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cenal cell carcinoma (RCC) accounts for 3% of all ma-
ignant tumors and is the sixth leading cause of cancer
defects in the United States. In 2007, an estimated
51,190 new cases of RCC will be diagnosed and 31,590
deaths will be attributed to renal cancer (Jemal et al.,
2007). The incidence of renal cancer at all stages has increased steadily
since 1973; unfortunately, increased detection of earlier-stage
disease has not coincided with a decrease in the number of pa-
tients diagnosed with advanced renal cancer (Hock, Lynch, &
Balaji, 2002). For patients with localized disease, nephrectomy
offered five-year survival rates from 90%–95% (Bui et al.,
2001), but median survival among patients with metastatic disease was
only 13 months (Cohen & McGovern, 2005).

Most renal cancers are sporadic: Risk factors include smoking
(associated with 24%–30% of all cases of RCC), obesity, sedentary
lifestyle, environmental and occupational exposure (e.g., asbes-
tos, cadmium, polycyclic hydrocarbons, solvents), and long-term
use of diuretics or phenacetin-containing analgesics (Linehan
et al., 2004). Patients with end-stage renal disease undergoing
dialysis, particularly those with cystic disease, also are at higher
risk for RCC (Denton et al., 2002). A small number of cases are
hereditary, associated either with the von Hippel-Lindau (VHL)
gene in clear cell renal cancer or the c-met gene on chromosome
7 in type 1 papillary renal cancer (Linehan et al.). Individuals with
VHL syndrome are at risk to develop tumors in multiple organs,
including several hundred clear cell tumors per kidney.

Pathogenesis of Renal Cancer

Loss of VHL gene function leads to increased expression of
genes associated with tumor growth and angiogenesis, especially

Diagnosis of renal cell carcinoma (RCC) frequently occurs at advanced stages, severely limiting the success of treatment,
and median survival is barely more than a year. Previously, treatment of renal cancer was limited to nephrectomy or im-
munotherapy (interleukin or interferon-α), which was effective in a small subset of patients but often was accompanied by
severe side effects. New orally administered targeted therapies have become available, offering broader benefits to patients
with advanced RCC. Sorafenib is an oral, multikinase inhibitor recently approved by the U.S. Food and Drug Administration
as treatment for advanced RCC based on its extension of median progression-free survival from 12–24 weeks. Oncology
nurses must ensure patient adherence and manage side effects of emerging treatments. This article reviews the manage-
ment of skin rash, hand-foot skin reaction, hypertension, diarrhea, and fatigue in patients receiving sorafenib. In addition,
a case study of a patient receiving sorafenib is presented.

At a Glance

- Sorafenib is the first tyrosine kinase inhibitor approved for
  the treatment of advanced renal cell carcinoma.
- Nursing assessment and interventions are critical for effective
  management of unique side effects, including hand-foot
  skin reaction.
- Effective side-effect management allows patients to main-
  tain therapeutic benefit and maximizes quality of life.

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