



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

[Music]

2  
00:00:13,930 --> 00:00:09,460

[Applause]

3  
00:00:19,269 --> 00:00:13,940

thank you it's an opportunity to present

4  
00:00:22,130 --> 00:00:19,279

Antony today Antony is based on

5  
00:00:24,679 --> 00:00:22,140

primarily microscope that's an atomic

6  
00:00:26,900 --> 00:00:24,689

force microscope it was named in honor

7  
00:00:29,720 --> 00:00:26,910

of the father of microbiology Anthony

8  
00:00:32,030 --> 00:00:29,730

and Meagan Leeuwenhoek I the clever

9  
00:00:35,930 --> 00:00:32,040

naming for that as due to brent chris

10  
00:00:40,130 --> 00:00:35,940

iners Hey midnite acronym searching

11  
00:00:46,120 --> 00:00:40,140

algorithm uh-huh the I'm gonna just get

12  
00:00:48,680 --> 00:00:46,130

started in here okay so I as you know

13  
00:00:50,090 --> 00:00:48,690

one of the primary goals of the Europe

14

00:00:53,030 --> 00:00:50,100

Lander science definition team was

15

00:00:55,040 --> 00:00:53,040

address to address characterizing bio

16

00:00:57,470 --> 00:00:55,050

signatures in Europe as near subsurface

17

00:00:59,660 --> 00:00:57,480

materials there are a couple of

18

00:01:02,389 --> 00:00:59,670

objectives that we address with a

19

00:01:04,520 --> 00:01:02,399

microscope the first is to identify and

20

00:01:06,650 --> 00:01:04,530

characterize morphological and/or

21

00:01:09,800 --> 00:01:06,660

textural features of Europe this near

22

00:01:12,139 --> 00:01:09,810

subsurface and the investigation that

23

00:01:13,880 --> 00:01:12,149

Anthony accomplishes here is to

24

00:01:17,539 --> 00:01:13,890

characterize particulates and samples

25

00:01:19,969 --> 00:01:17,549

that are 0.2 microns to 1500 microns in

26

00:01:21,950 --> 00:01:19,979

scale the second objective the antony

27

00:01:23,570 --> 00:01:21,960

addresses is to assess the structural

28

00:01:25,520 --> 00:01:23,580

and compositional nature of europe is

29

00:01:27,500 --> 00:01:25,530

near subsurface materials there are two

30

00:01:29,000 --> 00:01:27,510

investigations here one of them is to

31

00:01:31,310 --> 00:01:29,010

measure the mechanical properties of the

32

00:01:32,840 --> 00:01:31,320

sample and the second is to potentially

33

00:01:37,100 --> 00:01:32,850

determine the composition of

34

00:01:41,710 --> 00:01:37,110

particulates afm you may be familiar

35

00:01:44,240 --> 00:01:41,720

with it's been around for about 40 years

36

00:01:46,340 --> 00:01:44,250

works is a pretty simple instrument and

37

00:01:50,210 --> 00:01:46,350

it works by a cantilever interacting

38

00:01:51,590 --> 00:01:50,220

with a sample it raster's over the

39

00:01:54,530 --> 00:01:51,600

sample and it creates a

40

00:01:56,749 --> 00:01:54,540

three-dimensional topographic map if

41

00:01:58,149 --> 00:01:56,759

Emma's tip is often used in material

42

00:02:02,140 --> 00:01:58,159

science in the semiconductor industry

43

00:02:05,420 --> 00:02:02,150

but can also be used to detect biology

44

00:02:08,270 --> 00:02:05,430

particulates such as this cell were

45

00:02:10,520 --> 00:02:08,280

imaged in a 1999 paper if this came from

46

00:02:13,400 --> 00:02:10,530

Lake Vostok accretion ice you can get

47

00:02:17,030 --> 00:02:13,410

really high-resolution image at the sort

48

00:02:18,970 --> 00:02:17,040

of sub sub cell level even looking at

49

00:02:21,409 --> 00:02:18,980

textures that are on the cell surface

50

00:02:23,599 --> 00:02:21,419

but in addition to getting a 3d

51

00:02:25,280 --> 00:02:23,609

topographical map you

52

00:02:28,220 --> 00:02:25,290

also get nanomechanical properties

53

00:02:29,569 --> 00:02:28,230

things such as the sample stiffness the

54

00:02:31,039 --> 00:02:29,579

dissipation of energy when the

55

00:02:34,339 --> 00:02:31,049

cantilever interacts with the surface

56

00:02:37,339 --> 00:02:34,349

and adhesive properties and so you can

57

00:02:40,280 --> 00:02:37,349

see how sort of a diagram of the cartoon

58

00:02:44,270 --> 00:02:40,290

and the top figure here on the bottom is

59

00:02:46,940 --> 00:02:44,280

a scale showing the stiffness as a

60

00:02:48,649 --> 00:02:46,950

as measured as kiloPascals over eight

61

00:02:54,220 --> 00:02:48,659

orders of magnitude different between

62

00:02:56,959 --> 00:02:54,230

cells and hard materials like steel so

63

00:02:59,030 --> 00:02:56,969

AFM bio signature detection technology

64

00:03:02,809 --> 00:02:59,040

as we have thought about it builds on

65

00:03:06,140 --> 00:03:02,819

spaceflight heritage there are a number

66

00:03:09,229 --> 00:03:06,150

of assets to using AFM in space in

67

00:03:11,509 --> 00:03:09,239

addition to its submicron 3d resolution

68

00:03:14,330 --> 00:03:11,519

and the potential for nano mechanical

69

00:03:15,830 --> 00:03:14,340

property determinations the these

70

00:03:18,020 --> 00:03:15,840

instruments are lightweight they have a

71

00:03:19,939 --> 00:03:18,030

small footprint and low power

72

00:03:24,349 --> 00:03:19,949

consumption they can operate an air

73

00:03:27,080 --> 00:03:24,359

liquid and in vacuum the they've also

74

00:03:29,330 --> 00:03:27,090

been flown recently in space on two

75

00:03:31,429 --> 00:03:29,340

different missions one of them was flown

76  
00:03:34,399 --> 00:03:31,439  
on the Phoenix mission and the other on

77  
00:03:37,399 --> 00:03:34,409  
the Rosetta mission in both cases they

78  
00:03:39,229 --> 00:03:37,409  
imaged particulates on Mars on the

79  
00:03:41,420 --> 00:03:39,239  
Phoenix mission in particulates size

80  
00:03:43,490 --> 00:03:41,430  
distribution and soils on the Phoenix

81  
00:03:45,800 --> 00:03:43,500  
mint and the rosetta mission they

82  
00:03:49,129 --> 00:03:45,810  
measured cometary dust and an observed

83  
00:03:51,939 --> 00:03:49,139  
large particle aggregates mark Bentley

84  
00:03:54,439 --> 00:03:51,949  
wrote a paper post the Rosetta mission

85  
00:03:56,780 --> 00:03:54,449  
that was really on lessons learned from

86  
00:03:59,390 --> 00:03:56,790  
operating AFM's in space which we have

87  
00:04:02,030 --> 00:03:59,400  
adopted into the design going forward

88  
00:04:05,499 --> 00:04:02,040

and thinking about antony some of these

89

00:04:07,429 --> 00:04:05,509

are co registering if you use a no

90

00:04:09,800 --> 00:04:07,439

register ring an optical microscope

91

00:04:12,530 --> 00:04:09,810

image with the atomic force microscope

92

00:04:14,179 --> 00:04:12,540

image including an array of cantilevers

93

00:04:16,520 --> 00:04:14,189

that have different spring constants in

94

00:04:18,589 --> 00:04:16,530

different tip morphologies and then

95

00:04:20,209 --> 00:04:18,599

being able to operate the instrument

96

00:04:22,760 --> 00:04:20,219

both in open and closed loop formats

97

00:04:26,360 --> 00:04:22,770

which they was required on the Rosetta

98

00:04:29,149 --> 00:04:26,370

mission so one of the things that I

99

00:04:31,610 --> 00:04:29,159

think AFM has a real advantage to is

100

00:04:33,080 --> 00:04:31,620

detecting really small bio signatures we

101  
00:04:34,730 --> 00:04:33,090  
don't know to what to expect and going

102  
00:04:36,320 --> 00:04:34,740  
to the surface of Europa

103  
00:04:38,360 --> 00:04:36,330  
and we didn't know what to expect when

104  
00:04:41,150 --> 00:04:38,370  
we went to Lake Vita which is a lake in

105  
00:04:43,279 --> 00:04:41,160  
East Antarctica and the like and McMurdo

106  
00:04:44,540 --> 00:04:43,289  
Dry Valleys where the sand plan I don't

107  
00:04:45,499 --> 00:04:44,550  
know that you can see very well from the

108  
00:04:50,719 --> 00:04:45,509  
back that you can see they're really

109  
00:04:52,010 --> 00:04:50,729  
bright images here these these are what

110  
00:04:54,680 --> 00:04:52,020  
I would call kind of normal-sized

111  
00:04:57,710 --> 00:04:54,690  
bacterial cells and this background sort

112  
00:04:59,120 --> 00:04:57,720  
of starry night barely shows up when

113  
00:05:00,980 --> 00:04:59,130

you're looking at really what we have is

114

00:05:02,499 --> 00:05:00,990

like one of the brightest DNA stains

115

00:05:06,890 --> 00:05:02,509

under epi fluorescent microbes

116

00:05:09,290 --> 00:05:06,900

microscope if we look using an AFM you

117

00:05:11,420 --> 00:05:09,300

can see really really what these what

118

00:05:13,730 --> 00:05:11,430

these particles look like you can zoom

119

00:05:15,110 --> 00:05:13,740

in at higher resolution and and start

120

00:05:19,730 --> 00:05:15,120

looking at the surface textures of the

121

00:05:21,730 --> 00:05:19,740

cells so we have also been interested in

122

00:05:25,930 --> 00:05:21,740

can asking the question can we discern

123

00:05:28,760 --> 00:05:25,940

bio signatures from abiotic particles

124

00:05:32,000 --> 00:05:28,770

using AFM based on their nano mechanical

125

00:05:38,360 --> 00:05:32,010

properties this is a image that we

126

00:05:44,360 --> 00:05:38,370

collected where the we we mixed silica

127

00:05:46,100 --> 00:05:44,370

beads so if I go back here so the round

128

00:05:48,529 --> 00:05:46,110

features which you may or may not be

129

00:05:51,980 --> 00:05:48,539

able to see our silica beads that we

130

00:05:54,560 --> 00:05:51,990

mixed with bacterial cells and if we

131

00:05:56,899 --> 00:05:54,570

look at the forest curves from these

132

00:05:59,029 --> 00:05:56,909

particles we can see that this is on a

133

00:06:03,320 --> 00:05:59,039

glass slide so the black dots are the

134

00:06:05,060 --> 00:06:03,330

are plotted here on the graph the glass

135

00:06:09,200 --> 00:06:05,070

background is not different from the

136

00:06:11,930 --> 00:06:09,210

glass beads that are in blue and we have

137

00:06:13,750 --> 00:06:11,940

particulate organic matter that is as

138

00:06:18,710 --> 00:06:13,760

Roger calls the schmutz from the cells

139

00:06:20,510 --> 00:06:18,720

that is in green this is softer and then

140

00:06:25,159 --> 00:06:20,520

even softer yet are the cells that are

141

00:06:27,320 --> 00:06:25,169

in the and the figure we did a second

142

00:06:31,279 --> 00:06:27,330

kind of similar experiment but in this

143

00:06:32,870 --> 00:06:31,289

case due to a conversation we were

144

00:06:35,029 --> 00:06:32,880

inspired that we had with Mitch Schulte

145

00:06:37,219 --> 00:06:35,039

a few months ago he asked us a question

146

00:06:39,379 --> 00:06:37,229

could you actually discern the

147

00:06:43,070 --> 00:06:39,389

difference between carbonaceous

148

00:06:45,610 --> 00:06:43,080

chondrite meteorite and bio and cells

149

00:06:48,129 --> 00:06:45,620

and so we obtained some i-n

150

00:06:50,409 --> 00:06:48,139

crushed powdered meteorite sample we

151  
00:06:52,960 --> 00:06:50,419  
screened it through a 5 micron sieve and

152  
00:06:56,590 --> 00:06:52,970  
mixed it with three different bacterial

153  
00:06:59,080 --> 00:06:56,600  
cell types in this case the meteorite

154  
00:07:03,670 --> 00:06:59,090  
which is here in the particle in the

155  
00:07:06,460 --> 00:07:03,680  
middle of this aggregate of cells shows

156  
00:07:10,150 --> 00:07:06,470  
up to be quite stiffer not as stiff as

157  
00:07:13,480 --> 00:07:10,160  
the background of glass and then the

158  
00:07:16,000 --> 00:07:13,490  
cells have different they are softer yet

159  
00:07:17,350 --> 00:07:16,010  
and they have different curves for the

160  
00:07:19,210 --> 00:07:17,360  
different types of cells that we had in

161  
00:07:21,100 --> 00:07:19,220  
in this case we mixed an Antarctic

162  
00:07:25,480 --> 00:07:21,110  
bacterium called pseudo Vibrio with

163  
00:07:27,730 --> 00:07:25,490

bacillus and s worse in equine so we

164

00:07:30,610 --> 00:07:27,740

think we can discern particles of

165

00:07:33,969 --> 00:07:30,620

different types of properties from from

166

00:07:36,520 --> 00:07:33,979

life you could imagine then that if we

167

00:07:38,500 --> 00:07:36,530

went and got a library of different

168

00:07:40,360 --> 00:07:38,510

types of cells and particles that we

169

00:07:42,340 --> 00:07:40,370

might be able to just separate these in

170

00:07:44,140 --> 00:07:42,350

in three dimensions or in here in two

171

00:07:49,240 --> 00:07:44,150

dimensions on a on a graph based on

172

00:07:51,760 --> 00:07:49,250

their biophysical properties as Peter

173

00:07:53,860 --> 00:07:51,770

talked about a little bit in his talk

174

00:07:56,080 --> 00:07:53,870

but identifying morphological features

175

00:07:57,909 --> 00:07:56,090

of particulates and I see potentially

176  
00:08:00,790 --> 00:07:57,919  
salty samples is going to require sample

177  
00:08:03,010 --> 00:08:00,800  
processing and along these lines we

178  
00:08:06,730 --> 00:08:03,020  
partnered with the Monterey Bay Aquarium

179  
00:08:08,350 --> 00:08:06,740  
Research Institute to develop a concept

180  
00:08:12,190 --> 00:08:08,360  
for this instrument which would be a

181  
00:08:13,840 --> 00:08:12,200  
sample handling system Ambari has a lot

182  
00:08:16,180 --> 00:08:13,850  
of history and experience in operating

183  
00:08:20,140 --> 00:08:16,190  
remote autonomous platforms in the ocean

184  
00:08:21,940 --> 00:08:20,150  
on earth this is the 2g ESP that was

185  
00:08:25,330 --> 00:08:21,950  
originally funded by a NASA a step

186  
00:08:27,640 --> 00:08:25,340  
project with the the concept they would

187  
00:08:31,120 --> 00:08:27,650  
evolve this design into space based

188  
00:08:33,459 --> 00:08:31,130

operations and the - 2g ESP here is very

189

00:08:35,949 --> 00:08:33,469

large but this is actually a functional

190

00:08:37,959 --> 00:08:35,959

molecular biology platform where it's

191

00:08:40,540 --> 00:08:37,969

not only processing collecting samples

192

00:08:43,240 --> 00:08:40,550

and analyzing them but also doing real

193

00:08:45,730 --> 00:08:43,250

molecular biology DNA extractions PCR

194

00:08:48,280 --> 00:08:45,740

and and remotely communicating that

195

00:08:52,930 --> 00:08:48,290

information back to the lab they use

196

00:08:55,380 --> 00:08:52,940

titanium pucks which are inert for

197

00:08:59,250 --> 00:08:55,390

sample processing and

198

00:09:02,550 --> 00:08:59,260

this system is is again complete and

199

00:09:05,340 --> 00:09:02,560

remotely operated they have a new system

200

00:09:07,590 --> 00:09:05,350

that they have integrated into this

201  
00:09:11,130 --> 00:09:07,600  
submersible which is based on

202  
00:09:14,850 --> 00:09:11,140  
microfluidic cartridges which has a it

203  
00:09:18,090 --> 00:09:14,860  
rotates around and as actuator driven

204  
00:09:20,340 --> 00:09:18,100  
inside the head of the submersible so we

205  
00:09:23,130 --> 00:09:20,350  
have adopted elements from from both of

206  
00:09:26,340 --> 00:09:23,140  
these designs into the concept for

207  
00:09:28,980 --> 00:09:26,350  
Antony in which the titanium puck would

208  
00:09:32,160 --> 00:09:28,990  
be delivered to the instrument the

209  
00:09:36,500 --> 00:09:32,170  
central processing unit here then would

210  
00:09:38,880 --> 00:09:36,510  
accept the sample and process take it

211  
00:09:41,340 --> 00:09:38,890  
pressurize the sample in order to be

212  
00:09:43,680 --> 00:09:41,350  
able to melt it without adding heat or

213  
00:09:45,980 --> 00:09:43,690

very much heat and then it would be

214

00:09:49,020 --> 00:09:45,990

passed through different screen sizes

215

00:09:50,820 --> 00:09:49,030

and then those samples can then be

216

00:09:54,720 --> 00:09:50,830

rotated and passed by an optical

217

00:09:57,210 --> 00:09:54,730

microscope an afm and potentially a

218

00:10:00,240 --> 00:09:57,220

third instrument that is on the lander

219

00:10:03,330 --> 00:10:00,250

this all the design fits in the ten

220

00:10:06,240 --> 00:10:03,340

centimeter by 10 centimeter space that

221

00:10:08,430 --> 00:10:06,250

was described in the PIP so there are

222

00:10:09,990 --> 00:10:08,440

several elements that even though we're

223

00:10:11,280 --> 00:10:10,000

basing this on the heritage design of

224

00:10:13,200 --> 00:10:11,290

what has flown before there are a couple

225

00:10:15,090 --> 00:10:13,210

of upgrades that we can do 20 years

226

00:10:19,650 --> 00:10:15,100

later than when these developments were

227

00:10:22,230 --> 00:10:19,660

made the so piezo resistive probes can

228

00:10:27,840 --> 00:10:22,240

be upgraded as well as the actuator and

229

00:10:30,450 --> 00:10:27,850

fine XY scan range to move the sample we

230

00:10:32,190 --> 00:10:30,460

think calibration targets are needed for

231

00:10:34,920 --> 00:10:32,200

every sample processed so that you can

232

00:10:38,960 --> 00:10:34,930

get tip shape on each sample as well as

233

00:10:41,820 --> 00:10:38,970

the XYZ and mechanical force scales

234

00:10:43,950 --> 00:10:41,830

autonomous impressions are going to

235

00:10:46,350 --> 00:10:43,960

require image analysis some of this can

236

00:10:47,490 --> 00:10:46,360

be adopted from heritage algorithms that

237

00:10:49,800 --> 00:10:47,500

were developed from the Phoenix mission

238

00:10:51,990 --> 00:10:49,810

but we also have ideas for how to

239

00:10:55,080 --> 00:10:52,000

compress the data to preserve the high

240

00:10:57,710 --> 00:10:55,090

information aspects of it as well as for

241

00:11:00,480 --> 00:10:57,720

developing neural networks for particle

242

00:11:02,400 --> 00:11:00,490

classification this is based on work

243

00:11:05,610 --> 00:11:02,410

that was done at Oakridge and published

244

00:11:09,090 --> 00:11:05,620

several years ago with a project

245

00:11:11,400 --> 00:11:09,100

collaborator Steven Jesse we also have

246

00:11:14,640 --> 00:11:11,410

there's preliminary work that has been

247

00:11:17,280 --> 00:11:14,650

done with where you can use the AFM to

248

00:11:19,200 --> 00:11:17,290

do correlative mass spectrometry with

249

00:11:21,630 --> 00:11:19,210

particles by heating up the tip of the

250

00:11:23,760 --> 00:11:21,640

AFM you can volatilize the sample and

251  
00:11:25,260 --> 00:11:23,770  
bring it into a mass spectrometer and so

252  
00:11:26,700 --> 00:11:25,270  
this could potentially really give us a

253  
00:11:31,170 --> 00:11:26,710  
lot of information about particles being

254  
00:11:33,180 --> 00:11:31,180  
observed the advantages of Antony then

255  
00:11:35,430 --> 00:11:33,190  
are that this is a system as we've

256  
00:11:37,290 --> 00:11:35,440  
designed it has flexible architecture we

257  
00:11:39,680 --> 00:11:37,300  
can partner with other instruments it's

258  
00:11:42,060 --> 00:11:39,690  
integrated sample preparation system

259  
00:11:44,130 --> 00:11:42,070  
affords us to have high resolution image

260  
00:11:48,900 --> 00:11:44,140  
and proper image and property

261  
00:11:51,000 --> 00:11:48,910  
determinations the topographic maps will

262  
00:11:52,710 --> 00:11:51,010  
be useful information regardless of

263  
00:11:55,260 --> 00:11:52,720

whether the particles are biogenic or

264

00:11:57,750 --> 00:11:55,270

abiotic in nature and the Nano

265

00:12:00,780 --> 00:11:57,760

mechanical forces can potentially help

266

00:12:02,850 --> 00:12:00,790

us inform between life-forms and and

267

00:12:04,740 --> 00:12:02,860

those that are that are not life in

268

00:12:07,440 --> 00:12:04,750

terms of stiffness adhesion dissipation

269

00:12:08,730 --> 00:12:07,450

of energy and even in the absence of

270

00:12:10,290 --> 00:12:08,740

life I think that this kind of a

271

00:12:12,540 --> 00:12:10,300

platform can yield invaluable

272

00:12:14,160 --> 00:12:12,550

information about microscope materials

273

00:12:18,480 --> 00:12:14,170

and features in the near subsurface of

274

00:12:21,120 --> 00:12:18,490

Europa the the team here is large and

275

00:12:23,220 --> 00:12:21,130

great and I have learned a ton from

276

00:12:25,680 --> 00:12:23,230

working with all of them Roger approach

277

00:12:30,390 --> 00:12:25,690

is sort of the mastermind behind the AFM

278

00:12:33,630 --> 00:12:30,400

design crystal and is the chief and

279

00:12:37,110 --> 00:12:33,640

president of Ambari and with Jim birch

280

00:12:41,450 --> 00:12:37,120

and Doug market have designed the Europa

281

00:12:51,930 --> 00:12:41,460

generation ESP thank you

282

00:12:57,010 --> 00:12:55,480

yeah given that you're looking at very

283

00:12:58,720 --> 00:12:57,020

tiny scales with AFM how do you deal

284

00:13:00,790 --> 00:12:58,730

with searching for things that are

285

00:13:02,440 --> 00:13:00,800

extremely low concentration that you're

286

00:13:03,490 --> 00:13:02,450

likely to see 100 per milliliter or

287

00:13:06,840 --> 00:13:03,500

something like that see if the search

288

00:13:10,120 --> 00:13:06,850

throw a big haystack refer so we have an

289

00:13:12,160 --> 00:13:10,130

the the concept is that we would

290

00:13:15,190 --> 00:13:12,170

concentrate the sample onto a pretty

291

00:13:19,030 --> 00:13:15,200

into a pretty small surface but also

292

00:13:21,070 --> 00:13:19,040

that we would first inspect the the

293

00:13:23,590 --> 00:13:21,080

sample with an optical microscope and

294

00:13:27,370 --> 00:13:23,600

you have a larger scale and then be able

295

00:13:30,640 --> 00:13:27,380

to identify the high entropy areas of

296

00:13:32,560 --> 00:13:30,650

that image and then localize the AFM to

297

00:13:34,450 --> 00:13:32,570

where to go do the work and you can even

298

00:13:36,460 --> 00:13:34,460

still do nested scans like they've done

299

00:13:38,500 --> 00:13:36,470

on both of the other missions with AFM

300

00:13:40,480 --> 00:13:38,510

so you can start at larger and then go

301  
00:13:46,510 --> 00:13:40,490  
in and a higher resolution at a smaller

302  
00:13:48,040 --> 00:13:46,520  
area another way of distinguishing

303  
00:13:50,380 --> 00:13:48,050  
between whether you're looking at

304  
00:13:52,930 --> 00:13:50,390  
inorganic material or organic or bio

305  
00:13:55,120 --> 00:13:52,940  
material would be to do a tip enhance

306  
00:13:57,850 --> 00:13:55,130  
spectroscopy such as tip enhanced Raman

307  
00:14:00,400 --> 00:13:57,860  
or chip enhanced infrared and I know

308  
00:14:02,640 --> 00:14:00,410  
it's probably at a lower TRL but what do

309  
00:14:05,470 --> 00:14:02,650  
you think about the possibilities of

310  
00:14:07,420 --> 00:14:05,480  
eventually moving to such an enhancement

311  
00:14:10,330 --> 00:14:07,430  
of the technology yeah I think that's a

312  
00:14:13,570 --> 00:14:10,340  
super super super interesting an area to

313  
00:14:16,600 --> 00:14:13,580

develop for the future and to get that

314

00:14:18,790 --> 00:14:16,610

kind of instrumentation into the sort of