

USE OF PHYSICIAN PROFILES TO INFLUENCE PRESCRIBING OF TOPICAL CORTICOSTEROIDS

Ingrid S Sketris¹, George Kephart², Charmaine A Cooke², Chris D Skedgel³, Pam R McLean-Veysey⁴ on behalf of the Drug Evaluation Alliance of Nova Scotia (DEANS)

¹College of Pharmacy, Dalhousie University, ²Department of Community Health and Epidemiology, Dalhousie University, ³Department of Medicine, Dalhousie University, ⁴Capital District Health

Corresponding Author: ingrid.sketris@dal.ca

ABSTRACT

Background

Physician profiling is a tool used to attempt to affect changes in prescribing. The Drug Evaluation Alliance of Nova Scotia (DEANS) decided to implement a physician profiling project to determine if prescribing of topical corticosteroids could be altered.

Objectives

To evaluate a DEANS initiative utilizing physician prescribing profiles to shift prescribing of topical corticosteroids from higher to lower potency agents in beneficiaries of the Nova Scotia Seniors' Pharmacare Program.

Methods

Administrative claims from the Nova Scotia Seniors' Pharmacare program were used to identify prescriptions for topical corticosteroids. Prescriptions were summarized at the individual physician level, and aggregated by Anatomical Therapeutic Classification into weak, moderately potent, potent and very potent products. The number of prescriptions for topical corticosteroids was compared for the twelve-month period before and after mailing of the profiles. Overall results were aggregated by utilization and expenditures.

Results

The number of prescriptions for topical corticosteroids per physician profiled was 44.0 in 2000/2001 and 42.8 in 2001/2002 ($p=NS$) and the expenditures per physician profiled were \$838.94 in 2000/2001 and \$826.81 in 2001/2002 ($p=NS$). There was a small decrease in prescriptions dispensed for potent topical products over the profiling period (52.4% of prescriptions in 2000/2001 versus 51.5% of prescriptions in 2001/2002, $p=0.03$). Otherwise, changes in utilization or expenditures for topical corticosteroids were not statistically different between the profiling periods.

Conclusions

This project showed that mailing unsolicited individual-level profiles did not alter prescribing or expenditures for topical corticosteroids over a two-year period. Further work is needed to determine physician attitudes towards such projects.

Key Words: *Topical corticosteroids; physician prescribing profiles; drug utilization*

Gaps between evidence and clinical practice, along with variations in health care utilization affect quality of care and health expenditures.¹⁻⁸ Inappropriate medication prescribing has been identified as one of the five most important quality of care issues in terms of preventable morbidity and mortality, especially in the elderly.^{9,10}

One tool used to promote change in physician behaviour is physician profiling. Physician profiling is the process of providing a summary of a physician's past patient care activities in order to influence future clinical and administrative decisions.^{9, 11-14} Information can be obtained from clinical or administrative databases, or from surveys. Performance is typically compared to peers or to clinically 'acceptable' levels.^{5, 14-16}

Physician profiling has been used as a quality assessment tool to provide feedback on patient care.^{13,15,17-19} It assumes that notifying individuals or groups about deviations from peer behaviour or accepted clinical criteria will lead to improved physician performance.^{9,15,16,20} While physician profiling has been widely used to assess physician performance, evidence regarding whether it is effective in changing behaviour is conflicting.^{1,2,8,12,13, 21-23} Overviews of systematic reviews on the effectiveness of methods to change physician behaviour indicate that profiling sometimes had a significant effect on utilization, but that the clinical importance of the changes was moderate and that the generalizability of the studies was uncertain.^{1,12,23} These overviews also note that studies examining profiling often had methodological weaknesses and differed in design, content, data presentation, barriers to change, and study setting.

Topical Corticosteroid Profiling Initiative

Many topical corticosteroids are available on the market, varying by chemical entity, potency, formulation, combination, and price.²⁴⁻²⁶

The variety of products available makes it difficult for physicians to choose the most appropriate and cost-effective product for their patients. Topical corticosteroid products are classified by the World Health Organization (WHO) Anatomical Therapeutic Classification (ATC) system²⁷ as either weak, moderately potent, potent or very potent. For products within the

same potency category, other than differences in formulation and cost, there is little evidence for any therapeutic advantage of one product over another.^{28,29} Potent and very potent products are usually reserved for severe unresponsive skin conditions, are generally limited to 10-14 days of therapy due to the possibility of a higher incidence of adverse effects, such as tachyphylaxis, atrophy and thinning of the skin.^{26,28,30-32} Low to moderate potency corticosteroids are usually preferred for mild-moderate inflammatory conditions, for long-term use, or for use in patients with thin or sensitive skin, including the elderly.^{28,31,33}

One of the structures used to encourage appropriate drug use in Nova Scotia is the Drug Evaluation Alliance of Nova Scotia (DEANS), a program funded by the Nova Scotia Department of Health.³⁴ DEANS is comprised of a multi-disciplinary team, with expertise in family medicine, pharmacy, continuing pharmacy medical education, drug evaluation and epidemiology, with a mission to contribute to the health of Nova Scotians by encouraging appropriate and cost-effective drug use. The role of DEANS is to identify critical drug care issues; obtain and analyze information and data relevant to these issues; develop interventions to provide targeted, evidence-based information; and to evaluate the impact of initiatives on provider and consumer behaviours.

In order to determine the feasibility and impact of employing profiles to provide physicians with feedback on their prescribing practices for topical corticosteroids, DEANS initiated a physician profiling project in the Winter of 2001. The objective of the project was to determine if physician profiling could shift prescribing of topical corticosteroids from higher cost, higher potency products to lower cost, lower potency products. Topical corticosteroids were chosen for a number of reasons. The Nova Scotia Formulary Management Committee had completed a review of these products in 1999, had de-listed all but two topical combination corticosteroid products, and was currently considering adding several new products to the Nova Scotia Pharmacare Programs' benefit list. Although this delisting had decreased the number of agents available as insured benefits, there were still over 150 topical corticosteroid products listed as a benefit on the Nova Scotia Formulary. A drug

use evaluation²⁵ demonstrated that even after delisting specific topical corticosteroid combination products, approximately 50% of prescriptions dispensed to seniors were for potent topical corticosteroids, and drug expenditures for topical corticosteroids to the Senior Pharmacare Program were approximately \$900,000 for the fiscal year April 1, 1999 to March 31, 2000 (including pharmacists professional fees, but after deducting patient co-payments). It was also felt that physicians might appreciate a tool to help in choosing the most appropriate product for their patients.

METHODS

Study Setting

Nova Scotia is a province in Canada with approximately 942,000 inhabitants. The Nova Scotia government funds two main drug insurance programs: one for seniors (The Nova Scotia Seniors' Pharmacare Program, with approximately 95,000 beneficiaries) and one for persons receiving social assistance through the Department of Community Services (with approximately 65,000 beneficiaries). The seniors' program provides drug benefits to residents aged 65 years and over, who have opted to participate by payment of the required insurance premium and co-payments. It does not include seniors who have drug insurance from Federal Programs (the Royal Canadian Mounted Police, Veterans Affairs Canada, eligible First Nations or Inuit, Correctional Services of Canada, Retired Federal Employees) or those who use solely private drug insurance. This study was approved by the Health Sciences Human Research Ethics Board, Dalhousie University, Halifax, Nova Scotia.

Data for Profiles

Administrative claims from the Nova Scotia Seniors' Pharmacare program were utilized. To ensure confidentiality, all individual (patient and physician) identifiers are encrypted and all computing for this study was conducted on secure Population Health Research Unit (PHRU)³⁵ Dalhousie University computing facilities by PHRU staff.

Topical corticosteroids were identified using Drug Identification Numbers (DINs), a Canadian system used to identify unique drug products. A

DIN is assigned by Health Canada and uniquely identifies the drug product brand/trade name, manufacturer, name and strength of active ingredients, route of administration and pharmaceutical dosage form.³⁶ The completeness of the DIN list was ensured by using: the Health Canada Drug Product Database (DPD)³⁶ the Nova Scotia Formulary³⁷ the Anatomical Therapeutic Chemical Classifications (ATC) list²⁷ and the Compendium of Pharmaceuticals and Specialties (CPS)³⁸ for the study period. Prescriptions were summarized at the individual physician level, and were aggregated by DIN, according to the WHO ATC classification²⁷, into the number of weak (D07AA), moderately potent (D07AB), potent (D07AC) and very potent (D07AD) products prescribed.

Profile Development and Generation

The DEANS Management Committee developed the profile template. Profiles were designed to convey two main messages to prescribers: 1) The potency of the products prescribed by the physician and, 2) the distribution of low, medium, and high cost products prescribed within potency classes. Individual-level physician profiles were generated by PHRU using encrypted patient and physician identifiers. The profiles were sent from the university to the Nova Scotia Department of Health where the identifier was unencrypted for mailing purposes to maintain physician confidentiality. Individual-level physician prescribing profiles were sent by mail in June 2001 to all general practitioners in Nova Scotia who wrote at least one prescription for a topical corticosteroid between April 1, 2000 and March 31, 2001. A letter explaining the profile and a cost comparison chart (see next section) were included in the mailing. This letter provided background for the initiative, information on how to interpret the profiles, basic therapeutic information regarding topical therapy, and it indicated that physicians would be re-profiled in the following year. Re-profiling at the individual physician level was completed and mailed in June 2002 describing prescribing for the fiscal year April 01, 2001 to March 31, 2002 (Figure 1) No comparison to other physicians was provided. The DEANS Management Committee agreed that a 'norm' for prescribing was not appropriate given that the patient case-mix was not available.

FIGURE 1 Topical Corticosteroid Prescribing Profile for Seniors Pharmacare Beneficiaries

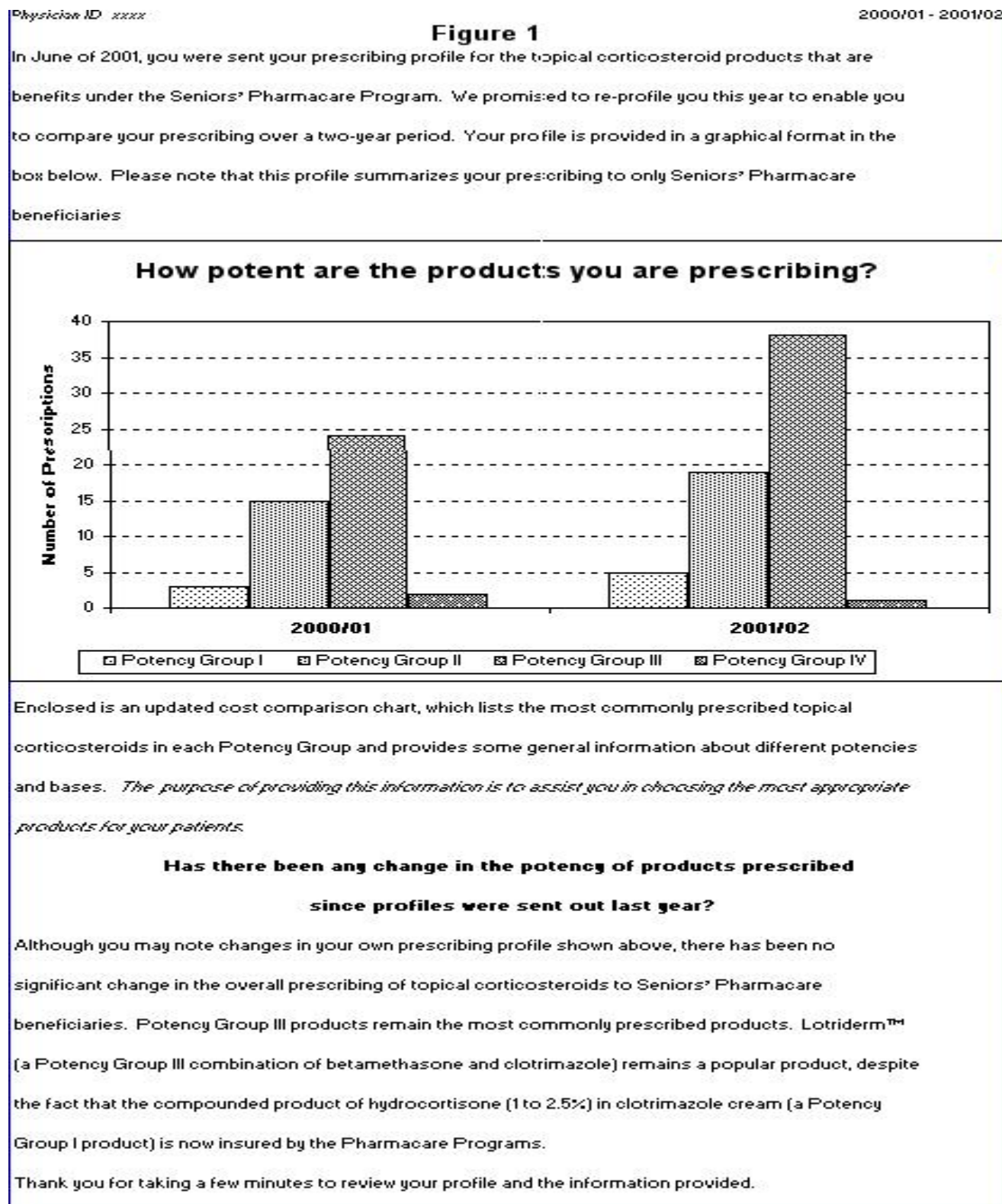


FIGURE 2 Comparison of topical corticosteroids covered in the Nova Scotia Pharmacare Programs

Group I- Least Potent			Group II – Moderate Potency		
Corticosteroid and strength	Brand examples	Cost per gm or mL	Corticosteroid and strength	Brand examples	Cost per gm or mL
Hydrocortisone USP 0.5% or 1%	Cortate Cortoderm Emo-Cort	2 - 20¢	Triamcinolone acetonide 0.025%	Triaderm	6 - 21¢
Hydrocortisone acetate 1%	Cortacet Dermaflex Hyderm	2 - 20¢	Triamcinolone acetonide 0.1%	Aristocort R Triaderm	6 - 21¢
Hydrocortisone 1% with urea 10%	Calmurid or Uremol HC	10 - 18¢	Hydrocortisone valerate 0.2%	Westcort Hydroval	12 - 17¢
Hydrocortisone 1% or 2.5% with camphor / menthol	Sarna HC	10 - 19¢	Triamcinolone acetonide 0.5%	Aristocort C	14 - 21¢
Hydrocortisone 1% to 2.5% in clotrimazole	Pharmacy compound	10 - 21¢	Desonide 0.05%	Desocort Tridesilon	17 - 39¢
Hydrocortisone 1% with a local anesthetic	Pramox HC	12 - 22¢	Clobetasone 0.05%	Eumovate	38 - 40¢
Hydrocortisone acetate 0.5%	Cortacet Hyderm	17 - 21¢	<p>Choosing a Potency</p> <p>In general, the least potent topical corticosteroid to control symptoms should be used.</p> <p>Low and moderate potency products effectively treat acute, inflammatory skin lesions. These products are preferred for areas of the body where the skin is thin (such as the groin and axilla) and for use on infants and the elderly.</p> <p>Potent and very potent products are often required for treating chronic, hyperkeratotic or lichenified lesions, such as psoriasis. They may also be required for areas where the skin is thick, such as the palms and soles.</p>		
Hydrocortisone 2.5% *	Emo-Cort	21 - 25¢			
Hydrocortisone 1% with silicone-type barrier	Barriere-HC Prevex HC	28 - 38¢			
<p>The Cost Ranges</p> <p>The cost ranges are broad because they include <u>all</u> package sizes and <u>all</u> formulations (cream, ointment, lotion, solution, gel and oil).</p> <p>Application Frequency</p> <p>Because the skin acts as a reservoir, most topical corticosteroids are efficacious when applied once or twice daily.</p>					

FIGURE 2 - Cont'd

Group III – Potent			Group IV – Very Potent		
Corticosteroid and strength	Brand examples	Cost per gm or mL	Corticosteroid and strength	Brand examples	Cost per gm or mL
Betamethasone valerate 0.05%	CelestodermV/2	2 - 10¢	Halcinonide 0.1% *	Halog	39 - 49¢
Betamethasone valerate 0.1%	Celestoderm V Valisone	2 - 29¢	Clobetasol 17-propionate 0.05%	Dermasone Dermovate	41 - 49¢
Fluocinolone acetonide 0.025%	Synalar	8 - 41¢	Halobetasol propionate 0.05%	Ultravate	65 - 75¢
Fluocinolone 0.01% * and ***	Derma-Smoothe Synalar	12 - 47¢	<p>Choosing a Base</p> <p>Ointments are occlusive and are generally more potent than creams and lotions. They are preferred for areas where the skin is thicker (palms or soles) or is dry, fissured or scaly.</p> <p>Creams are preferred for oozing lesions, acute and subacute dermatosis and intertriginous areas.</p> <p>Lotions are the least occlusive. They spread easily and are useful for large areas.</p> <p>Solutions and gels are non-greasy and are favoured for use on the scalp or hairy areas. Solutions and gels should be used cautiously on the face. Corticosteroids available in a solution format are indicated by *. Those available as gels are indicated by **.</p> <p>Oils are useful for dry, fissured or scaly scalp lesions. The only corticosteroid available as an oil is indicated by ***.</p>		
Betamethasone dipropionate 0.05%	Diprosone Topisone	20 - 22¢			
Fluocinonide 0.05% **	Lidemol Lidex, Topsylin Lyderm	25 - 65¢			
Betamethasone dipropionate 0.05% glycol	Diprolene Topilene	27 - 56¢			
Desoximetasone 0.05% **	Topicort gel & mild cream	28 - 34¢			
Desoximetasone 0.25%	Topicort	28 - 64¢			
Amcinonide 0.1%	Cyclocort	35 - 59¢			
Betamethasone valerate 0.1% with silicone Barrier	Prevex B	36 - 42¢			
Beclomethasone dipropionate 0.25%	Propaderm	39 - 68¢			
Mometasone furoate 0.1%	Elocom	40 - 56¢			
Diflucortolone valerate 0.1%	Nerisone	42 - 45¢			
Betamethasone dipropionate 0.05% with clotrimazole 1%	Lotriderm	55 - 62¢			

Community Pharmacy Education

In April 2001, all pharmacists in Nova Scotia were mailed the topical corticosteroid cost comparison chart, and a summary of the profiling project in order to assist in providing information to physicians. The summary information was published in the provincial Pharmacy Association bulletin.⁴⁰ As well, all pharmacies received a copy of the cost comparison chart and an outline of the initiative, published in the Pharmacare Bulletin.⁴¹

Data Analysis

The number of prescriptions for topical corticosteroids issued to Nova Scotia seniors covered by the Pharmacare Program was compared for the twelve-month period before and after mailing of the profiles. The claims submitted during the fiscal years April 1, 2000 to March 31, 2001 and April 1, 2001 to March 31, 2002 were examined to identify all topical corticosteroids dispensed to senior beneficiaries covered by the Pharmacare Program.

Overall results were aggregated by utilization and expenditures into the four key potency categories by ATC classification. The data were normally distributed and of equal variance, so the student's t-test was used to test the statistical significance of differences between overall usage and potency categories for both utilization and expenditures.

A modified Laspeyres index⁴² was also used to decompose any changes in topical corticosteroid prescribing into its component factors: price, volume, new drugs, exiting drugs and interaction effects by potency category. It is a forward-looking index that expresses prices and quantities in terms of the previous (base) period. The Laspeyres Index^{43,44} has been validated in similar types of evaluations. All calculations were done using SAS 8.1.⁴⁵

RESULTS

A total of 814 profiles were generated for all general practitioners in Nova Scotia who wrote at least one prescription for a topical corticosteroid between April 1, 2000 and March 31, 2001 for a senior covered by the Nova Scotia Seniors' Pharmacare Program. This represents approximately 85% of licensed/registered general practitioners in the province.⁴⁶ For the 2001/2002 fiscal year, re-profiles were sent to the 814 general practitioners who received the profiles in 2000/2001.

The number of prescriptions for topical corticosteroids per physician profiled was 44.0 in 2000/2001 and 42.8 in 2001/2002 ($p=0.10$), while the expenditures per physician profiled were \$838.94 in 2000/2001 and \$826.81 in 2001/2002 ($p=0.44$) (Table 1).

TABLE 1 Topical corticosteroid prescriptions dispensed to Nova Scotia Seniors' Pharmacare Beneficiaries by Year

Fiscal Year	#Prescriptions ¹	#Prescriptions/ Physician ¹	Total Expenditures ²	Expenditures/ Physician ²	Expenditures/ Prescription ²
2000-01	35,798	44.0	\$682,897.16	\$838.94	\$19.08
2001-02	34,873	42.8	\$673,023.34	\$826.81	\$19.30

¹Number of prescriptions written by the 814 physicians profiled in both 2000-01 and 2001-02

²Expenditures include the pharmacists' professional fees and patient co-payments

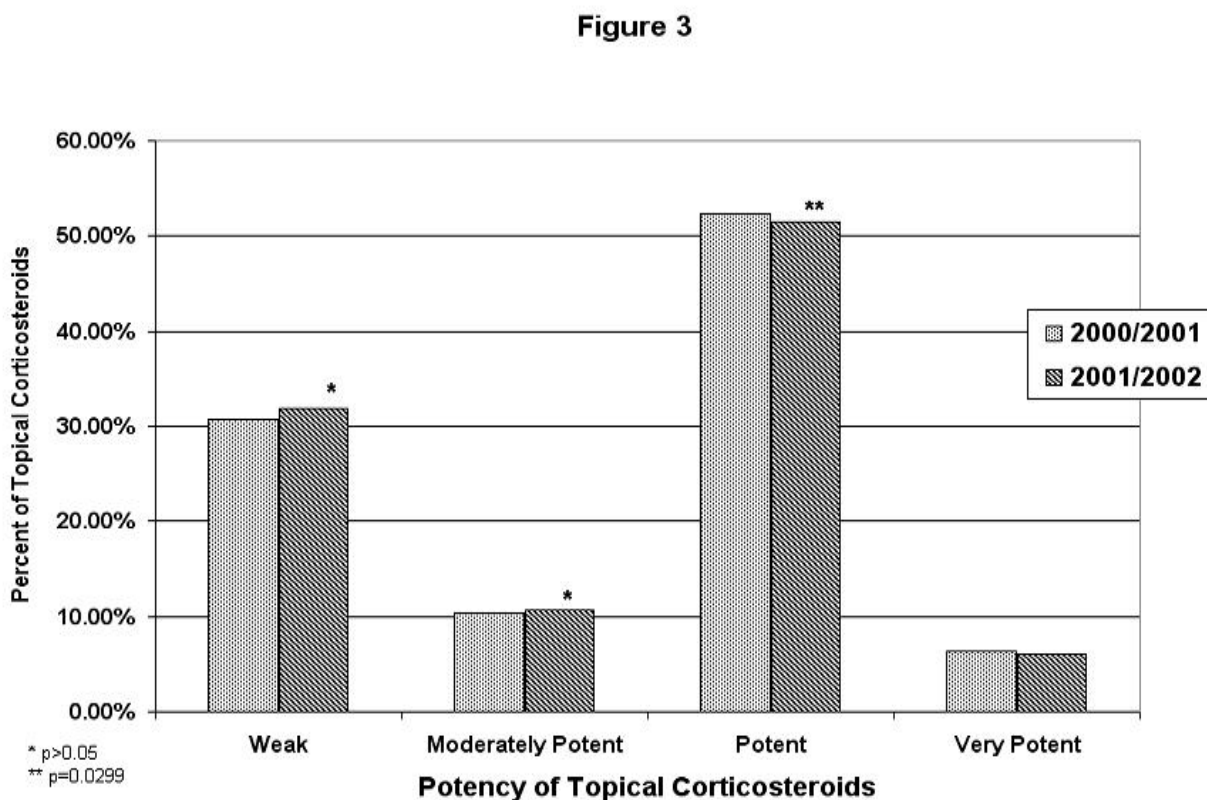
As seen in Figure 3, there was a small, but statistically significant, decrease in prescriptions for potent topical products over the profiling period (52.4% of prescriptions in 2000/2001 versus 51.5% of prescriptions in 2001/2002, $p=0.03$). This translates into an average decrease of one prescription for a potent product per physician (23.1 prescriptions/physician in 2000/2001 versus 22.1

prescriptions/physician in 2001/2002), or 817 fewer prescriptions for potent products over the profiling period. Otherwise, changes in utilization or in expenditures for topical corticosteroids were not statistically different within potency categories from the initial profiles to the re-profiles for general practitioners. When examining changes in prescribing and expenditures using the Laspeyres

Index, there was a seven percent increase in expenditures for weak potency agents driven primarily by an increase in prescribing volume, accompanied by declines in volume for moderately potent agents, potent agents and very potent agents. Declines in the price of prescribed agents within

potency categories also contributed to reducing total expenditures for topical corticosteroids over the profiling period. However, the Laspeyres Index showed that the overall reduction in total expenditures for topical corticosteroids was only \$8,922 (2.4 percent).

FIGURE 3 Topical Corticosteroid Utilization, Nova Scotia Seniors Pharmacare Program, 2000-2002



DISCUSSION

The form of physician profiling used in this study - mailed, unsolicited, centralized, government-sponsored, and involving aggregate data - did not substantially alter the potency or expenditures for topical corticosteroids dispensed to seniors in Nova Scotia. Although mild to moderate potency agents are recommended for the elderly^{31,33}, 52% of topical corticosteroids dispensed in 2000/2001 contained a potent agent; after mailing of the profiles, 51% of topical corticosteroids dispensed in 2001/2002 still contained a potent agent. This decrease, although statistically significant, may not be clinically relevant. The Laspeyres Index did show trends in changes in

prescribing that were consistent with the message of the profiling project, with expenditures for weak potency agents trending upward, while expenditures for more potent agents trending downward, driven primarily by volume changes. However, these changes were small overall.

Randomized controlled studies⁴⁷⁻⁵⁴ have examined changes in drug prescribing behavior using mailed physician profiles with or without the addition of mailed guidelines or prescribing information. Some studies⁴⁷⁻⁵⁰ reported no change in drug prescribing, while others⁵¹⁻⁵⁴ reported a positive change in drug prescribing. Our study was similar to three other studies that showed no change in drug prescribing when utilizing unsolicited mailing of

profiles to all physicians in a wide geographic area.⁵¹⁻⁵³ Studies that did report changes in prescribing focused primarily on benzodiazepine and sedative/hypnotics prescribing,⁴⁷⁻⁴⁹ or enrolled physicians who volunteered for the intervention.^{47,50} Nonetheless, physician profiling may produce small to moderate improvements in professional practice that may be important on a population level.²³

Behavioural theories suggest that the provision of information in isolation does not typically result in translating evidence into changes in prescribing practices.^{6,15,55} These theories also indicate that unsolicited prescriber feedback does not motivate physicians to change, nor does it address barriers to change.^{13,14,17,22,55} Strategies to alter prescribing, therefore, need to address these barriers, incorporating approaches targeting the individual, approaches utilizing social influences, and approaches focusing on the health care delivery system.^{1,2,14,15} A variety of these strategies may be needed to effect change, probably because each strategy affects a different part of the learning process.^{12,13,15,16,21}

Recent evidence indicates that multifaceted interventions may not have a benefit over single interventions.^{2,23} Nevertheless, systematic reviews of profiling initiatives do provide some guidance.^{1,2,14,23} The profiling intervention studied here, although easy to implement on a large scale, may have lacked some features that may be helpful in eliciting behaviour change. The profiles were non-solicited, and passive approaches to change behaviour have not generally been largely effective. A once-yearly message may not be frequent enough to alter prescribing, especially for a group of medications not frequently prescribed as in our physician population (less than four prescriptions for topical corticosteroids per month, on average). Profiles were aggregated at the physician level, so information on an individual patient was not available to physicians, nor was a specific actionable message related to an individual patient included in the intervention. Finally, behaviour change strategies, including physician profiling, may be more successful if incorporated into educational outreach programs where individual approaches and relational capital may assist in influencing behaviour changes.^{56,57} These may be important messages for other groups that plan to conduct large-scale profiling initiatives. As a result of this and other profiling initiatives at DEANS, future initiatives will focus on using

individual targeted patient profiles for educational endeavours when requested by the individual physician.

Strengths and Limitations of the Project

This profiling project used various intervention strategies. The profiles were not designed for use as a single tool, but were combined with written educational information on the therapeutics and costs of topical corticosteroids. A local dermatologist provided feedback on how to present the therapeutic and cost information on the comparison chart in a user-friendly manner and endorsed the information on the chart, along with specific recommendations on the use of less potent products in the elderly.

The multidisciplinary (pharmacy, medicine, epidemiology) and multisectoral (university, hospital, government, professional society) involvement brought in many skills and perspectives. The profiles used in our study also reflect a population-based approach as the Pharmacare Program covers most seniors and Canadian pharmacy administrative claims databases have been found to have good validity.^{58,59}

The profiles may not reflect overall prescribing patterns for topical corticosteroids as the Nova Scotia Seniors' Pharmacare database contains information only on those over 65 years, has no information on prescriptions written but not dispensed, physician samples, or prescriptions dispensed using private insurance. No adjustments were made for patient characteristics, such as age, sex or case-mix. Clinical outcomes were not measured, as the indication for prescriptions is not available in our database. Patients' previous failure with less potent therapy, or the use of potent agents for short bursts rather than for long-term therapy, was not documented. Therefore, it was not possible to determine the appropriateness of therapy. Although this project involved a local opinion leader, and aimed to promote appropriate therapy, the project's highlighting of cost differences in the profile and cost comparison chart design may have been seen primarily as a cost-containment measure. Administrative costs associated with implementing the profiling project were not assessed, nor were physician attitudes. Quasi-experimental designs, such as those employing time-series analysis, may aid in further evaluation of this project.

CONCLUSION

A multi-disciplinary team (DEANS) developed and implemented a profiling initiative focusing on prescribing of topical corticosteroids by Nova Scotia practitioners. An evaluation of the project showed that mailing of unsolicited individual-level profiles, along with a comparison chart, did not impact overall prescribing or expenditures for topical corticosteroids over a two-year period. Further work is needed to determine physician attitudes towards such projects, to examine individual-level changes in prescribing, and to examine prescribing on a monthly basis using a time series analysis.

Acknowledgement and Disclaimer

We would like to thank Dawn Frail, Nova Scotia Department of Health, for manuscript review. The observations and opinions stated are those of the investigators and do not represent the opinions of the Nova Scotia Department of Health.

Financial support

Dr. Ingrid Sketris holds a Canadian Health Services Research Foundation (CHSRF)/ Canadian Institutes of Health (CIHR) chair in Health Services Research, co-sponsored by Nova Scotia Health Research Foundation (NSHRF). Ingrid Sketris and George Kephart receive grants from the Nova Scotia Department of Health to assist in evaluations of DEANS initiatives. Charmaine Cooke was funded in part from a grant from the Nova Scotia Department of Health to Dalhousie University and in part from CHSRF/CIHR/NSHRF through Dr. Sketris' chair. This project has been funded in part by the Drug Evaluation Alliance of Nova Scotia (DEANS) and was carried out at Dalhousie University.

REFERENCES

1. Grimshaw JM, Shirran L, Thomas R et al. Changing provider behavior: an overview of systematic reviews of interventions. *Med Care* 2001 Aug;39(8 Suppl 2):II2-45.
2. Grimshaw JM, Eccles MP, Walker AE, Thomas RE. Changing physicians' behavior: what works and thoughts on getting more things to work. *J Contin Educ Health Prof* 2002;22:237-43.
3. Oxman AD, Thomson MA, Davis DA, Haynes RB. No magic bullets: a systematic review of 102 trials of interventions to improve professional practice. *CMAJ* 1995;53:1423-31.
4. Wennberg DE. Variation in the delivery of health care: the stakes are high. *Ann Intern Med* 1998;128:866-8.
5. Greco PJ, Eisenberg JM. Changing physicians' practices. *N Engl J Med* 1993;329:1271-3.
6. Rogers EM. Lessons for guidelines from the diffusion of innovations. *Jt Comm J Qual Improv* 1995;21:324-8.
7. Freemantle N, Eastaugh J. Using effectiveness studies for prescribing research, part 1. *J Clin Pharm Ther* 2002;27:383-9.
8. Grol R. Changing physicians' competence and performance: finding the balance between the individual and the organization. *J Contin Educ Health Prof* 2002;22:244-51.
9. Soumerai SB, Ross-Degnan D, Fortess EE, Abelson J. A critical analysis of studies of states drug reimbursement policies: research in need of discipline. *Milbank Q* 1993;71:217-52.
10. Institute of Medicine. Committee on Identifying Priority Areas for Quality Improvement. Karen Adams and Janet M. Corrigan, ed. *Priority Areas for National Action: Transforming Health Care Quality*. (2003) <http://www.nap.edu/books/0309085438/html/> (accessed May 12, 2004).
11. Bindman AB. Can physician profiles be trusted? *JAMA* 1999;281:2142-3.
12. Bero LA, Grilli R, Grimshaw JM, Harvey E, Oxman AD, Thomson MA. Closing the gap between research and practice: an overview of systematic reviews of interventions to promote the implementation of research findings. The Cochrane Effective Practice and Organization of Care Review Group. *BMJ* 1998;317:465-8.
13. Weiss KB, Wagner R. Performance measurement through audit, feedback, and profiling as tools for improving clinical care. *Chest* 2000;118(suppl 2):S53-8.
14. Foy R, MacLennan G, Grimshaw J, Penney G, Campbell M, Grol R. Attributes of clinical recommendations that influence change in practice following audit and feedback. *J Clin Epidemiol* 2002;55:717-22.
15. Grol R. Personal paper. Beliefs and evidence in changing clinical practice. *BMJ* 1997;315:418-21.
16. Johnston G, Crombie IK, Davies HT, Alder EM, Millard A. Reviewing audit: barriers and facilitating factors for effective clinical audit. *Qual Health Care* 2000;91:23-36.
17. Kassirer JP. The use and abuse of practice profiles. *N Engl J Med* 1994;330:634-6.
18. Carter AO, Strachan D, Appiah Y. Physician prescribing practices: What do we know? Where do we go? How do we get there? *CMAJ* 1996;154:1649-53.

19. Bloor K, Freemantle N. Lessons from international experience in controlling pharmaceutical expenditure. II: Influencing doctors. *BMJ* 1996;312:1525-7.
20. Davis DA, Thomson MA, Oxman AD, Haynes RB. Changing physician performance. A systematic review of the effect of continuing medical education strategies. *JAMA* 1995;274:700-5.
21. Smith WR. Evidence for the effectiveness of techniques to change physician behavior. *Chest* 2000;118(suppl 2):S8-17.
22. MacKinnon NJ, Lipowski EE. Opinions on provider profiling: telephone survey of stakeholders. *Am J Health Syst Pharm* 2000;57:1585-91.
23. Jamtvedt G, Young JM, Kristoffersen DT, Thomson O'Brien MA, Oxman AD. Audit and feedback: effects on professional practice and health care outcomes (Cochrane Review). In: *The Cochrane Library, Issue 1, 2004*. Chichester, UK: John Wiley & Sons, Ltd.
24. Stern RS. The pattern of topical corticosteroid prescribing in the United States, 1989-1991. *J Am Acad Dermatol* 1996;35(2Pt1):183-6.
25. Campbell CA, Cooke CA, Weerasinghe SD, Sketris IS, McLean-Veysey PR, Skedgel CD. Topical corticosteroid prescribing patterns following changes in drug benefit status. *Ann Pharmacother* 2003;37:787-93.
26. Boguniewicz M, Eichenfield LF, Hultsch T. Current management of atopic dermatitis and interruption of the atopic march. *J Allergy Clin Immunol* 2003;112(suppl 6):S140-50.
27. World Health Organization Collaborating Centre for Drug Statistics Methodology. *ATC Index with DDDs, 2002*. Oslo, Norway: World Health Organization, 2002.
28. Atherton DJ. Topical corticosteroids in atopic dermatitis. *BMJ* 2003;327:942-3.
29. Marek-Thompson TA, Bond CA. Dermatotherapy. In: Young LY, Koda-Kimble MA, eds. *Applied Therapeutics: the clinical use of drugs*. 7th ed. Vancouver, WA: Applied Therapeutics Inc, 2001:10-3.
30. Drugdex® Editorial Staff. Topical corticosteroids - dosing guidelines (Drug Consult). In: Hutchison TA & Shahan DR, eds. *DRUGDEX® System*. Greenwood Village, CO: MICROMEDEX, (Edition expires [06/2004]).
31. Lester RS. Atopic dermatitis. In: Gray J, ed. *Therapeutic Choices*. 4th ed. Ottawa: Canadian Pharmacists Association, 2003:706-711.
32. Zug KA, McKay M. Eczematous dermatitis: a practical review. *Am Fam Physician* 1996;54:1243-50.
33. Atopic dermatitis. In: Herfindal ET, Gourley DRH, eds. *Textbook of therapeutics: drug and disease management*. Philadelphia: Lippincott, Williams & Wilkins, 2000:976-983.
34. Nova Scotia Department of Health. Drug Evaluation Alliance of Nova Scotia (DEANS). <http://www.gov.ns.ca/heal/pharmacare/deans.htm> (accessed on 12 January 2005).
35. Population Health Research Unit (PHRU), Dalhousie University. <http://phru.medicine.dal.ca/> (accessed on 12 January 2005).
36. Health Canada. Drug Product Database. http://www.hc-sc.gc.ca/hpfb-dgpsa/tpd-dpt/index_drugs_dpd_e.html (accessed on 12 January 2005).
37. Department of Pharmaceutical Services. Nova Scotia Formulary. Halifax, NS: Nova Scotia Department of Health, 1999, 2000, 2001.
38. Gillis MC. *Compendium of Pharmaceuticals and Specialties*. 36th edition. Ottawa: Canadian Pharmaceutical Association, 2001.
39. Dalhousie University Faculty of Medicine. Department of Continuing Medical Education. <http://cme.medicine.dal.ca/> (accessed May 12, 2004).
40. Pharmacy Association of Nova Scotia. DEANS Launches an Initiative to Encourage Appropriate Prescribing of Topical Corticosteroids. *The Pharmacist* 2001;4(1):4.
41. Nova Scotia Pharmacare Program. Topical Corticosteroid Cost Comparison Chart. *Pharmacists' Bulletin* 2001; 01(9):2-3.
42. Patented Medicine Prices Review Board. A Description of the Laspeyres Methodology used to Construct the Patented Medicine Price Index (PMPI). Report #S9710. Patented Medicines Prices Review Board: Toronto, June 2000.
43. Getzen TE. Medical care price indexes: theory, construction & empirical analysis of the US series 1927-1990. *Adv Health Econ Health Serv Res* 1992;13:83-128.
44. Danzon PM, Chao LW. Cross-national price differences for pharmaceuticals: how large, and why? *J Health Econ* 2000;19:159-95.
45. SAS 8.1 for Open VMS 7.2-1. Version 8.1. Cary, North Carolina: SAS Institute Inc., 2000.
46. Canadian Institute of Health Information (CIHI). http://secure.cihi.ca/cihiweb/dispPage.jsp?cw_page=statistics_results_topic_physicians_e&cw_topic=Health%20Human%20Resources&cw_subtopic=Physicians (accessed on 12 January 2005).
47. Rokstad K, Straand J, Fugelli P. Can drug treatment be improved by feedback on prescribing profiles combined with therapeutic recommendations? A prospective, controlled trial in general practice. *J Clin Epidemiol* 1995;48:1061-8.
48. Anderson JF, McEwan KL, Hrudehy WP. Effectiveness of notification and group education in modifying prescribing of regulated analgesics. *CMAJ* 1996;154:31-9.

49. Smith DH, Christensen DB, Stergachis A, Holmes G. A randomized controlled trial of a drug use review intervention for sedative hypnotic medications. *Med Care* 1998; 36:1013-21.
50. Hux JE, Melady MP, DeBoer D. Confidential prescriber feedback and education to improve antibiotic use in primary care: a controlled trial. *CMAJ* 1999;161:388-92.
51. O'Connell DL, Henry D, Tomlins R. Randomised controlled trial of effect of feedback on general practitioners' prescribing in Australia. *BMJ* 1999;318:507-11.
52. Sondergaard J, Andersen M, Vach K, Kragstrup J, Maclure M, Gram LF. Detailed postal feedback about prescribing to asthma patients combined with a guideline statement showed no impact: a randomised controlled trial. *Eur J Clin Pharmacol* 2002;58:127-32.
53. Sondergaard J, Andersen M, Stovring H, Kragstrup J. Mailed prescriber feedback in addition to a clinical guideline has no impact: a randomised, controlled trial. *Scand J Prim Health Care* 2003;21:47-51.
54. Pimlott NJ, Hux JE, Wilson LM, Kahan M, Li C, Rosser WW. Educating physicians to reduce benzodiazepine use by elderly patients: a randomized controlled trial. *CMAJ* 2003;168:835-9.
55. Walker AE, Grimshaw J, Johnston M, Pitts N, Steen N, Eccles M. PRIME--PRocess modelling in ImpleMentation research: selecting a theoretical basis for interventions to change clinical practice. *BMC Health Serv Res* 2003;3:22.
56. Sketris IS, Kephart GC, Frail DM, Skedgel C, Allen MJ. The effect of deinsuring chlorpropamide on the prescribing of oral antihyperglycemics for Nova Scotia Seniors' Pharmacare beneficiaries. *Pharmacotherapy*. 2004 Jun;24(6):784-91.
57. Grimshaw J, McAuley LM, Bero LA, et al. Systematic reviews of the effectiveness of quality improvement strategies and programmes. *Qual Saf Health Care* 2003;12:298-303.
58. Tamblyn R, Lavoie G, Petrella L, Monette J. The use of prescription claims databases in pharmacoepidemiological research: the accuracy and comprehensiveness of the prescription claims database in Quebec. *J Clin Epidemiol* 1994;48:999-1009.
59. Levy AR, O'Brien BJ, Sellors C, Grootendorst P, Willison D. Coding accuracy of administrative drug claims in the Ontario Drug Benefit database. *Can J Clin Pharmacol* 2003;10:67-71.