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MEMORANDUM FOR:

SUBJECT: Summary of Project OFTEN Clinical Tests at Edgewood

1. Funds in the amount of $37,000 were transferred to Edgewood Arsenal on 17 February 1971 for the purpose of determining the clinical effects of EA §3167, a glycolate class chemical previously developed by Edgewood. Analysis of Edgewood file data had flagged this item as possessing unusual potential as an incapacitant, strongly suggesting the possibility of

2. The Soviets were known to be actively working in the glycolate area. Edgewood had partially investigated EA §3167 and found it to be effective in animals. In addition, there had been several laboratory accidents in which the agent had produced prolonged psychotic effects in laboratory personnel.

3. Since the routes of potential threat to U.S. VIP's and other key personnel, it was highly desirable that existing data on in humans previously acquired by Edgewood be extended to include the Simultaneously, plans were developed to implement countermeasures as required.

4. Preliminary laboratory work was undertaken to determine the solubility and of §3167. Additional work was undertaken to develop laboratory tests to identify the agent in blood. Further work was carried out on the masking effects of such common medicinals as aspirin, barbiturates, etc. The agent was found A good solvent was discovered. A detection test for §3167 was developed, but barbiturates were found to completely mask its presence.
Twenty human volunteer subjects, five prisoners (Holmesbury State Prison, Holmesbury, Pa.) and fifteen military volunteers in the "Hedon" program were tested. Both the treatments were found to be effective with symptoms lasting up to six weeks.

6. Concerning countermeasures, certain

7. In addition to the above project, in 1967, a contract through Edgewood was established for the collection of information on and samples of new psychopharmaceuticals developed in Europe and in 1967, and in 1969. The focus was on unpublished data and unusual new developments. Agency support of this action consisted of a pre-existing contract between Edgewood and for the collection of information on foreign chemical and pharmaceutical developments. Agency redirection, beginning in 1967, consisted of focusing on psychoactive drugs and on the collection of samples.

8. Agency support of both the clinical testing of EA #3167 and of the collection of information on and samples of foreign developments was terminated in January 1973. The transferred to Edgewood in 1972 for an enlarged foreign collection effort was withdrawn in January 1973. Expenditures for the human testing program were gradually reduced as subjects were cleared from the program during the necessary post-test follow-up observational and examination period. Agency involvement in the above activities was closely held at all times.