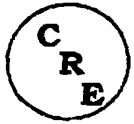


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TO: Mr. Richard Zazenski
FAX #: 303-799-8926
FROM: Jim Tozzi
DATE: November 27, 2000

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WORK ITEMS FOR PROPOSED RoC LISTING OF TALC

CRE

1. Prepare CRE white paper and Tozzi transmittal letter to RoC Subcomm.
 - draft to L. 11/27; final to RoC Subcomm. 11/29 (for receipt by 12/1)
2. Prepare detailed CRE comments to RoC Subcomm.
 - draft to L. 11/27; final to RoC Subcomm. 11/29 (for receipt by 12/1)
4. Prepare oral comments version (and written copy of oral comments) of CRE detailed comments for RoC Subcomm. meeting 12/13-15; attend talc portion of meeting and present CRE comments.
5. -- talc review by Subcomm. will probably take place 12/14; written version of oral comments are distributed at time of presentation
6. Consult and coordinate with L. on CRE and L. presentations (to make sure Luzenac raises all the issues necessary to make an adequate legal record) and network to explore obtaining favorable presentations at RoC meeting from FDA representative (on findings from 1994 workshop and 1995 publication) and ACGIH (American Council of Government and Industrial Hygienists).
7. -- both CRE and L. written comments are due to RoC Subcomm. 12/1; as noted above, written version of oral comments can be distributed at time of oral presentation
8. Prepare and submit CRE Freedom of Information Act request concerning nomination of talc and deliberations of RG1 and RG2 .
9. Obtain intelligence on what's happening in the RoC review committees, particularly when the issue gets to the NTP Executive Committee.
10. Monitor GSA FACA website following RoC Subcomm. meeting if they do not vote favorably to see whether there is still a FACA legal issue we want to pursue.
11. Review RoC Subcommittee outcome and prepare CRE comments for NTP Executive Committee.
12. Review potential for and advisability of litigation following RoC Subcomm. review . (Might be based, e.g., on failure of RoC Subcomm. to comply with FACA, failure to address significant issues raised in comments, and failure of NTP to follow its own

criteria for the “reasonably anticipated” category.)

13. Research possibility of having issue taken up by the Institute for Environmental Conflict Resolution.

Luzenac America

1. Prepare Luzenac comments for RoC Subcomm.
2. Consult with CTFA on obtaining FDA comments for RoC Subcomm. mtg.
3. Contact American Council of Government and Industrial Hygienists about sending someone to RoC Subcomm. meeting to make presentation on ACGIH finding of “not classifiable”.
4. Obtain presentations by other allies to RoC Subcomm.