



UDACITY

Introduction to Generative AI with AWS Project Documentation Report

Question	Your answer
Step 2: Domain Choice What domain did you choose to fine-tune the Meta Llama 2 7B model on? Choices: 1. Financial 2. Healthcare 3. IT	The domain I choose to fine-tune the Meta Llama 2 7B model on is: 2. Healthcare

Step 3: Model Evaluation Section

What was the response of the model to your domainspecific input in the model_evaluation.ipynb file?

Session Setup

```
import sagemaker, boto3, json
from sagemaker.session import Session

sagemaker_session = Session()
aws_role = sagemaker_session.get_caller_identity_arn()
aws_region = boto3.Session().region_name
sess = sagemaker.Session()
print(aws_role)
print(aws_region)
print(sess)

sagemaker.config INFO - Not applying SDK defaults from location: /etc/xdg/sagemaker/config.yaml
sarn:aws:iam::508781339756:role/service-role/SageMaker-udacitySagemakerRole
us-east-1
<sagemaker.session.Session object at 0x7f8a778d3970>
```

(model_id, model_version,) = ("meta-textgeneration-llama-2-7b","2.*",)

Model Definition and Deployment

```
from sagemaker.jumpstart.model import JumpStartModel
model = JumpStartModel(model_id=model_id, model_version=model_version, instance_type="ml.g5.2xlarge")
predictor = model.deploy()

For forward compatibility, pin to model_version='2.*' in your JumpStartModel or JumpStartEstimator definitions. Note that major version upgrades may have different EULA acceptance terms and input/output signatures.
Using vulnerable JumpStart model 'meta-textgeneration-llama-2-7b' and version '2.1.8'.
Using model 'meta-textgeneration-llama-2-7b' with wildcard version identifier '2.*'. You can pin to version '2.1.8' for more st able results. Note that models may have different input/output signatures after a major version upgrade.
```

• Model Evaluation

Inputs and Outputs: {

"Myeloid neoplasms and acute leukemias derive from"

Myeloid neoplasms and acute leukemias derive from

> a common progenitor and share a number of common genetic lesions.

Malignant myeloid neoplasms are a group of hematologic malignancies that includes myelodysplastic syndromes (MDS), myeloprolife rative neoplasms

.____

"Genomic characterization is essential for"

Genomic characterization is essential for

- > understanding the mechanisms of adaptation and speciation in this group.
- AB Background: The genus Sarcocystis contains 30 species that infect a wide range of vertebrates. These organisms are obligat e intracellular parasites of the phylum Cnidaria

• "Certain germline disorders may be associated with"

Certain germline disorders may be associated with

> an increased risk of developing breast cancer.

Genetic testing may be considered for people who have a family history of breast cancer.

Risk of breast cancer in BRCA-positive individuals

BRCA1 and BRCA2 are two genes that are responsible for repairing damaged DNA in cells. Mut

• "In contrast to targeted approaches, genome-wide sequencing"

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Risk of breast cancer in BRCA-positive individuals

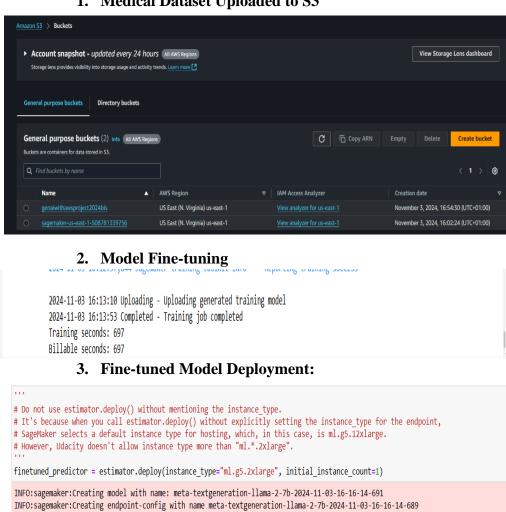
BRCA1 and BRCA2 are two genes that are responsible for repairing damaged DNA in cells. Mut

}

Step 4: Fine-Tuning Section

After fine-tuning the model, what was the response of the model to your domainspecific input in the model finetuning.ipynb file?

1. Medical Dataset Uploaded to S3



INFO:sagemaker:Creating endpoint with name meta-textgeneration-llama-2-7b-2024-11-03-16-16-14-689

4. Fine-tuned Model Evaluation:

Inputs and Outputs: {

"Myeloid neoplasms and acute leukemias derive from"

Myeloid neoplasms and acute leukemias derive from

> [{'generated text': ' a common myeloid progenitor. The genetic and epigenetic events that lead to the acquisition of the leuk emic phenotype are still poorly understood. We have identified a new transcription factor, TAL1/SCL, that is essential for norm al development of the hemat'}]

"Genomic characterization is essential for"

Genomic characterization is essential for

> [{'generated_text': ' the identification of novel genetic variations in diseases. Genetic variations are a major source of genetic diversity and are critical in understanding the evolutionary processes that shaped human genetic diversity.\nThe Human Genome Project (HGP) was a 13-year project that was completed in '}]

"Certain germline disorders may be associated with"

Certain germline disorders may be associated with

- > [{'generated_text': ' an increased risk of cancer, but it is unclear whether the risk is related to the disorder itself or to the treatment that patients with the disorder receive.\nIn a study published in the Journal of Clinical Oncology, researchers a t Dana-Farber Cancer Institute and the Broad Institute'}]
 - "In contrast to targeted approaches, genome-wide sequencing"

In contrast to targeted approaches, genome-wide sequencing

> [{'generated_text': ' is the only way to identify the entire spectrum of genetic variations that contribute to the risk of di sease. It also provides insights into the biological mechanisms underlying the disease and the potential for novel therapeutic targets.\nSeveral large-scale sequencing initiatives have been launched to generate the'}]

}

5. Fine-tuned model weights in 3s

