Two Carotenoid Oxygenases Contribute to Mammalian Provitamin A Metabolism**

Received for publication, July 11, 2013, and in revised form, October 7, 2013. Published, JBC Papers in Press, October 8, 2013, DOI 10.1074/jbc.M113.501049

Jaume Amengual, M. Airanthi K. Widjaja-Adhi, Susana Rodriguez-Santiago, Susanne Hessel, Marcin Golczak, Krzysztof Palczewski, and Johannes von Lintig †, ‡
From the Department of Pharmacology, Case Western Reserve University School of Medicine, Cleveland, Ohio 44106

Mammalian genomes encode two provitamin A-converting enzymes as follows: the β-carotene-15,15'-oxygenase (BCO1) and the β-carotene-9',10'-oxygenase (BCO2). Symmetric cleavage by BCO1 yields retinoids (β-15'-apocarotenoids, C20), whereas eccentric cleavage by BCO2 produces long-chain (>C20) apocarotenoids. Here, we used genetic and biochemical approaches to clarify the contribution of these enzymes to provitamin A metabolism. We subjected wild type, Bco1<sup>−/−</sup> and Bco2<sup>−/−</sup> <span>double knock-out mice to a controlled diet providing β-carotene as the sole source for apocarotenoid production. This study revealed that BCO1 is critical for retinoid homeostasis. Genetic disruption of BCO1 resulted in β-carotene accumulation and vitamin A deficiency accompanied by a BCO2-dependent production of minor amounts of β-apo-10'-carotenol (APO10ol). We found that APO10ol can be esterified and transported by the same proteins as vitamin A but with a lower affinity and slower reaction kinetics. In wild type mice, APO10ol was converted to retinoids by BCO1. We also show that a stepwise cleavage by BCO2 and BCO1 with APO10ol as an intermediate could provide a mechanism to tailor asymmetric carotenoids such as β-cryptoxanthin for vitamin A production. In conclusion, our study provides evidence that mammals employ both carotenoid oxygenases to synthesize retinoids from provitamin A carotenoids.

Vitamin A (all-trans-retinol, ROL) is critical for vision, embryonic development, cellular homeostasis, and immunity.

† To whom correspondence should be addressed: Dept. of Pharmacology, Albert Einstein College of Medicine, Bronx, NY 10461.
‡ John H. Hord Professor of Pharmacology.

The abbreviations used are: ROL, all-trans-retinol; APO10ol, β-apo-10'-carotenol; BC, β-carotene; BCO1, β-carotene-15,15'-monoxygenase 1; BCO2, β-carotene-9',10'-dioxygenase; ko/kO, BCO1<sup>−/−</sup>; Bco2<sup>−/−</sup> double mutant mice; RA, all-trans-retinoic acid; RBP, retinol-binding protein; hRBP, human RBP; RE, retinyl ester; RPE, retinal pigment epithelium; LRAT, lecithin:retinol acyltransferase; Tricine, N-[2-hydroxy-1,1-bis(hydroxymethyl)ethyl]glycine; BisTris, [2-(bis(2-hydroxyethyl)amino]-2-(hydroxymethyl)propane-1,3-diol; Bistris propane, 1,3-bis[(hydroxymethyl)amino]propane; RAR, retinoic acid receptor; VAD, vitamin A-deficient; VAS, vitamin A-sufficient; TLCK, tosyllysine chloromethyl ketone; qRT, quantitative RT.

Significance: Carotenoids are the major source for vitamin A in the human diet.

Background: Mammalian genomes encode two carotenoid oxygenases, but their contributions to vitamin A homeostasis remain undefined.

Results: Mammals employ symmetric and eccentric cleaving carotenoid oxygenases to convert different provitamin A carotenoids to vitamin A.

Conclusion: Both carotenoid oxygenases contribute to vitamin A production.

© 2013 by The American Society for Biochemistry and Molecular Biology, Inc. Published in the U.S.A.
uptake of vitamin A into peripheral tissues, an RBP receptor was identified to be encoded by the stimulated retinoic acid 6 gene \( \text{Stra6} \) (21).

Biochemical studies indicate that mammals also can convert BC into \( \beta \)-apocarotenoids differently from retinoids (22–24). This eccentric cleavage of BC was proposed as an alternative method for producing vitamin A or its biologically active derivative, RA (25). Recently, \( \beta \)-apo-14-apocarotenal and \( \beta \)-apo-13-apocarotenone also have been implicated as selective antagonists of RARs (26, 27). Mammalian genomes encode a second putative BC-metabolizing enzyme (28). In contrast to BCO1, this \( \beta \)-carotene-9',10'-oxygenase (BCO2) also can metabolize non-provitamin A compounds such as xanthophylls (29–31). BCO1 and BCO2 are localized in different intracellular compartments. BCO1 is a cytoplasmic protein, whereas BCO2 is found in mitochondria (17, 29). These differences in cellular localization may determine the substrate accessibility and reflect different physiological functions adopted by BCO1 and BCO2.

Whether BC can be metabolized by two different enzymes is of general interest for human health. Considering the vitamin A deficiency problem, it is clearly important to know what proportion of ingested BC can be converted to retinoids. Furthermore, the proposal that \( \beta \)-apocarotenoids function as naturally occurring retinoid antagonists has critical implications for the biological activities of BC as a provitamin (26). Thus, to study the role of the two carotenoid-cleaving enzymes in BC metabolism, we took advantage of previously generated knock-out mouse models for both enzymes (29, 32). Using single as well as compound mutants, we genetically dissected the function of these enzymes for provitamin A metabolism. Moreover, we analyzed the metabolic fate of apocarotenoids by performing both in vivo and in vitro experimental studies.

**EXPERIMENTAL PROCEDURES**

**Animals, Husbandry, and Experimental Diets**—Animal procedures and experiments were approved by the Case Western Reserve University Animal Care Committee and conformed to recommendations of both the American Veterinary Medical Association Panel on Euthanasia and the ARVO Statement for the Use of Animals in Ophthalmic and Vision Research. In all experiments, mice were maintained at 24°C in a 12:12-h light/dark cycle and had free access to food and water. The generation of Bco1\(^{-/-}\), Bco2\(^{-/-}\), and Lrat\(^{-/-}\) mice has been previously described (29, 32, 33). Double knock-out Bco1\(^{-/-}\); Bco2\(^{-/-}\) (ko/ko) mice were established by conventional cross-breding. Female Bco1\(^{-/-}\), Bco2\(^{-/-}\), ko/ko, and wild type (WT) control mice with a C57/BL6;129Sv mixed genetic background were used for the described experiments. During breeding and weaning periods (up to 5 weeks of age), all mice were maintained on breeder chow containing ~29,000 IU vitamin A/kg diet (Prolab RMH 3000). For BC intervention, mice then were fed a pelleted diet based on the AIN-93 formulation containing 50 mg of BC per kg dry pellet for 10 weeks as the sole source for vitamin A (data from this feeding study are presented in Figs. 1, 2, 5, and 6). As a control, female WT and Bco2\(^{-/-}\) mice \((n = 5)\) each were subjected to same diet for 10 weeks supplemented with 4000 IU of vitamin A instead of BC (experiment is presented in Fig. 7). The diet was prepared by Research Diets, Inc. (New Brunswick, NJ), by cold extrusion to protect BC from heat and incorporated a water-soluble formulation of beadlets (DSM Ltd., Sisseln, Switzerland). Because a previous study reported significant amounts of apocarotenoids in BC beadlets (34), we analyzed the diet for the presence of these compounds. However, \( \beta \)-apo-10'-carotenal content was below the detection limit (~1 pmol) of our LC-MS system (data not shown). Weight gain and food intake of mice were measured as described previously (35). After 10 weeks of dietary intervention, mice were anesthetized by intraperitoneal injection of a mixture containing ketamine (80 mg/kg body weight) and xylazine (20 mg/kg body weight) in 10 mM sodium phosphate, pH 7.2, with 100 mM NaCl, and blood was drawn directly from the heart after snipping the right atrium. Then mice were perfused with 10 ml of PBS (137 mM NaCl, 2.7 mM KCl, 4.3 mM NaH2PO4, 1.4 mM KH2PO4, pH 7.3) and sacrificed by cervical dislocation before tissue collection.

For studying esterification, 12-week-old Lrat\(^{-/-}\) and WT mice were gavaged with a pharmacological dose of \( \beta \)-10'-apo-carotenal (20 mg/kg of body weight) dissolved in canola oil as a vehicle; prior to this, mice were raised on a standard chow diet containing ~29,000 IU vitamin A/kg diet (Prolab RMH 3000) (the experiment is presented in Fig. 3). For assessing hepatic RBP release, 12-week-old female Bco1\(^{-/-}\) mice were fed a diet based on the AIN-93 formulation without vitamin A (Research Diets). After 5 weeks, mice were treated either with pharmacological doses of ROL (Sigma) or \( \beta \)-apo-10'-carotenol (APO10ol) (each at 30 mg/kg of body weight) dissolved in canola oil. Four hours after treatment, mice were anesthetized and perfused with PBS, and tissues were harvested as described above. For studying \( \beta \)-cryptoxanthin metabolism, mice were raised on KLIBA 3430 chow (Provimi Kliba AG, Kaiseraugst, Switzerland) containing 14 IU/g vitamin A for 12 weeks and were injected with pharmacological doses of \( \beta \)-cryptoxanthin (20 mg/kg dissolved in DMSO) three times in 24-h intervals. Twenty four h after the last injection, mice were sacrificed, and livers were removed for further analysis (experiments are presented in Fig. 10).

**Analysis of RBP Secretion by HepG2 Cells**—Human HepG2 cells were maintained in DMEM supplemented with 10% fetal bovine serum (FBS) and 1% penicillin/streptomycin sulfate and cultured at 37°C in 5% CO2. When cells reached confluence, the media were removed, and cells were washed twice with PBS. To mimic either vitamin A deficiency (VAS) or vitamin A sufficiency (VAS), cells were incubated with DMEM containing 14 IU/g vitamin A for 24 h. The amount and chemical form of vitamin A added to the media were chosen based on retinoid analyses of three different batches of FBS routinely used in our laboratory. After 48 h of incubation, medium was replaced with medium containing either ROL or APO10ol at a final concentration of 4 \( \mu \)M for 2 h. Ethanol (up to 0.1% v/v) was used as the vehicle control. Cells were harvested and frozen at −80°C until further analyses. For immunocytochemistry, HepG2 cells were seeded on coverslips after using the same treatment and media conditions described above. At collection time, the media were removed, and cells were imme-

**Carotenoid Oxygenases and Provitamin A Metabolism**
Carotenoid Oxygenases and Provitamin A Metabolism

diately fixed with 4% paraformaldehyde in PBS for 1 h. Then cells were washed in PBS with 0.1% Triton X-100 (Roche Diagnostics) (PBS-T) and blocked for another hour with PBS-T supplemented with 10% BSA and 5% goat serum (Sigma). Afterward, cells were incubated overnight in blocking buffer containing a rabbit anti-human RBP serum (DakoCytomation, Glostrup, Denmark) diluted 1:200 at 4 °C. Then cells were washed and incubated for 1 h at room temperature with a goat anti-rabbit secondary antibody-conjugated Alexa 488 fluorophore (Invitrogen) diluted 1:400. DAPI was used to stain nuclei. Confocal images were acquired with a Zeiss LSM 510 META laser scanning confocal microscope (Carl Zeiss MicroImaging, Jena, Germany) by using a multilane argon laser (excitation 488 nm) or a 405 diode laser (excitation 405 nm) with a 63 × C-Apochromat, NA 1.2-W objective.

mRNA Isolation and q-PCR Analysis—Total mRNA isolation was carried out with the TRIzol reagent (Invitrogen) according to the manufacturer’s instructions. RNA concentration and purity were measured with a Nano-drop spectrophotometer (ND-1000, Thermo Scientific, Marietta, OH). The retrotranscription kit (Applied BioSystems, Carlsbad, CA) was employed to reverse transcribe 0.5 μg of total RNA to cDNA. qRT-PCR was performed with TaqMan probes (Applied BioSystems) for Cyp26a1 (Mm00514486_m1), Stra6 (Mm00486457_m1), Lrat (Mm00469972_m1), Bco1 (Mm0151350_m1), and Bco2 (Mm00460051_m1). The β-actin (Mm01205647_g1) probe set was used as an endogenous control. All real time experiments were performed with an ABI Step-One Plus qRT-PCR instrument (Applied BioSystems).

Immunoblotting—Tissue samples were homogenized in M-PER mammalian protein extraction reagent (Thermo Scientific) following the manufacturer’s instructions. For RBP and LRAT determinations in liver, 20 μg of total protein was used. To determine RBP serum levels, 2 μl of serum was diluted with 40 μl of PBS containing complete mini EDTA-free protease inhibitor (Roche Diagnostics), and 4 μl of this solution was used for immunoblot analysis. To quantify RBP levels in HepG2 cells, 10–20 μg of total protein cell lysate were subjected to immunoblot analyses. Protein samples were separated by SDS-PAGE and then electroblotted onto PVDF membranes (Bio-Rad). Membranes were blocked with fat-free milk powder (5% w/v) dissolved in Tris-buffered saline (15 mM NaCl and 10 mM Tris/HCl, pH 7.5) containing 0.01% Tween 100 (TBS-T), washed, and incubated overnight at 4 °C with the appropriate primary antibody. For RBP detection, a rabbit anti-human RBP serum (DakoCytomation) was used at a 1:1000 dilution. For LRAT detection, a noncommercial anti-LRAT monoclonal antibody was employed at a dilution of 1:2000 (33). β-Actin antisemir (Cell Signaling, Boston) at a dilution of 1:1000 served as a loading control. Secondary antibodies were either horse-radish peroxidase-conjugated anti-rabbit IgG (Promega, Madison, WI) or anti-mouse IgG (Promega) used at a dilution of 1:5000. Immunoblots were developed with the ECL system (GE Healthcare).

HPLC Analyses of Apocarotenoids and Carotenoids—Non-polar compounds were extracted from either 100 μl of serum or 30 mg of tissue under a dim red safety light as described previously (32). The extraction solution was composed of 200 μl of methanol, 400 μl of acetone, and 500 μl of hexane. The organic phase was removed, and the extraction was repeated with 500 μl of hexane. For saponification, selected tissues were incubated with 100 μl of 12% pyrogallol (Sigma) in ethanol, 200 μl of 30% KOH in water, and 1 ml of ethanol for 2 h at 37 °C. After saponification, 2 ml of ethanol and 2 ml of H2O were added, and samples were extracted twice with 3 ml of ether/hexane (2:1, stabilized with 1% ethanol). After centrifugation for 5 min at 800 × g, organic layers were collected, pooled, and dried in a SpeedVac (Eppendorf, Hamburg, Germany). The residue was dissolved in 200 μl of HPLC solvent (hexane/ethyl acetate 90:10, v/v). HPLC was performed with a normal phase Zobas Sil (5 μm, 4.6 × 150 mm) column (Agilent Technologies, Santa Clara, CA). For isocratic chromatographic separation of apocarotenoid-oximes, 20% ethyl acetate was mixed with 80% hexane. For isocratic chromatographic separation of apocarotenoids, 5% ethyl acetate was mixed with 95% hexane. For retinyl ester (RE) separation, a linear gradient of 1% ethyl acetate in hexane over 15 min followed by 10 min with 20% of ethyl acetate in hexane was used. In both cases, the flow rate was 1.4 ml/min. For molar quantification of retinoids, the HPLC was previously scaled with the pattern compounds ROL, RE (Sigma), BC (Calbiochem), and APO10ol.

Mass Spectroscopy Analyses—Polar and nonpolar apocarotenoids were separated and extracted from HepG2 and liver samples for mass spectrometry by previously described methodology (36). Briefly, either HepG2 cells grown on 6-well plates or 750 mg of liver was homogenized in 0.9% saline solution in a final volume of 0.75 ml. To separate nonpolar from polar apocarotenoids, the solution was alkalinized with 0.025 M KOH in 1 ml of ethanol. Then 1 ml of acetonitrile was added, followed by extraction of nonpolar apocarotenoids with 10 ml of hexane. After addition of 60 μl of 4 mM HCl, polar retinoids were extracted with 10 ml of hexane. Lipid extracts were dried down in a SpeedVac, and residues were dissolved in hexane, ethyl acetate, and acetic acid (79:9:20:0.1 v/v) for further analysis.

Mass Spectrometry—MS-based detection of apocarotenoids was achieved with a LXQ linear ion trap mass spectrometer (Thermo Scientific, Waltham, MA) equipped with an atmospheric pressure chemical ionization interface coupled to an Agilent 1100 HPLC series and diode array detector (Agilent Technologies). Separation of compounds was achieved on a normal phase Zobas Sil (5 μm, 4.6 × 150 mm) column (Agilent Technologies) by isocratic flow of 20% ethyl acetate in hexane, 0.0175% acetic acid at rate of 1.4 ml/min. The HPLC effluent was directed into the MS via an atmospheric pressure chemical ionization probe operated in the positive ionization mode. Parameters of both ionization and detection were tuned with synthetic β-apocarotenoid standards to ensure the highest possible sensitivity. The detection limit (defined as a minimum of three times ion intensity over the background signal) was determined by titrations with known amounts of standards. The matrix had no gross influence on sensitivity, which was estimated to be ~1 pmol for all of the tested apocarotenoids. RE, ROL, retinal, and RA were purchased from Sigma. β-Apo-10′-, β-apo-12′-, and β-apo-14′-apocarotenonic acid as well as β-apo-10′-carotenol were a gift from BASF (Ludwigshafen, Germany). APO10ol was prepared from β-apo-10′-carotenol.
by BH₄ reduction. Retinoids and apocarotenoids were detected by a selected reaction monitoring mode using the following transitions: ROL and RE, m/z 269.2 → 213.2; retinol-oximes, m/z 300.3 → 208.2; RA, m/z 301.4 → 205.2; APO10ol and APO10ol esters, m/z 361.5 → 237.3; β-apo-10'-carotenol-oximes, m/z 392.4 → 285.2; β-apo-10'-carotenolic acid, m/z 393.4 → 347.3; β-apo-12'-carotenolic acid, m/z 367.4 → 349.3.0; and β-apo-14'-carotenolic acid, m/z 327.3 → 215.0.

ROL and APO10ol Uptake Assays—NIH3T3 cells expressing both LRAT and STRA6 were cultured in 12-well culture plates at a density of 1 × 10⁶ cells per well as described previously (37). When cells reached ~85% confluence, they were washed with PBS. Then serum-free medium containing human RBP (hRBP) with bound 2 μM ROL or APO10ol was added. As controls, separate sets of plates were treated with 2 μM of either free ROL or APO10ol. Cells were incubated at 37 °C. After collection, cells were washed twice with PBS, harvested in 250 μl of PBS, sonicated, and stored at −20 °C. The cell homogenate (200 μl) was also extracted with 200 μl of methanol, 400 μl of acetone, and 600 μl of hexane, and the entire organic phase was collected and dried down in a SpeedVac. Protein levels in the cell homogenate were measured by Bradford’s method (Bio-Rad), and esters of ROL or APO10ol were analyzed by normal phase HPLC as described above. All experiments were performed at least in duplicate.

Expression, Purification, and Refolding of Human Serum RBP—RBP expression and purification from *Escherichia coli* was accomplished essentially as described previously (38). Briefly, hRBP cDNA was cloned into a pET3a expression vector and expressed in BL-21 DE3 cells according to a standard protocol. Bacterial cells were harvested and lysed by osmotic shock. Insoluble material was pelleted by centrifugation, washed, and solubilized in 7 M guanidine hydrochloride and 10 mM dithiothreitol (DTT). After overnight incubation, insoluble material was removed by ultracentrifugation, and the supernatant was used for the hRBP refolding procedure. hRBP was refolded by dropwise addition of solubilized material into a mixture containing 1 mM ROL (Sigma) or APO10ol. Refolded holo-hRBP was dialyzed against 10 mM Tris/HCl buffer, pH 8.0, and loaded onto a DE53 anion exchange chromatography column (Whatman). Holo-hRBP was eluted with linear gradient of NaCl (0–1 M) in 10 mM Tris/HCl buffer, pH 8.0. Collected fractions were examined by SDS-PAGE and UV-visible spectroscopy to determine the protein/apocarotenoid ratio. Fractions containing holo-hRBP were pooled together and concentrated in a Centricon centrifugal filter device (cutoff 10,000 Da) (Millipore, Billerica, MA).

**LRAT Assays**—Bovine RPE was used to measure LRAT enzymatic activity. Bovine RPE microsomes were obtained from fresh RPE isolated from bovine eye cups with 0.05 M MOPS, 0.25 M sucrose, 1 mM DTT, pH 7. The extract was homogenized in a glass homogenizer and then centrifuged (12,000 × g, 30 min, 4 °C). The resulting supernatant was collected for further ultracentrifugation at 4 °C for 60 min (100,000 × g). The RPE microsomal fraction was resuspended in 5 mM BisTris, 5 mM DTT, pH 7, and stored in small aliquots at −80 °C. To remove pre-existing RE and other retinoids that could interfere with the enzyme assay, 500-μl aliquots of the RPE microsomal fraction were irradiated in a quartz cuvette for 20 min on ice with a ChromatoUVE transilluminator (model TM-15 from UVP Inc.) at low intensity. Then 20 μl of RPE microsomal fractions (15 μg of protein) were added to a total volume of 200 μl of 10 mM Bistris propane buffer, 1 mM DTT, and 1% BSA, pH 7.5. The reaction was initiated by addition of variable amounts of ROL or APO10ol delivered in 1 μl of dimethylformamide. The reaction mixture was vigorously vortexed and incubated at 37 °C with 550 rpm rotation in a thermomixer (Eppendorf thermomixer compact; Eppendorf, Germany). After incubation, the reaction was stopped by addition of 0.3 ml of methanol and 0.3 ml of acetone followed by total retinoid extraction into 0.5 ml of hexane. The hexane extraction was repeated, and the combined organic phases were dried in a SpeedVac (Eppendorf). Dried residues were dissolved in solvent, and their β-apocarotenoid composition was analyzed by HPLC as described above.

**Enzymatic Activity of Human BCO1**—Twenty milliliters of baculovirus containing a human BCO1 plasmid (28) was added to 800 ml of *Spodoptera frugiperda* 9 cell culture in a 2-liter baffled flask. Cell pellets (20–30 g) were resuspended in 50 ml of sample buffer containing 20 mM Tricine, pH 7.5, 1 mM tris(2-carboxyethyl)phosphine hydrochloride (Hampton Research, Aliso Viejo, CA), and 1 tablet of Complete EDTA-free protease inhibitor mixture (Roche Diagnostics). Cells were lysed in a glass tissue grinder, and the lysate was centrifuged at 100,000 × g for 1 h at 4 °C (Beckman Coulter Optima™ L-90K Ultracentrifuge). The supernatant was collected and used for assays of enzymatic activity as described previously (39). Briefly, APO10ol, β-8'-apocarotenol, and β-12'-apocarotenolic acid were mixed with 30 μl of 12% (w/v) n-octyl β-D-thioglucopyranoside micelles (3%), dissolved in ethanol, and dried in a SpeedVac. Then the enzyme solution (100 μl) or elution buffer only was added to the residues and vortexed for 20 s. Enzymatic assays were incubated at 28 °C with 300 rpm rotation in a thermomixer (Eppendorf) for 20 min under a dim red safety light. Reactions were stopped by the addition of 100 μl of NH₃OH and 200 μl of methanol. Lipids were extracted and analyzed by HPLC as described above. For enzyme tests with β-8'-apocarotenol, lipids were extracted without the addition of NH₃OH and analyzed by HPLC.

**Enzymatic Activity of Murine BCO2**—For heterologous expression of murine BCO2, a plasmid containing the BCO2 full-length cDNA was transformed in the *E. coli* strain XL1-blue (Stratagene Inc., La Jolla, CA) as detailed previously (28). Enzymatic activity of crude extracts (100 μl of total protein) was determined in the presence of β-cryptoxanthin (CaroteNature, Lupsingen, Switzerland) in n-octyl β-D-thioglucopyranoside micelles (3%) at 28 °C. Reactions were stopped after 8 min by the addition of 300 μl of methanol, and lipophilic compounds were extracted and subjected to HPLC analysis as described above.

**Statistical Analyses**—Values are expressed as means ± S.E. Statistical significance of differences was derived from the two-tailed Student’s t test with the threshold of significance set at *p* < 0.05. *V*ₘₐₓ and *K*ₘ values for APO10ol kinetics were calculated with software Origin 9 (OriginLab Corp., Northampton, MA).
RESULTS

**BC Accumulation in Mice Depends on BCO1**—We subjected 6-week-old and sex-matched WT, Bco1−/−, Bco2−/−, and ko/ko mice to dietary intervention with BC. During the 10-week feeding period, we found no differences in food intake and/or weight gain between genotypes (Fig. 1, A–C). Then mice were sacrificed, and blood and tissues were collected. HPLC analyses revealed that Bco1−/− and ko/ko mice had accumulated BC in the serum, liver, and lungs, but no significant differences were found in their total BC levels in serum and tissues. In contrast, WT and Bco2−/− mice showed about 100-fold lower levels of BC in tissues and serum when compared with Bco1−/− and ko/ko mice (Fig. 1, D–F). Again, no significant difference was observed in total BC levels between WT and Bco2−/− mice. Hence, we conclude that BCO2 did not affect bulk BC metabolism as indicated by comparable accumulation of BC in Bco1−/− and ko/ko mice.

**BCO2 Catalyzes β-Apocarotenoid Production in Vivo**—We next analyzed whether apocarotenoids other than retinoids were synthesized from BC by different mouse genotypes. After total hepatic lipid extraction and separation by HPLC, APO10ol was detected in Bco1−/− mice (Fig. 2A). APO10ol was identified by its retention time and mass as compared with an authentic standard (Fig. 2, B–D). The primary cleavage product β-apo-10′-carotenal was not detected by our LC-MS system (Fig. 2, E–G). Saponification of liver samples from Bco1−/− mice resulted in a 3-fold higher amount of APO10ol relative to nonsaponified liver samples, indicating that APO10ol exists largely in its esterified form (Fig. 2A). Also small amounts of APO10ol became detectable in saponified liver samples of WT mice. In contrast, APO10ol was absent in Bco2−/− and ko/ko mice, indicating that production of this compound depends on a functional Bco2 allele (Fig. 2A). Next, we investigated whether β-apo-carotenoid acids are formed from dietary BC in a BCO2-dependent manner. In Bco1−/− and ko/ko mice, apocarotenoids such as β-apo-12′- and β-apo-14′-carotenoid acid were not detected either in selected reaction monitoring or single ion monitoring modes (data not shown). However, trace amounts of β-10-apo-carotenoid acid became detectable in Bco1−/− mice (data not shown).

**LRAT Can Esterify APO10ol**—Our analyses revealed that APO10ol is the main eccentric cleavage product of BC in mice. In the liver of Bco1−/− and WT mice, this compound mainly existed in esterified form. The major hepatic enzyme for ROL esterification is LRAT (33). Because of the similar chemical structures of APO10ol and ROL, we tested whether APO10ol is a substrate for this enzyme by using bovine RPE microsomes as a source of LRAT. Upon incubation with APO10ol, we observed a time-dependent decrease of the substrate that corresponded to the formation of APO10ol esters (Fig. 3, A and B). To demonstrate that APO10ol esterification was indeed LRAT-dependent, we then performed assays in the presence of tosylsine chloromethyl ketone (TLCK). TLCK is serine protease inhibitor that also inhibits enzymes such as LRAT with a thiol group in their active sites (40–42). After preincubation of microsomes with TLCK, esterification of APO10ol was greatly decreased (Fig. 3C). Next, we determined $K_m$ and $V_{max}$ values for APO10ol and ROL, respectively. LRAT displayed a 3-fold higher $K_m$ value for APO10ol (3.85 μM) as compared with ROL (1.29 μM). But strikingly, the $V_{max}$ value was about 20-fold
lower for APO10ol (0.41 pmol/s) relative to ROL (9.3 pmol/s) (Fig. 3, D and E). To determine whether APO10ol can be esterified by LRAT in vivo, we tested this hypothesis in Lrat knockout mice. Lrat−/− and WT mice were gavaged with apo-10′-β-apocarotenol (20 mg/kg). After 3 h, they were sacrificed, and their livers were subjected to HPLC analysis. Livers of Lrat−/− mice showed significantly lower amounts of APO10ol esters as compared with WT controls. But small amounts of APO10ol esters were still formed in Lrat−/− mice, indicating that additional enzyme systems can esterify this compound (Fig. 3F).

APO10ol Can Trigger RBP Release—We showed that APO10ol, like ROL, existed in esterified form in mouse liver. Therefore, we next wondered whether APO10ol also can be released from hepatocytes like vitamin A. To experimentally

---

**FIGURE 2. APO10ol is the major long-chain β-apocarotenoid in mouse liver.** Six-week-old wild type (wt), Bco2−/−, Bco1−/−, and ko/ko mice were provided a controlled diet for 10 weeks with BC (50 mg/kg) as the sole source for β-apocarotenoid production. A, hepatic levels of free APO10ol (white bars) and total APO10ol (free + esterified) (dark gray bars). nd, not detectable. Values indicate means ± S.E. for at least five female animals per tissue and genotype. Statistical significance was tested by the two-tailed Student’s t test with results compared with the WT group. Threshold of significance was set at p < 0.05. B and C, MS/MS diffraction patterns of APO10ol (B) and β-apo-10′-carotenal (APO10al) (C). D and E, presence of APO10ol (D) and APO10al (E) extracted from livers of Bco1−/− (red trace) and ko/ko (blue trace) mice is indicated by an asterisk and identified by retention times and selected reaction monitoring (SRM) modes as compared with authentic standards (upper panel).
address this question, we employed human hepatoma HepG2 cells used as a reliable model to study RBP-mediated vitamin A release (43) and cultured these cells in serum-free medium supplemented with and without vitamin A. Cells then were treated with either ROL or APO10ol. By immunocytochemistry and confocal imaging we visualized cellular RBP levels. As expected, cells cultured in vitamin A-free medium accumulated the vitamin A carrier RBP (Fig. 4, A and B). ROL treatment of cells decreased intracellular RBP levels, indicative of its release (Fig. 4, A and B). Similarly, APO10ol treatment also decreased cellular RBP levels although this decrease was less pronounced.

We next subjected 12-week-old Bco1/H11002/H11002 mice to dietary vitamin A restriction for 5 weeks. We then gavaged these mice with a pharmacological dose (30 mg/kg) of either ROL or APO10ol. Mice were sacrificed 4 h later, and RBP protein levels were measured in their livers. Siblings maintained on a vitamin A-sufficient diet were used as controls. In vitamin A-deprived mice treated with vehicle only, RBP levels were significantly higher than in vitamin A-sufficient mice. Moreover, ROL treatment of vitamin A-deprived Bco1/H11002/H11002 mice decreased hepatic RBP levels (Fig. 4, C). Gavage with APO10ol also resulted in a reduction of RBP levels, indicating that APO10ol can trigger the release of RBP from the liver of these animals (Fig. 4, C).

**Cellular Uptake of APO10ol**—We next tested whether APO10ol bound to RBP can be taken up by target cells. Accord-
ingly, recombinant human RBP was produced in E. coli and refolded in the presence of ROL or APO10ol. Refolded RBP and free apocarotenoid fractions were purified by size exclusion chromatography to ensure the absence of free ROL or APO10ol in both protein preparations (Fig. 4, \( D \) and \( E \)) (38). We used NIH3T3 cells expressing STRA6 and LRAT previously established to measure ROL and APO10ol uptake (37, 38). These cells were incubated with \( 2\mu M \) free and RBP-bound APO10ol as well as with \( 2\mu M \) free and RBP-bound ROL. Uptake efficiency was measured as the amount of esterified APO10ol at different time points (Fig. 4\( F \)). Both RBP-bound ROL and APO10ol were taken up more rapidly than their unbound forms. However, RBP-bound ROL was taken up far more rapidly than its APO10ol counterpart. This difference might be explained in part by the enzymatic activity of LRAT that can catalyze ROL esterification more rapidly than APO10ol esterification (Fig. 3). Additionally, a lower affinity of RBP-bound APO10ol to STRA6 may slow down its transfer to LRAT and its accumulation as ester.

**BCO1, but Not BCO2, Is Critical for Vitamin A Homeostasis**—We showed that APO10ol is BCO2-dependently produced from BC and can be esterified and transported by similar mechanisms as ROL. Thus, to learn whether dietary APO10ol affects vitamin A homeostasis, we determined retinoid levels in the liver and lungs of different mouse mutants supplemented with BC and compared them with those of WT mice. Additionally, we measured dietary BC effects on the expression levels of key proteins for vitamin A homeostasis such as RBP, STRA6, LRAT, and CYP26a1. CYP26a1 is an RA-catabolizing enzyme widely used as a surrogate marker for RA status in tissues (44). Bco2\(^{-/-}\) mice displayed increased hepatic RE stores as compared with the other genotypes (Fig. 5\( A \)). LRAT and CYP26a1 expression also were significantly elevated as compared with WT animals (Fig. 5, \( B \), \( C \), and \( E \)). Furthermore, hepatic free ROL levels also were increased (Fig. 5\( D \)). In lungs of Bco2\(^{-/-}\) mice, no differences in RE levels were detectable when compared with WT mice (Fig. 6\( A \)). Additionally, LRAT mRNA expression levels and free ROL levels were not significantly altered as compared with WT mice (Fig. 6, \( B \) and \( D \)). Only a slight but significant increase of Stra6 mRNA level was observed (Fig. 6\( C \)). Thus, genetic disruption of BCO2 increased vitamin A status of the liver but not the lungs on the BC diet.

**FIGURE 4. APO10ol transport and uptake.** HepG2 cells were maintained at 37 °C in the presence (VAS) or absence (VAD) of vitamin A in the cell culture medium for 48 h. Cells then were incubated in 4 \( \mu M \) ROL or APO10ol with ethanol used as a vehicle control for 2 h. A, immunostaining for RBP (green). Nuclei were stained with DAPI (blue). Scale bar, 20 \( \mu m \). B, immunoblot analysis for RBP in protein extracts of HepG2 cells, using \( \beta\)-actin as the loading control. Ten \( \mu g \) of total protein were loaded per lane with \( \beta\)-actin used as a loading control. At least three mice were used per experimental group. D and \( E \), spectral characteristics of purified ROL-hRBP \( (D) \) and APO10ol-hRBP \( (E) \). F, NIH3T3 cells expressing human LRAT and STRA6 were incubated in either 2 \( \mu M \) of ROL-hRBP (open circles), APO10ol-hRBP (open triangles), free ROL (filled circles), or free APO10ol (filled triangles). Uptake efficiency was measured as RE and APO10ol ester formation per \( \mu g \) of protein at different time points. Values shown for cell culture experiments indicate means \pm S.E. from three independent experiments. Statistical significance was tested by the two-tailed Student's \( t \) test with results compared with free ROL (*) and APO10ol (*), respectively. Threshold of significance was set at \( p < 0.05 \).
In Bco1<sup>−/−</sup> mice, no significant differences were observed for hepatic RE stores as compared with WT mice (Fig. 5A). Similarly, LRAT expression was not altered in the liver of this mutant (Fig. 5, B and E). However, the expression level of Cyp26a1 was significantly decreased in Bco1<sup>−/−</sup> as compared with WT animals (Fig. 5C). Additionally, these mice showed increased hepatic RBP levels (Fig. 5E) and a substantial reduction of free hepatic ROL levels (Fig. 5D). Serum vitamin A and RBP levels were comparable between different genotypes (Fig. 5F). In the lungs of Bco1<sup>−/−</sup> mice, RE stores were significantly reduced as compared with WT mice (Fig. 6A). The decreased vitamin A status of this organ also was reinforced by the reduced Lrat and Stra6 mRNA levels (Fig. 6, B and C) and reduced free ROL levels in this organ (Fig. 6D). Thus, these analyses indicated that genetic disruption of BCO1 impaired vitamin A homeostasis, whereas genetic disruption of BCO2 resulted in increased hepatic but not pulmonary vitamin A status when BC was provided as the sole dietary source of this vitamin. To determine whether BCO2 impacted vitamin A status in BCO1 deficiency, we analyzed ko/ko mice. In livers of these mice, Cyp26a1 mRNA expression was reduced to a similar extent as in Bco1<sup>−/−</sup> mice (Fig. 5C). Additionally, hepatic free ROL and RBP protein levels were also comparable between the double and single BCO1 mutant (Fig. 5, D and E). Finally, pulmonary RE and free ROL levels (Fig. 6, A and D) as well as Lrat and Stra6 mRNA expression levels (Fig. 6, B and C) were all reduced to a similar extent as in Bco1<sup>−/−</sup> mice. Thus, we excluded that BCO2 alone can improve vitamin A status on BC diet.

**Increased Hepatic RA Status Is Associated with BCO1 Expression**—Our analyses revealed that hepatic RE stores were significantly elevated in Bco2<sup>−/−</sup> as compared with WT mice. Similarly, Lrat expression was not changed (Fig. 5B and E). However, the expression level of Cyp26a1 was significantly increased in Bco2<sup>−/−</sup> as compared with WT animals (Fig. 5C). qRT-PCR analysis of hepatic Cyp26a1 mRNA levels (Fig. 5C). Additionally, hepatic free ROL and RBP protein levels were also comparable between the double and single BCO1 mutant (Fig. 5, D and E). Finally, pulmonary RE and free ROL levels (Fig. 6, A and D) as well as Lrat and Stra6 mRNA expression levels (Fig. 6, B and C) were all reduced to a similar extent as in Bco1<sup>−/−</sup> mice. Thus, we excluded that BCO2 alone can improve vitamin A status on BC diet.
demonstrate more directly that the altered hepatic vitamin A status of $Bco2^{-/-}$ mice is causatively related to $Bco1$, we raised mice in a similar fashion as described above but on diet containing preformed vitamin A (4000 IU per kg) instead of BC. No differences should become apparent between $Bco1^{-/-}$ and WT mice if BC conversion caused this phenotype. In fact, our HPLC analysis revealed comparable levels of RE in the livers of $Bco2^{-/-}$ and WT mice (Fig. 7B). Additionally, expression of $Bco1$, $Cyp26a1$, and $Lrat$ was comparable between $Bco2^{-/-}$ and WT mice under this dietary regimen (Fig. 7C).

APO10ol Can Be Metabolized to Vitamin A by $Bco1$—$Bco1^{-/-}$ mice displayed higher levels of APO10ol than WT mice. Thus, we tested whether this difference might be related to further metabolic conversion of APO10ol by assaying the effect of purified recombinant human $Bco1$ on APO10ol. This enzymatic analysis revealed that $Bco1$ converted APO10ol to retinaldehyde (Fig. 8A). We also performed $Bco1$ enzymatic assays with other $apo$-apocarotenoids. These analyses revealed that $apo-8$- and $apo-12$-apocarotenoic acids were converted to retinal (Fig. 8, B and C), indicating that $Bco1$ can catalyze the conversion of $apo$-apocarotenoids with different chain lengths and oxidation states. We next determined the specific activity of the protein crude extract for different apocarotenoid substrates and compared it with the established substrate BC. This analysis revealed that apocarotenoids were converted to retinal with comparable efficiency as BC (Fig. 8). The specific activity was determined to be 183.5, 79.7, 27.6, and 80.4 pmol/min/mg, respectively, for APO10ol, $apo-8$- and $apo-12$-apocarotenoic acids, and BC. To confirm this finding in a cellular environment, we added APO10ol to a culture medium of human hepatoma HepG2 cells. After incubation, cells were harvested, and lipophilic compounds were extracted and analyzed, which revealed that various retinoids were produced from APO10ol, including ROL, RE, and retinaldehyde (Fig. 9A). We also tested by LC-MS analyses whether $apo-12$- and $apo-14$-apocarotenoic acids were synthesized from
APO10ol (Fig. 9B). Except for trace amounts of β-apo-12′-apocarotenoid acid, none of these compounds were detectable indicating that conversion to retinoids is the main metabolic fate of APO10ol in these cells.

Eccentric Cleavage by BCO2 Converts β-Cryptoxanthin to Vitamin A—We showed that APO10ol produced by BCO2 can be further converted to vitamin A via a second cleavage reaction carried out by BCO1. This stepwise cleavage pathway plays a minor role in BC metabolism. However, it could prove critical for the metabolism of asymmetric carotenoids such as β-cryptoxanthin. This carotenoid possesses a non-substituted β-ionone, necessary for vitamin A formation, and a 3-OH-β-ionone ring. Stepwise cleavage of β-cryptoxanthin by BCO2 and BCO1 would first remove the 3-OH-ring site and yield APO10ol. The latter compound would then be converted to vitamin A by BCO1. Thus, we incubated recombinant murine BCO2 with β-cryptoxanthin and extracted lipophilic compounds for HPLC analysis. In fact, β-apo-10′-carotenal was the main cleavage product of this carotenoid, and only small amounts of 3-OH-β-apocarotenal were formed (Fig. 10A). To analyze the metabolic fate of β-cryptoxanthin in an animal model, we injected 12-week-old female WT, Bco2−/−, and Bco1−/− mice (n = 3 each) with pharmacological doses of β-cryptoxanthin. After three consecutive daily injections, we analyzed β-cryptoxanthin levels in the livers of these mice. Bco2−/− mice showed a significant hepatic accumulation of β-cryptoxanthin as compared with Bco1−/− and WT mice (Fig. 10B). This accumulation suggested that BCO2 is critically required for the metabolism of β-cryptoxanthin in mouse liver.

DISCUSSION

There has been a long lasting controversy about symmetric versus eccentric cleavage of BC in mammalian vitamin A biol-
FIGURE 9. APO10ol is metabolized into retinoids by HepG2 cells. HepG2 cells were incubated with APO10ol (2 μM) (black line) or vehicle (ethanol) (gray line) for 12 h at 37 °C. Then cells were harvested, and lipophilic compounds were extracted and subjected to LC-MS analysis. A, analysis of nonpolar apocarotenoids. The presence of APO10ol, β-apo-10'-carotene-oximes (APO10alox), ROL, and all-trans-retinal-oximes (RALox) extracted from HepG2 cells treated with APO10ol are indicated by asterisks and were identified by retention times and selected reaction monitoring modes as compared with authentic standards (upper panel). Esters of APO10ol and ROL are evident in examined samples by the presence of distinct peaks at 2.2 min of elution. B, analysis of acidic apocarotenoids. The presence of RA, β-apo-12'-caroteneic acid (APO12 acid), APO14 acid, β-apo-14'-caroteneic acid; APO10 acid, β-apo-10'-carotenoic acid extracted from HepG2 cells treated with APO10ol or vehicle are indicated by asterisks and were identified by retention times and selected reaction monitoring modes as compared with authentic standards (upper panel).
Carotenoid Oxygenases and Provitamin A Metabolism

By a comprehensive study in single and double mutants for Bco1 and Bco2, we documented here the contributions of these enzymes to BC metabolism. This analysis revealed that BCO1 is the major BC-metabolizing enzyme in mice. Furthermore, it showed that BCO2 does not influence vitamin A-dependent processes. These conclusions are based on the following observations. Bco1<sup>−/−</sup> and ko/ko mice accumulated large amounts of hepatic BC, although no such accumulation took place in Bco2<sup>−/−</sup> and WT mice. Because no differences were observable between Bco1<sup>−/−</sup> and ko/ko animals, we conclude that BCO2 alone does not significantly contribute to BC homeostasis in mice. Nevertheless, small amounts of APO10ol were produced in Bco1<sup>−/−</sup> mice. Upon saponification, hepatic APO10ol content increased in Bco1<sup>−/−</sup> mice, and trace amounts of APO10ol also became detectable in WT mice. The lack of APO10ol in Bco2<sup>−/−</sup> and ko/ko mice clearly demonstrated that this production was BCO2-dependent. We did not detect other apocarotenoids in the liver of these mice, suggesting that BCO2 displays high regional selectivity for the C9, C10 double bond of BC. Although minor amounts of APO10ol were produced, our analysis failed to provide evidence that this production impacted vitamin A homeostasis and retinoid signaling. Bco1<sup>−/−</sup> and ko/ko mice showed similar reductions of retinoids and expression levels of key components of vitamin A metabolism, although the single knock-out mice expressed BCO2 and produced APO10ol. This finding also agrees with the results of previous studies in Bco1<sup>−/−</sup> mice. BC supplementation did not influence global transcriptional activity of white adipose tissue in Bco1<sup>−/−</sup> mice as compared with siblings maintained on the same diet without BC (35). When compared with nonsupplemented Bco1<sup>−/−</sup> mouse siblings, BC supplementation also had no gross effect on global gene expression profiles in other tissues, including the liver and the lungs (50). Together, these findings make it improbable that BCO2 is a component of a pathway for the production of β-apocarotenoid signaling molecules that agonize and/or antagonize the activity of nuclear receptors such as RARs.

Several studies showed that β-apocarotenoids can interact with proteins that bind vitamin A derivatives, including nuclear receptors (24). This finding is not surprising when it is considered that these compounds are chemically and structurally related to vitamin A. In fact, we also observed that APO10ol can interact with molecular components of vitamin A metabolism. APO10ol was esterified by LRAT, bound by the vitamin A carrier RBP, and taken up by target cells from its RBP-bound form by STRA6. Therefore, high concentrations of APO10ol presumably should compete with vitamin A for the same metabolic pathways. However, our study indicated instead that such ambiguity and competition between BC metabolites is limited in mammals. One possible explanation involves the substrate specificity of BCO2. In Bco1<sup>−/−</sup> mice, BC existed in 1000-fold excess over APO10ol, indicating that BC is a poor substrate for BCO2 in vivo. This conclusion is further corroborated by enzymatic assays with the asymmetric carotenoid, β-cryptoxanthin, that contains a 3-OH and a nonhydroxylated β-ionone ring site. Our results showed that BCO2 preferentially removed the 3-OH-β-ionone ring site from this carotenoid. This preference for hydroxylated ring sites of carotenoids also has been reported for BCO2 from ferrets (30). Moreover, structural analyses and enzymatic properties of carotenoid oxygenases indicate that these enzymes specifically interact with one ionone ring side of their carotenoid substrates (51–53). This specificity of BCO2 for carotenoids with 3-OH-β-ionone ring sites is also indicated by β-cryptoxanthin accumulation in BCO2-deficient mice as we found here and by accumulation of zeaxanthin and lutein as we demonstrated previously in this mouse mutant (29). The preference of BCO2 for 3-hydroxyionone rings also

---

**FIGURE 10.** BCO2 converts β-cryptoxanthin to β-apo-10'-apocarotenal. A, protein extract containing recombinant murine BCO2 was incubated with increasing concentrations of β-cryptoxanthin. Lipophilic compounds were extracted and subjected to HPLC analysis. Amounts of products (β-apo-10'-carotenal (filled diamonds) and 3-OH-β-apo-10'-carotenal (open squares)) are plotted against the substrate concentration. B, β-cryptoxanthin levels in the liver of 12-week-old female WT, Bco2<sup>−/−</sup>, and Bco1<sup>−/−</sup> mice. Values indicate means ± S.E. from three animals per genotype. Statistical significance compared with the WT group was tested by the two-tailed Student’s t test with p < 0.05 considered significant.
explains why no hydroxylated apocarotenoids accumulate in mice. These compounds are further converted by BCO2 (29, 30). In contrast, the accumulation of nonhydroxylated apocarotenoids such as APO10ol and probably also other long chain /H9252-8/H11032- and /H9252-12/H11032-apocarotenoids in mice is prevented by another mechanism. As we demonstrated here, not only APO10ol but also /H9252-8/H11032- and /H9252-12/H11032-apocarotenoids were readily converted by recombinant human BCO1 into retinoids. This conversion was also the major metabolic fate of APO10ol in human HepG2 cells. Such BCO1 action prevents the accumulation of long chain /H9252-apocarotenoids in blood and tissues that may interact with components of vitamin A metabolic pathways. Simultaneously, this BCO1 activity ensures that even long chain /H9252-apocarotenoids can be utilized for vitamin A production, thereby providing an explanation for the observed vitamin A activity of these compounds (24). Physiologically, this pathway may play a role in the in vivo situation and whether it involves subsequent action of both BCOs. The compartmentalization of carotenoid metabolism as well as the shuttling of carotenoids and their apocarotenoid derivatives between different cellular compartments are the focus of future investigations of our laboratories.

Acknowledgments—We thank Dr. Hansgeorg Ernst (BASF, Germany) for the gift of apocarotenoid standards and Dr. Leslie Webster for critical comments on the manuscript. We thank Darwin Babino for BCO1 enzyme production. We also thank Maryanne Pendergast and the Neurosciences Imaging Center for assistance with confocal microscopy.

REFERENCES

1. Hall, J. A., Grainger, J. R., Spencer, S. P., and Belkaid, Y. (2011) The role of retinoic acid in tolerance and immunity. *Immunity* 35, 13–22
2. von Lintig, J., Kiser, P. D., Golczak, M., and Palczewski, K. (2010) The biochemical and structural basis for trans-cis isomerization of retinoids in the chemistry of vision. *Trends Biochem. Sci.* 35, 400–410
3. Rhinn, M., and Dollé, P. (2012) Retinoic acid signalling during development. *Development* 139, 843–858
4. Noy, N. (2010) Between death and survival: retinoic acid in regulation of apoptosis. *Annu. Rev. Nutr.* 30, 201–217
Carotenoid Oxygenases and Provitamin A Metabolism

5. Chambon, P. (1996) A decade of molecular biology of retinoic acid receptors. *FASEB J.* 10, 940–954

6. Palczewski, K. (2006) G protein-coupled receptor rhodopsin. *Annu. Rev. Biochem.* 75, 743–767

7. Sommer, A., and Vyas, K. S. (2012) A global clinical view on vitamin A and carotenoids. *Am. J. Clin. Nutr.* 96, 12045–1206S

8. Grune, T., Lietz, G., Palou, A., Ross, A. C., Stahl, W., Tang, G., Thurnham, D., Yin, S. A., and Biesalski, H. K. (2010) β-Carotene is an important vitamin A source for humans. *J. Nutr.* 140, 2268S–2285S

9. von Lintig, J. (2010) Colors with functions: elucidating the biochemical and molecular basis of carotenoid metabolism. *Annu. Rev. Nutr.* 30, 35–56

10. Olson, J. A., and Hayashi, O. (1965) The enzymatic cleavage of β-carotene into vitamin A by soluble enzymes of rat liver and intestine. *Proc. Natl. Acad. Sci. U.S.A.* 54, 1364–1370

11. Goodman, D. S., and Huang, H. S. (1965) Biosynthesis of vitamin A with rat intestinal enzymes. *Science* 149, 879–880

12. Lietz, G., Lange, J., and Rimbach, G. (2010) Molecular and dietary regulation of β,β-carotene 15,15′-monooxygenase 1 (BCMO1). *Arch. Biochem. Biophys.* 502, 8–16

13. Quadro, L., Blaner, W. S., Salchow, D. J., Vogel, S., Piantedosi, R., Gouras, K. M., Kopec, R. E., Schwartz, S. J., Curley, R. W., Jr., and Harrison, E. H. (2011) A mitochondrial enzyme degrades carotenoids and protects against oxidative stress. *FASEB J.* 25, 948–959

14. Wyss, A., Wirtz, G., Woggon, W., Brugger, R., Wyss, M., Friedlein, A., Bachmann, H., and Hunziker, W. (2000) Cloning and expression of β,β-carotene 15,15′-dioxygenase. *Biochem. Biophys. Res. Commun.* 271, 334–336

15. Redmond, T. M., Gentleman, S., Duncan, T., Yu, S., Wiggert, B., Gantt, E., and Cunningham, F. X., Jr. (2001) Identification, expression, and substrate specificity of a mammalian β-carotene 15,15′-dioxygenase. *J. Biol. Chem.* 276, 6560–6565

16. Paik, J., Durning, A., Harrison, E. H., Mendelsohn, C. L., Lai, K., and Blaner, W. S. (2001) Expression and characterization of a murine enzyme able to cleave β-carotene. The formation of retinoids. *J. Biol. Chem.* 276, 32160–32168

17. Lindqvist, A., and Andersson, S. (2002) Biochemical properties of purified recombinant human β-carotene 15,15′-monooxygenase. *J. Biol. Chem.* 277, 23942–23948

18. O’Byrne, S. M., Wonguirirjoj, N., Libien, J., Vogel, S., Goldberg, I. J., Baehr, W., Palczewski, K., and Blaner, W. S. (2005) Retinoid absorption and storage is impaired in mice lacking lecithin:retinol acyltransferase (LRAT). *J. Biol. Chem.* 280, 35647–35657

19. D’Ambrosio, D. N., Clugston, R. D., and Blaner, W. S. (2011) Vitamin A metabolism: an update. *Nutrients.* 3, 63–103

20. Quadro, L., Blaner, W. S., Salchow, D. J., Vogel, S., Piantedosi, R., Gouars, P., Freeman, S., Cosma, M. P., Colantuoni, V., and Gottesman, M. E. (1999) Impaired retinoid repletion and availability in mice lacking retinol-binding protein. *EMBO J.* 18, 4633–4644

21. Kawaguchi, R., Yu, J., Honda, J., Hu, J., Whitegege, J., Ping, P., Wiita, P., Bok, D., and Sun, H. (2007) A membrane receptor for retinol-binding protein mediates cellular uptake of vitamin A. *Science* 315, 820–825

22. Wang, X. D., Tang, G. W., Fox, J. G., Krisny, N. I., and Russell, R. M. (1991) Enzymatic conversion of β-carotene into β-apo-carotenals and retinoids by human, monkey, ferret, and rat tissues. *Arch. Biochem. Biophys.* 285, 8–16

23. Tang, G. W., Wang, X. D., Russell, R. M., and Krisny, N. I. (1991) Characterization of β-apo-13-carotenone and β-apo-14′-carotenal as enzymatic products of the excenctive cleavage of β-carotene. *Biochemistry* 30, 9829–9834

24. Eroglu, A., and Harrison, E. H. (2013) Current understanding of provitamin A β-carotene function as antagonists of retinoic acid receptors. *J. Biol. Chem.* 288, 15886–15895

25. Eroglu, A., Hruszkewycz, D. P., Curley, R. W., Jr., and Harrison, E. H. (2010) The excenctive cleavage product of β-carotene, β-apo-13-carotenone, functions as an antagonist of RXRα. *Arch. Biochem. Biophys.* 504, 11–16

26. Eroglu, A., Hruszkewycz, D. P., dela Sena, C., Narayanasamy, S., Riedl, K. M., Kopec, R. E., Schwartz, S. J., Curley, R. W., Jr., and Harrison, E. H. (2012) Naturally occurring excenctive cleavage products of provitamin A β-carotene and apocarotenals. *J. Biol. Chem.* 287, 23790–23807

27. Amengual, J., Lobo, G. P., Golczak, M., Li, H. N., Klimova, T., Hoppel, C. L., Wyss, A., Palczewski, K., and von Lintig, J. (2011) A mitochondrial enzyme degrades carotenoids and protects against oxidative stress. *FASEB J.* 25, 948–959

28. Amengual, J., Lobo, G. P., Golczak, M., Li, H. N., Klimova, T., Hoppel, C. L., Wyss, A., Palczewski, K., and von Lintig, J. (2011) A mitochondrial enzyme degrades carotenoids and protects against oxidative stress. *FASEB J.* 25, 948–959
44. Liu, L., Tang, X. H., and Gudas, L. J. (2008) Homeostasis of retinol in lecithin:retinol acyltransferase gene knockout mice fed a high retinol diet. Biochem. Pharmacol. 75, 2316–2324
45. Wolf, G. (1995) The enzymatic cleavage of β-carotene: still controversial. Nutr. Rev. 53, 134–137
46. Sharma, R. V., Mathur, S. N., Dmitrovskii, A. A., Das, R. C., and Ganguly, J. (1976) Studies on the metabolism of β-carotene and apo-β-carotenoids in rats and chickens. Biochim. Biophys. Acta 486, 183–194
47. Sharma, R. V., Mathur, S. N., and Ganguly, J. (1976) Studies on the relative biopotencies and intestinal absorption of different apo-β-carotenoids in rats and chickens. Biochem. J. 158, 377–383
48. Fleshman, M. K., Lester, G. E., Riedl, K. M., Kopec, R. E., Narayanasamy, S., Curley, R. W., Jr., Schwartz, S. J., and Harrison, E. H. (2011) Carotene and novel apocarotenoid concentrations in orange-fleshed Cucumis melo melons: determinations of β-carotene bioaccessibility and bioavailability. J. Agric. Food Chem. 59, 4448–4454
49. Handelman, G. J., van Kuijk, F. J., Chatterjee, A., and Krinsky, N. I. (1991) Characterization of products formed during the autoxidation of β-carotene. Free Radic. Biol. Med. 10, 427–437
50. van Helden, Y. G., Heil, S. G., van Schooten, F. J., Kramer, E., Hessel, S., Amengual, J., Ribot, J., Teerds, K., Wyss, A., Lietz, G., Bonet, M. L., von Lintig, J., Godschalk, R. W., and Keijer, J. (2010) Knockout of the Bemo1 gene results in an inflammatory response in female lung, which is suppressed by dietary β-carotene. Cell. Mol. Life Sci. 67, 2039–2056
51. Messing, S. A., Gabelli, S. B., Echeverria, I., Vogel, J. T., Guan, J. C., Tan, B. C., Klee, H. J., McCarty, D. R., and Amzel, L. M. (2010) Structural insights into maize viviparous14, a key enzyme in the biosynthesis of the phytohormone abscisic acid. Plant Cell 22, 2970–2980
52. Schwartz, S. H., Tan, B. C., Gage, D. A., Zeevaart, J. A., and McCarty, D. R. (1997) Specific oxidative cleavage of carotenoids by VP14 of maize. Science 276, 1872–1874
53. Oberhauser, V., Voolstra, O., Bangert, A., von Lintig, J., and Vogt, K. (2008) NinaB combines carotenoid oxygenase and retinoid isomerase activity in a single polypeptide. Proc. Natl. Acad. Sci. U.S.A. 105, 19000–19005
54. Lobo, G. P., Isken, A., Hoff, S., Babino, D., and von Lintig, J. (2012) BCDO2 acts as a carotenoid scavenger and gatekeeper for the mitochondrial apoptotic pathway. Development 139, 2966–2977