

**INTEGRITY AGREEMENT  
BETWEEN THE  
OFFICE OF INSPECTOR GENERAL  
OF THE  
DEPARTMENT OF HEALTH AND HUMAN SERVICES  
AND  
JOHN T. LAROSSA, M.D.**

**I. PREAMBLE**

John T. Larossa, M.D. (Dr. Larossa) hereby enters into this Integrity Agreement (IA) with the Office of Inspector General (OIG) of the United States Department of Health and Human Services (HHS) to promote compliance by Dr. Larossa and his associates, employees, contractors, and agents with the statutes, regulations, program requirements, 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). This commitment to promote compliance applies to John T. Larossa, M.D., and any other entity that Dr. Larossa owns or in which Dr. Larossa has a control interest, as defined in 42 U.S.C. § 1320a-3(a)(3), and Dr. Larossa's and any such entity's Covered Persons as defined in Section II.C. Contemporaneously with this Agreement, Dr. Larossa is entering into a Settlement Agreement with the United States, and this Agreement is incorporated by reference into the Settlement Agreement.

**II. TERM OF THE AGREEMENT**

A. The date on which the final signatory of this IA executes this IA shall be known as the Effective Date. The period of compliance obligations assumed by Dr. Larossa under this IA shall be 5 years from the Effective Date. 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, VIII, IX, X, and XI shall expire no later than 120 days from OIG's receipt of: (1) Dr. Larossa's final Annual Report; or (2) any additional materials submitted by Dr. Larossa pursuant to OIG's request, whichever is later.

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

1. "Covered Persons" includes:

- a. Dr. Larossa and all associates and employees of Dr. Larossa; and
- b. all contractors, agents, and other persons who provide patient care items or services or who perform billing or coding functions on behalf of Dr. Larossa.

### III. INTEGRITY OBLIGATIONS

Dr. Larossa shall establish and maintain a Compliance Program that includes the following elements:

#### A. Compliance Contact

Within 30 days after the Effective Date, Dr. Larossa shall designate a person to be responsible for compliance activities ("Compliance Contact"). Dr. Larossa shall maintain a Compliance Contact for the term of this IA. The Compliance Contact shall be responsible for: (1) developing and implementing policies, procedures, and practices designed to ensure compliance with the requirements set forth in this IA and with Federal health care program requirements; (2) monitoring Dr. Larossa's day-to-day compliance activities; and (3) meeting all reporting obligations created under this IA.

Dr. Larossa shall report to OIG, in writing, any changes in the identity or job responsibilities of the Compliance Contact, or any actions or changes that would affect the Compliance Contact's ability to perform the duties necessary to meet the obligations in this IA, within 15 days after such change. The name, address, phone number, and a description of any other job responsibilities performed by the Compliance Contact shall be included in the Implementation Report.

#### B. Posting of Notice

Within the 90 days after the Effective Date, Dr. Larossa shall post in a prominent place accessible to all patients and Covered Persons a notice detailing his commitment to comply with all Federal health care program requirements in the conduct of his business.

This notice shall include the following information: (i) a means (e.g., telephone number or address) by which billing concerns and other issues may be reported anonymously; (ii) Dr. Larossa's commitment to maintain the confidentiality of the report; and (iii) notification that reporting concerns and issues will not result in retribution or retaliation by Dr. Larossa.

This notice shall also include the HHS OIG Fraud Hotline telephone number (1-800-HHS-TIPS) as a confidential means by which suspected fraud or abuse in the Federal health care programs might be reported.

A copy of this notice shall be included in the Implementation Report.

**C. Written Policies and Procedures**

Within 90 days after the Effective Date, Dr. Larossa shall develop, implement, and distribute written Policies and Procedures to all Covered Persons. In addition, Dr. Larossa shall make the promotion of, and adherence to, the written Policies and Procedures an element in evaluating the performance of all employees. The written Policies and Procedures shall, at a minimum, set forth:

1. Dr. Larossa's commitment to full compliance with all Federal health care program requirements, including his commitment to prepare and submit accurate claims consistent with such requirements;
2. the expectation that all of Dr. Larossa's Covered Persons shall be expected to comply with all Federal health care program requirements and with Dr. Larossa's own written Policies and Procedures as implemented pursuant to this Section III.C (including the requirements of this Agreement);
3. the responsibility and requirement that all Covered Persons report suspected violations of any Federal health care program requirements or of Dr. Larossa's own Policies and Procedures to the Compliance Contact and Dr. Larossa's commitment to maintain confidentiality and anonymity, as appropriate, and not to retaliate with respect to such disclosures;

4. the possible consequences to both Dr. Larossa and Covered Persons of failure to comply with Federal health care program requirements or with Dr. Larossa's written Policies and Procedures and the failure to report such noncompliance;
5. Dr. Larossa's commitment to remain current with all Federal health care program requirements by obtaining and reviewing program memoranda, newsletters, and any other correspondence from the carrier related to Federal health care program requirements;
6. the proper procedures for the accurate preparation and submission of claims in accordance with Federal health care program requirements;
7. the proper documentation of services and billing information; and
8. the proper coding of services provided.

Within 90 days after the Effective Date, each Covered Person shall certify in writing that he or she has received, read, understood, and shall abide by Dr. Larossa's written Policies and Procedures. New Covered Persons shall receive and review the written Policies and Procedures and shall complete the required certification within 30 days after becoming a Covered Person or within 90 days after the Effective Date, whichever is later.

At least annually (and more frequently if appropriate), Dr. Larossa shall assess and update, as necessary, the Policies and Procedures. Within 30 days after the effective date of any revisions, the relevant portions of any such revised Policies and Procedures shall be distributed to all Covered Persons. Appropriate and knowledgeable staff shall be available to explain the Policies and Procedures.

Copies of the written Policies and Procedures shall be included in the Implementation Report. Copies of any written Policies and Procedures that are subsequently revised shall be included in the next Annual Report along with a summary of any change or amendment to each Policy and Procedure required by this Section and the reason for each change.

**D. Training and Certification**

Within 90 days after the Effective Date and at least once each year thereafter, Dr. Larossa and Covered Persons shall receive at least three hours of training from an individual or entity, other than Dr. Larossa or another Covered Person. Persons providing the training shall be knowledgeable about the subject area and may be received from a variety of sources (e.g., CME classes, hospitals, associations, carriers).

New Covered Persons shall receive the training described above within 30 days after becoming a Covered Person or within 90 days after the Effective Date, whichever is later. The training for new Covered Persons may either be provided internally by a Covered Person who has completed the required annual training or externally by a qualified individual or entity. A new Covered Person shall work under the direct supervision of a Covered Person who has received such training, to the extent that the work relates to the delivery of patient care items or services and/or the preparation or submission of claims for reimbursement from any Federal health care program, until such time as the new Covered Person completes the training.

At a minimum, the initial, annual, and new employee training sessions shall include the following topics:

1. the requirements of Dr. Larossa's IA;
2. an overview of the Dr. Larossa's compliance program;
3. the accurate coding and submission of claims for services rendered and/or items provided to Federal health care program beneficiaries;
4. applicable reimbursement statutes, regulations, and program requirements and directives;
5. the written Policies and Procedures developed pursuant to Section III.C, above;
6. the policies, procedures, and other requirements applicable to the documentation of medical records;

7. the personal obligation of each individual involved in the coding and claims submission process to ensure that such claims are accurate;
8. the legal sanctions for the submission of improper claims or violations of the Federal health care program requirements; and
9. examples of proper and improper coding and claim submission practices.

Each Covered Person shall annually certify, in writing or in electronic format if the training is computerized, that he or she has received the required training. The certification shall specify the date the training was received. The Compliance Contact shall retain the certifications, along with all training materials. Dr. Larossa shall annually review the training, and, where appropriate, update the training to reflect changes in Federal health care program requirements, any issues discovered during the IRO Claims Review or other Review, and any other relevant information.

Dr. Larossa may provide the training required under this IA through appropriate computer-based training approaches. If Dr. Larossa chooses to provide computer-based training, he shall make available appropriately qualified and knowledgeable staff or trainers to answer questions or provide additional information to the individuals receiving such training.

The training materials shall be provided in the Implementation Report, and to the extent the training is revised, shall also be included in the Annual Reports. The certifications shall be made available to OIG, upon request.

**E. Review Procedures**

1. General Description.

a. *Engagement of Independent Review Organization.* Within 90 days after the Effective Date, Dr. Larossa shall engage an entity (or entities), such as an accounting, auditing, or consulting firm (hereinafter "Independent Review Organization" or "IRO"), to perform reviews to assist Dr. Larossa in assessing and evaluating his billing and coding practices and certain other obligations pursuant to this IA and the Settlement Agreement.

Each IRO engaged by Dr. Larossa shall have expertise in the billing, coding, reporting, and other requirements relevant to physician practices and in the general requirements of the Federal health care program(s) from which Dr. Larossa seeks reimbursement. Each IRO shall assess, along with Dr. Larossa, whether it can perform the IRO review in a professionally independent and/or objective fashion, as appropriate to the nature of the engagement; taking into account any other business relationships or other engagements that may exist.

The IRO(s) review shall evaluate and analyze Dr. Larossa's coding, billing, and claims submission to the Federal health care programs and the reimbursement received (Claims Review). If Dr. Larossa or an entity in which Dr. Larossa has an ownership or control interest in (as defined in 42 U.S.C. §1320a-3(a)(3)) submits cost reports, Dr. Larossa shall engage an IRO to analyze whether Dr. Larossa sought payment for certain unallowable costs (Unallowable Cost Review). The applicable requirements relating to the IRO are outlined in Appendix A to this Agreement, which is incorporated by reference.

b. *Frequency of Claims Review.* The Claims Review shall be performed annually and shall cover each of the Reporting Periods. The IRO(s) shall perform all components of each annual Claims Review.

c. *Frequency of Unallowable Cost Review.* If applicable, the IRO shall perform the Unallowable Cost Review for the first Reporting Period. If not applicable, Dr. Larossa shall sign a certification, as required by Section III.E.4 below, stating that he does not currently and has not submitted a cost report since this IA was executed.

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

2. Claims Review. The Claims Review shall include a Discovery Sample and, if necessary, a Full Sample. The applicable definitions, procedures, and reporting requirements are outlined in Appendix B to this IA, which is incorporated by reference.

a. *Discovery Sample.* The IRO shall randomly select and review a sample of 50 Medicare Paid Claims submitted by or on behalf of Dr. Larossa (Discovery Sample).

The Paid Claims shall be reviewed based on the supporting documentation available at Dr. Larossa's office or under Dr. Larossa's control and applicable billing and coding regulations and guidance to determine whether the claim submitted was correctly coded, submitted, and reimbursed.

i. If the Error Rate (as defined in Appendix B) for the Discovery Sample 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, Dr. Larossa should, as appropriate, further analyze any errors identified in the Discovery Sample. Dr. Larossa recognizes that OIG or other 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 the Discovery Sample or any other segment of the universe.)

ii. If the Discovery Sample indicates that the Error Rate is 5% or greater, the IRO shall perform a Full Sample and a Systems Review, as described below.

b. *Full Sample.* If necessary, as determined by procedures set forth in Section III.E.2.a, the IRO shall perform an additional sample of Paid Claims using commonly accepted sampling methods and in accordance with Appendix B. The Full Sample shall be designed to: (i) estimate the actual Overpayment in the population with a 90% confidence level and with a maximum relative precision of 25% of the point estimate; and (ii) conform with the Centers for Medicare and Medicaid Services' statistical sampling for overpayment estimation guidelines. The Paid Claims shall be reviewed based on supporting documentation available at Dr. Larossa's office or under Dr. Larossa's control and applicable billing and coding regulations and guidance to determine whether the claim submitted was correctly coded, submitted, and reimbursed. For purposes of calculating the size of the Full Sample, the Discovery Sample may



serve as the probe sample, if statistically appropriate. Additionally, the IRO may use the Items sampled as part of the Discovery Sample, and the corresponding findings for those 50 Items, as part of its Full Sample, if: (i) statistically appropriate and (ii) the IRO selects the Full Sample Items using the seed number generated by the Discovery Sample. OIG, in its sole discretion, may refer the findings of the Full Sample (and any related workpapers) received from Dr. Larossa to the appropriate Federal health care program payor, including the Medicare contractor (e.g., carrier, fiscal intermediary, or DMERC), for appropriate follow-up by that payor.

c. *Systems Review.* If the IRO's Discovery Sample identifies an Error Rate of 5% or greater, the IRO shall also conduct a Systems Review. Specifically, for each claim in the Discovery Sample and Full Sample that resulted in an Overpayment, the IRO shall perform a "walk through" of the system(s) and process(es) that generated the claim to 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.

d. *Repayment of Identified Overpayments.* In accordance with Section III.H.1 of this Agreement, Dr. Larossa shall repay within 30 days any Overpayment(s) identified in the Discovery Sample or the Full Sample (if applicable), regardless of the Error Rate, to the appropriate payor and in accordance with payor refund policies. Dr. Larossa shall make available to OIG any and all documentation and the associated documentation that reflects the refund of the Overpayment(s) to the payor.

3. Claims Review Report. The IRO shall prepare a report based upon the Claims Review performed (the "Claims Review Report"). Information to be included in the Claims Review Report is described in Appendix B.

4. Unallowable Cost Review.

a. Unless paragraph III.E.4.b, below, applies, the IRO shall conduct a review of Dr. Larossa's compliance with the unallowable cost provisions of the Settlement Agreement. The IRO shall determine whether Dr. Larossa has complied with his obligations not to charge

to, or otherwise seek payment from, Federal or State payors for unallowable costs (as defined in the Settlement Agreement) and his obligation to identify to applicable Federal or State payors any unallowable costs included in payments previously sought from the United States, or any State Medicaid program. This unallowable cost analysis shall include, but not be limited to, payments sought in any cost reports, cost statements, information reports, or payment requests already submitted by Dr. Larossa or any affiliates. To the extent that such cost reports, cost statements, information reports, or payment requests, even if already settled, have been adjusted to account for the effect of the inclusion of the unallowable costs, the IRO shall determine if such adjustments were proper. In making this determination, the IRO may need to review cost reports and/or financial statements from the year in which the Settlement Agreement was executed, as well as from previous years.

- b. If Dr. Larossa executes and submits to the OIG in the first Annual Report a certification that neither Dr. Larossa nor any entity in which Dr. Larossa has any ownership or control interest has ever submitted any cost report or other submission to a Federal health care program seeking reimbursement based on costs, then no Unallowable Cost Review will be required under this IA.

5. Unallowable Cost Review Report. If applicable, the IRO shall prepare a report based upon the Unallowable Cost Review performed. The Unallowable Cost Review Report shall include the IRO's findings and supporting rationale regarding the Unallowable Costs Review and whether Dr. Larossa has complied with his obligation not to charge to, or otherwise seek payment from, Federal or State payors for unallowable costs (as defined in the Settlement Agreement) and his obligation to identify to applicable Federal or State payors any unallowable costs included in payments previously sought from such payor.

6. Validation Review. In the event OIG has reason to believe that:  
(a) Dr. Larossa's Claims Review or any other review required by Section III.E (Review) fails to conform to the requirements of this IA; or (b) the IRO's findings or Claims Review results or other results are inaccurate, OIG may, at its sole discretion, conduct its own review to determine whether the Claims Review or other Review complied with the requirements of the IA and/or the findings or Claims Review results or

other results are inaccurate (Validation Review). Dr. Larossa shall pay for the reasonable cost of any such review performed by OIG or any of its designated agents so long as it is initiated within one year after Dr. Larossa's final submission (as described in Section II) is received by OIG.

Prior to initiating a Validation Review, OIG shall notify Dr. Larossa 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, Dr. Larossa may request a meeting with OIG to: (a) discuss the results of any Claims Review or other Review submissions or findings; (b) present any additional or relevant information to clarify the results of the Claims Review or other Review to correct the inaccuracy of the Claims Review or other Review; and/or (c) propose alternatives to the proposed Validation Review. Dr. Larossa agrees to provide any additional information requested by OIG under this Section in an expedited manner. OIG will attempt in good faith to resolve any Claims Review or other Review issues with Dr. Larossa 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.

7. Independence/Objectivity Certification. The IRO shall include in its report(s) to Dr. Larossa a certification or sworn affidavit that it has evaluated its professional independence and/or objectivity, as appropriate to the nature of the engagement, with regard to the Claims Review or other Review and that it has concluded that it is, in fact, independent and/or objective.

**F. Ineligible Persons**

1. Definitions. For purposes of this IA:

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 ambit 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://oig.hhs.gov>); and (ii) the General Services Administration's List of Parties Excluded from Federal Programs (available through the Internet at <http://www.epls.arnet.gov>)

2. Screening Requirements. Dr. Larossa shall ensure that all prospective and current associates, employees, contractors, and agents of Dr. Larossa are not Ineligible Persons. To ensure that such individuals and entities are not Ineligible Persons, Dr. Larossa shall require such individuals and entities to disclose immediately any debarment, exclusion, suspension, or other event that makes such individual or entity an Ineligible Person. Prior to engaging the services of such individuals and entities, Dr. Larossa shall screen such individuals and entities against the Exclusion Lists. In addition, Dr. Larossa shall:

a. Within 90 days after the Effective Date, review his list of individuals and entities identified in Section III.F.2 against the Exclusion Lists; and

b. Review his list of individuals and entities identified in Section III.F.2 against the Exclusion Lists annually.

Dr. Larossa shall maintain documentation demonstrating that: (1) he has checked the Exclusion Lists (e.g., print screens from search results) and determined that such individuals or entities are not Ineligible Persons; and (2) has required individuals and entities to disclose if they are an Ineligible Person (e.g., employment applications).

Nothing in this Section affects the responsibility of (or liability for) Dr. Larossa to refrain from billing Federal health care programs for services of the Ineligible Person.

3. Removal Requirement. If Dr. Larossa has notice that any individual or entity in one of the positions identified in Section III.F.2 has become an Ineligible Person, Dr. Larossa shall remove such individual or entity from responsibility for, or involvement with, Dr. Larossa's business operations related to the Federal health care programs and shall remove such individual or entity from any position for which the individual's or entity's compensation or the items or services rendered, ordered, or prescribed by

the individual or entity 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 individual or entity is reinstated into participation in the Federal health care programs.

4. Pending Charges and Proposed Exclusions. If Dr. Larossa has notice that an individual identified in Section III.F.2 is charged with a criminal offense that falls within the ambit of 42 U.S.C. § 1320a-7(a), or an individual or entity identified in Section III.F.2 is proposed for exclusion during his, her or its employment, involvement or contract term, Dr. Larossa shall take all appropriate actions to ensure that the responsibilities of that individual or entity has not and shall not adversely affect the quality of care rendered to any beneficiary, patient, or resident, or the accuracy of any claims submitted to any Federal health care program.

G. Notification of Government Investigation or Legal Proceedings

Within 30 days after discovery, Dr. Larossa shall notify OIG, in writing, of any ongoing investigation or legal proceeding known to Dr. Larossa conducted or brought by a governmental entity or its agents involving an allegation that Dr. Larossa 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. Dr. Larossa 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 proceedings, if any.

H. Reporting

1. Overpayments

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

b. *Reporting of Overpayments.* If, at any time, Dr. Larossa identifies or learns of any Overpayment, Dr. Larossa shall notify the payor (e.g., Medicare fiscal intermediary or carrier) within 30 days

after identification of the Overpayment and take remedial steps within 60 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. Also, within 30 days after identification of the Overpayment, Dr. Larossa shall repay the Overpayment to the appropriate payor to the extent such Overpayment has been quantified. If not yet quantified within 30 days after identification, Dr. Larossa shall notify the payor at that time of his 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, and for Medicare contractors shall include the information contained on the Overpayment Refund Form, provided as Attachment 1 to Appendix B to this IA.

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.

2. Reportable Events.

a. *Definition of Reportable Event.* For purposes of this IA, a "Reportable Event" means anything that involves:

- i. a substantial Overpayment; or
- ii. 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.

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

b. *Reporting of Reportable Event.* If Dr. Larossa determines (after a reasonable opportunity to conduct an appropriate review or investigation of the allegations) through any means that there is a Reportable Event, Dr. Larossa shall notify OIG, in writing, within 30 days after making the determination that the Reportable Event exists. The report to OIG shall include the following information:

i. If the Reportable Event results in an Overpayment, the report to OIG shall be made at the same time as the notification to the payor required in Section III.H.1, and shall include all of the information on the Overpayment Refund Form, as well as:

(A) the payor's name, address, and contact person to whom the Overpayment was sent; and

(B) the date of the check and identification number (or electronic transaction number) by which the Overpayment was repaid/refunded;

ii. a complete description of the Reportable Event, including the relevant facts, persons involved, and legal and Federal health care program authorities implicated;

iii. a description of Dr. Larossa's actions taken to correct the Reportable Event; and

iv. any further steps Dr. Larossa plans to take to address the Reportable Event and prevent it from recurring.

#### **I. Third Party Billing**

Future Contract with Third Party Biller. If, at any time during the term of this IA, Dr. Larossa contracts with a third party billing company to submit claims to the Federal health care programs, at least 30 days prior to executing the contract, Dr. Larossa shall submit a certification indicating whether he has an ownership or control interest (as defined in 42 U.S.C. § 1320a – 3(a)(3)) in the third party billing company and whether he is employed by or acts as a consultant to the third party billing company.

Within 30 days after Dr. Larossa contracts with the third party billing company, Dr. Larossa shall obtain a certification from the third party billing company that the company: (i) is presently in compliance with all Federal health care program requirements as they relate to the submission of claims to Federal health care programs; (ii) has a policy of not employing any person who is excluded, debarred, suspended or otherwise ineligible to participate in Medicare or other Federal health

care programs to perform any duties related directly or indirectly to the preparation or submission of claims to Federal health care programs; and (iii) provides the required training in accordance with Section III.D of the IA for those employees involved in the preparation and submission of claims to Federal health care programs.

If Dr. Larossa contracts with a new third party billing company during the term of this IA, Dr. Larossa shall, within 30 days of entering into such contract, obtain and send to OIG the certification described in this Section III.I.2.

#### **IV. NEW BUSINESS UNITS OR LOCATIONS**

In the event that, after the Effective Date, Dr. Larossa changes locations or sells, closes, purchases, or establishes a new business unit or location related to the furnishing of items or services that may be reimbursed by Federal health care programs, Dr. Larossa shall notify OIG of this fact as soon as possible, but no later than 30 days after the date of change of location, sale, closure, purchase, or establishment. This notification shall include the address of the new business unit or location, phone number, fax number, Medicare Provider number, other provider identification number and/or supplier number, and the corresponding contractor's name and address that issued each number. Each new business unit or location and all Covered Persons at each new business unit or location shall be subject to the applicable requirements in this IA.

Prior to Dr. Larossa becoming an employee or contractor with another party related to the furnishing of items or services that may be reimbursed by Federal health care programs, Dr. Larossa shall notify that party of this IA. This notification shall include a copy of the IA, a statement indicating the remaining term of the IA, and a summary of Dr. Larossa's obligations under the IA. In addition, Dr. Larossa shall notify OIG of such relationship in his next Annual Report.

#### **V. REPORTS**

##### **A. Implementation Report**

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



1. the Compliance Contact's name, address, and phone number, a description of any other job responsibilities performed by the Compliance Contact, and the date the Compliance Contact was appointed;
2. a copy of the notice Dr. Larossa posted in his office as required by Section III.B, a description of where the notice is posted, and the date the notice was posted;
3. a copy of the written Policies and Procedures required by Section III.C of this IA and the date these Policies and Procedures were implemented and distributed;
4. a copy of all training materials used for the training session(s) required by Section III.D, a description of the training, including a summary of the topics covered, the length of each session, and a schedule of when the training session(s) were held;
5. the name and qualifications of the IRO(s), a summary/description of all engagements between Dr. Larossa and the IRO, including, but not limited to, any outside financial audits, compliance program engagements, or reimbursement consulting, and the proposed start and completion dates of the first annual Claims Review and Unallowable Cost Review, if applicable;
6. a copy of the IRO's engagement letter, including the length of the engagement;
7. a certification from the IRO regarding its professional independence and/or objectivity with respect to Dr. Larossa;
8. a description of Dr. Larossa's process to screen Covered Persons to determine if they are ineligible;
9. a summary of personnel actions (other than hiring) taken pursuant to Section III.F, the name, title and responsibilities of any person who is determined to be an Ineligible Person under Section III.F, and the actions taken in response to the obligations set forth in Section III.F;
10. a list of all Dr. Larossa'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 Provider number(s), provider identification number(s), and/or supplier number(s), and the name and address of each contractor to which Dr. Larossa currently submits claims;

11. if Dr. Larossa became an employee or contractor with another party related to the furnishing of items or services that may be reimbursed by Federal health care programs, Dr. Larossa shall inform OIG of the name, location, relationship, and its responsibilities with respect Dr. Larossa's employment or contract;

12. a certification by the Compliance Contact that:

a. the written Policies and Procedures required by Section III.C of this IA have been developed, are being implemented, and have been distributed to all Covered Persons; and that all Covered Persons have executed the written Policies and Procedures certification in accordance with the time frame required by Section III.C;

b. all Covered Persons have completed the applicable training required by Section III.D; and that all Covered Persons have executed the applicable training certification(s) in accordance with the time frame required by Section III.D;

c. all associates, employees, contractors, and agents that were hired or engaged since the execution of the IA were screened against the Exclusion Lists and asked to disclose if they are excluded, debarred, suspended, or are otherwise considered an Ineligible Person, prior to entering into their relationship with Dr. Larossa, as required by Section III.F; and

d. all current associates, employees, contractors, and agents of Dr. Larossa were screened against the Exclusion Lists within 90 days after the Effective Date, as required by Section III.F, and the date(s) of the screening.

13. a certification signed by Dr. Larossa certifying (a) to the best of his knowledge, except as otherwise described in the Implementation Report, Dr. Larossa is in compliance with all of the requirements of this IA and (b)

Dr. Larossa has reviewed the Implementation Report and has made a reasonable inquiry regarding its content and believes that the information is accurate and truthful.

**B. Annual Reports**

Dr. Larossa shall submit to OIG Annual Reports with respect to the status of, and findings regarding, his compliance activities for each of the five Reporting Periods (Annual Report).

Each Annual Report shall, at a minimum, include:

1. any change in the name, address, phone number, or job responsibilities of Dr. Larossa's Compliance Contact;
2. any changes to the posted notice and the reason for such changes;
3. a copy of any new compliance-related Policies and Procedures;
4. a summary of any changes or amendments to the written Policies and Procedures required by Section III.C and the reason(s) for such changes (e.g., change in contractor policies);
5. a copy of all training materials used for the training session(s) required by Section III.D (to the extent they have not already been provided as part of the Implementation Report); a description of the training, including a summary of the topics covered; the length of each session; and a schedule of when the training session(s) was held;
6. a complete copy of all reports prepared pursuant to the IRO's Claims Review or other Review required by Section III.E (and, if applicable for the first Annual Report, a copy of the certification described in Section III.E.4.b);
7. if applicable, a certification by Dr. Larossa stating that he does not currently and has not submitted any cost reports to any Federal health care programs since this IA was executed;
8. Dr. Larossa's response and corrective action plan(s) related to any issues raised or recommendations made by the IRO, as a result of the Review(s)

performed pursuant to Section III.E;

9. a summary/description of all engagements between Dr. Larossa and the IRO, including, but not limited to, any outside financial audits, compliance program engagements, or reimbursement consulting, if different from what was submitted as part of the Implementation Report;
10. a certification from the IRO regarding its professional independence and/or objectivity to Dr. Larossa;
11. a description of Dr. Larossa's process to screen Covered Persons to determine if they are ineligible (to the extent it has changed from the Implementation Report);
12. a summary of personnel actions/other than hiring taken pursuant to Section III.F; the name, titles and responsibilities of any person who is determined to be an Ineligible Person under Section III.F; and Dr. Larossa's actions taken in response to the obligations set forth in Section III.F;
13. a summary describing any ongoing investigation or legal proceeding required to have been reported pursuant to Section III.G. 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;
14. a summary of Reportable Events (as defined in Section III.H) identified during the Reporting Period and the status of any corrective and preventative action relating to all such Reportable Events;
15. 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: Medicare, Medicaid, and other Federal health care programs;
16. a copy of the certification from the third party billing company required by Section III.I, if applicable;
17. a description of all changes to the most recently provided list of Dr. Larossa's locations (including addresses) as required by Section IV. Include the corresponding phone numbers; fax numbers, each location's

Medicare Provider Number(s), other provider identification number(s) and/or supplier number(s) if applicable, and the name and address of the contractor that issued each number;

18. if Dr. Larossa became an employee or contractor with another party related to the furnishing of items or services that may be reimbursed by Federal health care programs, Dr. Larossa shall inform OIG of the name, location, relationship, and the responsibilities with respect Dr. Larossa's employment or contract;

19. A certification, where appropriate, by the Compliance Contact that certifies that:

a. the written Policies and Procedures have been reviewed during the Reporting Period, as required by Section III.B, and that all Covered Persons have executed the written Policies and Procedures certification in accordance with the time frame required by Section III.C;

b. all Covered Persons have completed the applicable training required by Section III.D and that all Covered Persons have executed the applicable training certification(s) in accordance with the time frame required by Section III.D;

c. all associates, employees, contractors, and agents that were hired, engaged or otherwise involved with Dr. Larossa during the Reporting Period have been screened against the Exclusion Lists and asked to disclose if they are excluded, debarred, suspended, or are otherwise considered an Ineligible Person, prior to entering into their relationship with Dr. Larossa, as required by Section III.F;

d. all associates, employees, contractors, and agents (employed, engaged or otherwise involved with Dr. Larossa for the entire Reporting Period) were screened against the Exclusion Lists during the Reporting Period, in accordance with Section III.F and the date(s) they were screened;

e. Dr. Larossa has complied with his obligations under the Settlement Agreement: (i) 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; (ii) not to charge to or otherwise seek payment from Federal or State payors for unallowable costs (as defined in the Settlement Agreement); and (iii) to identify and adjust any past charges or claims for unallowable costs.

20. a certification signed by Dr. Larossa certifying that: (a) to the best of his knowledge, except as otherwise described in the applicable Report, Dr. Larossa is in compliance with all of the requirements of this IA; and (b) Dr. Larossa has reviewed the Annual Report and has made a reasonable inquiry regarding its content and believes that the information is accurate and truthful.

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. Designation of Information**

Dr. Larossa shall clearly identify any portions of his submissions that he 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. Dr. Larossa 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 IA 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

Compliance Contact: Judith Lubin  
2000 Washington Street  
Newton Lower Falls, MA 02462  
(617) 244-1311  
(617) 244-2819 (Facsimile)

Unless otherwise specified, all notifications and Reports required by this IA shall 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.

**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 Dr. Larossa's books, records, and other documents and supporting materials and/or conduct on-site reviews of any of Dr. Larossa's locations for the purpose of verifying and evaluating: (a) Dr. Larossa's compliance with the terms of this IA; and (b) Dr. Larossa's compliance with the requirements of the Federal health care programs in which he participates. The documentation described above shall be made available by Dr. Larossa to OIG or its duly authorized representative(s) at all reasonable times for inspection, audit, or reproduction. Furthermore, for purposes of this provision, OIG or its duly authorized representative(s) may interview any of Dr. Larossa's employees, contractors, or agents 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. Dr. Larossa shall assist OIG or its duly authorized representative(s) in contacting and arranging interviews with such individuals upon OIG's request. Dr. Larossa's employees may elect to be interviewed with or without a representative of Dr. Larossa present.

**VIII. DOCUMENT AND RECORD RETENTION**

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

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 Dr. Larossa prior to any release by OIG of information submitted by Dr. Larossa pursuant to its obligations under this IA and identified upon submission by Dr. Larossa as trade secrets, or information that is commercial or financial and privileged or confidential, under the FOIA rules. With respect to such releases, Dr. Larossa shall have the rights set forth at 45 C.F.R. § 5.65(d).

X. BREACH AND DEFAULT PROVISIONS

Dr. Larossa is expected to fully and timely comply with all of his IA obligations.

A. Stipulated Penalties for Failure to Comply with Certain Obligations

As a contractual remedy, Dr. Larossa and OIG hereby agree that failure to comply with certain obligations set forth in this IA (unless a timely written request for an extension has been requested and approved in accordance with Section B below) 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 \$750 (which shall begin to accrue on the day after the date the obligation became due) for each day Dr. Larossa fails to:
  - a. have a Compliance Contact in accordance with the requirements of Section III.A;
  - b. establish and/or post a notice in accordance with the requirements of Section III.B;
  - c. establish, implement, maintain, distribute and/or update the written Policies and Procedures in accordance with the requirements of Section III.C;
  - d. establish and implement a training program in accordance with the requirements of Section III.D;



- e. obtain a certification from the third party biller, send the third party biller certification to OIG in accordance with the requirements of Section III.I, or notify OIG within 30 days prior to Dr. Larossa obtaining an ownership or control interest (as defined in 42 U.S.C. § 1320a-3(a)(3)) in, or becoming employed by, or becoming a consultant to, any third party billing company;
  - f. engage an IRO in accordance with the requirements of Section III.E.1.a and Appendix A;
  - g. submit the IRO's annual Claims Review Report and any other required Review in accordance with the requirements of Section III.E and Appendix B;
  - h. obtain and/or maintain the following documentation: Policies and Procedures certifications in accordance with the requirements of Section III.C, training certification(s) in accordance with the requirements of Section III.D, and/or documentation of screening and disclosure requirements in accordance with the requirements of Section III.F.2;
  - i. screen current or prospective associates, employees, contractors or agents in accordance with the requirements of Section III.F; or require associates, employees, contractors or agents to disclose if they are debarred, excluded, suspended or are otherwise considered an Ineligible Person in accordance with the requirements of Section III.F; or
  - j. notify OIG of a Government investigation or legal proceeding, in accordance with the requirements of Section III.G.
2. A Stipulated Penalty of \$1,000 (which shall begin to accrue on the day after the date the obligation became due) for each day Dr. Larossa fails to submit the Implementation Report or the Annual Reports to OIG in accordance with the requirements of Section V by the deadlines for submission.
3. A Stipulated Penalty of \$750 for each day Dr. Larossa fails to grant access to the information or documentation as required in Section VII. (This Stipulated Penalty shall begin to accrue on the date Dr. Larossa fails

to grant access.)

4. A Stipulated Penalty of \$5,000 for each false certification submitted by or on behalf of Dr. Larossa as part of his Implementation Report, Annual Reports, additional documentation to a Report (as requested by OIG), or as otherwise required by this IA.

5. A Stipulated Penalty of \$750 for each day Dr. Larossa fails to comply fully and adequately with any obligation of this IA. OIG shall provide notice (Notice) to Dr. Larossa stating the specific grounds for its determination that Dr. Larossa has failed to comply fully and adequately with the Agreement obligation(s) at issue and steps the Dr. Larossa shall take to comply with the IA. (This Stipulated Penalty shall begin to accrue 10 days after the date Dr. Larossa 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 1-4 of this Section.

**B. Timely Written Requests for Extensions**

Dr. Larossa 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 IA. 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 Dr. Larossa 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 business days after Dr. Larossa 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 business 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 Dr. Larossa 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 Dr. Larossa of: (a) Dr. Larossa's failure to comply; and (b) OIG's exercise of its contractual right to demand payment of the Stipulated Penalties (this notification is referred to as the "Demand Letter").

2. Response to Demand Letter. Within 10 days of the receipt of the Demand Letter, Dr. Larossa shall either: (a) cure the breach to OIG's satisfaction and pay the applicable Stipulated Penalties; or (b) send in writing to OIG a request for 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 Dr. Larossa elects to request an ALJ hearing, the Stipulated Penalties shall continue to accrue until Dr. Larossa 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 IA and shall be grounds for exclusion under Section X.D.

3. Form of Payment. Payment of the Stipulated Penalties shall be made by certified or cashier's check, payable to: "Secretary of the Department of Health and Human Services," and submitted to OIG at the address set forth in Section VI.

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 Dr. Larossa has materially breached this IA, 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 IA**

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

a. a failure by Dr. Larossa to report a Reportable Event, take corrective action and make the appropriate refunds, as required in Section III.H;

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

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.E.

2. Notice of Material Breach and Intent to Exclude. The parties agree that a material breach of this IA by Dr. Larossa constitutes an independent basis for Dr. Larossa's exclusion from participation in the Federal health care programs. Upon a determination by OIG that Dr. Larossa has materially breached this IA and that exclusion is the appropriate remedy, OIG shall notify Dr. Larossa of: (a) Dr. Larossa's material breach; and (b) OIG's intent to exercise its contractual right to impose exclusion (this notification is hereinafter referred to as the "Notice of Material Breach and Intent to Exclude").

3. Opportunity to Cure. Dr. Larossa shall have 30 days from the date of receipt of the Notice of Material Breach and Intent to Exclude to demonstrate to OIG's satisfaction that:

a. Dr. Larossa is in compliance with the obligations of the IA cited by OIG as being the basis for the material breach;

b. the alleged material breach has been cured; or

c. the alleged material breach cannot be cured within the 30-day period, but that: (i) Dr. Larossa has begun to take action to cure the material breach; (ii) Dr. Larossa is pursuing such action with due diligence; and (iii) Dr. Larossa has provided to OIG a reasonable timetable for curing the material breach.

4. Exclusion Letter. If at the conclusion of the 30-day period, Dr. Larossa fails to satisfy the requirements of Section X.D.3, OIG may exclude Dr. Larossa from participation in the Federal health care programs. OIG shall notify Dr. Larossa in writing of its determination to exclude Dr. Larossa (this letter shall be referred to hereinafter 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 Dr. Larossa's receipt of the Exclusion Letter. The exclusion shall have national effect and shall

also apply to all other Federal procurement and non-procurement programs. Reinstatement to program participation is not automatic. After the end of the period of exclusion, Dr. Larossa may apply for reinstatement, by submitting a written request for reinstatement in accordance with the provisions at 42 C.F.R. §§ 1001.3001-3004.

### Dispute Resolution

1. Review Rights. Upon OIG's delivery to Dr. Larossa of its Demand Letter or of its Exclusion Letter, and as an agreed-upon contractual remedy for the resolution of disputes arising under this IA, Dr. Larossa 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 IA. 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 the 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.

2. Stipulated Penalties Review. Notwithstanding any provision of Title 42 of the United States Code or Chapter 42 of the Code of Federal Regulations, the only issues in a proceeding for Stipulated Penalties under this IA shall be: (a) whether Dr. Larossa was in full and timely compliance with the obligations of this IA for which OIG demands payment; and (b) the period of noncompliance. Dr. Larossa shall have the burden of proving his 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 IA and orders Dr. Larossa to pay Stipulated Penalties, such Stipulated Penalties shall become due and payable 20 days after the ALJ issues such a decision unless Dr. Larossa 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 Chapter 42 of the Code of Federal Regulations, the only issues in a proceeding for exclusion based on a material breach of this IA shall be:

- a. whether Dr. Larossa was in material breach of this IA;
- b. whether such breach was continuing on the date of the Exclusion Letter; and
- c. whether the alleged material breach could not have been cured within the 30-day period, but that: (i) Dr. Larossa had begun to take action to cure the material breach within that period; (ii) Dr. Larossa has pursued and is pursuing such action with due diligence; and (iii) Dr. Larossa provided to OIG within that period a reasonable timetable for curing the material breach and Dr. Larossa has followed the timetable.

For purposes of the exclusion herein, exclusion shall take effect only after an ALJ decision favorable to OIG, or, if the ALJ rules for Dr. Larossa, only after a DAB decision in favor of OIG. Dr. Larossa's election of his contractual right to appeal to the DAB shall not abrogate OIG's authority to exclude Dr. Larossa 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 Dr. Larossa 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. Dr. Larossa shall waive his right to any notice of such exclusion if a decision upholding the exclusion is rendered by the ALJ or DAB. If the DAB finds in favor of Dr. Larossa, he 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 IA agree that the DAB's decision (or the ALJ's decision if not appealed) shall be considered final for all purposes under this IA.

**XI. EFFECTIVE AND BINDING AGREEMENT**

Consistent with the provisions in the Settlement Agreement pursuant to which this IA is entered, and into which this IA is incorporated, Dr. Larossa and OIG agree as follows:

- A. This IA shall be binding on the successors, assigns, and transferees of Dr. Larossa;
- B. This IA shall become final and binding on the date the final signature is obtained on the IA;
- C. Any modifications to this IA shall be made with the prior written consent of the parties to this IA;
- D. OIG may agree to a suspension of Dr. Larossa's obligations under this IA in the event of Dr. Larossa's cessation of participation in Federal health care programs. If Dr. Larossa withdraws from participation in Federal health care programs and is relieved of his IA obligations by OIG, Dr. Larossa shall notify OIG 30 days in advance of Dr. Larossa's intent to reapply as a participating provider or supplier with any Federal health care program. Upon receipt of such notification, OIG shall evaluate whether the IA shall be reactivated or modified.
- E. All requirements and remedies set forth in this IA are in addition to, and do not effect, (1) Dr. Larossa's responsibility to follow all applicable Federal health care program requirements or (2) the Government's right to impose appropriate remedies for failure to follow applicable program requirements.
- F. The undersigned Dr. Larossa signatories represent and warrant that they are authorized to execute this IA. The undersigned OIG signatory represents that he is signing this IA in his official capacity and that he is authorized to execute this IA.

IN WITNESS WHEREOF, the parties hereto affix their signatures:

**John T. Larossa, M.D.**



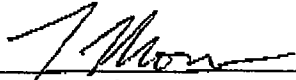
Dr. John T. Larossa  
2000 Washington Street  
Newton Lower Falls, MA

3/8/05

Date



**OFFICE OF INSPECTOR GENERAL OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES**

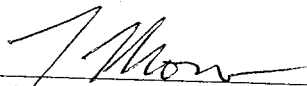


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Lewis Morris  
Chief Counsel to the Inspector General  
Office of Counsel to the Inspector General  
Office of Inspector General  
U. S. Department of Health and Human Services

3/29/05

\_\_\_\_\_  
Date

OFFICE OF INSPECTOR GENERAL OF THE DEPARTMENT OF HEALTH AND HUMAN SERVICES



\_\_\_\_\_  
Lewis Morris  
Chief Counsel to the Inspector General  
Office of Counsel to the Inspector General  
Office of Inspector General  
U. S. Department of Health and Human Services

3/29/05  
Date

## APPENDIX A INDEPENDENT REVIEW ORGANIZATION

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

### A. IRO Engagement.

Dr. Larossa 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/or objective fashion, as set forth in Paragraph D. Within 30 days after OIG receives written notice of the identity of the selected IRO, OIG will notify Dr. Larossa if the IRO is unacceptable. Absent notification from OIG that the IRO is unacceptable, Dr. Larossa may continue to engage the IRO.

If Dr. Larossa engages a new IRO during the term of the IA, this IRO shall also meet the requirements of this Appendix. If a new IRO is engaged, Dr. Larossa shall submit the information identified in Section V.A.5 of the IA to OIG within 30 days of engagement of the IRO. Within 30 days after OIG receives written notice of the identity of the selected IRO, OIG will notify Dr. Larossa if the IRO is unacceptable. Absent notification from OIG that the IRO is unacceptable, Dr. Larossa may continue to engage the IRO.

### B. IRO Qualifications.

The IRO shall:

1. assign individuals to conduct the Claims Review and, if applicable, Unallowable Cost Review engagement(s) who have expertise in the billing, coding, reporting and other requirements applicable to physicians and in the general requirements of the Federal health care program(s) from which Dr. Larossa 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 (e.g., CCA, CCS, CCS-P, CPC, RRA, etc.) 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 IA on a timely basis.

C. IRO Responsibilities.

The IRO shall:

1. perform each Claim Review and, if applicable, Unallowable Cost Review, in accordance with the specific requirements of the IA;
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., fiscal intermediary or carrier);
4. respond to all OIG inquiries in a prompt, objective, and factual manner; and
5. prepare timely, clear, well-written reports that include all the information required by Appendix B.

D. IRO Independence/Objectivity.

The IRO must perform the Claims Review in a professionally independent and/or objective fashion, as appropriate to the nature of the engagement, taking into account any other business relationships or engagements that may exist between the IRO and Dr. Larossa.

E. IRO Removal/Termination.

1. *Provider.* If Dr. Larossa terminates his IRO during the course of the engagement, Dr. Larossa must submit a notice explaining his reasons to OIG no later than 30 days after termination. Dr. Larossa must engage a new IRO in accordance with Paragraph A of this Appendix.

2. *OIG Removal of IRO.* In the event OIG has reason to believe that the IRO does not possess the qualifications described in Paragraph B, is not independent and/or 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 Dr. Larossa to engage a new IRO in accordance with Paragraph A of this Appendix.

Prior to requiring Dr. Larossa to engage a new IRO, OIG shall notify Dr. Larossa 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, Dr. Larossa may request a meeting with OIG to discuss any aspect of the IRO's qualifications, independence or performance

of its responsibilities and to present additional information regarding these matters. Dr. Larossa shall provide any additional information as may be requested by OIG under this Paragraph in an expedited manner. OIG will attempt in good faith to resolve any differences regarding the IRO with Dr. Larossa prior to requiring Dr. Larossa to terminate the IRO. However, the final determination as to whether or not to require Dr. Larossa to engage a new IRO shall be made at the sole discretion of OIG.

## APPENDIX B CLAIMS REVIEW

### A. Claims Review.

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

a. Overpayment: The amount of money Dr. Larossa has received in excess of the amount due and payable under any Federal health care program requirements.

b. Item: Any discrete unit that can be sampled (e.g., code, line item, beneficiary, patient encounter, etc.)

c. Paid Claim: A code or line item submitted by Dr. Larossa and for which Dr. Larossa has received reimbursement from the Medicare program.

d. Population: For the first Reporting Period, the Population shall be defined as all Items for which a code or line item has been submitted by or on behalf of Dr. Larossa and for which Dr. Larossa has received reimbursement from Medicare (i.e., Paid Claim) during the 12-month period covered by the first Claims Review.

For the remaining Reporting Periods, the Population shall be defined as all Items for which Dr. Larossa has received reimbursement from Medicare (i.e., Paid Claim) during the 12-month period covered by the Claims Review.

To be included in the Population, an Item must have resulted in at least one Paid Claim.

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 Sample 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 Items in the sample.

2. *Other Requirements.*

a. Paid Claims without Supporting Documentation. For the purpose of appraising Items included in the Claims Review, any Paid Claim for which Dr. Larossa cannot produce documentation sufficient to support the Paid Claim shall be considered an error and the total reimbursement received by Dr. Larossa for such Paid Claim shall be deemed an Overpayment. Replacement sampling for Paid Claims with missing documentation is not permitted.

b. Replacement Sampling. Considering the Population shall consist only of Paid Claims and that Items with missing documentation cannot be replaced, there is no need to utilize alternate or replacement sampling units.

c. Use of First Samples Drawn. For the purposes of all samples (Discovery Sample(s) and Full Sample(s)) discussed in this Appendix, the Paid Claims associated with the Items selected in each first sample (or first sample for each strata, if applicable) 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 Sample or Full Sample).

B. Claims Review Report. The following information shall be included in the Claims Review Report for each Discovery Sample and Full Sample (if applicable).

1. *Claims Review Methodology.*

a. Sampling Unit. A description of the Item as that term is utilized for the Claims Review.

b. Claims Review Population. A description of the Population subject to the Claims Review.

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

d. Sampling Frame. A description of the sampling frame, which is the totality of Items from which the Discovery Sample and, if any, Full Sample has been selected and an explanation of the methodology used to identify the sampling frame. In most circumstances, the sampling frame will be identical to the Population.

e. 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).

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

## 2. *Statistical Sampling Documentation.*

a. The number of Items appraised in the Discovery Sample and, if applicable, in the Full Sample.

b. A copy of the printout of the random numbers generated by the "Random Numbers" function of the statistical sampling software used by the IRO.

c. A copy of the statistical software printout(s) estimating how many Items are to be included in the Full Sample, if applicable.

d. A description or identification of the statistical sampling software package used to select the sample and determine the Full Sample size, if applicable.

## 3. *Claims Review Findings.*

### a. Narrative Results.

i. A description of Dr. Larossa'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 Sample, and the results of the Full Sample (if any).



b. Quantitative Results.

- i. Total number and percentage of instances in which the IRO determined that the Paid Claims submitted by Dr. Larossa (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 Dr. Larossa.
- iii. Total dollar amount of all Overpayments in the sample.
- iv. Total dollar amount of paid Items included in the sample and the net Overpayment associated with the sample.
- v. Error Rate in the sample.
- vi. A spreadsheet of the Claims Review results that includes the following information for each Paid Claim appraised: beneficiary health insurance claim number, date of service, procedure code submitted, procedure code reimbursed, allowed amount reimbursed by payor, correct procedure 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. (See Attachment 1 to this Appendix.)

4. *Systems Review.* Observations, findings, and recommendations on possible improvements to the system(s) and process(es) that generated the Overpayment(s).

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.



OVERPAYMENT REFUND

TO BE COMPLETED BY MEDICARE CONTRACTOR

Date: Contractor Deposit Control # Date of Deposit: Contractor Contact Name: Phone # Contractor Address: Contractor Fax:

TO BE COMPLETED BY PROVIDER/PHYSICIAN/SUPPLIER

Please complete and forward to Medicare Contractor. This form, or a similar document containing the following information, should accompany every voluntary refund so that receipt of check is properly recorded and applied.

PROVIDER/PHYSICIAN/SUPPLIER NAME ADDRESS PROVIDER/PHYSICIAN/SUPPLIER # CHECK NUMBER# CONTACT PERSON: PHONE # AMOUNT OF CHECK \$ CHECK DATE

REFUND INFORMATION

For each Claim, provide the following:

Patient Name HIC # Medicare Claim Number Claim Amount Refunded \$ Reason Code for Claim Adjustment: (Select reason code from list below. Use one reason per claim)

(Please list all claim numbers involved. Attach separate sheet, if necessary)

Note: If Specific Patient/HIC/Claim #/Claim Amount data not available for all claims due to Statistical Sampling, please indicate methodology and formula used to determine amount and reason for overpayment:

For Institutional Facilities Only:

Cost Report Year(s) (If multiple cost report years are involved, provide a breakdown by amount and corresponding cost report year.)

For OIG Reporting Requirements:

Do you have a Corporate Integrity Agreement with OIG? Yes No

Reason Codes:

- Billing/Clerical Error: 01 - Corrected Date of Service, 02 - Duplicate, 03 - Corrected CPT Code, 04 - Not Our Patient(s), 05 - Modifier Added/Removed, 06 - Billed in Error, 07 - Corrected CPT Code; MSP/Other Payer Involvement: 08 - MSP Group Health Plan Insurance, 09 - MSP No Fault Insurance, 10 - MSP Liability Insurance, 11 - MSP, Workers Comp.(Including Black Lung), 12 - Veterans Administration; Miscellaneous: 13 - Insufficient Documentation, 14 - Patient Enrolled in an HMO, 15 - Services Not Rendered, 16 - Medical Necessity, 17 - Other (Please Specify)

Integrity Agreement John T. Larossa, M.D.