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## **Exogenous chromosomes reveal how sequence composition drives chromatin assembly, activity, folding and compartmentalization**

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## 1 Abstract

2 Genomic sequences co-evolve with DNA-associated proteins to ensure the multiscale folding of  
3 long DNA molecules into functional chromosomes. In eukaryotes, different molecular complexes  
4 organize the chromosome's hierarchical structure, ranging from nucleosomes and cohesin-  
5 mediated DNA loops to large scale chromatin compartments. To explore the relationships  
6 between the DNA sequence composition and the spontaneous loading and activity of these  
7 DNA-associated complexes in the absence of co-evolution, we characterized chromatin  
8 assembly and activity in yeast strains carrying exogenous bacterial chromosomes that diverged  
9 from eukaryotic sequences over 1.5 billion years ago. We show that nucleosome assembly,  
10 transcriptional activity, cohesin-mediated looping, and chromatin compartmentalization can  
11 occur in a bacterial chromosome with a largely divergent sequence integrated in a eukaryotic  
12 host, and that the chromatinization of bacterial chromosomes is highly correlated with their  
13 sequence composition. These results are a step forward in understanding how foreign  
14 sequences are interpreted by a host nuclear machinery during natural horizontal gene transfers,  
15 as well as in synthetic genomics projects.

## 16        **Introduction**

17        Genome sequence composition, broadly defined by its GC%, polynucleotide frequencies,  
18        DNA motifs and repeats, varies widely between species as well as within individual genomes<sup>1,2</sup>.  
19        In eukaryotes, the sequence composition is known to correlate with: 1) chromatin composition,  
20        which includes nucleosome formation and binding of structural and functional proteins to DNA <sup>3</sup>,  
21        2) chromatin activity, such as transcription and replication <sup>4</sup>, and 3) functional 3D organization of  
22        the genome into loops and compartments <sup>5,6</sup>. For instance in mammals GC-rich regions are  
23        enriched in actively transcribed sequences and in chromatin loops mediated by the structural  
24        maintenance of chromosomes (SMC) cohesin, and genomic loci with different sequence  
25        composition tend to coalesce in distinct compartments <sup>7</sup>. These relationships between sequence  
26        and chromatin composition activities and folding reflect their continuous coevolution over  
27        millions of years.

28        Disruptive variations in sequence composition can also emerge naturally during evolution,  
29        e.g. through transfer of genetic material across species by horizontal gene transfers or  
30        introgression, in viral infections <sup>8-11</sup>, or even artificially, e.g. by introducing chromosome-long  
31        DNA molecules into chassis microbial strains or cell lines <sup>12-14</sup>. Such transfer can lead to the  
32        long-term integration of foreign DNA whose sequence composition strongly diverges from their  
33        host's genome (e.g. the introgression of a 12% GC-divergent 1Mb sequence in *Lachancea*  
34        *kluyveri* <sup>15</sup>). Once integrated, these sequences are organized and processed by chromatin-  
35        associated proteins of the host genome, obeying new rules under which they have not  
36        coevolved. How a eukaryotic host can successfully package, fold and regulate the activity of  
37        chromosome-long exogenous bacterial DNA sequences, and the importance of the sequence  
38        composition, remain largely unknown. Addressing this question would improve our  
39        understanding of the evolutionary processes and sequence determinants involved in chromatin  
40        folding and activity.

41 Here we explored whether and how foreign bacterial full-length chromosomes acquire  
42 chromatin features when integrated in a eukaryotic nucleus, and the relationship between their  
43 sequence composition and their chromatinization. To this end, we investigated the behavior of  
44 two individual bacterial chromosomes with different sequence composition, artificially introduced  
45 into the *S. cerevisiae* genome<sup>16,17</sup>. We profiled nucleosome assembly, polymerase and cohesin  
46 binding and activities, as well as the 3D chromosome organization during the cell cycle. We  
47 show that bacterial chromosomes with different sequence composition present different profiles  
48 of chromatin composition and activities, eventually leading to the spontaneous formation of  
49 active / inactive chromosomal compartments, similar to the ones observed in complex  
50 multicellular organisms. Using neural networks (NNs), we further show that eukaryotic sequence  
51 determinants are sufficient to predict the chromatin composition and activity of exogenous  
52 chromosomes integrated in the yeast genome. Consequently, this study suggests that the fate  
53 of any DNA molecule introduced into a given cellular context, including its organization from  
54 nucleosome positioning up to 3D folding and transcriptional activity, is governed by rules that  
55 are both deterministic and predictable.

56

## 57 **Results**

### 58 **Adaptation of supernumerary bacteria chromosomes to yeast**

59 To investigate large sequences which have not evolved in a eukaryotic context, we  
60 exploited *S. cerevisiae* strains carrying an extra 17<sup>th</sup> circular chromosome made either from the  
61 *Mycoplasma mycoides* subsp. *mycoides* (referred to as “Mmyco”) or *Mycoplasma pneumoniae*  
62 (“Mpneumo”) genomes containing a yeast centromeric sequence and an autonomous  
63 replication sequence (ARS)<sup>18</sup> (**Methods; Table S1**). While the GC content of *S. cerevisiae* is  
64 38%, the GC content of the Mmyco and Mpneumo chromosomes 24% (GC-poor) and 40% (GC-  
65 neutral), respectively (**Fig. 1a**). Dinucleotide composition, estimated by the dinucleotide odds

66 ratio  $\rho^*(XY)$ <sup>19</sup>, also largely differs between yeast and bacterial chromosomes (**Extended Data**  
67 **Fig. 1a**).

68 We linearized these chromosomes, added yeast telomeres at the extremities (**Fig. 1a**;  
69 **Extended Data Fig. 1b**; **Methods**), and investigated their replication and cohesion using split-  
70 dot assay and marker frequency analysis (**Extended Data Fig. 1c,d** and **Supplementary**  
71 **Results**; **Methods**). Overall, these chromosomes do not impose a significant fitness cost on  
72 their eukaryotic host, with each strain displaying similar growth rates compared to their parental  
73 strains in selective media (**Extended Data Fig. 1e**). They also have an estimated segregation  
74 rate similar to a centromeric plasmid<sup>20</sup> (**Extended Data Fig. 1f**).

75

### 76 **Spontaneous chromatinization of bacterial chromosomes in a eukaryotic context**

77 We first assessed whether integrated Mmyco and Mpneumo chromosomes were able to  
78 chromatinize by performing MNase-seq and H3 and H2A ChIP-seq experiments (Methods; **Fig.**  
79 **1b, c**; **Extended Data Fig. 2a**). Strikingly, both bacterial chromosomes were able to form  
80 nucleosomes and local nucleosome-depleted regions (NDR; **Fig. 1b, c**; **Extended Data Fig.**  
81 **2b-d**). Nucleosomes over the Mpneumo chromosome have structural features comparable to  
82 yeast nucleosomes, with a linker DNA of ~14 bp and a nucleosomal repeat length (NRL) of 160  
83 bp (**Fig. 1d,e**). In contrast, nucleosomes in the Mmyco chromosome have a longer linker DNA  
84 and thus are more spaced than in yeast or Mpneumo chromosomes, with an NRL of 177 bp  
85 (**Fig. 1d,e**). Two sequence features may explain this: first, poly(dA) and poly(dT) tracks, known  
86 to promote NDR formation, are over-represented along the Mmyco genome. Secondly, the  
87 AA/TT 10-bp periodicity, which facilitates DNA bending around histone cores, is reduced  
88 (**Extended Data Fig. 2e,f**). The analysis of MNase-seq time course experiments also revealed  
89 that Mmyco nucleosomal fragments are more rapidly degraded by MNase than yeast or  
90 Mpneumo nucleosomal fragments (**Extended Data Fig. 2c,d**), reflecting either the “fragile”  
91 nature of AT-rich nucleosomes or the MNase bias for these sequences<sup>21,22</sup>.

92 We further profiled the chromatin composition of Mmyco and Mpneumo chromosomes by  
93 performing RNA polymerase II (PolII) and cohesin (Scc1) ChIP-seq experiments (**Fig. 1f,**  
94 **Extended Data Fig. 2g-j**). We found that the Scc1 and PolII binding profiles along the  
95 Mpneumo chromosome appear similar to wild-type (WT) yeast chromosomes, with discrete  
96 Scc1 peaks (**Fig. 1f, Extended Data Fig. 2h,i**) preferentially located in nucleosome-depleted  
97 and PolII-enriched regions (**Extended Data Fig. 2j**). In contrast, Scc1 and PolII binding profiles  
98 in the Mmyco strain show significant differences. On the one hand, Scc1 is strongly enriched  
99 over the whole Mmyco chromosome, but does not form the distinct peaks observed on  
100 endogenous chromosomes (**Fig. 1f, Extended Data Fig. 2h,i**). Interestingly, Scc1 levels also  
101 appear strongly reduced at centromeres of endogenous *S. cerevisiae* chromosomes (**Extended**  
102 **Data Fig. 2g**), suggesting that the Mmyco chromosome titrates cohesins enriched over yeast  
103 centromeres. On the other hand, the PolII profile is greatly reduced along the Mmyco  
104 chromosome, compared with yeast chromosomes.

105 These results show that large exogenous bacterial chromosomes placed in a eukaryotic  
106 context spontaneously adopt eukaryotic chromatin composition features: histones, Pol II and  
107 cohesins all bind bacterial DNA irrespectively of their sequence composition. However, striking  
108 differences in the chromatin profiles led us to define two chromatin types: (1) “Y” (yeast-like)  
109 chromatin landscape, found over Mpneumo (whose 40% GC content is close to the native *S.*  
110 *cerevisiae* GC content), and (2) “U” (for Unconventional) chromatin, found over Mmyco (with a  
111 low 24% GC content), featuring less packed nucleosomes, a reduced Pol II enrichment, and a  
112 broad binding of cohesins across the entire chromosome.

113 Finally, we find that a convolutional neural network trained on yeast chromosome  
114 sequences was able to predict nucleosome, Scc1 and PolII binding profiles along integrated  
115 bacterial chromosomes (**Supplementary Results and Extended Data Fig. 3; Methods**). This  
116 indicates that the complex, underlying eukaryotic sequence determinants are successfully

117 reused to generate these different chromatin landscapes along integrated bacterial  
118 chromosomes.

119

## 120 **Transcriptional activity of bacterial genomes in a yeast context**

121 We next explored Y and U chromatin activities by first performing total, stranded RNA-seq  
122 (**Fig. 2a,b; Method**). Consistent with PolIII ChIP-seq profiles (**Fig. 1f**), we find that the Y  
123 chromatin type on Mpneumo is transcribed to levels similar to those of endogenous yeast  
124 chromosomes (**Fig. 2a**). However, Mpneumo transcription tracks are significantly longer than  
125 yeast genes (4.9kb vs 3.4kb, p-value < 2e-4, two-sided Student's t-test) and do not  
126 systematically display clear boundaries (**Fig. 2b**). They also do not preferentially occur over  
127 bacterial gene bodies nor seem to initiate at bacterial promoters, also consistent with an  
128 absence of PolIII enrichment at these loci (**Extended Data Fig. 4a-c**).

129 In sharp contrast, the Mmyco U chromatin type is only sparsely and lowly transcribed (**Fig.**  
130 **2a,b**), again in good agreement with the reduced levels of PolIII deposition (**Fig. 1f**). This  
131 transcriptional inactivity is not associated with the recruitment of the yeast silencing complex  
132 SIR Sir2/3/4<sup>23</sup> (**Extended Data Fig. 4d**).

133 Intriguingly, the transcription tracks orientation along bacterial chromosomes follows, on  
134 average, the orientation of the bacterial genes annotated along these sequences, although the  
135 transcription machinery has diverged billions years ago (**Fig. 2c**, see the orientation of genes  
136 and stranded tracks in **Fig. 2a,b**). This correlates with the over-representation of A (G)  
137 compared to T (C) on the coding strand, both detected in yeast genes and bacterial genes  
138 (**Extended Data Fig. 4e**, see **Discussion**).

139

## 140 **Inactive U chromatin forms a compartment into the nuclear space in G1**



141 To assess the 3D organization of Y and U chromatin types, we performed capture of  
142 chromosome conformation (Hi-C) <sup>24</sup> experiments on the Mmyco or Mpneumo strains arrested  
143 either in G1 or G2/M (**Fig. 3a, b**).

144 In G1-arrested cells, yeast chromosomes exhibit the expected Rab1 configuration, with  
145 clustered centromeres and subtelomeric regions, resulting in a dotted pattern in Hi-C contact  
146 maps <sup>25,26</sup>. For both Mmyco and Mpneumo chromosomes, the added yeast centromeric  
147 sequence clusters with the native yeast centromeres. However, the two bacterial chromosomes  
148 exhibit very distinct structural characteristics (**Fig. 3a**). The Y-type Mpneumo chromosome  
149 behaves as an endogenous *S. cerevisiae* chromosome (**Fig. 3a**), with similar trans-contacts  
150 (**Extended Data Fig. 5a**) and long-range cis-contacts (**Fig. 3c**), and a slope of the averaged  
151 contact probability ( $p$ ) as a function of the genomic distance ( $s$ ) close to -1.5, a value  
152 corresponding to the typical polymer structure observed in simulations (**Fig. 3c**) <sup>27</sup>. In addition,  
153 DNA-FISH labeling of Mpneumo DNA reveals an extended and contorted structure within the  
154 intranuclear space, confirming the intermixing of the chromosome with the actively transcribed  
155 native chromosomes (**Fig. 3e, Extended Data Fig. 5c,d**).

156 In contrast, in the U-type Mmyco chromosome, (1) only few contacts were detected with the  
157 endogenous chromosomes (**Extended Data Fig. 5a**), and predominantly with the 32 yeast  
158 subtelomeric regions (**Fig. 3a**, dotted rectangles; **Extended Data Fig. 5b**); (2) the slope of the  
159  $p(s)$  curve also changes dramatically towards a value of -1 at shorter distances, corresponding  
160 to a polymer globule (**Fig. 3c**) <sup>24</sup>; (3) DNA-FISH labeling of the Mmyco chromosome confirmed  
161 the globular conformation of the chromosome and revealed its preferential position at the  
162 nuclear periphery, reflecting its proximity with sub/telomeres (**Fig. 3e; Extended Data Fig.**  
163 **5c,d**). These results show that inactive, U chromatin forms a compartment in the nuclear space  
164 segregated from the active endogenous chromosomal set.

165

166 **Cohesin compacts similarly both exogenous bacterial chromosomes during G2/M**

167 We then generated Hi-C contact maps of both strains synchronized in G2/M. In yeast,  
168 chromosome compaction in G2/M-arrested cells reflect cohesin-mediated chromatin folding into  
169 arrays of loops, with cohesin peaks corresponding to loop anchors<sup>28,29</sup>. This compaction results  
170 in a typical p(s) curve that presents an inflection point around the average loop length<sup>25</sup>. The  
171 decrease of interchromosomal contacts (**Fig. 3b, Extended Data Fig. 5a**) and of p(s) (**Fig. 3d,**  
172 **Extended Data Fig. 5e,f**) observed for both native and bacterial chromosomes indicate a  
173 similar mitotic compaction of both Y and U chromatin types.

174 Yeast and Mpneumo chromosomes exhibit focal contact Scc1 enrichments corresponding  
175 to cohesin-anchored loops (black triangles on **Fig. 3f**)<sup>28</sup>. We found that cohesin accumulates at  
176 sites of convergent transcription not only, as expected, on yeast chromosomes<sup>30,31</sup>, but also on  
177 Mpneumo (**Fig. 2b**, green dotted lines, **Extended Data Fig. 5g**), with strong cohesin peaks  
178 associated with sharper transcriptional convergence (**Fig. 2b**, arrows; **Extended Data Fig.**  
179 **5h,i**). Using Chromosight, we identified 59 loops across the Mpneumo chromosome, spanning  
180 over slightly longer genomic distances (28 kb on average) than the loops found along yeast  
181 sequences (22 kb) (**Extended Data Fig. 5j; Methods**), and with strong Scc1 peaks positioned  
182 close to anchors, with a median distance of 566 bp (**Fig. 3f, h**). The dotted grid pattern in  
183 Mpneumo is reminiscent of multiple DNA loops observed along mammalian interphase  
184 chromosomes<sup>32</sup>, suggesting that some loop anchors have the ability to be involved in loops of  
185 various sizes, a phenomenon not observed across native *S. cerevisiae* chromosomes (**Fig. 3f**).

186 On the other hand, no visible discrete loops were observed along the Mmyco chromosome  
187 where cohesins cover inactive U chromatin regions without clear peaks, probably reflecting the  
188 absence of (convergent) transcription (**Fig. 3g**) (**Fig. 1f** and **Fig. 2b**). These results suggest that  
189 cohesin can bind DNA and promote compaction in absence of active transcription, but does  
190 not accumulate at discrete positions.

191 Cohesin-mediated loops along yeast chromosomes are formed through loop extrusion (LE),  
192 a process by which cohesin complexes organize DNA by capturing and gradually enlarging

193 small loops<sup>33,34</sup>. To test whether LE compacts bacterial genomes, we performed Hi-C in  
194 absence of Wpl1 (human Wapl), which impairs cohesin removal and results in longer loops  
195 readily visible by a shift of the inflection point in the p(s) curve<sup>28,34</sup>. In absence of Wpl1, we  
196 observed increased long-range contacts in all chromosomes, including Mmyco and Mpneumo  
197 **(Fig. 3i; Extended Data Fig. 5k)**, consistent with active cohesin-mediated LE proceeding on  
198 both Y and U chromatin.

199 Overall, these observations highlight two different spatial configurations of Mmyco and  
200 Mpneumo exogenous chromosomes. On the one hand, the inactive Mmyco chromosome in G1  
201 is compacted and positioned at the periphery of the nucleus, away from the active chromosome  
202 set. Upon S-phase and G2 arrest, the DNA is nevertheless compacted in a cohesin-dependent  
203 manner with no visible loops at discrete positions. On the other hand, the active Mpneumo  
204 chromosome behaves as endogenous yeast chromosomes, displaying discrete mitotic DNA  
205 loops in G2/M.

206

### 207 **Mosaic chimeric chromosomes display spontaneous, chromatin type-dependent** 208 **DNA compartmentalization**

209 The structural and functional features of the Y- and U-type chromatin are reminiscent of the  
210 euchromatin and heterochromatin compartments described along metazoan chromosomes<sup>24</sup>.  
211 We wondered whether they can coexist on a single chromosome. Using CRISPR-Cas9, we first  
212 fused the Mmyco chromosome with the native yeast chromosome XVI (XVIfMmyco  
213 chromosome). We then induced translocations to generate two other strains with chromosomes  
214 harboring alternating U and Y chromatin regions as small as 50 kb (XVIfMmycot1 and  
215 XVIfMmycot2; **Fig. 4a; Extended Data Fig. 6a-c** Methods). ChIP-seq profiles of RNA PolIII on  
216 Mmyco regions were similar before and after translocation, demonstrating that PolIII binding is  
217 independent of the broader regional context (**Extended Data Fig. 6d**). The overall Hi-C contact  
218 maps of XVIfMmyco strain synchronized in G1 showed little differences compared to the

219 parental Mmyco strain (but for the deleted and fused regions) (**Fig. 4b**, compare panel **i** with **ii**).  
220 The Y-type chr. XVI arm intermixes with the other 15 yeast chromosomes while the U-type  
221 Mmyco arm remains isolated from the rest of the genome, contacting subtelomeric regions (**Fig.**  
222 **4b**). In the translocated strains, however, the alternating chromatin type regions resulted in  
223 striking checkerboard contact patterns within XVIIfMmycot1 and XVIIfMmycot2 chromosomes  
224 (**Fig. 4b**, panels **iii** and **iv**). This pattern reflects the fact that U-type regions make specific  
225 contacts over long distances, bypassing the Y-type regions found in between (**Fig. 4c**;  
226 **Extended Data Fig. 6e** G1). Similarly, Y-type regions of chromosome XVI also are also  
227 involved in specific Y-Y contacts over longer distances (**Fig. 4c**). Intra U and Y regions contact  
228 decay is also different in the two chromatin types, U chromatin being prone to longer range  
229 contacts in cis (**Extended Data Fig. 6e**). We then induced another translocation to integrate a  
230 segment of U chromatin in a *trans* position at the end of chromosome XIII and generated the  
231 corresponding Hi-C map (**Fig. 4d**). The correlation and 4C-like plot revealed an increase in  
232 inter-chromosomal contacts between Y chromatin segments with distant segments of itself, and  
233 in alternance with decreased contacts with Mmyco U chromatin segments (and vice versa),  
234 showing that the inactive U chromatin compartment can contain segments positioned on  
235 different chromosomes (**Fig. 4d**). These results unambiguously show that in yeast alternated or  
236 inter-chromosomal segments of active and inactive chromatin are spontaneously segregated  
237 within the nuclear space in G1 (**Fig. 4e**, **Extended Data Fig. 6h**), a phenomenon reminiscent of  
238 eu- and heterochromatin compartments found in higher eukaryotes <sup>24</sup>.

239 In G2/M, the Scc1 deposition pattern, including centromere depletion, remained conserved  
240 along mosaic chromosomes (**Extended Data Fig. 6d**). Cohesin coating remained enriched over  
241 U-type Mmyco sequences, including over the 50 kb encompassed within 850 Mb of yeast DNA  
242 on XVIIfMmycot2 chromosome. These results show that cohesin is loaded along inactive Mmyco  
243 regions independently of the proximity with the centromere (as along WT chromosomes) and

244 independently of transcription, suggesting that the larger chromatin context plays little role in  
245 cohesin-mediated loading. In addition, Hi-C maps of the mosaic chromosomes in G2/M show  
246 that U-type compartments disappear upon cohesin-mediated compaction (**Extended Data Fig.**  
247 **6f**). The analysis of distance-dependent contact frequency shows that all chromosome regions  
248 behave similarly at this stage (**Extended Data Fig. 6e**, G2/M). The only noticeable difference  
249 resulting from the fusion/translocation events in G2/M Hi-C contact maps consists in a DNA loop  
250 appearing over the boundary between Mmyco and the endogenous yeast sequence (**Extended**  
251 **Data Fig. 6f,g**). The two anchors of the loop are a strongly cohesin-enriched region already  
252 present on chr. XVI sequence, and another region slightly enriched in cohesin already present  
253 on the Mmyco sequence (**Extended Data Fig. 6g**, green arrows).

254 Together with observations from the previous section, these results show that Y and U  
255 chromatin types are both prone to dynamic cohesin-dependent DNA compaction and loop  
256 formation in G2/M phase (**Fig. 4e**) and further suggest that loop extrusion can proceed through  
257 the Y/U chromatin junction to form a loop bridging the nearest cohesin enrichment sites on each  
258 side <sup>35,36</sup>. Therefore, as observed in multicellular organisms, LE-mediated metaphase  
259 chromosome compaction abolishes chromatin compartments to prepare chromosomes for  
260 segregation (**Fig. 4e**) <sup>33,37</sup>.

261

## 262 Discussion

263 *Chromatin composition and activity of bacterial DNA follows yeast sequence determinants*

264 Several studies have shown that exogenous or random DNA segments in the yeast nucleus  
265 with a GC% relatively similar to that of yeast chromosome are actively transcribed <sup>38-41</sup>. Here we  
266 explore the relationships between sequence composition and chromatin composition and  
267 activity by characterizing large, divergent bacterial DNA sequences in a eukaryotic cellular  
268 context into which they have not evolved. We find that *M. pneumoniae* and *M. mycoides*

269 chromosomes integrated in yeast adopt one of two archetypes of chromatin that we called Y  
270 and U for yeast-like and unconventional. In both types, nucleosomes can assemble albeit with a  
271 longer NRL in the U chromatin (177 vs 160 bp), which also correlates with a low transcription. In  
272 G1, transcriptionally inactive U chromatin segments spontaneously collapse into a globule  
273 leading to the formation of a new nuclear compartment, whereas Y chromatin intermixes with  
274 yeast endogenous DNA. In G2/M, SMC cohesin complexes bind to both chromatin types and  
275 compact DNA through the extrusion of DNA loops<sup>6</sup>. Along Y chromatin, these complexes  
276 accumulate at convergent transcription loci<sup>28,30</sup>, whereas along U chromatin, the absence of  
277 active transcription leads to a homogeneous enrichment pattern.

278 The chromatin composition and activity experimentally measured over these exogenous  
279 chromosomes can be predicted using convolutional neural networks solely trained on yeast  
280 sequences, showing that exogenous DNA composition obeys a set of host rules based on  
281 sequence determinants that include GC content, dinucleotide composition but also complex  
282 sequence features. Sequence composition is therefore an important driver of the chromatin  
283 composition of exogenous sequences.

284

285 *Prokaryotic DNA contains some determinants of transcription orientation in eukaryote*

286 Active transcription on bacterial chromosomes in yeast follows, on average, bacteria genes  
287 orientation (**Fig. 2**). Several mechanisms could account for this observation. Sequence  
288 determinants predating the divergence of eucaryotes and procaryotes, including GC and AT  
289 skews, are known to influence polymerase directionality<sup>47,48</sup>. Alternatively, bacteria codon  
290 usage bias could result in enrichment in Nrd1-Nab3-Sen1 (NNS) binding sites along the ncRNA  
291 strand, leading to transcription termination<sup>39,49</sup>. The Nonsense-Mediated Decay (NMD)  
292 complex, which destabilizes in the cytoplasm eukaryotic transcripts with long 3' UTR, could also  
293 be more frequently activated by bacteria RNA transcribed in an antisense orientation.  
294 Eventually, the resulting conserved orientation could facilitate the domestication of exogenous

295 sequences during horizontal transfer/introgression events between distant species. Whether  
296 these RNA molecules are translated, and peptides of bacterial origins exist in the yeast cell,  
297 remains to be determined. If so, they could provide a source of diversity and adaptation.

298

### 299 *Cohesin-mediated chromatin folding along non-transcribed DNA templates*

300 It is proposed that cohesins expand DNA loops by an active process of loop extrusion until  
301 they encounter an obstacle and/or a release signal <sup>6,33</sup>. For instance, the transcriptional  
302 repressor CTCF constitutes a cohesin roadblock to LE in mammals, but SMC-mediated loops  
303 have been identified in genomes lacking CTCF across domains of life. In yeast and other  
304 species, anchors are thought to be determined by a combination of convergent transcription,  
305 replication fork progression during S phase and/or the presence at these positions of stably  
306 bound cohesin promoting sister chromatid cohesion <sup>50-52</sup>. The barely transcribed Mmyco  
307 chromosome is compacted by cohesin at G2/M, without discrete loop anchoring, suggesting that  
308 transcription is neither necessary for loading nor a primary driver for translocation. We propose  
309 that cohesins can move freely along this template without encountering significant obstacles,  
310 making it a suitable model for studying potentially blocking sequences or molecules.

311

### 312 *Compartmentalization of foreign DNA in host's nucleus*

313 We show that the introduction a foreign DNA with a lower GC content is sufficient to  
314 spontaneously promote the formation of a distinct inactive chromatin type, with increased NRL  
315 and a compact globular structure, all features reminiscent of mammalian heterochromatin. This  
316 chromatin forms a new compartment at the periphery of the budding yeast nucleus (**Fig. 5e;**  
317 **Extended Data Fig. 6h**), like the metazoan compartment "B" formed by inactive,  
318 H3K9me3/HP1-mediated heterochromatin, which is absent in yeast. The mechanisms behind  
319 compartmentalization remain unknown, but we hypothesize that the unmixing between U and Y  
320 chromatin could be due either to NRL change, protein composition and/or activity. Importantly,

321 longer NRLs are found in heterochromatin in mammals, and are associated with histone H1<sup>53</sup>.

322 In contrast, Y chromatin intermixes with active yeast DNA.

323 Depending on its sequence composition, a DNA molecule will therefore either  
324 compartmentalize or intermix with the host DNA. These different fates of the two chromatin  
325 types raise interesting evolutionary considerations. In the context of the invasion of the genome  
326 by exogenous mobile elements, this could lead either to the spontaneous isolation of inactive  
327 foreign DNA or, on the contrary, to the co-option of a new set of active sequences that could  
328 represent a reservoir of genetic innovations. This sequence-dependent mechanism may have  
329 contributed to the heterochromatinization of AT-rich transposable elements integrated in  
330 mammalian genomes.

331

### 332 **Competing interests**

333 The authors declare no competing interests.

334

### 335 **Authors Contributions**

336 Conceptualization: CC, LM, JS, JM and RK. Methodology: LM, CC, JS, CL, JM, RK. Software:  
337 JS. Validation: CC, JS, LM. Investigation: LM, CC, with contributions from AP, AB, AgT MR and  
338 FB. Formal analysis: JS, AW, ER, CC, LM, FB, MR, AnT. Data Curation: JS, with contributions  
339 from CC, LM. Resources: GG, CL. Visualization: JS. Writing - original draft preparation: CC, JS,  
340 LM, JM and RK. Writing – Editing: all authors. Writing – Revisions: JS, LM, AW, JM, RK.  
341 Supervision: JM, RK. CC co-supervised a student. Funding acquisition: RK. Project  
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355 sequencing RNA samples.

## 356 **Figure legends**

357

### 358 **Figure 1. Chromatin composition of bacterial chromosomes in yeast.**

359 **a**, Schematic representation of the conversion from circular to linear chromosomes integrated in  
360 yeast. The purple and blue colors represent the *M. pneumoniae* (Mpneumo) and *M. mycoides*  
361 (Mmyco) bacterial sequence in all figures, respectively. Right: distribution of GC% for 1kb  
362 windows over yeast chromosome XVI, *M. pneumoniae* and *M. mycoides* chromosomes.

363 **b**, Nucleosomal track (grey, see **Methods**) and H2A and H3 ChIP-seq (shades of blue, IP vs  
364 input, log<sub>2</sub>) profiles obtained in the Mpneumo strain (*S. cerevisiae* + *M. pneumoniae*). 10kb-long  
365 genomic windows from the chromosome XVI (left: 250kb-260kb) and the Mpneumo  
366 chromosome (right: 710kb-720kb) are shown at the same scale. Genomic loci highlighted in  
367 yellow correspond to nucleosome-depleted regions.

368 **c**, Same as **b** in the Mmyco strain (*S. cerevisiae* + *M. mycoides*). 10kb-long genomic windows  
369 from chromosome XVI (left: 250kb-260kb) and the Mmyco chromosome (right: 989kb-999kb)  
370 are shown at the same scale.

371 **d**, Frequency of nucleosome linker DNA length in Mpneumo and Mmyco strains. For each  
372 strain, the distribution is calculated for each chromosome separately. The dashed line indicates  
373 14 bp and the dotted line indicates 25 bp.

374 **e**, Auto-correlation of the nucleosome track in Mpneumo and Mmyco strains, computed for each  
375 chromosome separately (lag ≤ 1000). Nucleosome repeat length (NRL) is indicated for yeast,  
376 Mpneumo and Mmyco chromosomes.

377 **f**, Scc1 (red) and RNA Pol II (green) ChIP-seq profiles (IP vs input, log<sub>2</sub>) obtained in the  
378 Mpneumo (left) or in the Mmyco (right) strains. For each strain, 60kb-long genomic windows  
379 from the bacterial chromosome and the chromosome XVI are shown, at the same scale. GC%  
380 in sliding 1kb windows is shown below the ChIP-seq profiles.

381

382 **Figure 2. Expression of exogenic bacterial sequences in budding yeast.**

383 **a**, Stranded RNA-seq profiles along yeast chromosome XVI or along bacterial chromosomes  
384 Mpneumo and Mmyco in the corresponding strains. Forward (pink) and reverse (turquoise)  
385 genes along yeast or bacterial sequences are indicated as transparent segments under the  
386 tracks. Pink and turquoise represent forward and reverse transcription, respectively.

387 **b**, Top: stranded RNA-seq profiles along a 60kb window along yeast chromosome XVI,  
388 Mpneumo or Mmyco. Bottom: Scc1 (cohesin) deposition profiles of the corresponding loci.  
389 Green dotted lines indicate identified loci of convergent transcription (see Methods).

390 **c**, Forward and reverse RNA-seq coverage of forward- and reverse-oriented yeast or bacterial  
391 genes. Scores were normalized by the length of each genomic feature.

392

393 **Figure 3. Folding of exogenic bacterial sequences within the yeast nucleus.**

394 **a, b**, Hi-C contact maps of representative endogenous and of Mmyco and Mpneumo bacterial  
395 chromosomes in G1 (**a**) and G2/M (**b**) (4kb resolution).

396 **c, d**, Contact frequency ( $\rho$ ) as a function of genomic distance ( $s$ ) plots of endogenous yeast  
397 chromosomes (long arms) and of Mmyco and Mpneumo bacterial chromosomes in G1 (**c**) and  
398 G2/M (**d**).

399 **e**, Left: FISH imaging. Representative field of either (top) Mpneumo or (bottom) Mmyco fixed  
400 cells labeled with DAPI (left panel) and hybridized with a fluorescent probe generated from  
401 either the Mpneumo or Mmyco chromosome, respectively. Right: For each probe, number of  
402 patches detected per nucleus and surface occupied by these patches relative to the whole  
403 nucleus surface (Methods).

404 **f, g**, Top: magnification of 150kb windows from Hi-C contact maps in G2/M from either an  
405 endogenous or the bacterial chromosome in Mpneumo (**f**) and Mmyco (**g**) strains (1kb

406 resolution). Bottom: Scc1 ChIP-seq deposition profile. Black arrowheads: loops. Cyan  
407 diamonds: Scc1 peaks positions.

408 **h**, Left: distance between chromatin loop anchors and the nearest Scc1 peak in Mpneumo (with  
409 and without a random shuffle of peak positions). Right: Scc1 peak strengths in yeast or  
410 Mpneumo chromosome, near (< 1kb) or outside loop anchors (p-values from two-sided  
411 Student's t-test).

412 **i**, Distance-dependent contact frequency in endogenous yeast and in bacterial chromosomes  
413 (left, Mpneumo; right, Mmyco), in WT (dashed) and in  $\Delta wpl1$  mutants.

414

415 **Figure 4. Compartmentalization of mosaic chromosomes composed of Y and U-type**  
416 **chromatin.**

417 **a**, Schematic representation of the CRISPR strategy used to generate the Mmyco mosaic  
418 chromosomes with alternating Y and U chromatin regions.

419 **b**, Top: G1 Hi-C contact maps of chr. XV, XVI, and bacterial chromosomes in the Mmyco strain,  
420 and in translocated derivatives from panel **a**) (4kb resolution). Bottom: correlation matrices of  
421 the corresponding contact maps.

422 **c**, Virtual 4C profiles of viewpoints (grey/blue arrows) located within yeast chr. XVI (in grey) or  
423 Mmyco chromosome segments (in blue), in XVIfMmyco (top), XVIfMmycot1 (middle) and  
424 XVIfMmycot2 (bottom) strains.

425 **d**, Left: Schematic representation of the CRISPR strategy used to generate the Mmyco mosaic  
426 XVIfMmycot3 strain, with alternating yeast and Mmyco segments in chromosomes XIII and XVI.  
427 Middle: G1 Hi-C maps of mosaic chromosomes XIII and XVI in XVIfMmycot3 strain. The color  
428 scales for the normalized contact map and the correlation matrix are the same as in **b**. Right:  
429 Virtual 4C profiles of viewpoints located within yeast chr. XIII (in grey) or within yeast chr. XVI (in  
430 grey) or Mmyco chromosome (in blue) segments in XVIfMmycot3 strain. Grey/blue arrows:Y  
431 and U chromatin viewpoints position.

432 **e**, Schematic of a chimeric chromosome composed of alternating Y- and U-type chromatin. Left:  
433 composition of Y- and U-type chromatin. Middle: Spatial and functional organization in G1.  
434 Right: Organization of Y and U-type chromatin segments in G2/M.

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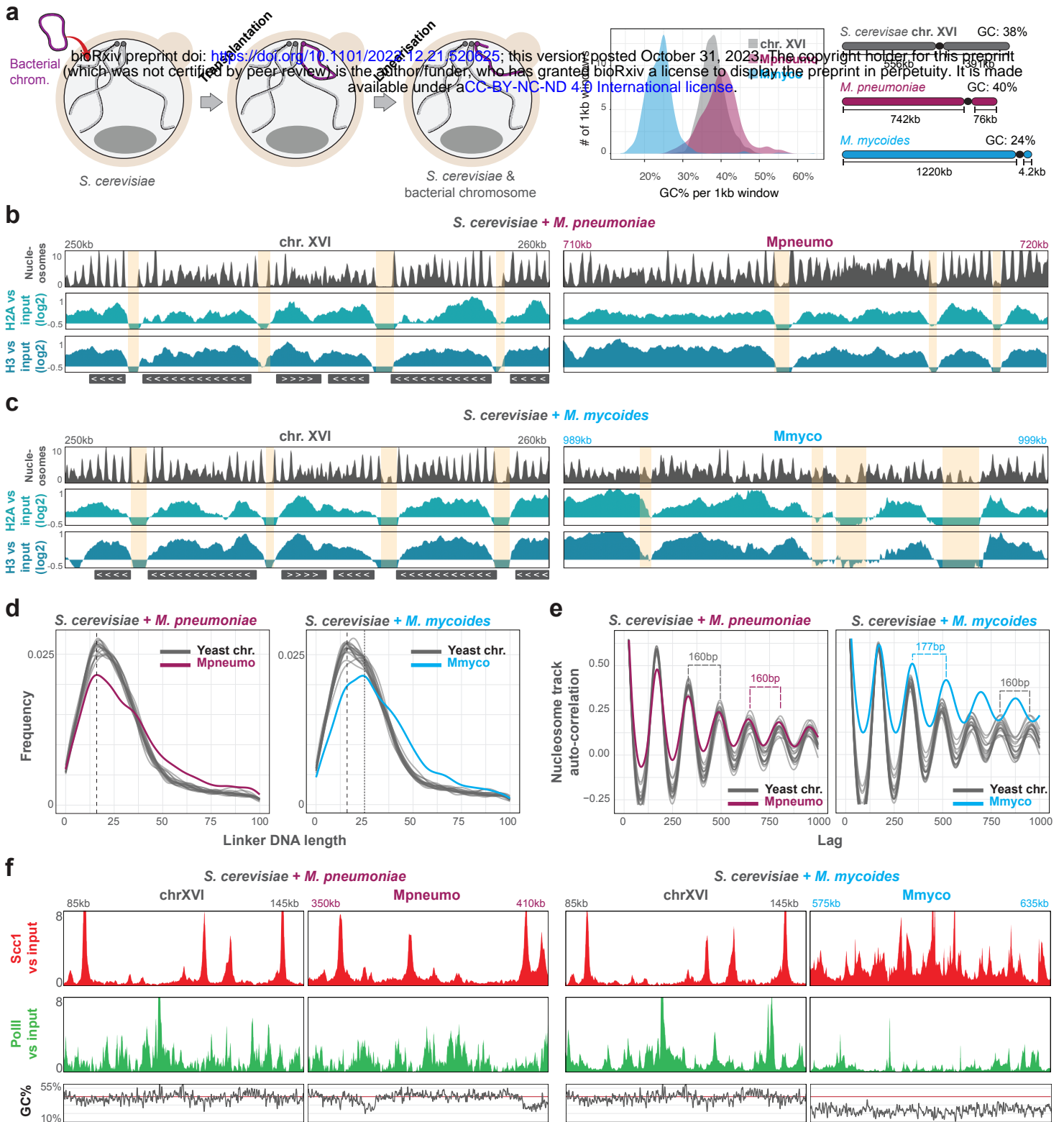


Fig. 1

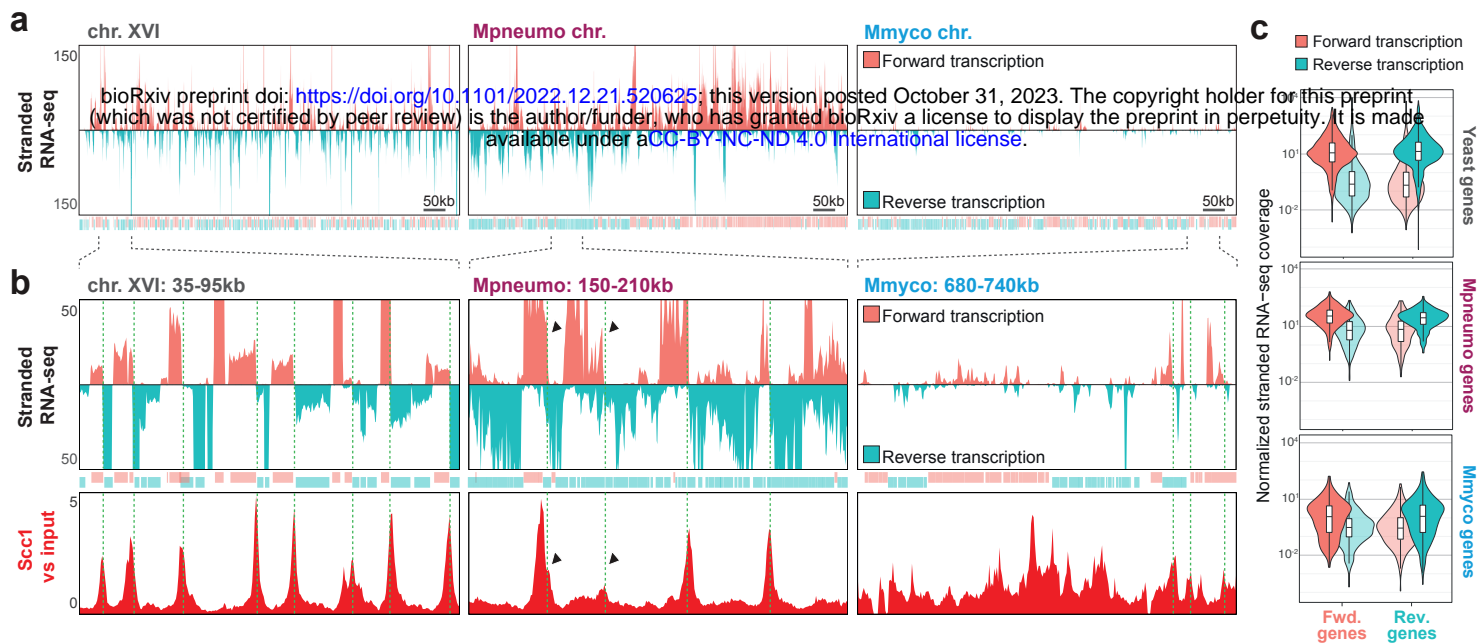


Fig. 2

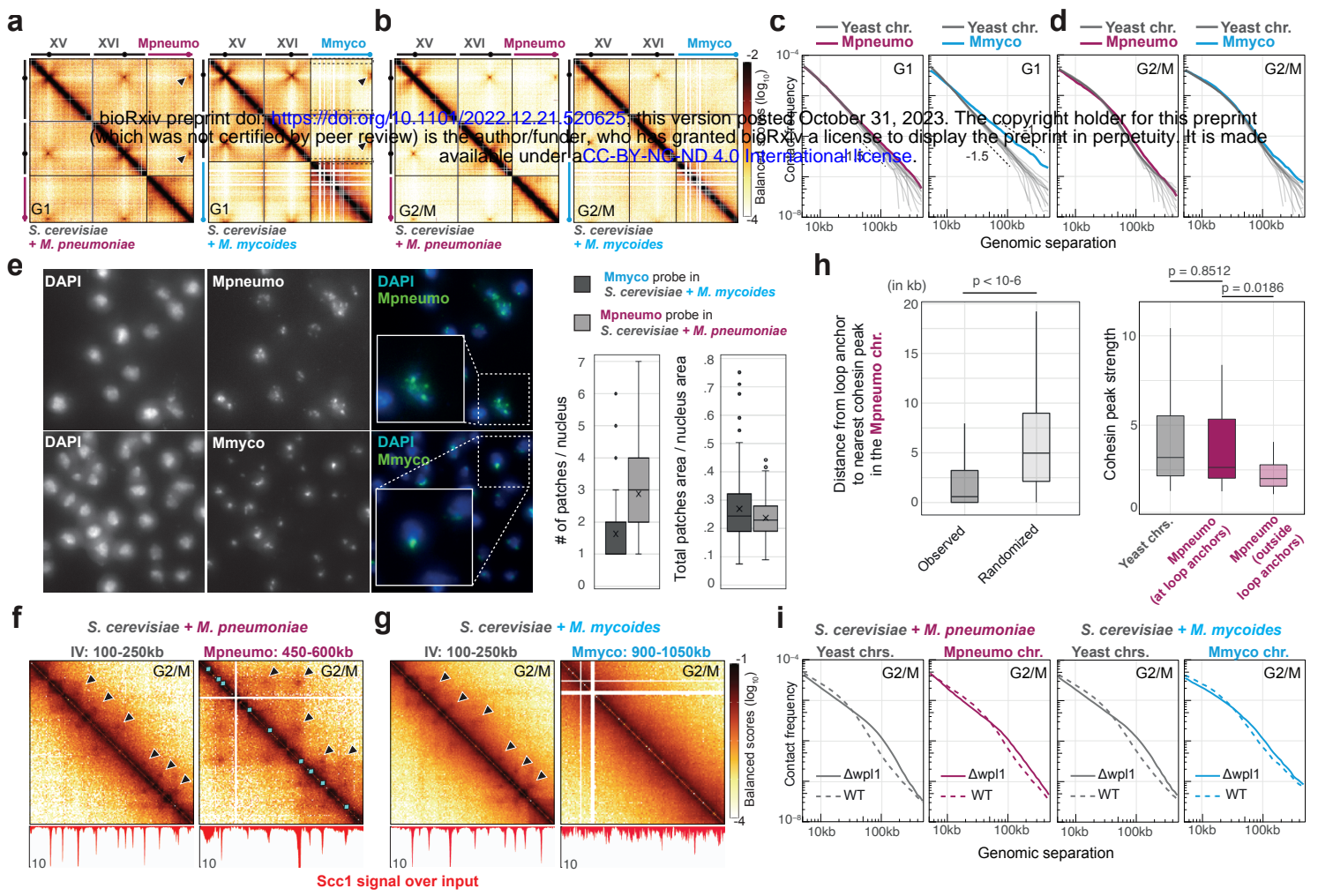


Fig. 3

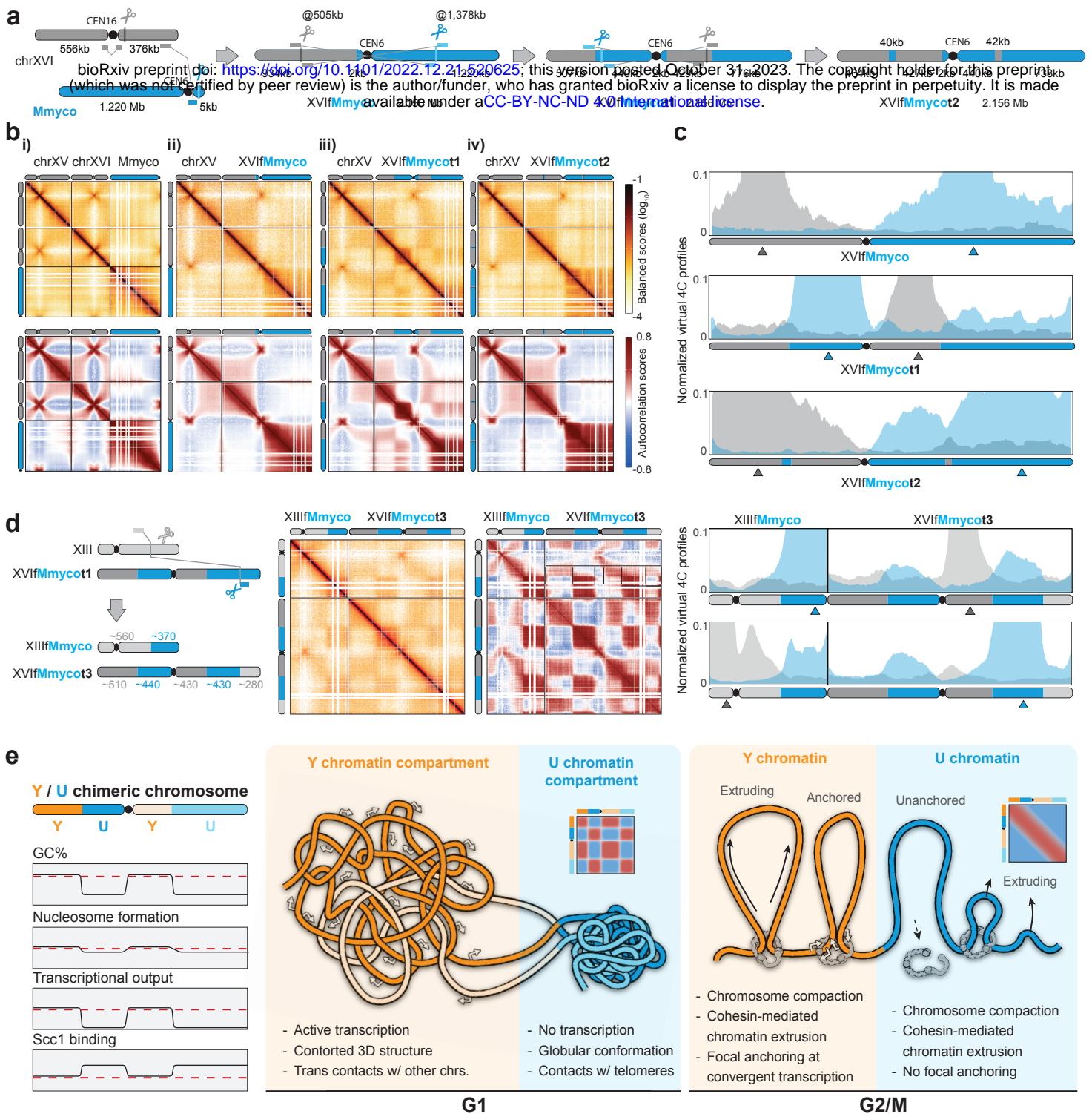


Fig. 4