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## Studies on Mechanisms of Augmentation of Liver Regeneration by Cyclosporine and FK 506

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Evidence could not be found of immune modulation of liver regeneration. The powerful immunosuppressive drug FK 506, which augments the response after partial hepatectomy in normal rats, had the same effect in T cell-deficient nude rats. The cytotoxicity of natural killer cells in treated nude rats was not significantly changed by FK 506 therapy. However, the serum of FK 506-treated nude rats increased hepatocyte proliferation when added to third-party hepatocyte cultures, suggesting that FK 506 had induced a serum growth factor in the nude rats or had suppressed an inhibitory factor. A hypothesis was advanced that FK 506 (and cyclosporine) affects hepatic growth by nonimmunological pathways. (HEPATOLOGY 1991;14:140-143.)

An unusual quality of two powerful immunosuppressive agents, FK 506 and cyclosporine, is augmentation of the regeneration response after partial hepatic resection (1-4). Two alternate explanations have been advanced for this phenomenon, one of which is the altered immune modulation of the regeneration response by these immunosuppressive agents that inhibit T-lymphocyte activation (4). The alternate hypothesis is that these drugs influence growth control responses by nonimmunological pathways, involving as yet poorly understood second messenger processes, signal transduction processes or both (4-6).

In this study regeneration was studied in nude rats first, to see whether these T-lymphocyte-deficient animals had normal regeneration, and second, to determine whether FK 506, which is a specific inhibitor of T-helper-lymphocyte activation, augmented the regeneration response. Failure to do so would support the immune modulation hypothesis. In addition, natural

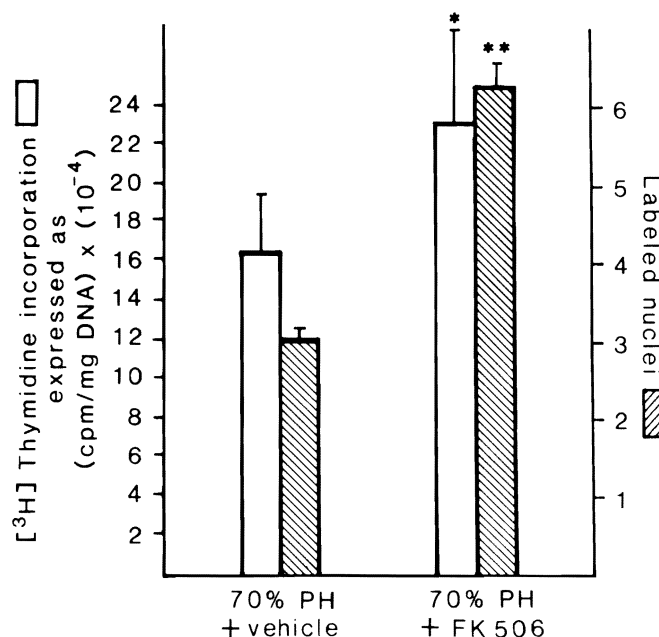


FIG. 1. [<sup>3</sup>H] thymidine incorporation and percentage of labeled hepatocyte nuclei 24 hr after 70% hepatectomy in nude (NIH-RNU) rats treated (right) or not treated (left) with FK 506. Mean ± S.E.M. (p < 0.05\* or <0.01\*\*). Ten animals in each group.

killer (NK) lymphocytes recently have been shown to have increased cytotoxicity against regenerating hepatocytes (7). Consequently, the effect of FK 506 on NK lymphocytes was determined in unoperated nude rats. The results of both kinds of experiments provided no evidence that either lymphocyte subpopulation plays a role in the control of regeneration.

### METHODS

#### *In Vivo Experiments*

Male nude rats (NIH-RNU, National Cancer Institute, Frederick, MD) were kept in pathogen-free temperature-controlled and light-controlled rooms and provided with food and water *ad libitum*. Before purchase, these rats were tested

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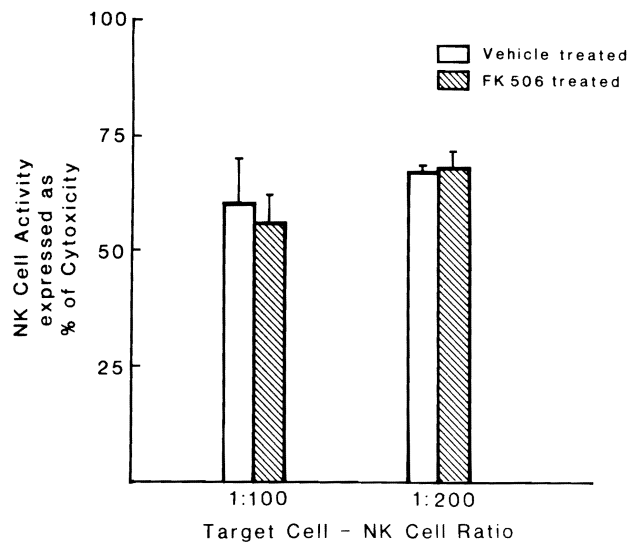


FIG. 2. Lytic activity of NK cells isolated from nude rats treated and untreated with FK 506. NK cells were added at a ratio of either 1:100 or 1:200 to target lymphoma cells. Each bar = four experiments. Mean  $\pm$  S.E.M.

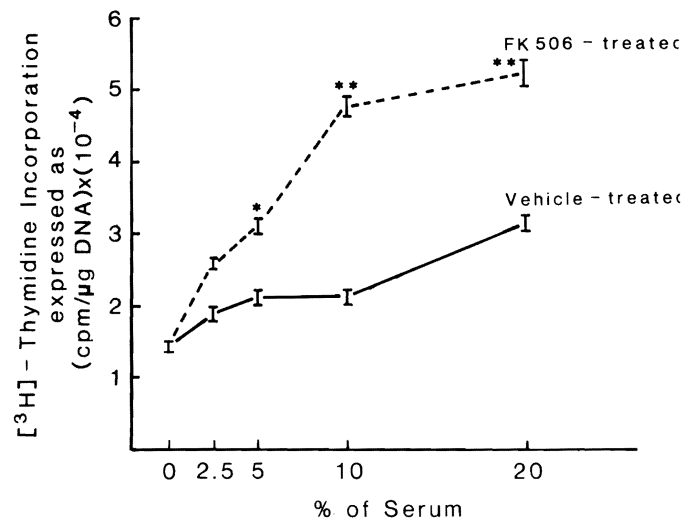


FIG. 3. Effect on DNA synthesis of cultured hepatocytes of different concentrations of rat serum obtained from nude rats treated or not treated with FK 506.  $n = 5$  in both experiments. ( $p < 0.05^*$  or  $< 0.01^{**}$ ).

TABLE 1. Experimental groups

Group	No. rats	Route	FK	Vehicle	Liver resection
1	10	IM	0	Saline	70%
2	10	IM	1 mg/kg	Saline	70%
3	4 <sup>a</sup>	IM	0	Saline	0 (no operation)
4	5 <sup>a</sup>	IM	1 mg/kg	Saline	0 (no operation)

<sup>a</sup>Serum and spleen were obtained at killing for *in vitro* experiments. IM = intramuscular.

by the National Cancer Institute and found to have absent T-lymphocytes and T-lymphocyte-mediated function. The nude rats were assigned to the four groups as shown in Table 1.

The 70% hepatectomies in groups 1 and 2 were performed with the rats under methopfan anesthesia between 7:30 and 9:00 AM by the method of Higgins and Anderson (8). Intramuscular drugs or saline vehicle was given daily during the three days before the surgery and just after completion of the operation. A further dose was given 24 hr after operation, 2 hr before killing. At the same time as the final dose, 50 mCi [<sup>3</sup>H-] thymidine/200 gm body wt was administered intraperitoneally. Liver DNA synthesis was judged by the incorporation of [<sup>3</sup>H-] thymidine and by the percent of labeled hepatocyte nuclei as previously described (4). The mortality rate was zero in treated and untreated groups.

### In Vitro Experiments

**NK Cell Activity.** In nude rats of groups 3 and 4 that had not been subjected to operation, the spleens were removed under sterile conditions at the time of death and NK cell preparations were made (9). Single cell suspensions prepared in RPMI 1640 in 5% FCS by this technique have viability exceeding 97% by trypan blue exclusion. These cells were tested for their ability to lyse the NK-sensitive Moloney virus-induced mouse T-cell

lymphoma YAC-1. The target tumor cell lines were grown in RPMI 1640 medium with 5% FCS and penicillin/streptomycin and subcultured two to three times per week until used.

The cytotoxicity of the NK cells was measured with a standard [<sup>51</sup>Cr]-release assay after labeling the target cells with 100 mCi of Na<sub>2</sub><sup>51</sup>CrO<sub>4</sub>/2 × 10<sup>6</sup> cells. NK cells were added to triplicate well round-bottom microplates (Costar, Cambridge, MA) that contained constant numbers of target cells to give an effector/target cell ratio of 1:100 or 1:200. After incubation at 37° C for 4 hr, the activity of the supernatant was determined in a gamma counter. Spontaneous isotope release (control) was obtained from wells receiving target cells and medium only, and total isotope release was obtained from wells receiving 1% Triton X-100 (Sigma Chemical Co., St. Louis, MO). The percent of cytotoxicity was calculated by: Percent cytotoxicity = 100 × [(Experimental release - Spontaneous release)/(Total release - Spontaneous release)].

**Serum Stimulatory Activity.** Sera at death were obtained from the FK 506-treated and untreated nonoperated nude rats in groups 3 and 4 and assayed for their hepatocyte stimulatory qualities. Target hepatocytes were isolated from 7-wk-old Fischer (F344) rats that weighed 180 to 200 gm. A modification of the two-step perfusion technique of Seglen (10) was used as described by Jirtle et al. (11). Viability was determined by trypan blue exclusion, and hepatocytes counted by hemocyt-

tometer were plated at a cell density of  $1.5 \times 10^5$ /well using 1.5 ml MEM + 5% FCS and  $10^{-7}$  molar insulin at 37° C in a 5% CO<sub>2</sub> atmosphere. Hepatocyte DNA synthesis was determined with [<sup>3</sup>H-] thymidine incorporation (4, 5). Serum or cellular proteins were determined by the method of Lowry et al. (12).

**Statistical Analysis.** Data were presented as mean  $\pm$  S.E.M. Unpaired Student's *t* test was used for statistical analysis of the data.

## RESULTS

### *In Vivo Experiments*

The nude rats not treated with FK 506 had a 16-fold increase in DNA synthesis and a commensurate increase of autographically labeled hepatocyte nuclei (Fig. 1). The regeneration response by both parameters was significantly greater when the animals received the 5-day course of preoperative treatment with FK 506 (Fig. 1).

### *In Vitro Experiments*

**NK Cell Activity.** The splenic NK cell cytolytic activity was vigorous when the cells were added at an effective target ratio of 100:1 or 200:1. This cytolytic activity was neither increased nor decreased by a 5-day course of FK 506 (Fig. 2), findings similar to those reported by Markus et al. (13) in normal Fischer rats.

**Hepatocyte Stimulatory Factors in Serum.** The serum of vehicle-treated unoperated nude rats (group 3) caused slight stimulation of F344 hepatocytes in culture (Fig. 3), an effect of normal rat serum that is well known (5, 14).

In contrast, the serum of unoperated nude rats submitted to a 5-day course of FK 506 (group 14) significantly increased the rate of replication when added to the hepatocyte culture medium. This was significant at all of the higher volume serum enrichments (Fig. 3). This stimulatory effect of the rat sera from FK 506-treated nude rats was similar to that previously seen in operated immunologically competent Fischer rats treated with FK 506 (5).

## DISCUSSION

These experiments suggest but do not prove that T helper or NK lymphocytes are unimportant in controlling liver regeneration, contrary to the speculation in the literature summarized elsewhere (4, 5, 7). In the nude rats in which the T cells were not present, the regeneration response was normal. Nevertheless, the FK 506, which targets T cells for its immunological action (15), augmented regeneration in the same way as in immunologically intact animals. The additional possibility (7) that NK cells were defunctionalized by the immunosuppressive therapy was essentially eliminated by our *in vitro* experiments and by *in vivo* experiments reported elsewhere (13).

Instead, the evidence, although circumstantial, is congruent with our previously enunciated hepatotrophic hypothesis that FK 506 and cyclosporine (16) affect basic growth control mechanisms that are independent

of immunological pathways. Both FK 506 and cyclosporine have cytosolic binding sites that, although distinct, contain peptidyl-prolyl isomerase (17, 18), an enzyme that facilitates the breakdown of oligopeptide bonds and facilitates protein folding (19). Cellular growth regulation as epitomized by its hepatotrophic qualities (6, 16; Francavilla A, Starzl TE, Porter K, et al. Manuscript submitted for publication, 1991), including regeneration (4, 5), is only one of the nonimmunological pleiotropic effects (20) of this ubiquitous immunophilin network after it binds FK 506 or cyclosporine (21).

The intriguing observation in our current experiments was the apparent development of hepatocyte stimulatory activity in the serum of FK 506-treated nude rats. The finding was similar to that recently reported in immunologically competent Fischer rats treated with this drug (5). At the moment, no clue exists as to the explanation for this induced stimulatory activity. The possibility has not been ruled out that what actually occurred instead of stimulation was removal or suppression of an inhibitory factor.

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