HOW TO SUCCESSFULLY PREPARE FOR SURVEYS AND RESPONDING TO AN ADVERSE SURVEY/LICENSURE ACTION

PACAH 2019 FALL CONFERENCE September 24, 2019



Presented by Tanya Daniels Harris, Esquire

INCREASE IN CMS/DOH OVERSIGHT OF NURSING HOMES

- Performance Audit of DOH Regulation and Oversight of Nursing Facilities July 26, 2016
 - Staffing
 - Disposition of Complaints
 - Inadequate Civil Money Penalties
- Pennsylvania DOH doubled its fines against nursing homes to approximately \$2.6 million in 2018

- Follow-up report issued by Auditor General on July 23, 2019.
 - Improve surveyor training to achieve consistency amongst surveys
 - DOH should exercise its authority to require additional nursing home staff where direct care is lacking
 - DOH to follow CMS' resident-centered guidance for handling complaints
 - Acknowledgment that DOH has increased the use of CMPs, but recommendation that DOH track effectiveness of fines to show that oversight tool leads to improved outcomes
 - Adoption of more stringent and clearly outlined policies for vetting nursing facility license applicants

- DOH Civil Penalty Assessment Guideline 12/19/16:
 - Factors to be considered when issuing civil penalties:
 - Statutory provisions authorizing civil penalties under HCFA
 - Recommendations contained in PA Auditor General's Performance Audit Report (July 2016)
 - DOH's interest in effective regulation to promote the highest possible quality of care and services for LTC residents in PA
 - Any facility with a survey exit date on or after 1/1/2017 may be subject, when warranted, to civil penalties calculated on a per violation per day basis pursuant to 35 P.S. § 448.817

 Updated DOH Civil Penalty Assessment Guideline – 3/30/18:

• Any facility with a survey exit date on or after 1/1/2017, may be subjected, when warranted, to civil penalties calculated on a *per instance* or per day basis, *or both*, pursuant to 35 P.S. § 448.817.

- Updated DOH Civil Penalty Assessment Guideline – 3/30/18:
 - When determining whether civil penalties are warranted, DOH will consider the facility's compliance history, including but not limited to the following:
 - Whether the facility's violations resulted in harm or death to a resident;
 - The facility's most current deficiency report;
 - The threat or potential threat to resident health and safety;
 - The number of residents at risk or affected by the noncompliance;
 - The facility's plan of correction;
 - Similar survey findings where sanctions were imposed; and
 - Repeat noncompliance in the same or similar regulatory categories.

SUMMARY OF DOH CIVIL PENALTIES IMPOSED*

Year	Range of Civil Penalties	Total Amount of CP's for the year
2016	\$1,000 - \$60,800	\$412,200
2017	\$1,500 - \$100,000	\$1,028,750
2018	\$1,500 - \$52,250	\$2,855,299
2019 (as of September)	\$500 - \$33,000	\$1,111,310

^{*}Chart based on sanctions disclosed on DOH's website as of 9/16/2019

OVERVIEW OF INCREASED ENFORCEMENT CMS-CMP Annual Adjustments

	Pre-inflation (prior to 2016)	2016	2017	2019
Category 2	Min. \$50	Min. \$103	Min. \$105	Min. \$107
Per Day	Max. \$3,000	Max. \$6,188	Max. \$6,289	Max \$6,417
Category 2	Min. \$1,000	Min. \$2,063	Min. \$2,097	Min. \$2,140
Per Instance	Max. \$10,000	Max. \$20,628	Max. \$20,965	Max. \$21,393
Category 3	Min. \$3,050	Min. \$6,291	Min. \$6,394	Min. \$6,525
Per Day	Max. \$10,000	Max. \$20,628	Max. \$20,965	Max. \$21,393
Category 3 Per Instance	Min. \$1,000	Min. \$2,063	Min. \$2,097	Min. \$2,140
	Max. \$10,000	Max. \$20,628	Max. \$20,965	Max. \$21,393

Commonwealth v. Golden Gate Nat'l Senior Care, LLC, 194 A.3d 1010 (Pa. 2018)

- In June, 2015, the AG of PA sued a NF chain for violation of the PA Unfair Trade Practices and the Consumer Protection Law, asserting claims for false advertising and fraud due to understaffing. In March, 2017, PA Commonwealth Court found that the marketing statements were mere "puffery" rather than material representations.
- The AG appealed to the PA Supreme Court, and in September, 2018, the PA Supreme Court reinstated the claims against the NF, stating that the NF made materially misleading statements about the nature and quality of care provided to their residents. Specifically, representations in brochures guaranteeing clean linens, fresh bedside ice water, and snacks at any time were contradicted by poor care at the facility. The Court found that the UFTPCPL violations can be asserted for any fraudulent practice, whether or not the practice is related to advertising or an inducement to enter the facility.
- Moreover, the PA Supreme Court recognized that statements made in Plans of Care and in Medicare/Medicaid claim submissions can be actionable under the Unfair Trade Practices Act if the provider fails to deliver the care outlined in the plans or provide the services claimed.

OVERVIEW OF RELEVANT SURVEY AND ENFORCEMENT UPDATES

Final Revised Policies re: Immediate Imposition of Federal Remedies

- Substantive revisions to the prior guidance include:
 - When the current survey identifies Immediate Jeopardy (IJ) that does not result in serious injury, harm, impairment, or death, the CMS Regional Offices may determine the most appropriate remedy.
 - Past Noncompliance deficiencies are **not** included in the criteria for immediate Imposition of Remedies.
 - For Special Focus Facilities (SFFs), S/S level "F" citations under tags F812, F813, or F814 are excluded from immediate imposition of remedies.

QSO 18-18-NH Posting Date - 6/15/2018

OVERVIEW OF RECENT SURVEY AND ENFORCEMENT UPDATES

REVISIONS TO APPENDIX Q, GUIDANCE ON IMMEDIATE JEOPARDY

- Revision creates a Core Appendix Q that will be used by surveyors of all provider and supplier types in determining when to cite immediate jeopardy.
- Drafted subparts to Appendix Q focus on immediate jeopardy concerns occurring in nursing homes and clinical laboratories since those provider types have specific policies related to immediate jeopardy.
- To cite immediate jeopardy, surveyors determine that (1) noncompliance (2) caused or created a <u>likelihood</u> that serious injury, harm, impairment, or death to one or more recipients would occur or recur; and (3) immediate action is necessary to prevent the occurrence or recurrence of serious injury, harm, impairment, or death to one or more recipients.
- Removal of concept of "culpability" and replaced with concept of noncompliance.
- A template has been developed to assist surveyors in documenting the information necessary to establish each of the key components of immediate jeopardy.

QSO 19-09-ALL Posting Date – 03/05/2019

PREPARING FOR A SURVEY

- New LTC Survey Process (Effective November 28, 2017)
 - One unified survey process that will utilize strengths from both the Traditional survey process and Quality Indicator Survey (QIS) process
 - Goal of being more effective and efficient
 - Focus is resident-centered
 - New survey process provides structure to ensure consistency while allowing surveyors autonomy
 - New survey process will be an automated process (i.e., computer software-based).

- Three Parts to New LTC Survey Process:
 - Initial Pool Process
 - Sample Selection
 - Investigation

- Survey Team Coordinator Offsite Preparation
 - CASPER 3 report for pattern of repeat deficiencies
 - Results of last standard survey
 - Complaints and Facility Reported Incidents (FRI) since last standard survey

- Facility Entrance
 - Team Coordinator coordinates an Entrance Conference
 - Entrance Conference Worksheet
 - Matrix
 - Initial brief visit to kitchen
 - Surveyors go to assigned areas

Entrance Conference Worksheet

DEPARTMENT OF HEALTH AND HUMAN SERVICES CENTERS FOR MEDICARE & MEDICAID SERVICES

ENTRANCE CONFERENCE WORKSHEET

INFORMATION NEEDED FROM THE FACILITY IMMEDIATELY UPON ENTRANCE Complete matrix for new admissions in the last 30 days who are still residing in the facility. 3. An alphabetical list of all residents (note any resident out of the facility). 4. A list of residents who smoke, designated smoking times, and locations. ENTRANCE CONFERENCE 5. Conduct a brief Entrance Conference with the Administrator. 6. Information regarding full time DON coverage (verbal confirmation is acceptable). 7. Information about the facility's emergency water source (verbal confirmation is acceptable). 8. Signs announcing the survey that are posted in high-visibility areas. 9. A copy of an updated facility floor plan, if changes have been made. 10. Name of Resident Council President. 11. Provide the facility with a copy of the CASPER 3. INFORMATION NEEDED FROM FACILITY WITHIN ONE HOUR OF ENTRANCE 12. Schedule of meal times, locations of dining rooms, copies of all current menus including therapeutic menus that will be served for the duration of the survey and the policy for food brought in from 13. Schedule of Medication Administration times. 14. Number and location of med storage rooms and med carts. 15. The actual working schedules for licensed and registered nursing staff for the survey time period. 16. List of key personnel, location, and phone numbers. Note contract staff (e.g., rehab services). 17. If the facility employs paid feeding assistants, provide the following information: a) Whether the paid feeding assistant training was provided through a State-approved training program by qualified professionals as defined by State law, with a minimum of 8 hours of training; The names of staff (including agency staff) who have successfully completed training for paid feeding assistants, and who are currently assisting selected residents with eating meals and/or c) A list of residents who are eligible for assistance and who are currently receiving assistance from paid feeding assistants. INFORMATION NEEDED FROM FACILITY WITHIN FOUR HOURS OF ENTRANCE 18. Complete the matrix for all other residents. The TC confirms the matrix was completed accurately. 19. Admission packet. 20. Dialysis Contract(s), Agreement(s), Arrangement(s), and Policy and Procedures, if applicable 21. List of qualified staff providing hemodialysis or assistance for peritoneal dialysis treatments, if 22. Agreement(s) or Policies and Procedures for transport to and from dialysis treatments, if applicable.

24. Hospice Agreement, and Policies and Procedures for each hospice used (name of facility designee(s)

25. Infection Prevention and Control Program Standards, Policies and Procedures, and Antibiotic Stewardship Program. ☐ 26. Influenza / Pneumococcal Immunization Policy & Procedures. 27. QAA committee information (name of contact, names of members and frequency of meetings). 28. QAPI Plan. 29. Abuse Prohibition Policy and Procedures. 30. Description of any experimental research occurring in the facility 31. Facility assessment 33. List of rooms meeting any one of the following conditions that require a variance: · Less than the required square footage INFORMATION NEEDED BY THE END OF THE FIRST DAY OF SURVEY 34. Provide each surveyor with access to all resident electronic health records – do not exclude any information that should be a part of the resident's medical record. Provide specific information on how surveyors can access the EHRs outside of the conference room. Please complete the attached for on page 4 which is titled "Electronic Health Record Information." INFORMATION NEEDED FROM FACILITY WITHIN 24 HOURS OF ENTRANCE 35. Completed Medicare/Medicaid Application (CMS-671). 36. Completed Census and Condition Information (CMS-672) 37. Please complete the attached form on page 3 which is titled "Beneficiary Notice - Residents Discharged Within the Last Six Months"

ENTRANCE CONFERENCE WORKSHEET

DEPARTMENT OF HEALTH AND HUMAN SERVICES CENTERS FOR MEDICARE & MEDICAID SERVICES

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23. Does the facility have an onsite separately certified ESRD unit?

who coordinate(s) services with hospice providers).

Entrance Conference Worksheet

DEPARTMENT OF HEALTH AND HUMAN SERVICES

ENTRANCE CONFERENCE WORKSHEET

Beneficiary Notice - Residents Discharged Within the Last Six Months

Please complete and return this worksheet to the survey team within 24 hours. Please provide a list of residents who were discharged from a Medicare covered Part A stay with benefit days remaining in the past 6 months. Please indicate if the resident was discharged home or remained in the facility. (Note: Exclude beneficiaries who received Medicare Part 8 benefits only, were covered under Medicare Advantage insurance, expired, or were transferred to an acute care

Resident Name	Discharge		arged to:		
Resident Name	Date	Home/Lesser Care	Remained in facility		
1.					
2.					
3.					
4.					
5.					
6.					
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DEPARTMENT OF HEALTH AND HUMAN SERVICES CENTERS FOR MEDICARE & MEDICAID SERVICES

ENTRANCE CONFERENCE WORKSHEET ELECTRONIC HEALTH RECORD (EHR) INFORMATION

Please provide the following information to the survey team before the end of the first day of survey.

in the hard copy if using split F	n where and how surveyors can access the following information in the EHR (of EHR and hard copy system) for the initial pool record review process. Surveyo
	embers have to residents' EHRs in a read-only format.
Example: Medications	EHR: Orders – Reports – Administration Record – eMAR – Confirm date range – Run Report
Example: Hospitalization	EHR: Census (will show in/out of facility)
	MDS (will show discharge MDS)
	Prog Note – View All - Custom – Created Date Range - Enter time period leading up to hospitalization – Save (will show where and why resident wasent)
1. Pressure ulcers	
2. Dialysis	
3. Infections	
4. Nutrition	
5. Falls	
6. ADL status	
Bowel and bladder	
8. Hospitalization	
9. Elopement	
Change of condition	
11. Medications	
12. Diagnoses	
13. PASARR	
Advance directives	
15. Hospice	

Please provide name and contact information for	· IT an	ıd back-up IT	for questions:
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IT Name and Contact Info: _

Back-up IT Name and Contact Info:

1/2018

Matrix for Providers

lesident Name	Resident Room Number	Date of Admission if Admitted within the Past 30 Days	N Alzheimer's / Dementia		Medications: Insulin (I), Anticoagulant (AC), Antibiotic (ABX), Diuretic (B), Opioid (O), Hypnotic (H), Antianxiety (AA), Antipsychotic (AP), Antidepressant (AD), Respiratory (RESP)	Pressure Uler(s) (highest stage I, II, III, IV, U, S), Facility Acquired (FA)	ο Worsened Pressure Ulcer(s) (any stage)	Excessive Weight Loss Without Prescribed Weight Loss Program	α Tube Feeding: Enteral (E) or Parenteral (P)	ه Dehydration	D Physical Restraints	Fall (f), Fall with Injury (FI), or Fall w/Major Injury (FMI)	7 Indwelling Catheter	Dialysis: Peritoneal (P), Hemo (H), in facility (F) or offsite (O)	Hospice	ក្នា End of Life Care / Comfort Care / Palliative Care	9 Tracheostomy	Ventilator	B Transmission-Based Precautions	ال Intravenous therapy	N Infections (M,WI, P, TB, VH, C, UTI)	2
tesident Name		1	2	3	4	5	6	7	8	9	10		12		14	15	16	17	18	19	20	2
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- Initial Pool Process
 - Sample size is determined by the facility census
 - 70% of the total sample is MDS pre-selected residents and 30% of the total sample is selected onsite by the survey team
 - Maximum sample size is 35 residents for larger facilities

- Initial Pool Process
 - First 8-10 hours onsite primarily spent completing initial pool process
 - Surveyors screen all residents in facility and narrow down to an initial pool of about 8 residents per surveyor
 - Surveyors complete an observation, interview (if appropriate) and limited record review for the initial pool residents to help identify those residents who should be in the sample

- Sample Selection
 - After completing the initial pool process, survey team chooses residents from initial pool to include in the sample based on concerns identified from the interview, observation and/or limited record review, and consideration of resident-specific data

- Investigation
 - After selecting the sample, the team spends the rest of the survey investigating all concerns that required further investigation for every resident in the sample. Facility task and closed record investigation are also conducted (although dining is observed the first day)
 - When investigations are complete, the team makes citation, severity and scope decisions for every tag identified by each surveyor.

- Facility Tasks to be Completed with all Surveys
 - Dining Observation
 - Infection Control
 - Beneficiary Protection Notification Review
 - Kitchen

- Medication Administration and Storage
- Resident Council Meeting
- Sufficient and Competent Nurse Staffing
- QAA/QAPI

- Critical Element Pathways
 - Pathways provide guidance to surveyors during the investigation process to determine compliance with the LTC Requirements of Participation. (NOTE: LTC Survey Pathways (total of 41) can be accessed via the following CMS website:

https://www.cms.gov/Medicare/Provider-Enrollmentand-Certification/GuidanceforLawsAndRegulations/ Nursing-Homes.html)

STEPS TO PREPARING FOR A SURVEY

- Understand new LTC survey process
- Review LTC Final Rule (effective 11/28/16) and revised interpretative guidance under Appendix PP of the State Operations Manual (effective 11/28/17)
- Ensure policies/procedures comply with LTC Final Rule
- Educate/Train facility staff regarding policies/procedures
- Train staff on what to expect during a survey

STEPS TO PREPARING FOR A SURVEY

- Conduct Mock Surveys
 - Facility staff vs. outside consultant
 - Utilization of Entrance Conference Worksheet, Facility Matrix and Critical Element Pathways as tools to assess compliance with LTC Final Rule and identify any systems, procedures and/or processes of care that need improvements
 - Address any compliance issues

HELPFUL LINKS:

New Survey Process - https://www.cms.gov/Medicare/Provider-Enrollment-and-Certification/GuidanceforLawsAndRegulations/Nursing-Homes.html

Revised F Tags - https://www.cms.gov/Medicare/Provider-Enrollment-and-Certification/GuidanceforLawsAndRegulations/Downloads/List-of-Revised-FTags.pdf

Appendix PP of SOM - https://www.cms.gov/Medicare/Provider-Enrollment-and-Certification/GuidanceforLawsAndRegulations/Downloads/
https://www.cms.gov/Medicare/Provider-Enrollment-and-Certification/GuidanceforLawsAndRegulations/Downloads/
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Responding to an Adverse Survey/Licensure Action

DOH SURVEY/ENFORCEMENT TRENDS FOR 2018

- Total Surveys 4,716
- Most Frequently Cited Tags
 - F684 (Quality of Care)
 - F689 (Fee of Accident Hazards/Supervision/Devices)
 - F88o (Infection Control)
 - F812 (Food Procurement, Store, Prepare, Serve)
 - F842 (Resident Records)
- Sanctions Issued
 - Provisional I License 3
 - Civil Penalty 184

Source of Information: PA Department of Health

SURVEY RESULTS IN DOH CITING DEFICIENCIES

- Statement of Deficiencies (2567)
- Plan of Correction
 - Required elements
 - Disclaimer language
- Potential for Imposition of Sanctions/Remedies

CRITERIA FOR PAST NONCOMPLIANCE

- All of the following criteria must be met for a survey team to cite past noncompliance with a specific survey data tag (i.e. Ftag or Ktag):
 - 1) The facility was not in compliance with the specific regulatory requirement(s) at the time the situation occurred;
 - 2) The noncompliance occurred after the exit date of the last standard (recertification) survey and before the survey (standard, complaint, or revisit) currently being conducted; and
 - 3) There is sufficient evidence that the facility corrected the noncompliance and is in substantial compliance at the time of the current survey for the specific regulatory requirements.

CRITERIA FOR PAST NONCOMPLIANCE (CONT'D)

- Strategies to evidence past noncompliance
 - Once deficient practice is identified, immediately develop and implement a plan of correction
 - QAPI committee to conduct audits
 - Documentation of corrective measures implemented (e.g. staff training, new/revised policies, ongoing monitoring via audits)
 - Documentation to evidence past noncompliance should be readily available to provide to surveyors if questioned about deficient practice

APPEAL OPTIONS

- IDR
- State IIDR
- Federal IIDR
- DOH Appeal
- CMS Appeal
- DAB Appeal
- Federal Court

INFORMAL DISPUTE RESOLUTION ("IDR")

- <u>Generally</u> The Federal Certification Survey Process provides an informal process to dispute survey findings with the State survey agencies. 42 C.F.R. §488.331.
- <u>Purpose</u> To challenge one or more deficiencies on the CMS-2567 that the facility believes was cited in error.
- <u>Timeline</u> Must submit IDR within the same 10-calendar day period the facility has for submitting an acceptable Plan of Correction.
- Other Failure to complete the IDR timely will not delay the effective date of any enforcement action against the facility.

IDR PROCESS

- Facilities <u>may not</u> use the IDR process to challenge:
 - Scope and severity (unless substandard quality of care or immediate jeopardy)
 - Remedy(ies) imposed by the enforcing agency
 - Failure of the survey team to comply with a requirement of the survey process
 - Alleged inconsistency of the survey team in citing one or more deficiencies among facilities; or the
 - Alleged inadequacy or inaccuracy of the IDR process

IDR PROCESS

- Documentation to support IDR
- IDR submitted to Department of Health for review
 - Decision of DOH final no appeal of final decision
- If IDR results in elimination of one or more deficiencies, the following applies:
 - Facility will receive a "clean" (new) CMS-2567
 - Any enforcement action imposed solely as a result of one or more deficiencies will be rescinded.

STATE INDEPENDENT INFORMAL DISPUTE RESOLUTION ("STATE IIDR")

- Pennsylvania's Long-Term Care Nursing Facility Independent Dispute Resolution Act (Effective 4/20/2012)
 - Establishes an independent informal review process for long-term care nursing facilities to dispute state and federal survey deficiencies
 - Quality Insights of Pennsylvania conducts the State IIDR process
 - State IIDR process conducted on a fee-for-service basis

STATE IIDR CON'T.

- Timeline State IIDR must be submitted within the same 10 calendar days that facility has to submit the POC
- To request a State IIDR, the nursing facility must submit:
 - Written IIDR request that identifies the deficiencies disputed and the reasons for the IIDR request
 - Supporting documentation
 - Copy of 2567
 - Indicate type of review requested: Desk review, telephone review or in-person review

STATE IIDR CON'T.

- QIP reviews the IIDR/supporting documentation and submits a written recommendation to the facility, with a copy to DOH, within 45 days of receipt of the IIDR request.
- If QIP sustains the deficiency, then QIP's written determination shall include the rationale for its decision and provide recommended action that the facility can implement to achieve compliance.
- If QIP reverses the deficiency and DOH disagrees,
 DOH has authority to nullify QIP's decision.

FEDERAL INDEPENDENT INFORMAL DISPUTE RESOLUTION ("FEDERAL IIDR")

- Federal IIDR applicable if:
 - The Centers for Medicare and Medicaid Services ("CMS") imposes civil money penalties against the nursing facility; and
 - The penalties are subject to being collected and placed in an escrow account pending a final administrative decision.
- CMS may collect and place imposed civil money penalties in an escrow account on whichever of the following occurs first:
 - The date on which the IIDR process is completed, or
 - The date which is 90 calendar days after the date of the notice of imposition of the civil money penalty

NOTE: If a facility utilizes the IDR or State IIDR process to challenge the survey findings, the facility cannot also utilize the Federal IIDR process for the same survey unless the IDR or State IIDR process (whichever is applicable) was completed prior to the imposition of the civil money penalty.

• Timeline:

- A request for a Federal IIDR must be submitted within 10 calendar days of the receipt of the letter from CMS regarding the imposition of the civil money penalties.
- The Federal IIDR shall be completed within 60 calendar days of a facility's request.

(Note: The Federal IIDR is deemed completed when a final decision from the IIDR process has been made, a written record has been generated and the State survey agency has sent written notice of this decision to the facility. The IIDR process is also considered to be completed if a facility does not timely request or chooses not to participate in the IIDR process.)

- During the Federal IIDR process, a facility may not challenge other aspects of the survey process, such as:
 - Scope or severity (unless substandard quality of care or immediate jeopardy)
 - Remedy(ies) imposed
 - Alleged failure of the survey team to comply with a requirement of the survey process
 - Alleged inconsistency of the survey team in citing deficiencies among other facilities; or the
 - Alleged inadequacy or inaccuracy of the IDR or IIDR process

- Request for Federal IIDR must include:
 - Copy of CMS letter indicating facility is eligible for an IIDR review
 - Written IIDR request that identifies the deficiencies disputed and reasons for the IIDR request
 - Supporting documentation
 - Names and contact information for residents involved in the deficiencies for which the facility seeks an IIDR review or the appropriate resident representative(s)

- Opportunity for Resident or Resident's Representative to comment:
 - Once a facility requests a Federal IIDR, the State must notify the involved resident or resident representative, as well as the State's long-term care ombudsman, that they have an opportunity to submit written comment

- The notice to the resident/resident's representative, at a minimum, must include:
 - A brief description of the findings of noncompliance for which the facility is requesting the IIDR, a statement about the CMP imposed based on those findings, and reference to the relevant survey date
 - Contact information for the State survey agency, or the approved IIDR entity or person regarding when, where and how potential commenters must submit their comments
 - A designated contact person to answer questions/concerns
 - For residents and/or resident's representatives, contact information for the State's long-term care ombudsman.

- Written Record re: Federal IIDR
 - The IIDR entity or person must generate a written record as soon as practicable but no later than within 10 calendar days of completing its review
 - Written record shall include:
 - List of each deficiency or survey findings that was disputed
 - A summary of the IIDR recommendation for each deficiency or finding at issue and the justification for that result
 - Documents submitted by the facility to dispute a deficiency
 - Any comments submitted by the State long-term care ombudsman and/or residents or resident representatives

- Federal IIDR Recommendation and Final Decision
 - Upon receipt of the IIDR written record, the State Survey Agency ("SSA") will review the IIDR recommendations and:
 - If SSA agrees with IIDR recommendations and no changes will be made to the disputed survey findings, the SSA will send written notice of the final decision to the facility within 10 calendar days of receiving the written record from the IIDR entity/person
 - If SSA disagrees with one or more of the recommendations of the IIDR entity/person, the complete written record will be sent to the applicable CMS Regional Office for review and final decision. SSA will then send written notice of final decision to the facility within 10 calendar days of receiving CMS' final decision.

- Federal IIDR Recommendation and Final Decision con't.
 - If SSA agrees with IIDR recommendation(s) or has received a final decision from the CMS Regional Office and changes will need to be made to the disputed survey findings, the SSA will, within 10 calendar days of receiving the written record:
 - Change deficiency(ies) citation content findings as recommended
 - Adjust scope and severity assessments if warranted by CMS policy
 - Annotate deficiency(ies) citations as "deleted" or "amended" where appropriate
 - Have a SSA manager/supervisor sign and date revised CMS-2567
 - Promptly recommend to CMS that any enforcement action(s) imposed solely because of deleted or altered deficiency citations be reviewed, changed or rescinded as appropriate; and
 - Provide written notice of the final decision to the facility

OVERVIEW

IDR	State IIDR	Federal IIDR
Submitted within same 10 calendar days that facility has to submit POC	Submitted within same 10 calendar days that facility has to submit POC	Submitted within 10 calendar days of the receipt of the CMS letter imposing CMP's
No Fee	Fee-for-Service basis	No Fee
Can only dispute federal deficiencies	Can dispute state and federal deficiencies	Can only dispute federal deficiencies
NO NOTICE to and NO OPPORTUNITY for comment by resident/resident's representative	NO NOTICE to and NO OPPORTUNITY for comment by resident/resident's representative	NOTICE to and OPPORTUNITY for comment by resident/resident's representative
DOH Reviews IDR	Quality Insights of PA reviews State IIDR but DOH is final decision- maker	Independent entity within the DOH reviews IIDR, but if SSA disagrees, CMS is final decision-maker

DOH SUMMARY OF IDR/IIDR RESULTS FOR 2018

• IDR

- 129 Tags disputed
- 43% deleted (55)
- 13% revised (17)

State IIDR

- 17 Tags disputed
- 18% deleted (3)
- 0% revised (0)

Federal IIDR

- 40 Tags disputed
- 0% deleted (0)
- 10% revised (4)

Source of Information: PA Department of Health

DOH APPEAL

- Possible Sanctions:
 - CMP
 - Provisional License
- Appeal of Adverse State Orders
 - File appeal within 30 days of the date of mailing of the Order
 - Appeal of sanction does not act as an automatic supersedeas
 - Must specifically deny the allegations

DOH APPEAL CON'T.

- Appeal of Adverse State Orders (continued)
 - Appeal filed with Health Policy Board
 - Hearing Officer to conduct hearing
 - Practical considerations
 - Possible admissions?
 - Probability of success

- Possible Sanctions
 - CMP
 - Denial of Payment for New Admissions or All Individuals
 - Loss of NATCEP
 - Termination

The facility must appeal within 60 days of receipt of notice of imposition of remedies from CMS. Procedural elements of the appeal process are as follows:

- 1. Notice of Appeal and request for hearing
- 2. Pre-hearing Procedural Order
 - a. Case readiness report
 - b. Document and witness exchange
 - c. Must identify evidence in exchange

Appeal Process con't.

- 3. Scheduling of hearing
 - a. CMS Motions to Dismiss
 - b. Timing
- 4. Hearing before Administrative Law Judge ("ALJ")
 - a. preparation clinical documentation
 - b. physical evidence
 - c. witnesses, identification of expert witnesses
 - d. oral and written summation
 - e. use of hearsay
 - f. burden of proof

Appeal Process con't.

- 5. Decision of ALJ
- 6. DAB Appeal
- 7. Specificity of Appeal. In order to preserve factual issues, appeals should be specific, including which survey and Tag numbers are being contested. The specific grounds for the dispute should be included and explanations of why the conclusions are incorrect. The focus should be on the alleged deficient practice in comparison to the regulatory requirement. Issues of timing, dates, chronological order should be noted.

What is Subject to Appeal

- 1. Only actual remedies not deficiencies alone
- 2. Severity and scope if related to IJ, Substandard Quality of Care, Loss of nurse aide training
- 3. Cannot appeal proposed or withdrawn remedies

Appeal Considerations

- 1. Nature of proposed remedy
 - a. Immediate Jeopardy
 - b. Resident death, abuse, serious injury
 - c. Second consecutive S/S "G"
 - d. Second revisit with any deficiencies (including new deficiencies) and 6 month mandatory termination date is approaching
 - e. Termination proposed.

Appeal Considerations con't.

- 2. Waiver of appeal in exchange for 35% discount on Civil Monetary Penalty
- 3. Can you win on merits?
- 4. Cost

CMS Proposed Rule Regarding Changes to Certain Survey, Certification and Enforcement Procedures

- CMS Proposed Rule (Requirements for Long-Term Care Facilities: Regulatory Provisions to Promote Efficiency and Transparency) – Issued July 16, 2019
- Proposed Changes to Survey, Certification & Enforcement Procedures
 - CMS proposes to revise §488.331(b)(1) (regarding informal dispute resolution (IDR)) by adding language to specify that the IDR process shall be completed within 60 days of the facility's request to dispute the survey findings if the request by the facility is timely.
 - CMS proposes to revise §488.331(b)(2) to specify that survey results are not to be uploaded into CASPER before the resolution of the IDR or independent informal dispute resolution (IIDR) processes.
 - CMS seeks to add new language to §488.431(a)(2) to specify that the facility must receive written notification of the results of the IIDR, including the rationale for the final decision.

CMS Proposed Rule Regarding Changes to Certain Survey, Certification and Enforcement Procedures (cont'd)

- CMS proposes to add language to §488.431(a)(4)(i) to clarify that, in order to be approved to conduct a federal IIDR, a component of an umbrella state agency must have specific understanding of Medicare and Medicaid program requirements.
- CMS proposes to revise §488.436(a) by eliminating the requirement for facilities to file a written waiver of the hearing, and instead, including language to state that the facility is deemed to have waived its rights to a hearing if the time period for requesting a hearing has expired and CMS has not received a timely request for a hearing. The accompanying 35% reduction of the civil money penalty would remain. So, if a provider fails to timely submit a request for a hearing, then such provider would be deemed to have waived their right to a hearing and would automatically be entitled to the 35% reduction in the CMP.

Tanya Daniels Harris, Esq.

LATSHA DAVIS & MARSHALL



717-620-2424

tharris@ldylaw.com

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