Class II Medical Devices – Terms of Market Authorization (TMA) Attestation

To: Ad Standards

From: _____ (the "Advertiser")
(Name of advertiser)

Re: _____, _____
(Script name) *(Media)*

Date: _____

I hereby attest the indication(s) for use registered with the Medical Devices Directorate (Health Canada) for _____,
(Device name)

_____ is/are as follows:
(Licence number)

(list out each indication individually)

I understand that:

- The onus is on the Advertiser to ensure the list of indications is current, accurate and complete (every indication registered must be included above).
- Ad Standards Clearance Services has not evaluated the sufficiency of studies, research, testing or other documentation in support of the indications as part of the preclearance process.
- The responsibility is on the Advertiser to ensure all indications are sufficiently supported and that the advertising otherwise complies with all applicable laws.
- Ad Standards' review of the advertising does not constitute legal advice.
- This attestation is valid for so long as the medical device licence remains active *and* the indications remain unchanged. If, at any point, updates, (including revisions and/or additions to the indications), are made, a new attestation will be required and will supersede the current document.

I confirm that to the best of my knowledge and belief, the foregoing are true statements as of the date first noted above.

Signed: ……
(I have authority to bind the Advertiser)

Name: _____

Title: _____

Email: _____