

## Effect of Long-term Administration of Immunomodulatory Food on Cancer Patients Completing Conventional Treatments

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**Key words :** Complementary medicine, Rice bran arabinoxylan derivative, Immunomodulatory food, Safety

### Summary

A study was conducted to investigate the effects of long-term administration of the immunomodulatory food modified arabinoxylan from rice bran (BioBran/MGN-3), on 16 cancer patients, mainly in stage IV with various conventional lesions, who had just undergone conventional cancer treatments, such as surgery, chemotherapy and radiotherapy. The main clinical observations were the safety and effect of BioBran/MGN-3 on the nutritional state of the patients, who were exhausted due to treatment. During the administration period, no decrease in body weight and leukocyte count or significant changes in leukogram were observed. Rather, the leukocyte count increased. In addition, most patients showed an increase in NK cell activity and a remarkable decrease in tumor markers.

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This paper is a translation of an article in *Clinical Pharmacology and Therapy*, Vol. 14/No. 3/May 2004.

## Introduction

In our clinic, complementary medicine is used in cancer patients who have completed surgery, chemotherapy, and irradiation therapy in order to improve QOL, prevent recurrence, and enhance life prolongation. We call the medicine "Ryo-yo." "Ryo" means treatment given in the clinic to enhance healing and immunity, and "yo" means daily care by the patients themselves to increase their self-healing capacity. For daily care, patients are trained in breathing, diet, and physical and mental health<sup>1)</sup>. The diet should be based on modern dietetics or grains and vegetables to enhance prophylactic effect. Functional foods are also used as part of the dietary therapy, but patients make their own decision about ingestion. Many functional foods are used to prevent decreased immunity and to reduce adverse reactions during cancer treatment. All our patients take up to 5 kinds of functional foods. Most contain ingredients equal or similar to those in foods taken every day. However, the form is as concentrates and capsules, granules, or tablets of partially purified ingredients in most cases. Thus, there is a possibility of ingesting larger quantities of some ingredients than those contained in foods. As it is reported that excessive ingestion of  $\beta$  carotene promotes lung cancer<sup>2)</sup>, sufficient attention should be paid to safety. In the present study, the effect of long-term administration of BioBran/MGN-3, most frequently used by our patients, was evaluated in 16 cancer patients with nutritional problems who had just completed conventional treatments, focusing particularly on the effect on leukocytes.

Table 1 Backgrounds of subjects

| Initials | Age | Sex    | Primary lesion                  | Study period                 |
|----------|-----|--------|---------------------------------|------------------------------|
| K.O.     | 56  | Male   | Stomach                         | January to July 2001         |
| I.R.     | 64  | Male   | Large intestine                 | March to September 2001      |
| M.T.     | 59  | Male   | Large intestine                 | March to September 2001      |
| K.K.     | 44  | Female | Breast                          | February to August 2001      |
| T.H.     | 58  | Female | Rectum                          | May to November 2001         |
| F.A.     | 46  | Female | Breast                          | July 2001 to January 2002    |
| T.S.     | 60  | Female | Stomach                         | August 2001 to February 2002 |
| K.H.     | 47  | Female | Breast                          | December 2001 to June 2002   |
| E.I.     | 44  | Male   | Biliary tract at hepatic portal | February to August 2002      |
| H.Y.     | 59  | Female | Large intestine                 | February to August 2002      |
| H.M.     | 77  | Female | Ovary                           | December 2001 to June 2002   |
| M.N.     | 72  | Female | Thyroid gland                   | January to July 2002         |
| Y.I.     | 44  | Male   | Lung                            | October 2001 to April 2002   |
| Y.H.     | 84  | Male   | Rectum                          | January to July 2002         |
| N.A.     | 39  | Female | Uterine cervix                  | March to September 2002      |
| K.M.     | 53  | Male   | Rectum                          | April to October 2002        |

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**Table 2 Changes in body weight (kg)**

| Initials | Before study | After study | Difference |
|----------|--------------|-------------|------------|
| K.O.     | 70.0         | 71.0        | +1.0       |
| I.R.     | 67.0         | 69.0        | +2.0       |
| M.T.     | 61.0         | 60.0        | -1.0       |
| K.K.     | 48.0         | 49.0        | +1.0       |
| T.H.     | 53.0         | 53.0        | 0          |
| F.A.     | 49.0         | 51.0        | +2.0       |
| T.S.     | 38.0         | 38.0        | 0          |
| K.H.     | 52.5         | 53.0        | +0.5       |
| E.I.     | 47.0         | 46.5        | -0.5       |
| H.Y.     | 50.0         | 51.0        | +1.0       |
| H.M.     | 44.0         | 44.0        | 0          |
| M.N.     | 46.5         | 47.0        | +0.5       |
| Y.I.     | 64.0         | 65.0        | +1.0       |
| Y.H.     | 59.0         | 60.0        | +1.0       |
| N.A.     | 45.0         | 46.5        | +1.5       |
| K.M.     | 68.0         | 68.0        | 0          |

**Table 3 Changes in leukocyte count and subsets**

| Initials | Leukocyte count (/mm <sup>3</sup> ) |                      |            | Neutrophil (%)        |                      |                | Lymphocyte (%)        |                      |                |
|----------|-------------------------------------|----------------------|------------|-----------------------|----------------------|----------------|-----------------------|----------------------|----------------|
|          | Before administration               | After administration | Difference | Before administration | After administration | Difference (%) | Before administration | After administration | Difference (%) |
| K.O.     | 5500                                | 6500                 | +1000      | 65.7                  | 76.2                 | +10.5          | 24.9                  | 19.5                 | -3.4           |
| I.R.     | 6100                                | 4400                 | -1700      | 69.8                  | 62.8                 | -7.0           | 24.1                  | 27.6                 | +3.5           |
| M.T.     | 3500                                | 4100                 | +600       | 56.4                  | 59.5                 | +3.1           | 27.2                  | 31.2                 | +4.0           |
| K.K.     | 3400                                | 3600                 | +200       | 60.8                  | 64.9                 | +4.1           | 22.7                  | 21.3                 | -1.4           |
| T.H.     | 5700                                | 5400                 | +300       | 51.9                  | 53.0                 | +1.1           | 42.0                  | 42.5                 | +0.5           |
| F.A.     | 2500                                | 3000                 | +500       | 57.0                  | 52.1                 | -4.9           | 24.5                  | 42.5                 | +18.0          |
| T.S.     | 3800                                | 4200                 | +400       | 40.0                  | 55.3                 | +15.3          | 56.0                  | 35.9                 | -20.1          |
| K.H.     | 4800                                | 4400                 | -400       | 80.0                  | 71.7                 | -8.3           | 11.0                  | 20.6                 | +9.6           |
| E.I.     | 2800                                | 3400                 | +600       | 57.9                  | 67.0                 | +9.1           | 25.6                  | 23.8                 | -1.8           |
| H.Y.     | 4200                                | 5400                 | +1200      | 50.5                  | 61.9                 | +11.4          | 33.7                  | 27.2                 | -6.5           |
| H.M.     | 3000                                | 3500                 | +500       | 54.6                  | 63.8                 | +9.2           | 30.3                  | 29.9                 | -0.4           |
| M.N.     | 7300                                | 6000                 | -1300      | 68.9                  | 62.2                 | -6.7           | 24.6                  | 28.7                 | +4.1           |
| Y.I.     | 3700                                | 5600                 | +1900      | 71.7                  | 82.0                 | +10.3          | 19.7                  | 11.5                 | -8.2           |
| Y.H.     | 5600                                | 5800                 | +200       | 64.0                  | 64.2                 | +0.2           | 25.2                  | 23.5                 | +1.7           |
| N.A.     | 5200                                | 4300                 | -900       | 80.0                  | 71.1                 | -8.9           | 13.5                  | 14.2                 | +0.7           |
| K.M.     | 5300                                | 5900                 | +600       | 44.8                  | 48.8                 | +4.0           | 35.7                  | 23.6                 | -12.1          |

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Table 4 Changes in leukocyte count and subsets from the normal ranges

|   | Leukocyte count       |                      | Neutrophil            |                      | Lymphocyte            |                      |
|---|-----------------------|----------------------|-----------------------|----------------------|-----------------------|----------------------|
|   | Before administration | After administration | Before administration | After administration | Before administration | After administration |
| L | 7                     | 4                    | -                     | -                    | 11                    | 13                   |
| N | 9                     | 12                   | 8                     | 5                    | 4                     | 3                    |
| H | -                     | -                    | 8                     | 11                   | 1                     | -                    |

Table 5 Categorization of changes in leukocyte count and subsets

|           | Leukocyte count | Neutrophil | Lymphocyte |
|-----------|-----------------|------------|------------|
| Increase  | 9               | 5          | 2          |
| No change | 4               | 5          | 10         |
| Decrease  | 3               | 6          | 4          |

Table 6 Changes in NK activity

| Patient's initials | Before administration | At 6 months of administration |
|--------------------|-----------------------|-------------------------------|
| K.O.               | 13.3                  | 31.2                          |
| I.R.               | 17.0                  | 47.5                          |
| M.T.               | 37.2                  | 50.3                          |
| K.K.               | 4.1                   | 45.3                          |
| T.H.               | 9.5                   | 34.8                          |
| F.A.               | 9.3                   | 39.5                          |
| T.S.               | 25.6                  | 38.6                          |
| K.H.               | 15.9                  | 63.6                          |
| E.I.               | 16.2                  | 39.4                          |
| H.Y.               | 9.6                   | 6.1                           |
| H.M.               | 22.5                  | 31.4                          |
| M.N.               | 31.4                  | 20.3                          |
| Y.I.               | 32.4                  | 32.0                          |
| Y.H.               | 30.3                  | 30.2                          |
| N.A.               | 20.3                  | 10.7                          |
| K.M.               | 8.2                   | 26.2                          |

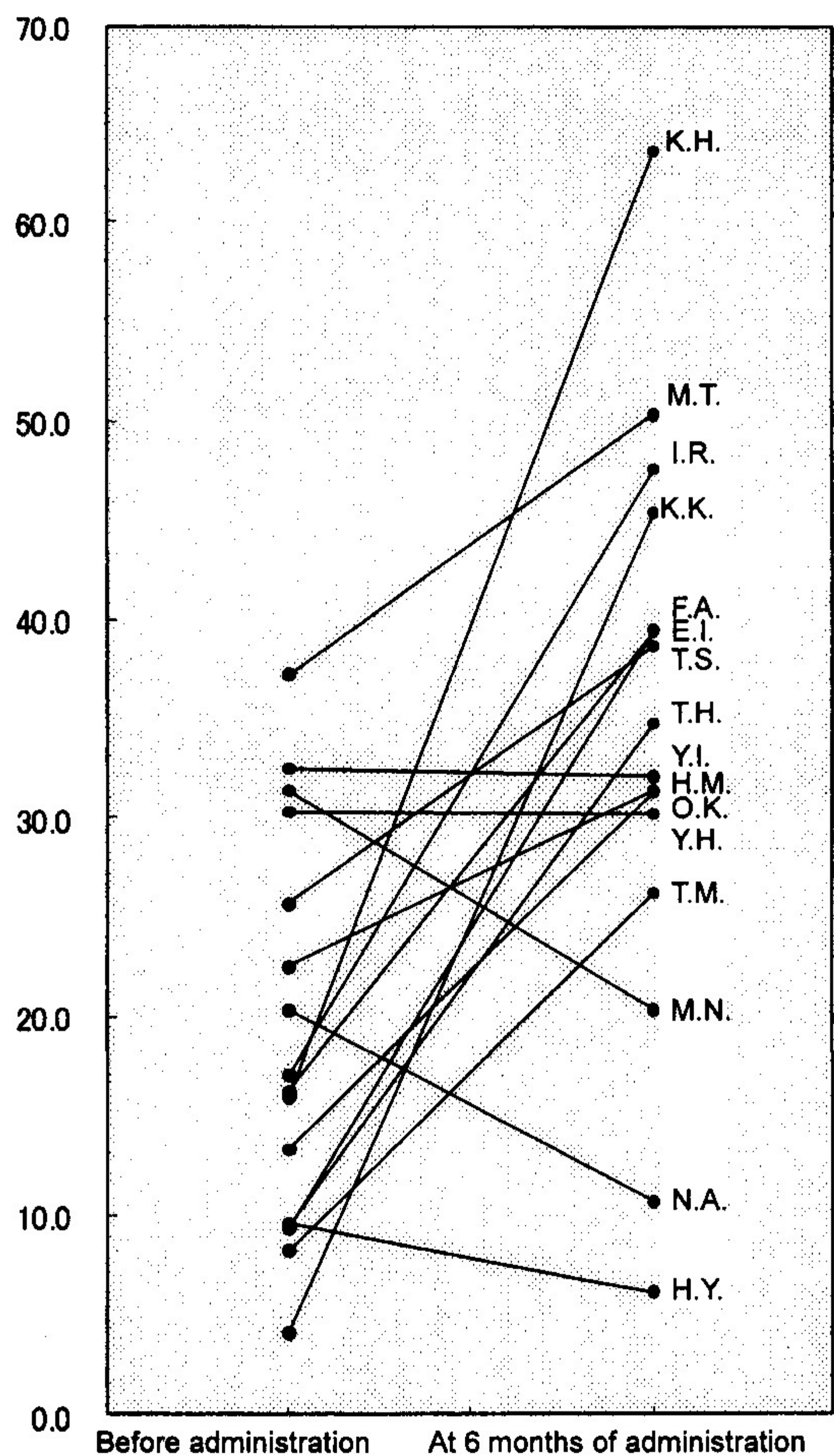


Fig. 1 Changes in NK activity

Methods

1. Patients and study period

The subjects were 16 cancer patients who met the criteria (1) to (3) below, and the study period was 6 months.

Table 1 shows the age, primary lesion, and study dates for each patient.

- 1) Cancer patients just after completion of surgery, irradiation therapy, and/or chemotherapy
- 2) Patients visiting this clinic for observation of outcome and care to improve QOL and prevent recurrence
- 3) Patients who consented to ingest BioBran/MGN-3 at 3 g/day.

2. Study items

The study items were body height and weight, leukocyte count and subsets (neutrophils, lymphocytes,

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monocytes, eosinophils, basophils, and band cells), NK cell activity, tumor markers, adverse reactions (abdominal pain, vomiting, and an enlarged feeling in the abdomen), and interruptions of administration and the reasons for interruption.

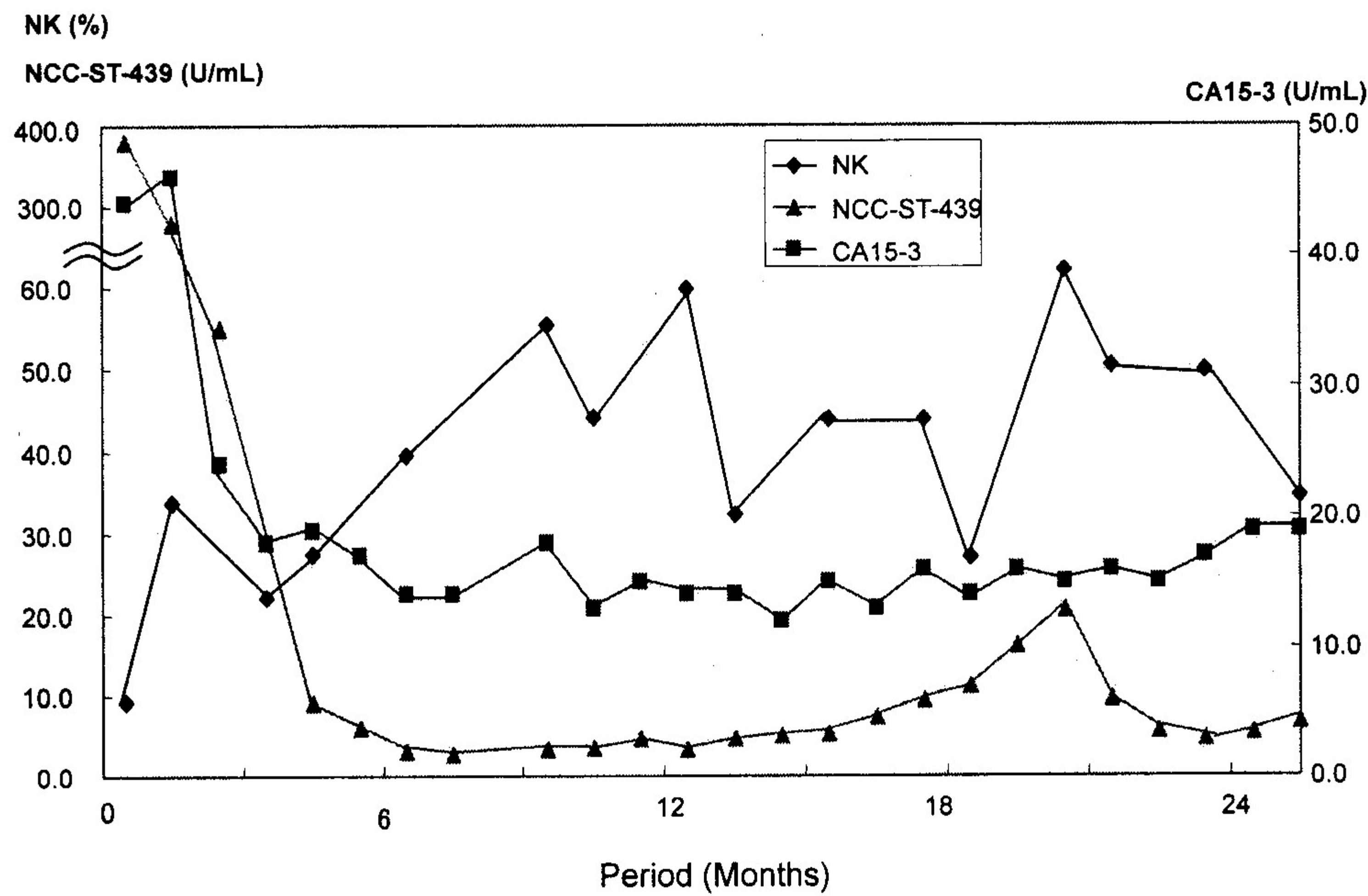


Fig. 2 Breast Cancer (Stage IV) F.A. (46) F.

Height was measured at the start of the study. Body weight and leukocyte count and subsets were monitored 3 times at the start of, during, and at the end of the study. NK activity and tumor markers were determined every month. Adverse reactions and interruptions of ingestion were checked throughout the study period.

3. Modified arabinoxylan from rice bran (BioBran/MGN-3)

The study material BioBran/MGN-3 is produced by partially hydrolyzing rice bran extract with enzymes. There are many reports on the physiological actions of MGN-3, the generic name of BioBran, including immunomodulation<sup>3-4)</sup>, active-oxygen scavenging<sup>5)</sup>, blood sugar control<sup>6)</sup> and reduction of adverse reactions to anticancer drugs<sup>7)</sup>.

Results

All 16 subjects completed the administration of BioBran/MGN-3 continuously during the study period.

1. Changes in body weight

Body weight increased in 10 patients, decreased in 2, and was unchanged in 4. The range of change was

within 4% for both increase and decrease. BioBran/MGN-3 had almost no effect on body weight. Table 2 shows these changes.

## 2. Changes in leukocytes

Changes in leukocyte counts and subsets were studied. Table 3 shows leukocyte counts and results for neutrophils and lymphocytes. The normal range is 4000-9000/mm<sup>3</sup> for leukocytes count, 40%-60% for neutrophils fraction, and 30%-45% for lymphocytes. Individual measurements before and after administration were divided into categories H (higher than the normal ranges), N (within the normal ranges), and L (lower than the normal ranges) (Table 4).

The changes in measurements were classified into the categories of increase (changes above 10% for leukocyte counts and 5% each for neutrophil and lymphocyte fractions), no change (changes within  $\pm 10\%$  and  $\pm 5\%$  each, respectively), and decrease (changes under -10% and -5% each, respectively) (Table 5).

The leukocyte count was generally low in the subjects of this study because they had just completed conventional treatments: it was below the normal range in 7 of 16 patients (44%).

After 6 months of BioBran/MGN-3 administration, leukocyte count increased in 9 of 16 patients, of whom 3 had a normal value. The fraction of neutrophils increased slightly, but no constant trend was observed. The lymphocyte fraction was low, and there was almost no change before and after administration. In one patient each, however, the value changed from a low level to the normal range and from a high level to the normal range. Overall, changes towards a healthy condition were observed, and no adverse changes were noted in the leukocyte profile for 6 months.

## 3. NK activity and tumor markers

The NK activity at the start of the study was  $\leq 30\%$  in 11 patients, 30%-50% in 3, and  $\geq 50\%$  in 2, and the proportion of patients with normal NK activity was 19%. After administration of BioBran/MGN-3, NK cell activity tended to increase, and was normal in 11 patients (69%). Tumor markers decreased in 10 (63%) after administration of BioBran/MGN-3.

Fig. 1 and Table 6 show changes in NK activity.

## 4. Adverse reactions

No adverse reactions to BioBran/MGN-3 were observed or reported by any of the subjects.

## 5. Cases with marked improvement in nutritional state

1) Patient: F.A., female, 46 years, recurrent breast cancer (stage IV)

The patient received a diagnosis of breast cancer in July 1998 and underwent surgery and hormonal treatment. After 2 years 6 months, she had metastases in the left iliac bone, lumbar vertebrae, and uterine body. A hysterectomy was performed and Taxol and Paraplatin administered for bone metastases. However, no improvement was observed, and metastases to the thoracic vertebrae and ribs occurred. She visited our clinic in July 2001, when the tumor markers CA15-3 and NCC-ST-439 were at the high concentrations of 44 U/mL and 369 ng/mL, respectively, and NK activity was at a low level of 9.3%. She had malaise, severe bone pain, and low QOL (PS2). She received our therapy while continuing administration of Paraplatin. BioBran/MGN-3 was taken at 3 g/day. NK cell activity increased to 33.7% at 1 month, and the levels of two tumor markers decreased rapidly at 2 months. By 7 months, pain due to bone metastases disappeared and

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malaise was reduced. Now, after 34 months (April 2004), she lives a normal life with QOL maintained (PS0) (Figure 2).

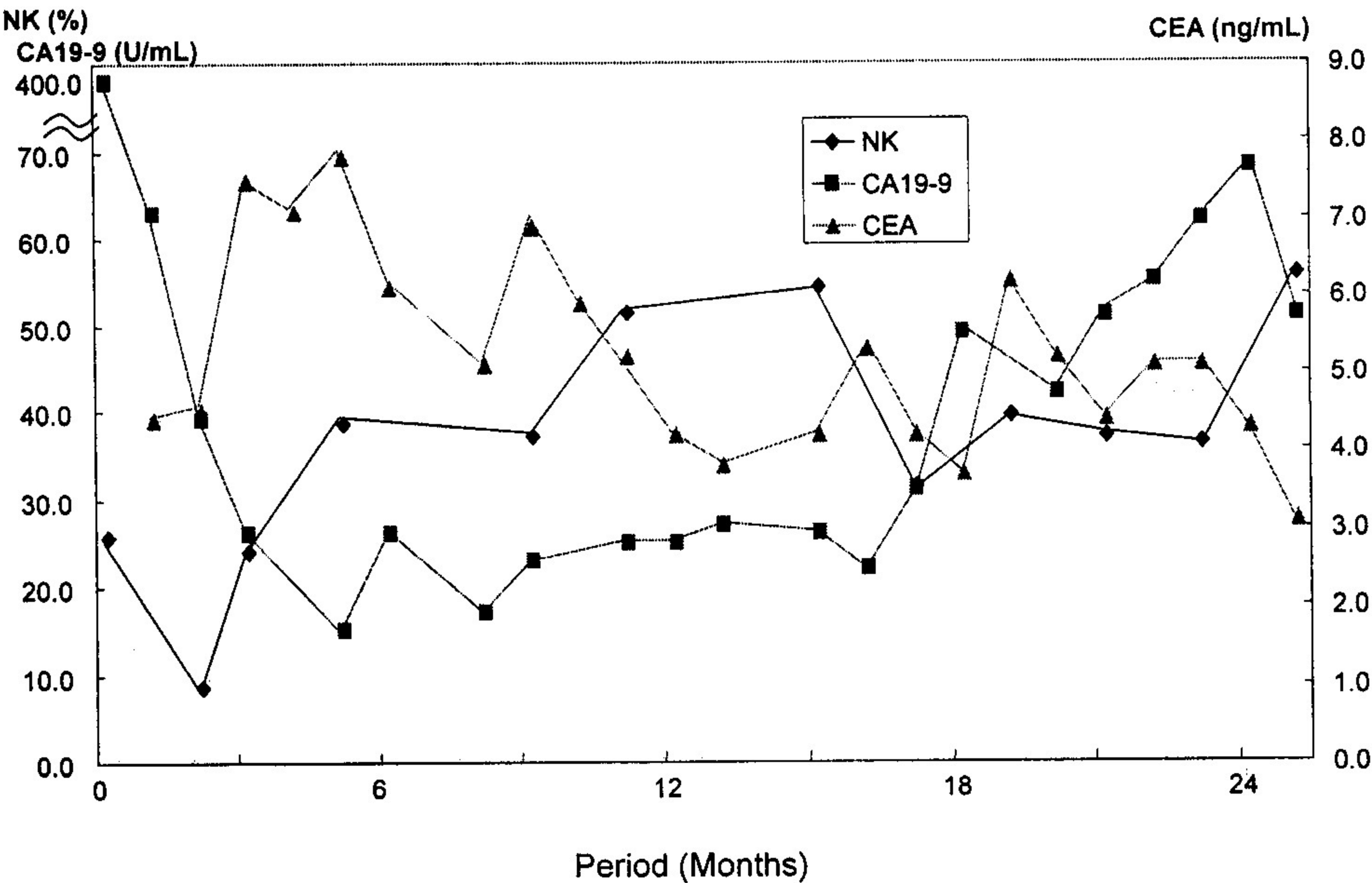


Fig. 3 Stomach Cancer (Stage IV) T.S. (60) F.

2) Patient: T.S., female, 60 years, stomach cancer (stage IV)

This patient underwent surgery for scirrhus carcinoma of the stomach, but curative resection was impossible because of cancerous peritonitis. She visited our clinic in August 2001 and complained of abdominal pain, an enlarged feeling in the abdomen, anemia, and anorexia (PS1). She was given the oral anticancer drug TS-1 and our hospital's therapy. BioBran/MGN-3 was taken at 3 g/day. The level of CA19-9 was 390 (U/mL) at the first visit and reduced to within the normal range by 3 months. The level of CEA increased, but began to fall at 6 months. Subjective symptoms gradually improved. Now, at 33 months (April 2004), her nutritional state is good, and she lives a normal life (PS0) (Fig. 3).

Discussion

During administration of BioBran/MGN-3, the patients' nutritional status was good, with no exacerbation in subjective and objective symptoms. Overall improvement was observed. The leukocyte count was low in many cases at the start of the study, but increased in almost all patients at the end of the study, and some achieved a normal value. Our clinic's complementary medicine maintains good physical conditions in high frequency after conventional cancer treatment. The conditions of patients in the present study were especially good, with no large difference in nutritional state between patients and healthy individuals. NK activity tended to increase: the number of patients with normal NK activity improved from 3 before the

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study to 11 after the study. These results supported data reported from other institutions<sup>8)</sup>. These phenomena were not clearly observed in patients who were not given BioBran/MGN-3.

Long-term administration of BioBran/MGN-3 had no adverse effects, such as compromised immunity, in cancer patients after conventional treatment, suggesting that BioBran/MGN-3 is useful as a dietary therapy that assists the improvement of the nutritional state.

### Conclusion

Long-term administration of BioBran/MGN-3 caused no subjective or objective adverse effects in cancer patients with decreased immunity. Improvement, rather than adverse changes, was observed in leukocyte counts and subsets. The NK cell activity was decreased at the baseline, but normalized after administration.

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