LEGISLATIVE RESEARCH COMMISSION

Managed Care Issues



REPORT TO THE 2000 SESSION OF THE 1999 GENERAL ASSEMBLY OF NORTH CAROLINA

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STATE OF NORTH CAROLINA LEGISLATIVE RESEARCH COMMISSION STATE LEGISLATIVE BUILDING RALEIGH, NC 27601



May 4, 2000

TO THE MEMBERS OF THE 1999 GENERAL ASSEMBLY (REGULAR SESSION 2000):

The Legislative Research Commission herewith submits to you for your consideration its 2000 report on managed care. The report was prepared by the Legislative Research Commission's Committee on Managed Care Issues pursuant to G.S. 120-30.17(1).

Respectfully submitted,

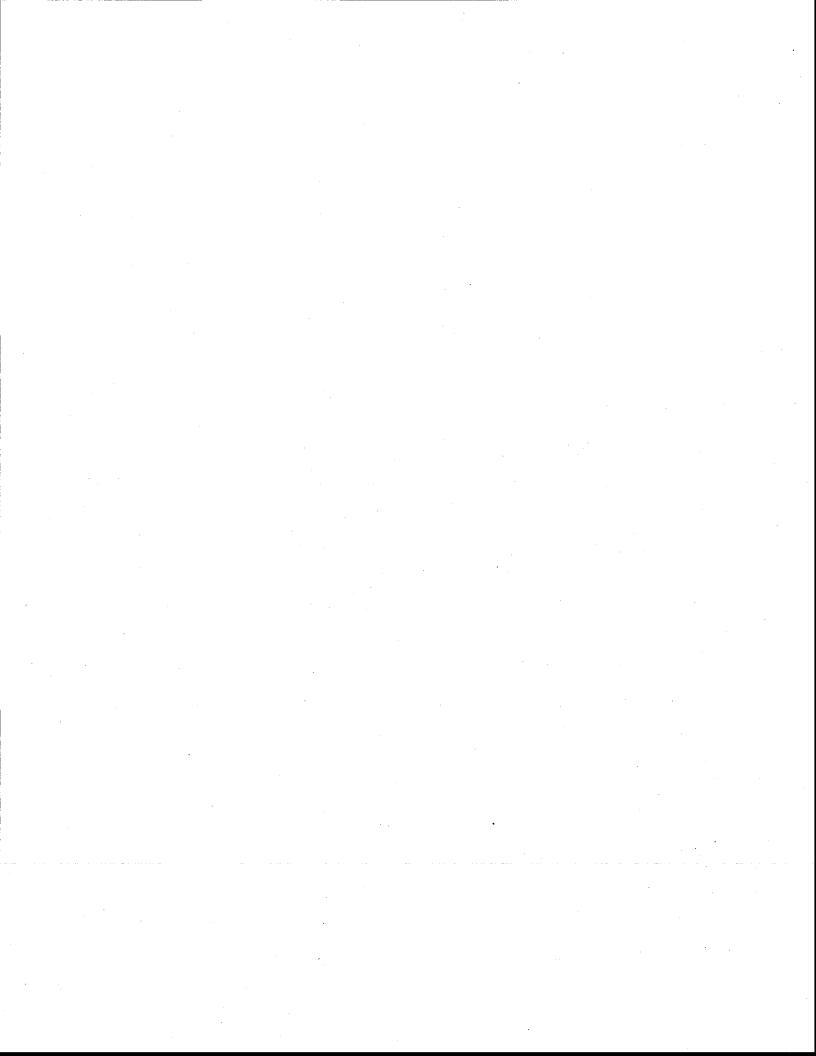
mer B Mark

James B. Black Speaker of the House

Marc Basnight

President Pro Tempore

Cochairs Legislative Research Commission



1999 - 2000

LEGISLATIVE RESEARCH COMMISSION

MEMBERSHIP

President Pro Tempore of the Senate Marc Basnight, Cochair

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Rep. James W. Crawford, Jr. Rep. Beverly M. Earle Rep. Verla C. Insko Rep. William L. Wainwright Rep. Steve W. Wood .

PREFACE

The Legislative Research Commission, established by Article 6B of Chapter 120 of the General Statutes, is the general purpose study group in the Legislative Branch of State Government. The Commission is cochaired by the Speaker of the House and the President Pro Tempore of the Senate and has five additional members appointed from each house of the General Assembly. Among the Commission's duties is that of making or causing to be made, upon the direction of the General Assembly, "such studies of and investigations into governmental agencies and institutions and matters of public policy as will aid the General Assembly in performing its duties in the most efficient and effective manner" (G.S. 120-30.17(1)).

The Legislative Research Commission, prompted by actions during the 1998 Session and 1999 Sessions, has undertaken studies of numerous subjects. These studies were grouped into broad categories and each member of the Commission was given responsibility for one category of study. The Cochairs of the Legislative Research Commission, under the authority of G.S. 120-30.10(b) and (c), appointed committees consisting of members of the General Assembly and the public to conduct the studies. Cochairs, one from each house of the General Assembly, were designated for each committee.

The study of managed care was authorized by Section 2.1 of Chapter 395 of the 1999 Session Laws (Regular Session, 1999). Part II of Chapter 395 allows for studies authorized by that Part for the Legislative Research Commission to consider

Senate Bill 1089 and H.J.R 1461 in determining the nature, scope and aspects of the study. Section 1 of Senate Bill 1089 reads in part:

:"Section 1. The Legislative Research Commission may study the following issues relating to managed care:

(1) Quality of care.

(2) Cost of care and cost-containment measures of managed care plans.

- (3) Provider selection and retention, including whether any provider willing to meet the terms, conditions, and standards, including the terms of reimbursement, of a managed care plan should be allowed to participate in the plan (SB 1090).
- (4) The establishment of a consumer's insurance advocate to appear on behalf of consumers in actions and proceedings involving insurance products and services, to publish consumer-oriented insurance information, to respond to consumer complaints, to advocate on behalf of State employees with respect to insurance products and services provided to those employees, and to report on its activities (SB 1108).
- (5) Requiring the Commissioner of Insurance to submit reports on the status of managed care to the General Assembly and Governor, including information on outcome data, utilization review, provider access, and related matters (SB 1089).
- (6) Providing funds to pay for prescription drugs for certain persons ineligible for Medicaid but whose income no more than two hundred percent (200%) above poverty level, and who has a life-threatening disease or condition for which the drugs have been prescribed (SB 1109).
- (7) Liability of providers and managed care plans.
- (8) Any other issues concerning the delivery of managed care."

The relevant portions of Chapter 395 are included in Appendix A.

The Legislative Research Commission authorized this study under authority of G.S. 120-30.17(1) and grouped this study in its Insurance and Managed Care Issues area under the direction of Representative Verla Insko. The Committee was chaired by Senator Allen Wellons and Representative Edd Nye The full membership of the Committee is listed in Appendix B of this report. A committee notebook containing the committee minutes and all information presented to the committee will be filed in the Legislative Library by the end of the 1999-2000 biennium.

COMMITTEE PROCEEDINGS

The Legislative Research Commission's Managed Care Issues Committee met five times prior to the 2000 Session of the 1999 General Assembly. The Committee was charged with studying managed care issues.

At its first meeting, the Committee heard a presentation from Ms. Barbara Morales Burke, Senior Deputy Commissioner with the Department of Insurance. Ms. Burke presented information concerning regulation and market performance in managed care in North Carolina. She stated that there are fewer managed care companies in the State now than in the past. Several companies have pulled out of the State. The State had ten (10) HMOs in 1992, and the number peaked in 1997 with 24 full service HMOs. Currently, 20 HMOs exist in the State and that number may go down by one or two due to mergers. Only three HMOs have been consistently profitable; the remainder lose money on a regular basis. In response to that, premiums increased in 1999 and 2000 for the first time in several years. A recent survey conducted by William Mercer, Inc., a large benefits consulting company, found that North Carolina is experiencing the largest increase in premiums of any state in the country. North Carolina's average increase was 9.9% compared to 7.5% nationally. Ms. Morales Burke stated that the higher increase in North Carolina could be due to HMOs keeping their rates on the low side for the past several years. Now, the HMOs facing financial losses and increased medical costs, and must increase rates.

Mr. Bill Stevens, Deputy Commissioner, Consumer Services Division within the Department of Insurance, spoke to the committee regarding the division and consumer HMO complaints. His division handles approximately a hundred thousand phone calls a year, regarding all types of complaints. Complaints raised by consumers include:

- Denial of claims.
- Delay of settlements.
- Medical necessity determination issues.
- Access to care issues.
- Concerns about premiums.

Ms. Nancy O'Dowd, Deputy Commissioner, Managed Care and Health Benefits Division within the Department of Insurance, spoke to the Committee on the activities of the Department in relation to the regulation of the managed cared industry in North Carolina. Ms. O'Dowd stated that with the passage of Senate Bill 594 during this past session of the General Assembly, HMOs are subject to G.S. 58-3-100(c) effective January 1, 2000. Under G.S. 58-3-100(c), HMOs are required to acknowledge claims in 30 days. This provision gives the Department enforcement authority to take action when it is determined that a carrier's claims payment practices are improper and unfair.

At its second meeting, Ms. Barbara Morales Burke summarized for the Committee information concerning the Federal Patient's Rights Legislation. Two bills both addressing the issue, Senate Bill 1344 and House Bill 2990, are currently in Congressional conference committee. She indicated that states, not Congress, have been the innovators and parties to act in the area of patient's rights in managed care issues.

Mr. Adam Searing, Project Director for the North Carolina Health Access Coalition, spoke to the Committee about accountability in health plan decision making. He stated that the Harvard School of Public Health and the Kaiser Family Foundation surveyed doctors and nurses nationally and asked them how often care is denied to their patients and does that denial impact the care the patients are receiving. Nine out of ten doctors said their patients are experiencing health plan denial for coverage of services and one-half of all nurses surveyed said that a health

plan denial of care resulted in a decline in their patients health. Mr. Searing stated that three states have passed laws allowing a patient to recover damages if a health plan makes a negligent decision. Thirty-seven other states have expressed an interest in looking at this issue.

Mr. Jim Kerr, an attorney with Smith Anderson Law Firm representing the North Carolina Medical Society, also addressed the Committee on the issue of health plan accountability. He said that the ERISA legislation was designed to prevent abuses in the administration of pension plans. The debate in the courts is whether Congress ever intended to provide a shield for negligent actions of health plans when ERISA was first adopted more than twenty years ago.

Ms. Peg O'Connell, Director of External Relations, Medical Review of North Carolina, Inc., made a presentation concerning the Medicare Independent Appeals and Grievance Review and the role of the Peer Review Organization. The Medical Review of North Carolina, Inc. works to assure health care services provided to the State's Medicare beneficiaries are medically necessary, are furnished in the appropriate setting, and meet professionally recognized standards of health care with respect to quality.

At its third meeting, Dr. Pam Silberman, Vice-President of the Institute of Medicine and a member of the Committee, spoke about the *Consumer Guide to Health Plan Selection*, a webbased guide that provides information to assist consumers and businesses in the selection of health care plans. This consumer guide project, started a year ago, provides information on:

- Understanding managed care.
- Consumer protection.
- Member responsibility.
- Questions to ask your health plan.

The guide provides a mechanism by which a consumer may compare the 13 primary HMOs in North Carolina, and also provides "hot links" to other managed care resources.

Mr. Henry Landsberger, with the American Association of Retired Persons (AARP), presented the results of a statewide survey sponsored by AARP. The survey, *What North Carolina's Citizens Think of Managed Care, and What They Want*, found support for both an ombudsman program and for a third stage independent appeals entity.

Mr. Paul Mahoney, Executive Director of the NC Association of Health Plans addressed the Committee on the issue of the uninsured. As a result of state and federal mandates, companies are providing greater benefits today than they did 10 years ago. However, with increased benefits come increasing costs pressures. This double-edged sword has resulted in companies providing more benefits to fewer people. Estimates indicate that for every one percent increase in premiums, 200,000 people drop coverage. Ten years ago, there were 37 million uninsured Americans. Today, there are 47 million uninsured persons. Mr. Mahoney also acknowledged there are significant problems in the delivery of health care. A recent report by the Institute of Medicine puts the annual death toll for medical errors between 44,000 and 100,000 a year.

Mr. Steve Keene, Director of Government Affairs, North Carolina Medical Society, spoke in support of an independent review process along with health plan liability legislation. He said the review process needs to be external and there should be a requirement that health plans follow the results of the review.

Representative Joe Hackney presented the fourth edition of HB1133, Health Insurance Liability, to the Committee. The bill provides a cause of action to individuals harmed or damaged by the decisions of a managed care entity. Representative Hackney stated HB1133 plugs a gap in the liability system with respect to health care decisions made by the HMOs. The need for such legislation is created due to the lack of substantial remedies for certain persons who are damaged by health care decisions made by the managed care entity.

Mr. Alan Hirsch, Senior Deputy Attorney General, Consumer Protection and Anti-Trust Division of the NC Department of Justice, conveyed to the Committee three issues his Department supports:

- The rights of states to make the determination as to the medical care their citizens receive.
- 2) Patients Bill of Rights.
- 3) Enforcement mechanisms to ensure patients receive what they need.

At the fourth meeting of the Committee, Ms. Elizabeth Ouzts, Executive Director of North Carolina Public Interest Research Group (NC PIRG), an environmental and consumer advocacy organization, spoke in favor of legislation:

- Establishing liability on behalf of managed care entities.
- Establishing independent external review.
- Creating an ombudsman program.

The Department of Insurance also briefed the Committee on proposed legislation regarding several aspects of managed care. The proposed bills were:

- 1) HMO Insolvency
- 2) Prompt Payment
- 3) External Review
- 4) HMO Liability.

Also presented to the Committee by the North Carolina Association of Health Plans were lists of proposed changes to the draft bills. The Committee discussed the bills and adopted the concepts presented in the proposed legislation from the Department of Insurance. However, the Committee desired to have the Department of Insurance work with the North Carolina Association of Health Plans to address issues raised by the North Carolina Association of Health Plans. The Committee requested that revised drafts be presented at the next meeting.

At its fifth meeting, the Committee heard presentations from Dr. Melvin T. Pinn, Jr. and Sharon Martin, RN, both speaking on Medicaid Managed Care topics. Dr. Pinn, medical director with the Wellness Plan of NC, briefed the Committee on his organization and its function. Ms. Martin, Medicaid case manager, spoke regarding the delivery of services to the clients and the necessity of community resource coordination.

The Committee was also presented with a draft copy of the report, including each proposed bill. The Committee discussed the draft bills as presented. Linda Attarian, Staff Counsel, presented one additional bill, entitled "Internal Review Panelists," to the Committee. Discussion on the proposed bill focused on the questionable need for clinical peers at the first level grievance review. Committee member's views on this were mixed, and all agreed that further study was warranted.

At its sixth meeting, the Committee discussed and approved an amended final report.

FINDINGS AND RECOMMENDATIONS

Upon discussion and debate, the Joint Legislative Research Commission's Committee on Managed Care Issues makes the following findings and recommendations:

<u>1. HEALTH CARE LIABILITY.</u>

A. Findings

Based upon the presentations and briefings, the Committee finds that current law in North Carolina fails to adequately provide concise remedies to consumers with respect to health care decisions made by managed care entities. In support of this finding, the Committee states the following:

- Substantial remedies for certain persons damaged by health care decisions made by a managed care entity do not exist currently in North Carolina.
- Trends in the industry indicate that, increasingly, health care treatment decisions are being made by individuals or entities that are not the treating physician.
- A wide variety of entities are integrating the functions of determining the treatment provided, providing the decided upon treatment, and paying for the treatment. This integration of functions is resulting in a breakdown of traditional distinctions of entities and in consumers being left unprotected by the legal system.
- Historically, the General Assembly has acted to protect consumers from egregious harms due to the improper actions of other individuals and entities.

B. Recommendations

Therefore, the Committee recommends the attached bill entitled "HMO Liability." In summary, the bill does as follows:

- Provides a legal remedy to consumers damaged by the health care treatment decisions made by a managed care entity.
- Establishes a standard of care for managed care entities.
- Establishes defenses that may be asserted by the managed care entity.
- Makes void indemnification and hold-harmless clauses in the contracts between the managed care entity and the health care provider.

2. PROMPT PAYMENT OF CLAIMS.

A. Findings

Based upon the presentations and briefings, the Committee finds that insurers are not consistently and cooperatively paying health care providers within a reasonable time frame for the services rendered to the insured. In support of this finding, the Committee states the following:

• It appears that the health care providers are often requested to submit claims numerous times.

- It appears that when inquiries are made of the insurer regarding specific claims, the health care provider or insured are told that the claim was lost or not submitted.
- Current law does not allow for interest to accrue on unpaid claims for approved health care services, and such a provision would provide incentive for the insurer to pay claims in a timely manner.

B. Recommendations

Therefore, the Committee recommends the attached bill entitled "Prompt Pay." In summary, the bill does as follows:

- Within 30 days of a claim being submitted by a health care provider or facility, an insured or the insured's legal representative, the insurer must do one of the following:
 - 1. pay the claim.
 - 2. send notice of denial.
 - 3. send notice of that the proof of loss is either inadequate or incomplete.
 - 4. send notice that the claim was not submitted on the proper form(s).
- Claims not processed in accordance with the proposed legislation will accrue interest of 18 percent per annum.

3. HMO INSOLVENCY.

A. Findings

Based upon the presentations and briefings, the Committee finds that current law in North Carolina needs to be adjusted to reflect the possibility of a healthcare maintenance organization becoming insolvent, and to provide for that entity's clients to be protected in the event of insolvency. In support of this finding, the Committee states the following:

- Under current law, other types of insurers are required to participate in a guaranty association in an effort to protect the consumer/customers of the entity in the event the entity is declared insolvent. Private pension plans are also encouraged to participate in such guaranty associations.
- Under current law, North Carolina does not require that HMOs participate in such a guaranty association, and as a result, consumers are left unprotected in the event of the insolvency of their HMO insurer.
- Historically, the General Assembly has acted to protect consumers from harm caused by such events.

B. Recommendations

Therefore, the Committee recommends the attached bill entitled "HMO Solvency." In summary, the bill does as follows:

• Allows the Commissioner of Insurance to make an assessment of no more than 2% of the HMO's average premiums received in NC during the 3 calendar years preceding the year in which an HMO was declared insolvent. The money collected is be used for: payment of claims, the continuation of coverage, and administrative purposes.

- Protects the insured from losing continuance coverage if the HMO is declared insolvent by providing \$300,000 expenditures on behalf of the insured and continued coverage of the lesser or one year or the remaining term of the insured's contract with the HMO.
- Establishes priority of claims.

4. EXTERNAL REVIEW.

A. Findings

Based upon presentations, the Committee finds that current law in North Carolina does not provide an orderly, mandated process whereby individuals not affiliated with the insurer and/or provider review insurance claims. In support of this finding, the Committee states the following:

- It would be beneficial to the consumer, the health care provider, and the insurer to establish a process to review, outside of the insurer's control and domain, contested claims.
- The Department of Insurance, the insurance industry, and the health care industry agree that an independent external review process is needed.
- It is desirable to establish a framework for review of claims that has a clinical peer review with an individual knowledgeable of the area to which the claim refers.

B. Recommendations

Therefore, the Committee recommends the attached bill entitled "External Review." In summary, the bill does as follows:

- Establishes an independent, external review process to be utilized after the exhaustion of the internal review process.
- Establishes time frames in which the review must be completed.
- Establishes qualifications for the individuals sitting in review capacity.

5. OMBUDSMAN PROGRAM.

Based upon presentations, the Committee agrees that such a program would be beneficial to the citizens of North Carolina. However, the Committee believes that the development of such a program should be discussed in further detail. Therefore, it is the intent of the Committee to make a report to the 2001 General Assembly regarding the establishment of an Ombudsman Program for Managed Care.

6. CLINICAL PEERS ON INTERNAL REVIEW.

A. Findings

Based upon presentations, the Committee finds that current law in North Carolina does not provide a clinical peer review during the internal grievance and review process. In support of this finding, the Committee states the following:

- It would be beneficial to the consumer, the health care provider, and the insurer to establish a process for the review of contested claims by clinical peers during the internal review process.
- The Department of Insurance, the insurance industry, and the health care industry agree that a clinical peer review process is needed.
- It is desirable to establish a framework for review of claims that has a peer review with an individual knowledgeable of the area to which the claim refers.

B. Recommendations

Therefore, the Committee recommends, by mixed vote, the attached bill entitled "Internal Review Panelists" for further discussion and consideration. In summary, the bill does as follows:

- Establishes a clinical peer review to be utilized in the internal grievance and review process.
- Establishes qualifications for the individuals sitting in review capacity.
- Requires that the clinical peer be licensed in North Carolina in the same capacity as the health care provider making the health care treatment decision at issue in the review.

APPENDIX A

<u>CHAPTER 395</u> 1999 Session Laws (1999 Session)

AN ACT TO AUTHORIZE STUDIES BY THE LEGISLATIVE RESEARCH COMMISSION, TO CREATE VARIOUS STUDY COMMISSIONS, TO DIRECT STATE AGENCIES AND LEGISLATIVE OVERSIGHT COMMITTEES AND COMMISSIONS TO STUDY SPECIFIED ISSUES, AND TO AMEND OTHER LAWS.

The General Assembly of North Carolina enacts:

PART I.----TITLE

Section 1. This act shall be known as "The Studies Act of 1999".

PART II.----LEGISLATIVE RESEARCH COMMISSION

Section 2.1. The Legislative Research Commission may study the topics listed below. When applicable, the bill or resolution that originally proposed the issue or study and the name of the sponsor is listed. Unless otherwise specified, the listed bill or resolution refers to the measure introduced in the 1999 Regular Session of the 1999 General Assembly. The Commission may consider the original bill or resolution in determining the nature, scope, and aspects of the study. The following groupings are for reference only:

(2) Insurance and Managed Care Issues:

- a. Managed care issues, including any willing provider, patients' rights, managed care entity liability, office of consumer advocacy for insurance, prompt payment of health claims, and related issues (S.B. 1089 Harris, H.J.R. 1461 Mosley).
- b. Mental health and chemical dependency parity (H.B. 713 Alexander; S.B. 836 Martin of Pitt).
- c. Health reform recommendations of the Health Care Planning Commission and its advisory committees (established by Section 1.2 of Chapter 529 of the 1993 Session Laws) that have not been implemented but are still needed and other health reform issues (Insko).

d. Pharmacy choice/competition (H.B. 1277 - Cole; S.B. 137 - Rand).

Section 21B.4. The Commission may make an interim report to the 1999 General Assembly, Regular Session 2000, upon its convening, and shall make its final report to the 2001 General Assembly upon its convening, and to the Governor. Upon submitting its final report, the Commission shall expire.

Section 21B.5. Upon approval of the Legislative Services Commission, the Legislative Services Officer shall assign appropriate professional staff from the Legislative Services Office of the General Assembly to assist with the study. The House of Representatives' and the Senate's Supervisors of Clerks shall assign clerical staff to the Commission, upon the direction of the Legislative Services Commission. The

Commission may meet in the Legislative Building or the Legislative Office Building upon the approval of the Legislative Services Commission.

Section 21B.6. The Speaker of the House of Representatives and the President Pro Tempore of the Senate shall each designate a cochair of the Commission. The Commission shall meet upon the call of the cochairs. A quorum of the Commission is 10 members. While in the discharge of its official duties, the Commission has the powers of a joint committee under G.S. 120-19 and G.S. 120-19.1. Members of the Commission shall receive per diem, subsistence, and travel allowances in accordance with G.S. 120-3.1, 138-5, or 138-6, as appropriate.

Section 21B.7. From funds appropriated to the General Assembly, the Legislative Services Commission shall allocate funds for the expenses of the Study Commission on Children With Special Needs.

APPENDIX B

COMMITTEE MEMBERSHIP

Sen. Allen Wellons, Co-Chair Sen. Charlie Dannelly Dr. James Elliot, Jr. Mr. Hank Estep Sen. Oscar Harris Dr. Pam Silberman Dr. Steven Michael Willen Rep. Edd Nye, Co-Chair Rep. W. Pete Cunningham Rep. Zeno Edwards Rep. Larry T. Justus Rep. Martin L. Nesbitt Ms. Elizabeth O'Keefe Mr. Thomas L. West

APPENDIX C

SESSION 1999

HMO LIABILITY THIS IS A DRAFT 25-APR-00 16:16:17

Short Title: Managed Care Entities Liable for Damages. (Public)

Sponsors:

Referred to:

1 A BILL TO BE ENTITLED 2 AN ACT TO PROVIDE THAT A MANAGED CARE ENTITY PROVIDING A HEALTH BENEFIT PLAN IS LIABLE FOR DAMAGES FOR HARM TO ITS INSUREDS OR 3 ENROLLEES CAUSED BY THE MANAGED 4 CARE ENTITY'S FAILURE TO EXERCISE ORDINARY CARE. 5 6 The General Assembly of North Carolina enacts: 7 8 Section 1. Chapter 90 of the General Statutes is 9 amended by adding a new Article to read: 10 "ARTICLE 1G. 11 "Health Care Liability. 12 "§ 90-21.50. Legislative findings and intent. (a) The General Assembly finds that a wide variety of entities 13 14 are integrating the functions of paying for health care, 15 determining what health care is paid for, and providing the care. 16 This integration of functions is breaking down traditional 17 distinctions. Increasingly, payor determinations are governing 18 health care and controlling decisions that in the past were the 19 exclusive domain of health care providers and patients. The 20 General Assembly further finds that this integration of functions

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SESSION 1999

		perative that managed care entities be held fully
2	responsible	for the consequences of their decisions, much as
		professionals have been held responsible for the
4		of their decisions.
5	(b) The	state's interest in regulating the business of
6	insurance as	provided in this Article is to protect insurance
7	purchasers an	nd their beneficiaries, including employees, their
8	dependents a	nd families, and any other patients covered by
9	private emplo	over-sponsored benefit plans, from the harm that may
10	occur when ma	anaged care entities, act improperly. To this end,
11	health care	providers rather than managed care entities are in
12	charge of pat	ient care.
13	(c) It is	the intent of the General Assembly in enacting this
14	Article to en	sure that adequate State law remedies exist for all
15	persons who a	are subject to the wrongful acts of those entities
16	that contrac	t to provide insurance for the health of North
17	Carolina cit:	izens. The existence of these remedies and the
18	deterrent eff	ects of these remedies are necessary to protect the
19	health and sa	fety of the residents of this State.
20	"§ 90-21.51.	Definitions.
21	As used in	this Article, unless the context clearly indicates
22	otherwise, th	
23	<u>(1)</u>	'Health benefit plan' means an accident and health
24		insurance policy or certificate; a nonprofit
25		hospital or medical service corporation contract; a
26	•	health maintenance organization subscriber
27		contract; a plan provided by a multiple employer
28		welfare arrangement; or a plan provided by another
29		benefit arrangement. 'Health benefit plan' does not
30		mean any plan implemented or administered by the
31		North Carolina or United States Department of
32		Health and Human Services, or any successor agency,
33		or its representatives. 'Health benefit plan' does
34		not mean any of the following kinds of insurance:
35		a. Accident.
36		b. Credit.
37		c. Disability income.
38		d. Long-term or nursing home care.
39		
40		e. Medicare supplement. f. Specified disease.
-		

HMO Liability

1		g.	Dental or vision.
2		h.	Coverage issued as a supplement to liability
3			insurance.
4		i.	Workers' compensation.
5		<u>i.</u> j.	Medical payments under automobile or
6			homeowners'.
7		k.	Hospital income or indemnity.
8		<u>k.</u> 1.	Insurance under which benefits are payable
9			with or without regard to fault and that is
10			statutorily required to be contained in any
11			liability policy or equivalent self-insurance.
12		m.	Short-term limited duration health insurance
13		<u></u>	policies as defined in Part 144 of Title 45 of
14	· ·		the Code of Federal Regulations.
15	(2)	'Heal	Lth care provider' means:
16		a.	An individual who is licensed, certified, or
17		<u> </u>	otherwise authorized under this Chapter to
18			provide health care services in the ordinary
19			course of business or practice of a profession
20			or in an approved education or training
21			program; or
22			A health care facility, licensed under
23			Chapters 131E or 122C of the General Statutes,
24			where health care services are provided to
25			patients;
26			th care provider' includes:
27			1. An agent or employee of a health care
28			facility that is licensed, certified, or
29			otherwise authorized to provide health
30			care services;
31			2. The officers and directors of a health
32		-	care facility; and
33			3. An agent or employee of a health care
34		-	provider who is licensed, certified, or
35			otherwise authorized to provide health
36			care services.
37	<u>(3)</u>	'Heal1	th care service' means a health or medical
38		proced	
39		provid	ler that:

· 1			a. Provides testing, diagnosis, or treatment of a
2			human disease or dysfunction; or
3			b. Dispenses drugs, medical devices, medical
4			appliances, or medical goods for the treatment
5			of a human disease or dysfunction.
6		(4)	'Health care treatment decision' means a
7			determination that:
8		,	a. Is made by a managed care entity;
9		•	b. Governs the extent to which health care
10			services are provided for, arranged for, paid
11			for, or reimbursed under a health benefit
12			plan; and
13			c. Affects the quality of the diagnosis, care, or
14			treatment provided under the health benefit
15			plan to an enrollee or insured of the health
16			benefit plan.
17		(5)	'Insured or enrollee' means a person that is
18		<u>`````</u>	insured by or enrolled in a health benefit plan
19			under a policy, plan, certificate, or contract
20			issued or delivered in this State by an insurer.
21		(6)	'Insurer' means any entity that is or should be
22		<u>a</u> dana sa	licensed under Articles 6, 7, 16, 49, 65, or 67 of
23			this Chapter.
24		(7)	'Managed care entity' means an insurer that:
25			a. Delivers, administers, or undertakes to
26			provide for, arrange for, or reimburse for
27			health care services, or assumes the risk for
28			the delivery of health care services; and
29			b. Has a system or technique to control or
30			influence the quality, accessibility,
31			utilization, or costs and prices of health
32			care services delivered or to be delivered to
33			a defined enrollee population.
34			'Managed care entity' does not include: (i) an
35			employer purchasing coverage or acting on
36			behalf of its employees or the employees of
37			one or more subsidiaries or affiliated
38			corporations of the employer, or (ii) a health
39			care provider.
40		(8)	'Ordinary care' means:
	-		

1	
2	degree of care that a carrier or managed care
3	entity of ordinary prudence would use under
4	
5	b. For a person that is an agent of employee of a
6	carrier or managed care entity, that degree of
7	care that a person of ordinary prudence in the
8	same profession, specialty, or area of
9	practice as the person would use in the same
10	or similar circumstances.
11	(9) 'Physician' means:
12	a. An individual licensed as a medical doctor
13	under Article 1 of this Chapter to practice
14	medicine in this State;
15	b. A professional association or corporation
16	comprising medical doctors and organized under
17	Chapter 55B of the General Statutes; or
18	c. A person or entity wholly owned by medical
19	doctors.
	"§ 90-21.52. Duty to exercise ordinary care; liability for
	damages for harm.
22	(a) Each managed care entity for a health benefit plan has the
	duty to exercise ordinary care when making health care treatment
	decisions and is liable for damages for harm to an insured or
	enrollee proximately caused by its failure to exercise ordinary
	care.
27	(b) In addition to the duty imposed under subsection (a) of
28	this section, each managed care entity for a health benefit plan
	is liable for damages for harm to an insured or enrollee
30 j	proximately caused by the health care treatment decisions made
31]	
32	(1) Its agents, ostensible agents, or employees; or
33	(2) Representatives that are acting on its behalf and
34	over whom it has the right to exercise influence or
35	control which results in the failure to exercise
36	ordinary care.
37	(c) It shall be a defense to any action brought under this
	section against a managed care entity for a health benefit plan
39 t	that:

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9	care provider to the insured or enrollee.
10	
11	care entity, a finding that a physician or health care provider
12	is an agent or employee of the managed care entity may not be
13	based solely on proof that the physician or health care provider
14	appears in a listing of approved physicians or health care
	providers made available to insureds or enrollees under the
16	managed care entity's health benefit plan.
17	(e) An action brought under this Article is not a medical
18	malpractice action as defined in Article 1B of this Chapter. A
	managed care entity may not use as a defense in an action brought
	under this Article any laws that prohibit the practice of
21	medicine by a corporate entity or by a health maintenance
22	organization.
23	(f) A managed care entity shall not be liable for the
24	independent actions of a health care provider, who is not an
	agent or employee of the managed care entity, when that health
26	care provider fails to exercise the standard of care required by
27	G.S. 90-21.12. A health care provider shall not be liable for
28	the independent actions of a managed care entity when the managed
29	care entity fails to exercise the standard of care required by
	this Article.
31	(g) Nothing in this Article shall be construed to create an
32	obligation on the part of a managed care entity to provide to an
33	insured or enrollee a health care service that is not covered
34	under its health benefit plan.
35	(h) A managed care entity may not enter into a contract with a
36	health care provider, or with an employer or employer group
37	purchasing organization, that includes an indemnification or hold
	harmless clause for the acts or conduct of the managed care
	entity. Any such indemnification or hold harmless clause is void
40	and unenforceable to the extent of the restriction.

l (i) A managed care entity shall not remove a physician or
2 health care provider from its plan or refuse to renew the
3 physician or health care provider with its plan for advocating on
4 behalf of an enrollee for appropriate and medically necessary
5 health care for the enrollee.
6 "§ 90-21.53. No liability under this Article on the part of an
7 employer or employer group purchasing organization that purchases
8 coverage or assumes risk on behalf of its employees or a
9 physician or health care provider.
10 (a) This Article does not create any liability on the part of
11 an employer or employer group purchasing organization that
12 purchases a health benefit plan or assumes risk on behalf of its
13 employees.
14 (b) This Article does not create any liability on the part of
15 an employer of an enrollee or insured or that employer's
16 employees, unless the employer is the enrollee's or insured's
17 managed care entity and makes coverage determinations under a
18 managed care plan. This Article does not create any liability on
19 the part of an employee organization, a voluntary employee
20 beneficiary organization, or a similar organization, unless such
21 organization is the enrollee's or insured's managed care entity
22 and makes coverage determinations under a managed care plan.
23 (c) This Article does not create any liability on the part of
24 a physician or health care provider in addition to that otherwise
25 imposed under existing law. No managed care entity held liable
26 under this Article shall be entitled to contribution under
27 Chapter 1B of the General Statutes from a physician or health
28 care provider.
29 "90-21.54. Separate trial required.
30 Upon motion of any party in an action that includes a claim
31 brought pursuant to this Article involving a managed care entity.
32 the court shall order separate discovery and a separate trial of
33 any claim, cross-claim, counterclaim, or third-party claim
34 against any physician or other health care provider.
35 <u>§ 90-21.55</u> . Punitive damages.
36 An action brought under this Article is subject to the
37 provisions and limitations of Chapter 1D of the General Statutes
38 for recovery of punitive damages.
39 90-21.55. Exhaustion of administrative remedies and appeals.

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-	(a) Except as provided in this section, no action shall be
	2 commenced under this Article until the plaintiff has exhausted
3	3 all internal and external administrative remedies established
4	under Parts 2 and 4 of Article 50 of Chapter 58 of the General
5	Statutes.
6	(b) The plaintiff may file a claim without exhausting all
7	internal and external administrative remedies established under
8	Parts 2 and 4 of Article 50 of Chapter 58 of the General Statutes
-9	if the plaintiff proves the following to the court:
10	(1) Harm to the plaintiff has already occurred because
11	of the conduct of the managed care entity or
12	because of an act or omission of an employee,
13	
14	managed care entity for whose conduct the managed
15	care entity is liable.
16	(2) The administrative review would not be beneficial
17	to the plaintiff.
18	(c) This Article does not prohibit a plaintiff from pursuing
19	other appropriate remedies for relief.
20	Section 2. G.S. 1A-1, Rule 42, reads as rewritten:
21	"Rule 42. Consolidation; separate trials.
22	(a) Consolidation When Except as provided in subdivision
23	(b)(2) of this section, when actions involving a common question
24	of law or fact are pending in one division of the court, the
25	judge may order a joint hearing or trial of any or all the
26	matters in issue in the actions; he may order all the actions
27	consolidated; and he may make such orders concerning proceedings
28	therein as may tend to avoid unnecessary costs or delay. When
29	actions involving a common question of law or fact are pending in
30	both the superior and the district court of the same county, a
31	judge of the superior court in which the action is pending may
32	order all the actions consolidated, and he may make such orders
	concerning proceedings therein as may tend to avoid unnecessary
34	costs or delay.
35	(b) Separate trials
36	(1) The court may in furtherance of convenience or to
37	avoid prejudice and shall for considerations of
38	venue upon timely motion order a separate trial of
39	any claim, crossclaim, cross-claim, counterclaim,
40	or third-party claim, or of any separate issue or

1		of any number of claims, crossclaims, cross-claims,
2		counterclaims, third-party claims, or issues.
3	<u>(2)</u>	Upon motion of any party in an action that includes
4		a claim commenced under Article 1G of Chapter 90 of
5		the General Statutes involving a managed care
6		entity as defined in G.S. 90-21.50, the court shall
7		the court shall
1		order separate discovery and a separate trial of
8		any claim, cross-claim, counterclaim, or third-
9		
-		party claim against a physician or other medical
10		provider."
11		
	Secu	on 3. This act becomes effective July 1, 2001,
12	and applies to	causes of action arising on and after that date.





Bill Summary HEALTH CARE LIABILITY

BILL ANALYSIS

Committee:	LRC Committee/Managed Care	Introduced by:		
Date:	April 27, 2000	Summary by:	Linda Attarian	
Version:	FINAL DRAFT		Committee Counsel	

SUMMARY: The Act would amend Chapter 90 of the North Carolina General Statutes, by adding a new article to establish a standard of care for managed care entities which administer, deliver, arrange for, provide for, or reimburse for health care services or assume the risk for the delivery of health care services and to provide for recovery for violations of that standard. The act prohibits the shifting or delegation of liability for the acts or conduct of managed care entities and ensures that certain other liability is not created. The act would become effective July 1, 2001.

CURRENT LAW:

Medical Malpractice: Article 1B of Chapter 90 establishes a standard of health care and a cause of action for individuals who have been harmed as a result of receiving or failing to receive health care services meeting that standard from a health care provider. The statute defines 'medical malpractice action' to mean " a civil action for damages for personal injury or death arising out of the furnishing or failure to furnish professional services in the performance of medical, dental, or other health care by a health care provider". The standard of health care is defined as the performance of health care practice "in accordance with the standards of practice among members of the same health care profession with similar training and experience situated in the same or similar communities at the time of the alleged act giving rise to the cause of action.

Corporate Practice of Medicine: The current law specifically states that health maintenance organizations and provider sponsored organizations are not health care providers and are not subject to the above medical malpractice standards. However, individual providers employed by or under contract with an HMO or PSO are subject to the medical malpractice standards.

BILL ANALYSIS:

1. Applicability:

The proposed legislation applies to *health care treatment decisions* made by *managed care entities*. A managed care entity is defined as an *insurer* that delivers, administers, or undertakes to provide for, arrange for, or reimburse for health care services, or assumes the risk for the delivery of health care services. An insurer is an entity that writes a health benefit plan and is or should be licensed under the provisions of Chapter 58, either as an insurance company, a service corporation, health maintenance organization, or a multiple employer welfare arrangement.

Managed care entities are defined in the act to specifically <u>exclude</u>: (1) employers who purchase coverage for or on behalf of their employees and (2) *health care providers*.

Page 2

A "*health care provider*" is defined to include (1) individuals who are licensed health care providers under Chapter 90 of the General Statutes and (2) health care facilities licensed under Chapters 131E (hospitals) and 122C (mental health facilities and hospitals) and their agents and employees, including any officers and directors of health care facilities.

The act defines a *health care treatment decision* as a decision that determines which and to what extent health care services will be provided and reimbursed under the plan. The decision must actually affect the quality of the diagnosis, care, or treatment provided under the plan to the enrollee or the insured.

The act applies to *health benefit plans* provided by entities that are or should be regulated under Chapter 58. The act does not apply to self-funded health plans regulated under ERISA.

2. Standard of Care/Liability:

Definition of ordinary care. The act places a duty upon the managed care entity to exercise ordinary care when making health care decisions. 'Ordinary care' is defined to mean that degree of care that a managed care entity of ordinary prudence would use under the same or similar circumstances. For a person who is an agent or employee of a managed care entity, the standard of care is that degree of care that a person of ordinary prudence in the same profession, specialty, or area of practice as the person would use in the same or similar circumstances.

Scope of duty: The act imposes liability upon a managed care entity for damages for harm to an insured or enrollee proximately caused by its failure to exercise ordinary care in making health care treatment decisions. In addition the managed care entity is also liable for damages proximately caused by the failure of its agents, employees, and representatives (over whom it has the *right* to exercise influence and control), to exercise ordinary care in making health care treatment decisions.

Liability of employer: An employer or other plan sponsor, or an employee of the employer or sponsor, will only be liable under the act for damages proximately caused by the failure to exercise ordinary care in making a health care treatment decision when the action is based on the employer, sponsor, or employee's exercise of authority to make a health care treatment decision.

Liability of health care provider or "physicians": The act does not place any liability on a physician or health care provider in addition to medical malpractice liability under current law. The act defines "physician" to include: (1) a licensed (NC) medical doctor; (2) a professional association or corporation comprising medical doctors; or 3) a person or entity wholly owned by medical doctors.

3. Defenses to Liability:

The managed care entity has a defense to an action brought under this act if (1) neither its employee, agent, or representative participated or had any influence or control over the health care treatment decision and (2) the managed care entity did not deny or delay payment for a recommended or prescribed treatment or health care service.

A managed care entity may not use as a defense in a action brought under this act any laws that prohibit the practice of medicine by a corporate entity or by an HMO.

4. Indemnification and hold-harmless clauses:

The act would make void and unenforceable any indemnification or hold-harmless clauses for the acts or conduct of the managed care entity.

5. Exhaustion of administrative remedies and appeals:

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A person would not be allowed to bring an action under the act unless they first sought an external review of the health care treatment decision as allowed under Part 4 of Article 50 of Chapter 58 of the General Statutes (See proposed recommendation entitled "Independent External Review."). This requirement would not apply upon a good faith showing that harm to the insured or enrollee has already occurred because of the conduct of the managed care entity or because of an act or omission of an employee, agent, ostensible agent, or representative of the managed care entity for whose conduct it is liable; or the review would not be beneficial to the insured or enrollee

6. Punitive Damages:

Actions brought under the act would be subject to the limitations and restrictions placed on punitive damage awards under Chapter 1D of the General Statutes. For example, the award for punitive damages may not exceed three times the compensatory damages or \$250,000, whichever is greater.

7. Separate Trial:

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In an action involving a managed care entity brought pursuant to this act, on motion of any party the court must order a separate trial of any claim, cross claim, counterclaim, or third party claim against any physician (includes physician -owned entities) or other health care provider.

North Carolina General Assembly

Research Division, 733-2578

8. Effective Date: The act becomes effective July 1, 2001.

APPENDIX D

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SESSION 1999

RM-017 THIS IS A DRAFT 25-APR-00 08:57:21

Short Title: Prompt Pay.

(Public)

D

Sponsors:

1

Referred to:

A BILL TO BE ENTITLED

2 AN ACT TO PROVIDE FOR THE PROMPT PAYMENT OF CLAIMS UNDER HEALTH 3 BENEFIT PLANS AND TO MAKE CONFORMING AMENDMENTS TO RELATED CLAIM 4 PAYMENT LAWS. 5 The General Assembly of North Carolina enacts: Section. 1. Article 3 of Chapter 58 of the General 6 7 Statutes is amended by adding a new section to read: "§ 58-3-225. Prompt claim payments under health benefit plans. 8 9 (a) As used in this section: 'Health benefit plan' means an accident and health 10 (1) 11

12hospital or medical service corporation contract; a13health maintenance organization subscriber14contract; a plan provided by a multiple employer15welfare arrangement; or a plan provided by another16benefit arrangement, to the extent permitted by the17Employee Retirement Income Security Act of 1974, as18amended, or by any waiver of or other exception to19that Act provided under federal law or regulation.20"Health benefit plan" does not mean any plan	11	insurance policy or certificate; a nonprofit
health maintenance organization subscriber contract; a plan provided by a multiple employer welfare arrangement; or a plan provided by another benefit arrangement, to the extent permitted by the Employee Retirement Income Security Act of 1974, as amended, or by any waiver of or other exception to that Act provided under federal law or regulation.	12	hospital or medical service corporation contract: a
14contract; a plan provided by a multiple employer15welfare arrangement; or a plan provided by another16benefit arrangement, to the extent permitted by the17Employee Retirement Income Security Act of 1974, as18amended, or by any waiver of or other exception to19that Act provided under federal law or regulation.	13	health maintenance organization subscriber
15welfare arrangement; or a plan provided by another16benefit arrangement; to the extent permitted by the17Employee Retirement Income Security Act of 1974, as18amended, or by any waiver of or other exception to19that Act provided under federal law or regulation.	14	contract; a plan provided by a multiple employer
16benefit arrangement, to the extent permitted by the17Employee Retirement Income Security Act of 1974, as18amended, or by any waiver of or other exception to19that Act provided under federal law or regulation.	15	welfare arrangement; or a plan provided by another
17Employee Retirement Income Security Act of 1974, as18amended, or by any waiver of or other exception to19that Act provided under federal law or regulation.	16	benefit arrangement, to the extent permitted by the
18amended, or by any waiver of or other exception to19that Act provided under federal law or regulation.	17	Employee Retirement Income Security Act of 1974, as
19 that Act provided under federal law or regulation.	18	amended, or by any waiver of or other exception to
20 "Health benefit plan" does not mean any plan	19	that Act provided under federal law or regulation.
	20	"Health benefit plan" does not mean any plan

1		implemented or administered by the North Carolina
2		or United States Department of Health and Human
3		Services, or any successor agency, or its
4		representatives. "Health benefit plan" also does
5		not mean any of the following kinds of insurance:
6		a. Credit.
7		b. Disability income.
8		c. Coverage issued as a supplement to liability
9		insurance.
10	х •	d. Hospital income or indemnity.
11		e. Insurance under which benefits are payable
12		with or without regard to fault and that is
13		statutorily required to be contained in any
14		liability policy or equivalent self-insurance.
15		f. Medical payments under motor vehicle or
16		homeowners' insurance policies.
17		g. Short-term limited duration health insurance
18		policies as defined in Part 144 of Title 45 of
19		the Code of Federal Regulations.
20		h. Workers' compensation.
21	<u>(2)</u>	
22		facility that is responsible for directly making
23		the claim with an insurer, an insured, or an
24		insured's legal representative.
25	_(3)	'Health care facility' means a facility that is
26		licensed under Chapter 131E or 122C of the General
27		Statutes in which health care services are provided
28		to patients.
29	<u>(4)</u>	'Health care provider' means an individual who is
30		licensed, certified, or otherwise authorized under
31		Chapter 90 of the General Statutes to provide
32		health care services in the ordinary course of
33		business or practice of a profession or in an
34		approved education or training program.
35	(5)	'Insurer' includes an insurance company subject to
36		this Chapter, a service corporation organized under
37		Article 65 of this Chapter, a health maintenance
38		organization organized under Article 67 of this
39		Chapter, or a multiple employer welfare arrangement

1 <u>subject to Article 49 of this Chapter, that write</u>
2 a health benefit plan.
3 (b) An insurer shall, within 30 days after receipt of a claim
4 send by electronic or paper mail to the claimant:
5 (1) payment of the claim,
6 (2) notice of denial of the claim,
7 (3) notice that the proof of loss is inadequate of
8 incomplete, or
9 (4) notice that the claim is not submitted on the form
10 required by the health benefit plan, by the
11 contract between the insurer and health care
12 provider or health care facility, or by applicable
13 law.
14 (c) If the claim is denied, the notice shall include the
15 specific reason or reasons for the denial. If the claim is
16 contested or cannot be paid because the proof of loss is
17 inadequate or incomplete, the notice shall contain the specific
18 reason or reasons why the claim has not been paid and an
19 itemization or description of all of the information needed by
20 the insurer to complete the processing of the claim. If a claim
21 is denied or contested in part, the insurer shall pay the
22 undisputed portion of the claim within 30 days after receipt of
23 the claim and send the notice of the denial or contested status
24 within 30 days after receipt of the claim. If a claim is
25 contested or cannot be paid because the claim was not submitted
26 on the required form, the notice shall contain the required form
27 and instructions to complete that form. Upon receipt of
28 additional information requested in its notice to the claimant,
29 the insurer shall continue processing the claim and pay or deny
30 the claim within 30 days after receiving the additional
31 information.
32 (d) If an insurer requests additional information under
33 subsection (c) of this section and the insurer does not receive
34 the additional information within 90 days after the request was
35 made, the insurer shall deny the claim and send the notice of
36 denial to the claimant in accordance with subsection (c) of this
37 section. The insurer shall include the specific reason or reasons 38 for depial in the notice including the fact that is f
38 for denial in the notice, including the fact that information 39 that was requested was not provided. The insurer shall inform
40 the claimant in the notice that the claim will be
40 the claimant in the notice that the claim will be re-opened if

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1 the information previously requested is submitted to the insurer 2 within one year after the date of the denial notice closing the 3 claim. (e) Health benefit plan claim payments that are not made in 4 5 accordance with this section shall bear interest at the rate of 6 18 percent (18%) per year, beginning on the date on which the 7 claim should have been paid. A payment is considered made on the 8 date upon which a check, draft, or other valid negotiable 9 instrument is placed in the United States Postal Service in a 10 properly addressed, postpaid envelope, or, if not mailed, on the 11 date of the electronic transfer or other delivery of the payment This subsection does not apply to claims for 12 to the claimant. 13 benefits that are not covered by the health benefit plan; nor 14 does this subsection apply to deductibles, co-payments, or other 15 amounts for which the insurer is not liable. (f) Insurers may require that claims be submitted within 180 16 17 days after the date of the provision of care to the patient by 18 the health care provider and, in the case of health care provider 19 facility claims, within 180 days after the date of the patient's 20 discharge from the facility. Failure to submit a claim within the 21 time required does not invalidate or reduce any claim if it was 22 not reasonably possible for the insured or the insured's legal 23 representative to file the claim within that time, provided that 24 the claim is submitted as soon as reasonably possible and in no 25 event, except in the absence of legal capacity of the insured, 26 later than one year from the time submittal of the claim is 27 otherwise required. (q) If a claim for which the claimant is a health care 28 29 provider or health care facility has not been paid within 60 days 30 after receipt of the initial claim, the insurer shall send a 31 claim status report to the insured. The report shall indicate 32 that the claim is under review and the insurer is communicating 33 with the health care provider or health care facility to resolve 34 the matter. While a claim remains unresolved, the insurer shall 35 send a claim status report to the insured every 30 days after the 36 previous report was sent. Any retroactive reductions of payments or demands for 37 (h) 38 refund of previous overpayments that are because retroactive 39 review-of-coverage decisions or payment levels shall be 40 reconciled for specific claims unless the insurer and health care

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1 provider or health care facility agree to other reconciliation Any retroactive demands by health care 2 methods and terms. 3 providers or health care facilities for payment because of 4 underpayments or nonpayments for covered services shall be 5 reconciled for specific claims unless the insurer and health care 6 provider or health care facility agree to other reconciliation 7 methods and terms. The period for which retroactive adjustments 8 may be made may be specified in the contract between the insurer 9 and health care provider or health care facility. (i) As used in this subsection, 'copayment or deductible' 10 11 means the portion of a charge for services covered by a health 12 benefit plan that, under the plan's terms, it is the obligation 13 of the insured to pay. No health care provider or health care 14 facility shall directly or indirectly seek payment or collection 15 of the claim, other than a copayment or deductible, from an 16 insured or an insured's legal representative while the claim is 17 being resolved under this section. No health care provider or 18 health care facility shall report an insured or an insured's 19 legal representative to any credit reporting agency while the

20 <u>claim is being resolved under this section. A violation of this</u> 21 <u>subsection by a health care provider or health care facility is a</u> 22 <u>violation of Article 2 of Chapter 75 of the General Statutes.</u>

(j) Every insurer shall maintain records of its activities under this section, including records of when each claim was, paid, denied, or pended, and the insurer's review and handling of each claim under this section, as well as documentation sufficient to demonstrate compliance with this section. The information to be included in these records and the maintenance of these records by the insurer, including electronic reproduction and storage, shall be governed by rules adopted by 1 the Commissioner.

32 (k) A violation of this section by an insurer subjects the 33 insurer to the sanctions in G.S. 58-2-70."

34 (1) An insurer is not in violation of this section nor subject 35 to interest payments under this section if its failure to comply 36 with this section is caused in material party by (i) the person 37 submitting the claim, or (ii) by matters beyond the insurer's 38 reasonable control, including an act of God, insurrection, 39 strike, fire, or power outages.

Section 2. G.S. 58-3-100(c) reads as rewritten:

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"(c) The Commissioner may impose a civil penalty under G.S. 1 2 58-2-70 if an HMO, service corporation, MEWA, or insurer fails to 3 acknowledge a claim within 30 days after receiving written notice 4 of the claim, but only if the notice contains sufficient 5 information for the insurer to identify the specific coverage Acknowledgement of the claim shall be made to the 6 involved. 7 claimant or his legal representative advising that the claim is 8 being investigated; or shall be a payment of the claim; or shall 9 be a bona fide written offer of settlement; or shall be a written 10 denial of the claim. A claimant includes an insured, a health 11 care provider, or a health care facility that is responsible for 12 directly making the claim with an insurer. This subsection does 13 not apply to insurers subject to G.S. 58-3-225." Section 3. G.S. 58-51-15(a)(7) reads as rewritten: 14 "(7) A provision in the substance of the following language: 15 PROOFS OF LOSS: Written proof of loss must be furnished to the 16 17 insurer at its said office in the case of a claim for loss for 18 which this policy provides any periodic payment contingent upon 19 continuing loss within 90 180 days after the termination of the 20 period for which the insurer is liable and in case of a claim for 21 any other loss within 90 180 days after the date of such loss. 22 Failure to furnish such proof within the time required shall not 23 invalidate nor reduce any claim if it was not reasonably possible 24 to give proof within such time, provided such proof is furnished 25 as soon as reasonably possible and in no event, except in the 26 absence of legal capacity, capacity of the insured, later than 27 one year from the time proof is otherwise required." Section. 4. If any section or provision of this act is 28 29 declared unconstitutional or invalid by the courts, it does not 30 affect the validity of the act as a whole or any part other than 31 the part so declared to be unconstitutional or invalid.

32 Section. 5. This act becomes effective July 1, 2001, 33 and applies to claims or services rendered on or after July 1, 34 2001.



Bill Summary PROMPT PAYMENT

BILL ANALYSIS

Committee:	LRC/Managed Care
Date:	April 27, 2000
Version:	FINAL DRAFT

Introduced by: Summary by: Linda Attarian Committee Counsel

SUMMARY: Section 1 of the act adds a new section to Article 3 to require a licensed insurer to pay an complete and uncontested claim submitted by a claimant within 30 days. If the claim is not paid within 30 days, interest at 18 percent will be added to the claim. The act requires the insurer to notify the claimant by email or in writing within the same 30 days if the claim is contested or denied. If a claim is denied or contested in part, the insurer must pay the undisputed portion of the claim within the same 30 days. The denial or contest notice is required to include the specific reasons supporting the denial or contest and an itemized list of any additional information required for the insurer to complete the processing for the claim. The insurer must pay the claim within 30 days after receiving the additional information. A violation of the act would subject the insurer to civil penalties, restitution or license suspension or revocation by the Commissioner of Insurance pursuant to G.S. 58-2-70.

Sections 2 and 3 make conforming changes to existing law. Section 4 is a severability clause. The bill would become effective on July 1, 2001.

CURRENT LAW: G.S. 58-3-100 authorizes the Commissioner of Insurance to impose a civil penalty if an insurer fails to acknowledge a claim within 30 days after receiving notice of the claim, but only if the notice contains sufficient information for the insurer to identify the specific coverage involved. Acknowledgment of the claim shall be made to the claimant or his legal representative by advising that the claim is being investigated; or shall be a payment of the claim; or shall be a bona fide written offer of a settlement; or shall be a written denial of the claim.

Uniform claim forms: G.S. 58-3-171 and rules adopted by the Commissioner require all claims submitted by institutional health care providers to health benefit plans to be submitted on the HCFA 1450 (UB 92), or a substantively similar claim form, and all claims submitted by noninstitutional health care providers to be submitted on the HCFA 1500, or a substantively similar claim form. Payors and health care providers that receive or generate claims or send payments by electronic means must accept or generate the appropriate ASC X12 Standard Format for their health care claims submission and remittance transactions. Additional information beyond that contained on the uniform form or format may be collected, but must meet certain requirements set by the Commissioner.

Current Remedies for claim settlement practices in violation of Chapter 58: G.S. 58-63-15(11) defines certain claim settlement practices which, if committed with sufficient frequency to indicate a general business practice, will constitute an unfair and deceptive act or practice in the practice of insurance. Allegations of such practice patterns are subject to investigation by the Commissioner, who may file charges and issue a cease and desist order. Violations of the cease and desist order will subject the insurer to a penalty (in addition to any other applicable penalties) of not less than \$1,000, but not more than \$5,000 for each violation. No private right of action is created under the Article. However, unfair and Page 2

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\$5,000 for each violation. No private right of action is created under the Article. However, unfair and deceptive acts in the insurance area are not regulated exclusively by Article 63, but are also actionable under § 75-1.1, which provides for a private right of action.

G.S. 58-2-70 authorizes the Commissioner of Insurance to seek appropriate remedies from any person who violates any provision of Chapter 58. The Commissioner has the authority to impose fines, petition the court to order appropriate restitution and suspend or revoke the violator's license.

BILL ANALYSIS: Section 1 of the bill amends Article 63 of Chapter 58 pertaining to the regulation of unfair trade practices in the business of insurance. The act adds a new section to Article 63 to require a licensed insurer to pay a clean claim submitted by a claimant for covered services within 30 days. A claimant includes a health care provider or facility, an insured or an insured's legal representative. Within 30 days of receipt of the claim, the insurer must send the claimant, by paper mail or electronic mail one of the following:

- 1. Payment of the claim.
- 2. A notice of denial.
- 3. A notice that the proof of loss is either inadequate or incomplete.
- 4. A notice that the claim was not submitted on the form required by the health plan or the contract between the provider or facility and the insurer.

Notice requirements: The notice of denial must include the reasons for the denial. The notice that the proof of loss is either inadequate or incomplete or that the claimed benefit or benefits are not covered under the plan must include the reasons why the claim has not been paid along with an itemization or description of the information needed to process the claim. If the claim is not on the form required by the health benefit plan or the contract between the health care provider or facility, the notice shall include the required forms and complete instructions as to the format to be used. If the claim is denied in part, the insurer must pay the undisputed portion and send notice of the denial or contest within 30 days of receiving the claim.

Time frame for payment after receiving the requested information: The insurer has 30 days in which to pay the claim after it receives the requested information. If the requested information is not received within 90 days of making the request, the insurer shall deny the claim, and must send the claimant notice of the denial. If the insurer receives the requested information within one year after the date of the notice of denial, the insurer must reopen the claim.

Interest accrual: Claims that are not processed according to the time frames discussed above will bear interest at 18 percent per year. The interest will begin on the date the claim should have been paid.

Timeframe within which to submit a claim: The act allows insurers to require claimants to submit claims within 180 calendar days of the last date the insured's health care provider provides health care services to the insured or from the date the insured is discharged from a health care facility. Failure to submit a claim within 180 days due to a reasonable impossibility does not invalidate or reduce any claim, provided that the claim is submitted as soon as reasonably possible. In all cases where the claimant has legal capacity, the claim must be submitted within 365 days of the time submittal would have otherwise been required.

Informing the insured on the status of a disputed claim: In cases where the claim is submitted by the insured's health care provider or facility, the insurer must send a claim status report to the insured if the

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Reconciling retroactive demands for overpayments and underpayments: Demands by an insurer for refunds for overpayments or demands by providers for additional payment because of underpayments or nonpayments for covered services must be reconciled for specific claims unless the parties agree on a different way to handle the movement of the money. This means that the party owed money must identify the specific service(s) and claim payment(s) that were involved in the error. Both parties would have to reflect the adjustment in the account/claims experience of the insured in question. In cases where a plan contracts with a provider, the parties could agree on a different way to handle the allocation of the refund or additional payment. Regardless of how the partites reconcile the transaction, the insurer must ensure that the patient's claims experience is adjusted to reflect the ultimate adjustment.

Payment/collection and credit protection: No provider or facility may report an insured or their legal representative to a credit reporting agency while a claim is in dispute. Further, no provider or facility may seek payment (other than copayment or deductible) from the insured while the claim in dispute. A violation will subject the provider or facility to the application of Article 2, Chapter 75 (Prohibited Acts by Debt Collectors). This act provides a private right of action, and authorizes the court to impose civil penalties up to \$2,000 for each violation.

Record keeping: Insurers must maintain records and other documentation to demonstrate compliance with the act according to adopted by the Commissioner. Such records include documentation of how the insurer reviewed and handled each claim, including the date it was paid, denied, or pended.

Violations of the act: A single violation of the act would subject the insurer to the sanctions provided for under G.S. 58-2-70.

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Sections 2 and 3 make conforming changes to existing law.

Section 4 is a severability clause.

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The act becomes effective on July 1, 2001, and applies to services rendered on or after that date.

APPENDIX E

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SESSION 1999

RM-016 THIS IS A DRAFT 25-APR-00 09:05:44

Short Title: HMO Insolvency.

(Public)

Sponsors:

1

Referred to:

A BILL TO BE ENTITLED

2 AN ACT TO PROTECT PERSONS ENROLLED IN AN [·] HMO FROM THE 3 CONSEQUENCES OF INSOLVENCY THE OF THAT HMO BY AUTHORIZING 4 ASSESSMENTS OF REMAINING HMOS IN THE STATE TO PAY FOR UNCOVERED 5 EXPENDITURES OF AND CONTINUATION OF COVERAGE FOR THE ENROLLEES. 6 The General Assembly of North Carolina enacts:

7 Section. 1. Article 67 of Chapter 58 of the General 8 Statutes is amended by adding a new section to read:

9 "§ 58-67-126. Insolvency protection; assessment.

10 (a) When an HMO in this State is declared insolvent by a court 11 of competent jurisdiction, the Commissioner may levy an 12 assessment on solvent HMOs doing business in this State to pay 13 claims for uncovered expenditures for enrollees who are residents 14 of this State and to provide continuation of coverage for 15 enrollees not covered under G.S. 58-67-120, G.S. 58-67-125, or 16 G.S. 58-67-130. Assessments against an HMO may not exceed two 17 percent (2%) of that HMO's average annual premiums received in 18 North Carolina on policies and contracts during the three 19 calendar years immediately preceding the year in which the 20 insolvent HMO was declared insolvent.

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(b) To provide the funds necessary to carry out the powers and 1 2 duties of the Commissioner under this section, the Commissioner 3 shall assess and notify in writing the HMOs at such time and for 4 such amounts, as the Commissioner finds necessary. Assessments 5 not paid within 30 days of the written notice shall accrue 6 interest at the rate of one percent (1%) per month, or any part made until necessary 7 thereof. Assessments shall not be to this the of section. 8 implement purposes Computation of 9 assessments under this section shall be made with a reasonable 10 degree of accuracy, recognizing that exact determinations may not 11 always be possible. 12 The Commissioner may use funds obtained under subsection (C) 13 (a) of this section to pay claims for uncovered expenditures for 14 enrollees of an insolvent HMO who are residents of this State, 15 provide for continuation of coverage for enrollees who are 16 residents of this State and are not covered under G.S. 58-67-120, 17 G.S. 58-67-125, or G.S. 58-67-130, and administrative costs. The 18 Commissioner may by rule prescribe the time, manner, and form for 19 filing claims under this section or may require claims to be 20 allowed by an ancillary receiver or the domestic liquidator or 21 receiver. A receiver or liquidator of an insolvent HMO shall claim the proceeding in 22 allow a in an amount equal to uncovered expenditures 23 administrative and paid under this 24 section. 25 (d) Any person receiving benefits under this section for 26 uncovered expenditures is deemed to have assigned the rights 27 under the covered health care plan certificates to the 28 Commissioner to the extent of the benefits received. The 29 Commissioner may require an assignment to it of such rights by 30 any payee, enrollee, or beneficiary as a condition precedent to 31 the receipt of any rights or benefits conferred by this section 32 upon that person. The Commissioner is subrogated to these rights 33 against the assets of an insolvent HMO held by a receiver or 34 liquidator of another jurisdiction. The assignment of subrogation rights of the Commissioner 35 (e) 36 and allowed claim under this section have the same priority 37 against the assets of the insolvent HMO as those possessed by the 38 person entitled to receive benefits under this section or for 39 similar expenses in the receivership or liquidation.

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1	(f) When assessed funds are unused following the completion of
2	the liquidation of a HMO, the Commissioner will distribute on a
3	pro rata basis any unused amounts received under subsection (a)
4	of this section to the HMOs that have been assessed under this
5	section.
6	(g) The aggregate coverage of uncovered expenditures under
7	this section shall not exceed \$300,000 with respect to one
8	individual. Continuation of coverage for an enrollee shall
9	continue for the duration of the contract period for which
10	premiums have been paid and continuation of coverage for an
11	enrollee who is confined in an inpatient facility shall continue
12	until his or her discharge or expiration of benefits. The
13	Commissioner may provide continuation of coverage on any
14	reasonable basis; including, continuation of the HMO contract or
15	substitution of indemnity coverage in a form determined by the
16	Commissioner.
17	
	the assessment of an HMO if, in the Commissioner's opinion,
19	
20	
	HMO is abated or deferred, in whole or in part, the amount by
	which the assessment is abated or deferred may be assessed
	against the other HMOs in a manner consistent with the basis for
	assessments set forth in this section. An HMO that fails to pay
	an assessment within 30 days after notice is subject to a civil
	penalty of not more than \$1,000 per day, or suspension or
	revocation of its license, or both.
28	(i) It is proper for any HMO, in determining its premium rates
	and policy owner dividends, to consider the amount reasonably
	necessary to meet its assessment obligations under this section."
31	Section 2. G.S. 58-30-220(2) reads as rewritten:
32	"(2) Claims or portions of claims for benefits under policies
	and for losses incurred, including claims of third parties under
	liability policies; claims of HMO enrollees and HMO enrollees'
	beneficiaries; <u>beneficiaries</u> , including situations where an
	enrollee or beneficiary is liable to a health care provider for
	services provided under the HMO plan; claims for unearned
	premiums; claims for funds or consideration held under funding
	agreements, as defined in G.S. 58-7-16; claims under life
40	insurance and annuity policies, whether for death proceeds,

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1 annuity proceeds, or investment values; and claims of domestic 2 and foreign guaranty associations, including claims for the 3 reasonable administrative expenses of domestic and foreign 4 guaranty associations; but excluding claims of insurance pools, 5 underwriting associations, or those arising out of reinsurance 6 agreements, claims of other insurers for subrogation, and claims 7 of insurers for payments and settlements under uninsured and 8 underinsured motorist coverages."

Section. 3. If any section or provision of this act is 9 10 declared unconstitutional or invalid by the courts, it does not 11 affect the validity of the act as a whole or any part other than 12 the part so declared to be unconstitutional or invalid. 13

Section 4. This act becomes effective January 1, 2001.





Bill Summary HMO INSOLVENCY

BILL ANALYSIS

Committee:	LRC/Managed Care Issues	Introduced by:	
Date:	April 27, 2000	Summary by:	Linda Attarian
Version:	FINAL DRAFT		Committee Counsel

SUMMARY: The act would amend Article 67 of Chapter 58 pertaining to the regulation of health maintenance organizations (HMOs) to provide a mechanism with which the Commissioner of Insurance may ensure that uncovered claims against an insolvent HMO are covered and health care coverage for enrollees is continued.

CURRENT LAW:

Warning of financial instability: § 58-67-105 authorizes the Commissioner of Insurance to order an HMO that is experiencing financial instability to take reasonable actions to rectify the situation. The Commissioner is authorized to adopt rules to set uniform standards and criteria for the early warning that the continued operation of any HMO might be hazardous to its enrollees, creditors, or the general public.

Net worth: § 58-67-110 requires each full service HMO to maintain a minimum net worth of not less than one million dollars (\$1,000,000). This amount is increased annually, depending on the amount of the HMO's contingency reserves. The statute further requires every full service HMO to have and maintain at all times an adequate plan, acceptable to the Commissioner, for protection against insolvency.

Insolvency protection: § 58-67-115 provides that unless the HMO maintains a special deposit or has adequate insurance or a guaranty arrangement, each contract between every HMO and a participating provider must include a hold harmless clause to ensure that in the event the HMO fails to pay for health care services, the subscriber or enrollee will not be liable to the provider for any amount the HMO is unable to pay the provider. The special deposit must be in cash or cash equivalent and is calculated according to specific circumstances. In all cases, the deposit is controlled by and administered by the Commissioner. If the HMO has a guaranty arrangement, it must be approved in writing by the Commissioner.

Continuation of benefits: § 58-67-120 provides that each HMO must have a plan for handling insolvency and for the continuation of benefits. The HMO must ensure that its enrollees continue to receive benefits for the duration of the contract period for which premiums have been paid, or if the enrollee is confined in an inpatient facility until the discharge or expiration of benefits. The Commissioner may require the HMO

- to acquire insurance to cover the expenses to be paid for benefits after an insolvency, or
- to contribute to an insolvency reserve, or
- to submit letters of credit.

Assignment of coverage: § 58-67-125 authorizes the Commissioner, in the event of an insolvency of an HMO, to order all other carriers that participated in the enrollment process with the insolvent HMO to offer enrollees of the insolvent HMO a 30-day enrollment period from the date of insolvency. The coverage and rates must be the same that the carrier had offered to the enrollees of the group at its last

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regular enrollment period. If no other HMOs participated in the enrollment process with the insolvent HMO or if the Commissioner determines that the other health benefit plan or plans lack sufficient health care delivery resources to assure that health care services will be available and accessible to all of the group enrollees of the insolvent HMO, then the Commissioner must allocate the insolvent HMO's group and nongroup enrollees among all other HMOs that operate within a portion of the insolvent HMO's service area.

Replacement coverage safeguards: § 58-67-130 provides that any insurer that has contracted with an HMO to provide replacement coverage within a period of 60 days from the date of discontinuance of the HMO contract or policy must immediately cover all enrollees who were validly covered under the previous HMO, regardless of any provisions of the contract relating to active employment or hospital confinement or pregnancy. Further, the contract for replacement coverage must provide for full benefits for conditions that proceeded the effective date of the succeeding insurer's contract.

Guaranty Associations: HMOs are not required to belong to guaranty associations. Guaranty associations cover claims against insolvent insurance companies by assessing the member companies an amount necessary to cover the claims. These associations can be found in Articles 48 (property and casualty) and 62 (life and health) of GS Chapter 58 and in Article 3 of GS Chapter 97.

BILL ANALYSIS: Section 1 of the act does not establish an HMO guaranty association, but instead authorizes the Commissioner to assess, as necessary, other HMOs for the unpaid obligations of the insolvent HMO. The Commissioner may waive or abate the assessment if the Commissioner determines the assessment would endanger the HMO's ability to continue coverage. The total of all assessments against a solvent HMO in one calendar year may not exceed two percent (2%) of the solvent HMO's average annual premiums during the three (3) calendar years preceding the year the insolvent HMO was declared insolvent. If an HMO fails to pay the assessment within 30 days of notice, it may be fined up to \$1,000 per day and have its license suspended or revoked. In addition, assessments will accrue interest at the rate of one percent (1%) per month. Assessments must be reasonably accurate and necessary to protect persons enrolled in an HMO from the consequences of insolvency.

The act authorizes the Commissioner to arrange for continuation of coverage of enrollees of an insolvent HMO. The Commissioner would be authorized to require other HMOs or other indemnity insurers to provide continuation coverage. The continuation coverage could not exceed \$300,000 for any one individual and would continue for the duration of the contract period for which premiums have been paid. Coverage of enrollees confined to inpatient facilities would continue until discharge.

Section 2 of the act clarifies the priority under Article 30 of Chapter 58 of claims for benefits and for losses incurred as a result of HMO insolvency. Medical care claims owed by HMO enrollees, including situations where an enrollee is liable to health care providers for services provided under a HMO plan would be paid like other claims under insurance and HMO contracts.

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Section 3 provides a severability clause.

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Section 4 provides that the act will become effective January 1, 2001.

APPENDIX F

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SESSION 1999

RM-020 THIS IS A DRAFT 27-APR-00 16:38:08

Short Title: External Review/Managed Care.

Sponsors:

Referred to:

1 2 A BILL TO BE ENTITLED 3 AN ACT TO PROVIDE STANDARDS FOR THE ESTABLISHMENT AND MAINTENANCE 4 OF EXTERNAL REVIEW PROCEDURES IN HEALTH INSURANCE AND MANAGED 5 CARE TO ASSURE THAT COVERED PERSONS HAVE THE OPPORTUNITY FOR AN 6 INDEPENDENT REVIEW OF A HEALTH BENEFIT PLAN COVERAGE DECISION 7 MADE BY THE INSURER OR MANAGED CARE PLAN; AND TO MAKE CONFORMING 8 AMENDMENTS TO EXISTING LAWS ON UTILAZTION REVIEW AND GRIEVANCES. 9 10 The General Assembly of North Carolina enacts: 11 12 Section 1. The title of Article 50 of Chapter 58 of the 13 General Statutes reads as rewritten: "ARTICLE 50. 14 15 General Accident and Health Insurance Regulations." 16 Section 2. Article 50 of Chapter 58 of the General 17 Statutes is divided into five Parts as follows: 18 Part 1. Miscellaneous Provisions Comprising G.S. 58-19 50-1 through G.S. 58-50-45 20 Part 2. Utilization Review, PPOs, and Grievances 21 Comprising G.S. 58-50-50 through G.S. 58-50-64

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(Public)

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4	Comprising G.S. 58-50-75 through G.S. 58-50-
5	95, as enacted in Section @ of this act.
6	Part 5. Small Employer Group Health Insurance Reform
7	Act Comprising G.S. 58-50-100 through G.S.
8	58-50-156.
9	Section 3. G.S. 58-50-151 is recodified as G.S. 58-51-
	116.
11	
	reads as rewritten:
	"(a) Definitions As used in this section and section, in
	G.S. 58-50-62, and in Part 4 of this Article, the term:"
15	-
16	Statutes is amended by adding a new Part to read:
17	<u>"PART 4.</u>
18	"Health Benefit Plan External Review.
19	"§ 58-50-75. Purpose, scope, and definitions.
	(a) The purpose of this Part is to provide standards for the
	establishment and maintenance of external review procedures to
	assure that covered persons have the opportunity for an
	independent review of an appeal decision upholding a
	noncertification or a second level grievance review decision
	upholding a noncertification, as defined in this Part.
	(b) This Part applies to all persons that provide or perform
	utilization review. With respect to second level grievance
	review decisions, this part applies only to second level
	grievance review decisions involving noncertification decisions.
30	
	in this Part:
32	(1) 'Covered benefits' or 'benefits' means those
33	benefits consisting of medical care, provided
34	directly through insurance or otherwise and
35	including items and services paid for as medical
36	care, under the terms of a health benefit plan.
37	(2) 'Disclose' means to release, transfer or otherwise
38	divulge protected health information to any person
39	other than the individual who is the subject of the
40	protected health information.

1	<u>(3)</u>	'Health information' means information or data,
2		whether oral or recorded in any form or medium, and
3		personal facts or information about events or
4		relationships that relates to: the past, present or
5		future physical, mental, or behavioral health or
6		condition of an individual or a member of the
7		individual's family; the provision of health care
8		services to an individual; or payment for the
9		provision of health care services to an individual.
10	(4)	'Independent review organization' or 'organization'
11		means an entity that conducts independent external
12		reviews of appeals of noncertifications and second
13		level grievance review decisions.
14	(5)	'Protected health information' means health
15		information that identifies an individual who is
16		the subject of the information; or with respect to
17		which there is a reasonable basis to believe that
18		the information could be used to identify an
19		individual.
20	"§ 58-50-77.	Notice of right to external review.
21		rer shall notify the covered person in writing of
22	the covered	person's right to request an external review and
23		ppropriate statements and information set forth in
24	this section	at the time the insurer sends written notice of a
25	decision on a	second-level grievance review in which the insurer
26	upheld its or	iginal noncertification as set forth in G.S. 58-50-
27	62.	
28	(b) The in:	surer shall include in the notice required under
29) of this section for a notice related to an appeal
30		r G.S. 58-50-61, a statement informing the covered
	person that:	
32	(1)	If the covered person has a medical condition where
33	· · ·	the timeframe for completion of an expedited review
34		of a grievance involving an appeal decision under
35		G.S. 58-50-61 would seriously jeopardize the life
36		or health of the covered person or would jeopardize
3.7		the covered person's ability to regain maximum
38		function, the covered person may file a request for
39		an expedited external review under G.S. 58-50-82 at
40	•	the same time the covered person files a request

1		for an expedited review of a grievance involving an
2		appeal decision under G.S. 58-50-61 and 58-50-62,
3		but that the organization assigned to conduct the
4		expedited external review will determine whether
5		the covered person shall be required to complete
6		the expedited review of the grievance before
- 7	1	conducting the expedited external review.
8	(2)	The covered person may file a grievance under the
9		insurer's internal grievance process under G.S. 58-
10		50-61 and 58-50-62, but if the insurer has not
11		issued a written decision to the covered person
12		within 45 days after the date the covered person
13		files the grievance with the insurer and the
14		covered person has not requested or agreed to a
15		delay, the covered person may file a request for
16		external review under G.S. 58-50-80 of this section
17		and shall be considered to have exhausted the
18		insurer's internal grievance process for purposes
19		of G.S. 58-50-79.
20	(c) The i	nsurer shall include in the notice required under
21) of this section for a notice related to a final
22		grievance review decision under G.S. 58-50-62, a
23		orming the covered person that:
24	(1)	If the covered person has a medical condition where
25		the timeframe for completion of a standard external
26		review under G.S. 58-50-80 would seriously
27		jeopardize the life or health of the covered person
28		or would jeopardize the covered person's ability to
29		regain maximum function, the covered person may
30		file a request for an expedited external review
31		under G.S. 58-50-82; or
32	(2)	If the second-level grievance review decision
33		concerns an admission, availability of care,
34		continued stay or health care service for which the
35		covered person received emergency services, but has
36		not been discharged from a facility, the covered
37.		person may request an expedited external review
38		under G.S. 58-50-82.
39	<u>(d) In ac</u>	dition to the information to be provided under
40	<u>subsections (</u>	b) and (c) of this section, the insurer shall

1	include a copy of the description of both the standard and
2	expedited external review procedures the insurer is required to
3	provide under G.S. 58-50-93, including the provisions in the
4	external review procedures that give the covered person the
	opportunity to submit additional information.
	(e) An insurer that has collected protected health information
7	under a valid authorization under this Part may use and disclose
8	the protected health information to a person acting on behalf of
9	or at the direction of the insurer for the performance of the
10	insurer's insurance functions: claims administration, claims
11	adjustment and management, fraud investigation, underwriting,
12	loss control, rate-making functions, reinsurance, risk
13	management, case management, disease management, quality
14	assessment, quality improvement, provider credentialing
15	verification, utilization review, peer review activities,
16	grievance procedures, policyholder service functions, and
17	internal administration of compliance, managerial, and
18	information systems. Additional insurance functions may be
19	allowed for the purpose of this subsection with the prior
20	approval of the Commissioner. The protected health information
21	shall not be used or disclosed for any purpose other than in the
22	performance of the insurer's insurance functions.
23	(f) Except for a request for an expedited external review
	under G.S. 58-50-82, all requests for external review shall be
25	made in writing to the Commissioner.
26	"§ 58-50-79. Exhaustion of internal grievance process.
27	(a) Except as provided in subsections (d) through (g) of this
28	section, a request for an external review under G.S. 58-50-80 or
29	G.S. 58-50-82 shall not be made until the covered person has
	exhausted the insurer's internal grievance process under G.S. 58-
31	50-61 and G.S. 58-50-62.
32	(b) A covered person shall be considered to have exhausted the
33	insurer's internal grievance process for purposes of this
34	
35	(1) Has filed a second level grievance involving a
36	noncertification appeal decision under G.S. 58-50-
37	<u>61 and 58-50-62.</u>
38	(2) Except to the extent the covered person requested
39	or agreed to a delay, has not received a written
40	decision on the grievance from the insurer within

1	45 days since the date the covered person filed the
2	grievance with the insurer.
3	(c) Notwithstanding subsection (b) of this section, a covered
4	person may not make a request for an external review of a
5	noncertification involving a retrospective review determination
6	made under G.S. 58-50-61 until the covered person has exhausted
7	the insurer's internal grievance process.
8	(d) At the same time a covered person files a request for an
9	expedited review of an appeal involving a noncertification as set
10	forth in G.S. 58-50-61(1), the covered person may file a request
11	the second secon
12	G.S. 58-50-82 if the covered person has a medical condition where
13	the timeframe for completion of an expedited review of the appeal
14	involving a noncertification set forth in G.S. 58-50-61(j) would
15	seriously jeopardize the life or health of the covered person or
16	would jeopardize the covered person's ability to regain maximum
17	function. An insurer may waive its right to conduct an expedited
18	review of an appeal and allow the covered person to proceed with
19	an expedited external review of the noncertification.
20	(e) Upon receipt of a request for an expedited external review
21	under subsection (d) of this section, the organization conducting
22	the external review in accordance with the provisions of G.S. 58-
23	50-82 shall immediately determine whether the covered person
24	shall be required to complete the expedited review process set
25	forth in G.S. 58-50-61(j) before it conducts the expedited
26	external review, unless the insurer has waived its right to
27	
28	(f) Upon a determination made under subsection (e) of this
29	section that the covered person must first complete the expedited
	appeal process under G.S. 58-50-61(j), the organization
	immediately shall notify the covered person and the insurer of
	this determination and that it will not proceed with the
33	expedited external review under G.S. 58-50-82 until completion of
34	the expedited appeal process and the covered person's grievance
35	at the completion of the expedited appeal process remains
	unresolved.
37	(g) A request for an external review of a noncertification may
38	be made before the covered person has exhausted the insurer's
39	internal grievance procedures under G.S. 58-50-61 and G.S. 58-50-

-	contraction the income correct to waive the exhaustion
	62 whenever the insurer agrees to waive the exhaustion
	requirement. (h) If the requirement to exhaust the insurer's internal
3	(h) If the requirement to exhaust the insurer's internal grievance procedures is waived under subsection (g) of this
4	section, the covered person may file a request in writing for a
5	section, the covered person may file a request in writing for a
	standard external review as set forth in G.S. 58-50-80.
	<pre>"§ 58-50-80. Standard external review. (a) Within 60 days after the date of receipt of a notice of a</pre>
8	noncertification appeal decision or a second level grievance
	review decision under G.S. 58-50-77, a covered person may file a
10	review decision under G.S. 58-50-77, a covered person may rice a
11	request for an external review with the Commissioner. (b) Upon receipt of a request for an external review under
12	subsection (a) of this section, the Commissioner immediately
13	shall notify and send a copy of the request to the insurer that
	made the decision which is the subject of the request. The
15	
16	insurer shall immediately submit to the Commissioner the information required for the preliminary review under subsection
17	
18	<pre>(c) of this section. (c) Within five business days after the date of receipt of a</pre>
19	request for an external review, the Commissioner shall complete a
20	preliminary review of the request to determine whether:
21	The second part of the second part of the
22 23	(1) The individual is or was a covered person in the health care health care
23 24	service was requested or, in the case of a
24 25	retrospective review, was a covered person in the
25 26	health benefit plan at the time the health care
20	service was provided.
28	
20 29	(2) The health care service that is the subject of the noncertification appeal decision or the second
30	level grievance review decision upholding a
31	noncertification reasonably appears to be a covered
32	service under the covered person's health benefit
33	plan.
34	(3) The covered person has exhausted the insurer's
35	internal grievance process under G.S.58-50-62(i)
36	unless the covered person is not required to
-37	exhaust the insurer's internal grievance process
38	under G.S. 58-50-79.
39	(4) The covered person has provided all the information
40	and forms required by the Commissioner that are
40	

1	necessary to process an external review, including		
2	the authorization form provided under G.S. 58-50-		
3	<u>77(e).</u>		
4	(d) Upon completion of the preliminary review under subsection		
5	(c) of this section, the Commissioner immediately shall notify		
6	the covered person in writing whether the request is complete and		
· 7	whether the request has been accepted for external review.		
8	(e) If the request is accepted for external review, the		
9	Commissioner shall:		
10	(1) Include in the notice provided under subsection (d)		
11	of this section a statement that the covered person		
12	may submit to the Commissioner in writing within		
13	seven days after the date of the notice additional		
14	information and supporting documentation that the		
15	organization shall consider when conducting the		
16	external review.		
17	(2) Immediately notify the insurer in writing of the		
18	acceptance of the request for external review.		
19	(3) Provide the covered person and the covered person's		
20	provider with a list of organizations approved		
21	under G.S. 58-50-85.		
22	(4) Inform the covered person that the covered person		
23	has the right to select the organization of his or		
24	her choice and notify the Commissioner within five		
25	days after receipt of the notice, and that if the		
26	covered person does not select an organization and		
27	inform the Commissioner of the selection within		
28	five days after receipt of the notice, the		
29	Commissioner will assign an organization to conduct		
30	the external review.		
31	(f) If the request is not complete, the Commissioner shall		
32	request from the covered person the information or materials		
33	needed to make the request complete. The covered person shall		
34	furnish the Commissioner with the requested information or		
35	materials within 90 days after the date of the insurer's decision		
36	for which external review is requested. If the request is not		
37	accepted for external review, the Commissioner shall inform the		
38	covered person and the insurer in writing of the reasons for its		
39	nonacceptance.		

1	(a) If the incurred door not colort an organization of his or			
1				
2	her choice and notify the Commissioner of the selection within			
3				
4	subsection (e) of this section, the Commissioner shall			
5	systematically assign an appropriate independent review			
6				
7				
	organization is not bound by any decisions or conclusions reached			
	during the insurer's utilization review process or the insurer's			
	internal grievance process under G.S. 58-50-61 and 58-50-62.			
11				
12	provided under subsection (e) of this section, the insurer or its			
	designee utilization review organization shall provide to the			
	assigned organization, the documents and any information			
15	considered in making the noncertification appeal decision or the			
	second level grievance review decision. Except as provided in			
	subsection (i) of this section, failure by the insurer or its			
	designee utilization review organization to provide the documents			
	and information within the time specified in this subsection			
20	shall not delay the conduct of the external review.			
21				
22				
	organization may terminate the external review and make a			
	decision to reverse the noncertification appeal decision or the			
26				
27	the decision under this subsection, the organization shall notify			
28	the covered person, the insurer, and the Commissioner.			
29				
	information and documents received under subsections (h) and (i)			
31	of this section and any other information submitted in writing by			
32	the covered person under subsection (e) of this section that has			
33	been forwarded to the organization by the Commissioner. Upon			
34	receipt of any information submitted by the covered person under			
35	subsection (e) of this section, at the same time the Commissioner			
36	forwards the information to the organization, the Commissioner			
37	shall forward the information to the insurer.			
38	(k) Upon receipt of the information required to be forwarded			
39				
40	its noncertification appeal decision or second level grievance			

		on that is the subject of the external review.
		on by the insurer of its noncertification appeal
3	decision or s	second level grievance review decision under this
4	subsection sh	all not delay or terminate the external review. The
5	external revi	ew shall be terminated if the insurer decides, upon
6	<u>completion</u>	of its reconsideration, to reverse its
7		ion appeal decision or second level grievance review
		provide coverage or payment for the requested health
9	care service	that is the subject of the noncertification appeal
10		econd level grievance review decision.
11	(1) Immed	iately upon making the decision to reverse its
12		ion appeal decision or second level grievance review
13		r subsection (k) of this section, the insurer shall
14	notify the cov	vered person, the organization, and the Commissioner
15	in writing of	its decision. The organization shall terminate the
16	external revie	ew upon receipt of the notice from the insurer sent
	under this sul	
18		dition to the documents and information provided
19	under subsect	tions (h) and (i) of this section, the assigned
20	organization,	to the extent the documents or information are
21		the organization considers them appropriate, shall
		following in reaching a decision:
23	(1)	The covered person's medical records.
24	(2)	The attending health care provider's
25		recommendation.
26	(3)	Consulting reports from appropriate health care
27		providers and other documents submitted by the
28		insurer, covered person, or the covered person's
29		treating provider.
30	(4)	The terms of coverage under the covered person's
31		health benefit plan with the insurer to ensure that
32		the organization's decision shall not be contrary
33		to the terms of coverage under the covered person's
34		health benefit plan with the insurer.
35	(5)	The most appropriate practice guidelines, which may
36		include generally accepted practice guidelines,
37		evidence-based practice guidelines, or any other
38		practice guidelines developed by the federal
39		government, national or professional medical

1		societies, boards and associations. Local practice
2		guidelines may be used when appropriate.
3	(6)	Any applicable clinical review criteria developed
4		and used by the insurer or its designee utilization
5		review organization.
6	(7)	
7	(n) With	
8		of the request for external review, the assigned
9		shall provide written notice of its decision to
10	uphold or rev	erse the noncertification appeal decision or second
11	level grieva	nce review decision to the covered person, the
12	insurer, and	the Commissioner.
13	(o) The or	ganization shall include in the notice sent under
14	subsection (n) of this section:
15	(1)	A general description of the reason for the request
16		for external review.
17	(2)	The date the organization received the assignment
18		from the Commissioner to conduct the external
19		review.
20	<u>(3)</u>	The date the organization received information and
21		documents submitted by the covered person and by
22		the insurer.
23	<u>(4)</u>	The date the external review was conducted.
24	<u>(5)</u>	The date of its decision.
25	<u>(6)</u>	The principal reason or reasons for its decision.
26	<u>(7)</u>	The clinical rationale for its decision.
27	(8)	References to the evidence or documentation,
28		including the practice guidelines, considered in
29		reaching its decision.
30	<u>(9)</u>	The professional qualifications and licensure of
31		the clinical peer reviewers.
32	(10)	Notice to the covered person that he or she is not
33		liable for the cost of the external review.
34	(p) Upon r	eceipt of a notice of a decision under subsection
35	(n) of this	section reversing the noncertification appeal
36	decision or s	econd level grievance review decision, the insurer
		hall approve the coverage that was the subject of
		ication appeal decision or second level grievance
	review decisio	
40	"§ 58-50-82.	Expedited external review.

1	(a) Excep	t as provided in subsection (g) of this section, a
2		on may make a request for an expedited external
3	review with	the Commissioner at the time the covered person
4	receives:	
5	(1)	An appeal decision upholding a noncertification if:
6		a. The noncertification appeal decision involves
7		a medical condition of the covered person for
8		which the timeframe for completion of an
9		expedited second level grievance review of a
10		noncertification set forth in G.S. 58-50-62(1)
11		would seriously jeopardize the life or health
12		of the covered person or would jeopardize the
13		covered person's ability to regain maximum
14		function; and
15		b. The covered person has filed a request for an
16		expedited appeal of a noncertification as set
17		forth in G.S. 58-50-61(1); or
18	<u>(2)</u>	
19		a noncertification under G.S. 58-50-62(h) or (i):
20		a. If the covered person has a medical condition
21		where the timeframe for completion of a
22		standard external review under G.S. 58-50-80
23		would seriously jeopardize the life or health
24		of the covered person or would jeopardize the
25		covered person's ability to regain maximum
26		function; or
27		b. If the second level grievance concerns a
28		noncertification of an admission, availability
29		of care, continued stay, or health care
30		service for which the covered person received
31		emergency services, but has not been
32		discharged from a facility.
33		he time the Commissioner receives a request for an
		ternal review, the Commissioner immediately shall:
35	<u>(1)</u>	Notify and provide a copy of the request to the
36		insurer that made the noncertification appeal
37		decision or second level grievance review decision
38	()	which is the subject of the request.
39	<u>(2)</u>	
40		meets the reviewability requirements set forth in

1		G.S. 58-50-80(c), assign an organization that has
2		been approved under G.S. 58-50-87. The organization
2		shall immediately determine whether the request
		should be reviewed on an expedited basis because
4		the timeframe for completion of a standard external
5		
6		review under G.S. 58-50-80 would seriously
7		jeopardize the life or health of the covered person
8		or would jeopardize the covered person's ability to
9		regain maximum function. The organization shall
10		then inform the covered person, insurer, and
11		Commissioner of its determination and conduct a
12		review and make a decision on the review within the
13		appropriate timeframe.
14	<u>(c) In read</u>	ching a decision, the assigned organization is not
15	bound by any	decisions or conclusions reached during the
		lization review process or internal grievance
17	process under	G.S. 58-50-61 and 58-50-62.
18	(d) At tl	he time the insurer receives the notice under
19	subsection (b) of this section, the insurer or its designee
20	utilization re	eview organization shall immediately provide or
21	transmit all	necessary documents and information considered in
22	making the f	inal noncertification decision to the assigned
23	organization e	electronically or by telephone or facsimile or any
		e expeditious method.
25	(e) In add:	ition to the documents and information provided or
	transmitted un	nder subsection (d) of this section, the assigned
27	organization.	to the extent the information or documents are
28	available and	the organization considers them appropriate, shall
		ollowing in reaching a decision:
30		The covered person's pertinent medical records.
31		The attending health care provider's
32		recommendation.
33		Consulting reports from appropriate health care
34		providers and other documents submitted by the
34 35		insurer, covered person, or the covered person's
36		treating provider.
30		The terms of coverage under the covered person's
37	<u>(4)</u>	health benefit plan with the insurer to ensure that
		the organization's decision shall not be contrary
39		the organization's decision shall not be contrary

1		to the terms of coverage under the covered person's
2		health benefit plan with the insurer.
3	(5)	The most appropriate practice guidelines, which may
4		include generally accepted practice guidelines,
5		evidence-based practice guidelines, or any other
6		practice guidelines developed by the federal
7		government, national or professional medical
8		societies, boards and associations. Local practice
9		guidelines may be used when appropriate.
10	(6)	Any applicable clinical review criteria developed
11		and used by the insurer or its designee utilization
12		review organization in making noncertification
13		decisions.
14	(7)	Medical necessity, as defined in G.S. 58-3-200(b).
15	(f) As expe	editiously as the covered person's medical condition
16	or circumstan	ces require, but not more than four days after the
17	date of recei	ot of the request for an expedited external review,
18	the assigned	organization shall make a decision to uphold or
19	reverse the	noncertification appeal decision or second level
		iew decision and notify the covered person, the
21		the Commissioner of the decision.
22	<u>(g) If t</u> l	ne notice provided under subsection (f) of this
23	<u>section was n</u>	ot in writing, within two days after the date of
24	providing tha	t notice, the assigned organization shall provide
		rmation of the decision to the covered person, the
26	insurer, and	the Commissioner; and include the information set
27	forth in G.S.	58-50-80(o). Upon receipt of the notice a decision
28	under subsec	ction (f) of this section reversing the
29		ion appeal decision or second level grievance review
		insurer immediately shall approve the coverage that
31		ct of the noncertification.
32		pedited external review may not be provided for
33		noncertifications.
34	<u>"§ 58-50-84.</u>	Binding nature of external review decision.
35		ernal review decision is binding on the insurer.
36		ternal review decision is binding on the covered
37		to the extent the covered person has other remedies
38		er applicable federal or state law.
39		ered person may not file a subsequent request for
40	<u>external rev</u>	iew involving the same noncertification appeal

1	decision or second level grievance review decision for which the		
	covered person has already received an external review decision		
3	under this Part.		
4	"§ 58-50-85. Approval of independent review organizations.		
5	(a) The Commissioner shall approve independent review		
6			
7	under this Part to ensure that an organization satisfies the		
8	minimum qualifications established under G.S. 58-50-87. The		
9	Commissioner shall develop an application form for initially		
10	approving and for re-approving organizations to conduct external		
11	reviews.		
12	(b) Any organization wishing to be approved to conduct		
	external reviews under this Part shall submit the application		
	form and include with the form all documentation and information		
	necessary for the Commissioner to determine if the organization		
16	satisfies the minimum qualifications established under G.S. 58-		
17	50-87.		
18	(c) The Commissioner may, in his discretion, determine that		
	accreditation by a nationally recognized private accrediting		
	entity with established and maintained standards for independent		
	review organizations that meet the minimum qualifications		
	established under G.S. 58-50-87 will cause an independent review		
	organization to be deemed to have met, in whole or in part, the		
	requirements of this section and G.S. 58-50-87. A decision by		
	the Commissioner to recognize an accreditation program for the		
	purpose of granting deemed status may be made only after		
	reviewing the accreditation standards and program information		
	submitted by the accrediting body. An independent review		
	organization seeking deemed status due to its accreditation shall		
	submit original documentation issued by the accrediting body to		
	demonstrate its accreditation.		
32	(d) The Commissioner may charge an application fee that		
	independent review organizations shall submit to the Commissioner		
	with an application for approval and re-approval.		
35	(e) An approval is effective for two years, unless the		
	Commissioner determines before expiration of the approval that		
	the independent review organization is not satisfying the minimum		
38	qualifications established under G.S. 58-50-87.		
39	(f) Whenever the Commissioner determines that an independent		
40	review organization no longer satisfies the minimum requirements		

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1	established un	nder (G.S. 58-50-87, the Commissioner shall terminate
2	the approval	of t	he independent review organization and remove
3	the independe	nt re	view organization from the list of independent
4	<u>review organi</u>	zatio	ns approved to conduct external reviews under
5	this Part that	t is r	maintained by the Commissioner under subsection
6	(g) of this se	ectio	n.
7	(g) The Co	mmiss	ioner shall maintain and periodically update a
8	list of approv	ved in	ndependent review organizations.
9	"§ 58-50-87.	Mi	nimum qualifications for independent review
10	<u>organizations</u>	<u>•</u>	
11	<u>(a) Asac</u>	ondit	ion of approval under G.S. 58-50-85 to conduct
12	<u>external revi</u>	ews,	an independent review organization shall have
13	and maintain	writ	ten policies and procedures that govern all
14	aspects of b	<u>oth</u> t	the standard external review process and the
15	expedited ext	ernal	review process set forth in G.S. 58-50-80 and
16	G.S. 58-50-82	that	include, at a minimum:
17	<u>(1)</u>	<u>A q</u>	uality assurance mechanism in place that
18		ensu	res:
19		<u>a.</u>	That external reviews are conducted within the
20			specified time frames and required notices are
21			provided in a timely manner.
22		b.	The selection of qualified and impartial
23			clinical peer reviewers to conduct external
24			reviews on behalf of the independent review
25			organization and suitable matching of
26			reviewers to specific cases.
27		с.	The confidentiality of medical and treatment
28			records and clinical review criteria.
29		<u>d.</u>	That any person employed by or under contract
30		· .	with the independent review organization
31			adheres to the requirements of this Part.
32	(2)	А	toll-free telephone service to receive
33		info	rmation on a 24-hour-day, seven-day-a-week
34		basi	s related to external reviews that is capable
35		of a	ccepting, recording, or providing appropriate
36		inst	ruction to incoming telephone callers during
37		othe	r than normal business hours.
38	(3)	Agre	e to maintain and provide to the Commissioner
39	<u>_</u>	the	information set out in G.S. 58-50-90.
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1		(4)	A program for credentialing clinical peer
2			reviewers.
3		(5)	Agree to contractual terms or written requirements
4			established by the Commissioner regarding the
5			procedures for handling a review.
6			linical peer reviewers assigned by an independent
7	review or	rgani	zation to conduct external reviews shall be medical
8	doctors d	or ot	ther appropriate health care providers who meet the
. 9	following	y min	imum qualifications:
10		<u>(1)</u>	Be an expert in the treatment of the covered
11			person's injury, illness, or medical condition that
12			is the subject of the external review;
13		(2)	Be knowledgeable about the recommended health care
14			service or treatment through recent or current
15			actual clinical experience treating patients with
16			the same or similar injury, illness, or medical
17			condition of the covered person;
18		<u>(3)</u>	If the covered person's treating provider is a
19			medical doctor, hold a non-restricted license from
20	•		the North Carolina Medical Board and, if a
21			specialist medical doctor, a current certification
22			by a recognized American medical specialty board in
23			the area or areas appropriate to the subject of the
24			external review;
25		(4)	If the covered person's treating provider is not a
26			medical doctor, hold a non-restricted North
27			Carolina license, registration, or certification in
28			the same allied health occupation as the covered
29			person's treating provider; and
30		(5)	<u>Have no history of disciplinary actions or</u>
31			sanctions, including loss of staff privileges or
32			participation restrictions, that have been taken or
33			are pending by any hospital, governmental agency or
34			unit, or regulatory body that raise a substantial
35			question as to the clinical peer reviewer's
36			physical, mental, or professional competence or
37			moral character.
38			lition to the requirements set forth in subsection
39			section, an independent review organization may not
40	own or c	ontro	ol, be a subsidiary of or in any way be owned or

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1	controlled by, or exercise control with a health benefit plan, a
	national, state or local trade association of health benefit
3	plans, or a national, state or local trade association of health
4	care providers.
5	(d) In addition to the requirements set forth in subsections
6	(a), (b), and (c) of this section, to be approved under G.S. 58-
7	50-85 to conduct an external review of a specified case, neither
8	the independent review organization selected to conduct the
9	external review nor any clinical peer reviewer assigned by the
10	independent organization to conduct the external review may have
11	a material professional, familial, or financial conflict of
12	interest with any of the following:
13	(1) The insurer that is the subject of the external
14	review.
15	(2) The covered person whose treatment is the subject
16	of the external review or the covered person's
17	authorized representative.
18	
19	the insurer that is the subject of the external
20	review.
21	(4) The health care provider, the health care
21 22	provider's medical group or independent practice
22 23	provider's medical group or independent practice association recommending the health care service or
22 23 24	provider's medical group or independent practice
22 23 24 25	provider's medical group or independent practice association recommending the health care service or treatment that is the subject of the external review.
22 23 24 25 26	provider's medical group or independent practice association recommending the health care service or treatment that is the subject of the external review. (5) The facility at which the recommended health care
22 23 24 25 26 27	<pre>provider's medical group or independent practice association recommending the health care service or treatment that is the subject of the external review. (5) The facility at which the recommended health care service or treatment would be provided.</pre>
22 23 24 25 26 27 28	<pre>provider's medical group or independent practice association recommending the health care service or treatment that is the subject of the external review. (5) The facility at which the recommended health care service or treatment would be provided. (6) The developer or manufacturer of the principal</pre>
22 23 24 25 26 27 28 29	<pre>provider's medical group or independent practice association recommending the health care service or treatment that is the subject of the external review. (5) The facility at which the recommended health care service or treatment would be provided. (6) The developer or manufacturer of the principal drug, device, procedure, or other therapy being</pre>
22 23 24 25 26 27 28 29 30	<pre>provider's medical group or independent practice association recommending the health care service or treatment that is the subject of the external review. (5) The facility at which the recommended health care service or treatment would be provided. (6) The developer or manufacturer of the principal drug, device, procedure, or other therapy being recommended for the covered person whose treatment</pre>
22 23 24 25 26 27 28 29 30 31	<pre>provider's medical group or independent practice association recommending the health care service or treatment that is the subject of the external review. (5) The facility at which the recommended health care service or treatment would be provided. (6) The developer or manufacturer of the principal drug, device, procedure, or other therapy being recommended for the covered person whose treatment is the subject of the external review.</pre>
22 23 24 25 26 27 28 29 30 31 32	 provider's medical group or independent practice association recommending the health care service or treatment that is the subject of the external review. (5) The facility at which the recommended health care service or treatment would be provided. (6) The developer or manufacturer of the principal drug, device, procedure, or other therapy being recommended for the covered person whose treatment is the subject of the external review. (e) In determining whether an independent review organization
22 23 24 25 26 27 28 29 30 31 32 33	<pre>provider's medical group or independent practice association recommending the health care service or treatment that is the subject of the external review. (5) The facility at which the recommended health care service or treatment would be provided. (6) The developer or manufacturer of the principal drug, device, procedure, or other therapy being recommended for the covered person whose treatment is the subject of the external review. (e) In determining whether an independent review organization or a clinical peer reviewer of the independent review</pre>
22 23 24 25 26 27 28 29 30 31 32 33 34	<pre>provider's medical group or independent practice association recommending the health care service or treatment that is the subject of the external review. (5) The facility at which the recommended health care service or treatment would be provided. (6) The developer or manufacturer of the principal drug, device, procedure, or other therapy being recommended for the covered person whose treatment is the subject of the external review. (e) In determining whether an independent review organization or a clinical peer reviewer of the independent review organization has a material professional, familial, or financial</pre>
22 23 24 25 26 27 28 29 30 31 32 33 34 35	<pre>provider's medical group or independent practice association recommending the health care service or treatment that is the subject of the external review. (5) The facility at which the recommended health care service or treatment would be provided. (6) The developer or manufacturer of the principal drug, device, procedure, or other therapy being recommended for the covered person whose treatment is the subject of the external review. (e) In determining whether an independent review organization or a clinical peer reviewer of the independent review organization has a material professional, familial, or financial conflict of interest for purposes of subsection (d) of this</pre>
22 23 24 25 26 27 28 29 30 31 32 33 34 35 36	<pre>provider's medical group or independent practice association recommending the health care service or treatment that is the subject of the external review. (5) The facility at which the recommended health care service or treatment would be provided. (6) The developer or manufacturer of the principal drug, device, procedure, or other therapy being recommended for the covered person whose treatment is the subject of the external review. (e) In determining whether an independent review organization or a clinical peer reviewer of the independent review organization has a material professional, familial, or financial conflict of interest for purposes of subsection (d) of this section, the Commissioner shall take into consideration</pre>
 22 23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 	<pre>provider's medical group or independent practice association recommending the health care service or treatment that is the subject of the external review. (5) The facility at which the recommended health care service or treatment would be provided. (6) The developer or manufacturer of the principal drug, device, procedure, or other therapy being recommended for the covered person whose treatment is the subject of the external review. (e) In determining whether an independent review organization or a clinical peer reviewer of the independent review organization has a material professional, familial, or financial conflict of interest for purposes of subsection (d) of this section, the Commissioner shall take into consideration situations where the independent review organization to be</pre>
 22 23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38 	<pre>provider's medical group or independent practice association recommending the health care service or treatment that is the subject of the external review. (5) The facility at which the recommended health care service or treatment would be provided. (6) The developer or manufacturer of the principal drug, device, procedure, or other therapy being recommended for the covered person whose treatment is the subject of the external review. (e) In determining whether an independent review organization or a clinical peer reviewer of the independent review organization has a material professional, familial, or financial conflict of interest for purposes of subsection (d) of this section, the Commissioner shall take into consideration situations where the independent review organization to be assigned to conduct an external review of a specified case or a</pre>
 22 23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38 39 	<pre>provider's medical group or independent practice association recommending the health care service or treatment that is the subject of the external review. (5) The facility at which the recommended health care service or treatment would be provided. (6) The developer or manufacturer of the principal drug, device, procedure, or other therapy being recommended for the covered person whose treatment is the subject of the external review. (e) In determining whether an independent review organization or a clinical peer reviewer of the independent review organization has a material professional, familial, or financial conflict of interest for purposes of subsection (d) of this section, the Commissioner shall take into consideration situations where the independent review organization to be</pre>

	may have an apparent professional, familial, or financial
2	relationship or connection with a person described in subsection
3	(d) of this section, but that the characteristics of that
4	relationship or connection are such that they are not a material
5	professional, familial, or financial conflict of interest that
6	results in the disapproval of the independent review organization
7	or the clinical peer reviewer from conducting the external
8	review.
9	"§ 58-50-89. Hold harmless for independent review organizations.
10	No independent review organization or clinical peer reviewer
	working on behalf of an organization shall be liable in damages
	to any person for any opinions rendered during or upon completion
	of an external review conducted under this Part, unless the
14	opinion was rendered in bad faith or involved gross negligence.
15	"§ 58-50-90. External review reporting requirements.
16	(a) An organization assigned under G.S. 58-50-80 or G.S. 58-
	50-82 to conduct an external review shall maintain written
	records in the aggregate and by insurer on all requests for
	external review for which it conducted an external review during
	a calendar year and submit a report to the Commissioner, as
21	required under subsection (b) of this section.
22	(b) Each organization required to maintain written records on
23	all requests for external review under subsection (a) of this
24	section for which it was assigned to conduct an external review
25	shall submit to the Commissioner, at least annually, a report in
26	the format specified by the Commissioner.
27	(c) The report shall include in the aggregate and for each
28	<u>insurer:</u>
29	(1) The total number of requests for external review.
30	(2) The number of requests for external review resolved
31	and, of those resolved, the number resolved
32	upholding the noncertification appeal decision or
33	second level grievance review decision and the
34	number resolved reversing the noncertification
35	appeal decision or second level grievance review
36	decision.
37	(3) The average length of time for resolution;
38	(4) A summary of the types of coverages or cases for
39	which an external review was sought, as provided in
40	the format required by the Commissioner;

1	(5) The number of external reviews under G.S. 58-50-
2	80(k) and (1) that were terminated as the result of
3	a reconsideration by the insurer of its
4	noncertification appeal decision or second level
5	grievance review decision after the receipt of
6	additional information from the covered person.
7	(6) Any other information the Commissioner may request
8	or require.
9	(d) The organization shall retain the written records required
10	under this section for at least three years.
11	(e) Each insurer shall maintain written records in the
12	aggregate and for each type of health benefit plan offered by the
13	
14	receives notice from the Commissioner under this Part. The
15	insurer shall retain the written records required under this
16	section for at least three years.
17	"§ 58-50-92. Funding of external review.
18	The insurer against which a request for a standard external
19	
20	
21	conducting the external review.
22	"§ 58-50-93. Disclosure requirements.
23	(a) Each insurer shall include a description of the external
24	review procedures in or attached to the policy, certificate,
25	
26	coverage it provides to covered persons.
27	(b) The description required under subsection (a) of this
28	section shall include a statement that informs the covered person
29	
30	external review of a noncertification appeal decision or a second
31	level grievance review decision upholding a noncertification with
32	the Commissioner. The statement shall include the telephone
33	number and address of the Commissioner.
34	(c) In addition to subsection (b) of this section, the
35	the set at inform the governed person that when filing a
55	statement shall inform the covered person that, when filing a
36	request for an external review, the covered person will be
36 37	request for an external review, the covered person will be required to authorize the release of any medical records of the
36 37 38	request for an external review, the covered person will be

SESSION 1999

1 "§ 58-50-94. Competitive selection of independent review 2 organizations. (a) The Commissioner shall prepare and publish requests for 3 4 proposals from independent review organizations that want to be 5 approved under G.S. 58-50-85. All proposals shall be sealed. The 6 Commissioner shall open all proposals in public. After the public opening, the Commissioner shall review 7 (b) 8 the proposals, examining the costs and quality of the services 9 offered by the independent review organizations, the reputation 10 and capabilities of the independent review organizations 11 submitting the proposals, and the provisions in G.S. 58-50-85 and 12 G.S. 58-50-87. The Commissioner shall determine which proposal 13 or proposals would satisfy the provisions of this Part. The 14 Commissioner shall make his determination in consultation with an 15 evaluation committee whose membership includes representatives of 16 insurers subject to Part 4 of Article 50, health care providers, 17 and insureds. In selecting the review organizations, in addition 18 to considering cost, quality, and adherence to the requirements 19 of the request for proposals the Commissioner shall consider the 20 desirability and feasibility of contracting with multiple review 21 organizations in order to allow insureds a choice of review 22 organizations, and shall ensure that at least one review 23 organization is available to and capable of reviewing cases 24 involving highly specialized services and treatments of any 25 nature. The Commissioner may reject any or all proposals. (c) An independent review organization may seek to modify or 26 27 withdraw a proposal only after the public opening and only on the 28 basis that the proposal contains an unintentional clerical error 29 as opposed to an error in judgment. An independent review 30 organization seeking to modify or withdraw a proposal shall 31 submit to the Commissioner a written request, with facts and 32 evidence in support of its position, before the determination 33 made by the Commissioner under subsection (b) of this section, 34 but not later than two days after the public opening of the 35 proposals. The Commissioner shall promptly review the request, 36 examine the nature of the error, and determine whether to permit 37 or deny the request. (d) The provisions of Article 3C of Chapter 143 of the General 38 39 Statutes do not apply to this Part. 40 "§ 58-50-95. Report by Commissioner.

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1	The Commissioner shall report semiannually to the Joint
2	Legislative Committee on Health Care Oversight regarding the
3	nature and appropriateness of reviews conducted under this Part.
4	The report should include the number of reviews, character of the
5	reviews, dollar amounts in question, and any other information
6	releant to the evaluation of the effectiveness of this Part."
7	Section 6. G.S. 58-50-61(a)(13) reads as rewritten:
8	"(13) 'Noncertification' means a determination by an insurer or
9	its designated utilization review organization that an admission,
10	availability of care, continued stay, or other health care
11	service has been reviewed and, based upon the information
12	provided, does not meet the insurer's requirements for medical
13	necessity, appropriateness, health care setting, level of care or
14	effectiveness, or does not meet the prudent layperson standard
15	for coverage of emergency services in G.S. 58-3-190, and the
16	requested service is therefore denied, reduced, or terminated. A
17	'noncertification' is not a decision rendered solely on the basis
18	that the health benefit plan does not provide benefits for the
19	health care service in question, if the exclusion of the specific
2.0	service requested is clearly stated in the certificate of
21	coverage. A 'noncertification' includes any situation in which
22	an insurer or its designated agent makes an evaluation or review
23	of medical information about a covered person's condition to
24	determine whether a requested treatment is experimental,
25	investigational, or cosmetic and the extent to which coverage
26	under the health benefit plan is affected by that decision."
27	Section 7. G.S. 58-50-61(a)(17)g. reads as rewritten:
28	"g. Retrospective review. Utilization review of medically
29	necessary services and supplies that is conducted after services
30	have been provided to a patient, but not the review of a claim
31	that is limited to an evaluation of reimbursement levels,
32	veracity of documentation, accuracy of coding, or adjudication
33	for payment. Retrospective review includes the review of claims
34	for emergency services to determine whether the prudent layperson
35	standard in G.S. 58-3-190 has been met."
36	Section 8. G.S. 58-50-61(i) reads as rewritten:
37	"(i) Requests for <u>Informal</u> Reconsideration. An insurer may
38	establish procedures for informal reconsideration of
39	noncertifications and if established, such procedures shall be in
40	writing. The reconsideration shall be conducted between the

		on's provider and a medical doctor licensed to
	-	icine in this State designated by the insurer.
3		er a written notice of noncertification has been
4		cordance with subsection (h) of this section. An
5	insurer shall	. not require a covered person to participate in an
6		onsideration before the covered person may appeal a
		ion under subsection (j) of this section. If, after
		nsideration the insurer upholds the noncertification
9	decision, the	insurer shall issue a new notice in accordance with
10	subsection (1	h) of this section. If the insurer is unable to
		formal reconsideration decision in fewer than 10
12	business day	vs, it shall treat the request for informal
		on as a request for an appeal, except that the
14	requirements	of subsection (k) of this section shall apply on or
15	before the 10	th business day after receipt of the request for an
16	informal reco	
17	Sect	ion 9. G.S. 58-50-62(a) is amended by adding a new
18	subsection to	
19		ormal Consideration of Grievances. If the insurer
20		cedures for informal considerations of grievances,
21	the procedure	s shall be in writing and the following requirements
22	apply:	
23	<u>(1)</u>	If the grievance concerns a clinical issue and the
24		informal consideration decision is not in favor of
25		the covered person, the insurer shall treat the
26		request as a request for a first-level grievance
27		review, except that the requirements of subdivision
28		(e)(1) of this section shall apply on the 10th
29		business day after receipt of the grievance; or
30	<u>(2)</u>	If the grievance concerns a non-clinical issue and
31		the informal consideration decision is not in favor
32		of the covered person, the insurer shall issue a
33		written decision that includes the information set
34		forth in G.S.58-50-62(c).
35	<u>(3)</u>	If the insurer is unable to render an informal
36		consideration decision within 10 business days of
.37		receipt of the grievance, the insurer shall treat
38		the request as a request for a first-level
39		grievance review, except that the requirements of
40		subdivision (e)(1) of this section shall apply on

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1	the 10th business day after receipt of the
2	grievance."
3	Section 10. G.S. 58-50-61(k)(5) reads as rewritten:
4	"(5) A statement advising the covered person of the covered
5	person's right to request a second-level grievance review and a
6	description of the procedure for submitting a second-level
7	grievance under G.S. 58-50-62. G.S. 58-50-62 if the insurer's
8	decision on the appeal is to uphold its noncertification."
9	Section 11. G.S. 58-50-62(e)(2)e. reads as rewritten:
10	"e. A statement advising the covered person of his or her
11	right to request a second-level grievance review and a
12	description of the procedure for submitting a second-level
13	grievance under this section. section if the insurer's decision
14	on the first level grievance review is not in favor of the
15	covered person."
16	Section 12. G.S. 58-50-62(h)(7) reads as rewritten:
17	"(7) A statement that the decision is the insurer's final
18	determination in the matter. In cases where the review concerned
19	a noncertification and the insurer's decision on the second-level
	review is to uphold its initial noncertification, a statement
	advising the covered person of his or her right to request an
22	external review and a description of the procedure for submitting
23	a request for external review to the Commissioner of Insurance."
24	Section 13. The Commissioner of Insurance shall report
	semiannually to the Joint Legislative Health Care Oversight
	Committee regarding the nature and appropriateness of reviews
	conducted under this Part. The report shall include the number
	of reviews, character of the reviews, dollar amounts in question,
29	and any other information relevant to the evaluation of the
30	effectiveness of the external review procedures establishes
31	pursuant to this act.
32	Section 14. If any section or provision of this act is
33	declared unconstitutional or invalid by the courts, it does not
34	affect the validity of the act as a whole or any part other than
35	the part so declared to be unconstitutional or invalid.
36	Section. 15. This act becomes effective July 1, 2001.



Bill Summary INDEPENDENT EXTERNAL REVIEW

BILL ANALYSIS

Committee:	LRC/Managed Care		
Date:	April 27, 2000		
Version:	FINAL DRAFT		

Introduced by: Summary by: Linda Attarian Committee Counsel

SUMMARY: The proposed legislation would add a new Part to Article 50 Chapter 58 of the General Statutes to establish an external, independent review process for consumers to obtain an external review of disputes regarding complaints and issues relating to the consumer's health benefit plan. A request for an external review would be made to the Commissioner of Insurance after exhausting all internal appeals. External reviews would be conducted by independent review organizations that are approved by the Commissioner of Insurance. The decision of the review organization must be made within 45 days or four days if necessary to avoid jeopardizing the health or life of the covered person. The decision would be binding upon the insurer. The insurer would pay the cost of the review. The act would become effective July 1, 2001, and would not be applicable to self-funded employer health plans regulated under ERISA.

CURRENT LAW: <u>Internal</u> Appeal and Grievance Procedures.

North Carolina law provides for two step internal appeal and grievance procedure that allows consumers to appeal denials of preauthorizations of covered services or other matters in dispute between the consumer and the health benefit plan. Consumers whose appeal of a preauthorization denial or whose first-level grievance review has been decided in favor of the insurer have a right to file a written grievance with the insurer and to have a panel investigate and make a determination regarding the grievance. Consumers also have a right to request the review, appeal and grievance through a person authorized to act on their behalf or through their health care provider.

Appeals of Utilization Review Decisions: Under current law, when a consumer requests authorization of a particular procedure or service or continued authorization of ongoing care, the insurer must make a determination within two business days of the request. If the insurer denies authorization (referred to as a noncertification), the consumer may informally appeal the denial. This informal appeal procedure allows the consumer to explain why the procedure should have been authorized. This appeal requires that the preauthorization denial be reviewed by at least one medical doctor, licensed in NC, who was not involved in the denial. The insurer must issue a written decision to the consumer and the health care provider within 30 days of the request for review. The insurer must provide an expedited review and issue a decision within four days when it is necessary to avoid jeopardizing the health of the patient.

First-Level Grievance Appeal: If the dispute concerns a matter of dissatisfaction other than a request for covered services, the consumer has a right to an informal review of the grievance. The current law requires that the insurer select someone with appropriate expertise, who was not involved in the matter, to evaluate the grievance. A written decision must be issued within 30 days of the request.

Second-Level Grievance Appeal: If the consumer is dissatisfied with the outcome of the informal, first level grievance appeal, or of the informal appeal of the denial of preauthorization, the consumer has a right to file a formal, second-level grievance appeal with the insurer. At this stage, the matter giving rise to the appeal must be evaluated by persons who were not previously involved in the matter, who are not

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employed by the insurer and who do not have a financial interest in the outcome of the appeal. However, these persons are appointed to serve on the panel by the insurer.

When the grievance concerns a utilization review matter or clinical issue, all members of the review panel must be health care professionals with appropriate expertise, including at least one clinical peer. One member of the panel may be an employee of the insurer if the panel is made up of three or more persons and the insurer included a clinical peer in the review of an appeal or first level grievance. The review must be held within 45 days of receipt of request and a written decision must be provided to the consumer seven days after the review meeting. An expedited review (within four days) must be provided if it is necessary to avoid jeopardizing the life or health of the patient.

BILL ANALYSIS:

Sections 1 through 4 of the bill divide Article 50 of Chapter 58 of the General Statutes, concerning general regulation of insurance, into five Parts and make conforming changes to existing language in the Article. Section 5 adds a new Part 4 to Article 50 of Chapter 58 to create a mechanism for independent, external review of an appeal decision upholding an initial noncertification decision or a second level grievance review decision that upheld an initial noncertification decision. The Part will apply to all persons who provide or perform utilization review.

Sections 6 amends the definition of "noncertification" under G.S. 58-3-190 to include (1) determinations and claims concerning whether a health care service provided in an emergency setting meets the prudent layperson standard for coverage; and (2) any review concerning whether a requested treatment is experimental, investigational, or cosmetic and the extent to which coverage under the health benefit plan is affected by that decision.

Section 7 amends the definition of "retrospective review" under G.S. 58-50-61(a)(17)g to specifically include reviews of claims concerning whether a health care service provided in an emergency setting meets the prudent layperson standard for coverage.

Section 8 amends G.S. 58-60-61(j), (Appeals of Noncertifications) to require informal reconsiderations of noncertifications to be conducted only after a written notice of the noncertification, meeting the requirements of G.S. 58-60-61 (h), has been issued. G.S. 58-60-61 (h) requires the notice to include the reasons for the noncertification, instructions on how to appeal the noncertifications, and instructions on how to request a written statement of the review criteria the insurer used in to make the noncertification. If the insurer is unable to reach an informal reconsideration decision in fewer than 10 business days, the informal reconsideration is to be treated as a formal appeal.

Section 9 adds a new subsection to G.S. 58-50-62 (Insurer grievance procedures), to provide similar requirements for procedures related to informal considerations of grievances as those outlined in Section 8.

Sections 10-12 makes clarifying and conforming amendments to current law.

Section 13 requires the Commissioner of Insurance to make biannual reports to the Joint Legislative Health Care Oversight Committee concerning the number and appropriateness of external appeals requested and conducted.

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Section 14 is a severability clause.

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Section 15 makes the bill effective July 1, 2001.

See attached chart for the key elements of Section 5, Independent External Review.

North Carolina Law Addressing External Grievance Review Procedures

1	Predominant State Regulator Involved in the External Review	Department of Insurance (DOI).
2	Entities Whose Decisions are Eligible for External Review	Health Insurer.
3	Who May Request External Review	Covered Person.
4	Decisions that are Eligible for External Review	1) A noncertification appeal decision and 2) a second level grievance review decision.
5	Dollar Threshold for External Review	None.
6	Cost Sharing Requirements	None.
7	Exhausting Internal Grievance Procedures	With some exceptions, a request for an external review shall not be made until the covered person has exhausted the insurer's internal grievance process. If the covered person has filed a grievance involving a noncertification appeal decision, but the insurer has not issued a written decision to the covered person within 45 days after the date it was filed and the covered person has not requested or agreed to a delay, the covered person is considered to have exhausted the insurer's internal grievance process.
8	Expedited Review	A covered person may make a request for an expedited external review with the Commissioner and if the Commissioner determines that the request meets the reviewability standards, an Independent Review Organization (IRO) will be assigned to complete the review on an expedited basis. If the IRO determines that the timeframe for a standard review would seriously jeopardize the life or health of the covered person or would jeopardize the covered person's ability to regain maximum function, the IRO must conduct the review and make the decision within four days.

9	General Description of Process	The request for external review is filed with the Department of Insurance. DOI notifies the health carrier of the filing of the request and provides a copy. DOI conducts a preliminary review to make sure the case is eligible. If the external review request is accepted, DOI notifies the insurer
	:	and covered person. The covered person has five days to select an approved Independent Review Organization (IRO). If the covered person hasn't selected an organization within five days, the
		Commissioner makes an assignment. The covered person and the insurer submit documents and information considered in making the internal review decision. The IRO conducts external
		review. IRO reverses or upholds determination, then notifies the covered person, Commissioner, and insurer. Except for a request for an expedited external review, all requests for external review must be made in writing.
10	Funding	The insurer against which a request for a standard review or an expedited external review is filed shall pay the cost of the IRO in conducting the external review.
11	Qualifications of Reviewer	The Commissioner approves IROs eligible to be assigned to conduct external reviews. The
		Commissioner develops an application form for initially approving and for re-approving organizations. Any IRO wishing to be approved submits the application form and includes with
		the form all documentation and information necessary for the Commissioner to determine if the
		organization satisfies the minimum qualifications established under G.S. 58-50-87.
		To be approved under G.S. 58-50-87, an IRO maintains written policies and procedures that
		govern all aspects of both the standard external review process and the expedited external review process that include, at a minimum:
		1. A quality assurance mechanism in place that ensures:
		a. That external reviews are conducted within the specified time frames and required notices are provided in a timely manner.
-		b. The selection of qualified and impartial clinical peer reviewers to conduct external reviews on behalf of the IRO and suitable matching of
		reviewers to specific cases.
		c. The confidentiality of medical and treatment records and clinical review criteria. AND
		d. That any person employed by or under contract with the IRO adheres to
		the requirements of this Part. 2. A toll-free telephone service to receive information on a 24-hour-day, seven-
	•	day-a-week basis related to external reviews that is capable of accepting,
		recording, or providing appropriate instruction to incoming telephone callers during other than normal business hours.

	Qualifications for	3. Agree to maintain and provide to the Commissioner the information set out in
	reviewers, cont.	G.S. 58-50-85.
		4. Agree to contractual terms or written requirements established by the Commissioner.
		5. A program for credentialing clinical peer reviewers.
		All clinical peer reviewers assigned by an IRO shall be physicians or other appropriate health care providers who meet the following minimum qualifications:
		1. Be an expert in the treatment of the covered person's injury, illness or medical condition.
		 Be knowledgeable about the recommended health care service or treatment through recent or current actual clinical experience treating patients with the same or similar injury, illness or medical condition of the covered person.
		 If the covered person's treating provider is a physician, a license from the North Carolina Medical Board, and if a specialist medical doctor, a current certification by a recognized American medical specialty board in the area(s) appropriate.
		 4. If the covered person's treating provider is not a medical doctor, hold an unrestricted license, registration, or certification in the same allied health occupation as the covered person's treating provider; and
		 Have no history of disciplinary actions or sanctions, including loss of staff privileges or participation restrictions, that have been taken or are pending by a
		hospital, governmental agency or unit, or regulatory body that raise a substantial question as to the clinical peer reviewer's physical, mental, or professional competence or moral character.
		competence of moral character.
2	Conflicts of Interest	Neither the IRO nor any clinical peer reviewer assigned by the IRO may have a material professional, familial, or financial conflict of interest with any of the following:
	· ·	1. The insurer.
		2. The covered person or the covered person's authorized representative.
		3. Any officer, director, or management employee of the insurer.
		4. The healthcare provider, the health care provider's medical group or independent
		practice association recommending the health care service or treatment.5. The facility at which the recommended healthcare service or treatment would be provided.
		6. The developer or manufacturer of the principal drug, device, procedure, or other therapy being recommended for the covered person.

	Conflicts of Interest, cont.	The IRO may not own or control, be a subsidiary of or in any way be owned or controlled by, or exercise control with a health benefit plan, a national, state or local trade association of health benefit plans, or a national, state or local trade association of health care providers
13	Standard of Review	 The IRO reviews all of the information and documents received. In addition to documents received, the IRO considers: the covered person's medical records, the attending health care professional's recommendation, consulting reports from appropriate health care professionals and other documents submitted, the terms of coverage under the covered person's health benefit plan, the most appropriate practice guidelines, including local practice guidelines, and any applicable clinical review criteria developed and used by the insurer or its designee utilization review organization. Medical necessity, as defined in G.S. 58-3-200(b). Within 45 days after the date of receipt of the request for external review, the IRO provides
		written notice of its decision to uphold or reverse the noncertification appeal decision or second level grievance review decision to the covered person, the insurer, and the Commissioner.
14	Time Frames	A covered person has 60 days after the date of receipt of a notice of a noncertification appeal decision or a second level grievance review decision to file a request with the Commissioner for an external review. Within 5 business days after the date of receipt of a request for an external review, the Commissioner has to complete a preliminary review of the request. If the request is not complete, the Commissioner will notify the covered person what information or materials are needed to review the request. The covered person has 90 days to furnish the materials. Once the request is accepted for review, the covered person has five days to select an IRO. The insurer or its designee utilization review organization has 7 days to forward documents and any information to the IRO. Within 45 days, the IRO has to provide written notice of its decision to uphold or reverse the decision to the covered person, the insurer, and the Commissioner. The decision of an expedited review must be made within four days.
15	Binding Nature of Decision	The decision is binding on the insurer. It is binding on the covered person except to the extent the covered person has other remedies available under applicable federal or state law. A covered person may not file a subsequent request for external review involving the same noncertification appeal decision or second level grievance review decision for which the covered person has already received an external review decision.

16	Attorney's Fees	Does not address attorneys.
17	Confidentiality Requirements	For purposes of conducting an external review, the insurer provides an authorization form by which the covered person authorizes the insurer to disclose protected health information, including medical records, concerning the covered person that are pertinent to the external review.
18	Liability of Reviewer	No IRO or clinical peer reviewer working on behalf of an IRO shall be liable in damages to any person for any opinions rendered during or upon completion of an external review, unless the opinion was rendered in bad faith or involved gross negligence.
19	Data Reporting	Each IRO is required to maintain for at least three years written records in the aggregate and by insurer on all requests for which it conducted an external review during a calendar year and submit a report, at least annually, to the Commissioner.
	· · · · · ·	Each insurer shall maintain for at least three years written records in the aggregate and for each type of health benefit plan offered by the insurer on all requests for external review that are filed with the insurer or that the insurer received notice of from the Commissioner.
20	Disclosure Requirements	Each insurer must provide each covered person a description of the plan's external review procedures, including a statement that informs the covered person of their right to file a request for external review.

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APPENDIX G

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SESSION 1999

00-DRM-001 (THIS IS A DRAFT AND IS NOT READY FOR INTRODUCTION)

Short Title: Internal Review Panelists.

(Public)

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Sponsors:

Referred to:

1

A BILL TO BE ENTITLED

2 AN ACT TO REQUIRE UTILIZATION REVIEW AND GRIEVANCE PROCEDURES PURSUANT TO G.S. 58-50-62 TO INCLUDE ON THE 3 REVIEW OR GRIEVANCE PANEL PROVIDERS LICENSED, CERTIFIED 4 OR REGISTERED IN NORTH CAROLINA IN THE SAME MEDICAL OR 5 ALLIED OCCUPATION AS THE PROVIDERS WHO ARE PARTIES TO 6 7 THE REVIEW OR GRIEVANCE. The General Assembly of North Carolina enacts: 8 9 Section 1. G.S. 58-50-61(d) reads as rewritten: 10 "§ 58-50-61. Utilization review. (d) Program Operations. -- In every utilization review program, an insurer 11 12 or URO shall use documented clinical review criteria that are based on sound 13 clinical evidence and that are periodically evaluated to assure ongoing efficacy. 14 An insurer may develop its own clinical review criteria or purchase or license

15 clinical review criteria. Criteria for determining when a patient needs to be 16 placed in a substance abuse treatment program shall be either (i) the diagnostic 17 criteria contained in the most recent revision of the American Society of 18 Addiction Medicine Patient Placement Criteria for the Treatment of Substance-19 Related Disorders or (ii) criteria adopted by the insurer or its URO. The 20 Department, in consultation with the Department of Health and Human

1 Services, may require proof of compliance with this subsection by a plan or 2 URO.

3 Qualified health care professionals shall administer the utilization review 4 program and oversee review decisions under the direction of a medical doctor. 5 A medical doctor licensed to practice medicine in this State shall evaluate the 6 clinical appropriateness of noncertifications. Compensation to persons involved 7 in utilization review shall not contain any direct or indirect incentives for them 8 to make any particular review decisions. Compensation to utilization reviewers 9 shall not be directly or indirectly based on the number or type of 10 noncertifications they render. In issuing a utilization review decision, an 11 insurer shall: obtain all information required to make the decision, including 12 pertinent clinical information; employ a process to ensure that utilization 13 reviewers apply clinical review criteria consistently; ensure that at least one 14 provider holding a valid North Carolina license, registration or certification in 15 the same medical or allied health occupation as the providers who are parties 16 to the review, or if the provider is a medical doctor, at least one clinical peer 17 of the party provider; and issue the decision in a timely manner pursuant to 18 this section."

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Section 2. G.S. 58-50-62 reads as rewritten:

20 "§ 58-50-62. Insurer grievance procedures.

(a) Purpose and Intent. -- The purpose of this section is to provide
standards for the establishment and maintenance of procedures by insurers to
assure that covered persons have the opportunity for appropriate resolutions of
their grievances.

(b) Availability of Grievance Process. -- Every insurer shall have a grievance process whereby a covered person may voluntarily request a review of any decision, policy, or action of the insurer that affects that covered person. The grievance process may provide for an immediate informal consideration by the insurer of a grievance. If the insurer does not have a procedure for informal consideration or if an informal consideration does not resolve the grievance, the grievance process shall provide for first- and second-level reviews of grievances; except that an appeal of a noncertification that has been reviewed under G.S. 58-50-61 shall be reviewed as a second-level grievance under this section.

35 (c) Grievance Procedures. -- Every insurer shall have written procedures for 36 receiving and resolving grievances from covered persons. A description of the 37 grievance procedures shall be set forth in or attached to the certificate of 38 coverage and member handbook provided to covered persons. The description 39 shall include a statement informing the covered person that the grievance 40 procedures are voluntary and shall also inform the covered person about the

1 availability of the Commissioner's office for assistance, including the telephone 2 number and address of the office.

3 (d) Maintenance of Records. -- Every insurer shall maintain records of each 4 grievance received and the insurer's review of each grievance, as well as 5 documentation sufficient to demonstrate compliance with this section. The 6 maintenance of these records, including electronic reproduction and storage, 7 shall be governed by rules adopted by the Commissioner that apply to insurers. 8 The insurer shall retain these records for three years or until the Commissioner 9 has adopted a final report of a general examination that contains a review of 10 these records for that calendar year, whichever is later.

11 (e) First-Level Grievance Review. -- A grievance may be submitted by a 12 covered person or his or her provider acting on the covered person's behalf.

13 (1) The insurer does not have to allow a covered person to attend 14 the first-level grievance review. A covered person may submit 15 written material. Within three business days after receiving a 16 grievance, the insurer shall provide the covered person with 17 the name, address, and telephone number of the coordinator 18 and information on how to submit written material.

(2) An insurer shall issue a written decision to the covered person and, if applicable, to the covered person's provider, within 30 days after receiving a grievance. The person or persons reviewing the grievance shall not be the same person or persons who initially handled the matter that is the subject of the grievance and, if the issue is a clinical one, at least one of whom shall be a medical doctor provider holding a valid North Carolina license, registration, or certification in the same medical or allied occupation as the providers who are parties to the grievance, or if the provider is a medical doctor, at least one clinical peer of the party provider with appropriate expertise to evaluate the matter. The written decision issued in a first-level grievance review shall contain:

a. The professional qualifications and licensure of the person or persons reviewing the grievance.

b. A statement of the reviewers' understanding of the grievance.

c. The reviewers' decision in clear terms and the contractual basis or medical rationale in sufficient detail for the covered person to respond further to the insurer's position.

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-		4	A reference to the evidence or documentation used as
1		d.	the basis for the decision.
2		~	A statement advising the covered person of his or her
3		e.	right to request a second-level grievance review and a
4			description of the procedure for submitting a second-
5			
6		a1	level grievance under this section. Grievance Review An insurer shall establish a second-
7	(f) Second-L	ever	Grievance Review All insurer shan establish a second
8	level grievance i	review	w process for covered persons who are dissatisfied with the
9	-	nce r	eview decision or a utilization review appeal decision.
10	(1)	An	insurer shall, within 10 business days after receiving a
11			lest for a second-level grievance review, make known to
12			covered person:
13		а.	The name, address, and telephone number of a person
14			designated to coordinate the grievance review for the
15			insurer.
16		b.	A statement of a covered person's rights, which include
17			the right to request and receive from an insurer all
18			information relevant to the case; attend the second-level
19			grievance review; present his or her case to the review
20			panel; submit supporting materials before and at the
21			review meeting; ask questions of any member of the
22			review panel; and be assisted or represented by a
23			person of his or her choice, which person may be
24			without limitation to: a provider, family member,
25			employer representative, or attorney. If the covered
26			person chooses to be represented by an attorney, the
27			insurer may also be represented by an attorney.
28	(2)	An	insurer shall convene a second-level grievance review
29		pan	el for each request. The panel shall comprise persons who
30		wer	e not previously involved in any matter giving rise to the
31		seco	ond-level grievance, are not employees of the insurer or
32		UR	O, and do not have a financial interest in the outcome of
33		the	review. A person who was previously involved in the
34		mat	ter may appear before the panel to present information or
35		ansy	wer questions. All of the persons reviewing a second-level
36		grie	vance involving a noncertification or a clinical issue shall
37		be	providers who have appropriate expertise, including at
38		leas	t one clinical peer. provider holding a valid North
39		Car	olina license, registration, or certification in the same
40		mee	lical or allied occupation as the providers who are parties

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1		to the grievance, or if the provider is a medical doctor, at			
2	least one clinical peer of the party provider. Provided,				
3	however, an insurer that uses a clinical peer on an appeal of a				
4	noncertification under G.S. 58-50-61 or on a first-level				
5		grievance review panel under this section An insurer may use			
6		one of the insurer's employees on the second-level grievance			
7		review panel in the same matter if the second-level grievance			
8		review panel comprises three or more persons.			
9	(g) Second-I	Level Grievance Review Procedures An insurer's procedures			
10	for conducting a	a second-level grievance review shall include:			
11	(1)	The review panel shall schedule and hold a review meeting			
12		within 45 days after receiving a request for a second-level			
13		review.			
14	(2)	The covered person shall be notified in writing at least 15			
15		days before the review meeting date.			
16	(3)	The covered person's right to a full review shall not be			
17		conditioned on the covered person's appearance at the review			
18		meeting.			
19	(h) Second-	Level Grievance Review Decisions An insurer shall issue a			
20	written decision	n to the covered person and, if applicable, to the covered			
21	person's provid	ler, within seven business days after completing the review			
22		ecision shall include:			
23	(1)	The professional qualifications and licensure of the members			
24		of the review panel.			
25	(2)	A statement of the review panel's understanding of the nature			
26		of the grievance and all pertinent facts.			
27	(3)	The review panel's recommendation to the insurer and the			
28		rationale behind that recommendation.			
29	(4)	A description of or reference to the evidence or			
30		documentation considered by the review panel in making the			
31		recommendation.			
32	(5)	In the review of a noncertification or other clinical matter, a			
33		written statement of the clinical rationale, including the			
34		clinical review criteria, that was used by the review panel to			
35		make the recommendation.			
36	(6)	The rationale for the insurer's decision if it differs from the			
37		review panel's recommendation.			
38	(7)	A statement that the decision is the insurer's final			
39		determination in the matter.			

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Notice of the availability of the Commissioner's office for (8) 1 assistance, including the telephone number and address of the 2 Commissioner's office. 3 (i) Expedited Second-Level Procedures. -- An expedited second-level review 4 5 shall be made available where medically justified as provided in G.S. 58-50-6 61(1), whether or not the initial review was expedited. The provisions of 7 subsections (f), (g), and (h) of this section apply to this subsection except for 8 the following timetable: When a covered person is eligible for an expedited 9 second-level review, the insurer shall conduct the review proceeding and 10 communicate its decision within four days after receiving all necessary 11 information. The review meeting may take place by way of a telephone 12 conference call or through the exchange of written information. (j) No insurer shall discriminate against any provider based on any action 13 14 taken by the provider under this section or G.S. 58-50-61 on behalf of a 15 covered person. (k) Violation. -- A violation of this section subjects an insurer to G.S. 58-2-16 17 70. (1997-519, s. 4.2.) Section 3. This act is effective when it becomes law.

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00-DRM-001



Bill Summary INTERNAL REVIEW PANELISTS

BILL ANALYSIS

Committee:	LRC/Managed Care Issues	Introduced by:	
Date:	April 27, 2000	Summary by:	Linda Attarian
Version:	FINAL DRAFT	· · · · · · · · · · · · · · · · · · ·	Committee Counsel

SUMMARY: The bill would amend G.S. 58-50-61 and 58-50-62, managed care grievance and appeals procedures, to require insurers to include on utilization review and grievance panels, providers who are licensed, certified or registered to practice in this State and who practices in the same medical or allied occupation as the providers who are parties to the review or grievance.

CURRENT LAW

Definition of clinical peer: The term "clinical peer" means a health care professional who holds a license in a state of the United States in the same or similar specialty and routinely provides the health care services subject to the review.

Utilization review panels: North Carolina law provides that all insurers establish a utilization review program to evaluate that health care services offered by the insurer are medically necessary and appropriate. Only "qualified health care professionals" may administer the utilization review program, and they must do so under the direction of a medical doctor. The law does not define "qualified health professionals." Thus the law does not prohibit insurers from including clinical peers or providers with the same license as the party to the review in the utilization review process, but it does not require their participation.

Grievance procedures concerning utilization review. North Carolina law establishes a two-step process for appeal and grievance when a health plan denies authorization for a particular type of health service pursuant to its utilization review program. Only a medical doctor licensed in North Carolina may make this denial. A denial of authorization may be appealed under G.S. 58-50-61, and the appeal panel must include a medical doctor licensed in North Carolina, but the law does not require the inclusion of a clinical peer at this stage of the appeal. If the panel upholds the denial, the insured may proceed to a second level appeal, which is conducted according to the provisions of G.S. 58-50-62. When the subject of the second level grievance is a clinical matter, the panel must be made up of health care professionals qualified to evaluate the matter, including at least one clinical peer. If an insurer uses a clinical peer on an appeal of a noncertification under G.S. 58-50-61 or on a first-level grievance review panel, then it may use one of its employees on the second-level grievance review panel in the same matter provided the panel contains three or more persons.

BILL ANALYSIS: Section 1 amends G.S 58-50-61(d) to require that, in issuing a utilization review decision, at least one provider holding a valid North Carolina license, registration, or certification in the same medical or allied occupation as the provider who is a party to the review is included in the utilization review decision making process. If one of the providers party to the review is a medical doctor, then the insurer must involve a clinical peer of that provider.

Section 2 amends G.S. 58-50-62 to require that an insurers' grievance panel includes in the first and second-level grievance review panels at least one provider holding a valid North Carolina license, registration, or certification in the same medical or allied occupation as the provider who are parties to the review. If one of the providers party to the grievance review is a medical doctor, then the insurer must

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include a clinical peer of that provider on the panels. Also, since the amended section requires the inclusion of a clinical peer on the appeal of a noncertification under G.S. 58-50-61 and on a first-level grievance review panel, the bill clarifies that an insurer may include one of the insurer's employees on the second-level review panel if it has three or more persons.

North Carolina General Assembly

Research Division, 733-2578

The act becomes effective when it becomes law.