Waste Acid Detoxification and Reclamation: Phase I - Project Planning and Concept Development

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WASTE ACID DETOXIFICATION AND RECLAMATION:
PHASE I-PROJECT PLANNING AND CONCEPT DEVELOPMENT

T. L. Stewart
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February 1988

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ABSTRACT

The objectives of the Waste Acid Detoxification and Reclamation project are to develop processes for reducing the volume, quantity, and toxicity of metal-bearing waste acids. The primary incentives for implementing these types of waste minimization processes are regulatory and economic (that is, they meet requirements in the Resource Conservation and Recovery Act and reduce the cost for treatment, storage, and disposal). Two precipitation processes and a distillation process are being developed to minimize waste from fuel fabrication operations, which comprise a series of metal-finishing operations. Waste process acids, such as HF-HNO₃ etch solutions containing Zr as a major metal impurity and HNO₃ strip solutions containing Cu as a major metal impurity, are detoxified and reclaimed by concurrently precipitating heavy metals and regenerating acid for recycle. Acid from a third waste acid stream generated from chemical milling operations will be reclaimed using distillation. This stream comprises HNO₃ and H₂SO₄ which contains U as the major metal impurity. Distillation allows NO₃⁻ to be displaced by SO₄²⁻ in metal salts; free HNO₃ is then vaporized from the U-bearing sulfate stream. Uranium can be recovered from the sulfate stream in a downstream precipitation step. These waste minimization processes were developed to meet Hanford's fuel fabrication process needs. However, precipitation and distillation operations are amenable to processes at other DOE facilities.
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INTRODUCTION

Since 1976, the U.S. Environmental Protection Agency (EPA) has encouraged generators of hazardous waste to prioritize waste management practices as follows: source reduction, recycle, waste stream treatment, and land disposal. This philosophy was incorporated into the 1984 Hazardous and Solid Waste Amendments (HSWA) to the Resource Conservation and Recovery Act (RCRA) that now requires hazardous waste generators to certify that a waste minimization program, which can include source reduction, recycling, or treatment, is being implemented at their facility. In general, the initial response by generators has been to conduct waste audits to better identify the type and sources of waste generated. Subsequent procedural changes within the operations can then be implemented to reduce waste; these changes can be made with low or no capital investment and with a minimal impact on operating costs. To further reduce the volume of waste generated, more capital-intensive measures for source reduction will need to be implemented. These expenditures can be justified based on reduced materials, disposal, and long-term liability costs.

The technology described in this document is a waste source reduction technique that can be applied to metal-bearing acids frequently generated during metal refining, metal finishing, and electroplating operations. The nuclear, aerospace, electronics, and steel and specialty metals industries commonly produce waste acid solutions such as HNO₃, HF, HCl, and H₂SO₄ contaminated with heavy metals such as Zr, Cu, Al, Ni, Cr, and U. Past treatment and disposal practices have included neutralization and discharge to solar evaporation ponds. This disposal technique is no longer environmentally acceptable. A waste management option is to reduce the volume, quantity, and toxicity of the waste acid at the source by concurrently removing heavy metals for reuse or disposal and regenerating acid for recycle.

The objective of the Waste Acid Detoxification and Reclamation (WADR) project is to adapt and demonstrate processes and process equipment that reduce the volume, quantity, and toxicity of metal-bearing waste acids. The goal is to use techniques to detoxify the acid by removing heavy metals and to reclaim
acid for recycle. The overall effect is a reduction of the volume of waste requiring disposal, subsequently reducing the potential hazard to the environment, the cost of raw materials, and the cost for disposal.

The approach to be taken to meet this objective includes conducting bench-scale tests to collect engineering data, designing and installing a pilot-scale system based on these data, and operating the pilot-scale system to demonstrate the process on actual spent acid solutions. The project duration is estimated to be five years (dependent on funding levels). The project was initiated in fiscal year (FY) 1986. At this time (FY 1988), bench-scale testing is near completion and the pilot-scale system has been designed and is being procured.

This project planning document will be used as a planning tool for completion of the work that has been initiated. It includes a description of the technology, waste stream information, by-product information, permit requirements, and cost/schedule information.
TECHNOLOGY DESCRIPTION

The processes being adapted for detoxification and reclamation of waste acids include two precipitation processes to remove heavy metals while concurrently regenerating acid for recycle and a distillation process to regenerate the acid solution for recycle. The waste streams chosen for demonstration of WADR consisted of Zr-bearing HNO₃-HF, Cu-bearing HNO₃, and U-bearing HNO₃-H₂SO₄. These spent acids are generated from chemical etching, stripping, and milling operations, respectively, that are used in the fabrication of fuel rods at the Hanford Site (see section on Waste Stream Information). Candidate methods for detoxification and reclamation of each waste stream were discussed in reports by Wick, Elmore, and Hartley (1983), Weakley (1984), and Stewart and Hartley (1985). These methods included precipitation, electrowinning, resin bed exchange, distillation, and membrane separation. Precipitation processes for Zr and Cu removal and a distillation process to recover HNO₃ and HF were chosen based on:

- technical and economic feasibility
- compatibility with disposal scenario
- compatibility with current operating procedures and available space
- availability of commercial equipment.

Further discussion on the candidate methods for detoxification and reclamation of spent acid and a description of the process will be discussed for each of the three streams in the following subsections.

DETOXIFICATION OF Zr-BEARING WASTE ACID

Precipitation methods based on the formation of insoluble fluorozirconates in mixed acid solutions containing 17 g/L HF and 400 g/L HNO₃ are proposed for detoxification of the etching solutions containing Zr. Several Na and Ca compounds can be used to remove Zr from this acid solution; CaF₂, CaO, CaCO₃, Ca(OH)₂, NaF, NaOH, and NaN₃ were identified as candidate precipitants. The use of NaF and NaOH for Zr removal are described in two patents (Megy et al. 1978 and Fenneman 1981, respectively). In addition, two producers of Zr products in the Northwest have experience with the use of NaF precipitation. Both have reported operating problems associated with
solid/liquid separation characteristics of the slurry, and equipment problems resulting from delayed precipitation in pipes, pumps, and storage tanks.

Precipitation of Na₂ZrF₆ using NaF was chosen over other possible candidates because the subsequent reaction did not introduce the potential for fouling from Ca, reduction in the quantity of HF through reaction with Ca, or reduction in HF concentration due to formation of H₂O using NaOH. Although operating problems had been reported using NaF, it was believed that further testing could help identify operating conditions that would avoid these problems. In addition, laboratory tests also indicated that NaF reduced the concentration of dissolved Zr at room temperature and increased the quantity of HF. The precipitation reaction is shown in Equation (1).

\[
H₂ZrF₆ + 2NaF(s) \rightarrow Na₂ZrF₆(s) + 2HF \tag{1}
\]

Bench-scale tests were needed to identify operating conditions such as reaction temperature, stoichiometric addition of NaF, stirring speeds, and length of crystallization period to ensure both acceptable removal of Zr and acceptable solid/liquid separation characteristics of the slurry to avoid the problems experienced by industry.

These bench-scale tests indicated that a reaction temperature of 20°C, stoichiometric addition of NaF at 0.83 g/g Zr, impeller tip speeds of 320 cm/sec, and a 24-h crystallization period could be used. The equipment used in this process included a conical bottom mix tank with a heat exchange unit and low/moderate shear impeller, diaphragm pump, and filter press. All wetted surfaces must be polypropylene, fluoropolymers, or polyvinyl chloride (for low-temperature service). The process includes the following steps: 1) charge the reaction/mix tank with spent acid from the etching operation, 2) add solid NaF using the mixer during reaction, 3) discontinue mixing during the 24-h crystallization period, 4) resuspend the slurry after crystallization, and 5) transfer the slurry to the filter press for solid/liquid separation.

The resulting filtrate can then be recycled as an enriched HF solution, and the zirconate salt (sodium hexafluorozirconate) could be sold to the aluminum industry as a by-product for alloying or could be disposed as a
hazardous waste after neutralization and dewatering. Eventually the concentration of heavy metals other than Zr will exceed acceptable operating levels and the etch solution will require further treatment. This spent solution will be detoxified using the precipitation process and mixed with the spent milling solution for reclamation using distillation.

DETOXIFICATION OF Cu-BEARING WASTE ACID

Methods studied for removing Cu from a solution of HNO₃ at concentrations of 3.5 to 4.5 M at the time of discharge included membrane separation, resin bed exchange, electrowinning, and precipitation. Although effective for separation, ion transfer membranes required large surface contact areas and high capital investment. Resin beds produced large aqueous waste streams, thereby reducing the incentive for detoxification. After initial screening and elimination of membrane and resin bed techniques, electrowinning and precipitation were evaluated in greater detail. Although high-current-density electrowinning is technically feasible in removing Cu metal from solution, it requires lower operating temperatures to adequately reduce the competing chemical dissolution reactions. The cost of a refrigeration system to obtain Cu separation and the low market value of Cu makes electrowinning economically unfeasible. A low-current-density electrowinning process that uses sulfamic acid to reduce the rate of the competing chemical dissolution was described in a patent (Mockrin and Hobin 1977). However, it could not be used because sulfamic acid introduces sulfate ions into the recycled strip solution, which causes premature milling of exposed U metal.

Precipitation reactions, however, can be used to form insoluble Cu compounds such as CuC₂O₄, Cu(OH)₂, CuS, and CuCO₃ which can be separated from solution. The addition of either Na₂C₂O₄ or H₂C₂O₄ to Cu-bearing acid solutions results in the formation of an insoluble copper oxalate. Based on technical considerations, these reactions were chosen over the formation of hydroxide, sulfide, or carbonate precipitants. Although both sodium oxalate and oxalic acid produce a CuC₂O₄ precipitant when added to Cu-bearing HNO₃ solutions, oxalic acid addition allows regeneration of HNO₃ as shown in Equation (2).
The proposed precipitation process has not been used commercially; therefore, bench-scale tests were required to determine operating conditions such as reaction temperature, initial Cu concentration, length of crystallization, use of seed slurry (to improve particle growth rates) to achieve both acceptable Cu removal and acceptable solid/liquid separation characteristics.

These bench-scale tests indicated that reaction temperatures of 60°C, initial Cu concentrations of 100 to 180 g/L, and a crystallization period of 24 h could be used. Furthermore, seed slurry did not improve the rate of particle growth and need not be used.

The equipment used for this process is the same as that used for Zr precipitation: a heated and stirred mix tank, diaphragm pump, and filter press. The process includes the following steps: 1) charge the mix tanks with spent acid from the stripping operation, 2) heat the spent acid, 3) add solid oxalic acid using the mixer during reaction, 4) discontinue mixing during the crystallization period, 5) resuspend the slurry, and 6) transfer the slurry via the pump to the filter press for solid/liquid separation.

As Cu is removed from solution, HNO₃ is regenerated and can, therefore, be recycled to the metal-finishing process. Irreversible dilution of the acid does occur during the metal stripping operation because of conversion of HNO₃ to NOₓ rather than Cu(NO₃)₂. Therefore, at a specified HNO₃ concentration (approximately 3 M HNO₃) the Cu-bearing HNO₃ would require discharge. This solution will be detoxified using the precipitation step and mixed with the spent milling solution for reclamation using distillation. The CuC₂O₄ precipitate will be neutralized and dewatered using absorbent material to meet waste packaging requirements. Heat generated during neutralization will thermally degrade CuC₂O₄ to a less toxic CuO for disposal.

**RECLAMATION OF U-BEARING WASTE ACIDS**

Two methods were considered for reclamation of U-bearing waste acids: membrane separation and distillation with H₂SO₄. Because of technical concerns,
the membrane separation method was eliminated from consideration during the screening process. Distillation with H$_2$SO$_4$ of the mixed acid stream containing 270 g/L HNO$_3$, 450 g/L H$_2$SO$_4$, and 62 g/L U was chosen for reclamtion of HNO$_3$ from free HNO$_3$ and nitrate salts present in the spent milling stream (i.e., U-bearing waste acid). In addition, this technique can be used to reclaim HNO$_3$ and HF-HNO$_3$ from the detoxified etch and strip solutions.

Normally, HNO$_3$ remains in the bottoms when an aqueous acid solution is distilled; however, in the presence of H$_2$SO$_4$, HNO$_3$ will distill to the overhead stream. This technique is used in the HNO$_3$ production industry to bind water and allow nitric acid to be concentrated above its azeotrope at 68.5 wt% HNO$_3$. Typically a mixture of about 50 to 60 wt% H$_2$SO$_4$ and aqueous HNO$_3$ is distilled to produce concentrated HNO$_3$ (90+ wt%) in the overhead vapor. The goal is to adapt this technology for use in reclaiming spent acid, primarily from the chemical milling operation, for recycle. The chemical milling stream contains 33 wt% H$_2$SO$_4$. Distillation of this stream results in an aqueous HNO$_3$ solution with a concentration suitable for metal-finishing operations (i.e., 57 wt%) that is free of impurities, because heavy metals including U, solids, and other contaminants are retained in the H$_2$SO$_4$ bottoms liquid. Uranium can be recovered in a downstream neutralization step and refined for reuse.

As shown in Figure 1, the acid mixture is fed to a HNO$_3$ boiler where water and HNO$_3$ are separated from H$_2$SO$_4$. Heavy contaminants and H$_2$SO$_4$ collect in a bottoms holding tank for later neutralization and separation, while greater than 50% of the original acid solution volume is vaporized to a packed distillation column. Traditional nitric acid rectification occurs in this acid column to concentrate the water and HNO$_3$ that was flashed off the reboiler. Water is vaporized to the condenser and HNO$_3$ collects in the reboiler and is transferred to a bottoms hold tank. The reclaimed HNO$_3$ is suitable for recycle to any of the metal-finishing operations and the distilled water can be used as make-up water for other processes.
FIGURE 1. Proposed Acid Distillation Process
This process can also be used to reclaim the HF-HNO₃ etch solution that can no longer be recycled due to concentration of minor metal impurities. Volatile HF is carried into the column with HNO₃ from the reboiler. To withstand the attack of a highly corrosive ternary mixture of HF-HNO₃-H₂SO₄, standard processing equipment cannot be used. Wetted surfaces of the process equipment must consist of a fluoropolymer suitable to high-temperature mixed-acid conditions. A recycling condenser/scrubber system is also included to retain any volatilized HF. When the concentration of acid in the scrubber water becomes elevated, this stream can also be processed through the distillation system.

DISCUSSION OF OVERALL PROCESS

Precipitation of primary metal contaminants and distillation of HNO₃ are used to detoxify and reclaim acids from the metal-finishing operations used in fuel fabrication. The annual volume of acid requiring neutralization and disposal can be reduced when these technologies are included in the metal finishing process. As shown in Figure 2, the precipitation reactions result

![Projected Volume Reductions](image-url)  

**FIGURE 2.** Projected Volume Reductions
in one-pass volume reductions greater than 90%. Eventually, metal contaminants other than Zr and Cu will concentrate above acceptable operating limits and these process streams will require distillation to recover HNO$_3$ and remove metal contaminants. Depending on the H$_2$SO$_4$ concentration, distillation can reduce the volume of acid discharged for treatment by as much as 60%. As a result the volume and quantity of acid requiring neutralization will be significantly reduced. In addition, removal of the heavy metals from these solutions reduces toxicity.

The 1984 HSWA amendments to RCRA encourage these types of waste minimization efforts as a means of hazardous waste management. Specifically, reduction of hazardous waste at the source, reuse or recovery of useful components, and minimization of the volume and toxicity of waste are emphasized as primary methods for eliminating hazardous wastes. This approach to waste management is supported by environmental groups and is generally considered socially acceptable. Under these guidelines, the WADR processes constitute methods of waste minimization and are therefore both acceptable and encouraged as a waste management practice. The process is not regulated as a treatment process because it uses internal process streams rather than the final waste stream.

In addition to regulatory compliance and social acceptance, WADR is economically justified. Figure 3 identifies projected payback periods based on installed equipment and operating costs and savings from recovered acid and reduced disposal cost. The payback period for Zr and Cu precipitation processes are 0.6 yr and 1.3 yr, respectively. Distillation of the U-bearing H$_2$SO$_4$-HNO$_3$ stream will yield a payback period of 2.2 yr.
FIGURE 3. Projected Payback Periods

Payback Period, yr.

0 1 2 3 4 5

HF-HNO₃ (Zr)  HNO₃ (Cu)  HNO₃·H₂SO₄ (U)

P.P. = \frac{IEC}{A.S · A.C.}
WASTE INFORMATION

The waste process streams chosen for demonstration of the detoxification and reclamation processes are generated at Hanford during nuclear fuel fabrication. Fuel fabrication is a series of metal-finishing operations in which acid solutions are used for chemical milling, etching, and stripping of fuel assembly materials. Currently, the acid solutions are discharged as the concentration of one major metal impurity exceeds acceptable operating limits and then are sent to downstream treatment and disposal, as shown in Figure 4. Until late 1985, the neutralized wastes were discharged to solar evaporation ponds.

When discharged, the process streams contain a high concentration of acid, a high concentration of one major metal impurity, and very dilute concentrations of several minor metal impurities introduced from the metal alloys used in fuel assemblies. Specifically, the chemical milling solution contains $\text{HNO}_3-\text{H}_2\text{SO}_4$, with $\text{U}$ as the major metal impurity. The etch solution contains $\text{HF}-\text{HNO}_3$ with $\text{Zr}$ as the major metal impurity, and the strip solution contains $\text{HNO}_3$ with $\text{Cu}$ as the major metal impurity. The average annual composition and quantity of the three process streams are shown in Table 1.

![Figure 4. Schematic of Current Fuel Fabrication Process](image-url)
TABLE 1. Average Annual Composition of Discharged Metal-Finishing Process Streams

<table>
<thead>
<tr>
<th>Stream</th>
<th>$\text{HNO}_3$</th>
<th>$\text{H}_2\text{SO}_4$</th>
<th>HF</th>
<th>U</th>
<th>Zr</th>
<th>Cu</th>
<th>$\text{H}_2\text{O}$</th>
<th>Volume, L/yr</th>
</tr>
</thead>
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<tr>
<td>Chemical Milling</td>
<td>19.9</td>
<td>33.2</td>
<td>--</td>
<td>4.6</td>
<td>--</td>
<td>--</td>
<td>42.3</td>
<td>129,700</td>
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<tr>
<td>Zr-Etching</td>
<td>40.5</td>
<td>--</td>
<td>2.0</td>
<td>--</td>
<td>3.6</td>
<td>--</td>
<td>53.8</td>
<td>161,000</td>
</tr>
<tr>
<td>Cu-Stripping</td>
<td>20.7</td>
<td>0.3</td>
<td>0.5</td>
<td>--</td>
<td>--</td>
<td>9.4</td>
<td>68.6</td>
<td>185,100</td>
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(a) Process streams contain trace quantities of Cr, Si, Fe, Ni, Al, etc.

Approximately 750,000 L/yr of waste slurry is generated after neutralizing the waste acid and then disposal separating the residual solids. Uranium is recovered from the chemical milling solution as a sodium diuranate after neutralization. The solids, primarily precipitated metal salts, are drummed and sent to a landfill, whereas the liquid wastes are transported to double-shell tanks for storage. Grouting of the liquid waste slurry has been proposed to reduce the mobility of nitrates, fluorides, and metals. Treatment of these streams must meet RCRA and Washington State regulations (WAC 173-303). However, the minimization processes proposed in this project, detoxification and reclamation, are not regulated because they are internal recycle operations that avoid waste generation.

There were several reasons for selecting the waste acid process streams from fuel fabrication for use in the WADR demonstration:

- The volume of the neutralized acid discharged from the process is on the order of $10^6$ L/yr and will contribute to the load on double-shell tank storage space. Therefore, reduction in the volume of this waste would have a positive impact.

- These wastes contain nitrates that are mobile in the environment and will require additional considerations or constraints on performance criteria for the final waste form. Therefore, it would be desirable to reduce or eliminate nitrates from the waste stream.
Fuel fabrication streams are amenable to detoxification and reclamation processes because their composition is relatively constant and the volume for each type of stream is great enough (i.e., on the order of 200,000 L/yr) to make this approach feasible.

Because HNO₃ is commonly used in the nuclear industry, demonstration of a method for reducing the volume of nitrate wastes would be transferrable to or supportive of the development of similar efforts for other waste process streams at DOE facilities. In addition to Hanford, low-level nitrate wastes are generated at Rocky Flats, Oak Ridge, Los Alamos, and Fernald. Approximately 10⁴ to 10⁵ kg NO₃⁻-N/yr are generated at each facility (Johnson et al., 1982). Similar types and volumes of nitrate wastes are also generated at Savannah River Plant, where fuel fabrication operations also exist. Distillation with H₂SO₄ could be considered at facilities where nitric acid reclamation through distillation was discounted because of expected impurity problems. In the proposed process, contaminants are separated from the acid rather than concentrated with the HNO₃. Facilities recovering nitric acid by other methods or treating nitrate wastes from nitric acid processes could consider acid distillation as an alternative.
BY-PRODUCT INFORMATION

The waste acid detoxification and reclamation process produces two new by-product streams and a waste stream with reduced volume. A solid fluorozirconate salt (Na$_2$ZrF$_6$) is produced from detoxification of the etching solution and a copper oxalate ($\text{CuC}_2\text{O}_4$) precipitate is formed from detoxification of the stripping solution. Treatment of these precipitants is discussed below. A neutralized sulfate waste stream contaminated with metals such as Si, Fe, Ni, Cr, and Al is also produced but with much less volume than the current neutralized waste stream. This process waste stream is stored in double-shell tanks until final disposal as a grout to minimize transport of residual metal constituents.

The Zr and Cu precipitates must be neutralized by mixing with solid NaOH to retard any residual acid. Heat generated during neutralization will thermally degrade CuC$_2$O$_4$ to a less toxic CuO. The Na$_2$ZrF$_6$ is used in the aluminum industry for alloying and could be sold for this purpose if generated from a process that did not contact nuclear material. After neutralizing, the precipitates can be disposed to secure landfills after proper packaging with absorbent materials. In the future, these solids may be considered radioactive mixed wastes. The current, applicable regulations for treating and disposing of these waste solids may include 40 CFR 261 (RCRA), WAC 173-303, and DOE 5820.2.
PERMIT REQUIREMENTS

The waste acid detoxification and reclamation process uses fuel fabrication internal process streams rather than waste streams. Therefore, WADR is not regulated as a treatment process and does not require special permits.
COST/SCHEDULE INFORMATION

The WADR concept will be developed from bench-scale evaluation through pilot-scale demonstration over a five year period, depending on availability of funding. Five tasks have been identified: 1) program management to provide coordination and reporting, 2) bench-scale studies to adapt precipitation and distillation techniques for waste acid detoxification and reclamation, 3) engineering development and design to determine process equipment requirements, 4) pilot-plant procurement and installation, and 5) pilot-plant operation to determine performance and economics of the proposed process. The overall cost is estimated as $1,660K for operation plus capital equipment costs of $59K, $120K, and $115K for FY 1987, 1988, and 1989, respectively. The work breakdown and associated costs are listed below.

FY 1986

Cost: $170K
Activity: Complete bench-scale precipitation tests and design of the bench-scale distillation test loop.
Status: Completed.

FY 1987

Cost: $250K
Activity: Conduct bench-scale distillation and materials of construction tests. Design pilot-plant equipment.
Status: Completed first phases of 100-day tests on construction materials and laboratory-scale distillation tests. Completed startup of bench-scale test loop and initiated distillation tests on this unit. Completed pilot-scale distillation equipment design.
FY 1988 (Current Fiscal Year)

Cost: $250K

Activity: Complete bench-scale distillation tests. Procure distillation pilot-plant equipment and continue pilot-plant design. Initiate installation.

Status: Continued bench-scale distillation tests and initiated procurement of pilot-plant equipment. Pilot-plant design is extended through FY 1989 because of lower than expected funding levels in FY 1988.

FY 1989

Cost: $440K

Activity: Complete startup of distillation pilot plant and initiate distillation demonstration. Design, procure, and initiate installation of precipitation pilot plant equipment.

FY 1990

Cost: $450K

Activity: Complete distillation and precipitation pilot plant operation. Complete economic evaluation and technology transfer documentation.
REFERENCES


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