

European Commission



**Draft Renewal Assessment Report prepared according to the Commission
Regulation (EU) N° 1107/2009**

BLOOD MEAL

Volume 3 – B.1 (AS)

Rapporteur Member State: Austria
Co-Rapporteur Member State: Lithuania

Version History

When	What
2018/02	Original dossier submission by applicant
2018/04	Revised dossier submission by applicant
2018/12	Draft RAR by RMS AT
2019/02	Draft RAR by RMS AT after commenting by Co-RMS LT

Table of contents

B.1. IDENTITY	4
B.1.1. IDENTITY OF THE ACTIVE SUBSTANCE	4
B.1.1.1. Common name proposed or ISO-accepted and synonyms	4
B.1.1.2. Chemical name (IUPAC and CA nomenclature).....	4
B.1.1.3. Producer's development code number.....	4
B.1.1.4. CAS, EEC and CIPAC numbers.....	4
B.1.1.5. Molecular and structural formula, molecular mass	4
B.1.1.6. Method of manufacture (synthesis pathway) of the active substance.....	5
B.1.1.7. Specification of purity of the active substance in g/kg	5
B.1.1.8. Identity and content of additives (such as stabilisers) and impurities.....	5
B.1.1.9. Analytical profile of batches.....	5
B.1.2. REFERENCES RELIED ON.....	5

B.1. IDENTITY**B.1.1. IDENTITY OF THE ACTIVE SUBSTANCE**

B.1.1.1. Common name proposed or ISO-accepted and synonyms	Blood meal
B.1.1.2. Chemical name (IUPAC and CA nomenclature)	
IUPAC (EFSA Journal 2011;9(10):2394)	<i>Not applicable</i>
CA	<i>Not applicable</i>
B.1.1.3. Producer's development code number	<div> <div></div> <div></div> <div></div> <div></div> <div>Telephone No: <div></div></div> <div>Fax No: <div></div></div> <div>E-mail address: <div></div></div> </div>
B.1.1.4. CAS, EEC and CIPAC numbers	
CAS	90989-74-5
EEC	292-731-9
CIPAC	909
B.1.1.5. Molecular and structural formula, molecular mass	
Molecular formula	<i>Not applicable</i>
Structural formula	<i>Not applicable</i>
Molecular mass	<i>Not applicable</i>

B.1.1.6. Method of manufacture (synthesis pathway) of the active substance	CONFIDENTIAL information - data provided separately in Volume 4
B.1.1.7. Specification of purity of the active substance in g/kg	<p>≥ 990 g/kg (haemoglobin)</p> <p>The following quality criteria are applied:</p> <ul style="list-style-type: none"> - Food grade quality blood collected in authorised slaughterhouses, - Destruction of pathogens and protein denaturation occur during blood processing - Blood of porcine origin. <p>Commission Regulation 142/2011, implementing Regulation 1069/2009 laying down health rules as regards animal by-products and derived products not intended for human consumption and implementing Council Directive 97/78/EC as regards certain samples and items exempt from veterinary checks at the border under that Directive, lays down provisions regarding to the quality criteria of such material for use in feed material or in organic fertilizers and soil improvers (rules for process, microbiologic requirements, ...). Regulation 853/2004 lays down specific hygiene rules for food of animal origin too.</p>
B.1.1.8. Identity and content of additives (such as stabilisers) and impurities	
<i>B.1.1.8.1. Additives</i>	CONFIDENTIAL information - data provided separately in Volume 4
<i>B.1.1.8.2. Significant impurities</i>	CONFIDENTIAL information - data provided separately in Volume 4
<i>B.1.1.8.3. Relevant impurities</i>	None
B.1.1.9. Analytical profile of batches	CONFIDENTIAL information - data provided separately in Volume 4

B.1.2. REFERENCES RELIED ON

Please refer to the different Volume 4, confidential parts.