



1  
00:00:00,790 --> 00:00:07,320

[Music]

2  
00:00:12,320 --> 00:00:09,200

[Applause]

3  
00:00:15,110 --> 00:00:12,330

thank you so much Madeline thank you for

4  
00:00:16,340 --> 00:00:15,120

staying this late and we're going to

5  
00:00:19,250 --> 00:00:16,350

talk about the evolution of complex

6  
00:00:21,500 --> 00:00:19,260

genetics as a part of a larger program

7  
00:00:23,740 --> 00:00:21,510

looking at the evolution of complexity

8  
00:00:25,760 --> 00:00:23,750

throughout life a couple of previous

9  
00:00:27,290 --> 00:00:25,770

speakers have talked about this I'm not

10  
00:00:31,220 --> 00:00:27,300

going to talk about the transitions

11  
00:00:33,260 --> 00:00:31,230

leading to complex life we look at

12  
00:00:34,640 --> 00:00:33,270

complex function particularly how the

13  
00:00:37,400 --> 00:00:34,650

complex organism such as ourselves

14

00:00:39,170 --> 00:00:37,410

evolve and not in terms of specific at

15

00:00:41,180 --> 00:00:39,180

that amaze you know has the human

16

00:00:44,090 --> 00:00:41,190

eyeball evolved but how does things like

17

00:00:46,490 --> 00:00:44,100

imaging vision evolve an imaging vision

18

00:00:48,620 --> 00:00:46,500

is evolved multiple times and through

19

00:00:51,500 --> 00:00:48,630

different independent routes things like

20

00:00:53,530 --> 00:00:51,510

powered flight also these complex

21

00:00:55,340 --> 00:00:53,540

functions have evolved independently

22

00:00:57,139 --> 00:00:55,350

multiple times once you've reached

23

00:01:00,260 --> 00:00:57,149

multicellularity which is a completely

24

00:01:02,900 --> 00:01:00,270

different issue so we want to know why

25

00:01:05,150 --> 00:01:02,910

is this happen and and therefore could

26

00:01:06,740 --> 00:01:05,160

it happen on other worlds with a life

27

00:01:09,139 --> 00:01:06,750

capable of doing that what's the

28

00:01:11,330 --> 00:01:09,149

requirements in order to happen life

29

00:01:13,789 --> 00:01:11,340

with complex function you have to be

30

00:01:16,010 --> 00:01:13,799

able to have differentiated cells cells

31

00:01:18,670 --> 00:01:16,020

that can perform lots of functions have

32

00:01:20,990 --> 00:01:18,680

specialized molecules within them and

33

00:01:23,270 --> 00:01:21,000

they have to be put in place in the

34

00:01:24,740 --> 00:01:23,280

right place and at the right time and in

35

00:01:26,300 --> 00:01:24,750

developmental programs after then we

36

00:01:27,679 --> 00:01:26,310

take it away again at the right place in

37

00:01:29,929 --> 00:01:27,689

the right time so this is a very

38

00:01:31,609 --> 00:01:29,939

complicated control process and that

39

00:01:34,039 --> 00:01:31,619

speaks to there being a complex genetic

40

00:01:36,500 --> 00:01:34,049

program underneath that to make that all

41

00:01:38,810 --> 00:01:36,510

work so we've been asking is this the

42

00:01:40,730 --> 00:01:38,820

evolution of this complex genetic

43

00:01:43,190 --> 00:01:40,740

program the capability to evolve a

44

00:01:47,270 --> 00:01:43,200

complex program itself a bottleneck in

45

00:01:49,550 --> 00:01:47,280

the evolution of complex life complex

46

00:01:51,350 --> 00:01:49,560

genetics is pretty much restricted to

47

00:01:54,219 --> 00:01:51,360

the eukaryotes of this sort of

48

00:01:56,870 --> 00:01:54,229

complexity so eukaryotes tend to have

49

00:01:58,940 --> 00:01:56,880

big genomes this is quite an old plot

50

00:02:01,609 --> 00:01:58,950

this is all the completely sequence

51  
00:02:03,440 --> 00:02:01,619  
genomes I could find our database these

52  
00:02:06,020 --> 00:02:03,450  
green guys here the eukaryotes along

53  
00:02:10,850 --> 00:02:06,030  
here we have gene and size in log scale

54  
00:02:12,380 --> 00:02:10,860  
and mega bases rafted Giga bases here is

55  
00:02:14,539 --> 00:02:12,390  
a huge amounts of information and

56  
00:02:16,130 --> 00:02:14,549  
eukaryotes can develop and have

57  
00:02:18,020 --> 00:02:16,140  
developed independently in multiple

58  
00:02:20,960 --> 00:02:18,030  
times very complex developmental

59  
00:02:22,530 --> 00:02:20,970  
programs by contrast prokaryotes and I

60  
00:02:24,179 --> 00:02:22,540  
don't apologize for lumping every

61  
00:02:25,740 --> 00:02:24,189  
else together in prokaryotes and it's

62  
00:02:27,660 --> 00:02:25,750  
old-fashioned there you go

63  
00:02:29,759 --> 00:02:27,670

tend to have much smaller genomes and

64

00:02:32,309 --> 00:02:29,769

when they do develop multicellularity it

65

00:02:34,589 --> 00:02:32,319

is simple this is limited a very limited

66

00:02:36,660 --> 00:02:34,599

number of cell types and it is quite

67

00:02:38,130 --> 00:02:36,670

often faculty to do so they can switch

68

00:02:40,830 --> 00:02:38,140

between unicellular and multicellular

69

00:02:45,149 --> 00:02:40,840

states you and I can't do that

70

00:02:47,369 --> 00:02:45,159

well I can't with the genomes do overlap

71

00:02:49,229 --> 00:02:47,379

in sizes there's there's not a distinct

72

00:02:50,699 --> 00:02:49,239

size gap where you have to jump from

73

00:02:52,649 --> 00:02:50,709

prokaryote to eukaryote and they also

74

00:02:54,449 --> 00:02:52,659

overlap in chemistry and this is

75

00:02:56,879 --> 00:02:54,459

something we spent rather too much time

76

00:02:58,440 --> 00:02:56,889

illustrating but I'll try to be brief if

77

00:03:01,140 --> 00:02:58,450

you look at all the chemistry that's

78

00:03:03,780 --> 00:03:01,150

involved in how genes are controlled in

79

00:03:06,119 --> 00:03:03,790

an organism so you have DNA binding to

80

00:03:08,129 --> 00:03:06,129

RNA RNA binding to proteins proteins

81

00:03:10,710 --> 00:03:08,139

binding to DNA there's a whole zoo of

82

00:03:12,659 --> 00:03:10,720

interactions and you can modulate any

83

00:03:14,250 --> 00:03:12,669

one of these by reacting them with

84

00:03:16,530 --> 00:03:14,260

different chemicals and then each of

85

00:03:19,740 --> 00:03:16,540

them can control different steps in the

86

00:03:21,539 --> 00:03:19,750

in the process of activating a gene so

87

00:03:22,770 --> 00:03:21,549

from the initial transcription all the

88

00:03:24,270 --> 00:03:22,780

way down here to breaking down the

89

00:03:26,719 --> 00:03:24,280

protein in the end of the process and

90

00:03:29,219 --> 00:03:26,729

you ask what combinations actually occur

91

00:03:31,409 --> 00:03:29,229

okay so you can have a targeting moiety

92

00:03:32,640 --> 00:03:31,419

a protein molecule over here and it

93

00:03:34,259 --> 00:03:32,650

binds to something else and then

94

00:03:36,240 --> 00:03:34,269

something happens so you've got

95

00:03:37,830 --> 00:03:36,250

targeting targeted and what happens and

96

00:03:39,809 --> 00:03:37,840

if you look at what happens in the

97

00:03:43,110 --> 00:03:39,819

eukaryotes the answer is pretty much

98

00:03:45,479 --> 00:03:43,120

everything and you look at what happens

99

00:03:48,240 --> 00:03:45,489

in the prokaryotes and the answer is

100

00:03:49,740 --> 00:03:48,250

pretty much everything all the

101

00:03:52,289 --> 00:03:49,750

combinations are there there isn't a

102

00:03:54,390 --> 00:03:52,299

distinct chemical pattern of this the

103

00:03:56,280 --> 00:03:54,400

chemistry is similar in all the major

104

00:03:58,259 --> 00:03:56,290

domains differently distributed

105

00:03:59,879 --> 00:03:58,269

different emphasis but there's not a

106

00:04:01,979 --> 00:03:59,889

fundamental chemical difference and

107

00:04:03,449 --> 00:04:01,989

these independently evolved they are not

108

00:04:04,710 --> 00:04:03,459

the same and we know that independent

109

00:04:06,420 --> 00:04:04,720

evolved because they're completely

110

00:04:09,030 --> 00:04:06,430

different sequences different structures

111

00:04:11,789 --> 00:04:09,040

and so on and so how our hypothesis

112

00:04:16,229 --> 00:04:11,799

which we developed is that the inherent

113

00:04:20,129 --> 00:04:16,239

difference is the control logic and that

114

00:04:22,860 --> 00:04:20,139

eukaryotes us have a gene structure that

115

00:04:25,020 --> 00:04:22,870

is by default off if you add a gene to a

116

00:04:26,610 --> 00:04:25,030

eukaryotic genome it will by default not

117

00:04:29,820 --> 00:04:26,620

do anything you have to do a lot of work

118

00:04:32,490 --> 00:04:29,830

and activity to turn it on prokaryotes

119

00:04:34,560 --> 00:04:32,500

have a default on logic if you add a bit

120

00:04:36,980 --> 00:04:34,570

of DNA to a prokaryotic genome it will

121

00:04:38,970 --> 00:04:36,990

tend to do something

122

00:04:40,470 --> 00:04:38,980

there's lots of everything I won't go

123

00:04:42,120 --> 00:04:40,480

into this we've written it very long and

124

00:04:45,540 --> 00:04:42,130

to be honest rather boring paper about

125

00:04:47,190 --> 00:04:45,550

this I should I should emphasize this is

126

00:04:50,010 --> 00:04:47,200

a style of control

127

00:04:52,170 --> 00:04:50,020

this isn't absolute okay see course you

128

00:04:53,910 --> 00:04:52,180

can you find examples of gene systems in

129

00:04:55,590 --> 00:04:53,920

prokaryotes that default off of course

130

00:04:58,770 --> 00:04:55,600

you can find ones in eukaryotes that are

131

00:05:00,630 --> 00:04:58,780

default on but it is a general style of

132

00:05:02,970 --> 00:05:00,640

control and our hypothesis which I'm not

133

00:05:07,350 --> 00:05:02,980

going to justify today extent of time is

134

00:05:09,810 --> 00:05:07,360

that this is related to why eukaryotic

135

00:05:12,120 --> 00:05:09,820

genomes can be amplified through gene

136

00:05:16,040 --> 00:05:12,130

duplication and made more complex in in

137

00:05:20,250 --> 00:05:16,050

it pre adapts you to be able to do that

138

00:05:21,690 --> 00:05:20,260

okay so is that true while we can't go

139

00:05:22,980 --> 00:05:21,700

back in with a time machine you know the

140

00:05:25,590 --> 00:05:22,990

snowflake thing is great but we'd have

141

00:05:27,090 --> 00:05:25,600

to run the snowflakes a yeast snowflake

142

00:05:29,010 --> 00:05:27,100

experience experiment for a million

143

00:05:30,750 --> 00:05:29,020

years to to test this that's not really

144

00:05:34,110 --> 00:05:30,760

practical with a five year project grant

145

00:05:36,390 --> 00:05:34,120

so we are so we we thought we model it

146

00:05:38,520 --> 00:05:36,400

and wanted as a model of the sort of

147

00:05:40,730 --> 00:05:38,530

complexity that occurs in genetic

148

00:05:43,710 --> 00:05:40,740

systems and evolving genetic systems

149

00:05:45,090 --> 00:05:43,720

which are spaghetti code and if you look

150

00:05:46,650 --> 00:05:45,100

at the sort of control maps that

151  
00:05:48,360 --> 00:05:46,660  
biochemists like to put up on their wall

152  
00:05:50,760 --> 00:05:48,370  
to show how much are they are they are

153  
00:05:52,350 --> 00:05:50,770  
completely spaghetti code control

154  
00:05:55,700 --> 00:05:52,360  
systems there's no layers there's no

155  
00:05:57,630 --> 00:05:55,710  
modularity so we wanted a model app and

156  
00:05:59,490 --> 00:05:57,640  
there's all the feedback between gene

157  
00:06:01,560 --> 00:05:59,500  
products and gene expression so this

158  
00:06:03,120 --> 00:06:01,570  
sort of highly abstract Network

159  
00:06:05,370 --> 00:06:03,130  
formulation didn't really work for us

160  
00:06:07,530 --> 00:06:05,380  
but equally we didn't want a model of

161  
00:06:09,240 --> 00:06:07,540  
detail chemistry this sort of thing down

162  
00:06:11,190 --> 00:06:09,250  
here proteins behind the DNA because if

163  
00:06:13,620 --> 00:06:11,200

you try to do that on a on a cell basis

164

00:06:15,390 --> 00:06:13,630

then well that would be too complicated

165

00:06:17,130 --> 00:06:15,400

we actually I wanted something that

166

00:06:19,260 --> 00:06:17,140

looked like the classic undergraduate

167

00:06:21,570 --> 00:06:19,270

textbook diagram of a jakob and mono

168

00:06:24,030 --> 00:06:21,580

control model for those who biochemists

169

00:06:27,870 --> 00:06:24,040

some a it looks like this okay so that's

170

00:06:30,840 --> 00:06:27,880

what we built so this is the model and

171

00:06:32,670 --> 00:06:30,850

it's conceptually quite simple I'll tell

172

00:06:35,520 --> 00:06:32,680

you about the coding later that's a

173

00:06:37,590 --> 00:06:35,530

disaster so we have a population of

174

00:06:39,300 --> 00:06:37,600

organisms in yes we do have five

175

00:06:42,870 --> 00:06:39,310

organisms at the moment in the model and

176  
00:06:44,960 --> 00:06:42,880  
those organisms contain genes and those

177  
00:06:48,060 --> 00:06:44,970  
genes can be active and produce

178  
00:06:49,470 --> 00:06:48,070  
transcripts we need selection in this

179  
00:06:52,650 --> 00:06:49,480  
this is an evolution area mode

180  
00:06:55,140 --> 00:06:52,660  
so for an organism to be fit those

181  
00:06:57,270 --> 00:06:55,150  
transcripts must match an environment

182  
00:06:59,180 --> 00:06:57,280  
and there are positive aspects of that

183  
00:07:01,800 --> 00:06:59,190  
environment things it must do and

184  
00:07:04,110 --> 00:07:01,810  
negative aspects things it must not do

185  
00:07:05,340 --> 00:07:04,120  
if it's just positive then you just

186  
00:07:09,480 --> 00:07:05,350  
select something that makes everything

187  
00:07:10,950 --> 00:07:09,490  
and then you is one ok how do you tell

188  
00:07:13,530 --> 00:07:10,960

whether you've got a transcript what the

189

00:07:16,230 --> 00:07:13,540

transcription of each gene is controlled

190

00:07:17,790 --> 00:07:16,240

by regulatory elements and you've got

191

00:07:20,100 --> 00:07:17,800

positive regulatory elements and

192

00:07:23,010 --> 00:07:20,110

negative regulator elements positive

193

00:07:24,720 --> 00:07:23,020

turn it on negative turn it on a note

194

00:07:26,310 --> 00:07:24,730

we've only got one type of sequence here

195

00:07:28,380 --> 00:07:26,320

we haven't got translation built into

196

00:07:30,690 --> 00:07:28,390

this I thought of doing that and then

197

00:07:32,610 --> 00:07:30,700

thought it's too complicated at this

198

00:07:34,650 --> 00:07:32,620

stage so if you wonder if you believe in

199

00:07:38,970 --> 00:07:34,660

the RNA world hypothesis this is a sort

200

00:07:40,220 --> 00:07:38,980

of RNA world type Gina how do you

201  
00:07:42,990 --> 00:07:40,230  
control the genes

202  
00:07:44,880 --> 00:07:43,000  
well pretty straightforward we've got

203  
00:07:47,040 --> 00:07:44,890  
positive elements and negative elements

204  
00:07:48,840 --> 00:07:47,050  
if the number of active positive

205  
00:07:51,570 --> 00:07:48,850  
elements outweighs the number of active

206  
00:07:54,360 --> 00:07:51,580  
negative elements it's on if it doesn't

207  
00:07:56,190 --> 00:07:54,370  
it's off keep it is want to be mean by

208  
00:07:57,960 --> 00:07:56,200  
active we want feedback between what the

209  
00:08:01,260 --> 00:07:57,970  
genome does and how it controls its

210  
00:08:04,530 --> 00:08:01,270  
genes so a genome produces a sequence

211  
00:08:07,530 --> 00:08:04,540  
here a transcript and we ask for each

212  
00:08:09,840 --> 00:08:07,540  
regulatory element here does that match

213  
00:08:14,430 --> 00:08:09,850

a transcript that's being made at the

214

00:08:16,290 --> 00:08:14,440

moment so here's a gene two on one off

215

00:08:18,720 --> 00:08:16,300

this one is produced let's say the

216

00:08:21,570 --> 00:08:18,730

transcript of this sequence here's

217

00:08:24,530 --> 00:08:21,580

another gene this regulatory element has

218

00:08:28,200 --> 00:08:24,540

sequence ACA that matches that bit there

219

00:08:30,990 --> 00:08:28,210

so that's all this one is see see see

220

00:08:34,409 --> 00:08:31,000

that doesn't match anything here that's

221

00:08:36,390 --> 00:08:34,419

not on this is BC that's not on this is

222

00:08:38,909 --> 00:08:36,400

see that matches that one there so

223

00:08:41,370 --> 00:08:38,919

that's on and you do this for all these

224

00:08:44,400 --> 00:08:41,380

regulator elements in all the genes

225

00:08:46,080 --> 00:08:44,410

matching all the transcripts and that

226  
00:08:49,260 --> 00:08:46,090  
tells you whether that particular gene

227  
00:08:52,500 --> 00:08:49,270  
is on or not two things to note this is

228  
00:08:53,820 --> 00:08:52,510  
a model of regulatory control of genes

229  
00:08:55,290 --> 00:08:53,830  
is not structural they're not saying

230  
00:08:57,270 --> 00:08:55,300  
this is an enzyme or something does

231  
00:09:01,950 --> 00:08:57,280  
something clever so it's a model of gene

232  
00:09:03,280 --> 00:09:01,960  
regulation and secondly this is a bit

233  
00:09:05,110 --> 00:09:03,290  
obvious but but

234  
00:09:07,210 --> 00:09:05,120  
said anyway because it's late in the day

235  
00:09:10,660 --> 00:09:07,220  
and you know your blood glucose levels

236  
00:09:13,780 --> 00:09:10,670  
are dropping if your regulatory sequence

237  
00:09:17,439 --> 00:09:13,790  
here is short it is more likely to match

238  
00:09:19,569 --> 00:09:17,449

something here so if you have a short

239

00:09:24,610 --> 00:09:19,579

regulatory sequence it's more likely to

240

00:09:26,920 --> 00:09:24,620

be active okay lots of variables in

241

00:09:28,240 --> 00:09:26,930

incomes of it my goodness you can have

242

00:09:29,949 --> 00:09:28,250

fun with this so we're not going to go

243

00:09:31,960 --> 00:09:29,959

through all that we're going to look to

244

00:09:33,879 --> 00:09:31,970

some summary statistics of what the

245

00:09:36,730 --> 00:09:33,889

output looks at and there are two things

246

00:09:39,939 --> 00:09:36,740

we're going to focus on one in

247

00:09:41,379 --> 00:09:39,949

particular if you have a lot of these

248

00:09:42,970 --> 00:09:41,389

negative elements and you could lose

249

00:09:45,249 --> 00:09:42,980

elements through selection they can be

250

00:09:46,960 --> 00:09:45,259

deleted so if an organism decides it

251  
00:09:48,370 --> 00:09:46,970  
only needs one regulatory element that's

252  
00:09:51,100 --> 00:09:48,380  
absolutely fine eventually it'll get

253  
00:09:52,689 --> 00:09:51,110  
that lots of negative elements compared

254  
00:09:55,449 --> 00:09:52,699  
to positive suggests you've got us of

255  
00:09:58,030 --> 00:09:55,459  
default off state now be more likely to

256  
00:09:59,920 --> 00:09:58,040  
be off than on and lots of short

257  
00:10:01,480 --> 00:09:59,930  
negative elements will mean you'll have

258  
00:10:09,629 --> 00:10:01,490  
a default off state they're more likely

259  
00:10:13,269 --> 00:10:11,680  
there are lots of outputs you can

260  
00:10:15,340 --> 00:10:13,279  
collect from this so you can measure

261  
00:10:17,769 --> 00:10:15,350  
average this is just an example run

262  
00:10:19,240 --> 00:10:17,779  
measure average dream gene length which

263  
00:10:20,889 --> 00:10:19,250

turns out to be quite interesting but I

264

00:10:22,840 --> 00:10:20,899

won't go into why you can measure the

265

00:10:24,160 --> 00:10:22,850

number of express genes number producing

266

00:10:27,639 --> 00:10:24,170

transcripts which turns out to be quite

267

00:10:29,319 --> 00:10:27,649

dull and I won't say why you start out

268

00:10:31,990 --> 00:10:29,329

with an entirely random genome and it

269

00:10:33,250 --> 00:10:32,000

adapts fairly rapidly and then sort of

270

00:10:36,730 --> 00:10:33,260

either plateaus or

271

00:10:38,050 --> 00:10:36,740

or slowly increases adaptation and you

272

00:10:39,910 --> 00:10:38,060

can look at the fitness and that turns

273

00:10:42,040 --> 00:10:39,920

out to be really confusing but I went to

274

00:10:45,250 --> 00:10:42,050

a why what I got to focus on is is this

275

00:10:47,350 --> 00:10:45,260

thing here this is for each gene pay

276

00:10:49,650 --> 00:10:47,360

attention now this is complicated the

277

00:10:52,470 --> 00:10:49,660

each gene you look at the shortest

278

00:10:54,819 --> 00:10:52,480

positive and the shortest negative

279

00:10:58,629 --> 00:10:54,829

regulatory element so that's the one

280

00:11:00,370 --> 00:10:58,639

most likely to be active and then you

281

00:11:02,170 --> 00:11:00,380

say for all the genes in the genome what

282

00:11:06,490 --> 00:11:02,180

is the average length of those short

283

00:11:09,129 --> 00:11:06,500

positive and short negative elements if

284

00:11:11,050 --> 00:11:09,139

the short negative ones are shorter on

285

00:11:13,449 --> 00:11:11,060

average they're more likely to be on

286

00:11:15,550 --> 00:11:13,459

that means the genes on average are more

287

00:11:17,490 --> 00:11:15,560

likely to be suppressed off you've got a

288

00:11:21,120 --> 00:11:17,500

default negative

289

00:11:22,860 --> 00:11:21,130

default off-mode if the Shh if the

290

00:11:24,870 --> 00:11:22,870

positive one said to be shorter they're

291

00:11:31,520 --> 00:11:24,880

more likely to be on and so you have a

292

00:11:34,020 --> 00:11:31,530

default on mode so in this example

293

00:11:36,120 --> 00:11:34,030

here's the average minimum regulatory

294

00:11:38,910 --> 00:11:36,130

element length blah blah blah rate is

295

00:11:42,360 --> 00:11:38,920

negative blue is positive the red is a

296

00:11:44,010 --> 00:11:42,370

little bit longer than the blue there's

297

00:11:46,410 --> 00:11:44,020

a lot of noise on that because I had the

298

00:11:47,790 --> 00:11:46,420

mutation turned up too high this is a

299

00:11:49,350 --> 00:11:47,800

great thing about doing life in

300

00:11:51,450 --> 00:11:49,360

computers you can turn mutation up and

301  
00:11:54,420 --> 00:11:51,460  
down and things like that so this is yes

302  
00:11:57,840 --> 00:11:54,430  
this is very weakly a prokaryotic

303  
00:12:00,690 --> 00:11:57,850  
default on mo but he's pretty weak it's

304  
00:12:02,760 --> 00:12:00,700  
pretty messy I mean yeah you wouldn't

305  
00:12:06,650 --> 00:12:02,770  
call that a good transit signal in it

306  
00:12:09,060 --> 00:12:06,660  
and a exoplanet good lecture with you so

307  
00:12:10,770 --> 00:12:09,070  
here's a few more is another one - doing

308  
00:12:12,480 --> 00:12:10,780  
the same thing I just dialed down the

309  
00:12:15,180 --> 00:12:12,490  
mutation rate so it's less noisy but

310  
00:12:17,160 --> 00:12:15,190  
it's it's weakly default on I wouldn't

311  
00:12:19,020 --> 00:12:17,170  
say anything this one can't decide what

312  
00:12:20,670 --> 00:12:19,030  
it's doing it's just wobbling all over

313  
00:12:22,740 --> 00:12:20,680

the place but these two are more

314

00:12:25,650 --> 00:12:22,750

interesting here we've started to evolve

315

00:12:29,760 --> 00:12:25,660

a clear pattern a signal of some sort

316

00:12:31,590 --> 00:12:29,770

this one the positive ones these are the

317

00:12:33,750 --> 00:12:31,600

elements that turn things on tend to be

318

00:12:37,230 --> 00:12:33,760

shorter suggest you go the prokaryotic

319

00:12:38,910 --> 00:12:37,240

type default on genetics evolving and

320

00:12:43,290 --> 00:12:38,920

this is the other way around you've got

321

00:12:45,180 --> 00:12:43,300

the negative evolving and you do this a

322

00:12:47,430 --> 00:12:45,190

lot for different combinations of

323

00:12:49,740 --> 00:12:47,440

parameters and different in our days of

324

00:12:51,960 --> 00:12:49,750

the week and things and you try to

325

00:12:54,330 --> 00:12:51,970

correlate are you getting default on or

326

00:12:56,220 --> 00:12:54,340

default off with all the different

327

00:12:59,130 --> 00:12:56,230

inputs and this is what we've got so far

328

00:13:00,990 --> 00:12:59,140

this is really preliminary this is just

329

00:13:03,540 --> 00:13:01,000

straightforward correlation coefficient

330

00:13:06,420 --> 00:13:03,550

greater than I think 32 is significant

331

00:13:09,150 --> 00:13:06,430

to 95% and those are colored in red here

332

00:13:11,010 --> 00:13:09,160

and one really surprising thing that

333

00:13:12,780 --> 00:13:11,020

left out I thought you know it's going

334

00:13:14,220 --> 00:13:12,790

to be you put complicated environments

335

00:13:15,780 --> 00:13:14,230

in the genomes gonna have to be

336

00:13:17,580 --> 00:13:15,790

complicated to respond to each and

337

00:13:20,070 --> 00:13:17,590

you'll have eukaryotic and you'll have

338

00:13:24,330 --> 00:13:20,080

kangaroos evolving it didn't care what

339

00:13:27,570 --> 00:13:24,340

the environment was wow that really

340

00:13:29,460 --> 00:13:27,580

annoyed me it was all about genome

341

00:13:30,780 --> 00:13:29,470

structure the correlations that were

342

00:13:31,380 --> 00:13:30,790

more significant was about genome

343

00:13:34,290 --> 00:13:31,390

structure

344

00:13:36,389 --> 00:13:34,300

and the thing that correlative would

345

00:13:38,190 --> 00:13:36,399

default off evolving that sort of thing

346

00:13:39,750 --> 00:13:38,200

where you've got a nice curve separating

347

00:13:44,699 --> 00:13:39,760

like this saying you've got a default of

348

00:13:46,259 --> 00:13:44,709

wealth having a lot of genes you could

349

00:13:47,970 --> 00:13:46,269

also solve your world alter the number

350

00:13:51,540 --> 00:13:47,980

of bases in your DNA in this - you could

351  
00:13:56,759 --> 00:13:51,550  
say let's have a seven base genome let's

352  
00:13:59,130 --> 00:13:56,769  
see Steve better do that few bases

353  
00:14:01,740 --> 00:13:59,140  
simpler genome correlated with default

354  
00:14:04,290 --> 00:14:01,750  
off short genes so this is simple

355  
00:14:08,060 --> 00:14:04,300  
genomes it's the number of genes that

356  
00:14:10,199 --> 00:14:08,070  
correlated weird so this isn't

357  
00:14:13,500 --> 00:14:10,209  
inconsistent with the original idea that

358  
00:14:15,480 --> 00:14:13,510  
having a lot of genes means it's or

359  
00:14:18,210 --> 00:14:15,490  
favors evolution of this Vieux Carre

360  
00:14:22,079 --> 00:14:18,220  
optic type default off genetics it's not

361  
00:14:25,230 --> 00:14:22,089  
entirely supportive of it it's about

362  
00:14:27,449 --> 00:14:25,240  
many regulatory elements not their size

363  
00:14:28,860 --> 00:14:27,459

or complexity and this is this sort of

364

00:14:30,329 --> 00:14:28,870

this is why I mentioned the RNA world

365

00:14:31,680 --> 00:14:30,339

you know because this sort of looks like

366

00:14:35,490 --> 00:14:31,690

that doesn't it lots and lots and lots

367

00:14:38,310 --> 00:14:35,500

of RNAs a little short ones and this

368

00:14:42,090 --> 00:14:38,320

hints that default off was the standard

369

00:14:45,630 --> 00:14:42,100

for first prototype so caveat it's only

370

00:14:47,100 --> 00:14:45,640

a model okay I wish more astrobiology

371

00:14:49,019 --> 00:14:47,110

talks would say it's only a model

372

00:14:52,680 --> 00:14:49,029

because quite often it is it's only a

373

00:14:57,600 --> 00:14:52,690

model in Excel I did this in Excel okay

374

00:14:59,220 --> 00:14:57,610

this is a bad idea and I'm very

375

00:15:00,870 --> 00:14:59,230

fortunate then Enrico Borriello it's

376

00:15:02,490 --> 00:15:00,880

Sarah Walker's groupid AC who is now

377

00:15:05,310 --> 00:15:02,500

recoding this into something this

378

00:15:07,920 --> 00:15:05,320

actually looks like code so conclusion

379

00:15:09,300 --> 00:15:07,930

genetics start ends differ I'll be

380

00:15:11,100 --> 00:15:09,310

developed a model of that that's not

381

00:15:13,530 --> 00:15:11,110

completely abstract and yet not

382

00:15:16,530 --> 00:15:13,540

chemically specific initial results hint

383

00:15:18,449 --> 00:15:16,540

and this is just a hint that more genes

384

00:15:20,880 --> 00:15:18,459

mean default off style

385

00:15:23,400 --> 00:15:20,890

it's about regulatory group number not

386

00:15:24,350 --> 00:15:23,410

about genome complexity and that hints

387

00:15:26,460 --> 00:15:24,360

that this might be a primitive

388

00:15:30,710 --> 00:15:26,470

characteristic in other words you and I

389

00:15:35,720 --> 00:15:30,720

are more similar to Luca than e.coli

390

00:15:38,040 --> 00:15:35,730

your thoughts welcome even yours and

391

00:15:39,689 --> 00:15:38,050

because this is really pretty Murray and

392

00:15:42,389 --> 00:15:39,699

we want your thoughts and input before

393

00:15:45,060 --> 00:15:42,399

we go on to the next stage those are the

394

00:15:49,230 --> 00:15:45,070

papers and one last call out

395

00:15:50,520 --> 00:15:49,240

mdps life I've published in life several

396

00:15:53,580 --> 00:15:50,530

times and served one of my colleagues

397

00:15:54,840 --> 00:15:53,590

and we published in another journal at

398

00:15:56,460 --> 00:15:54,850

all you'll be familiar with that I

399

00:15:59,360 --> 00:15:56,470

wouldn't name that we publishing quite a

400

00:16:03,360 --> 00:15:59,370

bit in this context and these guys are

401  
00:16:07,500 --> 00:16:03,370  
professional faster cheaper and open

402  
00:16:09,720 --> 00:16:07,510  
access so I'd have a look at em DPI's

403  
00:16:12,210 --> 00:16:09,730  
life as a potential and output for your

404  
00:16:13,350 --> 00:16:12,220  
next paper I think you'll be pleasantly

405  
00:16:15,600 --> 00:16:13,360  
surprised by the level of

406  
00:16:18,210 --> 00:16:15,610  
professionalism and speed with which

407  
00:16:19,610 --> 00:16:18,220  
they can get your stuff out thank you

408  
00:16:19,950 --> 00:16:19,620  
very much

409  
00:16:21,870 --> 00:16:19,960  
[Applause]