BIPHASIC CHANGE OF PROTON MAGNETIC RELAXATION TIMES DURING AZO-DYE HEPATOGENCY

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Summary.—For the first time, change in the proton longitudinal relaxation times ($T_1$) of rat tissues has been examined throughout the whole process of azo-dye hepatocarcinogenesis. Two maxima of the $T_1$ values were observed for liver, on Day 60 and after Day 120, and these changes correlated well with the changes in water content. The first peak was ascribed to the immature hepatocytes of hyperplastic nodules, and the second peak to the developed hepatoma cells. The significance of the change in $T_1$ values as a preneoplastic change is discussed.

Since the pioneering work of Damadian (1971), cancerous tissues have been well characterized by the prolonged proton longitudinal relaxation times ($T_1$s) of tissue fluid, when compared with normal tissues (Hollis et al., 1973). A number of papers have been published on this subject; in particular, Damadian et al. (1974) suggested the possibility of applying proton NMR for clinical use, to detect cancer at an early stage. Later studies revealed that the elevated $T_1$ of tumours could mostly be interpreted in terms of the increased water content of tissues (Inch et al., 1974a). Such an anomalous water content, however, may reflect an abnormal state of cell membrane, including Na$^+$K$^+$ATPase, which is closely correlated with cell growth (Kimelberg and Mayhew, 1975). In this regard, it is important to know how $T_1$ and water content may change during the course of chemical carcinogenesis. To this end, Floyd et al. (1975) carried out a feeding experiment with 3'-methyl-4-dimethylaminoazobenzene (3'-Me-DAB), and showed that the $T_1$ of blood serum and liver tissues of rats increased at 4 weeks. They ascribed the increased $T_1$ of liver tissues to the preneoplastic nature of hepatic nodules. However, in their experiment, only the early stage of hepatocarcinogenesis (i.e., 4 weeks) was covered, and a comparative study of the changes of $T_1$ and tissue histology was not made.

In the present work, correlation between the neoplastic change and $T_1$ of the liver, kidney, and blood serum of rats was studied for 150 days or more, and this is the first report of such studies covering the whole process of azo-dye hepatocarcinogenesis. Interestingly, we found that the $T_1$ and water content of liver tissues changed biphasically, corresponding to the formation of hyperplastic nodules and hepatoma.

MATERIALS AND METHODS

Male Sprague–Dawley rats weighing about 140 g were fed with either azo-dye diet (standard basal diet containing 0·06%, 3'-Me-DAB, prepared by CLEA Japan Inc., Tokyo) or a basal diet until Day 87. Thereafter, both groups received the basal diet. The incidence of hepatoma in azo-dye-fed rats was 80%, or more on Day 200. At 2-week intervals, 3 rats

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of each group were killed for the measurement of $T_1$ of liver, kidney, and serum. Two or 3 samples of the same tissues were examined for each rat. Blood was collected from the carotid artery under ether anaesthesia. Immediately after removal, liver and kidney tissues were cut into small pieces and blotted dry before placing in sample tubes. To avoid error due to heterogeneity of samples, a small volume of the tissues was packed into

![Graphs](image-url)

**Fig. 1.**—Changes of $T_1$ (A–C) and water content (D–F) of liver (A, D), kidney (B, E), and serum (C, F) during feeding of 3'-Me-DAB diet (×—×) and control diet (○—○). Means and s.d. ⊙ on Day 158 (A and D) indicates the value for the normal hepatic tissue adjoining the hepatoma. Each point represents data of 3–6 rats.
sample tubes of 5 mm outside diameter. In sampling tissues such as hyperplastic nodules or developed hepatoma, efforts were made to select the homogeneous portion. T1s were measured with a JEOL PFT-100 pulsed spectrometer operating at 100 MHz. Automatic T1 routine, as well as a JEOL PG-100 digital pulse programmer, was used to obtain T1 values automatically. Water content of tissues was determined by the difference in the weight before and after lyophilization of samples for 13 h or more. A portion of the tissues used for T1 measurement was always examined histologically, by staining with haematoxylin and eosin after fixation with 10% formaldehyde.

RESULTS AND DISCUSSION

Fig. 1, A, B, and C illustrates the changes in T1 of the liver, kidney, and blood serum, respectively, during the course of azo-dye hepatocarcinogenesis. The T1 of liver, the target organ of the azo dye, exhibited 2 maxima, on Days 60 and 160. Observation of the first maximum on Day 60 was further confirmed by conducting a second run of the feeding experiment. The T1s of kidney, a non-target organ, remained constant throughout. The T1 of blood serum of azo-dye-fed animals showed a change similar to those of liver tissues, but the pattern of change of T1 for control animals was quite different from that of liver. Thus, the T1 rose around Day 150, just as the case of treated animals. Although we have no explanation of this phenomenon, it might be that T1 of serum is especially variable and tends to fluctuate. In fact, standard deviation of serum T1 was too large to draw any conclusion. Fig. 1, D, E, and F indicates the change in water content of the same samples of liver, kidney, and blood serum used for the T1 measurements. As with the T1, the water content of liver showed 2 maxima, on Days 60 and 160. Clearly, the change of T1 in the course of azo-dye hepatocarcinogenesis is well correlated with the change of water content (Fig. 2). For the liver of azo-dye-fed rats, especially at the first peak (around Day 60), the standard deviation of T1 values and water content was not small, but the difference from that of the control liver was statistically significant (0.01 < P < 0.02 for T1 of liver on Day 60; 0.002 < P < 0.005 for water content of liver on Day 60, according to Student's t test). Such a fluctuation of data was mostly due to different responses of individual animals to 3'-Me-DAB. Instrumental error in determining T1 was estimated as 10%, at most, 15%. In the case of blood serum, error due to haemolysis may not be excluded completely, but it was not large.

In parallel with the 60 Day increase of T1, formation of hepatic nodules was observed macroscopically. Nodules had a solid, pale appearance, distinguished from normal hepatic regions. The histological examination confirmed regenerative nodules and adjoining cholangiolar (oval) cells (Farber, 1956). Fig. 1 shows that T1 decreases after the first peak. During this period, hyperplastic nodules were macroscopically obscured. Histologically, nodules still persist, yet proliferation of cholangiolar cells was not remarkable. The second increase in T1,
after Day 120, was accompanied by multiple formation of hepatoma, while normal hepatic regions adjoining hepatoma showed a low $T_1$ (Fig. 1 A). Thus, the change of $T_1$ accompanying the primary hepatoma was localized, not spreading to other normal tissues or organs. This is different from the situation in transplantable hepatoma (Hollis et al., 1974), in which the increased $T_1$ spread to a large extent to the surrounding normal liver tissues.

In respect of the early change of $T_1$, the present study confirmed the results of Floyd et al., (1975). Our new finding is that the elevated $T_1$ decreased before the appearance of hepatoma, thus resulting in a biphasic change in $T_1$. Detailed histological examinations revealed that the first peak of $T_1$ on Day 60 does not coincide with the disappearance of original hepatocytes, but with the proliferation of renewed hepatocytes. A similar biphasic pattern was also reported for the appearance of $\alpha$-foetoprotein in the course of azo-dye hepatocarcinogenesis (Watabe, 1971). Appearance of $\alpha$-foetoprotein, as well as increased $T_1$, is characteristic of proliferating immature hepatocytes, because both are shared, not only by hepatoma, but also by regenerating liver (Inch et al., 1974b; Abelev, 1968). These preneoplastic changes are reversible and transient in nature, which distinguishes them from irreversible and permanent changes such as for the isozyme pattern of aldolase (Endo et al., 1970).

In conclusion, hepatocarcinogenesis was characterized by 2 maxima of $T_1$, the first derived from preneoplastic hyperplasia and the second from genuine neoplasia. These 2 maxima must be distinguished carefully in the application of nuclear magnetic relaxation to clinical use.

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