

Reliability and subject specificity of personalized dynamical whole-brain models

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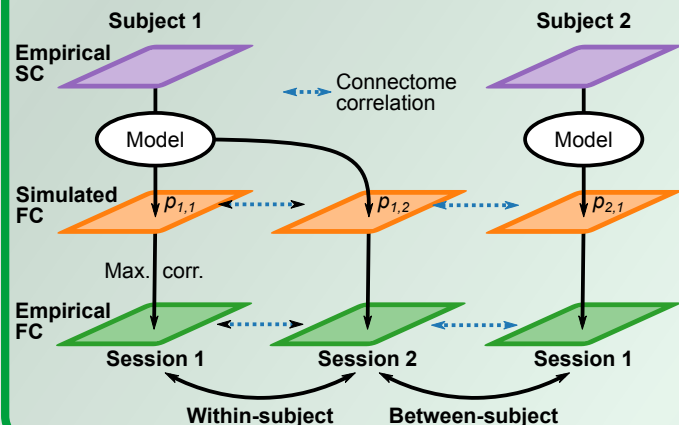


Main messages

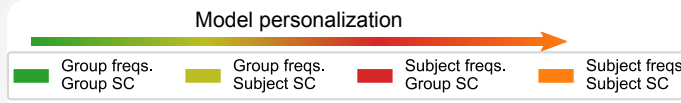
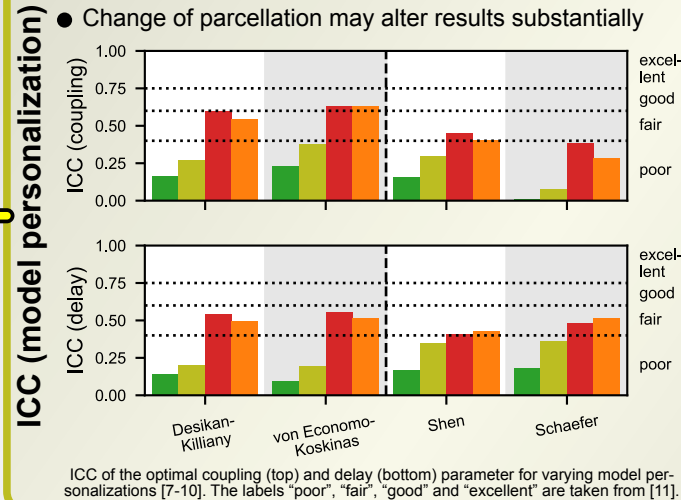
- The reliability of whole-brain dynamical models ranges from "poor" to "good"
- The reliability and subject specificity of modeling results may exceed those of empirical data
- Model personalization has a positive influence on the reliability and subject specificity
- Parcellations have a much larger effect on modeling results than on empirical data

Method

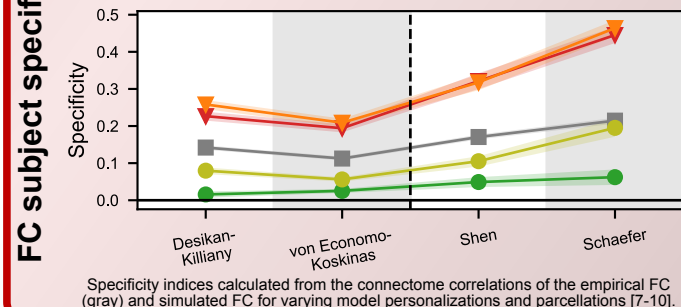
- How **reliably** and **specifically** are personalized dynamical whole-brain models fitted to the empirical data? [1,2]
- Empirical SC and FC (4 sessions) of 200 subjects in HCP [3,4]
- Model parameters optimized for every individual FC
- Reliability of model fit:** Intraclass correlation (ICC) of optimal model parameters and FC edges
- Reliability of FC:** within-subject connectome correlation
- Specificity** = within- — between-subject correlation
- Variation of model complexity:** networks of linear [5], phase oscillators and neural mass models [6]
- Variation of model personalization:** using varying amounts of subject-specific data for the phase oscillator



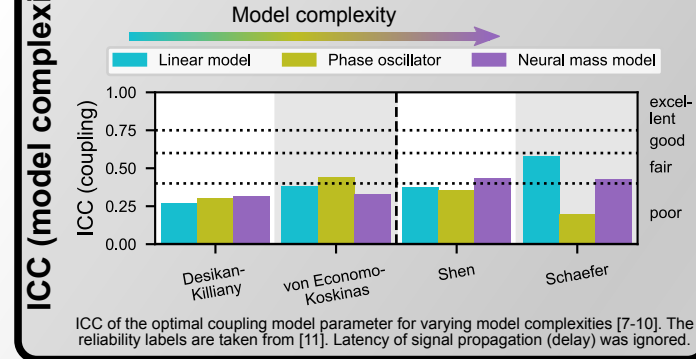
- Model parameters fitted with "poor" to "good" reliability
- Model personalization has positive influence on reliability
- Change of parcellation may alter results substantially



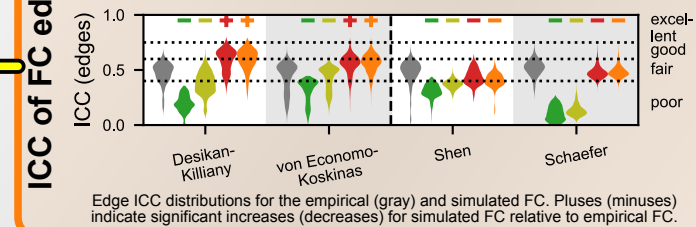
- Simulated FC can be more subject-specific than empirical FC
- Personalization positively affects subject specificity
- Effect of atlas larger for simulated than for empirical FC



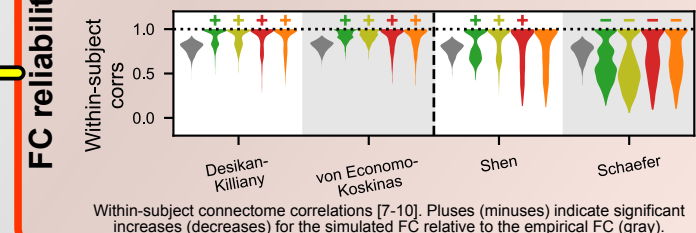
- Reduced variation across atlases for higher complexity
- Differences across complexities inconsistent across atlases



- ICC of edges may be higher for simulated than for empirical FC
- ICC of edges can be explained by ICC of parameters



- Reliability of simulated FC may exceed the empirical one
- Effect of atlas: potentially lower distributions with bimodalities



References (1) Honey et al. PNAS. 2009, 106(6), 2035-2040; (2) Bansal et al. Curr. Opin. Neurobiol. 2018, 52, 42-47; (3) Van Essen et al. NeuroImage. 2012, 62(4), 2222-2231; (4) Van Essen et al. NeuroImage. 2013, 80, 62-79; (5) Saggio et al. PLoS One. 2016, 11(8), e0157292; (6) Wilson et al. Biophys. J. 1972, 12(1), 1-24; (7) Desikan et al. NeuroImage. 2006, 31(3), 968-980; (8) Scholtens et al. NeuroImage. 2018, 170, 249-256; (9) Shen et al. NeuroImage. 2013, 82, 403-415; (10) Schaefer et al. Cereb. Cortex. 2018, 28(9), 3095-3114; (11) Cicchetti et al. Am. J. Ment. Defic. 1981, 86(2), 127-137

Acknowledgements

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