Flavor Programming During Infancy

Julie A. Mennella, PhD; Cara E. Griffin, BS; and Gary K. Beauchamp, PhD

ABSTRACT. Objective. Although individuals differ substantially in their flavor and food preferences, the source of such differences remains a mystery. The present experimental study was motivated by clinical observations that early experience with formulas establishes subsequent preferences.

Design. Infants whose parents had chosen to formula-feed them were randomized into 1 of 4 groups by the second week of life. One group was assigned to be fed a milk-based formula (Enfamil), whereas another was assigned to be fed (Nutramigen), a particularly unpleasant-tasting protein hydrolysate formula. The remaining groups were assigned to be fed Nutramigen for 3 months and Enfamil for 4 months; the timing of exposure differed between the groups. After 7 months of exposure, infants were videotaped on 3 separate days while feeding, in counterbalanced order, Enfamil, Nutramigen, and Alimentum, a novel hydrolysate formula.

Results. For each of the 4 interrelated measures of behavior (intake, duration of formula feeding, facial expressions, and mothers’ judgments of infant acceptance), previous exposure to Nutramigen significantly enhanced subsequent acceptance of both Nutramigen and Alimentum. Seven months of exposure led to greater acceptance than did 3 months.

Conclusions. The bases for clinical difficulties in introducing hydrolysate formulas during older infancy are clarified in this study. More broadly, variation in formula flavor provided a useful model for demonstrating experimentally the effects of long-term exposure differences on later acceptance. Such early variation, under more species-typical circumstances (eg, via exposure to different flavors in amniotic fluid and mothers’ milk), may underlie individual differences in food acceptability throughout the life span. Pediatrics 2004;113:840–845; protein hydrolysate, formula, taste, flavor, infants, programming, development, nutrition.

ABBREVIATIONS. E, Enfamil; N, Nutramigen; ANOVA, analysis of variance.

Sources of individual differences in food preferences and habits are one of the most fundamental mysteries of human behavior. Although genetic differences may underlie some of these differences,1,2 for omnivores such as humans, it is important that there not be too many genetically determined restrictions on what constitutes an acceptable food.3 Instead, as suggested by a growing body of data from other sensory and motor systems,4–6 experience may have important influences on later functioning and preferences. We hypothesized that there are sensitive periods during which the human infant is particularly likely to form flavor preferences and aversions that, in turn, serve as the foundation for lifelong food habits.7

Recently, we suggested that a particularly apt model system to explore potential early sensitive periods in flavor learning involves the inherent flavor variations characteristic of infant formulas and the ontogenic changes in acceptance/rejection of particular flavors.7–9 Within each of the 3 classes of commercially available formulas (ie, cow’s milk based, soy protein based, and hydrolyzed protein based), and in particular between the hydrolysate- and milk-based varieties, differences in sensory quality (flavor) are profound. Milk-based formulas are often described as having low levels of sweetness and being “sour and cereal-like,” whereas hydrolyzed protein-based formulas are of a most unpleasant character, with a bitter and sour taste profile, unpleasant odor volatiles, and a horrible after taste.10,11 The extreme unpalatability of hydrolysates, which supply protein nutrients in a “predigested” form, is likely caused by both its processing and composition, because many amino acids and small peptides taste sour and bitter and are characterized by unpleasant volatile components.12

There are striking developmental changes in infants’ acceptance of hydrolysate formulas. Infants <4 months old readily accept these formulas on first exposure, whereas older infants strongly reject them within the first few minutes of feeding.8,9,13 Clinicians report that infants who consume a hydrolysate formula from early infancy readily continue to accept it well after 5 months of age.7 These observations suggest that there is a profound change at ~4 months of age in the infants’ perception of these formulas and that early experience modifies later acceptance.

To test rigorously the suggestion derived from such cross-sectional data, we designed a longitudinal experimental study wherein infants were assigned randomly to different experience groups. This permitted precise control of exposure history and consequently is the ideal test of the hypothesis that prior exposure to a particular flavored formula impacts later acceptance of that and other formulas. More generally, this experimental study can provide a convenient and powerful model system for investigating the existence of sensitive periods in the development of human flavor preferences.
METHODS

Subjects

Mothers who had chosen previously to formula feed their term newborns were recruited by advertisements in local newspapers. When the infant was <3 weeks old and the mother’s decision not to lactate was well established, the mother-infant pairs (24% African American and 76% white) were randomized into 1 of 4 groups (Table 1) differing in the timing and type of formula (ie, Enfamil: milk-based formula [E]; Nutramigen: protein hydrolysate formula [N]) that the infant was fed during each month of the 7-month study. One group, EEEEEEE (hereafter referred to as the control group; n = 14), was assigned to a milk-based formula, Enfamil, whereas another group, NNNNNNN (n = 12), was assigned to the protein hydrolysate formula Nutramigen during the entire 7-month period of the study. The 2 other groups, NNNNEEEE (n = 15) and EENNNEE (n = 12), were assigned to feed Nutramigen for specified periods during their first 7 months of life, as indicated by the Ns and Es in the groups’ names. Infants were fed on demand and ad libitum. All procedures used in this study were approved by the Office of Regulatory Affairs at the University of Pennsylvania, and informed consent was obtained before entry into the study. Mothers were compensated for their participation in the study.

Monthly Procedures

At the start of the study and then again at the beginning of each 1-month cycle, mothers came to the Monell Center (Philadelphia, PA), where they were videotaped feeding their infants the formula consumed since the last visit. The infants’ weights and heights before feeding and the amount of formula consumed during this midday feed were recorded. These monthly evaluations were performed to ensure compliance with the study protocols, to chart infants’ acceptance of the assigned formulas, and to obtain accurate information on the introduction of cereal, fruits, and vegetables. The next month’s supply of formula (with the labels removed) then was distributed. Mothers were informed that the formula was either Enfamil or Nutramigen. Mothers completed questionnaires to evaluate their food and general neophobia at the start of each 4-week exposure period (± 1 week). Each infant was evaluated at the end of each month and at the end of the study at 7.5 months. The names of the groups refer to the protein [N] that the infant was fed during each month of the 7-month study. One group, EEEEEEE (hereafter referred to as the control group; n = 14), was assigned to a milk-based formula, Enfamil, whereas another group, NNNNNNN (n = 12), was assigned to the protein hydrolysate formula Nutramigen during the entire 7-month period of the study. The 2 other groups, NNNNEEEE (n = 15) and EENNNEE (n = 12), were assigned to feed Nutramigen for specified periods during their first 7 months of life, as indicated by the Ns and Es in the groups’ names. Infants were fed on demand and ad libitum. All procedures used in this study were approved by the Office of Regulatory Affairs at the University of Pennsylvania, and informed consent was obtained before entry into the study. Mothers were compensated for their participation in the study.

Test Procedures at the End of the 7-Month Exposure Period

By using methodologies established in our laboratory,8,9 infants were tested on 3 separate days within 1 week. At the same time of day and 4.1 ± 0.2 hours after their last formula feed, each infant was videotaped feeding the hydrolysate formula Nutramigen on 1 test day, the milk-based formula Enfamil on another day, and Alimentum (a novel hydrolysate formula to all infants) on yet another test day. Testing occurred under naturalistic conditions in which the infants determined the pacing and duration of feeding. The order of testing was counterbalanced between and within groups, and the mothers were not aware of the hypothesis or which formula was being fed to the infant during each of the 3 test sessions. The mothers refrained from talking or making faces during the feeding sessions to eliminate any potential influence of the mother’s verbal or facial responses on her infant’s behaviors15; replays of the videotapes verified that this indeed was the case. Mothers were instructed to feed the infants at their customary pace until the infant refused the bottle 3 consecutive times, using the criterion that the infant exhibited such behaviors as turning his or her head away, pushing the bottle away, crying, or becoming playful. The experimenter, who was unaware of the hypothesis of the study, sat behind the video camera, which was placed at the far corner of the testing room ~10 to 12 feet from the mother-infant dyad, and was out of view of the mother and her infant. Immediately after each feeding session, the mothers then rated how much they thought their infant liked the formula on a 9-point scale.17 Intermediate ratings were to be marked at the appropriate locations between the extremes such that ratings could range from 1 (did not like at all) to 9 (liked very much).

Trained raters who were unaware of the experimental conditions and hypothesis of the studies scored the videotaped records to determine the length of the formula feed and frequency of various facial expressions. During scoring, the sound was turned off so that the raters would not be influenced by the infants’ vocalizations. The Ekman and Friesen Facial Action Coding System,18 an anatomically based system that specifies facial movements in terms of minimally distinguishable actions of the facial muscles (termed action units), was used to code a variety of facial expressions. Based on previous research of the type of facial responses made by human infants and other primates to a variety of taste and olfactory stimuli that differed in hedonic valence,19-21 we determined the frequency of 4 negative facial expressions (ie, nose wrinkling, frowning, upper-lip raise, and gaping). Two observers individually scored the videotapes of 41 feedings selected at random. Reliability for scoring of each of the facial responses and the length of the feeding was >85% (P < .0001).

Statistical Analyses

For each infant, we determined the total intake (milliliters) and length (minutes) of each feed, the frequency of negative facial expressions during the first 2 minutes of feeding, and mothers’ ratings of their infants’ enjoyment of the formulas during each of the 3 test sessions conducted at the end of the 7-month exposure period. To determine whether there were significant differences among the 4 groups, we conducted separate repeated-measures analyses of variance (ANOVA) with group (n = 4) as the grouping factor and type of formula (n = 3) as the within-subjects factor. Significant effects in the ANOVA were probed by Tukey honest significant difference tests to determine whether the 3 exposure groups differed from each other as well as the control group. All summary statistics are expressed as mean ± standard error, and the level of significance was P < .05 for the ANOVA and P < .02 for the Tukey tests.

RESULTS

Subject Characteristics

There were no significant differences among the 4 groups in the ages of mothers and infants, the number of females/males, or the infants’ weights and lengths at the start of the study. There also were no significant group differences for any of the various measures of infant temperament, the age at which infants were introduced to solid foods, or maternal measures of food and general neophobia (all P values > .10).

TABLE 1. Description of Experimental Groups

<table>
<thead>
<tr>
<th>Infant Age at Start of Exposure, mo*</th>
<th>Experimental Groups</th>
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<tbody>
<tr>
<td>EEEEEE</td>
<td>NNNNN</td>
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<tr>
<td>0.5</td>
<td>Enfamil</td>
</tr>
<tr>
<td>1.5</td>
<td>Enfamil</td>
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<tr>
<td>2.5</td>
<td>Enfamil</td>
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<tr>
<td>3.5</td>
<td>Enfamil</td>
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<td>4.5</td>
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<tr>
<td>5.5</td>
<td>Enfamil</td>
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<tr>
<td>6.5</td>
<td>Enfamil</td>
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</tbody>
</table>

* Infants’ age at the start of each 4-week exposure period (± 1 week). Each infant was evaluated at the end of each month and at the end of the study at 7.5 months. The names of the groups refer to the month of life that infants were fed Nutramigen or Enfamil.
Infants’ Formula Acceptance Throughout the 7-Month Study

There were no significant differences between the groups in the infants’ acceptance of the formula that they were fed during the previous month throughout the 7-month study (F[3.49 df] = 0.91; P = .44; data not shown), thus indicating compliance with study procedures.

Infants’ Acceptance of Hydrolysate Formulas at 7.5 Months of Age

There was a significant interaction between groups and the infants’ acceptance of the 3 brands of formulas when tested at the end of the 7-month exposure period (intake: F[6.98 df] = 8.12 and P < .000000; duration of feed: F[6.98 df] = 5.60 and P < .000005). The 3 groups of Nutramigen-exposed infants drank significantly more and spent more time feeding Nutramigen and Alimentum when compared with those infants who were fed only Enfamil during the first 7 months of life (Table 2 and Fig 1; all P values < .05). However, exposure to Nutramigen for 7 months (group NNNEEEE) resulted in greater acceptance of the Nutramigen when compared with exposure for 3 months (groups NNNEEEE and EENNNNEE).

There was also a significant effect of group on the number of negative facial actions displayed while feeding the formulas (F[6.98 df] = 2.51; P < .05). Infants in groups NNNNNNN, NNNEEEE, and EENNNNEE made significantly less negative facial responses while ingesting Nutramigen when compared with infants exposed only to the milk-based Enfamil formula (Fig 1). Differences in facial responses made while feeding Nutramigen are also illustrated in Fig 2.

<table>
<thead>
<tr>
<th>Experimental Group</th>
<th>Intake, ml</th>
<th>Duration of feed, min</th>
<th>Frequency of negative facial action units during first 2 min of feed</th>
<th>Mothers’ ratings of infants’ acceptance of formula: (range: 1–9; 1 = did not enjoy at all)</th>
<th>No. of subjects (females/males)</th>
</tr>
</thead>
<tbody>
<tr>
<td>EEEEEE</td>
<td>160.9 ± 22.3</td>
<td>7.2 ± 1.3</td>
<td>3.9 ± 1.8</td>
<td>7.7 ± 0.6</td>
<td>14 (7.7)</td>
</tr>
<tr>
<td>NNNEEEE</td>
<td>237.1 ± 21.6*</td>
<td>12.0 ± 1.2</td>
<td>9.5 ± 1.5</td>
<td>8.5 ± 0.6*</td>
<td>15 (6.9)</td>
</tr>
<tr>
<td>EENNEEE</td>
<td>158.8 ± 24.1</td>
<td>7.3 ± 1.4</td>
<td>5.7 ± 1.3</td>
<td>7.3 ± 0.7*</td>
<td>12 (8.4)</td>
</tr>
<tr>
<td>NNNNNNN</td>
<td>119.1 ± 22.9</td>
<td>7.1 ± 1.4</td>
<td>6.4 ± 2.0</td>
<td>4.9 ± 0.7*</td>
<td>12 (5.7)</td>
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There was a significant group-by-formula interaction (F[6,96 df] = 9.96; P = .00000) in mothers’ perceptions of their infants’ behaviors that was consistent with the infants’ acceptance patterns (Table 2). Mothers of infants in the 3 groups who had previous exposure to Nutramigen were significantly more likely to report that their infant enjoyed feeding Nutramigen when compared with the infants who had never been exposed to Nutramigen. In addition, mothers of infants who were exposed only to Nutramigen during the first 7 months of life were significantly more likely to report that their infants liked feeding Alimentum but disliked feeding Enfamil when compared with the group never exposed to Nutramigen (P < .0001).

**DISCUSSION**

Previous exposure to a hydrolysate formula, Nutramigen, enhanced its later acceptance. Infants who were never fed Nutramigen during the first 7 months of life strongly rejected it when it was first offered at 7.5 months, whereas those who were regularly fed this formula during most of their infancy responded to it as if it were very acceptable at 7.5 months of age. Between these 2 extremes, infants fed Nutramigen either during their first 3 months of life or during months 3 through 5 were generally more accepting than those who had never been fed this formula but less so than those fed Nutramigen for the entire 7 months. No differences were evident between these 2 midexposure groups, providing no evidence for differential potency of exposure during different portions of early infancy.

**TABLE 2. Infants’ Feeding Behaviors and Mothers’ Perceptions After the 7-Month Exposure Period**

The amount of formula consumed, the duration of the feeding, the frequency of negative and positive facial actions displayed, and the mothers’ ratings of their infants’ acceptance of formula are shown for each of the 3 testing sessions in which 7.5-month-old infants consumed, in counterbalanced order, Enfamil, Nutramigen, and Alimentum. The groups differed in the type of formula consumed during the first 7 months of life. Group EEEEEE was assigned to be fed Enfamil and group NNNNNNN was assigned to be fed the protein hydrolysate formula Nutramigen during the entire 7-month period of the study. Groups NNNEEEE and EENNNNEE were assigned to feed Nutramigen for 3 months and Enfamil for 4 months; the timing of exposure differed between the groups. The values shown are means ± standard error.

* P < 0.05 for the comparison with the EEEEEE group.
† P < 0.05 for the comparison of group NNNEEEE or EENNNNEE with the NNNNNNN group.
Sensory Basis for Differential Responsiveness

The remarkable consistency among the 3 indices of acceptance (Fig 1) suggests that these measures reflect a common underlying factor that is most likely differential responses to the formulas based on their chemosensory attributes. Strengthening this conclusion are the data on expressive responses (mothers' judgments and analyses of facial expressions) that most likely directly reveal hedonic responses to sensory stimuli. Based on adult sensory profiling of these formulas, the 2 primary sensory pathways involved are likely taste and olfaction.7 Concerning taste, hydrolysate formulas taste more bitter and sour than most traditional milk-based formulas including Enfamil. Thus, it would not be surprising that hydrolysate formulas would be rejected more strongly, because newborns reject bitter and very sour stimuli.20–23 What makes this hypothesis questionable is that rejection does not occur until after 4 months of age. However, at least as it concerns bitterness perception, there are multiple classes of bitter compounds presumably recognized by members of a large family of bitter receptors.24 The bitterness of hydrolysate formulas, probably caused by bitterness of free amino acids and small peptides12 and possibly other substances produced during processing, may be detected by receptors that do not mature until the infant is several months old.

Concerning olfaction, that the cloying, unpleasant, nauseating flavor and aftertaste of hydrolysates can be reduced substantially if the olfactory component is eliminated by tasting hydrolysates with the nares held closed, implicates olfaction in the adults' and perhaps the infants' hedonic judgments. It has been suggested that although hedonic judgments for tastes are relatively "hard wired" or determined innately,25 hedonic responses to the olfactory components of flavor are influenced strongly by experience.26 Thus, the observed changes in response to the hydrolysate formulas with age and prior experience...
may depend on developmental and experiential changes (“olfactory imprinting”) in response to the volatile components of the formulas.

Persistence

There are indications that the effects of early experience to hydrolysate formulas may be long lived. In a prior study, we found that 4- to 5-year-old children who were fed hydrolysates during their infancy exhibited more positive responses to sensory attributes associated with them (eg, sour taste and aroma) several years after their last exposure to the formula when compared with same-aged children without such experience. In addition, clinical studies on adolescent children with phenylketonuria who went off their modified hydrolysate formula suggest that they can often, albeit with some difficulty, return to formulas on which they were reared as infants. Because these formulas are unpalatable to adolescents who have not had infant exposure, this relative ease of return could reflect the long-term effects of prior experience.

Generality

The effects observed in these studies are likely of substantially broad significance, revealing a fundamental feature of mammalian dietary learning. Studies of humans and other animals have shown that fetuses are bathed in flavored amniotic fluids that reflect in part the pregnant animals’ diet. After birth, mammals are exposed further to milk that is flavored by the diet the nursing mother is consuming. Consequently, even during very early development, mammals usually sample a large and varied set of flavor compounds. We suggest that these “natural” experiences with flavored amniotic fluid and milk serve to familiarize the young infant with flavors of the dietary constituents the mother consumes and establish them as acceptable and preferred. Culturally determined flavor preferences, one of the most enduring characteristics of an ethnic group, can be understood in the context of early flavor exposure. Flavors common to an ethnic group are experienced early in life, at a time when the pregnant or lactating mother is most likely being fed foods most characteristic of and most revered by that particular culture.

Clinical Implications

Some of the original impetus for this work came from questions raised by clinicians concerning the practical difficulties in introducing hydrolysate formulas to older infants as well as the long-term effects of early feeding. Our studies reveal the basis for these clinical impressions and suggest some remedies. First, if infants are to be placed on hydrolysate formulas, they should be introduced as early as possible and certainly before 4 months of age. A strategy of gradually introducing hydrolysates to older infants by mixing it with regular, milk-based formula at increasing proportions seems reasonable but to our knowledge has not been experimentally tested.

Second, caregivers who feed infants recognize hydrolysates are very unpalatable. They may feel that they are punishing their child by feeding such an offensively flavored formula. Our studies make it clear that this is not the case if infants are started early, however. The infant will not only accept it readily but will find it palatable.

Third, there is room to develop more palatable hydrolysate formulas. The first step in such a program should be to determine the specific sensory aspects of these formulas that are offensive to the naive infant and preferred by the exposed infant. Such information would help suggest a rational strategy for favorably modifying current formulations.

CONCLUSIONS

These data underscore one of the most fundamental differences in the experiences of the formula- and breastfed infants. To be sure, the nutrient composition of formula is held to standards and regulation that are based, in part, on the analyses of human milk. However, unlike breast milk, the flavors of all types of formulas are monotonous and lack sensory information on the dietary choices of the mother. Thus, the formula-fed infant is being deprived of rich and varied sensory experiences that at one time were common to all mammalian young. How this impacts on later food habits and flavor preferences remains to be determined, but it seems likely that the consequences could be profound.

ACKNOWLEDGMENTS

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REFERENCES

AGE OF PRIM PARA

“The average age at which American women are having their first child has climbed to a record 25.1, the government said. The rise reflects a drop in teen births and an increase in the number of women who are putting off motherhood until their 30s and 40s. The age of first-time American moms has risen steadily during the past three decades from an average of 21.4 in 1970. The latest figure, for 2002, was released by the Centers for Disease Control and Prevention.”


Submitted by Student
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