Teetha Teething Gel

NR 01175/0184

UKPAR

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TEETHA TEETHING GEL

NR 01175/0184

LAY SUMMARY

The Medicines and Healthcare products Regulatory Agency (MHRA) granted A Nelson & Co Limited a Homeopathic Marketing Authorisation for the homeopathic medicinal product Teetha Teething Gel (Homeopathic Marketing Authorisation number: NR 01175/0184) on 24 May 2012. This product is available without prescription and can be bought from pharmacies and other outlets.

Teetha Teething Gel is a homeopathic medicinal product used within the homeopathic tradition for the symptomatic relief of teething pain and the symptoms associated with teething, which are sore and tender gums, flushed cheeks and dribbling. These indications are based on homeopathic provings and published scientific literature. The active ingredients in Teetha Teething Gel are Chamomilla recutita 12c, Aconitum napellus 12c and Atropa belladonna 12c.

No new or unexpected safety concerns arose from this application and it was, therefore, decided that a Homeopathic Marketing Authorisation could be granted.

TEETHA TEETHING GEL

NR 01175/0184

SCIENTIFIC DISCUSSION

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INTRODUCTION

The MHRA granted a Homeopathic Marketing Authorisation for the homeopathic medicinal product Teetha Teething Gel on 24 May 2012. This product is on the General Sales List (GSL).

The application was submitted in accordance with Article 16.2 of Directive 2001/83 EC, as amended, under the National Rules Authorisation Scheme. Teetha Teething Gel contains the homeopathic stocks Chamomilla recutita (in a final dilution of 12c), Aconitum napellus (in a final dilution of 12c) and Atropa belladonna (in a final dilution of 12c). It is used within the homeopathic tradition for the symptomatic relief of teething pain and the symptoms associated with teething, which are sore and tender gums, flushed cheeks and dribbling.

Chamomilla recutita, Aconitum napellus and Atropa belladonna are established homeopathic remedies and their traditional use in homoeopathy is well documented. In support of this application to authorise Teetha Teething Gel details of homeopathic provings and published scientific literature have been provided.

PHARMACEUTICAL ASSESSMENT

HERBAL SUBSTANCE: **CHAMOMILLA RECUTITA**

Synonym: **Chamomile**
Part of plant used: **Whole fresh plant**

Manufacture of the Herbal Substance
The Chamomilla recutita is cultivated in the UK. The plants are organically grown without the use of any chemicals or pesticides in fields far away from contamination and pollution. The plants are harvested manually in mid summer, soon after flowering. The stage of vegetation at which the plant is harvested is in accordance with the German Homeopathic Pharmacopoeia (GHP) monograph. The plant material is then kept fresh in water until it is ready to be packed in bags and sent by overnight carrier to the manufacturer of the homeopathic stock ready for inspection and processing the following morning.

Control of the Herbal Substance
Chamomilla recutita is described in the GHP and the applicant refers to the tests specifications mentioned therein. As Chamomilla recutita is described in an official pharmacopoeia, the analytical tests do not require further validation.

Satisfactory Certificates of Analysis for Chamomilla recutita herbal substance have been provided.

Container Closure System
Satisfactory details of the container closure system used to store the herbal substance are provided.

Stability of the Herbal Substance
A shelf-life for the herbal substance is not necessary because it is used in the fresh state.

ACTIVE INGREDIENT (HOMEOPATHIC STOCK): **CHAMOMILLA RECUTITA MOTHER TINCTURE**

Extraction solvents: Ethanol 86 % (m/m)
Purified water
General properties: A golden yellow to yellowish green liquid with a characteristic aromatic odour and taste

Manufacture of Homeopathic Stock
A satisfactory description of the manufacturing process of the homeopathic stock and a corresponding flow diagram have been provided. Chamomilla recutita mother tincture is manufactured according to method 2a of the GHP. The in process controls

are satisfactorily detailed. The ethanol and purified water used to prepare the mother tincture comply with the specifications of the BP and the Ph Eur. Certificates of Analysis for all materials used in the manufacture of the herbal preparation have been provided.

Alternatively, GHP compliant mother tincture is purchased from an appropriately qualified supplier. Satisfactory Certificates of Analysis are provided.

Control of Homeopathic Stock
A satisfactory specification with appropriate tests and limits has been provided for the homeopathic stock.

Appropriate analytical procedures are used to control the quality of the homeopathic stock. Analytical methods have been validated, as appropriate.

Certificates of Analysis have been provided for batches of the homeopathic stock, demonstrating satisfactory compliance with the proposed specifications.

Container Closure System
Satisfactory details of the container closure system are provided and confirmation has been given that all components of the container closure system comply with current legislation relating to plastic materials and articles intended to come into contact with foodstuffs.

Stability of Homeopathic Stock
Stability studies have been carried out and the results support the proposed shelf life of the homeopathic stock.

HERBAL SUBSTANCE: **ACONITUM NAPELLUS**

Synonym: **Monkshood**
Part of plant used: **Whole fresh plant**

Manufacture of the Herbal Substance
The Aconitum napellus is cultivated in the UK. The plants are organically grown without the use of any chemicals or pesticides in fields far away from contamination and pollution. The plants are harvested manually in mid summer, at the start of flowering. The stage of vegetation at which the plant is harvested is in accordance with the German Homeopathic Pharmacopoeia (GHP) monograph. The plant material is then kept fresh in water until it is ready to be packed in bags and sent by overnight carrier to the manufacturer of the homeopathic stock ready for inspection and processing the following morning.

Control of the Herbal Substance
Aconitum napellus is described in the GHP and the applicant refers to the test specifications mentioned therein. As Aconitum napellus is described in an official pharmacopoeia, the analytical tests do not require further validation.

Satisfactory Certificates of Analysis for Aconitum napellus herbal substance have been provided.

Container Closure System
Satisfactory details of the container closure system used to store the herbal substance are provided.

Stability of the Herbal Substance
A shelf-life for the herbal substance is not necessary because it is used in the fresh state.

ACTIVE INGREDIENT (HOMEOPATHIC STOCK):	**ACONITUM NAPELLUS MOTHER TINCTURE**
Extraction solvents:	Ethanol 86 % (m/m) Purified water
General properties:	Greenish yellow tincture after production later changes to a brownish yellow colour

Manufacture
A satisfactory description of the manufacturing process of the homeopathic stock and a corresponding flow diagram have been provided. Aconitum napellus mother tincture is manufactured according to method 2a of the GHP. The in process controls are satisfactorily detailed. The ethanol and purified water used to prepare the mother tincture comply with the specifications of the BP and the Ph Eur. Certificates of Analysis for all materials used in the manufacture of the herbal preparation have been provided.

Alternatively, GHP compliant mother tincture is purchased from an appropriately qualified supplier. Satisfactory Certificates of Analysis are provided.

Control of Homeopathic Stock
A satisfactory specification with appropriate tests and limits has been provided for the homeopathic stock.

Appropriate analytical procedures are used to control the quality of the homeopathic stock. Analytical methods have been validated, as appropriate.

Certificates of Analysis have been provided for batches of the homeopathic stock, demonstrating satisfactory compliance with the proposed specifications.

Container Closure System
Satisfactory details of the container closure system are provided and confirmation has been given that all components of the container closure system comply with current legislation relating to plastic materials and articles intended to come into contact with foodstuffs.

Stability of Homeopathic Stock
Stability studies have been carried out and the results support the proposed shelf life of the homeopathic stock.

HERBAL SUBSTANCE: **ATROPA BELLADONNA**

Synonym: **Nightshade**
Part of plant used: **Whole fresh plant**

The Atropa belladonna is cultivated in the UK. The plants are organically grown without the use of any chemicals or pesticides in fields far away from contamination and pollution. The plants are harvested manually in late summer, at the end of flowering. The stage of vegetation at which the plant is harvested is in accordance with the German Homeopathic Pharmacopoeia (GHP) monograph. The plant material is then kept fresh in water until it is ready to be packed in bags and sent by overnight carrier to manufacture of the homeopathic stock ready for inspection and processing the following morning.

Control of the Herbal Substance
Atropa belladonna is described in the GHP and the applicant refers to the test specifications mentioned therein. As Atropa belladonna is described in an official pharmacopoeia, the analytical tests do not require further validation.

Satisfactory Certificates of Analysis for Atropa belladonna herbal substance have been provided.

Container Closure System
Satisfactory details of the container closure system used to store the herbal substance are provided.

Stability of the Herbal Substance
A shelf-life for the herbal substance is not necessary because it is used in the fresh state.

ACTIVE INGREDIENT (HOMEOPATHIC STOCK): **ATROPA BELLADONNA MOTHER TINCTURE**

Extraction solvents: Ethanol 86 % (m/m)
Purified water
General properties: Brown liquid with a characteristic odour

Manufacture of Homeopathic Stock
A satisfactory description of the manufacturing process of the homeopathic stock and a corresponding flow diagram have been provided. Atropa belladonna mother tincture is manufactured according to method 3a of the GHP. The in-process controls are satisfactorily detailed. The ethanol and purified water used to prepare the mother tincture comply with the specifications of the Ph Eur. Certificates of Analysis for all materials used in the manufacture of the herbal preparation have been provided.

Alternatively, GHP compliant mother tincture is purchased from an appropriately qualified supplier. Satisfactory Certificates of Analysis are provided.

Control of Homeopathic Stock
A satisfactory specification with appropriate tests and limits has been provided for the homeopathic stock.

Appropriate analytical procedures are used to control the quality of the homeopathic stock. Analytical methods have been validated, as appropriate.

Certificates of Analysis have been provided for batches of the homeopathic stock, demonstrating satisfactory compliance with the proposed specifications.

Container Closure System
Satisfactory details of the container closure system are provided and confirmation has been given that all components of the container closure system comply with current legislation relating to plastic materials and articles intended to come into contact with foodstuffs.

Stability of Homeopathic Stock
Stability studies have been carried out and the results support the proposed shelf life of the homeopathic stock.

HOMEOPATHIC MEDICINAL PRODUCT: TEETHA TEETHING GEL

Description and Composition of the Homeopathic Product
The finished product is a colourless, translucent oral gel containing a combination of Chamomilla recutita, Aconitum napellus and Atropa belladonna presented as centesimal dilutions of 12c. The excipients used to manufacture the homeopathic medicinal product are water, glycerol, xanthan gum, xylitol, hypromellose and ethanol. All excipients used are of pharmacopoeial quality, are considered to be compatible with the homeopathic stocks and do not influence the performance of the product. Certificates of Analysis for the excipients have been provided by the suppliers.

Manufacture of Homeopathic Product
A flow diagram outlining the various stages of the manufacturing process and the in process controls is provided. The dilutions of each homeopathic stock are mixed using an appropriate method. The production process is straightforward and considered validated.

Control of Homeopathic Product
The finished product specification is detailed and the tests and limits used were found to be satisfactory for a product of this nature.

Satisfactory details have been provided on all analytical procedures and these analytical procedures are valid.

Certificates of Analysis have been presented for batches of the drug product demonstrating insignificant inter-batch variation.

Container Closure System
The product is presented in a collapsible, sealed aluminium tube with a polypropylene cap incorporating a tamper evident seal and a piercing device used to pierce the tube seal. The tube contains 15g of gel. Where appropriate the components of the primary packaging system comply with current legislation relating to plastic materials and articles intended to come into contact with foodstuffs.

Stability of Homeopathic Product
Stability studies were conducted under ICH conditions on product batches in the container type proposed for marketing. The product has a shelf life of 24 months when stored in an unopened container; this is reduced to 28 days once the container is first opened. This is appropriate when the storage precaution "Do not store above 25 °C" is applied.

Summary of Product Characteristics, Labels and Patient Information Leaflet
The product literature for this product is pharmaceutically satisfactory.

A package leaflet has been submitted to the MHRA along with results of consultations with target patient groups ("user testing"), in accordance with Article 59 of Council Directive 2001/83/EC. The results indicate that the package leaflet is well-structured and organised, easy to understand and written in a comprehensive manner. The test shows that the patients/users are able to act upon the information that it contains.

CONCLUSION
There are no objections to the granting of a Homeopathic Marketing Authorisation from a quality point of view.

NON-CLINICAL AND CLINICAL SAFETY ASSESSMENT

SAFETY OVERVIEW

The applicant has presented an overview of the safety of the three homeopathic stocks and the combination of stocks in the finished product and concluded that it is highly unlikely that they will exert any acute or chronic toxic effects or any genotoxic effects.

The applicant has not provided any new safety data in support of their application. This is justified on the basis that the product is derived from stocks present in licensed medicinal products, diluted to at least 10^{24} and which are not materials of zoological origin.

As this product contains the excipient ethanol, appropriate warnings are included in the product SmPC, PIL and labels.

CONCLUSION

There are no objections to the granting of a Homeopathic Marketing Authorisation from a safety point of view.

CLINICAL ASSESSMENT (NON SAFETY)

LEGAL STATUS

Chamomilla is listed on the General sales List. Aconitum and belladonna are listed on the Prescription Only Medicines (POM) Order 1997. The Order provides an exemption from prescription control for homeopathic ingredients diluted to at least 12X/6C. Additionally, the POM Order Amendment 2003 provides an exemption from POM control for Aconitum and Belladonna at dilutions of 6X (3C) and above. Therefore, at 12C Aconitum and Belladonna are exempt from POM control and it is considered that Teetha Teething Gel is suitable for General Sales.

INDICATION

The applicant has proposed the following indication:

"A homeopathic medicinal product used within the homeopathic tradition for the symptomatic relief of teething pain and the symptoms associated with teething which are sore and tender gums, flushed cheeks and dribbling"

This indication is acceptable.

POSOLOGY AND METHOD OF ADMINISTRATION

The applicant has proposed the following

"For babies aged 3 months and over.
With a clean finger apply a pea sized amount of gel to the sore area on the baby's gums and teeth.
Use every 4 hours for up to 6 times a day.
This product is not recommended for use in babies under 3 months old.
If symptoms worsen or persist after 7 days of using this product, a doctor or qualified healthcare practitioner should be consulted. "

This is acceptable.

EVIDENCE SUPPORTING THE PROPOSED INDICATION

Schedule 1A Parts 1 and 3 of SI 2006 No. 1952 The Medicines for Human Use (National Rules for Homeopathic Products) Regulations 2006 specifies the data that must be provided to support the use of the product in the indications sought.

No new clinical data were submitted and none are required for an application of this type. In support of this application details of homeopathic provings and published scientific literature have been provided. These are adequate evidence to support the indications for which a national rules authorisation is sought.

CONCLUSION

It is considered sufficient evidence has been submitted to justify the homeopathic use of Teetha Teething Gel for the symptomatic relief of teething pain and the symptoms associated with teething, which are sore and tender gums, flushed cheeks and dribbling. There are no objections to the granting of a Homeopathic Marketing Authorisation from a clinical point of view.

OVERALL CONCLUSION AND RISK ASSESSMENT

QUALITY
The quality data submitted with this application are satisfactory.

NON-CLINICAL AND CLINICAL SAFETY ASSESSMENT
No new non-clinical data were submitted and none are required for an application of this type.

CLINICAL ASSESSMENT (NON SAFETY)
The applicant has provided literature references as supporting evidence to fulfil the requirements for this type of application. These references relate to the indications sought and are, therefore, acceptable.

The SmPC, PIL and labelling of the product are satisfactory.

RISK ASSESSMENT AND CONCLUSION
The quality of the product is acceptable and no new non-clinical or clinical safety concerns have been identified. A Homeopathic Marketing Authorisation may be granted.

TEETHA TEETHING GEL

NR 01175/0184

STEPS TAKEN FOR ASSESSMENT

1	The MHRA received an application under the Homeopathic National Rules Scheme (Article 16.2) on 31 May 2008
2	Following standard checks and communication with the applicant the MHRA considered the application valid on 10 June 2008
3	Following assessment of the application and a meeting of the Advisory Board on the Registration of Homeopathic products (ABRH) on 23 September 2008, the MHRA requested further information relating to the dossier
4	The applicant responded to the MHRA's request, providing further information on the dossier on 5 January 2009
5	The information provided in the applicant's response was reviewed by the ABRH and on 27 February 2009 the MHRA requested further information relating to the dossier
6	The applicant responded to the MHRA's request, providing further information on the dossier on 21 December 2009
7	The information provided in the applicant's response was reviewed by the ABRH and on 28 April 2010 the MHRA requested further information relating to the dossier
8	The applicant responded to the MHRA's request, providing further information on the dossier on 5 October 2010
9	Following assessment of the response the MHRA requested further information relating to the dossier on 29 February 2012
10	The applicant responded to the MHRA's request, providing further information on the dossier on 2 March 2012
11	A National Rules Marketing Authorisation was granted on 24 May 2012

SUMMARY OF PRODUCT CHARACTERISTICS

1 NAME OF THE MEDICINAL PRODUCT

Teetha Teething Gel

2 QUALITATIVE AND QUANTITATIVE COMPOSITION

Chamomilla recutita 12c
Aconitum napellus 12c
Atropa belladonna 12c

For full list of excipients, see section 6.1

3 PHARMACEUTICAL FORM

Oral gel.

Colourless translucent gel.

4 CLINICAL PARTICULARS

4.1 Therapeutic indications

A homeopathic medicinal product used within the homeopathic tradition for the symptomatic relief of teething pain and the symptoms associated with teething which are sore & tender gums, flushed cheeks and dribbling.

4.2 Posology and method of administration

For babies aged 3 months and over.

With a clean finger apply a pea sized amount of gel to the sore area on the baby's gums and teeth.

Use every 4 hours for up to 6 times a day.

This product is not recommended for use in babies under 3 months old.

If symptoms worsen or persist after 7 days of using this product, a doctor or qualified healthcare practitioner should be consulted.

4.3 Contraindications

Hypersensitivity to Chamomilla, Aconite or Belladonna preparations or any of the excipients.

4.4 Special warnings and precautions for use

This medicinal product contains small amounts of ethanol (alcohol), less than 100mg per dose.

The use of this product with other medicinal products containing ethanol should be avoided.

Do not use if seal is broken.

A doctor or a qualified healthcare practitioner should be consulted if symptoms worsen or persist for more than 7 days.

4.5 Interaction with other medicinal products and other forms of interaction
None known.

4.6 Fertility, pregnancy and lactation
The use of this product in pregnancy and lactation is not applicable. However, if use in pregnancy and lactation is required, the advice of a doctor should be sought.

Studies on the effects on fertility have not been performed

4.7 Effects on ability to drive and use machines
None known.

4.8 Undesirable effects
There are no known adverse effects. If any adverse effects are experienced, a doctor or pharmacist should be consulted.

4.9 Overdose
None known. Overdose with this medicine is unlikely to constitute a hazard and therefore symptomatic treatment only is necessary.

5 PHARMACOLOGICAL PROPERTIES

5.1 Pharmacodynamic properties
Not applicable.

5.2 Pharmacokinetic properties
Not applicable.

5.3 Preclinical safety data
Not applicable

6 PHARMACEUTICAL PARTICULARS

6.1 List of excipients
Water
Glycerol
Xanthan gum
Xylitol
Hypromellose

Ethanol

6.2 Incompatibilities
None known.

6.3 Shelf life
24 months
Once opened use within 28 days of opening.

6.4 Special precautions for storage
Do not store above 25^0C.

6.5 Nature and contents of container
Pack size 15g.
Collapsible, sealed aluminium tube with a polypropylene cap incorporating a tamper evident seal and a piercing device used to pierce the tube seal.

6.6 Special precautions for disposal
Pierce tube seal with point in top of cap before first use.
Apply with a clean finger to babies/children's gums and teeth.

7 MARKETING AUTHORISATION HOLDER

A Nelson & Co Limited
5-9 Endeavour Way
Wimbledon
London
SW19 8UH

8 MARKETING AUTHORISATION NUMBER(S)

NR 01175/0184

9 DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION

24/05/2012

10 DATE OF REVISION OF THE TEXT

24/05/2012

PATIENT INFORMATION LEAFLET

BACK

- If the medicine shows any signs of deterioration, consult your pharmacist before using it.
- Return any unused medicine to your pharmacist for safe disposal.

6. Further information

What is this medicine

Active Ingredients

Each 1g of oral gel contains: Chamomilla 12c, Belladonna 12c, Aconite 12c

Also contains Water, Glycerol, Xanthan Gum, Xylitol (E967), Hypromellose, Ethanol.

This homeopathic medicinal product contains small amounts of ethanol (alcohol), less than 100mg per dose.

What does the medicine look like

Teetha Teething Gel is a colourless translucent gel available in a 15g tube.

NR authorisation holder & manufacturer is:

A Nelson & Co Limited, 5-9 Endeavour Way, Wimbledon, London, SW19 8UH
NR 01175/0184

This leaflet is available on request in formats suitable for the blind and partially sighted.

You can help to make medicines safer by reporting any side effects to the yellow card scheme at www.mhra.gov.uk/yellowcard. Alternatively you can get a paper yellow card from your GP's surgery or pharmacy, or call free phone 0808 100 3352 (available 10am-2pm Monday-Friday).

Leaflet revision: 02/12 Art: 2293 WR106058

FRONT

A Homeopathic Medicinal Product

nelsons®

Teetha®
Teething Gel

Chamomilla 12c
Belladonna 12c Aconite 12c

The active ingredients in this product are used within the homeopathic tradition for the symptomatic relief of teething pain and symptoms related to teething:

- Chamomilla is traditionally used in homeopathy to help relieve teething pain.
- Belladonna is traditionally used in homeopathy to help relieve flushed cheeks and dribbling.
- Aconite is traditionally used in homeopathy to help with sore and tender gums.

Please read all of this leaflet carefully before giving this medicine, it contains important information.

Please keep this leaflet; you might need to read it again.

1. What is this medicine for

Nelsons Teetha Teething Gel is a homeopathic medicinal product used within the homeopathic tradition for the symptomatic relief of teething pain and the symptoms associated with teething, which are sore and tender gums, flushed cheeks and dribbling.

P2

P3

2. Before giving this medicine

- This medicine may not be suitable for some babies. Ask your doctor or pharmacist before giving it to the baby if you are not sure.
- Do not give this medicine to the baby if he/she is allergic to any of the ingredients, see section 6.
- Unless otherwise directed by your doctor, do not give this medicine to babies under 3 months.
- If the baby is being given any other medicines, including medicines prescribed by your doctor, or any bought without a prescription, talk to your doctor or pharmacist before giving the baby this medicine.
- This homeopathic medicinal product contains small amounts of ethanol (see section 6).
- The use of this product with other medicinal products containing ethanol should be avoided.

3. How to use this medicine

Directions

For babies aged 3 months and over only.

Check tube seal is not broken before first use. If it is do not use the gel.

- Pierce seal with point in top of cap.
- With a clean finger apply a pea sized amount of the gel to the sore area on the baby's gums and teeth.
- Use every 4 hours for up to 6 times per day.
- **Once opened** use within 28 days.
- **Do not** exceed the stated dose.
- **If you are unsure how to use** the gel consult your doctor or pharmacist.
- **If the symptoms worsen or do not show improvement** after 7 days, consult your doctor or qualified healthcare practitioner.
- **If you forget to give this medicine** to the baby, continue with the usual dose at the usual time. It does not matter if you have missed a dose.
- **If you accidentally give the baby too much** gel or if the baby accidentally swallows some gel, consult your doctor or pharmacist as soon as possible.

4. Possible side effects

This medicine is not expected to cause any side effects. However, as with all medicines, side effects can occur. If the symptoms become worse or they do not show improvement after 7 days, of if the baby experiences any adverse or unwanted effects, consult a doctor or pharmacist.

5. Storing this medicine

- Keep all medicines out of the reach and sight of children.
- Do not store above 25ºC.
- Do not use after expiry date on carton and tube.
- Once opened use within 28 days.

LABELLING

Label:

Carton:


nelsons
Teetha
Teething Gel
Chamomilla 12c Belladonna 12c Aconite 12c
15g oral gel
Active Ingredients
Each 1g oral gel contains:
Chamomilla 12c,
Belladonna 12c, Aconite 12c
Also contains Water, Glycerol, Xanthan Gum, Xylitol (E967), Hypromellose, Ethanol.
Precautions
Please read the enclosed leaflet before use.
• Do not give this medicine to the baby if he/she is allergic to any of the ingredients.
• Do not give to babies under 3 months old.
• Contains ethanol please see leaflet for more information.
• Keep all medicines out of sight and reach of children.
• Do not use after expiry date on carton and tube.
• Once opened use within 28 days. Date of first opening
• Do not store above 25°C.
Directions
For babies aged 3 months and over.
Check tube seal is not broken before first use. If it is do not use the gel.
• Pierce seal with point in top of cap.
• With a clean finger apply a pea sized amount of the gel to the sore area on the baby's gums and teeth.
• Use every 4 hours for up to 6 times a day.
• Do not exceed stated dose.
• If symptoms become worse or do not show improvement after 7 days or if the baby experiences any adverse or unwanted effects, consult a doctor or qualified healthcare practitioner.
5 000488 105247
Use by:
Lot No.
A HOMEOPATHIC MEDICINAL PRODUCT used within the homeopathic tradition for the symptomatic relief of teething pain and the symptoms associated with teething, which are sore and tender gums, flushed cheeks and dribbling.
SUGAR FREE
NEW
Teetha
nelsons
Teething Gel
Chamomilla 12c Belladonna 12c Aconite 12c
A HOMEOPATHIC MEDICINAL PRODUCT
Art 2292
WR103157
nelsons
NEW
Teetha
Teething Gel
Chamomilla 12c Belladonna 12c Aconite 12c
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SUGAR FREE
A HOMEOPATHIC MEDICINAL PRODUCT used within the homeopathic tradition for the symptomatic relief of teething pain and the symptoms associated with teething, which are sore and tender gums, flushed cheeks and dribbling
National rules authorisation holder & manufacturer:
A Nelson & Co Limited,
5-9 Endeavour Way,
Wimbledon, London,
SW19 8UH
NR 01175/0184
Copy 02/12
15g oral gel
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A Nelson & Co Limited.