

Social Influence and the Brain: Persuasion, Susceptibility to Influence and
Retransmission

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Abstract

Social influence is an important topic of research, with a particularly long history in the social sciences. Recently, social influence has also become a topic of interest among neuroscientists. The aim of this review is to highlight current research that has examined neural systems associated with social influence, from the perspective of being influenced as well as influencing others, and highlight studies that link neural mechanisms with real-world behavior change beyond the laboratory. Although many of the studies reviewed focus on localizing brain regions implicated in influence within the lab, we argue that approaches that account for networks of brain regions and that integrate neural data with data beyond the laboratory are likely to be most fruitful in understanding influence.

Introduction

Social influence is omnipresent, occurring through implicit observation of cultural norms, face-to-face and mediated interpersonal communication, as well as mass mediated communication. Even though individuals are often unaware of the power of social influence, research shows its effects on behavior in a wide variety of circumstances [1]. The mechanisms driving social influence thus remain of high interest in diverse fields including psychology, sociology, communications, health, political science, marketing, and economics.

Recently, neuroscientists have begun to contribute to our understanding of social influence, especially with respect to underlying mechanisms that are not necessarily accessible with traditional self-report methodologies (Figure 1; for reviews see: [2–4]). For example, neuroimaging enables examination of mental processes in real time and reduces the need to rely exclusively on participant introspection [5]. This review highlights recent advances in neuroscience research on social influence, examining the core processes believed to be associated with susceptibility to influence, as well as successfully influencing others. To connect the study of influence with the broader social and cognitive neuroscience literature, we summarize evidence for overlap between neural systems implicated in conflict detection, positive valuation, social cognition, and self-related processing in the context of social influence. We conclude with a discussion of new insights and methods within social and cognitive neuroscience and computational social science disciplines that promise to advance our understanding of influence moving forward.

Susceptibility to Social Influence

Building on a long history of social sciences studying compliance and conformity (for a review, see [1]), a growing body of research has documented neural correlates of attitude and

behavior change in response to social norms or peer pressure. Converging evidence emphasizes overlap with brain systems associated with conflict detection and valuation in susceptibility to social influence [4].

Conflict Detection and Distress of Misalignment with the Group

Social psychologists have suggested that one core function of compliance and conformity is to maintain group harmony [1]. This account suggests that attitude and behavior change in response to social influence require the ability, whether conscious or unconscious, to detect conflicts between one's current behavior, preference or choice and those of others. The perception of being misaligned with others may elicit distress [6,7], which can motivate behavioral and attitudinal adjustments to realign with the group [8]. In this context, conformity may be enacted to gain group acceptance or support, which are also key to survival in evolutionary contexts [9].

The dorsal anterior cingulate cortex (dACC) is one key brain region implicated in conflict monitoring and detection [10–17], and early studies of influence demonstrated that updating behavior in response to misalignment with the group is associated with increased activity within this region [6,18], as well as in anterior insula (AI), a region hypothesized to encode the discomfort of being misaligned with the group [6,7]. To further test the causal role of brain regions hypothesized to be involved in conflict monitoring and detection in social influence, researchers used transcranial magnetic stimulation (TMS) to downregulate the posterior medial frontal cortex (pmFC), overlapping with dACC, during a social influence task. This manipulation reduced conformity to social influence, possibly by interrupting key processes relevant to reinforcement learning, and hence social conformity [19].

Extending to behaviors beyond the neuroimaging lab, individual differences in reactivity to social exclusion within dACC, AI, and subgenual cingulate predicted susceptibility to risky social influence in teens in a driving context one week after data were collected within these hypothesized regions using fMRI [20]. Taken together, these studies are consistent with the idea that sensitivity to social conflict and distress in form of anticipated or actual ‘social pain’ may contribute to conformity, such that individuals may conform to avoid negative social consequences and promote social bonding [3,8].

Valuation

In addition to conflict detection, social influence may derive power from positive value placed on social relationships [21]. Expected or experienced reward of social belonging or approval from others is thought to motivate conformity [22]. The ventral striatum (VS) and ventromedial prefrontal cortex (VMPFC) are known to respond to a wide variety of rewarding stimuli, including primary and secondary rewards [23]; VMPFC is known to convert various types of value (e.g., monetary and social) into a common scale which allows individuals to anticipate overall benefits of a stimulus based on diverse types of information (e.g., [24]). In studies of social conformity, neural activity within VS and VMPFC have been implicated in updating preferences to be in line with group opinions [25–27], which may reflect anticipated social rewards of group alignment. Some authors have also interpreted this to suggest that participants internalize what is valued by peers and come to value attitude objects rated positively by others more highly.

One study that directly tested neural differences between public and private conformity, however, found that brain regions hypothesized to be involved in conflict detection (dACC) during compliance decisions were associated with public compliance, while amygdala and

hippocampus activity was associated with private opinion changes [28]. Additional research is needed to convincingly demonstrate whether neural activity in each of the brain systems reviewed above directly produce private acceptance of norms, or whether this activity reflects anticipated or actual reward (or distress) derived from alignment (or misalignment) with the group.

Moderators of neural conformity effects

The brain systems reviewed above do not work in isolation and neural activity during social influence can also be moderated according to social context. For example, research has demonstrated that neural underpinnings of social influence are modulated by message source variables (such as communicator celebrity and expertise) [29] and in-group versus out-group status [30]. Furthermore, research examining peer influence and risk behaviors among adolescents suggests that developmental factors modulate neural processes key to influence; for example, the mere presence of another peer is associated with increased activity in hypothesized reward regions (VS, orbitofrontal cortex) during the decision-making process in adolescents (compared to adults), which in turn is associated with increased risk-taking [31]. Likewise, social norms expressed by adolescent peer confederates (risky versus cautious) interact with individual differences in neural regions associated with response inhibition (including the right inferior frontal gyrus and basal ganglia) to predict later risk-taking behavior in adolescents, suggesting that neural resources may be used differently in different social contexts [32].

Together, these studies demonstrate the power of social variables (e.g., group closeness, peer presence, and expertise) to influence the relationship between neural processing and social influence outcomes. These results also highlight the importance of longitudinal research to capture changes that occur within individuals over development [33]. More broadly, these

studies also highlight a wider range of neural systems that interact depending on contextual variables to determine influence and highlight the complexity of the influence process. Given this complexity, we argue for the potential value in examining networks of brain regions when studying influence. Such approaches will allow development of more comprehensive, integrative models of influence in the brain (see Future Directions).

Predicting Behavior Change

Many of the studies reviewed above focus on proximal outcomes that can be measured in a neuroimaging context (e.g., preference shifts). Some of the studies reviewed above, however, allude to a growing trend to explore not only proximal outcomes, but also the extent to which activity in key brain regions can predict longitudinal behavior outside the laboratory [34]. Consistent with the idea that multiple social and contextual signals are integrated to produce behavior change, initial work predicting behavior change from brain activity examined the role of the VMPFC in persuasive message processing. VMPFC is known to integrate multiple types of value signals [23] from limbic and prefrontal regions [35], which may serve as a summary value signal in response to social influence. Indeed, individual differences in VMPFC activity during persuasive message exposure successfully predicted participants' changes in sunscreen use one week after the scanning session compared to baseline usage beyond the participants' self-reported attitudes toward sunscreen and intentions to change their behavior [36]. In addition, research examining the effectiveness of smoking quit messages found that increased activity in the VMPFC during ad exposure predicted reductions in smoking one month following the scanning session compared to baseline beyond a number of self-report measures collected [37]. The authors of these behavior change studies suggest that the VMPFC may integrate information about the value placed on message content with respect to one's own goals and motivations.

Differences in participants' average responses to campaigns within VMPFC have also predicted success of the campaigns at the population level above and beyond standard self-report measures [38–40], suggesting that VMPFC in small groups of people may index the value to larger groups as well. These studies highlight VMPFC's potential key role in influence and demonstrate the utility of the *brain-as-predictor* approach [34] by showing that neural data explains variance in real-world behavior above and beyond self-report measures and highlights specific psychological pathways to change (e.g., [40,41]).

Influencing Others

In addition to studying those being influenced by social information, neuroscientists have started to consider the perspective of the influencer. Although this line of inquiry is still in its infancy, existing studies highlight the importance of increased temporoparietal junction (TPJ) activity in communicators who effectively influence others [37,42,43]. The TPJ is commonly associated with considering the intentions and perspectives of others, called mentalizing [44,45].

For example, research has examined neural correlates associated with a salesperson's ability to effectively take the perspective of their customers as indicated by a "salesperson theory of mind scale". Specifically, increased activity in bilateral TPJ and the medial prefrontal cortex was correlated with an increased self-reported likelihood to mentalize about consumers' cognitive states, which in turn was associated with greater sales performance [43]. Similarly, neural activity in the right TPJ during an fMRI recommendation task was associated with greater success in convincing others of the value of one's own opinions during a retransmission task after the scan [37]. The authors suggest that those who were more successful in propagating their own preferences may have engaged in mentalizing (e.g., considering how to make relevant information useful for others) during initial idea encoding inside the scanner [37].

Finally, research has examined the intersection of social influence and making recommendations for others using an fMRI task in which participants make recommendations to others while being exposed to experimentally assigned feedback about the recommendations previously made by peers [42]. Consistent with conformity research reviewed above, participants displayed greater activity in VS and VMPFC when conforming to peer recommendations versus maintaining their initial recommendations. Furthermore, consistent with research reviewed above on successful retransmission of influence, individual differences in right TPJ was associated with using social feedback to update recommendations for others [42]. These findings highlight the intersection of brain systems implicated in social influence and successful sharing in contexts that are highly pervasive now, e.g., writing online reviews in the face of existing reviews [42].

Follow-up research has also begun to consider how the social environment might moderate the neural mechanisms implicated in social influence and sharing behavior, for example contextualizing neural data with tools from social network analysis (SNA). SNA tools examine the size, structure, and scope of participants' social networks. By quantifying patterns of social relationships, social network analysis can operationalize sociological concepts such as an individual's access to social capital, influence, support and brokerage [46], as well as individual differences in disposition [47]. One such social network characteristic that has been studied as a potential moderator of neural activity in the context of influencing others is *ego betweenness centrality*. Ego betweenness centrality is a measure of information brokerage capacity—the extent to which an individual connects otherwise unconnected individuals within their network, and hence is positioned to broker the spread of ideas and information [48]. Although those who are high and low in betweenness centrality both update recommendations for others in response to social feedback, the underlying neural processes differ [48]. Those higher in betweenness

centrality showed more mentalizing activity when making recommendations and updating them in response to peer feedback compared to those who are lower in ego betweenness centrality; the authors suggest that having access to more diverse points of view or more practice translating ideas between different groups may encourage use of the brain's mentalizing system in day to day life, and/or that those who tend to engage in more mentalizing may position themselves in greater brokerage roles [48]. This work highlights potential value in integrating new tools from computational social science (e.g., social network analysis) to study how the brain responds to influence (for reviews, see [49,50]).

Conclusion and Future Directions

Neuroimaging provides a unique view of the underlying mechanisms that contribute to attitude and behavior change in response to social influence that are difficult to access using traditional methodologies [5]. The current review highlights early studies demonstrating relations between neural systems associated with valuation, conflict detection and social influence, between neural systems implicated in integrating value signals with respect to one's own motivations and behavior change, and between neural systems associated with mentalizing and successful influence over others (Figure 2). As reviewed above, however, social context variables modulate both neural and behavioral responses to influence and it is clear that brain systems work together in complex ways that go beyond the foundational brain-mapping research in this area. Neuroimaging research must now examine more complex neural network patterns within and between key systems involved in influence. Some (of many possible) theoretical and methodological means to this end are suggested below.

First, data analysis approaches that move beyond traditional mean activation estimates will offer new perspectives on social influence, for instance, by examining neural networks

rather than individual regions [51,52]. Specific examples of this would include using techniques derived from graph theory [53–56], connectivity analysis [57], or cognitive architectures [58,59]. It is almost certain that social influence processing is not localized to specific brain areas and the incorporation of network methods may reveal new knowledge about complex interconnections between neural regions during social influence and their interactions with context and development [60,61]. Thereby, knowledge gained from the studies reviewed above can suggest key nodes to consider in neural network analysis.

Second, techniques such as TMS and tDCS can provide stronger evidence for causal relationships (i.e., regions or network nodes that are not only involved but necessary for influence to occur) [19,62]. In addition, taking advantage of alternative neuroimaging tools such as functional near-infrared spectroscopy can allow researchers to capture neural mechanisms of more natural, live social interactions, allowing for greater external validity of findings [63]. Methods such as inter-subject correlation analysis [64,65] can also aid in moving toward greater external validity of findings by allowing examination of influence in response to naturalistic media [66,67].

Finally, as the neuroscience of social influence remains a relatively new area of inquiry, researchers should continue to develop tasks suitable for neuroimaging environments that are optimized for methods that offer high degrees of promise (e.g., network connectivity analyses, multivariate pattern classification approaches), and that are optimized to characterize how influence is modulated across different populations and across development [33]. More broadly, the findings reviewed above and those to come will offer new insights into social influence processes and using this information in conjunction with findings from other methodologies

(e.g., self-report, analytic methods from computational social science) can help us develop a more holistic understanding of social influence.

References

1. Cialdini RB, Goldstein NJ: **Social influence: Compliance and conformity.** *Annu Rev Psychol* 2004, **55**:591–621.
2. Cascio CN, Dal Cin S, Falk EB: **Health Communications: Predicting Behavior Change from the Brain.** In *Social Neuroscience and Public Health*. Edited by Hall PA. Springer New York; 2013:57–71.
3. Falk EB, Way BM, Jasinska AJ: **An imaging genetics approach to understanding social influence.** *Front. Hum. Neurosci.* 2012, **6**:168.
4. Izuma K: **The neural basis of social influence and attitude change.** *Curr. Opin. Neurobiol.* 2013, **23**:456–462.
5. Lieberman MD: **Social cognitive neuroscience.** In *Handbook of Social Psychology*. Edited by Fiske ST, Gilbert DT, Lindzey G. McGraw-Hill; 2010:143–193.
6. Berns GS, Capra CM, Moore S, Noussair C: **Neural mechanisms of the influence of popularity on adolescent ratings of music.** *Neuroimage* 2010, **49**:2687.
7. Tomlin D, Nedic A, Prentice DA, Holmes P, Cohen JD: **The neural substrates of social influence on decision making.** *PloS One* 2013, **8**:e52630.
8. DeWall CN: **Forming a basis for acceptance: excluded people form attitudes to agree with potential affiliates.** *Soc. Infl.* 2010, **5**:245–260.
9. Lieberman MD, Eisenberger NI: **Pains and Pleasures of Social Life.** *Science* 2009, **323**:890.
10. Cacioppo JT, Hawkley LC, Crawford LE, Ernst JM, Burleson MH, Kowalewski RB, Malarkey WB, Van Cauter E, Berntson GG: **Loneliness and health: potential mechanisms.** *Psychosom. Med.* 2002, **64**:407.
11. Carter CS, Braver TS, Barch DM, Botvinick MM, Noll D, Cohen JD: **Anterior cingulate cortex, error detection, and the online monitoring of performance.** *Science* 1998, **280**:747–749.
12. Critchley HD, Tang J, Glaser D, Butterworth B, Dolan RJ: **Anterior cingulate activity during error and autonomic response.** *Neuroimage* 2005, **27**:885–895.
13. Eisenberger NI: **The neural bases of social pain: evidence for shared representations with physical pain.** *Psychosom. Med.* 2012, **74**:126.
14. Hawkley LC, Burleson MH, Berntson GG, Cacioppo JT: **Loneliness in Everyday Life.** *J. Pers. Soc. Psychol.* 2003, **85**:105.

15. Hawkley LC, Thisted RA, Masi CM, Cacioppo JT: **Loneliness Predicts Increased Blood Pressure.** *Psychol. Aging* 2010, **25**:132.
16. Kerns JG, Cohen JD, MacDonald AW, Cho RY, Stenger VA, Carter CS: **Anterior cingulate conflict monitoring and adjustments in control.** *Science* 2004, **303**:1023–1026.
17. Peters E, Riksen-Walraven JM, Cillessen AHN, de Weerth C: **Peer rejection and HPA activity in middle childhood: friendship makes a difference.** *Child Dev.* 2011, **82**:1906.
18. Klucharev V, Hytönen K, Rijpkema M, Smidts A, Fernández G: **Reinforcement Learning Signal Predicts Social Conformity.** *Neuron* 2009, **61**:140.
19. Klucharev V, Munneke M, Smidts A, Fernández G: **Downregulation of the posterior medial frontal cortex prevents social conformity.** *J. Neurosci.* 2011, **31**:11934.
20. Falk EB, Cascio CN, O'Donnell MB, Carp J, Tinney F, Bingham CR, Shope JT, Ouimet MC, Pradhan AK, Simons-Morton BG: **Neural Responses to Exclusion Predict Susceptibility to Social Influence.** *J. Adolesc. Health* 2014, **54**:S22eS31.
21. Baumeister RF, Leary MR: **The need to belong: Desire for interpersonal attachments as a fundamental human motivation.** *Psychol. Bull.* 1995, **117**:497–529.
22. Ruff CC, Fehr E: **The neurobiology of rewards and values in social decision making.** *Nat. Rev. Neurosci.* 2014, **15**:549–562.
23. Bartra O, McGuire JT, Kable JW: **The valuation system: A coordinate-based meta-analysis of {BOLD} fMRI experiments examining neural correlates of subjective value.** *NeuroImage* 2013, **76**:412 – 427.
24. Izuma K, Saito DN, Sadato N: **Processing of the incentive for social approval in the ventral striatum during charitable donation.** *J. Cogn. Neurosci.* 2010, **22**:621–631.
25. Campbell-Meiklejohn DK, Bach DR, Roepstorff A, Dolan RJ, Frith CD: **How the opinion of others affects our valuation of objects.** *Curr. Biol. CB* 2010, **20**:1165.
26. Mason MF, Dyer R, Norton MI: **Neural mechanisms of social influence.** *Organ. Behav. Hum. Decis. Process.* 2009, **110**:152.
27. Zaki J, Schirmer J, Mitchell JP: **Social influence modulates the neural computation of value.** *Psychol. Sci.* 2011, **22**:894.
28. Edelson M, Sharot T, Dolan RJ, Dudai Y: **Following the crowd: brain substrates of long-term memory conformity.** *science* 2011, **333**:108–111.
29. Klucharev V, Smidts A, Fernández G: **Brain mechanisms of persuasion: how “expert power” modulates memory and attitudes.** *Soc. Cogn. Affect. Neurosci.* 2008, **3**:353.

30. Stallen M, Smidts A, Sanfey AG: **Peer influence: neural mechanisms underlying in-group conformity.** *Front. Hum. Neurosci.* 2013, **7**.
31. Chein J, Albert D, O'Brien L, Uckert K, Steinberg L: **Peers increase adolescent risk taking by enhancing activity in the brain's reward circuitry.** *Dev. Sci.* 2011, **14**:F1–F10.
32. Cascio CN, Carp J, O'Donnell MB, Tinney FJ, Bingham CR, Shope JT, Ouimet MC, Pradhan AK, Simons-Morton BG, Falk EB: **Buffering Social Influence: Neural Correlates of Response Inhibition Predict Driving Safety in the Presence of a Peer.** *J. Cogn. Neurosci.* 2014.
33. Falk EB, Hyde LW, Mitchell C, Faul J, Gonzalez R, Heitzeg MM, Keating DP, Langa KM, Martz ME, Maslowsky J: **What is a representative brain? Neuroscience meets population science.** *Proc. Natl. Acad. Sci.* 2013, **110**:17615–17622.
34. Berkman ET, Falk EB: **Beyond Brain Mapping: Using Neural Measures to Predict Real-World Outcomes.** *Curr. Dir. Psychol. Sci.* 2013, **22**:45–50.
35. Hare, T. A., Malmaud, J., & Rangel, A: **Focusing Attention on the Health Aspects of Foods Changes Value Signals in vmPFC and Improves Dietary Choice.** *Journal of Neuroscience* 2011, **31**:11077–11087.
36. Falk EB, Berkman ET, Mann T, Harrison B, Lieberman MD: **Predicting persuasion-induced behavior change from the brain.** *J. Neurosci. Off. J. Soc. Neurosci.* 2010, **30**:8421.
37. Falk EB, Morelli SA, Welborn BL, Dambacher K, Lieberman MD: **Creating buzz: the neural correlates of effective message propagation.** *Psychol. Sci.* 2013, **24**:1234.
38. Cooper N, Tompson S, O'Donnell MB, Falk EB: **Brain activity in self- and value-related regions in response to online antismoking messages predicts behavior change.** *J. Media Psychol.* in press.
39. Falk EB, O'Donnell MB, Tompson S, Gonzalez R, Dal Cin S, Strecher V, An L: **Neural systems associated with self-related processing predict population success of health messages.** In *Mass Communication and Health*. 2014.
40. Falk EB, Berkman ET, Whalen D, Lieberman MD: **Neural activity during health messaging predicts reductions in smoking above and beyond self-report.** *Health Psychol. Off. J. Div. Health Psychol. Am. Psychol. Assoc.* 2011, **30**:177.
41. Falk, E. B., Berkman, E. T., & Lieberman, M. D: **From Neural Responses to Population Behavior: Neural Focus Group Predicts Population-Level Media Effects.** *Psychological Science* 2012, **23**:439–445.

42. Cascio, C. N., O'Donnell, M. B., Bayer, J. B., Tinney, F. J., & Falk, E. B: **Neural correlates of susceptibility to group opinions in online word-of-mouth recommendations.** *Journal of Marketing Research*, in press.
43. Dietvorst RC, Verbeke WJMI, Bagozzi RP, Yoon C, Smits M, van der Lugt A: **A salesforce-specific theory of mind scale: tests of its validity by multitrait-multimethod matrix, confirmatory factor analysis, structural equation models, and functional magnetic resonance imaging.** *J. Mark. Res.* 2009, **46**:6653–668.
44. Saxe R, Kanwisher N: **People thinking about thinking people. The role of the temporo-parietal junction in “theory of mind.”***NeuroImage* 2003, **19**:1835.
45. Saxe R, Powell LJ: **It's the Thought That Counts Specific Brain Regions for One Component of Theory of Mind.** *Psychol. Sci.* 2006, **17**:692–699.
46. Wasserman S: *Social network analysis: Methods and applications.* Cambridge university press; 1994.
47. Burt RS, Kilduff M, Tasselli S: **Social network analysis: foundations and frontiers on advantage.** *Annu. Rev. Psychol.* 2013, **64**:527–547.
48. O'Donnell MB, Bayer JB, Cascio CN, Falk EB: **Online social network structure modulates neural precursors of recommendations.** *Nat. Neurosci.* conditionally accepted.
49. O'Donnell MB, Falk EB: **Linking neuroimaging with functional linguistic analysis to understand processes of successful communication.** *Commun. Methods Meas.* in press.
50. O'Donnell MB, Falk EB: **Big data under the microscope: Using brains, networks and language to link individual and population level data.** *Ann. Am. Acad. Pol. Soc. Sci.* in press.
51. Bassett DS, Gazzaniga MS: **Understanding complexity in the human brain.** *Trends Cogn. Sci.* 2011, **15**:200–209.
52. Bassett DS, Wymbs NF, Rombach MP, Porter MA, Mucha PJ, Grafton ST: **Task-based core-periphery organization of human brain dynamics.** *PLoS Comput. Biol.* 2013, **9**:e1003171.
53. Bassett DS, Bullmore E: **Human brain networks in health and disease.** *Curr. Opin. Neurol.* 2009, **22**:340.
54. Bassett, D. S., & Bullmore, E. T: **Small-world brain networks.** *The Neuroscientist* 2006, **12**:512–523.

55. Bullmore E, Sporns O: **Complex brain networks: graph theoretical analysis of structural and functional systems.** *Nat. Rev. Neurosci.* 2009, **10**:186–198.
56. Robinson LF, Atlas LY, Wager TD: **Dynamic functional connectivity using state-based dynamic community structure: Method and application to opioid analgesia.** *NeuroImage* 2014, [no volume].
57. Friston KJ: **Functional and effective connectivity: a review.** *Brain Connect.* 2011, **1**:13–36.
58. Borst, J. P., Taatgen, N. A., van Rijn, H: **Validating Models of Complex, Real-Life Tasks Using fMRI.** In *In A. Johnson & R.W. Proctor (Eds.), Neuroergonomics: A Cognitive Neuroscience Approach to Human Factors and Ergonomics.* Palgrave MacMillan. in press.
59. Borst JP, Anderson JR: **Using the ACT-R Cognitive Architecture in combination with fMRI data.** *Introd. Model-Based Cogn. Neurosci. Springer N. Y.* 2014.
60. Barrett LF, Satpute AB: **Large-scale brain networks in affective and social neuroscience: towards an integrative functional architecture of the brain.** *Curr. Opin. Neurobiol.* 2013, **23**:361–372.
61. Lindquist KA, Wager TD, Kober H, Bliss-Moreau E, Barrett LF: **The brain basis of emotion: a meta-analytic review.** *Behav. Brain Sci.* 2012, **35**:121–143.
62. Nitsche MA, Paulus W: **Sustained excitability elevations induced by transcranial DC motor cortex stimulation in humans.** *Neurology* 2001, **57**:1899–1901.
63. Redcay E, Dodell-Feder D, Pearrow MJ, Mavros PL, Kleiner M, Gabrieli JDE, Saxe R: **Live face-to-face interaction during fMRI: A new tool for social cognitive neuroscience.** *NeuroImage* 2010, **50**:1639–1647.
64. Hasson, U., Ghazanfar, A. A., Galantucci, B., Garrod, S., & Keysers, C: **Brain-to-brain coupling: a mechanism for creating and sharing a social world.** *Trends in Cognitive Sciences* 2012, **16**:114–121.
65. Hasson, U., Malach, R., & Heeger, D. J: **Reliability of cortical activity during natural stimulation.** *Trends in Cognitive Sciences* 2010, **14**:40–48.
66. Dmochowski JP, Bezdek MA, Abelson BP, Johnson JS, Schumacher EH, Parra LC: **Audience preferences are predicted by temporal reliability of neural processing.** *Nat. Commun.* 2014, **5**.
67. Schmälzle, R., Häcker, F., Renner, B., Honey, C. J., & Schupp, H. T: **Neural Correlates of Risk Perception during Real-Life Risk Communication.** *The Journal of Neuroscience* 2013, **33**:10340–10347.

Annotated Bibliography

19. Falk EB, Cascio CN, O'Donnell MB, Carp J, Tinney F, Bingham CR, Shope JT, Ouimet MC, Pradhan AK, Simons-Morton BG: **Neural Responses to Exclusion Predict Susceptibility to Social Influence.** *J. Adolesc. Health* 2014, **54**:S22eS31. *

Demonstrates that susceptibility to social influence is determined by individual differences in brain regions implicated in sensitivity to social cues (hypothesized 'social pain' and 'mentalizing' networks). Those who showed more activity in dACC, AI and subACC, as well as rTPJ, PCC and DMPFC during social exclusion were more likely to engage in risky driving in the presence of a peer (compared to driving alone) in an independent driving simulator session one week later.

29. Stallen M, Smidts A, Sanfey AG: **Peer influence: neural mechanisms underlying in-group conformity.** *Front. Hum. Neurosci.* 2013, **7**. *

Highlights the sensitivity of social influence effects to specific social contexts. Conformity with an in-group was more related to increased striatal activity, among others, than conformity to an out-group, suggesting the importance of social context in understanding the neural processes involved in influence.

33. Berkman ET, Falk EB: **Beyond Brain Mapping: Using Neural Measures to Predict Real-World Outcomes.** *Curr. Dir. Psychol. Sci.* 2013, **22**:45–50. **

This review provides an overview of the brain-as-predictor approach, i.e. the idea that neural data collected in the laboratory can be used to predict long-term, ecologically valid outcomes in the real world. This is relevant for social influence given that unobtrusive neuroimaging can help to circumvent the limitations of introspection and predict influence outcomes that are difficult to predict using other measures.

36. Falk EB, Morelli SA, Welborn BL, Dambacher K, Lieberman MD: **Creating buzz: the neural correlates of effective message propagation.** *Psychol. Sci.* 2013, **24**:1234. *

This is an early examination of social influence from the perspective of the propagator. The authors show that higher activity in hypothesized mentalizing regions (bilateral TPJ) during first exposure to new ideas, predicts a person's success in propagating their evaluation of these ideas to others. Considerations of others' likes, wants, and needs may put individuals in a better position to convince others.

41. Cascio, C. N., O'Donnell, M. B., Bayer, J. B., Tinney, F. J., & Falk, E. B: **Neural correlates of susceptibility to group opinions in online word-of-mouth recommendations.** *Journal of Marketing Research*, in press. *

One of the first studies on the neural correlates of the dynamic interplay between influencing others and being influenced at the same time. Neural correlates similar to those found when both processes are studied in isolation, namely valuation (OFC, VS) and mentalizing (TPJ) were found to play a role in updating recommendations in response to peer recommendations.

47. O'Donnell MB, Falk EB: **Big data under the microscope: Using brains, networks and language to link individual and population level data.** *Ann. Am. Acad. Pol. Soc. Sci.* in press. **

Advocates a multi-method approach to the study of social influence, combining large-scale data from computational social science such as information about participants' social networks with individual-level neural data. The authors argue that computational social science can help to strengthen and contextualize neural approaches and offer two accessible examples of this approach. We believe that such methods will be highly valuable in future social influence research, as they allow us to make diverse links between detailed, mechanistic neural data and participant's social reality outside the lab.

49. Bassett DS, Wymbs NF, Rombach MP, Porter MA, Mucha PJ, Grafton ST: **Task-based core-periphery organization of human brain dynamics.** *PLoS Comput. Biol.* 2013, **9**:e1003171. **

Demonstrates how network approaches to the analysis of neural data can be leveraged to gain a more detailed, mechanistic understanding of cognitive processes that we advocate be applied to the study of social influence. The authors explore brain systems that enable robust learning of motor skills and are able to identify changes in connectivity patterns in two time-evolving neural communities in which changes over time trace participants' learning progress.

56. Barrett LF, Satpute AB: **Large-scale brain networks in affective and social neuroscience: towards an integrative functional architecture of the brain.** *Curr. Opin. Neurobiol.* 2013, **23**:361–372. **

Argues for social neuroscience research on the level of large-scale, domain-general neural networks rather than domain-specific modules. As outlined above, we believe that this will be a crucial approach in future research on social influence, because complex social interactions likely rely on multiple, related mechanisms at once rather than on isolated, specialized brain regions.

62. Schmälzle, R., Häcker, F., Renner, B., Honey, C. J., & Schupp, H. T: **Neural Correlates of Risk Perception during Real-Life Risk Communication.** *The Journal of Neuroscience* 2013, **33**:10340–10347. *

Presents an application of inter-subject correlation for the study of responses to naturalistic and influential media. The authors use inter-subject correlation analyses on neural time courses to assess similarity of neural responses to a 30-minute TV report about the H1N1 pandemic. Inter-subject correlation was moderated by participant's topic-related risk perceptions so that those with high risk perceptions showed stronger within-group similarity in ACC activity than those with low risk perceptions.

Figures

Figure 1. Example heuristic model demonstrating the use of multiple methodologies to understand unique variance in behavior change in response to social influence manipulations (modified from; [2]).

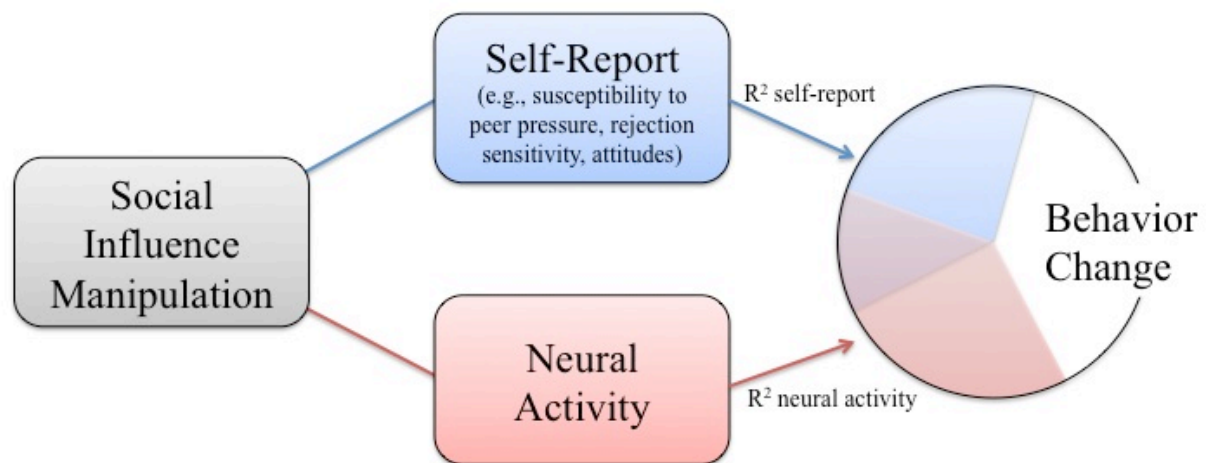


Figure 2. Hypothesized neural systems that may contribute to social influence. Valuation: VS = Ventral Striatum, VMPFC = ventral medial prefrontal cortex; Conflict detection and response: AI = anterior insula, dACC = dorsal anterior cingulate cortex; Self-related processing: MPFC = medial prefrontal cortex; and Mentalizing: DMPFC = dorsomedial prefrontal cortex, TPJ = temporal parietal junction

