
FORM:	55 – ESTROGEN ALONE SURVEY
Version:	2 – October 1, 2004
Description:	Self-administered; 8-page booklet; key entered at Clinical Center (CC).
When Used:	Mailed by CCs to all E-alone participants who were taking study pills at the time the E-Along intervention stopped in March 2004.
Purpose:	To gather data about symptoms and feelings, how participants managed them, and the use of medications/supplements for participants in the E-alone trial

GENERAL INSTRUCTIONS

1. The form is printed in both English (*Form 55*) and Spanish (*Form 55S*). Both in a key-enter format.
2. Place the participant's barcode label on the front page of the questionnaire and give to the participant to complete at home or in the CC during close-out.
3. In appointment reminder (letter or phone call) remind the participant to bring the completed questionnaire to her transition visit.
4. Review the form for completeness, looking for skipped pages. Ask the participant to complete pages she may have skipped. Do not review the questions with the participants.
5. Forward the form to Data Entry.
6. Data Entry: Review the form for completeness. Return to interview if one or more pages not completed. Scan the English version of the form. Key-enter the Spanish version of the form. Initial the first page of the questionnaire after data entry.
7. File in the participant's file.

Item Instructions

Date received	Date the CC receives the completed form.
Reviewed by	Standard 3-digit WHI employee ID.
Contact type	Mark appropriate box. (See common data items.)
Visit type	The contact at which the CC receives the completed form. Mark appropriate oval or box for visit type and number. (See common data items.)
Form administration	<p>Method used to administer form to participant:</p> <p>1 - Self: Participant completed form by herself.</p> <p>2 - Group: Participant completed the form with a group of other participants.</p> <p>3 - Interview: CC staff person completed <u>entire</u> form as interview.</p> <p>4 - Assistance: Participant needed partial assistance from CC staff or others to complete the form.</p>
1. Date study pills stopped	Month/Year (estimate if unsure).
2.1. - 2.42. Symptoms	Participant's view of severity of listed symptoms based on how much they interfere with her usual activities. Level of discomfort associated with each symptom listed.
3.1. – 3.25. Dealing with symptoms	Participant's view of how effective the behavior was in relieving her symptom(s). Level of effectiveness for each option listed.
4. Sexual activity	Don't want to answer/ No/Yes, self stimulation/Yes.
4.1. Reasons	Participant marks reasons for sexual inactivity.
4.2. – 4.3. Sexual interest	Participant rates experience of desire and arousal (includes don't want to answer).
5.1. – 5.6. Sexual experiences	Participant rates frequency of sexual experiences (includes don't want to answer).
6. Intercourse in last 3 months	No/Yes/Don't want to answer.
6.1. – 6.4. Experiences with intercourse in last 3 months	Participant rates frequency of intercourse and associated experiences (includes don't want to answer).
7. – 7.1. Bisphosphonates, currently	No/Yes /Don't know and marks preparations she is currently taking.
8. - 8.1. Natural hormones, currently	No/Yes /Don't know and marks all preparations she is currently taking.
9. - 9.1. Hormones since stopping study pills	No/Yes and marks reasons for taking hormones (mark all that apply).

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10. Current use of hormones and type No/Yes and marks type (mark all that apply; include don't know and other category).
11. - Current use of SERMS No/Yes (mark all types that apply; includes "other" category).
11.1.
12. Future research Scale representing participants interest in future research on a continuum, 0-4.
Comments Data Entry: Do not key-enter comments.