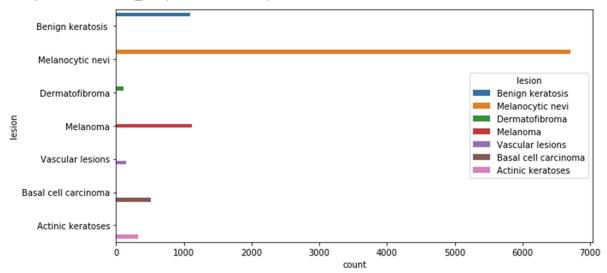
# Skin Disease Classification

```
from fastai import *
from fastai.vision import *
from fastai.callbacks.hooks import *
import numpy as np
import pandas as pd
import seaborn as sns
import matplotlib.pyplot as plt
from sklearn.metrics import auc,roc_curve
import os
print(os.listdir("../input"))
['hmnist_28_28_RGB.csv', 'hmnist_28_28_L.csv', 'ham10000_images_part_2', 'ham1
0000_images_part_1', 'hmnist_8_8_RGB.csv', 'HAM10000_metadata.csv', 'hmnist_8_
8 L.csv']
Exploratory Data Analysis
                                                                              In [2]:
# Paths and roots to the important files
path='../input/'
csv_file='../input/HAM10000_metadata.csv'
                                                                              In [3]:
df=pd.read_csv(csv_file).set_index('image_id')
df.head()
                                                                               ut[3]:
                                                                              In [4]:
# Categories of the diferent diseases
lesion_type_dict = {
    'nv': 'Melanocytic nevi',
    'mel': 'Melanoma',
    'bkl': 'Benign keratosis ',
    'bcc': 'Basal cell carcinoma',
    'akiec': 'Actinic keratoses',
'vasc': 'Vascular lesions',
    'df': 'Dermatofibroma'
}
                                                                              In [5]:
df.dx=df.dx.astype('category',copy=True)
df['labels']=df.dx.cat.codes # Convert the Labels to numbers
df['lesion'] = df.dx.map(lesion_type_dict)
df.head()
                                                                              Out[5]:
                                                                              In [6]:
print(df.lesion.value_counts())
Melanocytic nevi
                          6705
Melanoma
                          1113
Benign keratosis
                          1099
```

```
Basal cell carcinoma
                          514
Actinic keratoses
                          327
Vascular lesions
                          142
Dermatofibroma
                          115
Name: lesion, dtype: int64
                                                                           In [7]:
df.loc['ISIC_0027419','lesion']
                                                                           Out[7]:
'Benign keratosis '
Countplot
```

Here we notice tha we have data imbalance

```
In [8]:
fig, ax1 = plt.subplots(1, 1, figsize = (10, 5))
sns.countplot(y='lesion',data=df, hue="lesion",ax=ax1)
                                                                            Out[8]:
<matplotlib.axes. subplots.AxesSubplot at 0x7ff27a102550>
```



### Dataset

```
In [9]:
class CustomImageItemList(ImageItemList):
    def custom_label(self,df, **kwargs)->'LabelList':
        """Custom Labels from path"""
        file_names=np.vectorize(lambda files: str(files).split('/')[-1][:-4])
        get_labels=lambda x: df.loc[x,'lesion']
        #self.items is an np array of PosixPath objects with each image path
        labels= get labels(file names(self.items))
        y = CategoryList(items=labels)
        res = self._label_list(x=self,y=y)
        return res
                                                                            In [10]:
def get_data(bs, size):
    train_ds = (CustomImageItemList.from_folder('.../input', extensions='.jpg')
                    .random_split_by_pct(0.15)
                    .custom_label(df)
                    .transform(tfms=get_transforms(flip_vert=True), size=size)
                    .databunch(num_workers=2, bs=bs)
```

.normalize(imagenet\_stats)) return train\_ds In [11]: data=get\_data(16,224) In [12]: data.classes=list(np.unique(df.lesion)) data.c= len(np.unique(df.lesion)) In [13]: data.show\_batch(rows=3) Melanocytic nevi Melanocytic nevi Melanocytic nevi Benign keratosis Melanoma Melanoma Melanocytic nevi Melanocytic nevi Melanocytic nevi

## Model ResNet50

In [14]:
learner=create\_cnn(data,models.resnet50,metrics=[accuracy], model\_dir="/tmp/model/
")

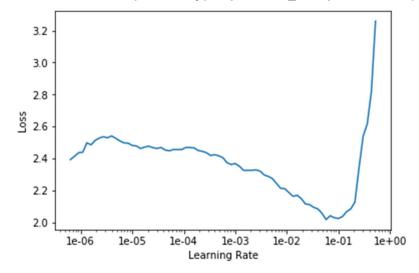
Downloading: "https://download.pytorch.org/models/resnet50-19c8e357.pth" to /t mp/.torch/models/resnet50-19c8e357.pth

100%| 102502400/102502400 [00:03<00:00, 26653549.20it/s]

In [16]:

learner.lr\_find()
learner.recorder.plot()

LR Finder is complete, type {learner\_name}.recorder.plot() to see the graph.



In [17]:

learner.fit\_one\_cycle(10, 3e-3)

60.00% [6/10 31:58<21:19]

e p o c h	tr ai n _1 o ss	va li d _l os s	a c c u r a c y
1	0. 8 4 1 1 9 4	0. 7 0 0 3 7 4	0 7 6 6 9 7 7
2	0. 7 3 7 2 5 6	0. 6 2 5 3 0 5	0 7 7 2 3 0 4

3	0. 7 3 8 6 8 0	0. 6 0 0 1 8 3	0 7 9 0 9 4 5
4	0. 6 2 4 4 8 9	0. 6 2 3 8 4 3	0 7 7 1 6 3 8
5	0. 5 7 5 1 6 1	0. 5 8 6 3 7 2	0 7 9 5 6 0 6
6	0. 4 7 6 0 6 6	0. 5 0 2 8 0 9	0 8 2 2 9 0 3

# 97.74% [520/532 04:32<00:06 0.5225]

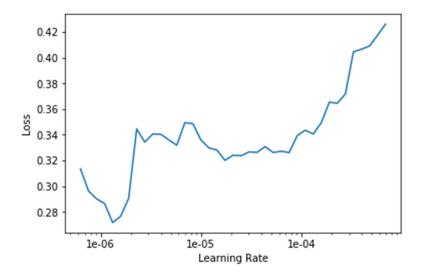
learner.unfreeze()

In [19]:

In [18]:

learner.lr\_find()
learner.recorder.plot()

LR Finder is complete, type {learner\_name}.recorder.plot() to see the graph.



lr=1e-6
learner.fit\_one\_cycle(3, slice(3\*lr,10\*lr))

Total time: 16:47

, 10. <del>4</del> 1			
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			a
e	tr	va	c
	ai	li	c c
P	n	d	u
O	1	1	r
p o c h	tr ai n _1 o ss	va li d _l os s	а
h	55	6	а с у
	88	5	
			У
	0	0	0
	0.	0.	
	3	4	8
	5	0	4
1	3	7	4
	8	8	6
	0. 3 5 3 8 8	0. 4 0 7 8 6 6	8
	0	6	7
	U	O	8 4 6 8 7 1
			0
	0.	0.	0
	3	3	•
	0	0	8
2	9	2	5
2	8	3	1
	0	0	5
	6	2	3
	0. 3 9 8 0 6 7	0. 3 9 3 0 2	8 5 1 5 3 1
			1
	0.	0.	
	3	3	0
	7	9	Č
3	, 4	5	
5	2	2	8
	3	2	5
	0. 3 7 4 3 8 3	0. 3 9 5 2 2	8 5 2 1
	3	9	1

In [20]:

```
9
```

learner.save('stage-1')

In [21]:

interp = ClassificationInterpretation.from\_learner(learner)

In [22]:

interp.plot\_confusion\_matrix(figsize=(10,8))

In [23]:

	,	Confusion matrix						
	Actinic keratoses -	25	2	9	1	8	1	0
В	asal cell carcinoma -	0	69	4	0	8	1	0
Actual	Benign keratosis -	8	2	105	1	42	15	0
	Dermatofibroma -	2	0	1	7	3	2	0
	Melanocytic nevi -	1	2	5	0	965	16	0
	Melanoma -	1	4	8	2	69	89	1
	Vascular lesions -	0	0	0	0	3	0	20
	'	Actinic keratoses -	Basal cell carcinoma -	Benign keratosis -	Dermatofibroma -	Melanocytic nevi –	Melanoma -	Vascular lesions -

In [24]:

interp.most\_confused()

Out[24]:

```
[('Melanoma', 'Melanocytic nevi', 69),
  ('Benign keratosis ', 'Melanocytic nevi', 42),
  ('Melanocytic nevi', 'Melanoma', 16),
  ('Benign keratosis ', 'Melanoma', 15),
  ('Actinic keratoses', 'Benign keratosis ', 9),
  ('Actinic keratoses', 'Melanocytic nevi', 8),
```

```
('Basal cell carcinoma', 'Melanocytic nevi', 8),
 ('Benign keratosis ', 'Actinic keratoses', 8),
 ('Melanoma', 'Benign keratosis ', 8),
 ('Melanocytic nevi', 'Benign keratosis ', 5),
 ('Basal cell carcinoma', 'Benign keratosis', 4),
 ('Melanoma', 'Basal cell carcinoma', 4),
 ('Dermatofibroma', 'Melanocytic nevi', 3),
 ('Vascular lesions', 'Melanocytic nevi', 3),
 ('Actinic keratoses', 'Basal cell carcinoma', 2), ('Benign keratosis', 'Basal cell carcinoma', 2),
 ('Dermatofibroma', 'Actinic keratoses', 2), ('Dermatofibroma', 'Melanoma', 2),
 ('Melanocytic nevi', 'Basal cell carcinoma', 2),
 ('Melanoma', 'Dermatofibroma', 2)]
Inference
                                                                             In [25]:
pred_data=get_data(16,224)
                                                                             In [26]:
pred data.classes=list(np.unique(df.lesion))
pred_data.c= len(np.unique(df.lesion))
                                                                             In [27]:
pred_data.single_from_classes(path, pred_data.classes)
/opt/conda/lib/python3.6/site-packages/fastai/data block.py:388: UserWarning:
Your training set is empty. Is this is by design, pass `ignore_empty=True` to
remove this warning.
  warn("Your training set is empty. Is this is by design, pass `ignore empty=T
rue` to remove this warning.")
/opt/conda/lib/python3.6/site-packages/fastai/data_block.py:391: UserWarning:
Your validation set is empty. Is this is by design, use `no_split()`
                  or pass `ignore_empty=True` when labelling to remove this war
ning.
  or pass `ignore_empty=True` when labelling to remove this warning.""")
                                                                             Out[27]:
ImageDataBunch;
Train: LabelList
y: CategoryList (0 items)
[]...
Path: ../input
x: ImageItemList (0 items)
[]...
Path: ../input;
Valid: LabelList
y: CategoryList (0 items)
[]...
Path: ../input
x: ImageItemList (0 items)
Path: ../input;
Test: None
```

In [28]:

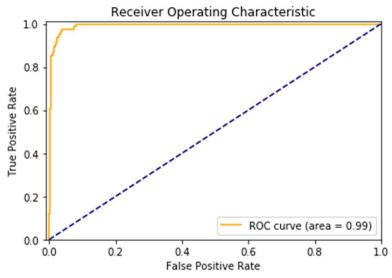
```
predictor = create_cnn(pred_data, models.resnet50, model_dir="/tmp/model/").load('
stage-1')
                                                                            In [29]:
img = open_image('../input/ham10000_images_part_2/ISIC_0029886.jpg')
img
                                                                            Out[29]:
                                                                            In [30]:
pred_class,pred_idx,outputs = predictor.predict(img)
pred_class
                                                                            Out[30]:
Category Melanocytic nevi
Predictions
                                                                            In [31]:
# Predictions of the validation data
preds_val, y_val=learner.get_preds()
Roc Curve
With the ROC curve we will mesuare how good it's our model
                                                                            In [32]:
# ROC curve
fpr, tpr, thresholds = roc_curve(y_val.numpy(), preds_val.numpy()[:,1], pos_label=
1)
# ROC area
pred_score = auc(fpr, tpr)
```

```
print(f'ROC area is {pred_score}')
ROC area is 0.9935846788045345

In [33]:
plt.figure()
plt.plot(fpr, tpr, color='orange', label='ROC curve (area = %0.2f)' % pred_score)
plt.plot([0, 1], [0, 1], color='navy', linestyle='--')
plt.xlim([-0.01, 1.0])
plt.ylim([0.0, 1.01])
plt.xlabel('False Positive Rate')
plt.ylabel('True Positive Rate')
plt.title('Receiver Operating Characteristic')
plt.legend(loc="lower right")

Out[33]:
```

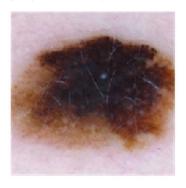
<matplotlib.legend.Legend at 0x7ff248dac860>



# Heatmap

x,y = data.valid\_ds[2]
x.show()
data.valid\_ds.y[2]

Category Melanocytic nevi



In [35]:

In [34]:

Out[34]:

def heatMap(x,y,data, learner, size=(0,224,224,0)):
 """HeatMap"""

```
# Evaluation mode
m=learner.model.eval()
# Denormalize the image
xb_{,-} = data.one_item(x)
xb_im = Image(data.denorm(xb)[0])
xb = xb.cuda()
# hook the activations
with hook_output(m[0]) as hook_a:
    with hook_output(m[0], grad=True) as hook_g:
        preds = m(xb)
        preds[0,int(y)].backward()
# Activations
acts=hook_a.stored[0].cpu()
# Avg of the activations
avg_acts=acts.mean(0)
# Show HeatMap
_,ax = plt.subplots()
xb_im.show(ax)
ax.imshow(avg_acts, alpha=0.6, extent=size,
          interpolation='bilinear', cmap='magma')
```

In [36]:

heatMap(x,y,pred\_data,learner)

