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## **Reporting Summary**

Nature Research wishes to improve the reproducibility of the work that we publish. This form provides structure for consistency and transparency in reporting. For further information on Nature Research policies, see Authors & Referees and the Editorial Policy Checklist.

Statistics		
For all statistical analyses, confirm that the following items are present in the figure legend, table legend, main text, or Methods section.		
n/a Confirmed		
The exact sample size (n) for each experimental group/condition, given as a discrete number and unit of measurement		
A statement on whether measurements were taken from distinct samples or whether the same sample was measured repeatedly		
The statistical test(s) used AND whether they are one- or two-sided Only common tests should be described solely by name; describe more complex techniques in the Methods section.		
A description of all covariates tested		
A description of any assumptions or corrections, such as tests of normality and adjustment for multiple comparisons		
A full description of the statistical parameters including central tendency (e.g. means) or other basic estimates (e.g. regression coefficient) AND variation (e.g. standard deviation) or associated estimates of uncertainty (e.g. confidence intervals)		
For null hypothesis testing, the test statistic (e.g. <i>F</i> , <i>t</i> , <i>r</i> ) with confidence intervals, effect sizes, degrees of freedom and <i>P</i> value noted Give <i>P</i> values as exact values whenever suitable.		
For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings		
For hierarchical and complex designs, identification of the appropriate level for tests and full reporting of outcomes		
Estimates of effect sizes (e.g. Cohen's $d$ , Pearson's $r$ ), indicating how they were calculated		
Our web collection on <u>statistics for biologists</u> contains articles on many of the points above.		
Software and code		
Policy information about <u>availability of computer code</u>		
Data collection Brainproducts (Brain Vision recorder), 4D Neuroimaging		
Data analysis MATLAB R2015B		
For manuscripts utilizing custom algorithms or software that are central to the research but not yet described in published literature, software must be made available to editors/reviewers.		

## Data

Policy information about availability of data

All manuscripts must include a data availability statement. This statement should provide the following information, where applicable:

- Accession codes, unique identifiers, or web links for publicly available datasets
- A list of figures that have associated raw data
- A description of any restrictions on data availability

Code availability.

The code used to generate the results that are reported in this study is available from the corresponding authors upon reasonable request

All data supporting the findings of this study are available from the corresponding authors upon reasonable request.

Field-spe	cific reporting
	ne below that is the best fit for your research. If you are not sure, read the appropriate sections before making your selection.
\times Life sciences	Behavioural & social sciences
	the document with all sections, see <u>nature.com/documents/nr-reporting-summary-flat.pdf</u>
Life scier	ices study design
All studies must dis	close on these points even when the disclosure is negative.
Sample size	The only criterion for patient selection was the presence of at least one electrode located in the hippocampus head or in the amygdala showing interictal epileptic activity ("interictal spikes") during underwent a simultaneous SEEG MEG recording study. Fourteen patients were studied (8 female).
Data exclusions	No electrode located in the hippocampus head or in the amygdala showing interictal epileptic activity
Replication	Replicability was tested across patients in these very rare datasets
Randomization	not applicable
Blinding	This is a retrospective study on simultaneous MEG-SEEG recordings; blinding is not relevant
materials & expenses expenses expenses and a linvolved in the linvolved in	ChIP-seq  cell lines  MRI-based neuroimaging  d other organisms  earch participants  a
	arch participants
Policy information  Population chara	about studies involving human research participants  cteristics Fourteen patients affected by pharmacoresistant epilepsy (8 female). Patients' mean age at recordings was 31.5 years, mean
Population chara	age at epilepsy onset was 18.2 years and mean epilepsy duration was 15 years.
Recruitment	Patients who underwent a simultaneous intracerebral stereotaxic EEG (SEEG) - MEG recordings in the setting of pre-surgical evaluation of focal drug-resistant epilepsies at the Clinical Neurophysiology and Epileptology Unit, APHM, Marseille, France.
Ethics oversight	Comité de Protection des Personnes, Sud-Méditerranée I, ID-RCB 2012-A00644-39
Note that full informa	tion on the approval of the study protocol must also be provided in the manuscript.
Clinical data	
•	about <u>clinical studies</u> d comply with the ICMJE <u>guidelines for publication of clinical research</u> and a completed <u>CONSORT checklist</u> must be included with all submissions.
Clinical trial regis	tration NCT02875964
Study protocol	https://clinicaltrials.gov/ct2/show/NCT02875964

Data collection

october2012-present

Primary objective: quantifying detectability of epileptic discharges; secondary objective: optimise signal processing methods