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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 our Editorial Policies and the Editorial Policy Checklist.

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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.				
X	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)				
x	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 <i>Give P values as exact values whenever suitable.</i>				
×	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				
x	Estimates of effect sizes (e.g. Cohen's <i>d</i> , Pearson's <i>r</i>), indicating how they were calculated				
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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

No software was used for data collection, as no new experimental or clinical data were collected.

Data analysis

All figures were produced using an agent-based modeling platform (Java) known as HAL: Hybrid Automata Library. Ductal branches were counted and quantified using opency-python (version 4.3.0) and scikit-image (version 0.17.0) for the Python programming language. The code for the model described in this paper is freely available open-source at: https://github.com/MathOnco/tissue-structure-modulates-evolution.

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 and reviewers. We strongly encourage code deposition in a community repository (e.g. GitHub). See the Nature Research guidelines for submitting code & software for further information.

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

No new experimental or clinical data were collected. Previously published data repurposed for parameterizing the model may be found at: https://doi.org/10.1016/i.cell.2017.12.007.

Field-spe	ecific reporting				
Please select the o	ne below that is the best fit for yo	ur research. If you are not sure, read the appropriate sections before making your selection.			
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Life scier	nces study desig	gn			
All studies must dis	sclose on these points even when t	the disclosure is negative.			
Sample size	Previously published clonal evolution and diversity measurement data of six synchronous patients with ductal carcinoma in situ breast cancer which progressed to invasive ductal carcinoma was used to parameterize the math model. This study is reference 20.				
Data exclusions	Four monoclonal tumors (P2,P3,P7,P9) were excluded in our model analysis, because the focus of the manuscript is on the timing and prevalence of tumor heterogeneity.				
Replication	Given the stochastic nature of the mathematical model, multiple simulations were performed to quantify mean-field averages for each parameterization. No new experimental or clinical data was collected during this study.				
Randomization	No new experimental or clinical data was collected during this study, and therefore randomization was not necessary.				
Blinding	No new experimental or clinical data was collected during this study, and therefore blinding was not necessary.				
Reporting for specific materials, systems and methods					
		materials, experimental systems and methods used in many studies. Here, indicate whether each material, enot sure if a list item applies to your research, read the appropriate section before selecting a response.			
Materials & exp	perimental systems	Methods			
n/a Involved in th	ne study	n/a Involved in the study			
X Antibodies		ChIP-seq			
x Eukaryotic	cell lines	Flow cytometry			
✗ ☐ Palaeontol	ogy and archaeology	MRI-based neuroimaging			

Palaeontology and archaeology Animals and other organisms

Human research participants

Dual use research of concern

Clinical data

X