CORPORATE INTEGRITY AGREEMENT
BETWEEN THE
OFFICE OF INSPECTOR GENERAL
OF THE
DEPARTMENT OF HEALTH AND HUMAN SERVICES
AND
U.S. PHYSICAL THERAPY, INC.
AND
OLD TOWNE PHYSICAL THERAPY LIMITED PARTNERSHIP

I. PREAMBLE

U.S. Physical Therapy, Inc. and Old Towne Physical Therapy Limited Partnership (Old Towne) hereby enter into this Corporate Integrity Agreement (CIA) with the Office of Inspector General (OIG) of the United States Department of Health and Human Services (HHS) to promote compliance with the statutes, regulations, and written directives of Medicare, Medicaid, and all other Federal health care programs (as defined in 42 U.S.C. § 1320a-7b(f)) (Federal health care program requirements). For purposes of this CIA, the term “USPh” shall mean U.S. Physical Therapy, Inc. and its subsidiaries, as well as any outpatient physical or occupational therapy clinic in which U.S. Physical Therapy, Inc. has a direct or indirect ownership or control interest (as defined in 42 U.S.C. § 1320a-3(a)(3)), excluding rehabilitation agencies and including Old Towne. Contemporaneously with this CIA, Old Towne is entering into a Settlement Agreement with the United States.

USPh represents that, prior to this CIA, USPh voluntarily established a Compliance Program that provides for a Compliance Officer, various compliance committees, maintenance of various policies and procedures, a compliance training and education program, a confidential disclosure reporting hotline, and auditing and monitoring activities aimed at ensuring USPh’s participation in the Federal health care programs conforms to all Federal and State laws and Federal health care program requirements. USPh shall continue its Compliance Program throughout the term of this CIA and shall do so in accordance with the terms set forth below. USPh may modify its Compliance Program, as appropriate, but at a minimum, USPh shall ensure that during the term of this CIA, it shall comply with the obligations set forth herein.
II. TERM AND SCOPE OF THE CIA

A. The period of the compliance obligations assumed by USPh under this CIA shall be five years from the effective date of this CIA. The “Effective Date” shall be the date on which the final signatory of this CIA executes this CIA. Each one-year period, beginning with the one-year period following the Effective Date, shall be referred to as a “Reporting Period.”

B. Sections VII, X, and XI shall expire no later than 120 days after OIG’s receipt of: (1) USPh’s final annual report; or (2) any additional materials submitted by USPh pursuant to OIG’s request, whichever is later.

C. The scope of this CIA shall be governed by the following definitions:

1. “Clinic” shall mean a particular USPh location where outpatient physical or occupational therapy services are provided.

2. “USPh Corporate Operations” shall mean the operations conducted by USPh that support each of the Clinics in which U.S. Physical Therapy, Inc. has a direct or indirect ownership or control interest. “USPh Corporate Operations” includes, but is not limited to, compliance, legal, business development, management, billing, and contracting functions.

3. “Old Towne” shall mean any Clinic controlled by Old Towne Physical Therapy Limited Partnership, and includes, but is not limited to, the three Old Towne Clinics located in Lewes, Millsboro, and Milford, Delaware.

4. “Covered Persons” includes:

   a. all owners who are natural persons (other than shareholders who: (1) have an ownership interest of less than 5% and (2) acquired the ownership interest through public trading), officers, and directors of U.S. Physical Therapy, Inc. or Old Towne;

   b. all employees of USPh who are engaged in or have responsibilities relating to the USPh Corporate Operations or Old Towne;
c. all employees, contractors, subcontractors, agents, and other persons who provide patient care items or services or who perform billing or coding functions on behalf of Old Towne, excluding vendors whose sole connection with Old Towne is selling or otherwise providing medical supplies or equipment to Old Towne and who do not bill the Federal health care programs for such medical supplies or equipment; and

d. for purposes of the requirements set forth in Section III.B.1 (Code of Conduct), Section III.B.2 (Policies and Procedures), Section III.F (Disclosure Program), and Section III.G (Ineligible Persons), all employees, contractors, subcontractors, agents, and other persons who provide patient care items or services or who perform billing or coding functions on behalf of USPh, excluding vendors whose sole connection with USPh is selling or otherwise providing medical supplies or equipment to USPh and who do not bill the Federal health care programs for such medical supplies or equipment.

Notwithstanding the above, this term does not include part-time or per diem employees, contractors, subcontractors, agents, and other persons who are not reasonably expected to work more than 160 hours during a Reporting Period, except that any such individuals shall become “Covered Persons” at the point when they work more than 160 hours during a Reporting Period.

5. “Relevant Covered Persons” includes all Covered Persons who:

a. are involved directly or indirectly in the delivery of outpatient physical or occupational therapy services or otherwise participate in the delivery of health care items or services payable by the Federal health care programs; or

b. perform billing or coding functions.

III. CORPORATE INTEGRITY OBLIGATIONS

USPh shall establish and maintain a Compliance Program that includes the following elements:
A. Compliance Officer and Committee

1. Compliance Officer and Compliance Contact. USP has appointed a Covered Person to serve as its Chief Compliance Officer (Compliance Officer) and shall maintain a Compliance Officer for the term of this CIA. The Compliance Officer shall be a member of senior management of USP, shall report directly to the Chief Executive Officer of USP, and shall not be or be subordinate to the General Counsel or Chief Financial Officer or have any responsibilities that involve acting in any capacity as legal counsel or supervising legal counsel functions for USP. The Compliance Officer shall be responsible for, without limitation:

   a. developing and implementing policies, procedures, and practices designed to ensure compliance with the requirements set forth in this CIA and with Federal health care program requirements;

   b. making periodic (at least quarterly) reports regarding compliance matters directly to the Board of Directors of USP, and shall be authorized to report on such matters to the Board of Directors at any time. Written documentation of the Compliance Officer’s reports to the Board of Directors shall be made available to OIG upon request; and

   c. monitoring the day-to-day compliance activities engaged in by USP as well as for any reporting obligations created under this CIA.

Any noncompliance job responsibilities of the Compliance Officer shall be limited and must not interfere with the Compliance Officer’s ability to perform the duties outlined in this CIA.

Within 90 days after the Effective Date, Old Towne shall appoint a Covered Person to serve as its Compliance Contact and shall maintain a Compliance Contact for the term of this CIA. Old Towne’s Compliance Contact shall be responsible for, without limitation:

   a. making periodic (at least monthly) reports regarding compliance matters at Old Towne to USP’s Compliance Officer; and
b. monitoring the day-to-day compliance activities engaged in by Old Towne.

USPh shall report to OIG, in writing, any changes in the identity or position description of the Compliance Officer or Old Towne’s Compliance Contact, or any actions or changes that would affect the Compliance Officer’s or Old Towne’s Compliance Contact’s ability to perform the duties necessary to meet the obligations in this CIA, within five days after such a change.

2. **Compliance Committee.** USPh has appointed a Compliance Committee and shall maintain a Compliance Committee for the term of this CIA. The Compliance Committee shall, at a minimum, include the Compliance Officer, a member of USPh’s Board of Directors, and other members of senior management necessary to meet the requirements of this CIA (e.g., senior executives of relevant departments, such as billing, clinical, therapy, human resources, medical records, audit, and operations). The Compliance Officer shall chair the Compliance Committee and the Committee shall support the Compliance Officer in fulfilling his/her responsibilities (e.g., shall assist in the analysis of USPh’s risk areas and shall oversee monitoring of internal and external audits and investigations). The Compliance Committee shall meet at least quarterly. The minutes of the Compliance Committee meetings shall be made available to OIG upon request.

USPh shall report to OIG, in writing, any changes in the composition of the Compliance Committee, or any actions or changes that would affect the Compliance Committee’s ability to perform the duties necessary to meet the obligations in this CIA, within 15 days after such a change.

3. **Board of Directors Compliance Obligations.** The Board of Directors (or a committee of the Board) of USPh (Board) shall be responsible for the review and oversight of matters related to compliance with Federal health care program requirements and the obligations of this CIA. The Board must include independent (i.e., non-executive) members.

The Board shall, at a minimum, be responsible for the following:

a. meeting at least quarterly to review and oversee USPh’s Compliance Program, including but not limited to the performance of the Compliance Officer and Compliance Committee;
b. submitting to the OIG a description of the documents and other materials it reviewed, as well as any additional steps taken, in its oversight of the Compliance Program and in support of making the resolution below during each Reporting Period; and

c. for each Reporting Period of the CIA, adopting a resolution, signed by each member of the Board summarizing its review and oversight of USPh’s compliance with Federal health care program requirements and the obligations of this CIA.

At minimum, the resolution shall include the following language:

“The Board of Directors has made a reasonable inquiry into the operations of USPh’s Compliance Program including the performance of the Compliance Officer and the Compliance Committee. Based on its inquiry and review, the Board has concluded that, to the best of its knowledge, USPh has implemented an effective Compliance Program to meet Federal health care program requirements and the obligations of the Corporate Integrity Agreement.”

If the Board is unable to provide such a conclusion in the resolution, the Board shall include in the resolution a written explanation of the reasons why it is unable to provide the conclusion and the steps it is taking to implement an effective Compliance Program at USPh.

USPh shall report to OIG, in writing, any changes in the composition of the Board, or any actions or changes that would affect the Board’s ability to perform the duties necessary to meet the obligations in this CIA, within 15 days after such a change.

B. Written Standards

1. **Code of Conduct.** Within 90 days after the Effective Date, USPh shall develop, implement, and distribute a written Code of Conduct to all Covered Persons. USPh shall make the performance of job responsibilities in a manner consistent with the Code of Conduct an element in evaluating the performance of all employees. The Code of Conduct shall, at a minimum, set forth:

a. USPh’s commitment to full compliance with all Federal health care program requirements, including its commitment
to prepare and submit accurate claims consistent with such requirements;

b. USPh’s requirement that all of its Covered Persons shall be expected to comply with all Federal health care program requirements and with USPh’s own Policies and Procedures;

c. the requirement that all of USPh’s Covered Persons shall be expected to report to the Compliance Officer, or other appropriate individual designated by USPh, suspected violations of any Federal health care program requirements or of USPh’s own Policies and Procedures;

d. the possible consequences to both USPh and Covered Persons of failure to comply with Federal health care program requirements and with USPh’s own Policies and Procedures, as well as the failure to report such noncompliance; and

e. the right of all individuals to use the Disclosure Program described in Section III.E, and USPh’s commitment to nonretaliation and to maintain, as appropriate, confidentiality and anonymity with respect to such disclosures.

Within 90 days after the Effective Date, USPh shall distribute the Code of Conduct to all Covered Persons. USPh shall review the Code of Conduct at least annually to determine if revisions are appropriate and shall make any necessary revisions based on such review. The Code of Conduct shall be distributed at least annually to all Covered Persons.

2. **Policies and Procedures.** Within 90 days after the Effective Date, USPh shall develop and implement written Policies and Procedures regarding the operation of its Compliance Program, including the Compliance Program requirements outlined in this CIA and USPh’s compliance with Federal health care program requirements. USPh shall enforce and comply with its Policies and Procedures and shall make such compliance an element of evaluating the performance of all employees.

USPh shall ensure that, at a minimum, its Policies and Procedures address:

a. the Compliance Program requirements outlined in this CIA;
b. the subjects relating to the Code of Conduct identified in Section III.B.1; and

c. the rules and regulations concerning the provision of physical and occupational therapy services to Federal health care program beneficiaries, including but not limited to:

   i. Medicare conditions of payment for outpatient physical and occupational therapy services, including the supervision requirements applicable to services furnished by physical therapist assistants (PTAs);

   ii. the scope of practice and supervision standards applicable to PTAs under the laws of each state in which USPh Clinics are located; and

   iii. the importance of complete and accurate documentation of medical records regarding the delivery of physical and occupational therapy services.

Within 90 days after the Effective Date, the Policies and Procedures shall be distributed to all Covered Persons. Appropriate and knowledgeable staff shall be available to explain the Policies and Procedures.

At least annually (and more frequently, if appropriate), USPh shall assess and update, as necessary, the Policies and Procedures. Within 30 days after the effective date of any revisions or addition of new Policies and Procedures, a description of the revisions shall be communicated to all affected Covered Persons and any revised or new Policies and Procedures shall be made available to all Covered Persons.

C. Training and Education

1. Training Plan. Within 90 days after the Effective Date, USPh shall develop a written plan (Training Plan) that outlines the steps USPh will take to ensure that: (a) all Covered Persons receive adequate training regarding USPh’s CIA requirements and Compliance Program, including the Code of Conduct; and (b) all Relevant Covered Persons receive adequate training regarding: (i) the Federal health care program requirements regarding the accurate coding and submission of claims for
outpatient physical and occupational therapy and other health care items or services rendered at USP Clinics; (ii) policies, procedures, and other requirements applicable to the provision of outpatient physical and occupational therapy and other health care items or services rendered at USP Clinics, as well as the proper medical documentation of such items or services; (iii) the personal obligation of each individual involved in the claims submission process to ensure that such claims are accurate; (iv) applicable reimbursement statutes, regulations, and program requirements and directives; (v) the legal sanctions for violations of the Federal health care program requirements; and (vi) examples of proper and improper claims submission practices.

The Training Plan shall include information regarding the training topics, the categories of Covered Persons and Relevant Covered Persons required to attend each training session, the length of the training, the schedule for training, and the format of the training. Within 30 days of OIG’s receipt of USP’s Training Plan, OIG will notify USP of any comments or objections to the Training Plan. Absent notification by the OIG that the Training Plan is unacceptable, USP may implement its Training Plan. USP shall furnish training to its Covered Persons and Relevant Covered Persons pursuant to the Training Plan during each Reporting Period.

2. **Board Member Training.** Within 90 days after the Effective Date, USP shall provide at least two hours of training to each member of the Board of Directors. This training shall address USP’s CIA requirements and Compliance Program (including the Code of Conduct), the corporate governance responsibilities of board members, and the responsibilities of board members with respect to review and oversight of the Compliance Program. Specifically, the training shall address the unique responsibilities of health care Board members, including the risks, oversight areas, and strategic approaches to conducting oversight of a health care entity. This training may be conducted by an outside compliance expert hired by the Board and should include a discussion of the OIG’s guidance on Board member responsibilities.

New members of the Board of Directors shall receive the Board Member Training described above within 30 days after becoming a member or within 90 days after the Effective Date, whichever is later.

3. **Certification.** Each individual who is required to attend training shall certify, in writing or in electronic form, that he or she has received the required training. The certification shall specify the type of training received and the date received. The Compliance Officer (or designee) shall retain the certifications, along with all course materials.
4. **Qualifications of Trainer.** Persons providing or preparing the training shall be knowledgeable about the subject area.

5. **Update of Training Plan.** USP shall review the Training Plan annually, and, where appropriate, update the Training Plan to reflect changes in Federal health care program requirements, any issues discovered during internal audits or the Claims Review, and any other relevant information. Any updates to the Training Plan must be reviewed and approved by OIG prior to the implementation of the revised Training Plan. Within 30 days of OIG’s receipt of any updates or revisions to USP’s Training Plan, OIG will notify USP of any comments or objections to the revised Training Plan. Absent notification from OIG that the revised Training Plan is unacceptable, USP may implement the revised Training Plan.

6. **Computer-Based Training.** USP may provide the training required under this CIA through appropriate computer-based training approaches. If USP chooses to provide computer-based training, it shall make available appropriately qualified and knowledgeable staff or trainers to answer questions or provide additional information to the individuals receiving such training.

D. **Review Procedures**

1. **General Description**

   a. **Engagement of Independent Review Organization.** Within 90 days after the Effective Date, USP shall engage an entity (or entities), such as an accounting, auditing, or consulting firm (hereinafter “Independent Review Organization” or “IRO”), to perform the reviews listed in this Section III.D. The applicable requirements relating to the IRO are outlined in Appendix A to this CIA, which is incorporated by reference.

   b. **Retention of Records.** The IRO and USP shall retain and make available to OIG, upon request, all work papers, supporting documentation, correspondence, and draft reports (those exchanged between the IRO and USP) related to the reviews.

2. **Outpatient Physical and Occupational Therapy Claims Review.** The IRO shall review USP’s coding, billing, and outpatient physical and occupational therapy claims submission to the Medicare program and the reimbursement received
(Claims Review) and shall prepare a Claims Review Report, as outlined in Appendix B to this CIA, which is incorporated by reference.

3. **Validation Review.** In the event OIG has reason to believe that: (a) USPh’s Claims Review fails to conform to the requirements of this CIA; or (b) the IRO’s findings or Claims Review results are inaccurate, OIG may, at its sole discretion, conduct its own review to determine whether the Claims Review complied with the requirements of the CIA and/or the IRO’s findings or Claims Review results are inaccurate (Validation Review). USPh shall pay for the reasonable cost of any such review performed by OIG or any of its designated agents. Any Validation Review of Reports submitted as part of USPh’s final Annual Report shall be initiated no later than one year after USPh’s final submission (as described in Section II) is received by OIG.

Prior to initiating a Validation Review, OIG shall notify USPh of its intent to do so and provide a written explanation of why OIG believes such a review is necessary. To resolve any concerns raised by OIG, USPh may request a meeting with OIG to: (a) discuss the results of any Claims Review submissions or findings; (b) present any additional information to clarify the results of the Claims Review or to correct the inaccuracy of the Claims Review; and/or (c) propose alternatives to the proposed Validation Review. USPh agrees to provide any additional information as may be requested by OIG under this Section III.D.3 in an expedited manner. OIG will attempt in good faith to resolve any Claims Review issues with USPh prior to conducting a Validation Review. However, the final determination as to whether or not to proceed with a Validation Review shall be made at the sole discretion of OIG.

4. **Independence and Objectivity Certification.** The IRO shall include in its report(s) to USPh a certification that the IRO has: (a) evaluated its professional independence and objectivity with respect to the reviews conducted under this Section III.D; and (b) concluded that it is, in fact, independent and objective, in accordance with the requirements specified in Appendix A to this CIA.

E. **Risk Assessment and Internal Review Process**

   Within 90 days after the Effective Date, USPh shall develop and implement a centralized annual risk assessment and internal review process to identify and address risks associated with the submission of claims for items and services furnished to Medicare and Medicaid program beneficiaries. The risk assessment and internal review process should include: (1) a process for identifying and prioritizing risks; (2) developing remediation plans in response to those risks, including internal auditing and monitoring of the identified risk areas; (3) tracking results to assess the effectiveness
of the remediation plans; and (4) conducting a comprehensive audit of all new Clinics that partner with USPh during the first year of the partnership, to include a review of Medicare plan of care certification, documentation, billing, and coding. The risk assessment and internal review process should require compliance, legal, and department leaders, at least annually, to evaluate and identify risks associated with the submission of claims for items and services furnished to Medicare and Medicaid program beneficiaries and develop and implement specific plans to address and mitigate the identified risks. The risk assessment and internal review work plans shall be developed annually. USPh shall implement the risk assessment and internal review work plans and track the implementation of the work plans. USPh shall maintain the risk assessment and internal review process for the term of the CIA. Copies of any internal audit reports developed pursuant to the risk assessment and internal review process shall be made available to OIG upon request.

F. Disclosure Program

USPh has established a Disclosure Program that includes a mechanism (e.g., a toll-free compliance telephone line) to enable individuals to disclose, to the Compliance Officer or some other person who is not in the disclosing individual’s chain of command, any identified issues or questions associated with USPh’s policies, conduct, practices, or procedures with respect to a Federal health care program believed by the individual to be a potential violation of criminal, civil, or administrative law. USPh shall appropriately publicize the existence of the disclosure mechanism (e.g., via periodic e-mails to employees or by posting the information in prominent common areas).

The Disclosure Program shall emphasize a nonretribution, nonretaliation policy, and shall include a reporting mechanism for anonymous communications for which appropriate confidentiality shall be maintained. Upon receipt of a disclosure, the Compliance Officer (or designee) shall gather all relevant information from the disclosing individual. The Compliance Officer (or designee) shall make a preliminary, good faith inquiry into the allegations set forth in every disclosure to ensure that he or she has obtained all of the information necessary to determine whether a further review should be conducted. For any disclosure that is sufficiently specific so that it reasonably: (1) permits a determination of the appropriateness of the alleged improper practice; and (2) provides an opportunity for taking corrective action, USPh shall conduct an internal review of the allegations set forth in the disclosure and ensure that proper follow-up is conducted.

The Compliance Officer (or designee) shall maintain a disclosure log and shall record each disclosure in the disclosure log within 48 hours of receipt of the disclosure.
The disclosure log shall include a summary of each disclosure received (whether anonymous or not), the status of the respective internal reviews, and any corrective action taken in response to the internal reviews.

G. Ineligible Persons

1. Definitions. For purposes of this CIA:

   a. an “Ineligible Person” shall include an individual or entity who:

      i. is currently excluded, debarred, suspended, or otherwise ineligible to participate in the Federal health care programs or in Federal procurement or nonprocurement programs; or

      ii. has been convicted of a criminal offense that falls within the scope of 42 U.S.C. § 1320a-7(a), but has not yet been excluded, debarred, suspended, or otherwise declared ineligible.

   b. “Exclusion Lists” include:

      i. the HHS/OIG List of Excluded Individuals/Entities (available through the Internet at http://www.oig.hhs.gov); and

      ii. the General Services Administration’s System for Award Management (available through the Internet at http://www.sam.gov).

2. Screening Requirements. USPh shall ensure that all prospective and current Covered Persons are not Ineligible Persons, by implementing the following screening requirements.

   a. USPh shall screen all prospective Covered Persons against the Exclusion Lists prior to engaging their services and, as part of the hiring or contracting process, shall require such Covered Persons to disclose whether they are Ineligible Persons.
b. USPh shall screen all Covered Persons against the Exclusion Lists within 90 days after the Effective Date and thereafter shall screen against the LEIE on a monthly basis and screen against SAM on an annual basis.

c. USPh shall implement a policy requiring all Covered Persons to disclose immediately any debarment, exclusion, suspension, or other event that makes that person an Ineligible Person.

Nothing in Section III.G affects USPh’s responsibility to refrain from (and liability for) billing Federal health care programs for items or services furnished, ordered, or prescribed by excluded persons. USPh understands that items or services furnished, ordered, or prescribed by excluded persons are not payable by Federal health care programs and that USPh may be liable for overpayments and/or criminal, civil, and administrative sanctions for employing or contracting with an excluded person regardless of whether USPh meets the requirements of Section III.G.

3. Removal Requirement. If USPh has actual notice that a Covered Person has become an Ineligible Person, USPh shall remove such Covered Person from responsibility for, or involvement with, USPh’s business operations related to the Federal health care programs and shall remove such Covered Person from any position for which the Covered Person’s compensation or the items or services furnished, ordered, or prescribed by the Covered Person are paid in whole or part, directly or indirectly, by Federal health care programs or otherwise with Federal funds at least until such time as the Covered Person is reinstated into participation in the Federal health care programs.

4. Pending Charges and Proposed Exclusions. If USPh has actual notice that a Covered Person has been charged with a criminal offense that falls within the scope of 42 U.S.C. §§ 1320a-7(a), 1320a-7(b)(1)–(3), or is proposed for exclusion during the Covered Person’s employment or contract term, USPh shall take all appropriate actions to ensure that the responsibilities of that Covered Person have not and shall not adversely affect the quality of care rendered to any beneficiary, patient, or resident, or any claims submitted to any Federal health care program.

H. Notification of Government Investigation or Legal Proceedings

Within 30 days after discovery, USPh shall notify OIG, in writing, of any ongoing investigation or legal proceeding known to USPh conducted or brought by a governmental entity or its agents involving an allegation that USPh has committed a
crime or has engaged in fraudulent activities. This notification shall include a description of the allegation, the identity of the investigating or prosecuting agency, and the status of such investigation or legal proceeding. USPh shall also provide written notice to OIG within 30 days after the resolution of the matter, and shall provide OIG with a description of the findings and/or results of the investigation or proceedings, if any.

I. Repayment of Overpayments

1. **Definition of Overpayments.** For purposes of this CIA, an “Overpayment” shall mean the amount of money USPh has received in excess of the amount due and payable under any Federal health care program requirements.

2. **Overpayment Policies and Procedures.** Within 90 days after the Effective Date, USPh shall develop and implement written policies and procedures regarding the identification, quantification, and repayment of Overpayments received from any Federal health care program.

3. **Repayment of Overpayments**

   a. If, at any time, USPh identifies or learns of any Overpayment, USPh shall repay the Overpayment to the appropriate payor (e.g., Medicare contractor) within 60 days after identification of the Overpayment and take remedial steps within 90 days after identification (or such additional time as may be agreed to by the payor) to correct the problem, including preventing the underlying problem and the Overpayment from recurring. If not yet quantified, within 60 days after identification, USPh shall notify the payor of its efforts to quantify the Overpayment amount along with a schedule of when such work is expected to be completed. Notification and repayment to the payor shall be done in accordance with the payor’s policies.

   b. Notwithstanding the above, notification and repayment of any Overpayment amount that routinely is reconciled or adjusted pursuant to policies and procedures established by the payor should be handled in accordance with such policies and procedures.
J. Reportable Events

1. Definition of Reportable Event. For purposes of this CIA, a “Reportable Event” means anything that involves:

   a. a substantial Overpayment;

   b. a matter that a reasonable person would consider a probable violation of criminal, civil, or administrative laws applicable to any Federal health care program for which penalties or exclusion may be authorized;

   c. the employment of or contracting with a Covered Person who is an Ineligible Person as defined by Section III.G.1.a; or

   d. the filing of a bankruptcy petition by USPh.

A Reportable Event may be the result of an isolated event or a series of occurrences.

2. Reporting of Reportable Events. If USPh determines (after a reasonable opportunity to conduct an appropriate review or investigation of the allegations) through any means that there is a Reportable Event, USPh shall notify OIG, in writing, within 30 days after making the determination that the Reportable Event exists.

3. Reportable Events under Section III.J.1.a. For Reportable Events under Section III.J.1.a, the report to OIG shall be made within 30 days of the identification of the Overpayment, and shall include:

   a. a complete description of all details relevant to the Reportable Event, including, at minimum, the types of claims, transactions or other conduct giving rise to the Reportable Event; the period during which the conduct occurred; and the names of entities and individuals believed to be implicated, including an explanation of their roles in the Reportable Event;

   b. the Federal health care programs affected by the Reportable Event;
c. a description of the steps taken by USP\textsuperscript{h} to identify and quantify the Overpayment; and

d. a description of USP\textsuperscript{h}'s actions taken to correct the Reportable Event and prevent it from recurring.

Within 60 days of identification of the Overpayment, USP\textsuperscript{h} shall provide OIG with a copy of the notification and repayment (if quantified) to the payor required in Section III.I.3.

4. \textit{Reportable Events under Section III.J.1.b}. For Reportable Events under Section III.J.1.b, the report to OIG shall include:

a. a complete description of all details relevant to the Reportable Event, including, at a minimum, the types of claims, transactions or other conduct giving rise to the Reportable Event; the period during which the conduct occurred; and the names of entities and individuals believed to be implicated, including an explanation of their roles in the Reportable Event;

b. a statement of the Federal criminal, civil, or administrative laws that are probably violated by the Reportable Event;

c. the Federal health care programs affected by the Reportable Event;

d. a description of USP\textsuperscript{h}'s actions taken to correct the Reportable Event and prevent it from recurring; and

e. if the Reportable Event has resulted in an Overpayment, a description of the steps taken by USP\textsuperscript{h} to identify and quantify the Overpayment.

5. \textit{Reportable Events under Section III.J.1.c}. For Reportable Events under Section III.J.1.c, the report to OIG shall include:

a. the identity of the Ineligible Person and the job duties performed by that individual;
b. the dates of the Ineligible Person’s employment or contractual relationship;

c. a description of the Exclusion Lists screening that USPh completed before and/or during the Ineligible Person’s employment or contract and any flaw or breakdown in the Ineligible Persons screening process that led to the hiring or contracting with the Ineligible Person;

d. a description of how the Reportable Event was discovered; and

e. a description of any corrective action implemented to prevent future employment or contracting with an Ineligible Person.

6. **Reportable Events under Section III.J.1.d.** For Reportable Events under Section III.J.1.d, the report to OIG shall include documentation of the bankruptcy filing and a description of any Federal health care program authorities implicated.

7. **Reportable Events Involving the Stark Law.** Notwithstanding the reporting requirements outlined above, any Reportable Event that involves only a probable violation of section 1877 of the Social Security Act, 42 U.S.C. § 1395nn (the Stark Law) should be submitted by USPh to the Centers for Medicare & Medicaid Services (CMS) through the self-referral disclosure protocol (SRDP), with a copy to OIG. The requirements of Section III.I.3 that require repayment to the payor of any identified Overpayment within 60 days shall not apply to any Overpayment that may result from a probable violation of only the Stark Law that is disclosed to CMS pursuant to the SRDP. If USPh identifies a probable violation of only the Stark Law and repays the applicable Overpayment directly to the CMS contractor, then USPh is not required by this Section III.J to submit the Reportable Event to CMS through the SRDP.

**IV. SUCCESSOR LIABILITY; CHANGES TO BUSINESS UNITS OR LOCATIONS**

**A. Sale of Business, Business Unit, or Location.**

In the event that, after the Effective Date, USPh proposes to sell any or all of its business, business units, or locations (whether through a sale of assets, sale of stock, or other type of transaction) that are subject to this CIA, USPh shall notify OIG of the proposed sale at least 30 days prior to the sale of its business, business unit, or location.
This notification shall include a description of the business, business unit, or location to be sold; a brief description of the terms of the sale; and the name and contact information of the prospective purchaser. This CIA shall be binding on the purchaser of the business, business unit, or location, unless otherwise determined and agreed to in writing by OIG.

B. Change or Closure of Business, Business Unit, or Location

In the event that, after the Effective Date, USPh changes locations or closes a business, business unit, or location related to the furnishing of items or services that may be reimbursed by Federal health care programs, USPh shall notify OIG of this fact as soon as possible, but no later than within 30 days after the date of change or closure of the business, business unit, or location.

C. Purchase or Establishment of New Business, Business Unit, or Location

In the event that, after the Effective Date, USPh purchases or establishes a new business, business unit, or location related to the furnishing of items or services that may be reimbursed by Federal health care programs, USPh shall notify OIG at least 30 days prior to such purchase or the operation of the new business, business unit, or location. This notification shall include the address of the new business, business unit, or location; phone number; fax number; the location’s Medicare and state Medicaid program provider number and/or supplier number(s); and the name and address of each Medicare and state Medicaid program contractor to which USPh currently submits claims. Each new business, business unit, or location and all Covered Persons at each new business, business unit, or location shall be subject to the applicable requirements of this CIA, unless otherwise specified in this CIA or agreed to in writing by OIG.

V. IMPLEMENTATION AND ANNUAL REPORTS

A. Implementation Report

Within 120 days after the Effective Date, USPh shall submit a written report to OIG summarizing the status of its implementation of the requirements of this CIA (Implementation Report). The Implementation Report shall, at a minimum, include:

1. the name, address, phone number, and position description of the Compliance Officer required by Section III.A.1, and a summary of other noncompliance job responsibilities the Compliance Officer may have;
2. the names and positions of the members of the Compliance Committee required by Section III.A.2;

3. the names of the Board members who are responsible for satisfying the Board of Directors compliance obligations described in Section III.A.3;

4. a copy of USPh’s Code of Conduct required by Section III.B.1;

5. a summary of all Policies and Procedures required by Section III.B.2 (copies of the Policies and Procedures shall be made available to OIG upon request);

6. the Training Plan required by Section III.C.1 and a description of the Board of Directors training required by Section III.C.2 (including a summary of the topics covered, the length of the training, and when the training was provided);

7. the following information regarding the IRO(s): (a) identity, address, and phone number; (b) a copy of the engagement letter; (c) information to demonstrate that the IRO has the qualifications outlined in Appendix A to this CIA; (d) a summary and description of any and all current and prior engagements and agreements between USPh and the IRO; and (e) a certification from the IRO regarding its professional independence and objectivity with respect to USPh;

8. a description of the risk assessment and mitigation process required by Section III.E;

9. a description of the Disclosure Program required by Section III.F;

10. a certification that USPh has conducted the screening required by Section III.G regarding Ineligible Persons, or a description of why USPh cannot provide such a certification;

11. a copy of USPh’s policies and procedures regarding the identification, quantification, and repayment of Overpayments required by Section III.I;

12. a list of all of USPh’s locations (including locations and mailing addresses); the corresponding name under which each location is doing business; the corresponding phone numbers and fax numbers; each location’s Medicare and state Medicaid program provider number and/or supplier number(s); and the name and address of each Medicare and state Medicaid program contractor to which USPh currently submits claims;
13. a description of USPh’s and Old Towne’s corporate structures, including identification of any parent and sister companies, subsidiaries, and their respective lines of business, as well as any individual owners of Old Towne; and

14. the certifications required by Section V.C.

B. Annual Reports

USPh shall submit to OIG annually a report with respect to the status of, and findings regarding, USPh’s compliance activities for each of the five Reporting Periods (Annual Report). Each Annual Report shall include, at a minimum:

1. any change in the identity, position description, or other noncompliance job responsibilities of the Compliance Officer; any change in the membership of the Compliance Committee described in Section III.A.2; and any change in the Board members who are responsible for satisfying the Board of Directors compliance obligations described in Section III.A.3;

2. the dates of each report made by the Compliance Officer to the Board (written documentation of such reports shall be made available to OIG upon request);

3. the Board resolution required by Section III.A.3, and a description of the documents and other materials reviewed by the Board, as well as any additional steps taken, in its oversight of the Compliance Program and in support of making the resolution;

4. a summary of any significant changes or amendments to USPh’s Code of Conduct or the Policies and Procedures required by Section III.B and the reasons for such changes (e.g., change in contractor policy);

5. a copy of USPh’s Training Plan developed under Section III.C and the following information regarding each type of training required by the Training Plan: a description of the training, including a summary of the topics covered; the length of sessions, a schedule of training sessions, a general description of the categories of individuals required to complete the training, and the process by which USPh ensures that all designated employees receive appropriate training. A copy of all training materials and the documentation to support this information shall be made available to OIG upon request;

U.S. Physical Therapy, Inc.
Old Towne Physical Therapy Limited Partnership
Corporate Integrity Agreement
6. a complete copy of all reports prepared pursuant to Section III.D, along with a copy of the IRO’s engagement letter;

7. USP$h$’s response to the reports prepared pursuant to Section III.D, along with corrective action plan(s) related to any issues raised by the reports;

8. a summary and description of any and all current and prior engagements and agreements between USP$h$ and the IRO (if different from what was submitted as part of the Implementation Report);

9. a certification from the IRO regarding its professional independence and objectivity with respect to USP$h$;

10. a description of the risk assessment and internal review process required by Section III.E, a summary of any changes to the process, and a description of the reasons for such changes;

11. a copy of USP$h$’s internal review work plans, and a list of all reviews completed during the Reporting Period pursuant to Section III.E;

12. a summary of the disclosures in the disclosure log required by Section III.F that relate to Federal health care programs (the complete disclosure log shall be made available to OIG upon request);

13. a certification that USP$h$ has completed the screening required by Section III.G regarding Ineligible Persons;

14. a summary describing any ongoing investigation or legal proceeding required to have been reported pursuant to Section III.H. The summary shall include a description of the allegation, the identity of the investigating or prosecuting agency, and the status of such investigation or legal proceeding;

15. a description of any changes to the Overpayments policies and procedures required by Section III.I, including the reasons for such changes;

16. a report of the aggregate Overpayments that have been returned to the Federal health care programs. Overpayment amounts shall be broken down into the following categories: outpatient Medicare, Medicaid (report each applicable state separately, if applicable), and other Federal health care programs. Overpayment amounts
that are routinely reconciled or adjusted pursuant to policies and procedures established by the payor do not need to be included in this aggregate Overpayment report;

17. a summary of Reportable Events (as defined in Section III.J) identified during the Reporting Period and the status of any corrective action relating to all such Reportable Events;

18. a summary describing any audits conducted during the applicable Reporting Period by a Medicare or state Medicaid program contractor or any government entity or contractor, involving a review of Federal health care program claims, and USPh’s response/corrective action plan (including information regarding any Federal health care program refunds) relating to the audit findings;

19. a description of all changes to the most recently provided list of USPh’s locations (including addresses) as required by Section V.A.12; the corresponding name under which each location is doing business; the corresponding phone numbers and fax numbers; each location’s Medicare and state Medicaid program provider number(s) and/or supplier number(s); and the name and address of each Medicare and state Medicaid program contractor to which USPh currently submits claims; and

20. the certifications required by Section V.C.

The first Annual Report shall be received by OIG no later than 60 days after the end of the first Reporting Period. Subsequent Annual Reports shall be received by OIG no later than the anniversary date of the due date of the first Annual Report.

C. Certifications

1. Compliance Officer and Chief Executive Officer. The Implementation Report and each Annual Report shall include a certification by the Compliance Officer and Chief Executive Officer that:

   a. to the best of his or her knowledge, except as otherwise described in the report, USPh is in compliance with all of the requirements of this CIA; and

   b. he or she has reviewed the report and has made reasonable inquiry regarding its content and believes that the information in the report is accurate and truthful.
2. **Chief Financial Officer.** The first Annual Report shall include a certification by the Chief Financial Officer that, to the best of his or her knowledge, Old Towne has complied with its obligations under the Settlement Agreement: (a) not to resubmit to any Federal health care program payors any previously denied claims related to the Covered Conduct addressed in the Settlement Agreement, and not to appeal any such denials of claims; (b) not to charge to or otherwise seek payment from federal or state payors for unallowable costs (as defined in the Settlement Agreement); and (c) to identify and adjust any past charges or claims for unallowable costs.

D. **Designation of Information**

USPh shall clearly identify any portions of its submissions that it believes are trade secrets, or information that is commercial or financial and privileged or confidential, and therefore potentially exempt from disclosure under the Freedom of Information Act (FOIA), 5 U.S.C. § 552. USPh shall refrain from identifying any information as exempt from disclosure if that information does not meet the criteria for exemption from disclosure under FOIA.

VI. **NOTIFICATIONS AND SUBMISSION OF REPORTS**

Unless otherwise stated in writing after the Effective Date, all notifications and reports required under this CIA shall be submitted to the following entities:

**OIG:** Administrative and Civil Remedies Branch  
Office of Counsel to the Inspector General  
Office of Inspector General  
U.S. Department of Health and Human Services  
Cohen Building, Room 5527  
330 Independence Avenue, SW  
Washington, DC 20201  
Telephone: (202) 619-2078  
Facsimile: (202) 205-0604
Unless otherwise specified, all notifications and reports required by this CIA may be made by certified mail, overnight mail, hand delivery, or other means, provided that there is proof that such notification was received. For purposes of this requirement, internal facsimile confirmation sheets do not constitute proof of receipt. Upon request by OIG, USPh may be required to provide OIG with an electronic copy of each notification or report required by this CIA in searchable portable document format (pdf), in addition to a paper copy.

VII. OIG INSPECTION, AUDIT, AND REVIEW RIGHTS

In addition to any other rights OIG may have by statute, regulation, or contract, OIG or its duly authorized representative(s) may examine or request copies of USPh’s books, records, and other documents and supporting materials and/or conduct on-site reviews of any of USPh’s locations for the purpose of verifying and evaluating: (a) USPh’s compliance with the terms of this CIA; and (b) USPh’s compliance with the requirements of the Federal health care programs. The documentation described above shall be made available by USPh to OIG or its duly authorized representative(s) at all reasonable times for inspection, audit, and/or reproduction. Furthermore, for purposes of this provision, OIG or its duly authorized representative(s) may interview any of USPh’s Covered Persons who consent to be interviewed at the individual’s place of business during normal business hours or at such other place and time as may be mutually agreed upon between the individual and OIG. USPh shall assist OIG or its duly authorized representative(s) in contacting and arranging interviews with such individuals upon
OIG’s request. USPh’s Covered Persons may elect to be interviewed with or without a representative of USPh present.

VIII. DOCUMENT AND RECORD RETENTION

USPh shall maintain for inspection all documents and records relating to reimbursement from the Federal health care programs and to compliance with this CIA for six years (or longer if otherwise required by law) from the Effective Date.

IX. DISCLOSURES

Consistent with HHS’s FOIA procedures, set forth in 45 C.F.R. Part 5, OIG shall make a reasonable effort to notify USPh prior to any release by OIG of information submitted by USPh pursuant to its obligations under this CIA and identified upon submission by USPh as trade secrets, or information that is commercial or financial and privileged or confidential, under the FOIA rules. With respect to such releases, USPh shall have the rights set forth at 45 C.F.R. § 5.65(d).

X. BREACH AND DEFAULT PROVISIONS

USPh is expected to fully and timely comply with all of its CIA obligations.

A. Stipulated Penalties for Failure to Comply with Certain Obligations

As a contractual remedy, USPh and OIG hereby agree that failure to comply with certain obligations as set forth in this CIA may lead to the imposition of the following monetary penalties (hereinafter referred to as “Stipulated Penalties”) in accordance with the following provisions.

1. A Stipulated Penalty of $2,500 (which shall begin to accrue on the day after the date the obligation became due) for each day USPh fails to establish and implement any of the following obligations as described in Sections III and IV:

   a. a Compliance Officer or Old Towne Compliance Contact;

   b. a Compliance Committee;

   c. the Board of Directors compliance obligations;

   d. a written Code of Conduct;
e. written Policies and Procedures;

f. the development and/or implementation of a Training Plan for the training of Covered Persons, Relevant Covered Persons, and Board Members;

g. a risk assessment and mitigation process as required in Section III.E;

h. a Disclosure Program;

i. Ineligible Persons screening and removal requirements;

j. notification of Government investigations or legal proceedings; and

k. policies and procedures regarding the repayment of Overpayments;

l. the repayment of Overpayments as required by Section III.I;

m. reporting of Reportable Events; and

n. disclosure of changes to business units or locations.

2. A Stipulated Penalty of $2,500 (which shall begin to accrue on the day after the date the obligation became due) for each day USP fails to engage and use an IRO, as required in Section III.D, Appendix A, and Appendix B.

3. A Stipulated Penalty of $2,500 (which shall begin to accrue on the day after the date the obligation became due) for each day USP fails to submit the Implementation Report or any Annual Reports to OIG in accordance with the requirements of Section V by the deadlines for submission.

4. A Stipulated Penalty of $2,500 (which shall begin to accrue on the day after the date the obligation became due) for each day USP fails to submit any Claims Review Report in accordance with the requirements of Section III.D and Appendix B.
5. A Stipulated Penalty of $1,500 for each day USPh fails to grant access as required in Section VII. (This Stipulated Penalty shall begin to accrue on the date USPh fails to grant access.)

6. A Stipulated Penalty of $50,000 for each false certification submitted by or on behalf of USPh as part of its Implementation Report, Annual Report, additional documentation to a report (as requested by OIG), or otherwise required by this CIA.

7. A Stipulated Penalty of $1,000 for each day USPh fails to comply fully and adequately with any obligation of this CIA. OIG shall provide notice to USPh stating the specific grounds for its determination that USPh has failed to comply fully and adequately with the CIA obligation(s) at issue and steps USPh shall take to comply with the CIA. (This Stipulated Penalty shall begin to accrue 10 days after USPh receives this notice from OIG of the failure to comply.) A Stipulated Penalty as described in this Subsection shall not be demanded for any violation for which OIG has sought a Stipulated Penalty under Subsections 1–6 of this Section.

B. Timely Written Requests for Extensions

USPh may, in advance of the due date, submit a timely written request for an extension of time to perform any act or file any notification or report required by this CIA. Notwithstanding any other provision in this Section, if OIG grants the timely written request with respect to an act, notification, or report, Stipulated Penalties for failure to perform the act or file the notification or report shall not begin to accrue until one day after USPh fails to meet the revised deadline set by OIG. Notwithstanding any other provision in this Section, if OIG denies such a timely written request, Stipulated Penalties for failure to perform the act or file the notification or report shall not begin to accrue until three days after USPh receives OIG’s written denial of such request or the original due date, whichever is later. A “timely written request” is defined as a request in writing received by OIG at least five days prior to the date by which any act is due to be performed or any notification or report is due to be filed.

C. Payment of Stipulated Penalties

1. Demand Letter. Upon a finding that USPh has failed to comply with any of the obligations described in Section X.A and after determining that Stipulated Penalties are appropriate, OIG shall notify USPh of: (a) USPh’s failure to comply; and
(b) OIG’s exercise of its contractual right to demand payment of the Stipulated Penalties. (This notification shall be referred to as the “Demand Letter.”)

2. **Response to Demand Letter.** Within 10 days after the receipt of the Demand Letter, USPh shall either: (a) cure the breach to OIG’s satisfaction and pay the applicable Stipulated Penalties or (b) request a hearing before an HHS administrative law judge (ALJ) to dispute OIG’s determination of noncompliance, pursuant to the agreed upon provisions set forth below in Section X.E. In the event USPh elects to request an ALJ hearing, the Stipulated Penalties shall continue to accrue until USPh cures, to OIG’s satisfaction, the alleged breach in dispute. Failure to respond to the Demand Letter in one of these two manners within the allowed time period shall be considered a material breach of this CIA and shall be grounds for exclusion under Section X.D.

3. **Form of Payment.** Payment of the Stipulated Penalties shall be made by electronic funds transfer to an account specified by OIG in the Demand Letter.

4. **Independence from Material Breach Determination.** Except as set forth in Section X.D.1.c, these provisions for payment of Stipulated Penalties shall not affect or otherwise set a standard for OIG’s decision that USPh has materially breached this CIA, which decision shall be made at OIG’s discretion and shall be governed by the provisions in Section X.D, below.

D. **Exclusion for Material Breach of this CIA**

1. **Definition of Material Breach.** A material breach of this CIA means:

   a. repeated violations or a flagrant violation of any of the obligations under this CIA, including, but not limited to, the obligations addressed in Section X.A;

   b. a failure by USPh to report a Reportable Event, take corrective action, or make the appropriate refunds, as required in Section III.J;

   c. a failure to respond to a Demand Letter concerning the payment of Stipulated Penalties in accordance with Section X.C; or
d. a failure to engage and use an IRO in accordance with Section III.D, Appendix A, and Appendix B.

2. Notice of Material Breach and Intent to Exclude. The parties agree that a material breach of this CIA by USP\textsubscript{h} constitutes an independent basis for USP\textsubscript{h}’s exclusion from participation in the Federal health care programs. The length of the exclusion shall be in OIG’s discretion, but not more than five years per material breach. Upon a determination by OIG that USP\textsubscript{h} has materially breached this CIA and that exclusion is the appropriate remedy, OIG shall notify USP\textsubscript{h} of: (a) USP\textsubscript{h}’s material breach; and (b) OIG’s intent to exercise its contractual right to impose exclusion. (This notification shall be referred to as the “Notice of Material Breach and Intent to Exclude.”)

3. Opportunity to Cure. USP\textsubscript{h} shall have 30 days from the date of receipt of the Notice of Material Breach and Intent to Exclude to demonstrate that:
   a. the alleged material breach has been cured; or
   b. the alleged material breach cannot be cured within the 30-day period, but that: (i) USP\textsubscript{h} has begun to take action to cure the material breach; (ii) USP\textsubscript{h} is pursuing such action with due diligence; and (iii) USP\textsubscript{h} has provided to OIG a reasonable timetable for curing the material breach.

4. Exclusion Letter. If, at the conclusion of the 30-day period, USP\textsubscript{h} fails to satisfy the requirements of Section X.D.3, OIG may exclude USP\textsubscript{h} from participation in the Federal health care programs. OIG shall notify USP\textsubscript{h} in writing of its determination to exclude USP\textsubscript{h}. (This letter shall be referred to as the “Exclusion Letter.”) Subject to the Dispute Resolution provisions in Section X.E, below, the exclusion shall go into effect 30 days after the date of USP\textsubscript{h}’s receipt of the Exclusion Letter. The exclusion shall have national effect. Reinstatement to program participation is not automatic. After the end of the period of exclusion, USP\textsubscript{h} may apply for reinstatement by submitting a written request for reinstatement in accordance with the provisions at 42 C.F.R. §§ 1001.3001–.3004.

E. Dispute Resolution

1. Review Rights. Upon OIG’s delivery to USP\textsubscript{h} of its Demand Letter or of its Exclusion Letter, and as an agreed-upon contractual remedy for the resolution of disputes arising under this CIA, USP\textsubscript{h} shall be afforded certain review rights comparable to the ones that are provided in 42 U.S.C. § 1320a-7(f) and 42 C.F.R. Part 1005 as if they
applied to the Stipulated Penalties or exclusion sought pursuant to this CIA. Specifically, OIG’s determination to demand payment of Stipulated Penalties or to seek exclusion shall be subject to review by an HHS ALJ and, in the event of an appeal, the HHS Departmental Appeals Board (DAB), in a manner consistent with the provisions in 42 C.F.R. § 1005.2–1005.21. Notwithstanding the language in 42 C.F.R. § 1005.2(c), the request for a hearing involving Stipulated Penalties shall be made within 10 days after receipt of the Demand Letter and the request for a hearing involving exclusion shall be made within 25 days after receipt of the Exclusion Letter. The procedures relating to the filing of a request for a hearing can be found at http://www.hhs.gov/dab/divisions/civil/procedures/divisionprocedures.html.

2. *Stipulated Penalties Review.* Notwithstanding any provision of Title 42 of the United States Code or Title 42 of the Code of Federal Regulations, the only issues in a proceeding for Stipulated Penalties under this CIA shall be: (a) whether USP\textsubscript{h} was in full and timely compliance with the obligations of this CIA for which OIG demands payment; and (b) the period of noncompliance. USP\textsubscript{h} shall have the burden of proving its full and timely compliance and the steps taken to cure the noncompliance, if any. OIG shall not have the right to appeal to the DAB an adverse ALJ decision related to Stipulated Penalties. If the ALJ agrees with OIG with regard to a finding of a breach of this CIA and orders USP\textsubscript{h} to pay Stipulated Penalties, such Stipulated Penalties shall become due and payable 20 days after the ALJ issues such a decision unless USP\textsubscript{h} requests review of the ALJ decision by the DAB. If the ALJ decision is properly appealed to the DAB and the DAB upholds the determination of OIG, the Stipulated Penalties shall become due and payable 20 days after the DAB issues its decision.

3. *Exclusion Review.* Notwithstanding any provision of Title 42 of the United States Code or Title 42 of the Code of Federal Regulations, the only issues in a proceeding for exclusion based on a material breach of this CIA shall be whether USP\textsubscript{h} was in material breach of this CIA and, if so, whether:

a. USP\textsubscript{h} cured such breach within 30 days of its receipt of the Notice of Material Breach; or

b. the alleged material breach could not have been cured within the 30-day period, but that, during the 30-day period following USP\textsubscript{h}’s receipt of the Notice of Material Breach:
   (i) USP\textsubscript{h} had begun to take action to cure the material breach;
   (ii) USP\textsubscript{h} pursued such action with due diligence; and (iii) USP\textsubscript{h} provided to OIG a reasonable timetable for curing the material breach.
For purposes of the exclusion herein, exclusion shall take effect only after an ALJ decision favorable to OIG, or, if the ALJ rules for USPh, only after a DAB decision in favor of OIG. USPh’s election of its contractual right to appeal to the DAB shall not abrogate OIG’s authority to exclude USPh upon the issuance of an ALJ’s decision in favor of OIG. If the ALJ sustains the determination of OIG and determines that exclusion is authorized, such exclusion shall take effect 20 days after the ALJ issues such a decision, notwithstanding that USPh may request review of the ALJ decision by the DAB. If the DAB finds in favor of OIG after an ALJ decision adverse to OIG, the exclusion shall take effect 20 days after the DAB decision. USPh shall waive its right to any notice of such an exclusion if a decision upholding the exclusion is rendered by the ALJ or DAB. If the DAB finds in favor of USPh, USPh shall be reinstated effective on the date of the original exclusion.

4. Finality of Decision. The review by an ALJ or DAB provided for above shall not be considered to be an appeal right arising under any statutes or regulations. Consequently, the parties to this CIA agree that the DAB’s decision (or the ALJ’s decision if not appealed) shall be considered final for all purposes under this CIA.

XI. EFFECTIVE AND BINDING AGREEMENT

USPh, Old Towne, and OIG agree as follows:

A. This CIA shall become final and binding on the date the final signature is obtained on the CIA.

B. This CIA constitutes the complete agreement between the parties and may not be amended except by written consent of the parties to this CIA.

C. OIG may agree to a suspension of USPh’s or Old Towne’s obligations under this CIA based on a certification by USPh or Old Towne that it is no longer providing health care items or services that will be billed to any Federal health care program and that it does not have any ownership or control interest, as defined in 42 U.S.C. § 1320a-3, in any entity that bills any Federal health care program. If USPh or Old Towne is relieved of its CIA obligations, USPh or Old Towne will be required to notify OIG in writing at least 30 days in advance if USPh or Old Towne plans to resume providing health care items or services that are billed to any Federal health care program or to obtain an ownership or control interest in any entity that bills any Federal health care program. At such time, OIG shall evaluate whether the CIA will be reactivated or modified.
D. The undersigned USP\textsubscript{h} and Old Towne signatories represent and warrant that they are authorized to execute this CIA. The undersigned OIG signatories represent that they are signing this CIA in their official capacities and that they are authorized to execute this CIA.

E. This CIA may be executed in counterparts, each of which constitutes an original and all of which constitute one and the same CIA. Facsimiles of signatures shall constitute acceptable, binding signatures for purposes of this CIA.
ON BEHALF OF U.S. PHYSICAL THERAPY, INC.
/Christopher Reading/

CHRISTOPHER READING
Chief Executive Officer
U.S. Physical Therapy, Inc.
/Julie E. Kass/

DATE
12/15/15

ON BEHALF OF
OLD TOWNE PHYSICAL THERAPY LIMITED PARTNERSHIP
/Christopher Reading/

CHRISTOPHER READING
President
Rehab Partners #3
/Julie E. Kass/

DATE
12/15/15

JULIE E. KASS
Ober Kaler
Counsel for U.S. Physical Therapy, Inc.

DATE
12/17/15

JULIE E. KASS
Ober Kaler
Counsel for Old Towne Physical Therapy Limited Partnership

DATE
12/17/15
APPENDIX A

INDEPENDENT REVIEW ORGANIZATION

This Appendix contains the requirements relating to the Independent Review Organization (IRO) required by Section III.D of the CIA.

A. IRO Engagement

1. USPh shall engage an IRO that possesses the qualifications set forth in Paragraph B, below, to perform the responsibilities in Paragraph C, below. The IRO shall conduct the review in a professionally independent and objective fashion, as set forth in Paragraph D, below. Within 30 days after OIG receives the information identified in Section V.A.7 of the CIA or any additional information submitted by USPh in response to a request by OIG, whichever is later, OIG will notify USPh if the IRO is unacceptable. Absent notification from OIG that the IRO is unacceptable, USPh may continue to engage the IRO.

2. If USPh engages a new IRO during the term of the CIA, this IRO shall also meet the requirements of this Appendix. If a new IRO is engaged, USPh shall submit the information identified in Section V.A.7 of the CIA to OIG within 30 days of engagement of the IRO. Within 30 days after OIG receives this information or any additional information submitted by USPh at the request of OIG, whichever is later, OIG will notify USPh if the IRO is unacceptable. Absent notification from OIG that the IRO is unacceptable, USPh may continue to engage the IRO.

B. IRO Qualifications

The IRO shall:

1. assign individuals to conduct the Claims Review who have expertise in the billing, coding, reporting, and other requirements applicable to outpatient physical and occupational therapy services and in the general requirements of the Federal health care program(s) from which USPh seeks reimbursement;

2. assign individuals to design and select the Claims Review sample who are knowledgeable about the appropriate statistical sampling techniques;

3. assign individuals to conduct the coding review portions of the Claims Review who have a nationally recognized coding certification and who have
maintained this certification (e.g., completed applicable continuing education requirements); and

4. have sufficient staff and resources to conduct the reviews required by the CIA on a timely basis.

C. IRO Responsibilities

The IRO shall:

1. perform each Claims Review in accordance with the specific requirements of the CIA;

2. follow all applicable Medicare rules and reimbursement guidelines in making assessments in the Claims Review;

3. if in doubt of the application of a particular Medicare policy or regulation, request clarification from the appropriate authority (e.g., Medicare contractor);

4. respond to all OIG inquires in a prompt, objective, and factual manner; and

5. prepare timely, clear, well-written reports that include all the information required by Appendix B to the CIA.

D. IRO Independence and Objectivity

The IRO must perform the Claims Review in a professionally independent and objective fashion, as defined in the most recent Government Auditing Standards issued by the United States Government Accountability Office.

E. IRO Removal/Termination

1. USPh and IRO. If USPh terminates its IRO or if the IRO withdraws from the engagement during the term of the CIA, USPh must submit a notice explaining its reasons for termination or the reason for withdrawal to OIG no later than 30 days after termination or withdrawal. USPh must engage a new IRO in accordance with Paragraph A of this Appendix and within 60 days of termination or withdrawal of the IRO.
2. **OIG Removal of IRO.** In the event OIG has reason to believe the IRO does not possess the qualifications described in Paragraph B, is not independent and objective as set forth in Paragraph D, or has failed to carry out its responsibilities as described in Paragraph C, OIG may, at its sole discretion, require USPh to engage a new IRO in accordance with Paragraph A of this Appendix. USPh must engage a new IRO within 60 days of termination of the IRO.

Prior to requiring USPh to engage a new IRO, OIG shall notify USPh of its intent to do so and provide a written explanation of why OIG believes such a step is necessary. To resolve any concerns raised by OIG, USPh may present additional information regarding the IRO’s qualifications, independence, or performance of its responsibilities. OIG will attempt in good faith to resolve any differences regarding the IRO with USPh prior to requiring USPh to terminate the IRO. However, the final determination as to whether or not to require USPh to engage a new IRO shall be made at the sole discretion of OIG.
APPENDIX B

CLAIMS REVIEW

A. Claims Review. The IRO shall perform the Claims Review annually to cover each of the five Reporting Periods. The IRO shall perform all components of each Claims Review.

1. Definitions. For the purposes of the Claims Review, the following definitions shall be used:

a. Overpayment: The amount of money USPh has received in excess of the amount due and payable under any Federal health care program requirements, as determined by the IRO in connection with the claims reviews performed under this Appendix B, and which shall include any extrapolated Overpayments determined in accordance with Section A.4 of this Appendix B.

b. Paid Claim: An outpatient physical or occupational therapy claim submitted by USPh and for which USPh has received reimbursement from the Medicare program.

c. Population 1: Population 1 shall be defined as all Paid Claims for services rendered at Old Towne during the 12-month period covered by the Claims Review.

d. Population 2: Population 2 shall be defined as all Paid Claims for services rendered at each of the 10 randomly selected non-Old Towne Clinics that employed or contracted with a physical therapist assistant (PTA) during the 12-month period covered by the Claims Review.

e. Error Rate: The Error Rate shall be the percentage of net Overpayments identified in the sample. The net Overpayments shall be calculated by subtracting all underpayments identified in the sample from all gross Overpayments identified in the sample. (Note: Any potential cost settlements or other supplemental payments should not be included in the net Overpayment calculation. Rather, only underpayments identified as part of the Discovery Samples shall be included as part of the net Overpayment calculation.)
The Error Rate is calculated by dividing the net Overpayment identified in the sample by the total dollar amount associated with the Paid Claims in the sample.

2. **Discovery Sample 1.** For each Reporting Period, the IRO shall randomly select and review a sample of 100 Paid Claims for services rendered at Old Towne (Discovery Sample 1). The Paid Claims shall be reviewed based on the supporting documentation available at USPh’s offices or under USPh’s control and applicable billing and coding regulations and guidance to determine whether the claim was correctly coded, submitted, and reimbursed.

   If the Error Rate (as defined above) for Discovery Sample 1 is less than 5%, no additional sampling is required, nor is the Systems Review required. (Note: The guidelines listed above do not imply that this is an acceptable error rate. Accordingly, USPh should, as appropriate, further analyze any errors identified in Discovery Sample 1. USPh recognizes that OIG or another HHS component, in its discretion and as authorized by statute, regulation, or other appropriate authority may also analyze or review Paid Claims included, or errors identified, in Discovery Sample 1 or any other segment of the universe.)

3. **Discovery Sample 2.** For each Reporting Period, the IRO shall review a sample of 500 Paid Claims submitted for services rendered at 10 randomly selected Clinics that employed or contracted with a PTA at any time during the applicable 12-month period, not including Old Towne (Discovery Sample 2). Specifically, the IRO shall randomly select and review a sample of 50 Paid Claims for each of the 10 randomly selected Clinics based on the supporting documentation available at USPh’s offices or under USPh’s control and applicable billing and coding regulations and guidance to determine whether the claim was correctly coded, submitted, and reimbursed.

   If the Error Rate (as defined above) for each of the 10 randomly selected Clinics is less than 5%, no additional sampling is required, nor is the Systems Review required. (Note: The guidelines listed above do not imply that this is an acceptable error rate. Accordingly, USPh should, as appropriate, further analyze any errors identified in Discovery Sample 2. USPh recognizes that OIG or another HHS component, in its discretion and as authorized by statute, regulation, or other appropriate authority may also analyze or review Paid Claims included, or errors identified, in Discovery Sample 2 or any other segment of the universe.)

4. **Full Sample.** If Discovery Sample 1 indicates that the Error Rate is 5% or greater, the IRO shall select an additional sample of Paid Claims for services rendered at Old Towne (Full Sample 1) using commonly accepted sampling methods. If Discovery Sample 2 indicates that the individual Error Rate is 5% or greater at any of the

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10 randomly selected Clinics, the IRO shall select an additional sample of Paid Claims for each of the Clinics at which the Error Rate was 5% or greater (Full Sample 2). The Paid Claims selected for Full Sample 1 or Full Sample 2 shall be reviewed based on supporting documentation available at USP’h’s offices or under USP’h’s control and applicable billing and coding regulations and guidance to determine whether the claim was correctly coded, submitted, and reimbursed.

For purposes of calculating the size of Full Sample 1 or Full Sample 2, the Discovery Samples may serve as the probe samples, if statistically appropriate. Additionally, the IRO may use the Paid Claims sampled as part of the Discovery Samples, and the corresponding findings for those Paid Claims, as part of its Full Samples, if: (1) statistically appropriate and (2) the IRO selects the Full Sample Paid Claims using the seed number generated by the Discovery Samples. The findings of the Full Samples shall be used by the IRO to estimate the actual Overpayment in Population 1 or Population 2 (on a per-Clinic basis) with a 90% confidence level and with a maximum relative precision of 25% of the point estimate. OIG, in its sole discretion, may refer the findings of the Full Samples (and any related workpapers) received from USP’h to the appropriate Federal health care program payor (e.g., Medicare contractor) for appropriate follow-up by that payor.

5. **Systems Review.** If the Discovery Sample at any Clinic identifies an Error Rate of 5% or greater, USP’h’s IRO shall also conduct a Systems Review. The Systems Review shall consist of the following:

a. a review of USP’h’s billing and coding systems and processes relating to outpatient physical and occupational therapy claims submitted to Federal health care programs (including, but not limited to, the operation of the billing system; the process by which claims are coded; safeguards to ensure proper coding, claims submission, and billing; and procedures to identify and correct inaccurate coding and billing); and

b. on a per-Clinic basis, for each outpatient physical or occupational therapy claim in the Discovery Samples and Full Samples that resulted in an Overpayment, the IRO shall review the system(s) and process(es) that generated the claim and identify any problems or weaknesses that may have resulted in the identified Overpayments. The IRO shall provide its observations and recommendations on suggested improvements to the system(s) and the process(es) that generated the claim.
6. **Other Requirements**

a. **Supplemental Materials.** The IRO shall request all documentation and materials required for its review of the Paid Claims selected as part of the Discovery Samples or Full Samples (if applicable), and USPh shall furnish such documentation and materials to the IRO prior to the IRO initiating its review of the Discovery Samples or Full Samples (if applicable). If the IRO accepts any supplemental documentation or materials from USPh after the IRO has completed its initial review of the Discovery Samples or Full Samples (if applicable) (Supplemental Materials), the IRO shall identify in the Claims Review Report the Supplemental Materials, the date the Supplemental Materials were accepted, and the relative weight the IRO gave to the Supplemental Materials in its review. In addition, the IRO shall include a narrative in the Claims Review Report describing the process by which the Supplemental Materials were accepted and the IRO’s reasons for accepting the Supplemental Materials.

b. **Paid Claims without Supporting Documentation.** Any Paid Claim for which USPh cannot produce documentation sufficient to support the Paid Claim shall be considered an error and the total reimbursement received by USPh for such Paid Claim shall be deemed an Overpayment. Replacement sampling for Paid Claims with missing documentation is not permitted.

c. **Use of First Samples Drawn.** For the purposes of all samples (Discovery Samples and Full Samples) discussed in this Appendix, the Paid Claims selected in each first sample shall be used (i.e., it is not permissible to generate more than one list of random samples and then select one for use with the Discovery Samples or Full Samples).

7. **Repayment of Identified Overpayments.** USPh shall repay within 30 days any Overpayment(s) identified in a Discovery Sample, regardless of the Error Rate, and (if applicable) a Full Sample, including the IRO’s estimate of the actual Overpayment in Population 1 or Population 2 (on a per-Clinic basis) as determined in accordance with Section A.4 above, in accordance with payor refund policies. USPh shall make available to OIG all documentation that reflects the refund of the Overpayment(s) to the payor.
B. **Claims Review Report.** The IRO shall prepare a Claims Review Report as described in this Appendix for each Claims Review performed. The following information shall be included in the Claims Review Report for each Discovery Sample and Full Sample (if applicable).

1. **Claims Review Methodology**


   b. **Claims Review Objective.** A clear statement of the objective intended to be achieved by the Claims Review.

   c. **Source of Data.** A description of the specific documentation relied upon by the IRO when performing the Claims Review (e.g., medical records; physician orders; certificates of medical necessity; requisition forms; local medical review policies (including title and policy number); CMS program memoranda (including title and issuance number); Medicare carrier or intermediary manual or bulletins (including issue and date); other policies, regulations, or directives).

   d. **Review Protocol.** A narrative description of how the Claims Review was conducted and what was evaluated.

   e. **Supplemental Materials.** A description of any Supplemental Materials as required by Section A.6.a, above.

2. **Statistical Sampling Documentation**

   a. A copy of the printout of the random numbers generated by the “Random Numbers” function of the statistical sampling software used by the IRO.

   b. A copy of the statistical software printout(s) estimating how many Paid Claims are to be included in the Full Samples, if applicable.

   c. A description or identification of the statistical sampling software package used to select the sample and determine the Full Sample sizes, if applicable.
3. **Claims Review Findings**

a. **Narrative Results**

i. A description of USPh’s billing and coding system(s), including the identification, by position description, of the personnel involved in coding and billing.

ii. A narrative explanation of the IRO’s findings and supporting rationale (including reasons for errors, patterns noted, etc.) regarding the Claims Review, including the results of the Discovery Samples, and the results of the Full Samples (if any).

b. **Quantitative Results**

i. Total number and percentage of instances in which the IRO determined that the Paid Claims submitted by USPh (Claim Submitted) differed from what should have been the correct claim (Correct Claim), regardless of the effect on the payment.

ii. Total number and percentage of instances in which the Claim Submitted differed from the Correct Claim and in which such difference resulted in an Overpayment to USPh.

iii. Total dollar amount of all Overpayments in the Discovery Samples and the Full Samples (if applicable).

iv. Total dollar amount of Paid Claims included in the Discovery Samples and the Full Samples and the net Overpayment associated with the Discovery Samples and the Full Samples.

v. Error Rate in the Discovery Samples and the Full Samples.

vi. A spreadsheet of the Claims Review results that includes the following information for each Paid Claim: Federal health care program billed, beneficiary
health insurance claim number, date of service, code
submitted (e.g., DRG, CPT code, etc.), code
reimbursed, allowed amount reimbursed by payor,
correct code (as determined by the IRO), correct
allowed amount (as determined by the IRO), dollar
difference between allowed amount reimbursed by
payor and the correct allowed amount.

vii. If any Full Samples are performed, the methodology
used by the IRO to estimate the actual Overpayment in
Population 1 or Population 2 (on a per-Clinic basis)
and the amount of such Overpayment.

c. Recommendations. The IRO’s report shall include any
recommendations for improvements to USPh’s billing and
coding system based on the findings of the Claims Review.

4. Systems Review Findings. The IRO shall prepare a Systems Review
Report based on the Systems Review performed (if applicable) that shall include the
IRO’s observations, findings, and recommendations regarding:

a. the strengths and weaknesses in USPh’s billing systems and
processes;

b. the strengths and weaknesses in USPh’s coding systems and
processes; and

c. possible improvements to USPh’s billing and coding systems
and processes to address the specific problems or weaknesses
that resulted in the identified Overpayments.

5. Credentials. The names and credentials of the individuals who: (1)
designed the statistical sampling procedures and the review methodology utilized for the
Claims Review and (2) performed the Claims Review.