The Anytown Medical Clinic

Coordinated Quality
Improvement
Program Plan

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1	Anytown Medical Clinic
2	Coordinated Quality Improvement Program Plan
3	
4	
5	I. Mission
6	
7	The mission of the Coordinated Quality Improvement Program is to improve the
8	quality of health care services rendered to patients and to identify, control and
9	reduce adverse health outcomes and medical malpractice events.
10	
11	In support of that mission, the Program will
12	
13	1. provide a coordinated framework for assessing, evaluating and improving
14	quality patient care and organizational functions
15	2. collect, maintain and analyze information pertinent to health care quality
16	improvement
17	3. educate and assist health care providers to lead and participate in
18	quality improvement activities such that health care quality is improved
19	and medical malpractice events are reduced.
20	
21	This framework will focus on patient satisfaction and quality care ensured
22	through involving participants in all aspects of providing services to patients.
23	
24	II. Coordinated Quality Improvement Program administration, authority
25	and accountability
26	
27	The Coordinated Quality Improvement Program will be administered by the
28	Quality Improvement Committee. The Committee shall be appointed by the

administrative leadership of the medical group and/or the Board of Directors.

The Committee derives its authority from the Board of Directors.

The ultimate accountability for quality improvement at the Anytown Medical Clinic rests with the Board of Directors (attached appendix). The Board of Directors has delegated to the Quality Improvement Committee the primary responsibility for quality improvement activities. It is the Board's expectation that quality improvement activities will involve all levels of the institution acting together collaboratively. The Board of Directors will review reports from the Committee, request further review or action as necessary, implement action plans and set policy for the institution and offer direction during planning activities of the Coordinated Quality Improvement Program.

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III. Quality Improvement Committee responsibilities

The Quality Improvement Committee provides guidance and inspiration for process development to design, assess, monitor and redesign program and service changes consistent with the goal of improving health care quality. The Committee is charged with the responsibility to implement and monitor system improvements to improve health care quality.

The Anytown Medical Center will have a written annual Quality Improvement Plan to identify and implement quality improvement measures consistent with the overall goals and resources of the Center. The plan will be based upon a calendar year and shall identify priorities for quality improvement activities. The plan will describe the core functions, structure, dimensions of focus and assigned responsibility for quality improvement activities. Creation, implementation and monitoring of the plan shall be the responsibility of the Quality Improvement Committee.

1	The	Quality Improvement Committee shall at a minimum have the following
2	resp	onsibilities:
3		
4	1.	Planning and structure of the Coordinated Quality Improvement Program
5		
6		The Program shall include three dimensions of focus: systems and
7		processes, resource utilization and risk management.
8		
9		Within each area of focus, quality improvement may consist of the
10		following activities:
11		issue identification and assessment
12		• prioritization
13		• process improvement
14		• monitoring
15		
16		A. Issue identification and assessment
17		
18		Issues may be identified via a number of different data sources:
19		• patient input
20		• patient comment forms
21		 quality control procedures
22		• utilization review
23		• incident reports
24		• credentialing
25		• clinician peer review
26		 cost accounting or productivity data
27		• referral data
28		 satisfaction surveys from patients and clinicians
29		e part to a company to the company of the company o
		\cdot

1 Issues may be assessed in accordance with process performance 2 compared internally over time and/or process performance 3 compared with external standards. The dimensions of process 4 performance may include whether the process is appropriate, 5 available, timely, effective, efficient, safe, and respectful. 6 7 B. Prioritization 8 9 After issue identification and assessment, areas of process 10 improvement shall be prioritized in accordance with the overall goals 11 and resources of the Clinic. 12 13 C. Process improvement 14 15 Improving health care quality may be accomplished by improving the 16 process of health care delivery. Process improvement may include: 17 18 1. Identifying a potential process improvement through issue 19 identification and assessment and prioritization 20 2. Formulating a plan of action 21 3. Testing the strategy of change 22 4. Assessing the results of the test strategy 23 5. Implementing the process improvement 24 25 D. Monitoring 26 27 System improvement shall be monitored to ensure that the goals of 28 the Program are being met. System improvement activities may be 29 modified as indicated by assessing the effects of the improvement 30 activity as related to the mission of the Program.

1			$\epsilon_{ij} = \epsilon_{ij} (b_j)^{(ij)}$
2	2.	Oversee and coordinate the Progr	ram;
3			
4		The Quality Improvement Committ	ee is responsible for overseeing and
5		coordinating the Coordinated Qua	lity Improvement Program. The
6		Committee will maintain managen	al oversight to ensure that Program
7		activities are consistent with the F	Program's mission and emphasize the
8		improvement of health care and th	ne reduction of adverse health outcomes
9		and medical malpractice events.	At its discretion, the Committee may
10		delegate Program activities to sub-	-committees, employees and agents of the
11		medical group while retaining ove	rsight responsibility.
12			
13	3.	Quality Improvement Committee	membership
14			
15		The Committee shall be comprised	of individuals who are broadly
16		representative of the medical grou	p and who are in a position to
17		effectively assess, evaluate, plan	and implement strategies pertinent to
18		patient care and organizational pr	ocesses. The membership of the
19		Committee shall include personne	from the following departments or job
20		responsibilities:	
21			
22		C	Co-Chairs
23		Med	ical Director
24		Quality Improvem	ent Program Coordinator
25			
26		Medical staff members	Other staff members
27		Family Practice	Medical records
28		Internal Medicine	Managed care coordinator
29		Pediatrics	Administration
30		Surgery	Clinical services

1		Obstetrics/gynecology	Others as needed
2		Others as needed	
3			
4	4.	Quality Improvement Committee s	sub-committees
5			
6		The Committee shall have the auth	ority and responsibility to appoint sub-
7		committees as necessary. All such	sub-committees shall report directly to
8		the Quality Improvement Committee	e .
9			
10	5.	Quality Improvement Committee r	neetings
11			
12		Meetings shall be held at least mor	nthly or more often as needed at the
13		discretion of the Committee.	
14			
15	6.	Confidentiality	
16			
17		Members, participants and visitors	of the Committee shall sign an
18		agreement of confidentiality relevan	nt to the functions of the Coordinated
19		Quality Improvement Program and	its activities.
20			
21	7.	Retrospective and prospective revi	ew of services to improve the quality of
22		health care and reduce medical m	alpractice events by measuring key
23		characteristics such as effectivene	ss, accuracy, timeliness and cost;
24			
25		Retrospective and prospective revie	w of services will be accomplished by:
26			
27		 review of incident report forms (attached appendix)
28		 providing and monitoring prever 	ntative health care
29		 clinician peer review 	
30		 patient satisfaction surveys 	

1		• outcomes research
2		 review of patient comment forms (attached appendix)
3		• contract review data from health care payors
4		
5	8.	Review the categories and methodologies of services offered and to be
6 7		offered in the future in order to improve health care outcomes;
8		Review of services offered currently and to be offered in the future will be
9		accomplished to ensure those services provide quality patient care in a
10		cost-effective manner that improves health care outcomes.
11		
12	9.	Collect, maintain, analyze and review Program information in order to
13		revise health care policies and procedures as necessary;
14		
15		Using the data sources noted in sections 1 or 7 above, the Committee will
16		collect, maintain, analyze and review data in order to
17		
18		 recommend further analysis or study;
19		 recommend revision of Clinic policy and/or procedure; or
20		 recommend training and/or education of clinicians and staff.
21		
22	10.	Report at least semi-annually on Program activities and actions to the
23		governing body of the medical group.
24		
25		The Committee shall report on Program activities and actions to the Board
26		of Directors at least semi-annually. An annual written report shall be
27		submitted by the Committee to the Board of Directors which includes a
28		report of quality improvement activities, actions, outcomes, trending of
29		clinical and service indicators and other information pertinent to the
30		mission of the Committee. Reports may be made more often as necessary

1		or on an ad-hoc basis. An annual summary of Program activities and
2		actions shall be made available to the governing body or other interested
3		parties.
4		
5	11.	The clinical staff members of the Quality Improvement Committee are
6		responsible for educating, counseling and monitoring the clinical staff of
7		the Clinic to ensure adherence to all applicable policies, procedures and
8 -		standards. The second of the public of the second of the s
9		
10	12.	The non-clinical staff members of the Quality Improvement Committee
11		are responsible for educating, counseling and monitoring the non-clinical
12		staff of the Clinic to ensure adherence to all applicable policies,
13		procedures and standards.
14		
15 16	13.	Reporting requirements
16		
17		The Committee will develop a process to assure compliance with any
18		reporting requirements to appropriate state, local or federal authorities.
19		Items to be reported include professional misconduct, malpractice payments
20		to patients and other items required by law.
21		
22	IV.	Coordinated Quality Improvement Program documentation
23		
24	Minu	tes shall be recorded at all meetings of the Quality Improvement
25	Comi	mittee and any appointed sub-committees. Any quality improvement
26	studi	es, recommendations of studies, incident reports, provider evaluations,
27	and o	other recorded information pertinent to the Coordinated Quality
28	Impro	ovement Program shall be attached or incorporated by reference into the
29	Comr	nittee minutes. It is the intent of the Committee that any Program

documentation shall be protected from legal discovery to the fullest extent allowed by law.

The Committee minutes will be available for internal review within the medical group on a need to know basis as determined by the Committee. Summaries of the minutes with identifying names removed may be divulged or provided to health care payors, governmental or regulatory agencies or other entities at the sole discretion of the Committee. Such disclosure will only be made when disclosure will not effect confidentiality requirements or impair any legal position of the medical group in any future legal or regulatory proceeding.

All Committee minutes, incident reports, provider evaluations and all other recorded information pertinent to the Coordinated Quality Improvement Program shall be clearly marked as follows: "Coordinated Quality Improvement Program work product".

V. Information collection and maintenance

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The Committee shall continually collect and maintain information concerning

- 1. experience with negative health care outcomes and injurious incidents; and
- 2. professional liability premiums, settlements, awards, and costs for injury 24 prevention, safety improvement and health care improvement activities.

Information on adverse health care outcomes and injurious incidents will be collected concurrently and maintained using incident reporting, peer review, patient comment forms and patient records of unexpected individual adverse health care outcomes.

Information on professional liability premiums, settlements, awards and costs for injury prevention, safety improvement and health care improvement activities will be jointly provided to the Committee by the malpractice insurer and administration of the medical group.

3. Safety improvements, health care system improvement activities and resource utilization

Information on safety improvements, health care system improvement activities and resource utilization will be provided to the Committee by the administration, employees, or agents of the medical group using whatever external or internal data sources are deemed appropriate by the Committee.

This information shall be periodically provided to the governing body and the providers of the medical group.

VI. Incident reporting

Accidents, injuries, negative health care outcomes, patient complaints and other information pertinent to health care quality improvement will be reported to the Committee as an incident under the Quality Improvement Program. An incident report form shall be used to ensure consistent reporting. Employee incident reports will be sent for review to the Center's safety officer. Patient incident reports will be sent for review to the Center's Medical Director. All incident reports will be reviewed and analyzed by the Quality Improvement Program Coordinator with appropriate action taken as necessary.

Maloccurrences that may result in adverse health outcomes or health care malpractice claims will be investigated and either resolved internally by the medical group, to the extent feasible, or reported to the malpractice insurer of the medical group for appropriate investigation and resolution.

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VII. Quality Improvement education

The Quality Improvement Committee shall sponsor quality improvement educational activities for health care providers at least annually or more often as needed at the discretion of the Committee. These educational activities will be in addition to other continuing education activities provided or facilitated by the Center. These educational activities may include, but are not limited to:

1. Quality improvement in health care

Education in health care quality improvement may be provided by local health care facilities; or providers, employees, agents or consultants of the medical group. The Quality Improvement Coordinator shall be responsible for ensuring that quality improvement data and information is communicated to the clinicians and staff of the Center.

2. Safety and injury prevention

Safety and injury prevention, infection control or hazardous materials education may be provided by local health care facilities; providers, employees, agents or consultants of the medical group; or the risk management educational programs of the medical group's malpractice insurer. Safety and injury prevention education is provided to all staff at the time of employment orientation, on an ad hoc basis and as a result of mandatory continuing education or activities that may be required by law or regulation.

1	3.	Responsibilities for reporting professional misconduct
2		
3		Education on responsibilities for reporting professional misconduct may be
4		provided by state and/or federal regulatory or enforcement agencies; or
5 6		providers, employees, agents or consultants of the medical group.
7	4.	Legal aspects of providing health care
8		
9		Education on legal aspects of providing health care may be provided by
10		local health care facilities or the risk management educational programs of
11		the medical group's malpractice insurer. All clinicians will comply with
12		State requirements for continuing medical education on risk management.
13		
14	5.	Improving communication with patients (1)
15		· · · · · · · · · · · · · · · · · · ·
16		Education on improving communication with patients may be provided by
17		local health care facilities; providers, employees, agents or consultants of
18		the medical group; or the risk management educational programs of the
19		medical group's malpractice insurer.
20		
21	6.	Causes, prevention and reduction of malpractice claims
22		
23		Education on the cause, prevention and reduction of malpractice claims
24		may be provided by the risk management educational programs of the
25		medical group's malpractice insurer.
26		
27	VIII.	Provider evaluation
28		
29	The A	anytown Medical Center has a physician and mid-level provider
30	crede	entialing process. Offers of employment are contingent upon satisfactory
		12

1	passage of the credentialing process. Credentialing included verification of	
2	licensure, DEA certification, education, residency, board certification or	
3	eligibility, and information from the National Practitioner Data Bank.	
4		
5	As appropriate to the scope of the provider's practice, every provider employed	
6	by the Center shall be periodically evaluated for mental and physical capacity,	· 7.
7	competence in delivering health care, verification of licensure, medical staff	
8	appointments, and reports from the National Practitioner Data Bank. Periodic	
9	evaluation of all providers will be done at least every two years.	
10		
11	The provider evaluation may include any of the following elements as deemed	
12	appropriate by the Quality Improvement Committee:	
13		
14	Patient satisfaction survey	
15	2. Evaluation by peer providers within the same clinical department and	
16	throughout the medical group as appropriate	
17	3. Evaluation by support staff as appropriate	
18	4. Verification of licensure	
19	5. Verification of current medical staff appointments	
20	6. Verification of reports to the National Practitioner Data Bank	
21	7. Evaluation of CME participation	
22	8. Evaluation of risk management issues	
23	9. Review of a representative sample of current charts	
24	10. Productivity (* (* 1948) * (* 1948)	
25		
26	IX. Complaint resolution	
27		

13

The Program will incorporate a procedure to investigate and resolve, in a timely

fashion and to the extent feasible, patient complaints pertaining to accidents,

injuries, treatment and other events that may result in adverse health

28

29

outcomes or claims of health care malpractice. The Quality Improvement Committee shall appoint a member to serve as a patient liaison for complaint resolution. Records shall be kept of patient complaints and the response thereto. These records shall come under the purview of the Program and are Program work product. Coordinated Quality Improvement Program information concerning Χ. providers Program information regarding a provider will be maintained within the medical group's personnel or credential file for that provider. Patient confidentiality will be maintained to the extent necessary to facilitate Program activities. All such information maintained on a provider pertinent to the Coordinated Quality Improvement Program shall be clearly marked as falling under the purview of the Program. Draft eight: 8/14/95 M.G. Lloyd

Appendix

Confidentiality agreement

I,	, acknowledge and agree that I am a
participant in the Coordinated	Quality Improvement Program (CQIP) conducted
under the auspices of the Qual	ity Improvement Committee at the Anytown
Medical Center. As a participal	nt, I understand and agree that maintaining
confidentiality is vital to the qu	ality improvement process.
I agree to respect and maintain	the confidentiality of all discussions, records,
and information, whether oral of	or recorded in any form or medium, produced in
connection with the CQIP. I fur	rther agree to not disclose any such information
except as required by the activi	ties of the CQIP or by law.
Signature	
Date	

Clinician credentialing application

Clinician name:			
Da	te of birth: Place of birth:		
Ple	ease attach the following items to this application:		
	Current Washington State license		
	Current DEA certificate		
	Professional school diploma		
	ECFMG (if applicable)		
	Residency certificate (if applicable)		
	Fellowship certificate (if applicable)		
	Board certification:		
	Report from the National Practitioner Data Bank (Call 1-800-767-6732 to obtain the report)		
	Additional credentialing information		
•	you answer yes to any of the listed questions, please explain further on a parate sheet of paper.		
1.	Has your license to practice or DEA certificate ever been denied, restricted, limited, suspended, subject to probationary terms, revoked, or voluntarily surrendered?		
	☐ Yes ☐ No		
2.	Have you ever been subject to disciplinary proceedings by any governmental agency, medical or professional society resulting in reprimand, censure, sanction, restriction or modification of your practice, either voluntary or involuntary, or are you currently the subject of an		

	administrative, judicial agency or society?	or disciplinary	proceeding or review by any such						
		Yes	□ No						
3.	ever been denied, restri- subject to probationary have disciplinary proces	eve your privileges or membership at any hospital or medical institution er been denied, restricted, suspended, reduced, modified, terminated, bject to probationary terms, voluntarily surrendered or not renewed or we disciplinary proceedings at any hospital or medical institution ever en instituted against you?							
	٥	Yes	□ No						
4.	Has your membership is ever been denied, suspensurrendered?	n any medical s ended, revoked,	society or professional organization restricted or voluntarily						
		Yes	□ No						
5.	ability to practice withir	n the scope of ex scope of your ca	ealth problems that may limit your xisting hospital medical staff apacity as a provider of medical						
	0	Yes	□ No						
	If so, are there any reas request?	onable accomm	odations that you may require or						
6.	Have you ever been convicted for an act committed in violation of ordinance other than traffic offenses?								
	ū	Yes	□ No						
7.		sional corporat	al malpractice ever been brought tion, your employer or your hospital I care provided by you.						
		Yes	□ No						

		ce ever been declined, cancelled, nor ch as a premium surcharge or
☐ Yes	3	□ No
Applicant's r	epresentat	tion (read carefully)
supplemental submission is co which are reasonably likely to Clinic in considering this appli agree that this application will	omplete and influence th cation have in part be t	ontained in this application and any it true and that no material facts he judgment of Anytown Medical been omitted. I understand and the basis of the employment Medical Clinic of any changes in the
Applicant's author	ization and	d release (read carefully)
an inquiry and investigation of competence, including any oth Medical Clinic, may be conduct authorized representatives. I elinvestigation and hereby authorized representatives and organizations in which I am or consultants or agents; any hos have ever held staff privileges of denied; any state licensing ages of higher learning; any attending carriers; prior employers; profecclinic or its duly authorized representatives and the member committees from any and all leas a result of any communication made, or any acts performed, in investigation initiated by the Arrepresentatives.	my profess or matters of ted by the A expressly convice the relative have been a pitals or me or have had ney; any uning or treating ssional assopresentative as Anytown ers or consugal liabilitie ons, reports n good faith	ease and exchange of information on between any professional a member; their insurance edical institutions at which I hold or an application for staff privileges niversity, college, school or institution g physicians; any prior insurance sociates; and the Anytown Medical es. I hereby release and discharge Medical Clinic, its duly authorized ultants or any established peer reviews which might otherwise be incurred as, disclosures and recommendations in, in connection with any inquiry or dical Clinic or its duly authorized
Applicant's signature		Date

Declaration

I declare under penalty of perjury ur	nder the laws of the State of Washington
that the information provided in this	Clinician Credentialing Application is
complete, true and correct.	3 11
*	
Date and place	Signature

Part La Son

The Anytown Medical Clinic 1995 Quality Improvement Plan

The Anytown Medical Clinic has a planned, systematic, organization-wide approach to assessing, measuring, and improving performance. Organizational performance activities are carried out collaboratively, and include appropriate departments and staff. Preparation of each year's plan will begin in December of the prior year. Monitoring activities will work through the calendar year with quarterly and annual reports submitted to the Quality Improvement Coordinator.

Monitoring activities, data collection and data sources

Monitoring activities will be derived from the clinics core functions. The core functions or processes of the Clinic are defined through management structuring and include the following:

Operations Services Information Services
Clinical Services Managed Care Services
Business Services Financial Services
Human Resource Services Medical Staff

Data gathering about the performance on each of these processes will be obtained internally and externally in order to compare to other data bases. Internal information may be gathered from the following sources:

- · Medical Record Review
- Case Reviews
- Patient Satisfaction/Comment Surveys
- Staff Meetings
- Clinical Indicator Measurements
- Risk Management Outcome Data
- Patient Grievance Data
- Audits from health care payors

External comparative information comes from peer group data including:

- MGMA Annual Cost Survey
- UMGA Bench-marking
- National Clinical Standards
- Clinical Health Plan Data

The aspects of care and service choices will be those which are deemed to be most important to the quality of patient care and service based upon available organizational resources. Areas of focus will be based upon the following elements:

- High risk to patients
- High volume aspects of care

All patient care and service that the Anytown Medical Clinic provides will be considered in determining priorities for ongoing monitoring. The key patient care and service functions at the clinic shall encompass the following:

- The appointment system
- Practice guidelines
- Infection control
- Managerial and governance activities, such as patient accounts, marketing
- Human Resources Regular staff and physician reviews, orientation, training, continuing education
- Patient/Family education
- Referral systems
- Telephone system (triage)
- Utilization management system
- Patient representation
- Patient conditions and diagnoses served
- Service accessibility times care and service are provided
- Return clinic visits

Quality improvement priorities for 1995

Clinic priorities will be reviewed on an annual basis. The 1995 priorities are:

*Service Excellence

Affects all patients
Positions clinic for improved marketability
Improves internal morale
Plays a role in reducing risk management

*Clinical Areas of focus

Immunizations follow-up Tracking system for radiologic risk area High cost pharmaceuticals Mammography,
Patient instruction
Asthma treatment modalities
Patient Recall System (paps, hemocult, etc)

Places patients at serious risk if not performed or performed well Places clinic at great risk if not done effectively & efficiently

Plays a role in increased costs to our patients
Standardizes treatment & assists in clinical staff education
Represents a wide age distribution
Represents HEDIS clinical outcome indicators

Quality improvement methodology

The approach for improvement for the 1995 organization priorities will be carried out in the following manner:

Process design is based upon the organization's

Mission

Strategic Plan

The needs and expectations of patients, staff, and physicians Current information about the performance of the processes & their outcomes from reference organizations

Data will be collected to assess processes, measure the level of performance & stability of important existing processes, to identify areas for possible improvement. Data will relate to the following dimensions of performance:

- Efficacy
- Appropriateness
- Availability
- Timeliness
- Effectiveness
- Continuity
- Safety
- Efficiency
- Respect & Caring

After the data is collected and priorites are set, process improvement shall be conducted as set forth in the CQIP plan.

Assigned Responsibility for quality improvement

The Administrative leaders are responsible for overseeing the design and fostering clinic wide approaches to improving quality, assigning quality

improvement responsibilities within the organization, and setting strategic priorities for quality assessment and improvement throughout the clinic.

Administrative leaders shall be defined as the Governing Board, The Medical Director, the CEO, the Director of Operations, the Clinical Services Manager, the Director of Human Resources, and the Director of Managed Care of the Anytown Medical Clinic.

The Quality Improvement Coordinator is responsible for coordinating, educating, monitoring and counseling the non-medical staff to ensure departmental implementation of the quality improvement process.

Clinical, Ancillary, and Support managers and service/program supervisors will be responsible for ensuring that the activities in their areas are encompassed by the quality improvement monitoring and evaluation activities.

It shall be the responsibility of every clinic employee to participate in improving organizational performance and improving quality of care.

INCIDENT REPORT Coordinated Quality Improvement Program work product Do not file or reference in medical record

Date of inci	dent:	Time:	Incide	Incident location:							
Affected pa	rty 🖸 Patient 🚨 Visitor 🖂] Employee □ Me	edical staff	Sex		 Vale		Female	Age:		
Affected pa	rty name and address:							Phone number:			
Description	of affected party's injury or comp	laint:									
Name and p	phone number of witnesses:				***************************************						
			er (des Mercelon) (f. des S Sentendo (f. des Santas (des Santas))		Van de	Devices	- P(67.5)-K				
Incident des	scription:										
Possible co	ntributing causes:		•								
Corrective a	action taken:										
Recommen	dations to prevent future incident										
Action	the state of the s	me)									
taken by person	☐ Employee supervisor notified	I (name)									
reporting incident											
Reported b	Othery		Title/depart	ment/c	elinic:				Date:		

INCIDENT REPORT Coordinated Quality Improvement Program work product Do not file or reference in medical record

	Action taker	by QI reviewer	oppeland Sign Son Signature and Artist				
□ No further action necessary □ Staff/clinician education □ Policy/procedure change □ Referred to Business Offi □ Attending clinician notified □ Staff/clinician counseling □ Referred to QI committee □ Maintenance notified □ Referred to Administration □ Referred to Medical Director □ Fee waiver/reimbursement □ Equipment repair or replain or replai							
oate: Insurance comp	any:	Contact r	person;				
ther actions taken:							
ther actions taken:							
dditional comments:							
	The Late						
		······································					
							
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