

## SUMMARY OF PRODUCT CHARACTERISTICS

### 1. NAME OF THE MEDICINAL PRODUCT

Temodal 5 mg, 20 mg, 100 mg, 140 mg, 180 mg, or 250 mg hard capsules

### 2. QUALITATIVE AND QUANTITATIVE COMPOSITION

Each hard capsule contains 5 mg, 20 mg, 100 mg, 140 mg, 180 mg or 250 mg temozolomide.

#### Excipient with known effect:

Each 5 mg hard capsule contains 132.8 mg of anhydrous lactose. Each 20 mg hard capsule contains 182.2 mg of anhydrous lactose. Each 100 mg hard capsule contains 175.7 mg of anhydrous lactose. Each 140 mg hard capsule contains 246 mg of anhydrous lactose. Each 180 mg hard capsule contains 316.3 mg of anhydrous lactose. Each 250 mg hard capsule contains 154.3 mg of anhydrous lactose. For the full list of excipients, see section 6.1.

### 3. PHARMACEUTICAL FORM

Hard capsule (capsule).

The 5 mg hard capsules have an opaque white body, an opaque green cap, and are imprinted with black ink. The cap is imprinted with "Temodal". The body is imprinted with "5 mg", the Schering-Plough logo and two stripes.

The 20 mg hard capsules have an opaque white body, an opaque yellow cap, and are imprinted with black ink. The cap is imprinted with "Temodal". The body is imprinted with "20 mg", the Schering-Plough logo and two stripes.

The 100 mg hard capsules have an opaque white body, an opaque pink cap, and are imprinted with black ink. The cap is imprinted with "Temodal". The body is imprinted with "100 mg", the Schering-Plough logo and two stripes. The 140 mg hard capsules have an opaque white body, a blue cap, and are imprinted with black ink. The cap is imprinted with "Temodal". The body is imprinted with "140 mg", the Schering-Plough logo and two stripes.

The 180 mg hard capsules have an opaque white body, an opaque orange cap, and are imprinted with black ink. The cap is imprinted with "Temodal". The body is imprinted with "180 mg", the Schering-Plough logo and two stripes.

The 250 mg hard capsules have an opaque white body and cap, and are imprinted with black ink. The cap is imprinted with "Temodal". The body is imprinted with "250 mg", the Schering-Plough logo and two stripes.

### 4. CLINICAL PARTICULARS

#### 4.1 Therapeutic indications

Temodal is indicated for the treatment of:

- adult patients with newly-diagnosed glioblastoma multiforme concomitantly with radiotherapy (RT) and subsequently as monotherapy treatment.
- children from the age of three years, adolescents and adult patients with malignant glioma, such as glioblastoma multiforme or anaplastic astrocytoma, showing recurrence or progression after standard therapy.

#### 4.2 Posology and method of administration

Temodal should only be prescribed by physicians experienced in the oncological treatment of brain tumours.

Anti-emetic therapy may be administered (see section 4.4).

#### Posology

*Adult patients with newly-diagnosed glioblastoma multiforme*

Temodal is administered in combination with focal radiotherapy (concomitant phase) followed by up to 6 cycles of temozolomide (TMZ) monotherapy (monotherapy phase).

#### Concomitant phase

TMZ is administered orally at a dose of 75 mg/m<sup>2</sup> daily for 42 days concomitant with focal radiotherapy (60 Gy administered in 30 fractions). No dose reductions are recommended, but delay or discontinuation of TMZ administration should be decided weekly according to haematological and non-haematological toxicity criteria. TMZ administration can be continued throughout the 42 day concomitant period (up to 49 days) if all of the following conditions are met:

- absolute neutrophil count (ANC)  $\geq 1.5 \times 10^9/l$
- thrombocyte count  $\geq 100 \times 10^9/l$
- common toxicity criteria (CTC) non-haematological toxicity  $\leq$  Grade 1 (except for alopecia, nausea and vomiting).

During treatment a complete blood count should be obtained weekly. TMZ administration should be temporarily interrupted or permanently discontinued during the concomitant phase according to the haematological and non-haematological toxicity criteria as noted in Table 1.

| Toxicity  | TMZ interruption <sup>a</sup>        | TMZ discontinuation   |
|---|--------------------------------------|-----------------------|
| Absolute neutrophil count   | $\geq 0.5$ and $< 1.5 \times 10^9/l$ | $< 0.5 \times 10^9/l$ |
| Thrombocyte count   | $\geq 10$ and $< 100 \times 10^9/l$  | $< 10 \times 10^9/l$  |
| CTC non-haematological toxicity (except for alopecia, nausea, vomiting) | CTC Grade 2                          | CTC Grade 3 or 4      |

a: Treatment with concomitant TMZ can be continued when all of the following conditions are met: absolute neutrophil count  $\geq 1.5 \times 10^9/l$ ; thrombocyte count  $\geq 100 \times 10^9/l$ ; CTC non-haematological toxicity  $\leq$  Grade 1 (except for alopecia, nausea, vomiting).

#### Monotherapy phase

Four weeks after completing the TMZ + RT concomitant phase, TMZ is administered for up to 6 cycles of monotherapy treatment. Dose in Cycle 1 (monotherapy) is 150 mg/m<sup>2</sup> once daily for 5 days followed by 23 days without treatment. At the start of Cycle 2, the dose is escalated to 200 mg/m<sup>2</sup> if the CTC non-haematological toxicity for Cycle 1 is Grade  $\leq 2$  (except for alopecia, nausea and vomiting), absolute neutrophil count (ANC) is  $\geq 1.5 \times 10^9/l$ , and the thrombocyte count is  $\geq 100 \times 10^9/l$ . If the dose was not escalated at Cycle 2, escalation should not be done in subsequent cycles. Once escalated, the dose remains at 200 mg/m<sup>2</sup> per day for the first 5 days of each subsequent cycle except if toxicity occurs. Dose reductions and discontinuations during the monotherapy phase should be applied according to Tables 2 and 3.

During treatment a complete blood count should be obtained on Day 22 (21 days after the first dose of TMZ). The dose should be reduced or administration discontinued according to Table 3.

| Dose level | TMZ dose (mg/m <sup>2</sup> /day) | Remarks                                       |
|------------|-----------------------------------|---|
| -1         | 100                               | Reduction for prior toxicity                  |
| 0          | 150                               | Dose during Cycle 1                           |
| 1          | 200                               | Dose during Cycles 2-6 in absence of toxicity |

| Toxicity  | Reduce TMZ by 1 dose level <sup>a</sup> | Discontinue TMZ          |
|---|---|--------------------------|
| Absolute neutrophil count   | $< 1.0 \times 10^9/l$                   | See footnote b           |
| Thrombocyte count   | $< 50 \times 10^9/l$                    | See footnote b           |
| CTC non-haematological Toxicity (except for alopecia, nausea, vomiting) | CTC Grade 3                             | CTC Grade 4 <sup>b</sup> |

- a: TMZ dose levels are listed in Table 2.  
 b: TMZ is to be discontinued if:
- dose level -1 (100 mg/m<sup>2</sup>) still results in unacceptable toxicity
  - the same Grade 3 non-haematological toxicity (except for alopecia, nausea, vomiting) recurs after dose reduction.

#### Adult and paediatric patients 3 years of age or older with recurrent or progressive malignant glioma:

A treatment cycle comprises 28 days. In patients previously untreated with chemotherapy, TMZ is administered orally at a dose of 200 mg/m<sup>2</sup> once daily for the first 5 days followed by a 23 day treatment interruption (total of 28 days). In patients previously treated with chemotherapy, the initial dose is 150 mg/m<sup>2</sup> once daily, to be increased in the second cycle to 200 mg/m<sup>2</sup> once daily, for 5 days if there is no haematological toxicity (see section 4.4)

#### Special populations

##### Paediatric population

In patients 3 years of age or older, TMZ is only to be used in recurrent or progressive malignant glioma. Experience in these children is very limited (see sections 4.4 and 5.1). The safety and efficacy of TMZ in children under the age of 3 years have not been established. No data are available.

##### Patients with hepatic or renal impairment

The pharmacokinetics of TMZ were comparable in patients with normal hepatic function and in those with mild or moderate hepatic impairment. No data are available on the administration of TMZ in patients with severe hepatic impairment (Child's Class C) or with renal impairment. Based on the pharmacokinetic properties of TMZ, it is unlikely that dose reductions are required in patients with severe hepatic impairment or any degree of renal impairment. However, caution should be exercised when TMZ is administered in these patients.

##### Elderly patients

Based on a population pharmacokinetic analysis in patients 19-78 years of age, clearance of TMZ is not affected by age. However, elderly patients (> 70 years of age) appear to be at increased risk of neutropenia and thrombocytopenia (see section 4.4).

##### Method of administration

Temodal hard capsules should be administered in the fasting state.

The capsules must be swallowed whole with a glass of water and must not be opened or chewed.

If vomiting occurs after the dose is administered, a second dose should not be administered that day.

### 4.3 Contraindications

Hypersensitivity to the active substance or to any of the excipients listed in section 6.1.

Hypersensitivity to dacarbazine (DTIC). Severe myelosuppression (see section 4.4).

### 4.4 Special warnings and precautions for use

#### Opportunistic infections and reactivation of infections

Opportunistic infections (such as *Pneumocystis jirovecii* pneumonia) and reactivation of infections (such as HBV, CMV) have been observed during the treatment with TMZ (see section 4.8).

#### Meningoencephalitis herpetic

In post marketing cases, meningoencephalitis herpetic (including fatal cases) has been observed in patients receiving TMZ in combination with radiotherapy, including cases of concomitant steroids administration.

#### *Pneumocystis jirovecii* pneumonia

Patients who received concomitant TMZ and RT in a pilot trial for the prolonged 42-day schedule were shown to be at particular risk for developing *Pneumocystis jirovecii* pneumonia (PCP). Thus, prophylaxis against PCP is required for all patients receiving concomitant TMZ and RT for the 42-day regimen (with a maximum of 49 days) regardless of lymphocyte count. If lymphopenia occurs, they are to continue the prophylaxis until recovery of lymphopenia to grade  $\leq 1$ .

There may be a higher occurrence of PCP when TMZ is administered during a longer dosing regimen. However, all patients receiving TMZ, particularly patients receiving steroids, should be observed closely for the development of PCP, regardless of the regimen. Cases of fatal respiratory failure have been reported in patients using TMZ, in particular in combination with dexamethasone or other steroids.

#### HBV

Hepatitis due to hepatitis B virus (HBV) reactivation, in some cases resulting in death, has been reported. Experts in liver disease should be consulted before treatment is initiated in patients with positive hepatitis B serology (including those with active disease). During treatment patients should be monitored and managed appropriately.

#### Hepatotoxicity

Hepatic injury, including fatal hepatic failure, has been reported in patients treated with TMZ (see section 4.8). Baseline liver function tests should be performed prior to treatment initiation. If abnormal, physicians should assess the benefit/risk prior to initiating temozolomide including the potential for fatal hepatic failure. For patients on a 42 day treatment cycle liver function tests should be repeated midway during this cycle. For all patients, liver function tests should be checked after each treatment cycle. For patients with significant liver function abnormalities, physicians should assess the benefit/risk of continuing treatment. Liver toxicity may occur several weeks or more after the last treatment with temozolomide.

#### Malignancies

Cases of myelodysplastic syndrome and secondary malignancies, including myeloid leukaemia, have also been reported very rarely (see section 4.8).

#### Anti-emetic therapy

Nausea and vomiting are very commonly associated with TMZ.

Anti-emetic therapy may be administered prior to or following administration of TMZ.

#### Adult patients with newly-diagnosed glioblastoma multiforme

Anti-emetic prophylaxis is recommended prior to the initial dose of concomitant phase and it is strongly recommended during the monotherapy phase.

#### Patients with recurrent or progressive malignant glioma

Patients who have experienced severe (Grade 3 or 4) vomiting in previous treatment cycles may require anti-emetic therapy.

#### Laboratory parameters

Patients treated with TMZ may experience myelosuppression, including prolonged pancytopenia, which may result in aplastic anaemia, which in some cases has resulted in a fatal outcome. In some cases, exposure to concomitant medicinal products associated with aplastic anaemia, including carbamazepine, phenytoin, and sulfamethoxazole/trimethoprim, complicates assessment. Prior to dosing, the following laboratory parameters must be met: ANC  $\geq 1.5 \times 10^9/l$  and platelet count  $\geq 100 \times 10^9/l$ . A complete blood count should be obtained on Day 22 (21 days after the first dose) or within 48 hours of that day, and weekly until ANC  $> 1.5 \times 10^9/l$  and platelet count  $> 100 \times 10^9/l$ . If ANC falls to  $< 1.0 \times 10^9/l$  or the platelet count is  $< 50 \times 10^9/l$  during any cycle, the next cycle should be reduced one dose level (see section 4.2). Dose levels include 100 mg/m<sup>2</sup>, 150 mg/m<sup>2</sup>, and 200 mg/m<sup>2</sup>. The lowest recommended dose is 100 mg/m<sup>2</sup>.

#### Paediatric population

There is no clinical experience with use of TMZ in children under the age of 3 years. Experience in older children and adolescents is very limited (see sections 4.2 and 5.1).

#### Elderly patients (> 70 years of age)

Elderly patients appear to be at increased risk of neutropenia and thrombocytopenia, compared with younger patients. Therefore, special care should be taken when TMZ is administered in elderly patients.

#### Male patients

Men being treated with TMZ should be advised not to father a child up to 6 months after receiving the last dose and to seek advice on cryoconservation of sperm prior to treatment (see section 4.6).

#### Lactose

This medicinal product contains lactose. Patients with rare hereditary problems of galactose intolerance, the Lapp lactase deficiency or glucose-galactose malabsorption should not take this medicine.

### 4.5 Interaction with other medicinal products and other forms of interaction

In a separate phase I study, administration of TMZ with ranitidine did not result in alterations in the extent of absorption of temozolomide or the exposure to its active metabolite monomethyl triazenoimidazole carboxamide (MTIC).

Administration of TMZ with food resulted in a 33 % decrease in C<sub>max</sub> and a 9 % decrease in area under the curve (AUC). As it cannot be excluded that the change in C<sub>max</sub> is clinically significant, Temodal should be administered without food.

Based on an analysis of population pharmacokinetics in phase II trials, co-administration of dexamethasone, prochlorperazine, phenytoin, carbamazepine, ondansetron, H<sub>2</sub> receptor antagonists, or phenobarbital did not alter the clearance of TMZ. Co-administration with valproic acid was associated with a small but statistically significant decrease in clearance of TMZ.

No studies have been conducted to determine the effect of TMZ on the metabolism or elimination of other medicinal products. However, since TMZ does not undergo hepatic metabolism and exhibits low protein binding, it is unlikely that it would affect the pharmacokinetics of other medicinal products (see section 5.2).

Use of TMZ in combination with other myelosuppressive agents may increase the likelihood of myelosuppression.

#### Paediatric population

Interaction studies have only been performed in adults.

### 4.6 Fertility, pregnancy and lactation

#### Pregnancy

There are no data in pregnant women. In preclinical studies in rats and rabbits receiving 150 mg/m<sup>2</sup> TMZ, teratogenicity and/or foetal toxicity were demonstrated (see section 5.3). Temodal should not be administered to pregnant women. If use during pregnancy must be considered, the patient should be apprised of the potential risk to the foetus.

#### Breast-feeding

It is not known whether TMZ is excreted in human milk; thus, breast-feeding should be discontinued while receiving treatment with TMZ.

#### Women of childbearing potential

Women of childbearing potential should be advised to use effective contraception to avoid pregnancy while they are receiving TMZ.

#### Male fertility

TMZ can have genotoxic effects. Therefore, men being treated with it should be advised not to father a child up to 6 months after receiving the last dose and to seek advice on cryoconservation of sperm prior to treatment, because of the possibility of irreversible infertility due to therapy with TMZ.

### 4.7 Effects on ability to drive and use machines

TMZ has minor influence on the ability to drive and use machines due to fatigue and somnolence (see section 4.8).

### 4.8 Undesirable effects

#### Clinical trial experience

In patients treated with TMZ, whether used in combination with RT or as monotherapy following RT for newly-diagnosed glioblastoma multiforme or as monotherapy in patients with recurrent or progressive glioma, the reported very common adverse reactions were similar: nausea, vomiting, constipation, anorexia, headache and fatigue. Convulsions were reported very commonly in the newly-diagnosed glioblastoma multiforme patients receiving monotherapy, and rash was reported very commonly in newly-diagnosed glioblastoma multiforme patients receiving TMZ concurrent with RT and also as monotherapy, and commonly in recurrent glioma. Most haematologic adverse reactions were reported commonly or very commonly in both indications (Tables 4 and 5); the frequency of grade 3-4 laboratory findings is presented after each table.

In the tables undesirable effects are classified according to System Organ Class and frequency. Frequency groupings are defined according to the following convention: Very common ( $\geq 1/100$ ); Common ( $\geq 1/1000$  to  $< 1/100$ ); Uncommon ( $\geq 1/1,000$  to  $< 1/100$ ); Rare ( $\geq 1/10,000$  to  $< 1/1,000$ ); Very rare ( $< 1/10,000$ ). Within each frequency grouping, undesirable effects are presented in order of decreasing seriousness.



### Newly-diagnosed glioblastoma multiforme

Table 4 provides treatment-emergent adverse events in patients with newly-diagnosed glioblastoma multiforme during the concomitant and monotherapy phases of treatment.

| System organ class  | TMZ + concomitant RT n=288*  | TMZ monotherapy n=224  |
|---|--|--|
| <b>Infections and infestations</b>                          |  |  |
| Common:   | Infection, <i>Herpes simplex</i> , wound infection, pharyngitis, candidiasis oral  | Infection, candidiasis oral  |
| Uncommon:   |  | <i>Herpes simplex</i> , herpes zoster, influenza-like symptoms   |
| <b>Blood and lymphatic system disorders</b>                 |  |  |
| Common:   | Neutropenia, thrombocytopenia, lymphopenia, leucopenia   | Febile neutropenia, thrombocytopenia, anaemia, leucopenia  |
| Uncommon:   | Febrile neutropenia, anaemia   | Lymphopenia, petechiae   |
| <b>Endocrine disorders</b>                                  |  |  |
| Uncommon:   | Cushingoid   | Cushingoid   |
| <b>Metabolism and nutrition disorders</b>                   |  |  |
| Very common:  | Anorexia   | Anorexia   |
| Common:   | Hyperglycaemia, weight decreased   | Weight decreased   |
| Uncommon:   | Hypokalaemia, alkaline phosphatase increased, weight increased   | Hyperglycaemia, weight increased   |
| <b>Psychiatric disorders</b>                                |  |  |
| Common:   | Anxiety, emotional lability, insomnia  | Anxiety, depression, emotional lability, insomnia  |
| Uncommon:   | Agitation, apathy, behaviour disorder, depression, hallucination   | Hallucination, amnesia   |
| <b>Nervous system disorders</b>                             |  |  |
| Very common:  | Headache   | Convulsions, headache  |
| Common:   | Convulsions, consciousness decreased, somnolence, aphasia, balance impaired, dizziness, confusion, memory impairment, concentration impaired, neuropathy, paresthesia, speech disorder, tremor | Hemiparesis, aphasia, balance impaired, somnolence, confusion, dizziness, memory impairment, concentration impaired, dysphasia, neurological disorder (NOS), neuropathy, peripheral neuropathy, paresthesia, speech disorder, tremor |
| Uncommon:   | Status epilepticus, extrapyramidal disorder, hemiparesis, ataxia, cognition impaired, dysphasia, gait abnormal, hypersthesia, hyposthesia, neurological disorder (NOS), peripheral neuropathy  | Hemiplegia, ataxia, coordination abnormal, gait abnormal, hypersthesia, sensory disturbance  |
| <b>Eye disorders</b>  |  |  |
| Common:   | Vision blurred   | Visual field defect, vision blurred, diplopia  |
| Uncommon:   | Hemianopia, visual acuity reduced, vision disorder, visual field defect, eye pain  | Visual acuity reduced, eye pain, eyes dry  |
| <b>Ear and labyrinth disorders</b>                          |  |  |
| Common:   | Hearing impairment   | Hearing impairment, tinnitus   |
| Uncommon:   | Otitis media, tinnitus, hyperacusis, Earache   | Deafness, vertigo, earache   |
| <b>Cardiac disorders</b>                                    |  |  |
| Uncommon:   | Palpitation  |  |
| <b>Vascular disorders</b>                                   |  |  |
| Common:   | Haemorrhage, oedema, oedema leg  | Haemorrhage, deep venous thrombosis, oedema leg  |
| Uncommon:   | Cerebral haemorrhage, Hypertension   | Embolism pulmonary, oedema, oedema peripheral  |
| <b>Respiratory, thoracic and mediastinal disorders</b>      |  |  |
| Common:   | Dyspnoea, coughing   | Dyspnoea, coughing   |
| Uncommon:   | Pneumonia, upper respiratory infection, nasal congestion   | Pneumonia, sinusitis, upper respiratory infection, bronchitis  |
| <b>Gastrointestinal disorders</b>                           |  |  |
| Very common:  | Constipation, nausea, vomiting   | Constipation, nausea, vomiting   |
| Common:   | Stomatitis, diarrhoea, abdominal pain, dyspepsia, dysphagia  | Stomatitis, diarrhoea, dyspepsia, dysphagia, mouth dry   |
| Uncommon:   |  | Abdominal distension, fecal incontinence, gastrointestinal disorder (NOS), gastroenteritis, haemorrhoids   |
| <b>Skin and subcutaneous tissue disorders</b>               |  |  |
| Very common:  | Rash, alopecia   | Rash, alopecia   |
| Common:   | Dermatitis, dry skin, erythema, Pruritus   | Dry skin, pruritus   |
| Uncommon:   | Skin exfoliation, photosensitivity reaction, pigmentation abnormal   | Erythema, pigmentation abnormal, sweating increased  |
| <b>Musculoskeletal and connective tissue disorders</b>      |  |  |
| Common:   | Muscle weakness, arthralgia  | Muscle weakness, arthralgia, musculoskeletal pain, myalgia   |
| Uncommon:   | Myopathy, back pain, musculoskeletal pain, myalgia   | Myopathy, back pain  |
| <b>Renal and urinary disorders</b>                          |  |  |
| Common:   | Micturition frequency, urinary incontinence  | Urinary incontinence   |
| Uncommon:   |  | Dysuria  |
| <b>Reproductive system and breast disorders</b>             |  |  |
| Uncommon:   | Impotence  | Vaginal haemorrhage, menorrhagia, amenorrhoea, vaginitis, breast pain  |
| <b>General disorders and administration site conditions</b> |  |  |
| Very common:  | Fatigue  | Fatigue  |
| Common:   | Allergic reaction, fever, radiation injury, face oedema, pain, taste perversion  | Allergic reaction, fever, radiation injury, pain, taste perversion   |
| Uncommon:   | Asthenia, flushing, hot flushes, condition aggravated, rigors, tongue discoloration, parosmia, thirst  | Asthenia, face oedema, pain, condition aggravated, rigors, tooth disorder  |
| <b>Investigations</b>                                       |  |  |
| Common:   | ALT increased  | ALT increased  |
| Uncommon:   | Hepatic enzymes increased, Gamma GT increased, AST increased   |  |

\*A patient who was randomised to the RT arm only, received TMZ + RT.

#### Laboratory results

Myelosuppression (neutropenia and thrombocytopenia), which is known dose-limiting toxicity for most cytotoxic agents, including TMZ, was observed. When laboratory abnormalities and adverse events were combined across concomitant and monotherapy treatment phases, Grade 3 or Grade 4 neutrophil abnormalities including neutropenic events were observed in 8 % of the patients. Grade 3 or Grade 4 thrombocyte abnormalities, including

thrombocytopenic events were observed in 14 % of the patients who received TMZ.

#### Recurrent or progressive malignant glioma

In clinical trials, the most frequently occurring treatment-related undesirable effects were gastrointestinal disorders, specifically nausea (43 %) and vomiting (36 %). These reactions were usually Grade 1 or 2 (0 – 5 episodes of vomiting in 24 hours) and were either self-limiting or readily controlled with standard anti-emetic therapy. The incidence of severe nausea and vomiting was 4 %.

Table 5 includes adverse reactions reported during clinical trials for recurrent or progressive malignant glioma and following the marketing of Temodal.

| Inflections and infestations                           |  |
|--|--|
| Rare:  | Opportunistic infections, including PCP                              |
| <b>Blood and lymphatic system disorders</b>            |  |
| Very common:   | Neutropenia or lymphopenia (grade 3-4), thrombocytopenia (grade 3-4) |
| Uncommon:  | Pancytopenia, anaemia (grade 3-4), leucopenia                        |
| <b>Metabolism and nutrition disorders</b>              |  |
| Very common:   | Anorexia   |
| Common:  | Weight decrease  |
| <b>Nervous system disorders</b>                        |  |
| Very common:   | Headache   |
| Common:  | Somnolence, dizziness, paresthesia                                   |
| <b>Respiratory, thoracic and mediastinal disorders</b> |  |
| Common:  | Dyspnoea   |



patients with newly-diagnosed glioblastoma multiforme demonstrated a statistically significant improvement in overall survival (OS) compared with RT alone (Figure 1).

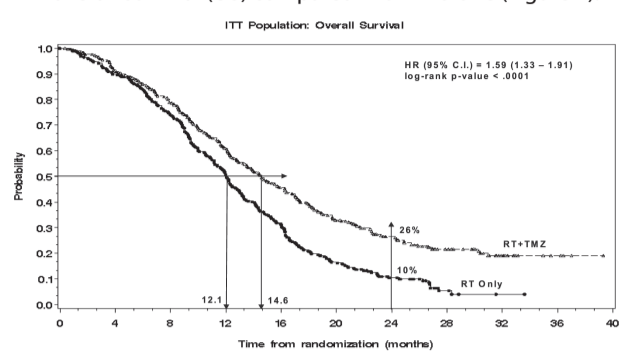


Figure 1 Kaplan-Meier curves for overall survival (intent-to-treat population)

The results from the trial were not consistent in the subgroup of patients with a poor performance status (WHO PS=2, n=70), where overall survival and time to progression were similar in both arms. However, no unacceptable risks appear to be present in this patient group.

#### Recurrent or progressive malignant glioma

Data on clinical efficacy in patients with glioblastoma multiforme (Karnofsky performance status [KPS]  $\geq$  70), progressive or recurrent after surgery and RT, were based on two clinical trials with oral TMZ. One was a non-comparative trial in 138 patients (29 % received prior chemotherapy), and the other was a randomised active-controlled trial of TMZ vs procarbazine in a total of 225 patients (67 % received prior treatment with nitrosourea based chemotherapy). In both trials, the primary endpoint was progression-free survival (PFS) defined by MRI scans or neurological worsening. In the non-comparative trial, the PFS at 6 months was 19 %, the median progression-free survival was 2.1 months, and the median overall survival 5.4 months. The objective response rate (ORR) based on MRI scans was 8 %.

In the randomised active-controlled trial, the PFS at 6 months was significantly greater for TMZ than for procarbazine (21 % vs 8 %, respectively – chi-square  $p = 0.008$ ) with median PFS of 2.89 and 1.88 months respectively (log rank  $p = 0.0063$ ). The median survival was 7.34 and 5.66 months for TMZ and procarbazine, respectively (log rank  $p = 0.33$ ). At 6 months, the fraction of surviving patients was significantly higher in the TMZ arm (60 %) compared with the procarbazine arm (44 %) (chi-square  $p = 0.019$ ). In patients with prior chemotherapy a benefit was indicated in those with a KPS  $\geq$  80.

Data on time to worsening of neurological status favoured TMZ over procarbazine as did data on time to worsening of performance status (decrease to a KPS of < 70 or a decrease by at least 30 points). The median times to progression in these endpoints ranged from 0.7 to 2.1 months longer for TMZ than for procarbazine (log rank  $p < 0.01$  to 0.03).

#### Recurrent anaplastic astrocytoma

In a multicentre, prospective phase II trial evaluating the safety and efficacy of oral TMZ in the treatment of patients with anaplastic astrocytoma at first relapse, the 6 month PFS was 46 %. The median PFS was 5.4 months. Median overall survival was 14.6 months. Response rate, based on the central reviewer assessment, was 35 % (13 CR and 43 PR) for the intent-to-treat population (ITT) n=162. In

43 patients stable disease was reported. The 6-month event-free survival for the ITT population was 44 % with a median event-free survival of 4.6 months, which was similar to the results for the progression-free survival. For the eligible histology population, the efficacy results were similar. Achieving a radiological objective response or maintaining progression-free status was strongly associated with maintained or improved quality of life.

#### Paediatric population

Oral TMZ has been studied in paediatric patients (age 3-18 years) with recurrent brainstem glioma or recurrent high grade astrocytoma, in a regimen administered daily for 5 days every 28 days. Tolerance to TMZ is similar to adults.

#### 5.2 Pharmacokinetic properties

TMZ is spontaneously hydrolyzed at physiologic pH primarily to the active species, 3-methyl-(triazene-1-yl)imidazole-4-carboxamide (MTIC). MTIC is spontaneously hydrolyzed to 5-amino-imidazole-4-carboxamide (AIC), a known intermediate in purine and nucleic acid biosynthesis, and to methylhydrazine, which is believed to be the active alkylating species. The cytotoxicity of MTIC is thought to be primarily due to alkylation of DNA mainly at the O' and N' positions of guanine. Relative to the AUC of TMZ, the exposure to MTIC and AIC is ~ 2.4 % and 23 %, respectively. *In vivo*, the  $t_{1/2}$  of MTIC was similar to that of TMZ, 1.8 hr.

#### Absorption

After oral administration to adult patients, TMZ is absorbed rapidly, with peak concentrations reached as early as 20 minutes post-administration (mean time between 0.5 and 1.5 hours). After oral administration of  $^{14}$ C-labelled TMZ, mean faecal excretion of  $^{14}$ C over 7 days post-dose was 0.8 % indicating complete absorption.

#### Distribution

TMZ demonstrates low protein binding (10 % to 20 %), and thus it is not expected to interact with highly protein-bound substances.

PET studies in humans and preclinical data suggest that TMZ crosses the blood- brain barrier rapidly and is present in the CSF. CSF penetration was confirmed in one patient; CSF exposure based on AUC of TMZ was approximately 30 % of that in plasma, which is consistent with animal data.

#### Elimination

The half-life ( $t_{1/2}$ ) in plasma is approximately 1.8 hours. The major route of  $^{14}$ C elimination is renal. Following oral administration, approximately 5 % to 10 % of the dose is recovered unchanged in the urine over 24 hours, and the remainder excreted as temozolomide acid, 5-aminoimidazole-4-carboxamide (AIC) or unidentified polar metabolites.

Plasma concentrations increase in a dose-related manner. Plasma clearance, volume of distribution and half-life are independent of dose.

#### Special populations

Analysis of population-based pharmacokinetics of TMZ revealed that plasma TMZ clearance was independent of age, renal function or tobacco use. In a separate pharmacokinetic study, plasma pharmacokinetic profiles in patients with mild to moderate hepatic impairment were similar to those observed in patients with normal hepatic function.

Paediatric patients had a higher AUC than adult patients; however, the maximum tolerated dose (MTD) was 1,000 mg/m<sup>2</sup> per cycle both in children and in adults.

#### 5.3 Preclinical safety data

Single-cycle (5-day dosing, 23 days non-treatment), 3- and 6-cycle toxicity studies were conducted in rats and dogs. The primary targets of toxicity included the bone marrow, lymphoreticular system, testes, the gastrointestinal tract and, at higher doses, which were lethal to 60 % to 100 % of rats and dogs tested, degeneration of the retina occurred. Most of the toxicity showed evidence of reversibility, except for adverse events on the male reproductive system and retinal degeneration. However, because the doses implicated in retinal degeneration were in the lethal dose range, and no comparable effect has been observed in clinical studies, this finding was not considered to have clinical relevance.

TMZ is an embryotoxic, teratogenic and genotoxic alkylating agent. TMZ is more toxic to the rat and dog than to humans, and the clinical dose approximates the minimum lethal dose in rats and dogs. Dose-related reductions in leukocytes and platelets appear to be sensitive indicators of toxicity. A variety of neoplasms, including mammary carcinomas, keratocanthoma of the skin and basal cell adenoma were observed in the 6-cycle rat study while no tumours or pre-neoplastic changes were evident in dog studies. Rats appear to be particularly sensitive to oncogenic effects of TMZ, with the occurrence of first tumours within 3 months of initiating dosing. This latency period is very short even for an alkylating agent.

Results of the Ames/salmonella and Human Peripheral Blood Lymphocyte (HPBL) chromosome aberration tests showed a positive mutagenicity response.

#### 6. PHARMACEUTICAL PARTICULARS

##### 6.1 List of excipients

**Capsule content:** anhydrous lactose, colloidal anhydrous silica, sodium starch glycolate type A, tartaric acid, stearic acid.

##### Capsule shell:

**5 mg:** gelatin, titanium dioxide (E 171), sodium laurilsulfate, yellow iron oxide (E 172), indigo carmine (E 132).

##### 20 mg:

gelatin, titanium dioxide (E 171), sodium lauril sulfate, yellow iron oxide (E 172)

##### 100 mg:

gelatin, titanium dioxide (E 171), sodium lauril sulfate, red iron oxide (E172).

##### 140 mg:

gelatin, titanium dioxide (E 171), sodium lauril sulfate, indigo carmine (E 132).

##### 180 mg:

gelatin, titanium dioxide (E 171), sodium lauril sulfate, yellow iron oxide (E 172), red iron oxide (E 172)

##### 250 mg:

gelatin, titanium dioxide (E 171), sodium lauril sulfate

##### Printing ink:

shellac, propylene glycol, purified water, ammonium hydroxide, potassium hydroxide, black iron oxide (E 172).

#### 6.2 Incompatibilities

Not applicable.

#### 6.3 Shelf life

Do not use after the expiry date shown on the packing.

#### 6.4 Special precautions for storage

##### Bottle presentation

Do not store above 30 °C. Store in the original bottle in order to protect from moisture. Keep the bottle tightly closed.

##### Sachet presentation

Do not store above 30 °C.

#### 6.5 Nature and contents of container

##### Bottle presentation

Type I amber glass bottles with polypropylene child-resistant closures containing 5 or 20 hard capsules. The carton contains one bottle.

##### Sachet presentation

Sachets are composed of linear low density polyethylene (innermost layer), aluminium and polyethylene terephthalate.

Each sachet contains 1 hard capsule and is dispensed in a cardboard carton. The carton contains 5 or 20 hard capsules, individually sealed in sachets.

Not all pack sizes or presentations may be marketed.

#### 6.6 Special precautions for disposal and other handling

Capsules should not be opened. If a capsule becomes damaged, contact of the powder contents with skin or mucous membrane must be avoided. If Temodal comes into contact with skin or mucosa, it should be washed immediately and thoroughly with soap and water.

Patients should be advised to keep capsules out of the sight and reach of children, preferably in a locked cupboard. Accidental ingestion can be lethal for children.

Any unused medicinal product or waste material should be disposed of in accordance with local requirements.

#### 7. MARKETING AUTHORISATION HOLDER

Merck Sharp & Dohme Limited Hertford Road, Hoddesdon Hertfordshire EN11 9BU United Kingdom

#### 8. MANUFACTURED BY

Orion Corporation Tengstrominkatu 8 Finland-20360 Turku

#### 9. PACKAGED AND RELEASED BY

Schering-Plough Labo N.V., Industriepark 30, B-2220 Heist-op-den-berg, Belgium

#### 10. DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION

Date of first authorisation: 26 January 1999 Date of latest renewal: 26 January 2009

#### 11. DATE OF REVISION OF THE TEXT

May 2018 Detailed information on this medicinal product is available on the website of the European Medicines Agency <http://www.ema.europa.eu>.

#### (THIS IS A MEDICAMENT)

Medicament is a product which affects your health, and its consumption contrary to instructions is dangerous for you. Follow strictly the doctor's prescription, the method of use, and the instructions of the pharmacist who sold the medicament.

The doctor and the pharmacist are experts in medicine, its benefits and risks.

Do not by yourself interrupt the period of treatment prescribed for you.

Do not repeat the same prescription without consulting your doctor.

Keep medicament out of reach of children

Council of Arab Health Ministers Union of Arab Pharmacists

