

**2 July 2026**  
**401-26**

## Administrative Assessment Report – Application A1357

Steviol glycosides (rebaudioside M and rebaudioside D) produced by enzymatic conversion using enzymes produced by *Escherichia coli* BL21

### 1. Application details

<b>Date received:</b> 8 May 2026 <b>Date due for completion of administrative assessment:</b> 29 May 2026 <b>Date completed:</b> 26 May 2026		
<b>Applicant:</b> Adorvia Biotechnology Co., Ltd, a subsidiary of Abiochem Biotechnology Co., Ltd		<b>Potentially affected Schedules</b> 3 and 18
<b>Brief description of application:</b> To seek approval for rebaudioside M and rebaudioside D (steviol glycosides) produced by enzymatic conversion using enzymes derived from <i>Escherichia coli</i> BL21 (DE3)		
<b>Procedure:</b> General Level 1	<b>Maximum total variable hours:</b> 180  <b>Reasons why:</b> The application is seeking a pre-market safety assessment of an already permitted food additive, produced using different enzymes, requiring a safety assessment of minor complexity.	<b>Estimated start date for assessment:</b> June 2026

### 2. Decision

<b>Application accepted</b> <b>Decision Date:</b> 26 May 2026 <b>If fees for ECCB are not received, date of rejection:</b> 25 June 2026
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### Consultation & assessment timeframe

<b>Proposed length of public consultation period:</b> 6 weeks	
<b>Proposed timeframe for assessment</b>	
<b>Notification date:</b> 2 July 2026	
Commence assessment (clock start)	Mid-June 2026
Public comment	Early November 2026
Board to consider approval	March 2027
Notification to Food Ministers' Meeting (FMM)	Mid-April 2027
Anticipated gazettal if no review requested	Late May 2027