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Preliminary analysis of claims data to understand
relationship between disease patterns and quality of care
and its implications for health insurance in India

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Abstract

This paper provides preliminary analysis of claims data of Mediciam insurance scheme to understand the relationship between disease pattern and the quality of health care. We use length of stay (LOS) and average length of stay (ALS) as one of the indicators of quality of care. We use the Diagnostic Related Grouping (DRG) based ALS as the benchmark to make this evaluation and comparison. It is observed that the reimbursements in insurance system are tied to hospital inputs and resource use and not to diagnostic related groups or outputs. Therefore the current system of reimbursements and provider payment system influences the length of stay and there is significant variation in ALS observed across disease groups and its sub-groups. There is no consistency observed in ALS as the severity of diseases under each group increases. This reflects lack of standards/protocols and unintended consequences of current practice of provider payment system. Implementing systems like Diagnosis Related Grouping would be an attempt to link it with outcomes. The paper provides insights into whether there is a significant mismatch in the premium that insurance companies charge in comparison to the risk insurer undertake while issuing policies. It was also found that after adjusting for the purchasing power parity, the claims data suggest that healthcare costs reimbursed for medical insurance to private providers in India are actually higher than healthcare costs reimbursed to providers of healthcare in the US under DRG system. The paper argues that under less regulated private healthcare providers market and health insurance market, cost based reimbursement is highly undesirable. The regulators should put in place a system of pre-determined rates for reimbursements in health insurance.

Preliminary analysis of claims data to understand relationship between disease patterns and quality of care and its implications for health insurance in India

Introduction

Voluntary medical insurance Mediciam is an indemnity based scheme, in which policyholders, on payment of a fixed premium, are covered for insurance up to a certain amount of sum assured. The policy is renewed every year. The premium is based on age of the policy holder and amount of sum assured. On hospitalisation, the policyholder is expected to first bear the complete cost of payment out-of-pocket and later they are reimbursed after verification of the claim submitted to the insurance company. It is only recently some insurance companies with the help of third party administrators are providing cash less facility.

This system of reimbursement is cost based and is rife with shortcomings, the primary among them being: (a) moral hazard which implies high probability of collusion between healthcare providers and patients to increase service provision and thereby affecting billing amounts, with lack of any overseeing authority to prevent the same; (b) absence of standardisation of medical diagnosis and treatment implying that even in the absence of moral hazard, due to a diverse array of diagnosis and treatment practices for the same disease across different healthcare providers all over India, verification of reimbursement claims took inordinately long periods of time, introducing inefficiency into the system; and (c) adverse selection implying that the system does not appear to have adequate screening for assessing the risk of contracting a particular disease. Currently, insurance premiums are a factor of two independent variables sum insured and age profile. The diversity of health insurance products which can take care of these problems and address the needs of consumers are lacking. For example, one finds that no premium is attached to the presence or absence of definite risk factors of an individual. This in turn increases risk of adverse selection to the insurance company. This is further accentuated by the fact there are serious information asymmetry problems. It seems that these shortcomings are perhaps the primary reasons for the reluctance of foreign insurance companies to enter into the Indian health insurance market, in spite of its huge potential. The estimated annual healthcare expenditure in India is Rs. 1030 billion (Bhat and Babu 2004). The presence of more players is likely to help the consumers as it increases the competition leading to increased and wider choice of products to consumers.

To facilitate development and move towards more mature insurance systems the Insurance Regulatory and Development Authority (IRDA) has recognised the need for third party administrators (TPAs) to act as an intermediary in the insurance sector, facilitating the coordination between insurance company, healthcare provider and consumer. This has entailed the concept of cashless hospitalisation. This system currently is facing a large number of teething problems, with opposition from healthcare providers due to tardiness in reimbursements.

Before liberalisation of insurance industry, Mediciam has been the sole health insurance product with a large number of exclusions and shortcomings, such as non-coverage of OPD based treatment. Not many variations in product offering exist which can meet need of different persons. There is a need for introducing novel insurance products, which deal with specific health concerns of a large section of the population. The variations could be in terms of variables such as: diseases covered, types of treatment covered, type of individual risks covered and so on. The introduction of such products would definitely increase risks such as adverse selection by narrowing the selection pool. This can in turn be supported by: (a) more rigorous screening procedures, (b) minimising the risk premium mismatch that occurs in blanket policies by customizing risk premium according to degree of perceived risk on screening, and (c) standardising norms and requirements of diagnosis and treatment.

The objective of this paper is to examine the relationship between disease pattern and quality of care. We use one simple indicator of quality of care and it has been measured in terms of average length of stay (ALS). We use the Diagnostic Related Grouping (DRG) based ALS as the benchmark to make this evaluation. Using this information the paper provides some insights into whether there is a significant mismatch in the premium that insurance companies charge in comparison to the benefits consumers gain and the risk insurer undertake while issuing policies. The paper makes an attempt to explore what modifications can be made with respect to present insurance scheme in terms of modifying the product characteristics so as to increase the benefit and decrease the risk without having implications for number of policies issued or amount of sum insured. The analysis presented in this paper may provide insights into the possibility of designing and introducing health insurance products.

Methodology and data analysis

The primary analysis presented in this paper is based on data of 621 claims, reimbursements, medical records and socio-economic indicators pertaining to policy initiation years 1997-1998 and 1998-1999 of the Ahmedabad branch of GIC's subsidiary. This data was manually collected in 2001-02. The raw data, mostly hand-written by doctors, contained a large amount of variation in describing the disease and illness. In some cases the information presented could not be clearly translated in exact illness. Some written prescriptions were having factual inaccuracies and at times the description was misleading. A large number of discrepancies in the medical terms used for the final diagnosis of the disease showed up in the data. For example, a case of Cerebrovascular Accident (commonly known as a Stroke) was variously termed as intracranial bleed, intracranial haemorrhage, stroke, UMN lesion and so on. Similar instances existed for most diseases, across all disease groups.

All diseases were segregated and renamed according to a standard nomenclature system followed by hospitals in the Municipal Corporation of Greater Mumbai. Information of the amount of insurance premium paid by policy holders also threw up some inconsistencies or omissions. As far as possible, such instances have been corrected, but data which could not be cleaned has been eliminated for the purpose of further analysis. A total of 579 cases were found to be valid for further analysis. The data were classified and presented in a standard format.

Data reclassification in Diagnosis Related Grouping (DRG): The data provided was in need of a mode of classification for analysis. Secondary research revealed a few systems for classification/grouping of disease, such as ICD-10 and DRG. Diagnosis Related Grouping (DRG) is a system created by the Federal Government of the USA in 1983 as a way to assess payment requirements for Medicare patients. A DRG is a 3 digit number that describes a particular medical diagnosis. Under the DRG system, a hospital receives the same fee for all patients diagnosed under a particular DRG disease, irrespective of how sick the patient is, how costly it is to treat the patient or how long the duration of stay of the patient is. This gives hospitals an incentive to reduce the cost of treatment and length of stay in the hospital. The DRG system has been adopted by many private insurance companies in the USA and Australia.

The DRG system provides a standardized diagnosis nomenclature, an *Average Length of Stay (ALS)* and a standard average *Cost Ceiling* for each disease. For example, Epistaxis (commonly known as bleeding from the nose) is classified as DRG 66, with an ALS of 3.2 hospital days and an average cost ceiling of USD 964.

Length of stay (LOS) and average length of stay (ALS) has been one important indicator to measure the hospital performance. It is considered to have significant influence on cost of care and can also be used as surrogate measure for cost. Generally hospitals having long ALS may be relatively inefficient in the use of resources and those with low ALS are considered to be efficient. Sometimes, however, LOS is assumed to relate to quality (Thomas, Guire and Horvat 1997). Reducing length of hospital stay (LOS) is a policy aim in many countries to regulate their health

care systems and is thought to indicate efficiency. For example, it is generally viewed that longer than expected LOS is indication of poor quality of care. In this study we use the LOS and ALS as indicators of quality of care.

Diseases from the primary data of 579 reimbursement cases were reclassified using the nomenclature followed by the Municipal Corporation of Greater Mumbai (MCGM). These were subsequently grouped according to disease groups such as Central Nervous System (CNS) for all diseases related to the brain and nervous system, Cardiovascular System (CVS) for all diseases related to the heart and the circulatory system and so on. Diseases with similar nomenclature were grouped together and each group then matched to its respective DRG nomenclature disease. The average of the actual length of hospitalization of each group was calculated, as also the average cost incurred. This enabled us to compare the Average Length of Stay and the Average Cost prescribed by the DRG classification system vis-à-vis the *Average Actual Length of Stay* undergone and *Average Costs* incurred by the patients in the 579 cases. An illustrative example is provided in Table 1.

Table 1: Illustrative example of ALS and average cost

| Diagnosis (Disease) | Cases | | | |
|----------------------------------|-------|------------------|-------------------|------------------|
| | 1 | 2 | 3 | 4 |
| | Fever | Fever with Cough | Fever with Chills | High Grade Fever |
| MCGM Diagnosis (Disease Group) | PUO* | PUO | PUO | PUO |
| Actual Length of Stay | 7 | 3 | 4 | 10 |
| Average Actual Length of Stay | | | | 6 |
| DRG Length of Stay | | | | 10 |
| Actual Cost of Treatment | 1000 | 1600 | 900 | 500 |
| Average Actual Cost of Treatment | | | | 1000 |

* Pyrexia of unknown origin (requiring hospitalisation)

Other variables used in the analysis of data include: (a) the presence of risk factors associated with the disease, (b) the ratio of the premium paid to sum assured, (c) age and gender of the patient (data regarding other socio-economic status was unavailable), and (d) days taken to settle the claim.

The classified data was subjected to a graphical analysis of average length of stay (DRG-ALS) and actual length of stay for each disease system. Diseases within each disease system are grouped in the ascending order of their average lengths of stay. The graphs on each disease system are provided in Exhibits 1 to 15 covering the following disease systems: Central Nervous System, Ophthalmology (Eye Diseases), Otorhinolaryngology (ENT), Respiratory System, Cardiovascular System, Gastrointestinal System, Hepatobiliary System (Liver and Pancreas Diseases), Orthopaedics (Bone diseases), Breast Diseases, Dermatology (Skin Disease), Endocrine System, Renal System (Kidney and Urinary Disease), Male Reproductive System, Obstetric & Gynaecological Disease, and Infectious Disease. The standard deviations of the actual duration of hospitalisation stay of each diagnosed disease group (within each disease system) are provided in Exhibit 16. The time duration (in days) between submissions of the indemnity claim (date of discharge from hospital) and claim reimbursement per disease group is provided Table 2. The table is arranged in descending order of claim settlement period.

Table 2: Disease systems and claim settlement period

| Disease System | Claim settlement period |
|---|-------------------------|
| Endocrine System (ES) | 182 |
| Central Nervous System (CNS) | 180 |
| Orthopaedics (OR) | 142 |
| Cardiovascular System (CS) | 140 |
| Breast Diseases (BD) | 138 |
| Otorhinolaryngology (ENT) | 135 |
| Respiratory System (RS) | 132 |
| Hepatobiliary System* (HS) | 129 |
| Obstetric & Gynaecological Disease (OG) | 126 |
| Renal System (RS) | 112 |
| Dermatology (D) | 107 |
| Infectious Disease (ID) | 103 |
| Gastrointestinal System (GS) | 101 |
| Male Reproductive System (MRS) | 101 |
| Ophthalmology (OP) | 99 |
| * Liver and Pancreas Diseases | |

From the above table, it is clear that while the claim reimbursement process is inordinately long across all disease systems, the average claim settlement period was 128 days. Descriptive Statistics for claim settlement period are provided in Exhibit 18. We carried out the following correlation analysis: (a) correlation between time taken to settle the claim and ALS, and (b) time taken to settle the claim and amount claimed across all disease groups. The results are presented in Table 3.

Table 3: Relationships

| Disease System | Claim Settlement Period (A) | ALS (B) | Amount Claimed Per Day (C) | Correlation A and B | Correlation A and C | Correlation B and C |
|----------------|-----------------------------|---------|----------------------------|---------------------|---------------------|---------------------|
| ES | 182 | 4.0 | 3398 | -0.24 | 0.32 | 0.61 |
| CNS | 180 | 5.3 | 4446 | -0.56 | 0.59 | 0.03 |
| OR | 142 | 4.0 | 4783 | -0.11 | 0.35 | 0.58 |
| CS | 140 | 7.0 | 6351 | 0.00 | 0.13 | 0.70 |
| BD | 138 | 1.9 | 10489 | -0.35 | -0.09 | 0.89 |
| ENT | 135 | 1.8 | 5596 | -0.13 | 0.11 | 0.68 |
| RS | 132 | 7.0 | 1640 | 0.41 | 0.34 | 0.45 |
| HS | 129 | 4.9 | 5183 | 0.09 | 0.38 | 0.74 |
| OG | 126 | 6.0 | 3170 | -0.25 | 0.22 | 0.32 |
| RS | 112 | 8.0 | 2032 | -0.09 | -0.30 | 0.72 |
| D | 107 | 3.9 | 2099 | -0.05 | 0.01 | 0.82 |
| ID | 103 | 4.0 | 1862 | -0.16 | 0.21 | 0.25 |
| GS | 101 | 4.0 | 2346 | 0.01 | -0.10 | 0.42 |
| MRS | 101 | 3.0 | 3934 | 0.10 | 0.41 | 0.70 |
| OP | 99 | 0.7 | 18397 | 0.20 | NA | 0.10 |

The correlation between ALS and amount claimed per day was very high and statistically significant from zero. This correlation was extremely high (>0.66) in case of 7 of the 15 disease systems under analysis. This indicates that the reimbursements are cost based. It was observed

that the correlation between ALS and claim settlement period and amount claimed and claim settlement period were not significant.

We carried out the correlation between Length of Stay (DRG) and Actual Length of Stay of our reimbursement cases. The results are provided in Table 4.

| Disease Group | Correlation Coefficient DRG and ALS |
|----------------------|--|
| ES | -0.07 |
| CNS | 0.20 |
| OR | 0.04 |
| CS | 0.31 |
| BD | 0.21 |
| ENT | 0.14 |
| RS | 0.43 |
| HS | 0.24 |
| OG | 0.29 |
| RS | 0.02 |
| D | -0.24 |
| ID | -0.04 |
| GS | 0.37 |
| MRS | 0.29 |
| OP | -0.02 |

There appeared to be no definite pattern in correlation between the DRG Length of Stay and actual observed ALS across similar disease groups. In reference to a DRG system which is well classified and standardised system the actual data do not indicate any consistent pattern across different groups of illnesses. The lack of correlations across different disease groups may also indicate difference in morbidity patterns in the US and India. However, the results signify the need for a new system of standardised reference system for Indian healthcare insurance claim verification. The disease-wise relationship between ALS as per DRG and actual ALS of our sample displayed a significant variation (see Exhibit 16).

The duration of time between discharge and claim settlement, one of the major cited problems with the current indemnity based health insurance process implemented by GIC appeared to be due to nature of disease itself, apart from the obvious systemic inefficiency in settling health insurance claims. This interpretation coincides with the observation that treatment of each different disease system, and even within each disease system varies greatly in terms of the following:

- Methods and cost of diagnosis,
- Duration of therapy (hospitalised and post hospitalisation),
- Methods and cost of therapy (medical, surgical, physical etc),
- In order to eliminate systemic inefficiency in claim processing, it is necessary to increase the accuracy of data provided for analysis of claims,
- Number and efficiency of personnel involved in claim reimbursement verification

There was high degree of correlation between length of stay and claim amount indicated that the longer time a patient spends in the hospital, the higher is the claim amount. While this sounds intuitively logical, it is pertinent to point out that this variable i.e., length of stay is most critical and is subject to manipulation by both consumers and providers to increase reimbursement claim amounts. More so this is also cost based and significant variation can take place in this across

different providers. This suggests towards the need for a standardised system of claim verification and reimbursement that should be based on properly researched data and medical practice.

Further the analysis of the data suggests that there is an immediate need for an insurance regulatory body IRDA and healthcare regulatory body such as the Medical Council of India or a leading healthcare providing organization such as AIIMS to formulate a set of rules and guidelines for standardised nomenclature of disease. Varied nomenclature of disease makes it extremely difficult for other medical practitioners and health insurance agencies to gauge the type, severity and /or complexity of a particular disease diagnosis, resulting in inability to assess the validity of insurance claims, in terms of necessity of hospitalisation, number of days hospitalised and cost of treatment accruing thereon. A standardised number of days of hospitalisation for each disease group, and a fixed payment for the same, similar to the DRG could be instituted and adopted by the MCI and recognised healthcare organisations. This, as mentioned before, will encourage healthcare providers to keep costs under control and provide quality healthcare at an affordable cost. At present, as displayed in the primary data findings, the large variance in days of hospitalisation results may result in highly inefficient system of insurance. This may result in high cost health care. Delay and difficulty in assessing health insurance claims also result in increased time gap between the dates of discharge to actual date of reimbursement (under the erstwhile indemnity form of health insurance). Disease System wise data regarding time gap between date of discharge and date of claim reimbursement is provided in Exhibit 17.

A standardised system of diagnosis and treatment for each disease, similar to ‘critical care pathway’ adopted at American hospitals such as Massachusetts General Hospital, Boston for CABG can be adopted by healthcare providers to minimise costs of healthcare delivery, especially in the case of commonly encountered ailments.

The data was further analysed based on the following three parameters: (a) gender, (b) presence or absence of risk factors contributing to disease, and (c) presence or absence of associated disease. The data segmentation is provided in Exhibits 18. This data was then subjected to observation for trends, regression and correlation analysis. The following observations were found to be significant. The significant findings and their possible interpretations are provided below.

Gender based morbidity patterns: Males appeared to have a higher predilection towards Gastrointestinal disease, Orthopaedic disease, Infectious disease, Ophthalmologic disease and Cardiovascular disease in that order. Females, on the other hand, showed a higher predilection towards Ophthalmologic, Orthopaedic, Gastronintestinal, Obstetric and Infectious diseases. Given the dominance of a particular set of diseases towards each gender, this indicates the need for similar analysis across a larger cross section of data to determine nationwide trends of gender based morbidity. Special gender based health insurance products could be an outcome of this. The average length of stay across all disease groups appears to be lower in the case of female patients. This may indicate a social bias, more than a higher morbidity pattern for male patients.

Fig 1: Distribution of Disease Groups

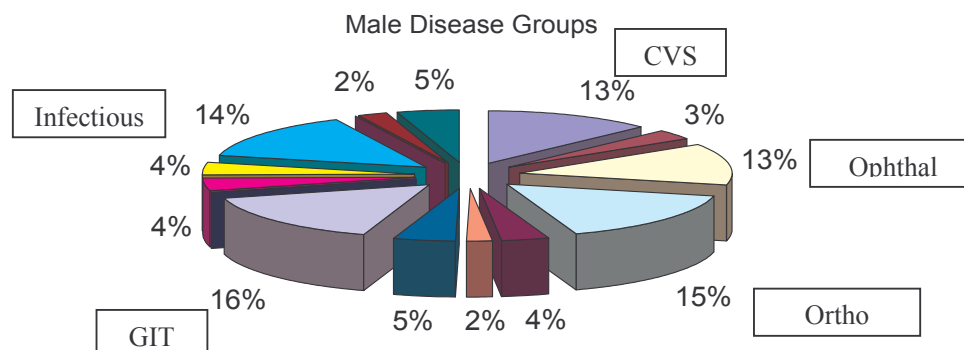
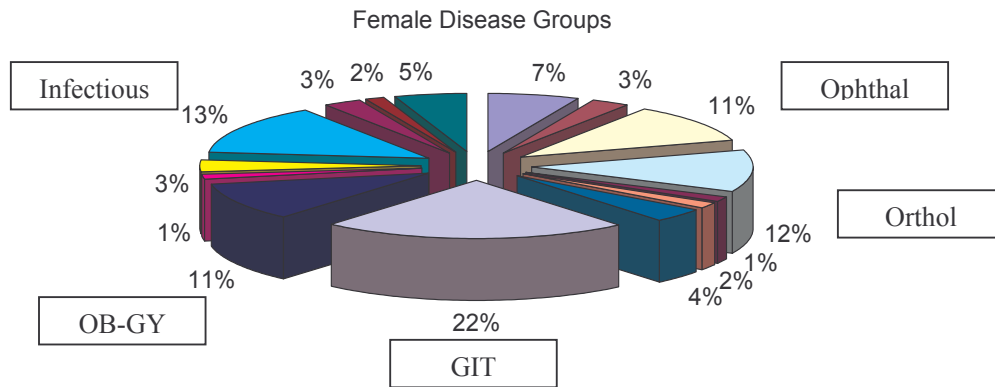
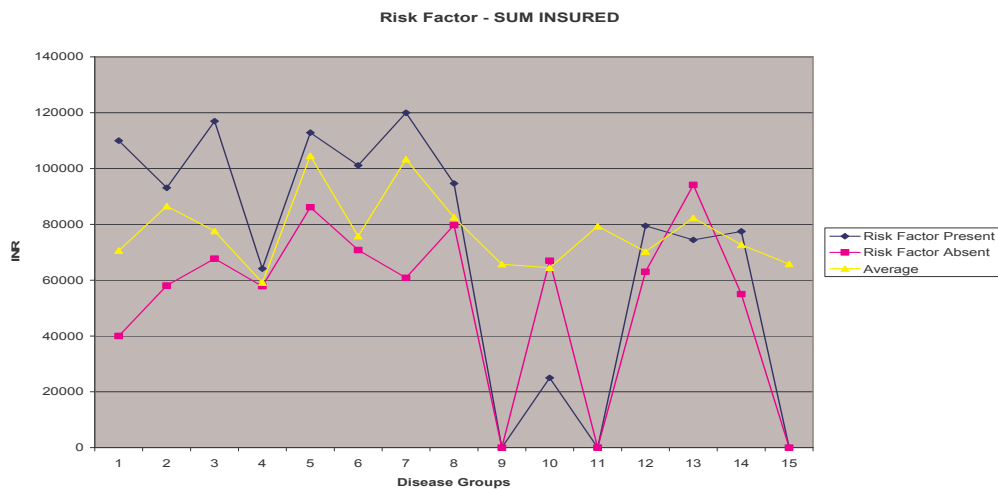


Fig 2: Distribution of Disease Groups – Female patients



Risk factor based analysis: It is observed that the difference in both the Sum Insured and the *Amount Claimed* between patients who possessed definite risk factors for a particular disease is higher across most disease groups. We plotted the sum assured and amount claimed after classifying the cases in high risk and low risk cases. The data suggested difference (see Figures 3 and 4). The premiums in Mediclaim health insurance scheme are determined by two factors and these are (a) age profile of the policyholder (premiums increase with increasing age, with 6 well defined risk groups (5-45, 46-55, 56-65, 66-70, 71-75 and 76 and above), and (b) amount of sum assured (lower the amount insured for, lower is the premium in proportion terms). An analysis of the primary data showed that within the same age group policy holder were charged the same premium/sum insured rate. The experience in disease pattern however had been different as they suffered from different ailments entailing different cost of treatment, claims and reimbursements. While this is a definite consequence of the fundamental ‘pooling of risk’ of insurance, there exist very definite cases in which the ailment of the individual could be predicted on the basis of the presence of *risk factors* for that particular disease. For example, within the Central Nervous System disease group, an individual in the 46-55 age group, who suffered from a hypodense lesion in the brain (for which no definite risk factor can be ascertained) has been charged the same premium as an individual who suffered from a cardiovascular accident (which has definite risk factors such as hypertension, diabetes and atherosclerosis).

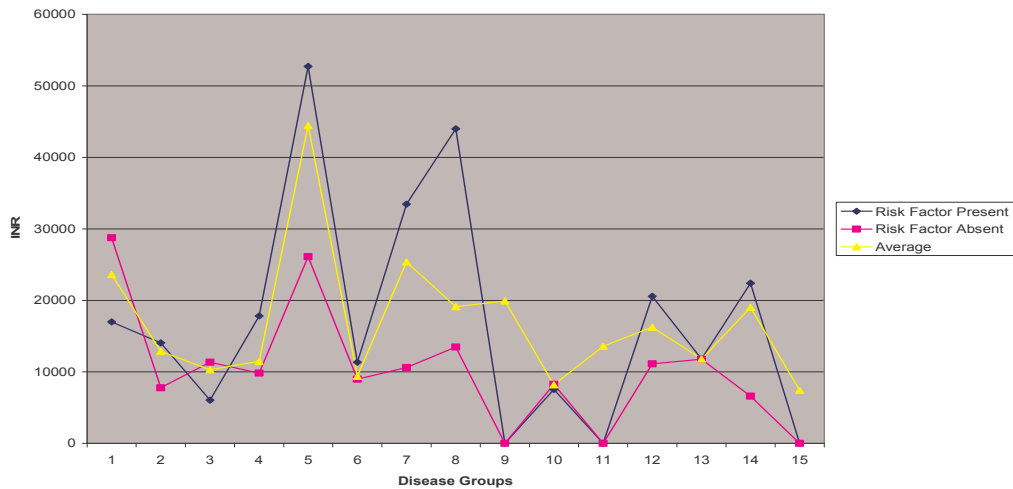
Fig 3



The amount paid by the insurance company as reimbursement was 7.8 times more in the case of the latter policy holders. Ideally, the insurer should have been able to charge a higher premium from the latter individual, given the definite presence of certain risk factors in their medical

history. Numerous other examples exist in the data. A complete analysis could not be made without the presence of medical records of each case. The development of insurance products and their coverage and pricing need to take this analysis into account. The claims data can be used to make comprehensive post-facto analysis of the mismatch between risk premium and claim amount. Other attributes, such as occupation of the policy holder, gender etc., should ideally be taken into consideration for assessing the health insurance premium to be paid by a potential policy holder.

Fig 4
Risk Factor - AMOUNT CLAIMED

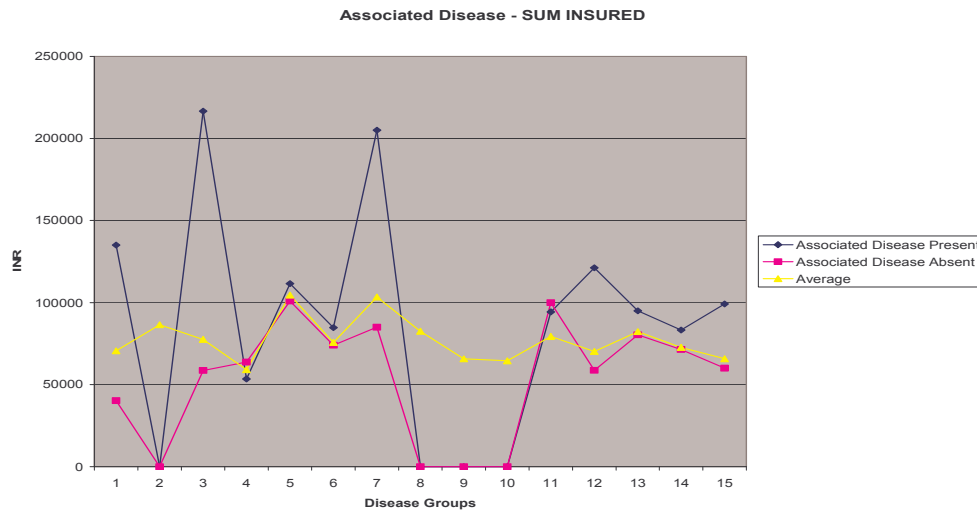


This observation leads to two possible inferences: (a) presence/knowledge of disease specific risk factors is incentive for patients to hide the same during filling of insurance forms, but at the same time insure themselves for higher amounts - an example of moral hazard in insurance, and (b) this can be avoided by providing for a comprehensive medical examination by an independent panel for all cases of sum insured above a particular amount in the absence of any compelling reason. Thus, while simple risk perception and disposability of income cannot be ruled out as causes for high sum insured, presence of disease specific risk factors must be ruled out

Associated disease based analysis: Similar to the above, the presence of associated disease increases the value of the sum insured across most disease groups (see Fig 5).

The data appear to indicate a similar inference as that provided in risk factors scenario as discussed above, that presence of associated disease tends to increase the tendency to insure for a higher amount.

Fig 5



We also carried out regression analysis of days taken to settle the claim and average length of stay. We used ALS as independent variable. Table 5 shows the results of regression analysis.

Table 5: Regression results

| Disease Group | R ² value |
|--|----------------------|
| Central Nervous System | 0.3114 |
| Ophthalmology (Eye Diseases) | 0.0412 |
| Otorhinolaryngology (ENT) | 0.0159 |
| Respiratory System | 0.1648 |
| Cardiovascular System | 0.0000 |
| Gastrointestinal System | 0.0002 |
| Hepatobiliary System (Liver and Pancreas Diseases) | 0.0053 |
| Orthopaedics (Bone diseases) | 0.0113 |
| Breast Diseases | 0.1213 |
| Dermatology (Skin Disease) | 0.0027 |
| Endocrine System | 0.0508 |
| Renal System (Kidney and Urinary Disease) | 0.0071 |
| Male Reproductive System | 0.0074 |
| Obstetric & Gynaecological Disease | 0.0633 |
| Infectious Disease | 0.0261 |

There appears to be no correlation between the dependent and the independent variable.

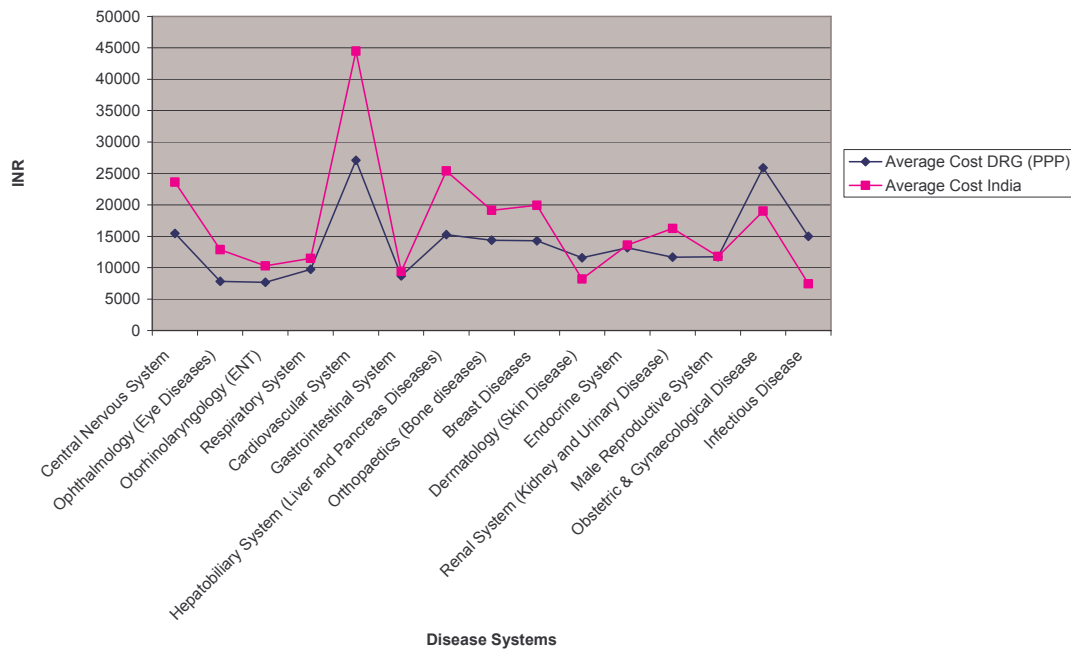
Comparison of the cost of treatment: We made a comparison of cost using DRG statistics and the actual amount of money claimed for hospitalisation in the Mediclaim scheme. Firstly, a comparison between average healthcare costs pre disease system (viz. CNS, Ophthalmology etc.) as prescribed by DRG (adjusted for Purchasing Power Parity) with the average costs of hospitalisation undergone by Indian patients claiming reimbursement from Mediclaim was made. Next, the analysis is performed one step further to study trends at a disease subsystem (e.g., Meningitis, Epilepsy and Cerebrovascular Accident in CNS) level.

As depicted in the chart below, healthcare costs in the USA and India appear to show similar patterns, with the most expensive disease systems, in descending order adjusted for purchasing power parity are shown in Table 6.

| Rank | USA | India |
|------|-------------------------------------|--------------------------------|
| 1 | Cardiovascular Disease | Cardiovascular Disease |
| 2 | Obstetrics & Gynaecological Disease | Hepatobiliary Disease |
| 3 | Central Nervous System Disease | Central Nervous System Disease |
| 4 | Hepatobiliary Disease | Breast Disease |
| 5 | Infectious Disease | Orthopaedic Disease |

From the table it appears that Cardiovascular System disease, Central Nervous System disease and Hepatobiliary System disease appear to be amongst the most expensive across both countries in terms of hospitalisation costs. Also visible from the chart is the fact that, after adjusting for purchasing power parity (assuming 1USD in the USA = 4.5 USD in India), it is observed that healthcare costs reimbursed for medical insurance to private providers in India are actually more than healthcare costs reimbursed to providers of healthcare in the US under DRG system (see Fig 6). This is contrary to popular perception that medical services are expensive in the US than in India. This also suggests that in India the cost based reimbursement is expensive and we need some standardisation of reimbursement system which should be based on pre-determined rates than cost based reimbursement. There also appears to be an economic anomaly arising due to various demand and supply factors the private providers face. Data pertaining to individual disease system and cost comparisons are provided in Exhibits 18 onwards.

Fig 6
DRG vs Indian Data Costs



Conclusion and implications

There is an immediate need for insurance regulatory body and healthcare regulatory body such as the Medical Council of India or a leading healthcare providing organisation such as AIIMS to formulate a set of rules and guidelines for standardised nomenclature of disease. Varied nomenclature of disease makes it extremely difficult for other medical practitioners and health insurance agencies to gauge the type, severity and/or complexity of a particular disease diagnosis, resulting in inability to assess the validity of insurance claims, in terms of necessity of hospitalisation, number of days hospitalised and cost of treatment accruing thereon. Third Part Administrators face a challenge in regulating costs of care in health insurance system. A standardised number of days of hospitalisation for each disease group, and a fixed payment for the same, similar to the DRG could be instituted and adopted by the MCI and recognised healthcare organisations. This, as mentioned before, encourages healthcare providers to cut costs and provide quality healthcare at an affordable cost. At present, as displayed in the primary data findings, the findings suggest the following.

The existing system promotes cost based reimbursement leading to high cost of healthcare. This will make insurance systems vulnerable to high cost and affect viability of schemes. In less regulatory health insurance regime like ours, cost based reimbursements in insurance will cause health care cost inflation. Hospital's revenue is function of cost. Once they spend more and if insurance is paying they get more. Health insurance therefore will encourage over use of resources making health care cost high. Capital costs will become dominant and there would less incentive to substitute capital for labour. In developing countries where *per se* the need for spending on health is high, high levels of private health expenditures through insurance pose serious challenge. The sheer size of these expenditures once it has risen to high levels can impede control of health expenditures itself. This also leads to exploitation of patients by healthcare providers for monetary gain. Delay and difficulty in assessing health insurance claims, resulting in increased time gap between the dates of discharge to actual date of reimbursement (under indemnity form of health insurance). Disease system wise data regarding time gap between date of discharge and date of claim reimbursement is provided in Exhibit 17.

A standardised system of diagnosis and treatment for each disease, similar to 'critical care pathway' adopted at American hospitals such as Massachusetts General Hospital, Boston for CABG can be adopted by healthcare providers to minimise costs of healthcare delivery, especially in the case of commonly encountered ailments such as Malaria and URTI. The American DRG system cannot be implemented directly in India, given the difference in morbidity patterns between the two countries. A modification of the same could however be implemented. This should take into account both statistical data collated over a significant period of time to arrive at reasonable *Length of Stay* and *Average Cost of Treatment* at different level hospitals (primary, secondary and tertiary)

The analysis of claims data suggests that the disease profile is the most important criterion governing: (a) duration, modality and cost of treatment, and (b) time delay between discharge and reimbursement of claim amount. It is suggested that new health insurance products, with disease specific coverage may be introduced. For example, the product option may include products which provide coverage for specific illnesses. Individuals, depending on their risk profiles, desirous of insuring themselves against specific diseases such as cardiovascular disease can be provided with a policy which requires them to pay a higher premium for the same. The premium amount would be a function of number of policy holders for that disease and data provided on standard costs of treatment for cardiovascular diseases. Similarly insurers can think of having bundled health insurance policies. Individuals perceived to be at risk of diseases belonging to different disease systems (e.g., employees in IT companies are at risk of developing lifestyle diseases such as hypertension, obesity, computer vision syndrome etc.) can avail of health insurance schemes targeted towards these diseases. A basic premium would cover care for these diseases, and an extra amount could be charged per disease not in the basket of diseases. Being

targeted towards segments *at risk* such as occupational segments, companies can avail of larger enrolment, and thus pooling of risk will occur. Similarly, insurers can think of region based health insurance policies. Morbidity patterns in India differ from region to region. Differential premiums across regions, with higher premiums charged in areas of higher susceptibility to a particular disease can be instituted. Customised health insurance policies can be other option to introduce the products in India. Current blanket insurance schemes can be replaced by customised insurance policies in which consumers pick and choose from a basket of disease and disease solutions, with varying risk premiums for each based on individual probability of morbidity and anticipated cost of treatment. Rigorous screening is of course necessary for implementation of such customised products. There are also opportunities to use treatment and disease information to make clinical determinations in connection with health care coverage decisions for policyholders. Policies can be written on selected clinical issues, especially addressing new technologies, new treatment approaches, and procedures. Actual clinical determinations in connection with coverage decisions can be made on a case-by-case basis. These are possibilities provided there are well accepted diagnostic and treatment parameters for a particular disease, both clinically and diagnostic test wise. This will also promote standardisation in healthcare benefits disbursement, something lacking in the current health insurance system.

There are also opportunities to develop health insurance products which cover some specific diseases. For example, these can include chronic diseases requiring prolonged OPD based therapy and support such as asthma, diabetes, low back pain and coronary heart disease. Alternatively, health insurance policies may include add-on options which provide concessional rates for alternative healthcare services focusing on preventive care and services such as ayurveda/ homeopathic systems of medicine, fitness clinics, massages etc. Insurance companies can tie-up with pharmaceutical companies and retail chains to provide medicines at lowest possible cost. This would be an innovative way of expanding the customer base by enticing potential policy holders with lifestyle products and benefits. There can also be people specific benefits in insurance schemes. These can include focus on schemes such as (a) women's health programmes, (b) mental health policies, (c) vision and dental programmes, and (d) students benefits schemes – designed specifically for students, covering adolescent health problems and commonly diseases faced by students. Individual/group plans are other opportunity. These plans can be designed specifically for groups of 20 to 50 people, offering them concessional rates and add-on benefits.

Exhibit 1: Average Length of Stay v/s Actual Length of Stay - CNS

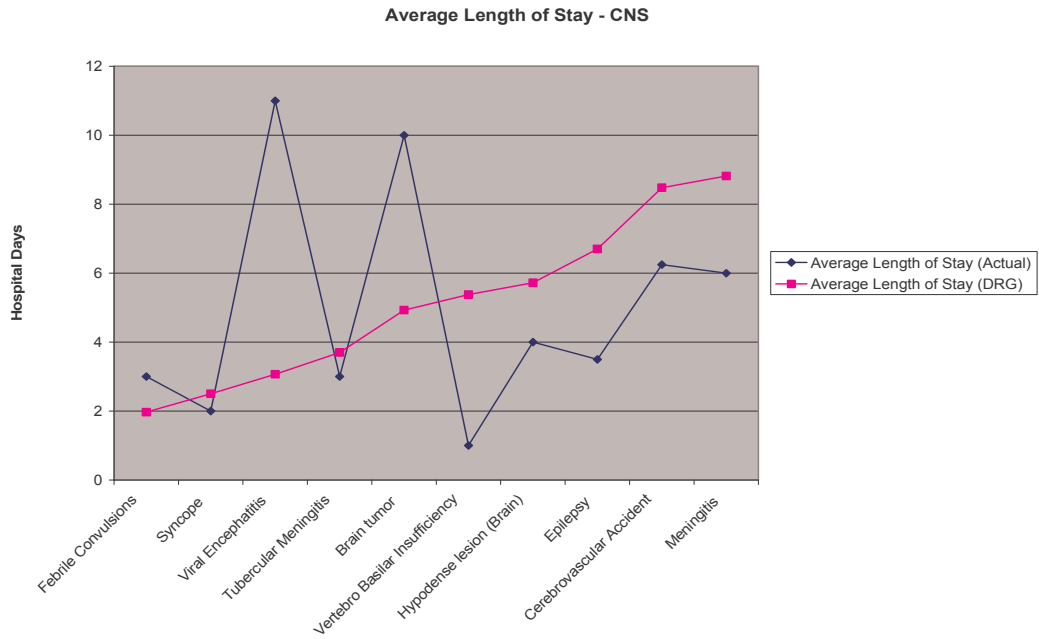


Exhibit 2: Average Length of Stay v/s Actual Length of Stay – Ophthalmology

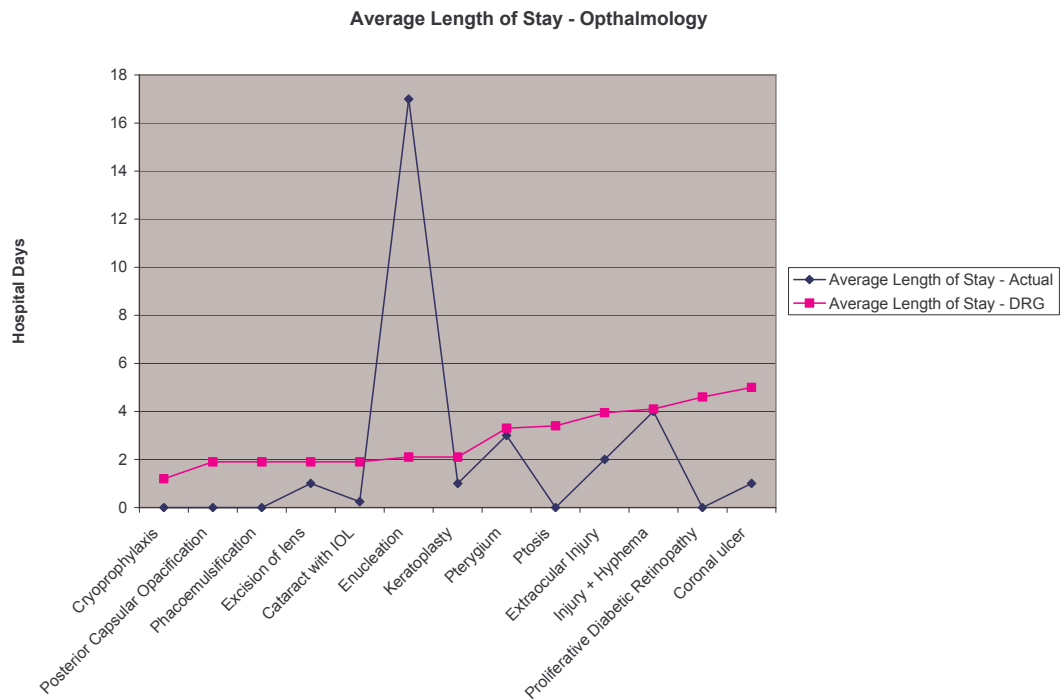


Exhibit 3: Average Length of Stay v/s Actual Length of Stay – ENT

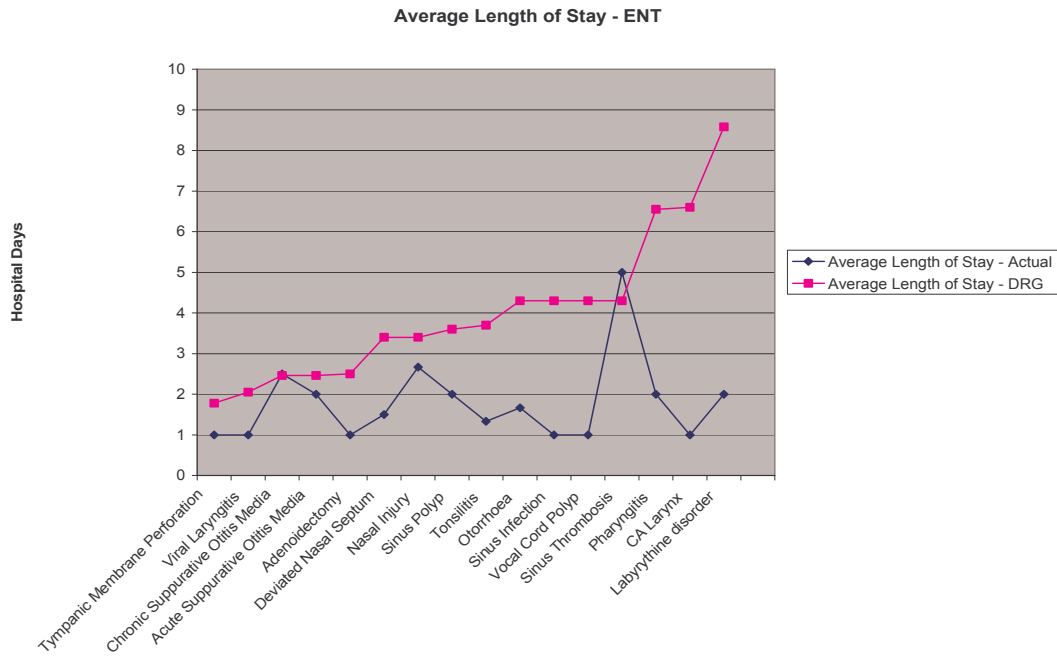


Exhibit 4: Average Length of Stay v/s Actual Length of Stay – Respiratory

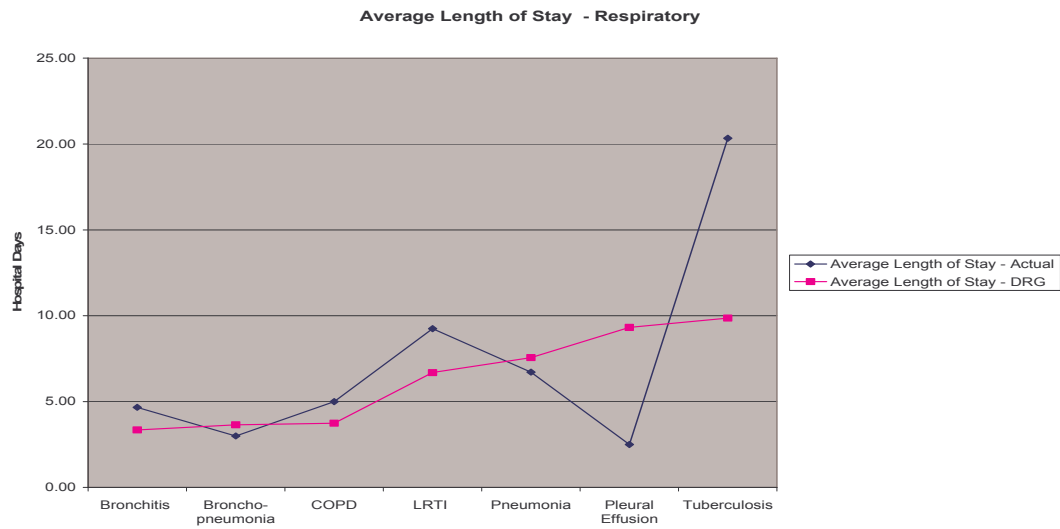


Exhibit 5: Average Length of Stay v/s Actual Length of Stay – CVS

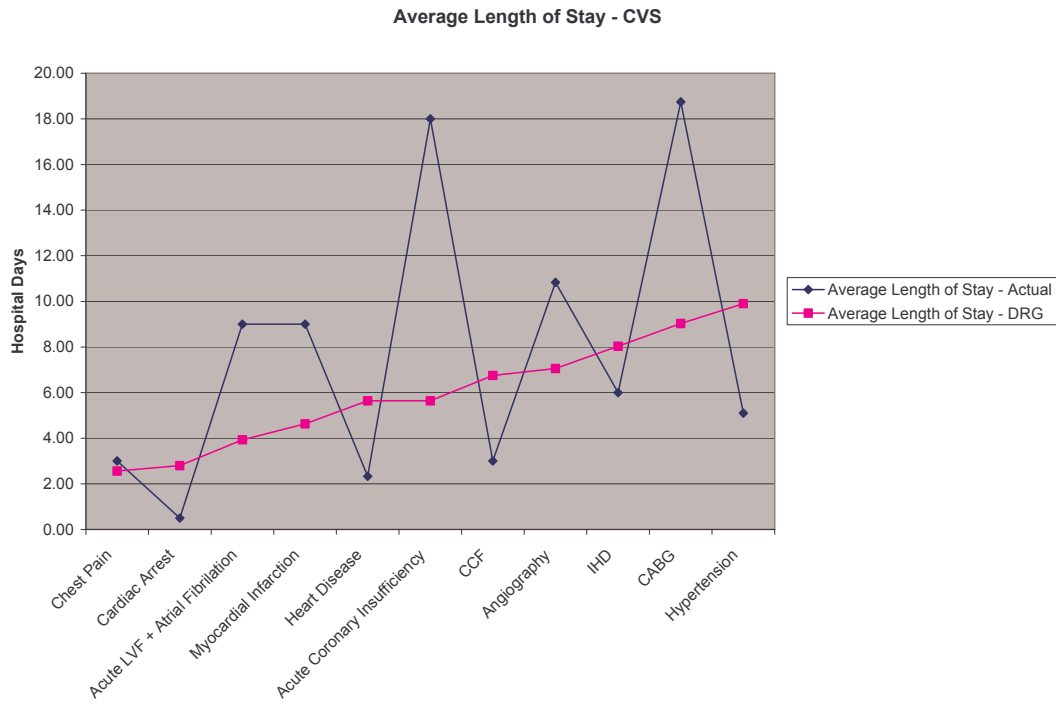


Exhibit 6: Average Length of Stay v/s Actual Length of Stay – GIT

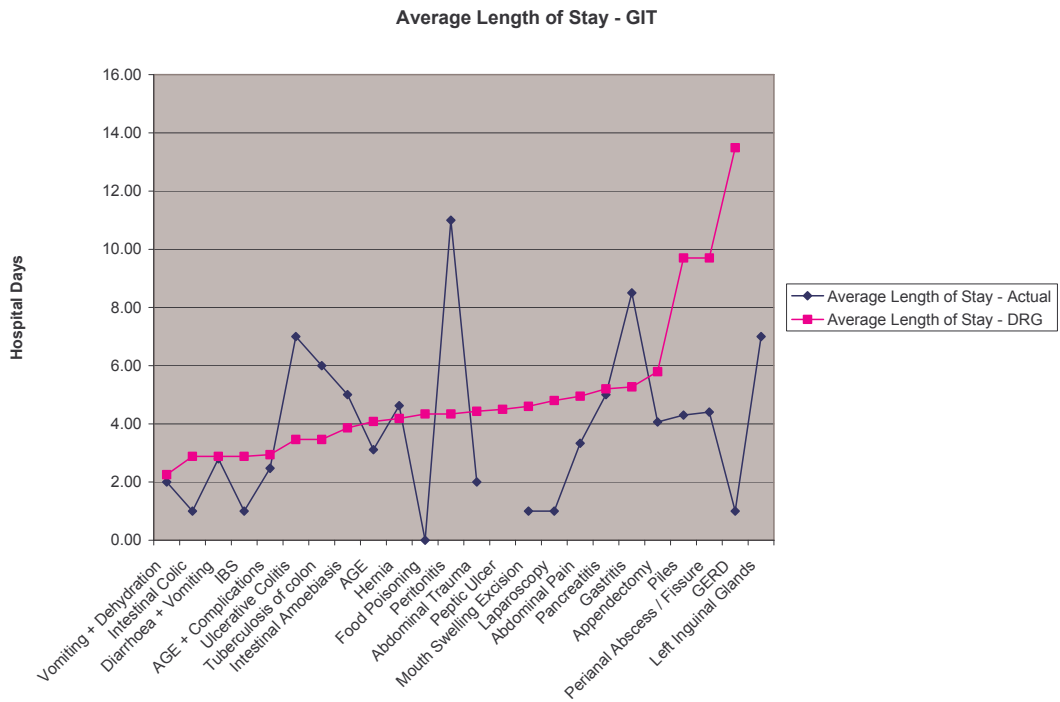


Exhibit 7: Average Length of Stay v/s Actual Length of Stay – Hepatobiliary System

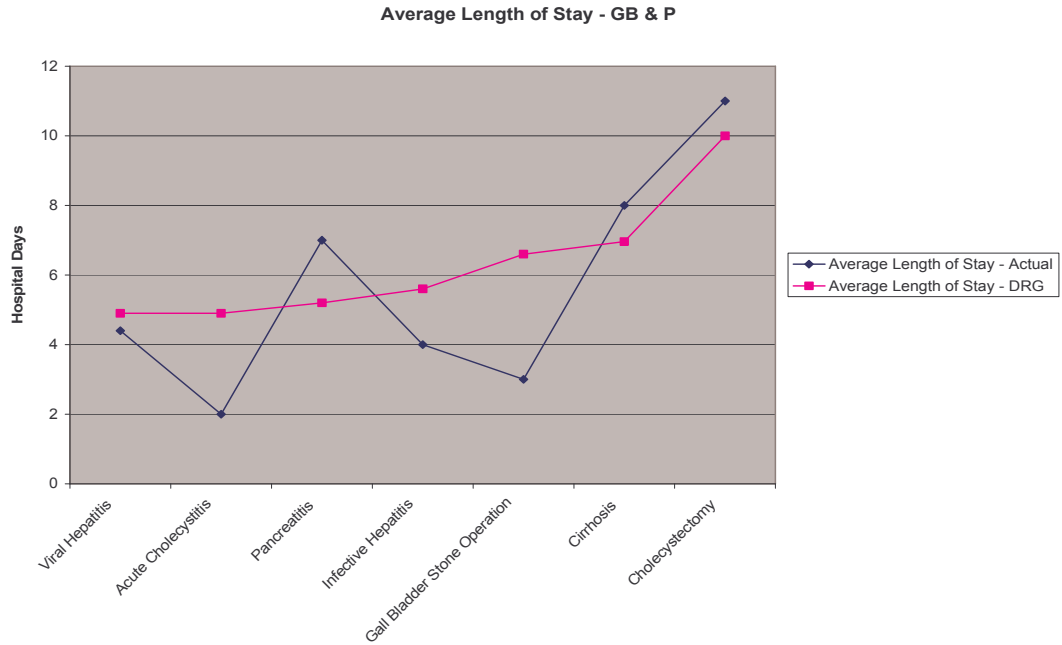


Exhibit 8: Average Length of Stay v/s Actual Length of Stay – Orthopaedics

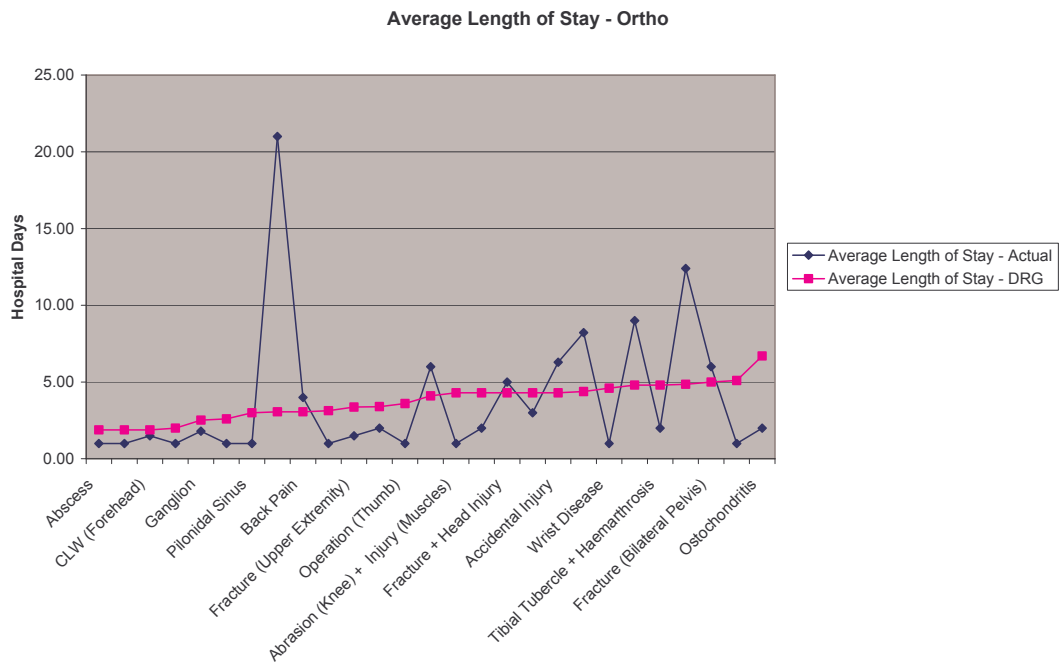


Exhibit 9: Average Length of Stay v/s Actual Length of Stay – Breast Disease



Exhibit 10: Average Length of Stay v/s Actual Length of Stay – Dermatology

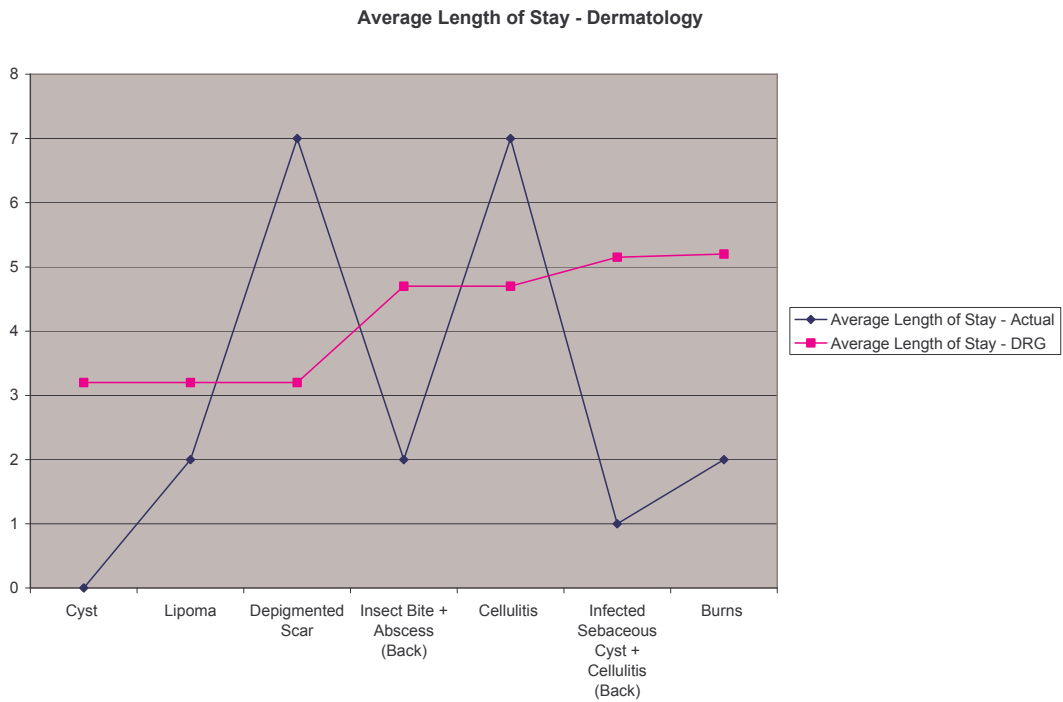


Exhibit 11: Average Length of Stay v/s Actual Length of Stay – Endocrine

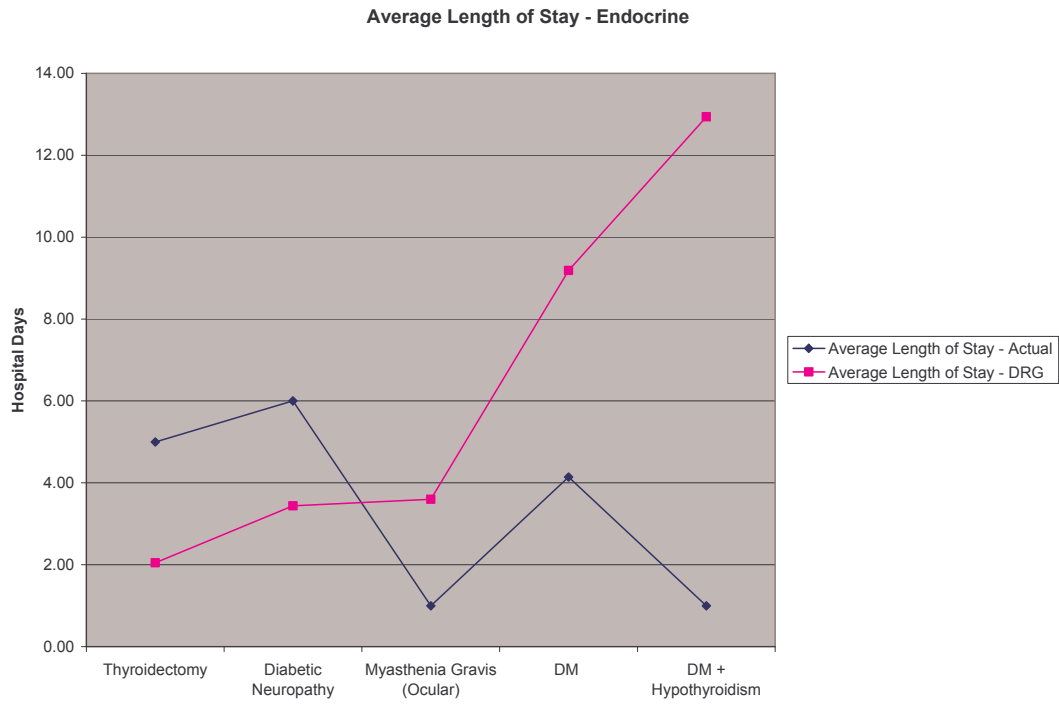


Exhibit 12: Average Length of Stay v/s Actual Length of Stay – Renal

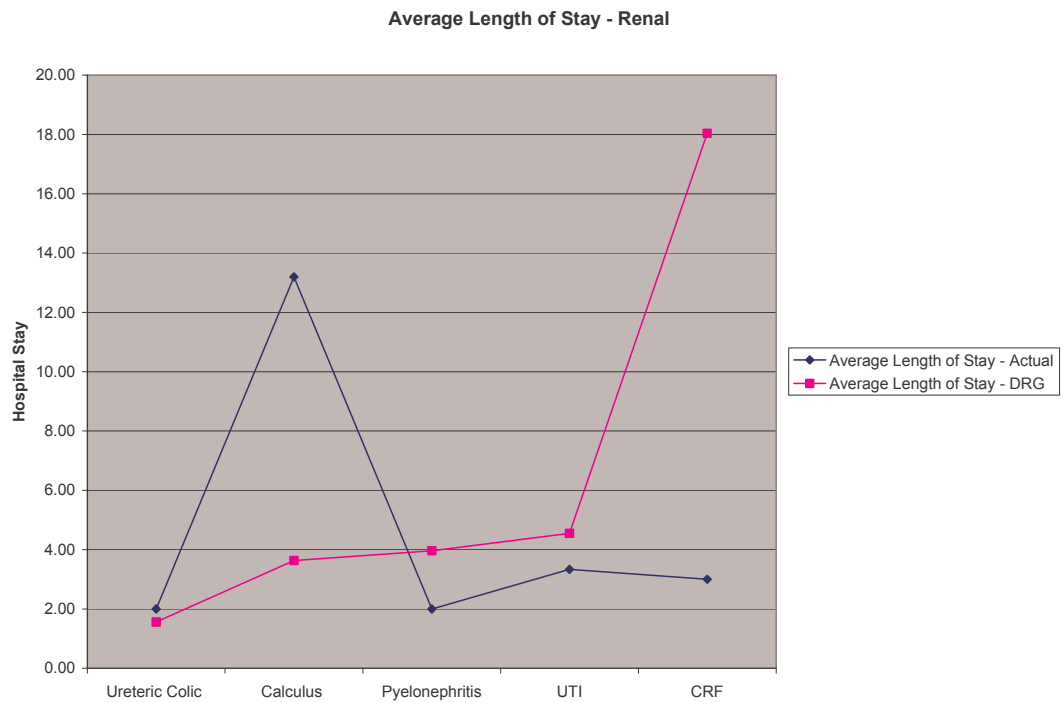


Exhibit 13: Average Length of Stay v/s Actual Length of Stay – Male Reproductive System

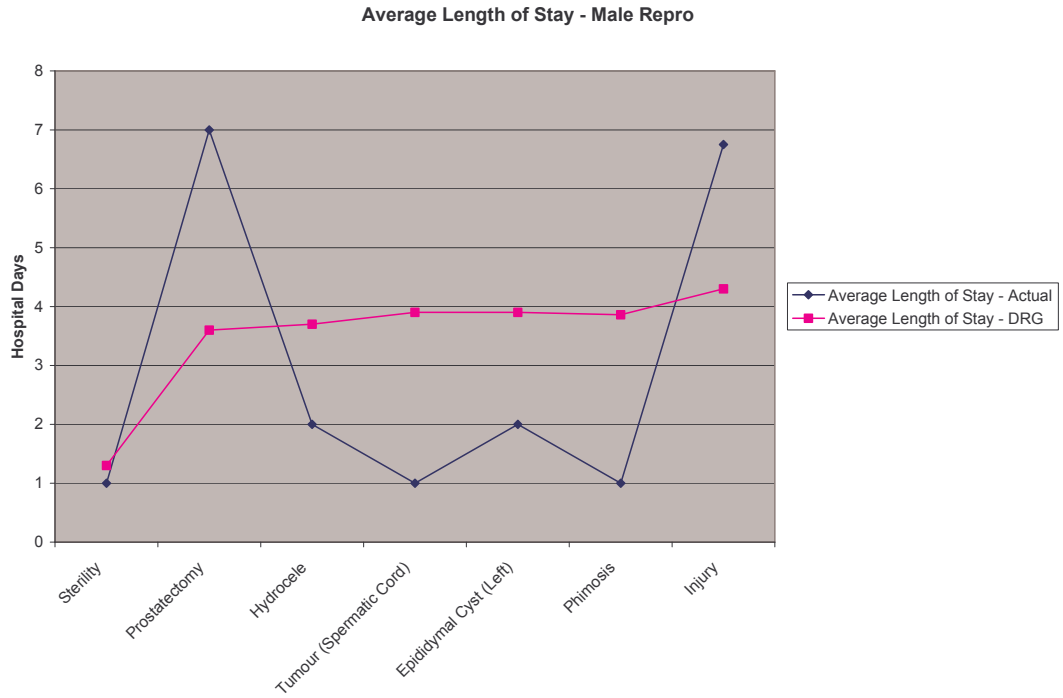


Exhibit 14: Average Length of Stay v/s Actual Length of Stay – Obstetrics & Gynaecology

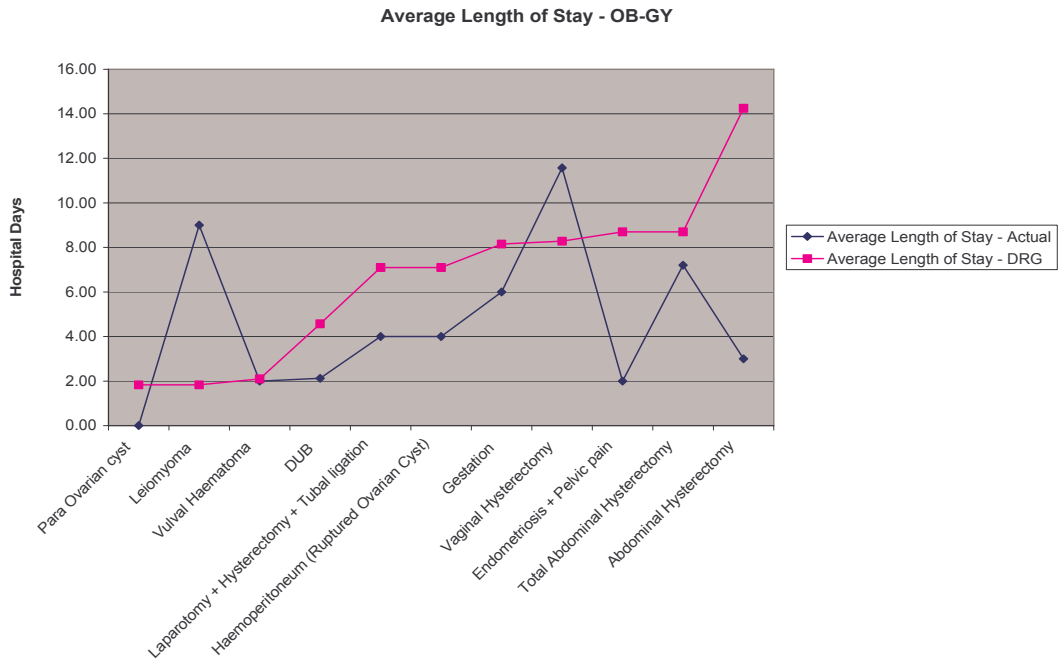


Exhibit 15: Average Length of Stay v/s Actual Length of Stay Infectious Disease

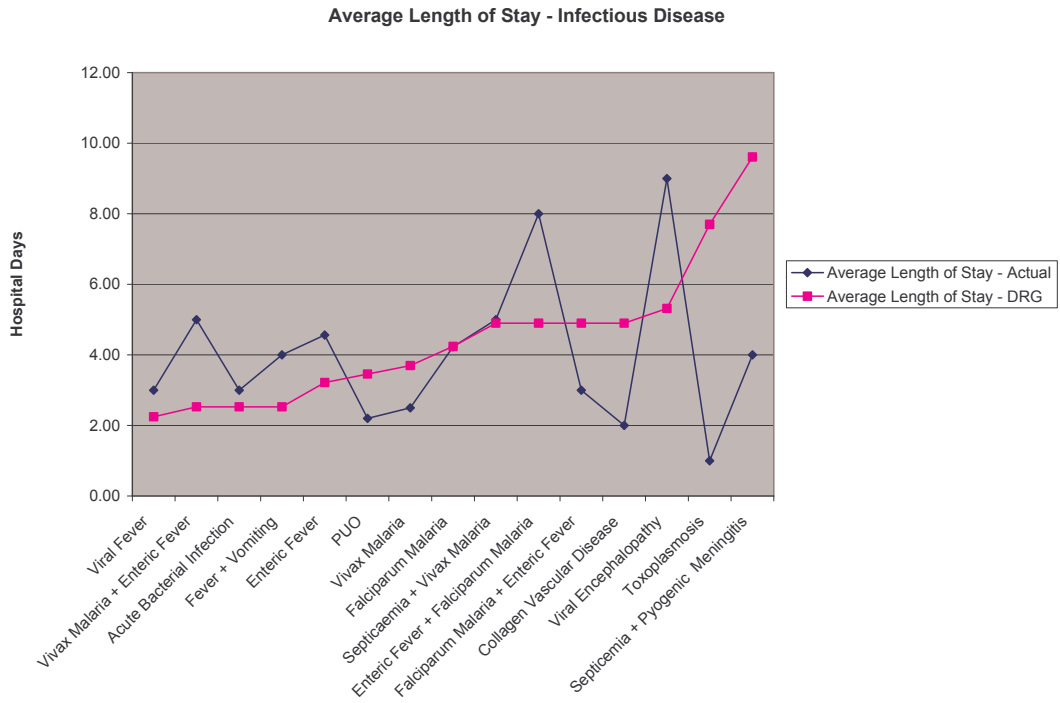


Exhibit 16: Disease Group: Actual Duration of Stay - Standard Deviations

| No | Disease System | Disease Group | Standard Deviation |
|----------------------|----------------|------------------------------------|--------------------|
| 1 | CNS | Febrile Convulsions | na |
| | | Syncope | na |
| | | Viral Encephatitis | 0 |
| | | Tubercular Meningitis | na |
| | | Brain tumor | na |
| | | Vertebro Basilar Insufficiency | na |
| | | Hypodense lesion (Brain) | na |
| | | Epilepsy | 0.71 |
| | | Cerebrovascular Accident | 3.86 |
| | | Meningitis | 4.24 |
| 2 | Ophthalmology | Cryoprophylaxis | na |
| | | Posterior Capsular Opacification | na |
| | | Phacoemulsification | na |
| | | Excision of lens | na |
| | | Cataract with IOL | 0.47 |
| | | Enucleation | Na |
| | | Keratoplasty | 0 |
| | | Pterygium | 4.24 |
| | | Ptosis | na |
| | | Extraocular Injury | 1.41 |
| | | Injury + Hyphema | na |
| | | Proliferative Diabetic Retinopathy | na |
| 3 | ENT | Coronal ulcer | na |
| | | Tympanic Membrane Perforation | na |
| | | Viral Laryngitis | na |
| | | Chronic Suppurative Otitis Media | 2.12 |
| | | Acute Suppurative Otitis Media | na |
| | | Adenoidectomy | na |
| | | Deviated Nasal Septum | 0.71 |
| | | Nasal Injury | 0.58 |
| | | Sinus Polyp | na |
| | | Tonsillitis | 0.58 |
| | | Otorrhoea | 1.15 |
| | | Sinus Infection | na |
| | | Vocal Cord Polyp | na |
| | | Sinus Thrombosis | na |
| | | Pharyngitis | na |
| CA Larynx | na | | |
| Labyrythine disorder | 0 | | |
| 4 | Respiratory | Bronchitis | 5.24 |
| | | Broncho-pneumonia | 1.73 |
| | | COPD | na |
| | | LRTI | 9.5 |
| | | Pneumonia | 6.52 |
| | | Pleural Effusion | 2.12 |
| | | Tuberculosis | 26.63 |
| 5 | CVS | Chest Pain | 4.40 |

| | | | |
|---|---------------|---|-------|
| | | Cardiac Arrest | 0.71 |
| | | Acute LVF + Atrial Fibrillation | na |
| | | Myocardial Infarction | 2.83 |
| | | Heart Disease | 3.21 |
| | | Acute Coronary Insufficiency | na |
| | | CCF | 2.83 |
| | | Angiography | 12.48 |
| | | IHD | 4.85 |
| | | CABG | 13.94 |
| | | Hypertension | 6.15 |
| 6 | GIT | Vomiting + Dehydration | 0.00 |
| | | Intestinal Colic | na |
| | | Diarrhoea + Vomiting | 1.30 |
| | | IBS | na |
| | | AGE + Complications | 1.01 |
| | | Ulcerative Colitis | na |
| | | Tuberculosis of colon | na |
| | | Intestinal Amoebiasis | na |
| | | AGE | 2.53 |
| | | Hernia | 2.72 |
| | | Food Poisoning | na |
| | | Peritonitis | na |
| | | Abdominal Trauma | na |
| | | Peptic Ulcer | na |
| | | Mouth Swelling Excision | na |
| | | Laparoscopy | na |
| | | Abdominal Pain | 1.53 |
| | | Pancreatitis | na |
| | | Gastritis | na |
| | | Appendectomy | 2.05 |
| | | Piles | 6.07 |
| | | Perianal Abscess / Fissure | 5.64 |
| | | GERD | na |
| | | Left Inguinal Glands | na |
| 7 | Hepatobiliary | Viral Hepatitis | 2.61 |
| | | Acute Cholecystitis | na |
| | | Pancreatitis | 2.83 |
| | | Infective Hepatitis | 4.12 |
| | | Gall Bladder Stone Operation | 1.41 |
| | | Cirrhosis | na |
| | | Cholecystectomy | na |
| 8 | Orthopaedics | Abscess | 0.00 |
| | | Ingrowing of Toe Nail | na |
| | | CLW (Forehead) | 0.71 |
| | | Operation for Manipulation of Left Shoulder | na |
| | | Ganglion | 1.10 |
| | | Dislocation | na |
| | | Pilonidal Sinus | na |
| | | PID | 25.46 |
| | | Back Pain | 2.24 |

| | | | |
|----|--------------------------|---|-------|
| | | Injury (Upper Extremity) | 0.00 |
| | | Fracture (Upper Extremity) | 0.97 |
| | | Arthritis | na |
| | | Operation (Thumb) | na |
| | | Medial Meniscus Injury | na |
| | | Abrasion (Knee) + Injury (Muscles) | na |
| | | Injury (HNF) | 1.85 |
| | | Fracture + Head Injury | na |
| | | Fracture (HNF) | na |
| | | Accidental Injury | 4.72 |
| | | Fracture (Lower Extremity) | 6.67 |
| | | Wrist Disease | 0.00 |
| | | Necrosis of Head of Femur | na |
| | | Tibial Tubercle + Haemarthrosis | na |
| | | Joint Replacement | 5.46 |
| | | Fracture (Bilateral Pelvis) | na |
| | | Muscular Sprain | na |
| | | Ostochondritis | na |
| 9 | Breast Disease | Breast Lump | 0 |
| | | Breast Hyperplasia | na |
| | | Breast Cancer | 3.70 |
| 10 | Dermatology | Cyst | na |
| | | Lipoma | na |
| | | Depigmented Scar | na |
| | | Insect Bite + Abscess (Back) | na |
| | | Cellulitis | 4.90 |
| | | Infected Sebaceous Cyst + Cellulitis (Back) | 0 |
| | | Burns | na |
| 11 | Endocrine | Thyroidectomy | Na |
| | | Diabetic Neuropathy | Na |
| | | Myasthenia Gravis (Ocular) | Na |
| | | DM | 1.68 |
| | | DM + Hypothyroidism | Na |
| 12 | Renal | Ureteric Colic | 0.00 |
| | | Calculus | 33.78 |
| | | Pyelonephritis | Na |
| | | UTI | 3.93 |
| | | CRF | Na |
| 13 | Male Reproductive | Sterility | 0 |
| | | Prostatectomy | na |
| | | Hydrocele | na |
| | | Tumour (Spermatic Cord) | na |
| | | Epididymal Cyst (Left) | na |
| | | Phimosis | 0 |
| | | Injury | 3.59 |
| 14 | Obstetrics & Gynaecology | Para Ovarian cyst | na |
| | | Leiomyoma | na |
| | | Vulval Haematoma | na |
| | | DUB | 2.30 |

| | | |
|----|---|-------|
| | Laparotomy + Hysterectomy + Tubal ligation | na |
| | Haemoperitoneum (Ruptured Ovarian Cyst) | na |
| | Gestation | na |
| | Vaginal Hysterectomy | 10.37 |
| | Endometriosis + Pelvic pain | na |
| | Total Abdominal Hysterectomy | 3.19 |
| | Abdominal Hysterectomy | na |
| 15 | Infectious Disease | |
| | Viral Fever | 1.41 |
| | Vivax Malaria + Enteric Fever | na |
| | Acute Bacterial Infection | na |
| | Fever + Vomiting | na |
| | Enteric Fever | 3.63 |
| | PUO | 1.30 |
| | Vivax Malaria | 1.73 |
| | Falciparum Malaria | 3.49 |
| | Septicaemia + Vivax Malaria | na |
| | Enteric Fever + Falciparum Malaria | na |
| | Falciparum Malaria + Enteric Fever | na |
| | Collagen Vascular Disease | na |
| | Viral Encephalopathy | na |
| | Toxoplasmosis | na |
| | Septicemia + Pyogenic Meningitis | na |

Exhibit 17: Date of Discharge to Date of Claim Reimbursement

| Disease System | Average Time Gap | Standard Deviation | Maximum | Minimum |
|----------------------------|------------------|--------------------|---------|---------|
| CNS | 177 | 96 | 387 | 23 |
| Ophthalmology | 99 | 56 | 276 | 29 |
| ENT | 135 | 73 | 261 | 50 |
| Respiratory | 132 | 74 | 295 | 38 |
| CVS | 140 | 72 | 358 | 43 |
| GIT | 101 | 59 | 429 | 21 |
| Hepatobiliary | 129 | 54 | 247 | 50 |
| Orthopaedic | 142 | 65 | 363 | 7 |
| Breast Disease | 138.4 | 86 | 268 | 35 |
| Dermatology | 107 | 46 | 214 | 50 |
| Endocrine | 182.4 | 72 | 308 | 117 |
| Renal | 112.2 | 79 | 361 | 26 |
| Male Reproductive | 101 | 40.1 | 186 | 46 |
| Obstetrics and Gynaecology | 126 | 78 | 420 | 51 |
| Infectious Disease | 103.6 | 64.9 | 363 | 25 |

Exhibit 18: Disease System Cost Comparison - DRG and Actual

| Disease System | Average Costs DRG | Average Cost DRG - PPP | RANK DRG | Average Costs India | RANK INDIA |
|---|-------------------|------------------------|----------|---------------------|------------|
| | USD | INR (PPP=4.5) | | INR | |
| Central Nervous System | 3436 | 15461 | 3 | 23617 | 3 |
| Ophthalmology (Eye Diseases) | 1738 | 7823 | 14 | 12878 | 9 |
| Otorhinolaryngology (ENT) | 1709 | 7691 | 15 | 10297 | 12 |
| Respiratory System | 2162 | 9729 | 12 | 11480 | 11 |
| Cardiovascular System | 6022 | 27101 | 1 | 44460 | 1 |
| Gastrointestinal System | 1934 | 8701 | 13 | 9385 | 13 |
| Hepatobiliary System | 3394 | 15274 | 4 | 25397 | 2 |
| Orthopaedics (Bone diseases) | 3197 | 14388 | 6 | 19130 | 5 |
| Breast Diseases | 3175 | 14286 | 7 | 19931 | 4 |
| Dermatology (Skin Disease) | 2575 | 11585 | 11 | 8187 | 14 |
| Endocrine System | 2927 | 13170 | 8 | 13594 | 8 |
| Renal System (Kidney and Urinary Disease) | 2595 | 11678 | 10 | 16257 | 7 |
| Male Reproductive System | 2608 | 11734 | 9 | 11802 | 10 |
| Obstetric & Gynaecological Disease | 5756 | 25902 | 2 | 19020 | 6 |
| Infectious Disease | 3333 | 14999 | 5 | 7446 | 15 |
| Standard Deviation | | 5730 | | 9426 | |

Exhibit 19: Disease Subsystem Cost Comparisons - DRG and Actual

| Disease System | Disease Subsystem | Average | Average | RANK DRG | Average | RANK INDIA | |
|-------------------------------------|---------------------------------------|----------------------------------|---------------------------------------|-------------|-----------------------|---------------|----|
| | | Costs DRG USD | Cost DRG - PPP INR (PPP=4.5) | | Costs India INR | | |
| CNS | Febrile Convulsions | 1818 | 8181 | 7 | 5304 | 10 | |
| | Syncope | 1656 | 7452 | 9 | 57260 | 2 | |
| | Viral Encephalitis | 1312 | 5904 | 10 | 5935 | 9 | |
| | Tubercular Meningitis | 5370 | 24165 | 1 | 53796 | 3 | |
| | Brain tumor | 5037 | 22666 | 3 | 91025 | 1 | |
| | Vertebro Basilar Insufficiency | 2560 | 11520 | 6 | 9320 | 6 | |
| | Hypodense lesion (Brain) | 3586 | 16137 | 5 | 7006 | 7 | |
| | Epilepsy | 1673 | 7530 | 8 | 6004.5 | 8 | |
| | Cerebrovascular Accident | 5156 | 23205 | 2 | 24412 | 4 | |
| | Meningitis | 4174 | 18783 | 4 | 16315 | 5 | |
| Ophthalmology | Cryoprophylaxis | 1809 | 8140 | 8 | 1679 | 12 | |
| | Posterior Capsular Opacification | 2167 | 9751.5 | 5 | 2033 | 11 | |
| | Phacoemulsification | 1548 | 6966 | 9 | 19000 | 3 | |
| | Excision of lens | 1548 | 6966 | 9 | 1287 | 13 | |
| | Cataract with IOL | 1548 | 6966 | 9 | 14082.11 | 4 | |
| | Enucleation | 6492 | 29214 | 1 | 25438 | 1 | |
| | Keratoplasty | 2167 | 9751.5 | 5 | 14042 | 5 | |
| | Pterygium | 2382 | 10719 | 4 | 2990 | 10 | |
| | Ptosis | 1847 | 8311.5 | 7 | 8200 | 6 | |
| | Extraocular Injury | 1471 | 6619.5 | 13 | 3054.5 | 8 | |
| | Injury + Hyphema | 1475 | 6637.5 | 12 | 3037 | 9 | |
| | Proliferative Diabetic Retinopathy | 3026 | 13617 | 3 | 7838 | 7 | |
| | Coronal ulcer | 4410 | 19845 | 2 | 19431 | 2 | |
| | Otorhinolaryngology | Tympanic Membrane Perforation | 2588 | 11646 | 4 | 5968 | 13 |
| | | Viral Laryngitis | 1115 | 5017.5 | 12 | 1939 | 16 |
| Chronic Suppurative Otitis Media | | 2177 | 9796.5 | 6 | 16139.5 | 3 | |
| Acute Suppurative Otitis Media | | 1123 | 5053.5 | 11 | 9072 | 6 | |
| Adenoidectomy | | 779 | 3505.5 | 16 | 4700 | 15 | |
| Deviated Nasal Septum | | 1265 | 5692.5 | 9 | 6584 | 10 | |
| Nasal Injury | | 1379 | 6205.5 | 8 | 9678 | 5 | |
| Sinus Polyp | | 2535 | 11407.5 | 5 | 20034 | 2 | |
| Tonsillitis | | 992 | 4464 | 14 | 6186.667 | 11 | |
| Otorrhoea | | 992 | 4464 | 14 | 14996 | 4 | |
| Sinus Infection | | 1522 | 6849 | 7 | 7147 | 8 | |
| Vocal Cord Polyp | | 4700 | 21150 | 2 | 5933 | 14 | |
| Sinus Thrombosis | | 2624 | 11808 | 3 | 36944 | 1 | |

| | | | | | | |
|--------------------------------|------------------------------|----------|----------|----|----------|----|
| | Pharyngitis | 1089 | 4900.5 | 13 | 6111 | 12 |
| | CA Larynx | 5332 | 23994 | 1 | 7094 | 9 |
| | Labyrinthine disorder | 1174 | 5283 | 10 | 7227.5 | 7 |
| Respiratory System | Bronchitis | 1256 | 5652 | 7 | 3425 | 6 |
| | Broncho-pneumonia | 2898 | 13041 | 4 | 1549 | 7 |
| | COPD | 2227 | 10021.5 | 5 | 14973 | 2 |
| | LRTI | 2078 | 9351 | 6 | 5565 | 4 |
| | Pneumonia | 3928 | 17676 | 1 | 5675 | 3 |
| | Pleural Effusion | 3679 | 16555.5 | 2 | 4587 | 5 |
| | Tuberculosis | 2916 | 13122 | 3 | 20117 | 1 |
| Cardiovascular System | Chest Pain | 1348 | 6066 | 9 | 8127.6 | 9 |
| | Cardiac Arrest | 12565 | 56542.5 | 2 | 24505.2 | 6 |
| | Heart Disease | 4579 | 20605.5 | 4 | 54931 | 4 |
| | Acute Coronary Insufficiency | 4579 | 20605.5 | 4 | 88551 | 3 |
| | CCF | 4579 | 20605.5 | 4 | 12810.5 | 8 |
| | Angiography | 12074.83 | 54336.75 | 3 | 119794 | 2 |
| | IHD | 3555.059 | 15997.76 | 7 | 26209.35 | 5 |
| | CABG | 20562 | 92529 | 1 | 192423.8 | 1 |
| | Hypertension | 3039.1 | 13675.95 | 8 | 16190.4 | 7 |
| Gastrointestinal System | Vomiting + Dehydration | 1161 | 5224.5 | 23 | 3978 | 20 |
| | Intestinal Colic | 1554 | 6993 | 16 | 5135 | 16 |
| | Diarrhoea + Vomiting | 1193 | 5368.5 | 20 | 3241.8 | 21 |
| | IBS | 2118 | 9531 | 10 | 5610 | 15 |
| | AGE + Complications | 1586 | 7137 | 14 | 4975.235 | 17 |
| | Ulcerative Colitis | 2598 | 11691 | 9 | 20234 | 2 |
| | Tuberculosis of colon | 3749 | 16870.5 | 4 | 7511 | 10 |
| | Intestinal Amoebiasis | 1193 | 5368.5 | 20 | 7305 | 12 |
| | AGE | 1286.926 | 5791.167 | 19 | 6915.667 | 14 |
| | Hernia | 1656.25 | 7453.125 | 13 | 17988.13 | 3 |
| | Food Poisoning | 3749 | 16870.5 | 4 | 1067 | 23 |
| | Peritonitis | 3749 | 16870.5 | 4 | 27028 | 1 |
| | Abdominal Trauma | 11357 | 51106.5 | 1 | 4348 | 18 |
| | Peptic Ulcer | 1457 | 6556.5 | 17 | 4300 | 19 |
| | Mouth Swelling | 2038 | 9171 | 11 | 1287 | 22 |
| | Excision | | | | | |
| | Laparoscopy | 4616 | 20772 | 3 | 9144 | 7 |
| | Abdominal Pain | 1193 | 5368.5 | 20 | 7306.667 | 11 |
| | Pancreatitis | 1955 | 8797.5 | 12 | 8329 | 9 |
| | Gastritis | 1373.5 | 6180.75 | 18 | 11466 | 6 |
| | Appendectomy | 2883.438 | 12975.47 | 8 | 17760.94 | 4 |
| | Piles | 1560 | 7020 | 15 | 8598.2 | 8 |
| | GERD | 8119 | 36535.5 | 2 | 7019 | 13 |
| | Left Inguinal Glands | 3401 | 15304.5 | 7 | 17234 | 5 |
| Hepatobiliary System | Viral Hepatitis | 1471 | 6619.5 | 7 | 6182.2 | 7 |
| | Acute Cholecystitis | 1696 | 7632 | 6 | 32789 | 3 |
| | Pancreatitis | 1955 | 8797.5 | 5 | 28441.75 | 4 |
| | Infective Hepatitis | 4544 | 20448 | 3 | 20812.8 | 6 |
| | Gall Bladder Stone | 5894 | 26523 | 1 | 28237 | 5 |

| | | | | | | |
|-----------------------|---|-------|---------|----|----------|----|
| | Operation | | | | | |
| | Cirrhosis | 4339 | 19525.5 | 4 | 60310 | 2 |
| | Cholecystectomy | 5894 | 26523 | 1 | 90325 | 1 |
| Orthopaedics | Abscess | 1652 | 7434 | 19 | 3638 | 20 |
| | Ingrowing of Toe Nail | 1892 | 8514 | 12 | 2473 | 24 |
| | CLW (Forehead) | 239 | 1075.5 | 26 | 3715 | 19 |
| | Operation for Manipulation of Left Shoulder | 4465 | 20092.5 | 8 | 3326 | 22 |
| | Ganglion | 1501 | 6754.5 | 22 | 7062 | 12 |
| | Dislocation | 2707 | 12181.5 | 9 | 15261 | 8 |
| | Pilonidal Sinus | 1567 | 7051.5 | 21 | 4862 | 17 |
| | PID | 4741 | 21334.5 | 6 | 11709 | 9 |
| | Back Pain | 1706 | 7677 | 18 | 10233.4 | 10 |
| | Injury (Upper Extremity) | 2108 | 9486 | 11 | 5372 | 16 |
| | Fracture (Upper Extremity) | 941 | 4234.5 | 24 | 6096.9 | 14 |
| | Arthritis | 4474 | 20133 | 7 | 8392 | 11 |
| | Operation (Thumb) | 1881 | 8464.5 | 14 | 3589 | 21 |
| | Medial Meniscus Injury | 1603 | 7213.5 | 20 | 15940 | 7 |
| | Abrasion (Knee) + Injury (Muscles) | 1789 | 8050.5 | 16 | 2033 | 25 |
| | Injury (HNF) | 1892 | 8514 | 12 | 5935 | 15 |
| | Fracture + Head Injury | 8739 | 39325.5 | 2 | 16034 | 6 |
| | Accidental Injury | 1804 | 8118 | 15 | 24158.43 | 4 |
| | Fracture (Lower Extremity) | 5838 | 26271 | 5 | 29845.11 | 3 |
| | Wrist Disease | 1734 | 7803 | 17 | 4145.25 | 18 |
| | Necrosis of Head of Femur | 7304 | 32868 | 4 | 134960 | 1 |
| | Tibial Tubercle + Haemarthrosis | 2288 | 10296 | 10 | 6760 | 13 |
| | Joint Replacement | 12234 | 55053 | 1 | 113953.4 | 2 |
| | Fracture (Bilateral Pelvis) | 7846 | 35307 | 3 | 20284 | 5 |
| | Muscular Sprain | 524 | 2358 | 25 | 1846 | 26 |
| | Ostochondritis | 1287 | 5791.5 | 23 | 3042 | 23 |
| Breast Disease | Breast Lump | 1824 | 8208 | 2 | 9157.5 | 2 |
| | Breast Hyperplasia | 1222 | 5499 | 3 | 7708 | 3 |
| | Breast Cancer | 4338 | 19521 | 1 | 28372.75 | 1 |
| Skin Disease | Cyst | 1267 | 5701.5 | 6 | 3852 | 7 |
| | Lipoma | 1267 | 5701.5 | 6 | 4550 | 5 |
| | Depigmented Scar | 2623 | 11803.5 | 3 | 7540 | 2 |
| | Insect Bite + Abscess (Back) | 2623 | 11803.5 | 3 | 4595 | 4 |
| | Cellulitis | 1484 | 6678 | 5 | 15862.33 | 1 |
| | Infected Sebaceous Cyst + Cellulitis | 4485 | 20182.5 | 2 | 4108 | 6 |
| | Burns | 5634 | 25353 | 1 | 5532 | 3 |

| | | | | | | | |
|--|----------------------------------|-------------------------------|----------|---------|----------|-------|---|
| Endocrine Disease | Thyroidectomy | 5538 | 24921 | 1 | 25346 | 1 | |
| | Diabetic Neuropathy | 3652 | 16434 | 3 | 17807 | 2 | |
| | Myasthenia Gravis (Ocular) | 3871 | 17419.5 | 2 | 12094 | 3 | |
| | DM | 2391.5 | 10761.75 | 4 | 11785.38 | 4 | |
| Renal System | Ureteric Colic | 1445 | 6502.5 | 5 | 6423.667 | 5 | |
| | Calculus | 2833.1 | 12748.95 | 3 | 20513.4 | 2 | |
| | Pyelonephritis | 3636 | 16362 | 2 | 6975 | 4 | |
| | UTI | 1894 | 8523 | 4 | 14309.33 | 3 | |
| | CRF | 6829 | 30730.5 | 1 | 24155 | 1 | |
| Male Reproductive | Sterility | 1029 | 4630.5 | 5 | 10471 | 5 | |
| | Prostatectomy | 2558 | 11511 | 4 | 40903 | 1 | |
| | Hydrocele | 740 | 3330 | 7 | 7305 | 6 | |
| | Tumour (Spermatic Cord) | 2790 | 12555 | 3 | 15131 | 3 | |
| | Epididymal Cyst (Left) | 1029 | 4630.5 | 5 | 12238 | 4 | |
| | Phimosis | 3135 | 14107.5 | 2 | 3411.2 | 7 | |
| | Injury | 3566 | 16047 | 1 | 15863.5 | 2 | |
| | Para Ovarian cyst | 2440 | 10980 | 6 | 10728 | 6 | |
| | Female Reproductive | Leiomyoma | 5530 | 24885 | 4 | 17377 | 4 |
| Vulval Haematoma | | 1853 | 8338.5 | 8 | 3415 | 10 | |
| DUB | | 748 | 3366 | 10 | 10162.38 | 7 | |
| Laparotomy + Hysterectomy + Tubal ligation | | 7929 | 35680.5 | 3 | 13783 | 5 | |
| Haemoperitoneum (Ruptured Ovarian Cyst) | | 5080 | 22860 | 5 | 10028 | 8 | |
| Gestation | | 1829 | 8230.5 | 9 | 9877 | 9 | |
| Vaginal Hysterectomy | | 9131 | 41089.5 | 2 | 24712.86 | 3 | |
| Endometriosis + Pelvic pain | | 2050 | 9225 | 7 | 28661 | 2 | |
| Total Abdominal Hysterectomy | | 10759 | 48415.5 | 1 | 30731.33 | 1 | |
| Infectious Disease | | Viral Fever | 1194 | 5373 | 11 | 5257 | 9 |
| | | Vivax Malaria + Enteric Fever | 4301 | 19354.5 | 2 | 7965 | 4 |
| | Acute Bacterial Infection | 4301 | 19354.5 | 2 | 5683 | 8 | |
| | Fever + Vomiting | 2297 | 10336.5 | 8 | 10387 | 2 | |
| | Enteric Fever | 2464.615 | 11090.77 | 7 | 7316.513 | 7 | |
| | PUO | 4301 | 19354.5 | 2 | 8323.6 | 3 | |
| | Vivax Malaria | 4301 | 19354.5 | 2 | 7512.931 | 6 | |
| | Collagen Vascular Disease | 2269 | 10210.5 | 9 | | | |
| | Viral Encephalopathy | 4301 | 19354.5 | 2 | 17876 | 1 | |
| | Toxoplasmosis | 2269 | 10210.5 | 9 | 2400 | 10 | |
| | Septicemia + Pyogenic Meningitis | 8834 | 39753 | 1 | 7694 | 5 | |

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