All-trans-retinoic acid distribution and metabolism in vitamin A-marginal rats

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Cifelli, Christopher J., and A. Catharine Ross. All-trans-retinoic acid distribution and metabolism in vitamin A-marginal rats. Am J Physiol Gastrointest Liver Physiol 291: G195–G202, 2006; doi:10.1152/ajpgi.00011.2006.—Retinoids, including all-trans-retinoic acid (RA), are considered to have anti-inflammatory properties and are used therapeutically for diseases of the skin and certain cancers. However, few studies have addressed the effects of disease states on RA metabolism. The present study was conducted to better understand the effects of exogenous RA, both in the absence and presence of inflammation, on the distribution and metabolism of a dose of [3H]RA. Female Sprague-Dawley rats fed a low vitamin A diet were pretreated with RA (po), a low dose of lipopolysaccharide (LPS, ip), or their combination. Twelve hours later, albumin-bound [3H]RA was injected intravenously, and tissue organic- and aqueous-phase [3H] was determined after 10 and 30 min. In liver and plasma, [3H]-labeled organic metabolites (e.g., 4-oxo- and 4-hydroxy-RA) were isolated by solid-phase extraction. LPS-induced inflammation significantly reduced plasma retinol by 47%, increased total [3H] in plasma at 10 min, and reduced total [3H] in liver at both times. In contrast, RA pretreatment did not affect plasma retinol, significantly increased total [3H] in plasma at both times, and did not affect liver total [3H]. However, by 30 min, RA significantly increased [3H]RA metabolism in plasma, liver, lung, and small intestine, as indicated by greater [3H]-labeled aqueous-phase and [3H]-labeled organic-phase metabolites. The results presented here demonstrate that, although LPS-induced inflammation affects the organ distribution of RA, the ability of RA to induce its own catabolism is maintained during inflammation. Thus we conclude that RA and LPS act independently to alter RA metabolism in vitamin A-marginal rats.

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designed a 2 × 2 factorial study to examine the effects of RA alone, LPS-induced inflammation alone, and both in combination on RA homeostasis. The results of these studies demonstrate that RA and LPS independently affect the distribution and the metabolism of RA in a tissue-specific manner and that RA and LPS are, for the most part, independent factors in the regulation of RA homeostasis.

MATERIALS AND METHODS

Animals and diet. All animal procedures were approved by the Institutional Animal Use and Care Committee of the Pennsylvania State University. Female Sprague-Dawley (Charles River Laboratories, Boston, MA) rats with litters of 12 female pups were fed a vitamin A-deficient diet from the time of arrival to reduce the accumulation of vitamin A from milk by their nursing offspring (18). The rats were housed in a room maintained at 22°C with a 12:12-h light-dark cycle, and food and water were available ad libitum. When the pups were 21 days old, they were weaned in pairs and fed a nutritionally complete AIN-93G diet (45) modified to contain 0.412 mg of retinyl palmitate per kilogram of diet (Research Diets, New Brunswick, NJ). This amount of vitamin A was shown previously to be adequate for normal growth but results in plasma retinol levels about half that of control-fed rats (51).

Experimental design. When the rats were between 54 and 60 days of age, they were randomly assigned to one of four treatment groups (n = 9/group): control, RA alone, LPS alone, or RA + LPS. Rats treated with RA were given an oral dose of 50 μg all-trans-RA (Sigma-Aldrich, St. Louis, MO) prepared in 90 μl of vegetable oil (65), and all other rats received an equal amount of oil only. Rats treated with LPS received an intraperitoneal injection of 50 μg/100 g body wt (BW) of Pseudomonas aeruginosa-LPS (List Biological Laboratories, Campbell, CA) (50, 51), whereas all other rats received an equal volume of sterile phosphate-buffered saline (PBS). Food was removed immediately after treatment until the end of the 12-h experimental period (50). Twelve hours posttreatment, the rats were lightly anesthetized by isoflurane-oxygen inhalation, and 0.15 ml/100 g BW was evaporated under argon until the end of the 12-h experimental period. For each group of rats, the rats were killed either 10 min (n = 9/group) or 30 min (n = 4/group) postinjection by carbon dioxide asphyxiation. Blood was drawn from the inferior vena cava in heparinized syringes, and the liver, lung, spleen, and small intestine were rinsed and placed in a modified cell pellet apparatus (Supelco). After collection, the samples were dried to constant weight under nitrogen for storage at 80°C before subsequent analysis. Because RA is rapidly metabolized, it is important to collect tissues quickly (11, 47); thus each rat was treated on a defined schedule so that all tissues could be collected and frozen within 5 min or less after euthanasia. The 10- and 30-min time periods chosen for this experiment were based upon previous reports (25) and pilot work (data not shown) which showed that >95% of the [3H]RA dose was cleared from plasma within 10 min and that metabolism of [3H]RA to polar metabolites could be readily detected by 30 min.

Dose preparation. All procedures involving retinoids were carried out in either dim or yellow light. The doses of [3H]RA for intravenous injection were prepared fresh immediately prior to injection. To prepare a 1-ml aliquot, 10 μCi of [3H]RA ([11,12(N)-3H]retinoic acid, specific activity 1.96 TBq/nmol; from PerkinElmer, Boston, MA) was combined with 0.23 μmol of all-trans-RA, and the solvent was evaporated under argon until <10 μl remained. Next, 5 μl of Tween 20 (Sigma-Aldrich) and 0.9 ml of 0.1% rat serum albumin (diluted in sterile PBS; Calbiochem, San Diego, CA) were added, and the mixture was vortexed. Then, 0.10 ml of whole rat plasma was added to the sample and immediately vortexed. The final concentra-

tion of RA in the dose was 0.23 mM, and the dose delivered to each rat was standardized at 0.15 ml (0.33 nmol RA) per 100 g BW. Following the preparation and administration of each dose, aliquots were analyzed by HPLC (9), which showed them to be >97% [3H]RA (data not shown).

Determination of tissue radioactivity. Aliquots of individual plasma, liver, lung, and small intestine samples were extracted using a modification of the method of Folch et al. (15, 16). One gram of minced tissue, or 1 ml of plasma, was placed in 20 ml of chloroform: methanol (2:1, vol/vol) overnight and filtered the next day, and the filtrate was washed four times with 4 ml of the appropriate Folch wash solution. Following each wash step, the resulting aqueous phases were removed and combined. After the final wash, methanol (100%) was added to the aqueous phase until the total volume was 25 ml per sample. Next, the washes were back extracted with 10 ml of hexane, mixed, and centrifuged at 850 g for 5 min. The hexane was removed and added to the Folch organic phase. This backwash was performed to remove any nonpolar compounds present in the aqueous phase. The organic phase from each sample was dried under argon in an analytical evaporator (Organonation Associates, Berlin, MA) to complete-

ness, and the lipophilic compounds were reconstituted in 5 ml of chloroform. Aliquots of the injected dose along with the organic and aqueous phases from each tissue were analyzed by liquid scintillation spectrometry using a Beckman LS-3801 counter (Beckman Coulter, Fullerton, CA).

Separation of RA from polar organic metabolites (4-oxo- and 4-hydroxy-retinoic acid). Reverse-phase solid-phase extraction (SPE) was employed to separate the parent compound ([3H]RA) from its 3H-labeled oxidation products (mainly 4-oxo- and 4-hydroxy-RA) (17, 48) in the Folch organic phases of liver and pooled plasma samples. All-trans-RA and 4-oxo-RA standards (1.34 nmol each) were added to 500 μl (equal to ~100 mg tissue) of Folch-washed organic phase and dried to completeness under argon at −37°C. The residue was reconstituted in 750 μl of HPLC-grade acetonitrile:water (65:35, vol/vol) containing 10 mM acetic acid. Supelclean LC-18 SPE columns (Supelco, Bellefonte, PA) were conditioned with 8 ml of 100% methanol followed by 2 ml of water. After conditioning the columns, the samples were loaded on to the column. Using the all-trans-RA and 4-oxo-RA standards as monitor separation, polar metabolites were eluted first in 3 × 4 ml of acetonitrile:water (65:35, vol/vol), whereas all-trans-RA was eluted with 3 × 4 ml of acetonitrile:water (80:20, vol/vol). The elution rate was maintained at ~1 ml/min with negative pressure using an elution vacuum apparatus (Supelco). After collection, the samples were dried to completeness and analyzed by liquid scintillation spectrometry as described above.

In this paper, the various fractions are described as follows: [3H]RA, material in the organic phase that eluted from the SPE column with the all-trans-RA standard; [3H]RA organic metabolites, material into the organic phase that eluted from the SPE column with 4-oxo-RA: 3H-labeled organic-phase metabolites, material that partitioned into the organic phase during the Folch extraction; 3H-labeled aqueous-phase metabolites, material that partitioned into the aqueous phase during the Folch extraction; total 3H-labeled polar metabolites, the sum of the [3H]RA organic metabolites and 3H-labeled aqueous-phase metabolites; and total 3H-labeled retinoids, the sum of 3H-labeled organic-phase metabolites and 3H-labeled aqueous-phase metabolites.

Statistics. All results are means ± SE. Two-way ANOVA was used to test for main effects and interaction of RA treatment and LPS treatment on [3H]RA metabolism. Significant differences found by ANOVA were then tested using the Fisher protected least-squares difference test to determine which groups differed significantly (P < 0.05) from one another.
RESULTS

Inflammation- and RA-induced changes in organ weight and plasma retinol. Rats grew normally and appeared in good health, and groups did not differ in body weight at the time of treatment (Table 1). Twelve hours later, after treatment and an overnight fast, all groups lost ≈8.2 g BW (Table 1). There were no differences due to treatment in the relative weights of the lungs and small intestine. The relative weight of the spleen was higher in LPS-treated rats, indicative of mild inflammation, and the relative weight of the liver of rats treated with RA + LPS was slightly increased (Table 1). There was no difference in liver vitamin A, which averaged from 17 to 21 nmol/g (Table 1). Plasma retinol (Table 1) was measured as an indicator of vitamin A status and to confirm the effect of LPS-induced inflammation on plasma retinol. Plasma retinol averaged 0.54 μM in the control group, which was similar to previous studies of rats with marginal vitamin A deficiency (51), but it was only 42% of that in vitamin A-sufficient rats (50). Plasma retinol was further reduced by 47% in rats with LPS-induced inflammation compared with the marginally vitamin A-deficient control group, whereas RA treatment did not affect plasma retinol in this 12-h study (Table 1). There was no difference due to time (10 min vs. 30 min) in the sum of the total Δ3H-labeled retinoids recovered in the plasma, liver, lung, and small intestine (data not shown). However, LPS decreased the sum of the total Δ3H-labeled retinoids recovered in the plasma, liver, lung, and small intestine compared with RA-treated rats (Table 1).

RA alone and in combination with LPS increases the total radioactivity in plasma and the proportion of Δ3H[RA] metabolites. In control rats, greater than 95% of the injected dose was eliminated from plasma within 10 min (Fig. 1A). In comparison, total Δ3H-labeled retinoids were higher in plasma in the RA, LPS, and RA + LPS groups at this time. Using two-way ANOVA, we showed both RA and LPS to be significant factors, but there was no interaction. By 30 min, total Δ3H-labeled retinoids in plasma were similar in the control group and the LPS-treated group but were higher in the RA group (Fig. 1B). Furthermore, rats treated with RA + LPS had significantly more (8.5%) of the injected Δ3H dose in plasma, which was approximately twice the amount in the control group (P < 0.01).

Total Δ3H-labeled polar metabolites in plasma, representing lipid-soluble metabolites of RA and its aqueous metabolites combined, were determined on a pooled sample from each treatment group at each time. Δ3H-labeled polar metabolites were increased by RA, compared with the control group, at both 10 and 30 min. At 10 min, about 60% of the total Δ3H-labeled retinoids recovered in plasma already consisted of

Table 1. Body weights, relative organ weights, and plasma retinol concentrations in RA- and LPS-treated rats

<table>
<thead>
<tr>
<th></th>
<th>Control</th>
<th>RA</th>
<th>LPS</th>
<th>RA + LPS</th>
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</thead>
<tbody>
<tr>
<td>BW at dosing, g</td>
<td>233.8±9.80</td>
<td>224.2±8.67</td>
<td>212.9±5.74</td>
<td>217.9±7.36</td>
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<tr>
<td>Change in BW, g/12 h</td>
<td>−6.8±1.0</td>
<td>−6.4±0.9</td>
<td>−8.0±1.0</td>
<td>−8.2±1.0</td>
</tr>
<tr>
<td>Liver weight, % BW</td>
<td>3.64±0.06</td>
<td>3.76±0.12</td>
<td>3.91±0.11</td>
<td>4.03±0.10</td>
</tr>
<tr>
<td>Lung weight, % BW</td>
<td>0.51±0.02</td>
<td>0.53±0.01</td>
<td>0.54±0.02</td>
<td>0.53±0.02</td>
</tr>
<tr>
<td>Spleen weight, % BW</td>
<td>0.22±0.01</td>
<td>0.22±0.01</td>
<td>0.27±0.01</td>
<td>0.29±0.01</td>
</tr>
<tr>
<td>Small intestine weight, % BW</td>
<td>2.29±0.09</td>
<td>2.41±0.09</td>
<td>2.43±0.13</td>
<td>2.43±0.14</td>
</tr>
<tr>
<td>Plasma retinol, μmol/l</td>
<td>0.54±0.06</td>
<td>0.44±0.06</td>
<td>0.30±0.04</td>
<td>0.29±0.04</td>
</tr>
<tr>
<td>Liver total retinol, nmol/g*</td>
<td>17±11</td>
<td>21±6</td>
<td>18±5</td>
<td>14±17</td>
</tr>
<tr>
<td>Total Δ3H-labeled retinoids, % ID†</td>
<td>30.0±1.23</td>
<td>33.3±1.38</td>
<td>26.5±1.16</td>
<td>28.1±1.15</td>
</tr>
</tbody>
</table>

Values are means ± SE; n = 9/group. RA, retinoic acid; LPS, lipopolysaccharide; BW, body wt. *For liver total retinol, values are means ± range of two representative rats per group. †For total Δ3H-labeled retinoids, values are sums of plasma, liver, lung, and small intestine total Δ3H-labeled retinoids, expressed as percent injected dose (ID). aStatistically different from control rats (P < 0.05). bStatistically different from RA-treated rats (P < 0.05).

Fig. 1. Retinoic acid (RA) and lipopolysaccharide (LPS) increase both the total radioactivity and amount of Δ3H-labeled metabolites present in the plasma at 10 and 30 min postinjection. Female Sprague-Dawley rats were treated with RA, LPS, or the combination of RA + LPS as described in MATERIALS AND METHODS, and [3H]RA (0.23 mM) bound to rat serum albumin was injected 12 h later. Tissues were collected 10 (n = 5) or 30 (n = 4) min later, and total plasma radioactivity (A, 10 min and B, 30 min) and [3H]RA and Δ3H-labeled metabolites (C, 10 min and D, 30 min) were determined by liquid scintillation spectrometry. Bars are means ± SE. Different letters above bars within panels indicate significant differences (P < 0.05, a < b < c).
Fig. 2. RA and LPS independently affect both the total radioactivity and amount of $^3$H-labeled metabolites present in the liver at 10 and 30 min postinjection. Liver was collected 10 ($n = 5$) or 30 ($n = 4$) min after injection of $[^3$H$]$RA and organic-phase and aqueous-phase radioactivity ($[^3$H$]$RA; 10 min; and $B$, 30 min), along with $[^3$H$]$RA and $^3$H-labeled metabolites ($C$, 10 min; and $D$, 30 min) was determined by liquid scintillation spectrometry. Bars are means ± SE. Different letters above bars within panels indicate significant differences ($P < 0.05$, a < b < c).

$^3$H-labeled polar metabolites, which increased to 75% at 30 min (Fig. 1, $C$ and $D$). In the plasma of LPS-treated rats, $^3$H-labeled polar metabolites were elevated 10 min postinjection but equal to the control level by 30 min. In rats treated with the combination of RA + LPS, the proportion of $^3$H-labeled polar metabolites was highest (Fig. 1, $C$ and $D$), representing ~68% and 82% of total $^3$H-labeled retinoids at 10 and 30 min, respectively. Thus RA and the combination of RA + LPS resulted in a two- to threefold increase in plasma total $^3$H, compared with the control group, and the increase was mostly in the form of polar metabolites.

LPS and RA independently affect the total amount and distribution of $^3$H-labeled retinoids in liver. At 10 min postinjection, the liver of control rats contained ~24% of the injected dose as total $^3$H-labeled retinoids (Fig. 2A). The liver of LPS- and RA + LPS-treated rats contained less of the total dose, suggesting that LPS-induced inflammation may delay the uptake of $[^3$H$]$RA by the liver. By 30 min, total $^3$H-labeled retinoids fell by 4.3% in the control group (Fig. 2B), and a similar reduction was observed in each treatment group (Fig. 2, $A$ and $B$). Using two-way ANOVA, we showed that LPS reduced the total $^3$H-labeled retinoids at both 10 and 30 min (Table 2; Fig. 2, $A$ and $B$).

The proportion of unmetabolized $[^3$H$]$RA and $^3$H-labeled polar retinoids was determined on liver samples from individual rats. LPS reduced the amount of unmetabolized $[^3$H$]$RA in liver at both 10 and 30 min (Fig. 2C; Table 2). Although there was no effect of RA on unmetabolized $[^3$H$]$RA in liver at 10 min, unmetabolized $[^3$H$]$RA was much lower in the liver of all RA-treated rats (RA alone and RA + LPS), compared with the control group, at 30 min (Fig. 2D; Table 2). The percent of total $^3$H-labeled polar metabolites did not differ among treatments at 10 min (Table 2), although the ratio of total $^3$H-labeled polar metabolites to $[^3$H$]$RA was greater in all RA- and LPS-treated rats compared with the control group. By 30 min, total $^3$H-labeled polar metabolites increased in the liver of all RA-treated rats, regardless of inflammation (Fig. 2D; Table 2). At this time, total $^3$H-labeled metabolites represented ~82% of the total radioactivity present in the liver in both of these groups. Thus, although LPS significantly affected the total amount of $^3$H present in the liver at both times, only RA affected the amount of $^3$H-labeled polar metabolites in the liver (Table 2).

### RA metabolism in extrahepatic tissues

Next, we examined two extrahepatic tissues, the lungs and small intestine, both of which are intimately involved in vitamin A homeostasis (53, 54), to determine their response to RA- and LPS-induced inflammation (Fig. 3). Because of the relatively small amount of total $^3$H recovered in the lungs and small intestine (see below), total $^3$H-labeled retinoids were only partitioned into an organic phase and an aqueous phase; thus the organic phase may contain unmetabolized $[^3$H$]$RA and $^3$H-labeled polar metabolites, whereas all $^3$H in the aqueous phase has undergone metabolism. Less than 1% of the injected dose of $[^3$H$]$RA was recovered in the lungs of control rats at either time. However, this still represented an enrichment of $1.34 \pm 0.12$ and $1.25 \pm 0.05$ at 10 and 30 min, respectively, based on the relative weight of the lung. RA treatment did not affect total $^3$H-labeled

<table>
<thead>
<tr>
<th>Organ</th>
<th>10 min</th>
<th>30 min</th>
</tr>
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<tbody>
<tr>
<td></td>
<td>RA</td>
<td>LPS</td>
</tr>
<tr>
<td>Liver</td>
<td>$0.23$</td>
<td>$0.002$</td>
</tr>
<tr>
<td>Organic-phase $^3$H</td>
<td>$0.03$</td>
<td>$0.0001$</td>
</tr>
<tr>
<td>Aqueous-phase $^3$H</td>
<td>$0.004$</td>
<td>$0.05$</td>
</tr>
<tr>
<td>$[^3$H$]$RA</td>
<td>$0.04$</td>
<td>$0.0002$</td>
</tr>
<tr>
<td>$^3$H-labeled polar metabolites†</td>
<td>$0.46$</td>
<td>$0.32$</td>
</tr>
<tr>
<td>Lung</td>
<td>$0.63$</td>
<td>$0.01$</td>
</tr>
<tr>
<td>Organic-phase $^3$H</td>
<td>$0.03$</td>
<td>$0.23$</td>
</tr>
<tr>
<td>Aqueous-phase $^3$H</td>
<td>$0.009$</td>
<td>$0.06$</td>
</tr>
<tr>
<td>Small intestine</td>
<td>$0.16$</td>
<td>$0.0004$</td>
</tr>
</tbody>
</table>

$^P$ value obtained via two-way ANOVA. *Amount of $^3$H that eluted with an all-trans-RA standard during solid-phase extraction (SPE). †Amount of $^3$H that eluted with a 4-oxo-RA standard during SPE.
The amount of LPS administered (50 μg/100 g BW) was similar to that used in previous experiments to elicit a moderate inflammatory response in rats, which included a significant reduction of ~7.5% in body weight over 24 h, a moderate increase in body temperature (0.6°C), and significant hyporeti-
nolemia (50, 51). However, the LPS treatment protocol we have used does not induce endotoxemia or septic shock, as are induced by 10- to 20-fold higher doses of LPS (24, 29). Inflammation-induced hyporeti-
olemia was confirmed in our experiment by a 47% reduction in plasma retinol. Inflammation was also evident by a small but significant increase in relative spleen weight (Table 1), which is consistent with immune cell activation by LPS as shown in other studies (36, 64). The dose of RA used in this study was shown previously to increase the expression of several retinoid-responsive genes in both vitamin A-deficient and vitamin A-sufficient rats (65) and, therefore, would be expected to have similar effects in vitamin A-marginal rats. By examining the distribution of intravenously administered [3H]RA in the plasma, liver, and extrahepatic organs of vitamin A-marginal rats, we determined that [3H]RA is cleared from plasma and metabolized very rapidly and that each organ differs in its pattern of RA metabolism. RA and LPS each significantly modified the metabolism of [3H]RA. However, for the most part, RA and LPS were independent determinants of RA metabolism.

**Plasma retinoid metabolites are predominantly regulated by RA.** Reports prior to this study had established that under normal physiological conditions, the plasma pools of RA turn over rapidly in both rats and humans (1, 25, 37, 58). Several investigators have shown that following the clearance of all-
trans-RA from plasma, this retinoid subsequently appears as various polar metabolites, including 4-hydroxy-RA, 4-oxo-RA, and retinoyl β-glucuronide (11, 17, 57). In the present study, greater than 95% of the injected dose of [3H]RA was cleared from plasma within 10 min, confirming the rapid clearance of RA from plasma observed in rats and other species. Although treatment with RA resulted in a higher percentage of [3H] in plasma, especially at 30 min, analysis of retinoids after reverse-
phase SPE showed that the increase is accounted for by the
RA-treated rats, regardless of concurrent inflammation. Labeled retinoid in our LPS-treated rats. It is evident that plasma \([3H]\)RA was maintained lower at 10 min than at 10 min in all four groups (Fig. 2). This represents an enrichment of 6.48 ± 0.50 based on the relative weight of the liver. The amount of total \(^3\)H-labeled retinoids recovered in liver was lower at 30 min than at 10 min in all four groups (Fig. 2). This reduction could be due in part to the loss from liver back into plasma (14, 34). Similarly, LPS administration reduces the amount of aqueous-phase radioactivity in both organs (Figs. 3 and 4). In contrast, only RA administration increased the amount of aqueous-phase radioactivity at 10 and 30 min in both organs, irrespective of LPS treatment. The increased aqueous-phase radioactivity could be caused by enhanced RA oxidation because both lung and small intestine contain members of the cytochrome P450 family capable of oxidizing RA (27, 30). Interestingly, in the small intestine, the amount of aqueous-phase radioactivity increased in each treatment group over time, suggesting the presence of biliary metabolites of RA in the small intestine by 30 min (11, 57, 62).

In conclusion, the results of our study demonstrate that LPS-induced inflammation affects the organ distribution of RA, which may reflect differences in either the rate of uptake or the recycling of RA in a tissue-specific manner. The greatest effect was observed in the liver, where LPS reduced the amount of radioactivity recovered by nearly 30%. However, despite differences in the initial distribution of RA, LPS did not affect the total amount of \(^3\)H-labeled metabolites present in the liver, lung, and small intestine by 30 min. The interval from 10 to 30 min may have been sufficient for all tissues to metabolize most of the RA into metabolites. In contrast, although RA administration did not influence tissue radioactivity levels, it did cause a significant increase in the recovery of \(^3\)H-labeled metabolites, irrespective of LPS. Taken together, our results suggest that acute inflammation can alter systemic RA homeostasis by disrupting the normal physiological distribution of RA. Nevertheless, the moderate degree of inflammation induced in our rat model did not greatly compromise the metabolism of \([3H]\)RA, whereas the conversion of \([3H]\)RA to \(^3\)H-labeled metabolites was significantly increased after treatment with RA.

**Inflammation-induced changes in hepatic \([3H]\)RA metabolism.** The liver is directly involved in the maintenance of retinol homeostasis by regulating its storage, mobilization, and utilization (52). Previous work has established that hepatic retinol mobilization is impaired during acute inflammation (19, 50). Since RA mediates most of the physiological functions of vitamin A, we were interested in understanding the effects of LPS-induced inflammation on the uptake and metabolism of \([3H]\)RA by liver. Significant amounts of RA are taken up by liver from plasma and presumably utilized or recycled (25). On average, 24% of the injected dose was recovered in the liver of control rats at 10 min (Fig. 2A). This represents an enrichment of 6.48 ± 0.50 based on the relative weight of the liver. The amount of total \(^3\)H-labeled retinoids recovered in liver was lower at 30 min than at 10 min in all four groups (Fig. 2). This reduction could be due in part to the loss from liver back into plasma of the parent compound \([3H]\)RA or more polar metabolites such as retinoyl β-glucuronide (2). Alternatively or additionally, the decline in liver radioactivity could be caused by the formation and excretion of polar and aqueous-phase metabolites of RA, which appear in the bile of normal rats 2 h after RA administration (11, 57, 62). In our study, LPS significantly reduced the total amount of \(^3\)H present in liver at both 10 and 30 min (Fig. 2, A and B). We speculate that the observed reduction in liver radioactivity could be a consequence of three primary causes. First, LPS-induced inflammation increases vascular permeability, which could alter organ blood flow and thus influence the uptake, utilization, and recycling of RA by the liver (7, 13). Second, acute inflammation is known to diminish the expression of fatty acid transport protein (FATP) and fatty acid translocase (FAT), both of which are involved in the uptake of long-chain fatty acids from plasma (14, 34). Similarly, LPS administration reduces the expression of cellular fatty acid binding protein, L-FABP (35). While it is currently believed that RA readily diffuses across plasma membranes in a nonsaturated manner, the effects of inflammation on different fatty acid transporters suggest that these or similar changes could affect RA homeostasis as well. Third, specific transporters involved in the uptake and secretion of chemicals by the liver are altered following LPS administration (5). Thus vascular alteration coupled with changes in membrane-bound transporters involved in fat and xenobiotic homeostasis, known to be affected by LPS, may have contributed to the LPS-induced decrease in liver \(^3\)H-labeled retinoid in our LPS-treated rats.

**RA-induced changes in hepatic \([3H]\)RA metabolism.** Pharmacokinetic studies have determined that long-term administration of RA results in a decrease in plasma RA and subsequent increase in RA metabolism (42, 46, 58). Specific cytochrome P450 enzymes are induced in response to high cellular concentrations of RA, which may work in concert to regulate the degradation of RA and to return the concentration of RA to normal cellular levels (17, 28, 48, 49, 65, 66). In our studies, pretreatment of marginally vitamin A-deficient rats with oral RA significantly increased the amount of \(^3\)H-labeled metabolites present in the liver. Furthermore, this induction in RA metabolism occurred irrespective of LPS administration. The inability of LPS to affect the amount of \(^3\)H-labeled metabolites in the liver was surprising since the detoxification capacity of the liver for drugs and xenobiotics is diminished during inflammation (20, 21), evidenced by an LPS-induced reduction in both the expression and enzymatic activity of various members of the cytochrome P450 family (56). Thus the results presented here suggest that the ability of RA to induce its own catabolism is maintained during periods of physiological stress, which would serve to prevent the elevation of RA concentrations when acute or chronic RA treatment is used therapeutically as an adjuvant or to treat different disease conditions.
ACKNOWLEDGMENTS

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