PRODUCT MONOGRAPH
INCLUDING PATIENT MEDICATION INFORMATION

Pr**SPRYCEL®**

dasatinib (as monohydrate)

Tablet, 20 mg, 50 mg, 70 mg, 80 mg, 100 mg and 140 mg, oral

Protein kinase inhibitor

Bristol-Myers Squibb Canada

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RECENT MAJOR LABEL CHANGES

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PART I: HEALTH PROFESSIONAL INFORMATION

1 INDICATIONS

SPRYCEL (dasatinib) is indicated for the treatment of adults with:

- Newly diagnosed Philadelphia chromosome positive (Ph+) chronic myeloid leukemia (CML) in chronic phase.

  Clinical effectiveness of SPRYCEL treatment in patients with newly diagnosed Ph+ CML in chronic phase is based on confirmed complete cytogenetic response rate (cCCyR) within 12 months. As of the 60 month cut-off date, overall survival, prevention of progression to advanced stage CML, or time-in cCCyR benefits have not been demonstrated (see 14 CLINICAL TRIALS).

- Ph+ chronic, accelerated, or blast phase chronic myeloid leukemia (CML) with resistance or intolerance to prior therapy including imatinib mesylate.

  Clinical effectiveness of SPRYCEL in CML is based on the rates of hematologic and cytogenetic responses in clinical trials with a minimum of 24 months of follow-up (see 14 CLINICAL TRIALS).

- Ph+ acute lymphoblastic leukemia (ALL) with resistance or intolerance to prior therapy.

  Clinical effectiveness in Ph+ ALL is based on the rates of hematologic and cytogenetic responses in clinical trials with a minimum of 24 months of follow-up (see 14 CLINICAL TRIALS).

SPRYCEL (dasatinib) should only be prescribed by a qualified physician who is experienced in the use of antineoplastic therapy.

1.1 Pediatrics

The safety and efficacy of SPRYCEL in patients <18 years of age have not been established. Therefore Health Canada has not authorized an indication for pediatric use. Nonclinical studies demonstrated greater toxicity in rat pups (See 7.1 Special populations).

1.2 Geriatrics

The safety profile of SPRYCEL in the geriatric population was similar to that in the younger population (see 7.1.4 Geriatrics).

2 CONTRAINDICATIONS

- Breastfeeding is contraindicated in women taking dasatinib (See 7.1.2 Breast-feeding).
- Use of SPRYCEL is contraindicated in patients with hypersensitivity to dasatinib or to any other component of SPRYCEL.

3 SERIOUS WARNINGS AND PRECAUTIONS BOX

Serious Warnings and Precautions

- SPRYCEL (dasatinib) should only be prescribed by a qualified physician who is experienced in the use of antineoplastic therapy.
- Myelosuppression: thrombocytopenia, neutropenia, and anemia (see 7 WARNINGS AND PRECAUTIONS, Hematologic).
- Hemorrhage, including fatal outcomes (see 7 WARNINGS AND PRECAUTIONS, Hemorrhage).
- Fluid retention, pleural effusion, pulmonary edema and—pericardial effusion (see 7 WARNINGS AND PRECAUTIONS, Respiratory).
- Congestive heart failure (see 7 WARNINGS AND PRECAUTIONS, Cardiovascular).
- Pulmonary arterial hypertension (See 7 WARNINGS AND PRECAUTIONS, Respiratory below)

4 DOSAGE AND ADMINISTRATION

4.1 Dosing Considerations

Dose reduction for concomitant use of strong CYP3A4 inhibitors

The concomitant use of strong CYP3A4 inhibitors and grapefruit juice with SPRYCEL should be avoided (see 9.4 Drug-Drug Interactions, Table 8). CYP3A4 inhibitors such as ketoconazole may increase SPRYCEL plasma concentrations. If possible, an alternative concomitant medication with no or minimal enzyme inhibition potential should be selected. If SPRYCEL must be administered with a strong CYP3A4 inhibitor, consider a dose decrease to:

40 mg daily for patients taking SPRYCEL 140 mg daily.

20 mg daily for patients taking SPRYCEL 100 mg daily.

20 mg daily for patients taking SPRYCEL 70 mg daily.

For patients taking SPRYCEL 60 mg or 40 mg daily, consider interrupting SPRYCEL until the inhibitor is discontinued. Allow a washout period of approximately 1 week after the inhibitor is stopped before reinitiating SPRYCEL.

The reduced doses of SPRYCEL are predicted to adjust the area under the curve (AUC) to the range observed without CYP3A4 inhibitors; however, clinical data are not available with these dose adjustments in patients receiving strong CYP3A4 inhibitors. If SPRYCEL is not tolerated after dose reduction, either discontinue the strong CYP3A4 inhibitor or stop SPRYCEL until the inhibitor is discontinued. Allow a washout period of approximately 1 week after the inhibitor is stopped before the SPRYCEL dose is increased.

4.2 Recommended Dose and Dosage Adjustment

- The recommended starting dosage of SPRYCEL (dasatinib) for chronic phase CML is 100 mg administered orally once daily (OD), either in the morning or in the evening.
- The recommended starting dosage of SPRYCEL for accelerated phase CML, or myeloid or

lymphoid blast CML, is 140 mg/day administered orally once daily (140 mg QD) either in the morning or in the evening.

- The recommended starting dosage of SPRYCEL for Ph+ ALL is 140 mg administered orally once daily (140 mg QD) either in the morning or in the evening.

Dosing recommendations in patients with imatinib resistant or intolerant CML and Ph+ ALL are based on the results of two randomized Phase III dose-optimization studies (see 14 CLINICAL TRIALS).

In clinical studies, treatment with SPRYCEL was continued until disease progression or until no longer tolerated by the patient. The effect of stopping treatment on long-term disease outcome after the achievement of a complete cytogenetic response ([CCyR]) or major molecular response (MMR) has not been investigated.

Dose Escalation

In clinical studies of adult CML and Ph+ ALL patients, dose escalation to 140 mg once daily (chronic phase CML) or 180 mg once daily (advanced phase CML and Ph+ ALL) was allowed in patients who did not achieve a hematologic or cytogenetic response at the recommended dosage.

Dose Adjustment for Adverse Reactions

Myelosuppression

In clinical studies, myelosuppression was managed by dose interruption, dose reduction, or discontinuation of study therapy. Hematopoietic growth factor has been used in patients with resistant myelosuppression. Guidelines for dose modifications are summarized in Table 1.

Table 1 - Dose Adjustments for Neutropenia and Thrombocytopenia

Chronic Phase CML (starting dose 100 mg once daily)	ANC* <0.5 × 10^9/L and/or Platelets <50 × 10^9/L	1. Stop SPRYCEL until ANC ≥1.0 × 10^9/L and platelets ≥50 × 10^9/L. 2. Resume treatment with SPRYCEL at the original starting dose. 3. If platelets <25 × 10^9/L and/or recurrence of ANC <0.5× 10^9/L for >7 days, repeat Step 1 and resume SPRYCEL at a reduced dose of 80 mg once daily for second episode. For third episode, further reduce dose to 50 mg once daily (for newly diagnosed patients) or discontinue SPRYCEL (for patients resistant or intolerant to prior therapy including imatinib).
Accelerated Phase CML, Blast Phase CML and Ph+ ALL (starting dose 140 mg once daily)	ANC* <0.5 × 10^9/L and/or Platelets <10 × 10^9/L	1. Check if cytopenia is related to leukemia (marrow aspirate or biopsy). 2. If cytopenia is unrelated to leukemia, stop SPRYCEL until ANC ≥1.0 × 10^9/L and platelets ≥20 × 10^9/L and resume at the original starting dose. 3. If recurrence of cytopenia, repeat Step 1 and resume SPRYCEL at a reduced dose of

Table 1 - Dose Adjustments for Neutropenia and Thrombocytopenia

		100 mg once daily (second episode) or 80 mg once daily (third episode). 4. If cytopenia is related to leukemia, consider dose escalation to 180 mg once daily.

*ANC: absolute neutrophil count

Non-hematological adverse reactions

If a moderate (Grade 2) non-hematological adverse reaction develops with SPRYCEL, treatment should be interrupted until the adverse reaction has resolved or returned to baseline. The same dose should be resumed if this is the first occurrence and the dose should be reduced if this is a recurrent adverse reaction.

If a severe (Grade 3 or 4) non-hematological adverse reaction develops with SPRYCEL use, treatment must be withheld until the event has resolved or improved. Thereafter, treatment can be resumed as appropriate at a reduced dose depending on the initial severity of the event. However, in patients diagnosed with pulmonary arterial hypertension (PAH), SPRYCEL should be permanently discontinued.

Patients with chronic CML who received 100 mg once daily, dose reduction to 80 mg once daily with further reduction from 80 mg once daily to 50 mg once daily, if needed, is recommended. For adult patients with advanced phase CML or Ph+ ALL who received 140 mg once daily, dose reduction to 100 mg once daily with further reduction from 100 mg once daily to 80 mg once daily, if needed, is recommended.

Hepatic impairment:

No clinical pharmacokinetic trials were conducted with a 70-100 mg dose of SPRYCEL in patients with decreased liver function. SPRYCEL should be used with caution in patients with moderate to severe hepatic impairment (see 7 WARNINGS AND PRECAUTIONS, Hepatic/Biliary/Pancreatic).

Renal impairment:

No clinical trials were conducted with SPRYCEL in patients with decreased renal function (trials excluded patients with serum creatinine concentration > 1.5 times the upper limit of the normal range). Since the renal clearance of dasatinib and its metabolites is < 4%, a decrease in total body clearance is not expected in patients with renal insufficiency.

Pediatrics (< 18 years of age):

The safety and efficacy of SPRYCEL in patients <18 years of age have not been established. Therefore Health Canada has not authorized an indication for pediatric use.

4.4 Administration

SPRYCEL can be taken with or without food. Tablets should not be crushed or cut; they should be swallowed whole.

4.5 Missed Dose

If a dose is missed, the next dose should be taken at the usual time. Do not double dose.

5 OVERDOSAGE

Experience with overdose of SPRYCEL in clinical studies is limited to isolated cases. The highest reported dosage ingested was 280 mg per day for 1 week in two patients and both developed a significant decrease in platelet counts. Since SPRYCEL is associated with severe myelosuppression (see 7 WARNINGS AND PRECAUTIONS and 8 ADVERSE REACTIONS), patients who ingested more than the recommended dosage should be closely monitored for myelosuppression and appropriate supportive treatment given.

For management of a suspected drug overdose, contact your regional poison control centre.

6 DOSAGE FORMS, STRENGTHS, COMPOSITION AND PACKAGING

Table 2 - Dosage Forms, Strengths, Composition and Packaging

Route of Administration	Dosage Form / Strength/Composition	Non-medicinal Ingredients
Oral	Tablet / 20 mg, 50 mg, 70 mg, 80 mg, 100 mg and 140 mg	Tablet core: croscarmellose sodium, hydroxypropyl cellulose, lactose monohydrate, magnesium stearate and microcrystalline cellulose. Film-coating: hypromellose, polyethylene glycol and titanium dioxide.

20 mg: white to off-white, biconvex, round, film coated tablet with “BMS” debossed on one side and “527” on the other.

50: white to off-white, biconvex, oval, film coated tablet with “BMS” debossed on one side and “528” on the other side.

70 mg: white to off-white, biconvex, round, film coated tablet with “BMS” debossed on one side and “524” on the other side

80 mg: white to off-white, biconvex, triangle, film coated tablet with “BMS” and “80” (BMS over 80) debossed on one side and “855” on the other side

100 mg: white to off-white, biconvex, oval, film coated tablet with “BMS 100” debossed on one side and “852” on the other side

140 mg: white to off-white, biconvex, round, film-coated tablet with “BMS” and “140” (BMS over 140) debossed on one side and “857” on the other side.

20 mg, 50 mg and 70 mg, are supplied in HDPE bottles containing 60 tablets and in blister packs of 60 tablets.

80 mg, 100 mg and 140 mg are supplied in HDPE bottles containing 30 tablets and in blister packs of 30 tablets.

7 WARNINGS AND PRECAUTIONS

Please see 3 SERIOUS WARNINGS AND PRECAUTIONS BOX.

Carcinogenesis and Mutagenesis

Please see 16 NON-CLINICAL TOXICOLOGY

Cardiovascular

The Phase III clinical study in patients with newly diagnosed CML in chronic phase excluded patients with uncontrolled or significant cardiovascular disease. The SPRYCEL arm (n=258) included 1.6 % of patients with prior cardiac disease and 24% with baseline cardiovascular risk factors. Cardiac adverse reactions of congestive heart failure/cardiac dysfunction, pericardial effusion, arrhythmias, palpitations, QT prolongation, and myocardial infarction (including fatal) were reported in patients taking SPRYCEL (see 8 ADVERSE REACTIONS). Severe pericardial effusion (1.2%) and arrhythmia (0.4%) were also reported in patients. Adverse cardiac events were more frequent in patients with cardiovascular risk factors or a previous medical history of cardiac disease (see 8 ADVERSE REACTIONS). Patients with risk factors or a history of cardiac disease should be evaluated at baseline and monitored carefully for clinical signs or symptoms consistent with cardiac dysfunction (such as chest pain, shortness of breath, and diaphoresis) during routine follow up.

In the Phase III clinical trials in patients with resistance or intolerance to prior imatinib therapy, patients were excluded from enrolment for a broad range of cardiac events or conditions. A significantly abnormal ECG at screening was also an exclusion criterion. No prospective evaluation of cardiac function was carried out.

In all clinical trials with patients resistant or intolerant to prior imatinib therapy, congestive heart failure/cardiac dysfunction was reported in 96 (4%) of subjects, of which 49 (2%) were considered to be severe. In some cases, the event was triggered by an acute volume load, including transfusion of blood products.

QT Prolongation

In vitro data suggest that dasatinib and its N-dealkylated metabolite, BMS-582691 have the potential to prolong cardiac ventricular repolarization (QT interval, see Safety Pharmacology).

In 865 patients with leukemia treated with SPRYCEL in Phase II clinical studies, the mean changes from baseline in QTcF interval were 4–6 msec; the upper 95% confidence intervals for all mean changes from baseline were <7 msec. Of the 2182 patients with resistance or intolerance to prior imatinib therapy who received SPRYCEL in clinical studies, 21 patients (<1%) experienced a QTcF >500 msec.

In the Phase III clinical study in patients with newly diagnosed CML in chronic phase, patients with baseline QTcF interval > 450 msec were excluded. After 5 years of follow-up, QTc prolongation was reported in one patient (<1%) who experienced a QTcF >500 msec and discontinued SPRYCEL treatment. SPRYCEL should be administered with caution in patients who have or may develop prolongation of QTc. These include patients with hypokalemia or hypomagnesemia, patients with congenital long QT syndrome, patients taking anti arrhythmic medicines or other medicinal products that lead to QT prolongation, and cumulative high-dose anthracycline therapy.

Hypokalemia or hypomagnesemia should be corrected prior to administration of SPRYCEL. (See 9 DRUG INTERACTIONS, Drugs that prolong QTc interval or induce torsade de pointes, 10.2 Pharmacodynamics, Electrocardiogram.)

Also refer to Respiratory below for information concerning fluid retention.

Endocrine and Metabolism

Lactose
SPRYCEL tablets 20 mg, 50 mg, 70 mg, 80 mg, 100 mg and 140 mg contain lactose in proportional amounts of 27 mg, 67.5 mg, 94.5 mg, 108 mg, 135 mg and 189 mg, respectively. SPRYCEL therefore contains 189 mg of lactose in the 140 mg daily dose of dasatinib and 135 mg in the 100 mg daily dose. Patients with rare hereditary problems of galactose intolerance, the Lapp lactase deficiency or glucose-galactose malabsorption should not take dasatinib.

Hematologic

Hemorrhage
Nonclinical studies have shown that dasatinib inhibits platelet aggregation *in vitro* and *in vivo* and increases bleeding time *in vivo* (see16 NON-CLINICAL TOXICOLOGY, Special Toxicity). Patients with a history of significant bleeding disorder unrelated to CML were excluded in SPRYCEL clinical studies. Patients taking concomitant medications that inhibit platelet function or anticoagulants were excluded in initial imatinib-resistant SPRYCEL (dasatinib) clinical studies. In subsequent trials, the use of anticoagulants, aspirin, and non-steroidal anti-inflammatory drugs (NSAIDs) was allowed concurrently with SPRYCEL if the platelet count was >50,000 per microliter. Caution should be exercised when SPRYCEL is to be concurrently administered with anticoagulants (see 9 DRUG INTERACTIONS).

In clinical studies in 2,712 CML or Ph+ ALL patients with a median duration of therapy of 19.2 months (range 0- 93.2 months), 272 (10%) patients experienced Grade 3-4 bleeding. Fifty-six (2%) patients experienced fatal bleeding. In 23 (1%) of these patients, fatal bleeding occurred more than 30 days after dasatinib discontinuation.

Intracranial hemorrhage occurred in 66 (2.4%) of 2,712 CML or Ph+ ALL patients, of which 27 (1%) cases were considered related to SPRYCEL. Intracranial hemorrhage was fatal in 25 (0.9%) of these patients, of which ten (0.4%) cases were considered related to SPRYCEL.

Gastrointestinal hemorrhage regardless of relationship to SPRYCEL occurred in 15 % of 2,712 CML or Ph+ ALL patients. The bleeding was severe in 6 % of these patients and generally required treatment interruptions and packed cell transfusions. Other episodes of severe bleeding occurred in 3% of patients.

Grade 3-4 hemorrhages were reported in 2.3% of 258 patients with newly diagnosed chronic phase CML (see 8 ADVERSE REACTIONS).

Myelosuppression
Treatment with SPRYCEL (dasatinib) is associated with thrombocytopenia, neutropenia, and anemia which occur earlier and more frequently in patients with advanced phase CML or Ph+ ALL than in patients with chronic phase CML. In a Phase III dose-optimization study in patients with chronic phase CML with resistance or intolerance to prior imatinib therapy with a minimum follow-up of 24 months, Grade 3 or 4 myelosuppression was reported less frequently in patients treated with 100 mg once daily (neutropenia 35%, thrombocytopenia 23% and anemia 13%) than in patients treated with 70 mg twice daily (neutropenia 45%, thrombocytopenia 38% and anemia 18%). Severe febrile neutropenia (including fatal outcomes) was reported in 2% of chronic phase patients and 14% of advanced phase CML patients.

In patients with advanced phase CML or Ph+ ALL treated with dasatinib complete blood counts (CBCs) should be performed weekly for the first 2 months and then monthly thereafter, or as clinically indicated.

In patients with chronic phase CML, CBCs should be performed every 2 weeks for 12 weeks, then every 3 months thereafter or as clinically indicated.

Myelosuppression was generally reversible and usually managed by withholding SPRYCEL temporarily or dose reduction (see 4.2 Recommended Dose and Dosage Adjustment and 8.4 Abnormal Laboratory Findings: Hematologic, Clinical Chemistry and Other Quantitative Data). In clinical studies in patients with resistance or intolerance to prior imatinib therapy, severe (CTC Grade 3 or 4) cases of anemia were managed with blood transfusions. Packed red blood cells were transfused in 30% of chronic phase CML patients and 79% of myeloid blast phase CML patients. Platelet transfusions were required in 17% of chronic phase CML patients and 66% of myeloid blast phase CML patients

Hepatic/Biliary/Pancreatic

The effect of hepatic impairment on the single-dose pharmacokinetics of dasatinib was assessed in 8 moderately hepatic impaired subjects who received a 50-mg dose and 5 severely hepatic-impaired subjects who received a 20-mg dose compared to matched healthy subjects who received a 70-mg dose of SPRYCEL. Hepatic impairment did not result in clinically meaningful change in dasatinib exposure at the doses studied. However no pharmacokinetic information is available from patients with hepatic impairment treated with a 70-100 mg dose of SPRYCEL (see 10.3 Pharmacokinetics, Special Populations and Conditions). Due to the limitations of this clinical study, caution is recommended in patients with hepatic impairment.

In nonclinical studies, increased liver weight and foci of hepatocellular alteration were observed in rats, and hepatocellular vacuolation was observed in monkeys following repeat dose administration of dasatinib (6 to 9 months). Increased ALT was observed in monkeys, and increased AST and/or decreased albumin were observed in rats and monkeys.

In clinical studies with 2,712 patients, 4 cases of hepatotoxicity, 4 cases of hepatocellular injury, 4 cases of hepatic steatosis, 2 cases of jaundice, 2 cases of liver disorder, 1 case of toxic hepatitis, 1 case of hepatic failure, 2 cases of abnormal hepatic function and 1 case of hepatitis were observed.

Immune

Hepatitis B virus reactivation

Reactivation of hepatitis B virus (HBV) has occurred in patients who are chronic carriers of this virus after receiving a BCR-ABL tyrosine kinase inhibitor (TKI), including SPRYCEL. Some cases resulted in acute hepatic failure or fulminant hepatitis leading to liver transplantation or death.

Patients should be tested for HBV infection before initiating treatment with SPRYCEL. Experts in liver disease and in the treatment of HBV should be consulted before treatment is initiated in patients with positive HBV serology (including those with active disease) and for patients who test positive for HBV infection during treatment. Carriers of HBV who require treatment with SPRYCEL should be closely monitored for signs and symptoms of active HBV infection throughout therapy and for several months following termination of therapy.

Monitoring and Laboratory Tests

In patients with chronic phase CML, complete blood counts (CBCs) should be performed every two weeks for 12 weeks, then every 3 months thereafter, or as clinically indicated. In patients with advanced phase CML or Ph+ ALL, CBC should be performed weekly for the first 2 months and then monthly thereafter, or as clinically indicated (see 7 WARNINGS AND PRECAUTIONS, Immune).

Hepatic function tests (AST, ALT and bilirubin), CK and renal function tests should be performed every two weeks for the first 2 months and then monthly thereafter or as clinically indicated (see 7 WARNINGS AND PRECAUTIONS: Hepatic Impairment and Musculoskeletal).

Musculoskeletal

Cases of rhabdomyolysis with acute renal failure have been reported. Patients with muscle symptoms (muscle aches/pains) should be investigated to rule out rhabdomyolysis (elevated creatine kinase, elevated serum creatinine, hyperkalemia, hyperphosphatemia, brown urine, elevated ALT and AST).

Renal

There are currently no clinical studies with SPRYCEL in patients with impaired renal function. The study in patients with newly diagnosed chronic phase CML excluded patients with serum creatinine concentration > 3 times the upper limit of the normal range, and studies in patients with chronic phase CML with resistance or intolerance to prior imatinib therapy excluded patients with serum creatinine concentration >1.5 times the upper limit of the normal range. Dasatinib and its metabolites are minimally excreted via the kidney. Since the renal excretion of unchanged dasatinib and its metabolites is <4%, a decrease in total body clearance is not expected in patients with renal insufficiency. The effect of dialysis on dasatinib pharmacokinetics has not been studied.

Reproductive Health: Female and Male Potential

- **Fertility**

  The effects of SPRYCEL on male and female fertility in humans are not known. Based on animal studies, SPRYCEL may impair fertility in females of reproductive potential (See 16 NON-CLINICAL TOXICOLOGY, Reproductive and Developmental Toxicology).

- **Teratogenic Risk**

  Dasatinib can cause fetal harm when administered to pregnant women. Knowledge of the potential effects of SPRYCEL on the sperm of male patients, and the level of maternal or fetal exposure from the semen of male SPRYCEL patients, is limited. Sexually active male patients or female patients of child bearing potential taking SPRYCEL should use highly effective contraception.

Respiratory

Pulmonary Arterial Hypertension

Serious cases of pulmonary arterial hypertension (PAH), confirmed by right heart catheterization, have been associated with SPRYCEL treatment in clinical trials and post-marketing reports. In these cases, PAH was reported after initiation of SPRYCEL therapy, including after more than one year of treatment. In the Phase III clinical study in patients with newly diagnosed CML in chronic phase, drug-related pulmonary hypertension was reported in 4.7% of dasatinib-treated patients (N= 12) compared to 0.4% of imatinib-treated patients. Additional evaluation by right heart catheterization to determine if PAH was present was only

performed in one case where PAH was not identified and pulmonary hypertension was not confirmed.

Patients should be evaluated for signs and symptoms of underlying cardiopulmonary disease prior to initiating SPRYCEL therapy. Patients who develop symptoms suggestive of PAH such as dyspnea and fatigue after initiation of therapy should be evaluated for more common etiologies including pleural effusion, pulmonary edema, anemia, or lung infiltration. If no alternative diagnosis is found, the diagnosis of PAH should be considered. If the symptoms are severe, SPRYCEL should be withheld during this evaluation. SPRYCEL should be permanently discontinued if PAH is confirmed (see 4.2 Recommended Dose and Dosage Adjustment, Dose Adjustment for Adverse Reactions). Follow up on patients with PAH should be performed according to standard practice guidelines. Improvements in hemodynamic and clinical parameters have been observed in patients with PAH following cessation of SPRYCEL therapy.

Fluid Retention

SPRYCEL is associated with fluid retention. Patients with pre-existing pleural effusion were excluded from Phase III studies.

In the Phase III dose-optimization studies in patients with resistance or intolerance to prior imatinib therapy, severe fluid retention was reported in 11% of patients, including severe pleural and pericardial effusion reported in 7% and 2% of patients, respectively. Severe ascites and generalized edema were each reported in <1% of patients. Other manifestations of fluid retention in these studies included pulmonary edema (3%), congestive heart failure/cardiac dysfunction (4%), and pericardial effusion (5%). Nineteen patients had severe pulmonary edema. In patients with chronic phase CML with resistance or intolerance to prior imatininb therapy, Grade 3 or 4 fluid retention events were reported less frequently in patients treated with 100 mg once daily (5%) than in patients treated with 140 mg once daily (9%) (See 8.2 Clinical Trial Adverse Reactions). In these studies, fluid retention events were typically managed by supportive care measures that include diuretics or short courses of steroids. Pleural effusion required oxygen in some cases and at least one thoracentesis in 64 (3%) patients.

In the Phase III study conducted with newly diagnosed chronic phase CML patients, grades 1-4 fluid retention and pleural effusion were reported in 22% and 10%, respectively, by 12 months of treatment (see 8.2 Clinical Trial Adverse Reactions). The median time to onset of pleural effusion was 28 weeks (range 4-88 weeks). With appropriate medical care, 23 patients (88% of those with pleural effusion) were able to continue on SPRYCEL. After 5 years follow-up, fluid retention and pleural effusion were reported in 43% and 29% of patients, respectively. The median time to first grade 1-2 pleural effusion was 114 weeks and to first grade 3-4 pleural effusions was 175 weeks. Dasatinib treatment was discontinued due to pleural effusion in 5.8% of all dasatinib-treated patients. Out of patients with a pleural effusion, dasatinib treatment was interrupted in 62% and dose reduced in 41%, and was also managed through the use of diuretics or other appropriate supportive care measures.

In all patients with newly diagnosed or imatinib resistant or intolerant patients with chronic phase CML (n=548), severe fluid retention occurred in 36 (7%) patients receiving SPRYCEL at the recommended dose. In patients with advanced phase CML or Ph+ ALL treated with SPRYCEL at the recommended dose (n=304), severe fluid retention was reported in 11% of patients, including severe pleural effusion reported in 8% of patients.

Patients who develop symptoms suggestive of pleural effusion or other fluid retention such as new or worsened dyspnea on exertion or at rest, pleuritic chest pain, or dry cough should be

evaluated promptly with chest X-ray or additional diagnostic imaging as appropriate (see 4 DOSAGE AND ADMINISTRATION and 8 ADVERSE REACTIONS). Consider treatment interruption, dose reduction, or treatment discontinuation.

Skin

Individual cases of severe mucocutaneous dermatologic reactions, including Stevens-Johnson syndrome and erythema multiforme, have been reported with the use of SPRYCEL. SPRYCEL should be permanently discontinued in patients who experience a severe mucocutaneous reaction during treatment if no other etiology can be identified.

7.1 Special Populations

7.1.1 Pregnant Women

Dasatinib can cause fetal harm when administered to pregnant women. There have been post-marketing reports of spontaneous abortion and fetal and infant anomalies from women who have taken SPRYCEL during pregnancy (see 8.5 Post-Market Adverse Reactions). Studies in animals have shown that at concentrations which are readily achievable in humans receiving therapeutic doses of SPRYCEL, fetal toxicity (enbryofetal lethality, skeletal abnormalities including malformations) was observed in both pregnant rats and rabbits. Fetal death was observed in rats (see 16 NON-CLINICAL TOXICOLOGY, Reproductive and Developmental Toxicology).

SPRYCEL therefore should not be used in women who are pregnant or contemplating pregnancy. Women of child bearing potential must be advised to use highly effective contraception (i.e. a method of birth control that results in a failure rate less than 1% per year when used consistently and correctly) during SPRYCEL treatment. If SPRYCEL is used during pregnancy, or if the patient becomes pregnant while taking SPRYCEL, the patient should be apprised of the potential hazard to the fetus.

7.1.2 Breast-feeding

It is unknown whether SPRYCEL is excreted in human milk. In an exploratory pre- and post-natal development study in rats, postnatal exposure to dasatinib through lactation resulted in pleural effusion and mortality in pups before postnatal age of 20 days at an exposure of 0.27 times the adult clinical dose (see 16 NON-CLINICAL TOXICOLOGY, Reproductive and Developmental Toxicology). Women who are taking SPRYCEL must not breastfeed (See 2 CONTRAINDICATIONS).

7.1.3 Pediatrics

The safety and efficacy of SPRYCEL in patients <18 years of age have not been established. Therefore Health Canada has not authorized an indication for pediatric use.

7.1.4 Geriatrics

In the newly diagnosed chronic phase CML study, 25 patients (10%) were 65 years of age and older and 7 patients (3%) were 75 years of age and older. Patients of 65 years and over had more serious adverse events reported (any or drug-related) compared to those under 65 years (40.7% vs. 29.7%, 16.7% vs. 12.1%, respectively). Of the 2,712 patients in clinical studies of SPRYCEL, 617 (23%) were 65 years of age and older and 123 (5%) were 75 years of age and older. While the safety profile of SPRYCEL in the geriatric population was similar to that in the younger population, patients aged 65 years and older are more likely to experience the commonly reported adverse reactions diarrhea, fatigue, cough, pleural effusion, dyspnea,

dizziness, peripheral edema, pneumonia, hypertension, arrhythmia, congestive heart failure, pericardial effusion, lower gastrointestinal hemorrhage, abdominal distension and more likely to experience the less frequently reported events pulmonary edema, lung infiltration, arthritis, and urinary frequency and should be monitored closely. No differences in cCCyR and MMR were observed between older and younger patients. However, in the two randomized studies in patients with imatinib resistant or intolerant chronic phase CML, the rates of major cytogenetic response (MCyR) at 2 years were lower among patients aged 65 years and older (42% MCyR in patients ≥ 65 years versus 56% MCyR in the rest of the study population and 47% MCyR in patients ≥ 65 years versus 68% MCyR in the rest of the study population in studies CA180017 and CA180034, respectively).

8 ADVERSE REACTIONS

8.1 Adverse Reaction Overview

The data described below reflect exposure to SPRYCEL at all doses studied from clinical studies in 2,712 patients, including 324 patients with newly diagnosed chronic phase CML and 2388 patients with imatinib intolerant or resistant chronic or advanced phase CML or Ph+ ALL. The median duration of therapy in 2,712 SPRYCEL treated patients was 19.2 months (range 0-93.2 months).

The majority of SPRYCEL-treated patients experienced adverse events at some time. Most events were mild to moderate. In the overall population of 2,712 SPRYCEL-treated subjects, 798 (29.4%) experienced adverse events leading to treatment discontinuation. Among the 258 patients in the Phase III newly diagnosed chronic phase CML study with follow up over a minimum of 60 months, serious adverse events, regardless of relationship to SPRYCEL, were reported in 35% of patients treated with SPRYCEL. A total of 69% of patients had dose interruption and 37% had dose reduction.

SPRYCEL was discontinued due to study drug toxicity in 14% of SPRYCEL-treated patients with a minimum of 60 months follow-up. The reasons for discontinuation were thrombocytopenia, leukopenia, pleural effusion, colitis, creatinine kinase increased, pericardial effusion, prolonged QTc interval, chest pain, optic neuritis, pulmonary hypertension, dyspnea, pleurisy, pneumothorax, acute myocardial infarction, abdominal discomfort, abdominal pain, colitis, diarrhea, peripheral edema, and acute renal failure.

Among the 1,618 SPRYCEL-treated subjects with chronic phase CML, adverse reactions leading to discontinuation were reported in 329 (20.3%) subjects, and among the 1,094 SPRYCEL-treated subjects with advanced phase disease (including Ph+ ALL), adverse reactions leading to discontinuation were reported in 191 (17.5%) subjects.

In a Phase III dose-optimization study in chronic phase CML patients resistant or intolerant to prior imatinib therapy with a minimum of 84 months follow-up, the rate of discontinuation for adverse reactions was 21% in patients treated with 100 mg once daily.

The median time to onset for Grade 1 or 2 pleural effusion events was 114 weeks (range 4-299 weeks). Fewer than 3% of pleural effusion events were Grade 3 or 4. With appropriate medical care, 58 patients (80% of those with pleural effusion) were able to continue on SPRYCEL (See 7 WARNINGS AND PRECAUTIONS, Respiratory).

With a minimum of 60 months of follow up, the most frequently adverse events reported in SPRYCEL-treated patients with newly diagnosed chronic phase CML were fluid retention (including pleural effusion, superficial edema, pulmonary hypertension, generalized edema,

pericardial effusion, congestive heart failure/cardiac dysfunction, pulmonary edema), diarrhea , infection (including bacterial, viral, fungal and non-specified), upper respiratory tract infection/inflammation, musculoskeletal pain, headache, cough, rash, pyrexia, and abdominal pain.

With a minimum of 84 months of follow up, in 165 patients with chronic phase CML resistant or intolerant to prior imatinib therapy treated with the recommended dose of 100 mg once daily, the most frequently reported adverse events, regardless of causality or severity, were diarrhea, fluid retention, headache, musculoskeletal pain, hemorrhage, pyrexia, fatigue, infection, skin rash, nausea, dyspnea, cough, upper respiratory tract infection/inflammation, vomiting, pain, abdominal pain, arthralgia, myalgia, pruritis and constipation.

8.2 Clinical Trial Adverse Reactions

Clinical trials are conducted under very specific conditions. The adverse reaction rates observed in the clinical trials, therefore, may not reflect the rates observed in practice and should not be compared to the rates in the clinical trials of another drug. Adverse reaction information from clinical trials may be useful in identifying and approximating rates of adverse drug reactions in real-world use.

Newly diagnosed patients with chronic phase CML

In the Phase III study in patients with newly diagnosed chronic phase CML the median duration of therapy was 60 months for both groups (range: < 1 to 73 months for the SPRYCEL group and <1 month to 75 months in the imatinib group); the median average daily dose was 99 mg and 400 mg, respectively.

All treatment-emergent adverse events (excluding laboratory abnormalities), regardless of relationship to study drug, that were reported in at least 5% of the patients are shown in Table 3.

A total of 26 (10%) SPRYCEL-treated patients died (11 of infections and 2 of myocardial infarction) and a total of 26 patients (10%) in the imatinib arm died (including 1 of myocardial infarction, 1 of pneumonia, 1 of fatal bleeding at time of disease progression and 2 of unknown cause/clinical deterioration and decrease in performance status).

Table 3 - Adverse Events Reported in ≥5% of Patients with Newly Diagnosed Chronic Phase CML - 60 month follow up

	SPRYCEL 100 mg QD n = 258		Imatinib 400 mg QD n = 258	
	All Grades	Grade 3/4	All Grades	Grade 3/4
SYSTEM ORGAN CLASS/ Preferred Term	**Percent (%) of patients**			
Any Adverse Event	95	27	95	24
Cardiovascular				
Pericardial effusion	5	1	2	0
Congestive heart failure/ cardiac dysfunction[a, *]	4	1	2	1
Gastrointestinal				

	SPRYCEL 100 mg QD n = 258		Imatinib 400 mg QD n = 258	
	All Grades	Grade 3/4	All Grades	Grade 3/4
SYSTEM ORGAN CLASS/ Preferred Term	**Percent (%) of patients**			
Diarrhea	40	2	35	2
Abdominal pain	22	1	17	<1
Vomiting	17	<1	21	<1
Nausea	15	0	29	0
Dyspepsia	11	0	12	0
Gastritis	10	<1	7	0
Mucosal inflammation (including mucositis/stomatitis)	9	<1	5	0
Constipation	8	0	3	0
Abdominal Distension	6	0	4	0
Ascites*	0	0	<1	0
General Disorders and Administration Site Conditions				
Pyrexia	23	1	20	<1
Fatigue	16	<1	16	0
Pain	16	1	15	<1
Asthenia	16	0	14	1
Face edema	12	0	38	0
Chest pain	11	0	5	0
Peripheral edema	9	0	13	<1
Generalized edema	5	0	9	0
Infections and Infestations				
Infection (including bacterial, viral, fungal, non-specified)	40	4	30	3
Upper respiratory tract infection/inflammation	38	1	38	1
Enterocolitis infection	11	0	6	<1
Investigations				
Weight increased	10	2	13	3
Metabolism and Nutrition Disorders				

	SPRYCEL 100 mg QD n = 258		Imatinib 400 mg QD n = 258	
	All Grades	Grade 3/4	All Grades	Grade 3/4
SYSTEM ORGAN CLASS/ Preferred Term	**Percent (%) of patients**			
Appetite disturbances	9	0	5	0
Musculoskeletal and Connective Tissue Disorders				
Musculoskeletal pain	31	<1	34	<1
Myalgia	14	<1	16	0
Arthralgia	14	0	16	<1
Muscle spasms	5	0	24	<1
Nervous System Disorders				
Headache	23	0	18	<1
Dizziness	11	<1	7	<1
Neuropathy (including peripheral)	10	<1	8	<1
Ophthalmologic				
Conjunctivitis	4	0	7	0
Psychiatric disorder				
Insomnia	8	0	6	0
Depression	2	0	5	<1
Respiratory, Thoracic and Mediastinal Disorders				
Pleural effusion	29	3	1	0
Cough	27	<1	11	0
Dyspnea	16	2	6	0
Pulmonary hypertension	5	1	<1	0
Pulmonary edema*	1	0	0	0
Skin and Subcutaneous Tissue Disorders				
Rash[b]	20	0	23	2
Pruritus	7	0	9	<1
Acne	6	0	2	0
Dermatitis including eczema	4	0	7	0
Pigmentation disorder	2	0	7	0
Hyperhidrosis	2	0	5	0

	SPRYCEL 100 mg QD n = 258		Imatinib 400 mg QD n = 258	
	All Grades	Grade 3/4	All Grades	Grade 3/4
SYSTEM ORGAN CLASS/ Preferred Term	**Percent (%) of patients**			
Vascular Disorders				
Hemorrhage	19	2	18	2
Other bleeding[c]	14	<1	15	2
Gastrointestinal bleeding	5	1	4	<1
CNS bleeding*	1	<1	<1	<1
Hypertension	11	<1	8	<1

[a] Includes cardiac failure, cardiac failure acute, cardiac failure congestive, cardiomyopathy, diastolic dysfunction, ejection fraction decreased, and left ventricular dysfunction.

[b] Includes erythema, erythema multiforme, heat rash, rash, rash erythematous, rash generalized, rash macular, rash papular, rash pustular, skin exfoliation, and rash vesicular.

[c] Includes conjunctival hemorrhage, ear hemorrhage, ecchymosis, epistaxis, eye hemorrhage, gingival bleeding, hematoma, hematuria, hemoptysis, hemorrhage, hemorrhage subcutaneous, intra-abdominal hematoma, menorrhagia, metrorrhagia, petechiae, scleral hemorrhage, uterine hemorrhage, and vaginal hemorrhage.

* Adverse events of special interest with <5% frequency.

Patients with imatinib intolerant or resistant CML or Ph+ ALL

All treatment-emergent adverse events (excluding laboratory abnormalities), regardless of relationship to study drug, that were reported in at least 5% of the patients treated with SPRYCEL at the recommended dose of 100 mg once daily in a Phase III clinical study of imatinib intolerant or resistant chronic phase CML are shown in

Table **4**.

In the Phase III dose-optimization study in patients with imatinib intolerant or resistant chronic phase CML, the median overall duration of therapy with 100 mg once daily was 30 months (range 1-93 months).

Table 4: Adverse Events Reported in ≥5% of Patients treated with 100 mg Once Daily dose in Clinical Studies of Imatinib Intolerant or Resistant Chronic Phase CML - 84 month follow up

SYSTEM ORGAN CLASS/ Preferred Term	Phase III	
	100 mg QD N=165	
	Percent (%) of patients	
	All Grades	Grade 3/4
Cardiology		
Arrhythmia (including tachycardia)	8	0
Palpitations	8	0

Pericardial effusion[a]	3	1
Congestive heart failure/cardiac dysfunction[a, b]	2	1
Gastrointestinal		
Diarrhea	42	4
Abdominal pain	24	2
Nausea	22	1
Constipation	18	2
Vomiting	14	1
Abdominal distension	12	0
Mucosal inflammation (including mucositis/stomatitis)	10	0
Dyspepsia	8	0
Ascites[b]	1	0
General Disorders and Administration Site Conditions		
Fatigue	37	4
Pain	27	1
Superficial edema[c]	26	1
Pyrexia	21	1
Chest pain	17	2
Asthenia	9	1
Chills	7	0
Generalized edema	5	1
Immune System Disorders		
Hypersensitivity (including erythema nodosum)	5	1
Infections and Infestations		
Infection (including bacterial, viral, fungal, non-specified)	48	6
Upper respiratory tract infection/inflammation	43	1
Pneumonia (including bacterial, viral, and fungal)	13	5
Enterocolitis infection	7	2
Herpes virus infection	5	1

Investigations		
Weight increased	11	1
Weight decreased	8	0
Metabolism and Nutrition Disorders		
Appetite Disturbances	10	0
Hyperuricemia	5	1
Musculoskeletal and Connective Tissue Disorders		
Musculoskeletal pain	48	3
Arthralgia	30	2
Myalgia	17	0
Muscle spasms	6	0
Arthritis	5	0
Nervous System Disorders		
Headache	48	1
Dizziness	16	2
Neuropathy (including peripheral neuropathy)	14	1
Ophthalmologic		
Visual disorder	7	0
Psychiatric disorder		
Insomnia	12	0
Depression	11	1
Anxiety	5	0
Respiratory, Thoracic and Mediastinal Disorders		
Dyspnea	34	2
Cough	34	1
Pleural effusion	28	5
Pulmonary hypertension[a]	2	1
Pulmonary edema[a]	1	0
Skin and Subcutaneous Tissue Disorders		
Skin rash	33	2
Pruritus	17	1
Hyperhidrosis	10	0
Alopecia	8	0
Dry skin	6	0

Acne	5	0
Vascular Disorders		
Hemorrhage	27	3
Gastrointestinal bleeding	6	1
CNS bleeding	0	0
Hypertension	9	0
Flushing	6	0

[a] Adverse events of special interest with <5% frequency.

[b] Includes ventricular dysfunction, cardiac failure, cardiac failure congestive, cardiomyopathy, congestive cardiomyopathy, diastolic dysfunction, ejection fraction decreased and ventricular failure

[c] Superficial edema is a grouped term composed of face edema, other superficial edema, and peripheral edema

With a minimum follow-up of 84 months, long-term cumulative safety data are available for the 100 mg once daily dose. Due to the allowance of switching to the 100 mg once daily dosing in the other three arms of the trial, safety results of these treatment groups are similar to the 100 mg once daily dose. Adverse events (all grades) that continued to occur in patients treated on the 100 mg once daily schedule at 2 and 7 years included: overall fluid retention (34% vs. 48%), pleural effusion (18% vs. 28%), and superficial edema (18% vs. 22%). Grade 3 or 4 pleural effusion among patients treated with 100 mg once daily at 2 and 7 years was 2% vs. 5%, respectively.

In the Phase III dose-optimization study exploring the once daily schedule of SPRYCEL (140 mg once daily) in patients with imatinib intolerant or resistant advanced diseases, the median duration of therapy was 13.62 months (range .03–31.15 months) for accelerated phase CML, 3.19 months (range .03–27.73 months) for myeloid blast CML, 3.55 months (range .10–22.08 months) for lymphoid blast CML, and 2.99 months (range .16–23.46 months) for Ph+ ALL.

Table 5: Adverse Events Reported in ≥5% of Patients treated with 140 mg daily dose in Clinical Studies of Imatinib Intolerant or Resistant Advanced Phase CML and Ph+ ALL

SYSTEM ORGAN CLASS/ Preferred Term	**Phase III**	
	140 mg QD n = 304	
	Percent (%) of patients	
	All Grades	Grade 3/4
Blood and Lymphatic Systems Disorders		
Febrile neutropenia	12	12
Cardiology		
Arrhythmia (including tachycardia)	13	1

Congestive heart failure/ cardiac dysfunction[a, b]	3	1
Pericardial effusion[a]	2	1
Gastrointestinal Disorders		
Diarrhea	44	6
Nausea	34	2
Vomiting	28	1
Abdominal pain	20	4
Mucosal inflammation (including mucositis/stomatitis)	17	1
Constipation	15	1
Dyspepsia	9	0
Ascites[a]	<1	<1
General Disorders and Administration Site Conditions		
Pyrexia	39	3
Fatigue	29	5
Superficial edema[c]	25	<1
Pain	24	2
Asthenia	13	3
Chest pain	13	1
Generalised oedema[a]	3	<1
Infections and Infestations		
Infection	46	14
Upper respiratory tract infection/inflammation	26	1
Pneumonia (including bacterial, viral, and fungal)	17	9
Sepsis (including fatal outcomes)	6	4
Enterocolitis infection	5	1
Injury, Poisoning and Procedural		
Contusion	6	<1
Investigations		

Weight decreased	17	1
Weight increased	11	1
Metabolism and Nutrition Disorders		
Appetite Disturbances	17	1
Musculoskeletal and Connective Tissue Disorders		
Musculoskeletal pain	38	7
Arthralgia	20	2
Myalgia	11	1
Nervous System Disorders		
Headache	37	4
Neuropathy (including peripheral neuropathy)	14	1
Dizziness	9	1
Psychiatric disorder		
Depression	8	0
Insomnia	6	0
Anxiety	6	1
Renal and urinary disorders		
Renal failure	6	5
Respiratory, Thoracic and Mediastinal Disorders		
Cough	29	0
Pleural Effusion	28	8
Dyspnea	28	6
Lung infiltration	5	2
Pulmonary oedema[a]	2	1
Pulmonary hypertension[a]	1	1
Skin and Subcutaneous Tissue Disorders		
Skin Rash	27	1
Pruritus	10	0
Hyperhidrosis	9	0
Dry skin	6	0

Vascular Disorders		
Hemorrhage	44	13
Gastrointestinal bleeding	17	9
CNS bleeding[a]	5	1
Hypertension	8	1
Hypotension	6	2

[a] Adverse events of special interest with <5% frequency.

[b] Includes ventricular dysfunction, cardiac failure, cardiac failure congestive, cardiomyopathy, congestive cardiomyopathy, diastolic dysfunction, ejection fraction decreased, and ventricular failure.

[c] Superficial edema is a grouped term composed of face edema, other superficial edema, and peripheral edema

8.3 Less Common Clinical Trial Adverse Reactions

The following additional adverse reactions, regardless of relationship to therapy or dosing regimen, were reported in patients in the SPRYCEL clinical studies (n = 2,712) at a frequency of <5%, unless otherwise noted. These reactions are presented by frequency category. Frequent reactions are those occurring in ≥1% of patients, infrequent reactions are those occurring in 0.1% – <1% of patients and rare reactions are those occurring in <0.1% of patients. These events are included based on clinical relevance.

Blood and Lymphatic System Disorders: *Frequent:* myelosuppression (including anemia, neutropenia, thrombocytopenia); *Infrequent:* coagulopathy, lymphadenopathy, lymphopenia; *Rare*: aplasia pure red cell, splenic calcification.

Cardiac Disorders: *Frequent:* angina pectoris, cardiomegaly, myocardial infarction (including fatal outcomes) *Infrequent*: electrocardiogram QT prolonged, pericarditis, ventricular arrhythmia (including ventricular tachycardia), acute coronary syndrome, cor pulmonale myocarditis, electrocardiogram T wave abnormal, troponin increased, cardiac arrest, coronary artery disease; *Rare*: arteriosclerosis coronary artery, restrictive cardiomyopathy, electrocardiogram PR prolongation, pleuropericarditis.

Congenital, Familial and Genetic Disorders: *Rare:* porokeratosis.

Ear and Labyrinth Disorders: *Frequent:* tinnitus, vertigo, hearing loss.

Endocrine Disorders: *Frequent:* hypothyroidism; *Infrequent:* hyperthyroidism, thyroiditis.

Eye Disorders: *Frequent:* conjunctivitis, dry eye, visual disorder; *Infrequent:* visual impairment, lacrimation increased; *Rare:* pterygium, retinal vascular disorder, photophobia.

Gastrointestinal Disorders: *Frequent*: dysphagia, gastroesophageal reflux disease, colitis (including neutropenic colitis), oral soft tissue disorder; *Infrequent*: anal fissure, esophagitis, anal fistula, upper gastrointestinal ulcer, pancreatitis, ileus; *Rare*: protein-losing gastroenteropathy, volvulus, pancreatitis acute.

General Disorders and Administration Site Conditions**:** *Frequent:* malaise, face edema (>5%), other superficial edema; Rare: gait disturbance.

Hepatobiliary Disorders: *Infrequen*t: cholecystitis, cholestasis, hepatitis; *Rare:* acquired dilatation intrahepatic duct.

Immune System Disorders: *Rare:* anaphylactic reaction.

Infections and Infestations: *Rare:* sialoadenitis

Injury, Poisoning and Procedural Complications: *Rare:* epicondylitis

Investigations: *Infrequent:* blood creatine phosphokinase increased, gamma-glutamyltransferase increased; *Rare*: clostridum test positive, coxsackle virus test positive, hepatitis C RNA increased, platelet aggregation abnormal, blood chloride increased.

Metabolism and Nutrition Disorders: *Frequent:* dehydration; *Infrequent:* hypoalbuminemia, diabetes mellitus, tumour lysis syndrome, hypercholesterolemia.

Musculoskeletal and Connective Tissue Disorders: *Frequent:* muscular weakness, musculoskeletal stiffness; *Infrequent:* tendonitis, rhabdomyolysis, muscle inflammation, osteonecrosis; *Rare:* chondrocalcinosis, osteochondrosis, gouty tophus.

Neoplasms Benign, Malignant and Unspecified: *Rare:* oral papilloma.

Nervous System Disorders: *Frequent:* dysgeusia, syncope, amnesia, tremor, convulsion, somnolence; *Infrequent:* cerebrovascular accident, transient ischemic attack, balance disorder, ataxia; *Rare*: VIIth nerve paralysis, cerebellar infarction, dementia, reversible posterior encephalopathy syndrome, optic neuritis, carotid artery stenosis.

Pregnancy, Puerperium and Perinatal Conditions: *Rare:* abortion

Psychiatric Disorders: *Frequent:* confusional state, affect lability; *Infrequent:* libido decreased; *Rare:* hypomania, seasonal affective disorder.

Renal and Urinary Disorders: *Infrequent:* proteinuria, renal impairment; *Rare*: nephrocalcinosis, bladder diverticulum, glomerulonephritis.

Reproductive System and Breast Disorders: *Frequent:* gynecomastia; *Infrequent:* menstrual disorder; *Rare:* orchitis non-infective, vaginal prolapse.

Respiratory, Thoracic, and Mediastinal Disorders: *Frequent:* asthma, lung infiltration, dysphonia, pneumonitis; *Infrequent:* bronchospasm, acute respiratory distress syndrome (including fatal outcomes), pulmonary embolism, oropharyngeal discomfort; *Rare:* pulmonary arterial hypertension, nasal septum deviation, rhinitis hypertrophic, reflux laryngitis, nasal septum performation.

Skin and Subcutaneous Tissue Disorders: *Frequent*: urticaria, skin ulcer, photosensitivity; *Infrequent*: bullous conditions, nail disorder, neutrophilic dermatosis, palmar-plantar erythrodysaesthesia syndrome, panniculitis, hair disorder; *Rare:* asteatosis, leukocytoclastic vasculitis, skin fibrosis.

Vascular Disorders: *Frequent:* thrombophlebitis; *Infrequent:* deep vein thrombosis, thrombosis, atherosclerosis; *Rare*: livedo reticularis, peripheral arterial occlusive disease, arterial occlusive disease, embolism, cerebral arteriosclerosis.

8.4 Abnormal Laboratory Findings: Hematologic, Clinical Chemistry and Other Quantitative Data

Clinical Trial Findings

Myelosuppression was commonly reported in all studies. However, the frequency of Grade 3 or 4 neutropenia, thrombocytopenia, and anemia was higher in patients with advanced phase CML or Ph+ ALL than in chronic phase CML. Most patients continued treatment without further progressive myelosuppression.

Newly diagnosed patients with chronic phase CML

Laboratory abnormalities reported in patients treated with SPRYCEL in the Phase III clinical study in patients with newly diagnosed CML are shown in Table 6. Myelosuppression was less frequently reported in newly diagnosed chronic phase CML, than in chronic phase CML patients with resistance or intolerance to prior imatinib therapy. In SPRYCEL-treated patients who experienced grade 3 or 4 myelosuppression, recovery generally occurred following brief dose interruptions and/or reductions and permanent discontinuation of treatment occurred in 2.3% of patients due to drug-related hematologic toxicities.

Table 6: CTC Grade 3/4 Laboratory Abnormalities in Patients with Newly Diagnosed Chronic Phase CML 60-month follow up

Laboratory Parameters	SPRYCEL (n=258)	Imatinib (n=258)
	Percent (%) of Patients	
Biochemistry Parameters		
Elevated Alkaline phosphatase	1	0
Hyperuricemia	4	1
Hypophosphatemia	7	31
Hypokalemia	0	3
Hypocalcemia	4	3
Hypomagnesemia	<1	2
Hyponatremia	3	2
Elevated SGPT (ALT)	<1	2
Elevated SGOT (AST)	<1	1
Elevated Bilirubin	1	0
Elevated Creatinine	1	1
Hematology Parameters		
Neutropenia	29	24
Thrombocytopenia	22	14
Anemia	13	9

CTC grades: neutropenia (Grade 3 ≥0.5–<1.0 × 10^9/L, Grade 4 <0.5 × 10^9/L); thrombocytopenia (Grade 3 ≥25–<50 × 10^9/L, Grade 4 <25 × 10^9/L); anemia (hemoglobin Grade 3 ≥65–<80 g/L, Grade 4 <65 g/L); elevated creatinine (Grade 3 >3–6 × upper limit of normal range (ULN), Grade 4 >6 × ULN); elevated bilirubin (Grade 3 >3–10 × ULN, Grade 4 >10 × ULN); elevated SGOT or SGPT (Grade 3 >5–20 × ULN, Grade 4 >20 × ULN); hypocalcemia (Grade 3 <7.0–6.0 mg/dL, Grade 4 <6.0 mg/dL); hypophosphatemia (Grade 3 <2.0–1.0 mg/dL, Grade 4 <1.0 mg/dL); hypokalemia (Grade 3 <3.0–2.5 mmol/L, Grade 4 <2.5 mmol/L).

Patients with imatinib intolerant or resistant CML or Ph+ ALL

Laboratory abnormalities that were reported in patients treated with SPRYCEL in clinical studies are shown in Table 7 for imatinib intolerant or resistant chronic or advanced phase CML and Ph+ ALL.

In patients who experienced severe myelosuppression, recovery generally occurred following brief dose interruptions and/or reductions. Occasionally permanent discontinuation of treatment was required.

Elevations of transaminases or bilirubin were reported in all disease phases, but were more common in patients with advanced disease. The numbers of patients who developed three or more simultaneous significant elevations of transaminases or bilirubin suggestive of hepatic toxicity were as follows: Chronic phase, 4; accelerated, 13; myeloid blast, 13; lymphoid blast, 7. Most events were managed with dose reduction or interruption. One patient required discontinuation of treatment due to abnormalities of liver function tests. Although causality has not been established, the occurrence of abnormal liver function tests on treatment should be followed closely and consideration given to discontinuing SPRYCEL.

Hypocalcemia:
Between 48% and 76% of patients experienced hypocalcemia at least once during this period. Grade 3 or 4 abnormalities were reported in 2, 7, 16, 13 and 9% of the patients in the chronic phase CML (n=1150), accelerated phase CML (n=502), myeloid blast phase CML (n=280), lymphoid blast phase CML (n=115) and Ph+ ALL (n=135), respectively. The percentage of patients with hypocalcemia who were treated with calcium supplements is 7% for chronic phase CML, 16% for accelerated phase CML, 28% for myeloid blast CML, 20% for lymphoid blast CML and 20% for Ph+ ALL.

Hypophosphatemia:
Between 41% and 50% of patients experienced hypophosphatemia at least once during this period. Grade 3 or 4 abnormalities were reported in 10, 13, 20, 19 and 21% of the patients in the chronic phase CML (n=1150), accelerated phase CML (n=502), myeloid blast phase CML (n=280), lymphoid blast phase CML (n=115) and Ph+ ALL (n=135), respectively.

In the Phase II randomized study, the frequency of Grade 3 or 4 neutropenia, thrombocytopenia, and anemia was 63%, 57%, and 20%, respectively, in the SPRYCEL group and 39%, 14%, and 8%, respectively, in the imatinib group. The frequency of Grade 3 or 4 hypocalcemia was 5% in the SPRYCEL group and 0% in the imatinib group.

Table 7: CTC Grades 3/4 Laboratory Abnormalities in Clinical Studies of CML: Patients with imatinib Resistant or Intolerant chronic phase CML, advanced phase CML or Ph+ ALL [a]

Laboratory Parameters	Chronic Phase[b] n=165	Accelerated Phase[c] n=157	Myeloid Blast Phase[c] n=74	Lymphoid Blast Phase[c] n=33	Ph+ ALL[c] n=40
Biochemistry Parameters					
Hypophosphatemia	10	13	12	18	16
Hypokalemia	2	7	11	15	8
Hypocalcemia	<1	4	9	12	5
Elevated SGPT (ALT)	0	2	5	3	8

Elevated SGOT (AST)	<1	0	4	3	3
Elevated Bilirubin	<1	1	3	6	3
Elevated Creatinine	0	2	8	0	0
Hematology Parameters*					
Neutropenia	35	58	77	79	67
Thrombocytopenia	23	63	78	85	72
Anemia	13	47	74	52	36

[a] Phase III dose optimization study results reported at 2-year study follow up
[b] CA180-034 study results at recommended starting dose of 100 mg once daily
[c] CA180-035 study results at recommended starting dose of 140 mg once daily

8.5 Post-Market Adverse Reactions

The following additional adverse reactions have been identified during post approval use of SPRYCEL. Because these reactions are reported voluntarily from a population of uncertain size, it is not always possible to reliably estimate their frequency or establish a causal relationship to drug exposure.

Cardiac disorders:	Atrial fibrillation/atrial flutter[a]
Infections and infestations	hepatitis B reactivation
Pregnancy disorders:	Fetal complications (including hydrops fetalis and fetal malformations)
Renal and urinary disorders:	Nephrotic syndrome
Respiratory, thoracic and mediastinal disorders:	Interstitial lung disease, pulmonary arterial hypertension[b]
Skin and subcutaneous tissue disorders:	Stevens-Johnson syndrome[c]
Vascular disorders	Thrombotic microangiopathy (TMA)

a. Typically reported in elderly patients or in patients with confounding factors including significant underlying or concurrent cardiac or cardiovascular disorders, or other significant comorbidities (eg, severe infection/sepsis, electrolyte abnormalities).
b. Some patients with PAH reported during SPRYCEL treatment were taking concomitant medications or had co-morbidities in addition to the underlying malignancy.
c. In the post-marketing setting, individual cases of Stevens-Johnson syndrome have been reported. It could not be determined whether these mucocutaneous adverse reactions were directly related to SPRYCEL or to concomitant medications.

9 DRUG INTERACTIONS

9.2 Drug Interactions Overview

Dasatinib is an inhibitor of CYP3A4 and may decrease the metabolic clearance of drugs that are primarily metabolized by CYP3A4. At clinically relevant concentrations, dasatinib does not inhibit CYP 1A2, 2A6, 2B6, 2C8, 2C9, 2C19, 2D6, or 2E1. Dasatinib is not an inducer of CYP enzymes.

9.4 Drug-Drug Interactions

The drugs listed in this table are based on clinical trials, due to the expected magnitude and seriousness of the interaction.

Table 8 - Established or Potential Drug-Drug Interactions

[Proper/Common name]	Source of Evidence	Effect	Clinical comment
Drugs that may increase dasatinib plasma concentrations			
CYP3A4 inhibitors (e.g., ketoconazole)	CT	In a study with solid tumors, 20-mg SPRYCEL once daily coadministered with 200 mg of ketoconazole BID (n=18) increased the dasatinib Cmax and AUC by four- and five-fold, respectively.	Substances that inhibit CYP3A4 activity (eg, ketoconazole, itraconazole, erythromycin, clarithromycin, ritonavir, atazanavir, lopinavir, grapefruit juice) may decrease metabolism and increase concentrations of dasatinib and should be avoided. In patients receiving SPRYCEL, coadministration of a potent CYP3A4 inhibitor is not recommended. Selection of an alternate concomitant medication with no or minimal CYP3A4 inhibition potential is recommended. If systemic administration of a potent CYP3A4 inhibitor cannot be avoided, a dose reduction to 20 or 40 mg daily should be considered and the patient should be closely monitored for toxicity(see 9.5 Drug-Food Interactions and 4.1 Dosing Considerations, Dose reduction for concomitant use of strong CYP3A4 inhibitors)

Drugs that may decrease dasatinib plasma concentrations			
CYP3A4 inducers (e.g. rifampicin)	CT	In healthy subjects (n=20), a single morning dose of SPRYCEL was administered following 8 days of continuous evening administration of 600 mg of rifampicin, a potent CYP3A4 inducer, the mean Cmax and AUC of dasatinib were decreased by 81% and 82%, respectively. In addition, more healthy male subjects experienced increases in QTcF of > 30msec from the baseline recordings when a single dose of dasatinib was administered 12 hours following rifampicin compared to when dasatinib was given alone (25% vs. 10%, n = 20). No subject experienced QTcF > 450 msec or a change from baseline > 60 msec. (see 7 WARNINGS AND PRECAUTIONS,Cardiovascular, and 16 NON-CLINICAL TOXICOLOGY, Safety Pharmacology)	Concomitant use of dasatinib and medicinal products that induce CYP3A4 (e.g. dexamethasone, phenytoin, carbamazepine, rifampicin, phenobarbital or Hypericum perforatum, also known as St. John's Wort) may substantially reduce exposure to dasatinib, potentially increasing the risk of therapeutic failure. Concomitant use of potent CYP3A4 inducers with dasatinib is not recommended. In patients in whom rifampicin or other CYP3A4 inducers are indicated, alternative agents with less enzyme induction potential should be used.

Antacids	CT	In healthy subjects (n=24), 30 mL of aluminum hydroxide/magnesium hydroxide was administered concomitantly with a 50 mg dose of SPRYCEL, a 55% reduction in dasatinib AUC and a 58% reduction in Cmax were observed However, no relevant change in dasatinib AUC or Cmax was observed following administration of 30 mL of aluminum hydroxide/magnesium hydroxide 2 hours prior to a single 50 mg dose of SPRYCEL	Concomitant use of dasatinib and aluminum hydroxide/magnesium hydroxide may reduce exposure to dasatinib. However, **aluminum hydroxide/magnesium hydroxide products may be administered up to 2 hours prior to, or 2 hours following the administration of dasatinib**
H2 antagonists or proton pump inhibitors:			
famotidine	CT	In healthy subjects (n=24), administration of a single 50 mg dose of SPRYCEL 10 hours following famotidine reduced the AUC and Cmax of SPRYCEL by 61% and 63%, respectively	Long-term suppression of gastric acid secretion by H2 antagonists or proton pump inhibitors (e.g. cimetidine, ranitidine, famotidine and omeprazole) is likely to reduce dasatinib exposure (see 9 DRUG INTERACTIONS). **The use of antacids should be considered in place of H2 antagonists or proton pump inhibitors in patients receiving SPRYCEL therapy.**

Drugs that may have their plasma concentration altered by dasatinib			
CYP3A4 Substrates (e.g. simvastatin)	CT	Single dose data from a study of 54 healthy subjects indicate that the mean Cmax and AUC of simvastatin, a prototypical CYP3A4 substrate, were increased by 37% and 20%, respectively, when simvastatin (80 mg) was administered in combination with a single 100 mg dose of SPRYCEL. In addition, three healthy subjects (n = 48) experienced QTcF of > 30 msec from the baseline ECG recordings following the concomitant use of a single dose of simvastatin and dasatinib. No subject experienced QTcF > 450 msec or a change from baseline > 60 msec. (See 7 WARNINGS AND PRECAUTIONS, Cardiovascular).	Caution is warranted when SPRYCEL is co-administered with a drug that potentially alters CYP3A4 activity, a QTc prolonger, or CYP3A4 substrates of narrow therapeutic index such as cyclosporine, macrolide antibiotics, benzodiazepine, pimozide, or ergot alkaloids (ergotamine, dihydroergotamine). The effect of a CYP3A4 substrate on the pharmacokinetic parameters of dasatinib has not been studied.

Legend: C = Case Study; CT = Clinical Trial; T = Theoretical

Drugs that prolong QTc interval or induce torsades de pointes

The concomitant use of SPRYCEL with medicinal products known to prolong QTc interval or medicinal products able to induce torsades de pointes should be avoided if possible. Medicinal products that are generally accepted to carry the risk of QT prolongation and torsades de pointes include but are not limited to the examples that follow: Class IA (e.g. disopyramide, procainamide), Class III (e.g. amiodarone, sotalol, ibutilide), or Class IC (e.g. flecainide), antiarrhythmic medicinal products, antipsychotics (e.g. chlorpromazine, haloperidol, pimozide), opioids (e.g. methadone), macrolide antibiotics (e.g. erythromycin, clarithromycin, quinolone antibiotics (e.g. moxifloxacin), antimalarials (e.g. chloroquine), GI stimulants or others (e.g. domperidone).

Antiemetics

No information is available on the safety of concomitant use of dasatinib with antiemetics (prochlorperazine, metochlopramide, 5-HT3 inhibitors).

9.5 Drug-Food Interactions

SPRYCEL should not be taken with grapefruit or grapefruit juice.

Data from a study of 54 healthy subjects administered a single, 100-mg dose of dasatinib 30 minutes following consumption of a high-fat meal indicated a 14% increase in the mean AUC of dasatinib. Consumption of a low-fat meal 30 minutes prior to dasatinib resulted in a 21% increase in the mean AUC of dasatinib. The observed food effects do not represent clinically relevant changes in exposure.

9.6 Drug-Herb Interactions

Concomitant use of dasatinib and St John's Wort *(Hypericum perforatum)* may substantially reduce exposure to dasatinib.

9.7 Drug-Laboratory Test Interactions

Interactions with laboratory tests have not been established.

10 CLINICAL PHARMACOLOGY

10.1 Mechanism of Action

Dasatinib inhibits the activity of the BCR-ABL kinase and SRC family kinases (LYN, HCK), along with a number of other kinases including c-KIT, ephrin (EPH) receptor kinases, and PDGFβ receptor. Dasatinib is a potent inhibitor of the BCR-ABL and SRC family kinases with potency at sub-nanomolar concentrations. It binds not only to the inactive but also to the active conformation of the enzyme.

10.2 Pharmacodynamics

Nonclinical pharmacodynamics

Extensive *in vitro* and *in vivo* studies demonstrated that dasatinib is a potent inhibitor of BCR-ABL and SRC family kinases along with a number of other kinases including c-KIT, ephrin (EPH) receptor kinases, and PDGFβ receptor. Dasatinib is active *in vitro* and *in vivo* in numerous nonclinical models of CML representing variants of both imatinib-sensitive and -

resistant diseases. Nonclinical studies show that dasatinib can overcome the imatinib resistance that results from divergent mechanisms including BCR-ABL kinase domain mutations, BCR-ABL overexpression, activation of alternate signaling pathways involving the SRC family kinases, and multidrug resistance gene overexpression.

Nonclinical studies demonstrate that dasatinib is capable of binding to the active conformation of BCR-ABL kinase domains, and is predicted to bind to the inactive form. Dasatinib is 300- to 1000-fold more potent than imatinib in killing human CML cells that harbor wild-type or mutant BCR-ABL *in vitro*. In a murine model of CML, dasatinib prevents the progression of chronic CML to blast phase. *In vivo*, dasatinib inhibits the growth and prolonged the survival of mice bearing xenografts of imatinib-sensitive (including an intracranial model) and one imatinib-resistant CML cell line.

In vitro, dasatinib is active in leukemic cell lines representing variants of imatinib sensitive and resistant disease. These nonclinical studies show that dasatinib can overcome imatinib resistance resulting from BCR-ABL overexpression, BCR-ABL kinase domain mutations (14/15 mutations with exception of T315I), activation of alternate signaling pathways involving the SRC family kinases (LYN, HCK), and multidrug resistance gene, *MDR1,* overexpression.

In vivo, in separate experiments using murine models of CML, dasatinib prevented the progression of chronic CML to blast phase and prolonged the survival of mice bearing patient-derived CML cell lines.

Electrocardiogram

In five Phase II clinical studies in patients with leukemia, repeated baseline and on-treatment ECGs were obtained at pre-specified time points and read centrally for 865 patients receiving SPRYCEL 70 mg BID. QT interval was corrected for heart rate by Fridericia's method. At all post-dose time points on day 8, the mean changes from baseline in QTcF interval were 4-6 msec, with associated upper 95% confidence intervals <7 msec. Of the 2182 patients who received SPRYCEL in clinical trials, 21 patients (<1%) experienced a QTcF >500 msec.
(See 7 WARNINGS AND PRECAUTIONS, Cardiovascular)

10.3 Pharmacokinetics

Nonclinical pharmacokinetics

The absorption, distribution, metabolism and excretion properties of dasatinib were evaluated in a series of *in vitro* and *in vivo* studies in mice, rats, rabbits, dogs and monkeys. Dasatinib had a good intrinsic membrane permeability *in vitro* and was rapidly absorbed following oral administration in all species and humans.

In rats and monkeys, systemic exposure was dose related with no apparent gender differences. No notable accumulation was observed after once-daily repeated dosing. After oral administration of [14C] dasatinib to rats, monkeys, and humans, drug-derived radioactivity was recovered primarily in the feces (>76%), with only a small portion of the dose (<7%) excreted in the urine. In all species tested, dasatinib was shown to undergo extensive metabolism, including hydroxylation, N-oxidation, N-dealkylation, oxidation to form a carboxylic acid, glucuronidation and sulfation. Dasatinib was the most abundant drug-related component in the plasma from these species, with multiple oxidative and conjugated metabolites also present. All metabolites identified in human plasma were also found in monkey plasma. The ADME profiles

of dasatinib in mice, rats, rabbits, dogs and monkeys as compared to humans suggest that these species were appropriate for safety assessment of dasatinib and its metabolites.

Multiple enzymes were involved in the metabolism of dasatinib with CYP3A4 playing a major role. The involvement of CYP3A4 was confirmed in clinical studies where the exposure of dasatinib was substantially decreased (> 80%) when it was administered 12 hours following 7-day treatment with rifampin, a potent inducer of CYP3A4. *In vitro* studies indicated that dasatinib was not an inducer of CYP enzymes. It inhibited CYP2C8 in a competitive manner and CYP3A4 in a time dependent manner. Based on the Cmax of dasatinib at the therapeutic dose, the probability of drug-drug interactions is low with co-administered drugs that are CYP2C8 substrates. However, there is a possibility of interaction with drugs that are CYP3A4 substrates given that clinical study with co-administration of dasatinib with simvastatin resulted in a moderate increase in the exposure of simvastatin and its acid.

Clinical pharmacokinetics

The pharmacokinetics of SPRYCEL (dasatinib) were also evaluated in 229 healthy subjects and in 84 patients with leukemia.

Absorption

Dasatinib is rapidly absorbed in patients following oral administration. Peak concentrations were observed between 0.25-6 hours. The overall mean terminal half-life of dasatinib is approximately 3 to 5 hours

Distribution

In patients, SPRYCEL has a large apparent volume of distribution (2505 L) suggesting that the drug is extensively distributed in the extravascular space.

Metabolism

Dasatinib is extensively metabolized in humans. In a study of 8 healthy subjects administered 100 mg of [^{14}C]-labeled dasatinib, unchanged dasatinib represented 29% of circulating radioactivity in plasma. Plasma concentration and measured *in vitro* activity indicate that metabolites of dasatinib are unlikely to play a major role in the observed pharmacology of the drug. CYP3A4 is a major enzyme responsible for the metabolism of dasatinib.

Elimination

Elimination is predominantly in the feces, mostly as metabolites. Following a single oral dose of [^{14}C]-labeled dasatinib, approximately 89% of the dose was eliminated within 10 days, with 4% and 85% of the administered radioactivity recovered in the urine and feces, respectively. Unchanged dasatinib accounted for 0.1% and 19% of the administered dose in urine and feces, respectively, with the remainder of the dose being metabolites.

Special Populations and Conditions

- **Pediatrics:** No clinical studies were conducted with SPRYCEL in pediatric populations. Health Canada has not authorized an indication for pediatric use.
- **Hepatic Insufficiency** The effect of hepatic impairment on the single-dose pharmacokinetics of dasatinib was assessed in 8 moderately hepatic impaired subjects who received a 50-mg dose and 5 severely hepatic-impaired subjects who received a 20-mg dose compared to matched healthy subjects who received a 70-mg dose of SPRYCEL. The mean Cmax and AUC of dasatinib adjusted for the 70-mg dose was decreased by 47% and 8%, respectively, in moderate hepatic impairment

compared to subjects with normal hepatic function. In severe hepatic impaired subjects, the mean Cmax and AUC adjusted for the 70-mg dose was decreased by 43% and 28%, respectively, compared to subjects with normal hepatic function. Hepatic impairment did not result in clinically meaningful change in dasatinib exposure at the doses studied. However no pharmacokinetic information is available from patients with hepatic impairment treated with a 70-100 mg dose of SPRYCEL. Due to limitations of this clinical study, caution is recommended in patients with hepatic impairment (See 7 WARNINGS AND PRECAUTIONS, Hepatic/Biliary/Pancreatic and 4.2 Recommended Dose and Dosage Adjustment, Hepatic Impairment).

- **Renal Insufficiency** No clinical studies were conducted with SPRYCEL in patients with decreased renal function. Less than 4% of SPRYCEL and its metabolites are excreted via the kidney. (See 7 WARNINGS AND PRECAUTIONS, Renal Impairment)

11 STORAGE, STABILITY AND DISPOSAL

SPRYCEL (dasatinib) tablets should be stored at room temperature between 15°–30° C. Keep out of reach and sight of children.

12 SPECIAL HANDLING INSTRUCTIONS

Procedures for proper handling and disposal of anticancer drugs should be considered. Several guidelines on this subject have been published. There is no general agreement that all of the procedures recommended in the guidelines are necessary or appropriate.

SPRYCEL (dasatinib) tablets consist of a core tablet (containing the active drug substance), surrounded by a film coating to prevent exposure of pharmacy and clinical personnel to the active drug substance. However, if tablets are crushed or broken, pharmacy and clinical personnel should wear disposable chemotherapy gloves. Personnel who are pregnant should avoid exposure to crushed and/or broken tablets.

PART II: SCIENTIFIC INFORMATION

13 PHARMACEUTICAL INFORMATION

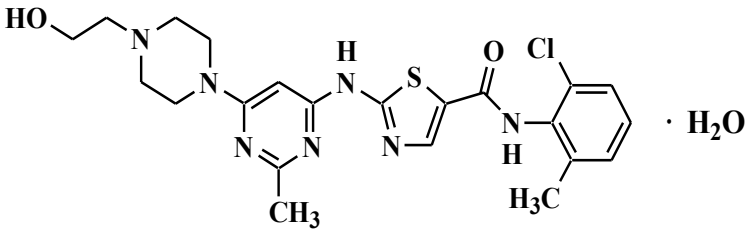
Drug Substance

Proper name: dasatinib

Chemical name: *N*-(2-chloro-6-methylphenyl)-2-[[6-[4-(2-hydroxyethyl)-1-piperazinyl]-2-methyl-4-pyrimidinyl]amino]-5-thiazolecarboxamide, monohydrate

Molecular formula and molecular mass: $C_{22}H_{26}ClN_7O_2S \bullet H_2O$; 488.01 (anhydrous free base)

Structural formula:



Physicochemical properties: Dasatinib is a white to off-white powder, which may contain lumps, and has a melting point of 280°–286° C. The drug substance is insoluble in water (0.008 mg/mL) at 24 ± 4° C. The pH of a saturated solution of dasatinib in water is about 6.0. Two basic ionization constants (pKa) were determined to be 6.8 and 3.1, and one weakly-acidic pKa was determined to be 10.9. The solubilities of dasatinib in various solvents at 24 ± 4°C are as follows: slightly soluble in ethanol (USP), methanol, polyethylene glycol 400, and propylene glycol; very slightly soluble in acetone and acetonitrile; and practically insoluble in corn oil. [text]

14 CLINICAL TRIALS

14.1 Trial Design and Study Demographics

Newly Diagnosed Chronic Phase CML

Table 9 - Summary of patient demographics for clinical trials in Newly Diagnosed Chronic Phase CML

Study #	Study design	Dosage, route of administration and duration	Study subjects (n)	Mean age (Range)	Sex
CA180-056	Open-label, multicenter	SPRYCEL 100 mg oral QD	259	46 (18-64)	44% women
		Imatinib oral 400 mg QD	260	47 (18-78)	37% women

An open-label, multicenter, international (Europe, South America and Asia-Pacific regions), randomized, Phase III study was conducted in adult patients with newly diagnosed chronic phase CML. Patients were randomized to receive either SPRYCEL 100 mg once daily or imatinib 400 mg once daily. The primary endpoint was the rate of confirmed complete cytogenetic response (cCCyR) within 12 months. Secondary endpoints included time-in cCCyR (measure of durability of response), time-to cCCyR, major molecular response (MMR) rate, time-to MMR, progression free survival (PFS), and overall survival (OS). The secondary endpoints were evaluated on a yearly basis. A pre-specified statistical comparison of these endpoints was conducted with data from up to 60 months of follow-up.

A total of 519 patients were randomized to a treatment group: 259 to SPRYCEL and 260 to imatinib. Baseline characteristics were well balanced between the two treatment groups with respect to age (mean age was 46 years for the SPRYCEL group and 47 years for the imatinib group with 10% and 11% of patients 65 years of age or older, respectively), gender (women 44% and 37%, respectively), and race (Caucasian 51% and 55%; Asian 42% and 37%, respectively). At baseline, the distribution of Hasford Scores was similar in the SPRYCEL and imatinib treatment groups (low risk: 33% and 34%; intermediate risk: 48% and 47%; high risk: 19% and 19%, respectively). The ECOG Performance Score was also similar in the SPRYCEL and imatinib treatment groups (ECOG 0 = 82% and 79%; ECOG 1 = 18% and 20%; and ECOG 2 = 0 and 1%, respectively). [**SNDS Control No. 140047; Mod. 2.7.6**]

With a minimum of 12 months follow-up, 84% of patients randomized to the SPRYCEL group and 81% of patients randomized to the imatinib group were still receiving first-line treatment. Discontinuation due to disease progression occurred in 3% of SPRYCEL-treated patients and 5% of imatinib-treated patients. With a minimum of 36 months follow-up, 71% of patients randomized to the SPRYCEL group and 69% of patients randomized to the imatinib group were still receiving first-line treatment. With a minimum of 60 months follow-up, 61% of patients randomized to the SPRYCEL group and 63% of patients randomized to the imatinib group were still receiving first-line treatment. Discontinuation due to disease progression occurred in 7% of SPRYCEL-treated patients and 8.5% of imatinib-treated patients.

Imatinib Resistant or Intolerant CML or Ph+ ALL

Table 10 - Summary of patient demographics for clinical trials in CML, resistant or intolerant to imatinib

Study #	Study design	Dosage, route of administration and duration	Study subjects (n)	Mean age (Range)	Sex
CA180-034	Randomized open-label, dose-optimisation (chronic phase CML patients)	SPRYCEL: 100 mg QD 140mg QD 50 mg BID or 70 mg BID; oral	QD administration: 334	54 (20 - 84)	54% women
			BID administration 336	54 (18 - 84)	52% women

CA180-035	Randomized open-label, dose-optimisation (advanced CML and Ph+ ALL patients)	SPRYCEL: 70 mg BID or 140mg QD; oral	70 mg: 305	53 (15 - 84)	44% women
			140 mg 306	52 (16 - 81)	44% women

Phase III dose-optimization study in chronic phase CML (Study CA180-034): A randomized, open-label study was conducted in patients with chronic phase CML to evaluate the efficacy of SPRYCEL administered once daily compared with SPRYCEL administered twice daily. The primary endpoint was MCyR in imatinib-resistant patients. The main secondary endpoint was MCyR by total daily dose level in the imatinib-resistant patients at 24-months follow-up. Other secondary endpoints included duration of MCyR and overall survival. A total of 670 patients, of whom 497 were imatinib resistant, were randomized to the SPRYCEL 100 mg once daily, 140 mg once daily, 50 mg twice daily, or 70 mg twice daily group. Median duration of treatment was 22 months.

Resistance to imatinib was defined as failure to achieve a CHR (after 3 months), MCyR (after 6 months), or CCyR (after 12 months); or loss of a previous molecular response (with concurrent ≥10% increase in Ph+ metaphases), cytogenetic response, or hematologic response.

Progression in the chronic phase CML was defined as any of the following events: loss of a CHR or MCyR; no CHR with an increase in white blood cell count; development of accelerated or blast phase CML; a ≥30% increase in the number of Ph+ metaphases; or death.

Phase III dose-optimization study in advanced phase CML and Ph+ ALL (Study CA180-035): A randomized, open-label study was conducted in patients with accelerated phase CML, myeloid blast phase CML, lymphoid blast phase CML, or Ph+ ALL to evaluate the efficacy of SPRYCEL administered once daily compared with SPRYCEL administered twice daily. The primary endpoint was the rate of MaHR. Secondary endpoints included the rate of MCyR, duration of MaHR, PFS, and overall survival. A total of 611 patients were randomized to the SPRYCEL 140 mg once daily or 70 mg twice daily group. Median duration of treatment was 14 months for accelerated phase CML, 3 months for myeloid blast CML, 4 months for lymphoid blast CML, and 3 months for Ph+ ALL.

Resistance to imatinib was defined as no hematologic response or a ≥50% increase in blasts in peripheral blood; loss of a hematologic response; progression to blast or accelerated phase CML with blasts in peripheral blood while on treatment with imatinib.

Progression was defined as follows:

- Accelerated phase CML: Loss of a CHR, NEL, or MiHR; development of blast phase CML; no decrease from baseline percent blasts in peripheral blood or bone marrow; development of extramedullary sites (other than spleen or liver); a ≥50% increase in blasts in peripheral blood; or death.

- Blast phase CML or Ph+ ALL: Loss of a CHR, NEL, or MiHR; no decrease from baseline percent blasts in peripheral blood or bone marrow; a ≥50% increase in blasts in peripheral blood; or death.

14.2 Study Results

Newly Diagnosed Chronic Phase CML

Efficacy results are presented in Table 11. A statistically significantly greater proportion of patients in the SPRYCEL group achieved a cCCyR compared with patients in the imatinib group within the first 12 months of treatment. This result was generally consistent across different subgroups, including age, gender, and baseline Hasford score. No statistically significant difference in the secondary endpoint, time-in cCCyR, was demonstrated between SPRYCEL and imatinib at the 60 month analysis. In accord with the pre-specified sequential testing strategy, formal statistical testing stopped after the treatment comparison for Time-in cCCyR was found to be not statistically significant. Therefore statistical comparisons with remaining secondary endpoints were not conducted.

Table 11 - Efficacy Results in Newly Diagnosed Patients with Chronic Phase CML (Study CA180-056)

Endpoints	SPRYCEL (n=259)	Imatinib (n=260)	p-value
	Response rate (95% CI)		
Cytogenetic Response within 12 months			
cCCyR[a]	76.8% (71.2–81.8)	66.2% (60.1–71.9)	p = 0.007*
within 24 months			
cCCyR[a]	80.3% (74.9-85.0)	74.2% (68.5-79.4)	----**
within 36 months			
cCCyR[a]	82.6% (77.5-87.0)	77.3% (71.7-82.3)	----**
within 60 months			
cCCyR[a]	83.0% (77.9-87.4)	78.5% (73.0-83.3)	----**
Major Molecular Response[b]			
12 months	52.1% (45.9–58.3)	33.8% (28.1–39.9)	p<0.00003*
24 months	64.5% (58.3-70.3)	50% (43.8-56.2)	----**
36 months	69.1% (63.1-74.7)	56.2% (49.9-62.3)	----**
60 months	76.4% (70.8-81.5)	64.2% (58.1-70.1)	----***
	Hazard Ratio (99.99% CI)		
	within 60 months (95% CI)		
Time-in cCCyR	0.79 [0.55, 1.13]		NS
	within 12 months (99.9% CI)		

Time-to cCCyR	1.55 (1.0–2.3)	p<0.0001*
Time-to MMR	2.01 (1.2–3.4)	p<0.0001*
	within 24 months (95% CI)	
Time-to cCCyR	1.49 (1.22–1.82)	___
Time-to MMR	1.69 (1.34–2.12)	___
	within 36 months (95% CI)	
Time-to cCCyR	1.48 (1.22–1.80)	___
Time-to MMR	1.59 (1.28–1.99)	___
	within 60 months (95% CI)	
Time-to cCCyR	1.46 (1.20–1.77)	----***
Time-to MMR	1.54 (1.25–1.89)	----***

[a] Confirmed complete cytogenetic response (cCCyR) is defined as a response noted on two consecutive occasions (at least 28 days apart).

[b] Major molecular response (at any time) was defined as BCR-ABL ratios ≤ 0.1% by RQ-PCR in peripheral blood samples standardized on the International Scale. Some subjects at the time of minimum follow up corresponding to a specific yearly database cutoff had been on treatment longer, and may have achieved an MMR beyond the corresponding 12, 24 or 36 months of treatment.

*Adjusted for Hasford Score and indicated statistical significance at a pre-defined nominal level of significance.
**Per protocol, formal statistical comparison of cCCyR and MMR rates was only performed at the time of the primary endpoint (cCCyR within 12 months).
***Based on hierarchical statistical testing procedure, formal testing was not done on this secondary endpoint since Time-in cCCyR was not significant.
CI = confidence interval.

NS= not statistically significant

Median time to cCCyR was 3.1 (3.0-3.1) months in 215 SPRYCEL responders and 5.8 (5.6-6.0) months in 204 imatinib responders based on 60-month data update. Median time to MMR (based on 60-month data update) was 9.3 months in 198 SPRYCEL responders and 15.0 months in 167 imatinib responders. The rates of cCCyR in the SPRYCEL and imatinib treatment groups, respectively, within 3 months (54% and 30%), 6 months (70% and 56%), 9 months (75% and 63%), 24 months (80% and 74%) and 36 months (83% and 77%), and 60 months (83 % and 79%) were consistent with the primary endpoint.

At 60 months follow-up in the SPRYCEL arm, the rate of MMR at any time in each risk group determined by Hasford score was 90% (low risk), 71% (intermediate risk) and 67% (high risk).

The rate of cCCyR at any time in each risk group determined by Hasford score was 94% (low risk), 77% (intermediate risk) and 78% (high risk).

The estimated progression-free survival rate at 60 months for dasatinib-treated subjects was 88.9% (95% CI = [84.0%, 92.4%]). The estimated overall survival rate at 60 months for dasatinib-treated subjects was 90.9% (95% CI = [86.6%, 93.8%]).

Disease progression (defined as ‘loss of complete hematologic response’, ‘loss of major cytogenetic response’, ‘rising WBC on two occassions at least one month apart’, ‘transformation to accelerated, blast phase of CML’ or ‘death’) was reported in 34 (13.0%)

patients treated with SPRYCEL and 39 (15%) patients with imatinib. Treatment failure (defined according to the 2006 European LeukemiaNet Guidelines, included disease progression, a lack of a hematologic response at 3 months, a lack of a complete hematologic response or CyR at 6 months, a lack of partial CyR at 12 months, or a lack of CCyR at 18 months) occurred in 10 (3.9%) of SPRYCEL-treated patients and 14 (5.4%) of imatinib-treated patients at 60 months. Transformation to accelerated or blast phase was reported in 8 (3.1%) SPRYCEL-treated patients and 15(5.8%) imatinib treated patients. Deaths were reported in 26 (10.1%) patients treated with SPRYCEL and 26 (10.1%) patients treated with imatinib.

BCR-ABL kinase domain sequencing was performed on blood samples from patients at the time of discontinuation or study closure. At 60 months follow-up, T315I, F317I/L, F3171/V299L and V299L mutations were detected in 15 patients who discontinued SPRYCEL treatment including 8 with T315I. Mutations including M244V, L387M, D276G/F359C, H396P/R, G250E, F359C/I/V, E255K, E355G, E255K/V, E355G/L248V, E255V/Y253H, F317L, and E450G were detected in 19 patients who discontinued imatinib. The T315I mutation confers resistance to treatment with dasatinib and other ABL tyrosine kinase inhibitors based on *in vitro* and clinical data.

Imatinib Resistant or Intolerant CML or Ph+ ALL

Phase III dose-optimization study in chronic phase CML (Study CA180-034): Efficacy was achieved across all SPRYCEL treatment groups with the once daily schedule demonstrating comparable efficacy (non-inferiority) to the twice daily schedule on the primary efficacy endpoint in imatinib resistant patients (difference in MCyR 1.9%; 95% confidence interval [-6.8%–10.6%]); however, the 100 mg once daily regimen demonstrated improved efficacy and tolerability. The main secondary endpoint of the study also showed comparable efficacy (non-inferiority) among imatinib-resistant patients between the 100 mg total daily dose and the 140 mg total daily dose (difference in MCyR -0.2%; 95% CI [-8.9%–8.5%]). Two year efficacy results are presented in Table 12.

Table 12 - Results of Efficacy of SPRYCEL in Phase III Dose-Optimization Study: Imatinib-Resistant or Intolerant Chronic Phase CML Patients (2-year results) [a] (Study CA180-034)

All Patients	n = 167
Imatinib-Resistant Patients	n = 124
Haematologic Response Rate[b] (%) (95% CI)	
CHR	92% (86-95)
Cytogenetic Response[c] (%) (95% CI)	
MCyR	
All Patients	63% (56-71)
Imatinib-Resistant Patients	59% (50-68)
CCyR	
All Patients	50% (42-58)
Imatinib-Resistant Patients	44% (35-53)

[a] Results reported in recommended starting dose of 100 mg once daily

[b] Haematologic response criteria (all responses confirmed after 4 weeks):
CHR (chronic CML): WBC ≤ institutional ULN, platelets < 450,000/mm^3, no blasts or promyelocytes in peripheral blood, < 5% myelocytes plus metamyelocytes in peripheral blood, basophils in peripheral blood < 20%, and no extramedullary involvement.

[c] Cytogenetic response criteria: complete (0% Ph+ metaphases) or partial (> 0%-35%). MCyR (0%-35%) combines both complete and partial responses.

A total of 378 out of 670 patients (56%) with chronic phase CML had abnormal blood count at entry; 317 out of the 378 (84%) patients achieved a CHR from an abnormal baseline (high WBC counts becoming normal and maintained for at least 4 weeks without any other concomitant therapy). A total of 554 out of 670 patients (83%) had abnormal cytogenetics at study entry.

Major molecular response (defined as BCR-ABL/control transcripts ≤0.1% by RQ-PCR in peripheral blood samples) was evaluated in a subset of assessed patients who had a CCyR.

Major molecular response was achieved in 72% (95% CI [58-83%] of imatinib-resistant patients in the SPRYCEL 100 mg once daily group.

Subjects on a BID dosing schedule were permitted to switch to a QD dosing schedule after 24 months of treatment. After 24 months of treatment cytogenetic response was not assessed; blood count with differential and molecular response were assessed once a year.

Based on the Kaplan-Meier estimates, the proportion of patients among those who achieved MCyR on 100 mg of SPRYCEL once daily and maintained MCyR for 18 months was 93% (95% CI: [88%-98%].

Based on the Kaplan-Meier estimates, the proportions of patients with PFS at 1 year were 88% (95% CI [82-94%]) of imatinib-resistant patients in the 100 mg once daily group. At 2 years, the estimated rates of PFS were 77% (95% CI [68-85%]) of imatinib-resistant patients in the 100 mg once daily group. At 5 years, the estimated rates of PFS were 49% (95% CI [39-59%]) of imatinib-resistant patients in the 100 mg once daily group. At 7 years, the estimated rates of PFS were 39% (95% CI [29-49%]) of imatinib-resistant patients in the 100 mg once daily group.

The estimated rates of overall survival at 1 year were 94% (95% CI [90-98%]) of imatinib-resistant patients in the 100 mg once daily group. At 2 years, the estimated rates of overall survival were 89% (95% CI [84-95%]) of imatinib-resistant patients in the 100 mg once daily group. At 5 years, the estimated rates of overall survival were 77% (95% CI [69-85%]) of imatinib-resistant patients in the 100 mg once daily group. At 7 years, the estimated rates of overall survival were 63% (95% CI [53-71%]) of imatinib-resistant patients in the 100 mg once daily group.

Efficacy was also assessed in patients who were intolerant to imatinib. In this population of patients who received 100 mg once daily, MCyR was achieved in 77%, CCyR in 67%, and major molecular response in 64%. Based on the Kaplan-Meier estimates, all imatinib-intolerant patients who achieved MCyR (100%) maintained MCyR for 1 year and 92% (95% CI: [80%-100%]) among those who achieved MCyR maintained MCyR for 18 months. The estimated rate of PFS in this population was 97% (95% CI: [92%-100%]) at 1 year, 87% (95% CI: [76%-99%]) at 2 years, 56% (95% CI [37%-76%]) at 5 years, and 50.9% (95% CI: [32.1%-67.0%]) at 7 years. The estimated rate of overall survival was 100% at 1 year, 95% (95% CI: [88%-100%]) at 2 years, 82% (95% CI: [70%-94%]) at 5 years, and 70.0% (95% CI: [52.2%-82.2%]) at 7 years.

Phase III dose-optimization study in advanced phase CML and Ph+ ALL (Study CA180-035):
Results described below are based on a minimum of 24 months follow-up.

The once daily schedule demonstrated comparable efficacy (non-inferiority) to the twice daily schedule on the primary efficacy endpoint (difference in MaHR 0.8%; 95% confidence interval [-7.1% - 8.7%]); however, the 140 mg once daily regimen demonstrated improved safety and tolerability. Response rates for patients in the 140 mg once daily group are presented in Table 13.

Table 13 - Efficacy of SPRYCEL in Phase III Dose-Optimization Study: Advanced Phase CML and Ph+ ALL (2 Year Results)[a] (Study CA180-035)

Endpoint	140 mg Once Daily			
	Accelerated (n=158)	Myeloid Blast (n=75)	Lymphoid Blast (n=33)	Ph+ ALL (n=40)
MaHR[b] (95% CI)	**66% (59-74)**	**28% (18-40)**	**42% (26-61)**	**38% (23-54)**
CHR[b] (95% CI)	**47% (40-56)**	**17% (10-28)**	**21% (9-39)**	**33% (19-49)**
NEL[b] (95% CI)	**19% (13-26)**	**11% (5-20)**	**21% (9-39)**	**5% (1-17)**
MCyR[c] (95% CI)	**39% (31-47)**	**28% (18-40)**	**52% (34-69)**	**70% (54-83)**
CCyR (95% CI)	**32% (25-40)**	**17% (10-28)**	**39% (23-58)**	**50% (34-66)**

[a] Results reported in recommended starting dose of 140 mg once daily.

[b] Hematologic response criteria (all responses confirmed after 4 weeks): Major hematologic response (MaHR) = complete hematologic response (CHR) + no evidence of leukemia (NEL).

CHR: WBC ≤ institutional ULN, ANC ≥1000/mm^3, platelets ≥100,000/mm^3, no blasts or promyelocytes in peripheral blood, bone marrow blasts ≤5%, <5% myelocytes plus metamyelocytes in peripheral blood, basophils in peripheral blood <20%, and no extramedullary involvement.

NEL: same criteria as for CHR but ANC ≥500/mm^3 and <1000/mm^3, or platelets ≥20,000/mm^3 and ≤100,000/mm^3.

[c] MCyR combines both complete (0% Ph+ metaphases) and partial (>0%-35%) responses.

CI = confidence interval

ULN = upper limit of normal range.

A total of 529 out of 611 patients (87%) with advanced phase CML or Ph+ ALL had abnormal blood count at entry; 238 out of the 529 (45%) patients achieved a MaHR from an abnormal baseline (high WBC counts becoming normal and maintained for at least 4 weeks without any other concomitant therapy)

A total of 526 out of 611 patients (86%) had abnormal cytogenetics at study entry.

In patients with accelerated phase CML treated with the 140 mg once daily regimen, the median duration of MaHR and the median overall survival was not reached; the median PFS was 25 months. In patients with myeloid blast phase CML, treated with the 140 mg once daily regimen, the median duration of MaHR was 8 months, the median PFS was 4 months and the median overall survival was 8 months. In patients with lymphoid blast phase CML, the median duration of MaHR was 5 months, the median PFS was 5 months, and the median overall survival was 11 months.

15 MICROBIOLOGY

No microbiological information is required for this drug product.

16 NON-CLINICAL TOXICOLOGY

General Toxicology:

Acute Toxicity

The single-dose oral toxicity of dasatinib was evaluated in rats at doses of 30, 100, and 300 mg/kg, and in monkeys at doses of 15, 25, and 45 mg/kg. In rats, dasatinib at 30 mg/kg was tolerated, and doses $\geq$ 100 mg/kg caused severe toxicity and death. Morbidity and mortality were attributed to gastrointestinal lesions resulting in fluid and electrolyte loss and impairment of mucosal integrity, bone-marrow and lymphoid depletion, and multifocal myocardial necrosis and hemorrhage. In monkeys, dasatinib was tolerated at doses up to 25 mg/kg, whereas a dose of 45 mg/kg resulted in severe toxicity and mortality at Days 1 and 2. Principal drug-related toxicities occurred in the skin (hemorrhage) at doses $\geq$ 15 mg/kg, GI and lymphoid-organ systems at doses $\geq$ 25 mg/kg, and kidney at 45 mg/kg.

Species/ Strain	Route of Administration	Duration of Dosing	Dose (mg/kg)	N/Dose/Sex	Findings
Rat / SD	Oral gavage	Single dose	30, 100, 300	10 M 10 F	≥ 30 mg/kg: Dose-related decreased food intake, mucous feces, soiled/rough haircoat, dehydration, chromodacryorrhea, and chromorhinorrhea. Decreased size and weight of the thymus, decreased spleen weights (M), increased liver weights (F), red discoloration, ulceration, hemorrhage, and/or edema in the stomach, bone marrow depletion, and lymphoid depletion in the thymus, spleen, and/or lymph nodes. Decreases in total leukocyte, lymphocyte, monocyte, and platelet counts; increases in fibrinogen, ALT and AST, and decreases in albumin, total protein, albumin/globulin ratio, ALP, potassium, calcium and phosphorus. ≥ 100 mg/kg: Mortality (55% at 100 mg/kg by Day 4, 100% at 300 mg/kg by Day 3). Prior to death, decreased activity, hunched posture, pallor, surface hypothermia, ptosis, tremors (F), and absence of feces (F). Hemorrhage and/or coagulative necrosis, macrophage infiltration, hemosiderosis, and fibrosis in the heart, Red/black discoloration of the intestines and lymph nodes, red discoloration of the ovaries, tan discoloration of the liver, and decreased size of the spleen. Enteropathy in the small intestine, hemorrhage or ulceration in the small intestine (F at 300 mg/kg), renal tubular dilatation and epithelial vacuolation, increases in urinary blood and bilirubin (M), lymphoid depletion in intestinal lymphoid nodules, single-cell necrosis in the liver (F), hemorrhage in the epididymides, and testicular degeneration.

Monkey / Cynomolgus	Oral gavage	Single dose	15, 25, 45	2 M 2 F	≥ 15 mg/kg: Decreased activity, surface hypothermia with decreased body temperature, dehydration, and hemorrhages at multiple sites (thorax, limbs, gingiva, head, neck and, in 1 monkey, retina). Increases in AST, decreases in total protein, globulins, and albumin, and increases or decreases in phosphorus. ≥ 25 mg/kg: Fecal changes (soft, liquid, bloody), pallor of mucous membranes, and decreased body weights and food intake. Lymphoid depletion in the spleen, lymph nodes, and lymphoid nodules of the stomach and intestines, and, in 1 monkey, edema in the stomach. Increases in ALT and urea nitrogen, and decreases in calcium, cholesterol, triglycerides, and γ-GT. 45 mg/kg: Mortality (100% by Days 1 or 2). Prior to death, emesis and increased muscle tone and tremors. Red or abnormal contents of the intestines (F), hemorrhage in the tongue, red discoloration and hemorrhage in the stomach and intestines, dilatation of cortical tubules of the kidney (F), increases in creatinine and potassium (F).

Short- and Long-Term Toxicity

Repeat-dose oral toxicity studies were conducted in rats for 2 weeks to 6 months, and in monkeys for 10 days to 9 months. Repeat-dose oral toxicity studies were conducted using a daily dosing regimen (2-week and 6-month studies in rats) or a 5-days on, 2-days off dosing schedule (1-month study in rats, and 10-day, 1-month, and 9-month studies in monkeys) to support a flexible clinical development plan. In both rats and monkeys, the principal drug-related toxicities were manifested in the GI and lymphoid-organ systems. Hematopoietic (bone marrow) toxicity was also a consistent finding in rats following single or repeated oral doses of dasatinib, and was accompanied by decreases in erythrocyte, lymphocyte, and platelet counts. In monkeys, minimal bone marrow toxicity occurred only in a small number of animals following repeat dosing, and was generally accompanied by decreases in erythrocyte and lymphocyte counts. In a 9-month monkey study, toxicity related to gastroenteropathy, lymphocytic depletion and others necessitated euthanasia of 50% of the animals at exposures that were only half of the systemic exposure in humans at a dose of 70 mg BID.

Species/ Strain	Route of Administration	Duration of Dosing	Dose (mg/kg)	N/Dose/Sex	Findings
Rat / SD	Oral gavage	2 weeks (daily dosing)	1, 15, 30	6 M 6 F	1 mg/kg: No drug-related changes. ≥ 15 mg/kg: Chromorhinorrhea, soiled/rough haircoat, dehydration, soft feces, and bloated/swollen abdomen (F at 15 mg/kg). Distention of the GI tract with gas, fluid, and/or ingesta or digesta. Enteropathy of the small and large intestines, edema of the large intestine, red discoloration of the mesenteric lymph nodes, decreased size of the thymus, and lymphoid depletion of the spleen, thymus, and lymph nodes. At 15 mg/kg, changes in erythrocyte parameters (decreases in erythrocyte counts, hemoglobin, and hematocrit, and increases in reticulocyte counts, MCV, and MCH), increased liver (F) and adrenal weights, and decreased kidney (M), thymus, and spleen weights. 30 mg/kg: Mortality (100%). Prior to death, decreased activity, surface hypothermia, pallor, diarrhea, hunched posture, ptosis, thin appearance, decreased body weight gain (F), body weight loss (M), and decreased food intake. Red discoloration of the small intestine (M), lymphoid depletion in the spleen and thymus, and bone-marrow haematopoietic depletion.

Rat / SD	Oral gavage	1 month (5-days on, 2-days off)	0.9, 15, 25	15 M 15 F	≥ 0.9 mg/kg: Decreased food consumption (M). ≥ 15 mg/kg: Changes in erythrocyte parameters (decreases in erythrocyte counts, hemoglobin, and hematocrit, and increases in MCV and MCH). Decreased body-weight gain (M) and spleen weights, and increases in liver weights (F). Enteropathy in the gastrointestinal track. Lymphoid depletion, edema, and/or hemorrhage in the thymus. 25 mg/kg: Mortality (43%) due to enteropathy/lymphoid depletion. Distention and red discoloration of the gastrointestinal tract, hemorrhage in the stomach, edema in the cecum, red discoloration of the mesenteric lymph node, lymphoid depletion in the spleen, and hypocellularity in the bone marrow accompanied with hematological changes.

Rat / SD	Oral gavage	6 months (daily dosing)	1.5, 4, 15/10/8	25 M 25 F	The high dose of 15 mg/kg was reduced to 10 mg/kg in Week 8 and then to 8 mg/kg in Week 17 due to gastrointestinal toxicity. ≥ 1.5 mg/kg/day: Increased heart weights. Gastrointestinal changes of villous blunting/fusion/branching and/or epithelial hyperplasia, increased vacuolation in the adrenal cortex, increased corpora lutea in the ovary and decreased incidence of acyclic ovaries, fluid-filled uteri and decreased squamous metaplasia of endometrial glands in the uterus. ≥ 4 mg/kg/day: The systemic exposure of dasatinib at 4 mg/kg was similar to that of humans at the therapeutic dose. Increased weights of ovaries, liver, adrenal glands, and thyroid/ parathyroid glands, and decreased weights of the pituitary gland. Fibrosis and crypt ectasia/abscesses in the cecum, and increased colloid in the thyroid. 15/10/8 mg/kg: Mortality (30%) at systemic exposure of dasatinib 2-4x that of humans at the therapeutic dose. In surviving animals, swollen abdomen, few or liquid feces, and fecal stained haircoat. Reversible bone marrow hypocellularity (minimal or moderate, 2 rats) or individual cell necrosis (minimal, 1 rat), changes in erythrocyte parameters (decreased erythrocyte counts, hemoglobin, and hematocrit, and increased MCV, MCH, and reticulocyte counts), and platelet parameters (increased platelet counts and decreases in platelet aggregation), increased neutrophil counts and fibrinogen, and decreased serum proteins (total protein, albumin, and globulins).

Dog / Beagle	Oral gavage	2 days	5	1 M 1 F	Dosing was discontinued after 2 days as a result of severe GI toxicity.
Monkey / Cynomolgus	Oral gavage	10 days	1, 10, 15 (5-days on, 2-days off), 25 (2-3 days), 62.5 (single dose)	1 M 1 F	≥ 1 mg/kg/day: Vomitus and fecal changes (soft, liquid, bloody, mucous). ≥ 15 mg/kg/day: Decreased food consumption, lymphoid depletion in the spleen and/or thymus, decreased spleen weights (15 mg/kg), and minimal enteropathy in the small intestine (10 and 15 mg/kg). Excretion of dasatinib in the urine increased from < 1% to up to 220-fold over the 10 day period in female monkeys. ≥ 25 mg/kg/day: Mortality (75%, both monkeys at 25 mg/kg and the female at 62.5 mg/kg; a male monkey was given a single dose of 62.5 mg/kg and discontinued). Prior to death, decreased activity, pale mucous membranes, hunched posture, and/or hypothermia. Red discoloration of the stomach (25 mg/kg) and small intestine (25 and 62.5 mg/kg), and red contents in the stomach and intestines (62.5 mg/kg). At 25 mg/kg, lymphoid depletion of intestinal lymphoid nodules and mesenteric lymph nodes and, at 62.5 mg/kg, edema, hemorrhage, and ulceration in the small intestine and tubular dilatation and degeneration in the kidney

Monkey / Cynomolgus	Oral gavage	1 month (5-days on, 2-days off)	1, 5, 15	4 M 4 F	1 mg/kg/day: No drug-related effects. ≥ 5 mg/kg/day: Fecal changes (liquid, nonformed, or no feces). 15 mg/kg/day: Vomitus, decreased body weight gain (F), and, in 1 M, hunched posture and thin, dehydrated appearance. Abnormal contents (gas and fluid) in the cecum and colon (F). Increases in ALT and decreases in albumin (M). Increases in liver weights and decreases in thymus weights (M). Splenic lymphoid depletion (M) and thymic lymphoid depletion.

Monkey / Cynomolgus	Oral gavage	9 months (5-days on, 2-days off)	1, 3/2, 10/6/4.5	6 M 6F	As a result of GI toxicity, the high dose of 10 mg/kg was reduced to 6 mg/kg in Week 3 and then again to 4.5 mg/kg in Week 12; the intermediate dose of 3 mg/kg was reduced to 2 mg/kg in Week 28. ≥ 1 mg/kg/day: Fecal changes (discolored, liquid, mucoid, nonformed and/or decreased), and low or no food consumption. Erosion/ulceration, acute to subacute inflammation, and epithelial flattening in the large intestine, and increased mineralization in the kidney. ≥ 3/2 mg/kg/day: Mortality (50%) primarily due to GI toxicity. Mean systemic exposure of dasatinib in the animals at 3/2 mg/kg/day reached only half the AUC of humans at the therapeutic dose (70 mg, BID). Prior to death, vomitus, hunched posture, hypoactivity, and decreased individual body weights. Decreased erythrocyte and lymphocyte counts, hemoglobin, hematocrit, albumin, sodium, potassium, and chloride, and increased total leukocyte and neutrophil counts, fibrinogen, urea nitrogen, and creatinine. Red foci in the large intestine and/or stomach. Lymphoid depletion in the thymus and spleen, and decreases in erythroid cells of the bone marrow. 10/6/4.5 mg/kg/day: Mortality (100%). None of the monkeys in this dosing group completed the nine month study due to unscheduled euthanasia that resulted from toxicity. Erosion/ulceration in the stomach (1 F), enlarged, gas-distended GI tract (1 M), and red, fluid contents in the stomach and small intestine (1 M).

Carcinogenicity:

In a 2-year carcinogenicity study, rats were administered oral doses of dasatinib at 0.3, 1, and 3 mg/kg/day. The highest dose resulted in a plasma drug exposure (AUC) levels generally equivalent to the human exposure at the recommended starting dose of 100 mg daily. A statistically significant increase in the combined incidence of squamous cell carcinomas and papillomas in the uterus and cervix of high-dose females (P = 0.0031) and of prostate adenoma in low-dose males (P = 0.0088; when the intermediate- and high-doses were excluded from the analysis due to increased incidence of mortality at these dose levels) was noted.

Genotoxicity:

Dasatinib was clastogenic *in vitro* to dividing Chinese hamster ovary cells with and without metabolic activation at concentrations ≥ 5 µg/mL. Dasatinib was not mutagenic when tested in *in vitro* bacterial cell assays (Ames test) and was not genotoxic in an *in vivo* rat micronucleus study.

Test / Test System	Route of Administration	Duration of Dosing	Concentration/ Dose	N/Dose/ Sex	Findings
Bacterial Mutagenicity Screening (Spiral Ames reverse mutation) *S. typhimurium*	*In vitro*	48 hr	21 - 5000 µg/plate, with and without rat S9 activation	NA	Not mutagenic.
Bacterial Mutagenicity Screening (Exploratory Ames reverse mutation) *S. typhimurium*	*In vitro*	48 hr	5 - 5000 µg/plate, with and without rat S9 activation	NA	Not mutagenic.
Bacterial Mutagenicity (Reverse mutation, definitive study) *S. typhimurium* and *E. coli*	*In vitro*	46-50 hr	12.5 - 400 µg/plate (*S. typhimurium*); 50-1600 µg/plate (*E. coli)*, with and without rat S9 activation	NA	Not mutagenic.

Cytogenetics Study Chinese hamster ovary cells	*In vitro*	4-20 hr	2.5 - 60 μg/mL, with and without activation	NA	Genotoxic effects: Chromatid and chromosome structural aberrations at ≥ 20 μg/mL (4 hr -S9), 5 μg/mL (4 hr +S9), and ≥ 5 μg/mL (20 hr -S9).
Oral Micronucleus Rat / SD	Oral gavage	3 days	10, 20, 40 mg/kg	5 M 5 F	Genotoxic effects: None.

Reproductive and Developmental Toxicology:

Dasatinib, when administered to pregnant rats during organogenesis at doses of 2.5, 5, 10, or 20 mg/kg, induced fetal toxicity (embryolethality with associated decreases in litter size, and fetal skeletal abnormalities, including malformations) at all doses, and maternal toxicity at doses ≥ 10 mg/kg. Maternal death occurred at 20 mg/kg. In a range-finding study in pregnant rabbits, dasatinib administered during organogenesis caused embryolethality of 13% at 6 mg/kg and 69% at 10 mg/kg. In the definitive embryo-fetal development study in rabbits, dasatinib did not cause maternal toxicity at 0.5, 2, or 6 mg/kg, whereas drug-related fetal skeletal alterations, including malformations, occurred at all doses.

In the oral study of fertility and early embryonic development in rats, dasatinib was not a reproductive toxicant in male rats at doses (≤ 10 mg/kg/day) that approximated human clinical exposures. In female rats, dasatinib did not affect mating or fertility at doses up to10 mg/kg/day, but induced embryo lethality at doses of ≥ 5 mg/kg/day (post-implantation losses of 14 to 48%, relative to 4% in controls) with associated decreases in litter size. Dasatinib is a selective reproductive toxicant in female rats at clinically relevant systemic exposures.

Dasatinib at doses of 5 and 10 mg/kg/day was given orally to female rats in 3 cohorts for which dosing was initiated on Gestation Day (GD) 16 (the end of organogenesis), GD 21 (the approximate onset of parturition), or Lactation Day (LD) 4 and continued up to LD 20. In all cohorts, in utero or lactational exposure to dasatinib in pups was associated with pleural effusion. For cohorts starting dasatinib on GD 16 or 21 at either dose, all groups were discontinued following 6 to 9 doses when more than 50% of pups had been euthanatized, found dead, or missing/presumed cannibalized. Among dams for which dosing initiated on LD 4, 34% of pups were lost due to mortality or moribundity at 10 mg/kg/day.

Study Type Species/Strain	Route of Administration	Duration of Dosing	Dose (mg/kg)	N/Dose/Sex	Findings
Embryofetal Development in Rats / SD	Oral gavage	10 days (GD 6 to 15)	2.5, 5, 10, 20	22 F	≥2.5 mg/kg: Embryolethality (17%) and associated decreases in litter size. Fetal skeletal abnormalities. ≥ 5 mg/kg: Embryolethality (77%). Fluid-filled thoracic and abdominal cavities, edema, microhepatia in fetus. ≥ 10 mg/kg: Embryolethality (100%). Decreased maternal food consumption. 20 mg/kg: Maternal mortality (22% during Days 12 - 15 of gestation). Decreased maternal body weight gain.
Range Finding Study in Rabbits / NZW	Oral gavage	13 days (GD 7 to 19)	1, 3, 6, 10	7 F	1 and 3 mg/kg: No drug-related effects. ≥ 6 mg/kg: Embryolethality (13%). Decreased maternal body weight gain and/or weight loss, and decreased food consumption. 10 mg/kg: Embryolethality (69%) and reduced number of litters with live fetuses at gestation day 29 (5/7).
Embryo-fetal Development in Rabbits / NZW	Oral gavage	13 days (GD 7 to 19)	0.5, 2, 6	22 F	No maternal toxicity. Delays in ossification of the fetal lumbar vertebrae (bifid arches) and pelvis (incompletely or unossified pubes), reduced ossification of hyoid (incompletely or unossified). 6 mg/kg: 21% of fetus resorption among rabbits with post-implantation loss.
Fertility and early embryonic development study in rats (Segment I)	Oral gavage	32 - 45 days 43 days	2.5, 5, 10	25 F 25 M	≤ 10 mg/kg: Dasatinib was not a reproductive toxicant in M and did not affect mating or fertility in F ≥ 5 mg/kg: Dasatinib induced embryo lethality (post implantation loss of 14 - 48%) in F and associated decreases in litter size.

Range finding pre- and post-natal development study in rats	Oral gavage	GD16, to LD 20 GD21 to LD 20 LD4 to LD 20	0, 5, 10	8F 8F 8F	5 mg/kg cohorts starting on GD 16 and 21: Profound pup mortality with associated decreases in litter sizes. Pleural effusion in 20 of 47 and 16 of 42 pups in cohorts starting on GD 16 and GD 21, respectively. 10 mg/kg all cohorts : Profound pup mortality with associated decreases in litter sizes. Pleural effusion in 30 of 30 and 25 of 57 pups in cohorts starting on GD 21 and LD4, respectively.

Special Toxicology:

Safety Pharmacology

Dasatinib had no significant effects in an *in vitro* ligand binding study. In the hERG/IKr assay, dasatinib inhibited hERG currents by 6, 37, and 77% at 3, 10, and 30 μm, respectively. The IC_{50} was 14.3 μM. In the Purkinje fiber assay, dasatinib prolonged APD_{50} by 26% and APD_{90} by 11% at 30 μM. Dasatinib at a single oral dose of 10 mg/kg in conscious, unrestrained monkeys (n = 6) elicited increases in blood pressure (6-15% in systolic and 8-21% in diastolic) for approximately 2 hours. In addition, mean QTc interval increases of 16-19 msec were observed between 1.5 – 2.5 hours post dose in the dasatinib-treated cohort compared to the vehicle control. Although these QTc changes were not statistically significant from control, an association of these changes with dasatinib treatment can not be excluded.

The N-dealkylated metabolite of dasatinib, BMS-582691 at 10 μM inhibited receptor-ligand binding to the adrenergic β_2, non-selective adrenergic α_2, non-selective serotonin 5-HT_1, serotonin 5-HT_{1A}, norepinephrine transporter, and dopamine transporter receptors, and to the sodium channel. In the hERG/IKr assay, BMS-582691 inhibited hERG currents with a calculated IC_{50} of 5.8 μM compared to 14.3 μM for dasatinib. In the Purkinje fiber assay, BMS-582691 at 30 μM. prolonged APD_{50} and APD_{90} by 10% and 9%, respectively, and reduced Vmax by 11%.

Study Type / Organ Systems Evaluated	Test System / Species/Strain	Route	Concentration/ Dose	N/Dose/ Sex	Findings
Receptor and Ion Channel Ligand Binding Study	Receptors, ion channels, and enzyme systems	*in vitro*	10 μM	--	No biologically significant effect on binding of ligands to receptors or ion-channels, or on acetylcholinesterase activity. BMS-582691 at 10 μM inhibited receptor-ligand binding to the adrenergic β_2 (50%), non-selective adrenergic α_2 (51%), non-selective serotonin 5-HT_1 (50%), serotonin 5-HT_{1A} (54%), norepinephrine transporter (54%), and dopamine transporter (87%) receptors, and to the sodium channel (84%)
hERG/IKr Channel Assay / Cardiovascular	HEK293 cells transfected with human hERG cDNA	*in vitro*	3, 10, 30 μM	--	Dasatinib: IKr currents were inhibited by 6, 37, and 77% at 3, 10 and 30 μM, respectively. The calculated IC_{50} was 14.3 μM. BMS-582691 inhibited IKr currents by 24, 72, and 95% at 3, 10 and 30 mM, respectively. The calculated IC_{50} was 5.8 μM
Rabbit Purkinje Fiber Action Potential Assay/ Cardiovascular	Rabbit Purkinje fibers	*in vitro*	3, 10, 30 μM	--	Dasatinib: APD_{50} and APD90 were prolonged by 26% and 11%, respectively, at 30 μM. BMS-582691: APD_{50} and APD_{90} were prolonged by 10% and 9%, respectively, and Vmax was reduced by 11%.
Single-Dose Safety Pharmacology / Cardiovascular	Monkey / Cynomolgus	Oral, single dose	10 mg/kg	3 M 3 F	Drug-related increases in systolic (6-15%) and diastolic (8-21%) blood pressure for approximately 2 hours and mean QTc increases of 16-19 msec between 1.5 – 2.5 hours following a single oral dose.

Other Toxicity Studies

The immunosuppressive potential of dasatinib was assessed in mouse models of T-cell proliferation (mixed lymphocyte response) and nonvascularized heart transplant rejection. The effects of dasatinib on *in vitro* platelet function were assessed in human, monkey, and rat plasma, and the effects on *in vivo* bleeding time were assessed in rats. The *in vitro* phototoxicity potential of dasatinib was assessed in mouse fibroblasts.

The effect of dasatinib on the cardiac sarcoplasmic reticulum and mitochondrial function is unknown. The potential for apoptosis in cardiomyocytes with dasatinib treatment has not been investigated, and no studies have been conducted with dasatinib to evaluate the potential signaling mechanism regulating cardiotoxicity.

Study Type / Test System	Route of Administration	Duration of Dosing	Dose (mg/kg)	N/Dose/Sex	Findings
Mixed Lymphocyte Response Assay/Mouse	Oral gavage	3 days	5, 20, 50	3 M	5 mg/kg: No effect on T-cell proliferation. ≥ 20 mg/kg: Dose-dependent inhibition of splenic T-cell proliferation.
Cardiac Transplant Study/Mouse	Oral gavage	30 days	15, 25, 50	4-5 M	15 mg/kg, twice daily (continuous daily dosing): Graft rejection not inhibited. 25 mg/kg, twice daily (5-days on, 2-days off schedule): Graft rejection not inhibited. 25 mg/kg, twice daily, (continuous daily dosing): Inhibition of graft rejection.
Platelet Function / Platelets from humans, cynomolgus monkeys, and rats	*In vitro*	--	0.05, 0.5, 5 μg/ mL	--	0.05 μg/mL: No effect. 0.5 and 5 μg/mL: Inhibition of the platelet aggregation response to ADP and collagen in human platelet-rich plasma, and inhibition of shear-induced aggregation of human platelets. 5 μg/mL: Decreased strength of human whole blood clots (29%); no effect on time to clot formation or rate of clot formation. In each species complete inhibition of the collagen response was observed with comparable IC50 values (μg/mL) for human (0.24 ± 0.06) and cynomolgus monkey (0.23 ± 0.06), and slightly but not significantly greater potency for rat (0.13 ± 0.01).

Bleeding Time and Platelet Function/Rat	Oral gavage or IV	Single oral dose or IV infusion	4, 8, 20 (mg/kg, oral) or 630, 1260,-2520 (μg/kg, IV)	5-9 M	Oral gavage: 4 mg/kg: No effect on mesenteric bleeding time, cuticle bleeding time, or ADP-induced platelet aggregation. 8 mg/kg: No effect on mesenteric bleeding time. The anticipated plasma concentration was not reached for evaluating the cuticle bleeding time and platelet aggregation. 20 mg/kg: 3-fold increase in cuticle bleeding time and inhibition of the platelet aggregation response (21 and 99%) induced by 10 μM ADP and 20 μg/mL collagen, respectively. IV infusion: Dasatinib produced dose-dependent increases in cuticle bleeding time at all doses (mean plasma concentrations as 61, 144, 273 ng/mL respectively) and proportion of vessels with re-bleeds and off scale bleeding at the high dose. A dose-dependent reduction in platelet aggregation (37%, 99% and 100%) was also observed at all doses.
Phototoxicity Assay/Mouse fibroblasts	*In vitro*	--	0.353-120 μg/mL	--	Results indicated that dasatinib is phototoxic *in vitro* to mouse fibroblasts.

PATIENT MEDICATION INFORMATION

READ THIS FOR SAFE AND EFFECTIVE USE OF YOUR MEDICINE

PRSPRYCEL®

dasatinib

Read this carefully before you start taking **SPRYCEL** and each time you get a refill. This leaflet is a summary and will not tell you everything about this drug. Talk to your healthcare professional about your medical condition and treatment and ask if there is any new information about **SPRYCEL**.

Serious Warnings and Precautions

Take SPRYCEL only under the care of a doctor who knows how to use anti-cancer drugs.

Serious and common side effects with SPRYCEL include:

- **Myelosuppression (thrombocytopenia, neutropenia, anemia)**: SPRYCEL can affect your body's ability to make blood cells. It can cause you to have low blood cell counts.
    - Neutropenia is a low white blood cell count. It can occur with and without a fever and can cause you to get infections.
    - Thrombocytopenia is low platelets in the blood. Platelets help with clotting.
    - Anemia is a low red blood cell count.

  Your doctor will do regular blood tests to monitor you for myelosuppression.
- **Bleeding**, which may result in death
- **Fluid retention**
- **Congestive heart failure (CHF):** This is when your heart doesn't pump as well as it should. Signs and symptoms of CHF are shortness of breath, swelling and weight gain, which are usually accompanied in almost all cases by fluid retention and pulmonary edema. Pulmonary edema is when fluid builds up in the lung.
- **Pulmonary Arterial Hypertension:** This is a condition where the blood pressure in the arteries of the lung is high.

What is SPRYCEL used for?

SPRYCEL is used to treat adults with certain types of leukemia including:

- Philadelphia chromosome positive (Ph+) chronic myeloid leukemia (CML) in chronic phase that has been recently diagnosed;
- Ph+ CML that is no longer benefiting from other available therapies for CML, including imatinib mesylate;

Ph+ acute lymphoblastic leukemia (ALL) that no longer responds to other therapies.

How does SPRYCEL work?

Leukemia is a cancer that affects different types of white blood cells. In patients with leukemia, these white blood cells are abnormal. They don't work properly and can multiply in an uncontrolled way.

SPRYCEL acts by stopping the activity of proteins in these abnormal white blood cells. This helps to slow the uncontrolled growth of the white blood cells.

What are the ingredients in SPRYCEL?

Medicinal ingredients: dasatinib

Non-medicinal ingredients: Croscarmellose sodium, hydroxypropyl cellulose, lactose monohydrate, magnesium stearate and microcrystalline cellulose. The tablet coating consists of hypromellose, titanium dioxide and polyethylene glycol.

SPRYCEL comes in the following dosage forms:
Tablet: 20, 50, 70, 80, 100 and 140 mg.

Do not use SPRYCEL if:
- You are allergic to dasatinib or to any other ingredients in SPRYCEL. Tell your healthcare provider if you think you have had an allergic reaction to any of these ingredients.
- You are breast-feeding. [text]

To help avoid side effects and ensure proper use, talk to your healthcare professional before you take SPRYCEL. Talk about any health conditions or problems you may have, including if you:
- Have a liver problem.
- Have a heart problem, such as an irregular heartbeat or a hereditary disorder of the heart's electrical activity, called long QT syndrome.
- Have or have previously had a hepatitis B infection. This is an infection of the liver. SPRYCEL could cause the hepatitis B virus to become active again, which can lead to death in some cases. Your doctor will check for signs of this infection before starting treatment with SPRYCEL. If the hepatitis B virus is found, you will be monitored closely during and for several months after treatment with SPRYCEL.
- Are lactose intolerant or have one of the following rare hereditary diseases:
  - Galactose intolerance
  - Lapp Lactase deficiency
  - Glucose-galactose malabsorption

  This is because lactose is a non-medicinal ingredient in SPRYCEL.
- Are taking medicines to thin the blood or prevent clots. SPRYCEL may cause bleeding.
- Have muscle aches/pains or weakness, or dark-colored urine. [text]

Other warnings you should know about:
Female patients:
- If you are pregnant or planning to become pregnant there are specific risks you must discuss with your healthcare professional.
- Do not become pregnant while taking SPRYCEL. It may harm your unborn baby or make you lose the pregnancy.
- Use highly effective methods of birth control while taking SPRYCEL. Your healthcare professional can tell you about the types of birth control available to you.
- If you do become pregnant while you are using SPRYCEL, tell your healthcare professional right away.
- SPRYCEL may affect your ability to have a child in the future. Talk to your healthcare professional if you have questions about this.

Male patients:
- Use highly effective methods of birth control each time you have sex with a woman during your treatment with SPRYCEL.

Blood tests:
During your treatment with SPRYCEL you will need to have blood tests done. These will be done about every 1 to 2 weeks for the first few months of your treatment. You will then need to have these tests repeated once every 1 to 3 months. These tests will tell your healthcare professional how SPRYCEL is affecting your blood. They will also show how well your liver and kidneys are working. [text]

Tell your healthcare professional about all the medicines you take, including any drugs, vitamins, minerals, natural supplements or alternative medicines.

The following may interact with SPRYCEL:

- Medicines used to treat irregular heart beat such as: disopyramide, procainamide, amiodarone, sotalol, ibutilide and flecainide.
- Medicines used to stabilize your mood like benzodiazepine, chlorpromazine, haloperidol and pimozide.
- Medicines used to treat chronic or severe pain like methadone.
- Medicines used to treat malaria like chloroquine.
- A medicine that stimulates stomach and bowel movement called domperidone.
- Medicines used to treat fungal infections, like ketoconazole and itraconazole.
- Medicines used to treat bacterial infections like erythromycin and clarithromycin, quinolone, moxifloxacin.
- A medicine used to treat HIV the virus that causes AIDS like ritonavir, lopinavir and atazanavir.
- A medicine used to treat tuberculosis called rifampicin.
- Medicines used to treat epilepsy like carbamazepine, phenytoin and phenobarbital.
- Medicines used to treat high cholesterol like simvastatin.
- A medicine used to prevent organ rejection or treat autoimmune conditions called cyclosporine.
- Medicines used to treat inflammation like dexamethasone.
- An herbal remedy used to treat depression called St. John's Wort.
- Medicines used to treat severe headaches or migraines like ergotamine and dihydroergotamine.

Do not eat or drink any products or juices that contain grapefruit or grapefruit juice. These can affect how SPRYCEL works.

Avoid taking medicines that neutralise stomach acids. Examples are antacids such as cimetidine, famotidine, ranitidine and omeprazole. If you must use these medicines, take them at least 2 hours before or 2 hours after taking SPRYCEL.

Tell your doctor if you are taking medicines to thin the blood or prevent clots like warfarin sodium or aspirin.

How to take SPRYCEL:
- Exactly as directed by your healthcare professional.
- Once per day, either in the morning or in the evening.
- With or without food, at about the same time each day.
- Swallow whole. Do not crush or cut tablets.

Usual dose:
Your dose of SPRYCEL will depend on the type of leukemia you have.
- Usual starting dose for chronic phase CML: 100 mg once a day.
- Usual starting dose for accelerated or blast crisis CML or Ph+ ALL: 140 mg once a day.

Your healthcare professional may interrupt or change your dose of SPRYCEL if:
- You are taking certain medications,
- You do not tolerate the treatment, or
- Your disease gets worse.

Overdose:
If you take too much SPRYCEL, you may experience side effects including low platelet counts.

If you think you, or a person you are caring for, have taken too much [Brand name], contact a healthcare professional, hospital emergency department, or regional poison control centre immediately, even if there are no symptoms.

Missed Dose:
If you miss a dose of SPRYCEL, wait until it is time for your next dose. Do not take two doses at the same time. Call your healthcare provider or pharmacist if you are not sure what to do.

What are possible side effects from using SPRYCEL?
These are not all the possible side effects you may have when taking [Brand name]. If you experience any side effects not listed here, tell your healthcare professional.

- Diarrhea
- Nausea
- Vomiting
- Stomach pain
- Fever
- Headache
- Fatigue
- Skin rash
- Shortness of breath
- Cough
- Upper respiratory tract infection
- Infection
- Pain
- Bone and extremity pain
- Muscle and joint aches

SPRYCEL can cause abnormal blood test results. Your doctor will decide when to test your blood and will interpret the results.

The following have been reported in patients using SPRYCEL: inflammation of the lungs, blood clots, irregular heart rhythm, and deaths from gastrointestinal bleeding. These may or may not have been related to SPRYCEL.

Serious side effects and what to do about them			
Symptom / effect	**Talk to your healthcare professional**		**Stop taking drug and get immediate medical help**
	Only if severe	**In all cases**	
COMMON			
Myelosuppression (low blood cell counts): such as **anemia** (low red blood cell counts), **neutropenia** (low white blood cell counts), or **thrombocytopenia** (low platelet counts)		√	
Bleeding (loss of blood or bruising without having an injury no matter how mild): bleeding; bruising; blood in vomit, stools or urine; or black stools; bleeding from the nose or gums, excessive period bleeding		√	
Fluid retention (build-up of water in your body, which can be in the lining of your lungs or around your heart): swelling anywhere on or in your body, weight gain; shortness of breath, especially after low levels of physical exertion; chest pain when taking a		√	

deep breath			
Heart problems (Irregular heart rate, heart attack): heartbeat that is abnormally slow, fast or forceful; shortness of breath; dizziness or feeling faint; chest pain accompanied with fatigue, nausea or cold sweats			√
Infections (bacterial or viral illness): fever, severe chills, discharge (fluid) with mucus or pus		√	
Myelosuppression (low blood cell counts): such as **anemia** (low red blood cell counts), **neutropenia** (low white blood cell counts), or **thrombocytopenia** (low platelet counts)		√	
UNCOMMON			
Liver damage (inflammation of the liver, increased liver enzyme levels on blood tests): yellow skin and/or eyes, nausea, loss of appetite, dark-coloured urine		√	
Rhabdomyolysis (breakdown of damaged muscle); muscle aches and pain, weakness, dark urine		√	
RARE			
Pulmonary arterial hypertension (increased blood pressure in the arteries supplying the lungs): shortness of breath, fatigue		√	
VERY RARE			
Stevens-Johnson syndrome (severe skin reaction): redness, blistering and/or peeling of the skin or mucous membranes (skin of lips, eyes, mouth, nasal passages, genitals) with fever, sore mouth or throat; can lead to death			√
Hepatitis B virus reactivation (an active viral infection of the liver): Weight loss, fever, abdominal pain, nausea and vomiting followed by jaundice (yellowing of the skin or whites of eyes)		√	
Erythema multiforme (severe skin reaction): raised red or purple skin patches with itching or burning, sores with puss			√
Thrombotic microangiopathy (damage to blood vessels): Bruising, bleeding, weakness, fever, fatigue and confusion.			√

If you have a troublesome symptom or side effect that is not listed here or becomes bad enough to interfere with your daily activities, tell your healthcare professional.

Reporting Side Effects
You can report any suspected side effects associated with the use of health products to Health Canada by:

- Visiting the Web page on Adverse Reaction Reporting (https://www.canada.ca/en/health-canada/services/drugs-health-products/medeffect-canada.html) for information on how to report online, by mail or by fax; or
- Calling toll-free at 1-866-234-2345.

NOTE: Contact your health professional if you need information about how to manage your side effects. The Canada Vigilance Program does not provide medical advice.

Storage:
Store at room temperature between 15°C to 30°C.

Keep out of reach and sight of children.

Do not use SPRYCEL after the expiry date written on the label, blister or carton after EXP.

If you want more information about SPRYCEL:

- Talk to your healthcare professional
- Find the full product monograph that is prepared for healthcare professionals and includes this Patient Medication Information by visiting the Health Canada website: (https://www.canada.ca/en/health-canada/services/drugs-health-products/drug-products/drug-product-database.html; the manufacturer's website https://bms.com/ca/en, or by calling 1-866-463-6267.

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