

Child Welfare Caregivers: An Evaluation of Access to Pediatric Health Care







•There are over 800,000 children in the foster care system

- 60% of foster children have chronic medical conditions (25% have 3 or more chronic conditions).
- And, yet foster children do not receive adequate health care.

(Casey Family Programs, n.d.; Halfon, Mendonca, & Berkowitz, 1995; Szilagyi, 1998)



The Adoption and Safe Families Act of 1997.

- -Expanded family preservation services.
- Increased the number of children in child welfare living at home.
- -Children living at home, in family maintenance, have similar health problems as those in foster care.



(Downs, Moore, McFadden, Michaud, & Costin, 2004)



What role do caregivers have in accessing and utilizing health care?

- –Caregivers may be the gatekeepers; however, little is known about their role.
- -Qualitative results from a focus group (N=55) suggest that foster caregivers may find barriers to accessing services.
- -There are no studies on pediatric access barriers for birth parents.

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(Pasztor el al., 2006)



•The goals of the present study were to:

- Examine the barriers caregivers report to accessing pediatric health care for children in the child welfare system.
- Determine if there are differences in access barriers based on caregiver type (birth parent, related foster parent, and unrelated foster parent).





•Design:

- Setting: The Community-Based Assessment and Treatment Clinic (CATC), a pediatric clinic only serving children in the child welfare system. It is located in East Los Angeles, CA next to a large county hospital.
- Format: A 20-minute interview conducted in either Spanish or English in a private room.
- Compensation: Subjects were awarded a gift certificates.



•Study Population:

- –Eligibility: Caregivers who are 1) over 18 years of age and 2) have a child who has received health care services in the US prior to their visit at CATC.
- -Sample: 237 reports from caregivers was collected between July 2006 and November 2006.





•Measure:

 An adapted survey tool developed by Flores to assess barriers to health care for Latino children.

Flores's 10 Common Access Problems

Difficulty understanding the doctor's explanations	Too difficult to make an appointment
Clinic hours inconvenient	Wait too long to see a doctor
Couldn't afford medical care	Transportation
Too far away	Rudeness of staff
Medical staff doesn't understand culture	Immigration problems



Sample Item

Before coming to the CATC clinic, I found it hard to bring my child in for medical care because the doctor's office hours were always at times when I was working and I couldn't take time off from work.

Strongly Agree-Agree-Uncertain-Disagree-Strongly Disagree



(Flores et al, 1998)



•Analyses:

- General linear models were used to evaluate the associations between caregiver type and accessto-care barriers.
- Covariates of child's age and caregiver ethnicity, citizenship, income, education, immigration status, and number of children were controlled for in each model.





•Descriptives of caregivers:

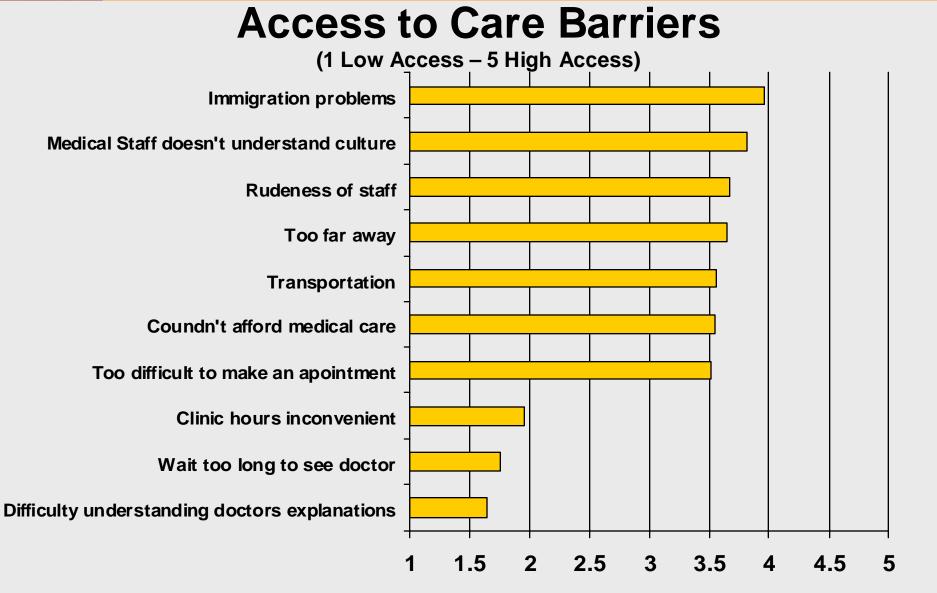
- 28% were birth parents, 25% related caregivers, and 46% were unrelated caregivers.
- 74% were Latina/o and 87% female.
- The average age of the child was 6.4 years.
- 86% of the caregivers rating their child's general health as excellent, very good or good and 14% rating their child's health status as poor or fair
- 78% of the caregivers had a regular physician for their child and 83% noted that their child's vaccinations were up-to-date.



- Significant differences in caregiver types:
 - Birth caregivers: younger, less likely to hold US citizenship, less likely to have a college education, and had a lower income.
 - Unrelated caregivers: more likely to be married, have more children, and less likely to be up-to-date on their vaccinations.

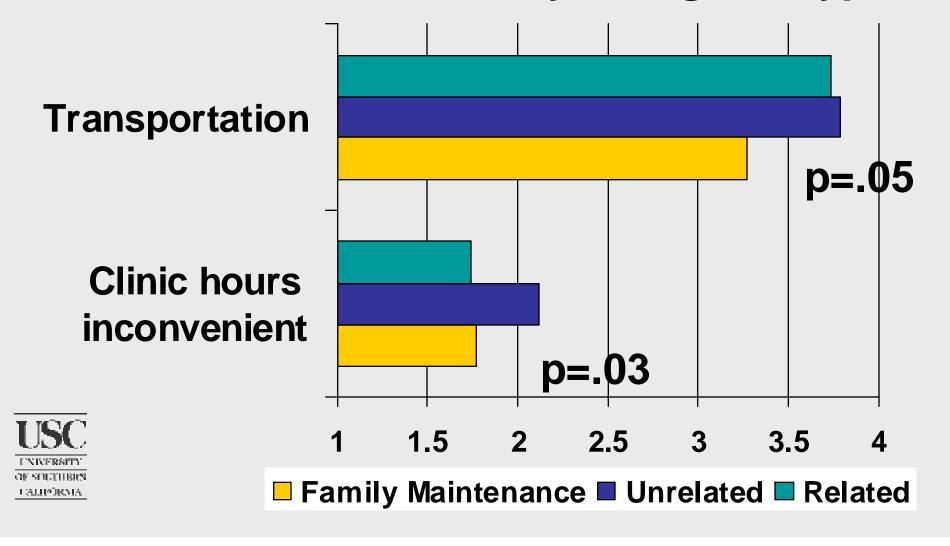






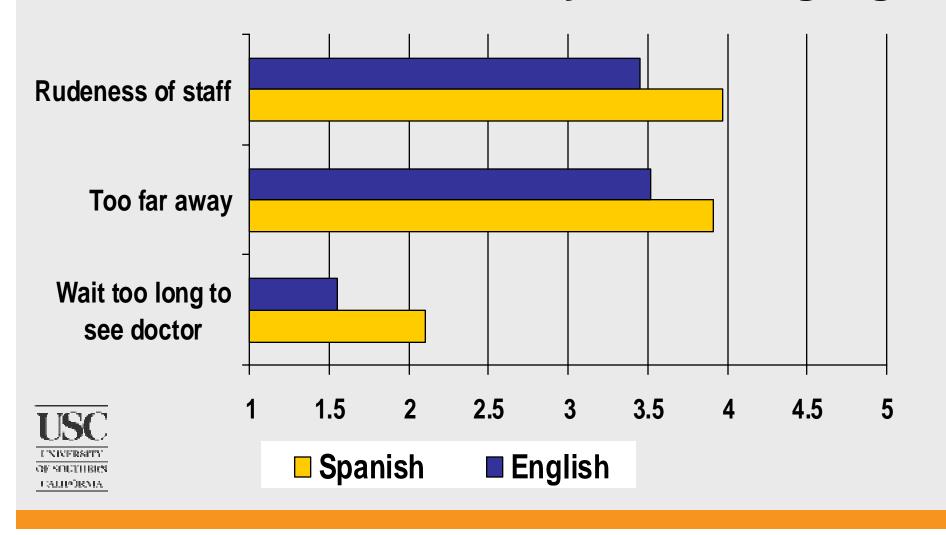


Access to Care Barriers by Caregiver Type





Access to Care Barriers by Form Language





•Limitations:

- Flores survey tool has no formal statistical reliability or validity data.
- Cross-sectional sample that might not generalize well to other child welfare populations.
- All caregivers interviewed were at a medical appointment; thus, they had accessed care. Those who were not at their appointments could not be included.
- Caregivers were asked to recall prior experiences.



•Discussion:

- Greatest barrier: Both English and Spanish speaking caregivers reported difficulty understanding doctor's explanations.
- While 94% of the caregivers reported health care coverage, caregivers still reported barriers to accessing care.
- Different medical care expectations for English and Spanish speaking caregivers.



•Implications:

- Based on the demographic differences of caregiver groups > Important to target birth parents.
- Because caregivers have unrealistically high perceptions of the health status of their children->
 Child welfare caseworkers need to carefully monitor adherence to well-child visits and illness-related appointments.





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