



COST DRIVER ANALYSIS
OF PROVINCIAL
DRUG PLANS

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BRITISH COLUMBIA

1990/91 - 1998/99

Federal/Provincial/Territorial

Working Group on Drug Prices

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EXECUTIVE SUMMARY

- The Federal Provincial Territorial (F/P/T) Task Force on Pharmaceutical Prices¹ was established to examine pharmaceutical pricing issues facing provincial drug plans and Canadians in general.
- This Study is an update which reports on pharmaceutical cost drivers in British Columbia's Pharmacare program over the period 1990/91 to 1998/99.
- An examination of cost drivers, produced by the Patented Medicine Prices Review Board (PMPRB) on behalf of the F/P/T Working Group on Drug Prices, provides both public and private drug plan managers, policy makers and other stakeholders, including consumers, with a better understanding of the major components that influence annual changes in pharmaceutical spending.
- The focus of the report was to disaggregate annual changes in the cost of drugs into five separate components: price effect, volume effect, entry of new drugs, exiting drugs and others. A further break out of cost drivers was done by therapeutic class, novelty and patent status.
- The British Columbia Pharmacare program has undergone major changes since its implementation with a view to contain increasing drug costs and encourage cost-effective prescribing. The implementation of the Low Cost Alternative Drug Program in 1994, and the Reference Based Pricing policy in 1995, coincided with the slower growth in Pharmacare expenditures over the period 1994/95 to 1996/97.
- Between 1990/91 and 1998/99 total drug expenditures increased by \$159 million (for plans under consideration). On average, between 1990/91 and 1998/99 per unit price changes seen by the province were responsible for -17% of the expenditure change, volume change or utilization was responsible 79%, entry of new drugs were responsible for 43%, and both exiting drugs and other factors were responsible for -1% and -5% of expenditures changes respectively. The findings suggest that utilization and entry of new drugs accounted for the largest increase in expenditures over the period with expenditures rising significantly despite a decrease in the average per unit price. The contribution of each of these factors change dramatically in 1994/95 with varying values from year to year; further work is required to understand the sensitivity of the model, the impact of cost containment policies and the entry and market penetration of new drug therapies.
- In 1998/99, drugs that existed in 1990/91 and newer drugs (drugs that were introduced after 1990/91) accounted for 34% and 66% respectively, in total drug expenditures.

¹ Presently known as F/P/T Working Group on Drug Prices

- In 1990/91, the proportion of total expenditures accounted for by patented drugs was 38%, by 1998/99, patented drugs accounted for 52% of total expenditures and were responsible for approximately 75% of the total increase in expenditures for the period under consideration.
- Among patented medicines, category 3 drugs made up the largest share of total patented drug expenditures. In 1998/99, drugs categorized as having little, moderate or no improvement (category 3) accounted for 56% of total patented drug expenditures. The share of line extension (category 1) and break through or substantial improvement (category 2) drugs were 31% and 8%, respectively.
- The study found that sales of new drugs increased rapidly in the first few years. For example, expenditures on new drugs during their first full year on the market grew, on average, by 43%. The growth in sales fell each year as the newer drugs matured. By the 4th-5th year, expenditures on these drugs fell, on average, by 5%. The tapering off in sales of these drugs occurred because new drugs continued to enter the market and experience high growth rates.
- In 1998/99, drugs in seven Anatomical Therapeutic Chemical (ATC) groups, including, Cardiovascular Systems, Central Nervous System, Alimentary Tract and Metabolism, Respiratory System, General Anti-infectives, Blood and Blood Forming Organs and Sensory Organs accounted for \$290.9 million or 91% of total expenditures.
- It was shown that over the period 1990/91 to 1998/99, drugs in the Cardiovascular Systems group contributed to the largest increase in the share of drug expenditures at 36%. This group was followed by drugs in the Nervous System, at 33%, Alimentary Tract and Metabolism at 11%, Respiratory Systems at 7% General Anti-infectives at 3%, Blood and Blood Forming Organs at 3% and Sensory Organs at 1%.
- In order to identify which disease groups are contributing proportionately more to increases in pharmaceutical expenditures, the analysis was broken down to the second level of their ATC classification. The study revealed that Lipid Reducing Agents in Cardiovascular Systems had the highest contribution to percentage increases in expenditures over the period 1990/91 to 1998/99. The second largest contributor was Psychoanaleptics in Central Nervous Systems followed by Agents Acting on the Renin-Angiotensin System in the Cardiovascular Systems. Each of these disease groups contributed 17%, 14%, and 12%, respectively, to increases in pharmaceutical expenditures over the period 1990/91 to 1998/99.

COST DRIVER ANALYSIS OF PROVINCIAL DRUG PLANS

BRITISH COLUMBIA 1990/91-1998/99

1.0 Introduction

In April 1997, the Task Force on Pharmaceutical Prices² prepared an overview paper which provided a description of the pharmaceutical sector in Canada, price and expenditure trends, and existing mechanisms used by private and public payers for regulating and/or influencing pharmaceutical prices.

The Task Force on Pharmaceutical Prices has made progress in the following areas:

- price trend analyses for the period 1990 to 1997 for prescription drug products covered by six provincial drug plans;
- an analysis of the relationship between price levels of generic and brand name drugs over the period 1990 to 1997;
- international price comparisons for the 1996 top selling non-patented single source drug products;
- comparisons of prices of non-breakthrough or non-substantial improvement (category 3) patented drugs introduced in 1995 and 1996 to other medicines in their therapeutic class; and,
- a comparison of prescription drug prices in six provincial drug plans (1990-1997).

This study updates a report on cost drivers of total pharmaceutical spending in British Columbia's Pharmacare program over the period 1990/91 to 1998/99³. Information on prices, quantities, total expenditures and market shares were obtained from the British Columbia Pharmacare database. Health Canada's Drug Product database was used to ensure that only those drugs defined by the *Food and Drug Act* were included. The Drug Product database was also used to identify all drug products by their respective ATC classification. Finally, the Patented Medicine Prices Review Board database was used to group drugs according to patent status and category.

² The Task Force has representatives from British Columbia, Alberta, Saskatchewan, Manitoba, Ontario, Nova Scotia, Health Canada and the Patented Medicine Prices Review Board. It was established to examine one of six pharmaceutical issues identified at the April, 1996 meeting of F/P/T Ministers of Health. The other issues included utilization, marketing, wastage, consumer education and research and development. The work is overseen by the Pharmaceutical Issues Committee, which reports to the Advisory Council on Health Services (ACHS).

³ The previous study was conducted on a calendar basis and price was calculated at the din level. This study is based on a fiscal year and price is calculated at the chemical level, ie. price for a chemical with an identical ingredient, strength, route, schedule and form. This change in definition was adapted in order to more fully capture the substitution within multi-source markets and refine the definition of a new drug.

The report is divided into the following sections: section 2 describes why a study of cost drivers provides important information to all stakeholders in the health care sector; section 3 describes the focal points of the cost driver analysis; section 4 reports on the growth of total drug costs in public and private drug plans for British Columbia over the period 1990/91 to 1998/99; section 5 presents the findings followed by a conclusion in section 6.

2.0 Why Study Cost Drivers?

An examination of cost drivers provides both public and private drug plan managers, policy makers and other stakeholders including consumers with a better understanding of the major components that influence annual increases and trends in pharmaceutical spending. During the 1990's, increases in the annual cost of drugs in Canada was, on average, approximately 10% per year⁴. This growth in total spending was occurring while average annual increases in overall prescription prices was less than 3%⁵. This demonstrates that changes in annual costs of pharmaceuticals are reflective of a combination of many factors. These factors are summarized in Figure 1.⁶

Figure 1

Factors Affecting Total Drug Expenditures	
1.	Changes in the total population
2.	Changes in the demographics and health status of the population (i.e. towards those with increased medication needs)
3.	Changes in the unit prices of drugs (both patented and non-patented)
4.	Changes in retail and wholesale mark-ups, and dispensing fees
5.	Changes in the prescribing habits of physicians (i.e. from older, less expensive medications to newer, relatively more expensive medications [\pm improved therapeutic effect] to treat the same underlying diagnosis)
6.	Changes in utilization of drugs on a per patient basis (i.e. more medications per patient per year)
7.	Trends towards using drug therapy instead of other treatments (e.g. as alternatives to surgery in some cases)
8.	New diseases to be treated and old diseases to be treated or better treated
9.	Extended patent protection, barriers to entry and reduction in competition

While it is difficult to quantify the relative effect that the above factors⁷ may have on increases in drug costs, some studies have attempted to do so.⁸ These studies have employed different methodologies to assess the impact of the different factors. The main findings from these studies are that price changes represent only one factor which influence changes in the total cost of

⁴ 1994 and 1996 had exceptionally low growth rates of approximately 3%

⁵ Statistics Canada, CANSIM, Series P200202

⁶ This figure was partly reproduced from the PMPRB's Discussion Paper, "*Examining the Role, Function and Methods of the Patented Medicine Prices Review Board.*", November 1997.

⁷ Another factor worth mentioning is the shift to community care over the last several years. In addition to replacing surgery, community based drug plans are experiencing utilization increases because more treatment is taking place in the community, that previously may have required hospitalization. An example of this trend is the growth in community based palliative care.

⁸ See for example Green Shield Canada "*A Report on Drug Costs*", 1994; Gorecki, P.K., "*Controlling Drug Expenditures in Canada, The Ontario Experience.*", 1991; Angus, D.E. et al. "*Sustainable Health Care for Canadians, 1995*"; and, Brogan Inc. (1998) "*Handbook on Private Drug Plans: 1993 - 1996.*"

drugs. Other important factors include utilization (i.e. changes in the amount of drugs consumed), therapeutic shifts, prescribing patterns and the influence from the introduction of new drugs.

3.0 Focus of Report

This analysis attempts to break out annual changes in the cost of drugs into the following major components:

- annual volume (utilization) changes of older and newer drugs;
- annual price changes of older and newer drugs⁹;
- annual influence from the introduction of new drugs (patented and non-patented); and,
- annual influence of newer drugs by therapeutic class or disease groups.

This analysis provides some insight into several factors outlined in Figure 1. Each of these factors is examined to assess their individual influence on annual drug cost changes. In other words, an evaluation of what percentage of the increase in annual cost of drugs is attributed to each of the above components will be done¹⁰. It is important to note that a more detailed review of price levels (rather than annual price change), substitution of older drugs; rapid market penetration of new drugs and trends in treatment costs are areas that need to be considered in much greater detail in further research and analysis.

A further disaggregation of cost drivers by therapeutic class allows an investigation of whether certain disease groups are experiencing proportionately greater increases in annual costs. Furthermore, an investigation of the extent to which new drugs are being substituted for older drugs and the relative cost of new drugs to older drugs can be done. Finally, breaking out the drugs into patented and non-patented drugs allows us to examine drugs by therapeutic novelty. In other words, to what extent is the introduction of new patented drugs that are line extensions (category 1), breakthrough or substantial improvement drugs (category 2) or, moderate, little or no improvement drugs (category 3) influencing annual changes in drug costs.

⁹ New drugs are defined at the chemical, dose, form and route level. Generic bioequivalent products are not considered as new drugs in the major component decomposition.

¹⁰ See Appendix 1 for methodology details and methodological and definitional changes from previous cost driver studies.

4.0 Trends in British Columbia Drug Expenditures

The British Columbia Pharmacare Program has undergone some major changes over the past few years and hence, a brief description of the Pharmacare Program may be helpful.

4.1 General Information

The British Columbia Pharmacare program was implemented on January 1, 1974. Pharmacare is administered by the Ministry of Health and Ministry Responsible for Seniors under the authority of the Pharmacists, Pharmacy Operations, and Drug Scheduling Act. All permanent residents of British Columbia are covered under Pharmacare. There are several plans under the program which contain various eligibility criteria. See Appendix 2 for detailed information on eligibility. The program covers prescription medications; designated permanent prosthetic appliances; insulin, syringes, needles for insulin dependent diabetics; ostomy supplies; mastectomy supplies; blood glucose testing strips for eligible holders of Certificates for Training; and orthotic bracing for children 19 years of age and under.

4.2 Major Changes Since Implementation of Pharmacare

- In 1977 the program expanded to include benefits beyond prescription drugs and beneficiaries beyond seniors and social assistance recipients
- In 1978 long-term care facility residents were added.
- Co-payments for seniors began in 1987.
- In 1988, a \$2000 maximum limit was placed on what an individual or family would pay.
- The annual deductible was increased to \$300 in 1988; \$325 in 1989; \$375 in 1991; \$400 in 1992; \$500 in 1993; \$600 in 1994; and \$800 in 1998.
- The Rural Incentive and Product Incentive Programs were introduced in 1990. The Product Incentive Program was phased out in 1994. The Rural Incentive program still remains.
- The Low Cost Alternative Drug Program was introduced in 1994.
- PharmaNet, a secure network linking pharmacies through a central database, was introduced in 1995. PharmaNet allows eligibility for recipients to be determined automatically.
- Reference-Based Pricing was introduced in 1995 for specific therapeutic categories of drug products. Pharmacare reimburses other drugs in the category up to the level of the reference-based product.

The implementation of the Low Cost Alternative Drug Program in 1994, and Reference- Based Pricing in 1995 coincided with the slower growth in Pharmacare expenditure over the period 1994 to 1996.

4.3 Cost Reimbursement of Prescription Drug Products

While on average manufacturers prices account for almost 65% of the retail price of a prescription drug, changes in retail and wholesale mark-ups and dispensing fees are also important components which contribute to changes in total pharmaceutical expenditure. Table 1 shows the changes in wholesale mark-ups and dispensing fees in Plan A and Plan C¹¹ from 1990/91 to 1998/99. Retail fees are not applicable for British Columbia Pharmacare.

Table 1

Year	Wholesale Mark-ups and Dispensing Fees British Columbia: 1990-1998		
	Wholesale Mark-up	Average Dispensing Fees	
		Plan A**	Plan C
1990	0% - 24%*	\$5.74	\$6.18
1991	0% - 24%*	\$5.78	\$6.24
1992	0% - 24%*	\$5.81	\$6.28
1993	0% - 24%*	\$5.90	\$6.35
1994	0% - 24%*	\$5.78	\$6.37
1995	0% - 9%	\$6.23	\$6.32
1996	0% - 9%	\$6.70	\$6.14
1997	0% - 7%	\$6.67	\$5.89
1998	0% - 7%	\$6.12	\$5.90

Source: Pharmacare Trends, British Columbia, 1997 and Pharmacare claims data.

* On average, wholesalers mark-up was approximately 12%

** The average dispensing fee for Plan A is an estimate due to the fact that it does not capture the fee for seniors until they go over the \$200 annual ceiling on fee.

Generally, Pharmacare reimburses pharmacists for the actual acquisition cost of the drug (the price paid by the pharmacy to the wholesaler or manufacturer). Prior to 1995, the wholesale mark-up in British Columbia varied between 0% and 24%, with an average mark-up of approximately 12%. On May 1, 1995, the maximum allowable wholesale mark-up was capped at 9%. On January 1, 1997, the maximum allowable wholesale mark-up was reduced to 7%.

After introducing the Low Cost Alternative Program, the wholesale up-charge maximum and the Reference Based Pricing policy, Pharmacare began reimbursing retail pharmacists subject to the following provisions:

- Pharmacare reimbursement is limited to the actual acquisition cost (AAC) for all drugs up to a maximum up-charge of 7% (9% in 1995 and 1996) above the manufacturers' list price. Reimbursement is based on best available price for a few high volume products.

¹¹ Expenditures in Plan A and Plan C was, on average, 75% of total Pharmacare Expenditure from 1990/91 to 1998/99.

- reimbursement is further limited to the actual acquisition cost of the lower cost alternatives for multiple source drugs;
- for all drugs falling under the Reference Based Pricing policy, reimbursement is limited to the cost of the identified reference-based product or products, a 7% max policy as well as applicable co-payments and deductibles.

Dispensing Fees - As shown in Table 1 the average dispensing fee in Plan A and Plan C remained fairly constant since 1990/91. Dispensing fees in Plan A increased from \$5.74 to \$6.12 per prescription between 1990/91 and 1998/99. Average dispensing fees in Plan C decreased from \$6.18 to \$5.90 per prescription over the period 1990/91 to 1998/99. The maximum dispensing fee for a regular prescription is \$7.55. The actual professional fee charged is determined by the individual pharmacy. Pharmacare will accept professional fees which do not exceed the provincial average by more than 15%.

Capitation Rates - prescription drugs intended for administration to residents of long-term care facilities are reimbursed on a capitation rate basis for pharmacy services. Service is defined in legislation and in agreement with the Pharmacare Program.

4.4 Total Retail Private and Public Drug Expenditures¹²

Since the early 1980s, drug expenditures in British Columbia, as in the rest of Canada, have been the fastest growing component of total health care spending¹³. Expenditures in British Columbia slowed down between 1994/95 and 1996/97, however, over the past two years, growth has returned to pre-94 levels. In 1997/98 expenditures grew by 9.6% and 10.2% in 1998/99. These rates are approximately twice the national average and faster than the annual rate of inflation, as measured by the Consumer Price Index (CPI) during this period.

In 1998 total retail spending on prescription drugs was \$1,041 million which was divided into public spending at \$619 million and private spending at \$422 million.¹⁴ The provincial drug plan portion or public (Expenditures of Prescription Drug Plans) portion was \$431 million or 70% of total public expenditures in 1998. Public (other) comprises the remaining 30% or \$187 million, which represents drug expenditures in hospitals and federal programs. Total retail spending, i.e. public and private spending including OTC drugs, was \$1,418 million in 1998. Total spending (public and private) on prescription drugs was 73% of total retail spending.

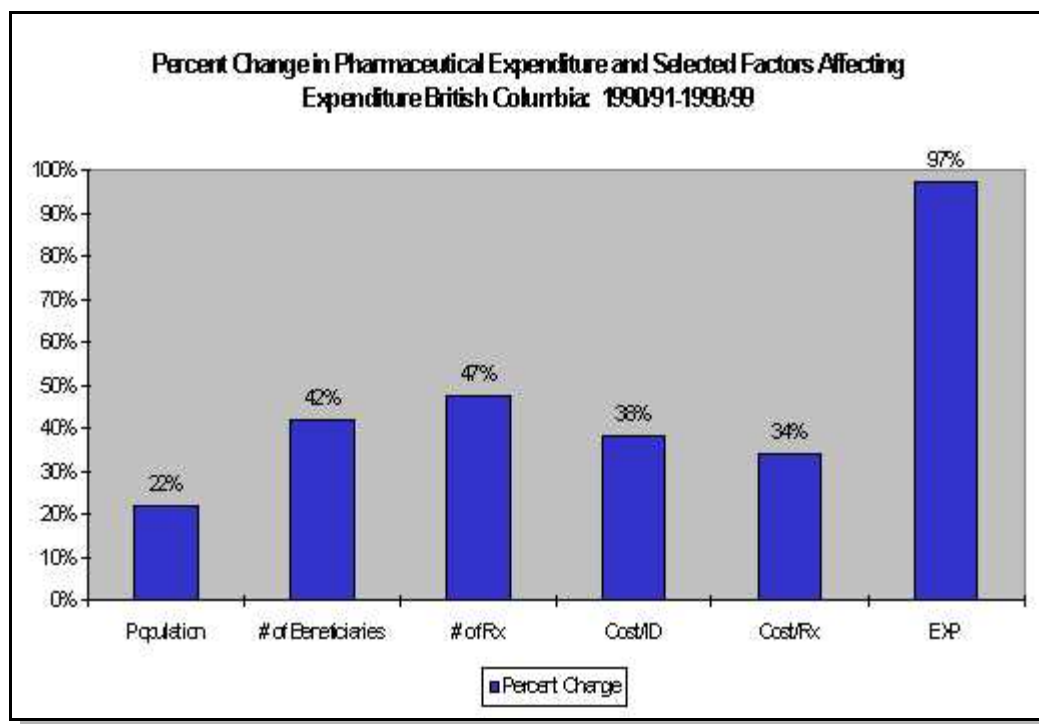
Over the years, the share of total public spending as a part of total spending has fallen. In 1990, total public spending accounted for 62% of total spending. In 1998, total public spending

¹² The figures used in this section are based on Health Canada and CIHI numbers. Expenditure levels used for 1998 are preliminary estimates.

¹³ Between 1993-1999 drug expenditures in Canada represented approximately 15% of total health expenditures. (CIHI, National Health Expenditure Trends 1975-1999)

¹⁴ Private spending includes co-pays and deductibles paid by beneficiaries of provincial prescription drug plans.

accounted for 59% of total spending. Nonetheless, the share of public spending in British Columbia has remained relatively high.



4.5 Factors Affecting Pharmaceutical Expenditures

Figure 2 summarises some of the important factors described in Figure 1, which may have contributed to total pharmaceutical expenditures, over the period 1990/91 to 1998/99:

- the number of beneficiaries increased by 42%. The large increase in the number of beneficiaries over this period may be partly attributed to the introduction of the PharmaNet system in 1995.¹⁵
- the number of prescriptions increased by 47% and the average cost per prescription increased by 34%. It is important to note that many factors may influence the cost per prescription. These include: manufacturers' unit price; wholesale and retail mark-ups; changes in prescribing habits of physicians (i.e. from older less expensive therapies to newer relatively more expensive ones); the trend towards using drug therapy instead of other treatments; and,

¹⁵ The Network provides a province wide profile of individual patient medication use, eliminates the need for patients to save receipts and submit claims, and automates billing and payment processing to pharmacies.

the inclusion of new diseases to be treated and new drugs for diseases in which drug therapy was not previously available; and, changes in the size of prescriptions.

A more complete analysis is required to evaluate the separate effect that each of these factors may have on changes in annual drug expenditures.

5.0 Analysis

5.1 Drug Expenditures in British Columbia's Pharmacare: 1990/91 to 1998/99

During the period 1990/91 to 1998/99, total Pharmacare expenditures¹⁶ on drug products in British Columbia increased from \$159.8 million to \$318.6 million. This amount differs from the total Pharmacare expenditures, for the following reasons:

- drugs were only included in this analysis if they could be matched to those drugs in the Health Canada Health Protection Branch (HPB) database;
- drug expenditures in this analysis excluded the universal plan since this data was not net of co-pays and deductibles paid by the client¹⁷; and,
- the expenditure figures do not include dispensing fees and non-drug expenditures such as diagnostic test strips.

5.2 Breakdown of Changes in Expenditure by Components

The change in total annual expenditures has been broken out into the following components: Price Effect, Volume Effect, Entry of New Drugs, Exiting drugs and Others¹⁸. Table 2 summarizes the relative contribution each of the above components have on the total annual change in expenditures.

On average, between 1991/92 and 1998/99 per unit price changes seen by the province were responsible for -17% of the expenditure change, volume change or utilization was responsible for 79%, entry of new drugs was responsible for 44%, and both exiting drugs and other factors were responsible for -1% and -5% of expenditures changes. The findings demonstrate that utilization and the entry of new drugs accounted for the largest increase in expenditures over the period.

¹⁶ Total expenditure refers to Plans A, B, C, D and F.

¹⁷ Co-payment and the deductible are based on both the recognized ingredient cost and dispensing fee.

¹⁸ Others represent the cross effect of price and volume. The cross effect is an interaction term between changes in prices and changes in quantity. That is, it is a measure of the correlation between price changes and the quantity changes. If a large change in price corresponds with a large change in quantity the cross effect will be significant. The negative sign indicates that the changes are moving in opposite directions and are significant in magnitude. A negative cross effect is recorded when a large decrease in price is accompanied by a large increase in quantity, or conversely, a large increase in price is accompanied by a large decrease in quantity.

The contribution of each of these factors change dramatically in 1994/95 with varying values from year to year. The change in each of the factors corresponds to the policy changes implemented by the program in 1994.

Between 1991/92 and 1993/94 the average per unit price effect was responsible for 10%¹⁹ of the expenditure change, volume change or utilization was responsible for 66%, entry of new drugs were responsible for 26%, and both exiting drugs and other factors were responsible for -1% and -2% of expenditure changes. Between 1995/96 and 1998/99 the average per unit price effect was responsible for -24% of the expenditure change, while volume changes or utilization was responsible for 80%, entry of new drugs were responsible for 54%, and both exiting drugs and other factors were responsible for -0% and -9% of expenditures changes. The change in contribution for each component provides some insight into the trends which are largely responsible for changes in expenditures. That is, increases in the utilization of drugs and the introduction of new drugs (mainly patented category 3 or me-too drugs) are the significant cost drivers and are becoming increasingly more important in explaining the increases in expenditure changes facing the province.

Table 2 illustrates that the impact of new drugs is significant in the year of introduction, but more significant in the second year where the average percent contribution to expenditure increases from 11% to 33%.

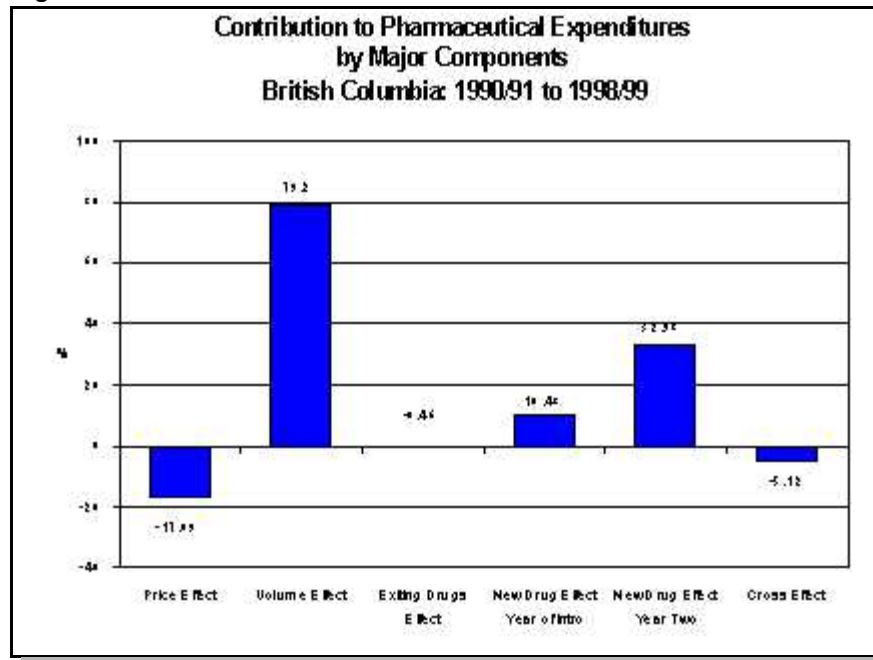
Table 2

Average Contribution to Pharmaceutical Expenditures by Major Components British-Columbia: 1990/1991 - 1998/1999						
Year	Price Effect (%)	Quantity ²⁰ Effect (Volume) (%)	Exiting Effect (%)	New Drug Effect Year of Introduction (%)	New Drug Effect Second Year (%)	Cross Effect (%)
1991/92	18.5	72.2	0	7.8	N/A	1.5
1992/93	17.1	50.9	0	6.3	25.1	0.6
1993/94	-12.5	76	-1.9	10.1	38.9	-10.5
1994/95	-1309.1	728.4	-0.8	60.3	380.7	40.5
1995/96	-10	85.7	-0.1	11.3	35.4	-22.3
1996/97	-63.1	48.5	-0.6	9.3	120.1	-14.3
1997/98	-20.4	90.4	-0.6	23.3	13.9	-6.6
1998/99	-21.2	78	-0.3	3.3	44.6	-4.5
Average	-17.1	79.2	-0.5	10.5	33	-5.12

¹⁹ It is important to note that this does not mean that prices declined by 17.1% over the time frame, a marginal decline in a popular drug may drive large negative price effects, as well, the introduction of LCA and generic substitution played an important role in reducing the cost of multiple source markets over the period of analysis.

²⁰ It is important to note that switching from older drugs to newer, more expensive therapies potentially overstates the quantity (volume) effect. Future work is required to better account for this substitution effect.

Figure 3



The findings presented above suggest that increases in utilization and coverage of new drugs significantly influence annual changes in expenditures. The expenditure decomposition provides a sense of the relative importance of changes in utilization of existing and newer drugs. It is important to keep in mind that the effects reported represent the relative impact each component had on changes in expenditure levels. The negative price effect in this analysis is greatly influenced by generic competition, which reduces the cost of the entire therapeutic class, and cost containment policies. Absolute price reductions at the DIN level, particularly of top selling newer drug products, are not the main source of the large negative price effect. Future analysis of price level of new drugs and changes in prescribing patterns toward newer therapies; changes in treatment costs and/or the price levels (rather than annual change); marketing strategies for new drugs, rate of new drug market penetration and displacement of older drugs, and impact of public policy would provide more insight into results presented above.

Table 3 breaks out annual total expenditures into “existing” drugs and “newer” drugs. Existing drugs are those drugs that were on the market in 1990/91, i.e., drugs that were introduced in 1990/91 or before. Newer drugs are those drugs that were introduced in 1991/92 or during subsequent years. It was found that expenditures on drugs that existed in 1990/91 fell by an average of 5% between 1990/91 and 1998/99, and expenditures on all drugs which includes both existing and newer drugs, increased by an average of 9% between the same time period. It can be seen from Table 1 that expenditures on all drugs had high growth rates throughout the years with the exception of period 1994/95 to 1996/97. Whereas, expenditures on existing drugs were

falling since 1994/95. As a result, the proportion of expenditures on newer drugs were increasing in total expenditures every year.

Table 3

Pharmaceutical Expenditure British Columbia: 1990/91-1998/99 (millions of dollars)						
Year	All Drugs (existing and newer drugs)			Existing Drugs in 90/91-98/99		
	Total Expenditure	Difference in Expenditure	% Growth Rates	Total Expenditure	Difference in Expenditure	% Growth Rates
1990/91	\$159.8	-	-	\$159.8	-	-
1991/92	\$195.1	\$35.3	22.1%	\$191.0	\$31.2	19.5%
1992/93	\$225.5	\$30.4	15.6%	\$208.2	\$17.2	9.0%
1993/94	\$248.0	\$22.5	10.0%	\$209.6	\$1.4	0.7%
1994/95	\$247.1	\$-0.9	-0.4%	\$168.9	\$-40.7	-19.4%
1995/96	\$256.0	\$9.0	3.6%	\$145.1	\$-23.9	-14.1%
1996/97	\$263.9	\$7.9	3.1%	\$128.6	\$-16.5	-11.4%
1997/98	\$289.2	\$25.3	9.6%	\$120.9	\$-7.7	-6.0%
1998/99	\$318.6	\$29.4	10.2%	\$109.0	\$-11.9	-9.9%

Table 3 shows that while expenditures in the “All Drug products” category were increasing on average at a rate of 9.01%, expenditures in the “Existing Drug products” category were falling on average at a rate of -5%. The growth of expenditures remained relatively small between 1994/95 and 1995/96. This coincides with the introduction of cost containment policies such as the Low Cost Alternative Program and the Reference Based Pricing Program introduced in 1994 and 1995. However, in the same years, spending on existing drug products fell by the largest amount. In 1997/98 expenditures in “All Drugs Products” category increased by 9.6% and by 1998/99 the growth rate returned to pre 1994/95 levels.

Figure 4 shows the contribution of each component in another way. As shown in Figure 4, pharmaceutical expenditures were increasing on average at a rate of 9.01% during the period 1990/91 to 1998/99. Figure 4 shows that utilization and new drugs were largely responsible for the growth in expenditures.

Figure 4

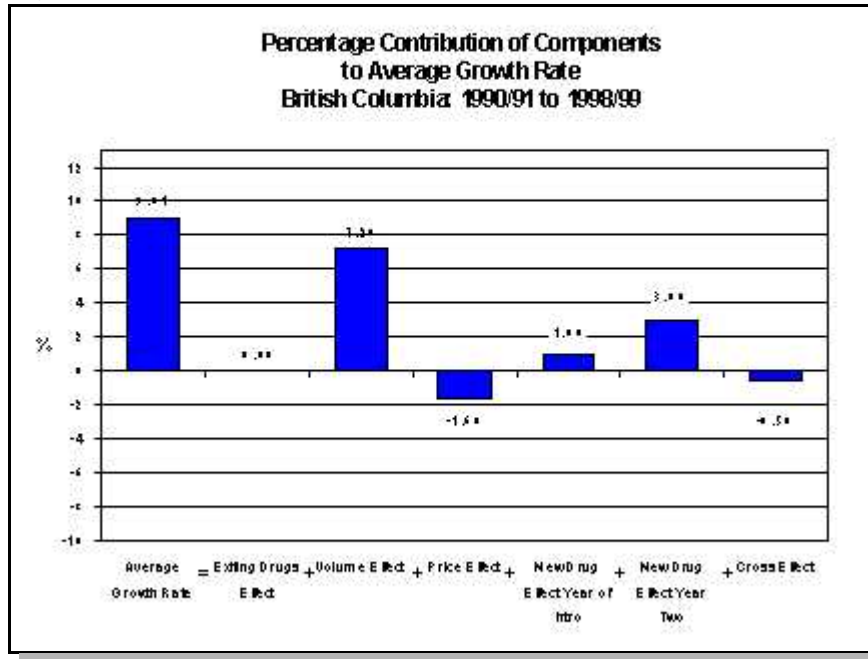


Figure 5 corresponds to Table 2; it shows the trends of expenditures on all, new and existing drug products. Expenditures on existing drug products fell by 32% between 1990/91 and 1998/99. Expenditures on new drug products, on the other hand, increased by over 5,000% and total expenditures rose by approximately 97% over the entire period of analysis.

Other than replacement of newer drug products for older drug products, the decrease in the price and utilization of existing drug products can account for some of the decrease in expenditure levels. While the number of prescriptions for older drugs decreased by 5% over the entire period, the claimed prices of older products fell by 20%; from \$25.10 in 1990/91 to \$19.96 in 1998/99 and the average period unit cost dropped from \$0.28 to \$0.23 respectively. The reverse is true for newer drugs, in 1992/93, the claimed ingredient cost of a newer prescription was \$53.17 with a corresponding per unit price of \$0.68; by 1998/99, the average claimed ingredient cost of a newer prescription was \$54.95 with a corresponding per unit price of \$0.70.²¹ By 1998/99 drugs that were introduced on to the formulary after 1990/91 accounted for approximately 40% of all prescriptions and 63% of the expenditures, up from 3.4% and 6.0% respectively in 1992/93.

²¹ These numbers are not deflated by an inflation factor.

Figure 5

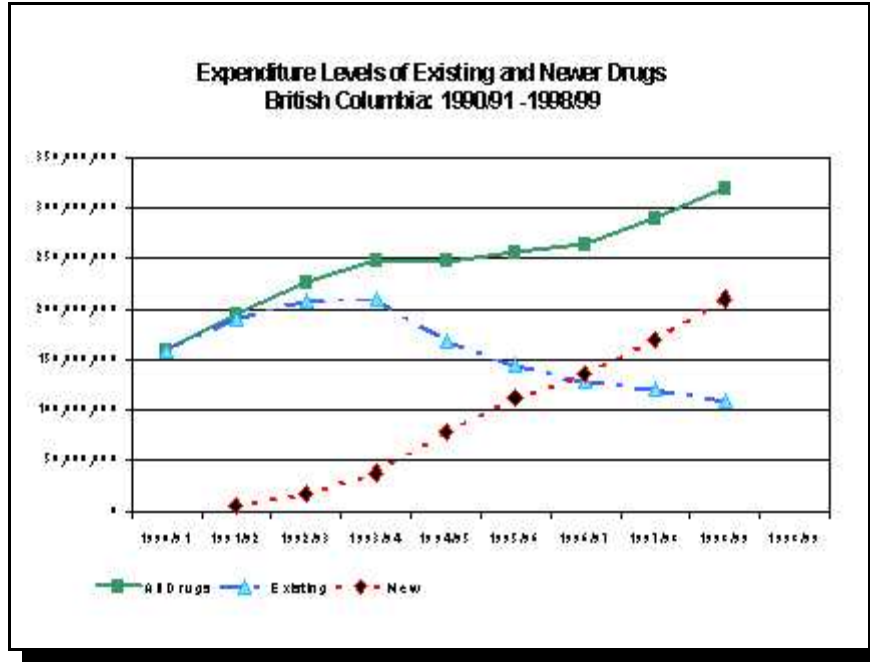
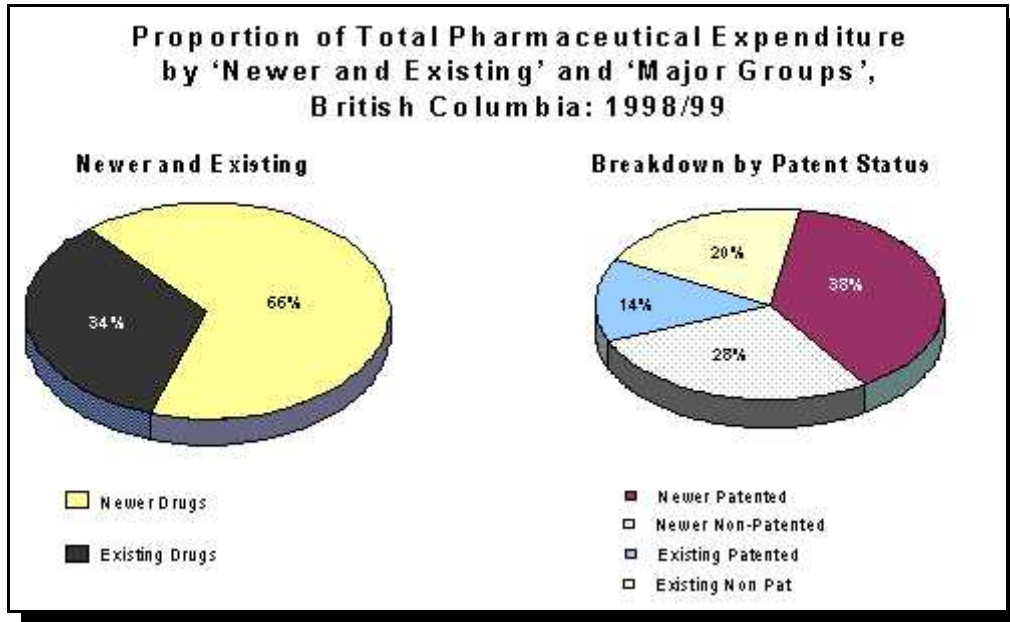


Figure 6 breaks out total pharmaceutical expenditures into patented and non-patented expenditures. In 1990/91, the proportion of patented and non-patented expenditures in total drug costs were 38% and 62%, respectively. Among patented expenditures, the proportion of older non-categorized patented drugs were 62%, category 1 drugs were 7%, category 2 drugs were 12%, and category 3 drugs were 19%. In 1990/91, the proportion of expenditures on patented drugs in total pharmaceutical expenditures was 38%. In 1998/99, the proportion of expenditures on patented drugs increased to 52%, while expenditures on non-patented drugs decreased to 48%. Among patented drugs, patented non-categorized drugs accounted for 5%, category 1 drugs accounted for 31%, category 2 drugs accounted for 8% and category 3 drugs accounted for 56% of total patented drug costs. The growth in patented drug expenditures is consistent with the impact of increased patent protection resulting from the passing of Bills C-22 and C-91 in 1987 and in 1993²².

²² This is consistent with overall growth in the share of patented drugs as reported by the PMPRB (1998). See S-9811, Trends in Patented Drug Prices.

Figure 6



5.3 Breakdown of Pharmaceutical Expenditure: (By Patent Status and Category)

Figure 7 shows the share of patented and non-patented drug products in total pharmaceutical expenditures. The patented portion is broken out into category 1 (line extensions of an existing drug product); category 2 (a breakthrough drug or substantial improvement over an existing drug product); category 3 (moderate, little or no improvement over an existing drug product) and older non-categorized patented drug products. However, it should be noted that, while the expenditures for category 1, category 2 and category 3 drug products are reported separately, they are often different brands, strengths and dosage forms of a single medicine. Category 1 products are sometimes a line extension of a category 2 or category 3 product and a category 3 drug product is often a moderate, little or no improvement over a category 2 product.²³

²³ For example, the Asthma medication Budesonide is available in many brands, strengths and dosage forms. Pulmicort Inhaler and Pulmicort Spacer, which are two different dosage forms of the brand Pulmicort, were introduced in 1988 as moderate improvements (category 3). Pulmicort Turbuhaler was introduced in 1990 as a line extension (category 1) and Pulmicort Nebuamp was introduced in 1992 as a breakthrough (category 2) product. Also, for example, Losec (20 mg/Cap) a brand of the medicine Omeprazole was introduced as a breakthrough (category 2) product in 1989. Losec (20 mg/Tab) was reintroduced in the same strength but different dosage form as a line extension (category 1) in 1996.

Figure 7

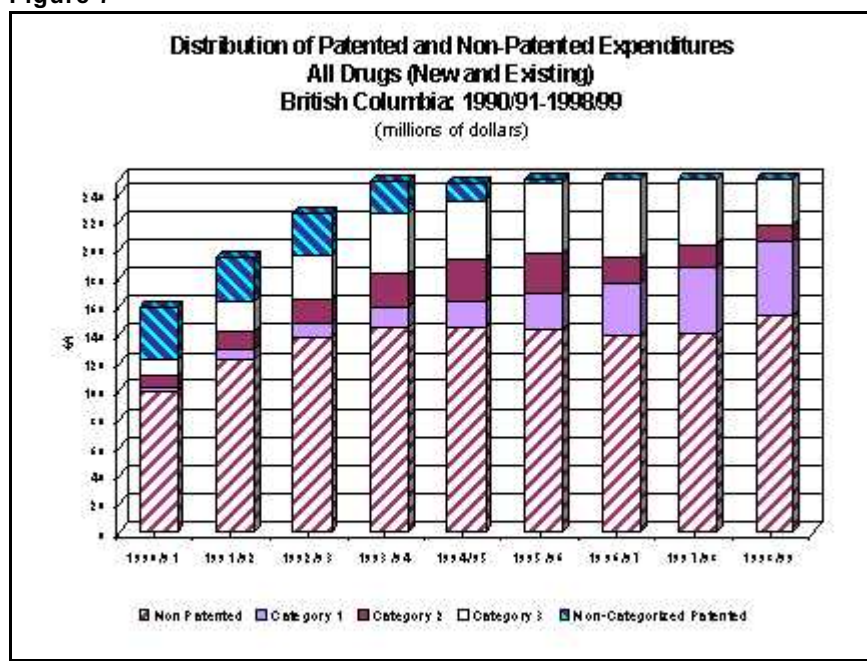


Figure 7 shows that in 1990/91 the proportion of patented and non-patented drug products in total expenditures were 38% (\$61 M) and 62% (\$ 99M) respectively. Of the 38% of expenditures accounted for by patented drug products, of category 1 drug products accounted for 7% (\$3.97 M), category 2 drug products accounted for 12% (\$7.36 M), category 3 drug products accounted for 19% (\$11.71 M), and older non categorized drug products accounted for 62% (\$37.63 M). In 1998/99, the proportion of patented and non-patented drug products in total expenditures were 52% (\$165.07 M) and 48% (\$153.57 M) respectively. Of the 52% of expenditures accounted for by patented drug products, category 1 products accounted for 31% (\$51.7 M), category 2 products accounted for 8% (\$12.4 M), category 3 products accounted for 56% (\$92.9 M), and older non-categorized patented products accounted for 5% (\$7.97 M) of total patented expenditures.

5.4 Growth of Expenditures on Newer Drug Products

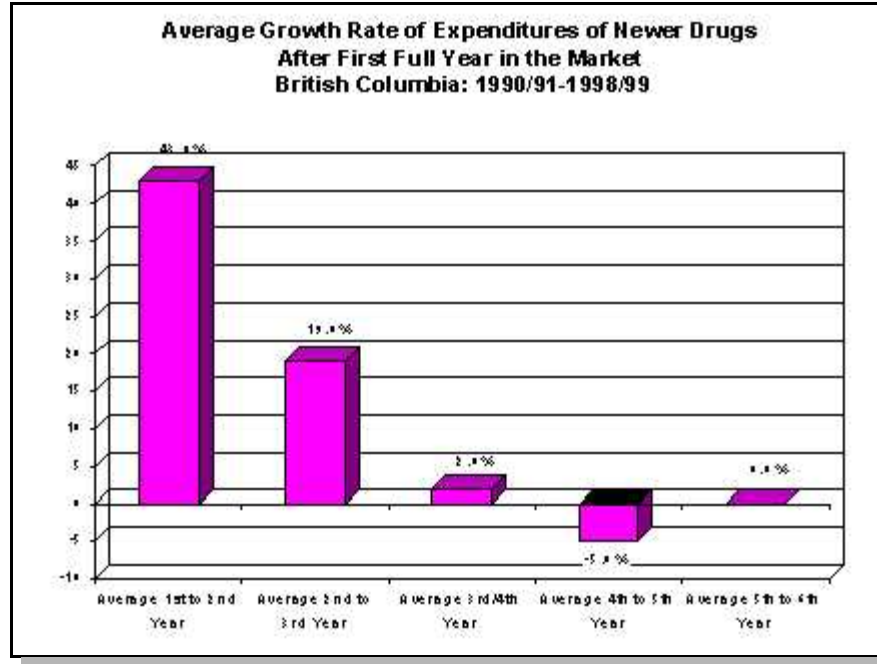
The information in Table 4 demonstrates how fast the market responds to new drugs. For example, expenditures on drugs introduced in 1991/92 were \$4.1 million and in 1992/93 increased to \$15 million. A similar increase in expenditures following the year of introduction can be observed in each of the eight years under consideration. After a sharp increase in the first two or three years, expenditures begin to fall off. However, it should be noted that, depending on the month of introduction, expenditures during the year of introduction represent expenditures of a “partial” year. For example, if a drug was introduced in July, of any year, the data on expenditures would represent expenditures for six months only.

Table 4

Expenditures on Newer Drug Products British-Columbia: 1990/1991 - 1998/1999 (millions of dollars)								
Year of Introduction	91/92	92/93	93/94	94/95	95/96	96/97	97/98	98/99
1991/1992	4.11	15.02	21.35	25.84	29.83	29.08	29.63	28.64
1992/1993	n/a	2.28	13.58	24.42	33.73	31.52	24.95	25.76
1993/1994	n/a	n/a	3.48	23.95	30.76	26.05	23.25	25.02
1994/1995	n/a	n/a	n/a	3.93	13.26	17.66	24.55	27.18
1995/1996	n/a	n/a	n/a	n/a	3.39	23.15	30.01	33.86
1996/1997	n/a	n/a	n/a	n/a	n/a	7.86	23.14	26.05
1997/1998	n/a	n/a	n/a	n/a	n/a	n/a	12.73	38.51
1998/1999	n/a	n/a	n/a	n/a	n/a	n/a	n/a	4.61
Total	4.11	17.3	38.4	78.14	110.98	135.33	168.27	209.64

In order to avoid over estimating the growth of new drugs by comparing expenditures between a “partial” year and a “full” year, the information in Table 3 is used to derive the average growth of expenditures on new drugs between each “full” year on the market, following the year of introduction.

Figure 8



In Figure 8, 1st-2nd Year represented the average growth of expenditures of new drugs between their first and second full year on the market. On average, the growth of expenditures between their first and second full year on the market was 43%. This growth fell each year, as the newer drugs matured. This is demonstrated in Figure 8 where the average growth of expenditures fell to 19% between the second and third year, 2% between the third and fourth year and eventually to -5% between the fourth and fifth year. The tapering-off of drug sales occurred because new drugs continued to enter the market and experience high growth rates.

5.5 Therapeutic Class Analysis

In order to identify which disease groups are contributing proportionately more to increases in pharmaceutical expenditures, the analysis is broken down to the second level of their ATC classifications. The second level of the ATC classification groups drugs of different pharmacological classes that have the same main therapeutic use. Sixteen therapeutic classes were identified based on their level of expenditures relative to other therapeutic classes. Table 5 shows the percentage contribution of the top sixteen therapeutic classes in total expenditures and their contribution to the changes in expenditures between 1990/91 and 1998/99.

Table 5



Percentage Contribution of Selected Therapeutic Classes to Total Expenditure British-Columbia: 1990/1991 - 1998/1999							
Therapeutic Class	Code	Contribution in 1990/91		Contribution in 1998/99		% of Total Expend. Change	Average Rate of Expend. Growth
		\$(in 000's)	% of total Expend.	\$(in 000's)	% of total Expend.		
Alimentary Tract and Metabolism	A	24,459	15.3%	42,626	13.4%	11.0%	7.0%
Antacids, drugs for treatment of peptic ulcer and flatulence	A02	17,121	10.7%	24,315	7.6%	5.0%	4.0%
Drugs used in diabetes	A10	3,642	2.3%	9,125	2.9%	3.0%	12.0%
Others		3,696	2.3%	9,186	2.9%	3.0%	12.0%
Blood and Blood Forming Organs	B	2,051	1.3%	6,865	2.2%	3.0%	16.0%
Antithrombotic agents	B01	1,865	1.2%	6,703	2.1%	3.0%	17.0%
Others		186	0.1%	162	0.1%	0.0%	-2.0%
Cardiovascular System	C	59,658	37.3%	116,365	36.5%	36.0%	9.0%
Cardiac therapy	C01	8,484	5.3%	10,823	3.4%	1.0%	3.0%
Beta blocking agents	C07	7,762	4.9%	10,090	3.2%	1.0%	3.0%
Calcium channel blockers	C08	19,315	12.1%	25,821	8.1%	4.0%	4.0%
Agents acting on the renin-angiotensin system (ACEI)	C09	11,762	7.4%	31,343	9.8%	12.0%	13.0%
Serum lipid reducing agents	C10	6,418	4.0%	34,076	10.7%	17.0%	23.0%
Others		5,917	3.6%	4,212	1.3%	-1.0%	-4.0%
General Antiinfectives for Systemic Use	J	7,535	4.7%	13,089	4.1%	3.0%	7.0%
Antibacterials for systemic use	J01	6,757	4.2%	10,558	3.3%	2.0%	6.0%
Others		778	0.5%	2,530	0.8%	1.0%	16.0%
Nervous System	N	22,935	14.4%	75,302	23.6%	33.0%	16.0%
Analgesics	N02	4,204	2.6%	9,037	2.8%	3.0%	10.0%
Antiepileptics	N03	2,288	1.4%	7,814	2.5%	3.0%	17.0%
Anti-Parkinson drugs	N04	4,564	2.9%	5,908	1.9%	1.0%	3.0%
Psycholeptics	N05	6,308	3.9%	24,347	7.6%	11.0%	18.0%
Psychoanaleptics	N06	5,141	3.2%	27,366	8.6%	14.0%	23.0%
Others		430	0.3%	831	0.3%	0.0%	9.0%
Respiratory System	R	13,043	8.2%	24,585	7.7%	7.0%	8.0%

Therapeutic Class	Code	Contribution in 1990/91		Contribution in 1998/99		% of Total Expend. Change	Average Rate of Expend. Growth
		\$ (in 000's)	% of total Expend.	\$ (in 000's)	% of total Expend.		
Anti-asthmatics	R03	11,261	7.0%	21,672	6.8%	7.0%	9.0%
Others		1,783	1.1%	2,914	0.9%	1.0%	6.0%
Sensory Organs	S	4,340	2.7%	5,933	1.9%	1.0%	4.0%
Ophthalmologicals	S01	4,061	2.5%	5,647	1.8%	1.0%	4.0%
Others		279	0.2%	286	0.1%	0.0%	0.0%
Top 16 ATCs - Level 2		120,953	75.7%	264,645	83.1%	90.5%	10.1%
Top 7 ATCs - Level 1		134,021	83.9%	284,765	89.4%	94.9%	0.1%
Total		159,816		318,635		100.0%	9.0%

It was found that the top sixteen therapeutic classes, which were approximately 15% of the total number of therapeutic classes (at second level), accounted for 80.5% of total pharmaceutical expenditures in 1998/99.

The first two columns of Table 5 show the percentage contribution of the top sixteen therapeutic classes in total expenditures, as well as the contribution of each of the seven ATC groups to which these sixteen therapeutic classes belong. The seven ATC groups are: Cardiovascular Systems, Nervous System, Alimentary Tract and Metabolism, Respiratory System, General Anti-infectives, Blood and Blood Forming Organs and Sensory Organs. Expenditures on these seven ATC groups were \$290.9 million or 91% of total expenditures in 1998/99.

Among the second level therapeutic classes, Lipid Reducing Agents in Cardiovascular Systems had the highest contribution to percentage increases in expenditures over the period 1990/91 to 1998/99. The second largest contributor was Psychoanaleptics in Central Nervous Systems followed by Agents Acting on the Renin-Angiotensin System in the Cardiovascular Systems. Each of these disease groups contributed 17%, 14%, and 12%, respectively, to increases in pharmaceutical expenditures over the period 1990/91 to 1998/99.

In 1990, Lipid Reducing Agents accounted for 4% of total expenditures; by 1998/99 it accounted for 10.7% of total expenditures. Psychoanaleptics rose from 3.2% in 1990/91 to 8.6% of total expenditures in 1998/99. Agents Acting on the Renin-Angiotensin System were 7.4% of total expenditures in 1990/91. By 1998/99 they increased to 9.8% of total expenditures.

The last column in Table 5 shows the contribution of each of the ATC groups and therapeutic classes to the total increase in expenditures between 1990/91 and 1998/99. Among the seven ATC groups, Cardiovascular System was the largest contributor to the increase in expenditures (36%), followed by Central Nervous System (33%) and Alimentary Tract and Metabolism (11%). Among the sixteen second level therapeutic classes, Lipid Reducing Agents, Psychoanaleptics and Agents Acting on the Renin-Angiotensin System were the major cost drivers. Their contributions to the increase in expenditures were 17%, 14% and 12%, respectively. Psycholeptics (11%) and Anti-asthmatics (7%) were among the other large contributors to the increase in expenditures. Nonsteroidal Anti-inflammatory drugs (NSAIDs)

and Diuretics helped reduce the increase in expenditures. Their contributions were -6% and -1%, respectively.

Table 6 reports on the average component contribution to expenditure change for the top 16 therapeutic classes. It is clear that there are significant differences between the classes. Generally speaking, the average trends reported in Table 2 are consistent with the average reported for the top 16 classes, however, there are some interesting exceptions. For instance, although price change contributes negatively to expenditure increases, i.e. price is decreasing on average over time, within Lipid reducing agents price change is responsible, on average, for 1% of the increase in expenditures. Within antibacterials for systemic use, price change is responsible, on average, for 12% of the increase in expenditure. The variations reported below suggest that therapeutic markets are different, understanding these differences and the reasons behind them is one of the future research challenges.

Table 6

Average Contribution to Pharmaceutical Expenditures by Major Components for Top 16 Therapeutic Classes British-Columbia: 1990/1991 - 1998/1999							
Therapeutic Class	Code	Average Price Effect (%)	Average Quantity Effect (%)	Average New Drug Effect (Year of Introduction) (%)	Average New Drug Effect (2nd Year) (%)	Average Exiting Drug Effect (%)	Average Cross (Other) Effect (%)
Antacids, drugs for treatment of peptic ulcer and flatulence	A02	-64	16	16	133	0	-1
Drugs used in diabetes	A10	-32	132	0	3	0	-3
Antithrombotic agents	B01	-1	86	3	9	0	4
Cardiac therapy	C01	-71	173	1	7	0	-9
Beta blocking agents	C07	-99	142	19	41	0	-3
Calcium channel blockers	C08	-109	107	8	65	0	29
Agents acting on the renin-angiotensin system	C09	-4	89	4	11	0	0
Serum lipid reducing agents	C10	1	67	6	29	0	-3
Antibacterials for systemic use	J01	12	111	13	15	-2	-49
Analgesics	N02	-7	74	9	26	0	-1
Antiepileptics	N03	-8	89	7	13	0	0
Anti-Parkinson drugs	N04	-39	35	44	55	0	6
Psycholeptics	N05	-13	63	22	27	0	1
Psychoanaleptics	N06	-15	92	5	19	0	-1
Anti-asthmatics	R03	-22	91	5	25	0	0
Ophthalmologicals	S01	-19	6	16	85	0	12
Total Average		-19	81	9	30	0	-1

Following is a detailed analysis of the impact of existing and newer drugs for Lipid Reducing Agents, Psychoanaleptics and Agents Acting on the Renin-Angiotensin System. Appendix 4 provides a detailed analysis of the remaining therapeutic classes.

Lipid Reducing Agents:

Expenditures in this therapeutic class had the largest growth rates. Table 5 shows that expenditures rose from \$6.4 million in 1990/91 to \$34.1 million in 1998/99.

In 1990/91, the proportion of expenditures on patented drugs accounted for 81% of total expenditures. Expenditures on patented products were distributed mainly between category 2 drugs, which represented 77% of patented expenditures, and older non classified drugs, which represented 19% of patented expenditures. In 1990/91 expenditures on patented category 3 drugs accounted for only 4% of total patented expenditures. By 1998/99, the proportion of expenditures on patented drugs fell to 78% of total expenditures and the proportion of expenditures on category 1 and category 3 drugs as part of total patented expenditures increased dramatically to 30% and 70% respectively.

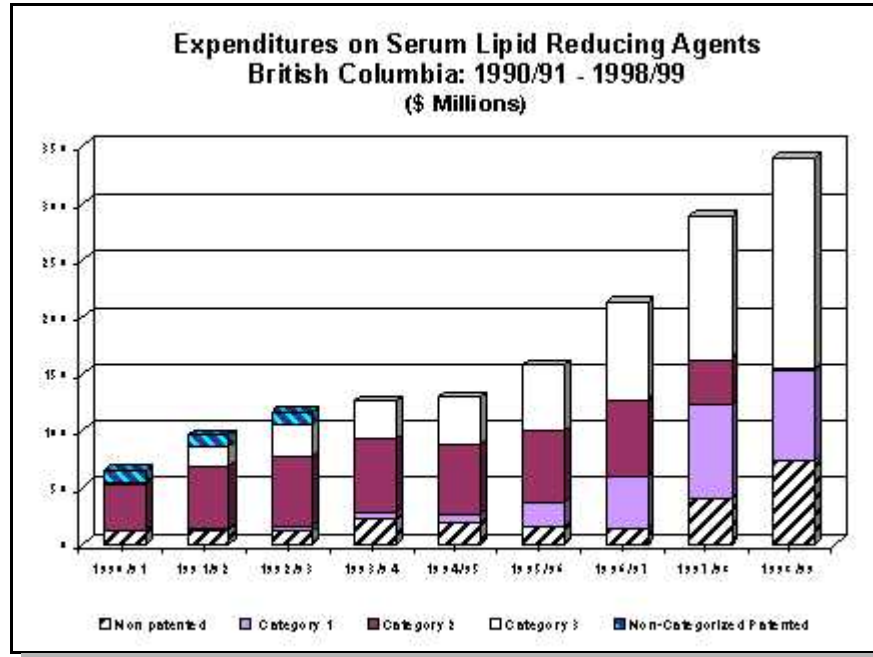
Table 7

Impact of Existing and Newer Drug Products by Major Disease Groups Serum Lipid Reducing Agents British-Columbia: 1990/1991 - 1998/1999 (thousands of dollars)										
Year of Introduction	CAT	90/91	91/92	92/93	93/94	94/95	95/96	96/97	97/98	98/99
1990/91	-	1,191	1,201	918	1,764	636	182	91	96	113
1990/91	1	29	188	368	521	637	787	1,058	880	30
1990/91	2	4,016	5,454	6,233	6,400	6,184	6,274	6,595	3,934	145
1990/91	3	218	1,632	2,786	3,386	3,978	5,393	8,034	10,696	11,035
1990/91	NC	964	1,038	1,103	0	0	0	0	0	0
1991/92	-	0	24	200	210	91	2	5	5	4
1991/92	1	0	4	18	39	43	52	79	120	132
1992/93	-	0	0	0	52	306	282	271	272	231
1993/94	-	0	0	0	224	838	630	442	347	251
1993/94	3	0	0	0	1	150	453	681	912	848
1993/94	NC	0	0	0	0	0	0	0	0	0
1994/95	-	0	0	0	0	2	234	256	205	162
1994/95	1	0	0	0	0	87	1,326	3,502	6,350	6,113
1995/96	-	0	0	0	0	0	176	220	173	133
1995/96	NC	0	0	0	0	0	0	0	0	0
1996/97	-	0	0	0	0	0	0	16	44	495
1996/97	1	0	0	0	0	0	0	86	993	1,625
1997/98	-	0	0	0	0	0	0	0	2,849	6,033
1997/98	3	0	0	0	0	0	0	0	1,036	6,650
1998/99	-	0	0	0	0	0	0	0	0	0
1998/99	3	0	0	0	0	0	0	0	0	74
1998/99	NC	0	0	0	0	0	0	0	0	0
Total Expenditures		6,418	9,541	11,627	12,597	12,952	15,790	21,335	28,911	34,076
Patented Expenditures		5,227	8,316	10,508	10,347	11,079	14,285	20,035	24,920	26,653
Non-Patented Expenditures		1,191	1,225	1,119	2,250	1,873	1,505	1,300	3,991	7,423

The top three selling drugs in this category were Provastatin, Atorvastatin and Lovastatin accounting for more than 70% of total expenditures.

Figure 9 corresponds to Table 7. It can be seen that over the years, total patented expenditures in this therapeutic class were dominated by category 3 drugs followed by category 1 drugs.

Figure 9



Psychoanaleptics:

Psychoanaleptics had the second highest contribution to the increase in total expenditures between 1990/91 and 1998/99. As shown in Table 8, total expenditures increased from \$5.1 million in 1990/91 to \$27.4 million in 1998/99. In 1990/91, 49% of the expenditures were on patented drugs. By 1998/99, patented drugs comprised 67% of total expenditures.

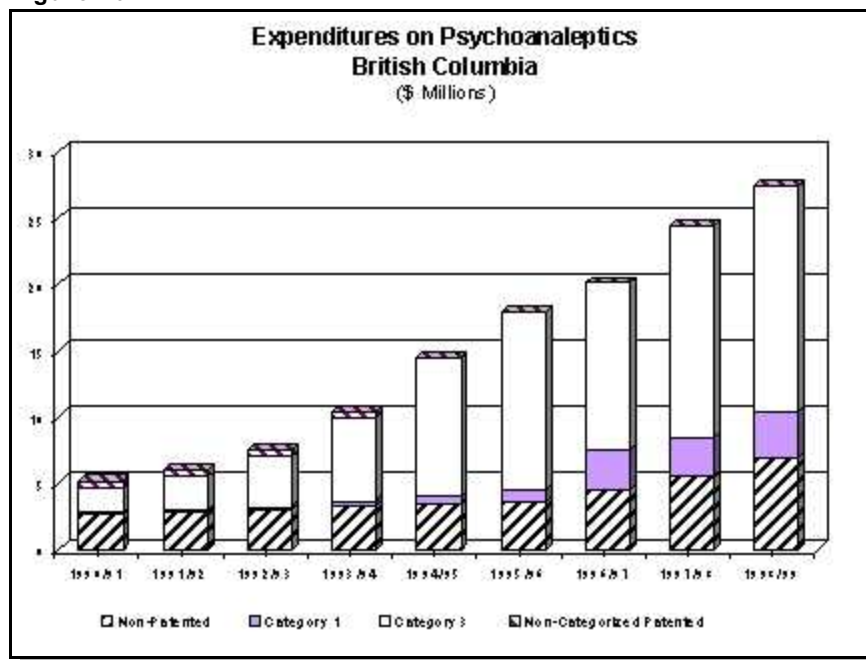
The top three selling drugs in this category were Paroxetine, Sertraline and Fluoxetine accounting for approximately 64% of total expenditures.

Table 8

Impact of Existing and Newer Drug Products by Major Disease Groups British-Columbia: 1990/1991 - 1998/1999 Psychoanaleptics (thousands of dollars)										
Year of Introduction	CAT	90/91	91/92	92/93	93/94	94/95	95/96	96/97	97/98	98/99
1990/1991	-	2,644	3,414	3,593	3,567	1,896	1,877	1,797	2,736	1,339
1990/1991	1	105	167	177	205	93	4	4	3	6
1990/1991	3	1,900	2,411	2,983	3,638	5,215	5,272	1,694	41	34
1990/1991	NC	491	0	0	0	0	0	0	0	0
1991/1992	-	0	2	11	15	35	34	34	35	29
1991/1992	3	0	8	694	1,484	2,153	2,494	2,702	3,142	3,472
1992/1993	-	0	0	46	166	407	341	282	237	192
1992/1993	3	0	0	62	535	939	1,335	1,545	1,789	1,668
1992/1993	NC	0	0	0	0	0	0	1	1	1
1993/1994	-	0	0	0	114	952	874	798	800	723
1993/1994	1	0	0	0	135	428	487	16	8	5
1993/1994	3	0	0	0	571	2,103	3,285	4,770	6,692	8,060
1994/1995	-	0	0	0	0	159	239	182	212	198
1994/1995	3	0	0	0	0	56	1,006	1,933	3,193	3,718
1995/1996	-	0	0	0	0	0	129	3,104	2,836	2,641
1995/1996	1	0	0	0	0	0	528	189	402	593
1996/1997	-	0	0	0	0	0	0	1,057	1,686	1,742
1997/1998	-	0	0	0	0	0	0	0	606	2,126
1997/1998	1	0	0	0	0	0	0	0	0	1
1997/1998	NC	0	0	0	0	0	0	0	0	0
1998/1999	-	0	0	0	0	0	0	0	0	149
1998/1999	1	0	0	0	0	0	0	0	0	664
1998/1999	3	0	0	0	0	0	0	0	0	6
Total Expenditures		5,141	6,002	7,566	10,430	14,437	17,906	20,106	24,419	27,366
Patented Expenditures		2,497	2,586	3,916	6,568	10,987	14,412	12,853	15,272	18,227
Non-Patented Expenditures		2,644	3,416	3,650	3,862	3,450	3,494	7,253	9,147	9,139

Figure 10 corresponds to Table 8. It reflects the high growth rates in expenditures in this therapeutic class. It also shows how over the years patented category 3 drugs had the largest share in total expenditures. The category 2 drugs are older drugs which were eventually replaced by new category 1 and 3 drugs.

Figure 10



Agents Acting on the Renin-Angiotensin System (ACEI):

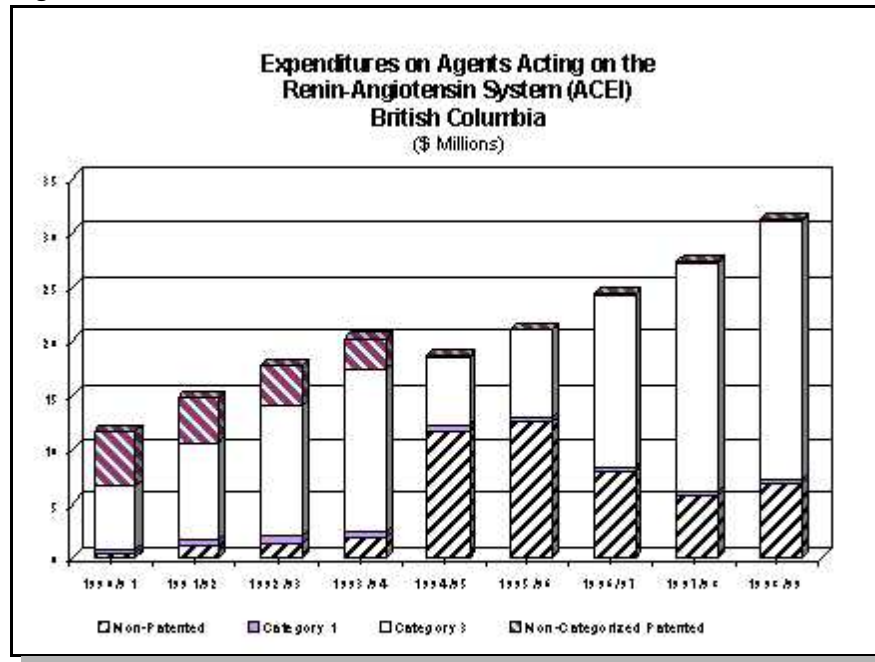
Total expenditures almost doubled since 1990/91. In 1990/91 expenditures were \$11.8 million and rose to \$31.3 million by 1998/99, despite the inclusion of Agents Acting on the Renin-Angiotensin System in the Reference Based Pricing Program in 1997. The greater proportion of expenditures were on patented drugs until 1994/95 and again in 1996/97 to 1998/99. In 1994/95 and 1995/96 the non-patented had a significant market share. The greatest increase in expenditures on patented drugs was accounted for by existing category 3 drugs in 1990/91 and new category 3 drugs introduced in 1992/93.

The top three drugs within this therapeutic class were Enapril Maleate with \$10.8 million, Quinapril at \$6.0 million and Ramipril at \$5.1 million. The total share of these three chemicals in total expenditures on Agents Acting on the Renin-Angiotensin System was 67%.

Table 9

Impact of Existing and Newer Drug Products by Major Disease Groups Agents Acting on the Renin-angiotensin System (ACEI) British-Columbia: 1990/1991 - 1998/1999 (thousands of dollars)										
Year of Introduction	CAT	90/91	91/92	92/93	93/94	94/95	95/96	96/97	97/98	98/99
1990/1991	-	352	980	980	1,102	1,747	1,463	1,360	1,230	1,379
1990/1991	1	378	672	679	625	514	428	314	254	0
1990/1991	3	5,823	8,885	11,701	13,850	4,521	5,600	12,140	13,452	13,283
1990/1991	NC	5,210	4,286	3,778	2,884	118	19	48	90	0
1991/1992	-	0	71	287	413	718	670	629	468	398
1991/1992	3	0	0	120	277	300	347	757	2,157	2,575
1992/1993	-	0	0	9	59	48	108	85	35	29
1992/1993	3	0	0	202	722	1,161	1,525	2,177	3,809	4,794
1993/1994	-	0	0	0	293	9,055	9,686	4,657	36	9
1993/1994	3	0	0	0	71	322	574	690	770	936
1994/1995	-	0	0	0	0	143	562	1,200	3,502	4,957
1994/1995	1	0	0	0	0	4	46	88	137	199
1994/1995	NC	0	0	0	0	0	0	0	0	0
1995/1996	-	0	0	0	0	0	17	52	330	347
1996/1997	-	0	0	0	0	0	0	7	77	124
1996/1997	3	0	0	0	0	0	0	141	966	1,964
1997/1998	-	0	0	0	0	0	0	0	2	3
1997/1998	3	0	0	0	0	0	0	0	26	114
1998/1999	-	0	0	0	0	0	0	0	0	3
1998/1999	1	0	0	0	0	0	0	0	0	26
1998/1999	3	0	0	0	0	0	0	0	0	205
Total Expenditures		11,762	14,895	17,755	20,297	18,650	21,045	24,340	27,342	31,343
Patented Expenditures		11,410	13,843	16,479	18,430	6,940	8,538	16,355	21,661	24,096
Non-Patented Expenditures		352	1,051	1,276	1,867	11,711	12,507	7,985	5,681	7,247

Figure 11



6.0 Conclusions

The study reports on the cost drivers of total pharmaceutical spending in British Columbia's Pharmacare program over the period 1990/91 to 1998/99.

During the period under review, expenditures increased from \$159.8 million to \$318.6 million for the plans used for analysis. Growth in spending was driven by higher utilization of existing drug products and by newer drug products introduced in 1991/92 or in subsequent years.

Between 1990/91 and 1998/99 total drug expenditures increased by \$159 million (for plans under consideration). On average, between 1990/91 and 1998/99 per unit price changes seen by the province were responsible for -17% of the expenditure change, volume change or utilization was responsible 79%, entry of new drugs were responsible for 43%, and both exiting drugs and other factors were responsible for -1% and -5% of expenditures changes respectively. The findings suggest that utilization and entry of new drugs accounted for the largest increase in expenditures over the period with expenditures rising significantly despite a decrease in the average per unit price. The contribution of each of these factors change dramatically in 1994/95 with varying values from year to year; further work is required to understand the sensitivity of the model, the impact of cost containment policies and the entry and market penetration of new drug therapies.

Change in total expenditures was broken out into five components: retail price effect, volume effect, new drug products effect, exiting drug products and others. Between 1990/91 and 1998/99 total drug expenditures increased by \$159 million.

The report also analyses the extent to which the top seven ATC groups are contributing to increases in pharmaceutical expenditures. In 1998/99, drugs in seven Anatomical Therapeutic Chemical (ATC) groups, including, Cardiovascular Systems, Nervous System, Alimentary Tract and Metabolism, Respiratory System, General Anti-infectives, Blood and Blood Forming Organs and Sensory Organs accounted for \$290.9 million or 91% of total expenditures.

The British Columbia Pharmacare Program underwent several changes since 1993/94 with a view to manage the growth in drug costs. Further analysis is necessary to fully understand the effect that those changes had on total pharmaceutical expenditures and utilization trends.

Appendix 1

Methodology

This study analyses the cost drivers in total pharmaceutical spending from 1990/91 to 1998/99 in British Columbia.

In order to conduct the analysis, information on prices, quantities and expenditures were obtained from the British Columbia Pharmacare Database. Health Canada's Drug Product Database was used to ensure that only those drugs defined by the Food and Drug Act were included. The Patented Medicine Prices Review Board data base was used to group drugs according to patent status.

Prices used in this study are claimed costs and include wholesale mark-ups but exclude dispensing fees.

This study reports expenditures by year of introduction of drugs. Year of Introduction is defined as the year of first sales recorded in British Columbia Pharmacare Database. Drugs with sales in 1990/91 or before, are termed as "existing" drugs while drugs with sales in 1991/91 and subsequent years are termed as "newer" drugs.

The study focuses on two aspects of expenditures change:

- the influence from existing drugs in terms of growth in price and quantity and exit
- the impact of new drugs in terms of replacement of older drugs

For this purpose, the annual change in pharmaceutical expenditures is broken down into five components: price effect, volume effect, entry of new drugs, exiting drugs and others. The following model was used to obtain the results.

$$TE_o = P_o Q_o \quad o = \text{base period} \dots \dots \dots (1)$$

$$\Delta TE_1 = P_1 Q_1 - P_o Q_o \quad 1 = \text{first period} \dots \dots \dots (2)$$

$$= P_o(Q_1 - Q_o) + Q_o(P_1 - P_o) + (P_1 - P_o)(Q_1 - Q_o) + P_{1n}Q_{1n} - P_o^o Q_o^o$$

Where:

TE = Total Expenditure

$P_o(Q_1 - Q_o)$ = Volume Effect

$Q_o(P_1 - P_o)$ = Price Effect

$(P_1 - P_o)(Q_1 - Q_o)$ = Interaction Term

$P_{1n}Q_{1n}$ = New Drug Expenditure Influence

$P_o^o Q_o^o$ = Exiting Drugs

$P_o(Q_1 - Q_o) + Q_o(P_1 - P_o) + (P_1 - P_o)(Q_1 - Q_o)$ = Existing Drug Influence, E_i

After the first period, 1, New drugs can be separated into Volume and Price influence on annual change in total expenditures:

$$\Delta TE = P_2 Q_2 - P_1 Q_1 \quad 2 = \text{Second Period} \dots \dots \dots (3)$$

$$= P_1(Q_2 - Q_1) + Q_1(P_2 - P_1) + (P_2 - P_1)(Q_2 - Q_1) + P_{2n}(Q_{2n} - Q_{1n}) + Q_{1n}(P_{2n} - P_{1n}) + (P_{2n} - P_{1n})(Q_{2n} - Q_{1n}) + P_{2n}^* Q_{2n}^*$$

Where,

$P_{2n}^* Q_{2n}^*$ = New Drugs in Period 2 = N_i^*

$P_1(Q_2 - Q_1)$ = New Drug Volume Influence

$Q_1(P_2 - P_1)$ = New Drug Price Influence

$(P_2 - P_1)(Q_2 - Q_1)$ = Interaction Term

$P_1(Q_2 - Q_1) + Q_1(P_2 - P_1) + (P_2 - P_1)(Q_2 - Q_1)$ = N_i , New Drug Influence

$$\therefore \Delta TE_i = E_i + \sum N_i + N_i^* \dots \dots \dots (4)$$

Divide (4) by ΔTE_i

$$\Delta TE_i / \Delta TE_i = 1 = E_i / \Delta TE_i + \sum N_i / \Delta TE_i + N_i^* / \Delta TE_i$$

Estimates the influence of each component

The previous study was conducted on a calendar basis and price was calculated at the din level, this study is based on a fiscal year and price is calculated at the chemical level, i.e. price for a chemical with an identical ingredient, strength, route, schedule and form. This change in definition was adapted in order to better capture the substitution within multi-source markets and better represent the contribution of each cost driver component in the model.²⁴

The impact of new drugs is tracked not only during the year of introduction, but also in the subsequent year. After the two periods, the effect of new drugs is recorded as part of the price, utilization and other effect.

The other major focus of the report was a breakdown of expenditures by therapeutic class and patent status over the period 1990/91 to 1998/99. This would enable us to:

- identify the extent to which each therapeutic class contributed to the increases in total Pharmacare expenditures over the period 1990/91 and 1998/99; This was done by calculating the difference between the level of expenditures of each therapeutic class between 1990/91 and 1998/99, and dividing the difference by the difference between the level of total expenditures between 1990/91 and 1998/99.
- identify the extent of substitution between new drugs and exiting drugs in each therapeutic class;
- identify the impact that category 1, 2 and 3 drugs have on the market.

²⁴ The previous version of cost drivers treated all new DIN's as new drugs, including generics.

Appendix 2

PROVINCIAL DRUG PLANS: BRITISH COLUMBIA

General Plan Information

Beneficiaries Covered

All permanent residents of British Columbia are covered under Pharmacare. There are several plans under the program which contain various eligibility:

- Plan A: seniors 65 or older.
- Plan B: residents in long term care facilities.
- Plan C: recipients of social assistance.
- Plan D: cystic fibrosis patients.
- Plan E: all other residents of British Columbia
- Plan F: medically dependent children.

Deductibles, Copayments and Professional Fees

- Plan A: Seniors pay 100% of the dispensing fee, up to a maximum of \$200 annually, with subsequent dispensing fee's paid by the province. 100% of the recognized ingredient cost is covered by Pharmacare.
- Plans B,C,D &F: Recipients under these plans pay no deductibles or copayments.
- Plan E: Recipients pay the first \$800 of their prescription costs and 30% of each prescription thereafter up to a maximum of \$2,000. Families with annual income less than \$20,000, have an annual deductible of \$600, with 100% coverage after the deductible

Pharmacare Coverage

Pharmacare covers:

- eligible drugs prescribed by a physician, dentist, or podiatrist
- insulin, needles, and syringes for diabetics
- blood glucose monitoring strips where home blood glucose monitoring has been deemed medically necessary and the individual has a valid certificate of training from an approved diabetic teaching centre. As of May 1996, there were 35,406 individuals with home blood glucose certificates of training registered with Pharmacare
- certain ostomy supplies
- designated permanent prosthetic appliances and children's orthotic devices

Pharmacare does not Cover

- bandages
- artificial sweeteners
- antacids, laxatives and other over-the-counter drugs
- wheelchairs, walkers, and other medical devices
- drug costs which have been fully reimbursed by another plan
- drugs or supplies obtained while outside of British Columbia

Appendix 3

Population Changes and Top Selling Drugs

The following table reports on population growth in British Columbia between 1990 and 1997 by age group. In 1991, the 30-39 age group represented the highest proportion of the total population, at 17.1%. This was followed by the 20-29 age group, at 15.3%, and the 40-49 age group and the 0-9 age group, at 13.7% each. In 1997, the 30-39 age group remained the largest group at 17.1% of the total population. The 40-49 age group increased to 15.6%. The 0-9 age group decreased to 12.6% and 20-29 age group decreased to 14.2% of the total population.

Between 1990 and 1997, the highest growth rate was achieved by the 80-90+ (45.8%) age group. This group was followed by 40-49 (44.2 %) and 50-59 (38.5%) age groups.

AGE GROUPS	YEAR				CHANGES 1990-1998 (in thousands)	% GROWTH 1990-1998
	1990		1998			
	Population (in thousands)	% of total	Population (in thousands)	% of total		
0-9	430.30	13.70	496.16	12.37	65.86	15.31
10-19	410.40	13.10	527.09	13.14	116.69	28.43
20-29	478.40	15.30	557.64	13.91	79.24	16.56
30-39	535.00	17.10	676.73	16.88	141.73	26.49
40-49	427.50	13.70	642.16	16.01	214.66	50.21
50-59	297.80	9.50	440.85	10.99	143.05	48.03
60-69	280.10	8.90	311.52	7.77	31.42	11.22
70-79	189.60	6.10	237.76	5.93	48.16	25.40
80-90+	82.60	2.60	120.00	2.99	37.40	45.28
Seniors(65+)	409.40	13.10	509.18	12.70	99.78	24.37
All Ages	3,131.70	100.00	4,009.92	100.00	878.22	28.04

Source: Statistics Canada, Annual Demographic Statistics, Catalogue Number 91-213-XPB, 1998.

■ Top 25 Drugs Patented and Non-Patented Drug Products British Columbia: 1998/99						
DIN	Brand	Ingredient	ATC	Year of Introduction	Expenditure 1997/98	Expenditure 1998/99
2190915	Losec 20mg	Omeprazole (omeprazole magnesium)	A	1995	\$12,906,228	\$14,751,090
1940481	Paxil tab 20mg	Paroxetine (paroxetine hydrochloride)	N	1993	\$6,041,080	\$7,229,754
893757	Pravachol tab 20mg	Pravastatin sodium	C	1990	\$5,512,836	\$5,773,801
2220172	Apo-lovastatin - tab 20mg	Lovastatin	C	1997	\$2,350,159	\$4,803,546
884340	Zocor tab 20mg	Simvastatin	C	1994	\$3,805,329	\$4,588,608
884332	Zocor tab 10mg	Simvastatin	C	1990	\$4,447,526	\$4,504,775
708879	Vasotec tab 5mg	Enalapril maleate	C	1990	\$4,615,112	\$4,399,638
2230711	Lipitor 10mg	Atorvastatin (atorvastatin calcium)	C	1997	\$657,441	\$4,149,526
2155907	Adalat xl - srt 30mg	Nifedipine	C	1991	\$4,371,698	\$4,147,411
2215055	Becloforte inhaler - aem inh 250mcg/aem	Beclomethasone dipropionate	R	1990	\$4,543,302	\$3,968,025
878928	Norvasc tab 5mg	Amlodipine (amlodipine besylate)	C	1992	\$3,275,684	\$3,928,583
2229285	Zyprexa - 10mg	Olanzapine	N	1997	\$1,704,150	\$3,724,447
1962817	Zoloft cap 50mg	Sertraline (sertraline hydrochloride)	N	1991	\$3,092,524	\$3,466,465
894745	Clozaril tab 100mg	Clozapine	N	1990	\$2,838,467	\$3,165,731
670901	Vasotec tab 10mg	Enalapril maleate	C	1990	\$3,140,277	\$3,144,974
870935	Sinemet cr 200/50	Levodopa	N	1991	\$2,582,629	\$2,920,277
1947672	Accupril tab 10mg	Quinapril (quinapril hydrochloride)	C	1991	\$2,156,729	\$2,575,431
836338	Prepulsid tab 10mg	Cisapride (cisapride monohydrate)	A	1990	\$2,569,551	\$2,473,000
576158	Atrovent aem 28.6mg/100gm	Ipratropium bromide	R	1990	\$2,140,699	\$2,333,715
2155966	Cipro 500 - tab 500mg	Ciprofloxacin (ciprofloxacin hydrochloride)	J	1990	\$2,067,228	\$2,255,929
851752	Pulmicort turbuhaler 200mcg/aem	Budesonide	R	1990	\$2,053,493	\$2,121,170
2230713	Lipitor 20mg	Atorvastatin (atorvastatin calcium)	C	1997	\$314,846	\$2,075,559
2229269	Zyprexa - 5mg	Olanzapine	N	1997	\$1,023,128	\$2,013,612
2176017	Didrocal -400mg tab and 1250mg tab(500mg ca)	Calcium carbonate	M	1995	\$1,169,277	\$1,967,795
2177587	Pms-fluoxetine - cap 20mg	Fluoxetine (fluoxetine hydrochloride)	N	1995	\$2,098,300	\$1,889,840
TOTAL					\$81,477,694.98	\$98,372,702.49

Top 10 Category 1 Patented Drug Products British Columbia: 1998/99						
DIN	Brand	Ingredient	ATC	Year of Introduction	Expenditure 1997/98	Expenditure 1998/99
2190915	Losec 20mg	Omeprazole (omeprazole magnesium)	A	1995	\$12,906,228	\$14,751,090
884340	Zocor tab 20mg	Simvastatin	C	1994	\$3,805,329	\$4,588,608
870935	Sinemet cr 200/50	Levodopa	N	1991	\$2,582,629	\$2,920,277
851752	Pulmicort turbuhaler 200mcg/aem	Budesonide	R	1990	\$2,053,493	\$2,121,170
2176017	Didrocal -400mg tab and 1250mg tab(500mg ca)	Calcium carbonate	M	1995	\$1,169,277	\$1,967,795
2177587	Pms-fluoxetine - cap 20mg	Fluoxetine (fluoxetine hydrochloride)	N	1995	\$2,098,300	\$1,889,840
2213613	Flovent inhalers - aem inh-ori 250mcg/aem	Fluticasone propionate	R	1996	\$1,041,600	\$1,700,577
2146959	Lipidil micro - cap 200mg	Fenofibrate	C	1994	\$2,544,855	\$1,524,203
2054817	Prepulsid tab 20mg	Cisapride (cisapride monohydrate)	A	1993	\$1,286,116	\$1,509,038
851760	Pulmicort turbuhaler 400mcg/aem	Budesonide	R	1991	\$1,159,564	\$1,182,325
2155990	Adalat xl - srt 60mg	Nifedipine	C	1992	\$1,055,523	\$1,066,906

**Expenditures on the Top 10 Category 2 Patented Drug Products
British Columbia: 1998/99**

DIN	Brand	Ingredient	ATC	Year of Introduction	Expenditure 1997/98	Expenditure 1998/99
2155966	Cipro 500 - tab 500mg	Ciprofloxacin (ciprofloxacin hydrochloride)	J	1990	\$2,067,228	\$2,255,929
2025299	Risperdal tab 2mg	Risperidone	N	1992	\$1,463,821	\$1,627,549
2025302	Risperdal tab 3mg	Risperidone	N	1993	\$1,347,972	\$1,347,575
2212161	Imitrex - tab 100mg	Sumatriptan (sumatriptan succinate)	N	1991	\$1,302,690	\$1,290,268
2155958	Cipro 250 - tab 250mg	Ciprofloxacin (ciprofloxacin hydrochloride)	J	1990	\$1,208,489	\$1,258,837
1978926	Pulmicort nebuamp sus inh 0.5mg/ml	Budesonide	R	1992	\$655,440	\$709,922
2025310	Risperdal tab 4mg	Risperidone	N	1993	\$686,648	\$708,496
2169649	Betaseron	Interferon Beta - 1B	L	1997	\$312,523	\$671,139
2010909	Proscar tab 5mg	Finasteride	G	1992	\$1,583,492	\$644,335
2213575	Zofran - tab 8mg	Ondansetron (ondansetron hydrochloride dihydrate)	A	1991	\$522,052	\$602,849

**Expenditures on the Top 10 Category 3 Patented Drug Products
British Columbia: 1998/99**

DIN	Brand	Ingredient	ATC	Year of introduction	Expenditure 1997/98	Expenditure 1998/99
1940481	Paxil tab 20mg	Paroxetine (paroxetine hydrochloride)	N	1993	\$6,041,080	\$7,229,754
893757	Pravachol tab 20mg	Pravastatin sodium	C	1990	\$5,512,836	\$5,773,801
884332	Zocor tab 10mg	Simvastatin	C	1990	\$4,447,526	\$4,504,775
708879	Vasotec tab 5mg	Enalapril maleate	C	1990	\$4,615,112	\$4,399,638
2230711	Lipitor 10mg	Atorvastatin (atorvastatin calcium)	C	1997	\$657,441	\$4,149,526
878928	Norvasc tab 5mg	Amlodipine (amlodipine besylate)	C	1992	\$3,275,684	\$3,928,583
2229285	Zyprexa - 10mg	Olanzapine	N	1997	\$1,704,150	\$3,724,447
1962817	Zoloft cap 50mg	Sertraline (sertraline hydrochloride)	N	1991	\$3,092,524	\$3,466,465
670901	Vasotec tab 10mg	Enalapril maleate	C	1990	\$3,140,277	\$3,144,974
1947672	Accupril tab 10mg	Quinapril (quinapril hydrochloride)	C	1991	\$2,156,729	\$2,575,431

Appendix 4

Therapeutic Class Analysis

Percentage Contribution of Selected Therapeutic Classes to Total Expenditure British-Columbia: 1990/1991 - 1998/1999			
Therapeutic Class	Contribution in 1990/1991 (\$ millions)	Contribution in 1998/1999 (\$ millions)	% of Total Expenditure Change
Cardiovascular System	59.57	116.36	35.76%
Nervous System	22.94	75.3	32.97%
Alimentary Tract and Metabolism	24.46	42.63	11.44%
Respiratory System	13.04	24.59	7.27%
General Antiinfectives for Systemic Use	7.54	13.09	3.50%
Musculo-skeletal System	5.71	10.82	-3.07%
Genito Urinary System and Sex Hormones	3.9	8.14	2.67%
Blood and Blood Forming Organs	2.05	6.86	3.03%
Sensory Organs	4.34	5.93	1.00%
Dermatologicals	3.11	5.08	1.24%
Antineoplastic and Immunomodulating Agents	0.92	4.66	2.36%
Systemic Hormonal Preparations, excl. Sex Hormones	1.26	3.14	1.18%
Antiparasitic Products, Insecticides and Repellents	0.23	0.57	0.22%
Unclassified	0.57	1.15	0.36%
Various	0.19	0.31	0.08%
Total	159.82	318.63	100.00%

Anatomical Therapeutic Chemical (ATC)

The Anatomical Therapeutic Chemical (ATC) classification system [and the Defined Daily Dose (DDD)] as a measuring unit are recommended by the WHO for drug utilization studies.

In the ATC classification system, the drugs are divided into different groups according to the organ or system on which they act and their chemical, pharmacological and therapeutic properties. Drugs are classified in groups at five different levels. The drugs are divided into fourteen main groups (1st level), with two therapeutic/pharmacological subgroups (2nd and 3rd levels). The 4th level is a therapeutic/pharmacological/chemical subgroup and the 5th level is the chemical substance.

Medicinal products are classified according to the main therapeutic use of the main active ingredient, on the basic principle of only one ATC code for each pharmaceutical formulation (i.e. similar ingredients, strength and pharmaceutical form). A medicinal product can be given more than one ATC code if it is available in two or more strengths or formulations with clearly different therapeutic uses.

ATC	Therapeutic Class	Subgroups*
A02	Antacids, drugs for treatment of peptic ulcer and flatulence	Antacids; H ₂ -receptor antagonists; Prostaglandins; Proton pump inhibitors; Combinations for eradication of <i>Helicobacter pylori</i> & Others such as sucralfate
A10	Drugs used in diabetes	Insulins and analogues; Biguanides; Sulfonamides; Alpha glucosidase inhibitors; Thiazolidinediones & Others such as repaglinide
B01	Antithrombotic agents	Vitamin K antagonists (warfarin); Heparin group (includes LMWH); Platelet aggregation inhibitors (clopidogrel, ticlopidine..., abciximab...); Enzymes (streptokinase, alteplase..) & Others (lepirudin)
C01	Cardiac Therapy	Cardiac glycosides (digoxin); Antiarrhythmics; Cardiac stimulants (adrenergic and dopaminergic agents, phosphodiesterase inhibitors); Vasodilators (organic nitrates) & Others such prostaglandins
C07	Beta blocking agents	Beta blocking agents; Beta blocking agents and Thiazides; Beta blocking agents and other diuretics; Beta blocking agents and Vasodilators & Beta blocking agents and Other antihypertensives
C08	Calcium channel blockers	Selective Calcium channel blockers with mainly vascular effects; Selective Calcium channel blockers with direct cardiac effects; Non-selective Calcium channel blockers & Calcium channel blockers and diuretics
C09	Agents acting on the renin-angiotensin system	ACEIs, plain; ACEIs, combinations; Angiotensin II antagonists, plain; Angiotensin II antagonists, combinations & Others
C10	Serum lipid reducing agents	HMG CoA reductase inhibitors; Fibrates; Bile acid sequestrants; Nicotinic acid and derivatives
J01	Antibacterials for systemic use	Tetracyclines; Amphenicols (chloramphenicol); Penicillins; Beta-lactamase inhibitors; Cephalosporins; Monobactams; Carbapenems; Sulfonamides and Trimethoprim; Macrolides and Lincosamides (clindamycin); Aminoglycosides; Quinolones & Others such as vancomycin, fusidic acid, metronidazole

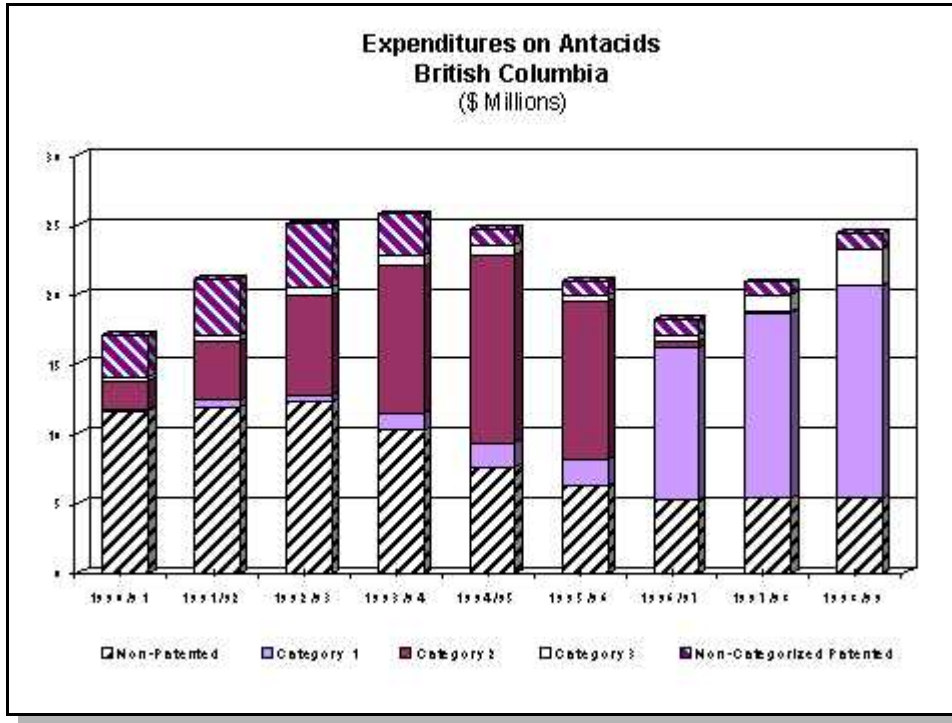
ATC	Therapeutic Class	Subgroups*
N02	Analgesics	Opioids (natural opium alkaloids such as morphine, codeine..; phenylpiperidines derivatives such as pethidine, fentanyl..; diphenylpropylamine derivatives such as methadone; pentazocine; morphinan derivative such as butorphanol and nalbuphine; opioids in combination with antispasmodics); Other analgesics and antipyretics (salicylic acid and derivatives, pyrazolones, anilides such as paracetamol); Antimigraine preparations (ergot alkaloids, selective 5HT ₁ -receptor agonists & other antimigraine preparations such as pizotifen, clonidine)
N03	Antiepileptics	Barbiturates and derivatives; Hydantoin derivatives; Oxazolidine derivatives; Succinimide derivatives; Benzodiazepine derivatives (clonazepam); Carboxamide derivatives; Fatty acid derivatives (valproic acid, vigabatrin) & Others (lamotrigine, topiramate, gabapentin)
N04	Anti-parkinson drugs	Anticholinergic agents; Dopaminergic agents [Dopa and dopa derivatives; Adamantane derivatives (amantadine); Dopamine agonists; MAO type B inhibitors (selegiline); Others (entacapone)]
N05	Psycholeptics	Antipsychotics (phenothiazines; butyrophenone derivatives; indole derivatives; thioxanthene derivatives; diphenylbutylpiperidine derivatives such as pimozide; diazepines, oxazepines and thiazepines such as clozapine, olanzepine & quetiapine; neuroleptics in tardive dyskinesia such as tetrabenazine; benzamides; lithium); Anxiolytics (benzodiazepine derivatives, carbamates, buspirone); Hypnotics and sedatives (barbiturates-plain, barbiturates-combinations, aldehydes and derivatives, benzodiazepine derivatives, piperidinedione derivatives, benzodiazepine related drugs such as zopiclone)
N06	Psychoanaleptics	Antidepressants; Psychostimulants and nootropics (centrally acting sympathomimetics, xanthine derivatives); Psycholeptics and psychoanaleptics in combination (antidepressants in combination with psycholeptics); Anti-dementia drugs
R03	Anti-asthmatics	Adrenergics, inhalants; Other anti-asthmatics, inhalants (glucocorticoids, anticholinergics, antiallergic agents); Adrenergics for systemic use; Other anti-asthmatics for systemic use (xanthines, xanthines and adrenergics, leukotriene receptor antagonists)

ATC	Therapeutic Class	Subgroups*
S01	Ophthalmologicals	Anti-infectives (antibiotics, sulfonamides, antivirals, other anti-infectives); Anti-inflammatory agents (corticosteroids, plain; corticosteroids and mydriatics in combination; anti-inflammatory agents, non-steroids); Anti-inflammatory agents and anti-infectives in combination; Anti-glaucoma preparations and miotics; Mydriatics and cycloplegics; Decongestants and antiallergics; Local anesthetics; Diagnostic agents; Surgical aids; Others such as artificial tears

* main one listed

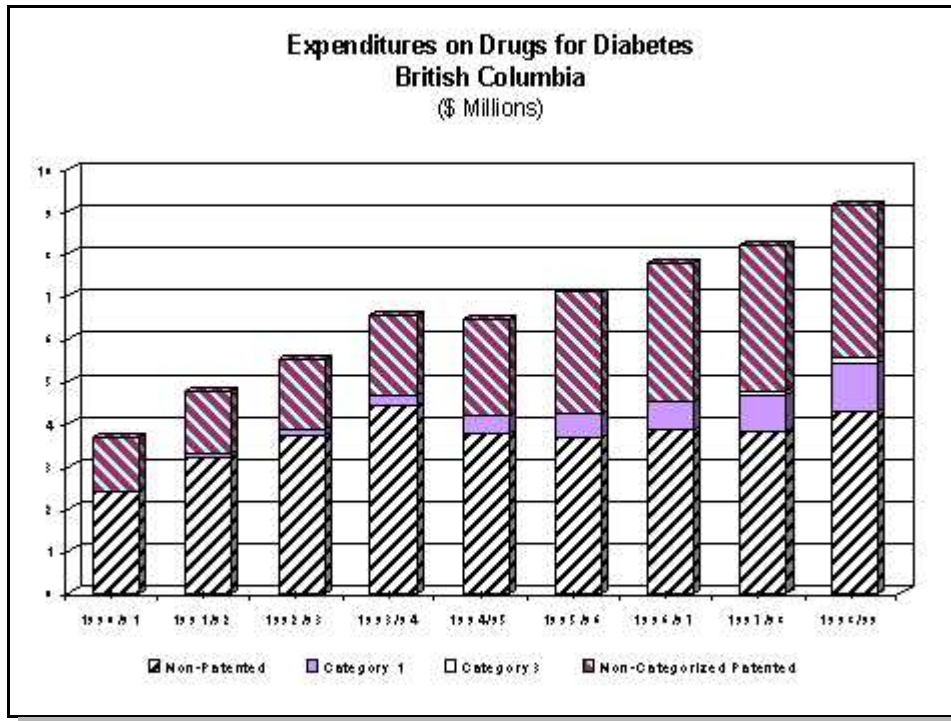
Impact of Existing and Newer Drug Products by Major Disease Groups
Antacids, Drugs for Treatment of Peptic Ulcer and Flatulence
British-Columbia: 1990/1991 - 1998/1999
(thousands of dollars)

Year of Introduction	CAT	90/91	91/92	92/93	93/94	94/95	95/96	96/97	97/98	98/99
1990/1991	-	11,618	11,981	12,031	9,399	6,339	6,240	5,437	4,872	4,303
1990/1991	1	211	373	405	381	576	277	89	104	93
1990/1991	2	1,920	4,327	7,189	10,636	13,515	11,382	396	121	41
1990/1991	3	289	383	591	782	718	463	217	198	167
1990/1991	NC	3,082	3,979	4,503	2,919	1,095	46	69	73	74
1991/1992	-	-	17	273	345	298	255	215	176	150
1992/1993	-	-	-	30	382	431	288	131	118	110
1992/1993	1	-	-	10	809	1,159	585	274	211	154
1993/1994	-	-	-	-	91	446	304	205	171	128
1994/1995	-	-	-	-	-	9	166	131	131	144
1995/1996	1	-	-	-	-	-	976	10,484	12,906	14,751
1995/1996	3	-	-	-	-	-	0	217	623	1,102
1995/1996	NC	-	-	-	-	-	0	-	-	0
1996/1997	-	-	-	-	-	-	-	193	745	925
1996/1997	3	-	-	-	-	-	-	87	160	158
1997/1998	-	-	-	-	-	-	-	-	121	656
1997/1998	1	-	-	-	-	-	-	-	12	55
1997/1998	3	-	-	-	-	-	-	-	144	1,196
1998/1999	-	-	-	-	-	-	-	-	-	14
1998/1999	1	-	-	-	-	-	-	-	-	95
Total Expenditures		17,121	21,060	25,030	25,745	24,637	20,932	18,144	20,885	24,315
Patented Expenditures		5,503	9,062	12,697	15,527	17,064	13,728	11,831	14,552	17,885
Non-Patented Expenditures		11,618	11,998	12,333	10,218	7,573	7,204	6,313	6,333	6,430

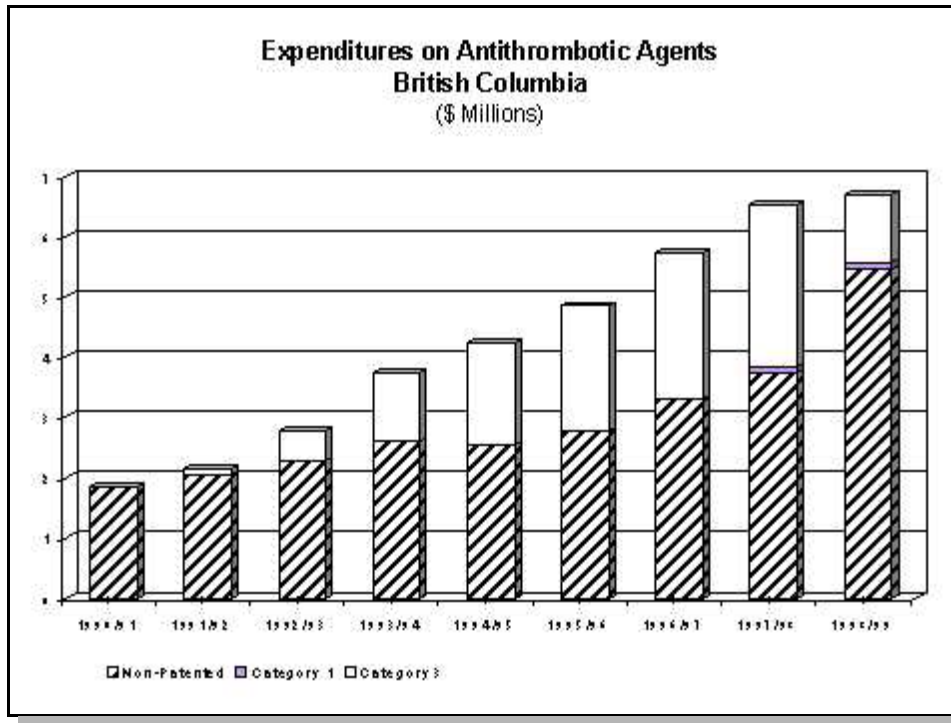


**Impact of Existing and Newer Drug Products by Major Disease Groups
Drug Used in Diabetes
British-Columbia: 1990/1991 - 1998/1999
(thousands of dollars)**

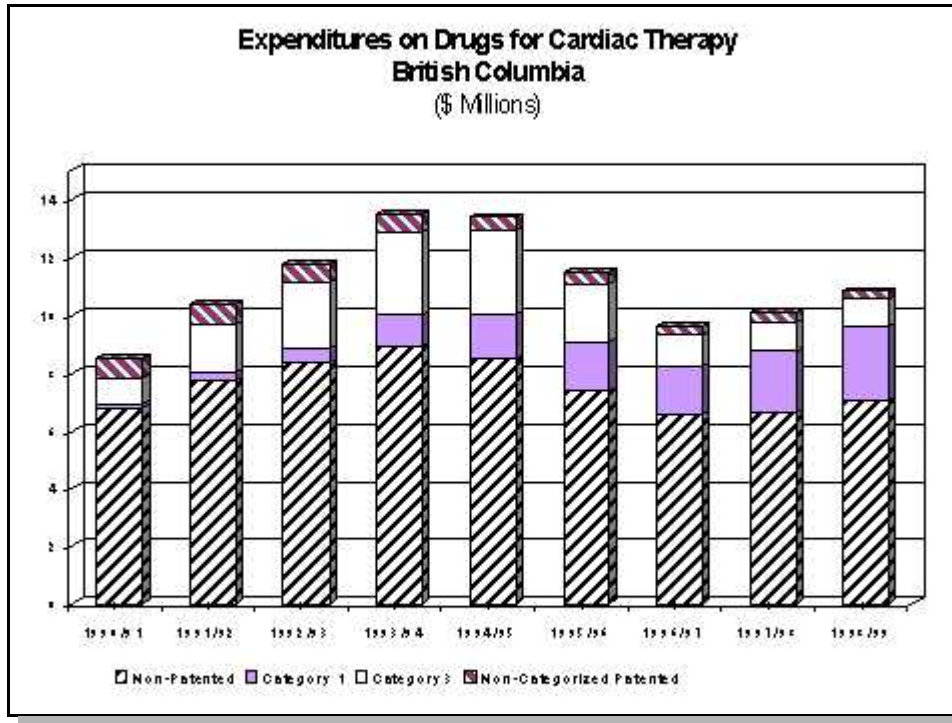
Year of Introduction	CAT	90/91	91/92	92/93	93/94	94/95	95/96	96/97	97/98	98/99
1990/1991	-	2,316	3,176	3,555	4,129	2,281	3,636	3,637	3,214	3,070
1990/1991	1	25	55	107	175	236	-	-	-	-
1990/1991	NC	1,251	1,475	1,672	1,799	1,853	514	519	488	464
1991/1992	-	-	42	149	153	273	328	414	476	495
1992/19923	-	-	-	3	142	244	383	372	397	509
1992/1993	1	-	-	13	46	96	-	-	-	-
1993/1994	-	-	-	-	1	972	1,159	828	834	860
1993/1994	1	-	-	-	16	63	76	129	191	251
1993/1994	NC	-	-	-	73	434	689	932	1,203	1,425
1994/1995	-	-	-	-	-	1	250	762	1,039	1,327
1995/1996	-	-	-	-	-	-	53	164	257	331
1995/1996	3	-	-	-	-	-	0	3	46	92
1996/1997	-	-	-	-	-	-	-	0	3	10
1996/1997	3	-	-	-	-	-	-	1	18	29
1997/1998	-	-	-	-	-	-	-	-	29	100
1998/1999	-	-	-	-	-	-	-	-	-	162
Total Expenditures		3,642	4,749	5,500	6,533	6,453	7,089	7,762	8,197	9,125
Patented Expenditures		1,276	1,531	1,792	2,108	2,681	1,279	1,584	1,946	2,262
Non-Patented Expenditures		2,366	3,218	3,708	4,424	3,772	5,809	6,177	6,251	6,863



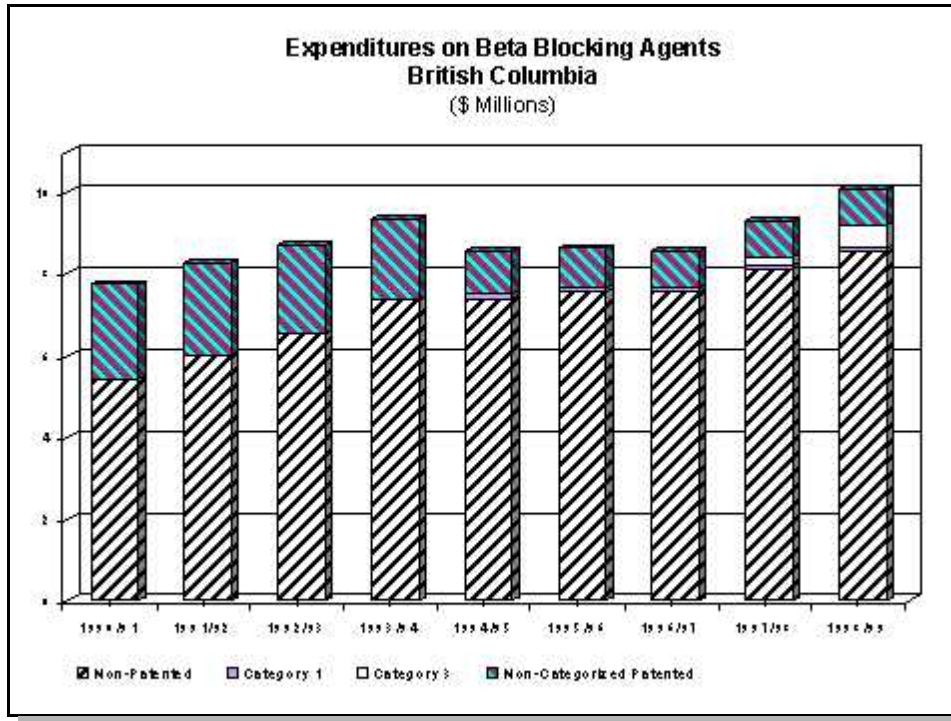
Impact of Existing and Newer Drug Products by Major Disease Groups Antithrombotic Agents British-Columbia: 1990/1991 - 1998/1999 (thousands of dollars)										
Year of Introduction	CAT	90/91	91/92	92/93	93/94	94/95	95/96	96/97	97/98	98/99
1990/1991	-	1,865	2,020	2,221	2,511	2,418	2,588	3,045	3,428	3,801
1991/1992	-	-	33	46	42	55	45	40	37	41
1991/1992	3	-	90	502	1,130	1,681	2,060	2,380	2,677	951
1992/1993	-	-	-	2	33	79	134	209	266	336
1994/1995	-	-	-	-	-	2	11	4	7	-
1995/1996	-	-	-	-	-	-	-	-	-	-
1995/1996	1	-	-	-	-	-	4	16	57	64
1996/1997	-	-	-	-	-	-	-	-	-	-
1996/1997	1	-	-	-	-	-	-	0	-	-
1996/1997	3	-	-	-	-	-	-	31	9	3
1997/1998	-	-	-	-	-	-	-	-	2	20
1997/1998	1	-	-	-	-	-	-	-	28	57
1997/1998	3	-	-	-	-	-	-	-	17	170
1998/1999	-	-	-	-	-	-	-	-	-	1,248
1998/1999	1	-	-	-	-	-	-	-	-	1
1998/1999	3	-	-	-	-	-	-	-	-	10
Total Expenditures		1,865	2,143	2,772	3,716	4,235	4,842	5,724	6,528	6,703
Patented Expenditures		0	90	502	1,130	1,681	2,064	2,427	2,789	1,257
Non-Patented Expenditures		1,865	2,053	2,269	2,586	2,554	2,778	3,297	3,739	5,445



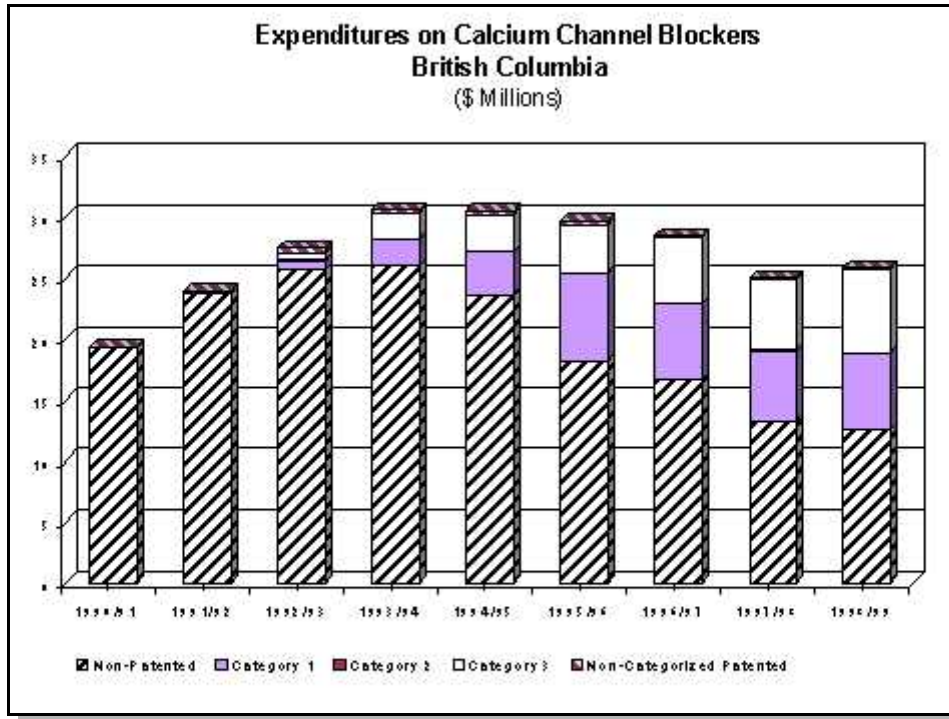
Impact of Existing and Newer Drug Products by Major Disease Groups Cardiac Therapy British-Columbia: 1990/1991 - 1998/1999 (thousands of dollars)										
Year of Introduction	CAT	90/91	91/92	92/93	93/94	94/95	95/96	96/97	97/98	98/99
1990/1991	-	6,889	8,446	8,816	9,574	8,942	7,759	6,838	6,862	7,141
1990/1991	1	1	81	431	705	981	1,025	962	1,160	1,395
1990/1991	3	901	1,686	2,297	2,889	2,893	2,061	1,093	1,043	965
1990/1991	NC	692	109	95	1	0	0	0	-	-
1991/1992	-	-	18	58	61	53	45	32	28	23
1991/1992	1	-	13	70	195	277	243	172	218	272
1991/1992	NC	0	-	0	0	-	-	-	-	-
1992/1993	-	-	-	3	16	25	34	37	42	41
1992/1993	1	-	-	1	3	5	7	11	19	28
1993/1994	1	-	-	-	12	193	255	233	275	372
1994/1995	-	-	-	-	-	1	6	4	0	0
1994/1995	1	-	-	-	-	14	51	74	99	55
1995/1996	-	-	-	-	-	0	4	13	16	17
1995/1996	1	-	-	-	-	-	10	168	282	329
1996/1997	-	-	-	-	-	-	-	0	0	-
1996/1997	1	-	-	-	-	-	-	6	38	58
1997/1998	-	-	-	-	-	-	-	-	7	42
1998/1999	-	-	-	-	-	-	-	-	-	86
Total Expenditures		8,484	10,353	11,772	13,456	13,385	11,500	9,643	10,089	10,823
Patented Expenditures		1,595	1,888	2,894	3,805	4,364	3,652	2,718	3,134	3,474
Non-Patented Expenditures		6,889	8,464	8,878	9,651	9,021	7,847	6,925	6,955	7,349



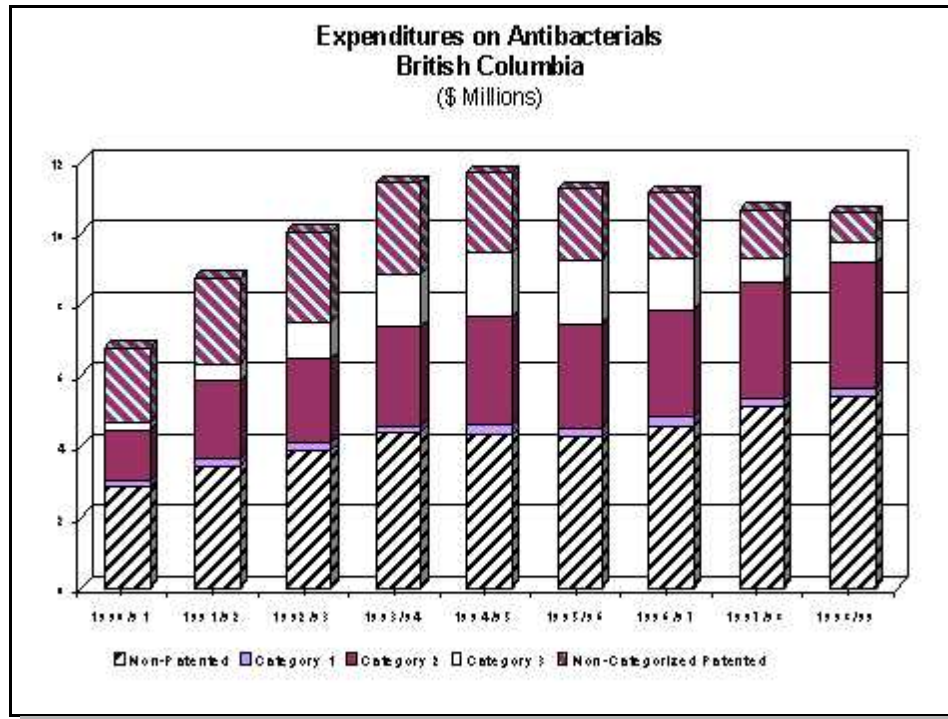
Impact of Existing and Newer Drug Products by Major Disease Groups Beta Blocking Agents British-Columbia: 1990/1991 - 1998/1999 (thousands of dollars)										
Year of Introduction	CAT	90/91	91/92	92/93	93/94	94/95	95/96	96/97	97/98	98/99
1990/1991	-	5,432	6,004	6,931	7,864	6,495	4,669	3,302	3,195	3,948
1990/1991	1	2	7	15	29	36	39	43	47	-
1990/1991	NC	2,328	2,251	1,745	1,060	873	832	797	783	170
1991/1992	-	-	0	1	1	6	7	9	9	13
1991/1992	1	-	1	-	-	-	-	-	-	-
1991/1992	NC	-	0	-	-	-	-	-	0	-
1992/1993	-	-	-	1	4	29	28	28	23	23
1993/1994	-	-	-	-	387	1,090	1,350	160	150	169
1994/1995	-	-	-	-	-	13	1,212	1,479	1,501	1,477
1995/1996	-	-	-	-	-	-	491	2,667	2,721	2,529
1996/1997	-	-	-	-	-	-	-	71	642	974
1997/1998	-	-	-	-	-	-	-	-	49	187
1997/1998	3	-	-	-	-	-	-	-	171	567
1998/1999	-	-	-	-	-	-	-	-	-	33
Total Expenditures		7,762	8,262	8,692	9,345	8,543	8,628	8,556	9,292	10,090
Patented Expenditures		2,329	2,258	1,760	1,089	910	871	839	1,001	737
Non-Patented Expenditures		5,432	6,004	6,932	8,256	7,633	7,757	7,717	8,290	9,353



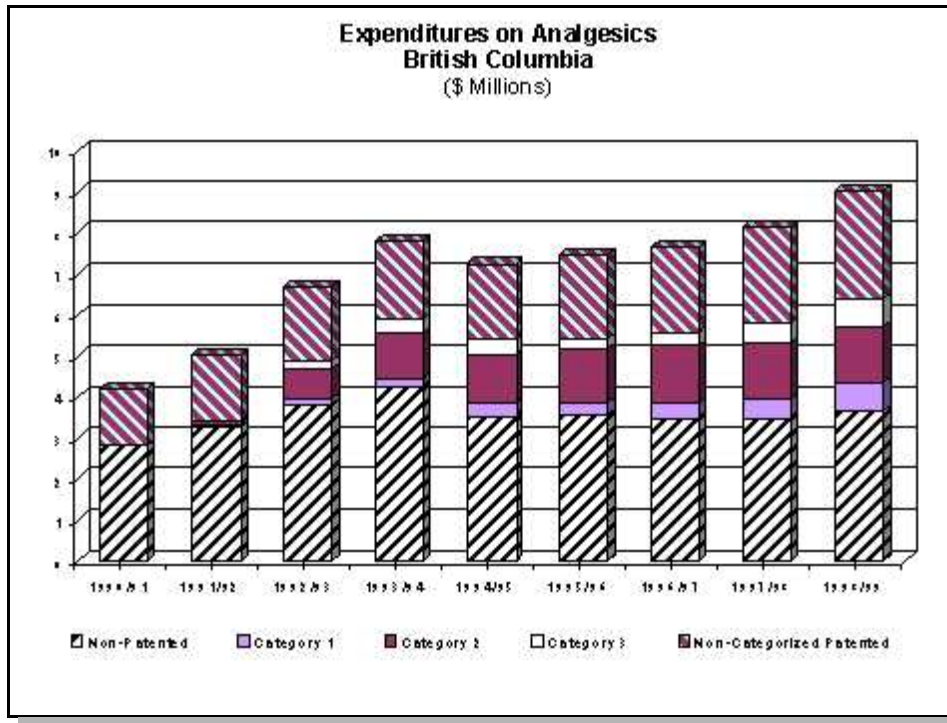
Impact of Existing and Newer Drug Products by Major Disease Groups Calcium Channel Blockers British-Columbia: 1990/1991 - 1998/1999 (thousands of dollars)										
Year of Introduction	CAT	90/91	91/92	92/93	93/94	94/95	95/96	96/97	97/98	98/99
1990/1991	-	19,287	23,673	25,419	24,285	19,448	4,436	2,853	993	760
1990/1991	2	0	1	7	8	8	9	22	37	41
1990/1991	NC	27	229	121	256	269	211	140	95	75
1991/1992	-	-	2	73	206	323	373	282	15	7
1991/1992	1	-	0	615	1,655	2,554	5,490	4,731	4,372	4,147
1991/1992	3	-	4	417	933	972	642	763	1,278	1,470
1991/1992	NC	-	19	281	164	53	24	14	6	5
1992/1993	-	-	-	146	1,469	3,123	9,885	7,484	131	84
1992/1993	1	-	-	157	548	914	1,692	1,521	1,056	1,067
1992/1993	3	-	-	161	1,099	2,056	3,317	4,697	4,526	5,432
1993/1994	-	-	-	-	38	526	2,801	2,411	697	724
1993/1994	1	-	-	-	0	5	14	33	100	148
1994/1995	-	-	-	-	-	180	663	765	1,048	1,125
1994/1995	1	-	-	-	-	12	24	10	14	11
1995/1996	-	-	-	-	-	-	42	740	1,576	1,599
1996/1997	-	-	-	-	-	-	-	2,051	7,353	5,239
1997/1998	-	-	-	-	-	-	-	-	1,634	3,715
1998/1999	-	-	-	-	-	-	-	-	-	13
1998/1999	1	-	-	-	-	-	-	-	-	159
Total Expenditures		19,315	23,928	27,395	30,663	30,444	29,628	28,515	24,932	25,821
Patented Expenditures		28	252	1,758	4,664	6,843	11,424	11,930	11,484	12,555
Non-Patented Expenditures		19,287	23,675	25,637	25,999	23,601	18,205	16,585	13,447	13,266



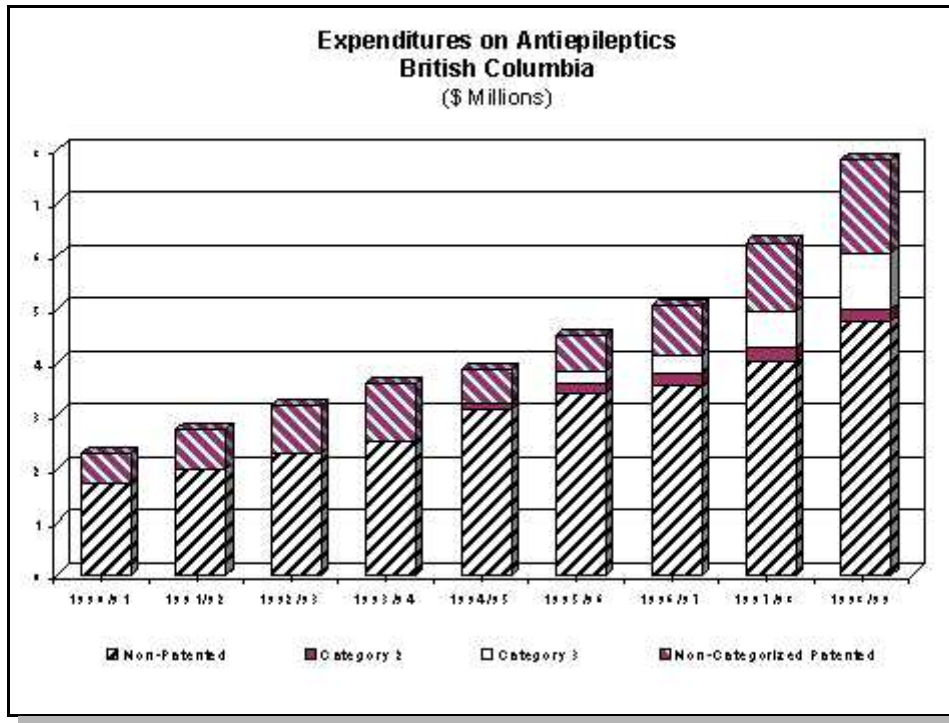
Impact of Existing and Newer Drug Products by Major Disease Groups										
Antibacterials for Systemic Use										
British-Columbia: 1990/1991 - 1998/1999										
(thousands of dollars)										
Year of	CAT	90/91	91/92	92/93	93/94	94/95	95/96	96/97	97/98	98/99
1990/1991	-	2,916	3,499	3,933	4,566	4,248	3,951	4,102	4,290	4,237
1990/1991	1	181	217	201	179	91	40	33	26	10
1990/1991	2	1,407	2,175	2,395	2,801	3,044	2,867	2,991	3,276	3,515
1990/1991	3	201	337	498	610	627	566	438	287	246
1990/1991	NC	2,053	2,380	2,387	2,320	2,099	1,837	1,704	1,264	796
1991/1992	-	-	0	6	11	10	9	8	5	8
1991/1992	1	-	0	-	-	-	-	0	0	0
1991/1992	3	-	114	214	207	175	159	153	137	44
1991/1992	NC	-	0	-	0	-	0	-	-	-
1992/1993	-	-	-	23	15	127	191	248	282	300
1992/1993	1	-	-	3	1	1	1	0	0	0
1992/1993	3	-	-	305	663	981	1,069	785	161	131
1992/1993	NC	-	-	0	-	-	-	-	-	-
1993/1994	-	-	-	-	7	215	165	135	142	142
1993/1994	1	-	-	-	0	0	-	-	-	-
1994/1995	-	-	-	-	-	38	215	239	250	265
1994/1995	1	-	-	-	-	19	110	147	93	100
1994/1995	3	-	-	-	-	0	-	-	7	0
1994/1995	NC	-	-	-	-	0	3	-	-	-
1995/1996	-	-	-	-	-	-	8	22	35	60
1995/1996	1	-	-	-	-	-	0	-	-	0
1995/1996	3	-	-	-	-	-	46	49	5	4
1995/1996	NC	-	-	-	-	-	3	2	8	-
1996/1997	-	-	-	-	-	-	-	32	263	304
1996/1997	3	-	-	-	-	0	-	2	21	33
1997/1998	-	-	-	-	-	-	-	-	5	87
1997/1998	3	-	-	-	-	-	-	-	54	85
1998/1999	-	-	-	-	-	-	-	-	-	186
1998/1999	1	-	-	-	-	-	-	-	-	2
1998/1999	3	-	-	-	-	-	-	-	-	4
Total Expenditures		6,757	8,724	10,025	11,381	11,676	11,243	11,089	10,611	10,558
Patented Expenditures		3,841	5,224	6,003	6,782	7,038	6,702	6,304	5,339	4,969
Non-Patented Expenditures		2,916	3,499	4,022	4,599	4,638	4,540	4,785	5,272	5,589



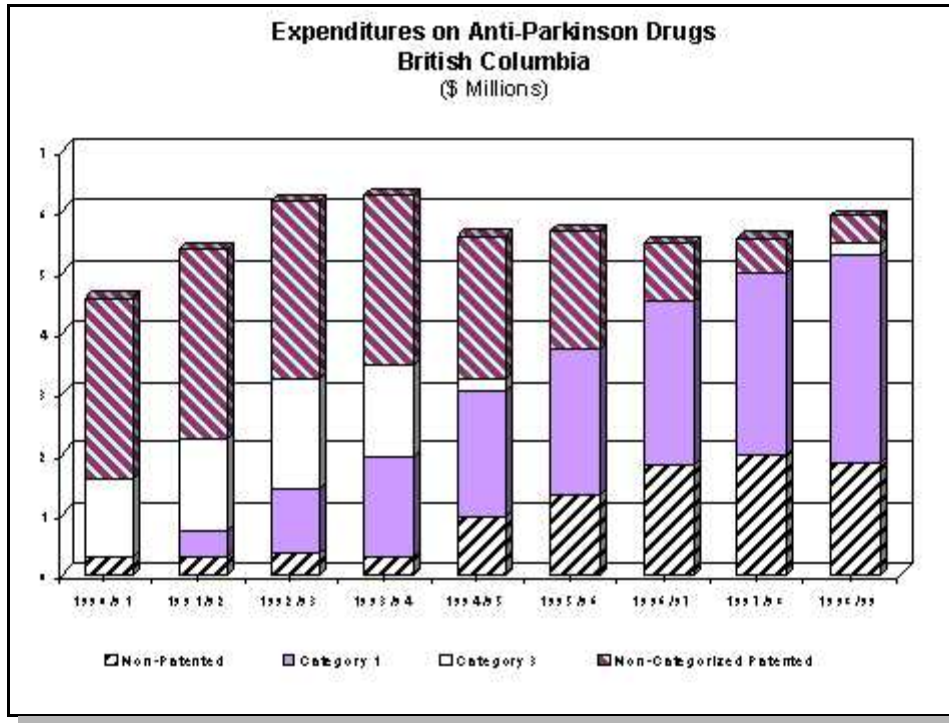
Impact of Existing and Newer Drug Products by Major Disease Groups Analgesics British-Columbia: 1990/1991 - 1998/1999 (thousands of dollars)										
Year of Introduction	CAT	90/91	91/92	92/93	93/94	94/95	95/96	96/97	97/98	98/99
1990/1991	-	2,791	3,238	3,491	3,778	4,939	5,056	5,162	5,214	5,565
1990/1991	1	0	95	147	198	-	-	-	-	-
1990/1991	NC	1,413	1,605	1,813	1,909	8	2	2	1	1
1991/1992	-	-	26	265	415	446	400	349	319	370
1991/1992	2	-	66	756	1,075	1,165	1,301	1,375	1,360	1,345
1992/1993	-	-	0	27	61	85	76	60	48	39
1992/1993	3	-	-	178	336	405	250	315	378	535
1993/1994	-	-	-	-	13	145	238	216	238	310
1994/1995	-	-	-	-	0	48	94	76	84	87
1994/1995	3	-	-	-	-	0	6	34	65	79
1995/1996	-	-	-	-	-	-	31	34	105	72
1995/1996	1	-	-	-	-	-	4	27	37	55
1996/1997	-	-	-	-	-	-	-	11	26	42
1996/1997	1	-	-	-	-	-	-	10	103	196
1996/1997	NC	-	-	-	-	-	-	0	76	92
1997/1998	-	-	-	-	-	-	-	-	37	3
1997/1998	1	-	-	-	-	-	-	-	3	97
1997/1998	3	-	-	-	-	-	-	-	47	87
1998/1999	-	-	-	-	-	-	-	-	-	47
1998/1999	1	-	-	-	-	-	-	-	-	15
Total Expenditures		4,204	5,030	6,676	7,784	7,241	7,457	7,671	8,142	9,037
Patented Expenditures		1,414	1,766	2,894	3,518	1,578	1,562	1,763	2,071	2,502
Non-Patented Expenditures		2,791	3,264	3,782	4,266	5,663	5,894	5,908	6,071	6,535



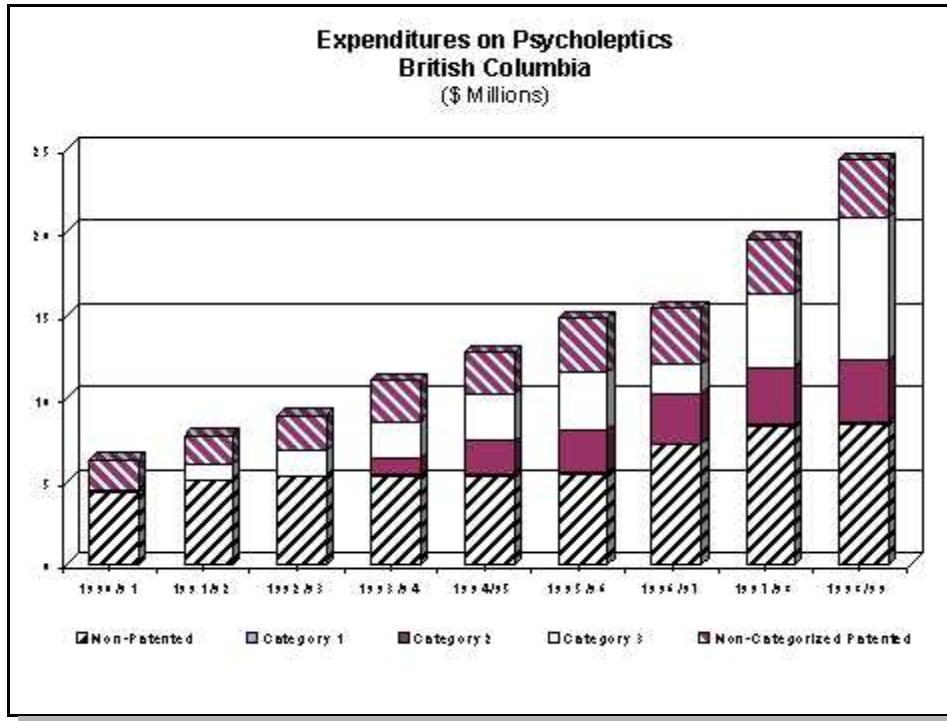
Impact of Existing and Newer Drug Products by Major Disease Groups										
Antiepileptics										
British-Columbia: 1990/1991 - 1998/1999										
(thousands of dollars)										
Year of Introduction	CAT	90/91	91/92	92/93	93/94	94/95	95/96	96/97	97/98	98/99
1990/1991	-	2,087	2,459	2,821	3,166	2,674	2,519	2,312	2,079	1,743
1990/1991	NC	201	282	357	431	501	668	911	1,293	1,745
1991/1992	-	-	0	0	-	-	0	0	0	0
1992/1993	-	-	-	0	0	0	0	0	0	0
1993/1994	-	-	-	-	16	509	515	385	365	338
1994/1995	-	-	-	-	-	45	343	398	917	1,590
1994/1995	2	-	-	-	-	154	215	231	-	-
1994/1995	3	-	-	-	-	2	190	308	470	580
1995/1996	-	-	-	-	-	-	52	427	603	622
1995/1996	3	-	-	-	-	-	9	24	45	58
1996/1997	-	-	-	-	-	-	-	60	283	302
1997/1998	-	-	-	-	-	-	-	-	27	215
1997/1998	3	-	-	-	-	-	-	-	172	444
1998/1999	-	-	-	-	-	-	-	-	-	178
Total Expenditures		2,288	2,741	3,178	3,614	3,885	4,511	5,057	6,255	7,814
Patented Expenditures		201	282	357	431	656	1,082	1,474	1,980	2,827
Non-Patented Expenditures		2,087	2,459	2,821	3,183	3,229	3,430	3,583	4,274	4,987



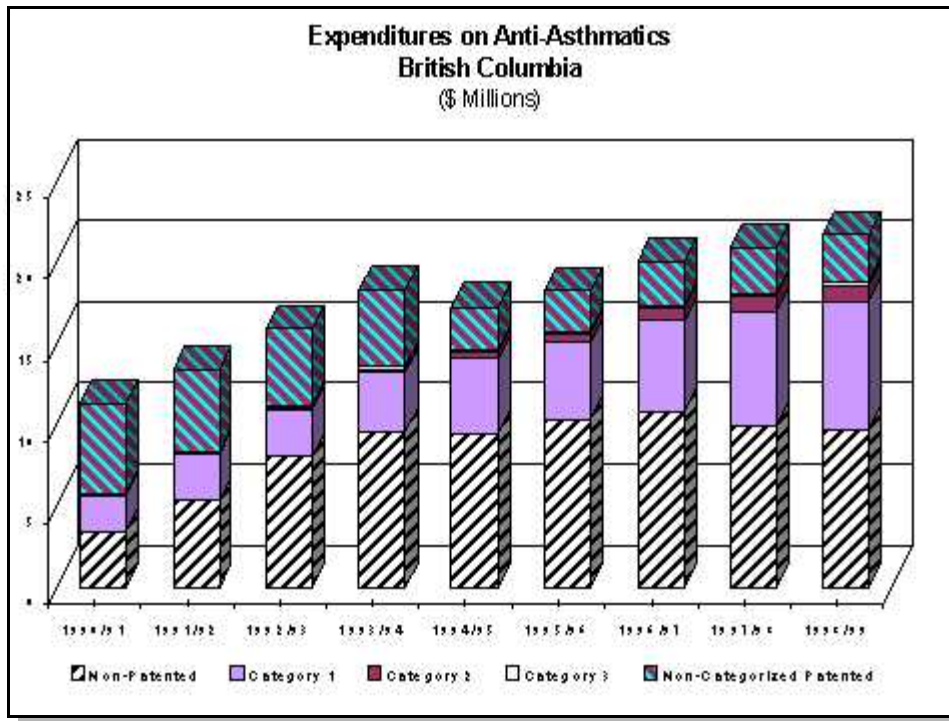
Impact of Existing and Newer Drug Products by Major Disease Groups										
Anti-Parkinson Drugs										
British-Columbia: 1990/1991 - 1998/1999										
(thousands of dollars)										
Year of Introduction	CAT	90/91	91/92	92/93	93/94	94/95	95/96	96/97	97/98	98/99
1990/1991	-	397	1,619	3,217	2,936	2,282	1,879	870	394	215
1990/1991	3	1,274	1,529	1,796	1,535	222	2	2	0	1
1990/1991	NC	2,894	1,823	-	-	-	-	-	-	-
1991/1992	-	-	0	0	0	-	-	-	-	-
1991/1992	1	-	414	1,083	1,615	1,928	2,146	2,350	2,583	2,920
1991/1992	NC	-	5	50	96	147	164	66	47	21
1992/1993	-	-	-	0	-	0	0	0	-	0
1993/1994	-	-	-	-	52	105	96	79	67	51
1993/1994	1	-	-	-	34	145	241	342	427	504
1994/1995	-	-	-	-	-	773	1,120	969	829	659
1995/1996	-	-	-	-	-	-	19	380	482	454
1995/1996	NC	-	-	-	-	-	13	152	225	291
1996/1997	-	-	-	-	-	-	-	278	480	542
1997/1998	-	-	-	-	-	-	-	-	10	39
1998/1999	-	-	-	-	-	-	-	-	-	-
1998/1999	3	-	-	-	-	-	-	-	-	209
Total Expenditures		4,564	5,391	6,146	6,268	5,603	5,680	5,487	5,545	5,908
Patented Expenditures		4,167	3,772	2,929	3,279	2,442	2,566	2,911	3,283	3,947
Non-Patented Expenditures		397	1,619	3,217	2,989	3,160	3,114	2,577	2,262	1,961



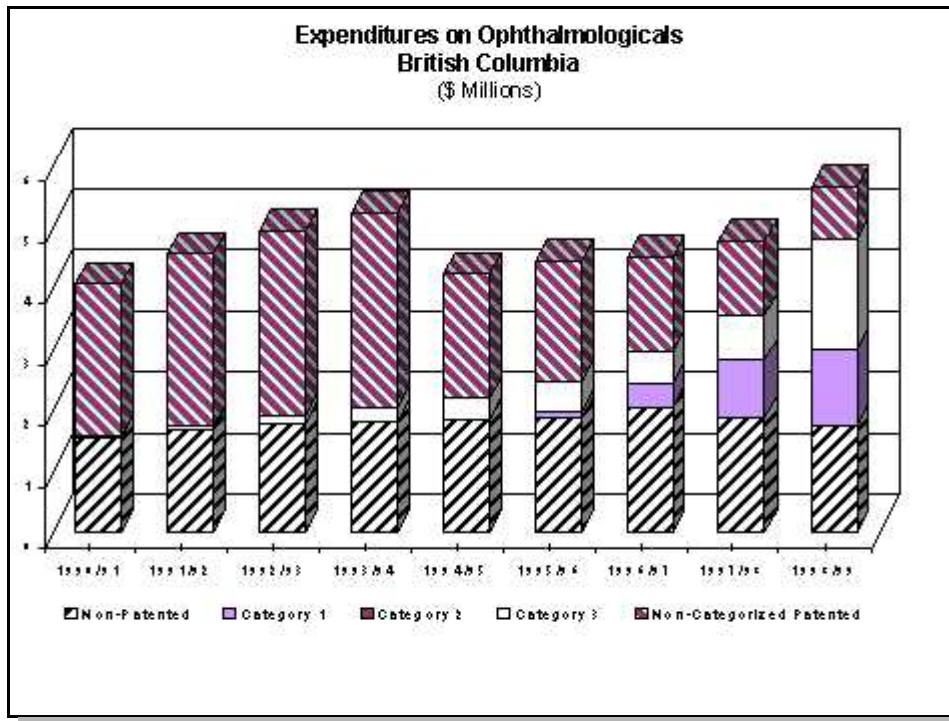
Impact of Existing and Newer Drug Products by Major Disease Groups										
Psycholeptics										
British-Columbia: 1990/1991 - 1998/1999										
(thousands of dollars)										
Year of Introduction	CAT	90/91	91/92	92/93	93/94	94/95	95/96	96/97	97/98	98/99
1990/1991	-	4,462	5,533	7,806	9,851	10,058	10,872	8,236	6,551	5,869
1990/1991	1	175	123	83	-	-	-	-	-	-
1990/1991	3	163	851	-	-	-	-	-	-	-
1990/1991	NC	1,608	1,271	1,048	54	61	63	59	65	65
1991/1992	-	-	5	30	38	60	65	79	103	123
1992/1993	-	-	-	3	46	93	105	101	92	26
1992/1993	2	-	-	0	373	723	986	1,245	1,464	1,628
1992/1993	3	-	-	0	2	1	1	3	8	0
1992/1993	NC	-	-	0	-	-	-	-	-	-
1993/1994	-	-	-	-	2	150	106	224	238	143
1993/1994	2	-	-	-	585	1,182	1,514	1,744	2,035	2,056
1993/1994	3	-	-	-	93	214	435	758	1,180	1,692
1994/1995	-	-	-	-	-	219	457	471	494	439
1995/1996	-	-	-	-	-	-	57	247	298	278
1996/1997	-	-	-	-	-	-	-	2,312	3,623	3,782
1997/1998	-	-	-	-	-	-	-	-	261	821
1997/1998	1	-	-	-	-	-	-	-	10	180
1997/1998	3	-	-	-	-	-	-	-	3,169	6,780
1998/1999	-	-	-	-	-	-	-	-	-	408
1998/1999	3	-	-	-	-	-	-	-	-	56
Total Expenditures		6,308	7,783	8,971	11,044	12,760	14,760	15,478	19,590	24,347
Patented Expenditures		1,846	2,245	1,132	1,107	2,180	2,998	3,810	7,930	12,457
Non-Patented Expenditures		4,462	5,538	7,839	9,937	10,580	11,761	11,668	11,660	11,890



Impact of Existing and Newer Drug Products by Major Disease Groups Anti-asthmatics British-Columbia: 1990/1991 - 1998/1999 (thousands of dollars)										
Year of Introduction	CAT	90/91	91/92	92/93	93/94	94/95	95/96	96/97	97/98	98/99
1990/1991	-	4,215	8,001	8,447	7,981	4,968	4,500	4,557	4,736	4,743
1990/1991	1	2,252	2,569	2,281	2,430	2,468	2,466	2,519	2,573	2,616
1990/1991	3	28	145	156	163	185	170	141	107	82
1990/1991	NC	4,766	1,718	1,310	1,037	265	226	195	121	99
1991/1992	-	-	930	3,284	5,030	5,826	6,747	6,497	4,722	3,995
1991/1992	1	-	11	246	521	737	911	1,083	1,160	1,182
1992/1993	-	-	-	18	717	1,718	1,778	1,861	1,725	1,444
1992/1993	1	-	-	55	-	-	-	-	-	-
1992/1993	2	-	-	56	238	408	560	804	980	1,064
1993/1994	-	-	-	-	105	316	423	498	385	42
1993/1994	NC	-	-	-	49	223	212	178	155	127
1994/1995	-	-	-	-	-	67	226	243	217	168
1995/1996	-	-	-	-	-	-	3	110	305	309
1995/1996	1	-	-	-	-	-	2	382	939	1,088
1995/1996	3	-	-	-	-	-	0	40	58	18
1996/1997	-	-	-	-	-	-	-	517	1,486	1,760
1996/1997	1	-	-	-	-	-	-	337	1,081	1,760
1997/1998	-	-	-	-	-	-	-	-	89	899
1997/1998	1	-	-	-	-	-	-	-	2	8
1997/1998	3	-	-	-	-	-	-	-	8	76
1998/1999	-	-	-	-	-	-	-	-	-	63
1998/1999	1	-	-	-	-	-	-	-	-	128
Total Expenditures		11,261	13,374	15,854	18,273	17,181	18,223	19,963	20,849	21,672
Patented Expenditures		7,046	4,443	4,104	4,440	4,286	4,547	5,679	7,184	8,248
Non-Patented Expenditures		4,215	8,931	11,750	13,833	12,895	13,676	14,284	13,665	13,424



Impact of Existing and Newer Drug Products by Major Disease Groups Ophthalmologicals British-Columbia: 1990/1991 - 1998/1999 (thousands of dollars)										
Year of Introduction	CAT	90/91	91/92	92/93	93/94	94/95	95/96	96/97	97/98	98/99
1990/1991	-	1,960	2,243	2,448	3,892	2,901	2,579	1,626	1,600	1,304
1990/1991	1	0	0	0	-	-	-	-	-	-
1990/1991	3	-	-	-	-	-	-	186	214	229
1990/1991	NC	2,100	2,296	2,400	960	541	547	521	25	1
1991/1992	-	-	3	18	24	48	48	51	49	41
1991/1992	3	-	5	39	55	78	184	173	167	211
1992/1993	-	-	-	8	179	425	662	841	777	674
1992/1993	2	-	-	0	0	2	2	-	-	-
1992/1993	3	-	-	7	78	144	100	121	158	229
1993/1994	-	-	-	-	7	20	17	22	26	22
1993/1994	NC	-	-	-	30	57	77	101	124	14
1994/1995	-	-	-	-	-	11	104	121	96	68
1994/1995	1	-	-	-	-	8	89	171	613	854
1995/1996	-	-	-	-	-	-	12	326	305	290
1995/1996	1	-	-	-	-	-	20	212	317	380
1996/1997	-	-	-	-	-	-	-	25	99	110
1997/1998	-	-	-	-	-	-	-	-	15	41
1997/1998	3	-	-	-	-	-	-	-	171	1,153
1998/1999	-	-	-	-	-	-	-	-	-	26
Total Expenditures		4,061	4,547	4,921	5,225	4,235	4,441	4,496	4,757	5,647
Patented Expenditures		2,101	2,301	2,446	1,123	830	1,018	1,485	1,789	3,071
Non-Patented Expenditures		1,960	2,246	2,475	4,102	3,405	3,422	3,011	2,968	2,576



Appendix 5

Glossary

Beneficiary

Someone who has made a claim to the British Columbia Pharmacare Program.

Category 1 Drugs

PMPRB din categorization - a new DIN of an existing or comparable dosage form of an existing medicines, usually a new strength of an existing drug (line extension).

Category 2 Drugs

PMPRB DIN categorization - the first drug product to treat effectively a particular illness or which provides a substantial improvement over existing drug products, often referred to as "breakthrough" or "substantial improvement".

Category 3 Drugs

PMPRB DIN categorization - a new drug or new dosage form of an existing medicine that provides moderate, little or no improvement over existing medicines.

Exiting Drug Effect

Exiting Drug Effect shows the amount by which expenditures decrease as a result of de-listing drugs from the formulary, discontinuation of the products by the manufacturer, or lack of claims during follow-up periods.

Existing Drug Products

In this Study, Existing Drug Products are defined as drug products that were already listed on the British Columbia benefit list on or before 1990/91.

New Drug Effect

New Drug Effect shows the amount by which expenditures increase as a result of listing new drugs on the formulary.

Newer Drug Products

In this Study, new drug products are defined as drug products that were listed on the British Columbia Pharmacare benefit list in 1991/92 or during subsequent years.

Price Effect

Price effect shows the impact of prices on expenditures by holding volume consumed constant. In other words, it is the amount by which expenditures would change if volume consumed did not change from the previous year.

Total Pharmaceutical Expenditures

Total Pharmaceutical Expenditures in this study include expenditures made by the British Columbia Pharmacare Program and any deductibles and co-payments made by its beneficiaries. Expenditures also include wholesale mark ups but do not include dispensing fees.

Volume Effect

Volume effect shows the impact of volume consumed on expenditures by holding prices constant. In other words, it is the amount by which expenditures would change if prices did not change from the previous year.