# #8

### COMPLETE

Collector: Web Link 1 (Web Link)

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Page 1: Disclosure of conflict of interest information

## Q1

Process:1. Complete the conflict of interest disclosure form and submit to the CPD provider organization or scientific planning committee, as directed.2. Disclosures must be made to the audience whether you do or do not have a relationship to disclose.3. Speakers must disclose conflicts verbally and in writing on a slide at the beginning of a presentation. All other individual's conflicts must be disclosed either in writing on a slide at the beginning of a presentation or be included in the written conference materials.4. Those responsible for developing or delivering content must ensure that the content and/or materials presented provide (where applicable) a balanced view across all relevant options related to the content area.5. The description of therapeutic options must utilize generic names (or both generic and trade names) and not reflect exclusivity and branding.

By clicking "NEXT," you confirm that you have read and agree with the above requirements.

## Page 2: Disclosure of conflict of interest form

## Q2

Name

marilia seelaender

### Q3

**Email** 

seelaend@icb.usp.br

Q4 Please select below: 08/13/2020

Today's Date

Q5 Moderator or session chair

What is your role in this CPD activity? Select all that apply.

Q6 Yes

Do you have a relationship with a for-profit and/or a not-forprofit organization to disclose?

Q7

Please indicate the for-profit or not-for-profit organization(s) with which you have/had a relationship over the previous two years and briefly describe the nature of that relationship. Answer format: NAME (description), NAME (description)

Any direct financial payments including receipt of honoraria

Funded grants or clinical trials

Nestlé do Brasil, technical report

Ajinomoto do Brasil, recipient of donation of amino

acids for clinical trial

Q8 N/A (I am not a speaker)

For speakers only: I intend to make therapeutic recommendations for medications that have not received regulatory approval (i.e. "off-label" use of medication).

Q9 N/A (I am not a speaker)

For speakers only: I acknowledge that the National Standard requires that any description of therapeutic options utilize generic names (or both generic and trade names), and not reflect exclusivity and branding.

Q10 I agree

By clicking "I agree" you are acknowledging that the above information is accurate and that you understand that this information will be publicly available.