



Defense Health Agency J-7 Continuing Education Program Office
Continuing Education Activities Disclosure Form

Name: \_\_\_\_\_

Title of Activity: \_\_\_\_\_

Date of Activity: \_\_\_\_\_

Please indicate your role: (Check all that apply) Presenter Planning Committee Moderator Reviewer

It is the policy of the Defense Health Agency, J-7, Continuing Education Program Office as accredited providers of continuing education (CE)/continuing medical education (CME) for health care professionals, to ensure balance, independence, objectivity, and scientific rigor in all CE/CME activities. In compliance with accreditation requirements, all persons in a position to influence the content of the CE/CME activity (i.e., planning committee members, reviewers, moderators and presenters) are required to disclose relevant financial/nonfinancial interests or other relationships they have with commercial interests, as well as, ineligible companies over the past 24months. A commercial interest is defined as any entity producing, marketing, re-selling, or distributing health care goods or services consumed by, or used on, patients. Providers of clinical service directly to patients are not considered to be commercial interests. Ineligible companies are defined as those whose primary business is producing, marketing, selling, re-selling, or distributing health care products used by or on patients.

Regarding your role in the educational activity (check one):

No, I do not have a relevant financial/nonfinancial interest or other relationship with a commercial interest to report in the last 24 months, as defined above.

Yes, I do have a relevant financial/nonfinancial interest or relationship with a commercial interest to report in the last 24 months, as defined above (provide information below). Note: there is no need to disclose the financial value of any affiliation.

Table with 3 columns: Type of Affiliation/Relationship, Name of Company(s), Has the Relationship Ended? (If financial relationship existed in the last 24 months, but has now ended, check below.)

Discussion of Off-label/Unapproved Use: CE/CME materials that provide information, in whole or in part, related to the off-label, unapproved (not FDA approved) use of products, procedures and/or devices must clearly acknowledge the unlabeled, experimental and/or investigational nature of their proposed uses to the audience.

Yes, I will be discussing off-labeled uses for purposes other than that for which the product use was approved by the FDA. I will inform the audience at the start of the CE/CME activity as required by accreditation guidelines.

If yes, list the product and manufacturer: \_\_\_\_\_

No, I will not be discussing off-label uses.

Not applicable to my role as a content reviewer, planner or moderator.

The information provided above must be disclosed to learners before the start of the activity.

Signature \_\_\_\_\_

Date \_\_\_\_\_