Solstice: Technical Architecture

Computer Vision + Multi-LLM Pipeline for Medical Document Verification

1 Technical Implementation

The Solstice pipeline processes medical documents through computer vision-based layout detection followed by multi-stage LLM analysis. Each stage builds on the previous, creating a traceable verification chain from raw PDF to final evidence report.

2 Pipeline Architecture

- 1. **Layout Detection**: Detectron 2 model identifies bounding boxes for text blocks, tables, figures, and captions
- 2. Content Extraction: Parser converts detected regions into structured JSON with text content and metadata
- 3. Claim Processing: User-provided claims are processed through orchestrated LLM pipeline
- 4. LLM Execution: Four specialized LLM calls analyze documents with specific prompts
- 5. Result Caching: All intermediate outputs stored in hierarchical cache structure for reproducibility

3 Implementation Details

3.1 Layout Detection Pipeline

The layout detection uses Detectron2 with ResNet-50 backbone. The model outputs bounding boxes with class labels:

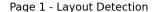
The detection pipeline:

- Processes PDFs page-by-page at 400 DPI resolution
- Runs inference with confidence threshold of 0.2
- Merges overlapping boxes to reduce duplicates
- Outputs JSON with bbox coordinates and class labels

3.2 Multi-Stage LLM Processing

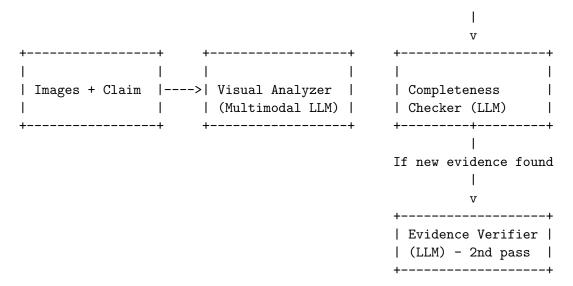
The claim verification uses four LLM calls (Claude-3.5-Sonnet):

++	+	+ +	+
1	1	1 1	1
Claim + Document -	> Evidence	> Evi	dence
1	Extractor	(LLM) Ver	ifier (LLM)
++	+	+ +	



Total elements: 15 Reading order: 15 elements Vaccine 29 (2011) 7733-7739 Contents lists available at ScienceDirect Vaccine 2. Text nal homepage: www.elsevier.com/locate/vaccine reotective efficacy of a trivalent recombinant hemagglutinin protein vaccine (FluBlok®) against influenza in healthy adults: A randomized, placebo-controlled إر<mark>انية المجتبعة المباركة الم</mark> Peter Patriarca^g, Manon Cox^e ity of Rochester, Rochester, NY, United States Baylor College of Medicine, Houston, TX, United States
University of Maryland, Baltimore, MD, United States
St. Louis University, St. Louis, MO, United States Protein Sciences Corporation, Meriden, CT, United States Blair & Co. Greenwich, CT. United States Biologics Consulting Group, Inc., Bethesda, MD, United State ARTICLE INFO TitlerRACT Article history Background: Development of influenza vaccines that do not use embryonated eggs as the substrate for Received 20 May 2011 Received in revised form 27 July 2011 Accepted 27 July 2011 vaccine production is a high priority. We conducted this study to determine the protective efficacy a recombinant, baculovirus-expressed seasonal trivalent influenza virus hemagglutinin (rHA0) vaccine Available online 9 August 2011 Methods: Healthy adult subjects at 24 centers across the US were randomly assigned to receive a single injection of saline placebo (2304 subjects), or trivalent FluBlok containing 45 mcg of each rHA0 component Keywords. (2344 subjects). Serum samples for assessment of immune responses by hemagglutination-inhibition (HAI) were taken from a subset of subjects before and 28 days after immunization. Subjects were followed Influenza vaccine Baculovirus expression Recombinant protein during the 2007-2008 influenza season and combined nasal and throat swabs for virus isolation were obtained from subjects reporting influenza-like illness. Clinical trials Results: Rates of local and systemic side effects were low, and the rates of systemic side effects were similar in the vaccine and placebo groups. HAI antibody responses were seen in 78%, 81%, and 52% of FluBlok recipients to the H1, H3, and B components, respectively. FluBlok was 44.6% (95% CI, 18.8%, 62.6%) effective in preventing culture-confirmed influenza meeting the CDC influenza-like illness case definition despite significant antigenic mismatch between the vaccine antigens and circulating viruses. Conclusions: Trivalent rHAO vaccine was safe, immunogenic and effective in the prevention of culture confirmed influenza illness, including protection against drift variants © 2011 Elsevier Ltd. All rights reserved. hat might also be responsible for pandemic influenza. It is . Title oduction usually necessary to adapt candidate vaccine viruses for high yield OFFICE URL WITH THE PROPERTY OF THE PROPERTY O growth in eggs, a process that can be time consuming, is not always effective influenza vaccines for many years, this system does have successful, and which can select receptor variants that may not be several important drawbacks. Vaccine manufacturing using eggs sentative of circulating influenza strains [1 requires specialized facilities, and the ability to scale up egg proession of proteins in insect cells using recombinant bac duction rapidly in response to an emergency is limited. In addition ulovirus has emerged as a promising technology for vaccine poultry are potentially vulnerable to the same subtypes of influenza production. New recombinant baculoviruses can be generated quickly from sequence data, protein expression is very efficient under the control of the baculovirus polyhedrin promoter, and post translational modifications of the protein are generally similar to Text http://dist.gov/Identifier: NCT00539981.
Corresponding author at: University of Rochester Medical Center, Room 3-6308 other eukarvotic systems. In previous studies, we have evaluated baculovirus-expressed recombinant influenza virus hemagglu-601 Elmwood Avenue, Rochester, NY 14642, United States. Tel.: +1 585 275 5871 tinins (rHA0s) as influenza vaccines in humans. Monovalent and fax: +1 585 442 9328. @urmc rochester edu (II Treanor) bivalent rHA0s have been well tolerated and immunogenic in IextiX/\$ - see front matter © 2011 Elsevier Ltd. All rights reserved. doi:10.1016/j.vaccine.2011.07.128

Figure 1: Detectron2 output showing bounding box detection on clinical trial paper. Colors indicate detected classes: text (blue), table (green), figure (red), caption (yellow).



1. Evidence Extractor:

- Input: Claim text + full document content
- LLM extracts quotes that support the claim
- Returns JSON with quotes and relevance explanations

2. Evidence Verifier:

- Input: Extracted evidence + original document
- Validates quote accuracy and relevance
- Checks for missing context or cherry-picking

3. Completeness Checker:

- Input: All previous LLM outputs
- Searches for additional supporting evidence
- If found, new evidence is passed to Evidence Verifier

4. Visual Analyzer:

- Input: Extracted tables/figures as PNG images + claim text
- Uses multimodal LLM to analyze if image supports the claim
- Returns binary support decision with extracted evidence

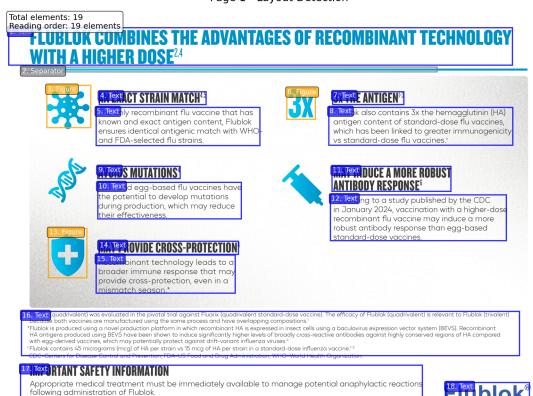
3.3 Cache Structure

All outputs are stored in a hierarchical cache:

```
data/cache/{document_name}/
    extracted/
        content.json  # Structured text content
        document.md  # Markdown representation
        figures/  # Extracted images
    agents/  # Named for legacy reasons, stores LLM outputs
        claims/
```

3.4 Marketing Pipeline Adaptation

Marketing materials require modified processing due to different layout patterns:



Page 1 - Layout Detection

Figure 2: Marketing layout detection shows emphasis on visual hierarchy and promotional elements. The pipeline adapts confidence thresholds for better detection of design-heavy layouts.

nza Vaccine

Key differences:

• Lower confidence threshold (0.1 vs 0.2) for creative layouts

Please see additional Important Safety Information throughout. Before administration, please see

- Additional post-processing for overlapping design elements
- Separate cache directory (data/marketing_cache/)
- Same LLM prompts used for both document types