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mutants display enhanced RNAi even in the first generation of treatment, it is possible that MET-2 down-regulates factors that promote heritable RNAi. This model is supported by recent work demonstrating that a feedback response between endogenous and exogenous RNAi pathways controls the duration of heritable RNAi [19].

These new findings raise several important questions concerning the relationship between small RNAs and chromatin modifications that will hopefully steer us towards a clearer understanding of transgenerational inheritance. C. elegans has been at the forefront in recent years in identifying roles for small RNAs in transgenerational gene silencing. The extent to which small RNAs affect transgenerational gene silencing in other species is poorly understood. Worms can propagate a small RNA signal at each generation via the activity of RNAdependent RNA polymerases, a class of enzymes lacking in mammals, and it is unclear if small RNA-based mechanisms of transgenerational silencing exist in humans [2]. However, in flies, which also lack RNA-dependent RNA polymerases, piRNAs are transgenerationally inherited to maintain piRNA production and transposon silencing from one generation to the next [20]. It will be important to identify the mechanisms underlying transgenerational inheritance across different species.

REFERENCES

- Holoch, D., and Moazed, D. (2015). RNAmediated epigenetic regulation of gene expression. Nat. Rev. Genet. 16, 71–84.
- Henikoff, S., and Greally, J.M. (2016). Epigenetics, cellular memory and gene regulation. Curr. Biol. 26, R644–R648.
- Rechavi, O., Houri-Ze'evi, L., Anava, S., Goh, W.S., Kerk, S.Y., Hannon, G.J., and Hobert, O. (2014). Starvation-induced transgenerational inheritance of small RNAs in C. elegans. Cell 158, 277–287.
- 4. Rechavi, O., Minevich, G., and Hobert, O. (2011). Transgenerational inheritance of an acquired small RNA-based antiviral response in C. elegans. Cell *147*, 1248–1256.
- Ashe, A., Sapetschnig, A., Weick, E.M., Mitchell, J., Bagijn, M.P., Cording, A.C., Doebley, A.L., Goldstein, L.D., Lehrbach, N.J., Le Pen, J., *et al.* (2012). piRNAs can trigger a multigenerational epigenetic memory in the germline of C. elegans. Cell *150*, 88–99.
- 6. Shirayama, M., Seth, M., Lee, H.C., Gu, W., Ishidate, T., Conte, D., Jr., and Mello, C.C.

(2012). piRNAs initiate an epigenetic memory of nonself RNA in the C. elegans germline. Cell 150, 65–77.

- Luteijn, M.J., van Bergeijk, P., Kaaij, L.J., Almeida, M.V., Roovers, E.F., Berezikov, E., and Ketting, R.F. (2012). Extremely stable Piwi-induced gene silencing in Caenorhabditis elegans. EMBO J. 31, 3422–3430.
- Lev, I., Seroussi, U., Gingold, H., Bril, H., Anava, S., and Rechavi, O. (2017). MET-2-dependent H3K9 methylation suppresses transgenerational small RNA inheritance. Curr. Biol. 27, 1138–1147.
- Fire, A., Xu, S., Montgomery, M.K., Kostas, S.A., Driver, S.E., and Mello, C.C. (1998).
 Potent and specific genetic interference by double-stranded RNA in Caenorhabditis elegans. Nature 391, 806–811.
- Vastenhouw, N.L., Brunschwig, K., Okihara, K.L., Muller, F., Tijsterman, M., and Plasterk, R.H. (2006). Gene expression: long-term gene silencing by RNAi. Nature 442, 882.
- Towbin, B.D., Gonzalez-Aguilera, C., Sack, R., Gaidatzis, D., Kalck, V., Meister, P., Askjaer, P., and Gasser, S.M. (2012). Step-wise methylation of histone H3K9 positions heterochromatin at the nuclear periphery. Cell 150, 934–947.
- Buckley, B.A., Burkhart, K.B., Gu, S.G., Spracklin, G., Kershner, A., Fritz, H., Kimble, J., Fire, A., and Kennedy, S. (2012). A nuclear Argonaute promotes multigenerational epigenetic inheritance and germline immortality. Nature 489, 447–451.
- Simon, M., Sarkies, P., Ikegami, K., Doebley, A.L., Goldstein, L.D., Mitchell, J., Sakaguchi, A., Miska, E.A., and Ahmed, S. (2014). Reduced insulin/IGF-1 signaling restores germ cell immortality to caenorhabditis elegans Piwi mutants. Cell Rep. 7, 762–773.

- Andersen, E.C., and Horvitz, H.R. (2007). Two C. elegans histone methyltransferases repress lin-3 EGF transcription to inhibit vulval development. Development *134*, 2991–2999.
- McMurchy, A.N., Stempor, P., Gaarenstroom, T., Wysolmerski, B., Dong, Y., Aussianikava, D., Appert, A., Huang, N., Kolasinska-Zwierz, P., Sapetschnig, A., et al. (2017). A team of heterochromatin factors collaborates with small RNA pathways to combat repetitive elements and germline stress. Elife 6.
- Phillips, C.M., Brown, K.C., Montgomery, B.E., Ruvkun, G., and Montgomery, T.A. (2015). piRNAs and piRNA-dependent siRNAs protect conserved and essential C. elegans genes from misrouting into the RNAi pathway. Dev. Cell. 34, 457–465.
- de Albuquerque, B.F., Placentino, M., and Ketting, R.F. (2015). Maternal piRNAs are essential for germline development following de novo establishment of endo-siRNAs in Caenorhabditis elegans. Dev. Cell. 34, 448–456.
- Sapetschnig, A., Sarkies, P., Lehrbach, N.J., and Miska, E.A. (2015). Tertiary siRNAs mediate paramutation in C. elegans. PLoS Genet. *11*, e1005078.
- Houri-Ze'evi, L., Korem, Y., Sheftel, H., Faigenbloom, L., Toker, I.A., Dagan, Y., Awad, L., Degani, L., Alon, U., and Rechavi, O. (2016). A tunable mechanism determines the duration of the transgenerational small RNA inheritance in C. elegans. Cell 165, 88–99.
- 20. Le Thomas, A., Stuwe, E., Li, S., Du, J., Marinov, G., Rozhkov, N., Chen, Y.C., Luo, Y., Sachidanandam, R., Toth, K.F., et al. (2014). Transgenerationally inherited piRNAs trigger piRNA biogenesis by changing the chromatin of piRNA clusters and inducing precursor processing. Genes Dev. 28, 1667–1680.

Human Memory: Brain-State-Dependent Effects of Stimulation

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A new study shows that direct stimulation of memory-relevant brain areas can enhance memory performance, but only when stimulation is applied during brain states associated with poor memory outcome — stimulation during optimal states results in a decrease in memory.

Zaphod Beeblebrox — a character in Douglas Adams, comic sci-fi novel "*The Hitchhiker's Guide to the Galaxy* [1]" — is in a rather confused state when his spaceship lands on planet 'Vogtsphere'. Conveniently, he has at hand a 'thinking





Figure 1. Illustration of the stimulation protocol and results of Ezzyat *et al.* [3]. (A) Intracranial EEG is recorded during a memory task. Brain activity during encoding is split into two classes (hits or misses) based on later memory performance. (B) A classifier is trained on the data to identify states that are associated with 'optimal' or 'poor' memory based on the spectral profile. Electrical stimulation during 'optimal states' reduces memory performance, whereas stimulation during 'poor states' increases memory performance.

cap', a device that electrically stimulates the brain in order to improve cognitive function. This intuition that electrical stimulation modifies brain function is not only evident in various sci-fi novels, but actually has a long standing history in cognitive neuroscience. For instance, over half a century ago Wilder Penfield [2] pioneered the technique of pre-surgical mapping, whereby brain areas that underlie specific cognitive and motor functions are mapped by applying electrical pulses to the brain tissue. A given area is assumed to be functionally relevant if electrical stimulation interferes with the associated cognitive or motor function, as manifested, for example, by interruptions or difficulties in the naming of objects during the stimulation of language areas (Wernicke's area, for example). But can this same stimulation technique be utilized in a way that does not disrupt but instead enhances cognitive performance? As they reported very recently in Current Biology, Ezzyat et al. [3] have developed a new approach to brain stimulation, obtaining results that show that brain stimulation is capable of improving memory, but only when applied during certain brain states.

The relationship between brain activity and memory is commonly studied with so-called subsequent memory experiments [4], wherein a list of items, for example words, is presented sequentially to a participant who then has to recall the items during a later test. Based on the participant's recall performance during the test phase, brain activity during the learning phase can be classified into 'subsequent hit trials' (items that were later recalled) or 'subsequent miss trials' (items that could not be recalled). Contrasting the internally generated brain activity between these two classes of items results in a so-called subsequent memory effect, which quantifies the difference between brain activity during subsequently recalled and forgotten items (Figure 1A). The former is typically associated with a pattern of increased high frequency

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power and decreased low frequency power, whereas the latter is characterized by a pattern of decreased high frequency power and increased low frequency power [5,6]. This subsequent memory effect suggests that the brain naturally fluctuates between states that do or do not facilitate memory formation. These 'optimal' and 'poor' memory states are characterized by distinct spectral profiles of electrical brain activity. Accordingly, Ezzyat et al. [3] hypothesized that the effect of electrical stimulation on memory performance may depend on whether stimulation is applied during 'optimal' or 'poor' memory states.

To examine this hypothesis, the authors tested patients with refractory epilepsy who were implanted with depth and surface grid electrodes for pre-surgical diagnostic purposes. The experiment followed a two-stage procedure. In the first stage (Figure 1A), patients performed a memory task where they learned a list of words, which they had to recall later. During this stage no stimulation was carried out; instead the patients' electrical brain activity was recorded and subsequent memory effects were identified by means of a pattern classifier algorithm. As hypothesized by the authors, a consistent picture emerged across patients, in which high frequency power increases and low frequency power decreases predicted later successful retrieval - indicative of an optimal memory state. Conversely, low frequency power increases and high frequency power decreases predicted later misses - indicative of a 'poor' memory state.

In the second stage (Figure 1B), the patients again performed the same memory task (with different words), this time receiving stimulation, in the form of electrical pulses at 50 Hz, for half of the words, whereas the other half served as a baseline control. The stimulated sites differed between patients but were mostly memory-relevant regions like the medial temporal lobe or dorsolateral prefrontal cortex. As expected, electrical stimulation led overall to a small but statistically significant memory decrease. However, when taking into account the brain state during which stimulation was applied, a drastically different pattern of results arose. When being stimulated during optimal memory states, patients showed

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worse memory compared to when not being stimulated; importantly, however, when the patients were stimulated during poor memory states, their memory improved significantly.

These are exciting results as they show that natural fluctuations in brain states, which are indicated by the frequency spectrum of the EEG, can account for the variable effects of brain stimulation. This could potentially resolve the question why stimulation studies with similar stimulation protocols show discrepant results, with some reporting improved memory [7] and others impaired memory during stimulation [8,9]. Furthermore, these results show that it is possible to increase cognitive performance, but only when stimulation is applied during states which indicate non-efficient information processina.

The study by Ezzyat et al. [3] opens up a number of interesting questions to be addressed by future experiments. There are three questions that we think are most pressing. First, is it possible to improve memory performance online by selectively stimulating during brain states correlated with poor processing? Notably, this is still an open question, as the authors stimulated throughout the memory task and obtained the results offline after splitting the data post-hoc into good and bad memory states. Addressing this question requires a closed-loop stimulation protocol [10,11] in which specific brain states are targeted online, that is, during a memory task, based on a priori defined brain states.

Second, what is the neurophysiological mechanism by which electrical stimulation during poor brain states boosts memory? One possibility is that a simultaneous increase in low frequency power and a decrease in high frequency power may reflect an inhibited state of a cortical region. An unspecific high frequency electrical stimulus could act as an excitatory drive to a given area that causes it to switch from a passive to an active state, thus mimicking a 'wake-up call' for the network. Interestingly, a study in animals showed that stimulation of the cortex during a passive state increases neural firing, whereas stimulation during an active state induces a decrease in neural firing [12]. These findings from animals fit perfectly with the opposing effects on memory reported by Ezzyat

et al. [3] and are consistent with the observation that electrical stimulation during poor memory states induced increased high frequency activity (which can be taken as a proxy of increased excitation).

Third, can this approach be utilized for non-invasive state-dependent brain stimulation in order to increase memory performance in healthy subjects? Transcranial magnetic (TMS) and transcranial electrical stimulation (TES) are currently the most used non-invasive brain stimulation techniques and hold the promise of becoming tomorrow's tools of cognitive enhancement [13,14]. However, each of these techniques has its limitations: in the case of TES, these are poor spatial resolution and attenuation of currents as they travel from the scalp to the brain; and in the case of TMS is poor ability to reach deep brain structures such as the hippocampus (but see [14]).

Nevertheless, the ability to boost memory via non-invasive stimulation might increase considerably if fluctuations between brain states is taken into account, as highlighted in the study by Ezzyat et al. [3]. Importantly, in order to follow this example EEG and/or MEG should be simultaneously recorded during magnetic [15,16] or electrical stimulation [17,18], which is not done routinely at the moment. Together, the new results open the way for the development of closed loop stimulation protocols in order to increase brain function, thus moving 'thinking caps' from the realm of science fiction into reality.

REFERENCES

- Adams, D. (1980). The Hitchhiker's Guide to the Galaxy, 1st American Edition (New York: Harmony Books).
- 2. Penfield, W., and Boldrey, E. (1937). Somatic motor and sensory representation in the cerebral cortex of man as studied by electrical stimulation. Brain 60, 389–443.
- Ezzyat, Y., Kragel, J.E., Burke, J.F., Levy, D.F., Lyalenko, A., Wanda, P., O'Sullivan, L., Hurley, K.B., Busygin, S., Pedisich, I., *et al.* (2017). Direct brain stimulation modulates encoding states and memory performance in humans. Curr. Biol. *27*, 1251–1258.
- Paller, K.A., and Wagner, A.D. (2002). Observing the transformation of experience into memory. Trends Cogn. Sci. 6, 93–102.
- Long, N.M., Burke, J.F., and Kahana, M.J. (2014). Subsequent memory effect in

intracranial and scalp EEG. Neuroimage 84, 488–494.

- 6. Hanslmayr, S., Staudigl, T., and Fellner, M.C. (2012). Oscillatory power decreases and long-term memory: the information via desynchronization hypothesis. Front. Hum. Neurosci. 6, 74.
- Suthana, N., Haneef, Z., Stern, J., Mukamel, R., Behnke, E., Knowlton, B., and Fried, I. (2012). Memory enhancement and deep-brain stimulation of the entorhinal area. N. Engl. J. Med. 366, 502–510.
- Merkow, M.B., Burke, J.F., Ramayya, A.G., Sharan, A.D., Sperling, M.R., and Kahana, M.J. (2016). Stimulation of the human medial temporal lobe between learning and recall selectively enhances forgetting. Brain Stimul. http://dx.doi.org/10.1016/j.brs.2016.12.011.
- Jacobs, J., Miller, J., Lee, S.A., Coffey, T., Watrous, A.J., Sperling, M.R., Sharan, A., Worrell, G., Berry, B., Lega, B., *et al.* (2016). Direct electrical stimulation of the human entorhinal region and hippocampus impairs memory. Neuron *92*, 983–990.
- 10. Brittain, J.S., Probert-Smith, P., Aziz, T.Z., and Brown, P. (2013). Tremor suppression by rhythmic transcranial current stimulation. Curr. Biol. 23, 436–440.
- Morrell, M. (2006). Brain stimulation for epilepsy: can scheduled or responsive neurostimulation stop seizures? Curr. Opin. Neuro. 19, 164–168.
- Allen, E.A., Pasley, B.N., Duong, T., and Freeman, R.D. (2007). Transcranial magnetic stimulation elicits coupled neural and hemodynamic consequences. Science 317, 1918–1921.
- **13.** Kadosh, R.C. (2013). Using transcranial electrical stimulation to enhance cognitive functions in the typical and atypical brain. Transl. Neurosci. *4*, 20–33.
- Wang, J.X., Rogers, L.M., Gross, E.Z., Ryals, A.J., Dokucu, M.E., Brandstatt, K.L., Hermiller, M.S., and Voss, J.L. (2014). Targeted enhancement of cortical-hippocampal brain networks and associative memory. Science 345, 1054–1057.
- Hanslmayr, S., Matuschek, J., and Feliner, M.C. (2014). Entrainment of prefrontal beta oscillations induces an endogenous echo and impairs memory formation. Curr. Biol. 24, 904–909.
- Thut, G., Veniero, D., Romei, V., Miniussi, C., Schyns, P., and Gross, J. (2011). Rhythmic TMS causes local entrainment of natural oscillatory signatures. Curr. Biol. 21, 1176– 1185.
- Helfrich, R.F., Knepper, H., Nolte, G., Struber, D., Rach, S., Herrmann, C.S., Schneider, T.R., and Engel, A.K. (2014). Selective modulation of interhemispheric functional connectivity by HD-tACS shapes perception. PLoS Biol. *12*, e1002031.
- Neuling, T., Ruhnau, P., Fusca, M., Demarchi, G., Herrmann, C.S., and Weisz, N. (2015). Friends, not foes: Magnetoencephalography as a tool to uncover brain dynamics during transcranial alternating current stimulation. Neuroimage *118*, 406–413.