ateness of the calls differs somewhat, but may be dependent upon a more stable population and greater familiarity than is present in the academic setting.

The care resulting in response to the after-hours calls shows private family physicians handling more problems by telephone and sending patients to the emergency room or seeing them personally less often. Again, however, the large clinic group in this study performed more closely to the residents, while the small clinic group created most of the savings in utilization.

Bergman and Rosenblatt conclude that "changes in after-hours utilization reflect practice maturation," but that "after-hours utilization bears a stable relationship to practice size and volume." Thus, they feel "further studies in other settings are needed." These results are useful as the first descriptive study of after-hours telephone

utilization in private family practice in the United States.

This study supports the hypothesis of constancy of utilization in most areas, although the frequency of calls per patient visit is lower in private practice. Physician-patient familiarity is greatest in small, private family practice clinics and may result in different utilization of afterhours services. Before concluding that the patterns of after-hours utilization can be generalized to the private practice setting, further studies should be done outside academic centers with a view toward the effect of the physician-patient relationship on that utilization.

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Social Identities of **Medical Students Oriented Toward Careers in Family Medicine**

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Underlying the design of this study is the assumption that medical students choosing to specialize in family medicine share a common pattern of social characteristics. Gouldner's work¹ on la-

tent and manifest social identities provides a theoretical basis for isolating these characteristics and determining whether they form a pattern that can distinguish family medicine students from their colleagues choosing other specialties.

Methods

This study identifies the manifest role as that of being a medical student. The manifest identities

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that pertain to this role are, therefore, associated with academic credentials and performance. Eight manifest identities were selected for inclusion in this study: the six scores on the Medical College Admissions Test (MCAT), and two undergraduate cumulative grade-point averages (science and total). For purposes of this study, latent identities were defined as characteristics associated with social roles other than being a medical student that may influence attitudes and decisions about specialty orientation, but are generally not considered to be uniformly "relevant" as criteria for admission to medical school. Twenty-one indicators of latent identities were selected for this study from findings of earlier research on family medicine students, other research findings relevant to specialty orientation, and related studies. These indicators related to the respondent's career plans, undergraduate institution, sibling relationships, parents, home town, employment patterns, spouse relationships, and ethnic and religious background.

The study was conducted at a public, northeastern medical school. The study population of 429 students was enrolled in good standing during the 1980-81 academic year. Data on latent identities were collected over a three-month period through a mailed questionnaire specifically designed for this study. Data on manifest identities were collected separately from the medical school's admission data. Manifest and latent social identities for each medical student were matched using social security numbers after data collection of latent identities was completed. A total of 305 students, 71 percent of the study population, responded to the questionnaire. Data analysis was performed by computer using program formats from the Statistical Analysis System Institute.

Results

Specialty orientation to family medicine at the time of the study was used as the classification variable in this study. The family medicine subgroup included 81 students, 27 percent of the respondents. The non-family-medicine subgroup had 224 respondents, accounting for 73 percent of the population.

Manifest Identities

The change in MCAT format in 1978 produced two sets of manifest identity data. The class entering in 1977 took the old MCAT. The remainder of the population took the new MCAT. Joint analysis of old and new MCAT formats is not practical.² For purposes of this study, it was assumed that if the non-MCAT manifest identities, eg, the two grade-point averages for the two subgroups, exhibited a significant correlation with grade-point averages, then the manifest identity data from the first-, second-, and third-year students would be generalizable to the fourth-year students. Analysis showed that the grade-point averages had a relationship significant at the .05 level. Therefore, the analysis of the manifest data was confined to the classes entering in 1978, 1979, and 1980. This subgroup included 231 respondents, of whom 67 were oriented to family medicine and 164 were not oriented to family medicine.

Analysis of variance showed that two of the eight identities (Biology Knowledge, and Skills Analysis: Reading) distinguished, in a statistically significant manner, the family medicine subgroup from the non-family-medicine subgroup. In both instances, the family medicine subgroup scored higher than their colleagues (Table 1).

Latent Identities

Chi-square analysis of discrete variables for family medicine vs non-family-medicine subgroups was used to identify subsets with sufficient power to distinguish, on an individual basis, the two subsets from each other in a statistically significant manner. Three subsets were identified as significant at the .05 level: specialty orientation at admission, type of undergraduate institution, and religious affiliation. Marital status and a younger sister sibling relationship were significant at the .10 level. A separate analysis of variance was done on the following three continuous variables in the latent identities: age at admission to medical school, number of siblings, and birth order. Age presented a statistical significance of .05.

This profile suggests that the family medicine subgroup is coming disproportionately from mar-

Variable	Oriented to Family Medicine (n=67) Mean/SD	Not Oriented to Family Medicine (n=164) Mean/SD
Cumulative-science	3.40/0.47	3.38/0.49
Cumulative-total	3.40/0.41	3.43/0.33
MCAT scores		
Biology knowledge*	9.98/1.70	9.44/1.71
Chemistry knowledge	9.68/2.04	9.91/1.90
Physics knowledge	9.37/2.14	9.55/2.03
Science knowledge	9.75/2.05	9.62/1.99
Skills analysis: reading**	9.46/1.94	8.95/1.55
Skills analysis: quantitative	9.47/2.03	9.30/1.92

ried students with Protestant religious affiliation. These students tend to be older than their colleagues, prefer family medicine at admission to medical school, and attend public undergraduate institutions. The subgroup also appears to have a higher percentage of younger sisters than their colleagues.

The two subgroups were then analyzed using a multinomial logistic analysis procedure. This procedure has been shown to be a preferred method for assessing the collective predictive capability for an array of normal, binomial, and multinomial variables when the variances of key variables are large and the sample is characterized by heterogeneity.³ All subsets that previously presented statistically significant ability to distinguish the two subgroups were included in the logistic regression. In addition, two variables that were slightly above the 0.1 level by chi-square analysis (sex, and younger brother sibling relationship) were also included in the analysis.

The logistic predictive model presented a chisquare value of 49.87 with three degrees of freedom. This model indicates that students oriented to family medicine are best identified through the collective influence of sex (higher frequency of male members than in the student population), type of undergraduate institution (greater frequency of attendance at public undergraduate institutions), and specialty orientation at admission (family medicine orientation rather than an orientation to other specialty subgroups).

Comment

The data suggest that students oriented toward family medicine in this medical school present a profile of latent identities that distinguishes them from students in other specialty subgroups. The study reflects a relatively small population of

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Tablets, containing 125, 250, or 500 mg methyldopa; Oral Suspension, containing 250 mg methyldopa per 5 ml and alcohol 1%

Contraindications: Active hepatic disease, such as acute hepatitis and active cirrhosis; if previous methyldopa therapy has been associated with liver disorders (see Warnings); hypersensitivity

Warnings: It is important to recognize that a positive Coombs test, hemolytic anemia, and liver disorders may occur with methyldopa therapy. The rare occurrences of hemolytic anemia or liver disorders could lead to potentially fatal complications unless properly recognized and managed. Read this section carefully to understand these reactions. With prolonged methyldopa therapy, 10% to 20% of patients develop a positive direct Coombs test, usually between 6 and 12 months of therapy. Lowest incidence is at daily dosage of 1 g or less. This on rare occasions may be associated with hemolytic anemia, which could lead to potentially fatal complications. One cannot predict which patients with a positive direct Coombs test may develop hemolytic anemia. Prior existence or development of a positive direct Coombs test is not in itself a contraindication to use of methyldopa. If a positive Coombs test develops during methyldopa therapy, determine whether hemolytic anemia exists and whether the positive Coombs test may be a problem. For example, in addition to a positive direct Coombs test there is less often a positive indirect Coombs test which may interfere with cross matching of blood. At the start of methyldopa therapy, it is desirable to do a blood count (hematocrit, hemoglobin, or red cell count) for a baseline or to establish whether there is anemia. Periodic blood counts should be done during therapy to detect hemolytic anemia. It may be useful to do a fired counts should be done done and a 6 and 12 months after the start of therapy. If Coombs-positive hemolytic anemia occurs, the cause may be methyldopa and the drug should be discontinued. Usually the anemia remits promptly. If not, corflosteroids may be given and other causes of anemia should be considered. If the hemolytic anemia is related to methyldopa, the drug should not be reinstituted. When methyldopa causes Coombs positivity alone or with hemolytic anemia, the red cell is usually coated with gamma globulin of the IgG (gamma G) class only. The positive Coombs test may not revert to normal until weeks to months after methyldopa is stopped. Should the need for transfusion arise in a patient receiving methyldopa, both a direct and an indirect

Coombs test should be performed on his blood. In the absence of hemolytic anemia, usually only the direct Coombs test will be positive. A positive direct Coombs test alone will not interfere with typing or cross matching. If the indirect Coombs test is also positive, problems may arise in the major cross match and the assistance of a hematologist or transfusion expert will be needed.

Fever has occurred within first 3 weeks of therapy, occasionally with eosinophilia or abnormalities in liver function tests, such as serum alkaline phosphatase, serum transaminases (SGOT, SGPT), bilirubin, cephalin cholesterol flocculation, prothrombin time, and bromsulphalein retention. Jaundice, with or without fever, may occur, with onset usually in the first 2 to 3 months of therapy. In some patients the findings are consistent with those of cholestasis. Rarely tatal hepatic necrosis has some parents the mindings are consistent with hose of cholestasts. Analy idah hepatic herosis has been reported. These hepatic changes may represent hypersensitivity reactions; periodic determination of hepatic function should be done particularly during the first 6 to 12 weeks of therapy or whenever an unexplained fever occurs. If fever and abnormalities in liver function tests or jaundice appear, stop therapy with methyldopa. If caused by methyldopa, the temperature and abnormalities in liver function characteristically have reverted to normal when the drug was discontinued. Methyldopa should not be reinstituted in such patients. Rarely, a reversible reduction of the white blood cell count with primary effect on granulocytes has been seen. Reversible thrombocytopenia has occurred rarely. When used with other antihypertensive drugs, potentiation of antihypertensive effect may occur. Patients should be followed carefully to detect side reactions or unusual manifestations of drug idiosyncrasy.

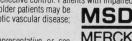
Pregnancy and Nursing: Use of any drug in women who are or may become pregnant or intend to nurse requires that anticipated banefits be weighed against possibile risks; possibility of tetal injury or injury to a nursing infant cannot be excluded. Methyldopa crosses the placental barrier, appears in cord blood, and appears in breast milk.

Precautions: Should be used with caution in patients with history of previous liver disease or dysfunction (see Warnings). May interfere with measurement of: urinary uric acid by the phosphotungstate method, serum creatinine by the alkaline picrate method, and SGOT by colorimetric methods. Since methyldopa causes fluorescence in urine samples at the same wavelengths as catecholamines, falsely high levels of urinary catecholamines may be reported. This will interfere with the diagnosis of pheochromocytoma. It is important to recognize this phenomenon before a patient with a possible pheochromocytoma. Urine exposed to air after willing the diagnosis of pheochromocytoma. Urine exposed to air after willing the diagnosis of pheochromocytoma. Urine exposed to air after willing the diagnosis of the diagnosis of a state the diagnosis of the diagno voiding may darken because of breakdown of methyldopa or its metabolites.

Stop drug if involuntary choreoathetotic movements occur in patients with severe bilateral cerebrovascular disease. Patients may require reduced doses of anesthetics; hypotension occurring during anesthesia usually can be controlled with vasopressors. Hypertension has recurred after dialysis in patients on methyldopa because the drug is removed by this procedure.

Adverse Reactions: Central nervous system: Sedation, headache, asthenia or weakness, usually early and transient; dizziness, lightheadedness, symptoms of cerebrovascular insufficiency, paresthesias, parkinsonism, Bell's palsy, decreased mental acuity, involuntary choreoathetotic movements; psychic disturbances, including nightmares and reversible mild psychoses or depression. Cardiovascular: Bradycardia, prolonged carotid sinus hypersensitivity, aggravation of angina pectoris. Orthostatic hypotension (decrease daily dosage). Edema (and weight gain) usually relieved by use of a diuretic. (Discontinue methyldopa if edema progresses or signs of heart failure anneast). *Gestrointecting barling discharged carotical in the distribution of the anneast*. relieved by use of a diuretic. (Discontinue methyldopa if edema progresses or signs of heart faiture appear.) *Gastrointestinal*: Nausea, vomiting, distention, constipation, flatus, diarrhea, colitis, mild dyness of mouth, sore or "black" tongue, pancreatitis, sialadenitis. *Hepatic*: Abnormal liver function tests, jaundice, liver disorders. *Hematologic*: Positive Coombs test, hemolytic anemia. Bone marrow depression, leukopenia, granulocytopenia, thrombocytopenia. Positive tests for antinuclear antibody; LE cells, and rheumatoid factor. *Allergic*: Drug-related fever, lupus-like syndrome, myocarditis. *Dermatologic*: Rash as in eczema or lichenoid eruption; toxic epidermal necrolysis. *Other*: Nasal stuffiness, rise in BUN, breast enlargement, gynecomastia, lactation, hyperprolactinemia, amenorrhea, impolence, decreased libido, mild arthralgia, myalgia.

Note: Initial adult dosage should be limited to 500 mg daily when given with antihypertensives other than thiazides. Tolerance may occur, usually between second and third months of therapy; increased dosage or adding a diuretic frequently restores effective control. Patients with impaired renal function may respond to smaller doses. Syncope in older patients may be related to increased sensitivity and advanced arteriosclerotic vascular disease;



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this may be avoided by lower doses.



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one medical school, but it has been supported by another study of residents in family medicine residency programs in the same state. These residents, who come from 25 different medical schools, presented a latent identity profile statistically similar to that of the students oriented to family medicine,⁴ suggesting that this profile has credibility beyond the study population per se.

If family medicine is to continue to grow, it must not only increase its supply of first-year postgraduate positions but, most important, also increase the proportion of the graduating medical students choosing to specialize in family medicine. Necessarily, this means either admitting to medical school more students attracted to the family medicine image or enhancing its appeal to the broader spectrum of students than is currently the case. This study suggests that family medicine is most appealing to male students who have attended public undergraduate institutions and who are oriented toward a career in family medicine before they enter medical school. The findings also point to students who are married, Protestant. and somewhat older than their peers. These results could be useful to other medical schools as a model for analysis of characteristics of medical students attracted to family medicine and other specialties and may also be helpful as they relate to selection criteria for entering medical students.

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