Inhibition of Human Tumor Xenograft Growth by Treatment with the Farnesyl Transferase Inhibitor B956

Takeki Nagasu, Kentaro Yoshimatsu, Cheryl Rowell, Michael D. Lewis, and Ana Maria Garcia

ABSTRACT

ras oncogenes are present in several types of cancers but are most frequently described in colon and pancreatic carcinomas. Consequently, ras is being targeted for drug development as a means to develop therapies for these types of cancer. The ras protein is posttranslationally modified by the addition of a farnesyl group, followed by cleavage of the COOH-terminal 3 amino acids and methylation of the prenylated cysteine. Because the posttranslational addition of farnesyl is obligatory not only for the remaining modifications to take place but also for ras control of cell growth, inhibitors of farnesylation are being developed as potential antitumor agents. In this report, a new peptidomimetic inhibitor of farnesyl transferase is described. This compound, B956, and its methyl ester B1086, inhibit the formation of colonies in soft agar of 14 human tumor cell lines expressing different ras oncogenes at concentrations between 0.2 and 60 μM. Higher concentrations of B956 (10—80 μM) were required to inhibit colony formation by 5 tumor cell lines without ras mutations. B956/B1086 at 100 mg/kg also inhibited tumor growth by EJ-1 human bladder carcinoma, HT1080 human fibrosarcoma, and to a lesser extent by HCT116 human colon carcinoma xenografts in nude mice. Furthermore, inhibition of tumor growth by B956 is shown to be correlated with inhibition of ras posttranslational processing in the tumor. Thus, peptidomimetic inhibitors of ras farnesylation have the potential to be developed as therapy for ras-dependent tumors.

INTRODUCTION

The ras proto-oncogene is involved in the control of cell proliferation and differentiation. In normal cells, ras switches between inactive GDP-bound and active GTP-bound states (1). Binding of growth factors like epidermal growth factor or platelet-derived growth factor to their receptors results in the activation of the ras proteins by promoting the exchange of bound GDP for GTP (2). The activated ras, by a still unknown mechanism, triggers a phosphorylation cascade involving raf, mitogen-activated protein kinase kinase, and mitogen-activated protein kinase, and ending in the activation of nuclear transcription factors. The final result is stimulation of cell growth or differentiation. Inactivation of ras occurs when the bound GTP is hydrolyzed to GDP, a reaction stimulated by GTPase-activating protein. The cycle is thus closed, and ras remains in inactive form until a new growth signal arrives (for reviews, see Refs. 3 and 4).

Oncogenic ras protein has lost the ability to switch between the inactive and active states and remains permanently activated. Although there are several ways in which ras can remain activated, the most common seems to be mutations at codons 12, 13, or 61, which result in loss of the ras-GTPase activity. Whereas it is not clear what triggers the specific mutations that activate ras, the result is a constitutively activated protein that constantly sends growth signals to the cell nucleus. The presence of oncogenic forms of ras has been detected in 20—30% of all types of human cancers, in up to 50% of colon cancer, and in more than 80% of pancreatic carcinomas (5). The high incidence of ras mutations in colon and pancreatic carcinomas has motivated a great deal of research on ways to inhibit the activity of the ras oncogene.

Signal transduction by ras is dependent on its plasma membrane localization (reviewed in Ref. 6). ras is localized at the plasma membrane after posttranslational farnesylation of the cysteine in the CAAX consensus sequence at the COOH terminus of the protein (7, 8). Farnesylation is followed by proteolytic cleavage of the last three amino acids, methylation of the now farnesylated cysteine, and, in the case of H- and N-ras, palmitylation of upstream cysteines. Several reports have indicated that farnesylation is critical for the subsequent modifications, as well as membrane localization and function, of the ras protein (9—12). Therefore, inhibition of farnesylation is being sought by several laboratories as a possible mechanism for inhibition of tumor growth (13). In this report, we describe in vivo data obtained with a potent new peptidomimetic inhibitor of farnesylation, B956. This compound inhibits the growth of tumors induced in nude mice by ras-transformed cells, including 3 human xenografts expressing different types of oncogenic ras. B956 also inhibits the ability of several human tumor cell lines to form colonies in soft agar, a common assay for tumorigenicity. Furthermore, it is shown that inhibition of tumor growth is correlated with inhibition of ras posttranslational processing in the tumor. Although differences in sensitivity to the inhibitor were observed in different cell lines, inhibition of ras farnesylation as antitumor drug therapy appears promising.

MATERIALS AND METHODS

Cells. NIH 3T3 cells transformed with H-ras (61L) (zH1, or pZIP-ras-F) have been described previously (11). NIH 3T3 cells transformed with K-ras (12V) (DK1) were prepared by infection with retrovirus carrying K-ras 4B (12V) clone. Infected cells were selected with G418, and single clones were isolated, grown, and tested for K-ras expression, growth in soft agar, and tumor formation in nude mice. A single clone (DK1) was chosen for these studies. EJ-1 cells were kindly provided by Dr. C. Der (University of North Carolina, Chapel Hill, NC). The remaining cell lines were obtained from American Type Culture Collection (Rockville, MD) or Dainippon Seiyaku (Osaka, Japan) as indicated in Table 1. All cells were maintained as recom
were injected i.p. Treatment was started on day 1 at the concentrations and was administered i.p. to 3 mice, at a concentration of 200 mg/kg every 12 h for vehicle. Tumor volume was measured on the days indicated, and body weight treatment schedules indicated in the figure legends. Control animals received B1086 were dissolved in saline containing 2% Tween 80 just before use and by Western blotting with F235—1.7.1 antibody (Oncogene Science).

Cell debris was removed by centrifugation at 12,000 X g for 5 mm, followed by cytosol and membrane fractionation by centrifugation at 100,000 X g for 30 min. Protein (50 pg/lane) was separated by SDS-PAGE and ras was identified by about 50% (11). The fact that non-ras-mutated cell lines are inhibited is not clear. In summary, transformants expressing either H- or N-ras oncogene seem to be the most sensitive, followed by cell lines expressing mutated N-ras. The more resistant tumor cell lines are the ones expressing mutant K-ras as well as cells without ras mutations. The IC<sub>50</sub> for inhibition of colony formation among the human cell lines expressing H-ras oncogene seem to be the most sensitive, followed by cell lines expressing mutated N-ras. The more resistant tumor cell lines are the ones expressing mutant K-ras as well as cells without ras mutations. The IC<sub>50</sub> for inhibition of colony formation among the human cell lines expressing H-ras oncogene fluctuates between 1.7 µM for the most sensitive (HCT116) and 50 µM for the least sensitive (HCT15). In contrast, H-ras transformed cell lines are inhibited by submicromolar concentration of B956, whereas for N-ras-transformed cells, 50% inhibition occurs at concentrations around 5 µM B956.

In contrast to other reports (14, 15), transformed cell lines without ras mutations were inhibited in their capacity to form colonies in soft agar at concentrations between 16 and 80 µM B956. Approximately one-half of the K-ras-transformed cell lines tested were inhibited at the same concentrations of B956. We have previously shown a similar result with another inhibitor, B581 (11). This compound at 100 µM inhibited colony formation of NIH 3T3 fibroblasts transformed either with theraf oncogene or mutant H-ras modified by geranylgeranylation or myristylation by about 50% (11). The fact that non-ras-mutated cell lines are inhibited in their colony-forming capacity is not unexpected if other farnesylated proteins, like lamins or other yet unidentified proteins, are considered. However, the involvement of these proteins in the control of cell growth is not clear. In summary, transformants expressing either H- or N-ras oncogenes are very sensitive to inhibition by the FTase inhibitor B956. In contrast, only 50% of K-ras mutant cell lines are inhibited at relatively low (i.e., below 10 µM) concentrations of inhibitor. The remaining 50% of the K-ras-transformed cells tested were inhibited by concentrations of B956 that also inhibit non-ras-mutated cell lines (i.e., between 10 and 100 µM).

B956 inhibits H- and K-ras posttranslational processing by 50% at 0.5 and 25 µM, respectively. In this report, these studies have been extended to the effect of B956 on anchorage-independent growth of 19 human tumor cell lines, as well as the 2 model cell lines zH1 and DK1. The panel of human tumor cell lines tested is summarized in Table 1, where the presence and identity of ras mutation is indicated. This group of 19 cell lines represents a variety of tissues, as well as ras mutations. Treatment with B956 resulted in inhibition of the number and size of colonies formed. The concentration of B956 that reduced the number of colonies by 50% is shown in Fig. 2. There is a similar inhibition pattern for the formation of colonies in soft agar as that seen for inhibition of ras processing (namely, cell lines expressing different types of ras show differences in sensitivity to the inhibitor). Tumor cell lines expressing H-ras oncogene seem to be the most sensitive, followed by cell lines expressing mutated N-ras. The more resistant tumor cell lines are the ones expressing mutant K-ras as well as cells without ras mutations. The IC<sub>50</sub> for inhibition of colony formation among the human cell lines expressing H-ras oncogene seemed to be the most sensitive, followed by cell lines expressing mutated N-ras. The more resistant tumor cell lines are the ones expressing mutant K-ras as well as cells without ras mutations. The IC<sub>50</sub> for inhibition of colony formation among the human cell lines expressing H-ras oncogene seemed to be the most sensitive, followed by cell lines expressing mutated N-ras. The more resistant tumor cell lines are the ones expressing mutant K-ras as well as cells without ras mutations. The IC<sub>50</sub> for inhibition of colony formation among the human cell lines expressing H-ras oncogene seemed to be the most sensitive, followed by cell lines expressing mutated N-ras.

**RESULTS**

**Inhibition of Anchorage-independent Growth of Human Tumor Cell Lines.** B956 (Fig. 1) is a new and potent inhibitor of FTase<sup>2</sup> activity in vitro. The concentration to inhibit H-ras farnesylation by 50% is in the nanomolar range (IC<sub>50</sub> = 11 nm<sup>3</sup>). In whole cells (zH1 or DK1),

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### Table 1 Human tumor cell lines

<table>
<thead>
<tr>
<th>Cell line</th>
<th>Tissue of origin</th>
<th>ras mutation (Ref.)</th>
<th>No. of cells/well</th>
</tr>
</thead>
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<tr>
<td>H520 (A)</td>
<td>Lung</td>
<td>27</td>
<td>2 X 10&lt;sup&gt;3&lt;/sup&gt;</td>
</tr>
<tr>
<td>COLO320DM (D)</td>
<td>Colon</td>
<td>28</td>
<td>1 X 10&lt;sup&gt;2&lt;/sup&gt;</td>
</tr>
<tr>
<td>COLO205 (A)</td>
<td>Colon</td>
<td>28</td>
<td>4 X 10&lt;sup&gt;3&lt;/sup&gt;</td>
</tr>
<tr>
<td>HT29 (A)</td>
<td>Colon</td>
<td>29</td>
<td>2 X 10&lt;sup&gt;4&lt;/sup&gt;</td>
</tr>
<tr>
<td>WiDr (D)</td>
<td>Colon</td>
<td>30</td>
<td>1 X 10&lt;sup&gt;2&lt;/sup&gt;</td>
</tr>
<tr>
<td>A-427 (A)</td>
<td>Lung</td>
<td>K-12D (33)</td>
<td>5 X 10&lt;sup&gt;3&lt;/sup&gt;</td>
</tr>
<tr>
<td>A549 (D)</td>
<td>Lung</td>
<td>K-12S (33)</td>
<td>5 X 10&lt;sup&gt;2&lt;/sup&gt;</td>
</tr>
<tr>
<td>SW480 (A)</td>
<td>Colon</td>
<td>K-12V (34)</td>
<td>4 X 10&lt;sup&gt;3&lt;/sup&gt;</td>
</tr>
<tr>
<td>SW620 (A)</td>
<td>Colon</td>
<td>K-12V (35)</td>
<td>1 X 10&lt;sup&gt;3&lt;/sup&gt;</td>
</tr>
<tr>
<td>DLD-1 (A)</td>
<td>Colon</td>
<td>K-13D (23)</td>
<td>1 X 10&lt;sup&gt;3&lt;/sup&gt;</td>
</tr>
<tr>
<td>HCT116 (A)</td>
<td>Colon</td>
<td>K-13D (23)</td>
<td>1 X 10&lt;sup&gt;3&lt;/sup&gt;</td>
</tr>
<tr>
<td>H460 (A)</td>
<td>Lung</td>
<td>K-61H (36)</td>
<td>1 X 10&lt;sup&gt;3&lt;/sup&gt;</td>
</tr>
<tr>
<td>HCT-15 (A)</td>
<td>Colon</td>
<td>K-61D (37)</td>
<td>2 X 10&lt;sup&gt;3&lt;/sup&gt;</td>
</tr>
<tr>
<td>MIA PaCa-2 (A)</td>
<td>Pancreas</td>
<td>K-12C (38)</td>
<td>1 X 10&lt;sup&gt;3&lt;/sup&gt;</td>
</tr>
<tr>
<td>HT-1080 (D)</td>
<td>Fibrosarcoma</td>
<td>N-61K (39)</td>
<td>5 X 10&lt;sup&gt;2&lt;/sup&gt;</td>
</tr>
<tr>
<td>THP-1 (A)</td>
<td>Leukemia</td>
<td>N-61D (40)</td>
<td>2.5 X 10&lt;sup&gt;3&lt;/sup&gt;</td>
</tr>
<tr>
<td>HL-60 (A)</td>
<td>Leukemia</td>
<td>N-61L (41)</td>
<td>1 X 10&lt;sup&gt;3&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

* Origin of cells: A, American Type Culture Collection (Rockville, MD); D, Dainippon Seiyaku (Osaka, Japan).

**Fig. 1. Structure of farnesyl transferase inhibitors.**

B1086 were dissolved in saline containing 2% Tween 80 just before use and were injected i.p. Treatment was started on day 1 at the concentrations and treatment schedules indicated in the figure legends. Control animals received vehicle. Tumor volume was measured on the days indicated, and body weight was determined twice weekly. Tumor volume was calculated according to the following equation: tumor volume (mm<sup>3</sup>) = (Length X Width<sup>2</sup>) / 2.

After 2-3 weeks, mice were sacrificed, and tumor weights were determined. Significance was determined by Student's t test.

**Inhibition of ras Posttranslational Processing.** Tumors were induced in 6 nude mice by zH1 cells as above. At day 6, when the tumor was visible, B956 was administered i.p. to 3 mice, at a concentration of 200 mg/kg every 12 h for a total of 7 injections. After determining tumor size, the mice were sacrificed, and the tumor was minced and treated with collagenase (1.2 mg/ml) and DNase (0.3 mg/ml). A single cell suspension was obtained by filtration through a nylon mesh. The cells were washed with cold PBS and lysed by sonication in 5 mm Tris-HCl (pH 7.5)-5 mm EDTA-1 mm phenylmethylsulfonyl fluoride. Cell debris was removed by centrifugation at 12,000 X g for 5 min, followed by cytosol and membrane fractionation by centrifugation at 100,000 X g for 30 min. Protein (50 µg/lane) was separated by SDS-PAGE and ras was identified by Western blotting with F235—1.7.1 antibody (Oncogene Science).

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2 The abbreviation used is: FTase, farnesyl protein transferase.

3 Unpublished observations.

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**Fig. 2. Soft agar colony formation by human tumor cell lines.** The indicated cell lines were grown in soft agar as described in "Materials and Methods." At 14-21 days, colonies were stained and counted. Cell lines are grouped according to the mutant form of ras they harbor: H-, N-, or K-ras, or normal (no ras mutation), as indicated on the abscissa. The concentration of B956 that inhibited the number of colonies to one-half is plotted as the ordinate for each cell line. Arrows, IC<sub>50</sub> for inhibition of colony formation by H-ras (61L) or K-ras (12V) NIH 3T3-transformed fibroblasts (zH1 and DK1 cells, respectively).
Inhibition of Tumor Growth Induced by ras-transformed 3T3 Fibroblasts. Nude mice were implanted s.c. with H-ras-transformed NIH 3T3 fibroblasts and treated systemically by two daily i.p. injections of 125 or 250 mg/kg B956 for 5 days. Tumors were measured every 3–4 days until mice were sacrificed at day 14. The results (Fig. 3) clearly show that B956 treatment inhibits tumor growth in a dose-dependent manner. Tumor growth resumes after a delay of several days, depending on the dose of B956 used. The rate of tumor growth after the delay is similar to that of untreated animals. No signs of toxicity were observed, and all the animals gained weight normally during the experiment.

Inhibition of ras Processing Associated with Inhibition of Tumor Growth. To determine whether the inhibition of tumor growth was in fact due to inhibition of ras processing, ras localization was determined after B956 treatment. For this purpose, mice given implantations of ZH1 cells were treated with B956 6 days after implantation, when the tumor was visible. After a 3-day treatment with the farnesyl transferase inhibitor, a significant reduction of tumor growth was observed (Fig. 4A). At this time, tumors from control and B956-treated animals were excised, homogenized, and separated into membrane and cytosol fractions. Fully processed ras localizes to the plasma membrane, whereas unprocessed ras, because of the lack of prenyl anchor, is not membrane associated but accumulates in the cytosol (16). As expected, all the ras protein was detected in the membrane fraction of tumor cells from control animals (Fig. 4B, Lane 1). In contrast, in the tumors from B956-treated animals, ras with a slower mobility accumulated in the cytosol (Fig. 4B, Lane 4), and almost no ras was detected in the membrane fraction (Fig. 4B, Lane 3). To our knowledge, this is the first demonstration that the decrease in tumor growth induced by a farnesyl transferase inhibitor is correlated with inhibition of ras farnesylation.

Inhibition of Growth of Human Tumor Xenografts. Preliminary experiments with the methyl ester prodrug B956, compound B1086, indicated that the latter was at least twice as potent at inhibiting tumor growth induced by 3T3 fibroblasts expressing the H-ras oncogene. Therefore, B1086 was used to test whether tumors induced by human tumor cell lines expressing oncogenic ras protein were also sensitive to FTase inhibitors. Nude mice were given implantations s.c. of human tumor cell lines expressing oncogenic ras protein: H-ras (HT-1080 fibrosarcoma; Fig. 5A), K-ras (HCT 116 colon carcinoma; Fig. 5C) oncogenes. Tumors induced by EJ-1 cells grew slowly and were very sensitive to treatment with B1086. The effect of the inhibitor was dose dependent. At the maximum dose of 100 mg/kg, no apparent tumor growth was observed, and 1 mouse of 5 had no visible tumor at the end of the experiment. Tumors induced by EJ-1 cells were found to be quite resistant to conventional anti-tumor drugs (such as cisplatin and doxorubicin; data not shown). However, the FTase inhibitor B1086 at the dose of 100 mg/kg inhibited tumor growth completely.

The effect of B1086 on HT-1080-induced tumors is not as striking as that observed against EJ-1 xenografts. However, there is a statistically significant difference between the tumor growth of treated and control animals. The tumors in B1086-treated animals grew more slowly, and the effect was dose dependent (Fig. 5B). At the end of the experiment, tumors of animals treated with 100 mg/kg were 40% smaller than those of control animals (Fig. 5B). HT-1080 xenografts induce cachexia to the animals; therefore, long experiments are not possible. Animals bearing tumors in the control group, as well as in the 25 and 50 mg/kg B1086 treatment groups, showed signs of necrosis and bleeding at the tumor site. However, animals treated with 100 mg/kg B1086 looked healthier than untreated animals.

The human tumor carcinoma cell line HCT116 was the least sensitive of the tumor xenografts tested (Fig. 5C), despite the fact that when grown in soft agar it was more sensitive to the inhibitor than were HT1080 cells (Fig. 2). Changes in growth properties in vitro and in vivo could account for this difference. Treatment with 100 mg/kg B1086 significantly reduced tumor size by only 20% (P < 0.05). Reduction of tumor growth rate was observed between days 11 and 17. By the end of the experiment (i.e., 21 days), no significant difference between the tumor size of treated and control animals was...
and i.p. administrations at the times indicated by arrows were started. Control
saturation at 17 days, probably because of the cachexia induced in the
observed (Fig. SC). The size of tumors in untreated animals reached
as shown. Points, tumor volume (mm³); bars, SD.
animals (U) received vehicle. Treated animals received 25 (•), 50 (A), or 100 (0) mg/kg
and Methods." At day I, mice were separated into control (n = 10) and treatment (n = 5)
HCTI 16 (C) cells were implanted s.c. into nude mice at day 0 as described in "Materials
4 Unpublished observations.
We found that tumor inhibition in vivo was comparable to that of L739,749 (Ref. 21; data not shown). It is also
shown that tumor growth by this FTase inhibitor is associated with inhibition of ras posttranslational processing in the
tumor (Fig. 4). As it has been indicated by others (21), the use of FTase inhibitors appears to be safe and has potential for use as
long-term therapy: after 18 daily administrations of B1086, no signs of toxicity were evident.

The effect of B1086 or B956 on tumor growth was transient. Once the
drug was removed, the tumors resumed growth at approximately the
same rate as untreated controls, indicating a cytocstatic and not a cytotoxic
effect. However, the results with EJ-1 xenografts (Fig. 5A) suggest that
with a more optimized treatment strategy, the inhibition of tumor growth
could be irreversible and complete. In this case, a longer treatment
schedule was devised to accommodate the slow growth rate of EJ-1 tumors.
At the end of the experiment, one mouse in the higher dose group was
tumor free, whereas the remaining 4 animals did not show apparent tumor
growth. Although it is possible that tumors would grow during a longer
observation period, this result suggests that an adequate treatment sched-
ule might eliminate tumor growth completely. Further support for this
idea comes from the experiment where treatment of animals with a
growing tumor resulted in a significant reduction of tumor growth (Fig.
4A). In most of our experiments, drug treatment is started soon after
tumor implantation because under this condition tumors are more sensi-
tive. However, this experiment shows that the growth of an already
present tumor can be slowed down and possibly stopped by FTase
inhibitors using an appropriate dosage regimen.

One piece of puzzling information is derived from data with a panel
of human tumor cell lines expressing different ras oncogenes. When
these cell lines are treated with B956 in a colony-forming assay,
differences in sensitivities are observed. We have shown previously
that increasing concentrations of inhibitor are required to inhibit
posttranslational processing of different prenylated proteins (18).
However, in this case, differences in sensitivities are observed among
cell lines expressing the same ras oncogene. This was especially clear
for cells expressing the K-ras oncogene, where a large number of cell
lines was tested. It is possible that a wide range in sensitivity to FTase
inhibitors will be observed in cells with H- or N-ras mutations if more
cell lines are studied. On the other hand, whereas other reports have
shown lack of effect of FTase inhibitors on non-ras-transformed cell
(i.e., raf-, mos-, or src-transformed cell lines; Refs. 14 and 17),
the data presented here indicate that about 50% of the K-ras-trans-
formed cell lines tested are as sensitive as non-ras-transformed cell
lines. This discrepancy could perhaps be eliminated if more cell lines
are studied in different laboratories.

Available data indicate that if a cell line expresses a ras oncogene,
its growth is governed by the presence of activated ras (23). However,
tumor cell lines are characterized by the presence of multiple mutations in different genes. It is possible that unidentified mutations downstream of ras could maintain cell growth, even if ras signaling is disrupted by FTase inhibitors. Alternatively, mutations in pathways parallel to ras could be sufficient to maintain the transformed phenotype. The above alternatives might explain why a ras-transformed cell line could be resistant to growth inhibition by farnesylases.

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