



Generating Electronic Health Records: an Investigation on Gender-Medicine and Rare Diseases

vorgelegt von

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dem Fachbereich VI

der Beuth Hochschule für Technik Berlin vorgelegte Masterarbeit
zur Erlangung des akademischen Grades

Master of Science (M.Sc.)

im Studiengang

Medieninformatik

Tag der Abgabe 6. Januar 2020

Gutachter

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Abstract

Gaining access to electronic health record data comes with numerous hardships, due to hindrances like, e.g., the risk of hurting a patient's privacy. However, EHR data plays a central role in medical research in many fields, e.g., investigating diseases, especially rare ones, preventative measures and developing medical software. The generation of synthetic data could solve this problem, if its quality is similar to real data. In our work we examine a Generative Adversarial Network (GAN) called medical Generative Adversarial Network (medGAN) that was proposed by (Choi et al., 2018). We compare the performance of the original model to a model that is trained with female and male patients separately. We show that the original model is superior to the newly introduced model. Further, we show that the original model is able to generate samples with rare diseases and that these samples can not be distinguished from real data by a medical doctor.

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Chapter 1

Introduction

1.1 Artificial intelligence in health care

Artificial intelligence (AI) tries to simulate human intelligence. Its applications are numerous and can also be found in the health care sector. A possible use case could be an assisting AI system, that provides up-to-date information about clinical practices to physicians and reduces errors, in a therapeutic and diagnostic context. (Jiang et al., 2017). Electronic health records (EHR) are the digital version of a patient's medical healthcare information over a lifetime. AI can extract and analyze patient data from such records and give alerts for possible health risks and, e.g., make predictions about diagnoses and the possible length of stay in the hospital. (Neill, 2013). Practically such a system could analyze a patient's data already during reception and provide a first diagnosis. Another exemplary use case is the application of AI in diagnosis of breast cancer detection. (Übeyli, 2007) Despite the many use cases, publicly available EHR data for research is rare. This is mainly to protect each patient's privacy. However, this especially poses a problem to the availability of data about rare diseases, leading to a lack of information and slowing down research. (Bremond-Gignac et al., 2015)

1.2 Generative Adversarial Networks

Since their introduction in 2014 by (Goodfellow et al., 2014), Generative Adversarial Networks have gained a substantial extent of attention. This is mainly due to their ability to augment synthetic image data is not distinguishable from real data. This quality proofs to be useful in the medical domain and to its applications count for example, the augmentation of medical images of liver lesions. After training with these images, the performance of a convolutional Neural Network that classifies liver lesions improved. (Frid-Adar et al., 2018) Explained briefly, a GAN consists of two neural networks, that are playing against each other. One creates fake samples, while the other one tries to distinguish these samples from real ones. During this process, both networks train each other. (Goodfellow et al., 2014) We will explain its functionality in detail in the following Chapter 2.

1.3 Goal

In this work, we aim to generate EHRs. For generation, we use an approach by (Choi et al., 2018), which produces high-dimensional discrete variables (e.g., binary and count features) using an altered Generative Adversarial Network, called medical Generative Adversarial Network (medGAN). As input dataset, we use an EHR dataset called MIMIC-III. In contrast to the model presented in (Choi et al., 2018), we additionally split the input dataset by gender. We want to investigate whether the samples the medical Generative Adversarial Network (medGAN)

Patient ID	272	414	780
A	0	0	1
B	1	1	0
C	0	0	0

Table 1.1: Example binary data entries with 3 digit ICD-9 codes

Patient ID	488.11	516.30	996.49
A	0	5	0
B	1	0	9
C	0	2	0

Table 1.2: Example count data entries with 5 digit ICD-9 codes

generates will be more realistic, by splitting the input data by gender. Further, we want to investigate whether medGAN can generate realistic samples with rare diseases.

1.4 Method

In our research, we use the same model as already mentioned in (Choi et al., 2018). We change the input training data by splitting up the data by gender. Subsequently, we train the network, using both, binary and count variables. We train the model with the full dataset and afterwards with the dataset of each gender. First, we quantitatively evaluate our measurements, using dimension-wise probability for binary output and dimension-wise average for count output. In our qualitative evaluation, we investigate our measurements for a correlation of Cardiovascular Disease (CVD) and Diabetes Mellitus (DM). Further, we will evaluate our measurements with the help of a medical doctor.

1.4.1 MedGAN

A central aspect of this work will be the application of the medical Generative Adversarial Network (medGAN), which is proposed by (Choi et al., 2018). It is a GAN, able to generate realistic synthetic patient records, that achieve comparable results to real data when used for predictive modeling tasks. (Choi et al., 2018) proof in their work that medGAN outperforms classical machine learning algorithms like Linear Regression, Random Forests , and Support Vector Machine. In Chapter 2, the motivation for creating medGAN will be explained. In Chapter 4, we will provide detailed information on the architecture of medGAN and will elaborate on recent improvements on medGAN.

1.4.2 Generating binary and count features

For our experiments, we generated both, records with binary and count variables. Records with binary data will show whether an ICD-9 code occurs or not, **Table 1.1** shows exemplary data. Records with count data will show how many times each ICD-9 code occurs for each patient, as depicted in **Table 1.2**

1.4.3 Privacy Risk

The risk of hurting a patient's privacy is one of the main reasons why Electronic Health Data is not publicly available. A commonly used method for providing access to patient data for researchers is de-identification, used, for example, on the MIMIC-III dataset. In this process, data gets cleansed and shifted, meaning that names, telephone numbers, and other personal information gets removed and dates altered. (Johnson et al., 2016) However, (Choi et al., 2018) explains that de-identification does not eliminate the risk of harming a patient's privacy, because identification of an individual is still possible, if an attacker has pieces of information about, e.g. diagnoses, lab tests, demographics, genomic variants or visits from other health care providers.

On the contrary, as the synthetic data from medGAN is artificially created, there is no direct connection between the real and the synthetic samples. Therefore identifying patients from the original dataset should, intuitively, not be possible if the artificially created samples get stolen because of a security breach or due to other reasons. In (Choi et al., 2018) assesses the privacy risk for the case that the synthetic samples get compromised and comes to the conclusion that a potential attacker poses no risk of breaching privacy, except for when he already has significant knowledge about various features of the target patient.

1.5 Outline

This work is divided into 6 chapters, starting with 'Introduction.', where we learned about the application of AI in health care, Generative Adversarial Networks and the extended version, medGAN, that we will apply in our work. In the next chapter 'Related Work', we will dive deeper into the topic by explaining all the essential concepts that find application in our work. In Chapter 3, 'Data Analysis', we will analyze the given MIMIC-III dataset. Subsequently, in Chapter 4, 'Electronic Medical Record Generation', we will perform our experiments and generate the EHR data. Afterwards, in 'Experimental Evaluation', we will assess the success of our experiments. Lastly, in Chapter 6 'Conclusion and Outlook', we will conclude whether our hypotheses held up during our experiments and will give an outlook on possible future work.

1.6 Summary

This chapter explained the motivation for using AI in the medical sector and the reason for scarcity of data in this field. For giving an insight into our method, we first provided a short definition of Generative Adversarial Networks and medGAN. We will explain GANs more thoroughly in the following Chapter 2, medGAN in Chapter 4. This chapter also depicts two examples of the representation of the generated data. We also addressed (Choi et al., 2018)'s work on investigating the privacy risk that synthetic samples pose.

Chapter 2

Related Work

2.1 Abstract

The following section will elaborate on the basic concepts that are needed to generate synthetic electronic health record data. First, we explain the concept of a Generative Adversarial Network. Further, we will define what exactly an Electronic Health record is. Subsequently, we will introduce the definition of Gender Medicine and Rare Diseases and their role in this work. Lastly, we will give information on medGAN and thoroughly explain its characteristics and architecture in Chapter 4.

2.2 Basic Concepts

2.2.1 Generative Adversarial Networks

(Goodfellow et al., 2014) proposed a new framework for generative models that learns the patterns in a given dataset, and generates new data that plausibly could originate from the original dataset. The model (Goodfellow et al., 2014) proposes, corresponds to a two-player minimax game: in which two independent neural networks train simultaneously. The generator **G** tries to learn the distribution of the given dataset and the discriminator **D** aims to distinguish between the data from the training set and the 'fake' by **G**. **G** generates the data from the learned distribution and takes random noise as input. In this game, **G** has the goal to maximize the probability of **D** making a mistake.

The two-player minimax game, as shown in (Goodfellow et al., 2014), can be described by the following value function:

$$\min_G \max_D V(D, G) = \mathbb{E}_{x \sim P_{data(x)}} [\log D(x)] + \mathbb{E}_{z \sim P_z(z)} [\log(1 - D(G(z)))] \quad (2.1)$$

For a better understanding of the process of the model, the following analogy can be helpful: "The generative model can be thought of as analogous to a team of counterfeiters, trying to produce fake currency and use it without detection, while the discriminative model is analogous to the police, trying to detect the counterfeit currency. Competition in this game drives both teams to improve their methods until the counterfeits are indistinguishable from the genuine articles." (Goodfellow et al., 2014)

Mainly due to their performance in generating synthetic image data that is not distinguishable from real images, Generative Adversarial Networks gained a considerable amount of attention since their introduction in 2014.

2.2.2 Electronic Health Records

The Healthcare Information and Management Systems Society defines an Electronic Health Record as "a longitudinal electronic record of patient health information generated by one or more encounters in any care delivery setting. Included in this information are patient demographics, progress notes, problems, medications, vital signs, past medical history, immunizations, laboratory data and radiology reports."¹ The term 'Electronic Medical Record' often is used synonymously, but a distinction between both terms has to be made. The Office of the National Coordinator for Health Information Technology defines an EMR as the digital version of a patient's paper record. It contains time tracking data, health parameters (for example blood values), and information about checkups. They are maintained for each practice. It's intended usage is for the hospital or doctor itself and it is not meant to be shared.² EHRs, on the other hand are meant to give a perspective on the overall health of each patient and to be shared for research, other healthcare providers and even the patients themselves. They contain a patient's life-long medical history, medications, diagnoses, treatment plans, vaccination dates, allergies, images, laboratory data (e.g. blood values) and test results.³ A problem with EHR data, is that interoperability is not always given due to missing standardization. (Johnson et al., 2016) In our work we will focus on generating diagnoses codes. MedGAN is also able to generate medication and procedure codes.

2.2.3 Gender Medicine

Medical research is dominated by the male gender, meaning that women are heavily underrepresented or sometimes even excluded from research studies not only in animal studies but also in human trials. (Baggio et al., 2013) But diseases differ between genders, not only in terms of prevention but also in clinical signs and therapeutic approach (Baggio et al., 2013) Sex-differences can also be found in the correlation of diseases. (Kautzky-Willer et al., 2010) shows in his research that "Sex-specific differences appear particularly relevant in the management of type 2 diabetes mellitus (T2DM), with women experiencing greater increases in cardiovascular morbidity and mortality than do men." (Kautzky-Willer et al., 2010) Gender, however, does not only include sex but also lifestyle-related diseases, stress, and behaviour, like, for example, regarding help-seeking actions. While we can not take the socio-cultural aspect of gender into account, the differences in sex are applicable. As "cardiovascular disease is the leading cause of death of both men and women" (Arain et al., 2009), we can find numerous occurrences in the MIMIC-III dataset. This allows us to investigate the co-occurrences of *Diabetes Mellitus (DM)* and *Cardiovascular Disease (CVD)*. Originally MedGAN was trained with male and female patients simultaneously, making no difference between them. In this work, we are separating the dataset in order to introduce a distinction between genders.

The previously mentioned gender-differences lead us to one of our hypotheses, that by training the network with samples of each gender separately, its performance improves and it is able to generate patients with gender-specific correlated diseases that seem realistic to a medical doctor.

2.2.4 Rare diseases

In Europe, a rare disease, also known as orphan disease, is defined as such when there are no more than five occurrences in 10.000 people. As defined by Orphanet, there are roughly 6,000 to 7,000 rare disorders, which include diseases, syndromes and anomalies. The characteristic of a disease is an altered state of health, presented "as a unique pattern of symptoms with a single

¹<https://www.himss.org/electronic-health-records> (01-17-2020)

²<https://www.healthit.gov/buzz-blog/electronic-health-and-medical-records/emr-vs-ehr-difference> (01-20-2020)

³See footnote 2

treatment".⁴ The majority of rare diseases have a genetic origin, but can also originate from an infection or due to unknown reasons.⁵ Specific codes for orphan diseases are only scarce in international classification systems like ICD-9 or ICD-10, which is used in this work, resulting in a deficit of information for researchers regarding orphan diseases. This problem finds its cause in the lack of research and a lack of awareness for rare diseases until recently, also resulting in patients without a diagnosis and unidentified illnesses.⁶ Therefore Orphanet developed a nomenclature, in which each rare disease receives a unique ORPHANumber. The Orphanet classification system can be accessed through their website⁷. This system increases the visibility of rare diseases and interoperability among health and research information systems.⁸

2.2.5 medGAN

The wide adoption of the electronic health record system by healthcare organizations (HCOs) promises advances in analyzing patient data and computational health. The records, however, are not easily accessible for researchers. Due to the fact that EHR data consists of personal and sensitive information, access is restricted in order to not induce a privacy risk. Further, to minimize the risk of data misuse, access to such data is regulated by the HCOs. (Choi et al., 2018) (Even researches that are in direct cooperation with a hospital, do not get access to patient data.) As (Choi et al., 2018) states, the process of getting access is a time-consuming act and without guarantee to gain it. The restricted access slows down research.)."⁹ (Choi et al., 2018) further describes, that despite processes like date-shifting and alteration and randomization of personal information, patients are still prone to re-identification. "To generate synthetic data (McLachlan et al., 2016; Buczak et al., 2010; Lombardo and Moniz, 2008)" (Choi et al., 2018), also does not solve the problem, as the resulting data is not sufficiently realistic to be used as training data. To overcome the limitations and risks of the above-stated methods, (Choi et al., 2018) introduced medGAN, which implements a Generative Adversarial Network that leverages an autoencoder to overcome its limitations: the GAN generates distributed representations of patient records, while the autoencoder decodes them into actual discrete records. This principle and the detailed architecture of medGAN will be further explained in the section 'Electronic Medical Record Generation'.

2.3 Summary

This chapter provided an overview of fundamental concepts that need to be understood to generate electronic medical records and to evaluate our measurements. It gave an insight into Generative Adversarial Networks, and how they will be used in this work and into the definition of Electronic Health Records and how they are distinguishable from Electronic Medical Records. The chapter proceeded with an explanation of gender-medicine and rare diseases and their role for the evaluation process in Chapter 5. It concluded in a detailed explanation of medGAN, that will be used in Chapter 4. In the following Chapter 3, we will investigate the MIMIC-III dataset.

⁴https://www.orpha.net/consor/cgi-bin/Education_AboutOrphanet.php (01-23-2020)

⁵See footnote 4

⁶See footnote 4

⁷<https://www.orpha.net/>

⁸See footnote 4

Chapter 3

Dataset and Analysis

3.1 Abstract

In this Chapter, we will learn about the MIMIC-III dataset, how it is structured and what contents it includes. Afterwards we will investigate the dataset for the most frequent diagnosis codes and give an overview about the correlation of diseases in the dataset. Further, we will examine the co-occurrence of Diabetes Mellitus (DM) and Cardiovascular Disease (CVD) for female and male patients.

3.2 Dataset

The MIMIC-III ('Medical Information Mart for Intensive Care') dataset is one of the biggest publicly available in the medical domain. It is developed and maintained by the MIT lab for computational physiology. It comprises de-identified health data of approximately 45,000 patients admitted to critical care units at Beth Israel Deaconess Medical Center in Boston. The data includes information such as vital signs, procedure codes, diagnostic codes, medications, laboratory measurements, observations, fluid balance, length of stay, survival data, and more.

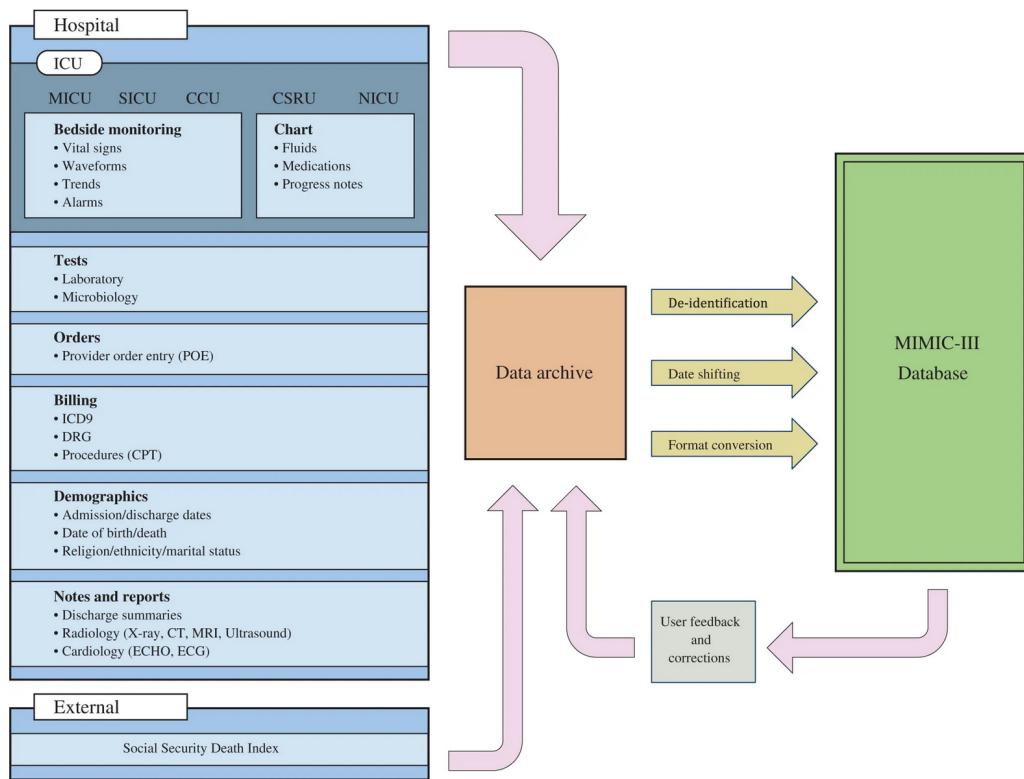
Figure 3.1 depicts the schema of MIMIC-III. On the left, in the blue section, we can see the data that the Beth Israel Medical Center collects. This data is stored in an archive. After applying de-identification, date shifting and format conversion, the EHR data gets into the MIMIC-III database, which is maintained and updated with the help of user feedback by the team of 'The Laboratory for Computational Physiology at Massachusetts Institute of Technology' (MIT) (Johnson et al., 2016)

3.3 Analysis

Before conducting our research, we performed an exploratory data analysis on the MIMIC-III dataset. The goal of this analysis was to discover patterns and correlations in the data to formulate hypotheses for further analysis. We used pandas and numpy for the analysis and matplotlib, to display the results. The total number of unique patients found in the dataset is 46520, consisting of 26121 male and 20399 female patients.

First, as a basic overview, we check for the Top10 most occurring ICD-9 codes in the dataset. As depicted in **Table 3.1**, *essential hypertension* has the highest frequency in MIMIC-III, followed by three more codes of circulatory diseases. *Diabetes mellitus (DM)* can also be found in the list of the most occurring codes.

Next, we have a look at the correlation heatmap (**Figure 3.2**) introduced in (Arya, 2019). This map divided ICD-9 codes into 18 groups. It shows, how many patients have a diagnosis of two groups at a same time and compares every group to each other. It shows the strongest correlation

**Figure 3.1:** Schematic overview over MIMIC-III

ICD Code	Diagnosis	Frequency	Patients affected	% of associated patients
401.9	Essential hypertension	20703	17613	37.86
414.0	Coronary atherosclerosis	15229	13480	28.98
428.0	Congestive heart failure	13111	9843	21.16
427.3	Atrial fibrillation	12891	10271	22.08
584.9	Acute kidney failure	9119	7687	16.52
250.0	Diabetes mellitus	9058	7370	15.82
272.4	Unspecified hyperlipidemia	8690	7465	16.05
518.8	Acute respiratory failure	7497	6719	14.04
599.0	Urinary tract infection	6555	5779	12.42
530.8	Esophageal reflux	6326	5272	11.33

Table 3.1: Top 10 of most occurring ICD-9 codes in the dataset

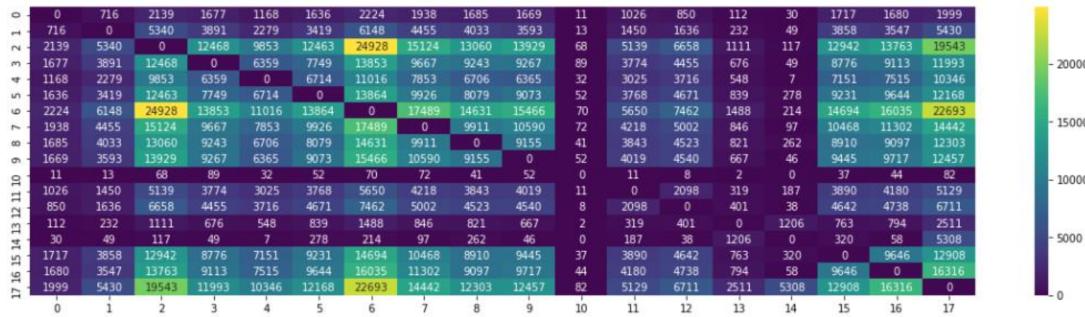


Figure 3.2: Correlation heatmap of ICD code groups (Arya, 2019)

Gender		diabetes	no diabetes	Total	Percentage
Female		1518	2142	3660	41.48%/58.42%
Male		2705	7115	9820	27.55%/72.45%

Table 3.2: Patients that have CAD and diabetes and patients, that have CAD and no diabetes

of ICD-9 codes from the groups two and six, which correspond to *endocrine, nutritional and metabolic diseases and immunity disorders* and *diseases of the circulatory system*. Codes from group two refer to the range of codes from 240-279 , codes from group six refer to codes in the range from 390 to 459. As shown in **Table 3.1**, codes from group six are the most frequent. From group two, we can find two codes, 250.00 *Diabetes mellitus* and 272.4, *unspecified hyperlipidemia*.

Subsequently, we filter all patients with coronary artery disease (Code 414.01) by diabetic diseases (Code 250.*) and compare the frequencies for male and female patients. 7280 female and 9174 male patients are affected by any form of diabetic disease. For coronary artery disease, 3660 female and 9820 male patients are affected.

In **Table 3.2** we can see that female patients with diabetic disease are significantly more often affected by coronary artery disease than male patients. The increased prevalence of cardiovascular diseases for women that have DM is a known risk factor, as already stated in Chapter 2.

3.4 Summary

This chapter gave an insight into the distribution of female and male patients and the most frequent ICD-9 codes in the MIMIC-III dataset. We also learned that statistically, significantly more female patients are affected by coronary artery disease if they also have diabetic disease. This correlation is not present in male patients. In our qualitative evaluation in Chapter 5, we will verify whether our model can reproduce these co-occurrences.

Chapter 4

Electronic Health Record Generation

4.1 Abstract

In this chapter, we will elaborate medGAN, and explain how it differentiates to a traditional GAN. Moreover, we will give an insight into recent improvements of medGAN and its different versions. Further, we will discuss the process of training and generating synthetic Electronic Health Record data on the cloudservice 'Colaboratory' and explain the architecture of medGAN, and our experimental setup. Also, we will introduce our hypotheses and describe how we used the data of the MIMIC-III dataset.

4.2 medGAN

One part of medGAN is a GAN, which consists of two independent neural networks that train each other. The first one is the *Generator (G)* that creates the synthetic data and presents it to the second one, the *Discriminator (D)*. During training real, and synthetic samples are presented to **D**, which tries to distinguish between both. **D** and **G** are both implemented as feedforward neural networks. As we learned in 2.2.1, the generator **G** "is trained by the error signal from the discriminator D via backpropagation, the original GAN can only learn to approximate discrete patient records $x \in Z|C|$ with continuous values." (Choi et al., 2018) To alleviate this limitation, they leveraged an autoencoder, which is reconstructing an dimensionality reduced approximate of the input. As (Choi et al., 2018) explains, during this process, the autoencoder learns the unique features of the samples. This mechanism has previously been used on image processing tasks.

"The objective of the autoencoder is, to minimize the reconstruction error:

$$\frac{1}{m} \left[\sum_{i=0}^m \|x_i - x'_i\|_2^2 \right] \quad (4.1)$$

$$\frac{1}{m} \left[\sum_{i=0}^m x_i \log x'_i + (1 - x_i) \log(1 - x'_i) \right] \quad (4.2)$$

where $x'_i = Dec(Enc(x_i))$

where m is the size of the mini-batch." (Choi et al., 2018)

An autoencoder consists of two elements: The Encoder (*Enc*), which compresses the input and the Decoder (*Dec*) that constructs the output. For binary variables, they used the *mean squared loss* and the *tanh* activation function for *Enc* and the *sigmoid* activation for *Dec*. Both, the *Enc* and the *Dec* are implemented as single-layer feedforward networks. The original input x it receives, is compressed into a 128 dimensional vector. The generator *G* consists of two

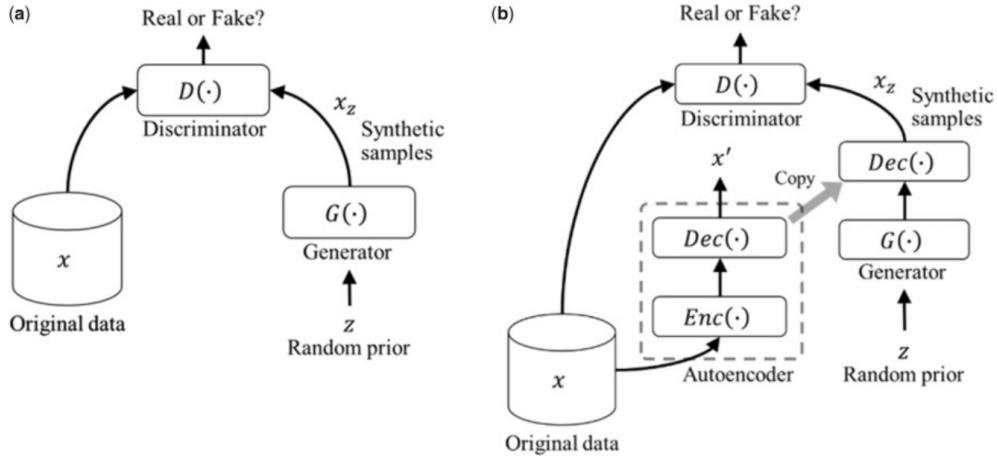


Figure 4.1 Depicts both, the architecture of a traditional GAN (a) and of a medGAN (b). The real data x originates from the EHR data from MIIMIC III. Z acts as the random prior for the generator G . An autoencoder (i.e, the encoder Enc and decoder Dec) is learned from x ; The same decoder Dec is used after the generator G to construct the discrete output. The discriminator D tries to differentiate real input x and discrete synthetic output $Dec(G(z))$. (Choi et al., 2018)

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hidden-layers with each 128 dimensions and is implemented as a feedforward network. For the batch normalization in G they use the scale parameter γ and the shift parameter β and set the moving average decay to 0.99. The discriminator D inherits the same structure, but the first hidden-layer consists of 256 dimensions. MedGAN is trained for 1,000 epochs with the mini-batch of 1000 records. (Choi et al., 2018)

Figure 4.1 Depicts both, the architecture of a traditional GAN (a) and of a medGAN (b). The real data x originates from the EHR data from MIIMIC III. Z acts as the random prior for the generator G . An autoencoder (i.e, the encoder Enc and decoder Dec) is learned from x ; The same decoder Dec is used after the generator G to construct the discrete output. The discriminator D tries to differentiate real input x and discrete synthetic output $Dec(G(z))$. (Choi et al., 2018)

One of the key underlying problems of GANs is *mode collapse*. *Mode collapse* happens when the generator learns a single or a few samples that the discriminator accepts and keeps generating the same sample(s). (Choi et al., 2018) approaches the problem of mode collapse, by introducing *minibatch averaging*, which functions similarly to *minibatch discrimination*. The general idea is that during training, D sees the minibatch of real samples during classifying fake samples and vice-versa. While minibatch discrimination compares the calculated distance of each given sample

and the samples of the minibatch, minibatch averaging calculates the average of the samples in the minibatch. In contrast to minibatch discrimination, minibatch averaging does not require any additional parameters and therefore, its impact on the training time is neglectable. (Choi et al., 2018) Additionally, batch normalization need to be applied, to increase the learning efficiency of \mathbf{G} . Otherwise \mathbf{D} will overpower \mathbf{G} . \mathbf{G} is implemented as a feedforward network with shortcut connections, as shown in **Figure 4.1**'s right hand side. (Choi et al., 2018)

4.2.1 SynthEHR (medBGAN, medWGAN)

Recently, (Baowaly et al., 2018) proposed two altered versions of medGAN that outperform their predecessor, however, just slightly.

Those two versions are:

medWGAN: This version substitutes the regular GAN with an improved *Wasserstein GAN (WGAN)*, that utilizes "an alternative method of weight clipping called gradient penalty, which entails penalizing the norm of the gradient of the discriminator (critic) with respect to its input" (Baowaly et al., 2018)

medBGAN: This version substitutes the regular GAN as well, but this time with a *boundary-seeking GAN (BGAN)*. This approach trains the generator to match the target distribution that converges toward the true distribution as the discriminator is optimized" (Baowaly et al., 2018)

4.2.2 Differentiation

MedWGAN and medBGAN both learn interdimensional relationships better than medGAN. We are trying to proof, that medGAN will improve on this point, if trained with female and male patients separately. If our hypotheses prove to be true and bring an improvement to medGAN, these improvements will also translate to altered versions of medGAN, because not the network itself is being changed but the input. In our tests, we separated the dataset and did not alter it. Henceforth we performed our tests only on the 'original' medGAN.

4.3 Process and Architecture

For generating the Electronic Medical Records (EMR), we used a Generative Adversarial Network (GAN) called medGAN, that was proposed in (Choi et al., 2018). As input data we use v1.4 of the MIMIC-III (Medical Information Mart for Intensive Care) dataset. We train the model with full-length 5 digits ICD-9 codes. Additionally, we train the model with generalized 3 digits long ICD-9 codes . For our experiments, we divided the dataset by gender and generated EMR data with binary and count variables for both mixed and separated patients. The code for medGAN is publicly accessible on github.¹ . It is implemented using TensorFlow. For training models, they chose the Adam-optimizer with a mini-batch size of 100 patients. (Choi et al., 2018) We trained the model using Colaboratory by Google², which is a Jupyter notebook environment, providing free Cloud computing for education and research. The machine is equipped with a Tesla T4 GPU from NVIDIA.

4.4 Experimental Setup

For training, we used the same setup as (Choi et al., 2018). We split the data into subsets with a ratio of 9:1 for training and validation subsets. Using the training subset, the autoencoder is

¹<https://github.com/mp2893/medgan> (01-23-2020)

²colab.research.google.com/

Input Variable	Female	Male	Mixed
binary	22min 53s	26min 49s	43min 29s
count	22min 21s	28min 8s	43min 12s

Table 4.1: Training durations with 3 digit codes

Input Variable	Female	Male	Mixed
binary	39min 42s	48min 12s	1h 26min 23s
count	39min 53s	48min 1s	1h 27min 43s

Table 4.2: Training durations with 5 digit codes

pretrained for 100 epochs. After each epoch, we report the training and validation loss. For binary variables we use the cross-entropy loss function, for count variables the mean squared error. Further, we use minibatch averaging and batch normalization. We conducted our data analysis in a Jupyter Notebook. Here, we use pandas to investigate the data and matplotlib to show our results. After finishing the training process, we select the epoch closest to 0.5, since that is when the discriminator is most confused and the generator makes the most convincing synthetic samples.

4.5 Hypotheses

In this work we are trying to proof the following three hypotheses:

First, the model can generate realistic patients and learns the distribution of ICD-9 codes, if it is trained with the MIMIC III dataset. Second, by training the network with female and male patients separately, its performance improves and it is able to generate patients with gender-specific correlated diseases that seem realistic to a medical doctor. Third, if the network is trained with the MIMIC-III dataset, it is able to generate patients affected by orphan diseases that can not be distinguished to a non-synthetic patient by a medical doctor

4.6 Data

We extracted the ICD-9 codes for female and male patients from the dataset's table 'DIAGNOSES_ICD.csv' separately and aggregated a patient's longitudinal record into a single fixed-size vector. The length of each vector equals the number of possible diagnoses for each patient. As mentioned in 'Process and Architecture', we use both, full-length and generalized ICD-9 codes. **Table 4.3** depicts the vector sizes for the output.

4.7 Models for Comparison

(Choi et al., 2018) compared medGAN with other popular generative methods like Random Noise, Independent Sampling. In their experiments, medGAN outperformed other methods. Therefore we will compare our results with medGAN. To assess the effectiveness of our method, we train the network both, with mixed and gender-separated patients.

4.8 EMR Generation

First, like in (Choi et al., 2018) we generated EMR data, without dividing the dataset. The results serve as baseline for the performance and will be compared with the separately generated patients.

Second, we divided our dataset by gender and generated patient data separately, resulting in 20399 female and 26121 male unique records.

4.9 Rare diseases

Wikidata provides a mapping of 4584 ICD-9 codes to the list of rare diseases of the Genetic and Rare Diseases Information Center (GARD) and OrphaNet IDs. To investigate the occurrence of rare diseases in MIMIC-III, we first generated a list containing all 6985 unique ICD-9 codes of the dataset. Then, we match the list from the mapping which contains a total of 962 codes, resulting in ten corresponding codes in the MIMIC-III dataset. 2408 diagnoses with orphan ICD-9 codes are present in the given dataset. The following section will elaborate our findings regarding rare diseases in our generated patients.

4.10 Summary

In this chapter, we learned about medGAN's architecture and unique characteristics. We also learned about other version of medGAN that were recently release. Further, we will elaborated on the process of training and generating synthetic Electronic Health Record data. We introduced our hypotheses and described how we use the data of the MIMIC-III dataset.

Chapter 5

Experimental Evaluation

5.1 Abstract

In this section we will evaluate our measurements in a quantitative and qualitative manner. We evaluate the measurements for 3 digit (generalized) and 5 digit (original-length) ICD-9 codes for both, binary and count variables. First comes the quantitative evaluation of our generated records. For this we choose two statistical methods: dimension-wise probability for binary variables and dimension-wise average for count variables. Second, we perform the qualitative evaluation. We begin with investigating the data like we did with the real dataset. In the next step, we are looking for correlations of diabetic-disease and heart-disease. Subsequently, we evaluate our measurements with the help of a medical doctor.

5.2 Quantitative Evaluation

In this section we evaluate the model's performance. For the quantitative evaluation of our measurements we choose the following statistical methods, as presented in (Choi et al., 2018).

Dimension-wise probability: This refers to the probability of each dimension (ICD9 code) in the binary dataset. The dimension-wise probability is computed using the following formula:

$$\text{Number of patients} = \frac{\text{Number of patients who had the disease}}{\text{Total number of patients}} \quad (5.1)$$

We calculate it for the binary data.

Dimension-wise average: This refers to the column average of each dimension (ICD9 code) in the count dataset. The dimension-wise average is calculated using the following formula:

$$\text{Dimension - wise average} = \frac{\text{Column sum}}{\text{Total number of records}} \quad (5.2)$$

We calculate it for the count data.

For both, dimension-wise probability and dimension-wise average, we present the outcomes in scatterplots. Each dot represents a diagnoses code. The x-axis represents the codes from the real data, the y-axis the codes from the synthetic data.

Subsequently we will present our findings.

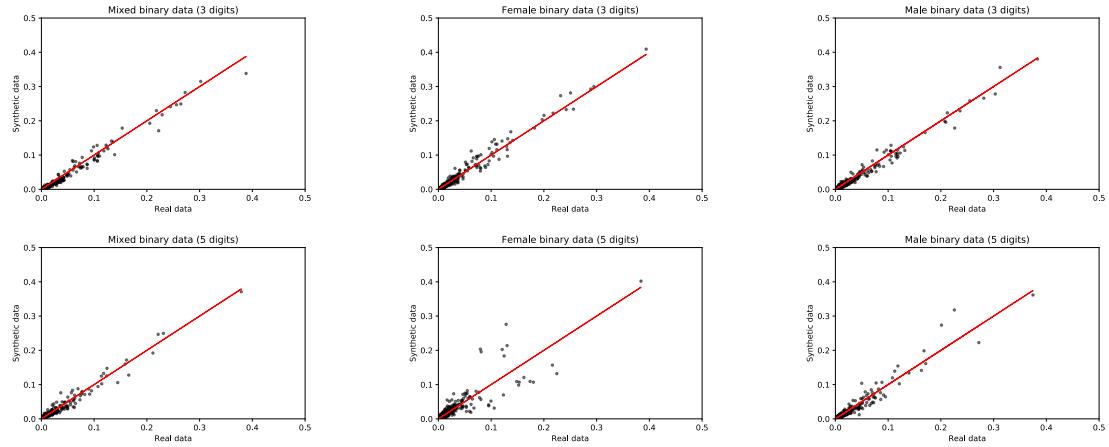


Figure 5.1: Scatterplots of dimension-wise probability

The performance of gender-based models decreased strongly for count data

In **Figure 5.1** we can see, after splitting up the dataset by gender, the performance decreased against our expectations for generating both, male and female samples, while male samples are not affected as heavily. In the best case scenario, all dots would accumulate along the regression line (red). However, we can see that the dots generated with the gender-specific models are more spread out than these for the model containing all data, meaning that the model did not learn the distribution as good as it did with the full input dataset. Because the plots are more spread, we see that especially for count data, the performance decreased, when using gender-based models in comparison to using the full dataset. This affects the model trained with female samples more heavily than the model trained with male samples.

The performance decreased slightly for binary data

Second, we see that the performance loss for binary data is just slightly. The model trained with male samples performed better than the model trained with female samples. This seems to be due to the higher number of input data.

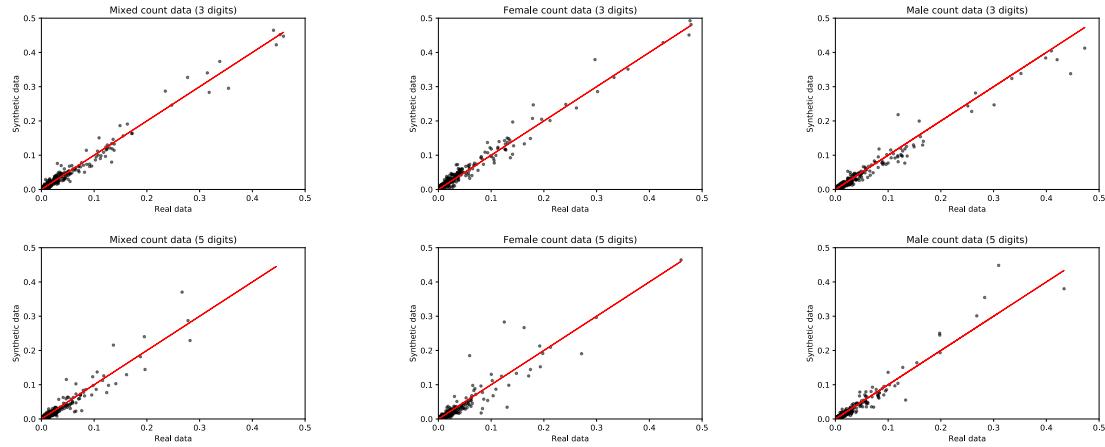
Figure 5.2 depicts, that also for count variables, the performance decreased after splitting up the dataset. This especially affects the model, trained with female data only. This seems to be due to the same reasons as already mentioned for binary data.

5.2.1 Evaluation Orphan diseases

For evaluating our measurements for generated patients with rare diseases, we depict our results in a table with numbers of total occurrences, since we want to find out whether the model learns to generate them at all. First, we choose the binary data for evaluation. Second we will show the results for the count data. For this case, the qualitative evaluation is of higher importance, since we have to find out whether the generated samples seem realistic to a medical doctor.

Orphan diseases:

After comparing the list of diagnoses of MIMIC-III with the list of Orphacodes from wikidata, we find 9 matching codes. Table 5.1 depicts the number of occurrences of each code in the original

**Figure 5.2:** Scatterplots of dimension-wise average

ICD-9 Code	No. of diagnoses
042 Human immunodeficiency virus [HIV] disease	538
075 Infectious mononucleosis	11
138 Late effects of acute poliomyelitis	73
193 Malignant neoplasm of thyroid gland	49
220 Benign neoplasm of ovary	25
317 Mild intellectual disabilities	82
515 Postinflammatory pulmonary fibrosis	544
570 Acute and subacute necrosis of liver	1067
8832 Open wound of finger(s), with tendon involvement	17

Table 5.1: rare disease diagnoses in MIMIC-III

ICD-9 Code	No. of diagnoses binary	No. of diagnoses count
042 Human immunodeficiency virus [HIV] disease	373	265
075 Infectious mononucleosis	0	0
138 Late effects of acute poliomyelitis	24	26
193 Malignant neoplasm of thyroid gland	0	40
220 Benign neoplasm of ovary	0	22
317 Mild intellectual disabilities	0	22
515 Postinflammatory pulmonary fibrosis	492	699
570 Acute and subacute necrosis of liver	870	494
8832 Open wound of finger(s), with tendon involvement	0	49

Table 5.2: successfully generated samples with rare diseases

ICD-9 Code	Percent of affected patients
401.9 Essential hypertension	37.01%
427.31 Atrial fibrillation	24.68 %
414.01 Coronary atherosclerosis	22.81%
428.0 Congestive heart failure	19.22%
272.4 unspecified hyperlipidemia	17.18%
250.00 Diabetes mellitus	16 %
599.00 Urinary tract infection	14.75 %
V29.0 suspected infectious condition of newborn	13.27%
584.9 Acute kidney failure	12.77%
V05.3 Need for prophylactic vaccination against viral hepatitis	12.54%

Table 5.3: Top 10 diagnoses for generated samples (both sexes).

ICD-9 Code	Percent of affected patients
401.9 Essential hypertension	20.47%
427.31 Atrial fibrillation	14.57%
244.9 Unspecified acquired hypothyroidism	11.92%
305.1 Tobacco use disorder	10.60%
424.1 Aortic valve disorders	9.68%
414.01 Coronary atherosclerosis	8.90%
V58.1 Antineoplastic chemotherapy and immunotherapy	8.70%
V05.3 Need for prophylactic vaccination against viral hepatitis	7.78%
250.00 Diabetes mellitus	7.45 %
428.0 Congestive heart failure	7.44%

Table 5.4: Top 10 diagnoses for generated samples (female).

dataset. In **Table 5.2** we see, that our model successfully generates samples with 8 out of the 9 rare disease codes if we use count data. For binary data only 4 out of 9 codes are generated successfully. We can also see that neither the binary nor the count data model successfully generates samples with *Infectious mononucleosis*, that only occurs 11 times in MIMIC-III. The count samples will be thoroughly evaluated by a medical doctor in our qualitative evaluation in the next section.

5.3 Qualitative Evaluation

Our qualitative evaluation consists of three parts. First, we repeat the steps of our data analysis and compare our synthetic samples with it. Second, we investigate whether the model learned gender-specific correlations of diseases and whether it learned to generate sample patients with rare diseases. This can be seen as a further test that proofs whether the model learned the interdimensional relationships. Third, we rate our measurements with the help of a medical doctor. Therefore we take 25 samples from the original dataset and 25 samples from our generated patients. Then, we present them in random order to a medical doctor and let him rate them on realisticness on a scale from 1 to 10, where 10 is the highest score and therefore a sample, that can not be distinguished from a real record. The samples are selected randomly, except for the rare diseases. Here we choose 5 samples each from the dataset and from our generated patients because of their scarcity.

ICD-9 Code	Percent of affected patients
401.9 Essential hypertension	36.16%
427.31 Atrial fibrillation	31.79 %
428.0 Congestive heart failure	27.35%
414.01 Coronary atherosclerosis	22.25%
584.9 Acute kidney failure	19.85%
272.4 unspecified hyperlipidemia	16.12%
V05.3 Need for prophylactic vaccination against viral hepatitis	15.43%
250.00 Diabetes mellitus	14.14%
V29.0 suspected infectious condition of newborn	13.92%
518.81 Acute respiratory failure	13.37%

Table 5.5: Top 10 diagnoses for generated samples (male)

Gender	Affected by DM	Not affected by DM	Total	Percentage
Male	2508	4973	7481	33.52/66.48
Female	1143	1868	3011	37.96/62.04

Table 5.6: Samples CAD diabetes / CAD no diabetes

5.3.1 Data analysis

We compare the Top 10 diagnoses of our generated samples (**Tables 5.3 to 5.5**) to these of the original dataset (**Table 3.1**). For the models trained with mixed and male data, 8 out of 10 ICD9 codes from the Top-10 list from the original dataset occur. For the model, trained with female data only 5 out of 10 ICD9 codes occur. This suggests, that both, the mixed and the male model, perform better in comparison to the female model. For the model trained with the full dataset, most percentages are close to these of the original dataset but there are some outliers, for example Coronary Atherosclerosis with the highest difference of 6.17This confirms our results from the quantitative evaluation.

5.3.2 Gender-specific

MedGAN as provided in its original form, already is, as stated by (Choi et al., 2018), able to generate patient EHR data, that can not be distinguished from real data by a medical doctor. The only exception is, when a patient's longitudinal record shows ICD-9 codes of both, female and male related diseases. (Choi et al., 2018) By splitting up the input dataset into female and male patients, we eliminate this exception. Therefore we evaluate whether the data still seems real after dividing input data into two groups. For our Evaluation, we choose the binary dataset that has been generated by training with 5 digit long codes. The reason is, that we want to compare it with the results from Chapter 3, and for example the code for CAD (414.01) can not be identified by using only 3 digits. First, as a basic check-up, we compare the top 10 occurring ICD-9 codes of our generated samples with those of the original dataset and look for any outliers. Subsequently, we check whether the model learned that female patients have an increased risk for coronary artery disease and cardiovascular disease, if they also have diabetic disease.

Further, we put the focus of our examination on diabetic and ischemic diseases. From our generated samples, 1965 females are affected by both types, while 1689 males are affected by both. When generating patients, without separation 1501 are affected. One of the known correlations is CAD (Cardiovascular Artery Disease) with Diabetes Mellitus. Women have an 3 to 6 fold increased risk if they have DM, while men have a 2 to 4 fold increased risk. (Juutilainen et al., 2004) In our generated data 728 female patients, 907 male patients and 569 mixed patients are affected by DM.

Looking at **Table 5.7** and **3.1** in comparison, we can see that the distribution for male and female samples with CAD and diabetes and these with no diabetes are similar. Therefore we conclude that the interdimensional relationship of cardiovascular disease with diabetic disease is not learned better, if we create gender-based models.

5.3.3 Qualitative evaluation with a medical doctor

Last, we evaluate our measurements with the help of a medical doctor. Therefore we choose 40 samples, 20 from the original dataset and 20 from our generated count datasets. For evaluation, we choose samples from the count dataset that we trained with 5 digit codes as input, as these are they best performing models according to our quantitative evaluation. From the samples with rare diseases, we choose 5 samples from the original dataset and 5 samples from the generated mixed dataset, again because of its superior performance . For the rest of the samples we choose 5 samples each, male, female, and mixed from the count datasets. We ask the medical doctor to act as the discriminator for our samples, as already shown in (Choi et al., 2018). From the 40 presented, he rejected three. One of the female samples, because it contained a code for chronic kidney disease and only one diagnosis for acute kidney failure, which is unlikely. One from the rare diseases samples, because a hypotension and a hypertension occurred together, which is highly unlikely. Another from the rare disease samples, got rejected because it had female-related and male-related codes. Apart from these outliers, the samples are indistinguishable to a real record.

5.4 Discussion

A discussion with the medical doctor thought us that the evaluation of a patient's record on realistic-ness solely with diagnoses codes is a challenging task. The reason for this is that diseases are not mutually exclusive and every diagnosis is possible. Also generalization to 3 digit ICD codes is not helpful, because essential information about each diagnosis is lost. For example, shortening the ICD-9-CM Diagnosis Codes for a symptom like hallucinations (ICD-9 780.1) will only occur as 'General symptoms' (ICD-9 780) after generalization.

5.5 Summary

In this chapter we evaluated our measurements in both, a quantitative and qualitative context. We learned from the quantitative evaluation, that separating the input data by gender does not improve the models performance in this context. The models trained with 3 digit ICD-9 codes outperformed those trained with 5 digit ICD-9 codes. Further the accuracy of the binary models is higher than the accuracy of the count models. In our qualitative evaluation, a medical doctor rated the majority of the synthetic samples as real. In our discussion with him, we learned that using 3 digit codes generalizes diseases and symptoms to strongly and makes the samples hard to interpret. This thought us that, while the 3 digit code models performed best in our quantitative evaluation, they are not superior in a qualitative manner.

Chapter 6

Conclusion and Outlook

6.1 Goal

Our goal was to proof, that medGAN can generate realistic patients if it is trained with the MIMIC III dataset and that the model learns the distribution of ICD-9 codes. Further we tried to elaborate if by training the network with female and male patients separately, it is able to generate patients with gender-specific correlated diseases that seem realistic to a medical doctor. Also we wanted to find out whether the model is able to generate patients affected by orphan diseases that can not be distinguished to a non-synthetic patient by a medical doctor despite the rare occurrences in the dataset.

6.2 Results from Evaluation

Our model successfully generates realistic patient data that can not be distinguished to real data, this also includes samples with rare diseases. However, the performance of medGAN did not increase after separating the input data by gender.

6.3 Outlook

6.4 Future work and Outlook

Application to other MedGAN versions Generating diagnosis and medication codes is the first step to generate high-quality synthetic EHR data. <https://www.aclweb.org/anthology/W19-5026/>
Problem: Utility of the data -> making it useful Generating synthetic EHR data is the first step to replacing real training data. The next step that would improve the utility of the samples could be the generation of clinical text with keyphrases. Our generated data could act as input data.

6.5 Summary

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Acknowledgements

I would like to express gratitude to Prof. Dr. Alexander Löser for the insightful discussions and valuable comments as well as his support and supervision of this thesis. I also would like to express my gratitude to my advisor Betty van Aken for her continuous guidance and supervision. Further I want to thank Dr. med. Sait Subasi for evaluating the generated data. Without their help, this thesis would not have been possible.

Appendix A

Bibliography

Appendix B

Listings

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Appendix C

Results

XXX
