Evening Primrose Extract

For Anti-Diabetes & Anti-Obesity

- Evening Primrose Extract - P
  (Food, Powder)
- Evening Primrose Extract - WSPS
  (Food, Water-Soluble Powder)
- Evening Primrose Extract - PC
  (Cosmetics, Powder, No binder)
- Evening Primrose Extract - LC
  (Cosmetics, Liquid)
1. Introduction

Evening Primrose is a dicotyledonous plant of genus *Oenothera*. The seeds have been used for edible oil and also known to be a medicine in Europe. Evening Primrose oil contains much &-linolenic acid, which has well known to relieve obesity, diabetes mellitus, hypercholesterolemia, premenstrual syndrome (PMS), and so on.

Recently, polyphenols in plant seeds have been paid attention. Polyphenols prevent oxidation of lipids and scavenge active oxygen as a trigger of cause of various diseases. ORYZA OIL & FAT CHEMICAL CO., LTD. has been studying for the physiological function of polyphenols of the Evening Primrose seeds. We have recently developed an EVENING PRIMROSE EXTRACT (EPE), in which polyphenols are highly concentrated. The product contains more polyphenols than any other products, showing a strong antioxidative activity. Further, we found that EVENING PRIMROSE EXTRACT has an anti-diabetic activity.
2. What is “EVENING PRIMROSE EXTRACT”?

2-1. EVENING PRIMROSE

EVENING PRIMROSE EXTRACT has been cultivated in North America or China to obtain oils from the seeds. Evening primrose is introduced to Japan as decorative plants which is also seen on the riverside or seashore. The following 4 species are known to be evening primrose: *Oenothera laciniata*, *Oenothera striata*, *Oenothera biennis* and *Oenothera erythrosepaa*.

2-2. Contents of Total Polyphenols

The contents of total polyphenols in EVENING PRIMROSE EXTRACT were superior high comparison to other plant extracts.

Fig. 1 Contents of total polyphenols of EVENING PRIMROSE EXTRACT
2-3. Components of EVENING PRIMROSE EXTRACT
EVENING PRIMROSE EXTRACT contains polyphenols such as gallic acid, ellagic acid, pentagalloylglucose, catechin, etc. Our research has shown that EVENING PRIMROSE EXTRACT would be effective for prevention of diabetes and obesity by test of carbohydrase inhibition *in vitro* and by sugar tolerance *in vivo*.

![Chemical structures of polyphenols](image)

2-4. Antioxidant Activity

<table>
<thead>
<tr>
<th>Assayed Items</th>
<th>Result</th>
<th>Assaying Method</th>
</tr>
</thead>
<tbody>
<tr>
<td>Superoxide dismutase activity</td>
<td>$3.5 \times 10^3$ units/g</td>
<td>ESR method</td>
</tr>
</tbody>
</table>

Tested by: Japan Food Research Center Foundation
Research results issue number: 302010023-001
EVENING PRIMROSE EXTRACT has antioxidant activity stronger than other plant extracts. It is said that taking foods which have such a strong SOD-like activity or radical scavenging activity is good for prevention of several disease from habit of lifestyle.

Fig. 3 SOD-like activity of several plant extracts

Fig. 4 DPPH radical scavenging activity of several plant extracts
2-5. Mechanism of Diabetes Mellitus

Diabetes mellitus is caused by the lack of insulin secretion and manifests chronic hyperglycemia. The disease is ascribed to the genetic background and environmental factor (a habit of life). The non-insulin dependent diabetes mellitus (NIDDM), which account for more than 90% of the incidence of diabetes mellitus, is caused by over-eating, ingestion of high-fat diet, lack of exercise, obesity, or aging. These factors decrease in responsiveness and sensitiveness of insulin. In particular, obesity is one of the highest risk factors of the diabetes mellitus among them. Therefore, it is important to control diet or to take exercise for prevention of the incidence of diabetes mellitus.

![Diagram of digestion and absorption of sugar and diabetes](image)

**Fig. 5** Relation between digestion and absorption of sugar and diabetes
3. Function of “EVENING PRIMROSE EXTRACT”

3-1. Inhibitory Effect on Sugar Degradation Enzyme

EVENING PRIMROSE EXTRACT has inhibitory efficacy on $\alpha$-amylase and $\alpha$-glucosidase. Especially, the inhibitory activity to $\alpha$-amylase showed stronger than other plant extracts.

Fig. 6 Inhibitory activity of EVENING PRIMROSE EXTRACT and several plant extracts on $\alpha$-amylase

Fig. 7 Inhibitory activity of EVENING PRIMROSE EXTRACT and several plant extracts on $\alpha$-glucosidase
The inhibitory activity of polyphenols contained in EVENING PRIMROSE EXTRACT on sugar degradation enzyme were investigated. Pentagalloylglucose showed the highest inhibitory contribution in monomeric polyphenols.

Table 1. Inhibitory activity of evening primrose polyphenols on sugar degradation enzyme

<table>
<thead>
<tr>
<th>Polyphenol</th>
<th>Inhibitory Activity</th>
<th>Sugar Degradation Enzyme</th>
<th>Other Enzymes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pentagalloylglucose</td>
<td>High</td>
<td>High</td>
<td>Low</td>
</tr>
<tr>
<td>Other polyphenol</td>
<td>Low</td>
<td>Low</td>
<td>High</td>
</tr>
<tr>
<td>Control</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
3-2. Suppressive Effect on Blood Plasma Glucose Level (in vivo)

EVENING PRIMROSE EXTRACT significantly suppressed the elevation of blood glucose level by sugar tolerance test in rats.

Fig. 8 Sugar tolerance test (starch)

Fig. 9 Sugar tolerance test (sucrose)
3-3. Effect of Anti-Diabetic Activity (in vivo)

Hereditary diabetic mice (Type 加盟店, KK-Ay/Ta Jcl) were fed a test diet containing 1% of EVENING PRIMROSE EXTRACT for 6 weeks. As shown in Fig. 7, the elevation of blood plasma glucose level of the test group were almost entirely suppressed compared with control group.

![Fig. 10 Effect of feeding on diabetes in KK-Ay mice](image)

3-4. Suppressive Effect on Blood Glucose Level in Normal Subjects

The post-prandial increases in blood glucose levels were significantly suppressed by EVENING PRIMROSE EXTRACT.

![Fig. 11 Sugar tolerance test in normal human subjects](image)
3-5. Suppressive Effect on Blood Glucose Level in Diabetic Subjects

We examined effects of single intake of an extract of seeds of EVENING PRIMROSE EXTRACT on postprandial blood glucose levels on 18 subjects suffering mild diabetes and borderline diabetes. In all subjects, the rises of postprandial blood glucose levels were reduced by EVENING PRIMROSE EXTRACT ingestion, compared to those when placebo taken. These results suggest that EVENING PRIMROSE EXTRACT may be a useful food ingredient for prevention of diabetes mellitus.

![Graph showing blood glucose levels](image)

*Fig. 12 Sugar tolerance test in diabetic patients*

<table>
<thead>
<tr>
<th>Table 2 Postprandial insulin levels in diabetic patients</th>
</tr>
</thead>
<tbody>
<tr>
<td>Time (h)</td>
</tr>
<tr>
<td>----------</td>
</tr>
<tr>
<td>0</td>
</tr>
<tr>
<td>1</td>
</tr>
<tr>
<td>2</td>
</tr>
</tbody>
</table>

*Significance levels: *p < 0.05, **p < 0.01*
3-6. Induction Effects of Apotosis in Human Stomach Cancer Cells by EVENING PRIMROSE EXTRACT

For the stomach cancer cells any a change was not found in controls. DNA fragmentation effect in cells was found by EVENING PRIMROSE EXTRACT addition.

Fig. 13 Morphological changes of human stomach cancer cells
(A) Non-treated cells (B) Cells treated with 2mg/ml EVENING PRIMROSE EXTRACT

Fig. 14 DNA fragmentation effect in human stomach cancer cells
Dose-dependency (A) and time-dependency (B) of induction of apoptosis by EVENING PRIMROSE EXTRACT in human stomach cancer cells.

(A) M: DNA marker
(1) EVENING PRIMROSE (0mg/ml)
(2) EVENING PRIMROSE (1mg/ml)
(3) EVENING PRIMROSE (2mg/ml)

(B) M: DNA marker
(1) EVENING PRIMROSE (0mg/ml)
(2) EVENING PRIMROSE (2mg/ml) 1days
(3) EVENING PRIMROSE (2mg/ml) 2days
(4) EVENING PRIMROSE (2mg/ml) 3days
3-7. Inhibition of Tyrosinase

EVENING PRIMROSE EXTRACT inhibits tyrosinase activity and appears to be applicable to foods for whitening.

![Fig. 15. Tyrosinase Inhibitory Activity](image)

3-8. Moisture Retention Test

After direct topical application of samples to human skin, moisture retention ability was measured. When only distilled water was applied, the water content returned to the pre-application level after about 5 minutes. However, moisture was retained for more than 20 minutes after application of EVENING PRIMROSE EXTRACT.

![Fig. 16 Moisture retention test of EVENING PRIMROSE EXTRACT](image)
4. Stability of EVENING PRIMROSE EXTRACT

4-1. Thermal Resistance

The pyrolysis of EVENING PRIMROSE EXTRACT does not occur at a normal food processing temperature for 60 minutes.

Fig. 17 Heat-resistance of EVENING PRIMROSE EXTRACT

4-2. pH Stability

Polyphenols in EVENING PRIMROSE EXTRACT remains stable specially at neutral to acid field of pH.

Fig. 18 Influence of pH on the polyphenols contents
5. Daily Dosage of EVENING PRIMROSE EXTRACT

It is recommended to take more than 50mg/time or 100~500mg of total EVENING PRIMROSE EXTRACT-P per day.

6. Acute Toxicity and Safety

6-1. Residual Agricultural Chemicals

<table>
<thead>
<tr>
<th>Assayed Items</th>
<th>Results</th>
<th>Detection Limits</th>
<th>Assay Method</th>
</tr>
</thead>
<tbody>
<tr>
<td>BHC</td>
<td>Not Detected</td>
<td>0.02ppm</td>
<td>Gas Chromatography</td>
</tr>
<tr>
<td>DDT</td>
<td>Not Detected</td>
<td>0.02ppm</td>
<td>Gas Chromatography</td>
</tr>
<tr>
<td>Aldrin</td>
<td>Not Detected</td>
<td>0.01ppm</td>
<td>Gas Chromatography</td>
</tr>
<tr>
<td>Dieldrin</td>
<td>Not Detected</td>
<td>0.01ppm</td>
<td>Gas Chromatography</td>
</tr>
<tr>
<td>Endrin</td>
<td>Not Detected</td>
<td>0.01ppm</td>
<td>Gas Chromatography</td>
</tr>
<tr>
<td>Parathion</td>
<td>Not Detected</td>
<td>0.05ppm</td>
<td>Gas Chromatography</td>
</tr>
<tr>
<td>Malathon</td>
<td>Not Detected</td>
<td>0.05ppm</td>
<td>Gas Chromatography</td>
</tr>
</tbody>
</table>

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6-2. Acute Toxicity

5 weeks old rats had been bred for two weeks after administering 5000mg/kg. No toxic effects were observed, thus the LD$_{50}$ (rat) is more than 5000mg/kg.

6-3. Chromosome Aberration Test

The chromosome aberration test was examined using cultured mammalian cells (CHL/IU).
EVENING PRIMROSE EXTRACT was considered to be non-induce for chromosome aberration.

6-4. Reverse Mutation Assay  □ AMES TEST □

Salmonella typhimurium strains and Escherichia coli strain were treated with suspensions of the material using the Ames plate incorporation method at five dose levels both with and without the addition of a rat liver homogenate metabolising system. No significant increases in the frequency of revertant colonies were recorded for any of the bacterial strains, with any dose of the test material (EVENING PRIMROSE EXTRACT), either with or without metabolic activation.
EVENING PRIMROSE EXTRACT was considered to be non-mutagenic under the conditions of this test.
6-5. Micronucleus Test in Mice

The micronucleus study was conducted using the intraperitoneal route in groups of seven mice (males) at the maximum tolerated dose (MTD) 200 mg/kg with 100 and 50 mg/kg as the two lower dose levels. Animals were killed 24 or 48 hours later, the bone marrow extracted and smear preparations made and stained. Polychromatic (PCE) and normochromatic (NCE) erythrocytes were scored for the presence of micronuclei. There was no evidence of a significant increase in the incidence of micronucleated polychromatic erythrocytes in animals dosed with the test material (EVENING PRIMROSE EXTRACT) when compared to the concurrent vehicle control groups. EVENING PRIMROSE EXTRACT was considered to be non-genotoxic under the conditions of the test.

7. Practical Applications of EVENING PRIMROSE EXTRACT

EVENING PRIMROSE EXTRACT has 3 forms:
- EVENING PRIMROSE EXTRACT-P (powder type)
- EVENING PRIMROSE EXTRACT-WSPS (water soluble type powder)
- EVENING PRIMROSE EXTRACT-LC (liquid type for cosmetics)

<table>
<thead>
<tr>
<th>Applications</th>
<th>Examples</th>
</tr>
</thead>
<tbody>
<tr>
<td>Drinks</td>
<td>Tea, Blended tea, Protein shakes, and Nutritional drinks.</td>
</tr>
<tr>
<td>Dried Foods</td>
<td>Soup, Dried noodles, Seasoning, Pasta, Cereal, Oatmeal, and Topping for pizza.</td>
</tr>
<tr>
<td>Confectionery</td>
<td>Candies, Gum, Cookies, Pudding, Jelly, Yogurt, Chocolate</td>
</tr>
<tr>
<td>Snacks</td>
<td>Rice crackers, Cookies, and Wafers.</td>
</tr>
<tr>
<td>Fermentative Foods</td>
<td>Bread and Yogurt</td>
</tr>
<tr>
<td>Others</td>
<td>Health foods, Nutraceutical foods, and Functional foods</td>
</tr>
<tr>
<td>Cosmetics</td>
<td>Base cosmetics (Lotion, Milk, Cream, and so on)</td>
</tr>
<tr>
<td></td>
<td>Body cosmetics (Body lotion, Body cream, and so on)</td>
</tr>
<tr>
<td></td>
<td>Cleansing cosmetics (Soap, and so on)</td>
</tr>
<tr>
<td></td>
<td>Makeup cosmetics (Lipstick, Foundation, and so on), etc.</td>
</tr>
</tbody>
</table>
8. Packaging

- EVENING PRIMROSE EXTRACT-P
  5kg  Interior packaging: aluminium-coated plastic bag
       Exterior packaging: 18L thin and cardboard box

- EVENING PRIMROSE EXTRACT-WSPS
  4kg  Interior packaging: aluminium-coated plastic bag
       Exterior packaging: 18L thin and cardboard box

- EVENING PRIMROSE EXTRACT-PC
  4kg  Interior packaging: cubic polyethylene container
       Exterior packaging: cardboard box

- EVENING PRIMROSE EXTRACT-LC
  5kg  Interior packaging: cubic polyethylene container
       Exterior packaging: cardboard box

9. Storing Method

Store in cool, dry place. Avoid humidity.

10. Indication of Evening Primrose Extract

- EVENING PRIMROSE EXTRACT-P
  Evening Primrose Seed Extract

- EVENING PRIMROSE EXTRACT-WSPS
  Evening Primrose Seed Extract

- EVENING PRIMROSE EXTRACT-PC
  INCI name: Oenothera Biennis Seed Extract

- EVENING PRIMROSE EXTRACT-LC
  INCI name: Butylene Glycol (and) Water (and) Oenothera Biennis Seed Extract

Please consult your nation’s regulations.
Methods of Experiments

Fig. 1 Contents of total polyphenols
Samples dissolved in methanol were assayed by using Folin-Denis Method.

Fig. 3 SOD-like activity of several plant extracts
Samples dissolved in ethanol were assayed by using SOD-Test Wako.

Fig. 4 DPPH radical scavenging activity of several plant extracts
DPPH radical scavenging activity was assayed by using DPPH method.

Fig. 6 Inhibitory activity of EVENING PRIMROSE EXTRACT and several plant extracts on $\alpha$-amylase
$\alpha$-amylase from human saliva was purchased from Wako Pure Chemical Ind., Ltd., Japan. $\alpha$-Amylase activity was assayed by using Amylase-Test Wako. In advance, sample were diluted with ethanol and 10 $\mu$l of the sample solution was added to the assay system. The inhibitory activity (%) was calculated by using the following equation: 100 - (% of Control).

Fig. 7 Inhibitory activity of EVENING PRIMROSE EXTRACT and several plant extracts on $\alpha$-glucosidase
Acetone powder of rat intestine was homogenized in phosphate buffer and centrifuged. The supernatant was used as a crude enzyme solution containing $\alpha$-glucosidase. $\alpha$-Glucosidase activity was determined by the fluorescent assay with 4-methylumbelliferyl-$\alpha$-D-glucopyranoside as substrate. The inhibitory activity (%) was calculated as described above.

Fig. 8 Sugar tolerance test (Starch)
Fig. 9 Sugar tolerance test (Sucrose)
After 24h without foods, six weeks old rats were loaded 2g/kg of soluble starch (or sucrose) and 0〜2500mg/kg of EVENING PRIMROSE EXTRACT. Every 30 minutes, blood samples were bled from the tail vein into heparinized capillary tubes, and the plasma glucose level were measured by Glucose B-Test Wako.

Fig. 10 Effect of feeding on diabetes in KK-Ay mice
Five weeks old male mice (Type $\alpha$ diabetes, KK-Ay/Ta Jcl, Clea Japan, Inc., Osaka) were fed freely giving water and basal diets (CE-2) for one week to accustom them to the surroundings. The mice were divided into two groups. One group (control group) was fed on basal diets, another group (experimental group) was fed on basal diets containing 1% of EVENING PRIMROSE EXTRACT for 8 weeks. The plasma glucose levels were assayed every week.
Fig. 11 Sugar tolerance test in normal human subjects
The blood glucose levels of the subjects after fasting for 11 hours were in the range of 70-110mg/dl. Subjects took meal (200g boiled rice) with or without EVENING PRIMROSE EXTRACT-P (200mg) at 9:00 am. Blood was withdrawn from finger at 0, 30, 60, and 90 min and measured blood glucose using a portable analyzer equipped glucose sensor.

Fig. 12 Sugar Tolerance Tests in Type II Diabetic Patients
Eighteen adult type II diabetic patients whose fasting blood glucose levels were ranged 110 to 180 mg/dl were obliged to the tests. A crossover diet loading study using a placebo as a control drug was conducted. Subjects were fasted after 9:00 pm the day before this study. The subjects took meal (200g boiled rice) with placebo capsules or capsules containing EVENING PRIMROSE EXTRACT at 8:50 am. Blood was withdrawn from finger at 0, 30, 60, and 90 min and measured blood glucose and serum insulin levels.

Fig. 13 Morphological Changes of Human Gastric Cancer Cells
Human gastric cancer cells were grown in RPMI 1640 medium with 10% heat-inactivated fetal bovine serum at 37°C in 95% air-5% CO2 atmosphere of saturated humidity. After 3-day cultivation in the presence of EVENING PRIMROSE EXTRACT (2 mg/ml), the cellular morphology was examined by an epifluorescence microscope equipped with a CCD camera digital imaging system.

Fig. 14 DNA Fragmentation Effect in Human Gastric Cancer Cells
Human gastric cancer cells were cultured as described in the Fig. 13 legend. Exponentially growing cells were placed at the initial density of $5 \times 10^5$ cells/ml in culture flasks. After cultivation in the presence of vehicle, EVENING PRIMROSE EXTRACT for 1, 2, or 3 days, the cells were collected and rinsed. DNA was extracted from the cells as described previously. Equivalent amounts of DNA were put into the well of 2% agarose gel.

Fig. 15 Tyrosinase Inhibitory Activity
EVENING PRIMROSE EXTRACT was dissolved in distilled water to make solutions of a series of concentrations. After addition of the extract to mushroom-derived tyrosinase solution, the reaction from L-tyrosine to dopaquinone was measured in terms of the absorbance of dopaquinone.

Fig. 16 Moisture Retention Tests of EVENING PRIMROSE EXTREACT
EVENING PRIMROSE EXTRACT was dissolved in distilled water to make 1% solution. One drop of the solution was topically applied to the medial side of the left brachium. The drop was spread over a 2-cm square area and absorbed to the skin. One minute later, the solution on the surface was wiped out with soft paper. Another one minute later, the epidermal moisture content was measured using a corneometer CM825. The atmospheric condition was 26°C with relative humidity of 38%.
The product is extracted with aqueous ethanol from the seeds of evening primrose (*Oenothera biennis*). It contains more than 60.0 % of polyphenol.

**Appearance**
- Brown-red powder with slight unique smell

**Polyphenols**
- Min. 60.0 % (Folin-Denis method)

**Loss on Drying**
- Max. 5.0 % (1 g, 105 °C, 2 hours)

**Purity Tests**
1. **Heavy Metals**
   - Max. 10 ppm (The Japanese Standards for Food Additives)
2. **Arsenic**
   - Max. 1 ppm (Standard Methods of Analysis in Food Safety Regulation)

**Standard Plate Counts**
- Max. $1 \times 10^3$ cfu/g (Analysis for Hygienic Chemists)

**Moulds and Yeasts**
- Max. $1 \times 10^2$ cfu/g (Analysis for Hygienic Chemists)

**Coliforms**
- Negative (Analysis for Hygienic Chemists)

**Composition**
- Ingredient: Evening Primrose Seed Extract
- Content: 100%
EVENING PRIMROSE EXTRACT Ver. 4.1TK

PRODUCT STANDARD

PRODUCT NAME

EVENING PRIMROSE EXTRACT-WSPS

(FOOD)

The product is extracted with water from the seeds of evening primrose (*Oenothera biennis*). It contains more than 50.0 % of polyphenol. This product is water soluble.

### Appearance
Brown-red powder with slight unique smell.

### Content of Polyphenols
Min. 50.0 % (Folin-Denis Method)

### Loss on Drying
Max. 5.0 % (1 g, 105 °C, 2 hours)

### Purity Tests

1. **Heavy Metals**
   Max. 10 ppm (The Japanese Standards for Food Additives)

2. **Arsenic**
   Max. 1 ppm (Standard Methods of Analysis in Food Safety Regulation)

### Standard Plate Counts
Max. $1 \times 10^3$ cfu/g (Analysis for Hygienic Chemists)

### Moulds and Yeasts
Max. $1 \times 10^2$ cfu/g (Analysis for Hygienic Chemists)

### Coliforms
Negative (Analysis for Hygienic Chemists)

### Composition

<table>
<thead>
<tr>
<th>Ingredient</th>
<th>Content</th>
</tr>
</thead>
<tbody>
<tr>
<td>Evening Primrose Seed Extract</td>
<td>100%</td>
</tr>
</tbody>
</table>
PRODUCT STANDARD

PRODUCT NAME

EVENING PRIMROSE EXTRACT-PC
(COSMETIC)

The product is extracted with aqueous ethanol from the seeds of evening primrose (Oenothera biennis). It contains more than 60.0% of polyphenols.

Appearance
Brown-red powder with slight unique smell

Polyphenols
Min. 60.0 % (Folin-Denis Method)

Loss on Drying
Max. 5.0 % (1 g, 105 °C, 2 hours)

Purity Tests
(1) Heavy Metals
Max. 10 ppm (The Second Method)
(2) Arsenic
Max. 1 ppm (The Third Method)

Standard Plate Counts
Max. 1 × 10^2 cfu/g (Analysis for Hygienic Chemists)

Moulds and Yeasts
Max. 1 × 10^3 cfu/g (Analysis for Hygienic Chemists)

Coliforms
Negative (Analysis for Hygienic Chemists)

Composition

<table>
<thead>
<tr>
<th>Ingredient</th>
<th>Content</th>
</tr>
</thead>
<tbody>
<tr>
<td>Oenothera Biennis Seed Extract</td>
<td>100%</td>
</tr>
</tbody>
</table>

PRODUCT STANDARD

PRODUCT NAME

EVENING PRIMROSE EXTRACT-LC

(COSMETIC)

This product is extracted with aqueous 1,3-butyleneglycol from the seeds of evening primrose (*Oenothera biennis*).

**Appearance**

Brown-red liquid with slight unique smell

**Certification Test**

1. The solution (1 ‰ 5) of this product turns dark blue to black when ferric chloride is added. (Tannins)

2. The solution (1 ‰ 5) of this product turns the color off when potassium permanganate is added. (Polyphenols)

**Polyphenols**

Min. 0.5 % (Folin-Denis Method)

**pH**

4.0 ~ 5.5 (10 % solution)

**Specific Gravity**

1.020~1.060 (25 ℃)

**Purity Tests**

1. Heavy Metals Max. 10 ppm (The Second Method)

2. Arsenic Max. 1 ppm (The Third Method)

**Standard Plate Counts**

Max. 1 ⨉ 10³ cfu/g (Analysis for Hygienic Chemists)

**Moulds and Yeasts**

Max. 1 ⨉ 10² cfu/g (Analysis for Hygienic Chemists)

**Coliforms**

Negative (Analysis for Hygienic Chemists)

**Composition**

<table>
<thead>
<tr>
<th>Ingredients</th>
<th>Contents</th>
</tr>
</thead>
<tbody>
<tr>
<td>Butylene Glycol</td>
<td>69 %</td>
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<tr>
<td>Water</td>
<td>30 %</td>
</tr>
<tr>
<td><em>Oenothera Biennis</em> Seed Extract</td>
<td>1 %</td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td>100 %</td>
</tr>
</tbody>
</table>

Ref: *The Japanese Standards of Quasi-Drug Ingredients.*
ORYZA OIL & FAT CHEMICAL CO., LTD., striving for the development of the new functional food materials to promote your health.

- From product planning to OEM - For any additional information or assistance, please contact us:

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