the following sequence parameters; (a) rotation of the asymmetric gradients through slice rotation [12], (b) variation of the flip angle [13], and (c) alteration of the off-resonance frequency [14]. For the analysis, averaged maximum peak (AMP) and baseline signal (BS) [15] from preprocessed data [16] were calculated. For (a), we found the characteristic angulation dependency of the dipole-dipole interaction as shown is (Fig. 3(A)). The plot represents the fitted function $|(3 \cdot cos^2[\varphi] - 1)|$ (adjusted R^2

test of goodness-of-fit resulted in R^2=0.9958) where φ takes the additional gradients in read and phase direction into account which relates to the slice angulation as $\varphi = \alpha - \tan^{-1}\left[\frac{\mathrm{Gt}_{cru} - \mathrm{Gt}_{rt}}{2\mathrm{Gt}_{sat} + \mathrm{Gt}_{cru} + \mathrm{Gt}_{st}}\right] = \alpha - 9.6°$. At the magic angle, the cardiac evoked signals disappeared completely which rules out any SQC component. Consequently, we can exclude that the signal was generated by changes in T$_1$ and T$_2$ relaxation, line narrowing, or magnetic field shifts. It also means that short-range ZQC weren't significantly altered. For (b), we found the predicted signal course for the ZQC flip angle dependency [8] which was fitted to the data (R^2=0.9964). Predicted maximum of AMP at 45° could be confirmed (Fig 3(B)). In contrast, the Ernst-angle [17] which is a good indication for the optimum angle for SQC is around 13° (for T$_1$ = 1.5s). For (c), we found a typical magnetization transfer contrast (MTC) change for the BS which depended on the off-resonance frequency (Fig. 3(C)). In contrast, the AMP intensity shows the remarkable immunity to MTC as expected for ZQC [18] with no significant changes in the same frequency range (Fig. 3(D)).

Figure 2. Whole-slice averaged signal time course (black line) which was selected by a mask over 12 heart cycles. Signal of the Superior sagittal sinus (grey line) as reference time frame demonstrates the instant breakdown of quantum coherence with the beginning outflow.

Furthermore, the cardiac-evoked ZQC (ceZQC) also showed unseen physiological properties of which we observed the following here:

(1) Temporal occurrence – In our experiments, the ceZQC always appeared between the arterial inflow and the venous outflow phase. The finding was established with the help of reference time frames from (a) finger pulse oximetry (supplementary material (Fig. 4)), (b) from the MRI data itself by using a voxel including the superior sagittal sinus (Fig. 2) and (c) from a data set with a longer repetition time (TR) of 70 ms, which also allowed the simultaneous observation of arteries (supplementary material (Fig. 5)). All three methods confirmed a perfect synchronization with the cardiac pulse.

(2) Spatial distribution – Further, we explored the distribution of the ceZQC over the entire brain. 9 slices (in 5 volunteers) were acquired at different positions, with slices from bottom to the top covering all anatomical regions. Test tubes with a water solution [19] were positioned close to the head for reference signals. We located ceZQC in brain tissue of all slices except around the periventricular area (probably due to movement induced by ventricular pulsation in those regions [20]) as shown in the supplementary material (Fig. 2). Regions like the skull and the attached test tubes showed no ceZQC excluding external signal sources. The sensitivity to movement could be confirmed in a hyperventilation challenge [21], where head movement was slightly altered. There, the signal declined immediately at the start of the challenge before any cerebral blood flow response could occur (supplementary material (Fig. 3)). The global aspect conformed with another interesting feature; the ceZQC signal could be restored while being averaged over the entire tissue component of the imaging slice (Figures 2 and 4(B)), which requires a traveling pulse that moves faster than the blood flow.

Figure 3. Variation of sequence parameters. Data shows signal averaged over 5 subjects. Error bars represent the standard deviation from the mean. **(A)** Signal intensity plotted against the slice gradient angulation α in respect to the magnetic main field. **(B)** Signal plotted against flip angle variation. ZQC prediction plotted in Black. **(C)** Signal intensity plotted against the frequency offset of the saturation slices of the averaged baseline signal and **(D)** averaged signal of the zigzag.

(3) Blood flow relation – In a breath-holding challenge [22], we found that the ceZQC amplitude did not

respond to the CO_2 challenge (Fig. 4(B)), despite the fact, that we observed an increase of blood flow in major blood vessels during the challenge (Fig. 4(A)). Furthermore, we found in a fMRI study [23], that local increase in blood perfusion during visual activation left the signal intensity undisturbed, too. We conclude from those observations that the signal intensity is not correlated to the cerebral blood flow directly. Further, we found that the AMP did not vary significantly with slice thickness [24] which confirmed the exclusion of movement and inflow as possible signal source (supplementary material (Fig. 6(A))). For the imaging repetition time (TR) [25], which defines the time-of-flight, no inflow effect could be detected either (supplementary material (Fig. 6(B))).

Figure 4. **(A)** Signal time course of an imaging voxel located next to the Superior sagittal sinus demonstrates the blood flow increase in response to the CO_2 challenge (breath-holding). In contrast to the vein signal, the corresponding ceZQC signal **(B)** showed no response to CO_2 activity. Breath-holding started at 140s. Volunteers were instructed to reduce any movement as long as possible (here until at 157s). From 157s, the signal breakdown was subject to movement.

(4) Conscious awareness – We found in seven participants from whom two had reported to have fallen asleep, that the ceZQC signal pattern changed dramatically as

shown in Fig. 5. This dramatic signal pattern changed gradually during the transition from awake to asleep as shown in the supplementary material (Fig. 7). In supplementary material at (Fig. 8), the results of a Recurrence Quantification Analysis and a Multifractal Detrended Fluctuation Analysis (for detailed description see Lopez-Perez et al. [26]) are plotted over all time points. The plots illustrate the difference between wakefulness and the slow decline during the falling asleep period. The analysis shows that periodicity, large and small fluctuations, repeating patterns and their predictability, and the stability of the system were changing over the observation period. In-depth analysis of sleep pattern and the effect of aging will be reported elsewhere [26]. For the final data acquisition, all participants had been asked to stay awake during the imaging protocol. Then, we no longer detected a sleep pattern.

Figure 5. Pattern observed in participant who had reported falling asleep. **(A)** Wake period. **(B)** Asleep, ceZQC signal declined with an increase of signal noise level. At 34 s, the peak resulted from short head movement.

In conclusion, our observation connects cardiac pulsation, which is a basic necessity for consciousness, with microscopic phenomena of the quantum world. A change of multiple quantum coherence, as observed here, is generally associated with a phase transition, which is consistent with our findings in a way that ceZQC was not caused by inflow or movement and further that it appeared and vanished in an instant and global fashion. However, a "classical" phase transition is always accompanied by a significant change of short-range multiple quantum coherence which was absent in our experiments. On the other hand, the existence of long-range (but without short-range) entanglement is the essence of topological order [27]. If we now consider that quantum coherence is equivalent to quantum entanglement for all practical purposes [28], then our observation suggests that we have found a topological-like phase in the conscious brain. Of course, it is not a topological phase as known from low temperature physics. However, active matter shows superfluid-like flow [29, 30]; similarly a topological-like

phase may evolve in the brain featuring long-range quantum entanglement. This topological-like phase could explain the link to conscious awareness; Taking into account that biological systems contain diversities of topological defects [31–34], which may also move in a cèilidh dance fashion [35], then it is intriguing to assume that such defects are manipulating the information held in the topological-like order [36]. Such a process may provide a way to perform topological-like quantum computations which would imply that consciousness, and probably cognition also, is based on quantum computing.

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[7] Two saturation pulses placed parallel to the imaging slice (Fig. 1) are added which allow to vary the long-range correlation of the ZQC.

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[9] Imaging protocols were approved by the Trinity College School of Medicine Research Ethics Committee. All participants of final data acquisition were asked to stay awake and stay still during the imaging protocol or to report any failure.

[10] The scanner (Philips, The Netherlands) was operated with a 32-channel array receiver coil.

[11] Initial experiments were carried out to establish a protocol that could deliver stable cardiac related signals over a range of subjects. The finalized parameters were as follows: FA = $45°$, TR = 45 ms and the TE = 5 ms with a voxel size was 3.5 x 3.5 x 3.5 mm, matrix size was 64x64, SENSE factor 3, bandwidth readout direction was 2148 Hz, saturation pulse thickness/distance 5/20mm.

[12] Slice angulation started from coronal $0°$ to axial $90°$ in the steps as [0, 10, 20, 30, 40, 45, 50, 55, 60, 65, 70, 80, 90]; Saturation gradients had a time integral (length x strength) of Gt_{sat} = 5.1 ms x 6.25 mT/m, the crusher gradients in read and slice direction of Gt_{cru} = 1.3 ms x 25 mT/m, the slice rephase gradient of Gt_{sr} = 0.65 ms x 25 mT/m, the slice termination gradient of Gt_{st} = 0.65 ms x 15 mT/m, and the total read dephase after EPI readout gradient of Gt_{rt} = 0.65 ms x 22.5 mT/m.

[13] The flip angle was varied from $5°$ to $60°$ in steps of $5°$($60°$ was the power limit by the specific absorption rate (SAR)).

[14] The off-resonance frequencies was varied as [2.62, 3.49, 4.36, 5.23, 6.11, 6.98, 7.84, 8.73, 9.60, 10.47, 12.22, 13.96, 15.71, 17.45] kHz.

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