How to Collect Data for Secondary Caregivers

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DO's

- Let participants know answering questions is voluntary.
- Let participants know the "why". FSS can say, "When you provide us with your partner's information, we can better understand how we can support your entire family."
- Let participants know that whatever they answer it is not punitive; there is no judgment, and no changes in services will occur.
- Leave "Secondary Caregiver Questions" sheet for them to complete & you to collect on your next visit.
- Ask for secondary caregiver information

DON'T

- Assume a participant does not know the information about the secondary caregiver.

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"I was asked about the household income. Then was explained that it was not punitive and nothing would change no matter how it was answered. (…) it was made clear I would not get in trouble."