Guidelines for Requesting an Increase in Enrollment in a Predoctoral Dental Education Program

TIMING OF REQUESTS AND RESPONSE: Approval of an increase in enrollment in predoctoral dental education programs must be reported to the Commission if the program’s total enrollment increases beyond the enrollment at the last site visit or prior approval of enrollment increase. Upon submission of the enrollment increase report, a substantial increase in program enrollment as determined by preliminary review by the discipline-specific Review Committee Chair will require prior approval by CODA.

RATIONALE FOR GUIDELINES: These Guidelines were drafted to focus upon adequacy of programmatic resources in support of additional student enrollees. Enrollment increases are tracked to ensure over time total enrollment does not exceed the resources of the program.

The Commission must review the request prior to implementation. It should be noted that the requirement for prior approval for an increase in enrollment is commensurate with the Commission’s Program Change policy under which previous enrollment increases were reported.

Programs should be cognizant of the impending need for enrollment increases through short- and long-term planning and proactively request permission for the increase. The Commission will not consider retroactive permanent requests. Additionally, the Commission will not consider inter-cycle requests unless there are documented extenuating circumstances.

Requests should be sent to the Commission on Dental Accreditation (211 E. Chicago Avenue, 19th floor, Chicago, IL 60611-2678) for initial review by the Review Committee Chair and, as needed, by the Predoctoral Dental Education Review Committee and subsequent review and approval by the Commission. The Predoctoral Dental Education Review Committee will review the request at the next regularly scheduled meeting. Reports submitted by June 1 will be considered at the Summer Commission meeting, and reports submitted by December 1 will be considered at the Winter Commission meeting.

POLICY ON MISSED DEADLINES: So that the Commission may conduct its accreditation program in an orderly fashion, all institutions offering programs accredited by the Commission are expected to adhere to deadlines for requests for program information. Programs/institutions must meet established deadlines to allow scheduling of regular or special site visits and for submission of requested information. Program information (i.e. self-studies, progress reports, annual surveys or other kinds of accreditation-related information requested by the Commission) is considered an integral part of the accreditation process. If an institution fails to comply with the Commission's request, or a prescribed deadline, it will be assumed that the institution no longer wishes to participate in the accreditation program. In this event, the Commission will immediately notify the chief executive officer of the institution of its intent to withdraw the accreditation of the program(s) at its next scheduled meeting.

Revised: 2/16; Reaffirmed: 8/15; 8/10, 7/07, 7/01, 5/88
FORMAT: The report must be clear and concise and must follow the “Required Documentation” and “Mechanics” sections illustrated within this guideline. Reports that fail to adhere to the stated guidelines may be returned to the program.

REQUIRED DOCUMENTATION (9 areas): Program directors (deans) must ensure that the proposed enrollment increases does not jeopardize the program’s ability to meet the Accreditation Standards.

In order to build and maintain calibration of evaluating requests for reportable enrollment increases, the following documentation must be submitted with the request for enrollment increase:

1. Date the program plans to increase enrollment.
2. The enrollment at the time of the most recent site visit.
3. The current enrollment in the program. Indicate the current enrollment in each year of the program and the projected enrollment. Indicate whether the proposed increase in enrollment or on a permanent increase.
4. The ratio of attendings/teaching staff to students before and after the proposed increase.
5. A schedule after the proposed increase is in effect (typical month)-and-a schedule to indicate the (pre-) clinic coverage assignments of the faculty.
6. Support staff available to students after the proposed enrollment increase.
7. (Pre-) Clinical facility/resources: operatories, student work/study areas, computer access, etc.
8. A description of the availability of adequate patient experiences to ensure the program’s goals and objectives for training to competencies will be achieved following the increased enrollment. Submit current (past two years) and projected numbers of patients by procedure type, including an accounting for the increased student enrollment. Additionally, provide minimum, mean, and maximum patient experiences by procedure type, for the preceding graduating class.
9. Explanation of how any off-campus sites may be involved in the proposed enrollment increase. Note: If new off-campus sites may be involved in the enrollment increase being reported, then the Policy and Guidelines for Off-Campus Sites must also be followed.

Omission of any of these nine (9) documentation areas may postpone Commission action on the request for increase in enrollment.
MECHANICS: The following **must** be observed in preparing the request.

1. **Cover page** must include
   a. name and address of the institution;
   b. program title;
   c. name, title, telephone number, e-mail address, and signature of individual preparing the request (this is typically the program director);
   d. name, title, and signature of the chief executive officer of the institution (the chief executive officer of the institution sponsoring the program must be copied on the letter transmitting the request to the Commission).

   **The electronic copy must include a signed cover/verification page and must conform to the Commission’s electronic submission guidelines.**

2. If documentation is extensive, a list of what is provided should be included. The actual items can be provided in **one (1) separate document that conforms to the electronic submission guidelines.**

3. **One (1) electronic copy** must be submitted following the Electronic Submission Guidelines. (Separate document) Failure to comply with these guidelines will constitute an incomplete report.

*Institutions/Programs are expected to follow Commission policy and procedure on privacy and data security, including those related to compliance with the Health Insurance Portability and Accountability Act (HIPAA). The Commission’s statement on HIPAA, as well as the Privacy and Data Security Summary for Institutions/Programs (PDF), are found in the Policies/Guidelines section of the Commission’s website at http://www.ada.org/en/coda/policies-and-guidelines/hipaa/. Programs that fail to comply with CODA’s policy will be assessed an administrative fee of $4000.*

**POLICY ON PREPARATION AND SUBMISSION OF DOCUMENTS TO THE COMMISSION:** All institutions offering programs accredited by the Commission are expected to prepare documents that adhere to guidelines set forth by the Commission on Dental Accreditation, including required verification signatures by the institution’s chief executive officer. These documents may include, but are not limited to, self-study, responses to site visit/progress reports, initial accreditation applications, reports of program change, and transfer of sponsorship and exhibits. The Commission’s various guidelines for preparing and submitting documents, including electronic submission, can be found on the Commission’s website or obtained from the Commission staff.

In addition, all institutions must meet established deadlines for submission of requested information. Any information that does not meet the preparation or submission guidelines or is
received after the prescribed deadlines may be returned to the program, which could affect the accreditation status of the program.

**Electronic Submission of Accreditation Materials:** All institutions will provide the Commission with an electronic copy of all accreditation documents and related materials, which conform to the Commission’s Electronic Submission Guidelines. Electronic submission guidelines will be provided to programs. Accreditation documents and related materials must be complete and comprehensive.

Documents that fail to adhere to the stated Guidelines for Submission will not be accepted and the program will be contacted to submit a corrected document. In this case, documents may not be reviewed at the assigned time which may impact the program’s accreditation status.

**Compliance with Health Insurance Portability and Accountability Act (HIPAA) (Excerpt):** The program’s documentation for CODA must not contain any patient protected health information (PHI) or sensitive personally identifiable information (PII). If the program submits documentation that does not comply with the policy on PHI or PII, CODA will assess an administrative processing fee of $4,000 per program submission to the institution; a program’s resubmission that continues to contain PHI or PII will be assessed an additional $4,000 administrative processing fee.

Adopted 1/20; Formerly Policy on Electronic Submission of Accreditation Materials, Commission Policy and Procedure Related to Compliance with the Health Insurance Portability and Accountability Act (HIPAA) and Policy on Preparation and Submission of Reports to the Commission

**ANNOUNCEMENT OF REVIEW RESULTS:** The Commission’s actions to approve or deny the request for reportable enrollment increases in predoctoral education programs, as are other accreditation actions, will be transmitted to the institutions/programs within 30 days following the Winter (January/February) or Summer (July/August) meetings.

**DENIAL OF REQUESTS:** Requests will be denied if the program cannot ensure continued compliance with the Accreditation Standards as demonstrated by documentation of the major program resource areas identified in the Guidelines for Enrollment Increases in Predoctoral Dental Education Programs.

**OTHER CHANGES IN ENROLLMENT:** Decreases in enrollment on a one-time-only basis or on a permanent basis must be reported to the Commission, but do not require prior approval. In the case of one-time-only decreases, programs are advised to maintain clinical experiences for the enrollment number for which they are approved.

**ASSISTANCE:** Commission staff is available to answer questions about request preparation. They may be contacted toll-free at (800) 621-8099, extension 2721. Requests should be sent to: Commission on Dental Accreditation, 211 E. Chicago Avenue, 19th floor, Chicago, IL 60611-2678.
Commission on Dental Accreditation  
Privacy and Data Security Reminders

Protect sensitive personally identifiable information (“PII”) such as social security numbers, drivers’ license numbers, credit card numbers, account numbers, etc.

Security Reminder: Personally Identifiable Information

Before submitting any documents to CODA or to a CODA site visitor, an institution must:

• Review for PII and patient identifiers.
• Fully and appropriately redact any PII and patient identifiers.
• Make sure the redacted information is unreadable in hard copy and electronic form. You must use appropriate redaction methods to ensure personal information cannot be read or reconstructed.

CODA does not accept PII or patient identifiers in any materials submitted by a program.

Security Reminder: Patient Identifiers

Before submitting any information about a patient to CODA or to a CODA site visitor, you must thoroughly redact all 18 patient identifiers listed on the next page.

Examples of information about a patient:

• Dental records
• Rosters of procedures (procedure logs)
• Chart review records (chart audit records)
• Information from affiliated teaching institutions, to include items listed above
• Brochures with patient images and/or information
• Presentations with patient images and/or information
• Course materials (exams, lecture materials) with patient images and/or information

If even one identifier is readable, do not submit the information to CODA.

CODA does not accept documents containing PII or patient identifiers from institutions. Any PHI/PII that is necessary for CODA accreditation may only be reviewed by CODA site visitors when they are on-site at the institution.

When redacting identifiers, you must ensure that the information is unreadable and cannot be reconstructed in both hard copy and electronic form. For example, certain information redacted on a hard copy can become readable when the hard copy is scanned. Instead, it may be effective to use opaque cover-up tape on the hard copy, scan, and then ensure the redacted information on the scanned version is not visible/readable through the redaction.
1. **Sensitive Information.** To protect the privacy of individuals and to comply with applicable law, the Commission on Dental Accreditation (“CODA” or “the Commission”) **prohibits all programs/institutions from disclosing in electronic or hard copy documents** provided to CODA other than on-site during a site visit, any of the following information (“Sensitive Information” or “PII”):
   - Social Security number
   - Credit or debit card number or other information (e.g., expiration date, security code)
   - Drivers’ license number
   - Account number with a pin or security code that permits access
   - Health insurance information, such as policy number or subscriber I.D.
   - Medical information, such as information about an individual’s condition or treatment
   - Mother’s maiden name
   - Taxpayer ID number
   - Date of birth
   - Any data protected by applicable law (e.g., HIPAA, state data security law)
   - Biometric data, such as fingerprint or retina image
   - Username or email address, in combination with a password or security question that permits access to an online account

2. **Patient Identifiers.** Before submitting information about a patient to CODA other than on-site during a site visit, a program/institution **must remove the following data elements** of the individual, and of relatives, household members, and employers of the individual (the “Patient Identifiers”):
   1. Names, including initials
   2. Address (including city, zip code, county, precinct)
   3. Dates, including treatment date, admission date, age, date of birth, or date of death [a range of dates (e.g., May 1 – 31, 2015) is permitted provided such range cannot be used to identify the individual who is the subject of the information]
   4. Telephone numbers
   5. Fax numbers
   6. E-mail addresses
   7. Social Security numbers
   8. Medical record numbers
   9. Health plan beneficiary numbers
   10. Account numbers
   11. Certificate/license numbers
   12. Vehicle identifiers and serial numbers, including license plate numbers
13. Device identifiers and serial numbers
14. Web Universal Resource Locators (URLs)
15. Internet Protocol (IP) address numbers
16. Biometric identifiers (e.g., finger and voice prints)
17. Full face photographic images and comparable images
18. Any other unique identifying number, characteristic, or code:
   - that is derived from information about the individual
   - that is capable of being translated so as to identify the individual, or
   - if the mechanism for re-identification (e.g., the key) is also disclosed

In addition, the information provided to CODA cannot be capable of being used alone or in combination with other information to identify the individual.

3. Redaction. When removing any Sensitive Information or Patient Identifier from paper or electronic documents disclosed to CODA, programs/institutions shall **fully and appropriately** remove the data such that the data cannot be read or otherwise reconstructed. Covering data with ink is not an appropriate means of removing data from a hard copy document and may sometimes be viewable when such documents are scanned to an electronic format.

4. Administrative fee. *If the program/institution submits any documentation that does not comply with the directives noted above, CODA will assess an administrative fee of $4000 to the program/institution; a resubmission that continues to contain prohibited data will be assessed an additional $4000 fee.*
   - CODA Site Visitors and Commission volunteers are only authorized to access Sensitive Information and Patient Identifiers:
     o Onsite during a site visit, and
     o That are necessary for conducting the accreditation site visit
   - CODA Site Visitors and Commission volunteers may not download or make hard copies or electronic copies of Sensitive Information or Patient Identifiers.

**NOTE:** If a document includes fictitious information, which may otherwise appear to be Sensitive Information or Patient Identifiers, the program is expected to clearly mark the document as “Fictitious Example”.

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Adopted August 2014; Updated 2.20